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Interventions for visual field defects in people with stroke (Review)

Pollock A, Hazelton C, Rowe FJ, Jonuscheit S, Kernohan A, Angilley J, Henderson CA, Langhorne P, Campbell P

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Interventions for visual field defects in people with stroke (Review)

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TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
SUMMARY OF FINDINGS	4
BACKGROUND	10
OBJECTIVES	11
METHODS	11
RESULTS	16
Figure 1.	17
Figure 2.	23
Figure 3.	29
DISCUSSION	30
AUTHORS' CONCLUSIONS	34
ACKNOWLEDGEMENTS	35
REFERENCES	36
CHARACTERISTICS OF STUDIES	45
DATA AND ANALYSES	97
Analysis 1.1. Comparison 1 Restitutive interventions versus control, placebo or no intervention, Outcome 1 Visual field.	98
Analysis 1.2. Comparison 1 Restitutive interventions versus control, placebo or no intervention, Outcome 2 Quality of life.	98
Analysis 2.1. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 1 Visual field.	99
Analysis 2.2. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 2 Extended activities of daily living.	100
Analysis 2.3. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 3 Reading.	100
Analysis 2.4. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 4 Quality of life.	101
Analysis 2.5. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 5 Scanning - cancellation.	101
Analysis 2.6. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 6 Adverse events.	102
Analysis 3.1. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 1 Functional Activities of Daily Living.	104
Analysis 3.2. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 2 Visual field.	104
Analysis 3.3. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 3 Extended activities of daily living.	104
Analysis 3.4. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 4 Reading.	105
Analysis 3.5. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 5 Falls.	105
Analysis 3.6. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 6 Quality of life.	106
Analysis 3.7. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 7 Scanning - cancellation.	106
Analysis 3.8. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 8 Adverse events.	106
Analysis 4.1. Comparison 4 Assessment or screening versus control, placebo or no intervention, Outcome 1 ADL.	107
ADDITIONAL TABLES	107
APPENDICES	138
WHAT'S NEW	144
CONTRIBUTIONS OF AUTHORS	145
DECLARATIONS OF INTEREST	145
SOURCES OF SUPPORT	145
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	146
INDEX TERMS	147

[Intervention Review]

Interventions for visual field defects in people with stroke

Alex Pollock¹, Christine Hazelton¹, Fiona J Rowe², Sven Jonuscheit³, Ashleigh Kernohan⁴, Jayne Angilley⁵, Clair A Henderson⁶, Peter Langhorne⁷, Pauline Campbell¹

¹Nursing, Midwifery and Allied Health Professions Research Unit, Glasgow Caledonian University, Glasgow, UK. ²Department of Health Services Research, University of Liverpool, Liverpool, UK. ³Vision Sciences, Glasgow Caledonian University, Glasgow, UK. ⁴Institute of Health & Society, Newcastle University, Newcastle upon Tyne, UK. ⁵Cornwall & Isles of Scilly Primary Care Trust, Newquay, UK. ⁶Police Division, The Scottish Government, Edinburgh, UK. ⁷Academic Section of Geriatric Medicine, ICAMS, University of Glasgow, Glasgow, UK

Contact address: Alex Pollock, Nursing, Midwifery and Allied Health Professions Research Unit, Glasgow Caledonian University, 6th Floor, Govan Mbeki Building, Cowcaddens Road, Glasgow, G4 0BA, UK. alex.pollock@gcu.ac.uk.

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ABSTRACT

Background

Visual field defects are estimated to affect 20% to 57% of people who have had a stroke. Visual field defects can affect functional ability in activities of daily living (commonly affecting mobility, reading and driving), quality of life, ability to participate in rehabilitation, and depression and anxiety following stroke. There are many interventions for visual field defects, which are proposed to work by restoring the visual field (restitution); compensating for the visual field defect by changing behaviour or activity (compensation); substituting for the visual field defect by using a device or extraneous modification (substitution); or ensuring appropriate diagnosis, referral and treatment prescription through standardised assessment or screening, or both.

Objectives

To determine the effects of interventions for people with visual field defects after stroke.

Search methods

We searched the Cochrane Stroke Group Trials Register, the Cochrane Eyes and Vision Group Trials Register, CENTRAL, MEDLINE, Embase, CINAHL, AMED, PsycINFO, and PDQT Database, and clinical trials databases, including ClinicalTrials.gov and WHO Clinical Trials Registry, to May 2018. We also searched reference lists and trials registers, handsearched journals and conference proceedings, and contacted experts.

Selection criteria

Randomised trials in adults after stroke, where the intervention was specifically targeted at improving the visual field defect or improving the ability of the participant to cope with the visual field loss. The primary outcome was functional ability in activities of daily living and secondary outcomes included functional ability in extended activities of daily living, reading ability, visual field measures, balance, falls, depression and anxiety, discharge destination or residence after stroke, quality of life and social isolation, visual scanning, adverse events, and death.

Data collection and analysis

Two review authors independently screened abstracts, extracted data and appraised trials. We undertook an assessment of methodological quality for allocation concealment, blinding of outcome assessors, method of dealing with missing data, and other potential sources of bias. We assessed the quality of evidence for each outcome using the GRADE approach.

Main results

Twenty studies (732 randomised participants, with data for 547 participants with stroke) met the inclusion criteria for this review. However, only 10 of these studies compared the effect of an intervention with a placebo, control, or no treatment group, and eight had data which could be included in meta-analyses. Only two of these eight studies presented data relating to our primary outcome of functional abilities in activities of daily living. One study reported evidence relating to adverse events.

Three studies (88 participants) compared a restitutive intervention with a control, but data were only available for one study (19 participants). There was very low-quality evidence that visual restitution therapy had no effect on visual field outcomes, and a statistically significant effect on quality of life, but limitations with these data mean that there is insufficient evidence to draw any conclusions about the effectiveness of restitutive interventions as compared to control.

Four studies (193 participants) compared the effect of scanning (compensatory) training with a control or placebo intervention. There was low-quality evidence that scanning training was more beneficial than control or placebo on quality of life, measured using the Visual Function Questionnaire (VFQ-25) (two studies, 96 participants, mean difference (MD) 9.36, 95% confidence interval (CI) 3.10 to 15.62). However, there was low or very-low quality evidence of no effect on measures of visual field, extended activities of daily living, reading, and scanning ability. There was low-quality evidence of no significant increase in adverse events in people doing scanning training, as compared to no treatment.

Three studies (166 participants) compared a substitutive intervention (a type of prism) with a control. There was low or very-low quality evidence that prisms did not have an effect on measures of activities of daily living, extended activities of daily living, reading, falls, or quality of life, and very low-quality evidence that they may have an effect on scanning ability (one study, 39 participants, MD 9.80, 95% CI 1.91 to 17.69). There was low-quality evidence of an increased odds of an adverse event (primarily headache) in people wearing prisms, as compared to no treatment.

One study (39 participants) compared the effect of assessment by an orthoptist to standard care (no assessment) and found very low-quality evidence that there was no effect on measures of activities of daily living.

Due to the quality and quantity of evidence, we remain uncertain about the benefits of assessment interventions.

Authors' conclusions

There is a lack of evidence relating to the effect of interventions on our primary outcome of functional ability in activities of daily living. There is limited low-quality evidence that compensatory scanning training may be more beneficial than placebo or control at improving quality of life, but not other outcomes. There is insufficient evidence to reach any generalised conclusions about the effect of restitutive interventions or substitutive interventions (prisms) as compared to placebo, control, or no treatment. There is low-quality evidence that prisms may cause minor adverse events.

PLAIN LANGUAGE SUMMARY

Interventions for visual field defects in people with stroke

Review question

Are there effective treatments for people who have visual field defects after stroke?

Background

Stroke can cause some people (20% to 57% of people with stroke) to lose the ability to see the entire space in front of them - often one complete half of the normal field of vision is lost. These problems with seeing are called visual field defects. Visual field defects can make it difficult for people to function normally - especially moving about freely, avoiding obstacles, reading, driving, and taking part in rehabilitation for other stroke-related problems.

We wanted to see whether treatments for visual field defects could improve stroke survivors' abilities in activities of daily living (our primary outcome of interest), or other (secondary) outcomes. Secondary outcomes that we were interested in included the size of the visual field defect, independence (in functional abilities), quality of life, ability to scan/search for objects, reading ability, balance and falls, depression and anxiety, and adverse events.

Study characteristics

We included 20 studies (involving 547 stroke participants) that investigated the effect of treatments for visual field defects. However, only 10 of these studies compared the effect of a particular treatment with no treatment. Three of these studies investigated a type of eye movement training designed to improve the lost visual field (a 'restitutive' intervention). Four of the included studies investigated the effect of scanning training, which involves training people to 'scan' across the space in front of them and into the 'lost' visual field, in order to better cope with their lost vision (a 'compensatory' intervention). Three of the included studies investigated the effect of wearing a special prism on a pair of glasses, which increases the amount a person can see on their affected side (a 'substitutive' intervention). One of

Interventions for visual field defects in people with stroke (Review)

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the studies investigated the effect of specialised assessment by an orthoptist (a hospital-based vision specialist), compared to standard care.

Search date

We searched for studies up to May 2018.

Key results

Only two studies presented data relating to how treatment can improve stroke survivors' abilities in activities of daily living, and there was a lack of consistency across studies that limited our ability to draw clear conclusions. There was insufficient evidence to draw any conclusions about the effectiveness of restitutive interventions as compared to control. There was low or very low-quality evidence that scanning training may help improve quality of life, but may have no effect on other outcomes (including adverse events). There was low or very-low quality evidence that prisms may have an effect on ability to scan (look) for objects, but may cause a range of minor adverse events (particularly headache) and may have no effect on other outcomes. Limitations with the evidence meant that we could not draw any conclusions about the benefits of assessment interventions.

Quality of the evidence

The quality of the evidence was low to very low, and in general was insufficient to reach conclusions about the effects of interventions for people with visual field defects.

SUMMARY OF FINDINGS

Summary of findings for the main comparison. Summary of findings: Restitutive interventions versus control

Restitutive interventions compared with control for visual field defects in patients with stroke

Patient or population: stroke survivors with visual field defects

Settings: any rehabilitation setting

Intervention: restitutive interventions

Comparison: control, placebo, or no intervention

Outcomes	Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
Functional ability in activities of daily living	(no data)	No studies	Insufficient evidence	
Visual field (TAP border position in degrees of visual angle from zero vertical meridian) After intervention	MD 1.02 (-1.37 to 3.41)	19 (1 study, Kasten 2007)	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> • Risk of bias - study had high ROB for 'other bias' (relating to potential financial interest in the intervention), study had uncertain ROB for allocation concealment and incomplete outcome data • Indirectness - included participants with diagnoses other than stroke • Imprecision - very small study population (n = 19)
Extended activities of daily living	(no data)	No studies	Insufficient evidence	
Reading ability	(no data)	No studies	Insufficient evidence	
Falls	(no data)	No studies	Insufficient evidence	
Quality of life (improved or not improved - derived from percentage of those who reported subjective improvements of vision)	OR 13.00 (2.07 to 81.48)	30* (1 study, Kasten 2007) *The data used in this analysis were derived from 30 of the original 38 participants, which included data from an additional 19 participants with optic nerve injury who had also received the same interventions in a separate (but parallel) trial. Partici-	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> • Risk of bias - study had high ROB for 'other bias' (relating to potential financial interest in the intervention), study had uncertain ROB for allocation concealment and incomplete outcome data • Indirectness - analysis contained data from a subset of participants from a separate trial, who were not relevant to this review • Indirectness - included participants with diagnoses other than stroke

		pants with optic nerve injury do not meet the inclusion criteria for this review.	<ul style="list-style-type: none"> Imprecision - very small study population (n = 19)
Scanning - cancellation	(no data)	No studies	Insufficient evidence

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

CI: confidence interval

MD: mean difference

n: number

OR: odds ratio

ROB: risk of bias

TAP: Tuebingen Automated Perimeter

Summary of findings 2. Summary of findings: Compensative interventions versus control

Compensative interventions compared with control for visual field defects in patients with stroke

Patient or population: stroke survivors with visual field defects

Settings: any rehabilitation setting

Intervention: compensative interventions

Comparison: control, placebo, or no intervention

Outcomes	Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
Functional ability in activities of daily living	(no data)	No studies	Insufficient evidence	
Visual field (Functional field score and relative change in visual field score, combined) After intervention	SMD -0.11 (-0.92 to 0.70) (no significant effect)	95 (2 studies, De Haan 2015 ; Rowe 2010)	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> Risk of bias - one study judged as high risk of bias for at least one domain Inconsistency - one study had baseline differences Inconsistency - I² = 75% Indirectness - studies explored very different compensatory interventions
Extended activities of daily living (Mobility questionnaire and change in Nottingham EADL, combined)	SMD 0.49 (-0.01 to 0.99) (no significant effect)	97 (2 studies, De Haan 2015 ; Rowe 2010)	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> Risk of bias - one study judged as high risk of bias for at least one domain

Interventions for visual field defects in people with stroke (Review)

After intervention				<ul style="list-style-type: none"> Indirectness - outcome measures were very different; for one study outcome was a mobility measure, rather than a general measure of EADL Indirectness - studies explored very different compensatory interventions
Reading ability (Reading speed; various tests) After intervention	SMD 0.26 (-0.05 to 0.58) (no significant effect)	162 (4 studies, Aimola 2011 ; De Haan 2015 ; Rowe 2010 ; Spitzyna 2007)	⊕⊕⊕⊕ low	Reasons for downgrades: <ul style="list-style-type: none"> Risk of bias - three studies judged as high risk of bias for at least one domain Indirectness - studies explored very different compensatory interventions
Falls	(no data)	No studies	Insufficient evidence	
Quality of life (National Eye Institute Visual Function Questionnaire (NEI - VFQ-25) total score) After intervention	MD 9.36 (3.10 to 15.62) (favours compensatory)	96 (2 studies, De Haan 2015 ; Rowe 2010)	⊕⊕⊕⊕ low	Reasons for downgrades: <ul style="list-style-type: none"> Risk of bias - two studies judged as high risk of bias for at least one domain Indirectness - studies explored very different compensatory interventions
Scanning - cancellation (cancellation tests - time to complete) After intervention	SMD -0.01 (-0.40 to 0.39) (no significant effect)	97 (2 studies, Aimola 2011 ; De Haan 2015)	⊕⊕⊕⊕ low	Reasons for downgrades: <ul style="list-style-type: none"> Risk of bias - two studies judged as high risk of bias for at least one domain Indirectness - studies explored very different compensatory interventions
Adverse events (number of participants with reported events during intervention period)	OR 5.18 (0.24 to 112.57) (favours control)	108 (2 studies, De Haan 2015 ; Rowe 2010) (NB. no events recorded in De Haan 2015 , which did not explicitly report adverse events as an outcome measure)	⊕⊕⊕⊕ low	Reason for downgrades: <ul style="list-style-type: none"> Inconsistency - no events from one study, means pooled result was not estimable for that study; large confidence intervals Indirectness - studies explored very different compensatory interventions

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

CI: confidence intervals

EADL: extended activities of daily living

MD: mean difference

NEI-VFQ-25: National Eye Institute Visual Function Questionnaire

OR: odds ratio

SMD: standardised mean difference

Summary of findings 3. Summary of findings: Substitutive interventions versus control

Substitutive interventions compared with control for visual field defects in patients with stroke

Patient or population: stroke survivors with visual field defects

Settings: any rehabilitation setting

Intervention: compensative interventions

Comparison: control, placebo, or no intervention

Outcomes	Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
Functional ability in activities of daily living (Barthel Index) After 4 weeks of treatment	Wearing prisms MD -4.00 (-17.86 to 9.86) (no significant effect)	39 (1 study, Rossi 1990)	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> • Risk of bias - study judged as high risk of bias for at least one domain • Indirectness - included data from participants with neglect • Imprecision - small study population (n = 39)
Visual field (change in visual field area & change in error scores, from baseline) After intervention	Not wearing prisms SMD 0.12 (-0.46 to 0.70) Wearing prisms SMD 1.12 (0.44 to 1.80)	85 (2 studies, Rossi 1990 ; Rowe 2010)	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> • Risk of bias - one study judged as high risk of bias for at least one domain • Indirectness - included data from participants with neglect • Indirectness - studies cannot be combined due to differences in testing (wearing/not wearing prisms)
Extended activities of daily living (Change in EADL from baseline; mobility improvement scores, in Logits) After intervention	Not wearing prisms SMD 0.20 (-0.44 to 0.85) Wearing prisms SMD 0.24 (-0.26 to 0.75)	99 (2 studies, Bowlers 2014 ; Rowe 2010)	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> • Risk of bias - one study judged as high risk of bias for at least one domain • Indirectness - one study outcome was a mobility measure, rather than a general measure of EADL • Indirectness - included participants with diagnoses other than stroke • Indirectness - studies cannot be combined due to differences in testing (wearing/not wearing prisms)
Reading ability	Not wearing prisms MD 2.80 (-7.13 to 12.73) (no significant effect)	45 (1 study, Rowe 2010)	⊕⊕⊕⊕ Low	Reasons for downgrades: <ul style="list-style-type: none"> • Imprecision - small study population (n = 45) • Imprecision - wide confidence intervals
Falls (number of falls)	Wearing prisms	39	⊕⊕⊕⊕ very low	Reasons for downgrades:

Interventions for visual field defects in people with stroke (Review)

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After intervention	OR 1.21, (0.26 to 5.76) (no significant difference)	(1 study, Rossi 1990)		<ul style="list-style-type: none"> Risk of bias - study judged as high risk of bias for at least one domain Indirectness - included data from participants with neglect Imprecision - small study population (n = 39)
Quality of life	Not wearing prisms	43	⊕⊕⊕⊕ Low	Reasons for downgrades: <ul style="list-style-type: none"> Imprecision - small study population (n = 43) Imprecision - wide confidence intervals
(Visual Function Questionnaire (VFQ-25))	MD 8.40 (-4.18 to 20.98)	(1 study, Rowe 2010)		
After intervention	(no significant effect)			
Scanning - cancellation	Wearing prisms	39	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> Risk of bias - study judged as high risk of bias for at least one domain Indirectness - included data from participants with neglect Imprecision - small study population (n = 39) Imprecision - wide confidence intervals
(line cancellation errors)	MD 9.80 (1.91 to 17.69)	(1 study, Rossi 1990)		
After intervention	(favours substitutive)			
Adverse events	OR 87.32 (4.87 to 1564.66)	59	⊕⊕⊕⊕ Low	Reason for downgrades: <ul style="list-style-type: none"> Inconsistency - large confidence intervals Imprecision - data from only one study
(number of participants with reported events during intervention period)	(favours control)	(1 study, Rowe 2010)		

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

EADL: extended activities of daily living

MD: mean difference

OR: odds ratio

SMD: standardised mean difference

VFQ-25: Visual function questionnaire

Summary of findings 4. Summary of findings: Assessment/screening interventions versus control

Assessment/screening interventions compared with control for visual field defects in patients with stroke

Patient or population: stroke survivors with visual field defects

Settings: any rehabilitation setting

Intervention: assessment/screening interventions

Comparison: control, placebo, or no intervention

Outcomes	Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
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Interventions for visual field defects in people with stroke (Review)

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Functional ability in activities of daily living (FIM) After intervention	MD -6.97 (-23.78 to 9.84) (no significant effect)	37 (1 study, Jarvis 2012)	⊕⊕⊕⊕ very low	Reasons for downgrades: <ul style="list-style-type: none"> • Risk of bias - study judged as high risk of bias for at least one domain • Imprecision - small study population (n = 37) • Imprecision - wide confidence intervals
Visual field	(no data)	No studies	Insufficient evidence	
Extended activities of daily living	(no data)	No studies	Insufficient evidence	
Reading ability	(no data)	No studies	Insufficient evidence	
Falls	(no data)	No studies	Insufficient evidence	
Quality of life	(no data)	No studies	Insufficient evidence	
Scanning - cancellation	(no data)	No studies	Insufficient evidence	

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

FIM: Functional Independence Measure

MD: mean difference

BACKGROUND

Description of the condition

Following stroke, a common visual problem is loss of one-half of the visual field in both eyes; this is called hemianopia or hemianopsia. As it affects the same side in both eyes, it is termed a homonymous hemianopia. For example, left hemisphere stroke causes the loss of the nasal field of the left eye and temporal (outer field) of the right eye, resulting in an inability to see to the right of the centre of the field of view. Visual field defects are common following stroke; the prevalence has been reported as being between 20% and 57% of people (Ali 2013; Rowe 2007; Rowe 2009). The extent of the loss within the visual field may vary, from the loss of the entire half of the visual field to the loss of only a portion of the affected half. It has been reported that 70% of those with visual field loss will maintain a small area of central vision (macular sparing) (Kerkhoff 1999).

The association between visual impairment and disability in activities of daily living has been well-established (Wolter 2006). Visual field defects can affect functional ability and quality of life following stroke (Dombovy 1986; Jongbloed 1986). Studies have demonstrated that people with visual field defects have an increased risk of falling (Ramrattan 2001), and that visual field loss is a predictor of poor functional status at discharge from a stroke unit (Kaplan 1982). People report walking into objects, tripping and falling, feeling unsafe, getting lost, and experiencing panic when in crowded or unfamiliar areas (Windsor 2008). Stroke survivors may struggle with reading, and with accomplishing everyday tasks such as shopping and handling their finances (Warren 2009), and they report severe difficulty returning to work, and marked loss of self-confidence (Rowe 2017).

Visual field loss may also impact on a person's ability to participate in rehabilitation, to live in their own home, and on depression, anxiety, social isolation, and quality of life following stroke (Hepworth 2016; Jones 2006). Visual field defects often co-exist with visual neglect or other perceptual problems. Differentiation between visual field defects and visual neglect can be difficult (Jones 2006).

Description of the intervention

There are many different treatment and management approaches available for people with visual field defects. This review considered any intervention that is specifically targeted at improving the visual field defect or improving the ability of the person to cope with the visual field loss.

Treatments for visual field defects can be described as restitution, compensation or substitution (Hämäläinen 2004; Kerkhoff 2000). In addition to these types of treatments, this review also considered assessment and screening interventions that are specifically targeted at people with visual field defects.

These interventions may include, but are not limited to, the following.

- Restitutive interventions: visual field training, contrast sensitivity training, fusional (binocular vision) training.
- Compensatory interventions: saccadic (fast, simultaneous) eye movement training, training in visual search strategies, training eye movements for reading, use of eye blinks or colour cues, training in activities of daily living.

- Substitutive interventions: prisms (Peli 2000; Rossi 1990), eye patches, adapted lighting, magnification, environmental modification.
- Assessment and screening interventions: standardised visual assessment, screening and referral for visual assessment and intervention.

These are all complex interventions and, therefore, there can be substantial variations, even within interventions of the same type. For example, there can be differences between interventions in relation to the mode of delivery (e.g. therapist-delivered, self-directed, or computer-based), personnel involved in delivery (e.g. vision specialists, such as orthoptists; stroke-care rehabilitation professionals, such as occupational therapists), and in the dose of the intervention (amount of training per day, or per week, and length of intervention period).

How the intervention might work

Interventions for visual field defects are proposed to work by either restoring the visual field (restitution); compensating (adapting) for the visual field defect by changing behaviour or activity (compensation); substituting for the visual field defect by using a device or extraneous modification (substitution); or ensuring appropriate diagnosis, referral and treatment prescription through standardised assessment or screening, or both.

Restitution

This includes the biochemical events that help restore functional neural (nervous system) tissue, for example, the reduction of oedema, absorption of blood, restoration of normal neuronal physiology and restoration of axon (part of a nerve cell) transport. In the past, researchers thought that restitutive approaches would have limited effect in visual rehabilitation (Kerkhoff 2000). However, in the last decade, reports have suggested that expansion of the visual field can be achieved with specific interventions after brain or optic nerve injury (Romano 2008; Sabel 2000; Sahraie 2006). Restitutive interventions include those where there is direct training of the impaired function or repetitive stimulation of the visual field. Commercially available treatments, including NovaVision® Visual Restorative Therapy, give people repeated exposure to stimuli targeting a vision deficit in a direct attempt to help activate the brain to restore vision (NovaVision 2009).

Compensation

Compensation aims to improve the mismatch between the skills of those affected and the demands placed on them by their environment by teaching them to compensate using a spared or intact function (Kerkhoff 1999; Kerkhoff 2000; Peli 2000). Interventions include teaching people compensatory visual search or scanning techniques, varying from simple training strategies to interventions using computerised scanning schedules and specially-developed commercially available tools (such as NVT 2009). When describing interventions for visual field defects, the term compensation may be used synonymously with the term adaptation, but we use the term compensation throughout this review.

Substitution

Substitution involves adaptation of visual components that have been lost or disrupted through the use of optical devices or environmental modifications (Kerkhoff 1999; Kerkhoff 2000; Peli 2000).

Optical devices can include prisms, which shift the image received into an area that can be perceived, and typoscopes, which provide a guide for eye movements when reading.

Assessment and screening interventions

These may work by ensuring that the visual field defect is appropriately diagnosed, enabling other interventions to be prescribed. In those who have co-existing visual field defects and visual neglect; determining the action of an effective intervention can be difficult. The co-existence of visual neglect could prevent interventions aimed at the visual field defect from working effectively. For example, teaching scanning techniques to people with visual field defects may help them learn to compensate by scanning to the affected field but may not be as effective in people with co-existing visual neglect.

Why it is important to do this review

The services available to people with visual problems following stroke are presently inconsistent. There are various treatment and management approaches that are available for people with visual field defects. However, these are used to varying degrees in clinical practice (Rowe 2014), and clinicians lack research-based guidance on the choice of treatment (Hanna 2017). There is a recognised need to identify the evidence base for treatments for visual problems following stroke. There is considerable controversy and debate about the effectiveness of restitutive interventions (Horton 2005a; Horton 2005b; Reinhard 2005; Sabel 2006). There are a number of published reviews of the literature relating to interventions for visual problems following stroke. Barrett 2009 and Riggs 2007 provided reviews of visual problems, which included a small section relating to visual field problems after stroke; both of these reviews were limited in their scope and methodological quality. Bouwmeester 2007 completed a systematic review of the effect of one intervention (visual training) on visual field defects in people with brain damage, including stroke. Lane 2008 provided a narrative review of evidence for interventions for visual field loss. Das 2010 provided a narrative review with an emphasis on restitutive interventions, and primarily discussed a range of cohort studies that used localised, repetitive perceptual training aimed at reversing visual field loss induced by cortical damage. Hanna 2017 provided a narrative review of evidence for visual impairments, including visual field defect, summarising evidence from earlier versions of this review and evidence from non-randomised studies. There is agreement amongst these reviews that there is a need for high-quality studies of the effectiveness of interventions for visual field defects. In order to determine the current evidence for the effectiveness of any treatment or management approaches for stroke patients with visual field defects, and to enable appropriate planning and prioritisation of future primary research, it is essential that there is an up-to-date high-quality systematic review of the existing evidence base.

OBJECTIVES

The key objective of this review was to determine the effects of interventions for people with visual field defects after stroke.

Research questions

- Do interventions for visual field defects improve functional ability following stroke?

- Are interventions for visual field defects more effective at improving functional ability in people with a visual field defect only than in those both with a co-existing visual field defect and visual perceptual problems?

Specific objectives

- To determine if in 1) all participants with visual field defects following stroke (with or without visual perceptual problems), 2) those with visual field defects and no visual perceptual problems, and 3) those with co-existing visual field defects and visual perceptual problems:
 - * restitutive interventions are more effective than control, placebo, or no intervention at improving functional ability in activities of daily living;
 - * compensative interventions are more effective than control, placebo, or no intervention at improving functional ability in activities of daily living;
 - * substitutive interventions are more effective than control, placebo, or no intervention at improving functional ability in activities of daily living;
 - * assessment and screening interventions are more effective than control, placebo, or no intervention at improving functional ability in activities of daily living;
 - * any one active intervention is more effective than any other active intervention at improving functional ability in activities of daily living.
- To determine if in 1) all participants with visual field defects following stroke (with or without visual perceptual problems), 2) those with visual field defects and no visual perceptual problems, and 3) those with co-existing visual field defects and visual perceptual problems:
 - * restitutive interventions are more effective than control, placebo, or no intervention at improving secondary outcomes;
 - * compensatory interventions are more effective than control, placebo, or no intervention at improving secondary outcomes;
 - * substitutive interventions are more effective than control, placebo, or no intervention at improving secondary outcomes;
 - * assessment and screening interventions are more effective than control, placebo, or no intervention at improving secondary outcomes;
 - * any one active intervention is more effective than any other active intervention at improving secondary outcomes.
- To explore the relationship between participant characteristics and the effect of interventions aimed at improving functional abilities in activities of daily living using subgroup analysis.
- To make specific recommendations for future research into the effectiveness of interventions for visual field defects based on a knowledge of the existing evidence base.

METHODS

Criteria for considering studies for this review

Types of studies

Randomised controlled trials (RCTs) and randomised controlled cross-over trials (the first phase analysed as a parallel-group trial).

Types of participants

Adult participants (over 18 years of age) after stroke (using the World Health Organization (WHO) definition of stroke, or a clinical definition if not specifically stated; that is, signs and symptoms persisting longer than 24 hours) who have been diagnosed as having a visual field defect.

Where studies included participants with visual field defects due to reasons other than stroke (e.g. traumatic brain injury), in addition to participants with visual field defects due to stroke, we included these studies. We documented the number of participants with each clinical diagnosis, and planned to use this information when exploring heterogeneity.

We defined a visual field defect as a homonymous loss of vision contralateral to the side of the lesion. We accepted a clinical diagnosis of visual field defect. We documented the method of diagnosing a visual field defect.

We excluded participants with monocular visual field defects due to retinal stroke.

Types of interventions

We included any intervention that was specifically targeted at improving the visual field defect or improving the ability of the participant to cope with the visual field loss. We classified interventions as either restitution, compensation, substitution, or assessment and screening (see [Description of the intervention](#)).

We compared interventions with no treatment, placebo, and control, within four specific preplanned comparisons:

- restitutive interventions versus no treatment, placebo, or control;
- compensatory interventions versus no treatment, placebo, or control;
- substitutive interventions versus no treatment, placebo, or control;
- assessment and screening interventions versus standard care.

We considered studies which compared one active intervention with another active intervention within a narrative synthesis. We did not plan to conduct any meta-analyses comparing one active intervention with another active intervention as we anticipated that there would be substantial variation in the interventions, and that it would not make sense to combine the results.

Two review authors (CH, AP) independently classified the types of interventions in each included trial as either restitution, compensation, substitution, or assessment and screening. We anticipated that we might experience some difficulties in the classification of some interventions, in particular, the classification of interventions as either restitutive or compensatory, and had planned to reach consensus through discussion, involving a third review author when necessary. If there was uncertainty about the action of a particular intervention, we planned to carry out sensitivity analyses to explore the effect of removing and including the relevant trial(s). However, the two independent review authors agreed on all classifications and did not require further discussion with a third review author.

Types of outcome measures

Where possible, we assessed the outcome at the end of the intervention period and at a follow-up point (ideally six months after the intervention had finished, but we accepted any follow-up point after the intervention period had finished, documenting the time point).

Primary outcomes

Functional ability in activities of daily living (ADL)

We included the following validated scales: Barthel Activities of Daily Living Index ([Mahoney 1965](#)), Functional Independence Measure (FIM) ([Smith 1990](#)), modified Rankin Scale (mRS) ([Wilson 2002](#)), Katz Index of Activities of Daily Living ([Katz 1963](#)), and Rehabilitation Activities Profile ([Van Bennekom 1995](#)). If more than one of these functional ability scales was reported, we used the scale appearing earliest in our list.

Secondary outcomes

We included the following secondary outcomes. We prestated outcome measurement tools/scales which we anticipated, and planned that if more than one of the scales or measures was reported, we would use the scale appearing earliest in our list. If additional tools/scales were reported, but none from our prestated list, we included these.

- Functional ability in extended activities of daily living (EADL): Nottingham Extended Activities of Daily Living scale ([Nouri 1987](#)), Lawton Instrumental Activities of Daily Living ([Lawton 1969](#)), Frenchay Activities Index ([Holbrook 1983](#)), Rivermead Activities of Daily Living (ADL) score ([Lincoln 1990](#)).
- Reading - reading ability: reading speed (text reading time), reading accuracy (Wide Range Achievement Test (WRAT) ([Wilkinson 2006](#)), Gray Oral Reading test ([Bryant 2011](#)).
- Visual field: visual field outcomes subdivided into 1) gross visual screening: confrontation tests, Harrington Flocks Visual Screener; 2) kinetic perimetry: Goldmann perimetry, Tangent Screen measures; 3) static perimetry: Humphrey Automated Perimetry, Tubinger Automated Perimetry (TAP), High resolution perimetry (HRP). For perimetry outcomes: when more than one measure had been taken with the same instrument we reported border position for the intact visual field and used it for analysis in preference to hit or detection rate.
- Balance: Berg Balance Scale ([Berg 1989](#)), Functional Reach ([Duncan 1990](#)), Get-Up and Go test ([Mathias 1986](#)), Standing Balance test, Step Test, or other standardised balance measure. We did not include measures of weight distribution or postural sway during standing as the relationship between ability to maintain balance and these outcomes is not established.
- Falls: number of reported falls, Falls Efficacy Scale ([Tinetti 1990](#)).
- Depression and anxiety: Hospital Anxiety and Depression scale ([Zigmond 1983](#)), Beck Depressive Inventory ([Beck 1987](#)), General Health Questionnaire ([Goldberg 1979](#)), Geriatric Depression Scale ([Cinnamon 2011](#)).
- Discharge destination or residence after stroke: dichotomous variable - discharged to previous place of residence (i.e. place of residence prior to stroke) or discharged to alternative destination.

- Quality of life and social isolation: EQ5D (Rabin 2001), Health-related quality of life scale (Williams 1999), Quality of Well Being scale (Kaplan 1993), SF36 (Garrett 1993).
- Visual scanning: cancellation techniques.
- Adverse events: any reported adverse events, excluding falls, death.
- Death.

Search methods for identification of studies

See the methods for the Cochrane Stroke Group [Specialised register](#). We searched for trials in all languages and arranged for the translation of trials where necessary.

Electronic searches

We searched the Cochrane Stroke Group Trials Register (May 2018), the Cochrane Eyes and Vision Group Trials Register (May 2018) and the following electronic bibliographic databases:

- Cochrane Central Register of Controlled Trials (CENTRAL; 2015, Issue 1) in the Cochrane Library (accessed May 2018) ([Appendix 1](#));
- MEDLINE Ovid (1950 to 16 May 2018) ([Appendix 2](#));
- Embase Ovid (1980 to 16 May 2018) ([Appendix 3](#));
- CINAHL EBSCO (Cumulative Index to Nursing and Allied Health Literature; 1982 to 16 May 2018) ([Appendix 4](#));
- AMED Ovid (Allied and Complementary Medicine; 1985 to 16 May 2018) ([Appendix 5](#));
- PsycINFO (1967 to 16 May 2018) ([Appendix 6](#));
- ProQuest Dissertations & Theses (PQDT) database (1861 to 22 March 2015) ([Appendix 7](#)).

Searching other resources

In an effort to identify further published, unpublished and ongoing trials we:

- searched the following registers of ongoing trials:
 - US National Institutes of Health Ongoing Trials Register ClinicalTrials.gov (www.clinicaltrials.gov; May 2018) ([Appendix 8](#));
 - World Health Organization International Clinical Trials Registry Platform (apps.who.int/trialsearch; May 2018) ([Appendix 8](#));
 - ISRCTN Registry (www.isrctn.org) (formerly known as the Current Controlled Trials; www.controlled-trials.com) (March 2015);
 - Health Service Research Projects in Progress (wwwcf.nlm.nih.gov/hsr_project/home_proj.cfm) (March 2015);
 - National Eye Institute Clinical Studies Database (clinicalstudies.info.nih.gov/cgi/protinstitute.cgi?NEI.0.html) (March 2015);
 - Stroke Trials Registry (www.strokecenter.org/trials/) (March 2015);
- handsearched the following journals and conference proceedings:

- *Australian Orthoptic Journal* (1959 to August 2018);
- *British Orthoptic Journal* (1939 to August 2003);
- *British and Irish Orthoptic Journal* (2004 to August 2018);
- International Orthoptic Association (IOA) (www.liverpool.ac.uk/orthoptics/research/search.htm) (1967 to August 2018);
- International Strabismological Association (ISA) (1966 to August 2018);
- Proceedings of the European Strabismological Association (ESA) (1969 to August 2018).

We searched the reference lists of included trials and review articles about vision after stroke and contacted experts in the field.

Data collection and analysis

One review author (CH or PC) ran all the electronic searches, downloaded references into bibliographic software, and removed duplicates. One review author excluded any titles which were obviously not related to stroke and vision (one of CH, PC, or AP). We obtained the abstracts for remaining references and two review authors (two of CH, AP, PC, SJ, AK) independently considered each of these abstracts, excluded any studies that were clearly not RCTs or cross-over trials, and excluded any studies where the intervention was not specifically aimed at improving the visual field defect or the participant's ability to cope with the visual field defect. The review authors resolved any disagreements through discussion, involving a third review author where necessary. We obtained the full papers for any studies included at this stage.

Selection of studies

Two review authors independently applied the selection criteria by considering and documenting the type of studies, type of participants, intervention, comparison intervention, and the outcome measures (two of AP, CH, SJ, AK). Each review author classified studies as 'include' or 'exclude'. If there was disagreement between these two review authors, they reached consensus through discussions involving a third review author.

We listed any excluded studies that included participants with visual field defects in the [Characteristics of excluded studies](#) table and provided the reason for exclusion. We did not list studies that were excluded because they included participants who did not have visual field defects (i.e. visual neglect, eye-movement disorders, age-related visual problems) in the [Characteristics of excluded studies](#) table unless the two review authors agreed that there was a clear reason to do so.

Data extraction and management

We used a pre-designed data extraction form to extract data from the included studies. Two review authors (two of AP, CH, SJ) independently documented the following.

- Methods: study design, method of randomisation.
- Participants: number of participants, inclusion criteria, time since stroke, type, nature and location of lesion. We documented the method of diagnosing the visual field defect and the type and extent of the visual field loss; the presence or absence of visual perceptual problems, and the method of diagnosis; and the country of origin of participants. We documented whether the included participants had visual field defects only (no visu-

al neglect), co-existing visual field defects and visual neglect, or whether the participants were a mixed group (some with and some without visual neglect). If there was a mixed group of participants, we documented whether data were available for the visual field defect-only group and the group with co-existent visual field and visual neglect. Where information was available, we documented the presence or absence of eye movement disorders or low vision, accepting a clinical diagnosis of these.

- **Interventions:** description of interventions given to each treatment group including, if relevant, the duration, intensity, frequency and dose. We classified the type of intervention as restitution, compensation, substitution, or assessment and screening; and the type of control as no treatment, placebo, control, or standard care. We documented the professional background of the person providing the intervention (e.g. occupational therapist, orthoptist).
- **Outcomes:** we documented the primary and secondary outcomes relevant to this review. If a study used a number of different methods of measuring the same outcome, we noted the outcome to be used for any subsequent analysis.
- **Notes:** we noted any important confounding variables. If more than two intervention groups were included in the study, we noted the method of including these groups in any subsequent analysis.

In addition, the review authors independently documented, if data allowed, the following demographics of the included participants: age, gender, place of residence, type of stroke, side of stroke, time since stroke, initial visual field defect, and initial functional ability. The review authors resolved any data extraction discrepancies through discussion.

Assessment of risk of bias in included studies

Two independent authors (two of AP, CH, SJ) assessed risk of bias by grading the following domains as 'low risk', 'high risk' or 'unclear risk' of bias for each included study. We documented this within the 'Risk of bias' tables.

Allocation concealment

Studies with adequate concealment included those that used central randomisation at a site remote from the study, computerised allocation in which records were in a locked readable file that could be assessed only after entering participant details, or the drawing of opaque envelopes. Studies with inadequate concealment included those using an open list or table of random numbers, open computer systems, or drawing of non-opaque envelopes. Studies with unclear concealment included those with no or inadequate information in the report.

Blinding

Adequate concealment included studies which stated that a masked outcome assessor was used, and that had masking of participants and key study personnel and did not identify any 'unmasking'. Inadequate concealment included studies that did not use masking of the outcome assessor, personnel, or participants, where there was incomplete masking, or where the report clearly identified that 'unmasking' occurred during the study. We documented concealment as unclear if a study did not state, or if there was insufficient information to judge, whether or not personnel, participants, and outcome assessors were masked. We acknowl-

edged that for some (but not all) interventions for visual field defects, masking of personnel or participants, or both, is not possible, and considered the potential for any lack of blinding of personnel or participants, or both, to introduce bias.

Incomplete outcome data

Studies adequately addressing incomplete outcome data either had: no missing outcome data; missing outcome data that were unlikely to be related to true outcome; missing outcome data that were balanced in numbers across intervention groups, with similar reasons for missing data across groups; a reported effect size (difference in means or standardised difference in means) among missing outcomes that were not enough to have a clinically relevant impact on observed effect size; or missing data that had been imputed using appropriate methods. Studies inadequately addressing incomplete outcome data either had: missing outcome data that were likely to be related to true outcome, with either imbalance in numbers or reasons for missing data across intervention groups; a reported effect size (difference in means or standardised difference in means) among missing outcomes enough to induce clinically relevant bias in the observed effect size; as-treated analysis done with substantial departure of the intervention received from that assigned at randomisation. We documented addressing of incomplete outcome data as unclear if there was insufficient reporting to allow this to be assessed, or if this was not addressed in the report.

Other bias

We assessed a study not to be free of other bias if it was assessed to have at least one important risk of bias, such as: a potential source of bias related to the specific study design used, an extreme baseline imbalance, a claim to have been fraudulent, financial association with the intervention, or some other problem. If there was insufficient information, or the information provided was unclear, we documented the risk of other bias as unclear.

We produced a 'Risk of bias' summary figure to illustrate the potential biases within each of the included studies.

Measures of treatment effect

We used the Review Manager software RevMan 5.3 ([RevMan 2014](#)) to carry out statistical analyses to determine the treatment effect of:

- restitutive interventions (compared to no treatment, control, placebo, or standard care);
- compensatory interventions (compared to no treatment, control, placebo, or standard care);
- substitutive interventions (compared to no treatment, control, placebo, or standard care);
- assessment and screening interventions (compared to standard care).

We used a random-effects model for all statistical analyses. For dichotomous variables we calculated and reported Peto odds ratios (ORs) with 95% confidence intervals (CIs). For continuous data, we calculated the treatment effect using standardised mean differences (SMDs) and 95% CIs where studies used different scales for the assessment of the same outcome, and using mean differences (MDs) and 95% CIs where all studies used the same method of measuring an outcome.

The primary outcome of functional ability in activities of daily living, and secondary outcomes of functional ability in extended activities of daily living, visual field data, balance, depression and anxiety, and quality of life and social isolation comprise either ordinal data from measurement scales, or continuous data. We analysed these as continuous variables.

Where reported outcomes had a measurement scale where a lower value is indicative of a better outcome (e.g. depression and anxiety scales) we multiplied the reported values by -1 so that in all analyses a higher value was indicative of a better outcome.

If studies reported change values and the baseline value was available, we calculated the value at follow-up (change value - baseline value). If studies reported change values and the baseline value was not available, we used these data in meta-analyses but planned sensitivity analyses to investigate the effect of including the data.

We planned to analyse falls, discharge destination, adverse events, and deaths as dichotomous variables.

Unit of analysis issues

We anticipated that the majority of trials would have a parallel-group design in which each individual participant was randomised to one of two, or more, treatment groups. Where studies had two or more active intervention groups eligible for inclusion within the same comparison (against a control, placebo, or no treatment group), we intended to 'share' the control group data between the multiple pair-wise comparisons in order to avoid double counting of participants within an analysis.

If studies used a randomised controlled cross-over design, we planned to analyse data from the first phase only. We did not anticipate that any studies would use a cluster-randomised design.

Dealing with missing data

If an included study did not report a particular outcome, we did not include that study in the analysis of that outcome.

If an included study had missing data (e.g. reported means but not standard deviations for the follow-up data), we took logical steps to enter an assumed value. Such steps included estimating a standard deviation (SD) based on a reported standard error and estimating a follow-up SD based on a baseline value. We performed calculations of SDs from standard errors and P values using methods described in section 7.7.3.3 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We planned to do sensitivity analyses to investigate the effect of entering assumed values. We also contacted authors in an attempt to obtain missing data.

Assessment of heterogeneity

We visually assessed heterogeneity by looking at the extent of overlap of the CIs on the forest plots. We considered the P value, considering that with $P < 0.1$ there was likely to be heterogeneity. We considered the I^2 statistic. We considered $I^2 > 50\%$ as substantial heterogeneity. If $I^2 > 50\%$, we explored the individual trial characteristics to identify potential sources of heterogeneity.

Assessment of reporting biases

We attempted to avoid reporting biases by using a comprehensive search strategy that included searching for unpublished studies

and searching trials registers. We planned to carry out sensitivity analyses to explore the effect of publication type.

Data synthesis

Two review authors (AP, CH, or SJ) independently extracted data from the included trials. One review author (AP) entered the data into RevMan 5.3 (RevMan 2014), and the other review author checked the entries. They resolved any disagreements through discussion, with reference to the original report.

Subgroup analysis and investigation of heterogeneity

We intended to explore heterogeneity by subgroup analyses to investigate the effect of:

- time after stroke (zero to three months, three to six months, more than six months);
- type of visual field defect (homonymous hemianopia, other) (We anticipated that the majority of the participants would have homonymous hemianopia. However, we documented the type of visual field defect and planned subgroup analyses to investigate the effect of including participants with types other than homonymous hemianopia);
- extent of visual field loss if homonymous (complete hemianopia, partial hemianopia, quadrantanopia);
- presence or absence of visual neglect (no visual neglect, all participants with co-existing visual field defects and visual neglect, mixed group of participants some with and some without visual neglect);
- macular sparing, macular splitting field loss;
- type of treatment (e.g. for compensatory interventions: saccadic eye movement, activities of daily living training; for substitutive interventions: prisms, patches, environmental modifications; for assessment and screening: by orthoptist, occupational therapist, doctor).

We planned to use an established method for subgroup analyses (Deeks 2001). We planned to carry out these subgroup analyses when there were six or more studies included in a single analysis, all with sufficient information to determine the subgroups.

Sensitivity analysis

We planned to carry out sensitivity analysis to explore the effect of the following methodological features.

- Allocation concealment: we planned to re-analyse data, excluding trials with inadequate or unclear allocation concealment.
- Masking of outcome assessor: we planned to re-analyse data, excluding trials without or with unclear masking of outcome assessor.
- Missing outcome data: we planned to re-analyse the data, excluding trials with inadequate or unclear methods of dealing with missing outcome data.
- Other bias: we planned to re-analyse the data, excluding trials assessed to have other bias, or unclear as to whether they had other bias.
- Type of intervention: we planned to re-analyse data, excluding trials where the classification of the type of intervention was uncertain.

- Publication type (peer-reviewed journal, conference abstract or proceedings, doctoral dissertation): we planned to re-analyse data including only those trials from peer-reviewed journals.

We planned to carry out these planned sensitivity analyses when there are six or more studies included in a single analysis.

GRADE assessment and 'Summary of findings' tables

We presented the results of the main preplanned comparisons of the review in 'Summary of findings' tables;

- restitutive interventions (compared to no treatment, control, placebo, or standard care): [Summary of findings for the main comparison](#);
- compensatory interventions (compared to no treatment, control, placebo, or standard care): [Summary of findings 2](#);
- substitutive interventions (compared to no treatment, control, placebo, or standard care): [Summary of findings 3](#);
- assessment and screening interventions (compared to standard care): [Summary of findings 4](#).

Within each 'Summary of findings' table, we summarised data for the primary outcome of interest (functional ability in activities of daily living), the six secondary outcomes for which we had identified the greatest volume of evidence in previous versions of this review (visual field, extended activities of daily living, reading ability, falls, quality of life, scanning - cancellation), and any data related to adverse events.

For each of the preplanned comparisons, we assessed quality of the evidence using the GRADE approach ([Guyatt 2011a](#)), considering each of the following criteria.

- Risk of bias due to flawed design or conduct of studies ([Guyatt 2011b](#)).
- Imprecision (e.g. when confidence intervals for treatment effect are wide) ([Guyatt 2011d](#)).
- Inconsistency (e.g. when point estimates vary widely, I^2 is large) ([Guyatt 2011e](#)).
- Indirectness (e.g. variations in participants, interventions, comparisons, and outcomes) ([Guyatt 2011f](#)).
- Publication bias (may be explored with the use of funnel plots and classed as not suspected, suspected, strongly suspected or very strongly suspected) ([Guyatt 2011c](#)).

We documented identified concerns relating to any of the above criteria, and downgraded the level of evidence accordingly (one downgrade for each concern, and a maximum of two downgrades for each of the listed criteria). If there were no downgrades the level of evidence was high quality, if there was one downgrade the level of evidence was moderate quality, if there were two downgrades the level of evidence was low quality, and if there were more than two downgrades the level of evidence was very low quality. We used the following definitions of evidence.

- High quality: when further research is very unlikely to change our confidence in the estimate of effect.
- Moderate quality: when further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
- Low quality: when further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
- Very low quality: when we are very uncertain about the estimate.

One review author (AP) assessed quality of evidence, reported identified concerns, and applied downgrades. Other review authors checked agreement with these judgements and resolved any disagreements through discussion.

RESULTS

Description of studies

See: [Characteristics of included studies](#); [Characteristics of excluded studies](#); [Characteristics of ongoing studies](#), [Characteristics of studies awaiting classification](#).

Results of the search

Results of the search are shown in [Figure 1](#). Our search strategy identified 17,224 records from the main electronic databases, and we identified a further eight studies through our wider search. One review author (AP, CH or PC) eliminated 15,658 titles that were clearly irrelevant, and two review authors (AP, CH) applied the inclusion criteria to the remaining 1574 abstracts, identifying 178 to be considered at full text. Of these, we identified a total of 20 studies for inclusion. In addition, we identified seven ongoing studies (see [Characteristics of ongoing studies](#)), and two studies that require further assessment (see [Characteristics of studies awaiting classification](#)).

Figure 1. Study flow diagram.

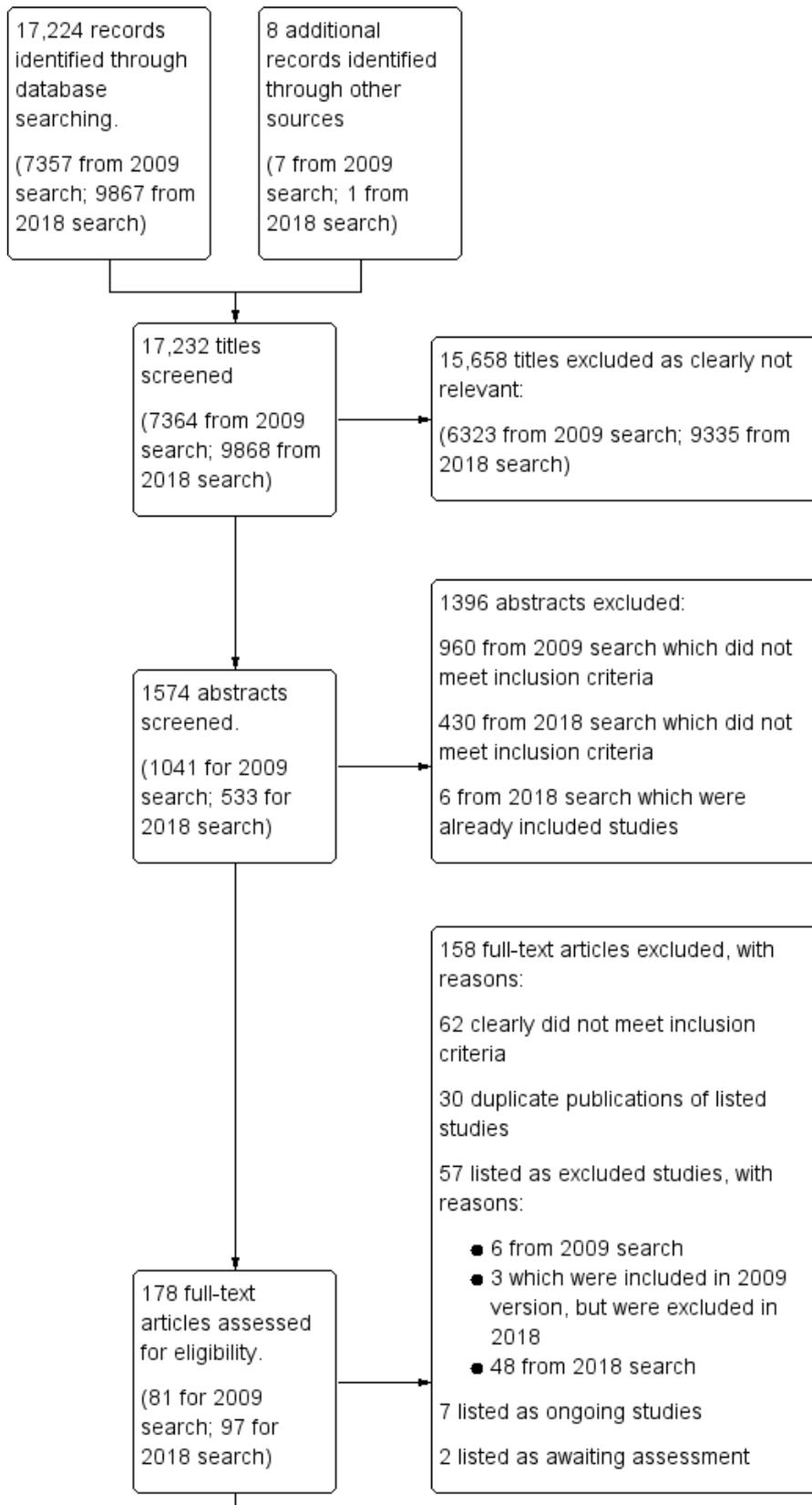
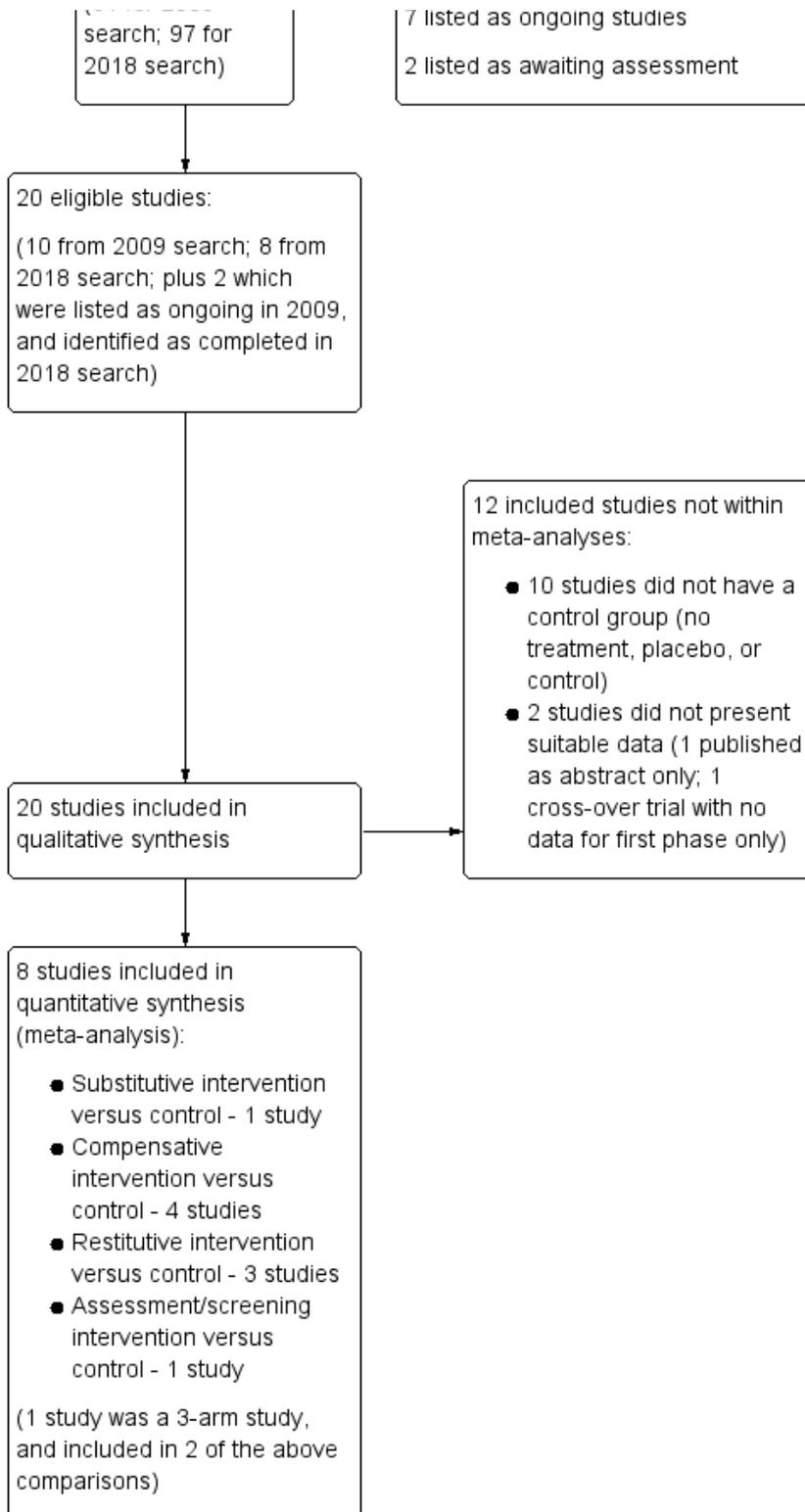


Figure 1. (Continued)



Included studies

We included 20 studies (732 randomised participants, with data available for 638, of whom 547 (85%) had a diagnosis of ischaemic or haemorrhagic stroke) in this updated review (Aimola 2011; Bainbridge 1994; Bowers 2014; De Haan 2015; Elshout 2016; Gall 2013; Jarvis 2012; Jobke 2009; Kasten 1998; Kasten 2007; Keller 2010; Modden 2012; Plow 2010; Poggel 2004; Rossi 1990; Roth 2009; Rowe 2010; Schuett 2012; Spitzyna 2007; Szlyk 2005).

The previous version of this review included 13 studies (344 randomised participants, 285 of whom were participants with stroke) (Bainbridge 1994; Carter 1983; Jobke 2009; Kasten 1998; Kasten 2007; Plow 2010; Poggel 2004; Rossi 1990; Roth 2009; Spitzyna 2007; Szlyk 2005; Weinberg 1977; Weinberg 1979). However, we have excluded three of these studies from this update (Carter 1983, Weinberg 1977; Weinberg 1979: see Excluded studies for reason for these exclusions).

There were only abstracts available for Bainbridge 1994 and Gall 2013. In the previous version, Plow 2010 was based on data from a conference abstract and ongoing trials register data only; however, for this update a full paper was available.

We present a brief overview of the studies below. Full descriptions of the included studies can be found in the Characteristics of included studies table and in Table 1 (Demographics of included studies: settings of included studies), Table 2 (Demographics of included studies: demographics of included participants), Table 3 (Demographics of included studies: visual problems of included participants), and Table 4 (Outcome measures within included studies).

Study design

Fifteen of the included studies were parallel-group randomised controlled trials (RCTs), and five were randomised cross-over studies (Bowers 2014; Elshout 2016; Jobke 2009; Schuett 2012; Szlyk 2005).

Sixteen of the included studies randomised participants to one of two treatment groups; three had three treatment groups (Kasten 2007; Modden 2012; Rowe 2010); and one was a cross-over AB/BA design, where A was an active treatment and B a placebo; however participants were also randomised to receive one of two different active treatments, each of which had a related sham treatment meaning that there were effectively four different treatment groups (active 1, active 2, sham 1 and sham 2) (Bowers 2014).

Comparison versus control

Ten of the 20 included studies had a control (no treatment, standard care, or placebo) group, comparing 10 active treatments with control (Rowe 2010 had two active treatment groups):

- three compared a restitutive intervention with control (Gall 2013; Elshout 2016; Kasten 1998);
- four compared a compensatory intervention with control (Aimola 2011; De Haan 2015; Rowe 2010; Spitzyna 2007);
- three compared a substitutive intervention with control (Bowers 2014; Rossi 1990; Rowe 2010);
- one compared assessment/screening with control (Jarvis 2012).

In Rowe 2010, the two active treatment groups (compensatory and substitutive interventions) were compared with each other.

Ten of the 20 included studies did not have a control group. Nine compared two different active treatments, and one had three active treatment groups (Modden 2012):

- four compared different restitutive interventions (Jobke 2009; Kasten 2007; Plow 2010; Poggel 2004);
- three compared different compensatory interventions (Keller 2010; Modden 2012; Schuett 2012);
- two compared different substitutive interventions (Bainbridge 1994; Szlyk 2005);
- two compared compensatory and restitutive interventions (Modden 2012; Roth 2009).

Interventions studied

Restitutive interventions

Nine studies (239 randomised participants) investigated the effect of restitutive interventions.

In eight of these studies, the restitutive intervention studied was a form of computer-based vision restoration therapy:

- Kasten 1998 and Elshout 2016 compared visual restitution therapy with a placebo intervention;
- Jobke 2009 and Kasten 2007 compared the effectiveness of two (or more) types of visual restitution therapy;
- Plow 2010 explored the effect of adding transcranial direct current stimulation (tDCS) to visual restitution therapy;
- Poggel 2004 explored the effect of adding attentional cueing to visual restitution therapy;
- Modden 2012 compared computerised restitution therapy with two different compensatory interventions;
- Roth 2009 compared 'flicker-stimulation training', which the authors described as a "potential" restitutive intervention, with a compensatory intervention.

In one of these nine included studies, the restitutive intervention studied was a form of non-invasive brain stimulation using alternating current stimulation. This was compared with a placebo intervention (Gall 2013).

Compensatory interventions

Eight studies (347 randomised participants) investigated the effect of compensatory interventions:

- Aimola 2011 compared computer-based compensatory training with a control;
- Spitzyna 2007 compared computer-based reading training ("optokinetic nystagmus inducing reading therapy", involving reading scrolling right to left text) with a control;
- De Haan 2015 compared a compensatory scanning training programme with a control;
- Rowe 2010 compared paper-based visual scanning training with a control (and with a substitutive intervention);
- Keller 2010 compared two types of compensatory training - audiovisual exploratory training and visual exploration training;
- Schuett 2012 compared two types of compensatory training - visual exploration training and reading training;

- [Modden 2012](#) compared computerised scanning training, an occupational therapy compensatory training program, and a computerised restitutive therapy;
- [Roth 2009](#) compared computer-based scanning training with restitutive training.

Substitutive interventions

Five studies (227 randomised participants) investigated the effect of substitutive interventions. In all five studies, the substitutive intervention studied was a type of prism:

- [Rossi 1990](#) compared 15 diopter Fresnel prisms with no prisms;
- [Rowe 2010](#) compared 40 diopter Fresnel prisms with no treatment, and with a compensatory intervention;
- [Bowers 2014](#) compared 57 diopter oblique prism glasses and horizontal prism glasses with five diopter sham prism glasses;
- [Bainbridge 1994](#) compared full-field 15 diopter Fresnel prisms with hemi-field 15 diopter Fresnel prisms;
- [Szlyk 2005](#) compared 20 diopter Fresnel prisms with 18.5 dioptre Gottlieb VFAS (Visual Field Awareness System) prisms.

Assessment and screening interventions

One study (64 randomised participants) investigated the effectiveness of an assessment and screening intervention on relevant outcomes:

- [Jarvis 2012](#) compared the effect of providing therapy staff with information from an orthoptic assessment with no intervention.

Populations studied

The reported diagnoses of the participants within the 20 included studies were as follows:

- 14 studies recruited participants with mixed diagnoses including stroke, trauma, surgery, and infections;
- five studies included participants with stroke only; and
- one study - the cause of the lesion was unclear.

(See details in [Characteristics of included studies](#) and [Table 2](#)). Despite the high number of studies including participants with mixed diagnoses, the majority of participants in the studies in this review did have stroke (across all studies, 85% (520/611) of participants with data reported had a diagnosis of stroke).

Fourteen of the 20 included studies included participants with visual field defects only (no visual neglect); seven of 14 studies clearly stated the method of diagnosis of visual neglect ([Bowers 2014](#); [De Haan 2015](#); [Elshout 2016](#); [Keller 2010](#); [Roth 2009](#); [Rowe 2010](#); [Schuett 2012](#)), four of the 14 studies stated that participants with neglect were excluded but did not state the method of diagnosis of visual neglect ([Kasten 1998](#); [Kasten 2007](#); [Modden 2012](#); [Poggel 2004](#)), and in the three remaining studies it was assumed (but not clearly stated) that participants with visual field defects only were included ([Jobke 2009](#); [Spitzyna 2007](#); [Szlyk 2005](#)). Four of the included studies included participants who had visual neglect in addition to, or instead of, visual field defects ([Aimola 2011](#); [Bainbridge 1994](#); [Jarvis 2012](#); [Rossi 1990](#)). In two studies, it was unclear whether the participants had visual neglect or not ([Gall 2013](#); [Plow 2010](#)). See [Table 3](#).

Visual field measurement

All 20 included studies reported a measurement of the visual field in order to inform participant inclusion or provide baseline information relating to visual field defect, or both. Seven reported one visual field measure ([De Haan 2015](#); [Gall 2013](#); [Jarvis 2012](#); [Modden 2012](#); [Plow 2010](#); [Schuett 2012](#); [Szlyk 2005](#)), 11 reported two visual field measures ([Aimola 2011](#); [Bainbridge 1994](#); [Bowers 2014](#); [Elshout 2016](#); [Jobke 2009](#); [Kasten 1998](#); [Keller 2010](#); [Rossi 1990](#); [Roth 2009](#); [Rowe 2010](#); [Spitzyna 2007](#)), and two reported three visual field measures ([Kasten 2007](#); [Poggel 2004](#)). [Spitzyna 2007](#) reported a second measure only where the first perimetric results had poor reliability.

In three studies, the perimetry equipment was unclear ([Aimola 2011](#); [Gall 2013](#); [Jobke 2009](#)), and in [Rowe 2010](#), either of two types of perimetry was reported (Esterman static programme or Goldmann kinetic).

Visual field measurements were categorised as:

- gross visual screening: five studies reported gross visual screening; two using the Harrington Flocks visual screener ([Bainbridge 1994](#); [Rossi 1990](#)), two using a confrontation method ([Bainbridge 1994](#); [Jarvis 2012](#)), one reporting the visual field assessment from the Test Battery of Attentional Performance ([Modden 2012](#)), and one a subjective topographic measure of perceived visual field defect ([Plow 2010](#));
- kinetic perimetry: 10 studies reported measures of kinetic perimetry; seven used Goldmann perimetry ([Bowers 2014](#); [De Haan 2015](#); [Elshout 2016](#); [Keller 2010](#); [Rowe 2010](#); [Spitzyna 2007](#); [Szlyk 2005](#)), one used Tubingen kinetic perimetry ([Schuett 2012](#)), one used tangent screen measures ([Rossi 1990](#)), and one used an unspecified kinetic perimeter ([Aimola 2011](#));
- static threshold perimetry: eight studies reported static perimetry measures; two used standard automated perimetry ([Gall 2013](#); [Jobke 2009](#)), four used Tubinger automated perimetry (TAP) ([Kasten 1998](#); [Kasten 2007](#); [Poggel 2004](#); [Roth 2009](#)), and two used Humphrey automated perimetry ([Elshout 2016](#); [Spitzyna 2007](#));
- static suprathreshold (inclusive of full field 120, Esterman, campimetry, tangent screen): five studies reported static suprathreshold measures; two reported Esterman measures ([Aimola 2011](#); [Rowe 2010](#)), one reported Humphrey full field 120 ([Bowers 2014](#)), one reported suprathreshold checks on a Goldmann perimeter ([Keller 2010](#)), and one reported high resolution campimetry ([Poggel 2004](#));
- resolution perimetry: six studies reported resolution perimetry measures; five used high resolution perimetry (HRP) ([Jobke 2009](#); [Kasten 1998](#); [Kasten 2007](#); [Plow 2010](#); [Poggel 2004](#)), and one used scanning laser ophthalmoscopy ([Roth 2009](#)).

Sample size

On average, included studies randomised 36 participants (standard deviation, 22 participants) into the trial prior to attrition. This ranged from just 10 participants ([Szlyk 2005](#)), to 87 participants ([Rowe 2010](#)). Only five of 19 studies recruited more than 50 participants: [Aimola 2011](#) (n = 70), [Bowers 2014](#) (n = 73), [De Haan 2015](#) (n = 54), [Jarvis 2012](#) (n = 64), and [Rowe 2010](#) (n = 87). A total of 732 participants were recruited across the 20 included studies, with data available for 638 participants, of whom 547 were stroke patients. See [Table 2](#) for recruitment numbers across all included studies.

Outcome measures

Table 4 summarises the outcome measures within the included studies, and highlights which studies had data which was suitable for inclusion in meta-analyses within this review.

Primary outcome

- Functional ability in activities of daily living. Three studies included a measure of functional ability: Rossi 1990 reported the Barthel Index, Jarvis 2012 reported the FIM, and Modden 2012 reported the extended Barthel Index (German version).

Secondary outcomes

- Visual field. As reported above, all 20 included studies used at least one measure of visual field at inclusion/baseline. However, two of the studies did not measure visual field as an outcome (i.e. measured following treatment) (Bowers 2014; Jarvis 2012); in one cross-over study, while it was measured as an outcome, no measurement was taken immediately after the cross-over (Schuett 2012); while in Aimola 2011 it was unclear whether this was measured as an outcome or just at baseline. The methods of assessing visual field as a study outcome are summarised in Table 4. Visual field outcomes of potential relevance to our meta-analyses included measures of: 1) gross visual screening: four studies (Bainbridge 1994; Rossi 1990; Modden 2012; Plow 2010); 2) kinetic perimetry: seven studies (Aimola 2011; De Haan 2015; Elshout 2016; Keller 2010; Rowe 2010; Spitzyna 2007; Szlyk 2005); 3) static threshold perimetry: eight studies (Jobke 2009; Kasten 1998; Kasten 2007; Elshout 2016; Gall 2013; Poggel 2004; Roth 2009; Spitzyna 2007); 4) static superthreshold perimetry: four studies (Keller 2010; Poggel 2004; Rossi 1990; Rowe 2010); and 5) resolution perimetry: six studies (Jobke 2009; Kasten 1998; Kasten 2007; Plow 2010; Poggel 2004; Roth 2009).
- Extended activities of daily living. Three studies reported a measure of extended activities of daily living (other than a measure of reading ability). Two were measures of functional mobility: Bowers 2014 quantified "Perceived difficulties with mobility" using "a 5-point rating scale (no difficulty to extreme difficulty) for seven situations (items) relevant to homonymous hemianopia, including at home, in stores, outdoors, in unfamiliar areas, in familiar areas, in crowded areas, and noticing objects off to the side when walking"; and De Haan 2015 used an independent mobility questionnaire. Rowe 2010 reported the Nottingham Extended Activities of Daily Living index.
- Reading. Ten studies reported measures of reading ability (Aimola 2011; De Haan 2015; Elshout 2016; Jobke 2009; Keller 2010; Modden 2012; Roth 2009; Rowe 2010; Schuett 2012; Spitzyna 2007). The measures used were the Radner reading test in three studies (De Haan 2015; Jobke 2009; Rowe 2010), tests of reading speed in seven studies (Aimola 2011; De Haan 2015; Elshout 2016; Keller 2010; Roth 2009; Schuett 2012; Spitzyna 2007), and measures linked to the correctness of reading in four studies (De Haan 2015; Jobke 2009; Modden 2012; Schuett 2012). Text reading time data were displayed graphically by Spitzyna 2007 but actual data were confirmed by correspondence with the author.
- Falls. One trial reported the number of falls (Rossi 1990).
- Quality of life. Ten trials reported measures of quality of life (Aimola 2011; De Haan 2015; Gall 2013; Jobke 2009; Kasten 1998; Kasten 2007; Keller 2010; Plow 2010; Roth 2009; Rowe 2010), with several reporting more than one type of measure. The Na-

tional Eye Institute Visual Functioning Questionnaire (NEI-VFQ) was used by five studies (Aimola 2011; De Haan 2015; Gall 2013; Jobke 2009; Rowe 2010), the Short-Form 12 (SF-12) by two studies (Gall 2013; Rowe 2010), the World Health Organization Quality of Life Bref (WHOQOL-Bref) by one (Roth 2009), the Impact of Visual Impairment profile (IVI) and Veterans Affairs Low Vision Visual Function Questionnaire (LV-LFQ) by one (Plow 2010), and the EQ-5D (standardised EuroQol health-related quality of life instrument) by one (Rowe 2010). A number of different, often self-designed, questionnaires of satisfaction, improvement and visual ability levels were also used (see Table 4). The quality of life data from Kasten 1998 were included as a dichotomous variable as the data presented in the published paper were a percentage of those who reported subjective improvements of vision; the results were for both the pre- and post-chiasmal group, with 30 of the 38 participants responding.

- Visual scanning. Eight trials reported measures of visual scanning as assessed by a range of scanning and cancellation techniques (Aimola 2011; Bainbridge 1994; De Haan 2015; Keller 2010; Modden 2012; Rossi 1990; Roth 2009; Schuett 2012). In addition, one reported the visual scan test from the Test Battery of Attentional Performance (Modden 2012), and one also used a video eye tracker (Roth 2009). Reported data included scanning/search time and number of errors or omissions (see Table 4). Where a study reported a range of scanning outcomes, we prioritised measures of scanning time for meta-analyses. De Haan 2015 used three different visual scanning tests, and reported a range of different data; we used the results of the "parallel search test" for all trials (target present and target absent), but explored the impact of using alternative data.

Excluded studies

We excluded 158 papers after assessment of the full paper (see Figure 1). Sixty-two of the 158 clearly did not meet the inclusion criteria, and 30 of the 158 were duplicate publications of excluded studies. Fifty-seven of the 158 papers required more in-depth appraisal prior to exclusion; we have provided our reasons for exclusion of these studies in the Characteristics of excluded studies table. We excluded the majority of these because the intervention was not specifically targeted at the ability of the participant to cope with visual field loss (32/57), or because the study was not randomised (22/57). We found one study did not include participants with stroke, one was focused on central alexia, and one was exploring agreement using a visual screening tool.

We included three studies in the previous (2011) version of this review (Carter 1983, Weinberg 1977 and Weinberg 1979), but they have been excluded in this update: in the 2011 version, we included studies that investigated the effectiveness of visual scanning training and techniques even if the population of participants had not been clearly defined as having visual field defects. Carter 1983, Weinberg 1977 and Weinberg 1979 included populations of participants with 'visual scanning' problems, who may have had either visual field defects or visual neglect, or both. For this latest update, we reconsidered and reversed this decision: we have excluded populations of participants with 'visual scanning problems', but no confirmed visual field defect, to ensure that all included studies are focused on stroke survivors with confirmed visual field defects.

Studies awaiting classification

Two studies are awaiting classification ([Ghandehari 2011](#); [Sand 2017](#)). [Ghandehari 2011](#) compared two different pharmacological interventions (Neuroaid and Piracetam) without a control group. The selection criteria, which were pre-stated for this review, did not clearly state whether pharmacological interventions were relevant for inclusion, but discussion amongst review authors supports the conclusion that it would be appropriate to include any trials of pharmacological interventions if they are specifically focused on improving outcomes in participants with visual field defects. However, it remains unclear whether this study is a randomised controlled trial or not, as there is inconsistent reporting of the study design, and we await clarification from the study authors. Prior to any future updates of this review, we will clarify the selection criteria and methods to ensure that the inclusion (or exclusion) of pharmacological interventions is addressed. [Sand 2017](#) is an ongoing study; however, it was not possible to determine from available information whether this was a randomised controlled trial or not.

Ongoing studies

Seven studies are listed as ongoing studies. Two ongoing studies compare a form of visual restitutive training with control ([Feldon 2017](#); [NCT02886663](#)); one is comparing a form of transcranial electrical stimulation with control ([Gall 2015](#)); one is comparing a pharmacological intervention focused on restitution of the visual field with a placebo ([NCT02737930](#)); and one is comparing two different modes of delivery of computer-based compensatory scanning interventions with a control ([ISRCTN16023965](#)). Two of these studies, listed as ongoing in the previous version of this review, are now complete ([Hayes 2010](#); [Komm 2009](#)), but we have been unable to obtain results from the authors. If results are not available at the time of the next update, we will exclude these studies.

Risk of bias in included studies

We have described the assessment of risk of bias for individual studies in the 'Risk of bias' tables in [Characteristics of included studies](#), summarised in [Figure 2](#).

Figure 2. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

	Allocation concealment (selection bias)	Blinding (performance bias and detection bias)	Incomplete outcome data (attrition bias)	Other bias
Aimola 2011				
Bainbridge 1994				
Bowers 2014				
De Haan 2015				
Elshout 2016				
...				

Figure 2. (Continued)

Eishouti 2016				
Gall 2013				
Jarvis 2012				
Jobke 2009				
Kasten 1998				
Kasten 2007				
Keller 2010				
Modden 2012				
Plow 2010				
Poggel 2004				
Rossi 1990				
Roth 2009				
Rowe 2010				
Schuett 2012				
Spitzyna 2007				

Figure 2. (Continued)

Spitzyna 2007				
Szlyk 2005				

Allocation

Although all 20 included studies were described as randomised controlled trials, only 10 of them reported sufficient information on the method of randomisation to assess whether the randomisation and allocation concealment methods were adequate, and, in two of these studies, we judged them to be at high risk of bias: in [Keller 2010](#) some of the allocation appeared to have been alternate, rather than random, and, in [Modden 2012](#), randomisation was through throwing of a dice, with no allocation concealment.

Blinding

In the majority of included studies, the nature of the intervention meant that it was not possible to mask (blind) participants or people involved in their care. We judged blinding as low risk of bias if there was evidence that the outcome assessor was masked to the treatment allocation of the participants - six of the 20 studies clearly reported having a masked outcome assessor. In eight of the 20 studies this information was unclear, and in six of the 20 studies we judged that there was a high risk of bias as the outcome assessor was not masked.

Incomplete outcome data

Lack of information about the management of incomplete outcome data, and whether or not there had been dropouts or participants excluded from the studies, made it difficult to assess risk of bias relating to incomplete outcome data for all the studies. In four of the 20 studies, we considered that adequate reasons for dropouts were provided, numbers of dropouts were even between groups, or intention-to-treat analysis was carried out. However, in five of the 20 studies where we felt that incomplete outcome data were not adequately addressed, and that dropouts were either unbalanced or might be related to the intervention studied (e.g. because people dropped out or were excluded because of low compliance), we judged risk of bias to be high. For the remaining 11 studies, we were unclear as to whether or not incomplete outcome data were adequately addressed.

Other potential sources of bias

Lack of information and details of methodology within the included studies generally made it difficult for us to assess the potential risk due to other biases. However, four of the studies of visual restitution therapy were carried out by researchers who have acknowledged a financial interest in this intervention (as stated in [Poggel 2004](#)) ([Jobke 2009](#); [Kasten 1998](#); [Kasten 2007](#); [Poggel 2004](#)), and [Bowers 2014](#) declared a financial conflict of interest relating to the prisms that were being investigated. We assessed this to potentially introduce a source of bias. We identified concerns that the increased attention given to the treatment group, as opposed to the control group, in [De Haan 2015](#) may create a high risk of bias. For two of the studies, only abstracts were available and there was insufficient information on which to assess bias ([Bainbridge 1994](#); [Gall 2013](#)). We judged the remaining 12 studies as unlikely to be at risk of other potential sources of bias, but based this assessment on the absence of information suggesting bias rather than the presence of information indicating that the study was free from bias.

Studies included in meta-analyses within this review

From the 20 studies included in this review, there are only eight studies included in meta-analyses within this review ([Aimola 2011](#); [Bowers 2014](#); [De Haan 2015](#); [Jarvis 2012](#); [Kasten 1998](#); [Rossi 1990](#);

[Rowe 2010](#); [Spitzyna 2007](#)). These eight studies randomised 428 participants. The studies relevant to our pre-stated comparisons (see [Objectives](#)), and included within the planned meta-analyses are:

1. Restitutive interventions versus no treatment, placebo, or control

- [Kasten 1998](#) (visual restitution therapy versus placebo, n = 19). We have included data from this study in relevant analyses.

2. Compensatory interventions versus no treatment, placebo, or control

- [Aimola 2011](#) (n = 52), [De Haan 2015](#) (n = 49), [Rowe 2010](#) (n = 70), and [Spitzyna 2007](#) (n = 22) are all relevant and have data included in relevant analyses.

3. Substitutive interventions versus no treatment, placebo, or control

- [Rossi 1990](#) (prisms versus no treatment, n = 39) and [Rowe 2010](#) (prisms versus control, n = 59). We have included data from these studies in relevant analyses.
- [Bowers 2014](#) (prisms versus placebo, n = 67). This is a cross-over study and data were principally presented after the cross-over; however, there were data for one outcome (extended activities of daily living) presented before the cross-over, with data suitable for inclusion in analysis.

4. Assessment or screening versus no treatment, placebo, or control

- [Jarvis 2012](#) (assessment versus control, n = 39). We have included data from this study in relevant analyses.

Studies not included in meta-analyses

Ten of the studies included in this review did not have a control group and are therefore not included in meta-analyses.

These 10 studies each compared one active intervention with another active intervention:

- [Jobke 2009](#), [Kasten 2007](#), [Modden 2012](#), [Plow 2010](#), [Poggel 2004](#), and [Roth 2009](#) all investigated restitutive interventions, and did not include a no-treatment, placebo, or control group.
- [Keller 2010](#), [Modden 2012](#), [Roth 2009](#), and [Schuett 2012](#) all investigated compensatory interventions, and did not include a no-treatment, placebo, or control group.
- [Bainbridge 1994](#) and [Szlyk 2005](#) both investigated substitutive interventions, and did not include a no-treatment, placebo, or control group.

The results from relevant outcomes from these studies are summarised in [Table 5](#) and [Table 6](#) and in a narrative synthesis below.

One study did have a relevant control group (alternating current stimulation versus placebo, n = 39) but was published as an abstract only and we have been unable to identify data suitable for inclusion ([Gall 2013](#)).

One study did have a relevant control group (visual restitution training versus placebo, n = 30) but was a randomised cross-over trial and we have been unable to obtain data from the first phase only, in order to include data in meta-analyses ([Elshout 2016](#)).

Effects of interventions

See: [Summary of findings for the main comparison](#) Summary of findings: Restitutive interventions versus control; [Summary of findings 2](#) Summary of findings: Compensative interventions versus control; [Summary of findings 3](#) Summary of findings: Substitutive interventions versus control; [Summary of findings 4](#) Summary of findings: Assessment/screening interventions versus control

Restitutive interventions versus no treatment, placebo, or control

There was one included study: [Kasten 1998](#) (see [Summary of findings for the main comparison](#)). Data were available for visual field and quality of life outcomes, but not for any other outcomes of interest to this review. Stroke survivors with visual neglect were not included in [Kasten 1998](#), therefore these analyses relate to participants with visual field defects only (no co-existing visual neglect).

Visual field

See [Analysis 1.1](#). Data from [Kasten 1998](#) (19 participants) showed that there was no statistically significant effect of a restitutive intervention as compared to control (MD 1.02, 95% CI -1.37 to 3.41) for the visual field outcome (confrontation). We judged this evidence to be of very low quality.

Quality of life

See [Analysis 1.2](#). Data from [Kasten 1998](#) showed that there was a statistically significant effect of a restitutive intervention as compared to control (OR 13.00, 95% CI 2.07 to 81.48). The data used in this analysis were derived from 30 randomised participants, and included data from participants with optic nerve injury who had also received the same interventions in a separate (but parallel) trial. A total of 38 participants were randomised, of whom 19 had stroke and 19 had optic nerve injury; data were only available for 30 of 38 of these participants for this outcome. Separate data were not available for participants with stroke only. We judged this evidence to be of very low quality.

Compensatory interventions versus no treatment, placebo, or control

Included studies: [Aimola 2011](#); [De Haan 2015](#); [Rowe 2010](#); [Spitzyna 2007](#) (see [Summary of findings 2](#)). Stroke survivors with visual neglect were not included in [De Haan 2015](#); [Rowe 2010](#); [Spitzyna 2007](#), while participants in [Aimola 2011](#) could have co-existing neglect: data are presented in subgroups relating to the inclusion of participants with neglect.

Visual field

See [Analysis 2.1](#). Data from two studies (95 participants) showed that there was no statistically significant effect of compensatory interventions compared with control (SMD -0.11, 95% CI -0.92 to 0.70, heterogeneity: $I^2 = 75%$) ([De Haan 2015](#); [Rowe 2010](#)). We judged this evidence to be of very low quality. Several factors could contribute to the substantial heterogeneity, including different inclusion criteria (e.g. participants had to be < 26 weeks post stroke in [Rowe 2010](#)) and very different interventions (see [Characteristics of included studies](#)). However a key factor to note, which limits confidence in these findings, is that there was a significant difference in baseline assessment between groups in [De Haan 2015](#).

Extended activities of daily living

See [Analysis 2.2](#). Data from two studies (97 participants) showed that there was no statistically significant benefit in favour of compensatory interventions compared with control (SMD 0.49, 95% CI -0.01 to 0.99, heterogeneity: $I^2 = 25%$) ([De Haan 2015](#), [Rowe 2010](#)). We judged this evidence to be of very low quality.

Reading

See [Analysis 2.3](#). Data from four studies (162 participants) showed that there was no statistically significant effect of compensatory interventions compared with control (SMD 0.26, 95% CI -0.05 to 0.58, heterogeneity: $I^2 = 0%$) ([Aimola 2011](#); [De Haan 2015](#); [Rowe 2010](#); [Spitzyna 2007](#)). We judged this evidence to be of low quality. Although some of the participants in [Aimola 2011](#) may have had neglect, there was no downgrade for indirectness as the test for subgroup differences demonstrated no significant differences between the studies including or not including participants with neglect ($P = 0.43$).

Quality of life

See [Analysis 2.4](#). Data from two studies (96 participants) showed that there was a statistically significant effect of compensatory interventions compared with control (MD 9.36, 95% CI 3.10 to 15.62), with minimal heterogeneity ($I^2 = 0%$), based on the results for the total score for the VFQ-25 assessment ([De Haan 2015](#); [Rowe 2010](#)). We judged this evidence to be of low quality.

Scanning - cancellation

See [Analysis 2.5](#). Data from two studies (97 participants) showed that there was no statistically significant effect of compensatory interventions compared with control (SMD -0.01, 95% CI -0.40 to 0.39), with minimal heterogeneity ($I^2 = 0%$) ([Aimola 2011](#); [De Haan 2015](#)). We judged this evidence to be of low quality. Although some of the participants in [Aimola 2011](#) may have had neglect, there was no downgrade for indirectness as the test for subgroup differences demonstrated no significant differences between the studies including or not including participants with neglect ($P = 0.55$). Substituting the [De Haan 2015](#) 'parallel search test' data for 'all trials' with other presented data, including the 'serial search test' data and 'dot counting test', or substituting the 'target present' data for the 'all trials' data, did not change the non-significant result.

Adverse events

See [Analysis 2.6](#). [Rowe 2010](#) collected and reported data relating to adverse events, stating that: "Two patients (6.7%) in the visual search training arm experienced seven adverse events (six fatigue and one headache). No adverse events were recorded for the standard care arm". [De Haan 2015](#) did not report adverse events as an outcome measure, but stated that: "No important harms caused by the training or the assessments were encountered, nor reported by the participants". Data from [Rowe 2010](#) (see [Analysis 2.6](#)), showed that there was no difference in the odds of a participant having an adverse event with compensatory scanning training, when compared to control (OR 15.18, 95% CI 0.24 to 112.57). We judged this evidence to be of low quality.

Substitutive interventions versus no treatment, placebo, or control

Included studies: [Bowers 2014](#); [Rossi 1990](#); [Rowe 2010](#) (see [Summary of findings 3](#)). Stroke survivors with visual neglect were not

included in [Bowers 2014](#) and [Rowe 2010](#), while participants in [Rossi 1990](#) could have co-existing neglect. Substitutive prisms were worn during the outcome assessment by participants in [Bowers 2014](#) and [Rossi 1990](#), but were not worn during outcome assessment by participants in [Rowe 2010](#). As the wearing of the substitutive intervention during outcome assessment should theoretically expand the size of the visual field, the data arising from these different approaches (wearing or not wearing prisms) were presented as subgroups, and no pooled total was calculated.

Functional activities of daily living (primary outcome)

See [Analysis 3.1](#). Data from one study (39 participants, wearing prisms during assessment; participants may have co-existing neglect) showed that there was no statistically significant effect of a substitutive intervention compared with control for the primary outcome of functional activities of daily living (MD -4.00, 95% CI -17.86 to 9.86) ([Rossi 1990](#)). We judged this evidence to be of very low quality.

Visual field

See [Analysis 3.2](#). Data from one study (46 participants; no neglect) showed that there was no statistically significant effect of a substitutive intervention compared with control when the substitutive intervention (prism) was not being worn (SMD 0.12, 95% CI -0.46 to 0.70) ([Rowe 2010](#)). Data from one study (39 participants; possibly co-existing neglect) showed that there was a statistically significant effect of a substitutive intervention compared with control when the substitutive intervention (prism) was being worn (SMD 1.12, 95% CI 0.44 to 1.80) ([Rossi 1990](#)). There was a statistically significant difference between the subgroups in which participants did and did not wear prisms during assessment ($P = 0.03$). We judged this evidence to be of very low quality.

Extended activities of daily living.

See [Analysis 3.3](#). Data from one study (48 participants, no neglect) showed that there was no statistically significant effect of a substitutive intervention compared with control when the substitutive intervention (prism) was not being worn (SMD 0.20, 95% CI -0.44 to 0.85) ([Rowe 2010](#)). Data from one study (61 participants, no neglect) showed that there was no statistically significant effect of a substitutive intervention compared with control for measures of extended activities of daily living, using a mobility score (SMD 0.24, 95% CI -0.26 to 0.75) ([Bowers 2014](#)). There was no statistically significant difference between the subgroups that did and did not wear prisms during assessment ($P = 0.92$). We judged this evidence to be of very low quality.

Reading

See [Analysis 3.4](#). Data from one study (45 participants, no neglect) showed that there was no statistically significant effect of a substitutive intervention compared with control when the substitutive intervention (prism) was not being worn (MD 2.80, 95% CI -7.13 to 12.73) ([Rowe 2010](#)). We judged this evidence to be of low quality.

Falls

See [Analysis 3.5](#). Data from one study (39 participants, possible co-existing neglect) showed that there was no statistically significant effect of a substitutive intervention compared with control for the risk of falls (OR 1.21, 95% CI 0.26 to 5.76) ([Rossi 1990](#)). We judged this evidence to be of very low quality.

Quality of Life

See [Analysis 3.6](#). Data from one study (43 participants, assessed not wearing prisms; no neglect) showed that there was no statistically significant effect of a substitutive intervention compared with control for a measure of quality of life (MD 8.40, 95% CI -4.18 to 20.98) ([Rowe 2010](#)). We judged this evidence to be of low quality.

Scanning (cancellation)

See [Analysis 3.7](#). Data from one study (39 participants, assessed wearing prisms; possibly co-existing neglect) showed that there was a statistically significant effect of a substitutive intervention compared with control for measures of scanning (MD 9.80, 95% CI 1.91 to 17.69) ([Rossi 1990](#)). We judged this evidence to be of very low quality.

Adverse events

[Rowe 2010](#) collected and reported data relating to adverse events, stating that "Eighteen patients (69.2%) in the Fresnel prisms arm experienced a total of 42 adverse events of which 28 were classified as headache. No adverse events were recorded in the standard care arm". The reported adverse events in the group wearing prisms were: headache (28 events in six participants); diplopia (five events in five participants); visual confusion (four events in three participants); difficulty with navigation (two events in two participants); dizziness (two events in one participant); optical glare/aberrations (one event in one participant). [Analysis 3.8](#) shows that there was an increased odds of a participant having an adverse event if they were wearing prisms (OR 87.32, 95% CI 4.87 to 1564.66). We judged this evidence to be of low quality.

Assessment or screening interventions versus no treatment, placebo, or control

Included study - [Jarvis 2012](#) (see [Summary of findings 4](#)).

Functional activities of daily living (primary outcome)

See [Analysis 4.1](#). Data from [Jarvis 2012](#) (37 participants) showed that there was no statistically significant effect of an assessment intervention compared with control for measures of functional activities of daily living (MD -6.97, 95% CI -23.78 to 9.84). We judged this evidence to be of very low quality.

Subgroup and sensitivity analyses

As there were no more than six trials in any single meta-analysis, as pre-stated in our protocol, we did not carry out the planned subgroup or sensitivity analyses. There were two exceptions to this:

- we carried out subgroup analysis to explore the effect of including studies in which participants may have had neglect in addition to visual field defects. This decision was made as there was one study in which some of the participants may have had neglect ([Aimola 2011](#)), and we considered the implications for including this group of participants of key clinical importance. Results of these analyses are reported above;
- we carried out subgroup analyses to explore different ways of doing assessment in studies investigating the use of prisms (i.e. assessment wearing or not wearing prisms). This was considered central to the clinical relevance of the pooled result. Results of these analyses are reported above.

One active intervention versus another active intervention (narrative synthesis)

As planned, no meta-analyses have been carried out to pool data from studies that compared one active intervention with another active intervention. A summary of results for relevant outcomes for these comparisons is provided in Table 5 and Table 6, and a brief narrative summary below.

Studies comparing two similar active interventions

(See Table 5)

One restitutive intervention versus another restitutive intervention

We found two studies that compared the effectiveness of two (or more) types of visual restitution therapy (Jobke 2009; Kasten 2007). Jobke 2009 compared extrastriate visual restitution therapy with conventional visual restitution therapy in a cross-over study including 18 participants. After the first phase of the study, the extrastriate visual restitution therapy group had improved in measures of visual field (detection performance in HRP) by 5.9% and the conventional visual restitution therapy group had improved by 2.9%. No significant changes were found for either group for Radner reading scores. Kasten 2007 compared three treatment groups: 1) standard visual restitution therapy with single stimulation, 2) visual restitution therapy with parallel co-stimulation, and 3) visual restitution therapy with moving co-stimulation. We found no statistically significant differences between groups for measures of the visual field (perimetry).

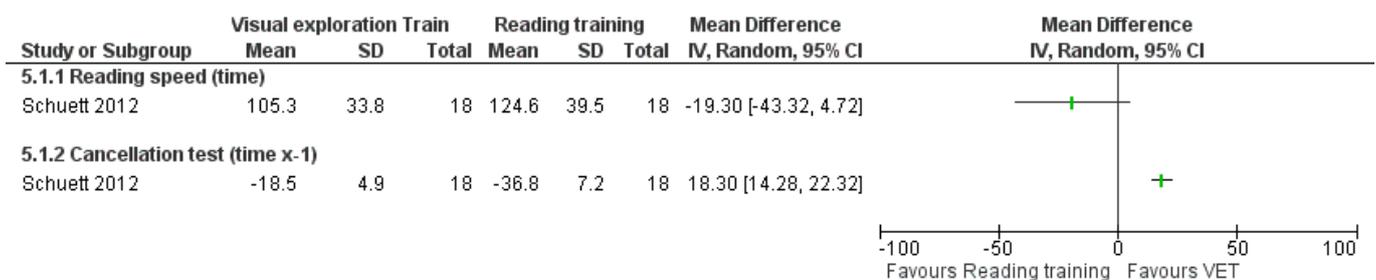
We found two studies that investigated the addition of another intervention to visual restitution therapy (Plow 2010; Poggel 2004). Plow 2010 investigated the effect of adding transcranial direct cur-

rent stimulation (tDCS) to visual restitution therapy and reported a greater increase in detection accuracy (perimetry) in the group receiving tDCS (9.37% increase compared to 5.55% increase in control group). Poggel 2004 compared standard visual restitution therapy with visual restitution therapy plus attentional cueing in 19 participants; results were primarily presented graphically and as whole-group data: the authors concluded that the visual field border increased significantly more in the participants in the attentional cueing group.

One compensative intervention versus another compensative intervention

We found three studies that compared different compensative interventions (Keller 2010; Modden 2012; Schuett 2012). Keller 2010 compared audiovisual training and visual exploration training in 10 participants, reporting that audiovisual training was better than visual exploration training for outcomes of activities of daily living, reading, and visual scanning. Schuett 2012 compared reading training and visual exploration training in 36 participants within a cross-over study. We used data reported from before the cross-over period to calculate effect sizes for Schuett 2012 (see Figure 3); this demonstrated that visual exploration training was significantly more beneficial than reading training at improving scanning outcomes, and there was a trend towards reading training improving reading outcomes more significantly than visual exploration training. Modden 2012 compared 45 participants within three groups; one receiving computer-based restitution therapy, one computer-based compensation therapy, and one standard occupational therapy. We judged the occupational therapy intervention to be a compensatory intervention. We found no differences between the groups receiving the computer-based compensatory therapy and occupational therapy interventions for measures of visual field enlargement, reading, or visual scanning.

Figure 3. One compensatory intervention versus another compensatory intervention. Effect sizes for Schuett 2012 (see Table 5)



One substitutive intervention versus another substitutive intervention

We found two studies that compared the effectiveness of two types of prism (Bainbridge 1994; Szlyk 2005). Bainbridge 1994 compared the effect of full-field and hemi-field prisms in 18 participants, and reported a significant effect in favour of full-field prisms for the cancellation test and Harrington Flocks Visual Field score. Szlyk 2005 compared Fresnel prisms and the Gottlieb Visual Field Awareness System in 10 participants and found no statically significant differences in outcomes between the two groups, although this was a cross-over study with no data presented for the period of time before the cross-over.

Studies comparing two different types of active intervention

(see Table 6)

Compensation intervention versus restitutive intervention

Two studies compared the effectiveness of a compensation intervention and a restitutive intervention (Modden 2012; Roth 2009). Roth 2009 compared explorative scanning training (a compensation intervention) with flicker stimulation training (a restitution intervention) in 29 participants, finding no significant differences between groups for key outcomes (although there were differences between groups at baseline for some outcomes). Modden 2012 compared 45 participants within three groups; one receiving computer-based restitution therapy, one computer-based compensa-

tion therapy, and one standard occupational therapy. In [Table 6](#), we compared the results of the groups receiving computer-based training. There were no significant differences reported for these groups for measures of visual field enlargement, activities of daily living, reading, or visual scanning.

Compensation intervention versus substitution intervention

One study compared the effectiveness of a compensative intervention (paper-based visual search training) and a substitutive intervention (Fresnel prisms) ([Rowe 2010](#)). This was a three-armed study, with a control (standard care) treatment; data from the active treatment arms compared to the control treatment have been included in meta-analyses within this review. We found no significant differences between the compensative and substitutive intervention for measures of visual field, extended activities of daily living, and reading. We found a statistically significant difference in favour of the compensative intervention for the measure of quality of life (using the VFQ-25).

DISCUSSION

Summary of main results

We found 20 studies (732 randomised participants, with data for 638 participants, 547 of whom were participants with stroke) that investigated interventions for visual field defects in people with stroke. However, only 10 of these studies compared the effect of an intervention with a placebo, control, or no treatment group, which were the comparisons of interest to the review question, and only eight had data suitable for inclusion within meta-analyses. Only two of these eight studies presented data relating to our primary outcome of functional abilities in activities of daily living, and there was a lack of consistency in outcome measures used across studies, which limited our ability to draw generalised conclusions.

Effect of restitutive interventions

Three studies (88 participants) compared a restitutive intervention with a control, but data were only available for one (19 participants). There was very low-quality evidence that visual restitution therapy had no effect on visual field outcomes, and a statistically significant positive effect on quality of life. However, the data relating to the quality of life outcome must be interpreted with caution as the data used for analysis combined the population of interest with an additional 19 participants with optic nerve injury who had been included in a separate (but parallel) trial. These participants had damage to the anterior visual pathway, a population which was specifically excluded from this review, and in none of the participants was the optic nerve damage due to stroke. We, therefore, do not believe that the findings based on data from this population are applicable to the population of patients with visual field loss due to post-chiasmal stroke. There is, therefore, insufficient evidence to draw any conclusions about the effectiveness of visual restitution therapy as compared to placebo, control, or no treatment. There was also some very limited evidence from two small studies which compared different types of restitutive interventions that there may be some benefits to adding either attentional cueing or tDCS to visual restitutive therapy, while a further two small studies found no difference in different modes of delivering visual restitution therapy.

Effect of compensatory interventions

Four studies (193 participants) compared a compensatory intervention with a control. There was low-quality evidence of a beneficial effect on measures of quality of life. However, there was low- or very low-quality evidence of no effect on measures of visual field, extended activities of daily living, reading, and scanning ability. Findings from a small study comparing two different types of compensatory therapy conflicts with the evidence of no effect on scanning outcome, demonstrating a beneficial effect of visual exploration therapy on scanning outcomes when compared to reading training. There is, therefore, some limited low-quality evidence that compensatory scanning training may improve an important outcome (quality of life) in people with visual field defects following stroke, but further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate. Studies comparing different active interventions provide some very limited evidence that there may be benefits associated with audiovisual training, rather than visual exploration training alone, and suggest that compensatory training, as delivered during standard clinical practice, may be as effective as computer-based training interventions.

Effect of substitutive interventions

Three studies (166 participants) compared a substitutive intervention (a type of prism) with a control. There was low- or very low-quality evidence that prisms did not have an effect on measures of activities of daily living, extended activities of daily living reading, falls, or quality of life, but that they may have an effect on scanning ability. Evidence from one study indicates that people using prisms may have a higher odds of adverse events, particularly headache. However, it is important to note a fundamental difference between these studies in that, in two of the studies, participants in the treatment group wore prisms during the outcome assessments ([Bowers 2014](#); [Rossi 1990](#)), while in one study none of the participants wore prisms during outcome assessment by participants ([Rowe 2010](#)). Evidence relating to measures of visual field demonstrated a significant difference between studies that measured outcome whilst wearing or not wearing the substitutive device, with a beneficial effect when the prisms were worn during testing but not without. However, due to the quality and quantity of evidence, we remain uncertain about the benefits of prisms.

Effect of screening/assessment interventions

One study (39 participants) compared the effect of assessment by an orthoptist to standard care (no assessment) and found very low-quality evidence that there was no effect on measures of activities of daily living. However, due to the quality and quantity of evidence, we remain uncertain about the benefits of assessment interventions.

Ten studies compared the effect of two or more different active interventions and did not include a control group.

In summary, this review has identified:

- a lack of evidence relating to the effect of interventions on our primary outcome (functional ability in activities of daily living);
- low-quality evidence that compensatory scanning training is more beneficial than placebo or control at improving quality of life, but not other outcomes;

- insufficient evidence to reach any generalised conclusions about the effect of restitutive interventions, substitutive interventions (prisms), or screening/assessment interventions as compared to placebo, control or no treatment;
- low-quality evidence that prisms may result in an increase in the number of people experiencing adverse events (particularly headache).

Overall completeness and applicability of evidence

Studies with control, placebo, or no treatment group

Although we identified 20 studies that investigated the effect of interventions for visual field defects after stroke, only 10 of these studies (and only eight with data) compared the effect of an intervention with a control, placebo, or no treatment group, which was the question of interest to this review. The remaining 10 studies compared different types of interventions, with nine of these studies comparing interventions of the same 'type' (i.e. comparing two substitutive interventions or two restitutive interventions); arguably, comparisons of different active interventions have little merit until such time as the benefits (and harms) of active interventions, as compared to control, placebo, or no treatment have been established. Three studies did compare the relative effects of two different types of intervention (i.e. compensatory versus restitutive, or compensative versus substitutive). Thus, although we made the decision to include all the studies which investigated the effectiveness of interventions for visual field defects, in fact only nine of these studies included comparisons that were directly relevant to the review question, focusing on a comparison with a control, placebo, or no treatment group. Five of the 20 included studies, including two of the 10 studies with a control/placebo comparison, were cross-over studies. The reporting of data from after the cross-over in studies with this design limited our ability to incorporate data within meta-analyses in this review, and there is a risk that outcomes from after the cross-over are affected by the treatments administered prior to the cross-over.

Restitutive interventions versus no treatment, placebo, or control

[Kasten 1998](#) was the one study comparing a restitutive intervention with a control or placebo, with data suitable for inclusion in our meta-analyses. This study included 19 participants, only 10 of whom had stroke. The study did not measure our primary outcome of interest (functional ability in activities of daily living). There was an uneven distribution of stroke patients between the two groups (with two stroke patients out of the nine participants in the treatment group and eight stroke patients out of 10 participants in the control group). This uneven distribution means that only two patients with stroke received an active intervention, providing evidence from which it would be inappropriate to generalise. Furthermore, for quality of life outcome data, the 19 participants in this study had been combined with the results of 19 participants in another similar study. However, the additional 19 participants all had optic nerve injury rather than post-chiasmal injury. Due to the nature of the participants included in this study, it would not be appropriate to make generalisations from this evidence to the population of stroke patients with visual field defects after stroke. Our confidence in the findings from this evidence is very low.

Compensatory interventions versus no treatment, placebo, or control

We were able to combine data from four studies (162 participants) comparing compensatory (scanning) interventions with control or placebo in meta-analysis. None of these studies measured our primary outcome of interest (functional ability in activities of daily living). Visual field defects in the participants included in these studies were confirmed using perimetry (see [Table 3](#)). Three of the studies included participants with visual field defects and no co-existing visual neglect; two of these studies excluded participants with neglect based on clinical testing ([De Haan 2015](#); [Rowe 2010](#)), while one only included participants with a left-sided cerebrovascular accident where patients rarely experience persistent neglect ([Spitzyna 2007](#)). One study included participants who had visual field defects but possibly also co-existing visual neglect ([Aimola 2011](#)); clinical testing confirmed that only three of the 52 participants had confirmed neglect. Participants included those from a mixed population (i.e. stroke and other neurological conditions) for three of the four studies; 41 of the 49 participants had stroke in [De Haan 2015](#), 39 of 52 had stroke in [Aimola 2011](#), and 13 of 22 had stroke in [Spitzyna 2007](#). All participants had stroke in [Rowe 2010](#). The majority of participants in these studies, therefore, had visual field defects following stroke and did not have co-existing neglect (as confirmed by clinical testing); therefore, it would be appropriate to generalise from these results to the population of stroke survivors with visual field defects and no neglect.

The nature of the scanning training in the four studies combined within the meta-analyses varied considerably; two investigated computer-based scanning training, one focused on visual exploration training ([Aimola 2011](#)), one on reading training using scrolling horizontal text ([Spitzyna 2007](#)), one was a training programme, primarily delivered face-to-face by an occupational therapist ([De Haan 2015](#)), and one was a self-delivered paper-based scanning exercise ([Rowe 2010](#)). It is likely that the nature of these interventions will result in varied scanning movements of the eye. As well as differences in the mode of delivery of the scanning training, there were also differences in the amount of training. These differences in the compensatory interventions reduce confidence in any findings, and limit ability to generalise from these findings. During the GRADE assessment of quality of evidence, we applied a downgrade to each comparison combining results from these studies due to the variations in the interventions studied. We, therefore, have low- to very-low confidence in the findings from this evidence.

Substitutive interventions versus no treatment, placebo, or control

Two studies compared a substitutive intervention (prisms) with a no treatment control ([Rossi 1990](#); [Rowe 2010](#)). [Rossi 1990](#) measured our primary outcome of interest (functional ability in activities of daily living). Both studies only included participants with stroke, but [Rossi 1990](#) included participants with visual neglect in addition to visual field defect. The studies both investigated the effect of Fresnel prisms. There was a fundamental difference between these studies relating to the assessment of outcome: [Rossi 1990](#) measured outcomes whilst participants in the intervention group wore the assigned prisms, whilst [Rowe 2010](#) measured outcomes while participants were not wearing any assigned prisms. It has been argued that the rationale for prisms is that they provide visual field expansion when in use, and that consequently outcomes from clinical trials exploring the effectiveness of prisms should be

measured with participants wearing the prisms (Bowers 2014). In contrast, Rowe 2010 measured outcomes without use of assigned prisms in order to preserve blinding of outcome assessor and enable direct comparison of study treatment groups. It was, therefore, not appropriate to pool data from these different studies of prisms, as the wearing of prisms during assessment in one study but not the other makes the results incomparable. This is an important issue which must be appropriately considered in future trials. Due to the methodological limitations of these studies, and the inability to combine results, we have low to very low confidence relating to this evidence.

Assessment/screening intervention versus control

One study explored the effect of implementing a full visual assessment by an orthoptist and sharing the results with hospital staff (Jarvis 2012). This study measured our primary outcome of interest (functional ability in activities of daily living). The evidence was judged to be very low quality, limiting our confidence in the findings from this study.

Quality of the evidence

For this updated review, we judged the quality of evidence using the GRADE approach. We judged all evidence included within meta-analyses to be of low or very low quality. Key factors contributing to downgrading of the evidence within these comparisons included:

Risk of bias

We identified concerns about the methodology for the majority of included studies, and there were often insufficient details available from incomplete reporting of methodological details. We judged only eight of the 20 studies to be at low risk of bias for allocation concealment, six of the 20 for blinding of outcome assessment, and four of the 20 for incomplete outcome assessment. Furthermore, five of the studies were carried out by researchers who have acknowledged a financial interest in intervention (Bowers 2014; Jobke 2009; Kasten 1998; Kasten 2007; Poggel 2004); we assessed this as potentially introducing a source of bias.

Imprecision

The number of participants within the included studies was small, ranging from 10 to 87 participants, with only five of the 20 studies including more than 50 participants. While there were a total of 732 randomised participants, variations in studies made it inappropriate for the majority of study data to be combined within analyses, and the maximum number of participants with data combined in a single analysis was 162 (Analysis 2.3). The small number of participants within the included studies and suitable for combination within meta-analyses limits the conclusions that can be drawn from this evidence.

Indirectness

A number of factors contributed to indirectness of the data included within meta-analyses. In particular:

- population: there was considerable heterogeneity between the populations recruited to individual studies. In addition to stroke-related differences, such as time post-stroke, initial impairment, and the presence of other stroke-related impairments (e.g. communication, mobility), this was confounded by the inclusion of participants with conditions other than stroke, and in

opposing the decision to either include or exclude participants with visual neglect. The variations in populations contributed to decisions to downgrade the quality of evidence, as this reduced our certainty in the reported findings;

- interventions: this review aimed to synthesise evidence relating to a wide range of different interventions for visual field defects following stroke and preplanned categories to support appropriate combination of evidence. However, we found substantial variations in the interventions within these different categories in relation to the details of the delivered interventions. In particular, the compensative interventions had considerable variation in the mode of delivery, with interventions varying from computer-based scanning training and reading training, paper-based scanning training, to face-to-face scanning and mobility training. What is being delivered in terms of the eye movements being trained with these different interventions is likely to vary considerably, and - while data from these 'scanning' interventions have been combined - the variations in interventions limit our ability to be confident about the pooled result;
- outcomes: as is highlighted in Table 4, there was a lack of consistency in the outcomes assessed by individual studies, and in the assessment tools used to do this. While there is an argument that outcome measures should be carefully selected according to the anticipated action of, or scientific rationale for, the intervention (Bowers 2017), the variations in interventions do not fully explain the lack of consistency between the outcome measures. Within the studies of prisms, the difference in choice of outcome measure was further confounded by opposing decisions relating to whether outcomes were assessed wearing or not wearing the prisms. The variations in outcome measures limited the ability to pool data from individual studies in a meaningful way, and where measures have been pooled, limited our certainty in the result.

In summary, we judged the quality of the evidence synthesised within this review to be low to very low, and this limits our confidence in the results. Future research needs to address the factors which contribute to this level of evidence, in order to produce results which are useful and meaningful.

Potential biases in the review process

Publication bias

Through a thorough searching process we are quite confident that we should have identified all relevant published studies. However, at the peer review stage of this review update, we were alerted to the fact that we had erroneously excluded a relevant study (Elshout 2016): while this was corrected prior to publication, it does highlight the potential for human error in our process of screening titles. A limitation of our search strategy for this update is that we have not searched a number of trials databases beyond March 2015 (Current Controlled Trials, Health Service Research Projects in Progress, National Eye Institute Clinical Studies Database, Stroke Trials Registry); this may have limited our ability to identify ongoing trials, but ought not to have impacted on our identification of completed trials. It must be acknowledged that there is a small possibility that there are additional studies (published and unpublished) that we did not identify. We had planned to explore the effect of publication type using sensitivity analyses; however, all data included in meta-analyses were from peer-reviewed journals.

Categorisation of interventions

Although we anticipated that we may experience difficulty in categorising the interventions studied into our predefined categories of restitutive, compensative, and substitutive interventions, this was not the case and the categorisation process was a clear and unambiguous process. This was because the studies we identified were primarily visual restitution therapy (restitutive), compensatory scanning training (compensative), or prisms (substitutive). We are, therefore, confident that our categorisation of interventions has not introduced bias into the review process. However, we did find substantial differences between the interventions within each category (see discussion above), and decisions to combine data from varied interventions may reduce applicability of these results. Future updates of this review should, therefore, consider and pre-plan which interventions it is clinically relevant to combine. Involvement of key stakeholders to inform this decision making for future updates would be an advantage.

Inclusion criteria: participants

In the previous version of this review, we reported that the inclusion criterion that was judged as most difficult to assess by the independent review authors was the participants. The particular difficulty encountered was with studies that did not appear to include the diagnosis of visual field defects as an inclusion criterion. In the previous version, we identified and included several studies that either used a clinical assessment of 'scanning' as an inclusion criterion without formally assessing or diagnosing either visual field defects or visual neglect, or which included participants with a right-sided cerebrovascular accident, making the assumption that these participants would have visual neglect (and possibly also visual field defects). For this update of the review, we addressed this difficulty by only including studies that reported a method for diagnosing visual field defects at the recruitment stage. This led to the exclusion of a number of studies that had previously been included, but in which the participants did not have confirmed visual field defects (Carter 1983; Weinberg 1977; Weinberg 1979). We made this change between the previous version and this updated version of the review in order to reduce potential bias in the review process; however, we acknowledge that this may have led to the exclusion of studies in which some participants had visual field defects that were not confirmed through clinical diagnosis (e.g. instead manifesting as a scanning problem).

Outcomes

Categorisation of some reported outcome assessments into our predefined outcomes of interest was difficult in some cases. For this update of the review, we added a table to report our categorisation of outcomes to ensure transparency in this process (Table 4). We also reconsidered categorisation of all outcomes from trials included in the previous version of the review, and made a number of changes through a process of consensus. For example, Plow 2010 reported the Veterans Affairs Low Vision-Visual Functional Questionnaire (LV-VFQ) which "assesses an individual's visual ability to perform ADLs across 4 domains, including reading, mobility, visual motor function, and visual processing". This measure arguably relates to both extended activities of daily living and quality of life. In the first version of this review, this was listed as a measure of EADL: however, this was changed to being listed as a measure of QoL outcome for subsequent updates, following consensus discussion between review authors.

The primary outcome for this review was functional ability in activities of daily living, measured using standardised scales. It has been argued that measurement of effectiveness in rehabilitation ought to take into account patients' individual goals (Turner-Stokes 2009). There is growing evidence that goal attainment scaling (a standardised method of scoring performance of patient-specific tasks) may provide a valid, reliable, sensitive method of evaluating outcomes that are of greatest importance to individual patients (Krasny-Pacini 2016). This scale was not considered for inclusion within this review, and could be considered for future updates.

As has been discussed under [Quality of the evidence](#), variations in outcomes and outcomes assessment tools between included studies created challenges for the synthesis of evidence within this review. The need to make judgements and decisions relating to categorisation and pooling of outcomes created potential for the introduction of bias into the review process. We have aimed for transparency in our decision making and reporting in an attempt to avoid the introduction of bias. However, to support the creation of meaningful evidence syntheses and meta-analyses, there is a need for consensus between stroke survivors, their families and carers, health professionals, and researchers in relation to core outcomes for trials relating to interventions for visual field defects after stroke as recommended by the [COMET Initiative](#).

Agreements and disagreements with other studies or reviews

Agreements and disagreements between this updated version and previous version

In the previous 2011 version of this review, we stated the key conclusions and implications for practice arising from the evidence as follows.

- There is limited evidence which supports the use of compensatory scanning training for patients with visual field defects (and possibly co-existing visual neglect) to improve visual field, scanning, and reading outcomes.
- There is insufficient evidence to reach a conclusion about the impact of compensatory scanning training on functional activities of daily living.
- There is also insufficient evidence to reach generalised conclusions about the benefits of visual restitution therapy (restitutive intervention) or prisms (substitutive intervention) for patients with visual field defects after stroke.

Key changes in the methods between different versions of this review include:

- updated searches in this updated version, increasing the number of included studies;
- amended inclusion criteria, leading to the exclusion of studies in which participants did not have confirmed visual field defects;
- the use of the GRADE approach to systematically assess quality of evidence in this updated version.

These changes have highlighted further uncertainty around previous limited evidence supporting the use of compensatory interventions and have introduced evidence relating to adverse events associated with substitutive interventions, but have not resulted in any changes in conclusions relating to restitutive or substitutive interventions.

- There is limited low-quality evidence that compensatory scanning training may improve an important outcome (quality of life) in patients with visual field defects following stroke, but further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate. There is insufficient evidence to reach a conclusion about the impact of compensatory scanning training on other outcomes.
- There remains insufficient evidence to reach generalised conclusions about the benefits of visual restitution therapy (restitutive intervention) or prisms (substitutive intervention), and there is insufficient evidence to reach conclusions about the effect of screening or assessment interventions for patients with visual field defects after stroke.
- There is now some low-quality evidence from one study that prisms may result in an increased number of adverse events, particularly headache.

Agreements and disagreements with other published reviews

The Royal College of Physicians updated the evidence included in the earlier version of this review, concluding that there is "insufficient evidence regarding the effectiveness of interventions aimed at improving function in people with visual field defects" (ISWP 2016). This updated review is in agreement with the conclusions drawn from the evidence in the guideline.

The Scottish Intercollegiate Guideline Network (SIGN) guidelines for stroke rehabilitation state that there is "limited poor quality evidence suggesting that visual scanning compensatory training techniques may be effective in improving functional outcomes after stroke" (SIGN 2011). This SIGN guideline (updated in 2010) is based on a number of other reviews (Barrett 2009; Bouwmeester 2007; Jones 2006). While the previous version of our review was in agreement with the recommendations made by the Royal College of Physicians and SIGN guidelines, our updated review has highlighted further uncertainty relating to the effect of compensatory interventions. The National Institute for Clinical Excellence (NICE) stroke guidelines recommends that "eye movement therapy" is provided to "people who have persisting hemianopia after stroke and who are aware of the condition" (NICE 2013): our updated review does not directly support this recommendation.

Our review is in agreement with the conclusions in other reviews that the evidence relating to the effectiveness of visual restoration therapy is inconsistent and of poor quality (Barrett 2009; Bouwmeester 2007), and that few studies have assessed functional ability in activities of daily living as an outcome.

Our review is in agreement with narrative reviews of evidence for interventions for visual problems after stroke (Lane 2008, Hanna 2017), which have concluded that there is a need for high-quality studies of the effectiveness of interventions for visual field defects.

AUTHORS' CONCLUSIONS

Implications for practice

There is limited low-quality evidence that compensatory scanning training may improve an important outcome (quality of life) in patients with visual field defects following stroke, but further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate. There is in-

sufficient evidence to reach a conclusion about the impact of compensatory scanning training on other outcomes.

There is insufficient evidence to reach generalised conclusions about the benefits of visual restitution therapy (restitutive intervention), prisms (substitutive intervention), or assessment or screening interventions for patients with visual field defects after stroke. Prisms may cause a range of adverse events, particularly headache.

Implications for research

Specific implications for research, based on the findings of this review, are outlined below.

Are further randomised controlled trials (RCTs) required?

RCTs are required to determine the effect of:

- compensatory scanning training compared to no treatment, placebo, or usual care;
- restitutive interventions compared to no treatment, control, or placebo;
- substitutive interventions compared to no treatment, control, or placebo;
- assessment or screening interventions compared to standard care.

Such RCTs must:

- have adequate power (i.e. with an appropriate power calculation undertaken based on existing trial evidence);
- have adequate allocation concealment, blinding of outcome assessor, and intention-to-treat analysis;
- clearly define trial participants, with particular care relating to the diagnosis and inclusion of patients with visual field defects or visual neglect, or both;
- consider the severity of the visual field defect and plan subgroup analyses, where appropriate;
- include measures of functional ability in activities of daily living;
- collect and report data relating to adverse events;
- report clear and usable data.

We recommend that future RCTs concentrate on answering the specific question relating to the effectiveness of interventions compared to control, placebo, no treatment, or usual care rather than comparisons with variations of the same 'type' or category of intervention, or comparisons of different doses, adjuncts to treatment, or modes of delivery. We believe that until such time as the benefits of interventions for visual field defects compared to control, placebo, no treatment, or usual care have been established (or refuted), it is not beneficial to compare the relative effects of different interventions. We recommend that future RCTs should include outcomes that are of importance to people affected by this problem, such as quality of life, confidence, and participation in social activities and relationships. We recommend that consensus is reached relating to the important outcome measures which all future trials of interventions for visual field defects in people with stroke should assess (i.e. a core outcome set), and that stroke survivors, carers, health professionals, and researchers are involved in developing this core outcome set.

A number of RCTs are currently ongoing. Once they are completed it will be important to update this review and to re-evaluate the need for further RCTs of interventions for visual field defects.

Are other primary research studies required?

We do not recommend other study designs aimed at investigating the effects of visual restitution therapy or prisms, although development or pilot studies may be justified to support the design of a full trial if such work has not previously been completed. Given the variation in compensatory interventions included in this review, we do recommend that other primary research studies are considered to explore an optimal scanning training intervention. Such studies may consider the evaluation of computerised as well as non-computerised interventions, the effective dosage, and the role of reading-specific training. However, this work ought to be done as a pre-quel to a well-designed RCT of the scanning intervention. Studies evaluating different types of scanning training should provide clear details of the treatment parameters, including the eye movements which are being trained by the intervention.

There is a need for further research to identify optimal outcome measures for use within future RCTs in this area. Such research may consider the relationship between functional measures of activities of daily living (ADL), measures of vision-related extended ADL (such as reading, driving, navigating within an environment), and laboratory measures of the visual field.

Are further systematic reviews required?

We do not recommend any further systematic reviews aimed at addressing the effectiveness of interventions for visual field defects after stroke. We do recommend that this review is updated when the ongoing RCTs are completed.

Summary of findings

- The quality of evidence summarised in this review is judged to be low to very low.
- Methodological quality of studies is, in general, poor or poorly reported, providing insufficient high-quality evidence on which to reach generalisable conclusions.

- Limited low-quality evidence suggests compensatory interventions may improve an important outcome (quality of life) in patients with visual field defects following stroke, but further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate. There is insufficient evidence to reach a conclusion about the impact of compensatory scanning training on functional activities of daily living, or other outcomes.
- There is insufficient evidence to reach generalised conclusions about the benefits of vision restoration therapy for patients with visual field defects after stroke.
- There is insufficient evidence to reach generalised conclusions about the benefits of prisms for patients with visual field defects after stroke; there is some low-quality evidence that prisms may cause adverse events.
- High-quality RCTs are needed to compare compensatory, restitutive, substitutive, and assessment interventions with placebo, control, no treatment, or usual care.

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* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES
Characteristics of included studies [ordered by study ID]

Aimola 2011

Methods	<p>Compensative intervention versus control, placebo, or no intervention</p> <p>Design: "A randomized, controlled, parallel-group design was used."</p> <p>Stratification: no</p> <p>Randomisation sequence: "participants were randomized equally to 2 groups ... using parallel trial allocation software"</p> <p>Comparisons: 2 groups: intervention versus control</p> <p>Allocation concealment: "Participants were informed about the training types but did not know to which group they were assigned."</p> <p>Blinding: not stated if outcome assessor was blinded</p> <p>Power calculation: no (feasibility study)</p> <p>Intention-to-treat analysis: no</p> <p>Other recruitment details: "recruited from local hospitals or as self-referrals"</p> <p>Patient and public involvement: not stated</p>
Participants	<p>Total study population: randomised 70; "final sample included in analyses consisted of 52 participants; 28 intervention and 24 control"</p> <p>Withdrawals: 18 "dropped out during the intervention period because of health problems (n = 7), death (n = 2), or low motivation (n = 9)".</p> <p>Method of diagnosing VFD: monocular automatic perimetry</p> <p>Characteristics of population: "chronic hemi-VFDs resulted from any post-chiasmatic lesion." Participant details are listed in Table 2.</p> <p>Type and severity of visual problems: participant details are listed in Table 3.</p> <p>Inclusion criteria: at least 18 years old, no previous access to any formal hemi-VFD rehabilitation (restoration, substitution or compensation)</p> <p>Exclusion criteria: "medical instability, inability to provide informed consent, visual loss as a consequence of prechiasmatic damage or a progressive condition, photosensitive epilepsy, oculomotor disorders, and severe cognitive impairment. Participants were not enrolled until at least 3 months after onset to minimize confounding by spontaneous recovery (range = 3-276 months)".</p>

Aimola 2011 (Continued)

Baseline comparison of treatment groups: no differences noted at baseline

This study included participants with diagnoses other than stroke.

This study included participants with visual field defects and co-existing visual neglect.

Method of diagnosing visual perceptual problems: Bells test

Interventions

Group 1: home-based compensatory training (n = 28)

Intervention: "the experimental training consisted of reading and exploration components; patients completed components sequentially with order randomized."

Intervention type: compensation. **Materials:** computer-based. **Where can materials be accessed?** not stated. **Procedures:** "In the visual exploration tasks patients had to find a target defined by a specific feature (color, shape, size) among an array containing distractors (i.e. a red letter among blue ones). In the reading task, patients had to detect a nonword target (ie, vowels) among a varying number of words (i.e. accent), presented in a single central horizontal line. In both task types participants responded to target presence using an appropriate computer-mouse press. Computer feedback on speed and accuracy of responses, and overall progress to date (i.e. difficulty level achieved and number of training sessions completed), was provided at the end of each block of trials." **Provided by:** "at the start of the intervention period the experimenter demonstrated the training, during which the computer was set up ... Participants were encouraged to train in a similar manner, though adherence cannot be guaranteed". **Delivery:** home-based (Table 1). **Regimen:** "Patients could perform a maximum of 14 blocks per day. Each block contained 120 trials. They completed 294 exploration and 196 reading blocks." **Tailoring:** "For both training components difficulty level was dynamically adjusted based on both accuracy and speed of previous performance. If patients were $\geq 90\%$ accurate then difficulty would increase to the next level, whereas with accuracy $< 75\%$ difficulty would drop to an easier level. In exploration tasks difficulty was increased by enlarging the spatial zone within which a target could appear and by making targets and distractors more similar. For the reading task the word length and number of distractor words increased (up to a maximum of 7). For both tasks, presentation time was directly related to previous response times, that is, the faster the participant, the shorter the successive presentation time". **Modification:** see tailoring. **Adherence:** "There were 18 withdrawals, 9 due to low motivation. A total of 89% of the low-motivation withdrawals were in the control group. Recommended training duration was 5 weeks. However, only 3 patients met this requirement; the mean completion time for the experimental training was 9.3 weeks (SD = 6.0)."

Group 2: control (n = 24)

Intervention: "This training consisted of a number of tasks requiring visual attention but no systematic exploration or large horizontal eye movements." **Intervention type:** placebo/attention control. **Materials:** computer-based. **Where can materials be accessed?** not stated. **Procedures:** "The randomly presented tasks included a Go/No-Go task, centrally presented sequential search, Sternberg task, and a "rabbit hunting" task." **Provided by:** "at the start of the intervention period the experimenter demonstrated the training, during which the computer was set up ... Participants were encouraged to train in a similar manner, though adherence cannot be guaranteed". **Delivery:** home-based (Table 1). **Regimen:** "Patients were instructed to complete 10 blocks per day in approximately 1 hour, with a total of 350 blocks." **Tailoring:** "Difficulty was adjusted dynamically depending on performance by reducing presentation time or increasing attentional load." **Modification:** see tailoring. **Adherence:** "There were 18 withdrawals, 9 due to low motivation. A total of 89% of the low-motivation withdrawals were in the control group. Recommended training duration was 5 weeks. However, only 3 patients met this requirement; the mean completion time for the experimental training was 9.3 weeks (SD = 6.0)."

Outcomes

See Table 4

Perimetry (Esterman)

Visual search (find the number)

Reading (corrected reading speed)

Tasks simulating ADL: 1) driving hazard perception (mean score per hazard), 2) obstacle avoidance (completion time), 3) visuomotor search (time)

Aimola 2011 (Continued)

Attention tasks: 1) sustained attention to response (mean percentage error score), 2) test of everyday attention

Subjective questionnaires: Visual Functioning Questionnaire (VFQ-25), Subjective Reasons Questionnaire

Time points when outcomes were assessed: before and after intervention

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	Stated "using parallel trial allocation software"
Blinding (performance bias and detection bias) All outcomes	Unclear risk	"Participants were informed about the training types but did not know to which group they were assigned." Not stated if outcome assessor was blinded.
Incomplete outcome data (attrition bias) All outcomes	High risk	"There were 18 withdrawals, 9 due to low motivation."
Other bias	Low risk	No other issues noted

Bainbridge 1994

Methods

Substitutive intervention versus substitutive intervention

Design: parallel RCT

Stratification: not stated

Randomisation sequence: not stated

Comparisons: Full Field plastic press-on 15-diopter Fresnel prisms (base towards the deficit) versus Half Field prisms cut to cover the respective homonymous field

Allocation concealment: not stated

Blinding: not stated

Power calculation: not stated

Intention-to-treat analysis: not stated

Other recruitment details: not stated

Patient and public involvement: not stated

Participants

Total study population: 18 (stroke only) participants who had hemianopsia or visual neglect, or both

Withdrawals: none

Method of diagnosing VFD: confrontation; method of diagnosing visual perceptual problems: confrontation

Characteristics of population: participant details are listed in [Table 2](#)

Bainbridge 1994 (Continued)

Type and severity of visual problems: participant details are listed in [Table 3](#)

Inclusion criteria: not stated

Exclusion criteria: not stated

Baseline comparison of treatment groups: not stated

Interventions	<p>Group 1: Full field prism (n = 10)</p> <p>Intervention type: substitution. Materials: Full field prism. Where can materials be accessed? not stated. Procedures: 15 dioptre full-field plastic press-on Fresnel prisms (base towards the deficit). Provided by: not stated. Delivery: face-to-face, individual, location not stated (Table 1). Regimen: used the prisms while awake for 4 weeks. Tailoring: not stated. Modification: not stated. Adherence: not stated</p> <p>Group 2: Hemi-field prism (n = 8)</p> <p>Intervention type: substitution. Materials: Hemi-field 15 diopter Fresnel prism. Where can materials be accessed? not stated. Procedures: Hemi-field prisms cut to cover the respective field. Provided by: not stated. Delivery: face-to-face, individual, location not stated (Table 1). Regimen: used the prisms while awake for 4 weeks. Tailoring: not stated. Modification: not stated. Adherence: not stated</p>	
Outcomes	<p>See Table 4</p> <p>Motor visual perception score</p> <p>Line Bisection Test</p> <p>Line Cancellation test</p> <p>Harrington-Flocks Visual Field score</p> <p>Time points when outcomes were assessed: baseline and at 4 weeks</p> <p>Adverse events? not stated</p>	
Notes	<p>Insufficient information to assess for any confounding variables (abstract)</p>	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Quote "patients were randomized". Insufficient information available in the abstract
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Insufficient information available in the abstract
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Insufficient information available in the abstract
Other bias	Unclear risk	Insufficient information available in the abstract

Bowers 2014

Methods	Substitutive intervention versus control, placebo or no intervention
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Interventions for visual field defects in people with stroke (Review)

Bowers 2014 (Continued)

Design: RCT (cross-over, with a "counterbalanced AB/BA design (AB = real first, BA = sham first)". The "real" prisms could be either "oblique" or "horizontal" peripheral prism glasses, meaning that there were 4 possible treatment allocations for each participant).

Stratification: see below (minimisation)

Randomisation sequence: participants were assigned using minimisation. "The first participant was assigned randomly, with each subsequent participant assigned in such a way as to minimize imbalances among the 4 treatment allocations. We could realistically balance for only 2 factors. Study site was the primary factor (because continuation rates varied significantly across sites in our first multicenter study) and side of hemianopia (right or left) was the second factor (because the side of the lesion could potentially affect performance with the prism glasses). We did not balance for age because it was not a significant factor affecting continuation rates in our previous study."

Comparisons: oblique peripheral prism glasses, horizontal peripheral prism glasses, sham prism glasses

Allocation concealment: yes

Blinding: "Double-masking was used, with participants and data collectors being masked as far as possible. In addition, the principal investigator who conducted data analyses was masked. However, it was impossible to mask all study personnel; there was an unmasked practitioner at each site who fitted the prism glasses and dealt with clinical aspects of patient care."

Power calculation: yes (data from previous trial used to calculate minimum sample size; led to "plan to enrol at least 68 participants")

Intention-to-treat analysis: no

Other recruitment details: "Data were collected at 13 study sites, including the Peli laboratory at Schepens, 11 vision rehabilitation clinics in the United States, and 1 in the United Kingdom. The clinics included university, hospital, and private practice clinics. Each site recruited a median of 7 participants (range, 3-12). Participants were recruited by practitioners at each study site."

Patient and public involvement: not stated

Participants

Total study population: 73 randomised

Withdrawals: 6 (before the start of the cross-over)

Method of diagnosing VFD: visual field mapping extended to at least 50° from fixation in all directions and was performed using Goldmann perimetry (V4e target), a Humphrey Field Analyzer 120-point full-field screening test, or similar tests, depending on the equipment available at each clinic.

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#)

Inclusion criteria: "Complete homonymous hemianopia of greater than 3 months' duration, no visual neglect (Bells test and Schenkenberg Line Bisection test), and no history of having worn peripheral prism glasses. In addition, participants had corrected monocular visual acuity of at least 20/50 in each eye, refractive error within the -5 diopter (D) to +5 D range, no strabismus, no significant cognitive decline (Short Portable Mental Status Questionnaire), and no balance problems or other deficits that could impair ability to walk or use the prism glasses. To ensure that study inclusion criteria were uniformly applied, screening data were sent to the principal investigator (A.R.B.), who determined eligibility".

Exclusion criteria: none listed

Baseline comparison of treatment groups: yes (mobility questionnaire)

Interventions

Group 1: Prism glasses (total n = 37)

Group 1a. Real oblique prism glasses (n = 19)

Bowers 2014 (Continued)

Group 1b. Real horizontal prism glasses (n = 18)

Intervention type: substitution. **Materials:** peripheral prism glasses of 57 prism dioptres (intervention) versus 5 prism dioptres (sham). **Where can materials be accessed?** From vision specialists; for the study these were manufactured by "Chadwick Optical, Inc. (White River Junction, Vermont)". **Procedures:** participant fitted and provided with prism glasses to be worn for the following 4 weeks. "Training in how to use the prism glasses was conducted by the practitioner." "Participants were taught to view through the central prism-free area of the spectacle lens at all times and to turn the head and eyes to fixate objects of interest that were initially detected from the prism image in peripheral vision. A simple "reach and touch" training exercise was used to familiarize participants with the relationship between the apparent and real positions of objects detected from the prism image; this exercise was also encouraged for home-training. Participants were given verbal and written instructions about how to use the prism glasses and were encouraged to wear them as much as possible each day. They were advised not to use the peripheral prism glasses for driving or prolonged reading." **Provided by:** "practitioner". **Delivery:** face-to-face, individual, clinic appointment to be provided with the glasses, then to be worn by participant (Table 1). **Regimen:** first pair of prisms glasses 4 weeks; then cross-over and use of second pair of prism glasses for another 4 weeks (see procedures). **Tailoring:** not stated. **Modification:** not stated. **Adherence:** not stated

Group 2: Sham prism glasses (total n = 36)
2a. Sham oblique prism glasses (n = 17)
2b. Sham horizontal prism glasses (n = 19)

Intervention: control. **Intervention type:** placebo control. **Materials:** sham prism glasses. **Where can materials be accessed?** For the study these were manufactured by "Chadwick Optical, Inc. (White River Junction, Vermont)". **Procedures:** as for real prisms. **Provided by:** "practitioner". **Delivery:** face-to-face, individual, clinic appointment to be provided with the glasses, then to be worn by participant (Table 1). **Regimen:** first pair of prisms glasses 4 weeks; then cross-over and use of second pair of prism glasses for another 4 weeks (see procedures). **Tailoring:** not stated. **Modification:** not stated. **Adherence:** not stated

Outcomes	See Table 4	
	The proportion of participants fitted with each type of prism glasses for whom the decision is to continue using the glasses	
	Difference in perceived mobility (i.e. rating of how helpful each type of prism glasses are in avoidance of obstacles when walking)	
	Time points when outcomes were assessed: at the end of each cross-over period	
Notes	Data presented for the first phase (i.e. before the cross-over) is presented for "real" versus "sham" prisms, i.e. with data from the oblique and horizontal groups combined: "there were no statistically significant differences between the oblique and horizontal groups for any of the outcome measures ... therefore, data were pooled across the 2 groups for the main analyses".	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	"The clinical coordinator at Schepens assigned participants to 1 of 4 possible treatment allocations (real oblique AB/BA and real horizontal AB/BA) using minimization". Allocation was done by central clinical co-ordinator.
Blinding (performance bias and detection bias) All outcomes	Low risk	Outcome assessor was blinded. The personnel fitting the prism were not blinded: "Double-masking was used, with participants and data collectors being masked as far as possible. In addition, the principal investigator who conducted data analyses was masked. However, it was impossible to mask all study personnel; there was an unmasked practitioner at each site who fitted the prism glasses and dealt with clinical aspects of patient care". In the debriefing,

Bowers 2014 (Continued)

61% (37/61) of participants reported that they thought that one pair of glasses might have been a sham; of these, 92% (34/37) correctly identified the sham.

Incomplete outcome data (attrition bias) All outcomes	Low risk	Reasons for dropouts were reported and were balanced across the groups.
Other bias	High risk	Authors have disclosed conflicts of interest: "Dr Peli has financial interest in a patent related to the peripheral prism glasses (assigned to Schepens Eye Research Institute). Ms Keeney has licensed that patent for Chadwick Optical, Inc. Chadwick Optical, Inc funded the study in part from National Institutes of Health (NIH) grant EY014723 through a subcontract with Schepens Eye Research Institute. Dr Peli was a paid consultant to Chadwick Optical, Inc on the design of the permanent prisms. No other financial interests were reported."

De Haan 2015

Methods	<p>Compensative intervention versus control, placebo or no intervention</p> <p>Design: RCT</p> <p>Stratification: minimisation ("minimised regarding gender, side of field defect (left or right), side of field defect (hemianopia versus quadrantanopia), age (younger versus older than 55), and time since onset (shorter versus longer than 12 months - because time since onset was assumed less important than the other variables, this variable was weighted less heavily than the others)</p> <p>Randomisation sequence: randomisation software</p> <p>Comparisons: 2 groups, training group and control group</p> <p>Allocation concealment: no, "author GH entered the characteristics of the patient into the randomisation software ... which resulted in allocation to the training group or the waiting list control group"</p> <p>Blinding: "Assessors were blinded to participants' group allocation".</p> <p>Power calculation: yes</p> <p>Intention-to-treat analysis: no; some participant data excluded from analysis due to deviations from study protocol ("too low compliance with training protocol", "time period between T1 and T2 longer than acceptable")</p> <p>Other recruitment details: patients were recruited at Royal Dutch Visio and Bartiméus, the two centers of expertise for blind and partially sighted people in the Netherlands.</p> <p>Patient and public involvement: not described</p>
Participants	<p>Total study population: 54 randomised; training group (n = 30), control group (n = 24)</p> <p>Withdrawals: at follow-up immediately after training; training group (n = 28), control group (n = 24). Additional participants excluded from analysis, so in analysis: training group (n = 26), control group (n = 23).</p> <p>Method of diagnosing VFD: specific method not clear: "patients underwent extensive and standardized ophthalmological and neuropsychological assessments at the centers mentioned above prior to participation in the study."</p> <p>Characteristics of population: participant details are listed in Table 2.</p> <p>Type and severity of visual problems: participant details are listed in Table 3.</p>

De Haan 2015 (Continued)

Inclusion criteria: "The main inclusion criterion was presence of a hemi-VFD, at least a quadrantanopia, restricted to one half of the visual field, due to acquired postchiasmatic brain injury ... To be included, patients required a minimum binocular visual acuity of Snellen 0.5 (6/12 or 20/40, LogMAR 0.3), a stable neurological and ophthalmological condition, non-disturbed eye and head motility, ability to walk at least 50 meters, and a Mini Mental State Examination (MMSE) score 24 out of 30."

Exclusion criteria: "Exclusion criteria were ocular diseases affecting the visual field or binocular visual acuity, signs of severe physical impairments or (neuro)psychological disorders. Neglect was excluded based on the Balloons, Drawings, Line Bisection and Rey Complex Figure Test."

Baseline comparison of treatment groups: yes

Interventions

Group 1: InSight-Hemianopia Compensatory Scanning Training (IH-CST) (n = 26)

Intervention type: compensation. **Materials:** "The full training protocol is available at Visio and Bartimeus for occupational therapists trained for applying the protocol". Electronic software was needed: "software was needed in order to implement the exercises ... this software was developed by the faculty of Behavioural and Social Science at the University of Groningen". **Procedures:** "The aim of the IH-CST is to teach patients with HVFD to compensate for their visual field defect during a wide range of mobility-related activities ... The compensation strategy taught in the IH-CST is to apply a scanning rhythm consisting of a triad of saccades. First a large saccade towards the blind side is made in order to receive information from the periphery. This is followed by a second saccade back towards the seeing side to prevent overcompensation. Third, a small saccade is made back to the starting point of looking straight forward ... The training program consists of exercises for improving awareness of the visual field defect and its consequences for daily life, exercises to learn the scanning rhythm, and practice of the scanning rhythm in daily life mobility situations". **Provided by:** occupational therapists: "Thirty occupational therapists ... experienced in working with brain injured patients were schooled in providing the IH-CST ...". **Delivery:** "Training according to the protocol was provided in Dutch at nine locations of Royal Dutch Visio and one location of Bartimeus in the Netherlands." "Homework assignments are included in the training protocol". (Table 1). **Regimen:** 15 individual sessions of 60 to 90 minutes each, 18.5 hours of face-to-face training in total during a period of 10 weeks. **Tailoring:** yes (stated "not applicable" in author's checklist, but stated that "... the patient proceeds to the next exercise once the predefined targets or an exercise are accomplished. This creates flexibility for individual needs and progress and can cause the training to take more or less than 15 sessions"). **Modification:** no. **Adherence:** "patients are asked to keep a diary of their practice at home and the therapist asks about the progress of the homework assignment at the beginning of every training session".

Group 2: Wait-list control (n = 23)

Intervention type: control. **Materials:** none. **Where can materials be accessed?** NA. **Procedures:** NA. **Provided by:** NA. **Delivery:** NA. (Table 1) **Regimen:** NA. **Tailoring:** NA. **Modification:** NA. **Adherence:** NA.

Outcomes

See Table 4

Tests for visual functions (Goldman perimetry)

Reading tests

Basic scanning tests

Hazard perception test

Tracking task

Obstacle course

Questionnaires for activity and participation

Time points when outcomes were assessed: participants in the training group were assessed the week before training (T1) and the week after training (T2); participants in the control group were assessed at the same time points (T1 and T2).

De Haan 2015 (Continued)

Notes

Trial registration details: ISRCTN Registry (ID ISRCTN16833414); Central Committee on Research Involving Human Subjects (CCMO; registration number NL31718.042.10)

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	Random allocation, using randomisation software. Researcher (GH) was aware of allocation, but was not involved in providing treatment or assessment
Blinding (performance bias and detection bias) All outcomes	Low risk	Outcome assessors were blinded. Personnel providing training and participants could not be blinded.
Incomplete outcome data (attrition bias) All outcomes	High risk	No intention-to-treat analysis. Final analysis completed only on those who completed training, and who complied with the training programme.
Other bias	High risk	Control group received no treatment or contact with personnel. Therefore, high risk of bias due to attention received by training group.

Elshout 2016

Methods

Restitutive intervention versus control, placebo or no intervention
Design: randomised cross-over trial

Stratification: not stated

Randomisation sequence: "For each cohort of 10 patients, numbers (J01–J40), the training stimuli (FloworPoint), and the order of training rounds [Test (defect) round first or Control (intact) round first] were randomly assigned to a patient number ... The patients were included in order of registration on our website. Thus, the assignment of a patient to a patient number (with corresponding training scheme: stimulus type and the training order) was determined prior to the first inclusion, completely random and not based on selection."

Comparisons: 2 groups: "(1) test (high contrast training of the affected field) and (2) control (low-contrast training of the intact visual field)"

Allocation concealment: see randomisation sequence

Blinding: "We applied both Humphrey (blinded) and Goldmann perimetry (not blinded, because of insufficient staffing)." Not stated for reading tests

Power calculation: not stated

Intention-to-treat analysis: no. "Three patients dropped out during the first training round for personal reasons and were excluded from analyses".

Other recruitment details: "This study was part of a larger project ... Forty stroke patients with visual field defects due to post-chiasmatic stroke were included following written informed consent. Patients throughout the Netherlands could sign up for our study voluntarily by filling in a form on our website." However after 3 cohorts of 10 patients, the training procedure was modified "because in some patients the control training reversed the increase of the visual field of the preceding defect training". This study therefore only reported the first 3 cohorts of participants.

Patient and public involvement: not stated

Participants

Total study population: 30 patients with post-chiasmatic lesions

Elshout 2016 (Continued)

Withdrawals: 3 patients dropped out "for personal reasons". They were excluded from analyses.

Method of diagnosing VFD: Goldmann perimetry measurement

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: "Patients in the chronic phase of stroke (> 10 months post CVA) were included if they showed no signs of visual neglect (line bisection test). Patient age was between 18 and 75 years, and they were able to undergo (f)MRI scanning."

Exclusion criteria: visual neglect (see above)

Baseline comparison of treatment groups: not stated

Interventions

Group 1: defect training (high contrast stimuli offered within the field defect along its border) (n = 15 for first phase, before cross-over)

Intervention type: restitution - high-contrast stimuli in affected hemifield. **Materials:** "Each patient received a training unit at home to create a controlled training environment. This unit consists of a container, to be placed on a table, with a top cover and side covers to present a dark visual surround for the training stimuli with the exclusion of stray light. Mini Mac computer, keyboard, and mouse, a support with 24" LED monitor, webcam, and chin/headrest were positioned inside the matte black container". **Where can materials be accessed?** not stated. **Procedures:** "Viewing distance was fixed at 40 cm. The subject's face was indirectly lighted with a TL light for eyetracking with the webcam. The computer was prepared with eyetracking software, and training programs that were adjusted to the particular visual field defect of the patient." "For both stimuli, the patient maintained fixation binocularly on a ring (diameter = 0.5°) at the center of the screen. During stimulus presentation (7s), patients shifted attention covertly (i.e. without shifting eye fixation) toward the stimulus and responded using the keyboard. Only the fixation point was shown during the intertrial interval of 2s". "To cue the stimulated target location and to perform a discrimination task, a line was presented simultaneously with the point extending from the fixation target into the trained hemifield. The meridional angle of the line differed by 10° from the training point. The patient made a forced-choice response whether the point stimulus was presented clockwise or counterclockwise relative to the presented line". **Provided by:** Not stated. **Delivery:** In patients home. ([Table 1](#)). **Regimen:** "1h a day, 5 days a week during 8 weeks to complete at least 40h of training per hemifield". "The length of one training session was on average 12min (depending on the number of trials set per session and amount of fixation errors). The number of trials in a session ranged between 60 and 100, depending on the shape and quality of the visual field defect. The stimuli were randomly presented for each session". **Tailoring:** The number of trials depended on the shape and quality of the visual field defect. "Point size was at least 0.2° in diameter (at 1° eccentricity) and was scaled with eccentricity: scale (E) = $(0 : 0006E^{\wedge}2 + 0:0448 E + 0:092) = 0:1374$ ". **Modification:** not stated. **Adherence:** was monitored: "Throughout the training, fixation was monitored via a low-cost commercial webcam and eyetracking software available in the public domain (www.inference.phy.cam.ac.uk/opengazer/) that was adapted to supply eye position data to the training program".

Group 2. Control (intact training) (stimuli presented in the intact field about the same eccentricities as for defect training) (n = 15 for first phase, before cross-over)

Intervention type: placebo/control. Although paper stated: "Following three cohorts of 10 patients, we modified the training procedure, because in some patients the control training reversed the increase of the visual field of the preceding defect training". **Materials:** as above. **Where can materials be accessed?** not stated. **Procedures:** similar to above defect training: "During intact training, stimuli were presented within the intact field at about the same eccentricities as for the patient's defect training ... To offer a challenging training during the intact training, the stimulus contrast was reduced ($C < 0.15$). "Within the entire visual field, a pattern of flow was shown that contracted on to a training location within the visual field (white points on a dark screen). The stimulus to be discriminated was placed on a black disk that covered the center of the contraction pattern, the diameter of which was eccentricity scaled with the same factor as for the point target. We used a minimal disk size of 1.7° at 1° eccentricity. The origin of the contraction pattern was the location cue for the flow discontinuity that had to be detected. The discontinuity stimulus (on the disk) was a flow pattern rotating clockwise or counter-

Elshout 2016 (Continued)

clockwise about the training location. The patient had to indicate the direction of rotation". **Provided by:** not stated. **Delivery:** in participant's home (Table 1). **Regimen:** as above. **Tailoring:** stimuli were presented at the same eccentricities as for the defect training, which depended on the shape and quality of the visual field defect. **Modification:** not stated. **Adherence:** as above

Outcomes	See Table 4 Perimetry: Humphrey (blinded) and Goldmann perimetry (not blinded because of insufficient staffing) Reading tests: reading speed
Notes	Supplementary material referred to within published paper, but not accessible via weblink. Requested from authors February 2019

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	Patients were included in the order in which they registered on the website. Order of treatment phases was randomly assigned, and allocated to patient number prior to patient registrations.
Blinding (performance bias and detection bias) All outcomes	High risk	No blinding for Goldmann Perimetry ("because of insufficient staffing"). Humphrey perimetry was blinded. Not stated for reading tests. Unclear who delivered the intervention and whether they were blind to the treatment phase.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No intention-to-treat analysis. 3 dropouts during first phase "for personal reasons", but unclear which treatment group they were in.
Other bias	Low risk	No other issues noted

Gall 2013

Methods	Restitutive intervention versus control, placebo or no intervention Design: RCT Stratification: not stated Randomisation sequence: not stated Comparisons: 2 groups Allocation concealment: not stated Blinding: stated "blinded" Power calculation: not stated Intention-to-treat analysis: not stated Other recruitment details: not stated Patient and public involvement: not stated
Participants	Total study population: 29 patients with post-chiasmatic lesions Withdrawals: not stated

Gall 2013 (Continued)

Method of diagnosing VFD: not stated

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: not stated

Exclusion criteria: not stated

Baseline comparison of treatment groups: not stated

Interventions	<p>Group 1: non-invasive brain stimulation using alternating current stimulation (ACS) (n = 15)</p> <p>Intervention type: restitution - alternating current stimulation. Materials: not stated. Where can materials be accessed? not stated. Procedures: not stated. Provided by: not stated. Delivery: face-to-face, individual, location unclear (Table 1). Regimen: 10 days. No other details reported. Tailoring: not stated. Modification: not stated. Adherence: not stated</p> <p>Group 2: sham stimulation (n = 14)</p> <p>Intervention type: placebo/control. Materials: not stated. Where can materials be accessed? not stated. Procedures: not stated. Provided by: not stated. Delivery: face-to-face, individual, location unclear (Table 1). Regimen: 10 days. No other details reported. Tailoring: not stated. Modification: not stated. Adherence: not stated</p>
Outcomes	<p>See Table 4</p> <p>Perimetric thresholds within areas of residual vision</p> <p>Subjectively perceived visual functioning/vision-related quality of life (composite score of National Eye Institute Visual Function Questionnaire 39)</p> <p>Health-related quality of life (Short Form Health Survey SF-12)</p> <p>Time points when outcomes were assessed: baseline and post-intervention (time point not stated)</p>
Notes	Insufficient information to assess for any confounding variables (abstract)
Risk of bias	
Bias	Authors' judgement Support for judgement
Allocation concealment (selection bias)	Unclear risk Insufficient information available in the abstract
Blinding (performance bias and detection bias) All outcomes	Unclear risk Insufficient information available in the abstract
Incomplete outcome data (attrition bias) All outcomes	Unclear risk Insufficient information available in the abstract
Other bias	Unclear risk Insufficient information available in the abstract

Jarvis 2012

Methods	Assessment or screening intervention versus control, placebo or no intervention
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Jarvis 2012 (Continued)

Design: pilot RCT

Stratification: none

Randomisation sequence: "randomized into one of two groups using a computer-generated randomization table"

Comparisons: 2 groups: Group A (control) and Group B (experimental)

Allocation concealment: not stated ("randomization process was administered by a researcher at the University of Liverpool, who was not involved in data collection but was involved in the later data analysis")

Blinding: participants were blinded ("participants were masked to group allocation"), "assessors were not masked" ("findings from the visual assessment were withheld from the therapy staff in group A (control group). In comparison the visual assessment details were made available to the therapy staff for participants in group B (experimental group)").

Power calculation: feasibility study ("prospective observation cohort study in the UK suggested that of all stroke patients referred with suspected visual impairment, 85% were found to have an identifiable visual impairment (Rowe 2007). On the basis of this preliminary data, this pilot study aimed to screen 100 patients in order to recruit a minimum of 70 participants").

Intention-to-treat analysis: no ("All data analysis was conducted based on the recruited patients to each group with full FIM data collection for both baseline and 6-week follow-up assessment").

Other recruitment details: "Participants were prospectively recruited between February 2008 and July 2009". "Specified members of the health care team (nurses, stroke physicians, physiotherapists and occupational therapists) on the stroke unit were required to screen patients against these criteria to identify potential participants. Staff used a screening form with questions to identify visual signs (head turning, strabismus, ptosis) and symptoms (diplopia, loss of vision, field loss). This was adapted from the screening form used in the Visual In Stroke (VIS) study (Rowe 2010)". "If visual impairment was noted, the screening form was sent to the orthoptist and hence, it doubled as a referral form". "Where necessary, for example when a potential participant had communication difficulties, adaptations were made to the consent process".

Patient and public involvement: not stated

Participants

Total study population: 64

Withdrawals: 13 died before full baseline assessment (Group A: 4; Group B: 9). Another 10 cases (Group A: 6, Group B: 4) withdrew (death, early discharge, no follow-up or DNA follow-up) at follow-up assessment.

Method of diagnosing VFD: full visual assessment undertaken by orthoptist. Battery of routine tests used as part of a previous study (Rowe 2009). It comprised tests of visual acuity - logMAR (Bailey 1976), ocular alignment - cover test (Pediatric Eye Disease Investigator Group 2009), ocular motility - saccadic, smooth pursuit and vergence assessment (Holmes 2001), stereopsis - Frisby test (Rosner 1984), visual field - confrontation (Cassidy 2001) and visual inattention - line bisection, star cancellation, and clock drawing. "The orthoptist did not suggest possible adaptive strategies to be undertaken, the focus was on alterations made by the therapists based only on the visual assessment information".

Characteristics of population: participant details are listed in Table 2.

Type and severity of visual problems: participant details are listed in Table 3.

Inclusion criteria: "stroke within 2–6 weeks of being recruited, had a decreased functional ability compared to pre-stroke functioning, had a post-stroke visual impairment, and were able to understand the research process"

Exclusion criteria: "unable to consent due to cognitive impairment or communication difficulties, or if they had a visual field impairment pre-existing their stroke"

Jarvis 2012 (Continued)

Baseline comparison of treatment groups: "no significant difference in the composition of both groups (χ^2 test) for gender ($P = 0.846$), age ($P = 0.113$), stroke type ($P = 0.564$), stroke area ($P = 0.499$), stroke laterality ($P = 0.396$) and handedness ($P = 0.268$)"

Interventions

Group 1: experimental (n = 24 full baseline assessment; n = 20 at follow-up assessment)

Intervention type: assessment or screening. **Materials:** modified screening form. **Where can materials be accessed?** Screening form (modified) available from the Visual In Stroke (VIS) study (Rowe 2010). **Procedures:** "all participants underwent a full visual assessment by an orthoptist. The findings from the visual assessment were ... made available to the therapy staff for participants in group B (experimental group)". "participants in both groups received occupational therapy and physiotherapy". "Therapy routinely included working to regain motor activity and increase ability to achieve valued functional tasks. The therapists used strategies such as visual scanning and cueing to the affected side as part of their practice". Treatment in the experimental group was informed by the results of the visual assessment. **Provided by:** orthoptists. No details provided about their training. **Delivery:** face-to-face, location inpatient, stroke unit (Table 1). **Regimen:** not stated. **Tailoring:** yes. Details not supplied. **Modification:** not stated. **Adherence:** not stated

Group 2: control (n = 27 full baseline assessment; n = 19 at follow-up assessment)

Intervention type: control/standard care. **Materials:** NA. **Where can materials be accessed?** NA. **Procedures:** described above. Treatment in the control group was not informed by results of the visual assessment ("all participants underwent a full visual assessment by an orthoptist. The findings from the visual assessment were withheld from the therapy staff in group A (control group)"). **Provided by:** OT, physiotherapy. No details provided about their training. **Delivery:** face-to-face, individual, location inpatient (Table 1). **Regimen:** NA. **Tailoring:** NA. **Modification:** NA **Adherence:** NA

Outcomes

See Table 4

Functional independence measure (FIM)

Timed 10 m walk

Non-validated questionnaire ("Therapist with the most contact with each participant was asked to complete a non-validated questionnaire giving qualitative information about their treatment approach." "Two versions of this questionnaire. The group A questionnaire asked the therapist to justify their treatment approach. The group B questionnaire required the therapist to comment on whether their treatment approach had been influenced by the visual assessment").

Time points when outcomes were assessed: "baseline and at 6 weeks after baseline (or on discharge if this occurred earlier)"

Notes

Review author Fiona Rowe was involved in this trial.

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	"computer generated randomization table. The randomization process was administered by a researcher at the University of Liverpool, who was not involved in data collection but was involved in the later data analysis. Participants were masked to group allocation, but the assessors were not masked in this trial".
Blinding (performance bias and detection bias) All outcomes	High risk	"Participants were masked to group allocation, but the assessors were not masked in this trial." "The qualitative study indicated an inherent bias had been introduced to this trial, due to the inability to blind carers, and assessors to group allocation. The health care team perceived that the presence of a full baseline vision assessment enhanced their awareness of the effect of visual deficits following stroke.

Jarvis 2012 (Continued)

		This was regardless of whether or not the full visual assessment details were available."
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No intention-to-treat analysis. Dropouts were clearly reported, but reason for not attending was not clear and could be related to the intervention.
Other bias	Low risk	No other concerns noted

Jobke 2009

Methods	Restitutive intervention versus restitutive intervention Design: RCT (cross-over) Stratification: no Randomisation sequence: no details reported ("randomly assigned them into 2 groups") Comparisons: 2 groups, extrastriate vision restoration therapy versus conventional vision restoration therapy Allocation concealment: not stated Blinding: "double-blind" Power calculation: no Intention-to-treat analysis: no Other recruitment details: none Patient and public involvement: no
Participants	Total study population: 21 (mixed population) participants, with visual field defects, lesions over 1 year old Group 1: 8 participants, Group 2: 10 participants. The study included a mixed population. Withdrawals: "Three patients had to be excluded from the analysis. One male patient (patient 19) used another vision therapy program not included in this study. Another patient (patient 20) was excluded because he discontinued the training for more than 4 weeks during the trial and a female patient (patient 21) showed poor fixation performance (70% rather than the minimally required 90%)." Method of diagnosing VFD: high-resolution perimetry and "common static perimetry diagnosis at their own ophthalmologists office". Method of diagnosing visual perceptual problems: not stated Characteristics of population: participant details are listed in Table 2 Type and severity of visual problems: participant details are listed in Table 3 . Inclusion criteria: at least 6 months prior experience of vision restoration therapy Exclusion criteria: none listed Baseline comparison of treatment groups: yes
Interventions	Group 1: extrastriate vision restoration therapy (VRT) (n = 8) Intervention type: restitution. Materials: extrastriate vision restoration therapy - not stated where this could be accessed/acquired

Jobke 2009 (Continued)

Procedures: stimulation of the "entire blind area with a massive moving spiral", plus standard VRT in areas of residual vision. "The spiral consisted of concentric circles which moved with a frequency of 2.5 Hz, and increased in width to generate the perception of motion". "Additionally, areas of residual vision (relative defects) were stimulated by standard VRT, i.e. the common single-point stimulation paradigm. Just like in standard VRT, the patient had to respond to the white target stimuli presented at random locations in areas of residual vision." **Provided by:** not stated. **Delivery:** training carried out at home by participants (Table 1). **Regimen:** extrastriate VRT, 1/2 hour daily for 90 days **Tailoring:** Yes. "The size of the training area varied from patient to patient according to the size of the visual field defect". **Modification:** Yes: "Every week the patients completed a self-administered diagnostic test at home and sent their data to the institute for updating the training region if necessary." **Adherence:** "Training breaks of longer than 2 weeks led to exclusion of the patient."

Group 2: conventional vision restoration therapy (VRT) (n = 10)

Intervention type: restitution. **Materials:** standard visual restoration therapy - not stated where this could be accessed/acquired. **Procedures:** standard vision restoration therapy consisted of single white target stimuli, presented at random in areas of residual vision. The participant had to respond to the white target stimuli presented at random locations in areas of residual vision. **Provided by:** not stated. **Delivery:** training carried out at home by participants (Table 1). **Regimen:** conventional vision restoration therapy, 1/2 hour daily for 90 days. **Tailoring:** see above. **Modification:** see above. **Adherence:** see above

Outcomes

See Table 4

High-resolution perimetry

Perimetry: common static perimetric diagnosis at own ophthalmologist

Near visual acuity: Radner reading test

NEI-VFQ

Zahlen-Verbindungs test: speed of connecting numbers in a paper-pencil test

Time points when outcomes were assessed: before and after each cross-over period

Notes

1. "All patients had used VRT for at least 6 months (range from 6-36 months) prior to study entry and were therefore experienced participants. We were aware that this might reduce the power of the therapy because of a possible 'ceiling effect'. To eliminate training bias, no patient carried out VRT during the 6 months preceding the study". But this also means that this was potentially a very selective group of patients, i.e. those who were motivated to continue using VRT after an initial experience. Therefore, it may be inappropriate to generalise from this study to the general population of patients with visual function defects. There was no data on the type or length of each participant's previous VRT (just the range of 6 to 36 months), and if there were previous changes in the functions measured in this test. In his previous paper on VRT, (Kasten 2001) concluded that patients can be split into 3 main types after scanning training: 1) those whose field increases both during and after training, 2) those whose field increases during training, but decrease afterwards, and 3) those who show no change at all. If true, this would impact on the results of this study, as types 1) and 3) do not benefit from further training.

2. Participants did not use their glasses for treatment and measurements. Measurements included a measure of near visual acuity; given the age of patients, most will be presbyopes, so will struggle with this task, dependent on age and prescription.

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Quote "random assignment", no further details given
Blinding (performance bias and detection bias)	Unclear risk	"Double-blind" was in the title, but no further information appeared in the paper

Jobke 2009 (Continued)

All outcomes

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	3 participants excluded from analysis and not included in demographic details. "One patient used another vision therapy program not included in this study. Another patient was excluded because he discontinued the training for more than 4 weeks during the trial and a patient showed poor fixation performance (70% rather than the minimally required 90%)." Some of these dropouts may have occurred because of an issue directly associated with the intervention studied Also, missing data for standard perimetry measures
Other bias	High risk	Researchers may have had financial interest in intervention being investigated. Poggel 2004 stated "BA Sabel and E Kasten are inventors. BA Sabel is a consultant and shareholder of NovaVision Inc, and has equity in excess of \$10,000."

Kasten 1998

Methods	<p>Restitutive intervention versus control, placebo or no intervention</p> <p>Design: RCT</p> <p>Stratification: not stated</p> <p>Randomisation sequence: not stated; correspondence with author did not provide any further detail</p> <p>Comparisons: visual restitution training (computer-based training) versus placebo (fixation training program)</p> <p>Allocation concealment: stated "randomly assigned (double-blind)"; no further details</p> <p>Blinding: "double-blind"</p> <p>Power calculation: no</p> <p>Intention-to-treat analysis: no (1 dropout from placebo group)</p> <p>Other recruitment details: not stated</p> <p>Patient and public involvement: not stated</p>
Participants	<p>Total study population: 19</p> <p>Withdrawals: 1 ("one patient from the placebo group failed to meet the requirement to train for 150 h. This patient dropped out of the study").</p> <p>Method of diagnosing VFD: high-resolution perimetry and Tubinger automated perimetry</p> <p>Characteristics of population: participant details are listed in Table 2.</p> <p>Type and severity of visual problems: participant details are listed in Table 3.</p> <p>Inclusion criteria: "patients had to have both a visual-field defect and post-chiasmatic or optic nerve damage as shown by CT, MRI, surgical records or ophthalmoscopic documentation of optic nerve atrophy".</p> <p>Exclusion criteria: insufficient fixation ability, neglect, non-optic nerve heteronymous visual field defect, disorders of the eye, no residual vision, no visual deficit, age > 75 years, age < 18 years, died, lesion age < 12 months, epilepsy or photosensitivity, cognitive deficits, not willing to participate in trial, no shows after initial screening</p>

Kasten 1998 (Continued)

Baseline comparison of treatment groups: baseline comparison was presented for combined groups of 19 post-chiasmatic injury participants and 19 participants with optic nerve injury (see notes below).

Interventions	<p>Group 1: visual restitution therapy (n = 9)</p> <p>Intervention type: restitution. Materials: computer-based training. Procedures: during vision restoration therapy, the participant maintained fixation on a central fixation spot, with their "visual border zone" stimulated by repetitive stationary stimuli of varying luminance presented in a random location within this border zone. Provided by: not stated. Delivery: training was carried out on a personal computer placed in a darkened room at home. (Table 1). Regimen: 1 hour per day, 6 days per week for 6 months (a total of 150 hours required). Tailoring: yes "compliance checks and adjustments of training difficulty level were done monthly". Modification: yes "we used an individually adapted training protocol, which was determined by the characteristics of the transition zone". Adherence: a total of 150 hours was required (compliance checks done monthly).</p> <p>Group 2: placebo (fixation training) (n = 10)</p> <p>Intervention type: placebo. Materials: computer-based training. Procedures: the fixation training program required "eye movements to stimuli near the foveal region for a comparable amount of time". Provided by: not stated. Delivery: training was carried out on a personal computer placed in a darkened room at home. (Table 1). Regimen: 1 hour per day, 6 days per week for 6 months (a total of 150 hours required). Tailoring: not stated. Modification: not stated. Adherence: a total of 150 hours was required (compliance checks done monthly).</p>
Outcomes	<p>See Table 4</p> <p>Visual field: high-resolution perimetry</p> <p>Visual field: Tubinger automated perimetry</p> <p>Visual acuity: Landolt ring to give minimum angle of resolution</p> <p>Quality of life questionnaire</p>
Notes	<p>1. Reports of this trial combined data from the trial of 19 post-chiasmatic injury participants with data from a trial of 19 participants with optic nerve injury. However, within the paper it did state "The data reported here are from two independent clinical trials each with an experimental and a control group. In the first trial, two groups of optic nerve injury patients ... In the second trial, patients with post chiasmatic injury were randomly assigned ...". It is the second trial which is included in this review. However, baseline characteristics (age, gender, size of visual field deficit) were only available for the 2 trials combined ... i.e. from 38 participants, 19 of whom had optic nerve injuries and who were not relevant to this trial/review.</p> <p>2. This study included a mixed population of patients and there was no separate data for stroke. There was an uneven distribution of stroke patients between the groups, with only 2/9 in Group 1 and 8/10 in Group 2.</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Stated that participants were randomly assigned (double-blind). No further details of method of randomisation. Correspondence with the author gave the further detail that while participating in the study the participants did not know which of the treatments were expected to produce effects.
Blinding (performance bias and detection bias) All outcomes	Low risk	Stated "(double-blind)", but no further information provided in the paper. Correspondence with the author stated that it was a different person performing the tests than the person who gave the training

Kasten 1998 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	"One patient who failed to meet the requirement to training for a total of 150 hours. This patient dropped out from the study". Data for this participant appeared to be included in baseline data but did not say if included in final assessment data; the implication was no. Visual acuity reported to be performed but no data were presented.
Other bias	High risk	Researchers may have financial interest in intervention being investigated. Poggel 2004 stated "BA Sabel and E Kasten are inventors. BA Sabel is a consultant and shareholder of NovaVision Inc, and has equity in excess of \$10,000".

Kasten 2007

Methods	Restitutive intervention versus restitutive intervention
	<p>Design: RCT</p> <p>Stratification: not stated</p> <p>Randomisation sequence: "Patients were assigned randomly to one of 3 groups", method of randomisation not stated. Correspondence with author did not provide any further detail.</p> <p>Comparisons: "Three experimental groups: Group I: parallel co-stimulation (n = 7); Group II: moving co-stimulation (n = 7); and Group III: single stimulation (n = 9)".</p> <p>Allocation concealment: not stated: ("Patients were informed that the effectiveness of three different programs for visual field training was being studied and that there was no placebo group".</p> <p>Blinding: not clear: Correspondence with the author stated that it was a different person performing the tests than the person who gave the training.</p> <p>Power calculation: no</p> <p>Intention-to-treat analysis: no ("A total of 5 patients had to be excluded after baseline examinations for one or more of the following reasons: fixation deficits, noncompliance to the training procedure, other treatments that may have had influenced their training, or illness during the trial.")</p> <p>Other recruitment details: not stated</p> <p>Patient and public involvement: no</p>
Participants	<p>Total study population: 28 considered; data available for 23</p> <p>Withdrawals: "A total of 5 patients had to be excluded after baseline examinations for one or more of the following reasons: fixation deficits, noncompliance to the training procedure, other treatments that may have had influenced their training, or illness during the trial."</p> <p>Method of diagnosing VFD: high-resolution perimetry and Tubinger automated perimetry</p> <p>Characteristics of population: participant details are listed in Table 2.</p> <p>Type and severity of visual problems: participant details are listed in Table 3</p> <p>Inclusion criteria: patients aged 18 to 75 years with a stable homonymous visual field defect resulting from CNS lesion, of greater than 6 months. Visual fields had to have matching deficits of both eyes on monocular perimetry plus structural damage of post-chiasmatic visual system as documented by MRI/CT or medical reports.</p> <p>Exclusion criteria: total blindness, central scotomata, lesions of the optic nerve or chiasma, insufficient fixation, other visual diseases, visual neglect, motor disturbances, cognitive deficits, psychotic episodes, or epilepsy. Method of diagnosing visual perceptual problems: "Zahlen-Verbbinbungd test" and "alters-Konzentrationstest"</p>

Kasten 2007 (Continued)

Baseline comparison of treatment groups: 23 causes of lesions were: 14 with vascular disease, stroke, cerebral haemorrhage or ischaemia; 8 with trauma; 1 with inflammation.

Interventions

Group 1: parallel co-stimulation (n = 7)

Intervention type: restitution (2 stimuli presented simultaneously). **Materials:** training programs ran on commercially available personal computers. **Procedures:** "visual field training was completed at home after initial examinations and instruction sessions were carried out in our laboratory". **Provided by:** not stated. **Delivery:** "All three training conditions had the following features in common: While the patient had to watch a fixation control, a target stimulus (adjustable between about 0.5° and 1.5° diameter, 100 cd/m²) was presented repeatedly in a predetermined intact part of the border region of the dark screen (adjustable between < 1 cd/m² up to 50 cd/m²). Upon detection of the target stimulus, the patient was required to respond as soon as possible by pressing the spacebar of the computer keyboard. Then the stimulus moved some degrees towards the blind area. If the patient still perceived the stimulus, this had to be acknowledged by pressing the spacebar again; the stimulus then moved again some degrees into the direction of the blind area and so forth. This procedure was repeated until the patient was unable to see the stimulus in the absolute blind field. At this "blind" position the stimulus was then presented 10 times. Thereafter, stimulation started at another predetermined position of the border region (see also "transition zone", [Kasten 1998b](#)), and the stepwise stimulation process moving from intact to blind visual field regions was repeated. Standardized variables throughout training procedures in all patients were size and luminance of stimuli and background luminance." ([Table 1](#)). **Regimen:** 30 minutes twice per day for three months. **Tailoring:** yes. "the trained area was individually adjusted to the patient's visual field border". "Other variables were adjusted individually according to the visual defect — for example, the location of fixation control, the location to start stimulation, the direction of stimulation, the frequency of fixation controls, and whether feedback upon responding was accompanied by a tone. Accordingly, training parameters were adjusted to the individual pattern of residual vision."

Modification: yes "Each month (if necessary even more often), results stored on the disc were analyzed, and the training area was readjusted, based on the progress of the patients". **Adherence:** "After each session, data were saved automatically on a disk to control compliance".

Group 2: moving co-stimulation (n = 7)

Intervention: restitution (2 stimuli presented, with 1 nearer the blind area "running" continually between intact and lost field). All other intervention details as Group 1

Group 3: single stimulus (n = 9)

Intervention: Restitution (only a single stimulus was presented). All other intervention details as Group 1

Outcomes

See [Table 4](#)

Visual field: high-resolution perimetry (number of hits, learning effects, fixation ability, false hits)

Visual field: Tubinger automated perimetry at 30 and 90 degrees (no of hits, fixation ability)

Eye movements: "Chronos Vision Eye Tracker"

Visual acuity

Subjective visual ability questionnaire

Attention: "Zahlen-Verbindungs Test" of visuo-spatial attention

"Alters-Konzentrationstest" attention test for older people

"testbatterie zur Aufmerksamkeitspruefung" ability to improve attention

Notes

1. This study included a mixed population of patients and there was no separate data for stroke.

2. Visual acuity at 40 cm was measured without glasses.

Kasten 2007 (Continued)

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Quote "participants were randomly assigned to one of three experimental groups", no further information on method given. Correspondence with the author gave the further detail that participating the patients did not know which of the treatments were expected to produce effects.
Blinding (performance bias and detection bias) All outcomes	Low risk	Correspondence with the author stated that it was a different person performing the tests than the person who gave the training.
Incomplete outcome data (attrition bias) All outcomes	High risk	"A total of 5 patients had to be excluded after baseline examinations for one or more of the following reasons: fixation deficits, non-compliance to the training procedure, other treatments that may influence training, illness during the trial". No data were provided for these participants.
Other bias	High risk	Researchers may have had financial interests in intervention being investigated. Poggel 2004 stated "BA Sabel and E Kasten are inventors. BA Sabel is a consultant and shareholder of NovaVision Inc, and has equity in excess of \$10,000".

Keller 2010

Methods	<p>Compensative intervention versus compensative intervention</p> <p>Design: parallel-group RCT</p> <p>Stratification: "To achieve a nearly balanced number of patients with quadrantanopia in each group, the assignment of quadrantanopic patients between treatment groups was alternated, resulting in 3 patients in the AVT group and 4 patients in the VST group".</p> <p>Randomisation sequence: random number algorithm</p> <p>Comparisons: 2 groups, audiovisual stimulation training (AVT) versus visual stimulation training (VST)</p> <p>Allocation concealment: not stated</p> <p>Blinding: no detail provided about blinding of patients, caregivers or assessors</p> <p>Power calculation: not stated</p> <p>Intention-to-treat analysis: not stated</p> <p>Other recruitment details: not stated</p> <p>Patient and public involvement: not stated</p>
Participants	<p>Total study population: 20 with unilateral lesions of the occipital lobe confirmed by MRI. Not all participants had VFD as a result of stroke. 18/20 stroke; 1/20 tumour and 1/20 TBI</p> <p>Withdrawals: none</p> <p>Method of diagnosing VFD: Goldmann perimeter examination. Visual field assessment was done using the kinetic method with suprathreshold checks. Measured before and after training. "Pre-post comparisons of visual field defects were classified as stable in all patients".</p> <p>Characteristics of population: participant details are listed in Table 2.</p>

Keller 2010 (Continued)

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: "left- or right-sided visual field deficits primarily caused by stroke" (as confirmed by MRI). Normal bilateral hearing measured by audiometry with no asymmetry between ears, normal or corrected binocular visual acuity.

Exclusion criteria: visual neglect or signs of aphasia ("Patients were initially tested with 3 different neglect tests (line bisection, Mesulam test, draw a clock face test) and the token test")

Baseline comparison of treatment groups: "Both groups were comparable with regard to demographic and clinical variables before treatment".

Interventions

Group 1: audiovisual exploration training (AVT) (n = 10)

Intervention type: compensation. **Materials:** red light-emitting diodes (LEDs), piezoelectric loudspeakers, white-noise generator, laptop computer using a custom-made software for this purpose. **Where can materials be accessed?** not stated. **Procedures:** "training sessions were conducted in a dimly illuminated room. The patients sat in chairs 60 cm in front of the apparatus, with their eyes adjusted to the center of the apparatus. Stimuli to the blind visual field and intact visual field were presented in random sequence. Patients were instructed to detect the presence of visual targets by pressing a response button as fast as possible. To prevent patients from reacting to false positives, 20% catch trials with solely acoustic stimulation were implemented in each training session. Whenever patients responded to a catch trial, a computer-generated 'nickering' of a horse sounded. Patients were explicitly instructed to execute eye movements with their head held straight forward. Training was carried out under binocular conditions". **Provided by:** not stated. **Delivery:** face-to-face, individual, location inpatient ([Table 1](#)). **Regimen:** 20 therapy sessions (each session lasting 30 minutes) over 3 weeks. **Tailoring:** yes. Based on type of VFD ("For patients with homonymous hemianopia, 70% of all stimuli were presented in the blind visual field. For patients with quadrantanopia, 70% of the stimuli were presented in the 2 upper or 2 lower rows of the affected side"). **Modification:** yes, "whenever a patient complained of tiredness, the training was interrupted for 1 minute". **Adherence:** not stated

Group 2: visual exploration training (VET) (n = 10)

Intervention type: compensation. **Materials:** as described for AVT. **Where can materials be accessed?** not stated. **Procedures:** "settings of stimuli presentation were identical as for the AVT, with the exception that the sound was turned off during all training sessions and catch trials were not needed". **Provided by:** not stated. **Delivery:** face-to-face, individual, location inpatient ([Table 1](#)). **Regimen:** 20 therapy sessions (each session lasting 30 minutes) over 3 weeks. **Tailoring:** yes. Based on type of VFD ("For patients with homonymous hemianopia, 70% of all stimuli were presented in the blind visual field. For patients with quadrantanopia, 70% of the stimuli were presented in the 2 upper or 2 lower rows of the affected side"). **Modification:** yes, "whenever a patient complained of tiredness, the training was interrupted for 1 minute". **Adherence:** not stated

Outcomes

See [Table 4](#)

Visual field: Perimetry - Goldmann kinetic and suprathreshold

Visual exploration test

Reading test

Search task

Evaluation of ADL

Electro-oculography

Time points when outcomes were assessed: before and after treatment

Definitions relevant to outcomes: "Changes in visual fields defects were established by comparing the pretraining visual field with the post-training visual field. Changes were classified as better, stable, or worse. Better or worse were defined by differences of more than 5° horizontally or vertically".

Keller 2010 (Continued)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	High risk	Random number algorithm used to assign participants. However, it was also stated that "To achieve a nearly balanced number of patients with quadrantanopia in each group, the assignment of quadrantanopic patients between treatment groups was alternated, resulting in 3 patients in the AVT group and 4 patients in the VT group". This suggests that allocation was not concealed for these participants.
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No information was given on blinding of participants, or outcome assessors. It is unlikely that personnel could be blinded.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No mention of any dropouts
Other bias	Low risk	No other concerns noted

Modden 2012

Methods	<p>Restitutive intervention versus compensative intervention</p> <p>Design: prospective randomised controlled, single-blind, single-center treatment study</p> <p>Stratification: no</p> <p>Randomisation sequence: "Patients were randomly assigned to receive either CT, RT, or OT. Randomization by throwing dice and allocation took place before starting with the initial assessment of neuropsychological tests".</p> <p>Comparisons: compensatory therapy (CT) versus restorative computerised training (RT) versus standard occupational therapy (OT)</p> <p>Allocation concealment: no</p> <p>Blinding: no ("All patients were recruited and assigned to treatment groups by a neuropsychologist. The same neuropsychologist also tested the patients before (time point T1) and after (time point T2) the treatment and was not blinded to the type of training. The training itself was performed by a psychological assistant or by the occupational therapists not involved in the study, and they provided the test results at T1 and T2.")</p> <p>Power calculation: yes</p> <p>Intention-to-treat analysis: no dropouts</p> <p>Patient and public involvement: not stated</p>
Participants	<p>Total study population: 45 patients randomised</p> <p>Withdrawals: 0</p> <p>Method of diagnosing VFD: "a perimetry test from the Test Battery of Attentional Performance, the latter having a sensitivity and specificity for visual field defects similar to the Goldmann perimetry."</p>

Modden 2012 (Continued)

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: homonymous hemianopia with a posterior cerebral artery stroke

Exclusion criteria: visual neglect, eye-movement disorders, neuropsychological disorders like aphasia, dysexecutive syndromes, memory deficits, or higher order motor impairments like apraxia

Baseline comparison of treatment groups: "At baseline, the 3 groups did not differ in demographic and neuropsychological measures".

Interventions

Group 1: restitution therapy (RT) (n = 15)

Intervention: RT - computer-based stimulation of visual field

Intervention type: restitutive. **Materials:** computer-based restitution therapy: "A therapy-integrated perimeter program (provided by Teltra company)". **Procedures:** "A therapy-integrated perimeter program (provided by Teltra company) created the exact measurement of the individual visual field border. Using that measurement, a series of colored targets appeared on a blue screen anywhere at 1 of 10 positions on the border line. A randomly presented first fixation target (a rotating arrow) announced the second stimulus target in the hemianopic border zone (basic principle of covert attention shift). The patients were instructed to respond (by pressing a key) to each stimulus target (colored and flickering frames, beams, and spots) as soon it was perceived. The program contained no adaptive difficulty levels. Eye movements were not allowed, and this was controlled by the assistant". **Provided by:** "The training itself was performed by a psychological assistant or by the occupational therapists not involved in the study". **Delivery:** face-to-face, individual, location rehabilitation centre: "The participants in both PC-based therapy groups were seated 60 cm away from the screen (19-inch monitor) and had to perform the tasks binocularly. As during testing, the head was fixed by a chin rest, the sessions were always controlled by the assistant to make sure that the instructions were followed." ([Table 1](#)). **Regimen:** 30 minutes/day for 15 sessions. **Tailoring:** no: "The program contained no adaptive difficulty levels." **Modification:** no. **Adherence:** stated "no-one dropped out because of problems with compliance".

Group 2: compensatory therapy (CT) (n = 15)

Intervention: CT - computer-based stimulation of visual field

Intervention type: compensatory. **Materials:** computer-based: "The 'Exploration' task (from RehaCom, provided by HASOMED GmbH, Magdeburg, Germany)". **Procedures:** "The 'Exploration' task was adapted individually according to the side of the hemianopia. On a dark background, different bright stimuli arranged in rows and columns were presented. A ring (diameter of 2 cm) moved line by line (interlaced) on a matrix unit over the field. The participant was instructed to follow the ring (starting point to an outmost fixation in the blind side) by eye movements and to identify a critical targeted icon. The targets were not always distributed homogeneously but were clustered in the blind side. Thus, the exploration in the hemianopic field was further promoted. The patients had to respond (by pressing a key) when the targeted icon was perceived in the circle. **Provided by:** "The training itself was performed by a psychological assistant or by the occupational therapists not involved in the study". **Delivery:** face-to-face, individual, location rehabilitation centre: "The participants in both PC-based therapy groups were seated 60 cm away from the screen (19-inch monitor) and had to perform the tasks binocularly. As during testing, the head was fixed by a chin rest. The sessions were always controlled by the assistant to make sure that the instructions were followed." ([Table 1](#)). **Regimen:** 30 minutes/day for 15 sessions. **Tailoring:** yes: "The program contained several difficulty levels. In levels 1 to 20, all lines were completely filled with symbols, whereas there were omissions in the rows of symbols in levels 21 to 30 to increase the difficulty." **Modification:** not clear. **Adherence:** stated "no-one dropped out because of problems with compliance".

Group 3. occupation therapy (OT) (n = 15)

Intervention: OT - standard occupational therapy

Intervention type: compensatory. **Procedures:** "After a standardized assessment of daily living activities, the therapy consisted of individually adapted stimulation of daily activity tasks to compensate via eye-, head-, and body movements. These compensation strategies included aspects of spatial and

Modden 2012 (Continued)

body perception, searching or arranging objects, pen and paper searching task, reading maps or newspapers, and self-care activities. The participant was instructed to perform systematic eye movements toward the lost visual field. The interventions were carried out in the treatment rooms, on the wards, in a kitchen or a bathroom, outside in the park, or in a supermarket." **Provided by:** occupational therapist. **Delivery:** face-to-face, individual, location rehabilitation centre: (Table 1). **Regimen:** 30 minutes/day for 15 sessions. **Tailoring:** yes: "After a standardized assessment of daily living activities, the therapy consisted of individually adapted stimulation of daily activity tasks". **Modification:** not clear. **Adherence:** stated "no-one dropped out because of problems with compliance".

Note: "Patients receiving RT and CT did not receive OT in the context of their standard rehabilitation treatment."

Outcomes	See Table 4 Visual field expansion - test battery of attentional performance visual field assessment Visual search performance - cancellation Reading performance - Weschler memory tests ADL - Extended Barthel Index Time points when outcomes were assessed: after completion of training
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Notes

Risk of bias		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	High risk	Randomisation was by throwing a dice: no allocation concealment.
Blinding (performance bias and detection bias) All outcomes	High risk	No blinding: "All patients were recruited and assigned to treatment groups by a neuropsychologist. The same neuropsychologist also tested the patients before (time point t1) and after (time point t2) the treatment and was not blinded to the type of training. The training itself was performed by a psychological assistant or by the occupational therapists not involved in the study, and they provided the test results at t1 and t2."
Incomplete outcome data (attrition bias) All outcomes	Low risk	No dropouts
Other bias	Low risk	No other concerns noted

Plow 2010

Methods	Restitutive intervention versus restitutive intervention Design: "Randomised controlled, double-blind pilot trial" Stratification: no Randomisation sequence: "Participants were randomly assigned using a predetermined enrollment sequence to 1 of 2 arms". Method of randomisation not stated. Correspondence with the author provided the information that "as subjects were admitted, they were randomly assigned to one of two groups based on an a priori generated randomization strategy."
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Plow 2010 (Continued)

Comparisons: 2 groups: vision restoration therapy with active transcranial direct current stimulation (VRT + tDCS) and vision restoration therapy with sham transcranial direct current stimulation (VRT + sham)

Allocation concealment: yes: "Participants and investigators analyzing visual field outcomes were blinded to the tDCS mode (active versus sham)."

Blinding: yes: "Participants and investigators analyzing visual field outcomes were blinded to the tDCS mode (active versus sham)."

Power calculation: not stated

Intention-to-treat analysis: no

Other recruitment details: "A total of 150 potential participants were screened. Following comprehensive neurological and ophthalmological screening, 12 patients ... were enrolled".

Patient and public involvement: not stated

Participants

Total study population: 12

Withdrawals: 4

Method of diagnosing VFD: high-resolution perimetry

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: "Patients with unilateral postchiasmal visual field loss following stroke or brain damage, who were in the chronic phase of recovery (> 3 months post-lesion)". Participants were "aged over 18, with deep hemianopic field loss as defined and confirmed by monocular perimetry along with established structural damage of the post-charismatic visual system as documented by standard neuroimaging techniques (CT or MRI), medical reports, or a combination of these. Participants also had cognitive, language and motor function sufficient to understand the experiments and follow instructions, had given informed written consent to participate in the study and had motivation to participate in the VRT program."

Exclusion criteria: "Any ocular visual pathology or contraindication to noninvasive brain stimulation and tDCS. Specific criteria drawn from safety guidelines pertaining to the use of noninvasive cortical stimulation include 1) the presence of any metallic, mechanical, or magnetic implant in the head or implantable device (e.g. cardiac pacemaker); 2) prior history of seizure or familial history of seizure disorder in a first degree relative, and 3) chronic use of neuroactive medication (e.g. neurostimulants, anti-convulsants, or antidepressants)."

Baseline comparison of treatment groups: not stated. Baseline data were provided, but no statistical comparison. Time since stroke did not appear to be comparable, with the VRT + tDCS group mean, 20.8 months (SD 26.6, range 3 to 72 months), and VRT + sham group mean, 58.7 months (SD 72.9, range 10 to 192 months).

This study appeared to include participants with visual field defects only (no visual neglect), although the method of ensuring no visual neglect was unclear

Method of diagnosing visual field defect: "monocular perimetry", no further details of method

Method of diagnosing visual perceptual problems: not stated

Interventions

Group 1: visual restoration therapy + transcranial Direct Current Stimulation (VRT + tDCS) (n = 6; data available for n = 4)

Intervention: computer-based stimulation of the visual field with active transcranial direct current stimulation, 30-minute sessions twice per day, 3 times per week for 3 months of vision restoration therapy and concurrent transcranial direct current stimulation of 2 mA/minute

Plow 2010 (Continued)

Intervention type: restorative. **Materials:** computer-based restitution therapy: "Vision Restoration Therapy (VRT; Novavision Inc, Boca Raton, Florida)", transcranial Direct Current Stimulation: "tDCS was applied using two 5 × 7 cm² saline-soaked sponge electrodes connected to a 9-V battery-driven stimulator (IOMED Inc, Salt Lake City, Utah)". **Procedures:** vision restoration therapy: "Briefly, participants were seated in front of a computer screen at a constant viewing distance and instructed to detect (signaled by a key press) the presence of a flashed light stimulus while maintaining fixation on a central target. Built-in fixation monitoring required patients to respond to a color change of the central fixation target occurring at random intervals. Target stimuli were presented primarily in the region of the transition zone (identified by a prior visual field test; see details on high-resolution perimetry), and the spatial parameters of customized therapy were determined based on weekly progress and results of monthly tests." Transcranial direct current stimulation: "tDCS was applied using two 5 × 7 cm² saline-soaked sponge electrodes connected to a 9-V battery-driven stimulator delivering a constant current of 2 mA for the entire duration of the training procedure. Following the 10-20 International EEG co-ordinate system, the anode was placed at the occipital pole and the cathode (reference) was positioned at the vertex. Electrodes were then secured using nonlatex rubber straps, and an identical montage was worn by all patients throughout training." **Provided by:** University eye clinic. **Delivery:** face-to-face, individual, location University eye clinic (Table 1). **Regimen:** "We used a contracted VRT regimen lasting 3 months (2 half-hour sessions, separated by a 30-minute rest interval, for 3 d/wk)." **Tailoring:** not stated. **Modification:** not stated. **Adherence:** not stated

Group 2: visual restoration therapy + sham transcranial Direct Current Stimulation (VRT + sham) (n = 6; data available for n = 4)

Intervention: computer-based stimulation of the visual field with sham transcranial direct current stimulation, 30-minute sessions twice per day, 3 times per week for 3 months of vision restoration therapy, transcranial direct current stimulation turned on for 30 seconds then ramped down to zero and turned off. **Intervention type:** restorative. **Materials:** see above. "Experimental blinding with respect to active or sham transcranial direct current stimulation was implemented according to standard protocol guidelines described previously". **Procedures:** see above. **Provided by:** see above. **Delivery:** see above (Table 1). **Regimen:** see above. **Tailoring:** not stated. **Modification:** not stated. **Adherence:** not stated

Outcomes

See Table 4

Visual field - high-resolution perimetry: position of visual field border and stimulus detection accuracy

Visual field - subjective topographic measure of perceived visual field deficit

Extended ADL - Veterans Affairs Low Vision-Visual Functional Questionnaire (LV-VFQ)

QoL - Vision Impairment (IVI) profile

Other - independent measure of fixation performance during training and HRP testing

Time points when outcomes were assessed: baseline and at completion of training (3 months)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	"Participants were randomly assigned using a predetermined enrollment sequence to 1 of 2 arms". Method of randomisation not stated. Correspondence with the author provided the information that "as subjects were admitted, they were randomly assigned to one of two groups based on an a priori generated randomization strategy."
Blinding (performance bias and detection bias) All outcomes	Low risk	Correspondence with the author stated "the investigators assessing the visual field results (primary outcome) were also blinded to patient group assign-

Plow 2010 (Continued)

ment. Once visual field assessment was complete, they were provided with an encrypted copy of the data without any identifying information".

Correspondence with the author stated "Blinding to stimulation (i.e. sham) was maintained by exploiting the inherent properties of tDCS. All patients wore the electrode montage regardless of the group they were relegated to. When the tDCS unit is turned on, current is slowly ramped up until the target current level is reached. During this time, patients will typically report a tingling or itching sensation beneath the surface of the anode electrode (overlying the occipital cortex). This sensation subsides shortly after a couple of minutes of habituation. Thus, in the experimental group, the current remains on but in the sham control group, the current is turned off. In either case, the patient is not aware of this happening. This fact is exploited for the purposes of experimental blinding since patients in the VRT + tDCS group can not perceive sensation of stimulation and neither can those patients who are in the VRT + sham tDCS group."

Incomplete outcome data (attrition bias) All outcomes	Low risk	There were 4 dropouts who were not included in data analysis. Reasons for dropouts were provided, and were even between the groups. The author supplied the following information: "Out of 12 patients who were enrolled into the study, 4 (i.e. 2 from each group), were excluded or could not complete the study. Here are the reasons: VRT + tDCS group: Patient 1 - excluded soon after randomization as she had an unrelated adverse event that excluded her from participation; Patient 2 - excluded from analysis due to technical issues that could not allow us to ascertain whether sufficient tDCS current was being delivered throughout the training period. VRT + sham tDCS group: Patient 1: did not receive allocated intervention due to onset of medication use that was contraindicated with tDCS; Patient 2 - discontinued due to onset of a medical condition that precluded her from further participation."
Other bias	Low risk	No other concerns noted

Poggel 2004

Methods	Restitutive intervention versus restitutive intervention
	Design: prospective RCT
	Stratification: no
	Randomisation sequence: stated participants were "assigned randomly" but no details given. Correspondence with author did provide any further detail.
	Comparisons: 2 groups - visual restoration therapy (VRT) with attentional cueing, VRT without attentional cueing
	Allocation concealment: no - "Patients were not told which type of training they received; a double-blind control was not possible owing to staff limitations."
	Blinding: no
	Power calculation: no
	Intention-to-treat analysis: no. 1 dropout, for whom no data was presented. Stated: "Patient 5 did not participate in the training procedure owing to time constraints and is not listed here."
	Other recruitment details: not stated
	Patient and public involvement: not stated

Poggel 2004 (Continued)

Participants

Total study population: 20 recruited (data available for 19)

Withdrawals: 1 ("Patient 5 did not participate in the training procedure owing to time constraints and is not listed here.")

Method of diagnosing VFD: high-resolution perimetry and standard Tübingen Automated Perimetry

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: homonymous visual field defects after postgeniculate lesions, age 18 to 75 years. Visual field size, assessed by high-resolution campimetry, had to be stable; that is, patients were included only if visual field size increased or decreased by < 2% over at least 4 weeks before the study started".

Exclusion criteria: optic nerve, retinal or other ophthalmic disorders; mental deficits; impaired attention and neglect; psychiatric disorders; epilepsy and diseases with obvious visual or cognitive effect

Baseline comparison of treatment groups: not stated

This study included patients with visual field defects only (no visual neglect).

Method of diagnosing visual perceptual problems: not stated

Interventions

Group 1: visual restitution therapy (VRT) with attentional cueing (n = 10)

Intervention: computer-based stimulation of visual field, with cueing

Intervention type: restitution

Materials: computer-based vision restoration therapy (Nova Vision, Magdeburg, Germany). **Procedures:** a high-resolution computer-based campimetric test (high-resolution perimetry [HRP]; Nova Vision, Magdeburg, Germany) was used to assess visual field size and to determine ARVs. Training stimuli appeared on a dark computer screen, each target increasing in brightness in four steps from dark gray (30 cd/m²) to bright white (96 cd/m²) over 2000 milliseconds. Stimulus size, fixation control, mode of response, and viewing distance were identical to those used for HRP. For vision restoration therapy with attentional cueing, the training stimulus was preceded by a large (12° x 12° visual angle) dim gray cue frame enclosing a predetermined segment in the upper visual field that included parts of the intact and blind fields as well as ARVs. The frame appeared for 200 milliseconds and was followed at a randomised interval (mean stimulus onset asynchrony: 1000 milliseconds, range 750 to 1250 milliseconds) by a training stimulus presented in the area that had been stimulated by the cue frame (the attention field). In each training session, patients received 500 training stimuli. In the experimental group, approximately one-third of the trials, that is, those where the target was to be presented in the attention field, were preceded by a cue. Targets were presented at random locations within the upper and the lower visual fields. Depending on the percentage of stimuli detected, the duration of each training session was approximately 30 to 35 minutes." **Provided by:** not stated. **Delivery:** training completed at home using computer ([Table 1](#)). **Regimen:** "Patients performed the training in six training units, each lasting approximately 1 month. A unit consisted of 56 sessions, with two sessions per day." **Tailoring:** yes: "Each patient's ARV was determined based on five HRP tests. The training area was then adjusted to the visual field border of that individual." **Modification:** not stated. **Adherence:** not stated ("Data from each session were saved on a disk")

Group 2: visual restitution therapy (VRT) with no attentional cueing (n = 9)

Intervention: computer-based stimulation of visual field

Intervention type: restitution

Materials: see above. **Procedures:** see above. **Provided by:** see above. **Delivery:** see above. ([Table 1](#)). **Regimen:** see above. **Tailoring:** see above. **Modification:** see above. **Adherence:** see above

Outcomes

See [Table 4](#)

Poggel 2004 (Continued)

Change in size of visual field:

Tubinger automated perimetry (TAP)

High resolution perimetry (HRP)

Time points when outcomes were assessed: at the end of each training unit (HRP), and after the 6th training unit (HRP and TAP)

Notes

Training and measurement were essentially the same thing.

 Authors referred to [Kasten 2001](#) as proof that control was not needed. They did not address his conclusions on effects of training when recruiting from an already trained group - see [Jobke 2009](#) above.

Outcome data was largely presented for all patients combined (control and experimental groups combined)

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	"Patients were not told which type of training they received; a double-blind control was not possible owing to staff limitations" - the paper did not state what methods (if any) were used to conceal the allocation from those enrolling and assigning participants and correspondence with the author did not provide any further information.
Blinding (performance bias and detection bias) All outcomes	High risk	"Patients were not told which type of training they received; a double-blind control was not possible owing to staff limitations".
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	1 participant did not participate, none of his data were included in final analysis. The co-author was not totally clear on the reason for non-participation, but believed it to be another stroke.
Other bias	High risk	Researchers had financial interest in intervention being investigated: "BA Sabel and E Kasten are inventors. BA Sabel is a consultant and shareholder of NovaVision Inc, and has equity in excess of USD 10,000".

Rossi 1990
Methods
Substitutive intervention versus control, placebo or no intervention
Design: RCT

Stratification: no

Randomisation sequence: "randomly assigned". Method of randomisation not stated

Comparisons: 2 groups; 15 diopter Fresnel prisms and control (no prisms)

Allocation concealment: no

Blinding: no, as the prism group wore their prisms during testing

Power calculation: no

Intention-to-treat analysis: not stated (not clear if any dropouts)

Rossi 1990 (Continued)

Other recruitment details: "39 patients with either homonymous heminaopia or unilateral visual neglect were recruited from an inpatient stroke rehabilitation unit".

Patient and public involvement: not stated

Participants

Total study population: 39

Withdrawals: none

Method of diagnosing VFD: tangent screen (inability to reliably detect a 1 cm red target on tangent screen examination in a homonymous field pattern) and Harrington Flocks visual field screener

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: homonymous hemianopia or unilateral visual neglect and a diagnosis of stroke based on clinical history, neurological exam and neuroimaging

Exclusion criteria: visual acuity worse than 20/200, unable to comprehend and co-operate with visual field assessment, disabling cardiac, pulmonary or rheumatologic problems

Baseline comparison of treatment groups: yes

The study included only stroke patients.

This study included patients with visual field defects and/or visual neglect.

Method of diagnosing visual perceptual problems: neglect was confirmed by inability to reliably detect bilateral tachistoscopically presented targets using the Harrington Flocks Visual Screener.

Interventions

Group 1: 15 diopter Fresnel prisms (n = 18)

Intervention: prisms

Intervention type: substitution

Materials: plastic, press-on Fresnel prisms (stated these are "commercially available"). **Procedures:** "Plastic press-on Fresnel prisms were trimmed to fit on the inside of the patients's spectacle lenses. Each prism was cut to the shape of a half circle overlaying only the affected hemifield, with the base of the prisms toward the affected field". **Provided by:** not stated (university neurology department), **Delivery:** face-to-face, individual, location inpatient ([Table 1](#)). **Regimen:** "Patients wore the prisms for all daytime activities. Prism position and cleanliness were checked daily by study personnel. Patients were restricted to ambulation 'with assistance only' for at least 1 day after the fitting of the prisms". Also "participated in a routine stroke rehabilitation program including physical, occupational, and speech therapy ... ADL training and table-top visual perception retraining tasks". **Tailoring:** no. **Modification:** no. **Adherence:** not stated (stated "In practice, most patients tolerated the Fresnel prisms well after a day or two")

Group 2: control (n = 21)

Intervention: no prisms

Intervention type: control

Materials: NA. **Procedures:** NA. **Provided by:** NA. **Delivery:** NA. **Regimen:** "participated in a routine stroke rehabilitation program including physical, occupational, and speech therapy ... ADL training and table-top visual perception retraining tasks". **Tailoring:** NA. **Modification:** NA. **Adherence:** NA

Outcomes

See [Table 4](#)

Modified Mini Mental Status Examination

Motor free visual perception test

Rossi 1990 (Continued)

Line bisection task
 Line cancellation task
 Harrington Flocks Visual Screener
 Tangent Screen Examination
 Barthel ADL score
 Frequency of falls

Time points when outcomes were assessed: baseline, 2 weeks and 4 weeks

Notes

1. The length of time that the prisms were worn may vary considerably for different patients, as this was for "all day time activities"
2. States that diagnosis of homonymous hemianopia precludes diagnosis of neglect - this is not current thinking so must question diagnostic criteria especially with respect to neglect

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Patients were "randomly assigned" but methodology not stated.
Blinding (performance bias and detection bias) All outcomes	High risk	"the study was not performed in a blinded fashion", however for Tangent Screen Examination "results were plotted onto templates and judged by observer unaware of the patient's group assignment ..."
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No incomplete data mentioned
Other bias	Low risk	No other sources of bias noted

Roth 2009

Methods

Compensative intervention versus restitutive intervention

Design: RCT

Stratification: no

Randomisation sequence: paper stated "an equal number of patients assigned randomly to either the EST or the FT group"; and correspondence with the author stated that "the patients were randomized according to a pre-existing list".

Comparisons: 2 groups; compensatory "exploratory saccadic training (EST)" and control ("flicker-stimulation training (FT), which is unlikely to affect visual-search behaviour")

Allocation concealment: no

Blinding: not stated

Power calculation: no

Intention-to-treat analysis: no

Roth 2009 (Continued)

Other recruitment details: participants, aged 18 to 80 years, with post-chiasmatic lesions of greater than 6 months duration and visual field defects that are isolated homonymous hemianopias or quadrantanopias. The study included a mixed population (mainly stroke).

Patient and public involvement: not stated

Participants

Total study population: 30 participants recruited (data analysed for 28)

Withdrawals: "Two FT patients dropped out because of illness and insufficient compliance".

Method of diagnosing VFD: not stated.

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: visual field defects must not cross the vertical midline, but reach within 5 degrees of it and there must be no other visual field defects, or visual acuity deficit.

Exclusion criteria: other visual field defects, visual acuity less than 0.6 (20/33), other diseases of eye or brain, motor impairments hampering computer use, other neurologic impairments (particularly epilepsy or hemineglect)

Baseline comparison of treatment groups: "The groups did not differ regarding age, diagnoses or duration of disease".

This study included a mixed population (mainly stroke). Group 1: 15 participants (12 stroke or haemorrhage, 3 other), Group 2: 15 participants (14 stroke or haemorrhage, 1 other)

This study included patients with visual field defects only (no visual neglect).

Method of diagnosing visual perceptual problems: clock drawing and line bisection test

Interventions

Group 1: explorative scanning training (n = 15)

Intervention: computer-based scanning training

Intervention type: compensation

Materials: laptop. **Where can materials be accessed?** "A custom software program (Borland Delphi 7.0) was used to generate a random array of digits....". **Procedures:** random digits were presented with equal distribution in blind and seeing fields: patients had to find these digits with positive feedback provided for a correctly found number. **Provided by:** not stated. **Delivery:** face-to-face, individual, location home-based training ([Table 1](#)). **Regimen:** training was performed for 30 minutes, twice per day, 5 days per week for 6 weeks at home, on a laptop set at 30 cm. **Tailoring:** no. **Modification:** no. **Adherence:** protocols were used to attempt to ensure standardisation and patients were instructed to avoid head movements. However, it is not known whether this standardisation and instructions were adhered to. There may, therefore, be some inconsistencies in the delivery of the intervention. Errors in aspects such as laptop position, etc. could impact on the position of the stimuli viewed and subsequently the explorative eye movements.

Group 2: flicker-stimulation training (n = 15)

Intervention: computer-based training, "designed to stimulate the blind hemifield"

Intervention type: restitution (note - author stated this is 'potential' restitution training)

Materials: laptop. **Where can materials be accessed?** "Borland Delphi 7.0 was used to generate the flicker stimulus". **Procedures:** letters flickering at 10 Hz were presented at 21.8 degrees eccentricity, with proportion 3:1 blind:seeing field. Participant to maintain central fixation but identify peripheral letter: positive feedback given of correctly identified letter. **Provided by:** not stated. **Delivery:** "The patients trained at home, using our laboratory's laptops to ensure standard training conditions (screens, fixed viewing distance, and visual-field area trained)" ([Table 1](#)). **Regimen:** training was performed for 30 minutes, twice per day, 5 days per week for 6 weeks at home, on a laptop set at 30 cm. **Tailoring:**

Roth 2009 (Continued)

no. **Modification:** no. **Adherence:** patients were instructed to fixate. There appears to have been no attempt to control or measure fixation.

Outcomes	See Table 4 Digit search task (response time) Natural search task (response time) Natural scene exploration and fixation stability (video eye tracker) Perimetry (Tubinger automated perimeter) Reading speed QoL - World Health Organisation questionnaire WHOQOL-BREF Time points when outcomes were assessed: baseline, post-treatment, follow-up
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Notes	Inclusion criteria stated visual field defect with "duration exceeding 6 months". However, 1 patient in group 2 was reported to have a 4-month-old homonymous hemianopia. Possibility of spontaneous recovery. Furthermore, this discrepancy casts doubt on the application of the selection criteria.
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Risk of bias		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Correspondence with the author stated that "the patients were informed that probably each method could be effective, one by eye movement training (EST), the other by visual field stimulation (FT). FT was supposed to be a potential restitution training. Therefore, the patients of both groups performed a potentially effective training - but with quite different approaches".
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No details in the paper or by contacting the author of whether or how assessor was blinded
Incomplete outcome data (attrition bias) All outcomes	High risk	Stated "two FT (Group 2) patients dropped out because of illness and insufficient compliance. Finally data from 15 EST and 13 FT patients were included". Data from these participants were included in baseline characteristics, but not included in final analyses.
Other bias	Low risk	No other cause of bias noted

Rowe 2010

Methods	<p>Substitutive intervention versus compensative intervention versus control, placebo or no intervention</p> <p>Design: "Randomized controlled, multicentre pilot trial"</p> <p>Stratification: "Randomization lists were generated using block randomization stratified by centre and degree of hemianopia (partial or complete) with treatment allocation ratio of 1:1:1."</p> <p>Randomisation sequence: "Participants were individually randomized to one of three treatment groups using a secure (24-hour) web-based randomization programme."</p> <p>Comparisons: 3 groups: Group 1 - Fresnel prisms; Group 2 - visual search training; Group 3 - control (standard care/information only)</p>
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Rowe 2010 (Continued)

Allocation concealment: treatment allocation was disclosed to the patient by the treating clinician: "The local PI (orthoptist) obtained the treatment allocation and subsequently assigned the participant to the treatment arm."

Blinding: outcome assessors for visual field assessment and reading speed were blind to treatment allocation. "due to the nature of the intervention, it is not possible to blind other study personnel or the participant."

Power calculation: no - pilot trial; designed to enable sample size calculation for future trials

Intention-to-treat analysis: "Outcome data were analysed according to the intention-to-treat principle."

Other recruitment details: "Participants were recruited from stroke units based in 15 United Kingdom (UK) National Health Service (NHS) trusts. Potentially eligible participants were identified by stroke research nurses, and screened for inclusion by a local principal investigator (a qualified orthoptist registered with the Health and Care Professions Council, UK). Participants eligible for inclusion, and providing consent, attended for a baseline assessment, which included assessment and documentation of patient demographics, visual signs and symptoms, visual acuity measures, any additional ocular problems, comorbidity, severity of stroke and level of disability".

Patient and public involvement: protocol stated: "This trial has involved a stroke survivor directly in the development of this protocol (JR) and will liaise closely with her for advice and direction throughout the conduct of the trial and in the dissemination process. Involvement of stroke survivors in oversight committees is also planned for this trial".

Participants

Total study population: 87 participants randomised. 71 participants at 26-week follow-up

Withdrawals: 2 "complete withdrawal" ("patients withdrawn from all data analysis and follow-up"). 9 "partial withdrawal" ("patients withdrawn from follow-up"). 5 loss to follow-up

Method of diagnosing VFD: "Stable homonymous hemianopia (partial or complete) induced by recent stroke, defined following WHO guidelines". "The visual field assessment will be conducted by a qualified Orthoptist at baseline and at the 6-week, 12-week and 26-week follow-up visits. An Esterman strategy is to be used for quantitative visual field assessment. This can be performed using either: The Esterman programme on Humphrey or Octopus perimetry; The III4e target on Goldmann with additional checks of static points in the central visual field."

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: "a. 18 years of age or older; b. best corrected visual acuity of 0.5 or better in each eye at distance; c. stable homonymous hemianopia (partial or complete) induced by recent stroke, defined following WHO guidelines, present over 2 weeks (to exclude rapid recovery cases) but less than 26 weeks prior to randomization; d. refractive error within ± 5 dioptres; e. willing and able to give consent for the study; f. prior to stroke able to read and understand English."

Exclusion criteria: "a. unable to consent due to severe cognitive impairment; b. assessed to have ocular motility impairment and/or visual inattention in addition to the visual field impairment; or c. had pre-existent visual field impairment due to previous stroke."

Baseline comparison of treatment groups: "There were no notable differences at baseline between three arms."

This study included patients with visual field defects only (no visual neglect).

Method of diagnosing visual perceptual problems: "as assessed by the orthoptist"

Interventions

Group 1: prisms (n = 27)

Intervention: Fresnel prisms (40 prism dioptre strength)

Intervention type: substitution

Rowe 2010 (Continued)

Materials: "Participants were assessed and given sector Fresnel prisms of 40 prism dioptre strength on their glasses (or plain glasses if not already worn). Separate prism segments were used as a mechanical displacement to expand the upper and lower quadrants." **Where can materials be accessed?** In the UK: NHS supplies for Fresnel prisms. **Procedures:** "The participant will be instructed to maintain central fixation through their glasses. They will then be instructed to use head movements to explore their field to the affected side when they become aware of an object of interest through the prism. The first prism will be placed at the participant's first visit; if possible the second prism will also be fitted at this time. However, if the participant is not comfortable with both prisms being fitted at once, the second prism can be placed at a second visit (2 weeks later, \pm 1 week) if no adaptation difficulties to the first prism have occurred. If adaptation difficulties have occurred the patient can continue with only the first prism and this will be captured on the case report forms." **Provided by:** orthoptist. **Delivery:** face-to-face, clinic location (patients could be in or outpatients). (Table 1) **Regimen:** "The prisms should be worn for a minimum of 2 h daily from prism affixation until 6-week follow-up visit as a minimum, after this the patient can elect to continue treatment if they wish." **Tailoring:** no. **Modification:** no. **Adherence:** "There were 73 protocol deviations in 58 patients (68.2% overall: 77% in the Fresnel prisms arm, 93% in the visual search arm and 34.5% in the standard care arm). The majority of deviations ($n = 41$, 56.2%) related to lack of compliance in the intervention arms (e.g. prism not worn a minimum of 2 hours daily for 6 weeks or visual exercises not carried out for 30 minutes daily for 6 weeks). Compliance level was similar across the intervention arms. Patients in the Fresnel prisms arm wore the prisms during 27 days on average."

Group 2. scanning training ($n = 30$)

Intervention: "visual search training"

Intervention type: compensatory (scanning) training

Materials: "Comprised an A4 landscape card with horizontal and diagonal numbered circles radiating out from a central fixation target." **Where can materials be accessed?** From author. **Procedures:** "The participant will be instructed to hold this at a distance of 8 inches from their eyes (to ensure a wide field of vision is utilised), glasses can be worn as required. Participants will be asked to transfer gaze quickly between printed targets on the A4 card. The targets are printed off centre to the right and left sides along the horizontal as well as oblique planes to ensure stimulation of a wide area in the blind and seeing parts of the visual field." **Provided by:** instructions provided by orthoptist. Training carried out at home by patients. "Participants will be instructed on the scanning exercises following randomisation to ensure their understanding of the procedure of doing this training. In addition, printed instructions will be provided with the visual training target card". **Delivery:** home (self-administered) (Table 1). **Regimen:** "Participants will be instructed to continually scan between the various targets for 30 min daily from baseline until their 6-week follow-up, after which they can elect to continue treatment if they wish." **Tailoring:** no. **Modification:** no. **Adherence:** "There were 73 protocol deviations in 58 patients (68.2% overall: 77% in the Fresnel prisms arm, 93% in the visual search arm and 34.5% in the standard care arm). The majority of deviations ($n = 41$, 56.2%) related to lack of compliance in the intervention arms (e.g. prism not worn a minimum of 2 hours daily for 6 weeks or visual exercises not carried out for 30 minutes daily for 6 weeks). Compliance level was similar across the intervention arms ... patients in the visual search training arm followed the visual search exercises 28 days on average."

Group 3: standard care ($n = 30$)

Intervention: advice only ("all three arms will receive the same information leaflets")

Intervention type: control

Materials: "Participants were given information leaflets from the UK Stroke Association and the UK Royal National Institute for the Blind about visual impairment following stroke." **Where can materials be accessed?** UK Stroke Association and the UK Royal National Institute for the Blind. **Procedures:** standard care. **Provided by:** NA. **Delivery:** NA (Table 1). **Regimen:** NA. **Tailoring:** NA. **Modification:** NA. **Adherence:** NA

Outcomes

See Table 4

Visual field assessment

Reading ability (Radner reading test)

Rowe 2010 (Continued)

Visual function questionnaire (VFQ 25-10)
 Rivermead mobility index
 Nottingham extended activities of daily living assessment
 ED-5Q (a standardised instrument for measuring health outcome)
 Short Form-12 (SF-12)
 Adverse events
Time points when outcomes were assessed: baseline, 6-week, 12-week, and 26-week follow-up

Notes Review authors Fiona Rowe and Alex Pollock were involved in this study.

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	"Participants were individually randomized to one of three treatment groups using a secure (24-hour) web-based randomization programme. Randomization lists were generated using block randomization stratified by centre and degree of hemianopia (partial or complete) with treatment allocation ratio of 1:1:1. The local PI (orthoptist) obtained the treatment allocation and subsequently assigned the participant to the treatment arm."
Blinding (performance bias and detection bias) All outcomes	Low risk	Participants and clinical personnel unable to be blinded due to the nature of the intervention. Blinded outcome assessors
Incomplete outcome data (attrition bias) All outcomes	Low risk	Incomplete data all accounted for. Intention-to-treat analysis (where possible)
Other bias	Low risk	No other cause of bias noted

Schuett 2012

Methods

Compensative intervention versus compensative intervention

Design: randomised cross-over trial

Stratification: no

Randomisation sequence: "randomly allocated"

Comparisons: 2 groups, Group A (VET followed by RT) and Group B (RT followed by VET)

Allocation concealment: not stated

Blinding: not stated

Power calculation: not stated

Intention-to-treat analysis: not stated

Other recruitment details: not stated

Patient and public involvement: not stated

Schuett 2012 (Continued)

Participants

Total study population: 36: Group A: 17/18 stroke and Group B: 17/18 stroke. The study included a mixed population.

Withdrawals: none

Method of diagnosing VFD: not stated

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: not stated (although the authors reported that "none of the patients had visual neglect as assessed by tests in accordance with the Behavioural Inattention Test (line bisection, letter and star cancellation, figure and shape copying, drawing from memory; [Halligan 1991](#))." and "All patients were native German speakers and had at least 5 years of education. All patients complained of moderate to severe difficulties in reading and visual exploration and showed impaired performance in both domains".

Exclusion criteria: cerebral visual disorders, including reduced visual acuity (< 0.90 for near and far binocular vision), impaired spatial contrast sensitivity (Vistech contrast sensitivity test, 1988), visual adaptation, disturbances of the anterior visual pathways or of the oculomotor system, macular disease (according to ophthalmologic examination), aphasia, premorbid reading disorders, pure alexia (vertical word reading test; [Zihl 1995](#); [Zihl 2011](#)), impairments of visual-lexical numerical processing (horizontal and vertical number reading; [Zihl 1995](#); [Zihl 2011](#)), or verbal memory deficits

Baseline comparison of treatment groups: none ("before treatment, there were no differences between groups either for demographic and clinical variables or for reading and visual exploration performance".) However, more participants with hemianopia in Group A compared with Group B ([Table 3](#))

This study included patients with diagnoses other than stroke.

This study included patients with visual field defects only (i.e no neglect).

Method of diagnosing visual perceptual problems: "Behavioural Inattention Test (line bisection, letter and star cancellation, figure and shape copying, drawing from memory; [Halligan 1991](#))."

Interventions

Group 1: visual exploration training (VET) (n = 18)

Intervention type: compensation

Intervention: "Visual exploration training: for improving visual exploration, we used standardized versions of the visual search paradigm (parallel and serial search mode)."

Materials: software-based reading and visual exploration training programmes (developed by [Zihl 2011](#)); training material was presented using a LCD monitor with a stimulus display.

Where can materials be accessed? not stated (but references provided). **Procedures:** "Patients were systematically trained to use larger saccadic eye movements to gain a quick complete visual overview as well as to develop and use a more efficient oculomotor visual exploration or scanning strategy that can be flexibly adapted to the visual-spatial structure of the respective scene or environment. Training material consisted of visual search displays extending 50° horizontally and 42° vertically. We used different target and distractor letters of varying similarity as stimuli. Stimulus size was 2.5°, and we used the same colours for the training material and the monitor background as in the reading training. Each training trial was composed of the presentation of a visual search display. Patients were instructed to fixate on a cross in the centre of the monitor and to search, after its offset, for a single target letter (e.g. 'T') among distractor letters (e.g. 'O's) as accurately and quickly as possible. In target-present trials, the patient was asked to press the left mouse button, in target-absent trials, the right mouse button. Presentation and, thus, visual search time was unlimited (exhaustive visual search)." **Provided by:** treatment was administered and supervised by the experimenter. No details about the training or experience of the experimenter. **Delivery:** not clear ([Table 1](#)). **Regimen:** "An individual training session lasted ~45 min consisting of 10-15 practice units (20 trials each) and short or, if required, longer breaks between units. Training was completed when patients reached a defined criterion (at least 90% correct responses) for any level of difficulty used. Patients required on average 12 training sessions, which

Schuett 2012 (Continued)

were carried out within 2–3 weeks for each patient. . . Visual exploration training: Group A: 12.3 sessions (SD 3.4)." **Tailoring:** "In addition to varying letter similarity during the course of training, visual search difficulty was also systematically increased by increasing the visual display size, i.e. the number of stimuli, not the display area (15–20-item displays). This training protocol was adjusted to individual reading performance and training progress. **Modification:** (see tailoring). **Adherence:** "Training was completed when patients reached a defined criterion (at least 90% correct responses) for any level of difficulty used".

Group 2: reading training (RT) (n = 18)

Intervention type: compensation

Intervention: reading training

Materials: "The treatment was performed using the software-based reading and visual exploration training programmes as developed by Zihl 2011; training protocol and procedure were similar to our previous studies (Schuett 2008; Zihl 1995; Zihl 2011). Training material was presented using a LCD monitor with a stimulus display extending 50° horizontally and 42° vertically at a viewing distance of 115 cm." **Where can materials be accessed?** not stated. **Procedures:** "single words of different lengths, ranging from 3 to 12 letters, were used as training material. Letter and digit size was 2.5, and width subtended 1°; spacing between letters (text material) was 0.4°. We used yellow for the training material and a dark blue for the background. These size and colour specifications have been shown to allow for comfortable reading and oculomotor training (Schuett 2008; Zihl 1995; Zihl 2011). Each training trial was composed of the time-limited presentation of one single word in the centre of the screen. Patients were instructed to perceive each word as a whole before reading it aloud by intentionally shifting their gaze, as quickly as possible, from the screen's centre to the beginning (in cases with left-sided visual field loss) or to the end (in cases with right-sided visual field loss) of each word. This paradigm allows reading-related eye movements to be trained and reinforced by the patient's normal internal visual feedback and feedback given by the experimenter. During the course of training, the length of the presented words was systematically increased from 3- to 13-letter words. When a patient was able to read at least 90% of the words of a given length correctly, presentation time was reduced from 1000 ms to 300–400 ms. The final training stage involved the randomized presentation of words of different lengths. By adopting this procedure, patients were forced to make quicker and more efficient saccades in order to perceive and read the whole word before its disappearance. In addition, patients learned to flexibly adjust the size of saccades according to word length". **Provided by:** treatment was administered and supervised by the experimenter. No details about the training or experience of the experimenter. **Delivery:** not clear (Table 1). **Regimen:** "an individual training session lasted ~45 min; it consisted of 10 practice units (30 trials each) and short or, if required, longer breaks between units. Training was completed when patients reached a defined criterion (at least 90% correct responses) for any level of difficulty used. Patients required on average 12 training sessions, which were carried out within 2–3 weeks for each patient ... Group B (12.6 sessions, SD 2.4)." **Tailoring:** "training protocol was adjusted to individual reading performance and training progress". **Modification:** not stated. **Adherence:** "Training was completed when patients reached a defined criterion (at least 90% correct responses) for any level of difficulty used".

Outcomes

See Table 4

Reading test (reading time and errors)

Visual exploration (cancellation task - time and errors)

Visual field assessment (Kinetic perimetry using a standard Tubingen perimeter)

Time points when outcomes were assessed: this was a cross-over trial. **Time points before the cross-over were:** T1, initial assessment; T2, before treatment; T3, after the first training component. (Additional time points - after the cross-over were: T4, after the second training component; T5, after a follow-up interval). "Visual field assessment was carried out before and after treatment (T2 and T4)."

Notes
Risk of bias

Schuett 2012 (Continued)

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Stated "randomly allocated". No further details on method of randomisation or concealment
Blinding (performance bias and detection bias) All outcomes	Unclear risk	It was unclear whether there was participant and/or outcome assessor blinding. However the "experimenter" delivered the intervention, and may therefore have taken the outcome measures.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No flow diagram, and no details of recruitment or retention
Other bias	Low risk	No other issues noted

Spitzyna 2007

Methods	<p>Compensative intervention versus control, placebo or no intervention</p> <p>Design: randomised cross-over trial.</p> <p>Stratification: "Because of the small number of patients entering the study, we wished to ensure that the two treatment groups did not become unbalanced on a few key variables so patients were allocated using a modified minimization technique where cumulative measures of two factors (text reading speed (≤ 90 or >90 wpm) and degrees of sparing of right foveal/parafoveal (0 or 2 degrees) vision) were used to minimize the difference between the two groups."</p> <p>Randomisation sequence: modified minimisation technique: "The weighting used for allocation to the group with the lowest total was one. We describe the minimisation as modified because the first subject was not allocated randomly to either group, but deterministically placed in Group 1."</p> <p>Comparisons: "Two-armed study with two therapy blocks in each arm: one group practiced reading moving text (MT) that scrolled from right-to-left, daily for two four week blocks (group 1), while the other had sham therapy (spot-the-difference) for the first block and then crossed over to MT for the second."</p> <p>Allocation concealment: personal communication with authors: "The allocation was concealed from the persons recruiting into the study but not from the tester. The recruiter(s) did not know that the first subject would be placed in group 1 but the tester did".</p> <p>Blinding: personal communication with authors: "The tester was the first author (a psychologist). She was not blind to the therapy. It was not possible to blind her to this as it was a behavioural therapy and she was 'dispensing' this at each time point."</p> <p>Power calculation: no</p> <p>Intention-to-treat analysis: no</p> <p>Other recruitment details: not stated</p> <p>Patient and public involvement: not stated</p>
Participants	<p>Total study population: 22 participants recruited, data for 18</p> <p>Withdrawals: "three dropped out before completing therapy".</p> <p>Method of diagnosing VFD: "Perimetry was performed twice, before and after completion of the therapy blocks (B and T4). Static fields were measured using the automated Humphrey field analyser II (Carl Zeiss Group, California, USA) analysis of the central 10 degrees of vision (central 10-2 threshold test). Dynamic fields were also measured using a Goldmann perimeter (Haag Streit, K�oniz, Switzerland) when</p>

Interventions for visual field defects in people with stroke (Review)

Spitzyna 2007 (Continued)

there were concerns over subject's performance with the automated procedure (a false positive or false negative response rate greater than 15%). The procedures for determining the amount of field sparing were as reported previously. In the event of a discrepancy between the static and dynamic fields or pre- and post-therapy changes, the opinion of the orthoptist who performed the tests (Ms Bronia Unwin) was sought. The precision of the static perimetry is two degrees and all the patients fell into one of two groups, either 0 or 2 degrees spared to the right of fixation."

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: not stated, but described as "participants with fixed homonymous right-sided visual field defect that interfered with reading (hemianopic alexia)." "All the patients had a fixed homonymous defect that had been present for at least 3 months".

Exclusion criteria: not stated

Baseline comparison of treatment groups: some discrepancies between baseline characteristics of groups. Length of time since symptom onset - all participants in Group 1 were over 1 year since onset of symptoms; 4 participants in Group 2 were less than 1 year since onset (3 are 6 months or less). Average age at time of symptom onset was 43 years for group 1 (range 5 to 67 years) and 63 years for group 2 (range 39 to 78 years). "There were no differences between the groups on all of the variables" for general neuropsychological assessments.

The study included a mixed population.

This study included patients with visual field defect only (no visual neglect).

Method of diagnosing visual perceptual problems: not stated

Interventions

Group 1: optokinetic nystagmus inducing reading therapy (n = 11)

Intervention type: compensation

Intervention: horizontal scrolling text from right to left to "induce small-field optokinetic nystagmus"

Materials: "The tapes were made by animating freely available text from a variety of Sherlock Holmes stories (www.citsoft.com/holmes3.html) across a computer screen from right-to-left (i.e. horizontal motion was from the patients' blind field into their seeing field), so-called Times Square presentation. After some minor preparations, such as deleting hyphens and adding a few spaces after each sentence, custom-written software was used to animate the text across the screen at a variable rate. The output from the graphics card was connected to the video-input on a VHS video recorder to produce the tapes. Tape speeds of: 85, 105, 143, 173, 205, 240, and 275 wpm were made with the text presented in Times New Roman font, 24-point, black on white, scrolling across the middle of the screen." **Where can materials be accessed?** not stated. **Procedures:** participants were instructed to read and try to follow a line of text scrolling horizontally from right to left in the middle of a computer screen at speeds of 85 to 275 words per minute. "Patients were instructed to read the story on the tapes and try to follow it, although no tests of comprehension were made to check this." **Provided by:** not stated. **Delivery:** not clear. Patient-led therapy. ([Table 1](#)). **Regimen:** "treatment blocks lasted for 4 weeks each" and participants were asked to record how long they spent on the tasks each day. "The aim was to achieve a minimum of 400 minutes of rehabilitation (20 sessions x 20 minutes) over approximately 4 weeks". **Tailoring:** "patients contacted one of us (G.A.S.) when they required a new (usually faster) tape." **Modification:** no. **Adherence:** "There were no differences between the two groups in total time spent on rehabilitation tasks across both treatment blocks in minutes (range), 914 (865-955); 901 (840-1000), t-test, t (10) 0.50, P = 0.628".

Group 2: control (n = 11)

Intervention type: control

Intervention: sham training; "spot the difference"

Spitzyna 2007 (Continued)

Materials: "'Spot-the-difference' tests were taken from a children's puzzle booklet; the original cartoons were altered to remove text. The number of differences on each page varied between eight and 12."

Where can materials be accessed? not stated. **Procedures:** "Patients were instructed to look for as many differences as possible between the two pictures, but were not told how many to expect, completing at least two cartoons over 20 min. Between 40 to 60 examples were selected randomly for each patient (out of a total of the 70 prepared tests), depending on their speed." **Provided by:** not stated. **Delivery:** not clear. Patient-led therapy. (Table 1). **Regimen:** "treatment blocks lasted for 4 weeks each" and participants were asked to record how long they spent on the tasks each day. "The aim was to achieve a minimum of 400 minutes of rehabilitation (20 sessions x 20 minutes) over approximately 4 weeks". **Tailoring:** not stated. **Modification:** no. **Adherence:** "There were no differences between the two groups in total time spent on rehabilitation tasks across both treatment blocks in minutes (range), 914 (865-955); 901 (840-1000), t-test, t (10) 0.50, P = 0.628"

Outcomes

See Table 4

Text reading speeds (passages from Neale analysis of reading)

Single word reading speeds (words taken from MRC psycholinguistic battery)

Eye movement characteristics - spatial characteristics of saccadic amplitude, incoming saccade amplitude and landing position

- temporal characteristics

Perimetry - Humphrey field analyser

- Goldmann perimetry

Time points when outcomes were assessed: T1, initial assessment; T2, before treatment; T3, after the first training component. (Additional time points - after the cross-over were: T4, after the second training component; T5, after a follow-up interval)

Notes

Did not state whether visual correction using glasses was used during training or testing.

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	Modified minimisation technique of sequence generation. Personal communication with authors: "The allocation was concealed from the persons recruiting into the study but not from the tester. The recruiter(s) did not know that the first subject would be placed in group 1 but the tester did".
Blinding (performance bias and detection bias) All outcomes	High risk	Personal communication with authors: "The tester was the first author (a psychologist). She was not blind to the therapy. It was not possible to blind her to this as it was a behavioural therapy and she was 'dispensing' this at each time point."
Incomplete outcome data (attrition bias) All outcomes	High risk	"Three patients dropped out before completing therapy" (all from group 2) and "data from these subjects were excluded from analyses"
Other bias	Low risk	No other potential bias noted

Szlyk 2005

Methods

Substitutive intervention versus substitutive intervention

Szlyk 2005 (Continued)

Design: randomised cross-over design

Stratification: not stated

Randomisation sequence: "randomly assigned" but method not given

Comparisons: participants in Group 1 received Gottlieb prism during the first 3-month phase of the study, and participants in Group 2 received Fresnel prisms during the first 3-month phase of the study. Participants then crossed over to receive the other treatment.

Allocation concealment: not stated

Blinding: not stated

Power calculation: not stated

Intention-to-treat analysis: not stated

Other recruitment details: not stated

Patient and public involvement: not stated

Participants

Total study population: 10 participants.

Withdrawals: no information (7 participants provided follow-up data at 2 years - 3 could not be contacted)

Method of diagnosing VFD: not stated.

Characteristics of population: participant details are listed in [Table 2](#).

Type and severity of visual problems: participant details are listed in [Table 3](#).

Inclusion criteria: stated "The patients were screened to include patients with only occipital lobe strokes". Participants described as having "Hemianopsia because of cerebral vascular accidents". All included participants were male, but unclear if this was an inclusion criterion.

Exclusion criteria: not stated.

Baseline comparison of treatment groups: yes

The study included a mixed population.

It was assumed that participants did not have neglect (as this is unlikely in occipital lesions).

Method of diagnosing visual perceptual problems: not stated - it was not stated whether the patients may have had visual neglect (although neglect is unlikely in occipital lesions)

Interventions

Group 1: Gottlieb Visual Field Awareness System (VFAS) (n = 5)

Intervention type: substitution

Intervention: Gottlieb VFAS prism

Materials: 18.5 dioptre Gottlieb VFAS prism

Where can materials be accessed? "Rekindle(R), Stone Mountain, GA, USA". **Procedures:** 18.5 dioptre Gottlieb VFAS prism drilled into one lens. Positioned just off pupil centre - generally on the same eye as side of field loss, on same side as field loss, base out. **Provided by:** low-vision specialist for laboratory and outdoor training, kinesiotherapist for on-road training. **Delivery:** laboratory and out-door training within university grounds, and on-road (driving) training on a road course within a medical centre ([Table 1](#)). **Regimen:** training of 4 x 2 to 3-hour sessions indoors with low vision specialist and 8 x 2-hour outdoor sessions behind the wheel. The lenses were then worn for 3 months. **Tailoring:** not stated. **Modification:** not stated. **Adherence:** no information (whether prism was still worn at time of follow-up was recorded as never/occasionally/frequently)

Szlyk 2005 (Continued)

Group 2: Fresnel prisms (n = 5)

Intervention type: substitution

Intervention: Press-On™ Fresnel 20 Diopter Prisms

Materials: Press-On™ Fresnel 20 Diopter Prisms. **Where can materials be accessed?** "3M Health Care, St. Paul, MN, USA". **Procedures:** 20 dioptre press-on Fresnel prisms attached to posterior surface of 1 spectacle lens. Positioned just off pupil centre - generally on the same eye as side of field loss, on same side as field loss, base out. **Provided by:** low-vision specialist for laboratory and outdoor training, kinesiologist for on-road training. **Delivery:** laboratory and out-door training within university grounds, and on-road (driving) training on a road course within a medical centre (Table 1). **Regimen:** training of 4 x 2 to 3 hour sessions indoors with low-vision specialist and 8 x 2-hour outdoor sessions behind the wheel. The lenses were then worn for 3 months. **Tailoring:** not stated. **Modification:** not stated. **Adherence:** no information (whether prism was still worn at time of follow-up was recorded as never/occasionally/frequently)

Outcomes

See Table 4

Goldmann visual field

Visual acuity

Contrast sensitivity

Lab assessment - indoor functional assessment

Outdoor function assessment

Driving skills assessment - indoor and on-road

Psychophysical assessment

- attentional visual acuity

- attentional motion sensitivity

Self-report (satisfaction)

Time points when outcomes were assessed: before and after each cross-over. Continued use of devices assessed at 2-year follow-up

Notes

1. Data were presented after the cross-over, for both groups combined - no first phase data were available.

2. Stated: "For each assessment task for each individual in the test-retest period, we computed the change in score from the initial baseline testing to the repeat baseline testing. We then averaged these change scores across subjects for each task". For each task if the change from baseline to training exceeded the test-retest change it was scored as 'improved', if it was less than or equal it was scored as 'no change'. The sum of improved tasks across the test battery was then computed.

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	Stated "randomly assigned into one of two experimental groups" but no details of method provided
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No details of blinding included

Szlyk 2005 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No incomplete data issues apparent, however, scores have been combined so it is difficult to tell if there are any missing outcomes.
Other bias	Low risk	No other cause of bias noted

AB/BA: refers to order of interventions within cross-over trial, where A and B denote different interventions and AB or BA the order of delivery

ACS: alternating current stimulation

ADL: activities of daily living

ARV: area of residual vision

AVT: audiovisual exploration training

cd/m²: candela per square meter (standard unit of luminance)

CNS: central nervous system

CT: computerised tomography

CVA: cerebrovascular accident

EEG: electroencephalogram

EQ-5D: standardised EuroQol health-related quality of life instrument

FIM: Functional Independence Measure

FT: flicker stimulation training

HRP: high resolution perimetry

HVFD: homonymous visual field defect

IH-CST: InSight Hemianopia - Compensatory Scanning Training

IVI: impact of visual impairment

LCD: liquid crystal display (high definition monitor)

LEDs: light emitting diodes

LV-VFQ: Low Vision - Visual Function Questionnaire

MMSE: Mini Mental State Examination

MRC: Medical Research Council

MRI: magnetic resonance imaging

MT: moving text

n: number

NA: not applicable

NEI-VFQ: National Eye Institute Visual Function Questionnaire

OT: occupational therapy

PI: principal investigator

QoL: quality of life

RCT: randomised controlled trial

RT: restitution therapy

SD: standard deviation

SF-12: Short-Form Health Survey

T1/T2/T3/T4/T5: outcome assessment timepoint 1, 2, 3, 4 or 5

TAP: Tubinger automated perimetry

TBI: traumatic brain injury

tDCS: transcranial direct current stimulation

TL: tube luminescent

VET: visual exploration training

VFAS: visual field awareness system

VFD: visual field defect

VFQ-25: visual functioning questionnaire

VIS: Visual In Stroke (study name)

VRT: vision restoration therapy

VST: visual stimulation training

WHOQOL-BREF:

wpm: words per minute

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Akinwuntan 2005	The protocol for this review was to include "any intervention that is specifically targeted at improving the visual field defect, or improving the ability of the participant to cope with the visual field loss". Although this randomised controlled intervention study did measure visual, neuro-physical and driving outcomes, the intervention targeted a person's decreased ability to drive, not specifically due to their visual field defect, but by all the "motor, visual, cognitive, and perceptual deficits ... experienced after stroke", and thus was ineligible.
Akinwuntan 2010	Not VFD
Akinwuntan 2012	Not VFD
Alber 2015	Not RCT
Balconi 2013	Not VFD
Beasley 2013	Not VFD
Bowers 2012	Not RCT
Braga 2018	Not RCT
Brandt 2009	Not RCT
Brigui 2014	Not RCT
Cameirao 2012	Not vision
Carter 1983	<p>***Was an included study in 2011 version of Cochrane review***</p> <p>Excluded from updated version because participants were not confirmed to have visual field defects: this study included 33 participants, all of whom had a diagnosis of stroke. The 2011 version stated: "this study did not diagnose visual field defects but instead relied on clinical identification of a 'visual scanning problem'. Visual scanning problems are likely to be due to visual neglect as well as visual field defects, thus we have to assume that the participants in this study had a combination of problems. However, it could be possible that the participants had visual neglect problems but not visual field defects, and vice versa."</p>
Carter 1988	The protocol for this review was to include "any intervention that is specifically targeted at improving the visual field defect, or improving the ability of the participant to cope with the visual field loss". The intervention in this study was cognitive skills retraining and ADL retraining; it included visual scanning training within a large battery of training activities, but this was not specifically targeted at visual field loss.
Chen 2013	Not stroke
ChiCTR-OON-15006688	Not RCT
Cho 2015	Not VFD
Courtney-Harris 2015	Aim of study was validation of a vision screening tool; and outcomes were focussed on level of agreement between tests.
Crotty 2009	Not VFD

Study	Reason for exclusion
Cutfield 2011	Not RCT
Dai 2014	Not RCT
Dargie 2012	Not RCT
Davis 2009	Not vision
Fedorov 2010	Not VFD
Ginsberg 2013	Not VFD
Gordon 1985	The method of sequence generation was described as "patients ... were assigned to either experimental or control condition, depending on the rehabilitation service to which the patient was assigned for treatment. All patients were accessed from two comparable inpatient services; experimental and control conditions were alternated every six months between the two services during the time which all patients on a given service were assigned to the same treatment condition". Although this was described as "quasi-random" it did not meet the level of randomisation of sequence generation that was needed for inclusion.
Hamel 2012	Not RCT
Hazelton 2013	Not RCT
Hollands 2013	Not VFD
Jo 2012	Visual neglect, not VFD
Kang 2009	Not VFD
Kerkhoff 2013	Visual neglect, not VFD
Kerry 2017	Focussed on central alexia ("an acquired reading disorder co-occurring with a generalised language deficit (aphasia)")
Ko 2011	Not RCT
Lane 2010	The method of sequence generation in this study of compensation interventions was stated as "the first 23 were assigned to Group A, and the last 23 to Group B" and thus was non-random.
Lee 2013a	Visual neglect, not VFD
Lee 2013b	Not RCT
Loverro 1988	The method of sequence generation in this study of bed orientation was stated as "assigned to rehabilitation hospital beds based on bed availability" and thus was non-random.
Machner 2012	Visual neglect, not VFD
Mancuso 2012	Visual neglect, not VFD
Markowitz 2010	This was a comment on a study: Bergsma DP and Van der Wildt G. British Journal of Ophthalmology 2010; 94:88–96, which was not a randomised controlled trial.
NTR5637	Not RCT

Study	Reason for exclusion
Olma 2013	Not RCT
Opolka 2013	Visual neglect, not VFD
Padula 2009	Not RCT
Park 2013	Not VFD
Rosenberg 2011	Not VFD
Sabel 2009	This study looked at visual restitution in optic nerve disorder. The consensus of the Advisory Group for this review was that the review should be limited to post-chiasmal visual field loss only, so this study was not included.
Saposnik 2013	Not VFD
Scholomov 2010	Not RCT
Taylor 2011	Not RCT
Teasell 2011	Visual neglect, not VFD
Thurtell 2010	Not VFD
Van Wyk 2011	Not VFD
Van Wyk 2016	Not VFD (eye movement disorders)
Weinberg 1977	<p>***Was an included study in 2011 version of Cochrane review***</p> <p>Excluded from updated version because participants were not confirmed to have visual field defects. Weinberg 1977 and Weinberg 1979 both included participants with the same inclusion criteria. The 2009 version stated: "These studies included only people with right-sided cerebrovascular accident. However, from the limited information available in the published papers, it does appear that the diagnosis of visual field defect (or visual neglect) was not an inclusion criterion for entry to these studies. The authors appear to assume that this patient group will have visual neglect as they have right-sided brain damage. It is worth noting that in these two studies the participants were divided into subgroups of 'mild' and 'severe' based on "presence of visual field defect on confrontation and performance on visual cancellation", and that we can therefore be confident that the 'severe' subgroup contained people with visual field defects, while it is not confirmed whether all the 'mild' subgroup had a visual field defect. Arguably, therefore, the 'severe' subgroups are more relevant to our review question than the 'mild' subgroups. There was substantial heterogeneity when we included the 'mild' subgroups in the analyses. Exploring this heterogeneity using subgroup analyses found that the 'severe' subgroup had a greater response to the intervention than the 'mild' subgroup, for all outcomes ... Arguably it was inappropriate to have included the 'mild' subgroups within any of the analyses, as it is possible that participants in this group did not have a visual field defect. However, due to the uncertainty we felt it was appropriate to include the 'mild' subgroup but to further explore the effect of removing them from the analyses."</p>
Weinberg 1979	<p>***Was an included study in 2011 version of Cochrane review***</p> <p>Excluded from updated version because participants were not confirmed to have visual field defects. Weinberg 1977 and Weinberg 1979 both included participants with the same inclusion criteria. The 2009 version stated: "These studies included only people with right-sided cerebrovascular accident. However, from the limited information available in the published papers, it does appear that the diagnosis of visual field defect (or visual neglect) was not an inclusion criterion for entry to these studies. The authors appear to assume that this patient group will have visual neglect as they have right-sided brain damage. It is worth noting that in these two studies the participants</p>

Study	Reason for exclusion
	<p>were divided into subgroups of 'mild' and 'severe' based on "presence of visual field defect on confrontation and performance on visual cancellation", and that we can therefore be confident that the 'severe' subgroup contained people with visual field defects, while it is not confirmed whether all the 'mild' subgroup had a visual field defect. Arguably, therefore, the 'severe' subgroups are more relevant to our review question than the 'mild' subgroups. There was substantial heterogeneity when we included the 'mild' subgroups in the analyses. Exploring this heterogeneity using subgroup analyses found that the 'severe' subgroup had a greater response to the intervention than the 'mild' subgroup, for all outcomes ... Arguably it was inappropriate to have included the 'mild' subgroups within any of the analyses, as it is possible that participants in this group did not have a visual field defect. However, due to the uncertainty we felt it was appropriate to include the 'mild' subgroup but to further explore the effect of removing them from the analyses."</p> <p>While we were confident that the 'severe' subgroup from this study contained some participants with visual field defects, this was not the focus of this trial, which was focused on participants with right brain damage only. Further, we cannot be certain that all participants in this group had visual field defects (there is a chance that some had severe visual neglect but no visual field defect).</p>
White 2010	Not VFD

ADL: activities of daily living
 RCT: randomised controlled trial
 VFD: visual field defect

Characteristics of studies awaiting assessment [ordered by study ID]

Ghandehari 2011

Methods	Methods described this as a "prospective, observational study". Results described as a "randomised controlled trial", and included a study flowchart. No details of randomisation were provided.
Participants	40 participants: "stroke patients with posterior cerebral artery territory infarction" with "pure homonymous hemianopia without neglect"
Interventions	Group 1: Neuroaid (MLC601) Group 2: Piracetam
Outcomes	Visual field (standard perimetry)
Notes	Study carried out in Iran during 2009-2010 Authors emailed in 2015, but no reply

Sand 2017

Methods	Conference abstract: "Patients were randomised ..." ClinicalTrials.gov: "observational" study
Participants	Stroke patients - not clear if VFD was an inclusion criteria
Interventions	Experimental: "training with vision teacher with individually adapted training program"

Sand 2017 *(Continued)*

	Control: no training
Outcomes	Visual field defect Visual function Quality of life National Institute of Health Stroke Scale (NIHSS)
Notes	ClinicalTrials.gov Identifier: NCT02307981 Unclear if this was an RCT or not

NIHSS: National Institute of Health Stroke Scale

RCT: randomised controlled trial

VFD: visual field defect

Characteristics of ongoing studies *[ordered by study ID]*
Feldon 2017

Trial name or title	Visual restoration for hemianopia
Methods	RCT
Participants	Adults (21 to 75 years); lesion in the occipital lobe of the brain (18 to 90 days previously); stable homonymous hemianopia
Interventions	Experimental: training in the blind field: A computer software and chin-rest necessary to perform visual training will be loaned to each participant to use at home. They will perform 1 to 2 daily training sessions in their home, consisting of 200 to 300 trials each. The visual task performed repetitively will involve discriminating the direction of motion of a small cloud of dots located at a predetermined location in the blind field. The computer program will automatically create a record of participant performance during each home training session. They will train daily (about 40 to 60 minutes total), 5 to 7 days per week, for at least 24 weeks. Control: training in the intact field (as above)
Outcomes	Primary outcome measure: 24-2 Humphrey PMD - Change in the perimetric mean deviation (PMD) from 24-2 Humphrey perimetry between baseline and post-training
Starting date	March 2018 (Estimated completion: May 2019)
Contact information	lisa_blanchard@urmc.rochester.edu
Notes	ClinicalTrials.gov Identifier: NCT03350919

Gall 2015

Trial name or title	Restoration of vision after stroke (REVIS)
Methods	"Randomised"

Interventions for visual field defects in people with stroke (Review)

Gall 2015 (Continued)

Participants	Adults aged 18 to 75 years; > 6 months post stroke; hemianopia or quadrantanopia confirmed by standard automated perimetry
Interventions	Experimental: active tDCS (transcranial direct current stimulation) Control: sham stimulation
Outcomes	Primary outcome: improved detection in the visual field, measured using high resolution perimetry Secondary outcome: change in extent of visual fields using standard automated perimetry
Starting date	April 2015 (study completion: April 2017)
Contact information	turgut.tatlisumak@hus.fi
Notes	ClinicalTrials.gov Identifier: NCT02405143

Hayes 2010

Trial name or title	Retraining following hemianopia in acquired brain injury following stroke
Methods	Comparison of 2 intensities of scanning training
Participants	20 participants
Interventions	Visual scanning training using a light panel
Outcomes	Includes quality of life measures and functional vision skills
Starting date	
Contact information	ahayes@nvtsystems.com.au
Notes	Unpublished. Recruitment has ended. ACTRN12610000494033

ISRCTN16023965

Trial name or title	Home-training for hemianopia (partial blindness)
Methods	RCT (3 groups)
Participants	Adults (aged at least 18) suffering with a nonprogressive visual field defect for at least 3 months caused by a brain injury
Interventions	Group 1 (intervention group 1): use the DREX program on a touchscreen tablet Group 2 (intervention group 2): use the DREX program on a computer Group 3 (control group): given their usual care and any treatments given by their doctors or therapists
Outcomes	"Reading and visual search abilities and also their quality of life"

Interventions for visual field defects in people with stroke (Review)

ISRCTN16023965 (Continued)

Starting date	June 2015 (study completion: December 2017)
Contact information	azuwan.musa@durham.ac.uk
Notes	ISRCTN16023965

Komm 2009

Trial name or title	Rehabilitation outcome in chronic neglect patients with associated visual field loss: results of a blinded randomized cross-over study
Methods	Masked randomised cross-over study
Participants	Stroke patients with persistent neglect and visual field loss 6 months post-onset
Interventions	Computer-aided visual restitution training versus compensatory eye movement training
Outcomes	Visual field and neglect measures
Starting date	
Contact information	bernhard.sabel@med.ovgu.de
Notes	Currently in press. Results not available for this review

NCT02737930

Trial name or title	Fluoxetine for visual recovery after ischemic stroke (FLUORESCE)
Methods	RCT
Participants	MRI-confirmed acute ischaemic stroke resulting in an isolated homonymous visual field loss Estimated enrolment: 40 participants
Interventions	Experimental: fluoxetine - 20 mg fluoxetine capsule by mouth once daily for 90 days Placebo comparator: placebo
Outcomes	Primary outcome measure: improvement in size of visual field deficit (degrees) Secondary outcome measures: Improvement in size of visual field deficit (square degrees) Improvement in parametric mean deviation Functional field score Visual Function Questionnaire-25 score Patient Health Questionnaire-9 score Modified Rankin Scale score

Interventions for visual field defects in people with stroke (Review)

NCT02737930 (Continued)

Post-stroke changes in cortical visual representation as measured by functional magnetic resonance imaging

Post-stroke changes in retinal nerve fibre layer thickness

Starting date	May 2016 (Estimated study completion: June 2020)
Contact information	bogachan_sahin@urmc.rochester.edu
Notes	ClinicalTrials.gov identifier: NCT02737930

NCT02886663

Trial name or title	Visual field restoration in patients with post-stroke homonymous hemianopsia (REVOIR)
Methods	Randomised cross-over design
Participants	Adults, 3 to 36 months post-stroke; presence of a homonymous visual field amputation after vascular retro-chiasmatic lesion of the occipital region (visual cortex alone or with optical radiations or with other associated occipital areas) Anticipated recruitment: 104 participants
Interventions	Experimental: immediate rehabilitation ("a stimulus, in the blind field of hemianopsic patients, to restore the vision of patients with homonymous hemianopsia consecutive to stroke") Control: delayed rehabilitation
Outcomes	Primary outcome measure: change in the sensitivity of detection of a visual stimulus after rehabilitation
Starting date	September 2016
Contact information	lsalomon@for.paris
Notes	ClinicalTrials.gov Identifier: NCT02886663

DREX: Durham Reading & Exploration

PMD: perimetric mean deviation

RCT: randomised controlled trial

tDCS: transcranial direct current stimulation

VFD: visual field defect

DATA AND ANALYSES
Comparison 1. Restitutive interventions versus control, placebo or no intervention

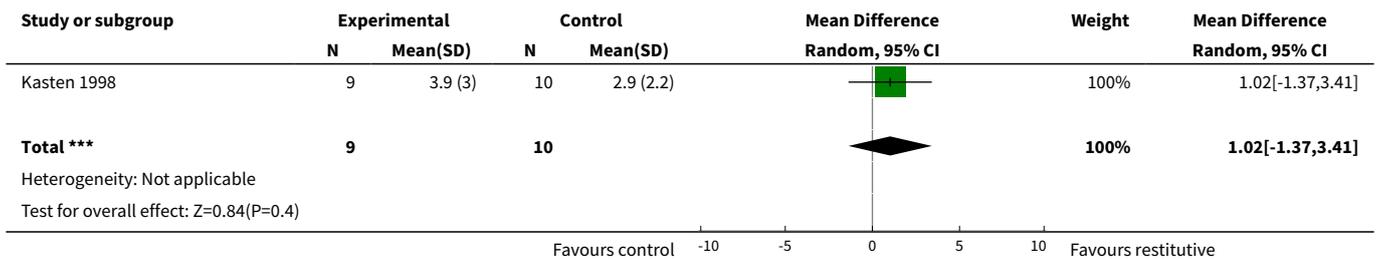
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Visual field	1	19	Mean Difference (IV, Random, 95% CI)	1.02 [-1.37, 3.41]

Interventions for visual field defects in people with stroke (Review)

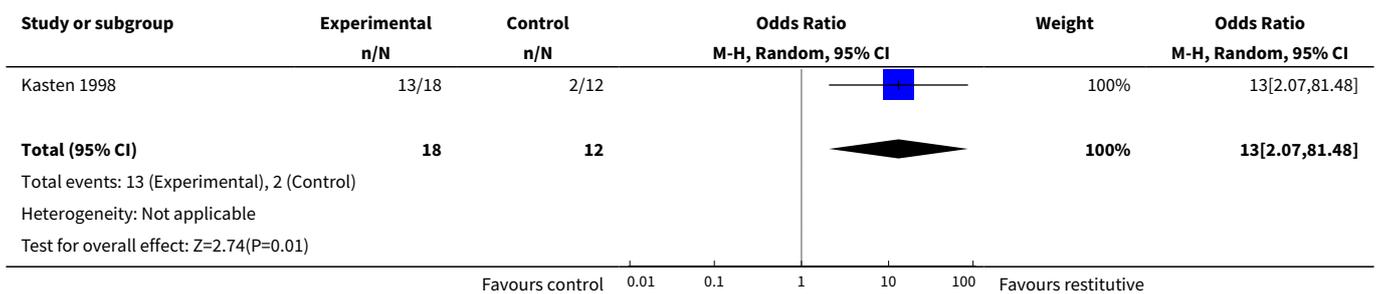
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Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
2 Quality of life	1	30	Odds Ratio (M-H, Random, 95% CI)	13.0 [2.07, 81.48]

Analysis 1.1. Comparison 1 Restitutive interventions versus control, placebo or no intervention, Outcome 1 Visual field.



Analysis 1.2. Comparison 1 Restitutive interventions versus control, placebo or no intervention, Outcome 2 Quality of life.

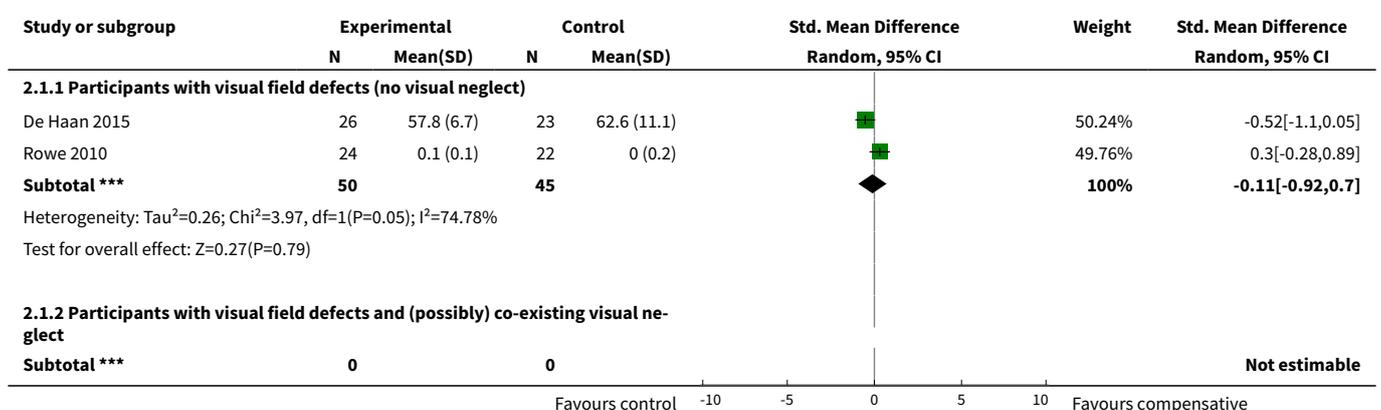


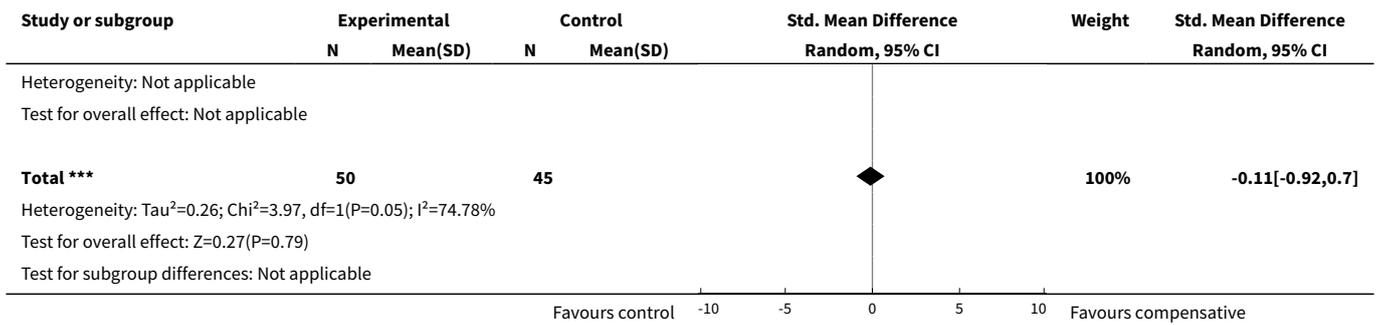
Comparison 2. Compensative interventions versus control, placebo or no intervention

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Visual field	2	95	Std. Mean Difference (IV, Random, 95% CI)	-0.11 [-0.92, 0.70]
1.1 Participants with visual field defects (no visual neglect)	2	95	Std. Mean Difference (IV, Random, 95% CI)	-0.11 [-0.92, 0.70]
1.2 Participants with visual field defects and (possibly) co-existing visual neglect	0	0	Std. Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
2 Extended activities of daily living	2	87	Std. Mean Difference (IV, Random, 95% CI)	0.49 [-0.01, 0.99]

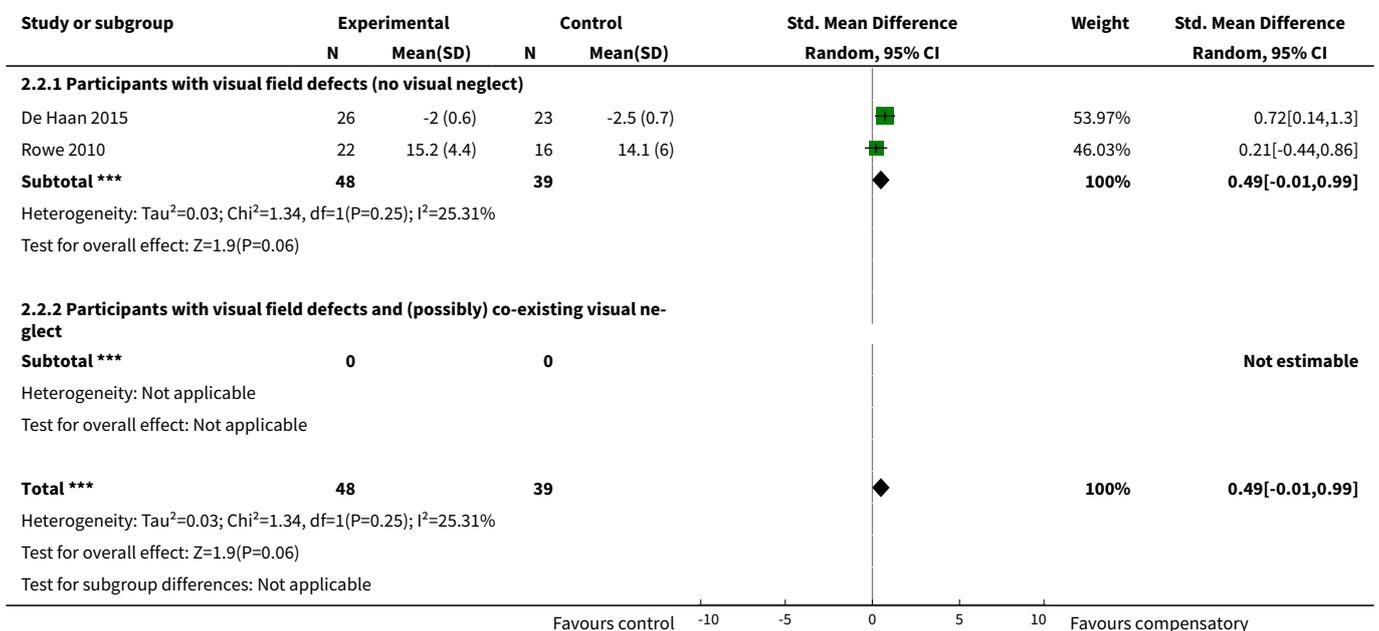
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
2.1 Participants with visual field defects (no visual neglect)	2	87	Std. Mean Difference (IV, Random, 95% CI)	0.49 [-0.01, 0.99]
2.2 Participants with visual field defects and (possibly) co-existing visual neglect	0	0	Std. Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
3 Reading	4	162	Std. Mean Difference (IV, Random, 95% CI)	0.26 [-0.05, 0.58]
3.1 Participants with visual field defects (no visual neglect)	3	110	Std. Mean Difference (IV, Random, 95% CI)	0.18 [-0.20, 0.56]
3.2 Participants with visual field defects and (possibly) co-existing visual neglect	1	52	Std. Mean Difference (IV, Random, 95% CI)	0.45 [-0.10, 1.00]
4 Quality of life	2	96	Mean Difference (IV, Random, 95% CI)	9.36 [3.10, 15.62]
4.1 Participants with visual field defects (no visual neglect)	2	96	Mean Difference (IV, Random, 95% CI)	9.36 [3.10, 15.62]
4.2 Participants with visual field defects and (possibly) co-existing visual neglect	0	0	Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
5 Scanning - cancellation	2	97	Std. Mean Difference (IV, Random, 95% CI)	-0.01 [-0.40, 0.39]
5.1 Participants with visual field defects (no visual neglect)	1	48	Std. Mean Difference (IV, Random, 95% CI)	0.12 [-0.45, 0.68]
5.2 Participants with visual field defects and (possibly) visual neglect	1	49	Std. Mean Difference (IV, Random, 95% CI)	-0.13 [-0.69, 0.44]
6 Adverse events	2	108	Odds Ratio (M-H, Random, 95% CI)	5.18 [0.24, 112.57]

Analysis 2.1. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 1 Visual field.

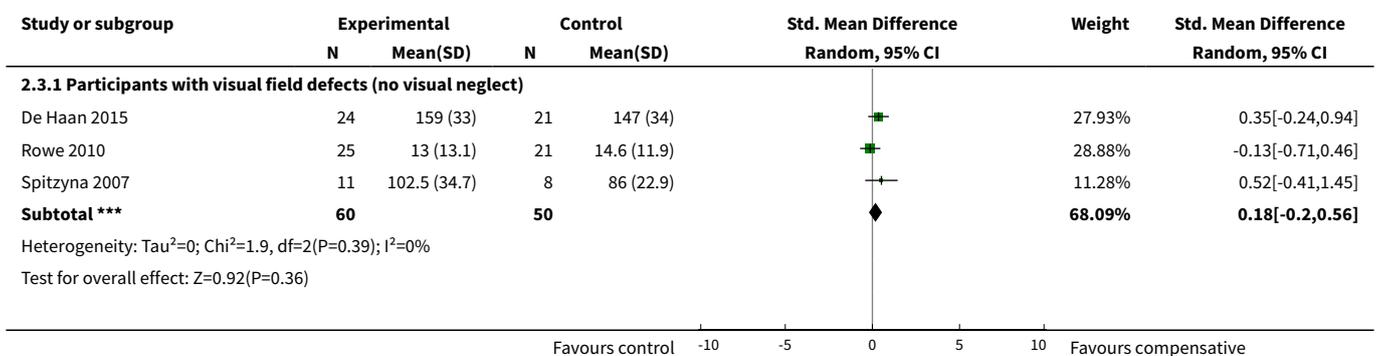


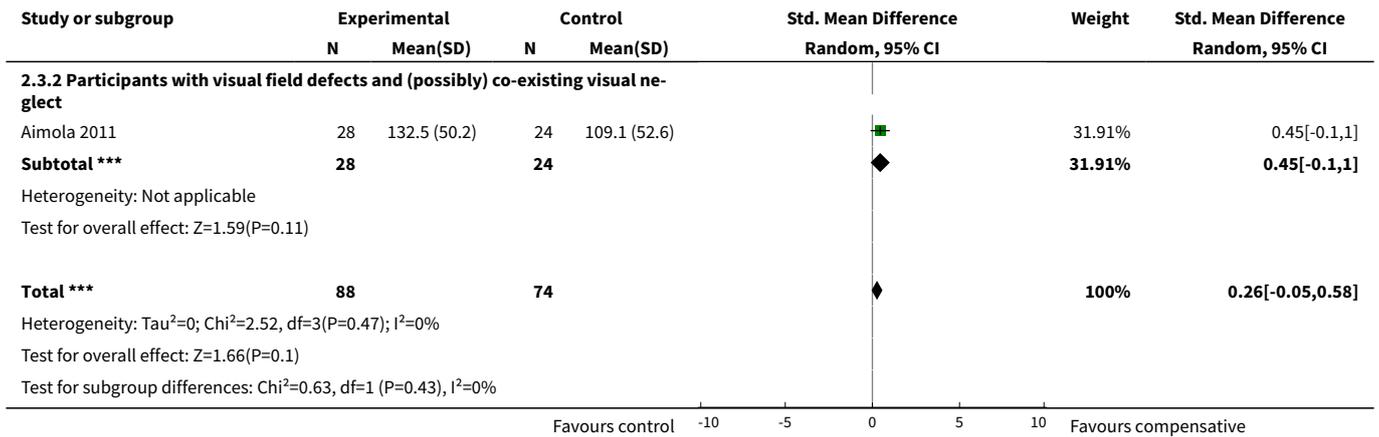


Analysis 2.2. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 2 Extended activities of daily living.

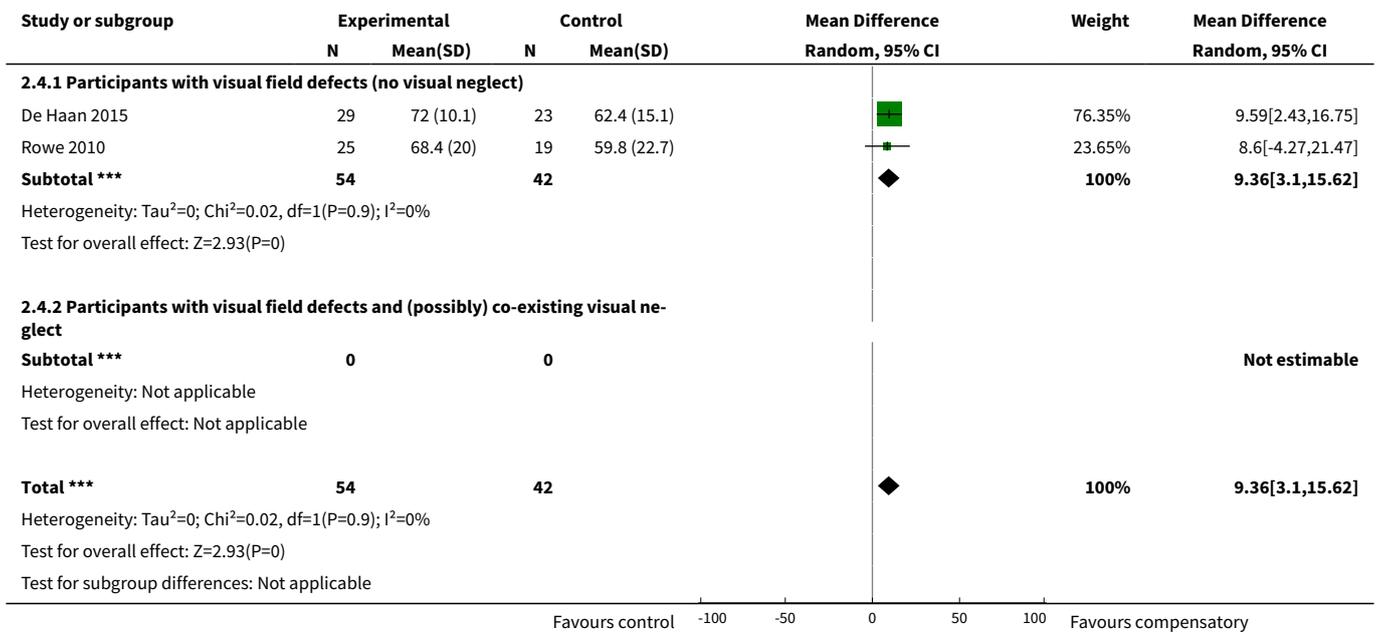


Analysis 2.3. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 3 Reading.

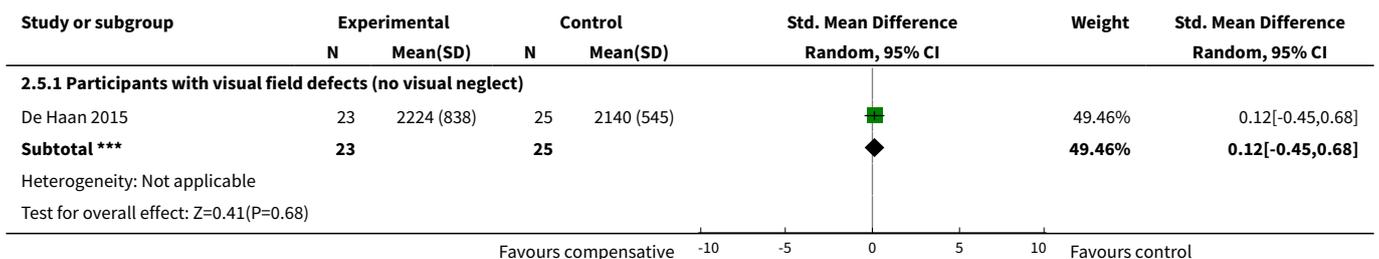


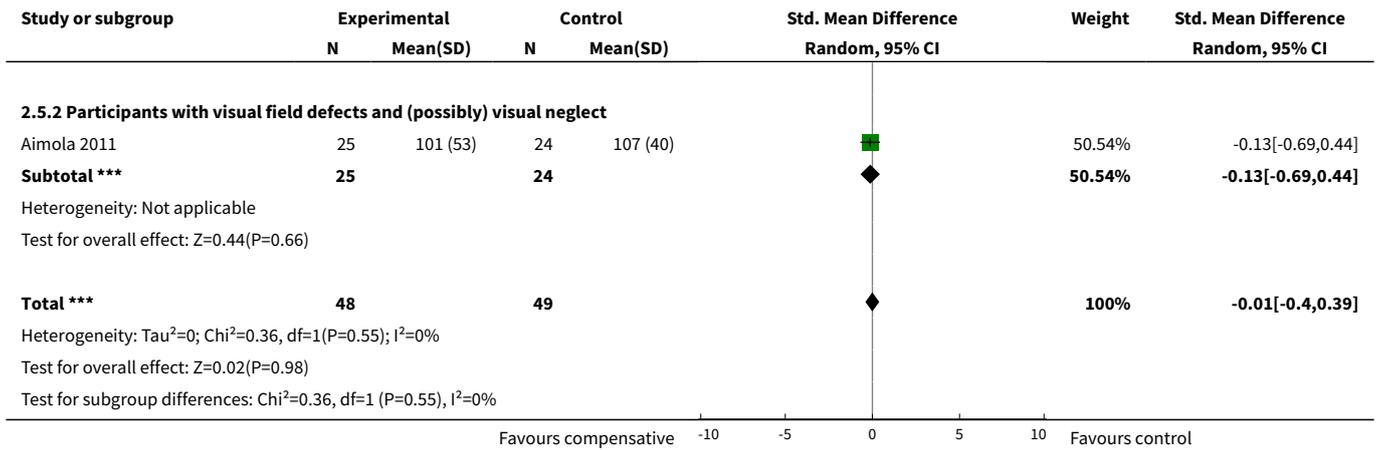


Analysis 2.4. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 4 Quality of life.

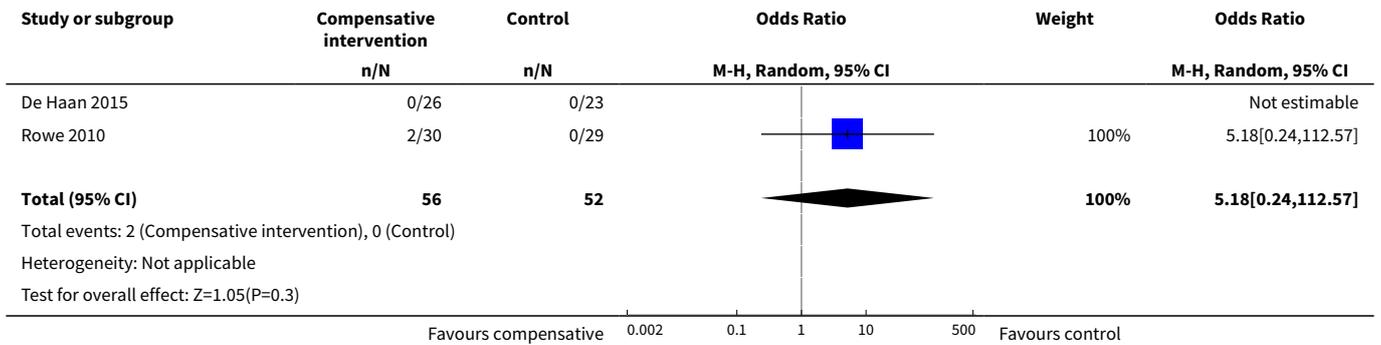


Analysis 2.5. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 5 Scanning - cancellation.





Analysis 2.6. Comparison 2 Compensative interventions versus control, placebo or no intervention, Outcome 6 Adverse events.

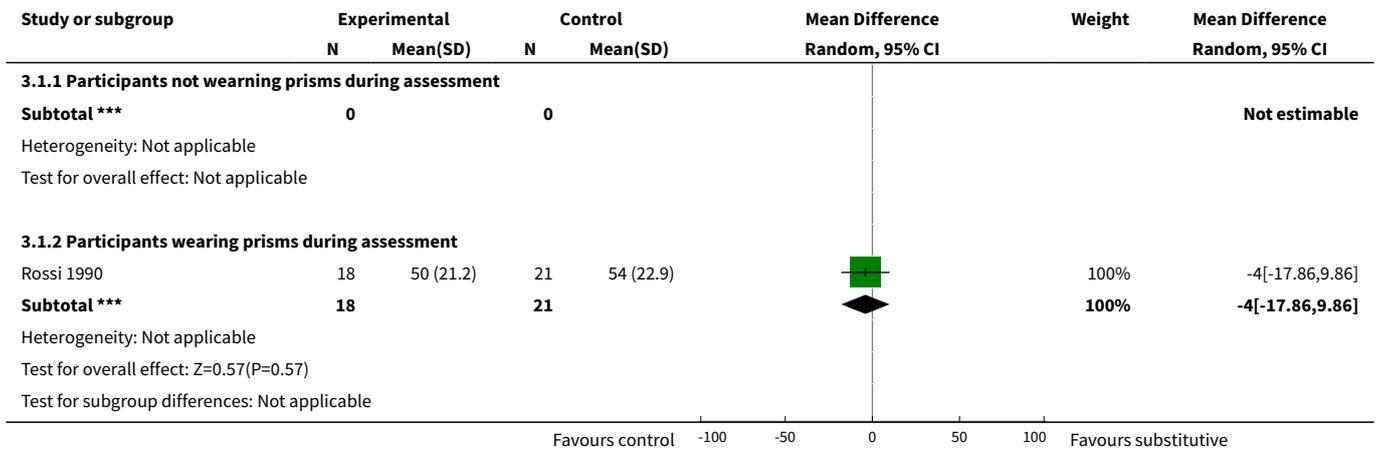


Comparison 3. Substitutive interventions versus control, placebo or no intervention

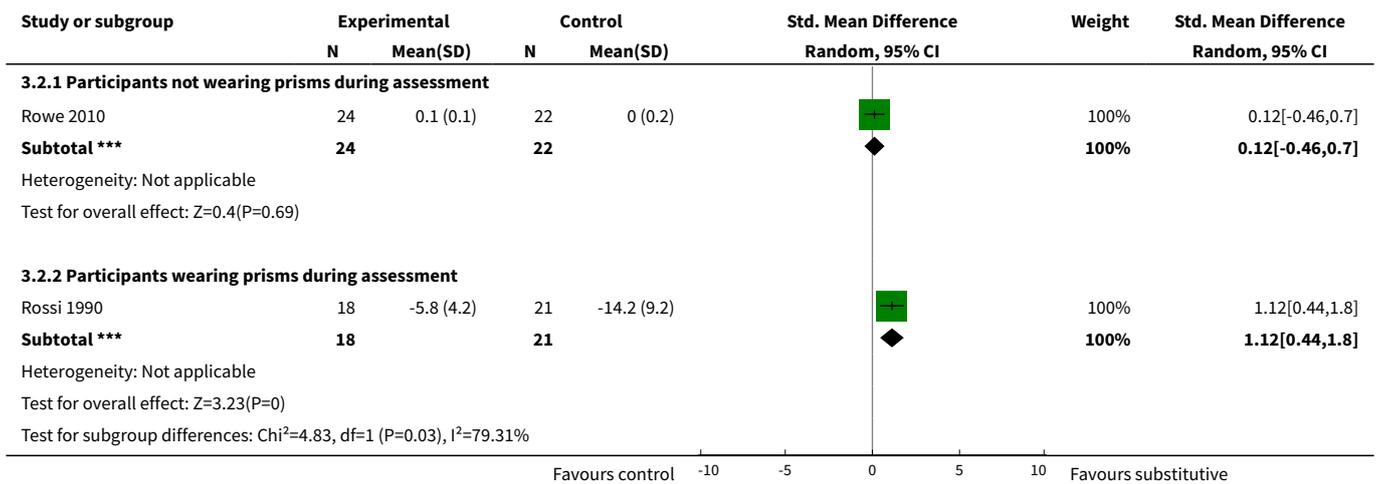
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Functional Activities of Daily Living	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
1.1 Participants not wearing prisms during assessment	0	0	Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
1.2 Participants wearing prisms during assessment	1	39	Mean Difference (IV, Random, 95% CI)	-4.0 [-17.86, 9.86]
2 Visual field	2		Std. Mean Difference (IV, Random, 95% CI)	Subtotals only
2.1 Participants not wearing prisms during assessment	1	46	Std. Mean Difference (IV, Random, 95% CI)	0.12 [-0.46, 0.70]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
2.2 Participants wearing prisms during assessment	1	39	Std. Mean Difference (IV, Random, 95% CI)	1.12 [0.44, 1.80]
3 Extended activities of daily living	2		Std. Mean Difference (IV, Random, 95% CI)	Subtotals only
3.1 Participants not wearing prisms during assessment	1	38	Std. Mean Difference (IV, Random, 95% CI)	0.20 [-0.44, 0.85]
3.2 Participants wearing prisms during assessment	1	61	Std. Mean Difference (IV, Random, 95% CI)	0.24 [-0.26, 0.75]
4 Reading	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
4.1 Participants not wearing prisms during assessment	1	45	Mean Difference (IV, Random, 95% CI)	2.80 [-7.13, 12.73]
4.2 Participants wearing prisms during assessment	0	0	Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
5 Falls	1		Odds Ratio (M-H, Random, 95% CI)	Subtotals only
5.1 Participants not wearing prisms during assessment	0	0	Odds Ratio (M-H, Random, 95% CI)	0.0 [0.0, 0.0]
5.2 Participants wearing prisms during assessment	1	39	Odds Ratio (M-H, Random, 95% CI)	1.21 [0.26, 5.76]
6 Quality of life	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
6.1 Participants not wearing prisms during assessment	1	43	Mean Difference (IV, Random, 95% CI)	8.40 [-4.18, 20.98]
6.2 Participants wearing prisms during assessment	0	0	Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
7 Scanning - cancellation	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
7.1 Participants not wearing prisms during assessment	0	0	Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
7.2 Participants wearing prisms during assessment	1	39	Mean Difference (IV, Random, 95% CI)	9.80 [1.91, 17.69]
8 Adverse events	1	59	Odds Ratio (M-H, Random, 95% CI)	87.32 [4.87, 1564.66]

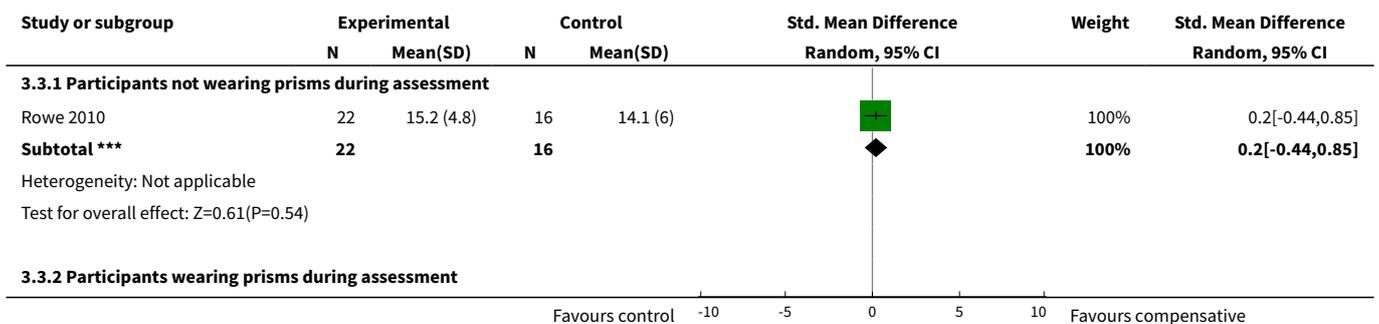
Analysis 3.1. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 1 Functional Activities of Daily Living.

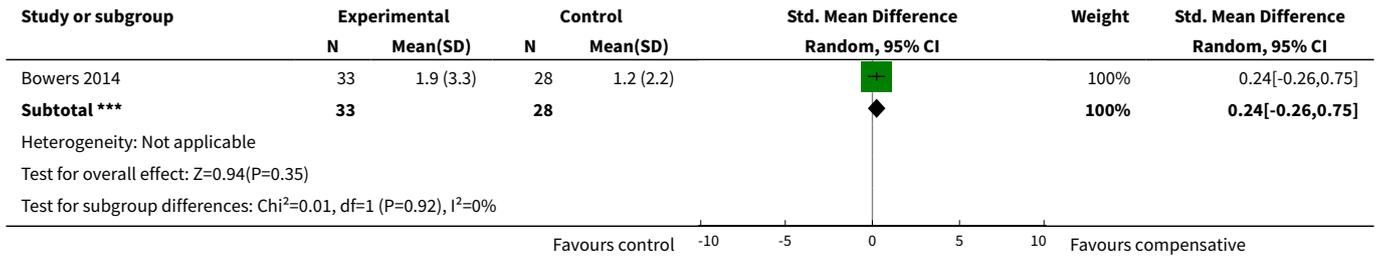


Analysis 3.2. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 2 Visual field.

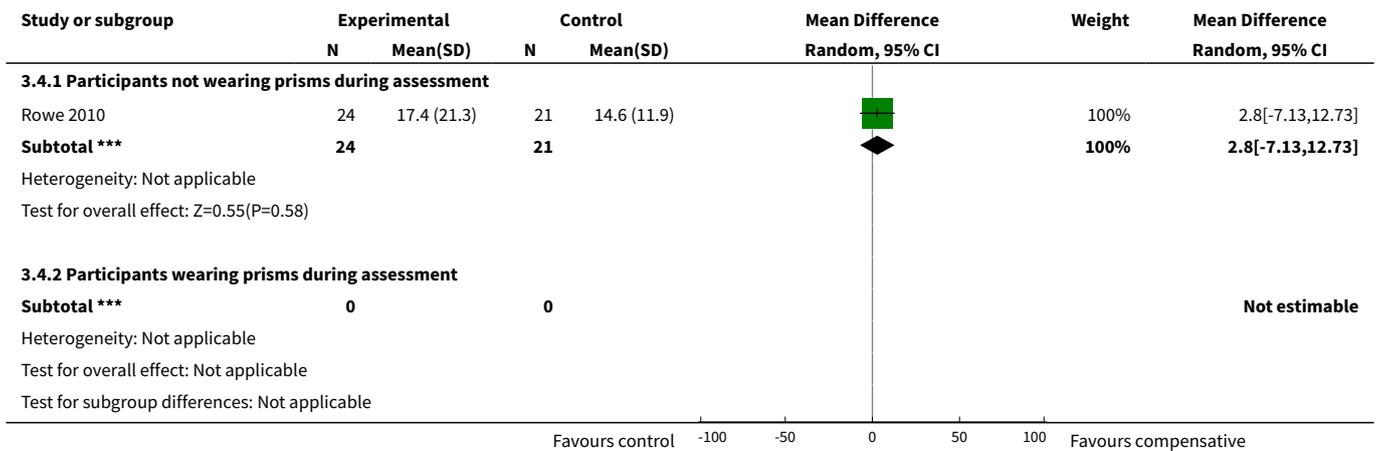


Analysis 3.3. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 3 Extended activities of daily living.

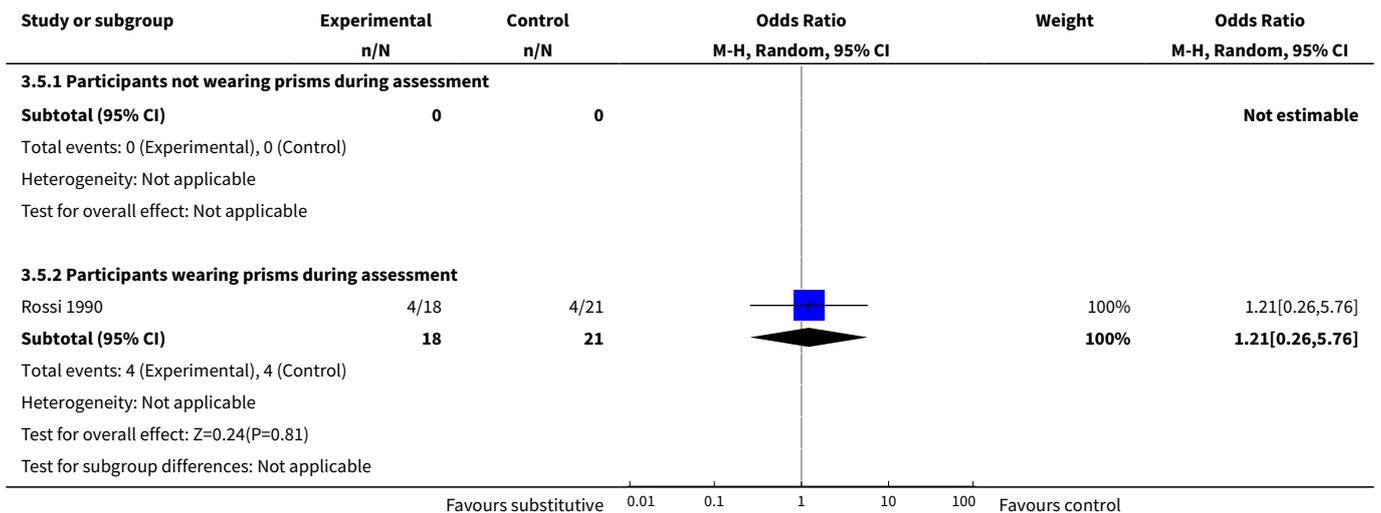




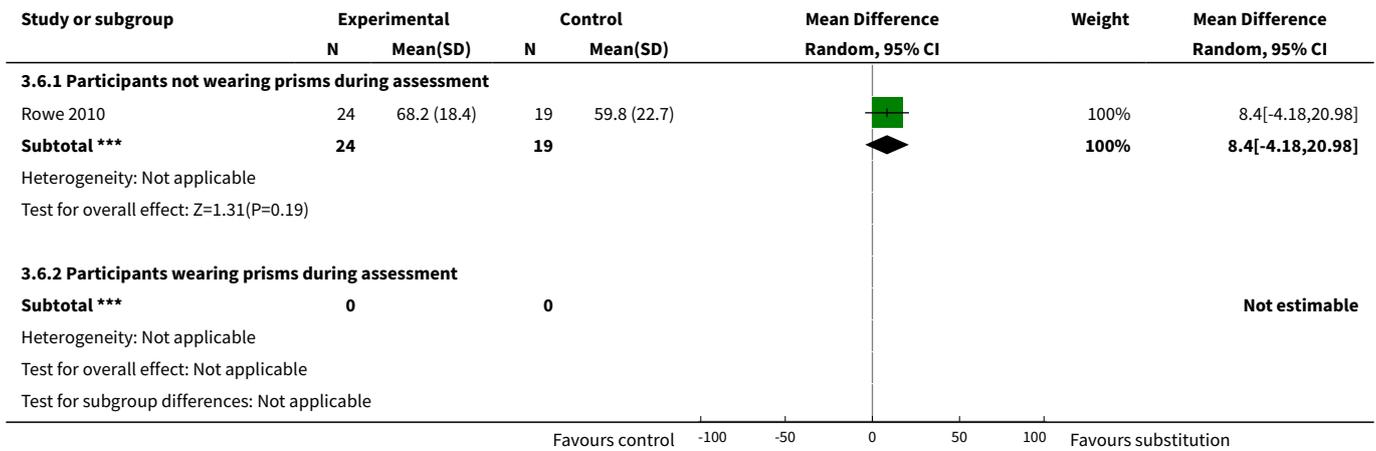
Analysis 3.4. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 4 Reading.



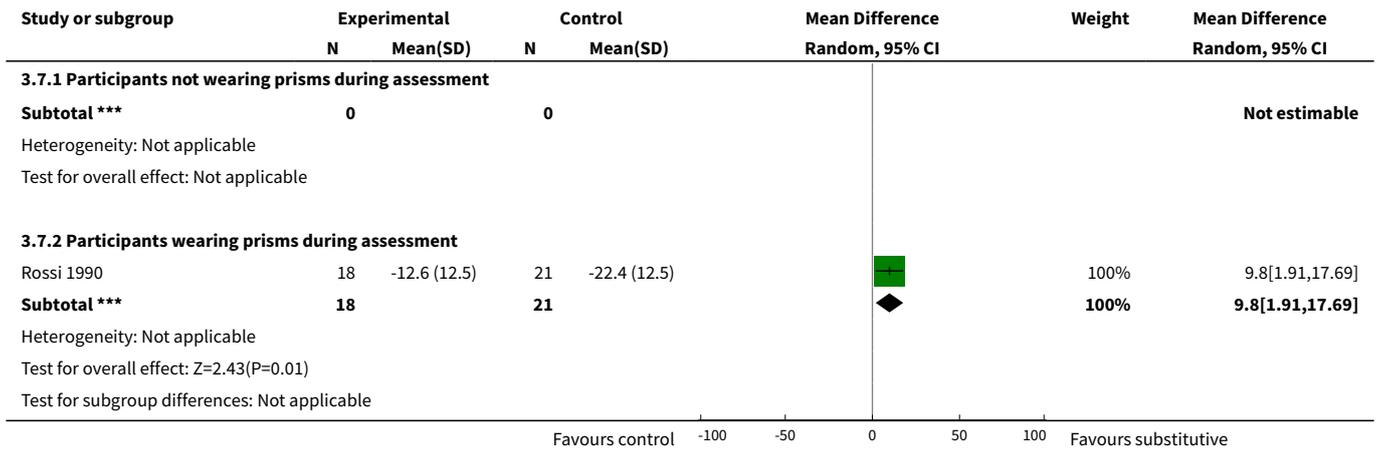
Analysis 3.5. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 5 Falls.



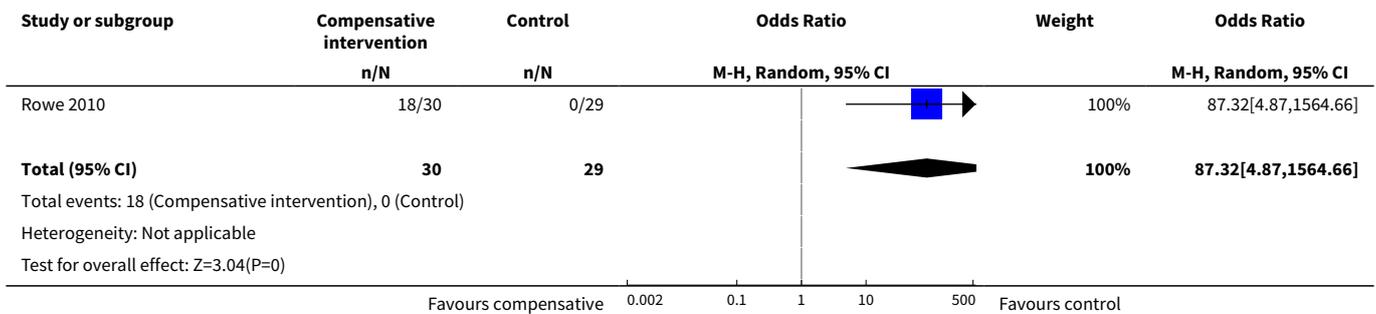
Analysis 3.6. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 6 Quality of life.



Analysis 3.7. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 7 Scanning - cancellation.



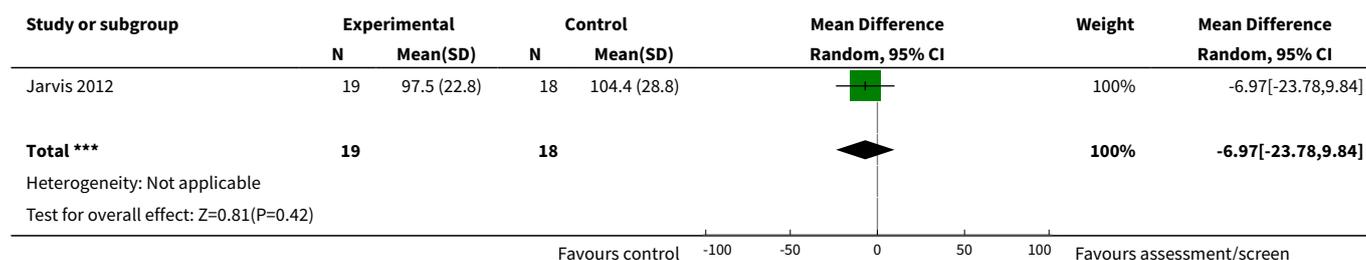
Analysis 3.8. Comparison 3 Substitutive interventions versus control, placebo or no intervention, Outcome 8 Adverse events.



Comparison 4. Assessment or screening versus control, placebo or no intervention

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 ADL	1	37	Mean Difference (IV, Random, 95% CI)	-6.97 [-23.78, 9.84]

Analysis 4.1. Comparison 4 Assessment or screening versus control, placebo or no intervention, Outcome 1 ADL.



ADDITIONAL TABLES

Table 1. Demographics of included studies: settings of included studies

Study	Country	Number of centres	Setting for intervention	Trial registration
Aimola 2011	UK	Multicentre ("from local hospitals or as self-referrals")	Community (participants' own homes)	UK Clinical Research Network Portfolio (UKCRN, ID 7144)
Bainbridge 1994	USA	Single	NS	NS
Bowers 2014	UK, USA	Multicentre (13 study sites)	University, hospital, private practice for fitting of prisms Then use at home (participants' own homes)	clinicaltrials.gov NCT00494676
De Haan 2015	Netherlands	2 ("Royal Dutch Visio and Bartiméus, the two centers of expertise for blind and partially sighted people in the Netherlands")	Training ... "was provided in Dutch at nine locations of Royal Dutch Visio and one location of Bartiméus in the Netherlands". Participants were also given homework assignments.	ISRCTN Registry ISRCTN16833414
Elshout 2016	Netherlands	Unclear ("Patients throughout the Netherlands could sign up for our study voluntarily by filling in a form on our website")	Community (participants' own homes)	NS

Table 1. Demographics of included studies: settings of included studies (Continued)

Gall 2013	Not clear	NS	NS	NS
Jarvis 2012	UK	Single	Stroke unit, Warring and Halton Hospitals, NHS Foundation Trust	NS
Jobke 2009	Germany	NS	NS	NS
Kasten 1998	Germany	NS	Community (participants' own homes)	NS
Kasten 2007	Germany	NS	Community (participants' own homes)	NS
Keller 2010	Germany	Single	Neurological clinic	NS
Modden 2012	Germany	Single	Rehabilitation centre (inpatients)	NS
Plow 2010	USA	Single	Outpatient (University clinic)	clinical- trials.gov NCT00921427
Poggel 2004	Germany	Single	Community (participants' own homes)	NS
Rossi 1990	USA	Single	Rehabilitation (inpatient)	NS
Roth 2009	Germany	NS	Community (participants' own homes)	NS
Rowe 2010	UK	Multicentre ("from stroke units based in 15 United Kingdom (UK) National Health Service (NHS) trusts")	Any (hospital, community)	Current Con- trolled Trials ISRCTN05956042.
Schuett 2012	Unclear. Authors from Austria, UK and Germany. "All participants were native German speakers."	NS	NS	NS
Spitzyna 2007	UK	NS	Community (participants' own homes)	NS
Szlyk 2005	USA	Single; university	Outpatient clinic	NS

NHS: National Health Service
NS: not stated
UK: United Kingdom
USA: United States of America

Table 2. Demographics of included studies: demographics of included participants

Study	Number of participants	Age	Gender	Time since stroke/lesion	Initial functional ability	Type of stroke/lesion	Side of stroke/lesion
Aimola 2011	70 participants recruited, 52 participants included in analyses	Group 1 Mean 61.4 years, SD 10.3	Group 1 9 F	NS	NS	Group 1 19 ischaemic stroke	Side of field defect
		Group 2 Mean 63.0 years, SD 10.9	Group 2 19 M 7 F 17 M			4 haemorrhagic, 4 traumatic brain injury 1 tumour	Group 1 L 15/R 13 Group 2 L11/R 13
Bainbridge 1994	18	NS	NS	NS	NS	NS	NS
Bowers 2014	73 randomised; 67 completed first phase (before cross-over); 61 completed second phase (after cross-over)	For 61 participants included after the cross-over: median 58 years (range 18 to 89)	For 61 participants included after the cross-over: M 40 F 21	For 61 participants included after the cross-over: median 18 months (range 3 to 396)	Overall baseline mobility difficulty, for 61 participants included after the cross-over: mean -0.17 (SD 2.31) logits for n = 31 using oblique prisms	For 61 participants included after the cross-over: hemianopia was caused by stroke for 47 (77%)	For 61 participants included after the cross-over: L hemianopia 39 (64%)

Table 2. Demographics of included studies: demographics of included participants (Continued)

					mean -0.06 (SD 1.89) logits for n = 30 using horizontal prisms		
De Haan 2015	54 randomised; data from 49 analysed (training group n = 26, control group n = 23)	Training group 55 ± 10.1 years Control group 57 ± 13.0 years	M 32 F 17	Training group 18 ± 22.5 months Control group 22 ± 24.6 months	NS	Ischaemic CVA 36 Haemorrhagic CVA 5 Traumatic brain injury 3 Penetrating head trauma 1 AVM extirpation 1 combined 3	L hemianopia 33 R hemianopia 16
Elshout 2016	40 recruited; data presented from first 3 cohorts of 10 only (n = 30); data from 27 analysed	Mean age 51.2 years (range 29 to 74)	M 22 F 5	Mean 26.3 months (range 11 to 111)	NS	5 haemorrhagic stroke 22 ischaemic stroke	L-sided field defect 14 R-sided field defect 13
Gall 2013	39 (alternating current stimulation n = 15, sham n = 14)	NS	NS	NS	NS	NS ("patients with post-chiasmatic visual pathway lesions")	NS
Jarvis 2012	64 randomised (experimental group n = 33, control n = 31)	Experimental: mean 70.4 years (SD 10.8) Control: mean 69.4 years (SD 14.5)	M 40 F 24	NS	NS	Ischaemic 56 Haemorrhage 7 Combined 1	R-sided stroke 41 L-sided stroke 19 Bilateral 4
Jobke 2009	21	Group 1 Mean 51.5 years, SD 14.8	Group 1 M 7	Group 1 Mean 89.0 months, SD 59.9	NS	Group 1 5 stroke/ischaemia	NS

Table 2. Demographics of included studies: demographics of included participants (Continued)

		Group 2	F 1	(range 67 to 225 months)		1 brain injury	
		Mean 47.3 years, SD 13.4	Group 2	Group 2		1 tumour	
			M 6	Mean 89.4, SD 57.6		1 surgery	
			F 4	(range 40 to 236 months)		Group 2	
						5 stroke/ischaemia	
						1 meningitis;	
						1 injury	
						3 surgery	
Kasten 1998	19	Group 1	Group 1	Group 1	NS	19 participants with post chiasmal injury; 10 were due to stroke, 4 due to trauma and 5 due to other reasons	NS
	(plus 19 with pre-chiasmal damage)	? Mean 47.7 years, ? SD 12.9	M 11	? Mean 6.8 months, ? SD 11.4			
		Group 2	F 8	Group 2			
	Data are presented for full group of 38	? Mean 55.3 years, ? SD 16.2	Group 2	? Mean 7.2 months, ? SD 6.3			
		It is assumed the data presented are mean and SD, but this was not stated	M 13	It is assumed the data presented are mean and SD, but this was not stated			
			F 6				
Kasten 2007	23	Group 1	Group 1	Group 1	NS	Group 1	NS
		Mean 41.1 years, SD 16.9	M 5	10 to 83 months, Mean 34.2*, SD 30.1*		4 stroke	
		Group 2	F 2	Group 2		1 trauma	
		Mean 39.3 years, SD 10.9	Group 2	13 to 477 months, Mean 92.7*, SD 170.6*		1 cerebral aneurysmal bleeding	
		Group 3	M 6	Group 3		1 hypoxia	
		Mean 44.3 years, SD 9.1	F 1	Group 3		Group 2	
			Group 3	10 to 143 months, Mean 47.6*, SD 54.4*		3 stroke	
			M 6			3 trauma	
			F 3			1 surgery	
						Group 3	
						3 stroke	

Table 2. Demographics of included studies: demographics of included participants (Continued)

						2 trauma	
						1 surgery	
						1 hypoxia	
						2 other	
Keller 2010	20	Group 1 Mean 54.7 years, SD 20.4	Group 1 M 6	Group 1 Mean 8.5 weeks, SD 6.7	NS	Group 1 9 vascular	Group 1 4 left hemi-anopia
		Group 2 Mean 63.6 years, SD 13.8	F 4	Group 2 Mean 4.2 weeks, SD 2.1		1 tumour	3 right hemi-anopia
			Group 2 M 6			Group 2 9 vascular	1 UL quadrantanopia
			F 4			1 traumatic	1 LL quadrantanopia
							1 UR quadrantanopia
							Group 2
							3 left hemi-anopia
							3 right hemi-anopia
							3 UL quadrantanopia
							1 LL quadrantanopia
Modden 2012	45	RT Group: Mean 58.3 ± 11.4 years	RT group: M 10	RT group: Mean 4.7 weeks	NS	RT Group: occipital 7	RT group: L stroke 7
		CT group: Mean 57.1 ± 8.3 years	F 5	CT group: Mean 4.9 weeks		temporo-occipital 2	R stroke 8
		OT group: Mean 59.0 ± 11.1 years	CT group: M 9	OT group:		temporomedial 5	CT group: L stroke 5
						parahippocampal 1	

Table 2. Demographics of included studies: demographics of included participants (Continued)

			F 6	Mean 4.3 weeks		CT Group:	R stroke 10
			OT group:	"Patients were recruited on average about 4 weeks after their stroke."		occipital 6	OT group:
			M 7			temporo-occipital 3	L stroke 5
			F 8			temporomedial 5	R stroke 10
						parahippocampal 1	
						OT Group*:	
						occipital 4	
						temporo-occipital 3	
						temporomedial 5	
						parahippocampal 1	
						*numbers presented in paper do not add up to 15 (?)	
Plow 2010	12	Mean 59.6 years, SEM 3.5 years	M 5 F 7	Mean 39.8 ± 16.2 months, range 3 to 192 months	NS	Stroke 8 (7 infarct, 1 haemorrhage) Surgical trauma 2	L-affected side 4 R-affected side 8
Poggel 2004	20 participants recruited. Baseline data only available for 19 (data for one dropout not reported)	Group 1 Mean 41.9 years Range 20 to 67 years Group 2 Mean 43.2 years Range 30 to 61 years	Group 1 M 6 F 3 Group 2 M 6 F 4	Group 1 Mean 49.1 months, SEM ?, Range 6.7 to 189.9 months Group 2 Mean 24.1 months, SEM 5.0, Range 6.8 to 58.3 months	NS	Group 1 vascular 1 infarct 8 cortical and radiations 4 cortical 5 Group 2 vascular 2 infarct 7, traumatic brain injury 1 cortical and radiation 5	Group 1 L 5/R 4 Group 2 L 5/R 5

Table 2. Demographics of included studies: demographics of included participants (Continued)

						cortical 3 radiation 2	
Rossi 1990	39	Group 1 Mean 72.6 years, SEM 1.8 Group 2 Mean 63.3 years, SEM 2.5	Group 1 M 10 F 8 Group 2 M 9 F 12	Group 1 Mean 4.4 weeks, SEM 0.3 Group 2 Mean 4.7 weeks, SEM 0.6	NS	Group 1 15 infarct 3 haemorrhage Group 2 18 infarct 3 haemorrhage	Group 1 16 R/2 L Group 2 13 R/8 L
Roth 2009	30 participants recruited (data available for 28; 2 dropouts)	Group 1 Mean 60.5 years, SD 11.0, Median 65 Group 2 Mean 60.3 years, SD 11.7, Median 63	Group 1 4 F 11 M Group 2 F 7 M 8	Group 1 Mean 39.20 months, SD 54.59, Median 26 Group 2 Mean 87.87 months, SD 186.66, Median 16	NS	Group 1 Stroke 11 Haemorrhage 1 Head injury 1 Abscess 1 AVM 1 Group 2 Stroke 11 Haemorrhage 3 Cyst 1	Affected side Group 1 L 8/R 7 Group 2 L 7/R 8
Rowe 2010	87 participants recruited (full results for 70 participants at 26 weeks)	Group 1 Mean 69.9 years, SD 12.9, median 68.8, IQR, 14.4 Group 2 Mean 70.9 years, SD 11.2, median 72.9, IQR, 15.2 Group 3 Mean 66.2 years, SD 11.3, median 68.2, IQR, 16.2	Group 1 4 F 22 M Group 2 13 F 17 M Group 3 9 F 20 M	Group 1 Mean 75.5 days, SD 45.3, median 64.5, IQR 78.0 Group 2 Mean 73.8 days, SD 49.2, median 69.0, IQR 97.0 Group 3 Mean 81.2 days, SD 48.0, median 67.0, IQR 61.0	Barthel Index score Group 1 Mean 97.5, SD 5.5, median 100.0, IQR 0.0 Group 2 Mean 92.7, SD 11.9, median 100.0, IQR 15.0 Group 3	Group 1 25 ischaemic 1 haemorrhage Group 2 28 ischaemic 2 haemorrhage Group 3 28 ischaemic 1 haemorrhage	Side of infarct Group 1 L 9/R 16/bilateral 1 Group 2 L 17/R 13/bilateral 0 Group 3 L 11/R 17/bilateral 1

Table 2. Demographics of included studies: demographics of included participants (Continued)

					Mean 93.3, SD 14.7, me- dian 100.0, IQR 5.0		
Schuett 2012	36	Group 1 Mean 64.0 years, SD 11.1, range 44 to 81 Group 2 Mean 63.7 years, SD 13.3, range 42 to 83	Group 1 3 F 15 M Group 2 3 F 15 M	Group 1 Mean 26.6 weeks, SD 14.5, range 6 to 57 Group 2 Mean 20.1 weeks, SD 18.8, range 4 to 74	NS	Group 1 17 posterior infarc- tion 1 tumour operation Group 1 17 posterior infarc- tion 1 tumour operation	Side of field loss Group 1 L 9/R 9 Group 2 L 7/R 11
Spitzyna 2007	22	Age at symptom onset Group 1 Range 5 to 67 years, mean 42.5*, SD 20.5* Group 2 Range 39 to 78 years, mean 63.1*, SD 12.2*	Group 1 M 6 F 5 Group 2 M 7 F 1	Time since symptoms onset Group 1 Range 1 to 37 years, mean 7.5*, SD 10.9* Group 2 Range 3 months to 5 years, mean 1.6*, SD 1.7*	NS	Group 1 3 infarct 1 tuberous sclerosis 2 traumatic brain in- jury, 2 tumour 2 haemorrhage 1 cyst Group 2 8 infarct	Group 1 All R Group 2 All R
Szlyk 2005	10	Group 1 Range 16 to 74 years, mean 50.6*, SD 22.5* Group 2 Range 34 to 73 years, mean 54.0, SD 14.4	Group 1 5 M Group 2 5 M	NS	NS	Group 1 4 CVA 1 tumour: all occipi- tal lobe Group 2 4 CVA 1 AVM: all occipital lobe	Group 1 L 3/R 2 Group 2 L 4/R 1

* Figures calculated from raw data supplied in papers

AVM: arteriovenous malformation

CVA: cerebrovascular accident

CT: compensatory training

F: female

IQR: interquartile range

L: left

LL: lower left

M: male

NS: not stated

OT: occupational therapy

R: right

RT: restitutive training

SD: standard deviation

SEM: standard error of the mean

UL: upper left

UR: upper right

Table 3. Demographics of included studies: visual problems of included participants

Study	Methods of visual field assessment	Type/extent of field loss	Macular sparing	Presence of neglect?
Aimola 2011	Unspecified kinetic perimeter Esterman measures of static superthreshold	Group 1 Hemianopia 20, quadrantanopia 8 Group 2 Hemianopia 20, quadrantanopia 4	Group 1 Mean 1.92° (SD 1.44) Group 2 Mean 2.45° (SD 1.85)	Yes: stated "Three patients (2 in the intervention group, 1 in the control group) had comorbid neglect as confirmed with the bells test".
Bainbridge 1994	Harrington Flocks Visual Screener Confrontation	Not stated	Not stated	Yes: no details of inclusion criteria or participants provided, but objective stated "To study the effect of ... on visual neglect or hemianopsia following stroke".
Bowers 2014	Goldmann perimetry	Not stated	Not stated	No: stated "no visual neglect". Visual neglect diagnosed with Bells test and Schenkenberg Line Bisection Test.
De Haan 2015	Goldmann perimetry	Training group Functional field score 58 ± 7.8 Quadrantanopia 5 (3 lower left, 1 upper left, 1 lower right) Hemianopia 21 Control group Functional field score 64 ± 11.4 Quadrantanopia 5 (3 lower left, 2 upper left) Hemianopia 18	Not stated	No: stated "Neglect was excluded based on the Balloons, drawings, Line Bisection and Rey Complex Figure Test."
Elshout 2016	Goldman perimetry Humphrey perimetry	Right field loss: hemifield 4, incomplete hemifield 5, quadrant 2, scotoma 1 Left field loss: hemifield 2, incomplete hemifield 9, quadrant 1, scotoma 2 Bilateral field loss Incomplete: 1	"All subjects had macular sparing of at least 2°"	No: patients with visual neglect were excluded (based on line bisection test)
Gall 2013	Standard automated perimetry	Not stated	Not stated	Not stated

Table 3. Demographics of included studies: visual problems of included participants (Continued)

Jarvis 2012	Confrontation	Ocular diagnosis: low vision 30 visual field loss 38 eye movement deficit 41 perceptual impairment 24 ("Note: patients may have had an isolated visual impairment or combined visual deficits")	Not stated	Yes: all patients with a "post-stroke visual impairment were eligible for inclusion".
Jobke 2009	Standard automated perimetry High resolution perimetry (HRP)	NB: It did not state whether participants had visual neglect or whether this was diagnosed. Group 1 2 diffuse, 2 full homonymous hemianopia, 1 partial homonymous hemianopia, 1 full quadrantanopia 2 partial quadrantanopia Group 2 4 diffuse, 2 full homonymous hemianopia, 2 partial homonymous hemianopia, 1 full quadrantanopia, 1 partial quadrantanopia	Group 1 7 sparing, 1 not sparing Group 2 10 sparing	Not stated
Kasten 1998	Tubinger automated perimetry (TAP) High resolution perimetry (HRP)	NB: data were presented for full group of 38 participants (including participants in parallel trial) Group 1 TAP 90° - border position, mean 3.51° (degrees of visual angle from zero vertical meridian), SEM 1.0 TAP 90° - number of misses, mean 53.0, SEM 9.1 Group 2 TAP 90° - border position, mean 3.43° (degrees of visual angle from zero vertical meridian), SEM 0.99 TAP 90° - number of misses, mean 69.2, SEM 11.2	Not stated	No: participants with neglect were excluded. Method of diagnosis of neglect not stated.
Kasten 2007	Tubinger automated perimetry (TAP) High resolution perimetry (HRP)	TAP 90° (number of blind stimuli positions) Group 1 Right eye - mean 46.6, SD 6.9, left eye - mean 43.9, SD 3.7 Group 2 Right eye - mean 50.3, SD 8.7, left eye - mean 43.1, SD 7.6 Group 3 Right eye - mean 32.9, SD 6.8, left eye - mean 37.9, SD 7.1	Not stated	No: participants with neglect were excluded. Method of diagnosis of neglect not stated

Table 3. Demographics of included studies: visual problems of included participants (Continued)

.Keller 2010	Goldmann perimetry Goldmann suprathresh- old	Group 1 4 left hemianopia 3 right hemianopia 1 UL quadrantanopia 1 LL quadrantanopia 1 UR quadrantanopia Group 2 3 left hemianopia 3 right hemianopia 3 UL quadrantanopia 1 LL quadrantanopia	Group 1 6 with 0° macular sparing 4 with < 5° macular sparing Group 2 6 with 0° macular sparing 4 with < 5° macular sparing	No: participants with neglect were excluded. 3 neglect tests were used: "line bisection, Mesu- lam test, draw a clock face test".
Modden 2012	Visual field assessment from the Test Battery of Attentional Performance	RT Group 10 hemianopia 5 quadrantanopia TAP alertness without cueing, ms; mean 304.2, SD 80.8 TAP conjunction search, omissions; mean 9.1, SD 9.0 CT Group 12 hemianopia 3 quadrantanopia TAP alertness without cueing, ms; mean 383.7, SD 205.2 TAP conjunction search, omissions; mean 10.7, SD 6.7 OT Group 10 hemianopia 5 quadrantanopia TAP alertness without cueing, ms; mean 308.1, SD 58.6 TAP conjunction search, omissions; mean 10.3, SD 5.6	RT Group 3/15 par- ticipants with less than 2° sparing CT Group 3/15 par- ticipants with less than 2° sparing plus 1 par- ticipant with no sparing OT Group 3/15 par- ticipants with less than 2° sparing	No: participants with neglect were excluded. Method of diag- nosis of neglect not stated.
Plow 2010	Subjective topographic measure of perceived vi- sual field de- fect	7 hemianopia 5 quadrantanopia	Not stated	Not stated

Table 3. Demographics of included studies: visual problems of included participants (Continued)

	High resolution perimetry (HRP)			
Poggel 2004	Tubinger automated perimetry (TAP)	Group 1 Upper attention field (size of area of residual vision, %), mean 18.2, SEM 4.0	Not stated	No: participants with neglect were excluded. Method of diagnosis of neglect not stated.
	High resolution campimetry	Lower probe field (size of area of residual vision, %), mean 21.3, SEM 3.1		
	High resolution perimetry (HRP)	Total visual field (size of area of residual vision, %), mean 7.3, SEM 1.9		
		Group 2 Upper attention field (size of area of residual vision, %), mean 16.9, SEM 2.4		
		Lower probe field (size of area of residual vision, %), mean 15.5, SEM 4.0		
		Total visual field (size of area of residual vision, %), mean 6.7, SEM 1.3		
Rossi 1990	Harrington Flocks Visual Screener	Group 1 Homonymous hemianopia 12 (Visual neglect 6)	Not stated	Yes: participants with "homonymous hemianopia or visual neglect were recruited". Method of diagnosis of neglect was Harrington Flocks Visual Screener.
	Tangent screen measures	Group 2 Homonymous hemianopia 15 (Visual neglect 6)		
Roth 2009	Tubinger automated perimetry (TAP)	Group 1 Homonymous hemianopia 12, quadrantanopia 3	Not stated	No: participants with neglect were excluded. Method of diagnosis of neglect was clock-drawing and line-bisection tests.
	Scanning laser ophthalmoscopy	Group 2 Homonymous hemianopia 12, quadrantanopia 3		
Rowe 2010	Goldmann perimetry	Group 1 Homonymous hemianopia left partial 8, Homonymous hemianopia right partial 3, Homonymous hemianopia left complete 9, Homonymous hemianopia right complete 6	Not stated	No: participants with neglect were excluded. Method of diagnosis was clinical assessment: "as assessed by the orthoptist".
	Esterman measures of static superthreshold	Group 2 Homonymous hemianopia left partial 5, Homonymous hemianopia right partial 9, Homonymous hemianopia left complete 8, Homonymous hemianopia right complete 8		
		Group 3		

Table 3. Demographics of included studies: visual problems of included participants (Continued)

Homonymous hemianopia left partial 8, Homonymous hemianopia right partial 5, Homonymous hemianopia left complete 10, Homonymous hemianopia right complete 6

Schuett 2012	Tübingen kinetic perimetry	Group 1	Group 1	No: participants with neglect were excluded. Method of diagnosis described as: "as assessed by tests in accordance with the Behavioural Inattention Test (line bisection, letter and star cancellation, figure and shape copying, drawing from memory; Halligan et al, 1991)."
		Hemianopia 15, quadrantanopia 1, paracentral scotoma 2	Mean 2.3° (SD 1.4)	
		Group 2	Group 2	
		Hemianopia 10, quadrantanopia 4, paracentral scotoma 4	Mean 2.3° (SD 1.2)	
Spitzyna 2007	Goldmann perimetry	Group 1	Macular sparing defined as 2° of sparing	No: only participants with right-sided homonymous hemianopic were included; therefore, presence of neglect was assumed unlikely.
	Humphrey automated perimetry	Full homonymous hemianopia 8, partial homonymous hemianopia 1, lower quadrantanopia 1, upper quadrantanopia 1		
		Group 2	Group 1	
		Full homonymous hemianopia 6, lower quadrantanopia 1, upper quadrantanopia 1	Sparing 5, non-sparing 6	
			Group 2	
			Sparing 3, non sparing 5	
Szlyk 2005	Goldmann perimetry	Group 1	Not stated	Not stated; however, although it was not stated whether the participants may have had visual neglect, neglect is unlikely in occipital lesions, and only participants with occipital lesions were included.
		Goldmann III4e, range 45.2 to 125, mean 59.12*, SD 22.07*		
		Goldmann V4e, range 48.8 to 115, mean 70.56*, SD 26.15*		
		Group 2		
		Goldmann III4e, range 46.8 to 123.8, mean 68.0*, SD 31.71*		
		Goldmann V4e, range 50.67 to 132, mean 73.73*, SD 33.10*		
		Figures were calculated for the affected side only.		

* Figures calculated from raw data supplied in papers

combined: combined etiology

HRP: high resolution perimetry

LL: lower left

LR: lower right

ms: milliseconds

SD: standard deviation

SEM: standard error of the mean

TAP: Tübingen automated perimeter

UL: upper left

UR: upper right

Table 4. Outcome measures within included studies

Study	Visual field Outcome category (measure)	Functional reading ability in EADL	Quality of life	Visual scanning	Ad-Other verse events	Outcomes with data included within meta-analyses
Aimola 2011	Kinetic Perimetry (unspecified kinetic perimeter) Static Superthreshold (Esterman measures of static superthreshold) (NB not clear if recorded as outcome or not; no results provided for visual field data)	Reading (corrected reading speed)	1. VFQ 25 2. VIQ - Visual Impairments questionnaire 3. Subjective Reasons questionnaire	1. visual search - find the number (computer-based) 2. visuomotor search - find items on a shelf	Tasks simulating ADL - 1. driving hazard perception (mean score per hazard), 2. obstacle avoidance (completion time), 3. visuomotor search (time) Attention tasks - 1. sustained attention to response (mean percentage error score), 2. test of everyday attention	Reading: Analysis 2.3 Visual search: time to complete Analysis 2.5 QoL: data not included as only available for individual questionnaire items
Bainbridge 1994	Gross visual screening (Harrington-Flocks Visual Field Score)			Line Cancellation Test	Motor Free Visual Perception Score Line Bisection Test	No data included in meta-analyses (as no control group). See Table 5
Bowers 2014		Mobility questionnaire			Question: "If the study were to end today, would you want to continue with these prism glasses (i.e. the prism glasses worn in that period)?"	Functional ability in EADL: Analysis 3.3
DeHaan 2015	Kinetic perimetry (Goldmann Perimetry, Functional Field score)	In- 1. Radner reading test; Mobility - Radner average reading	1. NEI-VFQ-25 (Visual Functional questionnaire) 2. Cerebral Vi-	1. visual scanning - dots test 2. visual search - letters (parallel search test) 3. visual search - let-	NotVisual acuity, contrast sensitivity, hazard perception, simulating driving/tracking task, obstacle course as an outcome measure,	Visual field: Functional Field score Analysis 2.1 Reading: Radner average reading speed Analysis 2.3 Visual scanning: Parallel search test, time Analysis 2.5 QoL: NEI VFQ Analysis 2.4

Table 4. Outcome measures within included studies (Continued)

	<p>toring na speed (wpm), (b) min- imal read- able text size (Lo- gRad)</p> <p>2. Text read- ing test; (a) text read- ing speed (wpm), (b) text correct an- swers</p>	<p>sual Dis- orders ques- tionnaire</p>	<p>ters (serial search)</p>	<p>but stat- ed no ad- verse events in ei- ther group</p>	<p>Functional ability in EADL: mobility questionnaire Analysis 2.2</p>
<p>Elshout 2016</p>	<p>Goldman perimetry Humphrey perimetry</p>	<p>Read- ing speed (words per minute) - 15 point Arial font, 88 and 165 words</p>			<p>No data included (as data not available for before the cross-over)</p>
<p>Gall 2013</p>	<p>Static Threshold Perimetry (Standard automated perimetry)</p>		<p>1. NEI VFQ 39 (vi- sion-re- lated)</p>		<p>No data included in meta-analyses (as no suitable data presented in abstract)</p>

Table 4. Outcome measures within included studies (Continued)

				2. SF-12 (health-related)		
Jarvi 2012	FIM				1. Functional mobility (timed walk) 2. Non-validated questionnaire giving qualitative information about their treatment approach	Functional ability in ADL: Functional Independence Measure Analysis 4.1
Jobke 2009	Static Threshold Perimetry (Standard automated perimetry) Resolution Perimetry (High resolution perimetry)	Radner reading test	NEI VFQ		Zahlen-Verbindungs test (ZVT) for measuring the speed of connecting numbers in a paper-pencil test	No data included in meta-analyses (as no control group). See Table 5 (for data available before the cross-over)
Kas- ten 1998	Resolution Perimetry (High resolution perimetry) Static Threshold Perimetry (Tubinger automated perimetry)		Quality of life questionnaire		Visual acuity: Landolt ring to give minimum angle of resolution Visual acuity: Landolt ring to give minimum angle of resolution	Visual field: Tubinger automated perimetry: border position in degrees of visual angle from zero vertical meridian Analysis 1.1 Quality of life Analysis 1.2
Kas- ten 2007	Resolution Perimetry (High resolution perimetry: number of hits, learning effects, fixation ability, false hits) Static Threshold Perimetry (Tubinger automated perimetry: no of hits, fixation ability)		Subjective visual ability questionnaire		1. Eye movements: "Chronos Vision Eye Tracker" 2. Visual acuity 3. "Zahlen-Verbindungs Test" of visuo-spatial attention 4. "Alters-Konzentrationstest" attention test for older people 5. "testbatterie zur Aufmerksamkeitspruefung" ability to improve attention	No data included in meta-analyses (as no control group). See Table 5 (for available data comparing group outcomes)
Keller 2010	Kinetic Perimetry (Goldmann perimetry) Static Superthreshold (Goldmann suprathreshold)	Reading time (standardised)	OT administered questionnaire (based	1. Visual exploration test (number of omissions)	Electro-oculography	No data included in meta-analyses (as no control group). See Table 5

Table 4. Outcome measures within included studies (Continued)

	read- ing test)	on Ker- hoff's self-eval- uation of ADL)	2. Search task (search time)		
Modden 2012	Gross Visual Screening (Test Battery of Attentional Performance: visual field assessment)	Reading - Wechsler memo- ry tests (errors)	1. Visual scan: from the Test Bat- tery of At- tentional Per- formance 2. Visual search: can- cellation tasks from Behavioural Inattention Test (BIT) (omissions)	Attention: Test Battery of Attentional Performance (alertness)	No data included in meta-analyses (as no control group). See Table 5
Plow 2010	Resolution Perimetry (High resolution perimetry: position of visual field border and stimulus detection accuracy) Gross Visual Screening (subjective topographic measure of perceived visual field deficit)	Impact of Vision Impair- ment (IVI) pro- file Low Vi- sion- Visual Func- tional Ques- tionnaire (LV-VFQ)		Measure of fixation perfor- mance	No data included in meta-analyses (as no control group). See Table 5
Poggel 2004	Static Threshold Perimetry (Tubinger automated perimetry (TAP)) Static Superthreshold (High resolution campimetry)				No data included in meta-analyses (as no control group). See Table 5

Table 4. Outcome measures within included studies (Continued)

	Resolution Perimetry (High resolution perimetry (HRP))					
Ross 2010	Barbur Visual Screening (Harrington InFlocks Visual Screener)		Number of falls	Line cancellation task	Modified Mini Mental Status Examination, Motor Free Visual Perceptual Test, Line Bisection Task	ADL: Barthel Index: Analysis 3.1 Visual Field: Analysis 3.2 Falls: number of falls Analysis 3.5 Visual scanning: cancellation Analysis 3.7
Roth 2009	Static Threshold Perimetry (Tubinger automated perimetry (TAP)) Resolution Perimetry (Scanning laser ophthalmoscopy)	Reading speed	QoL: World Health Organisation questionnaire WHO-QOL-BREF	1. Digit search task (response time) 2. Natural search task (table test) (response time) 3. Natural scene exploration 4. Fixation stability (video eye tracker)		No data included in meta-analyses (as no control group). See Table 5
Rowe 2010	Kinetic Perimetry (Goldmann perimetry) Static Superthreshold (Esterman measures of static superthreshold)	NotReading ability ex- (Rad-tender ed test) ac-tiv-i-ties of dai-ly liv-	1. VFQ 25-10 (vi-sion re-lated) 2. EQ-5D 3. SF-12		NurRivermead Mobility Index ber of par-tic-i-pants and num-ber of ad-verse events	Visual Field: relative change in visual field area Analysis 2.1 and Analysis 3.2 QoL: VFQ 25-10 Analysis 2.4 and Analysis 3.6 Adverse events: Analysis 2.6 ; Analysis 3.8

Table 4. Outcome measures within included studies (Continued)

		ing (NEADL)		
Schuetz 2012	Kinetic Perimetry (Tubingen kinetic perimetry) (NB not recorded immediately after first phase)	Reading (speed and errors)	Cancellation (speed and errors)	No data included in meta-analyses (as no control group). See Table 5
Spitzyna 2007	Kinetic Perimetry (Goldmann perimetry) Static Threshold Perimetry (Humphrey 10-2 central threshold programme) (NB not recorded immediately after first phase)	1. Text reading speeds 2. Single word reading speeds	Eye movement characteristics: - spatial characteristics of saccadic amplitude, incoming saccade amplitude and landing position - temporal characteristics	Reading (text reading speed): Analysis 2.3 Visual field -data not included as not collected before cross-over.
Szlyk 2005	Kinetic Perimetry (Goldman Perimetry)		Indoor functional assessment Outdoor functional assessment Driving skills assessment Psychophysical assessment Satisfaction Prisms use at 2 years	No data included in meta-analyses(as no control group). No data reported for before the cross-over.

ADL: activities of daily living

BIT: behavioural inattention test

EADL: extended activities of daily living

EQ-5D: standardised EuroQol health-related quality of life instrument

FIM: Functional Independence Measure

HRP: high resolution perimetry

IVI: impact of vision impairment

LogRad: a scale of reading acuity

LV-VFQ: low vision functional questionnaire

NB: note

NEADL: Nottingham Extended Activities of Daily Living

NEI-VFQ-25: National Eye Institution Visual Function Questionnaire

OT: occupational therapy
QoL: quality of life
SF-12: 12-Item Short Form Health Survey
TAP: tubinger automated perimetry
VFQ: visual function questionnaire
VIQ: visual impairment questionnaire
WHOQOL-BREF: World Health Organisation quality of life questionnaire
wpm: words per minute
ZVT: Zahlen-Verbindungs test

Table 5. Results of studies comparing two similar active interventions (i.e. two interventions from the same category)

Study	Interventions	Outcome	Mean (or other reported result if no mean available)	Standard deviation	Number of participants	Statistical test/ results	
Restitution: one restitution intervention versus another restitution intervention							
Jobke 2009	Extrastriate VRT	Visual field (high-resolution perimetry, HRP)	increase from baseline of 5.9% (percentage of HRP hits)		8	significant increase: $t = -5.262$, $P = 0.0005$	
	Standard VRT		increase from baseline of 2.9% (percentage of HRP hits)		10	significant increase: $t = -2.373$, $P = 0.021$	
Kasten 2007	Parallel co-stimulation	Visual field (high-resolution perimetry)	increase of 2.4% detected stimuli		7	No significant difference "confirmed by non-parametric Kruskal-Wallis ANOVA"	
	Moving co-stimulation		increase of 6.5% detected stimuli		7		
	Single stimulus		increase of 3.9% detected stimuli		9		
Plow 2010	VRT + tDCS	Visual field (high-resolution perimetry)	shift from baseline to post-test from $4.11^\circ \pm 1.50^\circ$ to $8.37^\circ \pm 2.29^\circ$; Wilcoxon signed-rank test = 0, $P = 0.068$		4	Mann-Whitney $U = 0$, $P = 0.021$ (significantly greater shift in the visual field border with VRT + tDCS than VRT alone)	
	VRT + sham tDCS		shift from baseline to post-test from $6.33^\circ \pm 2.59^\circ$ to $7.03^\circ \pm 2.51^\circ$; Wilcoxon signed-rank test = 1, $P = 0.144$		4		
	VRT + tDCS	Functional ability in ADL (LV-VFQ)	shift from baseline to post-test from 32.25 ± 5.30 to 28.25 ± 5.07 ; Wilcoxon signed-rank test = 0; $P = 0.068$		4		Mann-Whitney $U = 5.5$; $P = 0.468$ (non-significant)
	VRT + sham tDCS		shift from baseline to post-test from 28 ± 2.34 to 25.25 ± 1.11 ; Wilcoxon signed-rank test = 1; $P = 0.285$		4		
	VRT + tDCS	Functional ability in ADL (LV-VFQ) - 6-month follow-up	29.00	3.58	5		Wilcoxon signed-rank test = 4; $P = 0.343$

Table 5. Results of studies comparing two similar active interventions (i.e. two interventions from the same category) (Continued)

	VRT + sham tD-CS		26.80	2.11		
	VRT + tD-CS	Quality of life - 6-month follow-up	23.20	7.83	5	Wilcoxon signed-rank test = 2.5; P = 0.357
	VRT + sham tD-CS		16.8	4.62		
Poggel 2004	VRT + attentional cueing	Visual field (high-resolution perimetry) - percentage improvement, attention field	8.3	SEM 1.5	9	P = 0.001 (in favour of attentional cueing)
	VRT with no attentional cueing		2.9	SEM 0.8	10	
Compensation: one compensation intervention versus another compensation intervention						
Schuett 2012	Visual exploration training	Reading speed	105.3	33.8	18	Not reported; calculated as MD -19.30 (-43.32 to 4.72) (see Figure 3)
	Reading training		124.6	39.5	18	
	Visual exploration training	Cancellation test (exploration time)	18.5	4.9	18	Not reported; calculated as MD 18.30 (14.28 to 22.32) (see Figure 3)
	Reading training		36.8	7.2	18	
Keller 2010	Audiovisual exploration training (AVT)	Functional ability in ADL (ADL test total score)	1.5	(SE displayed on graph only)	10	ANOVA P = 0.036 (in favour of AVT)
	Visual exploration training		5.0	(SE displayed on graph only)	10	
	Audiovisual exploration training (AVT)	Reading time (seconds)	75	(SE displayed on graph only)	10	ANOVA P = 0.03 (in favour of AVT)
	Visual exploration training		178	(SE displayed on graph only)	10	

Table 5. Results of studies comparing two similar active interventions (i.e. two interventions from the same category) (Continued)

	Audiovisual exploration training (AVT)	Visual scanning (percentage hits)	85.3	(SE displayed on graph only)	10	ANOVA P = 0.01 (in favour of AVT)
	Visual exploration training		64.1	(SE displayed on graph only)	10	
Modden 2012	Computer-based compensation therapy	Visual field enlargement (visual field assessment from Test Battery of Attentional Performance)	2.9	4.0	15	Pre- to post-treatment significant field expansion (P = 0.013)
	Standard occupational therapy (compensation)		1.3	4.7	15	Pre- to post-treatment: no significant field expansion (P = 0.316)
	Computer-based compensation therapy (CT)	Functional ability in ADL (improvement in Extended Barthel Index)	3.3	3.6	15	"No significant treatment effects were found when comparing ... CT/OT".
	Standard occupational therapy (OT) (compensation)		1.8	2.0	15	
	Computer-based compensation therapy (CT)	Reading - Improvement in reading performance, reduction in number of errors (from baseline)	-0.9	1.1	15	"Compared with OT"... "CT did not significantly reduce reading errors."
	Standard occupational therapy (OT) (compensation)		-0.7	1.0	15	
	Computer-based compensation	Visual scanning - reduction in number of omissions from baseline, cancellation tasks of the Test Battery of Attentional Performance	-5.4	5.2	15	"Compared with OT"... "CT did not result in superior improvements".

Table 5. Results of studies comparing two similar active interventions (i.e. two interventions from the same category) (Continued)

	therapy (CT)					
	Standard occupational therapy (OT) (compensation)		-2.3	5	15	
Substitution: one substitution intervention versus another substitution intervention						
Bainbridge 1994	Full-field Fresnel Prisms	Visual Field (Harrington Flocks Visual Field Score)	2.9	2	10	States full-field more improved
	Hemi-field Fresnel Prisms		7.2	3	8	
	Full-field Fresnel Prisms	Scanning (Line cancellation test errors)	4.7	1.3	10	P < 0.01, Student's t-test (in favour of full-field prisms)
	Hemi-field Fresnel Prisms		0.3	0.6	8	
Szlyk 2005	18.5 diopetre Gottlieb Visual field awareness system prisms	Visual skills category assessment battery		"There was improvement within all categories with both of the prism systems ranging from 36% for mobility (with the Fresnel prisms) to 13% for recognition (with the Gottlieb VFAS)."	10 (data only available for after crossover)	"There were no statistically significant differences between improvements with the Gottlieb VFAS compared with the Fresnel prisms."
	Press-on TM 20 Diopter Fresnel prisms					

ADL: activities of daily living

ANOVA: analysis of variance (statistical test of)

AVT: audiovisual exploration training

CT: compensation therapy

HRP: high-resolution perimetry

LV-VFQ: Low Vision Visual Functioning Questionnaire

MD: mean difference

OT: occupational therapy

SE: standard error

SEM: standard error of the mean

tDCS: transcranial direct current stimulation

VFAS: visual field awareness system

VRT: visual restitution therapy

Table 6. Results of studies comparing two different types of active interventions (i.e. interventions from different categories)

Study	Interventions	Outcome	Mean (or other reported result if no mean available)	Standard deviation	Number of participants	Statistical test/results
Compensation intervention versus restitution intervention						
Modden 2012	Computer-based restitution therapy	Visual field enlargement (visual field assessment from Test Battery of Attentional Performance)	3.9	4.9	15	Pre- to post-treatment significant field expansion (P = 0.003)
	Computer-based compensation therapy		2.9	4.0	15	
	Computer-based restitution therapy (RT)	Functional ability in ADL (improvement in Extended Barthel Index)	1.5	2.8	15	"No significant treatment effects were found when comparing ... RT/CT".
	Computer-based compensation therapy (CT)		3.3	3.6	15	
	Computer-based restitution therapy (RT)	Reading: improvement in reading performance, reduction in number of errors (from baseline)	-0.9	2.4	15	"There were no differences between RT and CT."
	Computer-based compensation therapy (CT)		-0.9	1.1	15	
	Computer-based restitution therapy (RT)	Visual scanning: reduction in number of omissions from baseline, cancellation tasks of the Test Battery of Attentional Performance	-5.3	10.5	15	"... the improvement of the CT compared with the RT group did not meet the defined significance level after Bonferroni correction (P = .023)."
	Computer-based compensation therapy (CT)		-5.4	5.2	15	

Table 6. Results of studies comparing two different types of active interventions (i.e. interventions from different categories) (Continued)

	Intervention	Comparison	Mean (SD)	Mean (SD)	n	Notes	
Roth 2009	Explorative scanning training (EST) (compensation)	Visual field: Tübingen automated perimetry	44.4	13.1	15	"Neither the EST group nor the FT group showed any differences in their TAP or SLO outcomes, quantified as the total number of stimuli detected in the blind hemifield (lowest P = 0.204)."	
		Flicker stimulation training (FT) (restitution)	35.7	15.2	13		
	Explorative scanning training (EST) (compensation)	Quality of life (WHOQOL-BREF)	12.93	1.67	15		
		Flicker stimulation training (FT) (restitution)	13.23	1.3	13		
	Explorative scanning training (EST) (compensation)	Reading (reading speed)	99.7	34.7	15		"Although the EST and FT groups differed in their reading speeds at T1, this difference remained unchanged [main effect of group, F(1,26) = 133.074, P < 0.0001, interaction, F < 1]".
		Flicker stimulation training (FT) (restitution)	140.2	20.9	13		

Compensation intervention versus substitution intervention

Rowe 2010*	Fresnel prisms (substitution)	Visual Field (relative change in visual field area)	0.052	0.1396	24	ANOVA results: no significant differences between groups (P = 0.55, for comparison across 3 treatment groups)
	Visual search		0.0815	0.1488	24	

Table 6. Results of studies comparing two different types of active interventions (i.e. interventions from different categories) (Continued)

training (compensation)						
Fresnel prisms (substitution)	Extended activities of daily living (change in EADL from baseline)	15.2	4.8	22		"No evidence of differences ..."
Visual search training (compensation)		15.2	4.4	22		
Fresnel prisms (substitution)	Reading (change in Radner reading speed)	17.4	21.3	24		"No evidence of differences ..."
Visual search training (compensation)		13.0	13.1	25		
Fresnel prisms (substitution)	Quality of life (VFQ-25 total score)	68.2	18.4	24		"Visual function (using the VFQ 25-10) improved at 26 weeks in the visual search training arm (60 [SD 19] to 68.4 [SD 20]) when compared to the Fresnel prisms (68.5 [SD 16.4] to 68.2 [18.4]) and standard care arms (63.7 [SD 19.4] to 59.8 [SD 22.7]: Table 6, ANCOVA P = 0.05)."
Visual search training (compensation)		68.4	20.0	25		
Fresnel prisms (substitution)	Adverse events (number of participants with reported adverse events during study)	18 participants		26		"Given the extent and range of adverse events reported with prism wear, caution must be exercised if prescribing prism glasses as an intervention for homonymous hemianopia."
Visual search training		2 participants		30		

Table 6. Results of studies comparing two different types of active interventions (i.e. interventions from different categories) (Continued)

(compensation)

*Rowe 2010 also had a control (standard care) group, and data were included in relevant meta-analyses for compensatory and substitution interventions versus control.

ADL: activities of daily living

ANCOVA: analysis of covariance (statistical test of)

ANOVA: analysis of variance (statistical test of)

CT: compensation therapy

EADL: extended activities of daily living

EST: explorative scanning training

FT: flicker stimulation training

RT: restitution therapy

SD: standard deviation

SLO: Scanning Laser Ophthalmoscope

T1: outcome assessment timepoint 1

T2: outcome assessment timepoint 2

TAP: Tuebingen automated perimetry

VFQ: visual function questionnaire

WHOQOL-BREF: World Health Organization Quality of Life Instrument

APPENDICES

Appendix 1. Cochrane Central Register of Controlled Trials (CENTRAL) Search Strategy

1. MeSH descriptor Cerebrovascular Disorders, this term only
2. MeSH descriptor Basal Ganglia Cerebrovascular Disease explode all trees
3. MeSH descriptor Brain Ischemia explode all trees
4. MeSH descriptor Carotid Artery Diseases explode all trees
5. MeSH descriptor Intracranial Arterial Diseases explode all trees
6. MeSH descriptor Intracranial Arteriovenous Malformations explode all trees
7. MeSH descriptor Intracranial Embolism and Thrombosis explode all trees
8. MeSH descriptor Intracranial Hemorrhages explode all trees
9. MeSH descriptor Stroke explode all trees
10. MeSH descriptor Brain Infarction explode all trees
11. MeSH descriptor Vasospasm, Intracranial, this term only
12. MeSH descriptor Vertebral Artery Dissection, this term only
13. stroke or poststroke or post-stroke or cerebrovasc* or brain vascul* or cerebral vascul* or cva* or apoplex* or SAH
14. (brain* or cerebr* or cerebell* or intracran* or intracerebral) near/5 (isch?emi* or infarct* or thrombo* or emboli* or occlus*)
15. (brain* or cerebr* or cerebell* or intracerebral or intracranial or subarachnoid) near/5 (haemorrhage* or hemorrhage* or haematoma* or hematoma* or bleed*)
16. MeSH descriptor Hemiplegia, this term only
17. MeSH descriptor Paresis explode all trees
18. hemipleg* or hemipar* or paresis or paretic 1735
19. (#1 OR #2 OR #3 OR #4 OR #5 OR #6 OR #7 OR #8 OR #9 OR #10 OR #11 OR #12 OR #13 OR #14 OR #15 OR #16 OR #17 OR #18)
20. MeSH descriptor Eye explode all trees
21. MeSH descriptor Visually Impaired Persons explode all trees
22. MeSH descriptor Ocular Physiological Processes explode all trees
23. MeSH descriptor Diagnostic Techniques, Ophthalmological explode all trees
24. MeSH descriptor Optometry explode all trees
25. MeSH descriptor Orthoptics explode all trees

- 26.MeSH descriptor Eye Diseases, this term only
- 27.MeSH descriptor Vision Disorders, this term only
- 28.MeSH descriptor Eye Manifestations, this term only
- 29.MeSH descriptor Blindness, this term only
- 30.MeSH descriptor Diplopia explode all trees
- 31.MeSH descriptor Vision, Binocular, this term only
- 32.MeSH descriptor Vision, Monocular, this term only
- 33.MeSH descriptor Visual Acuity explode all trees
- 34.MeSH descriptor Visual Fields, this term only
- 35.MeSH descriptor Vision, Low, this term only
- 36.MeSH descriptor Perimetry, this term only
- 37.MeSH descriptor Ophthalmology, this term only
- 38.MeSH descriptor Vision Screening, this term only
- 39.MeSH descriptor Eye Diseases, Hereditary explode all trees
- 40.MeSH descriptor Eye Hemorrhage explode all trees
- 41.MeSH descriptor Lacrimal Apparatus Diseases explode all trees
- 42.MeSH descriptor Lens Diseases explode all trees
- 43.MeSH descriptor Ocular Hypertension explode all trees
- 44.MeSH descriptor Ocular Hypotension explode all trees
- 45.MeSH descriptor Ocular Motility Disorders explode all trees
- 46.MeSH descriptor Optic Nerve Diseases explode all trees
- 47.MeSH descriptor Orbital Diseases explode all trees
- 48.MeSH descriptor Pupil Disorders explode all trees
- 49.MeSH descriptor Refractive Errors explode all trees
- 50.MeSH descriptor Retinal Diseases explode all trees
- 51.MeSH descriptor Blindness, Cortical explode all trees
- 52.MeSH descriptor Hemianopsia explode all trees
- 53.MeSH descriptor Vitreoretinopathy, Proliferative explode all trees
- 54.MeSH descriptor Vitreous Detachment explode all trees
- 55.MeSH descriptor Scotoma, this term only
- 56.MeSH descriptor Abducens Nerve, this term only
- 57.MeSH descriptor Oculomotor Nerve, this term only
- 58.MeSH descriptor Trochlear Nerve, this term only
- 59.nystagmus or smooth pursuit or saccades or depth perception or stereopsis or gaze disorder* or retinal or retinopathy or macular degeneration or glaucoma or cataract* or ophthalmol* or optic nerve
- 60.intranuclear ophthalmoplegia or parinaud's syndrome or weber's syndrome or skew deviation or conjugate deviation
- 61.one near/3 half syndrome
- 62.(visual* or vision or eye or eyes or eyesight or sight) near/5 (problem* or disorder* or impair* or disabilit* or loss or disease* or defect* or manifestation* or screening or test* or examination*)
- 63.hemianop* or blindness or low vision or refractive errors or vitreoretinopathy or vitreous detachment or scotoma or diplopia or optometr* or ocular or orthoptic*
- 64.oscillopsia or visual tracking or fresnel prism*
- 65.III or IV or VI or third or fourth or sixth near/3 nerve palsy
- 66.(#20 OR #21 OR #22 OR #23 OR #24 OR #25 OR #26 OR #27 OR #28 OR #29 OR #30 OR #31 OR #32 OR #33 OR #34 OR #35 OR #36 OR #37 OR #38 OR #39 OR #40 OR #41 OR #42 OR #43 OR #44 OR #45 OR #46 OR #47 OR #48 OR #49 OR #50 OR #51 OR #52 OR #53 OR #54 OR #55 OR #56 OR #57 OR #58 OR #59 OR #60 OR #61 OR #62 OR #63 OR #64 OR #65)
- 67.(#19 AND #66)
- 68.MeSH descriptor Infant explode all trees
- 69.MeSH descriptor Child explode all trees
- 70.neonat* or child or children or childhood or juvenile or infan* or toddler
- 71.MeSH descriptor Neoplasms explode all trees
- 72.cancer* or carcinoma* or tumor* or tumour* or neoplasm*
- 73.(#68 OR #69 OR #70 OR #71 OR #72)

74.(#67 AND NOT #73)

Appendix 2. MEDLINE (Ovid) Search Strategy

1. cerebrovascular disorders/ or exp basal ganglia cerebrovascular disease/ or exp brain ischemia/ or exp carotid artery diseases/ or cerebral small vessel diseases/ or stroke, lacunar/ or exp intracranial arterial diseases/ or exp intracranial arteriovenous malformations/ or exp "intracranial embolism and thrombosis"/ or exp intracranial hemorrhages/ or stroke/ or exp brain infarction/ or vasospasm, intracranial/ or vertebral artery dissection/
2. (stroke or poststroke or post-stroke or cerebrovasc\$ or brain vasc\$ or cerebral vasc\$ or cva\$ or apoplex\$ or SAH).tw.
3. ((brain\$ or cerebr\$ or cerebell\$ or intracran\$ or intracerebral) adj5 (isch?emi\$ or infarct\$ or thrombo\$ or emboli\$ or occlus\$)).tw.
4. ((brain\$ or cerebr\$ or cerebell\$ or intracerebral or intracranial or subarachnoid) adj5 (haemorrhage\$ or hemorrhage\$ or haematoma\$ or hematoma\$ or bleed\$)).tw.
5. hemiplegia/ or exp paresis/
6. (hemiple\$ or hemipar\$ or paresis or paretic).tw.
7. 1 or 2 or 3 or 4 or 5 or 6
8. exp eye/
9. exp visually impaired persons/
10. exp ocular physiological processes/ or exp diagnostic techniques, ophthalmological/
11. Optometry/ or Orthoptics/
12. eye diseases/ or vision disorders/ or eye manifestations/ or blindness/ or diplopia/
13. vision, binocular/ or vision, monocular/ or exp visual acuity/ or visual fields/ or vision, low/ or perimetry/ or ophthalmology/ or vision screening/
14. exp ocular motility disorders/ or exp orbital diseases/ or exp pupil disorders/ or exp blindness, cortical/ or exp hemianopsia/ or scotoma/
15. abducens nerve/ or oculomotor nerve/ or trochlear nerve/
16. (nystagmus or smooth pursuit or saccades or depth perception or stereopsis or gaze disorder\$ or ophthalmol\$ or optic nerve).tw.
17. (intranuclear ophthalmoplegia or parinaud's syndrome or weber's syndrome or skew deviation or conjugate deviation or (one adj3 half syndrome)).tw.
18. ((visual\$ or vision or eye or eyes or eyesight or sight) adj5 (problem\$ or disorder\$ or impair\$ or disabilit\$ or loss or disease\$ or defect\$ or manifestation\$ or screening or test\$ or examination\$)).tw.
19. (hemianop\$ or blindness or low vision or scotoma or diplopia or optometr\$ or ocular or orthoptics\$).tw.
20. (oscillopsia or visual tracking or fresnel prism\$).tw.
21. ((III or IV or VI or third or fourth or sixth) adj3 nerve palsy).tw.
22. or/8-21
23. Randomized Controlled Trials as Topic/
24. random allocation/
25. Controlled Clinical Trials as Topic/
26. control groups/
27. clinical trials as topic/ or clinical trials, phase i as topic/ or clinical trials, phase ii as topic/ or clinical trials, phase iii as topic/ or clinical trials, phase iv as topic/
28. double-blind method/
29. single-blind method/
30. Placebos/
31. placebo effect/
32. cross-over studies/
33. randomized controlled trial.pt.
34. controlled clinical trial.pt.
35. (clinical trial or clinical trial phase i or clinical trial phase ii or clinical trial phase iii or clinical trial phase iv).pt.
36. (random\$ or RCT or RCTs).tw.
37. (controlled adj5 (trial\$ or stud\$)).tw.
38. (clinical\$ adj5 trial\$).tw.
39. ((control or treatment or experiment\$ or intervention) adj5 (group\$ or subject\$ or patient\$)).tw.
40. (quasi-random\$ or quasi random\$ or pseudo-random\$ or pseudo random\$).tw.
41. ((control or experiment\$ or conservative) adj5 (treatment or therapy or procedure or manage\$)).tw.
42. ((singl\$ or doubl\$ or tripl\$ or trebl\$) adj5 (blind\$ or mask\$)).tw.
43. (cross-over or cross over or crossover).tw.
44. (placebo\$ or sham).tw.
45. trial.ti.
46. (assign\$ or allocat\$).tw.
47. controls.tw.
48. or/23-47
49. 7 and 22 and 48

50. exp animals/ not humans.sh.
 51. 49 not 50
 52. (neonat\$ or child or children or childhood or juvenile or infant or toddler).ti.
 53. 51 not 52

Appendix 3. Embase Search Strategy

1. cerebrovascular disease/ or exp basal ganglion hemorrhage/ or exp brain hematoma/ or exp brain hemorrhage/ or exp brain infarction/ or exp brain ischemia/ or exp carotid artery disease/ or cerebral artery disease/ or exp cerebrovascular accident/ or exp cerebrovascular malformation/ or exp intracranial aneurysm/ or exp occlusive cerebrovascular disease/ or stroke unit/ or stroke patient/
2. (stroke or poststroke or post-stroke or cerebrovasc\$ or brain vasc\$ or cerebral vasc\$ or cva\$ or apoplex\$ or SAH).tw.
3. ((brain\$ or cerebr\$ or cerebell\$ or intracran\$ or intracerebral) adj5 (isch?emi\$ or infarct\$ or thrombo\$ or emboli\$ or occlus\$)).tw.
4. ((brain\$ or cerebr\$ or cerebell\$ or intracerebral or intracranial or subarachnoid) adj5 (haemorrhage\$ or hemorrhage\$ or haematoma\$ or hematoma\$ or bleed\$)).tw.
5. hemiparesis/ or hemiplegia/ or paresis/
6. (hemipleg\$ or hemipar\$ or paresis or paretic).tw.
7. 1 or 2 or 3 or 4 or 5 or 6
8. exp eye/ or exp eye disease/ or exp visual disorder/
9. exp visual system examination/ or eye examination/ or exp vision test/
10. exp ophthalmology/ or orthoptics/ or exp visual system/ or exp visual system function/ or depth perception/
11. exp visual aid/
12. abducens nerve/ or oculomotor nerve/ or trochlear nerve/
13. (nystagmus or smooth pursuit or saccades or depth perception or stereopsis or gaze disorder\$ or ophthalmol\$ or optic nerve).tw.
14. (intranuclear ophthalmoplegia or parinaud's syndrome or weber's syndrome or skew deviation or conjugate deviation or (one adj3 half syndrome)).tw.
15. ((visual\$ or vision or eye or eyes or eyesight or sight) adj5 (problem\$ or disorder\$ or impair\$ or disabilit\$ or loss or disease\$ or defect\$ or manifestation\$ or screening or test\$ or examination\$)).tw.
16. (hemianop\$ or blindness or low vision or scotoma or diplopia or optometr\$ or ocular or orthoptic\$).tw.
17. (oscillopsia or visual tracking or fresnel prism\$).tw.
18. ((III or IV or VI or third or fourth or sixth) adj3 nerve palsy).tw.
19. or/8-18
20. Randomized Controlled Trial/ or "randomized controlled trial (topic)"/
21. Randomization/
22. Controlled clinical trial/ or "controlled clinical trial (topic)"/
23. control group/ or controlled study/
24. clinical trial/ or "clinical trial (topic)"/ or phase 1 clinical trial/ or phase 2 clinical trial/ or phase 3 clinical trial/ or phase 4 clinical trial/
25. Crossover Procedure/
26. Double Blind Procedure/
27. Single Blind Procedure/ or triple blind procedure/
28. placebo/ or placebo effect/
29. (random\$ or RCT or RCTs).tw.
30. (controlled adj5 (trial\$ or stud\$)).tw.
31. (clinical\$ adj5 trial\$).tw.
32. ((control or treatment or experiment\$ or intervention) adj5 (group\$ or subject\$ or patient\$)).tw.
33. (quasi-random\$ or quasi random\$ or pseudo-random\$ or pseudo random\$).tw.
34. ((control or experiment\$ or conservative) adj5 (treatment or therapy or procedure or manage\$)).tw.
35. ((singl\$ or doubl\$ or tripl\$ or trebl\$) adj5 (blind\$ or mask\$)).tw.
36. (cross-over or cross over or crossover).tw.
37. (placebo\$ or sham).tw.
38. trial.ti.
39. (assign\$ or allocat\$).tw.
40. controls.tw.
41. or/20-40
42. 7 and 19 and 41
43. (exp animals/ or exp invertebrate/ or animal experiment/ or animal model/ or animal tissue/ or animal cell/ or nonhuman/) not (human/ or normal human/ or human cell/)
44. 42 not 43
45. (neonat\$ or child or children or childhood or juvenile or infant or toddler).ti.
46. exp Neoplasm/
47. (cancer\$ or carcinoma\$ or tumor\$ or tumour\$ or neoplasm\$).tw.
48. 45 or 46 or 47
49. 44 not 48

Appendix 4. CINAHL Search Strategy

- S1. MH "Cerebrovascular Disorders+" or MH "stroke patients" or MH "stroke units"
- S2. TI (stroke or poststroke or post-stroke or cerebrovasc* or brain vas*) or AB (stroke or poststroke or post-stroke or cerebrovasc* or brain vas*)
- S3. TI (brain* or cerebr* or cerebell* or intracran* or intracerebral) or AB (brain* or cerebr* or cerebell* or intracran* or intracerebral)
- S4. TI (ischemi* or ischaemi* or infarct* or thrombo* or emboli* or occlus*) or AB (ischemi* or ischaemi* or infarct* or thrombo* or emboli* or occlus*)
- S5. S3 and S4
- S6. TI (brain* or cerebr* or cerebell* or intracerebral or intracranial or subarachmoid) or AB (brain* or cerebr* or cerebell* or intracerebral or intracranial or subarachnoid)
- S7. TI (haemorrhage* or hemorrhage* or haematoma* or hematoma* or bleed*) or AB (haemorrhage* or hemorrhage* or haematoma* or hematoma* or bleed*)
- S8. S6 and S7
- S9. MH "Hemiplegia"
- S10. TI (hemipleg* or hemipar* or paresis or paretic) or AB (hemipleg* or hemipar* or paresis or paretic)
- S11. S1 or S2 or S5 or S8 or S9 or S10
- S12. MH "Eye+" or MH "Rehabilitation of Vision Impaired+" or MH "Optometry" or MH "Eye Diseases+"
- S13. MH "Visual Acuity+" or MH "Perimetry+" or MH "Ophthalmology+" or MH "Vision Screening+" or MH "Ocular Physiology+"
- S14. TI (orthoptics or vision, monocular or vision, binocular) or AB (orthoptics or vision, monocular or vision, binocular)
- S15. TI (vitreous detachment or hemianopsia or hemianopia or quadrantanopia) or AB (vitreous detachment or hemianopsia or hemianopia or quadrantanopia)
- S16. MH "Abducens Nerve" or MH "oculomotor nerve" or MH "troclear nerve" or MH "optic nerve" or MH "nystagmus, pathologic"
- S17. TI (smooth pursuit or saccades or gaze disorder* or retinal or retinopathy or ophthalmol*) or AB (smooth pursuit or saccades or gaze disorder* or retinal or retinopathy or ophthalmol*)
- S18. TI (hemianop* or blindness or low vision or refractive errors or vitreoretinopathy or vitreous detachment or scotoma or diplopia or optometry* or ocular or orthoptic*) or AB (hemianop* or blindness or low vision or refractive errors or vitreoretinopathy or vitreous detachment or scotoma or diplopia or optometry* or ocular or orthoptic*)
- S19. TI (oscillopsia or visual tracking or fresnel prism*) or AB (oscillopsia or visual tracking or fresnel prism*)
- S20. TI (intranuclear ophthalmoplegia or parinaud's syndrome or weber's syndrome or skew deviation or conjugate deviation) or AB (intranuclear ophthalmoplegia or parinaud's syndrome or weber's syndrome or skew deviation or conjugate deviation)
- S21. TI (visual* or vision or eye or eyes or eyesight or sight) or AB (visual* or vision or eye or eyes or eyesight or sight)
- S22. TI (problem* or disorder* or impair* or disability* or loss or disease* or defect* or manifestation* or screening or test* or examination*) or AB (problem* or disorder* or impair* or disability* or loss or disease* or defect* or manifestation* or screening or test* or examination*)
- S23. S21 and S22
- S24. TI (third or fourth or sixth) or AB (third or fourth or sixth)
- S25. AB nerve palsy or TI nerve palsy
- S26. S24 and S25
- S27. S12 or S13 or S14 or S15 or S16 or S17 or S18 or S19 or S20 or S23 or S26
- S28. S11 and S27
- S29. (MH "Random Assignment") or (MH "Random Sample+")
- S30. (MH "Crossover Design") or (MH "Clinical Trials+") or (MH "Comparative Studies")
- S31. (MH "Control (Research)") or (MH "Control Group")
- S32. (MH "Factorial Design") or (MH "Quasi-Experimental Studies") or (MH "Nonrandomized Trials")
- S33. (MH "Placebo Effect") or (MH "Placebos") or (MH "Meta Analysis")
- S34. (MH "Community Trials") or (MH "Experimental Studies") or (MH "One-Shot Case Study") or (MH "Pretest-Posttest Design+") or (MH "Solomon Four-Group Design") or (MH "Static Group Comparison") or (MH "Study Design")
- S35. (MH "Clinical Research") or (MH "Clinical Nursing Research")
- S36. PT clinical trial
- S37. PT systematic review
- S38. TI random* or AB random*
- S39. TI (singl* or doubl* or tripl* or trebl*) or AB (singl* or doubl* or tripl* or trebl*)
- S40. TI (blind* or mask*) or AB (blind* or mask*)
- S41. S39 and S40
- S42. TI (crossover or cross-over or placebo* or control* or factorial or sham) or AB (crossover or cross-over or placebo* or control* or factorial or sham)
- S43. TI (clin* or intervention* or compar* or experiment* or preventive or therapeutic) or AB (clin* or intervention* or compar* or experiment* or preventive or therapeutic)
- S44. TI trial* or AB trial*
- S45. S43 and S44
- S46. TI (counterbalance* or multiple baseline* or ABAB design) or AB (counterbalance* or multiple baseline* or ABAB design)

S47. TI (meta analysis* or metaanalysis or meta-anlysis or systematic review*) or AB (meta analysis* or metaanalysis or meta-anlysis or systematic review*)

S48. S29 or S30 or S31 or S32 or S33 or S34 or S35 or S36 or S37 or S38 or S41 or S42 or S45 or S46 or S47

S49. S28 AND S48

Appendix 5. AMED Search Strategy

1. cerebrovascular disorders/ or cerebral hemorrhage/ or cerebral infarction/ or cerebral ischemia/ or cerebrovascular accident/ or stroke/
2. (stroke or poststroke or post-stroke or cerebrovasc\$ or brain vasc\$ or cerebral vasc\$ or cva\$ or apoplex\$ or SAH).tw.
3. ((brain\$ or cerebr\$ or cerebell\$ or intracran\$ or intracerebral) adj5 (isch?emi\$ or infarct\$ or thrombo\$ or emboli\$ or occlus\$)).tw.
4. ((brain\$ or cerebr\$ or cerebell\$ or intracerebral or intracranial or subarachnoid) adj5 (haemorrhage\$ or hemorrhage\$ or haematoma\$ or hematoma\$ or bleed\$)).tw.
5. hemiplegia/
6. (hemipleg\$ or hemipar\$ or paresis or paretic).tw.
7. 1 or 2 or 3 or 4 or 5 or 6
8. exp eye/
9. eye disease/ or exp ocular motility disorders/ or exp vision disorders/ or optic nerve/
10. (nystagmus or smooth pursuit or saccades or depth perception or stereopsis or gaze disorder\$ or ophthalmol\$ or optic nerve).tw.
11. (intranuclear ophthalmoplegia or parinaud's syndrome or weber's syndrome or skew deviation or conjugate deviation or (one adj3 half syndrome)).tw.
12. ((visual\$ or vision or eye or eyes or eyesight or sight) adj5 (problem\$ or disorder\$ or impair\$ or disabilit\$ or loss or disease\$ or defect\$ or manifestation\$ or screening or test\$ or examination\$)).tw.
13. (hemianop\$ or blindness or low vision or scotoma or diplopia or optometr\$ or ocular or orthoptics\$).tw.
14. (oscillopsia or visual tracking or fresnel prism\$).tw.
15. ((III or IV or VI or third or fourth or sixth) adj3 nerve palsy).tw.
16. or/8-15
17. clinical trials/ or randomized controlled trials/ or random allocation/
18. research design/ or comparative study/
19. double blind method/ or single blind method/
20. placebos/
21. (random\$ or RCT or RCTs).tw.
22. (controlled adj5 (trial\$ or stud\$)).tw.
23. (clinical\$ adj5 trial\$).tw.
24. ((control or treatment or experiment\$ or intervention) adj5 (group\$ or subject\$ or patient\$)).tw.
25. (quasi-random\$ or quasi random\$ or pseudo-random\$ or pseudo random\$).tw.
26. ((control or experiment\$ or conservative) adj5 (treatment or therapy or procedure or manage\$)).tw.
27. ((singl\$ or doubl\$ or tripl\$ or trebl\$) adj5 (blind\$ or mask\$)).tw.
28. (cross-over or cross over or crossover).tw.
29. (placebo\$ or sham).tw.
30. trial.ti.
31. (assign\$ or allocat\$).tw.
32. controls.tw.
33. or/17-32
34. 7 and 16 and 33

Appendix 6. PsycINFO Search Strategy

1. cerebrovascular disorders/ or cerebral hemorrhage/ or exp cerebral ischemia/ or cerebral small vessel disease/ or cerebrovascular accidents/ or subarachnoid hemorrhage/
2. (stroke or poststroke or post-stroke or cerebrovasc\$ or brain vasc\$ or cerebral vasc\$ or cva\$ or apoplex\$ or SAH).tw.
3. ((brain\$ or cerebr\$ or cerebell\$ or intracran\$ or intracerebral) adj5 (isch?emi\$ or infarct\$ or thrombo\$ or emboli\$ or occlus\$)).tw.
4. ((brain\$ or cerebr\$ or cerebell\$ or intracerebral or intracranial or subarachnoid) adj5 (haemorrhage\$ or hemorrhage\$ or haematoma\$ or hematoma\$ or bleed\$)).tw.
5. hemiplegia/ or hemiparesis/
6. (hemipleg\$ or hemipar\$ or paresis or paretic).tw.
7. 1 or 2 or 3 or 4 or 5 or 6
8. exp "eye (anatomy)"/
9. optometry/ or ophthalmology/ or ophthalmologic examination/
10. exp eye disorders/ or exp eye movements/ or exp vision disorders/
11. binocular vision/ or monocular vision/ or visual acuity/ or visual field/ or peripheral vision/ or exp depth perception/
12. optic nerve/ or abducens nerve/
13. (nystagmus or smooth pursuit or saccades or depth perception or stereopsis or gaze disorder\$ or ophthalmol\$ or optic nerve).tw.

14. (intranuclear ophthalmoplegia or parinaud's syndrome or weber's syndrome or skew deviation or conjugate deviation or (one adj3 half syndrome)).tw.
15. ((visual\$ or vision or eye or eyes or eyesight or sight) adj5 (problem\$ or disorder\$ or impair\$ or disabilit\$ or loss or disease\$ or defect\$ or manifestation\$ or screening or test\$ or examination\$)).tw.
16. (hemianop\$ or blindness or low vision or scotoma or diplopia or optometr\$ or ocular or orthoptic\$).tw.
17. (oscillopsia or visual tracking or fresnel prism\$).tw.
18. ((III or IV or VI or third or fourth or sixth) adj3 nerve palsy).tw.
19. or/8-18
20. clinical trials/ or treatment effectiveness evaluation/ or placebo/
21. (random\$ or RCT or RCTs).tw.
22. (controlled adj5 (trial\$ or stud\$)).tw.
23. (clinical\$ adj5 trial\$).tw.
24. ((control or treatment or experiment\$ or intervention) adj5 (group\$ or subject\$ or patient\$)).tw.
25. (quasi-random\$ or quasi random\$ or pseudo-random\$ or pseudo random\$).tw.
26. ((control or experiment\$ or conservative) adj5 (treatment or therapy or procedure or manage\$)).tw.
27. ((singl\$ or doubl\$ or tripl\$ or trebl\$) adj5 (blind\$ or mask\$)).tw.
28. (cross-over or cross over or crossover).tw.
29. (placebo\$ or sham).tw.
30. trial.ti.
31. (assign\$ or allocat\$).tw.
32. controls.tw.
33. or/20-32
34. 7 and 19 and 33

Appendix 7. PQDT Search Strategy

stroke AND hemianopia

OR

stroke AND "visual field"

OR

cerebrovascular AND hemianopia

OR

cerebrovascular AND "visual field"

Appendix 8. Trials Registers Search Strategies

US National Institutes of Health Ongoing Trials Register: ClinicalTrials.gov

(www.clinicaltrials.gov):

(hemianopia OR visual field OR vision defect OR eye diseases) AND (Brain Infarction OR Intracranial Hemorrhages OR Carotid Artery Diseases OR Brain Ischemia OR Cerebral Hemorrhage OR Cerebrovascular Disorders OR Stroke) [DISEASE]

World Health Organization International Clinical Trials Registry Platform

(apps.who.int/trialsearch):

stroke AND hemianopia OR stroke AND visual field
 cerebrovascular AND hemianopia OR cerebrovascular AND visual field

WHAT'S NEW

Date	Event	Description
26 February 2019	New search has been performed	Review updated, with searches completed in May 2018. The review now includes 20 studies; previous (2009) version included 13 studies. In this updated version, we have excluded three of the studies from the 2009; therefore, this updated version in-

Date	Event	Description
		cludes 10 studies which were included in the previous (2009) version and 10 new studies, with a total of 547 participants with stroke.
11 October 2018	New citation required and conclusions have changed	The update of this review has led to changes in the conclusions relating to compensatory interventions (with greater uncertainty around previous, limited evidence of effectiveness), and has introduced new evidence relating to adverse events (particularly headache) associated with substitutive interventions.

CONTRIBUTIONS OF AUTHORS

For this version: Alex Pollock led the review update, identified relevant studies, provided methodological expertise, extracted data, entered data, carried out analyses, and wrote the final drafts. Pauline Campbell ran searches, contributed to screening of titles, provided methodological expertise, and read and commented on drafts. Christine Hazelton screened titles, identified relevant studies, extracted data, acted as a second review author, provided content expertise, and contributed to final drafts. Fiona Rowe acted as a second review author, provided content expertise, extracted and synthesised data relating to outcome measures, and contributed to final drafts. Sven Jonuscheit and Ashleigh Kernohan applied inclusion criteria to abstracts and full papers, identified relevant studies, extracted data, and read and commented on drafts. Jayne Angilley, Clair Henderson, and Peter Langhorne read and commented on drafts.

For the previous (2009) version: Alex Pollock led this review, provided methodological expertise, acted as a second review author, carried out analyses, and wrote the final drafts. Christine Hazelton ran searches, identified relevant studies, acted as first review author, extracted data, entered data, provided content expertise, and contributed to final drafts. Clair Henderson acted as a second review author for the identification of relevant studies. Baljean Dhillon, Heather Orr, Katrina Livingstone, Frank A Munro, Fiona Rowe, Uma Shahani, Jayne Angilley, and Peter Langhorne provided additional content expertise, read and commented on final drafts, and acted as additional reviewers where there was uncertainty or disagreement. Peter Langhorne also provided additional methodological expertise where there was uncertainty about the methodological design of studies.

DECLARATIONS OF INTEREST

Alex Pollock was involved in the VISION trial ([Rowe 2010](#)), which was funded by the Stroke Association. No other known interest.

Christine Hazelton has carried out non-randomised studies into the effectiveness of a number of scanning training interventions, including the intervention studied by [Roth 2009](#), and is developing further project proposals in this area. No other known interest.

Fiona Rowe was the chief investigator for the VISION trial ([Rowe 2010](#)), funded by the Stroke Association. She was a co-investigator for the [Jarvis 2012](#) trial, funded by the University of Liverpool.

Sven Jonuscheit: none known.

Ashleigh Kernohan: none known.

Jayne Angilley: none known.

Clair A Henderson: none known.

Peter Langhorne: none known.

Pauline Campbell: none known.

The work presented here represents the view of the review authors and not necessarily those of the funding bodies.

SOURCES OF SUPPORT

Internal sources

- No sources of support supplied

External sources

- Royal National Institute of Blind People (RNIB, Scotland), UK.
RNIB (Scotland) funded the first (2009) version of this review.
- Chief Scientist Office, UK.

Alex Pollock and Pauline Campbell are employed by the Nursing, Midwifery and Allied Health Professions Research Unit, which is funded by the Chief Scientist Office in Scotland.

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Participants

In the 2011 version of the review, we included studies that investigated the effectiveness of visual scanning training and techniques even if the population of participants had not been clearly defined as having visual field defects. This decision was made in consultation with a multidisciplinary group of clinicians (CH, JA, BD, KL, PL, FM, HO, FR, US); the decision was that it was clinically relevant and useful to include these studies. It is recognised that, in these studies, the visual scanning problems experienced by participants may have been due to either visual field defects, visual neglect, or a combination of both. The results of these studies were therefore relevant to the population of patients with 'visual scanning' problems, regardless of the physiological cause of the scanning problem. Data from these studies would not be combined with data arising from studies which include a population of patients with visual field defects but not visual neglect. In this update, this decision was reconsidered, and reversed: populations of participants with 'visual scanning problems', but no confirmed visual field defect have been excluded.

Comparisons

In the 2011 version, we included one additional comparison that we did not outline in the protocol. This was a comparison of compensative intervention versus restitutive interventions. Although not preplanned, discussion amongst the review authors led to the conclusion that this was a clinically relevant comparison and ought to be included in this review. However, for this updated review, we reached consensus that we should adhere to the pre-stated comparisons, but that we should include a summary of results data from any studies which compared two active interventions (i.e. studies with no control group). This has been added as [Table 5](#) and [Table 6](#) and a narrative synthesis included under [Effects of interventions](#).

Subgroup analyses

The following subgroup analyses were listed as planned in the protocol and previous versions of this review, but have been removed, as Cochrane guidance recommends restricting the number of subgroup analyses as large numbers can lead to spurious explanations of heterogeneity ([Higgins 2011](#)):

- age (under 60 years, 60 years and over);
- gender (men, women);
- side of stroke (left, right, bilateral);
- presence of age-related visual problems (presence, absence);
- presence of eye movement disorders (presence, absence);
- level of motor impairment (mild, moderate, severe);
- level of cognitive impairment (mild, moderate, severe).

Search strategy

Due to available time and resources for this latest update, and following a detailed consideration of the comprehensiveness of other databases:

- we searched, for the 2011 version of this review, the following databases, but did not search them for this updated version:
 - * British Nursing Index (searched 1985 to 31 December 2009);
 - * PsycBITE (Psychological Database for Brain Impairment Treatment Efficacy) at www.psycbite.com (last searched December 2009).
 - * Trials Central (www.trialscentral.org) (last searched February 2010).
- for the 2011 version, but not for this updated version, we also:
 - * searched the references supplied by commercial companies providing interventions aimed at restoration of visual field defect (including NovaVision®);
 - * performed citation tracking using Web of Science Cited Reference Search for all included studies;
 - * handsearched proceedings of Association for Research in Vision and Ophthalmology (www.arvo.org) (1969 to 2010); these abstracts are now covered by MEDLINE, so were not included in the 2018 handsearching activities.

INDEX TERMS**Medical Subject Headings (MeSH)**

Quality of Life; Randomized Controlled Trials as Topic; Stroke [*complications]; Vision Disorders [etiology] [*rehabilitation]; Visual Fields

MeSH check words

Humans