## COCHRANE CORNER

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# Interventions for Perceptual Disorders in Stroke: A Systematic Review

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Perception is the ability to recognize and interpret information from our senses. It is fundamental to an individual's ability to understand and interact with their environment. Disorders of perception are common after stroke, reducing quality of life. Research evidence relating to effectiveness of interventions is unclear. This Cochrane review update and expansion assessed the effectiveness of interventions for perceptual disorders after stroke.<sup>1</sup>

## **METHODS**

We searched key online databases including CENTRAL, MEDLINE, and Embase from inception to August 2021. We also searched trial and research registers and screened the reference lists of included studies.

We included randomized controlled trials (RCTs) of any intervention targeting perceptual disorders following stroke and affecting hearing, taste, touch, smell, somatosensation, or vision. We excluded deficits of sensation, for example, visual field loss or attention, for example, neglect.

One reviewer screened titles for eligibility. Two reviewers independently screened abstracts and full-text articles.

Data extraction and risk-of-bias assessment (using the Risk of Bias-1 tool) were conducted by one reviewer and checked by a second; evidence quality was appraised using the Grading of Recommendations, Assessment, Development, and Evaluations tool.

We compared the benefits of active interventions with no treatment, control, or alternative active interventions, on stroke survivors' activities of daily living, our primary outcome measure, and other outcomes. Meta-analysis used Review Manager software and a random-effects model.

We involved lived experience (4 people) and clinical expert (4 people) stakeholder groups throughout the review.

## RESULTS

Of 94434 records identified, we included 18 RCTs (541 participants, 535 [98.9%] stroke survivors).

## Hearing, Taste, and Smell

No RCTs were found.

#### Somatosensation

Interventions included robot-assisted gait training, standard physiotherapy, mirror therapy, and transcranial direct current stimulation.

One RCT (n=24) compared active intervention (transcranial direct current stimulation) to control. Activities of daily living were assessed via the Korean modified Barthel index. Analysis showed no difference between groups (mean difference, 10.08 [95% CI, -2.47 to 22.63]; *P*=0.12); the evidence was assessed as being very low quality.

Three RCTs compared one active intervention (computerized balance and movement training) with another

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active intervention (standard Pusher syndrome physiotherapy) (n=80 with Pusher syndrome). Activities of daily living were assessed using the Korean modified Barthel index. Analysis showed the computerized therapy was more effective than standard physiotherapy (mean difference, 10.19 [95% CI, 4.94–15.44]; P=0.0001); there was no heterogeneity (I<sup>2</sup>=0%) and very low-quality evidence.

## Touch

Interventions included pressure sense training and hand exercises with an assistive glove.

One RCT (n=24) compared one active intervention (hand exercises with robotic glove) and another active intervention (conventional hand exercises) using the modified Barthel index. Analysis showed no difference between the interventions (mean difference, -0.41 [95% CI, -12.31 to 11.49]). Evidence was very low quality.

## Vision

Interventions included repeated figure drawing, computer-based games, and therapist-led functional activities.

Two RCTs (n=96) comparing one active intervention with another measured activities of daily living using the modified Barthel index; data were not combined due to intervention differences.

## DISCUSSION

Limited evidence currently exists to determine the effectiveness of any intervention for perceptual disorders impacting any sensory modality.

Clinicians should continue to provide neurorehabilitation for perceptual disorders according to the current clinical guidelines.

High-quality trials are needed on interventions for perceptual disorders in stroke. Trials should have sufficient participant numbers, usual care comparisons, and measure longer term functional outcomes.

### **ARTICLE INFORMATION**

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