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Biosimilar infliximab use in paediatric IBD

Lisa Richmond,¹ Lee Curtis,¹ Victoria Garrick,¹ Pam Rogers,² Michelle Wilson,³ Rachel Tayler,¹ Paul Henderson,^{2,3} Richard Hansen,¹ David C Wilson,³ Richard K Russell¹

¹Department of Paediatric Gastroenterology, Royal Hospital for Children, Glasgow, UK

²Department of Paediatric Gastroenterology, Royal Hospital for Sick Children, Edinburgh, UK

³Child Life and Health, University of Edinburgh, Edinburgh, UK

Correspondence to

Lisa Richmond, Paediatric Clinical Research Facility, Neurosciences building, Queen Elizabeth University Hospital, Glasgow G51 4TF, UK; lisa.richmond@nhs.net

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ABSTRACT

Background Biosimilar infliximab became available in the UK in 2015. Paediatric experience to date on its use is limited. We prospectively evaluated the safety and efficacy of biosimilar infliximab (Remsima) in two paediatric gastroenterology networks in patients with inflammatory bowel disease.

Methods Prospective clinical data were collected from laboratory reports, electronic patient records and case notes of 40 patients starting Remsima for the first time. Disease activity scores together with blood and stool biomarkers were used to assess response.

Results Our data set highlights that Remsima was associated with a significant clinical and biochemical improvement ($p < 0.01$ or less for all parameters assessed) in Crohn's disease post induction. There were no significant safety issues noted. The total cost saving was £47 800, representing a 38% reduction from originator.

Conclusion We found that biosimilar infliximab is as effective as originator infliximab and its use is associated with significant cost savings.

INTRODUCTION

There are an increasing number of patients with paediatric inflammatory bowel disease (PIBD) commencing antitumour necrosis factor (TNF) therapy in the form of infliximab.¹ Infliximab is commonly used in PIBD for chronically active disease, which is not controlled by 'conventional' therapy, fistulising perianal Crohn's disease (CD) or for acute severe colitis as an alternative to colectomy.² Initial PIBD use was limited due to the lack of a specific paediatric licence for Remicade, its significant cost and the potential safety profile, specifically malignancy.

In early 2015, European Medicine Agency licence approval was granted for biosimilar infliximab use in PIBD from initial studies in adults with rheumatological conditions. Biosimilar infliximab is available for use in the UK either as Remsima or Inflectra. Since the initial studies satisfying the criteria for biosimilarity are in adult rheumatology and did not use standard inflammatory bowel disease dosing strategies, an urgent need developed for PIBD-specific data, focusing both on safety and clinical effectiveness.

There is also the added opportunity for significant cost savings, which may in turn allow easier access to therapy. A prospective safety, efficacy and cost-effectiveness evaluation of biosimilar infliximab (Remsima) was carried out in two paediatric gastroenterology networks between 2015 and 2016.

PATIENTS AND METHODS

All patients with PIBD from the regional services in the West and South East of Scotland (based at the Royal Hospital for Children, Glasgow, and the Royal Hospital for Sick Children, Edinburgh) who required infliximab were started on Remsima. Data were collected at treatment initiation and response reviewed, following three induction doses, at around week 12 post initiation. In line with the licence, standard dosing at 5 mg/kg at 0, 2 and 6 weeks was used with infusions given over 2 hours. Patients were routinely given hydrocortisone at 4 mg/kg (maximum dose 200 mg) prior to each Remsima infusion. No consistent phial sharing protocols were used for the duration of this evaluation. Prospective clinical data were collected from local laboratory reports, electronic patient records and case notes of patients starting Remsima. Our intention was that all patients were anti-TNF-naïve at initiation.

Clinical disease activity indices, namely the weighted Paediatric Crohn's Disease Activity Index³ and Paediatric Ulcerative Colitis Activity Index,⁴ were used to document disease activity at initiation and follow-up. For analysis purposes, calprotectin samples and C reactive protein values were limited to laboratory-reported maximum and minimum assay values ($>1800 \mu\text{g/g}$ and $<1 \text{ mg/L}$, respectively). An adequate infliximab trough was deemed to be in the range of 3–7 mg/L. Levels and antibodies were measured routinely and were not dependent on clinical status.

Data were expressed as median with range. Statistics were calculated using Microsoft Excel (Microsoft Corporation, Redmond, Washington, USA) and GraphPad V.3 software (GraphPad Software, La Jolla, California, USA). Ethical approval was not required as this was an evaluation of clinical practice. Average costings for a phial of biosimilar and originator infliximab (Remsima and Remicade, respectively) were obtained from the recent award of a national framework agreement by National Procurement Scotland (OJEU, contract award notice: <http://ted.europa.eu/udl?uri=TED:NOTICE:92352-2015:TEXT:EN:HTML>). The cost saving was then calculated by adding the total number of phials used and calculating the cost saving on each phial multiplied by this difference.

RESULTS

Forty consecutive patients (60% male) started Remsima between August 2015 and June 2016, equating to 190 infusions in total. Twenty-nine patients had CD and 11 ulcerative colitis (UC)/inflammatory bowel disease unclassified (IBDU). The median age of patients (IQR) at



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Short report

Table 1 Characteristics of patients at initiation and 12-week follow-up

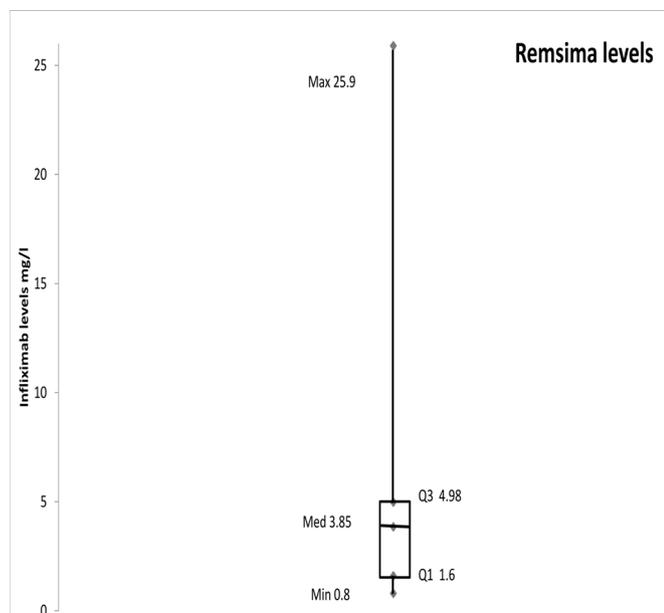
Clinical data	At initiation	At 12-week review	Comparative p Value
ESR, median (IQR)	16 (7.5–31.3) (n=40)	5 (2.25–10) (n=26)	p=0.0009
CRP, median (IQR)	5.5 (2–21.3) (n=40)	1 (1–2) (n=27)	p=0.0004
Albumin, median (IQR)	35 (30.8–38) (n=40)	38 (36–41) (n=27)	p=0.002
Calprotectin, median (IQR)	840 (618.3–1642.5) (n=34)	250 (60–760) (n=17)	p=0.008
wPCDAI, median (IQR)	27.5 (7.5–55) (n=29)	5 (0–20) (n=21)	p=0.002
PUCAI, median (IQR)	45 (40–55) (n=11)	23.8 (8.1–60) (n=6)	p=0.4
Disease classification (Crohn's) % (n/N)	Remission, 28% (8/29)	Remission, 67% (14/21)	p=0.002
	Mild, 31% (9/29)	Mild, 28% (6/21)	p=0.9
	Moderate, 28% (8/29)	Moderate, 0% (0/21)	p=0.02
	Severe, 13% (4/29)	Severe, 5% (1/21)	p=0.6

CRP, C reactive protein; PUCAI, Paediatric Ulcerative Colitis Activity Index; wPCDAI, weighted Paediatric Crohn's Disease Activity Index.

diagnosis was 12.7 years (10–14) and 13.7 years (13–16) at Remsima initiation.

The primary reasons for treatment in CD were active luminal disease 76% (22/29), perianal disease 14% (4/29) and with other indications 10% (3/29). For UC/IBDU, chronic refractory disease was the indication in 64% (7/11) and acute severe colitis in 36% (4/11). The majority (95%; 38/40) of patients were on coimmunomodulator therapy; 71% (27/38) on azathioprine/mercaptopurine and 29% (11/38) on methotrexate. All patients received intravenous hydrocortisone pre-Remsima dosing. At initiation 43% (17/40) were also on oral prednisolone. Remsima was associated with a significant clinical and biochemical improvement in CD postinduction (table 1). Infliximab levels and antibodies were recorded for 50% (20/40) of patients following induction (figure 1). The median trough level was 3.85 mg/L (IQR 1.60–4.98) and 10/20 (50%) samples had a level outside the targeted therapeutic range.

Only two patients had positive antibodies recorded, one of whom experienced an acute infusion reaction (AIR). This gives an AIR per infusion rate of 1/190 (0.5%) and AIR per patient 1/40 (2.5%) from this cohort. Immediately after the second

**Figure 1** Infliximab levels at week 14 following induction.

Remsima infusion began, the patient's face became flushed and the throat felt tight. The infusion was discontinued and therapy was subsequently changed from Remsima to adalimumab. Infliximab levels on this patient were recorded as 25.9 mg/L with strongly positive antibodies. This patient had in fact been exposed to Remicade 6 years previously with mild sensitivity. There were no additional safety issues recorded.

The average cost saving per phial during this period was approximately 38% reduction from the originator (Remicade) to the biosimilar (Remsima). For the total number of infusions carried out during this evaluation, we estimate a conservative cost saving of around £47 800 (€57 000).

DISCUSSION

The benefits of infliximab treatment for PIBD are well established and represent a cornerstone of PIBD practice for patients with inadequate or lost response to first-line and second-line therapy. Published studies to date have described outcomes using originator infliximab (Remicade). These data have shown that early clinical response and remission rates following an induction course of Remsima match those of historical patients treated with infliximab (Remicade).¹ To date, there is a paucity of data in the paediatric literature on biosimilar use, with none published from the UK⁵ and none as robust as the current study, which prospectively collected clinical and laboratory data as part of good clinical practice. Recent guidelines have generally advised caution in the use of biosimilar anti-TNFs until further data on their use have become available⁶; we hope this study will help support their wider introduction into paediatric practice.

Continued close monitoring of these patients is necessary to collect longer term data and to evaluate maintenance of remission, secondary loss of response and safety. The financial benefits of using these treatments are clear from this and other studies, but will only be upheld if long-term maintenance rates match these initial clinical response rates. As the emerging data for treatment-naïve patients are published, the logical next step is to look at switching those already established on originator infliximab to biosimilars. Early paediatric data on this are encouraging, but more detailed information on patient outcomes, including trough drug levels and antibodies, is awaited.⁷ The trough level data from this study showed a wide range of values with several below the quoted therapeutic range. Although we attempted to collect these data in all patients, there may be a slight bias in the samples we collected towards patients with signs of limited/no response to infliximab. This may be reflected in the fact that 40% of the results were subtherapeutic. Importantly there were no signs of significant allergenicity.

In summary, we have demonstrated that biosimilar infliximab is as safe and effective as originator infliximab in the short-term using data from two regional referral centres.

There are also significantly associated cost savings. These baseline data have now enabled us to confidently switch patients from originator to biosimilar, adopting the same prospective methodology to monitor effectiveness, safety and cost.

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Competing interests LR: conference fees and travel from Tillotts. VG: conference fees and travel from AbbVie. RT: speaker's fees and conference attendance from Nutricia. PH: lecture fees from Dr Falk. RH: speaker's fees, conference support or consultancy fees from Nutricia, Dr Falk, MSD Immunology and 4D Pharma. DCW: lecture fees, consultancy, travel support from AbbVie; consultancy from Takeda; and financial support for research by MSD. RKR: speaker's fees, travel support and participated in medical board meetings with AbbVie, Napp and Nestle. All other listed authors have no declared conflicts of interest.

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