EXTENDED REPORT

Randomised comparison of initial triple DMARD therapy with methotrexate monotherapy in combination with low-dose glucocorticoid bridging therapy; 1-year data of the tREACH trial

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ABSTRACT

Objectives To compare 1-year clinical efficacy of (1) initial triple disease-modifying antirheumatic drug therapy (ITDT) with initial methotrexate (MTX) monotherapy (iMM) and (2) different glucocorticoid (GC) bridging therapies: oral versus a single intramuscular injection in early rheumatoid arthritis.

Methods In a single-blinded randomised clinical trial patients were randomised into three arms: (A) iTDT (methotrexate+sulfasalazine+hydroxychloroquine) with GCs intramuscularly; (B) iTDT with an oral GC tapering scheme and (C) MTX with oral GCs similar to B. Primary outcomes were (1) area under the curve (AUC) of Health Assessment Questionnaire (HAQ) and Disease Activity Score (DAS) and (2) the proportion of patients with radiographic progression.

Results 281 patients were randomly assigned to arms A (n=91), B (n=93) or C (n=97). The AUC DAS and HAQ were respectively −2.39 (95% CI −4.77 to −0.00) and −1.67 (95% CI −3.35 to 0.02) lower in patients receiving iTDT than in those receiving iMM. After 3 months, treatment failure occurred less often in the iTDT group, resulting in 40% fewer treatment intensifications. The difference in treatment intensifications between the arms required to maintain the predefined treatment goal remained over time. No differences were seen between the two GC bridging therapies. Respectively 21%, 24% and 23% of patients in arms A, B and C had radiographic progression after 1 year. Patients receiving iTDT had more adjustments of their medication owing to adverse events than those receiving iMM.

Conclusions Treatment goals are attained more quickly and maintained with fewer treatment intensifications with iTDT than with iMM. However, no difference in radiographic progression is seen. Both GC bridging therapies are equally effective and, therefore, both can be used.

Trial registration number ISRCTN26791028.

INTRODUCTION

Recently EULAR updated the recommendations for the management of rheumatoid arthritis (RA). In patients with newly diagnosed RA the guidelines recommend that (1) the initial treatment strategy should encompass methotrexate (MTX) as monotherapy or in combination with other disease-modifying antirheumatic drugs (DMARDs), irrespective of glucocorticoids (GCs); (2) treatment is targeted to achieve remission, or low disease activity and (3) treatment should be adjusted if there is no improvement after 3 months or the target has not been reached by 6 months. When poor prognostic factors are present, biological agents should be considered if the initial treatment strategy has failed, or otherwise one could switch to another combination of DMARDs. Functional and radiological outcomes improve if current guidelines are upheld. Nevertheless, some major points for debate still exist.

First, 2010 criteria for RA are more and more often incorporated into daily practice. All current guidelines, however, were formulated using data from studies in patients fulfilling 1987 RA criteria. Thus trials comparing initial treatment strategies in the early phase of RA are needed for validation.

Second, several clinical trials concluded that initial combination therapy had better clinical efficacy than monotherapy; however, most rheumatologists have not implemented this in daily practice. Moreover, current guidelines do not recommend combination therapy for all patients with newly diagnosed RA. The principal motive for disregarding combination therapy was because (1) trials were biased by GCs, (2) patients were not DMARD naïve and (3) there were concerns about safety. Third, GCs have a rapid anti-inflammatory effect and are therefore used as bridging therapy to treat active disease in between initiation of DMARDs and the onset of their therapeutic effect. However, trials specifically comparing GC bridging therapies are sparse. More trials are, therefore, needed to investigate optimal dosage and tapering schemes.

Therefore, our aim was to compare in patients with very early RA the 1-year clinical efficacy of (1) initial triple DMARD therapy (iITDT) with initial MTX monotherapy (iMM), unbiased by GCs and (2) different GC bridging therapies: oral versus a single intramuscular injection.

PATIENTS AND METHODS

Patients

For this study, data were used from a clinical trial (ISRCTN26791028)—namely, treatment in the...
Randomisation and blinding

Patients were randomised into three groups A. iTDT (MTX, sulfasalazine and hydroxychloroquine) with GCs intramuscularly; B. iTDT with an oral GC tapering scheme; C. iMM with oral GCs similar to B. Concurrent treatment with non-steroidal anti-inflammatory drugs and intra-articular GC injections (maximum of two per 3 months) was allowed.

DMARD dosages were: MTX 25 mg/week orally (dosage reached after 3 weeks), sulfasalazine 2 g/day and hydroxychloroquine 400 mg/day (reduced to 200 mg/day after 3 months), GCs were given either intramuscularly (methylprednisolone 120 mg or triamcinolone 80 mg) or in oral tapering scheme (weeks 1–4: 15 mg/day, weeks 5–6: 10 mg/day, weeks 7–8: 5 mg/day and weeks 9–10: 2.5 mg/day). All patients received folic acid (10 mg/week) during MTX prescription. Osteoporosis prophylaxis (risendronate 35 mg/week and calcium/vitamin D combination 500/400 mg/IU/day) was given to patients in treatment arms B and C, during the first 3 months.

A treat-to-target approach was used, aiming for a DAS <2.4. If DAS was ≥2.4, medication was intensified. Intensification steps were in the order (1) MTX+etanercept (50 mg/week, subcutaneously), (2) MTX+adalimumab (40 mg/2 weeks, subcutaneously) and (3) MTX+abatacept (500–1000 mg/4 weeks, intravenously, dependent on weight). Treatment intensifications were the same for each treatment arm.

If DAS was <1.6 at two consecutive visits, medication was tapered. Hierarchically ordered tapering steps were: (1) biological agent, (2) sulfasalazine, (3) MTX and (4) hydroxychloroquine. Biological agent(s), MTX and sulfasalazine were gradually discontinued, whereas hydroxychloroquine was stopped immediately. A flare during tapering, defined as DAS ≥2.4, resulted in restarting full treatment, according to the stage in the protocol.

Outcomes and assessments

Patients were examined every 3 months for all outcomes, except for hand/foot radiographs, which were obtained at baseline and half-yearly.
Figure 1  Trial profile and protocol violations. Results are shown as number (%). Other reasons are: 3× no compliance, 1× pregnancy wish and 1× continuation of SASP after switch to etanercept. The figure shows the flowchart of the tREACH trial, whereas the table shows the protocol violations within the tREACH trial during the first year of follow-up. Other reasons for dropping out, in the flowchart, were incorrect randomisation and problems with communication. GCs, glucocorticoids; HCQ, hydroxychloroquine; IM, intramuscular; MTX, methotrexate; SASP, sulfasalazine; tREACH, treatment in the Rotterdam Early Arthritis Cohort.
All analyses were performed for patients in the high-probability stratum and two subgroups consisting of patients with RA according to 1987 and 2010 criteria. All statistical analyses were carried out using STATA V12.0. A p value <0.05 was considered statistically significant.

RESULTS

Patients

A total of 797 patients were assessed for eligibility and of those, 568 were included. In the high-probability stratum 281 patients were randomly assigned to treatment arm A (n=91), B (n=93) or C (n=97) (figure 1). Besides an mITT analysis, we also performed a per-protocol analysis. We excluded, from our per-protocol analysis, respectively 32 (35%), 37 (40%) and 32 (33%) patients randomised to arms A, B and C (figure 1). At baseline, the symptom duration and patients fulfilling 1987 criteria for RA differed significantly between arms (table 1).

Clinical outcome

The difference in AUC DAS between iTDT and iMM was −2.39 (95% CI −4.77 to −0.00, p=0.0497), and −0.91 (95% CI −3.17 to 1.34, p=0.42) between both GC bridging therapies. Adjusted differences were respectively −2.61 (95% CI −4.55 to −0.66, p=0.009) and −0.015 (95% CI −1.96 to 1.99, p=0.99). The largest difference in disease activity (states) between treatment arms was seen after 3 months, after which it gradually diminished (figure 2). After 12 months DAS was 0.08 (95% CI −0.34 to 0.19) lower in patients with iTDT than in those with iMM. Difference in DAS between the different GC bridging therapies was −0.20 (95% CI −0.45 to 0.04). Similar results were found in our multivariate analyses (data not shown). No differences in disease activity states were found after 12 months between iTDT and iMM, or between the two GC bridging therapies (table 2). DAS and its components over time per treatment arm are given in online supplement 4.

There was no significant difference in SHS after 12 months of treatment (table 2). Respectively 21%, 24% and 23% of patients in arms A, B and C had radiographic progression. The cumulative probability plots for the three treatment arms were superimposable (see online supplement 3).

Functional improvement was seen in all patients. Difference in AUC HAQ between iTDT and iMM was −1.67 (95% CI −3.35 to 0.02, p=0.052), and −0.46 (95% CI −2.04 to 1.12, p=0.57) between both GC bridging therapies (figure 2). Adjusted differences were, respectively, −1.30 (95% CI −2.45 to −0.14, p=0.028) and 0.33 (95% CI −0.84 to 1.51, p=0.38). No significant difference in functional ability was seen after 12 months (table 2). Secondary endpoints are shown in table 2 and figure 2.

We also performed a per-protocol analysis, which showed similar results (data not shown). The above-mentioned analyses were performed in both subgroups, and produced similar results (see online supplement 5 and 6).

Table 1   Baseline characteristics and clinical response after 12 months for each induction therapy group, according to intention-to-treat

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>A. MTX+SASP+HCQ+IM GCs (n=91)</th>
<th>B. MTX+SASP+HCQ+oral GCs (n=93)</th>
<th>C. MTX+oral GCs (n=97)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Demographic</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Age (years), mean (SD)*</td>
<td>53 (15)</td>
<td>54 (14)</td>
<td>54 (14)</td>
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<tr>
<td>Sex, female, n (%)</td>
<td>55 (60)</td>
<td>67 (72)</td>
<td>68 (70)</td>
</tr>
<tr>
<td>Disease characteristics</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Symptom duration (days), mean (SD)*</td>
<td>162 (98)</td>
<td>184 (92)</td>
<td>154 (83)</td>
</tr>
<tr>
<td>ACRA positive, n (%)</td>
<td>74 (81)</td>
<td>67 (72)</td>
<td>75 (77)</td>
</tr>
<tr>
<td>RF positive, n (%)</td>
<td>69 (76)</td>
<td>65 (70)</td>
<td>65 (67)</td>
</tr>
<tr>
<td>Fulfilment of RA criteria, n (%)</td>
<td>69 (76)</td>
<td>57 (61)</td>
<td>63 (65)</td>
</tr>
<tr>
<td>1987*</td>
<td>87 (96)</td>
<td>88 (95)</td>
<td>95 (98)</td>
</tr>
<tr>
<td>Disease activity</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DAS, mean (SD)</td>
<td>3.28 (0.82)</td>
<td>3.40 (1.07)</td>
<td>3.38 (0.97)</td>
</tr>
<tr>
<td>TJC44, median (IQR)</td>
<td>8 (4–14)</td>
<td>9 (5–15)</td>
<td>10 (4–14)</td>
</tr>
<tr>
<td>SJC44, median (IQR)</td>
<td>8 (5–12)</td>
<td>7 (4–12)</td>
<td>7 (4–12)</td>
</tr>
<tr>
<td>General health, median (IQR)†</td>
<td>52 (34–70)</td>
<td>55 (28–69)</td>
<td>53 (38–70)</td>
</tr>
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<td>ESR in mm/h, median (IQR)</td>
<td>27 (14–40)</td>
<td>22 (13–40)</td>
<td>24 (14–42)</td>
</tr>
<tr>
<td>CRP in mg/L, median (IQR)</td>
<td>8 (3.5–23)</td>
<td>6.5 (4.5–19)</td>
<td>11 (5–26)</td>
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<tr>
<td>Radiographs (hand/foot)</td>
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<td>Total SHS (0–488), median (IQR)</td>
<td>0.5 (0–2)</td>
<td>0.5 (0–2)</td>
<td>1 (0–2.5)</td>
</tr>
<tr>
<td>Erosion score (0–280), median (IQR)</td>
<td>0 (0–1)</td>
<td>0 (0–1)</td>
<td>0.5 (0–1)</td>
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<tr>
<td>JSN score (0–168), median (IQR)</td>
<td>0 (0–1)</td>
<td>0 (0–1)</td>
<td>0 (0–1.5)</td>
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<tr>
<td>Erosive disease, n (%)</td>
<td>10 (11)</td>
<td>6 (6)</td>
<td>8 (8)</td>
</tr>
<tr>
<td>Patient-reported outcomes¶</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HAQ, mean (SD)</td>
<td>0.98 (0.67) (n=84)</td>
<td>0.96 (0.64) (n=86)</td>
<td>1.06 (0.68) (n=92)</td>
</tr>
<tr>
<td>RADAI (0–10), mean (SD)</td>
<td>3.97 (1.83) (n=81)</td>
<td>3.94 (1.61) (n=82)</td>
<td>4.21 (1.82) (n=87)</td>
</tr>
</tbody>
</table>

* p=0.018 for B versus C.
† p=0.034 for A versus B.
‡ General health is measured with a Visual Analogue Scale from 0 to 100 mm.
§ Erosive disease is defined as having an erosion score >1 in three separate joints.
¶ Not everyone filled in a (complete) questionnaire and therefore n is different for HAQ and RADAI.

ACPA, anti-citrullinated protein/peptide antibodies; CRP, C-reactive protein; DAS, Disease Activity Score; ESR, erythrocyte sedimentation rate; GCs, glucocorticoids; HAQ, Health Assessment Questionnaire; HCQ, hydroxychloroquine; IM, intramuscular; JSN, joint space narrowing; MTX, methotrexate; RA, rheumatoid arthritis; RADAI, RA Disease Activity Index questionnaire; RF, rheumatoid factor; SASP, sulfasalazine; SHS, modified Sharp–Van der Heijde score; SJC44, swollen joint count (44 joints); TJC44, tender joint count (44 joints).
Medication
After 3 months 40% fewer biological agents were prescribed in the iTDT group than in the iMM group. This difference remained over time (figure 3). After 12 months, respectively 27% and 43% of patients, receiving iTDT and iMM, were using a biological agent (p=0.03). Moreover, for more patients receiving iMM their first biological agent had failed (16% vs 6%, p=0.03) (figure 3). In 117/281 (42%) of patients treatment could be tapered at one or more time points, and of those, 14/117 (12%) flared. Treatment could be tapered in all treatment arms, without differences in flare rates (figure 3). Biological usage did not differ between the two GC bridging therapies (figure 3). This analysis was also performed in both subgroups and showed similar results (see online supplements 5 and 6).

Adverse events (AEs)
No differences in serious AEs were seen (table 3). However, the proportion of patients with medication adjustments due to AEs differed significantly between iTDT and iMM (60/93 (65%) and 44/97 (45%), p=0.008). Besides switching to MTX subcutaneously, these differences vanished after stratification for drug (table 3). No differences were seen between the two GC bridging therapies. Most treatment adjustments occurred in the first 3 months (51/159, 32%). Among all patients gastrointestinal complaints and fatigue were the most commonly reported AEs, respectively 56% and 36% (table 3).

DISCUSSION
In this study, unbiased for GCs, we showed that the AUC DAS was significantly less in the iTDT group than in the iMM group. A trend was observed for the AUC HAQ. Treatment goals were attained more quickly and maintained with 40% fewer treatment intensifications in the iTDT group. Moreover, for more patients receiving iMM their first biological agent had failed, reducing therapeutic options. Disease activity, functional ability and radiographic progression, after 12 months, did not differ between each other.
Treatment alterations, including the prescription of 40% fewer biological agents. However, no differences in radiographic progression were seen, which was probably owing to early initiation of intensive treatment, resulting in less joint destruction and thus less radiological progression. Moreover, the early initiation of biological agents in the iMM group might have prevented/delayed the radiographic progression.

In our trial switching to biological agents was possible after 3 months, if the target had not been reached. In the iMM group one might argue that triple DMARD therapy rather than biological agents should be the first step-up, especially since there is no evidence that step-up to a biological agent rather than triple DMARD therapy results in better clinical efficacy. Moreover, the updated EULAR guidelines only recommend switching to biological agents after 3 months if (1) there is no improvement in disease activity and (2) poor prognostic factors are present (ie, autoantibody positivity and erosive disease).

Twenty of the 36 (56%) patients for whom iMM treatment had failed after 3 months were EULAR non-responders, defined as
having a DAS >2.4 and <0.6 decline in DAS from baseline, with poor prognostic factors. Therefore, we think intensification to biological agents after iMM treatment had failed was reasonable.

For economic reasons efficient use of biological agents is needed to be able to continue optimal rheumatic care in the future.29 With iTDT treatment, goals are attained more quickly and maintained with 40% fewer biological agents, reducing costs enormously. Better disease control improves worker productivity, and thus costs due to loss of productivity are also reduced.30 31 However, the cost–utility analysis of the tREACH trial still has to reconfirm this statement.

Patients and/or rheumatologists, however, may be averse to iTDT treatment, mainly because of the large amount of drugs that have to be taken. Medication adherence in RA is strongly influenced by a patient’s belief about the need for the drugs.32 These beliefs are moulded by rheumatologists through the information given about the disease and treatment approach.32 A personalised medicine approach would be ideal in this very early phase, especially since 60% respond well to iMM; conversely, therefore, some patients receiving iTDT are overtreated. Determination of early GC response after treatment initiation is a promising predictor, possibly leading to a more personalised medicine approach.33 Therefore, we think that future research should focus on developing a more personalised treatment approach, in which differentiation between patients who would thrive on iMM and those who need iTDT might be a first step.

In 42% of patients treatment could be tapered, and of those, 12% flared. Therefore, we think tapering DMARDs and/or biological agents is justified in patients with sustained remission. However, patients should still be monitored strictly during tapering. Data on tapering medication are sparse, especially in early RA.1 34 Future research is needed to determine (1) when to start tapering, (2) how to taper and (3) the optimal interval between taperings.

We found that intramuscular and oral GCs are equally effective as bridging therapy, but one single injection might be more feasible. However, duration of our GC tapering scheme was short (10 weeks) and the initial dosage was low (15 mg) in comparison with, for example, the COBRA regimen (respectively 28 weeks and 60 mg).9 Because GCs have disease-modifying traits with longlasting benefits even after withdrawal, a different GC oral tapering scheme might be superior.13 However, if we compare our iMM treatment with the COBRA-light strategy, intensification to biological agents after 6 months is equally indicated (respectively 36% and 41%).35 In the COBRA-light trial, however, the treatment goal is remission instead of low disease activity, which prevents useful direct comparison.35 Therefore, future research is needed for optimising GC bridging therapies.

Our study had certain limitations. Foremost, baseline imbalances occurred, despite randomisation, which is why we also performed an adjusted analysis. After adjustment similar results were found for the primary outcomes, but the difference in AUC HAQ became significant (favouring iTDT). Additionally, only research nurses, who assessed the DAS, were blinded to the allocated treatment arm. This design was chosen, since we wanted to mimic daily practice as far as possible. Single blinding, however, might be a potential source of bias, because of the aversion for iTDT by rheumatologists and/or patients (favouring iMM) or (un)intentional misinformation due to the

Figure 3  Withdrawal, flares and medication usage over time and after 12 months, stratified for induction therapy. Results are shown as number (%) unless stated otherwise. Other biological agents are: infliximab (A) and rituximab (B). Treatment could be tapered after 6 months. Therefore the total amount of possible taperings is the sum of all assessments at the last three visits per treatment arm. A flare is defined as a Disease Activity Score ≥2.4. The proportion is calculated by dividing the number of flares by the total number of taperings. *p=0.011 for B versus C. †p=0.031 for B versus C. ‡p=0.028 for A versus B. GCs, glucocorticoids; HCQ, hydroxychloroquine; IM, intramuscular; MTX, methotrexate; SASP, sulfasalazine.
rheumatologist’s beliefs (favouring iTDT). Moreover, for various reasons 101 (36%) patients were excluded from our per-protocol analysis, which was more than expected. Our exclusion percentage, however, was comparable with that of other trials.9

In conclusion, in our treat-to-target design, treatment goals were attained more quickly and maintained with fewer treatment intensifications, with iTDT than with iMM. However, no difference was seen in radiographic progression. Before choosing the initial treatment strategy, rheumatologists should be aware of the benefits and risks, but additionally, known prognostic factors and the patient’s wish should be taken into account. One single intramuscular GC injection and a low-dose oral GC tapering scheme would be sufficient as bridging therapy.


Contributors All authors contributed to the conception or design of the trial; or the acquisition, analysis, or interpretation of data for the work; drafting or revision of the manuscript; and final approval of the manuscript for publication.

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Competing interests None.

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Provenance and peer review Not commissioned; externally peer reviewed.

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