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

P H de Jong,1 J M Hazes,1 H K Han,2 M Huisman,3 D van Zeaben,3 P A van der Lubbe,4 A H Gerards,4 B van Schaeybroeck,5 P B de Sonnaville,6 M V van Krugten,6 J J Luime,1 A E Weel1,2

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).1 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 a(nother) combination of DMARDs. Functional and radiological outcomes improve if current guidelines are upheld.1 2 Nevertheless, some major points for debate still exist.

First, 2010 criteria for RA3 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.1 4 5 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.6–10 Moreover, current guidelines do not recommend combination therapy for all patients with newly diagnosed RA.1 4 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.11 12

Third, GCs have a rapid anti-inflammatory effect and are therefore used as bridging therapy to treat active disease in between initiation of DMARD(s) and the onset of their therapeutic effect.13 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 (iTDT) 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
If DAS was gradually discontinued, whereas hydroxychloroquine was the logical agent, (2) sulfasalazine, (3) MTX and (4) hydroxychloroquine was tapered. Hierarchically ordered tapering steps were: (1) biologic 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.

Randomisation and blinding

Patients were randomised, using variable block randomisation stratified for centre, by an independent call-centre. Trained research nurses, blinded to the allocated treatment arm throughout the study, examined patients and calculated the Disease Activity Score (DAS).

Design

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 an 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 1000 mg/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, depending 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) biologic agent, (2) sulfasalazine, (3) MTX and (4) hydroxychloroquine. Biologic 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.

Primary outcomes were (1) area under the curve (AUC) Health Assessment Questionnaire (HAQ) and DAS and (2) proportion of patients with radiographic progression. Secondary endpoints were disease activity (state), functional ability, EULAR response criteria,17 Boolean-defined remission criteria,18 self-assessed disease activity and medication usage over time and after 12 months of treatment.

DAS and its thresholds are used for disease state categorisation.16 Functional ability is measured with the HAQ.19 Higher HAQ scores indicate poorer function. Radiographic progression was measured with the modified Sharp–van der Heijde score (SHS).20 Radiographs were read chronologically by two out of five qualified assessors, who were blinded to the patient’s identity and treatment allocation.21 Mean SHS are reported.22 Weighted k between assessors was 0.36 with 98% agreement. The proportion of patients with radiographic progression, defined as SHS change >0.5 and >1.2 (the smallest detectable change) and >1.2 a year, was also calculated.23 EULAR response criteria are based on attained level and change in DAS (see online supplement 2).15 Boolean remission criteria are defined as having a tender joint count, swollen joint count, C-reactive protein (in mg/dL) and patient global assessment (0–10 scale) of ≤1.15 Self-assessed disease activity is measured with the RA Disease Activity Index questionnaire (RADA1).21 Higher RADA1 scores indicate more active disease.

Safety monitoring and toxicity

Safety monitoring occurred according to Dutch guidelines,24 which included laboratory tests at fixed intervals. Study medication was either stopped or the dosage lowered in accordance with the protocol if serious adverse events were seen by the attending rheumatologist. MTX could be given subcutaneously if patients had gastrointestinal complaints. If MTX had to be stopped for safety reasons, leflunomide (20 mg/day) was substituted.14

Statistical analysis

Sample-size calculation was based upon AUC HAQ, using data from the BeSt study7 where mean AUC HAQ of combination therapy and monotherapy respectively were 7.7 (SD 5.5) and 10.5 (SD 7.4). A target sample size of 270 patients per probability stratum and thus 90 patients per arm, was needed to detect the mentioned difference with a power of 80% and two-sided α=0.05. This size is sufficient to detect a difference of 6.1 AUC DAS and 20% difference in radiographic progression.14

Clinical efficacy was calculated in an intention-to-treat (mITT) and per-protocol analysis. In mITT analyses patients are analysed in the groups to which they were randomised, regardless of whether they received or adhered to the allocated intervention. For the primary, but not the secondary, outcomes missing values are imputed. Statistical comparison of the baseline characteristics and outcomes (after 12 months) between iTDT and iMM (arms B vs C) and both GC bridging therapies (arms A vs B) were made by Student t test, χ² test or Wilcoxon rank-sum test, as appropriate.

We used the AUC to compare DAS and HAQ over time between treatment arms, in which missing values at each time point were substituted with the mean value of the corresponding treatment arm. Radiographic progression was extrapolated or interpolated if the SHS was missing after 12 months.

We also performed adjusted analyses for our primary outcomes, in which we corrected for baseline imbalances, rheumatoid factor, anti-citrullinated protein/peptide antibodies and baseline HAQ, DAS or SHS, as appropriate, using multivariate analyses.
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.


Clinical and epidemiological research

Assessed for eligibility (n=797)
- Declined to participate n=115
- Not meeting criteria n=59
- Not eligible due to recruitment stop n=30
- Lost to follow-up n=25

Selected patients with high probability

Randomized (n=568)
- High n=281
- Intermediate n=243
- Low n=44

High probability population (n=281)

A. MTX + SASP + HCQ + im GCs (n=91)
- Drop out: n=3
  - Adverse event (n=1)
  - Other* (n=2)
  - Time-point skipped: n=3

Participants n=85
- Drop out: n=4
  - Patient refusal (n=4)
  - No compliance (n=1)

Participants n=81
- Drop out: n=4
  - Patient refusal (n=1)
  - No compliance (n=2)
  - Lung cancer (n=1)

Participants n=80
- Drop out: n=3
  - Patient refusal (n=3)

Participants n=77

B. MTX + SASP + HCQ + oral GCs (n=93)
- Drop out: n=3
  - Patient refusal (n=3)
  - Time-point skipped: n=1

Participants n=89
- Drop out: n=1
  - Patient refusal (n=1)
  - Time-point skipped: n=3

Participants n=86
- Drop out: n=1
  - Patient refusal (n=1)
  - No compliance (n=1)
  - Time-point skipped: n=2

Participants n=85

C. MTX + oral GCs (n=97)
- Drop out: n=6
  - Patient refusal (n=3)
  - Adverse event (n=2)
  - Time-point skipped: n=2

Participants n=90
- Drop out: n=2
  - Patient refusal (n=1)
  - No compliance (n=1)
  - Time-point skipped: n=2

Participants n=88
- Drop out: n=1
  - Patient refusal (n=1)
  - Deceased (n=1)

Participants n=87

Protocol violations
- Skipped ≥1 time point(s)
  - A. MTX + SASP + HCQ + im GCs (n=91) 16 (18)
  - B. MTX + SASP + HCQ + oral GCs (n=93) 12 (13)
  - C. MTX + oral GCs (n=97) 14 (14)
- Postponing/withholding biologicals
  - A. MTX + SASP + HCQ + im GCs (n=91) 7 (8)
  - B. MTX + SASP + HCQ + oral GCs (n=93) 11 (12)
  - C. MTX + oral GCs (n=97) 11 (11)
- Illicit treatment intensifications
  - A. MTX + SASP + HCQ + im GCs (n=91) 1 (1)
  - B. MTX + SASP + HCQ + oral GCs (n=93) 1 (1)
  - C. MTX + oral GCs (n=97) 1 (1)
- No tapering of treatment
  - A. MTX + SASP + HCQ + im GCs (n=91) 6 (7)
  - B. MTX + SASP + HCQ + oral GCs (n=93) 11 (12)
  - C. MTX + oral GCs (n=97) 5 (5)
- Other
  - A. MTX + SASP + HCQ + im GCs (n=91) 2 (2)
  - B. MTX + SASP + HCQ + oral GCs (n=93) 2 (2)
  - C. MTX + oral GCs (n=97) 1 (1)
- Total
  - A. MTX + SASP + HCQ + im GCs (n=91) 32 (35)
  - B. MTX + SASP + HCQ + oral GCs (n=93) 37 (40)
  - C. MTX + oral GCs (n=97) 32 (33)
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 V. 12.0. 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=0.58). 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>
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<tbody>
<tr>
<td>Demographic</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (years), mean (SD)</td>
<td>53 (15)</td>
<td>54 (14)</td>
<td>54 (14)</td>
</tr>
<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, median (IQR)</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>
<tr>
<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>
</tr>
<tr>
<td>Radiographs (hand/foot)</td>
<td></td>
<td></td>
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<tr>
<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>
</tr>
<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>
</tr>
<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, median (SD)</td>
<td>0.98 (0.67)</td>
<td>0.96 (0.64)</td>
<td>1.06 (0.68)</td>
</tr>
<tr>
<td>RADAI (0–10), mean (SD)</td>
<td>3.97 (1.83)</td>
<td>3.94 (1.61)</td>
<td>4.21 (1.82)</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
treatment alterations, including the prescription of 40% fewer medications due to AEs than the iMM group. Therefore, it is not the intensity, but the 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 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.

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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. 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. However, the cost–utility analysis of the tREACH trial still has to confirm 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. These beliefs are moulded by rheumatologists through the information given about the disease and treatment approach. 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. 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. 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). Because GCs have disease-modifying traits with longlasting benefits even after withdrawal, a different GC oral tapering scheme might be superior. 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%). In the COBRA-light trial, however, the treatment goal is remission instead of low disease activity, which prevents useful direct comparison. 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.

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.


Competing interests None.

Ethics approval Ethics committee of the Erasmus MC.

Provenance and peer review Not commissioned; externally peer reviewed.

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7. Goeicoe-Ruitman JP, de Vries-Bouwstra JK, Aalcaert CF, et al. Clinical and radiographic outcomes of four different treatment strategies in patients with early 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 was comparable with that of other trials.10 27

Table 3 Number (%) of patients with (serious) adverse events and treatment alterations due to side effects for each induction therapy group

<table>
<thead>
<tr>
<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>Adverse events (AEs)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Serious AEs*</td>
<td>5 (5)</td>
<td>10 (11)</td>
</tr>
<tr>
<td>Patients with ≥1 AEs</td>
<td>76 (84)</td>
<td>82 (88)</td>
</tr>
<tr>
<td>No. of AEs per patient, median (IQR)</td>
<td>2 (1–3)</td>
<td>2 (1–4)</td>
</tr>
<tr>
<td>Medication changes due to AEs</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Switch to MTX SC†</td>
<td>12 (13)</td>
<td>21 (23)</td>
</tr>
<tr>
<td>Lowering MTX dosage &lt;20 mg/week‡</td>
<td>17 (19)</td>
<td>10 (11)</td>
</tr>
<tr>
<td>Stop MTX</td>
<td>11 (12)</td>
<td>14 (15)</td>
</tr>
<tr>
<td>Stop SASP</td>
<td>11 (12)</td>
<td>8 (9)</td>
</tr>
<tr>
<td>Stop HCQ</td>
<td>4 (4)</td>
<td>5 (5)</td>
</tr>
<tr>
<td>Stop biological agents§</td>
<td>0 (0)</td>
<td>2 (2)</td>
</tr>
</tbody>
</table>

Observed AE¶

| | | |
| Malaise | 20 (22) | 19 (20) | 15 (15) |
| Fatigue | 23 (25) | 34 (37) | 40 (41) |
| Dizziness | 2 (2) | 10 (11) | 7 (7) |
| Headache | 10 (11) | 13 (14) | 13 (13) |
| Muscle weakness | 2 (2) | 8 (9) | 7 (7) |
| Hypertension | 2 (2) | 4 (4) | 0 (0) |
| Palpitations | 0 (0) | 4 (4) | 7 (7) |
| Oedema | 3 (3) | 3 (3) | 6 (6) |
| Dyspnoea | 0 (0) | 4 (4) | 7 (7) |
| Gastrointestinal complaints | 57 (63) | 59 (63) | 41 (42) |

Results shown are a number (%) unless stated otherwise.

*Serious AEs per treatment are respectively: arm (A) 4× hospitalisation (2× pneumonia, kidney stones and inguinal hernia surgery), 1× lung carcinoma; arm (B) 5× hospitalisation (MTX pneumonitis, severe constipation, transient ischaemic attack, gastroenteritis and observation chest pain), 1× deceased; arm (C) 6× hospitalisation (pneumonia, blood transfusion, syncope, cholecystectomy, inguinal hernia surgery and acute rheumatoid arthritis), 1× myocardial infarction, 2× colon carcinoma and 1× maculopathy.

†p=0.039 for B versus C.
‡p=0.028 for B versus C.
§The reasons for stopping biological agents are positive tuberculosis screening (3× in arm C), prostate carcinoma (1× arm B), allergic reaction (1× arm C) and recurrent infections (1× arm B).
¶Bone marrow depression is defined as an anaemia, thrombocytopenia or leucopenia with respectively a haemoglobin level, platelet count and white blood cells count below the lower limit of the normal range. High creatinine, raised liver enzymes and hyperglycaemia are defined as having respectively a creatinine, liver transaminases and glucose level above the upper limit of the normal range.GCs, glucocorticoids; HCQ, hydroxychloroquine; IM, intramuscular; MTX, methotrexate; SASP, sulfasalazine; SC, subcutaneous.
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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Ann Rheum Dis 2014 73: 1331-1339 originally published online May 1, 2014
doi: 10.1136/annrheumdis-2013-204788