Original article

Achieving comprehensive disease control in patients with early and established rheumatoid arthritis treated with adalimumab plus methotrexate versus methotrexate alone

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Abstract

Objective To evaluate the achievement of comprehensive disease control (CDC) following 1 year of treatment with adalimumab+methotrexate versus methotrexate alone and whether early achievement of remission (at week 24 or 26) is associated with CDC at week 52 in patients with either early or established rheumatoid arthritis (RA).

Methods Post hoc analyses were conducted in three clinical studies assessing treatment with adalimumab+methotrexate: DE019 (NCT00195702) enrolled patients with early RA who were methotrexate inadequate responders; OPTIMA (NCT00420927) and PREMIER (NCT00195663) enrolled methotrexate-naive patients with early RA. In OPTIMA, patients not achieving stable low disease activity at weeks 22 and 26 in the placebo+methotrexate group could receive open-label adalimumab+methotrexate for 52 weeks (Rescue ADA arm). CDC was defined as the simultaneous achievement of clinical remission (DAS28(CRP)<2.6), normal function (HAQ-DI<0.5) and absence of radiographic progression (∆mTSS<0.5).

Results Regardless of disease duration, significantly more patients receiving adalimumab+methotrexate achieved CDC compared with methotrexate alone. In the adalimumab+methotrexate group, a numerically greater proportion of patients with early RA (~25%) versus established RA (14%) achieved CDC at 1 year; achievement of CDC was notably greater among patients who met criteria for remission at week 24 or 26 (~50% of patients with early RA and 39% with established RA).

Conclusion Treatment with adalimumab+methotrexate increases the likelihood of achieving CDC in patients with either early or established RA. Clinical remission at week 24 or 26 is associated with achievement of CDC at week 52.

Trial registration number DE019 (NCT00195702), OPTIMA (NCT00420927), PREMIER (NCT00195663); Post-results.

Key messages

What is already known about this subject?

► Clinical remission is the primary goal of treating rheumatoid arthritis (RA); comprehensive disease control (CDC) additionally encompasses preservation of physical function and prevention of structural damage progression and is a realistic therapeutic goal in patients with early RA.

► It has been unclear whether achieving clinical remission can predict subsequent CDC.

What does this study add?

► In this analysis of three studies, clinical remission at 24 or 26 weeks was associated with achievement of CDC at 52 weeks.

► Combination treatment with adalimumab+methotrexate (vs methotrexate alone) increased the likelihood of achieving CDC in patients with either early or established RA.

How might this impact on clinical practice?

► These findings may help guide treatment decisions, including the utility of combination treatment in patients with early and established RA.

Introduction

Rheumatoid arthritis (RA) is a chronic and debilitating inflammatory disease with a population prevalence of approximately 1%.1 Progressive inflammation in the joints of patients can lead to irreversible joint damage2,3 and diminished function and disability.4 The primary goal of treating RA is to achieve a target of clinical remission, thereby maximising quality of life. Early diagnosis and treatment are important for achieving the elements of comprehensive disease control (CDC), including normalisation of physical function, achievement of clinical remission or low disease activity (LDA) and absence of structural damage progression through effective suppression of inflammation.5–7
CDC is unlikely to be achievable in a large proportion of patients with established RA because of increasing functional impairment, although it should constitute a realistic therapeutic goal in patients with early RA as suggested by treat-to-target recommendations, systematic literature reviews, and clinical studies. Treatment recommendations suggest initial treatment of active disease with methotrexate (MTX) monotherapy; after 6 months of unsuccessful treatment, a biological disease-modifying anti-rheumatic drug (DMARD), such as a tumour necrosis factor (TNF) inhibitor, should be added to the therapy of patients with poor prognostic factors. Although there is evidence that early response or remission in RA is strongly predictive of subsequent long-term remission, it is not clear to what extent clinical remission of RA can predict subsequent achievement of CDC.

The post hoc analyses detailed in this report aim to (1) evaluate attainment of CDC after 1 year of treatment with the combination of adalimumab+MTX and (2) assess whether the achievement of clinical remission at week 24 or 26 is associated with CDC at week 52 in three randomised controlled trials in patients with early or established RA.

**PATIENTS AND METHODS**

**Patients**

Patient data originated from three randomised clinical trials in RA: DE019 (NCT00195702), PREMIER (NCT00195663), and OPTIMA (NCT00420927). All of the studies enrolled adults aged ≥18 years with a diagnosis of active RA as defined by the 1987 revised criteria of the American College of Rheumatology (formerly American Rheumatism Association). As noted in the original publications, an institutional review board or ethics committee approved the studies at each site.

DE019 was a 1-year, phase 3, randomised, double-blind, placebo-controlled trial of adalimumab in patients with RA who had active disease despite a minimum of 3 months of treatment with MTX (MTX inadequate response (MTX-IR)) at a stable weekly dose of 12.5–25 mg (or ≥10 mg/week in those intolerant to higher doses of MTX) for ≥4 weeks. In addition, patients were required to have a swollen joint count based on 66 assessed joints (SJC66) ≥6, tender joint count based on 68 assessed joints (TJC68) ≥9, C-reactive protein (CRP) ≥1.0 mg/dL, and either had to be rheumatoid factor (RF) positive or have ≥1 joint erosion on X-ray. Traditional DMARDs other than MTX were discontinued at least 28 days before baseline evaluations.

PREMIER was a 2-year, phase 3, randomised, double-blind, active comparator–controlled trial of adalimumab in an MTX-naive population with RA disease duration <3 years. Patients were required to have SJC66≥8, TJC68≥10, either an erythrocyte sedimentation rate (ESR) of ≥28 mm/hour or CRP≥1.5 mg/dL, and either had to be RF positive or have ≥1 joint erosion on X-ray. Patients who had received treatment with MTX, cyclophosphamide, cyclosporine, azathioprine or ≥2 other DMARDS were excluded.

OPTIMA (Optimal Protocol for Treatment Initiation with MTX and Adalimumab) was a 78-week, phase 4, double-blind, randomised, placebo-controlled trial of adalimumab in patients with early RA with a duration ≤1 year and moderate or high disease activity not previously treated with MTX. Patients were required to have SJC66≥6, TJC68≥28, either an ESR of ≥28 mm/hour or CRP≥1.5 mg/dL, and either had to be RF positive, anticyclic citrullinated peptide positive, or have ≥1 joint erosion on X-ray. Patients who had received previous MTX or other biological treatment were excluded.

**Treatment regimens**

In DE019, patients were randomly assigned 1:1:1 to receive adalimumab 40 mg subcutaneously every other week (with placebo injections on alternate weeks), adalimumab 20 mg subcutaneously weekly (not included in this analysis) or placebo subcutaneously weekly. The patient’s dose of concomitant MTX at baseline remained constant during the 52-week study; if toxicity occurred, the dose could be reduced to not lower than 10 mg weekly.

In PREMIER, patients were randomised 1:1:1 to receive adalimumab 40 mg subcutaneously every other week plus weekly oral MTX, adalimumab 40 mg monotherapy every other week plus weekly placebo (not included in this analysis) or weekly MTX plus placebo. MTX therapy was initiated at a weekly dose of 7.5 mg for the first 4 weeks of the study, 15 mg weekly during weeks 4–8, then 20 mg weekly at week 9.

In OPTIMA, patients were randomised 1:1:1 to receive adalimumab 40 mg subcutaneously every other week plus weekly MTX, initiated at a dose of 7.5 mg and rapidly titrated to 20 mg by week 8, or placebo plus MTX for 26 weeks (period 1). On the basis of achievement of a stable LDA target (28-joint Disease Activity Score based on CRP (DAS28(CRP)) ≤3.2 at weeks 22 and 26), placebo+MTX patients not achieving stable LDA (MTX-IR patients) received open-label adalimumab+MTX during period 2 (Rescue ADA arm) for an additional 52 weeks. This analysis included patients in the Rescue ADA population who represent patients with early RA and MTX-IR.

**Clinical, functional and radiographic assessments of CDC**

CDC was considered the simultaneous achievement of (1) clinical remission, (2) normal physical function and (3) absence of radiographic progression. Clinical signs and symptoms of inflammation were assessed with the DAS28(CRP) or its components. The DAS28(CRP) is composed of four outcome variables: TJC and SJC (each based on 28-joint assessment), CRP and patient’s global assessment of disease activity on a visual analogue scale (VAS; 0–10 cm). Clinical remission was defined as DAS28(CRP)≤2.6 (OPTIMA and PREMIER trials). Additional post hoc analyses used the Simplified Disease Activity Index (SDAI), which is the sum of five outcome variables: TJC and SJC (each based on 28-joint assessment), patient and physician global...
assessment of disease activity (VAS 0–10 cm) and CRP; a score of ≤3.3 indicates remission.25 26 In patients with established RA (DE019 trial), achievement of LDA (defined as DAS28(CRP)≤3.2)23 27 28 rather than remission was used to define the clinical assessment component of CDC.

Physical function was assessed using the Health Assessment Questionnaire Disability Index (HAQ-DI; score range, 0–3).29 Normal physical function was defined as HAQ-DI<0.5.30 31 To quantify structural joint damage, radiographs of the hands/wrists and feet were obtained and scored using the modified total Sharp score (mTSS) method.32–34 Radiographic progression of structural joint damage was assessed as the change from baseline (or from week 24/26) in mTSS (ΔmTSS). Absence of radiographic progression was defined as ΔmTSS≤0.50.35

Statistical analyses

The intent-to-treat population included all patients who were randomised and received ≥1 dose of study drug. These post hoc analyses assessed the proportion of patients achieving CDC as well as its individual components following 1 year of treatment with adalimumab+MTX or placebo+MTX in DE019 and PREMIER, and open-label adalimumab+MTX in the Rescue ADA arm of OPTIMA. To explore which component may be a limiting factor to achievement of CDC, frequencies of patients achieving DAS28(CRP)<2.6 (or DAS28(CRP)≤3.2), HAQ-DI<0.5, or ΔmTSS≤0.5, and combinations of the targets were determined.

Last observation carried forward (LOCF) was used to replace missing clinical and functional data for patients with postbaseline assessments; non-responder imputation (NRI) was used for patients without any postbaseline assessments. Missing Sharp scores were imputed by linear extrapolation for patients with baseline and postbaseline X-rays. Patients without a postbaseline X-ray were considered as showing radiographic progression. Sensitivity analyses for radiographic non-progression and CDC (DAS28(CRP)<2.6 or ≤3.2), HAQ-DI<0.5 and ΔmTSS<0.5) based on observed data and LOCF and NRI was used for patients without any postbaseline assessment were used to assess the effect of missing data.

Logistic regression was used to assess the treatment effect on radiographic non-progression and CDC based on DAS28(CRP)<2.6 or ≤3.2. Generalised estimating equation was used to model the time-repeated measurement of LDA, remission and normal function. Fisher’s exact test was used to assess treatment effect on CDC based on SDAI≤3.3. There was no adjustment for multiple testing. Venn diagramming was used to display the combinations of clinical remission or LDA (yes/no), normal function (yes/no) and radiographic non-progression (yes/no) for CDC.

RESULTS

Baseline characteristics: overall study populations

Baseline demographics and disease characteristics for patients receiving adalimumab+MTX or placebo+MTX in DE019 and PREMIER were indicative of patients with established and early active RA, respectively (table 1). Despite an inadequate response to MTX at month 6, the majority of patients in the Rescue ADA arm of OPTIMA were in a moderate disease activity state at the time of open-label adalimumab initiation.

Achievement of CDC at 1 year: overall patient populations

In MTX-IR patients with established RA (DE019), CDC based on DAS28(CRP)<2.6 was achieved in 14% (n/N=28/207) of patients in the adalimumab+MTX arm compared with 3% (n/N=5/200) in the placebo+MTX arm (p<0.001; CDC based on SDAI≤3.3 was achieved in 5% (n/N=11/207) and 2% (n/N=4/207), respectively (figure 1A). When LDA (DAS28(CRP)≤3.2) was used for this population of patients with established disease, 19% (n/N=40/207) of patients receiving adalimumab+MTX achieved a CDC state at 1 year versus 5% (n/N=10/200) of patients receiving placebo+MTX (p=0.001 and p=0.45; figure 1A).

In patients with early RA (PREMIER), CDC based on DAS28(CRP)<2.6 was achieved in 25% (n/N=67/268) of patients in the adalimumab+MTX arm compared with 9% (n/N=22/257) in the placebo+MTX arm (p<0.001; figure 1B). Similarly, in patients with early aggressive RA who had inadequate response to MTX during the first 6 months of therapy with placebo+MTX (OPTIMA), CDC was achieved in 24% (n/N=82/348) of patients after 1 year of open-label adalimumab+MTX treatment (figure 1C). Similar trends were observed when CDC was based on LDA or SDAI remission (figure 1B,C).

Consistent with the comparative treatment data for CDC in DE019 and PREMIER, a greater percentage of patients in the adalimumab+MTX arms achieved each of the individual components for CDC (online supplementary figures 1 and 2). In the OPTIMA study, open-label adalimumab+MTX resulted in rapid improvements in clinical and functional components, with normal function being the more difficult target to reach (online supplementary figure 3).

Responder analysis: achievement of CDC and individual components

Post hoc analysis examining the frequency of responders to individual components of CDC (with the clinical component of remission based on DAS28(CRP)<2.6) is presented in Venn diagrams (figure 2). In addition to patients who achieved CDC by this definition, 6% (n/N=13/207) of patients in the adalimumab+MTX arm satisfied the remission and radiographic non-progression criteria at year 1 vs 2% (n/N=3/200) of patients in the placebo+MTX arm in the DE019 study; an additional 2% (n/N=5/207) and 1% (n/N=2/200) of patients in each treatment arm, respectively, also satisfied the remission and normal function criteria (figure 2A,B). In the PREMIER study, an additional 10% (n/N=28/268) of patients in the adalimumab+MTX arm and 10% (n/N=25/257) in the placebo+MTX arm achieved remission and normal function criteria, and 7% (n/N=19/268)
and 2% (n/N=4/257), respectively, achieved remission and radiographic non-progression criteria at year 1 (figure 2C,D). Finally, in the OPTIMA study, an additional 16% (n/N=54/348) satisfied the remission and radiographic non-progression criteria and 3% (n/N=10/348) achieved remission and normal function criteria at year 1 (figure 2E). More patients achieved the clinical component of CDC when using DAS28(CRP) ≤ 3.2 (online supplementary figure 4) and fewer patients achieved the clinical component when using SDAI ≤ 3.3 (figure 3), shifting the distribution of responses.

Characteristics of patients achieving remission at week 24 or 26

In the comparative studies (PREMIER (MTX-naive) and DE019 (MTX-IR)), patients receiving adalimumab+MTX were significantly more likely to achieve clinical remission at 24 or 26 weeks based on DAS28(CRP)<2.6 than patients receiving placebo+MTX (table 2). A comparison of remission criteria revealed that at week 24 or 26, more patients achieved a DAS28(CRP) score <2.6 than SDAI≤3.3 in all three studies (table 2). In line with previous observations, the mean values for DAS28(CRP), mTSS and HAQ-DI for patients in clinical remission at 6 months were generally slightly higher for patients classified as being in remission by DAS28(CRP) versus SDAI criteria, but this did not reach statistical difference.

Achievement of CDC at 1 year: patients meeting criteria for remission at week 24 or 26

In the DE019 study, 39% of patients in the adalimumab+MTX arm and 33% in the placebo+MTX arm who met criteria for remission (DAS28(CRP)<2.6) at week 24 achieved CDC at 1 year (figure 4A) versus 14% and 3% in the overall population (figure 1A). Assessing individual components of CDC, a numerically greater proportion of patients achieved DAS28(CRP)<2.6 and ΔmTSS≤0.5 with adalimumab+MTX compared with placebo+MTX; however, achievement of HAQ-DI<0.5 was comparable between the treatment groups. In the PREMIER study, approximately 50% of patients who met criteria for remission at week 26 achieved CDC at 1 year, regardless of treatment arm (vs 25% and 9% in the overall population; figures 1B and 4B). A numerically greater percentage of patients receiving adalimumab+MTX achieved DAS28(CRP)<2.6, whereas achievement of the other components of CDC was comparable between groups.

### Table 1  Baseline demographics and disease characteristics

<table>
<thead>
<tr>
<th>Characteristic*</th>
<th>DE019 (MTX-IR)</th>
<th>PREMIER (MTX-naive)</th>
<th>OPTIMA (MTX-IR†)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>ADA+MTX (n=207)</td>
<td>PBO+MTX (n=200)</td>
<td>ADA+MTX (n=268)</td>
</tr>
<tr>
<td>Age, year</td>
<td>56.1 (13.5)</td>
<td>56.1 (12.0)</td>
<td>51.9 (14.0)</td>
</tr>
<tr>
<td>Female, n (%)</td>
<td>158 (76)</td>
<td>146 (73)</td>
<td>193 (72)</td>
</tr>
<tr>
<td>Rheumatoid factor positive, n (%)</td>
<td>168 (81)</td>
<td>179 (90)</td>
<td>228 (88)</td>
</tr>
<tr>
<td>Duration of RA, year</td>
<td>11.0 (9.2)</td>
<td>10.9 (8.8)</td>
<td>0.8 (0.8)</td>
</tr>
<tr>
<td>Prior DMARD use, n (%)</td>
<td>207 (100)</td>
<td>200 (100)</td>
<td>87 (33)</td>
</tr>
<tr>
<td>Prior MTX dose, mg/week</td>
<td>17.4 (9.5)</td>
<td>16.8 (4.2)</td>
<td>0</td>
</tr>
<tr>
<td>Corticosteroid use, n (%)</td>
<td>90 (43)</td>
<td>99 (50)</td>
<td>96 (36)</td>
</tr>
<tr>
<td>Corticosteroid dose, mg/day</td>
<td>3.0 (5.8)</td>
<td>2.6 (6.0)</td>
<td>6.9 (2.5)</td>
</tr>
<tr>
<td>DAS28(CRP), range, 0–10</td>
<td>5.6 (0.9)</td>
<td>5.8 (0.9)</td>
<td>6.3 (0.9)</td>
</tr>
<tr>
<td>DAS28(CRP)&gt;5.1, n (%)</td>
<td>150 (73)</td>
<td>155 (78)</td>
<td>230 (89)</td>
</tr>
<tr>
<td>CRP, mg/dL</td>
<td>1.8 (2.3)</td>
<td>1.8 (2.1)</td>
<td>3.9 (4.2)</td>
</tr>
<tr>
<td>SJC (0–66 joints)</td>
<td>19.3 (9.8)</td>
<td>19.0 (9.5)</td>
<td>21.1 (11.2)</td>
</tr>
<tr>
<td>TJC (0–68 joints)</td>
<td>27.3 (12.7)</td>
<td>28.1 (13.8)</td>
<td>30.7 (14.2)</td>
</tr>
<tr>
<td>HAQ-DI, range, 0–3</td>
<td>1.5 (0.6)</td>
<td>1.5 (0.6)</td>
<td>1.5 (0.6)</td>
</tr>
<tr>
<td>mTSS, range, 0–398</td>
<td>72.1 (60.7)</td>
<td>66.4 (47.4)</td>
<td>18.1 (20.1)</td>
</tr>
<tr>
<td>JE, range, 0–230</td>
<td>41.4 (33.4)</td>
<td>37.2 (25.8)</td>
<td>11.0 (12.3)</td>
</tr>
<tr>
<td>JSN, range, 0–168</td>
<td>30.7 (29.2)</td>
<td>29.2 (24.5)</td>
<td>7.1 (9.6)</td>
</tr>
</tbody>
</table>

*All values are mean (SD), unless otherwise indicated.
†OPTIMA enrolled MTX-naive patients; the Rescue ADA arm included patients treated with placebo+MTX who did not achieve stable low disease activity (ie, MTX-IR patients) at week 22 or 26, and received open-label ADA+MTX for an additional 52 weeks.

ADA, adalimumab; CRP, C-reactive protein; DAS28(CRP), 28-joint Disease Activity Score based on CRP; DMARD, (non-biological) disease-modifying antirheumatic drug; HAQ-DI, Health Assessment Questionnaire Disability Index; IR, inadequate response; JE, joint erosion; JSN, joint space narrowing; mTSS, modified total Sharp score; MTX, methotrexate; PBO, placebo; RA, rheumatoid arthritis; SJC, swollen joint count; TJC, tender joint count.
Figure 1  Percentage of patients achieving comprehensive disease control based on DAS28(CRP)≤3.2, DAS28(CRP)<2.6 or SDAI≤3.3 at week 52 in (A) DE019, (B) PREMIER and (C) the Rescue ADA arm of OPTIMA. ADA, adalimumab; DAS28(CRP), 28-joint Disease Activity Score based on C-reactive protein; HAQ-DI, Health Assessment Questionnaire Disability Index; IR, inadequate response; ΔmTSS, change in modified total Sharp score; MTX, methotrexate; PBO, placebo; SDAI, Simplified Disease Activity Index. *p<0.05 from Fisher’s exact test, ADA+MTX versus PBO+MTX. ***p<0.001 from generalised estimating equation or logistic regression, ADA+MTX versus PBO+MTX.
In the open-label phase of OPTIMA, approximately 50% of patients who met criteria for remission at week 26 achieved CDC at 1 year (vs 24% in the overall population; figure 1C), with radiographic non-progression noted in approximately 90% of patients (figure 4C).

When remission was defined by SDAI $\leq 3.3$, findings were generally comparable, although the proportion of patients achieving CDC was somewhat higher in each of the studies (figure 4). Additionally, to increase the sample size of patients with established RA (DE019), another analysis was conducted using LDA as the clinical criterion at week 24 (ie, $n=46$ with DAS28(CRP) $< 2.6$ vs $n=82$ with DAS28(CRP) $\leq 3.2$) and found that the percentage of patients achieving CDC at week 52 was reduced for both treatment groups (online supplementary figure 5).

In each of the studies, a closer examination of radiographic data among patients who achieved DAS28(CRP) $< 2.6$ at week 24/26 found that the mean $\Delta mTSS$ from week 24/26 to week 52 was small (online supplementary figure 6). Notably, a subgroup of patients in all treatment
groups showed rapid radiographic progression (ie, $\Delta$mTSS≥1.5) between week 24/26 and week 52, with small differences between the treatment groups in early RA and larger differences in favour of combination therapy in established RA. Specifically, 14% of patients receiving adalimumab+MTX (n=44) had progression at week 52 vs 22% of patients receiving placebo+MTX (n=9) in DE019; for PREMIER, percentages were 14% (n=98) vs 17% (n=47), respectively. In OPTIMA, 5% of patients receiving adalimumab+MTX (n=120) had progression at week 52.

**DISCUSSION**
These post hoc analyses of the DE019, PREMIER and OPTIMA studies found that significantly higher rates of CDC were observed following 1 year of treatment with...
the combination of adalimumab+MTX than with placebo+MTX in MTX-IR patients with established RA and in patients with early RA who were MTX-naive. Overall, a numerically greater percentage of patients with early RA in PREMIER and OPTIMA met the criteria for clinical remission than patients with established RA in the DE019 study. Notably, a numerically greater percentage of patients who were in clinical remission at 24 or 26 weeks achieved CDC at 1 year versus patients in the overall population.

Given the clinical, functional and radiographic non-progression efficacy demonstrated by anti-TNFs plus MTX compared with MTX alone in patients with active RA,18 20 36 37 the finding that patients treated with adalimumab+MTX had better outcomes than those receiving MTX alone in both early and established RA is not surprising; however, the combination of these outcomes (ie, CDC) had not been previously assessed.

As might be expected in more advanced disease, CDC was achieved in a relatively low number of patients with established RA compared with patients with early RA. Achievement of CDC in patients with established RA was mostly limited by achievement of DAS28(CRP)<2.6 or SDAI≤3.3 as the clinical component criterion (figure 2 and 3), whereas HAQ-DI was the limiting factor when CDC was based on DAS28(CRP)<2.6 (online supplementary figure 4). Based on the responder analysis, clinical remission was also identified as the chief limiting factor for achieving CDC (using DAS28(CRP)<2.6 as the clinical component criterion) in patients with early RA in the PREMIER study, whereas achievement of normal physical function was limiting for patients with early RA in the OPTIMA study. In contrast, achieving CDC using SDAI≤3.3 as the clinical component criterion was the limiting factor in both PREMIER and OPTIMA, showing that the SDAI has a more stringent definition of remission than the DAS28(CRP). This is consistent with previous reports38 39 and aligns with proposals to change the cut-off values defining remission and LDA for DAS28(CRP) because the currently used values are too lenient.40 However, despite differences in disease activity level classifications, associations of SDAI and DAS28(CRP) remission criteria with functional status and joint damage progression show high accordance.38 39

Table 2 Patients in clinical remission at week 24/26 based on DAS28(CRP)<2.6 or SDAI≤3.3

<table>
<thead>
<tr>
<th></th>
<th>ADA+MTX</th>
<th>PBO+MTX</th>
<th>SDAI≤3.3</th>
<th>PBO+MTX</th>
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<tbody>
<tr>
<td><strong>DE019 (MTX-IR)</strong></td>
<td></td>
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</tr>
<tr>
<td>Proportion in remission at week 24</td>
<td>n=202</td>
<td>n=191</td>
<td>n=202</td>
<td>n=191</td>
</tr>
<tr>
<td>Characteristics of patients in remission at week 24, mean (SD)</td>
<td></td>
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</tr>
<tr>
<td>DAS28(CRP)</td>
<td>2.11 (0.33)</td>
<td>2.06 (0.40)</td>
<td>1.88 (0.32)</td>
<td>1.65 (0.08)</td>
</tr>
<tr>
<td>mTSS</td>
<td>57.4 (58.1)</td>
<td>41.4 (26.8)</td>
<td>45.2 (46.7)</td>
<td>41.6 (33.2)</td>
</tr>
<tr>
<td>HAQ-DI</td>
<td>0.40 (0.45)</td>
<td>0.51 (0.68)</td>
<td>0.33 (0.41)</td>
<td>0.50 (0.84)</td>
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<tr>
<td><strong>PREMIER (MTX-naive)</strong></td>
<td></td>
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<td></td>
</tr>
<tr>
<td>Proportion in remission at week 26</td>
<td>n=267</td>
<td>n=252</td>
<td>n=267</td>
<td>n=252</td>
</tr>
<tr>
<td>Characteristics of patients in remission at week 26, mean (SD)</td>
<td></td>
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</tr>
<tr>
<td>DAS28(CRP)</td>
<td>1.95 (0.35)</td>
<td>2.00 (0.35)</td>
<td>1.78 (0.29)</td>
<td>1.82 (0.27)</td>
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<tr>
<td>mTSS</td>
<td>17.2 (19.4)</td>
<td>23.1 (24.9)</td>
<td>15.5 (17.0)</td>
<td>21.9 (28.6)</td>
</tr>
<tr>
<td>HAQ-DI</td>
<td>0.25 (0.40)</td>
<td>0.22 (0.35)</td>
<td>0.18 (0.28)</td>
<td>0.18 (0.36)</td>
</tr>
<tr>
<td><strong>OPTIMA (MTX-IR†)</strong></td>
<td></td>
<td>NA</td>
<td></td>
<td>NA</td>
</tr>
<tr>
<td>Proportion in remission at week 26</td>
<td>n=348</td>
<td>NA</td>
<td>n=348</td>
<td>NA</td>
</tr>
<tr>
<td>Characteristics of patients in remission at week 26, mean (SD)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DAS28(CRP)</td>
<td>2.0 (0.37)</td>
<td>NA</td>
<td>1.71 (0.27)</td>
<td>NA</td>
</tr>
<tr>
<td>mTSS</td>
<td>13.4 (19.9)</td>
<td>NA</td>
<td>14.9 (23.3)</td>
<td>NA</td>
</tr>
<tr>
<td>HAQ-DI</td>
<td>0.47 (0.50)</td>
<td>NA</td>
<td>0.31 (0.40)</td>
<td>NA</td>
</tr>
</tbody>
</table>

*p<0.0001 and **p=0.001 vs ADA+MTX, Fisher’s exact test; the last observation carried forward method was used to impute missing data for patients with postbaseline DAS28(CRP).
†OPTIMA enrolled MTX-naive patients; the Rescue ADA arm included patients treated with placebo+MTX who did not achieve stable low disease activity (ie, MTX-IR patients) at week 22 or 26, and received open-label ADA+MTX for an additional 52 weeks.

The combination of adalimumab+MTX than with placebo+MTX in MTX-IR patients with established RA and in patients with early RA who were MTX-naive. Overall, a numerically greater percentage of patients with early RA in PREMIER and OPTIMA met the criteria for clinical remission than patients with established RA in the DE019 study. Notably, a numerically greater percentage of patients who were in clinical remission at 24 or 26 weeks achieved CDC at 1 year versus patients in the overall population.

Given the clinical, functional and radiographic non-progression efficacy demonstrated by anti-TNFs plus MTX compared with MTX alone in patients with active RA,18 20 36 37 the finding that patients treated with adalimumab+MTX had better outcomes than those receiving MTX alone in both early and established RA is not surprising; however, the combination of these outcomes (ie, CDC) had not been previously assessed.

As might be expected in more advanced disease, CDC was achieved in a relatively low number of patients with established RA compared with patients with early RA. Achievement of CDC in patients with established RA was mostly limited by achievement of DAS28(CRP)<2.6 or SDAI≤3.3 as the clinical component criterion (figure 2 and 3), whereas HAQ-DI was the limiting factor when CDC was based on DAS28(CRP)<2.6 (online supplementary figure 4). Based on the responder analysis, clinical remission was also identified as the chief limiting factor for achieving CDC (using DAS28(CRP)<2.6 as the clinical component criterion) in patients with early RA in the PREMIER study, whereas achievement of normal physical function was limiting for patients with early RA in the OPTIMA study. In contrast, achieving CDC using SDAI≤3.3 as the clinical component criterion was the limiting factor in both PREMIER and OPTIMA, showing that the SDAI has a more stringent definition of remission than the DAS28(CRP). This is consistent with previous reports38 39 and aligns with proposals to change the cut-off values defining remission and LDA for DAS28(CRP) because the currently used values are too lenient.40 However, despite differences in disease activity level classifications, associations of SDAI and DAS28(CRP) remission criteria with functional status and joint damage progression show high accordance.38 39
Radiographic non-progression was the least limiting factor for achievement of CDC in our analyses, similar to other studies in patients treated with a combination of MTX and a TNF inhibitor. This may be due to blockage of local TNF-α production and subsequent inflammatory response with TNF inhibitors compared with MTX alone. Notably, rapid radiographic progressors were seen in our study; however, absolute numbers were low, limiting conclusions.

A targeted treatment approach in OPTIMA (aimed at stable LDA at 6 months) enabled 24% of MTX-IR patients to achieve CDC following an open-label adalimumab+MTX treatment for an additional year. In PREMIER, 25% of MTX-naive patients with early RA who were receiving initial treatment with the combination of adalimumab+MTX achieved CDC at year 1. Although the numbers of patients who achieved CDC in PREMIER and DE019 may be lower than might have been achieved if these studies had allowed treatment adjustments for incomplete responders, the numbers indirectly support treat-to-target approaches to optimise long-term outcomes for patients with established as well as early RA disease.
The notable finding that achievement of remission at approximately 6 months was associated with CDC at 1 year is in line with the strong association between early treatment response and subsequent long-term remission that has been well documented in studies of patients with RA. Although no single response measure can reflect the full burden of RA, CDC provides a stringent and comprehensive assessment of patient outcomes following treatment; furthermore, remission as a sole outcome does reflect CDC. The components of CDC impact each other; for example, it has been demonstrated that joint damage is a consequence of disease activity and irreversible impairment of physical function a consequence of joint damage. Disability is also a consequence of both the current (reversible component) and past (irreversible component) disease activity, again highlighting the importance of achieving remission. Achievement of CDC has been associated with clinically meaningful improvements for patients in work-related outcomes, health-related quality of life, pain and fatigue; stringent remission is similarly associated with significant improvements in health-related quality of life and physical function. Thus, the potential for remission to identify patients likely to achieve CDC has important implications for disease management. Although there is evidence suggesting that an earlier time to remission is the strongest predictor of sustained response/remission in patients with RA, prospective clinical trials are needed to determine whether it will also be the strongest predictor of CDC.

The key limitation of the current report is that these were post hoc analyses of three individual clinical trials; however, although population differences prevented pooling of data, results were consistent among the trials. Nonetheless, the findings suggest that the current approach to comprehensive treatment deserves further study. Another limitation is that, although the components of CDC are established outcomes in clinical trials, the definition of CDC we used is not validated but is derived from treat-to-target recommendations. Debate remains regarding appropriate cut-off values for DAS28(CRP), defined in this study as ≤2.6 and ≤3.2 for remission and LDA, respectively; SDAI was also examined because it studies.51 In addition, the low number of patients was an issue in certain subgroup analyses among patients who achieved remission at week 24 or 26.

In conclusion, CDC appears to be a realistic treatment goal for many patients with early RA and established RA, and treatment with adalimumab+MTX appears to increase the likelihood of CDC. Achievement of remission at week 24 or 26 is associated with subsequent achievement of CDC at 1 year and thus may help guide treatment decisions.
Rheumatoid arthritis

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Achieving comprehensive disease control in patients with early and established rheumatoid arthritis treated with adalimumab plus methotrexate versus methotrexate alone

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