CONCISE REPORT

Efficacy of glucocorticoids, conventional and targeted synthetic disease-modifying antirheumatic drugs: a systematic literature review informing the 2016 update of The EULAR recommendations for the management of rheumatoid arthritis

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ABSTRACT

Objectives To perform a systematic literature review (SLR) informing the 2016 update of the recommendations for the management of rheumatoid arthritis (RA).

Methods An SLR for the period between 2013 and 2016 was undertaken to assess the efficacy of glucocorticoids (GCs), conventional synthetic disease-modifying antirheumatic drugs (csDMARDs) and targeted synthetic DMARDs (tsDMARDs) (tofacitinib and baricitinib) in randomised clinical trials.

Results For GCs, four studies were included in the SLR. Patients without poor prognostic factors experienced benefit when GCs were added to methotrexate (MTX). Lower doses of GCs were similar to higher doses. For csDMARDs, two new studies comparing MTX monotherapy with combination csDMARD were included in the SLR. In the REACH trial at the end of 12 months no difference between the groups in disease activity, functional ability and radiographic progression was seen, using principles of tight control (treat-to-target). In the CARERA trial, combination therapy with csDMARDs was not superior to MTX monotherapy and monotherapy was better tolerated. For tsDMARDs, tofacitinib and baricitinib were shown to be more effective than placebo (MTX) in different patient populations.

Conclusions Addition of GCs to csDMARD therapy may be beneficial but the benefits should be balanced against the risk of toxicity. Under tight control conditions MTX monotherapy is not less effective than combination csDMARDs, but better tolerated. Tofacitinib and baricitinib are efficacious in patients with RA, including those with refractory disease.

INTRODUCTION

The landscape of rheumatoid arthritis (RA) treatment has unquestionably changed dramatically during the last decade. The development and introduction to daily clinical practice of disease modifying antirheumatic drugs (DMARDs) as well as earlier diagnosis and treatment, and well defined goals of treatment, have contributed to this treatment revolution. Despite this progress, there are still unmet needs, and a better application of the currently available treatments as well as better treatment strategies are needed. Practical recommendations based on the existing evidence are appropriate tools for the rheumatologists. In 2013 a European League Against Rheumatism (EULAR) task force has revised the previous recommendations on RA treatment. A revision of the 2013 recommendations was now undertaken.

The aim of this review was to inform the new EULAR recommendations on the management of RA on efficacy of glucocorticoids (GCs), conventional synthetic DMARDs (csDMARDs) and two targeted synthetic DMARDs (tsDMARDs), tofacitinib and baricitinib based on new evidence accrued since 2013. The results of this and two other systematic literature reviews (SLRs) provided the task force with the current state of evidence.

METHODS

An SLR using MEDLINE, EMBASE and the Cochrane CENTRAL library was performed from January 2013 until February 2016, based on a pre-specified PICOS statement: P=population, I=interventions, C=comparators, O=outcomes and S=study design. The population was ‘adult RA patients’; the intervention was (1) GCs, (2) csDMARDs (methotrexate (MTX), leflunomide, sulfasalazine, hydroxychloroquine, intramuscular gold, auranofin, azathioprine, ciclosporine, minocycline, D-penicillamine, cyclophosphamide, chlorambucil, mycophenolate, tacrolimus), (as monotherapy or combination therapy) and (3) tsDMARDs (tofacitinib and baricitinib); the comparator was patients not receiving the abovementioned treatments; the outcome pertained to efficacy on disease activity, function, patient reported outcomes (PROs) and structural damage; and the study design always was ‘randomised controlled trials’ (RCTs). Risk of bias (RoB) was assessed using the Cochrane RoB assessment tool (Cochrane Handbook for Systematic Reviews of Interventions V5.1.0 March 2011 (cited September 2016); available from: http:// Handbook.cochrane.org/). ORs for dichotomous measures were...
determined to assess the magnitude of treatment effect. The DerSimonian and Laird random-effects model was used to pool the data when possible, allowing for both within-study and between-study variations. Statistical heterogeneity among studies was evaluated using the I² statistic and χ² test where a p value <0.10 was considered to be statistically significant. A value of above 50% for I² was considered to be high. Details about the search and the studies included can be found in the online supplementary material. The selected group of patients included in RCTs as well as the relatively short duration of RCTs, makes addressing long-term safety of drugs in RCTs difficult. For this reason, safety aspects of GCs and csDMARDs were addressed in a separate SLR based on observational studies coming from registries. Some safety issues regarding tsDMARDs will be discussed here, since real life data of tsDMARDs are still lacking.

RESULTS

Efficacy of addition of GCs to csDMARDs

Of 348 hits, 4 studies were included in the analysis (table 1). The selection of articles is shown in online supplementary figure S1. A small study by Menon et al showed greater efficacy of a combination of csDMARDs with intra-articular GCs than with csDMARDs alone in patients with RA with less than 2 years disease duration, but this was an open label study with high RoB. In the CareRA trial patients with early RA, but without poor prognostic factors, benefited from the addition of GCs (COBRA-slim) to MTX with no differences in safety observed. The primary end point of this study was not met, since the percentage of patients achieving remission at week 16 was only numerically but not significantly higher in the GC group (65.1% vs 46.8%, p=0.08). However, this substudy analysis did not have sufficient statistical power and had a high RoB, primarily due to lack of blinding.

A non-inferiority trial compared two different GC strategies; the COBRA-light strategy (prednisolone at 30 mg/day, tapered to 7.5 mg/day in 9 weeks) in combination with MTX; and the COBRA strategy, using prednisolone at 60 mg/day (tapered to 7.5 mg/day in 6 weeks) in combination with both MTX and sulfasalazine. The lower dose of GCs was efficacious in suppressing clinical disease activity and improving functional ability, but non-inferiority could not be claimed formally. The degree of radiographic progression was similar in the two groups (COBRA and COBRA-light). However, this study also had a high RoB (open design), and no comparison with application of conventional GCs was performed.

In a double-blind RCT with patients with established RA, low-dose prednisolone with modified release (‘chronotherapy’) added to existing DMARD treatment in patients with active disease had a significant effect on disease activity and health-related quality of life compared with placebo.

A pooled analysis could not be performed because of significant heterogeneity of the studies regarding designs, patient populations, doses and routes of administration of GCs, and outcome measures. The results of the newer RCTs are in accordance with the previously formulated standpoint that GC when added to csDMARD therapy may have beneficial effects. Safety aspects, as addressed in a separate SLR, have to be taken into consideration. Level of evidence (LOE): 1a.

Efficacy of csDMARDs and csDMARD combinations

In total 518 studies were screened. The selection of articles is shown in online supplementary figure S2. Only two new studies comparing MTX monotherapy with MTX in combination with another csDMARD without differences in GC usage were determined to assess the magnitude of treatment effect. The DerSimonian and Laird random-effects model was used to pool the data when possible, allowing for both within-study and between-study variations. Statistical heterogeneity among studies was evaluated using the I² statistic and χ² test where a p value <0.10 was considered to be statistically significant. A value of above 50% for I² was considered to be high. Details about the search and the studies included can be found in the online supplementary material. The selected group of patients included in RCTs as well as the relatively short duration of RCTs, makes addressing long-term safety of drugs in RCTs difficult. For this reason, safety aspects of GCs and csDMARDs were addressed in a separate SLR based on observational studies coming from registries. Some safety issues regarding tsDMARDs will be discussed here, since real life data of tsDMARDs are still lacking.

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monotherapy and in combination with MTX has been sug-
RA population after failure of both antitumour necrosis factor
were dosed orally using a weekly step-down scheme (30 –
GCs) (see online supplementary table S1). In these arms GCs
not bene
different in patients with early RA.15 One study compared a
of 10 mg/week and 20 mg/week MTX was not statistically dif-
mal dose.
formulated standpoint that combination of csDMARDs
1-year results of the tREACH, nor those of the CARERA study ,
—
we could not substantiate clear evidence in
monotherapy, again
in combination with MTX, was formally proven in different
1-year results of the tREACH, nor those of the CARERA study ,
—
primary end point, make the interpretation of the results
of many clinicians
monotherapy , again
expected to a daily dose of
unomide showed similar bene
le for the
higher doses (starting at 60 mg/day) as in the COBRA regimen,
although formal non-inferiority was not proven. In fact,
common” when a tight treat-to-target approach is
primary outcomes (MTX-naive, csDMARD and biological
DMARD (bDMARD) inadequate responders) compared with
whether the tsDMARDs (tofacitinib and baricitinib) are effective and safe in the short term. (LOE: 1A)
DISCUSSION
Overall, the results of this review confirmed the previous SLR and expanded the overall insights. Although the evidence on
efficacy of short-term GCs when added to csDMARDs is robust
and undisputed, there are still concerns regarding long-term
safety (such as infections, diabetes, osteoporosis, and gastro-
intestinal and cardiovascular events). Preliminary long-term
results of the CAMERA II trial showed a low occurrence of AEs
but suggested for the first time an increased cardiovascular risk
for the patients with early RA treated with 10 mg/day prednis-
one for at least 2 years.10 These results are still unpublished
(abstract in American College of Rheumatology 2015). A separate
SLR focusing on the safety of GCs has been performed in order to
inform the task force and enable the formation of the recommenda-
tions.5 GC safety aspects have also been addressed
in a separate paper prior EULAR activity.36 Clear consensus
regarding the dose and tapering of GCs is still lacking. New
data have suggested that short-term lower doses of GCs (starting
at 30 mg prednisone per day with rapid tapering), as in the
COBRA-light regimen, might be a feasible alternative to the
higher doses (starting at 60 mg/day) as in the COBRA regimen,
although formal non-inferiority was not proven. In fact, this
trial did not fulfil the inclusion criteria for the SLR, since there
was no comparator group (group without GCs according to the
PICO). However, we decided to include it in the SLR since the
question posed is highly clinically relevant.

Interestingly, the tREACH trial has suggested that the efficacy
of oral GCs as bridging treatments was not superior to intra-
muscular GCs. Two new studies were published regarding
chronotherapy and intra-articular GC therapy, thus answering
one of the research questions posed in 2013. The latter however
was a high RoB study.

Regarding the choice of csDMARD combination therapy over
monotherapy, again—and in contradiction with the perception
of many clinicians—we could not substantiate clear evidence in
favour of combination therapy with csDMARDs. Neither the
1-year results of the tREACH, nor those of the CARERA study,
showed clear evidence that MTX monotherapy is inferior to
combination therapy with csDMARDs when used in combina-
tion with GCs and when a tight treat-to-target approach is
employed. Importantly, monotherapy was generally better toler-
ated than combination therapy in these studies. Generally,
the complexity of the design of pragmatic trials and certain meth-
odoslogic issues, such as high dropout rates and change of
primary end point, make the interpretation of the results
challenging.

Efficacy of tsDMARDs (tofacitinib and baricitinib)
From the 134 hits on tofacitinib 9 were identified as RCTs (table 2).18–26 Efficacy of tofacitinib, both as monotherapy and
in combination with MTX, was formally proven in different
patient populations (MTX-naive, csDMARD and biological
DMARD (bDMARD) inadequate responders) compared with
placebo (background MTX). For baricitinib the literature search
yielded eight new RCTs (two of them had PROs as main study
outcomes) (table 3).27–34 Similar clinical efficacy of baricitinib
in monotherapy and in combination with MTX has been sug-
gested, but only the combination (baricitinib+MTX) signifi-
cantly inhibited radiographic progression.29 In the MTX-IR
(inadequate responder) RA-BEAM study, comparing adalimu-
mab+MTX versus baricitinib+MTX versus placebo+MTX,
showed small but significantly lower responses for adalimumab+
MTX versus baricitinib+MTX, but both were higher than
placebo+MTX (Disease Activity Score 28-C reactive protein
<2.6 19% vs 24% at 4%) at week 12.35

Importantly, baricitinib has now shown efficacy in a refractory
RA population after failure of both antitumour necrosis factor
(anti-TNF) and non-anti-TNF bDMARDs.16 All studies had low
RoB. The selection of articles for tofacitinib and baricitinib is
shown in online supplementary figures S3 and S4, respectively.

No meta-analysis could be performed due to the heterogen-
cy between the studies. The most commonly found laboratory
abnormalities with tofacitinib were mild decreases in neutrophil
and lymphocyte counts and mild increases in aminotransferase
and creatinine levels, while baricitinib was associated with
reductions in haemoglobin levels. The relative risks for serious
AEs with tofacitinib and baricitinib compared with placebo were
0.8 (95% CI 0.5 to 1.3) and 1.0 (95% CI 0.6 to 1.7), respect-
ively. However, a significantly increased risk of herpes infection
was seen (RR=3.1, 95% CI 1.1 to 8.5) with tofacitinib.

The results of the newer RCTs are in accordance with the previ-
ously formulated standpoint that the tsDMARDs (tofacitinib and
baricitinib) are effective and safe in the short term. (LOE: 1A)
<table>
<thead>
<tr>
<th>Study</th>
<th>Study design</th>
<th>N patients</th>
<th>Patient population</th>
<th>Primary outcome</th>
<th>Tofacitinib monotherapy or in combination with csDMARDs</th>
<th>Comparator arm</th>
<th>Result in tofacitinib arm (5 mg×2)</th>
<th>Result in tofacitinib arm (10 mg×2)</th>
<th>Result in comparator arm</th>
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<tr>
<td>Lee et al²³</td>
<td>ORAL start</td>
<td>958</td>
<td>DMARD-naive</td>
<td>ACR70 and mean change from baseline SHS at month 6</td>
<td>Monotherapy MTX</td>
<td>25.5% least-squares mean (±SE)=0.2±0.1</td>
<td>37.7% least-squares mean (±SE)=&lt;0.1±0.1</td>
<td>12% least-squares mean (±SE)=0.8±0.2</td>
<td>p&lt;0.001</td>
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<td>Kremer et al²²</td>
<td>ORAL sync</td>
<td>795</td>
<td>csDMARDs and/or bDMARDs IR</td>
<td>ACR20 at month 6 Change in HAQ at month 3 Remission at month 6</td>
<td>Combination with csDMARDs</td>
<td>Placebo with csDMARDs</td>
<td>52.1% –0.44 8.5%</td>
<td>56.6% –0.53 12.5%</td>
<td>30.8% –0.16 2.6%</td>
<td>&lt;0.001</td>
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<td>Van der Heijde et al²⁵</td>
<td>ORAL scan</td>
<td>800</td>
<td>MTX IR</td>
<td>ACR20 at month 6 Combination with MTX</td>
<td>Placebo with MTX</td>
<td>51.5% 61.8% 25.3%</td>
<td>&lt;0.001 for both tofacitinib doses vs placebo</td>
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<tr>
<td>Burmester et al²¹</td>
<td>ORAL step</td>
<td>399</td>
<td>TNFis IR and/or bDMARDs IR</td>
<td>ACR20 at month 3 ΔHAQ at month 3 Remission at month 3</td>
<td>Combination with MTX</td>
<td>Placebo with MTX</td>
<td>41.7% –0.43 6.7%</td>
<td>48.1% –0.46 8.8%</td>
<td>24.4% –0.18 1.7%</td>
<td>&lt;0.0011  &lt;0.0001†  &lt;0.05†</td>
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<td>Tanaka et al²⁴</td>
<td>ORAL step</td>
<td>318</td>
<td>csDMARDs and/or bDMARDs IR</td>
<td>ACR20 at month 3</td>
<td>Monotherapy Placebo</td>
<td>73.1% 84.9% 15.4%</td>
<td>&lt;0.0001 both groups vs placebo</td>
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<td>Strand et al²⁸</td>
<td>ORAL solo</td>
<td>611</td>
<td>csDMARDs and/or bDMARDs IR</td>
<td>Multiple PROs</td>
<td>Monotherapy Placebo</td>
<td>Tofacitinib &gt; placebo</td>
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<tr>
<td>Strand et al³⁹</td>
<td>ORAL step</td>
<td>399</td>
<td>≥1 TNFi*</td>
<td>Multiple PROs Combination with MTX</td>
<td>Placebo with MTX</td>
<td>Tofacitinib &gt; placebo</td>
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<tr>
<td>Wallenstein et al³⁵</td>
<td>2 Phase-IIb studies, double-blind</td>
<td>507</td>
<td>combination with MTX 384 (monotherapy)</td>
<td>MTX IR Multiple PROs Both in combination with MTX and in monotherapy</td>
<td>Placebo</td>
<td>Tofacitinib &gt; placebo (both as monotherapy and in combination with MTX)</td>
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<tr>
<td>Strand et al³⁶</td>
<td>ORAL standard</td>
<td>717</td>
<td>MTX IR</td>
<td>Multiple PROs Combination with csDMARDs</td>
<td>Placebo</td>
<td>Tofacitinib+MTX &gt; control group</td>
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*Since tofacitinib at 5 mg twice daily failed to be statistically significant for radiographic progression, and due to the step-down procedure applied to primary efficacy end points, significance was not declared for the HAQ DI score or remission (DAS28-ESR <2.6) for tofacitinib at 5 mg twice daily. For both groups versus placebo.

ACR, American College of Rheumatology; bDMARD, biological DMARD; csDMARD, conventional synthetic DMARD; DAS, Disease Activity Score; DMARD, disease-modifying antirheumatic drug; HAQ, Health Assessment Questionnaire; IR, inadequate responder; MTX, methotrexate; PROs, patient reported outcomes; RA, rheumatoid arthritis; SHS, van der Heijde modification of the total Sharp Score; TNFi, tumour necrosis factor inhibitor.
There is a clear need for studies addressing the optimal use of csDMARDs. No new studies fulfilling the inclusion criteria regarding dose and route of administration of MTX were identified. A previous SLR by Visser and van der Heijde had addressed this issue.

Tofacitinib is the first JAK inhibitor approved for the treatment of RA in many countries and baricitinib is under regulatory evaluation. This SLR confirmed that tofacitinib has beneficial effects on disease activity, physical function, radiographic progression and PROs, both in patients with early RA who are DMARD-naïve and in patients with established disease who have failed csDMARDs and/or bDMARDs. Baricitinib was found to be effective in MTX-naïve patients and also after failure of drugs with multiple modes of action. Data on long-term safety of this new class of DMARDs from real life observational studies are needed. Until then, rheumatologists are advised to take into account safety data obtained through RCTs and follow the labels of each drug, including AEs and lab monitoring.

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