CLINICAL SCIENCE

Low dose, add-on prednisolone in patients with rheumatoid arthritis aged 65+: the pragmatic randomised, double-blind placebo-controlled GLORIA trial

Maarten Boers 1,2, Linda Hartman 1,2, Daniela Opris-Belinski 3, Reinhard Bos 4, Marc R Kok 5, Jose AP Da Silva 6, Eduard N Griepp 7, Ruth Klaasen 8, Cornelia F Allaart 9, Paul Baudoin 10, Hennie G Raterman 11, Zoltan Szekanecz 12, Frank Buttgereit 13, Pavol Masaryk 14, L Thomas Klausch 1, Sabrina Paolino 15, Annemarie M Schilder 1, Willem F Lems 2, Maurizio Cutolo 15, For the GLORIA Trial consortium

Handling editor Josef S Smolen

► Additional supplemental material is published online only. To view, please visit the journal online (http://dx.doi.org/10.1136/annrheumdis-2021-221957).

For numbered affiliations see journal online (http://onlinelibrary.wiley.com/doi/10.1136/annrheumdis-2021-221957). To view, please visit the journal online (http://dx.doi.org/10.1136/annrheumdis-2021-221957).

ABSTRACT

Background Low-dose glucocorticoid (GC) therapy is widely used in rheumatoid arthritis (RA) but the balance of benefit and harm is still unclear.

Methods The GLORIA (Glucocorticoid LOw-dose in Rheumatoid Arthritis) pragmatic double-blind randomised trial compared 2 years of prednisolone, 5 mg/day, to placebo in patients aged 65+ with active RA. We allowed all cotreatments except long-term open label GC and minimised exclusion criteria, tailored to seniors. Benefit outcomes included disease activity (disease activity score; DAS28, coprimary) and joint damage (Sharp/van der Heijde, secondary). The other coprimary outcome was harm, expressed as the proportion of patients with ≥1 adverse event (AE) of special interest. Such events comprised serious events, GC-specific events and those causing study discontinuation. Longitudinal models analysed the data, with one-sided testing and 95% confidence limits (95% CL).

Results We randomised 451 patients with established RA and mean ± 2.1 comorbidities, age 72, disease duration 11 years and DAS28 4.5. 79% were on disease-modifying treatment, including 14% on biologics. 63% prednisolone versus 61% placebo patients completed the trial. Discontinuations were for AE (both, 14%), active disease (3 vs 4%) and for other (including covid pandemic-related disease) reasons (19 vs 21%); mean time in study was 19 months. Disease activity was 0.37 points lower on prednisolone (95% CL -0.23, p<0.0001); joint damage progression was 1.7 points lower (95% CL 0.04, p=0.02), with the largest contrast in (mostly non-severe) infections. Other GC-specific events were rare.

Conclusion Add-on low-dose prednisolone has beneficial long-term effects in senior patients with established RA, with a trade-off of 24% increase in patients with mostly non-severe AE; this suggests a favourable balance of benefit and harm.

Trial registration number NCT02585258.

INTRODUCTION

Rheumatoid arthritis (RA) is a systemic inflammatory disease characterised by pain, progressive disability and premature death. Both RA and its treatment cause comorbidity. Current treatment strategies have considerably improved the prognosis...
Rheumatoid arthritis

Summary box

How might this impact on clinical practice or future developments?
⇒ Results are immediately applicable to clinical practice and suggest add-on low-dose prednisolone has substantial long-term effects in senior patients with RA patients on optimum treatment, with a favourable balance of benefit and harm.

but come with safety issues and often high costs. In addition, many patients still have a smouldering progressive disease.1

Glucocorticoids (GC) were introduced in the 1950s, and chronic low-dose treatment is common in RA, but the balance between benefit and harm is still unclear, especially for chronic low-dose therapy. Meta-analyses show that GC therapy reduces disease activity and slows joint damage progression,2 3 so the debate mostly focuses on harm.4 Most experts agree that long-term GC therapy is harmful, and existing guidelines suggest to avoid or use GC only as ‘bridging’ therapy; however, such opinions are based on observational studies with high potential for bias.5 The limited data from trials (mostly in early RA) do not support strong claims of harm,4 but their generalisability is questioned. Pragmatic trials to overcome this6 have not been attempted. This lack of information results in a wide range of usage patterns,5 but overall, a high prevalence of chronic use.6 7

RA prevalence increases with age, peaking at age 70,8,9 so we can expect more RA in ageing populations. Seniors have the highest risk for treatment–associated harm, given comorbidity and its treatment.10 Regrettably, seniors are under-represented or even excluded from clinical trials that provide the evidence base for treatment of RA.11

In the 2-year pragmatic, placebo-controlled GLORIA (Glucocorticoid LOw-dose in Rheumatoid Arthritis) trial, we assessed the effectiveness and safety of prednisolone 5mg/day added to standard of care in senior patients with RA.

METHODS

GLORIA is an investigator-initiated, randomised, double-blind, placebo-controlled, multicentre pragmatic trial, performed in 28 clinical centres in seven EU countries, approved by country-specific regulatory bodies and medical ethical committees and executed in accordance with Good Clinical Practice and the Declaration of Helsinki. An independent Contract Research Organisation monitored the data. The first author prepared the manuscript; all authors vouch for the accuracy and completeness of the data and for the fidelity of the trial to the protocol.

Our pragmatic design approached routine standard of care and was tailored to senior patients: minimal eligibility criteria, routine assessments and procedures and minimal limitations on concurrent antirheumatic treatment. For full details (including published protocol14 and the statistical analysis plan, see online supplementary appendices).

Participants

Eligible patients aged 65 or above had RA15 16 with more than minimal disease activity, that is, with a 28-joint disease activity score (DAS28) ≥2.60 (after protocol amendment; initially ≥3.20). Exclusion criteria focused on uncontrolled conditions that might be adversely affected by GC therapy, current GC therapy and conditions with an absolute indication or contraindication for GC therapy. All patients provided written informed consent.

Procedures

We randomised patients (1:1) to receive prednisolone 5mg/day or placebo for 2 years. A web-based case record form allocated treatment based on minimisation,18 stratified for prior use of GC, modification of antirheumatic treatment at baseline and centre. Opaque capsules contained one prednisolone or placebo tablet; patients, care providers and assessors were blinded to allocation. Success of blinding was not assessed. Throughout the 2-year trial period, all patients received standard of care antirheumatic treatment, enhanced by the trial procedures and allowing most modifications (for limitations, see below). As part of this, we advised calcium 500mg/ vitamin D3 800 IU supplementation in all patients.

With exception of chronic oral GC, we allowed all cotreatment (and changes for RA, including disease-modifying antirheumatic drugs (DMARDs), non-steroidal anti-inflammatory drugs and short-term GC for flares and comorbidity within protocol-defined limits. Patients exceeding these limits but not placed on chronic GC therapy could remain in the trial. To emulate a short-term placebo-controlled trial, we requested (but did not mandate) stable antirheumatic therapy in the first 3 months; if deemed unavoidable, we requested to change treatment at baseline.

We measured medication adherence through counts of returned capsules, as electronic cap monitoring proved unreliable,19 and defined good adherence as ≥80% capsule intake.20 Outcomes requiring physical examination and routine blood sampling were assessed at baseline, 3, 6, 12, 18 and 24 months; patients reported outcomes at these times and additionally through telephone interviews at 9, 15 and 21 months. Imaging was performed at baseline and at 24 months.

Outcomes

The primary outcome for benefit was DAS28; for harm, the co-primary outcome was the total number of patients experiencing at least one adverse event (AE) of special interest (AESI). AESI included serious AE (SAE) according to the Good Clinical Practice definition, and the following (‘other AESI’):
► any AE (except worsening of disease) leading to discontinuation.
► Myocardial infarction, cerebrovascular or peripheral arterial vascular event.
► Symptomatic bone fracture.

We recorded and coded21 AE at every patient contact until 3 months from discontinuation or start of tapering and adjudicated all potential AESI on the blinded data.

Joint damage progression (radiographs of hands and forefeet) and bone health were secondary outcomes. We used the mean joint damage score22 of two assessors independently assessing at least one adverse event (AE) of special interest (AESI). Exclusion criteria focused on uncontrolled conditions that might be adversely affected by GC therapy, current GC therapy and conditions with an absolute indication or contraindication for GC therapy. All patients provided written informed consent.

Statistical procedures

For harm, we expected a base rate of 20%,24 and 800 patients would yield 80% power to detect an increase to 27.5%,
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relative risk 1.38. Slow recruitment and higher rates prompted a blinded interim analysis that suggested 450 patients would detect similar relative risks and a risk difference of 11%, with sufficient power to detect benefit, as suggested by the CAPRA2 (Circadian Administration of Prednisone in Rheumatoid Arthritis) study.\textsuperscript{25}

The safety population (for harm) comprised patients who took at least one capsule of study medication; the modified intention-to-treat population (for benefit) comprised patients in the safety population with at least one baseline and one follow-up assessment. To quantify early response after 3 months, we determined a ‘per-protocol’ population on the blinded data set: patients on stable antirheumatic treatment with complete data, at least 80% adherence, and no protocol violations in the first 3 months.

Before the analysis, we first addressed incomplete DAS28 data by imputation from adjacent values if only patient global assessment was missing, otherwise with single imputation by chained equations if DAS28-CRP or information from the Rapid-3 questionnaire was available. The mixed model analysis subsequently addressed data missing at random; to reduce complexity, time was treated as fixed factor. As sensitivity analysis, non-responder imputation addressed non-random missingness: patients completing the trial with DAS28 improvement less than 0.6, and patients with GC-related protocol violations or premature discontinuation were classified non-responder. For joint damage, we added a complete-case analysis and one that linearly extrapolated end point values from baseline, given disease duration and zero damage at disease initiation. Continuous remote and onsite checks against source data minimised missingness for harm.

We expected increased benefit and harm, so we applied a limited number of one-sided tests ($p<0.05$) to reject null hypotheses at maximum power. Furthermore, we predefined trial success, trade-off or failure on the basis of the primary

Figure 1  Description of analysis populations and patient disposition. For the early response analysis, extra criteria were applied as listed. ITT, intention to treat; PP, per protocol.
outcomes and damage progression (see statistical analysis plan).

For benefit, we designed mixed effects models adjusted for stratification factors. For disease activity, the main model estimated the mean effect of treatment over 2 years with possible time–treatment interactions as secondary analysis. For joint damage, the model did not converge, so we used linear regression, excluding the (non-significant) effect of site. For harm, we used generalised estimating equations to better estimate relative risks and their variance. We tested the three (correlated) bone health measures after Benjamini-Hochberg adjustment.26

We did not restrict concurrent antirheumatic treatment, so we expected confounding and loss of contrast due to (1) more treatment intensification in the placebo group for active disease or AEs and (2) more tapering in the prednisolone group for inactive disease. In the blinded data set, we looked for the first occurrence of such a lasting change in antirheumatic treatment between months 3 and 15 of the trial, in patients remaining in the trial for at least 3 months thereafter. Two separate Z-tests analysed the differences in proportions between the groups; as uncorrelated occurrences, these were tested at a Bonferroni-adjusted threshold26 of (one-sided) p<0.025.

R-software (V.4.0.2; gee_4.13–20, mice_3.11.0, lme4_1.1–26, lmerTest_3.1–3. 2021) performed the main analyses, and IBM SPSS statistics V.26 and Microsoft Excel (2016) the descriptives. The trial was registered at clinicaltrials.gov.

As noted above and previously reported,27 initial recruitment was slow, many patients proved ineligible due to low disease activity or current GC use. In addition, recruitment and retention of seniors proved challenging, an experience shared with other EU projects focused on this population. Network meetings organised as part of the GLORIA project have resulted in recommendations to improve this situation.28 We adjusted eligibility (see online supplementary appendix), sample size and added recruiting centres, but initiatives from our international patient panel to enhance recruitment and retention were hampered or prohibited by strict and varying ethical guidelines across countries. The COVID-19 pandemic compromised collection of important end point data.

### RESULTS

Between 27 June 2016 and 31 December 2018, we entered 451 patients in The Netherlands (286), Italy (60), Romania (56) and 49 in Portugal, Hungary, Germany and Slovakia (figure 1). Two patients never started study medication; five discontinued the study before the first follow-up assessment; 63% prednisolone and 61% placebo patients completed the 2-year trial. Discontinuations were similar in both groups: for AE (both 14%) and active disease (3 v 4%); the remainder mostly for ‘trial fatigue’ (ie, reasons related to the trial but not to the study medication) and COVID-related access issues (19 v 21%). Mean time on study drug was 19 (SD 8) months (online supplemental figure 1).

The groups were well balanced at baseline (table 1, online supplemental table 1). Patients were mean 72 years, predominately women, with established severe disease; mean DAS28 was 4.5. Most patients received treatment for RA and for multiple, often cardiovascular comorbidities: overall, a median of seven different drugs (table 1). During the trial, good adherence was found in 89% of prednisolone and 88% of placebo patients. At baseline, 61 patients changed DMARD treatment, and 26 during the first 3 months; a total of 60 v 67 patients had one or more changes postrandomisation.

#### Benefit outcomes

The coprimary and secondary end points of benefit were met. In both groups, disease activity declined in the first 3 months, stabilising at 1 year. Over 2 years, prednisolone resulted in mean 0.37
lower DAS28 than placebo (95% (CL) 0.23, p<0.0001; figure 2). Of the three stratification factors, only change of treatment at baseline significantly affected disease activity, adding 0.57 to the decrease in disease activity (95% CL 0.35). Secondary analyses suggested a larger effect of prednisolone initially, especially evident in the per-protocol population at 3 months (figure 2, table 2), and a smaller effect later on (figure 2). The pattern of benefit of prednisolone on disease activity was consistent across core set measures, response indices and achievement of minimal disease and remission (figure 2, tables 2 and 3, online supplemental table 2). In non-responder imputation, the numerical difference between the groups was no longer statistically significant (prednisolone 47%, placebo 40% responders, p=0.08).

At baseline, most patients had evidence of joint damage (table 3). Progression was significantly lower in the prednisolone group, confirmed by one of the two sensitivity analyses (complete case analysis, table 3), and numerically by the distribution of patients with negative or zero progression versus those with any or clinically
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We found significant evidence for confounding due to co-interventions: in the prespecified period, a total of 80 patients changed antirheumatic (DMARD excluding GC) treatment: for active disease, 30 prednisolone versus 48 placebo patients, and for AE 1 patient in each group (test for sum of patients with changes: one-sided p=0.02). This includes eight patients in each group who started or changed treatment with a biologic DMARD. In contrast, 29 prednisolone versus 18 placebo patients tapered treatment as a consequence of inactive disease (one-sided p=0.04, not significant at predefined threshold of 0.025). In addition, the number of patients receiving short-term GC for RA and the total number of administrations were somewhat greater in the placebo group, and placebo patients received such GC on average more than 3 months earlier (online supplemental table 3).

Harm outcomes

The coprimary end point of harm was met. Overall, 60% prednisolone versus 49% placebo patients experienced the harm outcome (adjusted relative risk 1.24, 95% CL 1.04, p=0.02; table 4); none of the stratification factors proved significant. Three respective two patients died. Most SAEs were classified as severe because of hospital admission, most ‘other AESI’ because they required treatment, or because the event was associated with study discontinuation (regardless of severity). The increase in AE was most marked for infections (table 4, online supplemental table 4). One prednisolone patient developed COVID-19 pneumonia.

Non-serious infections were rated as mild (41%) or moderate (56%) without clear differences between the groups. Discontinuations for AE were relatively rare and similar between groups. A minority of patients experienced the majority of AE (figure 3).

At baseline, about one-third of patients had osteoporosis (history or imaging) but only 13% were treated with antiresorptive drugs (table 1). Cotreatment with calcium and vitamin D was instituted in 81% of patients. During the trial symptomatic and asymptomatic fractures occurred at slightly higher rates in the prednisolone group, but the rate of new compression fractures was not significantly different: prednisolone, 19% versus 14% placebo (table 2).

### Table 2

<table>
<thead>
<tr>
<th>Prednisolone (n=156)</th>
<th>Placebo (n=148)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Baseline</strong></td>
<td><strong>Change</strong></td>
</tr>
<tr>
<td>DAS28</td>
<td></td>
</tr>
<tr>
<td>Model†</td>
<td></td>
</tr>
<tr>
<td>Adjusted</td>
<td>4.40 (1.04)</td>
</tr>
<tr>
<td>Unadjusted</td>
<td>4.40 (1.04)</td>
</tr>
<tr>
<td>DAS components</td>
<td></td>
</tr>
<tr>
<td>ESR</td>
<td>28.5 (20.2)</td>
</tr>
<tr>
<td>Tender joint count</td>
<td>4.9 (4.4)</td>
</tr>
<tr>
<td>Swollen joint count</td>
<td>2.9 (3.3)</td>
</tr>
<tr>
<td>Patient global ass.</td>
<td>5.6 (2.4)</td>
</tr>
<tr>
<td>Other core set</td>
<td></td>
</tr>
<tr>
<td>Pain</td>
<td>5.4 (2.5)</td>
</tr>
<tr>
<td>Fatigue</td>
<td>5.1 (2.6)</td>
</tr>
<tr>
<td>Physician global ass.</td>
<td>4.4 (2.0)</td>
</tr>
<tr>
<td>HAQ</td>
<td>1.2 (0.7)</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>9.1 (13.4)</td>
</tr>
<tr>
<td>SDAI</td>
<td>18.7 (8.7)</td>
</tr>
<tr>
<td>CDAI</td>
<td>17.7 (8.6)</td>
</tr>
<tr>
<td><strong>Response §</strong></td>
<td></td>
</tr>
<tr>
<td>EULAR Good</td>
<td>63 (41)</td>
</tr>
<tr>
<td>Moderate</td>
<td>48 (31)</td>
</tr>
<tr>
<td>None</td>
<td>43 (28)</td>
</tr>
<tr>
<td>ACR§</td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>53 (36)</td>
</tr>
<tr>
<td>50</td>
<td>30 (20)</td>
</tr>
<tr>
<td>70</td>
<td>12 (8)</td>
</tr>
<tr>
<td>State</td>
<td></td>
</tr>
<tr>
<td>Minimal disease</td>
<td>57 (37)</td>
</tr>
<tr>
<td>Boolean remission</td>
<td>5 (3.2)</td>
</tr>
</tbody>
</table>

Mean (SD) unless indicated otherwise.

Minimal disease activity: defined as DAS28 <2.60.

Remission: Boolean definition according to ACR-EULAR criteria.46

* Only DAS28 change estimate is based on the primary analysis model. Other results are unadjusted.

† Model does not offer change estimates, these are calculated from the point estimates and provided as reference for the observed/unadjusted DAS28 data.

‡ Difference estimate: mean [1-sided 95% confidence bound]; P value for difference in change:<0.0001.

§ Count (%).

ACR, American College of Rheumatology; CDAI, clinical disease activity index; CRP, C reactive protein; DAS28, Disease Activity Score 28 joints; ESR, erythrocyte sedimentation rate; EULAR, European League Against Rheumatism; HAQ, Health Assessment Questionnaire; NNT, number needed to treat; SDAI, simple disease activity index.
placebo 15%, adjusted relative risk 1.27 (95% CI 0.88; table 5). Over 2 years, spine bone density decreased by about 1% in prednisolone, but increased by 3% in placebo patients, resulting in a significant difference; hip bone density did not change (table 5).

Other GC-specific AESI was rare without relevant differences (table 4) and reports of worsening of pre-existent disease were infrequent (data not shown). Complaints of ecchymosis, haematoma and skin atrophy occurred predominantly in the prednisolone group (28 vs 3 AE). Weight gain was rare, and adrenal insufficiency was not reported. Unblinding occurred only once (GC stress schedule for elective surgery); a stress schedule was given for only two SAEs. One patient in each group underwent joint replacement surgery.

**DISCUSSION**

In patients with RA aged 65+ on standard care that allowed treatment optimisation, add-on low-dose prednisolone had beneficial long-term effects on disease activity and damage progression. The trade-off was an 11% increase in the number of patients with at least one AESI. Among events traditionally associated with GC, the increase comprised mostly mild to moderate infections requiring treatment. Although of concern, these should be interpreted in the light of the high-risk trial population, resembling patients in clinical practice. We suggest that our results constitute a benchmark for the upper limit of harm to be expected with this dose and duration. However, this assumes care by rheumatologists as given in this trial.

This trial is the first large pragmatic trial of GC added to standard of care in RA, the first large treatment trial in senior patients with RA, and one of the first to study and demonstrate long-term effects of GC on disease activity and damage progression in established RA, especially at the low dose of 5 mg/d (or equivalent) continued for 2 years. The mean DAS28 difference of 0.37 may appear modest, but our 3-month results point to more substantial benefit, and results were consistent across core set measures. In secondary analyses the second year contrast appeared smaller as physicians were allowed to continuously optimise treatment, confounding the comparison. Our early results resemble those of earlier studies, especially the CAPRA-2 trial (5 mg/day of modified release prednisolone vs placebo). They align with a recent double-blind trial on tapering of low-dose GC in patients with minimal disease on stable treatment with tocilizumab: tapering caused flares, documenting the value of GC on disease stability even in patients on biologics. Use of biologics was relatively rare in our study, most likely reflecting cautious physicians, and senior patients’ dislike of self-administered parenteral therapy. We do not think availability of expensive treatment played a major role; although still an issue in several of the participating countries, these countries contributed less than 20% of patients.
Despite widespread use, trials of GC are relatively rare, and few have been performed according to current quality standards. Most have studied smaller groups of patients with early RA with higher doses, usually for shorter periods, and mostly focused on benefits; almost all showed benefit, and none noted substantial risks of GC treatment. In addition, these trials and their extensions do show that many early patients with RA initially treated with GC are able to stop such treatment.\(^{31-33}\)

Our findings contrast with risks found in observational studies.\(^{34}\) However, observational studies are hard to interpret and often biased through confounding by indication (channeling bias): that is, preferentially treating more severely diseased patients with GC, then comparing their outcome with that of less severe patients not on GC. The bias increases with duration of follow-up and, if strong, cannot be adequately corrected by techniques such as propensity score matching.\(^{35-33}\) Paradoxically, the widespread perception of risk does not translate into action when GC is administered: many studies have shown that GC-associated comorbidity is often inadequately addressed.\(^{36-37}\) Even in our trial population, many patients with a diagnosis of osteoporosis were inadequately treated (table 1).

### Table 4  More AESI in prednisolone patients; safety population

<table>
<thead>
<tr>
<th></th>
<th>Prednisolone (n=224)</th>
<th>Placebo (n=225)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients with at least one AESI *</td>
<td>134 (60%)</td>
<td>111 (49%)</td>
</tr>
<tr>
<td>SAE only</td>
<td>25</td>
<td>25</td>
</tr>
<tr>
<td>Other AESI only</td>
<td>79</td>
<td>65</td>
</tr>
<tr>
<td>SAE and other AESI</td>
<td>30</td>
<td>21</td>
</tr>
<tr>
<td>By organ class (per 100 person-years) †</td>
<td>SAE Other AESI SAE Other AESI</td>
<td></td>
</tr>
<tr>
<td>Cardiac disorders‡</td>
<td>1.7</td>
<td>0.0</td>
</tr>
<tr>
<td>Eye disorders</td>
<td>0.0</td>
<td>3.1</td>
</tr>
<tr>
<td>Gastrointestinal disorders</td>
<td>1.7</td>
<td>1.7</td>
</tr>
<tr>
<td>Infections and infestations§</td>
<td>7.3</td>
<td>35.0</td>
</tr>
<tr>
<td>Injury, poisoning and procedural complications</td>
<td>0.8</td>
<td>4.0</td>
</tr>
<tr>
<td>Neoplasms benign, malignant and unspecified (incl cysts and polyps)§</td>
<td>2.5</td>
<td>1.1</td>
</tr>
<tr>
<td>Nervous system disorders</td>
<td>2.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Respiratory, thoracic and mediastinal disorders</td>
<td>2.0</td>
<td>2.5</td>
</tr>
<tr>
<td>Other</td>
<td>4.5</td>
<td>2.3</td>
</tr>
<tr>
<td>Total</td>
<td>23</td>
<td>55</td>
</tr>
<tr>
<td>By protocol-defined category</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Infection</td>
<td>26</td>
<td>124</td>
</tr>
<tr>
<td>Urinary tract</td>
<td>4</td>
<td>49</td>
</tr>
<tr>
<td>Pneumonia</td>
<td>2</td>
<td>17</td>
</tr>
<tr>
<td>Other</td>
<td>20</td>
<td>58</td>
</tr>
<tr>
<td>Cardiovascular†</td>
<td>8</td>
<td>2</td>
</tr>
<tr>
<td>Symptomatic fracture**</td>
<td>2</td>
<td>11</td>
</tr>
<tr>
<td>New onset</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>0</td>
<td>2††</td>
</tr>
<tr>
<td>Cataract</td>
<td>0</td>
<td>7</td>
</tr>
<tr>
<td>Glaucoma</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Others‡‡§§</td>
<td>43</td>
<td>43</td>
</tr>
<tr>
<td>Total</td>
<td>80</td>
<td>194</td>
</tr>
</tbody>
</table>

* Adjusted relative risk: 1.24, one-sided 95% bound 1.04, one-sided p: 0.02; number needed to harm: 9.5.
† Two deaths in placebo group on treatment (atrioventricular block, cardiac insufficiency). The protocol-defined category ‘Cardiovascular’ comprised myocardial infarction, cerebrovascular event, peripheral arterial vascular event.
‡ One death in prednisolone group on treatment (septic shock); another case excluded that occurred outside the assessment window of 3 months. This patient with septicemia was discharged alive and later reported (and initially included) as death with unknown date. Date of death was retrieved after database closure and found to be 5 months after discontinuation. The death of this patient was not counted in the primary analysis.
§ One death in prednisolone group (both, stage 4 pulmonary carcinoma; 1 respectively 2.5 months after premature discontinuation).
** See also table 5, bone health.
†† One patient in each group had a history of hyperglycemia.
‡‡ One patient admitted twice for cataract surgery, thus both classified as SAE.
§§ ‘Other’ SAE: events in other organ classes. ‘Other’ other AESI: non-serious AE outside of the above predefined categories, but associated with premature discontinuation. AESI, adverse event of special interest; comprises serious adverse events (SAE) and ‘other AESI’.
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the duration of our trial. For example, pooled data of trial patients continuing certolizumab showed worse rates of AE, SAE and especially infection compared with our active treatment group. The recently published ORAL surveillance study compared tofacitinib and adalimumab in patients aged 50+ with one cardiovascular risk factor; tofacitinib did not meet noninferiority criteria, with substantially increased rates (20%–60%) of death, cardiovascular events, thromboembolism, infection, and cancer. Regarding infection, observational studies of biologics also show increased infections in aged patients. So, with cautious interpretation because of the indirect comparisons, our harm results suggest that the risks of low-dose GC are not of special concern but should be viewed through the same lens as those of other DMARDs.

Strengths of this study include its sample size, focus on older patients seen in routine care, detailed documentation of AE and its initiation by investigators. The missing data at study end, mostly caused by the COVID-19 crisis, are a weakness, and necessitate caution in interpretation. For efficacy, we have confidence in our model approach, and our results are in line with the literature. For safety, the high rates of events in both groups over the observation period make it unlikely that the risk estimate would be substantially different in a more complete data set. The pragmatic design is both strength and weakness: results are immediately applicable to the target population, but long-term treatment benefits were probably underestimated due to confounding. We assume but did not test the success of

Figure 3 Patients with multiple adverse events (AE) have a major impact on the total number of events in both treatment groups. Infection events in top panels, all AE in bottom panels. Left panels show the distribution of patients by number of events/patient. This shows most patients experience no or only a few events. Right panels plot bivariate cumulative distributions of patients by AE. The numbers next to the series indicate the maximum number of events of the population at that point. The right panels show the impact of multiple events: for example, in the top right panel, 75% of patients (with 0 or 1 infection) contribute only 25% of all infections. Another example, in the bottom right panel, read in reverse, 50% of patients (with 2 respectively 3 or more events) contribute almost 90% of all AE. To facilitate interpretation, grey vertical bars indicate 10% on the vertical scale.
blinding. In theory, detectable effects of the active drug might especially inflate patient-reported outcomes. However, response was by no means guaranteed, placebo response was substantial, patients could also receive (open label) co-treatment, and the treatment contrast was seen across all measures. Other potential weaknesses include one-sided testing (but results are also significant on two-sided testing), and lack of power to detect small, but possibly relevant differences in any of the areas of concern. Two years is not long for a chronic disease, but longer trials are hardly feasible in this population. Also, effects of confounding cointerventions (including short-term GC use) would increasingly hamper interpretation. Longer term observational studies could still be of use, if they feature prospective high-quality data collection (including detailed documentation of disease activity over time, dose over time, the motivation for a certain dose and dose changes) and analyses with several prespecified models.

In current practice, many patients with RA are chronically treated with low-dose GC, in direct contradiction with guidelines that prescribe only short-term “bridge” therapy in view of the perceived long-term adverse effects. Our study adds substantial evidence to support practice rather than guidelines: add-on chronic prednisolone at 5 mg/day for up to 2 years is effective and not particularly dangerous compared with alternatives. With proper monitoring, prevention and treatment of harmful effects, especially infections and bone loss, titrating around this level will allow optimum suppression of disease activity.

In conclusion, add-on low-dose prednisolone has long-term effects in senior patients with RA on optimum treatment, with a favourable balance of benefit and harm.

### Author affiliations

1. Epidemiology & Data Science, Amsterdam University Medical Centers, Vrije Universiteit, Amsterdam, The Netherlands
2. Rheumatology, Amsterdam Rheumatology and Immunology Center, Amsterdam University Medical Centers, Vrije Universiteit, Amsterdam, The Netherlands
3. Rheumatology, Carol Davila University of Medicine and Pharmacy, Romania, Bucharest, Romania
4. Rheumatology, Medical Centre Leeuwarden, Leeuwarden, The Netherlands
5. Rheumatology and Clinical Immunology, Maastricht University, Maastricht, The Netherlands
6. Reumatologie, Faculdade de Medicina e Hospital da Universidade de Coimbra, Coimbra, Portugal
7. Rheumatology, Antonius Hospital, Sneek, The Netherlands
8. Rheumatology, Meander Medisch Centrum, Amersfoort, The Netherlands
9. Rheumatology, Leiden University Medical Center, Leiden, The Netherlands
10. Rheumatology, Reumazorg Flevoland, Emmeloord, The Netherlands
11. Rheumatology, Northwest Clinics, Alkmaar, The Netherlands
12. Rheumatology, Faculty of Medicine, University of Debrecen, Debrecen, Hungary
13. Rheumatology and Clinical Immunology, Charité – University Medicine Berlin, Berlin, Germany
14. National Institute of Rheumatic Diseases, Piestany, Slovakia
15. Laboratory of Experimental Rheumatology and Academic Division of Clinical Rheumatology, Department of Internal Medicine, University of Genova, Genova, Italy

### Correction notice

This article has been corrected since it was first published. The open access licence has been updated to CC BY.

### Acknowledgements

We thank all patients and trial collaborators for their participation in the trial. We thank M. El-Alii for additional analyses performed in the review phase.

### Collaborators

Contributors The corresponding author is guarantor and accepts full responsibility for the work and/or the conduct of the study, had access to the data, and controlled the decision to publish. He affirms that the manuscript is honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as originally planned (and, if relevant, registered) have been explained. All authors meet the ICMJE criteria: substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work; or drafting the work or revising it critically for important intellectual content; and final approval of the version to be published; and agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. MB: corresponding author, Concept, data curation, analysis, funding, data collection, methodology, supervision, visualization, writing draft, writing review. LH: data curation, analysis, data collection, writing draft, writing review. DG: funding, data collection, writing review. RB: data collection, writing review. MRK: data collection, writing review. JAPDS: concept, funding, data collection, writing review. EC: data collection, writing review. BK: data collection, writing review. CB: data collection, writing review. FB: Concept, data collection, writing review. PM: funding, data collection, writing review. TK: data curation, analysis, methodology, writing review. SP: data collection, writing review. AMS: data collection, writing review. WL: data collection, supervision, writing review. MC: Concept, funding, data collection, writing review.

Funding The trial is part of a larger project funded by the European Union’s Horizon 2020 research and innovation program under grant agreement Number 634886. The funder had no role in the design, collection, analysis or interpretation of the data, writing of the report or the decision to publish.


Patient and public involvement Patients and/or the public were involved in the design, conduct, or reporting, or dissemination plans of this research. Refer to the Methods section for further details.

Patient consent for publication Not applicable.

Ethics approval This study involves human participants and was approved by Ethics Committee (Medisch Ethische Toetsingscommissie) VU Medisch Centrum, Amsterdam, Netherlands. ID reference number: 2015.471NL55263.029.15. Participants gave informed consent to participate in the study before taking part.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request. All data of this trial will be made available to interested researchers upon reasonable request and at fair use cost.

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ORCID iDs Maarten Boers http://orcid.org/0000-0002-6969-283X Linda Hartman http://orcid.org/0000-0003-4583-962X Marc R Kok http://orcid.org/0000-0003-2394-6926 Jose AP Da Silva http://orcid.org/0000-0002-7282-6780 Zoltan Szekanez http://orcid.org/0000-0003-2534-550X Willem F Lems http://orcid.org/0000-0002-5396-0932

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