Study of active controlled monotherapy used for rheumatoid arthritis, an IL-6 inhibitor (SAMURAI): evidence of clinical and radiographic benefit from an x ray reader-blinded randomised controlled trial of tocilizumab

Norihiro Nishimoto, Jun Hashimoto, Nobuyuki Miyasaka, Kazuhiko Yamamoto, Shinichi Kawai, Tsutomu Takeuchi, Norikazu Murata, Désirée van der Heijde, Tadamitsu Kishimoto

Objective: To evaluate the ability of tocilizumab (a humanised anti-IL-6 receptor antibody) monotherapy to inhibit progression of structural joint damage in patients with RA.

Methods: In a multi-centre, x ray reader-blinded, randomised, controlled trial, 306 patients with active RA of <5 years’ duration were allocated to receive either tocilizumab monotherapy at 8 mg/kg intravenously every 4 weeks or conventional disease-modifying antirheumatic drugs (DMARDs) for 52 weeks. Radiographs of hands and feet were scored by the van der Heijde modified Sharp method.

Results: Patients had a mean disease duration of 2.3 years and a disease activity score in 28 joints of 6.5 at baseline. Mean total modified Sharp score (TSS) was 29.4, which was very high despite the relatively short disease duration. At week 52, the tocilizumab group showed statistically significantly less radiographic change in TSS (mean 2.3; 95% CI 1.5 to 3.2) than the DMARD group (mean 6.1; 95% CI 4.2 to 8.0; p<0.01). Tocilizumab monotherapy also improved signs and symptoms. The overall incidences of AEs were 89% and 82% (serious AEs: 18% and 13%; serious infections: 7.6% and 4.1%) in the tocilizumab and DMARD groups, respectively.

Conclusion: Tocilizumab monotherapy was generally well tolerated and provided radiographic benefit in patients with RA.

Methods

Patients

Eligible patients were age >20 years and fulfilled the American College of Rheumatology (ACR; formerly, the American Rheumatism Association) 1987 revised criteria for the classification of RA18 with a disease duration of ≥6 months and <5 years. In addition, they had ≥6 tender joints (of 49 evaluated), ≥6 swollen joints (of 46 evaluated), an erythrocyte sedimentation rate (ESR) of ≥30 mm/h and C-reactive protein (CRP) of ≥20 mg/l. All candidates had an inadequate response to at least one disease modifying antirheumatic drug (DMARD) or immunosuppressant. Use of anti-TNF agents and leflunomide were not allowed within 3 months prior to the first dose. Change in dose and type of DMARDs and/or immunosuppressants, plasma exchange therapies and surgical treatments were not allowed within 4 weeks. Oral corticosteroids (prednisolone, ≤10 mg per day) were allowed if the dosage had not been changed within 2 weeks. Eligible patients had white blood cell counts of at least 3.5×10^9/l, lymphocyte counts of at least 0.5×10^9/l and platelet counts of at least 100×10^9/l at enrolment. Patients were excluded if they had a medical history of a serious allergic reaction, significant concomitant diseases, or an active intercurrent infection requiring medication within 4 weeks before the first dose. Sexually active premenopausal women were required to have a negative urine pregnancy test at the entry and to use effective contraception during the study period.

Study protocol

This study was conducted at 28 sites in Japan. The study protocol was approved by the Ministry of Health, Labor and
Anti-IL-6R antibody therapy in RA

Enrolled 306

Randomisation

DMARDs group 148

Not treated 3

Completed 131 (90%)

Withdrawn 14 (10%)

AE: 5

Exacerbation of disease: 3

Refused treatment: 4

Protocol violation: 2

Tocilizumab group 158

Not treated 1

Completed 134 (85%)

Withdrawn 23 (15%)

AE: 17

Exacerbation of disease: 1

Refused treatment: 1

Protocol violation: 1

Anti-tocilizumab antibodies: 3

Welfare of Japan, and by the local ethical committee, and patients gave their written informed consent. This trial was registered with http://clinicaltrials.gov (NCT00144508).

Patients were randomly assigned to receive either tocilizumab monotherapy at 8 mg/kg intravenously every 4 weeks or conventional DMARD therapy for 52 weeks. A long-term placebo-controlled study in RA patients with highly active disease was not acceptable from an ethical point of view, and therefore DMARDs were used for the controls. The randomisation was performed by registering of patients at the patient registration centre with a centralised allocation method. For the tocilizumab group, DMARDs and/or immunosuppressants were discontinued from the start of the study. Oral corticosteroids (≤ 10 mg prednisolone per day) were allowed, but the dosage could not be increased during the study. Intra-articular corticosteroid injections were not allowed. Use of one non-steroidal anti-inflammatory drug (NSAID), including switching to another NSAID, was allowed. For the conventional DMARD group, the dose, type and combination of DMARDs and/or immunosuppressants, except for anti-TNF agents and leflunomide, could be varied according to disease activity at the discretion of the treating physician. Variations of NSAIDs and/or corticosteroids including intra-articular corticosteroid injections were also allowed. Surgical treatment and use of bisphosphonates was not allowed in either group. Safety was assessed through recording of adverse events, physical examinations, and standard laboratory tests for both groups.

Radiographic and clinical assessment

Posteroanterior radiographs of hands and anteroposterior radiographs of feet were performed at baseline, week 28 and week 52 or at the last visit for patients who withdrew from the study prior to week 52. For patients who dropped out before week 52, missing radiograph values at week 52 were estimated by linear extrapolation using data at baseline, week 28 and the early termination visit. Radiographs were scored using van der Heijde’s modified Sharp method, independently by two readers who were well trained and competent to score radiographs. The readers were blinded to the treatment group, chronologic order of the films and clinical response of each patient. Ten percent of the patients’ films were re-read for the analysis of intra-reader variability.

ACR20, 50 and 70 responses, and disease activity score in 28 joints (DAS28) were assessed for clinical improvement of RA using an intent-to-treat (ITT) analysis.

Statistical analysis

A sample size of 120 patients per treatment group was estimated to provide 80% power for detecting a significant (p≤0.05) difference in mean change score of radiographic findings between the tocilizumab and DMARD groups. We decided to recruit 150 patients per treatment group to allow for anticipated withdrawals. Radiographic endpoints, such as TSS, erosion score and joint space narrowing score, were assessed with a rank transformed analysis of covariance (ANCOVA) on the change scores that included factors for baseline score and baseline disease duration. The incidences of clinical improvements were analysed by the chi-square test.

All statistical analyses were two-sided and p values less than 0.05 were considered significant. All patients receiving at least one dose of study drug were included in the efficacy and safety analysis.

RESULTS

Characteristics of the patients

This study enrolled 306 patients in total (Figure 1). Four patients were withdrawn before treatment either due to their ineligibility or at the patients’ request. A total of 302 patients received study drugs. A total of 134 patients in the tocilizumab group and 131 patients in the DMARDs group completed 52 weeks treatment. Discontinuation occurred in 23 patients in the tocilizumab group and 14 patients in the DMARDs group. The reported reasons for withdrawal are shown in Figure 1.

Demographics and baseline disease characteristics did not differ between the two groups (Table 1).

Mean disease duration was 2.3 years. Patients had active disease, indicated by a DAS28 score of 6.5 and CRP of 48 mg/L at baseline. Moreover, TSS at baseline was 29.4, which was very high despite the relatively short disease duration. The mean estimated yearly progression rate, calculated from the baseline TSS divided by disease duration for each patient, was 13.3 Sharp units.

Treatment in the conventional DMARD group

At baseline, 67% of the patients in the DMARDs group received methotrexate (MTX): 37% received a combination of MTX and DMARDs, 30% received MTX monotherapy, and 22% received DMARDs and/or immunosuppressants other than MTX, besides corticosteroids. The dose of MTX was 7.1 ± 1.9 mg/week (mean ± SD) in patients treated with MTX. During the study, 123 patients (85%) received MTX: 81 (56%) received a
The combination of MTX and DMARDs, 42 (29%) received MTX monotherapy, and 20 (14%) received DMARDs and/or immunosuppressants other than MTX, besides corticosteroids. The dose of MTX was 8.0 ± 2.1 mg/week in patients treated with MTX (Japanese government recommends 6–8 mg/week of MTX based on the evidence from the Japanese clinical trial of MTX for RA).26 27 Besides MTX, salazosulfapyridine (41%), bucillamine (23%), mizoribine (8%) and D-penicillamine (8%) were frequently used in more than 5% of the patients.

Reliability of radiographic scoring
Intra-reader intraclass correlation coefficients for erosion, joint space narrowing and TSS were all 0.99 for both readers. Inter-reader intraclass correlation coefficients for erosion, joint space narrowing and TSS were 0.98, 0.96 and 0.98, respectively.

Radiographic evaluation of joint damage
Figure 2 shows the cumulative probability plots of the change from baseline to week 52 in the TSS. The space between the curves indicates different treatment effects with a considerable difference in favour of the tocilizumab group. The efficacy was more evident at week 52. In addition to the TSS and erosion score, joint space narrowing scores also showed significantly less change in the tocilizumab group than in the DMARDs group in an ANCOVA model (Table 2).

The mean changes in the TSS as well as erosion scores at week 28 were statistically significantly less in the tocilizumab group compared with the DMARDs group after adjustment for baseline values with a linear regression model (Table 2).


docilizumab monotherapy had negative TSS scores than those receiving conventional DMARDs (24 patients and 18 patients at week 52, respectively).

The mean changes in the TSS as well as erosion scores at week 28 were statistically significantly less in the tocilizumab group than in the DMARDs group with an ANCOVA model (Table 2).

Table 2 Change in radiographic scores

<table>
<thead>
<tr>
<th></th>
<th>Conventional DMARDs (n = 143)</th>
<th>8 mg/kg Tocilizumab (n = 157)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Week 28</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total Sharp score Mean (95% CI)</td>
<td>4.5 (3.1 to 6.0)</td>
<td>1.9 (1.2 to 2.6)*</td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>1.0 (0.0 to 5.0)</td>
<td>0.5 (0.0 to 2.0)</td>
</tr>
<tr>
<td>Erosion score  Mean (95% CI)</td>
<td>2.4 (1.6 to 3.2)</td>
<td>0.8 (0.4 to 1.2)†</td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>0.5 (0.0 to 2.5)</td>
<td>0.0 (0.0 to 1.0)</td>
</tr>
<tr>
<td>Joint space narrowing Mean (95% CI)</td>
<td>2.2 (1.4 to 2.9)</td>
<td>1.1 (0.7 to 1.6)</td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>0.0 (0.0 to 2.0)</td>
<td>0.0 (0.0 to 1.0)</td>
</tr>
<tr>
<td><strong>Week 52</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total Sharp score Mean (95% CI)</td>
<td>6.1 (4.2 to 8.0)</td>
<td>2.3 (1.5 to 3.2)†</td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>2.5 (0.0 to 7.0)</td>
<td>0.5 (0.0 to 3.0)</td>
</tr>
<tr>
<td>Erosion score  Mean (95% CI)</td>
<td>3.2 (2.1 to 4.3)</td>
<td>0.9 (0.3 to 1.4)‡</td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>1.0 (0.0 to 3.5)</td>
<td>0.0 (0.0 to 1.0)</td>
</tr>
<tr>
<td>Joint space narrowing Mean (95% CI)</td>
<td>2.9 (2.0 to 3.8)</td>
<td>1.5 (0.9 to 2.1)‡</td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>1.0 (0.0 to 4.0)</td>
<td>0.0 (0.0 to 1.7)</td>
</tr>
</tbody>
</table>

*p < 0.05, †p < 0.01, ‡p < 0.001.

P values were analysed with a rank transformed analysis of covariance (ANCOVA) on the change scores that included factors for baseline score and baseline disease duration, DMARDs, disease-modifying antirheumatic drugs; Tocilizumab, humanised anti-interleukin-6 receptor antibody; 95% CI, 95% confidence interval; IQR, interquartile range.
Clinical efficacy
At week 52, proportions of the patients achieving ACR20, ACR50, and ACR70 response were 78%, 64%, and 44% in the tocilizumab group and 34%, 13%, and 6% in the DMARD group, respectively, indicating the superiority of tocilizumab monotherapy to conventional DMARD therapy (p < 0.001, for each comparison) although clinical efficacy was assessed unblinded (Figure 3A).

Greater reduction in DAS28 scores and higher remission rates were also observed in the tocilizumab group than in the DMARDs group (Figure 3B). At week 52, clinical remission (defined as DAS28 < 2.6) was achieved in 59% of patients receiving tocilizumab, but only in 3% of patients receiving DMARDs (p < 0.001). Major clinical response (ACR70 response for 6 consecutive months) was achieved in 24% of patients receiving tocilizumab compared with only 2% of patients receiving DMARDs during the study period of 52 weeks.

Physical function and health-related quality of life
Tocilizumab monotherapy significantly improved MHAQ scores compared to conventional DMARDs (Figure 3C). A decrease of ≥ 0.22 units in HAQ scores represents significant clinical improvement and the minimum clinically important difference. Such improvement was seen in 40% of the patients treated with tocilizumab as early as week 4, the first scheduled study visit, and was even more evident at week 52 (68% in the tocilizumab group and 40% in the DMARDs group, p < 0.001).

Safety
The percentages of patients with adverse events were 89% and 82% in the tocilizumab and DMARD groups, respectively. Most of adverse events were mild or moderate. Table 3 shows frequent adverse events observed in at least 5% of the patients. Nasopharyngitis was the most common adverse event, but the incidences were similar in both groups.

Serious adverse events were reported in 18% and 13% in the tocilizumab group and DMARDs group, respectively. In the tocilizumab group, 12 serious infections were reported: 3 (1.9%) patients with pneumonia, 2 (1.3%) with upper respiratory tract infection, 2 (1.3%) with cellulitis, 1 (0.6%) each with gastroenteritis, herpes zoster, herpes simplex, perianal abscess and an unidentified infection. In the DMARD group, 8 serious infections were reported: 3 (2.1%) patients with gastroenteritis, 2 (1.4%) with pneumonia, and 1 (0.7%) each with upper respiratory tract infection, herpes zoster and sepsis. All the serious adverse events improved with appropriate treatment. There was no significant prolongation of infection by the tocilizumab treatment. Tuberculosis was not observed in this 1-year study without required screening or prophylactic use of any antituberculous drug.

Three malignancies were reported in the tocilizumab group: 2 patients with breast cancer (including 1 lobular carcinoma in situ) and 1 with colon cancer, which were improved or resolved by appropriate treatment (including surgery). No malignancies were reported in the DMARD group.

Drug-related infusion reactions were reported 14 times in 11 (7.0%) patients of the tocilizumab group: 3 with transient increase in blood pressure, 2 with injection site redness, 2 with headache, 2 with nausea, 2 with skin eruption, and 1 each with vomiting, pruritus, and malaise. All the infusion reactions were mild, and no patient withdrew from the study as a consequence.

Laboratory test abnormalities were reported in 61% and 31% of patients in the tocilizumab and DMARD groups, respectively. In the tocilizumab group, lipid metabolism-related reactions were common. Anomalous increases in total cholesterol (TC), triglycerides, and low-density lipoprotein cholesterol were reported in 38%, 17%, and 26% of the patients, respectively, and most of them were grade 1 according to the National Cancer Institute Common Toxicity Criteria. Twenty-seven patients were treated (HMG-CoA reductase inhibitor, 26 cases; fenofibrate, 1 case) and their cholesterol levels improved during the study. Tocilizumab monotherapy also raised high-density lipoprotein cholesterol (HDLc) levels to above the normal range.
in 24% of patients. The atherogenic index, calculated by (TC–
HDL)/HDL, did not change during the study period of
52 weeks. No cardiovascular complications were observed in
association with abnormal lipid profile.

Anti-tocilizumab antibodies were detected in 4 patients
(2.5%). Only one patient showed a skin eruption at the third
injection, while the other three were asymptomatic. They were
all withdrawn according to the study protocol.

**DISCUSSION**

This 52-week, x ray reader-blinded, randomised, controlled trial
demonstrated that tocilizumab monotherapy in patients with
active RA significantly inhibited the progression of structural
joint damage compared with conventional DMARDs therapy.
Note that even monotherapy with tocilizumab significantly
retarded the radiological progression. It is of interest whether
tocilizumab in combination with MTX would provide greater
benefit; this is being investigated in the European studies.

The results of this study confirmed that IL-6 plays a
pathological role in the joint destruction in RA. IL-6 blockade
can inhibit the osteoclast activation in RA. Additionally,
tocilizumab therapy reduced MMP-3 levels (data not shown),
which could also contribute to the radiographic benefit.

In addition to the radiographic benefits, tocilizumab mono-
therapy improved signs and symptoms as well as functional
evaluation with MHAQ. Although this was an open-label study
for clinical efficacy endpoints, the results of previous phase II
studies21 22 were confirmed. Moreover, significant improvement
in MHAQ scores indicates that tocilizumab improves patients’
daily living activity.

There was no decrease in the efficacy of tocilizumab during
the 1-year treatment. The benefit of a humanised antibody was
again emphasised in the repetitive treatment, as anti-tocilizu-
amb antibodies were detected in only 4 patients (2.5%) without
requiring the use of immunosuppressive agents such as MTX.
It could also be that tocilizumab that blocks IL-6 action to induce
B cell differentiation into antibody producing cells.

Tocilizumab monotherapy was generally well tolerated. There
was no specific infection related to tocilizumab therapy,
including tuberculosis, which is often a problem in anti-TNF
therapy,11 although patients had neither prophylactic medica-
tion nor screening in this study.

Three malignancies occurred in patients treated with
tocilizumab in this study. There are many reports describing
the relationship between IL-6 and malignant diseases, where
IL-6 is often a causative factor.11 Further analysis to compare
the incidence with an epidemiology surveillance data base
regarding malignancies in Japanese patients with RA will be
required to ascertain the carcinogenesis risk or the benefit of IL-
6 inhibition.

The increase in TC is also observed in anti-TNF therapy23
and therefore it could be secondary to the improvement in
inflammation. We have reported that tocilizumab treatment
improves wasting in Castleman’s disease, an atypical lympho-
proliferative disease with overproduction of IL-6 but not TNFα,
where tocilizumab monotherapy normalises hypocholesterole-
ma but seldom causes hypercholesterolemia.34 Therefore, IL-6
plays a role in regulating serum cholesterol levels.

As an increase in serum IL-6 levels has been reported as a
cardiovascular risk24 25 and as IL-6 could contribute to the
atherosclerosis,35 it is of interest whether tocilizumab therapy
reduces the incidence of cardiovascular events. This issue will
be also proven in a future epidemiological surveillance of
patients treated long-term with tocilizumab.

In conclusion, this study clearly demonstrates the superiority
of tocilizumab monotherapy in preventing joint damage to
conventional DMARDs in Japanese RA patients. However the
results need to be confirmed in the trials in western RA
patients.

**ACKNOWLEDGEMENTS**

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analysis of the study and preparation of this manuscript. This work was
supported by Chugai Pharmaceutical Co., Ltd., Tokyo, Japan.

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Competing interests: NN has served as a consultant to and/or received
honoraria from Chugai Pharmaceutical, the manufacturer of tocilizumab.
TK holds a patent for tocilizumab. The other authors have no competing
interests.

**Table 3** Adverse events observed in at least 5% of patients

<table>
<thead>
<tr>
<th></th>
<th>Conventional DMARDs (n = 145)</th>
<th>8 mg/kg Tocilizumab (n = 157)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rash</td>
<td>47 (32.4)</td>
<td>56 (35.7)</td>
</tr>
<tr>
<td>Diarrhoea</td>
<td>13 (9.0)</td>
<td>13 (8.3)</td>
</tr>
<tr>
<td>Headache</td>
<td>3 (2.1)</td>
<td>11 (7.0)</td>
</tr>
<tr>
<td>Stomatitis</td>
<td>13 (9.0)</td>
<td>9 (5.7)</td>
</tr>
<tr>
<td>Eczema</td>
<td>6 (4.1)</td>
<td>9 (5.7)</td>
</tr>
<tr>
<td>Nausea</td>
<td>2 (1.4)</td>
<td>9 (5.7)</td>
</tr>
<tr>
<td>Pruritus</td>
<td>2 (1.4)</td>
<td>9 (5.7)</td>
</tr>
<tr>
<td>Paronychia</td>
<td>1 (0.7)</td>
<td>9 (5.7)</td>
</tr>
<tr>
<td>Vomiting</td>
<td>5 (3.4)</td>
<td>8 (5.1)</td>
</tr>
<tr>
<td>Vertebral compression fracture</td>
<td>8 (5.5)</td>
<td>3 (1.9)</td>
</tr>
</tbody>
</table>

*Values are the number (%) of patients.

DMARDs, disease-modifying antirheumatic drugs; Tocilizumab, humanised
anti-interleukin-6 receptor antibody.

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