Tocilizumab: The evidence for its place in the treatment of juvenile idiopathic arthritis

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Introduction: Juvenile idiopathic arthritis (JIA) is one of the most common chronic diseases with childhood onset. It comprises different subtypes of which the systemic onset subtype is often resistant to treatment. With the advent of biological treatment with tumor necrosis factor-α (TNFα)-inhibitors, the clinical outcome of JIA has improved considerably, but only for subtypes other than systemic JIA. Substantial evidence shows that the proinflammatory cytokine interleukin-6 (IL-6) plays a pivotal role in systemic JIA. The blockage of IL-6 action by tocilizumab, a humanized anti-IL-6-receptor monoclonal antibody, could therefore be an effective treatment of systemic JIA.

Aims: The purpose of this article was to review the clinical trials of tocilizumab and to discuss its place in the treatment of JIA with the focus on the systemic onset of disease.

Evidence review: Two phase II studies and one phase III clinical trial of tocilizumab demonstrating the clinical efficacy and safety in systemic onset JIA have been published. Within those studies, sustained and high response rates of clinical improvement have been achieved with American College of Rheumatology Pediatric criteria (ACRPed) 30, 50, and 70 observed in 98%, 94%, and 90% of patients, respectively, after 48 weeks. One study regarding the clinical efficacy of tocilizumab for the treatment of oligo- and polyarticular JIA has been presented only as a conference abstract.

Place in therapy: The very promising results seen so far in patients with severe systemic JIA and acceptable tolerability gives tocilizumab a central role in the future therapy in controlling this disease. No other biological therapy has achieved similar high response rates when treating with tocilizumab 8 mg/kg every two weeks to patients with systemic onset JIA, but direct comparison of the efficacy of different biological agents are not yet available.

Keywords: tocilizumab, anti-IL-6-receptor antibody, biologics, systemic, juvenile idiopathic arthritis

Core evidence place in therapy summary for tocilizumab in the treatment of juvenile idiopathic arthritis

<table>
<thead>
<tr>
<th>Outcome measure</th>
<th>Evidence</th>
<th>Implications</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient-oriented evidence</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Improvement of symptoms</td>
<td>Clear</td>
<td>Reduction of joint pain and improvement of joint motion</td>
</tr>
<tr>
<td>Reduction of fever</td>
<td>Substantial</td>
<td>Rapid normalization of temperature</td>
</tr>
<tr>
<td>Tolerability</td>
<td>Clear</td>
<td>Few infusion reactions</td>
</tr>
<tr>
<td>Long-term safety</td>
<td>Limited</td>
<td>Upper respiratory tract infections observed but long-term observation are not at hand</td>
</tr>
</tbody>
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(Continued)
Scopes, aims, and objectives
Tocilizumab (Actemra®, Chugai Pharmaceutical Co., Ltd. and F Hoffmann-La Roche) is a humanized anti-interleukin-6 (IL-6)-receptor antibody used in the targeted therapy of rheumatoid arthritis (RA) and juvenile idiopathic arthritis (JIA). Tocilizumab blocks the activity of the proinflammatory cytokine, IL-6, which exerts a central role in both diseases. Within recent years, tocilizumab has been used for RA patients with treatment-resistant disease. The aim of this article was to review the clinical trials of tocilizumab for the use in systemic onset JIA and to discuss its role in the treatment strategy for this disease.

Methods
A review of the medical literature regarding tocilizumab was performed. Articles related to tocilizumab on PubMed (http://www.ncbi.nlm.nih.gov) using the search terms “tocilizumab” (117), “tocilizumab AND juvenile idiopathic arthritis” (26), and “anti-IL-6-receptor blockade AND juvenile idiopathic arthritis” (9) were selected for the review. The search was updated on February 20, 2009. Articles not written in English were excluded. Furthermore, the search term “tocilizumab AND rheumatoid arthritis” (79) was used to review clinical trials on adult patients with RA. In addition, selected abstracts from the Annual Meetings of the American College of Rheumatology (ACR) and of the European League Against Rheumatism (EULAR) in 2007 and 2008 were used.

Disease overview
Juvenile idiopathic arthritis is a collective term for different patterns of arthritis of unknown cause in children. All of them are defined as chronic arthritis lasting for more than six weeks in the absence of any known cause in a child aged under 16 years. JIA is classified according to the onset of the disease into seven subtypes: systemic, persistent oligoarticular, extended oligoarticular, rheumatoid factor-positive polyarticular, rheumatoid factor-negative polyarticular, psoriatic, and enthesitis-related arthritis subtypes. The disease is among the most frequent chronic diseases starting in childhood and in population-based studies using the International League of Associations for Rheumatology (ILAR) criteria, an annual incidence of 14–15 per 100,000 has been reported. Systemic onset JIA, representing about 10% of JIA, is a subtype quite distinct from the other subtypes characterized by a quotidian, spiking fever lasting more than two weeks, nonfixed erythematous skin rash, generalized lymphadenopathy, hepatosplenomegaly, or serositis. Arthralgia and myalgia can be prominent during fever attacks. The arthritis, which may not be present during the early phase of the disease, is often presented as a symmetrical polyarticular arthritis. Marked increases in the acute phase reactants are observed resulting in at times very high levels of C-reactive protein (CRP) and erythrocyte sedimentation rate (ESR). Leukocytosis, predominantly neutrophils, is commonly observed, as well as thrombocytosis, which may reflect the exacerbations of the disease.

The course of systemic onset JIA can be monocyclic, polycyclic, or persisting with destructive arthritis. The overall outcome of the disease is poor with a high risk of long-term functional impairment. A severe, life-threatening complication to systemic onset JIA is macrophage activation syndrome, a secondary form of hemophagocytic lymphohistiocytosis, presented with prolonged fever, pancytopenia, elevated liver transaminases, coagulopathy, and high levels of triglyceride and ferritin. Furthermore, growth failure, osteoporosis, secondary amyloidosis are other severe complications of the disease.

<table>
<thead>
<tr>
<th>Outcome measure</th>
<th>Evidence</th>
<th>Implications</th>
</tr>
</thead>
<tbody>
<tr>
<td>Disease-oriented evidence</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Reduction in synovitis</td>
<td>Clear</td>
<td>Improvement in number of swollen joints and joints with limitation in motion</td>
</tr>
<tr>
<td>Reduction of anemia</td>
<td>Clear</td>
<td>Rapid increase in hemoglobin</td>
</tr>
<tr>
<td>Reduction of inflammatory response during treatment</td>
<td>Substantial</td>
<td>Rapid decrease in CRP, ESR, neutrophils and platelet count</td>
</tr>
<tr>
<td>Maintenance of response during treatment</td>
<td>Clear</td>
<td>Long-term efficacy only during treatment</td>
</tr>
<tr>
<td>Economic evidence</td>
<td>Unclear</td>
<td>Long-term pharmacoeconomic studies missing</td>
</tr>
</tbody>
</table>

Abbreviations: CRP, C-reactive protein; ESR, erythrocyte sedimentation rate.
The role of IL-6 in the pathogenesis of JIA

Interleukin (IL)-6 is a 26 kD glycoprotein consisting of 184 amino acids which has previously been assigned different names such as B-cell differentiation factor (BCDF), Interferon-β2, a hybridoma/plastocytoma growth factor. Later, these names were replaced by IL-6. IL-6 is produced by a variety of cells of which several participate during inflammation such as T- and B-cells, monocytes, fibroblasts, and endothelial cells.

Function of IL-6 is mediated through two membrane proteins, the membrane-bound IL-6R, an 80 kD ligand-binding receptor, and gp130, a 130 kD signal transducing element. In addition there is a soluble form of IL-6R present in serum. When binding to IL-6, both soluble and membrane IL-6R connects to gp130 on the membrane whereby the intracellular signal activating the Jak-STAT pathway is mediated.8,9

During acute inflammation IL-6 stimulates hepatocytes to produce acute-phase proteins such as CRP, haptoglobin, fibrinogen, α1-antitrypsin, or serum amyloid A. IL-6 stimulates the production of hepcidin, an iron regulatory peptide hormone which reduces iron transport in the intestines and inhibits the release of iron from macrophages leading to anemia. Differentiation of osteoclasts is stimulated by IL-6 leading to joint destruction and osteoporosis.

Abnormal expression of the proinflammatory cytokines IL-1, IL-6, and tumor necrosis factor-α (TNFα) is seen in systemic JIA. It has been demonstrated by several studies that IL-6 is significantly elevated in the blood and synovial fluid of patients with systemic JIA.10–13 The level of IL-6 fluctuates with the characteristic spiking fever, which is not the case for the increased levels of TNFα. The high levels of IL-6, also being responsible for the impaired growth, thrombocytosis and chronic anemia, lead to the suggestion that systemic JIA is a IL-6-mediated disease.14 IL-6 and IL-1RA, being induced by IL-6, correlates with systemic JIA disease activity, which is not the case for IL-1 and TNFα.15 IL-6 form complexes with its soluble receptor, sIL-6R, which trigger gp130 dimerisation and signaling. Also, elevated concentration of sIL-6R has been found during active disease.16

In family studies the -174G allele of IL-6 gene has been confirmed as a susceptibility gene for systemic JIA, since it correlates with higher levels of IL-6.17,18

Outcome measures

The American College of Rheumatology Pediatric (ACRPed) criteria for the definition of improvement of the clinical outcome measures for JIA was proposed by Giannini and colleagues.19 A patient is considered to have responded to therapy (ACRPed30) if there is an improvement in at least three out of six variables (active joint count, number of joints with limited range of motion, parent/patient global assessment on a visual analogue scale, physician global assessment on a visual analogue scale, first hour ESR (mm/hour), Childhood Health Assessment Questionnaire) by at least 30% and worsening in not more than one variable by more than 30% (Table 1). For the ACRPed50 and ACRPed70, an improvement of at least 50% or 70%, respectively, must be achieved in three out of six of the above variables with worsening in not more than one variable by more than 30%.

Current therapeutic options

Treatment of patients with systemic JIA remains a challenge. Many different treatment regimens with immunosuppressants have been used. Although corticosteroids have been the mainstay of therapy for children with active systemic JIA, there is no evidence that systemic corticosteroids are disease modifying. Due to unacceptable, deleterious long-term side effects such as growth retardation, osteoporosis, and diabetes, reduction of corticosteroid dosage and prevention of prolonged therapy is mandatory. Instead of high-dose daily oral corticosteroids pulses with high-dose intravenous methylprednisolon (up to 30 mg/kg per dose, maximal 1000 mg have been proposed to minimize side effects.20 In a few patients, nonsteroidal anti-inflammatory drugs (NSAIDs) alone have been effective. More often methotrexate either given orally or subcutaneously have been used as a second-line medication in systemic JIA but has been less effective compared to the use in treating polyarticular JIA.21 Biological therapy with anti-TNFα agents have limited effect in systemic JIA with response rates as low as 30%.22,23 Pascual and colleagues24 demonstrated that sera from nine patients with systemic JIA provoked IL-1 synthesis

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Core set of criteria for evaluating change in treatment trials of children with juvenile idiopathic arthritis: American College of Rheumatology Pediatric response criteria19</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Number of active joints (swollen joints and/or two or three of the following: limitation of range of motion, pain on motion, and increased heat of joints).</td>
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<tr>
<td>2.</td>
<td>Number of joints with limited range of motion.</td>
</tr>
<tr>
<td>4.</td>
<td>Patient/parent assessment of overall well being.</td>
</tr>
<tr>
<td>5.</td>
<td>Functional ability (Childhood Health Assessment Questionnaire).</td>
</tr>
</tbody>
</table>

Notes: Improvement is at least 30% improvement from baseline in three of any six variables with no more than one of the remaining variables worsening by >30%.
in tissue culture of mononuclear cells. In seven of these nine
patients, treatment with anakinra, an anti-IL-1 receptor
antagonist, gave a complete clinical response. In an open-label
study, 20 patients with systemic JIA refractory to corticosteroids
were treated with anakinra 1–2 mg/kg/day. Only moderate
response rates were found with an ACRPed30, 50, and 70 of
55%, 30%, and 0%, respectively, after three months and after
six months 50%, 25% and 10%, respectively.25 A multicenter,
randomized double-blind trial of anakinra versus placebo in
systemic JIA with 12 patients in each group has been con-
ducted and preliminary results were presented.26 Achievement
of ACRPeds0 response along with resolution of fever and
normalization of CRP after one month was seen in 8/12 in
the anakinra group compared to 1/12 in the placebo group.
Because daily subcutaneous injections are required and are
painful, the practical use of anakinra has been limited.
A long-acting soluble receptor-based IL-1 blocker, rilonacept
(IL-1 Trap), has been investigated in a phase II trial in
systemic JIA. Preliminary results published in abstract form
from the open-label phase including 21 patients have revealed
improvement after four weeks with ACRPed30, 50, and
70 responses of 76.2%, 61.9%, and 33%, respectively.27 Fever
and/or rash was completely resolved. Sustained response
was recorded in 12 patients. Treatment with rilonacept was
generally safe and well tolerated.

**Tocilizumab**

The blockade of the biological functions of IL-6 may theoretically
be achieved through a blockade of the gp130 receptor
by neutralizing IL-6 or by the prevention of the IL-6/IL-6R
complex formation. Since the gp130 homodimer is shared
among other receptors than for IL-6, the blockade of gp130
may implicate a wide range of undesirable effects. Attempts
of trials with chimaeric anti-IL-6 monoclonal antibody have
failed to block IL-6 function.28,29

Tocilizumab, formerly myeloma receptor antibody
(MRA), is a humanized anti-IL-6R monoclonal antibody
(humanized by the technique of complementary-determining
region grafting from mouse anti-human IL-6R monoclonal
antibody) engrafted with a human IgG1 Fc to minimize
potential immunogenic responses in humans.30 It has a high
affinity to IL-6R and abrogates the IL-6 signaling by prevent-
ing the formation of the IL-6/IL-6R complex.

The clinical efficacy of tocilizumab in preliminary stud-
ies has previously been described in murine experimental
arthritis,31 in patients with RA,32–34 multicentric plasma cell
type or mixed type Castleman’s disease35 and adult-onset
Still’s disease.36

**Clinical efficacy of tocilizumab in adults with RA**

Tocilizumab was launched in April 2008 in Japan and
was approved by the European Agency for the Evaluation
of Medicinal Products (EMEA) in January 2009 for the
treatment of moderate and severe RA, but has not yet been
approved by the US Food and Drug Administration (FDA).
Results from randomized controlled trials of tocilizumab in
RA are summarized in Table 1. In the CHARISMA study,37
a phase II study evaluating the efficacy of tocilizumab in
359 patients with active RA and with an inadequate response
to methotrexate, patients were randomized to seven study
arms evaluating the effect of three doses of tocilizumab: 2, 4,
and 8 mg/kg with or without methotrexate plus methotrexate
alone. The infusion of tocilizumab was given at four-week
intervals. ACR20 responses were achieved in all tocilizumab
arms at doses of 4 and 8 mg/kg (Table 2).

Four phase III randomized, double-blind placebo-
controlled trials evaluating the effect of tocilizumab versus
placebo in a total of 2,467 patients with RA have been pub-
lished since 2008.38–41 In all studies with RA, tocilizumab was
infused at four-week intervals. The most significant results
were observed with doses of 8 mg/kg with achievement
of ACR20 responses of 50%–80.3%, ACR50 responses of
29%–49%, and ACR70 responses of 12.4%–29.5%. Remission,
defined as Disease Activity Score (DAS28) <2.6,
were observed in 27%–43.1% of patients receiving 8 mg/kg
doses compared with 7.6%–13% of patients receiving 4 mg/kg
in the OPTION and RADIATE trials.

In the SAMURAI study, a randomized X-ray double-
blind controlled trial, Nishimoto and colleagues42 evaluated
the ability of tocilizumab monotherapy against conventional
DMARDs to inhibit progression of structural joint dam-
age in RA patients. After 52 weeks, the tocilizumab group
showed significantly less radiographic change in the van
der Heijde score (modified Sharp score) (mean 2.3, 95%
confidence interval [CI]: 1.5–3.2; p < 0.01) compared to
the DMARD group (mean 6.1, 95% CI: 4.2–8.0; p < 0.01).
Erosion score was 0.9 and joint space narrowing was 1.5 in
the tocilizumab group versus 3.2 and 2.9 in the DMARD
group, respectively.

**Clinical efficacy of tocilizumab in systemic onset JIA**

Two phase II studies of the blockade of IL-6 signaling using
anti-IL-6 receptor antibody (MRA, tocilizumab) for systemic
JIA suggested that it was a highly effective treatment.43,44
Tocilizumab in juvenile idiopathic arthritis

Table 2 Randomized controlled trials of tocilizumab in rheumatoid arthritis

<table>
<thead>
<tr>
<th>Trial name</th>
<th>Author</th>
<th>Design</th>
<th>No of patients</th>
<th>Duration</th>
<th>Study arms tocilizumab in mg/kg/4 weeks</th>
<th>ACR20</th>
<th>ACR50</th>
<th>ACR70</th>
<th>DAS28 &lt; 2.6</th>
</tr>
</thead>
<tbody>
<tr>
<td>CHARIMA</td>
<td>Maini et al27</td>
<td>R-DB-PC</td>
<td>359</td>
<td>16 wk</td>
<td>0 + mtx</td>
<td>41</td>
<td>29</td>
<td>8</td>
<td>–</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>2/2 + mtx</td>
<td>31/64</td>
<td>6/32</td>
<td>2/14</td>
<td>–</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>4/4 + mtx</td>
<td>61/63</td>
<td>28/37</td>
<td>6/12</td>
<td>–</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>8/8 + mtx</td>
<td>63/74</td>
<td>41/53</td>
<td>16/37</td>
<td>–</td>
</tr>
<tr>
<td>OPTION</td>
<td>Smolen et al38</td>
<td>R-DB-PC</td>
<td>623</td>
<td>24 wk</td>
<td>0 + mtx</td>
<td>26</td>
<td>11</td>
<td>2</td>
<td>0.8</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>4 + mtx</td>
<td>48</td>
<td>31</td>
<td>12</td>
<td>13</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>8 + mtx</td>
<td>59</td>
<td>44</td>
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<tr>
<td>TOWARD</td>
<td>Genovese et al39</td>
<td>R-DB-PC</td>
<td>1220</td>
<td>24 wk</td>
<td>0 + DMARD</td>
<td>25</td>
<td>9</td>
<td>3</td>
<td>3</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>8 + DMARD</td>
<td>61</td>
<td>38</td>
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<td>30</td>
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<td>RADIATE</td>
<td>Emery et al40</td>
<td>R-DB-PC</td>
<td>499</td>
<td>24 wk</td>
<td>0 + mtx</td>
<td>10</td>
<td>4</td>
<td>1.3</td>
<td>1.6</td>
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<tr>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>4 + mtx</td>
<td>30</td>
<td>17</td>
<td>5</td>
<td>7.6</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>8 + mtx</td>
<td>50</td>
<td>29</td>
<td>12.4</td>
<td>30.1</td>
</tr>
<tr>
<td>SATORI</td>
<td>Nishimoto et al41</td>
<td>R-DB-PC</td>
<td>125</td>
<td>24 wk</td>
<td>placebo + mtx</td>
<td>25</td>
<td>10.9</td>
<td>6.3</td>
<td>1.6</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
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<td>toc 8 + placebo</td>
<td>80.3</td>
<td>49.2</td>
<td>29.5</td>
<td>43.1</td>
</tr>
</tbody>
</table>

Abbreviations: ACR, American College of Rheumatology; DAS, Disease Activity Score; DMARD, disease-modifying antirheumatic drug; mtx, methotrexate; R-DB-PC, randomized, double-blind, placebo-controlled.

Yokota and colleagues42 evaluated the safety and efficacy of tocilizumab in a dose-escalating protocol (MRA011JP) in 11 Japanese children (3–18 years) with active systemic onset JIA refractory to corticosteroids and/or DMARDs. Primarily Yokota and colleagues infused a 2 mg/kg/dose three times with a two-week interval. The dose was increased to 4 mg/kg and administered three times if CRP level were above 1.5 mg/dl after the first and second infusion and further to 8 mg/kg three times if CRP was still above 1.5 mg/dL. Tocilizumab abruptly reduced disease activity in 10 of the 11 children two weeks after the administration of the drug. Three children had sustained response to 2 mg/kg, five of the remaining eight children were stabilized with 4 mg/kg, and three children received 8 mg/kg before CRP was kept below 1.5 mg/dL. In total, ACRPed30 and ACRPed50 was achieved in 10 children and ACRPed70 was achieved in seven children.

In another open-labelled trial (LRO320), 15 of 18 enrolled Caucasian children with systemic JIA received a single dose of either 2 mg/kg (n = 4), 4 mg/kg (n = 6), or 8 mg/kg (n = 5) tocilizumab by infusion.44 All had active disease for at least three months with at least one active systemic feature, one active joint, and prednisolone doses above 0.2 mg/kg/day and in addition 12 also received methotrexate (<20 mg/m²/week). Although only a small number of patients were studied, there was a marked improvement in all patients within all three dose groups 48 hours after the infusion and at one-week follow-up, 11 patients achieved ACRPed30, eight patients achieved ACRPed50, and three achieved ACRPed70.

However, a prolonged clinical response was only observed for 4 mg/kg and 8 mg/kg with an ACRPed30 obtained in 6/9 patients. Along with the clinical improvement, a marked decrease in CRP and ESR after one week was followed by an increase shortly after.

While writing this review, only one randomized double-blind, placebo-controlled withdrawal study (MRA316JP) of the effect of tocilizumab for the treatment of Japanese children with systemic JIA has been published.43 The study included an open-label lead-in phase of six weeks, a double-blind, randomized, placebo-controlled withdrawal phase of 12 weeks, and an open-label extension phase of 48 weeks or more. Fifty-six children (21 boys, 35 girls) with active systemic JIA defined by an increase in CRP > 15 mg/L and inadequate clinical response to ≥0.2 mg/kg/day of prednisolone were enrolled in the open-label lead-in phase. They received 8 mg/kg tocilizumab three times every two weeks with a primary endpoint being ACRPed30 (Table 3). After six weeks, ACRPed30, 50, and 70 was achieved in 91%, 86%, and 68%, respectively. Forty-three patients continued to the double-blind withdrawal phase and were randomly assigned to either receive an infusion of tocilizumab 8 mg/kg or placebo every two weeks for 12 weeks. In the tocilizumab group 80% maintained an ACRPed30 or ACRPed50 response and a CRP concentration <15 mg/L after 12 weeks compared to 17% in the placebo group (p < 0.0001) (Table 4). Half of the children in the placebo group who did not maintain ACRPed30 relapsed within four weeks. In the open-labeled extension phase, 48 of the
50 patients continued to receive tocilizumab after 48 weeks. Achievement of ACRPed30, 50, and 70 was observed in 98%, 94%, and 90%, respectively. The median percentage change in ESR and CRP was −93.2% and −99.7%, respectively, along with a significant increase in hemoglobin.

In 2008, an ongoing two-part phase III multicenter study (WA18221) consisting of a part I 12-week randomized, double-blind, placebo-controlled two-arm study evaluating the efficacy and safety of tocilizumab in patients with systemic JIA was launched with a subsequent part II 92-week single-arm open-label extension study to examine the long-term use of tocilizumab. Pediatric rheumatology centers from Europe, North America, and South America are participating in the study. The patients are randomized to either 8 mg/kg for patients >30 kg every two weeks and 12 mg/kg for patients <30 kg or placebo infusions every two weeks. The patients (n = 108) are unequally randomized (tocilizumab/placebo ratio 2:1). The primary efficacy endpoints are the proportion of patients with at least 30% improvement in JIA core set (ACRPed30) at week 12 and absence of fever. The rationale for using the increased dose (12 mg/kg) for small children <30 kg was chosen as a consequence of the pharmacokinetic, safety, and efficacy data from the MRA316JP study.

**Clinical efficacy in oligo- and polyarticular onset JIA**

In a case report, a 26-year-old woman with oligoarthritis for 12 years was described having developed amyloidosis in the kidney and intestines. The patient responded well to the treatment of intravenous 8 mg/kg tocilizumab given every 3–4 weeks.46

One prospective, open-labeled, multicenter study has been performed in patients with polyarticular or oligoarticular onset JIA with a polyarticular course.47 The children were treated with 8 mg/kg tocilizumab every four weeks for 48 weeks. Concomitant treatments with other biologics, DMARDs, or immunosuppressants were not allowed. Nineteen patients were enrolled in the study with the primary efficacy endpoint being the proportion of patients achieving ACRPed30, 50, and 70 every 12 weeks from the baseline evaluation. They found a significant decrease in all core set parameters with an ACRPed30 of 94.1% after 12 weeks. The response continued until the evaluation after 48 weeks where 11 completed the assessments (ACRPed30, 100%; ACRPed50, 100%; ACRPed70, 90.9%).

### Safety and tolerability

In the phase II study, Yokota and colleagues,45 reported that the drug was generally well tolerated at all doses. Two patients had upper respiratory tract infections and three had pustules on the extremities. In four patients, cholesterol increases were observed, two patients showed elevated alanine aminotransferases, and two showed glucosuria. Woo and colleagues44 reported no evidence of dose-limiting toxicity. Fifty-nine adverse events were reported in 15 patients with mainly gastroenteritis and respiratory disorders. Three patients had increased alanine aminotransferases and one had urticaria. No serious bacterial infections were seen.

Two serious adverse events were reported in the lead-in phase of the phase III study (MRA316JP):45 one with anaphylaxis and one with gastrointestinal hemorrhage from colonic ulceration in a patient with previous history of diarrhea and rectal bleeding. In the double-blind phase, one patient had infectious mononucleosis with highly elevated liver transaminases and neutropenia and one developed herpes zoster.

To evaluate the long-term safety of tocilizumab, 128 patients with systemic JIA including patients who completed the MRA011JP and MRA316JP studies and an additional 61 patients were enrolled.48 Tocilizumab was administered as an infusion of 8 mg/kg every two weeks. The median duration of tocilizumab treatment was 78 weeks. Adverse events occurred in 120 patients (94%) at a rate of 787 per 100 patient-years. Serious adverse events occurred at an incidence rate of 37.2 per 100 patient years and serious infections at a rate of 14.5 per patients-years. The most frequent serious

### Table 3 Response to tocilizumab 8 mg/kg every two weeks in patients with systemic onset juvenile idiopathic arthritis in a six week open-label lead-in phase45

<table>
<thead>
<tr>
<th>Number of pts. (%)</th>
<th></th>
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<tbody>
<tr>
<td>ACRPed30</td>
<td>51/56 (91%)</td>
</tr>
<tr>
<td>ACRPed50</td>
<td>48/56 (86%)</td>
</tr>
<tr>
<td>ACRPed70</td>
<td>38/56 (68%)</td>
</tr>
</tbody>
</table>

**Abbreviation:** ACRPed, American College of Rheumatology Pediatric criteria.

### Table 4 Maintained ACRPed response after a 12-week double-blind phase continuing with either tocilizumab 8 mg/kg every two weeks or placebo45

<table>
<thead>
<tr>
<th></th>
<th>Tocilizumab</th>
<th>Placebo</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACRPed30</td>
<td>16/20 (80%)</td>
<td>4/23 (17%)</td>
<td>p &lt; 0.0001</td>
</tr>
<tr>
<td>ACRPed50</td>
<td>16/20 (80%)</td>
<td>4/23 (17%)</td>
<td>p &lt; 0.0001</td>
</tr>
<tr>
<td>ACRPed70</td>
<td>15/20 (75%)</td>
<td>3/23 (13%)</td>
<td>p &lt; 0.0001</td>
</tr>
</tbody>
</table>

**Abbreviation:** ACRPed, American College of Rheumatology Pediatric criteria.
infections were gastroenteritis (3.8/100 patient-years) and pneumonia (3.4/100 patient-years). In total, 14 patients were withdrawn from the study. Eight patients were withdrawn from the study due to adverse events which included macrophage activation syndrome, anaphylactoid reaction (two patients), cardiac amyloidosis, duodenal perforation, gastrointestinal hemorrhage, and infusion reactions (two patients). Furthermore, five patients were withdrawn from the study due to the development of antitocilizumab antibodies and one due to lack of efficacy.

**Place in therapy for tocilizumab in JIA**

With the advent of biological response modifiers, commonly referred to as biologics, the therapeutic options for JIA have improved considerably. The use of TNFα inhibitors for the treatment of JIA with a polyarticular course refractory to conventional methotrexate has shown efficacy in controlled clinical trials with etanercept, infliximab, and adalimumab. A long-term follow-up study with etanercept has shown a sustained effect and with relatively few serious adverse events and high tolerability.

However, children with systemic onset JIA may often be refractory to treatment with both conventional DMARDs and biologics. Response rates as low as 30% using etanercept have been shown and often unacceptably high doses of corticosteroids are used to control the disease. Only one phase III double-blind placebo-controlled withdrawal study with tocilizumab has been published so far in systemic JIA.

The study design, combining an open-label lead-in phase with a placebo-controlled withdrawal phase, has been used for studying biologics in JIA randomized controlled trials for the last decade and is considered ethically acceptable for the use in children. Tocilizumab infusions of 8 mg/kg every other week achieved high response rates with an ACR50 of 86%. Compared to this, tocilizumab 8 mg/kg given every four weeks to adult patients with RA resulted in an ACR50 of 29%–49% in four phase III studies.

Response to tocilizumab therapy is seen with few weeks and even days. The drug is generally well tolerated with an incidence of serious adverse events of 37/100 patient-year. The most frequent serious infections are gastroenteritis and pneumonia, at 3.8 and 3.4/100 patient-years, respectively.

If the ongoing double-blind placebo-controlled tocilizumab randomized controlled trial (WA18221) for systemic JIA shows comparable results as in the Japanese study, tocilizumab could be considered as a first-line drug in corticosteroid-resistant systemic JIA. However, direct comparison of treatment with tocilizumab and TNFα-inhibitors or other biologics like abatacept or IL-1R inhibitors in head-to-head or crossover studies could be warranted to elucidate its role in the therapeutic option of systemic JIA.

There is still very limited information regarding the pharmacoeconomic aspects of the treatment of children with JIA with tocilizumab.

In summary, tocilizumab is an effective drug for the treatment of systemic onset JIA. It represents an interesting option for patients refractory to or dependent on corticosteroid therapy. Full data are not yet available to place its role in the treatment of non-systemic JIA where other biologics have shown a higher efficacy. However, the significant effect on adult RA may indicate that tocilizumab could also have a promising role for polyarticular JIA.

**Disclosure**

The author reports no conflicts of interest in this work.

**References**


