Treatment of systemic-onset juvenile arthritis with canakinumab

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Abstract: Treatment of systemic-onset juvenile idiopathic arthritis is challenging, but the availability of cytokine antagonists targeting interleukin-1 and interleukin-6 have markedly advanced the therapeutic options. In this review, we focus on the current experience with canakinumab, an interleukin-1 monoclonal human antibody for the treatment of systemic-onset juvenile idiopathic arthritis and describe its efficacy and safety. Canakinumab is an important, safe, and valid drug in the treatment of systemic-onset juvenile idiopathic arthritis.

Keywords: anakinra, canakinumab, interleukin-1, interleukin-6, systemic-onset juvenile idiopathic arthritis

Juvenile idiopathic arthritis and systemic-onset juvenile idiopathic arthritis

Juvenile idiopathic arthritis (JIA) includes a group of diseases with arthritis lasting more than 6 weeks, of no apparent cause, and an onset prior to the age of 16 years. It is the most common chronic rheumatic disease in children. Seven JIA categories are defined (Table 1), with differences in clinical presentation, course, prognosis, and response to treatment with disease-modifying drugs. Differences in the biology and pathogenesis of the disease may be responsible for the variability in responses to medical treatment.

Systemic-onset juvenile idiopathic arthritis (soJIA) is characterized by the variable occurrence of chronic aggressive arthritis, intermittently high spiking fever, maculopapular rash (often described as salmon-like colored) during fever episodes, hepatomegaly and splenomegaly, lymphadenopathy, serositis, and a marked increase in acute-phase reactant levels. These findings make it unique among the categories of JIA.

SoJIA, which is the most severe category of JIA, accounts for about 4%–17% of JIA cases. The diagnosis is applied when children up to 16 years of age present with arthritis and fever of at least 2 weeks’ duration, a spiking appearance and spontaneous disappearance of fever documented for at least 3 days, and the presence of at least one of following: erythematous rash, generalized lymphadenopathy, hepatomegaly, splenomegaly, or serositis. Furthermore, diagnosis of soJIA requires exclusion of other diseases possibly related to the clinical findings.

Given the as yet unknown pathogenesis and extreme heterogeneity of soJIA, the course of the disease varies between individuals, ranging from a single appearance of the disease, to recurrent predominantly systemic disease courses, to a progressive polyarthritis that frequently leads to severe and destructive joint disease. Complications
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Table 1 Classification of juvenile idiopathic arthritis

<table>
<thead>
<tr>
<th>Subclassification</th>
<th>Category</th>
<th>Main extra-articular manifestations</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Systemic-onset arthritis (Still’s disease)</td>
<td>Fever, rash, hepatosplenomegaly, pericarditis, pleuritis, lymphadenopathy, vasculitis, short stature, dyspophy</td>
</tr>
<tr>
<td>2</td>
<td>Seronegative polyarthritis</td>
<td>Tenosynovitis, uveitis</td>
</tr>
<tr>
<td>3</td>
<td>Seropositive polyarthritis</td>
<td>Low-grade fever, tenosynovitis, rheumatoid nodules</td>
</tr>
<tr>
<td>4</td>
<td>Oligoarthritis</td>
<td>Chronic uveitis</td>
</tr>
<tr>
<td>5</td>
<td>Arthritis and enthesitis</td>
<td>Enthesitis, acute uveitis</td>
</tr>
<tr>
<td>6</td>
<td>Psoriasis and arthritis</td>
<td>Psoriasis, uveitis</td>
</tr>
<tr>
<td>7</td>
<td>Unclassified JIA</td>
<td>Variable</td>
</tr>
</tbody>
</table>

Abbreviation: JIA, juvenile idiopathic arthritis.

of soJIA include growth impairment, osteoporosis, and the macrophage activation syndrome, which is potentially lethal (Table 2).6–8

The aim of treatment is to reach inactive disease or even remission as defined in Table 3. Current treatments for soJIA have proved largely unsatisfactory.9–12 Management of the disease relies on corticosteroids. Children with soJIA do not respond well to disease-modifying agents such as methotrexate, and poor responses have also been reported with newer agents, such as anti-tumor necrosis factor.13 Several reports have suggested a major role for cytokines in the disease, such as interleukin (IL)-6 and, more recently, IL-1β.14–21

Role of interleukin-1

IL-1 is a proinflammatory cytokine that is produced by monocytes/macrophages and dendritic cells. Its stimulatory effects

Table 2 Criteria for macrophage activation syndrome in systemic-onset juvenile idiopathic arthritis

<table>
<thead>
<tr>
<th>Laboratory findings</th>
<th>Glutamic oxaloacetic transaminase (&gt;59 U/L)</th>
<th>Leukocytosis (&lt;4,000/μL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Central nervous system symptoms (seizures, coma, pain, irritability)</td>
<td>Hemorrhagia (purpura, hematoma, mucosa bleeding)</td>
<td>Hepatomegaly</td>
</tr>
<tr>
<td>Bone marrow puncture in bone marrow</td>
<td>Diagnosis</td>
<td></td>
</tr>
<tr>
<td>MAS needs at least two laboratory criteria or two clinical and/or laboratory criteria</td>
<td>Bone marrow puncture is necessary in uncertain cases only</td>
<td></td>
</tr>
</tbody>
</table>

Note: Data from Ravelli et al.6
Abbreviation: MAS, macrophage activation syndrome.

Table 3 Criteria for inactive disease in juvenile idiopathic arthritis

| “Wallace” criteria for inactive disease in oligoarticular (persistent and extended), polyarticular (RF− and RF+), and systemic JIA |
| Inactive disease |
| No joints with active arthritis |
| No fever, rash, serositis, splenomegaly, or generalized lymphadenopathy attributable to JIA |
| No active uveitis to be defined |
| ESR or CRP level within normal limits in the laboratory where tested; if both are tested, both must be normal |
| Physician’s global assessment of disease activity score of best possible on the scale used |

For systemic JIA, all criteria must be met

Abbreviations: RF, rheumatoid factor; eSR, erythrocyte sedimentation rate; CRP, C-reactive protein; JIA, juvenile idiopathic arthritis.

The biologic effects of IL-1 can be neutralized as outlined in Figure 1. The first published report of successful treatment of systemic JIA with IL-1 inhibition occurred in 2004 with the report of a remarkable response in two patients whose severe disease manifestations were previously refractory to other therapies.22 At about the same time, other researchers found that serum from children with soJIA induced the transcription of IL-1β-related genes in the peripheral blood mononuclear cells of healthy controls. Based on this finding, these investigators treated patients with soJIA with anakinra, an IL-1 inhibitor, and achieved a dramatic clinical response. For example, seven of nine patients who were refractory to prior therapies achieved remission. These data prompted clinicians to use anakinra for the treatment of soJIA in clinical practice, as reported in several case series. An early report showed a remarkable response to treatment with anakinra in ten of 21 patients. In these patients, it seemed that those with fewer involved joints might benefit more from anakinra therapy than those with arthritis in many joints.18 A larger retrospective case series of 46 patients with soJIA was limited to children who received anakinra as part of their initial glucocorticoid-sparing treatment regimen or even as
first-line monotherapy. Fifty-nine percent of these patients showed a good clinical response. Of ten patients who received anakinra as first-line therapy without any glucocorticoids, eight achieved a complete remission.

In addition to anakinra, other IL-1 inhibitors have been developed and subsequently studied for sJIA (Table 4). Currently, there are three different biological inhibitors of the IL-1 pathway available: anakinra, an IL-1 receptor antagonist with a relatively short half-life; canakinumab, a human IL-1β antibody with a long plasma half-life; and rilonacept, an IL-1 receptor fusion protein.

**Canakinumab in sJIA**

Canakinumab is a human monoclonal antibody (Table 4). Its half-life in plasma is 23–26 days, which is much longer than that of IL-1β. It binds IL-1β selectively and prevents binding of IL-1β to the IL-1 receptor. Furthermore, canakinumab does not bind IL-1α or the IL-1 receptor antagonist, and does not interact with the IL-1 receptor.

methotrexate (up to a maximum of 20 mg/m²/week) were permitted. Major exclusion criteria were concomitant treatment with another biologic or disease-modifying drug, diagnosis of macrophage activation syndrome within the previous 6 months, active tuberculosis, and a live vaccination within 3 months prior to enrollment.

The primary outcome criterion was achievement of at least an adapted PedACR30 response and no fever. Patients with continuing fever could be unblinded from day 3, and if they had received placebo, they were offered canakinumab in an open trial arm. Six patients (14.0%) on canakinumab and 37 (90.2%) on placebo discontinued prematurely, all due to an unsatisfactory response. On day 15, 36/43 (83.7%) patients in the canakinumab group and 4/41 (9.8%) of patients in the placebo group met the primary endpoint of a modified PedACR30 with simultaneous defervescence. The difference was highly significant (Figure 2). Furthermore, at day 15, 14 (32.6%) patients on canakinumab versus none on placebo reached inactive disease status. One patient in each group developed a macrophage activation syndrome, and also one patient in each group got a serious infection.

Patients in the above-mentioned first controlled trial could change to a second study with a withdrawal design. Seventy-one patients from this trial, ten from the above-mentioned Phase II trial, and 96 newly enrolled patients contributed to the total of 177 patients who received canakinumab 4 mg/kg, with a maximum of 300 mg every 4 weeks. In the open-label phase of the study, all patients received canakinumab. In this first part of the study, the primary outcome criterion was a reduction in the steroid dose taken by at least 25% of patients at entry to the study by week 20. For this, a standardized corticosteroid-tapering protocol was permitted from week 9 to week 28 patients who achieved at least an adapted PedACR50 response. Corticosteroids were used by 128 of 177 patients (72.3%) patients at study entry. Forty-two of 128 patients (32.8%) were able to stop corticosteroids on treatment with canakinumab and 57 of 128 patients could reduce their steroid doses. The mean daily dose of corticosteroids decreased from 0.34 mg/kg to 0.05 mg/kg. Thus, the primary goal of this part of the study was reached. The secondary endpoint of the study was the proportion of patients achieving an adapted PedACR30/50/70/90/100 response or inactive disease. At day 15, 81.3%, 74.1%, 58.3%, 34.5%, 18.0%, and 16.4% already fulfilled these response criteria (Figure 3).

If patients improved by reaching the goal of decreasing pre-existing steroid doses or discontinued steroids, they were then transferred into the second part of this study, ie, the double-blind, placebo-controlled withdrawal phase. One hundred

**Figure 1** Interaction of IL-1 and IL-1 receptor antagonist with the IL-1 receptor molecules.

**Notes:** The binding of IL-1 leads the IL-1 receptor to a steric approximation of the two chains of the receptor (IL1-R1 and IL1-RacP) and triggers activation of the T-cell. The natural IL1-RA binds to the IL1-R1, with no association with IL1-RacP. Thus, anakinra, a recombinant modified IL1RA, inhibits the activation of IL1-α and IL1-β. IL1-β is specifically bound by canakinumab. Rilonacept inhibits IL1-α and IL1-β.

**Abbreviations:** IL, interleukin; IL1-RA, IL-1 receptor antagonist; IL1-RacP, IL-1 receptor accessory protein.
Canakinumab in systemic-onset juvenile arthritis

Patients qualified and were admitted to this randomized part of the study in which occurrence of a predefined flare of the disease was the primary outcome criterion. In this part, the median time to onset of a disease relapse was 236 days (95% confidence interval 141–449) in the placebo cohort and was not determinable for canakinumab, since less than 50% of the patients ever had a relapse. This corresponds to a relative risk reduction of 63% (Figure 4). Patients who flared were immediately transferred to the open-label extension part of the study. The withdrawal phase was terminated after 37 patients had flared. Subsequently, all patients were transferred from an observational study to an ongoing open-label extension.

Thirty-nine (78%) patients remained flare-free in the canakinumab arm versus 24 (48%) in the placebo arm, with a statistically significant 63% relative risk reduction for experiencing a soJIA flare (hazard ratio 0.37; 95% confidence interval 0.17–0.78; \( P = 0.0043 \)). Corticosteroids were used by 128/177 (72.3%) of patients at study entry.

At the end of the second part of the second trial, 41/50 (82%) patients in the canakinumab group had a minimum adapted PedACR70 response and 31/50 (62%) had inactive disease status compared with 31/50 (62%) and 16/50 (32%), respectively, in the placebo group (Figure 5).

Canakinumab in other immune-mediated diseases

In a three-part, 48-week, double-blind, placebo-controlled, randomized withdrawal study of canakinumab in patients
with cryopyrin-associated autoinflammatory syndrome, treatment with subcutaneous canakinumab once every 8 weeks was associated with a rapid remission of symptoms in most patients. Following this study, canakinumab became approved as the first IL-1 blocker in 2011.

In a two-part, open-label, single-arm Phase II study, the long-term therapeutic effect of canakinumab was demonstrated in subjects aged 4–16 years with colchicine-resistant familial Mediterranean fever and a history of at least three documented attacks in the 3 months prior to enrolment. Treatment with canakinumab in an 8-year-old girl with a clinical picture of hyperimmunoglobulinemia D syndrome, that had started before the end of the first year of life, resulted in disappearance of febrile attacks and a considerable improvement in the patient’s quality of life during a 12-month follow-up period. In this case report, the drug was well tolerated, and no side effects were observed.

Figure 3 Response according to PedACR30, PedACR50, and PedACR100 criteria in a double-blind study in patients with systemic-onset juvenile idiopathic arthritis.

Notes: The primary endpoint was PedACR30 and resolution of fever, and was achieved significantly more often in the canakinumab group. (A) Fourteen patients had a PedACR100 response on day 15. (B) The effect was unchanged until the 29th day. *P<0.0001, **P=0.0001.

Abbreviation: PedACR, American College of Rheumatology Pediatric Criteria.
produced a rapid and highly effective clinical and serological benefit which was maintained for at least 4 months.26 A randomized, double-blind, placebo-controlled study of canakinumab in patients with these hereditary autoinflammatory disorders (tumor necrosis factor receptor-associated periodic fever syndrome, hyperimmunoglobulinemia D syndrome, familial Mediterranean fever) is currently recruiting.

In a 9-month study, monthly injection of canakinumab was an effective and well tolerated treatment for Schnitzler’s syndrome, a chronic disabling autoinflammatory disorder characterized by chronic urticaria, paraproteinemia, and severe systemic inflammation,27 as shown in other case reports.28

In a report of three adult patients with Behçet’s disease refractory to treatment with corticosteroids and different combinations of immunosuppressant agents who received canakinumab, prompt and sustained clinical efficacy was demonstrated, and canakinumab was seen as a therapeutic option for resistant or refractory Behçet’s disease.29 In this single-center, open-label pilot study, the effects of canakinumab on clinical signs and symptoms, quality of life, inflammation markers, and cytokine levels were investigated in ten patients with active urticarial vasculitis, and the efficacy and safety of canakinumab was documented.30

Khanna et al systematically reviewed the published data on the pharmacologic and nonpharmacologic agents used for the treatment of acute gouty arthritis, and found evidence to suggest the efficacy of treatment of acute gout with canakinumab, on a comedication with nonsteroidal anti-inflammatory drugs, selective cyclooxygenase type 2 inhibitors, corticosteroids, and colchicine.31

Safety of canakinumab in soJIA
As far as we can see at the moment, adverse events with canakinumab are manageable. In most cases, the adverse events are infections. Nevertheless, in one study, macrophage activation syndrome occurred in two patients, one of whom died.35 Macrophage activation syndrome is a well-known potential lethal complication in autoinflammatory diseases, and especially in soJIA. Despite this report, canakinumab can be considered as well tolerated. Even in patients with comorbidities like type 2 diabetes mellitus, the adverse events reported were manageable,35 although long-term observation and further studies are needed.

Conclusion
The more we know about the pathophysiology of inflammation and its pathways, the more specific targets we can find to control it. IL-1 is a global player in inflammation, and by blocking it we have a way to block inflammation very specifically with few side effects. With improving genetics and panels of ILs, we might in future even know in advance which patients to treat with which drug in the form of personalized medicine. Canakinumab seems to be one of the drugs for this approach.

In summary, canakinumab seems to be very effective in the treatment of soJIA. Its subcutaneous application is of benefit for patients who need to treated for a prolonged period of time.

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References


