

# An Economic Comparison of Treatment Strategies with Anakinra in Systemic Juvenile Idiopathic Arthritis (sJIA)

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**Introduction:** Systemic juvenile idiopathic arthritis (sJIA) is a rare, complex autoinflammatory disease with substantial morbidity, often characterized by fever, rash, and muscle pain, amongst other symptoms. Biologic agents, such as anakinra, have been successfully used to treat patients internationally, but their usage in some regions is limited to patients that have failed to achieve clinically inactive disease with corticosteroids and conventional synthetic disease-modifying anti-rheumatic drugs (csDMARDs). Use of anakinra early in the disease course leads to better clinical outcomes; however, longer-term costs for this treatment strategy have not been established. This study compares the economic implications of first-line versus later-line availability of anakinra for patients with sJIA.

**Methods:** Data for patients treated with first-line anakinra were identified from a single-center, prospective study and compared to a combination of published trial and economic evaluation information to facilitate a comparison to later-line anakinra (ie, following corticosteroids + csDMARDs). Costs were estimated for product acquisition and medical resource utilization (MRU), including planned outpatient visits and unplanned hospital admissions. Total costs over a 5-year horizon were compared.

**Results:** Total 5-year product acquisition cost for the first-line anakinra strategy was €24,021, and for later-line anakinra was €20,471. The corresponding MRU costs were €19,197 (first-line) versus €25,425 (later-line). Overall 5-year costs (product acquisition and MRU) were lower for the first-line strategy (€43,218 versus €45,896).

**Conclusion:** The use of anakinra for patients with sJIA in the first-line setting is efficacious to induce and sustain inactive disease, and the findings of this study show that this treatment strategy leads to cost savings through reduced medical expenditure.

**Keywords:** Still's disease, systemic juvenile idiopathic arthritis, anakinra, economic comparison, treatment strategies, first line

## Plain Language Summary

Anakinra, a treatment for patients with systemic juvenile idiopathic arthritis (sJIA), has been used in patients with sJIA for over a decade. Different regions treat with anakinra at different stages in the treatment pathway. In some regions, anakinra is used as a first-line treatment option, whereas in others, its use is reserved for patients who fail to respond to treatment with non-steroidal anti-inflammatory drugs (NSAIDs) and conventional synthetic disease-modifying antirheumatic drugs (csDMARDs), or, in other words, “later-line” use of anakinra. This study presents the first economic comparison of different treatment strategies in sJIA. We identified data from a range of different sources and estimated the costs of total drug and medical resource use associated with each alternative treatment regimen. We then compared

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the total costs accrued over a five-year period. The results of our analysis show first-line anakinra treatment is associated with cost savings versus later-line use, when considered in the context of the full cost implications of patient management.

## Introduction

Systemic juvenile idiopathic arthritis (sJIA) is a rare, complex autoinflammatory disease, with an incidence of 0.5–0.9 per 100,000 children.<sup>1–4</sup> It is associated with substantial morbidity, and is usually characterized by spiking fever, rash, muscle pain, arthritic symptoms, in addition to liver and spleen enlargement.<sup>5–7</sup> With an average age of diagnosis between 3 and 5 years, children with sJIA have reduced quality of life, caused by impaired physical and social functioning, which is especially detrimental in developing years.<sup>8,9</sup>

While sJIA is often considered a subtype of juvenile idiopathic arthritis (JIA), its systemic nature differentiates it from other, more common rheumatic diseases in several aspects. It is understood to be a polygenic autoinflammatory disease characterized by an excessive activation of the innate immune system and secretion of interleukin-1 (IL-1), -6, and -18. It is therefore different to other forms of JIA that are considered to be autoimmune where the pathology is predominantly driven by the adaptive immune system.<sup>10</sup> sJIA is also described as the most severe and potentially life-threatening disease in the group of JIA.<sup>10</sup> More recently, sJIA and adult-onset Still's disease (AOSD) have been described as the same disease entity with different ages of onset.<sup>11</sup>

Owing to its systemic nature, sJIA is one of very few rheumatic diseases associated with the development of macrophage activation syndrome (MAS) – a potentially-fatal complication, recognized as the most frequent life-threatening complication of Still's disease.<sup>12,13</sup> Therefore, in addition to substantial morbidity, sub-optimal management of sJIA is associated with an increased risk of mortality.

Traditionally, the management of sJIA was concerned primarily with reducing the risk of long-term complications associated with the disease (including issues related to long-term steroid use, and the requirement for articular surgery), as opposed to achieving a state of clinically-inactive disease (CID, or “remission” – that is, an absence of disease-related symptoms and normalization of laboratory markers). This was a direct consequence of the lack of efficacious treatments that had been specifically studied within sJIA populations, and so clinical practice

historically comprised the use of unlicensed treatments that had been studied in other rheumatic diseases. These unlicensed treatment options for sJIA include non-steroidal anti-inflammatory drugs (NSAIDs), corticosteroids, and conventional-synthetic disease-modifying anti-rheumatic drugs (csDMARDs). Moreover, the only randomized, placebo-controlled trial of methotrexate (csDMARD) in patients with sJIA did not demonstrate a statistically significant improvement in the systemic features of the disease.<sup>14</sup>

While traditional therapies were able to improve symptoms, their mechanistic properties did not allow for patients to achieve a state of CID. Following increased understanding of the role of IL-1/IL-6 in sJIA, biologic DMARDs (bDMARDs) were studied in sJIA populations in a number of clinician-led studies.<sup>14–17</sup> Anakinra (Kineret<sup>®</sup>, Sobi), an IL-1 receptor antagonist, has been used to treat patients with sJIA for over a decade across Europe,<sup>18</sup> and was licensed for the treatment of Still's disease (including sJIA) in April 2018.<sup>19,20</sup> With the availability of bDMARDs targeting IL-1/IL-6, treatment goals shifted from management of symptoms to achieving CID.

Clinical studies have shown that it is possible to achieve CID with bDMARDs, though outcomes are particularly impressive for patients treated with anakinra shortly after diagnosis.<sup>21</sup> Cytokine blockade early in the course of Still's disease may allow patients to achieve remission before the development of chronic, destructive, and often therapy-resistant arthritis.<sup>20–25</sup>

A number of clinical guidelines have been updated to reflect the expected benefits associated with earlier use of bDMARDs, acknowledging more recent clinical studies.<sup>11,26,27</sup> Cohorts of sJIA patients in the Netherlands and Italy have been treated with first-line anakinra, and a majority of patients were shown to achieve CID, many of whom were able to discontinue therapy in the longer term.<sup>21,25,28,29</sup> Earlier use of anakinra has been shown to be associated with clinical benefits; however, it is important to note that this strategy may have an impact on the overall cost of managing patients. To establish its cost-effectiveness, a comparison of hypothetical treatment pathways (ie, use of anakinra early versus later in the treatment pathway) is required.

To our knowledge, no comparison of treatment strategies in sJIA from an economic perspective have been previously conducted. This study aimed to formally compare alternative treatment strategies with anakinra (where it is available in a first-line versus later line setting) in the

management of sJIA to illustrate the differences in expected costs.

## Materials and Methods

### Strategies to be Compared

This study considers a comparison of two treatment strategies:

1. First-line anakinra: This strategy is concerned with the use of anakinra as soon as possible after diagnosis, theoretically prior to the use of any active treatment for sJIA (including corticosteroids). This strategy is aligned with the latest Single Hub and Access point for pediatric Rheumatology in Europe (SHARE) treatment guidelines,<sup>11</sup> the Dutch Pediatrics Association guidelines (Nederlandse Vereniging voor Kindergeneeskunde, NVK),<sup>26</sup> and a number of recent studies.<sup>21,25,28,29</sup>
2. Later-line anakinra: This strategy involves the use of anakinra after the use of NSAIDs, corticosteroids, and at least one csDMARD (such as methotrexate). Use of anakinra in this setting is aligned with the use of anakinra in earlier published studies (such as ANAJIS<sup>15</sup>) and is representative of current practice in the UK (per NICE TA238 and TA685 guidance, and a National Health Service clinical commissioning policy).<sup>30–32</sup>

## Evidence Base

Most countries treat sJIA patients based on a national policy (including the UK and the Netherlands).<sup>26,30,31</sup> Therefore, it is unlikely that a head-to-head comparison of alternative treatment strategies in the same country would be possible (eg, comparing strategies where anakinra is available in the first-line setting, versus only being available in a later-line setting). Accordingly, data to inform both treatment strategies were sought from the literature, with the expectation that supporting studies are likely to have been conducted in different geographies. Where literature was unavailable, information from published reports (eg, NICE guidance) were considered, else assumptions were made where necessary (which are described where applicable throughout).

For patients treated using a first-line anakinra strategy, data regarding the duration of treatment and proportion of patients that achieve CID were taken from ter Haar et al.<sup>21</sup> This study reports on a population of n=42 patients with

sJIA treated with anakinra monotherapy according to a treat-to-target strategy (before the use of corticosteroids), and followed for a median of 5.8 years.<sup>21</sup> At 1 year, 76% of patients (30/42) had CID and 52% (22/42) had CID off medication.<sup>21</sup> For those that were followed up to 5 years, 96% (24/25) had CID, and 72% (18/25) had CID off medication.<sup>21</sup>

In contrast to a first-line treatment strategy, there is no equivalent, comprehensive study available to consider the costs and outcomes for a later-line anakinra treatment strategy. However, tocilizumab (RoActemra<sup>®</sup>, Roche, an IL-6 inhibitor) was previously assessed by the National Institute for Health and Care Excellence (NICE) in this setting (TA238), which is broadly aligned with the use of anakinra for sJIA in a later-line setting.<sup>30</sup> Therefore, data and assumptions used to inform NICE TA238 were used to produce estimated costs for the later-line anakinra setting, supported with additional information from the literature where needed.

Patients managed with a later-line anakinra strategy were assumed to be treated with NSAIDs, corticosteroids, and a csDMARD (methotrexate) prior to the initiation of bDMARD therapy. Approximately a quarter of patients (25.5%) were assumed to have monocyclic disease course,<sup>33</sup> for which it was assumed possible for patients to achieve CID prior to the initiation of bDMARDs. The remaining patients were assumed to have chronic disease course. For NSAIDs + corticosteroids, it was assumed that approximately 30% of monocyclic patients would achieve remission within 6 weeks, and for methotrexate, 20% would achieve remission within 24 weeks (based on the Nordström et al study in AOSD, where it was assumed all patients that achieved “remission” had monocyclic disease course<sup>17</sup>).

Approximately 50% of patients (with either monocyclic or chronic disease course) were assumed to achieve CID with the first bDMARD used based on the anakinra arm of the Nordström et al. study.<sup>17</sup> In the base-case analysis, it was assumed all patients would currently receive tocilizumab as the first choice of biologic therapy after csDMARDs (based on its availability as a licensed treatment option for sJIA since August 2011). After failure on tocilizumab, patients would switch to anakinra. Given the lack of head-to-head comparisons between bDMARDs in sJIA patients, both products were assumed to have the same efficacy.

Discontinuation rates were calibrated based on the probabilities of achieving CID, background mortality,

and the expectation that nearly all patients will have discontinued NSAIDs + corticosteroids and methotrexate after 6 and 16 weeks, respectively. To do this, a probability of discontinuation per week was estimated assuming 95% would discontinue in these time periods. For bDMARDs, data from NICE TA238 (tocilizumab for sJIA) were taken, in which 12.6% of patients were assumed to discontinue bDMARD treatment per year.<sup>30</sup> It was also assumed that some patients would continue treatment with bDMARDs with CID. In the base-case analysis, this was assumed to be 50%.

## Analysis Methodology

A micro-costing analysis was undertaken to estimate the total costs related to product acquisition and medical resource use (MRU). Product acquisition costs were based on estimated durations of treatment from ter Haar et al (first-line) and TA238 (later-line).<sup>21,30</sup> Unit costs were taken from Dutch reference cost sources.<sup>34</sup>

MRU was assumed to be related to whether or not patients still had CID. In the base-case analysis, four cost items were considered: outpatient rheumatology appointments, outpatient hematology appointments, general practitioner (GP) visits, and unplanned hospitalizations. Resource use frequencies were sourced from NICE TA238, with a revision to the expected number of outpatient rheumatology appointments per year to align with approximately one appointment per month.<sup>30</sup> A table of parameters included in the analysis is provided in [Table 1](#).

The analysis takes the hypothetical perspective of a European cohort of patients, given that no single location is expected to consider a blend of both treatment strategies. Drug and MRU costs are presented in Euros over a 5-year horizon, in line with standard budget impact analysis convention, which are then combined to calculate the overall 5-year costs associated with each treatment strategy. Sensitivity analyses including additional test costs, costs associated with MAS and long-term complications of treatment were considered to explore key areas of uncertainty.

## Results

### Headline Analysis Results

The results of the base-case analysis show that the total 5-year drug costs for the first-line anakinra strategy were €24,021, versus €20,471 for later-line anakinra; and the corresponding MRU costs were €19,197 (first-line) versus

€25,425 (later-line) ([Figure 1](#)). Therefore, the overall 5-year costs (drug acquisition and MRU) were lower for the first-line strategy (€43,218 versus €45,896), demonstrating that first-line anakinra is cost saving (-€2,677) relative to a later-line strategy. In other words, while first-line anakinra was associated with increased spend related to drug acquisition, it led to cost savings associated with reduced medical resource use expenditure including, for example, a reduction in hospital stays.

## Sensitivity Analyses

A number of sensitivity analyses were conducted to further explore the likely differences in the costs associated with each treatment strategy.

The MRU items included in the base-case analysis were selected on the basis of these costs representing the majority of direct healthcare costs incurred. However, additional costs may be incurred in a real-world setting. A sensitivity analysis was conducted to include the costs of C-reactive protein and creatinine clearance tests as well as public transport for outpatient consultations. By including both of these transport-related additional costs, the savings associated with first-line anakinra use increased further to -€2714.

Patients with active disease are subject to a risk of developing MAS, yet these costs were not captured in the base-case analysis. To explore the impact of including MAS-related costs in the analysis, we considered a sensitivity analysis wherein patients with active disease had an average of 0.0525 MAS events per year (based on a study by Grom et al<sup>36</sup>). The cost of resolving MAS was assumed to be equivalent to a 14-day stay in intensive care, including diagnostics and medication. Including MAS-related costs, the base-case cost savings increased to -€3225.

It is also understood that patients who receive prolonged treatment with corticosteroids and/or csDMARDs (such as methotrexate) may develop long-term complications related to treatment.<sup>37</sup> In a sensitivity analysis, we considered an additional cost of a hospitalization each year for 18% of patients (ie, the difference in patients not in remission at 5 years, see [Table 1](#)) treated with a later-line strategy after 5 years for an additional 5 years, as a crude exploration of the potential impact of including longer-term complications on results. By including this cost, the cost savings increased to -€3249.

**Table 1** Economic Analysis Input Parameters

Input	Value	Reference
<p>Clinical parameters (first-line anakinra)            These parameters are used to establish which proportion of patients have CAD versus CID over the time frame of the economic analysis, for patients managed with first-line anakinra.</p>		
CID at 1 year	76%	ter Haar et al. <sup>21</sup>
CID at 3 years	94%	
CID at 5 years	96%	
CID off treatment at 1 year	52%	
CID off treatment at 3 years	71%	
CID off treatment at 5 years	72%	
<p>Clinical parameters (later-line anakinra)            These parameters are used to establish which proportion of patients have CAD versus CID over the time frame of the economic analysis, for patients managed with later-line anakinra. A multitude of sources is used for this treatment strategy, owing to a lack of data available specifically reporting on the experience of sJIA patients managed with later-line anakinra following initial treatment with NSAIDs and csDMARDs.</p>		
Monocyclic course	25.5%	Grevich and Shenoi <sup>33</sup>
CID w/ corticosteroids (monocyclic)	30% (in 6 weeks)	Assumption
CID w/ methotrexate (monocyclic)	20% (in 24 weeks)	Assumption based on Nordström et al. <sup>17</sup>
CID w/ corticosteroids (chronic)	0%	Assumption
CID w/ methotrexate (chronic)	0%	Assumption
CID w/ bDMARDs	50% (in 24 weeks)	Assumption based on Nordström et al. <sup>17</sup>
Discontinuation for NSAIDs + corticosteroids	95% off by 6 weeks	Assumption
Discontinuation for csDMARD ± NSAIDs + corticosteroids	95% off by 16 weeks	Assumption
Discontinuation for bDMARDs	12.6% per year	NICE TA238 <sup>30</sup>
Proportion on bDMARDs with CID	50%	Assumption
CID at 5 years	78%	Calculation based on input parameters above
Difference in CID at 5 years	18%	
<p>Drug costs per week            These parameters are used to inform drug acquisition cost results. These costs refer to the estimated cost per week.</p>		
NSAIDs + corticosteroids	€ 0.82	Z-Index B.V. <sup>35</sup>
csDMARD	€ 18.88	
Anakinra	€ 225.40	
Tocilizumab	€ 275.68	
<p>Annual medical resource use            These parameters are used to estimate the medical resource utilization associated with patients that have CAD versus CID, noting that in general patients with CAD require additional resource.</p>		

(Continued)

**Table I** (Continued).

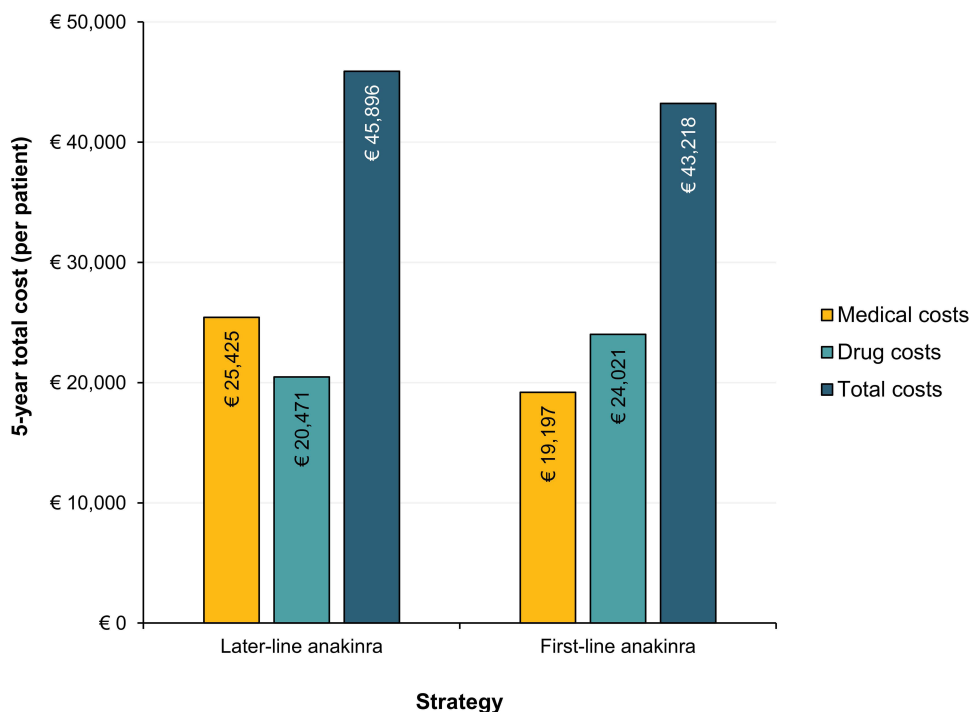
Input	Value	Reference
Rheumatology (CAD)	12.0	NICE TA238. <sup>30</sup> For rheumatology, increased from 10 reported in NICE TA238 to 12 reflecting an average of 1 appointment per month
Hematology (CAD)	12.0	
GP (CAD)	20.8	
Hospitalization (CAD)	22.1	
Rheumatology (CID)	4.0	Assumed 1 appointment every 3 months
GP (CID, on treatment)	3.5	NICE TA238 <sup>30</sup>
GP (CID, off treatment)	0.0	Assumption
Unit costs for medical resource use These parameters are used to assign costs based on the expected frequency of use (separated by CAD versus CID, as shown in the preceding rows).		
Rheumatology or hematology	€ 101.00	iMTA costing tool: <sup>34</sup> Outpatient visit, pediatrics
GP	€ 33.00	iMTA costing tool: <sup>34</sup> General practitioner, standard consultation
Hospitalization	€ 642.00	iMTA costing tool: <sup>34</sup> Nursing day, academic hospital

**Abbreviations:** bDMARD, biologic disease-modifying anti-rheumatic drug; csDMARD, conventional synthetic disease-modifying anti-rheumatic drug; CAD, clinically-active disease; CID, clinically-inactive disease; GP, General practitioner; iMTA, Institute for Medical Technology Assessment; NICE, National Institute for Health and Care Excellence; NSAID, non-steroidal anti-inflammatory drug; TA, technology appraisal.

## Discussion

This study demonstrates the economic implications of alternative treatment strategies (first-line versus later-line) with anakinra (in other words, either used in the

first-line setting, or in a refractory setting after failure to achieve CID with NSAID, corticosteroid, and csDMARD treatment [eg, methotrexate]). The results of the analysis demonstrate that first-line anakinra is



**Figure I** Base-case analysis results.

cost-saving relative to later-line use, which, when considered in conjunction with the improved clinical outcomes, gives further justification for the use of anakinra earlier in the course of sJIA.

The conclusion reached by the analysis presented may be surprising when considering the costs of biologic therapies in other rheumatic diseases. However, it is important to acknowledge that anakinra was first granted a marketing authorization in March 2002 (for its use in patients with rheumatoid arthritis), and has been used in clinical practice in the management of Still's disease for over a decade.<sup>20</sup> As such, the weekly cost of anakinra (€225, based on Dutch reference costs, last updated June 2020) is relatively low compared with other, more recent bDMARDs, such as canakinumab (Ilaris<sup>®</sup>, Novartis), which has an equivalent weekly cost of approximately €2750 (assuming one vial is used every 4 weeks, also last updated June 2020).

The clinical evidence in support of using anakinra earlier in the disease course is increasing over time. The SHARE guidelines advocate the use of anakinra as early as possible in sJIA to capitalize on the potential benefits of early IL-1 inhibition.<sup>11</sup> A recent study by Pardeo et al also demonstrated the probability of achieving CID with anakinra based on the disease duration from onset to start of anakinra – 91.9% of patients that started anakinra within 3 months of disease onset achieved CID off corticosteroids, versus 36.8% who started after 3 months of disease onset.<sup>25</sup> Accordingly, our study provides a timely assessment of the economic implications of earlier versus later use of anakinra.

In addition to anakinra, two other bDMARDs are licensed for the treatment of sJIA: tocilizumab (an IL-6 inhibitor) and canakinumab (an IL-1 $\beta$  inhibitor). However, unlike anakinra, both of these bDMARDs are only licensed for use after previous therapy with NSAIDs and corticosteroids, and tocilizumab is not licensed for the treatment of AOSD. Anakinra may be used within its licensed indication in patients with active systemic features of moderate to high disease activity without prior use of NSAIDs and corticosteroids. This distinction in the marketing authorization for bDMARDs is especially important in consideration of the emerging evidence for the role of targeted therapies in Still's disease, where anakinra can be started as soon as practically possible following diagnosis.

Economic analyses for sJIA treatments are limited in number, which is unsurprising given the rarity of sJIA. Our analysis makes use of some information reported in NICE TA238 (tocilizumab in sJIA).<sup>30</sup> In addition to the

previous NICE assessment of tocilizumab, Kittiratchakool et al recently undertook a cost-effectiveness analysis of tocilizumab in refractory sJIA.<sup>38</sup> This analysis only considers the potential use of tocilizumab in a relapsed setting, due to its availability in Thailand specifically in this setting. Conversely, our comparison considers the use of bDMARD therapy either in a refractory setting (ie, after NSAID, corticosteroid, and csDMARD treatment) or in a first-line setting – a comparison that (to our knowledge) has not been previously attempted.

We considered a synthesis of available information to compare the alternative treatment strategies. In doing so, we aimed to use the best available evidence to explore the differences in costs for each strategy, allowing for sensitivity analyses to be conducted concerning the key input parameters and associated assumptions. The ability to explore a range of sensitivity analyses for key settings and assumptions is an important advantage of our analytical approach, given that data available to quantify specific assumptions are either limited or in some cases not available (for example, the proportion of patients that achieve sustained CID with csDMARDs in an sJIA population).

Key uncertainties relating to lack of comparative data may be resolved with further evidence collection, though will unavoidably still be subject to various caveats related to differences in practices by geography and difficulties in enrolling patients within trials of treatments that are already routinely available. In particular, the first-line anakinra strategy considered in our economic analysis is based on data from a single-center study by ter Haar et al, though findings could have been different had the study been repeated in other settings or potentially within the context of a multi-center study. Therefore, while our study makes use of several assumptions to populate these missing data, further research and data collection are required to obtain a more robust estimate of the true costs associated with the management of sJIA.

In addition to the limited data available to inform our economic analysis, it is important to acknowledge that sJIA is a heterogeneous disease, and response to treatment may differ based on various patient characteristics.<sup>24,39,40</sup> These include time since diagnosis, disease subtype or severity, and treatment history.<sup>24,39–43</sup> Should additional data become available in the future, consideration of how outcomes compare across different patient subpopulations may be of interest from both a clinical and economic perspective.

The full benefits of first-line anakinra are not fully captured within our analysis, mainly due to the limitations of currently-available evidence. Benefits not captured within the main analysis include a reduced risk of developing MAS, the avoidance of long-run consequences relating to corticosteroid and/or csDMARD exposure, and productivity gains resulting from patients achieving CID. While we attempted to address these in part through sensitivity analysis, these potential benefits should also be considered when interpreting the results of our analysis. The findings presented within this study may be considered a conservative estimate of the potential cost savings associated with first-line anakinra.

## Conclusions

To our knowledge, this study presents the first economic comparison of treatment strategies in sJIA, based on current treatment pathways across Europe. There have been no published studies of medical management costs in Still's disease, based on whether patients have achieved CID and/or which treatment(s) they are currently managed with. The findings of the economic comparison illustrate the value of first-line anakinra in sJIA, when considered in the context of the full costs of patient management.

## Abbreviations

AOSD, Adult-onset Still's disease; bDMARD, Biologic disease-modifying anti-rheumatic drug; CAD, Clinically-active disease; CID, Clinically-inactive disease; csDMARD, Conventional-synthetic disease-modifying anti-rheumatic drug; GP, General practitioner; IL-1/IL-6, Interleukin-1/-6; JIA, Juvenile idiopathic arthritis; MAS, Macrophage activation syndrome; MRU, Medical resource use; NICE, National Institute for Health and Care Excellence; NSAID, Non-steroidal anti-inflammatory drug; NVK, Nederlandse Vereniging voor Kindergeneeskunde [Dutch Pediatrics Association]; SHARE, Single Hub and Access point for pediatric Rheumatology in Europe; sJIA, Systemic juvenile idiopathic arthritis.

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## Disclosure

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