

Paediatric rheumatology

On-demand corticosteroid use in the syndrome of undifferentiated recurrent fever: a literature review and results from the JIR-CLiPS survey study

E.D. Batu¹, S. Sener¹, M. Rodrigues², C. Vinit^{3,4}, F. Hofer⁵, K. Laskari⁶, R. Craveiro Costa⁷, M. Santos Faria⁸, G. Ozomay Baykal⁹, O. Boyarchuk¹⁰, O. Gilliaux¹¹, K. Pateras^{12,13}, H.E. Sonmez¹⁴, N. Toplak¹⁵, M. Gattorno¹⁶, M. Hofer¹⁷

¹Division of Paediatric Rheumatology, Department of Paediatrics, Hacettepe University Faculty of Medicine, Ankara, Turkey; ²Paediatric and Young Adult Rheumatology Unit, Centro Hospitalar Universitário São João, Porto Medical School, Porto, Portugal; ³Department of General Paediatrics, Jean Verdier Hospital, Bondy, France; ⁴Department of Paediatric Rheumatology, Robert Debre Hospital, Paris, France; ⁵Fondation RES, Lausanne, Switzerland; ⁶First Department of Propaedeutic and Internal Medicine, National and Kapodistrian University of Athens, Greece; ⁷Hospital Pediátrico, Unidade Local de Saúde de Coimbra, Coimbra, Portugal;

⁸Department of Rheumatology, Hospital Central do Funchal, SESARAM, Funchal, Portugal; ⁹Division of Paediatric Rheumatology, Department of Paediatrics, Umraniye Training and Research Hospital, Istanbul, Turkey; ¹⁰Department of Children's Diseases and Paediatric Surgery, I. Horbachevsky Ternopil National Medical University, Ternopil, Ukraine; ¹¹Department of Paediatrics, Clinique Notre Dame de Grâce, Gosselies, Belgium; ¹²Faculty of Public and One Health, University of Thessaly, Karditsa, Greece; ¹³Department of Statistics, Athens University of Economics and Business, Athens, Greece; ¹⁴Division of Paediatric Rheumatology, Department of Paediatrics, Kocaeli University Faculty of Medicine, Kocaeli, Turkey; ¹⁵Department of Allergology, Rheumatology and Clinical Immunology, UCH, UMC, Faculty of Medicine, University of Ljubljana, Slovenia; ¹⁶Department of Rheumatology and Autoinflammatory Diseases, IRCCS Istituto Giannina Gaslini, Genova, Italy; ¹⁷Department of Paediatrics, Hôpital Riviera-Chablais, Rennaz, Switzerland.

Abstract Objective

We aimed to analyse the strategies of physicians regarding corticosteroid use in the syndrome of undifferentiated recurrent fever (SURF) and examine the published data on this topic.

Methods

The JIR-CLiPS questionnaire which addresses physicians' practices about on-demand corticosteroid use in SURF was distributed via e-mail to potential respondents. We systematically reviewed the MEDLINE and Scopus databases and extracted the data about on-demand corticosteroid use in SURF.

Results

One hundred and thirty-seven physicians ($F/M=2.5$; 66.4% paediatric rheumatologists) from 45 countries responded to the survey. Around 70% of physicians prescribe corticosteroids for SURF flares. Most physicians (81.9%) do not use corticosteroids in SURF patients routinely, and this practice is more common among less experienced physicians ($p<0.001$). Prednisolone at a dose of 1 mg/kg (54.4%) was the most commonly preferred corticosteroid. The most common definition of response to corticosteroids was 'response within 12 hours' (51.6%). Most respondents (59.5%) consider changing treatment if corticosteroids cause a decrease in quality of life. We found 10 articles in the literature describing 239 SURF patients treated with on-demand corticosteroids. The most frequently preferred corticosteroid was prednisolone (63.8%). The response to corticosteroids was 70.8% and an increase in attack frequency was observed in almost 40% of patients.

Conclusion

On-demand corticosteroid use is not uncommon in the acute management of SURF attacks. However, most physicians do not use corticosteroids routinely and there is no consensus regarding the definition of response to treatment and when to change treatment neither in our survey results nor in the literature.

Key words

autoinflammatory disease, corticosteroid, steroid, syndrome of undifferentiated recurrent fever, treatment

Ezgi D. Batu, MD
 Seher Sener, MD
 Mariana Rodrigues, MD
 Caroline Vinit, MD
 François Hofer, MS
 Katerina Laskari, MD
 Ricardo Craveiro Costa, MD
 Margarida Santos Faria, MD
 Gulcan Ozomay Baykal, MD
 Oksana Boyarchuk, MD
 Olivier Gilliaux, MD
 Konstantinos Pateras, MD
 Hafize E. Sonmez, MD
 Natasa Toplak, MD
 Marco Gattorno, MD
 Michaël Hofer, MD

Please address correspondence to:

Ezgi Deniz Batu
 Division of Rheumatology,
 Department of Paediatrics,
 Hacettepe University Faculty of
 Medicine,
 Ankara 06100, Turkey.
 E-mail: egzidenizbatu@yahoo.com
 ORCID: 0000-0003-1065-2363

Received on November 5, 2024; accepted
 in revised form on March 5, 2025.

© Copyright CLINICAL AND
 EXPERIMENTAL RHEUMATOLOGY 2025.

Funding: this article is based upon work from COST Action CA21168-Improving outcome of Juvenile Inflammatory Rheumatism via universally applicable clinical practice strategies (JIR-CLIPS), supported by COST (European Cooperation in Science and Technology) <http://www.cost.eu>

Competing interests: M. Gattorno has received honoraria and speaker's bureau fees from Novartis, Sobi and Fresenius-Kabi.

The other authors have declared no competing interests.

Introduction

Systemic autoinflammatory diseases (SAIDs) are characterised by inflammation driven by innate immunity dysregulation (1). The most common monogenic SAIDs are familial Mediterranean fever (FMF), hyperimmunoglobulin D syndrome/mevalonate kinase deficiency (HIDS/MKD), cryopyrin associated periodic syndrome (CAPS), and tumour necrosis factor receptor associated periodic syndrome (TRAPS) (2, 3). On the other hand, periodic fever, aphthous stomatitis, pharyngitis, and adenopathy (PFAPA) syndrome is the most frequent multifactorial SAID especially among children (4).

Despite great advances in genetic analysis during the last decades, a monogenic aetiology cannot be identified in almost two-thirds of SAIDs (3). While chronic inflammation predominates the phenotype in some undifferentiated SAIDs (USAIDs), recurrent fever episodes are the main common feature for others. The subgroup of USAIDs characterised with febrile flares of inflammation is called the syndrome of undifferentiated recurrent fever (SURF) (5). SURF patients lack specific features of PFAPA syndrome and pathogenic variants or VUS on periodic fever genes (5). There are no widely accepted treatment recommendations for SURF patients.

On-demand corticosteroids form the mainstay of acute treatment in PFAPA episodes (4). In around 85–90% of PFAPA patients, single dose corticosteroids lead to an abrupt cessation of fever (4). On-demand corticosteroids could also work for some patients with HIDS/MKD (6). Although the response rate does not seem to be as high as that observed in PFAPA patients, on-demand corticosteroids may be beneficial during attacks in SURF patients. However, no previous study has focused on on-demand corticosteroid use in SURF patients.

Our aim was to evaluate the practices of physicians worldwide regarding on-demand corticosteroid use in SURF and to analyse the published data on this topic.

Materials and methods

Questionnaire about on-demand corticosteroid use in SURF

This study is an international, online,

cross-sectional survey study included in the JIR-CLIPS project. The general aim of this project is to analyse the real-life Clinical Practice Strategies (CLIPS) in five conditions: paediatric vasculitis (Kawasaki disease and immunoglobulin A vasculitis), paediatric lupus nephritis, and three autoinflammatory diseases as systemic juvenile idiopathic arthritis/adult-onset Still's disease, biological treatment in monogenic autoinflammatory diseases, and PFAPA/SURF. The SURF questionnaire was developed by the JIR-CLIPS PFAPA/SURF team, and it includes a total of 42 questions. Addressing the objectives of this study, we analysed the responses to nine questions of the survey that focused on corticosteroid use in SURF, in addition to the seven questions about the demographic features of the respondents (Supplementary Table S1). We evaluated the responses between January 27th and May 31st, 2024, but the survey is still accessible to new respondents. It is noteworthy that each respondent could fill the survey only once.

Data collection was in accordance with the General Data Protection Regulation (GDPR) law. And, ethical approval was not required for this study.

Systematic literature review

Two authors (EDB and SS) systematically searched the MEDLINE and Scopus databases from their inception until June 16th, 2024, according to the PRISMA guidelines (7). The following keywords were used during the literature search: syndrome of undifferentiated recurrent fever, SURF, undifferentiated systemic autoinflammatory disease, USAID, steroid, corticosteroid, glucocorticoid, prednisone, prednisolone, methylprednisolone, betamethasone, dexamethasone, deflazacort, and hydrocortisone. We analysed only English articles and hand-searched the references of the included articles. The studies that included data regarding on-demand corticosteroid use in SURF were included. Figure 1 shows the flowchart of the systematic literature review.

The following data were extracted from the included articles: number of patients, demographic and clinical features, type of corticosteroid, dose, the

number of doses per flare, treatment duration, response, and adverse events.

Statistical analysis

The descriptive analysis of the responses was performed using SPSS v. 25.0 (IBM, Armonk, NY, USA). Ratios and percentages were used to present categorical variables and continuous data were described in median and minimum-maximum values. The Chi-square test or Fisher's exact test were used to compare categorical variables, as appropriate. A *p*-value below 0.05 was regarded as statistically significant.

Results

Results of the JIR-CliPS survey

One hundred and thirty-seven of 298 physicians who responded the PFAPA/SURF survey answered questions regarding on-demand corticosteroid use in SURF (Table I). They were from 45 different countries (Supplementary Fig. S1) and 71.3% were female. The majority (85.4%) provided care for pediatric patients only. Most respondents were paediatric rheumatologists (66.4%), and 56.9% of them had ≥ 10 -year experience with patients who have recurrent fever (Table I).

Ninety-seven participants (70.8%) prescribe corticosteroids at the onset of a SURF flare. Prednisolone was the most commonly used corticosteroid (62.8%), followed by prednisone (52.6%). The most frequently chosen prednisone equivalent dose was 1 mg/kg (54.4%), followed by 0.67 mg/kg (10.4%).

Most physicians (81.9%) state that they do not use corticosteroids in SURF patients routinely (Table II). Among them, the most common factor influencing the decision to use corticosteroids during a flare was the severity of the attack (54.1%). The frequency of physicians using on-demand corticosteroids routinely in SURF management was higher among those with ≥ 10 -year experience compared to less experienced ones (24% vs. 10.3%; *p*<0.001) (Table II).

Most of the respondents prefer 1 or 2 doses of corticosteroids, depending on the response (78.7%) (Table II). The percentage of physicians preferring 1 or 2 doses over single dose was higher among those with ≥ 10 years of experi-

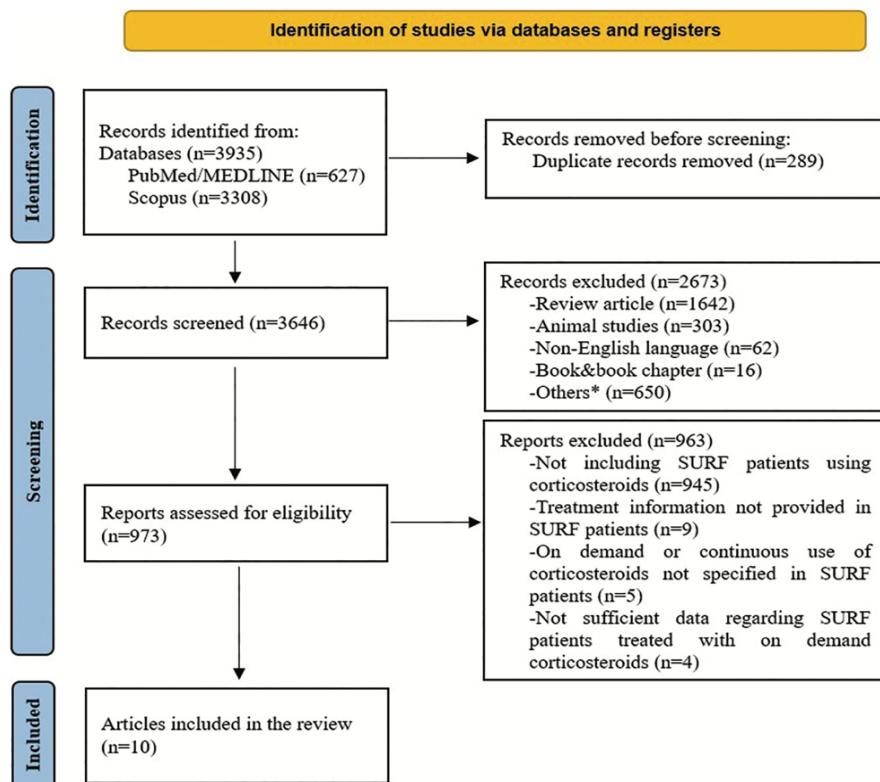


Fig. 1. The PRISMA flow diagram of literature screening regarding syndrome of undifferentiated recurrent fever (SURF) patients using on-demand corticosteroids.

*Letter, editorial, conference paper, guideline, short survey, note, commentary, poster.
SURF: syndrome of undifferentiated recurrent fever.

Table I. General characteristics of the participants who responded to the questions of the JIR-CliPS survey regarding corticosteroid use in SURF (n=137).

Demographic features	n (%)
Sex, female	97/136 (71.3)
Country	
Turkey	29 (21.2)
France	21 (15.3)
Brazil	12 (8.8)
Germany	9 (6.6)
United Kingdom	6 (4.4)
Other countries*	60 (43.8)
Institution type	
University hospital	88 (64.2)
Tertiary hospital	21 (15.3)
Hospital	21 (15.3)
Private practice	5 (3.6)
Others*	2 (1.5)
Taking care of inpatients or outpatients	
Both outpatients and inpatients	130 (94.9)
Only outpatients	7 (5.1)
Taking care of paediatric or adult patients	
Only children	117 (85.4)
Both children and adults	14 (10.2)
Only adults	6 (4.4)
Specialty	
Paediatric rheumatology	91 (66.4)
Paediatric rheumatology and immunology	14 (10.2)
Paediatrician	7 (5.1)
Others*	25 (18.2)
Experience in the care of patients with recurrent fever	
≥ 10 years	78 (56.9)
5-9 years	40 (29.2)
0-4 years	19 (13.9)

SURF: syndrome of undifferentiated recurrent fever.

*less than five respondents per individual country.

Table II. The responses to the survey questions regarding on-demand corticosteroid use in syndrome of undifferentiated recurrent fever (SURF).

Parameters, n (%)	All responders (n=137)	Responders with <10-year experience (n=59)	Responders with ≥10-year experience (n=78)	p-value*
How do you prescribe a treatment with a steroid dose at flare-onset?				
Routinely	24/133 (18.1)	6/58 (10.3)	18/75 (24)	0.042
Not routinely	109/133 (81.9)	52/58 (89.7)	57/75 (76)	0.042
Depending on attack severity	72/133 (54.1)	28/58 (48.3)	44/75 (58.7)	0.098
When antipyretics are not enough to control fever	41/133 (30.8)	17/58 (29.3)	22/75 (33.3)	0.435
Depending on attack frequency	38/133 (28.6)	16/58 (27.6)	22/75 (33.3)	0.357
Depending on family preferences	28/133 (21.1)	11/58 (18.9)	17/75 (22.7)	0.309
Only once, to confirm diagnosis	23/133 (17.3)	13/58 (22.4)	9/75 (12)	0.074
Others	11/133 (8.3)	5/58 (8.6)	6/75 (8)	0.981
How many doses per flare?				
1 or 2, depending on the response	100/127 (78.7)	36/54 (66.7)	64/73 (87.7)	0.015
Only 1 dose	27/127 (21.3)	18/54 (33.3)	9/73 (12.3)	0.015
Do you use any forms of steroids that are suitable for young children (e.g. suspension, drops, etc.)?				
Yes	101 (73.7)	32 (54.2)	69 (88.5)	<0.001
No	36 (26.3)	27 (45.8)	9 (11.5)	<0.001
What is the maximum steroid uses that you recommend per year?				
<5	50/136 (36.8)	27 (46.6)	23/77 (29.9)	0.217
5 to 10	66/136 (48.5)	25 (43.1)	41/77 (53.2)	0.138
>10	20/136 (14.7)	7 (12.1)	13/77 (16.9)	0.243
When do you consider that the patients responded to a steroid dose at flare-onset?				
Response within 3-4 hours	17/128 (13.3)	9/55 (16.4)	8/73 (10.9)	0.341
Response within 12 hours	66/128 (51.6)	23/55 (41.8)	43/73 (58.9)	0.103
Response within 24 hours	45/128 (35.1)	23/55 (41.8)	22/73 (30.1)	0.297
On which criteria would you consider that the patient did not respond?				
Need for more than 2 doses of steroid per episode	69/133 (51.9)	27/57 (47.4)	42/75 (56)	0.194
No improvement in fever within 24 hours	68/133 (51.1)	31/57 (54.4)	37/75 (49.3)	0.562
Improvement in fever but fever recurs within the same episode after the steroid dose	49/133 (36.8)	26/57 (45.6)	23/75 (31.9)	0.081
No improvement in fever within 12 hours	26/133 (19.5)	11/57 (19.3)	15/75 (20)	0.835
Need for more than 1 dose of steroid per episode	11/133 (8.3)	5/57 (8.8)	6/75 (8)	0.649
No improvement in fever within 3-4 hours	9/133 (6.8)	5/57 (8.8)	4/75 (5.3)	0.672
Others	4/133 (3.1)	1/57 (1.5)	3/75 (4)	0.153
If steroids at flare-onset decrease the intervals between the flares, based on which criteria would you consider another treatment?				
If decreased the quality of life	78/131 (59.5)	33/57 (57.9)	45/74 (60.8)	0.716
Flare interval shorter than 2 weeks	47/131 (35.9)	21/57 (36.8)	26/74 (35.1)	0.902
Decrease in attack intervals persisting for >3 months	40/131 (30.5)	19/57 (33.3)	21/74 (28.4)	0.357
Only if severe attacks	33/131 (25.2)	16/57 (28.1)	17/74 (22.9)	0.741
Flare interval shorter than 3 weeks	29/131 (22.1)	9/57 (15.8)	21/74 (28.4)	0.069
Flare interval shorter than 4 weeks	29/131 (22.1)	17/57 (29.8)	13/74 (17.6)	0.094
Decrease in attack intervals persisting for >6 months	27/131 (20.6)	11/57 (19.3)	16/74 (21.6)	0.643
Decrease in attack intervals persisting for >1 year	5/131 (3.8)	2/57 (3.5)	3/74 (4)	0.947
Others	7/131 (5.3)	2/57 (3.5)	5/74 (6.8)	0.183

*p-values are for the comparison between respondents with <10-year and ≥10-year experience.

ence than less-experienced respondents (87.7% vs. 66.7%; $p=0.015$) (Table II). The most common maximum number of corticosteroid doses per year was 5 to 10 (48.5%).

‘Response within 12 hours’ (51.6%) was the most frequent definition for response to corticosteroid dose at flare

onset (Table II). Around half of the physicians agreed that the patient was not responding to corticosteroids if the patient needed >2 doses of corticosteroids per flare (51.9%) or there was no improvement in fever within 24 hours (51.1%). When corticosteroid use leads to an increase in flare frequency, a de-

crease in patient’s quality of life is the most common reason for preferring a different treatment (59.5%).

Results of the literature review

We reviewed the literature and identified 10 articles containing 239 SURF patients treated with on-demand corti-

Table III. General characteristics of patients with syndrome of undifferentiated recurrent fever (SURF) treated with on-demand corticosteroids in the literature.

Characteristics	n (%) or median (min-max)
Total number of patients	498
Number of patients treated with CS	239
Age at diagnosis, years	3.7 (0-35)
Sex, female	12/32 (37.5)
Clinical features	
Fever	32/32 (100)
Abdominal pain	18/32 (56.3)
Headache	14/32 (43.8)
Nausea/vomiting	11/32 (34.4)
Pharyngitis/tonsillitis	10/32 (31.3)
Cervical lymphadenopathy	8/32 (25.8)
Aphthous stomatitis	8/32 (25.8)
Arthralgia	7/32 (21.9)
Ocular symptoms	6/32 (18.8)
Rash	6/32 (18.8)
Myalgia	3/32 (9.4)
Constitutional symptoms	3/32 (9.4)
Duration of febrile episode, days	4.3 (2-30)
Interval between febrile episodes, weeks	7.8 (2-25.7)
Type of CS	
Prednisolone	30/47 (63.8)
Betamethasone	16/47 (34.1)
Methylprednisolone	1/47 (2.1)
Doses of CS*	
0.5 mg/kg	1/81 (1.2)
0.5-1 mg/kg	46/81 (56.8)
≤1 mg/kg	29/81 (35.8)
>1 mg/kg	5/81 (6.2)
Response to CS in a SURF episode	
Improvement	136/192 (70.8)
No improvement	56/192 (29.2)
Concomitant therapy with CS	
Colchicine	15/25 (60)

CS: corticosteroid; SURF: syndrome of undifferentiated recurrent fever.

*prednisone equivalent.

costeroids (Fig. 1) (5, 8-16). The definitions of SURF and the inclusion/exclusion criteria for SURF patients are presented in Supplementary Table S2, while Supplementary Table S3 details these patients' characteristics.

Age and sex were reported in only four studies (11-14); the median (min.-max.) age of the patients was 3.7 (0-35) years and 37.5% of the patients were female (Table III). Among the patients whose clinical features were specified, all had fever (100%), and other common symptoms included abdominal pain (56.3%), headache (43.8%), nausea/vomiting (34.4%) and pharyngitis/tonsillitis (31.3%) (11-14).

In the five articles where corticosteroid type and dose were reported, the most

commonly used corticosteroid type was prednisolone (63.8%), while the most commonly preferred corticosteroid dose was 0.5–1 mg/kg prednisone equivalent (56.8%) (8, 10, 11, 13, 14). The number of corticosteroid doses per SURF flare was reported as 1 dose in only one article (13). The definition of response to corticosteroids during a SURF flare was given in only one article as 'resolution of symptoms with a single dose of steroid' (13). The corticosteroid response rate which was reported in seven articles was 70.8% (5, 8, 9, 12-15). An increase in attack frequency after on-demand corticosteroid use was mentioned only in one case reported by Harrison *et al.* (11) and in eight of 21 cases (38.1%) re-

ported by De Pauli *et al.* (8). Corticosteroid side effects were not addressed in any of the articles except one where the authors stated that no side effects were observed in a SURF patient using on-demand corticosteroids (11). Colchicine (60%) was the most common concomitant therapy (8, 11, 12, 14).

Discussion

To our knowledge, this is the first study in the literature that focus on on-demand corticosteroid use in SURF. Our results show that on-demand corticosteroid use is not uncommon in SURF management. However, most physicians do not use corticosteroids routinely and the most important factor affecting this treatment decision is the attack severity. On the other hand, quality of life of the patients is a significant consideration for physicians while deciding to switch to another treatment. The presented literature review revealed a response rate of 70.8% for on-demand corticosteroids in SURF.

SURF is defined as a subgroup of US-AIDs primarily characterised by recurrent fever episodes, while USAIDs include systemic inflammation that may affect multiple organ systems (5). Both are diagnosed by exclusion of other SAIDs, meaning other causes of recurring fever or inflammation must be ruled out before reaching a diagnosis (17). They both lack specific genetic or clinical markers, making classifying them under established autoinflammatory diseases challenging (17). Their management focuses on controlling inflammation and symptoms, treatment plans are often personalised.

There are no clear recommendations regarding SURF management since its aetiopathogenesis remains unknown. On-demand corticosteroids, colchicine, and biologic drugs (mainly anti-interleukin 1 agents) are among the main therapeutic options (5). Our results show that more than two-thirds of physicians use on-demand corticosteroids in SURF treatment. Previous studies showed that colchicine was one of the most frequently used drugs in SURF treatment with a complete response rate higher than 50% (5, 18). While colchicine is mainly used for prevent-

ing attacks, on-demand corticosteroids are a more acute treatment which aims to abort inflammatory flares. We know that successive corticosteroid use may cause an increase in attack frequency in PFAPA syndrome (4, 19). Whether the same impact is present in SURF patients remains to be elucidated. There are a few reports of increased attack frequency after on-demand corticosteroid use in SURF (8, 11). However, long term data is not present.

The corticosteroid response rate derived from the literature review was 70.8% in our study. In PFAPA syndrome, on the other hand, a response rate around 85–95% is observed (4, 19). Although it is challenging to make a direct comparison between these rates, inconsistent response to on-demand corticosteroids could be counted among the features differentiating SURF from PFAPA syndrome. PFAPA represents a more homogeneous phenotype with specific clinical features compared to SURF. SURF is a less well-defined and heterogeneous entity which may involve more complex or diverse immune mechanisms that do not respond to corticosteroids. Also, although a specific monogenic cause has not been identified in SURF patients, there could still be an overlap with monogenic SAIDs where corticosteroids are less effective than they are in PFAPA syndrome. Previous studies showed that more than half of SURF patients respond well to colchicine (5, 18). SURF patients with a good colchicine response may represent a more homogeneous subgroup of SURF. For instance, generalised lymphadenopathy was less frequently observed among colchicine-responsive SURF patients (18).

The main limitation of this study is inherent in the bias introduced by the characteristics of the respondents. Since the majority of the respondents were paediatric rheumatologists and physicians who take care of only children, the perspectives of general practitioners, paediatricians, and adult care specialists are not equivalently represented. However, paediatric rheumatology is currently the main subspecialty that focus on the care of SURF patients. Therefore, analysing paediatric rheu-

matologists' perspectives is valuable. Regarding literature review, possible overlap of cohorts from same center, underrepresentation of adult patients, the lack of specification of on-demand or continuous corticosteroid use, and the absence of a widely accepted definition for SURF were the main limitations. Also, some of the studies included from the literature review are from papers published before the definition of SURF in the literature. Furthermore, while most included papers defined SURF patients in similar terms, there is no widely accepted standard for determining the 'extent' of genetic testing required before classifying patients as having SURF. Thus, SURF still represents a heterogeneous group of disorders which makes it difficult to draw strict conclusions based on the literature data.

In conclusion, the results of this study may provide some guidance for the physicians taking care of SURF patients, in the absence of clear recommendations. The presented data may also serve as a reference to be communicated to the families while prescribing on-demand corticosteroids. Although the response rate is not as high as that observed in PFAPA syndrome, on-demand corticosteroids seem to work for more than two-thirds of SURF patients. Prospective studies with long-term follow-up will be invaluable and can shed more light to the benefits and risks of on-demand corticosteroid use in SURF treatment.

References

1. PARK H, BOURLA AB, KASTNER DL, COLBERT RA, SIEGEL RM: Lighting the fires within: the cell biology of autoinflammatory diseases. *Nat Rev Immunol* 2012; 12(8): 570-80. <https://doi.org/10.1038/nri3261>
2. BATU ED, BASARAN O, BILGINER Y, OZEN S: Familial Mediterranean fever: How to interpret genetic results? How to treat? A quarter of a century after the association with the MEFV gene. *Curr Rheumatol Rep* 2022; 24(6): 206-12. <https://doi.org/10.1007/s11926-022-01073-7>
3. DI DONATO G, D'ANGELO DM, BREDA L, CHIARELLI F: Monogenic autoinflammatory diseases: state of the art and future perspectives. *Int J Mol Sci* 2021; 22(12): 6360. <https://doi.org/10.3390/ijms22126360>
4. BATU ED: Periodic fever, aphthous stomatitis, pharyngitis, and cervical adenitis (PFAPA) syndrome: main features and an algorithm for clinical practice. *Rheumatol Int* 2019; 39(6): 957-70. <https://doi.org/10.1007/s00296-019-04257-0>
5. PAPA R, RUSMINI M, VOLPI S et al.: Next generation sequencing panel in undifferentiated autoinflammatory diseases identifies patients with colchicine-responder recurrent fevers. *Rheumatology (Oxford)* 2020; 59(2): 344-60. <https://doi.org/10.1093/rheumatology/kez270>
6. TER HAAR NM, JEYARATNAM J, LACHMANN HJ et al.: The phenotype and genotype of mevalonate kinase deficiency: a series of 114 cases from the Eurofever Registry. *Arthritis Rheumatol* 2016; 68(11): 2795-805. <https://doi.org/10.1002/art.39763>
7. PAGE MJ, MCKENZIE JE, BOSSUYT PM et al.: The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. *BMJ* 2021; 372: n71. <https://doi.org/10.1136/bmj.n71>
8. DE PAULI S, LEGA S, PASTORE S et al.: Neither hereditary periodic fever nor periodic fever, aphthae, pharyngitis, adenitis: Undifferentiated periodic fever in a tertiary pediatric center. *World J Clin Pediatr* 2018; 7(1): 49-55. <https://doi.org/10.5409/wjcp.v7.i1.49>
9. GERRITSMA AM, SUTERA D, CANTARINI L et al.: TNFRSF1A-pR92Q variant identifies a subset of patients more similar to systemic undifferentiated recurrent fever than TNF receptor-associated periodic syndrome. *Clin Exp Rheumatol* 2023; 41(10): 1998-2007. <https://doi.org/10.55563/clinexprheumatol/am4phc>
10. GÓMEZ-CAVERZASCHI V, YAGÜE J, ESPINOSA G et al.: Disease phenotypes in adult patients with suspected undifferentiated autoinflammatory diseases and PFAPA syndrome: Clinical and therapeutic implications. *Autoimmun Rev* 2024; 23(7-8): 103520. <https://doi.org/10.1016/j.autrev.2024.103520>
11. HARRISON SR, MCGONAGLE D, NIZAM S et al.: Anakinra as a diagnostic challenge and treatment option for systemic autoinflammatory disorders of undefined etiology. *JCI Insight* 2016; 1(6): e86336. <https://doi.org/10.1172/jci.insight.86336>
12. KOSUKCU C, TASKIRAN EZ, BATU ED et al.: Whole exome sequencing in unclassified autoinflammatory diseases: more monogenic diseases in the pipeline? *Rheumatology (Oxford)* 2021; 60(2): 607-16. <https://doi.org/10.1093/rheumatology/keaa165>
13. LUU I, NATION J, PAGE N et al.: Undifferentiated recurrent fevers in pediatrics are clinically distinct from PFAPA syndrome but retain an IL-1 signature. *Clin Immunol* 2021; 226: 108697. <https://doi.org/10.1016/j.clim.2021.108697>
14. MARQUES ML, CUNHA IM, ALVES S, GUEDES M, ZILHÃO C: Systemic autoinflammatory diseases in pediatric population. *Asia Pac Allergy* 2022; 12(3): e29. <https://doi.org/10.5415/apallergy.2022.12.e29>
15. TER HAAR NM, EIJKELBOOM C, CANTARINI L et al.: Clinical characteristics and genetic analyses of 187 patients with undefined autoinflammatory diseases. *Ann Rheum Dis* 2019; 78(10): 1405-11. <https://doi.org/10.1136/annrheumdis-2018-214472>
16. VITALE A, CAGGIANO V, SILVA I et al.: Axial spondyloarthritis in patients with recurrent

fever attacks: data from the AIDA network registry for undifferentiated autoinflammatory diseases (USAIDs). *Front Med (Lausanne)* 2023; 10: 1195995. <https://doi.org/10.3389/fmed.2023.1195995>

17. VYZHGA Y, WITTKOWSKI H, HENTGEN V *et al.*: Unravelling the clinical heterogeneity of undefined recurrent fever over time in the European registries on Autoinflammation. *Pediatr Rheumatol Online J* 2024; 22(1): 55. <https://doi.org/10.1186/s12969-024-00987-z>

18. SUTERA D, BUSTAFFA M, PAPA R *et al.*: Clinical characterization, long-term follow-up, and response to treatment of patients with syndrome of undifferentiated recurrent fever (SURF). *Semin Arthritis Rheum* 2022; 55: 152024. <https://doi.org/10.1016/j.semarthrit.2022.152024>

19. WANG A, MANTHIRAM K, DEDEOGLU F, LICAMELI GR: Periodic fever, aphthous stomatitis, pharyngitis, and adenitis (PFAPA) syndrome: A review. *World J Otorhinolaryngol Head Neck Surg* 2021; 7(3): 166-73. <https://doi.org/10.1016/j.wjorl.2021.05.004>