Multi-inflammatory syndrome and Kawasaki disease in children during the COVID-19 pandemic: A nationwide register-based study and time series analysis

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Abstract
Aim: We investigated whether the ongoing COVID-19 pandemic was associated with the occurrence of Kawasaki disease or with multi-inflammatory syndrome in children (MIS-C).
Methods: This national Finnish register-based study was based on laboratory-confirmed severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infections, MIS-C and Kawasaki disease cases. We performed a time series analysis on the occurrence of Kawasaki disease in 2016–2020.
Results: In 2020, there were 5170 laboratory-confirmed COVID-19 cases in children under 18 years of age and five fulfilled the MIS-C case definition. The occurrence of MIS-C was 0.97 per 1000 (95% confidence interval: 0.31-2.26) laboratory-confirmed SARS-CoV-2 infections in children. Our time series analysis showed that Kawasaki disease cases decreased during the COVID-19 pandemic. The seasonally adjusted incidence rate ratio was 0.49 (95% confidence interval: 0.32-0.74) when it was compared to pre-pandemic levels. This coincided with a reduced occurrence of respiratory infections, due to social distancing in the population.
Conclusion: This nationwide register-based study found that MIS-C was a rare complication of the SARS-CoV-2 infection. The occurrence of Kawasaki disease and respiratory infections decreased during the pandemic. This suggests that transmissible microbes may play an important role in Kawasaki disease and social distancing may have a protective effect.

KEYWORDS
epidemiology, Kawasaki disease, multi-inflammatory syndrome in children, social distancing, transmissible microbes

Abbreviations: MIS-C, multisystem inflammatory syndrome in children; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2.
INTRODUCTION

Since the global emergence of COVID-19, several centres have reported clusters of critically ill children with an inflammatory syndrome that appeared similar to Kawasaki disease. This is now commonly known as multisystem inflammatory syndrome in children (MIS-C).\(^1\)\(^2\) Surveillance data of childhood COVID-19 cases have indicated that children have lower rates of hospitalisation than adults. However, MIS-C has emerged as a novel clinical entity with a broad clinical spectrum, including life-threatening cardiogenic shock.\(^3\)\(^4\)

Several studies have reported typical symptoms of MIS-C in patient clusters.\(^1\)\(^5\)\(^-\)\(^9\) However, there have been limited epidemiological data about the estimated proportion of children who develop MIS-C after mild, but symptomatic, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infections. Furthermore, the overall incidence rate ratios of Kawasaki disease or Kawasaki disease-like diagnoses have rarely been compared before and during the COVID-19 pandemic using a time series analysis.\(^10\)\(^11\)

The aim of this register-based, nationwide study was to report the occurrence of MIS-C during the COVID-19 pandemic in 2020. Furthermore, we wanted to compare the relative changes in the occurrence of Kawasaki disease before and during the COVID-19 pandemic using a time series analysis.

PATIENTS AND METHODS

Study design

We conducted a register-based, nationwide study of the epidemiology of MIS-C in Finland in 2020 and of Kawasaki disease in 2016-2020, before and during the COVID-19 pandemic. Since August 2020, Finland has strongly recommended free SARS-CoV-2 polymerase chain reaction testing for all children with symptoms that suggest COVID-19, even if they are mild. On 16 March 2020, the Finnish Government announced a state of emergency, which included temporarily closing restaurants, food and beverage services and primary and secondary schools until 13 May. Preschools and day care centres were not closed. After the two-month closure period, schools remained mostly open for the rest of 2020.

The study was covered by the institutional research permit of the Finnish Institute for Health and Welfare, who performed the analyses. According to national legislation, informed consent was not required for this study, because it was performed with register-based data and no patients were contacted. Other authors used and analysed the summarised register-based data.

National registers and data retrieval

We used the Finnish Care Register for Health Care to identify children under 18 years of age with MIS-C from 1 January to 31 December 2020 and children with Kawasaki disease from 1 January 2016 to 31 December 2020. Finland’s Care Register for Health Care, maintained by the Finnish Institute for Health and Welfare, includes data on the diagnoses of patients who were discharged from hospitals or who visited outpatient clinics or family doctors at primary healthcare centres. We retrieved data for all patients with MIS-C aged 0–18 using the International Classification of Diseases, Tenth Revision (ICD-10) diagnosis code M35.8 and for patients with COVID-19 using the ICD-10 code U07.2. Information on patients with Kawasaki disease was retrieved using the ICD-10 code M30.3. We contacted all five university hospitals in Finland with paediatric intensive care units to double-check the MIS-C cases during the study period. This confirmed that all the cases had been recorded in the registers. We calculated the annual population-based incidence rate of Kawasaki disease from 2016 to 2020 using official government population data from Statistics Finland. The number of children with a laboratory-confirmed SARS-CoV-2 infection was retrieved from the National Infectious Diseases Register. All clinical microbiology laboratories in Finland are required to report all positive cases of SARS-CoV-2 to this Register. We also used the Register to demonstrate the impact of social distancing on the occurrence of other infections during the COVID-19 pandemic. To do this, we retrieved the numbers of influenza A, respiratory syncytial virus and Mycoplasma pneumoniae-positive respiratory samples, as well as positive Streptococcus pneumoniae blood cultures. Anonymised, summarised data from this Register are freely available to the public in Finland. The number of other respiratory infections was present in children and adolescents aged 0–19 years, because of the predefined age structure of the Register’s public database.

Case definition

We defined MIS-C as follows: the patient had a registry entry in the Finnish Care Register for Health Care for MIS-C or fulfilled the case definition of MIS-C provided by The American Centers for Disease Control and Prevention.\(^12\) They had also been discharged from, or died, in any of the five Finnish university hospitals with paediatric
intensive care wards. Kawasaki disease was defined as a patient with an entry for Kawasaki disease in the same Finnish Register. The child’s first visit for Kawasaki disease was identified, and subsequent, or duplicate, visits were excluded.

2.4 | Statistical analysis

We calculated the annual incidence rates, and 95% confidence intervals (95% CIs), for both Kawasaki disease and laboratory-confirmed SARS-CoV-2 cases by dividing the total numbers by children aged 0-18 in Finland in 2020, according to Statistics Finland. The MIS-C incidence rates, and 95% CIs, were calculated per 1000 laboratory-confirmed SARS-CoV-2 cases for the three-week periods from 1 January 2020 to 31 December 2020. This enabled us to examine the impact of the COVID-19 epidemic on the occurrence of Kawasaki disease.

We also performed an interrupted time series, which is a quasi-experimental design that evaluates changes on longitudinal data in relation to an intervention point. In this case, the intervention point was the start of the pandemic, including the need for social distancing. The outcome was the quarterly number of Kawasaki disease cases each year from 2016 to 2020. We chose the date 16 March as the cut-off point for our pre-pandemic and pandemic data, as that is when Finland declared a state of emergency. Our time series included an intercept time variable, which was 20 three-month periods, and a period variable. Seasonality was adjusted using a Fourier term, and over-dispersion of data was controlled by scaled parameters with Pearson’s chi-square test. The goodness of fit of model was evaluated by the Hosmer-Lemeshow chi-square test (p=0.448). Autocorrelation was graphically examined, but it was not detected after the data were adjusted for seasonality.

We used the most feasible program for each statistical analysis, because all the statistical analyses were not available in one program. The differences between the two proportions were tested with a standard normal deviate test available in StatsDirect statistical software 3 (StatsDirect Ltd). The data management and calculation of basic statistics were conducted using SPSS, version 27 (IBM Corp), and the 95% CI for rates and proportion differences were calculated using StatsDirect statistical software 3 (StatsDirect Ltd). ITS analysis was performed using Stata 16 (StataCorp LLC), and the incidence rates were created using OriginPro 2020 (OriginLab Corporation).

3 | RESULTS

From 1 January 2020 to 31 December 2020, there were 5,170 laboratory-confirmed SARS-CoV-2 infections in children under 18 years of age (Figure 1A). The first confirmed paediatric cases were reported on 24 February 2020. In Finland, schools were closed from 16 March 2020 to 13 May 2020, but these did not include preschool children or those in day care. Since August 2020, active polymerase chain reaction SARS-CoV-2 testing has been strongly recommended for all Finnish children with infectious symptoms that suggested COVID-19, as part of the national strategy.

The incidence of SARS-CoV-2 under 18 years of age was 470 per 100,000 (95% CI: 457-483). Five paediatric patients fulfilled the case definition of MIS-C from 1 March to 31 December 2020. There were no deaths. The annual population-based incidence of MIS-C was 0.45 per 100,000 in children under 18 years (95% CI: 0.15-1.06). The occurrence of MIS-C was 0.97 per 1,000 (95% CI: 0.31-2.26) when the rate was calculated against all laboratory-confirmed SARS-CoV-2 infections in that age group.

The annual population-based incidence rate of Kawasaki disease in children was 4.6 per 100,000 (95% CI: 3.5-6.1) in 2020 (Figure 1B). When we analysed the quarterly time periods from 2016 to 2020, the incidence ranged from 2.18 (95% CI: 1.40-3.25) in January to March 2018 to 0.55 (95% CI: 0.20-1.19) in both July to September 2020 and October to December 2020 (Table 1). Our time series analysis showed that the occurrence of Kawasaki disease decreased during the COVID-19 pandemic (Figure 2). The incidence rate ratio was 0.49 (95% CI: 0.29-0.72) when it was compared to the number of cases before the COVID-19 pandemic. Data from the National
TABLE 1 The population-based incidence of Kawasaki disease among children in Finland from 2016 to 2020. The COVID-19 pandemic began in March 2020

| Year | Incidence (95% CI) | Incidence (95% CI) | Incidence (95% CI) | Incidence (95% CI) | Incidence (95% CI) |
|------|--------------------|--------------------|--------------------|--------------------|--------------------|
| Q1   | 1.50 (0.88 to 2.41) | 1.69 (1.02 to 2.63) | 2.15 (1.38 to 3.19) | 1.90 (1.17 to 2.90) | 2.18 (1.40 to 3.25) |
| Q2   | 1.86 (1.49 to 2.84) | 1.06 (0.06 to 1.86) | 1.79 (1.09 to 2.76) | 1.26 (0.69 to 2.12) | 1.36 (0.76 to 2.25) |
| Q3   | 1.50 (0.88 to 2.41) | 1.51 (0.88 to 2.42) | 1.25 (0.68 to 2.10) | 1.99 (1.24 to 3.01) | 0.55 (0.20 to 1.19) |
| Q4   | 1.06 (0.55 to 1.85) | 1.15 (0.61 to 1.97) | 1.43 (0.81 to 2.33) | 1.44 (0.83 to 2.35) | 0.55 (0.20 to 1.19) |

Note: Q1 indicates January-March, Q2 indicates April-June, Q3 indicates July-September, and Q4 indicates October-December.

Infectious Disease Registry showed that the occurrence of influenza A, the respiratory syncytial virus, Mycoplasma pneumoniae and Streptococcus pneumoniae markedly decreased in children and adolescents aged 0-19 after March 2020 (Figure 3).

4 | DISCUSSION

This nationwide register-based study showed that MIS-C was a rare complication of the SARS-CoV-2 infection in children. The occurrence of Kawasaki disease markedly decreased during the COVID-19 pandemic, coinciding with a reduced number of respiratory infections in children. This was probably due to social distancing. The observed reduction suggests that transmissible microbes play an important role in the pathogenesis of Kawasaki disease.

MIS-C is a novel, serious and life-threatening disease associated with COVID-19 in children. There have been limited data on the proportion of children developing MIS-C after being infected with SARS-CoV-2. Most previous studies have presented the symptoms in selected patient series or clear temporal patient clusters during the peak of the pandemic. The data in our present study suggest that MIS-C is an infrequent complication of COVID-19, occurring in less than 0.1% of children in the population with laboratory-confirmed SARS-CoV-2. Despite this, our study may still markedly overestimate the proportion of children developing MIS-C after COVID-19, because not all children, especially those with asymptomatic infections, were tested for SARS-CoV-2. According to the Centers for Disease Control and Prevention, there were 3,185 patients and 36 deaths that met the case definition of MIS-C in the United States from mid-May 2020, when reporting began, to 29 March 2021. Data from a surge in cases in New York showed that the incidence of SARS-CoV-2 infections in those under 21 years of age was 322 per 100,000 and the incidence of MIS-C was two per 100,000 in the same age group. 

\[ \text{Incidence rate ratio (IRR)} = 0.3 \] compared to the incidence rate before the COVID-19 pandemic.

\[ \text{Seasonally adjusted incidence rate ratio (95% CI: } 0.29 \text{ to } 0.72) \]

\[ \text{Within the period of the COVID-19 pandemic. The start of the global pandemic in March 2020 is indicated with a vertical dash line.} \]

The aetiopathogenesis of Kawasaki disease
Koskela et al. has been extensively studied but is not fully understood. It appears to result from an interplay between genetic susceptibility and an infectious trigger, followed by an abnormal immune response. Ouldali et al. reported the emergence of Kawasaki disease in France during the COVID-19 pandemic and noted that the study population was multi-ethnic. The present study did not report the patients’ ethnicity, as that is not registered in medical records or national registers in Finland. Most Finnish residents are Northern European, but it is still possible that differences in ethnicity and genetic susceptibility may have influenced the occurrence of Kawasaki disease in our study population.

It has been suggested that respiratory viruses, such as influenza, the respiratory syncytial virus and enteroviruses, may trigger Kawasaki disease in genetically predisposed children. Esper et al. carried out a small case-control study and reported that a novel human coronavirus called New Haven coronavirus was detected more frequently in the respiratory secretions of patients with Kawasaki disease than in control patients. However, this finding has not been replicated in other studies. The idea that respiratory viruses are the main triggers of Kawasaki disease is supported by findings from its clear seasonal cycle, which is similar to that of respiratory infections, but the data are insufficient to prove its involvement with coronaviruses in general.

The present study had several strengths. This was a large nationwide study that used comprehensive data from high-quality national registers that also covered the periods before the COVID-19 pandemic. In addition, we performed a time series analysis, which has rarely been used to compare Kawasaki disease–like diagnoses before, and during, the COVID-19 pandemic. Since August 2020, Finland has strongly recommended free, national and easily available testing for all children with symptoms, no matter how mild they are. This means that we were able to estimate the total number of SARS-CoV-2 infections in the study population better than most previous reports of MIS-C patient clusters during the COVID-19 pandemic. In addition, we retrieved all MIS-C cases directly from the five university hospitals with paediatric intensive care units in Finland to ensure they had all been included in that data. We also present the changes in the epidemiology of common respiratory infections in the study population.

Our register-based study had some limitations. These included the lack of a specific ICD-10 code for MIS-C from the start of the COVID-19 emergency, which may have affected the accuracy of

**Figure 3** The occurrence of common respiratory viral infections (influenza A and RSV) (A-B), blood-culture positive Streptococcus pneumoniae infections (C) and Mycoplasma pneumoniae (D) in 0- to 19-year-old children and adolescents in Finland in 2016-2020. Data from the public National Infectious Diseases Register are presented. The beginning of COVID-19 pandemic is indicated with a vertical dash line. The 0 level has deliberately been elevated to show the absence of infections more clearly.
the register-based data. Finally, the low number of MIS-C cases did not allow us to carry out analyses to compare the characteristics of MIS-C and Kawasaki disease patients.

5 | CONCLUSION

This nationwide register-based study showed that MIS-C was a rare complication of the SARS-CoV-2 infection in children. The occurrence of Kawasaki disease decreased during the COVID-19 study period and coincided with a reduced number of respiratory infections due to social distancing. This finding suggests that transmissible microbes play an important role in the pathogenesis of Kawasaki disease.

CONFLICTS OF INTEREST
The authors have no conflicts of interest to declare.

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