Incidence of Kawasaki disease before and during the COVID-19 pandemic: a retrospective cohort study in Japan

Kyohei Iio,1,2 Kousaku Matsubara,1 Chisato Miyakoshi,2 Kunitaka Ota,3 Rika Yamaoka,4 Junji Eguchi,5 Osamu Matsumura,6 Takahiro Okutani,7 Ikuyo Ueda,8 Masahiro Nishiyama9

ABSTRACT

Background Epidemiological studies in Kawasaki disease (KD) have suggested infectious aetiology. During the COVID-19 pandemic, measures for mitigating SARS-CoV-2 transmission also suppress the circulation of other contagious microorganisms. The primary objective is to compare the number and incidence of KD before and during the COVID-19 pandemic in Japan, and the secondary objective is to investigate temporal association between the KD epidemiology and activities of SARS-CoV-2 and other viral and bacterial infections.

Methods A retrospective cohort study was conducted between 2016 and 2020 in Kobe, Japan. We collected information of hospitalised KD children in Kobe. Child population was identified through the resident registry system. Activity of COVID-19 and 11 other infectious diseases was derived from a public health monitoring system. Monthly change of KD incidence was analysed using a difference-in-difference regression model.

Results Throughout the study period, 1027 KD children were identified. KD had begun to decline in April 2020, coinciding with the beginning of the COVID-19 pandemic. The number of KD cases (n=66) between April and December 2020 was 40% of the average in the same period in 2016–2019 (165/year). Annual KD incidence was 315, 300, 353, 347 and 188/100 000 children aged 0–4 years in 2016–2020, respectively. The difference-value of KD incidence was significantly reduced in the fourth quarter in 2020 (−15.8, 95% CI −28.0 to −3.5), compared with that in 2016–2019. Sentinel surveillance showed a marked decrease of all infectious diseases except exanthema subitum after the beginning of the COVID-19 pandemic. There were 86 COVID-19 cases aged <10 years and no KD children associated with COVID-19.

Conclusion This study showed that the number and incidence of KD was dramatically reduced during the COVID-19 pandemic in Japan. This change was temporally associated with decreased activities of various infectious diseases other than COVID-19, supporting the hypothesis of infection-triggered pathogenesis in KD.

INTRODUCTION

Kawasaki disease (KD) is an acute systemic vasculitis that occurs in young children.1 Epidemiological studies describing incidence, age distribution and seasonality in KD suggest an infectious aetiology.2 3 However, despite intensive efforts over decades, the aetiology has remained to be elucidated. Some investigators proposed that KD is a consequence of an abnormal immunological response evoked by any of infectious agents in genetically susceptible individuals, and that these microorganisms may trigger the activation of onset only.2 4 5

Since December 2019, COVID-19 caused by SARS-CoV-2 has been reported globally.6–7 Accumulating evidence indicated that children appeared to be less affected and to exhibit milder manifestations than adults.5–7

What is known about the subject?

► Continuing nationwide surveys on Kawasaki disease (KD) show that KD incidence has consistently increased since 1970 in Japan.
► Existing research has hypothesised that KD is a consequence of an abnormal immunological response evoked by any of infectious agents in genetically susceptible individuals.
► Measures for avoiding disease transmission during the COVID-19 pandemic, such as use of masks and handwashing, suppress circulation of other contagious microorganisms.

What this study adds?

► The number and incidence of Kawasaki disease (KD) was dramatically decreased to less than 50% during COVID-19 pandemic, compared with that before the pandemic in Japan.
► The annual KD incidence of 188/100 000 children aged 0–4 years in 2020 was reduced to the nationwide estimates around 2006 in Japan.
► The epidemiological change in KD coincided with decreased activities in various infectious diseases other than COVID-19, supporting the hypothesis of infection-triggered pathogenesis in KD.
However, an exception is the newly emerging, multi-system inflammatory syndrome in children (MIS-C), which shares some clinical and laboratory features with those of KD. Children with MIS-C have been documented mainly in Europe and the USA, and little is known about MIS-C’s prevalence in Asia. Measures for mitigating SARS-CoV-2 transmission have been widely promoted in Japan such as use of masks, handwashing and school closures. These measures may also suppress the circulation of seasonal influenza and other contagious microorganisms, and may reduce KD under the presumption of its infection-triggered pathogenesis.

These backgrounds raise concerns on an epidemiological association between the number of cases of KD and the magnitude of the COVID-19 pandemic, especially in East Asia where the incidence of KD is the highest worldwide. Studies addressing this issue were limited by small sample sizes, being single-centre studies and being evaluated during only the initial stage of the COVID-19 pandemic. In the present study, we conducted a retrospective cohort study in Kobe, Japan. Our primary objective is to compare the number and incidence of KD before and during the COVID-19 pandemic, and secondary objective is to investigate temporal association between the KD epidemiology and activities of SARS-CoV-2 and other viral and bacterial infections.

METHODS

Study setting and design

Kobe city is located in west Japan with a population of approximately 1.5 million including approximately 200,000 children aged 0–15 years. In 2016, there were 14 hospitals in which children could be hospitalised and treated by paediatricians in Kobe, and four of them had been closed by the end of the study. The study covered all hospitals with paediatric departments in Kobe. Questionnaires were mailed or emailed to representative paediatricians at the 14 hospitals in Kobe and four hospitals in three cities adjacent to Kobe (two hospitals in Akashi and one each in Sanda and Ashiya).

Data collection

A retrospective cohort study was conducted between 1 January 2016 and 31 December 2020. The surveillance structured form included sex, age in months, date of onset of KD, date of admission, presence or absence of transfer to another institute during treatment and history on April 9, 2021 by guest. Protected by copyright.
With this model, we predicted the number of patients with KD using the data before March 2020, then visually compared it with that actually observed. We set four separate sampling chains, each consisting of 12 000 samples (including 2000 samples discarded for convergence). We evaluated sampling convergence by Gelman-Rubin statistics (R-hat) and by visually inspecting a trace plot. All statistical analyses were performed by the R software program, V.4.0.2 (R Foundation for Statistical Computing, Vienna, Austria). For Bayesian model analyses, we used the probabilistic programming language Stan (Stan development team).

Patient and public involvement
There were no patients or public involvement in the research design, process and research findings dissemination.

RESULTS
Of the 18 eligible facilities, all responded to the survey. A total of 1027 KD children were identified with a male/female ratio of 579/448. Age of onset ranged from 0 month to 13 years (median: 28 months, IQR: 15–47 months). Figure 1 shows the monthly changes of patients with KD in each year. Seasonality was observed during 2016–2019 with a peak generally in winter (December–January) and a reduction in May and June. However, such seasonality disappeared after April 2020, when KD began to decline. The annual number of 120 in 2020 was approximately half (53%) of the average (227/year) during the previous 4 years. When results were limited to the period between April and December, the reduction rate was more prominent (60%, 66 in 2020 vs 165 in 2016–2019). On the other hand, distributions of sex (male/female: 513/394 vs 66/54) and age of onset (median age (IQR): 28 (15–47) vs 29 (16–47) months) did not significantly differ between 2016–2019 and 2020. Notably, the beginning of the decline of KD in 2020 coincided with the onset of a COVID-19 outbreak in Kobe (figure 2): the first adult COVID-19 case was identified on March 2, and the first child aged <10 years on April 13. Thereafter, the first spike of the pandemic developed between April and May. Between late May and early June, COVID-19 cases were reduced; however, the second and third waves occurred in August and November–December, respectively (figure 2). Since the beginning of the pandemic, 86 children aged <10 years with COVID-19 were identified, accounting for 2.5% of the entire cases in 2020. There were 221 COVID-19 cases in the aged 10–19 years population. Of the 86 children aged <10 years, 27 children were hospitalised. All were asymptomatic or manifested mild symptoms and were discharged without sequelae. There were no cases of MIS-C and KD following COVID-19 within 2 months.

As previous nationwide surveys reported about KD in children aged 0–4 years, we examined changes of KD in this age group.22 23 The annual incidence of KD in children aged 0–4 years was 315, 300, 353, 347 and 188 (per 100 000 children aged 0–4 years) in 2016, 2017, 2018, 2019 and 2020, respectively. Figure 3 shows the result of a Bayesian time series analysis. We found that patients with KD had begun to decrease after April 2020, and continued to reduce until December, compared with the prediction on the basis of data before 2020. When limited to the period between April and December, the reduction rate of KD incidence was 53% in 2020 compared with the average incidence in 2016–2019 (111 vs 237/100 000 children aged 0–4 years). The result of difference-in-difference analysis is summarised in table 1. The incidence rate of KD significantly decreased in the fourth quarter in 2020 compared with 2016 to 2019 (−15.8, 95% CI (−28.0 to −3.5)). From visual assessment of figure 1, the assumptions for the difference-in-difference regression model were considered unviolated.

Several practices implemented to contain SARS-CoV-2 transmission have likely had an influence on the activity of other contagious organisms. Sentinel surveillance showed that the annual incidence of 11 infectious diseases, except exanthema subitum, was markedly lower than that in the previous 4 years. Weekly changes of four representative infections (RSV infection, influenza, herpangina and group A streptococcal pharyngitis) in children aged 0–4 years are shown in figure 4. RSV infections in 2020 were less than one-eighth the average in 2016–2019 (figure 4A). Influenza activity between November and December in 2020 remained at

Figure 1 Monthly change of Kawasaki disease cases in 2016–2020.

Figure 2 Daily case numbers of COVID-19 in 2020. Orange and blue bars indicate individuals with <10 years and ≥10 years of age, respectively.
interseasonal levels (figure 4B). In addition, a decrease in case numbers was observed in group A streptococcal pharyngitis (figure 4C) and herpangina (figure 4D).

DISCUSSION

This retrospective cohort study showed that in Japan KD children in 2020 were dramatically decreased to approximately half of the annual average in 2016–2019. When limited to the COVID-19 pandemic period, the reduction rate was more prominent. These changes were also found in KD incidence among children aged 0–4 years. A difference-in-difference regression model showed that the incidence in the fourth quarter in 2020 was significantly lower than that in 2016–2019. The KD incidence has shown a consistent increase since 1970 in Japan, which may be reflected by an improved awareness of the diagnosis and possible increase of multiple aetiological agents for KD.22 23 The abrupt reduction observed in this study was a marked contrast to this Japanese KD epidemiology. The KD incidence in 2020 (188/100 000 children aged 0–4 years) returned to the estimates around 2006.23 Notably, the reduction of KD was temporally associated with the COVID-19 pandemic as well as low activities in various infections other than SARS-CoV-2.

The present study has some strengths. It covered all hospitals even small institutes in Kobe and the adjacent three cities with complete response, and the number of children living in Kobe was based on a public registry. Children with KD are admitted and treated in hospitals in Japan; hence, these approaches allow us to accurately analyse KD incidence. Furthermore, the incidences reported in 2016, 2017 and 2018 in the present study were quite similar to the estimates in Japanese nationwide surveys22 23 (315, 300 and 353 vs 309, 317 and 359/100 000 children aged 0–4 years, respectively), which suggests that our cohort was representative. Simultaneously, we could assess the activity of SARS-CoV-2 and various other pathogens using a public health monitoring system.

KD has a varied incidence among children of different races and ethnicities with the highest incidence in East Asia, especially in Japan and South Korea.1 A seroprevalence study in a single Japanese institute showed no apparent association of COVID-19 and KD.17 An early epidemiological analysis in South Korea also failed to detect this association.19 We contrarily showed a temporal association between a reduction of KD and the COVID-19 pandemic. The discrepancy between the South Korean study and ours may be reflected by the evaluation period; our study was conducted throughout the year, while the former was limited to the early stage of the COVID-19 pandemic.19 We found no MIS-C cases during the

| Table 1 Estimated monthly number of Kawasaki disease patients aged 0–4 years and difference-in-difference value between 2020 and 2016–2019 |
|---------------------------------------------------------------|
| Estimated monthly no of cases per 100 000 children aged 0–4 years | Difference-in-difference value in 2020 vs 2016–2019 (95% CI) |
| January–March | 30.2 | 25.6 | −4.4 | (−16.6 to 7.8) |
| April–June | 22.8 | 13.8 | −9.0 | (−20.2 to 4.2) |
| July–September | 26.6 | 14.0 | −12.6 | (−28.0 to −3.5) |
| October–December | 29.6 | 9.2 | −20.4 | (−27.2 to −3.6) |
pandemic, in line with the other studies in Japan and South Korea. This may be explained by recent findings that MIS-C shows a predilection for individuals of black or Hispanic descent. In the USA, as of 20 February 2021, 2060 MIS-C cases were identified while COVID-19 cases in children aged 0–17 years were up to 2.39 million. In contrast, because of small number of COVID-19 cases (n=307) in the patients under 20 years of age in this study, we were not able to conclude that MIS-C was rare in Japanese children. A larger study is warranted to definitively assess these associations in Japan.

The exact reasons of the significant reduction of KD are unclear. It seems to be unlikely that SARS-CoV-2 itself directly suppressed the occurrence of KD, because only a small number of children aged <10 years were affected with this virus in Kobe. The most plausible explanation is that various measures to reduce the spread of SARS-CoV-2 also suppressed the transmission activities of other microorganisms, and thereby reduced KD. Weekly epidemics of 11 infectious diseases in Kobe paralleled nationwide surveillance reports. The disappearance of seasonality in KD in 2020 provides evidence for the contribution of multiple viruses to KD pathogenesis. Taken together, our findings support the hypothesis of infection-triggered pathogenesis in KD.

Several limitations are included this study. First, our findings indicated a temporal association between the decline in KD and the increase of SARS-CoV-2-affected individuals as well as a reduction of various infections. However, studies evaluating epidemiological time series could not fully account for this causality. Numerous infectious agents not examined in this study or other factors, such as environmental and behavioural changes, may contribute to KD occurrence. Second, our study was conducted in Japan. Whether similar findings are generalised to other regions with different genetic ethnicities, magnitude of COVID-19 pandemic, and protective measures remains to be determined.

CONCLUSION
The present study showed that KD significantly decreased during the COVID-19 pandemic in Kobe, Japan. Epidemiological analysis through the public surveillance system also showed a profound decrease of various types of infections, probably due to the implementation of protective measures against SARS-CoV-2 transmission. Our data support, at least in part, the hypothesis that many microorganisms are involved to trigger KD occurrence.

Author affiliations
1Department of Pediatrics, Kobe City Nishi-Kobe Medical Center, Kobe, Hyogo, Japan
2Department of Pediatrics, Kobe City Medical Center General Hospital, Kobe, Hyogo, Japan
3Department of Pediatrics, Konan Medical Center, Kobe, Hyogo, Japan
4Department of Pediatrics, National Hospital Organization Kobe Medical Center, Kobe, Hyogo, Japan
5Department of Pediatrics, Kobe City Medical Center West Hospital, Kobe, Hyogo, Japan
6Department of General Pediatrics, Hyogo Prefectural Kobe Children’s Hospital, Kobe, Hyogo, Japan
7Department of Pediatrics, Saiseikai Hyogo-ken Hospital, Kobe, Hyogo, Japan
8Department of Pediatrics, Japan Community Health Care Organization Kobe Central Hospital, Kobe, Hyogo, Japan
9Department of Pediatrics, Kobe University Graduate School of Medicine, Kobe, Hyogo, Japan

Acknowledgements The authors appreciate S. Nukina (Department of Pediatrics, Akashi Municipal Hospital), N. Yokoyama (Department of Pediatrics, Akashi Medical Center), A. Hirase (Department of Pediatrics, Kumon Hospital), T. Inoue (Department of Pediatrics, Sanda City Hospital), T. Monta (Department of Pediatrics, Ashiya Municipal Hospital), Y. Matsuura (Kobe Tokushukai Hospital), H. Yamashita (Director of hospital, Kobe Red Cross Hospital) and K. Tsukuda (Department of Pediatrics, Mahoishi Hospital), for providing the clinical records of the patients.

Contributors KM and CM conceptualised and designed the study. KI, KM and CM drafted the manuscript. CM carried out statistical analyses. All authors collected the data and critically reviewed the manuscript. All authors read and approved the final manuscript as submitted.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not required.

Ethics approval The study was approved by the Ethics Committee of Kobe City Nishi-Kobe Medical Center (receipt ID: 2020–37). Review board approval in each institution was not necessarily required for the assessment of anonymised surveillance.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available on reasonable request. KI and KM had full access to the data in the study and take responsibility for the accuracy and integrity of the data. Data are available on reasonable request.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/.

ORCID iD
Kyohei Ito http://orcid.org/0000-0002-7356-5307

REFERENCES
1 Burns JG, Glodé MP. Kawasaki syndrome. Lancet 2004;364:533–44.
2 Principi N, Rigante D, Esposito S. The role of infection in Kawasaki syndrome. J Infect 2013;67:1–10.
3 Rowley AH. Is Kawasaki disease an infectious disorder? Int J Rheum Dis 2018;21:20–5.
4 Nagao Y, Urabe C, Nakamura H, et al. Predicting the characteristics of the aetiological agent for Kawasaki disease from other paediatric infectious diseases in Japan. Epidemiol Infect 2016;144:478–92.
5 Swann OV, Holden KA, Turtle L, et al. Clinical characteristics of children and young people admitted to hospital with covid-19 in United Kingdom: prospective multicentre observational cohort study. BMJ 2020;370:m3249.
6 Wald ER, Schmitt KM, Gusland DY. A pediatric infectious disease perspective on COVID-19. Clin Infect Dis 2020;ciaa1095.
7 Viner RM, Myron OT, Bonell C, et al. Susceptibility to SARS-CoV-2 infection among children and adolescents compared with adults: a systematic review and meta-analysis. JAMA Pediatr 2021;175:e204573.
8 Riphagen I, Gomez X, Gonzalez-Martinez C, et al. Hyperinflammation and shock in children during COVID-19 pandemic. Lancet 2020;395:1607–8.

Ito K, et al. BMJ Paediatrics Open 2021;5:e001034. doi:10.1136/bmjpo-2021-001034

Open access
9 Verdoni L, Mazza A, Gervasoni A, et al. An outbreak of severe Kawasaki-like disease at the Italian epicentre of the SARS-CoV-2 epidemic: an observational cohort study. *Lancet* 2020;395:1771–8.

10 Toubiana J, Poirault C, Corsia A, et al. Kawasaki-like multisystem inflammatory syndrome in children during the covid-19 pandemic in Paris, France: prospective observational study. *BMJ* 2020;369:m2094.

11 Feldstein LR, Rose EB, Horwitz SM, et al. Multisystem inflammatory syndrome in U.S. children and adolescents. *N Engl J Med* 2020;383:334–46.

12 Jiang L, Tang K, Levin M, et al. COVID-19 and multisystem inflammatory syndrome in children and adolescents. *N Engl J Med* 2020;383:334–46.

13 Verdoni L, Mazza A, Gervasoni A, et al. An outbreak of severe Kawasaki-like disease at the Italian epicentre of the SARS-CoV-2 epidemic: an observational cohort study. *Lancet* 2020;395:1771–8.

14 Toubiana J, Poirault C, Corsia A, et al. Kawasaki-like multisystem inflammatory syndrome in children during the covid-19 pandemic in Paris, France: prospective observational study. *BMJ* 2020;369:m2094.

15 Feldstein LR, Rose EB, Horwitz SM, et al. Multisystem inflammatory syndrome in U.S. children and adolescents. *N Engl J Med* 2020;383:334–46.

16 Jiang L, Tang K, Levin M, et al. COVID-19 and multisystem inflammatory syndrome in children and adolescents. *N Engl J Med* 2020;383:334–46.

17 Choe S-A, An HS, Choe YJ. No temporal association between human coronavirus and Kawasaki disease: national data from South Korea. *J Med Virol* 2021;93:585–7.

18 Ayusawa M, Sonobe T, Uemura S, et al. Revision of diagnostic guidelines for Kawasaki disease (the 5th revised edition). *Pediatr Int* 2005;47:232–4.

19 Kobayashi T, Ayusawa M, Suzuki H, et al. Revision of diagnostic guidelines for Kawasaki disease (6th revised edition). *Pediatr Int* 2020;62:1135–6.

20 Makino N, Nakamura Y, Yoshi M, et al. Nationwide epidemiologic survey of Kawasaki disease in Japan, 2015-2016. *Pediatr Int* 2019;61:397–403.

21 Ae R, Makino N, Kosami K, et al. Epidemiology, treatments, and cardiac complications in patients with Kawasaki disease: the nationwide survey in Japan, 2017-2018. *J Pediatr* 2020;225:23–9.

22 Kim YJ, Park H, Choi YY, et al. Defining association between COVID-19 and the multisystem inflammatory syndrome in children through the pandemic. *J Korean Med Sci* 2020;35:e204.

23 Rowley AH. Multisystem inflammatory syndrome in children and Kawasaki disease: two different illnesses with overlapping clinical features. *J Pediatr* 2020;224:129–32.

24 CDC. COVID data Tracker. Available: https://covid.cdc.gov/covid-data-tracker/#demographics [Accessed 20 Feb 2021].

25 CDC. COVID data Tracker. Available: https://covid.cdc.gov/covid-data-tracker/#demographics [Accessed 20 Feb 2021].

26 Health Department-Reported cases of multisystem inflammatory syndrome in children (MIS-C) in the United States /CDC. Available: https://www.cdc.gov/mis-c/cases/index.html [Accessed 20 Feb 2021].

27 National Institutes Infectious Diseases. Infectious diseases Weekly report. Available: https://www.niid.go.jp/niid/ja/idwr.html [Accessed 15 Jan 2021].