

# COVID-19 Vaccine Acceptance Among Parents of Children With Multisystem Inflammatory Syndrome in Children

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**Abstract:** Data on COVID-19 vaccine acceptability among parents of children with multisystem inflammatory syndrome (MIS-C) are limited. In this cohort of children with MIS-C, enrolled in the Swissped RECOVERY trial (NCT 04826588), comparing intravenous immunoglobulins or methylprednisolone, who, in accordance with Swiss guidelines, were recommended for SARS-CoV-2 vaccination, 65% (73/112) of parents reported being vacci-

nated against SARS-CoV-2 before the MIS-C, while 70% were vaccinated after the MIS-C episode of their child. None of the children were vaccinated before the occurrence of the MIS-C, and only 9% (5/56) received the COVID-19 vaccine after the MIS-C. The predominant barriers to COVID-19 vaccination were concerns over potential side effects and insufficient support from their doctors. This emphasizes the crucial role of health care providers in promoting COVID-19 vaccination among children.

**Key Words:** vaccine acceptance, SARS-CoV-2, COVID-19 vaccine, MIS-C, PIMS

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Vaccination against SARS-CoV-2 has demonstrated remarkable efficacy in averting multisystem inflammatory syndrome (MIS-C) following SARS-CoV-2 infection in adolescents and children.<sup>1,2</sup> In Switzerland, SARS-CoV-2 vaccination was recommended for children 16 years of age and above starting from May 2021 onward, followed by an extension to children 12 years old and above from June 2021. Subsequently, from December 2021, children 5 years of age and above were included in the vaccination recommendation.<sup>3</sup> Several countries, including the United States,<sup>4</sup> Canada<sup>5</sup> and Switzerland, recommended SARS-CoV-2 vaccination for children who had experienced MIS-C. The rationale behind this recommendation rested on the potential to prevent recurrence of MIS-C through vaccination, considering that COVID-19 vaccination did not exhibit evidence of being a risk factor for MIS-C, while SARS-CoV-2 infection was unequivocally recognized as a risk factor for MIS-C. In addition, the vaccine was well tolerated in MIS-C children and was not associated to any relapse of MIS-C episode.<sup>6,7</sup> In other countries, the vaccination strategy for children post-MIS-C was debated. In France, for instance, SARS-CoV-2 vaccination was initially contraindicated in this population but their position was revised in February 2022 to recommend vaccination.<sup>8</sup> Limited research has been conducted on parental vaccination and attitudes toward vaccination of children who have experienced MIS-C.

## METHODS

This is an ancillary study nested in the prospective Swissped RECOVERY trial of the Swiss National Clinical Trials Portal (SNCTP000004720) and clinicaltrials.gov (NCT 04826588), an investigator-initiated randomized multicenter open-label 2-arm trial in which children hospitalized with MIS-C were randomly assigned 1:1 to intravenous immunoglobulins or intravenous methylprednisolone at 10 Swiss pediatric hospitals.<sup>9,10</sup> Cohort characteristics and outcomes have been published previously.<sup>9</sup> Included children were followed until 6 months after randomization, and their families received a questionnaire (Questionnaire, Supplemental Digital Content 2, <http://links.lww.com/INF/F338>) regarding COVID-19 vaccination at 6 months after hospital discharge.

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See collaborator group, Supplemental Digital Content 1, <http://links.lww.com/INF/F337>.

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For parents, the criterion for not being opposed to vaccination was having received at least 1 dose of the SARS-CoV-2 vaccine. This approach was adopted due to the absence of information regarding parental demographics, underlying health conditions, occupation or historical exposure, factors that would have enabled us to determine the requisite number of vaccine doses for considering vaccination as complete.

Data were summarized using the number (percentage) for categorical variables and the median (interquartile range) for continuous variables. Summary statistic comparisons between groups were performed using the Fisher exact test for categorical variables and the Wilcoxon rank-sum test for continuous variables. All analyses were based on the complete case data only and were performed using the statistical software R (version 4.0.3, R Development Core Team, Vienna, Austria<sup>11</sup>). A significance level of 5% was used for all statistical analysis.

## RESULTS

Between May 21, 2021 and April 15, 2022, a total of 75 patients were included in the intention-to-treat analysis. Detailed information on the cohort, including baseline characteristics, is presented in the original publication.<sup>9</sup> The vaccination questionnaire was completed for 56 children presenting with a MIS-C (74%; 56/75), resulting in 56 patients and 112 parents with available data for this analysis. We have included the patient characteristics from the original manuscript in Table, Supplemental Digital Content 3, <http://links.lww.com/INF/F339> (“Total patients”) to observe any important difference that could suggest a bias in our analysis. No big differences between the 2 populations were observed.

Before the occurrence of MIS-C in their offspring, 65% (73/112) of parents reported being vaccinated against SARS-CoV-2 and 70% (78/112) after the MIS-C episode. Among the 73 parents who had previously been vaccinated, 21% (15/73) received additional doses of the vaccine after the MIS-C episode of their child. In the group of parents who had initially chosen not to get vaccinated, 11% (4/37) got vaccinated (1 parent received a single dose, 2 received 2 doses and 1 received 3 doses) (Fig. 1). Most individuals within a couple shared the same vaccination status and there was no indication that the timing of the MIS-C occurrence influenced the vaccination status of the parents (Figure, Supplemental Digital Content 4, <http://links.lww.com/INF/F340>).

SARS-CoV-2 vaccination was recommended for children 5 years old and above in December 2021, while it had been recommended for children 12 years old and older since June 2021. In our study, none of the children were vaccinated before the occurrence of MIS-C, and only 9% (5/56) of children received vaccination after the MIS-C episode. The parents provided various reasons for not vaccinating their children before the MIS-C and after, which are summarized in Figure, Supplemental Digital Content 5, <http://links.lww.com/INF/F341>. Parents indicated that half of the children (27/54) were not eligible for vaccination before the MIS-C, and for 44% (24/54), vaccination was felt unnecessary by the parents. Safety concerns were reported for 41% (22/54) of children, and for 4% (2/54) and 6% (3/54), the parents’ decision not to vaccinate their child was influenced by relatives/close friends and their doctor, respectively. Following the MIS-C episode, the reasons cited by parents for not vaccinating their child remained unchanged with respect to their former concerns about safety. There was a decrease in the number of parents reporting eligibility as a reason and an increase in the percentage of parents who felt that vaccination was unnecessary for their children. Notably, 23% (11/47) of parents indicated that their doctor advised against vaccinating their children after the MIS-C episode.

We assessed selected clinical factors (Table, Supplemental Digital Content 3, <http://links.lww.com/INF/F339>) that could potentially impact the decision of parents to vaccinate or not vaccinate their children after experiencing MIS-C. Our findings indicate that the only significant factor influencing this decision is the age at time of randomization (Table, Supplemental Digital Content 3, <http://links.lww.com/INF/F339>).

## DISCUSSION

Sixty-five percent of parents were vaccinated against SARS-CoV-2 before the MIS-C episode of their child. This number is comparable to previous reports from Switzerland,<sup>12</sup> US,<sup>13</sup> Ireland and United Kingdom.<sup>14</sup> We observed that it was rare for parents who had not been vaccinated before the MIS-C episode to change their mind after witnessing their child being hospitalized with MIS-C as a severe complication of COVID-19. This is in contradiction with previous studies reporting that higher personal and family risk for COVID-19 was associated with stronger COVID-19 vaccination willingness.<sup>15,16</sup> Additionally, most couples shared the same vaccination status, indicating a consistent alignment within families and a lack of discrepancies in vaccination attitudes.

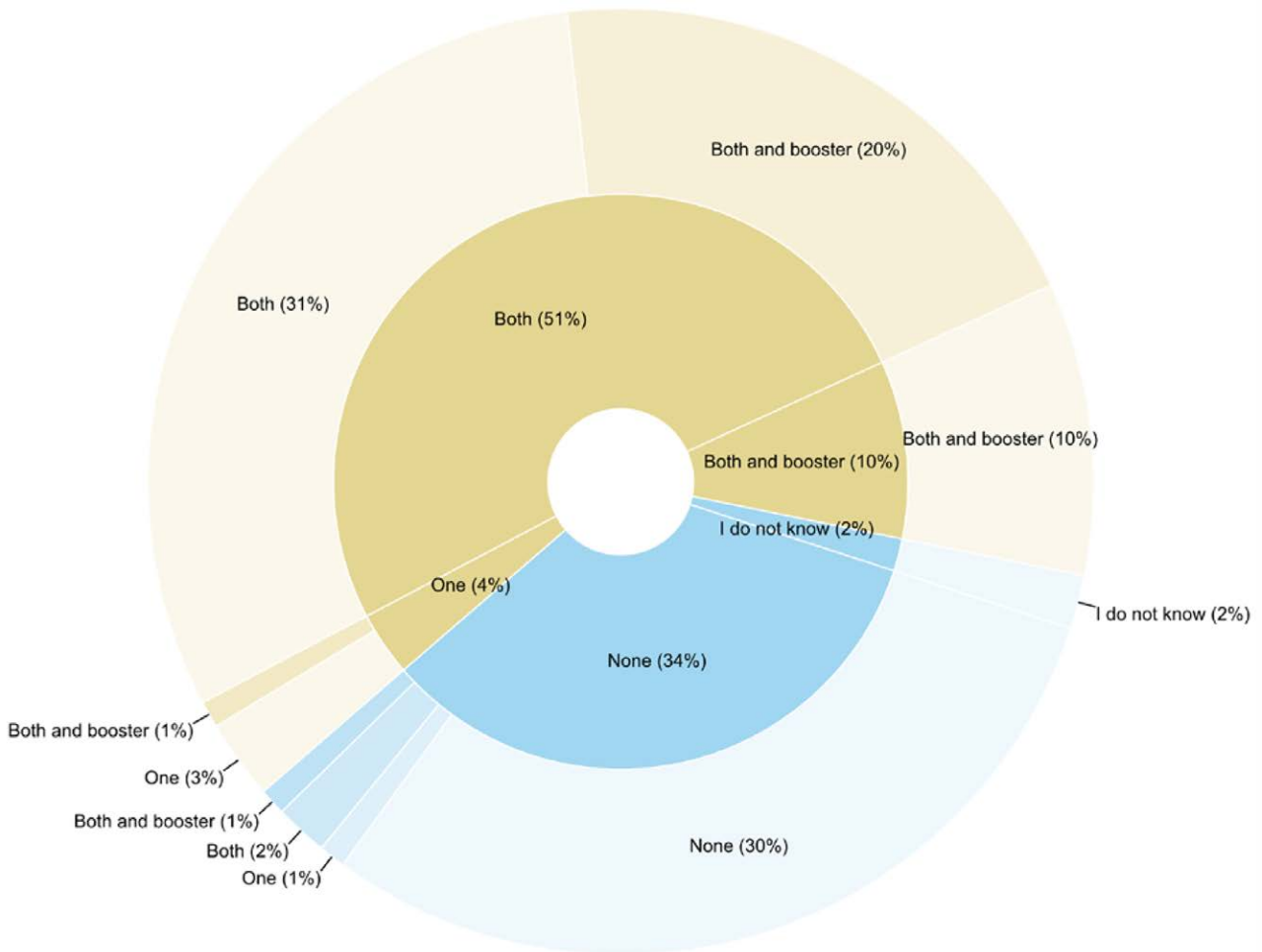
Concerning the vaccination of children, our study started 1 month after the COVID-19 vaccine was eligible for children 12 years of age and above in Switzerland and 7 months before the COVID-19 vaccine was eligible for children 5 years old and above. In our study, only 9% of the children had received the COVID-19 vaccine after their MIS-C episode. The sole differentiating factor between vaccinated and unvaccinated children was age, wherein vaccinated children had a higher median age in comparison to unvaccinated children, which could be explained by the fact that age recommendations for vaccination were progressively introduced, with younger patients being vaccinated later. However, given that only a small number of children were vaccinated at all in our study, this might not be relevant. In addition, most children in our study and in the original study were older than 5 years and so eligible for vaccination. Other studies also reported that parental willingness to vaccinate their children against SARS-CoV-2 decreased with the younger age of the child.<sup>17–19</sup> A significant concern expressed by parents both before and after the MIS-C episode was related to safety. The apprehensions included fears regarding potential side effects and the perception that the vaccine was not well researched, similar to other studies.<sup>6,20</sup> External factors and the environment also played a role in parents’ decision-making, including advice from family members or doctors. Indeed, other studies reported that pediatricians were among the most trusted source of information for COVID-19 vaccination.<sup>19,21</sup>

In conclusion, our study found that the acceptance of COVID-19 vaccines among parents of children with MIS-C was comparable to that of the general Swiss population, and it did not increase after the MIS-C episode in their child. A minority of MIS-C children received the COVID-19 vaccine after their MIS-C episode, primarily among the older age group. The main reasons cited for COVID-19 vaccine refusal were concerns about potential side effects and inadequate encouragement from doctors. These findings highlight the vital role of health care providers in promoting COVID-19 vaccination among children and in effectively addressing parental concerns to enhance vaccine acceptance.

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Parents vaccine status before and after hospitalization



**FIGURE 1.** Comparative pie chart illustrating parental vaccination status before their child’s MIS-C episode (inner circle) and 6 months post-MIS-C (outer circle). Due to rounding, the total percentage may sum to 101%. Light shades denote parental vaccination status before MIS-C, while darker shades represent status 6 months post-MIS-C. Notably, 2 parents indicated uncertainty regarding their SARS-CoV-2 vaccination. It is presumed that 1 parent completed the questionnaire, potentially explaining the inability to ascertain the other parent’s vaccination status in this instance.

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### REFERENCES

- Zambrano LD, Newhams MM, Olson SM, et al; Overcoming COVID-19 Investigators. Effectiveness of BNT162b2 (Pfizer-BioNTech) mRNA vaccination against multisystem inflammatory syndrome in children among persons aged 12–18 years - United States, July–December 2021. *MMWR Morb Mortal Wkly Rep.* 2022;71:52–58.
- Zambrano LD, Newhams MM, Olson SM, et al; Overcoming COVID-19 Investigators. BNT162b2 mRNA vaccination against coronavirus disease 2019 is associated with a decreased likelihood of multisystem inflammatory syndrome in children aged 5–18 years—United States, July 2021 - April 2022. *Clin Infect Dis.* 2023;76:e90–e100.
- F.C.o. Switzerland. COVID-19: vaccination recommended and possible for children from the beginning of January. 2021. <https://www.admin.ch/gov/en/start/documentation/media-releases.msg-id-86451.html#:~:text=beginning%20of%20January-,COVID%2D19%3A%20vaccination%20recommended%20and%20possible%20for%20children,from%20the%20beginning%20of%20January&text=Bern%2C%2014.12.,aged%20between%20five%20and%20eleven>.
- C. Centers for Disease. Interim Clinical Considerations for Use of COVID-19 Vaccines in the United States. 2021. <https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us-appendix.html>.
- C. Government. COVID-19 vaccine: Canadian Immunization Guide. 2021. <https://www.canada.ca/en/public-health/services/publications/healthy-living/canadian-immunization-guide-part-4-active-vaccines/page-26-covid-19-vaccine.html>.
- Epalza C, Prieto-Tato L, Pino R, et al; EPICO-AEP Working Group. Safety and acceptance of COVID-19 vaccination after multisystem inflammatory syndrome in children (MIS-C) in Spain. *J Pediatric Infect Dis Soc.* 2022;11:471–473.
- Minoia F, Lucioni F, Heshin-Bekenstein M, et al. Approaches to SARS-CoV-2 and other vaccinations in children with a history of multisystem inflammatory syndrome (MIS-C): an international survey. *Front Pediatr.* 2022;10:1030083.
- Conseil d'Orientation de la Stratégie Vaccinale/Conseil d'Orientation de la Stratégie Vaccinale du gouvernement français. Vaccination des enfants et adolescents ayant développé un PIMS, Note du 8 février. 2022. [https://sante.gouv.fr/IMG/pdf/cosv\\_-\\_note\\_du\\_8\\_fevrier\\_-\\_vaccination\\_des\\_enfants\\_ayant\\_developpe\\_un\\_pims.pdf](https://sante.gouv.fr/IMG/pdf/cosv_-_note_du_8_fevrier_-_vaccination_des_enfants_ayant_developpe_un_pims.pdf).
- Welzel T, Atkinson A, Schobi N, et al; Swissped RECOVERY Trial Group. Methylprednisolone versus intravenous immunoglobulins in children with paediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2 (PIMS-TS): an open-label, multicentre, randomised trial. *Lancet Child Adolesc Health.* 2023;7:238–248.
- Welzel T, Schobi N, Andre MC, et al; Swissped Recovery Trial. Multicenter randomized trial of methylprednisolone vs intravenous immunoglobulins to treat the pediatric inflammatory multisystem syndrome-temporally associated with SARS-CoV-2 (PIMS-TS): protocol of the Swissped RECOVERY Trial. *Front Pediatr.* 2022;10:905046.
- R Foundation for Statistical Computing. R Care Team: a language and environment for statistical computing. Austria; 2020.
- Wisniak A, Baysson H, Pullen N, et al; Specchio-COVID19 study group. COVID-19 vaccination acceptance in the canton of Geneva: a cross-sectional population-based study. *Swiss Med Wkly.* 2021;151:w30080.
- Malik AA, McFadden SM, Elharake J, et al. Determinants of COVID-19 vaccine acceptance in the US. *EClinicalMedicine.* 2020;26:100495.
- Hursh SR, Strickland JC, Schwartz LP, et al. Quantifying the impact of public perceptions on vaccine acceptance using behavioral economics. *Front Public Health.* 2020;8:608852.
- Blanchard-Rohner G, Caprettini B, Rohner D, et al. Impact of COVID-19 and intensive care unit capacity on vaccination support: evidence from a two-leg representative survey in the United Kingdom. *J Virus Erad.* 2021;7:100044.
- Peirola A, Posfay-Barbe KM, Rohner D, et al. Acceptability of COVID-19 vaccine among hospital employees in the Department of Paediatrics, Gynaecology and Obstetrics in the University Hospitals of Geneva, Switzerland. *Front Public Health.* 2021;9:781562.
- Galanis P, Vranka I, Siskou O, et al. Willingness, refusal and influential factors of parents to vaccinate their children against the COVID-19: a systematic review and meta-analysis. *Prev Med.* 2022;157:106994.
- Rane MS, Robertson MM, Westmoreland DA, et al. Intention to vaccinate children against COVID-19 among vaccinated and unvaccinated US Parents. *JAMA Pediatr.* 2022;176:201–203.
- Mondal P, Sinharoy A. The influence of pediatricians' recommendation on caregivers' COVID-19 vaccine acceptance for children: a nationwide cross-sectional survey study from USA. *Front Pediatr.* 2023;11:1149125.
- Shmueli L. Parents' intention to vaccinate their 5- to 11-year-old children with the COVID-19 vaccine: rates, predictors and the role of incentives. *BMC public health.* 2023;23:328.
- Yousaf AR, Kunkel A, Abrams JY, et al. COVID-19 vaccine reactogenicity and vaccine attitudes among children and parents/guardians after multisystem inflammatory syndrome in children or COVID-19 hospitalization: September 2021–May 2022. *Pediatr Infect Dis J.* 2023;42:252–259.