

Which children and young people are at higher risk of severe disease and death after SARS-CoV-2 infection: a systematic review and individual patient meta-analysis

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Systematic Review

Keywords: child, adolescent, COVID-19, SARS-CoV-2, severity, death, mortality

Posted Date: July 6th, 2021

DOI: <https://doi.org/10.21203/rs.3.rs-689808/v1>

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Abstract

Background We aimed to use individual patient data to describe pre-existing factors associated with severe disease, primarily admission to critical care, and death secondary to SARS-CoV-2 infection in children and young people (CYP) in hospital.

Methods We searched Pubmed, European PMC, Medline and Embase for case series and cohort studies that included all CYP admitted to hospital with 30 CYP with SARS-CoV-2 or 5 CYP with PIMS-TS or MIS-C. Eligible studies contained 1) details of age, sex, ethnicity or co-morbidities, and 2) an outcome which included admission to critical care, mechanical invasive ventilation, cardiovascular support, or death. Studies reporting outcomes in more restricted grouping of co-morbidities were eligible for narrative review. Authors of eligible studies were approached for individual patient data (IPD). We used random effects meta-analyses for aggregate study-level data and multilevel mixed effect models for IPD data to examine risk factors (age, sex, comorbidities) associated with admission to critical care and death. Data shown are odds ratios and 95% confidence intervals (CI).

Findings 81 studies were included, 57 in the meta-analysis (of which 22 provided IPD) and 26 in the narrative synthesis. Most studies had an element of bias in their design or reporting. Sex was not associated with critical care or death. Compared with CYP aged 1-4 years, infants had increased odds of admission to critical care (OR 1.63 (95% CI 1.40-1.90)) and death (OR 2.08 (1.57-2.86)). Odds of death were increased amongst CYP over 10 years (10-14 years OR 2.15 (1.54-2.98); >14 years OR 2.15 (1.61-2.88)).

Number of comorbid conditions was associated with increased odds of admission to critical care and death for COVID-19 in a dose-related fashion. For critical care admission odds ratios were: 1 comorbidity 1.49 (1.45-1.53); 2 comorbidities 2.58 (2.41-2.75); ≥ 3 comorbidities 2.97 (2.04-4.32), and for death: 1 comorbidity 2.15 (1.98-2.34); 2 comorbidities 4.63 (4.54-4.74); ≥ 3 co-morbidities 4.98 (3.78-6.65). Odds of admission to critical care were increased for all co-morbidities apart from asthma (0.92 (0.91-0.94)) and malignancy (0.85 (0.17-4.21)) with an increased odds of death in all co-morbidities considered apart from asthma. Neurological and cardiac comorbidities were associated with the greatest increase in odds of severe disease or death. Obesity increased the odds of severe disease and death independently of other comorbidities.

Interpretation Hospitalised CYP at greatest vulnerability of severe disease or death from SARS-CoV-2 infection are infants, teenagers, those with cardiac or neurological conditions, or 2 or more comorbid conditions, and those who are obese. These groups should be considered higher priority for vaccination and for protective shielding when appropriate. Whilst odds ratios were high, the absolute increase in risk for most comorbidities was small compared to children without underlying conditions.

Introduction

Children and young people (CYP) have suffered fewer direct effects of the COVID-19 pandemic than adults, as the vast majority of CYP experience few if any symptoms of SARS-CoV-2 infection.(1) However a small minority experience more severe disease (2) and small numbers of deaths have been documented.(3) As severe outcomes amongst CYP are uncommon, our understanding of which are at risk from SARS-CoV-2 is limited, in contrast to adults. Yet identification of CYP at the highest risk from infection and its sequelae is essential for guiding clinicians, families and policymakers about who can safely attend school, and for identifying groups to be prioritised for vaccination.

SARS-CoV-2 infection in CYP has two primary manifestations. The first is acute COVID-19 disease, an acute illness caused by current infection with the virus and often characterised by respiratory symptoms. The second is a delayed inflammatory condition referred to as Paediatric Inflammatory Multisystem Syndrome Temporally associated with SARS-CoV-2 (PIMS-TS) or Multisystem Inflammatory Syndrome in Children (MIS-C)(4-6). Postulated risk factors for developing more severe COVID-19 or PIMS-TS / MIS-C include existing co-morbid conditions, age, sex, ethnicity, socio-economic group, and geographical location (7-10). Existing systematic evaluations are not useful for guiding policy as reviews were undertaken early in the pandemic (11-13), included highly heterogeneous groups and a wide range of outcomes from very small studies(14), and failed to distinguish between acute COVID-19 and PIMS-TS/MIS-C. Rapid growth in the literature over the past year provides an opportunity to synthesize findings, and better inform policy decisions about vaccination and protective shielding of vulnerable CYP.

We undertook a high-quality systematic review and meta-analysis of the literature from the first pandemic year to identify which CYP were at increased risk of severe disease or death in CYP admitted to hospital with SARS-CoV-2 infection or PIMS-TS / MIS-C. The study was limited to hospitalised CYP to enable the baseline denominator characteristics to be more accurately defined, particularly co-morbidities, and because in itself, hospital admission may be an indicator of severity. We limited our review to pre-specified potential risk-factors (co-morbidities, age, sex, ethnicity and deprivation), plus a limited number of outcomes denoting severe disease (critical care admission, need for mechanical invasive ventilation or cardiovascular support) and death. These are relevant to policy on protective shielding and school attendance, and potential vaccination strategies for CYP.

Methods

The protocol for this systematic review and meta-analysis was published on PROSPERO (CRD42021235338) on the 5th February, 2021. We report findings according to the PRISMA 2020 guidelines(15) (Supplementary information 1).

Search

We performed a systematic search of four major databases: PubMed, European PMC, Scopus and Embase for relevant studies on COVID-19 in children and young people up to 21 years of age, published between the 1st January 2020 and the 29th January 2021 and updated the search on the 21st May 2021. Searches were limited to English only and included key search concepts relating to COVID-19 OR SARS-CoV-2 OR PIMS-TS OR MIS-C AND Child OR Young person OR neonate (full search strategy in supplementary information 1). References of published systematic reviews and included studies were checked for additional studies.

Study Selection

Two reviewers selected studies using a two-stage process. All titles and abstracts were reviewed independently in duplicate by a team of five reviewers to determine eligibility. Full texts of articles were reviewed if inclusion was not clear in the abstract. Disagreements were discussed between the two reviewers and a decision made about inclusion or exclusion of the study. We excluded studies if the data were duplicated elsewhere, as reported by the study authors, and prioritised the studies which gave comparative data on the risk factors and outcomes of interest; if both did so, we used the larger study.

Inclusion criteria were as follows:

1. Observational studies of any type of CYP under 21 years of age who had been admitted to hospital with a finding of SARS-CoV-2 infection at or during admission by reverse transcriptase polymerase chain reaction (RT-PCR) *OR* who had been identified clinically as having PIMS-TS or MIS-C. Studies were eligible from institutions or from defined populations.
2. Data were provided on any of the following potential risk factors: age, sex, ethnicity, co-morbidity and socioeconomic deprivation.
3. Studies that included all admitted CYP in a population or institution regardless of co-morbidity were eligible for inclusion in the meta-analysis if they included 30 children with COVID-19 or 5 children with PIMS-TS or MIS-C. 30 children with COVID-19 was selected as the minimum a-priori to account for the outcomes of admission to critical care and death being rare, with previous systematic reviews suggesting severe COVID-19 occurs in approximately 2.5% of children(16). Studies of a single pre-existing co-morbidity were included in the systematic review if they included 5 children but not included in the meta-analysis.
4. Studies which reported one of the following outcomes as a proxy for severe disease:
 - (1) Need for invasive ventilation during hospital stay (not including during anaesthesia for surgical procedures)
 - (2) Need for cardiovascular support (vasopressors, inotropes +/- ECMO)
 - (3) Need for critical/intensive care
 - (4) Death after diagnosis of SARS-CoV-2 infection or PIMS-TS/MIS-C.

We initially intended to include other identifiers indicative of severe disease including use of pharmacological therapy and length of stay in critical care, but discarded these during the search as they were rarely and inconsistently reported. In analyses, CYP who did not have an indicator of severe disease but had COVID-19 or PIMS-TS/MIS-C and were admitted to hospital were used as the comparator group.

Data Extraction for meta-analysis and Study Quality Assessment

Data on risk factors and outcome variables were extracted from individual studies by one reviewer using a pre-designed data collection form and extraction was cross-checked by a second reviewer in 10% of studies. Authors of studies from the first search (to January 2021) were contacted by email and asked to provide either additional aggregated data demonstrating the relationship between predictor and outcome variables or IPD. Time did not allow these to be requested for studies identified in the second search (to May 2021). IPD were shared by authors using a standardised data collection form and checked for consistency with the original publication. Any queries from sharing authors or the study team were discussed over email or by a virtual video call. Eligible studies not supplying individual patient data in a way that enabled the relationship between risk factors and outcomes to be analysed or that did not provide aggregate or individual patient data were excluded from the meta-analysis.

Assessment of Bias

We assessed the studies for bias using the Newcastle-Ottawa Scale (17) to assess the quality of observational studies. Studies were scored according to selection of participants, comparability, and outcome. The description of comparator cohorts was deemed present when analyses comparing two groups of outcomes were described within the publication.

Analysis

Meta-analyses were undertaken separately for COVID-19 and PIMS-TS/MIS-C to examine the association of each clinical outcome with sex (female sex was the reference group), age-group (1-4 years as reference group) and comorbidities (children without any comorbidity were the reference group). Children who were RT-PCR positive for SARS-CoV-2 but met the criteria for PIMS-TS or MIS-C were included in the latter group.

Meta-analyses were conducted in two ways. First, we undertook a random-effects meta-analysis of reported study-level data using RevMan 5 software (18) to estimate pooled odds-ratios for each outcome (death, intensive care admission, mechanical invasive ventilation and cardiovascular support). We refer to this analysis as the aggregate meta-analysis. Age categories were described as <1 year, 1-4 years, 5-9 years, 10-14 years and over 15 years and age 1-4 years used as the reference group. When studies reported a different age grouping, the group was used in the range which had the greatest cross-over of years. Co-morbidity data were compared using the presence and absence of individual co-morbidities. We calculated the I^2 statistic as a measure of heterogeneity and report prediction intervals. Funnel plots were examined to assess the evidence for publication bias. We then performed a sensitivity analysis by excluding the largest study of patients with COVID-19. The second set of meta-analyses were undertaken on the IPD, using multi-level logistic mixed-effects models in Stata 16 (StataCorp. College Station, TX) including a random effect for study, with models for co-morbidities adjusted for age and sex. After each model we calculated the predicted probability for each outcome amongst those with and without each comorbidity using the *margins* post estimation command. We did this to estimate risk difference for admission to critical care or death among CYP with comorbidities compared to those without. As a sensitivity analysis, a two-stage meta-analysis was conducted using study-level estimates calculated from the IPD data. A further sensitivity analysis for both the aggregate and IPD meta-analyses was performed by excluding one very large study.(19) Eligible studies which included only CYP with specific comorbidities were not included in the meta-analyses but included in a narrative synthesis. Data displayed are odds ratio (95% confidence interval) and absolute risk difference (95% confidence interval).

Results

Figure 1 shows the search flow. 23,050 reports were identified. After excluding duplicates and ineligible studies, 81 studies were included in the review. 57 studies were included in the meta-analysis, including a total of 21,412 children (see Table 1). Nine studies were from Asia, fifteen from Europe, one from Africa, twenty-one from North America and eight from South America. One study had global recruitment.

Data from 22 studies (40% of those in the meta-analysis) was included in the IPD meta-analyses, totalling 10,022 children. 26 studies reporting individual co-morbidities were eligible for inclusion in the narrative synthesis. Most studies eligible for inclusion in the meta-analysis were at considerable risk of bias (Figure 2).

We discuss findings from the aggregate and IPD meta-analyses for each set of risk factors and clinical outcomes below. Detailed data from included studies and pooled estimates from the aggregate meta-analyses are provided in Supplementary Table 1. Supplementary figures 1 and 2 show the sensitivity analysis with the largest study excluded. A two-stage meta-analysis using study-level estimates calculated from the IPD data is shown in supplementary figures 3 and 4.

Proportions of hospitalised children with COVID-19 admitted to critical care and who died in the aggregate analysis were 21.8% and 5.9% respectively and for PIMS-TS/MIS-C were 60.4% and 5.2%. In the IPD analysis, the proportion admitted to critical care with COVID-19 was 16.5% (6.7, 26.3) with death reported in 2.1% (-0.1, 4.3). For PIMS-TS/MIS-C, 72.6% (54.4, 90.7) were admitted to critical care and 7.41% (4.0, 10.8) died.

Demographic risk factors for admission to critical care and death

Sex was not associated with pooled risk of admission to critical care or death in either COVID-19 or PIMS-TS in either the aggregate or IPD analyses (Figure 3). Compared with 1-4 year olds, the aggregate analysis found a higher pooled risk of critical care admission amongst 10-14 year olds and a higher risk of death amongst infants for COVID-19. In contrast, the IPD analysis found higher risk of critical care and death amongst both infants and 10-14 year olds, plus a higher odds of death amongst those >14 years for COVID-19. For PIMS-TS/MIS-C, the aggregate analysis found higher odds of

critical care admission in all age-groups over 5 years, but no age-effects on risk of death. Numbers in the IPD analysis for PIMS-TS/MIS-C were very small, with no association of age-group with risk of death or critical care admission.

We were unable to assess the impact of ethnicity and socioeconomic position on clinical outcomes. The reporting of ethnicity data was highly variable and groupings were insufficiently similar across studies to allow meta-analysis. Socioeconomic position was reported by very few studies.

Association of co-morbidities and critical care and death in aggregate meta-analysis

The aggregate meta-analysis compared those with any or specific comorbidities with all other children in each study (Figure 4). The presence of any comorbidity increased odds of critical care and death in COVID-19, with pooled odds ratios of 2.56 (1.77, 3.71) for critical care and 4.16 (1.97, 8.80) for death, both with moderate to high heterogeneity. Pooled odds ratios for PIMS-TS/MIS-C were of a similar order but with wide confidence intervals.

Pooled odds of both critical care admission and death in COVID-19 were increased in children with the following co-morbidities: cardiovascular; gastrointestinal or hepatic; neurological; chronic kidney disease; endocrine conditions, including diabetes; and metabolic conditions, including obesity. Odds ratios for critical care ranged from 2.5 to 3.1 and for death from 2.9 to 13. The presence of asthma or trisomy 21 was not associated with either outcome, while respiratory conditions were associated with increased odds of critical care but not death. There was an increased odds of death but not of critical care admission in those with malignancy, haematological conditions and immunosuppression for non-malignant reasons.

Few individual comorbidities were associated with odds of critical care or death in PIMS-TS / MIS-C, with the exception of malignancy (OR for death 183 (2.61-12,815) and metabolic diseases including obesity (OR for critical care 1.45 (1.10-1.92)).

Association between co-morbidities and critical care and death in IPD meta-analysis

The IPD analysis compared those with each co-morbidity with children without any co-morbidity and additionally enabled analysis of risk associated with multiple comorbidities, obesity without other comorbidity, and trisomy 21 without cardiovascular disease. Figure 5 shows pooled OR for critical care and death for each comorbidity, and Figure 6 shows the risk difference estimated from the same models compared with children without comorbidities.

In IPD analysis, the presence of any comorbidity increased odds of critical care and death in COVID-19. The pooled odds ratio for admission to critical care was 1.64 (1.59-1.69), with risk difference being 4.6% (2.5, 6.7) greater than the 16.2% prevalence of critical care admission in those without comorbidities. The pooled odds of death from COVID-19 in those with any comorbidity was 2.49 (2.34-2.66), with a risk difference of 2.1% (-0.03, 4.2) above the 1.69% risk in those without comorbidity. For PIMS-TS/MIS-C, pooled odds of critical care was 12.44 (9.74-15.87) and risk difference 21.1% (4.4, 37.8) above baseline risk of 74.5%, and pooled odds of death was 11.23 (0.77, 163.22) with risk difference 21.0% (-3.4, 45.3) above baseline risk of death of 3.1%.

Number of comorbidities increased the odds of critical care and death in COVID-19 in an apparently dose-dependent fashion, with those with ≥ 3 comorbidities having a odds ratio of death of 4.98 (3.78, 6.56), twice that of the odds with one comorbidity. Small numbers with PIMS-TS / MIS-C meant that further analysis of co-morbidities could not be undertaken.

All individual comorbidities increased odds of admission to critical care except for malignancy and asthma, the latter associated with reduced odds (-0.6% (-1.0, -0.2)). Risk differences for critical care above the risk for the no comorbidities group were highest for cardiovascular, neurological, and gastrointestinal conditions, as well as for obesity. Obesity alone, without other conditions, increased risk difference to the same level as cardiovascular or neurological conditions, although numbers were small in the obesity analyses.

Odds of death in COVID-19 in the IPD analyses was elevated in all comorbidity groups except for asthma, where there was a reduced risk (-0.6% (-0.9, -0.3)). Risk difference additional to the no comorbidity group was highest for malignancy. Trisomy 21 increased risk of death in those with or without comorbid cardiovascular disease.

Narrative findings from studies of specific comorbidities

Twenty-six papers met the inclusion criteria for the narrative synthesis (Table 2), all reporting on the association of co-morbidity with acute COVID-19. Malignancy was the focus of sixteen of the studies, with rates of critical care admission in hospitalised patients ranging from 0-45% and of death in 0-47%. Six of the ten studies reporting deaths in this group of patients noted that some or all of the reported deaths were due to the underlying condition rather than SARS-CoV-2 infection.

Two studies focussed on hospitalised patients with sickle cell disease. There were fewer than fifteen patients in each study, with 17% of patients being admitted to critical care in one study and reported deaths in 0-10%. Two studies looking at non-malignant immunosuppression described no children requiring critical care admission or death and a study of children with Rheumatic diseases found a rate of critical care admission of 38%.

Chronic kidney disease was examined in two studies with small numbers of hospitalised patients, which describe a rate of critical care admission between 0-9% and of death between 0-6%. A study of CYP with cystic fibrosis found that 1 in 24 (4%) of those hospitalised were admitted to critical care and no deaths were described. Finally, two studies describe the association between pre-existing cardiac co-morbidity and outcome, which show a high proportion of children are admitted to critical care (43-71%) and that 14-29% are reported to die.

Discussion

We present the first individual patient meta-analysis of risk factors for severe disease and death in CYP from both COVID-19 and PIMS-TS/MIS-C, nested within a broad systematic review and meta-analysis of published studies from the first pandemic year. Studies were of mixed quality and most were open to substantial bias; yet our meta-analyses included data from 57 studies from 19 countries, including 8 low or middle-income countries (LMIC).

Across both the aggregate and IPD analyses, we found no association between sex and odds of severe disease or death for either COVID-19 or PIMS-TS/MIS-C. We found that odds of poor outcomes was 1.6 to 2-fold higher for infants than 1-4 year olds for COVID-19 alone, but that teenagers had elevated odds of severe COVID-19 (1.4 to 2.2-fold higher odds) and particularly PIMS-TS/MIS-C (2.5 to 8-fold greater odds).

The presence of underlying comorbid conditions were the strongest risk factors for critical care admission and death. The presence of any comorbidity increased odds of poor outcomes in COVID-19 for both the aggregate and IPD analyses, increasing absolute risk of critical care admission by 4.5% (a relative increase of 28%) and risk of death by 2.5% (125% relative increase), with an even greater 21% increase in risk of death for PIMS-TS/MIS-C (6.8-fold increase in risk). Whilst one comorbidity increased absolute risk of critical care by 3.6% and death by 1.5% in COVID-19, 2 or more comorbidities dramatically increased the absolute risk increase.

All comorbidities were associated with increased risk across the two analyses, with the exception of asthma. Increase in odds of poor outcomes in COVID-19 was highest amongst those with cardiovascular, respiratory, neurological, and gastrointestinal comorbidities, each increasing absolute risk of critical care by 8-11% and risk of death by 1-3.5%. Malignancy was associated with increased of death from COVID-19, but not critical care admission in both analyses, which is counter-intuitive and raises the possibility that this reflects the high mortality rate amongst cancer survivors who may have incidentally died with SARS-CoV-2 positivity. The aggregated analysis did not suggest increased risk in those with immunosuppression (outside malignancy) or with haematological conditions when compared to children without those comorbidities, but these groups were at increased risk in the IPD analysis.

The risk factors we identified for more severe COVID-19, older age and the presence of comorbidities, particularly cardiovascular, respiratory, neurological, and gastrointestinal comorbidities, and obesity, are highly similar to those risk factors now well described for adults.⁽²⁰⁾ This suggests that risk factors for severe COVID-19 are consistent across the life-course, but previously not well understood in CYP because of the rarity of severe disease. Our findings relate to risk factors for severe disease rather than risk factors for infection, as we included only hospitalised children. It is likely that our findings may over-estimate risks of critical care and death for CYP in high income countries, as the mortality rate in our analyses (2.1% of children with COVID and 7.41% of those with PIMS-TS/MIS-C) are very much higher than national mortality rates reported from these settings (21-23). This likely reflects inclusion of studies from LMIC and publication bias towards more severe cases. Despite this the additional absolute risks related to all comorbidities was small compared with those without comorbidities.

Comparison with the literature

Our finding of no differences in risk by sex is contrary to a large literature showing that males are more vulnerable to severe illness and death in childhood(24, 25). Whilst male sex is a known risk factor for more severe COVID-19 in adults, this excess risk arises only after middle age (26). We found that obesity, whether alone or with other conditions, markedly increased risk of critical care admission and death in the IPD analysis. Whilst numbers with obesity were very small, our findings are consistent with adult data showing obesity to be one of the strongest risk factors for severe disease in adults(27).

Previous reviews have not provided a systematic understanding of the associations of paediatric comorbidities and severe outcomes in children. Systematic reviews which were undertaken early in the pandemic highlighted some of the challenges around identifying comorbidities which were associated with severe disease, including pooled reporting of even common conditions such as asthma(28) and a focus on individual comorbidities without a comparator group(29). Our finding that children with trisomy 21 were at increased risk of critical care admission and death has not been described before, although it is consistent with previous adult data(30). We found that this risk appears to operate both through and independently of cardiovascular anomalies, indicating that all CYP with trisomy 21 are at some increased risk of severe disease.

Strengths and limitations

We undertook a high quality systematic review and meta-analysis, including use of individual patient data to provide more detailed analysis of comorbidities, including obesity and trisomy 21. Findings were largely consistent across the aggregate and the IPD analyses, with some exceptions. Findings from a sensitivity analysis excluding the largest study and from a two-stage meta-analysis of the IPD were highly similar to those described above.

Our data are subject to a number of limitations. Twenty-two of 57 studies (39%) provided individual patient data; systematic differences between these groups may have introduced bias. There were very small numbers with PIMS-TS/MIS-C in some analyses, particularly the IPD analyses. We were unable to examine ethnicity and socioeconomic position as risk factors due to lack of data in included studies.

Included studies were highly heterogenous and from a wide range of resource settings, and it is likely that findings were influenced by differing national approaches to hospitalisation of infected children and by differences in availability and use of resources including intensive care beds. A number of East Asian countries hospitalised all children who were SARS-CoV-2 positive, regardless of symptoms, whilst other countries limited hospitalisation to symptomatic children or those with significant illness. Policies on admission to and access to critical care likely also differed between countries(31). The novel nature of PIMS-TS/MIS-C also likely influenced critical care admission thresholds for this condition. Definitions of comorbidities were also heterogenous across studies and some of our comorbidity groups may be subject to misclassification bias. The definition of obesity in most studies related to severe or extreme obesity rather than the more common condition of being overweight, yet obesity was undefined in a number of studies.

We were unable to separate the increased risk for severe disease related to comorbidities from the underlying risks of illness and death seen in these comorbidities in uninfected children, as all children in our study had SARS-CoV-2. Yet studies of infected hospitalised cohorts are essential to understand how rare congenital or acquired comorbidities may influence risk of severe disease or death from SARS-CoV-2

Whilst our review examined comorbidities as risk factors in more detail than previous studies, there were limited data on sub-types of comorbidities, e.g. whether neurological problems were epilepsy or more complex neurodisability, and on combinations of comorbidities. Our finding that cardiovascular, neurological, and gastrointestinal conditions were associated with the highest risk of poor outcome, a risk similar to having 2 or more comorbidities, may reflect that these conditions were more likely to be comorbid with others. Given the low risk to children requiring hospital admission or critical care as a direct consequence of SARS-CoV-2 infection, it is likely that a significant number of reported cases were coincidental cases of SARS-CoV-2 seropositivity reflecting population prevalence.

Conclusions And Implications

When children are admitted to hospital with SARS-CoV-2 infection, those at greatest risk of severe disease or death are teenagers, those with cardiac or neurological conditions, or 2 or more comorbid conditions, and children who are significantly obese. These groups should be considered higher priority for vaccination and for protective shielding when appropriate. Our findings suggest that established risk factors for severe disease in adults also operate in the early life-course. Whilst odd ratios for poor outcomes were increased for nearly all comorbidities in children, the absolute increase in risk for most comorbidities was small compared to children without underlying conditions. This emphasises that our findings should be understood within the broader context that risk of severe disease and death from COVID-19 and PIMS-TS/MIS-C in hospitalised children is very low compared with adults.

Our study quantifies the additional risk related to comorbidities in infected children, however it is likely that some or all of this risk relates to the underlying condition rather than SARS-CoV-2 infection. Further population-based research using comparator groups which identify the risk of severe disease due to COVID-19 and the underlying risk due to comorbidity is required to develop a safe approach to shielding and vaccination for children.

Declarations

Acknowledgements

We thank the authors who shared patient level data to enable their inclusion in this study (Supplementary Information 1) and the Royal College of Paediatrics and Child Health who assisted with data extraction.

Contributions

Study Design: RH, NT, CS, JW, C T-S, ML, MC, EW, PJD, KL, ESD, SK, LF and RMV, Literature search, identification of papers and data extraction: RH, HY, NT, CS, JW, SK and LF, Data analysis: RH, CT-S and RV, First Draft: RH, Review and editing: All authors

Conflicts of interest

All authors declare no competing interests.

Funding

No funding obtained for these analyses.

Tables

Table 1. Study characteristics of 'All comer' studies for CYP with COVID-19 and PIMS-TS or MIS-C included in meta-analyses, grouped by region of origin

Data Source: / if extracted from paper; * if individual patient data shared, ** if individual patient data shared and includes unpublished data due to ongoing data collection, \$ if individual patient data extracted from paper, @ if aggregate data shared by authors. Admission to critical care - CC, Required mechanical invasive ventilation - mIV, Required cardiovascular support - CVS. Systematic Review - SR. uk - unknown.

Study		Population		Exposure	Risk Factors used in MA	Outcomes used in MA	Comparator Group(s)	CC n(%)	Death n(%)	Data Source
Author, Date, Country	Study Design	No of admitted children	Inclusion and Exclusion criteria	Criteria for diagnosis						
Asia										
COVID-19										
Du(32), May 2020, China	Retrospective Observational	182	<16 years Admitted	RT-PCR pos	Age	mIV n=3 Death n=1	Allergic vs non-allergic patients Pneumonia vs no pneumonia	uk	1 (0.5%)	/
Qian(33), July 2020, China	Retrospective cohort	127	1month - 16 years Patients admitted to hospital	RT-PCR pos	Age, sex, comorbidity, coinfection	CC n=7 Death n=0	Critical Disease (admission to CC/need for mIV/CVS) - only admission to CC analysed.	7 (5.5%)	0	/
Kyung Sung(34), July 202, South Korea	National prospective registry	101	All ages collected, only children <19 years inc	RT-PCR pos	Age, sex, comorbidities	CC n=0 mIV n=0 Death n=0	Comparison of disease d=severity	0	0	*
Alharbi(35), Dec 2020, Saudi Arabia	Retrospective observational	65 - C-19 6 - MIS-C	<15 years Community and hospitalised	RT-PCR pos MIS-C (CDC)	Sex, comorbidity	CC n=12 mIV n=5 CVS n=8 Death n=3	Community vs hospitalised, hospitalised vs critical care,	12 (17%)	3 (4%)	/
Bayesheva(36), Dec 2020, Kazakhstan	Retrospective Observational	549	<19 years	RT-PCR pos	Comorbidity, age, sex Obesity not defined	CC n=4 mIV n=1 Death n=0	Mild, moderate and severe disease	4 (0.7%)	0	*
Qian(37), April 2021, China	Retrospective cohort	127	1 month - 16 years	RT-PCR pos	Co-morbidities	Death	Mild, moderate, severe and critical	uk	2 (1.6%)	/
PIMS-TS / MIS-C										
Almoussa(38), Oct 2020, Saudi Arabia	Retrospective case series	10	<14 years Admitted to hospital	MIS-C (CDC)	Age, sex comorbidity	CC n=9 mIV n=1 CVS n=5 Death n=2	None	9 (90%)	2 (20%)	\$
Jain(39), Aug 2020, India	Retrospective and prospective observational	23	<15 years Hospitalised	MIS-C (WHO)	Sex, age	mIV n=9 CVS n=15 Death n=1	MIS-C with shock vs MIS-C without shock	uk	1 (4%)	*
Shahbaznejad (40), Oct 2020, Iran	Retrospective cohort	10	Patients admitted to hospital	PIMS-TS	Sex, Age	CC n=9 mIV n=3 CVS n=4 Death n=1	None	9 (90%)	1 (10%)	\$
Hasan (41), Feb 2021, Qatar	Retrospective cohort	7	Patients admitted to hospital	MIS-C (WHO)	Sex, Age	CC n=5 mIV n=1	None	5 (71%)	uk	\$
Europe										
COVID-19										
Armann(42), May 2020, Germany	Prospective Observational Registry	102	<20 years	RT-PCR pos	Age, sex, comorbidities Obesity not defined	CC n=15 mIV n=6 CVS n=8 Death n=1	None	15	1	*
Bellino(43), July 2020, Italy	Routine surveillance system	511	<18 years Admitted	RT-PCR pos	Age, sex, comorbidity	CC n=18 Death n=4	Outcomes compared by age. Multivariable logistic regression comparing predictor variables and outcomes	18	4	/
Giacomet(44), Oct 2020, Italy	Retrospective observational	127	<18 years Admitted	RT-PCR pos	Sex, comorbidity, ethnicity Obesity not defined	CC n=8 mIV, n=1	Asymptomatic, mild or moderate vs severe or critical. Admission to ICU/no ICU.	8	0	*
Gazzarino(45), May 2020, Italy	Retrospective observational	168	1 day - <18 years Admitted	RT-PCR pos	Age	mIV n=2	None	uk	uk	/
Ceano-Vivas(46), May 2020, Spain	Retrospective Observational	33	<18 years Presenting to hospital	RT-PCR pos	Sex, comorbidity, age Obesity: not defined	CC n=5 mIV n=1 CVS n=1 Death n=1	Admission to hospital	5	1	*

Study		Population		Exposure	Risk Factors used in MA	Outcomes used in MA	Comparator Group(s)	CC n(%)	Death n(%)	Data Source
Storch de Gracia(47), Oct 2020, Spain	Retrospective observational	39	< 18 years requiring hospital admission. Includes patients with MIS-C. Exclusion: pre-existing oncological disease, incidental or nosocomial SARS-CoV-2	RT-PCR post or IgG antibodies	Age	CC n=14	Uncomplicated vs complicated (fluids or vasopressors, high flow nasal cannulae / non-invasive ventilation / invasive ventilation, encephalopathy).	14	uk	/
M Korkmaz(48), June 2020, Turkey	Retrospective cohort	44	<18 years All patients attending ED	RT-PCR pos	Age	CC n=2	Admission to hospital vs discharge from ED, 5 years, >5 years	2	uk	/
Yayla(49), March 2021, Turkey	Retrospective Observational	77	<18 years Admitted	RT-PCR pos or antibodies	Comorbidity	CC n=1 mIV n=1 CVS n=1 Death n=1	Asymptomatic, mild, moderate, critical/severe	1	1	/
O Swann(7), Aug 2020, UK	Prospective Observational	579	< 19 years Admitted to hospital. (Patients with MIS-C were excluded from SR)	RT-PCR pos	Age, sex, comorbidities Obesity not defined	CC n=78 Death n=6	Admission to critical care, in-hospital mortality. <i>Details about patients with MIS-C could not be extracted and were excluded.</i>	78	6	/
Gotzinger(8), June 2020, Europe	Retrospective and prospective observational	582	<19 years Admitted and community	RT-PCR pos	Sex, comorbidity, age Obesity not defined	CC n=48 mIV n=25 CVS n=19 Death n=4	Admission to CC / no CC	48 (8.2%)	4 (0.7%)	@
Moraleda(50), July 2020, Spain	Retrospective case series	31	<18 years Admitted to hospital	RT-PCR, IgM or IgG positive or clinical MIS-C	Comorbidities	Death n=1	None	20 (65%)	1 (3%)	/
PIMS-TS / MIS-C										
Whittaker(4), June 2020, UK	Retrospective cohort	58	Patients admitted to hospital <18 years	PIMS-TS	Sex, comorbidity	CC n=32 mIV n=26 CVS n=27 Death n=1	Comparison with other childhood inflammatory disorders	32 (55%)	1 (1.7%)	@
Pang(51), UK	Retrospective selected cohort	5	Patients admitted to hospital <16 years	PIMS-TS	Sex, age, comorbidity, race	CC n=4 mIV n=4	Viral polymorphisms in admitted patients with and without PIMS-TS compared to community SARS-CoV-2 individuals	4 (80%)	4 (80%)	\$
Carbajal(52), Nov 2020, France	Retrospective observational	7	Hospitalised <18 years	MIS-C (CDC)	Sex, age	CC n=7 mIV n=3 CVS n=5 Death n=0	Kawasaki disease compared to MIS-C Comparison of MIS-C (CDC) vs MIS-C (WHO) vs PIMS-TS	7 (100%)	0	\$
Alkan(53), March 2021, Turkey	Retrospective observational	36	Hospitalised <18 years	MIS-C (CDC)	Age	CC	Mild, moderate and severe MIS-C	4 (11%)	0	/
Africa										
COVID-19										
van der Zalm(54), Nov 2020, South Africa	Retrospective Observational	62	<13 years Exclusion: MIS-C	RT-PCR pos	Age	CC n=11 mIV n=4 Death n=1	Outcomes compared based on age	11 (18%)	1 (1.6%)	/
North America										
COVID-19										
CDC(55), April 2020, USA	Voluntary national reporting	147	<18 years	RT-PCR pos	Age	CC n=15	Comparison with adults	15 (10%)	uk	/
Chao(56), Aug 2020, USA	Retrospective Observational	46	1 month - <22 years Admitted	RT-PCR pos	Sex, comorbidity Obesity: BMI >30 kg/m ²	CC n=13	Admission to critical care	13 (28%)	uk	/
Desai(57), Dec 2020, USA	Retrospective Observational	293	<18 years Presenting to hospital	RT-PCR pos	Sex, comorbidity	mIV n=27	Admission to hospital Admission to critical care	28 (9.5%)	Uk	/

Study		Population		Exposure	Risk Factors used in MA	Outcomes used in MA	Comparator Group(s)	CC n(%)	Death n(%)	Data Source
Fisler(58), Dec 2020, USA	Retrospective Observational	77	<21 years Admitted	RT-PCR pos	Sex, comorbidity Obesity: BMI 95 th percentile	CC n=30	Admission to critical care	30 (39%)	1 (1.2%)	/
Kainth(59), July 2020, USA	Retrospective observational	65	<22 years Admitted Symptomatic	RT-PCR pos	Sex, age, comorbidity	CC n=23	Subcategories of healthy infants, healthy children, immunocompromised children, chronically ill children and mild, moderate or severe disease.	23 (35%)	1 (1.5%)	/
Marcello(60), Dec 2020, USA	Retrospective observational	32	All ages included, data provided on children < 19 years	RT-PCR pos	Sex, comorbidity	Death n=1	Hospitalisation and death		1 (3.1%)	*
Kim(61), Aug 2020, USA	Population surveillance database	208 (completed data)	<18 years Hospitalised	RT-PCR pos	Age	CC n=69 mIV n=12 CVS n=10 Death n=1	Outcomes compared by age.	69 (33%)	1 (0.5%)	/
Moreira(10), Jan 2021, USA	Routinely collected data	445	All data complete <20 years All patients attending ED	RT-PCR pos	Age (0-9 years, 10-19 years), Gender, Race & ethnicity, comorbidity	CC n=69 Death n=12	Admission to hospital vs discharge from ED, Death	69 (16%)	12 (2.7%)	*
Richardson(62), April 2020, USA	Prospective cohort	110	Patients admitted to hospital No age restriction (patients included <19 years)	RT-PCR pos	Sex, comorbidities, Age, Race	CC n=37 mIV n=14 CVS n=0 Death n=1	Survival vs death	37 (34%)	1 (0.9%)	*
Verma(63), Jan 2021, USA	Retrospective Observational	82	<22 years Admitted	RT-PCR pos	Age, comorbidity Obesity: BMI 30 or 95 th percentile	CC n=23 mIV n=7 Death n=0	Admission to critical care	23 (28%)	0	/
Zachariah(64), June 2020, USA	Retrospective Observational	50	<22 years Admitted	RT-PCR pos	Sex, comorbidity	mIV n=9	Non-severe vs severe	uk	uk	/
Graff(65), April 2021, USA	Retrospective Observational	85	<21 years, all patients (admitted only in MA)	RT-PCR pos	Age, sex, race, comorbidity Obesity: BMI 95 th percentile	CC n=11	Non-severe vs severe	11 (13%)	1 (1.2%)	/
Preston(66), April 2021, USA	Routinely collected data	2430	<19 years, all patients (admitted only in MA)	Coded discharge with COVID-19	Age, sex, race, comorbidity	CC n=747 mIV n=172	Non-severe vs severe	747 (31%)	uk	/
PIMS-TS / MIS-C										
Abdel-Haq(67), Jan 2021, USA	Retrospective observational	33	<18 years Hospitalised	MIS-C (CDC)	Comorbidity Obesity not defined	CC n=22	Admission to critical care	22 (67%)	Uk	/
Capone(68), June 2020, USA	Retrospective observational	33	Hospitalised <18 years	MIS-C (CDC)	Sex	Death n=0	None	26 (79%)	0	/
Crawford(69), Feb 2021, USA	Retrospective observational	5	<19 years Hospitalised	MIS-C (CDC)	Sex, comorbidity, age Obesity not defined	CC n=4 mIV n=0 CVS n=5 Death n=0	None	4 (80%)	0	\$
Dufort(70), June 2020, USA	Emergency state reporting system	99	<21 years Hospitalised	MIS-C (NYSDOH)	Age	CC n=79 mIV n=10 CVS n=61 Death n=2	Clinical features and outcomes compared by age	79 (80%)	2 (2%)	/
Riollano-Cruz(71), USA	Retrospective cohort	15	Patients admitted to hospital <21 years	MIS-C (CDC)	Sex, comorbidity, age, Race	CC n=1 mIV n=3 CVS n=1 Death n=1	None	1 (6.7%)	1 (6.7%)	*

Study		Population		Exposure	Risk Factors used in MA	Outcomes used in MA	Comparator Group(s)	CC n(%)	Death n(%)	Data Source
Rekhtman(72), Feb 2021, USA	Prospective observational	19	Hospitalised <16 years	MIS-C (CDC)	Age, race, sex	CC n=12 mIV n=5 Death n=1	COVID-19 cohort compared to MIS-C cohort (with and without mucocutaneous disease)	12 (63%)	1 (5.3%)	*
Belay(73), April 2021, USA	Standardised reporting and retrospective notes review	1816	Hospitalised <21 years	MIS-C (CDC)	Age	CC n=1009 Death n=24	Outcomes compared based on age	1009 (56%)	24 (1.3%)	/
Abrams (74), May 2021, USA	Retrospective surveillance study	1080	Hospitalised <22 years	MIS-C (CDC)	Sex, comorbidity, Age, Race Obesity either documented by physician or BMI 95 th percentile for age and sex	CC	Admission to ICU vs no ICU	648 (60%)	18 (2%)	/
South America										
COVID-19										
OY Antunez-Montes(9), Jan 2021, Latin America	Prospective cohort	96 - C-19 67 - MIS-C	18 years All patients attending ED	RT-PCR pos MIS-C (CDC)	Sex, comorbidity, age, socioeconomic status, viral co-infections	CC n=43 mIV n=23 Death n=16	Admission to hospital, admission to PICU	43 (26%)	16 (10%)	/
Araujo da Silva(75), Jan 2021, Brazil	Retrospective cohort	50 - C-19 14 - MIS-C	Patients admitted to hospital. Clinical symptoms consistent with COVID-19.	RT-PCR pos MIS-C (WHO)	Age, gender, comorbidity Obesity not defined	CC n=38	Predominant vs non-predominant respiratory symptoms	38 (59%)	1 (1.6%)	*
Sousa(19), Oct 2020, Brazil	Routinely collected dataset	6948	<20 years, admission to hospital, Severe acute respiratory infection symptoms	RT-PCR pos	Sex, comorbidities, Age Obesity not defined	CC n=1867 mIV n=755 Death n=564	Outcomes of SARS-CoV-2 with other viral illnesses including influenza.	1867 (27%)	564 (8.1%)	**
Hillesheim(76), Oct 2020, Brazil	Prospective reporting to national surveillance system	6989	<20 years Admitted Excluded if incomplete information		Age, ethnicity, sex	mIV n=610 Death n=661	Survival vs death	610 (8.7%)	661 (9.5%)	/
Bolanos-Almeida(77), Jan 2021, Colombia	Retrospective Observational	597	<18 years	RT-PCR pos	Age, Sex	CC n=17 Death n=5	Mild, moderate and severe disease and death	17 (2.8%)	5 (0.8%)	*
Cairolí(78), Aug 2020, Argentina	Retrospective Observational	578	<21 years	RT-PCR pos	Age, sex, comorbidity Obesity: not defined	CC n=3 mIV n=1 CVS n=3 Death n=1	None	3 (0.5%)	1 (0.2%)	*
Sena(79), Feb 2021, Brazil	National Registry	315	<20 years	RT-PCR pos	Age	Death n=38	Outcomes compared by age and comorbidity (hospitalised and community).		38 (5.6%)	/
PIMS-TS / MIS-C										
Torres(80), Aug 2020, Chile	Retrospective and prospective cohort	27	Patients admitted to hospital <15 years	MIS-C (CDC)	Sex	CC n=16	Ward vs critical care admission	16 (59%)	0	/
Luna-Muñoz, 2021, Peru	Retrospective cohort	10	<13 years Hospitalised	MIS-C (CDC)	Age, Sex, comorbidity	mIV n=3 Death n=0	None	uk	0	/
Clark(81), Sept 2020, Global	Retrospective observational	55	<19 years Hospitalised	MIS-C (WHO)	Age, ethnicity	CC n=27	Comparison of cardiac abnormalities	27 (49%)	2 (3.6%)	\$

Table 2. Study characteristics of comorbidity studies for CYP with COVID-19 and PIMS-TS or MIS-C. Admission to critical care - CC.

Study		Population		Exposure	Comparator Group(s)	CC n(%)	Death n(%)	Other
Author, Date, Country	Study Design	No of admitted children	Inclusion and Exclusion criteria	Criteria for diagnosis				
Cystic Fibrosis								
COVID-19								
Bain(82), Dec 2020, Europe	Retrospective and prospective registry	24	<18 years	RT-PCR pos or clinical diagnosis	None	1 (4.2%)	0	
Heart Disease								
COVID-19								
Simpson(83), July 2020, USA	Case Series	7	<20 years	RT-PCR pos	None	3 (43%)	1 (14%)	Atrioventricular Septal Defect (AVSD) (n=2) Anomalous left coronary artery from pulmonary artery (n=1) Tetralogy of fallot (n=1) Hypertrophic cardiomyopathy (n=1) Dilated cardiomyopathy (n=1) Cardiac transplant (n=1) Comorbidities: Trisomy 21 (n=3), Obesity (n=2), Diabetes (n=1), Chronic Kidney Disease (n=1), Asthma (n=1)
Esmaeeli(84), April 2021, Iran	Case Series	7	<19 years Hospitalised	RT-PCR pos	None	5 (71%)	2 (29%)	Hypoplastic Left Heart (n=1) Truncus Arteriosus (n=1) Aortic Regurgitation (n=1) Ventricular Septal Defect (n=1) AVSD (n=1) Pulmonary Atresia (n=1) Unknown (n=1)
Cancer +/- stem cell transplant								
COVID-19								
Bisogno(85), July 2020, Italy	Retrospective and prospective case series	15	<18 years	RT-PCR pos	None	0	0	
De Rojas(86), April 2020, Spain	Retrospective case series	11	<19 years	RT-PCR pos	None	0	0	Leukaemia (n=8) Lymphoma (n=1) Bone / soft tissue (n=1) Solid organ (n=1)
Ebeid(87), Dec 2020, Egypt	Prospective observational study	15	u/k	RT-PCR pos	None	0	2 (13%)	Leukaemia (n=12) Lymphoma (n=1) Other (n=2) 5 symptomatic, 10 asymptomatic Deaths not due to COVID-19
Ferrari(88), April 2020, Italy	Retrospective and prospective case series	21	<18 years	RT-PCR pos	None	u/k	0	Leukaemia (n=10) Lymphoma (n=2) Other (n=9)
Gampel(89), June 2020, USA	Retrospective observational study	11	<18 years	RT-PCR pos	None	5 (45%)	0	Inpatient and outpatient Leukaemia/Lymphoma (n=6) Solid Tumour (n=8) Haematological diagnosis (n=3) Hematopoietic stem cell transplant (n=2)
Millen(90), Nov 2020, UK	Retrospective and prospective observational study	40	<16 years	RT-PCR pos	None	3 (8%)	1 (3%)	Inpatient and outpatient Leukaemia (n=28) Lymphoma (n=2) Soft tissue tumour (n=4) Solid organ tumour (n=10) CNS tumour (n=5) Other (n=5) 11/40 (28%) nosocomial infection Death not due to COVID-19
Montoya(31), July 2020, Peru	Case Series	33	<17 years	RT-PCR pos	None	3 (9%)	7 (21%)	Inpatient and outpatient Leukaemia (n=39) Lymphoma (n=5) CNS tumour (n=5) Other (n=27) 20/33 (61%) due to nosocomial infection 4/7 (57%) deaths not due to COVID-19

Study		Population		Exposure	Comparator Group(s)	CC n(%)	Death n(%)	Other
Palomo Colli(91), Dec 2020, Mexico	Case Series	30	<18 years	RT-PCR pos	None	2 (7%)	3 (10%)	Inpatient and Outpatient Leukaemia (n=24) Other (n=14) All deaths due to underlying condition
Radhakrishna(92), Sept 2020, India	Case Series	16	<18 years	RT-PCR pos	None	1 (6%)	0	Leukaemia (n=12) Other (n=3) 15/16 (94%) nosocomial infections
Sanchez-Jara(93), Nov 2020, Mexico	Retrospective observational study	15	<16 years	RT-PCR pos	None	u/k	7 (47%)	Leukaemia (n=15)
Madhusoodhan(94), April 2020, USA	Retrospective cohort study	28	<22 years	RT-PCR pos	None	u/k	4 (14%)	Inpatient and Outpatient Leukaemia (n=61) Lymphoma (n=3) Other (n=34) No deaths solely due to COVID-19
Kebudi(95), Jan 2021, Turkey	Retrospective cross-sectional study	38	<18 years	RT-PCR pos	None	9 (24%)	1 (3%)	Inpatient and Outpatient Leukaemia (n=26) Lymphoma (n=5) Other (n=20) No deaths solely due to COVID-19
Lima(96), Nov 2020, Brazil	Retrospective cohort study	35	<19 years	RT-PCR pos	None	10 (29%)	8 (23%)	5 deaths within 30 days, 8 within 60 days
Fonseca(97), Feb 2021, Colombia	Observational retrospective study	33	<18 years	RT-PCR pos	Comparison of diagnoses and admission to CC	7 (21%)	5 (15%)	2 deaths due to COVID-19 Leukaemia (n=14, 5 admitted CC) Lymphoma (n=4, 1 admitted CC) Other (n=9, 1 admitted CC)
Vincet(98), June 2020, Spain	Retrospective case series	5	<13 years	RT-PCR pos	None	2 (40%)	1 (20%)	3/5 (60%) nosocomial infections
Haematological								
COVID-19								
Arlet(99), June 2020, France	Prospective case series	12	<15 years	RT-PCR pos	Compared by age	2 (17%)	0	Sickle Cell Disease
Telfer(100), Nov 2020, England	Prospective case series	10	<20 years	RT-PCR pos	Compared by age	uk	1 (10%)	Sickle Cell Disease
Immunosuppression								
COVID-19								
Dannan(101), Oct 2020, United Arab Emirates	Case Series	5	<13 years	RT-PCR pos	None	0	0	Common Variable Immunodeficiency (n=1) Chemotherapy (n=1) Pyruvate kinase deficiency and splenectomy (n=1) Nephrotic Syndrome on Prednisone (n=1) Systemic Lupus Erythematosis on Prednisolone and Mycophenolate (n=1)
Perez-Martinez(102), August 2020, Spain	Retrospective case series	5	<15 years	RT-PCR pos	None	0	0	Hematopoietic stem cell transplant (n=1) Leukaemia (n=1) Liver Transplant (n=1) Kidney Transplant (n=1) C-ANCA vasculitis (n=1)
Chronic Kidney Disease								
COVID-19								
Melgosa(103), May 2020, Spain	Retrospective case series	8	<18 years	RT-PCR pos	None	0	0	Inpatient and Outpatient Renal Dysplasia (n=5) Nephrotic Syndrome (n=5) Uropathy (n=2) Other (n=4)
Malaris(104), Nov 2020, Global	Retrospective and prospective observational study	68	<20 years Under Paediatric Services CKD on immunosuppression	RT-PCR pos	None	6 (9%)	4 (6%)	Inpatient and Outpatient Kidney transplantation (n=53) Nephrotic Syndrome (n=30) Other (n=30)
Rheumatic Diseases								
COVID-19								
Villacis-Nunez(105), Jan 2021, USA	Retrospective case series	8	<22 years	RT-PCR pos	Need for hospitalisation	3 (38%)	0	Juvenile Idiopathic Arthritis (n=1) Systemic Lupus Erythematosis (n=5) Other (n=2)

Study	Population	Exposure	Comparator Group(s)	CC n(%)	Death n(%)	Other
Liver Disease and transplant						
COVID-19						
Kehar(106), Feb 2021, International	Retrospective observational study	21	Community and hospitalised <21 years	RT-PCR or antibody	Native liver disease vs liver transplant recipient	2 (9.5%) 1 (4.2%) Native liver disease (n=44) Liver transplant recipient (n=47)

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