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## Correspondence

## Low COVID-19 mortality in Spanish children

We thank Bhopal and colleagues for amending their recent letter,<sup>1</sup> which initially reported incorrect data for the number of COVID-19 deaths in children aged up to 19 years in Spain. The letter now reports 0.18 COVID-19 deaths per 100 000 in children aged 0–9 years (eight deaths) and 0.37 per 100 000 in children aged 10–19 years (18 deaths), up to Feb 10, 2021. We thought that readers of *The Lancet Child & Adolescent Health* might benefit from some clarity about how this situation arose and was rectified.

The primary source used by Bhopal and colleagues noted that the data were provisional and not consolidated.<sup>2</sup> In Spain, several clinical registries have been developed since the COVID-19 pandemic began, collecting data from more than 75 Spanish hospitals on the features and deaths of children with COVID-19.<sup>3,4</sup> These include the National Registry of COVID-19 in Spanish Children of the Spanish Paediatric Association (EPICO-AEP), the registry of children with SARS-CoV-2 infection admitted to Pediatric Intensive Care Units of the Spanish Society of Paediatric Intensive Care (SECIP), and the Hemato-Oncology Registry for COVID-19, among others.

We realised that the numbers of deaths described by Bhopal and colleagues were much higher than in the registries, in which ten deaths of children younger than 18 years were reported (appendix). After the publication of the letter by Bhopal and colleagues, Spanish researchers from the different registries reported this discordance to the Spanish regional and national agencies. Paediatricians and regional and national agencies coordinated to check the information regarding deaths with SARS-CoV-2 infection in children and young people. A programming error associated with the age of deceased patients was identified and resolved. Data from the National

Network of Public Health Surveillance (RENAVE) have been verified and updated to March 24, 2021: these data report nine deaths in children aged 0–9 years and 17 deaths in children aged 10–19 years.<sup>2</sup> Therefore, mortality was 0·21 per 100 000 in children aged 0–9 years and 0·34 per 100 000 in children aged 10–19 years. In agreement with the registries, most of the children reported to have died from COVID-19 in RENAVE data had pre-existing serious comorbidities.

Some discrepancies between the clinical registries and RENAVE might still be found for several reasons. In Spain, the process of reporting is complex. 17 different autonomous regions report COVID-19 deaths and positive SARS-CoV-2 cases to RENAVE, and not all reports are updated at the same time. Also, patients with multi-inflammatory syndrome might not be considered to have COVID-19 if the RT-PCR test is negative.

This event highlights several lessons. First, preliminary or provisional surveillance data are to be used with caution or they could lead to inaccurate conclusions. As Bhopal and colleagues state, the differences between countries might be partly due to differences in reporting mechanisms, especially with such small absolute numbers. Head-to-head comparisons between countries based on inaccurate or incomplete information could lead to false assumptions, which could trigger measures that have negative consequences for children. Second, updated clinical registry data are useful to monitor the pandemic and improve the interpretation of epidemiological surveillance information. Third, mechanisms for rechecking abnormally high or low rates of cases or deaths can be useful, both for epidemiologists and for authors.

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