We previously reported results from a seroprevalence study conducted in New Orleans, Louisiana, USA, which was hit hard early in the coronavirus disease (COVID-19) pandemic (1). Baton Rouge is a large metropolitan area roughly 80 miles northwest of New Orleans; at the time of this study, it was in the second phase of reopening after a stay-at-home order. Although the seroprevalence in New Orleans (6.9%) (1) was similar to prevalence recorded in Spain (5%), São Paulo, Brazil (4.7%), and New York, USA (6.9%) (2,3; B.H. Tess, unpub. data, https://doi.org/10.1101/2020.06.29.20142331), Baton Rouge had only 3,427 more cases as of August 2, 2020 (17,093 cases), than New Orleans did by May 16, 2020 (13,666 cases) (4). This latest study estimated severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infections in the greater Baton Rouge area (Ascension, East Baton Rouge, Livingston, and West Baton Rouge Parishes), with additional information on potential workplace exposures.

The protocol was approved by the Ochsner institutional review board and was designed to enroll and test ≤2,500 participants at 13 sites throughout Baton Rouge during July 15–31. Recruitment targeted a representative sample by using a method developed by Public Democracy (https://www.publicdemocracy.io) and described elsewhere (1,5). In contrast to the New Orleans study, in which persons tested were under a stay-at-home order, Baton Rouge was in phase 2 of reopening. A randomized subset of 500,000 Baton Rouge residents were targeted with digital ads for recruitment. Of those, 3,687 volunteers were recruited and restratified according to census designations; 2,309 were invited to participate, 2,179 enrolled and completed testing, and 2,138 were included in our final analysis. A total of 38 persons were excluded because they lived in ineligible ZIP codes, and 3 withdrew consent (Appendix Figure 1, https://wwwnc.cdc.gov/EID/article/27/1/20-3808-App1.pdf). All study materials were provided in English, Spanish, and Vietnamese. Participants were offered free transportation. Research staff verbally obtained consent from participants and electronically documented consent and survey responses. We then procured blood samples and nasopharyngeal swab specimens from participants.

We used US Food and Drug Administration Emergency Use Authorization–approved tests. Real-time reverse transcription PCR of nasopharyngeal swab specimens was performed by using the Abbott m2000 RealTime system (Abbott, https://www.molecular.abbott). Qualitative IgG blood tests were performed by using the ARCHITECT i2000SR (Abbott). The IgG test meets criteria established by the Centers for Disease Control and Prevention to yield high positive predictive value, which was validated by Ochsner Health laboratory and others (6,7). Study participants who tested positive on either or both tests were
assessed as having been infected with SARS-CoV-2. Point estimates and corresponding 95% CIs for proportions of SARS-CoV-2 exposure (PCR+ or IgG+ tests), point prevalence (PCR+, IgG−), and seroprevalence (IgG+ tests regardless of PCR test result) were estimated for the Baton Rouge area by using raw and census-weighted counts. Unadjusted odds ratios with Firth correction were calculated for all variables.

The sample was 63.6% female and 66.9% white; average age was 48.7 years (range 18–91) and average household size 2.84 persons. The census-weighted estimate of SARS-CoV-2 infections in the sample is 6.6% (6.0%, raw), with 3.0% positive for active viral shedding without detectable antibody, which translates to 16,536 contagious persons. By race and ethnicity, seroprevalence was highest (7.5%) in Black participants, compared with White non-Hispanic (1.8%), Asian non-Hispanic (1.7%), Hispanic of any race (1.6%), and other (2.7%) participants (Table).

The point prevalence and any SARS-CoV-2 infection were mapped by ZIP codes across the greater Baton Rouge area (Appendix Figure 2). Point prevalence and all infections were highly variable by ZIP code.

Marital status was associated with prevalence ($p = 0.0005$ by $\chi^2$ test). Single persons had the highest rate of infection (9.3%), compared with rates for married or cohabitating participants (5.0%), and were 1.9 times more likely to test positive (Figure). Work environment also affected prevalence ($p = 0.01$ by $\chi^2$ test); the lowest prevalence was in participants who worked from home part-time and went to a workplace part-time (3.7%). Those who worked primarily outside the home had the highest prevalence (8.2%) and were 2.3 times more likely to test positive than those who worked from home at least part-time. Infection rates varied by occupation ($p = 0.01$ by $\chi^2$ test); the lowest positivity was in office workers (3.0%) and increased odds of testing positive occurred in delivery, healthcare, and other public-facing jobs. However, based on seroprevalence, which also varied substantially by occupation ($p = 0.03$ by $\chi^2$ test), healthcare workers and public-facing workers bore the brunt of early infections, as demonstrated by higher odds of testing positive for antibodies (Figure).

We found the prevalence of SARS-CoV-2 infection in Baton Rouge to be 6.6% but with a heavy concentration of new, contagious infections (3.0%). Persons who were infected early possibly no longer had antibodies. This finding differed from our New Orleans study, which was performed after extensive lockdowns and estimated new infections at 0.9% (1). Some populations had higher rates of infection than others, including Black and Hispanic communities and public-facing workers or those who do not work from home.

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**Table. Prevalence of past and present severe acute respiratory syndrome coronavirus 2 infections by race and ethnicity across Baton Rouge, Louisiana, after phased reopening, July 2020**

<table>
<thead>
<tr>
<th>Race or ethnicity</th>
<th>Positive no./total no. (% of sample)</th>
<th>Residents &gt;18 y. no. (% of population)*</th>
<th>Any infection, raw,† % (95% CI)</th>
<th>Any infection, weighted,‡ % (95% CI)</th>
<th>Weighted point prevalence,§ % (95% CI)</th>
<th>Weighted seroprevalence,¶ % (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>128/2138 (100)</td>
<td>551,185 (100)</td>
<td>6.0 (5.0–7.1)</td>
<td>6.6 (5.7–7.7)</td>
<td>3.0 (2.3–3.7)</td>
<td>3.6 (2.8–4.4)</td>
</tr>
<tr>
<td>White alone</td>
<td>54/1431 (66.9)</td>
<td>332,445 (60.3)</td>
<td>3.8 (2.9–4.9)</td>
<td>4.2 (3.2–5.2)</td>
<td>2.4 (1.6–3.2)</td>
<td>1.8 (1.1–2.5)</td>
</tr>
<tr>
<td>Black or African</td>
<td>57/916 (24.1)</td>
<td>177,950 (32.3)</td>
<td>11.0 (8.5–14.1)</td>
<td>11.0 (8.5–14.1)</td>
<td>3.5 (1.9–5.1)</td>
<td>7.5 (5.2–9.8)</td>
</tr>
<tr>
<td>American alone</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Asian alone</td>
<td>2/59 (3.4)</td>
<td>13,630 (2.5)</td>
<td>3.4 (0.4–11.7)</td>
<td>3.5 (0.0–8.2)</td>
<td>1.7 (0.0–9.1)</td>
<td>1.7 (0.0–9.1)</td>
</tr>
<tr>
<td>Other#</td>
<td>1/28 (1.3)</td>
<td>7,025 (1.3)</td>
<td>3.6 (0.1–18.4)</td>
<td>2.7 (0.0–8.7)</td>
<td>0.0</td>
<td>2.7 (0.0–8.7)</td>
</tr>
<tr>
<td>Hispanic or Latino, any race</td>
<td>14/104 (4.9)</td>
<td>20,125 (3.7)</td>
<td>13.5 (7.6–21.6)</td>
<td>11.8 (5.6–18.0)</td>
<td>10.1 (4.3–15.9)</td>
<td>1.6 (0.0–4.0)</td>
</tr>
</tbody>
</table>

*Race or ethnicity is measured as single persons, married persons, or cohabitation (2.7%) participants (Table).

†Percentage of sample with a PCR-positive test, an IgG-positive test, or both.

‡Census-weighted percentage of PCR-positive test, IgG-positive test, or both calculated to match 2018 racial demographics by parish and combined.

§Census-weighted percentage of PCR-positive and IgG-negative tests calculated to match 2018 racial demographics by parish and combined.

¶Census-weighted percentage of IgG-positive tests calculated to match 2018 racial demographics by parish and combined.

#Other includes American Indian or Alaska Native, Pacific Islander, and multiracial persons.
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About the Author

Dr. Feehan is a research scientist at the Ochsner Clinic Foundation’s Infectious Disease Clinical Research Department. Her research focuses on the gut microbiome as a treatment modality for neurologic disease, but more immediately on the COVID-19 pandemic.

References

Despite robust research, knowledge about coronavirus disease (COVID-19) spread and effective control measures is still limited. Until recently, research has indicated that children rarely spread the infection to adults and are not the primary drivers of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) transmission (1).

We describe characteristics of the cluster of SARS-CoV-2 cases that emerged in a single nursery in Poland within 2 weeks of its reopening. We anonymized all data and collected no sensitive data. The Bioethics Committee of the Medical University of Warsaw approved the study protocol.

The nursery at issue was reopened after a nationwide lockdown on May 18, 2020. On May 31, a nursery worker reported family contact with a symptomatic SARS-CoV-2–infected person, and the nursery was closed. During the 14 days the nursery was open, a mean of 25 children attended the nursery daily. Children spent ≈8 hours there, divided into 3 groups, each cared for by 2 caregivers (Appendix, https://wwwnc.cdc.gov/EID/article/27/1/20-3849-App1.pdf). Neither children nor caregivers moved across multiple classes. Caregivers wore facemasks when in contact with children. Parents did not enter the building when dropping off and picking up children. Contacts between parents and nursery workers lasted <15 minutes, with facemasks on. Family members of different children did not mix.

The index case of SARS-CoV-2 infection (in a nursery worker with family contact) was confirmed on June 4. Subsequent PCR testing of nursery staff, children attending the facility, and family members (2 initial case-patients plus 104 other persons) (Appendix) revealed positive results in an additional 4 nursery workers (of whom 1 was also a parent of a child attending the facility), 3 children of the nursery workers, 8 children attending the facility, 3 siblings of those children, 8 parents, and 1 grandparent. The cluster involved a total of 29 persons; 8 were children attending the nursery, and 12 were children’s family members who did not enter the facility (Table). One child with a negative result had 2 parents with positive results. One child’s parent tested negative in this cluster but had tested positive within the previous 2 weeks, involved in another cluster.

The overall positivity rate in our cluster was 27%. COVID-19 prevalence in Poland is low. The number of tests conducted in the country was 124,194 in June, whereas the number of positive cases was 1,374, which corresponded to a positivity rate of 1% (2). Thus, local SARS-CoV-2 circulation in society is not sufficient to explain the positivity rate in our cluster.

We report a cluster of surprisingly high spread of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) associated with a single nursery in Poland. Our findings contrast with the presumed negligible role of children in driving the SARS-CoV-2 pandemic. Children 1–2 years of age might be effective SARS-CoV-2 spreaders.

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SARS-CoV-2 Cluster in Nursery, Poland

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We report a cluster of surprisingly high spread of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) associated with a single nursery in Poland. Our findings contrast with the presumed negligible role of children in driving the SARS-CoV-2 pandemic. Children 1–2 years of age might be effective SARS-CoV-2 spreaders.