Seroprevalence of SARS-CoV-2 in the West Bank region of Palestine: a cross-sectional seroepidemiological study

Nouar Qutob, Faisal Awartani, Zaidoun Salah, Mohammad Asia, Imad Abu Khader, Khaled Herzallah, Nadeen Balqis, Husam Sallam

ABSTRACT

Objectives Seroprevalence rates are important indicators to the epidemiology of COVID-19 and the extent of the pandemic given the existence of asymptomatic cases. The purpose of this study is to assess the seroprevalence rate in the Palestinian population residing in the West Bank.

Setting The study involved 1355 participants from 11 governorates, including 112 localities in the West Bank and 116 individuals visiting Palestinian medical laboratories.

Participants Blood samples were collected between 15th June 2020 and 30th June 2020 from 1355 individuals from randomly selected households in the West Bank, in addition to 1136 individuals visiting Palestinian medical laboratories between the 1st May 2020 and 9th July 2020 for a routine check-up.

Primary and secondary outcome measures Out of the 2491 blood samples collected, serological tests for 2455 adequate serum samples were done using an immunoassay for qualitative detection of antibodies against SARS-CoV-2. Seroprevalence was estimated as the proportion of individuals who had a positive result in the total SARS-CoV-2 antibodies in the immunoassay.

Results The random sample of Palestinians living in the West Bank yielded 0% seroprevalence with 95% and an adjusted CI (0% to 0.0043%), while the lab referral samples yielded an estimated seroprevalence of 0.354% with 95% and an adjusted CI (0.001325% to 0.011566%).

Conclusions Our results indicate that as of mid-June 2020, seroprevalence in Palestine persists low and is inadequate to provide herd immunity, emphasising the need to maintain health measures to keep the outbreak under control. Population-based seroprevalence studies are to be conducted periodically to monitor the SARS-CoV-2 seroprevalence in Palestinians and inform policymakers about the efficacy of their surveillance system.

INTRODUCTION

COVID-19 is an infectious respiratory disease caused by novel coronavirus SARS-CoV-2.1 Since its emergence in Wuhan, China in December 2019,2 SARS-CoV-2 has spread rapidly around the globe. A global pandemic was declared by the WHO.3 4 and new cases and deaths are being reported daily.5

Most authorities rely on PCR testing results to estimate the number of COVID-19 cases and make up-to-date decisions.6 Thus, numbers of patients tested positive for SARS-CoV-2 through PCR testing, symptomatic patients, those admitted to hospitals or deceased from COVID-19 are updated on a daily basis. However, the data may exclude a fraction of the population with previous mild or asymptomatic COVID-19 that has not been tested by PCR. The proportion of the population who have overcome the infection without being noticed can probably be approximated by testing for antibodies against SARS-CoV-2. Antibodies may confer immunity to repeat infection and a high proportion of immune individuals can attenuate the epidemic. Measures of anti-SARS-CoV-2 seroprevalence can also be used to estimate the clinical impact of COVID-19. To this effect, several serological surveys of SARS-CoV-2 have been done worldwide.7–16

There is lack of data on the percentage of undiagnosed Palestinian population with previous mild or asymptomatic COVID-19. Prevalence of COVID-19 infections among Palestinians residing in the West Bank remains unknown and many are concerned about this uncertainty. To this end, we conducted a population cross-sectional-based seroepidemiological study to assess the spread of SARS-CoV-2 throughout the West Bank. The study included 2491 individuals, designed to
The study also included 1136 participants from Medicare medical laboratories network referrals between 1st May 2020 and 9th July 2020 in 16 branches in the West Bank (online supplemental table 2).

Participants donated a blood sample for antibodies detection. Blood samples were centrifuged, and serum was separated, labelled and stored at −20°C at AAUP laboratory until it was used.

**Detection of antibodies**

Serological tests for 2455 adequate serum samples were done using an immunoassay for the qualitative detection of antibodies against SARS-CoV-2 (Elecsys Anti-SARS-CoV-2) in human serum by using the Cobas Analyzer cobas e 411 (Roche). The assay uses a recombinant protein representing the nucleocapsid (N) antigen for the determination of antibodies against SARS-CoV-2 with testing time of 18 min. The assay was reported to have a specificity of 100%, while the overall sensitivity was 83.87%, rising to 87.0(652,587),(761,589)% at 14 days after onset of symptoms, 87.7% 21 days after and 100% more than 40 days after.18 We included six samples from recovered cases with detected SARS-CoV-2 antibodies as a positive control.

**Statistical analysis**

We estimated seroprevalence as the proportion of individuals who had a positive result in the total SARS-CoV-2 antibodies in the immunoassay.

We used Wilson Method With Continuity Correction and Boundary Truncation (WCCBT) to construct 95% CI for the population parameter of seroprevalence.19

We calculated the adjusted CI. Since the specificity of the kit used for the antibodies test is 100% and 85%, respectively,18 we adjusted the CI for seroprevalence according to the following transformation: adjusted lower confidence limit=\( \text{max}[0, \frac{a−(1−sp)}/(sp+se−1)] \) and adjusted upper confidence limit=\( \text{min}[1, \frac{b−(1−sp)}/(sp+se+1)] \).20 The low representation in women represented in the random sample required reweighting by gender. As for the Medicare labs sample, the obtained estimates from the lab data are considered to be unbiased and did not require reweighting.

**Patient and public involvement**

The study involved 1355 participants from 11 governorates, including 112 localities, and included 1136 participants from Medicare laboratory referrals. The development of the research question and outcome measures was based on the public priorities to know the seroprevalence in the Palestinian community. Participants were notified of the results of the tests.

**RESULTS**

Of the 1395 eligible individuals residing in households selected using three-stage cluster sampling, 1355 participants agreed to participate in the study. The proportion of women was lower compared with men (137 and 1218, 137 and 1218 being representative by cities (1355 from randomly selected households and 1136 from laboratory referrals). Elecsys Anti-SARS-CoV-2 testing was done on 2455 adequate serum samples. Here, we describe the study design and the results of the first wave of the study.

**METHODS**

**Study design and participants**

The study conducted is a cross-sectional serological testing study aimed to investigate seropositivity for SARS-CoV-2 in the non-institutionalised Palestinian population residing in the West Bank.

The study involved 1355 participants from 11 governorates, including 112 localities (online supplemental table 1). We used three-stage cluster sampling to select 1395 households. The cluster of households or census track is considered to be a geographical location that is comprised of approximately 100 households. The process for conducting cluster sampling was carried as follows: (1) selecting a cluster of households, (2) selecting 10 households randomly from each cluster and (3) selecting a person at random from the selected household. The clusters were selected using probability proportional to size (PPS) sampling (table 1).

To select the number of clusters within each population location: (1) we calculated the sampling interval which equals the total number of households divided by the total number of clusters needed to be selected by the sample say, for example (m). So, the sampling interval SI=N/m, where N is the total number of households; (2) selected a random number R0 between 0 and SI; (3) calculated Ri as R0+i×SI, a cluster is selected in Li if Ri belongs to the interval (Ci−1, Ci).

Field work was carried out between 15th June 2020 and 30th June 2020 by a team of registered nurses, laboratory technicians, nursing students and laboratory technician students from the Arab American University (AAUP) following standardised health protocols.17 The random sample did not include children under the age of 15 and suffered from a low representation in women.

**Table 1** The PPS sampling algorithm

<table>
<thead>
<tr>
<th>Location</th>
<th># of households in the location</th>
<th>Cumulative</th>
</tr>
</thead>
<tbody>
<tr>
<td>L1</td>
<td>X1</td>
<td>C1=X1</td>
</tr>
<tr>
<td>L2</td>
<td>X2</td>
<td>C2=X1+X2</td>
</tr>
<tr>
<td>L3</td>
<td>X3</td>
<td>C3=X1+X2+X3</td>
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</tr>
<tr>
<td>Lk-1</td>
<td>Xk-1</td>
<td>Ck-1=Xk−1+Xk+⋯+Xk−1</td>
</tr>
<tr>
<td>Lk</td>
<td>Xk</td>
<td>Ck=X1+X2+⋯+Xk</td>
</tr>
</tbody>
</table>

PPS, probability proportional to size.

Field work was carried out between 15th June 2020 and 30th June 2020 by a team of registered nurses, laboratory technicians, nursing students and laboratory technician students from the Arab American University (AAUP) following standardised health protocols.17 The random sample did not include children under the age of 15 and suffered from a low representation in women.
The presence of antibodies against SARS-CoV-2, as evidenced by tables 3 and 4, none of the tested specimens revealed nor prevailing chronic diseases (online supplemental tables 3 and 4). None of the tested specimens revealed presence of antibodies against SARS-CoV-2. A 95% CI for the population parameter of seroprevalence was (0.0 to 0.0036). A 95% CI for the population parameter of seroprevalence was (0.0 to 0.0036), and an adjusted CI of (0 to 0.0043).

The study also included 1136 participants from Medicare laboratory referrals in 16 branches. The proportion of men was lower than women (395 and 741, respectively) including 71 in age group lower than 15, 173 in age group (15–24), 297 tests in age group (25–34), 290 in age group (35–49) and 305 in age group (50+) (figure 1). The majority of the participants did not report having symptoms in the last 3 months nor prevailing chronic diseases (online supplemental tables 3 and 4). None of the tested specimens revealed symptoms and chronic diseases were unattainable. Out of the 1136 tested participants, three men, aged 38, 58, and 1 woman, aged 40, revealed antibodies against SARS-CoV-2, with 95% CI (0.0011 to 0.0096) and an adjusted CI of (0.0013 to 0.0115).

All serum samples were adequate for testing. Out of the 1136 tested participants, 4 revealed antibodies against SARS-CoV-2 with 95% CI (0.0011 to 0.0096).

DISCUSSION

To our knowledge, this is the first SARS-CoV-2 seroprevalence study in the Palestine. The findings from this seroprevalence study for SARS-CoV-2 indicate that the estimated seroprevalence of the total SARS-CoV-2 antibodies persists low as of mid of June. The random sample of Palestinians living in the West Bank yielded 0% seroprevalence with 95% and an adjusted CI of (0% to 0.0043%), while the lab referral samples yielded an estimated seroprevalence of 0.354% with 95% and an adjusted CI (0.001325% to 0.011566%).

The low SARS-CoV-2 antibody prevalence in our population can be explained by the imposed total lockdown in the first 3 months when cases were under control and only reached around 800 cases by May. Our sampling dates ranged from 15th June 2020 to 30th June 2020 in the random sample and between 1st May 2020 and 9th July 2020 in the lab referral samples. The government of Palestine extended the state of emergency and imposed total lockdown in the first 3 months. The samples were collected after this strict lockdown which was accompanied by a drop in disease prevalence. COVID-19 cases were under control. The reported cases by the Palestinian Ministry of Health at the beginning of the data collection process by mid-June were 689 cases, and the reported cases rose to 2765 cases by the end of June. However, it takes 10–14 days for antibodies against SARS-CoV-2 to go up, therefore, our results indeed reflect the situation by mid of June. Also, it was previously reported that antibodies against SARS-CoV-2 may drop or even disappear in patients with mild COVID-19, which may have led to the inability to detect antibodies in a few previously positive cases.

Seroprevalence in Palestine is very close to that identified in Jordan (0% prevalence), a neighboring country, which was explained to be due to the strict closures implemented by the Jordanian government and the eventual curtailing of infections in Jordan. Also, seroprevalence in Maine and Montana was 0.4% and 0.5%, respectively, by end of August 2020. In comparison with other countries like Spain, Italy, Japan, Los Angeles, Germany and Switzerland, the seroprevalence in Palestine is low. Similarly, the seroprevalence among Palestinians residing in the West Bank was shown to be lower than that in Israel as reported in the first conducted serological study of 1700 tests. The study suggested that 2.5% of the Israeli population have had the coronavirus. It is, however, noteworthy that a comparison with other countries may be problematic due to the timing and the stage of the pandemic which may vary, affecting the seroprevalence estimates.

A key strength of our study is the random selection of household residents collected between 15th June 2020 and 30th June 2020. The clusters were selected using PPS sampling. This technique ensures getting self-weighting sample which can be used to produce unbiased estimators for the parameters of interest. The sample was supposed to be stratified by gender and age. However,
the female response was low and samples from women were more difficult to obtain due to a cultural boundary. In fact, women were less open to welcoming a team of nurses at home for research purposes. Also, the random sample did not include children under the age of 15 to avoid anxiety due to fear of needles in the study to eliminate personal fears due to blood withdrawal. To ensure that the estimated prevalence in the random sample was unbiased, we weighted the sample by gender. Except for cultural issues resulting in the low response rate from women in the random sample, we do not believe there were any issues that would lead participants to opt out from the study.

As for the lab referral samples, it represents the population of the lab referrals between the 1st May 2020 and 9th July 2020. The seroprevalence in this sample was 4 positive cases out of the 1136 samples tested. Since the seroprevalence within the lab referral population is close to zero, we used WCCBT to construct a 95% CI for the population parameter of seroprevalence (0.0011 to 0.0096). In comparison with household samples that were collected between 15th and 30th of June, laboratory referral samples include a good representation of women and children below the age of 15 (figure 2). Also, samples were collected starting from 1st May, which may better represent the small peaks of SARS-CoV-2-reported cases in Palestine and thus explain the presence of few positive samples in comparison with null in household samples. The sample was randomly selected by the lab data managers. Although women were not represented among the 1355 adults from Palestinian households, they were represented among the 1136 individuals visiting laboratories; out of the 1136, 741, more than 50% of the participants, were women (figure 2). Hence, the obtained estimates from the lab data are considered to be unbiased.

Our study only detected SARS-CoV-2 antibodies. It was previously reported that antibodies against SARS-CoV-2 may drop or even disappear in patients with mild COVID-19, which may have led to the inability to detect antibodies in a few previously positive cases. Also, it is important to recognise that cellular immunity may play a role in providing immunity against SARS-CoV-2 reinfection. Further studies aimed at testing cellular immunity are important. It is also noteworthy that previous studies have indicated that asymptomatic individuals were reported to have a weaker immune response to SARS-CoV-2 infection and a higher percentage of asymptomatic individuals became seronegative when compared with symptomatic individuals in the early recovering phases. The reduction in neutralising antibody levels may have implications for immunity strategy and serological surveys.

In conclusion, our study provides estimates of SARS-CoV-2 seroprevalence in Palestine. Our results reflect the seroprevalence as of mid of June. It is noteworthy that on 25th May, the government of Palestine eased the restrictions following a decline in cases which led to a surge in cases beginning of July, with the epicentre of the epidemic in Hebron accounting for over 70% of active cases implying a possible increase in seroprevalence after July. Reported cases rose to 2765 cases by the end of June. By December 2020, the cumulative number of positive cases grew to 150,046 cases. In this situation, health measures have to be taken to keep the outbreak under control, in order to monitor the SARS-CoV-2 seroprevalence in Palestine and inform policymakers about the efficacy of their surveillance system. With the surge in cases, conducting population-based seroprevalence studies on a regular basis is important.

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