Combined Point-of-Care Nucleic Acid and Antibody Testing for SARS-CoV-2 following Emergence of D614G Spike Variant

Graphical Abstract

Highlights
- Combined rapid antibody + nucleic acid detection correctly diagnoses SARS-CoV-2
- Rapid antibody tests detect immune responses against SARS-CoV-2 bearing D614G
- Rapid SARS-CoV-2 antibody tests do not cross-react with antibodies to seasonal CoV
- False positivity in SARS-CoV-2 finger prick blood antibody tests can be very low

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In Brief
Mlcochova et al. report that combined rapid nucleic acid amplification testing (NAAT) and finger prick blood antibody tests can substantially improve the diagnosis of COVID-19 as compared to NAAT alone and is able to detect the SARS-CoV-2 Spike D614G variant that dominates the pandemic.
Combined Point-of-Care Nucleic Acid and Antibody Testing for SARS-CoV-2 following Emergence of D614G Spike Variant

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SUMMARY

Rapid COVID-19 diagnosis in the hospital is essential, although this is complicated by 30%–50% of nose/throat swabs being negative by SARS-CoV-2 nucleic acid amplification testing (NAAT). Furthermore, the D614G spike mutant dominates the pandemic and it is unclear how serological tests designed to detect anti-spike antibodies perform against this variant. We assess the diagnostic accuracy of combined rapid antibody point of care (POC) and nucleic acid assays for suspected COVID-19 disease due to either wild-type or the D614G spike mutant SARS-CoV-2. The overall detection rate for COVID-19 is 79.2% (95% CI 57.8–92.9) by rapid NAAT alone. The combined point of care antibody test and rapid NAAT is not affected by D614G and results in very high sensitivity for COVID-19 diagnosis with very high specificity.

INTRODUCTION

As of August 2, 2020, >18.0 million people have been infected with severe acute respiratory syndrome-coronavirus-2 (SARS-CoV-2), with >690,000 deaths. The unprecedented numbers requiring SARS-CoV-2 testing has strained healthcare systems globally. There is no gold standard for the diagnosis of coronavirus disease 2019 (COVID-19). The detection of SARS-CoV-2 by nucleic acid amplification testing (NAAT) is largely done by real-time RT-PCR on nose/throat swabs in centralized laboratories. RT-PCR specimens are often batch analyzed, and the turnaround time for this test can be as long as 2–4 days in real-world settings. NAAT tests from a single nose/throat swab are negative in up to 50% of patients who have computed tomography (CT) changes consistent with COVID-19 and/or positive antibodies to SARS-CoV-2. The lack of detectable virus in upper airway samples is not only a serious barrier to making timely and safe decisions in the emergency department but it also leads to multiple swab samples being sent, frequently from the same anatomical site, resulting in additional strain on virology
laboratories. Nonetheless, NAAT remains important in identifying infectious individuals. In addition, in severely ill patients, tracheobronchial samples may be NAAT+, even when the nose/throat swab is negative.4,6

Multiple factors may contribute to negative results by NAAT, including test sensitivity, sampling technique, and timing of the sampling in the disease course.5 The viral load in the upper respiratory tract is detectable from ~4 days before symptoms and frequently wanes after 1 week post-symptom onset.6–8 Similarly, a case series from Germany found the detection rate by RT-PCR was <50% after 5 days since onset of illness.9 A proportion of patients develop secondary deterioration in clinical condition, requiring hospitalization and respiratory support, at a time when immune pathology rather than direct pathology related to viral replication is thought to be dominant.9,10

An antibody response to SARS-CoV-2 is detectable 6 days from infection and is almost always neutralizing.12,13 Antibody-based diagnosis of COVID-19 shows increasing sensitivity in the latter part of the infection course, when NAAT on nose/throat samples is more likely to be negative.4–11 As a result, the diagnosis of infection and the identification of infectivity would benefit from a combination of virologic and immunologic markers to inform patient initial triage and subsequent management. It is critical to determine whether a rapid point of care combined antibody and nucleic acid testing strategy could improve diagnosis.

We previously evaluated the diagnostic accuracy of the SAMBA (simple amplification-based assay) II SARS-CoV-2 rapid test compared with the standard laboratory RT-PCR and found similar accuracy, with a turnaround time of 2–3 h, even in real-world settings.15 Several studies have reported head-to-head comparisons of immunochromatographic lateral flow immunoassays (LFAs).16–18 These assays are inexpensive to manufacture and provide a binary positive/negative result, thereby lending themselves well to point-of-care (POC) testing. Even though they have variable performance and in general are negative in the early phase of infection, they become highly sensitive in the later stage of illness,15–17,18 and some are also highly specific.

In this study, we evaluated the diagnostic performance of a POC combination comprising NAAT and antibody testing against a composite reference standard of laboratory RT-PCR and a serum neutralization assay. Notably, SARS-CoV-2 viruses with a D-to-G mutation in Spike at position 614 have increased in prevalence globally.20 Cryoelectron microscopy (cryo-EM) studies suggest that D614 may play a role in Spike intermolecular stability,21 potentially contributing to increased infectivity.20 Given that POC antibody tests were designed to detect antibodies to the wild-type S protein, we also aimed to investigate whether SARS-CoV-2 infections with D614G Spike mutant virus could be diagnosed by POC antibody tests.

RESULTS

In phase one, 45 prospectively recruited participants in the COV-IDx study with suspected COVID-19 disease had nose/throat swabs specimens tested for nucleic acid and stored sera for antibody testing. Samples at hospital admission were collected at a median of 7 (interquartile range [IQR] 7–13) days after illness onset. The sera from 42.2% (19/45) participants showed neutralizing antibody response against SARS-CoV-2 Spike protein pseudotyped virus infection in a neutralization assay using a cut-off of 50% inhibition at 1:4 dilution (Figure 1A). The sera of 26 participants showed no neutralizing response (Figure 1B). The neutralization ability of participants’ sera was compared with an in-house ELISA immunoglobulin G (IgG) assay for Spike-specific antibodies based on a recently reported method (Figure S1), and significant association between positive results in both assays was demonstrated (Figure 1C, p < 0.0001). Figures 1D–1G show significant associations between the POC antibody test result and both ELISA (p < 0.0001) and neutralization (p < 0.0025) assays. POC antibody testing showed no cross-reactivity in sera obtained before the pandemic (Table S1). The neutralization assay also demonstrated a lack of cross-reactivity with SARS-CoV-1 on a limited subset of sera (Figure S2).

Results from the 4 IgG antibody assays used in this study were confirmed (4 or 3 concordant) in 38/45 samples, and, against this classification, neutralization (Figures 1A–1C), spike ELISA22 (Figures 1C, 1D, 1F, and S1), Surescreen, and COVIDIX Healthcare assays gave a correct result in 100%, 97.4%, 92.1%, and 86.8% of cases, respectively, justifying the choice of the neutralization assay as part of a composite reference standard.

A total of 53.3% (24/45) of participants had COVID-19 disease, as determined by the composite reference standard (lab RT-PCR and neutralization assay). The median age of the patients was 73.5 (IQR 54.0–86.5) years in those with SARS-CoV-2 infection by our composite reference standard and 63.0 (IQR 41.0–72.0) years in those without disease (Table 1). C-reactive protein (CRP) and procalcitonin were significantly higher in confirmed COVID-19 patients, and classical chest radiograph appearances were more common in confirmed COVID-19 patients (Table 1, p < 0.001). However, 6/24 (25%) had normal or indeterminate chest radiographs in the confirmed COVID-19 group.

As expected from the clinical study inclusion criteria, >80% of patients presented with influenza-like illness (ILI) with documented fever and ~33% had clinical or radiological evidence of pneumonia (Table 1). Highly experienced internal medicine physicians were caring for suspected COVID-19 cases at our institution, and this was partly due to the significant comorbidities in the local population that mandated a broad differential diagnostic approach in hospitalized individuals (Table 1). Among the patients with COVID-19, one suffered from rheumatoid arthritis and was immunosuppressed with prednisolone. Among the patients without COVID-19, 5 were immunosuppressed for the following conditions: psoriatic arthritis (usekinumab, anti-interleukin-12 [IL-12], IL-23), multiple myeloma (lenalidomide and dexamethasone), lymphoma (cyclosporin), hypersensitivity pneumonitis (mycophenolate and prednisolone), and renal transplant (mycophenolate and tacrolimus). No patients in the study were under treatment with the anti-B cell monoclonal antibody rituximab.

During the peak of the first wave, routine respiratory virus testing was halted at our institution due to the demands of SARS-CoV-2 testing and low seasonal prevalence of these pathogens. Multiplex PCR for other respiratory viral pathogens was performed in only 8 participants. Seven of these
participants were negative and one participant tested positive for influenza A. The overall COVID-19 diagnosis rate (positive predictive agreement) by rapid nucleic acid testing was 79.2% (95% confidence interval [CI] 57.8–92.9), decreasing from 100% (95% CI 59.0%–100%) for days 1–4 to 50.0% (95% CI 11.8–88.2) for days 9–28 post-symptom onset (Table 2; Figure S3). When IgG/IgM rapid tests were combined with NAAT, the overall positive predictive agreement increased to 100% (95% CI 85.8–100) (Table 2). Additional cases of COVID-19 detected in NAAT patients were identified by POC tests under investigation (Figure 2).

Among 21 COVID-19+ patients, there were 3 false positive results for 1 POC antibody test and 1 false positive result for the other, resulting in positive predictive values of 88.9% and 96.0%, respectively, for the 2 POC antibody/SAMBA II NAAT combinations.

Those with positive NAAT and sequence available were predominantly infected with strains containing the D614G mutation in Spike, downstream of the receptor-binding domain and located on the Spike surface (Figures 3A and 3B). A total of 14/24 (58.3%) patients deemed to be COVID-19+ by the reference composite standard were positive by both rapid NAAT and antibody testing, and 14/14 were infected with strains bearing D614G, indicating that POC serological tests were able to detect infections with this variant.
Table 1. Characteristics of Participants in Diagnostic Accuracy Study

<table>
<thead>
<tr>
<th></th>
<th>COVID-19+</th>
<th>COVID-19−</th>
<th>p*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male sex (%)</td>
<td>14 (58.3)</td>
<td>9 (42.9)</td>
<td>0.30</td>
</tr>
<tr>
<td>Median age, y (IQR)</td>
<td>73.5 (54.0–86.5)</td>
<td>63.0 (41.0–72.0)</td>
<td>0.03</td>
</tr>
<tr>
<td>Influenza-like illness with documented fever</td>
<td>20 (83.3)</td>
<td>17 (81.0)</td>
<td>0.84</td>
</tr>
<tr>
<td>Clinical or radiological evidence of pneumonia</td>
<td>10 (41.7)</td>
<td>7 (33.3)</td>
<td>0.57</td>
</tr>
<tr>
<td>Immunosuppressed</td>
<td>0.053</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>1 (4.2)</td>
<td>5 (23.8)</td>
<td>–</td>
</tr>
<tr>
<td>Cardiovascular disease</td>
<td>6 (25.0)</td>
<td>2 (9.5)</td>
<td>0.25</td>
</tr>
<tr>
<td>Chronic respiratory disease</td>
<td>5 (20.8)</td>
<td>6 (28.6)</td>
<td>0.73</td>
</tr>
<tr>
<td>Chronic renal disease</td>
<td>4 (16.7)</td>
<td>2 (9.5)</td>
<td>0.67</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>6 (25.0)</td>
<td>3 (14.3)</td>
<td>0.47</td>
</tr>
<tr>
<td>Median SpO2, % (IQR)</td>
<td>95.0 (92.5–96.0)</td>
<td>96.0 (94.0–98.0)</td>
<td>0.09</td>
</tr>
<tr>
<td>Median FiO2, % (IQR)</td>
<td>0.21 (0.21–0.24)</td>
<td>0.21 (0.21–0.21)</td>
<td>0.40</td>
</tr>
<tr>
<td>Median PaO2, kPa (IQR)</td>
<td>5.0 (3.0–9.1)</td>
<td>7.2 (3.8–9.0)</td>
<td>0.30</td>
</tr>
<tr>
<td>Median PaO2: FiO2 (IQR)</td>
<td>20.5 (13.3–32.9)</td>
<td>30.9 (18.1–36.2)</td>
<td>0.09</td>
</tr>
<tr>
<td>Median respiratory rate, breaths/min (IQR)</td>
<td>22.0 (19.0–27.5)</td>
<td>20.0 (17.0–23.0)</td>
<td>0.06</td>
</tr>
<tr>
<td>Median heart rate, beats/min (IQR)</td>
<td>86.0 (77.5–99.5)</td>
<td>88.0 (78.0−107.0)</td>
<td>0.44</td>
</tr>
<tr>
<td>Median systolic BP, mmHg (IQR)</td>
<td>139.5 (117.5–149.0)</td>
<td>135.0 (119.0–152.0)</td>
<td>0.90</td>
</tr>
<tr>
<td>Median duration of illness, days (IQR)</td>
<td>7 (1–8)</td>
<td>10 (3–14)</td>
<td>0.10</td>
</tr>
<tr>
<td>Median Hb, g/dL (IQR)</td>
<td>12.9 (12.0–13.8)</td>
<td>13.1 (11.6–14.1)</td>
<td>0.46</td>
</tr>
<tr>
<td>Median WCC, × 10⁶/L (IQR)</td>
<td>7.0 (5.0–8.0)</td>
<td>9.0 (7.0–14.0)</td>
<td>0.08</td>
</tr>
<tr>
<td>Median lymphocyte count, × 10⁹/L (IQR)</td>
<td>0.8 (0.5–1.2)</td>
<td>1.2 (0.8–1.5)</td>
<td>0.12</td>
</tr>
<tr>
<td>Median platelet count, × 10⁹/L (IQR)</td>
<td>213.5 (188.5–303.5)</td>
<td>271.0 (186.0–305.0)</td>
<td>0.59</td>
</tr>
<tr>
<td>Median ferritin, µg/L (IQR)</td>
<td>684.7 (206.2–1059.1)</td>
<td>112.3 (49.6–323.6)</td>
<td>0.02</td>
</tr>
<tr>
<td>Median CRP, mg/L (IQR)</td>
<td>72.0 (28.5–214.5)</td>
<td>12 (4.0–53.0)</td>
<td>0.004</td>
</tr>
<tr>
<td>Median procalcitonin, ng/mL (IQR)</td>
<td>0.2 (0.1–0.6)</td>
<td>0.0 (0.0–0.1)</td>
<td>0.03</td>
</tr>
</tbody>
</table>

COVID-19 status is based on composite reference standard test of nose/throat swab SARS-CoV-2 RT-PCR + serum neutralization of pseudovirus-bearing SARS-CoV-2 Spike. BP, blood pressure; Hb, hemoglobin; WCC, white blood cell count.

*Wilcoxon rank-sum test used except where indicated.

*Chi-square test.

(Figures 4A and 4B). Over time, the band intensity for IgM and IgG increased along with the serum neutralization activity. In the individual presenting 21 days into illness (Figures 4G–4I), only IgG was detected with rapid POC antibody testing, and, as expected, the band intensity did not increase over the following days.

In phase 2, we performed a prospective evaluation of combined testing in 128 patients presenting with possible COVID-19 from July 13 to 27, 2020. Their clinical presentation was less severe and diagnoses broader than in phase 1 (Table 3), with cardiovascular and gastrointestinal disease significantly represented and respiratory disease representing just 60% of cases—likely as a result of the increased appreciation of diverse presentations of COVID-19 disease. Patients did have significant comorbidities and ~10% were immunosuppressed, although without B cell-depleting agents (Table 3). By this time, the POC NAAT test had been validated in a head-to-head study against the lab RT-PCR and entered routine use, replacing the RT-PCR. Given the need to further assess the specificity of the POC antibody tests in routine clinical practice and with fresh blood rather than serum, we compared the performance of POC antibody tests on finger prick blood against serum neutralization (Figures 5A and 5B).

In this second phase, there was only one NAAT positive patient, who was also positive by both POC antibody tests and serum neutralization. There were three NAAT† individuals presenting with respiratory symptoms who had positive POC antibody tests by both COVIDIX and SureScreen, along with serum neutralization activity. The POC antibody tests showed 100% negative predictive agreement with serum neutralization and the kappa correlation between POC antibody tests and serum neutralization was extremely high, at 0.97.

DISCUSSION

Here, we have shown that POC NAAT testing in combination with antibody detection can significantly improve the diagnosis of COVID-19. The overall positive predictive agreement against the composite reference standard under clinical trial conditions was ~79% for rapid NAAT testing of nose/throat swab samples, reaching 100% with a combined approach of rapid NAAT testing.
and either of the 2 POC lateral flow-based antibody tests. The specificity of the combined approach was 85%–95% on stored serum under clinical trial conditions and 100% on finger prick blood in routine clinical care. As expected, nucleic acid detection in nose/throat samples was highest in those presenting within the first few days (100% in samples taken in the first 4 days after symptom onset). Conversely, antibody detection by LFA increased with time since symptom onset, with 100% efficacy beyond the 9th day post-symptoms. One study reported that combined lab-based RT-PCR with lab-based antibody testing could increase the sensitivity for COVID-19 diagnosis from 67.1% to 99.4% in hospitalized patients. However, in that study, this assessment of sensitivity was made using clinical diagnosis. A major strength of the present study is the use of an objective reference standard that included NAAT and serum neutralization, a phenotypic test for the functionality of antibodies. This assay was shown to be robust and accurate, using a recently described ELISA method for SARS-CoV-2 IgG detection that is now used globally. The D614G Spike mutant has spread globally. Wild-type Spike protein antigen is used in the development and validation of POC antibody tests, including those tested here. Of critical importance is the fact that both POC antibody tests (and ELISA) were able to detect antibody responses in patients infected with the D614G Spike mutant and that the band intensity of POC testing increased with neutralization activity in these individuals. Given that POC antibody tests are far cheaper and simpler to deploy, they will likely be used in low-resource settings that do not have access to NAAT. Demonstration that POC antibody LFA tests can detect the D614G spike mutant is therefore of importance.

The use of antibody tests for COVID-19 diagnosis in hospitals has been limited for a number of reasons. First, we know from SARS-CoV-1 that previous humoral immunity to HCoV OC43 and 229E can elicit a cross-reactive antibody response to N of SARS-CoV-1 that previous humoral immunity to HCoV OC43 and 229E can elicit a cross-reactive antibody response to N of SARS-CoV-1 in up to 14% of people tested in cross-sectional studies, and previous exposure to HCoV can rarely elicit a cross-reactive antibody response to the N and S proteins of SARS-CoV-2. Second, antibody tests do not achieve the same detection rates as nucleic acid-based tests early in infection, as humoral responses take time to develop following viral antigenic stimulation. However, by day 6 post-symptom onset, detection of IgG to Spike protein has been reported to reach 100% sensitivity, and this is useful in cases with immune-mediated inflammatory disease in which RT-PCR on respiratory samples is often negative—for example, in the recently described Kawasaki-like syndrome called PIMS (pediatric inflammatory multisystem syndrome).

In phase one (COVIDx trial), we tested stored sera rather than whole-blood finger prick, although this was intentional, given the caution needed in interpreting LFAs and concern regarding potential cross-reactivity of antibodies and poor specificity. Although SARS-CoV-2 ELISA testing of our pre-pandemic sera did reveal occasional N reactivity to SARS-CoV-2, likely due to cross-reactivity with seasonal CoV, these samples were negative on POC antibody testing. However, the specificity of the COVIDx test was estimated at only 85%, compared to a more acceptable 95% for SureScreen. We therefore carried out a prospective evaluation of POC antibody testing on finger prick blood in 128 suspected cases of COVID-19 to further evaluate the specificity of both tests in routine clinical practice. We found no false positives in patients whose sera were non-neutralizing. This is consistent with an estimated specificity of >99% with the SureScreen assay observed in an independent analysis using stored pre-pandemic sera. The greater incidence of false positive POC antibody tests, predominantly with COVIDx, on stored sera as compared to fresh finger prick blood may be due to processing and storage of sera, contamination of sera.

Table 2. Individual and Combined Diagnostic Accuracy of POC Rapid NAAT-Based and Antibody Tests According to Time from Initial Symptoms

<table>
<thead>
<tr>
<th>Test</th>
<th>Days 1–4</th>
<th>Days 5–8</th>
<th>Days 9–28</th>
<th>Overall</th>
</tr>
</thead>
<tbody>
<tr>
<td>SAMBA II NAAT</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive agreement</td>
<td>100</td>
<td>81.8</td>
<td>50.0</td>
<td>79.2</td>
</tr>
<tr>
<td>Predictive</td>
<td>(59.0–100)</td>
<td>(48.2–11.8)</td>
<td>(67.8–88.2)</td>
<td>92.9</td>
</tr>
<tr>
<td>SAMBA II NAAT and COVIDIX IgM and IgG</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive agreement</td>
<td>100</td>
<td>90.9</td>
<td>100</td>
<td>95.8</td>
</tr>
<tr>
<td>Predictive</td>
<td>(59.0–100)</td>
<td>(58.7–99.8)</td>
<td>(54.1–100)</td>
<td>99.9</td>
</tr>
<tr>
<td>SAMBA II NAAT and SureScreen IgM and IgG</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive agreement</td>
<td>42.9</td>
<td>90.9</td>
<td>100</td>
<td>79.2</td>
</tr>
<tr>
<td>Predictive</td>
<td>(9.9–81.6)</td>
<td>(58.7–99.8)</td>
<td>(54.1–100)</td>
<td>92.9</td>
</tr>
<tr>
<td>SAMBA II NAAT and SureScreen IgM and IgG</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive agreement</td>
<td>54.1</td>
<td>99.2</td>
<td>100</td>
<td>99.9</td>
</tr>
<tr>
<td>Predictive</td>
<td>(9.4–100)</td>
<td>(9.4–99.2)</td>
<td>(9.4–100)</td>
<td>99.9</td>
</tr>
</tbody>
</table>

Positivity predictive agreement is the percentage of positive test results in samples deemed positive by the composite reference standard. Negative predictive agreement is the percentage of negative test results in samples deemed negative by the composite reference standard. *43 of 45 patients had SureScreen antibody results.

The specificity of the combined approach was 85%–95% on stored serum under clinical trial conditions and 100% on finger prick blood in routine clinical care. As expected, nucleic acid detection in nose/throat samples was highest in those presenting within the first few days (100% in samples taken in the first 4 days after symptom onset). Conversely, antibody detection by LFA increased with time since symptom onset, with 100% efficacy beyond the 9th day post-symptoms.
with other blood products, or other causes, including patient factors that differed between the two phases. Nevertheless, now that we are in a low incidence period, it is advisable to perform confirmation testing using an alternative platform for either a single positive antibody or NAAT test, as is now the policy at our institution. One should note in particular that antibody tests may be negative in patients with immunosuppression, highlighting that patient factors can influence the interpretation of results and that alternative diagnoses should be considered.

We envisage a deployment approach whereby both test samples, finger prick whole blood and nose/throat swab, are taken at the same time on admission to the hospital. The finger prick antibody test result is available within 15 min. Due to the possibility of false positive results from POC serology testing, a positive POC antibody test result as the only positive marker should ideally be confirmed with a second rapid POC test/laboratory IgG/IgM test before movement to a COVID-19 area or recruitment into a clinical trial.

**Figure 2.** Venn Diagrams Comparing Positive and Negative Diagnostic Test Results in Hospitalized Patients

Testing by NAAT and POC antibody testing by (A) COVIDIX Healthcare IgM/IgG kit (n = 45) and (B) SureScreen IgM/IgG kit (n = 43).

**Figure 3.** Spike D614G Characterization in the Phase 1 Clinical Cohort

(A) Genome map of SARS-CoV-2, with overall topography of Spike expanded. FP, fusion peptide; HR1, heptad repeat 1; HR2, heptad repeat 2; IC, intracellular domain; NTD, N-terminal domain; RBD, receptor-binding domain; TM, transmembrane region. The aligned sequence of 10 amino acids on either side of D614 is shown for 16 participants for whom sequence data were available. A dot represents where the amino acid is unchanged from wild type, the mutant glycine is represented by G.

(B) Top view of SARS-CoV-2 Spike glycoprotein trimeric structure in a closed state, with position 614 in yellow in each protomer. Structure determined by cryoelectron microscopy. RCSB PDB: 6VXX.
Figure 4. Longitudinal Antibody Responses in Patients Infected with D614G Mutant SARS-CoV-2 Detected by Rapid Lateral Flow and Neutralization Assays

(A, D, and G) An immunochromatographic lateral flow rapid diagnostic test (POC antibody test-COVIDIX SARS-CoV-2 IgM IgG test) on longitudinal samples in individual patients detecting SARS-CoV-2 IgM and IgG bands. Band intensities were acquired using the ChemiDoc MP Imaging System and quantified using Image Lab software.

(B, E, and H) SARS-CoV-2 pseudotyped virus neutralization assay from longitudinal serum samples in individual patient examples. The assays were performed in duplicate. The error bars represent SEMs.

(C, F, and I) Comparison of IgG band intensities from lateral flow rapid diagnostic test with EC50 neutralization titers from SARS-CoV-2 pseudotyped virus neutralization assay in individual patients. The correlations were estimated by linear regression analysis.
treatment study. At our institution, additional diagnostic data from chest imaging and blood tests such as lymphocyte count and CRP are considered in clinical decision making when assessing patients for COVID-19. Further swabs for NAAT testing are also taken where possible.

**Limitations of Study**

This study was limited by the fact that it was conducted at a single center with relatively small numbers of individuals in the clinical study (phase 1), largely due to a lack of available stored serum. Phase 1 of the study used stored serum, in which there was a higher false positive rate than phase 2, in which whole blood was used. The implementation study (phase 2) had greater numbers and was able to effectively demonstrate the high specificity and the very low false positive rate of the POC antibody tests on whole blood. However, it was hampered by the low incidence of COVID-19 infection during the period it was undertaken. This limited the further evaluation of the sensitivity of the combined approach. There was also a lack of data on repeated sampling and sampling from deeper respiratory sites in those suspected cases who were NAAT+. Future larger studies are warranted.

**STAR METHODS**

Detailed methods are provided in the online version of this paper and include the following:
The authors declare no competing interests.

DECLARATION OF INTERESTS

E.G.K., A.B., and H.Z.


AUTHOR CONTRIBUTIONS

The Centre for Aid Reagents (CFAR), National Institute for Biological Standards and Control (NIBSC), thanks to the donation of Dr. Emma Bentley.

pCAGGS_SARS-CoV-2_Spike was obtained by the Centre for AIDS Reagents (CFAR), National Institute for Biological Standards and Control (NIBSC), thanks to the donation of Dr. Emma Bentley.

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Supplemental Information can be found online at https://doi.org/10.1016/j.xcrm.2020.100099.

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R.K.G. is supported by a Wellcome Trust Senior Fellowship in Clinical Science and Standards and Control (NIBSC), thanks to the donation of Dr. Emma Bentley.

REFERENCES


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RESOURCE AVAILABILITY

Lead Contact
Further information should be directed to and will be fulfilled by the Lead Contact, Ravindra K. Gupta rkg20@cam.ac.uk.

Materials Availability
This study did not generate new unique reagents.
Data and Code Availability
Raw anonymised data are available from the lead contact without restriction.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Clinical Study
The study was conducted in two phases: a clinical validation phase followed by an implementation phase. The study participants in phase one were part of the COVIDx trial, a prospective analytical study which compared SAMBA II SARS-CoV-2 point of care test to the standard laboratory RT-PCR test for the detection of SARS-CoV-2 in participants admitted to Cambridge University Hospitals NHS Foundation Trust (CUH) with a possible diagnosis of COVID-19. Consecutive participants were recruited during 12-hour day shifts over a duration of 4 weeks from the 6th of April 2020 to the 2nd of May 2020. We recruited adults (> 16 years old) presenting to the emergency department or acute medical assessment unit as a possible case of COVID-19 infection. This included any adult requiring hospital admission and who was symptomatic of SARS-CoV-2 infection, demonstrated by clinical or radiological findings. 45 participants who had available stored sera were included in this sub-study and underwent further antibody testing. Phase 2, from July 13th to 27th 2020, comprised a service evaluation of clinical practice whereby adults (> 16 years old) presenting to the emergency department or acute medical assessment unit as a possible case of COVID-19 infection were included. This included any adult requiring hospital admission and who was symptomatic of SARS-CoV-2 infection, demonstrated by clinical or radiological findings.

Cell lines
293T cells were cultured in DMEM complete (DMEM supplemented with 100 U/ml penicillin, 0.1 mg/ml streptomycin, and 10% FCS) and maintained at 37°C in 5% CO2.

Ethical approval
COVIDx (NCT04326387) was approved by the East of England - Essex Research Ethics Committee (REC ref: 20/EE/0109). Serum samples were obtained from patients attending Addenbrooke’s Hospital with a suspected or confirmed diagnosis of COVID19. Prospective combined point of care testing of suspected COVID-19 cases was done under CUH NHS Trust service evaluation 3163.

METHOD DETAILS

Test methods
NAAT tests
The standard laboratory RT-PCR test, developed by public health England (PHE), targeting the RdRp gene was performed on a combined nose/throat swab. This test has an estimated limit of detection of 320 copies/ml. In parallel, SAMBA II SARS-CoV-2 testing was performed on a combined nose/throat swab and inactivated in a proprietary buffer at the point of sampling. SAMBA II SARS-CoV-2 targets 2 genes- Orf1 and the N genes and uses nucleic acid sequence based amplification to detect SARS-CoV-2 RNA, with limit of detection of 250 copies/ml.

Pseudotype virus preparation
Viral vectors were prepared by transfection of 293T cells by using Fugene HD transfection reagent (Promega) as follows. Confluent 293T cells were transfected with a mixture of 11ul of Fugene HD, 1mg of pCAGGS_SARS-CoV-2_Spike or pCDNA D19Spike-HA, 1ug of p8.91 HIV-1 gag-pol expression vector, and 1.5mg of pCSFLW (expressing the firefly luciferase reporter gene with the HIV-1 packaging signal). Viral supernatant was collected at 48 and 72h after transfection, filtered through 0.45um filter and stored at –80°C. The 50% tissue culture infectious dose (TCID50) of SARS-CoV-2 pseudovirus was determined using Steady-Glo Luciferase assay system (Promega).

Pseudotype neutralisation assay
Spike pseudotype assays have been shown to have similar characteristics as neutralisation testing using fully infectious wild-type SARS-CoV-2. Virus neutralisation assays were performed on 293T cell transiently transfected with ACE2 and TMPRSS2 using SARS-CoV-2 Spike pseudotyped virus expressing luciferase. Pseudovirus was incubated with serial dilution of heat inactivated human serum samples from COVID-19 suspected individuals in duplicates for 1h at 37°C. Virus and cell only controls were also included. Then, freshly trypsinized 293T ACE2/TMPRSS2 expressing cells were added to each well. Following 48h incubation in a 5% CO2 environment at 37°C, the luminescence was measured using Steady-Glo Luciferase assay system (Promega).

Enzyme-linked immunosorbent assay (ELISA)
We developed an ELISA targeting the SARS-CoV-2 Spike and N proteins. Trimeric spike protein antigen used in the ELISA assays consists of the complete S protein ectodomain with a C-terminal extension containing a TEV protease cleavage site, a T4 trimerization foldon and a hexa-histidine tag. The S1/S2 cleavage site with amino acid sequence PRRAR was replaced with a single Arginine residue and stabilizing Proline mutants were inserted at positions 986 and 987. Spike protein was expressed and purified from Expi293 cells (Thermo Fisher). N protein consisting of residues 45-365 was initially expressed as a His-TEV-SUMO-fusion. After Ni-NTA purification, the tag was removed by TEV proteolysis and the cleaved tagless protein further purified on Heparin and gel filtration columns.
The ELISAs were in a stepwise process; a positivity screen was followed by endpoint titer as previously described. Briefly, 96-well EIA/RIA plates (Corning, Sigma) were coated with PBS or 0.1 μg per well of antigen at 4 °C overnight. Coating solution was removed, and wells were blocked with 3% skimmed milk prepared in PBS with 0.1% Tween 20 (PBST) at ambient temperature for 1 hour. Previously inactivated serum samples (56 °C for 1 hour) were diluted to 1:60 or serially diluted by 3-fold, six times in 1% skimmed milk in PBST. Blocking solution was aspirated and the diluted sera were added to the plates and incubated for 2 hours at ambient temperature. Diluted sera were removed, and plates were washed three times with PBST. Goat anti-human IgG secondary antibody Peroxidase (Fc-specific, Sigma) prepared at 1:3,000 in PBST was added and plates were incubated for 1 hour at ambient temperature. Plates were washed three times with PBST. ELISAs were developed using 3,5,3'-5'-tetramethylbenzidine (TMB, ThermoScientific); reactions were stopped after 10 minutes using 0.16M Sulfuric acid.

**COVIDIX 2019 SARS-CoV-2 IgG/IgM Test (COVIDIX Healthcare, Cambridge, UK)**

This colloidal-gold lateral flow immunoassay is designed to detect IgG and IgM to SARS-CoV-2. The test is CE marked. It was used according to the manufacturer’s instructions. 10μl of serum was added to the test well followed by 2 drops of the manufacturer’s proprietary buffer. In order to rule out cross reactivity of this test with seasonal coronavirus antibodies we tested 19 stored specimens from before 2020, some of which had N and S protein SARS-CoV-2 cross reactivity (Table S2).

**SureScreen SARS-CoV-2 IgG/IgM Test (SureScreen Diagnostics Ltd, Derby, UK)**

This colloidal-gold lateral flow immunoassay is designed to detect IgG and IgM to SARS-CoV-2. It was used according to the manufacturer’s instructions. The test has been CE marked and previously validated against a large panel of negative historical controls and in serum from confirmed PCR positive COVID-19 cases. 10μl of serum was added to the test well followed by 2 drops of the manufacturer’s proprietary buffer.

**Next generation sequencing of SARS-CoV-2 isolates in nose/throat swabs**

Samples with CT values above 33 were sequenced with a multiplex PCR approach according to the ARTIC version 2 protocol with version 3 primer set. Amplicons were sequenced using MinION flow cells version 9.4.1 (Oxford Nanopore Technologies, Oxford, UK). Genomes were assembled as previously described. The sequences are freely available from GISAID EpiCoVTM under accession IDs: EPI_ISL 433757, 433754, 433792, 433850, 433751, 433778, 433869, 433875, 433874, 433917, 433962, 433956, 434034, 438681, 438711 and 444331. The submitting laboratory is the COVID-19 Genomics UK (COG-UK) Consortium and the originating laboratory is Department of Pathology, University of Cambridge.

**QUANTIFICATION AND STATISTICAL ANALYSIS**

**Enzyme-linked immunosorbent assay (ELISA) quantification**

The optical density at 450 nm (OD450) was measured using a Spectramax i3 plate reader. The absorbance values for each sample were determined by subtracting OD values from uncoated wells. All data analyses were performed using Prism 8 version 8.4.2 (GraphPad). An OD cut off of 0.3 was used to define a positive IgG response to full length Spike protein.

**COVIDIX 2019 nCoV IgG/IgM Test band density**

For quantification of IgG and IgM band density in COVIDIX 2019 nCoV IgG/IgM Test, high resolution images of completed POC antibody test cassettes were acquired using ChemiDoc MP Imaging System (Bio-Rad) at 20min post-addition of the human serum. Band intensities were analyzed using Image Lab software (Bio-Rad).

**Quantification of neutralisation sensitivity**

The 50% inhibitory dilution (EC50) was defined as the serum dilution at which the relative light units (RLUs) were reduced by 50% compared with the virus control wells (virus + cells) after subtraction of the background RLUs in the control groups with cells only. The EC50 values were calculated with non-linear regression, log (inhibitor) versus normalized response using GraphPad Prism 8 (GraphPad Software, Inc., San Diego, CA, USA). The neutralisation assay was positive if the serum achieved at least 50% inhibition at 1 in 3 dilution of the SARS-CoV-2 spike protein pseudotyped virus in the neutralisation assay. The neutralisation result was negative if it failed to achieve 50% inhibition at 1 in 3 dilution.

**Assessment of neutralisation assay performance**

Four assays detecting IgG to COVID-19 were utilized in this study. 38 of the 45 samples were identified as concordant with at least three of the four assays and considered confirmed either negative or positive. Against this group of samples validated for content of COVID-19 IgG, each individual assay was assessed. Neutralisation, ELISA, SureScreen and COVIDIX assays gave a correct result in 100%, 97.4%, 92.1% and 86.8%, respectively, justifying the choice of the neutralisation assay as standard.

**Analyses**

The performance of SAMBA II SARS-CoV-2 test and COVIDIX SARS-CoV-2 IgG/IgM Test or SureScreen SARS-CoV-2 IgG/IgM Test for diagnosing COVID-19 were calculated alone and then in combination along with binomial 95% confidence intervals (CI). A composite reference standard was used - standard laboratory RT-PCR and a neutralisation assay. Descriptive analyses of clinical and
demographic data are presented as median and interquartile range (IQR) when continuous and as frequency and proportion (%) when categorical. The differences in continuous and categorical data were tested using Wilcoxon rank sum and Chi-square test respectively. Statistical analysis were conducted using Stata (version 13) and GraphPad Prism (version 8), with additional plots generated using GraphPad Prism. Venn diagrams were prepared using Venny. Structural modeling of location of D614G was done using Mol.

**ADDITIONAL RESOURCES**

COVIDx was registered with the ClinicalTrials.gov Identifier NCT04326387.