Antibody profiling and prevalence in US patients during the SARS-CoV2 pandemic

Hari Krishnan Krishnamurthy1†‡*, Vasanth Jayaraman1†‡, Karthik Krishna1, Karenah E. Rajasekaran2, Tianhao Wang1, Kang Bei1, John J. Rajasekaran1†, Inna Yaskin3, Alex J. Rai4, Rok Seon Choung5, Joseph A. Murray5

1 Vibrant Sciences LLC., San Carlos, CA, United States of America, 2 Vibrant America LLC., San Carlos, CA, United States of America, 3 Elite Medical Center, Sunnyvale, CA, United States of America, 4 Department of Pathology & Cell Biology, Columbia University Irving Medical Center, New York, NY, United States of America, 5 Division of Gastroenterology and Hepatology, Mayo Clinic, Rochester, MN, United States of America

† These authors are joint senior authors on this work.
* hari@vibrantsci.com (HKK); jjrajasekaran@vibrantsci.com (JJR)

Abstract

Background

Antibody diagnostics play an important role in disease detection and can potentially aid in monitoring of the immune responses to see if an individual has developed immunity. Developing high throughput diagnostics which does not involve handling of infectious material becomes imperative in the case of pandemics such as the recent outbreak of SARS-CoV2.

Methods

A protein microarray technology was used to detect the plurality of antibody response to four novel antigens namely S1 glycoprotein, Receptor binding domain (RBD), S2 glycoprotein and Nucleoprotein of the novel coronavirus named SARS-CoV2 using serum samples. A DBS card was additionally used to compare its performance with a venipuncture-based serum separator tube (SST) draw.

Results

The three main subclasses of antibodies IgM, IgA and IgG were analyzed to see the variations in immune responses in the affected population and compared to their microbial RT-PCR based NP swab results. The clinical sensitivity and specificity were determined to be 99.67% and 99.77%. In the matrix comparison study, which would enable patients to test without risk of transmitting the virus, DBS (Dried Blood Spot) matched with higher than 98% accuracy to a venipuncture-based serum separator tube (SST) collection.

Conclusion

Multiplex testing enables higher sensitivity and specificity which is essential while establishing exposure on a population scale. This flexible platform along with a discrete collection methodology would be crucial and broadly useful to scale up testing in current and future...
Introduction

Early in December 2019, the first pneumonia cases of unknown origin were reported in Wuhan, China [1, 2]. Subsequently this public health crisis that spread around the world has been identified to be caused by a novel coronavirus (2019-nCoV) or severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) [3, 4]. In most individuals that disease should resolve in itself but there are subgroups with higher risk of morbidity and mortality [5, 6]. In severe cases massive alveolar damage and progressive respiratory failure has been reported [7]. In comparison to the previous coronavirus outbreaks the novel coronavirus appears to have higher transmissibility while having a somewhat higher mortality rate than seasonal flu making for a very toxic combination [8]. Additionally, limited availability of PCR testing for viral illness and the likelihood of a significant proportion of infected individuals having no or trivial symptoms might accelerate disease spread [9].

Diagnostic testing is paramount in identification, containment and monitoring of infectious disease outbreaks [10]. Ideally, the collection process should be performed in quarantine while the diagnostic assay should be highly automated with throughput sufficient to handle several thousands if not millions of samples per day. Antibody diagnostics have been a preferred and accurate method in detection and monitoring of many infectious diseases. Most antibody tests rely on traditional ELISA methodology which has limited multiplexing capability and excessive antigen requirement which limits their ability to be used at scale during pandemics [10]. Traditional methods of specimen collection from an infected individual increases the risk of disease spread and reduces testing capability due to limited availability of specimen collection personnel during such pandemics. We present here a flexible platform for multiplex detection of antibodies from affected individuals enabling high sensitivity with high specificity of detection. We also show comparison of traditional serum separator tube (SST) collection by phlebotomist with dried blood spot-based self-collection by affected subjects to provide for enhanced diagnostics availability during current and future pandemics.

Materials and methods

Patient population

Study 1. To obtain the antibody profile of individuals with SARS-CoV2 microbiological confirmation from respiratory samples by PCR across multiple healthcare centers. 1. Gunnison Valley hospital (# of samples: 132), 2. Elite Medical Center (# of samples: 26). Additional RT-PCR positive samples were obtained from commercial biospecimen laboratories under their approved IRB’s. A cohort of samples that were collected prior to the outbreak were used as negative controls along with disease controls. The sample cohort information age, gender and sample type collected are summarized in Table 1.

Study 2. Traditional phlebotomy-based SST collection comparison to Dried Blood spot collection by subject was performed at healthcare centers across the US. A cohort of 158 clinically paired samples were used for clinical comparison along with 1418 samples which were...
prospectively collected across the geographical locations in the US to identify infection spread and also validate dried blood spot collection vs venipuncture-based SST collection.

Study approval and informed consent process
The study was conducted under the ethical principles that have their origins in the Declaration of Helsinki. The serum samples used for investigation studies such as disease controls were obtained from third-party specimen providers under individual IRBs. The collection of samples with NP swab positive and negative results for SARS-CoV2 was done under a multi-site central IRB (IRB # 1-1299372-1) approved by Western IRB. Informed consent form and patient questionnaire approved under the central IRB were used at all sites. Remnant de-identified samples that were collected prior to the disease outbreak were under IRB # 1-1098539-1 approved by Western IRB.

Pillar plate assembly
The test plate used by Vibrant America is in a 96-pillar plate format as shown in Fig 1. Briefly, silicon wafers are pre-processed to make a high binding surface capable of immobilizing proteins. The following antigens were included in the panel: S1 glycoprotein, Receptor binding domain, S2 glycoprotein, Nucleoprotein. The recombinant antigens were expressed in HEK293 cell lines using full length cDNA coding for the respective antigens fused with a hexa histidine purification tag. Individual wafers were immobilized with each antigen which were then diced into 0.7x0.7 mm² microchips using a stealth dicing process. The diced wafers were picked and
placed onto individual carrier tapes using a standard die sorting system. The carrier tapes were loaded onto a high-throughput surface mount technology (SMT) component placement system. The microchips were then picked and placed onto 96-pillar plates with each pillar containing a layout of 4 microchips—one for each antigen being probed.

**DBS sample processing**

DMPK-type C, Perkin Elmer 226 and Whatman 903 cards suitable for protein-based analysis were used in this study. These cards are compatible with automated punchers enabling rapid scaling depending on processing requirements. Briefly, four punches per card 6mm in diameter were made on the DMPK-C cards and collected in 96 well plates. The elution was done in phosphate buffered saline prior to dilution and performance of the assay.
Assay protocol

Serum samples were probed using 1:50 dilution and DBS samples using a 1:20 dilution on the pil-
lar plate and reacted for 15 minutes at room temperature. The plate was then washed with Tris-
buffered saline with Tween 20 (TBST) (Amresco) buffer 3 x 5 minutes each. The plate was incu-
bated with the secondary antibody (1:2000 dilution of Goat Anti-Human IgG HRP and Goat
Anti-Human IgM HRP and Goat Anti-Human IgA HRP individually) for 15 minutes at room
temperature. The plates were then washed with TBST buffer followed by washing with DI Water.
The plates were finally dried completely before adding chemiluminescent substrate (Clarity Max
from Bio-Rad) and scanned for five minutes on a standard Chemiluminescence Imager. For the
Enhanced IgM Assay, the serum was pre reacted with Goat anti-human IgG Fc fragment prior to
the remaining assay steps to increase the sensitivity of IgM and IgA detection. Assaying is per-
formed on automated liquid handlers enabling a throughput of 100,000 samples per day.

Data analysis

The raw chemiluminescent signals for all the probes were extracted from the images using an
in-house reporter software. The chemiluminescent signals were converted into intensity plots
after quantile normalization, background and spatial correction. The signal threshold was
defined for each antigen by calculating the mean +/- SD of the signal intensity for the same
antigen among the healthy controls collected prior to the infection outbreak. The raw data was
converted into arbitrary chemiluminescent units (CU) based on each individual antigen cut-
off for further analysis.

Results

Analytical performance

We evaluated the analytical performance characteristics of the CoV2 Antibody assay for the
following parameters: precision (repeatability/reproducibility), analytical specificity and lin-
earity. Precision (repeatability) was determined by having two test operators run a panel of 6
samples, 4 replicates daily over a period of 5 days for a total of 40 data points per sample. Panel
consisted of positive control, negative control, positive sample, negative sample, a sample with
concentration +20% above cut-off, and a sample with concentration -20% below cut-off. The
results are summarized in Table 2.

Furthermore, an interfering substance study was conducted to evaluate the potential inter-
ference of specific endogenous and exogenous substances to determine analytical specificity.
The interfering substances with levels tested include Bilirubin 40 mg/dl, Cholesterol 100 mg/
dl, Triglycerides 1000 mg/dl, Hemoglobin 1000 mg/dl, Rheumatoid Factor (RF) 2000 IU/ml,
HAMA 12.5ng/ml, Ribavirin 25mg/dl, Levofoxacin .5mg/dl, Azithromycin .5mg/dl, Ceftriax-
one sodium 25mg/dl, Oxymetazoline 1.25mg/dl, Sodium chloride 25mg/dl, EDTA 12.5mg/ml,
Acetaminophen 50mg/ml, Ibuprofen 50mg/ml, Budesonide 1.25mg/dl. No interference was
observed with any of the substances tested at the stated levels.

Linearity and recovery were tested by diluting positive samples across the assay measuring
range for each antigen in six serial dilutions with negative patient sera. Samples were assayed
by adding mixtures of positive sample and spiking varying concentration of negative sample to
obtain a serial dilution curve. Recoveries for spiked test samples were calculated by compari-
son to the measured recovery of undiluted results. All values represent the average of three rep-
licates tested. The observed values were evaluated against the calculated theoretical values and
a linear regression analysis was performed. R^2 regression values were greater than 0.98 for all
antigens tested. The results are summarized in Table 3.
Clinical sensitivity and specificity

The clinical study to determine the sensitivity and specificity was done using a panel containing retrospectively collected patient serum samples that have been previously confirmed to be positive by SARS-CoV-2 RT PCR along with healthy controls (samples collected prior to SARS-CoV-2 outbreak across age groups) and other disease controls including autoimmune and infectious diseases (as listed in Table 1). Importantly virus positive samples including for other common coronaviruses, influenza A, B, RSV, echovirus, rhinovirus, adenovirus, poliovirus, coxsackie virus, rhinovirus, hepatitis B and hepatitis C did not show cross reactivity to the four antigens specific to the novel coronavirus. Table 4 summarizes the sensitivity and specificity of the individual antigens across the subclasses of antibodies. The specificity of each antigen is relatively high, up to 99%, but the sensitivity is varied among tested antigens. Three antigens—S2 spike IgG, S1 spike IgM, and S2 spike IgM showed more than 80% of sensitivity to identify the cases. When combining all antigens and antibody subclasses together, the CoV2 antibody assay showed 99.67% sensitivity and 99.77% specificity.

Matrix comparison–SST vs DBS

Dried blood spots are a convenient method of collection and processing of samples in pandemic settings. They can be collected in isolation by the affected subject thereby limiting the chances of spreading the infection. Additionally, they do not require any centrifuge for pre-processing and are stable at room temperature for days. Central labs can process these specimens with automated punchers enabling scalable throughput. Specific filter papers for
different applications have been developed and validated including their use in measurement of antibodies to viruses. Prospective collection of 1418 samples at various clinical sites of both venipuncture-based SST collection and dried blood spot collection was performed and results summarized in S1 Fig along with 158 clinically paired samples. The reproducibility was greater than 98% across antigens and antibody subtypes.

### CoV2 antibody positivity in US patient population

This testing platform has the ability to detect three antibody subclasses–IgM, IgA and IgG to four key antigens. An FDA EUA has been granted and testing of the IgM and IgG antibodies is currently being performed across the US. More than 19200 individuals have tested from different geographies and their summary data is shown in Table 5. The overall positivity rate of any antibody for the tested population was 14.3%.

### Discussion

Serological assays play a critical role in identifying burden of disease and also to determine exposure and potential immunity against the infectious agent in a population. In a pandemic setting, discreet collection of samples and high-volume processing in central laboratories are essential for containment and understanding of disease burden. A dried blood spot-based

---

**Table 3. Linearity studies.**

| Spiked level (% positive) | S1 Spike IgG | S1 Spike IgA | S1 Spike IgM |
|--------------------------|-------------|-------------|-------------|
| Level 1 (100%)           | 100%        | 100%        | 100%        |
| Level 2 (75%)            | 103.4%      | 99.8%       | 99.5%       |
| Level 3 (50%)            | 98.2%       | 98.6%       | 104.0%      |
| Level 4 (25%)            | 103.7%      | 101.8%      | 96.7%       |
| Level 5 (12.5%)          | 99.1%       | 102.1%      | 100.9%      |
| Level 6 (6.25%)          | 95.5%       | 97.6%       | 98.2%       |

| Spiked level (% positive) | Nucleoprotein IgG | Nucleoprotein IgA | Nucleoprotein IgM |
|--------------------------|-------------------|-------------------|-------------------|
| Level 1 (100%)           | 100%              | 100%              | 100%              |
| Level 2 (75%)            | 99.1%             | 99.8%             | 100.5%            |
| Level 3 (50%)            | 104.1%            | 96.2%             | 95.7%             |
| Level 4 (25%)            | 103.5%            | 104.6%            | 102.3%            |
| Level 5 (12.5%)          | 98.9%             | 96.1%             | 98.1%             |
| Level 6 (6.25%)          | 95.1%             | 102.3%            | 98.8%             |

| Spiked level (% positive) | RBD IgG | RBD IgA | RBD IgM |
|--------------------------|--------|--------|--------|
| Level 1 (100%)           | 100%   | 100%   | 100%   |
| Level 2 (75%)            | 100.8% | 101.0% | 104.0% |
| Level 3 (50%)            | 97.2%  | 97.9%  | 97.8%  |
| Level 4 (25%)            | 104.0% | 97.0%  | 101.2% |
| Level 5 (12.5%)          | 100.2% | 97.9%  | 98.1%  |
| Level 6 (6.25%)          | 100.1% | 101.3% | 97.2%  |

| Spiked level (% positive) | S2 Spike IgG | S2 Spike IgA | S2 Spike IgM |
|--------------------------|-------------|-------------|-------------|
| Level 1 (100%)           | 100%        | 100%        | 100%        |
| Level 2 (75%)            | 100.8%      | 97.3%       | 100.4%      |
| Level 3 (50%)            | 102.0%      | 97.8%       | 100.5%      |
| Level 4 (25%)            | 104.0%      | 98.5%       | 101.5%      |
| Level 5 (12.5%)          | 97.7%       | 102.1%      | 95.6%       |
| Level 6 (6.25%)          | 97.6%       | 99.2%       | 101.5%      |

[https://doi.org/10.1371/journal.pone.0242655.t003](https://doi.org/10.1371/journal.pone.0242655.t003)
collection and high-volume testing using a flexible platform to test for antibodies of all subclasses against multiple antigens was validated in this study. The ability to add novel discovered antigens and continuously measuring the antibody levels across IgM, IgA and IgG subclasses enable real-time epidemiology studies in a disease outbreak using this platform.

Overall clinical sensitivity was 99.67% with a specificity of 99.77% while S1 and S2 Spike proteins were the two most sensitive markers observed. The sensitivity and specificities of the assay across each antibody subtype is shown in Table 6. Currently diagnosis of the SARS-CoV2 infection is predominantly based on NP swab-based confirmation of microbial DNA while serology adds value in supplemental testing [11]. Positivity on the RNA swab test is dependent on sufficient amount of sample present at the sampling site that can be amplified and detected using standard PCR. The time of sampling is also a crucial factor since viral replication and

Table 4. Clinical performance summary.

| Antigen      | Clinical Sensitivity (95% confidence) | Clinical Specificity (95% confidence) |
|--------------|---------------------------------------|--------------------------------------|
| S1 Spike IgG | 77.23% (72.18% - 81.59%)              | 99.85% (99.70% - 99.92%)             |
| RBD IgG      | 66.34% (60.84% - 71.42%)              | 99.79% (99.63% - 99.88%)             |
| S2 Spike IgG | 81.52% (76.76% - 85.48%)              | 99.83% (99.68% - 99.91%)             |
| NP IgG       | 70.96% (65.61% - 75.78%)              | 99.77% (99.60% - 99.87%)             |
| S1 Spike IgA | 52.15% (46.53% - 57.71%)              | 99.92% (99.80% - 99.97%)             |
| RBD IgA      | 41.25% (35.85% - 46.87%)              | 99.90% (99.78% - 99.96%)             |
| S2 Spike IgA | 41.58% (36.17% - 47.21%)              | 99.90% (99.78% - 99.96%)             |
| NP IgA       | 32.34% (27.33% - 37.80%)              | 99.94% (99.83% - 99.98%)             |
| S1 Spike IgM | 70.30% (64.92% - 75.16%)              | 99.85% (99.70% - 99.92%)             |
| RBD IgM      | 66.34% (60.84% - 71.42%)              | 99.83% (99.68% - 99.91%)             |
| S2 Spike IgM | 67.99% (62.54% - 72.99%)              | 99.81% (99.65% - 99.90%)             |
| NP IgM       | 63.73% (57.81% - 68.59%)              | 99.81% (99.65% - 99.90%)             |
| OVERALL IgG  | 99.01% (97.13% - 99.66%)              | 99.77% (99.60% - 99.87%)             |
| OVERALL IgA  | 61.72% (56.13% - 67.01%)              | 99.90% (99.78% - 99.96%)             |
| OVERALL IgM  | 87.46% (83.25% - 90.73%)              | 99.81% (99.65% - 99.90%)             |
| OVERALL      | 99.67% (98.15% - 99.94%)              | 99.77% (99.60% - 99.87%)             |

Table 6. Sensitivity and Specificity of SARS-CoV2 markers over time.

| Antigen      | < 7 days Sensitivity | 7–14 days Sensitivity | > 14 days Sensitivity | Total Sensitivity |
|--------------|----------------------|-----------------------|----------------------|------------------|
| S1 Spike IgG | 42.86%               | 56.73%                | 100.00%              | 77.23%           |
| RBD IgG      | 33.33%               | 28.85%                | 100.00%              | 66.34%           |
| S2 Spike IgG | 54.76%               | 64.42%                | 100.00%              | 81.52%           |
| NP IgG       | 35.71%               | 41.35%                | 100.00%              | 70.96%           |
| S1 Spike IgA | 52.38%               | 70.19%                | 40.13%               | 52.15%           |
| RBD IgA      | 40.48%               | 46.15%                | 38.22%               | 41.25%           |
| S2 Spike IgA | 40.48%               | 54.81%                | 33.12%               | 41.58%           |
| NP IgA       | 30.95%               | 29.81%                | 34.39%               | 32.34%           |
| S1 Spike IgM | 78.57%               | 66.35%                | 70.70%               | 70.30%           |
| RBD IgM      | 80.95%               | 62.50%                | 64.97%               | 66.34%           |
| S2 Spike IgM | 71.43%               | 60.58%                | 71.97%               | 67.99%           |
| NP IgM       | 59.52%               | 50.96%                | 72.61%               | 63.37%           |
| OVERALL IgG  | 92.86%               | 100.00%               | 100.00%              | 99.01%           |
| OVERALL IgA  | 66.67%               | 83.65%                | 45.86%               | 61.72%           |
| OVERALL IgM  | 97.62%               | 100.00%               | 76.43%               | 87.46%           |
| OVERALL      | 97.62%               | 100.00%               | 100.00%              | 99.67%           |

https://doi.org/10.1371/journal.pone.0242655.t004
load increases in specific window periods and sampling outside of this may lead to false negatives. Poor techniques in sample collection could also lead to false negatives.

When it comes to serological tests the main limitations that make them suboptimal tools for diagnosing those who are sick is that it takes time for the development of antibodies after infection. Thus, they may produce a false negative result in individuals who are acutely infected. On the other hand, they can be useful to indicate exposure in individuals who are asymptomatic or were symptomatic but PCR negative or were never tested. Nevertheless, it is important to note that the serological testing should not be used as the sole basis for diagnosis of an acute COVID19 infection.

Our study has some important limitations that should be taken into consideration while reviewing the data presented. The samples that were tested were ordered by physicians which would have inherent selection bias as to who is getting tested such as biased testing of individuals who were symptomatic. The prevalence data that we generated should hence not be considered to be from randomly selected individuals, additionally the number of tests performed is limited in certain states and the data should be looked at with caution in such cases. The antibodies that were measured in this study were against 4 recombinant antigens and we do not know if the individuals produced neutralizing antibody response.

### Table 5. Antibody prevalence in the US patient population.

| State | Total Samples Tested | Any Antibody Positive % | IgA Positive % | IgM Positive % | IgG Positive % |
|-------|----------------------|-------------------------|---------------|---------------|---------------|
| AZ    | 2713                 | 15.37%                  | 2.54%         | 8.99%         | 9.18%         |
| CA    | 5717                 | 13.91%                  | 2.57%         | 7.64%         | 9.06%         |
| CO    | 2780                 | 16.83%                  | 3.92%         | 9.24%         | 11.91%        |
| FL    | 513                  | 14.42%                  | 4.87%         | 7.02%         | 10.92%        |
| GA    | 343                  | 15.74%                  | 3.79%         | 10.50%        | 12.83%        |
| IL    | 659                  | 10.32%                  | 2.12%         | 5.46%         | 8.04%         |
| IN    | 369                  | 7.05%                   | 1.36%         | 3.79%         | 4.07%         |
| KS    | 217                  | 19.35%                  | 4.61%         | 7.37%         | 16.13%        |
| MA    | 337                  | 27.60%                  | 5.93%         | 11.57%        | 20.18%        |
| MD    | 731                  | 5.61%                   | 0.41%         | 2.33%         | 4.24%         |
| ME    | 116                  | 6.90%                   | 0.00%         | 5.17%         | 1.72%         |
| MI    | 172                  | 19.77%                  | 8.14%         | 10.47%        | 16.28%        |
| MN    | 155                  | 10.53%                  | 3.01%         | 3.76%         | 7.52%         |
| MO    | 496                  | 10.48%                  | 1.81%         | 4.44%         | 7.46%         |
| NC    | 220                  | 8.64%                   | 1.36%         | 5.45%         | 4.55%         |
| NE    | 243                  | 12.76%                  | 4.12%         | 5.35%         | 9.88%         |
| NJ    | 786                  | 24.81%                  | 10.31%        | 14.89%        | 19.85%        |
| NY    | 209                  | 28.71%                  | 11.48%        | 14.83%        | 24.88%        |
| OH    | 244                  | 12.30%                  | 2.87%         | 6.56%         | 7.79%         |
| OK    | 55                   | 9.09%                   | 0.00%         | 5.45%         | 5.45%         |
| OR    | 255                  | 10.59%                  | 2.75%         | 4.31%         | 8.24%         |
| PA    | 128                  | 10.16%                  | 1.56%         | 7.03%         | 3.91%         |
| TN    | 244                  | 6.97%                   | 0.41%         | 1.23%         | 6.15%         |
| TX    | 1190                 | 11.26%                  | 2.69%         | 8.32%         | 6.47%         |
| VA    | 198                  | 7.07%                   | 2.02%         | 4.04%         | 5.05%         |
| WA    | 123                  | 13.01%                  | 1.63%         | 4.07%         | 11.38%        |
| WI    | 64                   | 9.38%                   | 3.13%         | 7.81%         | 4.69%         |
| **Grand Total** | **19255** | **14.30%** | **3.20%** | **7.87%** | **9.79%** |

https://doi.org/10.1371/journal.pone.0242655.t005
Technology developments truly enable testing for multiple antibodies from small volumes of collected blood. The potential convenience and cost savings that it would bring to laboratory testing cannot be understated. Remote sampling such as with DBS cards, along with telemedicine would help providers to order tests not only in pandemic settings but also can be a great enabler in bringing healthcare costs down. Immediate application of such a platform to do multiplex testing would be in the monitoring of exposure to upper respiratory tract infections.

A subset of the NP swab positive individuals are being monitored for antibody levels and symptoms at regular intervals from the time they tested positive. Initial results show that the seroconversion from IgM to IgG which is marked by a reduction in IgM titer might be co-related with recovery since it is accompanied by a negative result in the subsequent NP swab test and also improvement in symptoms. This could possibly evolve into the criteria for confirmation of recovery from infection which would enable return to work of affected individual. Additionally, in patients that have had follow up testing for antibodies we noticed that the antibody levels remain high in individuals for 5 months and none of them have been re-infected with the virus indicating that antibodies are likely to generate immunity in affected individuals who have recovered.

In summary, we have developed and validated a highly accurate assay that enables the identification of individuals who have been infected with SARS CoV2. The assay meets the rigorous requirements of high sensitivity and specificity, scalability, high fidelity and reliability that will be critical to the control of the current pandemic. The utility of the self-collection method aligns with the dual needs of safe convenient testing with continued isolation for patients that may either be infectious or those who are sheltering at home to avoid infection exposure. If convalescent antibodies are demonstrated to be protective from future infection, then this testing could enable an easing of restrictions both at the individual and community level.

### Table 6. Clinical sensitivity and specificity.

| Vibrant COVID-19 Ab | Clinical Diagnosis–NP Swab Positive | Total | Analysis (95% Confidence) |
|---------------------|-----------------------------------|-------|---------------------------|
| Overall IgG/IgA/IgM |                                   |       |                           |
| Positive            | 302 12 314 | Sensitivity = 99.67% (98.15% - 99.94%) |
| Negative            | 1 5250 5251 | Specificity = 99.77% (99.60% - 99.87%) |
| Total               | 303 5262 5565 |                           |
| Overall IgG         |                                   |       |                           |
| Positive            | 300 12 312 | Sensitivity = 99.01% (97.13% - 99.66%) |
| Negative            | 3 5250 5253 | Specificity = 99.77% (99.60% - 99.87%) |
| Total               | 303 5262 5565 |                           |
| Overall IgA         |                                   |       |                           |
| Positive            | 187 5 192 | Sensitivity = 61.72% (56.13% - 67.01%) |
| Negative            | 116 5257 5373 | Specificity = 99.90% (99.78% - 99.96%) |
| Total               | 303 5262 5565 |                           |
| Overall IgM         |                                   |       |                           |
| Positive            | 265 10 275 | Sensitivity = 87.46% (83.25% - 90.73%) |
| Negative            | 38 5252 5290 | Specificity = 99.81% (99.65% - 99.90%) |
| Total               | 303 5262 5565 |                           |
Supporting information

S1 Fig. Venipuncture vs Dried Blood Spot. (PNG)

Acknowledgments

The authors thank the FDA and the staff of CDC for early guidance on validation requirements. We thank the healthcare providers and the study participants who made this research possible. Special thanks to Inna Yaskin, Jaclyn McCoy, Tina Wilson, Roanne Houck and Gunnison Valley Hospital for coordinating research efforts. Research reported in this publication was supported by Vibrant America LLC. The content is solely the responsibility of the authors and does not necessarily represent the official views of the CDC or FDA. We also would like to thank Miaomiao Sun and Ashish Gopalakrishnan for generating the Fig 1 illustration.

Author Contributions

Conceptualization: Hari Krishnan Krishnamurthy, Vasanth Jayaraman, Karthik Krishna, John J. Rajasekaran.

Data curation: Karenah E. Rajasekaran.

Formal analysis: Vasanth Jayaraman, Tianhao Wang.

Investigation: Hari Krishnan Krishnamurthy, Vasanth Jayaraman, Karthik Krishna, Karenah E. Rajasekaran, John J. Rajasekaran.

Methodology: Hari Krishnan Krishnamurthy, Karthik Krishna, Tianhao Wang.

Project administration: Hari Krishnan Krishnamurthy, Vasanth Jayaraman, John J. Rajasekaran.

Resources: Inna Yaskin.

Software: Kang Bei.

Supervision: Hari Krishnan Krishnamurthy, Vasanth Jayaraman, John J. Rajasekaran.

Validation: Karthik Krishna.

Visualization: Hari Krishnan Krishnamurthy.

Writing – original draft: Hari Krishnan Krishnamurthy.

Writing – review & editing: Hari Krishnan Krishnamurthy, Vasanth Jayaraman, John J. Rajasekaran, Alex J. Rai, Rok Seon Choug, Joseph A. Murray.

References

1. Zhu N, Zhang D, Wang W, Li X, Yang B, Song J, et al. A Novel Coronavirus from Patients with Pneumonia in China, 2019. N Engl J Med. 2020; 382(8):727–33. Epub 2020/01/25. https://doi.org/10.1056/NEJMoa2001017 PMID: 31978945; PubMed Central PMCID: PMC7092803.

2. Li Q, Guan X, Wu P, Wang X, Zhou L, Tong Y, et al. Early Transmission Dynamics in Wuhan, China, of Novel Coronavirus-Infected Pneumonia. N Engl J Med. 2020; 382(13):1199–207. Epub 2020/01/30. https://doi.org/10.1056/NEJMoa2001316 PMID: 31995857; PubMed Central PMCID: PMC7121484.

3. Bogoch II, Watts A, Thomas-Bachli A, Huber C, Kraemer MUG, Khan K. Pneumonia of unknown aetiology in Wuhan, China: potential for international spread via commercial air travel. J Travel Med. 2020; 27(2). Epub 2020/01/17. https://doi.org/10.1093/jtm/taaa006 PMID: 31943059; PubMed Central PMCID: PMC7107534.
4. Sohrabi C, Alsafi Z, O’Neill N, Khan M, Kerwan A, Al-Jabir A, et al. World Health Organization declares global emergency: A review of the 2019 novel coronavirus (COVID-19). Int J Surg. 2020; 76:71–6. Epub 2020/03/01. https://doi.org/10.1016/j.ijsu.2020.02.034 PMID: 32112977; PubMed Central PMCID: PMC7105032.

5. Chen N, Zhou M, Dong X, Qu J, Gong F, Han Y, et al. Epidemiological and clinical characteristics of 99 cases of 2019 novel coronavirus pneumonia in Wuhan, China: a descriptive study. Lancet. 2020; 395(10223):507–13. Epub 2020/02/03. https://doi.org/10.1016/S0140-6736(20)30211-7 PMID: 32007143; PubMed Central PMCID: PMC7135076.

6. Lu R, Zhao X, Li J, Niu P, Yang B, Wu H, et al. Genomic characterisation and epidemiology of 2019 novel coronavirus: implications for virus origins and receptor binding. Lancet. 2020; 395(10224):565–74. Epub 2020/02/03. https://doi.org/10.1016/S0140-6736(20)30251-8 PMID: 32007145; PubMed Central PMCID: PMC7159086.

7. Wang D, Hu B, Hu C, Zhu F, Liu X, Zhang J, et al. Clinical Characteristics of 138 Hospitalized Patients With 2019 Novel Coronavirus-Infected Pneumonia in Wuhan, China. JAMA. 2020. Epub 2020/02/08. https://doi.org/10.1001/jama.2020.1585 PMID: 32031570; PubMed Central PMCID: PMC7042881.

8. Zhao S, Lin Q, Ran J, Musa SS, Yang G, Wang W, et al. Preliminary estimation of the basic reproduction number of novel coronavirus (2019-nCoV) in China, from 2019 to 2020: A data-driven analysis in the early phase of the outbreak. Int J Infect Dis. 2020; 92:214–7. Epub 2020/02/03. https://doi.org/10.1016/j.ijid.2020.01.050 PMID: 32007643; PubMed Central PMCID: PMC7110798.

9. Huang L, Zhang X, Zhang X, Wei Z, Zhang L, Xu J, et al. Rapid asymptomatic transmission of COVID-19 during the incubation period demonstrating strong infectivity in a cluster of youngsters aged 16–23 years outside Wuhan and characteristics of young patients with COVID-19: A prospective contact-tracing study. J Infect. 2020. Epub 2020/04/14. https://doi.org/10.1016/j.jinf.2020.03.006 PMID: 32283156.

10. Lo YMD, Chiu RWK. Racing Towards the Development of Diagnostics for a Novel Coronavirus (2019-nCoV). Clin Chem. 2020; 66(4):503–4. Epub 2020/02/08. https://doi.org/10.1093/clinchem/hvaa038 PMID: 32031590; PubMed Central PMCID: PMC7108485.

11. Pan Y, Li X, Yang G, Fan J, Tang Y, Zhao J, et al. Serological immunochromatographic approach in diagnosis with SARS-CoV-2 infected COVID-19 patients. J Infect. 2020. Epub 2020/04/14. https://doi.org/10.1016/j.jinf.2020.03.051 PMID: 32283141.