Antibody Affinity Governs the Inhibition of SARS-CoV-2 Spike/ACE2 Binding in Patient Serum

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Cite This: https://doi.org/10.1021/acsinfecdis.1c00047

ABSTRACT: The humoral immune response plays a key role in suppressing the pathogenesis of SARS-CoV-2. The molecular determinants underlying the neutralization of the virus remain, however, incompletely understood. Here, we show that the ability of antibodies to disrupt the binding of the viral spike protein to the angiotensin-converting enzyme 2 (ACE2) receptor on the cell, the key molecular event initiating SARS-CoV-2 entry into host cells, is controlled by the affinity of these antibodies to the viral antigen. By using microfluidic antibody-affinity profiling, we were able to quantify the serum-antibody mediated inhibition of ACE2–spike binding in two SARS-CoV-2 seropositive individuals. Measurements to determine the affinity, concentration, and neutralization potential of antibodies were performed directly in human serum. Using this approach, we demonstrate that the level of inhibition in both samples can be quantitatively described using the dissociation constants (K_D) of the binary interactions between the ACE2 receptor and the spike protein as well as the spike protein and the neutralizing antibody. These experiments represent a new type of in-solution receptor binding competition assay, which has further potential applications, ranging from decisions on donor selection for convalescent plasma therapy, to identification of lead candidates in therapeutic antibody development, and vaccine development.

KEYWORDS: SARS-CoV-2, COVID-19, neutralizing antibodies, competition assay, in-solution binding, microfluidics

INTRODUCTION

The COVID-19 pandemic is causing not only a major public health crisis but also unprecedented economic challenges. One of the features of COVID-19 is the variability of disease outcomes, with a large majority of patients presenting mild or no symptoms, while others may become severely ill or die.1 Independently of the disease trajectory, the majority of acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infections trigger a response of the adaptive immune system.2−4 Essential for this immune response are antibodies that neutralize SARS-CoV-2 (NAbs) by preventing virus binding to the host-cell receptor angiotensin-converting enzyme 2 (ACE2).5−13 Effective NAbs target the receptor binding domain (RBD) of viral spike proteins and hence inhibit ACE2 binding.13 Such antibodies and their characterization are of great interest for therapeutic approaches such as plasmapheresis14 and as structural templates for rational vaccine design.15 Consequently, there is a need for assays that identify the most potent NAbs reliably and efficiently. The gold standard to detect NAbs is a virus-neutralization test (VNT) on live cells, which can be performed with either live viruses or pseudoviruses.16,17 While these assays are well established, VNTs are time-consuming, logistically challenging, can take several days for cell culture work and to obtain results, and necessitate stringent biosafety measures to be in place for handling live virus.18 Furthermore, it is challenging to standardize VNTs to allow consistency among different laboratories, studies, and patient cohorts.19 This situation has motivated the search for easier to use and faster alternatives in the form of surrogate VNTs (sVNTs). Such assays are typically based on an enzyme-linked immunosorbent assay (ELISA), which provides information on the titers of NAbs in a patient sample.20 Neither the standard VNTs nor the ELISA-based sVNTs can, however, provide information on whether virus neutralization is achieved by a high concentration of NAbs with weak binding affinities or a lower concentration of NAbs with tight binding affinities. This information, however, is of particular importance not just for understanding the fundamental physicochemical parameters that underlie neutralization but...
also has potential applications for the selection of suitable donors for plasmapheresis. Here, we investigate whether NAb efficacy can be predicted based on affinity by assessing the binding free energies that drive ACE2 displacement from the SARS-CoV-2 RBD. Using microfluidic antibody-affinity profiling (MAAP), we demonstrate that a ternary equilibrium of ACE2, SARS-CoV-2 spike S1 (S1), and antibodies from seropositive, recovered individuals predicts and quantifies the inhibition of ACE2–S1 binding. MAAP is an in-solution assay that determines both, the effective dissociation constants ($K_D$) of polyclonal serum antibodies and their concentrations.\textsuperscript{21,22} Based on these findings, we suggest that the ratio of the dissociation constants of ACE2–S1 binding and NAb–S1 binding ($r = K_{D,ACE2}/K_{D,NAb}$) together with the experimentally determined concentration of NAbs in serum are excellent measures to predict the inhibitory potential of the polyclonal antibody response. Such quantitative information on antibody-mediated inhibition of ACE2–S1 binding is crucial to drive decision making in donor selection for plasmapheresis, for lead identification in the development of therapeutic antibodies, and in vaccine development.

**RESULTS AND DISCUSSION**

Affinity Measurements in Prepandemic Control Serum Using Recombinant ACE2, S1, and a Monoclonal Neutralizing Anti-S1 Antibody. In this study, we investigate whether the affinity of anti-S1 antibodies present in human serum samples is a predictor for their neutralization efficacy. For this purpose, we established a novel assay using microfluidic diffusional sizing (MDS, Figure 1A) that utilizes a recombinant, monoclonal NAb (NAb1) spiked into SARS-CoV-2 prepandemic control serum in the presence of recombinant ACE2 and S1. The NAb-mediated inhibition of ACE2–S1 binding is physically determined by the ternary equilibrium of ACE2, SARS-CoV-2 spike S1 (S1), and NAb, which is described quantitatively by the $K_D$ values of the two binary equilibria, ACE2–S1 and NAb–S1 (refer to Materials and Methods for details). Thus, to validate the data generated on the ternary system, we determined the $K_D$ values of the two underlying binary reactions.

First, we measured the $K_D$ of ACE2–S1 binding in the presence of 90% prepandemic control serum (Figure 1B). As the S1 concentration was increased, more of the fluorescently labeled ACE2 was found in the protein complex, and the effective hydrodynamic radius ($R_h$) increased. Based on a 1:1 equilibrium binding model (eq 2), the $K_D$ of ACE2–S1 was
were determined by MDS, as displayed in Figure 1B and C. These two a

isotherm based on a competition binding model (eq 11). In

1 μ concentrations of free ACE2, S1, and NAb1 as well as of the

(\([ACE2]_0 = 10 \text{ nM}, [S1]_0 = 50 \text{ nM}, \text{and } 30 \text{ pM}

the ternary mixture used for the competition experiment

displace ACE2 from S1 (Figure 1D, red). The experimental

RBD and competes with ACE2 for S1 binding, and that the

prediction, demonstrating that NAb1 targets the spike

inhibition isotherm was in remarkably good agreement with

interaction.

globally (taken from Figure 1D). The global

from Figure 1B and C) and the ternary inhibition isotherm

Control Serum by a Monoclonal Neutralizing Anti-S1

whether NAb1-mediated inhibition of ACE2

concentrations of ACE2 and S1, cannot easily be in

a concentration of 10 nM and S1 at a concentration of 50 nM

addition of NAb1 to a mixture of

Microfluidic Antibody-Affinity Profiling and Neutraliz-

potency range for high-affinity monoclonal

Inhibition of ACE2–S1 Binding in Preparademic Control Serum by a

Monoclonal Neutralizing Anti-S1 Antibody. After having charac-

terized the binding between ACE2 and S1 as well as between NAb1 and S1, we investigated

whether NAB1-mediated inhibition of ACE2–S1 binding occurs in the ternary mixture of ACE2, S1, and NAB1. The addition of NAB1 to a mixture of fluorescently labeled ACE2 at a concentration of 10 nM and S1 at a concentration of 50 nM resulted in a reduction of \(R_b\) caused by complex formation of fluorescently labeled S1 and NAB1. The resulting \(K_D\) of NAB1–S1 was determined to be 5.1 nM (95% CI: 1.7–11.6), which is a typical affinity range for high-affinity monoclonal antibodies.24

For a quantitative analysis of these inhibition data, we
globally fitted both, the two binary binding isotherms (taken from Figure 1B and C) and the ternary inhibition isotherm (taken from Figure 1D). The global fit utilizes the \(K_D\) values of ACE2–S1 and NAB1–S1 as well as \(R_b\) of unbound ACE2 as global fitting parameters (Figure 2A). The \(K_D\) values obtained from the global fit were in agreement with those extracted from the individual binary fits, demonstrating that NAB1-mediated inhibition is driven by the binding free energy of the competing protein complexes ACE2–S1 and NAB1–S1. Next, we used \(K_{D,ACE2}\) and \(K_{D,NAb}\) to calculate equilibrium concentrations of free ACE2, S1, and NAB1 as well as of the complexes formed by ACE2–S1 and NAB1–S1. Based on those concentrations, we determined the S1-bound fraction of ACE2 and the NAB1-bound fraction of S1 to evaluate NAB1 efficacy (Figure 2B).

An effective NAB will block viral spike proteins from binding to ACE2, which is key to prevent host-cell infection. In vivo, this is accomplished through a combination of NAB–S1 affinity and NAB concentration, as the other parameters that govern the equilibrium, such as ACE2–S1 affinity and the concentrations of ACE2 and S1, cannot easily be influenced by the host. Thus, the most effective NABs will have \(K_D\) values considerably lower than the \(K_D\) of ACE2–S1 and concentrations considerably higher than their \(K_D\) for S1 binding. This condition is directly described by the ratio \(r = K_{D,ACE2}/K_{D,NAb}\), which is provided by our assay. For example, at \(r = 10\), under the condition that the concentration of NAB largely exceeds that of S1, the concentration of NAB could be ten times lower than the concentration of ACE2 to still bind the same amount of S1.

Microfluidic Antibody-Affinity Profiling and Neutralization Potency in Serum of SARS-CoV-2 Seropositive Individuals. After establishing the assay using a recombinant, monoclonal NAb, we applied the same approach to the quantification of the potency of polyclonal anti-S1 antibody responses in COVID-19 patient samples. To do so, we analyzed blood sera of two individuals who had tested seropositive for anti-SARS-CoV-2 IgG and IgM. Sample 1 was obtained from a 52-year old white female with mild COVID-19 symptoms, while sample 2 was derived from a 44-year old asymptomatic white male.

To determine the \(K_D\) of anti-S1 antibody binding to S1, we
mixed fluorescently labeled S1 at a constant concentration with

https://doi.org/10.1021/acsinfecdis.1c00047

ACS Infect. Dis. XXX, XXX, XXX–XXX
patient serum at various dilutions and determined $R_b$ as an indicator of complex formation (Figure 3A and C). Since the anti-S1 antibody concentration in the two patient sera was unknown, we extracted this information from the equilibrium binding isotherms by using the concentration of antibody-binding sites as a fitting parameter.  

To obtain well-constrained binding-site concentrations, we measured antibody-binding isotherms at three different concentrations of fluorescently labeled S1. To also assess the neutralization potential of the anti-S1 antibodies in the patient sera, we obtained inhibition isotherms from ternary mixtures of fluorescently labeled ACE2, S1, and patient serum at various concentrations (Figure 3B and D). Both seropositive samples inhibited binding of ACE2 to S1. While sample 1 reached complete inhibition at a serum concentration of 90%, ACE2 was not completely displaced from S1 in sample 2, which could either be due to weak affinity antibodies or a too low concentration of antibodies.

For each serum sample, we generated three binary binding isotherms and one inhibition isotherm, which we used as a combined set of data to extract $K_D$ values of antibody–S1 binding, concentrations of antibody-binding sites, and neutralization potency. As demonstrated with NAb1 (Figure 2A), we globally fitted binary antibody–S1 (eq 2) and ternary antibody-mediated inhibition (eq 16) using the $K_D$ of antibody–S1 binding as a shared parameter across all sets of data. The $K_D$ value of ACE2–S1 binding in human serum (7.0 nM) was used as a constant in the fit, based on the data shown in Figure 1B. In both patients, global analysis of binding and inhibition resulted in $K_D$ values in the subnanomolar range, which is in line with the values obtained by fitting the binding isotherms without the inhibition isotherm (Table S1 and Figure S1).

A MAAP study on a larger number of patient samples showed a similar range of affinities among anti-spike serum antibodies.  

While the agreement of Bayesian inference with the global fits was excellent, in both patient samples the posterior probability for the $K_D$ distribution provided strong constraints for the upper bound but weak constraints for the lower bound, thus providing an upper limit on the $K_D$ value and indicating a tight binding interaction. The best-fit values from the global fits converged to this upper limit, representing the higher end of $K_D$ values that describe the data. For sample
2, the antibody-binding affinity was tighter than for sample 1. The concentration of antibody-binding sites was 110 nM (CI 95%; 90–130) and 19 nM (CI 95%; 17–23) for sample 1 and sample 2, respectively, values which are within the concentration range observed in a MAAP study on a larger number of samples.21 In both serum samples, the $K_D$ values of antibody–S1 binding were considerably lower than the concentration of antibody-binding sites, a condition which is required for effective antibody binding. Moreover, in both samples the ratio $r = K_D,\text{ACE2}/K_D,\text{NAb}$ is 10 or higher, indicating that both individuals produced antibodies against S1 with good neutralization potential. The global fit describes the inhibition isotherms (Figure 3B and D) of both patient sera remarkably well, demonstrating that antibody-mediated inhibition of the ACE2–S1 interaction is driven by serum-antibody affinity. Furthermore, this indicates that most of the anti-S1 antibodies raised by these two individuals target the receptor binding motif of S1 and compete with ACE2 binding. Antibodies that bind to other noncompetitive epitopes would result in data that deviate from the combined analysis of binary and ternary equilibria used here.

## CONCLUSIONS

In this study, we have used microfluidic antibody affinity profiling to measure independently the binding affinities of the two molecular interactions involved in neutralizing SARS-CoV-2 cell entry, namely, that between ACE2–spike and spike–NAb. Furthermore, we have demonstrated that the knowledge of these thermodynamic parameters enables a quantitative description of the ability of patient antibodies to inhibit the binding of the viral spike protein to the host-cell ACE2 receptor. The fundamental parameters characterizing the binding interaction, the concentration of the antibodies and their affinity to the target measured in our assay, are independent of the concentration of probe molecules used and, as such, the results are comparable even on an absolute basis between individual experiments. The knowledge of both, affinity and concentration of anti-S1 antibodies provides valuable information that has the potential to support decision making in research as well as in clinical practice. For example, in convalescent plasma therapy, a donor with high-affinity, high-concentration anti-RBD antibodies would be most suitable as donor antibodies are diluted approximately by a factor of 10 during the procedure. Specifically, based on this analysis, sample 1 containing high-affinity antibodies at a concentration of 110 nM should be preferred for plasmapheresis over sample 2, which also contains high-affinity antibodies but at a significantly lower concentration of 19 nM. Furthermore, for the selection of antibodies as therapeutic candidates, it is crucial to be able to deconvolute antibody titers into the fundamental quantities of affinity and concentration. Indeed, a given antibody titer, as measured for example by conventional ELISA, could arise due to a low concentration of very tightly binding antibodies, which would be attractive candidates for further developments, or from a high concentration of weakly binding antibodies, which are less interesting as leads for further optimization for therapeutic use. Finally, the ability to measure antibody concentration and their affinity provides an added level of granularity in understanding the diverse immune responses characteristic of COVID-19 infections. This has the potential to evaluate and understand the potency of vaccines, which, in order to be optimally effective, must generate immune responses that lead to high-affinity virus-neutralizing antibodies at high concentrations.

## MATERIALS AND METHODS

### Fluorescent Labeling of Proteins.

SARS-CoV-2 spike S1 domain (Val 16–Arg 685; S1N-C52H4, ACROBiosystems) and the extracellular domain of ACE2 (Gln 18–Ser 740, AC2-H52H8, ACROBiosystems) were each recomposed in phosphate buffered saline (PBS) at pH 7.4 (Merck) at a concentration of 0.6 mg/mL according to the manufacturer’s instructions. For labeling, the pH of 50 μg of protein solution was adjusted to pH 8.3 using 1 M NaHCO$_3$ (Merck) followed by addition of Alexa Fluor 647 NHS ester (Thermo Fisher) at a molar dye-to-protein ratio of 3:1. Samples were incubated at 4 °C overnight, and free dye was removed by size-exclusion chromatography on an ÄKTA pure system (Cytiva) equipped with a Superdex 200 Increase 3.2/300 column (Cytiva) using PBS at pH 7.4 as a buffer. Labeled and purified proteins were stored at −80 °C in PBS pH 7.4 containing 10% (v/v) glycerol as cryoprotectant.

### Affinity Measurements of Binary Equilibrium by Microfluidic Diffusional Sizing.

To determine the affinity of ACE2–SARS-CoV-2 spike S1 binding, Alexa Fluor 647 labeled ACE2 (10 nM) was mixed with unlabeled SARS-CoV-2 spike S1 at increasing concentrations (100 pM to 3.2 μM) in the presence of 90% heat-inactivated human serum (HS667, Merck) and incubated at 4 °C overnight. To determine the affinity of a neutralizing antibody to SARS-CoV-2 spike S1, fluorescently labeled SARS-CoV-2 spike S1 (10 nM) was mixed with unlabeled NAb1(SAD-S35, ACROBiosystems) at increasing concentrations (3.8 pM to 250 nM) in the presence of 90% heat-inactivated human serum and incubated at 4 °C overnight. To measure complex formation by microfluidic diffusional sizing (MDS), 5 μL of sample were pipetted on a microfluidic chip, and analysis was performed at the 1.5–8.0 nm setting on a Fluidity One-W Serum instrument (center wavelength for excitation is 630 nm with a bandwidth of 38 nm and center wavelength for emission is 694 nm with a bandwidth of 44 nm; Fluidic Analytics). Serum autofluorescence was determined in the absence of labeled protein and used to correct MDS data measured of binding interactions. Error bars shown in figures are standard deviations from triplicate measurements. The equilibrium dissociation constant ($K_D$) was determined by nonlinear least-squares (NLSQ) fitting ( Prism, GraphPad Software) in terms of the following binary equilibrium:

\[ U + L \rightleftharpoons K_c U L \]  

\[ R_h = R_h,\text{free} + \left( R_h,\text{complex} - R_h,\text{free} \right) \times \left( \frac{K_D + n[U]_0 + [L]_0 - \sqrt{(K_D + n[U]_0 + [L]_0)^2 - 4n[U]_0[L]_0}}{2[L]_0} \right) \]  

with $R_h,\text{free}$, $R_h,\text{complex}$ being the effective hydrodynamic radii at equilibrium, of the unbound labeled species, and of the complex of unlabeled and labeled species, respectively. The parameter $n$ is equivalent to the number of its binding sites. Furthermore, $[U]_0$ and $[L]_0$ are total concentrations of unlabeled and labeled species, respectively.

### Measurement of Ternary Equilibrium (Recombinant Proteins) by Microfluidic Diffusional Sizing.

To assess inhibition of ACE2–S1 binding by a recombinant NAb, a
mixture of Alexa Fluor 647 labeled ACE2 and unlabeled S1 at concentrations of 10 nM and 50 nM, respectively, was mixed immediately with increasing concentrations of NAb (30 pM–1.0 μM) in the presence of 90% heat-inactivated human serum (HS667, Merck) and equilibrated at a temperature of 4 °C overnight. MDS data was obtained as described above.

The following ternary equilibrium model was used for data analysis:

\[
\text{Ab} + (\text{S1ACE2}) \xrightarrow{K_{D,ACE2}} \text{Ab} + \text{ACE2} + \text{S1} \xrightarrow{K_{D,AB}} (\text{S1Ab}) + \text{ACE2}
\]

In this ternary mixture, the two \(K_D\) values are defined as follows:

\[
K_{D,AB} \equiv \frac{[\text{Ab}][\text{S1}]}{[\text{S1Ab}]}
\]

\[
K_{D,ACE2} \equiv \frac{[\text{ACE2}][\text{S1}]}{[\text{S1ACE2}]}
\]

\[
[\text{Ab}], [\text{S1}], [\text{ACE2}], [\text{S1Ab}], \text{and [S1ACE2]} \text{are equilibrium concentrations of the antibody, S1, ACE2, S1–antibody complex, and S1–ACE2 complex, respectively.}
\]

\[
n_{\text{Ab}}[\text{Ab}]_0 = [\text{Ab}] + [\text{S1Ab}]
\]

\[
n_{\text{ACE2}}[\text{ACE2}]_0 = [\text{ACE2}] + [\text{S1ACE2}]
\]

\[
n \text{can be used to determine binding stoichiometry.}
\]

\[
[S1]_0 = [\text{S1}] + [\text{S1Ab}] + [\text{S1ACE2}]
\]

Substitution of eqs 6 and 7 into eqs 4 and 5, respectively, yields:

\[
[S1\text{Ab}] = \frac{n_{\text{Ab}}[\text{Ab}]_0[S1]}{K_{D,AB} + [\text{S1}]}
\]

\[
[S1\text{ACE2}] = \frac{n_{\text{ACE2}}[\text{ACE2}]_0[S1]}{K_{D,ACE2} + [\text{S1}]}
\]

Insertion of eqs 9 and 10 into eq 8 leads to:

\[
[S1]_0 = [\text{S1}] + \frac{n_{\text{Ab}}[\text{Ab}]_0[S1]}{K_{D,AB} + [\text{S1}]} + \frac{n_{\text{ACE2}}[\text{ACE2}]_0[S1]}{K_{D,ACE2} + [\text{S1}]}
\]

Rearrangement gives a cubic equation:

\[
[S1]^3 + p[S1]^2 + q[S1] + r = 0
\]

The coefficients are:

\[
p = K_{D,AB} + K_{D,ACE2} + n_{\text{Ab}}[\text{Ab}]_0 + n_{\text{ACE2}}[\text{ACE2}]_0 - [S1]_0
\]

\[
q = K_{D,ACE2}(n_{\text{Ab}}[\text{Ab}]_0 - [S1]_0) + K_{D,AB}(n_{\text{ACE2}}[\text{ACE2}]_0 - [S1]_0)
\]

\[
r = -K_{D,AB}K_{D,ACE2}[S1]_0
\]

The following root is the only physically meaningful solution:

\[
[S1] = \frac{-p + \frac{2}{3} \sqrt{p^3 - 3q \cos \frac{\theta}{3}}}{3}
\]

in which:

\[
\theta = \arccos \left( \frac{-2p^3 + 9pq - 27r}{2\sqrt{(p^3 - 3q)^3}} \right)
\]

Equation 16 describes the equilibrium concentrations of unbound S1 using the total concentrations of the individual components, two \(K_D\) values, and \(n\).

**Global Analysis of Binary and Ternary Binding Experiments.** To extract equilibrium concentrations and corresponding bound fractions of ACE2 and S1 in the ternary mixture with NAb, we globally fitted (NLSQ) binary (ACE2–S1 and S1–NAb) and ternary (ACE2–S1–NAb) experiments. The binary experiments were fitted in terms of eq 2, while the ternary experiment was fitted in terms of eqs 13–17. \(K_{D,ACE2}, K_{D,NAb}\), and \(R_{h,free}\) were set to be global fitting parameters, while all other parameters were fitted locally. The stoichiometry, \(n\), was set to 1, as concentrations refer to binding sites.

**Origin of Serum Samples.** Anti-SARS-CoV-2 seropositive human serum samples (convalescent) were obtained from BioIVT. BioIVT sought informed consent from each subject or from the subject’s legally authorized representative and appropriately documented this in writing. All samples were collected under IRB-approved protocols.

**Microfluidic Antibody-Affinity Profiling of Serum Antibodies (MAAP).** MAAP was used to determine concentration of antibody-binding sites, \([\text{Ab}]\), and \(K_D\) of antibodies in serum samples of SARS-CoV-2 seropositive individuals. To do so, fluorescently labeled SARS-CoV-2 spike S1 was mixed at constant concentrations of 10 nM, 40 nM, and 100 nM with serum at increasing concentrations (1–90%) to generate three equilibrium binding isotherms per serum sample. MDS data was obtained as described above, with the only differences being that samples were incubated for 1 h at 4 °C before measurement and that PBS-Tween (0.05%) supplemented with 0.75 mM human serum albumin (Merck) was used to dilute serum samples. Autofluorescence of serum samples was determined in the absence of labeled protein and used to correct MDS data measured of binding interactions. To determine the concentration of antibody binding sites and \(K_D\), global NLSQ fitting in terms of eq 2 was used, having \(R_{h,free}\) and \(R_{k,complex}\) \([U]_0 = [\text{Ab}]\), and \(K_D\) as global fit parameters.

**Measurement of ACE2–S1 Binding Inhibition by Serum Antibodies Using Microfluidic Diffusional Sizing.** For sample 1, Alexa Fluor 647 labeled ACE2 and unlabeled S1 at concentrations of 10 nM and 30 nM, respectively, were mixed immediately with increasing concentrations of serum and incubated at 4 °C for 1 h. For sample 2, Alexa Fluor 647 labeled ACE2 and unlabeled S1 at concentrations of 5 nM and 20 nM, respectively, were mixed immediately with increasing concentrations of serum and incubated at 4 °C for 1 h. MDS data was obtained as described in the previous section.

**ASSOCIATED CONTENT**

**Supporting Information**

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsinfecdis.1c00047.

Table S1: Summary of thermodynamic parameters and 95% confidence intervals; Figure S1: Microfluidic antibody affinity profiling (binary binding model) in serum obtained from two anti-SARS-CoV-2 seropositive individuals (PDF)
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The authors declare the following competing financial interest(s): T.P.J.K. is a member of the board of directors of Fluidic Analytics. A.A. is a member of the scientific advisory committee of Fluidic Analytics. V.K., M.M.S., and G.K. are consultants for Fluidic Analytics. S.F., M.A.P., V.D., A.S.M., A.I., A.Y.M., S.R.A.D., and H.F. are employees of Fluidic Analytics.

ACKNOWLEDGMENTS
The authors thank Marine Barbaroux, Thomas Barnes, and Richard Harrison (all Fluidic Analytics) for graphic design. V.K. was funded by an NIHR fellowship (PD-2016-09-065). T.P.J.K. is grateful for financial support by the Biotechnology and Biological Sciences Research Council and the European Research Council.

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