

# Correlation of Automated Chemiluminescent Method with Enzyme-Linked Immunosorbent Assay (ELISA) Antibody Titers in Convalescent COVID-19 Plasma Samples: Development of Rapid, Cost-Effective Semi-Quantitative Diagnostic Methods

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**Background:** We investigated the utility of an automated chemiluminescent SARS-CoV-2 IgG antibody assay platform in quantifying the amount of binding antibodies present in donated convalescent plasma.

**Methods:** A total of 179 convalescent plasma units were analyzed for the presence of SARS-CoV-2 IgG antibodies using the Beckman-Coulter chemiluminescent immunoassay (CLIA) platform. The equipment-derived numerical values (S/Co ratio) were recorded. Aliquots from the same units were subjected to enzyme-linked immunosorbent assay (ELISA) that detects IgG antibodies against the receptor-binding domain (RBD) of the SARS-CoV-2 S1 protein. The relationship between ELISA titers and CLIA S/Co values was analyzed using linear regression and receiver operating characteristics (ROC) curve.

**Results:** Twenty-one samples (11.7%) had S/Co values of less than 1.0 and were deemed negative for antibodies and convalescent plasma had S/Co values between >1.0 and 5.0 (70/179, 39.1%). Fifteen units (8.4%) had negative ELISA titer. The majority of the units (95/179, 53.1%) had titers  $\geq 1:1024$ . The sensitivities of ELISA to CLIA were comparable (90.5% vs 88.3%, respectively;  $p=0.18$ ). There was positive linear correlation between CLIA S/Co values and ELISA IgG titer ( $\rho = 0.75$ ; Spearman's rank = 0.82,  $p$ -value = <0.0001). The agreement between the two methods was fair, with a  $\kappa$  index of 0.2741. Using the ROC analysis, we identified a CLIA S/Co cutoff value of 8.2, which gives a sensitivity of 90% and a specificity of 82% in predicting a titer dilution of  $\geq 1:1024$ .

**Conclusion:** The utility of automated antibody detection systems can be extended from simply a screening method to a semi-quantitative and quantitative functional antibody analysis. CLIA S/Co values can be used to reliably estimate the ELISA antibody titer. Incorporation of chemiluminescent-based methods can provide rapid, cost-effective means of identifying anti-SARS-CoV-2 antibody titers in donated plasma for use in the treatment of COVID-19 infection.

**Keywords:** COVID-19, ELISA, chemiluminescence, antibody, titers

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## Introduction

Coronavirus disease 2019 (COVID-19), caused by a novel coronavirus, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), is the cause of a pandemic that has infected over 49.7 million people worldwide with a mortality

of 2.4% to date.<sup>1</sup> Due to the record-breaking economic and social impact of COVID-19, there is a need to optimize all resources to improve outcome and minimize the spread of the virus. A key goal is to increase the availability of treatment options for the severely ill COVID-19 patients, one of which is convalescent plasma.

Initial case reports from China have shown immediate beneficial clinical results in COVID-19 patients transfused with convalescent plasma.<sup>2-4</sup> The viral loads were decreased after transfusion, coinciding with increased levels of neutralizing COVID-19 antibody. In a match-controlled study done in the United States,<sup>5</sup> patients who received COVID-19 convalescent plasma (CCP) were more likely to have improved oxygen requirement by day 14 post-transfusion as compared to those who did not receive CCP. The recipients also had significantly longer survival, especially non-intubated patients.<sup>5</sup> Another study involving approximately 5000 patients observed better survival among recipients who were given CCP units that contain high antibody levels based on chemiluminescent intensity value.<sup>6</sup>

Recent investigations on recovered COVID-19 patients revealed low antibody levels among those with mild symptoms, especially among the healthy and younger individuals.<sup>7,8</sup> Healthy and younger individuals are most likely to be eligible for blood product donation. Another study observed rapid decline in IgG antibodies among recovered COVID-19 patients with mild symptoms from the time they were screened for donation up to plasmapheresis.<sup>9</sup> Although the FDA has issued emergency use authorization (EUA) for the use of convalescent plasma, it does not strictly implement antibody testing prior to transfusion of the blood product.<sup>10</sup> This has great implication on the quality of convalescent plasma units that are available in blood centers.

Neutralizing antibody assays, considered as the gold standard for detection of anti-COVID function, are assessed using native or pseudotype virus in cellular assay.<sup>11</sup> The disadvantages of this analysis include the limited number of facilities that have the capacity and expertise to perform this assay, lengthy turn-around time and high cost.<sup>12,13</sup> However, antibody levels can also be estimated using an antibody binding assay such as an enzyme-linked immunoassay (ELISA). A study on COVID-19 patients showed good correlation between ELISA antibody titer and neutralizing antibody levels.<sup>14</sup> In particular, the linear correlation with neutralizing antibody levels was seen in ELISA assays that detected

immunoglobulins (IgG) antibodies against the receptor-binding domain (RBD) of the viral spike protein S1.<sup>14</sup> Although easier to perform than neutralization assays, ELISA assays are limited by long turn-around time and higher costs.

Chemiluminescent immunoassay (CLIA) is considered as a qualitative antibody assay that detects binding antibodies to viral antigens, similar to the principle of an ELISA. The test relies on mixing patient samples with a known viral protein, buffer reagents, and specific enzyme-labeled antibodies that allow a light-based, luminescent read-out.<sup>15-17</sup> The amount of light (radiance) emitted from each sample is relative to the number of antibodies present in a patient sample.<sup>17</sup> This type of assay can assess the presence of multiple types of antibodies, including IgG, IgM, and IgA, is automated, has a short turn-around time (~45 minutes) and can be implemented with minimal cost in already existing clinical laboratory platforms.<sup>18</sup> However, this method has not been optimized to quantify the amount of antibodies. Currently, FDA has not authorized the use of automated chemiluminescent assays for screening of antibodies in potential donors of COVID-19 convalescent plasma.

At Maimonides Medical Center, antibody testing is performed using a Beckman Coulter platform that utilizes CLIA that detects IgG antibodies against the SARS-CoV-2 RBD. The intensity value retrieved from the platform, based on the sample-to-cut-off (S/Co) ratio, reflects the amount of anti-SARS-CoV-2 antibodies present in the sample. Since this platform utilizes the RBD of the virus, there is a good chance that the results of this assay may correlate with the number of neutralizing antibodies. However, to date, no study has been published correlating the S/Co value with an ELISA or neutralization titer in the setting of COVID-19. Such a correlation would be crucial to provide a more cost-efficient and rapid antibody testing process that is readily available and can be utilized for convalescent plasma donor screening.

## Methods

### Plasma Collection

Convalescent plasma was obtained from the Blood Donation Service at Maimonides Medical Center. Convalescent plasma was donated by recovered COVID-19 patients with confirmed diagnosis via clinical laboratory test and were symptom free for at least 28 days. Some CCP donors were initially screened for the presence of antibodies via Dynex Agility<sup>TM</sup> instrument through a reference laboratory (Table

**Table I** Characteristics of Volunteer Donors and Their Donated Plasma

Characteristics	Median (IQR)/N (%)
Age, years	39.9 (30.3–49.7)
Antibody test prior to donation	
Performed	91 (50.8)
Not performed	88 (49.2)
Antibody test result prior to donation (S/Co)	2.3 (1.7–3.1)
CCP chemiluminescent results	
0.02 to 1	21 (11.7)
>1-5	71 (39.1)
>5-10	32 (17.9)
>10-20	27 (15.1)
>20	29 (16.2)
Donated plasma ELISA titers	
Negative	15 (8.4)
1:32	2 (1.1)
1:64	7 (3.9)
1:128	10 (5.6)
1:256	19 (10.6)
1:512	31 (17.3)
1:1024	35 (19.6)
1:2048	22 (12.3)
1:4096	22 (12.3)
1:8192	5 (2.8)
1:16,384	8 (4.5)
1:32,768	3 (1.9)

1). An aliquot of plasma or serum was collected and stored at  $-20^{\circ}\text{C}$  for subsequent antibody testing at the time of plasma donation. This study was approved by the IRB/Research Committee at Maimonides Medical Center. The requirement to obtain informed consent from the subjects has been waived by the IRB in accordance with 45 C.F.R. § 46.116(d) and the guidelines outlined in the Declaration of Helsinki were followed.

The plasma or serum samples were analyzed for anti-SARS-CoV-2 RBD IgG antibodies using the Beckman Coulter chemiluminescent immunoassay (CLIA) in accordance with manufacturer instructions (<https://www.beckmancoulter.com/en/products/immunoassay/access-sars-cov-2-igg-antibody-test>). This CLIA test utilizes a recombinant SARS-CoV-2 protein specific for the receptor-binding domain (RBD) of the S1 protein. Results of this assay are based on the sample signal-to-cut-off (S/Co) ratio, with values  $<1.0$  and  $\geq 1.0$  corresponding to negative and positive results, respectively. The S/Co values reflect relative levels of anti-SARS-CoV-2 IgG antibodies.

## ELISA IgG

Assay controls and serum samples were diluted to 1:64 followed by 2-fold dilution up to 1:32,768 and added to a 96-well microtiter plate (Thermo Scientific Immulon, Waltham, MA, USA) that was coated with SARS-CoV-2 recombinant RBD (Mount Sinai Medical Center, New York, NY, USA). A secondary anti-human IgG (Fab specific) antibody labeled with horse radish peroxidase (Sigma-Aldrich, St. Louis, MO, USA) was added to each well to form a specific complex of antigen-antibody bound to the plate surface. The binding reaction was then enhanced visually with SIGMAFAST™ OPD (Sigma-Aldrich, St. Louis, MO, USA) substrate generating a yellow color for positive specimens. After application of the stop solution (3M Hydrochloric acid), the color changed from yellow to orange and optical density was measured at 490 nm. When the absorbance value was greater than the cut-off value ( $\text{OD}_{490} = 0.15$ ) at a minimum dilution of 1:64, the specimen was reported as a positive result and the corresponding titer reported.

## Statistical Analysis

The analysis population included COVID-19 convalescent plasma samples collected at the Blood Donation Service, therefore, no sample size calculations were performed. Correlation between CLIA and ELISA titer was analyzed using Spearman's rank, R square and  $\kappa$  index. The sensitivity and specificity of the two methods were analyzed using a 2x2 table. Linear relationship between the two assays was also assessed using a linear regression model, though due to violation of model assumptions, both CLIA and ELISA values were transformed to maintain the normality of residuals. The square-root of the CLIA assay was used to predict the log-transformed ELISA. The outputs of this model were then transformed back into raw units to show the relationship between the assays. A logistic regression model was created, as well, to predict the probability of a positive ELISA result (titer  $\geq 1:1024$ ). A CLIA S/Co cut-off value of 8.2 was determined using the ROC curve to meet 90% sensitivity and 82% specificity.

A two-sided P value of less than 0.05 was considered to indicate statistical significance. Statistical analyses were performed using SAS version 9.4 (Cary, NC) and GraphPad Prism 8 (GraphPad Software, La Jolla, CA).

## Results

A total of 179 donated convalescent plasma units were included in this study. The inclusion criteria were mild

COVID-19 symptoms and positive RT-qPCR tests on nasopharyngeal swab samples. Females and those younger than 18 years of age were excluded. The characteristics of the donors are summarized in Table 1.

Aliquots of all units were analyzed using the Beckman Coulter CLIA platform and the relative amount of antibodies as measured by the S/Co value was recorded. Twenty-one samples (21/179; 11.7%) had S/Co values of less than 1.0 and were deemed negative for antibodies. Fifteen units that had S/Co values >1.0 (15/179; 8.4%) had negative ELISA titers. The majority of the units had a titer of  $\geq 1:1024$  (95/179, 53.1%). The sensitivity of ELISA was comparable to the sensitivity of CLIA (90.5% vs 88.3%, respectively; p-value = 0.18; Table 2). Table 3 shows the distribution of CLIA S/Co values and ELISA IgG antibody titers of all donated convalescent plasma. Samples with S/Co values of less than 10 had ELISA IgG titers ranging from 0 (negative) to 1:4096. Those with S/Co values of 10–20 had titers varying from 1:128 to 1:4096. Samples with CLIA S/Co value higher than 20 had titers of 1:2048 to 1:32,768.

Using linear regression, there is positive linear correlation between CLIA S/Co values and ELISA IgG titer (Rho = 0.75; Spearman's rank = 0.82, p-value = <0.0001; Figure 1). The agreement between the two methods was fair, with a  $\kappa$  index of 0.2741. A linear regression model was developed using the square root of the S/Co value and logarithmic transformation of the equivalent ELISA IgG titer (Table 4). Using this linear regression model, an estimate of the equivalent ELISA IgG titer when given a CLIA S/Co value can be computed using the following formula:

$$\text{Raw Titer Dilution} = e^{4.8949+0.70612\sqrt{\text{CLIA } \frac{S}{Co} \text{ value}}}$$

Table 5 provides correlation of representative CLIA intensity to its expected titer concentration. Further, a predictive statistical file was created using this formula for more exact conversions from CLIA values to expected titer determinations (Supplementary Table). This file enables the user to enter any CLIA S/Co value and will

**Table 2** Sensitivity of the Antibody Tests Compared to SARS-CoV-2 Viral RNA RT PCR

	Chemiluminescent Assay	ELISA
Sensitivity	88.3%	90.5%
No.	158/179	162/179
95% CI	82.5–91.8%	83.6–92.6%

**Table 3** CLIA S/Co Values and ELISA Titers of Donated Convalescent Plasma

CLIA S/Co Values	N (%)	ELISA IgG Titer Equivalent
0.02 to 1	21 (11.7)	Neg to 1:512
>1 to 5	70 (39.1)	Neg to 1:4096
>5 to 10	32 (17.9)	Neg to 1:4096
>10 to 20	27 (15.1)	1:128 to 1:4096
>20	29 (16.2)	1:2048 to 1:32,768
ELISA IgG Titer	N (%)	CLIA S/Co Value Equivalent
Neg	15 (8.4)	0.02 to 6.25
1:32	2 (1.1)	0.37 to 0.5
1:64	7 (3.9)	0.09 to 9.71
1:128	10 (5.6)	0.25 to 11.51
1:256	19 (10.6)	0.05 to 7.66
1:512	31 (17.3)	0.81 to 19.65
1:1024	35 (19.6)	1.2 to 15.52
1:2048	22 (12.3)	2.2 to 26.62
1:4096	22 (12.3)	1.92 to 44.12
1:8192	5 (2.8)	23.48 to 34.26
1:16,384	8 (4.5)	28.08 to 63.4
1:32,768	3 (1.9)	28.82 to 52.64

**Abbreviations:** CLIA, chemiluminescent immunoassay, S/Co, sample to control intensity ratio, ELISA, enzyme-linked immunosorbent assay, Neg, negative, "0."

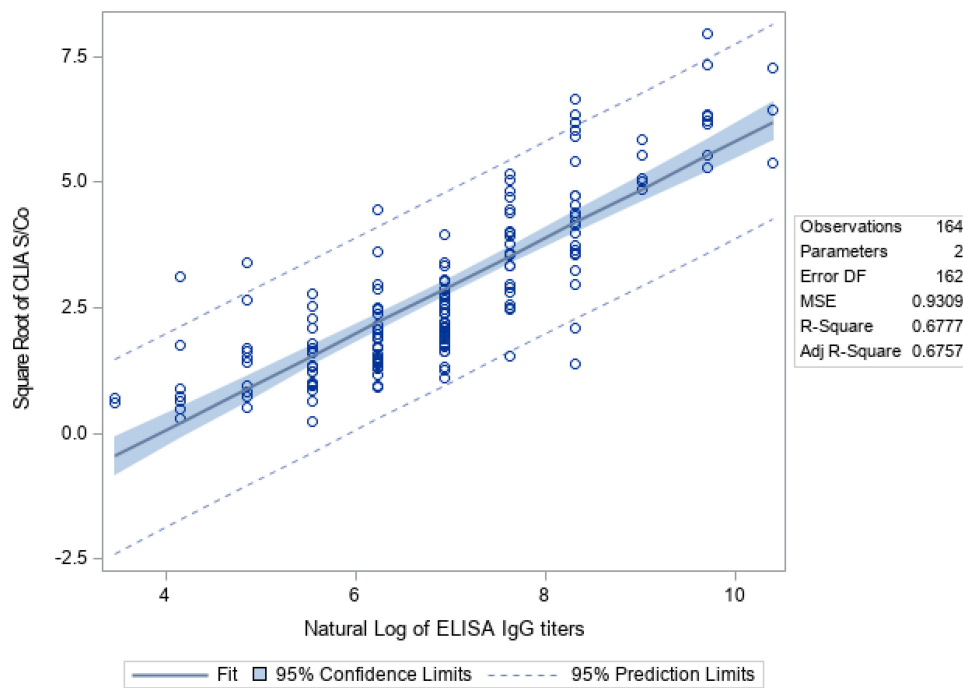
automatically generate the equivalent ELISA IgG titer with 95% confidence intervals.

A receiver operating characteristic (ROC) curve analysis was utilized to determine the optimal CLIA S/Co value that will reliably correspond to a high ELISA IgG titer (>1:1024). This analysis identified that a CLIA S/Co cut-off value of 8.2 gives a sensitivity of 90% and a specificity of 82% in predicting a titer dilution of more than 1:1024 (Figure 2).

## Discussion

Our results showed significant linear correlation between the CLIA S/Co values and ELISA IgG titers. This indicates that comparable antibody quantification can be reliably derived from the results of an automated antibody detection system. This observation agrees with the conclusion of a similar study that analyzed the quantification capability of 8 commercially available immunoassays, two of which were CLIA-based.<sup>19</sup> They found that the automated systems provided the highest sensitivities (up to 98%), and one of the CLIA platforms had the best overall quantitative correlation to the neutralization titer (Rho=0.729).

Neutralization assays are the gold standard for assessing specific immunity and a benchmark for



**Figure 1** Linear regression analysis on the relationship of CLIA S/Co values and ELISA IgG titers.

other antibody assays. These tests are very complex, require incubation times of 5–7 days and a biosafety level 3 laboratory, limiting the routine use of these assays on a large scale.<sup>20,21</sup> Previous studies have shown positive correlation between neutralizing titer and ELISA that detects IgG antibodies against the SARS-CoV-2 S1 protein<sup>19</sup> or IgG antibodies against the receptor-binding domain of the S1 protein.<sup>14</sup> Similar studies correlating CLIA S/CO with ELISA assessment have been reported in other infectious disease spaces such as hepatitis,<sup>22</sup> measles<sup>23</sup> mycoplasma<sup>24</sup> and HIV<sup>25</sup> for diagnosis as well as donor infectious disease screening, where applicable. In a similar manner, the results of this study demonstrated correlation between ELISA antibody titer and CLIA S/Co values. Since ELISA titers correlate with the amount of neutralizing antibody, our results extend

this association to support the use of automated CLIA-based systems as a fast and convenient method of estimating neutralizing antibody quantities in convalescent plasma.

Only about a third of convalescent plasma units collected at our center had S/Co values of at least 8.2 and antibody titers of more than 1:1024. Preliminary evidence with COVID-19 suggests that patients with mild symptoms may develop very low titer antibodies.<sup>14,26,27</sup> One study observed that 18% of convalescent COVID-19 donors had undetectable neutralizing titers in their plasma samples collected an average of 30 days after the onset of symptoms.<sup>28</sup> A larger study performed by NYBC showed that more than half of plasma donors had low neutralizing titers and that there was a large variation in antibody titers among donors.<sup>29</sup> There is an existing challenge to screen convalescent plasma donors for antibody titers, as

**Table 4** Linear Regression Model of the Root of Intensity as a Predictor of the Logarithmic Value of ELISA IgG Titer

Parameter Estimates							
Variable	DF	Parameter	Standard	t Value	Pr >  t	95% Confidence Limits	
		Estimate	Error				
Intercept	1	4.89496	0.12611	38.82	<0.0001	4.64593	5.14399
Root of CLIA S/Co	1	0.70612	0.03826	18.45	<0.0001	0.63056	0.78168

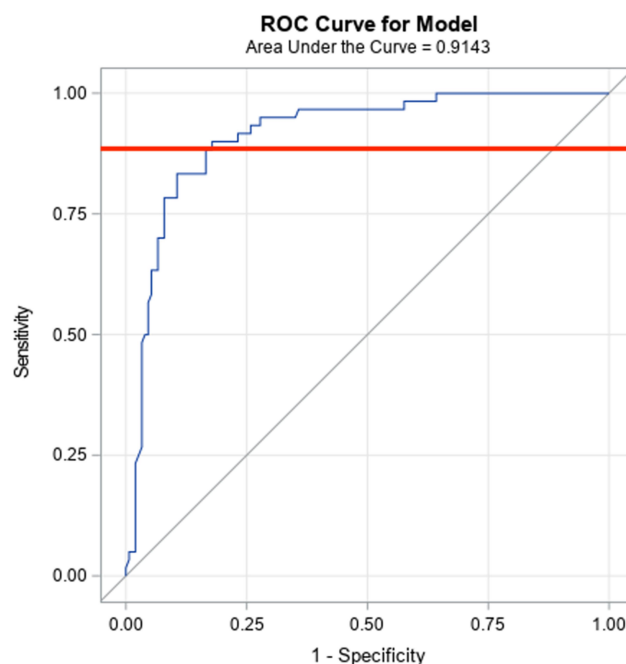
**Table 5** Representative CLIA Intensity to Expected Titer Concentrations

CLIA S/Co Values	Estimated ELISA IgG Antibody Titer		
	Estimated ELISA IgG Antibody Titer, Mean	Lower 95%	Upper 95%
0	1:133.6	1:104.2	1:171.4
1	1:270.7	1:195.7	1:374.5
2	1:362.7	1:254.1	1:517.7
3	1:454.0	1:310.5	1:663.7
4	1:548.5	1:367.6	1:818.4
5	1:648.0	1:426.6	1:984.2
6	1:753.4	1:488.1	1:1162.9
7	1:865.4	1:552.4	1:1355.8
8	1:984.5	1:619.8	1:1563.9
9	1:1111.3	1:690.6	1:1788.3
10	1:1246.3	1:765.0	1:2030.2
15	1:2058.5	1:1197.6	1:3538.4
20	1:3142.7	1:1747.4	1:5652.0
25	1:4562.2	1:2437.5	1:8538.9
30	1:6390.3	1:3293.3	1:12,399.7
35	1:8711.7	1:4343.2	1:17,474.0
50	1:19,692.3	1:8997.1	1:43,101.0
60	1:31,714.9	1:13,769.7	1:73,046.9
70	1:49,157.5	1:20,365.0	1:118,657.2

**Abbreviations:** CLIA, chemiluminescent immunoassay, S/Co, sample to control intensity ratio, ELISA, enzyme-linked immunosorbent assay.

evidence shows that the majority of mild COVID-19 patients may not develop an adequate level of anti-SARS - CoV-2 antibodies to provide a therapeutic benefit. Screening of donated convalescent plasma units for antibody is, therefore, an essential step to ensure the efficacy of convalescent plasma as a therapeutic product with viral-neutralizing capacity.

Testing for antibodies should be performed to first confirm the presence of antibodies. Subsequent identification of CCP units that provide higher concentrations of anti-COVID (neutralizing) antibodies may provide a more appropriate logic of selecting optimal CCP units for patient administration. The utility of automated antibody detection systems can be extended



**Figure 2** Receiver operating characteristic (ROC) curve showing an area under the curve (AUC) of 0.9143. The orange line denotes a CLIA S/Co cutoff value of 8.2.

from simply a screening method to a semi-quantitative and quantitative functional antibody analysis. CLIA S/Co values can be used to estimate the ELISA antibody titer, and this can immediately provide crucial information on dosing of convalescent plasma. As a semi-quantitative method, a CLIA S/Co cut-off value of 8.2 can be used to reliably detect convalescent plasma donors or units with more than 1:1024 IgG antibody titer (90% sensitivity; 82% specificity); this value is well above the recently published FDA guidance of  $S/Co \geq 3.3$  indicative of high titer CCP further relating the values to functional anti-viral titer-based activity using alternative instrumentation and/or methodologies.<sup>30</sup>

Application of additional approaches to assess the anti-COVID antibody concentration in CCP is paramount. In addition, methodological differences can introduce differences in results. For example, chemiluminescence methodology is able to determine total Ig (IgM, IgG, IgA) whereas ELISA determines only IgG. This may very well contribute towards the observed differences when a sample may have high percentage of IgM (giving high CLIA number), and low IgG (low titer) and vice versa. Although certain instruments have been promoted and approved to serve this need, many institutions and blood centers lack access to such instruments thereby causing a bottleneck in reference lab identification of

appropriate products for transfusion. Expansion to additional CLIA instrumentation can reduce the strain of CCP assessment and offer the ideal plasma product for patients who require such.

Furthermore, the methodology employed by CLIA instrumentation could be used to examine response to COVID-19 vaccination and, depending on target, provide distinction between immune response to COVID-19 infection from the immune response to vaccination.

The rapid turnover and cost benefit of such a CLIA based screening approach provides additional value. The testing can be easily implemented into existing chemiluminescent systems and provides opportunity costs for run time and personal skill set allocation. For example, in certain cases, time to result can be reduced from 180 minutes (ELISA) to 30 minutes (CLIA). In addition, adaptation can be automated in a CLIA setting and may require a lower degree of operator expertise (particularly if automated) than with ELISA.<sup>22</sup> This can beneficially impact the operational logic of the clinical laboratory to optimize efficiency of motion, human capital as well as resource allocation.

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## Disclosure

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