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Dual lateral flow optical/chemiluminescence immunosensors for the rapid detection of salivary and serum IgA in patients with COVID-19 disease

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ABSTRACT
To accurately diagnose COVID-19 infection and its time-dependent progression, the rapid, sensitive, and noninvasive determination of immunoglobulins A specific to SARS-CoV-2 (IgA) in saliva and serum is needed to complement tests that detect immunoglobulins G and M. We have developed a dual optical/chemiluminescence format of a lateral flow immunoassay (LFIA) immunosensor for IgA in serum and saliva. A recombinant nucleocapsid antigen specifically captures SARS-CoV-2 antibodies in patient specimens. A labelled anti-human IgA reveals the bound IgA fraction. A dual colorimetric and chemiluminescence detection enables the affordable and ultrasensitive determination of IgA to SARS-CoV-2. Specifically, a simple smartphone-camera-based device measures the colour signal provided by nanogold-labelled anti-human IgA. For the ultrasensitive chemiluminescence transduction, we used a contact imaging portable device based on cooled CCD, and measured the light signal resulting from the reaction of the HRP-labelled anti-human IgA with a H2O2/luminol/enhancers substrate. A total of 25 serum and 9 saliva samples from infected and/or recovered individuals were analysed by the colorimetric LFIA, which was sensitive and reproducible enough for the semi-quantification of IgA in subjects with a strong serological response and in the early stage of COVID-19 infection. Switching to CL detection, the same immunosensor exhibited higher detection capability, revealing the presence of salivary IgA in infected individuals. For the patients included in the study (n = 4), the level of salivary IgA correlated with the time elapsed from diagnosis and with the severity of the disease. This IgA-LFIA immunosensor could be useful for noninvasively monitoring early immune responses to COVID-19 and for investigating the diagnostic/prognostic utility of salivary IgA in the context of large-scale screening to assess the efficacy of SARS-CoV-2 vaccines.

1. Introduction

The SARS-CoV-2 (COVID-19) pandemic has highlighted the importance of rapid, specific, and accurate diagnostic tests in limiting the spread of infection and monitoring patients’ viral load and therapy. Around 200 diagnostic tests have been developed to detect the RNA of SARS-CoV-2 (through reverse-transcriptase polymerase chain reaction, rt-PCR). Other tests work by determining antibodies specific to the virus in serum following immune response. A common format for point-of-care detection of the immune response to a virus is to measure virus-specific antibodies (IgM and IgG, or in combination) in serum using the lateral flow immunoassay (LFIA), where gold nanoparticles are used to label biospecific reagents such as secondary antibodies.

A typical antibody response to exposure to antigen involves the primary humoral immune responses typified by the appearance of IgM within the first three to five days following the exposure, followed by IgG production within the first week (Morris and Gronowski, 2010). IgG persist after the virus is no longer detectable, indicating previous infection, while IgM are transient, so their presence is associated to a recent infection. However, the production of IgM has been reported as

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protein was selected among antigenic targets of the SARS coronavirus capturing reagent and anchored onto the detection membrane. The N structure in a previous work of the group because it is highly immunogenic and abundantly expressed (Zeng et al., 2020) and according to its antigenic and abundantly expressed (Zeng et al., 2020) and according to its

As a complement to IgG detection, one significant advantage of targeting IgA is the possibility of using saliva instead of blood for the analysis. Salivary anti-SARS-CoV-2 IgA have been shown to correlate with serum amounts (Randad et al., 2020). Salivary collection has several advantages over blood withdrawal, especially for point-of-care testing (Ong et al., 2020). Saliva can be collected easily by the patient, reducing the risk associated with contact between operator and patient. Furthermore, saliva collection is particularly suitable for babies and elderly people, and for cheap patient population screening in low-resource settings. However, at present, there are no rapid tests for detecting SARS-CoV-2-specific IgA in saliva.

The rapid and specific detection of serum and salivary IgA could deliver early and hopefully time-dependent information about the infection. In particular, due to inconsistent findings about the evolution of IgM levels during infection, a serological marker of recovery is needed to reduce the number of rRT-PCR analyses and support decision-making about ending quarantine. Moreover, a portable easy-to-use test for serum and salivary IgA could help evaluate the individual response to therapy or vaccination against the virus in large populations.

For IgA analysis, we developed an LFIA based on a SARS-CoV-2-specific antigen (the nucleocapsid protein N), which is used as the capturing reagent and anchored onto the detection membrane. The N protein was selected among antigenic targets of the SARS coronavirus structure in a previous work of the group because it is highly immunogenic and abundantly expressed (Zeng et al., 2020) and according to its reactivity to human sera from COVID-19 patients (Cavalera et al., 2020). The optical immunosensor includes an anti-human IgA (anti-IgA) labelled with gold nanoparticles (GNP) to reveal the IgA bound to the N antigen. The anti-SARS-CoV-2 IgA in the serum/saliva sample is captured by the N antigen and stained by the GNP-labelled anti-IgA, forming a coloured band at the test line. For signal transduction and quantification of the GNP-based LFIA, we imaged the coloured strip using a smartphone camera. We reported the results in RGB scale under optimized reading conditions using the smartphone flash, as previously reported (Calabria et al., 2017). To achieve higher detectability compared to the coloured GNP probe, we used the same LFIA format, but with chemiluminescence (CL) detection mediated by a horseradish peroxidase (HRP) labelled anti-IgA and an enhanced CL luminol-/H₂O₂/enhancer substrate (De Nardo et al., 2016). The CL signal to noise was improved compared to the previous system by adding the CL substrate solution directly to the control and test line area after the LFIA run, thus minimizing the nonspecific light signal. The CL emitted light is measured by an ultrasensitive cooled CCD in contact imaging mode, with the data reported in relative light units.

Both immunosensor platforms allow data recording and can be used for comparative evaluation within the same patient to monitor the presence of IgA in saliva and/or serum and connect this data to disease progression and a possible decrease in viral load. The noninvasive collection of saliva is a further strength of this test. This work describes the first dual LFIA platform for the rapid detection of salivary and serum IgA and illustrates its application for SARS-CoV-2 infection.

2. Materials and methods

2.1. Reagents and materials

Gold (III) chloride trihydrate (ACS reagent), mouse antihuman immunoglobulin A monoclonal antibody A (a-chain specific), protein A from Staphylococcus aureus (SpA), sucrose, and bovine serum albumin (BSA) were obtained from Merck group (Darmstadt, Germany). Tween 20 and other chemicals were purchased from VWR International (Milan, Italy). Nitrocellulose membranes with cellulose adsorbent pad, blood separator, and saliva-specific sample pads were purchased from MDI Membrane Technologies (Ambala, India). Glass fiber conjugate pads were obtained from Merck Millipore (Billerica, MA, USA). HRP-labelled mouse antihuman IgA were obtained from Invitrogen, Thermo Fisher Scientific (Rockford, IL). The Supersignal ELISA Fermo CL substrate for HRP was purchased from Thermo Fisher Scientific Inc. (Rockford, IL).

2.2. The IgA-LFIA strip

The LFIA strip for the colorimetric IgA-LFIA biosensor is schematized in Fig. 1a. The nucleocapsid (N) antigen (1 mg/ml in phosphate buffer 20 mM pH 7.4) and staphylococcal protein A (SpA, 0.5 mg/ml in phosphate buffer) were spotted at 1 μl/cm by means of a XYZ3050 platform (Biodot, Irvine, CA, USA) to form the test (TL) and control (CL) lines, respectively. The preparation of the recombiant nucleocapsid protein was previously described in Cavalera et al. (2020) and is detailed in the SI.

For the optical IgA-LFIA, gold nanoparticles (GNP) of ca. 30 nm diameter and SPR band centered at 525 nm were synthesised by tetra-aurothiuracil reduction and conjugated to a murine anti-human IgA by passive adsorption, as previously reported (De Nardo et al., 2017). Briefly, the anti-IgA was added to a pH-adjusted GNP solution (pH 8.5), in the proportion 10 μg per ml of GNP (optical density, OD 1). The unconvered GNP surface was saturated with BSA and the GNP-anti IgA were concentrated and recovered by centrifugation. GNP-labelled anti-IgA were pre-adsorbed in the conjugate pad (0.1 ml/cm). Sample pad, conjugate pad, the membrane, and adsorbent pad were overlapped, and strips were cut (4 mm-width). The strips were inserted into plastic cassettes.

For the chemiluminescence detection, strips were prepared as described above, except for the detection reagent (anti-human IgA-HRP, from Sigma-Aldrich), which was pre-adsorbed onto the conjugate pad as diluted 1:1000 with phosphate buffer. In addition, the membrane was saturated with 1% BSA after line deposition. In this case, the cassette was not used to maximise the contact between the strip and the CL reader.

2.3. Optical LFIA to detect IgA specific to SARS-CoV-2

Serum and saliva were diluted by 1:10 and 1:5 v/v with Tris-glycine buffer.
buffer 0.1 M (pH 8, with 0.2% casein and 1% Tween 20 added), respectively. 80 μl of diluted specimen were used and LFIA results were visually inspected at 15 min from sample application. For the (semi)-quantitative evaluation of TL colour, the LFIA strip was placed in front of the back-illuminated CMOS based camera, inside the mini dark box to exclude ambient light, and an additional lens was used to focus the T and C line image and standardize the reading using the smartphone flash illumination. A semicover and a mini dark box adaptable to any smartphones were made with 3D printing (Fig. 1b). Images were then digitally processed using an RGB scale to quantify the colour. The setup of the apparatus and image processing are described in further detail in the SI.

2.4. CL-LFIA to detect IgA anti SARS-CoV-2

For the chemiluminescence detection, we developed a simple device based on a cooled CCD camera with the LFIA strip in contact with the sensor using a fibre optic faceplate. The scheme of the device is reported in Fig. 2a and further detailed in the SI. The assay was carried out in a similar way to the colorimetric assay. At 15 min from sample application, the activity of the HRP labelled antibody was measured by overlying a transparent glass fibre pad on the detection membrane (Fig. 2b).
The pad contained freeze-dried sodium perborate, luminol, and p-iodophenol (Deng et al., 2016). Once in contact with the LFIA strip and following the addition of 20 μl of water, this delivered the CL substrate with production of light (Zangheri et al., 2015). The cooled CCD reader and device were placed in contact with the strip, imaged, and the CL analytical signal was quantified and expressed in relative light units (RLU).

3. Results and discussion

3.1. IgA-LFIA to detect IgA specific to SARS-CoV-2 in serum

We made an optical LFIA prototype to selectively detect anti-SARS-CoV-2 IgA and verified its ability to detect the target immunoglobulins in the serum of COVID-19-infected individuals. The immunosensor included the recombinant nucleocapsid protein from SARS-CoV-2 to capture antibodies specific to the virus, and an anti-IgA labelled with GNP as the probe (Fig. 1a). The test line (TL) was coloured in the presence of anti-SARS-CoV-2 IgA in the specimen because these interacted with the immobilized N antigen and were stained by the GNP probe. SpA, which captured the excess of the labelled anti-IgA used as the control line (CL) to confirm the validity of the assay. The diagnostic specificity of the IgA-LFIA was checked by analysing ten serum samples that did not contain any SARS-CoV-2-specific antibodies, as they were collected before the COVID-19 outbreak. No false positive results were recorded (false positive rate = 0/10). Regarding positive samples, we analysed 25 human sera from infected individuals as confirmed by rt-PCR. Blood samples were collected at variable times from the diagnosis, and some individuals (n = 17) recovered in the meantime. Of the 25 serum samples analysed by the IgA-LFIA, 15 showed colouring of the TL and were then assigned as IgA positive. Compared to the rt-PCR method, the false negative rate was calculated as 40.8% (10/25). Interestingly, plotting the number of IgA positive samples on the timeframe from infection diagnosis and recovery showed two clear patterns (Fig. S1a in the SI). Although the number of analysed samples is not sufficient to draw conclusions, we speculate that IgA increased during the second week of infection, peaked at the fourth week, and then declined. This trend is congruent with that reported by Padoan et al. (2020). We also observed that the IgA level in serum dramatically dropped a few days after recovery (Fig. S1b in the SI).

Repeatability of the IgA-LFIA was studied by analysing, in duplicate, three serum samples classified as positive and three as negative, then calculating the mean relative standard deviation (RSD%). Stability was investigated by analysing one positive and one negative serum sample at 0, 7, and 28 days from IgA-LFIA device construction. The quantification of TL colour of positive results by the smartphone camera showed that the colorimetric IgA-LFIA provided sufficiently repeatable and stable results (RSD% for duplicate analysis of three positive samples were below 15%, and the TL intensity varied within 10% over four weeks from IgA-LFIA construction).

3.2. IgA-LFIA to detect anti-sars-cov-2 IgA in saliva

The prototype IgA-LFIA for serum was then adapted and optimized for use with saliva specimens. To this end, the blood separator sample pad included in the original device was replaced by a new sample pad recommended by the manufacturer for application to saliva and oral fluid specimens.

In previous studies, we demonstrated that replacing the optical detection of GNP with chemiluminescence detection of HRP as a label increased the detectability of LFIA assays by a factor of ten (Zangheri et al., 2015). We also showed that detectability could be further increased by using a more sensitive CL reader based on cooled CCD instead of the smartphone camera (Calabria et al., 2017). Therefore, we modified the IgA-LFIA to enable CL detection. Although increasing the sensitivity, the CL detection required an additional step to add the CL substrate. The flow of the CL substrate across the strip produced a strong background light, which affected the signal-to-noise ratio and largely increased the analysis time. In order to overcome these limitations, we designed a semi-integrated system in which the dried CL substrate was embedded in a glass fibre pad. The pad was layered onto the detection zone of the strip, after completion of sample run, so that the CL substrate was quickly dissolved by the wet LFIA. To help the CL substrate resuspension, we also added a drop of water. This innovative and effective strategy was adapted from Deng J. et al. (2018), who set up a self-contained system, by which CL reagents were stored in dried form and were delivered directly on the detection zone by a microfluidic system aimed at revealing the enzymatic amplification of nucleic acids. Here, the addition of the CL in a very confined zone showed clear advantages over the traditional flow strategy as the CL substrate was
rapidly and specifically made available for the enzymatic reaction. The background light was strongly reduced thus increasing the signal-to-noise ratio. In addition, by avoiding running the CL substrate by capillarity, the assay was accelerated, completing in 15 min instead of in over 30 min.

The IgA-LFIA prototypes were applied to analyse nine salivary samples. Eight were from four COVID-19 infected individuals and were collected at two and four weeks from rt-PCR diagnosis (Table S1 in the SI). One sample was collected in February for another study from a donor who, in that period, showed symptoms compatible with COVID-19 (cough, fever, fatigue, difficulty breathing), but without confirmation of COVID-19.

The detectability of both colorimetric and CL IgA-LFIA was sufficient to reveal salivary IgA in three out of the five subjects with good agreement between the detection methods (Fig. 3a and b). In particular, the IgA-LFIA with both colour and CL detection revealed very high salivary IgA levels in subject #1, who was known to have a severe disease. Salivary IgA to SARS-CoV-2 were present at 2 and 4 weeks from diagnosis, and confirmed the tendency for levels to decrease over the time from infection observed for serum samples.

Subject #2, who reported moderate symptoms, showed IgA levels lower than #1. However, they were still detectable, especially by the ultrasensitive CL immunosensor. For subject #2, the intensity of the colour signal increased with time, while the CL detection confirmed the trend of IgA levels decreasing over time.

Subjects #3 and #4 provided results that were undetectable by colorimetric detection (a slight signal, close to the limit of detection, was shown for subject #3 only after 2 weeks from infection). In contrast, the CL detection highlighted the presence of salivary IgA in these subjects, at least in the early stage of the disease (week 2).

The improved detectability of the CL immunosensor was confirmed by the “unknown” sample. The results were at the limit of detection for colorimetric IgA-LFIA, but were clearly positive with CL detection (Fig. 3c).

For the subjects with a known clinical condition, the results obtained by the IgA-LFIA correlate with severity of symptoms and also with serum IgG and IgM levels, as measured by a reference assay (Table S1 in the SI). Moreover, the new IgA-LFIA suggested that the ‘unknown’ sample belonged to a subject that was presumably infected in an early phase of the pandemic. These few data suggest that salivary IgA can be quantified using both detection strategies.

The colorimetric immunosensor coupled with the smartphone reading enabled the one-step affordable determination of IgA levels in saliva. The introduction of a simple and portable tool to quantify the colour at the test line, which is related to amount of IgA, can help providing information on the severity and/or stage of the infection.

The CL detection using the portable cooled back illuminated CCD camera provided higher sensitivity and more accurate quantification compared to the smartphone BI-CMOS allowing to detect positivity in subjects with low serological response. Moreover, compared to previously reported CL-LFIA, we improved the signal-to-noise ratio by adding the CL substrate directly on the detection zone instead of flowing it...
across the LFIA strip. This modification of the protocol strongly reduced the light background and increasing detectability. The new protocol also halved the time required to complete the assay (15 min instead of 30 min), which is a major improvement for point-of-care testing. The combination with an optical/chemiluminescence transduction device also provides the option of connectivity for promptly communicating the patient’s infection and/or recovery status.

Although IgG and IgM specific to SARS-CoV-2 have been reported to be present in saliva, as well as IgA, and can then be considered as possible markers for the non-invasive detection of the serological response to the infection (Randad et al., 2020; Isbo et al., 2020), to the best of our knowledge, this is the only immunosensor for detecting salivary IgA reported to date. This is because almost all the rapid serology tests focus on serum IgM, IgG, and total immunoglobulins. However, the lack of information regarding the clinical meaning of salivary IgA measurements and the intrinsic variability of the composition of the oral fluids, which can affect the result especially in quantitative analysis, are current limitation of the method and needs further investigation. The validation of the developed IgA-LFIA by considering a larger number of samples is ongoing. Hopefully, results will be also double-checked by lab-scale tests to measure IgA (though methods for salivary IgA are still unavailable).

The availability of this rapid test may enable additional largescale studies on the significance of IgA as biomarkers of immune response to COVID-19 and the noninvasive screening to assess the efficacy of new vaccines. In the context of precision medicine, it could also support a personalized therapeutic intervention.

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CRediT authorship contribution statement
Aldo Roda: Conceptualization, Methodology, for chemiluminescence LFIA, Writing - original draft, preparation. Simone Cavalera: Investigation, for visual LFIA, Writing - original draft, preparation. Fabio Di Nardo: Methodology, for visual LFIA. Donato Calabria: Methodology, for chemiluminescence LFIA. Barbara Colitti: resource (ARS-CoV-2 recombinant nucleocapsid protein). Patrizia Simonini: Methodology, for chemiluminescence LFIA. Claudio Baggiani: Funding acquisition, and project management. Matilde Roda: Resource (clinical samples and diagnosis). Laura Anfossi: Conceptualization, Formal analysis, Writing - review & editing. All authors have read and agreed to the published version of the manuscript.

Declaration of competing interest
The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data
Supplementary data to this article can be found online at https://doi.org/10.1016/j.bios.2020.112765.

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