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Salivary SARS-CoV-2 antigen rapid detection: A prospective cohort study

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ABSTRACT

Background and aim: SARS-CoV-2 quick testing is relevant for the containment of new pandemic waves. Antigen testing in self-collected saliva might be useful. We compared salivary and naso-pharyngeal swab (NPS) SARS-CoV-2 antigen detection by a rapid chemiluminescent assay (CLEIA) and two different point-of-care (POC) immunochromatographic assays, with results of molecular testing.

Methods: 234 patients were prospectively enrolled. Paired self-collected saliva (Salivette) and NPS were obtained to perform rRT-PCR, chemiluminescent (Lumipulse G) and POC (NPS: Fujirebio and Abbott; saliva: Fujirebio) for SARS-CoV-2 antigen detection.

Results: The overall agreement between NPS and saliva rRT-PCR was 78.7%, reaching 91.7% at the first week from symptoms. SARS-CoV-2 CLEIA antigen was highly accurate in distinguishing positive and negative NPS (ROC-AUC = 0.939, 95\%CI:0.903–0.977), with 81.6% sensitivity and 93.8% specificity. This assay on saliva reached the optimal value within 7 days from symptoms onset (Sensitivity: 72%; Specificity: 97%). Saliva POC antigen was limited in sensitivity (13%), performing better in NPS (Sensitivity: 48% and 66%; Specificity: 100% and 99% for Espline and Abbott respectively), depending on viral loads.

Conclusions: Self-collected saliva is a valid alternative to NPS for SARS-CoV-2 detection by molecular, but also by CLEIA antigen testing, which is therefore potentially useful for large scale screening.

1. Introduction

Saliva testing for SARS-CoV-2, one of the strategies for COVID-19 diagnosis and monitoring, is advocated mainly for screening asymptomatic subjects in order to rapidly detect and isolate infected individuals and their contacts, thus limiting viral spread and containing further waves of the pandemic \cite{1–6}. Although the molecular detection of SARS-CoV-2 RNA in naso-pharyngeal swabs (NPS) is considered the “gold standard” technique for identifying symptomatic or asymptomatic individuals \cite{7}, it has limitations in both the analytical and the healthcare settings. From the analytical viewpoint, it is widely agreed that the sensitivity of rRT-PCR of NPS ranges from 70 to 90\% \cite{5,8}, reaching values around 50\% after the first two weeks of disease \cite{9,10}. Therefore, COVID-19 disease cannot be ruled out when a NPS result is negative, but the patient has clinical symptoms, and biochemical data and radiological findings that evidence a clinical scenario typical of the disease \cite{11}. In this context anti-SARS-CoV-2 antibodies should also be taken into account \cite{12}. From the healthcare organizational viewpoint, NPS testing calls for the involvement of healthcare workers and services for sample collection. This pre-requisite might be promptly met by the healthcare system when the demand is low, but not necessarily when it is high, as occurs in a pandemic. Any delay in testing puts individuals without a diagnosis at risk, consequently exposing the community to viral contagion.

Saliva testing might not only have the advantage of relieving health care resources, but also of reducing hazard exposure to healthcare workers during sampling; it might also limit the risk of viral spread incurred when numerous individuals queue for a long time waiting for...
testing, since saliva can be self-collected at home. However, none of these advantages support the use of saliva testing if its results are not as reliable as those of NPS. When rRT-PCR is used for SARS-CoV-2 testing, the reliability of saliva testing is reportedly equal to, or even higher than, NPS [5,10,13,14], although some studies report contradictory findings (i.e. saliva as less sensitive) probably depending on salivary viral load kinetic [3,9,14–16].

Saliva testing by rRT-PCR is reliable, but time consuming, calling for dedicated laboratory equipment for nucleic acid extraction and amplification, and personnel trained in molecular techniques [17]. These requirements, which might not be fulfilled by all laboratories, compromise the advantage of using the saliva sample. In front of a safe and rapid collection procedure, the overall testing process remains long not only because molecular testing takes time, but also because molecular laboratories might be limited in number, especially in low resource countries, thus increasing the turnaround time due to sample transportation and processing. In order to speed up testing while maximizing the number of tested individuals, the search for SARS-CoV-2 antigens rather than RNA, by immunometric techniques is now emerging, including point-of-care (POC) rapid immunochromatographic assays based on lateral flow technology [18]. The market now offers a number of SARS-CoV-2 antigen detection immunomagnetic assays, which are high throughput but require laboratory instrumentation [19], and ultra-rapid POC devices suggested for use in medical cabinets by general practitioners and nurses. Since they are simple to use, these devices are also considered potentially employable in auto-testing. However, simplicity is not synonymous with accuracy [20]. To date, no exhaustive data are present in the literature on antigen detection using saliva, an approach that might maximize an effective and timely COVID-19 diagnosis, encompassing the advantages of a) saliva self-collection and b) rapid viral protein detection, thus making wide-scale screening possible in many parts of the world.

The aim of this prospective study was to compare in saliva and NPS the diagnostic accuracy of molecular testing with SARS-CoV-2 antigen detection by a rapid chemiluminescent assay and two different point of care ultra-rapid immunochromatographic assays.

2. Materials and methods

2.1. Patients and samples

A total of 234 subjects were enrolled between 1st August and 30th November 2020. One hundred thirty-eight (52 females, 86 males, mean age ± SD: 55 ± 17 years) were COVID-19 inpatients, and 96 (47 females, 49 males, mean age ± SD: 42 ± 15 years) were outpatients screened for suspected SARS-CoV-2 (i.e. contact with a SARS-CoV-2 positive subject or with typical symptoms). After giving fully informed consent in writing (Local Ethic Committee Nr. 27444), patients were asked to collect a morning saliva sample (Salivette device, SARSTEDT AG & Co, Nümbrecht, Germany). After saliva sampling, trained nurses collected three NPS from each patient. A subset of 32 inpatients repeated saliva collection and testing 7 days after enrollment and, among them, 23 repeated also NPS.

Clinical data on each inpatient were also retrieved from the hospital information system (HIS).

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2.2. Laboratory testing

For molecular testing, saliva and NPS were analyzed by TaqPath COVID-19 RT-PCR kit (Applied Biosystems, USA) as detailed in Supplementary materials and methods [21].

Chemiluminescence immunoassay (CLEIA) was performed using a LUMIPULSE SARS-CoV-2 Ag kit on a LUMIPULSE G1200 automated analyzer (Fujirebio, Tokjo, Japan), following the manufacturer’s instructions. After the first result, available in 30 min, the diagnostic system provides 120 results per hour. ESPLINE rapid test and Panbio™ COVID-19 Ag Rapid Test (ABBOTT, Chicago, Illinois, USA) were the two evaluated POC lateral flow immunochromatographic assays.

2.3. Statistical analysis of data

The statistical analysis of data was made with Stata software ver. 13.1 (Lakeway drive, TX, US) by Wilcoxon rank test, Kruskal-Wallis rank test, Student’s t test, Fisher’s exact test, and multiple linear regression as detailed in Supplementary materials and methods.

3. Results

3.1. Clinical data

The clinical characteristics of the 138 inpatients are shown in Table 1.

While no significant differences were found between in- and outpatients (n = 96) for gender distribution (Fisher’s exact test: p = 0.106), the mean age of outpatients was significantly lower than that of inpatients (Student’s t test for unpaired data: t = 6.51, p < 0.001).

3.2. NPS and saliva molecular testing

At enrollment, NPS rRT-PCR results were positive among 84/138 (60.9%) inpatients and 3/96 (3.1%) outpatients, while saliva in 67/127 (52.8%) inpatients and in 4/96 (4.2%) outpatients. Positive NPS and saliva were more frequent among inpatients with a time lapse from symptoms of <7 days (37/38, 97.4% for NPS; 32/36, 88.9% for saliva), with respect to those with 7 to 14 days’ (40/74, 54.1% for NPS; 31/69, 44.9% for saliva) and those with more than 14 days’ (7/26, 26.9% for NPS, 4/22, 18.2% for saliva) (Fisher’s exact test: p < 0.001 for both NPS and saliva). Accordingly, the lowest median Ct values of NPS and saliva were found among inpatients with a time lapse from symptoms of <7 days (Supplementary Table S1).

Among the 96 outpatients, four had positive findings at NPS and/or saliva testing (Supplementary Table S2). The patient who was negative at NPS but positive at saliva (n. 4), repeated NPS two and ten days after enrollment, being positive in both cases.

On considering the overall inpatient population, agreement of 78.7% was found for rRT-PCR among paired NPS and saliva samples (n = 127, Cohen k = 0.569, SE = 0.086, p < 0.001). Agreement was 91.67% in those tested after 7 days after onset of symptoms (n = 36, Cohen k = 0.724, SE = 0.130, p < 0.001), 71.0% in those tested after 7 to 14 days (n = 69, Cohen k = 0.427, SE = 0.117, p < 0.001) and 81.8% in those tested after 14 days (n = 22, Cohen k = 0.571, SE = 0.188, p < 0.001).

Table 1

Clinical characteristics of inpatients.

| Days since onset of symptoms | ≤ 7 | 7–14 | greater than 14 | Pneumonia | Fever greater than 37.5 ºC | Dyspnea | Cough | Gastrointestinal | Other, minor | Therapy |
|-----------------------------|-----|------|-----------------|-----------|--------------------------|---------|-------|-----------------|------------|---------|
| Patients                    | n = 138 (%) | 38 (27.6) | 74 (53.6) | 26 (18.8) | 93 (67.4) | 97 (70.3) | 21 (15.2) | 46 (33.3) | 25 (18.1) | 55 (39.9) | 33 (23.9) | 13 (9.4) | 57 (41.3) | 35 (25.4) |

Steroids and oxygen 13 (9.4)
Steroids, oxygen and Remdesivir 35 (25.4)
after more than 14 days (n = 22, Cohen k = 0.488, SE = 0.206, p = 0.009).

3.3. NPS and saliva antigen testing

Fig. 1 shows the individual CLEIA antigen levels measured in NPS and saliva after subdividing patients on the basis of the time lapse between symptom onset and testing, data on analytical reproducibility being reported in Supplementary Table S3. SARS-CoV-2 antigen measured in NPS enabled distinction between positive and negative swabs classified on the basis of rRT-PCR with a high diagnostic accuracy (area under the ROC curve = 0.939, 95% CI: 0.903–0.977). Based on the threshold reported by the manufacturer (1.34 ng/L), the overall sensitivity and specificity were 81.6% (95% CI: 71.0–89.5%) and 93.8 (95% CI: 86.2–98.0%), respectively, with a positive likelihood ratio (LR) and negative LR of 13.2 (95% CI: 5.62–31.1) and 0.19 (95% CI: 0.12–0.32), respectively.

Salivary SARS-CoV-2 antigen allowed to distinguish between positive and negative samples classified on the basis of NPS rRT-PCR with a good diagnostic accuracy (area under the ROC curve = 0.805, 95% CI: 0.740–0.870) (Supplementary Table S4). For salivary antigen, the manufacturer’s suggested cut-off is 0.67 ng/L and the limit of quantification, 0.2 ng/L. With these two thresholds, sensitivity, specificity, positive and negative LR were calculated considering the patients overall and after subdividing them on the basis of duration of symptoms (Table 2).

Fig. 2 shows the individual levels of SARS-CoV-2 antigen measured in NPS (n = 23) and saliva (n = 32) and the corresponding Ct values obtained in a series of inpatients for whom two consecutive samples were available: one at enrollment and the other, after 7 days.

Multiple linear regression analyses were performed considering the Ct values (Orf1ab) of NPS and saliva as dependent variables, while antigen levels, clinical and demographic data as predictor variables (Supplementary Table S5). The Ct values were significantly correlated with antigen values independently from all clinical and demographic characteristics ($r^2 = 0.793$ for NPS and $r^2 = 0.598$ for saliva).

Fig. 3 shows the percentages of positive results for SARS-CoV-2 antigen by POC and CLEIA in NPS and saliva after subdividing samples based on viral load estimated with the Ct values at molecular analyses. POC sensitivity was satisfactory only for NPS with high viral loads (i.e. Ct values $< 25$), while CLEIA enabled the detection of viral particles with a good sensitivity also for Ct values above 25.

4. Discussion

There is a pressing need for novel strategies for the effective containment of a third wave of SARS-CoV-2 infection, particularly while awaiting vaccines. The early, reliable identification of SARS-CoV-2 infection appears to be the key to reducing community transmission yet the recommended diagnosis based on rRT-PCR analysis of NPS, although accurate, does not enable an early and prompt diagnosis.
Moreover, it calls personnel trained in NPS collection, analysis and interpretation. SARS-CoV-2 antigen determination in NPS by point-of-care immunochromatographic assays or by chemiluminescent assays developed with laboratory instrumentation has been proposed in order to overcome these limitations, and to facilitate large scale analyses [22]. This approach allows a reduction in the analytical time but does not

![Fig. 2. Kinetics of SARS-CoV-2 antigen and Ct values in NPS and saliva. Two consecutive samples (seven days apart) were available in a series of inpatients, who were subdivided on the basis of days from symptoms onset to enrollment in three groups: within 7 days (group A), between 7 and 14 days (group B), after 14 (group C). The p values reported were obtained after Wilcoxon rank test for paired data.](image)

![Fig. 3. Comparison of NPS and saliva SARS-CoV-2 rapid antigen testing. Percentages of positive results for SARS-CoV-2 antigen testing by means of rapid immunochromatographic assays (Abbott and Espline) and CLEIA in NPS and saliva after subdividing samples on the basis of viral load (Ct ranges) at molecular analyses.](image)
obviates the need to perform NPS by trained personnel within dedicated medical cabinets. This bottle-neck could be overcome by using self-collected saliva. To identify the best possible strategy for detecting the infection by antigenic rapid testing, in this second wave pandemic we studied two series of subjects representing real world scenarios: inpatients with COVID-19 disease and outpatients screened for SARS-CoV-2 due to a history of positive contact or suspect clinical signs. NPS and saliva, simultaneously collected from all patients enrolled in the study, were used as matrices for antigen detection employing immunochromatographic assays and rapid CLEIA, and for viral sequences identification by rRT-PCR. As expected, the prevalence of positive rRT-PCR findings in NPS was lower among outpatients (3%) than inpatients (61%); in this latter group, it progressively declined in parallel with duration of symptoms, as the viral load, assessed by the Ct value, declined. Based on this observation, our patients’ series was subdivided according to the duration of symptoms before enrollment: less than one week, one to two weeks, and more than two weeks. The molecular detection of viral sequences in NPS and saliva gave concordant results in a high percentage of cases at the onset (92%) and in the late phases (82%) of the infection. Conversely, patients enrolled one to two weeks after symptoms onset were more likely to have positive NPS results than positive saliva rRT-PCR results, in agreement with previous data in the literature [16]. These findings corroborate the hypothesis that buccal mucosa and salivary glands are the first sites of viral colonization, according to previous data the highest load occurring in the first week of symptom onset, followed by a progressive decline during the course of the disease [3,9,14]. As saliva is the first route for viral dissemination, it represents a suitable matrix for screening asymptomatic subjects, who are known to carry a viral load comparable to that of symptomatic patients [6,23]. CLEIA antigen testing in NPS enabled a highly accurate distinction between positive and negative swabs. The area under the ROC curve was higher than 0.9, and better than that reported earlier in a Japanese series by Hirotsu et al. [20]. Unlike the approach used in the cited study, we evaluated a large number of infected patients and for any patient one single sample, while Hirotsu et al. included serial samples from a limited number of infected patients. Moreover, CLEIA antigenic testing in NPS had a very high sensitivity not only in the first week, but also in the second week from symptoms onset paralleling molecular findings. With respect to CLEIA, rapid immunochromatographic assays in NPS appeared very reliable for high viral loads but much less so in the presence of less abundant viral loads (i.e. Ct values ranging from 25 to 30) or even worse, for higher Ct values (greater than 30), in agreement with data reported in the literature [19,24].

This must be considered a limitation of these rapid immunochromatographic assays taking also into account that Ct values ranging from 20 to 30 are considered normal findings [25]. We therefore suggest that CLEIA antigenic testing in NPS should be preferred to rapid immunochromatographic assays for obtaining accurate and fast results in the emergency setting to rapidly identify infected symptomatic patients before their hospitalization, on considering that in this context NPS collection does not represent a limitation. CLEIA antigenic testing, which requires a simple laboratory instrumentation usable by minimally trained personnel, enables the result to be obtained in 30 min and processes more than 100 samples per hour and can therefore be used by any emergency laboratory.

In the setting of screening asymptomatic subjects, salivary antigen might make the difference. In self-collected saliva, CLEIA antigen antigenic testing with molecular reflex testing for the detection of viral sequences in NPS and saliva gave concordant results in a high percentage of cases at the onset (92%) and in the late phases (82%) of the infection. Conversely, patients enrolled one to two weeks after symptoms onset were more likely to have positive NPS results than positive saliva rRT-PCR results, in agreement with previous data in the literature [16]. These findings corroborate the hypothesis that buccal mucosa and salivary glands are the first sites of viral colonization, according to previous data the highest load occurring in the first week of symptom onset, followed by a progressive decline during the course of the disease [3,9,14]. As saliva is the first route for viral dissemination, it represents a suitable matrix for screening asymptomatic subjects, who are known to carry a viral load comparable to that of symptomatic patients [6,23]. CLEIA antigen testing in NPS enabled a highly accurate distinction between positive and negative swabs. The area under the ROC curve was higher than 0.9, and better than that reported earlier in a Japanese series by Hirotsu et al. [20]. Unlike the approach used in the cited study, we evaluated a large number of infected patients and for any patient one single sample, while Hirotsu et al. included serial samples from a limited number of infected patients. Moreover, CLEIA antigenic testing in NPS had a very high sensitivity not only in the first week, but also in the second week from symptoms onset paralleling molecular findings. With respect to CLEIA, rapid immunochromatographic assays in NPS appeared very reliable for high viral loads but much less so in the presence of less abundant viral loads (i.e. Ct values ranging from 25 to 30) or even worse, for higher Ct values (greater than 30), in agreement with data reported in the literature [19,24].

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