

I BEST OF THE WEEK (24 – 30 gen 2022)

ARTICOLO	ABSTRACT	CONTENUTO E COMMENTO
<p>Møller IJB et al.</p> <p>Int J Infect Dis.</p> <p>Diagnostic performance, user acceptability, and safety of unsupervised SARS-CoV-2 rapid antigen detecting tests performed at home.</p> <p>https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8759098/pdf/main.pdf</p>	<p>Abstract</p> <p>Background: One strategy for reducing spread of COVID-19 is to contain the infection with broad screening, isolate infected individuals, and trace contacts. This strategy requires widely available, reliable SARS-CoV-2 testing. To increase testing, rapid antigen detection tests (RADTs) were developed for self-sampling, self-testing, and self-interpretation. This study examined diagnostic performance, user acceptability, and safety of nasal self-RADTs, compared to PCR testing.</p> <p>Methods: Self-RADT kits were distributed at a public COVID-19 test center in Aarhus, Denmark or delivered to participants. Participants reported test results and test preferences. During enrollment, participants reported occurrence and duration of symptoms consistent with COVID-19. Sensitivity and specificity of self-RADT, relative to oropharyngeal PCR testing, were calculated.</p> <p>Results: Among 827 participants, 102 showed positive PCR test results. Sensitivities of the self-RADTs were 65.7% (95%</p>	<p>In questo studio danese su 827 partecipanti viene valutata la capacità diagnostica del tampone antigenico rapido effettuato a domicilio rispetto al tampone molecolare, standard diagnostico di riferimento. Ne emerge una discreta sensibilità (62-65%) e un'elevatissima specificità (100%) ; inoltre, la maggior parte dei pazienti sembra preferire il test antigenico a domicilio rispetto al test molecolare effettuato da personale specializzato.</p> <p>Identificare un valido test rapido, poco costoso e di facile esecuzione da poter effettuare a livello capillare rappresenta una sfida per la « test-trace-isolate strategy » (TETRIS). Il test rapido analizzato in questo studio, nonostante sia di facilissima esecuzione, interpretazione e abbia un costo contenuto, è gravato da una percentuale di falsi negativi troppo elevata.</p>

	<p>CI: 49.2-79.2; DNA Diagnostic) and 62.1% (95% CI: 50.1-72.9; Hangzhou), and specificities were 100% (95% CI: 99.0-100; DNA Diagnostic) and 100% (95% CI: 98.9-100; Hangzhou). The sensitivities of both self-RADTs appeared higher in symptomatic participants than in asymptomatic participants. Two out of every three participants preferred self-RADT over PCR test.</p> <p>Conclusion: Self-performed RADTs were reliable, user acceptable, and safe among laypeople as supplement to professionally collected oropharyngeal PCR testing.</p>	
<p>Wolter N et al</p> <p>Early assessment of the clinical severity of the SARS-CoV-2 omicron variant in South Africa: a data linkage study</p> <p>The Lancet</p> <p>https://www.thelancet.com/action/showPdf?pii=S0140-</p>	<p>Background The SARS-CoV-2 omicron variant of concern was identified in South Africa in November, 2021, and was associated with an increase in COVID-19 cases. We aimed to assess the clinical severity of infections with the omicron variant using S gene target failure (SGTF) on the Thermo Fisher Scientific TaqPath COVID-19 PCR test as a proxy.</p> <p>Methods We did data linkages for national, South African COVID-19 case data, SARS-CoV-2 laboratory test data, SARS-CoV-2 genome data, and COVID-19 hospital admissions data. For individuals diagnosed with COVID-19 via TaqPath PCR tests, infections were designated as either SGTF or non-SGTF. The delta variant was identified by genome sequencing. Using multivariable logistic regression models, we assessed disease severity and hospitalisations by comparing individuals with SGTF versus non-SGTF infections diagnosed between Oct 1 and Nov 30, 2021, and we further</p>	<p>Studio retrospettivo nazionale di data linkages condotto in Sud-Africa con l'obiettivo di valutare la gravità clinica delle infezioni da variante omicron a confronto con infezioni da variante delta, utilizzando la perdita del gene S al test PCR per COVID-19 (SGTF) come proxy di variante omicron.</p> <p>I risultati dell'analisi suggeriscono che pazienti con SGTF hanno un ridotto rischio di ospedalizzazione e di malattia severa, probabilmente come risultato di una precedente immunità.</p>

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assessed disease severity by comparing SGTF-infected individuals diagnosed between Oct 1 and Nov 30, 2021, with delta variant-infected individuals diagnosed between April 1 and Nov 9, 2021.

Findings From Oct 1 (week 39), 2021, to Dec 6 (week 49), 2021, 161 328 cases of COVID-19 were reported in South Africa. 38 282 people were diagnosed via TaqPath PCR tests and 29 721 SGTF infections and 1412 non-SGTF infections were identified. The proportion of SGTF infections increased from two (3·2%) of 63 in week 39 to 21 978 (97·9%) of 22 455 in week 48. After controlling for factors associated with hospitalisation, individuals with SGTF infections had significantly lower odds of admission than did those with non-SGTF infections (256 [2·4%] of 10 547 vs 121 [12·8%] of 948; adjusted odds ratio [aOR] 0·2, 95% CI 0·1–0·3). After controlling for factors associated with disease severity, the odds of severe disease were similar between hospitalised individuals with SGTF versus non-SGTF infections (42 [21%] of 204 vs 45 [40%] of 113; aOR 0·7, 95% CI 0·3–1·4). Compared with individuals with earlier delta variant infections, SGTF-infected individuals had a significantly lower odds of severe disease (496 [62·5%] of 793 vs 57 [23·4%] of 244; aOR 0·3, 95% CI 0·2–0·5), after controlling for factors associated with disease severity.

Interpretation Our early analyses suggest a significantly reduced odds of hospitalisation among individuals with SGTF versus non-SGTF infections diagnosed during the same time

	<p>period. SGTF-infected individuals had a significantly reduced odds of severe disease compared with individuals infected earlier with the delta variant. Some of this reduced severity is probably a result of previous immunity.</p>	
<p>Shoji K et al</p> <p>Clinical characteristics and outcomes of COVID-19 in pregnant women: a propensity score matched analysis of the data from the COVID-19 Registry Japan</p> <p>Clinical Infectious Diseases</p> <p>https://academic.oup.com/cid/advance-article/doi/10.1093/cid/ciac028/6509063</p>	<p>Background</p> <p>Several studies have investigated whether pregnancy is a risk factor for developing severe COVID-19; however, the results remain controversial. In addition, the information regarding risk factors for developing severe COVID-19 in pregnant women is limited.</p> <p>Methods</p> <p>A retrospective cohort study analyzing the data from the nationwide COVID-19 registry in Japan was conducted. Propensity score matched analysis was performed to compare COVID-19 severity between pregnant and nonpregnant women. Multivariate analysis was also conducted to evaluate risk factors for developing moderate-to-severe COVID-19 in pregnant women.</p> <p>Results</p> <p>During the study period, 254 pregnant and 3752 nonpregnant women of reproductive age were identified. After propensity score matching, 187 pregnant women and 935 nonpregnant women were selected. A composite outcome of moderate-to-severe COVID-19 was more</p>	<p>Studio retrospettivo di coorte condotto utilizzando i dati del registro COVID-19 nazionale giapponese in cui 187 donne in gravidanza sono state matchate per gravità con 935 donne non in gravidanza per valutare i fattori di rischio di sviluppare malattia moderato-severa nelle donne in gravidanza.</p> <p>I risultati hanno dimostrato che la gravidanza in sè rappresenta un fattore di rischio per forme moderato-severe e che nelle donne in gravidanza la presenza di comorbidità e le fasi avanzate di gravidanza (secondo e terzo trimestre) rappresentano fattori di rischio per forme moderato-severe di COVID-19.</p>

frequently observed in pregnant women than that of nonpregnant women (n=18, 9.6% vs. n=46, 4.9%; $P=0.0155$). In multivariate analysis, the presence of underlying diseases and being in the second-to-third trimester of pregnancy were recognized as risk factors for moderate-to-severe COVID-19 in pregnant women (odds ratio [95% confidence interval]: 5.295 [1.21-23.069] and 3.871 [1.201-12.477], respectively).

Conclusions

Pregnancy could be a risk factor for moderate-to-severe COVID-19 for women in Japan. In addition to the presence of comorbidities, advanced pregnancy stages may contribute to greater risks for developing moderate-to-severe COVID-19 in pregnant women.