

Researchers develop protocol for a more accurate COVID-19 testing technique

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Researchers develop new protocol for COVID-19 testing technique Credit: NYUAD



One critical barrier in efforts to control the COVID-19 pandemic has been the relatively high false-negative rate of the most commonly-used Reverse Transcription Polymerase Chain Reaction (RT-PCR) testing methods.

Often, these methods are not able to detect lower viral loads, which are typically present in asymptomatic individuals but below the limit of detection (LoD) of standard one- or two-step RT-PCR methods. Researchers from NYU Abu Dhabi's Biology Program and Center for Genomics and Systems Biology (CGSB) have implemented a new threestep testing approach that promises to significantly—and costeffectively—improve testing accuracy.

Rather than combining the RT and qPCR reactions, the NYUAD researchers, led by Professor of Biology at NYU and NYUAD Kris Gunsalus and Assistant Professor of Biology at NYUAD Youssef Idaghdour employed a technique involving sequential RT, cDNA preamplification, and qPCR, using a microfluidics platform.

The researchers demonstrated reliable ultra-sensitive and quantitative detection of low SARS-CoV-2 viral loads (less than one copy/microliter) using synthetic viral RNA, clinical nasopharyngeal swab samples and saliva samples, including samples previously diagnosed as negative by clinical diagnostic testing. Their findings, reported in the journal *Processes* are that this microfluidic RT-PCR assay is a sensitive, quantitative, and cost-effective detection strategy which could markedly reduce the false negative rate of clinical diagnostic tests—a potentially valuable tool in SARS-CoV-2 active screening and early detection programs.





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"By adding a pre-amplification step and using microfluidic technology, we have demonstrated that this sensitive detection method can detect low viral loads, which is critical to enabling the most effective public health responses to the COVID-19 pandemic," said Idaghdour. "Our three-step approach can significantly reduce the false negative rate of standard RT-PCR-based diagnostic tests for SARS-CoV2 and other viral infections. This would allow public health officials to more readily identify and trace asymptomatic individuals, enhance the accuracy of air and environmental sampling for SARS-CoV-2, expand accurate detection to



saliva testing and help curtail the spread of the virus."



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More information: Xin Xie et al, Microfluidic Nano-Scale qPCR Enables Ultra-Sensitive and Quantitative Detection of SARS-CoV-2, *Processes* <u>doi.org/10.3390/pr8111425</u>

Provided by New York University



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