



Electrochemical detection of multiple RNA targets using MXene/duplex-specific nuclease: A path towards simultaneous detection of SARS-CoV-2 and H1N1 influenza virus

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Introduction

Rapid and sensitive detection of SARS-CoV-2 in an affordable fashion is vital for the early diagnostics of the COVID-19. Moreover, with the inevitable emergence of the upcoming seasonal influenza (flu), it is highly important to develop a biosensor capable of differentiating influenza virus from SARS-CoV-2 in order for the healthcare professionals to prescribe the most appropriate medications.

Concurrent detection of multiple RNAs

Sensor performance

SEM image of AuNP@MXene/Au







Selectivity and specificity analysis



- A novel synergetic signal amplification system is reported for multiple, rapid and attomolar quantification of microRNAs.
- MXene-Ti₃C₂T_x is synthesized and modified with 5 nm gold nanoparticles (AuNPs) as the electrochemical signal booster by almost 4 times of magnitude.
- DSN-based amplification system enhanced the sensitivity and provided single-mutation recognition ability.
- The biosensor could detect both target RNAs in 80 min with the detection limits of 204 aM and 138 aM and a wide linear range from 500 aM to 50 nM demonstrating promising features for the fabrication of practical devices

How to adapt the proposed biosensor for detection of long RNA genomes of SARS-CoV-2 and h1N1?

 We are currently working on the matter and appreciate any practical comment.

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