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Point-of-care diagnostics for SARS-CoV-2 wastewater-based epidemiology: a big leap toward miniaturization

Ahmed Donia,^a Nonsikelelo Precios Mthethwa-Hlongwa,^a Isaac Dennis Amoah,^{ab} Sheena Kumari^a and Faizal Bux^{*a}

The integration of point-of-care diagnostics in SARS-CoV-2 wastewater-based epidemiology signifies a substantial leap forward in disease surveillance and monitoring. Various innovative methods have been explored for the detection of SARS-CoV-2 in wastewater samples, each with its unique advantages and applications. Loop-mediated isothermal amplification (LAMP) has emerged as a prominent technique due to its sensitivity, rapidity, and simplicity. It amplifies pathogen genetic material without the need for a thermal cycler, making it suitable for point-of-care detection. Pairing microfluidic technology with LAMP enables swift and automated analysis directly on a chip. Additionally, paper-based devices offered a cost-effective and straightforward approach for LAMP-based detection, particularly beneficial in resource-limited settings. Combining LAMP with CRISPR/Cas technology enhances specificity and sensitivity, crucial for variant-specific detection like VarLOCK (variant-specific SHERLOCK). Aptamer-based electrochemical chips offer high specificity and stability, making them suitable for wastewater analysis. By integrating aptamer technology with filtration and purification systems, detecting SARS-CoV-2 on-site in wastewater becomes feasible, offering a practical solution for monitoring viral transmission. These methods showcase the diverse approaches in SARS-CoV-2 detection through wastewater-based epidemiology, promising effective disease surveillance. In this review, we will summarize these recent advancements in point-of-care diagnostics for SARS-CoV-2 wastewater-based epidemiology and explore their potential applications beyond SARS-CoV-2.

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Water impact

Integrating point-of-care diagnostics into SARS-CoV-2 wastewater surveillance is a novel method to enhance disease monitoring with rapid on-site detection, vital for early outbreak identification and timely public health responses. This innovation not only broadens its applications beyond COVID-19 surveillance but also offers potential for detecting other emerging and re-emerging pathogens in wastewater, bolstering public health efforts.

1. Introduction

A. Overview of wastewater-based epidemiology

Wastewater-based epidemiology (WBE) analyzes pollutants and biomarkers in untreated wastewater to gain insights into community behavior, substance use, and disease prevalence, offering real-time understanding of health conditions and environmental exposure. Initially introduced by Christian Daughton, WBE assesses drug residues in wastewater as indicators of population-level drug consumption.¹ Amidst the COVID-19 pandemic, WBE identifies SARS-CoV-2 RNA in wastewater, aiding global pandemic surveillance and

reinforcing adherence to World Health Organization (WHO) guidelines for wastewater management.² Wastewater monitoring of SARS-CoV-2 demonstrates superior cost-effectiveness compared to clinical testing, providing comprehensive population-level surveillance at a fraction of the cost. Wright and colleagues conducted a comparative cost analysis and found that wastewater-based epidemiology was more cost-effective, with data obtained at only 1.7% of the total cost of clinical testing (\$6042 for wastewater-based epidemiology *versus* \$338 000 for clinical testing).³

B. Importance of point-of-care diagnostics in wastewater surveillance

In areas with limited resources, conventional laboratory methods such as PCR and ELISA are often inaccessible or

^a Institute for Water and Wastewater Technology, Durban University of Technology, Durban, South Africa. E-mail: faizalb@dut.ac.za

^b Department of Environmental Science, University of Arizona, Tucson, AZ, USA

costly. Point-of-care (POC) devices offer a solution, providing on-site detection that replaces expensive methods with accessible, rapid, and cost-effective alternatives.⁴

Microfluidic devices, especially, excel at detecting contaminants in water and food. They are valued for their affordability, simplicity, minimal sample requirements, speed, and on-site usability.^{5,6} Different approaches have been explored, including PCR, loop-mediated isothermal amplification (LAMP), mass spectrometry, and electrochemistry. A microfluidic quantitative PCR (MFQPCR) system has been used to detect human viral and enteric bacterial pathogens in wastewater treatment plants, reaching concentrations as low as 2 copies per μL of cDNA/DNA⁷ and 100 cells per liter,⁸ respectively. For on-chip nucleic acid testing, chips are typically divided into three sections: sampling, sensing, and signaling. Essential components of microfluidic devices include micropumps, microvalves, micromixers, droplet generators, separators, traps, cell cultures, and gradient generators.⁹ Despite these advancements, challenges such as bubble generation and reagent evaporation in microfluidic PCR need to be addressed through improved modeling of microfluidic platforms for controlled fluid transfer.¹⁰

Electrochemistry emerges as a promising point-of-care method for various targets in wastewater, offering rapid, selective, and reproducible responses.¹¹ This integrates sensor electrodes from electrochemical biosensors with sustainable polymers and microfluidics for affordable sensing options. These devices, employing cost-effective polymers like plastics and thin-layer metal electrodes, inherently reduce production costs, which are further diminished through miniaturization. Biofuel cells, notably

powered by enzymes or microbes mediating glucose, oxygen, and other digestate conversions, have shown promise. Manufacturing plastic-based sensing platforms, utilizing printed circuit board technology, offers a feasible route. Leveraging this established industrial process, single board costs ranging from USD 0.20–0.52 per cm^2 , depending on layer complexity, contrast significantly with USD 10–20 per cm^2 for glass-based chips. Mass production can substantially lower these costs, promising the creation of genuinely cost-effective devices.¹²

The optical analytical technique encompasses methodologies like ultra-violet visible, fluorescence, chemiluminescence, or Raman spectroscopy, employed to monitor signal alterations. It includes a recognition component that engages with the target analyte and a transducer element that signals the interaction.¹³ Immunoprobes, typically colloidal gold, carbon, or colloidal selenium nanoparticles, are pivotal in colorimetric assays for detecting target analytes.^{14,15} They interact specifically with targets, inducing observable color changes, enabling qualitative or quantitative analysis. Advancements in nanotechnology have led to the widespread adoption of luminescent nanoparticles in point-of-care testing due to their optical stability and suitability for immunoassays, including quantum dots,¹⁶ dye-doped nanoparticles,¹⁷ and up-converting nanoparticles.¹⁸

C. Rationale for using wastewater surveillance for SARS-CoV-2 monitoring

SARS-CoV-2 may integrate into wastewater treatment plants due to its potential to adhere to fecal particles and subsequently infiltrate these facilities.¹⁹ Multiple studies from various countries have confirmed the presence of SARS-CoV-2 in sewage samples, with viral RNA levels showing an upward trajectory over time.^{20–23} A notable finding is the prolonged shedding of SARS-CoV-2 RNA *via* the digestive system compared to the respiratory tract, lasting up to 27.9 days from the onset of symptoms.²³ The presence of SARS-CoV-2 viral load in wastewater reflects both asymptomatic and symptomatic shedding, providing a more accurate representation of community infection levels.²⁴ Current surveillance primarily relies on diagnostic testing, which is limited in its ability to provide timely warnings for surges and is susceptible to various biases. In contrast, wastewater surveillance has emerged as a reliable, less biased, and predictive tool for monitoring SARS-CoV-2 infection rates in near-real-time.^{25–27} Wastewater surveillance has demonstrated its effectiveness by identifying lower-than-expected viral circulation in central Florida during the 2022/23 winter. Moreover, its capability to detect subtle changes in infection rates 30–46 days before clinical tests, even at low baseline levels of circulation, enables communities to confidently resume normal activities during low-risk periods while promptly implementing mitigation measures when the risk escalates.²⁸



Ahmed Donia

Dr Ahmed Donia is a postdoctoral research fellow at the Institute for Water and Wastewater Technology, Durban University of Technology, South Africa. He has a Ph.D. in Microbiology and Immunology from the Department of Biosciences, COMSATS University Islamabad, Islamabad campus, Pakistan. Dr Donia is an awardee of the TWAS (The World Academy of Sciences) full-time Ph.D. Fellowship. His Ph.D.

research focused on SARS-CoV-2/COVID-19. His Ph.D. thesis title was “Environmental Surveillance to Immunosurveillance & Rapid Approaches for Detection of SARS-CoV-2”. Dr Donia published many original and novel papers related to SARS-CoV-2/COVID-19 in internationally peer-reviewed journals. His research areas include wastewater-based epidemiology/surveillance, point-of-care testing, lab-on-a-chip technology, and antimicrobial resistance.

Critical review

Wastewater surveillance has been used for monitoring viruses other than SARS-CoV-2 such as poliovirus, norovirus, hepatitis A virus, influenza, respiratory syncytial virus (RSV) and enteroviruses. In efforts toward poliomyelitis eradication, Greece undertook enhanced laboratory surveillance of enteroviruses between 2008 and 2014. The Hellenic Polioviruses/Enteroviruses Reference Laboratory rigorously monitored high-risk populations, immigrants from poliovirus-endemic regions, and environmental samples. Out of 722 stool and 179 sewage water samples examined, no wild-type polioviruses were detected; however, two vaccine-derived strains were identified. Enteroviruses were found in

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25.3% of stool and 25.1% of sewage samples, predominantly species A, B, or C. Genetic analysis revealed close relationships between sewage and stool isolates, underscoring the importance of monitoring asymptomatic carriers to preserve Greece's polio-free status amid ongoing population movements.²⁹

Environmental surveillance in London, UK, was conducted to test sewage samples for investigation of the molecular properties of type 2 poliovirus isolates found in sewage to detect virus transmission in the community.³⁰ The findings revealed 118 genetically linked poliovirus isolates related to the serotype 2 Sabin vaccine strain in 21 of 52 sequential



**Nonsikelelo Precios
Mthethwa-Hlongwa**

Dr Nonsikelelo Precios Mthethwa-Hlongwa is a postdoctoral research fellow at Stellenbosch University, South Africa. Her research area focuses on molecular biology, microbiomics, and environmental and human health. She completed her PhD in Health sciences at Durban University of Technology SA and University of Minnesota, USA (as an exchange student). She is a recipient of a prestigious Fulbright Scholarship (2023). She has also received the prestigious NRF Next Generation Researchers Award (2022). She published eight papers in high-impact, peer-reviewed journals and presented her work at both national and international conferences.



Isaac Dennis Amoah

Prof Isaac Dennis Amoah is a global professor and leading researcher in environmental microbiology and health-related water microbiology at the University of Arizona. With a Ph. D. in Health Sciences from Durban University of Technology, he has spearheaded significant research in wastewater-based epidemiology and antimicrobial resistance. Prof Amoah has developed numerous undergraduate courses and established pivotal research collaborations between the University of Arizona and Chinese institutions. He has received several awards, including the Willie Grabow Young Investigator Award from the International Water Association, and has published extensively in esteemed journals. His work continues to influence environmental health policies and practices globally.



Sheena Kumari

Prof Sheena Kumari is an NRF C-rated researcher and full professor at DUT. With over 15 years of post-PhD experience in water and wastewater research, her expertise spans a wide range of areas. Her research interests encompass wastewater treatment, emerging pathogens and pollutants in water/wastewater, environmental health and risk assessment, and wastewater-based epidemiology. She is a recipient of the prestigious South African Young Woman in Science award in 2019 and has also earned several institutional awards recognizing her outstanding research contributions. She has successfully graduated over 30 postgraduate students, including PhD and Masters candidates over the years.



Faizal Bux

Prof Faizal Bux is the Director of the Institute for Water and Wastewater Technology and NRF – South African Chairs Initiative Chair holder. He has ~30 years of experience in the field of water and wastewater treatment technology. His research focus areas include wastewater treatment, algal biotechnology and wastewater-based epidemiology. As an NRF B-rated researcher, he has built a substantive track record gaining national and international accolades. He holds prominent positions internationally including fellowships to International Water Association (IWA) and Royal Society of Chemistry. He is also the Chair of the Water Institute of Southern Africa and a member of the Academy of Science of South Africa.

sewage samples collected in London between February 8 and July 4, 2022. Expansion of surveillance sites helped localize transmission to several boroughs in north and east London. All isolates lacked two key attenuating mutations, were recombinants with a species C enterovirus, and a subset (20 of 118) met the criterion for a vaccine-derived poliovirus, displaying six to ten nucleotide changes in the gene encoding the VP1 capsid protein. Environmental surveillance facilitated early detection of poliovirus importation and circulation in London, enabling a rapid public health response, including enhanced surveillance and an inactivated polio vaccine campaign among children aged 1–9 years. Whole-genome sequencing established linkage of isolates and confirmed transmission of a unique recombinant poliovirus lineage detected in Israel and the USA.³⁰ The detection of poliovirus in London's wastewater has sparked considerable public interest and media coverage. It underscores the ongoing risk of resurgence globally until the virus is eradicated. The simultaneous identification of genetically linked poliovirus strains in New York, USA, and Israel suggests the potential spread of the virus through international travel, reaching geographically distant regions.³¹

In response to a foodborne hepatitis A outbreak in Italy, urban sewages at wastewater treatment plants were monitored to investigate the spread of hepatitis A virus. From July 2012 to September 2013, 38 out of 157 sewage samples (24.2%) tested positive for Hepatitis A virus, including the IA variant linked to the outbreak. Notably, while clinical surveillance identified variants not found in wastewater, the majority of sequences belonged to genotype IB, indicating low/intermediate endemicity.³²

A study demonstrated the efficacy of wastewater-based surveillance as a valuable tool for monitoring viral circulation and serving as an early warning system, particularly for respiratory viruses like SARS-CoV-2, influenza, and RSV.³³ Over a 15 month period from September 2021 to November 2022, weekly sampling campaigns were conducted at two wastewater treatment plants in Barcelona, Spain, serving the entire population. The samples underwent concentration using the aluminum hydroxide adsorption-precipitation method and subsequent analysis through RNA extraction and RT-qPCR. In all cases, SARS-CoV-2 was detected, whereas the positivity rates for influenza virus and RSV were notably lower (10.65% for influenza A (IAV), 0.82% for influenza B (IBV), 37.70% for RSV-A, and 34.43% for RSV-B). SARS-CoV-2 gene copy concentrations were consistently higher, typically ranging approximately 1 to 2 logarithmic units above those of the other respiratory viruses. Distinct spikes in influenza A virus H3:N2 during February and March 2022, as well as RSV occurrences in the winter of 2021, were noted, aligning with the chronological pattern of infections documented in the clinical database of the Catalan Government. Overall, the findings underscore the utility of wastewater surveillance in providing novel insights into the abundance of respiratory viruses in the Barcelona area, demonstrating a favorable correlation with clinical data.³³

The importance of wastewater surveillance is extended to include pharmaceuticals/drugs.³⁴ Water pollution due to pharmaceuticals is an escalating concern, necessitating water quality monitoring for public health protection. Particularly, antidepressants, benzodiazepines, antiepileptics, and antipsychotics pose risks to aquatic life. A study developed a comprehensive method for detecting 105 pharmaceutical residues in small water samples, applied to screen samples from four Wastewater Treatment Plants in northern Italy. Samples were filtered, extracted, and analysed by Ultra-High-Performance Liquid Chromatography-Quadrupole Time-of-Flight High-Resolution Mass Spectrometry (UHPLC-QTOF-HRMS). Adequate sensitivity was achieved, detecting 23 of 105 targeted drugs in all samples. Retrospective analysis identified carbamazepine metabolites, including carbamazepine-10,11-epoxide, posing neurotoxic risks.³⁵

II. Point-of-care diagnostics: an overview

Point-of-care testing, which refers to diagnostic testing conducted directly at or near the patient's location, offers the advantage of providing rapid results to healthcare providers without the delay associated with transporting samples to a central laboratory.³⁶ The inception of point-of-care testing dates to 1962 with the development of a quick method to measure blood glucose levels.³⁷ However, it wasn't until the early 1990s that point-of-care testing gained significant momentum with the introduction of portable devices capable of measuring electrolytes in emergency departments.³⁸

Biosensors, acting as analytical tools, detect specific targets by converting biomolecular recognition events into measurable physicochemical signals. Developing simple, rapid, affordable yet highly sensitive biosensors for POC testing remains challenging. Comprising biological recognition and transducer components, biosensors interact with targets through various molecular interactions like antibody/antigen or enzyme/substrate binding, amplifying them into detectable signals. Transducers—electrochemical, optical, electronic, and mass-sensitive devices—convert these interactions into measurable outputs proportional to the target concentration.³⁹ Microfluidic systems, akin to miniaturized laboratories, streamline analytical processes and provide portability for POC testing. Crucial to developing miniaturized biosensors for POC testing are sensor specificity, driven by target recognition, and sensitivity, influenced by transducer and amplifier performance. These aspects enable the detection of diverse targets with high accuracy and efficiency.^{39–41}

Electrochemical devices can also be miniaturized. Miniaturized electrochemical sensors are employed for detecting trace amounts of targets such as small organic molecules, metal ions, and biomolecules. This is achieved by measuring changes in electrochemical signals (*e.g.*, current, voltage, potential, or impedance) arising from the oxidation/reduction of chemical/biological molecules using electrodes

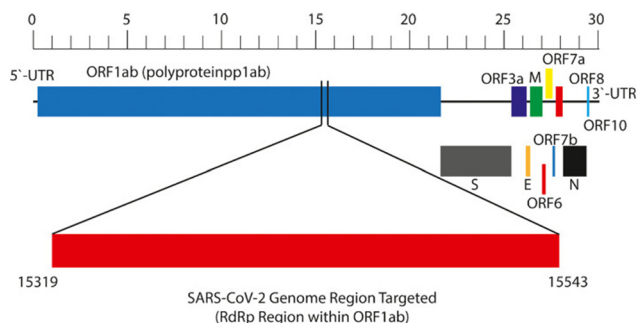


Fig. 1 Genomic location of the target region within the SARS-CoV-2 genome.⁵⁶

and electrochemical units. Typically, electrodes are modified to enhance sensor selectivity primarily through the attachment of specific recognition elements like aptamers, antibodies, and receptors. Miniaturized electrochemical sensors offer several advantages including simple instrumentation, user-friendliness, high sensitivity and selectivity, minimal sample pretreatment, short analysis time, portability, and cost-effectiveness.⁴²

Miniaturization technologies have been also applied for optical biosensor design.⁴³ Certain optical detection techniques, which are not readily adaptable to conventional sensing systems, such as fluorescence lifetime imaging, time-lapse microscopy, and multicolor analysis, have been effectively integrated as measurement methods in microfluidic sensors.^{44,45} Additionally, surface plasmon resonance, mass spectrometry, and surface acoustic wave are among the other methods successfully employed in the design of microfluidic sensors.^{43,46,47}

III. Recent advancements in point-of-care diagnostics for SARS-CoV-2 in wastewater

The target genes for SARS-CoV-2 are based on the conserved and specific open reading frame 1ab (ORF1ab), spike (S), RNA-dependent RNA polymerase (RdRp), envelope (E), and nucleocapsid (N) genes⁴⁸ (Fig. 1). The nucleic acid-based

detection method, notably reverse transcription-PCR (RT-PCR), offers high sensitivity and specificity, making it the gold standard for surveillance of SARS-CoV-2.⁴⁹ However, RT-PCR's reliance on expensive equipment, specialized reagents, and trained personnel in biosafety-compliant laboratories hinders its suitability for timely detection and surveillance. Hence, there's an urgent need for rapid and accessible SARS-CoV-2 detection and surveillance methods, leading to the emergence of real-time nucleic acid detection, particularly in the form of point-of-care testing. The point-of-care testing, performed at the sampling site using portable instruments, offers a valuable complement to RT-PCR in standard laboratories.⁵⁰ It provides quick results, enabling faster clinical decision-making and epidemic screening.^{51–54} Although real-time detection presents advantages such as accuracy, rapidity, portability, simplicity, and cost-effectiveness, it also introduces new challenges for clinical laboratories.⁵⁵

This leap toward miniaturization not only facilitates rapid results but also introduces the prospect of compact, portable diagnostic tools that can be easily deployed to wastewater treatment plants or community collection points. The miniaturization of point-of-care diagnostics for SARS-CoV-2 in wastewater-based epidemiology offers numerous advantages beyond speed and portability. By reducing the size of diagnostic devices, researchers can enhance cost-effectiveness, making these technologies more accessible for widespread use. This leap toward miniaturization also aligns with the growing trend of decentralized healthcare solutions, empowering communities to take a more active role in monitoring and managing public health.⁵⁷ Overall, the applying of point-of-care diagnostics in wastewater-based epidemiology marks a transformative step forward, not only in the battle against the current pandemic but also in the development of agile, community-centric approaches to infectious disease control.⁵⁸

Table 1 summarized the current studies about point-of-care diagnostics for SARS-CoV-2 wastewater-based surveillance.

A. LAMP

LAMP stands out as a contemporary isothermal nucleic acid amplification technique, gaining traction in point-of-care

Table 1 Point-of-care diagnostics studies for SARS-CoV-2 wastewater-based surveillance

Point-of-care diagnostics principle	Target gene/sequence/protein	Ref.
LAMP/microfluidic technology	N/ORF1a	59
LAMP	N	60
LAMP	N and E	61
LAMP	N and E	62
LAMP	RdRp	56
LAMP/electrochemical sensor	N and ORF1ab	63
Isothermal amplification/microfluidic technology/colorimetric sensor	N gene, E gene and Orf1a	64
Field-effect transistor	N and ORF1ab	65
Paper-based device (CRISPR/Cas12a and LAMP)	N, E, and S	66
VarLOCK (variant-specific SHERLOCK) ^a	S	67
Aptamer-based electrochemical chip	Spike protein	68

^a VarLOCK assays (variant-specific SHERLOCK): SHERLOCK: (Specific High-sensitivity Enzymatic Reporter unLOCKing).

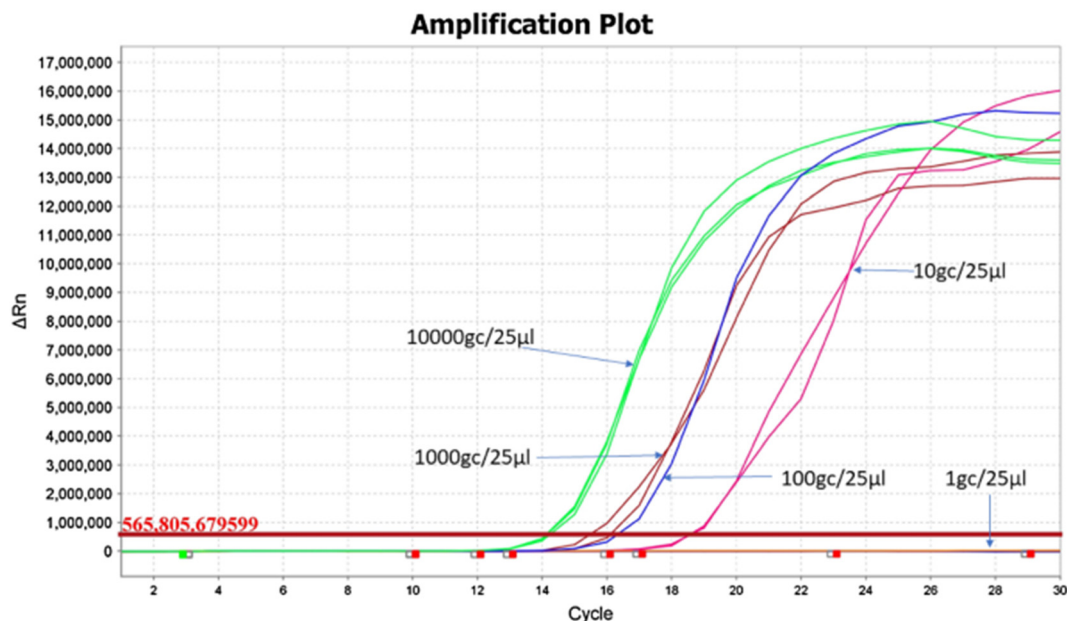


Fig. 2 Amplification plot for limit of detection assessment using the fluorescence method.⁶⁰

detection, especially during COVID-19 pandemic.^{59,60} It has several notable advantages, including sensitivity, rapidness, the absence of a need for a thermal cycler, and tolerance against sample inhibitors. Consequently, it emerges as a promising alternative to quantitative polymerase chain reaction (qPCR). It can amplify pathogen genetic material in under an hour, requiring a specific set of four to six primers to ensure precision and specificity.⁶⁹ To confirm the success of LAMP, various approaches can be employed, such as monitoring changes in fluorescence using intercalating dyes, employing DNA probes combined with gold nanoparticles, observing alterations in turbidity due to the formation of magnesium pyrophosphate precipitates, utilizing pH indicators, or conducting gel electrophoresis followed by UV detection.⁵⁹ We presented a broad insight into published point-of-care diagnostics used for SARS-CoV-2 wastewater-based epidemiology and the possibility to be used against other environmental pathogens.

Amoah and colleagues explored two optimized RT-LAMP protocols targeting N gene utilizing color change and fluorescence detection.⁶⁰ These protocols demonstrated a limit of detection of 10 copies/25 μ l reaction with positive amplification within 35 minutes. Fig. 2 showed that the limit of detection assessed using the fluorescence method is 10 copies/25 μ l reaction. Over a 4 week monitoring period of wastewater from four treatment plants, the colorimetric protocol identified a prevalence of 12.5%, increasing to 44% when the RNA template concentration was raised fivefold. The limit of RT-LAMP detection was high so this was one of the major limitations compared to droplet digit PCR and qPCR. It means when there is low concentration of virus in the samples, the LAMP was not effective as opposed to droplet digit PCR. The prevalence of the fluorescent RT-

LAMP was 31% and 47% when utilizing initial templates with concentrations of $92.7 (\pm 28.2) \text{ ng } \mu\text{l}^{-1}$ and $480 (\pm 134.5) \text{ ng } \mu\text{l}^{-1}$ of the extracted RNA, respectively. They recommended its utilization as a rapid molecular analysis tool to for COVID-19 detection within the catchment areas of wastewater treatment plants. This adaptable approach could be instrumental in future outbreaks, offering a swifter, more cost-effective, and less complex means of monitoring infections through wastewater-based epidemiology.⁶⁰

A recent study explored the use of tampons as passive swabs for sample collection and RT-LAMP to targeting the N and E genes in wastewater.⁶¹ The results of this workflow were available within three hours of sample collection. While this assay is approximately 20 times less analytically sensitive than RT-droplet digital PCR, during a building-level wastewater surveillance campaign conducted alongside independent weekly clinical testing of all students, the method demonstrated a three-day positive predictive value

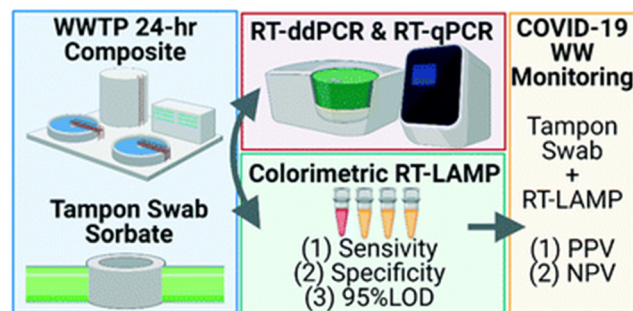


Fig. 3 Schematic representation of building-level wastewater surveillance using tampon swabs and RT-LAMP for rapid SARS-CoV-2 RNA detection.⁶¹

(PPV) of 75% (excluding convalescent cases) and a same-day negative predictive value (NPV) of 80% for incident COVID-19 cases (Fig. 3). These predictive values are comparable to those reported by wastewater monitoring using RT-qPCR. Despite lower analytical sensitivity, these findings suggest that the tampon swab and RT-LAMP workflow provides a cost-effective and rapid approach that could be effectively employed for scalable building-level wastewater surveillance for COVID-19, even in low-resource settings.⁶¹

A recent study compared the efficacy of RT-LAMP with RT-qPCR assays targeting the N and E genes of the SARS-CoV-2 genome in detecting RNA in untreated wastewater samples (Fig. 4).⁶² The RT-qPCR assays demonstrated consistent amplification down to 2×10^2 GC per reaction, with greater sensitivity at 2×10^1 GC per reaction for the US CDC N1 and N2 assays. In contrast, RT-LAMP exhibited lower sensitivity, detecting SARS-CoV-2 only at or above 2×10^3 GC per reaction. While RT-qPCR identified SARS-CoV-2 RNA in 29 out of 30 wastewater samples, RT-LAMP identified 27 positive samples, showing strong concordance with the US CDC N1 and E_Sarbeco RT-qPCR assays but lower agreement with the US CDC N2 assay. Despite lower sensitivity, RT-LAMP may offer advantages for wastewater surveillance, such as rapid results and simplicity. Logistic regression analysis indicated that RT-LAMP results correlated with RNA quantified by the US CDC N1 and E_Sarbeco assays, supporting the use of RT-LAMP as a specific and efficient method for screening wastewater samples for SARS-CoV-2, particularly in resource-limited settings.⁶²

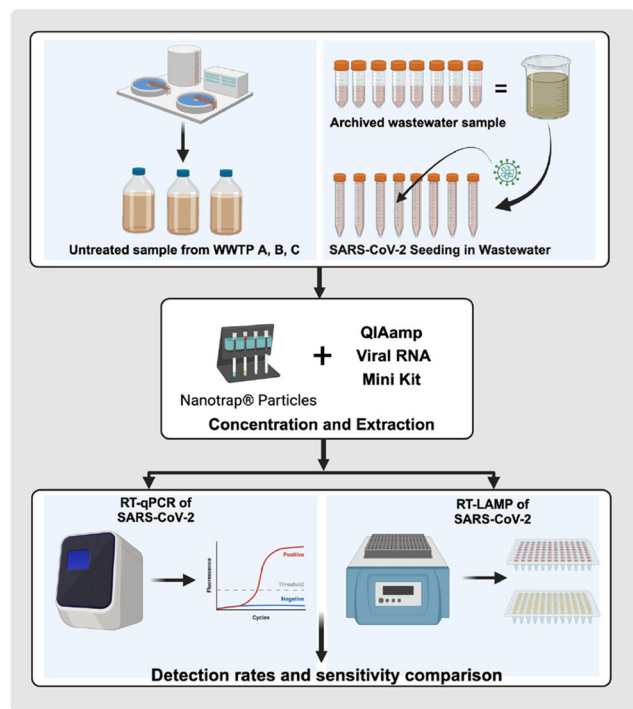
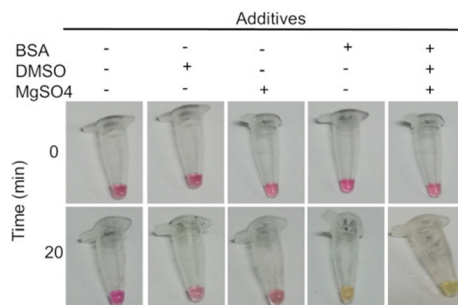


Fig. 4 Graphical abstract of comparison of RT-LAMP and RT-qPCR assays for detecting SARS-CoV-2 in wastewater.⁶²

A



B

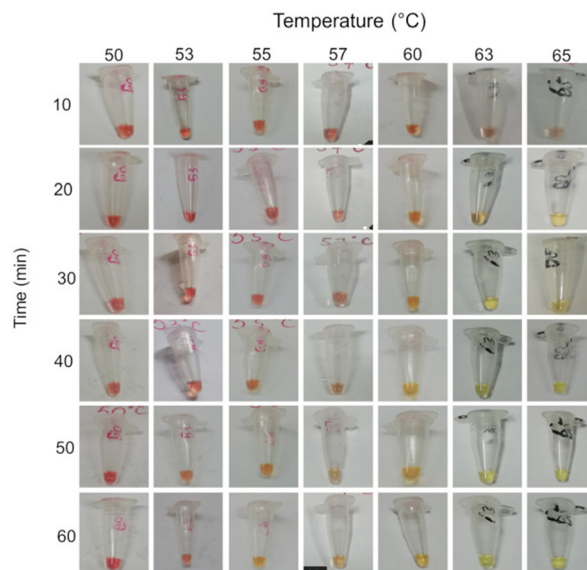


Fig. 5 Optimization of the LAMP assay.⁵⁶ The LAMP assay utilized a 225 nt synthetic RdRp gene fragment. (A) Results of the RdRp gene-based LAMP assay are shown with and without amplification-enhancing additives (BSA, MgSO₄, and/or DMSO). (B) Images display the RdRp gene-based LAMP assay results across different temperatures and time points. Pink indicates negative results, while yellow indicates positive results.

A study introduced a colorimetric RT-LAMP assay for the visual detection of SARS-CoV-2 in clinical and sewage samples.⁵⁶ The assay, based on the RNA dependent RNA polymerase (RdRp) gene, demonstrated a high success rate in detecting virus RNA in clinical samples and exhibited specificity in a short timeframe of 20 minutes, and its sensitivity was further enhanced by adding bovine serum albumin (BSA) (Fig. 5). This optimized assay was successfully applied to detect SARS-CoV-2 in sewage waters from virus hotspots in Lahore, Pakistan. To evaluate the specificity of the LAMP assay for SARS-CoV-2, they aligned each of the eight regions targeted by the primers with the reference genome sequences of six related coronaviruses. They conducted an in-silico analysis to assess the specificity of primers. The analysis revealed that most of the primer sequences exhibited high specificity for SARS-CoV-2, displaying no significant similarity to any of the six related coronaviruses included in the study. Although the F1c and LB regions displayed some similarity to

the Tor2 and HUK1 viruses, these similarities were determined to be insufficient to cause nonspecific amplifications. Therefore, owing to the inherent specificity of the LAMP assay—utilizing six primers targeting eight distinct regions in the target—and the meticulous design of the primers, they demonstrated that LAMP assay, used in this study, has the potential to differentiate SARS-CoV-2 from various related coronaviruses. Additionally, it can detect most common SARS-CoV-2 variants with equal efficiency.⁵⁶

LAMP/microfluidic technology. Donia and colleagues integrated microfluidic technology with RT-LAMP to detect SARS-CoV-2 from wastewater samples collected from COVID-19 hotspots.⁵⁹ They run RT-LAMP experiments within microfluidic devices, both with and without RNA extraction (Fig. 6), to target two specific genes (N and ORF1a) based on color change (pH change). Microchips were simply dipped into the RT-LAMP reaction solution loaded through capillary action, requiring no pump. Subsequently, the microchips were subjected to a temperature of 65 °C for a duration of 30 minutes on a pre-heated dry bath. They found that the N gene could be the most reliable target for detecting SARS-CoV-2 in wastewater. When they performed RT-LAMP on N and ORF1a genes directly in wastewater samples without RNA extraction, all samples remained pink, indicating a negative result. However, a change to a yellow color was observed in microchannels subjected to RT-LAMP targeting the N gene after RNA extraction. They found that results of RT-LAMP for these wastewater samples were largely consistent with those of RT-qPCR, with 6 (sample IDs: 1, 2, 4, 5, 6, and 7) out of 7 (85.7%) samples were positive for both RT-LAMP and RT-qPCR. Only one sample (sample ID 3) tested positive by RT-qPCR was negative by RT-LAMP (Fig. 7). This study demonstrates the successful detection of SARS-CoV-2 in wastewater samples using RT-LAMP within microfluidic chips. This opens the possibility of further modification of microfluidic devices to be preloaded with specific quantities of LAMP master mix and primers to detect other environmental pathogens. Possible modification of microfluidic devices is to fix heater with smartphone camera

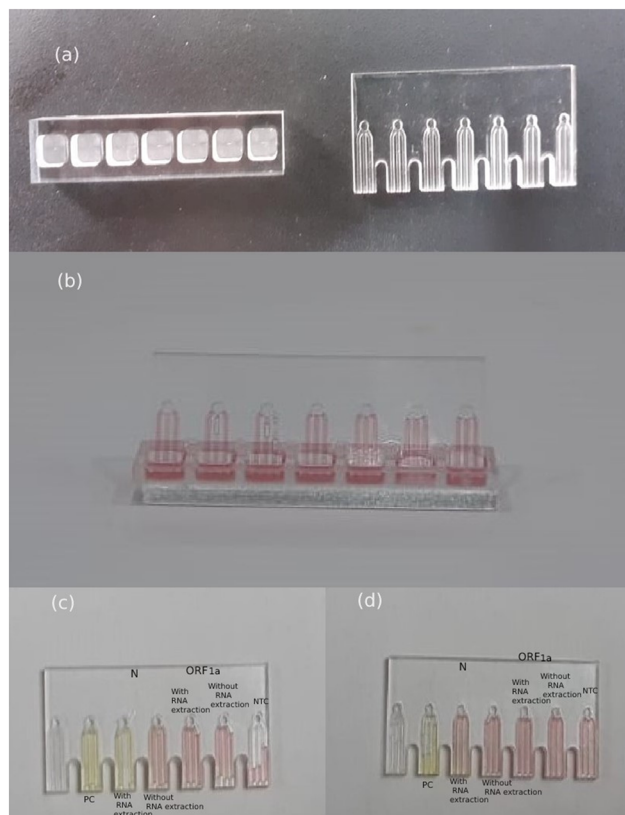


Fig. 7 Integration of microfluidic devices with RT-LAMP for the detection of SARS-CoV-2 in wastewater.⁵⁹ (a) Microfluidic device and wells. (b) Successful loading of RT-LAMP mixtures into the microfluidic device. (c) Sample ID 1 displayed a color change to yellow in the microchannels of the N gene after RNA extraction. (d) Sample ID 3 did not exhibit a color change, remaining red in the microchannels of the N gene after RNA extraction. NTC: negative control. PC: positive control.

to allow for quantitative detection through monitoring color/fluorescence/pH change.⁵⁹ One major benefit of RT-LAMP is its capacity to directly detect SARS-CoV-2 from clinical samples, bypassing the necessity for RNA extraction^{70,71} but RNA extraction was found to be necessary for detection of SARS-CoV-2 using RT-LAMP from wastewater samples.⁵⁹

LAMP/electrochemical sensor. Ramírez-Chavarría and colleagues presented a novel electrochemical sensor for specific detection of SARS-CoV-2 N and ORF1ab genes.⁶³ Utilizing screen-printed electrodes (SPEs), the sensor employs RT-LAMP and methylene blue as a redox probe for electrochemical detection. Employing end-point measurements, the sensor achieves a detection limit of $38 \times 10^{-6} \text{ ng } \mu\text{L}^{-1}$, while also tracking RT-LAMP reaction time courses for concentrations as low as $2.5 \times 10^{-6} \text{ ng } \mu\text{L}^{-1}$. The method involves mixing RNA with RT-LAMP primers and methylene blue for electrochemical transduction. A micro-volume of the sample is drop-cast onto custom screen-printed electrodes, where the RT-LAMP reaction occurs at 63 °C (Fig. 8). The resulting diffusion-controlled current is monitored by a portable potentiostat, reflecting the RT-LAMP reaction. Amplification and monitoring take around

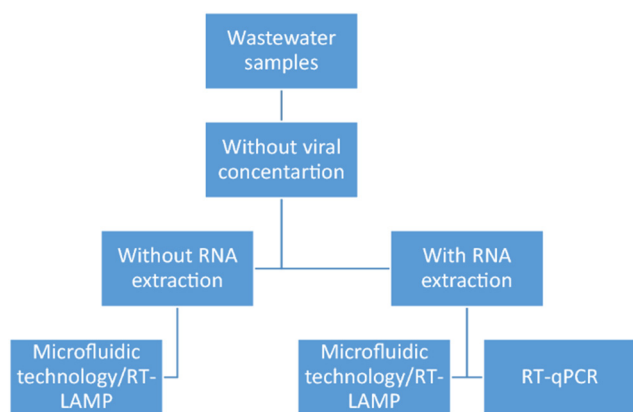


Fig. 6 Graphical abstract of integration of RT-LAMP and microfluidic technology for detection of SARS-CoV-2 in wastewater.⁵⁹

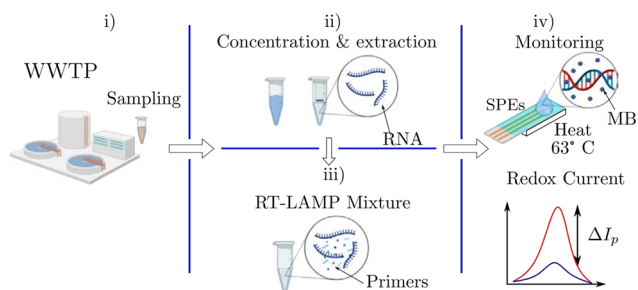


Fig. 8 Workflow of an RT-LAMP-based electrochemical sensor for wastewater samples.⁶³ i) sampling from the wastewater treatment plant. ii) Nucleic acid extraction and concentration. iii) RT-LAMP mixtures for genetic amplification. iv) Electrochemical monitoring of RT-LAMP products through redox current.

30 minutes, with $\% \Delta I_p$ peak current change encoding nucleic acid concentration. The complete experiment takes 2 hours. Validation includes simultaneous colorimetric assay on evaluated wastewater samples to confirm electrochemical monitoring performance.⁶³

Isothermal amplification/microfluidic technology/colorimetric sensor. A sensitive and multiplexed colorimetric

detection (SMCD) method was developed for early detection of SARS-CoV-2 and human enteric pathogens in wastewater.⁶⁴ The SMCD method combines on-chip nucleic acid extraction, two-stage isothermal amplification, and colorimetric detection on a 3D printed microfluidic chip. Recombinase polymerase amplification (RPA) (as a form of isothermal amplification) and synergetic enhanced colorimetric loop-mediated isothermal amplification (SEC-LAMP) were integrated within the microfluidic chip (Fig. 9). The colorimetric signal generated during SEC-LAMP can be easily observed by the naked eye or monitored in real-time using smartphone-based detection platform, eliminating the requirement for complex equipment. By integrating two-stage isothermal amplification, the method achieves sensitivities of 100 genome equivalent (GE) per mL for SARS-CoV-2 and 500 CFU mL⁻¹ for human enteric pathogens within an hour.⁶⁴

B. Field-effect transistor

Field-effect transistor (FET) biosensors present a promising avenue due to their low-cost fabrication, miniaturization, reliability, sensitivity, and label-free detection capabilities.⁷²

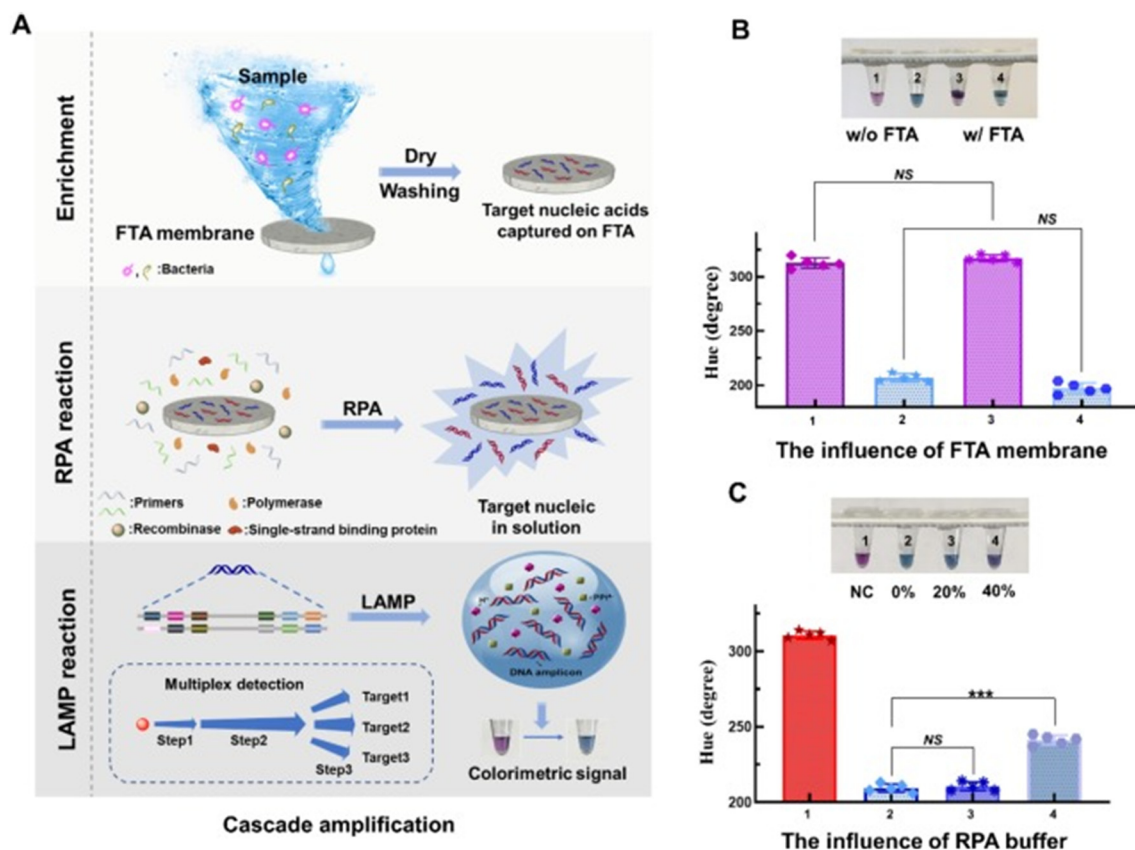


Fig. 9 Design and optimization of a Flinders Technology Associates (FTA)-based multiplexed colorimetric assay for sensitive pathogen detection in wastewater.⁶⁴ (A) Design of the FTA-based multiplexed colorimetric assay for detecting various pathogens. (B) Impact of the FTA membrane on the SEC-LAMP assay. Tubes 1–4: negative control without the FTA membrane, positive control (500 CFU per reaction, *Salmonella*) without the FTA membrane, negative control with the FTA membrane, and positive control (500 CFU per reaction, *Salmonella*) with the FTA membrane, respectively. (C) Influence of RPA reaction solution volume on the SEC-LAMP assay. Tubes 1–4: negative control, positive control (500 CFU per reaction, *Salmonella*), positive control with 20% (v/v) RPA reaction solution, and positive control with 40% (v/v) RPA reaction solution, respectively.

Extended-gate FET (EGFET) devices, in particular, offer a straightforward and cost-effective option for biosensing.⁷³ Despite technological progress, EGFET sensors face limitations in portable applications due to challenges in electrode integration and the need for labeled receptors for specificity.⁷⁴ To address these challenges, Alvarez-Serna and colleagues used an EGFET device for nucleic acid detection without electrode modification.⁶⁵ This device detected two segments of the SARS-CoV-2 genome (N and ORF1ab genes) through quantifying pH changes resulting from the isothermal amplification process, leveraging the selectivity of EGFET-based sensors to measure pH (Fig. 10). This sensor measured real wastewater samples for detection of SARS-CoV-2, with a limit of detection of $0.31 \times 10^{-3} \text{ ng } \mu\text{L}^{-1}$ for end-point measurement. Furthermore, the sensor can provide real-time-like results, with a robust response within just 15 minutes, even at concentrations as low as $0.37 \text{ ng } \mu\text{L}^{-1}$.⁶⁵ Firstly, they established the sensor's capability to detect pH fluctuations within the range where both negative and positive samples fall. Then, using the pH value of the negative template control (NTC) sample as a reference, they observed pH shifts in positive samples due to isothermal amplification. Sensor characterization was conducted using SARS-CoV-2 positive samples across seven concentrations *via* serial dilution. The calibration curve depicted a linear relationship between drain-to-source current (I_{DS}) and sample concentration. There is a straightforward relationship among the current I_{DS} and changes in pH; which in turn, are promoted by the RT-LAMP reaction.⁶⁵

C. Paper-based device

There has been growing interest in enhancing the specificity of analytical methods using CRISPR/Cas-based gene editing techniques. CRISPR/Cas technology offers

high-fidelity recognition down to single-base resolution and efficient signal transduction through *trans*-cleavage, making it a promising detection method.^{75,76} However, despite its advantages, the sensitivity of CRISPR/Cas alone may not be sufficient.⁷⁷ To address this, some studies have explored combining CRISPR/Cas with isothermal amplification methods.^{75,78} This combination utilizes the specificity of CRISPR/Cas to mitigate false positives in isothermal amplification while leveraging the high sensitivity of isothermal amplification to enhance the limit of detection (LOD) of CRISPR/Cas. One such combination is Cas12a-RT-LAMP, which has been employed for SARS-CoV-2 detection.^{79,80} For instance, Broughton and colleagues developed a lateral flow assay integrating Cas12a and RT-LAMP, achieving detection as low as 10 copies per μL in clinical samples.⁷⁹ Other approaches include visual detection methods utilizing the fluorescence characteristics of FAM, where Wang *et al.*⁸¹ and Pang *et al.*⁷⁸ integrated Cas12a *trans*-cleavage reaction and isothermal amplification into a single tube by creating two temperature regions in one device. Despite these advancements, the combination of CRISPR/Cas and isothermal amplification methods still requires further exploration and optimization.

Cao and colleagues introduced a new paper-based device that combines RT-LAMP with CRISPR/Cas12a for analyzing wastewater (Fig. 11).⁶⁶ Wastewater samples undergo filtration and lysis on a microporous membrane, after which they are transferred to a paper-based device for RNA purification and distribution. The device utilizes high-fidelity RT-LAMP to detect specific gene fragments (N, E,

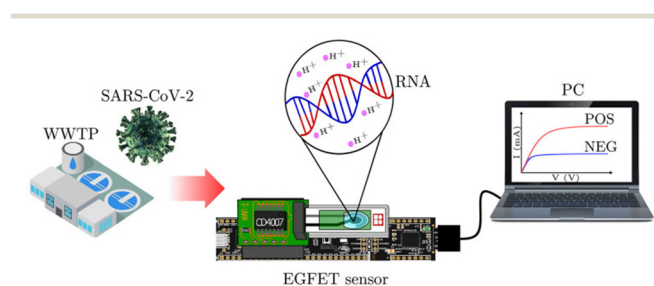


Fig. 10 Portable sensor monitors RT-LAMP for SARS-CoV-2 in wastewater.⁶⁵ Wastewater samples are collected from wastewater treatment plants. Next, the genetic material is extracted and concentrated. Then, the RNA is combined with RT-LAMP primers, and a 50 μL drop is applied onto the electrodes of the EGFET. The RT-LAMP reaction takes place at the electrode surface, with temperature control at 63 °C for approximately 30 minutes. Finally, the RT-LAMP products are monitored by observing changes in drain-to-source current (I_{DS}), which occur due to pH alterations as the reaction progresses. The pH value of the negative template control sample serves as the baseline value. Consequently, in the positive samples, one can observe pH changes attributed to the isothermal amplification process.

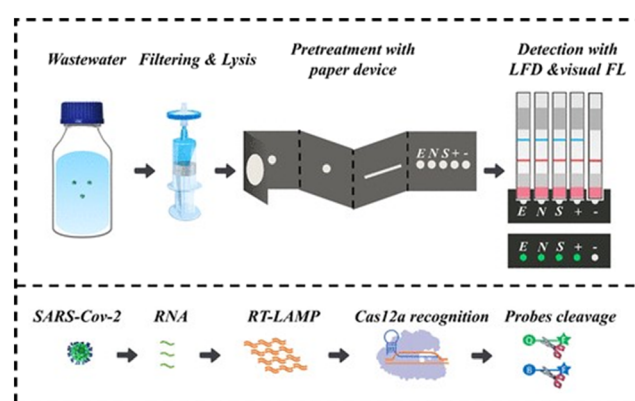


Fig. 11 Paper device for detection of SARS-CoV-2 in wastewater using CRISPR/Cas12a and RT-LAMP.⁶⁶ Wastewater samples are filtered and lysed on a microporous membrane, then transferred to a paper device for RNA purification and distribution. High-fidelity RT-LAMP detects SARS-CoV-2, which is visible by the naked eye following excitation by 480 nm light or through the use of lateral flow dipsticks. In the presence of SARS-CoV-2, RT-LAMP generates amplicons activating Cas12a-gRNA, leading to unspecific ssDNA probe cleavage. Fluorescence arises from FAM-BHQ1 separation in fluorescence method, while lateral flow method exhibits strong test-stained band due to FAM-biotin separation. Without SARS-CoV-2, probe remains intact, yielding no fluorescence.

and S) of SARS-CoV-2. Results are observable by the naked eye following excitation by 480 nm light or lateral flow dipsticks. When SARS-CoV-2 is present, RT-LAMP produces amplicons that activate Cas12a-gRNA, resulting in non-specific cleavage of single-stranded DNA probes. Fluorescence is observed through the separation of FAM-BHQ1 in fluorescence-based methods, while the lateral flow method shows a distinct test-stained band due to the separation of FAM-biotin. In the absence of SARS-CoV-2, the probe remains intact, resulting in no fluorescence. The device's design and components are illustrated in Fig. 11. Testing involved using SARS-CoV-2 spiked wastewater to determine limits of detection (LODs) and quantitative capabilities, followed by blind testing to confirm reliability. This paper-based device allowed for the simultaneous detection of the N, E, and S genes, having limit of detection 25, 310, and 10 copies per mL, respectively. Furthermore, the device enabled semiquantitative analysis within the range of 0 to 310 copies per mL, owing to the varying LODs of the three genes. Blind experiments confirmed the device's suitability for wastewater analysis, showing 97.7% sensitivity and 82% accuracy in semiquantitative assessment.⁶⁶

The device underwent a six-month application period from June to November 2021 for analyzing wastewater, aimed at further validating its performance. The Erqiao wastewater treatment plant in Huaguoyuan, among the largest communities in Asia, and the Jinyang wastewater treatment plant, situated in a new urban area in Guiyang, were selected as the designated locations for this evaluation. Following the analysis, no viruses were detected in the sewage over the course of six months, indicating that the

communities were experiencing a low-risk status, consistent with the observations of the Centers for Disease Control and Prevention (CDC). Consequently, the innovative device exhibits promising potential for sewage detection.⁶⁶

D. VarLOCK (variant-specific SHERLOCK)

CRISPR-based diagnostic tests offer a new platform for detecting pathogens. Techniques like SHERLOCK (Specific High-sensitivity Enzymatic Reporter unLOCKing) uses a two-step process involving target amplification and CRISPR-based nucleic acid detection,^{75,77} successfully applied to detect SARS-CoV-2 in clinical samples.⁷⁹ However, these methods are more complex than point-of-care tests due to RNA extraction and multiple liquid-handling steps, increasing the risk of sample cross-contamination. DETECTR, a recently emerging molecular diagnostic method, utilizes CRISPR-Cas12a along with isothermal nucleic acid amplification.⁷⁹

In 2017, Jennifer Doudna's team unveiled the CRISPR-Cas diagnostic tool named DNA endonuclease-targeted CRISPR trans reporter (DETECTR).⁷⁵ This technique leverages the collateral activity of the Cas12a protein, which is activated upon recognition of target RNA. Specifically, the LbCas12a variant of Cas12a demonstrates nonspecific collateral activity, breaking down neighboring DNA molecules after identifying target RNA. When utilized alongside single-stranded DNA reporters (probes) within a biological sample, Cas12a's detection of pathogenic nucleic acids triggers collateral activity, resulting in the destruction of DNA probes. These probes, resembling conventional TaqMan probes, emit a fluorescent signal upon degradation,

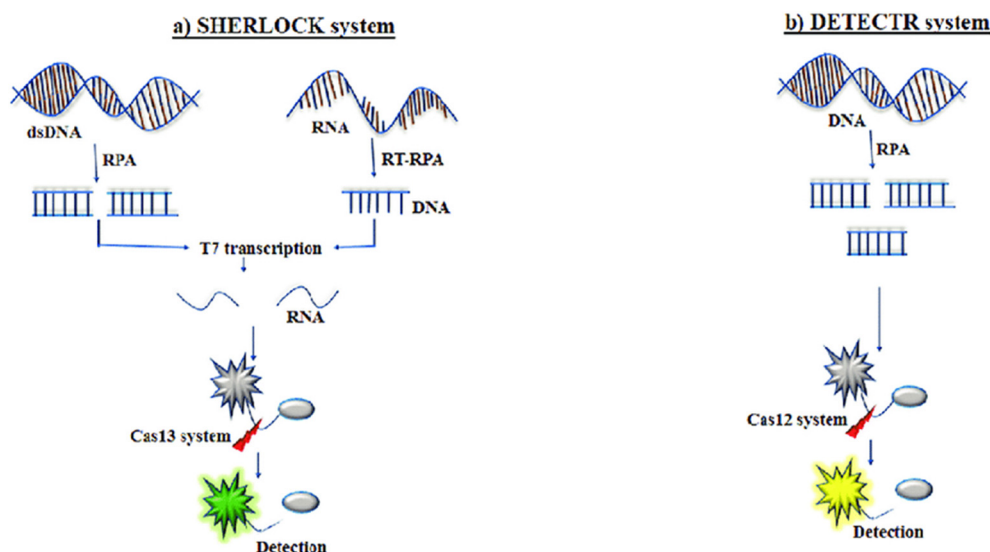


Fig. 12 Mechanism of SHERLOCK and DETECTR system. (a) SHERLOCK system. (b) DETECTR system.⁸³ (a) Recombinase polymerase amplification (RPA) or reverse transcription (RT)-RPA amplifies targeted double-stranded DNA (dsDNA) or RNA, respectively. This is followed by coupling RPA with T7 transcription to convert targeted RNA, allowing detection by the Cas13 system. The amplification steps, along with the reporter probe, facilitate specific high-sensitivity enzymatic reporter unlocking (SHERLOCK) to identify the targeted sequence. (B) DNA is amplified with RPA in the DNA endonuclease-targeted CRISPR trans reporter (DETECTR). The Cas12 system associates with the single-stranded DNA (ssDNA) of interest, initiating the DNase activity of the Cas12 system. This amplification step, in conjunction with the reporter probe, enables DETECTR to detect the targeted sequence.

which can be detected using a fluorimeter. To further enhance DETECTR's capabilities, an isothermal preamplification step (RPA) has been incorporated, improving analytical sensitivity, and obviating the necessity for costly equipment. SHERLOCK operates on similar principles to DETECTR but relies on the activity of the

Cas13 nuclease from *Leptotrichia wadei* (Fig. 12). Unlike Cas12a, Cas13 exclusively identifies and cleaves RNA rather than DNA.⁸²

Joung *et al.*⁸⁴ introduced STOP (SHERLOCK Testing in One Pot), a simple method for detecting SARS-CoV-2 with sensitivity comparable to RT-qPCR assays. STOP combines

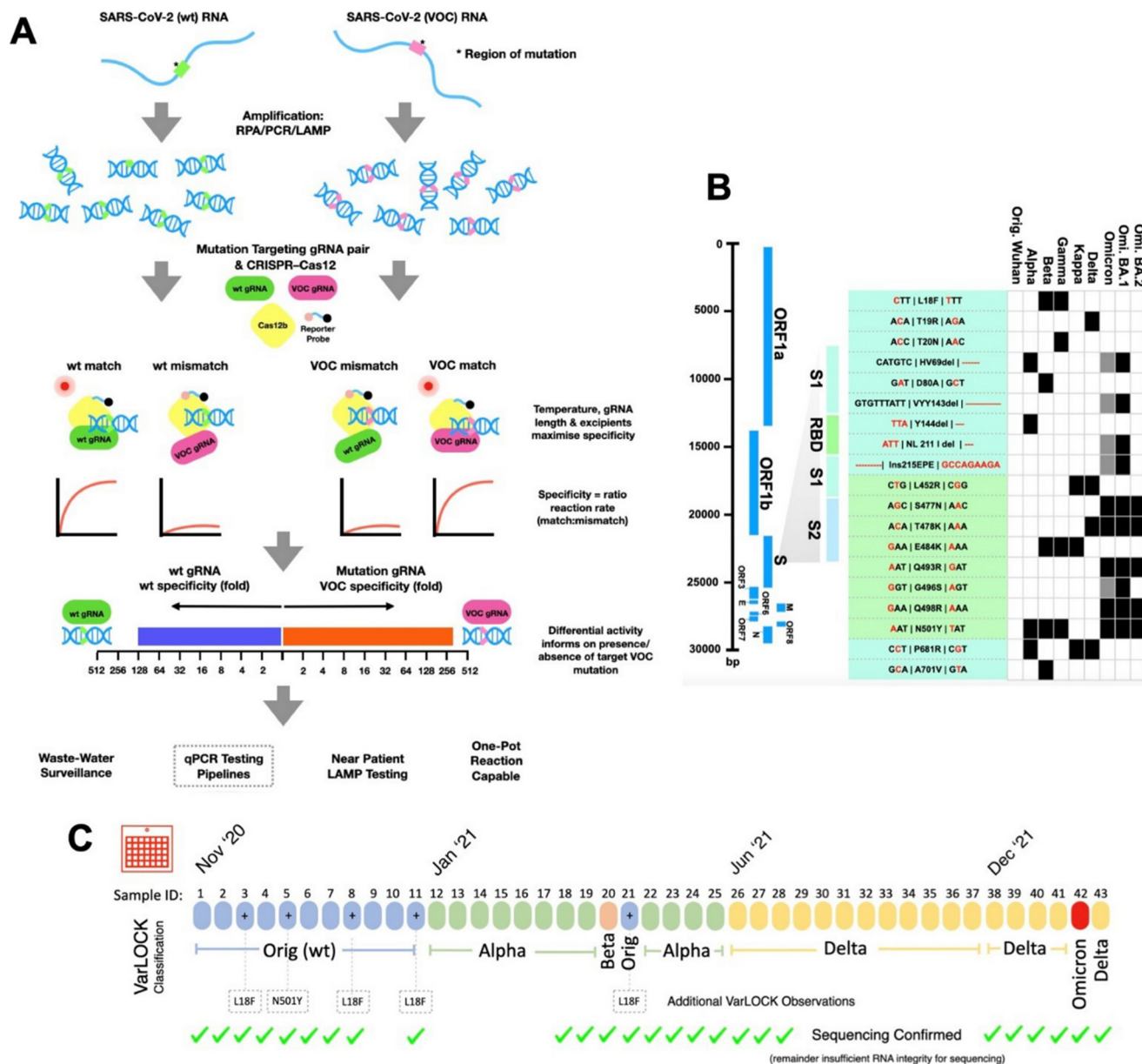


Fig. 13 SARS-CoV-2 variant of concern identification using the VarLOCK assay.⁶⁷ (A) The VarLOCK assay is illustrated as follows: SARS-CoV-2 RNA regions, whether containing the wildtype (green) or mutant (pink) sequences, are amplified using RPA, LAMP, or PCR. The resulting amplified DNA fragments are then subjected to the SHERLOCK assay with a pair of gRNAs specific to either the wildtype sequence (wt gRNA, green) or the mutant sequence (VOC gRNA, pink). The goal is to establish conditions where only a perfect match between the gRNA and the amplified DNA will activate the Cas12b protein's collateral nuclease activity, leading to the cleavage of a quenched fluorescent reporter. This cleavage results in an increase in fluorescence, allowing the nuclease activity to be monitored. The ratiometric response of an unknown SARS-CoV-2 variant sample indicates the presence or absence of the targeted mutation, thus enabling the identification of the VOC. (B) The schematic on the left shows the genome of SARS-CoV-2, with an expanded view of the spike protein encoding region, highlighting the locations of the RBD, S1, and S2. Nineteen mutation sites were selected for the VarLOCK assay, for which gRNA pairs were designed and optimized. The table displays the nucleotide sequences encoding the relevant amino acids, with nucleotide substitutions/deletions marked in red. The association of these mutations with various variants of concern is also indicated, creating a barcode-like identification matrix. (C) This approach, when applied to saliva samples, successfully identifies the original Wuhan, Alpha, Beta, Delta, and Omicron variants, with results confirmed by sequencing.

simplified RNA extraction, isothermal amplification, and CRISPR-based detection from nasopharyngeal or anterior nasal swab samples. Operating at a single temperature, STOP provides results in under an hour with minimal equipment.

Nan *et al.*⁶⁷ expanded on the CRISPR-Cas assay SHERLOCK to develop VarLOCK (variant-specific SHERLOCK), an innovative molecular diagnostic approach suitable for known variants of concern (VOCs) and newly detected variants (Fig. 13). VarLOCK swiftly and sensitively detects SARS-CoV-2 VOCs, applicable at point of care or in testing facilities of any size. It also estimates VOC proportions in pooled wastewater samples, complementing sequencing efforts for rapid identification of infected individuals and breaking infection chains. They optimized VarLOCK assays for various spike gene mutations of SARS-CoV-2 and validated them with samples from the Cardiff University Testing Service. VarLOCK's utility extends to national wastewater surveillance of SARS-CoV-2 variants, demonstrating adaptability to new and emerging VOCs.

They validated and implemented VarLOCK using patient samples from the Cardiff University COVID19 Testing Service. These assays can also be employed in national wastewater surveillance, facilitating identification and mapping of variant prevalence across different times and locations. Coupled with LAMP amplification, VarLOCK assays offer potential for widespread use in near-patient variant-specific diagnostic testing. They showcased VarLOCK's rapid adaptability to new variants, exemplified by the development of an Omicron-specific assay within two weeks of its classification by the WHO. This advancement could significantly enhance public health initiatives by enabling swifter identification of Omicron infections and providing a scalable approach for future variants of concern.⁶⁷

E. Aptamer-based electrochemical chip

Aptamers, highly specific nucleic acid-based recognition elements, are selected in labs to target molecules, including viral proteins.⁸⁵ They possess exceptional specificity, stability, and versatility for surface attachment and signal transmission, making them ideal for biosensors. During the COVID-19 pandemic, aptamers have been engineered to target various parts of the SARS-CoV-2 virus, such as the spike protein, nucleocapsid protein, and the whole virus, with very low dissociation constants. A universal aptamer has been developed capable of detecting not only the original SARS-CoV-2 strain but also several variants, including alpha, beta, gamma, epsilon, kappa, delta, and omicron. A dimeric aptamer, DSA1N5, was created by combining two previously selected aptamers, demonstrating a detection sensitivity of 100 copies per mL of SARS-CoV-2. Despite evaluating these aptamers in saliva and nasopharyngeal fluid, their potential for detecting SARS-CoV-2 in wastewater remains unexplored. While there have

been advancements in biosensing technologies for clinical samples, their adaptation to wastewater analysis is challenging due to low virus levels and matrix interference.^{86–88}

Facing this challenge, Sen and colleagues aimed to assess the suitability of aptamer technology for wastewater analysis and develop a prototype for aptamer-based wastewater monitoring. They began by examining the stability and specificity of their universal aptamer in wastewater using a dot blot method. Then, they integrated an electrochemical aptamer test with a water filtration, purification, and enrichment system to determine if this setup could achieve the required limit of detection and specificity for wastewater analysis. Finally, they used the integrated system to analyze actual wastewater samples and compared its performance with that of RT-qPCR.⁶⁸

They combined an aptamer-based electrochemical chip with a filtration, purification, and extraction system to create an on-site solution for wastewater analysis (Fig. 14). Using a dimeric aptamer effective against wild-type and variants like alpha, delta, and omicron, they confirmed its stability in wastewater (diluted to 50%) with minimal impact on binding affinity. The sensing chip demonstrates a detection limit of 1000 copies per L (1 copy per mL), enhanced by the filtration, purification, and extraction system. This integrated setup can detect trace virus levels (<10 copies per mL), moderate contamination (10–1000 copies per mL), or high levels (>1000 copies per mL) in native wastewater, offering a practical in-field wastewater analysis solution.⁶⁸

Current limitations and improvements of point-of-care diagnostics for wastewater. RT-LAMP stands out for its sensitivity, simplicity, and speed. Recent advancements have focused on employing RT-LAMP for detecting SARS-CoV-2, utilizing fluorescence detection, colorimetric assays, and electrochemical sensors. However, these methods often require bench-top instruments or lack automated measurement protocols.⁶⁵ One challenge with LAMP in quantification is its limited ability to precisely quantify target nucleic acids. Unlike quantitative PCR, which can provide accurate measurements of nucleic acid

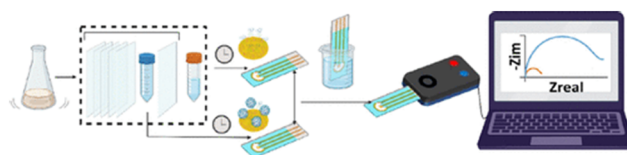


Fig. 14 Graphical abstract of aptamer-based electrochemical chip with a filtration, purification, and extraction system to create an on-site solution for wastewater analysis.⁶⁸ Samples undergo filtration, purification, and extraction to concentrate nucleic acids, including SARS-CoV-2 RNA. Dimeric aptamers on an electrochemical chip bind specifically to SARS-CoV-2 RNA or proteins, enhancing binding strength. Binding causes a structural change, altering electrochemical properties detected as redox current. Signal amplification by filtration, purification, and extraction precedes quantification, categorizing viral load levels.

concentrations, LAMP does not offer the same level of quantitative precision. This is primarily due to the exponential amplification nature of LAMP, which can lead to non-linear amplification kinetics and difficulties in establishing a direct correlation between the amount of starting material and the final amplification product. As a result, while LAMP is highly sensitive for detection purposes, its use in quantitative analysis may require additional calibration steps or complementary techniques for accurate quantification.⁸⁹ Recent study aimed to employ a quantitative LAMP approach to assess SARS-CoV-2 viral load, utilizing threshold time (TT) values as indicators. Analogous to conventional qPCR's cycle threshold (C_t) values, qLAMP's TT values exhibit a linear relationship with SARS-CoV-2 copy numbers. The RT-qLAMP assay efficiently quantifies viral loads within 15 minutes (fluorescent reading) or 25 minutes (colorimetric reading) for synthesized RNA, inactivated viral particles, and clinical samples. It successfully detects various SARS-CoV-2 variants and the human beta-actin gene, serving as a quality control indicator. RT-qLAMP results align closely with RT-qPCR, detecting all clinical samples within 10 minutes *via* fluorescent reading, even those with $C_t = 35$.⁹⁰ LAMP also encounters challenges in proof-of-concept and clinical diagnostics due to environmental control needs for microfluidic device development, skilled personnel required for integrated chip fabrication, and challenges in automating processes and primer selection. Ensuring primer selectivity and operational conditions further compounds these challenges.⁹¹

Microfluidics provides precise control over experimental environments using laminar flow, fluid mixing, and broader fluid behavior concepts. Microfluidic technology, enabling rapid on-site analysis with minimal reagent usage, reduces time, sample loss, contamination risks, and costs, while enhancing automation and reliability.⁹² However, microfluidic devices, which predominantly made from polydimethylsiloxane (PDMS), face challenges despite their advantages. PDMS processing is time-consuming, prone to swelling in certain solvents, difficult to seal, and exhibits selective molecule extraction. Finding affordable, solvent-compatible materials with versatile processing methods remains a significant obstacle for scaling up production for both research and commercial use. Alternative materials like thermoplastics and processing techniques such as hot embossing and 3D printing require further exploration to meet application requirements. Moreover, microfluidic devices often rely on external equipment, hindering accessibility and scalability. Integrating smaller, cost-effective components into machines could alleviate this issue. Streamlining sample preparation procedures and developing programmable microfluidic chips would further enhance accessibility and versatility, overcoming current limitations in reliability and compatibility between component chips.⁹³ Enhancing microfluidic accessibility hinges on developing cost-effective materials compatible with various solvents,

seamlessly transitioning from research to commercialization. Additionally, devising fluid movement techniques devoid of external pumps and microfluidic chips performing specific sample preparations is imperative.⁹⁴

CRISPR-Cas offers significant benefits for point-of-care testing in resource-limited environments. It provides real-time patient information, aiding in better treatment decisions and improving outcomes. CRISPR-based tests are affordable, sensitive, user-friendly, and independent of complex equipment, making them ideal for on-site point-of-care testing. With short detection cycles, low cost, high sensitivity, and versatile readout methods, CRISPR-Cas systems reduce dependency on PCR-based instruments, promising effective and accessible diagnostics.⁹⁵ Its distinctive ability to precisely target and cleave DNA sequences makes CRISPR-Cas technology a ground-breaking tool for diagnostics, genomic and molecular editing. Despite the notable progress, most CRISPR-Cas sensors primarily detect nucleic acids, limiting their broader applicability. Challenges include oversensitivity, off-target recognition, and false positives due to the system's tolerance for mismatched nucleotides, hindering widespread adoption.⁹⁶ Despite hurdles, CRISPR technology promises to revolutionize sensing and enhance POCT globally.⁹⁷

Aptamers, folded into 3D nanostructures *via* specific base pairing, offer increased sensitivity and selectivity. Their feasibility for chemical synthesis allows sequence-specific modifications to enhance binding affinity. Aptamers' self-amplification enhances biosensor sensitivity, while their binding-induced structure-switching improves selectivity. Adaptability to evolving biosensor requirements, such as real-time and in-field detection, is highlighted. Their modular nature facilitates multifunctional biosensor development, promising advancements beyond mere binding strength enhancement.⁹⁸ However, commercialization of aptamers and aptameric biosensors faces several challenges. Aptamer generation *via* SELEX technologies is time-consuming and labor-intensive, with a low success rate.⁹⁸ PCR during SELEX technology presents challenges due to the complexity of the reaction mixture and the generation of biased and artifactual products. Factors such as unequal primer hybridization, varying polymerase efficiency, and intertemplate hybridization contribute to this bias.⁹⁹ Addressing these challenges is crucial for the advancement of biosensors utilizing aptamers.

Point-of-care diagnostics represent a promising avenue for rapid and on-site analysis, particularly in the context of wastewater management. However, several limitations persist in their application to wastewater analysis, necessitating improvements to enhance their efficacy and reliability. One significant limitation lies in the complexity and heterogeneity of wastewater samples.¹⁰⁰ Unlike clinical samples, which are relatively standardized, wastewater can contain a myriad of substances, including organic matter, chemicals, and microbial contaminants.

Wastewater samples may contain compounds that cross-react with assay reagents or interfere with detection mechanisms, leading to false-positive or false-negative outcomes.¹⁰¹ The presence of interfering substances can compromise the accuracy of results obtained from point-of-care diagnostics systems. Moreover, the current lack of standardization and validation protocols specific to point-of-care diagnostics in wastewater analysis hinders their widespread adoption. Establishing standardized procedures for calibrating, validating, and quality-assuring point-of-care diagnostics platforms is imperative for ensuring the accuracy and comparability of results across different settings.¹⁰²

In summary, while point-of-care diagnostics holds immense potential for wastewater analysis, overcoming current limitations requires concerted efforts towards technological innovation, methodological refinement, and standardization initiatives. By addressing these challenges, point-of-care diagnostics can become indispensable tools for monitoring and managing wastewater resources effectively.

Conclusion

The integration of point-of-care diagnostics into SARS-CoV-2 wastewater-based epidemiology marks a notable advancement in COVID-19 surveillance. While RT-PCR has demonstrated high sensitivity, its complexity and reliance on specialized equipment pose challenges. Point-of-care testing methods, such as LAMP, microfluidic technology, field-effect transistors, biosensors, and CRISPR/Cas, offer rapid and accessible alternatives crucial for timely detection and surveillance efforts. Combining LAMP with microfluidic technology and paper-based devices further enhances their utility for on-site analysis. Furthermore, the development of aptamer-based electrochemical chips provides solutions with high specificity and stability, facilitating practical monitoring of viral transmission in wastewater. Despite current challenges associated with sample complexity and standardization, ongoing research aims to refine these point-of-care diagnostics to improve their effectiveness and reliability. In summary, point-of-care diagnostics offer a revolutionary approach to wastewater-based epidemiology, providing promising prospects for efficient disease surveillance and management beyond the scope of SARS-CoV-2.

Data availability

No primary research results, software or code have been included and no new data were generated or analysed as part of this review.

Conflicts of interest

The authors declare no competing financial interest.

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