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Development of a paper-based fluorescent carbon quantum dots MIPs sensor for selective detection of lumpy skin disease virus

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Lumpy skin disease (LSD) is a contagious viral disease caused by the Lumpy Skin Disease virus (LSDV), a member of the *Capripoxviridae* family. Traditional LSDV diagnostic procedures proved to have challenges in terms of cross reactivity as well as limited sensitivity and specificity. Herein, we combined molecularly imprinted polymers (MIPs) and quantum dots (QDs) technology to develop a paper-based turn on fluorescence sensor for rapid, sensitive and selective detection of LSDV. Under optimal conditions, the sensor showed linear enhancement in fluorescence intensity with the increase of LSDV concentration and exhibited a detection limit of $10^1 \log_{10}$ TCID₅₀ per ml. It also presented high specificity towards LSDV compared to other viruses *viz* sheep pox virus (SPV). Furthermore, the proposed sensor was successfully tested with spiked and real LSDV samples, proving its potential to serve as a sensitive selective sensor for LSDV diagnosis. Based on our knowledge, this is the first record of a paper-based diagnostic sensor for LSDV utilizing a CQDs-MIPs turn-on mechanism.

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1. Introduction

The OIE lists LSDV as one of the most hazardous animal diseases¹ causing considerable economic losses in terms of skin damage, reduced milk output, abortions and deaths in most of the infected ruminants.² The virion is a brick-shaped particle measuring 170–260 by 300–450 nm in diameter.³ Its capsid contains a linear, non-segmented, double-stranded DNA genome of ~150 kilobases. The genomes of sheep pox virus (SPV) and goat pox virus (GPV), members of capri pox family, exhibit significant resemblance of >97% nucleotide identity.⁴ Previously, they were classified solely based on their animal host. However, nowadays, it is possible to differentiate them by analyzing their genomic DNA.

Traditional laboratory techniques including virus isolation and serological testing are usually used for LSDV diagnosis.⁵ These strategies are generally challenging and time-consuming. Molecular techniques including reverse transcription PCR can detect LSDV in clinical samples with sufficient sensitivity.⁶ However, it is unsuitable for field diagnosis due to the requirement of specialized equipment. Novel quick point-of-care biosensing methods with high sensitivity and specificity are required for early LSDV detection leading to controlling infection spread.⁷

Nanotechnology can be used to create novel materials and products and is recognized as one of six main enabling technologies that will revolutionize society in the near future.⁸ CQDs are zero-dimensional carbon nanoparticles that are usually smaller than 10 nm and have a crystal constant of 0.34, which is equivalent to the *d*-spacing of graphite (002). The carbonyl, carboxyl, epoxy, and hydroxyl functional groups located on the carbon core of CQDs enhances their water solubility and provide a platform for simple functionalization with other substances.^{9,10} The presence of functional groups on the surface of CQDs enhances their biocompatibility, optical properties, and targeting ability.¹¹

Carbon Quantum dots (CQDs), a superior class of fluorescent nanostructures made from carbon materials, have garnered interest in biosensing applications due to their high biocompatibility, ease of synthesis, outstanding photoluminescence, and tailored surface functionalities. Yet, target selectivity is the main obstacle to CQD-based sensor development.^{12–14}

Recently, CQDs have been widely employed in different fields including biological imaging,¹⁵ biochemical sensing,^{16–19} biomedicine,²⁰ and catalysis.²¹ Furthermore, notable advancements were achieved in the identification of various molecules through the incorporation of CQDs.^{17–19} Various types of QDs were developed for the viral detection, for instance, cadmium telluride and carbon QDs were incorporated in immunosensor platform for the detection of human immune deficiency virus while molybdenum trioxide QDs were used for detection of influenza A virus.^{22–24}

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The selectivity of CQD-based sensors can be improved by Molecularly Imprinted Polymers, or MIPs. These sensors have proven to be very effective at detecting traces of chemicals in complex matrices.^{25,26} MIPs have advanced significantly as a result of the desire to enhance disease diagnosis and prevention through the nanoscale detection of harmful viruses and other biological targets. Although there are still many obstacles to overcome, imprinting techniques can quickly and accurately identify viruses, making them useful as novel sensing materials.²⁷

MIPs are able to recognize templates selectively²⁸ by creating templates with functional monomers and cross-linking monomers. After the eluting of the templates, the cavities with high specificity and affinity for the target molecule become apparent.^{29,30} MIPs components (template, monomers, cross-linker, polymerization initiator, and solvent) and the nature of their interactions with one another have a significant impact on the efficacy, affinity, and selectivity of the recognition sites of the MIPs. Therefore, choosing the right components for the MIPs during design is essential to achieving the required qualities for a particular application.³¹

MIPs have been employed extensively in the sensor area in recent years due to their inexpensive cost of preparation, simplicity, and durability.^{32,33} So, MIPs have been synthesized on the surface of CQDs to improve their poor selectivity as graphene QDs,³⁴ CdSe QDs,³⁵ CdTe QDs,³⁶ and CQDs.³⁷

To our knowledge, a paper-based fluorescent CQDs- MIP-sensor for LSDV determination has not yet been reported. In this study, we combined the high selectivity of MIPs with the strong stable fluorescence of CQDs to develop a simple method for LSDV detection.

2. Experimental section

2.1 Equipment

Attenuated total Reflectance-Fourier transform infrared spectroscopy (ATR-FTIR), PerkinElmer, the UK, Atomic force microscope by a Nanosurf C 3000 Atomic force microscope (AFM), Switzerland and all AFM images were operated in contact mode using a Nanosurf SNL-10 silicon tip, Field emission scanning electron Microscope (FE-SEM) (Quanta FEG250, Czech), transmission electron Microscope (TEM) (JEOL JEM-1400, USA), NANOTRAC WAVE || Dynamic light scattering (DLS), USA, X-ray photoelectron spectroscopy (XPS) (Escalab250Xi, Thermo Fisher, USA), Muffle furnace (Nbertherm, Germany) and BioTek Synergy HTX Multimode Reader, Agilent, USA were used in this study.

2.2 Synthesis of surface modified APTES-CQDs

CQDs was synthesized and surface functionalized by mixing 0.5 g of anhydrous citric acid (CAS No.: 77-92-9) with 10 ml of 3-Aminopropyl triethoxysilane (APTES) monomer (CAS No.: 919-30-2) according to.¹² The mixture was transferred into a 50 mL Teflon-lined autoclave and kept in a muffle at 210 °C for 5 h for carbonization and surface functionalization. The colour of the solution turned colourless confirming the synthesis of the

surface modified APTES-CQDs. The solution was then filtered through 0.45 µm filter membrane and stored at 4 °C for further use. DLS, UV-vis absorption microplate reader, XPS, ATR-FTIR and TEM were used for APTES-CQDs characterization. All used glassware was washed by detergent and tap water several times, then rinsed by deionized water and ethanol, then dried in a hot air oven.

2.3 Fabrication of the paper-based CQDs- MIP-sensor

MIPs were synthesized using the following polymeric mixture; 13.0 mg of acrylamide (CAS No.: 79-06-1), 10.4 µl of methacrylic acid (CAS No.: 79-41-4), 6.4 µl of methyl methacrylate (CAS No.: 80-62-6) and 6 µl of N-vinyl pyrrolidone (CAS No.:88-12-0) monomer were mixed with 48 mg of *N,N*-(1,2-dihydroxyethylene) bisacrylamide (CAS No.: 868-63-3) as a crosslinker. The prepared mixture was dissolved in 300 µl of DMSO (CAS No.: 67-68-5) containing 2.4 µl of 2, 5-BIS (*tert*-butyl peroxy)-2, 5-dimethylhexane (CAS No.: 78-63-7) as an initiator mixed with 10 µl of APTES-CQDs. MIPs synthesis and polymerization were conducted using similar methodology to Kassem *et al.*³⁷ with modifications including the use of CQDs as a cost-effective fluorescence source instead of N-fluorescein acrylamide monomer, and we also used APTES monomer to functionalize the CQDs surface to ensure proper binding of other monomers forming the MIPs.

Afterwards, the mixture was subjected to pre-polymerization at 80 °C for 1 h under stirring followed by 1 h at room temperature. 7 µl of 0.25 mg µl⁻¹ LSDV (provided by Vaccine and Serum Research Institute, Egypt) was pipetted on a sterile glass slide surface and allowed to dry at room temperature in a sterile, clean laminar flow cabinet in order to prepare the LSDV-template. The nitrocellulose membrane (NCM, HF180MC100, Millipore, Germany) was cut into 6 mm circular pieces using a normal sterile paper puncher. NCM pieces were dipped into the pre-polymerization solution and pressed directly onto the dried virus drop to prepare the CQDs-MIPs. Whereas to prepare the negative control CQDs-NIPs, the dipped NCM was pressed onto an empty sterile glass slide. Both NIPs and MIPs were exposed overnight to UV radiation with a wavelength of 254 nm until the NCM surface showed signs of a thin layer of polymerized film. Subsequently, they were submerged in 10% HCL (CAS No.: 7647-01-0, Sigma-Aldrich, UK) for ten seconds to denature and eliminate the viral molecules followed by their incubation for 2 h at 45 °C in deionized water under shaking conditions before being dried on a filter paper.

The chemical composition and binding of the functional groups between LSDV and CQDs-MIPs were evaluated using ATR-FTIR at the scanning range of 4000–400 cm⁻¹. The surface morphology of the CQDs-NIPs and CQDs-MIPs NCM were examined using AFM and FE-SEM.

2.4 LSDV-CQDs-MIP sensor validation

The LSDV-CQDs-MIP sensor's sensitivity was assessed by measuring the fluorescence intensity that was observed at various LSDV concentrations (10¹–10⁶ log₁₀ TCID₅₀ per ml). In



summary, a 96-well black fluorescent microplate containing CQDs-NIPs and CQDs-MIPs NCM was loaded with 20 μl of each LSDV dilution. To ensure that the viral particles were evenly distributed across the NCM, the reaction was incubated for 15 min under orbital shaking. The fluorescence intensity was then measured at 485/20 nm for excitation and 528/20 nm for emission.

Similar procedures were used to determine the assay's specificity, with the exception of the virus types, SPV, FMDV, and BVDV (they were provided by virology department, Animal Health Research Institute, Egypt) at $10^5 \log_{10}$ TCID₅₀ per ml. In order to assess the effectiveness of the sensor in identifying LSDV within the complex matrix of field real samples, 15 samples including viral-free blood buffy coat, serum, and skin nodules were spiked with varying amounts of LSDV (10^1 – $10^6 \log_{10}$ TCID₅₀ per ml). For confirmation and sensitivity comparison, Viral DNA was extracted following QIAamp DNA extraction kit instructions (Cat no. 51306, Qiagen, USA) then subjected to Real-Time PCR (qPCR) testing using LSDV dteqPCR kit (F100, GPS™, Spain) according to instruction manual. All experiments were run in triplicates.

3. Results and discussion

3.1 Characterization of surface modified APTES-CQDs

CQDs have emerged as a promising class of quantum dots. CQDs have garnered significant attention in recent years due to their numerous advantages. These include excellent photoluminescence properties, straightforward synthesis methods, cost-effective precursors, solubility, low toxicity, chemical stability, and ease of functionalization. As a result, they have been utilized in point-of-care sensors.^{38,39}

The synthetic pathways of CQDs can be classified into “top-down” and “bottom-up” techniques based on the carbon source utilized during the synthesis.⁴⁰ The “top-down” strategy, often uses physical or chemical processes, results in small-sized CQDs when a large-scale carbon source is used such as graphite rod, carbon fibers, and carbon nanotubes. On the other hand, “bottom-up” approach always begins with carbon sources of small molecules, such as folic acid, glucose, and citric acid.⁴¹ Bottom-up methodologies for synthesizing CQDs include hydrothermal,⁴² solvothermal,⁴³ microwave-assisted,⁴⁴ ultrasonic-assisted,⁴⁵ and combustion⁴⁶ procedures. The hydrothermal method usually results in CQDs with high quantum yield and strong fluorescence intensity when compared to those derived from the “top-down” approach. Hence, in recent years, this approach has mostly been used in the production of CQDs.^{41,47}

In our study, Surface modified APTES-CQDs were obtained *via* a simple one-pot hydrothermal process. The acylation reaction and surface silanization occurred directly in the presence of APTES. In this process, CQDs were synthesized from anhydrous citric acid which was carbonized under temperature of 210 °C for 5 h. The amine groups in APTES acylated with the carboxyl groups on the surface of the pyrolyzed organics to provide surface APTES-functionalized CQDs with fluorescent characteristics. Subsequently, CQDs-MIPs were successfully

obtained by the polymerization assay using a co-polymer mixture consisted of acrylamide, APTES-CQDs, methacrylic acid, methyl methacrylate and *N*-vinyl pyrrolidone and *N,N*-(1,2-dihydroxyethylene) bisacrylamide with adjusted ratios to enhance sensitivity and specificity of the developing sensor.³⁸

DLS measurement indicated average size of 25.3 ± 2.5 nm with PDI of 0.3 and surface charge of +28 mv (Fig. 1A). The positive charge relates to the presence of amine groups on APTES functionalized CQDs surface. The morphology of APTES-CQDs was characterized by TEM showing the presence of homogenous well dispersed spherical shaped particles with average size of 5.1 ± 0.8 nm (Fig. 1C1 and C2).

FT-IR spectra of APTES and APTES-CQDs are shown in (Fig. 1E) and Table 1. APTES results showed characteristic peaks at 2974, 2928, and 2886 cm^{-1} which corresponds to the stretching vibrations of O–H group. While the peaks at 1577, and 1484 cm^{-1} were attributed to the bending vibrations of the N–H group. Peaks at 1388 cm^{-1} was attributed to the bending vibrations of C–H group. In addition, the peaks at 1167 and 1080 cm^{-1} were attributed to the stretching vibration of C–O group. The peaks at 958 and 773 cm^{-1} indicated the bending vibration of C=C and C-CL groups respectively. Also, the peak at 682 cm^{-1} was attributed to the stretching vibration of the C=C group. Compared with APTES, APTES-CQDs FT-IR spectrum displayed two different absorption peaks at 1724 and 1299 cm^{-1} in addition to the shifting of most functional groups of O–H, N–H, C–H, C–O and C=C groups. The two different peaks at 1724 and 1299 cm^{-1} which are attributed to the stretching vibration of C=O and the bending vibration of the C–H group, together with the shifting of other functional groups, revealed the formation of an amide bond, which demonstrated the successful acylation reaction between CQDs and APTES, suggesting the successful synthesis of APTES-CQDs.

The elemental analysis of APTES-CQDs was conducted by XPS demonstrating four peaks at 533.25, 286.04, 401.73, and 103.91 eV referring to O 1s, C 1s, N 1s, and Si 2p, respectively (Fig. 1D). The results of XPS further confirmed that successfulness of the surface functionalization of CQDs with APTES. The optical properties of APTES-CQDs were examined with UV-vis absorption spectrum showing a strong absorption peak located at 300 nm (Fig. 1B). Furthermore, APTES-CQDs exhibited highest fluorescence at excitation and emission wavelengths of 485/20 nm and 528/20 nm, respectively using a filter-based microplate reader with different Excitation/Emission wavelengths (360/40–460/40, 360/40–528/20, 360/40–590/20, 485/20–528/20, 485/20–590/20), respectively.

The occurred hydrothermal reaction involved chemical interactions between various carbonaceous precursors under high temperature and pressure. This technique is well recognized as an environmentally friendly, regulated, and cost-effective solution.^{48,49} The synthesis of CQDs typically comprises four processes: breakdown, polymerization/aromatization, nucleation, and growth. Initially, the carbonaceous precursors undergo intermolecular dehydration and polymerization, forming the carbon skeleton and aromatic clusters. When clusters achieve critical super saturation, CQD nucleation occurs. During this process, aromatic clusters move



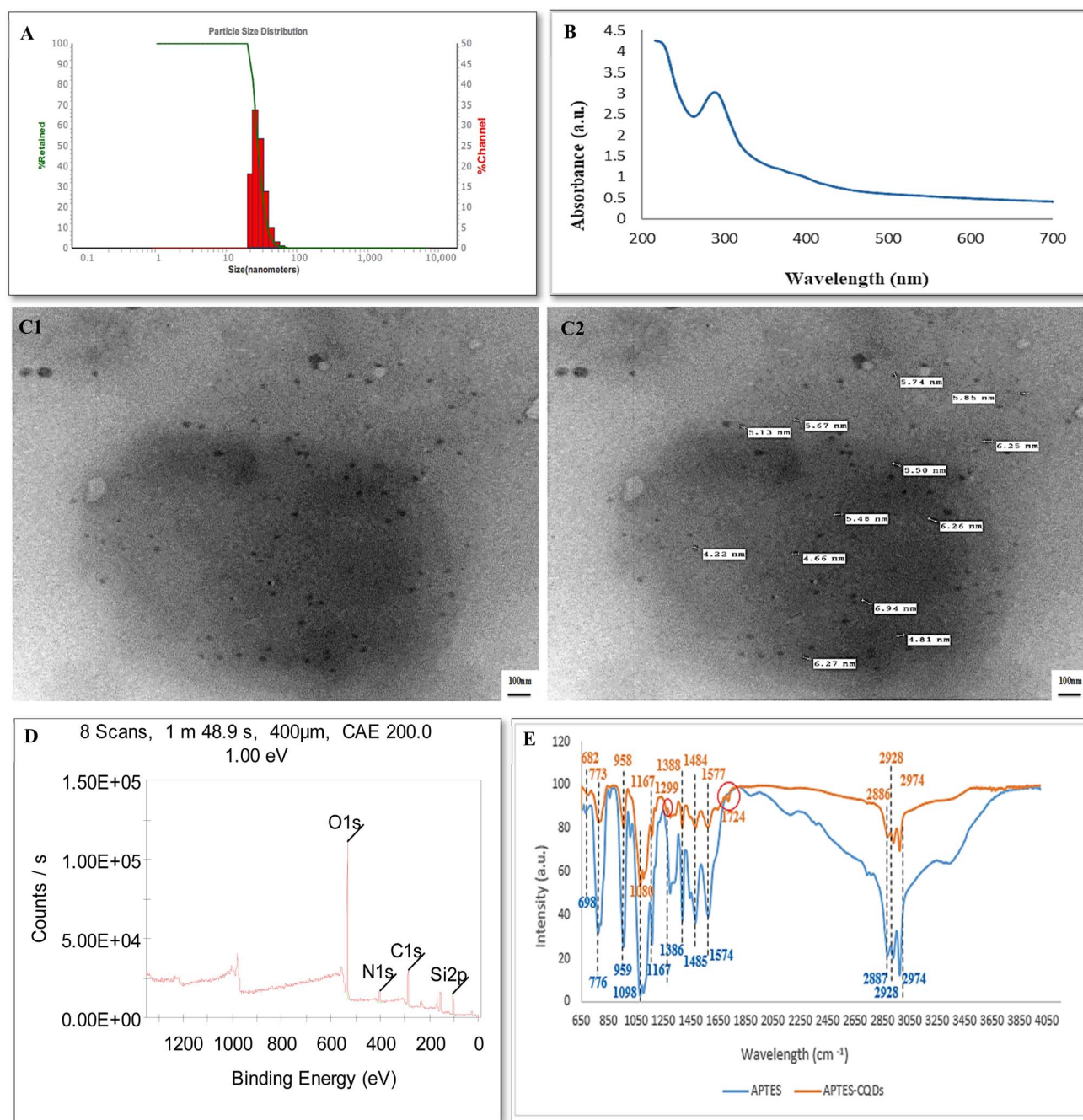


Fig. 1 Characterization of APTES-CQDs. (A) Particle size pattern of APTES-functionalized CQDs showing size distribution of 25.3 ± 2.5 nm by DLS. (B) UV-visible absorption spectra of APTES-functionalized CQDs. (C1 and C2) TEM image of APTES-functionalized CQDs with average size of 5.1 ± 0.8 nm; (D) XPS spectrum of APTES-functionalized CQDs; (E) FT-IR spectra of APTES and APTES-functionalized CQDs.

towards the surface of the particle, resulting in the functionalization of the surface of CQDs. As the reaction progresses, intermediates are gradually transformed into fully formed CQDs with a narrow particle size distribution.⁵⁰ The size of the formed CQDs produced depends on temperature, time, and reactants used in the process.⁵¹ Surface functionalization of CQDs is accomplished simultaneously during the hydrothermal process. This phase involves modifying the fluorophore, passivating the surface, and oxidizing the surface of all

synthesized CQDs. Most CQDs exhibit a strong hydrophilicity and their surfaces are typically treated with oxygen-containing functional groups.⁵² Usually, CQDs hydrophilicity is influenced by both the reactant and the synthesis process. For instance, CQDs derived from soluble precursors typically display significant hydrophilicity due to the retention of functional groups from the precursors on the CQD surface during their fabrication. Besides, the choice of reactants affects CQDs optical characteristics. Moreover, elemental analysis showed

Table 1 FT-IR spectra data of the APTES and APTES-functionalized CODs

Material	Absorption (cm ⁻¹)	Chemical bond	Vibration mode	Functional group	
APTES	2974	O-H	Stretching	Carboxylic acid	
	2928	O-H	Stretching	Carboxylic acid	
	2886	O-H	Stretching	Carboxylic acid	
	1577	N-H	Bending	Primary amine	
	1484	N-H	Bending	Secondary amine	
	1388	C-H	Bending	Aldehyde	
	1167	C-O	Stretching	Alcohol	
	1080	C-O	Stretching	Alcohol	
	958	C=C	Bending	Alkene	
	773	C-CL	Bending	Halo compound	
	682	C=C	Stretching	Alkene	
	APTES functionalized CQDs	2974	O-H	Stretching	Carboxylic acid
		2928	O-H	Stretching	Carboxylic acid
		2887	O-H	Stretching	Carboxylic acid
1724		C=O	Stretching	Carboxylic acid	
1574		N-H	Bending	Primary amine	
1485		N-H	Bending	Secondary amine	
1386		C-H	Bending	Aldehyde	
1299		C=O	Bending	Aromatic ester	
1167		C-O	Stretching	Alcohol	
1098		C-O	Stretching	Alcohol	
959		C=C	Bending	Alkene	
776		C-CL	Bending	Halo compound	
698		C=C	Stretching	Alkene	

that the reaction time and reactant type had a direct impact on the amount of nitrogen atoms doped into the carbon core. When comparing CQDs synthesized from secondary and tertiary amine to those made from citric acid and primary amine, the latter presented the highest nitrogen doping degree and the maximum photoluminescence intensity.⁵² Herein, surface functionalization of CQDs with APTES was confirmed through FTIR and XPS analysis.

3.2 Characterization of the CQDs-MIPs-NCM

Molecular imprinting involves the synthesis of polymers that possess molecular recognition sites that precisely correspond to the size, shape, and functional group of template molecules. Due to their low selectivity, CQDs should be paired with other technologies *viz* MIPs to enhance their selectivity while maintaining cost-effectiveness.^{53,54}

After successfully synthesizing the CQDs-MIPs, chemical composition and involved functional groups in the binding of LSDV with CQDs -MIPs were evaluated using ATR-FTIR at scanning range of 4000–400 cm⁻¹.⁴⁸ Hence, the FTIR spectra of CQDs -MIPs were compared before and after binding with LSDV (Fig. 2 and Table 2) to inspect the altered functional groups upon binding. For CQDs -MIPs before LSDV binding, the FTIR revealed peaks located at 3414, 2915, 2133, and 1690 cm⁻¹ that are referred to the stretching vibration of N-H, O-H, N=N=N, C=O groups, respectively. Also, peaks were detected at 1630, 1542, 1436 and 1409 cm⁻¹ indicate the bending vibration of N-H, N-H, O-H and O-H groups, respectively. While the peaks detected at 1315, 1272 and 1023 cm⁻¹ correspond to the stretching vibration of -NO₂, C-O and C-O groups, respectively.

In addition, peaks detected at 954 and 707 cm⁻¹ refer to the bending vibration of C=C group and the stretching vibration peak of C-CL, respectively. FTIR spectra of the CQDs -MIPs after rebinding with LSDV revealed shifting of the functional groups N-H, O-H and C=O represented by peaks at 3394, 2961 and 1695 cm⁻¹ with stretching mode, respectively. Also, shifting of two N-H groups were detected at 1631 and 1540 cm⁻¹. Likewise, O-H, -NO₂ and C-O groups shifting were detected at 1410, 1316 and 1270 cm⁻¹, respectively. Furthermore, new bonds were recorded at 1720 cm⁻¹ which referred to C=O groups with stretching mode. Additionally, the two functional groups, N=N=N and O-H, represented previously by peaks at 2133 and 1436 cm⁻¹ Disappeared. The occurred changes in the FTIR spectra peaks approve the successfulness of the formation of LSDV-CQDs-MIPs complex and the involvement of multiple

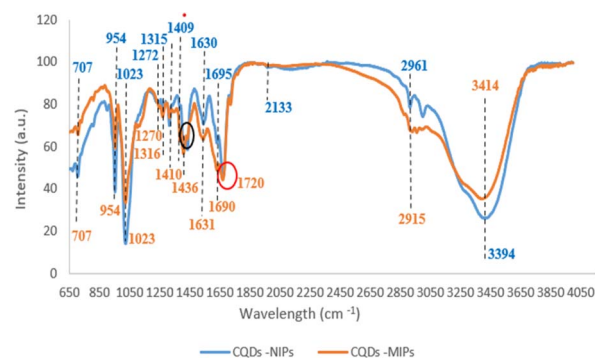


Fig. 2 FT-IR spectra of the CQDs-NIPs and CQDs-MIPs with LSDV.



Table 2 FT-IR spectra data of the CQDs-MIPs and CQDs-M IPs after LSDV rebinding

Material	Absorption (cm ⁻¹)	Chemical bond	Vibration mode	Functional group	
CQDs-MIPs before LSDV binding	3414	N-H	Stretching	Amine	
	2915	O-H	Stretching	Carboxylic acid	
	2133	N=N=N	Stretching	Azide	
	1690	C=O	Stretching	Primary amide	
	1630	N-H	Bending	Primary amine	
	1542	N-H	Bending	Secondary amine	
	1436	O-H	Bending	Carboxylic acid	
	1409	O-H	Bending	Carboxylic acid	
	1315	-NO ₂	Stretching	Aliphatic nitro	
	1272	C-O	Stretching	Aromatic ester	
	1023	C-O	Stretching	Alcohol	
	954	C=C	Bending	Alkene	
	707	C-Cl	Bending	Halo compound	
	CQDs-MIPs after LSDV rebinding	3394	N-H	Stretching	Amine
		2961	O-H	Stretching	Carboxylic acid
1720		C=O	Stretching	Carboxylic acid	
1695		C=O	Stretching	Primary amide	
1631		N-H	Bending	Primary amine	
1540		N-H	Bending	Secondary amine	
1410		O-H	Bending	Carboxylic acid	
1316		-NO ₂	Stretching	Carboxylic acid	
1270		C-O	Stretching	Aliphatic nitro	
1023		C-O	Stretching	Aromatic ester	
954		C=C	Bending	Alcohol	
707		C-Cl	Bending	Alkene	

functional groups indicates the presence of strong efficient binding of between the synthesized CQDs-MIPs and LSDV.

Using AFM, the topographical properties of CQDs-NIPs, CQDs-MIPs, and CQDs-MIPs-LSDV complex were observed and documented. These features included structure, size of pores, and degree of roughness and arrangement of imprinted sites on the imprinted polymeric layer. AFM (3D) images were recorded for the imprinted bare CQDs-MIPs following polymerization (Fig. 3A), CQDs-MIPs-LSDV complex after template removal (Fig. 3B) and CQDs-NIPs (Fig. 3C). Significantly, imprinted bare CQDs-MIPs layers demonstrated homogenous distribution across the surface, exhibiting height of 5.89 nm and a roughness of 1.314 ± 0.05 nm (Fig. 3A) with N grade of (01). The rebinding of LSDV to its specific empty sites on the bare MIP resulted in an increase in both; surface height from 5.89 to 17.8 nm and roughness from 1.314 ± 0.05 to 10.405 ± 0.04 nm with N grade of (0) demonstrating successful rebinding (Fig. 3B). Whereas, CQDs-NIPs exhibited a smooth surface with 1.95 nm height and 301.21 ± 0.5 pm roughness (Fig. 3C). FE-SEM images demonstrate the topographical morphology of the CQDs-MIPs empty cavities after template removal *versus* LSDV filled cavities and CQDs-NIPs (Fig. 4A-C). CQDs-NIPs thin film arranged uniformly on the surface of NCM (Fig. 4A1 and A2) with a range size of 500.8–650.7 nm. CQDs-MIPs directly after washing showed well distributed holes on the MIPs film with the range size of 850 nm–2.45 μ m (Fig. 4B1 and B2) which may indicate the agglomeration of virus clusters in some imprinting sites. In CQDs-MIPs-LSDV complex images, the holes appeared to be uniformly sealed with viral particles,

particularly in binding sites, following the rebinding of LSDV template (Fig. 4C1 and C2). AFM and FE-SEM images showed compatible results approving the successfulness of the imprinting process.

In this study, APTES-CQDs was copolymerized with other functional monomers as acrylamide, methacrylic acid, methyl methacrylate, N-vinyl pyrrolidone in the presence of bisacrylamide as a cross linker to imprint LSDV template forming recognition cavities inside the polymer structure. The results of ATR-FTIR, AFM, and FE-SEM were in accordance proving the successfulness of the imprinting process on the NCM surface.

3.3 LSDV-CQDs-MIP sensor validation

The change in fluorescence intensity in response to the addition of various LSDV concentrations (10^1 – 10^6 log₁₀ TCID₅₀ per ml) was evaluated in order to examine the sensitivity and recognition ability of the LSDV-CQDs-MIP sensor. Comparing its result to the negative control, a considerable increase in the fluorescence intensity was detected (Fig. 5A). Furthermore, the results showed direct correlation between measured fluorescence signal and LSDV concentration. When operating under optimum conditions, LSDV-CQDs-MIP sensor's detection limit was found to be 1 log₁₀ copies per ml, which is comparable to the C_T value of 37 of qPCR findings. In this study, we successfully performed a novel, rapid, accurate, sensitive, and specific tool for LSDV compatible with PCR findings and can be applied to clinical samples at a large scale in a short time, which will be helpful in the diagnosis of clinical samples in case of outbreaks. Within 15 minutes, our assay could identify LSDV with high



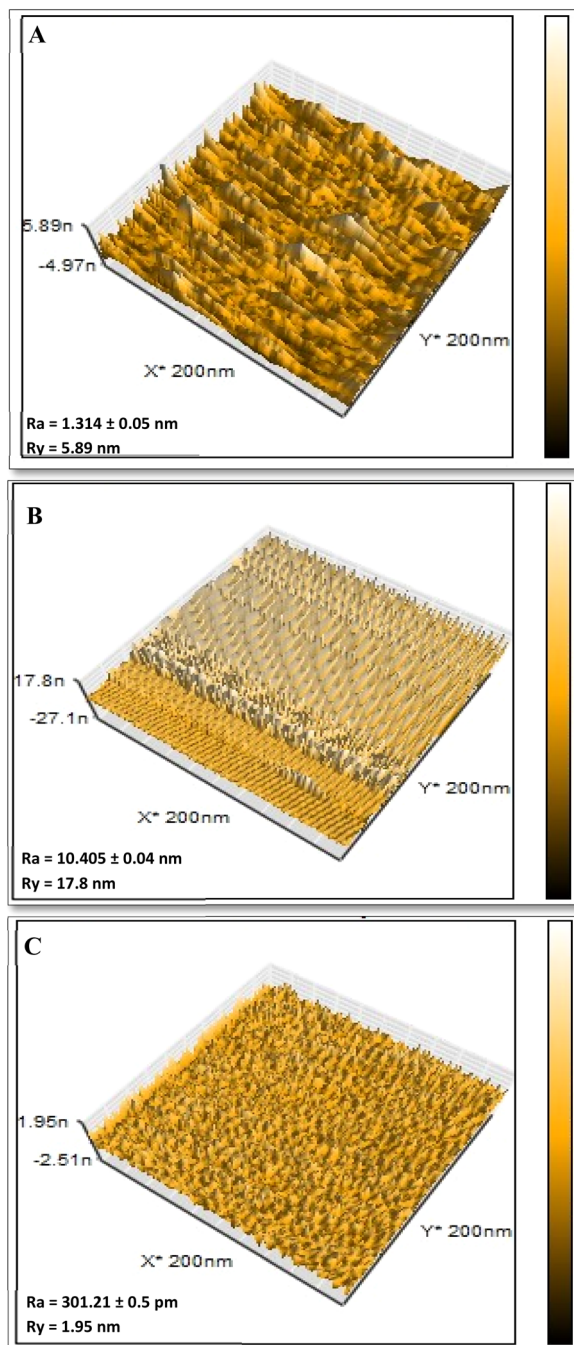


Fig. 3 AFM images (3D) of (A) of the imprinted bare CQDs-MIPs following polymerization at scale of 200 nm, (B) rebinding of LSDV with specific recognition sites on the bare CQDs-MIPs surface after removal of template at scale of 200 nm. (C) CQDs-MIPs without LSDV template virus showing smooth surface at scale of 200 nm. Ra is the surface roughness and Ry is the surface height.

fluorescence intensity. The presence of sufficient empty cavities for LSDV rebinding in a specific manner results in its rapid detection. Moreover, our sensor demonstrated the same sensitivity in detecting LSDV in complex sample matrixes, spiked or real (Fig. 5C). Fluorescence intensities just before (F0) and after (F) addition of LSDV template are shown in (Fig. 5D).

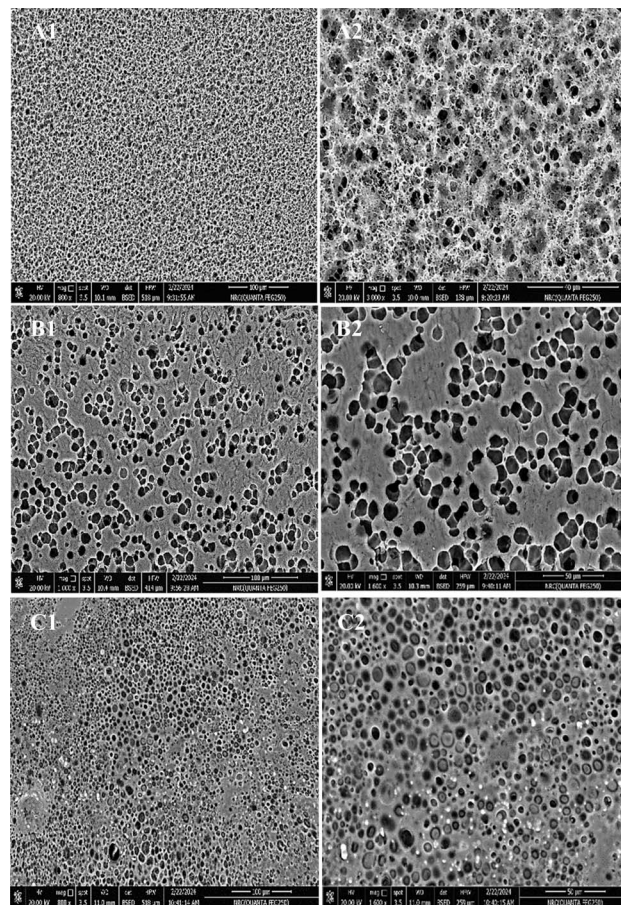


Fig. 4 FE-SEM images of (A1 and A2) the CQDs-NIPs formed without LSDV template virus, (B1 and B2) CQDs-MIPs after washing showed well distributed holes on the surface of MIPs film with the average size of 850 nm–2.45 μm . (C1 and C2) CQDs-MIPs after LSDV template rebinding at concentration of $10^6 \log_{10}$ TCID₅₀ per ml showed blocked holes with the viral particles specifically in the binding sites.

Additionally, the fluorescence intensity changes in response to the presence of SPV, FMDV, and BVDV at a concentration of $10^5 \log_{10}$ TCID₅₀ per ml was used to assess the specificity of the LSDV-CQDs-MIP sensor (Fig. 5B). Following the addition of samples, a significant reduction in the recorded fluorescence intensity was observed indicating a turn-off activity. This behaviour might be the consequence of these viruses' incompetence to fit as precisely into the empty imprinted cavities as LSDV due to the lack of the essential functional groups required to bind to the imprinted surface selective recognition site resulted in weaker or no attachment. Hence, it can distinguish between members of the *Capripoxviridae* family that attain identical morphological characteristics as well as other possible viral contaminants could be present within the sample. These findings support the developed sensor's eligibility for application as an effective tool for LSDV detection in a quantitative manner. It is the first sensor incorporating fluorescent CQDs and MIPs loaded on NCM, and there are few studies that design rapid diagnostic tools for LSDV. Although we mentioned all recent studies designed rapid diagnostic tools for LSDV as



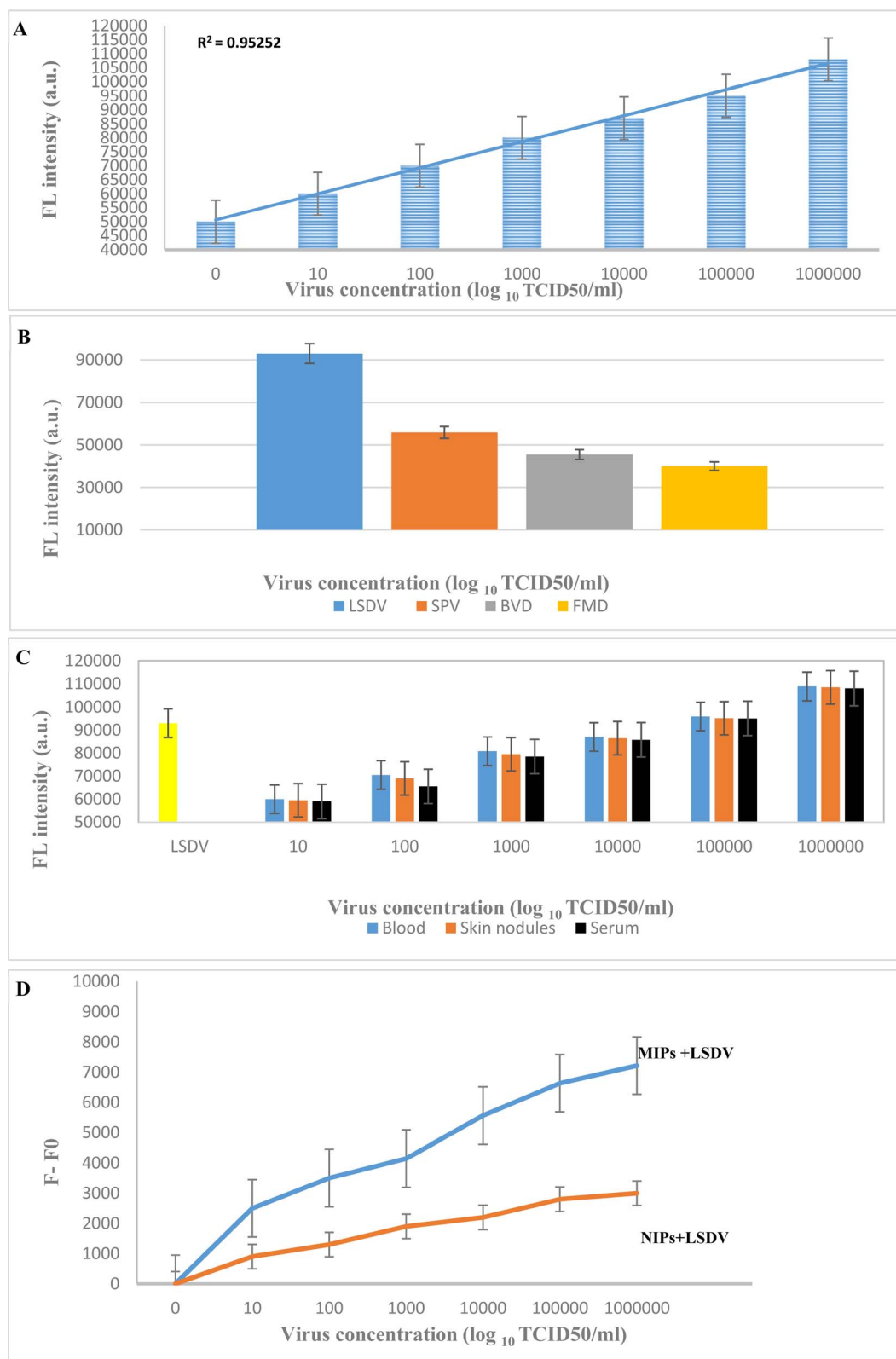


Fig. 5 (A) Sensitivity test of the developed LSDV-CQDs MIPs assay at different concentrations of LSDV (10^1 – 10^6 \log_{10} TCID₅₀ per ml), (B) Selectivity test of the LSDV-CQDs MIPs assay on SPV, BVDV and FMDV viruses at concentration of $10^5 \log_{10}$ TCID₅₀ per ml. (C) Application of the LSDV-CQDs MIPs assay on real blood, skin nodules and serum samples spiked with different concentrations (10^1 – 10^6 \log_{10} TCID₅₀ per ml) of LSDV template. (D) F_0 and F represent the fluorescent intensities before and after addition of LSDV template, respectively.



Table 3 Rapid diagnostic tools for LSDV

Assay	LOD	Advantages	Disadvantages	Reference
Real-time recombinase polymerase amplification (RPA)	15 copies per μL	High sensitivity High specificity Short detection time	Specific primer design High price of reagents Require specific instruments	55
A colorimetric sandwich-type lateral flow immunoassay (LFIA)	$10^{3.4}$ TCID ₅₀ per mL	High inter- and intra-assay repeatability Specificity	Qualitative nature not quantitative Critical antibody preparation step Time consuming in case of large scale of samples	56
Gold nanoparticles-lateral flow test	—	Accurate Rapid	Low sensitivity Monoclonal and polyclonal antibody preparation	57
Loop-mediated isothermal amplification (LAMP)	8 copies per μL	Simple Specific Accuracy similar to PCR	Complex primer design Require expensive kits	58

a Real-Time Recombinase Polymerase Amplification Assay⁵⁵ (RPA), a colorimetric sandwich-type lateral flow immunoassay (LFIA),⁵⁶ gold nanoparticles-lateral flow test⁵⁷ and Loop-mediated isothermal amplification (LAMP).⁵⁸ The advantages and disadvantage of these assays are in details in Table 3. Sensors and biosensors based on CQDs can work with variable mechanisms such as fluorescence static and dynamic quenching, fluorescence-resonance energy transfer (FRET), inner-filter effect (IFE), and photo-induced electron transfer (PET).³⁹ In FRET, the interaction between two fluorescent components results in the transfer of energy from a greater energy donor to a lower energy acceptor. Typically, the distance between the two fluorescent components should be around 10 nm.⁵⁹ Our proposed LSDV-CQDs-MIPs sensor depends on the FRET mechanism. In a previous study,³⁸ it was noticed that LSDV has a fluorescence intensity at Excitation/Emission wavelength (360/40–460/40). Since both LSDV and CQDs have fluorescence intensity, when they come in close contact due to the fitting of LSDV in the specific imprinted groove of CQDs labeled MIPs, their emission spectra overlapped. Hence, the high fluorescent signal. When the other viruses could not fit into the groove, a low fluorescent signal was detected. This proves the ability of the sensor to specifically detect LSDV. Our assay provides high specificity and sensitivity with LOD of 1 \log_{10} copies per mL. Moreover, it is cost effective due to low cost of required reagents and easy fabrication. Also, samples don't require extraction or extensive pre-processing and results can be detected in a short time within 15 min. Besides, it can be easily used to detect a large number of samples simultaneously. All these, gives our developed test the edge over the other rapid diagnostic test. Further studies should be conducted to determine the other factors that could affect the accuracy of the proposed sensor.

4. Conclusion

LSDV-CQDs-MIP is a novel cost-effective paper-based turn-on fluorescence sensor designed for accurate detection of LSDV within few minutes. Our sensor exhibited high specificity and sensitivity for LSDV, with a lower LOD of detection reaching 10^1

\log_{10} TCID₅₀ per mL with high accuracy and reproducibility. Moreover, it detected LSDV in various real blood, skin nodules, and serum samples successfully, so it can be applied to clinical samples on a large scale and considered a local product. The findings of this study highlighted the approach for developing a novel, facile, and low-cost sensor that can be fabricated for efficient detection of other microorganisms.

Ethical statement

This article does not contain any studies with human participants or animals performed by any of the authors.

Data availability

All the data and materials are accessible on request.

Conflicts of interest

Authors declared no conflict of interest.

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