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Carbon dot as fluorescence sensor for glutathione in human serum samples: a review

Kawan F. Kayani,^{id}*^{abc} Sewara J. Mohammed,^{id}^{de} Dlzar Ghafoor,^{bf}
Mohammed K. Rahim^b and Harez Rashid Ahmed^{id}^b

In recent times, glutathione (GSH) has emerged as a crucial player in various cellular functions and is implicated in human pathologies. The creation of simple and highly responsive sensors capable of identifying GSH would be immensely valuable for gaining a deeper understanding of disease mechanisms. Carbon dots (CDs) have attracted considerable interest owing to their remarkable fluorescence capabilities, excellent compatibility with biological systems, and ease of production. Consequently, numerous research endeavors have focused on developing highly efficient CD probes for the rapid and precise detection of GSH. However, a comprehensive overview addressing the effective synthesis of CDs and their practical utility in GSH detection is still needed to further enhance the widespread application of CDs. In this context, we present, for the first time, a detailed introduction to GSH and CDs. Following this, we classify the synthetic methods of CDs. Additionally, considering various sensing categories, we classify CD fluorescent probes into single emission probes of CDs, ratiometric sensing probes of CDs, and visual detection of CDs. Furthermore, we highlight existing shortcomings and potential avenues for future research to offer valuable guidance in the preparation of commendable CDs and the detection of GSH.

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1. Introduction

GSH, the most abundant natural thiol and an essential anti-oxidant in biological systems, plays a central role in safeguarding against harmful free radicals and toxins.^{1–3} GSH is crucial for maintaining redox homeostasis by balancing its disulfide form.⁴ Typically, GSH is predominantly found in its reduced form under normal conditions. However, situations leading to oxidative stress in pathophysiology can elevate GSSG concentrations through the oxidation of reduced GSH to its oxidized form, glutathione disulfide (GSSG).⁵ Irregular GSH levels are closely linked to various diseases, including Alzheimer's disease, cancer, liver damage, diabetes, and arthritis.⁶ Furthermore, GSH, a tripeptide composed of γ -L-glutamyl-L-cysteinyl-glycine, is

renowned for its essential biological roles. Fig. 1 illustrates several critical cellular functions associated with it.⁷

Hence, it is essential to devise a straightforward, effective, and sensitive method for detecting and monitoring the levels of GSH in serum samples.⁸ Several techniques have been devised for GSH assays, including EC,^{9,10} HPLC,^{11,12} SERS,¹³ MRI,¹⁴ ECL,¹⁵ COL,¹⁶ and FL spectroscopy.^{17–19} Among these, FL spectroscopy is distinguished by its notable benefits, including heightened sensitivity, user-friendly operation, real-time assay capability, and nondestructive detection of targets in various systems.²⁰

However, various fluorescent materials have been employed for the determination of GSH, including metal-organic frameworks (MOFs), semiconductor quantum dots (QDs), nanoclusters (NCs), and small molecule fluorescent sensors. These materials, however, come with certain drawbacks. For instance, MOFs face poor solubility issues, nanoclusters exhibit short-term stability issues, QDs pose toxicity concerns related to their metals, and sensors based on small molecules have poor selectivity.^{21–27} Carbon dots have garnered significant attention in various fields over the last few years.^{28–31}

CDs represent a novel category of carbon nanomaterials endowed with fascinating optical and photophysical characteristics.^{32–35} The chemical characteristics of CDs, especially in the domains of analyte sensing,^{36–38} CDs stand out as preferred sensing probes due to their distinctive features,

^a Department of Chemistry, College of Science, Charmo University, 46023 Chamchamal/Sulaimani, Kurdistan Region, Iraq.

E-mail: kawan.nasraddinn@univsul.edu.iq

^b Department of Chemistry, College of Science, University of Sulaimani, Qliasan St, 46002, Sulaymaniyah City, Kurdistan Region, Iraq

^c Department of Pharmacy, Kurdistan Technical Institute, Sulaymaniyah City, Iraq

^d Anesthesia Department, College of Health Sciences, Cihan University Sulaimaniya, Sulaimaniya 46001, Kurdistan Region, Iraq

^e Research and Development Center, University of Sulaimani, Qlyasan Street, Kurdistan Regional Government, Sulaymaniyah, 46001, Iraq

^f College of Science, Department of Medical Laboratory Sciences, Komar University of Science and Technology, 46001, Sulaymania, Iraq





Fig. 1 An overview of GSH functions in the organism. Adapted from ref. 7.

including their small size, solubility, tunable photoluminescence, nonblinking fluorescence, and biocompatibility.^{39–46} Among these features, the photoluminescence of CDs stands out, demonstrating both excitation-dependent and independent emission, multicolor emission, photostability, and both up-conversion and down-conversion FL. These attributes can be attributed to surface energy traps, quantum confinement, and defects within the structure of CDs.^{47,48}

In 2004, CDs were fortuitously uncovered during the separation and purification of SWCNs.⁴⁹ Since this discovery, the straightforward synthesis and distinctive properties of CDs have led to a significant increase in the number of publications utilizing diverse precursors and synthesis methods. The methods are primarily categorized as either top-down or bottom-up approaches.

Top-down techniques involve the breaking and separation of carbon sources such as graphite, carbon fibers, and coal. Various methods, including oxidative cleavage, hydrothermal, microwave, electrochemical, and ultrasonic-assisted techniques, are utilized in top-down approaches. Conversely, bottom-up methods entail the pyrolysis of small organic molecules like citric acid, glucose, fructose, and ascorbic acid. These molecules can be combined with others to introduce heteroatom doping atoms.^{50–53}

Due to its ability to interact with fluorescent species, compounds, or nanomaterials, GSH can be quantitatively assessed by monitoring changes in fluorescent signals. Recent advancements in fluorescent GSH sensing can be categorized based on the origin of the output signals into three types: (1) signals emanating from fluorescent compounds, where GSH induces alterations in fluorescence intensity or emission wavelength through chemical reactions. (2) Signals originating from fluorescent indicators, where GSH triggers fluorescence activation by interrupting FRET between indicators and certain sensing nanoplatforms. (3) Signals arising from fluorescent nanomaterials, where GSH causes changes in fluorescence performance by perturbing surface properties or nanostructures. Among various fluorescent sensors, carbon dots have garnered significant attention as one of the most promising sensors for GSH due to their well-known nanomaterial properties with tunable fluorescence.⁵⁴ The fluorescent behavior of carbon dots is

attributed to factors such as conjugated π -domains of the carbon core, surface functional groups, incorporation of organic fluorophores, doping elements, *etc.* By modulating these factors, GSH can regulate the fluorescence of carbon dots, thereby enabling their straightforward application as fluorescent GSH sensors.

In this review, with a focus on providing a distinct perspective, we will emphasize recent advancements in the application of CDs for detecting GSH in serum samples. This includes discussions on the synthesis and sensing mechanisms of CDs. Practical applications for GSH sensing encompass single probes, ratiometric probes, and visual detection modes in real serum samples. Finally, we will briefly discuss the obstacles and potential advancements in the exploration and advancement of applications involving carbon dots.

2. Synthesis method for carbon dots

Regarding carbon sources, the creation of carbon dots can be broadly categorized into two methods: 'top-down' and 'bottom-up'.⁵⁵ In the 'top-down' approach, larger structures are broken down into smaller ones through various physical or chemical processes. This method employs techniques such as laser ablation, electrochemical processes, arc discharge, and plasma treatment for synthesis. These methods encompass template approaches, supported synthetic strategies, pyrolytic processes, microwave-based techniques, chemical oxidation, reverse micelle processes, and various other approaches for bottom-up synthesis. In the 'bottom-up' approach, smaller building blocks or molecules are assembled to create larger structures. Biomass materials are gaining preference in CD synthesis due to their renewable characteristics.^{56–58}

CDs offer advantages over other nanoparticles due to the availability of multiple synthesis methods, some of which are environmentally friendly.⁵⁹ The 'bottom-up' method is especially beneficial as it is more eco-friendly, less time-consuming, and allows for easy surface modification and composition adjustment of CDs. This stands in contrast to the 'top-down' method, which typically necessitates lengthier processing times, more severe reaction conditions, and the utilization of



costly materials and equipment. Synthetic conditions such as time, temperature, and voltage can be optimized for specific CD preparation.⁶⁰ Post-synthetic purification methods, ranging from simple treatments like centrifugation, dialysis, and filtration to more precise techniques like electrophoresis and chromatography, are employed to obtain purified and monodisperse CDs with independent properties.⁶¹ Additionally, surface functionalization during or after synthesis significantly influences the properties and applications of CDs.⁶²

2.1 Top-down approach

Top-down method involves the controlled fragmentation of larger carbon-containing structures, such as CNT or graphene, into smaller nanoparticles through physical or chemical processes, such as laser ablation, arc discharge, or plasma treatment.⁶³ This approach allows for precise manipulation at the nanoscale to create uniform carbon dots with tailored size, morphology, and surface properties.⁶⁴ By selectively breaking down larger carbon structures, the top-down method offers versatility and control over the final characteristics of the carbon dots, making it a valuable technique for nanomaterial synthesis.⁶⁵ To be specific, chemical oxidation emerges as a straightforward and convenient method for mass-producing high-quality CDs without the need for complex equipment.

This method includes breaking down precursor carbon materials like carbon fibers, graphene oxide, or CNT by employing strong acids or oxidizing agents.⁶⁶ During the chemical oxidation process, potent oxidants are employed to disrupt the covalent bonds within the carbon source, gradually converting it into smaller molecules. The effectiveness of the reaction is primarily influenced by the oxidizability of the oxidant employed.

Additionally, the oxidative process introduces an abundance of oxygen-containing functional groups, such as $-OH$ and $-COOH$, which contribute to excellent hydrophilicity and customizable emission properties in the CDs.⁶⁷ To create environmentally friendly GCDs, H_2SO_4 was chosen as an oxidant, and green tea leaf residue served as the raw material. Initially, the tea leaf residue underwent drying at $80\text{ }^\circ\text{C}$ and grinding into powder, followed by calcination at $350\text{ }^\circ\text{C}$ for 2 hours. Subsequently, the resulting residue was dissolved in concentrated H_2SO_4 and carbonized for 20 hours. Ultimately, GCDs were achieved.⁴⁶ The ultrasonic synthesis method achieves reactions through the thermal effect of cavitation and ultra-high-frequency vibration, offering unparalleled advantages such as eco-friendliness, cost-effectiveness, strong penetration, and uniform effects.⁶⁸

This technique involves the generation of alternating high-pressure waves and low-pressure waves during ultrasound processing, resulting in the formation and collapse of small bubbles within the liquid. Consequently, macroscopic carbon materials are fragmented into nanoscale carbon dots due to the intense hydrodynamic shear forces originating from the cavitation of these small bubbles.^{69,70} The laser ablation technique involves using a high-energy laser to irradiate the material, causing rapid heating. Laser pulses generate small pits in the

sample, consuming the material and producing plasma that flows through the sample surface, ultimately ionizing it,^{71,72} The arc discharge method is a specialized technique used to fabricate carbon nanoparticles, specifically carbon dots. In this method, two graphite electrodes are submerged in a liquid medium, typically water or organic solvents, and subjected to an electric discharge. This discharge generates intense heat and pressure, causing the graphite electrodes to undergo vaporization and fragmentation. Consequently, carbon nanoparticles, including carbon dots, are formed within the solution,^{73,74}

2.2 Bottom-up approach

The bottom-up method involves the assembly of smaller carbon-containing molecules or building blocks to form larger nanoparticles through self-assembly or controlled growth processes.⁷⁵ This method enables the creation of carbon dots with complex structures and functionalities by building from molecular precursors.⁷⁶ The CDs produced through this method typically exhibit abundant functional groups, resulting in a high fluorescence quantum yield. However, the particle size distribution is uneven, necessitating additional steps for separation and purification.⁷⁷ Synthetic conditions such as time, temperature, and voltage can be optimized for specific CD preparation. The bottom-up method starts with molecular-scale components and builds them up to create the desired nanomaterial through various techniques such as pyrolysis. The controlled assembly and growth of carbon dots from organic precursors are facilitated through the use of microwave-assisted synthesis and hydrothermal synthesis,^{78,79}

The hydrothermal/solvothermal method involves a synthesis technique where the reaction mixture interacts with water or organic substances and a non-aqueous solvent under defined temperature and autogenous pressure conditions within a sealed system, such as an autoclave.⁸⁰ This approach is considered direct and efficient, facilitating polymerization and carbonization reactions. It has found widespread application in preparing various materials owing to factors such as the high reactivity of the reactants, ease of solution control, minimal environmental impact, and low energy consumption under hydrothermal conditions.⁸¹ The diverse range of reaction substrates, reaction temperatures, and reaction times significantly alters the properties of the resulting products,^{82,83} Pyrolysis, a method employed in the synthesis of CDs, entails the controlled thermal decomposition of carbon-rich precursors under inert conditions.⁸⁴ Initially, suitable carbonaceous materials, such as organic compounds or biomass derivatives, are carefully selected.⁸⁵ Subsequently, these materials undergo gradual heating in an oxygen-free environment to avoid combustion.⁸⁶ As the temperature escalates, chemical bonds within the precursors disintegrate, liberating volatile organic compounds and yielding carbon-rich intermediates. These intermediates then undergo further molecular reorganization and condensation, ultimately culminating in the formation of carbon dots with precise structural attributes, including size, morphology, and surface chemistry,^{87,88} The microwave method is widely utilized



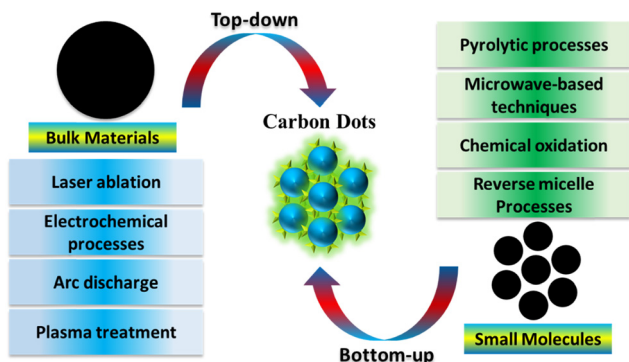


Fig. 2 Schematic diagram depicting the synthesis of CDs, incorporating both top-down and bottom-up approaches.

for directly carbonizing organic materials into carbon dots under microwave radiation, providing effectiveness and ease of operation, and minimal equipment requirements,^{89,90} The microwave method for synthesizing carbon dots involves using microwave irradiation to facilitate the conversion of carbon-rich precursors into carbon nanoparticles. Microwave treatment initiates the dehydration and pyrolysis of precursor substances, leading to their carbonization into small-sized CDs. As mentioned before, various methods have been devised for generating CDs for various applications, as illustrated in Fig. 2.

3. Sensing mechanisms

In theory, any alteration in fluorescence, be it in intensity, wavelength, anisotropy, or lifetime, linked to varying analyte

concentrations, holds the promise of being employed as a sensor.⁵¹

3.1 Static quenching

Fluorescence quenching describes a phenomenon that diminishes the overall intensity of fluorescent light. It is categorized into static quenching and dynamic quenching. In static quenching, a non-fluorescent complex is created between the fluorescent molecule and the quencher, leading to a reduction in the number of fluorescent molecules and a consequent decrease in fluorescence intensity. Notably, there is no observed change in fluorescence lifetime during the static quenching process,^{83,91,92} As shown in Fig. 3A.

3.2 Dynamic quenching

Dynamic quenching can be elucidated through energy transfer or charge transfer mechanisms. In the case of CDs, the excited state reverts to the ground state due to collisions with the quencher. For dynamic quenching to occur, the quencher must be in close proximity to the fluorescent molecule when it is in its excited state, initiating a reaction that accelerates the decay of the excited state. This leads to a shorter fluorescent lifetime while leaving the absorption spectrum unchanged.^{33,34} As shown in Fig. 3B.

3.3 Photoinduced electron transfer (PET)

PET is a deactivation mechanism characterized by an internal redox reaction occurring between the excited state of the fluorophore and another species capable of either donating or accepting an electron. In PET, an intricate complex is established between the electron donor and the electron

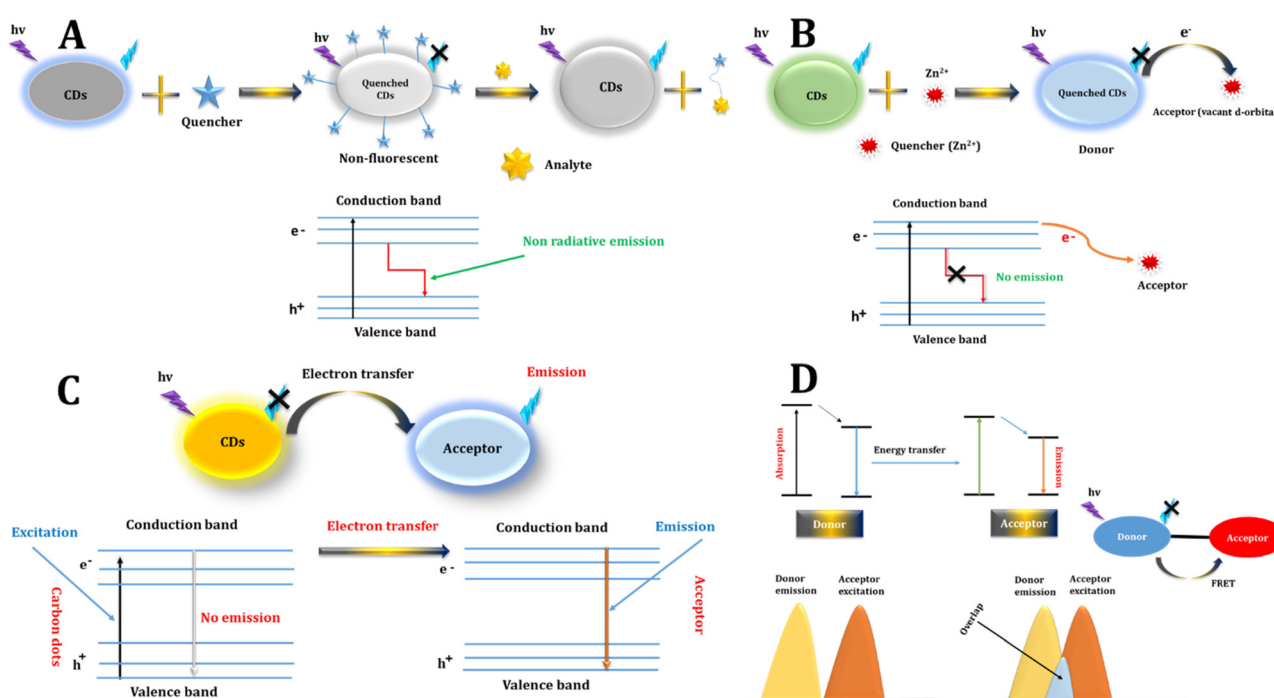


Fig. 3 Various FL sensing mechanisms include (A) static quenching, (B) dynamic quenching, (C) PET, and (D) FRET.



acceptor. This complex has the ability to revert to the ground state without emitting a photon, although in certain instances, exciplex emission may be detected. Ultimately, the additional electron present on the acceptor is restored to the electron donor to complete the process.^{31–33,35} As shown in Fig. 3C.

3.4 Photo-induced charge transfer (PCT)

This process entails the exchange of an electron between electron donor and acceptor functionalities to trigger FL. Partial charge transfer sensors involve the partial transfer of charges within a fully conjugated π system. The interaction between the donor and acceptor results in a modification of electron energy levels, leading to alterations in FL signals. PCT sensors are characterized by an integrated receptor and fluorophore, whereas PET sensors feature the separation of the electron donor moiety from the fluorophore by a spacer.⁵¹

3.5 Internal filter effect (IFE)

IFE is influenced by the spectral overlap between the fluorophore and absorber. Unlike FRET, IFE relies on radiative energy transfer, primarily affecting the donor's ground state through energy perturbation. Consequently, IFE results in only fluorescence quenching without any noticeable alteration in the fluorescence lifetime. Another notable difference from FRET is the absence of a distance requirement between the energy donor and absorber in IFE.^{32,36,37}

3.6 Aggregation induced emission (AIE)

AIE represents a luminescent phenomenon that stands in direct contrast to the Aggregation-Caused Quenching (ACQ) effect. In AIE materials, light emission does not occur in a sparse solution, yet strong fluorescence is evident at elevated concentrations or in a solid state. This behavior arises due to the dynamic intramolecular rotation of molecules in a diluted solution, which disperses their excited state energy (Table 1). However, in the aggregated state, intramolecular rotation becomes significantly constrained, resulting in the inhibition of nonradiative energy dissipation and the augmentation of FL.⁹³

3.7 Förster resonance energy transfer (FRET)

Resonance energy transfer is another valuable phenomenon for fluorescence sensors. In FRET, an initially excited molecule (donor) reverts to its ground state orbital, concurrently transferring energy to excite an electron on the acceptor.⁹⁴ This process involves the non-radiative energy transfer between donor and acceptor fluorophores, termed FRET pairs, occurring when the emission spectrum of CDs overlaps with the absorption spectrum of the quencher,^{92,95} FRET proceeds sans photon emission, owing to long-range dipole–dipole interactions between CDs and the quencher.⁹⁶ The efficiency of energy transfer hinges on factors such as the relative orientation of donor and acceptor dipoles, the degree of overlap between the fluorescence emission spectrum of the donor and the absorption spectrum of the acceptor, and the spatial separation between them. FRET manifests when the overlap between the emission spectrum of the donor and the absorption spectrum of the acceptor exceeds 30% and the distance is under 10 nm.⁹⁷ Through monitoring alterations in the fluorescence intensity or lifetime of the donor carbon dot, one can discern the presence or concentration of the acceptor molecule, thus enabling sensitive sensing applications. FRET with carbon dots garners preference due to their adaptable fluorescence properties, robust photostability, and biocompatibility, rendering them versatile tools across diverse sensing platforms.⁹⁸ As shown in Fig. 3D.

4. Application

Until this point, significant endeavors have been invested in creating diverse CDs to function as efficient fluorescence probes for detecting GSH. Due to their widespread sources, minimal toxicity, excellent photoluminescence, and biocompatibility,⁵⁰ CDs have garnered significant attention and found widespread use in chemical sensing applications. In this context, fluorescent CDs sensors are categorized into three types based on different sensing approaches: single probe of CDs, ratiometric probe of CDs, and visual detection of CDs.

Table 1 This table provides a comprehensive comparison of static quenching, dynamic quenching, PET, PCT, IEF, AIE, FRET in terms of their full forms, mechanisms, dependencies

Mechanisms	Full form	Mechanism	Dependency
Static quenching	Static quenching	Quencher forms a nonfluorescent complex with fluorophore, inhibiting fluorescence	Concentration of quencher, no dependence on time
Dynamic quenching	Dynamic quenching	Quencher transiently interacts with fluorophore, causing temporary decrease in fluorescence	Concentration of quencher, dynamic nature of interaction
PET	Photoinduced electron Transfer	Electron transfer from excited state of donor to acceptor	Extent of overlap between donor emission and acceptor absorption spectra
PCT	Photo-induced charge transfer	Charge transfer from donor to acceptor upon photoexcitation	Extent of overlap between donor emission and acceptor absorption spectra
IEF	Inner filter effect	Attenuation of fluorescence due to absorption by chromophores	Concentration and absorbance characteristics of chromophores
AIE	Aggregation induced emission	Emission enhancement upon aggregation of fluorophores	Aggregation state of fluorophores
FRET	Förster resonance energy transfer	Energy transfer between donor and acceptor fluorophores based on dipole–dipole interactions	Distance between donor and acceptor, spectral overlap, orientation



4.1 Single probe sensing

The fluorescence detection of CDs relies primarily on monitoring the variation in fluorescence intensity from a specific emission peak of an individual CD. This change occurs when exposed to a sole excitation wavelength, providing the outcome for target detection.

As an example, Zhang's team successfully developed a novel fluorescence strategy for detecting GSH. The designed probe consists of boron and nitrogen doped carbon dots (B,N-CDs) and MnO₂ nanosheets. In Fig. 4, it is evident that, with an increase in GSH concentration, the fluorescence of the CDs gradually intensifies. Furthermore, the determined LOD for GSH was 0.32 μM.⁹⁹

Khan's team¹⁰⁰ achieved the successful development of a novel carbon dots through hydrothermal synthesis, specifically creating a fluorescent CDs based sensor named GP-PEI-CDs. This sensor operates on an "on-off" FL strategy designed for detecting GSH. In Fig. 5, it is observed that the LOD for GSH was determined to be 38 nM, with a linear range spanning from 0 to 25 μM.

In this study, a novel NS-CQDs were successfully prepared. The fluorescence sensor exhibited quenching in the presence of Cu²⁺, and upon the introduction of GSH, the fluorescence sensor was restored. Moreover, this research effectively demonstrated the sensor's capability to selectively detect GSH. Specifically, the dynamic range for GSH spanned from 0.6–180.0 μM, with a noteworthy LOD of 100 nM, of significant importance, the NS-CQDs nanosensor exhibited reliable performance in monitoring GSH levels in serum.¹⁰⁰

Furthermore, Dong and colleagues utilized a hydrothermal method to prepare N,S-CQDs and gold nanoparticles. Through FRET, the resulting probe demonstrated high selectivity and a fast-responsive signal in GSH detection. The fluorescence probe for "turn-on" sensing of GSH, with a dynamic range spanning from 3.8–415.1 μM and a LOD of 210 nM. This sensor platform

was effectively employed for monitoring GSH in real samples, as illustrated in Fig. 6, yielding satisfactory results.¹⁰¹

Bano and colleagues effectively synthesized the N,S-CQD-MnO₂ nano-composite through a straightforward hydrothermal method. The detection capability of GSH in this system is noteworthy, as the MnO₂ nanosheet induces a significant reduction in the blue FL emission of N,S-CQDs. The reaction system also serves as an off-on nanoprobe for GSH detection. The presence of GSH in the system restores the fluorescence emission of N,S-CQDs by eliminating the FL resonance energy transfer. Additionally, the N,S-CQD-MnO₂ probe exhibits minimal cytotoxicity and excellent biocompatibility. The calculated LOD is determined to be 12 nM in a GSH dynamic range of 0.1–0.7 μM.¹⁰² As shown in Fig. 7.

In this section, we delve into more details about the most commonly utilized single sensing platforms based on CDs, as outlined in Table 2.

4.2 Ratiometric sensing

In 1993, Akkaya and colleagues introduced the concept of ratiometric sensors for detecting calcium in aqueous solutions.¹²³ Since then, there has been a growing trend in developing ratiometric sensors across various fields such as chemistry, biology, environmental studies, and food analysis. Typically, a ratiometric fluorescent sensor consists of both a response/indicator signal and a reference signal, creating a stable measurement system through self-calibration. The incorporation of an insensitive reference signal in ratiometric fluorescent measurements helps alleviate errors caused by interferences, while still capitalizing on the high selectivity and sensitivity of luminescence techniques for analyte detection.^{124,125} The fluorescence intensity of the indicator signal is divided by the intensity of a spectrally distinct reference signal, with the reference signal serving as a correction factor to enhance the reliability of the detection system. This

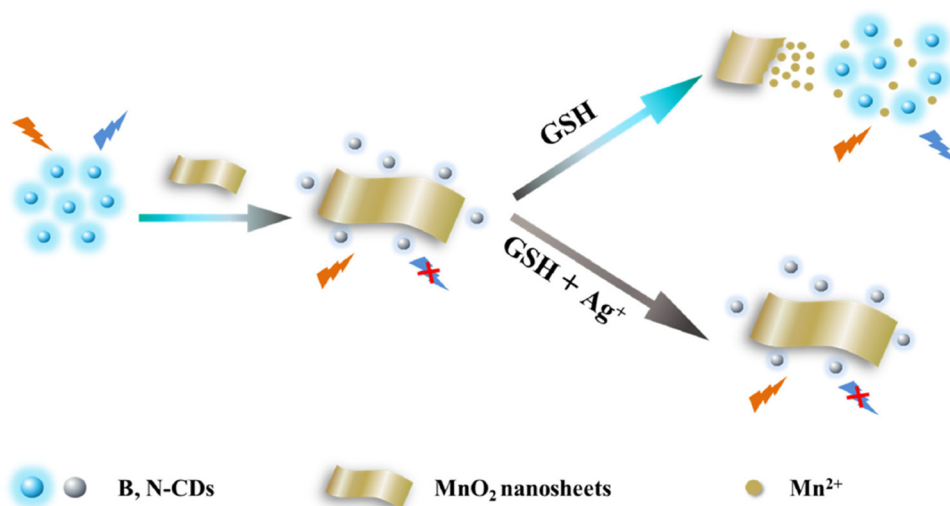


Fig. 4 Schematic illustration of the fluorometric sensing for GSH with permission. Copyright 2021, Elsevier.⁹⁹



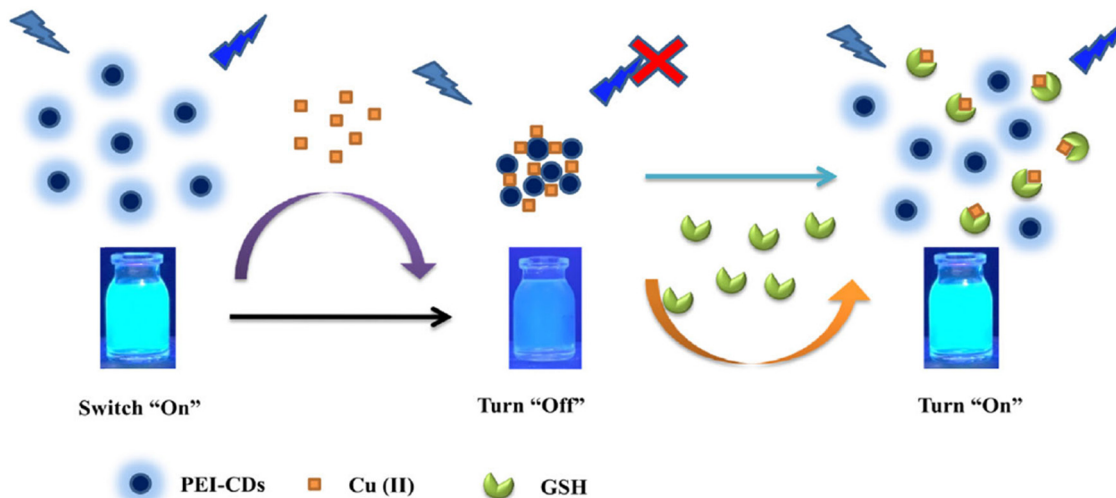


Fig. 5 A schematic illustration for GSH with permission. Copyright 2021, Springer Nature.¹⁰⁰

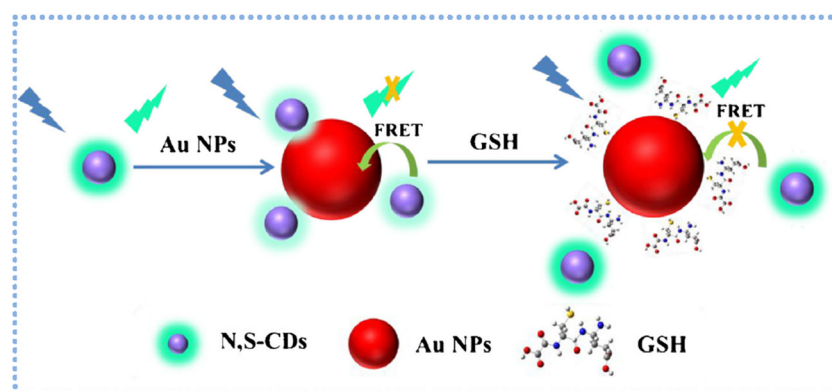


Fig. 6 Schematic principle for the determination of GSH based on FRET of N,S-CDs and Au NPs with permission. Copyright 2019, Springer Nature.¹⁰¹

approach facilitates accurate measurements that are less susceptible to variations in uncontrolled variables.¹²⁶ The construction of a novel ratiometric sensing method is promising and greatly significant for convenient detection of GSH in serum.

Han *et al.* reported ratiometric GSH probes based on FRET, as shown in Fig. 8.¹²⁷ The mechanism involved FRET between CQDs and 2,3-diaminophenazine (oxOPD). Moreover, the reaction involving the oxidation of Cu^{2+} and OPD could be hindered with the addition of GSH to the solution, thereby preventing the quenching of CQDs' fluorescence. The sensing system exhibited high sensitivity towards GSH within a range of 1.0–80.0 μM , with a corresponding LOD of 0.30 μM , respectively. Additionally, the proposed method could be applied to efficiently quantify GSH in samples.

In the conventional assay, CDs synthesized in a one-step process were employed as a ratiometric FL method for GSH detection. Varied concentrations of GSH (0–50 μM) were introduced into the CDs-RhB nanohybrid solution in the presence of 10 μM Hg^{2+} . Subsequently, Hg^{2+} was eliminated from the

surface of CDs, leading to a significant restoration of FL at 440 nm due to a competition mechanism as previously described. The FL intensity at 440 nm gradually recovered with increasing GSH concentrations, while the FL of RhB remained nearly constant. Consequently, a robust linear correlation between the I_{440}/I_{570} ratios and the GSH concentration in the range of 0–10 μM was evident. The estimated LOD was 20 nM.¹²⁸

Wang *et al.* also reported dual-emission carbon dots synthesized by one-pot hydrothermal method. Glutathione (GSH) can recover the fluorescence quenched by Hg^{2+} . Therefore, the sensing system showed high sensitivity toward GSH in a range of 1.0 to 10.0 μM , with a detection limit of 270 nM, respectively.¹²⁹

Alhazzani and colleagues also presented an innovative dual-emission ratiometric FL probe designed for detection of GSH. This was achieved by leveraging competitive interactions with a Ag-RF complex. The quantification of GSH levels without interference was enabled through the ratiometric measurement of FL intensities at 525 and 440 nm. This sensor exhibited



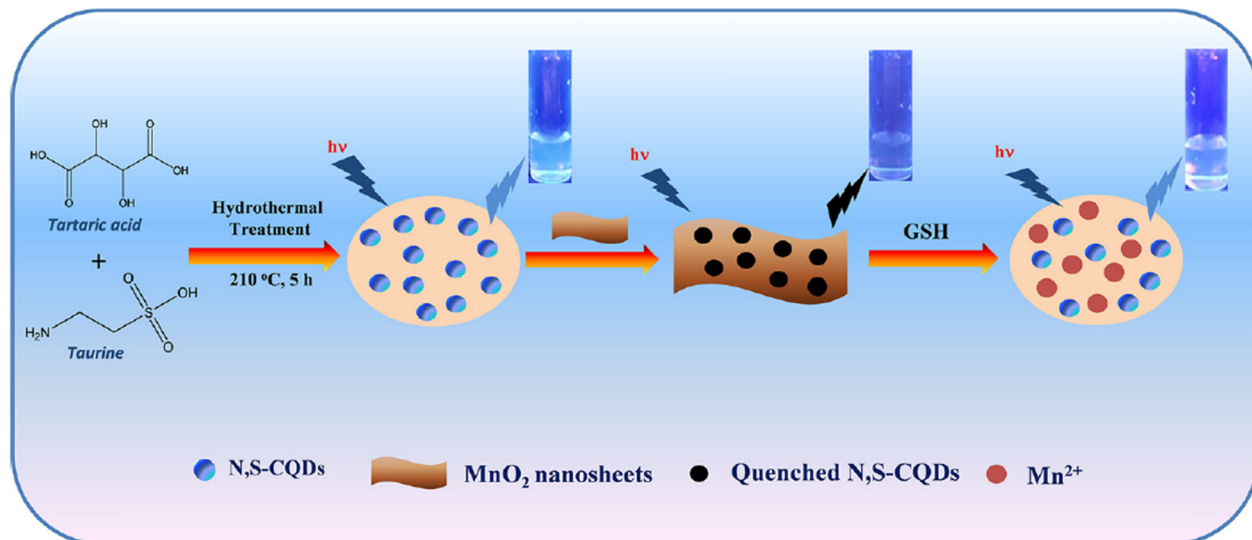


Fig. 7 Schematic representation of designed N,S-CQD-MnO₂ based nanoprobe for the 'off-on' detection of GSH with permission. Copyright 2019, Elsevier.¹⁰²

Table 2 List of selected CDs, precursors of preparation, emissive color, dynamic range, and LOD values

No.	Precursors of preparation	Color	Dynamic range (μM)	LOD (nM)	Ref.
1	Citric acid monohydrate, L-cysteine, and L-serine	Blue	0.01–5.0	3.6	103
3	Citric acid and ethanediamine	Blue	1–10	300	104
4	Tris(hydroxymethyl)-aminomethane and dimethylformamide	Blue, green, orange, and red	0.7–1.4, 0.4–1.1, 0.5–1.2, and 0.55–1.15	73 (b-CQDs), 88 (g-CQDs), 194 (o-CQDs), and 244 (r-CQDs)	105
5	Eggshell membrane	Blue	50–10	9.8	106
6	Citric acid and taurine	Blue	0–85	67	107
7	Grass carp	Blue	5206.3–117100	2505.53	108
8	Citric acid and triethylamine	Blue	0.02–0.3	7.52	109
9	Thylenediamine, polyethyleneimine or ethanolamine and DMF	Yellow	1–70	70	110
10	Starch soluble and citric acid	Green	0.5–100	350	111
11	(NH ₄) ₂ HPO ₄ and citric acid monohydrate	Blue	10–20	100	112
12	Citric acid anhydrous and 4,7,10-trioxa-1,13-tridecanediamine	Blue	30–400	460	113
13	Thiomalic acid and urea	Blue	0.1–4.0	59	114
14	Borax and boric acid	Blue	0.002–0.10	0.5	115
15	Citric acid	Blue	1.0–50	943	116
16	Uric acid	Blue	4–9	66	117
17	p-Hydroxybenzoic acid	Green	0.2–1000.0	30	118
18	Citric acid monohydrate and diammonium hydrogen phosphate	Blue	0.50–32	11.2	119
19	Citric acid and ethylenediamine	Blue	1000–100 $\times 10^3$	500 $\times 10^3$	120
20	para-Phenylenediamine	Red	12.5–800	700	121
21	Waste corncobs	Blue	50–150 $\times 10^3$	8.0133	122

excellent linearity over the range of 0.05–70 μM GSH with LOD of 15 nM. The sensor demonstrated precise analysis of GSH in serum samples with spiked concentrations (recoveries ranging from 96% to 98.5%), affirming its practical utility. Endowed with the benefits of simplicity, selectivity, and minimal sample volume needs, this fluorescent probe shows considerable potential as a tool for real-time, in-field monitoring of GSH, catering to both fundamental research and clinical investigations.¹³⁰

4.3 Visual detection and portable application

Visual detection is a simple method with observable results that can be easily distinguished by the naked eye. It offers several advantages, including the use of low-cost instruments,

rapid detection, user-friendly operation, minimal sample and reagent consumption, and suitability for on-site analysis,^{25,131} Scholars have been intrigued by visual detection for an extended period. The emergence of portable optical detection techniques for the qualitative and semi-quantitative analysis of various substances has piqued the interest of researchers across diverse fields. Visual detection enables the identification of analytes without the need for intricate external instrumentation.¹³² This is achieved by observing color changes during interactions with photoluminescence probes or sensors.¹³³ Consequently, the development of sensitive and selective techniques for visually detecting GSH in serum samples is of utmost importance.



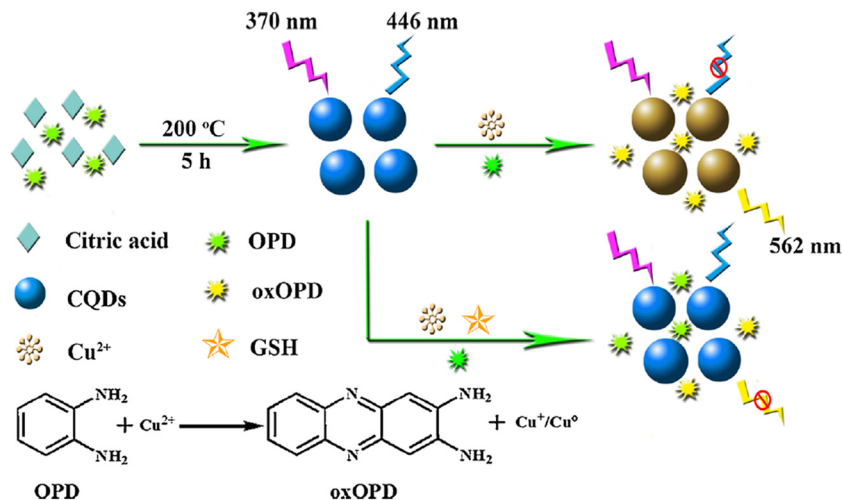


Fig. 8 Schematic illustration of ratiometric FL probe for GSH with permission. Copyright 2019, Elsevier.¹²⁷

A novel NCDs was synthesized using a solid-state method. The resulting NCDs in a water solution exhibit a highly selective “turn-off” FL response to Hg²⁺ compared to other interferences. Additionally, NCDs@Hg²⁺ demonstrates a remarkably selective “turn-on” FL response to the GSH as opposed to other amino acids, achieving a low LOD of 3.43×10^3 nM. Hydrogel films incorporating NCDs and NCDs@Hg²⁺ enable semi-quantitative detection of Hg²⁺ and GSH, respectively, observable to the naked eye under UV lamp illumination. This study validates the efficacy of NCDs as a practical material for the naked-eye detection of GSH in real-world samples.¹³⁴

Chu *et al.* introduced a portable sensing platform, comprising a ratiometric FL sensor integrated with a smartphone device, enabling sensitive, quantitative, and immediate determination of GSH levels in serum. The suppressed FL can rapidly recover upon interaction with GSH. Additionally, the red/blue values exhibited a close correlation with the GSH concentration. A strong linear relationship between the red/blue values and GSH concentration within the dynamic range

of 0–50 μ M was evident, as illustrated in Fig. 9. The estimated LOD was 1.84×10^3 nM.¹³⁵

He and his colleagues devised a system using developed CDs as a fluorophore and MnO₂ nanosheets as an absorber. The FL of CDs was effectively quenched by MnO₂ sheets through the IFE. Serving as the analyte, GSH could reduce MnO₂ sheets to Mn²⁺ ions, inhibiting the IFE and resulting in a fading of the solution color and the recovery of the FL signal. The results demonstrated that the proposed assay could visually distinguish 10 μ M GSH and quantitatively detect GSH within the dynamic range of 0.1–400 μ M. The LOD was 6.6 nM, evidenced by a color change from brown to colorless, facilitating the qualitative detection of GSH with the naked eye,¹¹² as depicted in Fig. 10.

However, in 2021, achieving exceptional selectivity and sensitivity became possible exclusively for GSH through a sensing platform based on PDA nanoparticles and r-CPDs using a ratiometric FL approach. In this sensing system, green-emitting PDA nanoparticles served as the response signal, while



Fig. 9 Schematic diagram of visual assay for GSH with permission. Copyright 2020, American Chemical Society.¹³⁵



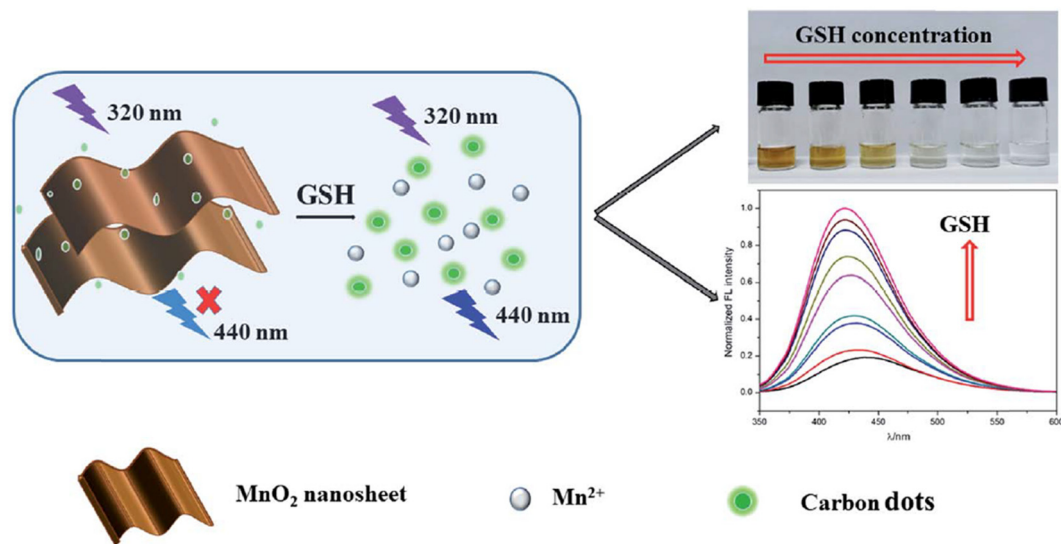


Fig. 10 Schematic illustration for GSH detection with permission. Copyright 2021, Royal Society of Chemistry.¹¹²

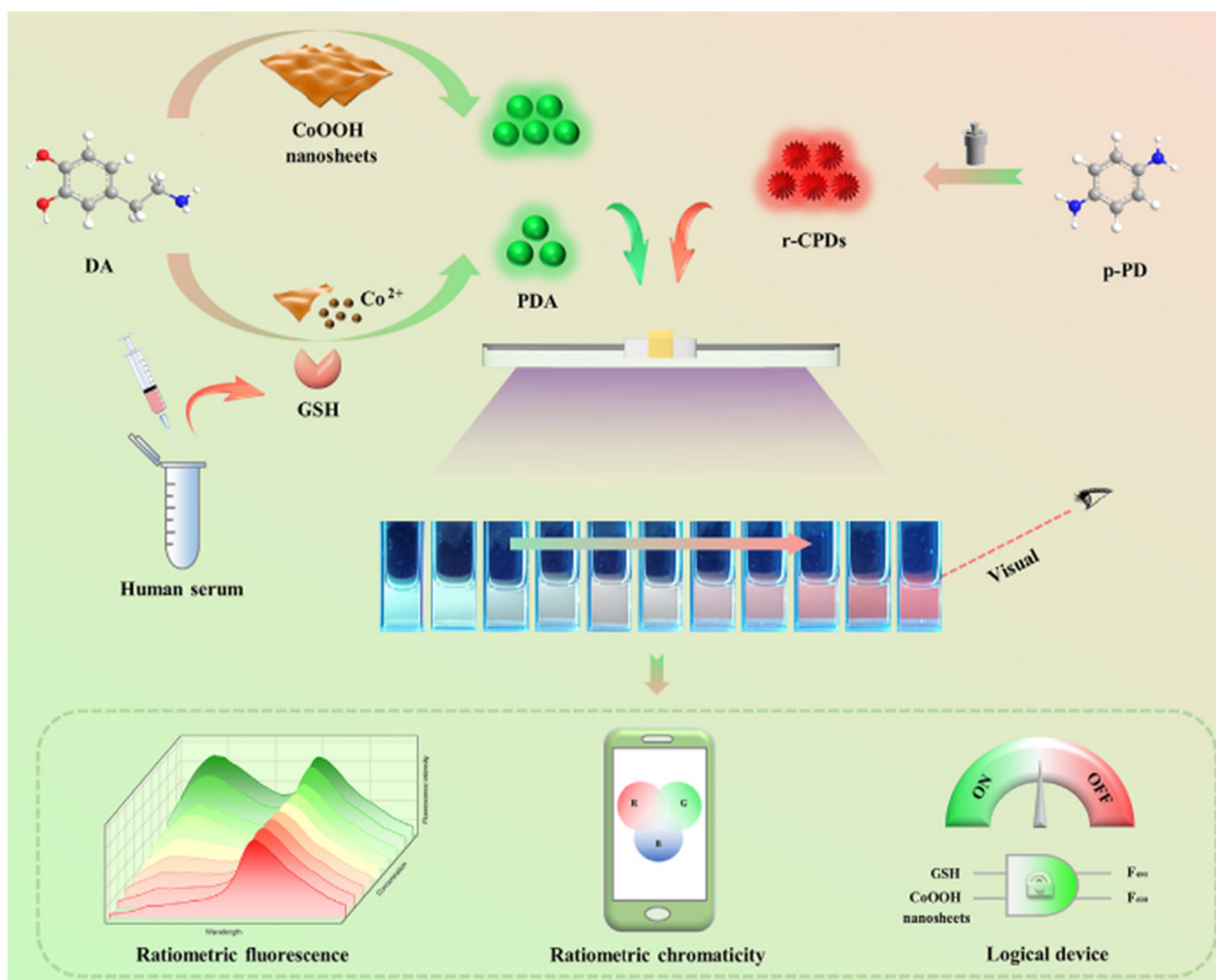


Fig. 11 Schematic diagram for the visual detection of GSH with permission. Copyright 2021, Elsevier.¹³⁶



r-CPDs functioned as the internal reference signal. Upon the addition of GSH, the green fluorescence of PDA nanoparticles decreased, while the red FL of the system remained relatively stable. This innovation established a detection threshold of 470 nM with a dynamic range of 1–100 μM .¹³⁶ These findings are visually depicted in Fig. 11.

5. Conclusions and prospects

Carbon dots, renowned for their outstanding PL and biocompatibility, have undergone extensive exploration as potential fluorescent probes for the swift and accurate detection of GSH. Consequently, as a result, we have gathered and emphasized reliable approaches for the effective synthesis of CDs and highlighted significant achievements in various sensing methods utilizing CDs for GSH detection. Despite notable progress, there are still several deficiencies and challenges that must be addressed.

The integration of diverse techniques renders CDs more adaptable in detection research. A significant step forward would involve combining CDs with FL immunoassay techniques, and molecular imprinting techniques to formulate efficient CDs with enhanced selectivity and sensitivity. Moreover, the practical utility of CDs should not be overlooked. Presently, the GSH detection using most CD probes remains confined to the laboratory settings. There is an urgent need to transition this technology for real-time detection and to devise simpler and faster detection regimens.

In summary, CD fluorescent probes have demonstrated promise in GSH detection; however, there is a need for continued efforts to explore innovative and dependable CDs to further improve detection efficiency. We expect that this review will stimulate increased interest to the preparation and optimization of CDs, with their widespread application in GSH detection flourishing in the near future.

List of abbreviations

CDs	Carbon dots
COL	Colorimetry
EC	Electrochemistry
HPLC	High performance liquid chromatography
SERS	Surface-enhanced Raman scattering
MRI	Magnetic resonance imaging
ECL	Electrochemiluminescence
FL	Fluorescence
SWCNs	Single-walled carbon nanotubes
NS-CQDs	Nitrogen and sulfur co-doped carbon quantum dots
Au NPs	Gold nanoparticles
FRET	Through Förster resonance energy transfer
NCDs	Nitrogen-doped carbon dots
PL	Photoluminescence
PDA	Polydopamine
r-CPDs	Red-carbonized polymer dots

Ag-RF	Silver-riboflavin
CNT	Carbon nanotube

Conflicts of interest

There are no conflicts to declare.

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