



# **Molybdenum Disulfide-Based Nanoprobes: Preparation and Sensing Application**

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**Abstract:** The use of nanoprobes in sensors is a popular way to amplify their analytical performance. Coupled with two-dimensional nanomaterials, nanoprobes have been widely used to construct fluorescence, electrochemical, electrochemiluminescence (ECL), colorimetric, surface enhanced Raman scattering (SERS) and surface plasmon resonance (SPR) sensors for target molecules' detection due to their extraordinary signal amplification effect. The MoS<sub>2</sub> nanosheet is an emerging layered nanomaterial with excellent chemical and physical properties, which has been considered as an ideal supporting substrate to design nanoprobes for the construction of sensors. Herein, the development and application of molybdenum disulfide (MoS<sub>2</sub>)-based nanoprobes is reviewed. First, the preparation principle of MoS<sub>2</sub>-based nanoprobes was introduced. Second, the sensing application of MoS<sub>2</sub>-based nanoprobes in future were discussed.

Keywords: molybdenum disulfide; nanoprobe; signal amplification; sensor; detection



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# 1. Introduction

As a powerful tool, a sensor has been employed to analyze chemical/biological molecules coupled with different detection methods, such as fluorescence, electrochemistry, electrochemiluminescence (ECL), colorimetry, surface enhanced Raman scattering (SERS) and surface plasmon resonance (SPR). To improve the analytical performance, many signal amplification strategies have been introduced into the construction of sensors, including DNA amplification technology, DNA walker, enzyme-assisted signal amplification and nanoprobes [1–5]. With the rapid development of nanomaterials, the nanoprobe has been considered as a promising signal amplification strategy to improve the performance of sensors.

Since gold nanoparticles (AuNPs) were introduced into the construction of nanoprobes [6,7], different kinds of nanomaterials have been extensively employed to construct nanoprobes due to their high surface area, excellent electrical and optical properties, high catalytic ability, excellent chemical stability and easy functionalization [8–12], such as noble metal nanoparticles [13,14], metal oxides [15], graphene and its derivative [16,17], transition metal dichalcogenides [18–20], and so on. The outstanding properties of nanomaterials allowed nanoprobes to easily load a large number of recognition and signal units, which can efficiently amplify the detection signal. Furthermore, the high biocompatibility of nanoprobes paves a way to analyze target molecules in vivo.

 $MoS_2$  is an emerging material star, which is a member of transition metal dichalcogenides. Due to its typical graphene-like layered nanostructure,  $MoS_2$  is also a potential candidate to construct the ideal nanoprobe due to its unique physical, chemical, and electronic properties, such as a large surface area, high conductivity, excellent quenching activity, accepted Raman enhancement effect and easy functionalization [21]. The recognition units or signal units assembled onto the MoS<sub>2</sub> nanosheet to form MoS<sub>2</sub>-based nanoprobes, which exhibited a high molecular recognition ability, excellent chemical stability, accepted biocompatibility and a strong signal amplification effect. Moreover, MoS<sub>2</sub>-based nanoprobes easily coupled with other signal amplification strategies to further amplify detection performances, including sensitivity, selectivity, reproducibility, stability, etc. Inspired by the rapid development of MoS<sub>2</sub>-based nanoprobes in sensing application, it is necessary to summarize its exciting advances (Figure 1). From this review, we hope to offer some useful suggestions to new researchers in the sensing field.

Application

# Molybdenum disulfide-based nanoprobes



**Figure 1.** Schematic diagram of preparation and sensing application of molybdenum disulfide-based nanoprobe.

# 2. Preparation of MoS<sub>2</sub>-Based Nanoprobes

Generally, a  $MoS_2$  nanosheet can load chemical/biological recognition units and signal molecules to form a nanoprobe via physical adsorption, chemical bond and noble metalmediated methods, respectively [22]. It should be noted that  $MoS_2$ -based nanoprobes prepared by different methods exhibited different advantages and disadvantages, which is listed in Table 1. According to the sensing application, the suitable nanoprobe coupled with analytical techniques often brings a better analytical performance, such as higher sensitivity, better selectivity and longer storage stability.

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Preparation Mechanism	Advantages	Disadvantages	References
physical interaction	simple, fast, facile, wide variety of binding molecules	unstable	[23–27]
chemical interaction	stable	The binding molecule needs to be modified, few choices of binding molecules	[28,29]
noble metal nanoparticles -mediated	simple, facile, stable, wide variety of binding molecules, properties enhanced	complicated preparation process	[30–36]

## 2.1. Physical Interaction

A MoS<sub>2</sub> nanosheet possesses a graphene-like layered nanostructure with a large surface area. As a result, it is easy to nonspecifically adsorb chemical or biological molecules via van der Waals force and electrostatic interactions. Notably, a MoS<sub>2</sub> nanosheet also exhibits different affinity towards single-strand (ss) and double-strand (ds) DNA. Based on these properties, MoS<sub>2</sub>-based nanoprobes including DNA-MoS<sub>2</sub>, aptamer-MoS<sub>2</sub> and peptide-MoS<sub>2</sub> probes, have been designed. For example, Zhu et al. firstly developed a fluorescence sensing platform by adsorbing DNA on the surface of a MoS<sub>2</sub> nanosheet as a nanoprobe [23]. A general platform for the construction of sensors was developed by combining the different affinity of the MoS<sub>2</sub> nanosheet towards ssDNA and dsDNA with its high fluorescence quenching efficiency. Five years later, Zhu and co-workers explored the possibility to construct MoS<sub>2</sub>-based fluorescence nanoprobes by adsorbing hairpin DNA [24]. Besides DNA, rhodamine B isothiocyanate (RhoBS) and antibodies also can be loaded on the surface of the MoS<sub>2</sub> nanosheet to form nanoprobes via physical adsorption and hydrophobic interactions, which can be used to determine silver ions and Escherichia coli by fluorescence and the SPR method, respectively [25,26].

#### 2.2. Chemical Interaction

Recognition and signal units assembled on the MoS<sub>2</sub> surface via chemical interaction is another efficient way to form MoS<sub>2</sub>-based nanoprobes. A popular method is to bind recognition and signal units with MoS<sub>2</sub> via classical thiol-metal coordination (typical Mo-S coordination). A typical example was given by Li et al., who designed a MoS<sub>2</sub>-based fluorescence nanoprobe for caspase-3 activity detection and images of cell apoptosis by efficiently conjugating two peptides with polydopamine-decorated MoS<sub>2</sub> nanosheets [28]. Since poly-cytosine (poly-C) DNA was proved as a high-affinity ligand for 2D nanomaterials [37], Xiao et al. [29] constructed a MoS<sub>2</sub>-based nanoprobe by assembling poly-C-modulated diblock molecular beacons on the MoS<sub>2</sub> surface. Experimental results suggested the length of poly-C could efficiently affect the analytical performance of the nanoprobe due to the regulation of the surface density [29].

#### 2.3. Noble Metal Nanoparticles-Mediated

As we know, noble metal nanoparticles have excellent advantages, including high catalytic activity, high electrical conductivity, large surface area and excellent biocompatibility, which have been widely used in sensing fields [38,39]. MoS<sub>2</sub> nanosheets have been proved as an ideal substrate to hybridize with noble metal nanoparticles [40,41]. As a result, the synergistic effect of noble metal nanoparticle-decorated MoS<sub>2</sub> nanocomposites brings faster electron transfer, higher catalytic activity, higher quenching efficiency and larger loading capacity, which have been considered as promising candidates to construct a nanoprobe. As a result, the designed nanoprobe not only retains the inherent characteristics of the hybrid element, but also brings better performance and enlarges its application fields. For instance, Su and co-worker prepared AuNP-decorated MoS<sub>2</sub> nanocomposites (MoS<sub>2</sub>-AuNPs) to construct electrochemical nanoprobes for biological molecules' detection with accepted results due to the signal amplification [30,32]. The recognition and signal units can efficiently co-immobilize on the MoS<sub>2</sub> surface via noble metal-mediated nanoparticles, such as an Au-S bond. Inspired by these exciting results, other noble metal nanoparticles were also successfully supported on the surface of molybdenum disulfide to construct a high-performance nanoprobe for sensing application [42–44].

#### 3. MoS<sub>2</sub>-Based Nanoprobes for Sensing Applications

MoS<sub>2</sub>-based nanoprobes can efficiently amplify the analytical performance due to their large loading amount, excellent electron transfer ability, high fluorescence quenching ability, and high Raman enhancement effect. As we know, different detection methods possess their inherent advantages and disadvantages (Table 2). Therefore, MoS<sub>2</sub>-based nanoprobes coupled with suitable analytical methods is a best way to construct sensors for obtaining

high-performance target molecules' detection. Herein, the recent progresses of MoS<sub>2</sub>-based nanoprobes coupled with electrochemical, ECL, colorimetric, SERS, fluorescence, and SPR methods is summarized (Table 3).

Detection Method	Advantages	Disadvantages	
fluorescence	easy design, simple, versatile, possible quantification	the need of large equipment, poor stability	
electrochemical	easy design, simple, fast, facile, quantification, miniaturization	complicated interface design, poor repeatability	
electrochemiluminescence	easy design, simple, fast, facile, quantification	complicated interface design, poor reproducibility	
colorimetric	simple, facile, no need of equipment	poor sensitivity, poor stability	
surface enhanced Raman scattering	fast, high sensitivity, high selectivity, quantification	poor reproducibility, the need of large equipment	
surface plasmon resonance	simple, high sensitivity	few application scenarios, the need of large equipment	

Table 2. Comparison of different detection methods.

## 3.1. Electrochemical Sensors

MoS<sub>2</sub>-based nanoprobe is a promising candidate to construct electrochemical sensors due to its high conductivity and high loaded capacity. To further improve the electronic properties of MoS<sub>2</sub>-based nanoprobes, the introduction of noble metal nanoparticles into nanoprobes has become a popular method. Therefore, gold nanoparticles (AuNPs), platinum nanoparticles (PtNPs), silver nanoparticles (AgNPs), and Au@AgPt nanocubes have been selected to form MoS<sub>2</sub>-based nanocomposites, which were further used to construct high-performance nanoprobes. For example, Su et al. used AuNPs-decorated MoS<sub>2</sub> nanocomposites to construct nanoprobes [32]. They utilized  $[Fe(CN)_6]^{3-/4-}$  and  $[Ru(NH_3)_6]^{3+}$  as signal molecules to design a dual-mode electrochemical sensor for microRNA-21 (miRNA-21) detection. As shown in Figure 2a, the  $MoS_2$ -based nanoprobes can efficiently amplify electrochemical responses by differential pulse voltammetry (DPV) and electrochemical impedance spectroscopy (EIS). Notably, the detection limit of this sensor obtained from EIS (0.45 fM) is lower than that obtained from DPV (0.78 fM), which is ascribed to the unique properties of 2D nanoprobes. This exciting finding opened a new way to construct electrochemical sensors. After three years, the same group developed a MoS<sub>2</sub>-based multilayer nanoprobe by using a DNA hybridization reaction (Figure 2b). Compared with a classical MoS<sub>2</sub>-based single-layer nanoprobe, the designed electrochemical sensor showed an ultrawide dynamic range (10 aM-1  $\mu$ M) and ultralow detection limit (38 aM) for miRNA-21 detection. The big structure of a MoS<sub>2</sub>-based multilayer nanoprobe and a large amount of negative DNA loaded on a multilayer nanoprobe both greatly hindered the electron transfer between  $[Fe(CN)_6]^{3-/4-}$  and the electrode surface, leading to the impedance value of this sensor obviously increasing with the addition of trace miRNA-21 [46]. To further amplify the detection performance, Bai's group coupled a MoS<sub>2</sub>-based nanoprobe with enzyme-assisted target recycling amplification to sensitively analyze the Sul1 gene. Due to the synergistic effect of two amplification strategies, the developed electrochemical sensor can determine 29.57 fM Sul1 gene with high selectivity [47]. Similarly, Ji et al. designed an electrochemical sensor for Pb<sup>2+</sup> analysis based on a MoS<sub>2</sub>-based nanoprobe and hemin/G-quadruplex DNAzyme [33]. The specificity of a DNAzyme combined with the high conductivity of MoS<sub>2</sub>-AuPt nanocomposites means this sensor has a lower detection limit for  $Pb^{2+}$  analysis (38 fg mL<sup>-1</sup>).

Method	Nanoprobe	Target	Linear Range	LOD	References
	MoS <sub>2</sub> -AuPt	Pb <sup>2+</sup>	$0.1 \text{ pg mL}^{-1} -1000 \text{ ng mL}^{-1}$	$38 \text{ fg mL}^{-1}$	[33]
	hemin/G-quadruplex-Tb- PdNPs/PDDA-G-MoS <sub>2</sub>	thrombin	0.0001-40 nM	0.062 pM	[45]
Electrochemistry	MoS <sub>2</sub> -AuNP MoS <sub>2</sub> -PANI-Au Au@Pd/MoS <sub>2</sub> @MWCNTs	microRNA-21 Sul1 HBeAg	10 aM–1 μM 40 fM–40 nM 0.1–500 pg mL <sup>-1</sup>	38 aM 29.57 fM 26 fg mL <sup>-1</sup>	[46] [47] [48]
	MoS <sub>2</sub> NFs/Au@AgPt YNCs	CEA	$10 \text{ fg mL}^{-1} -100 \text{ ng mL}^{-1}$	$3.09 \text{ fg mL}^{-1}$	[49]
	DPCN/MoS <sub>2</sub>	CTnI	$10 \text{ fg mL}^{-1} -100 \text{ ng mL}^{-1}$	$3.02 \text{ fg mL}^{-1}$	[50]
	MoS <sub>2</sub> @Cu <sub>2</sub> O-Au	AFP	$0.1 \text{ pg mL}^{-1} -50 \text{ ng mL}^{-1}$	$0.037 \text{ pg mL}^{-1}$	[51]
	ABEI-Ag-MoS <sub>2</sub> NFs/HP3	MUC1	$1 \text{ fg mL}^{-1} -10 \text{ ng mL}^{-1}$	$0.58~\mathrm{fg}~\mathrm{mL}^{-1}$	[52]
	MoS <sub>2</sub> @Au	Siglec-5	10–500 pM	8.9 pM	[53]
ECL	MoS <sub>2</sub> NF	concanavalin A	$1.0~{ m pg~mL^{-1}} - 100~{ m ng~mL^{-1}}$	$0.3 \text{ pg mL}^{-1}$	[54]
	MIL-101@Au -MoS <sub>2</sub> QDs MoS <sub>2</sub>	β-amyloid	$10^{-5} - 50 \ \mathrm{ng} \ \mathrm{mL}^{-1}$	$3.32~\mathrm{fg}~\mathrm{mL}^{-1}$	[55]
		CA19-9	$0.002-50 \text{ Um}L^{-1}$	$0.25 \ {\rm mU} \ {\rm mL}^{-1}$	[56]
	MoS <sub>2</sub> NSs	epididymis-specific protein 4	$10^{-6} - 10 \text{ ng mL}^{-1}$	$3 imes 10^{-7}~ng~mL^{-1}$	[57]
Colorimetry	MoS <sub>2</sub> -AuNPs Fe-doped MoS <sub>2</sub> MoS <sub>2</sub> @CNNS MoS <sub>2</sub> /GO MoS <sub>2</sub> -polypyrrole-Pd csDNA-Au-MoS <sub>2</sub>	CEA glutathione $H_2O_2$ glucose l-cysteine $Cd^{2+}$	0.005-10 ng mL <sup>-1</sup> 1-30 μM 2.0-50.0 μM 1-50 μM 1-10 μM 1-500 ng mL <sup>-1</sup>	0.5 pg mL <sup>-1</sup> 0.577 μM 0.02 μM 0.83 μM 0.08 μM 0.7 ng mL <sup>-1</sup>	[30] [34] [58] [59] [60] [61]
	TP/SYL3C-MoS <sub>2</sub>	circulating tumor cells	$5-10^4$ cells mL <sup>-1</sup>	2 cells mL <sup><math>-1</math></sup>	[62]
	MoS <sub>2</sub> /C-Au	$H_2O_2$ in living cells	$1 \times 10^{-5}$ – $2 \times 10^{-4}$ M	1.82 µM	[63]
SERS	R6G-tagged MoS <sub>2</sub> NF MoS <sub>2</sub> NFs@AuNPs/MBA Au NP@MoS <sub>2</sub>	CA19-9 CEA cell imaging	$\begin{array}{c} 5\times10^{-3}{-100}{\rm IU}{\rm mL}^{-1}\\ 0.0001{-}100.0{\rm ng}{\rm mL}^{-1}\\ -{-}\end{array}$	$\begin{array}{c} 3.43 \times 10^{-4} \ \text{IU} \ \text{mL}^{-1} \\ 0.033 \ \text{pg} \ \text{mL}^{-1} \\ \end{array}$	[64] [65] [66]
	MoS <sub>2</sub> -loaded MBs MoS <sub>2</sub> NSs MoS <sub>2</sub>	microRNA caspase-3 EpCAM	1 pM-10 nM 2-360 ng mL <sup>-1</sup> 3-54 nM	10 fM 0.33 ng mL <sup>-1</sup> 450 pM	[24] [28] [67]
Fluorescence	MoS <sub>2</sub>	PSA	$0-60 \text{ ng mL}^{-1}$	$0.2 \text{ ng mL}^{-1}$	[68]
1 Morescence	$MoS_2$	streptavidin	$0-600 \text{ ng mL}^{-1}$	$0.67 \text{ ng mL}^{-1}$	[69]
	DOX-SH/M-MoS <sub>2</sub> ND	glutathione cellular imaging	$0.1 \times 10^{-3}$ -100 × 10 <sup>-3</sup> M $0.1 \times 10^{-3}$ -4 × 10 <sup>-3</sup> M	$30  imes 10^{-9} \ \text{M}$	[70]
	$MoS_2$	ATP	0.067–26.7 μM	34.4 nM	[71]
	MoS <sub>2</sub> -NFP	programed cell death protein 1	$125-8000 \text{ pg mL}^{-1}$	$85.5 \text{ pg mL}^{-1}$	[72]
SPR	AuNPs-MoS <sub>2</sub>	miRNA-141	1–50 pM	0.5 fM	[73]

Table 3. MoS<sub>2</sub>-based nanoprobes for sensing applications.

Abbreviation: toluidine blue (Tb), poly (diallyldimethylammonium chloride) (PDDA), graphene (G), polyaniline (PANI), gold@palladium nanoparticles (Au@Pt), multiwalled carbon nanotubes (MWCNTs), hepatitis B e antigen (HBeAg), trimetallic yolk-shell Au@AgPt nanocubes (Au@AgPt YNCs), carcinoembryonic antigen (CEA), dendritic platinum-copper alloy nanoparticles (DPCN), cardiac troponin I (CTnI), alpha fetoprotein (AFP), N-(aminobutyl)-N-(ethylisoluminol) (ABEI), mucin 1 (MUC1), sialic acid-binding immunoglobulin (Ig)-like lectin 5 (Siglec-5), Materials Institute Lavoisier-101 (MIL-101), concanavalin A (ConA), quantum dots (QDs), carbohydrate antigen 19-9 (CA19-9), g-C3N4 nanosheets (CNNS), polypyrrole (PPy), thymolphthalein (TP), Rhodamine 6G (R6G), 4-mercaptobenzoic acid (MBA), molecular beacons (MB), epithelial cell adhesion molecule (EpCAM), prostate specific antigen (PSA), thiolated doxorubicin (DOX-SH), adenosine triphosphate (ATP), MoS2 modified nanofiber paper (MoS2-NFP).

A MoS<sub>2</sub>-based nanoprobe has been also employed to construct electrochemical immunosensors. For example, Li et al. constructed an immunosensor by using CeO<sub>2</sub>-MoS<sub>2</sub>-Pb<sup>2+</sup>-Ab<sub>2</sub> as a signal probe [36]. Ingeniously, Pb<sup>2+</sup> can adsorbs and aggregates on the surface of a CeO<sub>2</sub>-MoS<sub>2</sub> nanocomposite, which can not only anchor antibodies, but also generate and enhance electrical signals. This novel design of a MoS<sub>2</sub>-based nanoprobe achieved the purpose of the sensitive detection of CEA. To further improve the analytical performance, Su et al. [31] constructed an enzyme-assisted signal amplification strategy for carcinoembryonic antigen (CEA) analysis by taking the advantages of MoS<sub>2</sub>-AuNPs nanocomposites and the catalytic activity of enzymes (Figure 2c). In this work, MoS<sub>2</sub>-AuNPs can not only accelerate electron transfer due to its high conductivity, but also can load a large number of enzymes and antibodies to achieve multiple signal amplification. Therefore, the proposed immunosensor detected down to 1.2 fg mL<sup>-1</sup> CEA with high selectivity and good stability. Similarly, Gao et al. developed a signal probe by combining gold@palladium nanoparticle-loaded molybdenum disulfide with multi-walled carbon nanotubes (Au@Pd/MoS<sub>2</sub>@MWCNTs) to efficiently analyze the hepatitis B e antigen (HBeAg) [48]. With the addition of HBeAg, a classical sandwich immunosensor was formed (Figure 2d). The introduced signal probe contained Au@Pd nanoparticles, which can efficiently catalyze hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) to generate high electrochemical signal. Therefore, the sensor got a low detection limit of 26 fg mL<sup>-1</sup> with the help of signal probe amplification. Other MoS<sub>2</sub>-based electrochemical nanoprobes were also used to detect cardiac troponin I, HBsAg, and CEA due to their outstanding signal amplification effect, respectively [49,50,74].

#### 3.2. ECL Sensors

A few layers of MoS<sub>2</sub> the nanosheet possess a direct bandgap and a large surface. These properties made the MoS<sub>2</sub>-based nanoprobe a potential candidate to construct electrochemiluminescence (ECL) sensors. Usually, a MoS<sub>2</sub>-based nanoprobe is used as a co-reaction promoter to efficiently amplify the detection signal, called a "signal-on" detection mechanism. An example was offered by Li et al., who constructed a ECL sensor for mucin 1 (MUC1) analysis by coupling a target recycling signal amplification strategy and a MoS<sub>2</sub>-based nanoprobe [52]. The prepared MoS<sub>2</sub> nanoflowers can heavily load N-(aminobutyl)-N-(ethylisoluminol) (ABEI)-decorated AgNPs as signal amplifiers, which can catalyze ABEI-H<sub>2</sub>O<sub>2</sub> to improve the detection intensity. As shown in Figure 3a, the added MUC1 triggered the signal amplification process, leading to the designed ECL aptasensor having a wide linear range (1 fg mL<sup>-1</sup> to 10 ng mL<sup>-1</sup>) and low detection limit (0.58 fg mL<sup>-1</sup>) for MUC1 determination. Another ECL MoS<sub>2</sub>-based nanoprobe was constructed by MoS<sub>2</sub>@Au nanocomposites [53]. With the assistance of exonuclease III-driven DNA walker, a sensitive ECL sensor was developed for 8.9 pM sialic acid-binding immunoglobulin (Ig)-like lectin 5 analysis.

A MoS<sub>2</sub>-based nanoprobe was also used to construct "signal off" ECL sensors by utilizing the high quenching ability of MoS<sub>2</sub> nanostructures. For example, Yuan and coworker reported a ECL sensor for concanavalin A (Con A) determination according to the signal-off sensing mechanism [54]. The as-prepared MoS<sub>2</sub> nanoflowers highly quenched the ECL signal of the Ru complex, making the ECL response decrease with the increasing ConA concentration, ranging from 1.0 pg mL<sup>-1</sup>–100 ng mL<sup>-1</sup> (Figure 3b). According to the quenching properties of MoS<sub>2</sub>-based nanoprobes in ECL sensing application, several ECL sensors were constructed for beta-amyloid (A $\beta$ ), CA19-9 antigen and human epididymal specific protein 4 detection, respectively [55–57]. All experimental data suggested the introduction of MoS<sub>2</sub>-based nanoprobes can efficiently improve the analytical performances, such as linear range, detection limit, analytical time, etc.

#### 3.3. Colorimetric Sensors

Previous works proved that  $MoS_2$  nanostructures have peroxidase mimicking activity with high chemical and thermal stability [74]. For example, Zhao et al. found that sodium dodecyl sulfate-conjugated  $MoS_2$  nanoparticles (SDS- $MoS_2$  NPs) can efficiently catalyze a 3,3,5,5-tetramethylbenzidine (TMB) and hydrogen peroxide ( $H_2O_2$ ) reaction strategy, exhibiting peroxidase-like activity for the detection of glucose [75]. To improve the peroxidase-like activity of  $MoS_2$  nanostructures, the formation of  $MoS_2$ -based nanocomposites is a universal method. These nanocomposites offer the opportunity to develop high-performance colorimetric nanoprobes due to their better catalytic activity, such as MoS<sub>2</sub>-carbon nanotubes [76], MoS<sub>2</sub>-g-C<sub>3</sub>N<sub>4</sub> [58], MoS<sub>2</sub>-graphene oxide [59], MoS<sub>2</sub>-Au@Pt [77], etc. According to this concept, Peng et al. used a MoS<sub>2</sub>-graphene oxide (MoS<sub>2</sub>-GO) nanocomposite instead of a biological enzyme to colorimetricly detect H<sub>2</sub>O<sub>2</sub> and glucose [59]. The synergistic effect of MoS<sub>2</sub> and graphene oxide made this designed colorimetric sensor analyze H<sub>2</sub>O<sub>2</sub> and glucose in serum samples by the naked-eye (Figure 4a). Compared with graphene, noble metal nanostructures hybridized with a MoS<sub>2</sub> nanosheet can bring outstanding peroxidase-like activity. A typical example was offered by Su and co-workers, who designed a colorimetric sensor for cysteine analysis based on a MoS<sub>2</sub>-Au@Pt nanoprobe [77]. The enzyme-mimicking activity made this sensor show a wide linear range and low detection limit for cysteine detection. Moreover, this colorimetric sensor can determine cysteine in medical tables. Similarly, Singh et al. utilized the highly-efficient peroxidase-like activity of Fe-doped MoS<sub>2</sub> nanomaterials to colorimetricly detect glutathione in buffer and human serum [34]. The satisfactory results further proved the excellent application of MoS<sub>2</sub>-based nanoprobes in the colorimetric sensing field.



**Figure 2.** (a) Construction of dual-mode electrochemical sensor for miRNA-21 detection based on MoS<sub>2</sub>-based nanoprobes. Reprinted with permission from [32]. Copyright 2017, Elsevier. (b) Construction of multilayer MoS<sub>2</sub>-based nanoprobes for miRNA-21 analysis. Reprinted with permission from [46]. Copyright 2020, Royal Society of Chemistry. (c) Illustration of electrochemical immunosensor by coupling MoS<sub>2</sub>-based nanoprobe with triple signal amplification. Reprinted with permission from [31]. Copyright 2019, Elsevier. (d) Construction and application of MoS<sub>2</sub>-based nanoprobes for electrochemical analysis of HBeAg. Reprinted with permission from [48]. Copyright 2017, Elsevier.



**Figure 3.** (a) Development of ECL biosensor for mucin 1 analysis based on MoS<sub>2</sub>-based nanoprobe. Reprinted with permission from [52]. Copyright 2018, American Chemical Society. (b) Illustration of ECL biosensor for concanavalin detection A by using the high-efficient quenching ability of MoS<sub>2</sub> nanoflower. Reprinted with permission from [54]. Copyright 2017, Elsevier.



**Figure 4.** (a) Colorimetric analysis of glucose by coupling peroxidase-like MoS<sub>2</sub>-based nanoprobe and glucose oxidase. Reprinted with permission from [59]. Copyright 2016, Elsevier. (b) Construction of a MoS<sub>2</sub>-based colorimetric biosensor for carcinoembryonic antigen analysis. Reprinted with permission from [30]. Copyright 2018, American Chemical Society.

Besides peroxidase-like activity, another reason for  $MoS_2$ -based nanocomposites in the colorimetric sensing application is the high catalytic activity. By utilizing this property, Su et al. constructed a colorimetric nanoprobe by assembling an anti-CEA on the surface of a  $MoS_2$ -AuNPs nanocomposite [30]. The assembled amount of anti-CEA greatly influenced the catalytic activity of the  $MoS_2$ -AuNPs nanocomposite, which can be used to recognize and detect CEA by catalyzing the reaction of 4-nitrophenol (4-NP) and sodium borohydride (NaBH<sub>4</sub>). Corresponding with the solution color and adsorption intensity, the developed can analyze 5 pg mL<sup>-1</sup>–10 ng mL<sup>-1</sup> of CEA with high selectivity (Figure 4b). This potential colorimetric sensing application inspired more researchers to synthesize different kinds of  $MoS_2$ -based nanocomposites with high catalytic activity, such as AuNP or PtNP decorated Ni promoted  $MoS_2$  nanocomposites [78], multi-element nanocomposites composed by noble metal nanoparticles, polyaniline microtubes, and Fe<sub>3</sub>O<sub>4</sub> and MoS<sub>2</sub> nanosheets [79].

# 3.4. SERS Sensors

As a graphene-like 2D layered nanomaterial, a MoS<sub>2</sub> nanosheet also exhibits an excellent Raman enhancement effect due to the chemical enhancement mechanism [80]. Decoration with noble metal nanoparticles, the synergistic effect of chemical enhancement and electromagnetic enhancement makes the MoS<sub>2</sub>-noble metal nanoparticles' nanohybrids exhibit a better Raman enhancement effect. Therefore, MoS<sub>2</sub> and its nanocomposites are often employed as SERS-active substrates to construct sensors for target molecules' detection [81,82]. Besides SERS-active substrates, MoS<sub>2</sub>-based nanohybrids have also been used to construct nanoprobes for sensing application. For example, Jiang et al. [64] developed a MoS<sub>2</sub>-based immunosensor for the carbohydrate antigen 19-9's (CA19-9) detection by using a  $MoS_2$  nanosheet as a SERS-active substrate and a  $MoS_2$  nanoflower as a SERS tag (Figure 5). Expectedly, this sandwich design exhibited a desirable enhancement effect on CA19-9 analysis, resulting in a wide linear range  $(5 \times 10^{-4} - 1 \times 10^{2} \text{ IU} \cdot \text{mL}^{-1})$  and low detection limit ( $3.43 \times 10^{-4} \text{ IU} \cdot \text{mL}^{-1}$ ). More meaningfully, this designed immunosensor showed accepted results for CA19-9 detection in clinical patient serum samples, which was in agreement with the conventional chemiluminescent immunoassay. Similarly, Medetalibeyoglu et al. also reported a sandwich-type immunosensor for CEA detection by using 4-mercaptobenzoic acid assembled AuNPs-decorated MoS2 nanoflowers (MoS2 NFs@Au NPs/MBA) as SERS tag [65]. Coupled with Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> MXene-based SERS-active substrate, this immunosensor detected as low as 0.033 pg mL<sup>-1</sup> of CEA, with high selectivity, stability and repeatability. More interestingly, a MoS<sub>2</sub>-based SERS nanoprobe is also a powerful tool for label-free SERS imaging. For example, Fei et al. offered an example of a MoS<sub>2</sub>-based nanoprobe for SERS imaging in living 4T1 cells [66]. Experimental results suggested that a MoS<sub>2</sub>-based nanoprobe may be the promising alternative because of its intrinsic vibrational bands in the Raman-silence region of cells.



**Figure 5.** Illustration of MoS<sub>2</sub>-based immunosensor for CA19-9 detection. Reprinted with permission from [64]. Copyright 2021, Elsevier.

#### 3.5. Fluorescence Sensors

The tunable layer thickness of the MoS<sub>2</sub> nanosheet leads to its indirect to direct bandgap transition, which generates excellent optical properties. Especially, the outstanding quenching ability towards organic dyes suggests that a MoS<sub>2</sub> nanosheet can be employed as a nanoquencher to construct fluorescence sensors. Zhu et al. had given a first example of a fluorescence sensor for targetting DNA and other small molecules by using a MoS<sub>2</sub> nanosheet as a sensing probe [23]. The different affinity of the MoS<sub>2</sub> nanosheet towards ssDNA and dsDNA makes the labeled 5-carboxyfluorescein (FAM) close to or far from the surface of the MoS<sub>2</sub> nanosheet, resulting in the fluorescence signal recovering with the formation of dsDNA (Figure 6a). This exciting finding inspired more and more researchers to develop fluorescence sensors for target molecules' detection by using MoS<sub>2</sub>-based sensing nanoprobes. A typical design is coupling an aptamer with a MoS<sub>2</sub>-based nanoprobe to analyze nucleic acids, proteins, thrombin, metal ions, kanamycin, ochratoxin A, and so on [67,83–85]. For example, Kong et al. utilized the high-efficient quenching ability of a MoS<sub>2</sub> nanosheet to develop a fluorescence sensor for prostate specific antigen (PSA) analysis [68]. The structure of the aptamer was changed with the recognition of the PSA, leading to the aptamer-PSA product releasing from the MoS<sub>2</sub> nanosheet and the fluorescence recovering. Under optimal conditions, this designed sensor can detect as low as  $0.2 \text{ ng mL}^{-1}$  of PSA with high selectivity.

To further improve the analytical performance, several signal amplification strategies coupled with MoS<sub>2</sub>-based nanoprobes were introduced into the construction of fluorescent sensors. For example, Xiang et al. reported a fluorescence sensor for streptavidin (SA) detection by coupling exonuclease III (Exo III)-assisted DNA recycling amplification with MoS<sub>2</sub>-based nanoprobes [69]. As shown in Figure 6b, probe 1 was not degraded by Exo III because of the binding of SA and biotin. Subsequently, the protected probe 1 hybridized with probe 2, which can be digested by Exo III. The continually released FAM led to a strong fluorescence signal due to the signal amplification, producing a low detection limit of 0.67 ng mL $^{-1}$  for SA detection. Similarly, Xiao et al. combined duplex-specific nuclease (DSN)-mediated signal amplification with MoS<sub>2</sub>-based nanoprobes to develop a fluorescence for microRNA (miRNA) detection [24]. In the presence of miRNA, molecular beacons adsorbed onto the MoS<sub>2</sub> nanosheet changed to DNA-RNA heteroduplexes and were released from the  $MoS_2$  nanosheet due to the hybridization reaction. The formed DNA-RNA heteroduplexes were digested by the DSN and the target miRNA was released to trigger the next hybridization reaction. Under optimal conditions, this sensor showed a wide dynamic range (10 fM–10 nM), low detection limit (10 fM) and high selectivity for let-7a analysis. In the same year, Xiao et al. also constructed a poly-cytosine (poly-C)mediated  $MoS_2$ -based nanoprobe coupled with a DSN signal amplification strategy for miRNA detection [29]. The introduction of a unique poly-C tails design led to a lower detection limit (3.4 fM) than classical molecular beacon-loaded MoS<sub>2</sub>-based nanoprobes. Other signal amplification strategies have also been introduced into the construction of fluorescence sensors based on  $MoS_2$ -based nanoprobes, such as catalytic hairpin assembly (CHA), a hybrid chain reaction (HCR), rolling circle amplification (RCA), etc., [86–90].

A  $MoS_2$ -based fluorescence nanoprobe is also a potential tool for the detection of intracellular biomolecules due to its excellent biocompatibility, such as ATP, microRNA, etc., [91–93]. For example, Ju and co-worker assembled a chlorine e6 (Ce6) labelled ATP aptamer onto a MoS<sub>2</sub> nanoplate to develop an intracellular nanoprobe for ATP detection and imaging based on the favorable biocompatibility [94]. It was noted that this designed MoS<sub>2</sub>-based nanoprobe not only sensitively and selectively analyzed ATP in living cells, but also could achieve controllable photodynamic therapy. Inspired by this exciting work, Li et al. immobilized two peptides onto a polydopamine (PDA)-functionalized MoS<sub>2</sub> nanointerface to construct a fluorescence nanoprobe for caspase-3 activity detection [28]. Caspase-3 was activated with the cell apoptosis, leading to the cleavage of a peptide labeled with fluorescence dye and the trigger of "turn on" fluorescence imaging. According to this design, the developed fluorescence biosensor showed a lower detection limit of 0.33 ng mL<sup>-1</sup> compared with some previous reports. For the purpose of trace biomolecules analysis, Zhu et al. developed an ultrasensitive fluorescence sensor for intracellular miRNA-21 detection and imaging based on MoS<sub>2</sub> nanoprobes by assembling three Cy3-labelled molecular beacons onto  $MoS_2$  nanosheets [95]. As shown in Figure 6c, the added miRNA-21 triggered a CHA reaction to form "Y"-shaped DNA structures with multiple Cy3 molecules. This interesting design obtained an ultralow detection limit (75.6 aM) for miRNA-21 detection compared to a general strand displacement-based strategy (8.5 pM). The excellent analytical performance was also proved by the intracellular imaging of miRNA-21 in human breast cancer cells.

# 3.6. SPR Sensors

MoS<sub>2</sub> and its nanocomposites have been considered as ideal substrates for the construction of SPR sensors due to the unique properties of a MoS<sub>2</sub> nanosheet, such as high charge carrier mobility and easily functionalization of noble metal nanoparticles [25,96]. As expected, MoS<sub>2</sub>-based SPR sensors are widely used to rapidly, label-free detect biomolecules or real-time and in-situ monitor the biological reaction. For example, Chiu et al. assembled carboxyl-functionalized MoS<sub>2</sub> sheets (MoS<sub>2</sub>-COOH) onto a gold surface to construct a SPR immunosensor for monitoring a bioaffinity interaction [95]. Experimental data showed that the SPR angles can be amplified by the MoS<sub>2</sub>-COOH chip, which was almost 1.9 folds and 3.1 folds than  $MoS_2$  and traditional SPR chips when the bovine serum albumin (BSA) concentration was 14.5 nM. Unfortunately, most of the works focused on the development of MoS<sub>2</sub>-based SPR substrates. To explore the potential application of a MoS<sub>2</sub>-based nanoprobe in SPR sensing field, Wang and co-workers developed a SPR biosensor for microRNA-141 (miRNA-141) analysis based on MoS<sub>2</sub>-AuNPs nanocomposites [73]. As shown in Figure 7, a classical sandwich structure was formed in the presence of miRNA-141. The localized plasmon of AuNPs supported onto MoS<sub>2</sub> nanosheets easily generated the electronic coupling by associating with Au film. As a result, an ultralow detection limit of 0.5 fM for miRNA-141 detection was obtained due to this signal amplification effect. Moreover, this designed SRP biosensor exhibited high selectivity for miRNA-200 family members' determination.



**Figure 6.** (a) Cartoon of MoS<sub>2</sub>-based fluorescence sensor for DNA detection. Reprinted with permission from [23]. Copyright 2013, American Chemical Society. (b) Illustration of Exo III-assisted fluorescence biosensor for streptavidin detection based on MoS<sub>2</sub>-based nanoprobe. Reprinted with permission from [69]. Copyright 2015, Elsevier. (c) MoS<sub>2</sub>-based nanoprobe coupled with signal amplification strategy for ultrasensitive detection and imaging of miRNA-21 expression in living cells. Reprinted with permission from [95]. Copyright 2019, American Chemical Society.



**Figure 7.** Schematic diagram of SPR biosensor for miRNA-141 detection based on MoS<sub>2</sub>-based nanoprobe. Reprinted with permission from [73]. Copyright 2017, Elsevier.

#### 4. Conclusions and Perspective

During the past decade,  $MoS_2$  as an emerging material has aroused more and more scientists' interests to construct MoS2-based nanoprobes due to its inherent advantages, including the large-scale preparation, tunable bandgap, excellent biocompatibility, easy functionalization with inorganic/organic groups, and outstanding optoelectronic properties. The introduction of MoS<sub>2</sub>-based nanoprobes means sensors coupled with different analytical methods have been successfully employed in environmental monitoring, food safety, biochemical analysis, disease diagnosis, and even homeland safety. With the assistance of MoS<sub>2</sub>-based nanoprobes, the developed sensors exhibited high sensitivity, selectivity, and stability for the detection of chemical and biological molecules. Though great advances in sensing application were obtained, MoS<sub>2</sub>-based nanoprobes still face some challenges in practical application. First, high-quality and large-scale preparation of  $MoS_2$  nanosheets and their nancomposites should be solved. It is the basic to construct a high-performance MoS<sub>2</sub>-based nanoprobe. The high-quality of the MoS<sub>2</sub> nanosheet often brings a high-performance MoS<sub>2</sub>-based nanoprobe. Controllable and large-scale preparation of MoS<sub>2</sub> nanosheets can ensure the repeatability of MoS<sub>2</sub>-based nanoprobes. Second, the recognition unit or signal amplification unit should be efficiently assembled onto the MoS<sub>2</sub> nanosheet and its nanocomposites. The assembled amount and spatial configuration of the recognition unit or signal amplification unit greatly affects the analytical performance. Third, the preparation mechanism of MoS<sub>2</sub>-based nanoprobes should be further studied. It is important to design a high-efficient nanoprobe for the construction of sensors. Finally, the best combination of the MoS<sub>2</sub>-based nanoprobe and detection method is another important influence parameter for obtaining better analytical performance. We believed that a MoS<sub>2</sub>-based nanoprobe will eventually be used in practical applications in the future with our joint efforts.

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#### References

- Shao, H.; Lin, H.; Guo, Z.; Lu, J.; Jia, Y.; Ye, M.; Su, F.; Niu, L.; Kang, W.; Wang, S.; et al. A Multiple Signal Amplification Sandwich-Type SERS Biosensor for Femtomolar Detection of miRNA. *Biosens. Bioelectron.* 2019, 143, 111616. [CrossRef] [PubMed]
- Yang, B.; Zhang, S.; Fang, X.; Kong, J. Double Signal Amplification Strategy for Ultrasensitive Electrochemical Biosensor Based on Nuclease and Quantum Dot-DNA Nanocomposites in the Detection of Breast Cancer 1 Gene Mutation. *Biosens. Bioelectron.* 2019, 142, 111544. [CrossRef] [PubMed]
- Bai, Y.; Li, H.; Xu, J.; Huang, Y.; Zhang, X.; Weng, J.; Li, Z.; Sun, L. Ultrasensitive Colorimetric Biosensor for Brca1 Mutation Based on Multiple Signal Amplification Strategy. *Biosens. Bioelectron.* 2020, 166, 112424. [CrossRef] [PubMed]
- Zhang, Y.; Wang, L.; Luo, F.; Qiu, B.; Guo, L.; Weng, Z.; Lina, Z.; Chen, G. An Electrochemiluminescence Biosensor for Kras Mutations Based on Locked Nucleic Acid Functionalized DNA Walkers and Hyperbranched Rolling Circle Amplification. *Chem. Commun.* 2017, 53, 2910–2913. [CrossRef] [PubMed]
- He, W.; Qiao, B.; Li, F.; Pan, L.; Chen, D.; Cao, Y.; Tu, J.; Wang, X.; Lv, C.; Wu, Q. A Novel Electrochemical Biosensor for Ultrasensitive Hg<sup>2+</sup> Detection via a Triple Signal Amplification Strategy. *Chem. Commun.* 2021, 57, 619–622. [CrossRef]
- Aithal, S.; Mishriki, S.; Gupta, R.; Sahu, R.P.; Botos, G.; Tanvir, S.; Hanson, R.W.; Puri, I.K. SARS-Cov-2 Detection with Aptamer-Functionalized Gold Nanoparticles. *Talanta* 2022, 236, 122841. [CrossRef]
- Ma, X.; Song, S.; Kim, S.; Kwon, M.-s.; Lee, H.; Park, W.; Sim, S.J. Single Gold-Bridged Nanoprobes for Identification of Single Point DNA Mutations. *Nat. Commun.* 2019, 10, 836. [CrossRef]
- Bakirhan, N.K.; Ozcelikay, G.; Ozkan, S.A. Recent Progress on the Sensitive Detection of Cardiovascular Disease Markers by Electrochemical-Based Biosensors. J. Pharm. Biomed. Anal. 2018, 159, 406–424. [CrossRef]
- Crapnell, R.D.; Dempsey-Hibbert, N.C.; Peeters, M.; Tridente, A.; Banks, C.E. Molecularly Imprinted Polymer Based Electrochemical Biosensors: Overcoming the Challenges of Detecting Vital Biomarkers and Speeding up Diagnosis. *Talanta Open* 2020, 2, 100018. [CrossRef]
- Dhara, K.; Mahapatra, D.R. Review on Electrochemical Sensing Strategies for C-reactive Protein and Cardiac Troponin I Detection. *Microchem. J.* 2020, 156, 104857. [CrossRef]
- Mao, K.; Zhang, H.; Wang, Z.; Cao, H.; Zhang, K.; Li, X.; Yang, Z. Nanomaterial-Based Aptamer Sensors for Arsenic Detection. Biosens. Bioelectron. 2020, 148, 111785. [CrossRef] [PubMed]
- Nezami, A.; Dehghani, S.; Nosrati, R.; Eskandari, N.; Taghdisi, S.M.; Karimi, G. Nanomaterial-Based Biosensors and Immunosensors for Quantitative Determination of Cardiac Troponins. *J. Pharm. Biomed. Anal.* 2018, 159, 425–436. [CrossRef] [PubMed]
- Song, C.; Li, F.; Guo, X.; Chen, W.; Dong, C.; Zhang, J.; Zhang, J.; Wang, L. Gold Nanostars for Cancer Cell-Targeted SERS-Imaging and NIR Light-Triggered Plasmonic Photothermal Therapy (PPTT) in the First and Second Biological Windows. *J. Mater. Chem. B* 2019, 7, 2001–2008. [CrossRef] [PubMed]
- 14. Song, C.; Li, J.; Sun, Y.; Jiang, X.; Zhang, J.; Dong, C.; Wang, L. Colorimetric/SERS Dual-Mode Detection of Mercury Ion via SERS-Active Peroxidase-Like Au@AgPt NPs. *Sens. Actuators B Chem.* **2020**, *310*, 127849. [CrossRef]
- Sui, C.; Yin, H.; Wang, L.; Zhou, Y.; Ai, S. Electrochemiluminescence Biosensor for DNA Hydroxymethylation Detection Based on Enzyme-Catalytic Covalent Bonding Reaction of -CH<sub>2</sub>OH and Thiol Functionalized Fe<sub>3</sub>O<sub>4</sub> Magnetic Beads. *Biosens. Bioelectron.* 2020, 150, 111908. [CrossRef]
- Li, F.; Pei, H.; Wang, L.; Lu, J.; Gao, J.; Jiang, B.; Zhao, X.; Fan, C. Nanomaterial-Based Fluorescent DNA Analysis: A Comparative Study of the Quenching Effects of Graphene Oxide, Carbon Nanotubes, and Gold Nanoparticles. *Adv. Funct. Mater.* 2013, 23, 4140–4148. [CrossRef]
- 17. He, S.; Song, B.; Li, D.; Zhu, C.; Qi, W.; Wen, Y.; Wang, L.; Song, S.; Fang, H.; Fan, C. A Graphene Nanoprobe for Rapid, Sensitive, and Multicolor Fluorescent DNA Analysis. *Adv. Funct. Mater.* **2010**, *20*, 453–459. [CrossRef]
- Liu, X.; Ge, J.; Wang, X.; Wu, Z.; Shen, G.; Yu, R. Development of a Highly Sensitive Sensing Platform for T4 Polynucleotide Kinase Phosphatase and Its Inhibitors Based on WS<sub>2</sub> Nanosheets. *Anal. Methods* 2014, 6, 7212–7217. [CrossRef]
- 19. Qi, C.; Cai, S.; Wang, X.; Li, J.; Lian, Z.; Sun, S.; Yang, R.; Wang, C. Enhanced Oxidase/Peroxidase-Like Activities of Aptamer Conjugated MoS<sub>2</sub>/PtCu Nanocomposites and Their Biosensing Application. *RSC Adv.* **2016**, *6*, 54949–54955. [CrossRef]
- Du, C.; Shang, A.; Shang, M.; Ma, X.; Song, W. Water-Soluble VS<sub>2</sub> Quantum Dots with Unusual Fluorescence for Biosensing. Sens. Actuators B 2018, 255, 926–934. [CrossRef]

- Asif, M.; Aziz, A.; Wang, H.; Wang, Z.; Wang, W.; Ajmal, M.; Xiao, F.; Chen, X.; Liu, H. Superlattice Stacking by Hybridizing Layered Double Hydroxide Nanosheets with Layers of Reduced Graphene Oxide for Electrochemical Simultaneous Determination of Dopamine, Uric Acid and Ascorbic Acid. *Microchim. Acta* 2019, 186, 61. [CrossRef] [PubMed]
- 22. Chi, J.; Li, J.; Ren, S.; Su, S.; Wang, L. Construction and Application of DNA-Two-Dimensional Layered Nanomaterials Sensing Platform. *Acta Chim. Sin.* 2019, 77, 1230–1238. [CrossRef]
- Zhu, C.; Zeng, Z.; Li, H.; Li, F.; Fan, C.; Zhang, H. Single-Layer MoS<sub>2</sub>-Based Nanoprobes for Homogeneous Detection of Biomolecules. J. Am. Chem. Soc. 2013, 135, 5998–6001. [CrossRef] [PubMed]
- Xiao, M.; Man, T.; Zhu, C.; Pei, H.; Shi, J.; Li, L.; Qu, X.; Shen, X.; Li, J. MoS<sub>2</sub> Nanoprobe for MicroRNA Quantification Based on Duplex-Specific Nuclease Signal Amplification. ACS Appl. Mater. Interfaces 2018, 10, 7852–7858. [CrossRef]
- Kaushik, S.; Tiwari, U.K.; Pal, S.S.; Sinha, R.K. Rapid Detection of Escherichia Coli Using Fiber Optic Surface Plasmon Resonance Immunosensor Based on Biofunctionalized Molybdenum Disulfide (MoS<sub>2</sub>) Nanosheets. *Biosens. Bioelectron.* 2019, 126, 501–509. [CrossRef]
- Yang, Y.; Liu, T.; Cheng, L.; Song, G.; Liu, Z.; Chen, M. MoS<sub>2</sub>-Based Nanoprobes for Detection of Silver Ions in Aqueous Solutions and Bacteria. ACS Appl. Mater. Interfaces 2015, 7, 7526–7533. [CrossRef]
- Sun, X.; Fan, J.; Fu, C.; Yao, L.; Zhao, S.; Wang, J.; Xiao, J. WS<sub>2</sub> and MoS<sub>2</sub> Biosensing Platforms Using Peptides as Probe Biomolecules. *Sci. Rep.* 2017, 7, 10290. [CrossRef]
- Li, X.; Li, Y.; Qiu, Q.; Wen, Q.; Zhang, Q.; Yang, W.; Yuwen, L.; Weng, L.; Wang, L. Efficient Biofunctionalization of MoS<sub>2</sub> Nanosheets with Peptides as Intracellular Fluorescent Biosensor for Sensitive Detection of Caspase-3 Activity. *J. Colloid Interface Sci.* 2019, 543, 96–105. [CrossRef]
- Xiao, M.; Chandrasekaran, A.R.; Ji, W.; Li, F.; Man, T.; Zhu, C.; Shen, X.; Pei, H.; Li, Q.; Li, L. Affinity-Modulated Molecular Beacons on MoS<sub>2</sub> Nanosheets for MicroRNA Detection. ACS Appl. Mater. Interfaces 2018, 10, 35794–35800. [CrossRef]
- Su, S.; Li, J.; Yao, Y.; Sun, Q.; Zhao, Q.; Wang, F.; Li, Q.; Liu, X.; Wang, L. Colorimetric Analysis of Carcinoembryonic Antigen Using Highly Catalytic Gold Nanoparticles-Decorated MoS<sub>2</sub> Nanocomposites. ACS Appl. Bio Mater. 2019, 2, 292–298. [CrossRef]
- Su, S.; Sun, Q.; Wan, L.; Gu, X.; Zhu, D.; Zhou, Y.; Chao, J.; Wang, L. Ultrasensitive Analysis of Carcinoembryonic Antigen Based on MoS<sub>2</sub>-Based Electrochemical Immunosensor with Triple Signal Amplification. *Biosens. Bioelectron.* 2019, 140, 77–82. [CrossRef] [PubMed]
- Su, S.; Cao, W.; Liu, W.; Lu, Z.; Zhu, D.; Chao, J.; Weng, L.; Wang, L.; Fan, C.; Wang, L. Dual-Mode Electrochemical Analysis of microRNA-21 Using Gold Nanoparticle-Decorated MoS<sub>2</sub> Nanosheet. *Biosens. Bioelectron.* 2017, 94, 552–559. [CrossRef] [PubMed]
- Ji, R.; Niu, W.; Chen, S.; Xu, W.; Ji, X.; Yuan, L.; Zhao, H.; Geng, M.; Qiu, J.; Li, C. Target-Inspired Pb<sup>2+</sup>-Dependent DNAzyme for Ultrasensitive Electrochemical Sensor Based on MoS<sub>2</sub>-AuPt Nanocomposites and Hemin/G-quadruplex DNAzyme as Signal Amplifier. *Biosens. Bioelectron.* 2019, 144, 111560. [CrossRef] [PubMed]
- 34. Singh, P.; Ojha, R.P.; Kumar, S.; Singh, A.K.; Prakash, R. Fe-Doped MoS<sub>2</sub> Nanomaterials with Amplified Peroxidase Mimetic Activity for the Colorimetric Detection of Glutathione in Human Serum. *Mater. Chem. Phys.* **2021**, 267, 124684. [CrossRef]
- 35. Tao, Y.; Lao, Y.-H.; Yi, K.; Xu, Y.; Wang, H.; Shao, D.; Wang, J.; Li, M. Noble Metal-Molybdenum Disulfide Nanohybrids as Dual Fluorometric and Colorimetric Sensor for Hepatitis B Virus DNA Detection. *Talanta* **2021**, 234, 122675. [CrossRef]
- Li, W.; Qiao, X.; Hong, C.; Ma, C.; Song, Y. A Sandwich-Type Electrochemical Immunosensor for Detecting CEA Based on CeO<sub>2</sub>-MoS<sub>2</sub> Absorbed Pb<sup>2+</sup>. *Anal. Biochem.* 2020, 592, 113566. [CrossRef]
- 37. Lu, C.; Huang, Z.; Liu, B.; Liu, Y.; Ying, Y.; Liu, J. Poly-Cytosine DNA as a High-Affinity Ligand for Inorganic Nanomaterials. *Angew. Chem. Int. Ed.* **2017**, *56*, 6208–6212. [CrossRef]
- Guo, S.; Wang, E. Noble Metal Nanomaterials: Controllable Synthesis and Application in Fuel Cells and Analytical Sensors. *Nano* Today 2011, 6, 240–264. [CrossRef]
- 39. Hartland, G.V. Optical Studies of Dynamics in Noble Metal Nanostructures. Chem. Rev. 2011, 111, 3858–3887. [CrossRef]
- 40. Huang, X.; Zeng, Z.; Bao, S.; Wang, M.; Qi, X.; Fan, Z.; Zhang, H. Solution-Phase Epitaxial Growth of Noble Metal Nanostructures on Dispersible Single-Layer Molybdenum Disulfide Nanosheets. *Nat. Commun.* **2013**, *4*, 1444. [CrossRef]
- Yuwen, L.; Xu, F.; Xue, B.; Luo, Z.; Zhang, Q.; Bao, B.; Su, S.; Weng, L.; Huang, W.; Wang, L. General Synthesis of Noble Metal (Au, Ag, Pd, Pt) Nanocrystal Modified MoS2 Nanosheets and the Enhanced Catalytic Activity of Pd-MoS<sub>2</sub> for Methanol Oxidation. *Nanoscale* 2014, *6*, 5762–5769. [CrossRef] [PubMed]
- Su, S.; Sun, H.; Cao, W.; Chao, J.; Peng, H.; Zuo, X.; Yuwen, L.; Fan, C.; Wang, L. Dual-Target Electrochemical Biosensing Based on DNA Structural Switching on Gold Nanoparticle-Decorated MoS<sub>2</sub> Nanosheets. ACS Appl. Mater. Interfaces 2016, 8, 6826–6833. [CrossRef] [PubMed]
- 43. Najmaei, S.; Mlayah, A.; Arbouet, A.; Girard, C.; Leotin, J.; Lou, J. Plasmonic Pumping of Excitonic Photoluminescence in Hybrid MoS<sub>2</sub>-Au Nanostructures. *ACS Nano* **2014**, *8*, 12682–12689. [CrossRef] [PubMed]
- 44. Miro, P.; Ghorbani-Asl, M.; Heine, T. Two Dimensional Materials Beyond MoS<sub>2</sub>: Noble-Transition-Metal Dichalcogenides. *Angew. Chem. Int. Ed.* **2014**, *53*, 3015–3018. [CrossRef]
- Jing, P.; Yi, H.; Xue, S.; Chai, Y.; Yuan, R.; Xu, W. A Sensitive Electrochemical Aptasensor Based on Palladium Nanoparticles Decorated Graphene–Molybdenum Disulfide Flower-Like Nanocomposites and Enzymatic Signal Amplification. *Anal. Chim. Acta* 2015, 853, 234–241. [CrossRef]

- Su, S.; Sun, Q.; Ma, J.; Zhu, D.; Wang, F.; Chao, J.; Fan, C.; Li, Q.; Wang, L. Ultrasensitive Analysis of Micrornas with Gold Nanoparticle-Decorated Molybdenum Disulfide Nanohybrid-Based Multilayer Nanoprobes. *Chem. Commun.* 2020, 56, 9012–9015. [CrossRef]
- You, H.; Mu, Z.; Zhao, M.; Zhou, J.; Yuan, Y.; Bai, L. Functional Fullerene-Molybdenum Disulfide Fabricated Electrochemical DNA Biosensor for Sul1 Detection Using Enzyme-Assisted Target Recycling and a New Signal Marker for Cascade Amplification. Sens. Actuators B 2020, 305, 127483. [CrossRef]
- Gao, Z.; Li, Y.; Zhang, X.; Feng, J.; Kong, L.; Wang, P.; Chen, Z.; Dong, Y.; Wei, Q. Ultrasensitive Electrochemical Immunosensor for Quantitative Detection of HBeAg Using Au@Pd/MoS<sub>2</sub>@MWCNTs Nanocomposite as Enzyme-Mimetic Labels. *Biosens. Bioelectron.* 2018, 102, 189–195. [CrossRef]
- 49. Ma, E.; Wang, P.; Yang, Q.; Yu, H.; Pei, F.; Li, Y.; Liu, Q.; Dong, Y. Electrochemical Immunosensor Based on MoS<sub>2</sub> NFs/Au@AgPt YNCs as Signal Amplification Label for Sensitive Detection of CEA. *Biosens. Bioelectron.* **2019**, *142*, 111580. [CrossRef]
- 50. Zhao, H.; Du, X.; Dong, H.; Jin, D.; Tang, F.; Liu, Q.; Wang, P.; Chen, L.; Zhao, P.; Li, Y. Electrochemical Immunosensor Based on Au/Co-BDC/MoS<sub>2</sub> and DPCN/MoS<sub>2</sub> for the Detection of Cardiac Troponin I. *Biosens. Bioelectron.* **2021**, *175*, 112883. [CrossRef]
- Ma, N.; Zhang, T.; Fan, D.; Kuang, X.; Ali, A.; Wu, D.; Wei, Q. Triple Amplified Ultrasensitive Electrochemical Immunosensor for Alpha Fetoprotein Detection Based on MoS<sub>2</sub>@Cu<sub>2</sub>O-Au Nanoparticles. *Sens. Actuators B* 2019, 297, 126821. [CrossRef]
- Li, S.K.; Liu, Z.T.; Li, J.Y.; Chen, A.Y.; Chai, Y.Q.; Yuan, R.; Zhuo, Y. Enzyme-Free Target Recycling and Double-Output Amplification System for Electrochemiluminescent Assay of Mucin 1 with MoS<sub>2</sub> Nanoflowers as Co-Reaction Accelerator. ACS Appl. Mater. Interfaces 2018, 10, 14483–14490. [CrossRef] [PubMed]
- Fan, Z.; Yao, B.; Ding, Y.; Xie, M.; Zhao, J.; Zhang, K.; Huang, W. Electrochemiluminescence Aptasensor for Siglec-5 Detection Based on MoS<sub>2</sub>@Au Nanocomposites Emitter and Exonuclease III-powered DNA Walker. *Sens. Actuators B* 2021, 334, 129592. [CrossRef] [PubMed]
- Wang, B.; Zhong, X.; Chai, Y.; Yuan, R. An ECL Biosensor for Sensitive Detection of concanavalin A Based on the ECL Quenching of Ru Complex by MoS<sub>2</sub> Nanoflower. Sens. Actuators B Chem. 2017, 245, 247–255. [CrossRef]
- 55. Dong, X.; Zhao, G.; Li, X.; Fang, J.; Miao, J.; Wei, Q.; Cao, W. Electrochemiluminescence Immunosensor of "Signal-Off" for β-amyloid Detection Based on Dual Metal-Organic Frameworks. *Talanta* 2020, 208, 120376. [CrossRef] [PubMed]
- 56. Bahari, D.; Babamiri, B.; Salimi, A.; Hallaj, R.; Amininasab, S.M. A Self-Enhanced ECL-RET Immunosensor for the Detection of CA19-9 Antigen Based on Ru(bpy)<sub>2</sub>(phen-NH<sub>2</sub>)<sup>2+</sup> - Amine-rich Nitrogen-Doped Carbon Nanodots as Probe and Graphene Oxide Grafted Hyperbranched Aromatic Polyamide as Platform. *Anal. Chim. Acta* 2020, 1132, 55–65. [CrossRef] [PubMed]
- Zhang, S.; Chen, Y.; Huang, Y.; Dai, H.; Lin, Y. Design and Application of Proximity Hybridization-Based Multiple Stimuli-Responsive Immunosensing Platform for Ovarian Cancer Biomarker Detection. *Biosens. Bioelectron.* 2020, 159, 112201. [CrossRef]
- Ju, P.; He, Y.; Wang, M.; Han, X.; Jiang, F.; Sun, C.; Wu, C. Enhanced Peroxidase-Like Activity of MoS<sub>2</sub> Quantum Dots Functionalized g-C<sub>3</sub>N<sub>4</sub> Nanosheets Towards Colorimetric Detection of H<sub>2</sub>O<sub>2</sub>. *Nanomaterials* 2018, *8*, 976. [CrossRef]
- Peng, J.; Weng, J. Enhanced Peroxidase-Like Activity of MoS<sub>2</sub>/Graphene Oxide Hybrid with Light Irradiation for Glucose Detection. *Biosens. Bioelectron.* 2017, 89, 652–658. [CrossRef]
- 60. Chi, M.; Zhu, Y.; Jing, L.; Wang, C.; Lu, X. Fabrication of Ternary MoS<sub>2</sub>-Polypyrrole-Pd Nanotubes as Peroxidase Mimics with a Synergistic Effect and Their Sensitive Colorimetric Detection of L-Cysteine. *Anal. Chim. Acta* **2018**, *1035*, 146–153. [CrossRef]
- 61. Tao, Z.; Wei, L.; Wu, S.; Duan, N.; Li, X.; Wang, Z. A Colorimetric Aptamer-Based Method for Detection of Cadmium Using the Enhanced Peroxidase-Like Activity of Au-MoS<sub>2</sub> Nanocomposites. *Anal. Biochem.* **2020**, *608*, 113844. [CrossRef] [PubMed]
- Wang, X.; Cheng, S.; Wang, X.; Wei, L.; Kong, Q.; Ye, M.; Luo, X.; Xu, J.; Zhang, C.; Xian, Y. pH-Sensitive Dye-Based Nanobioplatform for Colorimetric Detection of Heterogeneous Circulating Tumor Cells. ACS Sens. 2021, 6, 1925–1932. [CrossRef] [PubMed]
- 63. Sun, H.; Gao, Y.; Hu, N.; Zhang, Y.; Guo, C.; Gao, G.; Ma, Z.; Ivan Ivanovich, K.; Qiu, Y. Electronic Coupling between Molybdenum Disulfide and Gold Nanoparticles to Enhance the Peroxidase Activity for the Colorimetric Immunoassays of Hydrogen Peroxide and Cancer Cells. *J. Colloid Interface Sci.* **2020**, *578*, 366–378. [CrossRef] [PubMed]
- Jiang, J.; Liu, H.; Li, X.; Chen, Y.; Gu, C.; Wei, G.; Zhou, J.; Jiang, T. Nonmetallic SERS-Based Immunosensor Byintegrating MoS<sub>2</sub> Nanoflower and Nanosheet Towards the Direct Serum Detection of Carbohydrate Antigen 19-9. *Biosens. Bioelectron.* 2021, 193, 113481. [CrossRef] [PubMed]
- 65. Medetalibeyoglu, H.; Kotan, G.; Atar, N.; Yola, M.L. A Novel Sandwich-Type SERS Immunosensor for Selective and Sensitive Carcinoembryonic Antigen (CEA) Detection. *Anal. Chim. Acta* 2020, *1139*, 100–110. [CrossRef] [PubMed]
- 66. Fei, X.; Liu, Z.; Hou, Y.; Li, Y.; Yang, G.; Su, C.; Wang, Z.; Zhong, H.; Zhuang, Z.; Guo, Z. Synthesis of Au NP@MoS<sub>2</sub> Quantum Dots Core@Shell Nanocomposites for SERS Bio-Analysis and Label-Free Bio-Imaging. *Materials* **2017**, *10*, 650. [CrossRef] [PubMed]
- Shi, J.; Lyu, J.; Tian, F.; Yang, M. A Fluorescence Turn-on Biosensor Based on Graphene Quantum Dots (GQDs) and Molybdenum Disulfide (MoS<sub>2</sub>) Nanosheets for Epithelial Cell Adhesion Molecule (EpCAM) Detection. *Biosens. Bioelectron.* 2017, 93, 182–188. [CrossRef]
- Kong, R.-M.; Ding, L.; Wang, Z.; You, J.; Qu, F. A Novel Aptamer-Functionalized MoS<sub>2</sub> Nanosheet Fluorescent Biosensor for Sensitive Detection of Prostate Specific Antigen. *Anal. Bioanal. Chem.* 2015, 407, 369–377. [CrossRef]
- Xiang, X.; Shi, J.; Huang, F.; Zheng, M.; Deng, Q.; Xu, J. MoS<sub>2</sub> Nanosheet-Based Fluorescent Biosensor for Protein Detection via Terminal Protection of Small-Molecule-Linked DNA and Exonuclease III-aided DNA Recycling Amplification. *Biosens. Bioelectron.* 2015, 74, 227–232. [CrossRef]

- Chen, S.-C.; Lin, C.-Y.; Cheng, T.-L.; Tseng, W.-L. 6-Mercaptopurine-Induced Fluorescence Quenching of Monolayer MoS<sub>2</sub> Nanodots: Applications to Glutathione Sensing, Cellular Imaging, and Glutathione-Stimulated Drug Delivery. *Adv. Funct. Mater.* 2017, 27, 1702452. [CrossRef]
- Fan, Y.-Y.; Mou, Z.-L.; Wang, M.; Li, J.; Zhang, J.; Dang, F.-Q.; Zhang, Z.-Q. Chimeric Aptamers-Based and MoS<sub>2</sub> Nanosheet-Enhanced Label-Free Fluorescence Polarization Strategy for Adenosine Triphosphate Detection. *Anal. Chem.* 2018, 90, 13708–13713. [CrossRef] [PubMed]
- 72. Peng, X.; Wang, Y.; Wen, W.; Chen, M.-M.; Zhang, X.; Wang, S. Simple MoS<sub>2</sub>-Nanofiber Paper-Based Fluorescence Immunosensor for Point-of-Care Detection of Programmed Cell Death Protein 1. *Anal. Chem.* **2021**, *93*, 8791–8798. [CrossRef] [PubMed]
- Nie, W.; Wang, Q.; Yang, X.; Zhang, H.; Li, Z.; Gao, L.; Zheng, Y.; Liu, X.; Wang, K. High Sensitivity Surface Plasmon Resonance Biosensor for Detection of microRNA Based on Gold Nanoparticles-Decorated Molybdenum Sulfide. *Anal. Chim. Acta* 2017, 993, 55–62. [CrossRef]
- Lin, T.; Zhong, L.; Guo, L.; Fu, F.; Chen, G. Seeing Diabetes: Visual Detection of Glucose Based on the Intrinsic Peroxidase-Like Activity of MoS<sub>2</sub> Nanosheets. *Nanoscale* 2014, 6, 11856–11862. [CrossRef] [PubMed]
- Zhao, K.; Gu, W.; Zheng, S.; Zhang, C.; Xian, Y. SDS-MoS<sub>2</sub> Nanoparticles as Highly-Efficient Peroxidase Mimetics for Colorimetric Detection of H<sub>2</sub>O<sub>2</sub> and Glucose. *Talanta* 2015, 141, 47–52. [CrossRef]
- Zheng, J.; Song, D.; Chen, H.; Xu, J.; Alharbi, N.S.; Hayat, T.; Zhang, M. Enhanced Peroxidase-Like Activity of Hierarchical MoS<sub>2</sub> -Decorated N-Doped Carbon Nanotubes with Synergetic Effect for Colorimetric Detection of H<sub>2</sub>O<sub>2</sub> and Ascorbic Acid. *Chin. Chem. Lett.* 2020, *31*, 1109–1113. [CrossRef]
- Wan, L.; Wu, L.; Su, S.; Zhu, D.; Chao, J.; Wang, L. High Peroxidase-Mimicking Activity of Gold@Platinum Bimetallic Nanoparticle-Supported Molybdenum Disulfide Nanohybrids for the Selective Colorimetric Analysis of Cysteine. *Chem. Commun.* 2020, 56, 12351–12354. [CrossRef]
- Akbarzadeh, E.; Bahrami, F.; Gholami, M.R. Au and Pt Nanoparticles Supported on Ni Promoted MoS<sub>2</sub> as Efficient Catalysts for P-Nitrophenol Reduction. J. Water Process Eng. 2020, 34, 101142. [CrossRef]
- 79. Li, Z.; Zhang, M.; Liu, L.; Zheng, J.; Alsulami, H.; Kutbi, M.A.; Xu, J. Noble Metal and Fe<sub>3</sub>O<sub>4</sub>Co-Functionalizedco-Functionalized Hierarchical Polyaniline@MoS<sub>2</sub> Microtubes. *Colloids Surf. A* **2020**, *605*, 125347. [CrossRef]
- Ling, X.; Fang, W.; Lee, Y.-H.; Araujo, P.T.; Zhang, X.; Rodriguez-Nieva, J.F.; Lin, Y.; Zhang, J.; Kong, J.; Dresselhaus, M.S. Raman Enhancement Effect on Two-Dimensional Layered Materials: Graphene, h-BN and MoS<sub>2</sub>. *Nano Lett.* 2014, 14, 3033–3040. [CrossRef]
- Su, S.; Zhang, C.; Yuwen, L.; Chao, J.; Zuo, X.; Liu, X.; Song, C.; Fan, C.; Wang, L. Creating SERS Hot Spots on MoS<sub>2</sub> Nanosheets with in Situ Grown Gold Nanoparticles. ACS Appl. Mater. Interfaces 2014, 6, 18735–18741. [CrossRef] [PubMed]
- 82. Majee, B.P.; Srivastava, V.; Mishra, A.K. Surface-Enhanced Raman Scattering Detection Based on an Interconnected Network of Vertically Oriented Semiconducting Few-Layer MoS<sub>2</sub> Nanosheets. *ACS Appl. Nano Mater.* **2020**, *3*, 4851–4858. [CrossRef]
- Wang, Y.; Ma, T.; Ma, S.; Liu, Y.; Tian, Y.; Wang, R.; Jiang, Y.; Hou, D.; Wang, J. Fluorometric Determination of the Antibiotic Kanamycin by Aptamer-Induced FRET Quenching and Recovery between MoS<sub>2</sub> Nanosheets and Carbon Dots. *Microchim. Acta* 2017, 184, 203–210. [CrossRef]
- 84. Lu, Z.; Chen, X.; Hu, W. A Fluorescence Aptasensor Based on Semiconductor Quantum Dots and MoS<sub>2</sub> Nanosheets for Ochratoxin a Detection. *Sens. Actuators B Chem.* **2017**, 246, 61–67. [CrossRef]
- 85. Deng, H.; Yang, X.; Gao, Z. MoS<sub>2</sub> Nanosheets as an Effective Fluorescence Quencher for DNA Methyltransferase Activity Detection. *Analyst* **2015**, *140*, 3210–3215. [CrossRef]
- 86. Zhang, F.; Wang, S.; Feng, J.; Zou, R.; Xiang, L.; Cai, C. MoS<sub>2</sub>-Loaded G-quadruplex Molecular Beacon Probes for Versatile Detection of MicroRNA through Hybridization Chain Reaction Signal Amplification. *Talanta* **2019**, *202*, 342–348. [CrossRef]
- Dong, H.; Tang, S.; Hao, Y.; Yu, H.; Dai, W.; Zhao, G.; Cao, Y.; Lu, H.; Zhang, X.; Ju, H. Fluorescent MoS<sub>2</sub> Quantum Dots: Ultrasonic Preparation, up-Conversion and Down-Conversion Bioimaging, and Photodynamic Therapy. ACS Appl. Mater. Interfaces 2016, 8, 3107–3114. [CrossRef]
- 88. Wang, Y.; Ni, Y. Molybdenum Disulfide Quantum Dots as a Photoluminescence Sensing Platform for 2,4,6-Trinitrophenol Detection. *Anal. Chem.* **2014**, *86*, 7463–7470. [CrossRef]
- 89. Xu, S.; Li, D.; Wu, P. One-Pot, Facile, and Versatile Synthesis of Monolayer MoS<sub>2</sub>/WS<sub>2</sub> Quantum Dots as Bioimaging Probes and Efficient Electrocatalysts for Hydrogen Evolution Reaction. *Adv. Funct. Mater.* **2015**, *25*, 1127–1136. [CrossRef]
- Gu, W.; Yan, Y.; Cao, X.; Zhang, C.; Ding, C.; Xian, Y. A Facile and One-Step Ethanol-Thermal Synthesis of MoS<sub>2</sub> Quantum Dots for Two-Photon Fluorescence Imaging. J. Mater. Chem. B 2016, 4, 27–31. [CrossRef]
- 91. Oudeng, G.; Au, M.; Shi, J.; Wen, C.; Yang, M. One-Step in Situ Detection of miRNA-21 Expression in Single Cancer Cells Based on Biofunctionalized MoS<sub>2</sub> Nanosheets. *ACS Appl. Mater. Interfaces* **2018**, *10*, 350–360. [CrossRef] [PubMed]
- Liu, Y.; Zhang, Y.; Zhang, W.; Wang, X.; Sun, Y.; Huang, Y.; Ma, P.; Ding, J.; Song, D. Ratiometric Fluorescent Sensor Based on MoS<sub>2</sub> QDs and AuNCs for Determination and Bioimaging of Alkaline Phosphatase. *Spectrochim. Acta Part A* 2021, 262, 120087. [CrossRef] [PubMed]
- Xu, Y.; Kang, Q.; Yang, B.; Chen, B.; He, M.; Hu, B. A Nanoprobe Based on Molybdenum Disulfide Nanosheets and Silver Nanoclusters for Imaging and Quantification of Intracellular Adenosine Triphosphate. *Anal. Chim. Acta* 2020, 1134, 75–83. [CrossRef]

- 94. Jia, L.; Ding, L.; Tian, J.; Bao, L.; Hu, Y.; Ju, H.; Yu, J.-S. Aptamer Loaded MoS<sub>2</sub> Nanoplates as Nanoprobes for Detection of Intracellular ATP and Controllable Photodynamic Therapy. *Nanoscale* **2015**, *7*, 15953–15961. [CrossRef]
- 95. Zhu, D.; Huang, J.; Lu, B.; Zhu, Y.; Wei, Y.; Zhang, Q.; Guo, X.; Yuwen, L.; Su, S.; Chao, J.; et al. Intracellular MicroRNA Imaging with MoS<sub>2</sub>-Supported Nonenzymatic Catassembly of DNA Hairpins. ACS Appl. Mater. Interfaces 2019, 11, 20725–20733. [CrossRef] [PubMed]
- Chiu, N.-F.; Lin, T.-L. Affinity Capture Surface Carboxyl-Functionalized MoS<sub>2</sub> Sheets to Enhance the Sensitivity of Surface Plasmon Resonance Immunosensors. *Talanta* 2018, 185, 174–181. [CrossRef]