

Fundamental Characteristics of Photodetectors and Applications of Two-Dimensional Materials in Photodetection

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Abstract. Two-dimensional (2D) nanomaterials, due to their atomic-level thickness, tunable bandgap, and strong light-matter interaction, have emerged as a transformative platform for high-performance photodetectors. In recent years, devices based on emerging materials such as graphene, transition metal dichalcogenides (TMDs), $\text{Bi}_2\text{O}_2\text{Se}$, and InSe have achieved record-breaking light response, detection rate, and ultrafast response times. This article reviews performance optimization strategies, including heterostructure design, defect and doping control, interface passivation, and novel device structures (such as self-powered and flexible devices). It systematically compares key performance indicators such as response rate, external quantum efficiency, specific detection rate, dark current, and time-domain response. It assesses the trade-off between gain and speed, as well as challenges in large-scale fabrication, device consistency, and multi-functional integration. Finally, it looks forward to new directions such as wafer-level direct growth, polarization-sensitive detection, plasma or cavity enhancement technology to promote the development of next-generation wide-spectrum, low-power, and flexible photodetectors.

Keywords: two-dimensional nanomaterials, photodetectors, performance optimization, heterostructure design.

1. Introduction

Since graphene, the field of atomically thin materials has experienced explosive growth in optoelectronics, driven by its exceptional electronic properties and mechanical flexibility. Graphene exhibits ultrahigh carrier mobility—exceeding $1 \times 10^5 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ at room temperature in clean, encapsulated samples—and remains among the most conductive materials known [1]. This high mobility facilitates rapid transport in photodetectors. However, the lack of a bandgap and low optical absorbance per monolayer (typically $<10\%$) limit graphene's standalone utility in light detection. In contrast, transition metal dichalcogenide (TMD) monolayers such as MoS_2 and WSe_2 possess direct band gaps (in the range of $\sim 1.2\text{--}2.0 \text{ eV}$), enabling strong photon absorption and high on/off ratios in photodiodes. These properties make them inherently more suitable for photovoltaic photodetection [2].

Emerging two-dimensional semiconductors further extend functionality. Materials like InSe and $\text{Bi}_2\text{O}_2\text{Se}$ demonstrate fast intrinsic response times and high sensitivity even in the near infrared red (NIR). $\text{Bi}_2\text{O}_2\text{Se}$, for example, exhibits responsivity of $\sim 65 \text{ A W}^{-1}$ at 1200 nm and ultrafast response

(~1 ps) at room temperature thanks to its narrow bandgap and relatively good crystallinity from chemical vapor deposition (CVD) synthesis [3,4]. Additionally, self-driven van der Waals heterostructures combining WSe₂ and Bi₂O₂Se enable broadband operation with fast, bias-free response [5].

Despite these advances, several critical challenges remain unresolved. Firstly, per-layer optical absorption is intrinsically low (<10 %), necessitating complex stacking or metamaterial strategies to enhance responsivity. Secondly, achieving wafer-scale, high-quality growth with uniform thickness and crystallinity remains hard—device-to-device variability is a significant barrier to commercialization. Thirdly, implementing high responsivity, high detectivity, fast response, and low dark current in the same device often leads to performance trade-offs. Photoconductive designs with high gain are typically slower, whereas ultrafast photodiode architectures may compromise responsivity. Furthermore, polarization-sensitive detection and multifunctional device integration (e.g., combining imaging with communication capabilities) remain underexplored areas, representing significant research gaps to be addressed for next-generation technologies.

Key performance metrics for 2D photodetectors include responsivity (R , in A W⁻¹), external quantum efficiency (EQE), specific detectivity (D^* , in Jones), dark current (I_d), and response time (τ). Responsivity relates to how effectively incident photons are converted to current. EQE represents the fraction of photons converted to charge carriers. Detectivity combines sensitivity with noise performance. Dark current contributes directly to noise and degrades detectivity. Response time determines the usable bandwidth of photodetectors.

Strategies to optimize these metrics span several approaches. The first approach is heterostructure integration. Creating built-in electric fields via vertical stacking (e.g., TMD/TMD, graphene/TMD) or lateral junctions to enhance carrier separation and broaden the spectral response into UV-infrared ranges. These designs exploit Type-II band alignment and built-in fields to simultaneously suppress dark current and boost responsivity [5]. The second approach is defect and doping engineering. Introducing controlled chalcogen vacancies, substitutional doping, or surface functionalization to tune carrier lifetime, doping levels, or built-in fields—strike a balance between gain, noise, and speed. The third approach is Interface passivation: Suppressing interfacial trap states via chemical treatments, encapsulation (e.g., h-BN layers), or surface oxides—to reduce dark current, suppress noise, and enhance stability. The last approach is device architecture design: Employing Schottky diodes, p-n junctions, photoconductors, and self-powered photodetector configurations to navigate the trade-offs between responsivity, speed, gain, and dark current.

This paper focuses on advances in scalability, device performance, and integration. Emphasis is placed on realistic pathways toward scalable, high-performance 2D photodetectors suitable for real-world applications. By synthesizing recent progress across materials optimization, heterostructure design, and architecture innovations, this review aims to identify performance bottlenecks and chart future directions. The analysis should guide materials scientists and device engineers toward creating next-generation 2D photodetectors that deliver flexible imaging, low-power optical communication, environmental sensing, and wearable detection capabilities.

2. Fundamental principles of two-dimensional materials in photodetectors

2.1. Theoretical foundations of the 2D photoelectric detector

The working principle of photodetectors based on two-dimensional materials mainly relies on the following three aspects: band theory, carrier dynamics, and light-material interaction.

The band theory is a crucial foundation for understanding the response and performance of devices. This theory encompasses two main light detection mechanisms: the photovoltaic effect and photoconductive effect [6]. The photovoltaic effect occurs in structures with an intrinsic electric field, such as Schottky junctions or p-n heterojunctions [7,8]. When two-dimensional materials are exposed to light, photons can excite electron-hole pairs. These charge carriers are rapidly separated under the drive of the intrinsic electric field, with electrons being attracted to the n-type region and holes flowing to the p-type region, thereby generating current in the absence of bias. Such photovoltaic structures typically exhibit fast response and low dark current [9]. For example, in the 2D MoS₂/p-Si heterostructure, due to the existence of the interface potential barrier, high-energy photons (such as visible or ultraviolet) can efficiently separate photogenerated electron-hole pairs and output photocurrent. The detection rate can reach 10¹³-10¹⁴ Jones, demonstrating the high-performance advantage of this mode [10]. The structure of the device is shown in Figure 1.

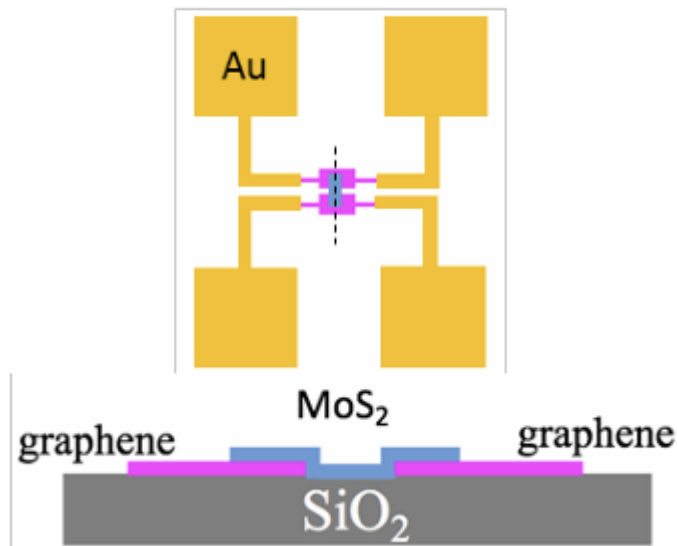


Figure 1. The structure of the Graphene/MoS₂/Graphene devices [10]

The photoconductive effect depends on the applied bias voltage, and the photoconductive device involves carrier dynamics, which determine the gain and speed of the device. Under bias voltage, the carriers are driven towards the electrode region, generating leakage current. Crucially, trap-mediated carrier localization, such as via defect states or interfacial traps, delays charge recombination, leading to a prolonged effective carrier lifetime (τ) [11]. Due to the average transport time (τ_t) of carriers being much shorter than the τ , the photogenerated carriers generated by each photon can participate in multiple cycles and be collected multiple times through the channel. The gain approximately satisfies the relationship $G \approx \tau / \tau_t$, and the responsivity can reach 10⁶ A/W [5,12]. Taking black phosphorus (BP) as an example, it is prone to form defects and has relatively low mobility channels. The device exhibits a remarkably strong photoelectric response exceeding 10⁶ A/W in the NIR spectral region, while its temporal response is limited to the microsecond-to-millisecond range owing to the substantial τ [8].

The light-matter interaction in a two-dimensional system can be quantified by the EQE, which is defined as the ratio of the number of collected charge carriers to the number of incident photons [7]:

$$EQE(\lambda) = \frac{I_{ph} h\nu}{e P_{opt}(\lambda)} \quad (1)$$

Among them, I_{ph} represents the photocurrent, $P_{opt}(\lambda)$ is the optical power at wavelength λ , $h\nu$ is the photon energy, and e is the electron charge. Despite the ultrathin nature (<1 nm), monolayer two-dimensional materials exhibit strong excitonic effects that enable external quantum efficiencies exceeding 40% in certain device architectures. After the material is combined with plasmas or optical resonators, the absorption capacity can be improved by more than ten times [7].

2.2. Typical spectral range of photodetectors

2.2.1. Ultraviolet-visible light (200-700 nm)

In the UV-visible wavelength range, photodetectors based on wide-bandgap two-dimensional materials, such as GaSe, Transition metal dichalcogenides (TMDs) and GaSe/MoS₂ composite junctions, are more favored due to their strong layer-dependent absorption and clear band-edge characteristics. The single-layer or few-layer fragments of GaSe have a direct bandgap of approximately 2.0 eV, enabling efficient photon absorption near 350 nm. For example, the type II heterojunction device of GaSe/MoS₂ achieves a response of ~ 42.6 A/W and a detection rate of $\sim 8 \times 10^{12}$ Jones at a wavelength of 300 nm, with its performance being three times that of a single-layer MoS₂ device [7]. Additionally, the simple-exfoliated GaSe device achieves a response of about 5 A/W under 405 nm laser irradiation of 42 mW/cm², with extremely low dark current (~ 21 pA) and a slow response speed (rise time of about 34 ms), mainly influenced by the photoconductive mechanism. Such devices can be fabricated through vapor-phase growth or exfoliation, but large-scale, high-quality industrial production remains challenging. Furthermore, WSe₂/graphene heterojunction has an EQE of over 50% under visible light [7,13].

2.2.2. Near infrared-short wave infrared (700-1700 nm)

The bandgap of the layer thickness of black phosphorus (BP) can be adjusted over the visible light range to approximately 4 μ m, and the carrier mobility can exceed 10^3 cm²/V·s. When BP is combined with PbS or PbSe quantum dots on two-dimensional materials such as MoS₂ to form a type II heterojunction interface, a responsivity of over 10^2 A/W can be achieved in the 1.55 μ m band, and the SWIR detection rate is approximately 10^{11} Jones. The BP/In₂Se₃ ferroelectric-controlled vdW heterojunction further shows tunable photo response and enhanced contrast due to built-in polarization fields, enabling gate-modulated responsivity in nearIR. BP devices can be fabricated through precise transfer or CVD growth, but they are prone to degradation in the air and require encapsulation treatment [7,8].

2.2.3. Terahertz (30-3000 μ m)

Graphene possesses a zero bandgap and high mobility, making it an ideal material for terahertz detection. Graphene FETs with integrated antenna structures (such as bow-tie or plasma slot structures) utilize the photovoltaic effect (PTE) to achieve a NEP $< 10^{-12}$ W/Hz^{1/2}, a response time < 3.3 ns, and a broadband (0.5 - 10 THz) detection capability. These room-temperature THz detectors have a dynamic range covering four orders of magnitude and are suitable for high-resolution spectroscopy, imaging, and communication. The manufacturing process relies on CVD-grown graphene combined with nano-scale antenna patterns, although it has the potential for large-area scalability, it requires high alignment accuracy [8,14].

2.3. Performance characteristics of photodetectors

Based on the above materials and structures, high-performance two-dimensional photodetectors possess the properties of wideband detection, high responsivity (R), specific detectivity (D^*), and fast response speed (τ or bandwidth f_c). These figures of merit are interdependent and governed by the underlying physics and device architectures.

Firstly, through heterostructure stacking or hybridization, a wide spectrum operation from UV to THz can be achieved, demonstrating multi-mode detection capabilities [7,8].

Secondly, responsivity (R) expresses how effectively incident light power is converted into photocurrent. Its fundamental relation is [15]:

$$R(\lambda) = EQE \frac{\lambda q}{hc} = \frac{I_{ph}}{P_o} \quad (2)$$

Where λ is the wavelength, q is the electronic charge, h is Planck's constant, c is the speed of light, I_{ph} is the photocurrent, and P_o is the amount of optical power incident on the detector. In practice, R values range widely: photodiode devices exceed 10^3 A/W, BP devices can reach 10^6 A/W, and graphene/organic systems achieve $>5 \times 10^5$ A/W in the UV-visible range [7,8].

Thirdly, in terms of specific detectivity (D^*), it normalizes detectivity to device area and bandwidth, capturing noise-limited sensitivity. It is defined as [15]:

$$D^* = \frac{\sqrt{A}}{NEP} = \frac{R(\lambda) \sqrt{A \Delta f}}{I_N} \quad (3)$$

Where A is the sensing area, Δf is the bandwidth, NEP is the noise equivalent power, and I_N is the total current noise. Van der Waals (VDW) heterostructures and 2D/Si devices often exceed 10^{12} Jones, FePSe₃/MoS₂ reaches 1.5×10^{13} , and MoS₂/GaAs heterojunction can reach above 10^{14} Jones [16]. Finally, response speed (τ or f_c) is governed by carrier transit time, RC time constant [17]:

$$f_{RC} = \frac{1}{2\pi RC} \quad (4)$$

Where R is the load resistance and C is the capacitance. Oxygen sulfide devices achieve picosecond levels, waveguide structures have a bandwidth of tens of GHz, while BP devices have a slower response time (in the millisecond range). In summary, material selection, heterostructure design, and device structure determine the trade-offs in performance, providing customized optimization solutions for imaging, optical communication, and sensing applications [6,8].

3. Comprehensive analysis and discussion on the performance of photodetectors

3.1. Performance analysis and challenges

A systematic comparison of optimization strategies for two-dimensional material photodetectors reveals that there are performance trade-offs among various methods. Van der Waals Type-II heterostructures, such as MoS₂/WSe₂ and InSe/graphene, utilize the built-in electric field to achieve efficient carrier separation, with their detectivity D exceeding 10^{13} Jones and the responsivity reaching 10^3 - 10^4 A/W [8]. However, due to the trap capture mechanism, the response time is typically in the microsecond to millisecond range. In contrast, plasma-enhanced devices concentrate the light field through metal nanostructures, enabling sub-nanosecond response while maintaining a

responsivity of nearly 10^3 A/W. Nonetheless, the manufacturing process is complex and is limited by the effective area [18]. The fabrication process exhibits considerable complexity and is restricted by the effective area.

Ferroelectric-2D hybrid structures, such as $\text{ZnIn}_2\text{S}_4/\text{TMD}$, can achieve self-powered operation, with a D value of approximately 10^{12} Jones, a response rate of $>10^3$ A/W, and operation without bias [19]. Nevertheless, critical challenges remain in terms of cycle stability and material fatigue, which require further optimization. Hybrid dimensional structures, such as PbS QD/graphene, CNT/MoS₂, provide broadband UV–SWIR detection and gains up to 10^5 , yet uniformity and trap-limited response (about 100 μs) restrict their throughput in high-speed applications [20]. A comparative summary is shown in Table 1 below.

The common challenges faced by all platforms include dark current suppression, noise management, and spectral coverage issues. High-gain photovoltaic conductive devices typically exhibit elevated $1/f$ and generation-recombination (g-r) noise. Without strict noise equivalent power (NEP) characterization, D is prone to being overestimated [21]. To extend to mid-infrared, long-wave infrared, and terahertz frequency bands, additional hybrid or low-temperature operation is required, increasing system complexity and power consumption [22]. The large-scale and highly consistent fabrication of defect-free 2D layers and their heterojunctions remains a bottleneck for commercialization [6].

Table 1. Comparative Performance of 2D Photodetector Optimization Strategies

Strategy	D (Jones)	Responsivity (A/W)	Speed (μs)	Key Challenges
Type-II Heterostructure [8,18,22]	$>10^{13}$	$10^3\text{-}10^4$	$1\text{-}10^3$	Trap-limited speed, synthesis scale
Plasmonic Enhancement [18,23,24]	$\sim 10^{14}$	$\sim 10^3$	$<10^{-3}$	Fabrication complexity, area coverage
Ferroelectric Hybrid [19,25]	$\sim 10^{12}$	$>10^3$	$1\text{-}10^{-3}$	Fatigue, material stability
Mixed-Dimensional [20]	$\sim 10^{13}$	$10^2\text{-}10^3$	~ 100	Uniformity, noise

3.2. Optimization methods for two-dimensional materials

3.2.1. Optimization via material structural engineering

Current performance optimization paradigms in materials engineering primarily involve the implementation of van der Waals heterostructures, organic hybrid architectures, two-dimensional/silicon heterojunctions, and defect passivation methodologies. Firstly, the vertically stacked structure with type-II band steps (such as MoS₂/WSe₂) can effectively separate charge carriers and broaden the spectral response to the ultraviolet-near infrared (UV-NIR) range [26]. Secondly, the organic-2D hybrid system combines the high mobility of graphene with organic semiconductors, leveraging their respective absorption and injection advantages to achieve a response rate greater than 10^3 A/W in the ultraviolet region [7]. Moreover, the 2D/silicon heterojunction can achieve a detection rate $D > 10^{12}$ Jones at room temperature by combining mature silicon processes [6,8]. Finally, by chemically treating (such as super-acid treatment of MoS₂) or hexagonal boron nitride encapsulation for defect passivation, the trap recombination can be reduced, and the NEP of dark current and noise can be decreased [6,7].

3.2.2. Optimization via device structure

In terms of device structure, by arranging metal nanostructures (such as Au nanowires/nanoparticles) near the two-dimensional layer to concentrate electromagnetic fields, a localized plasmon resonance can be formed with the two-dimensional layer. For example, periodic arrays of gold nano-disks or nanorods with diameters of approximately 18 nm, 60 nm, or 110 nm placed at controlled interparticle gaps (~ 2 nm) and surface-to-layer spacing produce intense nearfield “hotspots” at disk edges and between particles, tuned to ~ 530 – 630 nm resonance; these arrays yield photocurrent enhancements of over 800–1,500 % compared to bare material, while avoiding excessive parasitic capacitance. This can enhance absorption while maintaining response speed [8,14]. In waveguide integration, superimposing the two-dimensional layer onto a silicon waveguide enables a bandwidth greater than 50 GHz and a micron-scale size. Additionally, the thermoelectric or thermophotovoltaic effects of two-dimensional ferroelectric materials, such as ZnIn_2S_4 , can be utilized to convert the temperature change, which is caused by light absorption, into an instantaneous electric potential. This method can achieve self-powered detection under zero bias, and has an inherent gain mechanism [6,8].

3.3. Application-oriented optimization

Imaging and sensing applications demand ultrahigh detectivity and minimal noise. MoS_2/Si photodiodes, enhanced by surface passivation and resonant cavities, achieve D above 10^{13} Jones, enabling high-contrast, low-light imaging at room temperature [2]. Organic-2D hybrids further boost UV sensitivity, with responsivities nearing 10^{13} A/W under low-power illumination, facilitating flame detection and environmental monitoring [23].

In optical communications, speed is paramount. Ternary oxyselenide photodetectors ($\text{Bi}_2\text{O}_2\text{Se}$) deliver picosecond-scale response times (>100 GHz bandwidth) with moderate responsivity (~ 65 A/W), making them ideal for high-bit-rate interconnects [22]. Waveguide-integrated graphene devices also achieve >50 GHz operation within silicon photonics platforms, balancing CMOS compatibility with ultrafast performance [6].

Flexible and wearable electronics require mechanical robustness and low power. Printed CNT/graphene hybrid sensors maintain responsivity $>10^2$ A/W under bending radii <5 mm, with self-powered architectures based on pyro-phototronic effects in ferroelectric 2D layers, ideal for health monitoring and e-skin applications [25]. However, their detectivity ($\sim 10^{10}$ Jones) and spectral range remain limited compared to rigid counterparts.

3.4. Performance optimization direction of photodetectors

To overcome scale and performance barriers, scalable synthesis of large-area, transfer-free 2D heterostructures via roll-to-roll CVD and direct growth on flexible substrates is under intense development. Advances in in situ passivation and layer-by-layer growth promise to reduce defects and maintain built-in field strengths across wafer scales [8].

Multifunctional integration—combining photodetection with memory, polarization sensitivity, and thermal sensing—has emerged as a promising research frontier. Ferroelectric-2D hybrids already demonstrate nonvolatile photogating, while anisotropic materials such as ReS_2 and BP enable on-chip polarization-encoded imaging and encryption [27]. Quantum photodetectors leveraging single-photon avalanche in TMDC heterostructures and QD/2D hybrids aim to achieve

single-photon sensitivity and timing resolution below 100 ps, critical for quantum communications and light detection and ranging (LIDAR) systems [24].

Progress in industrial-scale synthesis, multifunctional integration, and quantum-level sensitivity positions 2D-material photodetectors as critical enablers for next-generation optoelectronics, from ultrafast communications to quantum information processing.

4. Conclusion

This paper summarizes the methods for optimizing the performance of photodetectors based on two-dimensional nanomaterials and their mixed systems, and summarizes four main approaches: heterostructure engineering, defect and doping control, interface passivation, and innovative device architectures. Heterostructures, especially Type-II van der Waals stacking and hybrid-dimensional structures, achieve high detection rates by efficiently separating charge carriers and covering a wide spectral range; defect/doping engineering provides an effective means to regulate carrier lifetime and transmission performance, thereby achieving intentional trade-offs between responsivity and speed; interface passivation, such as encapsulation and chemical treatment, can significantly reduce trap states and dark current, directly improving noise performance and device stability; device structure innovation (plasma enhancement, waveguide integration, ferroelectric/thermoelectric hybridization, etc.) offers orthogonal approaches to enhance absorption, accelerate response, or achieve self-power supply. Overall, selectively combining these methods for specific target performance often approaches the ideal performance curve more closely than a single strategy.

These technological advancements have significant practical implications. The optimized two-dimensional photodetectors can offer advantages in terms of volume, weight, and flexibility that traditional bulk materials or III-V devices cannot match, thereby providing new solutions in areas such as wearable biosensing, foldable imaging arrays, portable environmental sensing, and on-chip optical interconnects. Moreover, two-dimensional materials can be precisely controlled at the atomic level to align the energy bands, reduce trap density, and modulate interface electric fields, enabling the detection function to be designed in coordination with electronic/optical components, thus achieving multi-functional integration such as sensing and storage, polarization selection, or on-chip spectral filtering. The development of low-power or self-powered systems also makes two-dimensional detectors suitable for energy-constrained scenarios such as the Internet of Things and distributed sensing.

To advance the laboratory progress to industrialization, future research should prioritize the following: First, achieve scalable and repeatable growth and integration processes to ensure the quality of grain boundaries and interfaces; second, establish standardized testing and reporting standards to make performance comparisons comparable and operational; third, promote the systematic integration of multi-functional devices, coupling detection, signal processing or memory functions onto a single chip; fourth, deeply explore quantum-level and polarization-sensitive detection mechanisms based on anisotropic or ferroelectric two-dimensional compounds. At the same time, long-term stability, packaging, and manufacturing costs are also key factors for technology implementation. Through collaborative efforts in material synthesis, device design, and system integration, two-dimensional photodetectors are expected to outperform existing solutions in several application scenarios and become the next generation of flexible, low-power, and wide-spectrum adaptable core components in optoelectronics.

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