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Self-biased visible-NIR photodetection enabled via a dual-heterojunction n-MoS₂/p-CuO/n-Si design

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In this study, a comprehensive dual-junction (n-MoS₂/p-CuO/Si and p-CuO/n-Si) evaluation of a self-biased heterostructure was conducted for photodetector applications. Owing to the integration of both junctions, the proposed design offered dual-response functionality, under zero bias, corresponding to the visible (625 nm) and NIR (720 and 808 nm) regions. At zero applied bias, the n-MoS₂/p-CuO/Si heterojunction exhibited a responsivity (R_{λ}) of 21.04/30.50 mA W⁻¹ and a detectivity (D^*) of $1.0 \times 10^{14}/1.5 \times 10^{14}$ Jones at incident wavelengths of 625/720 nm; this highlights the self-biased nature of the fabricated design. The attained values were found to be dramatically increased under a 3 V bias, with R_{λ} values of 0.144 and 0.124 A/W for the n-MoS₂/p-CuO/Si and p-CuO/n-Si heterostructures, respectively. The observed figures-of-merit consistently reduced as the incident light intensity increased, indicating a strong negative correlation, which was further confirmed by the R^2 value approaching unity ($R^2 = 1$). The time-resolved features confirmed response/recovery times of 0.27/0.36 and 0.41/0.48 s, respectively, for the addressed heterostructures, highlighting the suitability of this design for efficient, bias-free photodetection over Vis-NIR wavelengths.

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1. Introduction

Photodetector-based optoelectronics, specifically those fabricated using Si technology, are being broadly considered for optical and electronic applications, UV/Vis/IR optical communication, flame detection, environmental and biological sensing, and imaging because of their CMOS technology compatibility, easy fabrication process, and low cost.^{1–3} Nevertheless, the characteristic restrictions of Si technology, such as constrained absorption in the Vis-NIR range and indirect optical bandgap, have prompted the incorporation of complementary semiconductor metal oxides to boost the spectral response and subsequently the device performance.^{4–6} Thus, heterojunction-based Si optoelectronics, dual-band photodetectors in particular, are being further investigated among research and industrial societies due to their benefits compared with their homojunction counterparts; these include high photoresponsive performance, almost zero power consumption, and wide spectral response. Further, dual-band photodetectors have demonstrated excellent potential in this regard, allowing enhanced charge separation effectiveness

through wide spectral absorption at zero applied bias. Accordingly, various dual-band optoelectronics have been developed with relatively high photo-responsive properties, including high R_{λ} , excellent D^* , fast rise/fall times, and high on/off ratio; these include CuO-TiO₂/Si, CuO/Cu₂O/Si, ZnO/ZnTe/Si, and WSe₂/WS₂/p-Si.^{7–10} Notably, the combination of p-type CuO, a narrow optical bandgap metal oxide semiconductor with relatively robust absorption within the visible spectrum, along with an Si wafer, has been broadly evaluated. CuO, with p-type conductivity, is known to be favorable for its band alignment with Si, enabling well-organized carrier separation and transport across the formed heterojunction, thereby allowing an enhanced photo-response behavior even under low applied bias.^{11–13} Moreover, the integration of a two-dimensional (2D) layered material, like MoS₂, could alter the photodetector functionality. MoS₂ exhibits solid light-matter interaction, excellent scalability, and relatively high carrier mobility,^{14–16} when n-MoS₂ is configured onto a p-CuO/n-Si stack, the proposed candidate can facilitate a dual-heterostructure design, enabling spectral lengthening to the NIR range, effective carrier collection, and self-driven operation via a built-in electric field. In this context, we report a multi-band (p-n and n-p) dual-junction optoelectronic design. Moreover, a methodical fabrication procedure for the intended dual-band, fast-response, self-driven heterostructure is elucidated. In detail, the proposed heterojunction (n-MoS₂/p-CuO) resulted in pronounced figure-of-merit profiles along a broad range at zero applied bias voltage.

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2. Experimental procedure

The intended dual-heterostructured device (n-MoS₂/p-CuO/n-Si) was fabricated *via* a pulsed laser deposition (PLD) process along with selective masking in order to attain a spatially patterned design. In detail, a commercially presented n-type Si wafer (Sigma Aldrich, 100, 1–10 Ω cm) was utilized as the main substrate. The Si wafer was thoroughly cleaned using the RCA standard procedure to remove the undesired contaminants; then, the wafer was diluted with HF to eliminate native oxides. To create the spatially patterned dual-heterojunction foundation (Fig. 1a), the wafer was partially covered, allowing two-thirds to be exposed to the generated plasma plume. Herein, a high-purity Cu target was irradiated at 6 J cm⁻² using a 2nd harmonic Q-switched Nd:YAG laser (532 nm) with a repetition rate of 10 Hz. The temperature of the Si wafer was maintained at 200 °C in an oxygen environment of 10⁻³ mbar, allowing the deposition of a CuO layer; the attained substrate was then allowed to cool naturally to room temperature. Thereafter, two-thirds of the attained geometry was covered, allowing the deposition of MoS₂ on one half of the CuO layer (Fig. 1a). In detail, the MoS₂ target was ablated by the same laser source (6.5 J cm⁻²) to obtain the n-MoS₂/p-CuO/n-Si heterostructure; the MoS₂ film covered one half of the deposited CuO film, as demonstrated in Fig. 1a, forming distinct dual-heterojunctions. In order to enhance the interfacial quality and crystallinity of the deposited layers, the attained geometry was thermally treated in air at 250 °C for 45 minutes. Finally, Ag electrodes were deposited by thermal evaporation to define the active

device area and ensure reliable electrical contact within the dual-heterojunction architecture.

The structural investigation was conducted using X-ray diffraction techniques (Bruker, AXSD8, XRD), while the optical investigation was performed *via* UV-Vis spectrophotometry (UV-3600, Shimadzu). The topographies of the deposited layers were examined using field-emission scanning electron microscopy (SU8030, Hitachi, FE-SEM). Simultaneously, a room-temperature photoresponsive evaluation was conducted utilizing a source measure unit (Keithley 2400, SMU) in conjunction with a narrow optical-bandpass filter/s (ThorLabs), allowing a wavelength scan from 405 nm to 808 nm. The incident light intensity, transmitted through the filter, was measured at the light incident spot with an LX2-illuminance meter (Japan, Sanwa). Moreover, the time-dependent features were projected by considering 10% to 90% of the entire attained photocurrent (I_{ph}) by histogram search, while the incident light pulse-width was fixed at ~12.5 s.

3. Results and discussion

Fig. 1b depicts the XRD diffraction patterns of the deposited CuO and MoS₂ layers, in which the addressed layers showed polycrystalline orientation. Specifically, the XRD spectrum of CuO exhibits five distinct diffraction planes (110), (101), (111), (-202), and (020), which are characteristic of the monoclinic CuO structure, in agreement with JCPDs card no. 80-1919; these planes were identified at 2θ values of 32.426°, 35.407°, 38.641°, 48.684°, and 58.221°, respectively. Simultaneously,

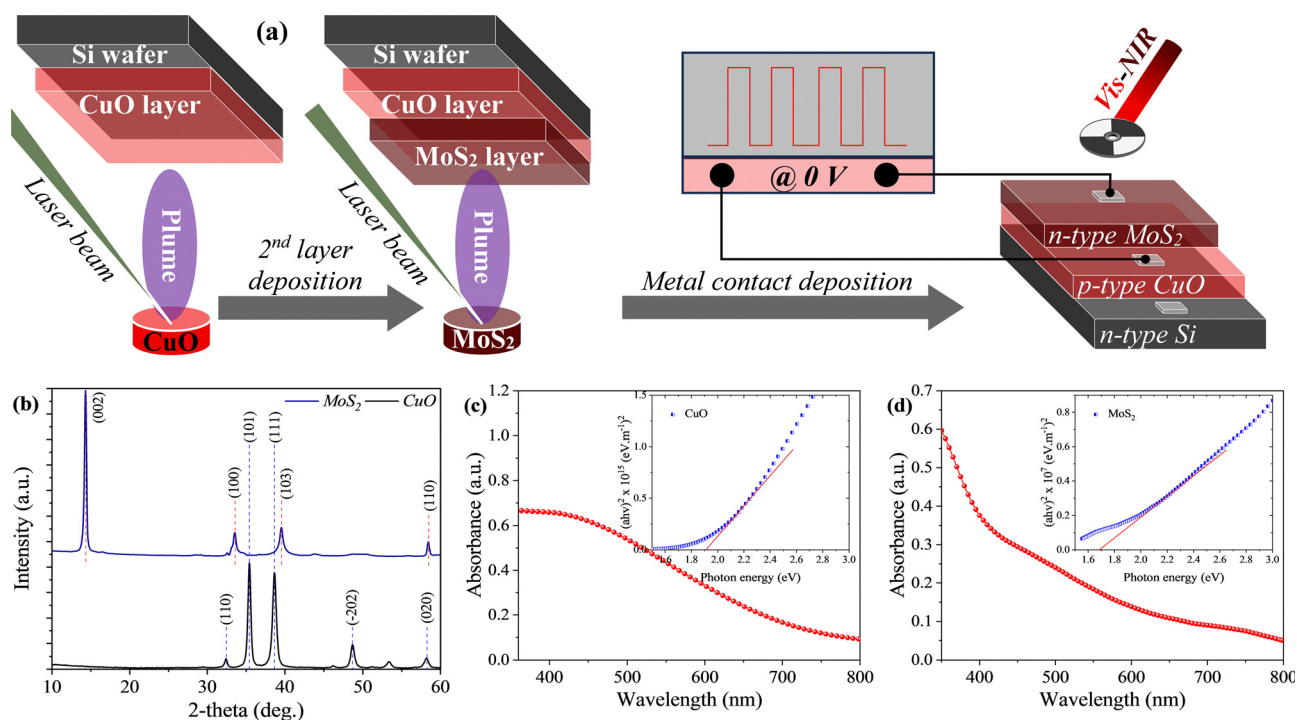


Fig. 1 (a) Schematic of the pulsed laser deposition process for the design of the fabricated device and the measurement set-up. (b) XRD patterns and UV-Vis spectra and optical bandgap (inset) of the deposited (c) CuO and (d) MoS₂ layers.



four prominent distinct XRD patterns for the MoS₂ layer were observed at around $2\theta \approx 14.194^\circ$, 33.536° , 39.542° , and 58.440° , which correspond to the (002), (100), (103), and (110) planes, respectively. The positions and sharpness of these reflections indicate well-aligned, high crystallinity structures that closely match the hexagonal 2H-MoS₂ phase, as confirmed by JCPDS card no. 37-1492. Optical analysis of the deposited CuO film (Fig. 1c) revealed a broad absorption curve along the visible range, which indicates good light-harvesting ability in the visible spectrum. The optical bandgap of the addressed layer (CuO), estimated according to the Tauc relation,^{17,18} exhibited a value of 1.91 eV (see the inset of Fig. 1c). Similarly, MoS₂ (Fig. 1d) demonstrated a smooth increase in absorption from the visible to the UV region; however, the absorption curve lacks an excitonic peak characteristically detected in monolayer MoS₂, which, in turn, suggests bulk-like/multilayer structure formation during the deposition using the PLD method. The optical bandgap (see the inset of Fig. 1d) was found to be 1.68 eV, corresponding to an incident wavelength of ~ 738 nm. The topography analysis of the CuO layer (Fig. 2a) revealed the formation of relatively compressed and uniformly distributed granular particles, suggesting a polycrystalline structure along random crystallite orientations within the > 50 μm scanned area. The particle distribution appears moderately heterogeneous, with isolated bright regions across the scanned area, which could be due to surface roughness and/or localized agglomeration. Such morphological singularities are frequently obtained in CuO films attained using the PLD technique.^{19,20} In comparison to the CuO morphology, MoS₂ (Fig. 2b) exhibited a slightly more homogeneous distribution of grain-like features and a smoother surface. The reduction in the bright agglomerate phenomenon indicates a reduced surface roughness and higher film continuity. Such morphology is usual for layered MoS₂, where the interactions of van der Waals (vdW) forces allow compact growth and planar stacking.²¹

The current–voltage (I – V) characteristics of the proposed dual-heterojunction (n-MoS₂/p-CuO/Si and p-CuO/n-Si) under both dark and light settings are shown in Fig. 3a. An unsymmetrical nonlinear diode-like conduct is observed with a clear

rectification profile under both dark and illuminated states with respect to the utilized wavelength. In detail, the heightened carriers' diffusion and their recombination with injected charge result in dark current suppression. As the applied bias increases, the diffusion current dominates, leading to a significant dark current growth; in this case, the depletion region becomes thinner while the built-in electric field is nearly debilitated, which, in turn, allows diffusion current domination. The rectification ratios were found to be 10.06 and 7.83, respectively, at 5 V under dark conditions for the named heterojunctions; the attained variation in the rectification ratio could be because of the alteration in carrier recombination as well as defect states at different interfaces. Such interface quality enhancement, in the case of n-MoS₂/p-CuO/Si, may result in higher carrier transport phenomenon and subsequently higher rectifying singularity.²² It is worth pointing out that similar behavior was noted at different incident wavelengths with respect to the considered heterojunction. Further, the ideality factor (n) was estimated in accordance with Cheung's model, considering a semi-linear plot of the forward dark current²³ (see inset of Fig. 3a). The proposed heterojunctions exhibited n values of 1.7 and 2.1 for n-MoS₂/p-CuO/Si and p-CuO/n-Si, respectively. Such a deviation from $n = 1$ suggests the presence of interface recombination and/or trap-assisted tunneling effects. Fig. 3b and c show the wavelength-dependent photoresponse of the proposed junctions as a function of time-resolved profiles under a 3 V applied bias. Owing to the narrow bandgap of the utilized active layers (CuO and MoS₂), the fabricated heterojunctions exhibited rather broad responsive detection capability. However, two dominant wavelengths (625 and 720 nm) were identified for the n-MoS₂/p-CuO/Si and p-CuO/n-Si heterojunctions, respectively, corresponding to efficient photoinduced carrier generation under the specific incident wavelengths, the absorption edge of the active layers as pre-demonstrated in Fig. 1(c) and (d); this in turn validates the dual function nature of the proposed design. The distinguished photoresponse at 808 nm, in the case of p-CuO/n-Si heterojunction, is mainly due to the window effect of the utilized Si wafer.²⁴ Such wavelength-dependent behavior could

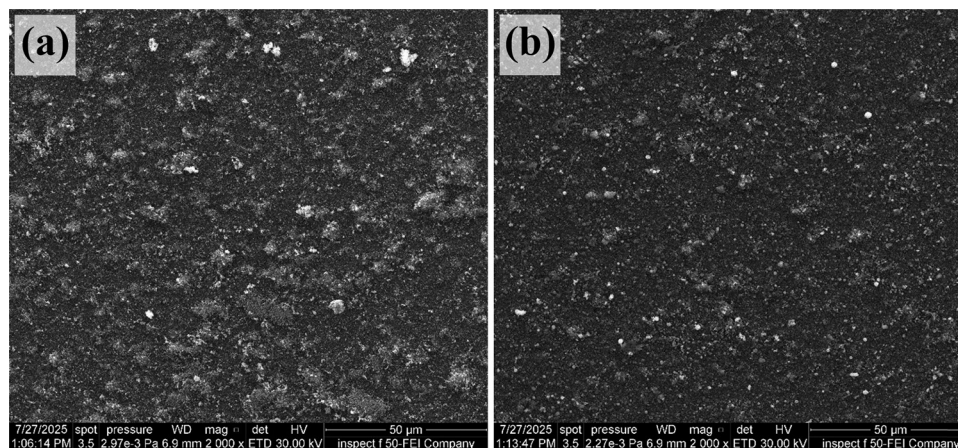


Fig. 2 FE-SEM images of the deposited (a) CuO and (b) MoS₂ layers; scale bar of 50 μm .



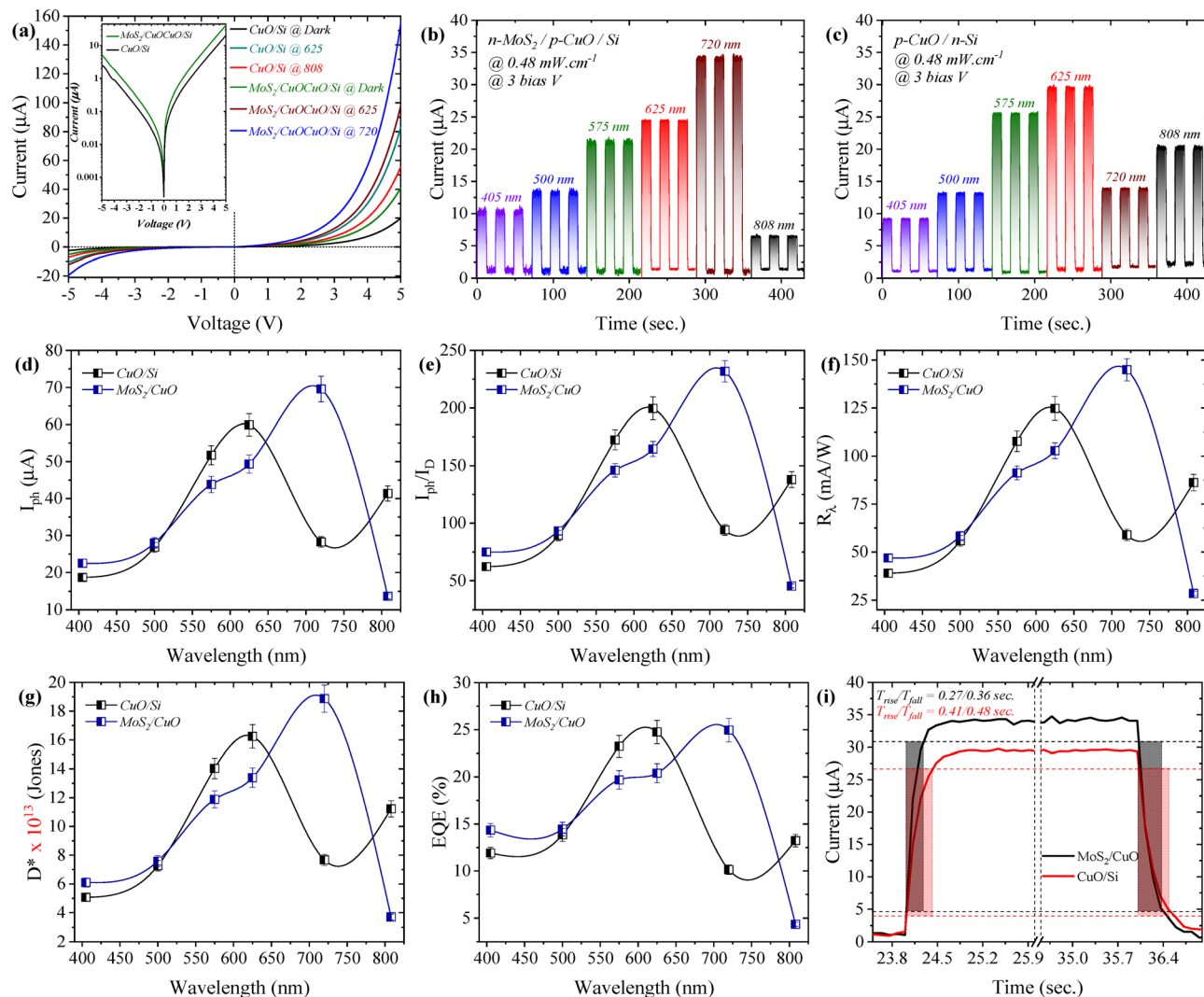


Fig. 3 Photo-responsive analysis of the fabricated n-MoS₂/p-CuO/Si and p-CuO/n-Si photodetectors at a 3 V bias: (a) *I*-*V* characteristics, with the inset of semi-log dark *I*-*V*, (b) time-based profile of n-MoS₂/p-CuO/Si, (c) time-based profile of p-CuO/n-Si, (d) *I*_{ph}, (e) *I*_{ph}/*I*_D, (f) *R*_λ, (g) *D*^{*}, (h) EQE, and (i) response/recovery time of CuO/Si and MoS₂/CuO.

be clearly indicated from the *I*_{ph} profile (Fig. 3c). Four critical figures-of-merit are demonstrated in Fig. 4e-h: light-to-dark ratio [*I*_{ph}/*I*_D], responsivity [*R*_λ = *I*_{ph} - *I*_D/*P*_{in}], detectivity [*D*^{*} = *R*_λ*A*^{1/2}/(2*eI*_D)^{1/2}], and external quantum efficiency [EQE = (*I*_{ph}/*e*)(*P*_{in}/*hν*)].^{25,26} Specifically, *R*_λ (Fig. 3f) demonstrated values of 0.144 and 0.124 A W⁻¹ at 720 and 625 nm for n-MoS₂/p-CuO/Si and p-CuO/n-Si, respectively, while *D*^{*} values of 1.9 × 10¹⁴ and 1.6 × 10¹⁴ Jones (Fig. 3g) and EQE of 25 and 24.7 (Fig. 3h) were obtained at the same addressed parameters, respectively. The proposed geometries exhibited response/recovery times (Fig. 3i) of 0.27/0.36 and 0.41/0.48 s, respectively, indicating faster responses compared to the recovery; these were considered under incident wavelengths of 720 and 625 nm, respectively. This, in turn, strongly indicates a faster generation phenomenon compared to recombination.^{27,28} The attained sub-second rise/fall time, in both junctions, could be attributed to the interfacial dynamics as well as carrier trapping/de-trapping within the proposed dual-heterojunction. In detail, both PLD-based CuO and MoS₂ layers

are polycrystalline (Fig. 1b) along defect states and grain boundaries, which, in turn, causes a localized carrier phenomenon along with prolonged lifetime. Additionally, such structural imperfections might also induce defects/traps, which directly influence the response speed.^{29,30} Results evidently suggest that the performance of n-MoS₂/p-CuO/Si outweighs that of p-CuO/n-Si due to the ability of the n-MoS₂/p-CuO heterojunction in reducing the defect states and/or carrier recombination.

The light intensity-dependent responses of the proposed junctions at the optimum wavelengths and 3 V applied bias are presented in Fig. 4a and b; this investigation was conducted in terms of the time-resolved characteristics. The considered junctions responded linearly to the light-intensity increment, with an *R*² value close to unity (*R*² ≈ 1). A more detailed examination was conducted into this behavior, considering the *I*_{ph}/*I*_D ratio (Fig. 4c), where the considered heterojunction demonstrated linear correlation to the utilized intensity at different wavelengths, suggesting a dynamic role of the active layers.



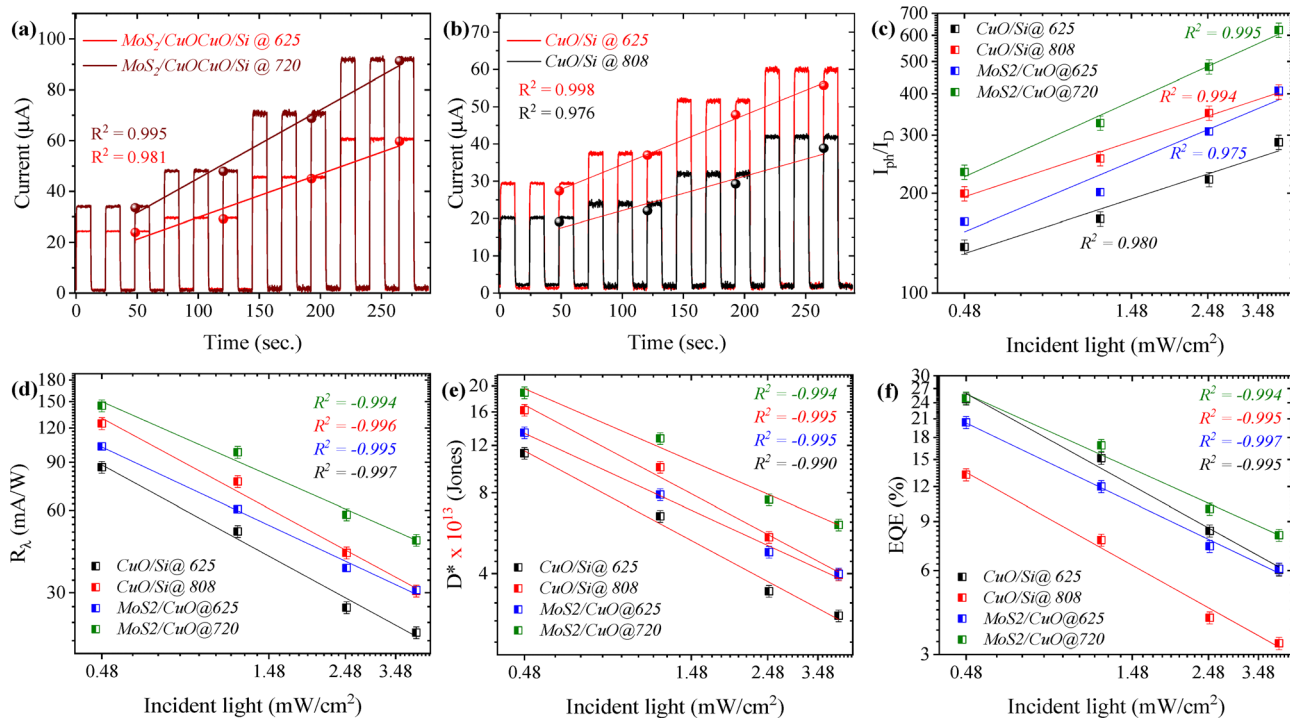


Fig. 4 Intensity-dependent profile: time characteristics of (a) $n\text{-MoS}_2/p\text{-CuO}/\text{Si}$, (b) $p\text{-CuO}/n\text{-Si}$. (c) I_{ph}/I_D , (d) R_{λ} , (e) D^* , and (f) EQE values achieved at a 3 V applied bias.

Other figures-of-merit, including R_{λ} , D^* , and EQE, exhibited clear inverse trends, as shown in Fig. 4d–f. These characteristics tend to saturate as the incident intensity increases; in detail, R_{λ} decreased from 0.144 to 0.047 A W^{-1} as the light intensity, for $n\text{-MoS}_2/p\text{-CuO}/\text{Si}$ at a wavelength of 720 nm, increased from 0.48 to

4 mW cm^{-2} . Such an occurrence is mainly because the relationship between R_{λ} and P_{in} is inversely proportional ($R_{\lambda} \propto P_{\text{in}}^{-1}$),³¹ while D^* and EQE are mathematically dependent on R_{λ} . Irrespective of the incident wavelength, $n\text{-MoS}_2/p\text{-CuO}/\text{Si}$ demonstrated higher figures-of-merit than $p\text{-CuO}/n\text{-Si}$.

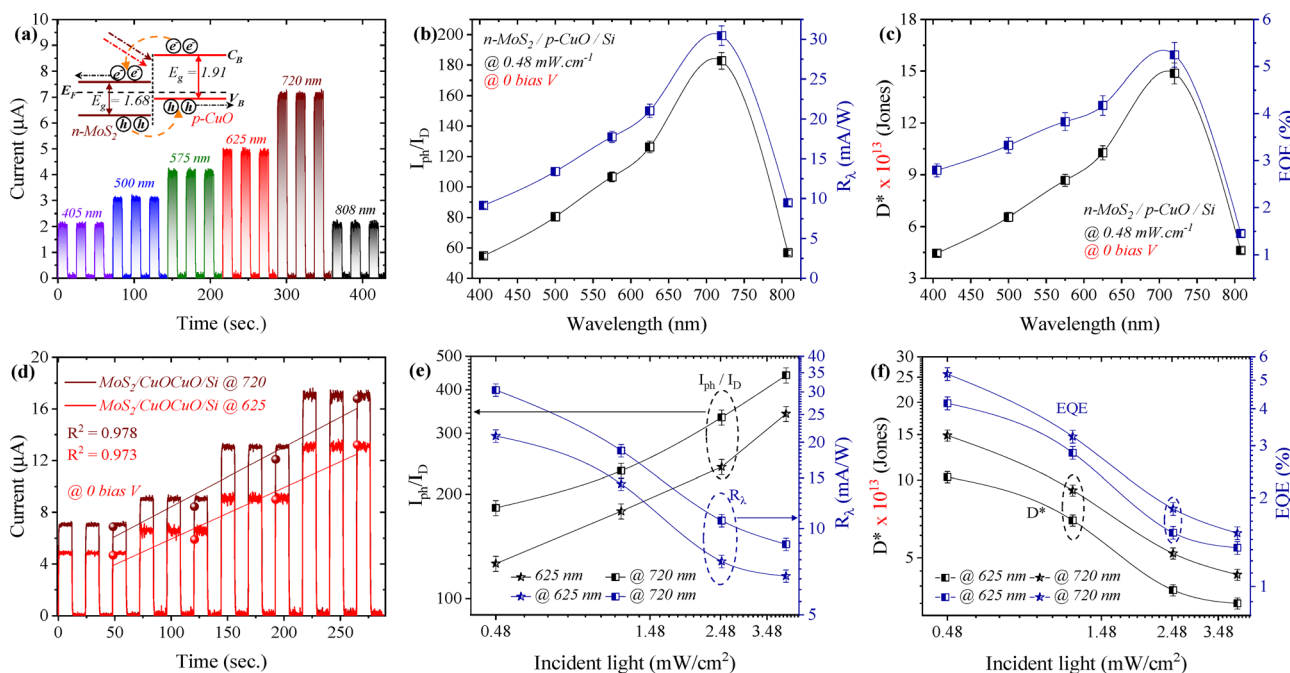


Fig. 5 Self-biased Vis-NIR configuration. Wavelength-based profiles: (a) time-resolved characteristics, (b) I_{ph}/I_D and R_{λ} , (c) D^* and EQE. Intensity-based profiles: (d) time-resolved characteristics, (e) I_{ph}/I_D and R_{λ} , and (f) D^* and EQE.



The responsive characteristics, under zero-bias configuration, of the investigated n-MoS₂/p-CuO/Si heterojunction were investigated systematically at 0.48 mW cm⁻² (Fig. 5). Within this particular configuration, the n-Si wafer is left electrically floated; this condition results in the emergence of a photogating phenomenon at the n-MoS₂/p-CuO/Si heterostructure's bottom interface. The detected effect not only induces a noticeable photovoltaic response within the proposed heterostructure by facilitating the separation of photogenerated carriers, where zero bias is applied, but also leads to a distinguished augmentation in photoconductive gain.⁷ The time-based light response, at zero applied bias, could be explained *via* the band alignment (see the inset of Fig. 5a). In detail, charge transport and carrier dynamics in the n-MoS₂/p-CuO/Si heterojunction are governed by specific aspects, including defect concentration, interface state, and band alignment. Particularly, light-induced electron excitation results in so-called band-bending, which, in turn, increases the built-in electric field and subsequently smooths the photo-generated carriers' separation and collection. Incident light induces additional band-bending (see the inset of Fig. 5a), thereby enhancing the electron-hole separation and strengthening the internal electric field, which, in turn, allows more efficient carrier extraction. Therefore, the attained band alignment suppresses the electron-hole recombination and increases the carrier collection efficiency at zero applied bias.

The light-response characteristics exhibited a well-oriented wavelength-dependent profile, indicating the good performance of the proposed heterostructure in the self-driven mode (Fig. 5a), with a considerably low dark current of 90 nA; the fluctuation in the measured dark current mainly resulted from the resolution limitation of our setup. The self-driven feature was further inspected in terms of the related figures-of-merit. The demonstrated figures-of-merit (Fig. 5b and c) delivered a considerable R_{λ} of 30.50 mA W⁻¹, D^* of 1.5×10^{14} Jones, and EQE of 5.25 at zero applied bias, indicating the self-driven nature of the proposed geometry. In terms of light intensity dependence, the n-MoS₂/p-CuO/Si heterojunction was also investigated at zero applied bias under illumination at the optimum wavelengths (625 and 720 nm); the light response (Fig. 5d) demonstrated a linear increase, which indicates the superior light sensitivity of the proposed junction. The I_{ph}/I_D showed a linear increase as a function of light intensity increment (Fig. 5e, black curves). Other figures-of-merit showed a linear decrease as a function of incident light intensity increase (Fig. 5e and f).

4. Conclusion

A self-driven optoelectronic dual-heterojunction (n-MoS₂/p-CuO/n-Si) was effectively fabricated through consecutive PLD techniques, integrating spatially decorated films. The photodetector exhibited a multi-spectrum response, under zero biasing, at wavelengths of 625, 720 and 808 nm, crossing both Vis and NIR regions. At 0 V bias, the fabricated n-MoS₂/p-CuO/

n-Si heterostructure exhibits R_{λ} of 21.04/30.50 mA W⁻¹ and D^* of $1.0 \times 10^{14}/1.5 \times 10^{14}$ Jones at incident wavelengths of 625/720 nm, confirming the self-driven feature. Such metrics were enhanced at an applied bias of 3 V, with R_{λ} values of 0.144 and 0.124 A W⁻¹ for n-MoS₂/p-CuO/Si and p-CuO/n-Si, respectively, attributed to lower interface recombination and improved carrier separation. Additionally, time-resolved features revealed rise/fall times of 0.27/0.36 and 0.41/0.48 s, respectively, for the addressed heterojunctions.

Conflicts of interest

There are no conflicts to declare.

Data availability

All data used are presented in the manuscript.

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