

Supporting Information

Boosting Two-dimensional MoS₂/CsPbBr₃

Photodetectors via Enhanced Light Absorbance and Interfacial Carrier Separation

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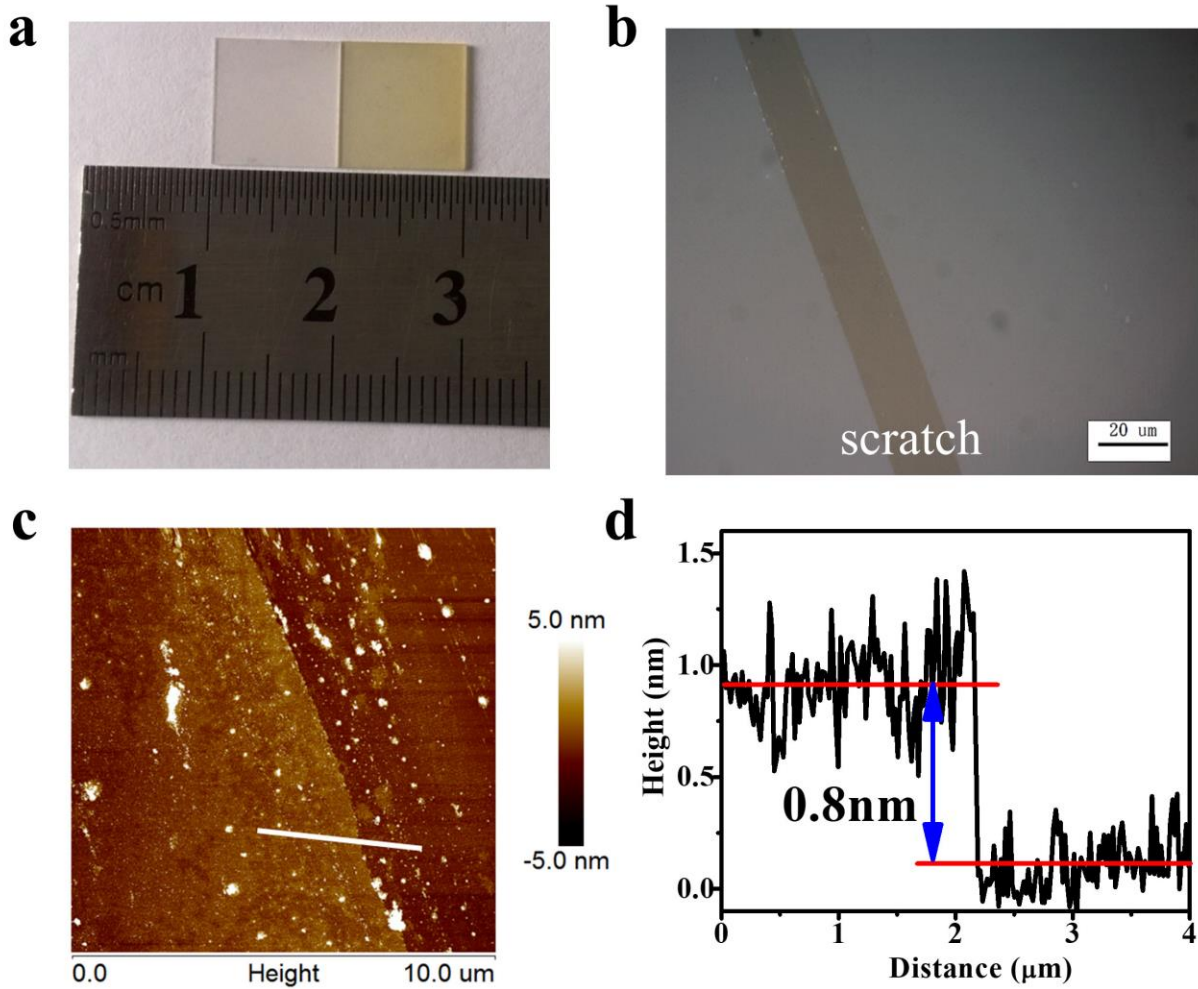


Figure S1. Large-area uniformity of the synthesized MoS₂ film. a) Photograph of as-grown MoS₂ monolayer, left: bare sapphire substrate, Right: fully covered MoS₂ monolayer on sapphire substrate. b) Optical image of the MoS₂ monolayer. c) AFM image of the obtained MoS₂ monolayer. d) Height profile corresponding to the white line shown in c).

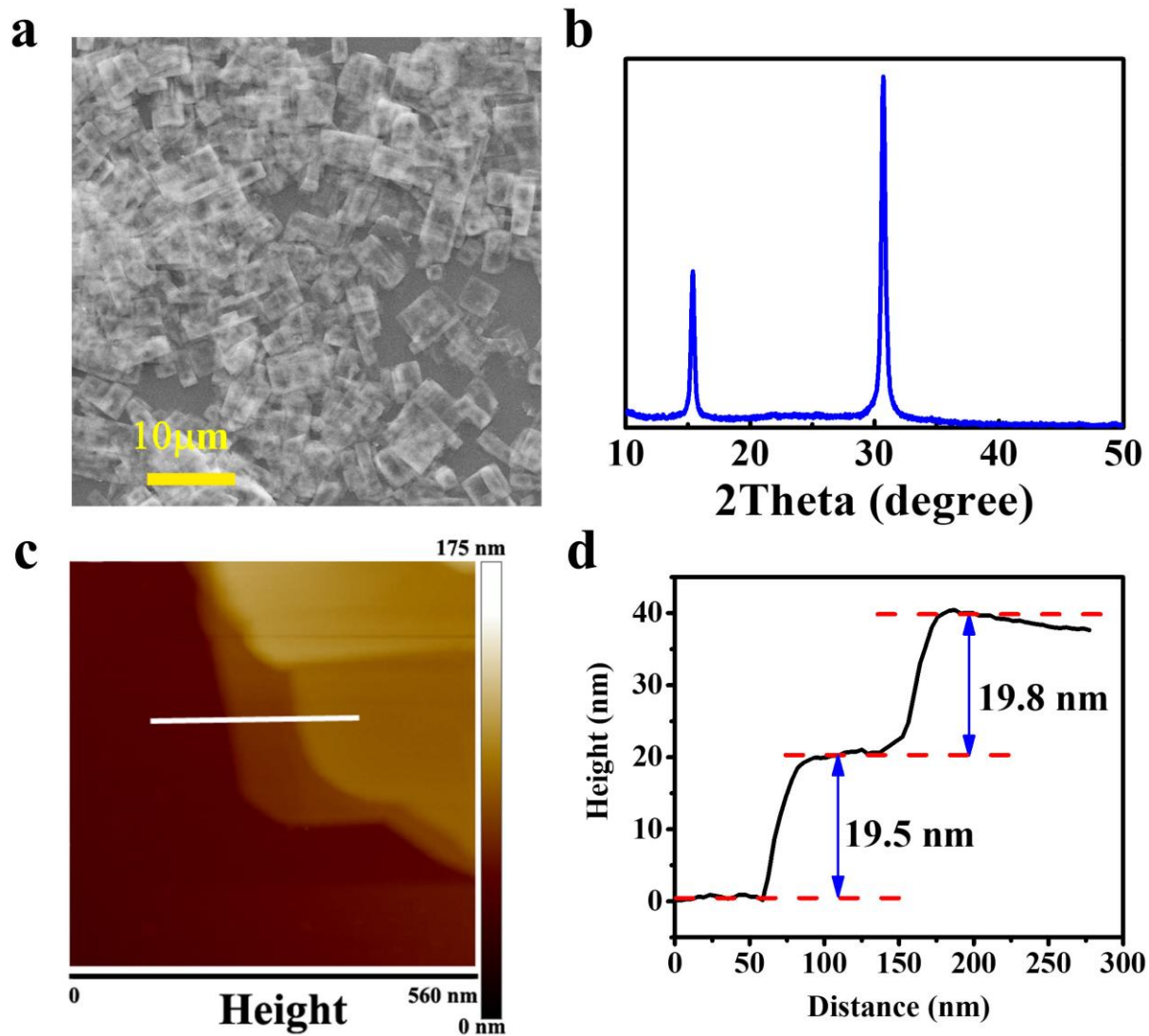


Figure S2. a) SEM image and b) XRD pattern of the CsPbBr₃ nanosheets. c) AFM image of the obtained CsPbBr₃ nanosheets. d) Height profile for the white line shown in c).

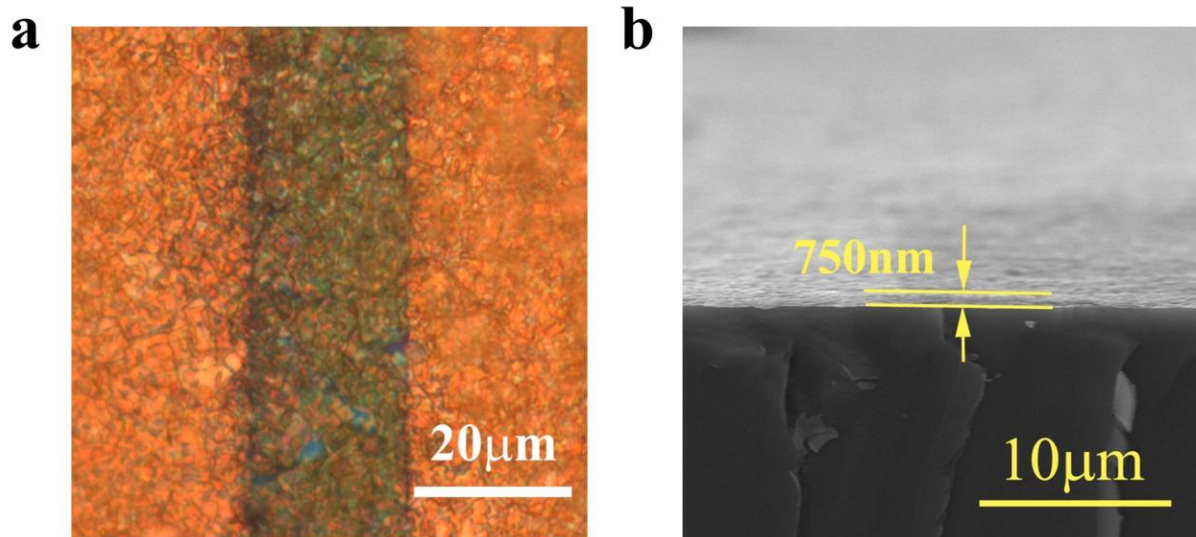


Figure S3. a) Optical image of the hybrid $\text{MoS}_2/\text{CsPbBr}_3$ photodetector. b) Cross-sectional SEM image of the coated CsPbBr_3 nanosheets.

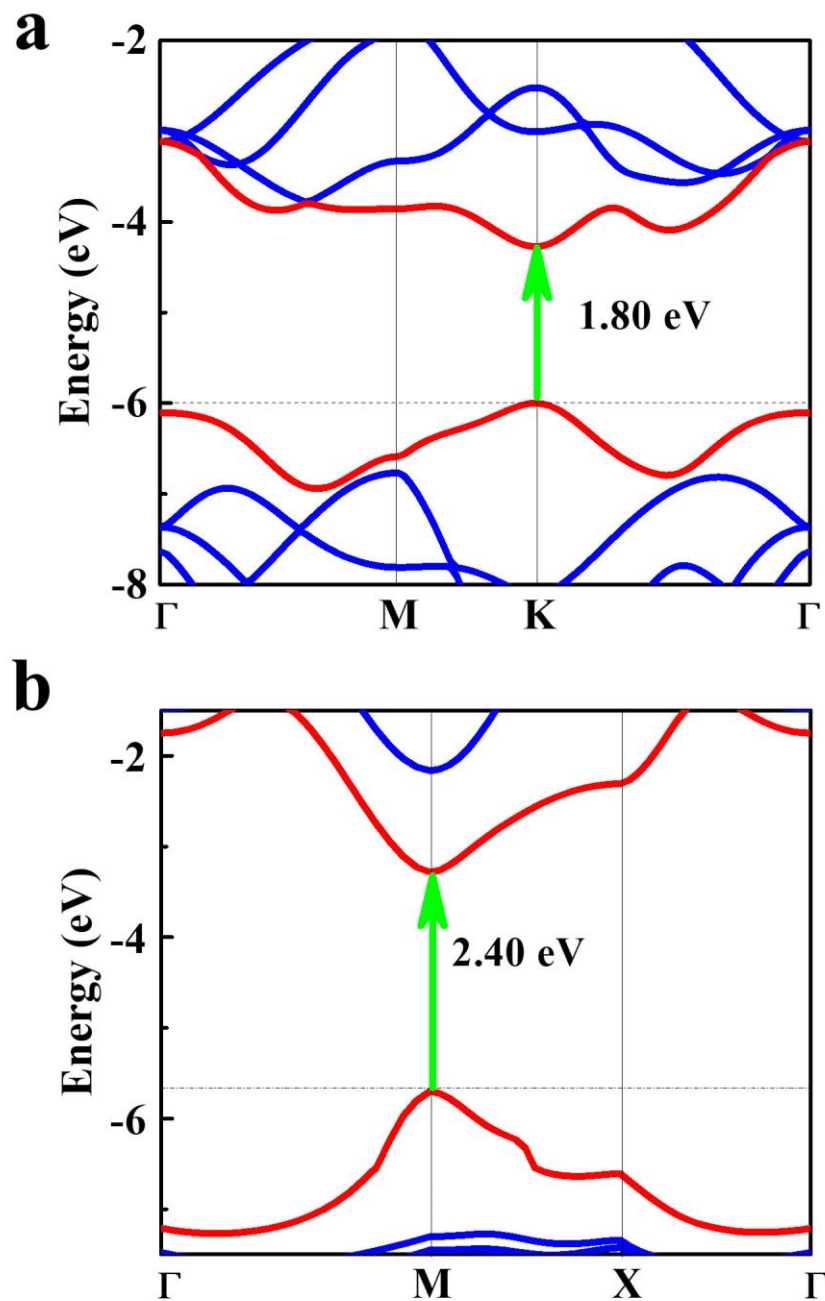


Figure S4. Calculated band structure of a) MoS₂ and b) CsPbBr₃.

Our calculated results (MoS₂ with band gap 1.8 eV, and CsPbBr₃ with band gap 2.4 eV) agree well with those experimentally determined values^{S1-3}.

PL lifetime of pure CsPbBr₃ and MoS₂/CsPbBr₃ perovskite structure

Figure 3b shows the time-resolved PL decay transients measured at 515 nm for pure CsPbBr₃ and hybrid MoS₂/CsPbBr₃.

The experimental curves of the PL decay could be fitted well using the biexponential functions:

$$A(t) = A_0 + A_1 * \exp\left(\frac{-t}{\tau_1}\right) + A_2 * \exp\left(\frac{-t}{\tau_2}\right) \quad (S1)$$

The average lifetime (τ) is calculated from

$$\tau = \frac{\sum A_i \tau_i^2}{\sum A_i \tau_i} \quad (S2)$$

Table S1. PL lifetimes of CsPbBr₃ and MoS₂/CsPbBr₃.

	A ₁	τ_1 (ns)	A ₂	τ_2 (ns)	R ²	τ (ns)
CsPbBr ₃	5.70	2.06	0.81	13.33	0.987	7.46
MoS ₂ /CsPbBr ₃	30.54	1.42	0.66	4.99	0.992	1.67

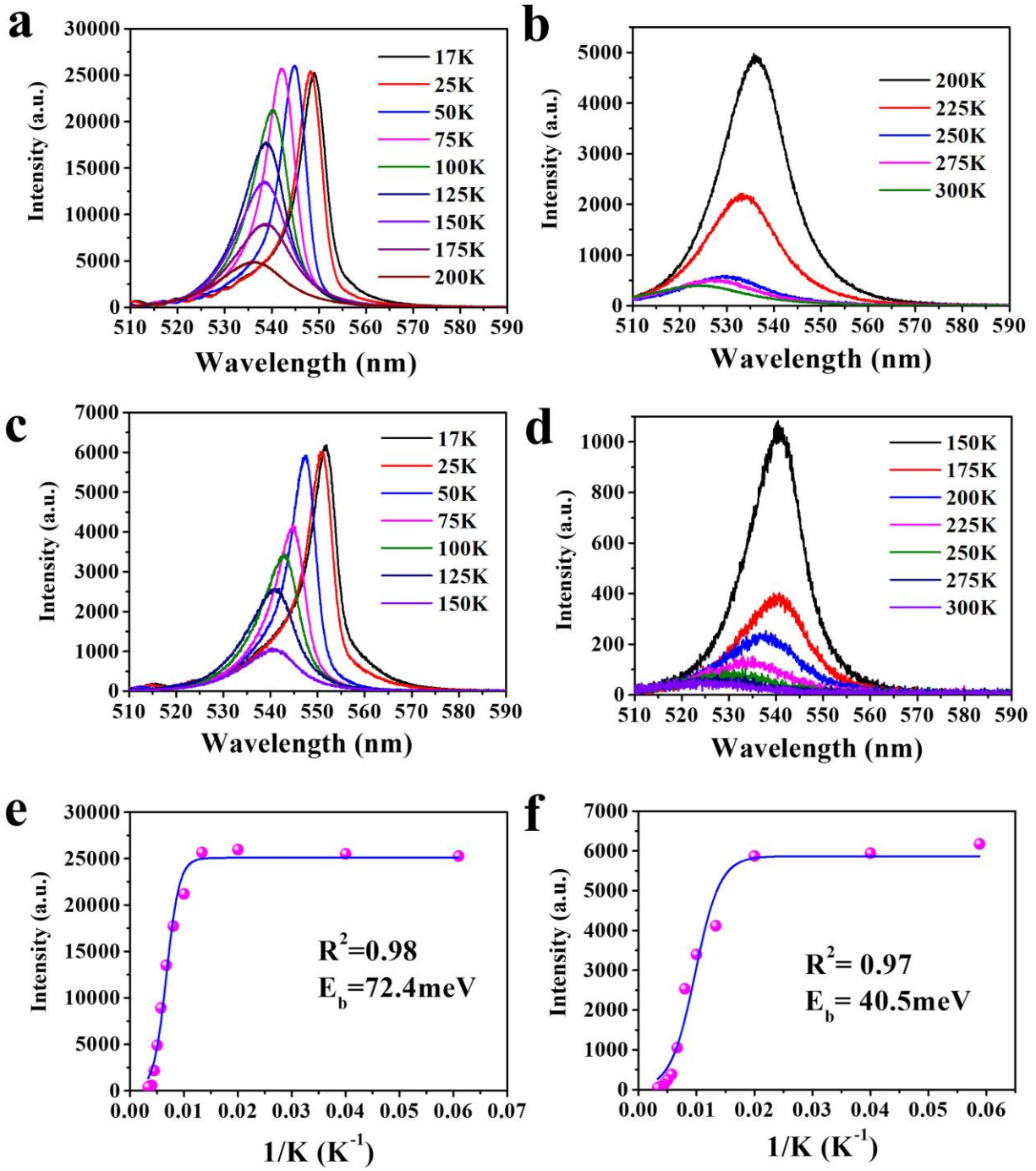


Figure S5. Temperature dependent PL spectra of a),b) CsPbBr₃, and c),d) MoS₂/CsPbBr₃. (e) Temperature dependent PL intensity of CsPbBr₃. f) Temperature dependent PL intensity of MoS₂/CsPbBr₃.

As shown in Figure S5, the PL intensity of the CsPbBr₃ can decrease with increasing the temperature. The exciton binding energy can be calculated by the following equation:

$$I(T) = \frac{I_0}{1 + Ae^{-E_b/k_b T}} \quad (S3)$$

where, I_0 is the PL intensity at 0 K, k_b is the Boltzmann constant, E_b is the exciton binding energy.

The calculated exciton binding energy of CsPbBr₃ and MoS₂/CsPbBr₃ are 72.4 meV and 40.5 meV, respectively. The decrease of the exciton binding energy indicates that the generated excitons in the hybrid structure tend to separate more easily during the device operation process, which is beneficial to increase the photocurrent of the photodetector based on CsPbBr₃/MoS₂.

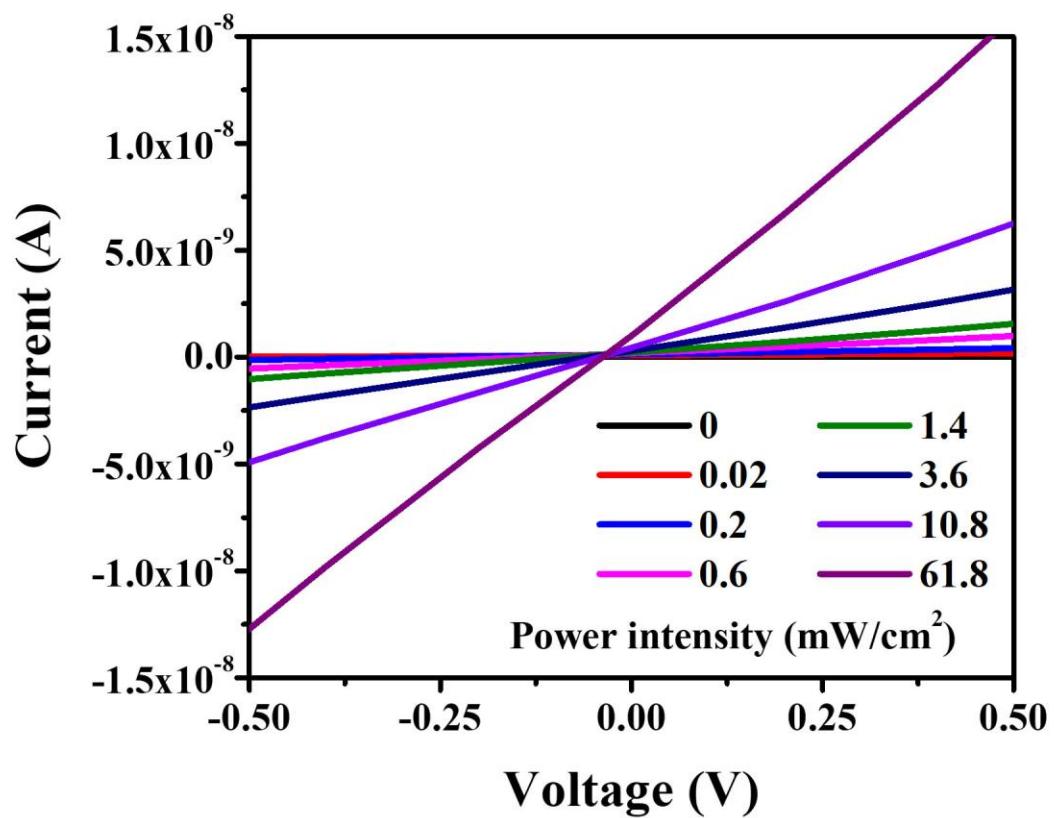


Figure S6. Drain-source characteristic of MoS₂/CsPbBr₃ device in the darkness and under different illumination intensities.

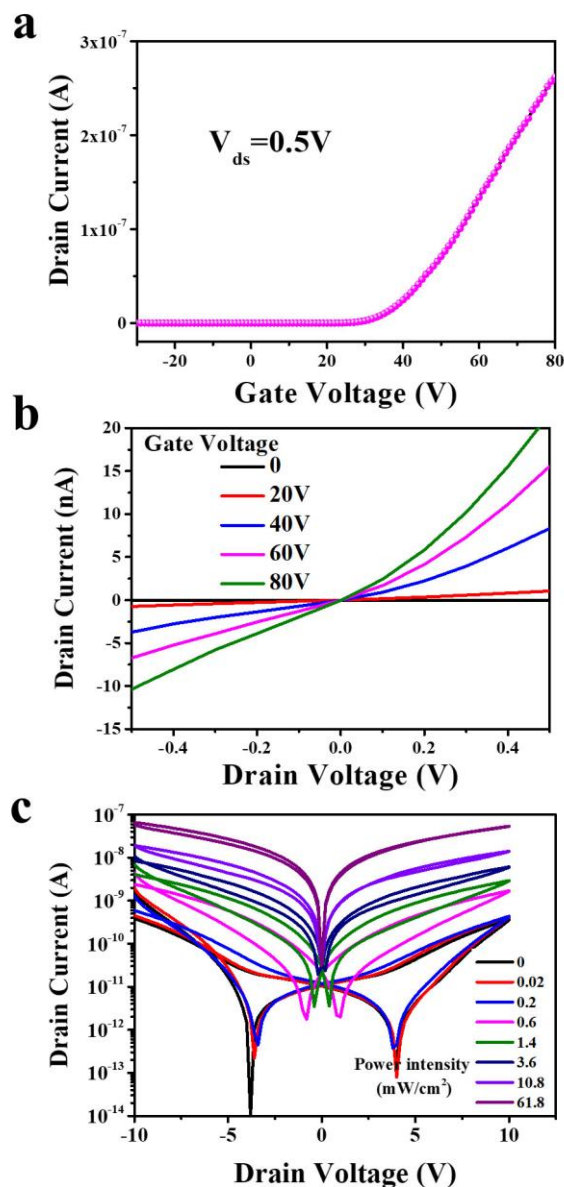


Figure S7. Electrical characteristics of MoS₂ transistor. a) Transfer characteristics. b) Output characteristics with different gate voltage. c) Drain-source characteristic in darkness and under different illumination intensities.

Due to the wet-transfer process, water or organic residues could exist on the surface of MoS₂ and at the MoS₂/SiO₂ interface, leading to the reduction of conductivity and large hysteresis of the devices. This is in consistence with previous reports in literature.^{S4-6}

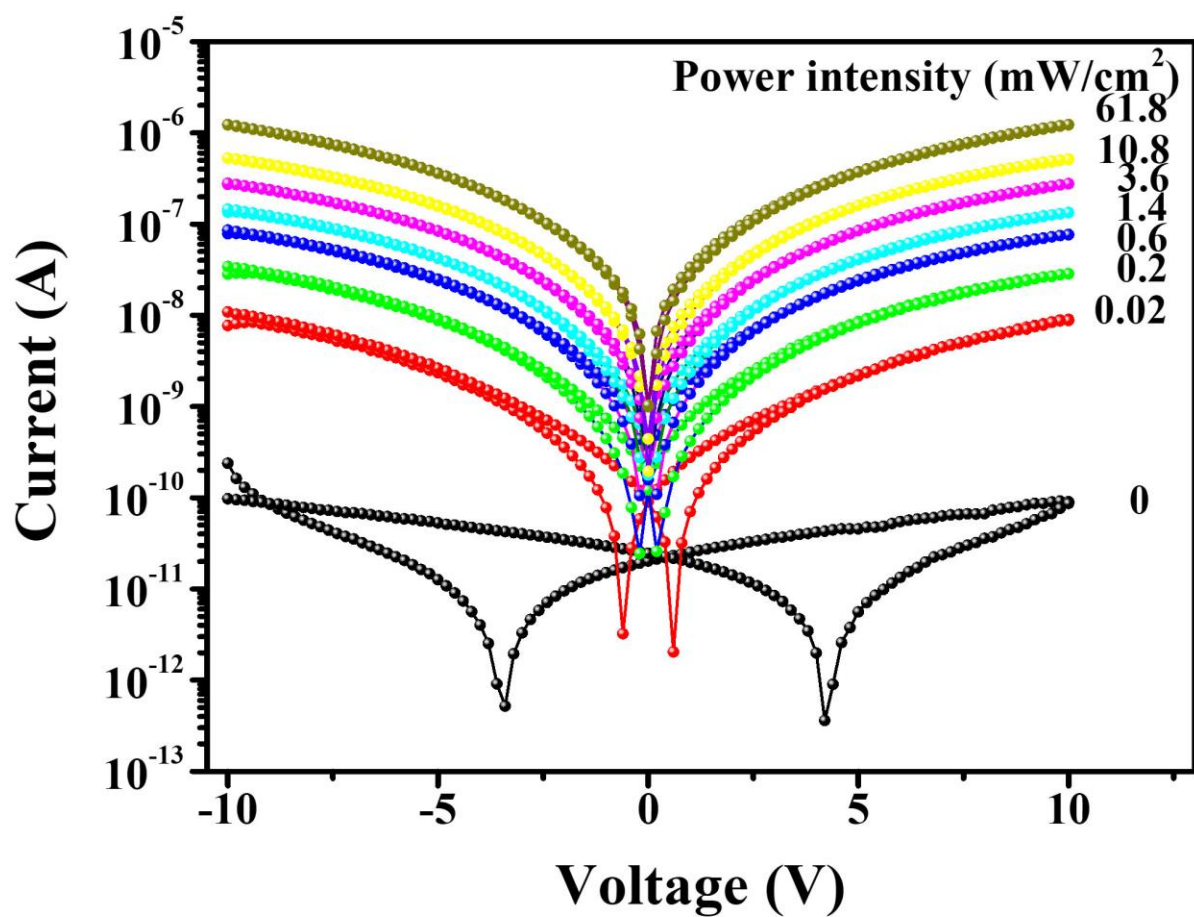


Figure S8. Drain-source characteristic of CsPbBr₃ photodetector in darkness and under different illumination intensities.

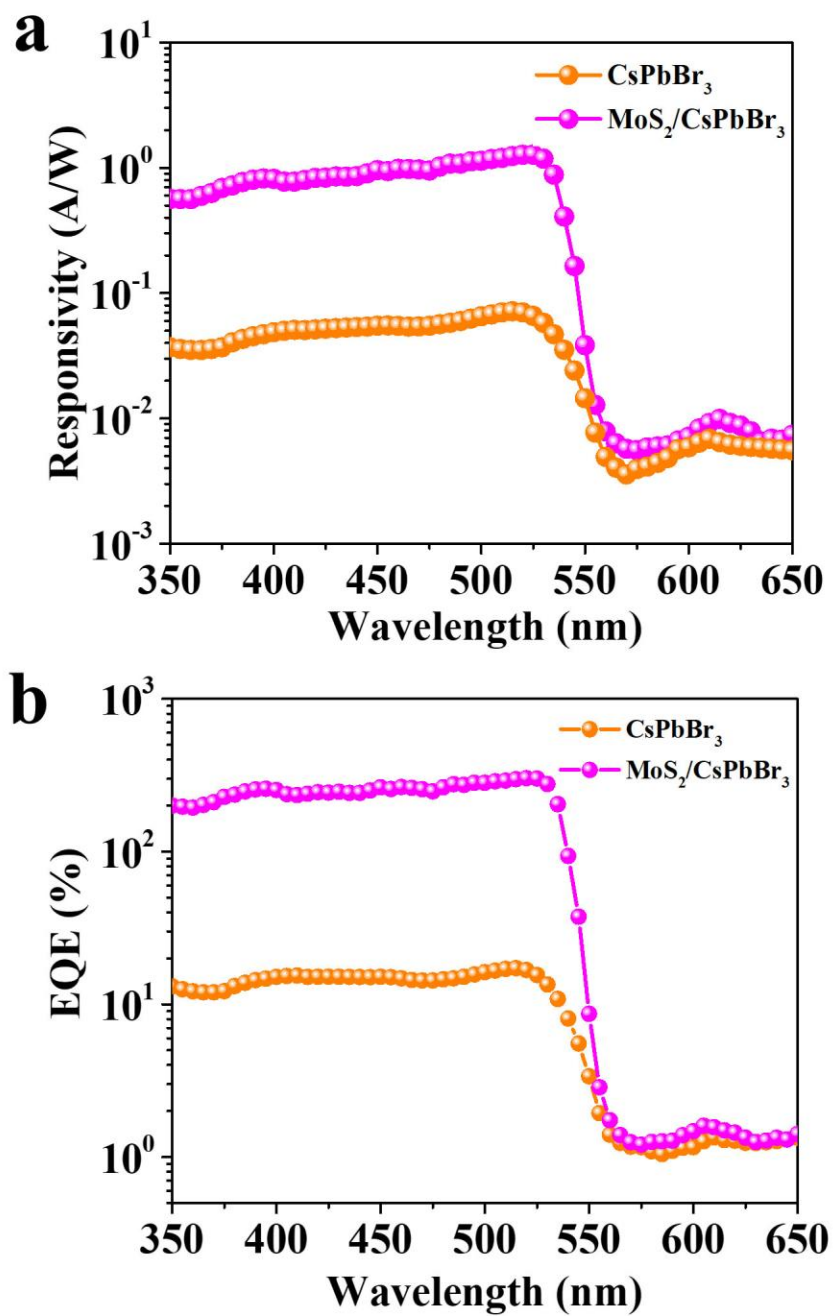


Figure S9. a) Photoresponsivity and b) EQE of CsPbBr₃ photodetectors with or without MoS₂ as a function of illumination wavelength.

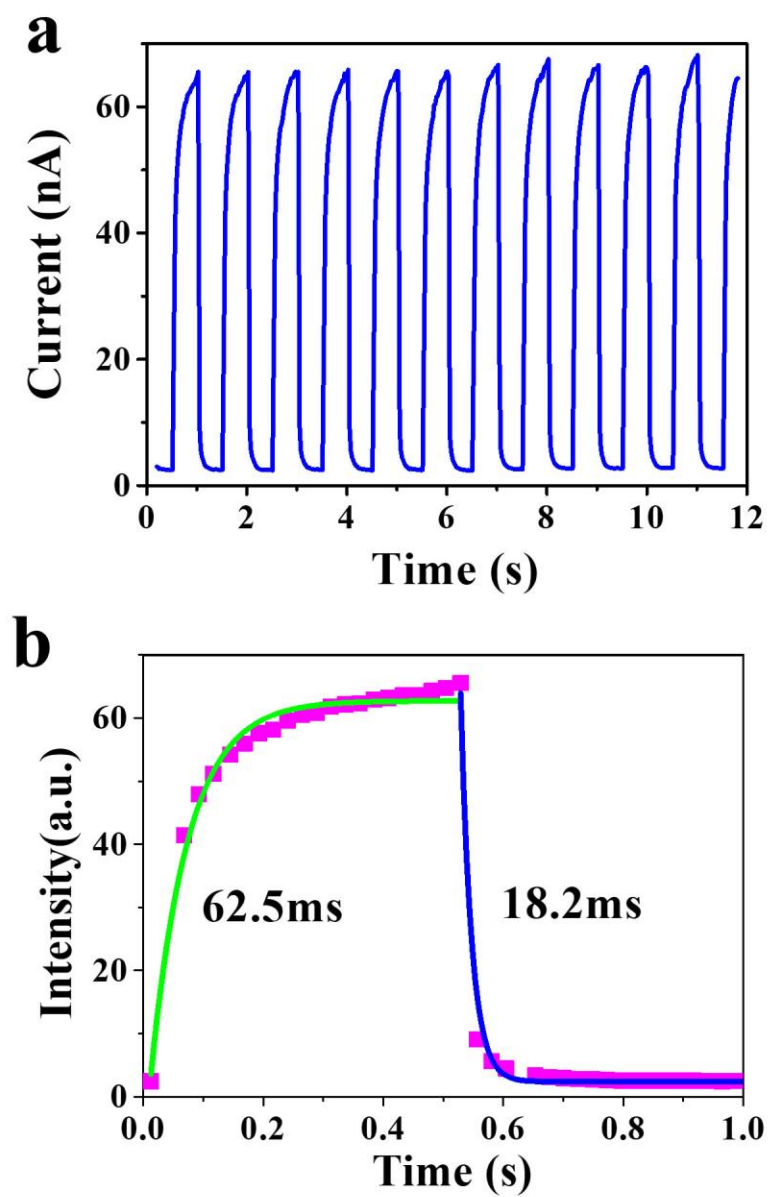


Figure S10. Time-resolved photoresponse of the CsPbBr₃ photodetector under 442 nm laser illumination. a) Photoswitching behavior of the hybrid device at different voltages and incident optical power intensity. b) Temporal photocurrent response of the hybrid device with rising time (62.5 ms) and decay time (18.2 ms).

Table S2. Performance comparing of photodetectors on 2D/Perovskite.

Device structure	Incident	Responsivity (A/W)	Detectivity (J)	Rise /fall	Ref.
MoS ₂	150pW @561nm	880		4s /9s	S7
MoS ₂	4mW cm ⁻² @633nm	0.12	10 ¹¹	-	S8
WSe ₂	10nW @532nm	0.15	-	-	S9
InSe	0.66 mW cm ⁻² @405nm	12.3	5.47×10 ¹⁰	50ms /60ms	S10
CH ₃ NH ₃ PbI ₃ /MoS ₂ /APTES	4.63pW @520nm	2.11×10 ⁴	1.38×10 ¹⁰	6.17s /4.5s	S11
CH ₃ NH ₃ PbI ₃ /1T-MoS ₂	0.14mW @500nm	3.3×10 ⁵	7×10 ¹¹	0.45s /0.75s	S12
CH ₃ NH ₃ PbI ₃ /2H- MoS ₂	0.14mW @500nm	142	2.6 × 10 ¹¹	25ms /50ms	S12
MAPbI ₃ /Graphene	3.3 pW @633nm	2.6×10 ⁶	-	55s /75s	S13
CH ₃ NH ₃ PbI ₃ /Graphene	0.002 mW @515nm	115	3×10 ¹²	-	S14
CH ₃ NH ₃ PbPbBr ₂ I/Graphene	223 μW @405nm	6.0×10 ⁵	-	120ms /750ms	S15
CH ₃ NH ₃ PbI ₃ /WSe ₂	70 mW cm ⁻² @532nm	1.1×10 ⁵	2.2×10 ¹²	2.1s /3s	S16
CH ₃ NH ₃ PbI ₃ /Graphene	1 μW @520nm	180	10 ⁹	87ms /540ms	S17
CsPbBr _{3-x} I _x /Graphene	0.07 mW cm ⁻² @405nm	8.2×10 ⁸	2.4×10 ¹⁶	0.81s /3.65s	S18
CH ₃ NH ₃ PbI ₃ /WS ₂	0.2 μW cm ⁻² @505nm	17	2×10 ¹²	2.7ms /7.5ms	S6
CsPbBr ₃	4.65 mW cm ⁻² @532nm	0.00471	16.84×10 ⁸	0.2 ms /1.3 ms	S2
CsPbBr ₃ /Au	4.65 mW cm ⁻² @532nm	0.01004	4.56×10 ⁸	0.2 ms /1.2 ms	S2
CsPbBr ₃ /MoS ₂	20μW cm ⁻² @442nm	4.4	2.5×10 ¹⁰	0.72ms /1.01ms	This work

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