Coupling between Pyroelectricity and Built-In Electric Field Enabled Highly Sensitive Infrared Phototransistor Based on InSe/WSe₂/ P(VDF-TrFE) Heterostructure

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ABSTRACT: The assorted utilization of infrared detectors induces the demand for more comprehensive and high-performance electronic devices that work at room temperature. The intricacy of the fabrication process with bulk material limits the exploration in this field. However, two-dimensional (2D) materials with a narrow band gap opening aid in infrared (IR) detection relatively, but the photodetection range is narrowed due to the inherent band gap. In this study, we report an unprecedented attempt at the coordinated use of both 2D heterostructure (InSe/ WSe₂) and the dielectric polymer (poly(vinylidene fluoridetrifluoroethylene), P(VDF-TrFE)) for both visible and IR photodetection in a single device. The remnant polarization due to the ferroelectric effect of the polymer dielectric enhances the



photocarrier separation in the visible range, resulting in high photoresponsivity. On the other hand, the pyroelectric effect of the polymer dielectric causes a change in the device current due to the increased temperature induced by the localized heating effect of the IR irradiation, which results in the change of ferroelectric polarization and induces the redistribution of charge carriers. In turn, it changes the built-in electric field, the depletion width, and the band alignment across the p-n heterojunction interface. Consequently, the charge carrier separation and the photosensitivity are therefore enhanced. Through the coupling between pyroelectricity and built-in electric field across the heterojunction, the specific detectivity for the photon energy below the band gap of the constituent 2D materials can reach up to 10^{11} Jones, which is better than all reported pyroelectric IR detectors. The proposed approach combining the ferroelectric and pyroelectric effects of the dielectric as well as exceptional properties of the 2D heterostructures can spark the design of advanced and not-yet realized optoelectronic devices.

KEYWORDS: pyroelectricity, 2D materials, $InSe/WSe_2$, van der Waals heterostructure, P(VDF-TrFE), ferroelectric gating, an infrared detector

INTRODUCTION

Infrared (IR) photodetectors have attracted attention among researchers due to their extensive applications in remote sensing, optical communication, biomedical imaging, military surveillance, and chemical sensing.^{1,2} According to the wavelength range, the IR (0.8–30 μ m) region can be divided into the near-infrared (NIR), short-wave infrared (SWIR), mid-wave infrared (MWIR), long-wave infrared (LWIR), and far-infrared (FIR) range.¹ Due to the thermal noise and complicated device fabrication steps involved with IR photodetectors based on three-dimensional (3D) crystal structures so far, studies were focused on ultrathin two-dimensional (2D) materials for IR detection.^{3,4} Mentioning 2D materials, graphene was the prime choice to experiment for IR photodetection by inducing a desirable band gap through doping.^{1,5} Xia et al. demonstrated a graphene-based photodetector at 80 V gate bias for detecting 1.55 μ m wavelength IR irradiation.⁶ Consecutively, the next widely studied 2D transition metal dichalcogenide (TMD) MoS₂ was assembled with other TMDs to form a type II heterostructure (MoS₂/WS₂, MoS₂/MoTe₂) where the interband transition aids in IR absorption.^{7,8} The MoTe₂/graphene/SnS₂ heterostructure to detect SWIR rays was demonstrated by Li et al.⁹ Other 2D semiconductors like black phosphorous (0.3–2.0 eV), tellurium (0.3–1.0 eV), platinum diselenide (PtSe₂, 0–1.2

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eV), and palladium diselenide (PdSe₂, 0.1–1.2 eV) were studied to show promising results on IR photodetection.^{1,3,10–12} The PdSe₂/MoS₂ heterostructure was reported by Long et al. for LWIR detection.³ Besides assembling 2D layers, zero-dimensional (0D) materials like quantum dots, nanoparticles, and one-dimensional (1D) carbon nanotubes (CNT) were also combined with 2D materials for IR detection.^{13–15} Hybrid structures like the graphene/CNT layer and MoS₂/HgTe quantum dots were also reported to detect the SWIR range.^{15,16}

Despite assuring outcomes with specific 2D materials and their hybrid structures, the range for IR photodetection is limited according to the inherent nature of the band gap of the material. In this case, if the thermal heating effect of IR rays is taken advantage of, the range for IR detection can be increased, which will widen the application areas as well. The heat caused by IR rays causes a change in temperature over the irradiated area to which certain materials exhibit a change in net polarization in the material due to the pyroelectric effect.¹⁷ Poly(vinylidene fluoride) (PVDF) is one such material to possess a pyroelectric effect.¹⁸ PVDF copolymerizes with trifluoroethylene (TrFE) to form P(VDF-TrFE), which is a ferroelectric polymer that exhibiting both pyroelectric and piezoelectric effects.¹⁹ Employing P(VDF-TrFE) as a dielectric to a device aids in IR detection by measuring the change in current caused by the change in dielectric polarization, and it can also be used as a ferroelectric transistor.

In this work, we will make an unprecedented attempt to demonstrate the coupling between pyroelectricity and a builtin electric field for a highly sensitive IR phototransistor consisting of P(VDF-TrFE) and a 2D van der Waals heterostructure to harvest both the optoelectronic property of 2D heterojunction for visible light photodetection and the pyroelectric effect of P(VDF-TrFE) for IR detection. Although there are few reports on ferroelectric gated 2D material transistors, there have been no studies on a pyroelectric gated 2D heterostructure. Here, we chose the type II InSe/WSe₂ heterostructure for our study, and there are limited reports on InSe-based heterostructures.^{20,21} InSe is an n-type semiconductor with a band gap of 1.27 eV for a thickness of more than 6 nm and an electron mobility of $10^3 \text{ cm}^2/(\text{Vs})$, whereas WSe₂ is a p-type semiconductor with a direct and indirect optical transition based on the number of layers and has a carrier mobility of $10^2 \text{ cm}^2/(\text{Vs})$.^{22–25} The InSe/WSe₂ heterostructure top-gated with P(VDF-TrFE) dielectric shows an ambipolar transfer behavior, and the ferroelectric polarization results in a relatively higher photoresponsivity ($\lambda = 633$ nm) in the polarized state (10^4 A W^{-1}) than the unpolarized state. During IR irradiation (λ = 980, 1064 nm) in the polarized state, due to the pyroelectric effect in P(VDF-TrFE), a significant change in current was observed for different laser power intensities, which is indicative of the enhanced IR detection. The underlying mechanism of enhanced photosensitivity arises from the existence of the built-in electric field across the heterojunction. Under IR irradiation, the increased temperature will induce the change of the PVDF polarization as well as the built-in electric field, which enables the alteration of depletion width, changes the band alignment across the heterojunction, and varies the charge carrier separation. As a result, the photosensitivity is enhanced, and the specific detectivity for the photon energy below the band gap of the constituent 2D materials can reach up to 10¹¹ Jones, which is better than all the reported pyroelectric IR detectors.

RESULTS AND DISCUSSION

Figure 1a represents the schematic illustration of the InSe/ WSe_2 device on a SiO₂/Si substrate with ferroelectric P(VDF-



Figure 1. Device structure on a solid-state substrate. (a) Schematic representation of the P(VDF-TrFE)-coated $InSe/WSe_2$ heterostructure on a SiO₂/Si substrate. (b) Optical microscope image of the assembled heterostructure after electrode deposition. (c) AFM image of the InSe/WSe₂ heterostructure; the inset shows the height profile indicating the thickness of InSe and WSe₂ layers individually. (d) Raman spectra were recorded on the individual InSe, WSe₂ layers, and the assembled InSe/WSe₂ heterojunction.

TrFE) film as the top dielectric layer. InSe and WSe₂ crystals were grown by the vertical Bridgman technique and chemical vapor transport (CVT) method, respectively. The growth procedures were detailed in our previous report.²⁶ The asgrown InSe crystal belongs to the $\hat{\beta}$ phase and has a hexagonal structure.^{26,27} The layered WSe₂ crystals also have a hexagonal crystal structure.²⁶ Both the crystals were exfoliated into ultrathin flakes by a mechanical exfoliation technique. The exfoliated WSe₂ flakes were transferred to the SiO₂/Si substrate with a poly(dimethylsiloxane) (PDMS) stamp. In the same manner, InSe flakes were precisely assembled on top of WSe₂ to form a heterojunction. The assembled heterostructure was coated with P(VDF-TrFE) solution after depositing electrodes and then annealed. The detailed device fabrication steps involved along with the schematic are given in Supplementary Note 1. The optical microscope image of the stacked InSe/WSe₂ heterojunction device is shown in Figure 1b. The atomic force microscopy (AFM) image with the height profile is represented in Figure 1c. The thicknesses of InSe and WSe₂ flakes were determined to be \sim 21 and \sim 19 nm, respectively. To obtain ambipolar or anti-ambipolar transport behavior in a heterostructure where comparable electron and hole contributions play a major role, it is necessary to have similar levels of intrinsic carrier density in both p- and n-type material.^{28,29} Otherwise, the asymmetry can lead to the domination of either of the carrier resulting in unipolar



Figure 2. Transfer characteristics of the ferroelectric gated transistor. (a) Transfer curves are measured at constant V_{ds} with P(VDF-TrFE) as the top-gated dielectric and the corresponding logarithmic plot is represented in panel (b). (c, d) Schematic illustration of the ferroelectric gated InSe/WSe₂ heterostructure transistor at two different polarization states.



Figure 3. InSe/WSe₂ heterostructure as a ferroelectric gated phototransistor. (a) $I_{ds}-V_{ds}$ characteristic curves of the device in the dark condition at the normal and polarized states. (b) $I_{ds}-V_{ds}$ curves measured at the ferroelectric polarized state under 633 nm laser illumination with different power intensities. (c, d) Responsivity and photogain of the device as a function of laser power intensity at two different ferroelectric polarized states.



Figure 4. Schematic of the pyroelectric detector in a ferroelectric gated $InSe/WSe_2$ heterostructure. (a) Representation of the pn junction with ptype WSe_2 and n-type InSe without ferroelectric gating. (b) Cross-sectional illustration of the pn junction with ferroelectric P(VDF-TrFE) topgating polarized in the upward direction. (c) Schematic depiction of a decrease in polarization in the device when irradiated with a NIR laser because of the pyroelectric effect.

transport behavior.^{28,29} So, to match the n-type carrier density in InSe, we used a multilayered p-WSe₂ nanoflake. The P(VDF-TrFE) film coating with the top-gate electrode is shown in Figure S2. The Raman spectrum of the individual InSe, WSe₂ layers, and their heterojunction is represented in Figure 1d. The strong vibrational modes at 115 (A_{1g}^1 mode), 178 (E_{1g}^2 mode), and 228 cm⁻¹ (A_{1g}^2 mode) in the separate InSe and the combined modes at 250 (E_{2g}^1 and A_{1g} mode) and 302 cm⁻¹ (B_{2g}^1 mode) in the individual WSe₂ seem to be superimposed in the InSe/WSe₂ heterojunction, which is in accordance with the previous reports.^{26,27}

Initially, before coating P(VDF-TrFE), the InSe/WSe₂ heterostructure exhibited anti-ambipolar behavior as per our previous report.²⁶ However, after coating P(VDF-TrFE) over the heterojunction, due to the doping effect and change in threshold voltages, the device exhibits ambipolar behavior.³⁰ The P(VDF-TrFE) solution was spin-coated at 3500 rpm for 20 s to obtain a thickness of 300 nm, which is well above the critical thickness (100 nm)³¹ required to avoid a leakage problem. The dual sweep transfer characteristics of the InSe/ WSe₂ heterostructure device are represented in Figure 2a. The nA range current can be attributed to the remote carrier scattering occurring at the semiconductor/P(VDF-TrFE) interface due to the polymer coating and the existence of the depletion region across the p-n junction. As mentioned earlier, the device exhibits ambipolar behavior, and two types of hysteresis loop are observed: one is anti-clockwise on the nside and the other is clockwise on the p-side. Additionally, the polarization direction changes when it reaches the coercive voltage ($V_c = \pm 20$ V), which is clear in the log plot (Figure 2b). The on/off ratio of the device estimated from Figure 2b is $\sim 10^3$, which is a considerable value in the class of 2D heterostructure-based transistors.^{3,32,33} Due to the ferroelectric gating with P(VDF-TrFE), three polarization states can be

observed in the device. First is the normal state without any polarization, second is the polarization up state (P-up), and third is the polarization down state (P-down), which can be obtained by poling the ferroelectric polymer with a certain voltage for 30 seconds (P-up (-50 V) and P-down (+50 V)). The schematic illustration of the polarization up and down states is depicted in Figure 2c,d, respectively.

The photodetecting performance of the device was studied by carrying out electrical measurements with laser illumination of 633 nm at different power intensities. The $I_{ds}-V_{ds}$ characteristics of the device measured in dark conditions at normal and P-up states are shown in Figure 3a. Note that if we consider a unipolar ferroelectric gated transistor, either of the up and down polarization yields higher current saturation in the ON state and the other saturates in the OFF state. So, there will be a drastic difference in the photoresponse obtained in the P-up and P-down polarization states. However, in our present study, we obtain an ambipolar transfer curve with p-WSe2 and n-InSe heterostructure. As we can observe from Figure 2a,b, the drain source current at the P-up and P-down polarized state saturates at a higher value compared with the normal state. So, there will be a minimal difference in the photoresponse measured at P-up and P-down states. Hence, we chose to compare either of the polarized states (P-up here) with the normalized state throughout the study than with the P-down state for clarity. The electrical measurement of the device at normal and P-up states with laser illumination is represented in Figures S3a and 3b, respectively. It can be observed that the photocurrent generation is higher in the polarized state than in the normal state. The photoresponsivity (R) of the device in both states was estimated with the photocurrent (I_{ph}) , incident laser power (P), and the active device area (A) using eq $1.^{26,34}$



Figure 5. Ferroelectric P(VDF-TrFE) gated InSe/WSe₂ heterostructure as a pyroelectric detector. (a) and (b) $I_{ds}-V_{ds}$ characteristic curves of the device showing a change in current when irradiated with 980 and 1064 nm laser, respectively. (c) Responsivity of the device to laser with wavelengths 633, 980, and 1064 nm as a function of the incident laser power intensity.

$$R = \frac{I_{\rm ph}}{P \times A} \tag{1}$$

The maximum responsivity of the $InSe/WSe_2$ device obtained is $\sim 10^4$ A W⁻¹ in the P-up state, which is twice the responsivity obtained in the normal state, i.e. $\sim 5.3 \times 10^3$ A W^{-1} (Figures 3c and S3b). Similarly, the photogain (η) of the device determined using the equation $\eta = (Rhc)/(e\lambda)$ (h is Planck's Constant, c is the speed of light, e is the electronic charge, and λ is the wavelength of the illuminated light)^{27,34} is 1.05×10^4 in the normal state and 2.23×10^4 in the P-up state (Figure 3d). The photo-switching characteristics of the device at the normal and P-up states are given in Figure S3c. In both the states, the device exhibits a stable photo-switching ability. The photoresponse time of the device under the normal state is 250 ms and under the P-up state <200 ms. The P-up state exhibits higher responsivity and photogain than the normal state because of the polarization effect. In the polarized state (Figure 2c), the holes in the InSe and WSe_2 get bound to the remnant polarization field, which reduces the depletion region and increases the conduction. Besides, the Schottky barrier formed between the material and electrode due to the difference in the Fermi level is also greatly reduced, which aids in conduction.³⁴ When the laser is illuminated over the device area, a large number of electron-hole pairs are generated in the material, and one of the carriers, preferably holes, here gets trapped in the remnant polarization field. This acts as a local gating while the other carrier flows through the device, which aids in the enhancement of photocurrent.³⁵ In addition, the device also exhibits a significant photovoltaic effect, as shown in Figure S4, where a considerable negative short-circuit current (I_{sc}) is observed with an increase in laser power intensity. The P-up state exhibits a higher short-circuit current than the normal state, indicating that the charge carrier separation is more in the P-up state, which is following the

mechanism explained earlier. The above behavior provides firm evidence to support the fact that the $InSe/WSe_2$ area contributes more to the better photoresponse in the P-up state as the p-n junction is crucial for charge carrier separation, which aids in enhanced photocurrent.

To demonstrate IR detection in InSe/WSe₂ devices, the pyroelectric effect of the P(VDF-TrFE) is utilized. First, we exclude several possible mechanisms responsible for the IR photoresponse. The lasers 980 and 1064 nm used in this study have energies (1.26, 1.16 eV) less than the band gap of fewlayered InSe (1.27 eV) and WSe_2 (1.36 eV). So, the possible origin of photoresponse to IR rays through the photoexcitation for the band-to-band transition can be ruled out. Another alternative interpretation for the photoresponse may be attributed to the photoexcitation from defect states to band transition. However, this possibility can also be excluded for the magnitude of the photoresponse is in the same order of the above band gap excitation, and the density of defect states should be much less than that of the conduction and valence bands. Hence, it clearly proves the role of the pyroelectric effect in IR detection in our study. The mechanism for IR detection using the pyroelectric effect is illustrated in Figure 4. At a reverse bias condition and thermal equilibrium without the application of any gate voltage, there is a considerable depletion region formed (assume that the length is L) with ptype WSe₂ and n-type InSe, as shown in Figure 4a. At the same reverse bias condition and thermal equilibrium but at the P-up state without NIR irradiation, the holes in the p-WSe2 and n-InSe get bound to the polarization electric field, as depicted in Figure 4b. So, the depletion region is reduced $(L - \Delta L)$ and the current flows through the device (I_{dark}) . However, at a similar condition with IR irradiation, there is a localized heating effect over the polymer increasing the temperature (T+ ΔT). So, the polarization effect decreases $(P - \Delta P)$, and the number of holes bound by the polarization electric field decreases. Consecutively, the depletion region increases (L + ΔL), and more electron-hole separation takes place, increasing the measured current $(I_{ds} + \Delta I_{ds})$, as illustrated in Figure 4c. This overall mechanism was experimentally observed when the InSe/WSe2 device with P(VDF-TrFE) gating was irradiated with IR lasers with wavelengths of 980 and 1064 nm. The I_{ds} - $V_{\rm ds}$ characteristics of the device under 980 and 1064 nm laser irradiation at different laser power intensities are shown in Figure 5a,b, respectively. There is an increase in the photocurrent observed with an increase in IR irradiation intensity, as observed from the represented graphs. The increase in temperature (ΔT) for different IR laser intensities for 980- and 1064-nm lasers is shown in Figure S5. The maximum responsivity obtained with 980 nm laser irradiation is 9.6 \times 10² Å W⁻¹ at a laser power of 9.2 μ W cm⁻² whereas, for 1064-nm laser irradiation, it is 7.9×10^2 A W⁻¹ at 0.17 mW cm^{-2} . Notably, the responsivity of the pyroelectric IR detection for the photon below the band gap of 2D materials is in the same order of magnitude as that of the normal state under the excitation of the photon energy above the band gap without the effect of pyroelectricity. To study further, the pyroelectric current was measured as a function of frequency (f) at a constant IR irradiation with 980 and 1064 nm laser, as shown in Figure S6. It can be observed that the pyroelectric current measured, $R_{py}(\omega)$, decreases with an increase in frequency, which can be fitted with the following model³⁶

$$R_{\rm py}(\omega) = \frac{X(\omega)}{\sqrt{(1+\omega^2\tau^2)}}$$
(2)

where $X(\omega)$ is a polynomial term associated with the thermal effect in the pyroelectric polymer, ω is the frequency, and τ is the pyroelectric response time constant. The curves fitted with eq 2 are in accordance with the experimental data, as shown in Figure S6. The pyroelectric response time (τ) obtained for 980 and 1064 nm laser irradiation is 240 and 290 ms, respectively. According to our proposed mechanism, as the IR detection in our reported device is mainly due to the pyroelectric effect of P(VDF-TrFE), any IR laser that causes a localized heating effect upon irradiating with P(VDF-TrFE) dielectric would change its polarization, resulting in a change in the channel current in IR detection.

In addition to responsivity and photogain, specific detectivity is one of the most important figures of merit to justify the sensitivity of a photodetector. The larger the detectivity, the better it is for obtaining signals out of noises. As shown in Supplementary Note 3, the calculated detectivity for 633 nm laser illumination can reach up to about 10¹³ Jones, and IR radiation with the photon energy below the band gap of the constituent 2D materials is up to 10¹¹ Jones of magnitude (Supplementary Note 5 and Figure S7), which is much better than all the previous reports of pyroelectric IR detectors, including 2D materials.^{37–39} For example, the pyroelectric IR detector base on graphene/PVDF has a detectivity of only 7.42 \times 10⁶ Jones.³⁹ The origin of the enhanced photosensitivity can be attributed to the uniqueness of the coupling between the pyroelectricity and the built-in electric field across the p-njunction. Under IR irradiation, the increased temperature will induce the change of the PVDF polarization and, in turn, change the built-in electric field across the heterojunction. Consequently, due to the charge carriers' flow reaching a new equilibrium state, the depletion width is changed and the band alignment across the heterojunction is rearranged, which

enables us to increase the charge carrier separation and enhances the photocurrent.

CONCLUSIONS

To conclude, we have demonstrated a highly sensitive photodetector based on a new mechanism of the coupling between the pyroelectric and built-in electric field for the device consisting of the 2D InSe/WSe₂ heterojunction and P(VDF-TrFE), which can detect a wide range of wavelengths covering the visible to IR spectrum. The ferroelectric polarization in the dielectric leads to an excellent photodetection in the visible range with a photoresponsivity of 10^4 A W^{-1} . On the other hand, using the heterojunction as the channel, the coupling between the pyroelectric effect of the polymer and the built-in electric field across the p-n junction extensively promoted highly sensitive IR detection. The increased temperature induces a polarization change in the irradiated area and creates charge carriers flow to establish a new equilibrium state and changes the depletion width. The specific detectivity of the IR detection is in the order of 10¹¹ Jones, which is much better than all published pyroelectric IR detectors, such as the devices based on PVDF/graphene 2D materials. Thus, the coordinated benefaction from both the dielectric and the 2D heterostructure opens up multifarious possibilities for novel optoelectronic devices. For instance, with the overlap of the absorption spectrum of PVDF and black body radiation at 300 K, 34 it is foreseeable that our approach may introduce not-yet realized applications in the fields ranging from bio-imaging to military defense.

EXPERIMENTAL DETAILS

Device Fabrication. The dry transfer technique was followed to stack up the InSe/WSe2 heterostructure. Both the crystals were mechanically exfoliated into thin flakes using a scotch tape. Initially, exfoliated WSe2 flakes were transferred to the SiO2/Si substrate with a PDMS stamp. In the same manner, exfoliated InSe flakes were transferred exactly to the top of the WSe2 flake using a customdesigned micromanipulator fitted to an optical microscope (Olympus, BX 51M) with a CCD camera (Leica DFC495). After determining the thickness of the stacked InSe/WSe2 heterostructure using the AFM technique, electrodes (Cr/Au-7/70 nm) were deposited by the thermal evaporation technique while masking the heterojunction using a Cu grid with mesh sizes ranging in the microns. For ferroelectric gating, the P(VDF-TrFE) solution was spin-coated on the device at 3500 rpm for 20 s. The pyroelectric and ferroelectric properties of P(VDF-TrFE) films were demonstrated in previous reports.¹⁸ After baking the device in a vacuum oven at 135 °C for 2 h, Al/Au was deposited on the heterojunction area by using a custommade shadow mask to serve as an electrode for top-gating the device.

Characterization Details. The thickness of the InSe and WSe₂ flakes was determined using a nanosurf Flex AFM (Easyscan 2V.3.1). The Raman Spectrum of the heterojunction was recorded using a micro-Raman spectrometer (Triax 550) equipped with an optical microscope (Olympus CX41) and detector (Jobin Won Horiba S Drive-500 Syncernity). The wavelength of the laser used for excitation is 532 nm. For the electrical measurements, Keithley 2636 and 2400 were employed. To measure the photoresponse of the device, an He–Ne laser with a wavelength of 633 nm was used. For IR detection, a continuous-wave 980 nm laser was used.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsami.2c22876.

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Heterostructure assembly; microscopic images; photoresponse of the device; photovoltaic effect, and pyroelectric effect (PDF)

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Notes

The authors declare no competing financial interest.

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Supporting Information

Coupling Between Pyroelectricity and Built-in Electric Field Enabled Highly Sensitive Infrared Phototransistor Based on InSe/WSe₂/P(VDF-TrFE) Heterostructure

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Supplementary Note 1, Heterostructure assembly

The schematic illustration of each step involved in the InSe/WSe₂ heterostructure device fabrication with P(VDF-TrFE) coating as a dielectric layer is shown in **Figure S1**. At first, WSe₂ flakes were exfoliated and transferred to the substrate using a PDMS stamp. After removing the PDMS stamp, the exfoliated InSe was precisely placed on the WSe₂ flake using the custom-designed micromanipulator. The electrodes were deposited for the assembled heterostructure and the P(VDF-TrFE) copolymer solution is spin-coated on the device. After annealing the device, electrodes were deposited for top-gating.



Figure S1. Schematic depiction of various steps involved in the device fabrication which includes assembling InSe/WSe₂ heterostructure, depositing electrodes, coating P(VDF-TrFE) dielectric on top, and depositing gate electrode.

Supplementary Note 2, Microscopic images

The optical microscope image of the InSe/WSe₂ heterostructure device after P(VDF-TrFE) coating and top electrode deposition are represented in **Figure S2**.



Figure S2. Optical microscope image of the P(VDF-TrFE) coated top-gated InSe/WSe₂ heterostructure.

Supplementary Note 3, Photoresponse of the device

The I_{ds} - V_{ds} characteristics of the device under laser (λ = 633 nm) illumination with different power intensities in the normal state are shown in **Figure S3a**. The photocurrent increases with an increase in laser power. The photoresponsivity of the device is estimated to be in the order of 10³ A W⁻¹. Another significant parameter that determines the detection sensitivity is the detectivity (D*) which can be expressed in terms of noise equivalent power (NEP) as,¹

$$D^* = \frac{(A \times \Delta f)^{1/2}}{NEP} \tag{1}$$

where A is the active device area and Δf is the measuring bandwidth. NEP signifies the minimum optical signal that can be distinguished from noise by the photodetector and is directly proportional to the root mean square noise current (I_n) and inversely proportional to the responsivity (R) of the device which can be expressed as NEP = I_n / R . The noise current is directly proportional to the dark current as follows,

$$I_n^2 = 2eI_{dark}\Delta f \tag{2}$$

where e is the electronic charge. By substituting the expression for NEP and noise current, equation (1) can be simplified and expressed as given in equation (3).

$$D^* = \frac{(A)^{1/2} \times R}{(2eI_{dark})^{1/2}}$$
(3)

The detectivity (D*) of the InSe/WSe₂ heterostructure device in the normal state calculated at different laser power intensities is shown in **Figure S3b**. The maximum detectivity obtained is in the order of 10^{14} Jones at a lower laser power. To double-check the accuracy of the calculated detectivity, spectral noise was used to estimate the detectivity following our previous publication.² The spectral noise density $S_n = 1.1 \times 10^{-11} \text{ A} / \text{Hz}^{0.5}$ was obtained from the noise-portion of Fast-S4

Fourier-Transform of the photocurrent. Based on the measured responsivity, the noise-equivalent power NEP = $S_n / R = 2 \times 10^{-15} \text{ W} / \text{Hz}^{0.5}$. This value yields a specific detectivity as follows,

$$D^* = \frac{\sqrt{A\Delta f}}{NEP} = 1 \times 10^{12}$$
 Jones

where A is the active area of the detector and Δf is the bandwidth. The obtained result was not too much different from the calculation based on the dark current as shown above. The photoresponse time of the device under normal state is 250 ms and the P-up state is < 200 ms (**Figure S3c**).



Figure S3. (a) $I_{ds}-V_{ds}$ curves measured at normal state under 633 nm laser illumination with different power intensities. (b) Responsivity and detectivity of the device as a function of laser power intensity at a normal state. (c) The photo-switching characteristics of the device.

Supplementary Note 4, Photovoltaic Effect

The electrical measurements carried out for $InSe/WSe_2$ heterostructure at the normal and polarized (P-up) state under laser illumination exhibit a significant photovoltaic effect. A notable negative short-circuit current (I_{sc}) is observed in the device at zero bias as shown in **Figure S4**. The P-up state exhibits a higher short-circuit current than the normal state which indicates that the charge carrier separation is more in the P-up state due to the localized gating effect by the bound carriers to the remnant polarization field.



Figure S4. Short-circuit current (I_{sc}) at zero bias as a function of incident laser power intensity at normal and ferroelectric polarized state.

Supplementary Note 5, Pyroelectric Effect

The change in temperature at the IR laser-irradiated region was estimated using the following equation.

$$I_{ir} = \varepsilon \sigma (T^4 - T_A^4)$$

Here, I_{ir} is the intensity of IR radiation, ε is the emissivity of the surface (0.7 for P(VDF-TrFE) film), σ is the Stefan-Boltzmann constant (5.6703 × 10⁻⁸ W m⁻²K⁻⁴), T is the surface temperature and T_A is the ambient temperature. The increase in temperature is determined as $\Delta T = T - T_A$ and the ΔT for different IR laser intensity for 980 nm and 1064 nm laser is shown in **Figure S5**.



Figure S5. The increase in temperature (Δ T) at the irradiated area of the device as a function of IR laser power intensity.

To support the near-infrared (NIR) photodetection, the pyroelectric current was measured as a function of frequency (f) at a constant NIR irradiation with 980 and 1064 nm laser as depicted in **Figure S6**. The pyroelectric current measured decreases with an increase in frequency and it matches well with the fitted model.



Figure S6. (a) and (b) Change in current due to pyroelectric effect measured as a function of frequency (f) when irradiated with 980 nm and 1064 nm laser, respectively.

The specific detectivity of IR irradiation estimated using equation (1) with NEP calculated from spectral noise density (S_n) (**Supplementary Note 3**) was shown in **Figure S7** as a function of laser power intensity. The calculated detectivity for 980 nm and 1064 nm laser irradiation is in the order of 10^{11} Jones. Thus, the specific detectivity of our IR detector is much better than the previously reported values.^{3–5}



Figure S7. The specific detectivity (D*) of the device for IR irradiation as a function of laser power intensity.

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