

Cite this: *Mater. Adv.*, 2023,
4, 596

Reduced-graphene oxide decorated γ - In_2Se_3 /Si heterostructure-based broadband photodetectors with enhanced figures-of-merit

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Recently, two-dimensional (2D) semiconductor-based broadband photodetectors have gained tremendous attention due to their immense potential applications in high-performance optoelectronic devices. In this report, we demonstrate a high-performance photodetector based on reduced-graphene oxide (rGO) decorated p- γ - In_2Se_3 /n-Si heterostructures. The integration of rGO with a p- In_2Se_3 /n-Si heterostructure results in an enhanced responsivity of $\sim 9.5 \text{ A W}^{-1}$ with detectivity of $\sim 2.39 \times 10^{12} \text{ cm Hz}^{1/2} \text{ W}^{-1}$ at -3 V under the illumination of 685 nm light with an intensity of 0.1 mW cm^{-2} . In addition, the rGO/ In_2Se_3 /Si heterostructure shows high sensitivity and fast response/recovery times (40/90 μs) with a broadband response ranging from visible to infra-red wavelengths, which makes it a suitable candidate for an efficient broadband photodetector. The enhanced figures-of-merit of the rGO/ γ - In_2Se_3 /Si heterostructure could be due to the increased optical absorption of incident light and effective separation of the photogenerated charge carriers because of the top rGO layer. Here, the rGO layer acts as an efficient hole transporting layer due to its high hole mobility, which effectively reduces the recombination of the photogenerated charge carriers.

Received 21st September 2022,
Accepted 4th December 2022

DOI: 10.1039/d2ma00918h

rsc.li/materials-advances

1. Introduction

Photodetectors based on two-dimensional (2D) layered semiconductor materials have gained incredible research interest in recent years due to their immense potential applications in high performance optoelectronic devices.^{1–4} 2D layered semiconductors provide an alternative and viable approach to encounter the challenges faced by Si-based optoelectronic devices because of their exceptional and unique characteristics. Si-based photodetectors are sensitive in the near IR region because of their narrow band gap ($\sim 1.1 \text{ eV}$).^{5,6} In addition, Si-based photodetectors show low photo-responsivity. To demonstrate Si-based broadband photodetectors with high photo-responsivity, it is essential to integrate Si with 2D-layered semiconductors with appropriate band gaps. Such hybrid 2D/3D heterostructures could pave the way toward the realization of high-performance broadband photodetectors.^{7–11} As an important class of 2D-layered materials, IIIA metal chalcogenides

(where, metal = In, Ga; chalcogenide = S, Se, Te) have shown great potential in electronic and optoelectronic devices due to their interesting electrical and optical properties. Among these, In_2Se_3 has gained substantial attention as a promising 2D material because of its direct wide bandgap, high absorption coefficient in the visible range and efficient generation of electron-hole pairs under light illumination,^{12–14} which makes it an ideal candidate to integrate with Si for the demonstration of a broadband photodetector with high photo-responsivity. In_2Se_3 exists in three different crystalline phases denoted by α , β and γ .^{15–17} Among them γ - In_2Se_3 is a 2D material having a hexagonal crystal structure with a distorted wurtzite type atomic arrangement¹⁸ and having a wide direct bandgap of $\sim 1.8 \text{ eV}$.^{19–21} Hence, by integrating γ - In_2Se_3 with Si, a photodetector with broadband detection can be achieved. In addition, the integration of In_2Se_3 with Si creates a depletion region at the interface, which generates a large built-in electric field at the In_2Se_3 /Si interface. This internal electric field facilitates the separation of photogenerated charge carriers and suppresses the recombination probability. Another approach to further achieve high performance broadband photodetectors is by utilizing a van der Waals heterostructure (2D/2D hybrid configuration) on a Si substrate. By integrating van der Waals heterostructures on Si, the device performance is further enhanced because of the perfect interface with low defects at the 2D/2D interface.^{22,23} In the present studies, we have

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integrated reduced-graphene oxide (rGO)/In₂Se₃ as a 2D/2D heterostructure on Si and demonstrate a broadband photo-detector with high photo-responsivity and fast response/recovery times.

As of now, there are few reports on the photodetection properties of In₂Se₃ nanostructure-based photodetectors.^{23–27} Gedrim *et al.* demonstrated a 2D α -In₂Se₃ nanosheet-based photodetector with photo-responsivity of $3.95 \times 10^2 \text{ A W}^{-1}$ and an external quantum efficiency (EQE) of $1.63 \times 10^5\%$ under the illumination of 300 nm at a bias voltage of 5 V.²³ Moreover, the response time and specific detectivity of the detector were $1.8 \times 10^2 \text{ s}$ and $2.26 \times 10^{12} \text{ cm Hz}^{1/2} \text{ W}^{-1}$, respectively. Here, α -In₂Se₃ nanosheets were obtained through the exfoliation of commercial grade α -In₂Se₃ onto a SiO₂/Si substrate. Li *et al.* fabricated a photodetector based on a single In₂Se₃ nanowire, which showed a photo-responsivity of $1.5 \times 10^2 \text{ A W}^{-1}$ at 500 nm with an EQE of $3.7 \times 10^4\%$ at an applied bias of 5 V.²⁴ Here, the authors synthesized In₂Se₃ nanowires at around 950 °C using a high temperature tubular furnace. In another report, Zhai *et al.* demonstrated a fast, reversible and stable response of In₂Se₃ nanowire-based photodetectors with photo-responsivity and EQE of $\sim 89 \text{ A W}^{-1}$ and $\sim 22\,000\%$, respectively, under an incident wavelength of 500 nm at an applied bias of 3 V, where In₂Se₃ nanowires were synthesized at 900 °C using a high temperature tubular furnace. Here, the high photo-responsivity and fast photoresponse are attributed to the superior single-crystal quality and large surface-to-volume ratio of In₂Se₃ nanowires.²⁵ Chen *et al.* demonstrated an In₂Se₃ nanoflower/Si heterostructure-based self-powered and broadband photodetector ranging from UV to IR with a responsivity of 5.67 A W^{-1} and detectivity of $5.66 \times 10^{13} \text{ cm Hz}^{1/2} \text{ W}^{-1}$ under an illumination of 820 nm at -1 V bias voltage.²⁶ In the above reports, the authors have synthesized In₂Se₃ in various nanostructured forms, like nanosheets, nanowires, nanoflakes, nanoflowers, *etc.* using mechanical/chemical exfoliation or high temperature chemical vapor deposition techniques, where the size of the nanostructure is limited to a few microns. Though the performance of those nanostructure-based photodetectors is excellent, the adopted synthesis techniques have a major limitation of being scaled for mass production. In this article, a large area In₂Se₃ thin film has been realized using a simple technique for photodetection applications. Here, a large area γ -In₂Se₃ thin film has been deposited using thermally evaporating indium on a Si substrate followed by selenisation at much lower temperature and a high performance γ -In₂Se₃/Si heterostructure based broadband photodetector has been demonstrated.

To the best of our knowledge, there are few reports on In₂Se₃ thin film-based high performance broadband photodetectors.^{17,28} Shao *et al.* have grown a large area β -In₂Se₃ thin film on a c-sapphire substrate using a molecular beam epitaxy system and demonstrated a graphene/ β -In₂Se₃ heterostructure-based photodetector with responsivity and EQE of 1.17 A W^{-1} and 93.6%, respectively, under the illumination of 1550 nm at an applied bias of 0.35 V.²⁸ Wu *et al.* have fabricated γ -In₂Se₃/n-Si-based broadband photodetectors by depositing γ -In₂Se₃

on an n-Si substrate using RF magnetron sputtering. A photo-responsivity of 0.57 A W^{-1} with a specific detectivity of $2.6 \times 10^{12} \text{ cm Hz}^{1/2} \text{ W}^{-1}$ was obtained under 808 nm light illumination with rise/decay times of 35/115 μs .¹⁷ Herein, we have demonstrated γ -In₂Se₃ (p-type)/n-Si heterostructure-based high performance broadband photodetectors with a photo-responsivity of 4.6 A W^{-1} and specific detectivity of $1.22 \times 10^{12} \text{ cm Hz}^{1/2} \text{ W}^{-1}$ under the illumination of 685 nm at an applied bias of -3 V . In addition, we have decorated the γ -In₂Se₃/n-Si heterostructure with reduced-graphene oxide (rGO) and observe an enhanced photo-responsivity of 9.5 A W^{-1} with a specific detectivity of $\sim 2.39 \times 10^{12} \text{ cm Hz}^{1/2} \text{ W}^{-1}$ and response/recovery times of 40/90 μs under an illumination of 685 nm at an applied bias of -3 V , which are better than the other reported values with similar device geometries. The enhanced device performance of the rGO/ γ -In₂Se₃/n-Si heterostructure is attributed to the effective separation of photogenerated charge carriers due to the efficient hole transporting nature of the top rGO layer, which reduces the recombination probability of the charge carriers.

II. Experimental procedure

A commercial n-doped Si (100) substrate was used for the deposition of γ -In₂Se₃ thin film. The carrier concentration and electrical conductivity of the Si substrate as estimated using Hall measurement were $\sim 7.5 \times 10^{14} \text{ cm}^{-3}$ and $\sim 1.8 \times 10^{-1} (\Omega \text{ cm})^{-1}$, respectively. Here, the deposition of the γ -In₂Se₃ thin film on the n-Si (100) substrate was carried out by thermally depositing indium metal on the Si substrate followed by its selenisation. Prior to the selenisation, a circle of indium metal of diameter 1000 μm was deposited on the Si substrate using a standard photolithography process followed by the thermal evaporation and lift off techniques. Subsequently, selenisation of the indium metal was carried out in a tubular furnace at 450 °C for 1 h in the presence of a continuous flow of argon gas. The sequential steps involved in the deposition of In₂Se₃ on Si followed by the fabrication of the rGO/In₂Se₃/Si heterostructure are shown in Fig. 1(a)–(e).

To fabricate the rGO/In₂Se₃/Si heterostructure, rGO was drop-cast on the In₂Se₃/Si heterostructure. Here, rGO was synthesized using the modified Hummers' method.²⁹ The photodetection measurements of the In₂Se₃/Si and rGO/In₂Se₃/Si heterostructures were carried out by taking one contact from the top of the In₂Se₃ (or rGO) layer and another from the bottom of the Si substrate, as shown in Fig. 1(c) and (e), respectively. A circle of aluminium metal of diameter 400 μm on the In₂Se₃ (or rGO) layer was used as the top electrode. Fig. 1(f) shows an optical image of the fabricated γ -In₂Se₃/Si heterostructure-based photodetector. Here, the effective area of the device is around 0.659 mm^2 (excluding the area of the metal electrode). Oriel's QEPVSI system equipped with a 300 W Xenon bulb was used for the photodetection measurements of the heterostructure. The light intensity of the system was varied by changing the width of the entrance slit. The structural characterizations of the



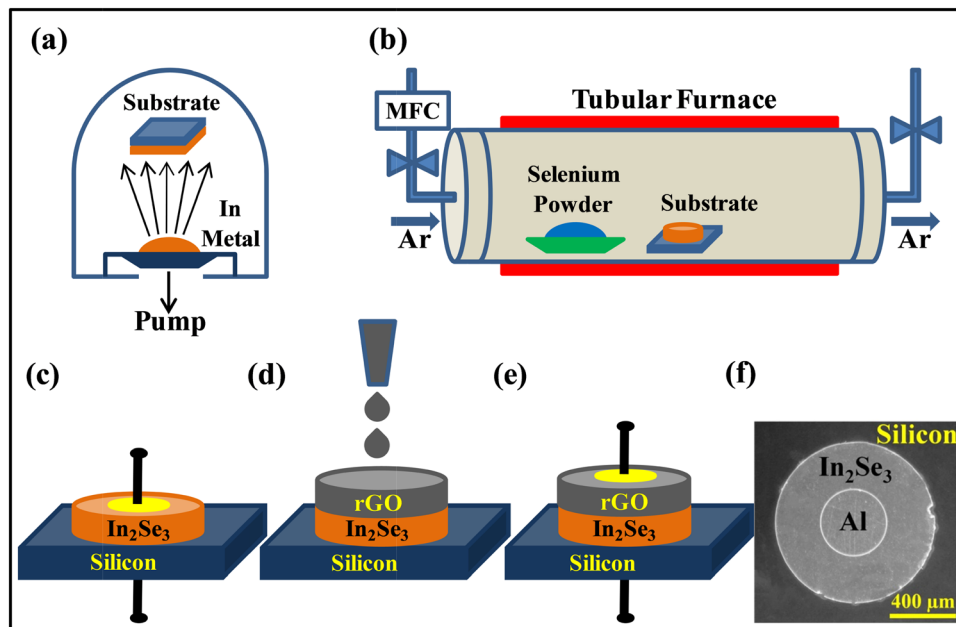


Fig. 1 The complete process step sequences for the deposition of γ - In_2Se_3 on the Si substrate followed by the fabrication of the $\text{rGO}/\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure-based photodetector, (a) deposition of indium metal on the Si substrate using a thermal evaporation system, (b) selenisation of the indium metal using a tubular furnace, (c) fabrication of the $\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure-based photodetector, (d) decoration of the $\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure with rGO using the drop-casting technique, (e) fabrication of the $\text{rGO}/\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure-based photodetector, and (f) an optical image of the fabricated $\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure-based photodetector.

$\text{In}_2\text{Se}_3/\text{Si}$ and $\text{rGO}/\text{In}_2\text{Se}_3/\text{Si}$ heterostructures were carried out using X-ray diffraction (XRD) and Raman spectroscopy. The surface morphology and composition of the In_2Se_3 thin film were determined using field emission scanning electron microscopy (FESEM) and energy dispersive X-ray spectroscopy (EDS), respectively. A Hitachi (U-2900) absorption spectrophotometer was used to estimate the optical band gap of the $\gamma\text{-In}_2\text{Se}_3$ film. Room temperature Hall measurement was carried out to determine the carrier concentration of the In_2Se_3 thin film and Si substrate using an Ecopia HMS 5000 system.

III. Results and discussion

Fig. 2(a) shows the XRD pattern of the as-deposited In_2Se_3 thin film on the Si (100) substrate. All diffraction peaks as marked with an asterisk are consistent with the ICSD standard card of the γ -crystalline phase of In_2Se_3 , confirming the deposition of the γ -phase of In_2Se_3 thin film. To further examine the γ -phase of In_2Se_3 , micro-Raman spectroscopy has been carried out on the $\text{rGO}/\text{In}_2\text{Se}_3/\text{Si}$ heterostructure. As displayed in Fig. 2(b), Raman spectroscopy shows the clear characteristics peaks (as denoted with an asterisk) associated with the different

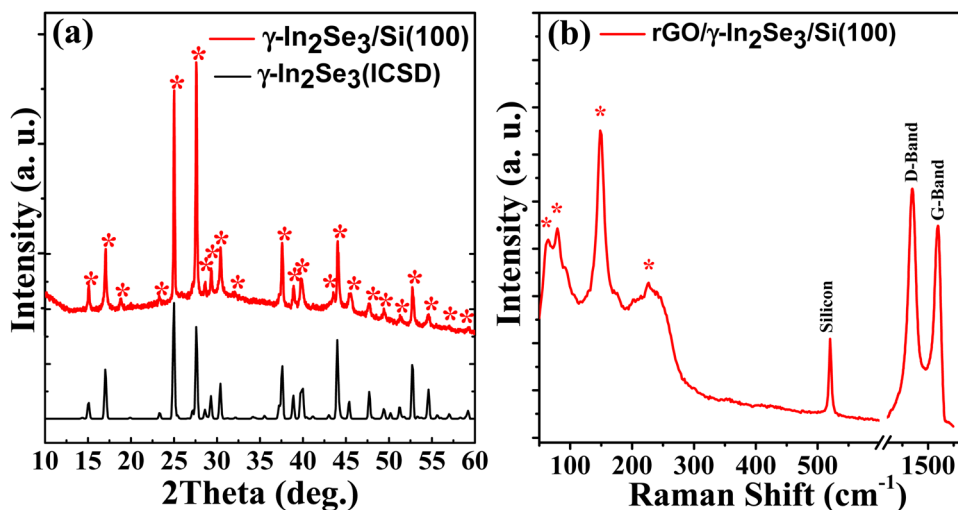


Fig. 2 (a) XRD of the $\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure and (b) room-temperature μ -Raman spectroscopy of the $\text{rGO}/\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure.



vibrational modes of $\gamma\text{-In}_2\text{Se}_3$. The peak at 520 cm^{-1} is associated with the E_2 (high) mode of the Si substrate. In addition, the Raman spectrum shows two peaks at around ~ 1346.4 and $\sim 1591.9\text{ cm}^{-1}$, which are assigned to the D- and G-band of rGO, respectively.^{30,31}

Fig. 3(a) shows the optical absorption spectra of the as-deposited $\gamma\text{-In}_2\text{Se}_3$ thin films with and without rGO coating on a soda-lime glass (SLG) substrate. Here, the SLG substrate was used in order to avoid the optical absorption on the Si substrate. The deposition of the $\gamma\text{-In}_2\text{Se}_3$ thin film on the SLG substrate was carried out using the same techniques as used for $\gamma\text{-In}_2\text{Se}_3$ on the Si substrate. With the rGO coating on the $\gamma\text{-In}_2\text{Se}_3$ thin film, the optical absorption is increased considerably in the wavelength range of 400–1100 nm as compared to without the rGO coating. A plot of $h\nu$ vs. $(\alpha h\nu)^2$, as shown in Fig. 3(b), yields a direct bandgap of 1.81 eV, which is very close to the reported value of $\gamma\text{-In}_2\text{Se}_3$.^{19–21} Fig. 4(a) shows the FESEM image of $\gamma\text{-In}_2\text{Se}_3$ on the Si substrate, which shows the presence of uniform nanoscale particulates on the surface of the film. In order to measure the thickness of the $\gamma\text{-In}_2\text{Se}_3$ thin film, cross-sectional SEM was carried out, as shown in the inset

of Fig. 4(a). The thickness of $\gamma\text{-In}_2\text{Se}_3$ was found to be around 320 nm. Fig. 4(b) shows EDS of $\gamma\text{-In}_2\text{Se}_3$ on the Si substrate, which shows the presence of In and Se without any impurities. The average ratio of In to Se is approximately 2:3, indicating that the $\gamma\text{-In}_2\text{Se}_3$ film is stoichiometric.

Fig. 5(a) and (b) show the I - V characteristics of the $\gamma\text{-In}_2\text{Se}_3/\text{Si}$ (detector-1) and $\text{rGO}/\gamma\text{-In}_2\text{Se}_3/\text{Si}$ (detector-2) heterostructures, respectively, in the dark and under the illumination of 685 nm light with a power density of 0.1 mW cm^{-2} . Here, both heterostructures exhibit rectifying I - V characteristics which demonstrate a well-defined photodiode. Under dark conditions, the rectification ratio (on/off ratio) of detector-1 and detector-2 is 30.5 and 24.1, respectively, at an applied bias of $\pm 3\text{ V}$. Under the illumination of 685 nm, the detectors show an enhanced reverse bias current with insignificant detection under forward bias conditions. The dark current (I_{dark}) of detector-1 and -2 is approximately 1.83×10^{-7} and $3.26 \times 10^{-7}\text{ A}$, respectively, at an applied bias of -3 V . On the other hand, the current under an illumination (I_{illu}) of 685 nm is 2.29×10^{-6} and $5.81 \times 10^{-6}\text{ A}$, resulting in a photocurrent ($I_{\text{ph}} = I_{\text{illu}} - I_{\text{dark}}$) of 2.10×10^{-6} and $5.48 \times 10^{-6}\text{ A}$ for detector-1

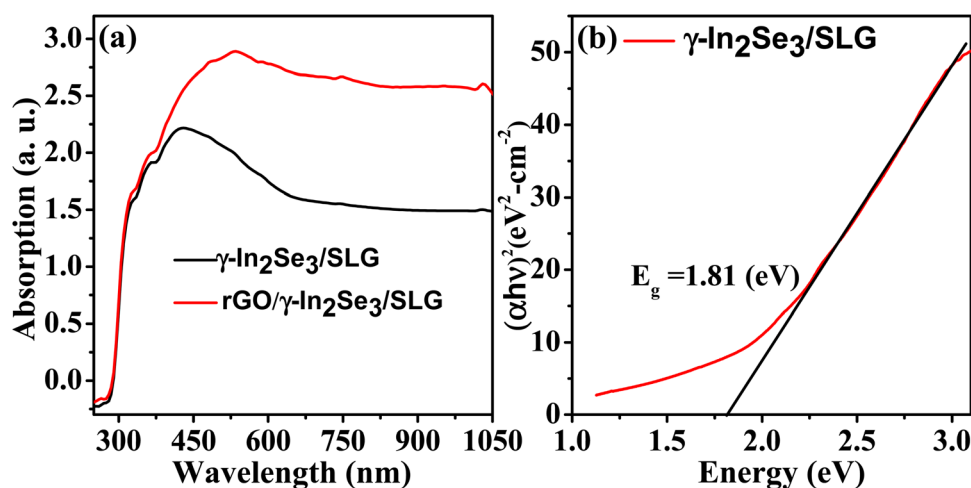


Fig. 3 (a) The optical absorption spectra of $\gamma\text{-In}_2\text{Se}_3$ and $\text{rGO}/\gamma\text{-In}_2\text{Se}_3$ thin films on the SLG substrate and (b) a plot of $h\nu$ vs. $(\alpha h\nu)^2$ of the $\gamma\text{-In}_2\text{Se}_3$ thin film on the SLG substrate.

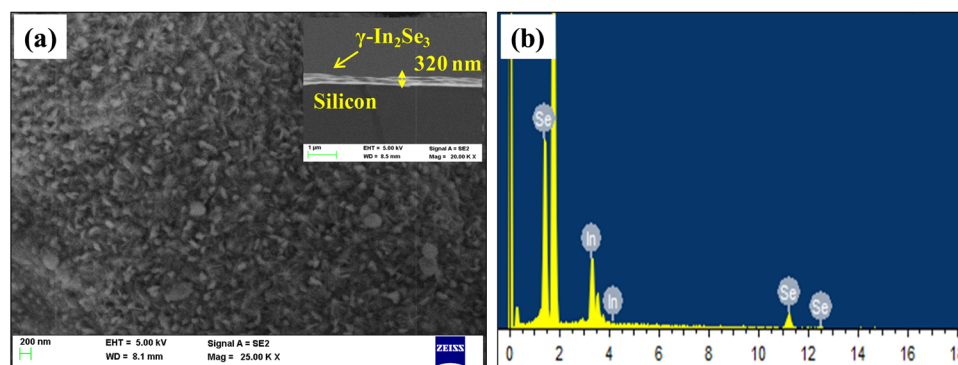


Fig. 4 (a) FESEM image of the $\gamma\text{-In}_2\text{Se}_3$ thin film on the Si substrate. The inset shows the cross-sectional SEM image of the $\gamma\text{-In}_2\text{Se}_3/\text{Si}$ heterostructure and (b) EDS of the $\gamma\text{-In}_2\text{Se}_3$ thin film deposited on the Si substrate.

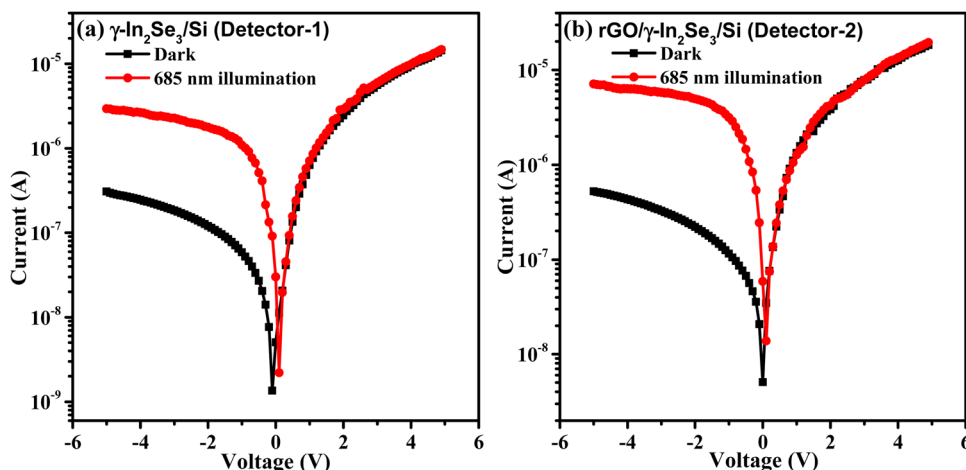


Fig. 5 Room temperature I - V characteristics of (a) γ - $\text{In}_2\text{Se}_3/\text{Si}$ (detector-1) and (b) rGO/γ - $\text{In}_2\text{Se}_3/\text{Si}$ (detector-2) heterostructure-based photodetectors in the dark and under an illumination of 685 nm with a power density of 0.1 mW cm^{-2} .

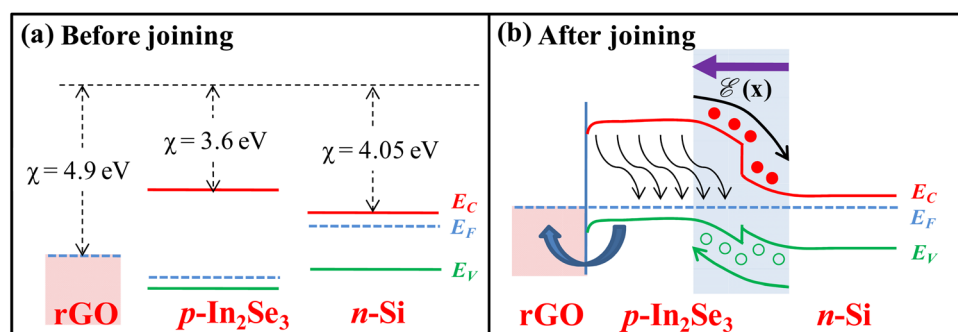


Fig. 6 (a) The energy band diagram of rGO, $\text{p-In}_2\text{Se}_3$ and n-Si before joining, and (b) an equivalent band diagram of the $\text{rGO}/\text{p-In}_2\text{Se}_3/\text{n-Si}$ heterostructure.

and detector-2, respectively. Thus, the decoration of the $\text{In}_2\text{Se}_3/\text{Si}$ heterostructure with rGO improved the photocurrent of the detectors, which could be due to the increase in optical absorption of the incident light.

The improved photocurrent in the rGO decorated $\text{In}_2\text{Se}_3/\text{Si}$ heterostructure is explained using a band diagram of the $\text{rGO}/\text{In}_2\text{Se}_3/\text{Si}$ heterostructure. Fig. 6(a) shows the energy band diagram of rGO, In_2Se_3 and Si before joining them together. Here, the electron affinity (χ) of rGO, γ - In_2Se_3 and Si is taken as 4.9,³² 3.6^{33–35} and 4.05 eV,^{6,36} respectively, based on the previous reports. The band gap of Si is taken as 1.12,³⁶ whereas the bandgap of γ - In_2Se_3 is 1.81 eV as estimated using absorption spectroscopy as shown in Fig. 3(b). The doping concentration of Si and γ - In_2Se_3 is estimated using Hall measurement and is found to be $\sim 7.5 \times 10^{14} \text{ cm}^{-3}$ for Si with n-type behavior and $\sim 2.28 \times 10^{16} \text{ cm}^{-3}$ for γ - In_2Se_3 with p-type behavior. Fig. 6(b) shows an equivalent band diagram of the heterostructure after joining individual layers together. A depletion region is created at the interface of the γ - In_2Se_3 and Si heterostructure, leading to the generation of a built-in electric field at the γ - $\text{In}_2\text{Se}_3/\text{Si}$ interface with the direction from Si to γ - In_2Se_3 . In addition, the $\text{In}_2\text{Se}_3/\text{Si}$ heterostructure produces a type-I band alignment³⁷ at the interface with conduction and valence band offsets of

0.45 and 0.24 eV, respectively. Under reversed bias conditions, when the heterostructure is illuminated by light, the photo-generated electron-hole pairs in the depletion region are effectively separated in opposite directions due to the presence of a high built-in electric field across the γ - $\text{In}_2\text{Se}_3/\text{Si}$ interface. Here, the photo-generated electrons are swept towards the Si side and at the same time the holes are swept towards the γ - In_2Se_3 layer and subsequently the holes are extracted by the top rGO layer, giving rise to high photocurrent in the heterostructure. Here, rGO acts as an efficient hole transporting layer, where the recombination probability of the photo-generated charge carriers reduced drastically. Hence, the photocurrent increases significantly in the rGO decorated γ - $\text{In}_2\text{Se}_3/\text{Si}$ heterostructures.

The figures-of-merit like responsivity, detectivity and sensitivity are calculated to evaluate the performance of the detectors. The responsivity (R_λ) of the photodetectors, as defined by the ratio of the photocurrent to the intensity of the illuminated light, is calculated by using the relation,^{6,38–40} $R_\lambda = I_\lambda/P_\lambda S$, where I_λ is the photocurrent ($I_{\text{illumination}} - I_{\text{dark}}$), P_λ is the light intensity and S is the effective area of the heterostructure. Fig. 7(a) shows the voltage dependent responsivity of detector-1 and detector-2 under an illumination of 685 nm. The responsivity of the detectors



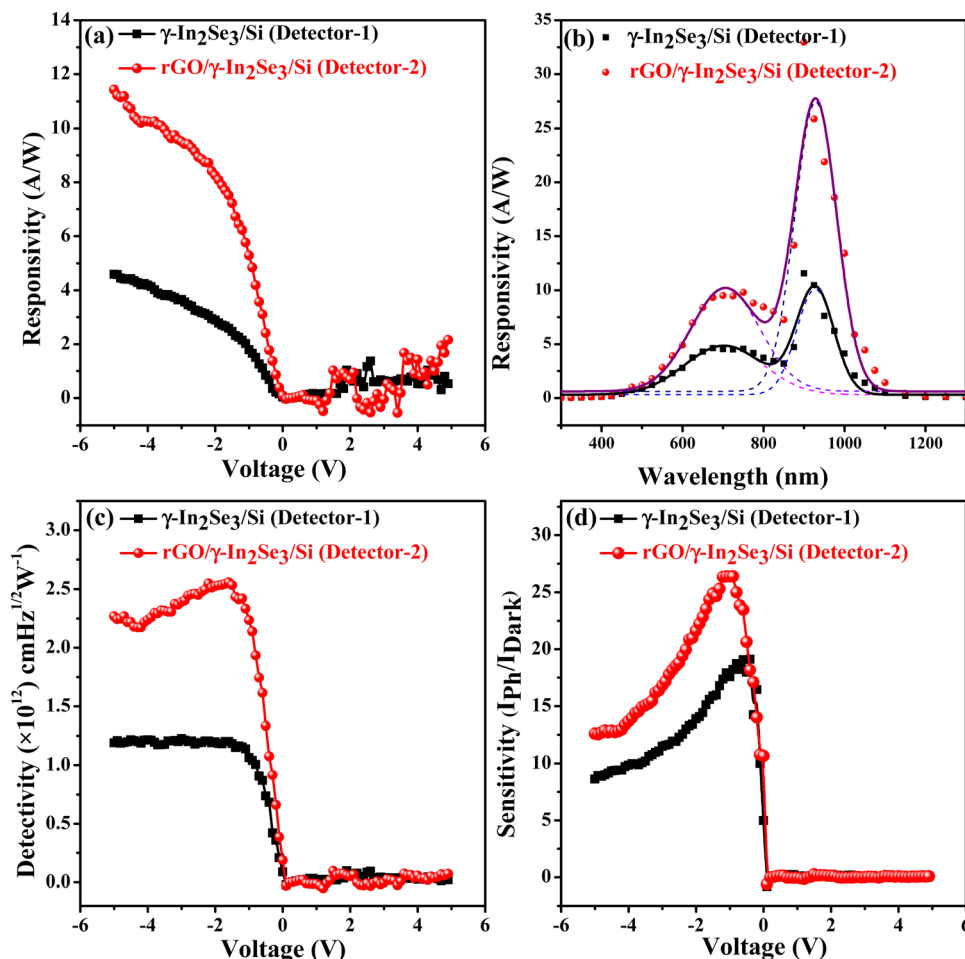


Fig. 7 (a) The voltage dependent photo-responsivity of the detectors under an illumination of 685 nm, (b) the spectral response of the detectors in the wavelength range of 300–1300 nm, measured at an applied bias of -3 V with light intensity of 0.1 mW cm^{-2} , (c) the voltage dependent specific detectivity of the detectors under an illumination of 685 nm, and (d) the voltage dependent sensitivity of the detectors under an illumination of 685 nm.

increases with applied reverse bias voltage, which is due to the increase in photocurrent. At higher applied reverse bias voltage, the photo-generated charge carriers are effectively separated due to the presence of the high internal electric field at the $\text{In}_2\text{Se}_3/\text{Si}$ interface, which resulted in high photocurrent. Moreover, the responsivity of detector-2 is significantly higher than that of detector-1 because of the top rGO layer, which further reduces the recombination of the photo-generated charge carriers due to its efficient hole transporting properties. The values of R_λ were found to be 4.6 and 9.5 A W^{-1} for detector-1 and detector-2, respectively, under the illumination of 685 nm at an applied bias of -3 V . Fig. 7(b) shows the spectral response of the detectors in the wavelength range of $300\text{--}1300 \text{ nm}$, measured at an applied bias of -3 V with light intensity of 0.1 mW cm^{-2} . The detectors showed a peak detection at around $\lambda = 685$ and 900 nm , corresponding to the optical absorption edge of In_2Se_3 and Si , respectively. The values of R_λ for detector-1 and detector-2 were ~ 9 and 27 A W^{-1} , respectively, under the illumination of 900 nm at an applied bias of -3 V . In addition, the detectivity (D^*) of the photodetectors is estimated using

the relation,^{38,41} $D^* = R_\lambda / (2eJ_d)^{1/2}$, where R_λ is the responsivity of the photodetectors and J_d is the dark current density. Fig. 7(c) shows the voltage dependent detectivity of detector-1 and detector-2 under the illumination of 685 nm , which shows a maximum detectivity at an applied bias of -1.5 V . The D^* values of detector-1 and detector-2 were found to be $\sim 1.18 \times 10^{12}$ and $\sim 2.55 \times 10^{12} \text{ cm}^2 \text{ Hz}^{1/2} \text{ W}^{-1}$, respectively, at an applied bias of -1.5 V . Moreover, the sensitivity of the detectors, as defined by the ratio of photocurrent to the dark current, is calculated using the relation,^{39,41} $\text{Sensitivity} = I_{\text{ph}}/I_{\text{dark}}$, where I_{ph} and I_{dark} are the photocurrent and dark current of the detector, respectively. The voltage dependent sensitivity of the detectors is shown in Fig. 7(d), which shows a maximum sensitivity at an applied bias of -1 V . The sensitivities of the detectors were found to be 17.5 and 26.3 for detector-1 and detector-2 at an applied bias of -1 V , respectively.

In order to further investigate the photo-response characteristics of the detectors, the temporal response of the devices has been investigated. Fig. 8(a) depicts the temporal response of the detectors under the illumination of 685 nm at an



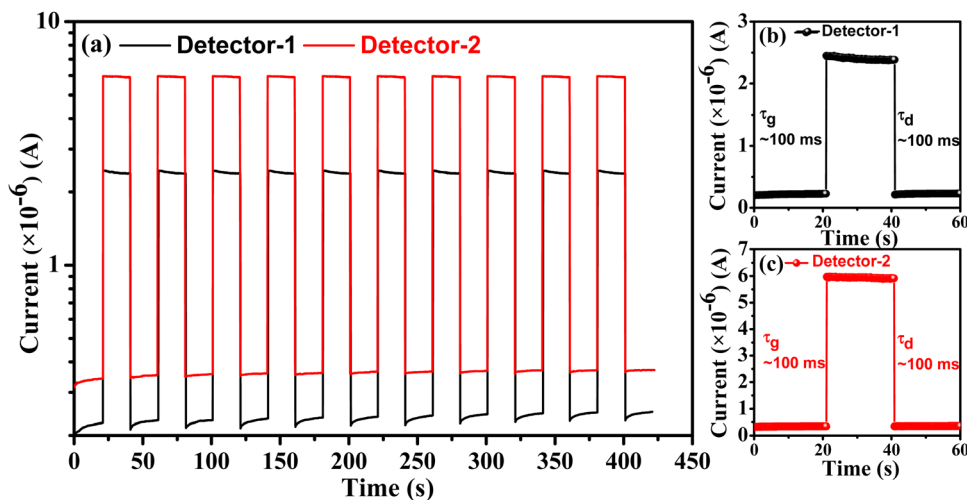


Fig. 8 (a) The temporal response of the detectors under the illumination of 685 nm at an applied bias of -3 V on semi-log scale, (b) zoomed-in view of one ON/OFF cycle of detector-1, and (c) zoomed-in view of one ON/OFF cycle of detector-2.

applied bias of -3 V. The temporal response remains identical after several cycles, confirming the excellent stability of the photodetectors. At the onset of illumination of 685 nm, the current increases and saturates within the response time. After the illumination is tuned off, the current returns to its initial value within the recovery time. Here, the response and

recovery times of the detectors were found to be ~ 100 ms, which is not accurate as the time interval between two data points is ~ 100 ms.

In order to estimate the precise response and recovery times, we have used an ultrafast time measurement setup, which is shown in Fig. 9(a). The transient response of the detectors

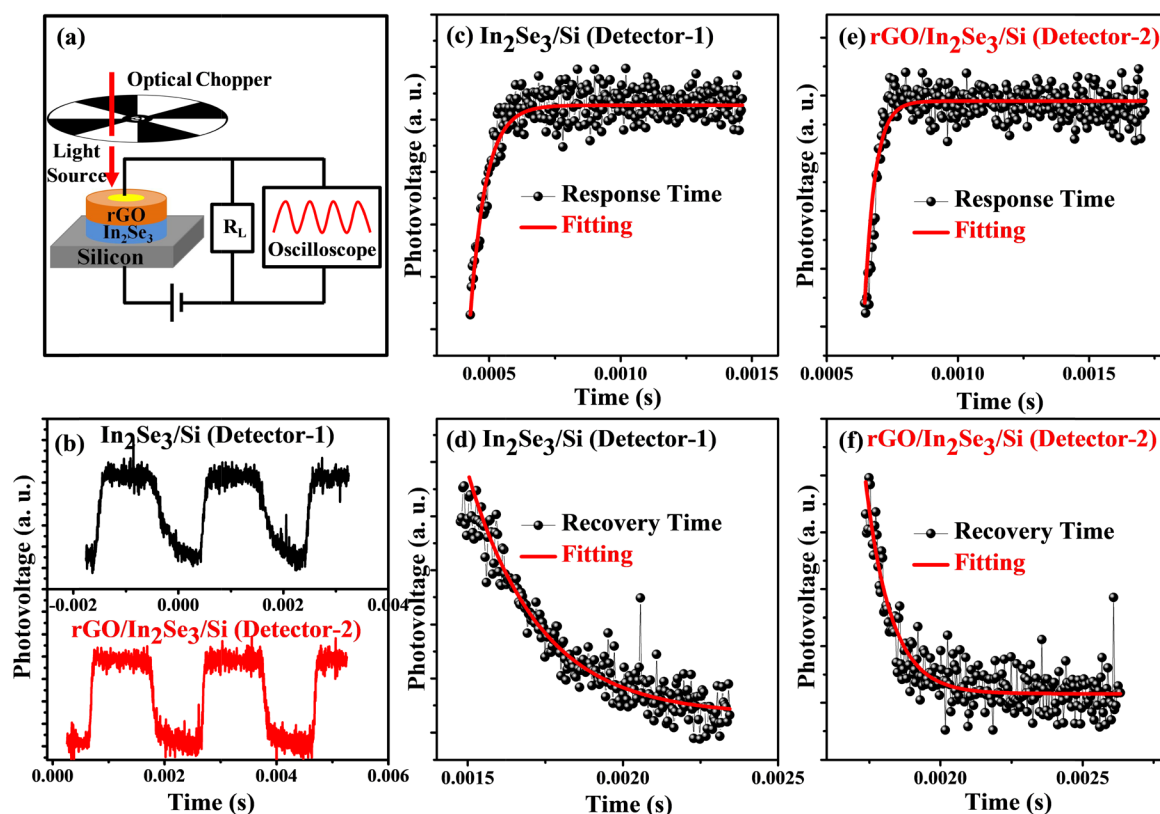


Fig. 9 (a) An ultrafast time measurement setup, (b) the transient response of the detectors under a pulsed light source of 685 nm with a frequency of 500 Hz, and (c)–(f) the response and recovery of detector-1 and detector-1 under a pulsed light source of 685 nm with a frequency of 500 Hz.



Table 1 Comparison of figures-of-merit of our device with the various 2D-layered heterostructure-based photodetectors reported in the literature, considering the significance of large area deposition of 2D materials for large scale production

Photodetector	Bias voltage (V)	λ (nm)	R_{λ} (A W^{-1})	D^* (Jones)	τ_g/τ_d (ms)
$\gamma\text{-In}_2\text{Se}_3/\text{n-Si}$ ¹⁷	0	808	0.57	2.6×10^{12}	0.035/0.115
Graphene/ In_2Se_3 ²⁸	0.35	1550	1.17	—	1.76/1.86
$\text{In}_2\text{S}_3/\text{CIGS}$ ⁴³	−1	870	2.06	2.3×10^{11}	—
$\text{n-In}_2\text{S}_3/\text{p-Si}$ ⁴⁴	2	750	0.68	2.5×10^{11}	—
$\text{MoS}_2/\text{p-Si}$ ⁴⁵	0	532	0.253	1×10^9	85/136
$\text{n-MoS}_2/\text{n-Si}$ ⁴⁶	−2	650	11.9	2.1×10^{10}	0.030/0.071
MoS_2 Schottky MSM ⁴⁷	10	532	0.57	1×10^{10}	0.07/0.11
MoS_2 Schottky MSM ⁴⁸	3	532	0.55	—	0.2/1.7
WS_2/Si ⁴⁹	0	980	0.224	1.5×10^{12}	0.016/0.029
rGO/ $\text{In}_2\text{Se}_3/\text{Si}$	−3	$\lambda = 685$	$R_{\lambda} = 9.5$	2.39×10^{12}	0.04/0.09
(This work)		$\lambda = 900$	$R_{\lambda} = 27$		

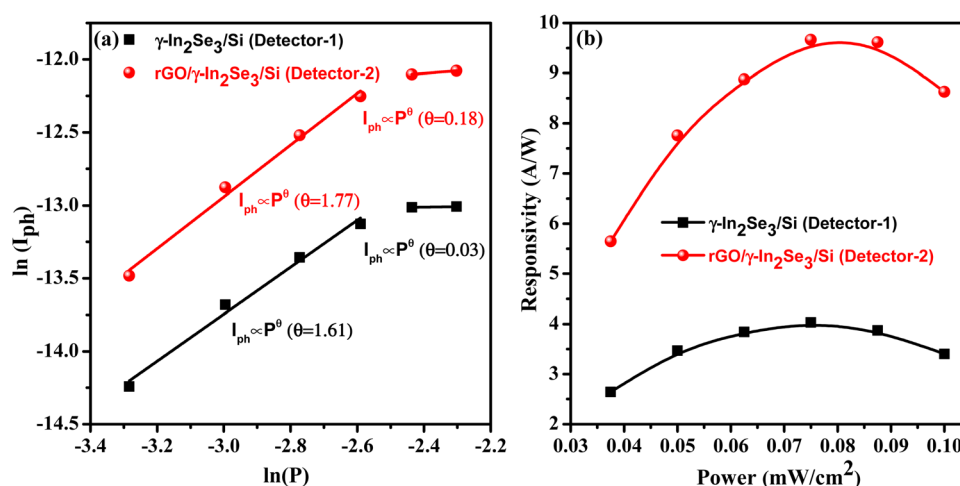


Fig. 10 (a) The variation of the photocurrent of the detectors with illuminated light intensity at an applied bias of -3 V and (b) the variation of the photo-responsivity of the detectors with illuminated light intensity at an applied bias of -3 V.

using an ultrafast time measurement under a pulsed light source of 685 nm with a frequency of 500 Hz is shown in Fig. 9(b). The time constants of the detectors were estimated by fitting the transient response with the following equations,^{41,42}

$$V(t) = V_{\text{dark}} + A\{1 - \exp[(t_0 - t)/\tau_g]\} \quad (1)$$

and

$$V(t) = V_{\text{dark}} + A\{1 - \exp[(t_0 - t)/\tau_d]\} \quad (2)$$

where V_{dark} is the dark photo-voltage, A is the scaling constant, t_0 is the time when the illumination was switched ON or OFF and τ_g/τ_d are the response/recovery times, respectively. The fitting of the transient responses with eqn (1) and (2) for detector-1 and detector-2 is shown in Fig. 9(c)–(f). In detector-1 and -2, the response/recovery times were 60/230 μs and 40/90 μs , respectively. In the present case, the fast temporal response in detector-2 could be due to the efficient hole transporting properties of the top rGO layer and the presence of a high internal electric field at the $\text{In}_2\text{Se}_3/\text{Si}$ interface. In Table 1, we have summarized the key figures-of-merit of our device and compared it with the various 2D-layered/Si heterostructure-based photodetectors reported in the literature,

considering the significance of large area deposition of 2D materials for large scale production.

The variation of the photocurrent with illuminated light intensity at an applied bias of -3 V is shown in Fig. 10(a). The photocurrent of the detectors increases with increasing light intensity with a power law,⁵⁰ $I_{ph} \propto P^\theta$, where I_{ph} is the photocurrent of the detectors and θ is an exponent which determines the conduction process of the photo-generated charge carriers. In Fig. 10(a), we observed two linear regions (region-I and region-II). In region-I (where, the power density of the illuminated light is low), the values of θ are 1.61 and 1.77 for detector-1 and detector-2, respectively. The high value of θ (greater than unity) in region-I suggests the effective separation of photogenerated charge carriers across the depletion region,⁵¹ which resulted in an enhance photocurrent and thereby the responsivity of the detector is increased as illustrated in Fig. 10(b). In region-II (where, the power density of the illuminated light is high), the values of θ are 0.03 and 0.18 for detector-1 and detector-2, respectively. Here, the non-unity value of θ in region-II suggests a complex recombination or scattering of the photogenerated charge carriers due to more population density of the photogenerated charge carriers,^{52,53}

which resulted in a decrease in the responsivity of the detectors, as shown in Fig. 10(b).

IV. Conclusion

In conclusion, we have demonstrated a high performance rGO decorated γ -In₂Se₃/Si heterostructure-based broadband photodetector ranging from 400–1200 nm. The rGO/ γ -In₂Se₃/Si heterostructure showed an enhanced photo-responsivity of ~ 9.5 and ~ 27 A W⁻¹ under the illumination of 685 and 900 nm, respectively, at an applied bias of -3 V. In addition, the heterostructure showed high sensitivity with fast response/recovery times (40/90 μ s). The enhanced figures-of-merit of the rGO/ γ -In₂Se₃/Si heterostructure are due to the increase in optical absorption of incident light and efficient hole transporting properties of the top rGO layer, which effectively reduces the recombination of the photogenerated charge carriers.

Author contributions

Prof. S. B. Krupanidhi, Prof. K. K. Nanda, Dr Sujit Das and Dr Basanta Roul generated the idea and designed experiments. Dr Basanta Roul and A. M. C. performed experiments. Dr Basanta Roul, M. K., and K. L. K., prepared the manuscript. All authors participated in the discussion of the experimental results and revision of the manuscript.

Conflicts of interest

The authors declare no conflicts of interest.

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