



# Surface-Plasmon-Induced Ag Nanoparticles Decorated In<sub>2</sub>O<sub>3</sub> Nanowires for Low Noise Photodetectors

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## Abstract

Silver (Ag) nanoparticles (NPs) were synthesized by glancing angle deposition (GLAD) technique on indium oxide (In<sub>2</sub>O<sub>3</sub>) nanowires (NWs) over n-type Si substrate. The In<sub>2</sub>O<sub>3</sub> NWs and Ag NPs were morphologically characterized by field emission scanning electron microscopy (FESEM) and electron dispersive spectroscopy (EDS). The complete growth of In<sub>2</sub>O<sub>3</sub> NWs was observed by high-resolution transmission electron microscopy (HRTEM) and corresponding selected area electron diffraction (SAED) pattern was also studied. The structural analysis was done by high-resolution X-ray diffraction (HRXRD), and relevant peaks were identified to calculate the crystalline size. The HRXRD patterns displayed the peak for Ag NPs and monoclinic crystal structure of Ag<sub>3</sub>O<sub>4</sub>. The optical properties were analyzed by photoluminescence (PL) emission spectrums. The presence of Ag NPs over In<sub>2</sub>O<sub>3</sub> NWs reduced the PL intensity. Atomic force microscopy (AFM) was also studied to estimate the surface roughness for both the samples. The semi-logarithmic *I-V* (*ln(I)-V*) characteristics revealed the enhancement in photoconduction for the n-Si/In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs device at  $-4.5$  V using a 100-W tungsten filament source. The total  $\sim 2.6$  fold enhancement in photosensitivity were recorded for the n-Si/In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs device at an applied voltage of  $-2.4$  V. This n-Si/In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs device possessed high detectivity and low noise equivalent power (NEP) as compared with the bare n-Si/In<sub>2</sub>O<sub>3</sub> NWs device.

**Keywords** GLAD · In<sub>2</sub>O<sub>3</sub> nanowires · Ag nanoparticles · Surface plasmon · Photodetector

## Introduction

In the field of optoelectronics technology, indium oxide (In<sub>2</sub>O<sub>3</sub>) is a most frequently used direct bandgap ( $\sim 3.6$  eV) semiconductor material [1, 2]. Many nanocrystalline oxides materials like TiO<sub>2</sub>, ZnO In<sub>2</sub>O<sub>3</sub> etc. are broadly used because of their remarkable physical and chemical properties [3–6]. Among all the metal oxides, In<sub>2</sub>O<sub>3</sub> shows flawless advantages in optoelectronics applications like solar cells [7], sensors [8], photodetectors [9], organic light-emitting diodes (OLEDs) [10] etc., due to very good electrical transparency, synthesis method, large-area uniformity and mechanical flexibility. There are numerous physical techniques that have been introduced to fabricate the In<sub>2</sub>O<sub>3</sub> nanostructures like sputtering [11], laser ablation [12], electron beam evaporation [13] and thermal evaporation [14], which make it too costly to be

commercially produced. Earlier, It was reported that the plasmonic metal NPs like Cu [15], Au [16] and Ag [17] reduces the reflection of incident photons and it exhibits robust surface plasmon resonance (SPR) behaviour. Nevertheless, to controlling SPR, the behaviour size, shape and separation between two NPs assume a noteworthy role. Glancing angle deposition or GLAD is an attractive option for fabricating In<sub>2</sub>O<sub>3</sub> photodetectors because, in addition to being a reliable, sophisticated technique with in situ characterization and control, it is a cheap, one-step process. It has been already reported that Au/TiO<sub>2</sub> nanorod arrays having imminent applications in the efficient solar energy conversion [18]. NWs and NPs are very appealing for photodetectors for their high surface-to-volume ratio compared with their sizes and high controllability in growth with advanced techniques [19, 20]. In this research paper, the authors were fabricated In<sub>2</sub>O<sub>3</sub> NWs ( $\sim 400$  nm) and Ag NPs ( $\sim 10$  nm) based In<sub>2</sub>O<sub>3</sub> NWs over n-type silicon (Si) substrate using a double step glancing angle deposition (GLAD) cum electron beam (e-beam) evaporation technique. It has been shown that the plasmonic Ag NPs-based device is superior compared with the In<sub>2</sub>O<sub>3</sub> NW device in terms of photoconduction. Therefore, simply employing structural phenomenon, the optical properties of metal NPs can be

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transformed [21]. Gorurla et al. reported that Au-ZnO plasmonic device has greater photoresponsivity than Ag-GaN-based device [22]. Li et al. also reported that plasmonic enhancement by metal NPs deals with novel opportunities to progress the performance of optoelectronic devices [23]. So, the fabrication of metal NPs is relatively a very challenging task. Fabrication of metal NPs was reported by J. Fu et al. using GLAD technique [24]. Mehta et al. reported  $\text{In}_2\text{O}_3:\text{Ag}$  nanocomposite layers by chemical capping and dip-coating method for the application of gas sensing properties [25]. Zhou et al. also reported that the GLAD technique was an incredible and active technique for monitoring the morphology, thickness and shape of the nanostructured samples [26]. So, Ag NPs oriented  $\text{In}_2\text{O}_3$  NWs device is one of the effective and unique structure to develop photodetector in the optoelectronics applications, and to the best of our knowledge, no such reports have been published regarding the combined NPs-NWs structure using GLAD-assisted e-beam evaporation technique, thus making it more attractive with commercial off-the-shelf devices.

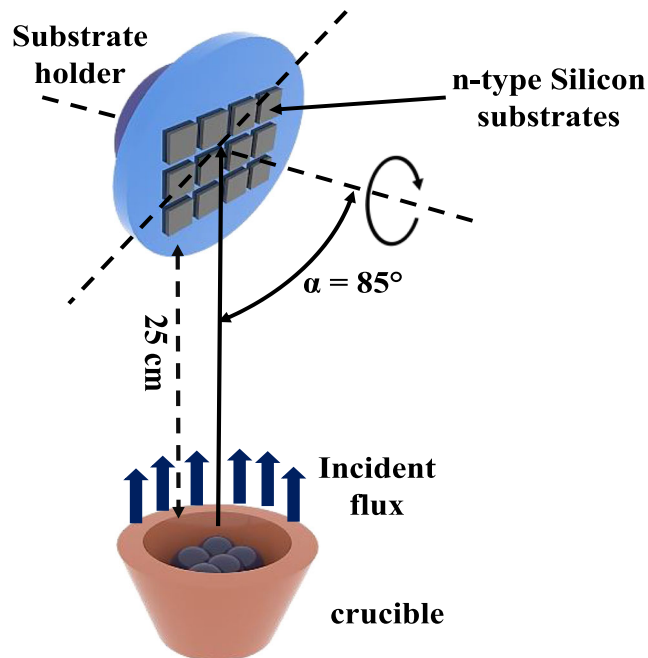
In this research paper, an easy and inimitable fabrication technique has been illustrated to synthesize Ag NPs coated  $\text{In}_2\text{O}_3$  NWs by GLAD technique. This technique was controlled inside the e-beam evaporator system. The growth of  $\text{In}_2\text{O}_3$  NWs and Ag NPs was observed and relevant information was extracted through FESEM and HRTEM analysis. Manifestation of Ag NPs and formation of  $\text{Ag}_3\text{O}_4$  compound was identified by HRXRD investigation. Further, the PL intensity was studied and subsequent degradation in PL intensity was observed. The surface roughness was calculated using AFM analysis. With all morphological and structural characteristics, the photoconduction properties were analysed through In contact, and relevant results were studied. The improvement in photoconduction was observed for the  $\text{In}_2\text{O}_3$  NWs/Ag NPs device and the improved detectivity, low noise equivalent power response was also recorded.

## Experimental

$\text{In}_2\text{O}_3$  NWs ( $\sim 400$  nm) (Highly pure 99.999%, MTI Corporation, USA) and Ag NPs ( $\sim 10$  nm) were deposited over the pre-cleaned Si wafer (Highly pure 99.999%, n-type,  $\langle 100 \rangle$ , 4" dia  $\times$  0.525 mm, bulk resistivity:  $10 \Omega \text{ cm}$ , sheet resistance:  $190 \Omega/\text{sq.}$ ) using a double-step GLAD technique by e-beam (Hind High Vacuum Co. (p) Ltd., 15F6) evaporator. The n-Si substrates were cleaned with tetrachloroethylene ( $\text{C}_2\text{Cl}_4$ ), methanol ( $\text{CH}_3\text{OH}$ ) and DI water ( $\text{H}_2\text{O}$ ). The solution of hydrofluoric acid (HF) + DI water ( $\text{H}_2\text{O}$ ) was used at a volume ratio of 1:50 to etch native oxide from the n-Si substrate. The vacuum chamber pressure of  $\sim 2 \times 10^{-5}$  mbar and the deposition rate of  $1.2 \text{ \AA s}^{-1}$  are maintained during entire experimental work. The highly pure  $\text{In}_2\text{O}_3$  and Ag

pellets were placed perpendicularly  $\sim 25$  cm from the substrate in the vacuum chamber with an azimuthal angle of  $85^\circ$  and GLAD rotation of  $\sim 130$  and  $\sim 460$  rpm was maintained for both the depositions, respectively.  $\text{In}_2\text{O}_3$  and Ag pellets were evaporated between the melting current of  $\sim 10$  mA to 40 mA. Here, Fig. 1 shows the full GLAD setup for the fabrication of  $\text{In}_2\text{O}_3$  NWs and Ag NPs inside the e-beam evaporator. Total five successful depositions were carried out to deposit the  $\sim 400$  nm of vertical  $\text{In}_2\text{O}_3$  NWs. Indium (In) (Highly pure 99.999%, MTI Corporation, USA) contact has been fabricated on the GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs sample at a melting current less than 30 mA through the metal mask hole. The area of deposited In contact was  $\sim 1.96 \times 10^{-7} \text{ m}^2$  and thickness of the contacts was  $\sim 33$  nm. The two devices thus fabricated are n-Si/ $\text{In}_2\text{O}_3$  NW/In and n-Si/ $\text{In}_2\text{O}_3$  NW/Ag NP/In.

The samples were morphologically characterized by field emission scanning electron microscopy (FESEM, Carl-ZEISS) and energy dispersive spectroscopy (EDS, Carl-ZEISS). The growth of NWs was characterized by high-resolution transmission electron microscopy (HRTEM, JEM-2100) and corresponding selected area electron diffraction (SAED) was done on the sample. The atomic force microscopy (AFM, Bruker Multimode 8) and high-resolution X-ray diffraction (HRXRD, Bruker D8 Advanced, Cu-K $\alpha$  radiation,  $1.542 \text{ \AA}$ ) were done to study the structural characterization. The room temperature ( $T = 300 \text{ K}$ ) PL study was done on the samples using Hitachi F-7000 Fluorescence Spectrophotometer with an excitation wavelength of 350 nm to study the optical characteristics. The photoconduction analysis was done using a Keysight B2902A source and



**Fig. 1** Experimental setup of GLAD for the fabrication of  $\text{In}_2\text{O}_3$  NWs and Ag NPs inside the e-beam evaporator

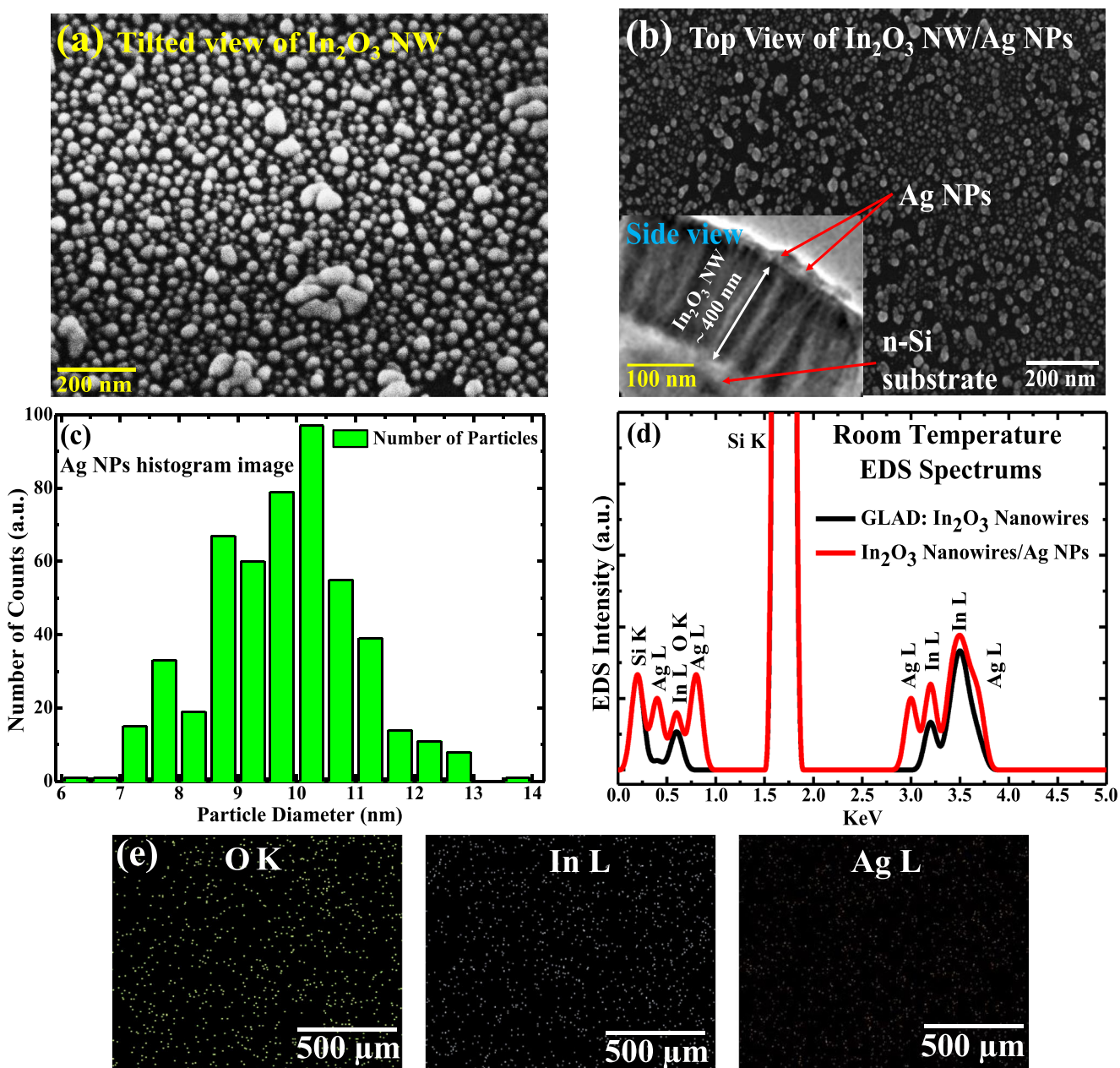
measurement unit (SMU) on the devices using a 100-W tungsten filament source.

## Results and Discussions

### FESEM and EDS Analysis

The FESEM images were displayed to study the morphological and structural characteristics of the fabricated samples. Figure 2(a), (b) and inset Fig. 2(b) show the 55° tilted view of In<sub>2</sub>O<sub>3</sub> NWs, top view of Ag NPs and side view of Ag NPs

decorated In<sub>2</sub>O<sub>3</sub> NWs. The careful observation of FESEM images confirms the Ag NPs were densely filled and evenly distributed all over the surface. Likewise, the 55° tilted view also reveals the much attired and uneven formation of In<sub>2</sub>O<sub>3</sub> NWs over the n-type substrate. The individual NWs and Ag NPs are interconnected to each other due to the high substrate temperature [20]. The average diameter of the fabricated single In<sub>2</sub>O<sub>3</sub> NW was calculated around ~24 nm from the Fig. 2(a). In GLAD technique, the Ag NPs were fabricated due to self-atomic shadowing effect on the arbitrarily deposited seeds. Though, this type of physical evaporation technique is a very classy technique, due to GLAD rotation speed (~130



**Fig. 2** (a) Tilted view FESEM images of In<sub>2</sub>O<sub>3</sub> NWs, (b) Top view of the FESEM images of Ag NPs (inset side view of Ag NPs decorated In<sub>2</sub>O<sub>3</sub> NWs), (c) Ag NPs size histogram image, (d) EDS analysis and (e) Colour mapping

and ~460 rpm for In<sub>2</sub>O<sub>3</sub> NWs and Ag NPs), deposition angle, evaporation rate, substrate temperature etc. [13, 27].

According to Fick's law, diffusion coefficient ( $D_{\text{coeff}}$ ) can be found from the equation;  $D_{\text{coeff}} \propto \exp(-E_d/K_b T_s)$ , where,  $E_d$  is called diffusion energy,  $K_b$  is Boltzmann constant and  $T_s$  is the temperature of the substrate. The substrate temperature (room temperature) effects in a large diffusion coefficient, which may be the reason for the formation of In<sub>2</sub>O<sub>3</sub> NWs over the substrate [20]. R. Lahiri et al. reported that the formation of different length In<sub>2</sub>O<sub>3</sub> NWs due to inherent atomic shadowing property by GLAD technique [28]. A low deposition rate of 1.2 ÅS<sup>-1</sup> and substrate temperature (room temperature) were maintained for the deposition of Ag NPs, thus formed the connected Ag NPs throughout the sample surface due to the large diffusion coefficient of Ag molecules [20]. Figure 2(c) indicated the particle size histogram image, which is extracted from Fig. 2(b). This histogram image shows that the Ag NPs consists of different NPs size ranging from ~6 to ~14 nm. The maximum number of particles diameter was measured between the sizes of ~10 to ~11 nm. Q. Zhou et al. and G. Wang et al. reported that the high substrate temperature (room temperature) indicated thick and connected nanostructures likewise low substrate temperature (approximately -40 °C) indicated thin and separated nanostructures [29, 30].

To inspect the presence of deposited materials, room temperature EDS analysis was performed. Figure 1(d) established the EDS spectra for GLAD: In<sub>2</sub>O<sub>3</sub> NWs and In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs. The elements like silicon (Si), indium (In), silver (Ag) and oxygen (O) molecules are present on the samples. Apart from In, Ag and O, no other extra peak for any other element has been found. EDS spectrum also shows the emission from In L, Ag L and O K shells with the colour mapping of EDS images of the samples depicted in Fig. 2(e), where oxygen (O) shows the green colour, indium (In) shows the blue colour and silver (Ag) shows the orange colour. Moreover, there was a high peak emission from K shell to Si substrate.

## HRTEM and HRXRD Analysis

To observe the growth of vertical NWs and examine the microstructural morphologies of the same HRTEM (JEM-2100), analysis has been done. A pull-out portion of single NW was depicted in Fig. 3(a), which shows the classic growth In<sub>2</sub>O<sub>3</sub> NWs having an average top diameter of ~25 nm and bottom diameter of ~16 nm. The dusky part of the single NW was due to the existence of silicon, and the dazzling part indicates the presence of oxygen molecules. Figure 3(b) displays the selected area electron diffraction (SAED) pattern which did not contain any fringe spacing and that revealed the amorphous nature of the NW [31]. The 'd' spacing of the crystal plane was found from the SAED pattern, and it was obtained as 2 Å, 1.1 Å and 0.8 Å (JCPDS 653170).

To investigate the structural characteristics, HRXRD profile was obtained using Bruker D8 Advanced using Cu-K $\alpha$  radiation source, the wavelength of 1.54 Å in the  $2\theta$  range of 30 to 75° for GLAD: In<sub>2</sub>O<sub>3</sub> NWs and In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs, displayed in Fig. 3(c). The bare GLAD: In<sub>2</sub>O<sub>3</sub> NWs shows the amorphous nature of the sample, which was also revealed in HRTEM analysis. The HRXRD diffraction peaks for GLAD: In<sub>2</sub>O<sub>3</sub> NWs were emerging at  $2\theta = 38.26^\circ$ ,  $44.47^\circ$  and  $64.85^\circ$  with the corresponding plane of (400), (422) and (444), respectively. All the peaks were matching with the JCPDS data (JCPDS card no. 06–0416) [32]. The In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs shows same diffraction peak with a new diffraction peak at  $2\theta = 38.06^\circ$  due to the presence of Ag NPs which corresponds (111) plane (JCPDS card no. 04–0783) [33, 34]. Overall 50% magnified graph of Fig. 3(c) shows a clear image of a diffraction peak at  $2\theta = 38.06^\circ$  displayed in Fig. 3(d). Likewise, its revealed another diffraction peak at  $2\theta = 50.68^\circ$  corresponds to (031) plane (JCPDS card no. 84–1261) for the monoclinic crystal structure of Ag<sub>3</sub>O<sub>4</sub> [35]. The formation of the Ag–O compound shows (031) diffraction plane which indicated the monoclinic crystal structure, which was formed due to oxygen attached with the Ag NPs by attracting the oxygen molecules [36]. The average crystalline size was calculated ~38 and ~87 nm for the bare GLAD: In<sub>2</sub>O<sub>3</sub> NWs and In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs, respectively, using Debye Scherrer equation, which was shown in equation 1.

$$D = (0.9\lambda)/(\beta^*\cos\theta) \quad (1)$$

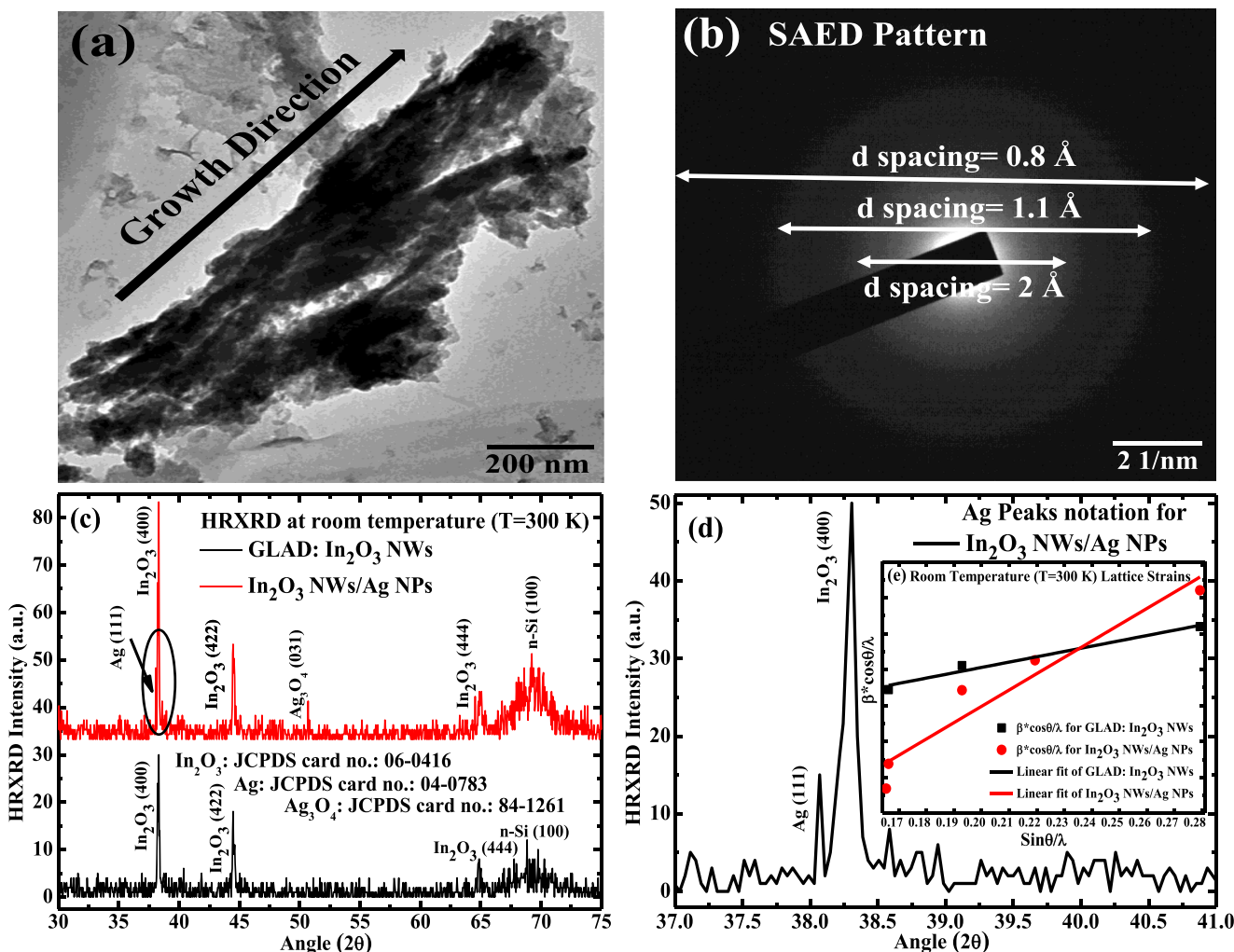
where  $D$  is crystalline size,  $\lambda$  is the wavelength of the incident X-ray ( $K_\alpha$ ),  $\beta$  is the full-width half maximum (FWHM) in radians of Gaussian distribution of the HRXRD data and  $2\theta$  is the Bragg's angle measured in radians. The authors have reported the method of extraction of crystalline size [37]. The average crystalline size was increased for In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs due to the difference in atomic radius of Ag (144 pm) and In (167 pm). The tendency for agglomeration of Ag NPs during the process in the high ambient temperature of the chamber was maybe the reason for the increase in dimension. Lattice strain was also calculated ~0.0073 and ~0.0221 for both the same samples displayed in Fig. 3(e) (inset of Fig. 3(d)) using Williamson-Hall ( $W-H$ ) plot equation, which was shown in equation 2.

$$\beta^*\cos\theta/\lambda = k/D + 4\epsilon\sin\theta/\lambda \quad (2)$$

where  $k$  is the shape factor,  $\epsilon$  is lattice strain and  $2\theta$  is the Bragg's diffraction angle. Correspondingly in that case, the lattice strain was increased, and it offers the work hardening process in the material [38].

## PL Emission

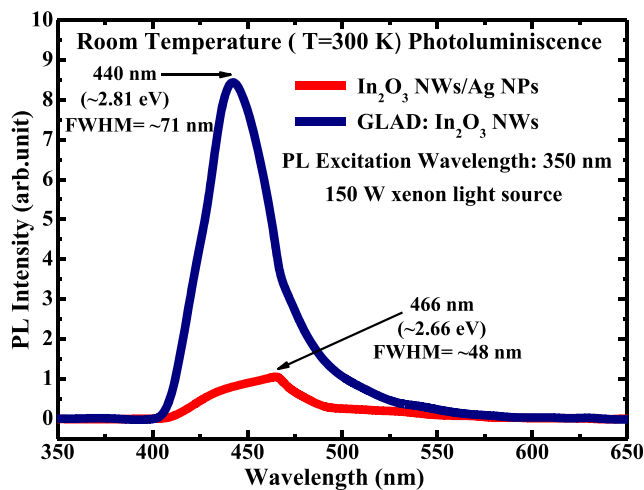
Room temperature PL emission has been done between the wavelengths of 350 and 650 nm using an excitation



**Fig. 3** (a) HRTEM analysis of  $\text{In}_2\text{O}_3$  NWs, (b) SAED pattern of  $\text{In}_2\text{O}_3$  NWs, (c) HRXRD spectrums of GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs (d) 50% magnified spectrums of  $\text{In}_2\text{O}_3$  NWs/Ag NPs and (e) Lattice strain of GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs (inset of Fig. 3(d))

wavelength of 350 nm by Hitachi F-7000 Fluorescence Spectrophotometer with a Xenon light source of 150 watt for the bare GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs samples presented in Fig. 4.

The GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs indicated the PL emission spectrums at 440 nm ( $\sim 2.81$  eV) and 466 nm ( $\sim 2.66$  eV) with the full-width half maximum (FWHM) of  $\sim 71$  nm and  $\sim 48$  nm, respectively.  $\text{In}_2\text{O}_3$  shows the PL emission spectrum between 400 and 500 nm due to the existence of oxygen vacancies in the material [39]. The blue region emission arises at 440 nm for the GLAD:  $\text{In}_2\text{O}_3$  NWs attributed new energy levels in the forbidden energy gap. This type of emission occurred due to the recombination of electron-hole pairs at the surface of the samples. Zheng et al. reported that the PL emissions of 429 and 460 nm occurred for the  $\text{In}_2\text{O}_3$  NWs samples [40]. It was apparent that the high PL intensity manifested a higher rate of recombination of electro-hole pairs and the low PL intensity indicates a lower rate of recombination of electro-hole pairs. Additionally, The Ag NPs created a



**Fig. 4** PL emission spectrum for GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs using an excitation wavelength of 350 nm

precise bond with the GLAD:  $\text{In}_2\text{O}_3$  NWs and formed an inorganic silver compound particle like  $\text{Ag}_3\text{O}_4$ . This type of inorganic silver compound was observed in the room temperature ( $T = 300$  K) through HRXRD analysis. These  $\text{Ag}_3\text{O}_4$  particles act as trap particles to capture the photo-induced electrons in the forbidden energy gap and thus defeat the recombination of electron and holes [41]. In the experiment, a decrease in  $\sim 8.5$ -fold PL intensity was observed for the  $\text{In}_2\text{O}_3$  NWs/Ag NPs sample due to the aforementioned reason. Here, with the reduction in PL intensity, a  $\sim 26$ -nm red shift had been observed for the  $\text{In}_2\text{O}_3$  NWs/Ag NPs samples as compared with that of the bare GLAD:  $\text{In}_2\text{O}_3$  NWs samples due to the SPR effect. Kim et al. reported a similar type of emission due to oxygen-related defect states [39]. Fong et al. also reported that the Ag NPs attached with  $\text{In}_2\text{O}_3$  NWs and produced SPR effect, which shifted the resonance peak to longer wavelength [42]. Conclusively, the presence of Ag NPs over  $\text{In}_2\text{O}_3$  NWs shifted the resonance peak and enhanced the emission at  $\sim 466$  nm. Therefore, the PL emission analysis delivered the SPR effect using the Ag NPs over the  $\text{In}_2\text{O}_3$  NWs.

### AFM Analysis

Figure 5(a) and 5(b) shows the three dimensional (3D) AFM images for the bare GLAD:  $\text{In}_2\text{O}_3$  NWs (height:  $\sim 400$  nm) and  $\text{In}_2\text{O}_3$  NWs/Ag NPs samples. The overall scan area of  $10 \mu\text{m} \times 10 \mu\text{m}$  and resolution of  $2 \mu\text{m}$  were maintained for both the samples. The Ag NPs were milled up with  $\text{In}_2\text{O}_3$  NWs and created Ag NPs– $\text{In}_2\text{O}_3$  layer because of the electrostatic charge-charge interaction between metal-semiconductor junctions [43]. The root mean square surface roughness ( $R_p$ ) was calculated  $\sim 6.10$  nm for the GLAD:  $\text{In}_2\text{O}_3$  NWs sample and  $\sim 5$  nm for  $\text{In}_2\text{O}_3$  NWs/Ag NPs sample, which will be reduced by  $\sim 1.22$ -fold. The packing density of  $\text{In}_2\text{O}_3$  nanostructures was predisposed by the newly formed Ag NPs over

$\text{In}_2\text{O}_3$  NWs that leads to the reduction in surface roughness. Antonio Ruiz Puigdollers and others also reported that the same theory on the formation of Ag and Au clusters over the different materials [44].

### Photoconduction Analysis

To study the electrical behaviour of bare GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs samples, indium (In) metallization was performed in the e-beam chamber. Figure 6(a) and 6(b) shows the 3-D layout design of the bare n-Si/ $\text{In}_2\text{O}_3$  NWs/In (device-1) and n-Si/ $\text{In}_2\text{O}_3$  NWs/Ag NPs/In (device-2) devices. Figure 6(c) displays the room temperature semi-logarithmic  $I$ - $V$  ( $\ln(I)$ - $V$ ) characteristics for both the devices using B2902A source and measurement unit (SMU) and a 100-W tungsten filament source. In this case, almost no changes in dark current conduction and trifling changes in light current conduction had been observed under the forward bias condition. Likewise, under reverse bias mode, a low dark current conduction was observed for device-2, moreover light current for device-2 ( $\sim 4$  mA) was higher than device-1 ( $\sim 3.2$  mA). The enhancement in light current may be due to trapping of electrons at the interface of the  $\text{In}_2\text{O}_3$  NWs/Ag NPs followed by trapping and de-trapping process of electrons [45, 46]. Trapping of electrons at the interface was started under dark current measurement likewise de-trapping of electrons was started under light current measurement. The enhancement of current attributed to the abovementioned de-trapping process and due to oxygen-related trap states, it reduces the depletion region in the structures and allows the tunnelling of electrons [47]. Lee et al. also reported a similar type of observation in their research work [45]. The ideality factor ( $\eta$ ) was calculated  $\sim 20$  and  $\sim 11$  for device-1 and device-2, respectively, from the slope and intercept of semi-logarithmic forward bias of  $I$ - $V$  characteristics, represented

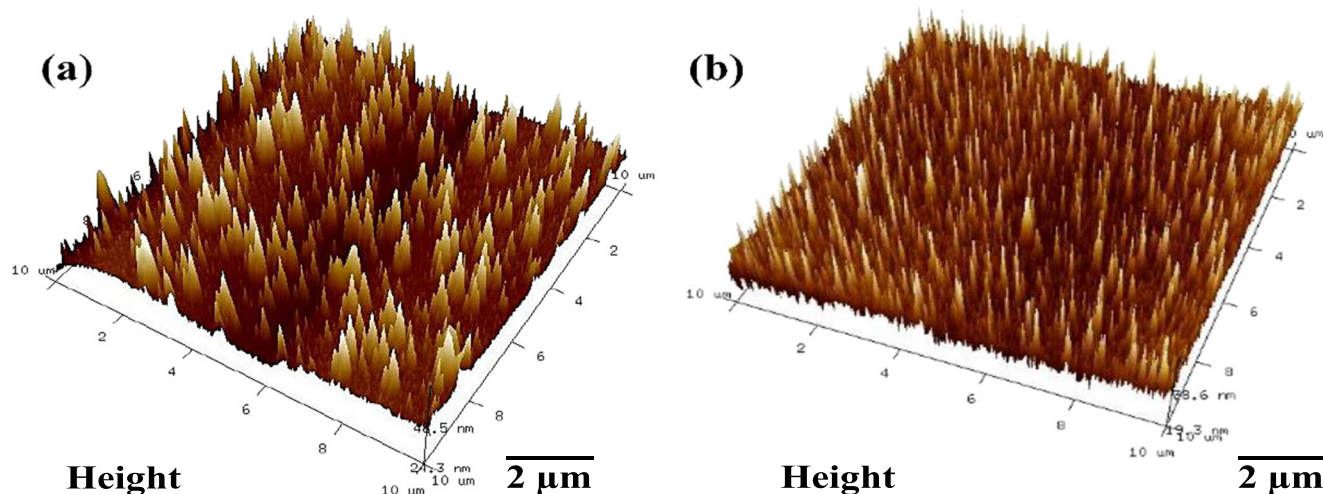
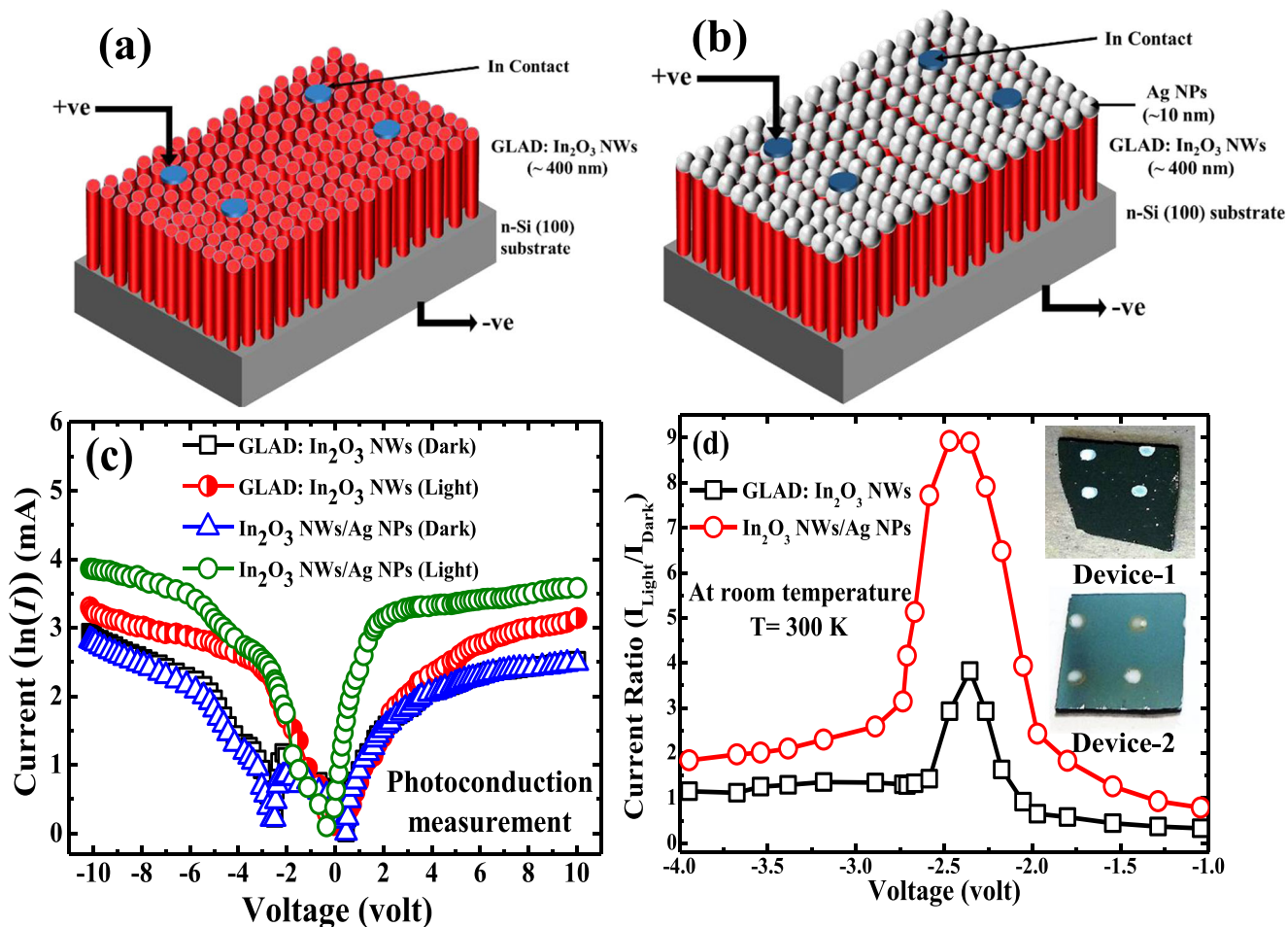


Fig. 5 AFM images of (a) GLAD:  $\text{In}_2\text{O}_3$  NWs and (b)  $\text{In}_2\text{O}_3$  NWs/Ag NPs



**Fig. 6** 3-D Layout design of (a) GLAD: In<sub>2</sub>O<sub>3</sub> NWs device (device-1), (b) In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs (device-2), (c) Semi-logarithmic current (*I*)-voltage (*V*) characteristics for GLAD: In<sub>2</sub>O<sub>3</sub> NWs and In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs, (d) Current ratio (*I*<sub>light</sub>/*I*<sub>dark</sub>)-voltage graph for both the devices

in Fig. 5(c). The method of extraction of the ideality factor for a diode was reported by Devi et al. using equation-3,

$$\eta = \frac{q}{kT} \frac{1}{\frac{\partial(\ln(I))}{\partial V}} \tag{3}$$

where *k* is called Boltzmann constant, *T* is absolute temperature, *q* is an electron charge and  $\frac{\partial(\ln(I))}{\partial V}$  is the slope of the semi-logarithmic *I*-*V* curve [48, 49].

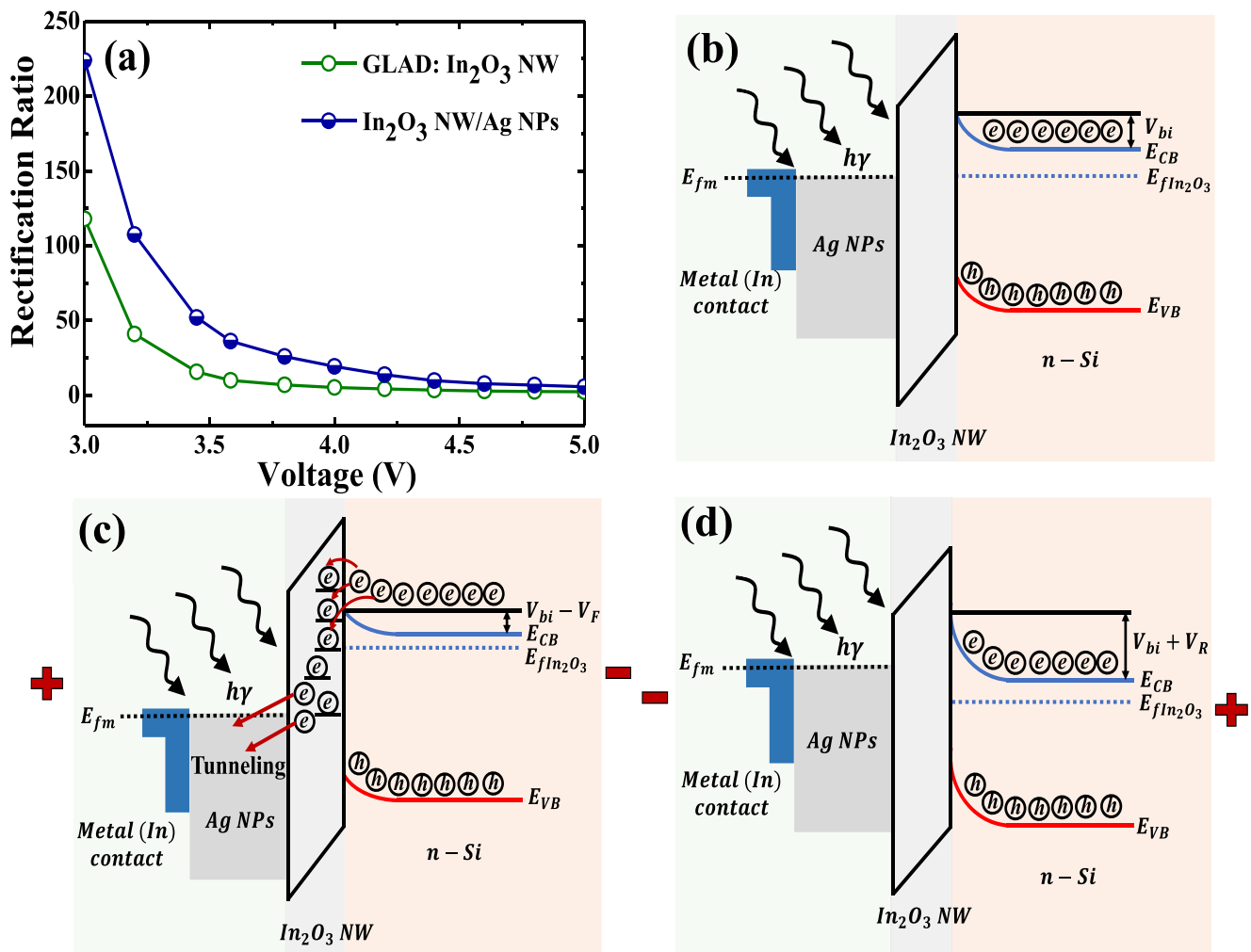
The photosensitivity (*I*<sub>light</sub>/*I*<sub>dark</sub>) for both the devices was calculated in reverse bias mode from the applied voltage of − 1 to − 4 V, displayed in Fig. 6(d). At − 2.4 V applied bias, ~ 2.6-fold enhancement was observed for the device-2 as compared with that of the device-1. This enhancement in photosensitivity was due to the SPR effect and a sufficient number of dangling bonds at the surface of In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs device, which can easily attract the oxygen molecules and hence the maximum electron tunnelling happened [47]. The SPR effect and dangling bonds enhanced the oxygen molecules over the

surface of device-2, and thus, photosensitivity was increased. A clear colour change from deep blue to light blue confirms the formation of Ag NPs on the device-2 (Fig. 6(d), inset).

The barrier height was calculated of ~ 0.6 eV and ~ 0.5 eV for bare GLAD: In<sub>2</sub>O<sub>3</sub> NWs device and In<sub>2</sub>O<sub>3</sub> NW/Ag NPs device, respectively, at zero bias voltage using the following equation.

$$J_0 = A^* T^2 \exp \left[ \frac{-q\phi_B}{kT} \right] \tag{4}$$

where *J*<sub>0</sub> is the reverse saturation current corresponding to zero bias, *A*<sup>\*</sup> is the Richardson constant, *k* is the Boltzmann constant,  $\phi_B$  is the zero-bias barrier height. The decrease in barrier height was attributed to the reduction in the depletion region and tunnelling of charge carriers through the device [50]. The rectification ratio (*R*<sub>ratio</sub>) of the two devices was calculated to investigate the device performance, depicted in inset Fig. 7. The *R*<sub>ratio</sub> was defined as  $\frac{|I_F|}{|I_R|}$ , where *I*<sub>F</sub> and *I*<sub>R</sub> are forward and reverse dark current. At ~ 3 V, the *R*<sub>ratio</sub> was found to be maximum (approximately twofold) in case of



**Fig. 7** (a) Rectification ratio for GLAD: In<sub>2</sub>O<sub>3</sub> NWs and In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs devices. In<sub>2</sub>O<sub>3</sub> NWs/Ag NPs device band diagram for (b) zero bias, (c) forward bias and (d) reverse bias under light illumination

In<sub>2</sub>O<sub>3</sub> NW/Ag NPs device compared with the bare In<sub>2</sub>O<sub>3</sub> NW device, which shows better rectification capability. A high  $R_{\text{ratio}}$  exhibited good rectifying behaviour in dark illumination and attributed high electron mobility in the In<sub>2</sub>O<sub>3</sub> NW/Ag NPs device [51]. Figure 7(b-d) shows the band diagram for zero bias, forward bias and reverse bias under light illumination condition, respectively.

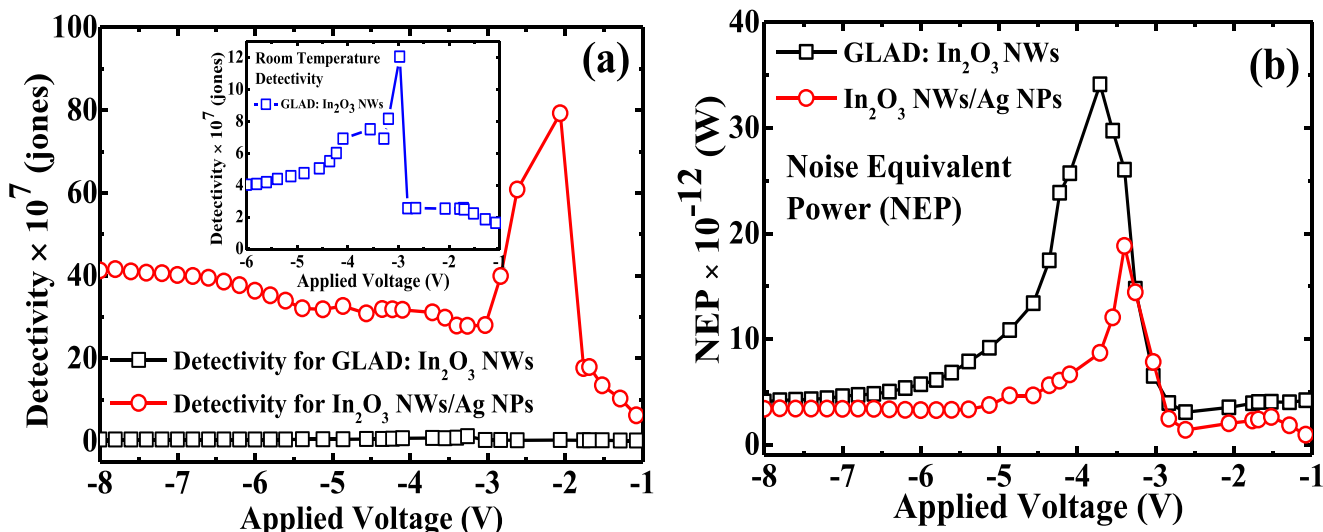
In case of forward bias, the n-Si and In<sub>2</sub>O<sub>3</sub> NW junction was connected to negative voltage whereas In and Ag NPs junction was connected to positive voltage. Under light illumination, the charge carriers can be highly separated by n-Si and In<sub>2</sub>O<sub>3</sub> NW junction, and there was a flow of photogenerated electrons through Ag NPs/In<sub>2</sub>O<sub>3</sub> NW junction which can be collected through In and Ag NPs junction. Furthermore, when the positive voltage was applied at In contact, In and Ag/In<sub>2</sub>O<sub>3</sub> interface states can trap the electrons at the conduction band (E<sub>CB</sub>), as seen in Fig. 7(c). This reduces the width of the depletion region, barrier height and hence trap-assisted tunnelling of photogenerated electrons happened, which enhanced the

photosensitivity [19]. Moreover, Ag has higher electronegative value (electronegativity: 1.93) thus attracts the oxygen molecules in Ag NPs/In<sub>2</sub>O<sub>3</sub> NW device. The attraction of oxygen by Ag from In<sub>2</sub>O<sub>3</sub> induces oxygen-assisted defect states in the Ag NPs/In<sub>2</sub>O<sub>3</sub> NW device [52]. In case of reverse bias (Fig. 7(d)), the n-Si and In<sub>2</sub>O<sub>3</sub> NW junction was connected to positive voltage whereas In and Ag NPs junction was connected to a negative voltage. As a result under light illumination, the photocurrent was enhanced, which may be due to the separation of photoexcited carriers by Ag NPs/In<sub>2</sub>O<sub>3</sub> NW junction as well as it leads to increase in the width of depletion region, and therefore, no tunnelling has occurred [19].

The authors were also plotted specific detectivity ( $D^*$ ) and noise equivalent power (NEP) from  $\ln(I)$ - $V$  characteristics using the below equations to determine the detectors properties, displayed in Fig. 8(a) and 8(b), respectively.

$$D^* = \frac{R}{\sqrt{2e J_{\text{dark}}}} \quad (5)$$





**Fig. 8** (a) Detectivity vs. voltage graph for the GLAD:  $\text{In}_2\text{O}_3$  NWs and  $\text{In}_2\text{O}_3$  NWs/Ag NPs device. (b) Detectivity vs. voltage graph for the GLAD:  $\text{In}_2\text{O}_3$  NWs (inset of Fig. 7(a)). (c) NEP vs. voltage graph for both the same devices

$$\text{NEP} = \frac{\sqrt{A}}{D^*} \tag{6}$$

where  $R$  is the responsivity of the devices,  $e$  is the charge of an electron and  $J_{\text{dark}}$  is current density under dark illumination,  $A$  is the particular device area [20]. The maximum detectivity of  $\sim 78 \times 10^7$  jones (Fig. 8(a)) for the device -2 was observed from the applied voltage of -8 to -1 V, which possesses  $\sim 6.5$ -fold boost-up as compared with that of device-1 (Fig. 8(a), inset). The augmentation in detectivity was achieved due to the low dark current and SPR effect in the device-2. Wang et al. reported the enhancement in detectivity was due to low dark current levels in the Ag NPs surrounded ZnO thin film photodetector [53]. The low NEP was observed  $\sim 19 \times 10^{-12}$  W (Fig. 8(b)) for the device-2 at the same applied voltage, and it retains the  $\sim 1.8$ -fold low NEP as compared with that of device-1. In our case, the calculated NEP was lower than the previously reported values [54, 55]. The comparison of different outcomes of this work with other reported works based on their performance was depicted in Table 1. Additionally, the traps reduces the occurrence of generation-recombination, which may decrease the NEP and enhances

$D^*$ . Moreover, the low NEP results in the high detectivity and increased in photocurrent for the Ag NPs based device, which provides a flexible opportunity to use  $\text{In}_2\text{O}_3$  NWs/Ag NPs device as a low noise photodetector for optoelectronics applications.

### Conclusion

In this paper, the authors successfully synthesized Ag NPs over  $\text{In}_2\text{O}_3$  NWs using a double-step GLAD technique. From HRTEM image, the average diameter of  $\sim 25$  nm and bottom diameter of  $\sim 16$  nm of single NW have been extracted. The ‘ $d$ ’ spacing of 2 Å, 1.1 Å and 0.8 Å was calculated from SAED pattern, and it displays the amorphous nature of the  $\text{In}_2\text{O}_3$  NWs. HRXRD profile exhibits the formation of monoclinic  $\text{Ag}_3\text{O}_4$  compound over the  $\text{In}_2\text{O}_3$  NWs, which indicated the Ag–O compound with (031) diffraction plane. The PL emission of  $\text{In}_2\text{O}_3$  NWs/Ag NPs displayed the red shifting of spectrum at 466 nm ( $\sim 2.66$  eV) due to the scattering of SPR effect by  $\text{Ag}_3\text{O}_4$  compound over the  $\text{In}_2\text{O}_3$  NWs. The mitigation of  $\sim 1.22$ -fold root mean square surface

**Table 1** Comparison of different outcomes of this work with other reported works based on their performance

| Device structure                                | Device type                        | Detectivity (jones)                       | NEP (watt)            | References  |
|---|------------------------------------|---|-----------------------|-------------|
| n-Si/ $\text{In}_2\text{O}_3$ NW/Ag NPs         | Metal-oxide-semiconductor detector | $78 \times 10^7$<br>$= (7.8 \times 10^8)$ | $19 \times 10^{-12}$  | this device |
| $\text{Al}_2\text{O}_3/\text{AlGaIn}/\text{Si}$ | MIS detector                       | $4.52 \times 10^8$                        | $48 \times 10^{-12}$  | [54]        |
| GaN /Ni   | Schottky barrier                   | $1.57 \times 10^7$                        | $9.95 \times 10^{-8}$ | [55]        |
| AlGaIn/GaN                                      | MSM detector                       | $2.85 \times 10^7$                        | $1.62 \times 10^{-8}$ | [56]        |
| Al/SiO <sub>2</sub> /GaN                        | MIS detector                       | $2.03 \times 10^8$                        | $2.19 \times 10^{-9}$ | [57]        |

roughness was observed for the  $\text{In}_2\text{O}_3$  NWs/Ag NPs, which affects the packing density in the  $\text{In}_2\text{O}_3$  nanostructures. The n-Si/ $\text{In}_2\text{O}_3$  NWs device and n-Si/ $\text{In}_2\text{O}_3$  NWs/Ag NPs devices were created using In top contact through metallization process. At  $-4.5$  V, the n-Si/ $\text{In}_2\text{O}_3$  NWs/Ag NPs/In device shows higher light current due to the coupling of SPR effect of Ag NPs as compared with the n-Si/ $\text{In}_2\text{O}_3$  NWs/In device under a 100-W tungsten filament source. The enhancement in light current attributed de-trapping process due to the oxygen-related trap states which reduces the depletion region in the structures and allows the tunnelling of electrons. The ideality factor of  $\sim 20$  and  $\sim 11$  was calculated for the n-Si/ $\text{In}_2\text{O}_3$  NWs/In device and n-Si/ $\text{In}_2\text{O}_3$  NWs/Ag NPs/In devices. The maximum  $\sim 2.6$ -fold enhancement in photosensitivity was observed at an applied bias of  $-2.4$  V due to the SPR effect in the n-Si/ $\text{In}_2\text{O}_3$  NWs/Ag NPs/In device. The maximum specific detectivity of  $\sim 78 \times 10^7$  jones and low NEP of  $\sim 19 \times 10^{-12}$  W were observed for the devices from the applied voltage of  $-8$  to  $-1$  V, which possess  $\sim 6.5$ -fold enhancement in detectivity and  $\sim 1.8$ -fold low NEP in the Ag NPs decorated device. Therefore, the whole technical experimental process signifies that it can be used as an efficient Ag NP-based plasmonic photodetector for optoelectronics applications with better photosensitivity in the existing technology.

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