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Indium tin oxide and indium phosphide heterojunction nanowire array solar cells

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Heterojunction solar cells were formed with a position-controlled InP nanowire array sputtered with indium tin oxide (ITO). The ITO not only acted as a transparent electrode but also as forming a photovoltaic junction. The devices exhibited an open-circuit voltage of 0.436 V, short-circuit current of 24.8 mA/cm², and fill factor of 0.682, giving a power conversion efficiency of 7.37% under AM1.5 G illumination. The internal quantum efficiency of the device was higher than that of the world-record InP cell in the short wavelength range. © 2013 AIP Publishing LLC.

Photovoltaic devices based on semiconductor nanowire (NW) arrays have drawn interest as a possible way of reducing costs and materials while maintaining the power conversion efficiency of solar cells.1–8 A size- and position-optimized array of NWs can absorb an amount of light almost equal to that of a bulk structure with an antireflection coating because of its near-field-optics enhanced absorption and antireflection effect.9,10 However, photovoltaics based on NW arrays need to have a transparent electrode put on their NWs before the collecting electrode is formed, because each NW independently functions as a solar cell. In many cases, indium tin oxide (ITO) has been used as the transparent electrode since it has high conductivity and transmittance.1–7,11 We have focused on the ITO because it can act not only as a transparent conductive oxide but also as forming a photovoltaic junction with groups IV and III-V semiconductors.12–14 In this study, we experimentally demonstrated an ITO/InP heterojunction NW array photovoltaic device. The p-doped InP NW array was epitaxially grown using selective area growth (SAG), which has the capability of position control. Photovoltaic junctions were formed by ITO sputtering on the p-InP NW array. The resulting heterojunction NW-array cell had low reflectance and excellent quantum efficiency in the short wavelength region.

The NW growth and the fabrication process for the NW array solar cells were as previously reported.15 We grew p-type InP NWs by selective-area MOVPE using a 20-nm-thick SiO₂ mask deposited onto p-type InP (111)A substrates (N_A ~ 5 x 10¹⁸ cm⁻²). The SiO₂ pattern was designed to be a triangular periodic array of openings with a diameter, d₀, of 200 nm and pitch, a, of 400 nm in 1.2 x 1.2 mm² regions. The NWs were grown using metal-organic vapor phase epitaxy. After the growth, the NW array was embedded by spin-coating benzocyclobutene (BCB, Dow Chemical). The BCB layer was etched by reactive ion etching (RIE) with CF₄/O₂, exposing the tips of the NWs. A transparent ITO (SnO₂/In₂O₃; 10:90 wt. %) layer was then rf sputtered at a rate of 1.1 Å/s for 45 min onto the NW array at room temperature and annealed in N₂ at 400 °C for 15 min. The sputtered ITO film typically has a sheet resistance of 10 Ω/□ and an optical transmittance higher than 85% in the wavelength range between 400 nm and 900 nm. U-shaped Ag and alloying Au-Zn electrodes were, respectively, formed on the ITO layer and back of the substrate. The active device area was 0.62 mm² and included approximately 4.5 x 10⁶ NWs.

Figure 1(a) shows 20°-tilted scanning electron microscopy (SEM) image of p-InP NWs grown on patterned p-InP(111)A substrates. The NWs were well-defined and highly uniform in length, diameter, shape, and position. The sidewalls of the hexagonal NW were {−211} facet surfaces vertical to the substrate, as shown in Fig. 1(b). The average height and diameter of NWs were 1.0 µm and 170 nm, respectively. The geometric fill factor (FF), i.e., the ratio between the footprint area of the NW and the area of the unit cell, was 18%. The SEM image in Fig. 1(c) shows a cross-section of the NWs after completion of the device processes. The sputtered ITO layer had a dome-shaped morphology, which functioned as a light coupling within the NW array.

Figure 2(a) shows the current density versus voltage (J–V) characteristics of the fabricated ITO/p-InP heterojunction solar cell. They were measured under AM1.5G (1 sun) illumination at 25.0 °C. The illumination intensity was calibrated using a standard silicon photodiode (Bunkoukeiki BS-500). The inset shows the J–V characteristics of the cell in the dark. The device had good rectifying properties and a rectification ratio greater than 10⁶ at ±1.0 V in the dark, which means that a photovoltaic junction formed in it. These rectification properties can be attributed to SAG, which guarantees high crystal quality by preventing any metal from being incorporated during epitaxial growth.18 There are several models that could be used to explain the behavior of the ITO/p-InP junction, including heterostructure, buried n⁺/p homojunction, metal-insulator-semiconductor, and...
metal-semi-insulating-semiconductor, but the buried $n^+/p$ homojunction model is compelling because the deposition of an ITO layer induces an $n^+$-defect layer near the surface.

The device exhibited an open circuit voltage ($V_{OC}$) of 0.436 V, a short circuit current density ($J_{SC}$) of 24.8 mA/cm$^2$, and a $FF$ of 0.682, for an overall power conversion efficiency ($\eta$) of 7.37%. The current density and efficiency were determined from the active area, 82% of which was the space between the NWs. This high $J_{SC}$ for a NW-array-based device resulted from the combination of near-field-optics enhanced absorption and anti-reflection effects. The ideality factor and series resistance determined from the illumination $J-V$ characteristics were 1.6 and 0.21 $\Omega$-cm$^2$.

To analyze the absorption properties, we investigated the quantum efficiency of the heterojunction cell, as shown in Fig. 2(b). In the wavelength range between 370 nm and 730 nm, the internal quantum efficiency (IQE) of the device exceeded 80%, despite the device covering only 18% of the substrate surface. The IQE at shorter wavelengths was much higher than the previous reported InP NW solar cell, and the device had peak IQE of 0.943 at 490 nm without employing a window layer for reflecting minority carriers moving toward the front ITO layer. Moreover, the IQE of our heterojunction cell was significantly greater than that of the current world record holder for InP planar cells in the range of 300 nm and 570 nm. There were two reasons for this. First, the wide-band-gap ITO layer functioned as a window layer, which reflects minority carriers, for the InP layer; thus, IQE increased because of a reduction in the amount of surface recombination. Second, the tips of the InP NWs, whose average length was 300 nm, were covered by hemispherical caps of ITO. A photovoltaic junction formed not only at the top but also on the sides of the NWs, and this resulted in an improvement in the separation and collection efficiency of the photogenerated carriers. In addition, more than 98% of the incident light with wavelengths less than 500 nm was absorbed within a depth of 300 nm in the InP, and hence, the IQE at shorter wavelengths increased. The effective reflectance of the device after taking account of the solar spectrum distribution was 6.2%, and it should be emphasized that the device did not have specially designed antireflection coatings. It is thought the lower index of refraction in the NW array structure made for a stepped-index antireflection coating.

The absorption on the low-energy side started to rise from about 870 nm, which is substantially blue-shifted relative to the bulk bandgap of InP (925 nm). To examine the course of the blue-shift, we measured the crystal structure of the InP NW by using high-resolution transmission electron microscopy (HR-TEM) at an acceleration voltage of 300 kV along the (-110) incidence, as shown in Fig. 3(a). Figures 3(b) and 3(c) show the corresponding selective area electron diffraction (SAED) patterns. In the HR-TEM image, the crystal structure in the NW region is clearly different from the substrate region. The SAED pattern at the NW indicates the grown p-InP NW is a pure wurtzite structure without a zincblende structure (Fig. 3(b)); in contrast, zincblende spots can be seen in the SAED pattern at the interface between the NWs and the substrate (Fig. 3(c)). The wurtzite structure of the NWs was confirmed by making a reciprocal space mapping results of X-ray diffraction, as shown in Fig. 3(d). The (10–1.5) and (331) peaks correspond to wurtzite and zincblende structures. This is consistent with our previous results whose samples were grown under similar growth-temperature and V/III ratio conditions. It is known that bandgap energy of wurtzite InP is about 80 meV higher than that of zincblende. This suggests that the absorption edge on the low-energy side of the heterojunction device is blue-shifted. To clarify the bandgap shift in the p-InP NWs, the wires were studied with micro-photoluminescence (PL) at room temperature (Fig. 3(e)). The PL spectrum of the InP substrate exhibits a peak at 920 nm (=1.35 eV), while the
spectrum of the $p$-InP NWs had a peak at 885 nm (=1.40 eV). The PL peak of the NWs was blue-shifted, but it was less than the theoretical predicted 80 meV. This could be explained by the free-to-bound excitons stemming from zinc-doping. In addition, the HR-TEM image shows that the $p$-InP NWs had no twin and misfit dislocations, suggesting that the grown NWs were high-quality wurtzite crystals. This resulting low rate of bulk recombination was another source of IQE improvement.

In conclusion, we demonstrated a heterojunction solar cell that combines a $p$-doped InP NW array and a transparent ITO layer. The position-controlled and catalyst-free NWs were grown using SAG. The fabricated device showed a power conversion efficiency of 7.37% under AM1.5G with a $J_{SC}$ of 24.8 mA/cm$^2$. Furthermore, the ITO/InP heterojunction cell in the wavelength range below 570 nm. This showed higher IQE than that of the current world-record-hold-

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