A high-performance TiO$_2$ nanowire UV detector assembled by electrospinning

Yuting Wang, Jing Cheng, Muhammad Shahid, Mengfei Zhang and Wei Pan*

Here, a transparent, flexible and nanoscale TiO$_2$ ultraviolet (UV) photodetector has been fabricated by electrospinning. Well aligned TiO$_2$ nanowires were collected on a flexible mica substrate. Then, the nano-device was simply assembled by depositing interdigitated platinum electrodes on the surface. Upon UV illumination, the photosensitivity of this device is up to over three orders of magnitude with a relatively fast and stable response speed under 254 nm and 365 nm UV light. This nanosensor retains a high photo-dark current ratio, fast response time and stable durability during bending tests, indicating an excellent reversibility and stability of the flexible TiO$_2$ nanowires. The highly flexible photosensor demonstrates a good potential candidate for wearable optoelectronic applications.

1. Introduction

Flexible or stretchable electronic devices with high mechanical stability are considered key components for next generation wearable smart electronics.\textsuperscript{1–3} Although inorganic-based semiconducting nanomaterials present excellent UV sensitivity and fast response/recovery speed,\textsuperscript{4} the poor flexibility of ceramic materials hinders their wide application.\textsuperscript{5} Flexible photodetectors based on bendable plastic substrates have attracted great attention because of their potential applications in optoelectronics, sensors, and transducers, etc.\textsuperscript{6,7} Among various ceramic oxides, titanium dioxide, as an n-type semiconductor with a wide bandgap (anatase 3.2 eV and rutile 3.0 eV), is well known for its versatile characteristics in photocatalysis, solar cells, and diodes etc.\textsuperscript{8–11} In particular, TiO$_2$ has been widely used in photocatalysis because of its outstanding photocatalytic activity and electronic properties.\textsuperscript{12–17} Many nanotechniques have been developed to fabricate one-dimensional (1D) TiO$_2$, such as nanorods,\textsuperscript{18} nanowires,\textsuperscript{19} nanobelts,\textsuperscript{20} and nanotubes.\textsuperscript{21} Three crystal phases of TiO$_2$ are well known as rutile, anatase, and brookite.\textsuperscript{22} Although the bandgap energy for anatase phase (~3.2 eV) is larger than the rutile (~3 eV), it shows high photocatalytic activity and efficiency of the charge carrier photogeneration when combining anatase and rutile mixtures of TiO$_2$ nanocrystalline with a radio of 80% and 20%.\textsuperscript{23,24} As we known, few research has reported about the photoresponse performance of TiO$_2$-based photodetector.

Electrospinning is a facile approach to produce one-dimensional (1D) nanostructures with great scalability. The diameter and morphology of the electrospin nanofibers could be simply controlled by adjusting the density of precursor solutions, applied voltage, working distance and spinning rate.\textsuperscript{25} Moreover, this nanomaterial synthesize technology shows advantages of low cost, high efficiency and convenient assembly. Electrospun nanofibers demonstrate outstanding electric, optic, mechanical and magnetic flexibility properties, with considerable potential for novel applications in different fields.\textsuperscript{26–29} However, most of the reports for electrospun nanofibers concentrated on the 1D nanostructures. More detailed research on the fabrication of two-dimensional (2D) networks has not been reported so much. Actually, the collected electrospun nanowires could be automatically formed as a nanoweb on the plate substrate. Therefore, in this study, electrospinning was used for fabricating ultralong nanowires and flexible 2D nanowebs by simply annealing these 2D networks in air.

Here, we report a high-performance 2D nanoweb photosensor composed of ultralong 1D-structured TiO$_2$ nanowires. The nanoweb-based photodetector shows high photo-dark current ratio, excellent mechanical flexibility and durability. Possible mechanisms of the improved mechanical performance of this TiO$_2$ nanosensor have been discussed in detail.

2. Experimental section

TiO$_2$ nanowires were synthesized using an electrospinning approach. Tetrabutyl titanate (6 wt% of the ceramic precursor) was first dissolved in the mixture of ethanol–acetic acid (3.3 : 1, volume rate). After stirring for 30 min, a concentration of 6 wt% polyvinylpyrrolidone (PVP, MW = 1 300 000) was further mixed to control the viscosity. Afterward, the prepared solution was stirred at room temperature for 2 h to achieve complete dissolution and mixing. After dissolution of all the components, the obtained transparent precursor solution was placed in a hypodermic syringe at the positive voltage of 15 kV with a constant flow rate of 0.5 mL h$^{-1}$. The distance between the needle tip and
the aluminum foil collector was 15 cm. Finally, nanofiber webs
were removed from the collector and annealed in air at different
temperatures, 550 °C, 600 °C, 700 °C, and 800 °C for 1 h with the
heating rate of 5 °C min⁻¹, respectively.

The morphologies of TiO₂ nanowires and flexible nanoweb
were observed by scanning electron microscopy (SEM, JEOL
JSM-6460LV, Tokyo, Japan) and transmission electron micros-
copy (TEM, JEOL-2011, Tokyo, Japan). A diffractometer (XRD, D/
max-2500, Rigaku Co., Tokyo, Japan) was used to examine the
crystal structures of these samples. The current–voltage (I–V)
characteristics of TiO₂ networks were measured by a Keithley
wires. (e and f) TEM and HRTEM images of the TiO₂ nanowire annealed
structure and a smooth surface of the annealed TiO₂ nanowire.
A selected-area electron diffraction (SAED) pattern shown in the
inset of Fig. 1e reveals that the nanofibers are comprised of
polycrystalline and fine grains with no preferential orientation.
Fig. 1f shows a high-resolution transmission electron micros-
copy (HRTEM) image with distances of crystal lattice fringes of
3.5 Å that can be attributed to the spacing of (101) lattice plane
of the anatase TiO₂.

Ultra-long TiO₂ nanowires can be further assembled to a two-
dimensional nanopaper. This free-standing nanopaper could be
bended without any cracking. SEM measurements were used to
determine the strength of TiO₂ nanoweb. Fig. 2a and b show the
distinct boundary of the folded nanopaper. The TiO₂ nanofibers
are not broken or got any fracture when bended in half. More-
over, spring- and wave-like structures have been spontaneous
formed during electrospinning. After annealing, the spring-like
nanowires could maintain their curly and wavy geometries,
attributing to the superior mechanical flexibility of TiO₂ nano-
materials. Fig. 2d shows that the freestanding TiO₂ networks are
successfully folded in half and remain intact. The mechanistic
explanation is that the TiO₂ nanowires remain continuous and
undergo nano-scaled deformation to relax the applied stress
upon folding.

To investigate the crystal structure of the nanofibers, X-ray
diffraction (XRD) analyses of TiO₂ nanofiber-based powders
were observed after annealing at 500 °C, 550 °C, 600 °C, 700 °C,
and 800 °C for 1 h in Fig. 3. The as-spun nanofibers are amor-
phous in nature, and the polycrystalline nanofibers are formed
after calcination at 500 °C. We note that the annealing temper-
ate could distinctly affect the phase compositions of the TiO₂
nanowires. As the sample is calcined at 500 °C, a phase occurs
with a low and wide characteristic peak corresponding to the
typical anatase crystal, and this phase completely disappears at
800 °C, leaving pure rutile phases. The XRD patterns also show
a mixture of anatase and rutile phases in the annealing range of
500 to 700 °C. It proves that the different ratio of these two phases
in TiO₂ composite could be simply controlled by annealing at
different temperatures. Contrasting the strength of the anatase
(101) peak and the rutile (110) peak in Fig. 3, the concentration of
anatase phase in hybrid TiO₂ nanofibers can be estimated as
92%, 83%, 75%, 17%, and 6%, corresponding to the annealing
temperature change from 500 to 800 °C, respectively. It was re-
ported that anatase TiO₂ shows more photocatalytic activity than
its rutile phase due to its large surface area, thus, in this
study, the high-performance TiO₂ nanosensor is composed of
a majority of anatase phases combined with little amount of
rutile phases. With the high reactivity of anatase TiO₂ together
with the lower band gap of rutile TiO₂, the process of the

3. Result and discussion

Intuitively, the flexible photosensor consists of semi-conductive
nanowires and platinum electrodes, shown in Fig. 1a. By elec-
trospinning, continuous aligned TiO₂ nanowires are first
collected on a flexible and transparent mica substrate. Pt ele-
trodes were deposited using a mask by high quality vacuum
evaporation. Fig. 1b shows a photograph of the flexible sensor
attached to a nonplanar surface. The morphology and
curved surface. (c and d) SEM images of highly oriented TiO₂ nano-
trodes were deposited using a mask by high quality vacuum
heat. The TEM image in Fig. 1e also shows a fibrillar
structure and a smooth surface of the annealed TiO₂ nanowire.

**Fig. 1** Highly oriented TiO₂ nanowire photosensor. (a) Schematic
representation of the device assembled by aligned TiO₂ array and
coated Pt electrodes. (b) Photograph of the photosensor attached to
curved surface. (c and d) SEM images of highly oriented TiO₂ nano-
wires. (e and f) TEM and HRTEM images of the TiO₂ nanowire annealed
to 550 °C. The inset shows the SAED pattern.
photoinduced electrons and holes transfer between conduction and valence bands can be highly facilitated in the composite sample.

The mechanism of phase-dependent photoresponse activity in phase junction TiO$_2$ nanowires is drawn in Fig. 4a. The band gaps of anatase and rutile phases of TiO$_2$ are 3.2 and 3.0 eV, respectively. This means that when the two phases are in nano-contact there will be a phase junction between the band edges. The phase junctions can promote charges transfer from anatase to rutile phases, similar to the mechanism of P25. When the photosensor is illuminated under UV light with photon energy higher than the band gap of anatase TiO$_2$, electrons in the...
valence band can be excited to the conductive band with the generation of the same amount of holes. Generally these electron–hole pairs are easily recombined that reducing the photoresponse of semiconductors. However, the existing two phases make it possible for the photogenerated charges to migrate from anatase to rutile phases and to efficiently separate the photoinduced electron–hole pairs.

The developed single TiO$_2$ nanowire shows a low dark current of approximately pA level even at an applied voltage of 10 V, as shown in Fig. 4b. A significant increase in the current is observed under 254 nm UV light illumination (500 µW cm$^{-2}$) with a high photocurrent-to-dark current ratio of at approximately three orders of magnitude. The photocurrent of the sample annealed at 550 ºC could reach up to 47 nA and 38 nA under 254 nm and 365 nm UV light at an applied voltage of 10 V, corresponding to a photo-dark current ratio of 760 and 600, respectively. For comparison, the photo-dark current ratio of the anatase (annealed at 500 ºC) and rutile (annealed at 800 ºC) TiO$_2$ nanowires are 160 and 1.1, respectively. The performance of the composite TiO$_2$ nanowires is comparable with other reported ZnO-based and TiO$_2$-based photodetectors. Compared with reported flexible ZnO nanowires, our TiO$_2$ network photosensor shows much better flexibility. For the ZnO photosensors, they can withstand a bending radius of 10 mm or 15 mm. But the photosensor would keep stable only after releasing the strain. In our case, the photoresponse tests were measured while the sensor was under bending.

The flexibility of TiO$_2$ photodetector annealed at 550 ºC was studied by bending to different curvature radii and repeated bending tests. It is noteworthy that the $I$–$V$ characteristics keep linear and remain stable under UV light illumination, as shown in Fig. 5a. When bended to a radius from 100 to 5 mm, the responsivity shows a little decrease upon illumination at 254 nm. Even at the smallest curvature radius of 5 mm, the responsivity shows a high retention of 85% and 65% at 10 V and 1 V, respectively (Fig. 5c). The retention of responsivities at curvature radius of 100 mm and 10 mm was tested to be 96% and 78% during cycling under bending (Fig. 4b), respectively. This excellent flexible performance could be corresponding to the SEM results and visible bending tests in Fig. 2. As the TiO$_2$ nanowire could retain a continues nanostructure without any cracking, the electrical properties of them could also keep stable under bending. The response time of photodetector is another major issue for its practical application. Fig. 5d shows the response time of photodetector upon 254 nm illumination at 1 V and 10 V. When the applied voltage is 1 V, this response time decreases slightly under bending. With a higher voltage of 10 V excites the photosensor, this photoresponse time could be remained stable even after bending to a curvature radium of 5 mm. These results demonstrate the great mechanical flexibility and durability of this TiO$_2$ photodetector.

Fig. 5 (a) $I$–$V$ curves of the flexible TiO$_2$ nanopaper photodetector annealed at 550 ºC under dark or UV light illumination. (b) Photo-switching characteristics of the photosensor under dark and UV light. Plot of sensitivity (c) or response time (d) versus bending curvature radium of the photosensor under 254 nm at 1 or 10 V bias.
continuous and undergo nano-scaled deformation to relax the applied stress upon bending. Further observations under XRD demonstrate that the crystal structure of this sensor is composed of approximately 83% anatase and 17% rutile phases. The energy band gap of TiO$_2$ nanoweb is calculated to be 3.15 eV, which is a little lower than that of anatase but higher than rutile phase, corresponding to the XRD results. Both of these results could be correlated to the efficient electron–hole transfer between the two phases in the composites, which lead to the superior UV responsibility of our photosensor.

Acknowledgements

This study was supported financially by the Natural Science Foundation of China (Grant No. 51323001).

References