Supporting Information

Recycled Silicon Solar Cells-Derived Nanostructured *p*-Black Silicon for High Performance NO₂ Gas Sensor Applications

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Characterization Techniques:

p-Black Silicon derived from recycled solar cells were spectroscopically and microscopically characterized by various techniques. X-Ray diffraction (XRD) patterns were acquired using a Power X-ray diffractometer (Ultima IV, Rigaku, Japan) with Cu K α radiation of wavelength 1.5418 Å at a scanning rate of 0.02°/sec in the 2 θ range of 20°–80°. Micro Raman studies on *p*-Black Silicon was performed using confocal Raman microscope (WiTec, 300 Alpha, Germany) with He-Cd laser ($\lambda = 532$ nm) as the excitation light source and 600 g/mm gratings. The structure and morphology of the *p*-Black Silicon was examined using Field Emission Scanning Electron Microscopy (ZEISS EVO 18, US), with in-built energy-dispersive X-ray spectrometer (Oxford Instruments, INCA, UK).

Evaluation of NO₂ gas sensing properties

The sensor device was fabricated using DC magnetron sputtering of gold thin films with ~ 300 nm thickness directly on the etched *p*-B-Silicon substrate using a laser machined stainless steel mask. The gas sensing properties of the materials were analyzed using a custom-built gas sensor test station consisting of a stainless-steel double-walled test chamber equipped with temperaturecontrolled hot stage, sensor holder, mass flow controllers (MFC, Alicat, USA), digital multimeter (Agilent 34401A, USA) connected with a data acquisition system interfaced with Labview software. During measurements, NO₂ gas was mixed with dry nitrogen to achieve the desired concentrations and the flow rate was maintained as 300 sccm using mass flow controllers (Alicat, U.S.A.). A temperature controller (Eurotherm, 2420, U.K.) was used to maintain the working temperature of the sensor mounted inside the sensing chamber. Constant gas pressure was maintained in the sensing chamber (710 Torr) throughout the testing, which was measured using Baratron 722B Absolute Capacitance Manometer (MKS Instruments, Singapore). In order to perform the interference studies, NH₃, H₂S, ethanol, and acetone were individually exposed to the sensor device by keeping the constant concentration using Owlstone gas generator unit (OVG-4, U.K.) the sensor response was measured. 35W HID kit Xenon source was used for light illumination.

| Conc. of NO ₂ (ppm) | <i>p</i> -B-Silicon | | |
|-----------------------------------|---|----------------------|----------------------|
| | Response [(R _g - R _a)/R _a] S (%) | Response time (s) | Recovery time (s) |
| 1 | 10.99 | 14.2 | 78.4 |
| 2 | 17.20 | 17 | 116.4 |
| 3 | 32.80 | 17.6 | 162 |
| 4 | 40.40 | 24.8 | 246.2 |
| 5 | 43.08 | 25.2 | 282.6 |

Table S1 NO₂ Sensing properties of *p*-B-Silicon under lower concentration.



Figure S1 (a) Dynamic NO2 sensing characteristics of p-B-Silicon and (b) NO₂ response as a function of concentration.

| Conc. of NO ₂ (ppm) | <i>p</i> -B-Silicon | | | |
|-----------------------------------|--|----------------------|----------------------|--|
| | Response $[(R_g - R_a)/R_a]$ S (%) | Response time (s) | Recovery time (s) | |
| 5 | 50.50 | 10 | 32.2 | |
| 10 | 106.63 | 11.4 | 92.2 | |
| 15 | 153.13 | 12.2 | 191 | |
| 20 | 211.19 | 14.2 | 279.8 | |
| 25 | 258.69 | 17.4 | 305.6 | |

Table S2 NO₂ Sensing properties of *p*-B-Silicon under higher concentration.



Figure S2 Dynamic NO₂ response-recovery graph of *p*-B-Silicon (a) recycled silicon wafer and (b) commercial silicon wafer.



Figure S3 Dynamic NO₂ sensing characteristics of p-B-Silicon (a) without Xenon light illumination and (b) under Xenon light illumination.



Figure S4 Relative humidity interference studies of p-B-Silicon (a) NO₂ response-recovery graph and (b) relative humidity (RH) versus sensitivity graph.