

Supporting Information

Highly sensitive room temperature liquid petroleum gas (LPG) sensor with fast response based on TiO₂-reduced graphene oxide (r-GO) composite

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S I. FT-IR spectrum of the composite

Figure S1 provides the FT-IR spectrum of the TiO₂-rGO composite. The peaks observed at the lower wavenumbers such as 545 cm⁻¹ and 670 cm⁻¹ correspond to vibrational modes of the TiO₂ molecule; ν (Ti-O) or ν (Ti-O-Ti) to be more exact (Verma et al., 2017). The intense and broad peak observed at ~3440 cm⁻¹ could be attributed to the O-H stretching vibrations of the residual functional groups of the reduced graphene oxide or of surface adsorbed moisture (Chen et al., 2019). The peak observed at ~1025 cm⁻¹ corresponds to the epoxide C-O-C or the alkoxy C-OH functional group of rGO. A relatively sharp peak observed at ~1650 cm⁻¹ could be assigned to the bending vibrational modes from the aromatic C=C of the graphene skeleton. An intense peak observed at ~2358 cm⁻¹ is generally associated with the asymmetric stretching of CO₂ adsorbed from the atmosphere. The transmittance peak observed at 2921 cm⁻¹ can be attributed to the combination band of TiO₂ (Verma et al., 2017). Its neighboring weak transmittance peak observed at ~2860 cm⁻¹ could be attributed to C-H stretching vibrations.

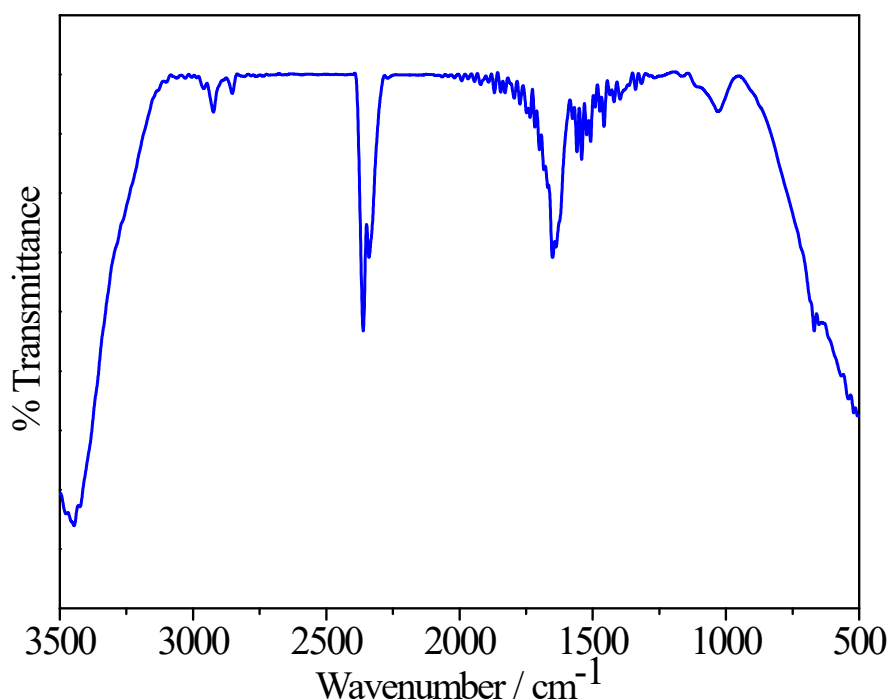


Figure S1 FT-IR transmittance spectrum of the TiO₂-rGO composite.

S II. Elemental composition from EDS

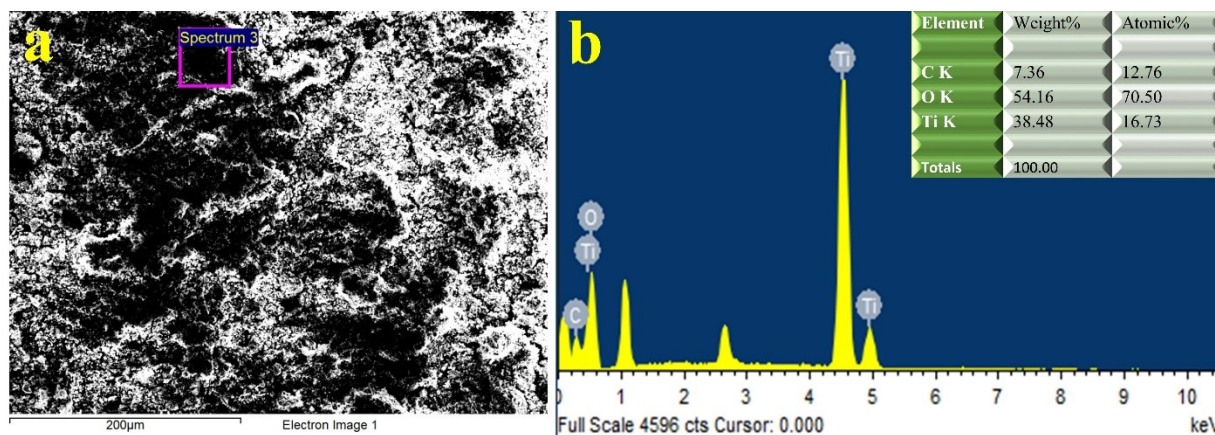


Figure S2 (a) SEM micrograph of the rGO-TiO₂ composite; (b) EDS spectra of the composite revealing the presence of C, O and Ti.

Figure S2 provides the EDS spectrum of the rGO-TiO₂ composite. S2a provides the associated SEM micrograph and S2b provides the EDS spectrum where we can clearly see the presence of C (coming from reduced graphene oxide), O (coming from TiO₂ and the oxygenated functional groups of rGO), and Ti (coming from TiO₂). The spectrum was obtained by scanning an area of ~2500 μm² of the sample. The atomic percentage observed indicated a Ti/O/C ratio of 16.7:70.5:12.3. However, because of the localized nature of electron microscopic measurements, the ratio most probably does not indicate the real atomic ratio within the composite. Additionally, there were peaks of elements such as Au, Sn etc that would accompany the spectrum due to the experimental requirements. These peaks were not indexed not accounted for while calculating the elemental ratio.

S III. Electron microscopy images of graphene oxide (GO)

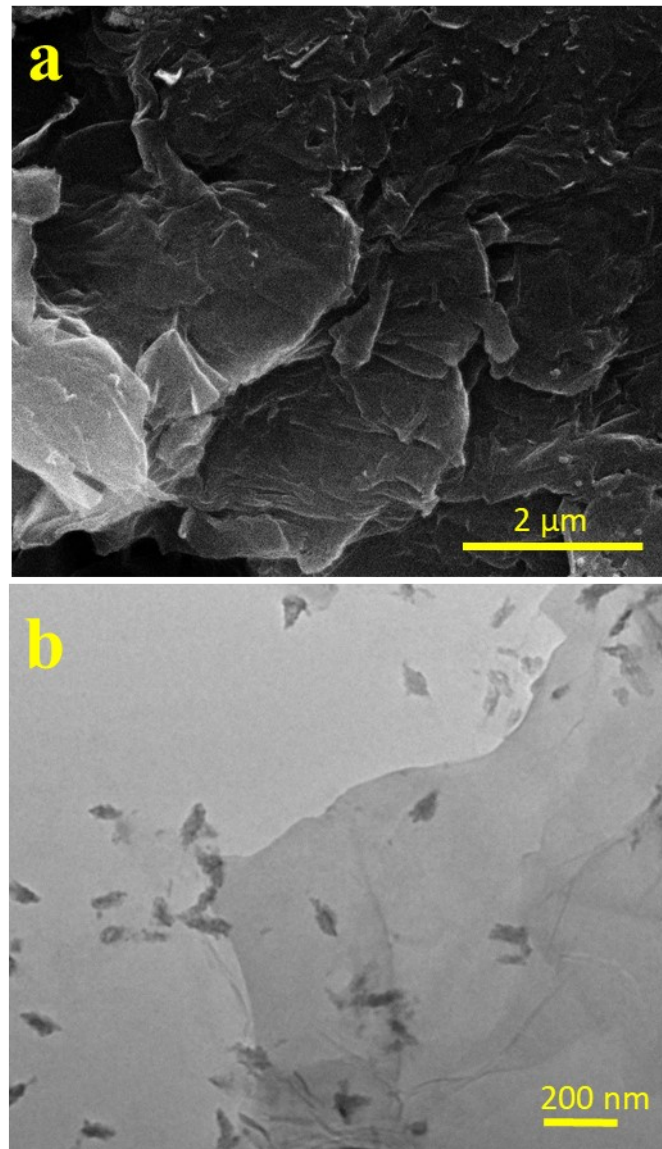


Figure S3 (a) FESEM micrograph and (b) TEM micrograph of the as synthesized GO particles. Both the images depict the sheet like morphology of GO. The smaller particles observed in (b) are the broken sheets due to sonication during sample preparation for TEM

S IV. TEM micrographs of reduced graphene oxide (rGO)

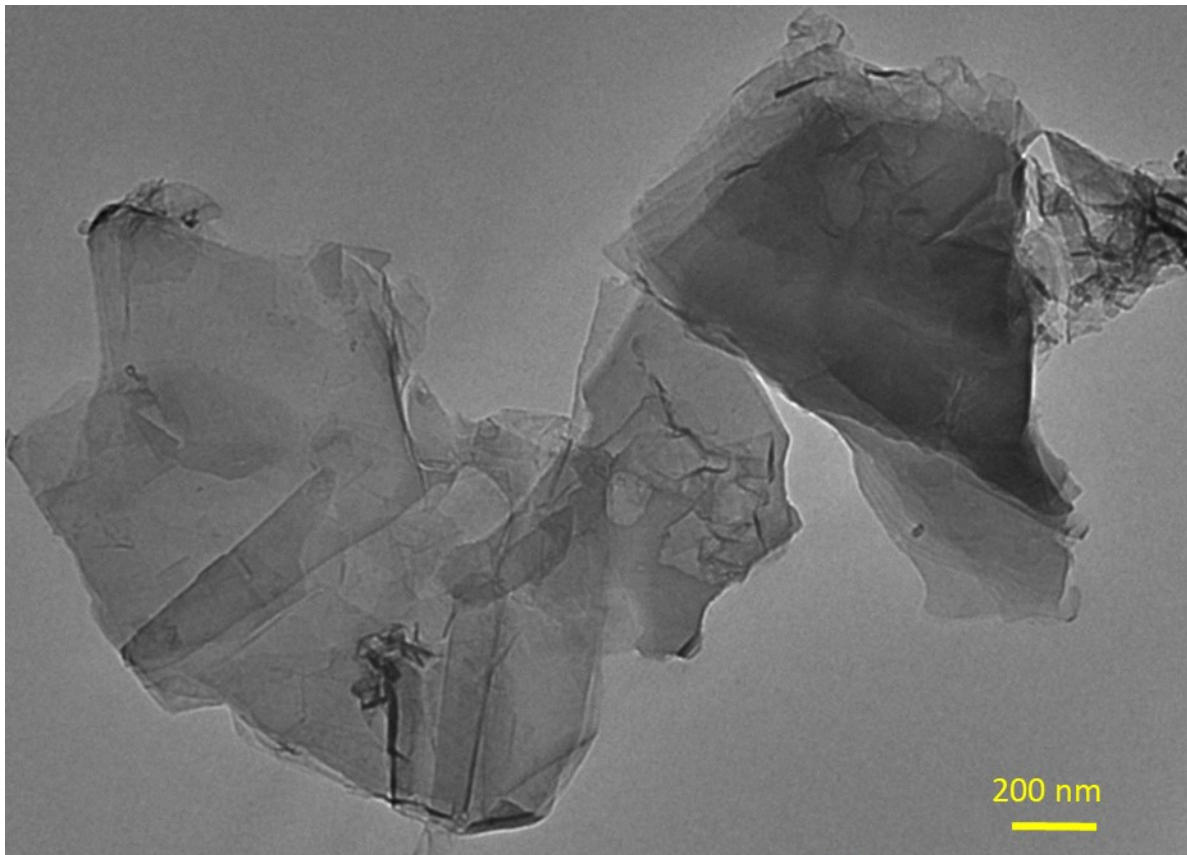


Figure S4 TEM micrograph of the reduced graphene oxide (rGO) displaying its sheet-like morphology. The wrinkles visible are characteristic of its sheet morphology.

S V. Elemental mapping of the TiO₂-rGO composite

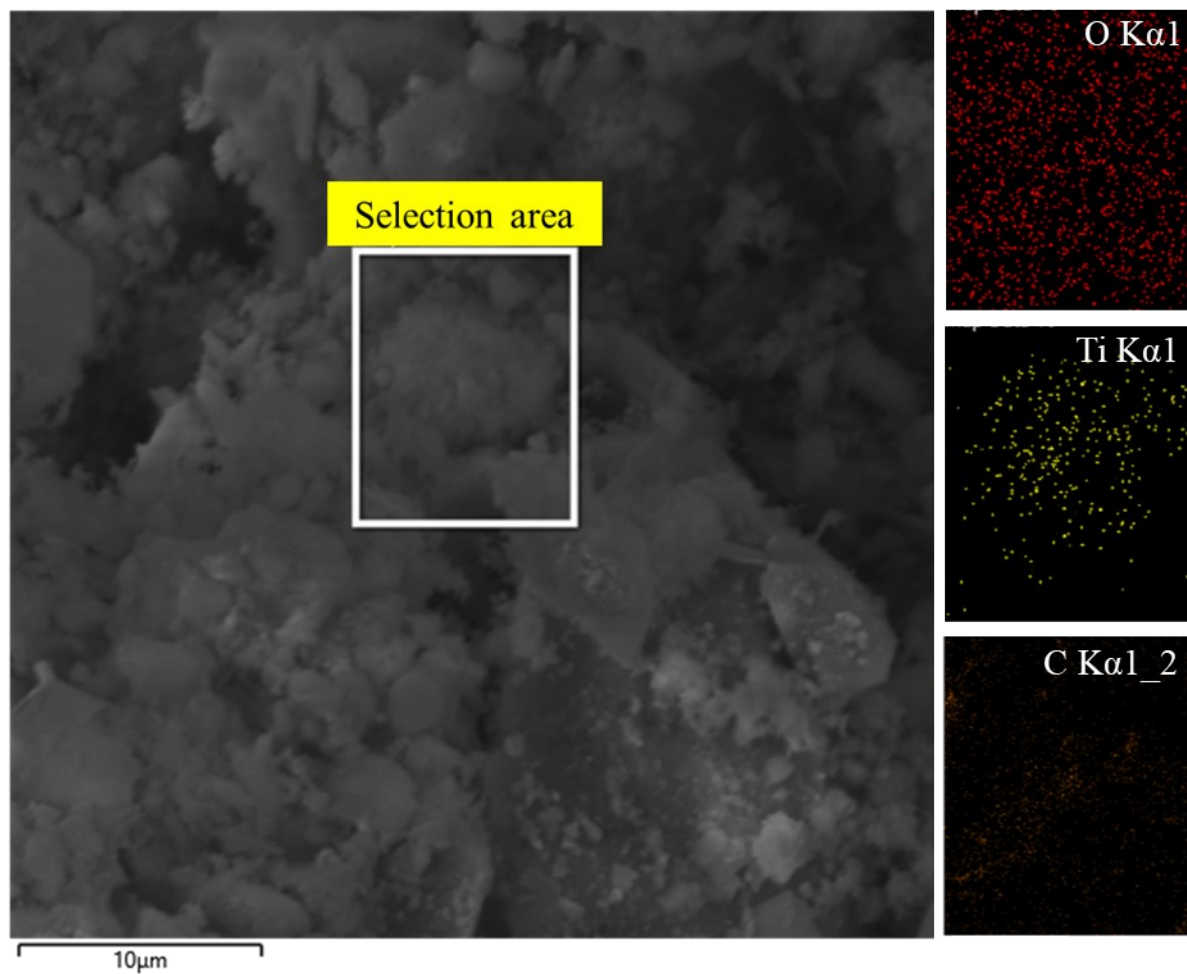


Figure S5 Elemental mapping of the TiO₂-rGO composite obtained from a portion of the micrograph. The mapping clearly shows how the rGO sheet (carbon) is all around the TiO₂.

S VI. Hydrodynamic particle size of the rGO-TiO₂ composite

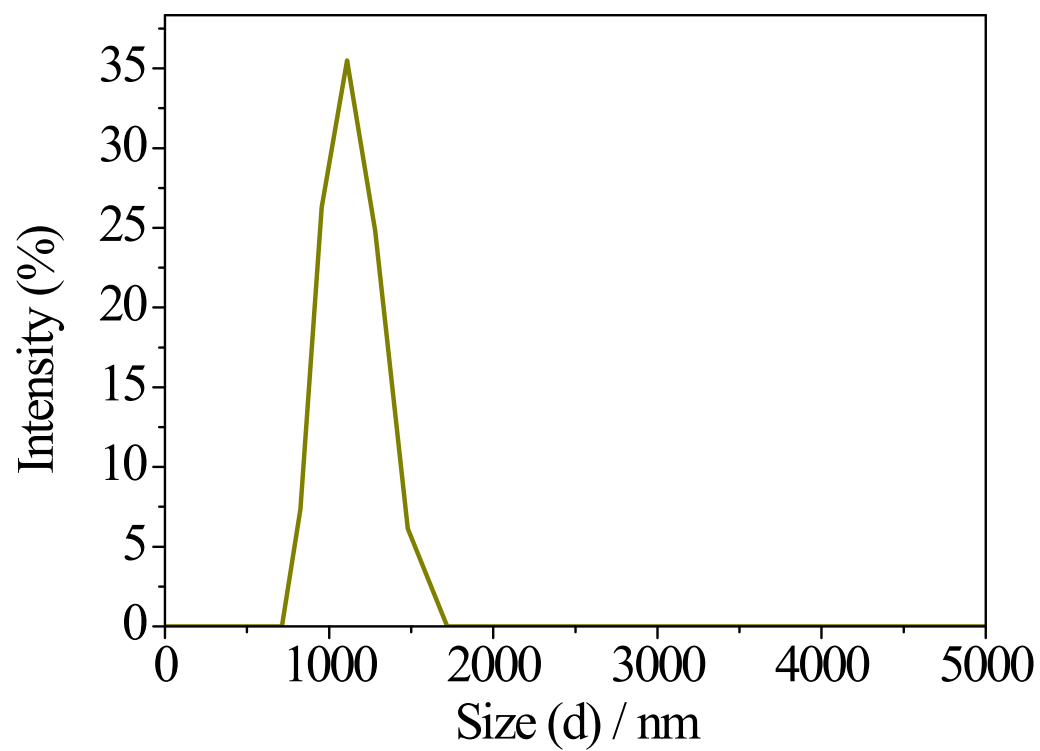


Figure S6 Intensity based particle size distribution (obtained from DLS technique) of the TiO₂-rGO composite material

S VII. A comparison between the sensing performance of r-GO, TiO₂ and the rGO-TiO₂ composite

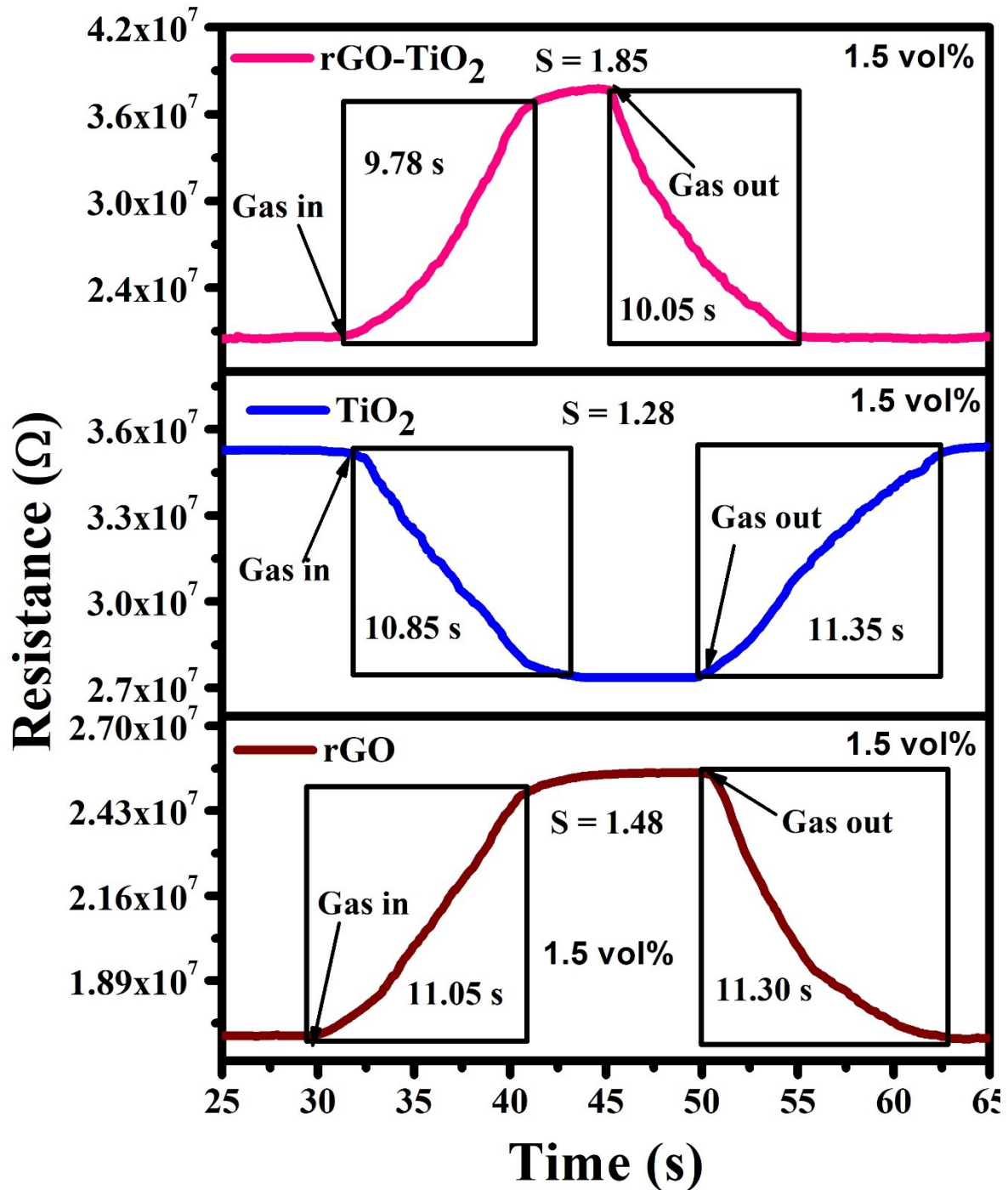


Figure S7: Transient sensor response of the TiO₂-rGO composite sensor, TiO₂ sensor and rGO sensor against exposure to 1.5 vol% LPG