

Supplementary information

Ultrathin Au nanowires supported on RGO/TiO₂ as an efficient photoelectrocatalyst

A Leelavathi,^a Giridhar Madras^b and N Ravishankar^{c*}

^a Centre for Nanoscience and Engineering, Indian Institute of Science, Bangalore-560012, India.

^b Department of Chemical Engineering, Indian Institute of Science, Bangalore-560012, India.

^c Materials Research Centre, Indian Institute of Science, Bangalore-560012, India.

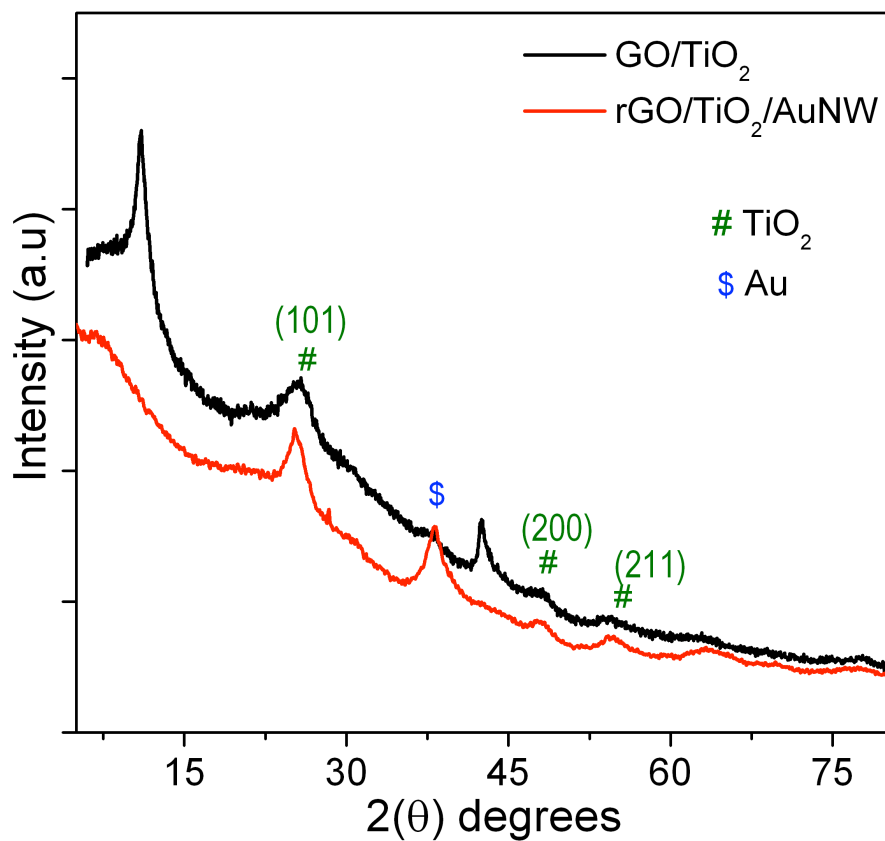


Fig. S1 pXRD patterns of GO/TiO₂ before and after growth of Au nanowires. GO, basal spacing decreased from 8 Å to ~3.7 Å after incubating in nanowire growth solution, which validates that the room temperature reduction of GO.¹ Indexed phase corresponds to anatase TiO₂.

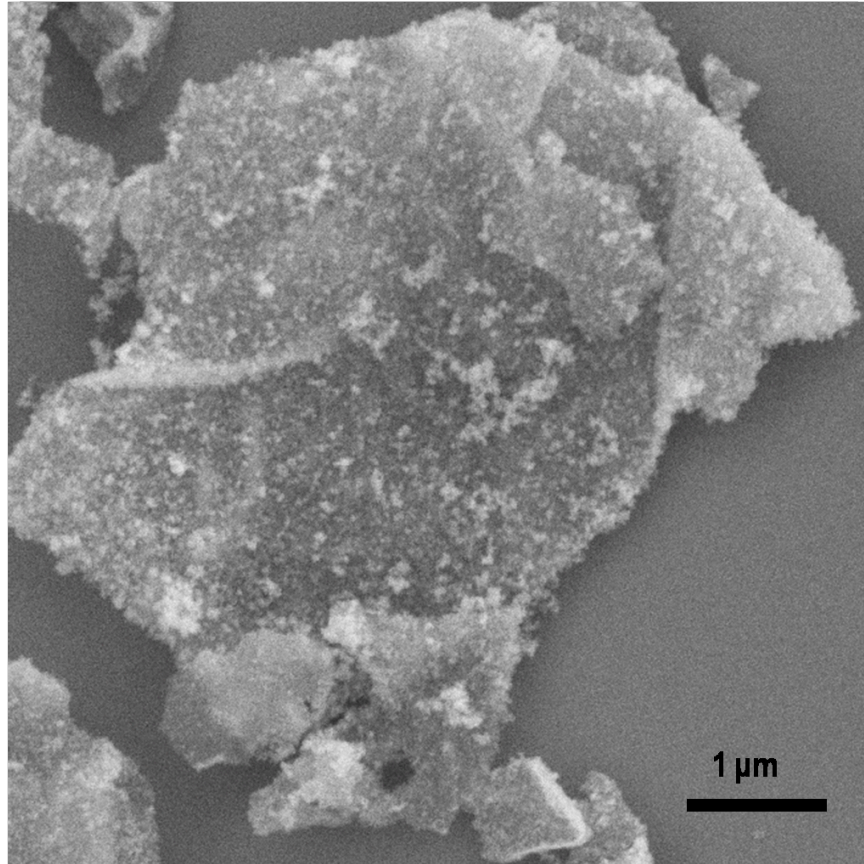


Fig. S2 SEM micrograph of GO/TiO₂, indicating that the heterogeneous nucleation of TiO₂ on sheets.

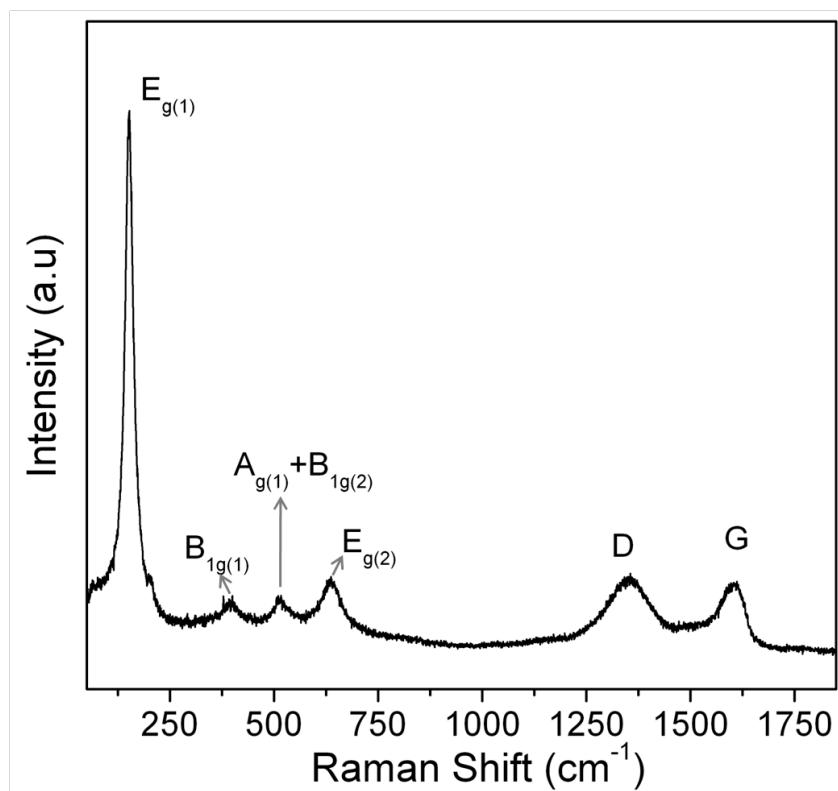


Fig. S3 Raman spectrum of GO/TiO₂ synthesized through microwave, using titanil nitrate precursor. Shows the typical features of GO with the presence of D and G band. Appearance of Raman modes around 150 cm⁻¹ (E_{g(1)}), 396 cm⁻¹ (B_{1g(1)}), 513 cm⁻¹ (A_{1g}+B_{1g(2)}) and 639 cm⁻¹ (E_{g(2)}) confirms the presence of anatase phase TiO₂ on GO sheets.²

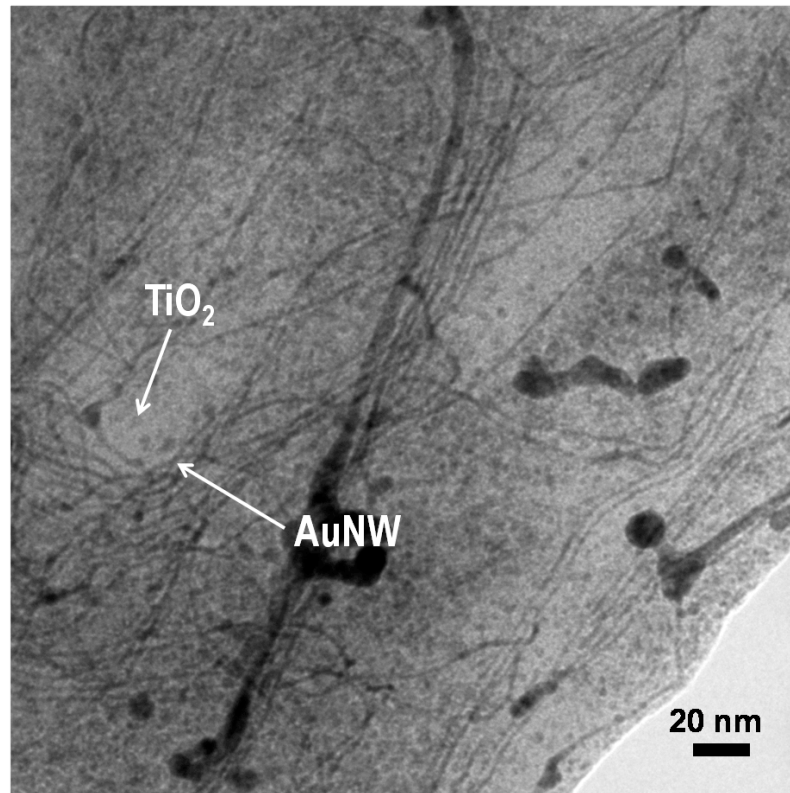


Fig. S4 TEM micrograph of rGO/TiO₂/AuNW, displays that the nanowires are dispersed on the support rGO/TiO₂.

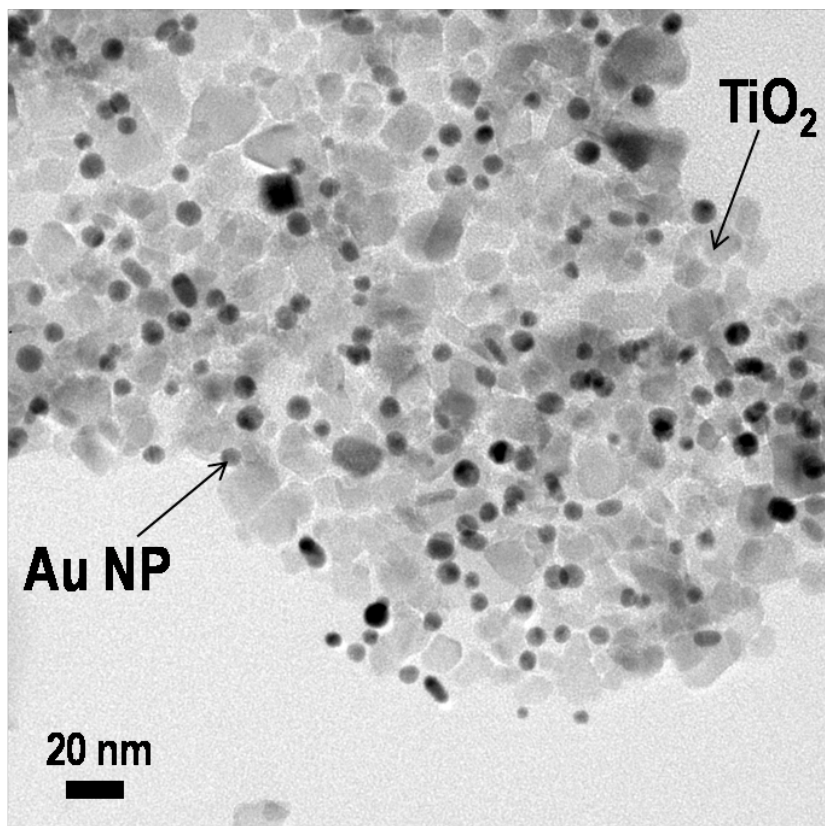


Fig. S5 Bright field micrograph displays that unmodified TiO₂ favors for the Au nanoparticles formation.

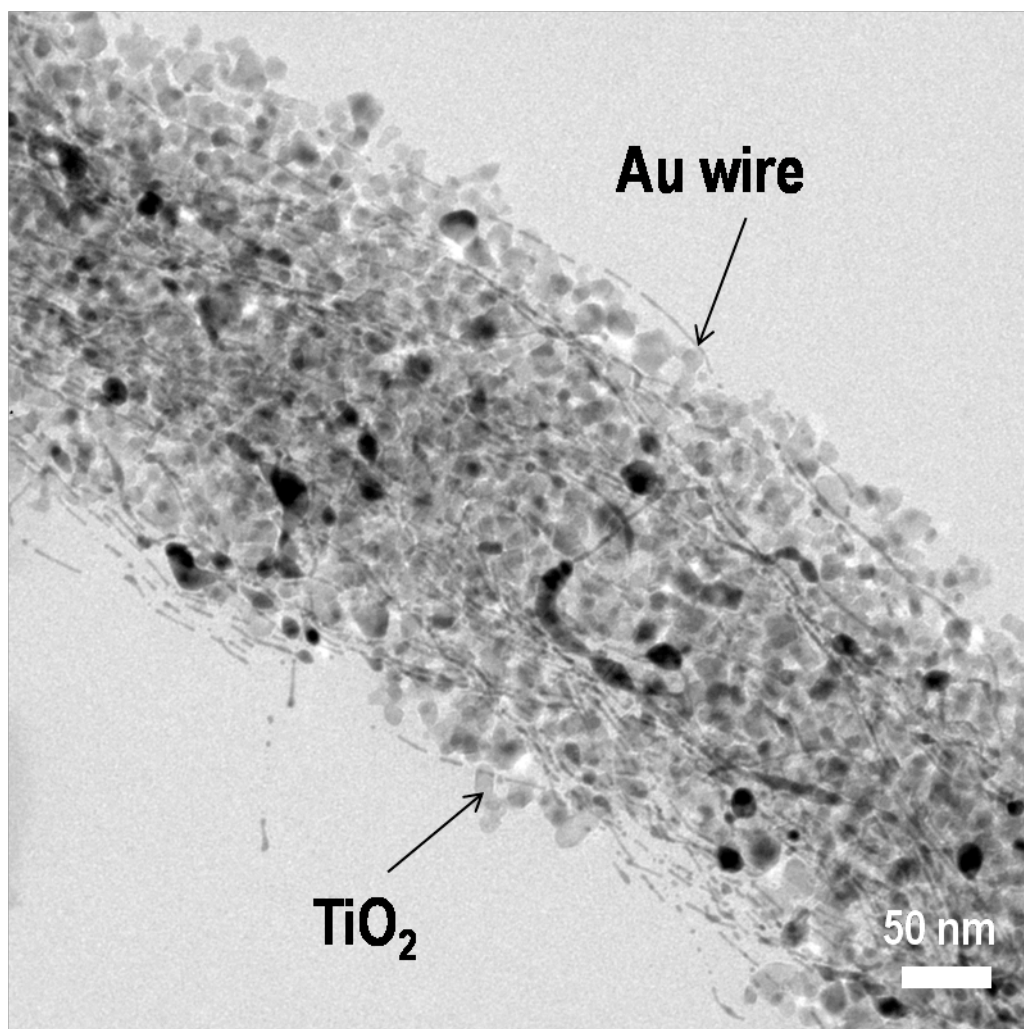


Fig. S6 Bright field TEM image reveals the growth of nanowires on amine modified TiO₂ nanoparticles.

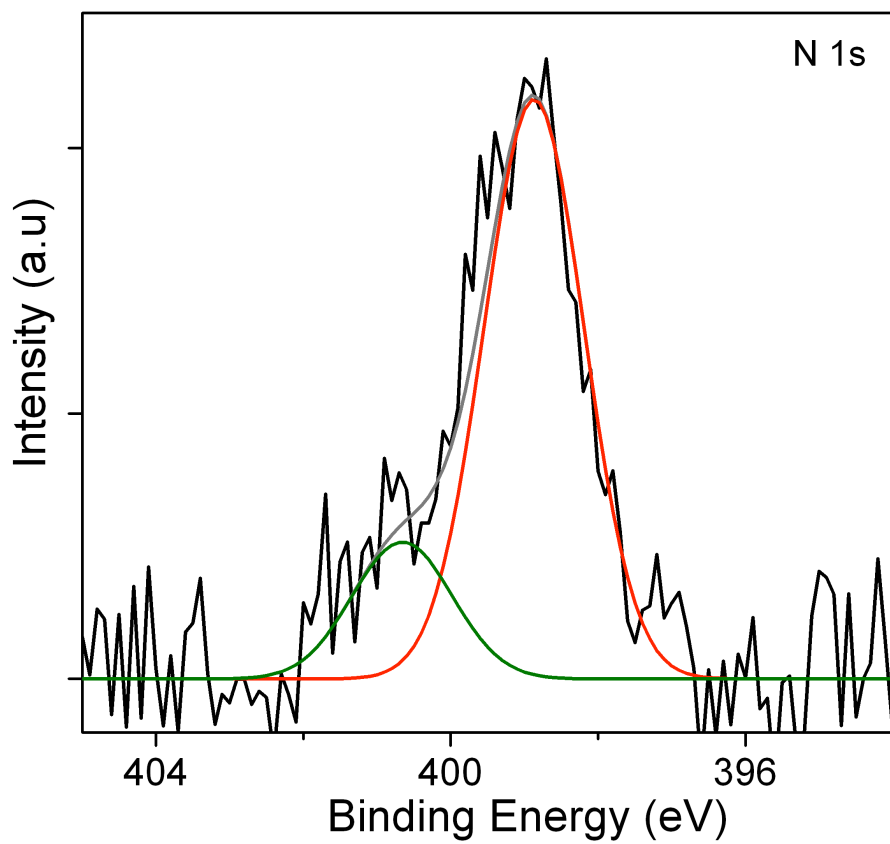


Fig. S7 Deconvoluted N 1s spectra of rGO/TiO₂/Au nanowires, displaying the existence of nitrogen species.

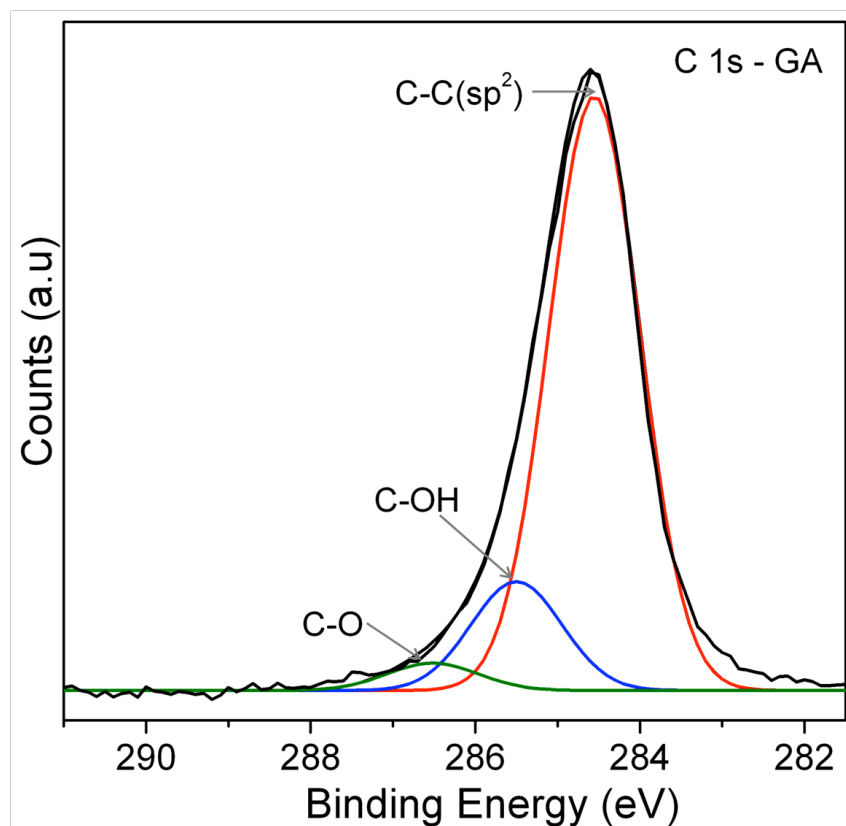


Fig. S8 XPS C 1s of rGO/AuNW(GA), showing the room temperature reduction of GO after incubating with Au nanowires growth solution for 6 h.

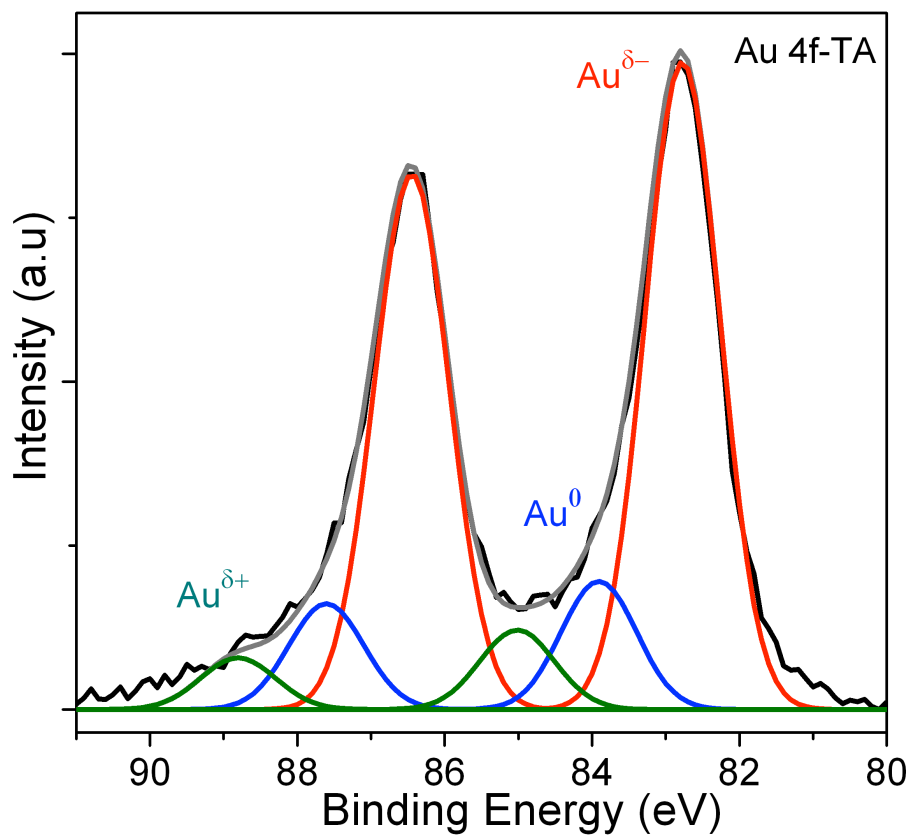


Fig. S9 Deconvoluted Au 4f of TiO₂/AuNW (TA) showing the presence of Au anionic species, predominantly electron-rich Au surface.

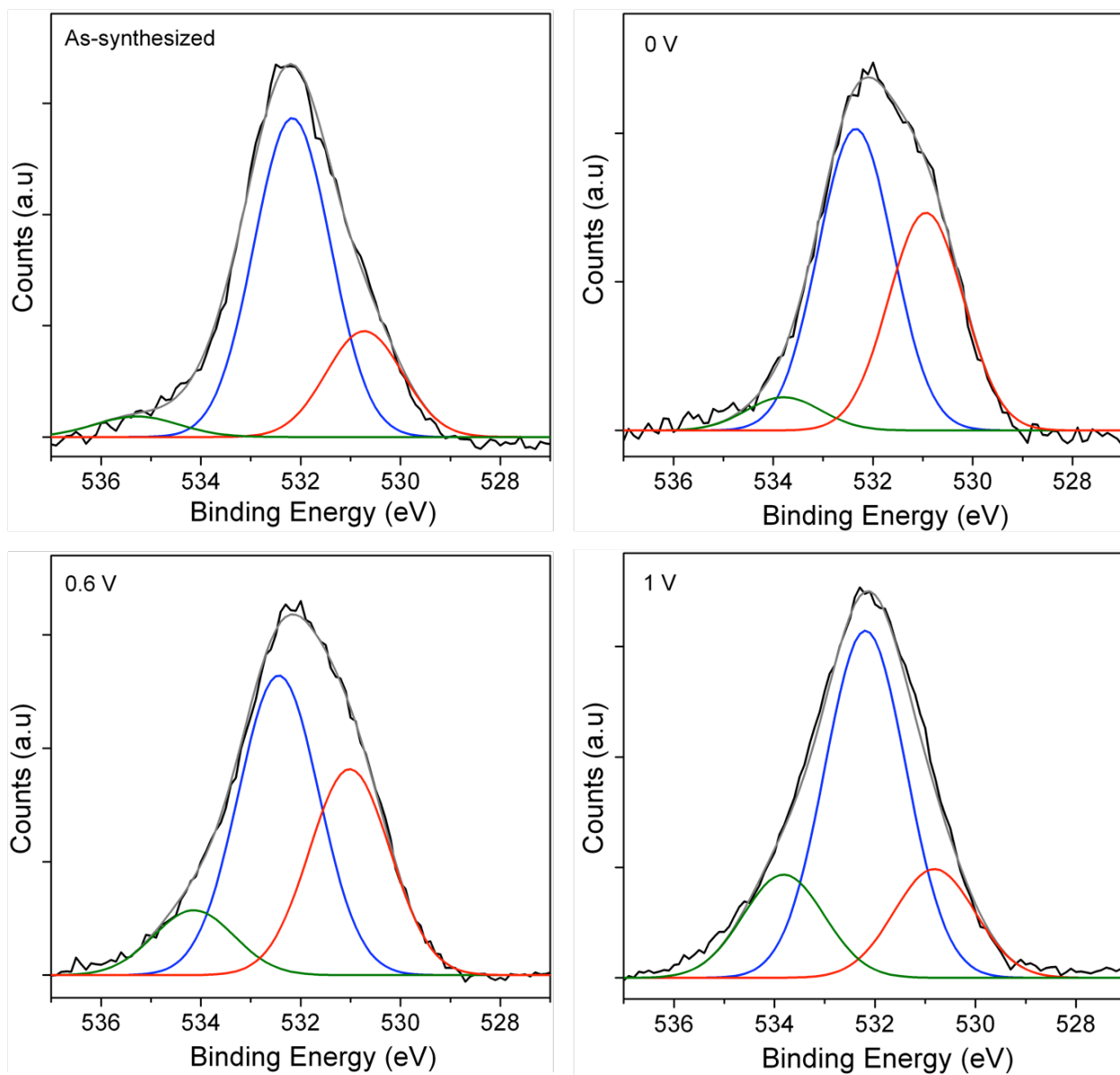


Fig. S10 O 1s spectra of rGO/AuNW before (as-synthesized) and after CV scans at different positive potential limit. The fitted peaks are C=O at 530.7 eV, C-OH 532.2 at eV and -O-C at 534.4 eV. Indicates that compared to other oxygenated species, relative content of C=O (amide) drastically varies with respect to positive potential limit. The increase in C=O content at 0.6 V may be the reason for featureless voltammogram (absence of redox features).

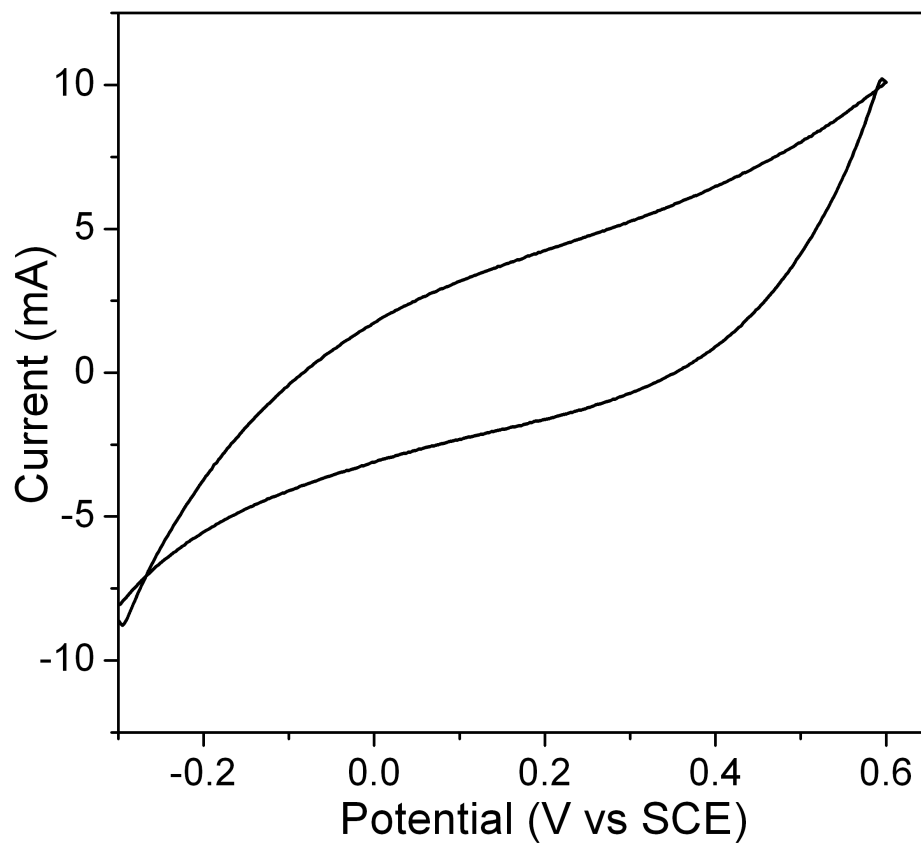


Fig. S11 CV of GO/TiO₂ measured in 1M NaOH containing 1.5 M ethanol, showing the absence of ethanol oxidation peak.

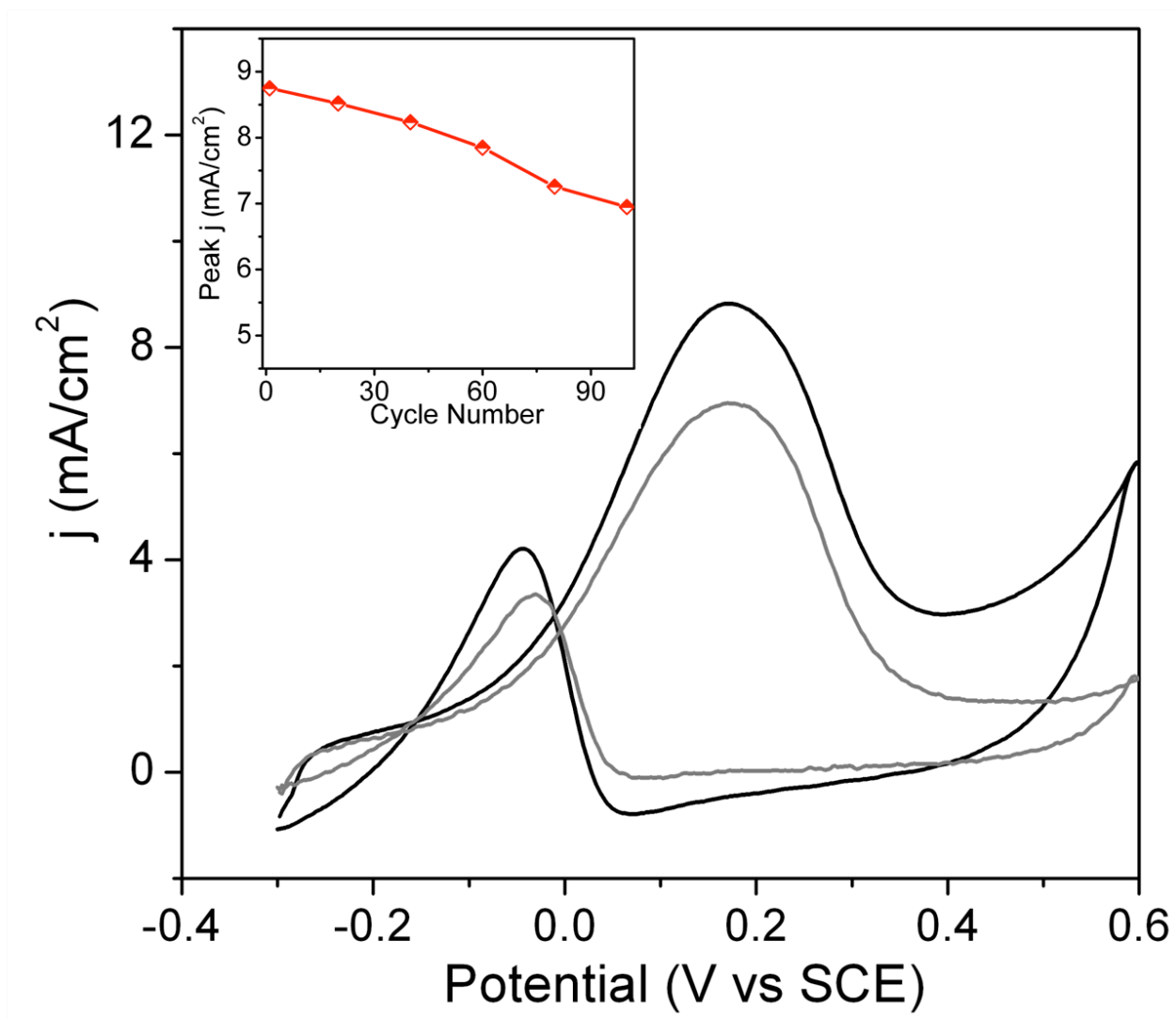


Fig. S12 CV stability test of rGO/TiO₂/AuNW, in 1 M NaOH aqueous solution containing 1.5 M ethanol, 1st and 100th cycle, inset displays peak current density versus cycle number at a scan rate 40 mV/s.

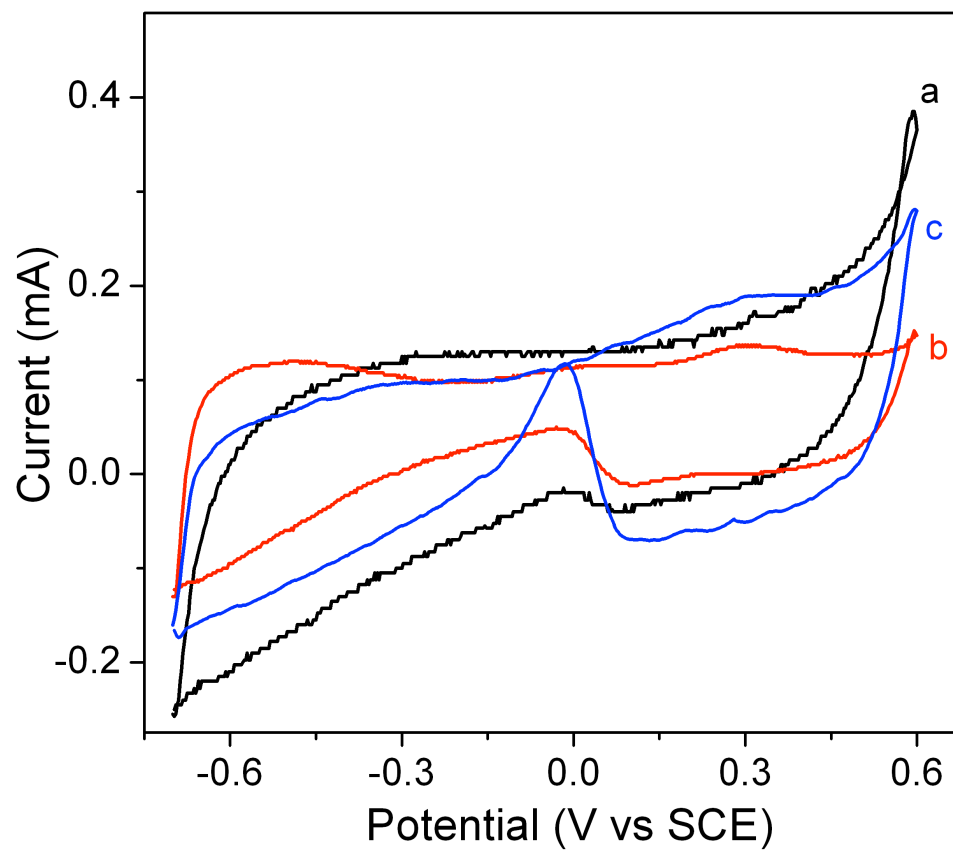


Fig. S13 CVs for CO oxidation obtained in CO saturated 1M NaOH on (a) TiO_2/AuNW , (b) rGO/AuNW and (c) $\text{rGO}/\text{TiO}_2/\text{AuNW}$.

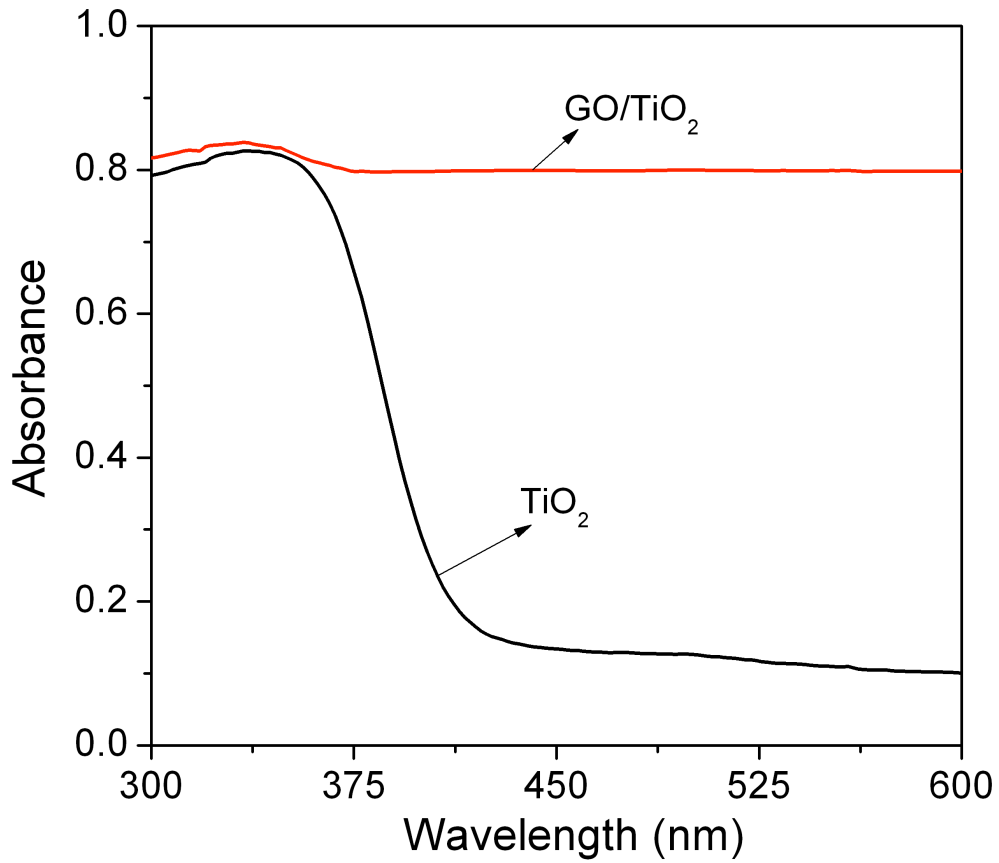


Fig. S14 Diffuse reflectance spectra of TiO₂ and GO/TiO₂. Enhanced absorption in the visible region is attributed to the presence of GO. Expected to boost the utilizing efficiency of solar light.

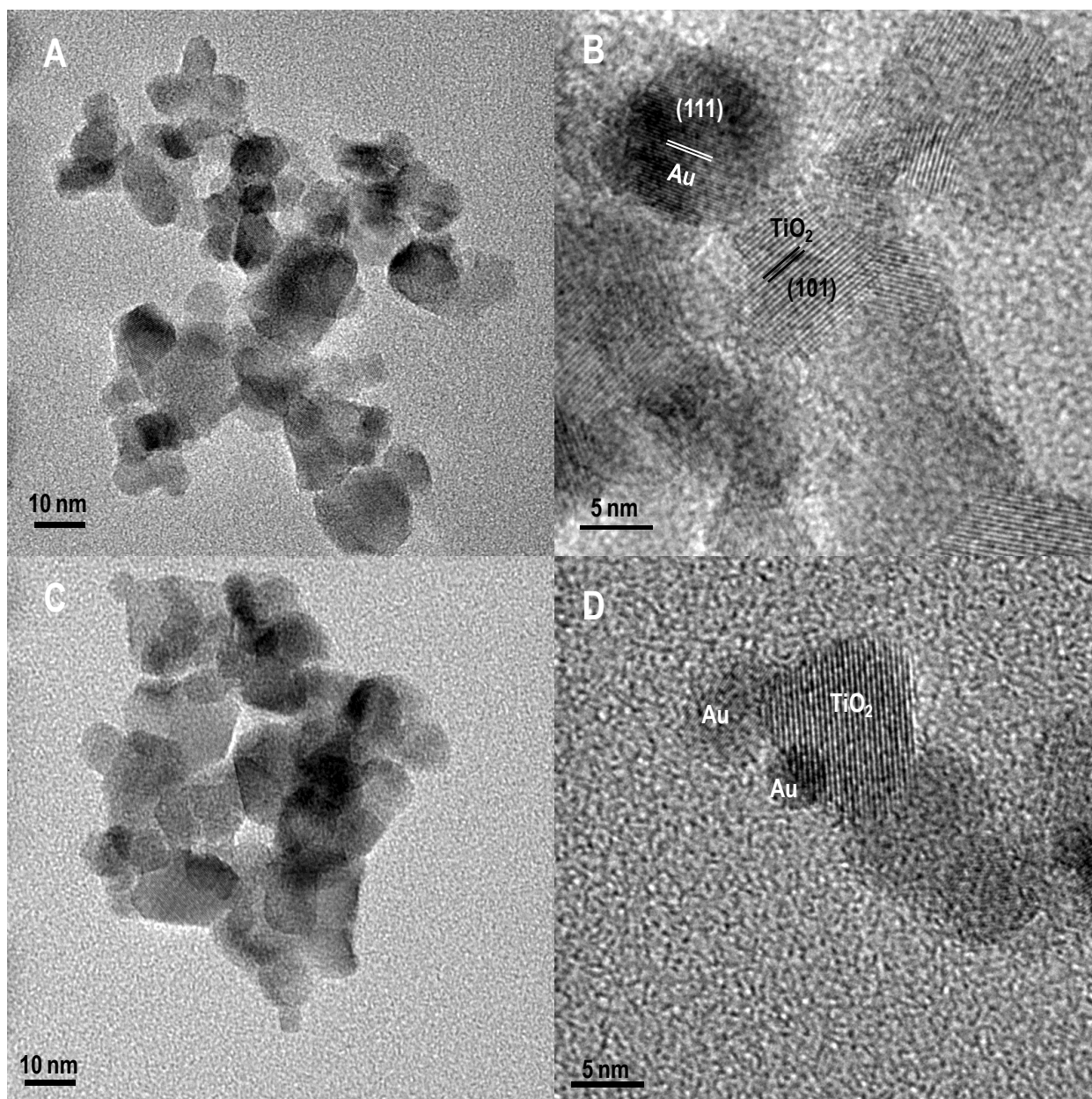


Fig. S15 TEM micrographs (A, B) and (C, D) corresponds to Au nanoparticles nucleated on unmodified and amine modified TiO₂, respectively. Marked d-spacing represents the (101) plane of TiO₂ and (111) plane of Au.

References

1. I. K. Moon, J. Lee, R. S. Ruoff and H. Lee, *Nat. Commun.*, 2010, **1**, 73.
2. T. Ohsaka, F. Izumi and Y. Fujiki, *J. Raman Spectrosc.*, 1978, **7**, 321-324.