## **Supplementary Material**

## Recyclable Textiles Functionalized with Reduced Graphene Oxide@ZnO for Removal of Oil Spills and Dye Pollutants

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## *Synthesis of PVP-capped ZnO nanoparticles*

Briefly, 0.27 g of zinc acetate was dissolved in ethanol and heated to 60 °C under constant stirring for 2 hours. Then, 0.5 g of PVP was added to the solution. A sodium hydroxide solution was prepared separately by dissolving 0.098 g of sodium hydroxide in 50 mL ethanol. The sodium hydroxide solution was added drop-wise into the zinc acetate solution. The solution mixture was then stirred continuously at 60 °C for up to 2 hours.

## Preparation of GO dispersion

Briefly, 10 g graphite flake and 7.5 g sodium nitrate were added to 300 mL of sulfuric acid (98%), and then 40 g of potassium permanganate was added in 1 hour. The mixture was stirred at room temperature for 3 days. Then 1 L of hydrogen peroxide solution (1% in water) was added into the mixture. Subsequently, the mixture was filtered and washed with deionized water. The resulting black cake was re-dispersed in de-ionized water to give a dark brown dispersion, which was subjected to dialysis for one week. The brown suspension was dried at 40 °C under vacuum. GO suspension was obtained by sonicating the as-prepared solid in water. The resulting homogenous brown dispersion was used for the assembly of GO@ZnO films.

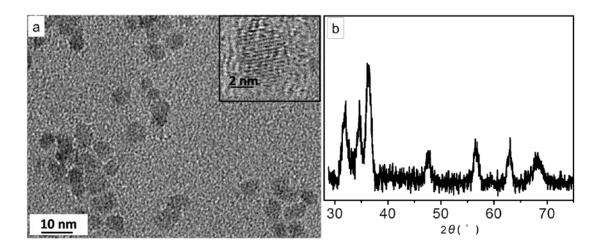


Fig. S1. (a) TEM image of PVP-capped ZnO nanoparticles. (b) XRD pattern of PVP-capped ZnO nanoparticles.

XRD spectra confirmed that the ZnO nanoparticles had the wurtzite-type crystal structure.

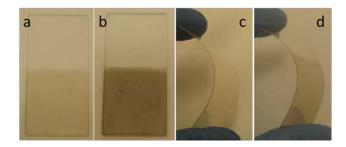


Fig. S2. Optical images of a (GO@ZnO)<sub>5</sub> multilayer film prepared on a quartz substrates (a) before and (b) after 3 hours of UV photoreduction; on a flexible PET substrate (c) before and (d) after 3 hours of UV photoreduction.

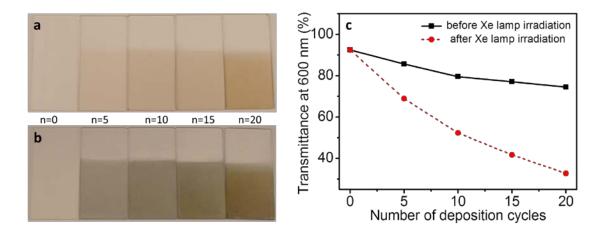


Fig. S3. Photographs of (GO@ZnO)<sub>n</sub> prepared on quartz slides with different numbers of layers: (a) before and (b) after the UV-induced photoreduction process for 3 hours. (c) Optical transmittance at 600 nm as a function of the number of layers before and after photoreduction for 3 hours.

After the photoreduction process, the colour of the slides darkened due to the formation of RGO as shown in Fig. S3b. Fig. S3c shows that the UV-vis transmittance of the hybrid films after photoreduction.

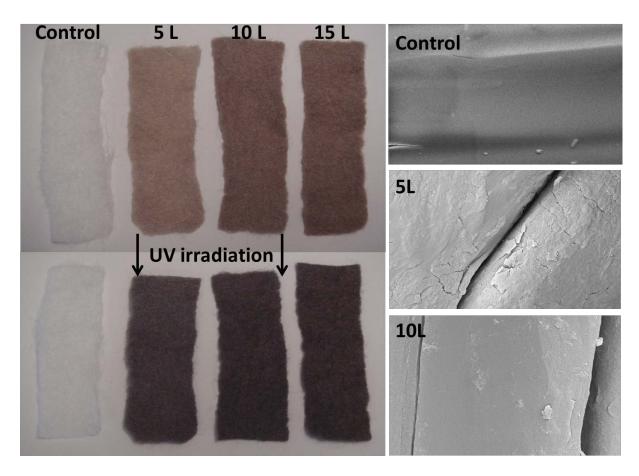


Fig. S4. Appearances of (GO@ZnO)n/PET fabrics (n=0, 5, 10 and 15) before and after UV-photoreduction for 3 hours. SEM images of (RGO@ZnO)n coated on PET fabrics and PET single fibre with n=0, 5 and 10.

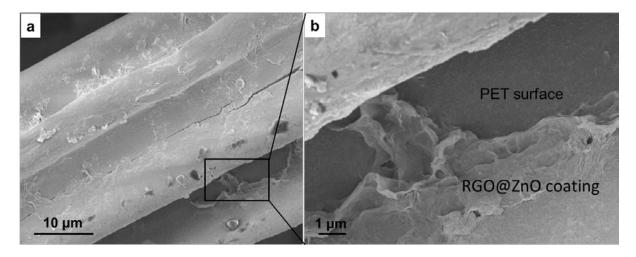


Fig. S5. SEM images of (GO@ZnO)<sub>15</sub>/PET fabrics.

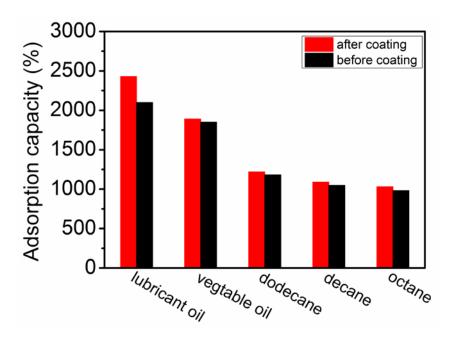


Fig. S6. Adsorption capacity of PET fabric before and after coating with RGO@ZnO in terms of weight gain.

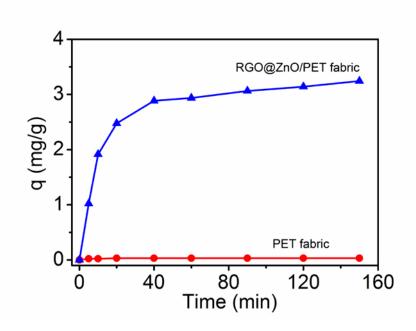


Fig. S7. Dye adsorption kinetics of RhB on RGO@ZnO/PET fabric. C (RhB) $_{initial}$  = 12 ppm T=293K

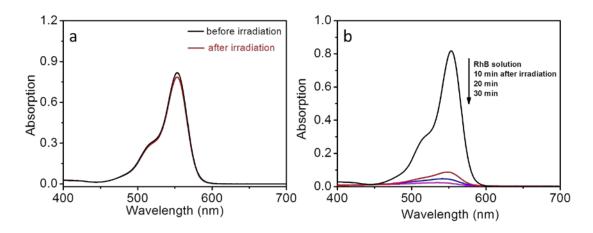


Fig. S8. (a) UV-vis spectra of RhB solution before and after simulated sunlight irradiation for 30 min without (RGO@ZnO)<sub>10</sub>/PET fabric. (b) The UV-vis spectra of the original RhB solution and the RhB solution after simulated sunlight irradiation for different times with the presence of (RGO@ZnO)<sub>10</sub>/PET fabric.