## **Supplementary information**

## **A facile synthesis of porous graphene for efficient water and wastewater treatment**

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## **Materials and reagents**

NaNO<sub>3</sub> (Product No. S5506), H<sub>2</sub>SO<sub>4</sub> (95.0-98.0%, Product No. 320501), KMnO<sub>4</sub> (Product No. 223468),  $H_2O_2$  (30 wt%, Product No. 216763), hydrazine (35 wt%, product No. 309400), sodium fluoride (product No.: 7681-49-4), sodium (meta) arsenite (product No.: S7400), potassium nitrate salt (product No.: 542040) were purchased from Sigma-Aldrich. HCl (36 wt%, product No.: 7647-01-0), was purchased from Acros. Graphite flake (GFs, product No.17346-25) and 0.2 micron membrane filter were purchased from Thermo and Fisher Scientific suppliers. Fluoride kit (Product No.: 90734, test fluorure was purchased from MACHEREY-NAGEL GmbH & Co. KG, Germany).



**Figure S1:** Photographs of the stages in producing PG. (a) Oxidation and exfoliation of graphite in the presence of oxidizing agents. (b) GO dispersed in water prior to reduction. (c) Reduction of GO using hydrazine as reducing agent followed by thermal treatment results in the as-prepared PG in powder form.



**Figure S2:** Exfoliation of graphite flakes. (a-d) SEM images of the raw material graphite flakes. Images of the residual graphite sediment, showing many disintegrated and exfoliated graphite flakes.



**Figure S3:** (a-c) SEM images of as-prepared GO. The flaky texture reflects the layered structure of GO.

**Table S1**: Parameters obtained from nitrogen desorption isotherms.

**SA<sup>a</sup>**  $\left[\text{m}^2 \text{g}^{-1}\right]$  **PV**<sup>b</sup>  $[\text{cm}^3 \text{g}^{-1}]$  **PD<sup>c</sup>**  $[\text{nm}]$ 

653 m<sup>2</sup>/<sub>g</sub> 
$$
\begin{array}{c} 4.904e-01 \\ cc/g \end{array}
$$
 1.5037 nm

a BET surface area.

b Pore volume: DFT desorption cumulative volume of the pores.

c Pore diameter: average pore diameter determined by the BJH method.

Micropore Volume : 0.187 cc/g

Total pore volume  $= 4.904e-01$  cc/g for

Average pore Radius =  $1.50370e+01 \text{ Å} = 1.5037 \text{ nm}$ 



**Figure S4**: Fluoride kit testing of fluoride removal from drinking water. (a) The standard concentration of fluoride in aqueous solutions using fluoride kit varying from 0, 2, 5, 10, 20, 50, 100mg/l fluoride in water. (b) 100mg/l fluoride sample tested using PG over the exposure time of 5, 10, 15, 20, 30, 60 mins, 24 hrs and 48 hrs. It is found that adsorption process has been completed within 60 mins using PG as adsorbent.



Figure S5: Photographs showing the removal process of pump oil. (a) 1 g oil in 50 ml water, (b) immersion of 2.5 mg PG contained in a porous bag for oil removal, and (c) the oilabsorption after 10 sec.



**Figure S6**: Time evolution of MB adsorption properties of PG after the fifth regeneration cycle. (a) Adsorbance as a function of wavelength. (b) Adsorption capacity. The data show the complete removal of MB within 240 mins.



**Figure S7**: Recyclability of PG for the adsorption of MB and RB.



**Table S2:** Analysis of the results of adsorption experiments of arsenic, RB and MB on PG

**Table S3.** Comparison of the adsorption capacities and performance of different contaminants from water between PG and previous reported different adsorbents.





**Figure S8**: Elemental mapping of fluoride adsorption using an SEM. (a) electron micrograph of PG after adsorption. (b-d) Mapping of carbon, oxygen and fluoride over the area highlighted in  $(a)$ .



**Figure S9**: Raman spectroscopy of PG before adsorption and after regenration PG

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