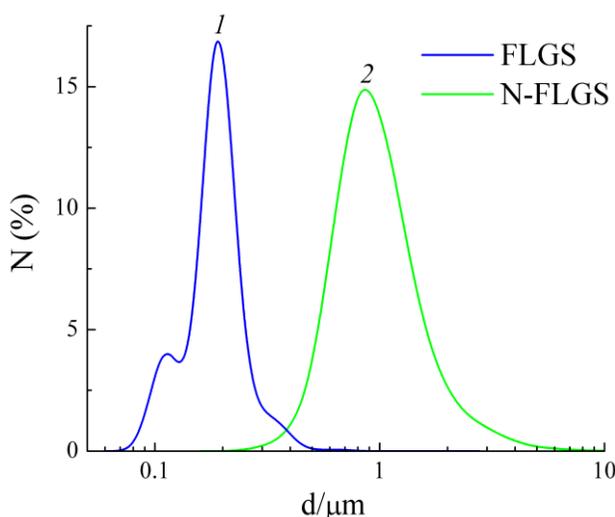


## Effect of graphene surface functionalization on the oxygen reduction reaction in alkaline media

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### 1. Size distribution of FLGS particles

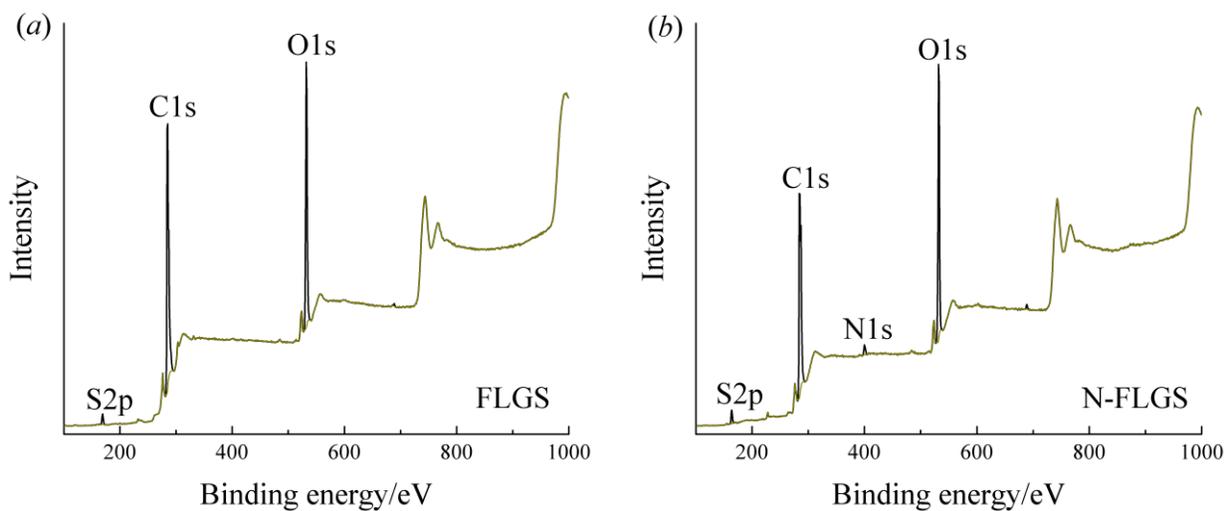
The size distribution of FLGS particles was obtained using an LS 13 320 XR laser particle size analyzer (Beckman Coulter, USA).



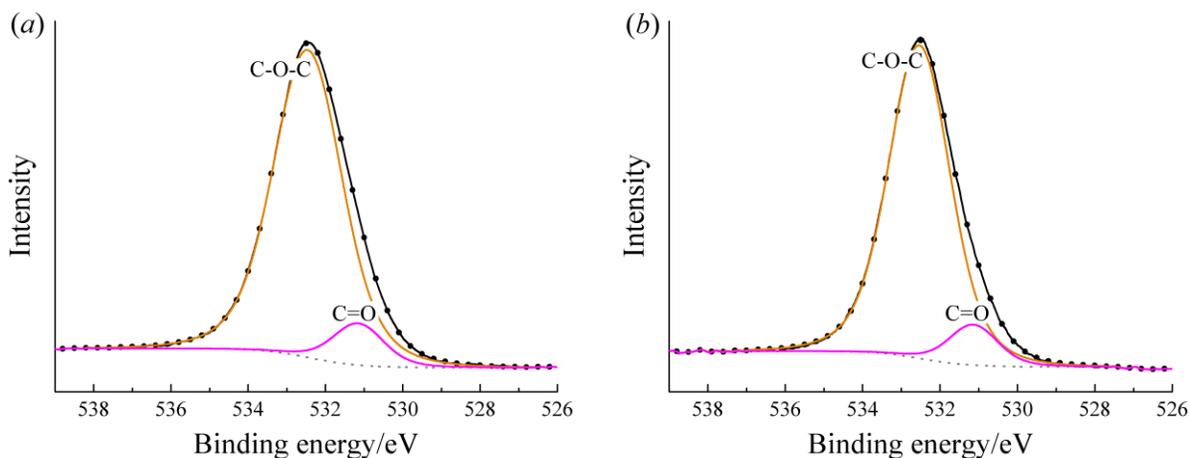
**Figure S1** Size distribution of FLGS (1) and N-FLGS (2) particles.

### 2. XPS analysis of FLGS surface

The XPS spectra were obtained using a Specs PHOIBOS 150 MCD spectrometer (Specs, Germany) with Mg  $K\alpha$  radiation (1253.6 eV). Pressure in a working chamber of a spectrometer was within  $4 \cdot 10^{-8}$  Pa. The probed area was 300–700  $\mu\text{m}^2$ , depth of analysis was 1–2 nm.



**Figure S2** Survey XPS spectra of FLGS (a) and N-FLGS (b).



**Figure S3** O1s high resolution XPS spectra of FLGS (a) and N-FLGS (b)

**Table S1** Surface elemental composition of FLGS and N-FLGS samples.

sample	C 1s, % (284 eV)	O 1s, % (532 eV)	N 1s, % (400 eV)	S 2p, % (169 eV)
FLGS	79.9	18.8	-	1.3
N-FLGS	75.4	20.7	1.4	2.5
sample	C=O, % (531.4 eV)	C-O-C, % (532.3 eV)	pyrrolic-N, % (399.0 eV)	pyridinic-N, % (400.0 eV)
FLGS	1.7	17.1	-	-
N-FLGS	1.9	18.8	0.7	0.7

### 3. Oxygen reduction reaction on GC, FLGS and N-FLGS in air-saturated 0.1 M KOH solution

To determine the number of electrons participating in ORR at the covered with FLGS and the pristine GC electrodes, the  $j,E$ -dependencies were measured for various  $\omega$  [Figures S4(a), S4(b) and S4(c)]. The values of  $n$  were calculated from the  $j,E$ -curves analysis with the Koutecký-Levich equation:<sup>S1</sup>

$$\frac{1}{j} = \frac{1}{j_k} + \frac{1}{j_d} \quad (1)$$

$$j_k = nFkc^0 \quad (2)$$

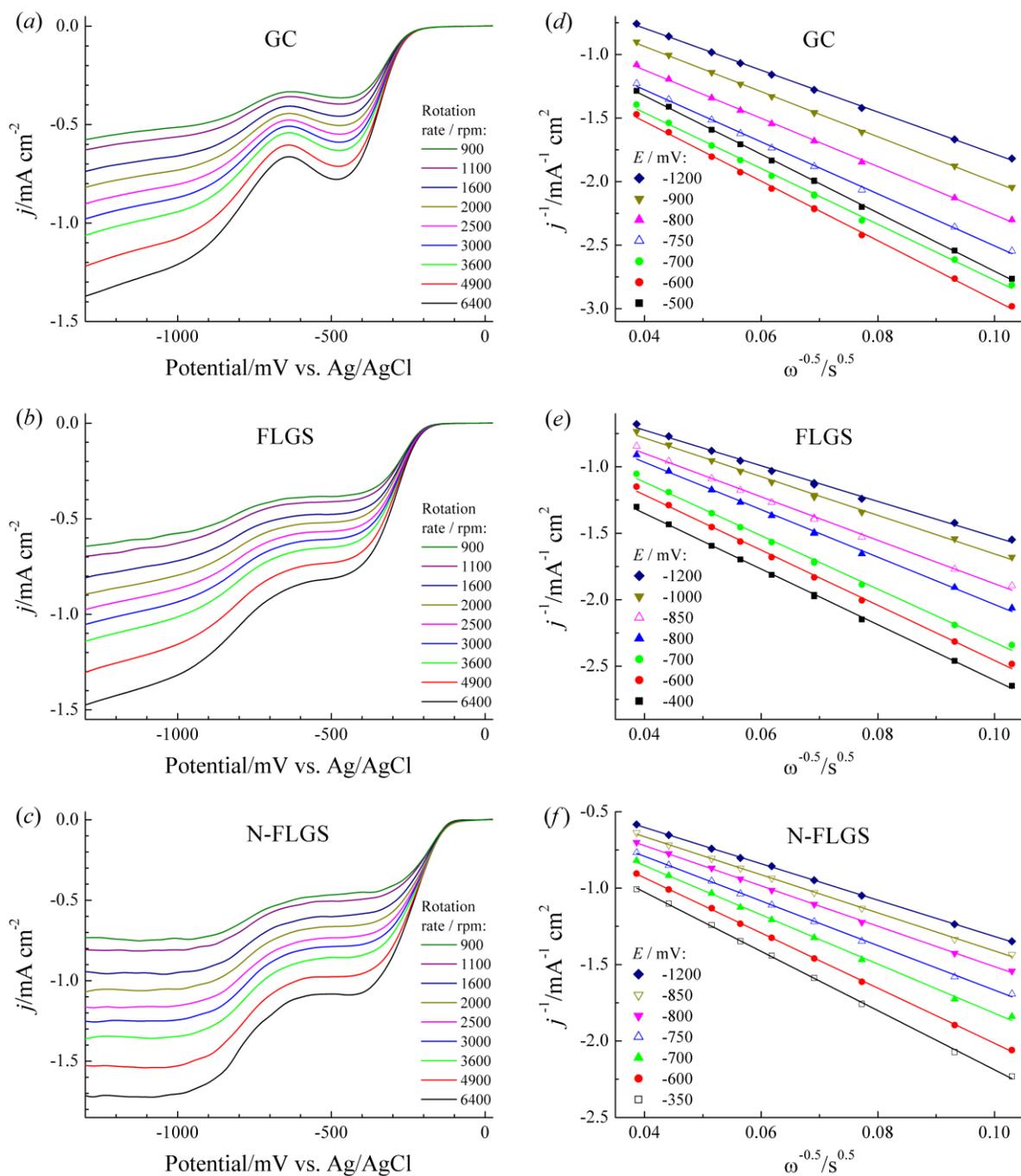
$$j_d = 0.62nFD^{2/3}\omega^{1/2}\nu^{-1/6}c^0, \quad (3)$$

where  $j_k$  and  $j_d$  are densities of the kinetic and the limiting diffusion current, respectively,  $[j] = \text{mA}\cdot\text{cm}^{-2}$ ;  $k$  is a rate constant of oxygen reduction reaction,  $[k] = \text{cm}\cdot\text{s}^{-1}$ ;  $\omega$  is an angular velocity of electrode rotation,  $[\omega] = \text{rad}\cdot\text{s}^{-1}$ ;  $F$  is Faraday constant,  $F = 96485 \text{ C}\cdot\text{mol}^{-1}$ ;  $D$  is a coefficient of oxygen diffusion in a 0.1 M KOH solution,  $D = 1.9\cdot 10^{-5} \text{ cm}^2\cdot\text{s}^{-1}$ ;  $\nu$  is kinematic viscosity of the 0.1 M KOH,  $\nu = 0.01 \text{ cm}^2\cdot\text{s}^{-1}$ ;  $c^0$  is bulk concentration of dissolved oxygen,  $c^0 = 0.24\cdot 10^{-3} \text{ mol}\cdot\text{L}^{-1}$  in the 0.1 M KOH solution.<sup>S2,S3</sup>

The values of the limiting diffusion current density were evaluated according to the Levich equation<sup>S1</sup> (equation 3).

#### References

- S1 A. J. Bard and L. R. Faulkner, *Electrochemical Methods: Fundamentals and Applications*, 2nd edn., Wiley, New York, 2001.
- S2 L. T. Qu, Y. Liu, J. B. Baek and L. M. Dai, *ACS Nano*, 2010, **4**, 1321.
- S3 G. Jürmann and K. Tammeveski, *J. Electroanal. Chem.*, 2006, **597**, 119.



**Figure S4** (a) (b) and (c) are  $j$ , $E$ -curves for pristine GC and GC electrodes covered with FLGS and N-FLGS, respectively, at various electrode rotation rates; (d), (e) and (f) are the corresponding  $j$ , $\omega$ -dependencies in the Koutecký-Levich coordinates at selected potentials.