

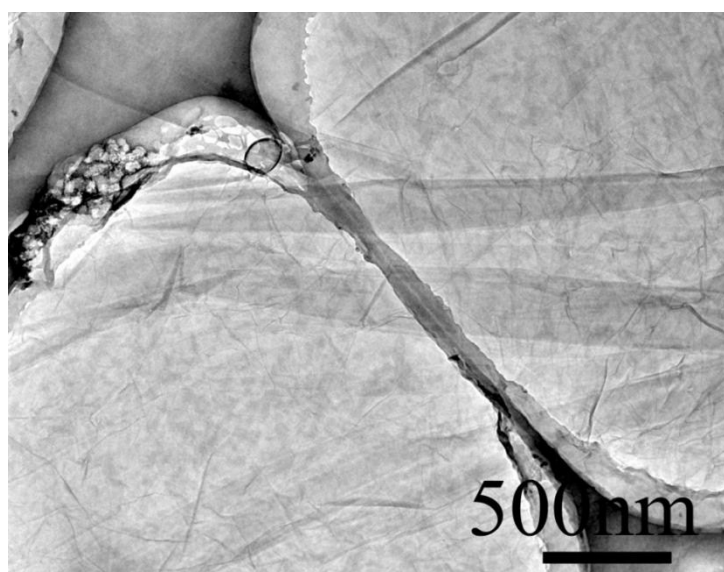
# A Novel Gas Sensor Based on Porous $\alpha$ -Ni(OH)<sub>2</sub> Ultrathin Nanosheet/Reduced Graphene Oxide Composites for Room Temperature Detection of NO<sub>x</sub>

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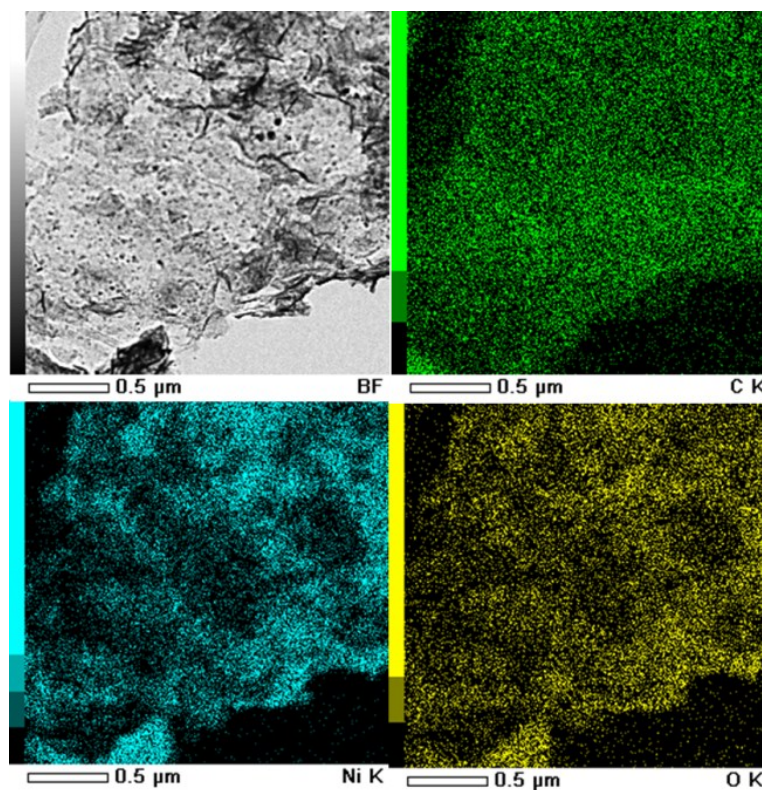
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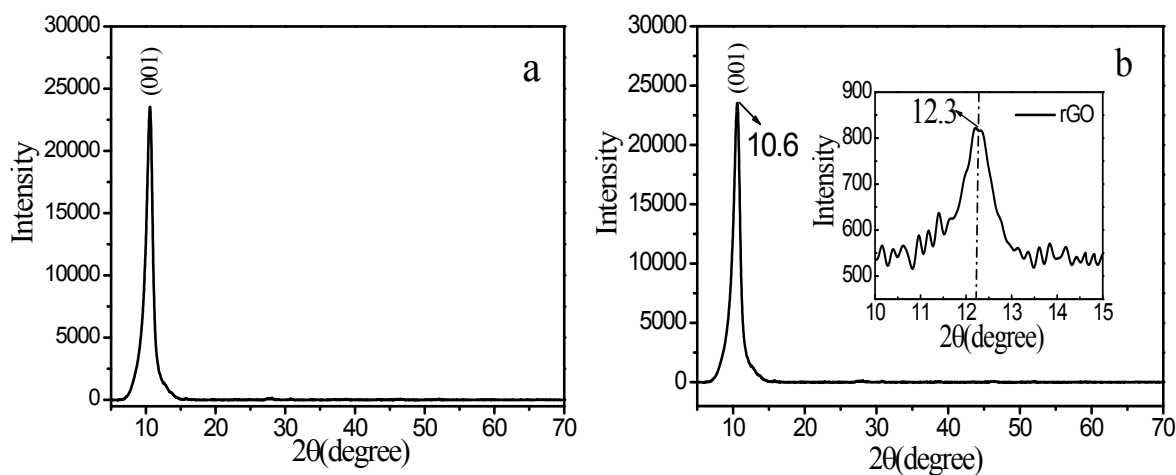


**Figure S1.** TEM image of the prepared GO by modified Hummers method.

**Figure S1** shows the TEM image of the prepared GO by modified Hummers method. The GO is thin and transparent.



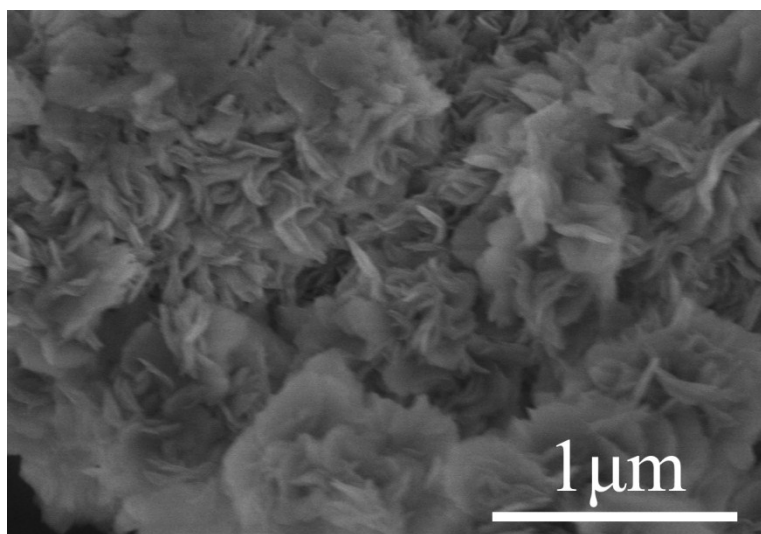
**Figure S2.** Elemental mapping of C2 sample: a) Bright field image; b), c) and d) corresponding C, Ni and O elemental mapping.



**Figure S3.** The XRD patterns of (a) the as-prepared GO by a modified Hummers method, and (b) the synthesized rGO obtained by reflux in the absence of the  $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ .

The weak diffraction peak at  $2\theta = 12.2^\circ$  can be indexed to the (001) plane of graphite oxide, and the

broad diffraction peak at  $2\theta$  of  $24.5\text{-}27.5^\circ$  can be indexed to the (002) planes of the graphite (G) in Figure. S3a. Figure. S3b shows the XRD pattern of the rGO. In Figure. S3b, the (001) peak of graphite oxide was greatly reduced, and the very broad (002) peak of graphite can be seen, suggesting that the graphite component is very poorly ordered along the stacking direction.



**Figure S4.** SEM image of the pure  $\alpha\text{-Ni(OH)}_2$  obtained by reflux.

**Table S1.** Data of sensitivity of  $\alpha\text{-Ni(OH)}_2/\text{rGO}$  nanocomposites with different mass ratio of  $\text{Ni(NO}_3)_2\cdot\text{H}_2\text{O}/\text{GO}$  exposed to  $\text{NO}_x$  with decreasing concentrations.

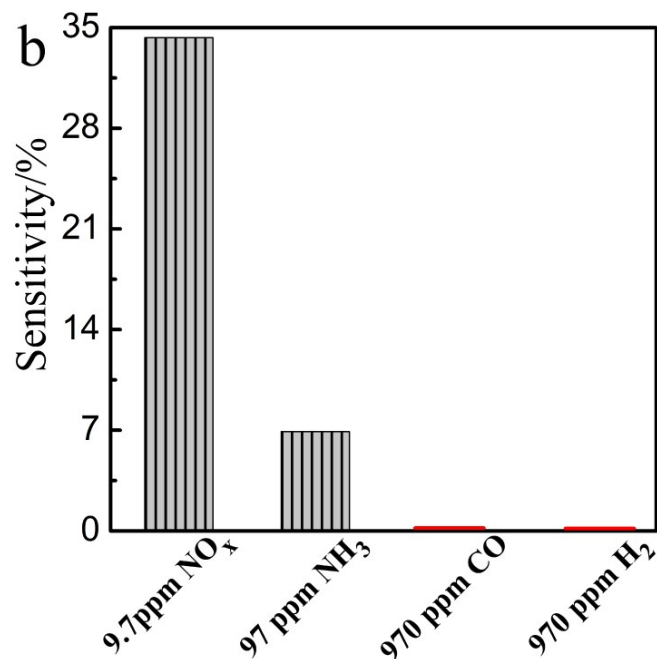
Concentration of $\text{NO}_x$ ppm	S/%	S/%	S/%	S/%
	C1	C2	C3	C4
97.0	29.3	64.4	50.0	12.4
68.1	22.8	61.5	44.5	11.3
48.5	19.0	56.5	41.0	10.0

29.1	17.0	49.1	33.9	8.6
9.7	11.8	40.7	28.6	6.7
4.85	9.0	31.0	23.5	--
2.91	6.7	23.2	17.8	--
0.97	4.3	16.5	11.2	--

**Table S2.** Data of response time of  $\alpha$ -Ni(OH)<sub>2</sub>/rGO nanocomposites with different mass ratio of

Ni(NO<sub>3</sub>)<sub>2</sub>·H<sub>2</sub>O/GO exposed to NO<sub>x</sub> with decreasing concentrations.

Concentration of NO <sub>x</sub> ppm	t/s	t/s	t/s	t/s
	C1	C2	C3	C4
97.0	9	10	12	16
68.1	10	12	14	18
48.5	10	12	16	18
29.1	11	13	15	20
9.7	10	12	16	21
4.85	12	13	16	--
2.91	16	20	40	--
0.97	48	40	56	--



**Figure S5.** The bar graphs for gas selectivity of the C2 sensor to NO<sub>x</sub>, NH<sub>3</sub>, CO and H<sub>2</sub>

**Table S3.** Fitted impedance parameters of samples

Samples	$R_{\Omega}(\Omega)$	$C(\text{F cm}^{-2})$	$R_{ct}(\Omega)$
$\alpha\text{-Ni(OH)}_2$	579.5	$5.57 \times 10^{-6}$	$2.74 \times 10^3$
C2	772	$1.91 \times 10^{-6}$	$3.10 \times 10^3$
rGO	661.8	$2.15 \times 10^{-8}$	$0.26 \times 10^3$

The result of the impedance of the  $\alpha\text{-Ni(OH)}_2$ , C2 and rGO electrodes are shown in Table S3. The results demonstrate that the C2 sample have the Rct of  $3.10 \times 10^3 \Omega$ , which exhibits much higher conductivity than the pure  $\alpha\text{-Ni(OH)}_2$  sample. Therefore, the C2 sample can improve the electron transportation from the sensor to the adsorbate's NO<sub>x</sub> surface, and consequently.