Supplementary Information

Layered mesoporous Mg(OH)₂/GO nanosheet composite for efficient

removal of water contaminants

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Digital pictures of GO synthesized by Hummers' method and colloid after laser ablated were shown in Figure S1. Mg(OH)₂ and Mg(OH)₂/GO composite colloids were both turbid compared with transparent GO solution. Mg(OH)₂ turns to be turbid because of the nanoflakes nature of Mg(OH)₂, but the reason of Mg(OH)₂/GO composite colloids become cloudy is electrostatic interaction between positively charged and negatively charged GO. Figure S2 shows the hexagonal stack morphology and XRD pattern of Mg(OH)₂ nanoflakes. All peaks coincide with PDF No. 00-44-1482, except that the strongest peak is not (101) but (001) and all peaks shows non-uniform broadening, which is caused by various disorders, such as stack faults or interstratifications in layered materials.¹ It is obvious that MB is a cationic dye from the MB molecular structure, as shown in Figure S3. Digital pictures of MB solution after different contact time were shown in Figure S4. And the color becomes lighter gradually. Table S1-S4 shows pseudo-second-order model adsorption kinetics constants, intro-particle diffusion model constants, equilibrium adsorption isotherm fitting parameters, and specific data of heavy metal ions adsorption respectively.



Figure S1. Digital pictures of (a) GO solution, (b) as-prepared $Mg(OH)_2/GO$ composite, (c) pure $Mg(OH)_2$.



Figure S2. SEM morphology and XRD pattern of the pure Mg(OH)₂ nanoflakes.



Figure S3. The molecule structure of MB



Figure S4. Digital pictures for the MB solution (100 mg L^{-1} , 20mL) and those after treatment with Mg(OH)₂/GO composite (20mg) at different time.

Table 51. 1 seudo-second-order model adsorption kineties constants									
$q_{e,cal} (mg g^{-1})$	$x_2 (\times 10^{-4} \text{ g mg}^{-1} \text{ min}^{-1})$	$V_0 (g mg^{-1} min^{-1})$	\mathbb{R}^2						
87.7	3.88	2.99	0.998						
Table S2. Intro-particle diffusion model constants and correlation coefficients $k_{\rm er}$ (mg g ⁻¹ min ^{-1/2}) $C_{\rm er}$ \mathbf{P}^2									
12.07	18.03	1							
2.03	50.95 0.9		40						
0.49	65.3	0.96	57						
	$\frac{q_{e,cal} (mg g^{-1})}{87.7}$ article diffusion mo k _{di} (mg g^{-1} min 12.07 2.03 0.49	accord order model adsorption kniede. $q_{e,cal} (mg g^{-1})$ $k_2 (\times 10^{-4} g mg^{-1} min^{-1})$ 87.7 3.88 article diffusion model constants and cor $k_{di} (mg g^{-1} min^{-1/2})$ C_i 12.07 18.03 2.03 50.95 0.49 65.3	article diffusion model constants and correlation coefficient k_{di} (mg g ⁻¹) k_2 (×10 ⁻⁴ g mg ⁻¹ min ⁻¹) V_0 (g mg ⁻¹ min ⁻¹) 87.7 3.88 2.99 article diffusion model constants and correlation coefficient k_{di} (mg g ⁻¹ min ^{-1/2}) C_i R^2 12.07 18.03 1 2.03 50.95 0.94 0.49 65.3 0.96						

 Table S1. Pseudo-second-order model adsorption kinetics constants

Linear fitting			
Langmuir isotherm -	$q_m (mg g-1)$	$k_L (L mg^{-1})$	R ²
	555.6	0.00966	0.967
Freundlich isotherm -	1/n	$k_{\rm F}$	R ²
	0.36	39.8	0.993
Non-linear fitting			
Langmuir isotherm –	$q_m (mg g-1)$	$k_L (L mg^{-1})$	R ²
	609.5	0.0048	0.967
Freundlich isotherm –	1/n	k _F	R ²
	0.39	34.2	0.992

Table S3. Equilibrium adsorption isotherm fitting parameters

Table S4. Equilibrium adsorption capacity and removal efficiency data of heavy metal ion adsorbed by Mg(OH)₂/GO nanoflowers

Metal species	Zn^{2+}	Pb ²⁺	Cu ²⁺	Ni ²⁺
Equilibrium adsorption capacities (mg g ⁻¹)	327.7	344.4	215.5	174.7
Removal capacity (%)	99.9	99.6	62.1	57.2

Reference

 Liang, Changhao, et al. Pulsed-laser ablation of Mg in liquids: surfactant-directing nanoparticle assembly for magnesium hydroxide nanostructures. Chemical Physics Letters 389.1 (2004): 58-63.