

## Preparation of Phosphorylated Graphene Oxide-Chitosan Composite for Highly Efficient Removal of U(VI)

Yawen Cai<sup>1</sup>, Shitong Yang<sup>1</sup>, Shuao Wang\*<sup>1</sup>

<sup>1</sup> School for Radiological and Interdisciplinary Sciences (RAD-X), Soochow University, Suzhou, 215123

\*Email:shuaowang@suda.edu.cn

Uranium is not only a strategic resource for the nuclear power but also a highly toxic contaminant in the environment.<sup>1</sup> Although a series of traditional capturing materials including zeolite, metal-organic framework, mesoporous silica, and carbon-based nanomaterials have been developed, the combined advantages of decent stability, ultrafast removal kinetics, high sorption capacity, great selectivity, and potential recyclability have yet to be integrated into a single material.<sup>2-6</sup> Herein, a new synthesis strategy was developed to synthesize a novel phosphorylated graphene oxide (GO)-chitosan (CS) composite (denoted as GO-CS-P) for U(VI) removal. Batch experiments and spectroscopic analysis were performed to explore the removal performance and mechanism of GO-CS-P towards U(VI). The results showed that with the presence of the GO-CS-P composite, the concentration of U(VI) in the aqueous solution can be reduced to an extremely low level within a short time of 15 min. In addition, the maximum sorption capacity of U(VI) on GO-CS-P was calculated to be ~779.44 mg/g, which is higher than those on GO, GO-CS and a series of currently reported adsorbents. Moreover, GO-CS-P also exhibited high selectivity in capturing U(VI) from a mixture containing multiple competing metal ions. According to the desorption experiments, Fourier transform infrared spectroscopy (FTIR), X-ray absorption spectroscopy (XAS) and X-ray photoelectron spectroscopy (XPS) analysis, the highly efficient immobilization of U(VI) in GO-CS-P was predominantly controlled by inner-sphere surface complexation with a minor contribution of surface reduction. These observations indicate that the GO-CS-P composite can be used for nuclear fuel partition and uranium-bearing wastewater remediation.

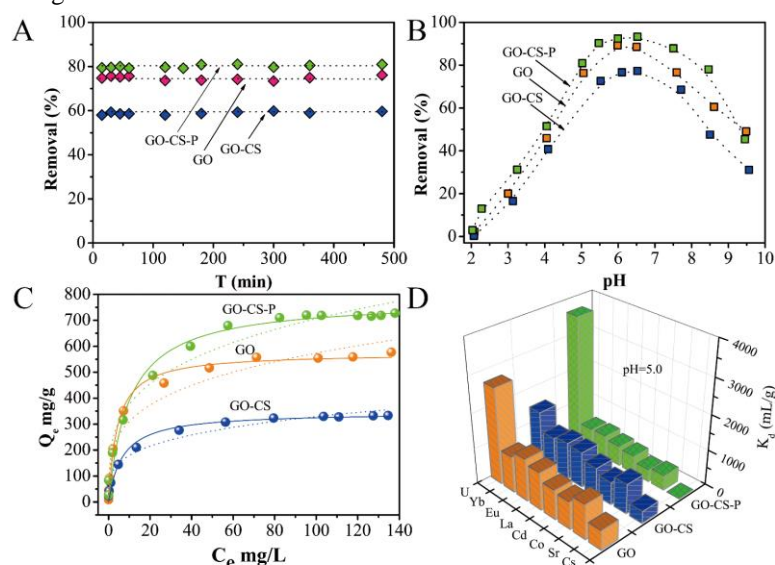


Fig.1. (A) Time-dependent sorption behaviors of U(VI) onto GO, GO-CS and GO-CS-P.  $T = 293$  K,  $\text{pH} = 5.0$ ,  $m/V = 0.05$  g/L,  $C_{\text{U(VI)initial}} = 5.0 \times 10^{-5}$  mol/L,  $I = 0.01$  mol/L  $\text{NaNO}_3$ ; (B) Effect of solution pH on the removal of U(VI) by GO, GO-CS and GO-CS-P.  $T = 293$  K,  $m/V = 0.05$  g/L,  $C_{\text{U(VI)initial}} = 5.0 \times 10^{-5}$  mol/L,  $I = 0.01$  mol/L  $\text{NaNO}_3$ ; (C) Sorption isotherms, Langmuir and Freundlich model fits of U(VI) on GO, GO-CS and GO-CS-P.  $T = 293$  K,  $\text{pH} = 5.0$ ,  $m/V = 0.05$  g/L,  $I = 0.01$  mol/L  $\text{NaNO}_3$ . Symbols represent the experimental data, solid lines represent Langmuir model fits and dash lines represent Freundlich model fits; (D) Sorption selectivity of GO, GO-CS and GO-CS-P towards multiple metal ions.  $T = 293$  K,  $\text{pH} = 5.0$ ,  $m/V = 0.05$  g/L,  $I = 0.01$  mol/L  $\text{NaNO}_3$ ,  $C_0 = 5.0 \times 10^{-4}$  mol/L for a single metal ion.

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