

Ultra-rapid removal of per- and poly fluorinated alkyl substances from water using cyclodextrin polymer networks

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Abstract: Herein, we developed a β -cyclodextrin (β -CD)-based polymer (β -CD-TriPod) crosslinked with tripodal amine to demonstrate the synergetic effects in superior adsorption of both short- and long-chain per- and polyfluoroalkyl substances (PFASs). Kinetics studies showed rapid adsorption ($\sim 100\%$ for nine PFASs at $1 \mu\text{g L}^{-1}$, except PFBA, and $>86\%$ at $200 \mu\text{g L}^{-1}$ individually) within two minutes. Isotherm results showed exceptional adsorption affinity and capacity, with $K_L = 0.310 \pm 0.180 \text{ L mg}^{-1}$, $q_m = 246.20 \pm 14.80 \text{ mg g}^{-1}$ for PFBS, and $K_L = 0.980 \pm 0.260 \text{ L mg}^{-1}$, $q_m = 587.10 \pm 54.50 \text{ mg g}^{-1}$ for PFOS, significantly outperforming traditional activated carbons (ACs) and resins. The adsorbent also exhibited excellent regeneration and reusability, maintaining stable performance ($>94\%$) over five consecutive adsorption-desorption cycles. Additionally, it performed effectively in PFASs-spiked real industrial wastewater with 55-100% removal efficiencies, regardless of the presence of co-contaminants. The adsorption mechanism confirmed the combined role of hydrophobic inclusion within β -CD cavities and electrostatic interactions with amines groups using elemental mapping, composition and FTIR techniques. Overall, this work demonstrates advanced molecular design strategies for rapid PFASs removal, establishing β -CD-TriPod as a highly regenerable and promising adsorbent for the rapid and efficient treatment of PFASs-contaminated water and industrial wastewater.

Keywords: Short-chain PFASs; β -CD polymerization; Adsorption affinity; Wastewater treatment.