Targeted Arsenic Sequestration from Natural Waters with a Functionalized Adsorbent

Patricia Pachedo Umpire

Escuela de Ingeniería Ambiental, Universidad Católica San Pablo, Quinta Vivanco s/n, Urb. Campiña Paisajista, Arequipa, Peru

Katherine Cáceres Huaylla

Escuela de Ingeniería Ambiental, Universidad Católica San Pablo, Quinta Vivanco s/n, Urb. Campiña Paisajista, Arequipa, Peru

Juliana Santos dos Santos

Escuela de Ingeniería Ambiental, Universidad Católica San Pablo, Quinta Vivanco s/n, Urb. Campiña Paisajista, Arequipa, Peru

Javier Montalvo Andiaa

Escuela de Ingeniería Ambiental, Universidad Católica San Pablo, Quinta Vivanco s/n, Urb. Campiña Paisajista, Arequipa, Peru

Abstract:

Groundwater contamination by arsenic is widespread, with arsenite [As(III)] prevailing as the dominant—and most toxic—redox species. A two-stage treatment train was evaluated, comprising pre-oxidation of As(III) to arsenate [As(V)] followed by adsorption onto surface-modified activated carbon. Equilibrium sorption on the unmodified carbon was well described by the Langmuir isotherm $q_{max} = 1.99 \text{ mg g}^{-1}$, whereas the modified carbon exhibited Freundlich behavior, $K = 0.49055 \text{ L mg}^{-1}$, consistent with a heterogeneous surface. Time-dependent uptake for both sorbents was best represented by the pseudo-first-order kinetic model. Relative to direct adsorption alone, coupling oxidation with adsorption significantly enhanced overall arsenic removal, 34.68%. These results indicate that integrating an oxidation step with adsorption on modified carbon provides an efficient, environmentally benign, and cost-effective approach for treating waters containing elevated As(III)/As(V).

Keywords:

Arsenic, arsenite oxidation, modified activated carbon, adsorption isotherms, pseudo-first-order kinetics, water treatment.