



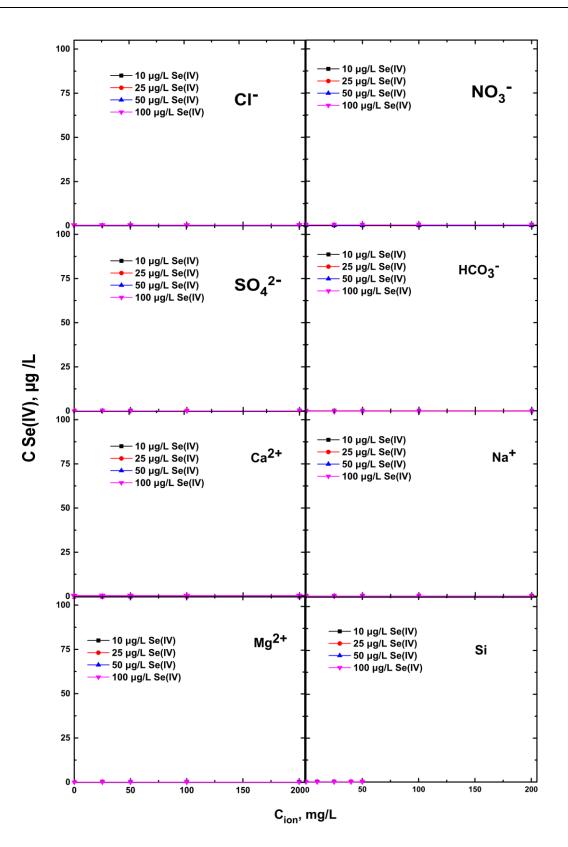
## Speciation and Determination of Selenium Oxyanions at the Drinking Water Pollution Concentration Levels

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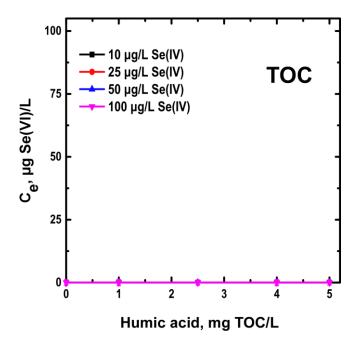
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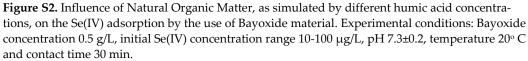
## Text S1

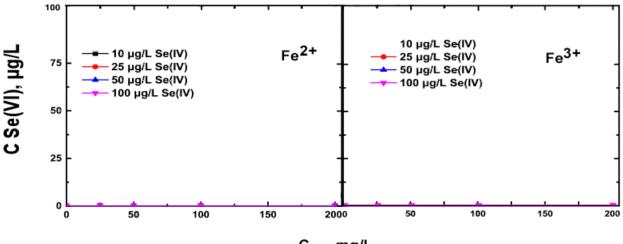
- The iron content of adsorbents was determined by the dissolution of sample in HCl and the measurement of iron by flame atomic absorption spectroscopy, using a Perkin Elmer AAnalyst 800 instrument.
- The surface area of the adsorbents was measured by nitrogen gas adsorption at liquid N<sub>2</sub> temperature (77 K), using a micropore surface area analyzer, according to the Brunauer-Emmett-Teller (BET) model.
- The determination of isoelectric point (IEP) in a water dispersion of oxy-hydroxide solid was performed by the respective curve of zeta-potential against the pH value at 20 ± 1°C. The sample initially prepared by suspending 50 mg of fine oxy-hydroxide powder into 1 L of electrolyte solution (0.01 M NaNO<sub>3</sub>). Then, several volumes of 100 mL were equilibrated at different pH values (4-10) for 60 min by adding either HNO<sub>3</sub> or NaOH solutions (1 N) under stirring. The electrophoretic velocity of at least 20 particles was subsequently reported via a digital camera, using a Rank Brothers Micro-Electrophoresis Apparatus Mk II device and the ζ-potential value was calculated accordingly.
- The surface charge density of the adsorbents and the point of zero charge (ZPC) were estimated by using the potentiometric mass titration method [1].



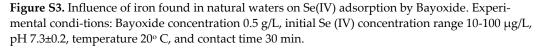
**Figure S1.** Influence of common water ions on Se(IV) residual concentration (due to adsorption) by the use of Bayoxide material. Experimental conditions: Bayoxide concentration 0.5 g/L, initial Se(IV) concentration range 10-100  $\mu$ g/L, pH 7.3±0.2, temperature 20° C and contact time 30 min.











## Supplementary reference

 Kosmulski M. Surface charging and points of zero charge, Surface Charging and Points of Zero Charge. 1<sup>st</sup> ed.; CRC Press, Boca Raton 2009. doi:10.1201/9781420051896