Removal of TcO₄⁻ using zero-valent manganese

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After the FDNPP accident, Tc-99 is recovered by anionexchange resin in ALPS from the contaminated water. Since anion-exchange resin is concerned about gas generation by radiolysis, an alternative recover of Tc-99 should be challenged for final disposal. Zero-valent metals have been widely studied for reductive remediation of toxic metals due to their suitable redox potentials[1-3]. We here studied the removal of dissolved Tc-99 in aqueous water by using zerovalent metals for the reduction of $Tc^{7+}_{(aq)}$ to $Tc^{4+}_{(s)}$. We examined the removal of ReO_4^- (as an alternative of TcO_4^-) using zero valent Mn, Fe, Co, and Cu (Mn⁰, Fe⁰, Co⁰, Cu⁰, respectively), and chemical species change of the Re after the treatment by Mn⁰.

Removal experiments of ReO₄⁻ were conducted that a 6.67 g/L of zero-valent metal was suspended in 30 mL of 0.01 mg/L ReO₄⁻ stock solution under anaerobic condition.

Removal ratios of ReO₄⁻ after the treatment of the metals are obtained ~99% by Mn⁰, ~60% by Fe⁰ ~30% by Co⁰, and ~2% by Cu⁰, indicating the highest removal of ReO₄⁻ by Mn⁰ among the metals used. Time course of the removal ratio of ReO₄⁻ by Mn⁰ showed that ReO₄⁻ was abruptly increased up to 98% within 2h at pH5.5. Re L_{III}-edge XANES spectra indicated chemical species change of ReO₄⁻ to Re³⁺ after the treatment by Mn⁰. XRD spectra of Mn⁰ after the treatment was correspond to Mn(OH)₂.

These results indicate that dominant removal mechanism of ReO_4^- by Mn^0 was the formation of insoluble Re oxides by collaborated redox reactions of Re^{7+} to Re^{3+} with Mn^0 to Mn^{2+} . Our results strongly suggest that Mn^0 is an adequate reductant for the removal of TcO_4^- from aqueous solution.

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