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Stability of chain and cyclic monoamide resins for selective separation of U(VI) against γ -ray irradiation in HNO₃

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In order to develop resins with selectivity to U(VI) in HNO₃ media, we have synthesized several polymer beads with the structure of a monoamide as the functional group. For wider applications of these resins including treatment of highly-radioactive solutions, it is necessary to investigate their stability under irradiation conditions. In this study, resins with a chain-type monoamide and a cyclic-type one as the functional group were irradiated by γ -ray in HNO₃ and adsorptivities to FP ions were examined. The structure change in the resins by irradiation was also studied.

N-methyl-N-vinylacetamide (VMAA) [1] was used the chain-type monoamide resin. For the cyclic type resin, that with a 6-membered ring of 1-(4-vinylbenzyl)piperidin-2-one (VBPP) was newly synthesized. The resins and HNO₃ of up to 6 M were mixed in a Pyrex sample tube at a ratio of 5 cm³/g, respectively, and used as the samples for γ -ray irradiation. Irradiation by the Co-60 source was performed up to 0.95 MGy at room temperature under ambient atmosphere. The irradiated resins were separated from the supernatant and washed using distilled water to avoid further degradation of the resins by HNO₃. Adsorptivities of the irradiated resins to FP ions were examined in HNO₃ by the batch method similarly to the previous studies[1-3]. The resulting resins and the supernatants were also analyzed by IR and NMR, respectively.

For irradiated VMAA, the K_d values for Pd(II) and Re(VII) were found to become lower than those of the neat ones. This indicate that the degradation properties are similar to those of chain-type monoamide extractants[4]. On the other hand, the K_d values of the irradiated VBPP for Pd(II) and Zr(IV) were found to be increased at lower concentration of HNO₃ with increasing dose. Polyvinylpyrrolidone (PVPP), which is a cyclic monoamide resin and has a 5-membered pyrrolidone ring, changes the chemical structure by γ -ray irradiation in HNO₃, where the degradation starts with the ring opening and the final products have been found to be a primary-amine-type anion exchange resin and oxalic acid[5]. This study suggests that the major route of change in the chemical structure of VBPP is basically identical to that of PVPP and that the resulting resin structures differ according to the number of carbon atoms in the cyclic monoamides.

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