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Separation of no-carrier-added ^{109}Cd from natural silver target using RTIL 1-butyl-3-methylimidazolium hexafluorophosphate

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The room temperature ionic liquid (RTIL), 1-butyl-3-methylimidazolium hexafluorophosphate $[\text{C4mim}][\text{PF6}]$ has found application in separation of a range of metal ions replacing volatile and toxic traditional organic solvents in liquid-liquid extraction (LLX) systems. Despite of disadvantage of probable release of HF in acidic reaction, the RTIL $[\text{C4mim}][\text{PF6}]$ is used widely in developing green processes in analytical chemistry as it is hydrophobic in nature, lacks vapor pressure, and can be synthesized easily. Favorable half-life of ^{109}Cd (461.4 d) allows its use in the field of medical science, environmental science as well as in technology. Several reports are also available on the production and separation of no-carrier-added (NCA) ^{109}Cd . This paper reports the separation of NCA ^{109}Cd from the natAg target using $[\text{C4mim}][\text{PF6}]$ in LLX. The ^{109}Cd was produced by bombarding a natAg foil (25.4 mg/cm² thick) by 30 MeV α -particles at the Variable Energy Cyclotron Centre, Kolkata, India. After the decay of all short-lived products, NCA ^{109}Cd was separated from the bulk Ag using $[\text{C4mim}][\text{PF6}]$ as extractant in combination with HNO_3 where ammonium pyrrolidine dithiocarbamate (APDC) was used as a complexing agent. At the optimum condition, 3 M HNO_3 , 0.2 mL 0.1 M APDC and 1.25 mL $[\text{C4mim}][\text{PF6}]$, bulk Ag was extracted to the IL phase binding with APDC, leaving $\sim 85\%$ NCA ^{109}Cd in the aqueous phase. The ionic liquid was also recovered by washing the IL phase with 5 M HNO_3 . The reported separation technique is simple, fast and in concurrence with green chemistry approach.

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