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Universal chromatographic system – paper impregnated with N,N,N',N'-Tetraoctyl Diglycolamide is an useful tool for various elements separation and may serve as validated QC method for radionuclide generators eluates and nuclides separation.

The <sup>227</sup>Ac is produced in a nuclear reactor and decays according to a scheme:

$$^{226}Ra(n,\gamma)^{227}Ra \xrightarrow{\beta^{-}}{42min} ^{227}Ac$$

$$^{227}Ac \xrightarrow{\beta^{-}}{22 y} ^{227}Th \xrightarrow{\alpha}{223}Ra \xrightarrow{\alpha}{11 d} ^{219}Rn \xrightarrow{\alpha}{4s} ^{215}Po \xrightarrow{\alpha}{211}Pb \xrightarrow{\beta^{-}}{36 m} ^{211}Bi \xrightarrow{\alpha}{207}Tl \xrightarrow{\beta^{-}}{5 m} ^{207}Pb$$

Thus <sup>227</sup>Ac may be separated from <sup>226</sup>Ra as well as <sup>223</sup>Ra from <sup>227</sup>Ac and <sup>227</sup>Th. Single 5 × 20 cm DGA-sheet was cut of 20 x20 cm bulk sheet and used for the separation of *AEM*. Mobile phase was the 1M HNO<sub>3</sub>. Developed chromatographic sheet was measurement on a standard radio-TLC scanner (see **Fig.1**). Further, the sheet was re-measured repeatedly during one hour in the same position to allow to decay/growth the <sup>211</sup>Pb in the respective peaks of <sup>211</sup>Pb and <sup>223</sup>Ra. Finally the sheet was measured after the decay of <sup>227</sup>Th and <sup>223</sup>Ra in their respective peaks (one year after the separation) and so at the <sup>227</sup>Ac peak site the ingrowth of <sup>227</sup>Th and <sup>223</sup>Ra allowed its detection. In pure <sup>223</sup>Ra solution, no activity should remain in first half of the chromatogram.

**Fig. 1.** Separation of *AEM* using 5 × 20 cm DGA-sheet with 1M HNO<sub>3</sub> as a mobile phase. <sup>227</sup>Th remains on start, <sup>227</sup>Ac has a retention factor  $\approx$  0.2, <sup>211</sup>Pb  $\approx$  0.7 and <sup>223</sup>Ra  $\approx$  0.9.



**Fig. 2**. Left: γ-spectrum of peak 2. (<sup>227</sup>Ac in equilibrium with daughter nuclei); Right: Decay/ingrowth of <sup>211</sup>Pb in peaks 3./4.



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