Activation cross sections of α -particle-induced reactions on scandium in the energy range of 22–51 MeV[†]

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Titanium radioisotopes have important applications in medical diagnostics. The positron emitter $^{45}\mathrm{Ti}~(T_{1/2}=184.8~\mathrm{min})$ is utilized in positron emission tomography (PET). It can be produced in carrier-free form through charged-particle-induced reactions on calcium or scandium. Scandium, a monoisotopic element ($^{45}\mathrm{Sc}$: 100%), may offer a distinct advantage for producing $^{45}\mathrm{Ti}$.

In this study, we focused on α -particle-induced reactions on scandium. A literature survey using the EXFOR database identified six experimental studies of α -particle-induced reactions on scandium. However, the available data are limited and inconsistent, with no prior studies addressing the $^{45}\mathrm{Sc}(\alpha,x)^{45}\mathrm{Ti}$ reaction. Therefore, we conducted a new experiment to provide reliable cross sections in the energy range of 22–51 MeV.

The experiment was performed at the AVF cyclotron at RIKEN using stacked-foil activation and high-resolution γ -ray spectrometry to determine the production cross sections.

The stacked target comprised pure metallic foils of 45 Sc (99.9% purity, 7.71 mg/cm²), nat Ti (99.5% purity, 9.13 mg/cm²), and 27 Al (>99% purity, 4.99 mg/cm²). Nine sets of Sc-Al-Ti-Al foils were arranged in a target holder that also functioned as a Faraday cup. The target stack was irradiated for 30 min with a 50.9-MeV α -particle beam at an average intensity of 201 electric nA (enA). The incident beam energy was determined using the time-of-flight method, $^{2)}$ while the SRIM code³⁾ was used to estimate energy degradation and uncertainty propagation through the stacked foils. The beam intensity was determined from the charge collected by the Faraday cup.

The γ -ray spectra of each irradiated Sc foil and its corresponding Al catcher foil were recorded using a high-resolution HPGe detector (ORTEC GMX30P4-70, 30% relative efficiency). Five measurements were performed per foil to accommodate the different half-lives of the products. Nuclear reaction and decay data for γ -ray spectrometry were obtained from the online databases NuDat 3.0^4) and LiveChart. ⁵

The $^{nat}\mathrm{Ti}(\alpha,x)^{51}\mathrm{Cr}$ monitor reaction was used to assess and correct the primary beam parameters and target thicknesses. The derived cross sections for this

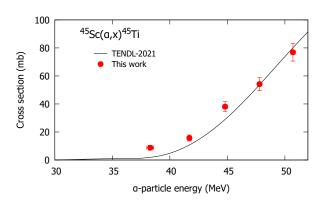


Fig. 1. Cross sections of ${}^{45}\mathrm{Sc}(\alpha,x){}^{45}\mathrm{Ti}$ reaction.

reaction were compared with the IAEA-recommended values. To improve agreement with the reference values, corrections were applied to the measured parameters: the beam energy was adjusted by +0.2 MeV, the beam intensity by -2%, and foil thicknesses by -2% (Sc), -1% (Ti), and -1% (Al) within their respective uncertainties. The corrected parameters, beam energy (51.1 MeV), beam intensity (197 enA), and thicknesses (7.56 mg/cm² for Sc, 9.04 mg/cm² for Ti, and 4.94 mg/cm² for Al) were used for cross-section analysis.

Production cross sections of 45 Ti ($T_{1/2}=184.8$ min) were determined using 511-keV annihilation γ rays. In addition to direct formation, contribution by the indirect route through the decay of its parent nuclei, 45 V ($T_{1/2}=547$ ms), is possible. 45 V has decayed completely before the γ -ray measurements. Because the 511-keV γ rays are also emitted by co-produced, longer-lived positron emitters such as 48 V ($T_{1/2}=15.974$ d), 44g Sc ($T_{1/2}=3.97$ h), and 43 Sc ($T_{1/2}=3.891$ h), their contributions to the photopeak were subtracted to obtain net counts from 45 Ti.

The cumulative cross sections, derived from the net counts in the spectra, are presented in Fig. 1 along-side TENDL-2021 predictions.⁶⁾ No prior experimental data on this reaction have been found in the literature. The theoretical values show good agreement with our experimental results.

Production cross sections were also determined for ^{48}V , $^{47,\,46m+g,\,44m,\,44g,\,43g}\text{Sc}$, and $^{43,\,42}\text{K}$ in this work. The results are expected to contribute to the further development of nuclear medicine.

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