

Activation cross sections of alpha-induced reactions on natural tungsten for ^{186}Re and ^{188}Re production

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Radioisotopes (RI) are used for diagnosis and therapy in nuclear medicine. ^{186}Re is a β emitter with a half-life of 3.72 days, a maximum β energy of 1.07 MeV, average penetration ranges of 1.1 mm in soft tissue and 0.5 mm in bone, and a 9.47% γ -ray emission at 137 keV. ^{188}Re is a β emitter with a half-life of 17 hours, maximum β energy of 2.12 MeV, and 15.61% γ -ray emission at 155 keV.^{1,2} Both isotopes can be used for theranostics (therapy and diagnosis).

We focused on a process to produce $^{186,188}\text{Re}$ through alpha-induced reactions of natural tungsten because we could find data for only one 43 MeV.³ Therefore, we measured the excitation function of the $^{\text{nat}}\text{W}(\alpha, x)^{186,188}\text{Re}$ reactions up to 51 MeV.

The excitation functions of the $^{\text{nat}}\text{W}(\alpha, x)^{186,188}\text{Re}$ reaction were measured by the stacked-foil method, activation method and high-resolution γ -ray spectroscopy. $^{\text{nat}}\text{W}$ foils (purity: 99%, Goodfellow Co., Ltd., UK) were stacked with $^{\text{nat}}\text{Ti}$ foils (purity: 99%, Goodfellow Co., Ltd., UK) for monitoring the beam parameters and degrading the beam energy. The thicknesses of the W and Ti foils were 15.03 and 2.23 mg/cm², respectively.

The irradiation was performed at the RIKEN AVF cyclotron. A 51 MeV alpha beam with an average intensity of 209.7 pA was irradiated on the target for 2 h. The incident beam energy was measured by the time-of-flight method using plastic scintillator monitors.⁴ The beam energy degraded in the stacked target was calculated using the SRIM code available online.⁵ The γ -ray spectra of the activated foils were measured by an HPGe detector. Nuclear decay data were taken from the online NuDat 2.7 database.⁶

From the net peak areas of the 137.16- and 155.04-keV γ -rays, the activation cross sections for the $^{\text{nat}}\text{W}(\alpha, x)^{186,188}\text{Re}$ reaction were deduced using the standard activation formula

$$\sigma = \frac{T_{\gamma} \lambda}{\varepsilon_d \varepsilon_{\gamma} \varepsilon_t N_t N_b (1 - e^{-\lambda t_b}) e^{-\lambda t_c} (1 - e^{-\lambda t_m})}$$

where N_t denotes the surface density of target atoms; N_b the number of bombarding particles per unit time; T_{γ} the number of counts in the photo-peak; ε_d the detector efficiency; ε_{γ} the γ -ray abundances; ε_t the

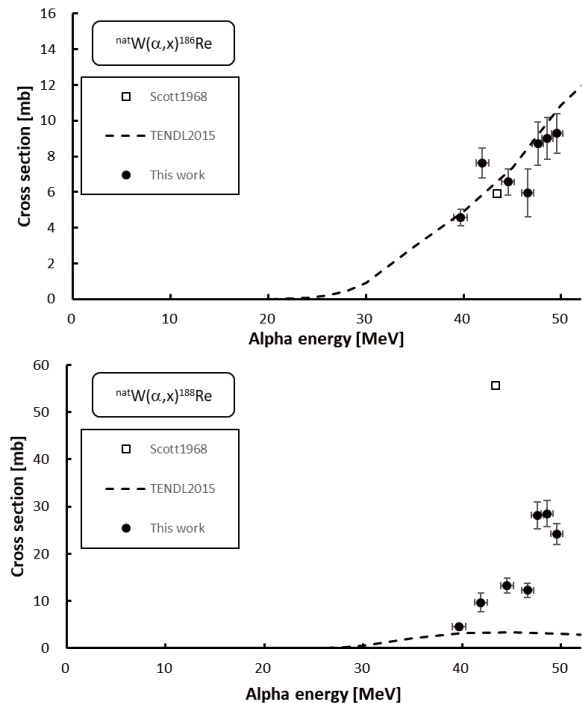


Fig. 1. Excitation functions of the $^{\text{nat}}\text{W}(\alpha, x)^{186,188}\text{Re}$ reactions. The result is compared with a previous study³⁾ and TENDL-2015.⁷⁾

measurement dead time, which is the ratio of live time to real time; λ the decay constant; t_b the bombarding time; t_c the cooling time; and t_m the acquisition time.

We found that our $^{\text{nat}}\text{W}(\alpha, x)^{186}\text{Re}$ result is in good agreement with previous data obtained by NE. Scott *et al.*³⁾ and the theoretical calculation (TENDL-2015).⁷⁾

On the other hand, the $^{\text{nat}}\text{W}(\alpha, x)^{188}\text{Re}$ result shows disagreements with the other data. TENDL-2015⁷⁾ underestimates the cross section at all energies.

References

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