Correlated measurement of mass and decay of fusion evaporation products for ${}^{51}V+{}^{159}Tb$ reactions via MRTOF+ α -TOF detector

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The SHE-Mass-II facility¹⁾ is a system with a multireflection time-of-flight mass spectrograph (MRTOF- MS^{2}) coupled with the gas-filled recoil ion separator GARIS-II³⁾ in the E6 experimental room for the mass measurement of fusion evaporation products, such as very low yield nuclei in the superheavy region.

Recently, we developed and installed a novel detector named α -TOF⁴⁾, which simultaneously records the time-of-flight (TOF) signal and successive α -decay. The α -TOF detector has the capability to significantly reduce the background level.

The experiment was performed using ${}^{51}V+{}^{159}Tb$ reactions. A ${}^{51}V$ beam was accelerated up to 6.0 MeV/nucleon by RRC. The beam energy on the target was reduced by an aluminum degrader to approximately 4.8 MeV/nucleon. A 460- μ g/cm²-thick ${}^{159}Tb$



Fig. 1. (a) Singles TOF spectrum. (b) TOF spectrum in coincidence with 207 Ra α -decays obtained using a time gate of 5s.

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Fig. 2. Two-dimensional mapping of TOF versus correlated α -decay energies.

target was prepared by a sputtering method on a 3.0 μm Ti backing foil.

The fusion evaporation residues (ERs) were separated from the primary beam and efficiently transported using GARIS-II. After decelerating ERs using a Mylar foil, the ERs were captured in a cryogenic high-purity He gas catcher, and the thermalized ions were extracted by an RF-carpet and transported to the MRTOF-MS via multiple RF ion traps. We observed the $^{206, 207}$ Fr, $^{206, 207}$ Ra and 204 Rn isotopes extracted as doubly charged ions. In this measurement, we focused on $^{206, 207}$ Ra isotopes having relatively short half-lives, *i.e.*, $T_{1/2} = 240$ ms for 206 Ra and $T_{1/2} = 1.38$ s for 207 Ra. Figure 1 shows a part of the TOF spectrum at A =

Figure 1 shows a part of the TOF spectrum at A = 207. When we gated the ToF spectrum by the α -ray energy of 207 Ra with a coincident time of 5 s (3.5 half-lives), we clearly discriminated the decay-correlated TOF events (Fig. 1(b)).

Figure 2 shows a part of the two-dimensional spectrum for TOF and the correlated α -decay energies. From the correlation mapping, we obtained the information of TOF, as well as decay properties such as α -energy and decay time, atom by atom. Further analysis is in progress.

References

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