

Determination of half-life of ^{89}Rh by measuring in-flight decay

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The in-flight decay of ^{89}Rh was measured to deduce its half-life using two-fold particle-identification (PID) sections, the second stage of the BigRIPS separator and the ZeroDegree spectrometer. Moreover, its production cross section was deduced.

Inconsistent results were observed for the half-life of ^{89}Rh . One was $\gtrsim 1.5 \mu\text{s}$ estimated from Ref. 1). In this experiment at the LISE spectrometer in GANIL, ^{89}Rh was produced using a $^{112}\text{Sn} + \text{natNi}$ reaction and its yield was consistent with the yield systematics of the neighboring RIs. Thus, its half-life was considered to be comparable to or longer than the time of flight (TOF) in the spectrometer ($\sim 1.5 \mu\text{s}$). The other was $< 120 \text{ ns}$, investigated at RIBF, RIKEN.²⁾ Using a $^{124}\text{Xe} + \text{Be}$ reaction, no events of ^{89}Rh were obtained at F11, the final focal plane of the ZeroDegree spectrometer. The production yield at the F0 target was estimated from the yield systematics, and in-flight decay during the flight from F0 to F11 was assumed. Subsequently, the upper limit of the half-life was deduced to be 120 ns. Thus, direct measurement of the decay was required to determine the half-life.

We measured the in-flight decay of ^{89}Rh at RIBF. The proton-rich RI beams including ^{89}Rh were produced from a 120-particle nA 345-MeV/nucleon ^{124}Xe beam impinging on a 4-mm Be target. The fully-stripped ^{89}Rh was identified³⁾ in the second stage of the BigRIPS. With a 4.9 hours primary-beam dose, 17 events were obtained as shown in Fig. 1. They were re-identified in the following spectrometer, ZeroDegree, whether they maintained ^{89}Rh until F11 or decayed to ^{88}Ru by proton emissions. The PID method in the ZeroDegree spectrometer was basically the same with the one in the second stage. In the PID, the mass-to-charge ratios (A/q) could not be deduced for the events that ^{89}Rh decayed to ^{88}Ru , because their magnetic-rigidity values changed before and after the decays. In contrast, the atomic number Z at F11 was properly deduced to be 44, because the velocity almost unchanged by the decay. Figure 2 shows the Z spectra of the all events and the ^{89}Rh events at F7. Among 17 events of ^{89}Rh at F7, eight events were located around $Z = 44$, suggesting their decays to ^{88}Ru . The remaining nine events were ^{89}Rh until F11. From the numbers of ^{89}Rh nuclei at F7 and F11 and the γ -corrected TOF of 258 ns, its half-life was determined to be $281 \pm 101 \text{ ns}$ (preliminary).

The production cross section of ^{89}Rh was deduced from its half-life, γ -corrected TOF from F0 to F7, the

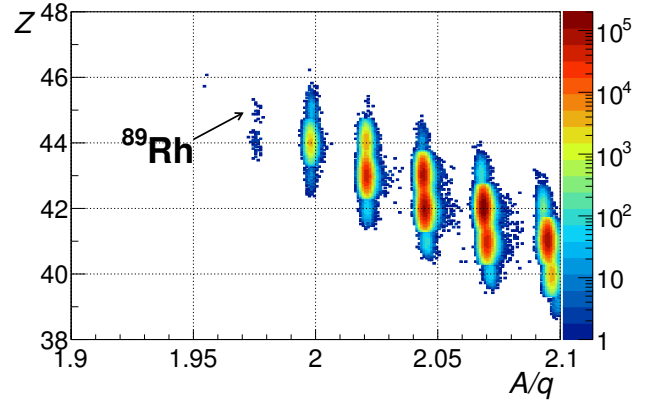


Fig. 1. (Preliminary) Z versus A/q PID plots in the second stage of the BigRIPS for ^{89}Rh produced by the reaction of $^{124}\text{Xe} + \text{Be}$ (4 mm) at 345 MeV/nucleon. Seventeen events of ^{89}Rh were obtained.

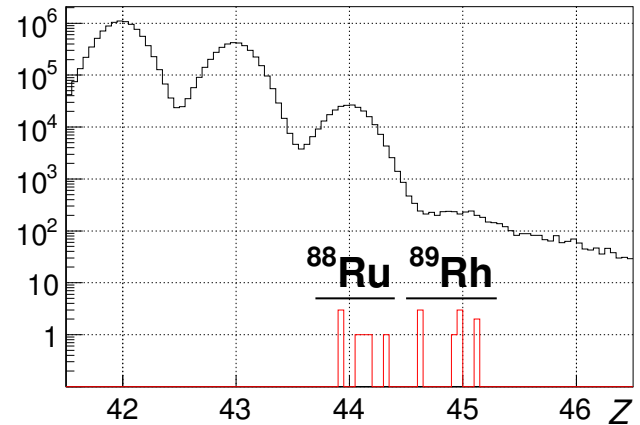


Fig. 2. (Preliminary) Z spectra in the ZeroDegree spectrometer for all events (black) and ^{89}Rh events at F7 (red). Till F11, eight nuclei of ^{89}Rh decayed to ^{88}Ru .

observed events at F7, and the transmission in the BigRIPS. It was $(3.81^{+2.46}_{-0.80}) \times 10^{-10} \text{ mb}$ (preliminary), and was $\sim 1/20$ of expected cross section from the systematics of the other isotopes with $N = Z - 1$. This small cross section might cause the inconsistent short half-life estimated in the Ref. 2).

More elaborate analysis is now in progress.

References

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