

Commissioning of helium gas-cell + MRTOF-MS system for studies of multi-nucleon transfer reactions

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The multi-nucleon transfer (MNT) reactions have recently garnered interest for producing neutron-rich exotic isotopes that cannot be reached by conventional methods, for example, fragmentation, in-flight fission, and complete fusion. We recently proposed a plan to probe the existence of the $N = 152$ sub-shell closure in the lighter actinide region by using the MNT reactions with a ^{248}Cm target. However, there are technical difficulties peculiar to the MNT reactions; their products have wide energy and angular distributions. To solve this, we proposed the upgradation of the experimental setup composed of a cryogenic helium gas cell (GC) and a multi-reflection time-of-flight mass spectrograph (MRTOF-MS) used for the mass measurements of super-heavy nuclei.¹⁾

The setup is located downstream of the focal plane of the gas-filled recoil separator GARIS-II; the schematic view of the setup is shown in Fig. 1. The GARIS-II is used in vacuum mode to transport the primary beam to the target. The isotopes of interest are produced via

MNT reactions by irradiating the primary beam on a fixed target, which is mounted in a target cassette directly before the GC. A Ti degrader is placed immediately after the target to adjust the reaction products' stopping ranges inside the GC. A beam dump is also set on the cassette exit to stop the unreacted primary beam from entering the GC. A small beam spot is required to collect the MNT reaction products efficiently, and some additional primary beam adjustment equipment is installed in the GARIS-II focal plane chamber to this end.

The first commissioning experiment was performed with the MNT reaction of $^{136}\text{Xe} + ^{\text{nat}}\text{Pb}$. A 1.6 mg/cm^2 $^{\text{nat}}\text{Pb}$ target was prepared through evaporation on a $3 \mu\text{m}$ Ti backing. The energy of the ^{136}Xe primary beam was 6.5 MeV/nucleon on the target. The doubly charged $A = 209$ and 207 isobar series were observed in the experiment. The counting rate of $^{209}\text{Bi}^{2+}$ was 0.21 cps with a primary beam intensity of 0.9 particle nA at the FC. This is approximately 3 times lower than the predictions from the GRAZING code.²⁾ The experimental value may be explained by the fraction of the primary beam lost owing to the second slit between the FC and the target. Figure 2 shows the GC extraction yield of the doubly charged $A = 207$ isobar series as a function of the primary beam intensity. It shows linearity up to at least 1 particle nA at the FC. These results prove the proposed setup is suitable for the planned study with the MNT reactions.

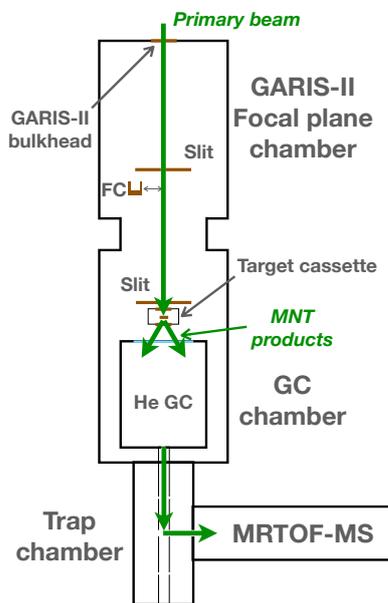


Fig. 1. Schematic view of the experimental setup.

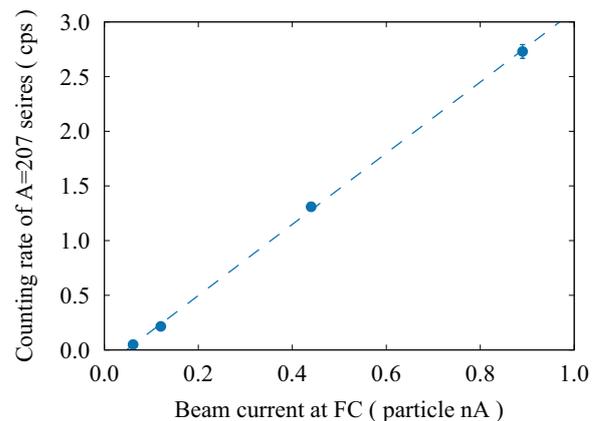


Fig. 2. Gas cell extraction efficiency as a function of primary beam current. The dashed line is a guide for the eye.

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