

Direct mass measurement for ^{103}Sn and its impact on rp -process endpoint SnSbTe cycle

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The nuclide ^{103}Sn was produced by projectile fragmentation of ^{112}Sn at a energy of 400 MeV/u, and its mass was directly measured by a storage-ring based isochronous mass spectrometry at HIRFL-CSR facility in Lanzhou, China. The new mass value deviated the literature one, which was indirectly determined from β -decay spectrum of ^{103}In , by about 2.5σ , and the mass precision was improved by a factor of two.

Since the decay energies of proton or α particle were precisely measured in the decay chain of $^{112}\text{Cs}(p)^{111}\text{Xe}(\alpha)^{107}\text{Te}(\alpha)^{103}\text{Sn}$, masses of other three nuclides will also shift following the change of ^{103}Sn mass. The mass surfure of the four isotopic chain seem to be more smooth according to the systematic behavior of two-neutron separation energies.

Astrophysical network calculations indicate that the rp process in x-ray bursts ends at a SnSbTe cycle and cannot proceed past tellurium isotopes due to the low α separation energies of neutron-deficient tellurium isotopes. The main reaction flow of this cycle is $^{103}\text{Sn}(\beta+)^{103}\text{In}(p, \gamma)^{104}\text{Sn}(\beta+)^{104}\text{In}(p, \gamma)^{105}\text{Sn}(p, \gamma)^{106}\text{Sb}(p, \gamma)^{107}\text{Te}(\gamma, \alpha)^{103}\text{Sn}$. The change of ^{107}Te mass will influence on the competition between $^{106}\text{Sb}(p, \gamma)^{107}\text{Te}$ and $^{106}\text{Sb}(\beta+)^{106}\text{Sn}$, and thus determine the amount of the SnSbTe cycle. However, since the reaction rate of $^{106}\text{Sb}(p, \gamma)^{107}\text{Te}$ is about 3 orders of magnitude stronger than that of $^{106}\text{Sb}(\beta+)^{106}\text{Sn}$. Our new mass will not change amount of the SnSbTe cycle too much.