

⁵³Fe

In “Radioactive Isotopes of Iron”, Livingood and Seaborg reported the production of ⁵³Fe in 1938 ([1938Li06](#)). The isotope was produced in the reaction ⁵⁰Cr(α ,n)⁵³Fe with 16 MeV α -particles accelerated by the Berkeley cyclotron. The decay curves of the produced radioactivity were measured with a quartz fiber electroscope following chemical separation. The authors “believe the 9-minute activity to be due to Fe⁵³ rather than to Fe⁵⁵ because: (1) it is not produced by deuteron or slow neutron bombardment of Fe, (2) it is produced by fast neutrons on Fe, (3) attempts to produce ⁵⁵Fe by other reactions have not disclosed a 9 minute activity.” The half-life was determined to be 8.9(2) m. In 1937, Ridenour and Henderson had observed a 9-minute activity; however, they were unable to make the unique mass assignment and attributed it to either the reaction ⁵⁰Cr(α ,n)⁵³Fe or the reaction ⁵²Cr(α ,n)⁵⁵Fe ([1937Ri01](#)). In an even earlier publication, they had preferred the later assignment ([1937He06](#)).

Adapted from reference ([2010Sc18](#))

- [1937He06](#) W. J. Henderson and L. N. Ridenour, Phys. Rev. **52**, 40 (1937).
[1937Ri01](#) L. N. Ridenour and W. J. Henderson, Phys. Rev. **52**, 889 (1937).
[1938Li06](#) J. J. Livingood and G. T. Seaborg, Phys. Rev. **54**, 51 (1938).
[2010Sc18](#) A. Schuh, A. Fritsch, M. Heim, A. Shore, and M. Thoennessen, At. Data Nucl. Data Tables **96**, 817 (2010).

Please cite this abstract as: “FRIB Nuclear Data Group, *Discovery of Nuclides Project*, Isotope Database, doi:[10.11578/frib/2279152](https://doi.org/10.11578/frib/2279152)”