

FULLY AUTOMATED PRODUCTION OF Zr-89 USING IBA NIRTA AND PINCTADA SYSTEMS

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Background

⁸⁹Zr

⁸⁹Zr ($t_{1/2} = 3.3d$) has a near ideal half-life for antibody-based imaging, and the low energy of its positron (395.5keV) results in PET images with high spatial resolution. The most popular reaction to produce ⁸⁹Zr is the ⁸⁹Y(p,n)⁸⁹Zr nuclear reaction. Yttrium targets could be either a foil, sputtered onto a support or Y₂O₃ pellets.

The motivation of the present work was the fully automated production of ⁸⁹Zr using commercially available automated systems. We also describe a newly designed and tested platinum cradle, capable of holding a metallic foil and being directly transferable/compatible between the IBA NIRTA target and IBA Pinctada Metal dissolution/purification module.

Materials & Method

The commercially available Nirta Solid Target from IBA was coupled to our 18/9 IBA cyclotron using a 2-meter external beam line. A fully automated pneumatic solid target transfer system (STTS) designed by TEMA Sinergie was used to deliver the irradiated targets to a dedicated hotcell. The newly designed platinum cradle holding the yttrium foil (8mm \varnothing foils and 0.1mm to 0.250mm possible thickness) is shown in image 1.

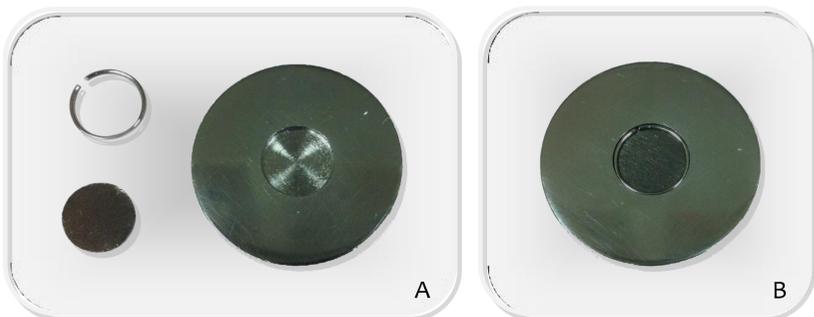


Image 1: (A) Disassembled Pt target foil cradle, Y-foil and Pt containment ring. (B) Assembled target ready for irradiation.

Typical irradiation parameters were 14.9MeV at 20-30 μ A for 1.5-3.0 hours (90° angle of incidence). No visible damage or deformation of the Y-foil was observed at these beam intensities. The irradiated

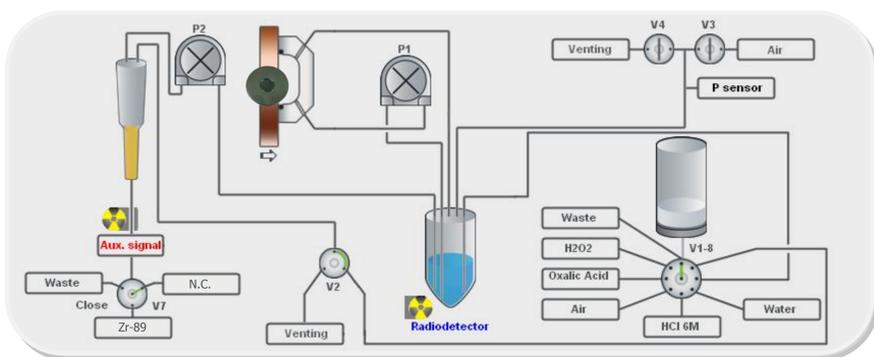


Image 2: Schematic of IBA Pinctada Metal dissolution/purification module setup for ⁸⁹Zr production.



cradle, containing the ⁸⁹Zr target was then loaded directly into the IBA Pinctada Metal module (1st Gen., see image 2) for dissolution/purification without disassembly (foil dissolved on Pt cradle).

We used the dissolution/purification method described by Holland et al. 2009 without modification.

Image 3: Bio-Rad 10mL column filled with 100mg hydroximate resin (insoluble black material visible on top of resin after use).

Results

Image 4 shows the radioactive elution profile from the hydroximate resin column. ⁸⁹Zr was eluted from the resin and collected in 1.0-1.2mL 1.0M oxalic acid at a flow rate of 0.7mL/min (start and duration of ⁸⁹Zr collection determined by radioactive profile rather than fixed elution volumes).

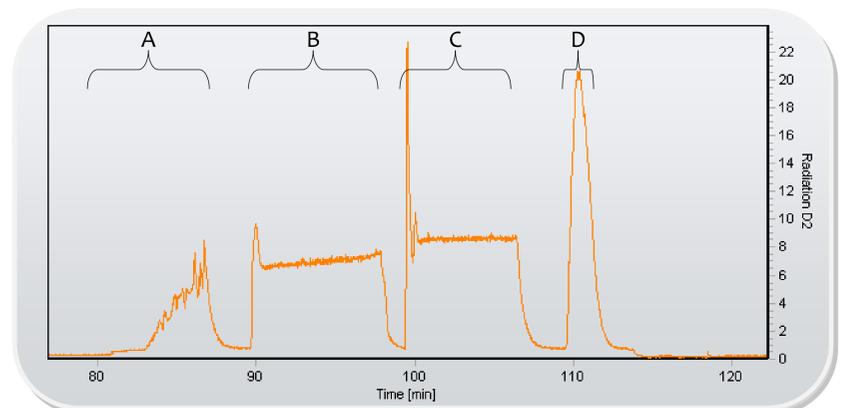


Image 4: Radioactive elution profile from hydroximate resin column. A. Loading dissolved foil (in 2mL 6M HCl + 5mL H₂O) onto resin, B. Washing resin with 10mL 2M HCl, C. Washing resin with 10mL H₂O, D. Elution of ⁸⁹Zr with 1.0M oxalic acid in 1.0-1.2mL final volume.

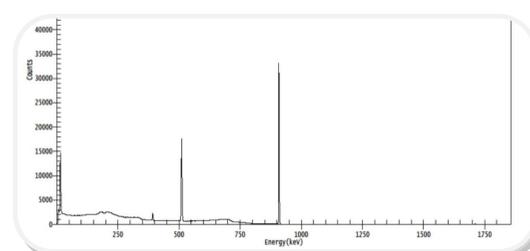


Image 5: Spectrum of γ -ray emissions from a purified ⁸⁹Zr sample 2hrs after EOB

Since yttrium has one stable isotope only, relatively pure ⁸⁹Zr is produced at low energy (14.9MeV).

Table 1 lists current production yields with various Y-foil thicknesses used.

Y-foil thickness	Irradiation	Zr-89 Product EOS MBq (mCi)	D.C. EOB Yield MBq (mCi) / μ Ah
8mm \varnothing			
0.127mm	20 μ A 1.5h	381 (10.3)	13.0 (0.35)
0.254mm (2x0.127)	25 μ A 2.0h	1110 (30.0)	22.6 (0.61)
0.127mm	28 μ A 1.4h	451 (12.2)	11.5 (0.31)
0.250mm	30 μ A 3.0h	1980 (53.5)	22.6 (0.61)

Table 1: Production summary

Conclusion

In these preliminary productions, average purified ⁸⁹Zr yield for 0.127mm thick Y-foils was 12.2MBq(0.33mCi)/ μ Ah and 22.6MBq (0.61mCi)/ μ Ah for 0.250mm thickness was achieved, in comparison to an average of 52.7MBq(1.43mCi)/ μ Ah reported by Holland et al. 2009 (at 10° angle of incidence, 0.1mm thick foil).

Acknowledgements: The funding provided for the construction of our solid targetry facility is an ANSTO/Austin/Ludwig partnership

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