

# Mass Production of $^{64}\text{Cu}$ with $^{64}\text{Ni}(p,n)^{64}\text{Cu}$ Nuclear Reaction

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## Introduction

$^{64}\text{Cu}$  ( $T_{1/2} = 12.7\text{h}$ ,  $\beta^-$  decay: 40%,  $\beta^+$  decay: 19%, E.C. decay: 41%) is one of the most useful radioisotope in nuclear medicine due to its multiple decay mode and the intermediate half-life. Several nuclear reactions, i.e.,  $^{64}\text{Ni}(p,n)^{64}\text{Ni}$ ,  $^{68}\text{Zn}(p,\alpha n)^{64}\text{Cu}$  and  $^{64}\text{Ni}(d,2n)^{64}\text{Cu}$  have been investigated for  $^{64}\text{Cu}$  production [1,2]. The highest production yield could be obtained with proton irradiation on the enriched  $^{64}\text{Ni}$  target. Therefore for mass and routine production, the  $^{64}\text{Ni}$  target fabrication by using electroplating [3], the reliable chemical separation of  $^{64}\text{Cu}$  from the irradiated  $^{64}\text{Ni}$  target and the effective recovery process for the recycling of very expensive enriched material ( $^{64}\text{Ni}$  enrichment : 96%, \$20,000/g) and so on are absolutely necessary to be established. In this work, we report our mass production method of  $^{64}\text{Cu}$  with enriched  $^{64}\text{Ni}$  and Cyclone-30 accelerator.

## Methods

$^{64}\text{Cu}$  was produced with high current cyclotron via  $^{64}\text{Ni}(p,n)^{64}\text{Cu}$  nuclear reaction at 200 $\mu\text{A}$ , 18MeV proton beam. Nickel target was prepared by electro-plating of enriched  $^{64}\text{Ni}$  (25% of enrichment) on Au coated Cu cooling plate. After proton beam irradiation, Ni target was dissolved with circulation of 50ml of 5N HCl on the dissolving device (home made) and 90°C heating. Water was added to  $^{64}\text{Ni}$  solution to dilute the normality of hydrochloric acid to 0.5N. Radiochemical separation of  $^{64}\text{Cu}$  from Ni target solution was performed with 0.01% dithizone in  $\text{CCl}_4$  solvent extraction and back extraction with 7N HCl [4]. Purification of back extracted  $^{64}\text{Cu}$  solution was carried out with AG1-x8 (Bio-Rad) anion exchange resin. For  $^{64}\text{Ni}$  recycling,  $^{64}\text{Ni}$  from the aqueous phase of solvent extraction and the electrolyte of electroplating was recovered by using AG1-x8 anion and AG50w-x8 (Bio-Rad) cation resin [5].

## Results

With the electroplating cell designed by ourselves and the electrolyte, consisting of 1.5g  $^{64}\text{Ni}$  (25% enrichment), 1.0g boric acid and 2.0g NaCl in 90ml distilled water, the smooth and uniformed Ni target (thickness : > 50mg/cm<sup>2</sup>, area: 1 x 10cm<sup>2</sup>) was obtained with applying 200mA of constant current on the cathode for 5hrs. The cathode current efficiency was about 50%. There was no damage on Ni surface during more than 200 $\mu\text{A}$  proton beam irradiation. The chemical separation yield of  $^{64}\text{Cu}$  with solvent extraction and anion exchange resin was more than 90% and the radionuclidic purity was more than 99% 1 day after bombardment. The  $^{64}\text{Ni}$  recovery yield was quantitative and measured with  $^{57}\text{Ni}$  activity produced with  $^{58}\text{Ni}(p,2p)^{57}\text{Ni}$  nuclear reaction and AA spectroscopy.

## Conclusion

$^{64}\text{Cu}$  production yield was about 9mCi/ $\mu\text{Ah}$  corrected on 96% enrichment at EOB with  $^{64}\text{Ni}(p,n)^{64}\text{Cu}$  nuclear reaction and Cyclone-30. The chemical separation yield and the radionuclidic purity of the final  $^{64}\text{Cu}$  solution was more than 90% and 99%, respectively. The  $^{64}\text{Ni}$  recovery yield performed with ion exchange resin was more than 98%.

## References

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