Production of ¹²⁴I and ⁶⁴Cu on an 18/9 MeV cyclotron as a starting material for radiopharmaceuticals preparation.





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Iodine-124 ($T_{1/2} = 4.18$ d) and Cu-64 ($T_{1/2} = 12.7$ hrs) are two very important radionuclides for radiopharmaceuticals production in preclinical research using positron emission tomography (PET). The method for producing ¹²⁴I was based on a dry distillation technique of ¹²⁴I from a solid [¹²⁴Te]TeO₂ target. The platinum target disk was used as a base for TeO₂ melt and irradiated on COSTIS target station installed on the solid [-16] target. The plantinin target uses was used as a base for the Po₂ linet and irradiated of COS1 target station instance on the end of the external beam line of the IBA Cyclone 18/9 cyclotron. The target station was equipped with a 25 μ m aluminum or 250 μ m Nb window foil in front of the target, which results in a final beam energy of 17.7 or 13.5 MeV respective. The TERIMO module was placed in a GMP class "D" research laboratory in a suitable hot cell. Isotope ⁶⁴Cu (I₂₊= 17.6 %, E₂₊max=653 keV, I₂= 38.5 %, E₂-max=579.4 keV) is a positron emitters and with half-life 12.7 hrs. The ⁶⁴Ni(p,n)⁶⁴Cu reaction route is used for ⁶⁴Cu preparation because its entrance channel is accessible at low energies and a yield of the reaction is quite high. Disadvantage of the reaction used is a high price of the enriched ⁶⁴Ni. The gold or platinum target were used for a thick ⁶⁴Ni target preparation by electro deposition. Since the external beam line of the cyclotron has no beam diagnostic devices, several aluminum plates were irradiated in the COSTIS target station with a 5 µA proton beam for 5 min with different settings for the beam focusing quadrupole magnets. After 15 minutes decay time the plates were scanned by a TLC scanner along the horizontal and vertical central axes of the plates in order to visualize the beam shape. The settings providing the most homogeneous beam spot on the target were selected and used further for the actual target irradiations. The radionuclidic purity of the product was determined by γ-spectrometry.

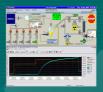




Irradiated 124TeO₂ target



GMP class "D" research laboratory in a hot cell

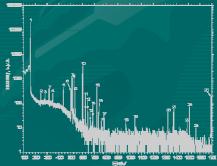


Terimo - scheme from the computer screen

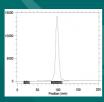
y-lines of the spectra with their energy and intensity

			-				
Peak	Nuclide	E, keV	Intensity, %	Peak	Nuclide	E, keV	Intensity, %
1	¹²³ I	158.97	83.3	14	123I	687.95	0.0267
2	123 I	247.96	0.071	15	124I	722.78	10.35
3	123 I	281.03	0.079	16	123I	735.78	0.062
4	123I	346.35	0.126	17	¹²³ I	783.59	0.059
5	123 I	440.02	0.428	18	124I	968.22	0.435
6	123 I	505.33	0.316	19	124I	1045.0	0.441
7	¹²⁴ I (annih.)	511.0	46.0	20	124I	1325.50	1.561
8	123 I	528.96	1.39	21	¹²⁴ I	1376.0	1.75
9	123 I	538.54	0.382	22	¹²⁴ I	1488.9	0.199
10	¹²⁴ I	602.72	62.9	23	¹²⁴ I	1509.49	3.13
11	123I	624.57	0.083	24	124I	1559.8	0.165
12	¹²⁴ I	645.82	0.988	25	¹²⁴ I	1691.02	10.88
13	¹²⁴ I	662.4	0.056				

y-spectra of the 124 product at EOS

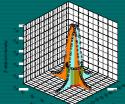


Paper chromatography 124I- Rf=0.783



Yield of 124I in the EOB time

Energy [MeV]	¹²⁴ l Yield EOB [MBq/μAh]	Yield ratio of 123 / 124			
17.7→13.5	4,99	29,2			
13.5→10.5	13,9	6,4			





The electrochemical cell



onto gold disk





EDX of electrodeposited Ni

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