

FEASIBILITY STUDY FOR PRODUCTION OF IODINE-131 USING DIOXIDE OF TELLURIUM-130

ABDESSAMAD DIDI*, AHMED DADOUCH, HASSANE EL BEKKOURI

Laboratory of Integration System and Technology Advanced (LISTA), Department of Physics, Faculty of Science Dhar Mahraz, University of Sidi Mohamed Ben Abdellah Fez Morocco
Email: abdessamad.didi1@usmba.ac.ma

Received: 21 Jun 2016 Revised and Accepted: 12 Aug 2016

ABSTRACT

Objective: Currently, nuclear medicine is becoming increasingly important, through the discovery of several medical radioisotopes, which are used in diagnosis, treatment, and medical imaging. Among the most important radionuclide which is commonly used is iodine-131, with a half-life of 8.02 d. Iodine-131 is one of the mainly essential elements in nuclear medicine. Since their first use, several studies have been conducted to meet the world need of hospital specialists in nuclear medicine. The purpose of this study was to participate in a lawsuit about the feasibility of producing ^{131}I .

Methods: using neutron activation of the dioxide of tellurium (TeO_2) under a neutron flux which varies between $5 \cdot 10^{11}$ and 10^{13} $\text{n/cm}^2\text{s}$ for 4, 6 and 8 hours** per irradiation cycle during 5 d, and used the Fortron90 Code to calculate the activity of iodine-131.

Results: The result of the activity of iodine-131 found about 4,634 Curie with an irradiation of 4 hours** per day and 9.381 Curie with an activation of 8 hours** per day.

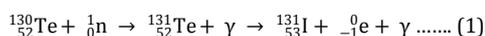
Conclusion: Production of iodine-131 can be very effective if an acceptable capsule is used for different masses of tellurium and a neutron flux in a nuclear reactor.

Keywords: Iodine-131, Tellurium, Neutron activation, Thyroid, Cancer, Nuclear medicine, Becquerel

© 2016 The Authors. Published by Innovare Academic Sciences Pvt Ltd. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>)
DOI: <http://dx.doi.org/10.22159/ijpps.2016v8i11.13595>

INTRODUCTION

Radioisotopes used for many years in many fields of medicine [1-14], industry [15], food and science, [16-17]. Accelerators and nuclear reactors are frequently used for the production of radioisotopes, at the moment, the demand for medical isotopes increases, such as iodine-131, [18-32], but it raises problems of cost and transportation of the producer country to consumer countries, which requires a new strategy for countries who had research reactors to produce iodine-131 [33]. Such as Morocco, which has personal skills, researchers and academics in the nuclear field. Furthermore, Morocco is also a well-developed infrastructure which uses the nuclear research reactor TRIGA-MARC II 2-MW installed at the National Centre of Energy and Science and Nuclear Techniques in Rabat (CNESTEN). This facility meets the global regulatory requirements, nuclear safety, radiation protection and environmental protection. At this time, some studies are made to the feasibility of the production ^{131}I , B. El Bakkari [34], A. S. Elom Achoribo [35], O. Yu. Kochnov [36], M. A. El-Absy [37], D. IAEA CDOC[38]. There are two main methods for the production of ^{131}I , the first way is the fission of ^{235}U [36] and the second way is a nuclear reaction (1) of ^{130}Te , the last method is the one which attracts my interest.



With $t_{1/2}(^{131}\text{Te}) = 25$ min and $t_{1/2}(^{131}\text{I}) = 8.02$ d

Research reactors cannot function for a long time because they should be stopped sporadically to make them cold, so a discontinuous operation is obligatory for the production of iodine-131. We call it cyclical irradiation, [40-42]. It is an irradiation of samples for a moment which can take hours. After the shutdown of the irradiation for some hours; we repeat this method for many cycles until the attainment of our objective.

The main aim of this theoretical study was to search the feasibility of producing ^{131}I . We are interested in a range of neutron flux in the channel of irradiation from $5 \cdot 10^{11}$ to 10^{13} $\text{n/cm}^2\text{s}$ which is generated by research nuclear reactor and capsulated targets of TeO_2 with the mass from 1 g to 150 g. By using an analytical method and the cyclic neutron activation technical, we calculate

the following activities of 4, 6 and 8 hours** of irradiation per day to achieve our goal.

METHODS AND MATERIALS

Production of-131

Productions of iodine-131 are generated by the following equation,

$$\frac{dN(^{131}\text{I})}{dt} = \sigma(^{130}\text{Te})\phi N(^{130}\text{Te})(1 - e^{-\lambda(^{131}\text{Te})t}) - \lambda(^{131}\text{I})N(^{131}\text{I}) \dots \text{Equation 1}$$

With,

σ Cross section microscopic (barn)

ϕ Neutron fluxes $\text{n/cm}^2\text{s}$

λ Radiative constant h^{-1}

At $t=0$ are found that $N(^{131}\text{I}) = N(^{131}\text{Te}) = 0$

By integrating equation 1 can give the number of formal's core in each irradiation cycle which is in Equation 2, [34] [36][43-48].

$$N(^{131}\text{I}) = \sigma(^{130}\text{Te})\phi N(^{130}\text{Te}) \left(\frac{1 - e^{-\lambda(^{131}\text{I})t_{\text{irrad}}}}{\lambda(^{131}\text{I})} + \frac{e^{-\lambda(^{131}\text{Te})t_{\text{irrad}}} - e^{-\lambda(^{131}\text{I})t_{\text{irrad}}}}{\lambda(^{131}\text{Te}) - \lambda(^{131}\text{I})} \right) e^{-\lambda(^{131}\text{I})t_{\text{decay}}} + \frac{(1 - e^{-\lambda(^{131}\text{Te})t_{\text{irrad}}})e^{-\lambda(^{131}\text{Te})t_{\text{decay}}} - e^{-\lambda(^{131}\text{I})t_{\text{decay}}}}{\lambda(^{131}\text{I}) - \lambda(^{131}\text{Te})} \dots \text{Equation 2}$$

Or t_{irrad} : Irradiation time

t_{decay} : Decay time at the end of irradiation.

$$\text{And } N(^{130}\text{Te}) = \frac{m(^{130}\text{Te})N_A}{M(^{130}\text{Te})} \dots \text{Equation 3}$$

$m(^{130}\text{Te}) = 6.02 \cdot 10^{23}$ mol^{-1} , Avogadro's number;

$m(^{130}\text{Te})$ the weight in grams of tellurium;

$M(^{130}\text{Te})$ the atomic mass of irradiated Tellurium isotope.

We can extract activity of ^{131}I in each irradiation cycle using equation 4

$$A(^{131}\text{I}) = \lambda(^{131}\text{I}) N(^{131}\text{I}) \dots\dots \text{Equation 4}$$

With A activity by Curie,

As it was motioned at the beginning of the irradiation (^{130}Te) in the form of a cyclic radiation from each study, with 4, 6 and 8 hours** respectively per day for 5 d, the fig. 1 explains the steps of irradiation in each cycle.

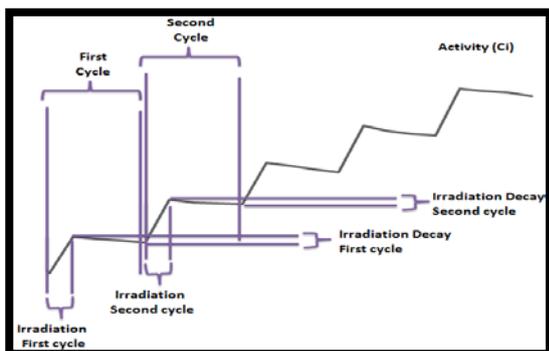


Fig. 1: Evolution of activity with a cyclic irradiation

By accumulation of activity for each cycle is,

For the 1st cycle:

$$A(\text{cycle 1}) = A_{\text{irrad time of first cycle}}(^{131}\text{I}) - A_{\text{decay (8h to 24h) of first cycle}}(^{131}\text{I})$$

For the 2nd cycle:

$$A(\text{cycle 2}) = A(\text{cycle 1}) + A_{\text{irrad time of second cycle}}(^{131}\text{I}) - A_{\text{decay time of second cycle}}(^{131}\text{I})$$

For the 3rd cycle:

$$A(\text{cycle 3}) = A(\text{cycle 2}) + A_{\text{irrad time of third cycle}}(^{131}\text{I}) - A_{\text{decay time of third cycle}}(^{131}\text{I})$$

For the 4th cycle:

$$A(\text{cycle 4}) = A(\text{cycle 3}) + A_{\text{irrad time of fourth cycle}}(^{131}\text{I}) - A_{\text{decay time of fourth cycle}}(^{131}\text{I})$$

For the 5th cycle:

$$A(\text{cycle 5}) = A(\text{cycle 4}) + A_{\text{irrad time of fifth cycle}}(^{131}\text{I}) - A_{\text{decay time of fifth cycle}}(^{131}\text{I})$$

Finally, we found the global activity of ^{131}I ,

After the final calculation, the results obtained are shown schematically in the form of figures.

RESULTS AND DISCUSSION

From the simulations obtained in five irradiation cycles for each cycle with 4, 6 and 8 hours** of irradiation in a neutron flux $5 \cdot 10^{11}$, 10^{12} , $5 \cdot 10^{12}$ and $10^{13}\text{n/cm}^2\text{s}$, we have observed clearly that the activity increases during 4, 6 and 8 hours**.

After the stoppage of the first irradiation, the activity decreases slightly in the hours after (radioactive decay of Iodine-131). When we start the second cycle, the activity increases in the irradiation period and it decreases in the cooling period.

This process continues to increase and decrease until the end of the fifth round to achieve the maximum activity, fig. from 4 to 15 show the variation of the activity of ^{131}I at different target weights (5g,

10g, 30g, 50g, 100g and 150g) and attached to a neutron flux ranges from $5 \cdot 10^{11}$ to $10^{13} \text{n/cm}^2\text{s}$, for irradiation time 4, 6 and 8 hours** in each cycle. This work was well validated with the study of A. S. Elom Achoribo [35] and B. El Bakkari [34].

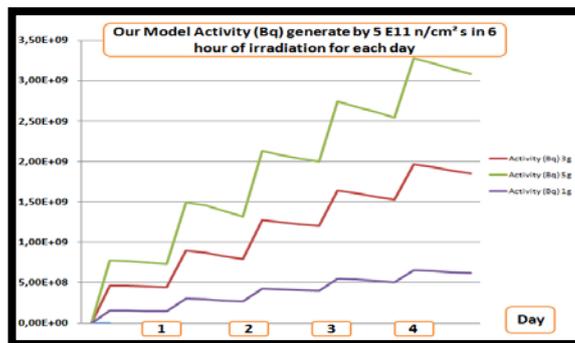


Fig. 2: Validation of our model, evolution of activity (Bq) in 6h of irradiation in each day with $5 \cdot 10^{11} \text{n/cm}^2\text{s}$

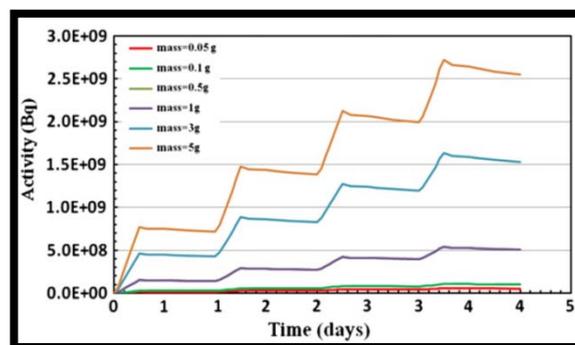


Fig. 3: Evolution of activity [35]

To validate our results we compared them with the results of AS Elom Achorido [35] (fig. 3). For 1g, 3g and 5g, with a neutron flux of $5 \cdot 10^{11}\text{n/cm}^2\text{s}$, masses of TeO_2 irradiated for 6 hours** per day (fig. 2), the results show the validation of our work. In our study, we are interested in 5g, 10g, 30g, 50g, 100g and 150g.

Productions ^{131}I with 4 hours of irradiation for each cycle**

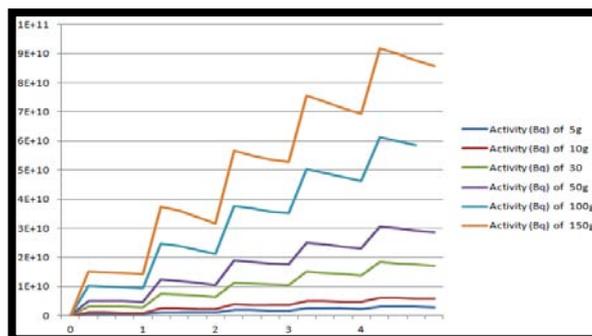


Fig. 4: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of $5 \cdot 10^{11} \text{n/cm}^2\text{s}^{-1}$ for an irradiation time of 4h during each day

After four hours of irradiation by cycle, the results which are shown in fig. 4, 5, 6 and 7 clarify that the activity increases with irradiation time, then a slight decrease at the end of irradiation, then it begins to increase after the second irradiation. For example, the neutron flux of $5 \cdot 10^{11}\text{n/cm}^2\text{s}$ for masses of 5g, 10g, 50g, 100g and 150g this activity is $2.86 \cdot 10^9$, $5.72 \cdot 10^9$, $1.71 \cdot 10^{10}$, $2.86 \cdot 10^{10}$, $5.72 \cdot 10^{10}$ and $8.57 \cdot 10^{10}$ Becquerel respectively. if we increase the fluxes to $10^{13} \text{n/cm}^2\text{s}$, and for the same masses, the activity is equal to $3.80 \cdot 10^{10}$, $1.14 \cdot 10^{11}$, $3.43 \cdot 10^{11}$, $5.72 \cdot 10^{11}$, $1.14 \cdot 10^{12}$ and $1.71 \cdot 10^{12}$ Becquerel, respectively.

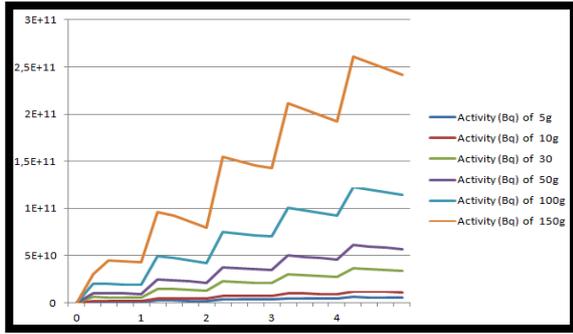


Fig. 5: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of 10^{12} n/cm²s⁻¹ for an irradiation time of 4h during each day

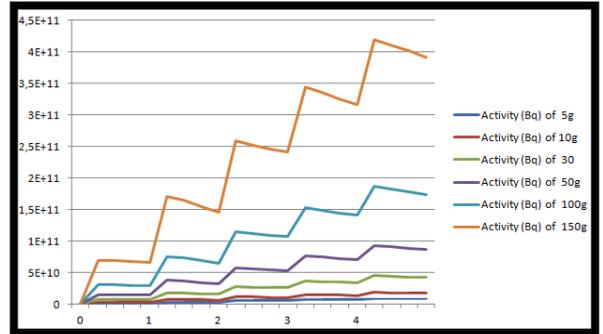


Fig. 9: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of 10^{12} n/cm²s⁻¹ for an irradiation time of 6h during each day

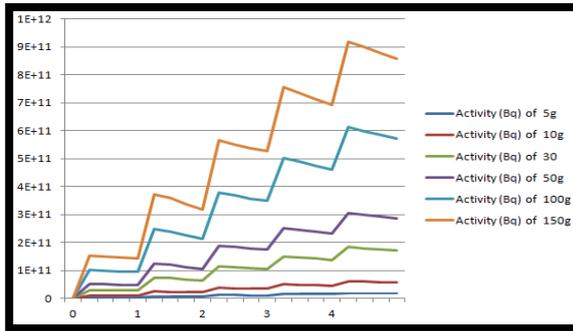


Fig. 6: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of $5 \cdot 10^{12}$ n/cm²s⁻¹ for an irradiation time of 4h during each day

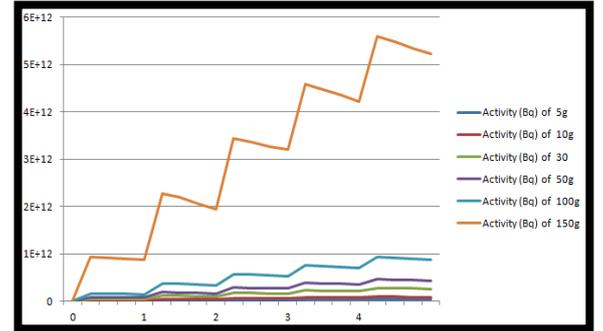


Fig. 10: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of $5 \cdot 10^{12}$ n/cm²s⁻¹ for an irradiation time of 6h during each day

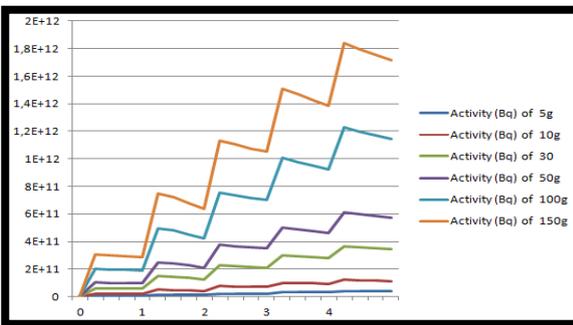


Fig. 7: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of 10^{13} n/cm²s⁻¹ for an irradiation time of 4h during each day

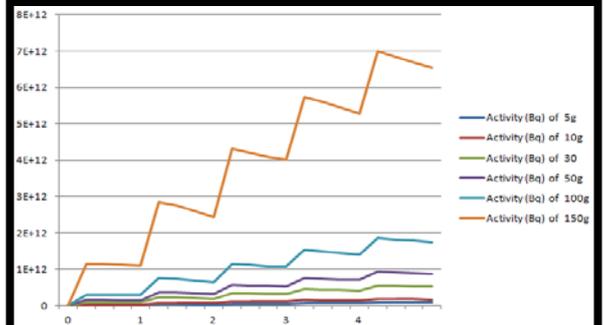


Fig. 11: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of 10^{13} n/cm²s⁻¹ for an irradiation time of 6h during each day

Productions ¹³¹I with 6 hours of irradiation for each cycle**

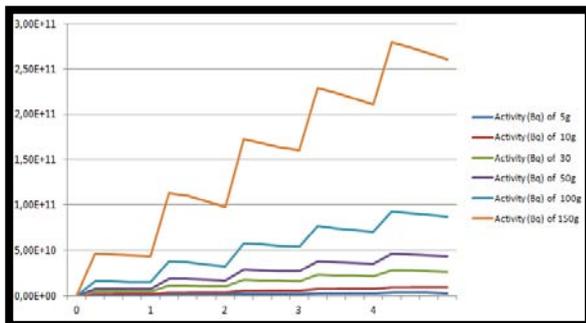


Fig. 8: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of $5 \cdot 10^{11}$ n/cm²s⁻¹ for an irradiation time of 6h during each day

The schemes illustrated in fig. 8, 9, 10 and 11 explain that the activity clearly related to the level of neutron flux and the mass of samples in addition to the microscopic Cross section of target of tellurium. After the analysis of the evolution of the activity, we noticed that the activity increases every day and every increase of irradiation followed directly with a slight decrease. For example, the neutron flux of $5 \cdot 10^{11}$ n/cm²s and masses of 5g, 10g, 30g, 50g, 100g and 150g, activity was $3.08 \cdot 10^9$, $8.71 \cdot 10^9$, $2.61 \cdot 10^{10}$, $4.35 \cdot 10^{10}$, $8.71 \cdot 10^{10}$ and $2.61 \cdot 10^{11}$ Becquerel respectively. If we increase the flux to 10^{13} n/cm²s, and for the same masses, activity equals $8.71 \cdot 10^{10}$, $1.74 \cdot 10^{11}$, $5.23 \cdot 10^{11}$, $8.71 \cdot 10^{11}$, $1.74 \cdot 10^{12}$ and $6.53 \cdot 10^{12}$ Becquerel, respectively.

Productions ¹³¹I with 8 hours of irradiation for each cycle**

Fig. 12, 13, 14 and 15 show the evolution of ¹³¹I with activation of 8 hours** per day for 5 d. The activity of iodine-131 depends on irradiation time, the neutron flux and the mass of target capsules. For example in fig. 12, the neutron flux equal to $5 \cdot 10^{11}$ n/cm²s, the activity becomes at the end of the cyclic irradiation $5.79 \cdot 10^9$, $1.16 \cdot 10^{10}$, $3.47 \cdot 10^{10}$, $5.79 \cdot 10^{10}$, $1.16 \cdot 10^{11}$ and $1.74 \cdot 10^{11}$ Becquerel,

respectively for 5 g, 10 g, 30 g, 50 g, 100 g et 150 g. Therefore, an increase in the neutron flux to 10^{13} n/cm²s and for the same masses in fig. 15, the activity increases respectively 1.16 10¹¹, 2.31 10¹¹, 6.94 10¹¹, 1.16 10¹², 2.31 10¹² and 3.47 10¹² Becquerel.

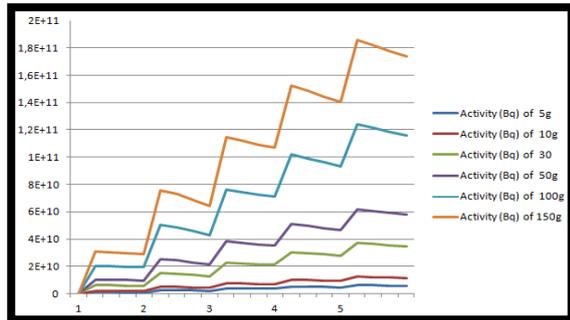


Fig. 12: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of $5 \cdot 10^{11}$ n/cm²s for an irradiation time of 8h during each day

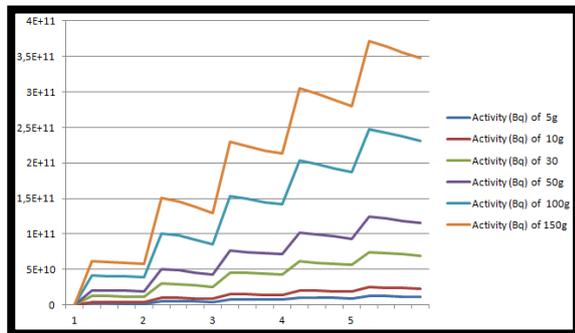


Fig. 13: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of 10^{12} n/cm²s for an irradiation time of 8h during each day

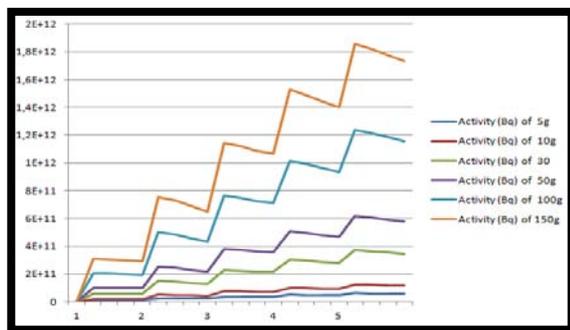


Fig. 14: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of $5 \cdot 10^{12}$ n/cm²s for an irradiation time of 8h during each day

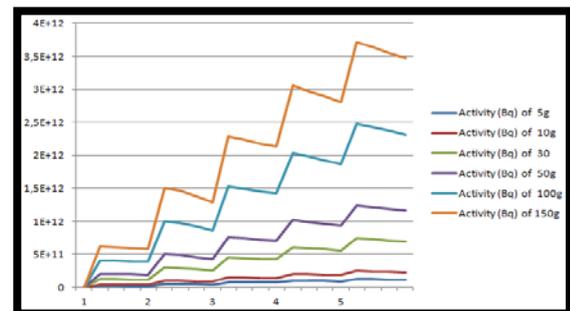


Fig. 15: Theoretical evolution activity (Bq) for I-131 production at a neutron flux of 10^{13} n/cm²s for an irradiation time of 8h during each day

Profile of activity

Maximum feasible activities have been based on the capsules used; when we increase the mass of TeO₂, it causes an increase in the activity. maximum activity in irradiation period 4, 6 and 8 hours** for the mass of 150 g give 2.317 Curie, 3.530 Curie, 4.69 Curie respectively in a flux of $5 \cdot 10^{11}$ n/cm²s, if the neutron flux increases to 10^{12} n/cm²s, for the identical weight (150g) and for the same irradiation period 4, 6 and 8 hours** per day, we note that the activity increases respectively in the following values, 4.6345 Ci, 7.0616 Ci and 9.3818 Curie, fig. 16 and 17.

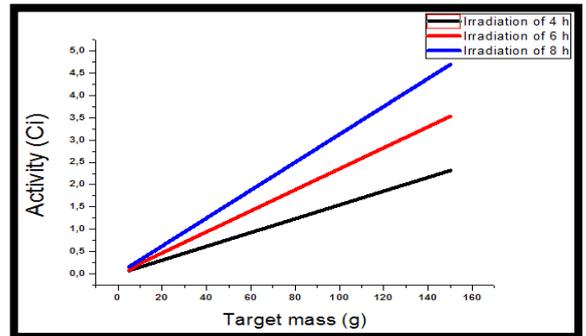


Fig. 16: Increased activity of Iodine-131 as a function of time of irradiation and the mass of the tellurium-130 target in a neutron flux of $5 \cdot 10^{11}$ n/cm²s

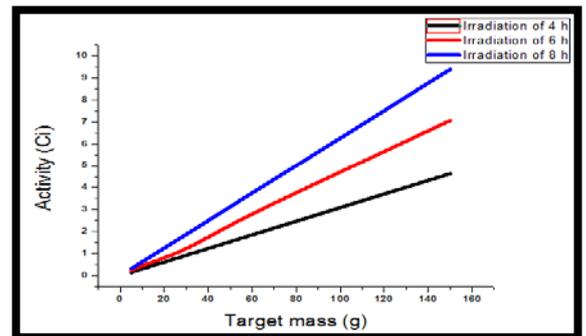


Fig. 17: Increased activity of Iodine-131 as a function of time of irradiation and the mass of the tellurium-130 target in a neutron flux of 10^{12} n/cm²s

CONCLUSION

A feasibility study of the production of iodine-131 is done by using a neutron flux from $5 \cdot 10^{11}$ to 10^{13} n/cm². Then the results were that the mass of 150 g and a flux of $5 \cdot 10^{11}$ n/cm²s the activity gives (2.317 Curie in 4 hours** of irradiation and 4.69 Curie in 8 hours** of irradiation), for the same mass of 160 g in a flux of 10^{12} n/cm²s gives (4.634 Curie in 4 hours** of irradiation and 9.381 Curie in 8 hours** of irradiation). These results are compared with study of B. El Bekkouri [34] and AS Elom Achoribo [35]. We noticed that the neutron standpoint, an acceptable capsule for different masses of tellurium, and an irradiation in a neutron flux in the reactor, approximately in the center, can give continuation to the production of iodine-131.

CONFLICTS OF INTERESTS

Declared none

REFERENCES

- Mbodj M, Guerrouj H, Amjad I, Ben Rais NA. Contribution of radio-iodine 131 in the treatment of Grave's Basedow disease in the department of nuclear medicine of Ibn sina hospital in rabat. J Med Nucl 2009;33:592-8.

2. Schlienger JL, Goichot B, Grunenberge F. Iode et fonction thyroïdienne. *La Revue Méd Int* 1997;9:709-16.
3. Maged Abdel Galil Hamed, Ahmed Fathy Abdel Ghany, Noha Mohamed Osman. The diagnostic usefulness of FDG-PET/CT in detecting tumor recurrence not evident in whole body I-131 scan in differentiated thyroid carcinoma. *Egyptian J Radiol Nucl Med* 2014;45:361-5.
4. René Caquet. Temps de lyse des euglobulines, 250 examens de laboratoire. 11th édition; 2010. p. 211-2.
5. Wémeau JL. Chapitre 17-Iode et thyroïde Les maladies de la thyroïde; 2010. p. 141-8.
6. Taïeb D, Guille DA, Mundler LO. Guidelines for radionuclide imaging of pheochromocytoma and paraganglioma. *J Med Nucl* 2008;32:101-10.
7. Vitaux F. Thyroid gland irradiations and thyroid cancers critical bibliographic journal. *J Med Nucl* 2007;31:350-5.
8. Intidhar El Bez. Cancer de la thyroïde et ablation par iode 131 sous thyroïde: quand doser la thyroglobuline? *Ann Endocrinol* 2013;74:156.
9. Spagnoli V, Azzalini L, Tadros VX, Picard F, Ly HQ. Contrast-induced nephropathy: an update. *Ann Cardiol Angeiol* 2016;65:87-94.
10. Guerrouj H, Elamrani M, Ghfiri, Ben Rais N. Apport de l'iode 131 dans le traitement de l'adénome thyroïdien toxique. *J Med Nucl* 2012;36:561-4.
11. Boisserie G, Hasboun D. Utilisation de l'imagerie multimodalité en radiothérapie. *Cancer/Radiothérapie* 2001;5(1 Suppl 1):15-35.
12. Belkacémi Y, Tsoutsou PG, Comet B, Kerrou K, Lartigau E. Évaluation de la radiosensibilité tumorale par l'imagerie fonctionnelle et métabolique: de la recherche à l'application clinique. *Revue de la littérature. Cancer/Radiothérapie* 2006;10:124-33.
13. Delmaire C. Imagerie des métastases cérébrales. *Cancer/Radiothérapie* 2015;19:16-9.
14. Bhavna Shah. composites from agricultural detritus for pollution remedy. *Int J Pharm Pharm Sci* 2016;3:4-49.
15. Yongchun G, Jijin G, Huabai T, Yuewen Y. Miniature Neutron Source Reactor General Description. China Institute of Atomic Energy. Peking; 1992.
16. Hillaire-Marcel G. Isotopes and food. the Terrestrial Environment, B, Elsevier, Amsterdam; 1986. p. 507-48.
17. Kelly SD, Food authenticity and traceability, A volume in Woodhead Publishing Series in Food Science, Technology and Nutrition; 2003. p. 156-83.
18. Szumowski P, Rogowski F, Abdelrazek S, Kociura-Sawicka A, Sokolik-Ostasz A. Iodine isotope ¹³¹I therapy for toxic nodular goitre: treatment efficacy parameters. *Nucl Med Rev Cent East Eur* 2012;15:713.
19. Stokkel MP, Handkiewicz Junak D, Lassmann M, Dietlein M, Luster M. EANM procedure guidelines for therapy of benign thyroid disease. *Eur J Nucl Med Mol Imaging* 2010;37:2218-28.
20. Kaniuka S, Lass P, Sworczak K. Radioiodine an attractive alternative to surgery in large non-toxic multinodular goitres. *Nucl Med Rev Cent East Eur* 2009;12:239.
21. Fast S. Prestimulation with recombinant human thyrotropin (rhTSH) improves the long-term outcome of radioiodine therapy for multinodular nontoxic goiter. *J Clin Endocrinol Metab* 2012;97:2653-60.
22. Giusti M. Long-term outcome after radioiodine therapy with adjuvant rhTSH treatment: comparison between patients with nontoxic and pre-toxic large multinodular goiter. *Endocrine* 2014;45:221-9.
23. Verelst J, Bonnyns M, Glinier D. Radioiodine therapy in voluminous non-toxic goiter. *Acta Endocrinol* 1990;122:417-21.
24. Huysmans DA, Buijs WC, van de Ven MT. Dosimetry and risk estimates of radioiodine therapy for large, multinodular goiters. *J Nucl Med* 1996;37:2072-9.
25. Sun XS. Radiation therapy in thyroid cancer. *Cancer/Radiothérapie* 2013;17:233-43.
26. Schlumberger M, Chevillard S, Ory K, Dupuy C, Le Guen B, de Vathaire F. Cancer de la thyroïde après exposition aux rayonnements ionisants. *Cancer/Radiothérapie* 2011;15:394-9.
27. Raoul JL. Traitement des carcinomes hépatocellulaires par injection intra-artérielle de radio-isotopes. *Cancer/Radiothérapie* 2011;15:64-8.
28. Cecconi A, Blotta A, Ntreta M, Busutti L. La radiothérapie métabolique par iode 131 et la radiothérapie transcutanée dans une population avec long suivi. *Cancer/Radiothérapie* 2007;1:429.
29. Dutrillaux B. Les cancers radio-induits. *Cancer/Radiothérapie* 1998;2:541-8.
30. Mallet F. Faisabilité et toxicité d'une séance unique de curiethérapie de haut débit de dose suivie d'une irradiation externe dans le cancer localisé de la prostate: étude rétrospective de la polyclinique de courlancy. *Cancer/Radiothérapie* 2010;14:11-8.
31. Mohammed Azharuddin. Evaluation of anti-thyroid activity of ficus racemosa linn bark in male rats. *Int J Pharm Pharm Sci* 2015;7:118-22.
32. Abdul Aziz Ramadan, Hasna Mandil, Jenan Sabouni. Determination of atorvastatin calcium in pure and its pharmaceutical formulations using iodine in acetonitrile by uv-visible spectrophotometric method. *Int J Pharm Pharm Sci* 2015;7:427-33.
33. Gerbaulet AP. Quel avenir pour la curiethérapie? *Cancer/Radiothérapie* 1999;3(1, Suppl 1):11-27.
34. El Bakkari B. analysis of I-131 production in the Moroccan TRIGA research reactor. *Ann Nucl Energy* 2015;78:140-5.
35. Elom Achoribo AS. Feasibility study for production of I-131 radioisotope using MNSR research reactor. *Appl Radiat Isot* 2012;70:76-80.
36. Yu O Kochnov, Kolesov VV, Fomin RV, Jerdev GM. Assessment of the increasing in 131-I production due to improved tellurium target in the WWR-c reactor core. *Nucl Eng Sci Technol* 2015;1:213-7.
37. El-Absy MA, El-Garhy MA, El-Amir MA, Fasih TW, El-Shahat MF. Separation and purification of 131 I from neutron irradiated tellurium dioxide targets by wet-distillation method. *Separation Purification Technol* 2010;71:1-12.
38. International Atomic Energy Agency. Manual for reactor produced radioisotopes, IAEA/TECDOC-1340. IAEA: Vienna, Austria; 2003. p. 121-4.
39. Daniel Cestau. Production of Iodine-131 from Low Enriched Uranium Targets International Meeting on Reduced Enrichment for Research and Test Reactors, Cape Town, South Africa; 2006.
40. Tout RE, Chatt A. The effect of sample matrix on selection of optimum timing parameters in cyclic neutron activation analysis. *Anal Chim Acta* 1981;133:409-19.
41. Parijat Pandey, Mandeep Dahiya. A brief review on inorganic nanoparticles. *Int J Pharm Pharm Sci* 2014;6:34-41.
42. Shefali arora, Shilpi Agarwal, Shailey Singhal. Anticancer activities of thiosemicarbazides/thiosemicarbazones. *Int J Pharm Pharm Sci* 2014;6:34-41.
43. Lieser KH. Nuclear and radiochemistry: fundamentals and applications, second ed. Wiley-WCH, New York; 2001.
44. Mirzadeh S, Walsh P. Numerical evaluation of the production of radionuclides in a nuclear reactor (Part I). *Appl Radiat Isot* 1998;49:379-82.
45. Kassakov M. Analyse par activation neutronique de substances ayant des sections efficaces macroscopiques élevées pour l'absorption de neutrons thermiques. M. Sc. A. Ecole Polytechnique de Montreal, Montréal; 2006.
46. Revel G. Analyse par activation. Technique de l'ingénieur, traite analyse et caractérisation; 2009. p. 1-21.
47. Abdessamad Didi. Calculating concentrations of elements in sample and compare with standard certified results of the International Atomic Energy Agency (IAEA) Soil-7. *Der Pharm Chem* 2016;8:250-5.
48. DIDI. New design of thermal neutron flux distribution of Am-Be neutron source irradiation in paraffin moderator using MCNP-6. *Mor J Chem* 2016;4:285-8.

How to cite this article

- Abdessamad Didi, Ahmed Dadouch, Hassane EL Bekkouri. Feasibility study for production of iodine-131 using dioxide of tellurium-130. *Int J Pharm Pharm Sci* 2016;8(11):327-331.