

YIELDS OF CYCLOTRON PRODUCED MEDICAL ISOTOPES: A COMPARISON OF
THEORETICAL POTENTIAL AND EXPERIMENTAL RESULTS *

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Abstract

Experimentally obtained yields of most of the medical radioisotopes, produced with cyclotrons through different nuclear reactions at various bombarding energies in laboratories around the world, are presented. These yields are compared with those calculated using experimentally measured cross sections (where available) at similar bombarding conditions. Where experimental cross sections are unavailable, empirically constructed excitation functions have been used.

The information provided in this paper would be a valuable aid in selecting the most suitable nuclear reaction and bombarding conditions for producing a particular radioisotope and in assessing various losses of the isotope during chemical processing of the irradiated target.

Introduction

Production of radioisotopes for diagnostic studies is by far the most well known medical application of cyclotrons, and most medical cyclotrons are occupied in the production of various isotopes for research and routine work in nuclear medicine and nuclear biology. Generally neutron-deficient, carrier free, and shorter-lived isotopes, which cannot be produced in a reactor, are produced with cyclotrons. However, at the same time, neutron enriched isotopes can also be produced with cyclotrons, if required, through reactions of the type (d,p), (³He,p), (α,p) etc.

Method and Results

Cyclotron production of most of the isotopes in current use has been summarized in Table I.

Thick-target yields at saturation of most isotopes through different nuclear reactions have been calculated¹ using experimentally measured (for ¹¹C, ¹³N, ¹⁵O, ¹⁸F and ⁵²Fe) or empirically constructed excitation functions² and range-energy data³. In the calculations, the isotopic abundance of the particular isotope contributing to the nuclear reaction has been taken into consideration. However, the matrix which slows down the incoming particle beam in the target is assumed to consist of only the element taking part in the nuclear reaction, which is true when elemental rather than compound targets are being used. The calculated yields are designed as guide-lines only to optimize actual production and may be in error by as much as a factor of two in the case of empirically constructed excitation functions². For comparison between the actually measured and calculated yields, the saturation yields would have to be converted into yields at time 't' where 't' is the time for actual bombardment, by using the factor (1 - e^{-λt}).

In compiling the Table, an attempt has been made to include most of the published data regarding the production of various isotopes, in biomedical use, using different nuclear reactions. However, it is possible that some particular isotope publications might have been inaccessible or inadvertently omitted. Only the first or the first significant published paper for radioisotope production under any particular bombarding conditions, such as energy, target material etc., has been included.

The operating costs of a small cyclotron in a developed country are estimated to be around US\$60-90 per hour of useful machine time. Keeping this figure in mind, one can calculate the expected cost per millicurie of any particular isotope. However, the man-hours required for the chemical processing of the irradiated target must also be included in the overall cost estimate.

TABLE 1

| Isotope | Reaction | Energy (MeV) | Target | Production yield μCi/μAh, unless otherwise indicated | Ref. | Calculated yield at saturation ¹ | |
|-----------------|-------------------------------------|--------------|-------------------------------------|---|------|---|--------------|
| | | | | | | mCi/μA | Energy (MeV) |
| ¹¹ C | ¹¹ B(p,n) | | | ----- | --- | 405 | 11.5 |
| | ¹⁰ B(d,n) | 14 | B ₂ O ₃ | 5 mCi/min CO - 100 mCi per litre of H ₂ carrier gas - | 4 | 80 | 14 |
| | ¹¹ B(d,2n) | | | 70 mCi per 35 ml of He CO ₂ - 50 mCi in 35 ml of He | | | |
| | ¹² C(³ He,α) | 15-18 | CaC ₂ | 2500 | 5 | 87 115 | 15 18 |
| ¹⁴ N | ¹⁴ N(p,α) | 15 | LiNH ₂ | 18 mCi/18 min of H ¹¹ C N | 6 | --- | --- |
| | " | 18 | 5% H ₂ in N ₂ | 90 mCi/μAh | 7 | --- | --- |
| ¹³ N | ¹² C(d,n) | 14 | Graphite | 30 mCi/ml in gas form 100-300 μCi per ml in solution | 8 | 300 | 14 |
| | ¹⁴ N(³ He,α) | 30 | N ₂ | 15 mCi/μAh | 9 | 38 | 30 |

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| Isotope | Reaction | Energy (MeV) | Target | Production yield $\mu\text{Ci}/\mu\text{Ah}$, unless otherwise indicated | Ref. | Calculated yield at saturation ¹ $\text{mCi}/\mu\text{A}$ | Energy (MeV) |
|--------------------------|---|---------------------|--|---|------|--|--------------|
| ^{13}N | $^{16}\text{O}(\text{p},\alpha)$ | 18 | H_2O | 24 $\text{mCi}/\mu\text{Ah}$ as NH_3 | 10 | --- | --- |
| ^{15}O | $^{14}\text{N}(\text{d},\text{n})$ | 5 | 4% O_2 in N_2 | 120 mCi of O_2 per litre of carrier gas, 40 mCi/ml of H_2O | 4 | 12.5 | 5 |
| | $^{14}\text{N}(^3\text{He},\text{d})$ | --- | ---- | ----- | --- | 26 | 17 |
| | $^{16}\text{O}(\text{p},\text{pn})$ | 27 | O_2 | 1.5 mCi of ^{15}O per 15 ml of CO_2 - 6 mCi/min of 0-15 labelled CO_2 at 15 μA | 11 | --- | --- |
| ^{18}F | $^{16}\text{O}(\alpha,\text{d})$ | 30 | H_2O | 1100 | 12 | 5.4 | 30 |
| | (α,pn) | 40 | " | 11000 | 13 | 27 | 40 |
| | $(\alpha,2\text{n})$ | 40 | O_2 | 14300 | 13 | --- | --- |
| | | 65 | H_2O | 19000 | 14 | --- | --- |
| | $^{16}\text{O}(^3\text{He},\text{p})$ | 22 | H_2O | 6000 | 15 | 12 | 9.5 |
| | $(^3\text{He},\text{n})$ | 30 | H_2O | 8500 | 13 | 32 | 30 |
| | | --- | ---- | ----- | --- | 43 | 35 |
| | $^{19}\text{F}(^3\text{He},\alpha)$ | --- | ---- | ----- | --- | 8 | 30 |
| | $^{20}\text{Ne}(\text{d},\alpha)$ | 8 | Neon Gas | 10000 | 16 | 52 | 8 |
| | | --- | ---- | ----- | --- | 100 | 16 |
| ^{28}Mg | $^{27}\text{Al}(\alpha,3\text{p})$ | 45 | Al | 6.2 | 17 | --- | --- |
| | | 140 | Al | 35 | 18 | --- | --- |
| ^{43}K | $^{40}\text{Ar}(\alpha,\text{p})$ | 17 | Argon | 57 | 19 | 9 | 17 |
| ^{52}Fe | $^{50}\text{Cr}(\alpha,2\text{n})$ | 30 | natural Cr | 3.3 | 20 | .007 | 30 |
| | $^{52}\text{Cr}(^3\text{He},3\text{n})$ | 45.5 | " | 50 | 21 | 9 | 45 |
| | $^{55}\text{Mn}(\text{p},4\text{n})$ | 23 | MnO_2 | 0.7 | 22 | 0.18 | 23 |
| | | 65 | | 160 | 23 | 90 | 65 |
| ^{62}Zn | $^{63}\text{Cu}(\text{p},2\text{n})$ | 38 \rightarrow 18 | Cu-foils 1.6 g/cm^2 | 6000 | 24 | --- | --- |
| ^{67}Ga | $^{60}\text{Ni}(\alpha,2\text{n})$ | 30 | natural Ni | 100 | 25 | 6 | 30 |
| | $^{65}\text{Cu}(\alpha,2\text{n})$ | 30 | natural Cu | 160 | 26 | 8 | 30 |
| | $\text{Zn}(\text{p},\text{xn})$ | 22 | natural Zn | 430 | 27 | 95 | 22 |
| | $\text{Zn}(\text{d},\text{xn})$ | 8 | " | 100 | 28 | 2 | 8 |
| | " | " | " | 30 | 22 | --- | --- |
| | " | 16 | " | 340 | 29 | --- | --- |
| | " | 16 | enriched ^{66}Zn (90%) | 946 | 29 | --- | --- |
| ^{77}Br | $^{75}\text{As}(\alpha,2\text{n})$ | 28 | As_2O_5 | 160 | 30 | 25 | 28 |
| | " | 28 | As_2O_3 | 290 | 31 | --- | --- |
| | " | 28 | As metal | 170 | 31 | --- | --- |
| | $^{79,81}\text{Br}(\alpha,6\text{n})$ | 100 | NaBr | 322 | 32 | --- | --- |
| | (α,pxn) | | | | | | |
| | (d,pxn) | 55 \rightarrow 20 | " | 550 | 33 | --- | --- |
| | (d,xn) | " | " | 640 | 33 | --- | --- |
| $^{81\text{m}}\text{Kr}$ | | | Decay product of ^{81}Rb | | | | |
| $^{85\text{m}}\text{Kr}$ | $^{84}\text{Kr}(\text{d},\text{p})$ | 15 | Kr | 790 | 34 | 60 | 15 |
| ^{81}Rb | $^{79}\text{Br}(\alpha,2\text{n})$ | 30 | NaBr | 2000 | 35 | 18 | 30 |
| | | 50 | " | 2900 | 36 | | |
| | $^{81}\text{Br}(^3\text{He},\text{n})$ | 22 | NaBr | 30 | 37 | 0.13 | 22 |
| $^{82\text{m}}\text{Rb}$ | $^{81}\text{Br}(^3\text{He},2\text{n})$ | 22 | NaBr | 80 | 37 | 1.2 | 22 |
| ^{83}Rb | $^{83}\text{Kr}(\text{p},\text{n})$ | 22 | Natural Kr-gas | 7 | 38 | 60 | 22 |
| ^{85}Sr | $^{85}\text{Rb}(\text{d},2\text{n})$ | 13 | RbCl | 15 | 39 | 70 | 13 |
| ^{87}Y | $^{85}\text{Rb}(\alpha,2\text{n})$ | 32 | RbCl | 174 | 40 | 30 | 32 |

| Isotope | Reaction | Energy | Target | Production yield μCi/μAh, unless otherwise indicated | Ref. | Calculated yield at saturation ¹ | | |
|--|---------------------------------------|-------------------------|--|--|-------|--|--------------|-----|
| | | | | | | mCi/μA | Energy (MeV) | |
| ¹¹¹ In | ¹⁰⁹ Ag(α,2n) | 30 | natural Ag | 200 | 41 | 9 | 30 | |
| | Cd(p,xn) | 15 | natural Cd | 140 | 22 | 35 | 15 | |
| | ¹¹¹ Cd(p,n) | 16 | enriched ¹¹¹ Cd | 515 | 42 | 150 | 16 | |
| | ¹¹² Cd(p,2n) | 22 | natural Cd | 1035 | 43 | 300 | 22 | |
| | ¹¹⁰ Cd(d,n) | 12 | natural Cd | 117 | 43 | 20 | 12 | |
| ¹²³ I | ¹²¹ Sb(α,2n) | 25 | natural Sb | 150 | 44 | 3 | 25 | |
| | | 25-36 | enriched ¹²¹ Sb | 900 | 45 | 6-30 | 25-36 | |
| | ¹²¹ Sb(³ He,n) | 23 | natural Sb | 24 | 25 | 0.22 | 23 | |
| | ¹²² Te(d,n) | 6-9 | enriched ¹²² Te | 100 | 45 | 0.3 | 9 | |
| | ¹²³ Te(p,n) | 15.5 | enriched ¹²³ Te | 450 | 46 | 140 | 16 | |
| | ¹²⁴ Te(p,2n) | 30 | enriched ¹²⁴ Te | 40000 | 47 | 1300 | 30 | |
| | ¹²³ Xe | ¹²² Te(α,3n) | 46 | enriched ¹²² Te | 5000 | 48 | 70 | 46 |
| ¹²³ Te(³ He,3n) | | 30 | enriched ¹²³ Te | 750 | 49 | 20 | 30 | |
| " | | 32 | enriched (¹²³ Te (88%)) | 4954 | 50 | 25 | 32 | |
| " | | 35 | " | 6441 | 50 | 30 | 35 | |
| ¹²⁴ Te(³ He,4n) | | 38 | enriched ¹²⁴ Te (96%) | 1000 | 50 | 8.5 | 38 | |
| " | | 52 | " | 7155 | 50 | 33 | 52 | |
| ¹²⁷ I(p,5n) | | | 60 → 50 | NaI powder 1.5g/cm ² | 5600 | 24 | --- | --- |
| | | | 52 | CH ₂ I ₂ (Circulating liquid) | 1000 | 51 | --- | --- |
| | | | 53 | " " | 2200 | 52 | --- | --- |
| | | | 72 | KI & NaI | 11500 | 53 | --- | --- |
| | ¹²⁷ I(d,6n) | 78 | NaI | 8000 | 54 | --- | --- | |
| ¹²⁷ Cs | ¹²⁷ I(³ He,3n) | 22 | NaI | 500 | 55 | 6 | 22 | |
| ¹²⁹ Cs | ¹²⁷ I(α,2n) | 30 | NaI | 170 | 56 | 18 | 30 | |
| | | 35 | " | 300 | 57 | 30 | 35 | |
| | | 36 | " | 700 | 58 | 30 | 36 | |
| ¹⁵⁷ Dy | ¹⁵⁹ Tb(p,3n) | 30 | Terbium foil | 23000 | 59 | 500 | 30 | |
| | | | TbCl ₃ ·6H ₂ O | 2500 | 60 | --- | --- | |
| | ¹⁵⁵ Gd(α,2n) | 30 | Gd | 80 | 61 | 3 | 30 | |
| ¹⁹⁷ Hg | ¹⁹⁷ Au(p,n) | 12.5 | Gold | 14 | 62 | 2 | 12.5 | |
| ^{197m} Hg | ¹⁹⁷ Au(p,n) | 12.5 | Gold | 15 | 62 | 10 | 12.5 | |
| ²⁰³ Pb | ²⁰³ Tl(p,n) | 15 | Tl metal | 50 | 63 | 1 | 15 | |
| | ²⁰³ Tl(d,2n) | 16 | Tl ₂ O | 100 | 4 | 6 | 16 | |
| ²⁰⁴ Bi | ²⁰⁶ Pb(p,3n) | 32 | Pb | 2000 | 64 | 180 | 32 | |
| ²⁰⁶ Bi | ²⁰⁷ Pb(p,2n) | 22 | Pb | 700 | 65 | 110 | 22 | |
| | ²⁰⁶ Pb(d,2n) | 16 | Pb | 30 | 4 | 6 | 16 | |

| Isotope | Reaction | Energy (MeV) | Target | Production yield $\mu\text{Ci}/\mu\text{Ah}$, unless otherwise indicated | Ref. | Calculated yield at saturation ¹ $\text{mCi}/\mu\text{A}$ | Energy (MeV) |
|-------------------|-------------------------|--------------|------------|---|------|--|--------------|
| ^{201}Tl | $^{203}\text{Tl}(p,3n)$ | 30 | natural Tl | 700 | 66 | --- | --- |
| | | 14 | natural Hg | 180 | 67 | --- | --- |
| | Hg(p,n) | 16 | " | 350 | 67 | --- | --- |
| | | 20 | " | 350 | 67 | --- | --- |

References

1. Chaudhri, M. A. To be published.
2. Lange, J. and Muenzel, H. Report KFK-767 (1968) Gesellschaft fuer Kernforschung m.b.H. Karlsruhe.
3. Williamson, C.F., Boujot, J.P. and Picard, J. Rapport C.E.A. - R3042 (1966).
4. MRC Cyclotron Unit. List of Radioactive Products, MRC-Cyclotron Unit, Hammersmith Hospital, London (1969).
5. Myers, W.G. J. Nucl. Med. 13 (1972).699.
6. Lamb, J.F., James, R.W. and Winchell, H.S. Int. J. Appl. Radiat. Isotopes 21.(1970).475.
7. Christman, D.R., Finn,R.D., Karlstron, K.I. and Wolf, A.P. Int. J. Appl. Radiat. Isotopes 26.(1975).435.
8. Buckingham, P.D. and Clark, J.C. Int. J. Appl. Radiat. Isotopes, 23.(1972).5.
9. Krasnov, N.N., Dmitriev, P.P., Dmitrieva, Z.P., Konstantinov, I.O. and Molin, G.A. Atomnaya Energiya, 28,(1970)503-504.
10. Suzuki, K. and Iwata, Ren. Int. J. Appl. Radiat. Isotopes 28.(1977) 663-665.
11. Beaver, J.E., Finn, R.D. and Hupf, H.B. Int. J. Appl. Radiat. Isotopes.27(1976) 195-197.
12. Clark, J.C. and Silvester, D.J. Int. J. Appl. Radiat. Isotopes 17 (1966).151.
13. Nozaki, T., Tanaka, Y., Shimamura, A. and Karasawa. Int. J. Appl. Radiat. Isotopes 19.(1968).27.
14. Yano, Y., Van Dyke, D.C., Verdon, T.A. and Anger, H.O. J. Nucl. Med. 12.(1971).815.
15. Tilbury, R.S., Dahl, J.R., Mamacos, J.P. and Laughlin, J.S. Int. J. Appl. Radiat. Isotopes 21.(1970).277.
16. Harper, P.V., Lembares, N. and Krisek, J. J. Nucl. Med. 12.(1971).362.
17. Nozaki, T., Furukawa, M., Kume, S. and Seki, R. Int. J. Appl. Radiat. Isotopes 26.(1973).17-20.
18. Weinreich, R., Qaim, S.M., Michael, H. and Stoecklin, G. J. Radioanal. Chem. 30(1976).53.
19. Clark, J.C., Thakur, M.L. and Watson, I.A. Int. J. Appl. Radiat. Isotopes 23.(1972).329.
20. Thakur, M.L., Nunn, A.D. and Waters, S.L. Int. J. Appl. Radiat. Isotopes 22(1971).481.
21. Greene, W.M., Lebowitz, E., Richards, P. and Hillman, M. Int. J. Appl. Radiat. Isotopes 21.(1970).719.
22. Dahl, J.R. and Tilbury, R.S. Int. J. Appl. Radiat. Isotopes 23.(1972).431.
23. Saha, G.B. and Farrer, P.A. Int. J. Appl. Radiat. Isotopes 22.(1971).495.
24. Suzuki, K. and Iwata, R. Int. J. Appl. Radiat. Isotopes 28.(1977) .663-665.
25. Thakur, M.L. and Nunn, A.D. Radiochem. Radioanal. Lett. 2.(1969).301.

26. Silvester, D.J. and Thakur, M.L. *Int. J. Appl. Radiat. Isotopes* 21.(1970).630.
27. Hupf, H.B. and Beaver, J.E. *J. Appl. Radiat. Isotopes* 21.(1970).75.
28. Porter, J., Kawana, M., Krizek, H., Lathrop, K.A. and Harper, P.V. *J. Nucl. Med.* 11.(1970).352.
29. Neirinckx, R.D. *Int. J. Appl. Radiat. Isotopes* 27.(1976).1.
30. Helus, F. *Radiochem. Radioanal. Lett.* 3.(1970).45.
31. Nunn, A.D. and Waters, S.L. *Int. J. Appl. Radiat. Isotopes* 26.(1975).731.
32. Helus, F., Maier-Borst, W., Lambrecht, R.M. and Wolf, A.P. *Proc. 7th Int. Conf. on Cyclotrons and their Applications* (Birkhaeuser, Basel,1975).474-477.
33. Qaim, S.M., Stoecklin, G., and Weinreich, R. *Int. J. Appl. Radiat. Isotopes* 28.(1977).947-953.
34. Glass, H.I., Arnot, R.N., Clark, J.C. and Allan, R.N. "Cerebral Blood Flow", Springer Verlag, Heidelberg (1969).p.63.
35. Clark, J.C., Jones, T. and Mackintosh. "Radioaktiv Isotope in Klinik und Forschung", Urban and Schwarzenberg, Muenchen (1970), p.444.
36. Yano, Y., McRae, J. and Anger, H.O. *J. Nucl. Med.* 11.(1970).674.
37. Watson, I.A. *Radiochem. Radioanal. Lett.* 4.(1970).7.
38. Moghissi, A.A. and Hupf, H.B. *Int. J. Appl. Radiat. Isotopes* 22.(1971).218.
39. Hurby, J.J., in the "Uses of Cyclotrons in Chemistry, Metallurgy and Biology", Ed. Amphlett, C.B., Butterworths, London (1970). p.149.
40. Steyn, J., Myer, B.R. and Barendsma, J.M.J. *Int. J. Appl. Radiat. Isotopes* 22.(1971).55.
41. Thakur, M.L. and Nunn, A.D. *Int. J. Appl. Radiat. Isotopes* 23.(1972).139.
42. Brown, L.C. and Beets, A.L. *Int. J. Appl. Radiat. Isotopes* 23.(1972).57.
43. MacDonald, N.S., Neely, H.H., Wood, R.A., Takahashi, J.M., Wakakuwa, S.I. and Birdsall, R.L. *Int. J. Appl. Radiat. Isotopes* 26.(1975).631.
44. Silvester, D.J., Sugden, J. and Watson, I.A. *Radiochem. Radioanal. Lett.* 2.(1969).17.
45. Sodd, V.J., Blue, J.W. and Scholz, K.L., in the "Uses of Cyclotrons in Chemistry, Metallurgy and Biology", Ed. Amphlett, C.B., Butterworths, London (1970),p.125.
46. Hupf, H.B., Eldridge, J.S. and Beaver, J.E. *Int. J. Appl. Radiat. Isotopes* 19.(1968).345.
47. Cauwe, F., Deutsch, J.P., Favarat, D., Prieels, P. and Cogneau, M. *Int. J. Appl. Radiat. Isotopes* 25.(1974).187.
48. Lambrecht, R.M. and Wolf, A.P. *Radiat. Res.* 52.(1972).32.
49. Lebowitz, E., Greene, M.W. and Richards, P. *Int. J. Appl. Radiat. Isotopes* 21.(1970).489.
50. Guillume, M., Lambrecht, R.M. and Wolf, A.P. *Int. J. Appl. Radiat. Isotopes* 26.(1975).703.
51. Lindner, L. Iodine - 123 in Western Europe, *Juel. Conf.* 20 (1976).139-144.
52. Cunninghame, J.G., Morris, B., Nichols, A.L. and Taylor, N.K. Iodine - 123 in Western Europe, *Juel. Conf.* 20.(1976).145-153.
53. Loepfe, E. Iodine - 123 in Western Europe, *Juel. Conf.* 20. (1976).137-138.
54. Weinreich, R. Iodine - 123 in Western Europe, *Juel. Conf.* 20. (1976).49-69.
55. Watson, I.A. and Tilbury, R.S. *J. Nucl. Med.*11.(1970).373.
56. Jones, T., Clark, J.C., Kocak, N., Cox, A.G. and Glass, H.I. *Brit. J. Radio.* 43.(1970).537.

57. Yano, Y., van Dyke, D., Budinger, T., Anger, H.O. and Chu, P.
J. Nucl. Med. 11.(1970).663.
58. Sodd, V.J., Blue, J.W. and Scholz, K.L. Phys. Med. Biol.
16.(1971).587.
59. Lebowitz, E. and Greene, M.W. Int. J. Appl. Radiat. Isotopes
22.(1971).789.
60. Yano, Y., van Dyke, D.C., Verdon, T.A. and Anger, H.O.
J. Nucl. Med. 12.(1971).815.
61. Thakur, M.L. and Waters, S.L., private communications.
62. Wilkins, P.E., Beach, L.A. and Marlow, K.W. Radiochim. Acta
17.(1972).110.
63. Stark, V.J., Harper, P.V., Lathrop, K.A., Krizek, H., Rowed, D.W.,
Lembares, N. and Hoffer, P.B. J. Nucl. Med. 13.(1972).468.
64. Lebowitz, E., Greene, M.W., Kinsley, M. and Richards, P.
J. Nucl. Med. 12.(1971).376.
65. Brown, L.C. and Callahan, A.P. Int. J. Appl. Radiat. Isotopes
26.(1975).213.
66. Leibowitz, E., Greene, M.W., Fairchild, R., Bradley-Moors, P.R.,
Atkins, H.L., Ansari, A.N., Richardson, P. and Belgrave, E.
J. Nucl. Med. 16.(1975).151-155.
67. Comar, D. and Crouzel, C. C.E.A. Bull. Inform. Sci. Tech. No. 220.
(1976).9-18.
