

NEUTRON ACTIVATION ANALYSIS

D. De Soete R. Gijbels J. Hoste

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Neutron Activation Analysis

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PREFACE

Neutron activation analysis is a technique applied in many different fields of research and technology such as, for example, biology, geochemistry, solid state physics or criminology. It was the author's aim therefore to give sufficient coverage of the underlying principles and practical problems to enable non-specialists in radiochemistry and analytical chemistry to apply activation analysis in their own field of interest. Consequently, fairly detailed treatment is given to such topics as cross sections, neutron fluxes, neutron sources, growth and decay of radionuclides, counting techniques, chemical separations, systematic errors, sampling problems, etc. Purely instrumental techniques, including computer handling of data and statistical interpretation of the results, are also covered. Tables provide the necessary data on thermal cross sections, resonance integrals, fission neutron cross sections, 14 MeV neutron cross sections, gamma ray energies of nuclides formed by neutron activation and gamma ray attenuation coefficients. A selection of methods is not presented, as the choice depends not only on the specific problem under investigation but also on the neutron source and detection equipment available. Instead, there is a compilation of the literature amounting to over 3,000 references, referring to both trace element and matrix.

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| Figure 6.37,
6.38 | Palms, J. M. <i>et al.</i> , <i>IEEE Trans Nucl. Sci.</i> , N5-15 (3), 397 (1968), The Institute of Electrical and Electronics Engineers. |
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| Figures 7.3, 7.4 | Diebolt, J., <i>Contribution à l'Analyse par Activation des Gazes Rares</i> , Thesis, University of Grenoble (France), 1963. |
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| Figure 8.6 | Girardi, F. and Sabbioni, E., <i>J. Radioanal. Chem.</i> , 1 , 176 (1968), Elsevier. |
| Figures 8.15, 8.17 | Samsahl, K. <i>et al.</i> , <i>Anal. Chem.</i> , 40 , 183 (1968), American Chemical Society. |

- Figure 8.19 Jervis, R. E. and Wong, K. Y., *Nuclear Activation Techniques in the Life Sciences*, p. 137, International Atomic Energy Agency, Vienna, 1967.
- Figure 8.20 Van den Winkel, P. *et al.*, *Nuclear Activation Techniques in the Life Sciences*, p. 159, International Atomic Energy Agency, Vienna, 1967.
- Figure 8.21 Tera, F. and Morrison, G. H., *Anal. Chem.*, **38**, 960 (1966), American Chemical Society.
- Figure 8.23 Op de Beeck J., *Anal. Chim. Acta*, **40**, 224 (1968), Elsevier.
- Figure 8.24 Op de Beeck, J. and Hoste, J., *Proceedings of the Analytical Chemical Conference at Budapest*, 1966. Vol. II, p. 256, Institute for General and Analytical Chemistry, Technical University, Budapest.
- Figure 8.25 Meinke, W. W., *Modern Trends in Activation Analysis, Proceedings of International Conference, Texas A. & M. University* 1961, p. 37.
- Figure 8.26, 8.27 Girardi, F. *et al.*, *Radiochemical Methods of Analysis*, vol. II p. 5-9, International Atomic Energy Agency, Vienna, 1965.
- Figure 8.28 Samsahl, K., *Anal. Chem.*, **39**, 1481 (1967), American Chemical Society.
- Figure 8.29 Samsahl, K., *The Analyst*, **93**, 102 (1968), Society for Analytical Chemistry.
- Figure 9.1 Aubouin, G., *Radiochim. Acta*, **1**, 122 (1963), Akademische Verlagsgesellschaft, Frankfurt/Main.
- Figure 9.2 Borg, D. C. *et al.*, *Int. J. Appl. Radiation and Isotopes*, **11**, 21 (1961), Pergamon.
- Figures 9.3, 9.4 Wölfe, R. *et al.*, *Z. anal. Chem.*, **233**, 224, 244 (1968), Springer-Verlag, Heidelberg.
- Figure 9.5 Van Grieken, R. *et al.*, *Anal. Chim. Acta*, **43**, 200 (1968), Elsevier.
- Figure 9.6 De Neve, R. *et al.*, *Anal. Chim. Acta*, **40**, 380 (1968), Elsevier.
- Figure 9.13 Heath, R. L., *Scintillation Spectrometry Gamma-ray Spectrum Catalogue*, AEC Report IDO-16880-1, vol. I (1964).
- Figure 9.14 Gardner, D. G. *et al.*, *J. Chem. Phys.*, **31**, 983 (1959), American Institute of Physics.
- Figure 9.15 Isenhour, T. L. *et al.*, *Modern Trends in Activation Analysis, Proceedings of International Conference, Texas A. & M. University*, 1965, p. 127.

- Figure 10.11 Junod, E., Report CEA-R 2980, vol. I (A and B), 1966, Commissariat à l'Energie Atomique, France.
- Figure 11.1 Kim, J. I. *et al.*, *Anal. Chim. Acta*, **33**, 129 (1965), Elsevier.
- Figure 11.4 Heath, R. L., *Semiconductor Nuclear-Particle Detectors and Circuits*, Publication 1593, Committee on Nuclear Science, National Academy of Sciences, National Research Council, Washington, D.C. (1969), p. 656.
- Figure 11.6 Kawashima, T., *Radiochim. Acta*, **4**, 110 (1965), Akademische Verlagsgesellschaft, Frankfurt/Main.
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- Table 3.4 Zijp, W. L., RCN-40, 1965, Stichting Reactor Centrum Nederland.
- Table 6.2 Schulze, W., *Neutronenaktivierung als Analytisches Hilfsmittel*, 1962, F. Enke Verlag, Stuttgart.
- Table 6.7 Adams, F. and Dams, R., *J. Radioanal. Chem.*, **3**, 123 (1969), Elsevier.
- Table 7.5 Gleit, C. E. and Holland, W. D., *Anal. Chem.*, **34**, 1456 (1962), American Chemical Society.
- Table 8.5 Bowen, H. J. M., and Gibbons, D., *Radioactivation Analysis*, 1963, Clarendon Press, Oxford.
- Table 9.1 Wölfe, R. *et al.*, *Z. anal. Chem.*, **233**, 248 (1968), Springer-Verlag, Heidelberg.
- Table 9.6 Isenhour, T. L. *et al.*, *Modern Trends in Activation Analysis, Proceedings of International Conference, Texas A. & M. University*, 1965, p. 126.
- Appendix 5, Tables 1 and 2 Dams, R. and Adams, F., *Radiochim. Acta*, **10**, 5, 8 (1968), Akademische Verlagsgesellschaft, Frankfurt/Main.

PHYSICAL CONSTANTS

<i>Symbol</i>	<i>Denomination</i>	<i>Magnitude</i>
a.m.u.	atomic mass unit	931.48 MeV (¹² C scale) 931.16 MeV (¹⁶ O scale)
<i>c</i>	light velocity in vacuum	$2.99793 \cdot 10^{10}$ cm s ⁻¹
<i>e</i>	electron charge	$4.803 \cdot 10^{-10}$ esu
<i>h</i>	Planck's constant	$6.625 \cdot 10^{-27}$ erg s
$\hbar$	$= h/2\pi$	$1.0544 \cdot 10^{-27}$ erg s
<i>k</i>	Boltzmann's constant	$1.38047 \cdot 10^{-16}$ erg deg ⁻¹
<i>m_n</i>	neutron mass	1.008665 a.m.u. (¹² C scale) $1.6757 \cdot 10^{-24}$ g
<i>m₀</i>	electron rest mass	$5.486 \cdot 10^{-4}$ a.m.u. (¹² C scale) $9.107 \cdot 10^{-28}$ g
<i>m₀c²</i>	electron rest energy	0.5110 MeV
<i>m_p</i>	proton rest mass	1.007273 (¹² C scale) $1.6724 \cdot 10^{-24}$ g
<i>N_A</i>	Avogadro's number	$6.023 \cdot 10^{23}$ atoms mole ⁻¹ (¹² C scale)

LIST OF SYMBOLS

<i>Symbol</i>	<i>Denomination</i>	<i>Symbol</i>	<i>Denomination</i>
α	—alpha particle		—number of disintegrations
	—conversion coefficient		—distance
	—quotient of distribution coefficients	ϵ	energy needed for a hole-electron pair creation (eV)
	—correction factor in the internal standard method	ϵ_x	excitation energy of nucleus x (MeV)
A	—mass number	E	energy (MeV, keV, eV, erg, . . .)
	—atomic weight	E_b	Coulomb barrier energy (MeV)
	—activity (rate or number of disintegrations, — counts)	E_{ca}	cadmium cut-off energy (eV)
	—ampere	E_{eff}	effective threshold energy (MeV)
$\text{\AA}$	Ångstrom (10^{-8} cm)	E_{max}	maximum energy in beta spectrum (MeV)
β	beta particle	E_0	energy corresponding to neutron velocity v_0 (0.025 eV)
β^-	negatron	E_r	energy at the maximum of a resonance peak (eV)
β^+	positron	E_T	threshold energy (MeV)
B	background	E_x	recoil energy of nucleus x (MeV)
b	barn $\doteq$ unit of cross section (cm^2)	eV	electron volt ($1.60 \cdot 10^{-12}$ erg)
c	number of counts	φ	neutron flux (beam or multidirectional) ($\text{n cm}^{-2} \text{s}^{-1}$)
cm	centimeter (10^{-2} m)	$\bar{\varphi}$	equivalent fission flux ($\text{n cm}^{-2} \text{s}^{-1}$)
C	—signal + background	$\varphi(E)$	φ at neutron energy E ($\text{n cm}^{-2} \text{s}^{-1}$)
	—capacity (Farad)	φ_e	epicadmium neutron flux per unit of lethargy = conventional epicadmium flux ($\text{n cm}^{-2} \text{s}^{-1}$)
	—compound nucleus	$\varphi_e(E)$	φ_e at neutron energy E ($\text{n cm}^{-2} \text{s}^{-1}$)
Ci	Curie ($3.7 \cdot 10^{10}$ dps)		
C_i	convolution integer		
CR	cadmium ratio		
CR_x	cadmium ratio of element (isotope) x		
CT	clock time		
δ	—density (g cm^{-3})		
	—residual standard deviation		
D	disintegration rate (dps, . . .)		
D_A	distribution coefficient for species A		
DT	total dead time		
d	—absorption thickness (mg cm^{-2})		

<i>Symbol</i>	<i>Denomination</i>	<i>Symbol</i>	<i>Denomination</i>
φ_s^{eff}	φ_s corrected for resonance absorption at finite dilution ($n \text{ cm}^{-2} \text{ s}^{-1}$)	f_s	self-absorption coefficient for beta rays
φ_0	conventional thermal neutron flux = $n v_0$ ($n \text{ cm}^{-2} \text{ s}^{-1}$)	ft	comparative half-life of beta decay
$\varphi_{reactor}$	conventional reactor neutron flux ($n \text{ cm}^{-2} \text{ s}^{-1}$)	f_{th}	thermal neutron absorption factor in solids
φ_{th}	conventional thermal neutron flux below E_{Cd} = $n_{th} v_0$ ($n \text{ cm}^{-2} \text{ s}^{-1}$)	Γ	nuclear level width (eV)
$\varphi_{th}(E)$	thermal neutron flux at energy E ($n \text{ cm}^{-2} \text{ s}^{-1}$)	Γ_s	partial Γ (eV)
F	-14 MeV neutron flux ($n \text{ cm}^{-2} \text{ s}^{-1}$)	γ	gamma ray
	-Fano factor	g	gram
	-decay correction factor for measurement of short lived activities (starting point of measuring interval)	h	-photopeak height (cm, activity) -height -hour
FDT	fractional dead time	I	resonance integral at infinite dilution (barn)
F_n	fraction of the disintegration of the nuclides of the n^{th} step, producing nuclides of the $(n+1)^{\text{th}}$ step in a disintegration chain	I'	I corrected for $1/v$ contribution (barn)
		I_{abs}	absorption resonance integral at infinite dilution (barn)
		I_{act}	activation resonance integral at infinite dilution (barn)
		IDT	instantaneous dead time
		I^{eff}	effective resonance integral = I corrected for resonance absorption = I at finite dilution (barn)
FWHM	full width at half maximum of a (photo)peak (eV, keV. . .)	I_n	I for the n^{th} resonance peak (barn)
f	-total neutron absorption factor in solids	I_{tot}	= $\sum^n I_n$ (barn)
	-decay correction factor for the measurement of short lived activities (exact time within the measuring interval)	I_x	I for the nuclear reaction of type x (barn)
	-fission	I'_x	I' for the nuclear reaction of type x (barn)
	-fraction	$I_{1/v}$	resonance integral at infinite dilution obtained by integration of $\sigma_{1/v}$ (barn)
f'	total neutron absorption factor in solution	κ	-linear absorption coefficient for pair production (cm^{-1}) -dielectric constant
$f(E)$	fission neutron flux at energy E ($n \text{ cm}^{-2} \text{ s}^{-1}$)		
f_s	epicadmium neutron absorption factor in solids		

<i>Symbol</i>	<i>Denomination</i>	<i>Symbol</i>	<i>Denomination</i>
K_D	distribution constant	mCi	milli Curie (10^{-3} Curie)
k	-reactor reproduction factor	mg	milligram (10^{-3} gram)
-	-constant	ml	milliliter = 10^{-3} liter
keV	kilo electron volt (10^3 eV)	m μ	milli micron = 10^{-7} cm (10^{-3} micron)
kg	kilogram (10^3 g)	mm	millimeter (10^{-3} m)
λ	-radioactive decay constant (s^{-1})	ms	milli second (10^{-3} second)
	-wave length (m μ , Å)	mV	-milli electron volt (10^{-3} eV)
l	mean free path (cm)		-millivolt (10^{-3} volt)
L	-liter		-frequency (s^{-1})
	-ligand		-neutrino
L_c	critical limit		-number of neutrons liberated per fission
L_D	detection limit	N	-number of neutrons in the nucleus
L_q	quantitative determination limit		-number of target nuclei per cm ³
LT	live time		-normality (g eq. L ⁻¹)
μ	-total mass absorption coefficient (cm ² mg ⁻¹)	n	-thermal neutron density from energy 0 to ∞ (n cm ⁻³)
	-micron (10^{-4} cm)		-neutron
μ'	total linear absorption coefficient (cm ⁻¹)	$n(E)$	neutron density at energy E (n cm ⁻³)
μA	micro ampere (10^{-6} A)	ng	nano gram (10^{-9} gram)
μb	microbarn (10^{-6} barn)	ns	nano second (10^{-9} second)
μCi	micro Curie (10^{-6} Curie)	n_{th}	thermal neutron density below E_{cd} (n cm ⁻³)
μg	micro gram (10^{-6} gram)	$\bar{n}_{th}$	average n_{th} in the sample at finite dilution (n cm ⁻³)
μl	micro liter (10^{-6} liter)	$n(v)$	neutron density at velocity v (n cm ⁻³)
μ_n	mobility of electrons in n -type semiconductor material (cm ² V ⁻¹ s ⁻¹)	P	-probability
			-peak to total ratio
μ_p	mobility of holes in p -type semiconductor material (cm ² V ⁻¹ s ⁻¹)	p	proton
		Q	-reaction energy (MeV, a.m.u., . . .)
μs	micro second (10^{-6} second)		-quality criterion of a counter
M	-molarity (mole L ⁻¹)	q	branching factor
	-factor of merit of a counter = $S/2\sqrt{B}$	ρ	resistivity (ohm cm ⁻¹)
MeV	million electron volt (10^6 eV)	R	-reaction rate (s^{-1})
m	-minute		count rate (cps, cpm, . . .)
	-mass		
	-meter		
mA	milli ampere		
meq	milli equivalent		
mb	millibarn (10^{-3} barn)		

<i>Symbol</i>	<i>Denomination</i>	<i>Symbol</i>	<i>Denomination</i>
	-particle range	σ_c	cross section for compound nucleus formation (= σ_{abs}) (barn)
	-resolution	σ_{coh}	coherent scatter cross section (barn)
	-radius	σ_{col}	collision cross section (barn)
	-distance	$\sigma(D)$	standard deviation for a signal equal to the detection limit
	-resistance (ohm)	$\sigma(E)$	cross section at neutron energy E (barn)
R_A	-nuclear radius (Fermi unit = 10^{-13} cm)	σ_{eff}	effective cross section (barn)
	-recovery factor of species A	σ_{el}	elastic scatter cross section (barn)
$R(V, E)$	response function of a detector	σ_f	fractional standard deviation (= coefficient of variation) = $\sigma\%/100$
r	-distance	σ_{fa}	free atom scatter cross section (barn)
	-radius	σ_{inel}	inelastic scatter cross section (barn)
Σ	macroscopic cross section = σN (cm^{-1})	$\sigma_{14 \text{ MeV}}$	cross section for 14 MeV neutrons (barn)
Σ_R	macroscopic removal cross section of sample for 14 MeV neutrons (cm^{-1})	$\sigma_{n.e.}$	non elastic scatter cross section (barn)
$\Sigma_{R(i)}$	macroscopic removal cross section of element i for 14 MeV neutrons (cm^{-1})	$\sigma_{n,x}$	reaction cross section (barn)
σ	-effective microscopic target area (cm^2)	$\sigma(0)$	standard deviation of background for zero signal
	-cross section (barn)	σ_0	cross section at neutron velocity v_0 (barn)
	-linear absorption coefficient for Compton effect (cm^{-1})	$\bar{\sigma}_0$	average elemental cross section at neutron velocity v_0 (barn)
	-standard deviation for an infinite population	σ_p	potential scattering cross section (= $4\pi R_A^2$) (barn)
$\bar{\sigma}$	average cross section in a fission neutron spectrum (barn)	$\sigma(Q)$	standard deviation for a signal at the quantitative determination limit
$\sigma\%$	percentage standard deviation (= percentage coefficient of variation)	σ_r	resonance cross section (barn)
σ_{abs}	absorption cross section (= σ_c) (barn)		
σ_{act}	isotopic activation cross section (barn)		
$\bar{\sigma}_{act}$	average elemental activation cross section (barn)		

<i>Symbol</i>	<i>Denomination</i>	<i>Symbol</i>	<i>Denomination</i>
$\sigma_r(E)$	resonance cross section at neutron energy E (barn)	$T_{1/2}$	-total time = $\Delta t_c + \Delta t_s$ half-life = $\ln 2/\lambda$ (y, h, m, s, . . .)
σ_{reactor}	cross section for a reactor neutron spectrum (barn)	T_m	Maxwellian temperature ($^{\circ}\text{K}$)
$\sigma_{\text{reactor}}^{\text{eff}}$	effective $\sigma_{\text{reactor}} = \sigma_{\text{reactor}}$ corrected for absorption at finite dilution (barn)	T_0	293.6 $^{\circ}\text{K}$
$\sigma_{R(t)}$	microscopic elemental removal cross section for 14 MeV neutrons (cm^2)	T_R	reactor period
$\bar{\sigma}_s$	average scatter cross section (barn)	t	-decay time (h, m, s, . . .)
σ_T	total cross section (barn)	$t_{1/2}$	-thickness (cm)
σ_{th}	average cross section for neutron energies up to E_{Cd} (barn)		-temperature ($^{\circ}\text{C}$)
$\sigma(v)$	cross section at neutron velocity v (barn)		time necessary to establish half of the equilibrium distribution in isotopic exchange irradiation time (h, m, s, . . .)
$\sigma(\bar{v})$	cross section at neutron velocity $\bar{v}$ (barn)	t_b	effective thickness for resonance neutron absorption (cm)
$\sigma_{1/v}$	cross section in the epicalm region, disregarding resonance peaks (barn)	t_{eff}	number of standard deviations
S	-saturation factor -surface, area -signal	V	-volume (L, ml, μl , . . .)
$S_{B/A}$	separation factor of species B from A = enrichment factor of A = depletion factor for B	$V(d)$	-tension (volt)
s	-standard deviation for a finite population -second	v	-volt
θ	-isotopic abundance -angle (degree, radian)	$\bar{v}$	electrical field
τ	-mean life (s, . . .) -dead time (μs , . . .) -resolving time (μs , . . .) -linear absorption coefficient for photoelectric effect (cm^{-1})	v_1	-neutron velocity (cm s^{-1})
T	-absolute temperature ($^{\circ}\text{K}$)	v_0	average neutron velocity in a Maxwell-Boltzmann distribution (cm s^{-1})
			neutron velocity at E_{Cd} (cm s^{-1})
			most probable neutron velocity in a Maxwell-Boltzmann distribution (2200m/s at 20°C = 0.025 eV)
		W_i	statistical weight
		w	weight (g, mg, . . .)
		y	-year
		ζ	-fission yield
		Z	Fermi potential in a semiconductor (eV)
		z	-atomic number
			-residual counting efficiency

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