DEEP PENETRATION OF NEUTRONS IN HYDROGENOUS SHIELDS

by

B. CHINAGLIA and G. BOSIO

(SORIN)

(Topical Report)

1972
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The experimental results refer to:

a) Al-27 (n, α) reaction rate in water, measured up to 2.1 m from the Avogadro RSI reactor core;

b) several detector reaction rates covering the energy range from thermal to the Al-27 (n, α) threshold in an iron-water lamination up to 1.5 m from a disk source (EURACOS facility of CCR Euratom Ispra).
The investigated thicknesses are greater than those previously reported in the literature for similar media and calculation errors which might be relevant for reactor shield design can be detected by comparison with the present experiment. The experimental results are therefore given in full detail together with the source description to permit their use for meaningful tests of shielding codes.

As examples of such tests the following results have been obtained for the shielding codes QAD and SABINE:

- the Moment Method data for water above ~ 6 MeV calculated up to 1.6 m (water density 1 g/cm$^2$) reported in ORNL 3487 are confirmed within 10%; exponential extrapolation of the attenuation kernel beyond this distance yields a slight under-estimate of the flux at 2.1 m. When the neutron dose is considered it is expected that the QAD predictions obtained using the exponential fitting of the above data are somewhat lower than the true values at this distance, but the discrepancy should not exceed a factor 2;

- the SABINE thermal flux is in excellent agreement with the experimental data in the iron-water mixture. Fast fluxes are somewhat underestimated beyond 1 m in iron-water and, to a greater extent, in plain water. This effect seems to derive from the removal cross sections of water and a small decrease in their values yields reasonable agreement with the experimental data.

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ABSTRACT

The penetration of fission neutrons in hydrogenous shields has been measured by means of activation detectors in well defined geometrical conditions.

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KEYWORDS

NEUTRONS
ATTENUATION
SHIELDING MATERIALS
WATER
IRON
MIXTURER
MEASURED VALUES
ACTIVATION
RADIATION DETECTORS
RADIATION SOURCES
DISTANCE
COMPUTER CALCULATIONS
ERRORS
Index *)

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*) Manuscript received on July 20, 1972
1. Introduction

It is well known that rather large errors may affect the prediction of the neutron flux far from the source (more than $\sim 10$ relaxation lengths). The failure of the diffusion theory for shield calculation is the major example evidenced since long time, but even more exact solutions are based on, or imply, assumptions and approximations which may be uncorrect at deep penetration (for instance the number of angular interval divisions in a $S_N$ code).

In any case, whatever computational method is used, the answer is dependent on the accuracy of basic data (cross section) and their average values within finite energy intervals. The error deriving from uncorrect cross section sets increases with the distance (for a pure exponential attenuation the relative error is proportional to the optical path) and can reach high values for real reactor shields.

The various effects due to the method, mesh intervals, cross sections are correlated and the basis for assessing the validity of a code is an experiment in known conditions.

This work has the scope to give some deep penetration results obtained during an experimental study of the neutron attenuation in water and iron-water mixtures. These media have been extensively investigated in the past with the main purpose to determine the detailed space-energy distribution in the first part of a shield and most of the published data* refer to penetration depths ranging from 50 to 100 cm.

* A survey of experimental results may be found in Ref.1.
Such depths however are well below the total thickness of a shield. On the other hand in hydrogenous media the absorption and energy degradation process is such that the attenuation of the total flux is controlled by a fast neutron component whose mean energy is increasing with the distance.

Therefore the flux behaviour at deep penetration could not be extrapolated from the short distance data, where the most energetic neutrons have a minor importance.

This work was originated for a particular problem and some results which refer to the comparison between experiment and calculation with the Shielding codes Sabine (7) and QAD-P5 (8) are presented. The main emphasis is however given to the experimental results and the source and shield characteristics are described in full detail to permit the use of the data for other tests.

2. Experiments

2.1 - The plain water medium.

For the plain water medium the source is the core of the Avogadro RS1 swimming pool reactor and the measurements have been made on vertical and horizontal lines passing through the center of the core.

The core consist of MTR type fuel elements having 19 plates (10 plates for control elements) of Al/U alloy 0,127 cm thick separated by 0,287 cm of water as shown in fig.1. For the source description the active zone of each element may be considered a homogeneous mixture containing the following volu-
The power distribution is known from thermal flux measurements. Fig. 2 shows the reactor loadings n.271 I and n.279 I which have been used for the present experiments; the numbers in each box representing an element are proportional to the thermal flux at the center of the element (between the two central plates at half the active length). The vertical profile is nearly identical for all the elements and a mean value is given in fig. 3a. The position of the control rods during the experiments is also shown in fig. 2.

The neutron attenuation has been measured by the use of aluminium foils with the Al$^{27}$ (n, α) reaction, along a vertical line passing through the water hole of the reactor loading n.271 I and a horizontal line for reactor loading n.279 I as shown in fig. 2.

The choice of the detector has been dictated by several reasons:

- A high threshold detector is not sensitive to photoneutrons produced in water. The photoneutron flux is known to overcome the flux of fission born neutrons at low (thermal) energies beyond a distance of 2m in water, and the experimental results
would be of doubtful usefulness at this penetration in the case of low energy detectors.

- The low density of aluminum and the high γ energy of the reaction product Na$^{24}$ make possible the use of rather thick detectors without strong self-absorption during the activity counting. The detection limits can therefore be lowered by the use of large size detectors.

- The half period of 15 hours permits to reach easily activity values near to saturation and to make use of the decay after irradiation for lowering the very high counting rate ratio ($\sim 10^{12}$) between the most and less active detectors.

- The behaviour of high energy neutrons is particularly important and the Al$^{27}$ (n, α) reaction may provide a direct test of the energetic component of the spectrum which controls the deep penetration of neutrons.

High purity (> 99,999) aluminum disks of various dimensions (diameter: 1.5 to 7 cm, thickness 0.1 to 0.2 cm) have been used as detectors. During irradiation they were wrapped in aluminum foil inside cadmium boxes to reduce surface contamination and activation of impurities from thermal neutrons. Furthermore after irradiation each detector was etched in NaOH solution to eliminate any surface contamination. The activity measurement has been performed with calibrated scintillation detectors (3"x3" NaI (Tl) crystals) at various source-detector distances using appropriate correction factors for the finite size of the disks. Further details of the experimental procedure are given in Ref.2.

The results are given in table I and also plotted in fig.4. The reactor power for the vertical axis, reactor loading n.271 I, was 5 Mw and for the horizontal axis, reactor loading n.279 I, 7 Mw.
In Ref. 2 and 3 an analysis of the errors has been made, and the following values have been obtained:

a) Counting statistics: only the less active detectors are affected by bad counting statistics. The standard deviation is given in table 1.

b) Reactor power: 10%.

c) Distance from the core: the position of the detectors with respect to the supporting frame is known with good precision, so that the distance between the various detectors has a negligible error; the position of the frame with respect to the source is defined with smaller precision. For the vertical axis measurements the uncertainty of the fuel length within the elements leads to an error of $\pm 1$ cm, which corresponds to about 10% on the activity of detectors far from the source. For the horizontal axis some uncertainty arises from the curvature of fuel plates and the resulting error on the activity is $\pm 3\%$.

d) Perturbation due to the supporting frame: a negligible perturbation is evaluated.

e) Spurious contribution from activation of impurities (taking into account the contribution of photoneutrons to the low energy flux): less than $10^{-3}$ for the more distant detector.

f) Contribution to Al$^{27}$ (n, a) reaction from photoneutrons: absent (an upper limit of $5 \cdot 10^{-4}$ is evaluated with largely conservative assumptions).

g) Absolute calibration of $\gamma$ counter and correction factors: $\leq 5\%$

The resulting standard deviation is about 12% for horizontal and 15% for vertical axis measurements and it must be observed
that only the statical error (a), which is negligible up to about 1.5 m, affects the shape of the curve of fig.4.

2.2 - Iron - water lamination.

The source is the EURACOS facility of CCR-Euratom Ispra which is described in detail in Ref.4. Its essential features are:
- disk source, 0.8 m diameter, 90% enriched Uranium-235;
- well defined neutron spectrum;
- power: (1.25 ± 0.10) Kw.

The experimental shield configuration is an array of iron, water and air slabs as shown in fig.5. The iron slabs (transversal dimensions 1.45 m x 1.45 m) are immersed in water inside an iron walled tank which during irradiation fills the void of the concrete tunnel of the EURACOS facility, as shown in the insert of fig.6.

Streaming along the unavoidable gap between tank and concrete walls was greatly reduced by the bends of the tank walls. A streaming component was however detected as shown later and gave rise to a spurious contribution to the measurements at the end of the tank.

Starting from the source, the sequence of materials along the axis of fig.5 is given by the following table:

<table>
<thead>
<tr>
<th>M</th>
<th>air</th>
<th>Fe</th>
<th>H₂O</th>
<th>Fe</th>
<th>H₂O</th>
<th>Fe</th>
<th>H₂O</th>
<th>Fe</th>
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<td>1</td>
<td>11</td>
<td>3</td>
<td>3.5</td>
<td>8</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>t</td>
<td>11.9</td>
<td>3.2</td>
<td>32.5</td>
<td>13</td>
<td>10</td>
<td>13</td>
<td>22.4</td>
<td>1</td>
</tr>
</tbody>
</table>

M : material

t : thickness of material M (in cm).
Reaction rates of various detectors were measured along the axis of the shield by irradiating foils or pellets fixed to lucite holders (in water) or to iron plugs (inside the iron slabs). The reactions and the physical form of the detectors are given below:

\[ ^{164}\text{Dy} (n, \gamma) : \text{Al/Dy alloy (10\% or 30\%); disks.} \]

\[ ^{197}\text{Au} (n, \gamma) : \text{pure gold or Al/Au 0.1\% alloy; disks.} \]

\[ ^{115}\text{In} (n,n') : \text{pure In; disks.} \]

\[ ^{32}\text{S} (n, p) : \text{pure S; fused pellets.} \]

\[ ^{27}\text{Al} (n, \alpha) : \text{pure Al; disks.} \]

All the above detectors, except Dy, were irradiated under cadmium; the Cd thickness used for Au was 1 mm (cut-off at \( \sim 0.44\) eV). Absolute saturation activities per gram element were obtained by means of calibrated counters. Further details on the experimental procedure are given in Ref. 5.

The results are given in Table II and fig. 7 as a function of the distance \( z \) measured from the inner face of the tank wall, expressed in reaction rates (R.R.) for In, S, Al detectors. For Dy the data have been converted to thermal flux \( \Phi_{th} \) (the conversion is based on intercalibration of Dy and Au detectors in a slowing down spectrum in water, using \( \sigma_{Au} = 98\) b) and from Au/Cd the epithermal flux per unit lethargy \( \Phi_{u} \) has been calculated, using a value 1550 b for the resonance integral.

In addition to the statistical error \( \varepsilon \) reported in Table 2, the values are affected by uncertainty in the source power (8\%), position (\( \pm 0.1\) cm, which for fast neutron detectors is equivalent to \( \sim 3\% \)) and calibration of the counters (\( \pm 5\% \)).

Near the end of the tank the data have a spurious contribution due to neutron streaming along the gaps; fig. 6 shows the
epithermal streaming flux measured around the tank. A correction has been made for this effect assuming that the streaming flux entering the end wall is attenuated exponentially in water with a relaxation length of 10 cm, as shown in fig.7. The corrected values are given in Table 2.

3. Calculation

Some calculation results obtained with QAD and Sabine are given in figs. 4 and 7. Before discussing the comparison with the experimental data, it is useful to describe briefly the geometry and cross section sets used as inputs to the codes.

3.1 - Geometry.

The simple disk geometry of the Euracos source was described exactly in both codes. The Avogadro reactor core source was approximated by one or more regions having power distributions obtained by averaging the values of fig. 2 along the appropriate axis as described below.

**Reactor loading n.271-I.**

QAD: a single homogeneous source region was considered, bounded by planes:

\[
\begin{align*}
  x &= 0 & x &= 50,7 \text{ cm} \\
  y &= 0 & y &= 46,8 \text{ cm} \\
  z &= 0 & z &= 68,4 \text{ cm}
\end{align*}
\]

where \( z \) is a vertical axis and the origin is taken at the left lower corner of the core shown in fig.2.

In this rather coarse description the water hole C4 is considered a source element.

**Sabine**: no calculation has been made.
Reactor loading n.279-I.

The greater accuracy reached in the measurements along the horizontal axis suggested to use a more accurate description of the source for this case.

QAD: the core was divided in 3 regions:

1) rows 2, 3, 4, 5 power 5,344 Mw
2) " 6 " 0,929 Mw
3) " 7 " 0,727 Mw

The boundary planes are shown in fig.8b and the power distributions are given in fig.9.

Sabine: the core was divided in 2 cylindrical regions as shown in fig.8C. The power distribution along the z axis and the radial direction are given in fig.3a and 3b.

3.2 - Cross sections.

In QAD the attenuation kernel in water is fitted by $\ e^{-\ A_1 d_i}$ where $d$ is the water thickness. The coefficients $A_i$ used for the present calculation for the various energy groups are given in Table III.

Three different removal cross section sets for water have been used for the Sabine calculation, they are shown in Table IV.

The activation cross section of the threshold detectors were derived from the best literature data.

The average values within the QAD groups are given in Table V and refer to a pure fission spectrum.

3.3 - Boundary conditions.

For the Sabine code the following boundary conditions have been used:

reactor: a value derived from in core measurements and a cal-
culated spectrum\(^{(9)}\) has been assigned to every group flux at the inner boundary. Transversal buckling:
\[ B^2 = 3.6 \cdot 10^{-3} \text{ cm}^{-2} \text{ (from measurements).} \]

**Euracos:** \( J^+ = 0 \) at the inner boundary for the first and the thermal group, for the other groups calculation with \( J^+ = 0 \) and \( J^+ > 0 \) \( (\Phi / \Phi') = (2+i) D \) for \( i \geq 23; i = \text{group index} \) gave similar results except near the source. At the outer boundary: zero incoming current.

4. **Discussion and conclusion**

The data reported in the previous sections have been obtained in well defined conditions and can therefore be used for meaningful tests of calculation methods. The investigated thickness (2.1 m in water, 1.40 m effective thickness in iron-water) is sufficiently large to put in evidence errors which might be relevant in reactor shield calculation. Some implications concerning the examined codes are given in the following.

4.1 - Water.

The comparison based on the \( \text{Al (n, a)} \) reaction in water shows that the calculated data are somewhat lower than the experimental ones at deep penetration.

4.1.1 - QAD.

The QAD results are examined firstly. Two geometry errors affect the data at short distances:
- the first is the homogenization in reactor loading 271-I;
  a rough evaluation indicates that, if in reactor loading n.271-I the water hole is taken properly into account, the predictions near the core are to be lowered by about 30%;
- the second effect is characteristic of the integration
method, which for any volume element of the source approximates the mean value of the attenuation kernel with the value calculated at the geometrical center of the element itself. This error depends on the distance and on the size of the volume elements and in the present case the flux should not be overestimated by more than 5% at large distance (above 1 m).

An error derives from the use of constant cross sections based on a fission spectrum as used in the QAD groups. A correction factor may be calculated by weighing the cross sections over the Moment Method spectrum(6). This error is however of little importance (below 10% in the range of distances investigated).

The values of the fitting coefficients $A_i$ play a more important role. When the fluxes given by these coefficients are compared with the data of ref.6, it is found that the fitting is rather poor at deep penetration; furthermore beyond 1,6 m no data are available and one must extrapolate the exponential behaviour of the last fitted zone. This procedure is certainly not conservative.

A correction factor has been calculated for taking into account the two effects of cross section and incorrect fitting. This factor is the ratio of the $A_1(n, a)$ reaction rate calculated at various distances $d$ in water from a point source with the exact detailed shape of Ref.6 to that given by QAD for the same conditions. As a result, the QAD data of fig.4 should be multiplied by:

$$
0.85 \quad \text{for } z \sim 30 \text{ cm}
$$

$$
1.10 \quad " \quad 70 \text{ cm}
$$

$$
1.30 \quad " \quad 140 \text{ cm}
$$

where the hypothesis has been made that the correction factor
evaluated at a distance d from a point source applies to a distance \( \zeta = (d+20) \text{ cm} \) for the core source.

The above corrections bring in excellent agreement experimental data and QAD predictions indicating that above 6 Mev the Moment Method results of Ref. 6 are confirmed by the present experiments within 10%. The use of a constant cross section and of the fitting coefficients of Table III does not introduce large errors and even at about 2 m the ratio experimental over predicted reaction rate is lower than 2.

These findings indicate that at lower energies one should expect a better agreement or even conservative predictions, since the fluxes obtained with the coefficients \( A_i \) below 6 Mev are in much better agreement or overestimate those of the Moment Method.

4.1.2 - SABINE

A preliminary observation concerns the procedure followed to obtain the threshold detector reaction rates. Mean group activation cross sections are calculated for each group using the removal flux as weighing function. In the case of \( {^{27}\text{Al}}(n,\alpha) \) reaction and water medium this procedure leads to a slight overestimate of the reaction rate since within the first group the total and removal flux spectra are quite different, the latter being strongly peaked at high energy.

This effect can be evaluated on the basis of the more realistic spectrum shape given by the Moment Method calculation\(^{(6)}\).

Assuming that the spectrum at a distance \( \zeta \) from the core has the same shape calculated at a distance \( \zeta + 20 \text{ cm} \) from a point source, the Sabine predictions of fig. 4 should be lowered by:
and therefore the discrepancies with the experimental points should be slightly increased.

The present comparison indicates that the removal cross section sets B and C of table IV are more adequate than set A for the description of the high energy neutron flux in water. This conclusion could be extrapolated to lower energy neutrons, since according to the calculation scheme the flux at any energy in water follows the removal flux attenuation far from the source.

4.2 - Iron - water lamination.

4.2.1.- QAD

As well known, when a shield is composed by one or more materials in addition to the reference material, use is made in the QAD code of an equivalent thickness

\[ t'_M = t_M \frac{\Sigma_M}{\Sigma_0} \]

where:

- \( t_M \) = thickness of material \( M \)
- \( \Sigma_M \) = removal cross section of material \( M \)
- \( \Sigma_0 \) = removal cross section of the reference material.

In the present case the following values have been used:

- \( \Sigma_0 = 0.101 \text{ cm}^{-1} \) (water)
- \( \Sigma_M = 0.168 \) " (iron)
The comparison with experimental data has been limited to threshold detector reaction rates, calculated with activation group cross sections averaged over a fission spectrum.

An excellent agreement is found everywhere (see fig. 7) for high energy neutrons (above ~3 MeV, sulphur and aluminum data) and also for lower energy neutrons (above ~1 MeV, indium data) far from the iron slabs, as expected from the application of a removal cross section.

4.2.2. - Sabine

The calculation shown in fig. 7 has been made with the removal cross section set A of table IV.

The tendency to underestimate the fluxes is again observed, although the discrepancies with the experimental data are lower than in the previous experiment.

Threshold detector and epicadmium gold activation are somewhat underestimated whilst the thermal flux closely fits the experimental points (except at the interior of the iron slabs) over the whole shield.

A discussion of these findings is outside the scope of this work; it may be observed that a better agreement would be obtained by the use of the water removal cross sections B and C of table IV and that in a lamination where the water thickness is comparable to the iron thickness and does not exceed 30 cm the thermal diffusing flux never attains equilibrium with the removal flux and therefore in this range its attenuation does not depend strongly on the removal neutron attenuation.
Acknowledgments.

Part of this work has been performed at the Euratom CCR of Ispra and the authors wish to thank dr. Nicks, Perlini, and Ponti of the Euratom staff for assistance during the experiments and calculations.

REFERENCES


7) C. Ponti, H. Preusch and H. Schubart "SABINE, a one dimensional bulk shielding program" Eur 3636 e (1967).


9) M. Iman Atom Kernenergie 12-7 p. 35 (1967).
FIGURES:

1/ Fuel elements (standard and control) of Avogadro RS1 reactor.

2/ Scheme of the reactor loadings used for the water experiments. Numbers inside elements are proportional to the thermal flux values.

3/ Vertical and radial power distribution (average values).

4/ Attenuation of the Al$^{27}$ (n,α) reaction rate in water.

5/ Scheme of the experimental set-up used for iron-water experiments.

6/ Neutron (fast and epithermal) streaming around the gap shown in the insert.

7/ Attenuation of thermal, epithermal fluxes and threshold reaction rates in an iron-water mixture.

8/ a) Scheme of the core
    b) Scheme of the Geometry used for QAD
    c) Scheme of the Geometry used for SABINE.

9/ Power distribution used for QAD.
### TABLE I

**Al \((n,\alpha)\) reaction rates \((s^{-1}g^{-1})\)**

Reactor loading

<table>
<thead>
<tr>
<th></th>
<th>271-I; 5 Mw</th>
<th></th>
<th>279-I; 7 Mw</th>
<th></th>
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<tr>
<td>(z) (cm)</td>
<td>R.R. ((-1s^{-1}g^{-1}))</td>
<td>(\varepsilon^*)</td>
<td>(z) (cm)</td>
<td>R.R. ((-1s^{-1}g^{-1}))</td>
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<td>8.80/7**</td>
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<td>2</td>
<td>2.85/8</td>
</tr>
<tr>
<td>11</td>
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<td>8</td>
<td>8,7</td>
<td>8.68/7</td>
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<tr>
<td>41</td>
<td>4.17/5</td>
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<td>3.93/7</td>
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<td>4.40/3</td>
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<td>7.1 /1</td>
<td>&lt;5</td>
<td>23,7</td>
<td>9.76/6</td>
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<td>4.2 /0</td>
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<td>28,7</td>
<td>5.04/6</td>
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<td>60</td>
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\(\varepsilon^* = \) Statistical counting error

\((**): n/x = n \cdot 10^x\)
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<th>Au(n, y) /Cd</th>
<th>In(n, y)</th>
<th>S (n, p)</th>
<th>Al (n a)</th>
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<td>R (s^-1 g^-1)</td>
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**TABLE - II**

Euracos results (thermal flux and reaction rates). Values with * are corrected for streaming; n/x = n.10^*.
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<th>Energy (MeV)</th>
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<th>$A_1$</th>
<th>$A_2$</th>
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<td>-6.7966/2</td>
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<tr>
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<td>-9.6949/2</td>
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**TABLE - IV**

Water removal cross sections used for SABINE

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TABLE - V

Activation cross sections averaged over QAD groups

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<th>Group</th>
<th>Energy (MeV)</th>
<th>In(n,n') ($\text{cm}^2 \cdot \text{g}^{-1}$)</th>
<th>S(n,p) ($\text{cm}^2 \cdot \text{g}^{-1}$)</th>
<th>Al(n, ) ($\text{cm}^2 \cdot \text{g}^{-1}$)</th>
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<td>1,18/-3</td>
<td>6,20/-3</td>
<td>2,169/-3</td>
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<td>9-7</td>
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<td>5,97/-3</td>
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<tr>
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<td>5,49/-3</td>
<td>1,043/-4</td>
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<td>4,63/-3</td>
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<td>1,65/-3</td>
<td>4,48/-3</td>
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<td>1,91/-3</td>
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<tr>
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<td>9</td>
<td>0,835-0,5</td>
<td>6,84/-4</td>
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<td>0,5-0,1</td>
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Fig. 1 - Fuel elements (standard and control) of Avogadro RS1 reactor.
Fig. 2 - Scheme of the reactor loading used for the water experiments. Numbers inside elements are proportional to the thermal flux values.
Fig. 3 - Vertical and radial power distribution (average values).
Fig. 4 - Attenuation of the $\text{Al}^{27} (n, \alpha)$ reaction rate in water.
Fig. 5 - Scheme of the experimental set-up used for iron-water experiments.
Fig. 6 - Neutron (fast and epithermal) streaming around the gap shown in the insert.
Fig. 7 - Attenuation of thermal, epithermal fluxes and threshold reaction rates in an iron-water mixture.
Fig 8 - a) Scheme of the core
b) Scheme of the Geometry used for QAD
c) Scheme of the Geometry used for SARTNE
Fig. 9 - Power distribution used for QAD.
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