UNITED STATES ATOMIC ENERGY COMMISSION

YIELD OF PHOTONEUTRONS FROM U^{235} FISSION PRODUCTS IN HEAVY WATER

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FINAL REPORT

YIELD OF PHOTONEUTRONS FROM U235 FISSION PRODUCTS IN HEAVY WATER

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with Appendix

NEUTRON YIELD FROM MULTIPLY SCATTERED COMPTON PHOTONS

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ABSTRACT

The half-lives and yields of the photoneutrons created in heavy water by U^{235} fission product gamma rays have been measured. Seven half-lives have been found: 2.5 sec., 41.2 sec., 2.38 min., 7.7 min., 27.3 min., 1.65 hr., 4.37 hr., and 53 hr. The shortest one and longest one of these are least reliable. Eighty-five percent of the photoneutrons appear in the two shortest half-lives, the 2.5 sec. component being three times as intense as the 41 sec. component. The total saturated activity of the photoneutrons for an infinite amount of heavy water was calculated from measurements with a 10" radius sphere to be about 16.5% of the saturated delayed neutron activity. The data indicate that there must be of the order of two gamma rays, of energy above 2.17 Mev, emitted per fission by fission products with half-lives longer than one sec.

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I. Introduction

In a pile consisting of uranium and heavy water, gamma rays of
energy greater than 2.17 Mev given off by the fission products will
create neutrons in the heavy water by photo-disintegration of the deuteron.
These photoneutrons will have the same periods as the gamma rays which
create them. They will, therefore, act in all respects like the delayed
neutrons themselves, and may exert an appreciable effect on the kinetic
behavior of the pile. In order to estimate the effect of the photo-
neutrons upon pile periods, their number as a function of time must be
known, as in the case of the delayed neutrons. The purpose of this ex-
periment was to measure the number of photoneutrons created in heavy
water by U^{235} fission products as a function of the length of time after
bombardment. Since the photoneutrons are, for short times after bombard-
ment, accompanied by delayed neutrons, the yields were measured relative
to those of the delayed neutrons. For this purpose, the yields of delayed
neutrons as given by Hughes, Dabbs and Cahn in CP-3094 were used. The re-
sults of CP-3094 have been collected in Table I for convenience.

II. Apparatus

A schematic diagram of the apparatus is shown in Fig. 1. It con-
sisted of a ten inch radius, 1/8 inch wall, aluminum sphere immersed in
a large tank of oil. A "rabbit" containing a sample of enriched uranium
oxide could be transferred from the center of the pile to the center of the
sphere in about 0.25 sec. by means of a pneumatic tube. The sphere could
be filled or emptied of heavy water in about one minute by means of a
siphon arrangement.

The neutron distribution in the oil was studied by means of fission
chambers, each containing about 64 mg of 90% enriched uranium. Fission
chambers were chosen in preference to BF_3 counters because of their greater
insensitivity to the large gamma ray "background" being emitted by the
sample. The ion chamber could be placed at chosen reproducible radial
positions in the tank by means of a track and carriage arrangement.

Since some of the periods involved are fractions of a sec., it was
necessary to keep the initial counting rates quite high in order to obtain
good statistics throughout the run. To keep corrections for missed counts
low, the fastest most readily available amplifier was used. A modified
Simpson proportional counter amplifier and scaling circuit were adapted
and found to have a resolving time of about 20 microseconds. This re-
solving time was determined by exposing a sample of uranium first at high
pile power, then at low power, and plotting the ratio of the activities
as a function of time as described in CP-3094. All of the data were then
corrected for missed counts using this over-all experimentally determined
"resolving time".

Fission pulses from the amplifier were fed into a scale of 256
The data was recorded by photographing the scaling-circuit interpolation
bulbs with an open shutter oscilloscope camera. The film moved past the
neon bulbs in such a manner that each bulb appeared as a dashed line on
the film for the period the bulb was lit. A time signal was also impressed
on the film every 0.1 sec by means of a neon bulb and a revolving shutter
driven by a synchronous motor. The length of the exposure was recorded
automatically on the film by another neon bulb which lighted at the start
and went out at the end, by connections with the electropneumatic control
switches. A reproduction of one of the films is shown on Fig. 2.

III. Method

The method employed to measure the source activity is based on the
following theory. Consider a point source of Q neutrons/sec, having an
arbitrary energy distribution (above thermal energy), surrounded by an
oil bath large enough to absorb all the neutrons. It is known that most
of the absorption is due to hydrogen nuclei in the oil, and that the
H-capture cross section is very small above thermal energy. Then the
rate of production of fast neutrons must equal the rate at which they
are captured, or $Q = \int (nv)_t \, \sigma_c \, dV$, where $(nv)_t$ is the flux of thermal
neutrons at any point in the oil. σ_c is the macroscopic cross section
for capture at thermal energy, and the integral is extended over the oil
volume. Measurements with a fission chamber detector give a counting
rate "Y" at any point in the oil which is proportional to $(nv)_t$, provided

correction is made for the epicadmium counting rate. In the case of spherical symmetry, $\psi \propto \int Y\, R^2\, dR$ In our case, since there is negligible neutron absorption in the heavy water sphere, the integral is taken from R_0 to ∞, where R_0 is the sphere radius.

The ion chamber was fixed in a given radial position, an exposure was made, and the resulting neutron counting rate recorded on the moving film as a function of time "t" after the end of the exposure. When the activity had decreased to an acceptable value, the ion chamber was moved to the next radial position and another exposure made with the same sample. A single series usually comprised exposures in four to eight positions. A complete experiment required three complete series: (a) with heavy water in the sphere, in which case both delayed and photoneutrons are present; (b) with the sphere empty, so that the activity is that due to delayed neutrons alone (c) with the sphere empty and the fission chamber surrounded by cadmium, in order to measure the delayed neutrons with epi-cadmium energies. With heavy water in the sphere the number of epi-cadmium neutrons was negligible.

The constancy of the pile power level and the detecting instruments was checked by repeating the initial exposure at the end of each experiment. Instrument sensitivity alone was checked by means of a Ra-α-Be source. Variations remained within one percent, which simplified the experiment by making it unnecessary to use an exposure monitor.

For each exposure, a plot was made from the film data of the
counting rate at the particular radial position as a function of "t".
From the group of curves forming a series, other curves were constructed
of (counting rate x R^2) vs R, for successive fixed values of "t". Here,
R is the radial distance of the detector from the source at the center of
the sphere. The source activity at any time is then proportional to the
area under the corresponding curve.

Examples of these (counting rate x R^2) vs R curves are shown in Fig.
3. With heavy water in the sphere, the curves on a semi-log scale are
linear with a relaxation length (inverse slope) which is constant and
equal to approximately 1.25 inches. This indicates that the neutrons
have been well thermalized before they escape into the surrounding oil.
When the sphere is empty, the semi-log plot of the neutron distribution
shows a decided initial curvature due to slowing down in the oil.

In general, the desired source activity due to photoneutrons alone,
at a time "t_1", is the difference between the areas under the correspond-
ing distribution curves, with and without heavy water, after correction
for epicadmium activity. By computing a number of points, a curve may
be drawn of source photoneutron activity as a function of time. We assume
that the data can be represented by a formula of the form (1), where A_p
is the photoneutron activity of the source as a function of time "t"
after the end of the exposure.

$$A_p(t) = \sum_i a_i(1 - e^{-T/\tau_i}) e^{-t/\tau_i} \qquad (1)$$

T the exposure time, T_i the mean life, and a_i the yield of the

i'th radioactive period for infinite exposure time. If the experimen-

tal curve is linear on a semi-log plot at sufficiently large values of

"t", the longest half-life period can be determined. The half-lives

of shorter periods are evaluated by the well-known "peeling off" process.

A similar formula for $A_d(t)$, the delayed neutron activity, can be fitted

to the volume integral curve from the data taken with the sphere empty.

By evaluation at $t = \bullet$, $T = \infty$, we can then express the yields a_i of

each photoneutron period as fractions of the total calculated delayed

neutron yield $A_d(t = \bullet, T = \infty)$.

At small values of "t" the photoneutron activity of the source

is as little as one-tenth of the delayed neutron activity. That is, the

difference of the areas under two curves of the type shown in Fig. 3,

with and without heavy water, is only '0% of either area. Thus an error

of 1 percent in either area causes a 10% error in the result, and it is

desirable to obtain the maximum number of experimental points in order to

increase accuracy. At larger values of "t", however, the delayed neutron

activity becomes a small and finally vanishing fraction of that due to

the photoneutrons, which contain several components of comparatively long

mean life. In this case, a sufficiently accurate value of the volume

integral of the counting rate with heavy water can be obtained from the

counting rate at a single radial position, as is clear from Fig. 3.

Since the slope is approximately constant and known, a single point de-

termines the area under a curve, where "t" must be extended to many

hours, this procedure results in a considerable saving of time.

IV. Experimental Results

(A) Periods and Uncorrected Yields

Fig. 4 is a plot of the photoneutron activity, measured at a single radial position with D_2O in the sphere, as a function of time. The sample was 2.7 gm of roughly 60% 25 exposed in the pile for 3.75 hours. Within the interval 4 hours to 26 hours, the gross activity, Curve #1, is well represented by three periods of half-life 53 hr., 4.37 hr., and 1.65 hr. The longest period may be considerably in error because it was followed over a range of only 2 in activity, and the activity at the end was only a few times background. The other two periods were followed over a factor of 100, and should be accurate. The relative yields were computed by extrapolation of the individual terms to zero time and infinite exposure.

Fig. 5 illustrates a 30 min exposure of a 0.7 gm sample, also measured at a fixed radial position. Curve #1 is the gross activity. A formula representing the three previously determined photoneutron periods was fitted to Curve #1 at t = 6.5 hr. by the adjustment of a constant multiplier, which allows for the different sample size and pile power level. Curve #2 is the result of subtracting from #1 these three longer periods. It indicates a fourth period of 27.3 minutes half-life, with signs of a still shorter period at t < 1 hr.

Fig. 6 illustrates a 5 min exposure of a .7 gm sample. Curve #1 is the crude data. Curve #2 is the result of subtracting off the four

previously determined periods by means of a formula adjusted to fit at
t = 129 min. A fifth period of half-life 7.7 min is demonstrated.

Thus far we have determined the yields of 5 photoneutron periods
relative to each other. In the next experiment, the yields are related
to that of the delayed neutrons. Fig. 7 shows results of a 3 min ex-
posure with a 0.7 gm sample, from t = 3 min to 1 hour. In the first part
of this interval the delayed neutron activity is not negligible. It was
measured in the following manner. A series of 7 exposures was made, with-
out D_2O in the sphere, varying the radial distance R of the detector from
the source and measuring with a mechanical recorder the number of counts
in the interval t = 2.5 to t = 4.0 min. This time integral of the de-
layed neutrons was plotted as a (counting rate x R^2) vs R curve, the
area N under the curve being proportional to the time and volume integral
of the delayed neutron activity of the source. A six term formula similar
to Eq. (1) was set up for $A_d(t)$, using the constants for the delayed
neutron periods as given in Table I, and integrated from t = 2.5 to 4.0
min. Letting $N = C_1 \int A_d(t)dt$, the constant C_1 was determined for this
experiment. $C_1 A_d(t)$ then represents the volume integrated delayed
neutron activity as a function of time.

Curve #1 of Fig. 7 shows the gross source activity with D_2O in the
sphere. This was actually computed from the counting rate measured at
a single radial position, an approximation previously discussed. If we
subtract from this the delayed neutron activity $C_1 A_d(t)$, computed as above,
the result is Curve #2, representing photoneutrons alone. Curve #3 is the
result of subtracting off the 5 previously determined photoneutron periods,
fitting the formula at t = 53 min. A sixth period is indicated of half-
life 2.38 min.

Since we ha.e a formula representing the volume integrated activity of the delayed neutrons and one for 6 photoneutron periods, we can now extrapolate to t = o and infinite exposure time and determine the photo-neutron yields relative to the delayed neutrons. Table II lists these six periods with their yields A relative to the yield of the 22 sec delayed neutron period as 1.00.

In the next experiment, no short cuts were permissible because of the need for accuracy. Fig. 8, Curve #1, shows the gross integrated activity, with D_2O in the sphere, resulting from a 30 sec exposure of a 0.15 gm en-riched sample. Each point on the curve represents an integral of (counting rate) x R^2 vs R, taken from curves of the type shown in Fig. 3. Curve #2 shows, for the same exposure, the volume integrated activity without D_2O, due to delayed neutrons alone. Curve #2 has been corrected for epi-cadmium activity, which amounted to about 2%. Curve #3, Fig. 8, shows the total photoneutron activity, Curve #1 minus #2. It is seen that the difference is only 10% of the total activity at t = 30 sec.

The delayed neutron formula was fitted to the experimental Curve #2, and the saturated delayed neutron activity computed for this experiment. From previous results, a formula expressing the photoneutron activity could be set up for the 2.4 min and longer periods. Subtracting these off from Curve #3, the result was Curve #4, which represents a new period of half-life 41 sec and yield A = 0.090 (see Table II). There is evidence for a still shorter period below t = 30 sec., but in this experiment the counting rate became so high in this region that the data are not reliable.

A series of 30 sec exposures was made with a 1.5 gm sample of un-
enriched uranium. Curve #1, Fig. 9, is the total integrated activity
with D_2O. Curve #2, that without. Curve #3 their difference. The de-
layed neutron formula, using the values in Table I, was fitted to Curve
#2 at t = 10 sec. It is of interest as a check on Table I that the formula
then agreed with the experimental results to 1% or better from t = .5 sec
to t = 12 sec and with the time integrated activity from t = 30 to t = 50
sec.

A formula for the photoneutron activity due to the 7 previously de-
termined periods was obtained by fitting it to the observed time integrated
activity in the interval t = 30 to t = 50 sec. This was subtracted from
Curve #3, Fig. 9, to yield the new period, Curve #4, of half-life 2.5 sec
and A = 0.225 (See Table II).

(B) Estimated Accuracy

The accuracy to be expected in obtaining one of the areas under
curves of the type in Fig. 3 is about one percent, due to statistical
probable errors alone. Since the short period photoneutron yields were
obtained as the difference of two areas, amounting to about 10 percent
of either, these yields are uncertain by at least 10 percent. The yields
of the longer periods, except for the 53 hr period, should be somewhat
more accurate.

If the heavy water was 99.87 percent pure, as stated by the manu-
facturer, the absorption of neutrons in it would have been negligible.

Since contamination might have been introduced during the course of the experiment, the neutron absorption of the heavy water was tested in the following manner: An Sb-Be photoneutron source was placed at the center of the sphere and the neutron distribution curves in the oil were obtained, with and without heavy water. Since the γ-rays from Sb are less than 2.17 mev, they cannot create any photoneutrons in the heavy water, and the areas under the (counting rate) x R^2 vs R curves should be equal in the two cases. Actually the areas were found to be the same within about 2 percent.

(C) Corrections to Experimental Yields

In Table II, Column A represents photoneutron yields (saturated activities) for our sphere of heavy water relative to the 22 sec delayed neutron period yield as 1.00. In order to correct these yields to the case of an infinitely large sphere of D_2O and complete γ-ray absorption, it is necessary to know the energy of the γ-rays. Conclusions about the initial energy of the photoneutrons could not be made from the data of this experiment. The large amount of heavy water moderator slowed the neutrons down to such an extent that the shape of the distribution curve in the oil where the measurements were taken was no longer a measure of the initial neutron energy. Hence, we have tried to correlate our periods with those reported by Hughes, Spaatz and Cahn in CP-3472, April 25, 1946. Table III shows this correlation. Although the agreement in the half-lives is only fair, it is sufficient to set up a correspondence between

our periods and theirs. Accordingly, we have used their values for the γ-ray energies. In the case of the two longest periods, as shown in Table II, we have assumed entirely arbitrarily an energy $\bar{E}_\gamma = 3$ mev for purposes of computation.

In D_2O and for γ-ray energies between 2 and 5 mev the photoelectric effect and pair production are negligible in comparison with the Compton effect. If we assume that every Compton scattering of a γ-ray quantum by an electron reduces the quantum energy below the γ.n threshold in D_2O, the fraction of the quanta absorbed in a sphere of radius R cm, from a central point source, is $F_t = 1 - e^{-(\mu_s + \mu_\gamma)R}$ where μ_s is the linear "absorption" coefficient/cm for Compton scattering and μ_γ is the coefficient for the γ.n reaction in D_2O. For this experiment the effective thickness of heavy water is 23.3 cm. $\mu_s = \sigma_s \times 3.35 \times 10^{23}/cm^3$ for D_2O, where σ_s, the Compton scattering cross section/electron varies from 1.27×10^{25} cm^2 at 2.5 mev to 1.06×10^{-25} at 3.5 mev. $\mu_\gamma = 6.7 \times 10^{22} \times \sigma_\gamma/cm^3$, where σ_γ, the γ.n cross section/deuteron varies from zero at 2.17 mev to about 14×10^{-28} cm^2 at 3.5 mev. In Table II are listed values of $1/F_t$ the correction factor by which the yields A must be multiplied to give the yields for an infinite sphere of D_2O, for the assumed energies $\bar{E}_\gamma$.

A small correction, $\alpha = 1.06$, is listed in the next column of Table II. This compensates for the γ-ray absorption in the 0.050 inch wall thickness of the pneumatic tube, the fixed 0.125 inch thick Al tube

through the center of the sphere, and in the 3/16 inch thick wall of the bakelite rabbit.

Column β in Table II gives an additional correction necessary because not every Compton scattering process reduces the energy of a γ-ray quantum below the γ,n threshold, so that the flux of γ-rays of energy above the threshold is everywhere higher than we would calculate using the full Compton scattering cross section. β/F_t is then the ratio of the number of γ,n neutrons produced in an infinite D_2O sphere to that produced in a sphere of radius 23.3 cm. taking into account multiple scattering. The theory of this correction, developed by Soodak and Greuling, is given in Appendix A.

Column Ac in Table II is A x $\propto \beta/F_t$, the corrected yield of photoneutrons in an infinite sphere of D_2O. Column A_d is 100 Ac/4.54, the percent yield relative to the total number of delayed neutrons (See Table I, total of Col. A). Column A_f is 2.43 x 0.00755 A /100, the absolute yield of photoneutrons per fission, taking ν (25) = 2.43 and the delayed neutron yield from Table I.

(D) Gamma Ray Yields

We now compute the number of γ-rays per fission necessary to produce the photoneutrons of each period, with yields A_f in Table II. The fraction of the total γ-ray quanta from a point source which are absorbed in the γ,n reaction in an infinite D_2O sphere is

$$F_\gamma = \int_0^\infty e^{-(\mu_s + \mu_\gamma)R} \mu_\gamma \, dR = \frac{\mu_\gamma}{\mu_\gamma + \mu_s}$$

provided each scattering process is assumed to reduce the energy be-
low the threshold. If we take into account the fact that some scattered
quanta still have energy above the threshold, the actual photoneutron
production will be higher by a factor E, which will be small near the
threshold and increase with the initial energy of the γ-ray. In Table
II are listed, for each period, the values of $1/F_{\gamma}$ and of E. The ab-
solute yield of γ-ray quanta per fission, A_{γ}, will be

$$A_{\gamma} = A_f/F_{\gamma}E ,$$

V   Discussion of Results

(A) Summary

A summary of the results obtained is given in Table II. An explana-
tion of the meaning of each column is given with the table. The re-
sults may be summarised as follows:

(1) The photoneutron periods observed show half-lives from 2.5 sec to
53 hrs.

(2) Eighty-five percent of the photoneutrons appear in the shortest
half-lives of 2.5 sec and 30 sec, the 2.5 sec half-life component be-
ing three times as intense as the 30 sec half-life.

(3) The total saturated activity of the photoneutrons for an infinite
amount of heavy water is about 16% of the saturated delayed neutron
activity.

(B)   The Effect of Possible Parent-Daughter Contributions

It should be pointed out that in the analysis of the data as des-
cribed above it has been assumed that the hard gamma ray emitters are
initial fission fragments. The interpretation of the data is made some-
what uncertain because of the possibility that the hard photon giving
the photoneutron may come from a daughter of the initial fragment. In
such cases the data may yield the true period, but the amplitude deduced
by extrapolating the decay curve back to zero time will be in error.
If the daughter has a very much longer life than its parent, the error
will not be large. If the two have comparable lifetimes the saturated
activities given may be a considerable overestimate. If both the parent
and daughter give hard gamma rays, both amplitudes will be in error, but
the sum of the two exponential terms will give a correct description of
the activity as a function of time.

(C)   Number of Photons Per Fission

From the number of photoneutrons per fission, the number of photons
per fission was calculated from considerations of the relative proba-
bilities of Compton and γ,n collisions. The results for each period are
given in the column A_{γ} of Table II. The sum of the values in this column
is 2.3 photons per fission. For the sixth period listed, Hughes, Spaatz
and Cahn give two values for the γ-ray energy, one from photoneutron energy
measurements (2.25 mev) and one from direct γ-ray absorption measurements
(2.4 mev). The γ,n cross section is changing very rapidly in this region,
so that the calculated number of photons per fission is very sensitive to

which of these two values is used. Even if the value giving the lesser number of photons per fission is used, the total number of photons per fission adds up to 1.7. As a total, this number may not be unreasonable. However, the 0.62 photons per fission given for the single 2.5 sec period is much too high to be due to a single isotope. The same conclusion could be made also for the 41 sec and the 2.4 min periods, since the highest fission yields observed are about six percent. These considerations suggest that there may be many fission products having periods of several seconds to several minutes, with γ-rays above 2.2 mev. If close together, such periods could not be resolved in this experiment.

If the suggestion that 25 may have many short period γ-rays is true, and since these short periods have been shown to account for 90% of the photoneutrons, then the results of this report might be used to speculate about photoneutron yield of 23, for which there are no observations available at the present time, by the following considerations: The 2.5 sec and 41 sec periods, which account for 90% of the photoneutrons, give about 1 photon per fission. Since maximum fission product yields are about 6%, there are about 17 fission chains contributing the hard γ-rays in the case of 25 if we assume one photon per fission chain. The effect of one of these is, then, at most 5 to 10% and the shift in the maximum of the fission product yield curve of a few mass number units, in going from 25 to 23, is not likely to have any considerable effect. The number of photoneutrons from 23 might, therefore, be expected to be roughly the same as for 25. The number of delayed neutrons

for 23 has been reported to be about one-third that for 25 (CP-3147).
Consequently, it seems that for 23 the number of photoneutrons may be
comparable to the number of delayed neutrons.

(D) Comparison with CP-3472

This work was done simultaneously with that of Hughes, Spaatz and
Cahn at the Argonne Laboratory (CP-3472). Our results are compared with
theirs in Table III. The agreement between the periods is fair, con-
sidering the large number of delayed and photoneutron periods involved.
The main disagreement in the periods is that our data showed a 4.4 hr
period not reported by them. It seems very probable that really accurate
determination of the periods, especially the shorter ones, would neces-
sitate chemical separation of individual fission products, or groups of
products. The greatest disagreement on yield values lies in the region
where the measurements are made difficult because the photoneutrons
represent only a small part of all the neutrons present. The 30 sec
period of CP-3472 is three times as intense as our 41 sec period. The
sum of the values listed in Column A of Table II gives the result that
the total saturated photoneutron activity is 16% of the saturated delay-
ed neutron activity. CP-3472 gives 28% for this ratio.

T A B L E   I

Delayed Neutrons from 25

τ	$\tau_{\frac{1}{2}}$	E	A	B
80.2	55.6	.25	0.153	2.5×10^{-4}
31.8	22.0	.56	1.00	16.6
6.50	4.51	.45	1.28	21.3
2.19	1.52	.62	1.45	24.1
0.62	0.43	.42	0.51	8.5
0.072	0.06		0.15	2.5
Totals			4.54	75.5×10^{-4}

τ = period, $\tau_{\frac{1}{2}}$ = half life, E = energy in mev.

A = saturated yield relative to the yield of the 22 sec period as 1.00

B = absolute yield relative to the total number of neutrons emitted per fission

Data from CP-3094 (Hughes, Dabbs & Cahn 7-30-45) and CF-3408 (Hall, Dec. 1945).

Table II

Photoneutron Yields and Periods

Ref. No.	$\tau_{\frac{1}{2}}$	τ	A	E_γ	$1/F_t$	α	β	A_c	% A_d	A_f x 10^4	$1/F_\gamma$	B	A_γ
1	53h	76h	.00058	(3)	1.695	1.06	1.085	.00074	.0165	.0298	476	1.14	.00125
2	4.37h	6.3h	.00119	(3)	1.695	1.06	1.085	.00232	.0515	.0942	476	1.14	.00394
3	1.65h	2.4h	.0093	2.62	1.62	1.06	1.052	.0166	.370	.676	570	1.09	.0416
4	27.3m	39.3m	.0082	2.66	1.63	1.06	1.056	.0149	.328	.600	643	1.09	.0554
5	7.7m	11.1m	.0124	3.0	1.695	1.06	1.085	.0242	.554	.975	476	1.14	.0407
6	2.4m	3.44m	.0276	2.65	1.63	1.06	1.064	.0604	1.11	2.03	660	1.09	.121
7	41s	59.5s	.090	2.25	1.52	1.06	1.009	.147	5.24	5.93	2450	1.015	1.43
8	2.5s	3.6s	.226	3.4	1.77	1.06	1.11	.469	10.31	18.9	589	1.18	.623

Total 15.96% 29.2x10⁻⁴ 2.3

$\tau_{\frac{1}{2}}$ = half life. τ = mean life

A = uncorrected photoneutron yield, relative to a yield of 1.00 for the delayed neutron 22 sec period

E_γ = γ-ray energies, in mev. from CP-3472. Values in parentheses are arbitrary

α = Correction for γ-ray absorption in the rabbit wall and in the Al tubes at the center of the sphere

β = Correction for multiple scattering (see Appendix A)

A_c = A x α x β/F_t, the corrected yield in an infinite D_2O sphere

A_d = 100 A_c/4.54 = percent yield of photoneutrons relative to total delayed neutrons

A_f = 2.43 x .00755 x A_d/100 = absolute photoneutron yield per fission

$\frac{1}{F_\gamma}$ = Number of photons per photoneutron in ∞ D_2O on the basis that single Compton collision knocks γ-energy below threshold

B = Correction factor to account for γ's not knocked below threshold on single collision

A_γ = Absolute number of photons per fission

TABLE III

Comparison of Results, This Paper & CP-3472

	This Paper			Hughes, Spaatz & Cahn			
Ref. No.	$T_{\frac{1}{2}}$	A_c		$T_{\frac{1}{2}}$	A_c	E_γ	
1	53 h	.00074		24 h	.00032	-	
2	4.37 h	.00232		-		-	
3	1.65 h	.0168		2.0 h	.0091	2.62	
4	27.3 m	.0149		32 m	.0118	2.66	
5	7.7 m	.0242		6.5 m	.0210	3.0	
6	2.4 m	.0504		90 s	.0720	2.65	
7	41 s	.147		30 s	.456	2.25	
8	2.5 s	.469		6.7 s	.655	3.4	

$T_{\frac{1}{2}}$ = half-life

A_c = corrected yield, infinite D_2O absorber, relative to the yield
of the 22 sec delayed neutron period as 1.00

E_γ = γ-ray energy in mev

APPENDIX A

Neutron Yield from Multiply Scattered Compton Photons

E. Greuling

The neutron yield resulting from the photodisintegration of deuterium in a sphere of heavy water can be calculated by making use of the following simple assumptions:

(a) Every Compton scattered photon of incident energy greater than 2 mev suffers negligible change in direction. Actually 4 mev photons are scattered by less than 30° if their resulting energy is above the γ,n threshold, 2.2 mev.

(b) The differential cross-section for scattering of photons of energy E' into the energy range E to $E + dE$ is independent of the lower energy E, and is equal to

$$\sigma(E' \rightarrow E)dE \approx \frac{2\pi r_0^2}{E'^2}\, dE, \quad 4.3 \leqq E' \leqq 7 \qquad (1)$$

Here $r_0 = e^2/mc^2$, and E' and E are the incident and scattered photon energy in mc^2 units. The maximum deviation of the correct Klein-Nishina formula from equation (1) is 3.5% which occurs for $E' = 7$ and $E = 4.3$.

(c) The total Compton cross-section, as a function of γ-ray energy, E between 4.3 and 7 mc^2, differs by less than 2% from the approximate formula.

$$\sigma(E) = 2\pi r_0^2 \left[1/(\alpha E) + 0.069\right]$$

$$\text{where } \alpha = 5/4. \qquad (2)$$

Consider a point source of γ-rays of energy E_0 at the center of a sphere of D_2O. The number of photons per unit source strength crossing a spherical surface at r that have suffered no collisions is $e^{-\mu_0 r}$ where μ_0 is the reciprocal mean free path of photons of energy E_0. According to Equ. (2)

$$\mu = 2\pi r_0^2 N \left[1/(\alpha E) + 0.089 \right] + \mu_\gamma(E) \qquad (3)$$

where N is the number of electrons per cm^3, and $\mu_\gamma(E)$ is the γ,n inverse absorption mean free path for photons of energy E. Hereafter $\mu_\gamma(E)$ is dropped from Equ. (3) because it at most amounts to only a 0.3% correction on μ.

Let $F(r,E)dE$ be the number of photons of energy E to $E + dE$ that cross the spherical surface at r after having suffered at least one collision. The balance between scattered photons entering and leaving the cell of extension dEdr is, upon inserting Equ. (1), given by:

$$\frac{\partial F}{\partial r} + \mu(E)F = \frac{2\pi r_0^2 N}{E_0} \int_E^{E_0} \frac{E_0 dE'}{E'^2} F(r,E') + \frac{2\pi r_0^2 N}{E_0^2} e^{-\mu_0 r} \qquad (4)$$

If one defines a new variable,

$$s = \mu - \mu_0 \approx \frac{2\pi r_0^2 N}{\alpha} (1/E - 1/E_0), \qquad (5)$$

and substitutes into equation (4) the form,

$$F = \frac{2\pi r_0^2 N}{E_0^2} e^{-\mu_0 r} \varphi(r,s). \qquad (6)$$

one obtains the following integro-differential equation for φ :

$$\frac{\partial \varphi}{\partial r} + s\,\varphi = 1 + d\int_0^s ds'\,\varphi\,(r,s') \;. \qquad (7)$$

Clearly the assumption of forward scattering (a) requires F,
and therefore φ, to vanish as r approaches zero. By taking the Laplace
transform of equation (7) with respect to the variable r, i.e.

$$L\,(\ldots) \equiv \int_0^\infty dr\, e^{-rs}\,(\ldots),$$ one obtains:

$$(s + s)\,f\,(s,s) = 1/s + d\int_0^s ds'\,f\,(s,s')\,. \qquad (8)$$

Where $f\,(s,s) = L\,\varphi(r,s)$.

Differentiating (8) with respect to s one obtains $\partial/\partial s\left[\ln f(s,s)\right]$
$= (d-1)/(s + s)$, which has the solution $f(s,s) = A(s,d)(s + s)^{d-1}$.
The constant of integration $A(s,d)$, is determined by substituting this
solution into Equ. (8). In this manner one obtains:

$$f(s,s) \;=\; s^{-(d + 1)}(s + s)^{d - 1} \qquad (9)$$

To obtain φ (r,s) the inverse Laplace transform of f(s,s) is required[*].

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* See "Collected Papers of Geo. A. Campbell", pp 443 and 320

$$\mathcal{P}(r,s) = L^{-1} f(s,s) = r \; {}_1F_1 (1 - q \; ; \; 2; \; -sr) \; . \tag{10}$$

Here ${}_1F_1 (a;b;x) = \sum_{n=0}^{\infty} \dfrac{\Gamma(a + n) \cdot \Gamma(b) \cdot x^n}{\Gamma(a) \cdot \Gamma(b + n) \cdot n \, !}$ is the confluent hyper-

geometric series or the so-called modified Kummer function.

The yield of γ,n neutrons in a sphere of radius R is made up of two parts. The fraction arising from the first collisions of the primary photons is

$$F_\gamma(1 - e^{-x}) . \tag{11}$$

Here F_γ is the fraction of the primary photons that would be absorbed in the γ,n reaction upon first collision in an infinite sphere of D_2O, $\mu_\gamma (E_0)//\mu_0$, and x is the sphere radius in units of the primary photon mean free path; $x = \mu_0 R$. The yield of second and higher collisions is given by $\int_{E_1}^{E} dE \mu_\gamma(E) \int_0^R dr F(r,E)$ which becomes, upon introducing the expressions (6) and (10)

$$F_\gamma \cdot D \tag{12}$$

Where $D = \left[\sigma(E_0 \rightarrow E)/\sigma(E_0) \right] \Big\rvert_{E_1}^{E_0} dE(\mu_\gamma(E)//\mu_\gamma(E_0)) \sum_{n=0}^{\infty} \dfrac{\Gamma(n-\tfrac{1}{4})(-B)^n}{\Gamma(-\tfrac{1}{4}) n \, !} P_n(x)$

Here B is an energy dependent function which is, according to the approximation (3),

$$B = (\mu(E) - \mu_0)//\mu_0 \approx (E_0/ E - 1)/(1 + .069 \, d \, E_0) . \tag{13}$$

and $P_n(x)$ is the integrated Poisson collision distribution of constant mean free path forward collisions; $n = 0$ corresponds to second collisions, $n = 1$ to third, etc.

$$P_n(x) = \int_0^x dx e^{-x} x^{n+1}/(n+1)! = 1 - e^{-x} \sum_{i=0}^{n+1} x^i/i! \qquad (14)$$

The lower energy limit E_1 in (12) is the γ,n threshold, $4.5 \, mo^2$.

The γ,n yield from all collisions in an infinite sphere of D_2O is the sum of (11) and (12) for $x = \infty$ namely, $P_\gamma \cdot E$ where

$$E = 1 + \left[\sigma(E_0 \to E)/\sigma_0\right] \int_{E_1}^{E_0} dE \left[\mu_\gamma(E)/\mu_\gamma(E_0)\right] (1 + B)^{\frac{1}{3}} \qquad (15)$$

Neglecting all except first collisions, the ratio between the finite and infinite sphere yield is

$$P_t = 1 - e^{-x} \qquad (16)$$

Taking into account second and higher collisions, one obtains the ratio between the infinite and finite sphere yield by multiplying the first collision yield ratio, $1/P_t$, by a factor β where

$$\beta = E/(1 + D/P_t) \qquad (17)$$

The quantities D and E defined by equations (12) and (15) were obtained by numerical integration over the γ-ray energies above the γ,n threshold. Dropping terms beyond n = 2 in the series appearing in Equ. (12) introduced an error in D of less than 0.2%.

PILE FACE

OIL BATH

PREAMPLIFIER

PNEUMATIC
TUBE

D_2O

RABBIT

10" R

CO_2

3"

FISSION
CHAMBER

$\frac{1}{8}$" ALUMINUM
SPHERE

¢

¢

4' D

FIG. I
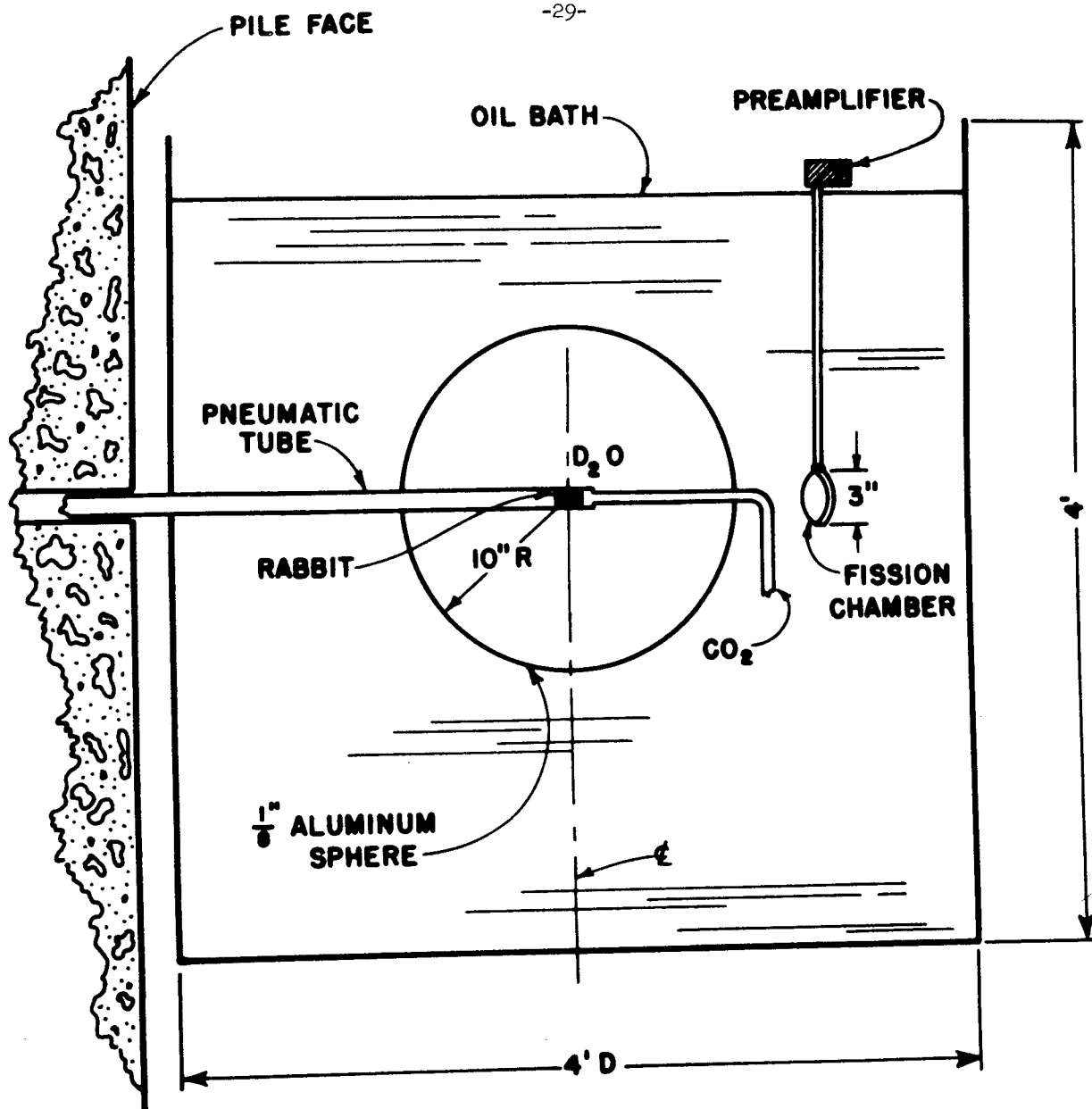
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9-24-46

SCHEMATIC DIAGRAM OF APPARATUS

SAMPLE OF FILM-RECORDED DATA

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9-24-46

Rabbit cut switch on

Rabbit arrives

Time scale, sec.

—Fig. 2—

Fig. 3

ACTIVITY x R² vs. R CURVES AT DIFFERENT TIMES "t" AFTER
THE END OF A 30sec. EXPOSURE.

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WITHOUT D₂O

WITH D₂O

$t = 0.5$ sec.

2.0 sec.

4 sec.

6 sec.

8 sec.

10 sec.

12 sec.

R INCHES

$(\text{ACTIVITY}) \times r^2; \ \frac{\text{FISSIONS}}{\text{sec.}} (\text{INCHES})^2$

Fig. 4

3.74 HOUR, EXPOSURE WITH D_2O

CURVE No.1 UNANALYZED DATA

No.2 $T_{\frac{1}{2}}$ = 53 hr.
No.3 $T_{\frac{1}{2}}$ = 4.37 hr.
No.4 $T_{\frac{1}{2}}$ = 1.65 hr.

Fig. 5
30min. EXPOSURE WITH D₂O
CURVE No.1 UNANALYZED DATA

• No.2 7·$t_{\frac{1}{2}}$ = 27.3 m. POINTS ARE
UNANALYZED DATA MINUS
(53 h + 4.37 h + 1.65 h PERIODS.)

No.1

No.2, 7,$t_{\frac{1}{2}}$ = 27.3 min.

Dr. 3011
9—24—46

ACTIVITY, COUNTS/sec

TIME "t" IN HOURS

Fig. 6
5 MINUTE EXPOSURE
No 1 UNANALYZED DATA
No 2 $T\frac{1}{2} = 7.7$ min. POINTS = VALUES OF
CURVES No.1 MINUS 4 LONGER
PERIODS.

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9-24-46

No.1

No.2, $T\frac{1}{2} = 7.7$ min.

TIME "t" MINUTES

ACTIVITY, COUNTS/sec

Fig. 7
3 MINUTE EXPOSURE

No. 1 TOTAL ACTIVITY, WITH D_2O
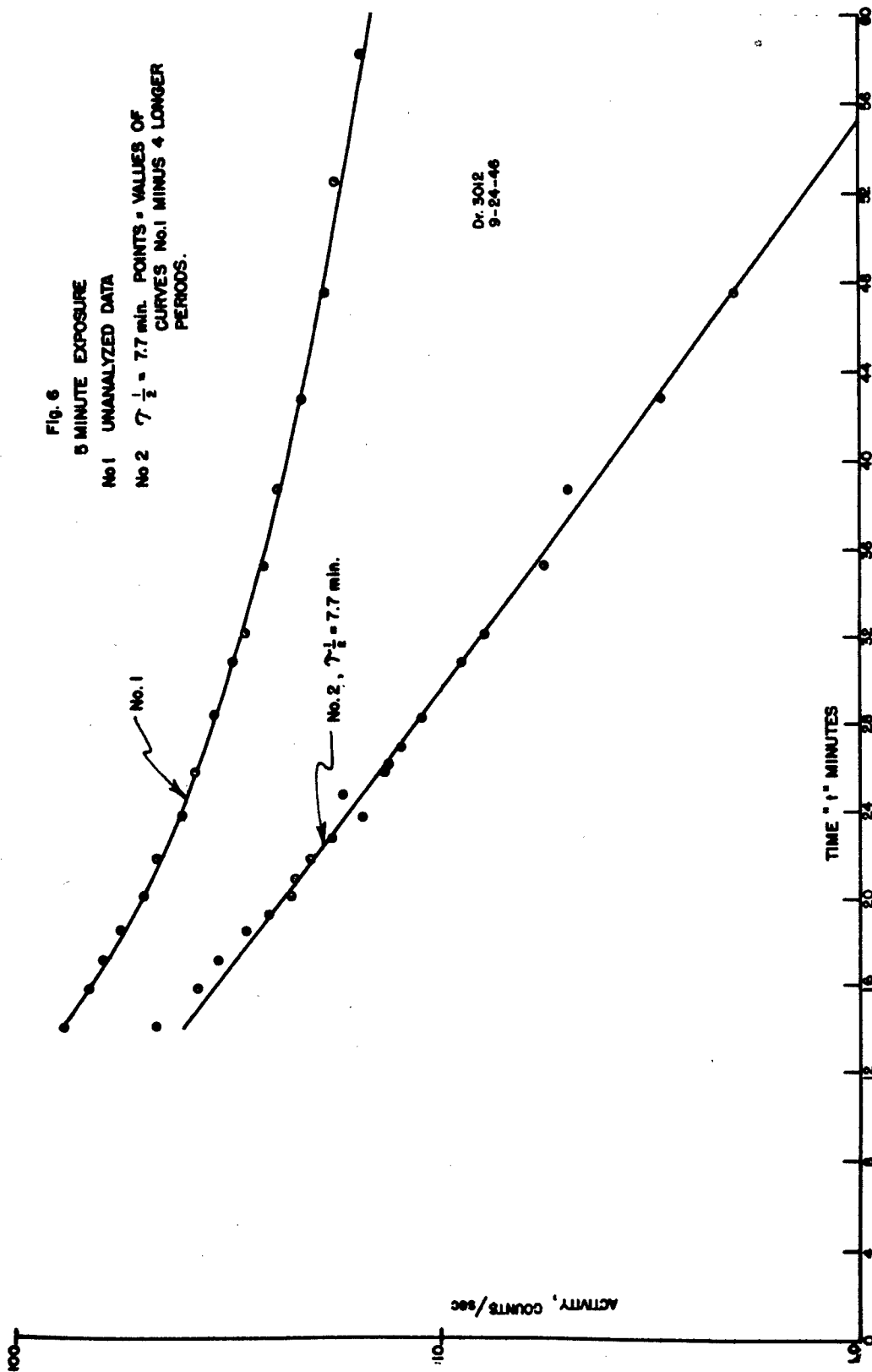No. 2 No. 1 MINUS CALCULATED DELAYED NEUTRON ACTIVITY
No. 3 $\gamma_{\frac{1}{2}} = 2.4$ min. POINTS FROM No. 2, MINUS ACTIVITIES
        OF 5 LONGER PERIODS.

Dr. 3013
9-24-45

No. 1
No. 2
No. 3

TIME "t", MINUTES

ACTIVITY, COUNTS/sec.

Fig 6
30 sec EXPOSURE VOLUME INTEGRALS
No.1 TOTAL ACTIVITY WITH D₂O
No.2 TOTAL ACTIVITY WITHOUT D₂O
No.3 PHOTONEUTRONS ALONE = No.1 MINUS No.2
No.4 $T_{\frac{1}{2}}$ = 41 sec, No. 3 MINUS 6 LONGER PERIODS

Fig. 9
30 sec. EXPOSURE VOLUME INTEGRALS

No. 1 TOTAL ACTIVITY WITH D_2O
No. 2 " " WITHOUT D_2O
No. 3 PHOTONEUTRONS ALONE, = No. 1 MINUS No. 2
No. 4 $\gamma \frac{1}{2} = 2.5$ sec, No. 3 MINUS 7 LONGER PERIODS

Dr. 3015
9-24-46

$$\int r^2 (\text{ACTIVITY}) \, dr, \frac{\text{sec.}}{\text{FISSION}} (\text{INCHES})^3$$

TIME "t" SECONDS