Lawrence Berkeley National Laboratory
Recent Work

Title
EFFECTIVE PHOTON ENERGIES OF HIGH-ENERGY PHOTO-NUCLEAR REACTIONS

Permalink
https://escholarship.org/uc/item/650233wn

Author
Eyges, Leonard

Publication Date
1950-09-07
UNIVERSITY OF CALIFORNIA
Radiation Laboratory

Contract No. W-7405-eng-48

UNCLASSIFIED

EFFECTIVE PHOTON ENERGIES OF HIGH ENERGY PHOTO-NUCLEAR REACTIONS

Leonard Eyges

September 7, 1950

Berkeley, California
DISCLAIMER

This document was prepared as an account of work sponsored by the United States Government. While this document is believed to contain correct information, neither the United States Government nor any agency thereof, nor the Regents of the University of California, nor any of their employees, makes any warranty, express or implied, or assumes any legal responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by its trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof, or the Regents of the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof or the Regents of the University of California.
<table>
<thead>
<tr>
<th>INSTALLATION</th>
<th>Number of Copies</th>
</tr>
</thead>
<tbody>
<tr>
<td>Argonne National Laboratory</td>
<td>8</td>
</tr>
<tr>
<td>Armed Forces Special Weapons Project</td>
<td>1</td>
</tr>
<tr>
<td>Atomic Energy Commission - Washington</td>
<td>2</td>
</tr>
<tr>
<td>Battelle Memorial Institute</td>
<td>1</td>
</tr>
<tr>
<td>Brush Beryllium Company</td>
<td>1</td>
</tr>
<tr>
<td>Brookhaven National Laboratory</td>
<td>4</td>
</tr>
<tr>
<td>Bureau of Medicine and Surgery</td>
<td>1</td>
</tr>
<tr>
<td>Bureau of Ships</td>
<td>1</td>
</tr>
<tr>
<td>Carbide and Carbon Chemicals Division (K-25 Plant)</td>
<td>4</td>
</tr>
<tr>
<td>Carbide and Carbon Chemicals Division (Y-12 Plant)</td>
<td>4</td>
</tr>
<tr>
<td>Chicago Operations Office</td>
<td>1</td>
</tr>
<tr>
<td>Columbia University (J. R. Dunning)</td>
<td>1</td>
</tr>
<tr>
<td>Columbia University (G. Failla)</td>
<td>1</td>
</tr>
<tr>
<td>Dow Chemical Company</td>
<td>1</td>
</tr>
<tr>
<td>H. K. Ferguson Company</td>
<td>1</td>
</tr>
<tr>
<td>General Electric, Richland</td>
<td>3</td>
</tr>
<tr>
<td>Harshaw Chemical Corporation</td>
<td>1</td>
</tr>
<tr>
<td>Idaho Operations Office</td>
<td>1</td>
</tr>
<tr>
<td>Iowa State College</td>
<td>2</td>
</tr>
<tr>
<td>Kansas City Operations Branch</td>
<td>1</td>
</tr>
<tr>
<td>Kellex Corporation</td>
<td>2</td>
</tr>
<tr>
<td>Knolls Atomic Power Laboratory</td>
<td>4</td>
</tr>
<tr>
<td>Los Alamos Scientific Laboratory</td>
<td>3</td>
</tr>
<tr>
<td>Mallinckrodt Chemical Works</td>
<td>1</td>
</tr>
<tr>
<td>Massachusetts Institute of Technology (A. Gaudin)</td>
<td>1</td>
</tr>
<tr>
<td>Massachusetts Institute of Technology (A. R. Kaufmann)</td>
<td>1</td>
</tr>
<tr>
<td>Mound Laboratory</td>
<td>3</td>
</tr>
<tr>
<td>National Advisory Committee for Aeronautics</td>
<td>1</td>
</tr>
<tr>
<td>National Bureau of Standards</td>
<td>3</td>
</tr>
<tr>
<td>Naval Medical Research Institute</td>
<td>1</td>
</tr>
<tr>
<td>Naval Radiological Defense Laboratory</td>
<td>2</td>
</tr>
<tr>
<td>New Brunswick Laboratory</td>
<td>1</td>
</tr>
<tr>
<td>New York Operations Office</td>
<td>3</td>
</tr>
<tr>
<td>North American Aviation, Inc.</td>
<td>1</td>
</tr>
<tr>
<td>Oak Ridge National Laboratory</td>
<td>8</td>
</tr>
<tr>
<td>Patent Branch - Washington</td>
<td>1</td>
</tr>
<tr>
<td>RAND Corporation</td>
<td>1</td>
</tr>
<tr>
<td>Sandia Corporation</td>
<td>2</td>
</tr>
<tr>
<td>Santa Fe Operations Office</td>
<td>2</td>
</tr>
<tr>
<td>Sylvania Electric Products, Inc.</td>
<td>1</td>
</tr>
<tr>
<td>Technical Information Division (Oak Ridge)</td>
<td>15</td>
</tr>
<tr>
<td>Armament Division, Deputy for Research and Development (Capt. Glenn Davis)</td>
<td>1</td>
</tr>
<tr>
<td>Assistant for Atomic Energy, Deputy Chief of Staff (Col. Robert E. Greer)</td>
<td>1</td>
</tr>
<tr>
<td>Chief of Documents and Disseminations Branch (Col. J. E. Mallory)</td>
<td>1</td>
</tr>
<tr>
<td>USAF Assistant for Research Director of Research and Development&lt;br&gt;Deputy Chief of Staff, Development (Colonel B. G. Holzman)</td>
<td>1</td>
</tr>
<tr>
<td>Electronic Systems Division (Mr. E. C. Trafton)</td>
<td>1</td>
</tr>
<tr>
<td>Chief of Scientific Advisors (Dr. Theodore Von Karman)</td>
<td>1</td>
</tr>
<tr>
<td>USAF, Eglin Air Force Base (Major A. C. Field)</td>
<td>1</td>
</tr>
<tr>
<td>USAF, Kirtland Air Force Base (Col. Marcus F. Cooper)</td>
<td>1</td>
</tr>
</tbody>
</table>
### INSTALLATION:

<table>
<thead>
<tr>
<th>Installation</th>
<th>Number of Copies</th>
</tr>
</thead>
<tbody>
<tr>
<td>USAF, Maxwell Air Force Base (Col. F. N. Moyers)</td>
<td>1</td>
</tr>
<tr>
<td>USAF, NEPA Office</td>
<td>2</td>
</tr>
<tr>
<td>USAF, Offutt Air Force Base (Col. H. R. Sullivan, Jr.)</td>
<td>1</td>
</tr>
<tr>
<td>USAF Surgeon General, Medical Research Division (Col. A. P. Gagge)</td>
<td>1</td>
</tr>
<tr>
<td>USAF, Wright-Patterson Air Force Base (Rodney Nudenberg)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Atomic Energy Branch (Lt. Col. A. W. Betts)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Army Field Forces (Captain James Kerr)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Commanding General, Chemical Corps</td>
<td></td>
</tr>
<tr>
<td>Technical Command (Col. John A. MacLaughlin thru Mrs. Georgia S. Benjamin)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Chief of Ordnance (Lt. Col. A. R. Del Campo)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Commanding Officer, Watertown Arsenal (Col. Carrowll H. Deitrick)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Director of Operations Research (Dr. Ellis Johnson)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Office of Engineers (Allen O'Leary)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Office of the Chief Signal Officer (Curtis T. Clayton thru Maj. George C. Hunt)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Army, Office of the Surgeon General (Col. W. S. Stone)</td>
<td>1</td>
</tr>
<tr>
<td>U. S. Geological Survey (T. B. Nolan)</td>
<td>2</td>
</tr>
<tr>
<td>U. S. Public Health Service</td>
<td>1</td>
</tr>
<tr>
<td>University of California at Los Angeles</td>
<td>1</td>
</tr>
<tr>
<td>University of California Radiation Laboratory</td>
<td>5</td>
</tr>
<tr>
<td>University of Rochester</td>
<td>2</td>
</tr>
<tr>
<td>University of Washington</td>
<td>1</td>
</tr>
<tr>
<td>Western Reserve University</td>
<td>2</td>
</tr>
<tr>
<td>Westinghouse Electric Company</td>
<td>4</td>
</tr>
</tbody>
</table>
EFFECTIVE PHOTON ENERGIES OF HIGH ENERGY PHOTO-NUCLEAR REACTIONS

Leonard Eyges

Radiation Laboratory, Department of Physics
University of California, Berkeley, California

September 7, 1950

ABSTRACT

We have attempted to use shower theory to evaluate the effective energies of the photo nuclear reactions measured by Strauch. It seems that these energies can be determined most accurately from the area under the transition curve, the so-called 'track length'. A theoretical formula for the track length is discussed. The shape of the transition curve at small thicknesses can also be calculated quite accurately and serves as a rough check on the effective energies as derived from the track length. A comparison with experiment of the theoretical shape of the whole transition curve is given and, as one would expect, the agreement is not very good.
EFFECTIVE PHOTON ENERGIES OF HIGH ENERGY PHOTO-NUCLEAR REACTIONS

Leonard Eyges
Radiation Laboratory, Department of Physics
University of California, Berkeley, California

September 7, 1950

Introduction

In this paper we try to use shower theory to evaluate some of Strauch's results on high-energy photo-nuclear reactions. As Strauch has described, the cross sections for these reactions have a more or less sharp maximum for some photon energy. For most of our calculations it will be adequate to assume that the width at this maximum is very small, i.e., that the reactions take place for only one photon energy, which we shall call $W_e$, the 'effective energy'. The effect of this approximation is discussed later. If it were easy to make accurate calculations with present shower theory there would be no problem; one would simply calculate shower curves for various energies $W_e$, and for some $W_e$ obtain a fit with the experimental curve. For the energies that we are interested in, however, around 20 Mev, it is well known that shower theory cannot be relied upon to predict an accurate cascade curve, mainly because the cross sections for pair production and bremsstrahlung vary considerably over the range of energies of interest, which is from about 20 to 300 Mev. We must, therefore, look to some quantity that can be calculated more accurately than the shape of the entire transition curve and yet one that gives us the information we want.

It is clear that one does not really have to know the whole transition curve in order to find the energy it corresponds to. If we consider transition

---

1 K. Strauch, Phys. Rev. (in press) 1951
curves corresponding to different energies, but to the same initial conditions, then at any thickness there is a unique correlation between the energy and the height of the curve. Thus, any one point on the transition curve determines, in principal, the energy. Of course, this is no real help, for if we could calculate an arbitrary point accurately, we could calculate the detailed shape. There is, however, a particular point on the transition curve which can be calculated rather more accurately than any other point, namely the height of the maximum. The reason is, as Rossi and Greisen\(^2\) have pointed out, that at the maximum of the shower one can take into account approximately the variation of the pair production cross section with energy. This enhances the accuracy considerably. Thus, if the shower curve corresponding to an energy \(W_e\) shows a maximum, one might hope to determine \(W_e\) by the position and height of the maximum. There lies the difficulty. Although some of Straugh's curves have a maximum, those corresponding to higher energies do not. We must find a different method if we wish it to be universally applicable.

For very large thicknesses multiplication becomes unimportant, and the "shower" curves simply become the exponential absorption curve of photons of energy \(W_e\).\(^*\) One might hope to determine the absorption coefficient from the slope of the experimental curve and from this get the energy \(W_e\). There is no difficulty in principal with this idea; in practice one must go to such large thicknesses before pure absorption sets in that the intensity becomes unusably small.

The beginning of Straugh's shower curves have a characteristic shape. There is a drop at very small thicknesses due to the absorption of photons of

\(^2\) B. Rossi and K. Greisen, Rev. Mod. Phys. 13, 274 (1941).

\(^*\) If we take the finite width of the reaction cross sections into account, the shower curve at large thicknesses really becomes the absorption curve of the photons of lowest energy that can produce the reaction.
energy $W_e$; multiplication soon sets in however, and the curve becomes less steep and may even rise again. The initial slope is entirely due to self absorption and this slope, in principal, determines the energy. Unfortunately, this slope is very difficult to measure with any accuracy. On the other hand, one might hope that since the first part of the shower curve, up to perhaps a half radiation length, is mainly an absorption curve, multiplication processes being secondary, one might be able to calculate this multiplication with sufficient accuracy to predict the behaviour of the beginning of the curve with reasonable accuracy. This expectation is fulfilled. One can calculate the shower curve to almost a radiation length with considerable accuracy. Unfortunately, there is an experimental limitation. It is difficult to get good statistics on the beginning part of the curve. Therefore, one cannot obtain a very accurate value for $W_e$ in this way.

Finally, it is possible to determine the effective energy $W_e$ from the area under the transition curve, the so-called 'track length'. This quantity has the advantage that one can take the variation of the pair production cross section with energy into account, just as in calculating the height at the maximum. Moreover, it has an advantage over the latter quantity: although not all of Strauch's transition curves have a maximum, all have an area. It is the track length that we have mainly used in calculating $W_e$, although we have also used the initial behaviour of the shower curve, up to almost a radiation length, as a rough check.

The rest of the paper is in four parts: in section I we have calculated the photon spectrum to be expected from the synchrotron target; in section II we have calculated the track length as a function of energy, and applied our formula to Strauch's results; in section III we have calculated
the detailed shape of the curves at their beginning; finally, in section IV, we have calculated, as best we could, the detailed shape of the transition curves, neither expecting nor getting very good agreement, with experiment.

For convenient reference, we will write here the usual shower equations, using the notation of Rossi and Greisen unless otherwise indicated.

\[
\frac{\partial n(E, t)}{\partial t} = 2 \int_0^1 \gamma(E, u) \psi(E, u) \frac{du}{u} - \int_0^1 \pi(E, t) \phi(E, v) dv
\]

\[
+ \int_0^1 \frac{1}{1-v} n(E, t) \phi(E, \frac{E}{1-v}, v) dv + \varepsilon \frac{\partial \pi(E, t)}{\partial E}
\]

\[
\frac{\partial \gamma(W, t)}{\partial t} = \int_0^1 \pi(W, t) \phi(W, \frac{W}{v}, v) \frac{dv}{v} = \sigma(W) \gamma(W, t)
\]

Here \(n(E, t)\) is the number of electrons of energy \(E\) at thickness \(t\) and \(\gamma(W, t)\) the same for photons. \(\psi(W, u)\) is the probability per radiation length that a photon of energy \(W\) produce a pair, one particle of which has fractional energy \(u\). \(\phi(E, v)\) is the probability per radiation length that an electron of energy \(E\) emit a photon with fractional energy \(v\). The usual shower theory deals with high energies where \(\psi\) and \(\phi\) are functions of \(u\) and \(v\) only, but for our purposes, keeping the dependence on \(W\) and \(E\) explicit facilitates discussion. \(\varepsilon\) in the above equations is the critical energy, and thicknesses are, of course, measured in radiation units. We also depart slightly from Rossi and Greisen by letting \(\sigma(W)\) be the total absorption coefficient for photons of energy \(W\). This will be discussed later. If we call \(\sigma_c(W)\) the absorption coefficient for the Compton effect and \(\sigma_p(W)\) that due to pair production, then \(\sigma(W) = \sigma_p(W) + \sigma_c(W)\).
I. Photon Spectrum from Target

The 322 Mev electrons from the beam of the synchrotron are allowed to fall on a target of Pt, 0.020 in. thick, producing the beam of photons used in the experiment. If the target were infinitely thin the distribution of photon energies \( W \) should be given by \( \phi (E_0, v) \) where \( E_0 = 322 \) Mev and \( v = \frac{W}{E_0} \). Actually, the finite thickness of the target introduces a correction, which we shall calculate in this section. First, we should like to make explicit an assumption inherent in our use of the function \( \phi (E_0, v) \). This function gives the energy distribution of photons produced by an electron of energy \( E_0 \), integrated over the angles between the electron and the photon. At first sight, one might think that the appropriate function for our purposes should be the energy distribution of photons produced in essentially the same direction as the electron. In passing through the target, however, the electrons are multiply scattered and in the present geometry the effect of these deviations due to scattering will be taken care of to a good approximation by using the integrated function \( \phi (E_0, v) \).

We can get the photon spectrum by simply putting into the shower equations a power series expansion corresponding to the correct initial conditions, i.e.

\[
\pi (E_0, E, t) = \phi (E_0 - E) + P(E_0, E)t + \ldots \\
\gamma (E_0, W, t) = Q (E_0, W)t + R(E_0, W)t^2 + \ldots
\]

Equating to zero various powers of \( t \) in the shower equation, the unknown functions \( P, Q \) and \( R \) are found in succession as easily evaluable integrals. \( Q(E_0, W) \) comes out to be just \( \phi (E_0, v) \), as it must. If we use the approximate expression \( \phi (E_0, v) \approx 1/v \) to calculate the small correction term \( R(E_0, W) \)

\[3\] L. I. Schiff, Phys. Rev. 70, 87 (1946)
we get

\[ R(E_0, W) = -\frac{1}{2} \left[ \sigma(W) - \ln \left( \frac{W}{E_0} \right) \right] \]

Thus our corrected spectrum from the target is:

\[ \phi(E_0, v) - \frac{t}{2} \left( \sigma(W) - \ln (1-v) \right) \]

This corrected spectrum is plotted in Figure 7 of Strauch's paper. The physical interpretation of this spectrum is clear. For \( v \) small, \( \ln(1 - v) \approx 0 \) and the important term in the correction is just the absorption of photons in the target. The correction term \( -\frac{\sigma(W) t}{2} \) represents this self absorption. For large \( v \) the term in \( \ln(1 - v) \) is important. This term diminishes the number of high energy photons. This represents a double radiation process: there is an overwhelming probability for emitting a low energy quantum in which the energy of the electron is diminished below 322 Mev; therefore, it can no longer emit a quantum with this upper limit. Thus, the effect of the finite target thickness is to diminish appreciably the number of very high energy photons.

After leaving the target, the beam must pass through a quartz donut about 5/8 in. thick. The main effect of this on energies above 18 Mev is to reduce the intensity uniformly, since the absorption coefficient is small and varies slowly with energy in this region. We shall, therefore, neglect this correction.

II. The Track Length

The most accurate calculation of the track length of photons to date is the numerical work by Richards and Nordheim\(^4\) in which collision loss of electrons and the Compton effect are taken into account, as well as the varia-

\(^4\) J. Richards and L. Nordheim, Phys. Rev. 74, 1106 (1948)
tion with energy of the radiation and pair production cross sections. These
are not very convenient for our purpose, however, since they are made for a
single incident photon. To apply them to the present problem, one would have
to integrate numerically their results over the photon spectrum emerging from
the synchrotron. Moreover, aside from the labor involved in this, there is
the difficulty that Richards and Nordheim's results hold only when the single
incident photon has an energy much larger than the energy \( W_e \) of the photons
one is considering; in these circumstances it is not clear how to carry out
the integration over the photon beam from the synchrotron, since, of course,
it contains photons with energy arbitrarily close to \( W_e \).

Fortunately, the photon energies \( W_e \) with which we have to deal are
always greater than about 18 Mev, i.e., about two and a half times the criti-
cal energy in lead. In this case, it is possible to make a slight adaptation
of the formulae for the track length in Rossi and Greisen so that they apply
with considerable accuracy. Before we do this, there is a somewhat peculiar
feature of the usual track length formulae which we should like to discuss.

For the sake of discussion, suppose we are interested in the track length of
photons of energy \( W \), due to an initial spectrum which goes as \( \frac{1}{W} \) up to some
maximum energy \( W_o \). Now, following Nordheim and Hebb\(^5\) the track length of
photons of energy \( W \), due to an arbitrary initial spectrum \( \gamma (W_o, W, 0) \) is

\[
Z(W_o, W) = \frac{1}{\sigma_{2ni}} \int_{s=1}^{s=1} \frac{A(s)M(s, \sigma)}{A(s) - B(s)C(s)} \cdot ds
\]

(3)

where

\[
M(s, \sigma) = \int_{W_o}^{\infty} \frac{1}{W} \gamma (W_o, W, 0) dW
\]

\(^5\) L.W. Nordheim and M.H. Hebb, Phys. Rev. 56, 494 (1939)
For a $\frac{1}{W}$ spectrum up to $W_0$, $M(s, 0) = \frac{W_0 s}{s}$ and we therefore have

$$Z(W_0, W) = \frac{1}{\alpha W \cdot 2\pi i} \int_{\delta - i \infty}^{\delta + i \infty} \frac{A(s)}{A(s) - B(s) C(s)} \frac{e^{\gamma s}}{s} ds,$$  \hspace{1cm} (4)

where $\gamma = \ln \frac{W_0}{W}$ and the integration path is to the right of all the singularities of the integrand. Here $A(s), B(s), C(s)$ are defined as in Rossi and Greisen. Now the integrand of (4) has simple poles at $s = 1, -2.6, -3.6, \ldots$. If we evaluate the residues at these poles and divide the result by the initial spectrum $\frac{dW}{W}$ to get the track length relative to the number of photons at $t = 0$ we get

$$Z_{\text{relative}}(W_0, W) = \frac{1}{\sigma} \left[ 0.437 \frac{W_0}{W} - 0.02 \left( \frac{W}{W_0} \right)^{2.6} - 0.005 \left( \frac{W}{W_0} \right)^{3.6} + \ldots \right],$$  \hspace{1cm} (5)

Now it is clear that (5) becomes incorrect when $W$ approaches $W_0$. The relative number of photons of energy $W$, when $W$ is very close to $W_0$, will be given by $e^{-\sigma t}$, since there will be essentially no multiplication. Hence, $Z_{\text{rel.}}$ will be just $\frac{1}{\sigma}$; and for smaller $W$ where there is multiplication $Z_{\text{rel.}}$ must be greater than $\frac{1}{\sigma}$. This condition fails to hold for (5) when $W \approx 0.41 W_0$. It is not clear to us why (5) is incorrect for $W \sim W_0$, when (4) is almost certainly correct. It may be that the integrand in (4) has singularities off the real axis, although Nordheim and Hebb have made a search near $s = 1$, and we also have made a rather perfunctory search, without finding any. We are concerned with this point, not because we want to use a formula like (5) for $W$ close to $W_0$ -- in Strauch's experiments $\frac{W}{W_0}$ is always fairly small -- but we really would like to know whether (5) can be considered correct for

* Note that $B(s)$ as defined in Reference (2) has a factor $\sigma$ in it, so that the integrand in (4) is really independent of $\sigma$.

**This was called to my attention by Dr. Strauch.
small, where the higher order terms are negligible. This is not obvious, since the fact that \( W \sim W_o \) throws suspicion on it.

Some light can be shed on this question in the following way. If one uses the Carlson-Oppenheimer approximation to the shower equations, it is easy to show that this is equivalent to using

\[
A(s) = \frac{2s}{s+1}, \quad B(s) = \frac{2\sigma}{s+1}, \quad C(s) = \frac{1}{s}.
\]

Then

\[
Z_{rel}(W_0, W) = \frac{1}{\sigma \cdot 2\pi i} \oint_{\delta = 1} \frac{e^{ys}}{s^2 - 1} ds = \frac{0.500}{\sigma} \left( \frac{W_0}{W} + \frac{W}{W_0} \right).
\]

This is a most reasonable result since \( Z_{rel} \) as given by (6) is always greater than \( \frac{1}{\sigma} \) and approaches \( \frac{1}{\sigma} \) as \( W \) approaches \( W_o \). Unfortunately, (6), however reasonable in appearance, cannot really be trusted for \( W \) close to \( W_o \), since the Carlson-Oppenheimer approximation is not very good in this region. For smaller \( W \), Equation (6) shows that \( Z_{rel} \) is proportional to \( \frac{W_o}{W} \) with a correction term of order \( \left( \frac{W}{W_o} \right)^2 \), which is small. This is probably a trustworthy qualitative conclusion in general, since the Carlson-Oppenheimer approximation is not bad for \( W < W_o \).

We have also tried to check (5) in the following way: we have calculated the shower curve as a function of thickness for various values of \( y = \ln \frac{W_0}{W} \) and integrated these numerically to find the track length; we have used an expansion in powers of \( t \) for small \( t \) and the usual saddle point method for larger \( t \). This method gives reasonable results; e.g., as \( W \) approaches \( W_o \) the relative track length approaches \( \frac{1}{\sigma} \). The main

---

difficulty is that it is not very accurate since the saddle point method can be off by 10 or 15 percent for the smaller values of $y$ and $t$. We have corrected for the inaccuracies of the saddle point method as best we could by comparing the answers it gives with the quite accurate results given by the power series in $t$, for those $t$ for which one can get an answer by both methods. Comparing our answers for $Z_{re1}$ by this method with (5) it appears that (5) is correct for $y = 3$, is a few percent low for $y = 2$, about 30 percent low for $y = 1$ and, of course, off by a factor $\frac{1}{0.41}$ for $y = 0$. We have done the same sort of calculation for an initial spectrum

$$\frac{1}{W} \left[ \frac{4}{3} \left( 1 - \frac{W}{W_0} \right) + 0.38 \left( \frac{W}{W_0} \right)^2 \right],$$

which is a rough approximation to the spectrum from the synchrotron, and find that the formula corresponding to (5) is more accurate than for the $\frac{1}{W}$ spectrum, being off, e.g. only by a few percent for $y = 1$. All in all, then, it seems clear that although formulae like (5) are not correct for $W$ close to $W_0$, they are probably all right for $W << W_0$.

Now we turn to the real problem of interest, that of calculating as accurately as possible the track length of photons using the initial spectrum given by (2). We are interested in energies from about 18 Mev up. Energy loss of electrons by ionization is not negligible in this range, but, as Rossi and Greisen have shown, one can correct for this by using an asymptotic expansion in powers of $\frac{E}{W}$. The variation with energy of the pair production cross section can also be included in the manner indicated by Rossi and Greisen, i.e., by writing

$$\psi(W_0, u) du = \sigma_p(W) du$$

and considering that the unknown function in the shower equations for the track track length is not $Z(W_0, W)$ but $\sigma_p(W) Z(W_0, W)$. The above approxima-

\textsuperscript{7} Reference 2, p. 293.
tion for $V(W, u)$ means that the pair spectrum is taken to be flat, i.e. the
probability for producing an electron of any energy is independent of energy.
This is quite a good approximation in the range $18 - 322$ MeV. The variation
of the radiation cross section with energy can be included by taking some
average expression appropriate to the region $18 - 322$ MeV. The expression
we have chosen is

$$
\Phi(v) = \frac{1}{v} \left[ \frac{4}{3} (1-v) + \frac{3}{4} v^2 \right]
$$

Reference to Rossi and Greisen will show that this seems a reasonable
approximation except for $v$ close to unity, where $\Phi(v)$ is relatively
small anyway.

If one carries out the calculations according to the above sketch,
using the boundary condition that the incident spectrum is that given by (2)
one gets

$$
Z(W_0, W) = \frac{0.346 W_0}{\sigma_p(W) \cdot W^2 \cdot \nu(\xi/W)}
$$

(9)

where

$$
\nu(\xi/W) = 1 + 0.71 \frac{\xi}{W} = 0.32 \left( \frac{\xi}{W} \right)^2 + 0.41 \left( \frac{\xi}{W} \right)^3 + \cdots
$$

We have not included the dubious negative powers of $\frac{W_0}{W}$ that one gets from
evaluating (4) at the poles on the negative real axis, since the discussion
at the beginning of this section implies that they are negligible for the
energies we will be interested in. The factor $0.346$ appears in (9) instead
of the usual $0.437$ for two reasons. First, $W(1, 0)$, as evaluated by numeri-
cal integration of the spectrum given by (2), turns out to be $0.816 \frac{W_0}{W}$
instead of $\frac{W_0}{W}$ as before. Second, we have used the $A(s), B(s),$ and $C(s)$
corresponding to the $\Phi(v)$ and $\Psi(u)$ given above instead of those calculated
with the usual asymptotic $\Phi(v)$ and $\Psi(u)$, as in Rossi and Greisen. This
alters the residue at \( s = 1 \). This change is rather insensitive to the choice of \( \phi(v) \). E.g., if we use the \( A(s) \) corresponding to \( v \phi(v) = \frac{4}{3} (1-v)v^2 \) and to \( \frac{4}{3} (1-v) + \frac{v^2}{2} \), the factor 0.346 changes to 0.355 and 0.330 respectively. Thus, using the average \( \phi(v) \) given by (8) introduces only a small error.

At the lowest energies for which we wish to use (9) the cross section for Compton effect is about 15 percent of that for pair production, and is therefore not negligible. To take this into account accurately one would have to supplement Equations (1) by a term that describes the production of electrons with energy greater than \( W \) in the Compton effect, and a term that describes the photons with energy greater than \( W \) that get an energy \( W \) in a Compton scattering. This is difficult and we shall not attempt it, since the effect of these terms is probably small anyway. In addition to these effects the Compton effect acts to absorb the photons of energy \( W \) that we are interested in. One takes this roughly into account in the following way. One replaces \( \sigma_p(W) \) in (7) by \( \sigma(W) \) and as mentioned before uses the total absorption coefficient \( \sigma(W) \) in (1b). A glance at equations (1b) shows that this means we take the absorption of photons of energy \( W \) into account correctly, but that we falsify the spectrum of electrons with energy greater than \( W \) since replacing \( \sigma(W) \) by \( \sigma_p(W) \) makes the pair cross section too large. But a photon of 13 Mev is produced on the average by an electron of, say, twice that energy, where the Compton effect is very small anyway, so it doesn't matter that we have allowed \( \Psi(W,u) \) to include the Compton effect. Also, and this is probably a stronger argument, we have checked (9) by comparing the analogous formula for a single incident photon with the numerical results of Richards and Nordheim and found agreement within 5 percent from 14 Mev up, if we take \( \sigma(W) \) and not \( \sigma_p(W) \).
In evaluating the experiments, one wants not the track length given by (9), but \( \frac{Z_{\text{relative}}(W_0, W)}{Z_{\text{relative}}(W_0, W)} \), the track length relative to the number of photons initially present in \( dW \).

If we use the notation

\[
\gamma(W_0, W, 0) = \frac{f(W_0, W)}{W}
\]

and make the change from \( \sigma'(W) \) to \( \sigma'(W) \) just mentioned, the relative track length is given by

\[
Z_{\text{rel.}}(W_0, W) = \frac{0.346 W_0}{\sigma'(W) W \nu(\frac{\tilde{E}}{W}) f(W_0, W)}
\]

In applying (10) we have, as Strauch has explained, increased the radiation length by 10 percent over the value given in Rossi and Greisen and decreased the pair production cross section per cm by 10 percent. It is obviously difficult to estimate the error in (10), but if forced to guess we would say that it is probably good to 15 percent at 18 MeV, and perhaps 10 percent at twice the energy. As we have discussed earlier in this section, (10) must break down for \( W \sim W_0 \), but if we can extrapolate from our previous results it should still be reasonably accurate for \( \frac{W}{W_0} < \frac{1}{3} \).

As Strauch has noted, a kind of internal check on the experiments and theory can be had by carrying out an experiment on a given element with two different maximum energies \( W_{01} \) and \( W_{02} \) of the photon beam. The ratio of the relative track lengths is then

\[
\frac{Z_{\text{rel.}}(W_{01}, W)}{Z_{\text{rel.}}(W_{02}, W)} = \frac{W_{01} f(W_{02}, W)}{W_{02} f(W_{01}, W)}
\]
This theoretical expression for the ratio should be quite accurate, since most of the approximations involved in the derivation of (10) effectively cancel in forming it. Equation (11) has been checked for the two reactions $\text{Cu}^{62}(\gamma, n) \text{Cu}^{62}$ and $\text{C}^{12}(\gamma, n) \text{C}^{11}$, which were done at maximum energies 322 and about 200 Mev. The results are given in Table 1 of Strauch's paper.

III. Small Thicknesses

If we wish to find the shape of the transition curve for small thickness, an expansion in powers of $t$ suggests itself. As we have seen, the transition curve must drop at the very beginning, since absorption of the photons in the incident beam is a first order effect proportional to $t$, and the production of photons is at least of second order. This suggests using an expansion of the form

$$\gamma(W_0, W, t) = e^{-\sigma(W)t} \left[ \gamma(W_0, W, 0) + K(W_0, W)t + L(W_0, W)t^2 + \cdots \right] \quad (12a)$$

$$\pi(W_0, E, t) = \left[ M(W_0, E)t + N(W_0, E)t^2 + \cdots \right] \quad (12b)$$

This expansion obviously satisfies the boundary conditions. The factor $e^{-\sigma(W)t}$ in (12a), of course, represents the absorption of photons of energy $W$ initially in the beam and the coefficients $K(W_0, W), L(W_0, W)$ etc. describe their subsequent multiplication.

These functions can be found by putting (12a) and (12b) into the shower equations (1) and equating to zero successive powers of $t$. The functions then come out as simple, but sometimes tedious integrals. Alternately, one can find, in much the same manner, the Mellin transforms of the shower equations as a series in $t$ and invert this transform, evaluating the complex integrals that result by the method of residues. Both methods,
of course, lead to the same result, which we first write down and then discuss. As before, the quantity of interest is not \( \gamma(W_0, \bar{W}, t) \) but

\[
\gamma_{rel.}(W_0, \bar{W}, t) = \frac{\gamma(W_0, \bar{W}, t)}{\gamma(W_0, \bar{W}, 0)}
\]

Our results are

\[
\gamma_{rel.}(W_0, \bar{W}, t) = e^{-\sigma(W)t} \left[ 1 + \frac{\overline{\sigma_p} F(y)t^2}{f(W_0, \bar{W})} + \frac{G(y)t^3}{3f(W_0, \bar{W})} + \ldots \right]
\]

where

\[
y = \ln \frac{W_0}{\bar{W}}, \quad \overline{\sigma_p} = \sigma_p \left( \sqrt{\frac{W_0}{\bar{W}}} \right)
\]

\[
F(y) = \frac{16}{9} (3e^{-y} - 3 + y + y^2 \frac{e^{-y}}{2} + 2y e^{-y}) + 1.8 \left( \frac{1}{2} + \frac{e^{-2y}}{2} \right) e^{-y} e^{-2y}
\]

\[
+ 0.45 (e^{-y} - e^{-2y} + y e^{-2y})
\]

\[
G(y) = \overline{\sigma_p^2} (y - 1 + e^{-y}) = \overline{\sigma} H(y) = \overline{\sigma_p} \frac{\xi}{W_0} (e^y - 1)
\]

\[
H(y) = 1.645 - \sum_{n=1}^{\infty} \frac{e^{-ny}}{n^2} + (1-e^{-y}) \ln(1-e^{-y}) = y e^{-y}
\]

The following approximations have been used in calculating the above result. As an analytic approximation to the initial spectrum, we have used \( \overline{nY(W_0, \bar{W}, 0)} = \left[ \frac{4}{3} (1 - \frac{\bar{W}}{W_0}) + 0.6 \frac{\bar{W}^2}{W_0^2} \right] \). In calculating the term in \( t^2 \), we have used, as for the track length, \( \overline{nY(v)} = \frac{4}{3} (1-v) + \frac{3}{4} v^2 \), and \( \overline{\gamma(W, u)du} = \text{const.} u \). In calculating the small term in \( t^3 \), we have, for simplicity, used \( \overline{nY(v)} = 1/v \). In the terms in \( t^2 \) and \( t^3 \) describing the multiplication, there is the factor \( \overline{\sigma_p} \left( \sqrt{\frac{W_0}{\bar{W}}} \right) \). This enters in the following way: multiplication takes place because electrons of some average energy between \( W_0 \) and \( \bar{W} \) are created and then radiate photons of energy \( \bar{W} \); for this average energy we have chosen the geometrical mean.
of $W_0$ and $W$; hence, the pair production cross section at this energy is
\[ \sigma_p(\sqrt{W_0/W}). \]
We have checked this approximation by writing $\sigma_p(W)$ as a linear function of $\ln(W)$, which is a fair representation in the energy range of interest here, and then calculating the coefficient of $t^2$ using the initial spectrum $\gamma(W_0, W_0) = \frac{1}{W}$, and the approximation $\phi(v) = 1/v$. One can then carry out the integrations and it turns out that to a very close approximation one gets the same results for the coefficient of $t^2$ as if he had started from the beginning with the average value $\sigma_p(\sqrt{W_0/W})$.

For $W = W_0$ it is clear that (13) must become $e^{-\sigma(W)t}$, since there can no longer be any multiplication. Thus, for this case ($y = 0$), $F(y)$ and $G(y)$ must vanish. This provides a useful check on the calculations. In the terms in $t^3$ there enters a correction due to collision loss. This correction diverges as $W$ goes to zero, but for $W > E_0$ it is probably correct.

We have used the above formula to calculate the beginning shapes of the transition curves for the reactions $\text{Cu}^{63}(\gamma, n) \text{Cu}^{62}$ and $\text{C}^{12}(\gamma, n) \text{C}^{11}$, using for the respective 'resonance' energies the values 18 Mev and 27 Mev derived from the track length. The results are shown in Figure 1. The experiments, of course, do not give very reliable results at these thicknesses since one is trying to measure changes in intensity of the order of a few percent and very long counting periods are needed to get good statistical accuracy. Within the experimental errors, however, the theory seems to give fair agreement. If anything, the theoretical curve seems to be too low. The theoretical curves would be raised if one assumed that the effective energies were somewhat lower than 18 and 27 Mev, but the poor accuracy of the experiments does not justify this.

One should note that the expansions given above are quite accurate.
where they apply. This is because the main phenomenon at small thicknesses is simply the absorption of original photons and one knows accurately the absorption coefficient for this. The shower theory enters, of course, in giving the coefficients $F(y)$ etc. for the higher order terms, but it turns out that these are not at all sensitive to the approximate expression for the cross sections one chooses for radiation and pair production.

IV. Miscellany

We have also calculated as best we could, the detailed shape of the transition curves for the $^{63}$Cu ($\gamma$,n) $^{62}$Cu and $^{12}$C ($\gamma$,n) $^{11}$C reactions using the usual saddle point method. As in calculating the track lengths, one can take some reasonable average value for the radiation cross section and take into account ionization loss by using the asymptotic expansions as given in Rossi and Greisen. One cannot, however, take into account the variation of the pair production cross section in even the approximate way in which it was done in calculating the track length. Since the pair production varies by almost a factor of two between maximum and minimum energies with which we deal, i.e., between 322 and 18 Mev, considerable uncertainty is introduced into the results. Nonetheless, we thought it might be of some interest to present them.

Since one cannot take into account the variation of the absorption cross section $\sigma(W)$ with energy, one must choose some average value in carrying out the calculations. The question arises as to what is the most reasonable value for this. We have chosen to use $\sigma(W_{e})$ for the following reasons: this value is roughly correct for the track length and, for the same reasons that apply there, for the maximum of the shower curve; also, for very large thicknesses, the cascade curve approaches a pure absorption
curve with absorption coefficient $\sigma(W_e)$.

The saddle point method leads to the following expression

$$\gamma_{rel}(W_0, W, t) = \frac{1}{\sqrt{2\pi}} \frac{H_2(s)}{f(W_0, W)} \frac{e^{\lambda_1'(s) t}}{\sqrt{\lambda_1''(s) t + 1/s^2}} \frac{M(s, 0)}{\sqrt{\lambda_1''(s) t + 1/s^2}}$$

where

$$t = \frac{1/s - \gamma}{\lambda_1'(s)}$$

Here, the functions $\lambda_1(s), H_2(s)$ and $\nu(s, \frac{E}{W})$ are defined in terms of $A(s), B(s)$ and $C(s)$ as in Rossi and Greisen, but in actually calculating the latter functions we have used $\nu(u)du = \sigma(W_e)du$ and the $\phi(v)$ given by (6). $M(s, 0)$ is calculated numerically from the curve in Figure (7) of Strauch's paper. Using the above expression, we have calculated the transition curves for the reactions $^{63}\text{Cu}(\gamma, n) ^{62}\text{Cu}$ and $^{12}\text{C}(\gamma, n) ^{11}\text{C}$. The curves are shown in Figure 2. Whether the agreement is better or worse than one should expect is a moot question. The agreement for Cu with a resonance energy of 18 Mev is not as good as that for C for which the resonance energy is 27 Mev. This is not implausible qualitatively, since the various approximations involved in taking average cross sections and neglecting Compton effect are somewhat more serious at 18 than at 27 Mev. One might perhaps have expected better agreement at the maximum, for the reasons given in the introduction.

It may be, however, that the errors in the saddle point method are not negligible. It is altogether possible that the saddle point method gives too low a value by perhaps 10 percent near the maximum; if this is true the shower theory proper is in better agreement with the experiment than evaluation by the saddle point method would seem to imply.
In all the work thus far, we have assumed that the \((\gamma, n)\) cross sections are infinitely sharp, i.e., if we call \(\Sigma(W)\) the cross section as a function of energy, that \(\Sigma(W) = \delta(W - W_e)\) where \(W_e\) is the 'resonance' energy. What then is the effect of the finite width?

Suppose for illustration that \(\Sigma(W)\) is constant, and has a square shape centered about a value \(W_e\), and with width \(\Delta\) i.e.,

\[
\Sigma(W) = \text{constant} \quad W_e - \frac{\Delta}{2} < W < W_e + \frac{\Delta}{2}
\]

\[
\Sigma(W) = 0 \quad \text{otherwise}.
\]

It is then easy to see, e.g. that if \(\frac{\Delta}{2} \ll W_e\) that Equation (9) is replaced approximately by the following one

\[
Z_{\text{rel.}}(W_0, W) = \frac{0.346 \ W_0 \ (1 + \frac{\Delta^2}{6 W_0^2})}{\sigma(W) \ W_e \ \Gamma(W_0, W) \ \Gamma(W)}
\]

The effect of the finite width is quite small. E.g., if \(W_e = 20, \Delta = 10\), this effect increases \(Z_{\text{rel.}}\) by about 4 percent. One can also see that the effect of the finite width varies with depth in the shower. The spectrum of photons goes as \(\frac{1}{W^{s+1}}\) where \(s\) is 0 at the beginning of the shower, is unity at the maximum and increases slowly with thickness thereafter.

Thus, the effect of the finite width in raising the shower curve increases slowly with thickness. Taking it into account would therefore slightly increase the discrepancy between experiment and theory shown in Figure 2.

I should like to thank Professor Robert Serber for several interesting discussions and helpful suggestions.

This work was performed under the auspices of the Atomic Energy Commission.

Information Division
9-12-50 cg
Fig. 1a - Relative intensity in Pb of photons causing $^{12}(\gamma\gamma)\gamma$.

Fig. 1b - Relative intensity in Pb of photons causing $^{63}(\gamma\eta)\gamma$.

Mu 780
FIG. 2—RELATIVE INTENSITY IN Pb OF PHOTONS CAUSING $\gamma \eta$ $\gamma \gamma$ AND $\gamma \eta$ $\gamma \gamma$