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SOLID CHEMICAL RADIATION DOSIMETER

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PREFACE

This report presents the accumulated results of investigations carried out to determine the usefulness of the HAP solid chemical system for charged particle dosimetry. Protons from a 160 meV cyclotron and beta rays from a strontium -90 plaque were the radiations studied.

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A. Proton dosimetry using the HAP system.

B. Beta-ray dosimetry using the HAP system.
PROTON DOSIMETRY USING THE HAP SYSTEM*

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The solid phantom chemical dosimeter, which contains halogenated compounds and azo dyes in a paraffin matrix, has been used primarily to measure ionizing electromagnetic radiation. Some limited applications of the HAP system as a method of visualizing and determining dose distribution associated with charged particles have been previously reported. This article presents more precise quantitative data about specific uses of these translucent tissues—approximating formulations in proton dosimetry. In addition to presenting the effect of specific gravity and effective atomic number, there are studies to illustrate dose distribution in phantoms irradiated with proton beams of different sizes. Analyses of the Bragg peak as well as the shaft of the proton beam are included in the experiments.

MATERIAL AND EQUIPMENT

Chloroform or bromoform was the halogenated compound dissolved in a paraffin matrix with one of three azo dyes: p-dimethylaminoazobenzene (PDAB), 4-phenylazodiphenylamine, (PADA), or p-phenylazoaniline (PPAA). Hexylresorcinol and p-phenylazophenol (PPAP) were employed as chemical modifiers or stabilizers.

The proton source was the Harvard synchrocyclotron with an external beam maximum intensity of about $10^{11}$ protons per second with an energy of approximately 160 meV. One-half of the protons could be focused by a quadrupole lens to pass through an area of one centimeter square. Calibration dosimetry was carried out with nitrogen-filled
ionization chambers and a Faraday Cup. In water the full range of the protons was 17.5 centimeters. When actually used the range of the beam was attenuated by various thicknesses of interposed water-equivalent plastic.

The equipment for quantitative determinations of color intensity in the HAP dosimeter was the same as described in detail in a preceding paper. It consisted of a constant-voltage-regulated fluorescent light source, a photoelectric cell to pick up the light transmitted through the dosimeter, and a galvanometer to measure the light intensity.

RESULTS

A. Effect of specific gravity and Z on sensitivity.

Disks of HAP dosimeter, 0.8 cm. thick and 12 cm in diameter, contained $1.2 \times 10^{-4}$ molal of p-dimethylaminoazobenzene with different molal concentrations of chloroform or bromoform. These small round slabs were placed at the front or beam entrance end of 10 cm. more of HAP dosimeter, which constituted additional phantom material. The experimental disks measured the radiation intensity of a 2 x 2 cm. square proton beam as it entered the tissue-approximating material which was also the dosimeter.

Chloroform and bromoform systems of equal specific gravity exhibited color changes of similar intensity to a given dose of radiation. For example, a 0.4 molal bromoform formulation and a 1.5 molal chloroform system had identical densities of 0.95 and about the same sensitivities (Figure 1), in spite of calculated effective Z values of 14 for the bromoform system and 9 for the chloroform formulation. Under these
conditions 2 had little effect on sensitivity.

B. Effect of chemical modifiers on sensitivity.

When hexylresorcinol in a $5 \times 10^{-3}$ molal concentration was added to a chloroform system containing p-dimethylaminoazobenzene sensitivity was definitely enhanced as is pointed out by Figure 2. The addition of a $6 \times 10^{-4}$ molal concentration of p-phenylazophenol to the same hexylresorcinol--chloroform--p-dimethylaminoazobenzene formulation further increased the sensitivity. To change the relative transmittance of the dosimeter from 100 per cent to 50 per cent required 2700 rad for chloroform alone, 920 rad when hexylresorcinol was included and only 730 rad when p-phenylazophenol was an additional modifier.

C. Effects of beam size on surface dose.

Protons passing through portals of 1 x 1 cm., 2 x 2 cm., and 4 x 2 cm. were used to irradiated dosimeter disks as described above under section A. The measured surface dose was not affected by the size of the proton beam. These results indicated little or no back scatter of protons to the surface, for the energies used in this study.

D. Effect of beam size on the geometry and dose at the Bragg peak.

For the remaining proton beam experiments the HAP system was used in the form of 20 x 20 cm. slabs that were 9.5 mm. thick. The protons were so directed into the sides of several closely-packed slabs that the central one, recording the whole shaft and Bragg peak, could be removed, the color intensities quantified and dose curves plotted (Fig. 3).

After irradiation the per cent relative transmittance was measured along the center of the beam. Dose distribution was determined from a standard curve (Figure 4) obtained by irradiating HAP disks of the same
composition with known amounts of protons. Figure 5 shows the dose distribution for three different-sized proton beams. The ratio of absorbed dose at the Bragg peak to the dose at the entering or proximal-most end of the beam was calculated from these data. As presented in the table in Figure 6, there was less dose differential between the shaft and the peak with narrower beams.

Figure 5 also shows that the position of the Bragg peak along the axis of irradiation in the HAP dosimeter was not affected by changes in proton beam size. However, the relative flaring or expansion of the proton beams at their distal ends was greater for small beams. The ratios of beam width at the Bragg peak to the beam width at the entering surface of the phantom dosimeter are given in the table in Figure 7. A very narrow beam manifested the spread of the distal end so much that no point increase or peak dose was found. For such beam sizes the dose distribution indicated the absence of any significant Bragg peak (Fig. 8).

E. Effect of Z on the position of the Bragg peak.

Chloroform formulations of the HAP dosimeter were calculated to have an effective Z of 7 for 0.5 molal, 8 for 1.0 molal and 9 for 1.5 molal concentrations. For bromoform systems the effective Z was determined to be 12 for 0.2 molal and 18 for 0.8 molal concentrations. These different concentrations of chlorinated and brominated hydrocarbons with effective Z ranging from 7 to 18 did not alter significantly the position of the Bragg peak as shown in Tables I and II. In all instances the distal end of the proton beam was found to be at a 12.2 or 12.3 cm. depth within the phantom dosimeter.

In another similar experiment 3 centimeters of water was interposed at the front or beam entrance end of the HAP dosimeter slab. Measurements
Potsaid et al. showed that the Bragg peak was now at a 9.2 cm. depth. Figure 9 illustrates that the distal end of the proton beam in the combined phantom material was at a depth of 12.2 centimeters. This evidently means that the energy loss coefficient for protons in water, certain plastics, and specific HAP dosimeter formulations is about the same.

REFERENCES


-5-
### TABLE I

**EFFECT OF CHLOROFORM CONCENTRATION ON POSITION OF THE BRAGG PEAK**

<table>
<thead>
<tr>
<th>Chloroform (Molal)</th>
<th>0.5</th>
<th>1.0</th>
<th>1.5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Distance from surface (cm.)</td>
<td>12.3</td>
<td>12.2</td>
<td>12.2</td>
</tr>
</tbody>
</table>

### TABLE II

**ROLE OF EFFECTIVE ATOMIC NUMBER ON POSITION OF THE BRAGG PEAK**

<table>
<thead>
<tr>
<th>Z (effective)</th>
<th>8.17</th>
<th>11.82</th>
<th>17.74</th>
</tr>
</thead>
<tbody>
<tr>
<td>Halogenated cpd (Molal)</td>
<td>1.0 chloroform</td>
<td>0.2 bromoform</td>
<td>0.8 bromoform</td>
</tr>
<tr>
<td>Specific gravity</td>
<td>0.93</td>
<td>0.93</td>
<td>1.0</td>
</tr>
<tr>
<td>Distance from surface (cm.)</td>
<td>12.3</td>
<td>12.3</td>
<td>12.2</td>
</tr>
</tbody>
</table>
EFFECT OF $Z$ ON PROTON DOSE MEASUREMENTS

FIGURE 1
EFFECT OF ATOMIC NUMBER ON SENSITIVITY
A has $5.0 \times 10^{-3}$ molal hexylresorcinol. 
B has the same concentration of hexylresorcinol plus $6.0 \times 10^{-4}$ molal p-phenylazophenol (PPAP).
FIGURE 3.

Photograph of the terminal end of a 160 meV proton beam as demonstrated by the HAP dosimeter.
In addition to the chloroform this 9.5 mm thick formulation contained $1.2 \times 10^{-4}$ molal p-dimethylaminoazobenzene (PDAB), $6.0 \times 10^{-4}$ molal p-phenylazophenol (PPAP) and $5.0 \times 10^{-3}$ molal hexylresorcinol.
EFFECT OF PROTON BEAM SIZE ON DOSE ALONG THE CENTRAL AXIS

CHLOROFORM - PDAB-PPAP-HEXYLRESORCINOL SYSTEM
Bragg peak at 12.2 cm depth
Thickness 9.5 mm

FIGURE 5
DOSE DISTRIBUTION ALONG THE CENTRAL AXES OF PROTON BEAMS
EFFECT OF BEAM SIZE ON RATIO OF DOSE AT BRAGG PEAK TO DOSE AT ENTRY SURFACE

<table>
<thead>
<tr>
<th>BEAM SIZE</th>
<th>1 x 1 cm</th>
<th>2 x 2 cm</th>
<th>4 x 2 cm</th>
<th>5 cm circle</th>
</tr>
</thead>
<tbody>
<tr>
<td>B/A DOSE RATIO</td>
<td>1.50</td>
<td>2.00</td>
<td>2.00</td>
<td>3.10</td>
</tr>
</tbody>
</table>

FIGURE 6

SURFACE-BRAGG PEAK DOSE RATIO AS Affected BY PORTAL SIZE
EFFECT OF BEAM SIZE ON RATIO OF BEAM WIDTH AT BRAGG PEAK TO WIDTH AT ENTRY SURFACE

<table>
<thead>
<tr>
<th>BEAM SIZE</th>
<th>1 x 1 cm</th>
<th>2 x 2 cm</th>
<th>4 x 2 cm</th>
<th>5 cm Circle</th>
</tr>
</thead>
<tbody>
<tr>
<td>WIDTH IN CM.</td>
<td>A</td>
<td>1.0</td>
<td>2.0</td>
<td>4.0</td>
</tr>
<tr>
<td>B</td>
<td>1.3</td>
<td>2.4</td>
<td>4.6</td>
<td>5.5</td>
</tr>
<tr>
<td>B / A RATIO</td>
<td>1.30</td>
<td>1.20</td>
<td>1.15</td>
<td>1.10</td>
</tr>
</tbody>
</table>

FIGURE 7
RATIO OF SURFACE-BRAGG PEAK BEAM WIDTH AS AFFECTED BY PORTAL SIZE
Photographic patterns of dose distribution of 160 meV protons resulting from entrance beam widths of A (4.8 mm), B (3.2 mm), C (1.6 mm), and D (0.9 mm), showing a gradual spread of the beam within the HAP dosimeter with an ultimate loss of the Bragg Peak for the narrowest portals of entry. Arrow indicates the beam direction.
COMPARISON OF STOPPING POWER
OF HAP DOSIMETER AND WATER FOR 160 MEV PROTONS

![Graph showing the comparison between HAP dosimeter and water as absorbers of protons.]

**Figure 9**

The solid dosimeter compared to water as an absorber of protons.
BLTRAY DOSIMETRY USING THE HAP SYSTEM*

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This work was supported in part by NASA Grants NsG-719 and NsG-262
In earlier papers the solid chemical dosimeter containing halogenated compounds and azo dyes in a paraffin matrix had been shown to be a valuable method of visualizing radiation effects due to charged particle ionization.\textsuperscript{1-3}

This part of the report pursues applications of similar techniques to dose problems associated with beta ray sources and covers effects of light-source size in the measuring device and of dosimeter thickness on sensitivity. Patterns of beta ray scatter as well as the density on the range of beta rays in the HAP system are included.

MATERIAL AND EQUIPMENT

Chloroform or bromoform was the halogenated compound dissolved in a paraffin matrix with the azo dye, 4-phenylazodiphenylamine (PADA). Hexylresorcinol was employed as a chemical modifier.

The beta ray source was a strontium-90 skin-therapy applicator having a 5 mm. active diameter on a 12.7 mm. diameter overall base and a dose rate of 1250 rep per minute at the surface.

The equipment for quantitative determinations of color intensity consisted of a constant-voltage-regulated fluorescent light source, a photoelectric cell to pick up the light transmitted through the dosimeter, and a galvanometer to measure the light intensity. It was the same as described in a previous paper except that a more sensitive photoelectric cell was used for the beta ray dose measurements.
RESULTS

A. Effect of light-source aperture size on sensitivity.

The HAP dosimeter was made into 2 mm. thick disks containing 0.8 molal bromoform and 2.3 x 10^{-4} molal 4-phenylazodiphenylamine and irradiated with various doses by the strontium-90 applicator. Color changes in the disks were then quantified. All components and geometries of the light source and light measuring device were kept constant except the size of the aperture at the light source. According to the results as presented in Figure 1, dosimeter sensitivity was greater for the smallest light window. To obtain 70% relative transmittance required a dose of 9500 rep for a 10 mm. diameter circular measuring light beam and only 4650 rep for a 0.7 mm. square light beam.

B. Effect of dosimeter thickness on sensitivity.

Disks containing 1.0 molal chloroform, 2.3 x 10^{-4} molal 4-phenylazodiphenylamine and 5.0 x 10^{-3} molal hexylresorcinol were exposed for different periods to the strontium-90 source. Color change was measured as before by determining the ability of each disk to transmit light. In these experiments, both the light-source window and the aperture at the phototube were 0.7 mm. square. Figure 2 shows that for an accurate measure of dose sensitivity was improved up to 5 mm. thickness of HAP dosimeter. When thicker disks were used as dosimeters the sensitivity was not much improved. Primarily this reflected the fact that beta rays from a strontium-90 applicator have a maximum range of only 8 mm. in soft-tissue phantom materials.
C. Effect of density on the range of beta rays in the HAP dosimeter.

Two millimeter thick slabs having different concentrations of chloroform and bromoform, but all with 2.3 x 10^{-4} molal 4-phenylazodiphenylamine and 5.0 x 10^{-3} molal hexylresorcinol constituted the dosimeters for these studies. One sheet of dosimeter was placed between 10 mm. thick paraffin slabs and the whole sandwich was held firmly between plastic plates. The sheet and slabs were then irradiated end on for 0.33, 0.66, 1.0, 3.0, 6.0, 12, 24, and 36 hours. This arrangement permitted the full depth of beta ray penetration to be recorded on the central sheet of dosimeter as would be evidenced by a measurable color change. Figure 3 relates the exposure time to the depths at which radiation-induced color changes stopped and where 100 per cent relative transmittance reappeared along the central axis of irradiation. The maximum depth of beta ray penetration in a 1.0 molal concentration of chloroform was found to be 8 mm. and was reached in 24 hours of exposure.

The absolute beta ray range in HAP dosimeters having different values of density was then determined for the strontium-90 applicator. These were determined in the same way as described above and are presented in Table I. Without much question the maximum range of penetration is lower for dosimeters having the highest densities.

D. Effect of beta ray scatter.

Figure 4 is a photograph of the horizontal planar and vertical cross section patterns obtained for a chloroform system with 4-phenylazodiphenylamine and hexylresorcinol. Measurements indicated that
the greatest diameter of color change was at a depth of 2 millimeters. Area of color changes plotted against exposure time are seen in Figure 5, which shows saturation in 24 to 36 hours. These saturation diameters were 18.0 mm. for 1 molal chloroform systems, 17.5 mm. for 2 molal chloroform and 0.2 molal bromoform systems and 14.0 mm. for 0.8 molal bromoform systems.

Beta ray back scatter was studied using 1 mm. thick HAP formulations containing 1.0 molal chloroform, $2.3 \times 10^{-4}$ molal 4-phenylazodiphenylamine and $5.0 \times 10^{-3}$ molal hexylresorcinol. These thin sheets of dosimeter were irradiated with different thicknesses of phantom placed behind as illustrated in Figure 6. Relative transmittance was determined for each 4 minute exposure. Also included in Figure 6 is the equivalent exposure dose for each point on the curve. These data indicate that saturation occurred from back scattering beta rays with a 5 mm. thickness of phantom and that the amount of back scattering was 30% of the given exposure.

REFERENCES


**TABLE I**

**ATTENUATION OF STRONTIUM-90 RADIATION BY DIFFERENT HAP FORMULATIONS**

<table>
<thead>
<tr>
<th>CHLOROFORM SYSTEM</th>
<th>BROMOFORM SYSTEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Molal</td>
<td>Density</td>
</tr>
<tr>
<td>0.5</td>
<td>0.93</td>
</tr>
<tr>
<td>1.0</td>
<td>0.94</td>
</tr>
<tr>
<td>2.0</td>
<td>0.98</td>
</tr>
<tr>
<td>4.0</td>
<td>1.03</td>
</tr>
</tbody>
</table>
FIGURE 1

Effect of light-source aperture size on sensitivity.
FIGURE 2

Effect of Dosimeter thickness on sensitivity
FIGURE 3.

Relationship of depth of penetration to radiation exposure time.
FIGURE 4.

Photograph of actual dosimeter sections showing the range of lateral scatter as well as the depth penetration.
FIGURE 5.
The area of color change plotted against exposure time.
RANGE OF $^{90}\text{Sr}$ BETA RAY BACKSCATTER
CHLOROFORM 1.0 MOLAL
PADA 2.3 $\times$ 10$^{-4}$ MOLAL
HEXYLRESORCINOL 5.0 $\times$ 10$^{-3}$

FIGURE 6.
Schematic diagram of dosimeter geometry used in studying beta ray backscatter.