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POSITRON RING CAMERAS FOR EMISSION-COMPUTED TOMOGRAPHY

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ABSTRACT

A properly designed circular ring of NaI(Tl) crystals and suitable reconstruction algorithms can provide rapid, high-resolution transverse section images of positron-labeled compounds in the human body. The spatial resolution is limited by the crystal center-to-center spacing and by the positron range in tissue. Using 280 crystals with 10-mm center-to-center spacing and isotopes such as 68Ga, 11C, 18F, or 52Fe, a resolution of 7.5 mm full width at half maximum (FWHM) is possible. Improvements in this technology will involve larger numbers of smaller crystals, more efficient crystals (possibly bismuth germanate), or faster detectors (such as loaded organic scintillators).

1. INTRODUCTION

The first camera specifically designed to produce transverse section images of positron-labeled compounds in the human body was built by Bankowitz, Robertson, and co-workers at Brookhaven National Laboratory in 1962.\(^1\) This innovative instrument employed 32 NaI(Tl) detectors in a circular pattern and was built about 10 years before the mathematical techniques for accurate tomographic reconstruction had been developed.\(^3\) It is now in use at the Montreal Neurological Institute.\(^4,5\)

More recently, a 48-crystal hexagonal system has been built by Ter-Pogossian and co-workers at Washington University at St. Louis\(^6-10\) and a 64-crystal ring system has been built by Cho and co-workers at UCLA\(^11-13\). In addition, the planar detector positron camera systems of Brownell\(^14,15\) and of Maehlhehner\(^16,17\) have been rotated to permit 360° transverse-section tomographic imaging, and the wire chamber-converter positron camera of Perez-Mendez and co-workers can be used for computed longitudinal tomography.\(^18\)

During the past 15 years, beginning with the work of Kuhl, techniques have also been developed for 360° transverse-section imaging of single-gamma radionuclides.\(^19-24\)

2. RING SYSTEMS

2.1 Comparison

In Table I we compare the dimensions, spatial resolution, and sensitivity of three circular positron cameras, the BNL system,\(^1,2\) the UCLA system,\(^11-13\) and our 280-crystal system now under construction.\(^25,26\) The spacing between crystal centers ranges from 30 mm to 10 mm and the resolution FWHM in the reconstructed images is about 75% of this spacing. This is understandable since (a) circular ring systems sample the space at linear intervals of one-half the distance between crystal centers and (b) the sampling theorem states that in order to reliably recover a spatial frequency component, it must be sampled at least twice per cycle. Thus the finest bar pattern that a circular ring system can resolve has a repeat distance approximately equal to the repeat distance of the detectors. Other factors such as (1) the range of positrons in tissue\(^29-30\) and (2) the annihilation angle uncertainty of about 8 mrad FWHM that arises because the positron-electron system does not annihilate at rest\(^29-32\) are less important. For the case of \(^{68} \text{Ga} \) (beta end point 1.9 MeV) in a 100-cm diam ring, these factors contribute about 3 mm FWHM and 2 mm FWHM, respectively. A higher resolution system with a detector center spacing of 3 mm, for example, might approach a resolution of 3 mm FWHM in transverse section, but only by using the lower-energy positron emitters such as \(^{18} \text{F}, \) \(^{11} \text{C}, \) or \(^{52} \text{Fe}. \)

2.2 The Donner 280-Crystal Positron Ring Camera

We are now constructing a system for the high-resolution imaging of positron-labeled compounds in the human body. This system has been physically simulated by two opposing groups of eight crystals viewing positron-filled phantoms rotated on a turntable between them. See Ref. 26 for details and phantom images. We summarize below some of the properties expected for the full ring, based on measurements made with the 16-crystal system.

a. The spatial resolution is best at the center of the ring, where the point-spread function is circular with 7.5 mm FWHM. At 10 cm from the ring center, the point-spread function is elliptical with an 8 mm x 12.5 mm FWHM.

b. The coincident event rate for a 200 μCi point source in air at the center of the ring is 30 000 per sec, including 1700 random coincidences per sec. The coincident event rate for 200 μCi per 1 cm section distributed in a 20 cm cylinder of water is 8600 per sec, including 1400 random coincidences per sec (detection of two unrelated photons within the coincidence resolving time) and 1200 scattered coincidences per sec (detection of two photons from the same positron where one or both have scattered). These random and scattered backgrounds are distributed over a 30-cm field of view and do not present a problem when the activity occupies a small portion of the field.

3. NEW SCINTILLATOR MATERIALS FOR POSITRON COINCIDENCE DETECTION

3.1 Bismuth Germanate (Bi\(_4\)Ge\(_3\)O\(_{12}\))

Recently a new scintillator material (bismuth germanate) has been developed that has several advantages over NaI(Tl) for the detection of annihilation photons\(^33-35\) (Table II). Compared to NaI(Tl), bismuth germanate has a much higher density and atomic number, and thus has a higher gamma-ray detection efficiency. However, bismuth germanate has a much lower scintillation yield than NaI(Tl) and its energy and time resolutions are poorer.\(^15,33-35\)
Table 1. Positron ring systems for transverse section computed tomography.

<table>
<thead>
<tr>
<th>Institution</th>
<th>BNL</th>
<th>UCLA</th>
<th>Donner (LBL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>References</td>
<td>1, 2</td>
<td>11-13</td>
<td>25, 26</td>
</tr>
<tr>
<td>Number of Crystals</td>
<td>32</td>
<td>64</td>
<td>280</td>
</tr>
<tr>
<td>Crystal face</td>
<td>32 mm diam</td>
<td>20 mm diam</td>
<td>8 mm x 30 mm</td>
</tr>
<tr>
<td>Spacing between crystal centers</td>
<td>39 mm</td>
<td>23 mm</td>
<td>10 mm</td>
</tr>
<tr>
<td>Image resolution</td>
<td>~ 40 mm</td>
<td>17 mm&lt;sup&gt;b&lt;/sup&gt;</td>
<td>7.5 mm</td>
</tr>
<tr>
<td>Number of projection integrals measured</td>
<td>~ 2 x 270</td>
<td>2 x 864</td>
<td>14 000</td>
</tr>
<tr>
<td>Diam of crystal ring</td>
<td>40 cm</td>
<td>47 cm</td>
<td>90 cm</td>
</tr>
<tr>
<td>Field of view</td>
<td>~ 30 cm</td>
<td>~ 30 cm</td>
<td>50 cm</td>
</tr>
<tr>
<td>Sensitivity (counts/sec/µCi)&lt;sup&gt;a&lt;/sup&gt;</td>
<td>240</td>
<td>imaging</td>
<td>140</td>
</tr>
<tr>
<td>Status</td>
<td>imaging</td>
<td>in vivo</td>
<td>imaging (phantom imaging)</td>
</tr>
</tbody>
</table>

<sup>a</sup>Coincidence rate for a point source in air at the ring center and a 100 keV threshold.

<sup>b</sup>L. H. Cho (private communication).

Table II. Comparison between NaI(Tl) and Bi<sub>4</sub>Ge<sub>3</sub>O<sub>12</sub> crystals.

<table>
<thead>
<tr>
<th></th>
<th>NaI(Tl)</th>
<th>Bi&lt;sub&gt;4&lt;/sub&gt;Ge&lt;sub&gt;3&lt;/sub&gt;O&lt;sub&gt;12&lt;/sub&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Density (g/cm&lt;sup&gt;3&lt;/sup&gt;)</td>
<td>3.67</td>
<td>7.13&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Atomic number</td>
<td>11, 53</td>
<td>83, 32, 8</td>
</tr>
<tr>
<td>Relative scintillation output</td>
<td>1.0</td>
<td>0.08&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Scintillation peak wavelength (nm)</td>
<td>420</td>
<td>480&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Index of refraction at peak wavelength</td>
<td>1.8</td>
<td>2.15&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Scintillation decay time (nsec)</td>
<td>230</td>
<td>300&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Energy resolution at 662 keV</td>
<td>7% FWHM</td>
<td>15% FWHM&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Time resolution&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.4 nsec FWHM</td>
<td>7.1 nsec FWHM</td>
</tr>
<tr>
<td>Detection efficiency&lt;sup&gt;c&lt;/sup&gt; (20-mm wide crystals)</td>
<td>59% (41%)&lt;sup&gt;d&lt;/sup&gt;</td>
<td>90% (88%)</td>
</tr>
<tr>
<td>Detection efficiency (8-mm wide crystals)</td>
<td>48% (31%)</td>
<td>82% (80%)</td>
</tr>
<tr>
<td>Detection efficiency (4-mm wide crystals)</td>
<td>42% (25%)</td>
<td>72% (70%)</td>
</tr>
</tbody>
</table>

<sup>a</sup>As reported in Ref. 34.

<sup>b</sup>For a 100 keV threshold as reported in Ref. 13.

<sup>c</sup>Calculated detection efficiency for 511 keV photons incident on a bank of rectangular crystals 5 cm deep as shown in Fig. 1. Only single-crystal detections are included.

<sup>d</sup>First percentage: 100 keV threshold on each crystal; percentage in parenthesis: 511 keV threshold.

To determine the relative detection efficiency of NaI(Tl) and Bi<sub>4</sub>Ge<sub>3</sub>O<sub>12</sub>, we wrote a Monte Carlo computer code that traced the interactions of 511 keV photons through groups of crystals. One crystal was uniformly illuminated along a line across its face by a beam parallel to the 5 cm dimension (Fig. 1). The detection efficiencies given in Table II include only those photons that deposited more than the threshold energy (100 keV or 511 keV) in the illuminated crystal and less than the threshold energy in each of the other crystals.

Using 8-mm wide crystals and a 100 keV threshold, we find that bismuth germanate has a 1.7-fold advantage in detection efficiency over NaI(Tl), and thus a 2.9-fold advantage in coincident detection efficiency. The use of narrower 4-mm wide crystals reduces the efficiencies for both materials by a factor of about 0.88, but the ratio between them remains nearly the same.

The photopeak efficiency for NaI(Tl) is relatively low and photopeak energy selection is generally not used in positron imaging. By comparison, bismuth germanate has a very high photopeak efficiency and when used with a high threshold (300 to 400 keV), would have the advantage of rejecting about half of the tissue-scattered annihilation photons while rejecting very few unscattered photons.

We now estimate the true and accidental event rates for rings of NaI(Tl) and Bi<sub>4</sub>Ge<sub>3</sub>O<sub>12</sub> crystals 0.8 x 3 x 5 cm deep. We assume a crystal ring radius c = 40 cm, a shielding slit width S = 2 cm and depth T = 20 cm, and a 200 µCi/cm line source lying on the ring axis in a 20 cm diam cylinder of water (Fig. 2). From Ref. 25 we have:

True unscattered coincident event rate:

\[ C_0 = \frac{\rho a S^2 e^{-1/c} L^2}{4c} \]
Single counting rate:

$$C_s = \frac{\rho a S^2}{\tau} \left[ e^{-\mu L/2} + e^{-\mu L} \right]$$

Accidental event rate:

$$C_a = \frac{f c_s^2 \tau}{2}$$

where $\rho = 200 \mu\text{Ci/cm}$, $a = 37,000$ annihilations per sec per $\mu\text{Ci}$, attenuation factor $e^{-\mu L} = 0.15$, and $f$ = the fraction of crystals in coincidence with each crystal = 0.22. This value of $f$ assures that all points within a 30-cm diam circle are sampled at all angles. The estimated values of $C_o$, $C_s$, and $C_a$, as well as the definitions and assumed values of $e$, $e_p$, $P_p$, and $\tau$ are given in Table III.

In summary, for the positron ring systems of Table III, bismuth germanate has the following advantages over NaI(Tl):

a. For a given amount of activity, the true coincident rate is nearly three times larger and the accidental/true coincidence ratio is nearly two times smaller.

b. For a given accidental/true coincidence ratio, the event rate is about six times larger.

In addition, the reduced penetration of off-axis photons in bismuth germanate provides better spatial resolution toward the edges of the field of view.

3.2 Loaded Organic Scintillators

The ideal scintillator for the coincident detection of annihilation photons would have high density, high atomic number, emit a large number of scintillation photons in a short decay time, operate at room temperature, and be inexpensive. Unfortunately, the high-efficiency scintillators such as NaI(Tl), CsI(Na), and $\text{Bi}_4\text{Ge}_3\text{O}_{12}$, are slow and the fast organic scintillators have low detection efficiency. One notable exception is liquid xenon, $^{37,38}$ which has a detection efficiency similar to NaI(Tl) and is about 10 times faster, but requires an operating temperature of about -100°C.

One possibility for an improved scintillator was recently discussed by Lyons, $^{39}$ based on work with perylene and bromo-perylene. $^{40}$ These compounds are fast organic fluors, whose uv fluorescence is not quenched even by large amounts of iodinated compounds such as 1-iodopropane.

Accordingly, we investigated the light output corresponding to 662 keV Compton electrons for various concentrations of perylene in toluene and for various admixtures of iodinated hydrocarbons. The light output increased with increasing concentration of perylene in toluene and saturated at about $10^{-4}$ molar. At this concentration the light output increased threefold when argon gas was bubbled through the mixture to de-oxygenate it. The output level reached 70% of pilot B $^{41}$ and 20% of NaI(Tl). The output was reduced 5 to 10 fold by only 0.2% admixtures (by volume) of either 1-iodopropane, iodo-
Table III. Comparison between NaI(Tl) and Bi$_4$Ge$_3$O$_{12}$ crystal rings with 8-mm wide crystals.

(see text for details)

<table>
<thead>
<tr>
<th>Quantity</th>
<th>Symbol</th>
<th>NaI(Tl)</th>
<th>Bi$_4$Ge$<em>3$O$</em>{12}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Detection threshold</td>
<td>$E_p$</td>
<td>100 keV</td>
<td>350 keV</td>
</tr>
<tr>
<td>Detection efficiency for 511 keV photons (single-crystal interactions only)</td>
<td>$\epsilon$</td>
<td>0.48</td>
<td>0.80</td>
</tr>
<tr>
<td>Probability of a 511 keV photon scattering on passing through 10 cm water and retaining an energy above $E_p$</td>
<td>$P_p$</td>
<td>0.6</td>
<td>0.3</td>
</tr>
<tr>
<td>Average detection efficiency for above scattered photons</td>
<td>$\epsilon_p$</td>
<td>0.8</td>
<td>0.9</td>
</tr>
<tr>
<td>True unscattered coincident event rate</td>
<td>$C_0$</td>
<td>6400/sec</td>
<td>17 800/sec</td>
</tr>
<tr>
<td>Single counting rate (sum over all crystals)</td>
<td>$C_s$</td>
<td>980 000/sec</td>
<td>850 000/sec</td>
</tr>
<tr>
<td>Coincidence resolving time (full width)</td>
<td>$\tau$</td>
<td>15 nsec</td>
<td>30 nsec</td>
</tr>
<tr>
<td>Accidental coincident event rate (distributed over a 30 cm field of view)</td>
<td>$C_a$</td>
<td>1600/sec</td>
<td>2400/sec</td>
</tr>
<tr>
<td>Accidental/true coincidence ratio</td>
<td>$C_a/C_0$</td>
<td>0.25</td>
<td>0.13</td>
</tr>
</tbody>
</table>

methane, diiodomethane, iodoform, or iodobenzene. Larger admixtures reduced the pulse height still further. All the mixtures had strong UV fluorescence. Thus we found (unfortunately) that while iodinated hydrocarbons do not quench perylene for UV excitation, they do quench it for ionizing radiation.

It is hoped that further work in this area will result in a convenient, fast, high-density scintillator.

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REFERENCES


36. In Ref. 25 the factor 1/2 was omitted.
41. Manufactured by Nuclear Enterprises Inc., San Carlos, CA.
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