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POSITRON EMISSION CAMERA FOR TRANSVERSE SECTION TOMOGRAPHY

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Authors
Derenzo, Stephen E.
Zaklad, Haim
Budinger, Thomas F.

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POSITRON EMISSION CAMERA FOR TRANSVERSE SECTION TOMOGRAPHY

Stephen E. Derenzo, Haim Zaklad, and Thomas F. Budinger

DONNER LABORATORY

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1. INTRODUCTION

We propose an imaging device for the rapid, non-invasive, high resolution (6 mm FWHM), three-dimensional reconstruction of the myocardium in end-systole and end-diastole. The instrument can also be used for transverse axial tomography of the head and abdomen. The major emphasis is to image the accumulation of $^{82}$Rb (positron emitter) in the myocardium after i.v. injection. This isotope has a 75-sec half-life, and can be readily obtained from a portable generator containing $^{82}$Sr, which decays to $^{82}$Rb with a 25-day half-life. Studies of blood flow are also feasible using $^{68}$Ga or $^{82}$Rb. The device we have designed incorporates multiple crystals in a ring and a data collection scheme based on coded light pipes. Transverse tomography of body sections from 5 to 15 mm thick are imaged in 1 to 5 min without rotation. Both transmission and emission modes are envisioned in order to effect compensation for attenuation in quantitative studies and comparison of anatomy to function using algorithms already developed.

More than 100,000 Americans die each year from fatal heart attacks. An essential aspect of prevention is early diagnosis and understanding of the pathophysiologic processes in the myocardium. The instrument we propose presents a method for the non-invasive study of myocardial disease associated with diminution in flow. The instrument also allows the rapid assessment of dyskinesia and can be used in conjunction with a number of isotopes but is basically for use with short half life positron emitters. Good examples are
Rubidium$^{82}$ and Galium$^{68}$ which can be obtained from a generator remote from a cyclotron. $^{18}$F for imaging acute myocardial infarctions is another important application. This device has the potential for imaging not only the heart in end-systole and end-diastole, but for assessing brain flow with 10 second frame times and for imaging distribution of isotopes or density (5% accuracy) in other organs such as the pancreas, prostate, and bladder.
2. BACKGROUND

Early studies by Love and Burch (1957) with Rubidium-86 and Carr and co-workers (1964) with $^{131}$Cs suggested that these alkaline metals are extracted by the myocardium in much the same way as potassium, and can be used to infer coronary flow in accordance with the Sapirstein principle (Sapirstein, 1958). In the past four years there has been an intense effort to develop noninvasive techniques for delineating normal or abnormal myocardial perfusion using the radio-nuclides of potassium, rubidium, and cesium as well as coronary injection techniques using labeled albumin or particle micropheres and xenon. The background work in assessing coronary flow and myocardial functional integrity is shown in Table 1.

These efforts, while notable for establishment of the clinical efficacy of imaging the myocardial accumulation of isotopes, have not given us a substantially useful tool for noninvasive evaluation of coronary flow or myocardial integrity. The reason appears not to be in the selection of isotope so much as in the instrumentation for gated three-dimensional distribution delineation. Our previous work with the positron emitter $^{82}$Rb (Budinger, Yano and Hoop, 1975; and Volkelman et al., 1972) suggests that this isotope given in 3-5 mCi amounts can provide sufficient data for transverse section reconstruction of the human heart in gated studies. Previous work in instrumentation for three-dimensional imaging of nuclide distribution has been limited to the head. This work has been with a rectilinear scanner (Kuhl and Edwards, 1963; Kuhl and Edwards, 1968; Kuhl et al., 1973)
and some studies with phantoms and patients have been made with the gamma camera (Budinger et al., 1974). Recently using $^{129}$Cs, we have been able to image transverse sections of the human myocardium (Fig. 1) and the myocardium of the dog (Fig. 2). The reconstruction of transverse sections was carried out using five different algorithms and six methods for the compensation of attenuation were incorporated in these and other studies (Budinger and Gullberg, 1974). These studies done with the gamma scintillation camera are inadequate because of low statistics, and clumsy patient rotation schemes (many have been explored). Using $^{81}$Rb, we explored the feasibility of gating studies of the myocardium (see Fig. 3) with good results with one view; however, we have not yet combined gating with multiple views due to lack of adequate parallel hole collimation for the 450 - 520 keV photons of $^{81}$Rb and the 600 - 1400 keV emmanations of $^{82m}$Rb contamination present in some preparations.

This problem might be overcome by use of lower energy nuclides such as $^{201}$Tl, which just now is becoming available (Lebowitz et al., 1974). However, this isotope has a 75-hr half-life, and does not provide the characteristics of a positron emitter for transverse section imaging.

Even the best spatial resolution imaging device cannot give in one view a satisfactory representation of the three-dimensional distribution of the isotopes. While tomographic imaging can be accomplished using a moving, focusing collimator (Anger, 1968; Mathieu and Budinger, 1974), or a coded aperture technique (Barrett et al., 1973; Chan et al., 1974; Budinger and Macdonald, 1975),
HUMAN MYOCARDIUM TRANSVERSE SECTIONS

Tc-99m TRANSMISSION

1. RIGHT ATRIUM
2. RIGHT VENTRICLE
3. INTRAVENTRICULAR SEPTUM
4. ANTERIOR PAPILLARY MUSCLE
5. LEFT VENTRICLE

Cs-129 EMISSION CORRECTED FOR TRANSMISSION

XBB 749-6039
DOG MYOCARDIUM
TRANSVERSE SECTION RECONSTRUCTION

TRANSMISSION
Tc-99m

EMISSION
Se-75 METHIONINE

PROJECTION

TRANSVERSE SECTION

XBB 748-4532A

Fig. 2
the result is an image of the plane of interest with continuous planes superposed as a heterogeneous out-of-focus background.

A positron camera can be used in a fashion which allows focusing on one plane (Brownell and Burnham, 1972); however, the adjacent planes will also be present in the image, but out of focus. This type of imaging selected planes with the positron camera is equivalent to back projection or simple superposition.

The Brownell camera at Massachusetts General Hospital has been used for transverse section imaging on phantoms and dogs with good results (Chesler, 1973). These studies involve rotation of the subject between the camera planes. Thus multiple injections are required for sufficient data due to the short half-life of the positron emitters $^{13}$NH$_3$, $^{82}$Rb, and $^{38}$K.

The Anger positron camera, employing a scintillation camera in coincidence with a crystal array, has been in operation for a number of years, but the count rate limitations (~ 1000 sec$^{-1}$) obviate its usefulness for the studies we envision.

A positron camera employing two gas-filled multiwire proportional chambers is being developed at the University of California San Francisco Medical Center and Lawrence Berkeley Laboratory. This system uses lead honeycomb converters and each of the two detector planes has efficiencies from 2.5% to 10% depending upon the number of chambers and converters used. The coincidence rate in air is 330 events per sec. (30-$\mu$Ci source) (10% accidentals). The spatial resolution is 6-7 mm FWHM (Lim et al., 1975). No collimation is provided.
and energy selection is not possible. It appears that in a 20-cm cylinder of tissue, approximately 70% of the coincident events will not provide useful spatial information (Derenzo, 1975).

Another two-plane positron camera system employing two Anger scintillation cameras is being developed at Searle Radiographics, Inc. (Muehllehner, 1975). The resolution is 10 mm FWHM and counting rates in excess of 4000 sec$^{-1}$ are possible in 12 cm water.

Efficient transaxial tomography using the above systems involves patient or detector rotation, and dynamic flow studies are not possible except indirectly through use of the equilibrium image method proposed by Brownell, Ter-Pogossian and others (Jones et al., 1974).

Ring detector systems have been built for positron mode transverse section imaging. Studies are underway at Brookhaven National Laboratory using a 32-detector ring which is 40 cm in diameter. Each detector is a NaI(Tl) crystal of 3.2 cm diameter (Robertson et al., 1973). The resolution and sensitivity of this innovative device are limited by the large size and small number of crystals.

At the Washington University School of Medicine a ring of detectors has been built for positron imaging and transverse section tomography (Ter-Pogossian et al., 1975; Phelps et al., 1975). The device at present is a hexagonal ring of 48 NaI(Tl) detectors, each 5 cm × 5 cm diameter. Preliminary work demonstrates excellent depth insensitivity, quantitative image reconstruction, good contrast, and 14 mm FWHM image resolution. The detectors were stopped down to 2.5 cm diameter in order to achieve this resolution, resulting in a
loss in sensitivity. Gated imaging might be difficult because of the need to rotate the apparatus through many positions for every section to be imaged.

In summary, previous work using various radiopharmaceuticals and imaging devices for assessing the integrity of the human myocardium have laid the important ground work for the device we propose here. We wish to construct a device capable of performing rapid, high resolution images of the heart. In addition this device has application to head and abdomen, without modification.
3. RATIONALE

3.1 Myocardial Accumulation of Rubidium. The overall objective of ascertaining myocardial perfusion by the accumulation of rubidium involves a basic assumption that the amount of nuclide present in the muscle is directly proportional to the flow. This principle (Sapirstein, 1956, 1958) asserts that the fraction of isotope present in each organ is directly related to the fraction of cardiac output that organ receives. This is true to a first approximation for potassium, rubidium, and $^{13}$NH$_3$ where the extraction efficiency is high, and the Sapirstein principle can yield valuable clinical data if we have some means of reaching inside the myocardium to make analyses. The problem is that we can only estimate the isotope concentration because what is normally imaged is a projection of the three-dimensional distribution on a two-dimensional plane. The distribution of isotope in the wall as well as its gradient through the wall are important clinical observations for which there exists no in vivo techniques (Yipintsoi et al., 1973).

3.2 Need for Multiple Views or Equivalent Imaging. Activity in underlying and overlying tissues is superimposed on the myocardial activity and results in poor contrast for single view images; that is, the target-to-nontarget ratio in the object is reduced from a true ratio of 7:1 to less than 2:1 in the projected image. For positive imaging of infarctions using $^{99m}$Tc-tetracycline or $^{99m}$Tc-pyrophosphate, the situation is not as serious because the true ratios are closer to 12:1 and 20:1, respectively. Yet even with these latter radiopharmaceuticals,
methods of avoiding the superposition of rib and sternum activity are most important, and multiple views are needed for quantitative data.

Another reason for the desirability of heart transverse sections is to quantitate the amount of isotope present in order to grade the quantity of accumulated isotope. Using $^{81}\text{Rb}$, which has both positron annihilation gammas and 190 keV photons from $^{81m}\text{Kr}$, it is possible to measure flow by examining the ratio of Rb to Kr. A high ratio indicates relatively greater flow over a low ratio. Our previous experience with this isotope has led to inconsistent results, probably due to the accumulation in the lungs. Transverse section imaging would avoid this problem.

Thus, there are four reasons for going to the design and instrumentation given here:

1. Selectively remove background from under- and overlying tissue.
2. Aid in detection of intramyocardial defect by transverse section display of data.
3. Provide a quantitative method of assessing the concentration of isotope of a μCi per gram basis.
4. Provide a means for evaluating nuclide flow into and out of the myocardium.

3.3 Positron Mode Imaging Rationale. A severe limitation in nuclide imaging has been the very low statistics. The data are limited by the dose to the patient and instrument performance. Whereas, millicurie amounts ($3.7 \times 10^7$ disintegrations per sec) are injected, the presence of scattering and very poor transmission of collimators ($3 \times 10^{-4}$ to $3 \times 10^{-5}$ for isotopes of interest) leave us with little data to work with, particularly in the beating heart where image motion requires
ten times more disintegrations than for no motion imaging. An analysis of this situation leads to the conclusion that in order to achieve our objective, we must get rid of the collimator and use an isotope with a very short half-life to reduce the dose. Positron mode imaging is clearly the answer, and \(^{38}\text{K},^{81}\text{Rb},\) and \(^{13}\text{NH}_3\) are already-proven isotopes.

3.4 Choice of Isotope. Radionuclides of the chemically related alkaline elements potassium, rubidium, and cesium and their analogs have been shown to be of clinical value in the direct assessment of the distribution of myocardial blood flow using noninvasive scintigraphic techniques. High resolution positron scintigraphic techniques appear particularly attractive not only for localization and sizing of myocardial infarctions, but for direct assessment of the distribution of regional myocardial blood flow.

However, of the many radioisotopes of potassium analogs, relatively few are positron emitters. Among these are: \(^{38}\text{K}, 7.7\) min; \(^{79}\text{Rb}, 21\) min; \(^{81}\text{Rb}, 4.6\) hr; \(^{82}\text{Rb}, 1.2\) min; \(^{82m}\text{Rb}, 6.3\) hr; \(^{84}\text{Rb}, 33\) d; and \(^{13}\text{N-ammonia}, 10\) min. Of these nuclides, \(^{79}\text{Rb},^{82}\text{Rb},\) and \(^{13}\text{N-ammonia}\) are relatively free from other gamma rays and are sufficiently short lived to be useful in acute clinical situations in which the patient’s condition is changing rapidly, thus enabling repeated studies, particularly to assess the efficacy of various modes of therapy. Studies using most of these radionuclides require a charged-particle accelerator near the hospital or clinic. An exception is \(^{82}\text{Rb},\) which can be obtained from a long-lived \(^{82}\text{Sr}\) generator (25-d half-life) and is therefore available,
at will, for application in acute clinical situations. In addition, its short half-life offers low radiation exposure to the patient (Budinger, et al., 1974) (Fig. 4).

Both $^{82}$Rb and $^{68}$Ga (available from 280D $^{68}$Ge) can be used to delineate hemodynamic flow through the heart chambers and the brain. $^{68}$Ga is suitable because of its 68 min half-life and availability as a chelate with EDTA. In this form it will remain in the intravascular space.

3.5 Rationale for Geometry and Data Handling Configuration. Previous attempts to image positrons dating from 1951 (Wrenn, et al. 1951, and Brownell and Sweet, 1953) have been somewhat limited by detector saturation. Camera devices also saturate at count rates below those needed for adequate data gathering. This is certainly the case for the Anger positron camera whose efficiency in air is $\sim$ 30 cts sec$^{-1}$ μCi$^{-1}$, but only about 250 μCi can be in the field of view at one time.

The MGH camera (Burnham et al., 1972) can tolerate 3 mCi or more, and has been used to give good myocardial images with $^{82}$Rb (Budinger et al., 1974).

Our approach is to examine all the way around the heart at one time, with a ring of 288 crystals (Figs. 5 & 6). A section 5 to 15 mm thick will be imaged on each injection.
In order to achieve adequate statistics to present a transverse image of the myocardium after a single injection of $^{82}\text{Rb}$ (75 sec half-life), we envision the following:

(a) the heart is 10 cm in the cephalad-caudad dimension, and the myocardial volume is 150 cm$^3$;

(b) the cardiac rate is 72 min$^{-1}$;

(c) the scattering medium is equivalent to a 20 cm diam cylinder of water;

(d) 20 mCi of $^{82}\text{Rb}$ is injected i.v. and 3.5% of this is extracted by the myocardium;

(e) eight separate images are taken, each corresponding to a 0.1 sec interval of the 0.8 sec cardiac cycle;

(f) data are taken for a period of 200 sec after an initial 100 sec wait for the blood pool clearance of $^{82}\text{Rb}$;

(g) as in Table 3, Case I, the imaging sensitivity is 31 events sec$^{-1}$ [µCi per axial cm$^2$]. The resulting initial activity density is 0.7 µCi per $4 \times 4 \times 10$ mm image element and for each of the eight images typically 100 events will be collected per image element.
4. PHYSICAL DESCRIPTION OF THE SYSTEM

The patient is positioned on a specially designed bed that can be inserted into the ring detector (Fig. 6). The bed can be adjusted vertically and is cantilevered from one end. The other end in the field of view of the detector is made to minimize interaction with the gamma rays. (Thin aluminum tubes and canvas back.) The overall dimension of the ring assembly is 170 cm OD and 60 cm ID, and an additional insert can be used to reduce the backgrounds by reducing the ID to 40 cm. The assembly can be tilted 20° from the vertical to allow for an oblique view of the heart.

The detector consists of an array of NaI(Tl) crystals measuring 0.8 cm wide × 3 cm high × 5 cm deep mounted in a ring 80 cm in diameter (Fig. 7). The total number of crystals is 288. The ring is broken into 8 segments of 36 crystals each. Lead shielding is mounted on the inner circle of the detector extending inward to a diameter of 60 cm. An additional collimator (typically 40 cm ID) is shown in broken lines.

A source containing $^{133}$Ba (not shown) is mounted on a rail inside the collimator opening and driven by a motor. The source generates a 60° fan beam, and travels 180° of the periphery. The position of the source is digitized and the counts collected by the detecting crystals are stored for a subsequent computation of the attenuation distribution in the section.
Fig. 7
Co-linear gamma rays due to positron annihilation in the patient scintillate in NaI crystals. Light generated in a crystal exits into coded light-pipes (example in Fig. 8). Nine photomultipliers per segment view this decoding network utilizing a specially designed code to generate a binary word corresponding to the position. A coincidence circuit (example in Fig. 9) checks for coincidence between any crystal and the opposing 62 crystals. This allows a possibility of 8,928 coincident crystal pairs and provides a nearly uniform geometrical sensitivity over the central 30 cm. If coincidence has been found, the sum of the PMT outputs is energy selected and the data is transferred to storage. The stored positional information is processed using a three-dimensional reconstruction program that corrects for the attenuation in the tissue (generated by the traveling source) to reconstruct an image of the section.

4.1 Shielding. The purpose of the lead shielding is two-fold:

1. Provide a mechanism for varying the slice thickness.

2. Absorb gammas originating from tissue external to the field of view.

The shield consists of two concentric rings extending over the crystal ring with an OD = 106 cm and ID = 40 cm; this determines the slice thickness (see Fig. 10). The internal diameter of 40 cm can be varied by removing a part of the assembly and inserting an alternate to fit different patient sizes.
Light Pipe
Connection Scheme

<table>
<thead>
<tr>
<th>Crystal position</th>
<th>PMT</th>
<th>PMT</th>
<th>PMT</th>
<th>PMT</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>$2^0$</td>
<td>$2^1$</td>
<td>$2^2$</td>
<td>$2^3$</td>
</tr>
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<td>1</td>
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</tr>
<tr>
<td>6</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>

Fig. 8
Fig. 9
For a collimator opening $S$ ($S = 1, 2$ or $3$ cm) the crystals can receive radiation from a region considerably larger than $S$ (depending on the dimensions of the collimator). On the other hand, valid colinear coincidences can arise only from a region of thickness $\sim S/2$. The activity external to the section (thickness $S$) is a major cause of undesired background (see section 5.7) motivating us to extend the collimator as close as possible to the patient.

The collimator is assumed to be about 6 cm thick (10 attenuation lengths). As shown in section 5.7 the overall leakage is $10^{-4}$ or 5 counts/sec/(µCi per axial cm) which is a small fraction of the total.

4.2 Choice of NaI(Tl) Crystals. A "standard" crystal manufactured by Harshaw Corp. for the Baird-Atomic $\gamma$-camera measures 9.5 mm $\times$ 9.5 mm $\times$ 38 mm long (Fig. 5). To increase the sensitivity and to offer the option of varying the slice thickness, a 8 mm $\times$ 30 mm $\times$ 50 mm crystal appears more desirable.

4.3 Crystal Position Coding. The detector ring is divided into eight segments of 36 crystals. A position code is created by coupling each crystal to a unique pair of PMTs (Fig. 8). This allows a large number of crystals to be read out by a small number of PMTs (Alvarez, 1960). Moreover, it is possible to reject events where a significant amount of energy has been deposited in more than one crystal of a segment, since in such cases more than two PMTs will respond. This can occur when a gamma ray scatters in the crystals or when two gamma rays are detected in the same segment within the coincidence resolving time.
In general, the maximum number of distinct combinations is given by

\[ \frac{n!}{m! (n-m)!} \]

where \( n \) is the number of PMTs serving each segment and \( m \) is the number of PMTs to which each crystal is coupled. In the example of Fig. 8 the code for a segment containing 6 crystals used \( n = 4 \) and \( m' = 2 \).

The number of PMTs is minimized when \( m \approx n/2 \), while the number of lightpipes is minimized when \( m \) is small. The use of binary and Gray codes will reduce the number of PMTs somewhat further, but will not permit the rejection of multiple detections. The fast digital coding of crystal positions allows each segment to be placed in coincidence with several opposing segments at high rates using a small number of coincidence circuits. Then crystal address codes can be compared to place each crystal in coincidence with a predetermined number of opposing crystals.

For 8 segments of 36 crystals each, the choice of \( n = 9 \) and \( m = 2 \) allows 72 PMTs to serve 288 crystals via 576 lightpipes and 12 coincidence circuits.

4.4 Energy and Time Resolution. The energy resolution of a scintillation detector depends on the amount of scintillation light produced, the light collection efficiency, and the quantum efficiency of the PMT at the relevant wavelengths. A 5 cm diam \( \times \) 5 cm NaI(Tl) crystal can achieve an energy resolution of 8% FWHM for 511 keV gamma rays, but any reduction in the light reaching the PMT (such as by the use of narrow crystals or by coupling the crystal to the PMT via a long lightpipe) will degrade the resolution.
Grenier et al. report an energy resolution of 23% FWHM at 122 keV and 10% FWHM at 662 keV for a 9 × 9 × 38 mm NaI(Tl) crystal coupled directly to a PMT; and a resolution of 50% FWHM at 122 keV when the same crystal was coupled to the PMT via a 61 cm long plastic lightpipe that transmitted 20% of the light. Their data show that, as expected, the width of the photopeak is inversely proportional to the square root of the number of photons reaching the PMT. As the crystals listed in the example of Table 2 are similar in size and would be coupled to the PMTs via similar lightpipes, we can thus expect that at 511 keV the energy resolution will be approximately 50% FWHM × (122 keV/511 keV)²/² = 24% FWHM. Any non-uniformity among the lightpipes will result in a further degradation of the resolution.

The time resolution in NaI(Tl) depends on the same statistical fluctuations that determine the energy resolution. Braunsfurth et al. report a time jitter distribution with a FWHM of 1.3 nsec for 511 keV gamma ray pairs in NaI(Tl), using a system with a good light collection efficiency. We expect that the time jitter distribution of a ring detector system with an energy resolution of 24% FWHM will have a FWHM of 1.3 nsec × (24%/8%) = 3.9 nsec. A 20 nsec coincidence resolving time is assumed in Tables 2 and 3.
4.5 **Readout.** The readout circuits must perform the following:

1) Determine which PMTs have pulses and convert this information into the binary address of the gamma ray interaction point.

2) Reject events where more than two PMTs in a segment have pulses (i.e., where more than one crystal has received a significant amount of energy). For increased sensitivity at the expense of spatial resolution, it may be desirable to include events where three PMTs have pulses (i.e., where two neighboring crystals have received significant energy). This requires that the code be chosen so that all neighboring crystal pairs generate a unique 3-PMT code.

3) Determine whenever two crystal segments have detected gamma rays within a 10 to 20 nsec time interval.

4) Add the PMT outputs for each coincident crystal segment and reject the event if the pulse height does not fall within a preselected window.

5) Reject the event if another gamma ray is detected (in either coincident crystal segment) within the 200 to 800 nsec required to determine the pulse height.

6) Handle gross counting rates of $2 \times 10^5$ sec$^{-1}$ per crystal segment.

7) Collect fan-beam-$\gamma$-ray transmission data for the attenuation correction. Figure 7 shows the block diagram of a circuit that meets many of the above requirements.
4.6 Attenuation Corrections. Corrections for attenuation are needed to obtain good transaxial reconstructions. One possibility is the use of a continuous source of positron emitter around the outside of the patient, as suggested by Phelps et al. (1975). Since the attenuation depends only on the line of flight of the two gamma rays and not on the position of the point of annihilation, the attenuation correction for each pair of coincident detectors is the ratio of the transmitted flux before and after the patient is positioned in the system.

A second possibility is the use of a fan beam source of 7.9-yr $^{133}$Ba (principal gamma energy 356 keV) that travels 180 degrees around the patient while the opposing 62 crystals collect transmission data. As the attenuation cross section is greater at 356 keV than at 511 keV by a factor of 1.16 for water, muscle and bone (Hubbell, 1969), the attenuation correction for 511-keV pairs is determined as the 0.86 power of the ratio of the transmitted flux of 356 keV gamma rays before and after the patient is positioned in the system. It is necessary to know the attenuation to an accuracy of only 10% rather than the 1% accuracy of the EMI scanner, and approximately 100 times fewer counts are needed. Thus the dose is reduced from 1-3 rad to 50-150 mrad. This second method permits greater ease of source collimation to reduce patient exposure, greater data rates, and the use of an isotope of long half life.
5. PROPERTIES OF THE SYSTEM FOR THE DETECTION OF COINCIDENT
ANNIHILATION RADIATION

5.1 Crystal Detection Efficiency. A large slab of NaI(Tl) 5 cm
thick detects 511 keV gamma rays with a gross detection efficiency of
82% and a photofraction of 0.79 (Anger, 1966). We have measured the
pulse height spectrum for the smaller 0.8 cm × 3 cm × 5 cm deep
crystals described here and find that the gross detection efficiency
is also 80%, but the photofraction has a lower value of 0.25 due to
the loss of Compton scattered gamma rays from the sides of the crystals.
In Table 3 we have used a detection efficiency of 50% (100 keV
threshold and single crystal interactions only) and a photopeak
efficiency of 20%.

5.2 Axial Response Function. Since the two annihilation gamma rays
are emitted in nearly opposite directions, true coincident events can
only arise from a disc-shaped region of space (called a "section")
within the detector circle. The thickness of the section is effectively
equal to one-half the corresponding dimensions of the shielding.

A schematic of the detector circle and the collimator circle
used to block emissions from outside the section may be seen in Fig. 10.
The variable Z is the coordinate along the axis of the detector ring.
(Z = 0 at the midplane.) The variable R indicates the distance to
that axis.
The sensitivity is maximal at the median place of the section (Z = 0) and falls to zero at Z = ± S/2 (the top and bottom surfaces of the section). The dependence of the sensitivity along an axis parallel to the Z axis is called the Axial response Function.

Consider a point at a distance R from the central axis of the detector circle. Consider also two detectors located on opposite sides of the point so that the line between them forms an angle θ with a line connecting the point with R = 0 (Fig. 10). Let B₁ be the distance to the closer detector and B₂ be the distance to the farther detector

\[ B₁ = \sqrt{C² - R² \sin² θ} - R \cos θ \]
\[ B₂ = \sqrt{C² - R² \sin² θ} + R \cos θ \]  

(1)
The sensitivity is uniform in the central interval - Z₁ < Z < Z₁ and falls to zero linearly in the intervals - S/2 < Z < -Z₁ and Z₁ < Z < S/2 (Fig. 11), where

\[ Z₁ = \left( \frac{S}{2} \right) \left( \frac{B₂ - B₁}{B₂ + B₁} \right) \]  

(2)

Writing the coincident detection probability at Z, R, θ as f(Z, R, θ), we have

\[ f(Z, R, θ) = \begin{cases} \left( \frac{S+2Z}{S-2Z₁} \right) \left( \frac{ε^2S}{2B₂} \right), & \frac{S}{2} < Z < -Z₁ \\ \frac{ε^2S}{2B₂}, & Z₁ < Z < Z₁ \\ \left( \frac{S-2Z}{S-2Z₁} \right) \left( \frac{ε^2S}{2B₂} \right), & Z₁ < Z < S/2 \\ 0, & Z < -\frac{S}{2} \text{ or } Z > S/2 \end{cases} \]  

(3)

where ε is the detector efficiency.
Fig. 11
The axial response function normalized to $f(Q,R,\theta)$ and averaged over $\theta$ is given by:

$$F_A(Z,R) = \frac{1}{2\pi} \int_0^{2\pi} \frac{f(Z,R,\theta)}{f(0,R,\theta)} \, d\theta$$  \hspace{1cm} (4)$$

This quantity is plotted in Fig. 12 for the case $C = 40$ cm, $S = 2$ cm for $R = 0$ cm, 10 cm and 20 cm.

5.3 Sensitivity. The event rate for the simultaneous detection of two annihilation gamma rays may be calculated as the following product:

$[\text{contributing activity}] \times [\text{probability of photon #1 reaching a detector}] \times [\text{probability of photon #2 reaching a detector}] \times [\text{product of detector efficiencies}]$. This formulation is used below for both un scattered gamma rays (Eq. 5) and scattered gamma rays (Eqs. 15 & 18).

The unscattered coincident event rate $C_0$ of a ring detector is approximated by:

$$C_0 = \left[ \frac{\rho a S}{2} \right] \left[ \frac{S e^{-\mu L}}{2c} \right] \left[ \varepsilon^2 \right] = \left[ \frac{\rho a S^2 e^{-\mu L}}{4c} \varepsilon^2 \right]$$  \hspace{1cm} (5)$$

where

$\rho = \text{activity density in } \mu\text{Ci per axial cm}$

$a = 37,000 \text{ annihilations per sec per } \mu\text{Ci of } \beta^+ \text{ (conversion factor)}$

$S = \text{exposed dimension of detector ring along ring axis (determined by width of shielding slit) (see Fig. 10)}$

$e^{-\mu L} = \text{probability that neither gamma ray scatters in the tissue}$

$\mu = \text{narrow beam attenuation coefficient for } 511\text{-keV gamma rays in tissue (in water } \mu = 0.10 \text{ cm}^{-1})$

$L = \text{tissue thickness}$

$c = \text{radius of detector ring}$

$\varepsilon = \text{detection efficiency for } 511\text{-keV gamma rays}$. 
Fig. 12

Axial response vs. Position (cm)

- 20 cm (FWHM=14 mm)
- 10 cm (FWHM=12 mm)
- 0 cm (FWHM=10 mm)

Position (cm)

XBL749-4227
The first bracket is the contributing activity; the second bracket is the probability that both gamma rays reach the detector ring without having scattered; and the third bracket is the probability that both gamma rays are detected and pass the pulse height threshold.

For the case 1 of Table 3, $C = 40 \text{ cm}$, $S = 2 \text{ cm}$, $E = 50\%$, $\mu L = 2$ (20 cm diam cylinder of tissue) the sensitivity is:

$$C_0 = \left(\frac{37,000 \times 2}{2}\right) \left(\frac{2e^{-2}}{2 \times 40}\right)(0.50)^2$$

$$= 31 \text{ events/sec/}[\mu \text{Ci of } \beta^+ \text{ per axial cm}]$$

We assume that the source distribution lies within a distance $R$ from the ring axis given by:

$$R = c \sin (f\pi/2)$$

(7)

where each detector (of a total of $N$) is in coincidence with the $fN$ opposing detectors. For a circular ring the geometrical sensitivity is nearly uniform within the radius $R$. We also assume that the detectors are densely packed around the ring and that the shielding blocks only gamma rays external to the section being imaged.

In Table 3 we have calculated the sensitivity $C_0/\rho$ and many other parameters for a particular positron ring detector system and a tissue thickness of 20 cm ($\mu L = 2$). In Case I the event rate is 6000 events per sec for $\rho = 200 \mu \text{Ci per axial cm}$. Thus if the section is to be resolved into a grid of $40 \times 40$ pixels, 375 events can be collected per pixel over a 100-sec time period.
The greater imaging sensitivity of the positron ring detector over conventional single gamma imaging arises primarily from a greater solid angle. The ring detector has a solid angle factor of $S/2c$ which is $2.5 \times 10^{-2}$ for the example of Table 3. In single gamma imaging only a small area of the crystal ($\pi d^2/4$ for a resolution distance $d$) receives gamma rays from sources at a typical imaging distance $D$. For low photon energy, the resulting solid angle factor is approximately $(\pi d^2/4)/(4\pi D^2) = 5.6 \times 10^{-5}$ for $d = 6$ mm and $D = 20$ cm. Thus in these examples the positron ring detector has approximately 450 times greater solid angle than single gamma imaging with the additional benefit of transaxial tomography without rotation.

5.4 Effect of the Range of Positrons in Tissue. The line spread function for positrons in water has been measured for a variety of isotopes (Cho et al., 1975). For the worst case ($^{82}$Rb, 3 MeV beta end point), the range of the positrons introduces an rms of about 1.5 mm.

5.5 Effect of Deviations from 180° Line of Flight. In most solids and liquids, annihilation occurs before the positron has fully thermalized. Typical energies are 10 to 20 eV and result in angular distributions about 180° with FWHM of 6-9 mr. Specific examples include water at +4°C (FWHM = 8.5 mr, FWO.1M = 18 mr) (Colombino et al., 1965) and calcium metal (FWHM 6.5 mr, FWO.1M = 13 mr) (Stewart 1957). For the positron ring system described here (crystal radius 40 cm) an angular uncertainty with FWHM of 8 mr corresponds to a position uncertainty with FWHM 1.6 mm.
5.6 Transaxial Resolution Function. Each point within the section will be viewed by many pairs of detectors and for points not on the central axis of the detector circle, each pair will have a different resolution. This difference is due to the fact that the effective size of a detector depends on the viewing angle. Detector pairs disposed along a diameter (such as III and IV in Fig. 13) view point \( p \) with better resolution than those disposed along a chord (such as I and II in Fig. 13).

Let us assume that each detector has a length \( L \) (in the \( R \) direction), a height \( H \) (in the \( Z \) direction) and a width \( W \). The detector circle radius is \( C \). For a parallel group of rays having an angle \( \theta \) with respect to the length of the detector (Fig. 14), the path length \( D(x, \theta) \) through the detector is given by the following equations:

For \( \theta < \theta_c \)

\[
D(x, \theta) = \begin{cases} 
0 & \text{for } x < 0 \text{ or } x > x_{\text{max}} \\
\frac{x}{\sin \theta \cos \theta} & \text{for } 0 < x < L \sin \theta \\
\frac{L}{\cos \theta} & \text{for } L \sin \theta < x < W \cos \theta \\
\frac{x_{\text{max}} - x}{\sin \theta \cos \theta} & \text{for } W \cos \theta < x < x_{\text{max}}
\end{cases}
\]  

(8)

where \( x_{\text{max}} = L \sin \theta + d \cos \theta \)

and \( \theta_c = W/L \)
For $\theta > \theta_c$

\[
D(x, \theta) = 0 \quad \text{for } x < 0 \text{ or } x > x_{\text{max}}
\]

\[
D(x, \theta) = \frac{x}{\sin \theta \cos \theta} \quad \text{for } 0 < x < W \cos \theta
\]

\[
D(x, \theta) = \frac{W}{\sin \theta} \quad \text{for } W \cos \theta < x < L \sin \theta
\]

\[
D(x, \theta) = \frac{x_{\text{max}} - x}{\cos \theta \sin \theta} \quad \text{for } L \sin \theta < x < x_{\text{max}}
\]

The absorption path $A(x, \theta)$ due to passage through other detectors is given by the following equations:

\[
A(x, \theta) = 0 \quad \text{for } x < W \cos \theta
\]

\[
A(x, \theta) = \frac{x - W \cos \theta}{\cos \theta \sin \theta} \quad \text{for } W \cos \theta < x < x_{\text{max}}
\]

We write the photopeak detection probability for a gamma ray at angle $\theta$ as

\[
P(x, \theta) = \left[ \text{EXP}(-A(x, \theta)\mu_A) \right] \left[ 1 - \text{EXP}(-D(x, \theta)\mu_D) \right]
\]

where $\mu_A$ is the coefficient for absorbing more than 511 keV - $E_p$, $\mu_D$ is the coefficient for absorbing more than $E_p$, and $E_p$ is the effective pulse height threshold for each crystal. The first bracket is the probability that a 511 gamma ray will emerge from a thickness $A(x, \theta)$ of NaI and retain an energy greater than $E_p$, while the second bracket is the probability that a 511 gamma ray will deposit more than $E_p$ on passing through a thickness $D(x, \theta)$ of NaI. For a threshold $E_p = 410$ keV, we estimate $\mu_A = 0.25 \text{ cm}^{-1}$ and $\mu_D = 0.12 \text{ cm}^{-1}$. 

The line spread function $g(R,x)$ at a distance $R$ from the ring axis is given by

$$g(R,x) = \int_0^{2\pi} \frac{d\theta}{2\pi} \int_{-\infty}^{\infty} dx' P(x + x', \theta) P(x - x', \theta) \left( \frac{S^2/2}{B_1 + B_2} \right)$$

where $\sin \theta = \frac{R}{c} \sin \phi$, and $x' = x' B_2 / B_1$. It is assumed that in the interval $0 < x' < x_{\max}$, $\theta$ is essentially constant. This is true whenever $W \ll B_1$.

The function $g(R,x)$ is plotted in Fig. 15 for $R = 0$ cm, 10 cm, and 20 cm under the assumptions: $W = 0.8$ cm, $L = 3.8$ cm, $C = 40$ cm, $\nu_A = 0.25$ cm$^{-1}$, $\nu_D = 0.12$ cm$^{-1}$. Although the detection probability $P(x,\theta)$ is not a symmetric function in $x$, we see that $g(R,x)$ is quite symmetric about $x = 0$. The FWHM of $g(R,x)$ is plotted in Fig. 16 as a function of $R$.

5.7 Shielding and Backgrounds. Shielding is essential for blocking gamma rays emitted by activity external to the section being imaged. Furthermore, varying the width of the shielding slit allows the section thickness to be varied. True unscattered coincident events can arise only from a positron activity $^{\frac{\rho S}{2}}$ within the section, but scattered coincident events and single counts (both of which contribute to the background) can arise from an activity that is effectively $^{\frac{\rho S}{T}}$, where $T$ is the depth of the shielding slit (Figs. 10, 17, 18).

There are two additional methods available for rejecting scattered gamma rays. The coincidence requirement (i.e., that each of the $N$ detectors be in coincidence with the $fN$ opposing detectors) establishes an approximate upper limit $\theta_f = f\pi$ on the scattering angle. This
Fig. 15

-20 cm (FWHM=8.0 mm)
-10 cm (FWHM=5.5 mm)
-0 cm (FWHM=4 cm)
(a) Axial response for colinear $\gamma$-ray pairs

\[
\begin{align*}
\text{(area } &= \frac{S}{2} \text{)} \\
&
\end{align*}
\]

(b) Axial response for single $\gamma$-rays or non-collinear pairs

\[
\begin{align*}
\text{(area } &= \frac{SC}{T} \text{)} \\
\text{Axial response for single } \gamma \text{-rays or non-collinear pairs}
\end{align*}
\]

Fig. 17
approximation is realistic, as the size of the scattering medium is usually small compared to the diameter of the detector ring. The second method is the use of a pulse height threshold that rejects scattered gamma rays with an energy below $E_p$. In single Compton scattering this corresponds to a maximum scattering angle $\Theta_p$ (AIP Handbook, 1972, Eq. 8 e-5):

$$\cos(\Theta_p) = 2 - \frac{511 \text{ keV}}{E_p}$$  \hfill (13)

For plot, see Fig. 19.

For scattered coincident events the effective maximum scattering angle $\Theta_m$ is the smaller of $\Theta_f$ and $\Theta_p$. (For single counts the maximum scattering angle is always $\Theta_p$). The effective minimum scattered gamma ray energy $E_{m}$ corresponding to $\Theta_m$ is given by:

$$E_m = \frac{511 \text{ keV}}{2 - \cos(\Theta_m)}$$  \hfill (14)

**Scattered Coincident Backgrounds.** The rate of scattered coincident events where only one member of the annihilation pair scatters is approximated by:

$$C_1 = \left(\frac{\rho a S c}{T}\right) \left[\frac{S(1-P_s')}{c}\right] \left[\frac{P_s g_1 S}{c}\right] \left[\epsilon \epsilon_m\right] = \frac{\rho a S^3 g_1 c \epsilon \epsilon_m P_s (1-P_s)}{T_c}$$  \hfill (15)

where $P_s = 1 - e^{-\mu L/2}$, the probability a 511 keV gamma ray will scatter on passing through half the tissue,

$\epsilon_m$ is the average detection efficiency for gamma rays between energy $E_m$ and 511 keV, and $g_1$ is a solid angle factor. The first bracket is the contributing activity; the second bracket is the probability that either gamma ray reaches a detector without scattering;
the third bracket is the probability that the other gamma ray scatters and reaches one of the opposing (coincident) detectors; and the fourth bracket is the probability that both gamma rays are detected and pass the pulse height threshold. This expression assumes that the activity is near the center of the scattering medium, that the scattering takes place close to the axis of the ring, and that single scattering predominates.

The dimensionless quantity \( g_1 \) describes the fraction of Compton scattering angles that fall within the angular strip \( S/c \) wide and \( 2\theta_m \) long:

\[
g_1 = \frac{\int_{-\theta_m}^{\theta_m} \int_0^{2\pi} \frac{d\sigma}{d\Omega} \sin \theta \, d\theta \, d\Omega}{\int_0^{2\pi} \int_0^{\pi} \frac{d\sigma}{d\Omega} \sin \theta \, d\theta}
\]

where \( \frac{d\sigma}{d\Omega} \) is the differential cross section for Compton scattering of 511 keV gamma rays on free electrons, given by the Klein-Nishina formula (AIP Handbook, 1972, Eq. 8e-13) (for plot see Fig. 20):

\[
\frac{d\sigma}{d\Omega} = 11.91 \times 10^{-26} \text{ cm}^2/\text{str/electron} \left( \frac{1 - \cos \theta + \cos^2 \theta - \frac{1}{3} \cos^3 \theta}{(2 - \cos \theta)^3} \right)
\]

and \( \theta \) is the \( \gamma \)-ray scattering angle. See Table 4 for values of \( g_1 \) as a function of \( \theta_m \).

The rate of scattered coincident events \( C_2 \) where both members of the annihilation pair have scattered is approximated by:

\[
C_2 = \left[ \frac{\rho a S c}{T} \right] \left[ \frac{S_p}{2c} \right] \left[ \frac{p_s g_2 S}{c} \right] \left[ \varepsilon_m^2 \right] = \frac{\rho a S^3 g_2 \varepsilon_m^2 p_s^2}{2 T c}.
\]

The first bracket is the contributing activity, the second bracket is the probability that one gamma ray scatters and reaches the detectors;
Fig. 20

\( \frac{d\sigma}{d\Omega} \) (10^{-16} cm²/sr)

\( \gamma \)-ray scattering angle (deg)
the third bracket is the probability that the other member of the pair also scatters and reaches one of the opposing (coincident) detectors; and the fourth bracket is the probability that both gamma rays are detected and pass the pulse height threshold. Assuming that single scattering predominates for each gamma ray, \( g_2 \) is given by

\[
g_2 = \frac{\int_{-\theta_m}^{\theta_m} \frac{d\sigma^*}{d\Omega} d\theta}{\int_0^{\pi} \frac{d\sigma^*}{d\Omega} \sin\theta d\theta}
\]

where \( \frac{d\sigma^*}{d\Omega} \) is the autocorrelation of \( \frac{d\sigma}{d\Omega} \). See Table 4 for approximate values of \( g_2 \) as a function of \( \theta_m \). The total background rate \( C_b \) from scattered coincident events is given by:

\[
C_b = C_1 + C_2 = \frac{\rho \sigma S^3}{2Tc} \left[ 2g_1 \epsilon \epsilon_m \epsilon_p (1-p_s) + g_2 \epsilon_m^2 \epsilon_p^2 \right].
\]

Note that the backgrounds \( C_1 \) and \( C_2 \) are not associated with time resolution but with scatter geometry and energy resolution. As expected, the ratio of scattered coincident events \( C_b \) to unscattered coincident events \( C_0 \) is lowest for "good scatter geometry" (i.e., small S/T).

**Associated Gamma Rays.** Many important positron emitting isotopes (such as \(^{82}\)Rb and \(^{68}\)Ga) have accompanying gamma rays that are emitted essentially in time coincidence with the 511 keV gamma ray pair. The rate of background events \( C_\gamma \) where an annihilation gamma ray is detected in coincidence with an accompanying gamma ray is approximated by:

\[
C_\gamma = \left[ \frac{2 \rho a S c}{T} \right] \left[ \frac{f_s^2}{4c^2} \right] [F \epsilon] [\epsilon m \epsilon]
\]

where \( F_\gamma \) is the probability of emission of the accompanying gamma ray.
and $\varepsilon_y$ is its detection efficiency. This expression is an overestimate in that it includes all detected gamma rays, whether they have scattered in tissue or not (i.e., it does not consider that scattered gamma rays have a reduced chance of producing pulses in the pulse height window). The first bracket is the rate of emission of 511 keV gamma rays, the second bracket is the probability that a 511 keV gamma ray reaches the detector ring while an accompanying gamma ray reaches one of the opposing (coincident) detectors, and the third bracket is the probability that the accompanying gamma ray is emitted and both gamma rays are detected. For the example of Table 2 and $^{82}$Rb,

$$C_y = \left(\frac{2 \times 200 \times 37}{20} \times \frac{0.22 \times 2}{4 \times 40^2}\right) \times (0.1 \times 0.3 \times 0.43) = 53 \text{ counts/sec} \quad (22)$$

**Single detection backgrounds.** The single counting rate $C_s$ is given by:

$$C_s = \left[\frac{2 \rho \alpha S \text{Sc}}{T}\right] \left[\frac{S}{2c}\right] \left[\varepsilon(1-P_s) + \varepsilon_p P_p\right] = \frac{\rho \alpha S^2}{T} \left[\varepsilon(1-P_s) + \varepsilon_p P_p\right] \quad (23)$$

where $P_p$ is the probability that a gamma ray will scatter in tissue and retain an energy greater than $E_p$ (i.e., the pulse height threshold) and $\varepsilon_p$ is the average detection efficiency for such gamma rays. The first bracket is the contributing activity; the second bracket is the solid angle factor of the detector ring; and the third bracket is the sum of the probability for not scattering and being detected and the probability for scattering and being detected. For the single Compton scattering of 511-keV gamma rays,

$$P_p \approx P_s \left[\frac{5/2 + 3 \ln \alpha + 1/\alpha - 3\alpha - \alpha^2/2}{40/9 - 3 \ln 3}\right] \quad (24)$$

where $\alpha = E_p/511$ keV and $1/3 \leq \alpha \leq 1$. The quantity in brackets is the
probability that the scattered photon will have an energy greater
than $E_p$, derived by integrating Eq. 8e-21 of Gray, The AIP Handbook,
1972.

Accompanying gamma rays can also contribute a background counting
rate $C_{SY}$:

$$ C_{SY} = \left( \frac{\rho a S \epsilon}{1} \right) \left[ \frac{S}{2c} \right]^2 \left[ F \right] \left[ E_{\gamma} \right] $$

(25)

This expression includes all detected accompanying gammas, whether they
have scattered in tissue or not. For the example of $^{82}$Rb and Table 3,

$$ C_{SY} = \left[ \frac{200 \times 37,000 \times 2 \times 40}{20} \right] \left[ \frac{2}{2 \times 20} \right] [0.1 \times 0.3] $$

$$ = 44,400 \text{ counts/sec}.$$

Two other single detection backgrounds are also present, the rate
of PMT noise pulses within the pulse height window (negligible and not
considered further), and $C_L$ the rate for the process in which a gamma
ray penetrates or scatters in the lead shielding, is detected and
passes the pulse height threshold. In calculating $C_L$, we assume that
the detectors are shielded by 10 attenuation lengths of lead (6.1 cm)
in all directions where appreciable activity is present. The probability
of penetration (no interaction in the lead) is $e^{-10} = 4.5 \times 10^{-5}$.
To include the gamma rays that scatter in the lead and escape, we use
the buildup factor of 2.3 from Goldstein et al. (1954). As shown in the
same reference, 75% of the scattered photons escaping 10 attenuation
lengths of lead have energies above 400 keV. Thus the overall
leakage probability (penetration plus scatter) is $P_L = 1.0 \times 10^{-4}$.

We note that the product of the detection efficiency and the solid angle
subtended by the detectors is very nearly the same for points within 30 cm of the midplane. We therefore use the detection efficiency and solid angle associated with the midplane. In this case we also assume that the available activity extends over 60 cm of the central axis.

\[ C_L = (2\alpha \times 60) \left( \frac{S}{2\sigma} \right) (\varepsilon) (P_L) \]  

(27)

In the example of Table 3,

\[ C_L = (2 \times 200 \times 37,000 \times 60) \left( \frac{2}{2 \times 40} \right) (.50) (1.0 \times 10^{-4}) \]

= 1100 counts/sec.

Accidental Coincidence Background. The accidental coincidence rate \( C_a \) is given by:

\[ C_a = fC_s^2t/2 \]

(29)

where \( t \) is the time resolution. It is of importance to calculate the maximum rate \( \hat{C}_o \) for a certain fraction \( n = C_a/C_o \) of accidental coincidences:

\[ \hat{C}_o = \frac{nT^2 \varepsilon^4 (1-P_s)^4}{8 ftc^2 \left[ \varepsilon(1-P_s) + \varepsilon_p P_p \right]^2} \]

(30)

As expected, the maximum event rate is enhanced by good time resolution, good shielding, and good detection efficiency. It is important to reduce \( f \) to the point where all (or almost all) of the activity lies within the radius \( R \) (Eq. 7). Reducing \( f \) below this value will reduce the signal \( C_o \) approximately as \( f^2 \) (without effecting the single counting rate \( C_s \)) and actually cause \( \hat{C}_o \) to decrease.
For the example of Case I of Table 3 and for a 20% accidental coincidence rate ($\eta = 0.20$), we have

\[ \hat{C}_o = 3500 \text{ events per sec} \]

and

\[ \rho = 110 \mu\text{Ci per axial cm.} \]

The profound effect that the scattering medium has on the maximum event rate can be seen by noting what these values would be in the absence of the scattering medium (i.e., $P_S = P_p = 0$):

\[ \hat{C}_o = 350,000 \text{ events per sec} \]

and

\[ \rho = 1500 \mu\text{Ci per axial cm.} \]

One should note that: (a) increasing $T$ (whenever possible) by extending the collimator closer to the patient reduces all backgrounds without reducing the sensitivity; (b) increasing the section thickness $S$ increases the sensitivity as $S^2$, but the scattered coincident background increases as $S^3$; (c) the maximum event rate $\hat{C}_o$ (as limited by accidentals) is not a function of $S$. In Table 3 we have calculated these backgrounds for a particular activity level and for several pulse height windows. While these backgrounds are 30-45% of the total number of events, they are fairly uniform over the entire image and contribute little relative to regional concentrations of activity (e.g., in the myocardium).
5.8 **Deadtime.** Deadtime arises from the limited speed of both the scintillation process and the electronic circuits. After the interaction of a gamma ray in NaI(Tl), 800 nsec is required to collect 90% of the available scintillation light (91% of the light is emitted with an exponential time constant of 218 nsec and 9% is emitted with an exponential time constant of 1340 nsec) (Tidd et al., 1973). To preserve pulse height information to within 10% at high counting rates, it is necessary to reject all counts that occur within approximately 800 nsec of another count. This requirement results in a paralyzing deadtime of 800 nsec for each coded segment of the ring. The circuits that perform the pulse height selection may also introduce a paralyzing deadtime, but by proper design it is possible to limit the overall paralyzing deadtime \( t_p \) to 1 µsec (Blatt et al., 1968). The circuits that handle the time coincidence and position information introduce a non-paralyzing deadtime \( t_n \) of \(< 200 \) nsec, but this does not result in an appreciable loss of counts, provided the circuits are properly designed and \( t_n \leq t_p \).

In Cases II and III of Table 3, where a pulse height selection is imposed, the fraction of time that each 36 crystal segment is available is \( e^{-Gt_p} \), and the fraction of counts \( f_c \) lost per segment is given by:

\[
f_c = 1 - e^{-Gt_p} .
\]  
\[(31)\]

\( G \) is the gross counting rate for each segment \((1.3 \times 10^5)\), \( t_p = 1 \) µsec, and the fraction of counts lost \( (f_c) \) is 12%. As a coincident event
requires that two segments be available for counting (probability = $e^{-2Gt_p}$),
the fraction of coincident events $f_e$ lost is given by

$$f_e = 1 - e^{-2Gt_p}$$

and equal to 23% for this example.

In Case I of Table 3 no pulse height selection is imposed, and we assume that the paralyzing deadtime can be reduced to 200 nsec. In this case the fraction of counts lost ($f_c$) is 2.5% and the fraction of events lost ($f_e$) is only 5%, but the backgrounds are larger (see Table 3).
6. DATA ANALYSIS

6.1 Interface. The procedure involved in the reconstitution of the coincidence events involves interfacing a small digital computer to the output from the imaging device. The data position information is configured in 16-bit digital words to identify which of the 8,928 crystal pairs were in coincidence for the particular event. The data are stored in one of two modes, histogram or list, depending upon the final configuration for the microprocessor which will be used to configure the digital words. A number of schemes have been envisioned. One scheme is to calculate the projection angle and ray position by simply noting the relative angular positions of the coincident detectors \( \theta_1, \theta_2 \) with respect to some reference line. The angle of the projection plane \( 90^\circ \) orthogonal to the line of position somewhere on which the event occurred is given by

\[
\theta_1 + \frac{\theta_2 - \theta_1}{2}
\] 

(33)

With this projection angle and a knowledge of the position of one of the detectors on the periphery, it is straightforward to calculate the ray \( P_{k\theta} \) associated with the event. Data are then histogrammed in terms of equivalent projections by adding for each angle \( \theta \) the events that belong in similar \( k \)'s. By this means a projection of data will be reconstituted and appear similar to data which would have been obtained if 144 views were made using a parallel hole collimator.
6.2 Reconstruction Algorithms. Once the data have been accumulated, both transmission and emission, the reconstruction of the transverse section will proceed by one of five techniques:

1. iterative least squares
2. Fourier methods
3. convolution hardwire technique
4. direct hardwire technique
5. Marr's technique.

We have explored various algorithms using emission and transmission (Budinger and Gullberg, 1974a) and are still not satisfied which technique would be optimum for our system. The reason for this is that new devices such as hardwire two-dimensional Fourier transform packages might favor the Fourier two-dimensional technique over the Fourier technique, or the availability of charged coupled computing devices (Tiemann et al., 1974) and read only memory, or microprocessors might lead to the algorithm employing hardwired electronics rather than a digital computer. We have been actively working in this field for a number of years, and see a great potential in a microprocessor approach.

Comparison of some of the techniques is shown as Fig. 21. Figure 22 shows a difference between the result of reconstruction of 36 projections on the letters L O V on a 64 × 64 array using the back projection of filtered projections (convolution) technique and the iterative least squares technique. The Fourier filter technique does not handle noise well but is 80 times faster to implement than the iterative least squares technique. We have explored techniques of applying a Butterworth filter in digital form to the back projection
Fig. 21
FILTERED BACK PROJECTION

LEAST SQUARES

Fig. 22
of filtered projections technique with some improvement. The noise associated with positron coincidence detection scheme might be of less importance than that which we have experienced with conventional gamma imaging. Thus the filtered back projection technique with roll-off filters we have explored might be adequate for this reconstruction. Shepp and Logan (1974) have also shown a significant noise suppression of 6 db in the reconstruction using a spatial average (3-point weighting average) with the convolution technique.

**Direct Matrix Method.** Iteration procedures are designed to minimize the square of the weighted differences between the observed data and the expected distribution of events with a given concentration of radionuclide,

\[
\widetilde{R}(A) = \sum_{k,\theta} \left[ \frac{P_{k\theta} - R_{k\theta}(A)}{\sigma_{k\theta}} \right]^2
\]

(34)

where \( P_{k\theta} \) is the measured projection,

\( \sigma_{k\theta}^2 \) is the variance in \( P_k \)

\( R_{k\theta}(A) \) is the expected projection for concentration \( A \).

\( R_{k\theta}(A) \) is a linear function of \( A \) and can be written as

\[
R_{k\theta}(A) = \sum_i A_i F_{ki}^{k\theta}
\]

(35)

where \( i \) sums over pixels. Representing \( k \) and \( \theta \) by the single index \( j \) we get,

\[
R_j(A) = FA = \sum_i F_{ji} A_i
\]

(36)

\[
\widetilde{R}(A) = \sum_j \left[ \frac{P_j - R_j}{\sigma_j} \right]^2 = \sum_j \left[ \frac{P_j - \sum_i F_{ji} A_i}{\sigma_j} \right]^2
\]

(37)
This can be written in vector notation as follows,
\[ \tilde{R}(A) = (P - FA)^T \phi^{-1} (P - FA) \]
where \( \phi^{-1} \) is the inverse of the covariance matrix in the measured projections \( P \) and is diagonal since the measured projections are assumed independent. \( \tilde{R}(A) \) is minimized when
\[ F^T \phi^{-1} FA = F^T \phi^{-1} P, \quad (38) \]
or
\[ A = (F^T \phi^{-1} F)^{-1} F^T \phi^{-1} P \quad (39) \]

Another unbiased estimator is obtained by substituting unity for the variance of the measured values \( P \), namely
\[ A = (F^TF)^{-1} F^T P. \quad (40) \]

With good statistics, the results from these two estimators will be close.

For a given experimental situation (i.e., a fixed number of angles and fixed field of view) the matrix \( F \) is fixed, independent of the distribution of radionuclide under study. The vector \( F^T P \) (usually called the back projection) is the convolution of the source by an appropriate function (\( 1/r \) for an infinity of rotation angles). \( F^TF \) represents this convolution.

Then, given the back projection of a certain distribution of radionuclide, one can envision operating on it with the precalculated matrix \( (F^TF)^{-1} \) to obtain the reconstruction. In fact, \( F^TF \) has very large dimensions, and calculating its inverse on the largest computers does not seem feasible at this time. (For a 64 x 64 array of pixels, \( F^TF \) has dimensions of 4096 x 4096). We intend to investigate the possibility of calculating an approximation to the matrix \( (F^TF)^{-1} \).
For continuous rotation angles $F^TF$ falls off as $1/r$, and it is expected that the inverse $(F^TF)^{-1}$ also falls off as the distance between pixels increases. Therefore, we intend to explore the use of an approximation to $(F^TF)^{-1}$ which is also a convolution, but whose elements are zero for pixels separated by some minimum distance. This procedure has been tried on various phantoms with results which compare favorably with the iterative least squares technique.

**Marr's Positron Technique.** Another technique we are considering was proposed by Marr (1974). For N detectors, the number of coincidence lines of position is $N(N-1)/2$. This value includes near neighbor detector combinations, which in practice would not be used. These projection sample points are represented by

$$\{g(I,J) \mid 1 \leq I \leq N, 1 \leq J \leq N - 1\}$$

where I and J refer to detectors. Note that $g(I,I)$ cannot be a data sample point since a detector cannot record a coincidence with itself. The reconstructed image $\rho(x,y)$ for this data set is given by the following equations

$$\rho(x,y) = \sum_{n=0}^{M} \sum_{k=0}^{[(M-n)/2]} \left( \alpha_{n,k} \sin n\psi + \beta_{n,k} \cos n\psi \right) r^n Q_{n,k}(r^2) \quad (41)$$

where

$$\beta_{0,k} = \frac{2k+1}{N^2} \sum_{I,J} \sin \left( 2k+1 \right) \frac{\pi J}{N} g(I,J) \quad \ldots \quad 0 \leq k \leq \frac{N-2}{2}, \quad (42)$$

$$\left\{ \alpha_{n,k} \right\} = \frac{2(n+2,k+1)}{N^2} \sum_{I,J} \sin \left( (n+2) \frac{\pi J}{N} \right) \times \left\{ \sin \right\} \frac{\pi J}{N} \left( 2I+J-2 \right) g(I,J) \quad (43)$$

$$\ldots \quad 1 \leq n \leq N - 2, \quad 0 \leq k \leq \left[ \frac{N-n-2}{2} \right],$$
\[ Q_{n,k}(t) = \frac{1}{k!} \frac{d}{dk} \left\{ t^{n+k} (t-1)^k \right\}, \] (44)

and where \( M \) is some specified value such that \( M \leq N - 2 \). The Radon transform of \( \rho(x,y) \) fits the data in a least squares sense.
7. RUBIDIUM GENERATOR AND DOSE

Two $^{82}$Rb generators have been fabricated with about 1 mCi of $^{82}$Sr produced at the Lawrence Berkeley Laboratory 88-inch cyclotron using the reaction $^{85}$Rb(p,4n)$^{82}$Sr (Yano and Anger, 1968). A $^{82}$Rb generator with 6 mCi capacity can be prepared using the same principles of eluting $^{82}$Rb from a weakly acidic cation exchange resin. The eluant is 10-30 cc of 2-3% NaCl. Experience has shown that we have only $10^{-5}$ contamination from $^{82}$Sr.

The total body dose is approximately 10 mrads for 6 mCi $^{82}$Rb. This dose is in good part due to the energetic positrons. The positron dose (in rads) is calculated from:

$$\text{Dose } \beta^+ = 73.8 \ C_0 \ \text{T}_{1/2} \ \int_0^{E_{\text{max}}} \ N(E) \ E \ dE$$  \hspace{1cm} (45)

where $C_0$ is the concentration in tissue ($\mu$Ci/gm) immediately after injection; assuming an equal distribution, i.e., 6000 microcuries, 70,000 gms. $T_{1/2}$ is the half life in days, and $N(E)$ is the density distribution for positrons of energy $E$. For $^{82}$Rb this integral is 1.4 MeV. The half life in days is $8.68 \times 10^{-4}$. Thus the positron dose is 8 millirads.

The gamma ray dose is calculated from the absorbed fraction method introduced by G. Brownell and gives only one fourth the dose given from the positrons. Thus the total dose is 10 millirads for a 6 millicurie injection.
**TABLE 1.**

**CARDIAC IMAGING**

Photoscanning by rectilinear scanner or imaging with the gamma camera.

<table>
<thead>
<tr>
<th>Isotope</th>
<th>Limitations</th>
<th>Investigators</th>
</tr>
</thead>
<tbody>
<tr>
<td>$^{131}$Cs</td>
<td>Low energy (29.4)Xe x-rays absorbed by photoelectric process in ribs and sternum. Long half-life.</td>
<td>Carr et al., 1964</td>
</tr>
<tr>
<td>$^{129}$Cs</td>
<td>High whole-body dose per amount injected and limitation in resolution due to 375 keV photons. Expense of isotope. High dose, high energy and high cost.</td>
<td>Bing, 1967</td>
</tr>
<tr>
<td>or $^{43}$K</td>
<td></td>
<td>Collier et al., 1968</td>
</tr>
<tr>
<td>$^{134m}$Cs</td>
<td>Short half-life relative to optimum scanning time and availability.</td>
<td>Yano et al., 1970</td>
</tr>
<tr>
<td>$^{43}$K</td>
<td>High energy photons (620, 373) and high cost. Rapid heart clearance.</td>
<td>Romhilt et al., 1973</td>
</tr>
<tr>
<td>$^{13}$NH$_3$ (gamma)</td>
<td>High energy gammas (511 keV) from positron annihilation leads to low resolution because of difficulties in collimation. Need for cyclotron nearby.</td>
<td>Smith et al., 1970</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Bennett et al., 1972</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Chandra et al., 1973</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Holman et al., 1973a</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Hurley et al., 1971</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Zaret et al., 1973</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Strauss et al., 1973</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ishii et al., 1973</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Smith et al., 1972</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Botti et al., 1973</td>
</tr>
<tr>
<td></td>
<td></td>
<td>MacIntyre et al., 1972</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Poe, 1972</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Harper et al., 1972</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Monohan et al., 1972</td>
</tr>
<tr>
<td>Isotope</td>
<td>Limitations</td>
<td>Investigators</td>
</tr>
<tr>
<td>---------------------</td>
<td>-----------------------------------------------------------------------------</td>
<td>----------------------------------------</td>
</tr>
<tr>
<td>$^{13}$NH$_3$ (positron)</td>
<td>Need for positron camera and hospital cyclotron.</td>
<td>Hoop et al., 1973</td>
</tr>
<tr>
<td>$^{81}$Rb</td>
<td>High energy (450-520) leads to low resolution due to difficulty in collimation.</td>
<td>Budinger et al., 1973</td>
</tr>
<tr>
<td>$^{82}$Rb (positron)</td>
<td>Requires positron camera of high count rate capacity for studies in man.</td>
<td>Martin et al., 1974</td>
</tr>
<tr>
<td>Thallium</td>
<td>Relative to other isotopes limitation is long half-life, 75 hrs., obviating repeated studies. Compares well in other respects.</td>
<td>Yano and Anger, 1968</td>
</tr>
<tr>
<td>$^{131}$I fatty acids</td>
<td>Availability and statistics. Low statistics per patient dose.</td>
<td>Bonte et al., 1973</td>
</tr>
<tr>
<td>$^{99m}$Tc-tetracycline</td>
<td>Unavailability of radiopharmaceutical with sufficient activity. Accumulates in fresh infarcts only 3-7 days.</td>
<td>Holman et al., 1973b</td>
</tr>
<tr>
<td>$^{99m}$Tc-pyrophosphate</td>
<td></td>
<td>Cook et al., 1974</td>
</tr>
<tr>
<td>Coronary Injection</td>
<td>Invasive, see $^{43}$K above.</td>
<td>Holman et al., 1973a</td>
</tr>
<tr>
<td>$^{43}$K</td>
<td>Invasive.</td>
<td>Ashburn et al., 1971</td>
</tr>
<tr>
<td>Microspheres</td>
<td>Invasive.</td>
<td>Weller et al., 1972</td>
</tr>
<tr>
<td>$^{131}$I, $^{99m}$Tc</td>
<td>Invasive.</td>
<td>Jansen et al., 1973</td>
</tr>
<tr>
<td>Xe</td>
<td>Invasive.</td>
<td>Cannon et al., 1973</td>
</tr>
</tbody>
</table>

Table 1. Cardiac Imaging (continued)
Table 1. Cardiac Imaging (continued)

<table>
<thead>
<tr>
<th>Isotope</th>
<th>Limitations</th>
<th>Investigators</th>
</tr>
</thead>
<tbody>
<tr>
<td>Precordial Counting</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Use of NaI(T) probe or positron coincidence systems.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>$^{42}$K, $^{86}$Rb</td>
<td>No or limited spatial resolution capabilities. Variable and unknown background contribution.</td>
<td>Love et al., 1954 Love et al., 1957 Donato et al., 1964</td>
</tr>
<tr>
<td>$^{84}$Rb (positron)</td>
<td>No or limited spatial resolution.</td>
<td>Levi and DeOliveira, 1961 Bing et al., 1964 Cohen et al., 1965 Luebs et al., 1966 Cohen et al., 1967 McHenry et al., 1967 Teb et al., 1969</td>
</tr>
</tbody>
</table>
TABLE 2.
PRELIMINARY SPECIFICATIONS

1. Physical

<table>
<thead>
<tr>
<th>Specification</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>External Ring Diameter</td>
<td>170 cm</td>
</tr>
<tr>
<td>Crystal Ring Diameter</td>
<td>80 cm</td>
</tr>
<tr>
<td>Minimum Lead Collimator Diameter</td>
<td>40 cm</td>
</tr>
<tr>
<td>Crystal Size</td>
<td>0.8 cm x 3 cm x 5 cm</td>
</tr>
<tr>
<td>Total Number of Crystals</td>
<td>288</td>
</tr>
<tr>
<td>Sections</td>
<td>8</td>
</tr>
<tr>
<td>Crystals per Section</td>
<td>36</td>
</tr>
<tr>
<td>Photomultiplier Tubes per Section</td>
<td>9</td>
</tr>
<tr>
<td>Total Number of PMTs</td>
<td>72</td>
</tr>
<tr>
<td>Crystals in Coincidence with Opposite</td>
<td>62</td>
</tr>
<tr>
<td>Effective Diameter of Full Sensitivity</td>
<td>~ 30 cm</td>
</tr>
<tr>
<td>Weight of Ring Assembly and Shield</td>
<td>600 kg</td>
</tr>
</tbody>
</table>

2. Performance:

<table>
<thead>
<tr>
<th>Specification</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Resolution at Center</td>
<td>6 mm FWHM</td>
</tr>
<tr>
<td>Slice Thickness at Center</td>
<td>5 to 15 mm FWHM</td>
</tr>
<tr>
<td>Pulse Pair Resolving Time</td>
<td>$10^{-8}$ sec</td>
</tr>
</tbody>
</table>
### TABLE 3.
PROPERTIES OF A RING DETECTOR SYSTEM\(^a\)
FOR THREE ASSUMED ENERGY RESOLUTIONS

<table>
<thead>
<tr>
<th>Quantity</th>
<th>Symbol</th>
<th>Units</th>
<th>Case I</th>
<th>Case II</th>
<th>Case III</th>
</tr>
</thead>
<tbody>
<tr>
<td>Energy resolution (FWHM) at 511 keV</td>
<td>( \Gamma )</td>
<td>%</td>
<td>none</td>
<td>30</td>
<td>15</td>
</tr>
<tr>
<td>Energy threshold</td>
<td>( E_p )</td>
<td>keV</td>
<td>100</td>
<td>410(^b,c)</td>
<td>460(^b,c)</td>
</tr>
<tr>
<td>Approximate maximum scattering angle corresponding to ( E_p ) (Eq. 13)</td>
<td>( \theta_p )</td>
<td>degrees</td>
<td>180</td>
<td>41</td>
<td>27</td>
</tr>
<tr>
<td>Effective maximum scattering angle</td>
<td>( \theta_p )</td>
<td>degrees</td>
<td>40</td>
<td>40</td>
<td>27</td>
</tr>
<tr>
<td>Minimum scattered gamma energy corresponding to ( \theta_m ) (Eq. 14)</td>
<td>( E_m )</td>
<td>keV</td>
<td>415</td>
<td>415</td>
<td>460</td>
</tr>
<tr>
<td>Average detection efficiency for gamma ray energies from ( E_m ) to 511 keV</td>
<td>( \epsilon_m )</td>
<td>%</td>
<td>65</td>
<td>25</td>
<td>25</td>
</tr>
<tr>
<td>Detection efficiency for 511 keV gammas</td>
<td>( \epsilon )</td>
<td>%</td>
<td>50</td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td>Probability that a 511 keV gamma ray will scatter and retain an energy above ( E_p ) (Eq. 24)</td>
<td>( P_p )</td>
<td>%</td>
<td>63</td>
<td>20</td>
<td>10</td>
</tr>
<tr>
<td>Average detection efficiency for gamma ray energies from ( E_p ) to 511 keV</td>
<td>( \epsilon_p )</td>
<td>%</td>
<td>80</td>
<td>25</td>
<td>25</td>
</tr>
<tr>
<td>Geometry factor - one gamma scatters (Eq. 16)</td>
<td>( g_1 )</td>
<td>--</td>
<td>0.31</td>
<td>0.31</td>
<td>0.23</td>
</tr>
<tr>
<td>Geometry factor - both gammas scatter (Eq. 19)</td>
<td>( g_2 )</td>
<td>--</td>
<td>0.19</td>
<td>0.19</td>
<td>0.13</td>
</tr>
<tr>
<td>Rate for unscattered coincident events (Eq. 5)</td>
<td>( C_0 )</td>
<td>events per sec</td>
<td>6260</td>
<td>1000</td>
<td>1000</td>
</tr>
<tr>
<td>Sensitivity</td>
<td>( C_0/\rho )</td>
<td>events per sec per [( \mu )Ci/cm]</td>
<td>31</td>
<td>5</td>
<td>5</td>
</tr>
</tbody>
</table>
Table 3. Properties of a ring detector system for three assumed energy resolutions (continued)

<table>
<thead>
<tr>
<th>Quantity</th>
<th>Symbol</th>
<th>Units</th>
<th>Case I</th>
<th>Case II</th>
<th>Case III</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coincidence rate for one gamma scatter (Eq. 15)</td>
<td>$C_1$</td>
<td>events per sec</td>
<td>1735</td>
<td>267</td>
<td>198</td>
</tr>
<tr>
<td>Coincidence rate for two gamma scatter (Eq. 18)</td>
<td>$C_2$</td>
<td>events per sec</td>
<td>1190</td>
<td>176</td>
<td>120</td>
</tr>
<tr>
<td>Singles rate for entire ring (Eq. 23)</td>
<td>$C_s$</td>
<td>$10^5$ events per sec</td>
<td>10.2</td>
<td>1.83</td>
<td>1.46</td>
</tr>
<tr>
<td>Accidental coincidence rate for entire ring (Eq. 29)</td>
<td>$C_a$</td>
<td>events per sec</td>
<td>2290</td>
<td>74</td>
<td>47</td>
</tr>
<tr>
<td>Total event rate ($C_0 + C_1 + C_2 + C_a$)</td>
<td></td>
<td>events per sec</td>
<td>11475</td>
<td>1517</td>
<td>1365</td>
</tr>
<tr>
<td>Background fraction</td>
<td></td>
<td>%</td>
<td>45</td>
<td>34</td>
<td>27</td>
</tr>
<tr>
<td>Paralyzing deadtime</td>
<td>$t_p$</td>
<td>nsec</td>
<td>200</td>
<td>800</td>
<td>800</td>
</tr>
<tr>
<td>Fraction of events lost due to deadtime (Eq. 32)</td>
<td>$f_e$</td>
<td>%</td>
<td>5</td>
<td>23</td>
<td>23</td>
</tr>
</tbody>
</table>

a) Physical specifications of assumed ring detector system are as follows:

- $c$ = detector ring radius = 40 cm
- $S$ = shielding slit width = 2 cm
- $T$ = shielding slit depth = 20 cm
- $P_s$ = probability of scattering on entering from the center of a 20 cm diam cylinder of water = 63%
- $f = 0.22$ (for 288 crystals, each would be in coincidence with the opposing 62 crystals).
- $\theta_f = 40^\circ$ maximum scattering angle imposed by coincidence requirement (approximate)
Table 3. Properties of a ring detector system for three assumed energy resolutions (continued)

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\rho$ = activity density</td>
<td>200 $\mu$Ci per axial cm</td>
</tr>
<tr>
<td>$t$ = time resolution</td>
<td>20 nsec</td>
</tr>
<tr>
<td>Dimension of crystals along gamma line of flight</td>
<td>5 cm</td>
</tr>
<tr>
<td>Dimension of crystals along ring axis</td>
<td>3 cm</td>
</tr>
<tr>
<td>Dimension of crystals transverse to ring axis</td>
<td>0.8 cm</td>
</tr>
<tr>
<td>$R$ = radius of uniform geometrical sensitivity</td>
<td>14 cm</td>
</tr>
<tr>
<td>$G$ = gross singles rate per segment</td>
<td>$1.3 \times 10^5$ counts per sec</td>
</tr>
</tbody>
</table>

b Corresponds to a 7% photopeak loss, assuming that the photopeak has a Gaussian distribution.

c This event loss has not been included elsewhere in this table.
TABLE 4.

THE GEOMETRICAL FACTORS $g_1$ AND $g_2$

AS A FUNCTION OF THE MAXIMUM EFFECTIVE SCATTERING ANGLE $\theta_m$

<table>
<thead>
<tr>
<th>$\theta_m$</th>
<th>$g_1$</th>
<th>$g_2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>0°</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>10°</td>
<td>0.10</td>
<td>0.05</td>
</tr>
<tr>
<td>20°</td>
<td>0.18</td>
<td>0.10</td>
</tr>
<tr>
<td>30°</td>
<td>0.26</td>
<td>0.15</td>
</tr>
<tr>
<td>40°</td>
<td>0.31</td>
<td>0.19</td>
</tr>
<tr>
<td>50°</td>
<td>0.36</td>
<td>0.23</td>
</tr>
<tr>
<td>60°</td>
<td>0.40</td>
<td>0.26</td>
</tr>
</tbody>
</table>
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FIGURE CAPTIONS

1. Transverse section tomography of the human myocardium, using $^{129}$Cs and multiple views taken by a Scintillation Camera with parallel channel collimation.

2. Transverse section tomography of the myocardium of the dog, as per Fig. 1.

3. Gated single view of the myocardium using $^{81}$Rb and a Scintillation Camera with pinhole.

4. Comparison of $^{82}$Rb to $^{13}$N-ammonia myocardial imaging in a dog.

5. NaI(Tl) crystal in 0.25 mm thick stainless steel package manufactured by Harshaw Chemicals. Dimensions are 9.5 mm x 9.5 mm x 38 mm.

6. Schematic showing patient positioned inside the detector ring.

7. Example of assembly of crystals (inside lead collimator) connected to light pipes and photomultiplier tubes (PMTs). Signals from PMTs are checked for validity of code (to insure that detection occurred in one crystal only) before energy selection.

8. Example showing coding scheme for 6 crystals and 4 photomultipliers. Expansion to 36 crystals requires 5 additional photomultipliers.

9. Schematic of coincidence coding for 6 segment example. Each crystal is required to be in time coincidence with a pre-selected number of opposing crystals.
10. Schematic showing detector circle, collimator and geometrical quantities used in Section 5.2

11. Axial response function - calculated coincident response for a fixed angle $\phi$ (see Fig. 10) as an off-axis positron source is moved parallel to the Z axis.

12. Axial response function (averaged over all gamma ray directions) for various distances to the center of the detector circle.

13. Schematic showing that for off-axis points the transaxial resolution depends on the gamma ray direction.

14. Schematic showing a portion of the detector ring and the geometrical variables used in Section 5.6.

15. Transaxial point spread function (averaged over all gamma ray directions) for various distances to the center of the detector ring.

16. FWHM of transaxial point spread function (averaged over all gamma ray directions) vs. the distance to the center of the detector circle.

17. Axial response functions (on axis sources) for co-linear coincident gamma ray pairs and for single gamma rays (or non-co-linear coincident pairs).

18. Illustration showing how Compton Scattering can contribute background to the coincident event rate. Although two photons are in time coincidence, they are not colinear and provide erroneous spatial information. Unscattered coincident events can arise only from region between the dashed lines.
19. Energy lost by a gamma ray during Compton scattering as a function of the gamma ray scattering angle.

20. Cross section per unit solid angle for the Compton scattering of 500 keV gamma rays as a function of the gamma ray scattering angle.

21. Comparison of four techniques for three-dimensional reconstruction — back projection, simultaneous iterative reconstruction, least-squares, and back projection of filtered projections (BPFP) — using lucite phantoms and $^{99m}$Tc.

22. Least-squares technique reconstruction of a phantom consisting of the letters "L" "0" "V" and the discrepancy between the true image and the reconstruction.
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