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Develop Real-Time Dosimetry Concepts and Instrumentation for Long-Term Missions

Technical Progress
February 1981 to February 1982
L. A. Braby

May 1982

Prepared for
National Aeronautics and Space Administration
Lyndon B. Johnson Space Center
under a Related Services Agreement
with the U.S. Department of Energy

Pacific Northwest Laboratory
Operated for the U.S. Department of Energy
by Battelle Memorial Institute

Battelle
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Pacific Northwest Laboratory
Richland, Washington 99352
**CONTENTS**

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>FIGURES</td>
<td>iii</td>
</tr>
<tr>
<td>TABLES</td>
<td>iii</td>
</tr>
<tr>
<td>INTRODUCTION</td>
<td>1</td>
</tr>
<tr>
<td>DETECTOR DEVELOPMENT</td>
<td>1</td>
</tr>
<tr>
<td>ELECTRONIC SYSTEM</td>
<td>2</td>
</tr>
<tr>
<td>EVALUATION OF RADIATION QUALITY</td>
<td>6</td>
</tr>
<tr>
<td>FUTURE DIRECTION</td>
<td>15</td>
</tr>
<tr>
<td>REFERENCES</td>
<td>16</td>
</tr>
</tbody>
</table>
FIGURES

1. Detector Gas Gain at a Constant Anode Voltage as a Function of Time After Sealing the Detector ...................................... 2
2. Gas Gain Versus Anode Voltage ............................................. 3
3. Flow Diagram of the Subroutine and Block Diagram of Hardware for Adjusting the Anode Voltage to Maintain Constant Gas Gain .................................................. 4
4. Block Diagram of a System Using Two Detectors to Cover a Wide Range of Event Sizes .................................................. 5
5. Detailed Block Diagram of the Electronics for the High Gain Detector ................................................................. 6
6. Analog to Digital Converter Using CMOS Successive Approximation Circuit ................................................................. 7
7. Prototype Portable Instrument with the High Gain Detector ................................................................. 8
8. Typical Curves for the Density of Dose in Lineal Energy for Neutron and Mixed Field Irradiations ............................................. 11
9. Mean and Standard Deviation for Forty Repetitions of $\bar{Q} = 0.8 + 0.14 \bar{Y}_D$ at Each Value of the Dose .................................................. 12
10. Mean and Standard Deviation for 40 Samples of $\bar{Q}$ Determined by a Fourier Transform Deconvolution of the LET Distribution at Each Dose ............................................. 13

TABLES

1. Power Consumption ............................................................. 8
2. Mean Value for 40 Calculations of Quality ............................................. 13
4. Milestones ................................................................. 15
INTRODUCTION

Major objectives in the process of developing a rugged portable instrument to evaluate dose and dose equivalent have been achieved. A tissue-equivalent proportional counter simulating a 2 micrometer spherical tissue volume has operated satisfactorily for over a year. The basic elements of the electronic system have been designed and tested. And finally, the most suitable mathematical technique for evaluating dose equivalent with a portable instrument has been selected. Design and fabrication of a portable prototype, based on the previously tested circuits, is underway.

DETECTOR DEVELOPMENT

The 5.7 cm detector has been operated for over 15 months with the original gas filling. Figure 1 shows the gas gain for a constant anode voltage during that time. The long-term drift, less than one percent per month, and short-term (less than 24 hour) variations of ±2 percent can easily be controlled by adjusting the anode voltage, (see Figure 2). These adjustments will be made automatically by a subroutine, illustrated in Figure 3, which compares the actual position of a calibration peak with the position of that peak at the proper gas gain. The difference is used to calculate a new high-voltage supply setting. The actual voltage is provided by a high-voltage supply referenced to the output of a digital-to-analog converter.

The detector will be operated with a combination of gas gain and electronic gain which results in a calibration factor of 0.12 keV/μm/channel and a useful range of 0.36 to 300 keV/μm. Since the dose mean of the single-event distribution for X and γ rays varies from 0.7 to 3.0 keV/μm, and for neutrons from 30 to 130 keV/μm, the system is expected to detect photon as well as neutron dose. However, the stainless steel vacuum chamber surrounding the detector will distort the response as a function of photon energy. Also, the photon-induced events cannot be distinguished from very low-energy neutron events.
Thus, though the system is expected to accurately measure dose due to photons and low-energy neutrons, estimates of the mean quality factor become less certain when low-energy neutrons are abundant.

Specifications for the second detector, intended to detect high-energy heavy particles, have been completed. It will use the same basic design and materials but will be 12.7 cm in diameter in order to provide approximately five times the counting rate.

ELECTRONIC SYSTEM

The electronics for a system using two detectors is outlined in Figure 4. Each detector is supported by a dedicated high-voltage supply, amplifiers and analog-to-digital converters (ADC's), but the two detectors share a multichannel analyzer (MCA) and microcomputer. The ADC's output is used as a MCA memory address. Each event results in a "one" being added to the memory content at
the corresponding address. As illustrated in Figure 5, the entire MCA content can be periodically transferred to the microcomputer memory. The dosimetric quantities can then be computed without interrupting further data collection. An absolute time clock in the microcomputer is used to initiate the dose calculation, data storage, calibration and other functions. A CMOS successive approximation ADC chip and other CMOS components are used in the ADC circuit, in Figure 6, to minimize power consumption.
FIGURE 3. Flow Diagram of the Subroutine and Block Diagram of Hardware for Adjusting the Anode Voltage to Maintain Constant Gas Gain

Preliminary versions of all of this circuitry have been assembled and are being tested in the laboratory prototype described previously (PNL-3747, Braby 1981). Power consumption, an important feature in a portable system, is listed in Table 1 for the current versions of the main components. These designs have established the physical and electronic requirements for a prototype portable instrument. Mechanical work on this portable prototype (see Figure 7) has been
completed. As individual circuits are refined and power consumption is reduced, cards will be assembled for use in this prototype. For convenience in making calibration measurements, it has been equipped with operator controls such as acquire, reset, and record. These functions will be controlled by the micro-computer when the system is complete. During testing, the operating program will be in random access memory and can be easily altered and re-entered via an RS232 port. For routine operation, the same program will be installed in a read-only memory to avoid having to re-enter the program after a power loss.
There are two basic ways of determining the quality factor based on the measured probability density of energy imparted, \( f(\epsilon) \), or related functions such as \( f(y) \), the density of \( y \) where \( y \) is related to the mean cord length \( T \) by \( y = \epsilon / T \), and the density of dose in \( y \), \( d(y) = yF(y) \). The first method involves direct application of the lineal energy, usually in the form of \( d(y) \), and the second method involves the deconvolution of the LET distribution from
FIGURE 6. Analog to Digital Converter Using CMOS
Successive Approximation Circuit
### TABLE 1. Power Consumption

<table>
<thead>
<tr>
<th>Circuit</th>
<th>Power Watts</th>
<th>Number Required in Two-Detector System</th>
<th>Total Watts</th>
</tr>
</thead>
<tbody>
<tr>
<td>preamp</td>
<td>0.2</td>
<td>2</td>
<td>0.4</td>
</tr>
<tr>
<td>amplifier</td>
<td>1.1</td>
<td>2</td>
<td>2.2</td>
</tr>
<tr>
<td>H.V. supply</td>
<td>1.6</td>
<td>2</td>
<td>3.2</td>
</tr>
<tr>
<td>A.D.C.</td>
<td>0.75</td>
<td>3</td>
<td>2.25</td>
</tr>
<tr>
<td>MCA</td>
<td>0.05</td>
<td>1</td>
<td>0.05</td>
</tr>
<tr>
<td>microcomputer</td>
<td>0.35</td>
<td>1</td>
<td>0.35</td>
</tr>
<tr>
<td>memory</td>
<td>~0.2</td>
<td>1</td>
<td>0.2</td>
</tr>
<tr>
<td>display</td>
<td>~0.1</td>
<td>1</td>
<td>0.1</td>
</tr>
<tr>
<td>tape recorder</td>
<td>1</td>
<td>~5% duty cycle</td>
<td>0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>8.80</td>
</tr>
</tbody>
</table>
the $y$ distribution (see for example Rossi, 1968). If one assumes that energy-loss straggling and delta-ray effects are insignificant and that all of the tracks are long compared to the site diameter so that there is no stopping or starting within the detector volume (Kellerer 1969) and if one further assumes a functional form for the relationship between LET and quality factor, one can derive

$$Q = 0.8 + 0.14 \overline{y}_D$$  \hspace{1cm} (1)

where $\overline{y}_D$ is the dose mean of the lineal energy.

A second way of applying $y$ directly,

$$Q = \int_0^\infty y^{1.5} d(y)dy,$$  \hspace{1cm} (2)

was proposed by Rossi (1977) as a new definition for quality factor. The quantity $y^*$ is a saturated version of the lineal energy intended to take into account the fact that RBE generally decreases for LET values above about 150 keV/µ. This formulation of Equation 2 was intended to provide very large values of the radiation quality, in line with biophysics models which suggest that the quality factor should be 10 to 20 times higher than the current definition. However, this same method can be applied to give quality factors in line with current definitions by substituting 0.6 for the exponent 1.5 in Equation 2.

The simplest of the methods for unfolding LET distribution from the single-event distribution measured in the spherical detector is

$$d(L_{\infty}) = \frac{L_{\infty}d}{z} \left( f(\varepsilon) - \varepsilon \frac{df(\varepsilon)}{d\varepsilon} \right)$$  \hspace{1cm} (3)

where $d(L_{\infty})$ is the density of absorbed dose in LET, \(d\) is the site diameter, and $f(\varepsilon)$ is the density of energy imparted. This method is based on the fact

\[ \text{(a) In ICRU 19 and 33 the symbol } D_{L_{\infty}} \text{ is used for } d(L_{\infty}). \]
that a spherical detector produces a triangular cord-length distribution and
the contribution of any given LET to the overall distribution can be found
from the slope of the distribution for energy imparted corresponding to that
value of LET. Once the dose distribution in LET has been derived, the defini-
tion for the mean quality factor given in ICRU 19 and 33 is;

\[
\bar{Q} = \int_0^\infty Q \, d(L_\infty) \, dL_\infty \int_0^\infty d(L_\infty) \, dL_\infty
\]  

and can be utilized to get the mean quality factor.

Another method for unfolding the dose distribution of LET was proposed
by Kellerer (1972). This involves generating the Fourier transform of both
the distribution of energy imparted and the track-length distribution. The
quotient of these two is the transform of the LET distribution. The reverse
transform provides the desired distribution, \( d(L_\infty) \). While this process sounds
complicated, algorithms for the fast Fourier transform are available even for
small microprocessor-based computers and this unfolding technique can easily
be applied in a microcomputer with 8K bytes of memory.

Each of these methods was tested using simulated spectra corresponding to
specified doses. A smooth distribution for a specific neutron energy, derived
from the literature, was entered as data. Random number generators were then
used to generate a simulated measurement with the appropriate statistical vari-
ation for the specified dose. The algorithm being tested was used to calculate
the quality factor for that simulated distribution and the result was stored.
Then a new distribution for the same dose and initial distribution was gener-
ated and another value of the quality factor was calculated. This procedure
was repeated 40 times for each dose. Finally a mean and standard deviation of
the quality factor for those 40 simulated measurements was calculated.

All of the methods of calculating quality involved the dose distribution
either of \( y \) or LET. That is, they involved the frequency of events multiplied
by the event size. This adds substantial weight to high values of \( y \) or LET so
that, even though there are relatively few events at the high values of \( y \), these
values have a large effect on the estimate of the quality. Figure 8 gives several microdosimeter distributions for neutrons and mixed fields in terms of \( yd(y) \) plotted versus the log of \( y \) so that equal areas under the curve represent equal doses. These smoothed curves were used as the starting point for calculations of the distributions to be expected from experimental measurements at various dose levels.

Figure 9 shows the mean and standard deviation of \( \bar{Q} \) determined by the method in Equation 1 (Equation 2 results in similar lines but with smaller standard deviations). The value of \( \bar{Q} \) is constant for each radiation as a function of dose, but the standard deviation for the estimate of \( \bar{Q} \) increases rapidly from less than a percent at \( 10^{-2} \) rads to over 9 percent at \( 10^{-5} \) rads for the

![Typical Curves for the Density of Dose in Lineal Energy for Neutron and Mixed Field Irradiations](image)

**FIGURE 8.** Typical Curves for the Density of Dose in Lineal Energy for Neutron and Mixed Field Irradiations
half MeV neutrons. Figure 10 is the equivalent result using the Fourier transform method. This has the unfortunate nature that as the dose decreases, the mean of the 40 samples of $\bar{Q}$ decreases abruptly, and the standard deviation increases rapidly. In fact it is not possible to get an estimate of $\bar{Q}$ at $10^{-5}$ rads for the 15 MeV neutrons or even at $10^{-4}$ rads with a mixed-neutron gamma ray field. This is due to high-frequency components in the transform resulting from the noise in $F(e)$. These high-frequency components lead to negative values for the density of LET.

Table 2 gives the mean value for 40 calculations of quality as defined by Equations 1 through 4 for each of the $y$ distributions illustrated in Figure 8 at the dose of $10^{-2}$ rads, and also gives the ICRP report 21 maximum

FIGURE 9. Mean and Standard Deviation for 40 Repetitions of $Q = 0.8 + 0.14 \bar{y}_D$ at Each Value of the Dose
FIGURE 10. Mean and Standard Deviation for 40 Samples of $\bar{\Phi}$ Determined by a Fourier Transform Deconvolution of the LET Distribution at Each Dose.

TABLE 2. Mean Value for 40 Calculations of Quality

<table>
<thead>
<tr>
<th>Equations</th>
<th>Mixed Field</th>
<th>0.55 MeV</th>
<th>2.2 MeV</th>
<th>15 MeV</th>
</tr>
</thead>
<tbody>
<tr>
<td>1) $\bar{a}$</td>
<td>2.6</td>
<td>11.7</td>
<td>12.5</td>
<td>6.4</td>
</tr>
<tr>
<td>2) $\bar{a}_y^{1.5}$</td>
<td>69.3</td>
<td>476</td>
<td>461</td>
<td>184</td>
</tr>
<tr>
<td>$\gamma^{0.6}$</td>
<td>2.9</td>
<td>10.9</td>
<td>11.2</td>
<td>6.6</td>
</tr>
<tr>
<td>3) $\bar{c}$</td>
<td>4.7</td>
<td>13.6</td>
<td>16.4</td>
<td>9.7</td>
</tr>
<tr>
<td>numerical filtering</td>
<td>3.7</td>
<td>14.8</td>
<td>16.4</td>
<td>8.0</td>
</tr>
<tr>
<td>4) $\bar{c}_e$</td>
<td>2.6</td>
<td>11.3</td>
<td>11.2</td>
<td>5.4</td>
</tr>
<tr>
<td>ICRP21(a)</td>
<td></td>
<td>11</td>
<td>9.2</td>
<td>6.4</td>
</tr>
</tbody>
</table>

(a) ICRP report maximum quality for monoenergetic neutron sources.
quality for the monoenergetic neutron sources. Since there is no direct comparison between the spectra used in these calculations and the ICRP report calculations, this cannot be taken as a good indication of the accuracy of these calculational methods, but does give a rough idea of the accuracy which might be obtained. Clearly even for 15 MeV neutrons, which should show substantial straggling, the method outlined in Equation 1 produces a surprisingly satisfactory value for the quality factor. Equation 2, using the original exponent of 1.5 is intended to produce very large values of quality and in fact does. However, substituting the exponent 0.6 this method provides equally good, if not better values of the quality than Equation 1 did. The method involving Equation 3 for the deconvolution of LET from \( y \) distribution is somewhat less successful. It tends to overestimate the quality in almost all situations. Some numerical filtering to smooth the differentiation process reduces the overestimations somewhat but does not eliminate the problem. The Fourier transform method produces values of the quality which are consistent with the methods based directly on \( y \) distributions but is less satisfactory in that it loses precision much more rapidly. Table 3 summarizes the relative standard deviation of the value of \( \bar{Q} \) at a dose of \( 10^{-4} \) rads for the four equations, giving conventional values of \( \bar{Q} \). This shows that the method of Equation 2 produces the least variation between measurements at low doses.

<table>
<thead>
<tr>
<th>Equations</th>
<th>Mixed Field</th>
<th>0.55 MeV</th>
<th>2.2 MeV</th>
<th>15 MeV</th>
</tr>
</thead>
<tbody>
<tr>
<td>1) ( Q_a )</td>
<td>11.2</td>
<td>2.9</td>
<td>4.4</td>
<td>9.7</td>
</tr>
<tr>
<td>2) ( Q_b(y^{0.6}) )</td>
<td>6.9</td>
<td>1.1</td>
<td>1.3</td>
<td>3.8</td>
</tr>
<tr>
<td>3) ( Q_c )</td>
<td>20.3</td>
<td>3.0</td>
<td>1.8</td>
<td>10.8</td>
</tr>
<tr>
<td>4) ( Q_e )</td>
<td>--</td>
<td>1.5</td>
<td>1.4</td>
<td>12.9</td>
</tr>
</tbody>
</table>

Based on these calculations for monoenergetic neutrons, it is evident that the method of Equation 2 (with the exponent set at 0.6) has several advantages over the other methods of calculating \( Q \). It is relatively simple compared to unfolding \( d(L) \), it produces mean values in agreement with the other methods and ICRP recommendations, the mean value does not decrease with dose, and the
estimates show the least variation at low doses. In addition, this method is least likely to overestimate the quality factor for very high LET particles. The method of Equation 1 would be a good second choice, its accuracy is good and it does not change significantly with dose, but the precision is significantly poorer.

FUTURE DIRECTION

All of the major components needed for the portable instrument have now been demonstrated. The portable prototype will be assembled as refinements in individual circuits are completed. This instrument will utilize the method of Equation 2 to evaluate dose equivalent, and will be tested in a variety of different radiation fields. Milestones for the next twelve months are listed in Table 4.

<table>
<thead>
<tr>
<th>Date</th>
<th>Milestone</th>
</tr>
</thead>
<tbody>
<tr>
<td>June 1, 1982</td>
<td>Complete portable prototype hardware</td>
</tr>
<tr>
<td>July 1, 1982</td>
<td>Complete electronic testing and initiate testing with external radiation sources</td>
</tr>
<tr>
<td>October 1, 1982</td>
<td>Complete initial testing with monoenergetic neutrons</td>
</tr>
<tr>
<td>November 15, 1982</td>
<td>Test low-gain detector with high-energy heavy particles</td>
</tr>
<tr>
<td>January 15, 1983</td>
<td>Install operating program in read-only memory</td>
</tr>
</tbody>
</table>

NOTE: Some previous milestones were delayed approximately four months due to a period of no-cost extension of the contract.
REFERENCES


