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Contribution of High Charge and Energy (HZE) Ions During Solar-Particle Event of September 29, 1989

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Symbols

BFO	blood forming organ
С	coefficient (see eq. (3))
CAM	computerized anatomical man
CRT	cosmic ray telescope
DFE	degree of iron (FE) enhancement
Ε	kinetic energy, MeV/amu
GOES	Geostationary Operational Environmental Satellite
GSFC	Goddard Space Flight Center
Η	dose equivalent, cSv
HZE	high charge and energy
IMP	Interplanetary Monitoring Platform
LET	linear energy transfer
NCRP	National Council on Radiation Protection and Measurements
p(E)	momentum, $\sqrt{E(E+1876)}$
p ₃₀	momentum corresponding to proton energy of 30 MeV, 239.15 MV
SEPB _z	solar energetic particle baseline for atomic charge
SPE	solar-particle event
VLET	very low energy telescope
Ζ	atomic charge
$\alpha(E)$	enhancement index
β	proton velocity relative to velocity light
Φ	integral fluence, particles/cm ²
$\phi_p(E)$	differential fluence, protons/(cm ² -MeV)

 $\phi_Z(E)$ energy spectra of atomic charge

Abstract

The solar-particle event (SPE) of September 29, 1989, produced an iron-rich spectrum with energies approaching 1 A GeV with an approximate spectral slope parameter of 2.5. These high charge and energy (HZE) ions challenge conventional methods of shield design and assessment of astronaut risks. In the past, shield design and risk assessment have relied on proton shielding codes and biological response models derived from X-ray and neutron exposure data. Because the HZE spectra decline rapidly with energy and HZE attenuation in materials is limited by their penetration power, details of the mass distributions about the sensitive tissues (shielding materials and the astronaut's body) are important determining factors of the exposure levels and distributions of linear energy transfer. Local tissue environments during the SPE of September 29, 1989, with its HZE components are examined to analyze the importance of these ions to human SPE exposure. Typical space suit and lightly shielded structures leave significant contributions from HZE components to certain critical body tissues and have important implications on the models for risk assessment. A heavily shielded equipment room of a space vehicle or habitat requires knowledge of the breakup of these ions into lighter components, including neutrons, for shield design specifications.

Introduction

Solar-particle events (SPE's) have long been recognized as a potential hazard to human operations in space. Recently, the SPE of August 4-7, 1972, was analyzed, and the inclusion of dose rate effects on risk assessment was found to reduce health effects of exposure by a factor of 3 in comparison with a single high-dose-rate exposure (ref. 1). In that calculation, only incident protons were considered, as that was the data set available for analysis. A large SPE, occurring on September 29, 1989, contained an iron-rich spectrum with energies approaching 1 A GeV and an approximate energy power index of 2.5 (ref. 2). This SPE is the largest high-energy event of the space era (1959 to present), and 10 times this event matches the ground level data of the SPE of February 23, 1956. It suggests a 10-scaled event as a worst case event for a design guide of future deep space mission.

To analyze the importance of HZE ions to human SPE exposure and for a design guide for future deep space missions, the event-integrated fluences of the SPE of September 29, 1989, are constructed from the limited measured data. The proton spectrum above 30 MeV is well described by a model developed by Nymmik (refs. 3 and 4). Below that energy, an exponential distribution of protons developed by Shea and Smart is used (ref. 5). The enhancements of the high charge and energy (HZE) ion abundances, which may present new and significant problems for radiation protection, are calculated relative to solar energetic particle baseline (SEPB) abundances (ref. 6). By using the HZETRN code systems (ref. 7), transport properties of the shielding materials and an astronaut's body tissues are calculated. The typical shield configurations are assumed to be equivalent aluminum structures. The astronaut geometry is taken from the computerized anatomical man (CAM) model (ref. 8). The risk assessment and the risk contributions of particle groups are made with the full spectrum of the SPE of September 29, 1989, to characterize local tissue environments.

Energy Spectra of SPE of September 29, 1989

Analytic representation of proton spectra is made according to the model of Nymmik (refs. 3 and 4) for energies above 30 MeV and the exponential rigidity spectra of Shea and Smart (ref. 5) for energies below 30 MeV. From the model of Nymmik (refs. 3 and 4), the event-integrated proton fluence above 30 MeV of this event is given by

$$\int_{30}^{\infty} \phi_p(E) \ dE = 1.39 \times 10^9 \text{ protons/cm}^2 \quad (1)$$

where E is the kinetic energy, and $\phi_p(E)$ is the differential fluence. For the proton differential fluence above 30 MeV, $\phi_p(E)$ is given as a power law by

$$\phi_p(E) \ dE = \frac{C}{\beta} \times \left[\frac{p(E)}{p_{30}}\right]^{-4.5} \ dE \qquad (2)$$

where β is the proton velocity relative to the velocity of light, the momentum $p(E) = \sqrt{E(E + 1876)}$, and p_{30} is the momentum corresponding to a proton energy of 30 MeV, which is equivalent to 239.15 MV. The coefficient C is calculated from equation (2) as

$$C = \frac{1.39 \times 10^9}{\int_{30}^{\infty} \left[\frac{239.15}{p(E)}\right]^{4.5} \frac{1}{\beta} dE} = 2.034 \times 10^7 \quad (3)$$

The differential fluence below 30 MeV is calculated by using an exponential distribution (ref. 5) because there is a flattening of the spectra below 30 MeV based upon observations of this event (ref. 9). This calculation gives a very good empirical description of data below 30 MeV, and the integral energy spectra above 1 MeV of this event are well described with measured data of NOAA (ref. 10) as shown in figure 1.

Empirical representations of oxygen and iron spectra are given by Tylka, Dietrich, and Boberg (ref. 2) as shown in figures 2 and 3. In these figures, the mean spectral slopes are 3.9 and 2.5 for oxygen and iron particles, respectively. To obtain the abundance distributions from He to Ni (Z = 2 to 28), oxygen is chosen as the element for the SEPB to which the relative abundances of those ions are normalized because its statistics are the best for the ions heavier than He (ref. 6). The defined SEPB values in reference 6 are reproduced in table 1, with which the energy spectra from He to N (Z = 2 to 7) are given as

$$\phi_{Z}(E) = \operatorname{SEPB}_{Z} \times \phi_{\Omega}(E) \tag{4}$$

where $\phi_{\Omega}(E)$ is the energy spectra of oxygen.

In some events, an enhancement of heavy ion abundances is seen relative to an SEPB that increases with increasing atomic number. These enhancements have some correlation with spectral slope (ref. 6). The SEPB Fe/O ratio is 0.066, but for the SPE of September 29, 1989, the fluence ratio for Fe/O is approximately 0.2, which demonstrates that this event is enriched in Fe threefold. This enhancement is viewed as a distinct physical process superposed on some underlying pattern for ions heavier than oxygen, and their spectra are classified by the degree of Fe enhancement (DFE) with increasing atomic number Z beyond oxygen as follows:

DFE(Z, E) =
$$\left(\frac{Z}{8}\right)^{\alpha(E)}$$
 (5)

Here, the enhancement index $\alpha(E)$ is given from the energy spectra of oxygen and iron as

$$\alpha(E) = \frac{\ln\left[\frac{\phi_{\text{Fe}}(E)}{\phi_{\text{O}}(E)}\left(\frac{1}{0.066}\right)\right]}{\ln\left(\frac{26}{8}\right)}$$
(6)

The degree of enhancement at 100 A MeV of each atomic charge number beyond oxygen (eq. (5)) is shown in figure 4 for its abundance relative to the baseline abundance. With the degree of enhancements and the SEPB abundances, the data for ions from F to Mn (Z = 9 to 25) are obtained as

$$\phi_Z(E) = \text{SEPB}_Z \phi_O(E) \left(\frac{Z}{8}\right)^{\alpha(E)}$$
 (7)

For Co (Z = 27) and Ni (Z = 28) energy spectra, one thirtieth and one tenth of Fe ion spectra are taken, respectively, which are the relative abundances in cosmic ray fluence. The relevant integrated fluence spectra of the SPE of September 29, 1989, are shown in figure 5. These spectra are used as input to the transport code systems, HZETRN (ref. 7), for the risk analysis of the SPE of September 29, 1989.

Critical Regions Considered Inside Shield for SPE Protection

Particles arriving at some remote location from the Sun are diffusing through the interplanetary media and show some anisotropy because in the leading edge of the expanding radiation field, back-scattered particles are absent. Following the first 20 to 30 min after initial particle arrival, isotropy is usually achieved; therefore, the radiation fields that are incident on the spacecraft are assumed to be isotropic. Simple spacecraft geometry is chosen in which an astronaut is assumed to be at the center of a large spherical shell of uniform material. Three typical shield representations for a spacecraft are space suit, pressure vessel, and equipment room. These shields are assumed as equivalent aluminum structures: space suit with a thickness of 0.3 g/cm², pressure vessel with a thickness of 1 g/cm², and equipment room with a thickness of 5 g/cm² (ref. 11).

Inside a shield enclosure, three critical organs considered in this study are skin, ocular lens, and blood forming organ (BFO). In order to determine spatially the positions of these critical sites, the computerized anatomical man (CAM) model, which contains 2400 separate geometric tissue regions of several different elemental compositions and densities (ref. 8), is used. The detailed mass distributions of body geometry from 512 rays are used to generate fluences of the various components of solar particles at each site inside a shield.

Risk Assessment at Local Tissue Environment

The incident fluences from the SPE of September 29, 1989, inside typical space shields are calculated with the HZETRN code system (ref. 7). The interior radiation environment is transported through the inhomogeneous and geometrically complex body to calculate the specific local tissue environment. The radiation risks in terms of conventional dosimetry, dose and dose equivalent with quality factor (ref. 12), are estimated at local tissue environments. These are shown in tables 2 and 3.

The dose equivalent contributions of particle groups to the skin inside shields are shown in figure 6. Skin is the least dependent on individual body masses, and the fractional contribution of protons increases as a function of equivalent aluminum shielding because of the breakup of multiple charged ions and secondary production from HZE nuclear interactions with mainly shield materials, as evident from figure 6. At the ocular lens, the fractional contributions of all the particle groups show the same trend as for skin but with a lesser degree of change as shown in figure 7. This trend is caused by an increased shielding by body masses than for skin. The fractional contributions of all the particle groups at BFO are nearly independent of the shield thickness as shown in figure 8. It is reasoned that BFO is the most dependent on individual body masses and all the radiation components are attenuated to some degree (ref. 13) before reaching the specific target site of BFO. The most body-shielding effect is achieved at BFO among the sensitive sites considered. The risk contribution is further analyzed to study the detailed local tissue environment and is presented as follows.

The differential spectra of dose equivalents in figures 9 through 11 show the detailed risks calculated for local tissue environments. In these figures, the height of each curve is proportional to the contribution for that energy interval. At skin and ocular lens within a shield, most HZE particles have not reached the end of their range, but after passing through a shield and body masses they have been slowed down to higher LET and fragmented into smaller HZE's that have higher quality factors. At BFO, which is rather deep into the body, most HZE particles have been substantially stopped. The energy levels at the peak contribution of dose equivalent are shifted to higher energy levels for all the particle groups, as the sensitive site shifts to deeper within the body. This change is because HZE particles with high penetration power become the most effective radiation constituent at that location. Also noticeable are still significant risk contributions from HZE particles to these tissues inside a space suit and pressure vessel, as shown in figures 9 and 10.

In figures 9 through 11, two sources of secondary particles for atomic charge number 1 and 2 are identified. Differential dose equivalent contributions at low energy correspond to targetlike fragments, and those at high energy are from projectilelike fragments (ref. 14). Relatively abundant low-energy protons and alpha particles have been produced by nuclear interactions of all primaries in shielding materials and body, and they contribute substantially to the calculated dose equivalent. Detailed local tissue environments are determined by the mass distributions of shielding materials and body geometry.

In table 3, dose equivalent calculations of the SPE are given along with the NCRP 30-day limit (ref. 15). These limits are set to preclude both acute and late nonstochastic effects by not exceeding the threshold levels to these effects. From the table, total dose equivalent at each local tissue is significantly reduced as shield thickness is increased from space suit, to pressure vessel, and to equipment room. But dose equivalent in the equipment room only is within the acceptable NCRP 30-day limit at each local tissue. Therefore, only the equipment room provides sufficient shielding to avoid acute or nonstochastic late effects in sensitive organs by the NCRP 30-day limit.

For the current risk estimation, the dose equivalent is calculated by using the defined quality factors (ref. 12). These quality factors are derived principally for carcinogenesis and mutagenesis, and the actual risk estimates for acute exposure may be somewhat lower than the current estimation. The current estimation is approximated at best, since quality factors for early radiation effects are not defined (ref. 12). Furthermore, not only track structure effects on injury cross sections are poorly understood, but there is a lack of detailed spectral data for alpha particles and heavier ions emitted during the SPE. Because the full heavy ion spectra of the SPE of September 29, 1989, are constructed from the limited measured data, the current risk estimation for this SPE will be changed undoubtedly due to these uncertainties.

The current risk estimation in table 3 shows that skin dose equivalent inside a space suit would be very large (≈300 Sv) for an event of similar spectral characteristics and 10 times intensities of the SPE of September 29, 1989, such as the SPE of February 23, 1956. Even a dose rate reduction factor of 2 to 3 (ref. 1) would leave the exposures high compared with the threshold for moist desquamation of 30 Gy (30 Sv) for gamma rays (ref. 15). The mortality threshold is 1.5 Gy (1.5 Sv) for gamma rays, which is much lower than 4.2 Sv to BFO inside a space suit for a "February 23, 1956 SPE." In addition to the possibility of skin infection with an already depressed immune system, the coincident BFO exposure may provide a serious medical problem to the astronauts. Clearly, more shielding is required even within an equipment room to protect the critical tissues during most of SPE's. An understanding of the breakup of HZE ions into lighter components, including neutrons, is required for shield design specification because all HZE particles are not stopped as has been shown in figures 9 through 11.

The buildup of neutrons is harmful even inside an equipment room because they are an important source of the proton and He ion exposures within the tissues. Indeed, for the contributions of dose equivalents of Z = 1 and 2 at low energy near 0.1 to several MeV/ amu, these particles result from neutrons produced in the shielding materials.

Concluding Remarks

The event-integrated fluences of the solar-particle event (SPE) of September 29, 1989, are calculated according to the descriptions of the observed spectra. This SPE was threefold enriched in Fe relative to the baseline composition of SPE. The risk estimations . clearly show that critical organs respond differently to their local environments. To obtain accurate local tissue environments, the detailed mass distributions of shielding materials and the sensitive tissues are required, and they are important determining factors for accurate exposure level assessments and linear energy transfer (LET) distributions. The result shows that high charge and energy (HZE) ions play a limited role inside a lightly shielded space suit and for the least shielded organs by the NCRP 30-day limit during the SPE of September 29, 1989.

The detailed risk analysis according to the differential spectra of dose equivalent shows that even a heavily shielded equipment room cannot stop completely all energetic particles, which may break up into lighter components including neutrons from the body. Therefore, a well-shielded region and medical treatment must be considered when a large SPE occurs during deep space mission.

For accurate risk estimations and shield design specifications, the spectral data for alpha particles and heavier ions emitted during SPE's are required as well as quality factors for early radiation effects. Improved understanding of track structure effects on injury cross sections and accurate nuclear databases including neutrons are required to decrease the risk estimate uncertainties.

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[From ref. 6]							
Particle	SEPB _z	Particle	SEPBZ				
H/O	3500 ± 500	Na/O	0.0083 ± 0.0015				
He/O	53 ± 5	Mg/O	0.183 ± 0.010				
Li/O	<0.001	Al/O	0.0115 ± 0.018				
Be/O	$<5 \times 10^{-4}$	Si/O	0.147 ± 0.009				
B/O	$<5 \times 10^{-4}$	P/O	0.0014 ± 0.0006				
C/0	0.454 ± 0.018	S/O	0.0229 ± 0.0025				
N/O	0.129 ± 0.008	Ar/O	0.0016 ± 0.0007				
0/0	1.00 ± 0.031	Ca/O	0.0076 ± 0.0016				
F/O	<5 × 10 ⁻⁴	(Ti+Cr)/O	0.0024 ± 0.0009				
Ne/O	0.128 ± 0.008	Fe(group)/O	0.066 ± 0.006				

Table 1. Solar Energetic Particle Baseline

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Table 2. Dose for SPE of September 29, 1989

	Dose, cGy, at—									
Atomic Charge	Skin inside—			Ocular lens inside—			BFO inside—			
	Space suit	Pressure vessel	Equipment room	Space suit	Pressure vessel	Equipment room	Space suit	Pressure vessel	Equipment room	
Z = 1	780.9	296.0	39.9	377.2	193.5	36.8	27.6	23.4	13.1	
<i>Z</i> = 2	105.2	10.2	0.7	16.7	4.8	0.6	0.4	0.4	0.2	
$3 \le Z \le 10$	2.9	0.3	≈ 0	0.5	0.1	≈ 0	≈0	≈ 0	≈ 0	
$11 \le Z \le 20$	0.7	0.1	≈ 0	0.2	0.1	≈ 0	≈ 0	≈ 0	≈ 0	
$21 \le Z \le 28$	0.3	0.1	≈ 0	0.1	≈ 0	≈ 0	≈ 0	≈ 0	≈ 0	
Total	890.0	306.6	40.6	394.5	198.4	37.4	28.0	23.8	13.3	

	Dose, cGy, at									
Atomic Charge	Skin inside—			Ocular lens inside—			BFO inside—			
	Space suit	Pressure Vessel	Equipment Room	Space suit	Pressure Vessel	Equipment Room	Space suit	Pressure Vessel	Equipment Room	
Z = 1	1738.3	553.7	57.7	668.1	317.8	50.1	37.8	31.4	16.8	
Z = 2	1149.3	81.9	6.6	132.3	33.1	4.9	4.0	3.5	2.4	
$3 \le Z \le 10$	53.2	5.3	0.2	9.3	2.4	0.2	0.1	0.1	≈ 0	
$11 \le Z \le 20$	9.0	1.6	0.1	2.4	0.9	0.1	0.1	0.1	≈ 0	
$21 \le Z \le 28$	2.4	0.8	0.2	1.0	0.5	0.1	0.1	0.1	≈ 0	
Total	2952.2	643.1	64.7	813.1	354.7	55.4	42.2	35.1	19.3	
NCRP 30-day limit (ref. 15)		150			100			25		

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Table 3. Dose Equivalent for SPE of September 29, 1989



Figure 1. Analytic representation of proton spectra for SPE of September 29, 1989.



Figure 2. Empirical representation of oxygen spectra for SPE of September 29, 1989 (ref. 2).



Figure 3. Empirical representation of iron spectra for SPE of September 29, 1989 (ref. 2).



Figure 4. Prediction of September 29, 1989, solar event abundances relative to the SEPB abundances for each atomic charge number Z.



Figure 5. Event-integrated integral fluence spectra of SPE of September 29, 1989.



Figure 6. Dose equivalent at skin for SPE of September 29, 1989.



Figure 7. Dose equivalent at ocular lens for SPE of September 29, 1989.



Figure 8. Dose equivalent at BFO for SPE of September 29, 1989.



Figure 9. Differential spectra of dose equivalent per logarithmic interval of energy per nucleon at different regions inside space suit.



Figure 10. Differential spectra of dose equivalent per logarithmic interval of energy per nucleon at different regions inside pressure vessel.



Figure 11. Differential spectra of dose equivalent per logarithmic interval of energy per nucleon at different regions inside equipment room.

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