Space Radiation and Risks to Human Health

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To live and work safely in space with acceptable risks from radiation

- The space radiation environment poses both **acute & chronic** risks to crew health and safety, with clinically relevant implications for the lifetime of the crew.
- “Safely” means that **risk limits** are sufficiently understood, defined, and not exceeded.
- Because of unique properties of space radiation, estimating risks from space radiation exposure carries **large uncertainties** that constrain mission planning, with impacts on mission duration, crew selection and development of effective mitigation strategies.
- NASA needs to close the knowledge gap on a broad-range of biological questions before radiation protection goals can be met for exploration.
- The Human Research Program (HRP), Space Radiation Program Element (SRPE) led by JSC is committed to solving the space radiation problem with the goal of uncertainty reduction in risk estimates for Mars missions (<50%) to support agency goals in vehicle, mission, and crew selection.
Space Radiation Environment

Solar Particle Events (SPE)
- Consist of low to medium energy protons
- Risk of acute radiation effects for unshielded exposure to large event
- Effectively blocked by shielding but optimization required to reduce weight
- Accurate event alert and dosimetry are essential for crew safety
- Improved understanding of radiobiology needed to perform risk model optimization

Galactic Cosmic Rays (GCR)
- Penetrating protons and highly charged, energetic atomic nuclei (HZE particles)
- Abundances and energies in space environment understood, but large biological uncertainties limit ability to accurately evaluate risks
- Heavy ions are not effectively shielded (break up into lighter, more penetrating fragments)
- Uncertainties cloud understanding of effectiveness of possible mitigations
**X-rays and Gamma Radiation**

- X-rays are photons (electromagnetic radiations) emitted from electron orbits, such as when an excited orbital electron "falls" back to a lower energy orbit.

- Gamma rays are photons emitted from the nucleus, often as part of radioactive decay.

- Considered low linear energy transfer
Space radiation is composed of high-energy protons and heavy ions (HZEs) along with secondary protons, neutrons, and heavy ions produced via interaction with shielding materials.

- **Linear energy transfer (LET)**
  - Energy loss per unit path length as particles pass through matter.

- High LET defined as LET > 10 keV/µm in tissue.

- HZE nuclei traversal causes unique damage to biomolecules, cells, and tissues.

- **Distinct damage**
  - DNA and oxidative damage (ROS) and spatial patterns distinct from terrestrial radiation (x-rays and gamma rays).

- **Distinct biological effects and health risks?**

- No human data to estimate risk
  - Must use animal and cellular models.

1 GeV/u $^{56}$Fe nucleus LET $\sim$ 150 keV/µm

Qualitative differences due to track “core” and correlated tissue damage along a particle path.

(Plante, 2011)
**Linear Energy Transfer (LET)**

- Radiation dose is the amount of energy per unit of biological material

- The LET is related to Biological Damage

- The average energy locally imparted to a medium by a charged particle of specified energy, per unit distance traversed. Measure of the ion pairs generated/cm of tissue traversed

- Gamma and X-Rays are low-LET and are sparsely ionizing producing uniform ionization across a tissue

- HZE Particle Radiation from GCR are high-LET.

- The localized DNA damage caused by dense ionizations from high-LET radiations is more difficult to repair than the diffuse DNA damage caused by the sparse ionizations from low-LET radiations

- The higher the LET, the higher the Q-quality factor in determining dose equivalent (Severts, where 1 Sv = 100 rem)
The Space Radiation Problem

GCR and secondary fragmentation produced by interaction w/ shielding materials

- Unique damage to biomolecules, cells, and tissues occurs from HZE ions
- No human data to estimate risk from heavy ion damage- large uncertainty
- Animal models must be applied or developed to estimate cancer and other risks
- Shielding has excessive costs and will not eliminate galactic cosmic rays (GCR)

Track structure of deposited energy (of what’s seen biologically at left)

HZE ion traversals in cells and DNA breaks

Single HZE ions in photo-emulsions
Leaving visible images
Space Safety Requirements

• Congress has chartered the National Council on Radiation Protection (NCRP) to guide Federal agencies on radiation limits and procedures
  – NCRP guides NASA on astronaut dose limits
• Crew safety
  – limit of 3% fatal cancer risk at 95% Confidence Level
  – prevent radiation sickness during mission
  – new exploration requirements limit Central nervous system (CNS) and Heart disease risks from space radiation
• Mission and Vehicle Requirements
  – shielding, dosimetry, and countermeasures
• NASA programs must follow the ALARA principle to ensure astronauts do not approach dose limits

Cell fusion caused by radiation

Space Radiation in breast cancer formation
Future Manned Missions

**International Space Station**
- 2013-2020: 6-person crews, 180 days (nominal); 2-person crew 360 days in planning
- Approach limits for acceptable radiation risks after 1 to 3 missions

**Lagrange Points**
- Design Reference Mission currently being formulated
- Outside Earth’s magnetosphere and radiation belts
- Galactic cosmic ray risks are major concern

**Near Earth Objects**
- Design Reference Mission currently being formulated
- Outside Earth’s magnetosphere and radiation belts
- Galactic cosmic ray risks are major concern

**Mars**
- 2030 and beyond: 6-person crews, up to 1000 days
- Long deep space transit times
- Risks exceed NASA Permissible Exposure Limits (PELs) for cancer, and pose significant non-cancer risks
Integrated Radiation Protection Strategy Enables Human Mars Exploration

Long-Term Commitment across Research and Technology Required...

Mission and Architecture Systems Analysis

- Near Earth Asteroid Systems
- Mars NTV
- In-situ Resource Utilization
- Active Shielding Concepts

Environmental Modeling, Monitoring, and Prediction

- Predictive Models
- Precursor Data — MSL RAD
- On-board Dosimetry — ISS TBPC

Crew Selection and Operations

- Human Digital Twin

Radiobiology and Biological Countermeasures

- NASA Space Radiation Lab at Brookhaven National Laboratory
- X-ray vs. Heavy Ion Track Damage to DNA
- Leukemia Induction with GCR — Mouse Model

Innovative Multi-Purpose Shield Solutions

- Heavy Ion Testing of Inflatable Shield Prototype
- Water Filled Composite Shield Solutions
- Reconfigurable Personal Shielding
- Hydrogen Storage BNNT

Advances benefit homeland security, cancer therapy, Earth observing and communication satellites, and commercial air safety

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### Space Radiation Risks

**Risk of Radiation Carcinogenesis**
- Morbidity and mortality risks
  - Major drivers for PELs and therefore major research emphasis for program

**Risk of Acute & Late Central Nervous System Effects from Radiation Exposure**
- Possible in-flight changes in motor function, cognition, and behavior
- Late neurological disorders
  - SRPE aligning with Behavioral Health and Performance Element for possible future collaboration on acute risk monitoring

**Risk of Degenerative Tissue or Other Health Effects from Radiation Exposure**
- Degenerative changes in the heart, vasculature, and lens
- Diseases related to aging, including digestive, respiratory disease, premature senescence, endocrine, and immune system dysfunction
  - SRPE monitoring HHC radiation research on heart risks

**Risk of Acute Radiation Syndromes due to Solar Particle Events**
- Prodromal effects (nausea, vomiting, anorexia, and fatigue), skin injury, and depletion of the blood-forming organs
- Acute radiation syndrome (ARS) from solar particle events (SPE) avoided by Alert-Dosimetry and Shielding
Foundations of SRP Research Plans

• External review by National Council on Radiation Protection (NCRP), National Academy of Sciences, and standing Radiation Discipline Working Group (RDWG)

• Simulate space radiation at the NASA Space Radiation Laboratory (NSRL)
  • Located at DoE’s Brookhaven National Lab (Long Island NY)

• Six NASA Specialized Centers of Research (NSCOR’s) studying the biology of space radiation risks

• Broad program of directed research at over 40 US Universities including collaborative research with US Department of Energy (DoE)

• Collaborate with NASA’s Science Missions on advanced SPE alert & Mars robotic missions

• Long-term goal to improve knowledge to develop individual risk assessments, countermeasures
NASA Space Radiation Lab (NSRL)

MEDICAL Dept.
- Long-Term Laboratories
- Logistic and Administrative Support
- Animal Care Facility
- Liaison Scientist
- X-Ray Source

BIOLOGY Dept.
- Management
- Logistic Support
- Gamma Source

CAD-AGS-NSRL
- Beam Line
- Target Area
- Dosimetry
- Staging Area
- Liaison Scientist

RHIC
Sources of Uncertainty

- Radiation quality effects on biological damage
  - Qualitative and quantitative differences of Space Radiation compared to x-rays
- Dependence of risk on dose-rates in space
  - Biology of DNA repair, cell regulation
- Space dosimetry and organ doses
- Predicting solar particle events
  - Temporal and size predictions
- Extrapolation from experimental data to humans
- Individual radiation-sensitivity
  - Genetic, dietary and “healthy worker” effects
Radiation Carcinogenesis
Risk Summary & Research Strategy

Risk of Radiation Carcinogenesis

• Morbidity and mortality risks for a wide variety of cancers [including lung, breast, colon, stomach, esophagus, the blood system (leukemias), liver, bladder, skin, and brain]

• Evidence for space radiation cancer risks are well known and described most recently by:
  − NCRP Reports 132 and 152
  − BEIR VII Report
  − UNSCEAR 2006 Report

• Cancer risk is a major driver for Space Radiation PELs and therefore a major research emphasis of program

• PELs guide mission, vehicle, and crew selection requirements

• The SRPE established a scientific approach that follows a progression of activities designed to feed the development of an integrated risk model with acceptable uncertainty for exploration missions, followed by activities targeting risk mitigation and monitoring
Research to reduce uncertainties in risk projection models are expected to increase NASA’s ability to select crew, extend mission duration, and reduce cost through possible reductions in shielding requirements.
Major Findings on Cancer Risk from NSRL

First experiments at NSRL were in Oct, 2003 and many publications are in preparation. Findings to date include:

- A low RBE for Leukemia from Iron due to high efficiency of apoptosis
- A high RBE for solid cancer is emerging
- Major differences in signaling pathways between high and low LET and high and low dose
The **Acute CNS risks** include altered cognitive function including short-term memory, reduced motor function, and behavioral changes, which may affect performance and human health.

- The NCRP recommends that all clinically significant acute risks must be avoided
- Further development of evidence base for acute and late CNS risks required to define areas of significant concern and establish limits

→ **SRPE in process of aligning with Behavioral Health and Performance Element for possible future collaboration on CNS Acute risk monitoring**

The **Late CNS risks** are possible neurological disorders such as Alzheimer’s disease (AD), dementia, cerebrovascular disease or premature aging.

- AD is fatal, with mean time from early stages to death approx. 8 yrs
- Inclusion in the overall acceptable REID probability for space missions if AD risk established
CNS Risks from Galactic Cosmic Rays

- In-flight acute performance changes and late effects similar to Alzheimer’s disease are a concern for GCR
- Retinal flashes observed by Apollo astronauts suggests single heavy nuclei can disrupt brain function
- Central nervous system (CNS) damage by x-rays is not observed except at very high doses
- NASA research on performance of animals is establishing threshold doses for acute CNS risks
  - studies have quantified rate of neuronal degeneration, plaque formation and changes in dopamine function related to late CNS risks
- Studies will be extended to other GCR components to establish acute CNS Permissible Exposure Limits, and to derive a risk limitation approach to neurological disorders

Reduction in number of neurons (neurodegeneration) for increasing Iron doses in mouse hippocampus

Oxidative Stress (Lipid peroxidation: 4-Hydroxynonenal) is increased in Mouse Hippocampus 9 Months After 2 Gy of **Fe Irradiation**
Radiation and Non-Cancer Effects

- Early Acute risks are very unlikely:
  - Low or modest dose-rates for SPE’s insufficient for risk of early death
  - SPE doses are greatly reduced by tissue or vehicle shielding

- Radiation induced Late Non-Cancer risks are well known at high doses and recently a concern at doses below 1 Sv (100 rem)
  - Significant Heart disease in Japanese Survivors and several patient and Reactor Worker Studies
  - Dose threshold is possible making risk unlikely for ISS Missions (<0.2 Sv); however a concern for Mars or lunar missions due to higher GCR and SPE dose
  - Qualitative differences between GCR and gamma-rays are a major concern

Vasculature damage: $\mu$m of vessel per cell after protons or Fe (PI-C. Geard Columbia U)

- Controls
- Iron Irradiated

\[
\begin{array}{|c|c|}
\hline
\text{Dose, Gy} & \text{$\mu$m vessels per cell} \\
\hline
0 & \\
1 & \\
2 & \\
3 & \\
\hline
\end{array}
\]

- Protons
- Iron

\[
\begin{array}{|c|c|}
\hline
\text{Dose, Gy} & \text{$\mu$m vessels per cell} \\
\hline
0 & \\
1 & \\
2 & \\
3 & \\
\hline
\end{array}
\]
Degenerative Risks
Risk Summary

Risk of Degenerative Tissue Effects:
- Cardiovascular and circulatory changes
- Cataract formation

Other Health Effects:
- Diseases related to aging, including digestive, respiratory disease, premature senescence, endocrine, and immune system dysfunction

Driving Evidence:
- Astronaut data (cataracts)
- Radiotherapy, environmental disasters, and atomic bomb survivor data
  - Data is confounded by life-style factors to larger extent than cancer
- Most prior work focused on high dose effects, high fat diets or other protocols that are atypical for astronauts

Risk Projections:
- Preliminary risk assessment models being formulated
- Current exposure limits set as dose thresholds; recent studies suggest there may be low dose and dose-rate effects
Mitigation Approaches

- Time in the Solar Cycle
- Radiation Shielding
  - Amounts and material types
  - Design Optimization
- Accurate Risk Quantification / Uncertainty reduction
- Crew Selection
  - Age, gender, lifestyle factors, etc,
  - Individual Sensitivity (genetic factors)
- Biological Countermeasures (BCMs)
  - Radioprotectors / Mitigators
- Biomarkers predictive of radiation induced diseases
  - Future individualized risk assessment
  - Early detection and prognostic monitoring

BCM: Pharmaceuticals

Shield Design and Optimization

Variation of Solar Activity

Individual Susceptibility

α-lipoic acid

amifostine

Potential Impact of Individual Genetic Susceptibility and Previous Radiation Exposure on Radiation Risk for Astronauts

BCM: Pharmaceuticals
Summary

Space radiation is a major challenge to exploration:
- Risks are high limiting mission length or crew selection
- Large mission cost to protect against risks and uncertainties
- New findings may change current assumptions

NASA approach to solve these problems:
- Probabilistic risk assessment framework for ISS and Exploration Trade Studies
- Ground-based research focused on uncertainty reduction at NASA Space Radiation Laboratory (NSRL)
- Collaborative research with DoE, and ESA
- Ongoing external reviews by authoritative bodies
- Well defined deliverables to Cx, ISS, and CHMO
Thank You!

As seen through the Hubble telescope, thousands of stars are forming in the cloud of gas and dust known as the Orion nebula. More than 3,000 stars of various sizes appear in this image. Some of them have never been seen in visible light.
BACK-UP SLIDES
Physical Quantities

- **HZE** = High charge (Z) and energy (E) nuclei (ions)
- **Fluence (F)** = number of particles per unit area (#/cm²)
- **Linear energy transfer (LET)** = rate of energy loss per unit distance in bulk matter (keV per micron)
- **Range** = average distance traveled before ion stops
- **Absorption cross section** = probability of nuclear reaction expressed as an Area
  - Dose (D) = Energy absorbed in bulk matter \( D = F \times \text{LET} \) (1 Gy = 100 rad)
- **Relative Biological Effectiveness (RBE)** = ratio of doses of ions to gamma rays to produce Equal biological effect
- **Quality Factor (Q)** = committee assigned value of RBE for human radiation protection
- **Dose Equivalent (H)** = D x Q (1 Sv = 100 rem) (organ shielding)
- **Risk** = *model quantity can not be measured*
<table>
<thead>
<tr>
<th>Syndrome</th>
<th>Dose*</th>
<th>Prodromal Stage</th>
<th>Latent Stage</th>
<th>Manifest Illness Stage</th>
<th>Recovery</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hematopoietic</td>
<td>&gt; 0.7 Gy (&gt; 70 rads) (mild symptoms may occur as low as 6.3 Gy or 30 rads)</td>
<td>• Symptoms are anorexia, nausea and vomiting.</td>
<td>• Stem cells in bone marrow are dying, although patient may appear and feel well.</td>
<td>• Symptoms are anorexia, fever, and malaise.</td>
<td>• In most cases, bone marrow cells will begin to repopulate the marrow.</td>
</tr>
<tr>
<td>(Bone Marrow)</td>
<td></td>
<td>• Onset occurs 1 hour to 2 days after exposure.</td>
<td>• Stage lasts 1 to 6 weeks.</td>
<td>• Primary cause of death is infection and hemorrhage.</td>
<td>• There should be full recovery for a large percentage of individuals from a few weeks to two years after exposure.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Stage lasts for minutes to days.</td>
<td></td>
<td>• Survival decreases with increasing dose.</td>
<td>• Death may occur in some individuals at 1.2 Gy (120 rads).</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>• Most deaths occur within a few months after exposure.</td>
<td>• In the LD50/60 is about 2.5 to 5 Gy (250 to 500 rads).</td>
</tr>
<tr>
<td>Gastrointestinal (GI)</td>
<td>&gt; 1.0 Gy (&gt; 1000 rads) (some symptoms may occur as low as 0 Gy or 600 rads)</td>
<td>• Symptoms are anorexia, severe nausea, vomiting, cramps, and diarrhea.</td>
<td>• Stem cells in bone marrow and cells lining GI tract are dying, although patient may appear and feel well.</td>
<td>• Symptoms are malaise, anorexia, severe diarrhea, fever, dehydration, and electrolyte imbalance.</td>
<td>• Death is due to infection, dehydration, and electrolyte imbalance.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Onset occurs within a few hours after exposure.</td>
<td>• Stage lasts less than 1 week.</td>
<td>• Death occurs within 2 weeks of exposure.</td>
<td>• Death occurs within 2 weeks of exposure.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Stage lasts about 2 days.</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiovascular (CV/ Central Nervous System (CNS))</td>
<td>&gt; 5.0 Gy (5000 rads) (some symptoms may occur as low as 20 Gy or 2000 rads)</td>
<td>• Symptoms are extreme nervousness and confusion; severe nausea, vomiting, and watery diarrhea; loss of consciousness; and burning sensations of the skin.</td>
<td>• Patient may return to partial functionality.</td>
<td>• Symptoms are return of watery diarrhea, convulsions, and coma.</td>
<td>• Ne recovery is expected.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Onset occurs within minutes of exposure.</td>
<td>• Stage lasts for hours but often is less.</td>
<td>• Great occurs 5 to 6 hours after exposure.</td>
<td>• Death occurs within 2 days of exposure.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Stage lasts for minutes to hours.</td>
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</tr>
</tbody>
</table>

* The absorbed doses quoted here are "gamma equivalent" values. Neutrons or protons generally produce the same effects as gamma, beta, or X-rays but at lower doses. If the patient has been exposed to neutrons or protons, consult radiation experts on how to interpret the dose.

† The LD50/60 is the dose necessary to kill 50% of the exposed population in 60 days.

‡ The LD100 is the dose necessary to kill 100% of the exposed population.
Solar Cycle Effects

• The solar cycle is approximately 11-years in length, however variations in length of ±2 y can occur
  – Doses from SPEs are highest at solar maximum when solar activity is highest
  – Doses from GCR are highest at solar minimum when the solar wind is strongest
• Each cycle will have varying modulation conditions and number and sizes of SPE
• The prediction of solar conditions temporal patterns are uncertain for future solar cycles
Value Of Uncertainty Reduction:
Cost of research to reduce uncertainties 2-fold, much less than cost of shielding in space or reducing mission length.