

# Space Radiation Environment and Human Exploration

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# Radiation Exposure



## Background

- Cosmic rays
- Air (radon)
- Building material
- Water
- Food
- Earth

## Occupational and Other

- Air and Space travel
- Testing fallout /contamination
- TV & luminous watches
- Nuclear power plants (20%)
- Radioactive waste
- Diagnosis & therapy\*



\*Diagnostic radiology > 200 million procedures/year (USA). Two billion procedures world-wide. High dose/partial body.

Most of our understanding of radiation effects comes following a single acute exposure



## Hiroshima, 1945

Killed outright	100,000
Survivors studied	86,572
Expected cancers (no bomb)	21,643
Observed cancers	22,064*
Increase	421*

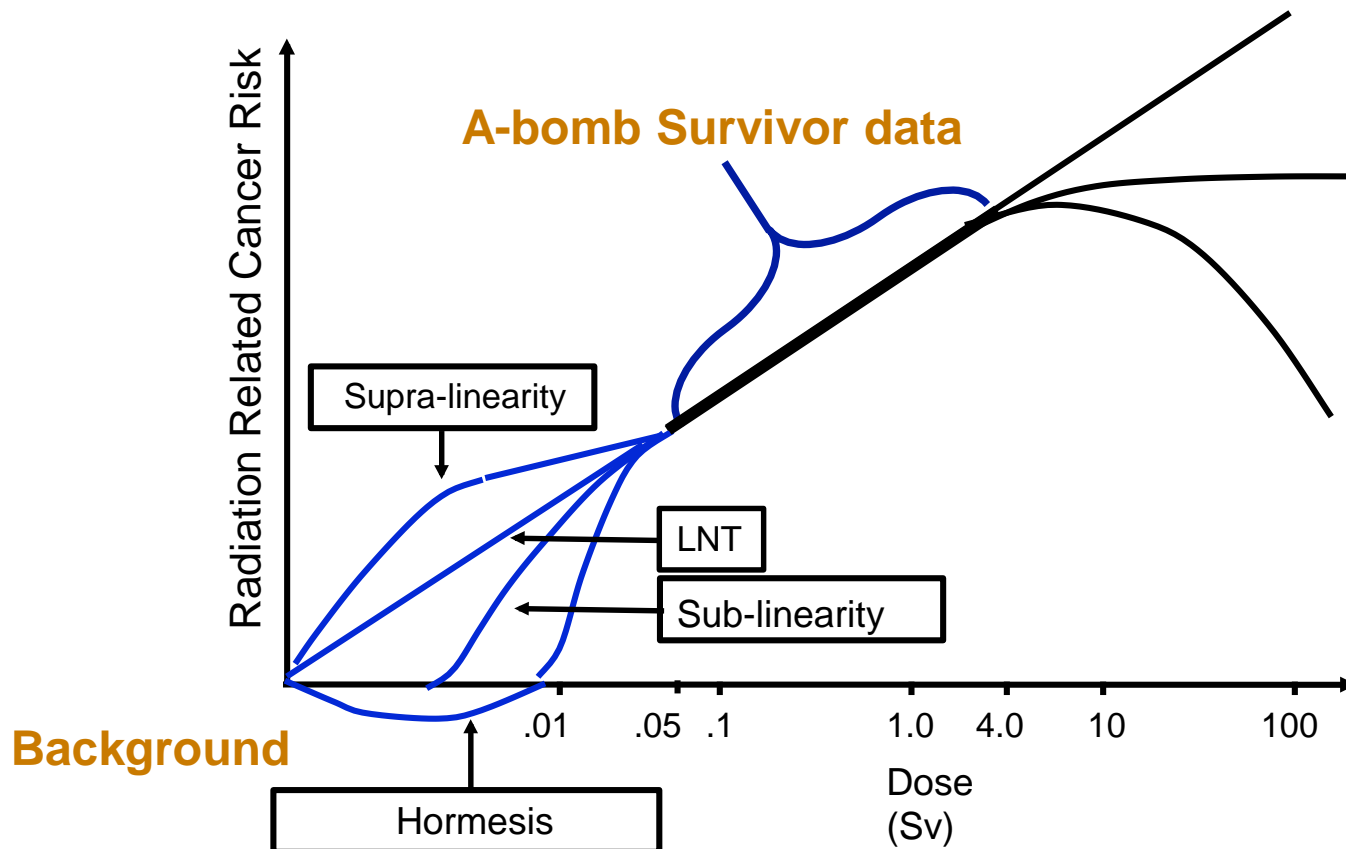
Increased non cancer deaths:  
Circulatory, digestive and  
respiratory diseases

\*Data not updated

# What is the issue?



The shape of the dose response at low radiation doses (< 100 mGy) and low dose rates for radiation-induced cancer is a critical judgement for radiation policy and risk assessment.





# Specific Questions being Addressed



Are the mechanisms of action the same for low and high doses of radiation?

Do we need to change current paradigms in radiation biology?

Is the LNT hypothesis an accurate scientific description for the dose-response relationship for cancer (or other risks) in the low dose region?

How do non-targeted effects influence cancer and non-cancer risks after irradiation?

**How does this translate to space radiation exposures?**



# The Space Radiation Environment

**Solar particle events (SPE)** (generally associated with Coronal Mass Ejections from the Sun):

medium to high energy protons

largest doses occur during maximum solar activity

not currently predictable

MAIN PROBLEM: develop realistic forecasting and warning strategies

**Trapped Radiation:**

medium energy protons and electrons

effectively mitigated by shielding

mainly relevant to ISS

MAIN PROBLEM: develop accurate dynamic model

**Galactic Cosmic Rays (GCR)**

high energy protons

**highly charged, energetic atomic nuclei (HZE particles)**

not effectively shielded (break up into lighter, more penetrating pieces)

abundances and energies quite well known

MAIN PROBLEM: biological effects poorly understood but known to be most significant space radiation hazard

# GCR High Energy



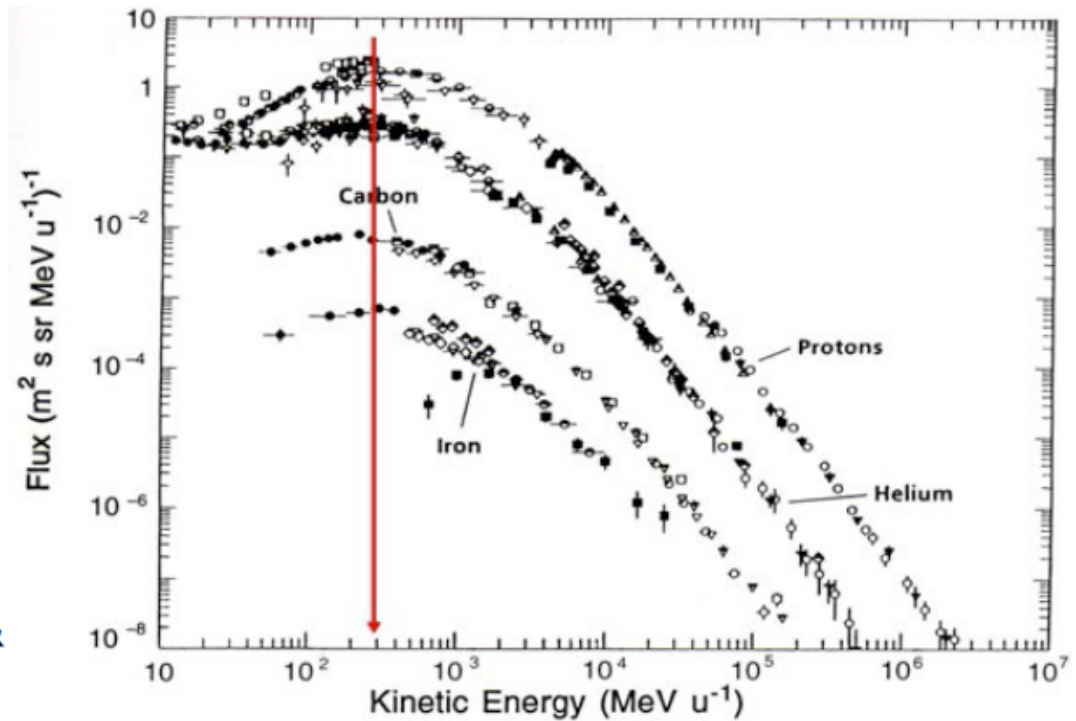
## Space radiation problem

## GCR (primary) composition

- 98% nuclei, 2%  $e^+ e^-$
- Nuclear component:
  - 87% Hydrogen
  - 12% Helium
  - 1% heavy nuclei

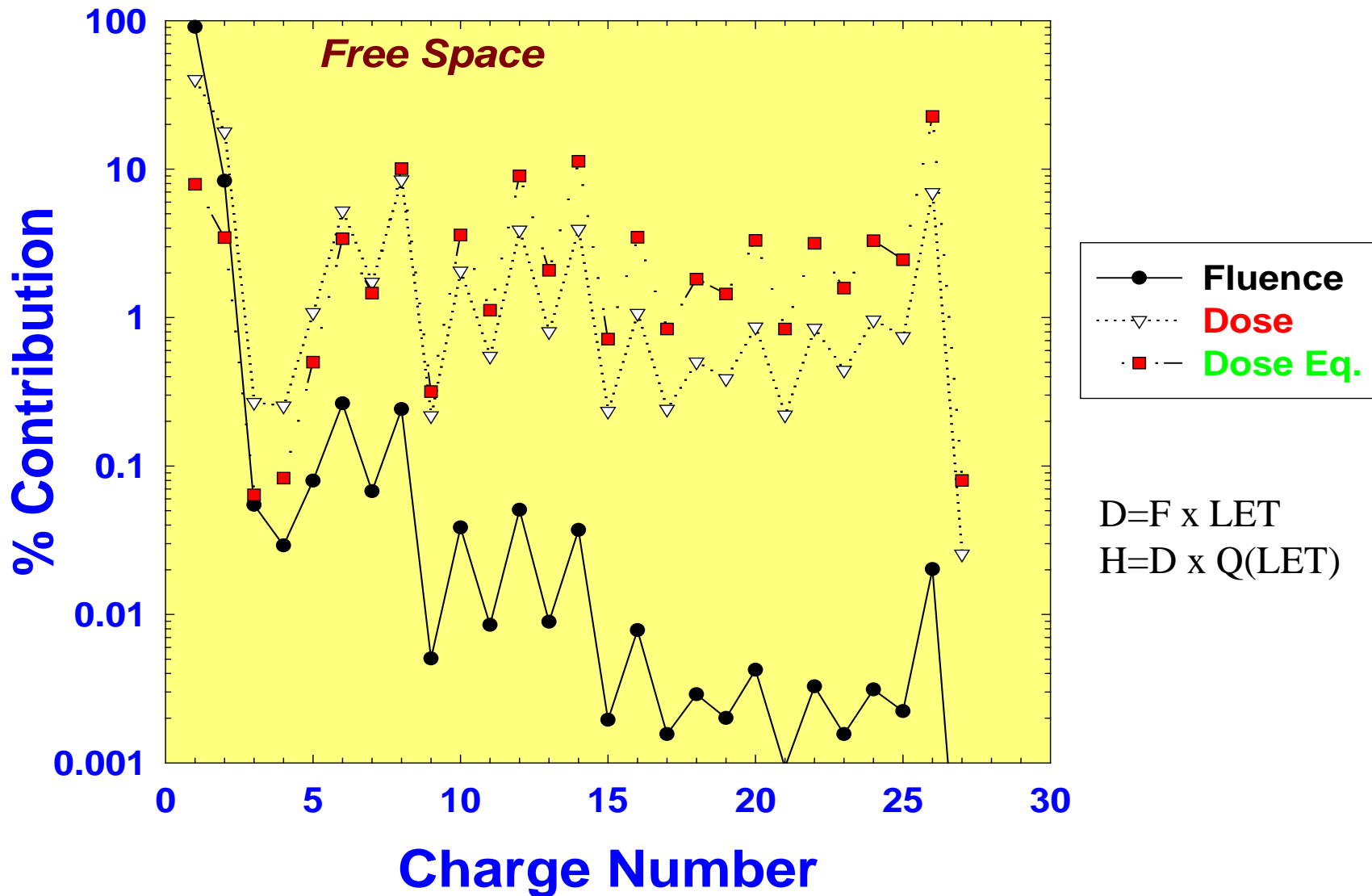
## GCR origin

- Emitted in stellar wind & flares & accelerated by supernova shock waves (within our Galaxy)



[Simpson, Ann. Rev. Nucl. Part. Sci. 33, 323, 1983]

# GCR Charge Contributions





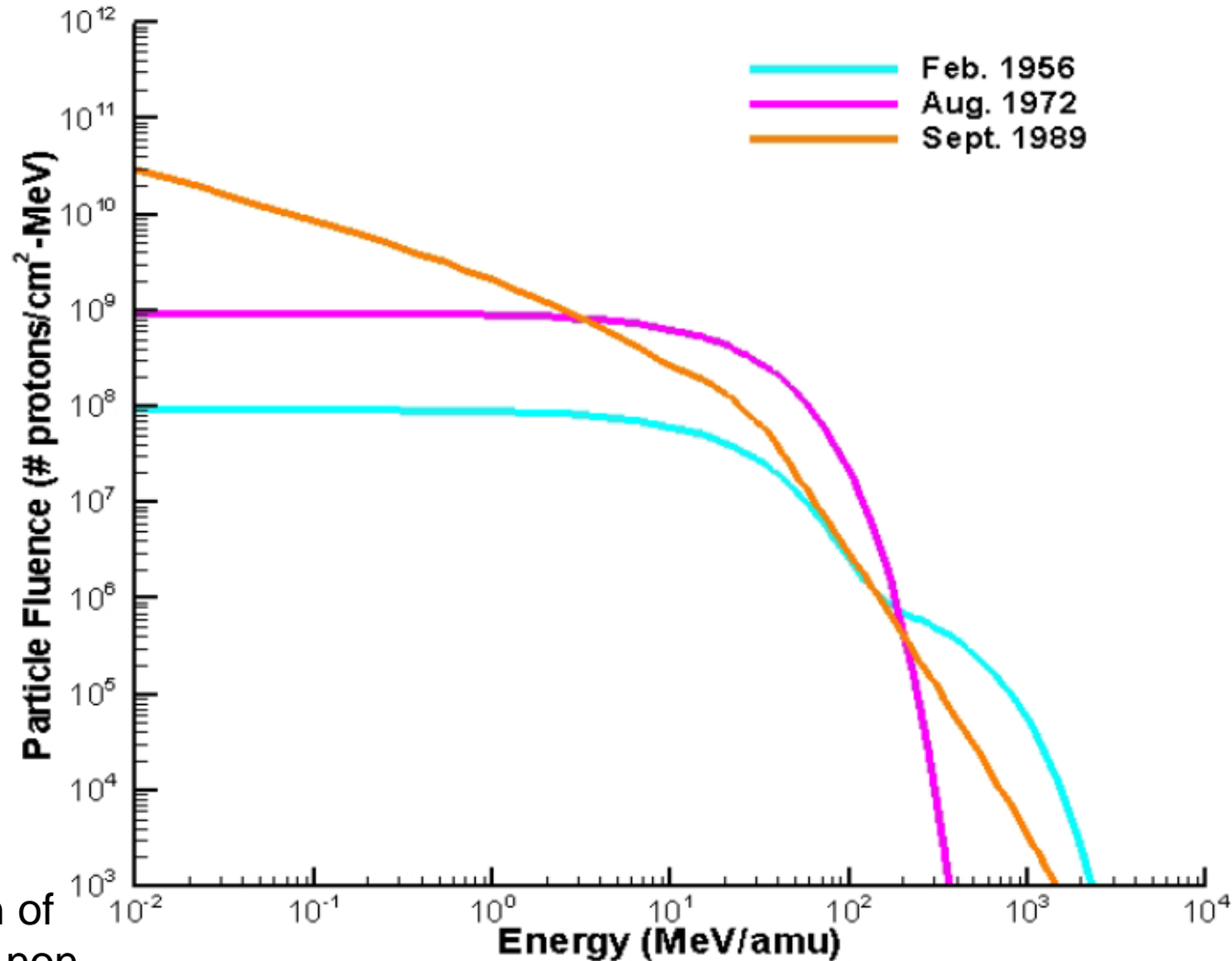
# Solar Particle Events



e, p & some heavy nuclei

$< 1 \text{ GeV/N}$

( $v \sim 0.9c$ )



Energy/dose deposition of protons in tissue highly non-linear

SPE Proton spectra [Cloudslev et al., AIAA, 2006]

# The Space Radiation Problem



- The space radiation environment is unique
  - High energy protons and heavy ions (HZE)
  - Secondary protons
  - Neutrons
  - Heavy ions produced in shielding
- Unique damage to biomolecules, cells, and tissue
- No human data to estimate risk from heavy ions
- Solar particle events cannot currently be predicted with sufficient warning
- Shielding not practical (weight, cost, will not eliminate HZE due to fragmentation)

**Primarily a low dose/low dose rate problem**

# Biological Temporal and Spatial Scales



- Biological time and spatial scales as well as linear or non-linear response modes must be known relative to GCR fluxes (type and rate) to properly simulate:
  - **protracted exposures are essential**
- Temporal:
  - Initial damage and radiation chemistry per nucleus <1 sec
  - DNA Damage repair 10 minutes to ~1 day
  - Apoptosis and differentiation ~0.5 to several days
  - Cell turn-over rates- Tissue specific from ~1 to 30 days
  - Tumor latency (post initial damage) in humans 2 to 30 yrs
  - Tumor latency in mice 100 to 800 days
  - CNS Adaptation time to GCR insult??
  - Lifespan in astronauts ~80 yrs
  - Lifespan in mice ~1000 days
  - Mars mission 1000 days

# Spatial Scales- continued



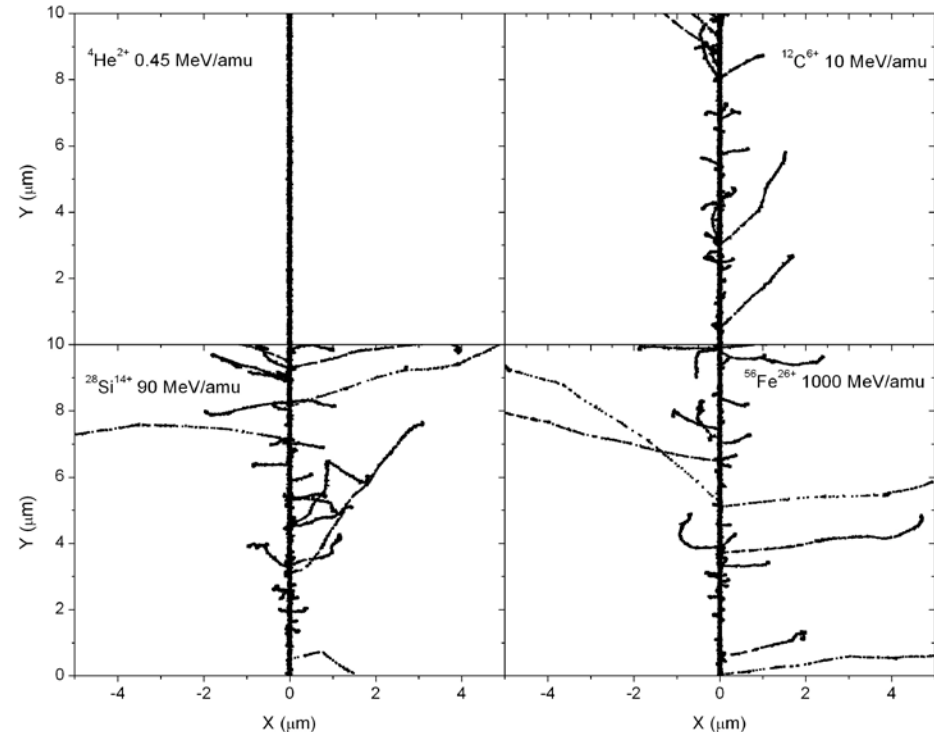
- Space Radiation (GCR proton hits/day ; HZE)
  - DNA damage repair foci  $\sim 0.5 \mu\text{m}^2$  (0.002 per day /  $1 \times 10^{-6}$ )
  - Chromosome cross sections  $\sim 2 \mu\text{m}^2$  (0.01 per day /  $2 \times 10^{-4}$ )
  - Cell Nucleus  $\sim 200 \mu\text{m}^2$  (1 per day/ 0.02)
  - Interacting cell matrix  $\sim 1000 \mu\text{m}^2$  (>5 per day/ >0.1)
  - Whole Tissue >Many per day
- Space radiation simulation is anchored in knowing space conditions relative to scientific question being addressed
  - A “low dose” for certain biological questions, may be a “high dose” for another: Microdosimetry
- Most ground based experiments to date have been performed in “track segment mode” where samples are irradiated with largely a single species with constant LET across sample

# Track Structure of Space Radiation



- Linear energy transfer (LET) does not adequately describe energy deposition at biomolecular or cellular scales
- Full characterization of HZE nuclei-
  - Charge,  $Z$  defines density of ionization along track ( $Z^2$  dependence)
  - Kinetic energy of  $\beta$  defines width of track corresponding to maximum distance of energy deposition laterally from track

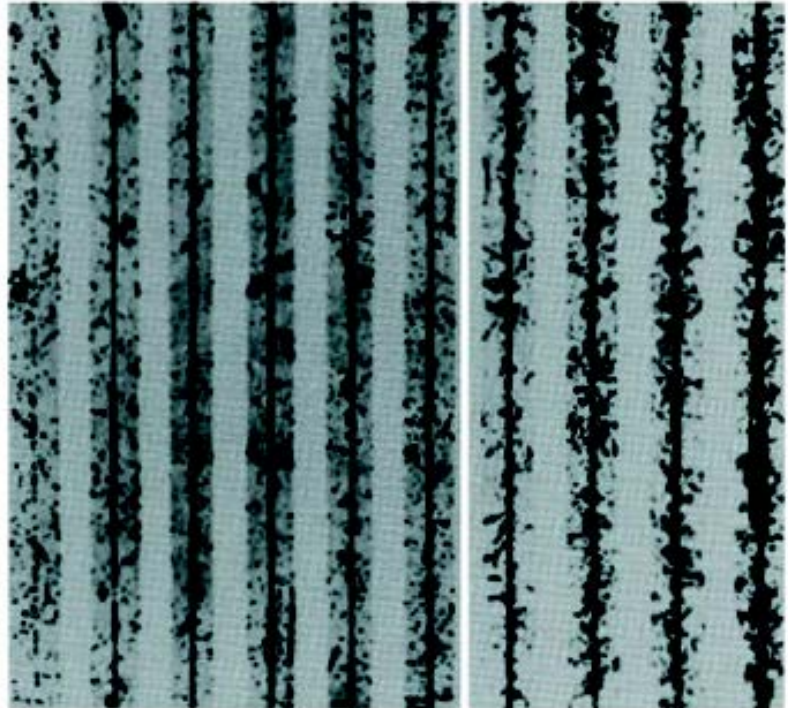
Nuclei of LET = 150 keV/ $\mu$ m



Monte-Carlo simulation of ionization patterns (He 0.45 MeV/u; C (10 MeV/u), Si (90 MeV/u) and Fe (1 GeV/u))



# Charged particle tracks in nuclear emulsions



H Z=1   He Z=2   Li Z=3   Be Z=4   B Z=5   C Z=6   Si Z=14   Cs Z=20   Ti Z=22   Fe Z=26

100  $\mu\text{m}$

(Cucinotta & Durante, Lancet Oncol 2006)

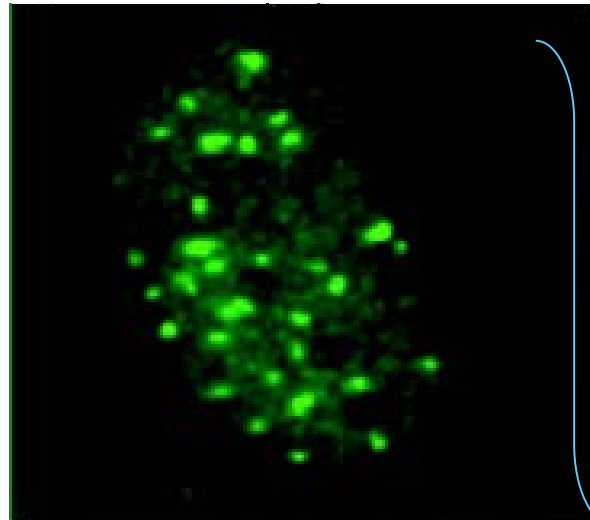
*Magnification*

# Fluorescent foci marking DSB in cell nuclei

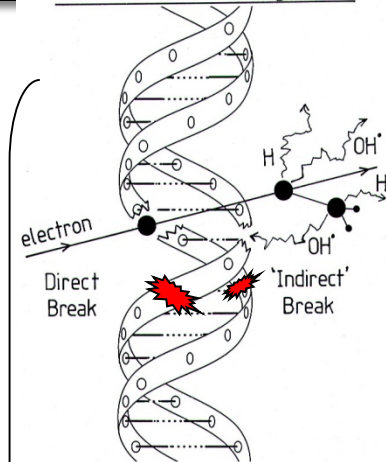
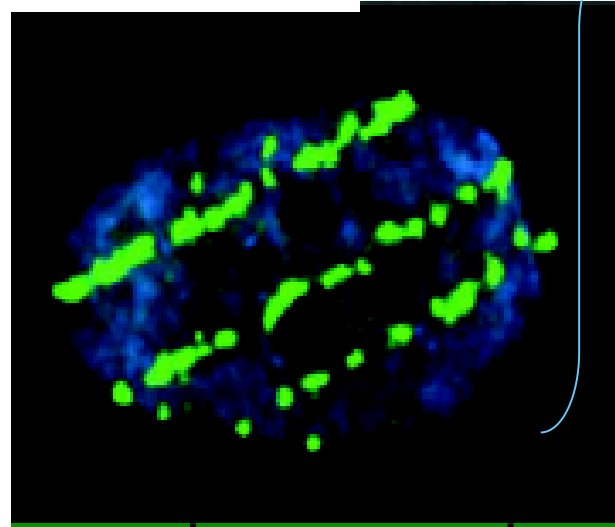


A small 'clustered damage' (simple dsb) resulting from a local cluster of ionizations within a single track:

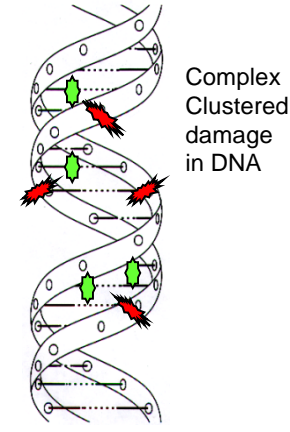
Gamma-ray irradiation:



Fe ion irradiation:



Radical diffusion distances in cells are very small (< 4 nanometres).



2 nm

X 20                      10  $\mu\text{m}$                       X 1600

# Understanding Radiation Risk



Current radiation health effects assume:

- 1) the primary mode of action is linearly related to dose and
- 2) that the individual cell is the unit of risk.

Non-targeted effects and other low dose effects suggest responses occur non-uniformly over time at the multi-cellular scale

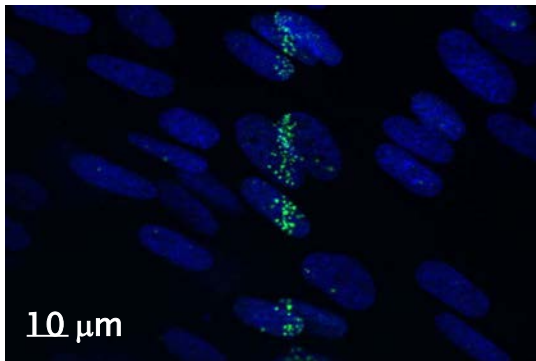
## Not all radiation is equal: RBE

### Space Radiation - High-LET

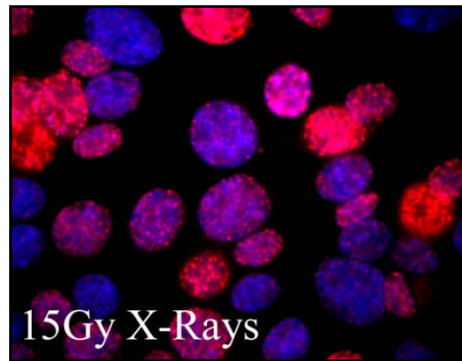
- Galactic cosmic rays (HZE)
- Solar Particle events
- Trapped radiation



Role of radiation quality and track structure?



High-LET



Low-LET

# Dose Equivalents



	Dose Equivalent (mSv)
Chest x-ray	0.1
USA annual background	4
<b>Public annual limit</b> (above background)	<b>1</b>
International Airline crews	4
<b>Radiation worker annual limit</b>	<b>20</b>
No observed effects (Abomb)(instant)	200
Blood count changes	>200
Acute Radiation Syndrome (ASR)	1,000
Death (instantaneous dose)	3,000
Mir, ISS (with shield) annual	150
Large solar flare (free space)	10,000

ICRP cancer risk estimate: 5% per Gy ~ 5% per Sv (for Q=1)

1 in 20,000 risk of fatal cancer per 1mSv dose (lifetime)



# Categories of Space Radiation Risks

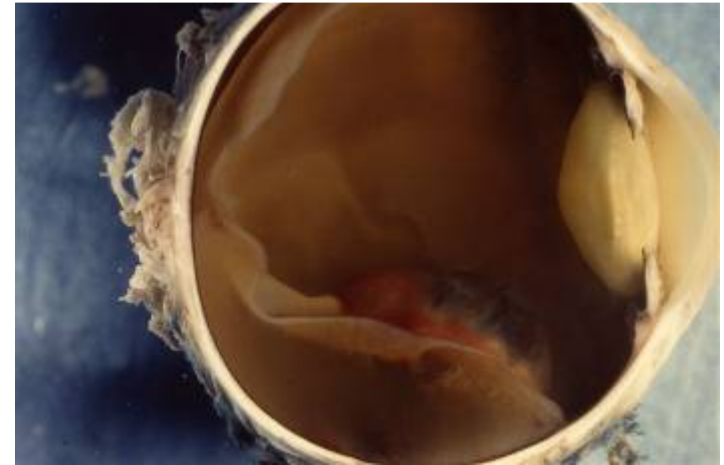


## Four categories of risk of concern to NASA:

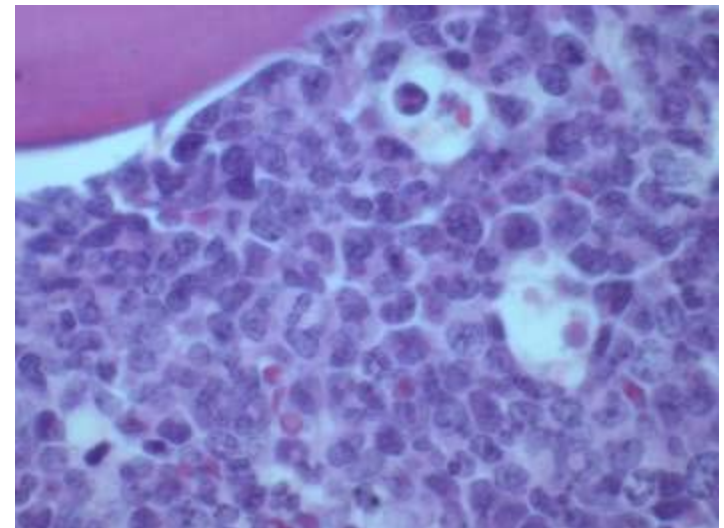
- ***Carcinogenesis (morbidity and mortality risk)***
- ***Acute and Late Central Nervous System (CNS) risks***
  - ✓ immediate or late functional changes
- ***Chronic & Degenerative Tissue Risks***
  - ✓ cataracts, heart-disease, etc.
- ***Acute Radiation Risks*** – sickness or death

## Differences in biological damage of heavy nuclei in space with x-rays, limits Earth-based data on health effects for space applications

- **New knowledge on risks must be obtained**



*Lens changes in cataracts*

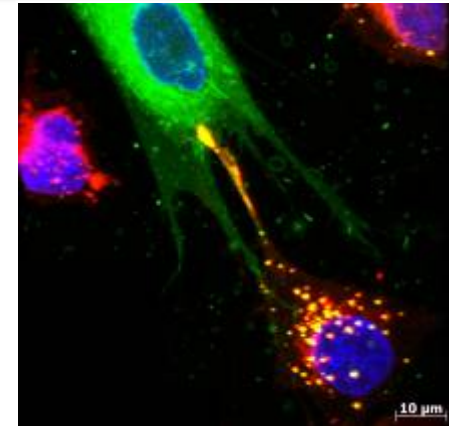


*First experiments for leukemia induction with GCR*

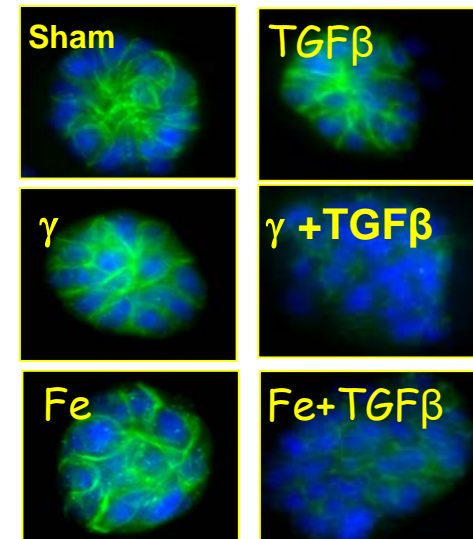
# Space Safety Requirements



- 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
  - prevent radiation sickness during mission
  - new exploration requirements limit brain 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*



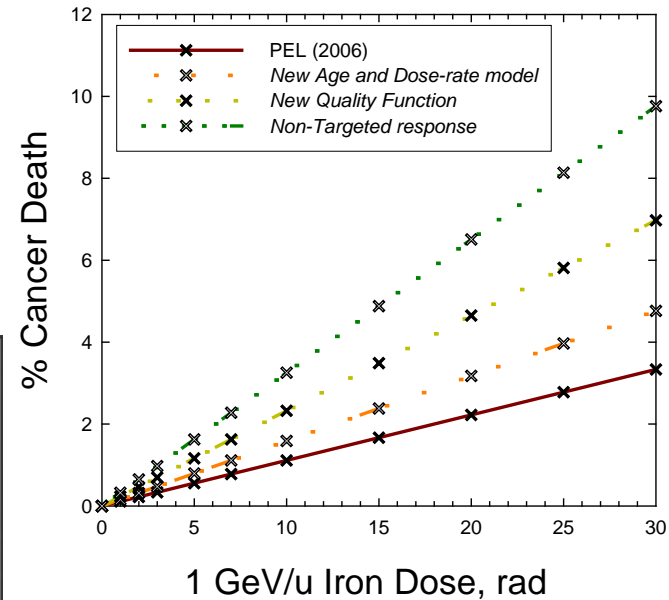
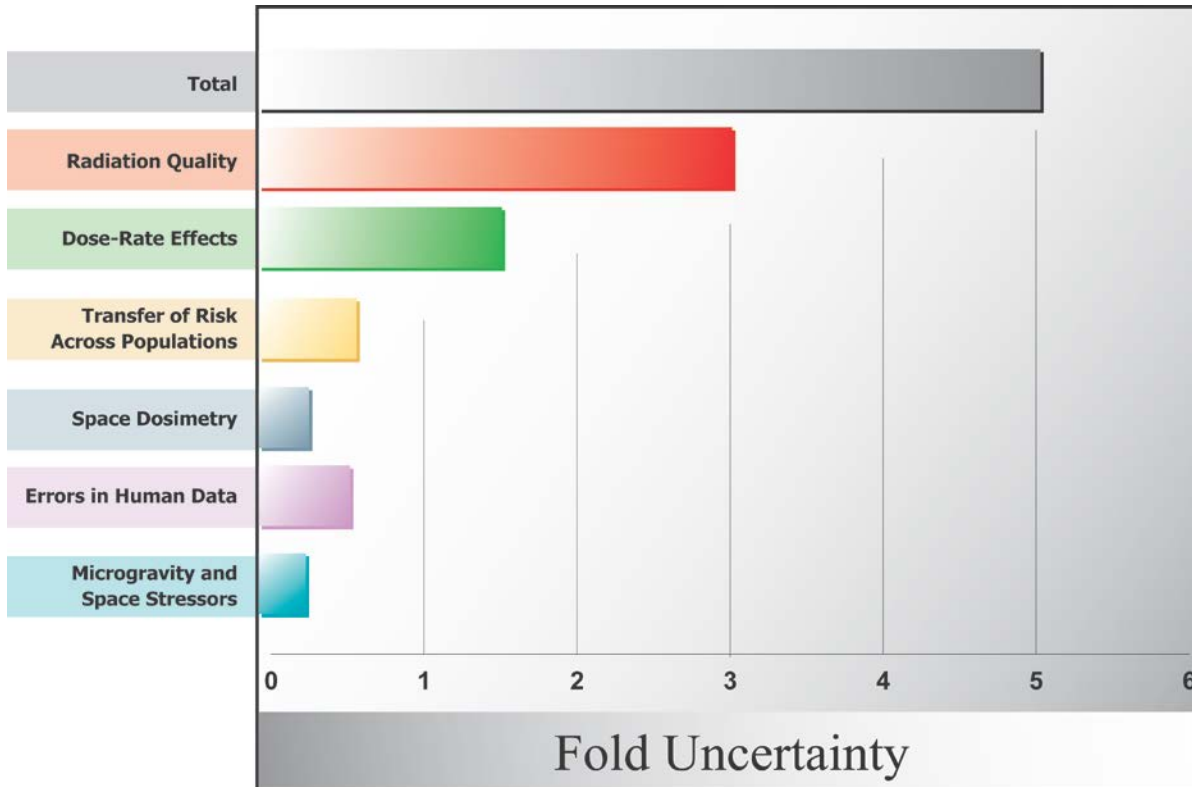
*Space Radiation in breast cancer formation*



# Cancer Uncertainty Assessments



- Uncertainties assessments use probability distribution functions (PDF) to represent range of data of different factors in risk model
- NASA is Unique amongst government agencies in that Uncertainty assessments are part of radiation protection program



Risk per unit Dose from HZE nuclei will potentially be assessed higher in near future:  
 - Age, RBE, dose-rate, and latency for solid tumors

# Estimates of “Safe Days” in Space for GCR



Research progress shows significant increase in “Safe Days” to be within acceptable risks for a Mars mission

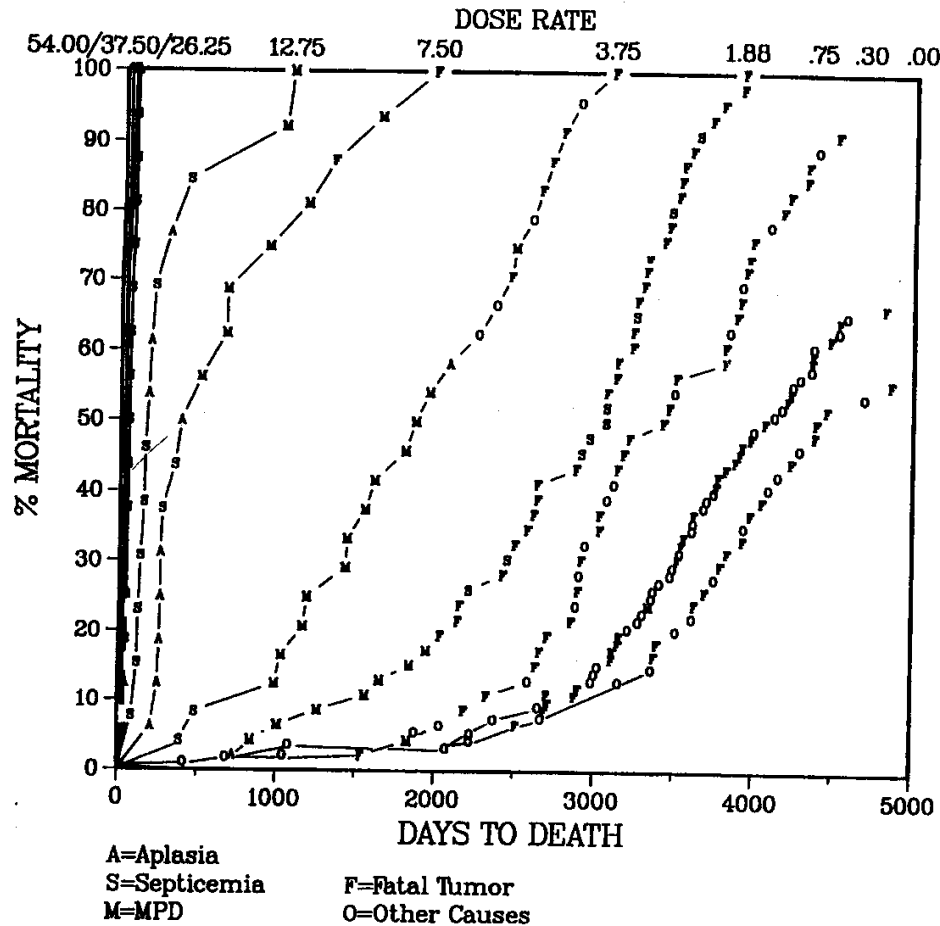
- Uncertainties are being reduced through ground based research

**Table: Estimates of “Safe Days” in deep space under heavy shielding where NASA limit is not exceeded**

Age, yr	Females		Males	
	Previous (2001)	Current (2006)	Previous (2001)	Current (2006)
30	54	112	91	142
35	62	132	104	166
40	73	150	122	186
45	89	182	148	224
50	115	224	191	273

Note: missing reference for this table

# Dose Rate effects



**Figure 2.** Relationship between dose rate, time to death and causes of death in adult beagles continuously irradiated (22 h day<sup>-1</sup>) at several dose rates (rads day<sup>-1</sup>; average absorbed dose).

# Conclusions and Questions?



- Relative to low LET exposures, we know comparatively little about the ultimate health risks of HZE and SPE exposures.
  - Relevant doses and dose rates
  - Protracted exposures
  - Combined exposures (microgravity)
- How does the heterogeneous nature of the energy distribution effect the response? Can microdosimetry predict response?
- Much of the past emphasize was on cancer risk but at low dose are other risk more important?
  - CNS – low dose (0.1 cGy) induced cognitive impairment
  - Cardio vascular risk, visual impairments
- Does microgravity act synergistically, antogonistically or they are uncoupled?
- Role of ground, LEO, and BLEO, the integration of large data sets and systems approaches in defining mechanisms of response.

# Acknowledgments



Many of the slides presented were shared by colleagues for teaching purposes. Please do not distribute without permission.

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# Low dose exposure situations



Environmental clean-up – Hanford (>\$110 Billion to date)

Nuclear Accidents – Fukushima (160,000 evacuated, 20 mGy/yr)

Rad worker exposures

Flight Crews and Astronauts (limits to the Mars mission?)

Potential Terrorist Attacks (dirty bombs, IND)

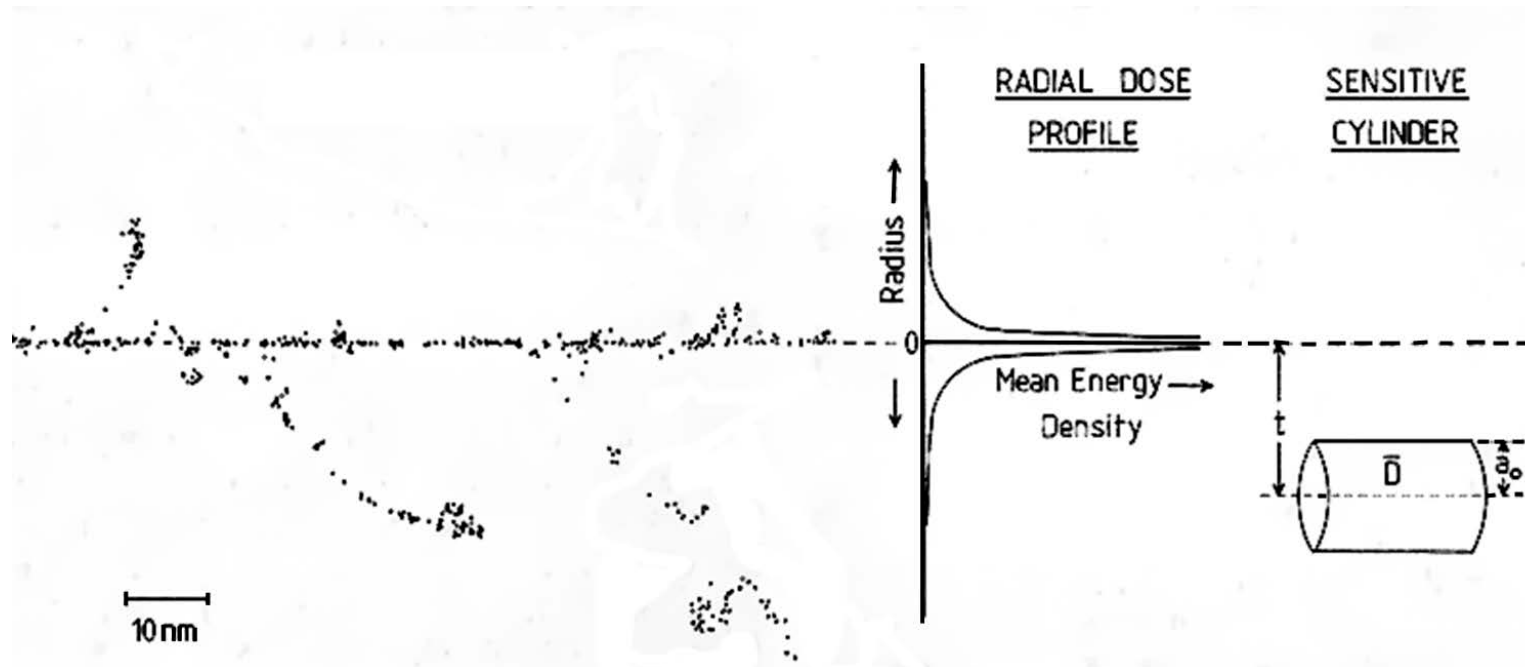
Security issues (airport backscatter machines)

High natural background exposures – Radon, geographical locations in Karala (India) Yanjing (China)

Medical Diagnostics – >90 million CT scans annually 5-100 mSv each (acute exposure v protracted exposure LDRt)

# Radial Dose Distribution of a Track

An Average the spectrum of ionization patterns as function of distance from a Track to a molecule



Ionization patterns



Radial dose from Ion deposited in Small cylinder