Miniaturized BeO-OSL detectors for dosimetry in cell cultures and organisms: Applicability and challenges

Diploma thesis Michael Anders, Radiation Physics Group TU Dresden, October 2009



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Advantages of small dosimeters:

- allow dose measurements at one defined point, also in very inhomogenous environments
- smaller perturbation of the radiation field (and of the dose distribution)
- match the typical size of cell cultures and small organisms ( $\leq 1 \text{ mm}$ )



#### But:

- small detectors are difficult to handle and probably also to find after application in an organism
- detectors have to remain inert in biological environments
- new questions about the analyzation method (a very low signal level is expected)



Almost impossible to get a dose value directly from biological tissue (e.g. by analyzing damaged proteins, chemical changes or increased temperature).

→ Idea: Replace local tissue with a material, where an easy measurable value is proportional to the applied dose

Examples: - increasing optical density of a film

- light emitted by scintillating materials
- luminescence light of materials like LiF, Al<sub>2</sub>O<sub>3</sub>, BeO
- electric current in an ionisation chamber

But: Probe material has different interaction with radiation field because of different interaction coefficients

 $\rightarrow$  leads to different dose values!





Possible solution: Theory of ideal probes – avoiding or keeping the dose gradient outside the probe

Equilibrium probes: - shouldn't affect the primary radiation field noticeably (have to be small!)

- require a secondary electron equilibrium within the whole probe volume
- only the field of primary photons determine the energy dose

Equilibrium probes are not usable for electron radiation, but are very good photon dose detectors.

How to provide a secondary electron equilibrium within the whole probe volume?

- Find an equivalent probe wall material!





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How to calculate the energy dose value in the original material?

Same radiation field – same fluence, different interaction coefficients lead to

$$D_{\rm B} = \frac{\left(\overline{\mu}_{\rm tr} / \rho\right)_{\rm B}}{\left(\overline{\mu}_{\rm tr} / \rho\right)_{\rm A}} D_{\rm A}$$

 $(D_A \text{ is the probe dose value})$ 

Similar approach for electron radiation (Bragg-Gray-Theory)

- thin probes to avoid affection of primary electron field
- very thin wall to keep secondary generation electrons out
- $\rightarrow$  Probe is a pure electron dose detector, then:

$$D_{\mathrm{B}} = rac{\left(\overline{S}/
ho
ight)_{\mathrm{B}}^{\mathrm{col}}}{\left(\overline{S}/
ho
ight)_{\mathrm{A}}^{\mathrm{col}}} D_{\mathrm{A}}$$

to be tested: miniaturized BeO-detectors (1 mm height and diameter)

- sintered from BeO powder
- density about 2.85 g/cm<sup>3</sup>
- very stable and hard (9 at Mohs scale)
- insoluble in water
- melting point about 2530 °C
- electric isolator
- effective atomic number of 7.12
- have property of optical stimulated luminescence

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Need for specific calibration in some radiation fields because detectors won't be ideal probes!

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Why can BeO be used as dosimetric detector material?

- property of optically stimulated luminescence (OSL)
   with a dose proportional measurement effect
- easy measurable value (UV light emission)
- material is reusable many times after deleting the energy dose
- effective atomic number of 7.12 is similar to effective atomic numbers of biological materials
   muscle: 7.64 water: 7.51 adipose: 6.46 air: 7.78



Optically stimulated luminescence:

Lattice defects and impurities produce additional energy levels, some of them with very long electron storage periods.



#### How to collect the emitted light?

BeOmax

BeO*max* system, developed at the Radiation Physics Group



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Problem: background signal – sources:

- changing preamplifier offset
- preamplifier noise
- background radiation
- $\rightarrow$  leads to a lowest measurable dose of 0.4 mGy (100 measurements of the background effect)

Problem: every BeO detector has its own sensitivity  $\rightarrow$  every detector has to be calibrated individually

Problem: short-time fading after irradiation (10% within 30 minutes)

→ detectors have to remain in a dark environment between irradiation and measurement



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Detectors have to remain inert in:

- water and aqueous solutions
- acids and bases
- organic solvents
- bodily liquids
- culture mediums used for cell growth

Results after treatment of detectors with a lot of substances: Detectors are inert, even in hot concentrated acids (except hydroflouric acid) and don't change their sensitivity.

Test for applicability in radioactive solutions:



#### Irradiation of detectors in 90 ml of a solution containing 1 GBq of <sup>90</sup>Y



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Increasing detector dose because of detector surface contamination!



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## 1. Decreasing detector size

BeO powder is highly cancerogen when inhalated!

→ Grinding of the detectors to reduce their size must be done under safety precautions!

Partner: Karlsruhe Beryllium Handling Facility



## Effects of modifying the dosimetry system



**Result:** 





- reduced height of about 57%
- slightly increased lowest measurable dose (0.6 mGy)
- very difficult to handle

## 2. Changing surrounding material during measurement

When using aluminium, only the bottom of the detector is stimulated  $\rightarrow$  An opaque material allows additional stimulation of the detector side



Aluminium and Optisol, an opaque Teflon-based polymer

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## Effects of modifying the dosimetry system

- Result: increased sensitivity (about 50%)
  - decreased lowest measurable dose (below 0.3 mGy)
  - improved measurement repeatability



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#### Further results and outlook





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### Further results and outlook

#### Dose characteristic: beta irradiation (<sup>90</sup>Y)









- Done: characterisation of the dosimetry system consisting of miniaturised BeO-detectors and the BeO*max* device
  - detectors can be used in biological and radioactive environments
  - knowledge about application and reusability
  - link to dosimetry theory to avoid wrong dose values
- To do: using a photon counter instead of a photo multiplier tube to improve sensitivity and to decrease the lowest measurable dose
  - improved stimulation unit
  - effects of an even further miniaturisation?
  - research on use of the detectors in bright environments

### Thank you for your attention!





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aboratory Underground Nuclear Astrophysics

# The LUNA experiment at the Gran Sasso Laboratory in Italy

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# Nuclear Astrophysics research at LNGS

LUNA...

- ... is a collaboration of several institutions in Italy, Germany and Hungary (about 25 people)
- ... is the world's only deep underground experiment using an accelerator
- ... has successfully measured data of important reactions of primordial and nucleosynthesis in stars
- ... uses now a 400 kV linear ion accelerator

"My" experiment is the  ${}_{1}^{2}H + {}_{2}^{4}He \rightarrow {}_{3}^{6}Li + \gamma$  reaction...



# The challenges of Nuclear Astrophysics

...at energies below the Coulomb barrier, so with very low cross sections. Measuring them is sometimes like...





## The LUNA 400 kV accelerator





# The shielded gas target setup



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# Thank you again for your attention!