

Inactivation of Human Cells by Heavy-ion Beams and Microdosimetric Estimates

Y. Kase (1), T. Kanai (2), Y. Matsumoto (2), Y. Furusawa (2), M. Sakama (1) and T. Asaba(1)

(1) Tokyo Institute of Technology, Japan (2) National Institute of Radiological Sciences, Japan (y_kase@nirs.go.jp)

In the field of heavy-ion therapy and space radiation protection, the prediction of biological effects of heavy charged particles is an important problem. In microdosimetry, tissue-equivalent proportional counter (TEPC) has been often used to estimate the biological effect of ionizing radiations. We measured the radiation quality of various ion beams using a spherical TEPC, while the inactivation of in vitro human cells was investigated by colony assay.

The ion beams were provided by the Heavy-ion Medical Accelerator (HIMAC) in Chiba, NIRS, Japan. The microdosimetric spectra of photon and proton-, helium-, carbon-, neon-, silicon- and iron-ions (LET =range from 0.5 to 880 keV/micron) were measured using a spherical walled TEPC with a tissue-equivalent diameter of 1 micron at various depths in a plastic phantom. In the measurement of cell surviving curves, human salivary gland (HSG) tumor cells were used as a human tumor cell line, and GM05389 cells were used as a normal human fibroblast cell line.

The linear terms of linear quadratic model with a fixed quadratic term for surviving curves of each cell types were plotted as a function of the dose-mean lineal energy measured by the TEPC. We found that these plots agreed with the microdosimetric kinetic (MK) model [R. B. Hawkins, 1994] for ion beams with a LET value of less than 450 keV/micron as long as the saturation-corrected dose-mean lineal energy was used instead of dose-mean lineal energy with non-Poisson correction in the MK model. Therefore, in heavy ion therapy, this microdosimetric estimate of the revised MK model can help determining a clinical dose in the physical definition.