METABOLISM AND GENETIC ACTION OF BIOGENIC TRITIUM IN MAMMALS

M.I. Balonov, T.V. Zhesko, M.E. Chetchueva, M.D. Pomerantseva and L.K. Ramaya Institute of Radiation Hygiene 8 Mira ul., Leningrad 197101, U.S.S.R

ABSTRACT

In the process of biosynthesis the general metabolic behaviour of H-labeled precursors of biopolymers consists in stable binding of 5-80% of tritium activity by mammalian tissues and simultaneous chipping of 90-10% of the label and its incorporating as THO into the body liquids. When injected into the body of mice and rats, H-glucose and H-amino acids will cause the tissue dose exceeding that from the equal quantity of THO by 1.1 - 10 times. The nonreplaceable amino acid L-lisin-H will cause the greatest dose and within the space of 3-5 months increase the frequency of dominant lethal mutations

(DLM) in male mice by 2-4 times as compared with THO.

Th-desoxinucleosides are incorporated into chromosomes during S-phase of the cell cycle for about an hour. The retention function of the TdR and the CdR depends on their dosage owing to reparation of DNA damages and labeled cells destruction. The injection of activity in several fractions in 10-15 hours conduces to prolonged DNA-bound tritium retention in tissues and increased radiobiological effect than does a single injection of the same dosage. Dependence of DLM and reciprocal translocations frequency in spermatogonia on the dosage of the TdR is non-linear. The effect per unit activity administered is the greatest at 37 kBq/g (1 µCi/g) and reduces by 2-4 times or by one order of magnitude with increasing dosage up to 0.37 MBq/g and 1.11 MBq/g respectively. By the genetic tests in male mice injection of 37 kBq/g of the TdR and of the CdR is equivalent to 0.2 - 0.4 and 0.05 - 0.01 Gy of gamma-radiation respectively.

The materials of the investigation have been used for

developing radiation standards.