A Comparison of Uptake of 67Ga-Citrate and 37Co-Bleomycin in Tumor Using a Semiconductor Detector

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The purpose of this comminication is to report the comparison of uptake in tumorbearing mice and clinical cases of ⁶⁷Ga-citrate and ⁵⁷Co-bleomycin. The method used in this study is a new method using a Ge (Li) semiconductor detector. This method made it possible to dectct ⁶⁷Ga and ⁵⁷Co simultaneously and to estimate their relative uptakes, and thus minimize the differences due to individual variations in experimental animals.

The tumor uptake of ⁵⁷Co-bleomycin reached a maximim at 6 hours after injection, and then the activity decreased rapidly. The ⁶⁷Gacitrate reached a maximum at 24 hours after injection and then the activity gradually decreased.

The accumulation of ⁵⁷Co-bleomycin in bone, muscle, intestine, liver, and kidney was always lower than that of ⁶⁷Ga-citrate.

From this results, it was postulated that ⁵⁷Co-bleomycin was more specific than ⁶⁷Ga-citrate for the detection of tumor.

The acculmulation of 67Ga-citrate was

always slight more than that of ⁵⁷Co-bleom-vcin.

Clinically, the scintigram obtained with ⁵⁷Co-bleomycin is superior to that of ⁶⁷Ga-citrate, because the uptakes of ⁵⁷Co-bleomycin in liver and bone was lower than that of ⁶⁷Ga-citrate.

The accumulation of ⁶⁷Ga-citrate in metastatic lymph node of patients with lung, breast cancer and reticulosarcoma was 4—20 times higher than that of ⁵⁷Co-bleomycin.

Especially, the accumulation of ⁶⁷Ga-citrate in reticulosarcoma was remarkably higher than that of ⁵⁷Co-bleomycin. The accumulation of ⁶⁷Ga-citrate in normal liver was 24 times higher than that of ⁵⁷Co-bleomycin. These results were quite similar that of animal experiments.

After treatment of radiation therapy of ⁶⁰Co (7,000 R), the accumulation of ⁶⁷Ga-citrate in lung cancer strongly decreased, whereas that of ⁵⁷Co-bleomycin not so much decreased.