

²⁰³Pb FOR BONE SCANNING

In their Letter to the Editor, Rao and Goodwin (1) reported the potential applicability of ²⁰³Pb radionuclide for skeletal imaging. This radionuclide was investigated in many centers in the United States and the radiation dosimetry based on biologic distribution in animals was highly discouraging for the use of ²⁰³Pb acetate. The critical organs for ²⁰³Pb acetate are the RES (reticuloendothelial system) organs, particularly the spleen. For example, the authors reported the image of a rabbit taken 3 days after intravenous administration of 800 μCi of ²⁰³Pb-acetate. Usually a rabbit weighs about 5 kg. Even assuming a maximum weight of 7 kg, the dose administered was 800 μCi. The corresponding tracer

dose to a 70-kg human would be 8 mCi. Scanning may have to be done 3 days after administration of ²⁰³Pb in order to obtain a high target-to-nontarget ratio and for which the tracer dose administered will be 8 mCi. According to the authors' own calculations the whole body and skeleton would receive an estimated radiation dose of 0.03 and 0.06 rads, respectively, from 100 μCi of ²⁰³Pb. For an 8 mCi dose the estimated radiation dose to the whole body would be 2.4 rads and the skeleton receives 4.8 rads.

Using the nuclear parameters of ²⁰³Pb presented in Table 1, radiation dose estimates to several organs were made. The whole-body dose was 3.09 rads/8 mCi. Assuming a 50% deposition in the skeleton, the skeleton receives 6.74 rads/8 mCi administered dose. Assuming 8% of the injected dose (8 mCi) goes to the liver (2), the radiation dose to the liver was estimated to be 6.0 rads. However, when a 70-kg patient receives an 8 mCi dose, the radiation dose to the spleen is high (29.2 rads) even assuming only 5% of the lead acetate goes to this organ.

In the case of a rare-earth metal the activity in the liver, spleen, and kidneys is reduced by approximately 50% with HEDTA chelate compared with a weak chelate such as citrate (3). As the radiation dose to the RES organs, particularly spleen, is more than 29 rads with ²⁰³Pb acetate, the use of ²⁰³Pb HEDTA will not result in a relatively low radiation dose to the patient.

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TABLE 1. NUCLEAR PARAMETERS OF ²⁰³Pb

Radiation (i)	Mean number/disint. (η _i)	Mean energy (MeV) (E _i)	Δ _i (gm-rad) (μCi-hr)
Electron capture	—	—	*
Gamma—1	0.8077	0.2792	0.4803
K int. con. electrons	0.1316	0.1936	0.0543
L int. con. electrons	0.0397	0.2649	0.0224
M int. con. electrons	0.0124	0.2762	0.0073
Gamma—2	0.0382	0.4013	0.0326
K int. con. electrons	0.0055	0.3158	0.0037
L int. con. electrons	0.0009	0.3870	0.0007
M int. con. electrons	0.0003	0.3984	0.0003
Gamma—3	0.0084	0.6805	0.0122
X-rays: Kα	0.6859	0.0719	0.1053
Kβ	0.1993	0.0840	0.0354
Lα	0.1787	0.0103	0.0039
Lβ	0.1675	0.0122	0.0044
Lγ	0.0225	0.0143	0.0007
Auger			
Electrons: KLL	0.0221	0.0570	0.0027
KLX	0.0127	0.0683	0.0018
KXY	0.0020	0.0796	0.0003
LMM	0.5842	0.0084	0.0104
MXY	1.7681	0.0029	0.0109

* For reference only—does not contribute to dose.

REFERENCES

1. RAO DV, GOODWIN PN: ²⁰³Pb: A potential radionuclide for skeletal imaging. *J Nucl Med* 14: 872, 1973
2. SPIERS FW: *Radioisotopes in the Human Body*, New York, Academic Press Inc, 1968, p 67
3. ROSOFF B, SIEGEL E, WILLIAMS GL, et al: Distribution and excretion of radioactive rare-earth compounds in mice. *Int J Appl Radiat Isot* 14: 129-135, 1963

THE AUTHORS' REPLY

We appreciate the interest of Syed regarding ²⁰³Pb dosimetry. First of all, when we injected 800 μCi in the rabbit to obtain a quick image, we did not mean

that a proportionate dose should be given to a patient. We feel that 2 mCi or less would be sufficient to obtain a good scan on a normal patient.