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## Ionic interaction of [<sup>11</sup>C]- $N, \alpha$ -dimethylbenzylamine (DMBA) in rodent brain

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The [S] enantiomer of  $[^{11}C]$ -*N*, $\alpha$ -dimethylbenzylamine (DMBA) was synthesized by N-methylation of [S]- $\alpha$ -methylbenzylamine, and its biodistribution in mice was measured.  $[^{11}C]$ -[S]-DMBA was rapidly distributed into the brain, heart and lungs, and considerable long-term retention in the brain was observed. The radioactive metabolizes in the plasma were analyzed by liquid chromatography. Kinetic analysis using unmetabolized  $[^{11}C]$ DMBA in the plasma as the input function was performed employing a simplified two-compartment model. The estimated distribution volumes (DV) of  $[^{11}C]$ DMBA in the brain and heart were 6.05 and 3.95, respectively. The right striatum of the rat brain was lesioned with ibotenic acid 2 weeks before the tracer experiment. Both *in vitro* and *in vivo* autoragiographic studies were performed, and revealed significant reduction of the radioactivity levels in the lesioned striatum. On the other hand, the regional cerebral blood flow, as measured by  $[^{14}C]$ iodoantipyrine, was not significantly altered in the lesioned striatum. These results indicate that the ionic binding component for DMBA exists mainly in neural cells rather than in glial cells.  $[^{11}C]$ DMBA might be a useful radiotracer for detection of neural cell loss in the brain.

Key words: dimethylbenzylamine (DMBA), brain, neuron, ibotenic acid