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| Title            | Metabolism of 24, 25-dihydrolanosterol analogs by partially purified cytochrome P-450 <sub>14</sub> DM from rat liver microsomes |
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| Sub Title        |  |
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| Publisher        | 共立薬科大学   |
| Publication year | 1989   |
| Jtitle           | 共立薬科大学研究年報 (The annual report of the Kyoritsu College of Pharmacy). No.34 (1989. ) ,p.100- 100                                   |
| JaLC DOI         |  |
| Abstract         |  |
| Notes            | 抄録   |
| Genre            | Technical Report   |
| URL              | https://koara.lib.keio.ac.jp/xoonips/modules/xoonips/detail.php?koara_id=AN00062898-00000034-0100                                |

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## Metabolism of 24,25-Dihydrolanosterol Analogs by Partially Purified Cytochrome P-450<sub>14DM</sub> from Rat Liver Microsomes\*

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園田よし子, 関川善夫, 佐藤良博

27-Nor-24,25-dihydrolanosterol (27-nor-DHL), 26,27-dinor-24,25-dihydrolanosterol (26,27-dincr-DHL), and 25,26,27-trinor-24,2S-dihydrolanosterol (25,26,27trinor-DHL), analogs of 24,25-dihydrolanosterol (DHL) which have no C-27 carbon, C-26, 27 carbons and C-25, 26, 27 carbons, were converted to the corresponding 14-demethylated products using a reconstituted monooxygenase system from rat liver microsomes which contained cytochrome P-450<sub>14DM</sub> catalyzing lanosterol 14demethylation and NADPH-cytochrome P-450 reductase in the presence of NADPH and molecular oxygen. Each metabolite showed a relative retention time (Rrt) of 0.72 with respect to each substrate in high-performance liquid chromatography (HPLC) on a reversed-phase column. Comparison of each gas chromatography-mass spectrum and Rrt value with those of the metabolite of DHL, 4,4-dimethyl-5α-cholesta-8,14-dien-3 B-ol, indicated that the metabolites could be inferred to be 27-nor-4,4-dimethyl-5  $\alpha$ -cholesta-8,14-dien-3 $\beta$ -ol, 26,27-dinor-4,4-dimethyl-5 $\alpha$ -cholesta-8,14-dien-3 $\beta$ -ol, and 25,26,27-trinor-4,4-dimethyl-5 $\alpha$ -cholesta-8,14-dien-3 $\beta$ -ol. However, 24,25,26,27tetranor- and 23,24,25,26,27-pentanor analogs of DHL and 20-iso-24,25dihydrolanosterol were not metabolized by the reconstituted enzyme system.

<sup>\*</sup> 本報告は Chem. Pharm. Bull. 37, 718 (1989) に発表.