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P11 COMPARATIVE PHARMACOKINETIC STUDY OF TWO-POSTCOITAL CONTRACEPTIVE AGENTS CONTAINING LEVONORGESTREL

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ABSTRACT

The pharmacokinetic study was carried out on levonorgestrel tablets from two different sources (Hugarian-and Thai-made). Both preparations contained 0.75 mg levonorgestrel and had been used for postcoital contraception. subjects were given a single oral dose of 0.75 mg tablets in a crossover disign. Plasma levonorgestrel concentration were determined via radio-immunoassay technique. The pharmacokinetic analysis of the levonorgestrel data was performed on the basis of a two-compartment open model uning a M.K. model programme. Absorption and bioavailability of the Hungarian-made tablets were slighly higher as evidenced by higher serum concentrations of levonorgestrel in the first 2 hours. Thaimade tablets showed a slighly greater area under the concentration-time curve (AUC) during the first 24 hours. The volume distribution (Vd), the clearance (CL) and elimination half-life $(T_{1/2})$ of both preparations were closely. Comparing the pharmacokinetic parameters (C_{max}, T_{max}, T_{1/2}, CL, Vd, AUC) by using t-test were not statistically significant difference. It implied that both preparations (Hungarian-and Thai-made) were bioequivalent with each other.