SYNTHESIS OF TRANSDERMAL PERMEATION ACCELERANTS ON

THE BASIS OF PIPERIDIN-3-KARBOXYLIC ACID

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Transdermal permation enhancers are chemical compounds which facilitate the drug

delivery through the skin. They influence the stratum corneum, as the outer layer of the

epidermis. Basic requirements for the enhancer are safety, drug compatibility,

biodegradability, and that they must be non-toxic and non-irritating.

The model structure was effective trancarbame 12, according to which have been

synthetized its cyclic analogues:

Hydrobromide of piperidine-3-carboxylic acid decylester

Hydrobromide of piperidine-3-carboxylic acid dodecylester

N-acetylderivative of piperidine-3-carboxylic acid decylester

N-acetylderivative of piperidine-3-carboxylic acid dodecylester

• 3-(decyloxycarbonyl)piperidinium-3-(decyloxycarbonyl)piperidine-1-carbamate

3-(dodecylexycarbonyl)piperidinium-3-(dodecyloxycarbonyl)piperidine-1-carbamate

Those compounds have been characterized by common spectral methods.

The transdermal permation activity has been consequently evaluated on the porcine skin

in the Franz cells, using theophylline as the model penetrating drug.

The activity has been evaluated using the HPLC method and determined by the

Microsoft Excel. The results demonstrated higher activity, both of N-acetyl derivatives of

piperidin-3-carboxylic acids, and of carbamates of those esters, then transcarbam 12 and

coumpunds, which i have synthetized, have.

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