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SYNTHESIS AND DESULFURATION OF THIANAPHTHEN-3-CARBOXYLIC ACID DERIVATIVES

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SYNTHESIS AND DESULFURATION OF THIANAPHTHEN-3-CARBOXYLIC ACID DERIVATIVES

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Recently, we have been interested in the synthesis of some heterocyclic compounds as potential antiinflammatoric agents¹. Derivatives of thianaphthen-3-carboxylic acid might also show such activity. Therefore, a rational synthetic pathway for these compounds was developed:

Last step in the synthesis i.e. benzoylation, gave two isomers in the ratio 3,5:1, total yield 36%. To prove the structure of isomers obtained, 5-benzoyl derivative was synthesized in multistep procedure starting with p-thiocresol:

The compounds thus obtained were treated with Raney nickel undergoing desulfuration and yielding derivatives of phenylpropionic acid. Isomer ratio and position of electrophilic attack on thianaphthen-3-carboxylic acid methyl ester will be discussed, as well as the products obtained under different desulfuration conditions.

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