Transformations of Steroid Esters by Fusarium culmorum

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The course of transformations of the pharmacological steroids: testosterone propionate, 4-chlorotestosterone acetate, 17β -estradiol diacetate and their parent alcohols in *Fusarium culmorum* AM282 culture was compared. The results show that this microorganism is capable

estrone – the main metabolite of estradiol – was absent in the reaction mixture. The alcohols resulting from the hydrolysis underwent oxidation at C-17 and hydroxylation. The same products (6β - and 15α -hydroxy derivatives) as from testosterone were formed by transformation of testosterone propionate, but the quantitative composition of the mixtures obtained after transformations of both substrates showed differences. The 15α -hydroxy deriv-

The presence of the chlorine atom at C-4 markedly reduced 17β -saponification in 4-chlorotestosterone acetate. Only 3β , 15α -dihydroxy- 4α -chloro- 5α -androstan-17-one (the main product of transformation of 4-chlorotestosterone) was identified in the reaction mixture. 6β -Hydroxy-4-chloroandrostenedione, which was formed from 4-chlorotestosterone, was not detected in the extract obtained after conversion of its ester.

atives were obtained from the ester in considerably higher yield than from the parent alcohol.

of regioselective hydrolysis of ester bonds. Only 4-ene-3-oxo steroid esters were hydrolyzed at C-17. 17β -Estradiol diacetate underwent regioselective hydrolysis at C-3 and as a result,

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