

Abstract

The work is focused on the synthesis of deuterium-labelled neuroactive steroids useful as inner standards for determination of their pharmacokinetics and bioavailability. The starting material for the syntheses was commercial 11α -hydroxyprogesterone **16**. The target compound, 20-Oxo-[9,12,12- $^2\text{H}_3$]5 β -pregnan-3 α -yl L-glutamyl 1-ester (**47**), contains 3 deuterium atoms in positions 9α , 12α , 12β .

Alternative target with a 18-functionalized group (**34**) was also studied. It will be used for an analogue with deuterium atoms on carbon C-18.