DESIGN OF NOVEL THERAPEUTIC STRATEGIES FOR ALCOHOL ADDICTION

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ABSTRACT

Background: Baclofen® is an FDA-approved GABAB agonist, used for the treatment of spasticity since the early seventies.

Objective: The aim of this study was to synthesize a new analog of BF.

Methods: The main problem of baclofen® is that it is administered as a racemate. However, in our work we synthesize the R-(−) enantiomer. In our study, we used a stereoselective method for the synthesis of peptide analog and the study of their inclusion complex with CD, at the University of Tlemcen, Algeria in 2017.

Results: For evaluation of anti-addiction activity, we have developed a new model in vivo, and the results are that our synthetized baclofen® analogues showed an anxiolytic effect. Regarding the toxicity, our results showed that our analogues have less toxic effect than baclofen; it reduces the activity of TGO enzymes, and TGP. From a histological point of view, it has no effect on the liver structure in addition to having a protective effect against lesions liverworts induced by alcohol.

Conclusion: Our conclusion begs an open question: If baclofen® can cure any form of addiction such as smoking, alcoholism and other addictions to addictive substances, and if its beneficial effect is already proven, approved, and used as an antircraving agent in several countries in the world, why until now has the Algerian health community not used baclofen® in the treatment of numerous addictions?

KEYWORDS: Alcohol addiction, Baclofen®

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