## Design, Synthesis and Biological Evaluation of Triptorelin Analogs Containing

## **Tetrazole Moiety**

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## Abstract

The design and synthesis of novel bioisosteric analogues of triptorelin acetate containing tetrazole moiety at Cor N-terminus of the peptides is described. Tetrazole acetic acid was synthesized through the reaction of an alkyl cyanoacetate and sodium azide, after that, it was coupled to the N-terminus of triptorelin instead of pyroglutamic acid. In another approach, tetrazole acetohydrazide was prepared and conjugated to the Cterminus of triptorelin instead of the amide bond. The synthesized peptides containing tetrazole moiety at the C- or N-terminus of the peptide sequence are peptidomimetics of triptorelin acetate. The docking results of the designed derivatives showed the same interaction of triptorelin with the receptor but with a higher score. The peptide that has tetrazole moiety in its C-terminus, restricted the testosterone level during four weeks of *in vivo* study and led to keep testosterone at a moderate level during the first week compared to control