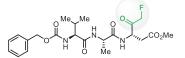
α-Fluoroketones for Drug Discovery

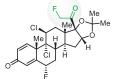
Introduction

 α -Fluoroketones, particularly α -fluoromethyl ketones (FMK), act as a robust and selective chemical handle for irreversible binding of bioactive molecules to cysteine residues in target proteins. FMK molecules are present in various compound classes, including several commercial peptidic caspase inhibitors related to Z-VAD-FMK, halosteroid trainide, and more. Additionally, α -fluoroketones are frequently utilized in drug discovery projects as advanced ketone building blocks suitable for Claisen-Schmidt condensation and reductive amination, that produce fluorinated bioactive molecules with enhanced properties. Try our α -fluoroketones in our next project!

FMK inhibitor of p90 ribosomal protein S6 kinase Nat. Chem. Biol. 2007, 3, 156



Z-VAD(OMe)-FMK Caspase inhibitor J. Exp. Med. 1996, 184, 2445



Tralonidesynthetic glucocorticoid corticosteroid
Syntex Corp

Reactions

Reductive amination

J. Med. Chem. 2020, 63, 12574 anticoagulant

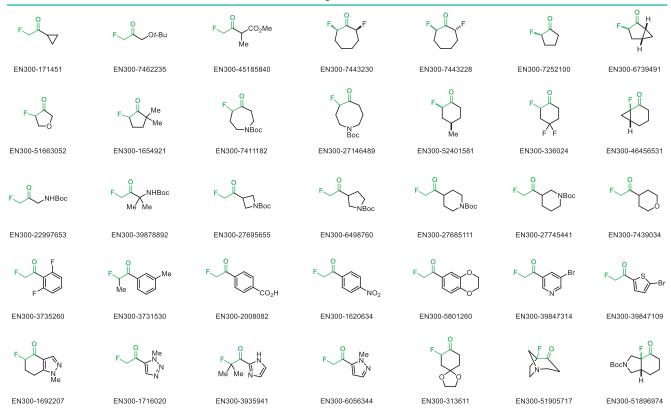
Claisen-Schmidt condensation

Eur. J. Med. Chem. 2020, 193, 112216 against chronic inflammation

NO inhibition rate 70% at 10 μM pre-treatment in RAW 264.7 cells

We offer: over 100 α -fluoroketones from stock on 5-10 gram scale.

BAY1217224



References

M. Cohen et al. Science 2005, 308, 1318.
 S. Dhani et al. Cell Death Dis. 2021, 12, 949.

- 3. A. Hillisch et al. J. Med. Chem. 2020, 63, 12574.
- 4. Y. Yang et al. Eur. J. Med. Chem. 2020, 193, 112216.



