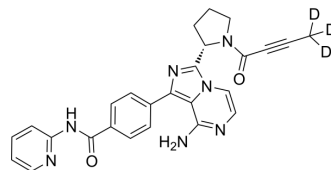


Acalabrutinib-d₃

Cat. No.:	HY-W757743
Molecular Formula:	C ₂₆ H ₂₀ D ₃ N ₇ O ₂
Molecular Weight:	468.53
Target:	Isotope-Labeled Compounds
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description

Acalabrutinib-d₃ (ACP-196-d₃) is the deuterated form of Acalabrutinib (HY-17600). Acalabrutinib (ACP-196) is an orally active, irreversible, highly selective second-generation BTK inhibitor. Acalabrutinib covalently binds to Cys481 in the ATP-binding pocket of BTK. Acalabrutinib shows strong targeting and efficacy in mouse models of chronic lymphocytic leukemia (CLL).

REFERENCES

- [1]. Wu J, et al. Acalabrutinib (ACP-196): a selective second-generation BTK inhibitor. *J Hematol Oncol.* 2016 Mar 9;9:21.
- [2]. Herman SE, et al. The Bruton's tyrosine kinase (BTK) inhibitor acalabrutinib demonstrates potent on-target effects and efficacy in two mouse models of chronic lymphocytic leukemia. *Clin Cancer Res.* 2016 Nov 30.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA