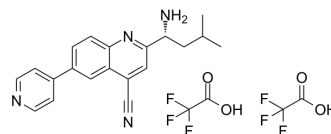


## AAK1-IN-3 TFA

Cat. No.:	HY-144302A
CAS No.:	2761587-27-1
Molecular Formula:	C <sub>24</sub> H <sub>22</sub> F <sub>6</sub> N <sub>4</sub> O <sub>4</sub>
Molecular Weight:	544.45
Target:	AAK1
Pathway:	Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	AAK1-IN-3 TFA, a quinoline analogue, is a brain-penetrant adaptor protein 2-associated kinase 1 (AAK1) inhibitor with an IC <sub>50</sub> of 11 nM. AAK1-IN-3 has the potential for neuropathic pain research <sup>[1]</sup> .
<b>In Vitro</b>	AAK1-IN-3 (compound (R)-31) TFA is brain-penetrant kinase inhibitors with a B/P ratio of 1.3 <sup>[1]</sup> . AAK1-IN-3 TFA inhibits AAK1 in HEK293 cells (IC <sub>50</sub> =108 nM) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>In Vivo</b>	AAK1-IN-3 (compound (R)-31; 30 mg/kg; sc; single dose) TFA causes a 46% reduction of μ2 phosphorylation in C57BL6 mice <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Richard A Hartz, et al. Bicyclic Heterocyclic Replacement of an Aryl Amide Leading to Potent and Kinase-Selective Adaptor Protein 2-Associated Kinase 1 Inhibitors. J Med Chem. 2022 Mar 10;65(5):4121-4155.

**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