

Product Name : CLK inhibitor T3

Synonyms : T3

Cat No. : M13352

CAS Number : 2109805-56-1

Molecular Formula : C28H30N6O2

Formula Weight : 482.59

Chemical Name : 4-(2-methyl-1-(4-methylpiperazin-1-yl)-1-oxopropan-2-yl)-N-(6-(pyridin-4-yl)imidazo[1,2-a]pyridin-2-yl)benzamide

CLK inhibitor T3 (T3) is a highly potent, selective, and cell-based stable CDC-like kinase (CLK) inhibitor with 0.67, 15 and 110nM for CLK1,2 and 3, respectively; displays 200-300-fold selectivity over other dual specificity kinases such as DYRK1A

N NH NH NH

DescriptionTronk for CLK 1,2 and 3, respectively, displays 200-300-load selectivity over other dual specificity kinases such as DYRK and DYRK1B; induces dose-dependent reduction in exon recognition and exhibits an overlapping, but greater effect on

transcriptome splicing compared to KH-CB19.

Pathway : Cell Cycle/DNA Damage

Target : CLK

Receptor : CLK

Solubility : —

SMILES : O=C(NC1=CN2C=C(C3=CC=NC=C3)C=CC2=N1)C4=CC=C(C)(C)(C)C(N5CCN(C)CC5)=O)C=C4

Storage : (-20°C)

Stability : ≥2 years

Reference :

1. Funnell T, et al. Nat Commun. 2017 Feb 23;8(1):7.