

# **GNE-6776**

# Catalog No: tcsc0031103

Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

**Size:** 100mg

**Specifications** 

CAS No:

2009273-71-4

Formula:

 $C_{20}H_{20}N_4O_2$ 

**Pathway:** Cell Cycle/DNA Damage

**Target:** 

Deubiquitinase

### Purity / Grade:

>98%

#### Solubility:

DMSO : ≥ 310 mg/mL (889.78 mM)

#### **Observed Molecular Weight:**

348.4

## **Product Description**

GNE-6776 is a selective **USP7** inhibitor.

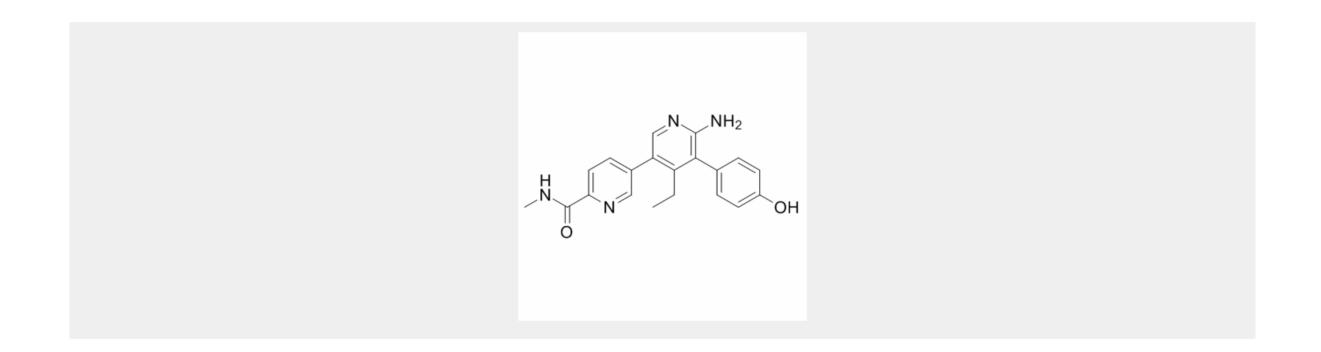
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#### IC50 & Target: USP7<sup>[1]</sup>

*In Vitro:* GNE-6776 non-covalently targets USP7 12 Å distant from the catalytic cysteine. GNE-6776 attenuates ubiquitin binding and thus inhibits USP7 deubiquitinase activity. GNE-6776 interacts with acidic residues that mediate hydrogen-bond interactions with the ubiquitin Lys48 side chain. GNE-6776 targets cellular USP7, MDM2, and p53 signalling pathways.GNE-6776 selectively inhibits recombinant USP7 relative to 36 other deubiquitinases. GNE-6776 remains selective even at 100  $\mu$ M, a more than sixfold higher concentration than used in cellular assays. GNE-6776 significantly inhibits USP7 while remaining selective against 44-47 other detected deubiquitinases<sup>[1]</sup>.

*In Vivo:* Although efficacious exposure is only transiently achieved, GNE-6776 causes modest, although significant, EOL-1 xenograft growth delay<sup>[1]</sup>.



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