

## CUDC-101

Cat. No. CEI-0982

Lot. No. (See product label)

### Introduction

#### Description

CUDC-101 is a potent inhibitor of histone deacetylase (HDAC) and the receptor kinases epidermal growth factor receptor (EGFR) and human epidermal growth factor receptor 2 (HER2), with IC<sub>50</sub> values of 4.4, 2.4, and 15.7 nM, respectively. It inhibited EGFR and Her2 phosphorylation, reduced cell proliferation and induced apoptosis in HCC827 non-small cell lung cancer (NSCLC) xenografts. CUDC-101 inhibited EGFR and induced upregulation of acetylated histone H3 in a dose-dependent fashion.

### Product Information

<b>Appearance</b>	White solid.
<b>CAS No.</b>	1012054-59-9
<b>Molecular Formula</b>	C <sub>24</sub> H <sub>26</sub> N <sub>4</sub> O <sub>4</sub>
<b>Molecular Weight</b>	434.5 Da
<b>Purity</b>	>99%
<b>Targets</b>	HDAC, EGFR, HER2
<b>IC<sub>50</sub></b>	HDAC 、 EGFR 、 HER2 : 4.4, 2.4, and 15.7 nM respectively
<b>Solubility</b>	Soluble in DMSO at 28 mg/ml; very poorly soluble in ethanol; very poorly soluble in water; maximum solubility in plain water is estimated to be about 20-50 μM; buffers, serum, or other additives may increase or decrease the aqueous solubility.

### Storage and Shipping Information

<b>Stability</b>	Store at or below -20°C. Solid form is stable at least 12 months from date of receipt, when stored as directed. Do not store aqueous solutions for more than one day.
------------------	---