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## Catalog Number: CM03077

产品信息

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CAS号: 1207293-36-4

分子式: C<sub>29</sub>H<sub>28</sub>N<sub>4</sub>O<sub>2</sub>

主要靶点:

Apoptosis|Aurora Kinase|MEK

主要通路: 凋亡|表观遗传|细胞周期|MAPK信 号通路

分子量: 464.56 溶解度:

H2O:<1 mg/mL,Ethanol:1 mg/mL (2.15 mM),DMSO:18 mg/mL (38.7 mM)

靶点活性

aurora kinases (AK):15 nM|MEK:4nM

体外活性

BI-847325 shows growth-inhibitory effects on BRAF-mutant and vemurafenib-resistant melanoma cells with IC50 ranging from 0.3 nM to 2  $\mu$  M, and prevents colony formation in six BRAF-mutant melanoma cell lines. BI-847325 also induces apoptosis by reducing Mcl-1 expression. [1]

In mice bearing 1205Lu and 1205LuR xenografts, BI-847325 (75 mg/kg, p.o.) causes significant tumor suppression without significant alteration in the body weights. [1]

细胞实验

Cells are plated at a density of 2.5  $\times$  103 cells per 100  $\,^{\,\mu}$  L and left to grow overnight before being treated with increasing concentrations of BI-847325 for 72 hours. The metabolic activity is determined using Alamar blue reagent as per the manufacturer's protocol.(Only for Reference)

BI-847325 is a selective dual inhibitor of MEK and aurora kinases (AK) with IC50 values of 4 and 15 nM for human MEK2 and

AK-C, respectively.

储存

Powder: -20°C for 3 years | In solvent: -80°C for 2 years