abcam

Product datasheet

Z-D(OMe)E(OMe)VD(OMe)-FMK, Cell permeable caspase-3 inhibitor ab120488

2 References 2 图像

概述

产品名称 Z-D(OMe)E(OMe)VD(OMe)-FMK, Cell permeable caspase-3抑制剂

描述 Cell permeable caspase-3抑制剂

生物学描述 A potent, cell-permeable, and irreversible inhibitor of caspase-3. Also inhibits caspase-6,

caspase-7, caspase-8, and caspase-10. Once in the cell, endogenous esterase activity hydrolyzes the methyl groups to form the biologically active form. Therefore, when using with isolated, purified or recombinant caspase enzymes, pre-treatment with an esterase is required.

CAS编号 210344-95-9

化学结构 Z-Asp(OMe)-Glu(OMe)-Val-Asp(OMe)-FMK

性能

分子量 668.67

分子式 C₃₀H₄₁FN₄O₁₂

序列 DEVD (Modifications: N-terminal benzyloxycarbonyl; C-terminal FMK; Asp-1 = Asp(OMe); Glu-2 =

Glu(OMe); Asp-4 = Asp(OMe))

PubChem识别号 16760394

存放说明 Store at -20°C. Store under desiccating conditions. The product can be stored for up to 12

months.

溶解度概述 Soluble in DMSO to 20 mM

处理 Wherever possible, you should prepare and use solutions on the same day. However, if you need

to make up stock solutions in advance, we recommend that you store the solution as aliquots in tightly sealed vials at -20 $^{\circ}$ C. Generally, these will be useable for up to one week. Before use, and

prior to opening the vial we recommend that you allow your product to equilibrate to room temperature for at least 1 hour.

Need more advice on solubility, usage and handling? Please visit our **frequently asked**

questions (FAQ) page for more details.

SMILES FCC(=0)[C@H](CC(=0)OC)NC(=0)[C@@H](NC(=0)[C@H](CCC(=0)OC)NC(=0)[C@H]

(CC(=O)OC)NC(=O)OCc1ccccc1)C(C)C

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应用

The Abpromise guarantee Abpromise™承诺保证使用ab120488于以下的经测试应用

"应用说明"部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

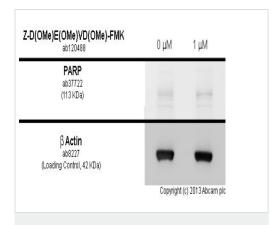
应用	Ab评论	说明
Functional Studies		Use at an assay dependent concentration.

图片



Chemical Structure - Z-D(OMe)E(OMe)VD(OMe)-FMK, Cell permeable caspase-3 inhibitor (ab120488)

2D chemical structure image of ab120488, Z-D(OMe)E(OMe)VD(OMe)-FMK, Cell permeable caspase-3 inhibitor



Functional Studies - Z-D(OMe)E(OMe)VD(OMe)-FMK, Cell permeable caspase-3 inhibitor (ab120488)

HeLa cells were incubated at 37 °C for 1h with vehicle control (0 μ M) and different concentrations of Z-D(OMe)E(OMe)VD(OMe)-FMK (ab120488). After this incubation 10 μ M of camptothecin (ab120115) was added to all samples and the cells were incubated for further 24h. Increased expression of full length PARP (ab37722) in camptothecin induced apoptotic HeLA cells correlates with an increase in Z-D(OMe)E(OMe)VD(OMe)-FMK concentration, as described in literature.

Whole cell lysates were prepared with RIPA buffer (containing protease inhibitors and sodium orthovanadate), 10 μ g of each were loaded on the gel and the WB was run under reducing conditions. After transfer the membrane was blocked for an hour using 5% BSA before being incubated with **ab37722** at 1 μ g/ml and **ab8227** at 1 μ g/ml overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP (**ab97051**

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES, NOT FOR USE IN HUMANS"

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