

DAPK

Death associated protein kinase

DAPK (Death-associated protein kinase) is the founding member of a newly classified family of Ser/Thr kinases, whose members not only possess significant homology in their catalytic domains, but also share cell death-associated functions. The realization that DAPk is a tumor suppressor gene, whose expression is lost in multiple tumor types, has spurred a flurry of interest in the kinase family and produced an impressive body of literature concerning its function, regulation, and connection to disease. The DAPk family has been linked to several cell death-related signaling pathways, and functions other than cell death have also been proposed.

DAPK Inhibitors

3MB-PP1

3MB-PP1, a bulky purine analog, is a Polo-like kinase 1 (Plk1) inhibitor. 3MB-PP1 blocks mitotic progression and cell division arise through target Plk1 in in cells expressing analog-sensitive Plk1 alleles.

Purity: >98%

Clinical Data: No Development Reported

Size: 1 mg, 5 mg

Cat. No.: HY-102069

DAPK Substrate Peptide TFA

DAPK Substrate Peptide TFA is a synthetic peptide substrate for death associated protein kinase (DAPK), with a K_m of 9 μM .

KKRPQRRYSNVF (TFA salt)

Cat. No.: HY-P1344A

Purity: 99 33%

Clinical Data: No Development Reported

Size: 5 mg, 10 mg

DRAK2-IN-1

DRAK2-IN-1, compound 16, is a potent, selective and ATP-competitive DRAK2 inhibitor with IC50 and K₂values of 3 nM and 0.26 nM, respectively. DRAK2-IN-1 also has inbitory effect on DRAK1 $(IC_{50} = 51 \text{ nM}).$

Purity: >98%

Clinical Data: No Development Reported

Size: 1 mg, 5 mg



TC-DAPK 6

Cat. No.: HY-15513

TC-DAPK 6 is a potent, ATP-competitive, and highly selective DAPK inhibitor (IC₅₀=69 and 225 nM against DAPK1 and DAPK3, respectively, with 10 μM ATP).

Purity: 95.03%

Clinical Data: No Development Reported Size: 10 mM × 1 mL, 10 mg, 50 mg

HS38

HS38 is a potent, selective, and ATP-competitive inhibitor of death-associated protein kinase 1 (DAPK1) and zipper-interacting protein kinase (ZIPK, also called DAPK3), with K_ds of 300 nM and 280 nM, respectively. HS38 is also a PIM3

inhibitor with an IC₅₀ of 200 nM.

98.01% **Purity:**

Clinical Data: No Development Reported 5 mg, 10 mg, 25 mg

Cat. No.: HY-15847

TNIK-IN-3

Cat. No.: HY-145293

TNIK-IN-3 is a potent, selective and orally active inhibitor of Traf2- and Nck-interacting protein kinase (TNIK), with an IC $_{s0}$ of 0.026 $\mu M.$ TNIK-IN-3 could also inhibit Flt4 (IC₅₀=0.030 μ M), Flt1 (IC₅₀=0.191 μ M) and DRAK1 (IC₅₀=0.411 μ M).

>98% Purity:

Clinical Data: No Development Reported

Size: 1 mg, 5 mg