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## DATASHEET

### MK-2206 dihydrochloride

#### Product overview

<b>Name</b>	MK-2206 dihydrochloride
<b>Cat No</b>	HB5934
<b>Alternative names</b>	MK-2206 2HCl
<b>Biological action</b>	Inhibitor
<b>Purity</b>	>98%
<b>Description</b>	Highly selective, pan-Akt inhibitor. Autophagy and apoptosis inducer. Orally bioavailable.

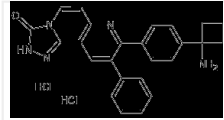
#### Biological Data

<b>Biological description</b>	Highly selective, allosteric pan-Akt inhibitor (IC <sub>50</sub> values are 8, 12 and 65 nM at Akt1, Akt2 and Akt3 respectively) which shows no inhibitory activity at 250 other protein kinases. Orally bioavailable and blood brain barrier permeable. Binds to Akt and inhibits it in a non-ATP competitive manner. Shows anti-cancer (antineoplastic) antiproliferative and anti-tumor activity. Synergistically inhibits cell proliferation of human cancer cell lines to enhance anti-tumor efficacy when used in combination with other anticancer agents. Induces cell cycle arrest, autophagy and apoptosis.
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#### Solubility & Handling

<b>Storage instructions</b>	-20 °C
<b>Solubility overview</b>	Soluble in DMSO (100 mM)
<b>Important</b>	This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use

#### Chemical Data

<b>Chemical name</b>	8-[4-(1-aminocyclobutyl)phenyl]-9-phenyl-2H-[1,2,4]triazolo[3,4-f][1,6]naphthyridin-3-one;dihydrochloride
<b>Molecular Weight</b>	480.4
<b>Chemical structure</b>	
<b>Molecular Formula</b>	C <sub>25</sub> H <sub>21</sub> N <sub>5</sub> O.2HCl
<b>CAS Number</b>	1032350-13-2
<b>PubChem identifier</b>	46930998
<b>SMILES</b>	C1CC(C1)(C2=CC=C(C=C2)C3=C(C=C4C(=N3)C=CN5C4=NNC5=O)C6=CC=CC=C6)N.C1.C1
<b>InChi</b>	InChI=1S/C25H21N5O.2ClH/c26-25(12-4-13-25)18-9-7-17(8-10-18)22-19(16-5-2-1-3-6-16)15-20-21(27-22)11-14-30-23(20)28-29-24(30)31;/h1-3,5-11,14-15H,4,12-13,26H2,(H,29,31);*2*1H
<b>InChiKey</b>	HWUHTJIKQZZBRA-UHFFFAOYSA-N

#### References

**MK-2206, an allosteric Akt inhibitor, enhances antitumor efficacy by standard chemotherapeutic agents or molecular targeted drugs in vitro and in vivo.**

Hirai H et al (2010) Molecular cancer therapeutics 9

**PubMedID**

20571069

**eEF-2 kinase dictates cross-talk between autophagy and apoptosis induced by Akt Inhibition, thereby modulating cytotoxicity of novel Akt inhibitor MK-2206.**

Cheng Y et al (2011) Cancer research 71

**PubMedID**

21307130

**Abstract #DDT01-1: MK-2206: A potent oral allosteric AKT inhibitor**

Yan Li (2009) New Drugs on the Horizon 1 69

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