

Product Name: KN-92 hydrochloride Revision Date: 01/10/2021

Product Data Sheet

HCI

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KN-92 hydrochloride

Cat. No.:	A3530		
CAS No.:	1431698-47-3		
Formula:	C24H26Cl2N2O3S		
M.Wt:	493.45		
Synonyms:	KN 92 hydrochloride;KN92 hydrochloride		
Target:	Membrane Transporter/Ion Channel		
Pathway:	P2X purinergic receptor		
Storage:	Store at -20°C		

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Solvent & Solubility

	≥24.65 mg/mL in DN	\geq 24.65 mg/mL in DMSO; insoluble in H2O; \geq 11.73 mg/mL in EtOH with gentle warming and ultrasonic				
In Vitro	Preparing Stock Solutions	Mass Solvent Concentration	1mg	5mg	10mg	
	Slock Solutions	1 mM	2.0265 mL	10.1327 mL	20.2655 mL	
	PEBIO	5 mM	0.4053 mL	2.0265 mL	4.0531 mL	
		10 mM	0.2027 mL	1.0133 mL	2.0265 mL	

Please refer to the solubility information to select the appropriate solvent.

Biological Activity

Shortsummary

Inactive derivative of KN-93, control compound

IC₅₀ & Target

In Vitro

Cell Viability Assay	and the second
Cell Line:	NIH 3T3 fibroblasts
Preparation method:	The solubility of this compound in DMSO is > 10 mM. General tips for obtaining
	a higher concentration: Please warm the tube at 37 °C for 10 minutes and/or
	shake it in the ultrasonic bath for a while. Stock solution can be stored below
	-20°C for several months.
Reacting conditions:	72 h, 4-24 μM

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	Applications:	KN-92 is an inactive derivative of KN-93. KN-92 is usually used as a control in		
		studies to elucidate the effect of KN-93. KN-93 inhibits fibroblast CaMK-II		
		activity and cell growth, whereas KN-92 had no effect on CaMK-II activity or cell		
		growth.		
	Animal experiment			
In Vivo	Animal models:	AC3-I and AC3-C transgenic mice		
	Dosage form:	20 µmol/kg, intraperitoneal		
	Applications:	Treatment with KN-93 in WT mice resulted in a dose-dependent improvement		
		in left ventricular function compared to WT mice treated with KN-92. Surviving		
		myocytes from infarcted wild-type mice without treatment or treated with control		
		drug KN-92 exhibited severely disordered Ca2+ homeostasis. In contrast,		
		Ca2+ homeostasis was preserved after myocardial infarction in wild-type mice		
		treated with KN-93.		
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may		
	BIO	slightly differ with the theoretical value. This is caused by an experimental		
	PErson	system error and it is normal.		
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Product Citations

See more customer validations on www.apexbt.com.

References



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[1]. Tombes R M, Grant S, Westin E H, et al. G1 cell cycle arrest and apoptosis are induced in NIH 3T3 cells by KN-93, an inhibitor of CaMK-II (the multifunctional Ca2+/CaM kinase)[J]. Cell growth & differentiation: the molecular biology journal of the American Association for Cancer Research, 1995, 6(9): 1063.

[2]. Zhang R, Khoo M S C, Wu Y, et al. Calmodulin kinase II inhibition protects against structural heart disease[J]. Nature medicine, 2005, 11(4): 409-417.

Caution

FOR RESEARCH PURPOSES ONLY. NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.

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