

Hello Bio, Inc.  
304 Wall St., Princeton, NJ 08540 USA

T. 609-683-7500  
F. 609-228-4994

customercare-usa@hellowbio.com



## DATASHEET

KT 5720

### Product overview

<b>Name</b>	KT 5720
<b>Cat No</b>	HB0361
<b>Biological action</b>	Inhibitor
<b>Purity</b>	>98%
<b>Description</b>	Potent, selective PKA inhibitor

### Images



### Biological Data

<b>Biological description</b>	Potent and selective protein kinase A inhibitor ( $K_i = 60$ nM). Arrests skin fibroblasts in the G <sub>1</sub> stage. Reverses multi-drug resistance in leukaemia and attenuates hyperpolarization-activated cyclic nucleotide-gated (HCN) channels in the dorsal root ganglion (DRG). Cell-permeable.
-------------------------------	--

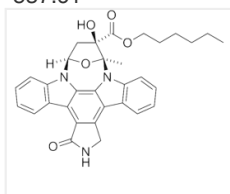
### Solubility & Handling

<b>Storage instructions</b>	-20 °C (desiccate)
<b>Solubility overview</b>	Soluble in DMSO and in methanol
<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>	(9 <i>R</i> ,10 <i>S</i> ,12 <i>S</i> )-2,3,9,10,11,12-Hexahydro-10-hydroxy-9-methyl-1-oxo-9,12-epoxy-1 <i>H</i> -diindolo[1,2,3- <i>fg</i> :3',2',1'- <i>kl</i> ]pyrrolo[3,4- <i>i</i> ][1,6]benzodiazocine-10-carboxylic acid, hexyl ester
<b>Molecular Weight</b>	537.61

**Chemical structure**



<b>Molecular Formula</b>	C <sub>32</sub> H <sub>31</sub> N <sub>3</sub> O <sub>5</sub>
<b>CAS Number</b>	108068-98-0
<b>PubChem identifier</b>	3844
<b>SMILES</b>	CCCCCOC(=O)C1(CC2N3C4=CC=CC=C4C5=C6C(=C7C8=CC=CC=C8N(C7=C53)C1(O2)C)CN C6=O)O
<b>InChi</b>	InChI=1S/C32H31N3O5/c1-3-4-5-10-15-39-30(37)32(38)16-23-34-21-13-8-6-11-18(21)25-26-20(17-33-29(26)36)24-19-12-7-9-14-22(19)35(28(24)27(25)34)31(32,2)40-23/h6-9,11-14,23,38H,3-5,10,15-17H2,1-2H3,(H,33,36)
<b>InChiKey</b>	ZHEHVZXPVFXKEY-UHFFFAOYSA-N
<b>MDL number</b>	MFCD00132118

---

## References

### **K-252 compounds, novel and potent inhibitors of protein kinase C and cyclic nucleotide-dependent protein kinases.**

Kase H *et al* (1987) *Biochem Biophys Res Commun* 142(2)

**PubMedID** [3028414](#)

### **In vitro and in vivo reversal of MDR1-mediated multidrug resistance by KT-5720: implications on hematological malignancies.**

Galski H *et al* (2006) *Leuk Res* 30(9)

**PubMedID** [16542724](#)

### **Multiple kinase arrest points in the G1 phase of nontransformed mammalian cells are absent in transformed cells.**

Gadbois DM *et al* (1992) *Proc Natl Acad Sci U S A* 89(18)

**PubMedID** [1528872](#)

### **Novel role of KT5720 on regulating hyperpolarization-activated cyclic nucleotide-gated channel activity and dorsal root ganglion neuron excitability.**

Cheng Q *et al* (2013) *DNA Cell Biol* 32(6)

**PubMedID** [23713946](#)

---