

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

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

customercare-usa@hellobio.com



## DATASHEET

AM281

### Product overview

<b>Name</b>	AM281
<b>Cat No</b>	HB2312
<b>Biological action</b>	Antagonist
<b>Purity</b>	>98%
<b>Description</b>	Potent, selective CB <sub>1</sub> receptor antagonist / inverse agonist

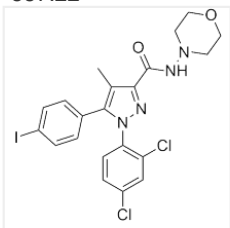
### Biological Data

<b>Biological description</b>	AM 281 is a potent, selective CB <sub>1</sub> receptor antagonist / inverse agonist which is selective for CB <sub>1</sub> over CB <sub>2</sub> receptors (K <sub>i</sub> values are 12 nM and 4200 nM respectively). AM 281 modulates locomotor activity and shows neuroprotective actions in septic shock.
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### Solubility & Handling

<b>Storage instructions</b>	+4 °C (desiccate)
<b>Solubility overview</b>	Soluble in DMSO (50 mM with heating)
<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>	1-(2,4-Dichlorophenyl)-5-(4-iodophenyl)-4-methyl-N-4-morpholinyl-1H-pyrazole-3-carbox amide
<b>Molecular Weight</b>	557.22
<b>Chemical structure</b>	

<b>Molecular Formula</b>	C <sub>21</sub> H <sub>19</sub> Cl <sub>2</sub> IN <sub>4</sub> O <sub>2</sub>
<b>CAS Number</b>	202463-68-1
<b>PubChem identifier</b>	4302962
<b>SMILES</b>	CC1=C(N(N=C1C(=O)NN2CCOCC2)C3=C(C=C(C=C3)Cl)Cl)C4=CC=C(C=C4)I
<b>Source</b>	Synthetic
<b>InChi</b>	InChI=1S/C21H19Cl2IN4O2/c1-13-19(21(29)26-27-8-10-30-11-9-27)25-28(18-7-4-15(22)12-17(18)23)20(13)14-2-5-16(24)6-3-14/h2-7,12H,8-11H2,1H3,(H,26,29)
<b>InChiKey</b>	AJFFBPZYXRNAIC-UHFFFAOYSA-N
<b>MDL number</b>	MFCD01861180
<b>Appearance</b>	White solid

### References

Design and synthesis of the CB<sub>1</sub> selective cannabinoid antagonist AM281: a potential human SPECT ligand.

Lan et al (1999) AAPS PharmSci. 1(2)

**PubMedID** [11741201](#)

**Effect of the cannabinoid receptor SPECT agent, AM 281, on hippocampal acetylcholine release from rat brain slices.**

Gifford et al (1997) Neurosci Lett. 238(1-2)

**PubMedID** [9464661](#)

**Cannabinoid antagonist AM 281 reduces mortality rate and neurologic dysfunction after cecal ligation and puncture in rats.**

Kadoi et al (2005) Crit Care Med 33(11)

**PubMedID** [16276190](#)

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