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DATASHEET

Flumazenil

Product overview

Name	Flumazenil
Cat No	HB0915
Alternative names	Ro 15-1788
Biological action	Antagonist
Purity	>99%
Description	GABA _A receptor antagonist

Images



Biological Data

Biological description	GABA _A receptor antagonist, non-selective for $\alpha 1$, $\alpha 2$, $\alpha 3$ or $\alpha 5$ subunits. Binds at the benzodiazepine site. Used as a reversal agent for benzodiazepines. Reverses tolerance to benzodiazepines and restores the neuroprotective actions of diazepam in ischemia models.
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Solubility & Handling

Storage instructions	+4 °C (desiccate)
Solubility overview	Soluble in DMSO (25mM, gentle warming) or ethanol (5mM)
Important	This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use.

Chemical Data

Chemical name	8-Fluoro-5,6-dihydro-5-methyl-6-oxo -4 <i>H</i> -imidazo[1,5-a][1,4]benzodiazepine-3-carboxylic acid, ethyl ester
Molecular Weight	303.29
Chemical structure	The chemical structure of Flumazenil ethyl ester, showing a benzodiazepine core with a fluorine atom at the 8-position, a methyl group at the 5-position, and an ethyl ester group at the 3-position.

Molecular Formula	C ₁₅ H ₁₄ FN ₃ O ₃
CAS Number	78755-81-4
PubChem identifier	3373
SMILES	CCOC(=O)C1=C2CN(C)C(=O)C3=CC(F)=CC=C3N2C=N1
InChIKey	OFBIFZUFASYRE-UHFFFAOYSA-N

References

Regional differences in the inhibition of mouse in vivo [3H]Ro 15-1788 binding reflect selectivity for alpha 1 versus alpha 2 and alpha 3 subunit-containing GABAA receptors.

Atack JR *et al* (1999) *Neuropsychopharmacology* 20(3)

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Effects of diazepam and flumazenil on forebrain ischaemia in a rat model of benzodiazepine tolerance.

Iwata M *et al* (2012) *Br J Anaesth* 109(6)

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Rapid and reliable sedation induced by diazepam and antagonized by flumazenil in zebra finches (*Taeniopygia guttata*).

Prather JF (2012) *J Avian Med Surg* 26(2)

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