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## DATASHEET

PEAQX (NVP-AAM 077)

### Product overview

<b>Name</b>	PEAQX (NVP-AAM 077)
<b>Cat No</b>	HB2841
<b>Alternative names</b>	NVP-AAM 007
<b>Biological action</b>	Antagonist
<b>Purity</b>	>98%
<b>Customer comments</b>	<i>It works! PEAQX (NVP-AAM 077) blocks a considerable portion of the NMDA-mediated current in whole cell patch set-up. <b>Verified customer, Rutgers University</b></i>
<b>Description</b>	Potent, competitive NMDA receptor antagonist

### Images



### Biological Data

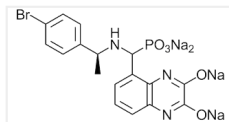
<b>Biological description</b>	Potent and competitive NMDA receptor antagonist. Binds at the glutamate site.
	Shows some selectivity (~7-10-fold) for GluN1/2A (NR1/2A) over GluN1/2B (NR1/2B) subunit containing receptors (IC <sub>50</sub> values are 31 and 215 nM at GluN1/2A and GluN1/2B respectively).
	Shows anticonvulsant activity. Active <i>in vivo</i> .

### Solubility & Handling

<b>Storage instructions</b>	-20°C
<b>Solubility overview</b>	Soluble in water (10mM)
<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>	[[[(1S)-1-(4-Bromophenyl)ethyl]amino][1,2,3,4-tetrahydro-2,3-dioxo-5-quinoxaliny]methyl] phosphonic acid tetrasodium salt
<b>Molecular Weight</b>	542.14

**Chemical structure****Molecular Formula** $C_{17}H_{13}BrN_3Na_4O_5P$ **CAS Number**

459836-30-7

**PubChem identifier**

101043065

**SMILES**CC(C1=CC=C(C=C1)Br)NC(C2=C3C(=CC=C2)N=C(C(=N3)[O-])[O-])P(=O)([O-])[O-].[Na+].[Na+].[Na+].[Na+]**InChi**InChI=1S/C17H17BrN3O5P.4Na/c1-9(10-5-7-11(18)8-6-10)19-17(27(24,25)26)12-3-2-4-13-14(12)21-16(23)15(22)20-13;;;/h2-9,17,19H,1H3,(H,20,22)(H,21,23)(H2,24,25,26);;;/q;4\*+1/p-4**InChiKey**

QBVOZIDKUFSQL-UHFFFAOYSA-J

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## References

### 5-Phosphonomethylquinoxalinediones as competitive NMDA receptor antagonists with a preference for the human 1A/2A, rather than 1A/2B receptor composition.

Auberson et al (2002) Bioorg Med Chem Lett 12(7)

**PubMedID** [11909726](#)

### Equilibrium constants for (R)-[(S)-1-(4-bromophenyl)-ethylamino]-[2,3-dioxo-1,2,3,4-tetrahydroquinoxalin-5-yl)-methyl]-phosphonic acid (NVP-AAM077) acting at recombinant NR1/NR2A and NR1/NR2B NMDA receptors: Implications for studies of sy

Frizelle et al (2006) Mol Pharmacol. 70(3)

**PubMedID** [16778008](#)

### The effects of an acute challenge with the NMDA receptor antagonists, MK-801, PEAQX, and ifenprodil, on social inhibition in adolescent and adult male rats.

Morales and Spear (2014) Psychopharmacology (Berl). 231(8)

**PubMedID** [24043344](#)

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