

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

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

customercare-usa@hellobio.com



## DATASHEET

L-AP4

### Product overview

|                          |  |
|--------------------------|--|
| <b>Name</b>              | L-AP4  |
| <b>Cat No</b>            | HB0370   |
| <b>Description</b>       | L-AP4 is a selective group III mGluR agonist which inhibits synaptic transmission.   |
| <b>Biological action</b> | Agonist  |
| <b>Purity</b>            | >99%   |
| <b>Customer comments</b> | <i>Great product - L-AP4 arrived in time for experiments. Excellent documentation and ease of use. Data looks promising from experiments - happy customer!</i> <b>Verified customer, The University of Newcastle</b> |

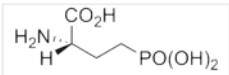
### Biological Data

|                               |  |
|-------------------------------|--|
| <b>Biological description</b> | <p>L-AP4 is a selective group III mGluR agonist (EC<sub>50</sub> values are 0.9, 252, 0.06-0.6 μM at mGlu<sub>4</sub>, mGlu<sub>7</sub>, mGlu<sub>8</sub> receptors respectively and &gt;1000 μM at other mGluRs) and also a weak NMDA receptor agonist.</p> <p>L-AP4 is a potent synaptic depressant.</p> <p>L-AP4 reduces glutamate release, inhibits glutamate-mediated EPSPs in the hippocampus, olfactory cortex and spinal cord and inhibits long term potentiation (LTP) in vivo.</p> <p>Additionally, L-AP4 also inhibits GABA release and reversibly reduces GABA-mediated inhibitory post-synaptic potential (IPSP) amplitude.</p> <p>L-AP4 also shows neuroprotective properties.</p> |
|-------------------------------|--|

### Solubility & Handling

|                             |   |
|-----------------------------|---|
| <b>Storage instructions</b> | Room temperature  |
| <b>Solubility overview</b>  | Soluble in water (5mM) and in 0.1M NaOH (100mM)   |
| <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>      | L-(+)-2-Amino-4-phosphonobutyric acid   |
| <b>Molecular Weight</b>   | 183.1   |
| <b>Chemical structure</b> |  |
| <b>Molecular Formula</b>  | C <sub>4</sub> H <sub>10</sub> NO <sub>5</sub> P                                    |
| <b>CAS Number</b>         | 23052-81-5  |
| <b>PubChem identifier</b> | 179394  |
| <b>SMILES</b>             | C(CP(=O)(O)O)[C@@H](C(=O)O)N  |
| <b>Source</b>             | Synthetic   |
| <b>InChi</b>              | InChI=1S/C4H10NO5P/c5-3(4(6)7)1-2-11(8,9)10/h3H,1-2,5H2,(H,6,7)(H2,8,9,10)/t3-m/s1  |
| <b>InChiKey</b>           | DDOQBQRIEWHWBT-VKHMYYEASA-N   |

**MDL number**  
**Appearance**

MFCD00083244  
White solid

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## References

### **L-AP4, a potent agonist of group III metabotropic glutamate receptor, decreases central action of angiotensin II.**

Fedosiewicz-Wasiluk M *et al* (2002) *Pol J Pharmacol* 54(5)

**PubMedID** [12593528](#)

### **The agonist selectivity of a class III metabotropic glutamate receptor, human mGluR4a, is determined by the N-terminal extracellular domain.**

Tones MA *et al* (1995) *Neuroreport* 7(1)

**PubMedID** [8742431](#)

### **Changes of mGluR4 and the effects of its specific agonist L-AP4 in a rodent model of diffuse brain injury.**

Zhou F *et al* (2003) *J Clin Neurosci* 10(6)

**PubMedID** [14592619](#)

### **Unveiling the functions of presynaptic metabotropic glutamate receptors in the central nervous system.**

Schoepp DD (2001) *J Pharmacol Exp Ther* 299(1)

**PubMedID** [11561058](#)

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