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## DATASHEET

Endomorphin-1

### Product overview

<b>Name</b>	Endomorphin-1
<b>Cat No</b>	HB2410
<b>Biological action</b>	Agonist
<b>Description</b>	Potent, selective $\mu$ -opioid receptor agonist

### Biological Data

<b>Biological description</b>	Potent, selective $\mu$ -opioid receptor agonist ( $K_i$ values are 360 pM and >1000 nM at $\mu$ , $\delta$ and $\kappa$ receptors respectively). Endogenous peptide. Shows >4000 and 15,000-fold selectivity for $\mu$ over $\delta$ and $\kappa$ receptors. Shows antinociceptive activity.
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### Solubility & Handling

<b>Storage instructions</b>	-20°C (desiccate)
<b>Solubility overview</b>	Soluble in water
<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>	Tyr-Pro-Trp-Phe-NH <sub>2</sub>
<b>Molecular Weight</b>	610.67
<b>Chemical structure</b>	
<b>Molecular Formula</b>	C <sub>34</sub> H <sub>38</sub> N <sub>6</sub> O <sub>5</sub>
<b>CAS Number</b>	189388-22-5
<b>PubChem identifier</b>	5311080
<b>SMILES</b>	<chem>C1CC(N(C1)C(=O)C(CC2=CC=C(C=C2)O)N)C(=O)NC(CC3=CNC4=CC=CC=C43)C(=O)NC(CC5=CC=CC=C5)C(=O)N</chem>
<b>InChiKey</b>	ZEXLJFNSKAHNFH-SYKYGTKKSA-N

### References

#### A potent and selective endogenous agonist for the mu-opiate receptor.

Zadina et al (1997) Nature 386

**PubMedID** [9087409](#)

#### Opposite effects of neuropeptide FF on central antinociception induced by endomorphin-1 and endomorphin-2 in mice.

Wang et al (2014) PLoS 9

**PubMedID** [25090615](#)

#### Endomorphin-1 and endomorphin-2: pharmacology of the selective endogenous mu-opioid receptor agonists.

Horvath (2000) Pharmacol Ther 88

