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**产品名称: 精氨酰-脯氨酰-赖氨酰-脯氨酰-谷氨酰-谷氨酰-苯丙氨酰-苯丙氨酰
-SAR-亮氨酸-蛋氨酸[O2]-胺**
产品别名: [Sar9, Met(O2)11]-Substance P

生物活性:

Description	[Sar9, Met(O2)11]-Substance P is a tachykinin NK1 receptor selective agonist.																													
IC₅₀ & Target	NK1 receptor[1]																													
In Vitro	[Sar9, Met(O2)11]-Substance P and peptide (10-100 pmol per rat, i.c.v.) are equipotent in increasing mean arterial blood pressure (MAP) and heart rate (HR), yet they have dissimilar time-course. Both agonists increase dose-dependently face washing and sniffing while [Sar9, Met(O2)11]-Substance P is the sole to produce grooming[1].																													
Solvent&Solubility	<p>In Vitro: H₂O : 50 mg/mL (35.88 mM; Need ultrasonic)</p> <table border="1"><thead><tr><th rowspan="2">Preparing</th><th>Solvent</th><th>Mass</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr><tr><th>Concentration</th><th></th><th></th><th></th><th></th></tr></thead><tbody><tr><td>Stock Solutions</td><td>1 mM</td><td>0.7175 mL</td><td>3.5877 mL</td><td>7.1754 mL</td></tr><tr><td></td><td>5 mM</td><td>0.1435 mL</td><td>0.7175 mL</td><td>1.4351 mL</td></tr><tr><td></td><td>10 mM</td><td>0.0718 mL</td><td>0.3588 mL</td><td>0.7175 mL</td></tr></tbody></table>				Preparing	Solvent	Mass	1 mg	5 mg	10 mg	Concentration					Stock Solutions	1 mM	0.7175 mL	3.5877 mL	7.1754 mL		5 mM	0.1435 mL	0.7175 mL	1.4351 mL		10 mM	0.0718 mL	0.3588 mL	0.7175 mL
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实验参考:

Kinase Assay	Rats initially receive an i.c.v. injection of artificial cerebrospinal fluid (aCSF; 1 µl) followed 60 min later by a single dose of either [Sar9, Met(O2)11]-Substance P (10 pmol (n=9), 25 pmol (n=9), 65 pmol (n=8) or 100 pmol (n=8)) or peptide (10 pmol (n=12), 25 pmol (n=9), 65 pmol (n=6) or 100 pmol (n=6)) to construct a complete dose-response curve. Each rat is selected randomly and injected with only one of the two agonists for the remainder of the protocol. Increasing doses of [Sar9, Met(O2)11]-Substance P or peptide are given at 24 h intervals on day 1 (10 pmol), day 2 (25 pmol), day 3 (65 pmol) and day 4 (100 pmol). Control rats (n=18) receive only the vehicle (aCSF) each day of experiment. Peptides are administered in a volume of 1 µL of vehicle followed by 5 µL flush volume of aCSF which corresponds to the void volume of the catheter. Each dose is calculated per rat in 1 µL solution[1].
References	[1]. Cellier E, et al. Characterization of central and peripheral effects of peptide with the use of five tachykinin NK1 receptor antagonists in the rat. Br J Pharmacol. 1999 Jun;127(3):717-28.