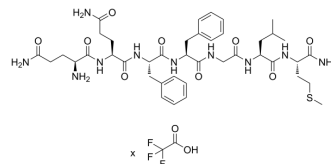


## Substance P (5-11) (TFA)

<b>Cat. No.:</b>	HY-P3890A
<b>Molecular Formula:</b>	C <sub>41</sub> H <sub>60</sub> N <sub>10</sub> O <sub>9</sub> S.xC <sub>2</sub> HF <sub>3</sub> O <sub>2</sub>
<b>Sequence:</b>	Gln-Gln-Phe-Phe-Gly-Leu-Met-NH <sub>2</sub>
<b>Sequence Shortening:</b>	QQFFGLM-NH <sub>2</sub>
<b>Target:</b>	Neurokinin Receptor
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling
<b>Storage:</b>	Sealed storage, away from moisture and light Powder    -80°C    2 years -20°C    1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	H <sub>2</sub> O : 100 mg/mL (Need ultrasonic)
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### BIOLOGICAL ACTIVITY

<b>Description</b>	Substance P (5-11) TFA TFA, the C-terminal heptapeptide of Substance P ( <a href="#">Substance P (HY-P0201)</a> ), is a neuropeptide. Substance P (5-11) TFA TFA binds to NK-1 tachykinin receptor <sup>[1]</sup> .
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<b>In Vitro</b>	When a fraction from rabbit brain enriched in glial cells was incubated with [ <sup>3</sup> H]Substance P (5-11) TFA TFA, an uptake of [ <sup>3</sup> H]Substance P (5-11) TFA TFA is observed. When Substance P is released from nerve terminals, it is hydrolysed into Substance P (5-11) TFA TFA, which is in turn accumulated into glial cells as well as nerve terminals and that this high affinity uptake mechanism may play an important role in terminating the synaptic action of Substance P <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
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### REFERENCES

[1]. A Inoue, et al. Active uptake system for substance P carboxy-terminal heptapeptide (5-11) into a fraction from rabbit enriched in glial cells. Jpn J Pharmacol. 1984 Oct;36(2):137-45.

**Caution: Product has not been fully validated for medical applications. For research use only.**

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