

## [DAla4] Substance P (4-11)

Cat. No.:	HY-P3883
CAS No.:	81381-50-2
Molecular Formula:	C <sub>44</sub> H <sub>65</sub> N <sub>11</sub> O <sub>10</sub> S
Molecular Weight:	940.12
Sequence Shortening:	{D-Ala}-QQFFGLM-NH <sub>2</sub>
Target:	Neurokinin Receptor
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

### BIOLOGICAL ACTIVITY

Description	[DAla4] Substance P (4-11) is an analog of Substance P ( <a href="#">Substance P (HY-P0201)</a> ) that inhibits the binding of <sup>125</sup> I-Bolton Hunter-conjugated Eledoisin ( <a href="#">Eledoisin (HY-P0006)</a> ) (IC <sub>50</sub> of 0.5 μM) and <sup>125</sup> I-Bolton Hunter-conjugated Substance P (IC <sub>50</sub> of 0.15 μM) to rat brain cortex membranes <sup>[1]</sup> .
In Vitro	Eledoisin and Substance P are members of a class of peptides termed tachykinins. They share a similar spectrum of biological activities. [DAla4] Substance P (4-11) ([D-Ala <sup>0</sup> -(5-11)]SP) inhibits the binding of <sup>125</sup> I-Bolton Hunter-conjugated Eledoisin and <sup>125</sup> I-Bolton Hunter-conjugated Substance P to rat brain cortex membranes <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. M A Cascieri, et al. Demonstration of two distinct tachykinin receptors in rat brain cortex. J Biol Chem. 1985 Feb 10;260(3):1501-7.

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

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