



# SZABO SCANDIC

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### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

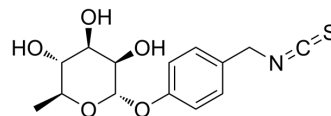
[mail@szabo-scandic.com](mailto:mail@szabo-scandic.com)

[www.szabo-scandic.com](http://www.szabo-scandic.com)

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## Moringin

<b>Cat. No.:</b>	HY-N8264
<b>CAS No.:</b>	73255-40-0
<b>Molecular Formula:</b>	C <sub>14</sub> H <sub>17</sub> NO <sub>5</sub> S
<b>Molecular Weight:</b>	311.35
<b>Target:</b>	TRP Channel
<b>Pathway:</b>	Membrane Transporter/Ion Channel; Neuronal Signaling
<b>Storage:</b>	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



### BIOLOGICAL ACTIVITY

<b>Description</b>	Moringin is a potent and selective TRPA1 ion channel natural agonist with an EC <sub>50</sub> of 3.14 μM. Moringin does not activate or activates very weakly the vanilloids somatosensory channels TRPV1, TRPV2, TRPV3 and TRPV4, and the melastatin cooling receptor TRPM8. Moringin has hypoglycemic, antimicrobial, anti-inflammatory, anticancer and neuroprotection activities <sup>[1]</sup> [2].
<b>In Vitro</b>	In SH-SY5Y human neuroblastoma cells, Moringin (16.4 μM; 24-72 h) significantly reduces SH-SY5Y cell growth in a time and concentration-dependent manner <sup>[2]</sup> . Moringin (1.64-8.2 μM; 24 h) increases the expression of p53, p21, and Bax at both the protein and transcriptional level in SH-SY5Y cells. Moringin significantly increases the gene expression of both caspase 3 and 9 and enhanced their cleavage, thereby initiating an intrinsic apoptotic cascade <sup>[2]</sup> . Moringin inhibits nuclear translocation of NF-κB <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>In Vivo</b>	In experimental autoimmune encephalomyelitis (EAE) mice, Moringin (10 mg/kg; intraperitoneally daily for 5 week) pretreatment normalizes the aberrant Wnt-β-catenin pathway, resulting in GSK3β inhibition and β-catenin upregulation, which regulates T-cell activation (CD4 and FoxP3), suppresses the main inflammatory mediators (IL-1β, IL-6, and COX2), through activation of PPARγ. Moringin increases antioxidant Nrf2 expression in EAE mice <sup>[3]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

- [1]. Gigliola Borgonovo, et al. Moringin, A Stable Isothiocyanate from *Moringa oleifera*, Activates the Somatosensory and Pain Receptor TRPA1 Channel In Vitro. *Molecules*. 2020 Feb 22;25(4):976.
- [2]. Santa Cirmi, et al. Moringin from *Moringa Oleifera* Seeds Inhibits Growth, Arrests Cell-Cycle, and Induces Apoptosis of SH-SY5Y Human Neuroblastoma Cells through the Modulation of NF-κB and Apoptotic Related Factors. *Int J Mol Sci*. 2019 Apr 19;20(8):1930.
- [3]. Sabrina Giacoppo, et al. Moringin activates Wnt canonical pathway by inhibiting GSK3β in a mouse model of experimental autoimmune encephalomyelitis. *Drug Des Devel Ther*. 2016 Oct 4;10:3291-3304.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA