**Proteins** 

# (E)-Naringenin chalcone

Cat. No.: HY-N3007A CAS No.: 25515-46-2 Molecular Formula: C<sub>15</sub>H<sub>12</sub>O<sub>5</sub> Molecular Weight: 272.25 Target: Others

Pathway: Others

Please store the product under the recommended conditions in the Certificate of Storage:

**Product** Data Sheet

## **BIOLOGICAL ACTIVITY**

Description

(E)-Naringenin chalcone is an orally active anti-allergic agent. (E)-Naringenin chalcone also has antioxidant, antiinflammatory activities. (E)-Naringenin chalcone can improve adipocyte functions. (E)-Naringenin chalcone inhibits histamine release from rat peritoneal mast  $cell^{[1][2][4]}$ .

In Vitro

- (E)-Naringenin chalcone (25-125  $\mu$ g/mL, 10 min) inhibits histamine release from rat peritoneal mast cells, with an IC<sub>50</sub> value of 68  $\mu$ g/mL<sup>[1]</sup>.
- (E)-Naringenin chalcone (25-50 μM, 8 days) improves adipocyte (3T3-L1) functions by enhancing adiponectin production<sup>[2]</sup>.
- (E)-Naringenin chalcone (25-100 μM, 24 h) stimulates the activity of PPARγ in NIH-3T3 cells<sup>[2]</sup>.
- (E)-Naringenin chalcone (0-200 µM 24 h) inhibits the production of TNF-alpha, MCP-1, and nitric oxide (NO) by LPSstimulated RAW 264 macrophages<sup>[4]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

RT-PCR<sup>[2]</sup>

Cell Line:	3T3-L1 adipocytes
Concentration:	25-100 μΜ
Incubation Time:	8 days
Result:	Increased adiponectin mRNA levels up to 8.0-fold in a dose-dependent manner.

# Western Blot Analysis<sup>[4]</sup>

Cell Line:	RAW 264 macrophages
Concentration:	0,25, 50, 100, 200 μΜ
Incubation Time:	24 h
Result:	Suppressed the degradation of IkB- $\alpha$ .

In Vivo

- (E)-Naringenin chalcone (0.8 mg/kg, oral administration) shows anti-allergic effect in type I allergic mice<sup>[1]</sup>.
- (E)-Naringenin chalcone (0.8 mg/kg, oral administration) suppresses allergic asthma by inhibiting the type-2 function of CD4 T cells in allergic airway inflammatory mice<sup>[3]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Mouse type I allergic model $^{[1]}$
Dosage:	0.8 mg/kg
Administration:	Oral administration
Result:	Inhibited the ear-swelling response, with the inhibitory ratio of 46.7%.
Animal Model:	Allergic airway inflammation induced in mice <sup>[3]</sup>
Dosage:	0.8 mg/kg
Administration:	Oral administration, daily
Result:	Decreased cell numbers of the infiltrating leukocyte and eosinophils.
	Decreased Muc5ac and gob-5 expression in the lungs.

### **REFERENCES**

- [1]. Taichi Yamamoto, et al. Anti-allergic activity of naringenin chalcone from a tomato skin extract. Biosci Biotechnol Biochem. 2004 Aug;68(8):1706-11.
- [2]. Taro Horiba, et al. Naringenin chalcone improves adipocyte functions by enhancing adiponectin production. Mol Cell Endocrinol. 2010 Jul 29;323(2):208-14.
- [3]. Chiaki Iwamura, et al. Naringenin chalcone suppresses allergic asthma by inhibiting the type-2 function of CD4 T cells. Allergol Int. 2010 Mar;59(1):67-73.
- [4]. Shizuka Hirai, et al. Inhibitory effect of naringenin chalcone on inflammatory changes in the interaction between adipocytes and macrophages. Life Sci. 2007 Sep 29;81(16):1272-9.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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