
Product Data Sheet

Product Name: ONO-8711 dicyclohexylamine

Cat. No.: GC67726

Chemical Properties

Cas. No.

Formula $C_{34}H_{53}ClN_2O_4S$

M.Wt 621.31

Solubility DMSO : 50 mg/mL (80.48 mM); ultrasonic and warming and heat to 60°C)

Storage 4°C, away from moisture

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure

Background

ONO-8711 dicyclohexylamine is a selective and orally active **EP1** competitive antagonist with K_i value of 0.6 nM and 1.7 nM for human and mouse EP1 respectively. ONO-8711 dicyclohexylamine effectively reduces tumor incidence and multiplicity in mouse models of colon, breast, and oral cancer^[1].

ONO-8711 (10 and 30 μ M; 30 min) blocks the contractions induced by sulprostone in human pulmonary veins in a non-competitive manner^[2].

ONO-8711 inhibits PGE₂-induced increase in cytosolic Ca²⁺ concentration with IC₅₀s of 0.21 μ M, 0.05 μ M, and 0.22 μ M for the mouse, human, and rat receptors, respectively^[3].

ONO-8711 (400 or 800 p.p.m.; p.o.; for 20 weeks) suppresses cancer incidence and delays occurrence of breast tumors^[3].

Animal Model: Female Sprague-Dawley rats (induced breast cancer by gavage of 85 mg/kg PhIP 4 times for 2 weeks)

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

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Dosage: 400 or 800 p.p.m.

Administration: p.o.; for 20 weeks

Result: Did not induce any symptoms of toxicity at 800 p.p.m. Delayed occurrence of breast tumors for 2 or 4 weeks at 400 or 800 p.p.m., respectively. Significantly suppressed cancer incidence compared with the control diet group at 800 p.p.m. (56% versus 79%, $P < 0.05$).

[1]. Watanabe K, et al. Role of the prostaglandin E receptor subtype EP1 in colon carcinogenesis. *Cancer Res.* 1999 Oct 15;59(20):5093-6.

[2]. Norel X, et al. Vasoconstriction induced by activation of EP1 and EP3 receptors in human lung: effects of ONO-AE-248, ONO-DI-004, ONO-8711 or ONO-8713. *Prostaglandins Other Lipid Mediat.* 2004 Oct;74(1-4):101-12.

[3]. Kawamori T, et al. Chemopreventive effects of ONO-8711, a selective prostaglandin E receptor EP(1) antagonist, on breast cancer development. *Carcinogenesis.* 2001 Dec;22(12):2001-4.

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