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## Product Data Sheet

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Product Name: Calindol (hydrochloride)

Cat. No.: GC12620

**Chemical Properties**

Cas. No. 729610-18-8

Chemical Name N-[(1R)-1-(1-naphthalenyl)ethyl]-1H-indole-2-methanamine, monohydrochloride

SMILES C[C@H](C1=C(C=CC=C2)C2=CC=C1)NCC3=CC4=CC=CC=C4N3.ClFormula  $C_{21}H_{20}N_2 \cdot HCl$ 

M.Wt 336.9

Solubility  $\leq 15\text{mg/ml}$  in ethanol;  $50\text{mg/ml}$  in DMSO;  $50\text{mg/ml}$  in dimethyl formamideStorage Store at  $-20^\circ\text{C}$ General tips For obtaining a higher solubility, please warm the tube at  $37^\circ\text{C}$  and shake it in the ultrasonic bath for a while. Stock solution can be stored below  $-20^\circ\text{C}$  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**

Calindol is an activator of the calcium-sensing receptor (CaSR).

The extracellular calcium-sensing receptor (CaSR), a G protein-coupled receptor (GPCR), senses extracellular calcium  $[Ca^{2+}]_e$  and regulates calcium ion homeostasis. The CaSR has a long amino terminal tail typical of family 3 of GPCRs including the metabotropic glutamate receptors, the c-aminobutyric acid B receptor and certain pheromone and taste receptors. CaSR cDNA has been cloned from such tissues as the parathyroid, thyroid, kidney as well as brain. The CaSR at the surface can detect and respond to small changes of circulating  $[Ca^{2+}]_e$  in the particular case of the parathyroid cell.

In vitro: Previous study found that calindol could display improved calcimimetic activity compared to its N2-(2-chloro-(or 4-fluoro-)benzyl)-N1-(1-(1-naphthyl)ethyl)-3-phenylpropane-1,2-diamine analogs as well as stereoselectivity. In the presence of 2 mM  $Ca^{2+}$ , calindol was able to stimulate  $[(3)H]$ inositol phosphates accumulation with an

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

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EC(50) of 1.0+/-0.1 or 0.31+/-0.05  $\mu$ M in cells expressing the rat or the human CaSR, respectively. Therefore, such calcimimetic activities of this novel compound were shown to be because of a specific interaction with the CaSR [1].

In vivo: Up to now, there is no animal in vivo data reported.

Clinical trial: So far, no clinical study has been conducted.

Reference:

[1] Kessler, A. ,Faure, H.,Petrel, C., et al. N2-benzyl-N1-(1-(1-naphthyl)ethyl)-3-phenylpropane-1,2-diamines and conformationally restrained indole analogues: Development of calindol as a new calcimimetic acting at the calcium sensing receptor. *Bioorganic & Medicinal Chemistry Letters* 14(12), 3345-3349 (2004).

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