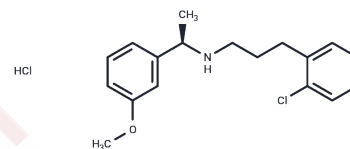


Tecalct Hydrochloride

Chemical Properties

CAS No. :	177172-49-5
Formula:	C ₁₈ H ₂₃ Cl ₂ NO
Molecular Weight:	340.29
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA.



Biological Description

Description	Tecalct Hydrochloride (R-568 hydrochloride) is an allosteric and positive modulator of the calcium-sensing receptor(CaSR). Tecalct Hydrochloride increases the sensitivity to the activation of extracellular Ca ²⁺ .
Targets(IC50)	CaSR
In vitro	Tecalct Hydrochloride (0.1-100 nM) shifts the concentration-response curve for extracellular Ca ²⁺ to the left and decreases the EC ₅₀ value to 0.61 mM. Tecalct Hydrochloride (0.1-100 μM) increases Ca ²⁺ in a dose-dependent and stereoselective manner[3].
In vivo	In male Sprague-Dawley rats with renal insufficiency, Tecalct Hydrochloride (1.5 and 15 mg/kg, oral) inhibits PT cell proliferation and reduces serum PTH level in a dose-dependent manner. Tecalct Hydrochloride reduces the number of BrdU-positive PT cells by 20% at a low dose (1.5 mg/kg), and by 50% at a high dose (15 mg/kg)[1].

Solubility Information

Solubility	DMSO: 150 mg/mL (440.8 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
In vivo Formulation	10% DMSO+40% PEG300+5% Tween 80+45% Saline: 2 mg/mL (5.88 mM),Sonication is recommended. <i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i>

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.9387 mL	14.6933 mL	29.3867 mL
5 mM	0.5877 mL	2.9387 mL	5.8773 mL
10 mM	0.2939 mL	1.4693 mL	2.9387 mL
50 mM	0.0588 mL	0.2939 mL	0.5877 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Wada, M., et al. The calcimimetic compound NPS R-568 suppresses parathyroid cell proliferation in rats with renal insufficiency. Control of parathyroid cell growth via a calcium receptor. *Journal of Clinical Investigation* 100(12), 2977-2983 (1997).

Nemeth, E.F., et al. The parathyroid calcium receptor: a novel therapeutic target for treating hyperparathyroidism. *Pediatr.Nephrol.* 10(3), 275-279 (1996).

Nemeth, E.F., et al. Calcimimetics with potent and selective activity on the parathyroid calcium receptor. *Proceedings of the National Academy of Sciences of the United States of America* 95(7), 4040-4045 (1998).

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