Proteins

Minoxidil

Cat. No.: HY-B0112 CAS No.: 38304-91-5 Molecular Formula: C₉H₁₅N₅O Molecular Weight: 209.25

Target: Potassium Channel; Endogenous Metabolite

Pathway: Membrane Transporter/Ion Channel; Metabolic Enzyme/Protease

-20°C Storage: Powder 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

Ethanol: 7.14 mg/mL (34.12 mM; Need ultrasonic)

H₂O: 1 mg/mL (4.78 mM; ultrasonic and warming and heat to 50°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.7790 mL	23.8949 mL	47.7897 mL
	5 mM	0.9558 mL	4.7790 mL	9.5579 mL
	10 mM	0.4779 mL	2.3895 mL	4.7790 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- 1. Add each solvent one by one: 50% PEG300 >> 50% saline Solubility: 5 mg/mL (23.89 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: PBS Solubility: 1.96 mg/mL (9.37 mM); Clear solution; Need ultrasonic and warming and heat to 60°C
- 3. Add each solvent one by one: 10% EtOH >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 0.71 mg/mL (3.39 mM); Clear solution
- 4. Add each solvent one by one: 10% EtOH >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 0.71 mg/mL (3.39 mM); Clear solution
- 5. Add each solvent one by one: 10% EtOH >> 90% corn oil Solubility: ≥ 0.71 mg/mL (3.39 mM); Clear solution
- 6. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 0.5 mg/mL (2.39 mM); Clear solution
- 7. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 0.5 mg/mL (2.39 mM); Clear solution
- 8. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 0.5 mg/mL (2.39 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Minoxidil (U10858) is an ATP-sensitive potassium (K_{ATP}) channel opener, a potent oral antihypertensive agent and a peripheral vasodilator that promotes vasodilation also affects hair growth. Minoxidil is also a potent inhibitor of soybean lipoxygenaseare with an IC ₅₀ of 20 μ M ^{[1][2][3]} .		
IC ₅₀ & Target	IC50: 20 μ M (soybean lipoxygenaseare) $^{[1]}$; ATP-sensitive potassium channel $^{[2]}$		
In Vitro	Minoxidil (1-00 μ M; 24 hours; RAMEC cells) treatment shows very low cytotoxicities in the whole area of concentrations examined (from 1 μ M to 100 μ M) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
In Vivo	Minoxidil (0.01 mmoL/kg body weight; intraperitoneal injection; for 3.5 hours; fisher 344 rats) treatment inhibits carrageenan-induced rat paw oedema with an inhibitory potency (49%) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Fisher 344 rats (150-200 g) with oedema ^[1]	
	Dosage:	0.01 mmol/kg body weight	
	Administration:	Intraperitoneal injection; for 3.5 hours	
	Result:	Inhibition of the carrageenin-induced oedema.	

CUSTOMER VALIDATION

- Cell Rep. 2021 Jun 8;35(10):109225.
- Sys Rev Pharm. 2021;12(1):402-410.

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REFERENCES

- [1]. Hadjipavlou-Litina D, et al. Synthesis and evaluation of the antioxidative potential of minoxidil-polyamine conjugates. Biochimie. 2013 Jul;95(7):1437-49. doi: 10.1016/j.biochi.2013.03.009. Epub 2013 Mar 28.
- [2]. Davies GC, et al. Novel and established potassium channel openers stimulate hair growth in vitro: implications for their modes of action in hair follicles. J Invest Dermatol. 2005 Apr;124(4):686-94.
- [3]. Cohen RL, et al. Direct effects of minoxidil on epidermal cells in culture. J Invest Dermatol. 1984 Jan;82(1):90-3.

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

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