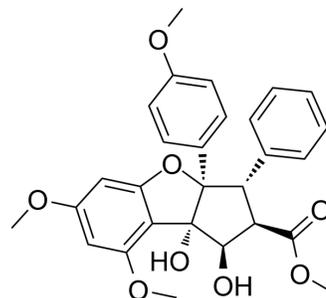


Aglafoline

Cat. No.:	HY-19354		
CAS No.:	143901-35-3		
Molecular Formula:	C ₂₈ H ₂₈ O ₈		
Molecular Weight:	492.52		
Target:	Others		
Pathway:	Others		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro

Ethanol : 100 mg/mL (203.04 mM; Need ultrasonic)
 DMSO : 21.43 mg/mL (43.51 mM; Need ultrasonic)

	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.0304 mL	10.1519 mL	20.3037 mL
	5 mM	0.4061 mL	2.0304 mL	4.0607 mL
	10 mM	0.2030 mL	1.0152 mL	2.0304 mL

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

In Vivo

- Add each solvent one by one: 10% EtOH >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 2.5 mg/mL (5.08 mM); Clear solution
- Add each solvent one by one: 10% EtOH >> 90% corn oil
Solubility: ≥ 2.5 mg/mL (5.08 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 2.14 mg/mL (4.35 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.14 mg/mL (4.35 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Aglafoline inhibits in a selective and concentration-dependent manner the aggregation and ATP release reaction induced in washed rabbit platelets by PAF (platelet-activating factor). The IC₅₀ values of Aglafoline on PAF (3.6 nM)-induced platelet aggregation were about 50 μM. Target: PAF in vitro: Aglafoline also inhibits [3H]PAF (3.6 nM) binding to washed rabbit platelets with an IC₅₀ value of 17.8 ± 2.6 μM. The concentration-response curve of PAF-induced platelet

aggregation was shifted to the right by Aglafoline with pA₂ and pA₁₀ values of 5.97 and 5.04, respectively. Although thromboxane B₂ formation caused by collagen and thrombin was partially suppressed by Aglafoline, thromboxane B₂ formation caused by ionophore A23187 and arachidonic acid was not affected. Aglafoline inhibited the [3H]inositol monophosphate formation caused by PAF but not that caused by collagen or thrombin in the presence of indomethacin (20 μM). [1]in vivo: The cAMP content of washed rabbit platelets was not affected by Aglafoline. Rat femoral intravenous administration of Aglafoline (10 mg/kg) did not affect blood pressure. However, Aglafoline (10 mg/kg) both prophylactically and therapeutically antagonized PAF (2.5 μg/kg)-induced hypotensive shock in rats. Intravenous PAF (30 ng/kg) caused severe bronchoconstriction in guinea pigs. This effect was completely blocked by Aglafoline. This implies Aglafoline is an effective PAF antagonist not only in vitro, but also in vivo.[1]

CUSTOMER VALIDATION

- Mol Cell. 2019 Feb 21;73(4):738-748.e9.

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REFERENCES

[1]. Ko FN, et al. PAF antagonism in vitro and in vivo by aglafoline from *Aglaiia elliptifolia* Merr. Eur J Pharmacol. 1992 Jul 21;218(1):129-35.

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

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