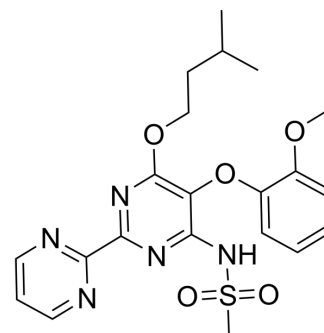


ET receptor antagonist 1

Cat. No.:	HY-155737
Molecular Formula:	C ₂₁ H ₂₅ N ₅ O ₅ S
Molecular Weight:	459.52
Target:	Estrogen Receptor/ERR
Pathway:	Vitamin D Related/Nuclear Receptor
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	ET receptor antagonist 1 (compound 16h) is an orally active ET receptor antagonist (IC ₅₀ =0.18 nM), which can be used for research in pulmonary arterial hypertension (PAH). ET receptor antagonist 1 attenuates monocrotaline (HY-N0750) induced PAH in rat model ^[1] .
IC₅₀ & Target	IC ₅₀ : 0.18 nM (ET receptor) ^[1]
In Vivo	ET receptor antagonist 1 (compound 16h) (150 mg/kg/d, 300 mg/kg/d; po; 48 h following MCT administration, for 21-26 d) results significant reduction in mPAP, and decreases the levels of HIF1α, ANP, and TNNI3 in MCT-exposed rats. And shows antioxidant profile and inhibits lipid peroxidation ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Panchal J, et al. Development of novel bosentan analogues as endothelin receptor antagonists for pulmonary arterial hypertension. Eur J Med Chem. 2023 Nov 5;259:115681.

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

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