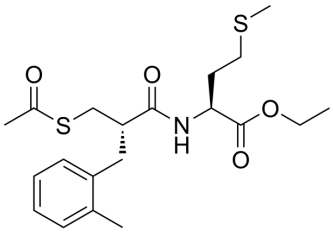
SCH 42495

Cat. No.:	HY-101682
CAS No.:	136511-43-8
Molecular Formula:	$C_{20}H_{29}NO_4S_2$
Molecular Weight:	411.58
Target:	Neprilysin
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	SCH 42495 is an orally active neutral metalloendopeptidase (NEP) inhibitor with antihypertensive effect. SCH 42495 is the orally active ethylester proagent of SCH 42354[1].
IC_{50} & Target	NEP[1]
In Vitro	SCH 42354 selectively inhibits hydrolysis of leu-enkephalin and ANF (IC_{50} of 8.3 and 10.0 nM, respectively) in vitro[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	SCH 42495 (30 mg/kg; oral gavage; twice daily) causes a significant reduction in the pulmonary vascular remodelling and ventricular hypertrophy in hypoxic rats after 10 days[2]. Treatment with SCH 42495 (30 mg/kg; oral gavage; twice daily) leads to a decrease in cardiovascular remodelling secondary to chronic hypoxia in rats[2]. SCH 42495 (oral doses of 1, 3, or 10 mg/kg) produces significant reductions in blood pressure in DOCA-N a hypertensive rats of 22±6, 43±7, and 62±12 mm Hg, respectively[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Hypoxic rats[2]
Dosage:	30 mg/kg
Administration:	Oral gavage; twice daily for 10 days
Result:	Caused a significant reduction in the pulmonary vascular remodelling and ventricular hypertrophy. Led to a decrease in cardiovascular remodelling secondary to chronic hypoxia.

REFERENCES

[1]. Watkins RW, et al. Atrial natriuretic factor potentiating and hemodynamic effects of SCH 42495, a new, neutral metalloendopeptidase inhibitor. Am J Hypertens. 1993 May;6(5 Pt 1):357-68.

[2]. Thompson JS, et al. Effects of the neutral endopeptidase inhibitor, SCH 42495, on the cardiovascular remodelling secondary to chronic hypoxia in rats. Clin Sci (Lond). 1994 Jul;87(1):109-14.

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