TRPC3/6-IN-1

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BIOLOGICAL ACTIVITY							
Description	TRPC3/6-IN-1 is a potent selectivity blocker of the canonical transient receptor channels (TRPC3/6), has block potency for hTRPC3 and hTRPC6 with IC ₅₀ values of 1260 nM and 500 nM, respectively. TRPC3/6-IN-1 can be used for the research of chronic models of heart failure ^[1] .						
IC ₅₀ & Target	hTRPC3 1260 nM (IC ₅₀)						
In Vitro	TRPC3/6-IN-1 has block potency for hTRPC3 and hTRPC6 with IC ₅₀ values of 1260 nM and 500 nM, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.						
In Vivo	TRPC3/6-IN-1 (compound 3) has good pharmacokinetic in rats ^[1] . Pharmacokinetic Parameters of TRPC3/6-IN-1 in rats ^[1] .						
	Compd Dos	e(iv)(mg/kg) C _{max}	(iv)(ng/ml) CL(i	v)(ml/min/kg)	MRT(iv)(h)	DNAUC(iv)(h•kg/L)	%F
	3	0.93	250	120	0.3	0.14	~1
	MCE has not independently confirmed the accuracy of these methods. They are for reference only.						

REFERENCES

[1]. David G Washburn, et al. The discovery of potent blockers of the canonical transient receptor channels, TRPC3 and TRPC6, based on an anilino-thiazole pharmacophore. Bioorg Med Chem Lett. 2013 Sep 1;23(17):4979-84.

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

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