EPAC 5376753

MedChemExpress

Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway: Storage:	HY-111446302826-61-5C15H8Cl2N2O3S367.21RasGPCR/G ProteinPlease store the product under the recommended conditions in the Certificate of Analysis.	
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BIOLOGICAL ACTIVITY		
Description	EPAC 5376753 is an allosterically inhibitor of Epac which inhibits Epac1 with an IC $_{50}$ of 4 μ M in Swiss 3T3 cells.	
IC ₅₀ & Target	IC50: 4 μM (Epac) ^[1]	
In Vitro	Epac is a key mediator of cAMP signaling. EPAC 5376753, the thiobarbituric acid modification of 5225554, has a greater ability to decrease the conformational changes of Epac necessary for its activation and to inhibit Epac signaling in cells. 5376753 selectively inhibits Epac-mediated migration in fibroblasts ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

PROTOCOL	
FROTOCOL	
Cell Assay ^[1]	Swiss 3T3 cells are treated with EPAC 5376753 (1, 10, 25, 50, 75, 100 μ M) or 1% dimethyl sulfoxide for 48 h. Cell viability is assessed using the CellTiter-Glo assay ^[1] .
	MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Brown LM, et al. Allosteric inhibition of Epac: computational modeling and experimental validation to identify allosteric sites and inhibitors. J Biol Chem. 2014 Oct 17;289(42):29148-57.

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

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Product Data Sheet