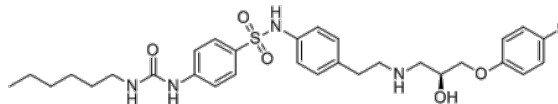


**Product Name** : L755507  
**Cat. No.** : PC-38661  
**CAS No.** : 159182-43-1  
**Molecular Formula** : C<sub>30</sub>H<sub>40</sub>N<sub>4</sub>O<sub>6</sub>S  
**Molecular Weight** : 584.732  
**Target** : c-Myc  
**Solubility** : 10 mM in DMSO



## Biological Activity

L755507 is a potent, selective  $\beta$ <sub>3</sub>-adrenoceptor ( **$\beta$ <sub>3</sub>-AR**) agonist, exhibits robust concentration-dependent increase in cAMP accumulation in CHO-K1 cells expressing human  $\beta$ <sub>3</sub>-adrenoceptors (pEC<sub>50</sub>=12.3), also effectively inhibits the **c-Myc-MAX** heterodimerization.

L755507 increases phosphorylation of extracellular signal-regulated kinase 1/2 (Erk1/2) with pEC<sub>50</sub> of 11.7, couples to both G<sub>s</sub> and G<sub>i</sub> to activate adenylate cyclase and MAPK signaling.

L755507 efficiently restricts the growth of diverse Myc-expressing cells with low micromolar IC<sub>50</sub> values, decreases expression of c-Myc target genes.

L755507 binds to the c-Myc peptide and thereby stabilizes the helix-loop-helix conformation of the c-Myc transcription factor.

## References

Sato M, et al. *Mol Pharmacol*. 2008 Nov;74(5):1417-28.

Hutchinson DS, et al. *Br J Pharmacol*. 2002 Apr;135(8):1903-14.

Singh A, et al. *J Biol Chem*. 2021 Jul;297(1):100903.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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