



**LXR-623** 

**Catalog No: tcsc1721** 

	Available Sizes
Size: 5r	ng
Size: 10	Omg
<b>Size:</b> 50	Omg
Size: 10	00mg
Size: 20	00mg
S	Specifications
<b>CAS No</b> 875787-	
Formul	
<b>Pathwa</b> Metabol	<b>iy:</b> ic Enzyme/Protease
Target:	
<b>Purity</b> / >98%	' Grade:
<b>Solubil</b> i DMSO :	ity: ≥ 47 mg/mL (111.17 mM)
<b>Alterna</b> WAY 253	itive Names: 2623





## **Observed Molecular Weight:**

422.78

## **Product Description**

LXR-623 is a brain-penetrant partial  $\mathbf{LXR}\alpha$  and full  $\mathbf{LXR}\beta$  agonist, with  $\mathbf{IC}_{50}$ s of 24 nM and 179 nM, respectively.

IC50 & Target: IC50: 24 nM (LXR- $\alpha$ ), 179 nM (LXR- $\beta$ )<sup>[2][3]</sup>

In Vitro: LXR-623 potently kills U87EGFRvIII and GBM39 cells in vitro while completely sparing NHAs. LXR-623 also increases ABCA1 protein and decreases LDLR protein levels in all three cell lines. LXR-623 suppresses LDLR expression, increases expression of the ABCA1 efflux transporter, and induces substantial cell death in all of the GBM samples tested. LXR-623 (5  $\mu$ M) also induces GBM cell death through activation of LXR $\beta$ <sup>[1]</sup>. LXR-623 treatment of human PBMC in vitro significantly increases transcription of ABCA1 and ABCG1<sup>[4]</sup>.

In Vivo: LXR-623 (400 mg/kg, p.o.) crosses the blood-brain barrier, induces target gene expression, and achieves therapeutic levels in GBM cells in the brain with minimal activity in the periphery. LXR-623 inhibits tumor growth, promotes tumor cell death, and prolongs the survival of mice bearing intracranial patient-derived GBMs<sup>[1]</sup>. LXR-623 (1.5, 5 mg/kg/day) significantly reduces progression of atherosclerosis in animals compared with the placebo group<sup>[2]</sup>. WAY-252623 (15 and 50 mg/kg) results in a significant reduction of atherosclerosis in a dose-dependent manner. WAY-252623 (20, 60, and 120 mg/kg/day, p.o.) displays neutral lipid effects in this CETP-expressing Syrian hamster<sup>[3]</sup>. Moreover, LXR-623 (50 mg/kg) induces gene expression in rodent peripheral blood cells in rat. LXR-623 (0, 15 and 50 mg/kg) dose-dependently upregulates transcription of ABCA1 and ABCG1 in monkey whole blood cells proportional to dose<sup>[4]</sup>.

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!