

# Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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# Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

## SZABO-SCANDIC HandelsgmbH

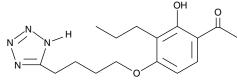
Quellenstraße 110, A-1100 Wien T. +43(0)1 489 3961-0 F. +43(0)1 489 3961-7 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com

# Product Information

### LY171883

Item No. 70710

CAS Registry No.:	88107-10-2	
Formal Name:	1-[2-hydroxy-3-propyl-4-[4-(1H-tetrazol-5-	
	yl)butoxy]phenyl]-ethanone	
MF:	$C_{16}H_{22}N_4O_3$	
FW:	318.4	N'
Purity:	≥98%	\ N
Stability:	≥2 years at room temperature	
Supplied as:	A crystalline solid	
UV/Vis.:	λ <sub>max</sub> : 218, 284 nm	



#### Laboratory Procedures

For long term storage, we suggest that LY171883 be stored as supplied at room temperature. It should be stable for at least two years.

LY171883 is supplied as a crystalline solid. A stock solution may be made by dissolving the LY171883 in an organic solvent purged with an inert gas. LY171883 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide. The solubility of LY171883 in these solvents is approximately 25 mg/ml.

Further dilutions of the stock solution into aqueous buffers or isotonic saline should be made prior to performing biological experiments. Ensure that the residual amount of organic solvent is insignificant, since organic solvents may have physiological effects at low concentrations. For greater aqueous solubility, LY171883 can be directly dissolved in 0.5 M Na<sub>2</sub>CO<sub>3</sub> (15 mg/ml) and then diluted with PBS (pH 7.2) to achieve the desired concentration or pH. We do not recommend storing the aqueous solution for more than one day.

LY171883 is a selective, potent, orally active antagonist of the leukotriene  $D_4$  receptor.<sup>1</sup> Dissociation constants ( $K_B$ ) for LY171883 on guinea pig ileum and parenchyma are 0.07 and 0.34 µM, respectively.<sup>1</sup> LY171883 is an inhibitor of phosphodiesterase obtained from human polymorphonuclear leukocytes (IC50 of 22.6 µM) and various guinea pig tissues  $(IC_{50}s$  range from 6.9-209  $\mu$ M).<sup>1</sup> At a concentration of 50-100  $\mu$ M, LY171883 binds to the PPAR $\gamma$  nuclear receptor, inducing adipogenesis in cultured NIH3T3 fibroblasts.<sup>2,3</sup>

#### References

- 1. Fleisch, J.H., Rinkema, L.E., Haisch, K.D., et al. LY171883, 1-<2-hydroxy-3-propyl-4-<4-(1H-tetrazol-5-yl) butoxy>phenyl>ethanone, an orally active leukotriene D4 antagonist. J. Pharmacol. Exp. Ther. 233, 148-157 (1985).
- Forman, B.M., Tontonoz, P., Chen, J., et al. 15-Deoxy- $\Delta^{12,14}$ -prostaglandin J<sub>2</sub> is a ligand for the adipocyte 2. determination factor PPARy Cell 83, 803-812 (1995).
- Tontonoz, P., Hu, E., Spiegelman, B.M. Stimulation of adipogenesis in fibroblasts by PPARy2, a lipid-activated 3. transcription factor. Cell 79, 1147-1156 (1994).

#### **Related Products**

For a list of related products please visit: www.caymanchem.com/catalog/70710

#### Cayman Chemical

#### **Mailing address**

1180 E. Ellsworth Road Ann Arbor, MI 48108 USA

Phone (800) 364-9897 (734) 971-3335

Fax (734) 971-3640

#### E-Mail

custserv@caymanchem.com

#### Web

www.caymanchem.com

WARNING: This product is for laboratory research only: not for administration to humans. Not for human or veterinary DIAGNOSTIC OR THERAPEUTIC USE.

#### MATERIAL SAFETY DATA

This material should be considered hazardous until information to the contrary becomes available. Do not ingest, swallow, or inhale. Do not get in eyes, on skin, or on clothing. Wash thoroughly after handling. This information contains some, but not all, of the information required for the safe and proper use of this material. Before use, the user must review the complete Material Safety Data Sheet, which has been sent *via* email to your institution.

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