PRODUCT INFORMATION



LXR-623

Item No. 21117

CAS Registry No.: 875787-07-8

2-[(2-chloro-4-fluorophenyl)methyl]-3-(4-fluorophenyl)-Formal Name:

7-(trifluoromethyl)-2H-indazole

Synonym: WAY-252623 MF: C₂₁H₁₂CIF₅N₂

422.8 FW: **Purity:** ≥98%

UV/Vis.: λ_{max} : 209, 318 nm Supplied as: A crystalline solid

Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



LXR-623 is supplied as a crystalline solid. A stock solution may be made by dissolving the LXR-623 in the solvent of choice, which should be purged with an inert gas. LXR-623 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide. The solubility of LXR-623 in these solvents is approximately

LXR-623 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, LXR-623 should first be dissolved in ethanol and then diluted with the aqueous buffer of choice. LXR-623 has a solubility of approximately 0.3 mg/ml in a 1:2 solution of ethanol:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

LXR-623 is a liver X receptor (LXR) modulator that acts as a partial agonist at LXRQ and a full agonist at LXR β (IC₅₀s = 179 and 24 nM, respectively).^{1,2} It is orally bioavailable and readily passes the blood-brain barrier. 1-3 LXR-623 is anti-atherogenic in animal models of atherosclerosis, showing synergistic effects when combined with simvastatin (Item Nos. 10010344 | 10010345). 1,2,4

References

- 1. DiBlasio-Smith, E., Arai, M., Quinet, E.M., et al. Discovery and implementation of transcriptional biomarkers of synthetic LXR agonists in peripheral blood cells. J. Transl. Med. 6, 59 (2008).
- Quinet, E.M., Basso, M.D., Halpern, A.R., et al. LXR ligand lowers LDL cholesterol in primates, is lipid neutral in hamster, and reduces atherosclerosis in mouse. J. Lipid. Res. 50(12), 2358-2370 (2009).
- Villa, G.R., Hulce, J.J., Zanca, C., et al. An LXR-cholesterol axis creates a metabolic co-dependency for brain cancers. Cancer Cell. 30(5), 683-693 (2016).
- Giannarelli, C., Cimmino, G., Connolly, T.M., et al. Synergistic effect of liver X receptor activation and simvastatin on plaque regression and stabilization: An magnetic resonance imaging study in a model of advanced atherosclerosis. Eur. Heart J. 33(2), 264-273 (2012).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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