PRODUCT INFORMATION



BMS 779788

Item No. 27339

	0,	918348-67-1	
Form	al Name:	2-[1-(2-chlorophenyl)-1-methylethyl]-α,α-	\times \times \times \times
		dimethyl-1-[3'-(methylsulfonyl)[1,1'-biphenyl]-	
		4-yl]-1H-imidazole-4-methanol	V_N (
MF:		C ₂₈ H ₂₉ CIN ₂ O ₃ S	
FW:		509.1	
Purity	y:	≥98%	
UV/V	/is.:	λ _{max} : 258 nm	<u> </u>
Supp	lied as:	A crystalline solid	
Stora	ge:	-20°C	
Stabi	lity:	≥4 years	
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Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

BMS 779788 is supplied as a crystalline solid. A stock solution may be made by dissolving the BMS 779788 in the solvent of choice, which should be purged with an inert gas. BMS 779788 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide. The solubility of BMS 779788 in these solvents is approximately 30 mg/ml.

Description

BMS 779788 is a partial agonist of liver X receptor α (LXR α) and LXR β (K_is = 68 and 14 nM, respectively, in a radioligand binding assay).¹ It activates LXR α and LXR β with EC₅₀ values of 230 and 250 nM, respectively, in a transactivation assay. BMS 779788 increases expression of ATP-binding cassette transporter (ABCA1) in HeLa cells and isolated human and mouse whole blood ($EC_{50}s = 33$, 1,200, and 120 nM, respectively). It also increases expression of Abca1 and Abcg2 in isolated mouse blood cells but does not increase plasma or hepatic triglycerides when administered at a dose of 10 mg/kg.

References

1. Kick, E., Martin, R., Xie, Y., et al. Liver X receptor (LXR) partial agonists: Biaryl pyrazoles and imidazoles displaying a preference for LXRB. Bioorg. Med. Chem. Lett. 25(2), 372-377 (2015).

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

SAFETY DATA

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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