PRODUCT INFORMATION



Alobresib

Item No. 30726

CAS Registry No.:	1637771-14-2	
Formal Name:	2-cyclopropyl-5-(3,5-dimethyl-4-isoxazolyl)-α,α-	
	di-2-pyridinyl-1H-benzimidazole-7-methanol	
Synonym:	GS-5829	тарана Н
MF:	C ₂₆ H ₂₃ N ₅ O ₂	
FW:	437.5	
Purity:	≥95%	
Supplied as:	A solid	
Storage:	-20°C	9
Stability:	≥4 years	N N
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Laboratory Procedures

Alobresib is supplied as a solid. A stock solution may be made by dissolving the alobresib in the solvent of choice, which should be purged with an inert gas. Alobresib is soluble in the organic solvent DMSO at a concentration of approximately 10 mM.

Description

Alobresib is an inhibitor of bromodomain and extra-terminal domain (BET) proteins.^{1,2} It inhibits proliferation of MEC-1 chronic lymphocytic leukemia cells (CLL; IC₅₀ = 46.4 nM).¹ Alobresib (400 nM) decreases the levels of Blk and Myc and increases the levels of IkBa in patient-derived CLL cells co-cultured with nurse-like cells (NLC). It inhibits proliferation of USPC-ARK-1 and USPC-ARK-2 uterine serous carcinoma (USC) cells with IC₅₀ values of 31 and 27 nM, respectively.² Alobresib (10 and 20 mg/kg, p.o.) reduces tumor growth in a USPC-ARK-2 USC mouse xenograft model.

References

- 1. Kim, E., Hacken, E.T., Sivina, M., et al. The BET inhibitor GS-5829 targets chronic lymphocytic leukemia cells and their supportive microenvironment. Leukemia 34(6), 1588-1598 (2020).
- 2. Bonazzoli, E., Predolini, F., Cocco, E., et al. Inhibition of BET bromodomain proteins with GS-5829 and GS-626510 in uterine serous carcinoma, a biologically aggressive variant of endometrial cancer. Clin. Cancer Res. 24(19), 4845-4853 (2018).

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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