PRODUCT INFORMATION



CID-5721353

Item No. 22967

CAS Registry No.: 301356-95-6

Formal Name: 2-[5-(5-bromo-1,2-dihydro-2-oxo-3H-

> indol-3-ylidene)-4-oxo-2-thioxo-3thiazolidinyl]-butanedioic acid

MF: $C_{15}H_9BrN_2O_6S_2$

457.3 FW: ≥98% **Purity:** Supplied as: A solid Storage: -20°C Stability: ≥4 years

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

Laboratory Procedures

CID-5721353 is supplied as a solid. A stock solution may be made by dissolving the CID-5721353 in the solvent of choice, which should be purged with an inert gas. CID-5721353 is soluble in organic solvents such as acetonitrile and DMSO.

Description

CID-5721353 is an inhibitor of the transcriptional repressor B cell lymphoma 6 (Bcl-6; $K_i = 147 \mu M$). It is selective for Bcl-6 over Kaiso, hypermethylated in cancer 1 (HIC1), and promyelocytic leukemia zinc finger (PLZF) at 50 μM. CID-5721353 is cytotoxic against Bcl-6-dependent SU-DLH-6, SU-DLH-4, Farage, OCI-LY7, OCI-LY1, and OCI-LY10 diffuse large B cell lymphoma (DLBCL) cells (GI_{50} s = 0.024-0.936 mM) but not Bcl-6-independent Toledo or OCI-LY4 DLBCL cells (GI₅₀ = >15 mM for both). It increases expression of the Bcl-6 target genes ATR, TP53, CD69, p21, and CD44 in SU-DLH-6 and SU-DHL-4 cells when used at a concentration of 50 μM. CID-5721353 (50 mg/kg) reduces tumor growth in OCI-LY7 and SU-DHL-6 mouse xenograft models.

Reference

1. Cerchietti, L.C., Ghetu, A.F., Zhu, X., et al. A small-molecule inhibitor of BCL6 kills DLBCL cells in vitro and in vivo. Cancer Cell 17(4), 400-411 (2010).

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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