PRODUCT INFORMATION



YCH1899

Item No. 39737

Formal Name: 4-(3-(3-(5-bromofuran-2-yl)-

> 5,6,7,8-tetrahydro-[1,2,4] triazolo[4,3-a]pyrazine-7carbonyl)-4-fluorobenzyl) phthalazin-1(2H)-one

MF: C₂₅H₁₈BrFN₆O₃

FW: 549.4 **Purity:** ≥98% Supplied as: A solid Storage: -20°C Stability: ≥3 years

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

Laboratory Procedures

YCH1899 is supplied as a solid. A stock solution may be made by dissolving the YCH1899 in the solvent of choice, which should be purged with an inert gas. YCH1899 is soluble in DMSO.

Description

YCH1899 is an inhibitor of poly(ADP-ribose) polymerase 1 (PARP1) and PARP2 ($IC_{50}s = <0.001$ nM for both).¹ It is selective for PARP1 and PARP2 over PARP3, -4, -5A, -5B, -6, -7, -10, and -12 $(IC_{50}s = 1-14.1 \text{ nM})$. YCH1899 inhibits the proliferation of Capan-1 cells ($IC_{50} = 0.1 \text{ nM}$), as well as Capan-1 cells resistant to the PARP inhibitors talazoparib (BMN 673; Item No. 19782) or olaparib (Item No. 10621; $IC_{50}s = 1.13$ and 0.89 nM, respectively). It inhibits the proliferation of BRCA1 mutant HCC1937 breast cancer cells (IC₅₀ = 4.54 nM). In vivo, YCH1899 (12.5 and 25 mg/kg) reduces tumor growth in an olaparib-resistant MDA-MB-436 breast cancer mouse xenograft model.

Reference

1. Sun, Y., Yang, H., Yuan, J., et al. YCH1899, a highly effective phthalazin-1(2H)-one derivative that overcomes resistance to prior PARP inhibitors. J. Med. Chem. 66(17), 12284-12303 (2023).

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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