ABT-263

Small Molecules

B cell lymphoma 2 family inhibitor



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Catalog #100-1173 25 mg

Product Description

ABT-263 is a chemotherapeutic agent and an inhibitor of B cell lymphoma 2 (Bcl-2), B cell lymphoma-extra large (Bcl-xL), and Bcl-2-like protein 2 (Bcl-W) ($K_i = < 1 \text{ nM}, < 0.5$, and < 1 nM, respectively). Bcl-2 and Bcl-xL are anti-apoptotic proteins that regulate cell death by interacting with pro-apoptotic proteins like Bcl-2-like protein 11 (BlM) to induce or inhibit apoptosis (Tse et al.). ABT-263 displaces Bcl-xL from binding to BlM to allow cells to undergo apoptosis (Lagares et al.).

Alternative Names: Navitoclax

CAS Number: 923564-51-6

Chemical Formula: $C_{47}H_{55}CIF_3N_5O_6S_3$ Molecular Weight: 974.6 g/mol

Purity: \geq 98%

Chemical Name: 4-[4-[[2-(4-chlorophenyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl]-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl-1-piperazinyl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]methyl-1-piperazinyl-1-yl]-N-[[4-[[(1R)-3-(4-morpholinyl)-5,5-dimethyl-1-cyclohexen-1-yl]-N-[[4-[[(1R)-3-(4-morpholinyl)-6,5-dimethyl-1-cyclohexen-1-yl]-N-[[4-[[4-[[4-[(1R)-4-(4-[4-[4-[4-[4-[4-[4

1-[(phenylthio)methyl]propyl]amino]-3-[(trifluoromethyl)sulfonyl]phenyl]sulfonyl]-benzamide

Structure:

Properties

Physical Appearance: A white powder

Storage: Product stable at -20°C as supplied. As a precaution, STEMCELL recommends storing all small molecules away

from direct light. For long-term storage, store with a desiccant. Stable as supplied for 12 months from date of

receipt.

Solubility: • DMSO \leq 25 mM

For example, to prepare a 10 mM stock solution in DMSO, resuspend 10 mg in 1.03 mL of DMSO.

Prepare stock solution fresh before use. Information regarding stability of small molecules in solution has rarely been reported; however, as a general guide we recommend storage in DMSO at -20°C. Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

Compound has low solubility in aqueous media. For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. Avoid final DMSO concentration above 0.1% due to potential

cell toxicity.

Small Molecules ABT-263



Published Applications

IMMUNOLOGY

- · Reduces expression of mitochondrial antiviral signaling protein (MAVS) in macrophages (Kim et al.).
- · Induces apoptosis in influenza A virus-infected cells by activating the caspase 9 mediated mitochondrial apoptosis pathway (Kakkola et al.).
- · Inhibits tumor growth in small cell lung cancer xenograft models (Aguilar et al.).
- · Acts as a selective senolytic in vitro by inducing apoptosis in senescent tumor cells after induction into senescence by chemotherapy or radiation (Saleh et al.).
- · Promotes apoptosis in senescent murine Lewis lung carcinoma (LLC) cells (Saleh et al.).
- · Induces G1/G0-phase arrest, apoptosis, and autophagy in human esophageal cancer cells (EC109, HKESC-2, and CaES-17) in vitro (Lin et al.).

References

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Lagares D et al. (2017) Targeted apoptosis of myofibroblasts with the BH3 mimetic ABT-263 reverses established fibrosis. Sci Transl Med 9(420): eaal3765.

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Tse C et al. (2008) ABT-263: a potent and orally bioavailable Bcl-2 family inhibitor. Cancer Res 68(9): 3421-8.

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