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Inhibitors, Screening Libraries, Proteins

Survivin

Survivin is a member of the inhibitor of apoptosis (IAP) family. The survivin protein functions to inhibit caspase activation, thereby leading to negative regulation of apoptosis or programmed cell death. This has been shown by disruption of survivin induction pathways leading to increase in apoptosis and decrease in tumour growth. Survivin expression is highly regulated by the cell cycle and is only expressed in the G2-M phase. Survivin localizes to the mitotic spindle by interaction with tubulin during mitosis and may play a contributing role in regulating mitosis. Survivin is highly expressed in most cancers and associated with chemotherapy resistance, increased tumor recurrence, and shorter patient survival, making antisurvivin therapy an attractive cancer treatment strategy.

Survivin Inhibitors & Antagonists

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| <p>Cucurbitacin IIa (Hemslecin A)</p> <p>Cucurbitacin IIa is a triterpene isolated from <i>Hemsleya amalilis</i> Diels, induces apoptosis of cancer cells, reduces expression of survivin, reduces phospho-Histone H3 and increases cleaved PARP in cancer cells.</p> <p>Purity: 99.27% Clinical Data: No Development Reported Size: 5 mg, 10 mg, 20 mg</p> | <p>GDP366</p> <p>GDP366, a dual inhibitor of survivin and Op18, induces cell growth inhibition, cellular senescence and mitotic catastrophe in human cancer cells.</p> <p>Purity: 99.73% Clinical Data: No Development Reported Size: 5 mg, 10 mg, 25 mg, 50 mg, 100 mg</p> |
| <p>Isolinderalactone</p> <p>Isolinderalactone suppresses human glioblastoma growth and angiogenic activity through the inhibition of VEGFR2 activation in endothelial cells. Isolinderalactone suppresses the expression of B-cell lymphoma 2 (Bcl-2), survi.</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 5 mg, 10 mg, 25 mg</p> | <p>LQZ-7I</p> <p>LQZ-7I is a survivin-targeting inhibitor. LQZ-7I inhibits survivin dimerization. LQZ-7I orally effectively inhibits xenograft tumor growth and induces survivin loss in tumors.</p> <p>Purity: 99.81% Clinical Data: No Development Reported Size: 10 mM × 1 mL, 5 mg, 10 mg, 25 mg, 50 mg</p> |
| <p>Sepantronium bromide (YM-155)</p> <p>Sepantronium bromide (YM-155) is a survivin inhibitor with an IC_{50} of 0.54 nM.</p> <p>Purity: 98.91% Clinical Data: Phase 2 Size: 10 mM × 1 mL, 5 mg, 10 mg, 50 mg, 100 mg</p> | <p>Sepantronium hydrochloride (YM-155 hydrochloride)</p> <p>Sepantronium hydrochloride (YM-155 hydrochloride) is a novel survivin suppressant with an IC_{50} of 0.54 nM for the inhibition of survivin promoter activity.</p> <p>Purity: >98% Clinical Data: Phase 2 Size: 1 mg, 5 mg</p> |
| <p>Shepherdin (79-87)</p> <p>Shepherdin (79-87) is amino acids 79 to 87 fragment of Shepherdin. Shepherdin is a peptidomimetic antagonist of the complex between Hsp90 and Survivin. Anticancer activity.</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg</p> | |