

TRAIL Receptor DR5 Activator – Bioymifi

Chemical Name: (Z)-5-(5-((3-(4-bromophenyl)-2-imino-4-oxothiazolidin-5-ylidene)methyl)furan-2-yl)isoindoline-1,3-dione



Molecular Weight:	494.32
Formula:	C ₂₂ H ₁₂ BrN ₃ O ₄ S
Purity:	≥97%
CAS#:	1420071-30-2
Solubility:	DMSO up to 100 mM
Storage	Powder: 4°C 1 year
	DMSO: 4°C 3 month
	-20°C 1 year

Biological Activity:

Bioymifi is a potent and selective small molecule agonist of DR5, identified by a high-throughput chemical screening for compounds that promote cell death in synergy with a small-molecule mimetic of Smac. Bioymifi directly targets DR5, specifically binds the ECD of DR5 (Kd \sim 1.2 μ M), and induces the formation of DR5 aggregates and DR5 activation. Bioymifi induces caspase-8–dependent apoptosis, which occurs through a DR5-dependent extrinsic pathway but independent of TRAIL. Bioymifi is capable of acting as a single agent to induce DR5 clustering and aggregation, leading to apoptosis without the need for a Smac mimetic in a variety of cancer cell lines, even in U2OS and HT29 cell lines. Bioymifi could be a potential lead compound for the development of small-molecule TRAIL mimics targeting DR5 for cancer therapy.

How to Use:

In vitro: Bioymifi was used at 5-10 µM final concentration in vitro and in cellular assays.

In vivo: n/a

Reference:

1. Wang G, et al. Small-molecule activation of the TRAIL receptor DR5 in human cancer cells. (2012) Nat Chem Biol. 9(2):84-89.

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