

SL-54. Inhibitors of Menin-MLL Interaction

Chromosomal translocations in the mixed lineage leukemia gene (MLL) are observed in many patients diagnosed with acute myeloid leukemia [1]. A potential therapeutic strategy for MLL1-dependant leukemia is to block the interaction between menin and MLL using small molecule ligands [1]. Discovery of drug-like, high-affinity small molecule inhibitors of the menin–MLL interaction with an appropriate pharmacological profile for clinical development is a significant challenge [2]. MI-503 (below) is a recently reported lead compound demonstrating substantial *in vivo* efficacy in animal models [2].

A 2D similarity search through ASINEX's compound collection has resulted in several thienopyrimidine-based analogs, close analogs of MI-503. Such analogs could be interesting chemical probes for MLL-directed research.



Signature Library 54

Formats	Supplementary Information
80 compounds per plate	SL#54_Menin-MLL.sdf
0.1 mg; 1 mg; 2 mg dry film/powder	
0.1 μmol; 1 μmol DMSO solutions	

References:

1. *Exp Hematol.* 2014 December; 42(12): 995–1012. doi:10.1016/j.exphem.2014.09.006

2. Cancer Cell 27, 589-602; April 13, 2015. 10.1016/j.ccell.2015.02.016

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