

## **SL-04. Oncology Phenotypic Screening**

Deregulation of the balance between cell proliferation and death can lead to the formation of tumor malignancies. Signaling transduction pathways can regulate this process via the interaction of multiple functional extracellular and intracellular proteins such as kinases (e.g. EGRF, VEGF, Aurora A). Targeting these proteins leads to cytotoxic activity against various cells of tumor origin.

The ASINEX oncology-oriented phenotypic library includes compounds based on several scaffolds with proven cytotoxicity (sub- $\mu$ M EC<sub>50</sub> in MTT assay) against a number of

cancer cell lines: MV-411 (acute monocytic leukemia), HCT-116 (human colon cancer), MCF-7 (human breast adenocarcinoma), A-172 (Human glioblastoma), COLO-320 (colorectal adenocarcinoma), U-937 (histiocytic lymphoma), A-375 (malignant melanoma), BXPC-3 (pancreas adenocarcinoma), U118-MG (malignant glioma), and LN-229 (glioblastoma). Some active compounds were screened in a biochemical assay showing nM level of IC<sub>50</sub> against several kinases: Aurora A, Haspin, VEGFR, EGFR, PDGFR.



## Signature Library 04

Formats	Supplementary Information
80 compounds per plate	EC <sub>50</sub> MTT test
0.1 mg; 1 mg; 2 mg dry film/powder	Solubility data in PBS
0.1 µmol; 1 µmol DMSO solutions	SL#4_MTT_04-16.sdf

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