

http://www.ime.fraunhofer.de/en/businessareasMB/screeningport.html

Germany

high-capacity screening site

Hamburg

eu :: openscreen an official partner site of









THE PEOPLE

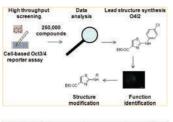
Dr. Sheraz Gul (sheraz.gul@ime.fraunhofer.de)

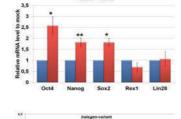
Dr. Bernhard Ellinger (bernhard.ellinger@ime.fraunhofer.de)

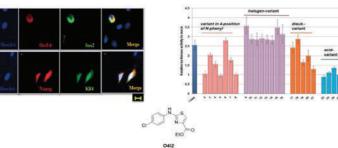
Dr. Björn Windshügel (bjoern.windshuegel@ime.fraunhofer.de)

THE PROJECTS

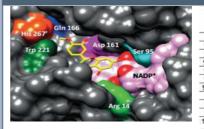
drug-iPS (BMBF Systems Biology, MedSys)

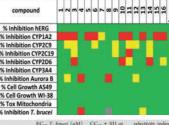


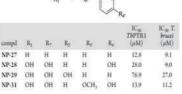


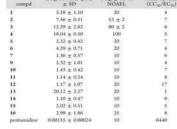


NMTrypl (EU FP7)





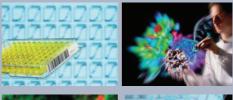




THE HARDWARE

- 1. Label free screening including multimode capacity (Enspire EPIC technology, PerkinElmer Inc), SPR screening (384 well, MASS-1, Sierra Sensors), live cell imaging (IncuCyte ZOOM, Essen Instruments), single molecule spectroscopy diffusion and fluorescence intensity analysis (Clarina, Evotec AG), ion channel and transporter protein screening in cell free formats (longate, Nanion), brightfield and fluorescence cell imaging in multimode format (EnSight, PerkinElmer), multimode capacity (EnVision, PerkinElmer Inc) and HCS (Opera, PerkinElmer Inc).
- 2. AlphaScreen, TR-FRET, fluorescence intensity, confocal high content imaging, absorbance, luminescence, reporter gene, cellular biosensors, caspase induction, cAMP, immunocytochemistry, cell migration, translocation, ion sensing (FluxOR, Fluo-4 and Fura-2), ELISA, Proximity Ligation Analysis, cell free electrophysiology and fluorescence polarisation.
- 3. Kinase, GPCR, ion channel, HDAC, PDE, deubiquitinylase, PPI, transporter, efflux pump, protease, nuclear receptor, DNA binding protein and toxicity (CYP450, gene toxicity, hERG, mitochondrial toxicity).
- 4. LOPAC, ENZO, Enamine, Evotec, MRCT, HYPHA Discovery, CHEMBRIDGE, Sanofi, MMV and Kailash.

THE SOFTWARE







THE OUTPUT

- 1. ETHYL 2-(I4-CHLOROPHENYL)AMINO)THIAZOLE-4-CARBOXYLATE (O4I2) AND DERIVATIVES ARE POTENT INDUCERS OF OCT3/4. Cheng, et al. Journal of Medicinal Chemistry, 2015, 58, 5742-5750.

 2. IDENTRICATION OF 2-I4-I(I4-METHOXYPHENYLIMETHOXY)-PHENYL]ACETONITRILE (O4I1) AND DERIVATIVES AS POTENT OCT3/4 INDUCERS. Cheng, et al. Journal of Medicinal Chemistry, 2015, 58, 4976-4983.

 3. PROFILING OF FLAVONOL DERIVATIVES FOR THE DEVELOPMENT OF ANTI-TRYPANOSOMATIDIC DRUGS. Borsari, et al. Journal of Medicinal Chemistry, 2015, 98, 7958-7616.

 4. ETHOXYLATED 2-HYDROXYCHALCONES AS ANTIPARASTIC HIT COMPOUNDS. Borsari, et al. European Journal of Medicinal Chemistry, 2017, 126, 1129-1135.

 5. CHROMAN-4-ONE DERIVATIVES TARGETING PTERIDINE REDUCTASE 1 AND SHOWING ANTI-PARASTIC ACTIVITY. DIPISA et al. Molecules, 2017, 22, 1426.

ative Medicines Initiative - K4DD, ND4RR-TRANSI OCATION and FRISC

THE FUTURE

2017: 3 small. 2 medium and 2 large screens.

2022: 4-5 small, 4-5 medium and 3 large screens

iPS cell technologies for drug discovery, including access to patient derived cell lines for in-vitro disease models.

2. Structure based drug design & virtual screening



Fraunhofer

