CONFIDENTIAL, contact the Imaging Platform to collaborate on the findings herein MAPK1_WT.2 - in Canonical MAPK How similar is this gene to the other genes? 0.9 0.4 Correlation of the gene to the other genes 0.3 RBPJ_WT.2 XBP1_WT.2 CASP8_WT.1 JUN_WT.2 GLI1_WT PPARGC1A_1 PRKACG 0.0 TGFBR1_K232R
STK3_WT.2

KRAS_WT.1

STK3_WT.1

STK3_WT.1

STK3_WT.2

DVL3_WT AKT3_WT.3
JAG1_WT
DDIT4_WT
MAP3K7_WT

What groups of morphological features are distinguishing in the cluster relative to the untreated samples? (maximum of absolute m-score for the features belonging to the same category; m-score defined as median of a feature z-score across genes in the cluster) Black means no feature is available in the category

AreaShape -Cells Cytoplasm Nuclei Texture -Intensity 10 RadialDistribution DNA AĞP ER RNA Mito

Correlation

between

compound

the gene

scored

against the

gene using

L1000

profiling

correlation of the

compound signature

(95th DMSO

replicate correlation

is 0.51)

Chemical

structure

RNA

ER

common names (where

available); blue/red colored

box means the matching

compound is

positively/negatively

correlated with the cluster

Which individual morphological features are distinguishing in the gene relative to the untreated samples? Blue/Red means the feature has a positive/negative z-score. Size is proportional to the z-score value.

Distinguishing individual features for the compound relative to Number of PubChem assays in which

untreated samples. Black means a mismatch; i.e. active (= high

z-score in magnitude) in the compound, and either inactive (=

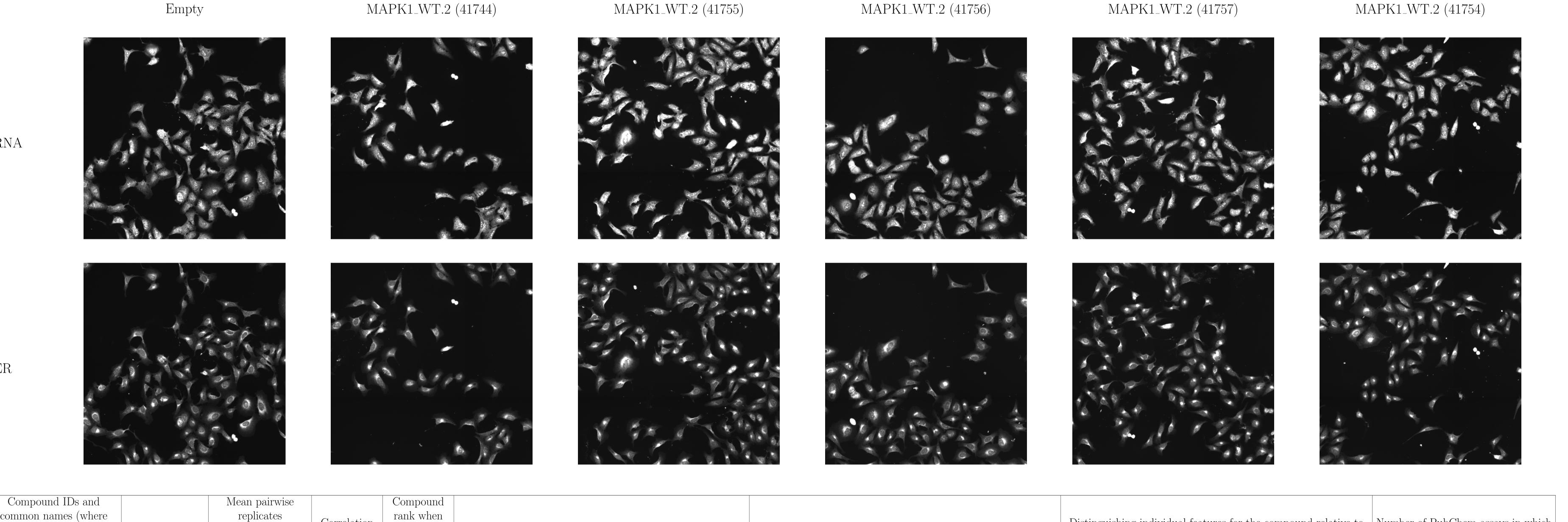
small z-score in magnitude) or oppositely active in the gene

the compound was tested; assays in

which the compound was active are

itemized





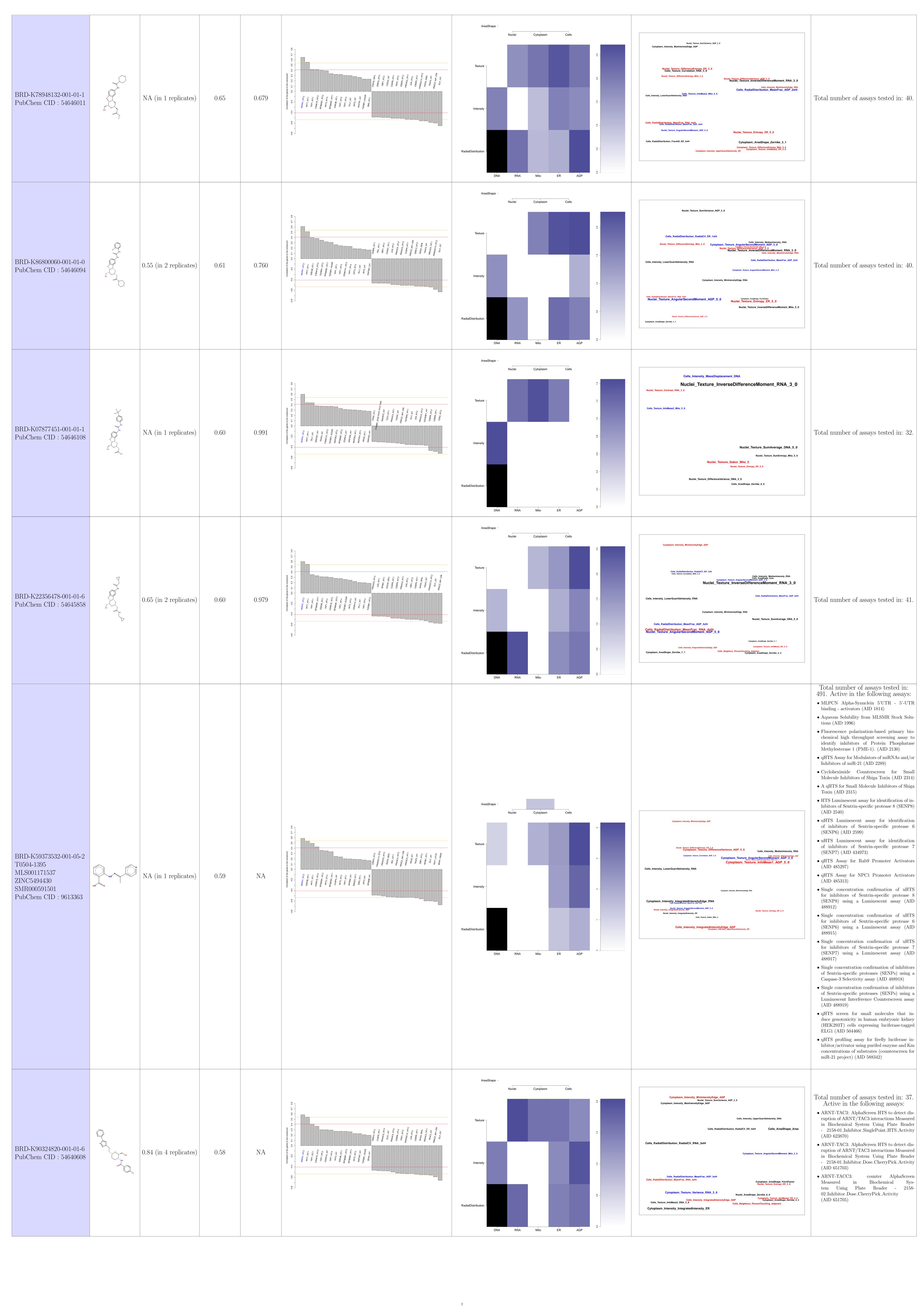
Common distinguishing feature categories in the compound and

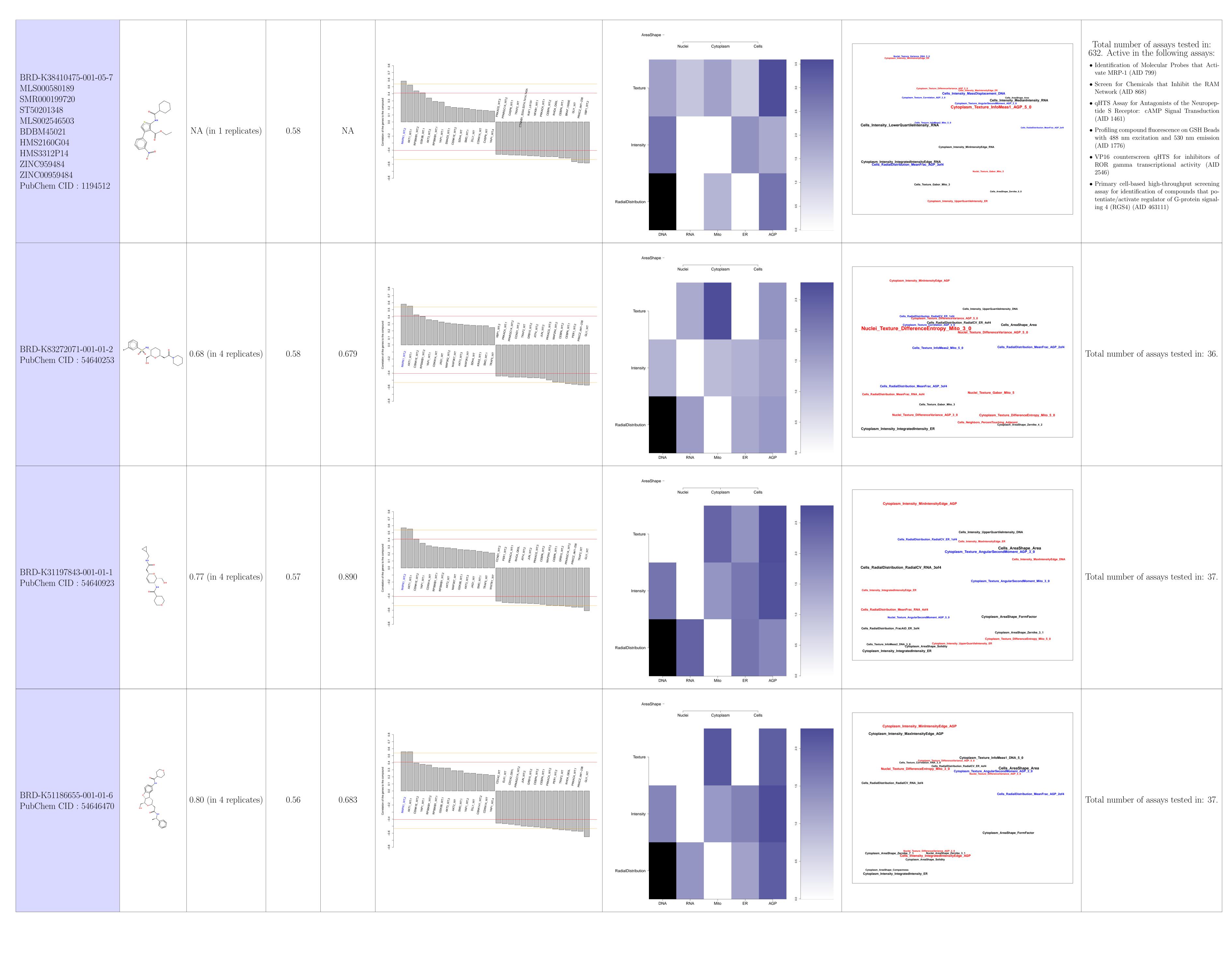
the gene relative to the untreated samples

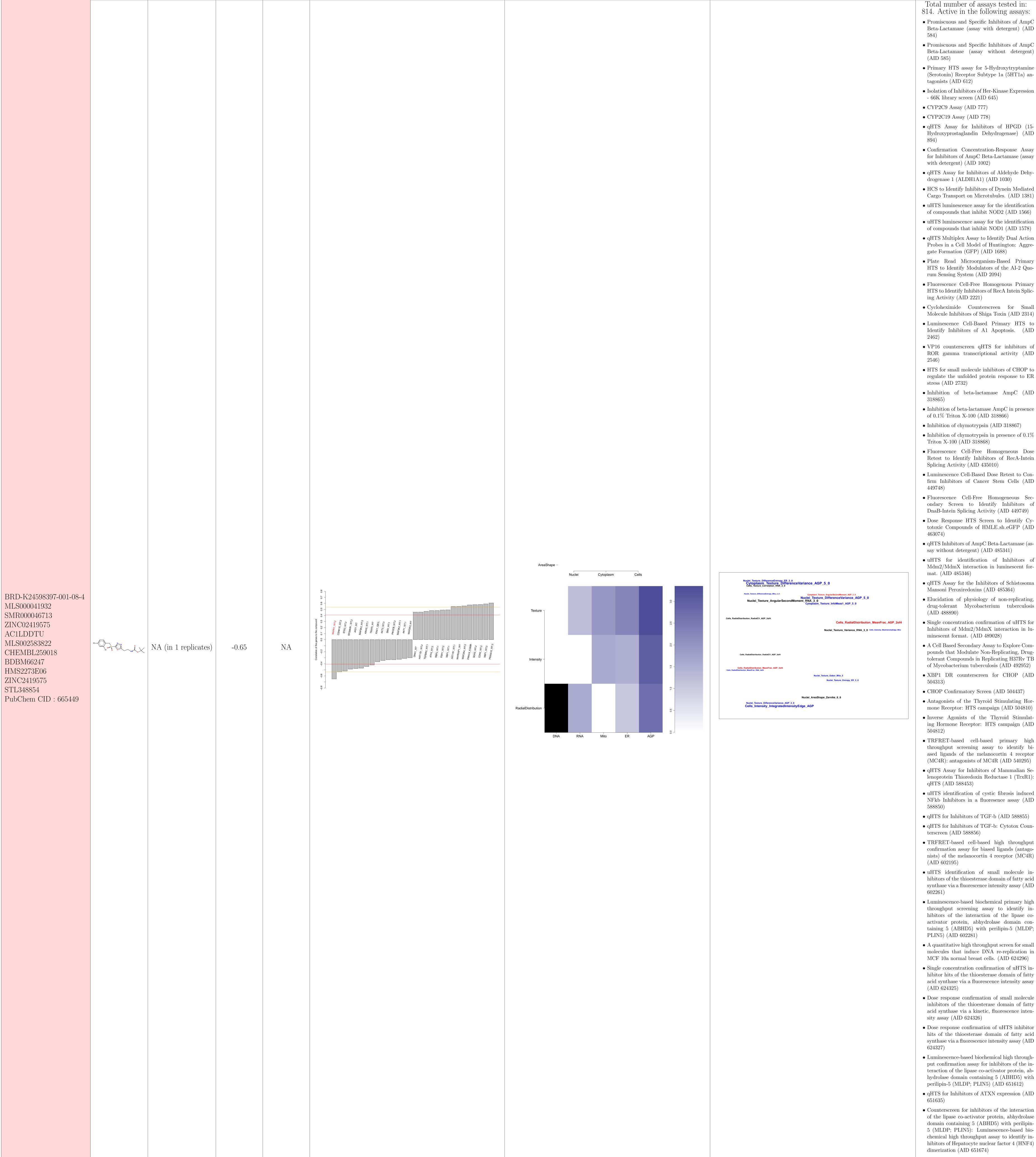
How similar is the compound signature to the genes in this

experiment? (Yellow and red lines correspond to top/bottom

1st and 5th percentile DMSO correlation to all the genes)







814. Active in the following assays: • Promiscuous and Specific Inhibitors of AmpC

Beta-Lactamase (assay with detergent) (AID

Beta-Lactamase (assay without detergent) • Primary HTS assay for 5-Hydroxytryptamine

(Serotonin) Receptor Subtype 1a (5HT1a) antagonists (AID 612)

• Isolation of Inhibitors of Her-Kinase Expression - 66K library screen (AID 645)

• CYP2C9 Assay (AID 777)

• CYP2C19 Assay (AID 778) • qHTS Assay for Inhibitors of HPGD (15-

Hydroxyprostaglandin Dehydrogenase) (AID

with detergent) (AID 1002)

• qHTS Assay for Inhibitors of Aldehyde Dehydrogenase 1 (ALDH1A1) (AID 1030)

• HCS to Identify Inhibitors of Dynein Mediated Cargo Transport on Microtubules. (AID 1381) • uHTS luminescence assay for the identification

of compounds that inhibit NOD2 (AID 1566) • uHTS luminescence assay for the identification of compounds that inhibit NOD1 (AID 1578)

• qHTS Multiplex Assay to Identify Dual Action Probes in a Cell Model of Huntington: Aggregate Formation (GFP) (AID 1688)

• Plate Read Microorganism-Based Primary HTS to Identify Modulators of the AI-2 Quorum Sensing System (AID 2094)

HTS to Identify Inhibitors of RecA Intein Splicing Activity (AID 2221) • Cycloheximide Counterscreen for Small

Molecule Inhibitors of Shiga Toxin (AID 2314) • Luminescence Cell-Based Primary HTS to Identify Inhibitors of A1 Apoptosis. (AID

• VP16 counterscreen qHTS for inhibitors of ROR gamma transcriptional activity (AID

• HTS for small molecule inhibitors of CHOP to regulate the unfolded protein response to ER

• Inhibition of beta-lactamase AmpC (AID

• Inhibition of beta-lactamase AmpC in presence of 0.1% Triton X-100 (AID 318866)

• Inhibition of chymotrypsin (AID 318867) • Inhibition of chymotrypsin in presence of 0.1%

Triton X-100 (AID 318868) • Fluorescence Cell-Free Homogeneous Dose Retest to Identify Inhibitors of RecA-Intein

• Luminescence Cell-Based Dose Retest to Confirm Inhibitors of Cancer Stem Cells (AID

• Fluorescence Cell-Free Homogeneous Secondary Screen to Identify Inhibitors of DnaB-Intein Splicing Activity (AID 449749)

• Dose Response HTS Screen to Identify Cytotoxic Compounds of HMLE_sh_eGFP (AID

say without detergent) (AID 485341) • uHTS for identification of Inhibitors of Mdm2/MdmX interaction in luminescent for-

mat. (AID 485346) • qHTS Assay for the Inhibitors of Schistosoma

• Elucidation of physiology of non-replicating, drug-tolerant Mycobacterium tuberculosis

• Single concentration confirmation of uHTS for

Inhibitors of Mdm2/MdmX interaction in luminescent format. (AID 489028) • A Cell Based Secondary Assay to Explore Com-

pounds that Modulate Non-Replicating, Drugtolerant Compounds in Replicating H37Rv TB of Mycobacterium tuberculosis (AID 492952) • XBP1 DR counterscreen for CHOP (AID

• Antagonists of the Thyroid Stimulating Hormone Receptor: HTS campaign (AID 504810)

• Inverse Agonists of the Thyroid Stimulating Hormone Receptor: HTS campaign (AID

throughput screening assay to identify biased ligands of the melanocortin 4 receptor (MC4R): antagonists of MC4R (AID 540295)

• qHTS Assay for Inhibitors of Mammalian Selenoprotein Thioredoxin Reductase 1 (TrxR1):

qHTS (AID 588453) • uHTS identification of cystic fibrosis induced

• qHTS for Inhibitors of TGF-b (AID 588855)

• qHTS for Inhibitors of TGF-b: Cytotox Coun-

• TRFRET-based cell-based high throughput confirmation assay for biased ligands (antago-

nists) of the melanocortin 4 receptor (MC4R)

• uHTS identification of small molecule inhibitors of the thioesterase domain of fatty acid synthase via a fluorescence intensity assay (AID

• Luminescence-based biochemical primary high throughput screening assay to identify inhibitors of the interaction of the lipase co-

PLIN5) (AID 602281) • A quantitative high throughput screen for small

molecules that induce DNA re-replication in MCF 10a normal breast cells. (AID 624296) • Single concentration confirmation of uHTS inhibitor hits of the thioesterase domain of fatty

• Dose response confirmation of small molecule inhibitors of the thioesterase domain of fatty acid synthase via a kinetic, fluorescence inten-

sity assay (AID 624326) • Dose response confirmation of uHTS inhibitor hits of the thioesterase domain of fatty acid synthase via a fluorescence intensity assay (AID

• Luminescence-based biochemical high throughput confirmation assay for inhibitors of the interaction of the lipase co-activator protein, ab-

hydrolase domain containing 5 (ABHD5) with perilipin-5 (MLDP; PLIN5) (AID 651612) • qHTS for Inhibitors of ATXN expression (AID

• Counterscreen for inhibitors of the interaction of the lipase co-activator protein, abhydrolase domain containing 5 (ABHD5) with perilipin-5 (MLDP; PLIN5): Luminescence-based biochemical high throughput assay to identify inhibitors of Hepatocyte nuclear factor 4 (HNF4)

• qHTS of D3 Dopamine Receptor Antagonist: qHTS (AID 652054)

• qHTS for induction of synthetic lethality in tumor cells producing 2HG: qHTS for the HT-1080-NT fibrosarcoma cell line (AID 686970) • qHTS for induction of synthetic lethality in tumor cells producing 2HG: qHTS for the HT-

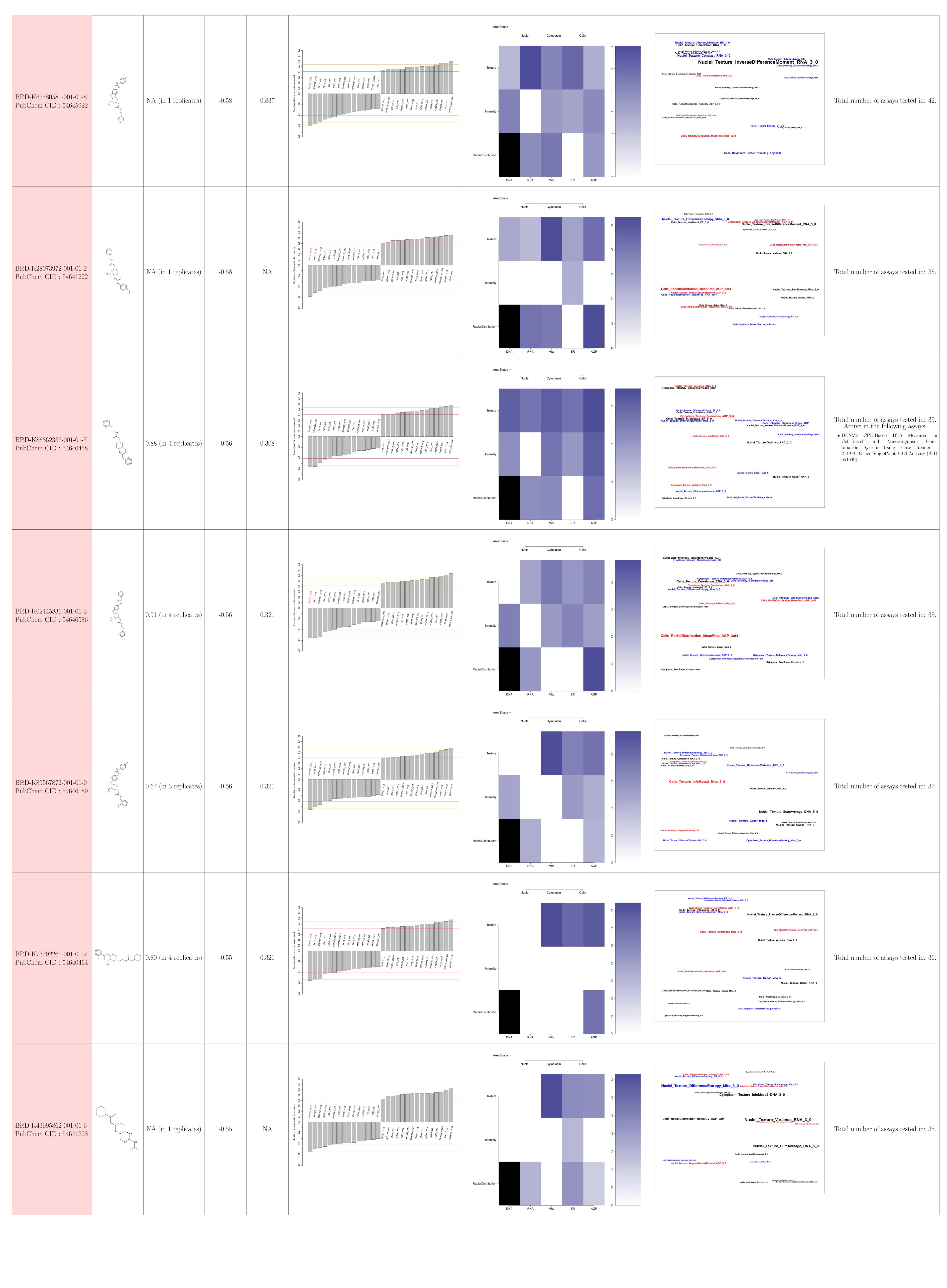
• qHTS for Inhibitors of human tyrosyl-DNA phosphodiesterase 1 (TDP1): qHTS in cells in absence of CPT (AID 686978)

1080-IDH1KD cell line (AID 686971)

• qHTS for Inhibitors of human tyrosyl-DNA phosphodiesterase 1 (TDP1): qHTS in cells in presence of CPT (AID 686979)

• qHTS for Inhibitors of Inflammasome Signaling: IL-1-beta AlphaLISA Primary Screen

(AID 743279)



BRD-K35413741-001-01-1 PubChem CID: 54640338	NH O H	0.67 (in 4 replicates)	-0.54 0.039	S	AreaShape - Nuclei Cytoplasm Cells Congliser Interfacty Minimizer Figure AP 2.0 Codes Reductive Management AP 2.0 Codes Reductive Management AP 2.0 Nuclei Texture Difference Variance AP 2.0 Nuclei Texture Internetive Management AP 2.0 Codes Texture Interfacty Management AP 2.0	Total number of assays tested in: 35.
				9 -	RadialDistribution – RadialDistribution – DNA RNA Mito ER AGP RadialDistribution ER AGP RadialDistribution = Cytoplasm_Intensity_Cytoplasm_Int	
BRD-K26108750-001-01-6 PubChem CID: 54640710	N N N N N N N N N N N N N N N N N N N	0.66 (in 4 replicates)	-0.54 0.186	1-08	Nuclei Cytoplasm Colls Nuclei Texture DifferenceEnropy EE 3 0 Colls Texture DifferenceEnropy EE 3 0 Nuclei Texture DifferenceEnropy EE 3 0 Colls Texture DifferenceEnropy EE 3 0 Nuclei Texture DifferenceEnropy EE 3 0 Colls Texture DifferenceEnropy EE 3 0 Nuclei Texture DifferenceEnropy EE 3 0 Nuclei Texture DifferenceEnropy EE 3 0 Colls Texture DifferenceEnropy EE 3 0 Nuclei Texture DifferenceEnropy EE 3 0 Colls Texture DifferenceEnropy EE 3 0 Nuclei Texture DifferenceEnropy EE 3 0 Nucl	Total number of assays tested in: 37.