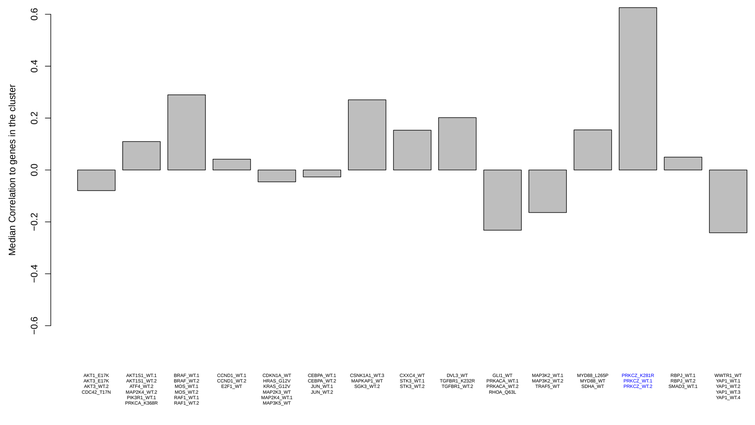
How similar is this cluster to the other clusters?



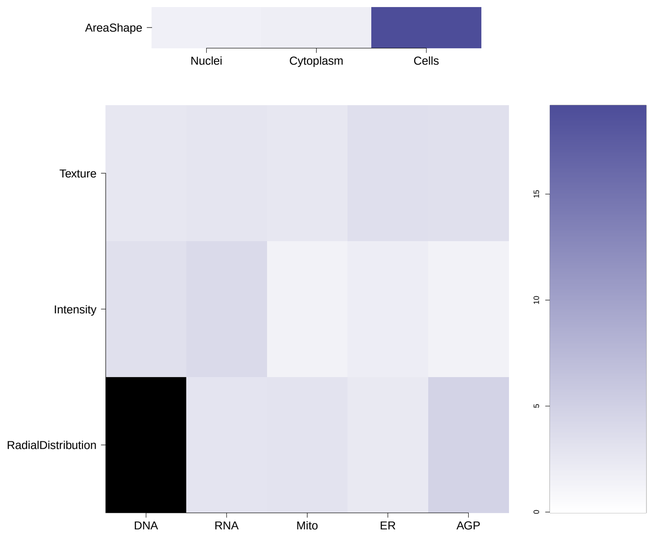
Top 5 genes negatively correlated to the cluster

	Expert Annotation			
Treatment	Pathway	Regulation Type	Mean Correlation	Standard Deviation
WWTR1_WT	Canonical Hippo	Inhibitor	-0.50	0.10
STK11_WT.1	Canonical TOR	Inhibitor	-0.43	0.06
NFKBIB_WT	Canonical NFkB	Inhibitor	-0.40	0.14
TGFB1_WT	Canonical TGFbeta	Activator	-0.35	0.11
DIABLO_WT	Canonical Apoptosis	Inhibitor	-0.33	0.04

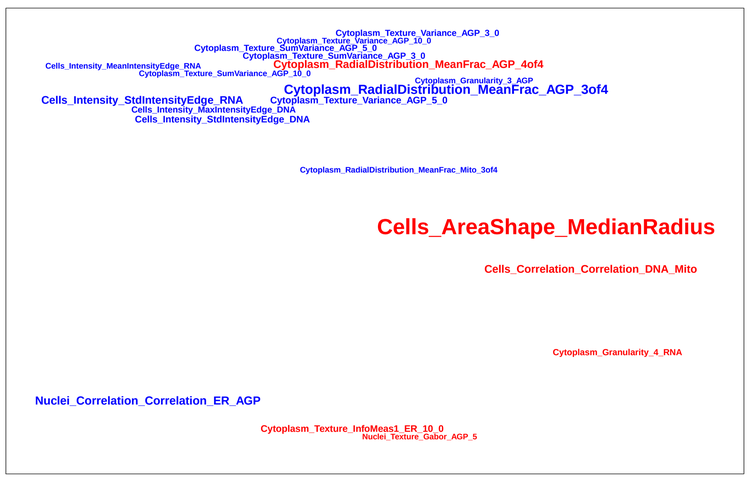
Genes in the cluster along with the pathways as annotated by experts

	Expert Annotation	
Treatment	Pathway	Regulation Type
PRKCZ_WT.1	Canonical PKC	Activator
PRKCZ_WT.2	Canonical PKC	Activator
PRKCZ_K281R	Canonical PKC	Inhibitor

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



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



How strongly are genes within the cluster correlated?

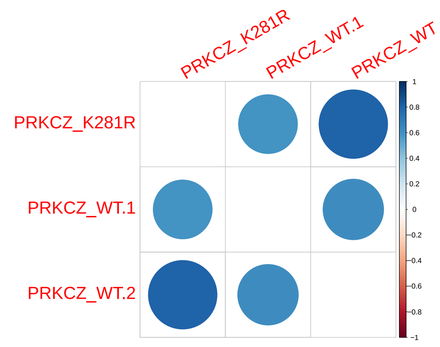
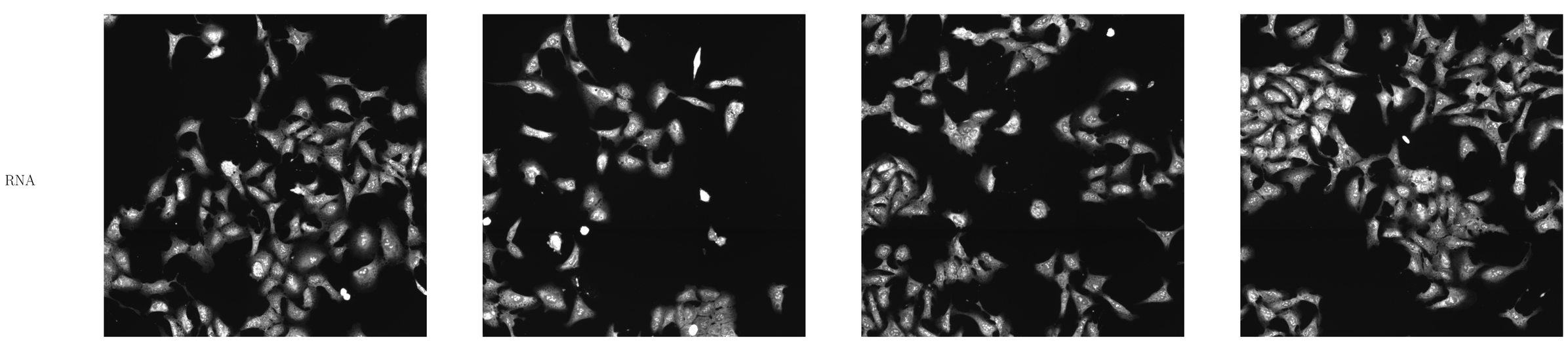
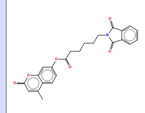
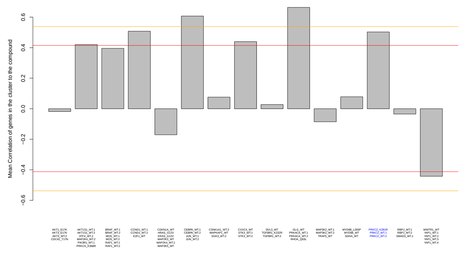
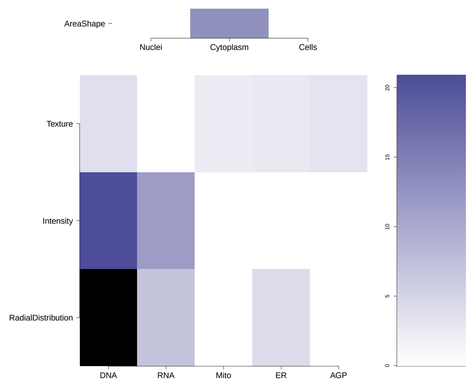
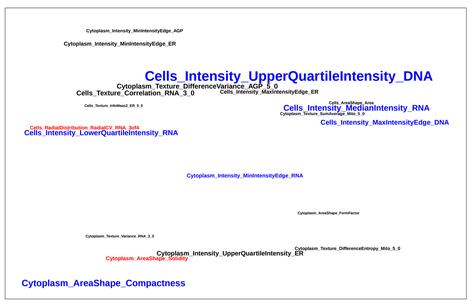
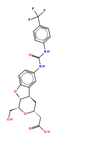
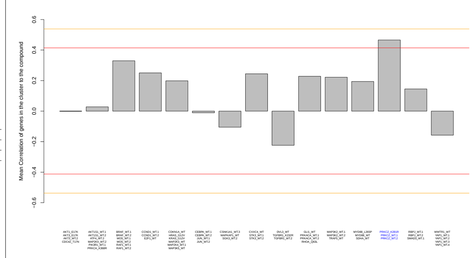
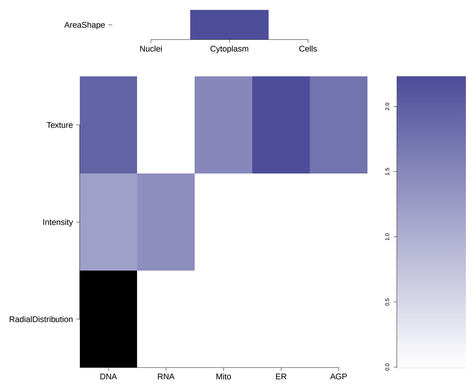
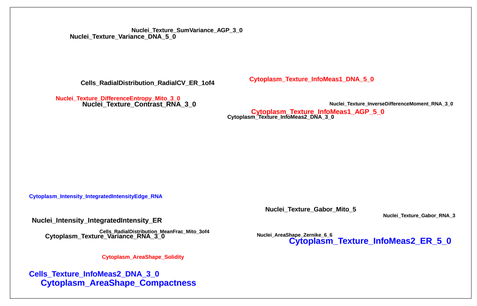
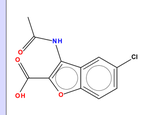
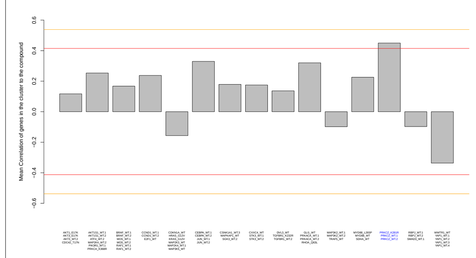
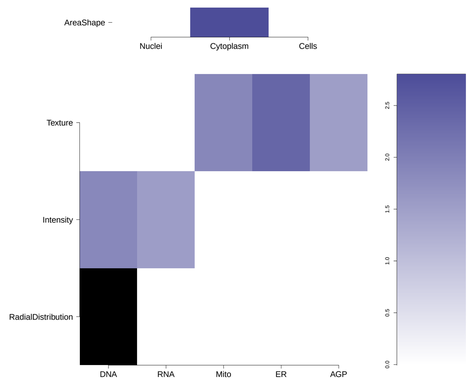
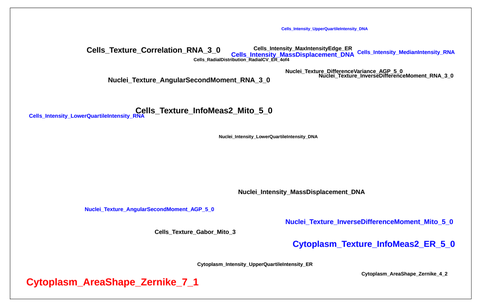
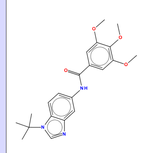
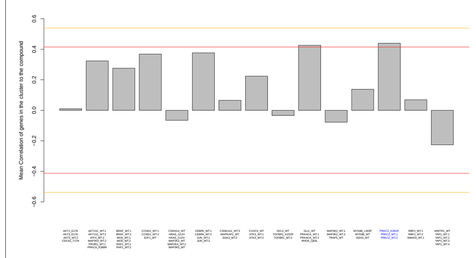
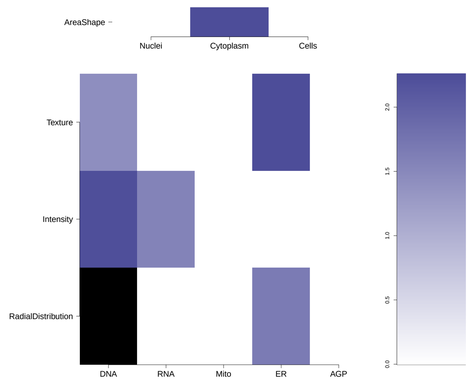
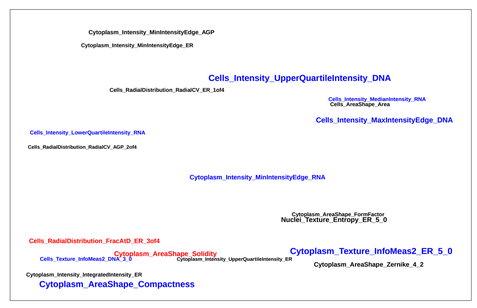
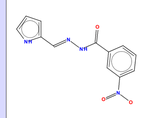
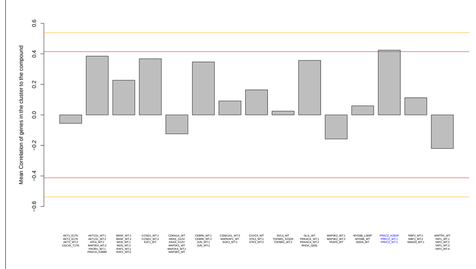
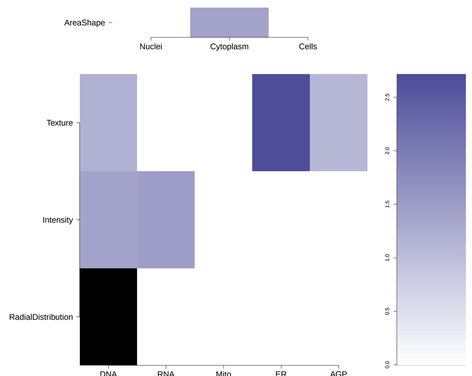
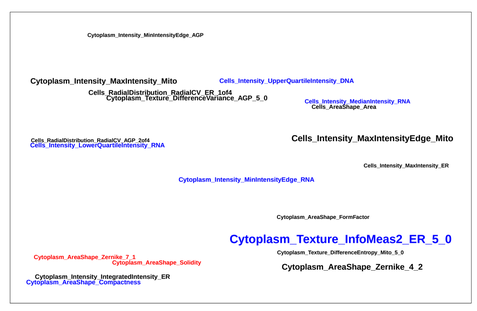
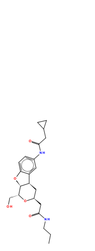
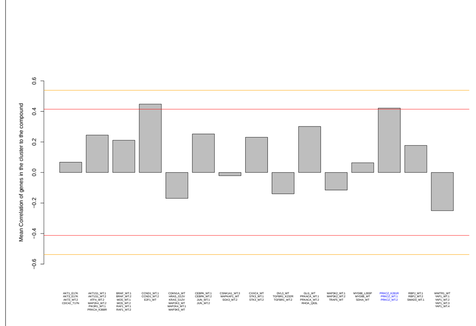
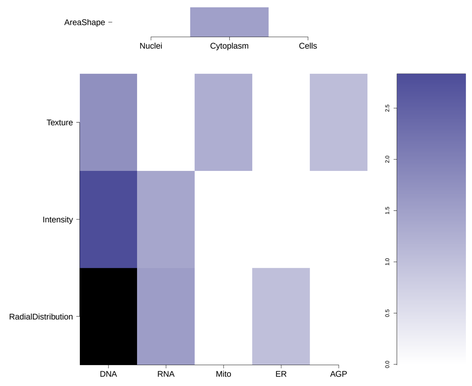
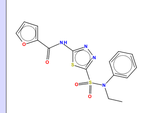
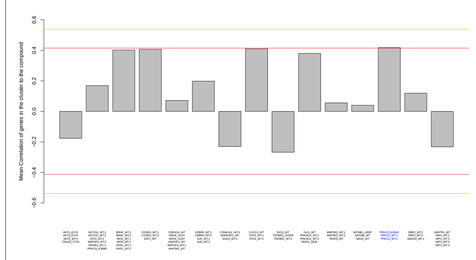
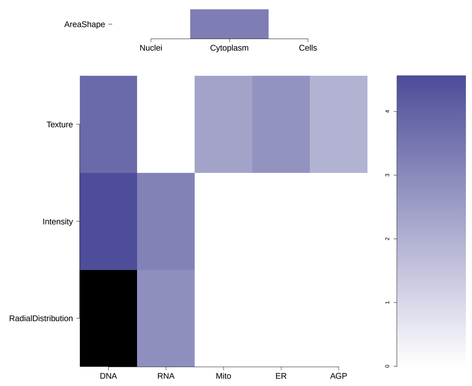
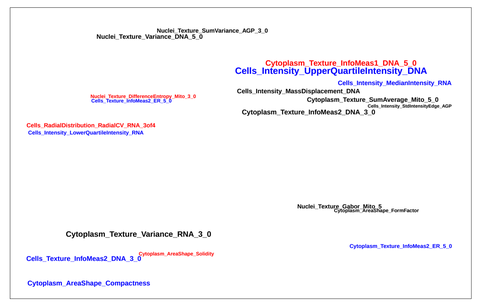


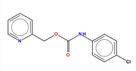
Plate : 41744 - Genes in the Cluster (Channels are sorted based on their dominance in the grid plot)



Compound IDs and common names (where available); blue/red colored box means the matching compound is positively/negatively correlated with the cluster	Chemical structure	Mean pairwise replicates correlation of the compound signature (95th DMSO replicate correlation is 0.52)	Mean ± standard deviation correlation between compound and each gene in cluster; Tables contain data for individual genes	Mean compound rank when scored against genes in cluster using L1000 profiling ± standard deviation; Tables contain data for individual genes	How similar is the compound signature to the gene clusters in this experiment? (Yellow and red lines correspond to top/bottom 1st and 5th percentile DMSO correlation to all the genes)	Common distinguishing feature categories in the compound and genes in the cluster relative to the untreated samples	Distinguishing individual features for the compound relative to 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 cluster	Number of PubChem assays in which the compound was tested; assays in which the compound was active are itemized
BRD-K28901743-001-05-3 ZINC01748812 AC1LTAWC MLS000552933 ZINC1748812 CCG-15676 STL331422 BAS 00558059 SMR000175471 ST50181975 PubChem CID : 1555494		NA (in 1 replicates)	0.50 ± 0.13 Treatment / Score PRKCZ.K281R 0.65 PRKCZ.WT.1 0.40 PRKCZ.WT.2 0.45	NA				Total number of assays tested in: 626. Active in the following assays: • Screen for Chemicals that Extend Yeast Lifespan (AID 775) • uHTS identification of small molecule inhibitors of tim10-1 yeast via a luminescent assay (AID 463190) • Single concentration confirmation of small molecule inhibitors of tim10-1 yeast via a luminescent assay (AID 463213) • Fluorescence-based biochemical primary high throughput screening assay to identify inhibitors of the fructose-bisphosphate aldolase (FBA) of M. tuberculosis (AID 588726) • Fluorescence Intensity-based biochemical primary high throughput screening assay to identify activators of kallikrein-7 (K7) zymogen (AID 652039) • Fluorescence Intensity-based biochemical primary high throughput confirmation assay to identify activators of kallikrein-7 (K7) zymogen (AID 686949) • Counterscreen for activators of kallikrein-7 (K7) zymogen: Fluorescence intensity-based biochemical high throughput counterscreen assay for activators that optically interfere with measurement of EDANS-DABCYL fluorescence (AID 686952)

BRD-K19061245-001-01-9 PubChem CID : 54646112		NA (in 1 replicates)	0.47 ± 0.06 Treatment | Score PRKCZ.K281R | 0.46 PRKCZ.WT.1 | 0.53 PRKCZ.WT.2 | 0.41	0.903 ± 0.104 Treatment | Score PRKCZ.K281R | 0.932 PRKCZ.WT.1 | 0.990 PRKCZ.WT.2 | 0.787				Total number of assays tested in: 39.
BRD-K73953291-001-06-2 MLS000035731 AC1LDESP HMS1675M03 ZINC380820 STK731381 BAS 01507272 SMR000010547 PubChem CID : 646952		NA (in 1 replicates)	0.45 ± 0.07 Treatment | Score PRKCZ.K281R | 0.46 PRKCZ.WT.1 | 0.37 PRKCZ.WT.2 | 0.52	NA				Total number of assays tested in: 774. Active in the following assays: • Primary cell-based high throughput screening assay to measure STAT3 activation (AID 871) • A screen for compounds that inhibit cell wall-associated teichoic acid synthesis in Staphylococcus aureus (AID 463173) • Luminescence Cell-Free Homogeneous Dose Retest to Confirm Inhibitors of GSK-3 alpha (AID 463203) • Primary qHTS for delayed death inhibitors of the malarial parasite plastid, 96 hour incubation (AID 504834)
BRD-K60865828-001-05-7 BAS 05022416 AC1LL2O5 MLS001211773 HMS1615G11 HMS2838A09 ZINC789744 STK090735 ZINC00789744 SMR000523157 VU0417912-1 PubChem CID : 1077764		0.55 (in 4 replicates)	0.44 ± 0.05 Treatment | Score PRKCZ.K281R | 0.50 PRKCZ.WT.1 | 0.40 PRKCZ.WT.2 | 0.42	NA				Total number of assays tested in: 470. Active in the following assays: • Primary cell-based screen for identification of compounds that inhibit the two-pore domain potassium channel KCNK3 (AID 602410) • Confirmation assay for identification of compounds that inhibit the two-pore domain potassium channel KCNK3 [Primary Screening] (AID 651638) • qHTS Assay for Inhibitors of Hepatitis C Virus (HCV) (AID 651820)
BRD-K47430271-001-05-0 SMR000184928 Ambcb5227425 MLS000595187 AC1NX317 HMS2509F11 LS-38042 119034-11-6 PubChem CID : 5719122		0.56 (in 4 replicates)	0.42 ± 0.08 Treatment | Score PRKCZ.K281R | 0.50 PRKCZ.WT.1 | 0.34 PRKCZ.WT.2 | 0.43	NA				Total number of assays tested in: 632. Active in the following assays: • VP16 counterscreen qHTS for inhibitors of ROR gamma transcriptional activity (AID 2546) • qHTS for inhibitors of ROR gamma transcriptional activity (AID 2551) • qHTS Assay for Inhibitors of Hepatitis C Virus (HCV) (AID 651820)
BRD-K23220352-001-01-4 PubChem CID : 54647935		0.55 (in 3 replicates)	0.42 ± 0.05 Treatment | Score PRKCZ.K281R | 0.47 PRKCZ.WT.1 | 0.41 PRKCZ.WT.2 | 0.38	NA				Total number of assays tested in: 37.
BRD-K19608696-001-04-5 MLS001121487 HMS1859H04 HMS2253I17 ZINC6818267 ZINC06818267 SMR000626594 E157-5383 PubChem CID : 16017323		NA (in 1 replicates)	0.42 ± 0.10 Treatment | Score PRKCZ.K281R | 0.50 PRKCZ.WT.1 | 0.44 PRKCZ.WT.2 | 0.31	NA				Total number of assays tested in: 508. Active in the following assays: • qHTS Screen for Compounds that Selectively Target Cancer Cells with p53 Mutations: Cytotoxicity of p53ts Cells at the Nonpermissive Temperature (AID 902) • qHTS Screen for Compounds that Selectively Target Cancer Cells with p53 Mutations: Cytotoxicity of p53 Null Cells at the Nonpermissive Temperature (AID 904) • A screen for compounds that inhibit cell wall-associated teichoic acid synthesis in Staphylococcus aureus (AID 463173) • qHTS Assay for the Inhibitors of Schistosoma Mansoni Peroxiredoxins (AID 485364) • qHTS profiling assay for firefly luciferase inhibitor/activator using purifed enzyme and Km concentrations of substrates (counterscreen for miR-21 project) (AID 588342) • Absorbance-based biochemical primary high throughput screening assay to identify inhibitors of Methionine sulfoxide reductase A (MsrA) (AID 651718) • Absorbance-based biochemical high throughput confirmation assay to identify inhibitors of Methionine sulfoxide reductase A (MsrA) (AID 651822)
BRD-K48818351-001-01-7 PubChem CID : 54641364		NA (in 1 replicates)	-0.56 ± 0.05 Treatment | Score PRKCZ.K281R | -0.56 PRKCZ.WT.1 | -0.61 PRKCZ.WT.2 | -0.51	NA				Total number of assays tested in: 37.

BRD-K05680206-001-06-8
ST51024277
AC1LNZAM
SMR000114553
MLS000550223
HMS2353E22
ZINC984086
ZINC00984086
PubChem CID : 1211462

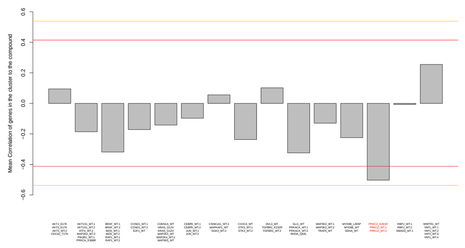


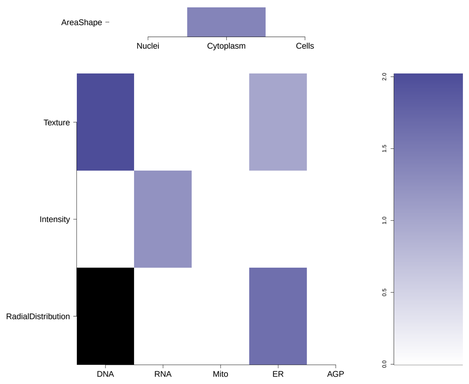
0.80 (in 2 replicates)

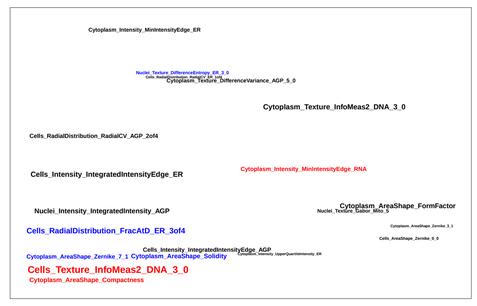
-0.50 ± 0.10

Treatment	Score
PRKCZ.K281R	-0.58
PRKCZ.WT.1	-0.54
PRKCZ.WT.2	-0.39

NA



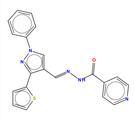




Total number of assays tested in: 679. Active in the following assays:

- Primary cell-based high throughput screening assay to measure STAT1 activation (AID 932)
- qHTS Assay for Enhancers of SMN2 Splice Variant Expression (AID 1458)
- MLPCN Alpha-Synuclein 5'UTR - 5'-UTR binding - activators (AID 1814)
- qHTS Assay for Modulators of miRNAs and/or Inhibitors of miR-21 (AID 2289)
- Cycloheximide Counterscreen for Small Molecule Inhibitors of Shiga Toxin (AID 2314)
- A qHTS for Small Molecule Inhibitors of Shiga Toxin (AID 2315)
- Fluorescence-based counterscreen for orexin 1 receptor (OX1R) antagonists: cell-based assay to identify antagonists of the parental CHO cell line (AID 463079)
- qHTS Assay for Rab9 Promoter Activators (AID 485297)
- qHTS Assay for NPC1 Promoter Activators (AID 485313)
- Heat Shock Factor-1 (HSF-1) Measured in Cell-Based System Using Plate Reader - 2038-01_Activator_SinglePoint_HTS_Activity (AID 504408)
- MITF: Counter assay: A375 proliferation Measured in Cell-Based System Using Plate Reader - 2084-03_Inhibitor_Dose_CherryPick_Activity_Set2 (AID 540335)
- MITF: Counter assay: A375 proliferation Measured in Cell-Based System Using Plate Reader - 2084-03_Inhibitor_Dose_DryPowder_Activity (AID 540346)
- qHTS profiling assay for firefly luciferase inhibitor/activator using purifed enzyme and Km concentrations of substrates (counterscreen for miR-21 project) (AID 588342)
- Screen for inhibitors of the SWI/SNF chromatin remodeling complex (esBAF) in mouse embryonic stem cells with Luciferase reporter assay Measured in Cell-Based System Using Plate Reader - 2141-01_Inhibitor_SinglePoint_HTS_Activity (AID 602393)
- uHTS identification of small molecule inhibitors of the mitochondrial permeability transition pore via an absorbance assay (AID 602449)
- qHTS Assay to Identify Small Molecule Activators of BRCA1 Expression (AID 624202)
- Single concentration confirmation of uHTS inhibitor hits of the mitochondrial permeability transition pore via a fluorescent based assay (AID 624504)
- Wnt/Beta-catenin HTS Measured in Cell-Based System Using Plate Reader - 2161-01_Activator_SinglePoint_HTS_Activity (AID 743398)
- Wnt/Beta-catenin HTS Measured in Cell-Based System Using Plate Reader - 2161-01_Activator_Dose_CherryPick_Activity (AID 1053144)

BRD-K08606983-001-05-1
STK138991
AC1OBPFE
SMR000259826
MLS000390788
ZINC15989497
T0500-2826
PubChem CID : 6900173

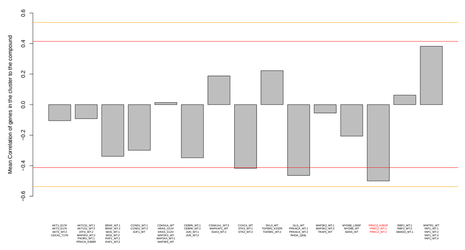


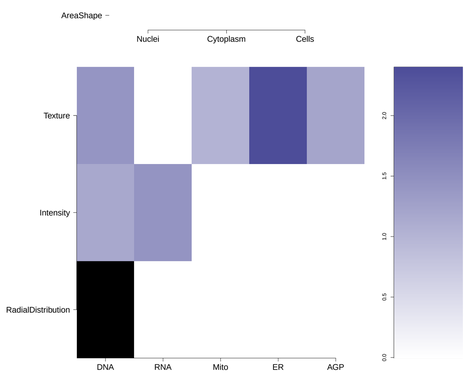
NA (in 1 replicates)

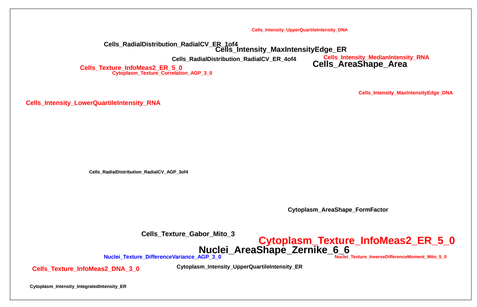
-0.50 ± 0.08

Treatment	Score
PRKCZ.K281R	-0.50
PRKCZ.WT.1	-0.48
PRKCZ.WT.2	-0.44

NA







Total number of assays tested in: 631. Active in the following assays:

- Cycloheximide Counterscreen for Small Molecule Inhibitors of Shiga Toxin (AID 2314)
- A qHTS for Small Molecule Inhibitors of Shiga Toxin (AID 2315)
- VP16 counterscreen qHTS for inhibitors of ROR gamma transcriptional activity (AID 2546)
- qHTS for inhibitors of ROR gamma transcriptional activity (AID 2551)
- uHTS identification of small molecule antagonists of the CCR6 receptor via a luminescent beta-arrestin assay (AID 493098)
- qHTS profiling assay for firefly luciferase inhibitor/activator using purifed enzyme and Km concentrations of substrates (counterscreen for miR-21 project) (AID 588342)
- uHTS identification of small molecule modulators of myocardial damage (AID 588492)
- Fluorescence-based cell-based primary high throughput screening assay to identify antagonists of the human trace amine associated receptor 1 (TAAR1) (AID 624466)
- uHTS identification of inhibitors of cullin neddylation in a TR-FRET assay (AID 651699)

BRD-K01990668-001-06-6
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ZINC00426457
AC1LD48Q
MLS000077450
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HMS1417B06
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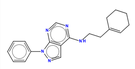
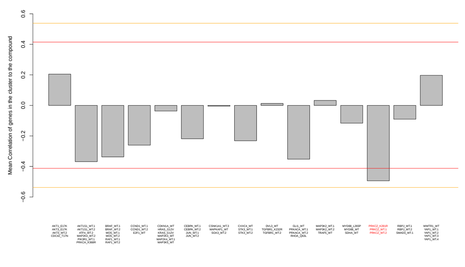


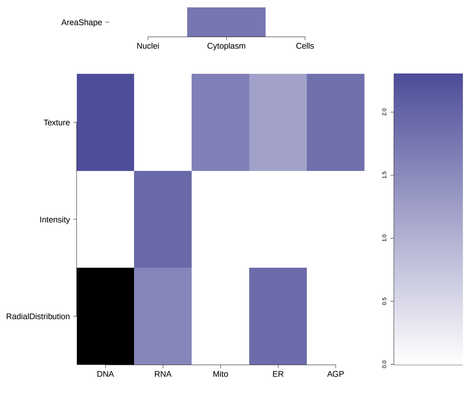
0.78 (in 3 replicates)

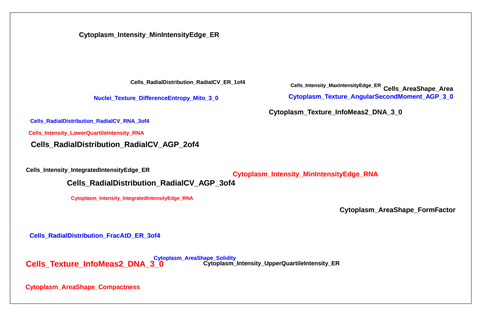
-0.49 ± 0.10

Treatment	Score
PRKCZ.K281R	-0.61
PRKCZ.WT.1	-0.46
PRKCZ.WT.2	-0.41

NA

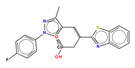
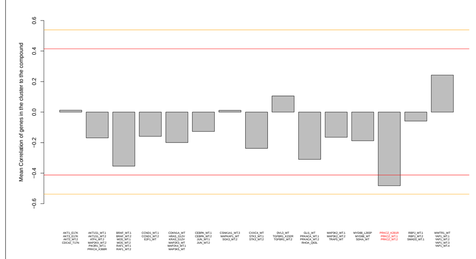
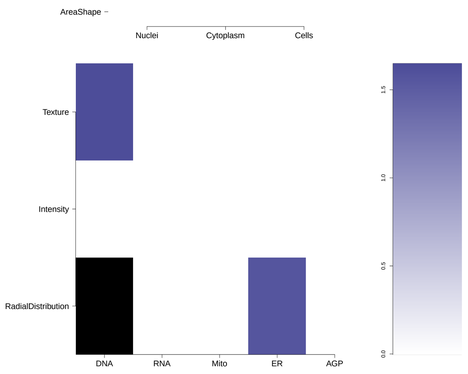
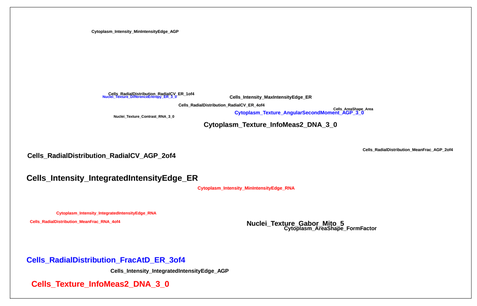
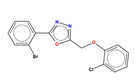
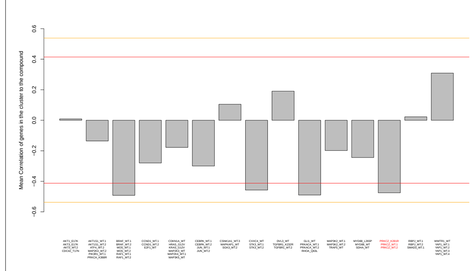
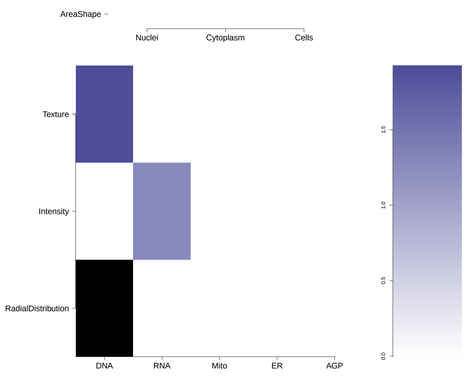
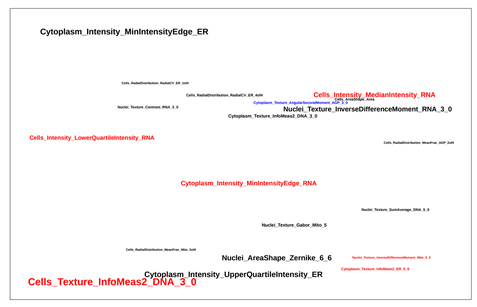
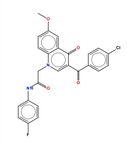
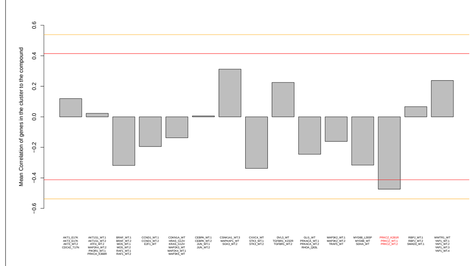
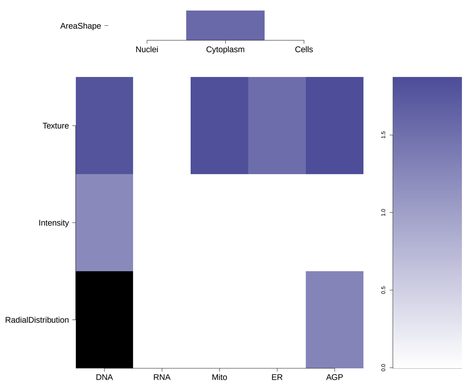
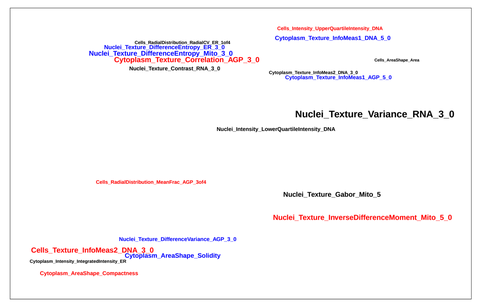
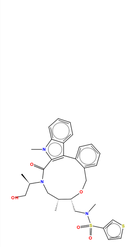
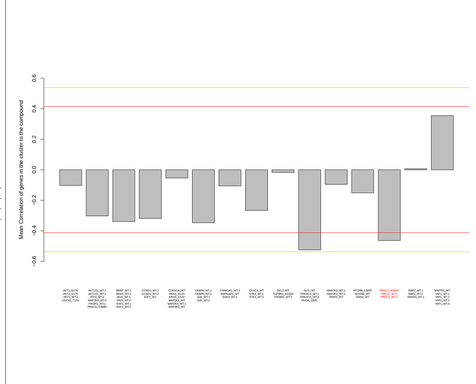
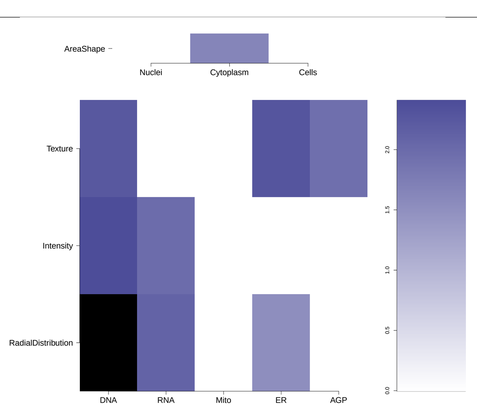
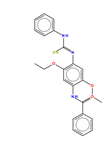
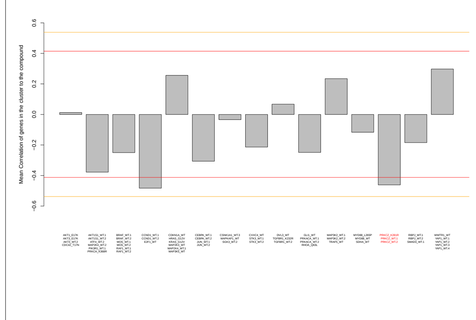
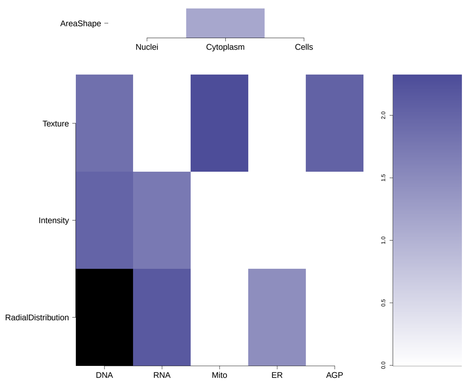
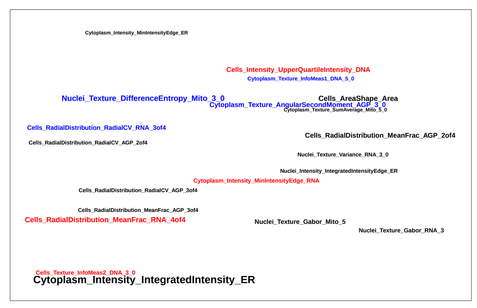






Total number of assays tested in: 697. Active in the following assays:

- CYP2C9 Assay (AID 777)
- High Throughput Screen to Identify Compounds that increase expression of NF-kB in Human Neuronal Cells - Primary Screen (AID 1239)
- qHTS Assay for Enhancers of SMN2 Splice Variant Expression (AID 1458)
- MLPCN Alpha-Synuclein 5'UTR - 5'-UTR binding - activators (AID 1814)
- Cycloheximide Counterscreen for Small Molecule Inhibitors of Shiga Toxin (AID 2314)
- A qHTS for Small Molecule Inhibitors of Shiga Toxin (AID 2315)
- uHTS Luminescent assay for identification of inhibitors of Sentrin-specific protease 6 (SENP6) (AID 2599)
- HTS for small molecule inhibitors of CHOP to regulate the unfolded protein response to ER stress (AID 2732)
- uHTS Luminescent assay for identification of inhibitors of Sentrin-specific protease 7 (SENP7) (AID 434973)
- qHTS Assay for Rab9 Promoter Activators (AID 485297)
- qHTS Assay for NPC1 Promoter Activators (AID 485313)
- Heat Shock Factor-1 (HSF-1) Measured in Cell-Based System Using Plate Reader - 2038-01_Activator_SinglePoint_HTS_Activity (AID 504408)
- qHTS screen for small molecules that induce genotoxicity in human embryonic kidney (HEK293T) cells expressing luciferase-tagged ELG1 (AID 504466)
- Primary qHTS for delayed death inhibitors of the malarial parasite plastid, 96 hour incubation (AID 504834)
- MITF Measured in Cell-Based System Using Plate Reader - 2084-01_Activator_SinglePoint_HTS_Activity (AID 588334)
- qHTS profiling assay for firefly luciferase inhibitor/activator using purifed enzyme and Km concentrations of substrates (counterscreen for miR-21 project) (AID 588342)
- Screen for inhibitors of the SWI/SNF chromatin remodeling complex (esBAF) in mouse embryonic stem cells with Luciferase reporter assay Measured in Cell-Based System Using Plate Reader - 2141-01_Inhibitor_SinglePoint_HTS_Activity (AID 602393)
- Luminescence-based cell-based primary high throughput screening assay to identify activators of the function of SWI/SNF related, matrix associated, actin dependent regulator of chromatin, subfamily a, member 2 (SMARCA2, BRM) (AID 652017)
- Counterscreen for activators of the function of SWI/SNF related, matrix associated, actin dependent regulator of chromatin, subfamily a, member 2 (SMARCA2, BRM): Luminescence-based cell-based high throughput screening assay to identify non-selective compounds using the VP16 reporter assay (AID 686939)
- qHTS for Inhibitors of PLK1-PDB (polo-like kinase 1 - polo-box domain): Primary Screen (AID 720504)

BRD-K71327258-001-05-4 T5822519 MLS000565087 AC1O37PV HMS2452A07 ZINC8672173 SMR000152273 PubChem CID : 6249835		0.61 (in 2 replicates)	-0.48 ± 0.08 Treatment | Score PRKCZ.K281R | -0.56 PRKCZ.WT.1 | -0.49 PRKCZ.WT.2 | -0.40	NA				Total number of assays tested in: 698. Active in the following assays: • Identification of Novel Modulators of Cl- dependent Transport Process via HTS: Primary Screen (AID 1456) • Primary biochemical high throughput screening assay to identify inhibitors of VIM-2 metallo-beta-lactamase (AID 1527) • Epi-absorbance primary biochemical high throughput screening assay to identify inhibitors of IMP-1 metallo-beta-lactamase (AID 1556) • Identification of Novel Modulators of Cl- dependent Transport Process via HTS: Retesting of KCC2 cells with Ouabain (AID 1717) • Epi-absorbance-based confirmation biochemical high throughput screening assay to identify selective inhibitors of VIM-2 metallo-beta-lactamase. (AID 1860) • Fluorescence-based primary cell-based high throughput screening assay to identify antagonists of the G-protein coupled receptor 7 (GPR7). (AID 1861) • qHTS for Antagonists of gsp, the Etiologic Mutation Responsible for Fibrous Dysplasia/McCune-Albright Syndrome: qHTS (AID 624288) • Fluorescence-based biochemical primary high throughput screening assay to identify molecules that bind r(CAG) RNA repeats (AID 651821) • Fluorescence-based biochemical high throughput confirmation assay to identify molecules that bind r(CAG) RNA repeats (AID 652065) • Counterscreen for molecules that bind rCAG RNA repeats: fluorescent based biochemical counterscreen assay for inhibitors of the DNA-based (5'CAG/3'GTC) TO-PRO-1 dye complex (AID 652068)
BRD-K29738272-001-05-7 MLS000058143 SMR000067553 ZINC02628750 MLS002634971 AC1M19D5 BDBM77989 HMS2339H20 ZINC2628750 T5270429 PubChem CID : 2091123		0.75 (in 2 replicates)	-0.48 ± 0.08 Treatment | Score PRKCZ.K281R | -0.52 PRKCZ.WT.1 | -0.52 PRKCZ.WT.2 | -0.38	NA				Total number of assays tested in: 780. Active in the following assays: • Primary Cell-based High Throughput Screening assay for activators of the nuclear receptor Steroidogenic Factor 1 (SF-1) (AID 522) • Primary Cell-based High Throughput Screening assay for activators of the Retinoic Acid Receptor-related orphan receptor A (RORA) (AID 560) • Primary Antimicrobial Assay for E. coli BW25113 and 8710;tolC::kan Protocol for 384-well HTS (AID 573) • Antimicrobial Assay for E. coli BW25113 and 8710;tolC::kan - Dose Response (AID 617) • qHTS Assay for Inhibitors of HPGD (15-Hydroxyprostaglandin Dehydrogenase) (AID 894) • Primary cell-based high throughput screening assay to measure STAT1 activation (AID 932) • HTS Assay for Activators of Cytochrome P450 2A9 (AID 1024) • MLPCN Alpha-Synuclein 5'UTR - 5'-UTR binding - activators (AID 1814) • qHTS Assay for Modulators of miRNAs and/or Inhibitors of miR-21 (AID 2289) • Cycloheximide Counterscreen for Small Molecule Inhibitors of Shiga Toxin (AID 2314) • A qHTS for Small Molecule Inhibitors of Shiga Toxin (AID 2315) • uHTS identification of small molecule inhibitors of tim10-1 yeast via a luminescent assay (AID 463190) • Single concentration confirmation of small molecule inhibitors of tim10-1 yeast via a luminescent assay (AID 463213) • qHTS Assay for NPC1 Promoter Activators (AID 485313) • HTS Assay for Allosteric Antagonists of the Human D2 Dopamine Receptor: Primary Screen for Antagonists (AID 485344) • Dose Response confirmation of uHTS small molecule inhibitors of tim10-1 yeast via a luminescent assay (AID 493003) • Dose Response confirmation of uHTS small molecule inhibitors of tim10-1: a luminescent TIM10 yeast counterscreen (AID 504542) • Dose Response confirmation of uHTS small molecule inhibitors of tim10-1: a luminescent tim23-1 yeast counterscreen. (AID 504544) • MITF Measured in Cell-Based System Using Plate Reader - 2084-01_Activator_Dose_CherryPick_Activity (AID 540258) • MITF Measured in Cell-Based System Using Plate Reader - 2084-01_Activator_SinglePoint_HTS_Activity (AID 588334) • qHTS profiling assay for firefly luciferase inhibitor/activator using purifed enzyme and Km concentrations of substrates (counterscreen for miR-21 project) (AID 588342) • Screen for inhibitors of the SWI/SNF chromatin remodeling complex (esBAF) in mouse embryonic stem cells with Luciferase reporter assay Measured in Cell-Based System Using Plate Reader - 2141-01_Inhibitor_SinglePoint_HTS_Activity (AID 602393) • qHTS Assay to Identify Small Molecule Activators of BRCA1 Expression (AID 624202)
BRD-K08564028-001-04-2 F1602-0395 AC1M21WZ MLS000724735 HMS2255H24 ZINC2688545 ZINC02688545 SMR000237570 PubChem CID : 2135408		NA (in 1 replicates)	-0.47 ± 0.13 Treatment | Score PRKCZ.K281R | -0.51 PRKCZ.WT.1 | -0.58 PRKCZ.WT.2 | -0.33	NA				Total number of assays tested in: 648. Active in the following assays: • Activator for delta FosB/delta FosB homodimer Measured in Biochemical System Using Plate Reader - 2072-01_Activator_SinglePoint_HTS_Activity (AID 493131) • uHTS identification of DNMT1 inhibitors in a Fluorescent Molecular Beacon assay (AID 588458) • uHTS luminescent assay for identification of compounds that enhance the survival of human induced pluripotent stem cells when cultured as single cells (AID 602274) • Dose response confirmation of uHTS hits that enhance the survival of human induced pluripotent stem cells when cultured as single cells in a luminescent assay (AID 623861) • Dose response confirmation of uHTS hits that enhance the survival of human induced pluripotent stem cells when cultured as single cells in a fluorescent-based, imaging assay (AID 624145)
BRD-K13515789-001-01-2 PubChem CID : 54638030		0.83 (in 3 replicates)	-0.46 ± 0.10 Treatment | Score PRKCZ.K281R | -0.57 PRKCZ.WT.1 | -0.43 PRKCZ.WT.2 | -0.39	0.851 ± 0.043 Treatment | Score PRKCZ.K281R | 0.886 PRKCZ.WT.1 | 0.864 PRKCZ.WT.2 | 0.803				Total number of assays tested in: 38.
BRD-K17013498-001-05-7 ST51025861 MLS000556066 HMS2413G21 ZINC8666980 ZINC08666980 ASN 03271100 SMR000172598 PubChem CID : 1172870		NA (in 1 replicates)	-0.46 ± 0.09 Treatment | Score PRKCZ.K281R | -0.57 PRKCZ.WT.1 | -0.38 PRKCZ.WT.2 | -0.44	NA				Total number of assays tested in: 649. Active in the following assays: • qHTS Assay for Inhibitors of 15-hLO-2 (15-human lipoxygenase 2) (AID 881) • Leishmania major promastigote HTS (AID 1063) • Primary cell-based high-throughput screening assay to identify agonists of the transient receptor potential channel ML3 (TRPML3) (AID 1448) • VP16 counterscreen qHTS for inhibitors of ROR gamma transcriptional activity (AID 2546) • qHTS for inhibitors of ROR gamma transcriptional activity (AID 2551) • HTS using DiI-HDL to assay lipid transfer in ldlA[SR-BI] cells Measured in Cell-Based System Using Plate Reader - 2085-01_Inhibitor_SinglePoint_HTS_Activity (AID 488896) • Primary qHTS for delayed death inhibitors of the malarial parasite plastid, 96 hour incubation (AID 504834) • Counterscreen of compound fluorescence effects on High-throughput multiplex microsphere screening for inhibitors of toxin protease (AID 624483) • qHTS for Inhibitors of human tyrosyl-DNA phosphodiesterase 1 (TDP1): qHTS in cells in absence of CPT (AID 686978) • qHTS for Inhibitors of human tyrosyl-DNA phosphodiesterase 1 (TDP1): qHTS in cells in presence of CPT (AID 686979)

BRD-K07726175-001-06-4
MLS000101513
SMR000017682
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ZINC02723956
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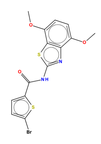
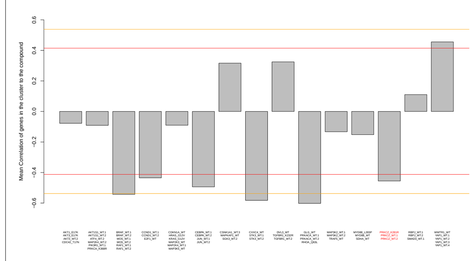


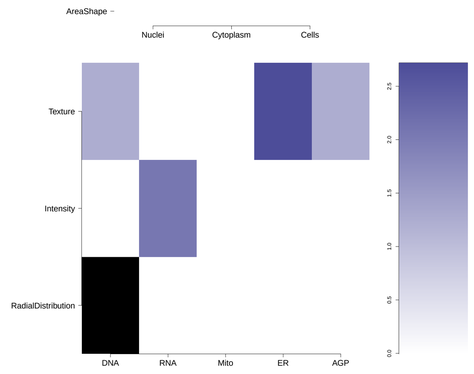
NA (in 1 replicates)

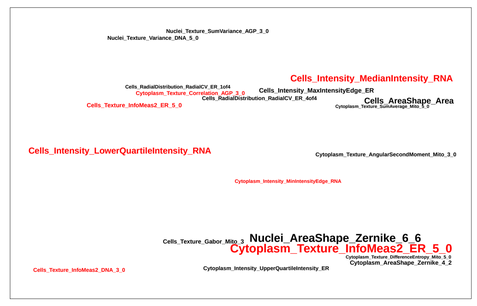
-0.46 ± 0.09

Treatment	Score
PRKCZ.K281R	-0.56
PRKCZ.WT.1	-0.41
PRKCZ.WT.2	-0.40

NA







Total number of assays tested in: 760. Active in the following assays:

- Primary Cell-based High Throughput Screening assay for activators of the nuclear receptor Steroidogenic Factor 1 (SF-1) (AID 522)
- HTS of Estrogen Receptor- alpha Coactivator Binding inhibitors (AID 629)
- HTS for Estrogen Receptor-beta Coactivator Binding inhibitors (AID 633)
- Primary cell-based high-throughput screening assay to identify agonists of Galanin Receptor 2 (GALR2) (AID 803)
- uHTS of Mcl-1/Bid interaction inhibitors (AID 1021)
- uHTS of Mcl-1/Noxa interaction inhibitors (AID 1022)
- Dose Response Confirmation for Mcl-1/Bid Interaction Inhibitors (AID 1418)
- Identification of Novel Modulators of Cl- dependent Transport Process via HTS: Primary Screen (AID 1456)
- qHTS Assay for Promiscuous and Specific Inhibitors of Cruzain (without detergent) (AID 1476)
- HTS for small molecule inhibitors of CHOP to regulate the unfolded protein response to ER stress (AID 2732)
- uHTS identification of DNMT1 inhibitors in a Fluorescent Molecular Beacon assay (AID 588458)
- qHTS for Stage-Specific Inhibitors of Vaccinia Orthopoxvirus: Venus Reporter Primary qHTS (AID 720580)