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Compound Libraries



Top academic journals such as **Science, Nature, and Cell** have published over **206** articles featuring Selleck Products.

Nature, 2019, 572(7768):254-259	Nature, 2016, 539(7627):54-58	Cell, 2019, 176(4):869-881
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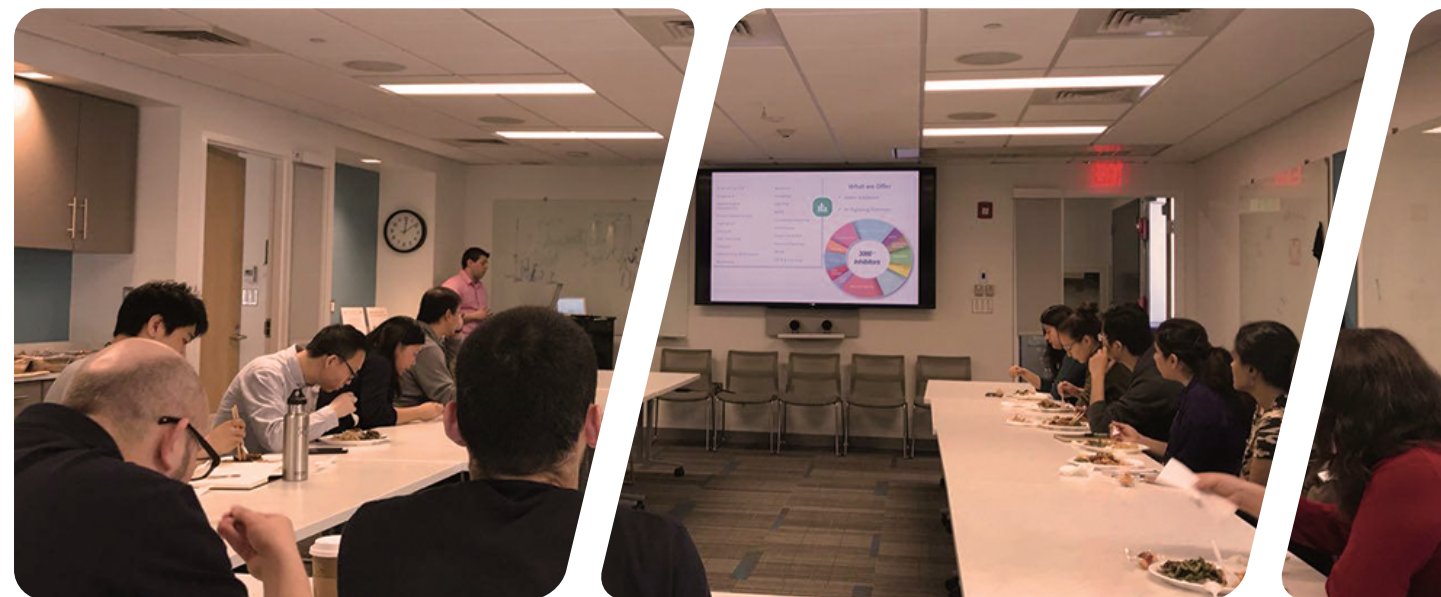
Q1 Why choose Selleck Libraries?

Put your research on the fast track to publication! Selleckchem customers publish on average within **two years**. Compared to other methods, our libraries save you time and help increase article impact range. With a Selleckchem library, you'll have a **faster** and **more comprehensive** way to get your article published.

Institute	Group	Duration	Journal	IF
Tsinghua University	YN Du	2y	Nature Materials	39.74
UCSF	JR Chan	1y8m	Nature Medicine	29.89
Roche Innovation Center Basel	CA Cowan	2y4m	Nature Cell Biology	20.06
BC Cancer Agency	S Dedhar	3y	Nature Communications	12.12
Baylor College of Medicine	XH Zhang	1y10m	Nature Communications	12.12
Lund University	JU Kazi	2y8m	Cancer Letters	6.34
University of Wisconsin	X Zhao	1y6m	Stem Cells	5.6
Wuhan Institute of Virology	GF Xiao	1y11m	Journal of Virology	4.66
University of Bergen	BT Gjertsen	1y7m	Pharmacological Research	4.48
University of Nevada	BS Ferguson	1y	Molecular Nutrition & Food Research	4.48
Moffitt Cancer Center	E Schönbrunn	3y	ChemMedChem	3.23
...				
Average		2 years		12.97

Q2 Need help with the library?

We provide free technical advice and assistance with experimental design.



Q3 Pressed for time?

Our partners can provide a platform for screening and collaboration.



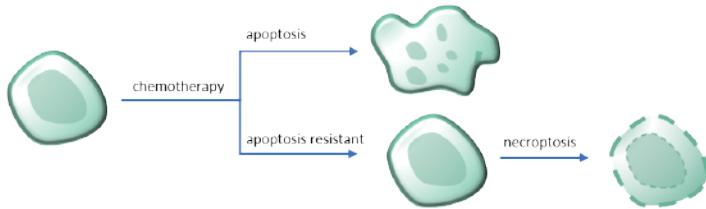
Disease Research Applications of Compound Libraries: Case Study I

Inhibition of Aurora Kinase A Induces Necroptosis in Pancreatic Carcinoma

The author purchased the Selleck's Kinase Inhibitor Library in May of 2015. This article (PMID:28764929) was published in the *Journal of Gastroenterology* (IF= 19.23) in November of 2017.

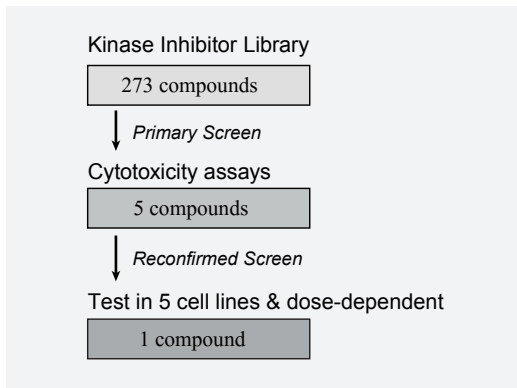
Background: Inducing cell necrosis is a new way to treat pancreatic cancer

The anti-apoptotic mechanism of pancreatic cancer cells themselves makes them difficult to kill. Therefore, a cell death mode, which is independent of the form of apoptosis, cell necrosis has caused widespread concern. The development of drugs that induce tumor cell necrosis has become a new idea for the treatment of pancreatic cancer.



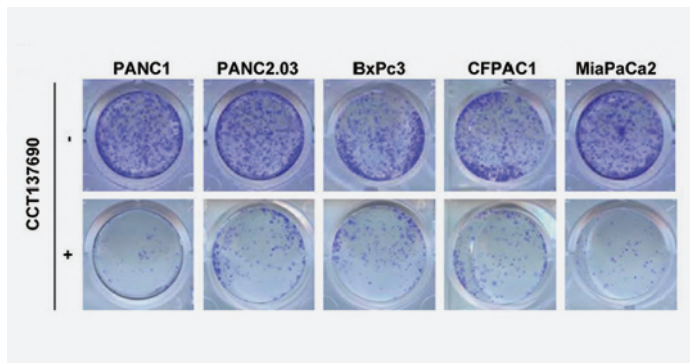
Experiment Design

1. High-throughput drug screening



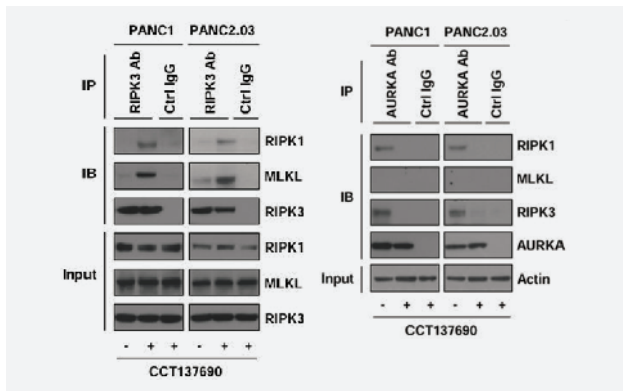
Compound Library: Selleck Kinase Inhibitor Library
Model: Human pancreatic ductal adenocarcinoma cell PANC1
Indicators: Cell viability and cell death assays confirm that CCT137690 kills human pancreatic ductal adenocarcinoma cells

2. Phenotype research



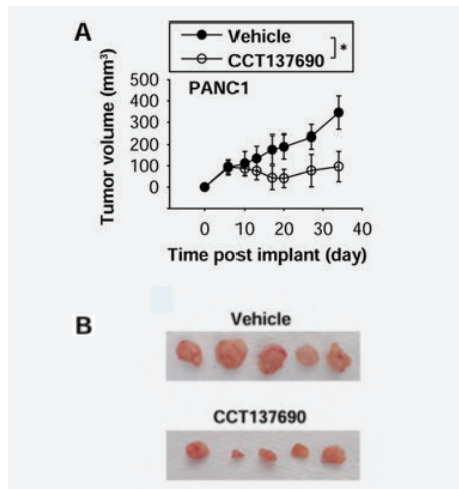
1. Cell morphology study
2. Cloning formation experiment

3. Mechanism research



AURKA can bind to the RIPK1/RIPK3 necrotic complex to inhibit necrosis. Tumor cell necrosis is activated after inhibition by CCT137690.

4. *in vivo* efficacy verification



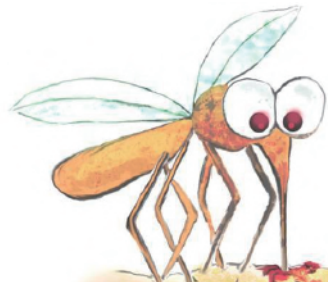
Disease Research Applications of Compound Libraries: Case Study II

Screening of FDA-Approved Drugs for Inhibitors of Japanese Encephalitis Virus Infection

The author purchased the Selleck's FDA-approved Drug Library in September of 2015, and published three articles in four years. The article (PMID:28814523) was published in the *Journal of Virology* (IF:4.324) in August of 2017.

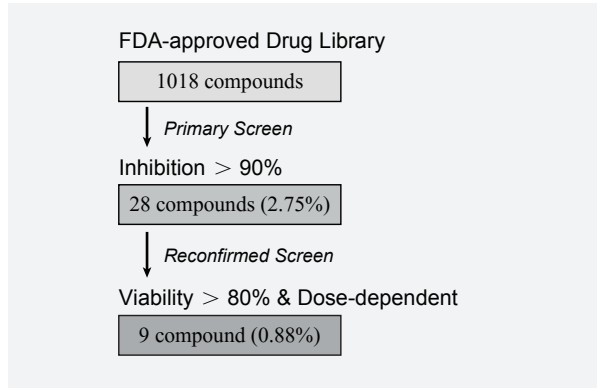
Background: Japanese encephalitis lacks any form of treatment.

Japanese encephalitis is caused by Japanese encephalitis virus (JEV). Except for vaccination, there are still no specific drugs or treatments for JEV.



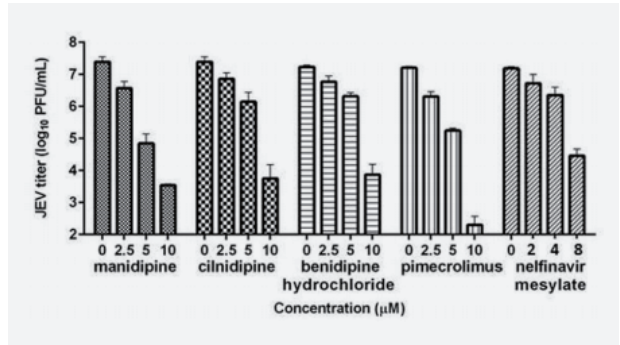
Experiment Design

1. High-throughput drug screening



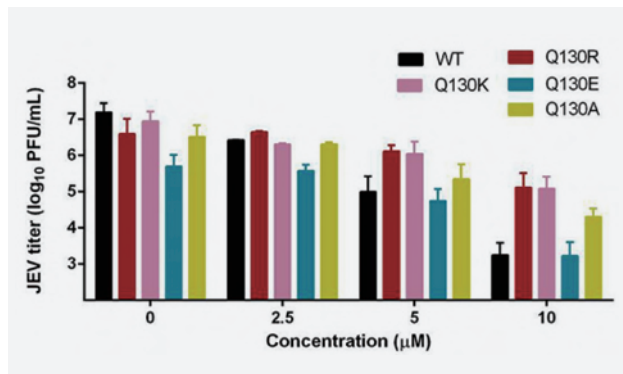
Compound Library: Selleck FDA-approved Drug Library
Model: JEV infectious clone with luciferase reporter
Indicators: inhibition> 90%, viability> 80%, dose–response relationship establishment, five hit drugs were confirmed, of which three were calcium channel inhibitors

2. Phenotype research



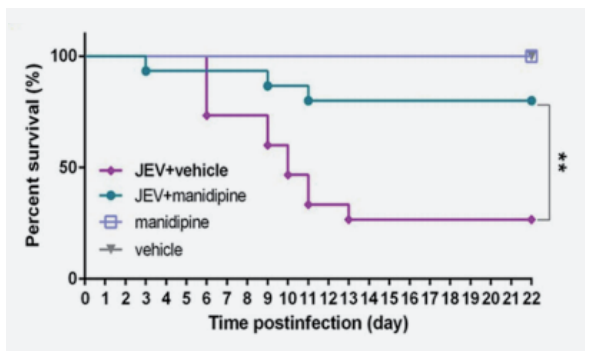
1. Dose-effect relationship verification
2. Time-effect relationship verification
3. Antiviral spectrum study

3. Mechanism research



1. Using other calcium channel inhibitors can have similar effects
2. Identify the drug's target site for JEV

4. *in vivo* efficacy verification



FDA-approved Drug Library [Cat.No. L1300](#)

- A unique collection of **2,576** FDA-approved APIs
- Biosafety and effectiveness validated by clinical trials

Size (Pre-dissolved in DMSO or Water)

100 µL/well (2 mM or 10 mM solution)



Customize Your Library



Specific Compounds



Quantities



Plate map



Format (Dry/solid or DMSO solution)

Target Class	Number of FDA-approved Compounds
Others	500
Anti-infection	300
Adrenergic Receptor	90
5-HT Receptor	80
ACR	70
DNA/RNA Synthesis	60
Histamine Receptor	50
Immunology & Inflammation related	40
Dopamine Receptor	30
CCK	20
Sodium Channel	20
Calcium Channel	20
AdoPhag	20
Glucocorticoid Receptor	20
Topoisomerase	20
PKMS	20
ECER	20
PKC	20
Vitamin	20
P450 (e.g. CYP17)	20
Reverse Transcriptase	20
EGFR	20
PDGFR	20
ADK	20
GABA Receptor	20
KCI Protease	20
N/A	20
C-Kit	20
Microtubule Associated	20
Dehydrogenase	20
JAK	20
DPR-4	20
Androgen Receptor	20
Proton Pump	20
PKA	20
HMG-CoA Reductase	20
Opioid Receptor	20
RGS	20
HIV Protease	20
Thrombin	20
CKK	20
Carbonic Anhydrase	20
S-alpha Reductase	20
PKK	20
ILK2	20
Bcr-Abl	20
SGLT	20
Retinoid Receptor	20
mTOR	20
MKO	20
FLT3	20
DNK	20
Transferrin	20
Other targets	260

Journals Citing of this Library

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Assay Drug Dev Technol,2019,17(1):14-16	Nat Med,2014,20(8):954-60
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ACS Infect Dis,2018,4(4):635-645	Oncotarget,2015,6(3):1531-43
J Lipid Res,2018,59(10):1916-1926	Pharmacol Res,2016,113(Pt A):216-227
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J Cell Physiol,2018,233(5):3713-3722	Biosens Bioelectron,2015,68:699-704
J Cell Physiol,2018,233(1):422-433	Curr Protoc Chem Biol,2014,4:177-191
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Clin Cancer Res,2016,10.1158/1078-0432	Neuropsychopharmacology,2019,10.1038/s41386-018-0266-7
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Preclinical and Clinical Compound Library [Cat.No. L3900](#)

- A unique collection of **2,578** preclinical and clinical compounds
- Includes drug molecules with preclinical studies, clinical drug molecules, and drugs already on the market

Size (Pre-dissolved in DMSO or Water)

100 µL/well (2 mM or 10 mM solution)



Customize Your Library



Specific Compounds



Quantities



Plate map



Format (Dry/solid or DMSO solution)

Target Class	Number of Clinical Compounds
Others	430
Anti-infection	230
Adrenergic Receptor	100
5-HT Receptor	90
ACR	80
DNA/RNA Synthesis	70
Histamine Receptor	60
Immunology & Inflammation related	50
Dopamine Receptor	40
CCK	30
Sodium Channel	20
Calcium Channel	20
AdoPhag	20
Glucocorticoid Receptor	20
Topoisomerase	20
PKMS	20
ECER	20
PKC	20
Vitamin	20
P450 (e.g. CYP17)	20
Reverse Transcriptase	20
EGFR	20
PDGFR	20
ADK	20
GABA Receptor	20
KCI Protease	20
N/A	20
C-Kit	20
Microtubule Associated	20
Dehydrogenase	20
JAK	20
DPR-4	20
Androgen Receptor	20
Proton Pump	20
PKA	20
HMG-CoA Reductase	20
Opioid Receptor	20
RGS	20
HIV Protease	20
Thrombin	20
CKK	20
Carbonic Anhydrase	20
S-alpha Reductase	20
PKK	20
ILK2	20
Bcr-Abl	20
SGLT	20
Retinoid Receptor	20
mTOR	20
MKO	20
FLT3	20
DNK	20
Transferrin	20
Other targets	250

Journals Citing of this Library

Sci Adv,2019,5(6):eaav9784	PLoS One,2016,11(2):e0149848
Neuropsychopharmacology,2019,10.1038/s41386-018-0266-7	Sensors (Basel),2016,16(3)
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Kaohsiung J Med Sci,2019,10.1002/kjm2.12051	Cell Cycle,2015,14(1):109-22
Cell Rep,2018,22(5):1185-1199	PLoS One,2015,10(11):e0142655
Cancer Sci,2018,109(4):1220-1229	Oncotarget,2015,6(3):1531-43
Sci Rep,2018,8(1):13106	Head Neck,2015,37(12):1722-32
J Cell Physiol,2018,233(1):422-433	Nat Med,2014,20(8):954-60
Anal Chem,2017,89(12):6678-6685	Mol Cancer Res,2014,12(5):703-13
Antimicrob Agents Chemother,2017,62(1)	Oncotarget,2014,5(15):6512-25
Pharmacol Res,2016,113(Pt A):216-227	Curr Protoc Chem Biol,2014,4:177-191

Metabolism Compound Library [Cat.No. L3700](#)

- A unique collection of **1,383** metabolically related active compounds, including inhibitors and human endogenous metabolites
- Covers multiple metabolic pathways such as glucose metabolism, lipid metabolism, protein hydrolysis, and nucleotide metabolism
- Covers more than 200 targets such as FXR, DPP-4, glucokinase, HSP, PDE, PPAR, etc.

Size (Pre-dissolved in DMSO or Water)

100 µL/well (2 mM or 10 mM solution)

**Customize Your Library**

Specific Compounds

Quantities

Plate map

Format (Dry/solid or DMSO solution)



Journals Citing of this Library

Sci Adv ,2019,5(6):eaav9784	PLOS One ,2016,11(2):e0149848
Neuropsychopharmacology ,2019,10.1038/s41386-018-0266-7	Sensors (Basel) ,2016,16(3)
Breast Cancer Res ,2019,21(1):37	Clin Cancer Res ,2015,21(5):1172-82
Neurobiol Aging ,2019,76:24-34	Biosens Bioelectron ,2015,68:699-704
Bioengineered ,2019,10(1):98-107	Mol Cancer Ther ,2015,14(5):1213-23
Kaohsiung J Med Sci ,2019,10.1002/kjm2.12051	Cell Cycle ,2015,14(1):109-22
Cell Rep ,2018,22(5):1185-1199	PLOS One ,2015,10(11):e0142655
Cancer Sci ,2018,109(4):1220-1229	Oncotarget ,2015,6(3):1531-43
Sci Rep ,2018,8(1):13106	Head Neck ,2015,37(12):1722-32
J Cell Physiol ,2018,233(1):422-433	Nat Med ,2014,20(8):954-60
Nat Med ,2017,23(4):405-408	Mol Cancer Res ,2014,12(5):703-13
Anal Chem ,2017,89(12):6678-6685	Oncotarget ,2014,5(15):6512-25
Antimicrob Agents Chemother ,2017,e01674-17	Curr Protoc Chem Biol ,2014,4:177-191
Pharmacol Res ,2016,113(Pt A):216-227	

Kinase Inhibitor Library [Cat.No. L1200](#)

- A unique collection of **1,188** kinase-related active compounds
- Includes small molecules with biological research, clinical drug molecules, and drugs already on the market
- Targets include EGFR, PI3K, Aurora Kinase, CDK and MEK, etc.

Size (Pre-dissolved in DMSO or Water)

100 µL/well (2 mM or 10 mM solution)

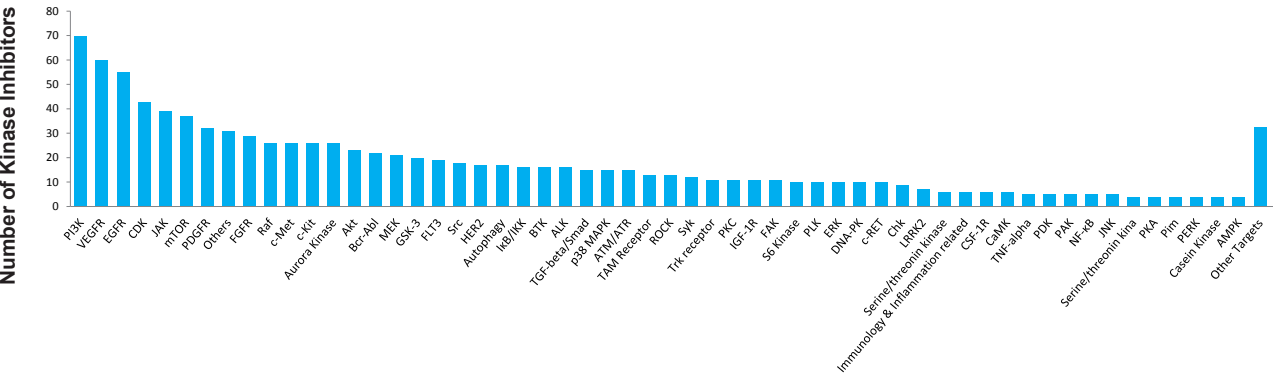
**Customize Your Library**

Specific Compounds

Quantities

Plate map

Format (Dry/solid or DMSO solution)



Journals Citing of this Library

Nat Commun ,2019,10(1):2197	J Biol Chem ,2017,292(41):17037-17045
Sci Adv ,2019,5(6):eaav9784	Front Plant Sci ,2017, 10.3389/fpls.2017.00852.
NPJ Genom Med ,2019,4:7	ChemMedChem ,2017,12(22):1857-1865
Neuropsychopharmacology ,2019,10.1038/s41386-018-0266-7	Biochem Biophys Res Commun ,2017,493(1):58-63
J Exp Clin Cancer Res ,2019,38(1):56	J Mol Cell Cardiol ,2016,97:106-13
Breast Cancer Res ,2019,21(1):37	Pharmacol Res ,2016,113(Pt A):216-227
Development ,2019,146(14)	ChemMedChem ,2016,11(11):1137-44
Neurobiol Aging ,2019,76:24-34	ChemMedChem ,2016,10.1002/cmdc
Anal Biochem ,2019,569:46-52	Bioorg Med Chem ,2016,24(19):4647-51
Bioengineered ,2019,10(1):98-107	PLOS One ,2016,11(2):e0149848
Kaohsiung J Med Sci ,2019,10.1002/kjm2.12051	Sensors (Basel) ,2016,16(3)
Methods Mol Biol ,2019,1888:115-126	Clin Cancer Res ,2015,21(5):1172-82
Methods Mol Biol ,2019,1888:21-43	Biosens Bioelectron ,2015,68:699-704
Cell Rep ,2018,22(5):1185-1199	J Control Release ,2015,204:20-9
Mol Cancer Ther ,2018,10.1158/1535-7163.MCT-18-0179	Oncogene ,2015,34(32):4199-210
Stem Cell Res Ther ,2018,9(1):319	Mol Cancer Ther ,2015,14(5):1213-23
Cancer Sci ,2018,109(4):1220-1229	Breast Cancer Res Treat ,2015,149(3):715-26
Antimicrob Agents Chemother ,2018,10.1128/AAC.01744-18	Cell Cycle ,2015,14(1):109-22
Hum Gene Ther ,2018,29(8):886-901	BMC Cancer ,2015,15:239
Sci Rep ,2018,8(1):13106	PLOS One ,2015,10(11):e0142655
J Cell Physiol ,2018,233(1):422-433	Head Neck ,2015,37(12):1722-32
Cell Stress ,2018,10.15698/cst2018.04.131	Oncotarget ,2015,6(3):1531-43
FEBS Open Bio ,2018,9(1):82-91	Nat Med ,2014,20(8):954-60
Antiviral Res ,2018,158:226-237	Mol Cancer Ther ,2014,13(2):353-63
Nat Med ,2017,23(4):405-408	Mol Cancer Res ,2014,12(5):703-13
Gastroenterology ,2017,153(5):1429-1443	ACS Chem Biol ,2014,9(5):1160-71
Nat Commun ,2017,8:15289	PLOS One ,2014,9(7):e102741
Cell Rep ,2017,21(9):2639-2646	Mol Med Rep ,2014,10(6):3348-56
Cancer Lett ,2017,405:73-78	Oncotarget ,2014,5(15):6512-25
Anal Chem ,2017,89(12):6678-6685	Curr Protoc Chem Biol ,2014,4:177-191
J Mol Biol ,2017,429(13):2042-2045	PLOS One ,2013,8(5):e63240
Biochim Biophys Acta ,2017,1861(4):947-957	PLOS One ,2013,8(4):e60334
Antimicrob Agents Chemother ,2017,62(1)	

Natural Product Library [Cat.No. L1400](#)

- A unique collection of **2,116** natural products
- Small toxic side effects, easy to absorb
- Pharmacological activity and medicinal value reported

Size (Pre-dissolved in DMSO or Water)

100 µL/well (2 mM or 10 mM solution)



Customize Your Library



Specific Compounds



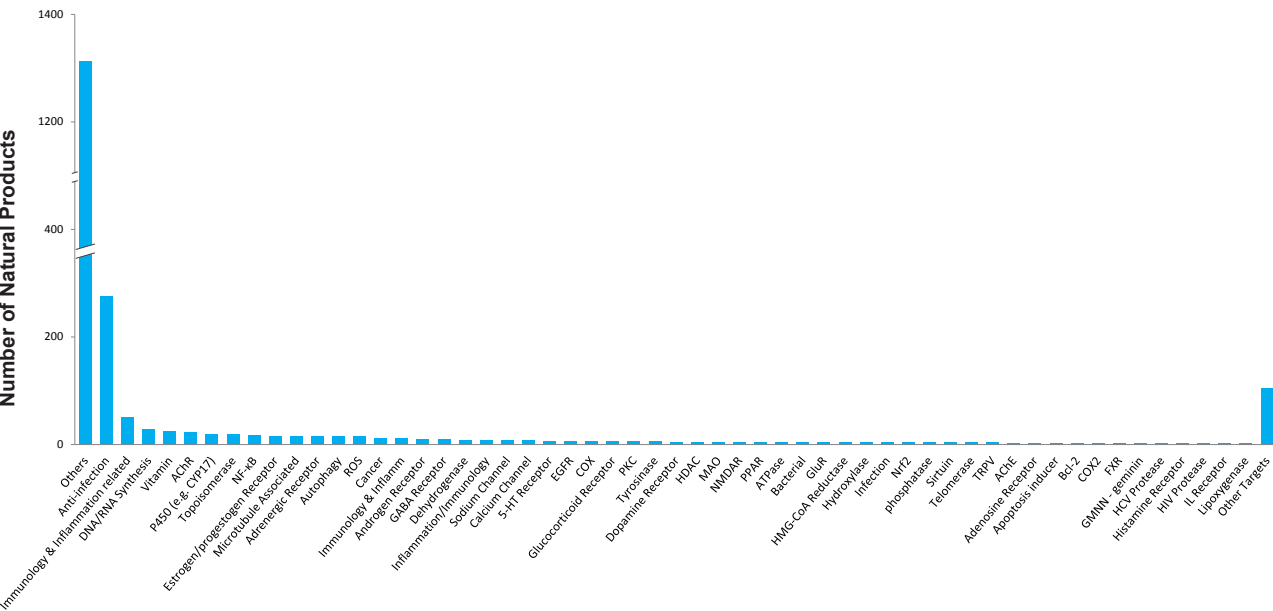
Quantities



Plate map



Format (Dry/solid or DMSO solution)



Journals Citing of this Library

Sci Adv ,2019,5(6):eaav9784	Nat Med ,2017,23(4):405-408
Cell Rep ,2019,26(6):1544-1556	Anal Chem ,2017,89(12):6678-6685
Neuropsychopharmacology ,2019,10.1038/s41386-018-0266-7	Mol Nutr Food Res ,2017,61(4)
Cancer Lett ,2019,459:50-58	Antimicrob Agents Chemother ,2017,e01674-17
Protein Cell ,2019,10(6):417-435	PeerJ ,2017,5:e3283
Breast Cancer Res ,2019,21(1):37	Pharmacol Res ,2016,113(Pt A):216-227
Oncogenesis ,2019,8(3):14	PLoS One ,2016,11(2):e0149848
Neurobiol Aging ,2019,76:24-34	Biochem Biophys Res Commun ,2016,473(4):775-80
Sci China Life Sci ,2019,10.1007/s11427-018-9477-3	Sensors (Basel) ,2016,16(3)
Bioengineered ,2019,10(1):98-107	Cytotechnology ,2016,68(4):1633-40
Kaohsiung J Med Sci ,2019,10.1002/kjm2.12051	Clin Cancer Res ,2015,21(5):1172-82
bioRxiv ,2019,10.1101/517086	Biosens Bioelectron ,2015,68:699-704
Cell Rep ,2018,22(5):1185-1199	Mol Cancer Ther ,2015,14(5):1213-23
Mol Neurobiol ,2018,10.1007/s12035-018-0953-8	PLoS Negl Trop Dis ,2015,9(6):e0003878
Cancer Sci ,2018,109(4):1220-1229	Cell Cycle ,2015,14(1):109-22
Antiviral Res ,2018,S0166-3542(18)30404-2	PLoS One ,2015,10(11):e0142655
Sci Rep ,2018,8(1):13106	Oncotarget ,2015,6(3):1531-43
Int J Biol Sci ,2018,10.7150/ijbs	Head Neck ,2015,37(12):1722-32
Int J Biol Sci ,2018,14(11):1521-1534	Nat Med ,2014,20(8):954-60
J Biol Chem ,2018,10.1074/jbc.RA118.006986	Mol Cancer Res ,2014,12(5):703-13
J Cell Physiol ,2018,233(1):422-433	Oncotarget ,2014,5(15):6512-25
Int J Biol Macromol ,2018,116:173-181	Curr Protoc Chem Biol ,2014,4:177-191
Sci Rep ,2018,8(1):14178	

Immunology/Inflammation Compound Library [Cat.No. L4100](#)

- A unique collection of **1,091** active compounds associated with immune inflammation
- Includes molecules with biological research, clinical drug molecules, and drugs already on the market
- Targets include COX, Nrf2, CXCR, ROS, TLR, p38 MAPK, NOS, etc.

Size (Pre-dissolved in DMSO or Water)

100 µL/well (2 mM or 10 mM solution)



Customize Your Library



Specific Compounds



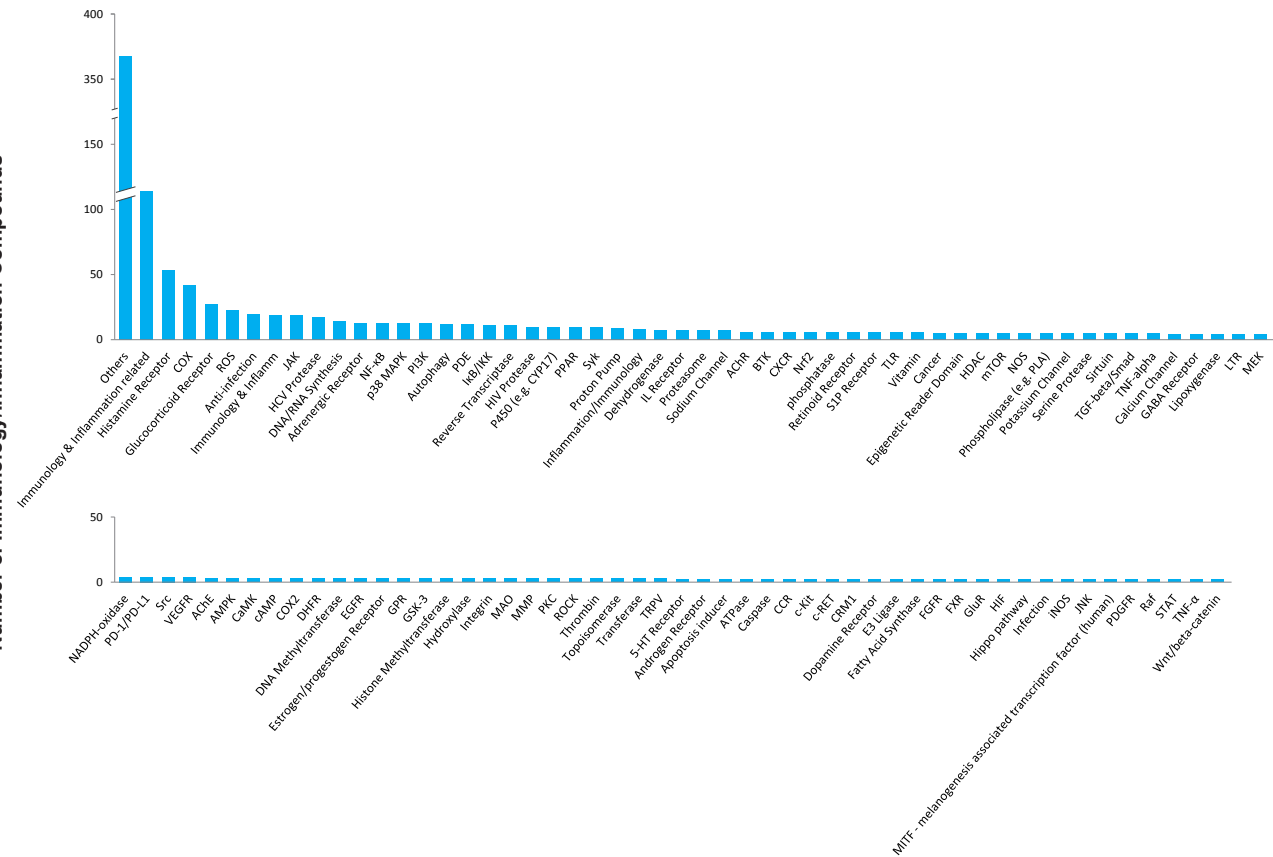
Quantities



Plate map



Format (Dry/solid or DMSO solution)



Journals Citing of this Library

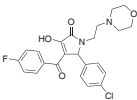
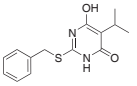
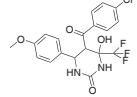
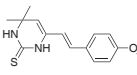
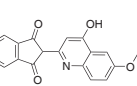
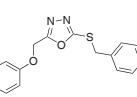
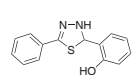
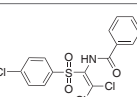
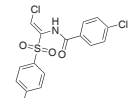
Sci Adv ,2019,5(6):eaav9784	PLoS One ,2016,11(2):e0149848
Neuropsychopharmacology ,2019,10.1038/s41386-018-0266-7	Sensors (Basel) ,2016,16(3)
Breast Cancer Res ,2019,21(1):37	Clin Cancer Res ,2015,21(5):1172-82
Neurobiol Aging ,2019,76:24-34	Biosens Bioelectron ,2015,68:699-704
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Antimicrob Agents Chemother ,2017,62(1)	Oncotarget ,2014,5(15):6512-25
Pharmacol Res ,2016,113(Pt A):216-227	Curr Protoc Chem Biol ,2014,4:177-191

Express-Pick Library (Provided by Pfizer) [Cat.No. L3600](#)

- **4,208** innovative compounds of different parent core structures
- Compounds from Pfizer
- Diverse structures and pharmacophores increase effective binding to biological targets

Size (Pre-dissolved in DMSO)	
100 µL/well	(10 mM solution)

Express-Pick Library Composition

Structure	Name	ID	Formula	Molecular Weight
	WAY-216050	1	C ₂₃ H ₂₂ ClFN ₂ O ₄	444.8831832
	WAY-312414	2	C ₁₄ H ₁₆ N ₂ O ₂ S	276.35404
	WAY-217028	3	C ₁₉ H ₁₆ ClF ₃ N ₂ O ₄	428.7895496
	WAY-312092	4	C ₁₅ H ₁₈ N ₂ OS	274.38122
	WAY-312398	5	C ₂₀ H ₁₅ NO ₄	333.3374
	WAY-311274	6	C ₁₇ H ₁₅ ClN ₂ O ₂ S	346.8312
	GI-523285	7	C ₁₄ H ₁₂ N ₂ OS	256.32288
	WAY-312818	8	C ₁₅ H ₁₀ Cl ₃ NO ₃ S	390.6688
	WAY-312684	9	C ₁₆ H ₁₃ Cl ₂ NO ₃ S	370.25032
...				

Other Compound Libraries

HTS Library for Drug Discovery (Provided by Pfizer) Cat.NO. L5000 Contains a unique collection of 99,040 innovative and diverse compounds
Human Endogenous Metabolite Compound Library Cat.No. L4500 A unique collection of small molecules of 408 endogenous metabolites
Drug Repurposing Library Cat.No. L3800 2,833 drug collections that have been marketed or have passed clinical phase I
Fragment Library Cat.No. L1600 A unique collection of 1,015 fragments of small molecules
Angiogenesis Related compound Library Cat.No. L5200 A unique collection of 113 bioactive compounds associated with angiogenesis
Anti-cancer Compound Library Cat.No. L3000 A unique collection of 1,966 anti-tumor compounds in clinical trials
Cambridge Cancer Compound Library Cat.No. L2300 A unique collection of 247 high value-added anticancer compounds
Anti-cancer Metabolism Compound Library Cat.No. L5700 A unique collection of 198 bioactive compounds for cancer metabolism research
Small Molecule Immuno-Oncology Compound Library Cat.No. L4800 A unique collection of 92 tumor immune small molecules
Cell Cycle compound library Cat.No. L5100 A unique collection of 121 bioactive compounds associated with the cell cycle
DNA Damage/DNA Repair compound Library Cat.No. L4600 179 unique collections of bioactive compounds for DNA damage and repair studies
Antibiotics compound Library Cat.No. L5300 A unique collection of 308 antibiotics
Anti-infection Compound Library Cat.No. L3100 A unique collection of 938 compounds with anti-infective activity
Anti-alzheimer's Disease Compound Library Cat.No. L5900 490 unique collections of active compounds for studying Alzheimer's disease
Anti-diabetic Compound Library Cat.No. L2900 A unique collection of 155 bioactive compounds associated with the development of diabetes
Inhibitor Library Cat.No. L1100 A unique collection of 2,790 latest inhibitors
Highly Selective Inhibitor Library Cat.No. L3500 A unique collection of 320 highly selective small molecule inhibitors
Covalent Inhibitor Library Cat.No. L5800 A unique collection of 81 covalent inhibitors

Other Compound Libraries

Tyrosine Kinase Inhibitor Library <i>Cat.No. L1800</i> A unique collection of 395 active compounds associated with tyrosine kinases
Protease Inhibitor Library <i>Cat.No. L2500</i> a unique collection of 229 protease-related bioactive compounds
Epigenetics Compound Library <i>Cat.No. L1900</i> A unique collection of 464 bioactive compounds associated with epigenetic research
Histone modification compound library <i>Cat.No. L4900</i> A unique collection of 112 bioactive compounds associated with histone modifications
Ubiquitination Compound Library <i>Cat.No. L6000</i> a unique collection of 153 bioactive compounds associated with ubiquitination
Stem Cell Signaling Compound Library <i>Cat.No. L2100</i> A unique collection of 297 bioactive compounds associated with stem cell signaling pathways
Autophagy Compound Library <i>Cat.No. L2600</i> A unique collection of 830 bioactive compounds associated with autophagy signaling pathways
Apoptosis Compound Library <i>Cat.No. L3300</i> A unique collection of 700 biologically active compounds that target apoptotic proteins such as Bcl-2, Caspase, p53, TNF-alpha, Mdm2 or survivin
Neuronal Signaling Compound Library <i>Cat.No. L4000</i> 1,111 unique collection of biologically active compounds associated with neural signaling pathways
CNS-Penetrant Compound Library <i>Cat.No. L4700</i> A unique collection of 307 bioactive compounds that cross the blood-brain barrier
Ion Channel Ligand Library <i>Cat.No. L2700</i> A unique collection of 232 bioactive compounds associated with ion channel ligands
PI3K/Akt Inhibitor Library <i>Cat.No. L2800</i> A unique collection of 174 biologically active compounds associated with the PI3K signaling pathway
MAPK Inhibitor Library <i>Cat.No. L3400</i> A unique collection of 95 biologically active compounds associated with the MAPK signaling pathway
GPCR Compound Library <i>Cat.No. L2200</i> A unique collection of 746 bioactive compounds associated with GPCRs
JAK/STAT compound library <i>Cat.No. L5400</i> A unique collection of 51 biologically active compounds associated with the JAK-STAT signaling pathway
NF-κB Signaling Compound Library <i>Cat.No. L5500</i> A unique collection of 41 biologically active compounds associated with the NF-κB signaling pathway
TGF-beta/Smad compound library <i>Cat.No. L5600</i> A unique collection of 61 bioactive compounds associated with the TGF-β/Smad signaling pathway

※ Please visit www.selleckchem.com to inquire prices of other compound libraries.



FAQ

Q1 Are there patent issues after a successful screening?

The compounds do not have any patent issues. When we “discover a hit”, it refers to a novel discovery for a new indication of the drug. In fact, drug repurposing is a very popular research area: Gefitinib, for example, was approved by the FDA for treatment of NSCLC (non-small cell lung cancer) but is now involved in over 400+ clinical trials involving several types of cancers as well as many other diseases. This includes lung cancer, head and neck cancer, salivary gland cancer, esophageal cancer, breast cancer, kidney cancer, ovarian cancer, colon cancer, and many other diseases. R&D can certainly apply for a patent, but it is not a concern for Selleck nor the original drug company for matters of re-purposing and publication.

Q2 How long does it take, using Selleck Libraries, to publish an article and what kind of article?

According to Research Journals, it takes an average of 2 years to publish an article while the impact factor ranges from 3 to 39.

Q3 Interested in a Selleck Library, but don’t have the time or energy for experiments?

Selleck provides free user training through lectures, technical support and other ways to help. For those without time, we can provide references to our network of technical service providers.

Q4 Why do some Selleck Libraries contain fewer compounds than other companies?

Selleck libraries neither contain any controlled products or other unrelated products nor do we add both salt and non-salt versions in the same library collection. We only consider effective molecules: Selleck libraries contain more compounds, which are updated every three months and are the best choice.