## **Resource Summary Report**

Generated by <u>NIF</u> on May 13, 2025

# **LINCS Project**

RRID:SCR\_016486 Type: Tool

### **Proper Citation**

LINCS Project (RRID:SCR\_016486)

## **Resource Information**

URL: http://www.lincsproject.org/

Proper Citation: LINCS Project (RRID:SCR\_016486)

**Description:** Project to create network based understanding of biology by cataloging changes in gene expression and other cellular processes when cells are exposed to genetic and environmental stressors. Program to develop therapies that might restore pathways and networks to their normal states. Has LINCS Data Coordination and Integration Center and six Data and Signature Generation Centers: Drug Toxicity Signature Generation Center, HMS LINCS Center, LINCS Center for Transcriptomics, LINCS Proteomic Characterization Center for Signaling and Epigenetics, MEP LINCS Center, and NeuroLINCS Center.

#### Abbreviations: LINCS

**Synonyms:** LINCS, Library of Integrated Network based Cellular Signatures, LINCS Program

**Resource Type:** portal, consortium, data or information resource, project portal, organization portal, database

Defining Citation: PMID:29199020

**Keywords:** data integration, network biology, gene expression, L1000, MCF10A, MEMA, P100, LINCS program, LINCS project, systems biology, systems pharmacology, FASEB list

Related Condition: cancer, heart disease, neurodegenerative disorder

Funding: NIH Common Fund ; NHLBI U54 HL127624; NHLBI U54 HL127366; NHLBI U54 HL127365; NHGRI U54 HG008100; NHGRI U54 HG008097; NHGRI U54 HG008098; NINDS U54 NS091046

Availability: Free, Freely available

Resource Name: LINCS Project

Resource ID: SCR\_016486

Alternate IDs: SCR\_016487

**Record Creation Time:** 20220129T080330+0000

Record Last Update: 20250513T061751+0000

#### **Ratings and Alerts**

No rating or validation information has been found for LINCS Project.

No alerts have been found for LINCS Project.

#### Data and Source Information

Source: SciCrunch Registry

#### **Usage and Citation Metrics**

We found 40 mentions in open access literature.

Listed below are recent publications. The full list is available at <u>NIF</u>.

Chaib S, et al. (2024) The efficacy of chemotherapy is limited by intratumoral senescent cells expressing PD-L2. Nature cancer, 5(3), 448.

lida M, et al. (2024) A network-based trans-omics approach for predicting synergistic drug combinations. Communications medicine, 4(1), 154.

Pognan F, et al. (2023) The evolving role of investigative toxicology in the pharmaceutical industry. Nature reviews. Drug discovery, 22(4), 317.

Parra-Rivas LA, et al. (2023) Serine-129 phosphorylation of ?-synuclein is an activitydependent trigger for physiologic protein-protein interactions and synaptic function. Neuron, 111(24), 4006. Namba S, et al. (2022) From drug repositioning to target repositioning: prediction of therapeutic targets using genetically perturbed transcriptomic signatures. Bioinformatics (Oxford, England), 38(Suppl 1), i68.

Iwata M, et al. (2022) Regulome-based characterization of drug activity across the human diseasome. NPJ systems biology and applications, 8(1), 44.

Iwata M, et al. (2022) Pathway trajectory analysis with tensor imputation reveals druginduced single-cell transcriptomic landscape. Nature computational science, 2(11), 758.

Lu L, et al. (2022) Recent computational drug repositioning strategies against SARS-CoV-2. Computational and structural biotechnology journal, 20, 5713.

Fustero-Torre C, et al. (2021) Beyondcell: targeting cancer therapeutic heterogeneity in single-cell RNA-seq data. Genome medicine, 13(1), 187.

Sinkala M, et al. (2021) Integrated molecular characterisation of the MAPK pathways in human cancers reveals pharmacologically vulnerable mutations and gene dependencies. Communications biology, 4(1), 9.

Stanfill AG, et al. (2021) Enhancing Research Through the Use of the Genotype-Tissue Expression (GTEx) Database. Biological research for nursing, 23(3), 533.

Han L, et al. (2021) Modeling drug response using network-based personalized treatment prediction (NetPTP) with applications to inflammatory bowel disease. PLoS computational biology, 17(2), e1008631.

He B, et al. (2021) A Review of Current In Silico Methods for Repositioning Drugs and Chemical Compounds. Frontiers in oncology, 11, 711225.

Liu Z, et al. (2021) Unraveling Gene Fusions for Drug Repositioning in High-Risk Neuroblastoma. Frontiers in pharmacology, 12, 608778.

Mapelli SN, et al. (2020) A Novel Prostate Cell Type-Specific Gene Signature to Interrogate Prostate Tumor Differentiation Status and Monitor Therapeutic Response (Running Title: Phenotypic Classification of Prostate Tumors). Cancers, 12(1).

Liu K, et al. (2020) Broad-Spectrum Profiling of Drug Safety via Learning Complex Network. Clinical pharmacology and therapeutics, 107(6), 1373.

Kunc V, et al. (2020) On tower and checkerboard neural network architectures for gene expression inference. BMC genomics, 21(Suppl 5), 454.

Sailem HZ, et al. (2020) KCML: a machine-learning framework for inference of multi-scale gene functions from genetic perturbation screens. Molecular systems biology, 16(3), e9083.

Chen X, et al. (2020) iGMDR: Integrated Pharmacogenetic Resource Guide to Cancer Therapy and Research. Genomics, proteomics & bioinformatics, 18(2), 150.

Zhu L, et al. (2020) Drug Repositioning for Noonan and LEOPARD Syndromes by Integrating Transcriptomics With a Structure-Based Approach. Frontiers in pharmacology, 11, 927.