Resource Summary Report

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Database of Interacting Proteins (DIP)

RRID:SCR 003167

Type: Tool

Proper Citation

Database of Interacting Proteins (DIP) (RRID:SCR_003167)

Resource Information

URL: http://dip.doe-mbi.ucla.edu/

Proper Citation: Database of Interacting Proteins (DIP) (RRID:SCR_003167)

Description: Database to catalog experimentally determined interactions between proteins combining information from a variety of sources to create a single, consistent set of proteinprotein interactions that can be downloaded in a variety of formats. The data were curated. both, manually and also automatically using computational approaches that utilize the the knowledge about the protein-protein interaction networks extracted from the most reliable, core subset of the DIP data. Because the reliability of experimental evidence varies widely, methods of quality assessment have been developed and utilized to identify the most reliable subset of the interactions. This CORE set can be used as a reference when evaluating the reliability of high-throughput protein-protein interaction data sets, for development of prediction methods, as well as in the studies of the properties of protein interaction networks. Tools are available to analyze, visualize and integrate user's own experimental data with the information about protein-protein interactions available in the DIP database. The DIP database lists protein pairs that are known to interact with each other. By interact they mean that two amino acid chains were experimentally identified to bind to each other. The database lists such pairs to aid those studying a particular protein-protein interaction but also those investigating entire regulatory and signaling pathways as well as those studying the organization and complexity of the protein interaction network at the cellular level. Registration is required to gain access to most of the DIP features. Registration is free to the members of the academic community. Trial accounts for the commercial users are also available.

Abbreviations: DIP

Synonyms: , Database of Interacting Proteins, DIP, Database of Interacting Proteins (DIP)

Resource Type: analysis service resource, storage service resource, data or information resource, production service resource, data analysis service, database, data repository, service resource

Defining Citation: PMID:14681454

Keywords: blast, cellular network, ligand-receptor complex, ligand, network, protein, protein interaction, protein ligand, protein-protein interaction, protein receptor, receptor, sequence, interaction, regulatory pathway, signaling pathway, protein binding, bio.tools, FASEB list

Funding Agency: NIGMS

Availability: Account required, Creative Commons Attribution-NoDerivs License, Trial accounts for commercial users are available, Terms of Use, The community can contribute to this resource

Resource Name: Database of Interacting Proteins (DIP)

Resource ID: SCR_003167

Alternate IDs: OMICS_01905, nif-0000-00569, biotools:dip

Alternate URLs: https://dip.doe-mbi.ucla.edu/dip/Main.cgi, https://bio.tools/dip

Ratings and Alerts

No rating or validation information has been found for Database of Interacting Proteins (DIP).

No alerts have been found for Database of Interacting Proteins (DIP).

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We found 143 mentions in open access literature.

Listed below are recent publications. The full list is available at <u>FDI Lab - SciCrunch.org</u>.

Wang X, et al. (2023) Cross-Talk between N6-Methyladenosine and Their Related RNAs Defined a Signature and Confirmed m6A Regulators for Diagnosis of Endometriosis. International journal of molecular sciences, 24(2).

Luo L, et al. (2023) Identification of kukoamine a as an anti-osteoporosis drug target using network pharmacology and experiment verification. Molecular medicine (Cambridge, Mass.),

Jiang Y, et al. (2022) Senkyunolide H protects PC12 cells from OGD/R-induced injury via cAMP-PI3K/AKT signaling pathway. Journal of ethnopharmacology, 282, 114659.

Sadeghi M, et al. (2022) IncRNA-miRNA-mRNA ceRNA Network Involved in Sheep Prolificacy: An Integrated Approach. Genes, 13(8).

Asim MN, et al. (2022) ADH-PPI: An attention-based deep hybrid model for protein-protein interaction prediction. iScience, 25(10), 105169.

Kang P, et al. (2021) A Network Pharmacology and Molecular Docking Strategy to Explore Potential Targets and Mechanisms Underlying the Effect of Curcumin on Osteonecrosis of the Femoral Head in Systemic Lupus Erythematosus. BioMed research international, 2021, 5538643.

Li Y, et al. (2021) Robust and accurate prediction of protein-protein interactions by exploiting evolutionary information. Scientific reports, 11(1), 16910.

Xu Y, et al. (2021) Identification of the Key Role of NF-?B Signaling Pathway in the Treatment of Osteoarthritis With Bushen Zhuangjin Decoction, a Verification Based on Network Pharmacology Approach. Frontiers in pharmacology, 12, 637273.

Xu H, et al. (2021) A comprehensive review of integrative pharmacology-based investigation: A paradigm shift in traditional Chinese medicine. Acta pharmaceutica Sinica. B, 11(6), 1379.

Jin Z, et al. (2021) Protective effect of Qingre Huoxue decoction against myocardial infarction via PI3K/Akt autophagy pathway based on UPLC-MS, network pharmacology, and in vivo evidence. Pharmaceutical biology, 59(1), 1607.

Yan VKC, et al. (2021) Drug Repurposing for the Treatment of COVID-19: A Knowledge Graph Approach. Advanced therapeutics, 4(7), 2100055.

Yoshioka H, et al. (2021) Overexpression of miR-1306-5p, miR-3195, and miR-3914 Inhibits Ameloblast Differentiation through Suppression of Genes Associated with Human Amelogenesis Imperfecta. International journal of molecular sciences, 22(4).

Yang S, et al. (2021) Mechanisms and molecular targets of the Yu-Ping-Feng powder for allergic rhinitis, based on network pharmacology. Medicine, 100(35), e26929.

Chen S, et al. (2021) Establishment of an anti-inflammation-based bioassay for the quality control of the 13-component TCM formula (Lianhua Qingwen). Pharmaceutical biology, 59(1), 537.

Yao H, et al. (2021) Network Pharmacology-Based Approach to Comparatively Predict the Active Ingredients and Molecular Targets of Compound Xueshuantong Capsule and Hexuemingmu Tablet in the Treatment of Proliferative Diabetic Retinopathy. Evidence-based complementary and alternative medicine: eCAM, 2021, 6642600.

Ma F, et al. (2021) Applications and analytical tools of cell communication based on ligand-receptor interactions at single cell level. Cell & bioscience, 11(1), 121.

Muzio G, et al. (2021) Biological network analysis with deep learning. Briefings in bioinformatics, 22(2), 1515.

Potrony M, et al. (2021) DNA Repair and Immune Response Pathways Are Deregulated in Melanocyte-Keratinocyte Co-cultures Derived From the Healthy Skin of Familial Melanoma Patients. Frontiers in medicine, 8, 692341.

Kuleshov MV, et al. (2021) KEA3: improved kinase enrichment analysis via data integration. Nucleic acids research, 49(W1), W304.

Wu L, et al. (2021) Identification of IFN-Induced Transmembrane Protein 1 With Prognostic Value in Pancreatic Cancer Using Network Module-Based Analysis. Frontiers in oncology, 11, 626883.