

# Resource Summary Report

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## SIDER

RRID:SCR\_004321

Type: Tool

### Proper Citation

SIDER (RRID:SCR\_004321)

### Resource Information

**URL:** <http://sideeffects.embl.de/>

**Proper Citation:** SIDER (RRID:SCR\_004321)

**Description:** Database containing information on marketed medicines and their recorded adverse drug reactions. The information is extracted from public documents and package inserts. The available information include side effect frequency, drug and side effect classifications as well as links to further information, for example drug-target relations. The SIDER Side Effect Resource represents an effort to aggregate dispersed public information on side effects. To our knowledge, no such resource exist in machine-readable form despite the importance of research on drugs and their effects. The creation of this resource was motivated by the many requests for data that we received related to our paper (Campillos, Kuhn et al., Science, 2008, 321(5886):263-6.) on the utilization of side effects for drug target prediction. Inclusion of side effects as readouts for drug treatment should have many applications and we hope to be able to enhance the respective research with this resource. You may browse the drugs by name, browse the side effects by name, download the current version of SIDER, or use the search interface.

**Abbreviations:** SIDER

**Synonyms:** Side Effect Resource, SIDER: Side Effect Resource

**Resource Type:** data or information resource, database

**Defining Citation:** [PMID:20087340](#)

**Keywords:** medicine, drug, side effect, adverse drug reaction, drug-target, phenotype, bio.tools, FASEB list

**Availability:** Except as otherwise noted, Creative Commons Attribution-NonCommercial-ShareAlike License, v3, Commercial use requires permission

**Resource Name:** SIDER

**Resource ID:** SCR\_004321

**Alternate IDs:** r3d100012791, nlx\_33359, OMICS\_01588, biotools:sider

**Alternate URLs:** <https://bio.tools/sider>, <https://doi.org/10.17616/R3J226>

**Record Creation Time:** 20220129T080223+0000

**Record Last Update:** 20260829T182926+0000

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## Ratings and Alerts

No rating or validation information has been found for SIDER.

No alerts have been found for SIDER.

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## Data and Source Information

**Source:** [SciCrunch Registry](#)

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## Usage and Citation Metrics

We found 410 mentions in open access literature.

**Listed below are recent publications.** The full list is available at [NIF](#).

Cai C, et al. (2026) A unified multi-scale deep learning framework for molecular property prediction that bridges molecular structures and fingerprinting. Communications chemistry, 9(1).

Hong H, et al. (2026) A hierarchical interaction message net for accurate molecular property prediction. Communications chemistry, 9(1).

Patidar P, et al. (2026) An integrative analysis of genetic factors reveals dysregulation of cytokines, immune signaling, and T cell activity as the underlying immune mechanisms in autoimmune immune thrombocytopenia. Thrombosis journal, 24(1), 25.

Nygren S, et al. (2026) RAG-based architectures for drug side effect retrieval using compact LLMs. Scientific reports, 16(1).

Zhang Y, et al. (2026) MolGramTreeNet: A multimodal molecular property prediction model via grammar tree-constrained molecular representation. iScience, 29(3), 114928.

Xie C, et al. (2026) MMPCS: multi-view molecular pretraining based on consistency information and specific information. Bioinformatics (Oxford, England), 42(2).

Zhang W, et al. (2026) Task-specific pre-training for molecular property prediction. Briefings in bioinformatics, 27(1).

Wu S, et al. (2026) Enhanced drug disease association prediction through multimodal data integration and meta path guided global local feature fusion. Scientific reports, 16(1).

Pala MA, et al. (2026) Graph-Aware AURALSTM: An Attentive Unified Representation Architecture with BiLSTM for Enhanced Molecular Property Prediction. *Molecular diversity*, 30(2), 1709.

Zheng W, et al. (2026) Signal detection of drug-induced esophageal ulcer across 20 years of real-world study: Focus on 49 high-risk medicines. *The Journal of international medical research*, 54(3), 3000605261426764.

Fan H, et al. (2026) A Bond-Level Sequence Framework for Molecular Representation Learning with Structural Constraints. *Molecules (Basel, Switzerland)*, 31(11).

Zhao J, et al. (2026) MolProphecy: Bridging medicinal chemists' knowledge and molecular pre-trained models via a multi-modal framework. *Journal of advanced research*, 85, 577.

Khodadadi AghGhaleh M, et al. (2026) ConvAHKG: Action-based hybrid knowledge graph with a dual-channel convolutional approach for drug repurposing. *Scientific reports*, 16(1).

Rjoob K, et al. (2026) A multimodal vision knowledge graph of cardiovascular disease. *Nature cardiovascular research*, 5(1), 18.

Huang T, et al. (2026) Explainable drug side effect prediction in central neural system via biologically informed graph neural network. *Translational psychiatry*, 16(1).

Mun V, et al. (2026) CheMLT-F: multitask learning in biochemistry through transformer fusion. *Journal of cheminformatics*, 18(1).

Yang L, et al. (2026) Multimodal Collaborative Modeling of Molecular Structures and Biomedical Text for Accurate Drug-Drug Interaction Extraction. *Biomedicines*, 14(6).

Jeon W, et al. (2025) Predicting Drug-Side Effect Relationships From Parametric Knowledge Embedded in Biomedical BERT Models: Methodological Study With a Natural Language Processing Approach. *JMIR medical informatics*, 13, e67513.

He M, et al. (2025) Prediction of adverse drug reactions based on pharmacogenomics combination features: a preliminary study. *Frontiers in pharmacology*, 16, 1448106.

Abbasi SH, et al. (2025) Acute cardiovascular effects associated with the use of prescription medications: A Danish nationwide screening study. *British journal of clinical pharmacology*, 91(7), 1947.