# Resource Summary Report

Generated by PRECISE-TBI on Aug 2, 2026

## MedGen

RRID:SCR_000111
Type: Tool

### Proper Citation

MedGen (RRID:SCR_000111)

### Resource Information

**URL:** http://www.ncbi.nlm.nih.gov/medgen/

**Proper Citation:** MedGen (RRID:SCR_000111)

**Description:** A database of organized information related to human medical genetics, such as attributes of conditions with a genetic contribution.

**Abbreviations:** MedGen

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

**Defining Citation:** PMID:32329672

**Keywords:** medical genetics, medical, genetics, disease, clinical

**Funding:** NLM 1ZIHLM200888-05

**Resource Name:** MedGen

**Resource ID:** SCR_000111

**Alternate IDs:** nlx_156941, OMICS_01549

**Record Creation Time:** 20220129T080159+0000

**Record Last Update:** 20260801T190853+0000

### Ratings and Alerts

No rating or validation information has been found for MedGen.

No alerts have been found for MedGen.

### Data and Source Information

**Source:** SciCrunch Registry

## Usage and Citation Metrics

We found 6 mentions in open access literature.

**Listed below are recent publications.** The full list is available at PRECISE-TBI .


Hayman GT, et al. (2016) The Disease Portals, disease-gene annotation and the RGD disease ontology at the Rat Genome Database. Database : the journal of biological databases and curation, 2016.

Pratte-Santos R, et al. (2016) Analysis of chromosomal abnormalities by CGH-array in patients with dysmorphic and intellectual disability with normal karyotype. Einstein (Sao Paulo, Brazil), 14(1), 30.

Bornstein AT, et al. (2016) Tracking medical genetic literature through machine learning. Molecular genetics and metabolism, 118(4), 255.

Sayito??lu M, et al. (2016) Clinical Interpretation of Genomic Variations. Turkish journal of haematology : official journal of Turkish Society of Haematology, 33(3), 172.

, et al. (2015) Database resources of the National Center for Biotechnology Information. Nucleic acids research, 43(Database issue), D6.

Landrum MJ, et al. (2014) ClinVar: public archive of relationships among sequence variation and human phenotype. Nucleic acids research, 42(Database issue), D980.