

# Resource Summary Report

Generated by SPARC SAWG on Sep 17, 2026

## deCODE genetics

RRID:SCR\_003334

Type: Tool

### Proper Citation

deCODE genetics (RRID:SCR\_003334)

### Resource Information

**URL:** <http://www.decode.com/>

**Proper Citation:** deCODE genetics (RRID:SCR\_003334)

**Description:** A biopharmaceutical company applying its discoveries in human genetics to develop drugs and diagnostics for common diseases. They specialize in gene discovery - their population approach and resources have enabled them to isolate key genes contributing to major public health challenges from cardiovascular disease to cancer. The company's genotyping capacity is now one of the highest in the world. They have a large population-based biobank containing whole blood and DNA samples with extensive relevant phenotypic information from around 120.000 Icelanders. In the company's work in more than 50 disease projects, their statistical and informatics departments have established themselves in data processing and analysis. deCODE genetics is widely recognized as a center of excellence in genetic research.

**Abbreviations:** deCODE

**Synonyms:** Islensk Erfdagreining EHF, Islensk Erfdagreining

**Resource Type:** commercial organization

**Keywords:** biopharmaceutical, genetics, drug, diagnostic, genotyping, phenotype, data processing, analysis, genetic variant, risk factor, genome, blood, dna, biobank, single nucleotide polymorphism

**Related Condition:** Schizophrenia, Cardiovascular disease, Cancer, Type 2 diabetes, Atrial fibrillation, Heart attack

**Availability:** Free, Freely available

**Resource Name:** deCODE genetics

**Resource ID:** SCR\_003334

**Alternate IDs:** nif-0000-31959, ISNI: 0000 0004 0618 6889, grid.421812.c, Wikidata: Q493712

**Alternate URLs:** <https://ror.org/04dzdm737>

**Record Creation Time:** 20220129T080218+0000

**Record Last Update:** 20260912T195556+0000

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## Ratings and Alerts

No rating or validation information has been found for deCODE genetics.

No alerts have been found for deCODE genetics.

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## Data and Source Information

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

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## Usage and Citation Metrics

We found 67 mentions in open access literature.

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

Nie H, et al. (2026) Intracellular Osteopontin of Macrophages Promotes Carotid Plaques Formation by Inducing Foam Cells and Releasing Proinflammatory Cytokines. Journal of the American Heart Association, 15(6), e041336.

Zhang H, et al. (2026) Mendelian randomization implicates circulating plasma proteins in gout risk and identifies candidate therapeutic targets. International journal of immunopathology and pharmacology, 40, 3946320261428859.

Cong L, et al. (2026) Causal effects of mental disorders on stroke subtypes: A proteome-wide Mendelian randomization and mediation analysis. Medicine, 105(23), e49108.

Luo X, et al. (2026) Mitochondrial Hub Genes in Rheumatoid Arthritis Identified by Machine Learning and Mendelian Randomization with Experimental Validation. Journal of inflammation research, 19, 576901.

Li H, et al. (2025) Multi-ancestry sequencing-based genome-wide association study of C-reactive protein in 513,273 genomes. Nature communications, 16(1), 3892.

Lv R, et al. (2025) Investigating causal relationships between plasma proteins and lung adenocarcinoma: result from proteomics and Mendelian randomization study. Discover oncology, 16(1), 42.

Hu M, et al. (2025) Identifying Key Biomarkers Related to Immune Response in the Progression of Diabetic Kidney Disease: Mendelian Randomization Combined With Comprehensive Transcriptomics and Single-Cell Sequencing Analysis. Journal of

inflammation research, 18, 949.

Li R, et al. (2025) Mendelian randomization study on the causal relationships among fasting blood glucose, plasma proteins, and squamous cell lung cancer. *Discover oncology*, 16(1), 588.

Chen Q, et al. (2025) Exploring the causal relationship between iron status and subarachnoid hemorrhage based on two sample mendelian randomization. *Scientific reports*, 15(1), 15549.

Chen W, et al. (2025) Causal Effect of Obstructive Sleep Apnea on Sick Sinus Syndrome: A Bidirectional Mendelian Randomization Study. *Nature and science of sleep*, 17, 689.

Akdeniz BC, et al. (2025) Polygenic Hazard Score for Predicting Age-associated Risk of Alzheimer's Disease in European Populations: Development and Validation. *medRxiv : the preprint server for health sciences*.

Fu J, et al. (2025) CCNE1 Exerts a Protective Effect on Parkinson's Disease by Regulating Ferroptosis-Related Proteins. *Brain and behavior*, 15(12), e71110.

Yue H, et al. (2025) Identification of Potential Therapeutic Targets for Sepsis Using Mendelian Randomization and Integrated eQTL/pQTL Analysis. *International journal of general medicine*, 18, 6137.

Sun H, et al. (2025) A multi-omics target study for glioblastoma multiforme (GBM) based on Mendelian randomization analysis. *IBRO neuroscience reports*, 18, 400.

Xu G, et al. (2025) Causality of Childhood and Adult Body Mass Index on Sick Sinus Syndrome: A Mendelian Randomization Study. *Cureus*, 17(3), e80913.

Malomane DK, et al. (2025) Patterns of population structure and genetic variation within the Saudi Arabian population. *bioRxiv : the preprint server for biology*.

Fan W, et al. (2025) Mendelian randomization analysis of plasma proteins reveals potential novel tumor markers for gastric cancer. *Scientific reports*, 15(1), 3537.

Fu J, et al. (2025) Assessing the causal relationship between the plasma proteome and epilepsy: A Mendelian randomization study. *Epilepsia open*, 10(6), 1966.

Chen J, et al. (2025) Identification of novel therapeutic targets for liver fibrosis, cirrhosis, and hepatocellular carcinoma via dual-omics analysis and preliminary exploration of regulatory mechanisms. *Clinical and experimental hepatology*, 11(4), 360.

Xu H, et al. (2024) Exploring the causal effect of complement and IgA nephropathy-a Mendelian randomization study. *Renal failure*, 46(2), 2436632.