

Resource Summary Report

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Cardiovascular Disease Knowledge Portal

RRID:SCR_016536

Type: Tool

Proper Citation

Cardiovascular Disease Knowledge Portal (RRID:SCR_016536)

Resource Information

URL: <http://www.broadcvdi.org/>

Proper Citation: Cardiovascular Disease Knowledge Portal (RRID:SCR_016536)

Description: Platform for analysis of the genetics of cardiovascular disease.Used for searching and analysis of human genetic information linked to myocardial infarction, atrial fibrillation and related traits while protecting the integrity and confidentiality of the data.

Resource Type: disease-related portal, data or information resource, database, topical portal, portal

Keywords: genetic, data, cardiovascular, disease, human

Related Condition: cardiovascular disease, myocardial infarction, atrial fibrillation

Funding: Accelerating Medicines Partnership in Type 2 Diabetes ;
National Institute of Cardiovascular Diseases and Stroke

Availability: Free, Available for download, Google ID required, Tutorial available

Resource Name: Cardiovascular Disease Knowledge Portal

Resource ID: SCR_016536

Record Creation Time: 20220129T080331+0000

Record Last Update: 20260729T044420+0000

Ratings and Alerts

No rating or validation information has been found for Cardiovascular Disease Knowledge Portal.

No alerts have been found for Cardiovascular Disease Knowledge Portal.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 28 mentions in open access literature.

Listed below are recent publications. The full list is available at [kravitz2](#) .

Abou Daya F, et al. (2025) Identifying links between cardiovascular disease and insomnia by modeling genes from a pleiotropic locus. *Disease models & mechanisms*, 18(5).

Wei S, et al. (2025) Mendelian randomization provides a multi-omics perspective on the regulation of genes involved in ribosome biogenesis in relation to cardiac structure and function. *Clinical epigenetics*, 17(1), 42.

Rämö JT, et al. (2023) The Cardiovascular Impact and Genetics of Pericardial Adiposity. *medRxiv : the preprint server for health sciences*.

Mou X, et al. (2023) Causal influence of muscle weakness on cardiometabolic diseases and osteoporosis. *Scientific reports*, 13(1), 19974.

Khurshid S, et al. (2023) Clinical and genetic associations of deep learning-derived cardiac magnetic resonance-based left ventricular mass. *Nature communications*, 14(1), 1558.

Jiang W, et al. (2023) Mendelian Randomization Analysis Provides Insights into the Pathogenesis of Serum Levels of Branched-Chain Amino Acids in Cardiovascular Disease. *Metabolites*, 13(3).

Li M, et al. (2023) Genetic Evidence for Causal Association Between Atrial Fibrillation and Dementia: A Mendelian Randomization Study. *Journal of the American Heart Association*, 12(16), e029623.

Actkins KV, et al. (2023) Sex modifies the effect of genetic risk scores for polycystic ovary syndrome on metabolic phenotypes. *PLoS genetics*, 19(5), e1010764.

Pirruccello JP, et al. (2022) Deep learning enables genetic analysis of the human thoracic aorta. *Nature genetics*, 54(1), 40.

Hindy G, et al. (2022) Rare coding variants in 35 genes associate with circulating lipid levels-A multi-ancestry analysis of 170,000 exomes. *American journal of human genetics*, 109(1), 81.

Chen L, et al. (2022) Obstructive sleep apnea and atrial fibrillation: insights from a bidirectional Mendelian randomization study. *BMC medical genomics*, 15(1), 28.

Zhang M, et al. (2022) Causal associations of circulating adiponectin with cardiometabolic diseases and osteoporotic fracture. *Scientific reports*, 12(1), 6689.

Zhang T, et al. (2021) The associations of plasma phospholipid arachidonic acid with cardiovascular diseases: A Mendelian randomization study. *EBioMedicine*, 63, 103189.

He B, et al. (2021) Causal Effect of Serum Magnesium on Osteoporosis and Cardiometabolic Diseases. *Frontiers in nutrition*, 8, 738000.

Li H, et al. (2021) Exploring the Pleiotropic Genes and Therapeutic Targets Associated with Heart Failure and Chronic Kidney Disease by Integrating metaCCA and SGLT2 Inhibitors' Target Prediction. *BioMed research international*, 2021, 4229194.

Lumbers RT, et al. (2021) The genomics of heart failure: design and rationale of the HERMES consortium. *ESC heart failure*, 8(6), 5531.

Liu Y, et al. (2021) Genome-wide association study of neck circumference identifies sex-specific loci independent of generalized adiposity. *International journal of obesity* (2005), 45(7), 1532.

Chan II, et al. (2021) The total and direct effects of systolic and diastolic blood pressure on cardiovascular disease and longevity using Mendelian randomisation. *Scientific reports*, 11(1), 21799.

Moksnes MR, et al. (2021) Genome-wide association study of cardiac troponin I in the general population. *Human molecular genetics*, 30(21), 2027.

Kwok MK, et al. (2021) Mendelian randomization study on atrial fibrillation and cardiovascular disease subtypes. *Scientific reports*, 11(1), 18682.