

Resource Summary Report

Generated by PRECISE-TBI on Sep 17, 2026

ParseCNV

RRID:SCR_000355

Type: Tool

Proper Citation

ParseCNV (RRID:SCR_000355)

Resource Information

URL: <http://parsecnv.sourceforge.net/>

Proper Citation: ParseCNV (RRID:SCR_000355)

Description: Software that takes CNV calls as input and creates SNP based statistics for CNV occurrence in cases and controls then calls CNVRs based on neighboring SNPs of similar significance.

Resource Type: software resource

Defining Citation: [PMID:23293001](#)

Keywords: standalone software

Availability: Free, Available for download, Freely available

Resource Name: ParseCNV

Resource ID: SCR_000355

Alternate IDs: OMICS_02566

Record Creation Time: 20220129T080201+0000

Record Last Update: 20260912T195507+0000

Ratings and Alerts

No rating or validation information has been found for ParseCNV.

No alerts have been found for ParseCNV.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 1 mentions in open access literature.

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

Dajani R, et al. (2015) CNV Analysis Associates AKNAD1 with Type-2 Diabetes in Jordan Subpopulations. Scientific reports, 5, 13391.