# Resource Summary Report

Generated by PRECISE-TBI on Aug 1, 2026

## ParseCNV

RRID:SCR_000355
Type: Tool

### Proper Citation

ParseCNV (RRID:SCR_000355)

### Resource Information

**URL:** http://parsecnv.sourceforge.net/

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**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:** 20260725T190443+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.