# Resource Summary Report

Generated by PRECISE-TBI on Sep 18, 2026

## WaveCNV

RRID:SCR_012244
Type: Tool

### Proper Citation

WaveCNV (RRID:SCR_012244)

### Resource Information

**URL:** https://github.com/WaveCNV

**Proper Citation:** WaveCNV (RRID:SCR_012244)

**Description:** Cancer specific CNV caller for Next Generation sequence.

**Abbreviations:** WaveCNV

**Resource Type:** software resource

**Defining Citation:** PMID:24192544

**Keywords:** cancer

**Resource Name:** WaveCNV

**Resource ID:** SCR_012244

**Alternate IDs:** OMICS_00353

**Record Creation Time:** 20220129T080309+0000

**Record Last Update:** 20260912T195746+0000

### Ratings and Alerts

No rating or validation information has been found for WaveCNV.

No alerts have been found for WaveCNV.

### Data and Source Information

**Source:** SciCrunch Registry

## Usage and Citation Metrics

We found 2 mentions in open access literature.

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

Chien J, et al. (2015) TP53 mutations, tetraploidy and homologous recombination repair defects in early stage high-grade serous ovarian cancer. Nucleic acids research, 43(14), 6945.

Holt C, et al. (2014) WaveCNV: allele-specific copy number alterations in primary tumors and xenograft models from next-generation sequencing. Bioinformatics (Oxford, England), 30(6), 768.