

Resource Summary Report

Generated by [NIF](#) on Aug 10, 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: 20260808T190009+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 [NIF](#) .

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.