

# Resource Summary Report

Generated by [SPARC SAWG](#) on Jul 30, 2026

## SomaticCall

RRID:SCR\_001196

Type: Tool

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### Proper Citation

SomaticCall (RRID:SCR\_001196)

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### Resource Information

**URL:** <http://www.broadinstitute.org/science/programs/genome-biology/computational-rd/somaticcall-manual>

**Proper Citation:** SomaticCall (RRID:SCR\_001196)

**Description:** Software program that finds single-base differences (substitutions) between sequence data from tumor and matched normal samples. It is designed to be highly stringent, so as to achieve a low false positive rate. It takes as input a BAM file for each sample, and produces as output a list of differences (somatic mutations). Note: This software package is no longer supported and information on this page is provided for archival purposes only.

**Abbreviations:** SomaticCall

**Resource Type:** software resource

**Keywords:** somatic mutation, substitution, sequence, bam, mutation

**Related Condition:** Tumor, Cancer, Normal

**Resource Name:** SomaticCall

**Resource ID:** SCR\_001196

**Alternate IDs:** OMICS\_02155

**Record Creation Time:** 20220129T080206+0000

**Record Last Update:** 20260725T190457+0000

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### Ratings and Alerts

No rating or validation information has been found for SomaticCall.

No alerts have been found for SomaticCall.

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## Data and Source Information

**Source:** [SciCrunch Registry](#)

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## Usage and Citation Metrics

We have not found any literature mentions for this resource.