

# Resource Summary Report

Generated by [Kravitz](#) on Sep 10, 2026

## exomeSuite

RRID:SCR\_000129

Type: Tool

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### Proper Citation

exomeSuite (RRID:SCR\_000129)

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### Resource Information

**URL:** <http://sourceforge.net/projects/exomesuite/>

**Proper Citation:** exomeSuite (RRID:SCR\_000129)

**Description:** THIS RESOURCE IS NO LONGER IN SERVICE. Documented on July 31,2025. A software application designed to analyze variant call files from next generation sequencing data to identify variants causing disease.

**Resource Type:** software resource

**Defining Citation:** [PMID:24603341](#)

**Keywords:** standalone software, c, matlab

**Availability:** THIS RESOURCE IS NO LONGER IN SERVICE

**Resource Name:** exomeSuite

**Resource ID:** SCR\_000129

**Alternate IDs:** OMICS\_04839

**Record Creation Time:** 20220129T080159+0000

**Record Last Update:** 20260905T132413+0000

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### Ratings and Alerts

No rating or validation information has been found for exomeSuite.

No alerts have been found for exomeSuite.

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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.