

Resource Summary Report

Generated by [RRID](#) on Aug 4, 2026

SomaticSeq

RRID:SCR_024891

Type: Tool

Proper Citation

SomaticSeq (RRID:SCR_024891)

Resource Information

URL: <https://github.com/bioinform/somaticseq>

Proper Citation: SomaticSeq (RRID:SCR_024891)

Description: Software accurate somatic mutation detection pipeline implementing stochastic boosting algorithm to produce somatic mutation calls for both single nucleotide variants and small insertions and deletions. NGS variant calling and classification.

Resource Type: software resource, software toolkit

Defining Citation: [PMID:26381235](#)

Keywords: NGS variant calling and classification, somatic mutation detection, somatic mutation calls, single nucleotide variants, detect somatic mutations,

Funding: NIGMS R01 GM109836;
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Resource Name: SomaticSeq

Resource ID: SCR_024891

License: BSD-2-Clause

Record Creation Time: 20240117T050246+0000

Record Last Update: 20260805T044537+0000

Ratings and Alerts

No rating or validation information has been found for SomaticSeq.

No alerts have been found for SomaticSeq.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 6 mentions in open access literature.

Listed below are recent publications. The full list is available at [RRID](#) .

Li D, et al. (2025) Augmenting precision medicine via targeted RNA-Seq detection of expressed mutations. NPJ precision oncology, 9(1), 182.

Iser F, et al. (2024) Cerebrospinal Fluid cfDNA Sequencing for Classification of Central Nervous System Glioma. Clinical cancer research : an official journal of the American Association for Cancer Research, 30(14), 2974.

Starrett GJ, et al. (2024) Multiomics Profiling Distinguishes Sebaceous Carcinoma from Benign Sebaceous Neoplasms and Provides Insight into the Genetic Evolution of Sebaceous Carcinogenesis. Clinical cancer research : an official journal of the American Association for Cancer Research, 30(21), 4887.

Amin SB, et al. (2020) Comparative Molecular Life History of Spontaneous Canine and Human Gliomas. Cancer cell, 37(2), 243.

Sahraeian SME, et al. (2019) Deep convolutional neural networks for accurate somatic mutation detection. Nature communications, 10(1), 1041.

Fang LT, et al. (2015) An ensemble approach to accurately detect somatic mutations using SomaticSeq. Genome biology, 16(1), 197.