

Resource Summary Report

Generated by [RRID](#) on Aug 2, 2026

ShortFuse

RRID:SCR_001107

Type: Tool

Proper Citation

ShortFuse (RRID:SCR_001107)

Resource Information

URL: <https://bitbucket.org/mckinsel/shortfuse>

Proper Citation: ShortFuse (RRID:SCR_001107)

Description: THIS RESOURCE IS NO LONGER IN SERVICE. Documented on September 23,2022. A software package with tools for identifying fusion transcripts from RNA-Seq data. It is written in C++, and has dependencies on packages from Python 2.

Resource Type: data analysis software, software application, software resource, sequence analysis software, data processing software

Defining Citation: [PMID:21330288](#)

Keywords: fusion transcripts, rna, sequence data, python 2, c++, sequence analysis software, bio.tools

Availability: THIS RESOURCE IS NO LONGER IN SERVICE

Resource Name: ShortFuse

Resource ID: SCR_001107

Alternate IDs: biotools:shortfuse, OMICS_01355

Alternate URLs: <https://bio.tools/shortfuse>

Record Creation Time: 20220129T080205+0000

Record Last Update: 20260801T190139+0000

Ratings and Alerts

No rating or validation information has been found for ShortFuse.

No alerts have been found for ShortFuse.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 1 mentions in open access literature.

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

Latysheva NS, et al. (2016) Discovering and understanding oncogenic gene fusions through data intensive computational approaches. Nucleic acids research, 44(10), 4487.