## **Resource Summary Report**

Generated by ASWG on May 12, 2025

# **UnSplicer**

RRID:SCR\_000226

Type: Tool

### **Proper Citation**

UnSplicer (RRID:SCR\_000226)

#### **Resource Information**

URL: http://exon.gatech.edu/paul/unsplicer/index.htm

**Proper Citation:** UnSplicer (RRID:SCR\_000226)

**Description:** An RNA-seq alignment program that provides alignment of short reads to a reference genome. The program requires two inputs that are provided by the output of GeneMark-ES: HMM model parameters and ab initio gene predictions. UnSplicer is a sister pipeline to TrueSight.

Resource Type: software resource

**Defining Citation:** PMID:24259430

**Keywords:** RNA, sequencing, alignment, short reads, genome, genemark-es, gene

prediction

**Funding:** 

Availability: Open Source

Resource Name: UnSplicer

Resource ID: SCR\_000226

Alternate IDs: OMICS\_01806

**Record Creation Time:** 20220129T080200+0000

Record Last Update: 20250420T013935+0000

## **Ratings and Alerts**

No rating or validation information has been found for UnSplicer.

No alerts have been found for UnSplicer.

### **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 ASWG.

Burns PD, et al. (2014) UnSplicer: mapping spliced RNA-Seq reads in compact genomes and filtering noisy splicing. Nucleic acids research, 42(4), e25.