DELLY
RRID:SCR_004603
Type: Tool

Proper Citation

DELLY (RRID:SCR_004603)

Resource Information

URL: https://tobiasrausch.com/delly/

Description: Integrated structural variant prediction software that can detect deletions, tandem duplications, inversions and translocations at single-nucleotide resolution in short-read massively parallel sequencing data. It uses paired-ends and split-reads to sensitively and accurately delineate genomic rearrangements throughout genome.

Resource Name: DELLY

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Resource Type: Resource, software resource

Keywords: structural variant, genomic rearrangement, deletion, tandem duplication, inversion, translocation

Resource ID: SCR_004603

Parent Organization: European Molecular Biology Laboratory

References: PMID:22962449

Website Status: Last checked down

Alternate IDs: OMICS_00313

Alternate URLs: https://github.com/dellytools/delly/

Abbreviations: DELLY
Ratings and Alerts

No rating or validation information has been found for DELLY.

No alerts have been found for DELLY.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We found 109 mentions in open access literature.

Listed below are recent publications. The full list is available at scicrunch.


