## Running workflow "hisat2-alignment-paired-dataset-collections"



# Input data format **FASTQ** Single end or paired reads? Collection of paired reads Paired reads Output dataset 'output' from step 1 Paired-end options Use default values Write unaligned reads (in fastq format) to separate file(s) Not available. Write aligned reads (in fastq format) to separate file(s) Not available. Source for the reference genome to align against Use a genome from history Select the reference genome Output dataset 'output' from step 2 Primary alignments Not available. Maximum number of seeds that will be extended Not available. Report secondary alignments Not available. Alignment options Specify alignment parameters **Function type** Constant Constant term (y)

Not available.

#### Coefficient (z)

Not available.

#### Ignore quality values

Not available.

Not available.
Skip reverse strand of reference Not available.
Input options Specify input parameters
Skip the first N reads or pairs in the input Not available.
<b>Stop after aligning N reads</b> Not available.
<b>Trim 5' end</b> Not available.
<b>Trim 3' end</b> Not available.
Scoring options Specify scoring parameters
Function type Constant
Constant term (y) Not available.
Coefficient (z) Not available.
Set match bonus 2
Maximum mismatch penalty 6
Minimum mismatch penalty 2
Disallow soft-clipping
Not available.
Not available.  Ambiguous read penalty  1
Ambiguous read penalty
Ambiguous read penalty  1  Maximum soft-clipping penalty
Ambiguous read penalty  1  Maximum soft-clipping penalty  2  Minimum soft-clipping penalty
Ambiguous read penalty  1  Maximum soft-clipping penalty  2  Minimum soft-clipping penalty  1  Read gap open penalty
Ambiguous read penalty  1  Maximum soft-clipping penalty  2  Minimum soft-clipping penalty  1  Read gap open penalty  5  Read gap extend penalty
Ambiguous read penalty  Maximum soft-clipping penalty  Minimum soft-clipping penalty  Read gap open penalty  Read gap extend penalty  Reference gap open penalty
Ambiguous read penalty  Maximum soft-clipping penalty  Minimum soft-clipping penalty  Read gap open penalty  Read gap extend penalty  Reference gap open penalty  Reference gap open penalty

Skip forward strand of reference

### Penalty for canonical splice sites Not available. Penalty for non-canonical splice sites 3 Penalty for long introns with canonical splice sites Constant Constant term (y) Not available. Coefficient (z) Not available. Penalty for long introns with noncanonical splice sites Natural logarithm [f(x) = y + z \* log(x)]Constant term (y) -8 Coefficient (z) 1 Minimum intron length 20 Maximum intron length 500000 Specify strand-specific information FR Unstranded Disable spliced alignment True Minimum fragment length for valid paired-end alignments Not available. Maximum fragment length for valid paired-end alignments GTF file with known splice sites Selection is Optional \$ Transcriptome assembly reporting Use default reporting. Paired alignment parameters Specify paired alignment parameters Minimum fragment length Not available. Maximum fragment length Disable finding alignments for individual mates Not available. Disable looking for discordant alignments Not available. Mates not dovetail Not available. Mates cannot contain others

Not available.

### Mates cannot overlap

Not available.

Send results to a new history

### tools used in this workflow:

"tool\_id": null,
"tool\_id": null,

"tool\_id": "toolshed.g2.bx.psu.edu/repos/iuc/hisat2/hisat2/2.0.5.1",