

STAR manual 2.5.4b

Alexander Dobin
dobin@cshl.edu

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1 Getting started.

1.1 Installation.

STAR source code and binaries can be downloaded from GitHub: named releases from <https://github.com/alexdobin/STAR/releases>, or the master branch from <https://github.com/alexdobin/STAR>. The pre-compiled STAR executables are located `bin/` subdirectory. The `static` executables are the easisest to use, as they are statically compiled and are not dependents on external libraries.

To compile STAR from sources run `make` in the source directory for a Linux-like environment, or run `make STARforMac` for Mac OS X. This will produce the executable 'STAR' inside the source directory.

1.1.1 Installation - in depth and troubleshooting.

STAR is compiled with gcc c++ compiler and depends only on standard gcc libraries. Some generic instructions on installing correct gcc environments are given below.

Ubuntu.

```
$ sudo apt-get update
$ sudo apt-get install g++
$ sudo apt-get install make
```

Red Hat, CentOS, Fedora.

```
$ sudo yum update
$ sudo yum install make
$ sudo yum install gcc-c++
$ sudo yum install glibc-static
```

SUSE.

```
$ sudo zypper update
$ sudo zypper in gcc gcc-c++
```

Mac OS X.

Current versions of Mac OS X Xcode are shipped with Clang replacing the standard gcc compiler. Presently, standard Clang does not support OpenMP which creates problems for STAR compilation. One option to avoid this problem is to install gcc (preferably using `homebrew` package manager). Another option is to add OpenMP functionality to Clang.

1.2 Basic workflow.

Basic STAR workflow consists of 2 steps:

1. Generating genome indexes files (see [Section 2. Generating genome indexes](#).
In this step user supplied the reference genome sequences (FASTA files) and annotations (GTF file), from which STAR generate genome indexes that are utilized in the

2nd (mapping) step. The genome indexes are saved to disk and need only be generated **once** for each genome/annotation combination. A limited collection of STAR genomes is available from <http://labshare.cshl.edu/shares/gingeraslab/www-data/dobin/STAR/STARgenomes/>, however, it is strongly recommended that users generate their own genome indexes with most up-to-date assemblies and annotations.

2. Mapping reads to the genome (see [Section 3. Running mapping jobs](#)).

In this step user supplies the genome files generated in the 1st step, as well as the RNA-seq reads (sequences) in the form of FASTA or FASTQ files. STAR maps the reads to the genome, and writes several output files, such as alignments (SAM/BAM), mapping summary statistics, splice junctions, unmapped reads, signal (wiggle) tracks etc. Output files are described in [Section 4. Output files](#). Mapping is controlled by a variety of input parameters (options) that are described in brief in [Section 3. Running mapping jobs](#), and in more detail in [Section 9. Description of all options](#).

STAR command line has the following format:

```
STAR --option1-name option1-value(s) --option2-name option2-value(s) ...
```

If an option can accept multiple values, they are separated by spaces, and in a few cases - by commas.

2 Generating genome indexes.

2.1 Basic options.

The basic options to generate genome indices are as follows:

```
--runThreadN NumberOfThreads
--runMode genomeGenerate
--genomeDir /path/to/genomeDir
--genomeFastaFiles /path/to/genome/fasta1 /path/to/genome/fasta2 ...
--sjdbGTFfile /path/to/annotations.gtf
--sjdbOverhang ReadLength-1
```

--runThreadN option defines the number of threads to be used for genome generation, it has to be set to the number of available cores on the server node.

--runMode genomeGenerate option directs STAR to run genome indices generation job.

--genomeDir specifies path to the directory (henceforth called "genome directory" where the genome indices are stored. This directory has to be created (with `mkdir`) before STAR run and needs to have writing permissions. The file system needs to have at least 100GB of disk space available for a typical mammalian genome. It is recommended to remove all files from the genome directory before running the genome generation step. This directory path will have to be supplied at the mapping step to identify the reference genome.

--genomeFastaFiles specifies one or more FASTA files with the genome reference sequences. Multiple reference sequences (henceforth called chromosomes) are allowed for each fasta file. You can rename the chromosomes names in the `chrName.txt` keeping the order of the chromosomes in the file: the names from this file will be used in all output alignment files (such as `.sam`). The tabs are not allowed in chromosomes names, and spaces are not recommended.

`--sjdbGTFfile` specifies the path to the file with annotated transcripts in the standard GTF format. STAR will extract splice junctions from this file and use them to greatly improve accuracy of the mapping. While this is optional, and STAR can be run without annotations, using annotations is **highly recommended** whenever they are available. Starting from 2.4.1a, the annotations can also be included on the fly at the mapping step.

`--sjdbOverhang` specifies the length of the genomic sequence around the annotated junction to be used in constructing the splice junctions database. Ideally, this length should be equal to the $ReadLength-1$, where *ReadLength* is the length of the reads. For instance, for Illumina 2x100b paired-end reads, the ideal value is $100-1=99$. In case of reads of varying length, the ideal value is $max(ReadLength)-1$. **In most cases, the default value of 100 will work as well as the ideal value.**

Genome files comprise binary genome sequence, suffix arrays, text chromosome names/lengths, splice junctions coordinates, and transcripts/genes information. Most of these files use internal STAR format and are not intended to be utilized by the end user. It is strongly **not recommended** to change any of these file with one exception: you can rename the chromosome names in the chrName.txt keeping the order of the chromosomes in the file: the names from this file will be used in all output files (e.g. SAM/BAM).

2.2 Advanced options.

2.2.1 Which chromosomes/scaffolds/patches to include?

It is strongly recommended to include major chromosomes (e.g., for human chr1-22,chrX,chrY,chrM,) as well as un-placed and un-localized scaffolds. Typically, un-placed/un-localized scaffolds add just a few MegaBases to the genome length, however, a substantial number of reads may map to ribosomal RNA (rRNA) repeats on these scaffolds. These reads would be reported as unmapped if the scaffolds are not included in the genome, or, even worse, may be aligned to wrong loci on the chromosomes. Generally, patches and alternative haplotypes should **not** be included in the genome.

Examples of acceptable genome sequence files:

- **ENSEMBL:** files marked with .dna.primary.assembly, such as: ftp://ftp.ensembl.org/pub/release-77/fastq/homo_sapiens/dna/Homo_sapiens.GRCh38.dna.primary_assembly.fa.gz
- **NCBI:** "no alternative - analysis set": ftp://ftp.ncbi.nlm.nih.gov/genbank/genomes/Eukaryotes/vertebrates_mammals/Homo_sapiens/GRCh38/seqs_for_alignment_pipelines/GCA_000001405.15_GRCh38_no_alt_analysis_set.fna.gz

2.2.2 Which annotations to use?

The use of the most comprehensive annotations for a given species is strongly recommended. Very importantly, chromosome names in the annotations GTF file have to match chromosome names in the FASTA genome sequence files. For example, one can use ENSEMBL FASTA files with ENSEMBL GTF files, and UCSC FASTA files with UCSC GTF files. However, since UCSC uses chr1, chr2, ... naming convention, and ENSEMBL uses 1, 2, ... naming, the ENSEMBL and UCSC FASTA

and GTF files cannot be mixed together, unless chromosomes are renamed to match between the FASTA and GTF files.

For mouse and human, the Gencode annotations are recommended: <http://www.gencodegenes.org/>.

2.2.3 Annotations in GFF format.

In addition to the aforementioned options, for GFF3 formatted annotations you need to use `--sjdbGTFtagExonParentTranscript Parent`. In general, for `--sjdbGTFfile` files STAR only processes lines which have `--sjdbGTFfeatureExon (=exon by default)` in the 3rd field (column). The exons are assigned to the transcripts using parent-child relationship defined by the `--sjdbGTFtagExonParentTranscript (=transcript_id by default)` GTF/GFF attribute.

2.2.4 Using a list of annotated junctions.

STAR can also utilize annotations formatted as a list of splice junctions coordinates in a text file: `--sjdbFileChrStartEnd /path/to/sjdbFile.txt`. This file should contain 4 columns separated by tabs:

```
Chr    \tab  Start    \tab  End    \tab  Strand=+/-/.
```

Here Start and End are first and last bases of the introns (1-based chromosome coordinates). This file can be used in addition to the `--sjdbGTFfile`, in which case STAR will extract junctions from both files.

Note, that the `--sjdbFileChrStartEnd` file can contain duplicate (identical) junctions, STAR will collapse (remove) duplicate junctions.

2.2.5 Very small genome.

For small genomes, the parameter `--genomeSAindexNbases` must be scaled down, with a typical value of $\min(14, \log_2(\text{GenomeLength})/2 - 1)$. For example, for 1 megaBase genome, this is equal to 9, for 100 kiloBase genome, this is equal to 7.

2.2.6 Genome with a large number of references.

If you are using a genome with a large ($>5,000$) number of references (chromosomes/scaffolds), you may need to reduce the `--genomeChrBinNbits` to reduce RAM consumption. The following scaling is recommended: `--genomeChrBinNbits = $\min(18, \log_2(\text{GenomeLength}/\text{NumberOfReferences}))$` . For example, for 3 gigaBase genome with 100,000 chromosomes/scaffolds, this is equal to 15.

3 Running mapping jobs.

3.1 Basic options.

The basic options to run a mapping job are as follows:

```
--runThreadN NumberOfThreads
--genomeDir /path/to/genomeDir
--readFilesIn /path/to/read1 [/path/to/read2]
```

`--genomeDir` specifies path to the genome directory where genome indices were generated (see [Section 2. Generating genome indexes](#)).

`--readFilesIn` name(s) (with path) of the files containing the sequences to be mapped (e.g. RNA-seq FASTQ files). If using Illumina paired-end reads, the *read1* and *read2* files have to be supplied. STAR can process both FASTA and FASTQ files. Multi-line (i.e. sequence split in multiple lines) FASTA (but not FASTQ) files are supported.

If the read files are compressed, use the `--readFilesCommand` *UncompressionCommand* option, where *UncompressionCommand* is the un-compression command that takes the file name as input parameter, and sends the uncompressed output to stdout. For example, for gzipped files (*.gz) use `--readFilesCommand` *zcat* OR `--readFilesCommand` *gunzip -c*. For bzip2-compressed files, use `--readFilesCommand` *bunzip2 -c*.

Multiple samples can be mapped in one job. For single-end reads use a comma separated list (no spaces around commas), e.g. `--readFilesIn` *sample1.fq,sample2.fq,sample3.fq*. For paired-end reads, use comma separated list for read1 /space/ comma separated list for read2, e.g.: `--readFilesIn` *sample1read1.fq,sample2read1.fq,sample3read1.fq sample1read2.fq,sample2read2.fq,sample3read2.fq*.

3.2 Advanced options.

There are many advanced options that control STAR mapping behavior. All options are briefly described in the [Section 9. Description of all options](#).

3.2.1 Using annotations at the mapping stage.

Since 2.4.1a, the annotations can be included on the fly at the mapping step, without including them at the genome generation step. You can specify `--sjdbGTFfile` */path/to/ann.gtf* and/or `--sjdbFileChrStartEnd` */path/to/sj.tab*, as well as `--sjdbOverhang`, and any other `--sjdb*` options. The genome indices can be generated with or without another set of annotations/junctions. In the latter case the new junctions will be added to the old ones. STAR will insert the junctions into genome indices on the fly before mapping, which takes 1-2 minutes. The on the fly genome indices can be saved (for reuse) with `--sjdbInsertSave` *All*, into *_STARgenome* directory inside the current run directory.

3.2.2 ENCODE options

An example of ENCODE standard options for long RNA-seq pipeline is given below:

`--outFilterType` *BySJout*
reduces the number of "spurious" junctions

`--outFilterMultimapNmax` *20*
max number of multiple alignments allowed for a read: if exceeded, the read is considered unmapped

`--alignSJoverhangMin` *8*
minimum overhang for unannotated junctions

`--alignSJDBoverhangMin 1`
 minimum overhang for annotated junctions

`--outFilterMismatchNmax 999`
 maximum number of mismatches per pair, large number switches off this filter

`--outFilterMismatchNoverReadLmax 0.04`
 max number of mismatches per pair relative to read length: for 2x100b, max number of mismatches is $0.04 \times 200 = 8$ for the paired read

`--alignIntronMin 20`
 minimum intron length

`--alignIntronMax 1000000`
 maximum intron length

`--alignMatesGapMax 1000000`
 maximum genomic distance between mates

3.3 Using shared memory for the genome indexes.

The `--genomeLoad` option controls how the genome is loaded into memory. By default, `--genomeLoad NoSharedMemory`, shared memory is not used.

With `--genomeLoad LoadAndKeep`, STAR loads the genome as a standard Linux shared memory piece. The genomes are identified by their unique directory paths. Before loading the genome, STAR checks if the genome has already been loaded into the shared memory. If the genome has not been loaded, STAR will load it and will keep it in memory even after STAR job finishes. The genome will be shared with all the other STAR jobs. You can remove the genome from the shared memory running STAR with `--genomeLoad Remove`. The shared memory piece will be physically removed only after all STAR jobs attached to it complete. With `--genomeLoad LoadAndRemove`, STAR will load genome in the shared memory, and mark it for removal, so that the genome will be removed from the shared memory once all STAR jobs using it exit. `--genomeLoad LoadAndExit`, STAR will load genome in the shared memory, and immediately exit, keeping the genome loaded in the shared memory for the future runs.

If you need to check or remove shared memory pieces manually, use the standard Linux command `ipcs` and `ipcrm`. If the genome residing in shared memory is not used for a long time it may get paged out of RAM which will slow down STAR runs considerably. It is strongly recommended to regularly re-load (i.e. remove and load again) the shared memory genomes.

Many standard Linux distributions do not allow large enough shared memory blocks. You can fix this issue if you have root privileges, or ask your system administrator to do it. To enable the shared memory modify or add the following lines to `/etc/sysctl.conf`:

```
kernel.shmmax = Nmax
kernel.shmall = Nall
```

Nmax, *Nall* numbers should be chosen as follows:

$Nmax > GenomeIndexSize = Genome + SA + SAindex$ (31000000000 for human genome)

$Nall > GenomeIndexSize / PageSize$

where PageSize is typically 4096 (this can be checked with `getconf PAGE_SIZE`). Then run:
`/sbin/sysctl -p`
This will increase the allowed shared memory blocks to 31GB, enough for human or mouse genome.

4 Output files.

STAR produces multiple output files. All files have standard name, however, you can change the file prefixes using `--outFileNamePrefix /path/to/output/dir/prefix`. By default, this parameter is `./`, i.e. all output files are written in the current directory.

4.1 Log files.

Log.out: main log file with a lot of detailed information about the run. This file is most useful for troubleshooting and debugging.

Log.progress.out: reports job progress statistics, such as the number of processed reads, % of mapped reads etc. It is updated in 1 minute intervals.

Log.final.out: summary mapping statistics after mapping job is complete, very useful for quality control. The statistics are calculated for each read (single- or paired-end) and then summed or averaged over all reads. Note that STAR counts a paired-end read as one read, (unlike the samtools flagstat/idxstats, which count each mate separately). Most of the information is collected about the UNIQUE mappers (unlike samtools flagstat/idxstats which does not separate unique or multi-mappers). Each splicing is counted in the numbers of splices, which would correspond to summing the counts in **SJ.out.tab**. The mismatch/indel error rates are calculated on a per base basis, i.e. as total number of mismatches/indels in all unique mappers divided by the total number of mapped bases.

4.2 SAM.

Aligned.out.sam - alignments in standard SAM format.

4.2.1 Multimappers.

The number of loci **Nmap** a read maps to is given by **NH:i:Nmap** field. Value of 1 corresponds to unique mappers, while values >1 corresponds to multi-mappers. **HI** attributes enumerates multiple alignments of a read starting with 1 (this can be changed with the `--outSAMattrIHstart` - setting it to 0 may be required for compatibility with downstream software such as Cufflinks or StringTie).

The mapping quality MAPQ (column 5) is 255 for uniquely mapping reads, and $\text{int}(-10 \cdot \log_{10}(1 - 1/\text{Nmap}))$ for multi-mapping reads. This scheme is same as the one used by TopHat and is compatible with Cufflinks. The default MAPQ=255 for the unique mappers maybe changed with `--outSAMmapqUnique` parameter (integer 0 to 255) to ensure compatibility with downstream tools such as GATK.

For multi-mappers, all alignments except one are marked with 0x100 (secondary alignment) in the FLAG (column 2 of the SAM). The unmarked alignment is selected from the best ones (i.e.

highest scoring). This default behavior can be changed with `--outSAMprimaryFlag AllBestScore` option, that will output all alignments with the best score as primary alignments (i.e. 0x100 bit in the FLAG unset).

By default, the order of the multi-mapping alignments for each read is not truly random. The `--outMultimapperOrder Random` option outputs multiple alignments for each read in random order, and also randomizes the choice of the primary alignment from the highest scoring alignments. Parameter `--runRNGseed` can be used to set the random generator seed. With this option, the ordering of multi-mapping alignments of each read, and the choice of the primary alignment will vary from run to run, unless only one thread is used and the seed is kept constant.

The `--outSAMmultNmax` parameter limits the number of output alignments (SAM lines) for multimappers. For instance, `--outSAMmultNmax 1` will output exactly one SAM line for each mapped read. Note that `NH:i:` tag in STAR will still report the actual number of loci that the reads map to, while the number of reported alignments for a read in the SAM file is `min(NH, --outSAMmultNmax)`.

4.2.2 SAM attributes.

The SAM attributes can be specified by the user using `--outSAMattributes A1 A2 A3 ...` option which accept a list of 2-character SAM attributes. The implemented attributes are: `NH HI NM MD AS nM jM jI XS`. By default, STAR outputs `NH HI AS nM` attributes.

`NH HI NM MD` have standard meaning as defined in the SAM format specifications.

`AS` is the local alignment score (paired for paired-end reads).

`nM` is the number of mismatches per (paired) alignment, not to be confused with `NM`, which is the number of mismatches in each mate.

`jM:B:c,M1,M2,...` intron motifs for all junctions (i.e. `N` in CIGAR): 0: non-canonical; 1: GT/AG, 2: CT/AC, 3: GC/AG, 4: CT/GC, 5: AT/AC, 6: GT/AT. If splice junctions database is used, and a junction is annotated, 20 is added to its motif value.

`jI:B:I,Start1,End1,Start2,End2,...` Start and End of introns for all junctions (1-based).

`jM jI` attributes require samtools 0.1.18 or later, and were reported to be incompatible with some downstream tools such as Cufflinks.

4.2.3 Compatibility with Cufflinks/Cuffdiff.

For unstranded RNA-seq data, Cufflinks/Cuffdiff require spliced alignments with `XS` strand attribute, which STAR will generate with `--outSAMstrandField intronMotif` option. As required, the `XS` strand attribute will be generated for all alignments that contain splice junctions. The spliced alignments that have undefined strand (i.e. containing only non-canonical unannotated junctions) will be suppressed.

If you have stranded RNA-seq data, you do not need to use any specific STAR options. Instead, you need to run Cufflinks with the library option `--library-type` options. For example, `cufflinks ... --library-type fr-firststrand` should be used for the standard dUTP protocol, including

Illumina’s stranded Tru-Seq. This option has to be used only for Cufflinks runs and not for STAR runs.

In addition, it is recommended to remove the non-canonical junctions for Cufflinks runs using `--outFilterIntronMotifs RemoveNoncanonical`.

4.3 Unsorted and sorted-by-coordinate BAM.

STAR can output alignments directly in binary BAM format, thus saving time on converting SAM files to BAM. It can also sort BAM files by coordinates, which is required by many downstream applications.

`--outSAMtype` BAM Unsorted

output unsorted `Aligned.out.bam` file. The paired ends of an alignment are always adjacent, and multiple alignments of a read are adjacent as well. This ”unsorted” file can be directly used with downstream software such as HTseq, without the need of name sorting. The order of the reads will match that of the input FASTQ(A) files only if one thread is used

`--runThread 1`, and `--outFilterType --BySJout` is **not** used.

`--outSAMtype` BAM SortedByCoordinate

output sorted by coordinate `Aligned.sortedByCoord.out.bam` file, similar to `samtools sort` command.

`--outSAMtype` BAM Unsorted SortedByCoordinate

output both unsorted and sorted files.

4.4 Splice junctions.

`SJ.out.tab` contains high confidence collapsed splice junctions in tab-delimited format. Note that STAR defines the junction start/end as intronic bases, while many other software define them as exonic bases. The columns have the following meaning:

column 1: chromosome

column 2: first base of the intron (1-based)

column 3: last base of the intron (1-based)

column 4: strand (0: undefined, 1: +, 2: -)

column 5: intron motif: 0: non-canonical; 1: GT/AG, 2: CT/AC, 3: GC/AG, 4: CT/GC, 5: AT/AC, 6: GT/AT

column 6: 0: unannotated, 1: annotated (only if splice junctions database is used)

column 7: number of uniquely mapping reads crossing the junction

column 8: number of multi-mapping reads crossing the junction

column 9: maximum spliced alignment overhang

The filtering for this output file is controlled by the `--outSJfilter*` parameters, as described in Section 9.15. Output Filtering: Splice Junctions.

5 Chimeric and circular alignments.

To switch on detection of chimeric (fusion) alignments (in addition to normal mapping), `--chimSegmentMin` should be set to a positive value. Each chimeric alignment consists of two "segments". Each segment is non-chimeric on its own, but the segments are chimeric to each other (i.e. the segments belong to different chromosomes, or different strands, or are far from each other). Both segments may contain splice junctions, and one of the segments may contain portions of both mates. `--chimSegmentMin` parameter controls the minimum mapped length of the two segments that is allowed. For example, if you have 2x75 reads and used `--chimSegmentMin 20`, a chimeric alignment with 130b on one chromosome and 20b on the other will be output, while 135 + 15 won't be.

5.1 STAR-Fusion.

STAR-Fusion is a software package for detecting fusion transcript from STAR chimeric output. It is developed and maintained by Brian Haas (@Broad Institute), whose effort was inspired by earlier work done by Nicolas Stransky in the landmark publication "The landscape of kinase fusions in cancer" by Stransky et al., Nat Commun 2014, in addition to very nice work done by Daniel Nicorici with his FusionCatcher software. Please visit its GitHub page for instructions and documentation: <https://github.com/STAR-Fusion/STAR-Fusion>.

5.2 Chimeric alignments in the main BAM files.

Chimeric alignments can be included together with normal alignments in the main (sorted or unsorted) BAM file(s) using `--chimOutType WithinBAM`. In these files, formatting of chimeric alignments follows the latest SAM/BAM specifications.

5.3 Chimeric alignments in Chimeric.out.sam .

When chimeric detection is switched on, STAR will output normal alignments into `Aligned.*.sam/bam`, and will output chimeric alignments into a separate file `Chimeric.out.sam`. Some reads may be output to both normal SAM/BAM files, and `Chimeric.out.sam` for the following reason. STAR will output a non-chimeric alignment into `Aligned.out.sam` with soft-clipping a portion of the read. If this portion is long enough, and it maps well and uniquely somewhere else in the genome, there will also be a chimeric alignment output into `Chimeric.out.sam`. For instance, if you have a paired-end read where the second mate can be split chimerically into 70 and 30 bases. The 100b of the first mate + 70b of the 2nd mate map non-chimerically, and the mapping length/score are big enough, so they will be output into `Aligned.out.sam` file. At the same time, the chimeric segments 100-mate1 + 70-mate2 and 30-mate2 will be output into `Chimeric.out.sam`.

5.4 Chimeric alignments in Chimeric.out.junction

In addition to `Chimeric.out.sam`, STAR will generate `Chimeric.out.junction` file which maybe more convenient for downstream analysis. The format of this file is as follows. Every line contains one chimerically aligned read, e.g.:

```
chr22    23632601      +      chr9    133729450      +      1        0        0
SINATRA-0006:3:3:6387:5665#0    23632554    47M29S  133729451      47S29M40p76M
```

The first 9 columns give information about the chimeric junction:

column 1: chromosome of the donor

column 2: first base of the intron of the donor (1-based)

column 3: strand of the donor

column 4: chromosome of the acceptor

column 5: first base of the intron of the acceptor (1-based)

column 6: strand of the acceptor

column 7: junction type: -1=encompassing junction (between the mates), 1=GT/AG, 2=CT/AC

column 8: repeat length to the left of the junction

column 9: repeat length to the right of the junction

Columns 10-14 describe the alignments of the two chimeric segments, it is SAM like. Alignments are given with respect to the (+) strand

column 10: read name

column 11: first base of the first segment (on the + strand)

column 12: CIGAR of the first segment

column 13: first base of the second segment

column 14: CIGAR of the second segment

Unlike standard SAM, both mates are recorded in one line here. The gap of length *L* between the mates is marked by the *p* in the CIGAR string. If the mates overlap, *L*<0.

For strand definitions, when aligning paired end reads, the sequence of the second mate is reverse complemented.

For encompassing junctions, i.e. junction type: -1=junction is between the mates, columns 2 and 5 represent the bounds on the chimeric junction loci. For the 1st mate, it will be the genomic base following the last 3' mapped base. For the 2nd mate (which is reverse complemented to have the same orientation as 1st mate), it will be the genomic base preceding the 5' mapped base. For example, if there is a chimeric junction that connects chr1/+strand/base1000 to chr2/+strand/base2000,

and read 1 maps to chr1/+strand/bases800-900, and read 2 (after reverse complementing) maps to chr2/+strand/bases2100-2200, then columns 2 and 5 will have 901 and 2099.

To filter chimeric junctions and find the number of reads supporting each junction you could use, for example:

```
cat Chimeric.out.junction |
awk '$1!="chrM" && $4!="chrM" && $7>0 && $8+$9<=5 {print $1,$2,$3,$4,$5,$6,$7,$8,$9}' |
sort | uniq -c | sort -k1,1rn
```

This will keep only the canonical junctions with the repeat length less than 5 and will remove chimeras with mitochondrion genome.

When I do it for one of our K562 runs, I get:

181	chr1	144676873	-	chr1	147917466	+	1	0	1
29	chr5	69515744	-	chr5	34182973	-	1	3	1
28	chr1	143910077	-	chr1	149459550	-	1	1	0
27	chr22	23632601	+	chr9	133729450	+	1	0	0
20	chr12	90313405	-	chr21	40684813	-	1	2	0
20	chr22	23632601	+	chr9	133655755	+	1	0	1
20	chr9	123636256	-	chr9	123578959	+	1	1	4
15	chr16	85589970	+	chr6	16762582	+	1	3	2
15	chr3	197348574	-	chr3	195392936	+	1	1	0
14	chr18	39584506	+	chr18	39560613	-	1	2	0

Note that line 4 and 6 here are BCR/ABL fusions. You would need to filter these junctions further to see which of them connect known but not homologous genes.

6 Output in transcript coordinates.

With `--quantMode TranscriptomeSAM` option STAR will output alignments translated into transcript coordinates in the `Aligned.toTranscriptome.out.bam` file (in addition to alignments in genomic coordinates in `Aligned.*.sam/bam` files). These transcriptomic alignments can be used with various transcript quantification software that require reads to be mapped to transcriptome, such as RSEM or eXpress. For example, RSEM command line would look as follows:

```
rsem-calculate-expression ... --bam Aligned.toTranscriptome.out.bam
/path/to/RSEM/reference RSEM
```

. Note, that STAR first aligns reads to entire genome, and only then searches for concordance between alignments and transcripts. I believe this approach might offer certain advantages compared to the alignment to transcriptome only, by not forcing the alignments to annotated transcripts.

By default, the output satisfies RSEM requirements: soft-clipping or indels are not allowed. Use `--quantTranscriptomeBan Singleend` to allow insertions, deletions and soft-clips in the transcriptomic alignments, which can be used by some expression quantification software (e.g. eXpress).

7 Counting number of reads per gene.

With `--quantMode GeneCounts` option STAR will count number reads per gene while mapping. A read is counted if it overlaps (1nt or more) one and only one gene. Both ends of the paired-end read are checked for overlaps. The counts coincide with those produced by htseq-count with default parameters. This option requires annotations (GTF or GFF with `-sjdbGTFfile` option) used at the genome generation step, or at the mapping step. STAR outputs read counts per gene into `ReadsPerGene.out.tab` file with 4 columns which correspond to different strandedness options:

column 1: gene ID

column 2: counts for unstranded RNA-seq

column 3: counts for the 1st read strand aligned with RNA (htseq-count option `-s yes`)

column 4: counts for the 2nd read strand aligned with RNA (htseq-count option `-s reverse`)

Select the output according to the strandedness of your data. Note, that if you have stranded data and choose one of the columns 3 or 4, the other column (4 or 3) will give you the count of antisense reads. With `--quantMode TranscriptomeSAM GeneCounts`, and get both the `Aligned.toTranscriptome.out.bam` and `ReadsPerGene.out.tab` outputs.

8 2-pass mapping.

For the most sensitive novel junction discovery, I would recommend running STAR in the 2-pass mode. It does not increase the number of detected novel junctions, but allows to detect more splices reads mapping to novel junctions. The basic idea is to run 1st pass of STAR mapping with the usual parameters, then collect the junctions detected in the first pass, and use them as "annotated" junctions for the 2nd pass mapping.

8.1 Multi-sample 2-pass mapping.

For a study with multiple samples, it is recommended to collect 1st pass junctions from all samples.

1. Run 1st mapping pass for all samples with "usual" parameters. Using annotations is recommended either at the genome generation step, or mapping step.
2. Run 2nd mapping pass for all samples, listing `SJ.out.tab` files from all samples in `--sjdbFileChrStartEnd /path/to/sj1.tab /path/to/sj2.tab`

8.2 Per-sample 2-pass mapping.

Annotated junctions will be included in both the 1st and 2nd passes. To run STAR 2-pass mapping for each sample separately, use `--twopassMode Basic` option. STAR will perform the 1st pass mapping, then it will automatically extract junctions, insert them into the genome index, and, finally, re-map all reads in the 2nd mapping pass. This option can be used with annotations, which can be included either at the run-time (see #1), or at the genome generation step.

`--twopass1readsN` defines the number of reads to be mapped in the 1st pass. The default and most sensitive approach is to set it to -1 (or make it bigger than the number of reads in the sample) - in which case all reads in the input read file(s) are used in the 1st pass. While it can reduce mapping time by $\sim 40\%$, it is not recommended to use a small portion of the reads in the 1st step, since it will significantly reduce sensitivity for the low expressed novel junctions. The idea to use a portion of the reads in the 1st pass was inspired by Kim, Langmead and Salzberg in Nature Methods 12, 357360 (2015).

8.3 2-pass mapping with re-generated genome.

This is the original 2-pass method which involves genome re-generation step in-between 1st and 2nd passes. Since 2.4.1a, it is recommended to use the on the fly 2-pass options as described above.

1. Run 1st pass STAR for all samples with "usual" parameters. Genome indices generated with annotations are recommended.
2. Collect all junctions detected in the 1st pass by merging `SJ.out.tab` files from all runs. Filter the junctions by removing likelie false positives, e.g. junctions in the mitochondrion genome, or non-canonical junctions supported by a few reads. If you are using annotations, only novel junctions need to be considered here, since annotated junctions will be re-used in the 2nd pass anyway.
3. Use the filtered list of junctions from the 1st pass with `--sjdbFileChrStartEnd` option, together with annotations (via `--sjdbGTFfile` option) to generate the new genome indices for the 2nd pass mapping. This needs to be done only once for all samples.
4. Run the 2nd pass mapping for all samples with the new genome index.

9 Description of all options.

For each STAR version, the most up-to-date information about all STAR parameters can be found in the `parametersDefault` file in the STAR source directory. The parameters in the `parametersDefault`, as well as in the descriptions below, are grouped by function:

Special attention has to be paid to parameters that start with `--out*`, as they control the STAR output.

In particular, `--outFilter*` parameters control the filtering of output alignments which[] you might want to tweak to fit your needs.

Output of chimeric alignments is controlled by `--chim*` parameters.

Genome generation is controlled by `--genome*` parameters.

Annotations (splice junction database) are controlled by `--sjdb*` options at the genome generation step.

Tweaking `--score*`, `--align*`, `--seed*`, `--win*` parameters, which requires understanding of the STAR alignment algorithm, is recommended only for advanced users.

Below, allowed parameter values are typed in magenta, and default values - in blue.

9.1 Parameter Files

`--parametersFiles`

default: -

string: name of a user-defined parameters file, "-": none. Can only be defined on the command line.

9.2 System

`--sysShell`

default: -

string: path to the shell binary, preferably bash, e.g. /bin/bash.

-

the default shell is executed, typically /bin/sh. This was reported to fail on some Ubuntu systems - then you need to specify path to bash.

9.3 Run Parameters

`--runMode`

default: `alignReads`

string: type of the run:

`alignReads`

map reads

`genomeGenerate`

generate genome files

`inputAlignmentsFromBAM`

input alignments from BAM. Presently only works with `-outWigType` and `-bamRemoveDuplicates`.

`liftOver`

lift-over of GTF files (`-sjdbGTFfile`) between genome assemblies using chain file(s) from `-genomeChainFiles`.

`--runThreadN`

default: 1

int: number of threads to run STAR

`--runDirPerm`

default: `User_RWX`

string: permissions for the directories created at the run-time.

`User_RWX`

user-read/write/execute

`All_RWX`

all-read/write/execute (same as `chmod 777`)

`--runRNGseed`

default: `777`

int: random number generator seed.

9.4 Genome Parameters

`--genomeDir`

default: `./GenomeDir/`

string: path to the directory where genome files are stored (for `-runMode alignReads`) or will be generated (for `-runMode generateGenome`)

`--genomeLoad`

default: `NoSharedMemory`

string: mode of shared memory usage for the genome files. Only used with `-runMode alignReads`.

`LoadAndKeep`

load genome into shared and keep it in memory after run

`LoadAndRemove`

load genome into shared but remove it after run

`LoadAndExit`

load genome into shared memory and exit, keeping the genome in memory for future runs

`Remove`

do not map anything, just remove loaded genome from memory

`NoSharedMemory`

do not use shared memory, each job will have its own private copy of the genome

`--genomeFastaFiles`

default: `-`

string(s): path(s) to the fasta files with the genome sequences, separated by spaces. These files should be plain text FASTA files, they **cannot** be zipped. Required for the genome generation (`-runMode genomeGenerate`). Can also be used in the mapping (`-runMode alignReads`) to add extra (new) sequences to the genome (e.g. spike-ins).

--genomeChainFiles

default: -

string: chain files for genomic liftOver. Only used with `-runMode liftOver`.

--genomeFileSizes

default: 0

uint(s)>0: genome files exact sizes in bytes. Typically, this should not be defined by the user.

--genomeConsensusFile

default: -

string: VCF file with consensus SNPs (i.e. alternative allele is the major (AF>0.5) allele)

9.5 Genome Indexing Parameters - only used with `-runMode genomeGenerate`

--genomeChrBinNbits

default: 18

int: $=\log_2(\text{chrBin})$, where chrBin is the size of the bins for genome storage: each chromosome will occupy an integer number of bins. For a genome with large number of contigs, it is recommended to scale this parameter as $\min(18, \log_2[\max(\text{GenomeLength}/\text{NumberOfReferences}, \text{ReadLength})])$.

--genomeSAindexNbases

default: 14

int: length (bases) of the SA pre-indexing string. Typically between 10 and 15. Longer strings will use much more memory, but allow faster searches. For small genomes, the parameter `-genomeSAindexNbases` must be scaled down to $\min(14, \log_2(\text{GenomeLength})/2 - 1)$.

--genomeSAsparseD

default: 1

int>0: suffix array sparsity, i.e. distance between indices: use bigger numbers to decrease needed RAM at the cost of mapping speed reduction

--genomeSuffixLengthMax

default: -1

int: maximum length of the suffixes, has to be longer than read length. -1 = infinite.

9.6 Splice Junctions Database

`--sjdbFileChrStartEnd`

default: -

string(s): path to the files with genomic coordinates (chr <tab> start <tab> end <tab> strand) for the splice junction introns. Multiple files can be supplied and will be concatenated.

`--sjdbGTFfile`

default: -

string: path to the GTF file with annotations

`--sjdbGTFchrPrefix`

default: -

string: prefix for chromosome names in a GTF file (e.g. 'chr' for using ENSEMBL annotations with UCSC genomes)

`--sjdbGTFfeatureExon`

default: exon

string: feature type in GTF file to be used as exons for building transcripts

`--sjdbGTFtagExonParentTranscript`

default: transcript_id

string: tag name to be used as exons' transcript-parents (default "transcript_id" works for GTF files)

`--sjdbGTFtagExonParentGene`

default: gene_id

string: tag name to be used as exons' gene-parents (default "gene_id" works for GTF files)

`--sjdbOverhang`

default: 100

int>0: length of the donor/acceptor sequence on each side of the junctions, ideally = (mate.length - 1)

`--sjdbScore`

default: 2

int: extra alignment score for alignmets that cross database junctions

`--sjdbInsertSave`

default: **Basic**

string: which files to save when sjdb junctions are inserted on the fly at the mapping step

Basic

only small junction / transcript files

All

all files including big Genome, SA and SAindex - this will create a complete genome directory

9.7 Input Files

`--inputBAMfile`

default: **-**

string: path to BAM input file, to be used with `-runMode inputAlignmentsFromBAM`

9.8 Read Parameters

`--readFilesType`

default: **Fastx**

string: format of input read files

Fastx

FASTA or FASTQ

SAM SE

SAM or BAM single-end reads; for BAM use `-readFilesCommand samtools view`

SAM PE

SAM or BAM paired-end reads; for BAM use `-readFilesCommand samtools view`

`--readFilesIn`

default: **Read1 Read2**

string(s): paths to files that contain input read1 (and, if needed, read2)

`--readFilesPrefix`

default: **-**

string: prefix for the read files names, i.e. it will be added in front of the strings in `--readFilesIn`

-: no prefix

`--readFilesCommand`

default: -

string(s): command line to execute for each of the input file. This command should generate FASTA or FASTQ text and send it to stdout

For example: `zcat -` to uncompress .gz files, `bzcat -` to uncompress .bz2 files, etc.

`--readMapNumber`

default: -1

int: number of reads to map from the beginning of the file

-1: map all reads

`--readMatesLengthsIn`

default: `NotEqual`

string: `Equal/NotEqual` - lengths of names, sequences, qualities for both mates are the same / not the same. `NotEqual` is safe in all situations.

`--readNameSeparator`

default: /

string(s): character(s) separating the part of the read names that will be trimmed in output (read name after space is always trimmed)

`--clip3pNbases`

default: 0

int(s): number(s) of bases to clip from 3p of each mate. If one value is given, it will be assumed the same for both mates.

`--clip5pNbases`

default: 0

int(s): number(s) of bases to clip from 5p of each mate. If one value is given, it will be assumed the same for both mates.

`--clip3pAdapterSeq`

default: -

string(s): adapter sequences to clip from 3p of each mate. If one value is given, it will be assumed the same for both mates.

--clip3pAdapterMMp

default: 0.1

double(s): max proportion of mismatches for 3p adapter clipping for each mate. If one value is given, it will be assumed the same for both mates.

--clip3pAfterAdapterNbases

default: 0

int(s): number of bases to clip from 3p of each mate after the adapter clipping. If one value is given, it will be assumed the same for both mates.

9.9 Limits

--limitGenomeGenerateRAM

default: 31000000000

int>0: maximum available RAM (bytes) for genome generation

--limitIObufferSize

default: 150000000

int>0: max available buffers size (bytes) for input/output, per thread

--limitOutSAMoneReadBytes

default: 100000

int>0: max size of the SAM record (bytes) for one read. Recommended value: $>(2*(LengthMate1+LengthMate2+100)*outFilterMultimapNmax)$

--limitOutSJoneRead

default: 1000

int>0: max number of junctions for one read (including all multi-mappers)

--limitOutSJcollapsed

default: 1000000

int>0: max number of collapsed junctions

--limitBAMsortRAM

default: 0

int>=0: maximum available RAM (bytes) for sorting BAM. If =0, it will be set to the genome index size. 0 value can only be used with `-genomeLoad NoSharedMemory` option.

--limitSjdbInsertNsj

default: 1000000

int>=0: maximum number of junction to be inserted to the genome on the fly at the mapping stage, including those from annotations and those detected in the 1st step of the 2-pass run

9.10 Output: general

`--outFileNamePrefix`

default: `./`

string: output files name prefix (including full or relative path). Can only be defined on the command line.

`--outTmpDir`

default: `-`

string: path to a directory that will be used as temporary by STAR. All contents of this directory will be removed!

- the temp directory will default to `outFileNamePrefix_STARtmp`

`--outTmpKeep`

default: `None`

string: whether to keep the temporary files after STAR runs is finished

`None`

remove all temporary files

All .. keep all files

`--outStd`

default: `Log`

string: which output will be directed to stdout (standard out)

`Log`

log messages

`SAM`

alignments in SAM format (which normally are output to `Aligned.out.sam` file), normal standard output will go into `Log.std.out`

`BAM.Unsorted`

alignments in BAM format, unsorted. Requires `-outSAMtype BAM Unsorted`

`BAM.SortedByCoordinate`

alignments in BAM format, unsorted. Requires `-outSAMtype BAM SortedByCoordinate`

`BAM.Quant`

alignments to transcriptome in BAM format, unsorted. Requires `-quantMode TranscriptomeSAM`

`--outReadsUnmapped`

default: **None**

string: output of unmapped and partially mapped (i.e. mapped only one mate of a paired end read) reads in separate file(s).

None

no output

Fastx

output in separate fasta/fastq files, Unmapped.out.mate1/2

--outQScnversionAdd

default: **0**

int: add this number to the quality score (e.g. to convert from Illumina to Sanger, use -31)

--outMultimapperOrder

default: **Old_2.4**

string: order of multimapping alignments in the output files

Old_2.4

quasi-random order used before 2.5.0

Random

random order of alignments for each multi-mapper. Read mates (pairs) are always adjacent, all alignment for each read stay together. This option will become default in the future releases.

9.11 Output: SAM and BAM

--outSAMtype

default: **SAM**

strings: type of SAM/BAM output

1st word:

BAM

output BAM without sorting

SAM

output SAM without sorting

None

no SAM/BAM output

2nd, 3rd:

Unsorted

standard unsorted

SortedByCoordinate

sorted by coordinate. This option will allocate extra memory for sorting which can be specified by `-limitBAMsortRAM`.

`--outSAMmode`

default: Full

string: mode of SAM output

None

no SAM output

Full

full SAM output

NoQS

full SAM but without quality scores

`--outSAMstrandField`

default: None

string: Cufflinks-like strand field flag

None

not used

intronMotif

strand derived from the intron motif. Reads with inconsistent and/or non-canonical introns are filtered out.

`--outSAMattributes`

default: Standard

string: a string of desired SAM attributes, in the order desired for the output SAM

NH HI AS nM NM MD jM jI XS ch

any combination in any order

Standard

NH HI AS nM

All

NH HI AS nM NM MD jM jI ch

None

no attributes

`--outSAMattrIHstart`

default: 1

int>=0: start value for the IH attribute. 0 may be required by some downstream software, such as Cufflinks or StringTie.

`--outSAMunmapped`

default: **None**

string(s): output of unmapped reads in the SAM format

1st word:

None

no output

Within

output unmapped reads within the main SAM file (i.e. Aligned.out.sam)

2nd word:

KeepPairs

record unmapped mate for each alignment, and, in case of unsorted output, keep it adjacent to its mapped mate. Only affects multi-mapping reads.

`--outSAMorder`

default: **Paired**

string: type of sorting for the SAM output

Paired: one mate after the other for all paired alignments

PairedKeepInputOrder: one mate after the other for all paired alignments, the order is kept the same as in the input FASTQ files

`--outSAMprimaryFlag`

default: **OneBestScore**

string: which alignments are considered primary - all others will be marked with 0x100 bit in the FLAG

OneBestScore

only one alignment with the best score is primary

AllBestScore

all alignments with the best score are primary

`--outSAMreadID`

default: **Standard**

string: read ID record type

Standard

first word (until space) from the FASTx read ID line, removing /1,/2 from the end

Number

read number (index) in the FASTx file

--outSAMmapqUnique

default: 255

int: 0 to 255: the MAPQ value for unique mappers

--outSAMflagOR

default: 0

int: 0 to 65535: sam FLAG will be bitwise OR'd with this value, i.e. FLAG=FLAG — outSAMflagOR. This is applied after all flags have been set by STAR, and after outSAMflagAND. Can be used to set specific bits that are not set otherwise.

--outSAMflagAND

default: 65535

int: 0 to 65535: sam FLAG will be bitwise AND'd with this value, i.e. FLAG=FLAG & outSAMflagOR. This is applied after all flags have been set by STAR, but before outSAMflagOR. Can be used to unset specific bits that are not set otherwise.

--outSAMattrRGline

default: -

string(s): SAM/BAM read group line. The first word contains the read group identifier and must start with "ID:", e.g. -outSAMattrRGline ID:xxx CN:yy "DS:z z z".

xxx will be added as RG tag to each output alignment. Any spaces in the tag values have to be double quoted.

Comma separated RG lines corresponds to different (comma separated) input files in -readFilesIn. Commas have to be surrounded by spaces, e.g.

-outSAMattrRGline ID:xxx , ID:zzz "DS:z z" , ID:yyy DS:yyyy

--outSAMheaderHD

default: -

strings: @HD (header) line of the SAM header

`--outSAMheaderPG`

default: -

strings: extra @PG (software) line of the SAM header (in addition to STAR)

`--outSAMheaderCommentFile`

default: -

string: path to the file with @CO (comment) lines of the SAM header

`--outSAMfilter`

default: None

string(s): filter the output into main SAM/BAM files

[KeepOnlyAddedReferences](#)

only keep the reads for which all alignments are to the extra reference sequences added with `-genomeFastaFiles` at the mapping stage.

[KeepAllAddedReferences](#)

keep all alignments to the extra reference sequences added with `-genomeFastaFiles` at the mapping stage.

`--outSAMmultNmax`

default: -1

int: max number of multiple alignments for a read that will be output to the SAM/BAM files.

-1

all alignments (up to `-outFilterMultimapNmax`) will be output

`--outBAMcompression`

default: 1

int: -1 to 10 BAM compression level, -1=default compression (6?), 0=no compression, 10=maximum compression

`--outBAMsortingThreadN`

default: 0

int: ≥ 0 : number of threads for BAM sorting. 0 will default to $\min(6, \text{-runThreadN})$.

9.12 BAM processing

`--bamRemoveDuplicatesType`

default: `-`

string: mark duplicates in the BAM file, for now only works with (i) sorted BAM feeded with inputBAMfile, and (ii) for paired-end alignments only

`-`

no duplicate removal/markings

`UniqueIdentical`

mark all multimappers, and duplicate unique mappers. The coordinates, FLAG, CIGAR must be identical

`UniqueIdenticalNotMulti`

mark duplicate unique mappers but not multimappers.

`--bamRemoveDuplicatesMate2basesN`

default: `0`

int>0: number of bases from the 5' of mate 2 to use in collapsing (e.g. for RAMPAGE)

9.13 Output Wiggle

`--outWigType`

default: `None`

string(s): type of signal output, e.g. "bedGraph" OR "bedGraph read1_5p". Requires sorted BAM: `-outSAMtype BAM SortedByCoordinate`.

1st word:

`None`

no signal output

`bedGraph`

bedGraph format

`wiggle`

wiggle format

2nd word:

`read1_5p`

signal from only 5' of the 1st read, useful for CAGE/RAMPAGE etc

`read2`

signal from only 2nd read

`--outWigStrand`

default: `Stranded`

string: strandedness of wiggle/bedGraph output

Stranded

separate strands, str1 and str2

Unstranded

collapsed strands

--outWigReferencesPrefix

default: -

string: prefix matching reference names to include in the output wiggle file, e.g. "chr", default "-" - include all references

--outWigNorm

default: RPM

string: type of normalization for the signal

RPM

reads per million of mapped reads

None

no normalization, "raw" counts

9.14 Output Filtering

--outFilterType

default: Normal

string: type of filtering

Normal

standard filtering using only current alignment

BySJout

keep only those reads that contain junctions that passed filtering into SJ.out.tab

--outFilterMultimapScoreRange

default: 1

int: the score range below the maximum score for multimapping alignments

--outFilterMultimapNmax

default: 10

int: maximum number of loci the read is allowed to map to. Alignments (all of them) will be output only if the read maps to no more loci than this value.

Otherwise no alignments will be output, and the read will be counted as "mapped to too many loci" in the Log.final.out .

`--outFilterMismatchNmax`

default: 10

int: alignment will be output only if it has no more mismatches than this value.

`--outFilterMismatchNoverLmax`

default: 0.3

float: alignment will be output only if its ratio of mismatches to *mapped* length is less than or equal to this value.

`--outFilterMismatchNoverReadLmax`

default: 1.0

float: alignment will be output only if its ratio of mismatches to *read* length is less than or equal to this value.

`--outFilterScoreMin`

default: 0

int: alignment will be output only if its score is higher than or equal to this value.

`--outFilterScoreMinOverLread`

default: 0.66

float: same as outFilterScoreMin, but normalized to read length (sum of mates' lengths for paired-end reads)

`--outFilterMatchNmin`

default: 0

int: alignment will be output only if the number of matched bases is higher than or equal to this value.

`--outFilterMatchNminOverLread`

default: 0.66

float: same as outFilterMatchNmin, but normalized to the read length (sum of mates' lengths for paired-end reads).

`--outFilterIntronMotifs`

default: None

string: filter alignment using their motifs

None

no filtering

RemoveNoncanonical

filter out alignments that contain non-canonical junctions

RemoveNoncanonicalUnannotated

filter out alignments that contain non-canonical unannotated junctions when using annotated splice junctions database. The annotated non-canonical junctions will be kept.

--outFilterIntronStrands

default: **RemoveInconsistentStrands**

string: filter alignments

RemoveInconsistentStrands

remove alignments that have junctions with inconsistent strands

None

no filtering

9.15 Output Filtering: Splice Junctions

--outSJfilterReads

default: **All**

string: which reads to consider for collapsed splice junctions output

All: all reads, unique- and multi-mappers

Unique: uniquely mapping reads only

--outSJfilterOverhangMin

default: **30 12 12 12**

4 integers: minimum overhang length for splice junctions on both sides for: (1) non-canonical motifs, (2) GT/AG and CT/AC motif, (3) GC/AG and CT/GC motif, (4) AT/AC and GT/AT motif. -1 means no output for that motif

does not apply to annotated junctions

--outSJfilterCountUniqueMin

default: **3 1 1 1**

4 integers: minimum uniquely mapping read count per junction for: (1) non-canonical motifs, (2) GT/AG and CT/AC motif, (3) GC/AG and CT/GC motif, (4) AT/AC and GT/AT motif. -1 means no output for that motif

Junctions are output if one of outSJfilterCountUniqueMin OR outSJfilterCountTotalMin conditions are satisfied

does not apply to annotated junctions

--outSJfilterCountTotalMin

default: 3 1 1 1

4 integers: minimum total (multi-mapping+unique) read count per junction for: (1) non-canonical motifs, (2) GT/AG and CT/AC motif, (3) GC/AG and CT/GC motif, (4) AT/AC and GT/AT motif. -1 means no output for that motif

Junctions are output if one of outSJfilterCountUniqueMin OR outSJfilterCountTotalMin conditions are satisfied

does not apply to annotated junctions

--outSJfilterDistToOtherSJmin

default: 10 0 5 10

4 integers ≥ 0 : minimum allowed distance to other junctions' donor/acceptor

does not apply to annotated junctions

--outSJfilterIntronMaxVsReadN

default: 50000 100000 200000

N integers ≥ 0 : maximum gap allowed for junctions supported by 1,2,3,,N reads

i.e. by default junctions supported by 1 read can have gaps ≤ 50000 b, by 2 reads: ≤ 100000 b, by 3 reads: ≤ 200000 . by ≥ 4 reads any gap $\leq \text{alignIntronMax}$

does not apply to annotated junctions

9.16 Scoring

--scoreGap

default: 0

int: splice junction penalty (independent on intron motif)

--scoreGapNoncan

default: -8

int: non-canonical junction penalty (in addition to scoreGap)

--scoreGapGCAG

default: -4

GC/AG and CT/GC junction penalty (in addition to scoreGap)

--scoreGapATAC

default: -8

AT/AC and GT/AT junction penalty (in addition to scoreGap)

--scoreGenomicLengthLog2scale

default: -0.25

extra score logarithmically scaled with genomic length of the alignment:
 $\text{scoreGenomicLengthLog2scale} * \log_2(\text{genomicLength})$

--scoreDelOpen

default: -2

deletion open penalty

--scoreDelBase

default: -2

deletion extension penalty per base (in addition to scoreDelOpen)

--scoreInsOpen

default: -2

insertion open penalty

--scoreInsBase

default: -2

insertion extension penalty per base (in addition to scoreInsOpen)

--scoreStitchSJshift

default: 1

maximum score reduction while searching for SJ boundaries in the stitching step

9.17 Alignments and Seeding

`--seedSearchStartLmax`

default: 50

int>0: defines the search start point through the read - the read is split into pieces no longer than this value

`--seedSearchStartLmaxOverLread`

default: 1.0

float: seedSearchStartLmax normalized to read length (sum of mates' lengths for paired-end reads)

`--seedSearchLmax`

default: 0

int>=0: defines the maximum length of the seeds, if =0 max seed length is infinite

`--seedMultimapNmax`

default: 10000

int>0: only pieces that map fewer than this value are utilized in the stitching procedure

`--seedPerReadNmax`

default: 1000

int>0: max number of seeds per read

`--seedPerWindowNmax`

default: 50

int>0: max number of seeds per window

`--seedNoneLociPerWindow`

default: 10

int>0: max number of one seed loci per window

`--seedSplitMin`

default: 12

int>0: min length of the seed sequences split by Ns or mate gap

`--alignIntronMin`

default: 21

minimum intron size: genomic gap is considered intron if its length \geq alignIntronMin, otherwise it is considered Deletion

`--alignIntronMax`

default: 0

maximum intron size, if 0, max intron size will be determined by $(2^{\text{winBinNbits}}) * \text{winAnchorDistNbins}$

`--alignMatesGapMax`

default: 0

maximum gap between two mates, if 0, max intron gap will be determined by $(2^{\text{winBinNbits}}) * \text{winAnchorDistNbins}$

`--alignSJoverhangMin`

default: 5

int >0 : minimum overhang (i.e. block size) for spliced alignments

`--alignSJstitchMismatchNmax`

default: 0 -1 0 0

4*int \geq 0: maximum number of mismatches for stitching of the splice junctions (-1: no limit).

(1) non-canonical motifs, (2) GT/AG and CT/AC motif, (3) GC/AG and CT/GC motif, (4) AT/AC and GT/AT motif.

`--alignSJDBoverhangMin`

default: 3

int >0 : minimum overhang (i.e. block size) for annotated (sjdb) spliced alignments

`--alignSplicedMateMapLmin`

default: 0

int >0 : minimum mapped length for a read mate that is spliced

`--alignSplicedMateMapLminOverLmate`

default: 0.66

float>0: alignSplicedMateMapLmin normalized to mate length

--alignWindowsPerReadNmax

default: 10000

int>0: max number of windows per read

--alignTranscriptsPerWindowNmax

default: 100

int>0: max number of transcripts per window

--alignTranscriptsPerReadNmax

default: 10000

int>0: max number of different alignments per read to consider

--alignEndsType

default: Local

string: type of read ends alignment

- Local
 - standard local alignment with soft-clipping allowed
- EndToEnd
 - force end-to-end read alignment, do not soft-clip
- Extend5pOfRead1
 - fully extend only the 5p of the read1, all other ends: local alignment
- Extend5pOfReads12
 - fully extend only the 5p of the both read1 and read2, all other ends: local alignment

--alignEndsProtrude

default: 0 ConcordantPair

int, string: allow protrusion of alignment ends, i.e. start (end) of the +strand mate downstream of the start (end) of the -strand mate

1st word: int: maximum number of protrusion bases allowed

2nd word: string:

- ConcordantPair
 - report alignments with non-zero protrusion as concordant pairs
- DiscordantPair
 - report alignments with non-zero protrusion as discordant pairs

`--alignSoftClipAtReferenceEnds`

default: **Yes**

string: allow the soft-clipping of the alignments past the end of the chromosomes

Yes

allow

No

prohibit, useful for compatibility with Cufflinks

9.18 Windows, Anchors, Binning

`--winAnchorMultimapNmax`

default: 50

int>0: max number of loci anchors are allowed to map to

`--winBinNbits`

default: 16

int>0: $=\log_2(\text{winBin})$, where winBin is the size of the bin for the windows/clustering, each window will occupy an integer number of bins.

`--winAnchorDistNbins`

default: 9

int>0: max number of bins between two anchors that allows aggregation of anchors into one window

`--winFlankNbins`

default: 4

int>0: $\log_2(\text{winFlank})$, where win Flank is the size of the left and right flanking regions for each window

`--winReadCoverageRelativeMin`

default: 0.5

float \geq 0: minimum relative coverage of the read sequence by the seeds in a window, for STARlong algorithm only.

`--winReadCoverageBasesMin`

default: 0

int>0: minimum number of bases covered by the seeds in a window , for STARlong algorithm only.

9.19 Chimeric Alignments

`--chimOutType`

default: `SeparateSAMold`

string(s): type of chimeric output

1st word:

`SeparateSAMold`

output old SAM into separate Chimeric.out.sam file

`WithinBAM`

output into main aligned BAM files (Aligned.*.bam)

2nd word:

`WithinBAM HardClip`

hard-clipping in the CIGAR for supplemental chimeric alignments
(default if no 2nd word is present)

`WithinBAM SoftClip`

soft-clipping in the CIGAR for supplemental chimeric alignments

`--chimSegmentMin`

default: `0`

int \geq 0: minimum length of chimeric segment length, if ==0, no chimeric output

`--chimScoreMin`

default: `0`

int \geq 0: minimum total (summed) score of the chimeric segments

`--chimScoreDropMax`

default: `20`

int \geq 0: max drop (difference) of chimeric score (the sum of scores of all chimeric segments) from the read length

`--chimScoreSeparation`

default: `10`

int \geq 0: minimum difference (separation) between the best chimeric score and the next one

`--chimScoreJunctionNonGTAG`

default: `-1`

int: penalty for a non-GT/AG chimeric junction

`--chimJunctionOverhangMin`

default: 20

int \geq 0: minimum overhang for a chimeric junction

`--chimSegmentReadGapMax`

default: 0

int \geq 0: maximum gap in the read sequence between chimeric segments

`--chimFilter`

default: `banGenomicN`

string(s): different filters for chimeric alignments

`None`

no filtering

`banGenomicN`

Ns are not allowed in the genome sequence around the chimeric junction

`--chimMainSegmentMultNmax`

default: 10

int \geq 1: maximum number of multi-alignments for the main chimeric segment.
=1 will prohibit multimapping main segments.

9.20 Quantification of Annotations

`--quantMode`

default: -

string(s): types of quantification requested

-

none

`TranscriptomeSAM`

output SAM/BAM alignments to transcriptome into a separate file

`GeneCounts`

count reads per gene

`--quantTranscriptomeBAMcompression`

default: 1 1

int: -1 to 10 transcriptome BAM compression level, -1=default compression (6?), 0=no compression, 10=maximum compression

`--quantTranscriptomeBan`

default: `IndelSoftclipSingleend`

string: prohibit various alignment type

`IndelSoftclipSingleend`

prohibit indels, soft clipping and single-end alignments - compatible with RSEM

`Singleend`

prohibit single-end alignments

9.21 2-pass Mapping

`--twopassMode`

default: `None`

string: 2-pass mapping mode.

`None`

1-pass mapping

`Basic`

basic 2-pass mapping, with all 1st pass junctions inserted into the genome indices on the fly

`--twopass1readsN`

default: `-1`

int: number of reads to process for the 1st step. Use very large number (or default -1) to map all reads in the first step.