Assemble Transcripts with StringTie Element

StringTie is a fast and highly efficient assembler of RNA-Seq alignments into potential transcripts. It uses a novel network flow algorithm as well as an optional de novo assembly step to assemble and quantitate full-length transcripts representing multiple splice variants for each gene locus.

Parameters in GUI

Parameter	Description	Defaultvalu
Reference annotations	Use the reference annotation file (in GTF or GFF3 format) to guide the assembly process (-G).	
	The output will include expressed reference transcripts as well as any novel transcripts that are assembled.	
Reads orientation	Select the NGS libraries type: unstranded, stranded fr-secondstrand (fr), or stranded fr-firststand (rf).	Unstranded
Label	Use the specified string as the prefix for the name of the output transcripts (-I).	STRG
Min isoform fraction	Specify the minimum isoform abundance of the predicted transcripts as a fraction of the most abundant transcript assembled at a given locus (-f).	0.1
	Lower abundance transcripts are often artifacts of incompletely spliced precursors of processed transcripts.	
Min assembled ranscript length	Specify the minimum length for the predicted transcripts (-m).	200
Min anchor length for unctions	Junctions that don't have spliced reads that align them with at least this amount of bases on both sides are filtered out (-a).	10
Min junction coverage	There should be at least this many spliced reads that align across a junction (-j).	1
	This number can be fractional since some reads align in more than one place.	
	A read that aligns in n places will contribute 1/n to the junction coverage.	
Trim transcripts based on coverage	By default StringTie adjusts the predicted transcript's start and/or stop coordinates based on sudden drops in coverage of the assembled transcript.	True
	Set this parameter to "False" to disable the trimming at the ends of the assembled transcripts (-t).	
Min coverage for assembled transcripts	Specifies the minimum read coverage allowed for the predicted transcripts (-c).	2.5
issembled transcripts	A transcript with a lower coverage than this value is not shown in the output.	
	This number can be fractional since some reads align in more than one place. A read that aligns in n places will contribute 1/n to the coverage.	
Min locus gap separation	Reads that are mapped closer than this distance are merged together in the same processing bundle (-g).	50 bp
Fraction covered by multi-hit reads	Specify the maximum fraction of multiple-location-mapped reads that are allowed to be present at a given locus (-M).	0.95
	A read that aligns in n places will contribute 1/n to the coverage.	
Skip assembling for sequences	Ignore all read alignments (and thus do not attempt to perform transcript assembly) on the specified reference sequences (-x).	
	The value can be a single reference sequence name (e.g. "chrM") or a comma-delimited list of sequence names (e.g. "chrM,chrX,chrY").	
	This can speed up StringTie especially in the case of excluding the mitochondrial genome, whose genes may have very high coverage in some cases,	
	even though they may be of no interest for a particular RNA-Seq analysis.	
	The reference sequence names are case sensitive,	
	they must match identically the names of chromosomes/contigs of the target genome against which the RNA-Seq reads were aligned in the first place.	
Multi-mapping correction	Enables or disables (-u) multi-mapping correction.	Enabled
/erbose log	Enable detailed logging, if required (-v). The messages will be written to the UGENE log (enabling of "DETAILS" and "TRACE" logging may be required) and to the dashboard.	False
Number of threads	Specify the number of processing threads (CPUs) to use for transcript assembly (-p).	8
Output transcripts file	StringTie's primary output GTF file with assembled transcripts.	Auto
Enable gene abundance output	Select "True" to generate gene abundances output (-A). The output is written to a tab-delimited text file. Also, the file URL is passed to an output slot of the workflow element.	False
	I .	

Parameters in Workflow File Type: stringtie

Parameter	Parameter in the GUI	Туре
reference-annotations	Reference annotations	string
reads-orientation	Reads orientation	string
label	Label	string
min-isoform-fraction	Min isoform fraction	numeric
min-isoform-fraction	Min assembled transcript length	numeric
min-anchor-length	Min anchor length for junctions	numeric
min-junction-coverage	Min junction coverage	numeric
trim-transcripts	Trim transcripts based on coverage	bool
min-coverage	Min coverage for assembled transcripts	numeric
min-locus-gap	Min locus gap separation	numeric
multi-hit-fraction	Fraction covered by multi-hit reads	numeric
skip-sequences	Skip assembling for sequences	string
multi-mapping-correction	Multi-mapping correction	bool
verbose-log	Verbose log	bool
threads	Number of threads	numeric
transcripts-output-url	Output transcripts file	string
gene-abundance-output	Enable gene abundance output	bool

Input/Output Ports The element has 1 input port.

Name in GUI: Input BAM file(s)

Name in Workflow File: in

Slots:

Slot in GUI	Slot in Workflow File	Туре
Source URL	url	string

And 1 output port:

Name in GUI: StringTie output data

Name in Workflow File: out

Slots:

Slot in GUI	Slot in Workflow File	Туре
Output URL	url	string