



## Salmon.Indexer Documentation

**Description:** Indexing a transcriptome in order to perform quantification with Salmon and Alevin. See: [The Salmon User Guide](#) for detailed usage guidelines.

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**Contact:** Algorithm and scientific questions: <https://github.com/COMBINE-lab/salmon/issues>.  
Module specific issues: <https://genepattern.org/help>

**Summary:** In order to quantify gene expression using Salmon or Alevin, the transcriptome must first be indexed. This module performs this indexing step.

**Source Publication:** Patro R, Duggal G, Love MI, Irizarry RA, Kingsford C. Salmon provides fast and bias-aware quantification of transcript expression. Nat Methods. 2017 Apr;14(4):417-419. doi: 10.1038/nmeth.4197. Epub 2017 Mar 6. PMID: 28263959; PMCID: PMC5600148.

### Parameters:

Name	Description
GTF gz	A GTF file containing the genomic features to be indexed. Must be gzipped.
Transcriptome fa gz	A FASTA file of the transcript sequences corresponding to the organism's GTF file. Optional: if a Fasta file is not provided the module will attempt to use gffread to extract the sequences from the Genomic fasta file. Must be gzipped.
Genome fa gz	A FASTA file of the genomic sequences corresponding to the organism's genome assembly used for the transcript annotation. Must be gzipped.
kmer	An odd numbered integer. Used to index the transcriptome, used as the minimum acceptable length of a valid match. (Default: 31, for reads <70bp, consider adjusting this to a smaller value.)
Index Mode	Options for generating a decoy-aware transcriptome index Build a full decoy index (full), or a partial decoy index (partial). See parameter description for details.
Use Gencode	Gencode uses "I" characters in their FASTA record IDs, "true" allows salmon.index to be aware of these characters when processing gene IDs from records
output index name	The base name for the output indexed transcriptome

### Output Files:

Name	Description
<GTF.basename>.k<kmer>.salmon_ <Index.Mode>_decoy_index.tar.gz	A gzipped file containing the salmon index for downstream quantification.
<GTF.basename>.fa.gz	If a transcriptome fasta was not provided, this output is generated from the provided GTF file and genome fasta file using GFFread.

# GenePattern

**Module Language:** Shell script

**Source Repository:** <https://github.com/genepattern/Salmon.Indexer/releases/tag/v0.4>

**Docker image:** [genepattern/salmon-indexer:beta](https://hub.docker.com/r/genepattern/salmon-indexer)

Version	Comment
0.4	Initial release.