

# Biostrings Quick Overview

Hervé Pagès  
Fred Hutchinson Cancer Research Center  
Seattle, WA

March 13, 2024

Most but not all functions defined in the **Biostrings** package are summarized here.

Function	Description
length	Return the number of sequences in an object.
names	Return the names of the sequences in an object.
[	Extract sequences from an object.
head, tail	Extract the first or last sequences from an object.
rev	Reverse the order of the sequences in an object.
c	Combine in a single object the sequences from 2 or more objects.
width, nchar	Return the sizes (i.e. number of letters) of all the sequences in an object.
==, !=	Element-wise comparison of the sequences in 2 objects.
match, %in%	Analog to <code>match</code> and <code>%in%</code> on character vectors.
duplicated, unique	Analog to <code>duplicated</code> and <code>unique</code> on character vectors.
sort, order	Analog to <code>sort</code> and <code>order</code> on character vectors, except that the ordering of DNA or Amino Acid sequences doesn't depend on the locale.
relist, split, extractList	Analog to <code>relist</code> and <code>split</code> on character vectors, except that the result is a <code>DNAStringSetList</code> or <code>AAStringSetList</code> object. <code>extractList</code> is a generalization of <code>relist</code> and <code>split</code> that supports <i>arbitrary</i> groupings.

Table 1: Low-level manipulation of `DNAStringSet` and `AAStringSet` objects.

Function	Description
alphabetFrequency	Tabulate the letters (all the letters in the alphabet for <code>alphabetFrequency</code> , only the specified letters for <code>letterFrequency</code> ) in a sequence or set of sequences.
letterFrequency	Only the specified letters for <code>letterFrequency</code> ) in a sequence or set of sequences.
uniqueLetters	Extract the unique letters from a sequence or set of sequences.
letterFrequencyInSlidingView	Specialized version of <code>letterFrequency</code> that tallies the requested letter frequencies for a fixed-width view that is conceptually slid along the input sequence.
consensusMatrix	Computes the consensus matrix of a set of sequences.
dinucleotideFrequency	Fast 2-mer, 3-mer, and k-mer counting for DNA or RNA.
trinucleotideFrequency	
oligonucleotideFrequency	
nucleotideFrequencyAt	Tallies the short sequences formed by extracting the nucleotides found at a set of fixed positions from each sequence of a set of DNA or RNA sequences.

Table 2: Counting / tabulating.

Function	Description
reverse	Compute the reverse, complement, or reverse-complement, of a set of DNA sequences.
complement	
reverseComplement	
translate	Translate a set of DNA sequences into a set of Amino Acid sequences.
chartr	Replace letters in a sequence or set of sequences.
replaceAmbiguities	
subseq, subseq<-	Extract/replace arbitrary substrings from/in a string or set of strings.
extractAt, replaceAt	
replaceLetterAt	Replace the letters specified by a set of positions by new letters.
padAndClip, stackStrings	Pad and clip strings.
strsplit, unstrsplit	strsplit splits the sequences in a set of sequences according to a pattern. unstrsplit is the reverse operation i.e. a fast implementation of sapply(x, paste0, collapse=sep) for collapsing the list elements of a <i>DNAStringSetList</i> or <i>AAStringSetList</i> object.

Table 3: Sequence transformation and editing.

Function	Description
matchPattern	Find/count all the occurrences of a given pattern (typically short) in a reference sequence (typically long). Support mismatches and indels.
countPattern	
vmatchPattern	Find/count all the occurrences of a given pattern (typically short) in a set of reference sequences. Support mismatches and indels.
vcountPattern	
matchPDict	Find/count all the occurrences of a set of patterns in a reference sequence. (whichPDict only identifies which patterns in the set have at least one match.) Support a small number of mismatches.
countPDict	
whichPDict	
vmatchPDict	[Note: vmatchPDict not implemented yet.] Find/count all the occurrences of a set of patterns in a set of reference sequences. (whichPDict only identifies for each reference sequence which patterns in the set have at least one match.) Support a small number of mismatches.
vcountPDict	
vwhichPDict	
pairwiseAlignment	Solve (Needleman-Wunsch) global alignment, (Smith-Waterman) local alignment, and (ends-free) overlap alignment problems.
matchPWM	Find/count all the occurrences of a Position Weight Matrix in a reference sequence.
countPWM	
trimLRPatterns	Trim left and/or right flanking patterns from sequences.
matchLRPatterns	Find all paired matches in a reference sequence i.e. matches specified by a left and a right pattern, and a maximum distance between them.
matchProbePair	Find all the amplicons that match a pair of probes in a reference sequence.
findPalindromes	Find palindromic regions in a sequence.

Table 4: String matching / alignments.

Function	Description
readBStringSet readDNAStringSet readRNAStringSet readAAStringSet	Read ordinary/DNA/RNA/Amino Acid sequences from files (FASTA or FASTQ format).
writeXStringSet	Write sequences to a file (FASTA or FASTQ format).
writePairwiseAlignments	Write pairwise alignments (as produced by pairwiseAlignment) to a file (“pair” format).
readDNAMultipleAlignment readRNAMultipleAlignment readAAMultipleAlignment	Read multiple alignments from a file (FASTA, “stockholm”, or “clustal” format).
write.phylip	Write multiple alignments to a file (Phylip format).

Table 5: I/O functions.

Function	Description
stringDist	Computes the matrix of Levenshtein edit distances, or Hamming distances, or pairwise alignment scores, for a set of strings.

Table 6: Miscellaneous.