## NAME

InfoSequenceFiles.pl - List information about sequence and alignment files

#### **SYNOPSIS**

InfoSequenceFiles.pl SequenceFile(s) AlignmentFile(s)...

InfoSequenceFiles.pl [-a, --all] [-c, --count] [-d, --detail infolevel] [-f, --frequency] [--FrequencyBins number | "number, number, [number,...]"] [-h, --help] [-i, --I gnoreGaps yes | no] [-I, --longest] [-s, --shortest] [--SequenceLengths] [-w, --workingdir dirname] SequenceFile(s)...

#### **DESCRIPTION**

List information about contents of <code>SequenceFile(s)</code> and <code>AlignmentFile(s)</code>: number of sequences, shortest and longest sequences, distribution of sequence lengths and so on. The file names are separated by spaces. All the sequence files in a current directory can be specified by <code>\*.aln, \*.msf, \*.fasta, \*.fta, \*.pir</code> or any other supported formats; additionally, <code>DirName</code> corresponds to all the sequence files in the current directory with any of the supported file extension: <code>.aln, .msf, .fasta, .fta, and .pir</code>.

Supported sequence formats are: ALN/CLustalW, GCG/MSF, PILEUP/MSF, Pearson/FASTA, and NBRF/PIR. Instead of using file extensions, file formats are detected by parsing the contents of SequenceFile(s) and AlignmentFile(s).

## **OPTIONS**

#### -a, --all

List all the available information.

#### -c, --count

List number of of sequences. This is default behavior.

## -d, --detail InfoLevel

Level of information to print about sequences during various options. Default: 1. Possible values: 1, 2 or 3.

# -f, --frequency

List distribution of sequence lengths using the specified number of bins or bin range specified using FrequencyBins option.

This option is ignored for input files containing only single sequence.

## --FrequencyBins number | "number,number,[number,...]"

This value is used with -f, --frequency option to list distribution of sequence lengths using the specified number of bins or bin range. Default value: 10.

The bin range list is used to group sequence lengths into different groups; It must contain values in ascending order. Examples:

```
100,200,300,400,500,600 200,400,600,800,1000
```

The frequency value calculated for a specific bin corresponds to all the sequence lengths which are greater than the previous bin value and less than or equal to the current bin value.

#### -h, --help

Print this help message.

## -i, --I gnoreGaps yes | no

Ignore gaps during calculation of sequence lengths. Possible values: yes or no. Default value: no.

# -I, --longest

List information about longest sequence: ID, sequence and sequence length. This option is ignored for input files containing only single sequence.

#### -s, --shortest

List information about shortest sequence: ID, sequence and sequence length. This option is ignored for input files containing only single sequence.

#### --SequenceLengths

List information about sequence lengths.

## -w, --WorkingDir dirname

Location of working directory. Default: current directory.

#### **EXAMPLES**

To count number of sequences in sequence files, type:

```
% InfoSequenceFiles.pl Sample1.fasta
% InfoSequenceFiles.pl Sample1.msf Sample1.aln Sample1.pir
% InfoSequenceFiles.pl *.fasta *.fta *.msf *.pir *.aln
```

To list all available information with maximum level of available detail for a sequence alignment file Sample1.msf, type:

```
% InfoSequenceFiles.pl -a -d 3 Sample1.msf
```

To list sequence length information after ignoring sequence gaps in Sample1.aln file, type:

```
% InfoSequenceFiles.pl --SequenceLengths --IgnoreGaps Yes
Sample1.aln
```

To list shortest and longest sequence length information after ignoring sequence gaps in Sample1.aln file, type:

```
% InfoSequenceFiles.pl --longest --shortest --IgnoreGaps Yes
Sample1.aln
```

To list distribution of sequence lengths after ignoring sequence gaps in Sample1.aln file and report the frequency distribution into 10 bins, type:

```
% InfoSequenceFiles.pl --frequency --FrequencyBins 10
--IgnoreGaps Yes Sample1.aln
```

To list distribution of sequence lengths after ignoring sequence gaps in Sample1.aln file and report the frequency distribution into specified bin range, type:

```
% InfoSequenceFiles.pl --frequency --FrequencyBins
"150,200,250,300,350" --IgnoreGaps Yes Sample1.aln
```

## **AUTHOR**

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## SEE ALSO

 $Analyze Sequence Files Data.pl,\ Extract From Sequence Files.pl,\ Info Amino Acids.pl,\ Info Nucleic Acids.pl$ 

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