

Package ‘PhyloWGA’

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Title PhyloWGA: a user-friendly framework for chromosome-aware phylogenetic interrogation of whole ge-nome alignments

Version 0.0.0.9000

Description PhyloWGA Contains a suite of functions of phylogenetic analyses and interrogation of WGAs

License What license it uses

Encoding UTF-8

LazyData true

RoxygenNote 7.1.0

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Chromo.Crawl	<i>Chromo.Crawl: function to crawl over an alignment of chromosomes while attempting to concatenate adjacent genomic windows</i>
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Description

This function writes the results to an output file. Additionally, this function provides a directory where the resulting supergenes are placed.

Usage

```
Chromo.Crawl(
  string.PathParentDir,
  numeric.WindowSize,
  numeric.StepSize,
  string.PathToFastaFile,
  numeric.NumberOfCores
)
```

Arguments

```
string.PathParentDir
    Path to parent dir for all analyses
numeric.WindowSize
    Size of each chromosomal window
numeric.StepSize
    Spacing of each chromosomal window
string.PathToFastaFile
    Path to fasta file
numeric.NumberOfCores
    Number of cores
```

Value

Results Results written to outputfiles

Examples

```
#####
# Read example chromosome alignment #
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####
# Conduct.ChromoPhylome #
#####
Chromo.Crawl(string.PathParentDir = '~/Desktop/',
              numeric.WindowSize = 1000,
              numeric.StepSize = 1000,
              string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)
```

Chromo.Phylome

Chromo.Phylome: function to infer a chromosome-specific set of phylogenetic trees

Description

This function returns a list of the phylogenetic tree models, one for each window set by the user-defined window and step size input parameters

Usage

```
Chromo.Phylome(  
  string.PathParentDir,  
  string.PathToFastaFile,  
  numeric.WindowSize,  
  numeric.StepSize,  
  string.Commands_iqtree  
)
```

Arguments

<code>string.PathParentDir</code>	Path to parent dir for all analyses
<code>string.PathToFastaFile</code>	Path to fasta file
<code>numeric.WindowSize</code>	Size of each chromosomal window
<code>numeric.StepSize</code>	Spacing of each chromosomal window
<code>string.Commands_iqtree</code>	String used for nucleotide substitution model for IqTree. Leave as "" for model selection

Value

Results Results writted to output files

Examples

[illegible]

Chromo.Phylome.Custom *Chromo.Phylome.Custom: function to infer a chromosome-specific set of phylogenetic trees*

Description

This function returns a list of the phylogenetic tree models, one for each window set by the user-defined window and step size input parameters

Usage

```
Chromo.Phylome.Custom(
  string.PathParentDir,
  string.PathToFastaFile,
  matrix.WindowCoordinates,
  string.Commands_iqtree
)
```

Arguments

```
string.PathParentDir
    Path to parent dir for all analyses
string.PathToFastaFile
    Path to fasta file
string.Commands_iqtree
    String used for nucleotide substitution model for IqTree. Leave as "" for
    model selection
numeric.WindowSize
    Size of each chromosomal window
numeric.StepSize
    Spacing of each chromosomal window
```

Value

Results Results writted to output files

Define.PhyloWGA_Experiment *Define.PhyloWGA_Experiment: function to generate a matrix of window coordinates for a given alignment*

Description

This function returns a matrix of the window coordinates for an alignment

Usage

```
Define.PhyloWGA_Experiment(
  numeric.WindowSize,
  numeric.StepSize,
  numeric.TotalLength
)
```

Arguments

numeric.WindowSize
 Number of bp for each window, window length

numeric.StepSize
 Number of bp separating each window

numeric.TotalLength
 Total number of bp for the entire alignment

Value

matrix.WindowCoordinates Matrix of k x 2 dimensions, each row indicating the start and end coordinates for the respective window

Examples

```
#####
# Specify experimental parameters #
#####
matrix.ExperimentalParams <- Define.PhyloWGA_Experiment(numeric.WindowSize = 1000, numeric.StepSize = 500, nu
```

Execute.PhylogeneticCongruence_concatepillar

Execute.PhylogeneticCongruence_concatepillar: function to execute concatepillar on a pair of alignments

Description

This function returns a list of results from concatepillar

Usage

```
Execute.PhylogeneticCongruence_concatepillar(
  matrix.WindowAlignment_01,
  matrix.WindowAlignment_02,
  string.PathParentDir,
  numeric.NumberOfCores
)
```

Arguments

matrix.WindowAlignment_01
 Matrix of alignment for the first window

matrix.WindowAlignment_02
 Matrix of alignment for the second window

string.PathParentDir
 Path to parent dir for all analyses

numeric.NumberOfCores
 Number of cores

Value

List List of results from concatepillar

Examples

```
#####
# Load depends #
#####
library(PhyloWGA)
library(ape)

#####
# Read example chromosome alignment #
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####
# Get alignment length #
#####
numeric.AlignmentLength <- Get.AlignmentLength(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)

#####
# Get sequence names #
#####
vector.SequenceNames <- Get.SequenceNames(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)

#####
# Define experimental parameters #
#####
matrix.ExperimentalParams <- Define.PhyloWGA_Experiment(numeric.WindowSize = 1000, numeric.StepSize = 1000, n

#####
# Extract window alignment #
#####
matrix.WindowAlignment_01 <- Extract.WindowAlignment(string.PathToFastaFile = String.Path_ExampleChromosomeA
               vector.SequenceNames = vector.SequenceNames,
               vector.Coordinates_ExperimentalLocus = matrix.ExperimentalParams[5,])

#####
# Prune taxa that are only missin data from matrix #
#####
matrix.Pruned_WindowAlignment_01 <- Prune.MissingDataTaxa(matrix.WindowAlignment = matrix.WindowAlignment_01)

#####
# Extract window alignment #
#####
matrix.WindowAlignment_02 <- Extract.WindowAlignment(string.PathToFastaFile = String.Path_ExampleChromosomeA
               vector.SequenceNames = vector.SequenceNames,
               vector.Coordinates_ExperimentalLocus = matrix.ExperimentalParams[6,])

#####
# Prune taxa that are only missin data from matrix #
#####
```

```

matrix.Pruned_WindowAlignment_02 <- Prune.MissingDataTaxa(matrix.WindowAlignment = matrix.WindowAlignment_02

#####
# Estimate model with iqtree #
#####
Execute.PhylogeneticCongruence_concatepillar(matrix.Pruned_WindowAlignment_01, matrix.Pruned_WindowAlignmen

```

Execute.PhylogeneticInference_iqtree

Execute.PhylogeneticInference_iqtree: function to estimate a phylogenetic tree model for a window with iqtree

Description

This function returns a list containing parameter estimates and a phylogenetic tree estimate

Usage

```

Execute.PhylogeneticInference_iqtree(
  matrix.WindowAlignment,
  string.PathParentDir,
  string.Commands_iqtree
)

```

Arguments

```

matrix.WindowAlignment
    Matrix of the alignment for a window
string.PathParentDir
    Path used for estimating trees with iqtree

```

Value

List List containing (1) list of parameter estimates and (2) phylogenetic tree estimate

Examples

```

#####
# Load depends #
#####
library(PhyloWGA)
library(ape)

#####
# Read example chromosome alignment #
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####

```

```

# Get alignment length #
#####
numeric.AlignmentLength <- Get.AlignmentLength(string.PathToFastaFile = String.Path_ExampleChromosomeAlignme

#####
# Get sequence names #
#####
vector.SequenceNames <- Get.SequenceNames(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)

#####
# Define experimental parameters #
#####
matrix.ExperimentalParams <- Define.PhyloWGA_Experiment(numeric.WindowSize = 1000, numeric.StepSize = 1000, n

#####
# Extract window alignment #
#####
matrix.WindowAlignment <- Extract.WindowAlignment(string.PathToFastaFile = String.Path_ExampleChromosomeAlign
vector.SequenceNames = vector.SequenceNames,
vector.Coordinates_ExperimentalLocus = matrix.ExperimentalParams[5,])

#####
# Prune taxa that are only missin data from matrix #
#####
matrix.Pruned_WindowAlignment <- Prune.MissingDataTaxa(matrix.WindowAlignment = matrix.WindowAlignment, nume

#####
# Estimate model with iqtree #
#####
Execute.PhylogeneticInference_iqtree(matrix.WindowAlignment = matrix.Pruned_WindowAlignment, string.PathPar

```

Extract.WindowAlignment

Extract.WindowAlignment: function to extract an alignment matrix given an input vector of sequences, coordinates, and a path to a fasta file

Description

This function returns matrix consisting of an extracted alignment

Usage

```

Extract.WindowAlignment(
  string.PathToFastaFile,
  vector.SequenceNames,
  vector.Coordinates_ExperimentalLocus
)

```


Arguments

`string.PathToFastaFile`
String defining the path to the input fasta file

`vector.SequenceNames`
Vector of sequence names to be extracted

`vector.Coordinates_ExperimentalLocus`
Vector containing two coordinates for the window (start and end)

Value

`matrix.Window_Alignment` Matrix containing the alignment for the input window coordinates from the fasta file

Examples

```
#####
# Load depends #
#####
library(PhyloWGA)
library(ape)

#####
# Read example chromosome alignment #
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####
# Get alignment length #
#####
numeric.AlignmentLength <- Get.AlignmentLength(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)

#####
# Get sequence names #
#####
vector.SequenceNames <- Get.SequenceNames(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)

#####
# Define experimental parameters #
#####
matrix.ExperimentalParams <- Define.PhyloWGA_Experiment(numeric.WindowSize = 1000, numeric.StepSize = 1000, n

#####
# Extract window alignment #
#####
matrix.WindowAlignment <- Extract.WindowAlignment(string.PathToFastaFile = String.Path_ExampleChromosomeAlign
                                                    vector.SequenceNames = vector.SequenceNames,
                                                    vector.Coordinates_ExperimentalLocus = matrix.ExperimentalParams[5,])

write.dna(x = matrix.WindowAlignment, file = '~/Desktop/ExampleWindow.fasta', colsep = "", format = "fasta")
```

Get.AlignmentLength	<i>Get.AlignmentLength: get the total length of an alignment in the input fasta file</i>
---------------------	--

Description

This function returns the number of bp found in the input fasta file

Usage

```
Get.AlignmentLength(string.PathToFastaFile)
```

Arguments

string.PathToFastaFile
String defining the path to the input fasta file

Value

numeric.Total_AlignmentLength Number of base pairs in the input fasta file

Examples

```
#####
# Read example chromosome alignment #
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####
# Get alignment length #
#####
numeric.AlignmentLength <- Get.AlignmentLength(string.PathToFastaFile = String.Path_ExampleChromosomeAlignme
```

Get.SequenceNames	<i>Get.SequenceNames: get a vector of the sequence names provided in a fasta file</i>
-------------------	---

Description

This function returns a vector of sequence names

Usage

```
Get.SequenceNames(string.PathToFastaFile)
```

Arguments

string.PathToFastaFile
String defining the path to the input fasta file

Value

vector.SequenceNames Vector containing the set of sequence names obtained from fasta file

Examples

```
#####
# Read example chromosome alignment #
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####
# Get alignment length #
#####
vector.SequenceNames <- Get.SequenceNames(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)
```

Organize.ParallelPhyloWGA

Organize.ParallelPhyloWGA: function to partition and organize subdirectories for parallel WGA analyses

Description

This function returns a XXX

Usage

```
Organize.ParallelPhyloWGA(
  numeric.NumberSubsets,
  string.PathToFastaFile,
  string.PathParentDir,
  string.Analysis,
  numeric.WindowSize,
  string.Commands_iqtree,
  numeric.StepSize,
  numeric.NumberOfCores
)
```

Arguments

numeric.NumberSubsets
Number of subsets for the original alignment and analyses

string.PathToFastaFile
String defining the path to the input fasta file

string.PathParentDir
Path to parent dir for all analyses

numeric.WindowSize
Size of each chromosomal window

numeric.StepSize
Spacing of each chromosomal window

numeric.NumberOfCores
Number of cores

Value

XXX XXX

Prune.MissingDataTaxa *Prune.MissingDataTaxa: function to pruned matrix (high-missing data sequences removed)*

Description

This function returns a matrix that has been pruned of sequences with missing data above a given threshold

Usage

```
Prune.MissingDataTaxa(matrix.WindowAlignment, numeric.MissingDataThreshold)
```

Arguments

matrix.WindowAlignment
Matrix of input chromosomal window alignment

numeric.MissingDataThreshold
Numeric (proportion) of missing data permitted of any one sequence

Value

matrix.Pruned_WindowAlignment Matrix of window alignment with sequences with

Examples

```
#####
# Load depends #
#####
library(PhyloWGA)
library(ape)

#####
# Read example chromosome alignment #
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####
# Get alignment length #
#####
numeric.AlignmentLength <- Get.AlignmentLength(string.PathToFastaFile = String.Path_ExampleChromosomeAlignme

#####
# Get sequence names #
#####
vector.SequenceNames <- Get.SequenceNames(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)
```

```
#####
# Define experimental parameters #
#####
matrix.ExperimentalParams <- Define.PhyloWGA_Experiment(numeric.WindowSize = 1000, numeric.StepSize = 1000, n

#####
# Extract window alignment #
#####
matrix.WindowAlignment <- Extract.WindowAlignment(string.PathToFastaFile = String.Path_ExampleChromosomeAlign
vector.SequenceNames = vector.SequenceNames,
vector.Coordinates_ExperimentalLocus = matrix.ExperimentalParams[5,])

#####
# Prune taxa that are only missin data from matrix #
#####
matrix.Pruned_WindowAlignment <- Prune.MissingDataTaxa(matrix.WindowAlignment = matrix.WindowAlignment, nume
```

Read.OutputFile_iqtree

Read.OutputFile_iqtree: function to extract a list of parameter estimates from an iqtree outputfile

Description

This function returns a list of the parameter estimates

Usage

```
Read.OutputFile_iqtree(string.Path_OutputFile_iqtree)
```

Arguments

```
string.Path_OutputFile_iqtree
String defining path to the output iqtree file
```

Value

List List of containing estimates of the GTR+G model

Examples

```
#####
# Load depends #
#####
library(PhyloWGA)
library(ape)

#####
# Read example chromosome alignment #
```

```
#####
String.Path_ExampleChromosomeAlignment <- system.file("extdata", "Example_Chr10.fasta", package="PhyloWGA")

#####
# Get alignment length #
#####
numeric.AlignmentLength <- Get.AlignmentLength(string.PathToFastaFile = String.Path_ExampleChromosomeAlignme

#####
# Get sequence names #
#####
vector.SequenceNames <- Get.SequenceNames(string.PathToFastaFile = String.Path_ExampleChromosomeAlignment)

#####
# Define experimental parameters #
#####
matrix.ExperimentalParams <- Define.PhyloWGA_Experiment(numeric.WindowSize = 1000, numeric.StepSize = 1000, n

#####
# Extract window alignment #
#####
matrix.WindowAlignment <- Extract.WindowAlignment(string.PathToFastaFile = String.Path_ExampleChromosomeAlign
vector.SequenceNames = vector.SequenceNames,
vector.Coordinates_ExperimentalLocus = matrix.ExperimentalParams[5,])

#####
# Prune taxa that are only missin data from matrix #
#####
matrix.Pruned_WindowAlignment <- Prune.MissingDataTaxa(matrix.WindowAlignment = matrix.WindowAlignment, nume

#####
# Estimate model with iqtree #
#####
Execute.PhylogeneticInference_iqtree(matrix.WindowAlignment = matrix.Pruned_WindowAlignment, string.PathPare

#####
# Read output file #
#####
Read.OutputFile_iqtree(string.Path_OutputFile_iqtree = '~/Desktop/PhylogeneticInference_iqtree/WindowAlignm
```

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