

Package ‘FIREVAT’

March 23, 2019

Type Package

Title FIREVAT, FInding REliable Variants without ArTifacts

Description FIREVAT is a variant filtering tool for cancer sequencing data, which uses mutational signatures to identify sequencing artifacts and low-quality variants.

Version 0.3.1

Authors Andy Jinseok Lee, Hyunbin Kim

Maintainer Andy Jinseok Lee <jinseok.lee@ncc.re.kr>, Hyunbin Kim <khb7840@ncc.re.kr>

Imports data.table,
stringi,
bedr,
GA,
jsonlite,
yaml,
MutationalPatterns,
deconstructSigs,
lsa,
BSgenome.Hsapiens.UCSC.hg19,
BSgenome.Hsapiens.UCSC.hg38,
ggpubr,
caTools,
ggrepel,
gridExtra,
ggplot2,
rmarkdown,
gtable,
dplyr,
foreach,
doParallel

URL <https://github.com/cgab-ncc/FIREVAT>

License GPL-2

Encoding UTF-8

LazyData true

RoxygenNote 6.1.1

Suggests knitr

VignetteBuilder knitr

R topics documented:

AnnotateVCFObj	3
CheckIfVariantRefinementIsNecessary	4
Chromosome.Names	5
ComputeZScore	5
ComputeZScoreEquiValue	6
DecimalCeiling	6
Default.Obj.Fn	7
DefaultFilterToBinary	7
EnumerateTriNucCounts	8
Euc.Exp.Weighted.Obj.Fn	8
Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.1	9
Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.2	9
Euc.Obj.Fn	10
Exp.Weighted.Obj.Fn.1	10
Exp.Weighted.Obj.Fn.2	11
Exp.Weighted.Refined.Seq.Art.Only.Obj.Fn	11
FilterVCF	12
GenerateConfigObj	12
GetCOSMICMutSigs	13
GetCOSMICMutSigsEtiologiesColors	13
GetCOSMICMutSigsNames	14
GetGASuggestedSolutions	14
GetOptimizedSignatures	15
GetParameterLowerUpperVector	16
GetPCAWGMutSigs	16
GetPCAWGMutSigsEtiologiesColors	17
GetPCAWGMutSigsNames	17
InitializeVCF	17
MakeFilter	18
MutaliskParseVCFObj	18
MutPatParseRefMutSigs	19
MutPatParseVCFObj	19
ParameterToBits	20
ParseConfigFile	21
PCAWG.All.Sequencing.Artifact.Signatures	21
PCAWG.Known.Sequencing.Artifact.Signatures	22
PCAWG.Likely.Sequencing.Artifact.Signatures	22
PCAWG.Possible.Sequencing.Artifact.Signatures	22
PCAWG.Target.Mutational.Signatures	23
PerformStrandBiasAnalysis	23
PlotMutaliskResults	24

PlotMutationTypes	25
PlotOptimizationIterations	26
PlotSignaturesContProbs	27
PlotTable	28
PlotTriNucSpectrum	28
PlotVCFStatsBoxPlots	29
PlotVCFStatsHistograms	30
PrepareAnnotationDB	31
PrepareArtifactAnnotationTable	32
PrepareArtifactStrandBiasTable	32
PrepareArtifactualMutsOptimizationIterationsPlot	33
PrepareFilterCutoffsTable	33
PrepareGeneticAlgorithmParametersTable	34
PrepareIdentifiedSignaturesPlot	34
PrepareMLEReconstructedSpectrumsPlot	35
PrepareNucleotideSubstitutionTypesPlot	35
PrepareObservedSpectrumsPlot	36
PrepareOptimizationResultsTable	36
PrepareOptimizedVCFStatisticsPlot	37
PrepareRefinedAnnotationTable	37
PrepareRefinedMutsOptimizationIterationsPlot	38
PrepareRefinedStrandBiasTable	38
PrepareResidualSpectrumsPlot	39
PrepareTrinucleotideSpectrumsTable	39
QueryAnnotatedVCF	40
ReadOptimizationIterationReport	40
ReadVCF	41
ReportFIREVATResults	41
RunFIREVAT	42
RunMutalisk	44
RunMutaliskHelper	46
RunMutPat	47
TriNuc.Mutation.Type.Hex.Colors	48
UpdateFilter	48
WriteFIREVATResultsToTSV	49
WriteVCF	49
Index	50

AnnotateVCFObj

*AnnotateVCFObj***Description**

Annotates a vcf.obj using df.variants.of.interest (from [PrepareAnnotationDB](#))

Usage

```
AnnotateVCFObj(vcf.obj, df.annotation.db, columns.to.include,
               include.all.columns = FALSE)
```

Arguments

`vcf.obj` [ReadVCF](#)

`df.annotation.db`
A data.frame from [PrepareAnnotationDB](#). This data.frame must have the columns 'CHROM', 'POS', 'REF', 'ALT'

`columns.to.include`
A character vector of columns to include. Note that existing columns in `vcf.obj` will not be affected.

`include.all.columns`
A boolean value. If TRUE, then annotates `vcf.obj` with all columns present in `df.variants.of.interest`. If FALSE, `columns.to.include` must be supplied.

Value

An annotated `vcf.obj`

CheckIfVariantRefinementIsNecessary

CheckIfVariantRefinementIsNecessary

Description

Checks if variant refinement is necessary by identifying mutational signatures related to sequencing artifact in the `vcf.obj` (set of original unrefined point mutations).

Usage

```
CheckIfVariantRefinementIsNecessary(vcf.obj, bsg, df.mut.pat.ref.sigs,
                                     target.mut.sigs, sequencing.artifact.mut.sigs,
                                     init.artifact.stop = 0.05, verbose = TRUE)
```

Arguments

`vcf.obj` A list from [ReadVCF](#)

`bsg` [BSgenome.Hsapiens.UCSC](#) object

`df.mut.pat.ref.sigs`
A data.frame from [MutPatParseRefMutSigs](#)

`target.mut.sigs`
A character vector of target mutational signatures from reference mutational signatures.

sequencing.artifact.mut.sigs	A character vector of sequencing artifact mutational signatures from reference mutational signatures.
init.artifact.stop	Numeric value less than 1. If the sum of sequencing artifact weights in vcf.obj is less than or equal to this value then this function returns judgment = FALSE, otherwise returns judgment = TRUE.
verbose	If TRUE, provides process detail. Default value is TRUE.

Value

- A list with the following elements
- judgmentA boolean value
 - seq.art.sigs.weights.sumA numeric value. Sum of sequencing artifact weights.

Chromosome.Names	Constant
------------------	----------

Description

Chromosome names for FIREVAT. Chromosome names should be given in the format of "chr" + chromosome number.

Usage

Chromosome.Names

Format

An object of class character of length 25.

ComputeZScore	ComputeZScore
---------------	---------------

Description

Returns a z-score of x given a distribution of values

Usage

ComputeZScore(values, x)

Arguments

values	a numeric vector
x	a numeric value

Value

a numeric value corresponding to the z-score of x

ComputeZScoreEquiValue
<i>ComputeZScoreEquiValue</i>

Description

Returns a numeric value that is equivalent to the specified z.score in the distribution of 'values'

Usage

ComputeZScoreEquiValue(z.score, values)

Arguments

z.score	numeric value
values	numeric vector

Value

a numeric value corresponding to the specified z.score in the 'values' distribution

DecimalCeiling	<i>DecimalCeiling</i>
----------------	-----------------------

Description

Returns the ceiling of a decimal value e.g. value = 0.15, decimal = 0.1 returns 0.2

Usage

DecimalCeiling(value, decimal)

Arguments

value	numeric value (decimal)
decimal	numeric value (e.g. 0.1, 0.001)

Value

a numeric value

Default.Obj.Fn	<i>Default.Obj.Fn</i>
----------------	-----------------------

Description

Calculates the default objective value for FIREVAT GA optimization.

Usage

```
Default.Obj.Fn(C.refined, A.refined, C.artifactual, A.artifactual)
```

Arguments

C.refined	A numeric value between 0 and 1.
A.refined	A numeric value between 0 and 1.
C.artifactual	A numeric value between 0 and 1.
A.artifactual	A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

DefaultFilterToBinary	<i>Transform default filtering parameters to a binary vector</i>
-----------------------	--

Description

This function transforms default filtering parameter to binary vector which can be used as a suggested solution in GA algorithm.

Usage

```
DefaultFilterToBinary(vcf.filter, params.bit.len)
```

Arguments

vcf.filter	A list generated in MakeFilter
params.bit.len	A list with bit lengths of filtering parameters which is generated from ParameterToBits

Value

A binary vector

EnumerateTriNucCounts *EnumerateTriNucCounts*

Description

Returns C>A, C>G, C>T, T>A, T>C, T>G counts

Usage

EnumerateTriNucCounts(spectrum)

Arguments

spectrum a numeric vector with 96 numeric values

Details

Please note that this function assumes that 'spectrum' is sorted (i.e. 1:16 → C>A; 17:32 → C>G; 33:48 → C>T; 49:64 → T>A; 65:80 → T>C; 81:96 → T>G)

Value

a numeric vector of length 6 corresponding to the counts of each trinucleotide change (C>A, C>G, C>T, T>A, T>C, T>G)

Euc.Exp.Weighted.Obj.Fn
Euc.Exp.Weighted.Obj.Fn

Description

Calculates the Euclidean-distance of logarithmically weighted objective value for FIREVAT GA optimization.

Usage

Euc.Exp.Weighted.Obj.Fn(C.refined, A.refined, C.artifactual, A.artifactual)

Arguments

C.refined A numeric value between 0 and 1.
A.refined A numeric value between 0 and 1.
C.artifactual A numeric value between 0 and 1.
A.artifactual A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.1

Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.1

Description

Calculates the Euclidean-distance of logarithmically weighted objective value for FIREVAT GA optimization.

Usage

Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.1(C.refined, A.refined, C.artifactual, A.artifactual)

Arguments

C.refined	A numeric value between 0 and 1.
A.refined	A numeric value between 0 and 1.
C.artifactual	A numeric value between 0 and 1.
A.artifactual	A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.2

Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.2

Description

Calculates the Euclidean-distance of logarithmically weighted objective value for FIREVAT GA optimization.

Usage

Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.2(C.refined, A.refined, C.artifactual, A.artifactual)

Arguments

C.refined	A numeric value between 0 and 1.
A.refined	A numeric value between 0 and 1.
C.artifactual	A numeric value between 0 and 1.
A.artifactual	A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

Euc.Obj.Fn	<i>Euc.Obj.Fn</i>
------------	-------------------

Description

Calculates the Euclidean-distance based objective value for FIREVAT GA optimization.

Usage

Euc.Obj.Fn(C.refined, A.refined, C.artifactual, A.artifactual)

Arguments

- C.refined A numeric value between 0 and 1.
- A.refined A numeric value between 0 and 1.
- C.artifactual A numeric value between 0 and 1.
- A.artifactual A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

Exp.Weighted.Obj.Fn.1	<i>Exp.Weighted.Obj.Fn.1</i>
-----------------------	------------------------------

Description

Calculates the exponentially weighted objective value for FIREVAT GA optimization.

Usage

Exp.Weighted.Obj.Fn.1(C.refined, A.refined, C.artifactual, A.artifactual)

Arguments

- C.refined A numeric value between 0 and 1.
- A.refined A numeric value between 0 and 1.
- C.artifactual A numeric value between 0 and 1.
- A.artifactual A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

Exp.Weighted.Obj.Fn.2 *Exp.Weighted.Obj.Fn.2*

Description

Calculates the exponentially weighted objective value for FIREVAT GA optimization.

Usage

Exp.Weighted.Obj.Fn.2(C.refined, A.refined, C.artifactual, A.artifactual)

Arguments

C.refined	A numeric value between 0 and 1.
A.refined	A numeric value between 0 and 1.
C.artifactual	A numeric value between 0 and 1.
A.artifactual	A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

Exp.Weighted.Refined.Seq.Art.Only.Obj.Fn
Exp.Weighted.Refined.Seq.Art.Only.Obj.Fn

Description

Calculates the Euclidean-distance of logarithmically weighted objective value for FIREVAT GA optimization.

Usage

Exp.Weighted.Refined.Seq.Art.Only.Obj.Fn(C.refined, A.refined,
C.artifactual, A.artifactual)

Arguments

C.refined	A numeric value between 0 and 1.
A.refined	A numeric value between 0 and 1.
C.artifactual	A numeric value between 0 and 1.
A.artifactual	A numeric value between 0 and 1.

Value

A numeric value between 0 and 1.

FilterVCF

*FilterVCF***Description**

Filter vcf based on the filter Filtering parameters are saved in config.obj Split vcf.obj into vcf.obj.filtered & vcf.obj.artifact based on vcf.filter

Usage

```
FilterVCF(vcf.obj, vcf.filter, config.obj, include.array = NULL,
         force.include = FALSE, verbose = TRUE)
```

Arguments

vcf.obj	A list from ReadVCF
vcf.filter	A list from MakeMuTect2Filter
config.obj	A list from ParseConfigFile
include.array	A boolean vector
force.include	A boolean value. If TRUE, then uses 'include.array'
verbose	If true, provides process detail

Value

A list with the following elements

- 1) Mutations which passed filteringvcf.obj.filtered = vcf.obj (list with data, header, genome)
- 2) Mutations which did not pass filteringvcf.obj.artifact = vcf.obj (list with data, header, genome)

GenerateConfigObj

*Generate config.obj by checking vcf header***Description**

This function generate config.obj by checking vcf header. Users should fill in the information needed in console. In current version, only Integers & Float values can be used in config.obj for running FIREVAT.

Usage

```
GenerateConfigObj(vcf.obj, save.config = TRUE,
                 config.path = "../temp/FIREVAT_configure.json")
```

Arguments

- vcf.obj A list from [ReadVCF](#)
- save.config If true, save config.obj to config.path
- config.path File path to write config.obj (json or yaml)

Value

config.obj

GetCOSMICMutSigs	<i>GetCOSMICMutSigs</i>
------------------	-------------------------

Description

Returns a data.frame of the COSMIC mutational signature reference file from the data directory

Usage

GetCOSMICMutSigs()

Value

a data.frame of the COSMIC reference mutational signatures

GetCOSMICMutSigsEtiologiesColors
<i>GetCOSMICMutSigsNames</i>

Description

Returns all COSMIC mutational signature etiologies and colors

Usage

GetCOSMICMutSigsEtiologiesColors()

Value

data.frame with following columns: signature, group and color.

GetCOSMICMutSigsNames *GetCOSMICMutSigsNames*

Description

Returns all COSMIC mutational signature names

Usage

GetCOSMICMutSigsNames()

Value

a character vector

GetGASuggestedSolutions
 GetGASuggestedSolutions

Description

Computes suggested solutions

Usage

```
GetGASuggestedSolutions(vcf.obj, bsg, config.obj, lower.upper.list,
  df.mut.pat.ref.sigs, target.mut.sigs, sequencing.artifact.mut.sigs,
  objective.fn, original.muts.seq.art.weights.sum, verbose = TRUE)
```

Arguments

vcf.obj	A list from ReadVCF
bsg	BSgenome.Hsapiens.UCSC object
config.obj	A list from ParseConfigFile
lower.upper.list	A list from GetParameterLowerUpperVector
df.mut.pat.ref.sigs	A data.frame from MutPatParseRefMutSigs
target.mut.sigs	A character vector of the target mutational signatures from reference mutational signatures.
sequencing.artifact.mut.sigs	A character vector of the sequencing artifact mutational signatures from reference mutational signatures.

objective.fn Objective value derivation function.

original.muts.seq.art.weights.sum
 A numeric value. 'seq.art.sigs.weights.sum' from CheckIfVariantRefinementIs-Necessary

verbose If TRUE, provides process detail. Default value is TRUE.

Value

A list with the following elements

- judgmentA boolean value
- seq.art.sigs.weightsA numeric value. Sum of sequencing artifact weights.

GetOptimizedSignatures

GetOptimizedSignatures

Description

This function fetches the last row from the optimization iteration log and returns the target and artifactual mutational signatures for the type of mutations ('refined' or 'artifactual')

Usage

```
GetOptimizedSignatures(data, mutations.type = "refined",
  signatures = "all")
```

Arguments

data A list of main data from [RunFIREVAT](#)

mutations.type A string for type of mutations ('refined' or 'artifact')

signatures A string ('all', 'target', 'artifact')

Value

A data.frame with the columns 'signature' and 'weight'

GetParameterLowerUpperVector

GetParameterLowerUpperVector

Description

Return a lower/upper vector needed to conduct FIREVAT GA real-valued optimization.

Usage

```
GetParameterLowerUpperVector(vcf.obj, config.obj, vcf.filter,
                             multiplier = 100)
```

Arguments

vcf.obj	A list from ReadVCF
config.obj	A list from ParseConfigFile
vcf.filter	A list from MakeMuTect2Filter
multiplier	A multiplier for convert fraction to integer (default = 100)

Details

vcf.obj\$data: if $\max(\text{vcf.obj\$data}[[\text{param}]]) < 1$, then multiply multiplier to the vector

Value

A list with the elements

- lower.vector A numeric vector. Each element is the minimum value of each parameter
- upper.vector A numeric vector. Each element is the maximum value of each parameter
- vcf.obj vcf.obj with updated data

GetPCAWGMutSigs

GetPCAWGMutSigs

Description

Returns the PCAWG mutational signatures data

Usage

```
GetPCAWGMutSigs()
```

Value

a data.frame of the PCAWG mutatioanl signatures

GetPCAWGMutSigsEtiologiesColors
<i>GetPCAWGMutSigsEtiologiesColors</i>

Description

Returns the PCAWG mutational signatures etiologies and colors

Usage

GetPCAWGMutSigsEtiologiesColors()

Value

a data.frame with the columns 'signature', 'group', 'color'

GetPCAWGMutSigsNames	<i>GetPCAWGMutSigsNames</i>
----------------------	-----------------------------

Description

Returns the PCAWG mutational signatures names

Usage

GetPCAWGMutSigsNames()

Value

a character vector of the PCAWG mutational signatures names

InitializeVCF	<i>InitializeVCF</i>
---------------	----------------------

Description

Initialize VCF with FIREVAT config file This functions selects point mutations and appends filter values to vcf.obj\$data

Usage

InitializeVCF(vcf.obj, config.obj, verbose = TRUE)

Arguments

- vcf.obj A list from ReadVCF
- config.obj A list from ParseConfigFile
- verbose If true, provides process detail

Value

- A list with the following elements
- vcf.obj.filteredvcf.obj (high-quality vcf)
 - vcf.obj.artifactvcf.obj (low-quality vcf)

MakeFilter	<i>MakeFilter</i>
------------	-------------------

Description

Creates a vcf filter from config.obj

Usage

MakeFilter(config.obj)

Arguments

- config.obj A list from ParseConfigFile (any filter with "use_in_filter" value declared as FALSE is not considered)

Value

A list with the filter parameters

MutaliskParseVCFObj	<i>MutaliskParseVCFObj</i>
---------------------	----------------------------

Description

Parses a vcf.obj and prepares it to run Mutalisk.

Usage

MutaliskParseVCFObj(vcf.obj)

Arguments

- vcf.obj A list from ReadVCF

Value

A data.frame

MutPatParseRefMutSigs *MutPatParseRefMutSigs*

Description

Parses a df.ref.mut.sigs and prepares it to run Mutational Patterns.

Usage

```
MutPatParseRefMutSigs(df.ref.mut.sigs, target.mut.sigs,
  signature.start.column.index = 4,
  mutation.type.header = "SomaticMutationType")
```

Arguments

`df.ref.mut.sigs`
A data.frame of reference mutational signatures

`target.mut.sigs`
A character vector of target mutational signatures names

`signature.start.column.index`
= An integer value (e.g. column index corresponding to 'SBS1')

`mutation.type.header`
= A string value (name of header corresponding to column containing 'A[C>A]A' data))

Value

A data.frame of the format `deconstructSigs::signatures.cosmic`

MutPatParseVCFObj *MutPatParseVCFObj*

Description

Parses a vcf.obj and prepares it to run Mutational Patterns.

Usage

```
MutPatParseVCFObj(vcf.obj, bsg, sample.id = "sample")
```

Arguments

vcf.obj	A list from ReadVCF
bsg	BSgenome.Hsapiens.UCSC.hg19::BSgenome.Hsapiens.UCSC.hg19 or BSgenome.Hsapiens.UCSC.hg38
sample.id	A string value

Value

A data.frame with the column sample.id and row names corresponding to 96 substitution types

ParameterToBits	<i>ParameterToBits</i>
-----------------	------------------------

Description

Calculate the number of bits needed to conduct FIREVAT GA binary optimization.

Usage

ParameterToBits(vcf.obj, config.obj, vcf.filter, multiplier = 100)

Arguments

vcf.obj	A list from ReadVCF
config.obj	A list from ParseConfigFile
vcf.filter	A list from MakeMuTect2Filter
multiplier	A multiplier for convert fraction to integer (default = 100)

Details

vcf.obj\$data: if $\max(\text{vcf.obj\$data}[[\text{param}]]) < 1$, then multiply multiplier to the vector

Value

A list with the elements

- params.bit.lenA numeric vector. Each element is the bit length of each parameter value
- vcf.objA vcf.obj ([ReadVCF](#)) with updated data

ParseConfigFile	<i>ParseConfigFile</i>
-----------------	------------------------

Description

This function returns config.obj from JSON or YAML config file. - Check if the config file is in JSON format or YAML format - Return config.obj

Usage

ParseConfigFile(config.path, verbose = TRUE)

Arguments

config.path	A string for config file path
verbose	If true, provides process detail

Value

config.obj: list of parameters

Examples

```
## Not run:
ParseConfigFile("example.variant.caller.json")
ParseConfigFile("example.variant.caller.json", verbose=False)

## End(Not run)
```

PCAWG.All.Sequencing.Artifact.Signatures
<i>Constant</i>

Description

PCAWG mutational signatures reported to be associated with sequencing artifacts

Usage

PCAWG.All.Sequencing.Artifact.Signatures

Format

An object of class character of length 17.

PCAWG.Known.Sequencing.Artifact.Signatures
Constant

Description

PCAWG mutational signatures reported to be associated with sequencing artifacts

Usage

PCAWG.Known.Sequencing.Artifact.Signatures

Format

An object of class character of length 1.

PCAWG.Likely.Sequencing.Artifact.Signatures
Constant

Description

PCAWG mutational signatures reported to be associated with sequencing artifacts

Usage

PCAWG.Likely.Sequencing.Artifact.Signatures

Format

An object of class character of length 5.

PCAWG.Possible.Sequencing.Artifact.Signatures
Constant

Description

PCAWG mutational signatures reported to be associated with sequencing artifacts

Usage

PCAWG.Possible.Sequencing.Artifact.Signatures

Format

An object of class character of length 11.

PCAWG.Target.Mutational.Signatures
Constant

Description

PCAWG target mutational signatures reported to be unrelated to sequencing artifacts

Usage

PCAWG.Target.Mutational.Signatures

Format

An object of class character of length 49.

PerformStrandBiasAnalysis
PerformStrandBiasAnalysis

Description

Performs strand bias analysis

Usage

```
PerformStrandBiasAnalysis(vcf.obj, ref.forward.strand.var,  
  ref.reverse.strand.var, alt.forward.strand.var, alt.reverse.strand.var,  
  perform.fdr.correction = TRUE, fdr.correction.method = "BH")
```

Arguments

vcf.obj	ReadVCF
ref.forward.strand.var	A string value.
ref.reverse.strand.var	A string value.
alt.forward.strand.var	A string value.
alt.reverse.strand.var	A string value.
perform.fdr.correction	If TRUE, then performs false discovery rate correction
fdr.correction.method	A string value. FDR correction method (Refer to p.adjust() function)

Value

An updated vcf.obj

PlotMutaIiskResults	<i>PlotMutaIiskResults</i>
---------------------	----------------------------

Description

Plots MutaIisk results

Usage

```
PlotMutaIiskResults(mutaIisk.results, signatures, trinuc.max.y,
  trinuc.min.y, mut.type.max.y, title)
```

Arguments

mutaIisk.results	A list obtained from RunMutaIisk
signatures	A character vector of mutational signatures names
trinuc.max.y	A numeric value (maximum y-axis value)
trinuc.min.y	A numeric value (minimum y-axis value)
mut.type.max.y	A numeric value
title	A string value

Value

A ggplot object

Examples

```
## Not run:
df.ref.mut.sigs <- GetPCAWGMutSigs()
target.mut.sigs <- GetPCAWGMutSigsNames()
vcf.obj <- ReadVCF(vcf.file = "../data/sample/P-233-CT.final.vcf")
mutaIisk.results <- RunMutaIisk(vcf.obj = vcf.obj,
                              df.ref.mut.sigs = df.ref.mut.sigs,
                              target.mut.sigs = target.mut.sigs)
p <- PlotMutaIiskResults(mutaIisk.results = mutaIisk.results)
print(p)

## End(Not run)
```

PlotMutationTypes	<i>PlotMutationTypes</i>
-------------------	--------------------------

Description

Plots a horizontal barplot of mutation types

Usage

```
PlotMutationTypes(mutation.types = c("C>A", "C>G", "C>T", "T>A", "T>C",
  "T>G"), mutation.types.values, mutation.types.colors, max.y.val, title,
  convert.to.percentage = T, show.legend = T, font.size.small = 8,
  font.size.med = 14, plot.margin = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))
```

Arguments

mutation.types	Mutation types; Default = c("C>A", "C>G", "C>T", "T>A", "T>C", "T>G")
mutation.types.values	Mutation count for each mutation type
mutation.types.colors	A color vector for indicating mutation types
max.y.val	y axis maximum value
title	Plot title
convert.to.percentage	if True convert y values to percentage (x 100); Default = T
show.legend	If True, show legend; Default = T
font.size.small	Small font size; Default = 8
font.size.med	Medium font size; Default = 14
plot.margin	Margin vector for drawing plot; Default = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))

Value

A ggplot object

Examples

```
## Not run:
p <- PlotMutationTypes(mutation.types = c("C>A", "C>G", "C>T", "T>A", "T>C", "T>G"),
  mutation.types.values = c(0.3, 0.3, 0.1, 0.1, 0.1, 0.1),
  mutation.types.colors = TriNuc.Mutation.Type.Hex.Colors,
  max.y.val = 0.5,
  convert.to.percentage = T,
  show.legend = T,
  font.size.small = 8,
  font.size.med = 14,
```

```

plot.margin = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))
print(p)

## End(Not run)

```

PlotOptimizationIterations

PlotOptimizationIterations

Description

Plots multiple scatter plots into one figure

Usage

```

PlotOptimizationIterations(df, columns.to.plot, x.axis.var, x.axis.title,
  x.max, save.file, title, y.axis.title = "", y.max = 1,
  point.size = 1, connect.dots = T, plot.legend = T,
  legend.ncol = 1, font.size.med = 14, font.size.large = 16,
  plot.margin = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))

```

Arguments

df	A data.frame (from reading "FIREVAT_Optimization_Logs.tsv")
columns.to.plot	A character vector (of column names to plot)
x.axis.var	x axis variable
x.axis.title	x axis title
x.max	x axis maximum value
save.file	Filename (including full path) to which the plot will be saved
title	Plot title
y.axis.title	y axis title; Default = ""
y.max	y axis maximum value; Default = 1
point.size	Point size; Default = 1
connect.dots	If True draws dots for each iteration; Default = True
plot.legend	If True write legend of plot; Default = T
legend.ncol	legend.n Default = 1
font.size.med	Medium font size; Default = 14
font.size.large	Large font size; Default = 16
plot.margin	Margin vector for plot; Default = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))

Value

A ggplot object

`PlotSignaturesContProbs`*PlotSignaturesContProbs*

Description

Plots a horizontal barplot of identified mutational signatures

Usage

```
PlotSignaturesContProbs(df.identified.mut.sigs, df.ref.sigs.groups.colors,  
  title, convert.to.percentage = T, font.size.small = 8,  
  font.size.med = 14, plot.margin = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))
```

Arguments

<code>df.identified.mut.sigs</code>	A data.frame of identified mutational signatures
<code>df.ref.sigs.groups.colors</code>	A data.frame with 'signature', 'group', and 'color' columns
<code>title</code>	Plot title
<code>convert.to.percentage</code>	If true, convert y values to percentage (x 100); Default = T,
<code>font.size.small</code>	Small font size; Default = 8,
<code>font.size.med</code>	Medium font size; Default = 14,
<code>plot.margin</code>	Margin vector for drawing plot; Default = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))

Value

A ggplot object

Examples

```
## Not run:  
g <- PlotSignaturesContProbs(sigs = c(mutalisk.results$identified.mut.sigs),  
  sigs.probs = c(mutalisk.results$identified.mut.sigs.probs),  
  df.ref.sigs.groups.colors = GetPCAWGMutSigsEtiologiesColors())  
print(g)  
  
## End(Not run)
```

PlotTable	<i>PlotTable</i>
-----------	------------------

Description

Plots basic statistics table

Usage

```
PlotTable(df, padding = 20, font.size = 14)
```

Arguments

- df = A data.frame where the first column is header and the second column is data value
- padding Padding size; Default = 20
- font.size Font size; Default = 14

Value

A plot

PlotTriNucSpectrum	<i>PlotTriNucSpectrum</i>
--------------------	---------------------------

Description

Plots the spectrum of 96 trinucleotide distribution (C>A, C>G, C>T, T>A, T>C, T>G) Please note that this function assumes that both sub.types and spectrum are sorted in the following order: C>A, C>G, C>T, T>A, T>C, T>G

Usage

```
PlotTriNucSpectrum(sub.types, spectrum, max.y.val, min.y.val, y.axis.title,
  draw.top.strip = T, draw.x.axis.labels = T, draw.y.axis.labels = T,
  draw.y.axis.title = T, font.size.small = 8, font.size.med = 14,
  plot.margin.top = 0.5, plot.margin.bottom = 0.5,
  plot.margin.left = 0.5, plot.margin.right = 0.5, title)
```

Arguments

sub.types	A character vector (types of 96 trinucleotide substitutions)
spectrum	A numeric vector (96 elements)
max.y.val	y axis maximum value
min.y.val	y axis minimum value
y.axis.title	y axis title
draw.top.strip	If True then draws top strip; Default = T
draw.x.axis.labels	If True then draws x axis labels; Default = T
draw.y.axis.labels	If True then draws y axis labels; Default = T
draw.y.axis.title	If True then draws y axis title; Default = T
font.size.small	Small font size; Default = 8
font.size.med	Medium font size; Default = 14
plot.margin.top	Top margin; Default = 0.5
plot.margin.bottom	Bottom margin; Default = 0.5
plot.margin.left	Left margin; Default = 0.5
plot.margin.right	Right margin; Default = 0.5
title	Plot title

Value

A ggplot object

PlotVCFStatsBoxPlots *PlotVCFStatsBoxPlots*

Description

Plots multiple (original, refined, artifact vcf) boxplots for single filter parameter

Usage

```
PlotVCFStatsBoxPlots(original.vcf.stat.values, refined.vcf.stat.values,
  artifact.vcf.stat.values, xlab, axis.font.size = 10,
  label.font.size = 10, title.font.size = 12)
```

Arguments

<code>original.vcf.stat.values</code>	A numeric vector corresponding to the original vcf.obj values of single filter parameter
<code>refined.vcf.stat.values</code>	A numeric vector corresponding to the refined vcf.obj values of single filter parameter
<code>artifact.vcf.stat.values</code>	A numeric vector corresponding to the artifact vcf.obj values of single filter parameter
<code>xlab</code>	A string value (x-axis label)
<code>axis.font.size</code>	An integer value (axis font size)
<code>label.font.size</code>	An integer value (label font size)
<code>title.font.size</code>	An integer value (title font size)

Value

A ggboxplot

PlotVCFStatsHistograms

PlotVCFStatsHistograms

Description

Plots multiple VCF stats histograms into one figure

Usage

```
PlotVCFStatsHistograms(plot.values, x.axis.labels, stat.y.max.vals,
  stat.x.max.vals, sample.id, save.file, title, cutoff.values,
  plot.boxplot = F, plot.cutoff.line.color = "#D4012E",
  plot.cutoff.value.lines = F, bin.width = 1, ncol = 4, nrow = 3,
  font.size.med = 10, font.size.large = 12, plot.margin = unit(c(0.5,
  0.5, 0.5, 0.5), "cm"))
```

Arguments

<code>plot.values</code>	A list of multiple numeric vectors
<code>x.axis.labels</code>	A character vector of x axis labels
<code>stat.y.max.vals</code>	A numeric vector of max y-axis values

stat.x.max.vals	A numeric vector of max x-axis values
sample.id	A string value of sample ID
save.file	A string value of file to which the resulting plot will be saved
title	A string value of plot title
cutoff.values	A numeric vector of cutoff values
plot.boxplot	A boolean value (default = False)
plot.cutoff.line.color	A hex string value (default = "#D4012E")
plot.cutoff.value.lines	A boolean value (default = False)
bin.width	An integer value (default = 1; histogram bin width)
ncol	An integer value (default = 4; ggarrange ncol)
nrow	An integer value (default = 3; ggarrange nrow)
font.size.med	An integer value (default = 10)
font.size.large	An integer value (default = 12)
plot.margin	A list (default = unit(c(0.5, 0.5, 0.5, 0.5), "cm"))

Value

A list with the following elements

- f = A ggarrange object
- graphs = A list of length 3; each element is a ggplot histogram

PrepareAnnotationDB	<i>PrepareAnnotationDB</i>
---------------------	----------------------------

Description

Prepares df.genes.of.interest from a vcf.obj ([ReadVCF](#)) of COSMIC or ClinVar vcf for [AnnotateVCFObj](#)

Usage

```
PrepareAnnotationDB(annotation.vcf.obj)
```

Arguments

annotation.vcf.obj
vcf.obj of COSMIC or ClinVar vcf file

Value

A data.frame of annotation.vcf.obj

PrepareArtifactAnnotationTable
<i>PrepareArtifactAnnotationTable</i>

Description

Prepares artifactual mutations annotation (filtered, queried) table

Usage

PrepareArtifactAnnotationTable(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

PrepareArtifactStrandBiasTable
<i>PrepareArtifactStrandBiasTable</i>

Description

Prepares artifactual mutations strand biased variants table

Usage

PrepareArtifactStrandBiasTable(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

`PrepareArtifactualMutsOptimizationIterationsPlot`*PrepareArtifactualMutsOptimizationIterationsPlot*

Description

Prepares artifactual mutations optimization iterations plot

Usage

```
PrepareArtifactualMutsOptimizationIterationsPlot(data)
```

Arguments

`data` A list of elements returned from [RunFIREVAT](#)

Value

A ggplot object

`PrepareFilterCutoffsTable`*PrepareFilterCutoffsTable*

Description

Prepares filter cutoffs table for reporting

Usage

```
PrepareFilterCutoffsTable(data)
```

Arguments

`data` A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

PrepareGeneticAlgorithmParametersTable
<i>PrepareGeneticAlgorithmParametersTable</i>

Description

Prepares Genetic Algorithm parameters table

Usage

PrepareGeneticAlgorithmParametersTable(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

PrepareIdentifiedSignaturesPlot
<i>PrepareIdentifiedSignaturesPlot</i>

Description

Prepares identified signatures plot for reporting

Usage

PrepareIdentifiedSignaturesPlot(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A ggarrange object

`PrepareMLEReconstructedSpectrumsPlot`*PrepareMLEReconstructedSpectrumsPlot*

Description

Prepares MLE reconstructed spectrums plot

Usage

```
PrepareMLEReconstructedSpectrumsPlot(data)
```

Arguments

`data` A list of elements returned from [RunFIREVAT](#)

Value

A ggarrange object

`PrepareNucleotideSubstitutionTypesPlot`*PrepareNucleotideSubstitutionTypesPlot*

Description

Prepares nucleotide substitution types plot

Usage

```
PrepareNucleotideSubstitutionTypesPlot(data)
```

Arguments

`data` A list of elements returned from [RunFIREVAT](#)

Value

A ggarrange object

PrepareObservedSpectrumsPlot
<i>PrepareObservedSpectrumsPlot</i>

Description

Prepares observed spectrums plot

Usage

PrepareObservedSpectrumsPlot(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A ggarrange object

PrepareOptimizationResultsTable
<i>PrepareOptimizationResultsTable</i>

Description

Prepares optimization results table

Usage

PrepareOptimizationResultsTable(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

PrepareOptimizedVCFStatisticsPlot
<i>PrepareOptimizedVCFStatisticsPlot</i>

Description

Prepares optimized VCF statistics plot

Usage

PrepareOptimizedVCFStatisticsPlot(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A ggarrange object

PrepareRefinedAnnotationTable
<i>PrepareRefinedAnnotationTable</i>

Description

Prepares refined mutations annotation (filtered, queried) table

Usage

PrepareRefinedAnnotationTable(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

PrepareRefinedMutsOptimizationIterationsPlot
PrepareRefinedMutsOptimizationIterationsPlot

Description

Prepares refined mutations optimization iterations plot

Usage

PrepareRefinedMutsOptimizationIterationsPlot(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A ggplot object

PrepareRefinedStrandBiasTable
PrepareRefinedStrandBiasTable

Description

Prepares refined mutations strand biased variants table

Usage

PrepareRefinedStrandBiasTable(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

PrepareResidualSpectrumsPlot
<i>PrepareResidualSpectrumsPlot</i>

Description

Prepares residual spectrums plot

Usage

PrepareResidualSpectrumsPlot(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A ggarrange object

PrepareTrinucleotideSpectrumsTable
<i>PrepareTrinucleotideSpectrumsTable</i>

Description

Prepares trinucleotide spectrums table

Usage

PrepareTrinucleotideSpectrumsTable(data)

Arguments

data A list of elements returned from [RunFIREVAT](#)

Value

A data.frame

QueryAnnotatedVCF	<i>FilterAnnotatedVCF</i>
-------------------	---------------------------

Description

Annotates a `vcf.obj` using `df.variants.of.interest` (from ([PrepareAnnotationDB](#)))

Usage

```
QueryAnnotatedVCF(vcf.obj.annotated, filter.key.value.pairs,
  filter.condition = "AND")
```

Arguments

- `vcf.obj.annotated`
[AnnotateVCFObj](#)
- `filter.key.value.pairs`
A list with the key as the column name and value as the filtering values. E.g.
`list("CLNSIG" = c("Pathogenic", "Pathogenic/Likely_pathogenic"))`
- `filter.condition`
'AND' or 'OR'.

Value

A `vcf.obj`

ReadOptimizationIterationReport	<i>ReadOptimizationIterationReport</i>
---------------------------------	--

Description

Read optimization iteration report

Usage

```
ReadOptimizationIterationReport(data)
```

Arguments

- `data` A list of elements returned from [RunFIREVAT](#)

Value

A `data.frame` of FIREVAT optimization logs

ReadVCF	<i>ReadVCF</i>
---------	----------------

Description

Reads a .vcf file

Usage

```
ReadVCF(vcf.file, genome = "hg19", split.info = FALSE,
        check.chromosome.name = TRUE)
```

Arguments

- vcf.file (full path of a .vcf file)
- genome ('hg19' or 'hg38')
- split.info A boolean value. If TRUE, then makes the INFO column in the vcf as a separate column. Default value is FALSE.
- check.chromosome.name A boolean value. If TRUE, then check whether converts 'MT' to 'M' and adds 'chr' to the CHROM column. Default value is TRUE.

Value

A list with elements 'data', 'header', 'genome'

ReportFIREVATResults	<i>ReportFIREVATResults</i>
----------------------	-----------------------------

Description

Reports FIREVAT results in html format (generated from Rmd)

Usage

```
ReportFIREVATResults(data)
```

Arguments

- data A list of main data from [RunFIREVAT](#)

Value

An updated data list

RunFIREVAT

*RunFIREVAT***Description**

Runs FIREVAT using configuration data. Filters point mutations in the user-specified vcf file based on mutational signature identification and outputs the refined and artifact vcf files as well as meta-data related to the refinement process.

Usage

```
RunFIREVAT(vcf.file, vcf.file.genome, config.file, df.ref.mut.sigs,
  target.mut.sigs, sequencing.artifact.mut.sigs, num.cores, output.dir,
  mode = "ga", init.artifact.stop = 0.05,
  objective.fn = Default.Obj.Fn, use.suggested.soln = TRUE,
  ga.type = "real-valued", ga.pop.size = 200, ga.max.iter = 200,
  ga.run = 50, ga.pmutation = 0.25, mutalisk = TRUE,
  mutalisk.method = "all", mutalisk.random.sampling.count = 20,
  mutalisk.random.sampling.max.iter = 10,
  perform.strand.bias.analysis = TRUE,
  strand.bias.perform.fdr.correction = TRUE,
  strand.bias.fdr.correction.method = "BH",
  ref.forward.strand.var = NULL, ref.reverse.strand.var = NULL,
  alt.forward.strand.var = NULL, alt.reverse.strand.var = NULL,
  annotate = TRUE, df.annotation.db = NULL,
  annotated.columns.to.display = NULL,
  annotation.filter.key.value.pairs = NULL,
  annotation.filter.condition = "AND", write.vcf = TRUE,
  report = TRUE, save.rdata = TRUE, save.tsv = TRUE,
  report.format = "html", verbose = TRUE)
```

Arguments

vcf.file	String value corresponding to input .vcf file. Please provide the full path.
vcf.file.genome	Genome assembly of the input .vcf file. The value should be either 'hg19' or 'hg38'.
config.file	String value corresponding to input configuration file. For more details please refer to ...
df.ref.mut.sigs	A data.frame of the reference mutational signatures
target.mut.sigs	A character vector of the target mutational signatures from reference mutational signatures.
sequencing.artifact.mut.sigs	A character vector of the sequencing artifact mutational signatures from reference mutational signatures.

num.cores	Number of cores to allocate
output.dir	String value of the desired output directory
mode	String value. The value should be either 'ga' or 'manual'.
init.artifact.stop	Numeric value less than 1. If the sum of sequencing artifact weights in the user-specified original VCF file (i.e. vcf.file) is less than or equal to this value then FIREVAT does not perform variant refinement. Default value is 0.05. Note that this option does not apply if 'mode' is 'manual'.
objective.fn	Objective value derivation function. Default: Default.Obj.Fn.
use.suggested.soln	Boolean value. If TRUE, then FIREVAT passes the default values of filter variables declared as 'use_in_filter' in the config file to the 'suggestions' parameter of the Genetic Algorithm package. If FALSE, then FIREVAT supplies NULL to the GA package 'suggestions' parameter. FIREVAT also computes baseline performance of each filter variable and uses fittest population from each variable as a suggested solution.
ga.type	String value. The value should be either 'binray' or 'real-valued'.
ga.pop.size	Integer value of the Genetic Algorithm 'population size' parameter. Default: 200. This value should be set based on the number of filter parameters. Recommendation: 40 per filter parameter.
ga.max.iter	Integer value of the Genetic Algorithm 'maximum iterations' parameter. Default: 200. This value should be set based on the number of filter parameters. Recommendation: same as 'ga.pop.size'.
ga.run	Integer value of the Genetic Algorithm 'run' parameter. Default: 50. This value should be set based on the 'ga.max.iter' parameter. Recommendation: 25 percent of 'ga.max.iter'.
ga.pmutation	Float value of the Genetic Algorithm 'mutation probability' parameter. Default: 0.25.
mutalisk	If TRUE, confirm mutational signature analysis with Mutalisk. Default: TRUE.
mutalisk.method	Mutalisk signature identification method. Default: 'random.sampling'. The value can be either 'all' or 'random.sampling'. 'all' uses all target.mut.sigs to identify mutational signatures. 'random.sampling' randomly samples from target.mut.sigs to identify mutational signatures.
mutalisk.random.sampling.count	Mutalisk random sampling count. Default: 20. The number of signatures to sample from target.mut.sigs
mutalisk.random.sampling.max.iter	Mutalisk random sampling maximum iteration. Default: 10. The number of times Mutalisk randomly samples from target.mut.sigs before determining the candidate signatures.
perform.strand.bias.analysis	If TRUE, then performs strand bias analysis.
strand.bias.perform.fdr.correction	If TRUE, then performs false discovery rate correction for strand bias analysis.

strand.bias.fdr.correction.method
A string value. Default value is 'BH'. Refer to 'p.adjust()' function method.

ref.forward.strand.var
A string value.

ref.reverse.strand.var
A string value,

alt.forward.strand.var
A string value,

alt.reverse.strand.var
A string value,

annotate
A boolean value. Default value is TRUE.

df.annotation.db
A data.frame. Please refer to [PrepareAnnotationDB](#)

annotated.columns.to.display
A character vector.

annotation.filter.key.value.pairs
A list.

annotation.filter.condition
'AND' or 'OR'.

write.vcf
If TRUE, write original/refined/artifact vcfs. Default: TRUE.

report
If TRUE, generate report. Default: TRUE.

save.rdata
If TRUE, save rdata. Default: TRUE.

save.tsv
If TRUE, save tsv. Default: TRUE.

report.format
The format of FIREVAT report. We currently only support 'html'.

verbose
If TRUE, provides process detail. Default: TRUE.

Value

A list with the following elements

- f = A ggarrange object
- graphs = A list of length 3; each element is a ggplot histogram

RunMutalisk

RunMutalisk

Description

Identifies mutational signatures using Mutalisk

Usage

```
RunMutalisk(vcf.obj, df.ref.mut.sigs, target.mut.sigs,
  random.sampling.candidate.mut.sigs = c(), method = "random.sampling",
  n.sample = 20, n.iter = 10, verbose = TRUE)
```

Arguments

<code>vcf.obj</code>	A list (from <code>firevat_vcf::ReadVCF</code>)
<code>df.ref.mut.sigs</code>	A data.frame of reference mutational signatures
<code>target.mut.sigs</code>	A character vector of target mutational signatures names to identify from
<code>random.sampling.candidate.mut.sigs</code>	A character vector of mutational signatures names that gets appended to the list of candidate mutational signatures so that these are always considered.
<code>method</code>	A string value (must be either 'random.sampling' or 'all'). The method 'random.sampling' samples (without replacement) 'n.sample' number of signatures 'n.iter' number of times and runs the candidate signatures one last time. The method 'all' uses all target.mut.sigs
<code>n.sample</code>	An integer value ('random.sampling' method parameter) Number of signatures to choose for each iteration of random sampling).
<code>n.iter</code>	An integer value ('random.sampling' method parameter). Number of iterations to perform random sampling.
<code>verbose</code>	If true, provides process details

Value

A list with the following elements

- `num.point.mutations` An integer value - count of total point mutations
- `sub.types` A character vector of length 96
- `sub.types.spectrum` A numeric vector of length 96
- `num.mut.sigs` An integer value (count of unique mutational signatures identified)
- `identified.mut.sigs` A character vector where each element is a mutational signature identified
- `identified.mut.sigs.probs` A numeric vector where each element is the weight of mutational signature identified. The ordering follows `identified.mut.sigs`
- `identified.mut.sigs.spectrum` A numeric vector of length 96
- `residuals` A numeric vector of length 96
- `rss` A numeric value (residual sum of squares)
- `cos.sim.score` A numeric value (cosine similarity score between observed mutational spectrum and reconstructed mutational signatures)
- `all.models.sigs` A list where each element is a model; a model is a list of signatures identified
- `all.models.sigs.probs` A list where each element is a model; a model is a list of contribution probabilities
- `all.models.cos.sim.scores` A list where each element is a model; a model is a list of cosine similarity scores

RunMutaliskHelper	<i>RunMutaliskHelper</i>
-------------------	--------------------------

Description

Helper function for RunMutalisk

Usage

```
RunMutaliskHelper(vcf.trinucleotide.data, df.ref.mut.sigs, target.mut.sigs)
```

Arguments

`vcf.trinucleotide.data`
 A data.frame (from firevat_mutalisk::MutaliskParseVCFObj)

`df.ref.mut.sigs`
 A data.frame of reference mutational signatures

`target.mut.sigs`
 A character vector of target mutational signatures names

Value

A list with the following elements

- `num.point.mutations` An integer value - count of total point mutations
- `sub.types` A character vector of length 96
- `sub.types.spectrum` A numeric vector of length 96
- `num.mut.sigs` An integer value (count of unique mutational signatures identified)
- `identified.mut.sigs` A character vector where each element is a mutational signature identified
- `identified.mut.sigs.probs` A numeric vector where each element is the weight of mutational signature identified. The ordering follows `identified.mut.sigs`
- `identified.mut.sigs.spectrum` A numeric vector of length 96
- `residuals` A numeric vector of length 96
- `rss` A numeric value (residual sum of squares)
- `cos.sim.score` A numeric value (cosine similarity score between observed mutational spectrum and reconstructed mutational signatures)
- `all.models.sigs` A list where each element is a model; a model is a list of signatures identified
- `all.models.sigs.probs` A list where each element is a model; a model is a list of contribution probabilities
- `all.models.cos.sim.scores` A list where each element is a model; a model is a list of cosine similarity scores

RunMutPat

*RunMutPat***Description**

Identifies mutational signatures using Mutational Patterns

Usage

```
RunMutPat(mut.pat.input, df.mut.pat.ref.sigs, target.mut.sigs,
          verbose = TRUE)
```

Arguments

`mut.pat.input` A list from [MutPatParseVCFObj](#)
`df.mut.pat.ref.sigs` A data.frame returned by [MutPatParseRefMutSigs](#)
`target.mut.sigs` A character vector of target mutational signatures names
`verbose` If true, provides process details

Value

A list with the following elements

- `tumor.mutation.types.spectrum` A numeric vector of length 96 - 'observed' spectrum
- `identified.mutation.types.spectrum` A numeric vector of length 96 - 'identified' spectrum
- `residuals` A numeric vector of length 96 - residuals
- `mutation.types` A character vector of length 96
- `identified.mut.sigs` A character vector where each element is a mutational signature identified
- `identified.mut.sigs.contribution.weights` A numeric vector where each element is the weight of mutational signature identified. The ordering follows `identified.mut.sigs`
- `cosine.similarity.score` A numeric value

Examples

```
## Not run:
vcf.obj <- ReadVCF(vcf.file = "../data/sample/HNT-082-BT.final.call.vcf", genome = "hg19")
df.ref.mut.sigs <- GetPCAWGMutSigs()
target.mut.sigs <- GetPCAWGMutSigsNames()
RunMutPat(vcf.obj = vcf.obj,
          df.ref.mut.sigs = df.ref.mut.sigs,
          target.mut.sigs = target.mut.sigs)

## End(Not run)
```

TriNuc.Mutation.Type.Hex.Colors
<i>Constant</i>

Description

Hex codes for the mutation types (for plotting purposes)

Usage

TriNuc.Mutation.Type.Hex.Colors

Format

An object of class character of length 6.

UpdateFilter	<i>UpdateFilter</i>
--------------	---------------------

Description

Update filter based on optim parameter values

Usage

UpdateFilter(vcf.filter, param.values)

Arguments

- | | |
|--------------|--|
| vcf.filter | A list from MakeFilterFromConfig |
| param.values | A numeric vector contains filtering value (same length with length(vcf.config.filter)) |

Value

Updated vcf.filter (list)

`WriteFIREVATResultsToTSV`*WriteFIREVATResultsToTSV*

Description

Writes FIREVAT results to a csv file

Usage

```
WriteFIREVATResultsToTSV(firevat.results)
```

Arguments

`firevat.results`

List returned from RunFIREVAT

`WriteVCF`*WriteVCF*

Description

Writes a vcf.obj to a .vcf file

Usage

```
WriteVCF(vcf.obj, save.file)
```

Arguments

`vcf.obj` (from the function ReadVCF)

`save.file` (full path including filename)

Index

*Topic **datasets**

- Chromosome.Names, [5](#)
- PCAWG.All.Sequencing.Artifact.Signatures, [21](#)
- PCAWG.Known.Sequencing.Artifact.Signatures, [22](#)
- PCAWG.Likely.Sequencing.Artifact.Signatures, [22](#)
- PCAWG.Possible.Sequencing.Artifact.Signatures, [22](#)
- PCAWG.Target.Mutational.Signatures, [23](#)
- TriNuc.Mutation.Type.Hex.Colors, [48](#)
- AnnotateVCFObj, [3](#), [31](#), [40](#)
- CheckIfVariantRefinementIsNecessary, [4](#)
- Chromosome.Names, [5](#)
- ComputeZScore, [5](#)
- ComputeZScoreEqiValue, [6](#)
- DecimalCeiling, [6](#)
- Default.Obj.Fn, [7](#)
- DefaultFilterToBinary, [7](#)
- EnumerateTriNucCounts, [8](#)
- Euc.Exp.Weighted.Obj.Fn, [8](#)
- Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.1, [9](#)
- Euc.Exp.Weighted.Seq.Art.Only.Obj.Fn.2, [9](#)
- Euc.Obj.Fn, [10](#)
- Exp.Weighted.Obj.Fn.1, [10](#)
- Exp.Weighted.Obj.Fn.2, [11](#)
- Exp.Weighted.Refined.Seq.Art.Only.Obj.Fn, [11](#)
- FilterVCF, [12](#)
- GenerateConfigObj, [12](#)
- GetCOSMICMutSigs, [13](#)
- GetCOSMICMutSigsEtiologiesColors, [13](#)
- GetCOSMICMutSigsNames, [14](#)
- GetGASuggestedSolutions, [14](#)
- GetOptimizedSignatures, [15](#)
- GetParameterLowerUpperVector, [16](#)
- GetPCAWGMutSigs, [16](#)
- GetPCAWGMutSigsEtiologiesColors, [17](#)
- GetPCAWGMutSigsNames, [17](#)
- InitializeVCF, [17](#)
- MakeFilter, [7](#), [18](#)
- MutaliskParseVCFObj, [18](#)
- MutPatParseRefMutSigs, [19](#), [47](#)
- MutPatParseVCFObj, [19](#), [47](#)
- ParameterToBits, [7](#), [20](#)
- ParseConfigFile, [21](#)
- PCAWG.All.Sequencing.Artifact.Signatures, [21](#)
- PCAWG.Known.Sequencing.Artifact.Signatures, [22](#)
- PCAWG.Likely.Sequencing.Artifact.Signatures, [22](#)
- PCAWG.Possible.Sequencing.Artifact.Signatures, [22](#)
- PCAWG.Target.Mutational.Signatures, [23](#)
- PerformStrandBiasAnalysis, [23](#)
- PlotMutaliskResults, [24](#)
- PlotMutationTypes, [25](#)
- PlotOptimizationIterations, [26](#)
- PlotSignaturesContProbs, [27](#)
- PlotTable, [28](#)
- PlotTriNucSpectrum, [28](#)
- PlotVCFStatsBoxPlots, [29](#)
- PlotVCFStatsHistograms, [30](#)
- PrepareAnnotationDB, [3](#), [4](#), [31](#), [40](#), [44](#)
- PrepareArtifactAnnotationTable, [32](#)
- PrepareArtifactStrandBiasTable, [32](#)

PrepareArtifactualMutsOptimizationIterationsPlot,
33

PrepareFilterCutoffsTable, 33

PrepareGeneticAlgorithmParametersTable,
34

PrepareIdentifiedSignaturesPlot, 34

PrepareMLEReconstructedSpectrumsPlot,
35

PrepareNucleotideSubstitutionTypesPlot,
35

PrepareObservedSpectrumsPlot, 36

PrepareOptimizationResultsTable, 36

PrepareOptimizedVCFStatisticsPlot, 37

PrepareRefinedAnnotationTable, 37

PrepareRefinedMutsOptimizationIterationsPlot,
38

PrepareRefinedStrandBiasTable, 38

PrepareResidualSpectrumsPlot, 39

PrepareTrinucleotideSpectrumsTable, 39

QueryAnnotatedVCF, 40

ReadOptimizationIterationReport, 40

ReadVCF, 4, 13, 20, 23, 31, 41

ReportFIREVATResults, 41

RunFIREVAT, 15, 32–41, 42

RunMutalisk, 24, 44

RunMutaliskHelper, 46

RunMutPat, 47

TriNuc.Mutation.Type.Hex.Colors, 48

UpdateFilter, 48

WriteFIREVATResultsToTSV, 49

WriteVCF, 49