

Package ‘CRISPRcleanR’

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Type Package

Title Unsupervised correction of gene independent cell responses to CRISPR-cas9 targeting

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Description CRISPRcleanR is an R package for identifying and correcting gene independent responses to CRISPRcas9 targeting, in genome-wide pooled sgRNA drop-out screens. CRISPRcleanR uses an unsupervised approach based on the segmentation of single-guide RNA (sgRNA) fold change values across the genome, without making any assumption on the copy number status of the targeted genes. CRISPRcleanR reports sgRNA fold changes and normalised sgRNA read counts, and is therefore compatible with downstream analysis tools, and works with multiple sgRNA libraries.

Depends stringr, DNACopy, pROC

RoxygenNote 6.0.1

R topics documented:

| | |
|---|----|
| BAGEL_essential | 2 |
| BAGEL_nonEssential | 3 |
| CCLE.gisticCNA | 3 |
| ccr.cleanChrm | 4 |
| ccr.correctCounts | 7 |
| ccr.geneMeanFCs | 9 |
| ccr.genes2sgRNAs | 10 |
| ccr.get.CCLEgisticSets | 11 |
| ccr.get.gdsc1000.AMPgenes | 12 |
| ccr.get.nonExpGenes | 13 |
| ccr.GWclean | 15 |
| ccr.logFCs2chromPos | 18 |
| ccr.NormfoldChanges | 19 |
| ccr.PrecisionRecallCurve | 21 |
| ccr.VisDepAndSig | 23 |
| CL.subset | 25 |
| EPLC.272HcorrectedFCs | 25 |
| EssGenes.DNA_REPLICATION_cons | 27 |

| | |
|--|-----------|
| EssGenes.KEGG_rna_polymerase | 27 |
| EssGenes.PROTEASOME_cons | 28 |
| EssGenes.ribosomalProteins | 29 |
| EssGenes.SPLICEOSOME_cons | 29 |
| GDSC.CL_annotation | 30 |
| GDSC.geneLevCNA | 31 |
| KY_Library_v1.0 | 32 |
| RNAseq.fpkms | 33 |
| Index | 34 |

| | |
|-----------------|---|
| BAGEL_essential | <i>Reference Core fitness essential genes</i> |
|-----------------|---|

Description

A list of reference core fitness essential genes assembled from multiple RNAi studies used as classification template by the BAGEL algorithm to call gene depletion significance [1].

Usage

```
data(BAGEL_essential)
```

Format

A vector of strings containing HGNC symbols of reference core fitness essential genes.

References

[1] BAGEL: a computational framework for identifying essential genes from pooled library screens. Traver Hart and Jason Moffat. BMC Bioinformatics, 2016 vol. 17 p. 164.

See Also

[BAGEL_nonEssential](#)

Examples

```
data(BAGEL_essential)
head(BAGEL_essential)
```

| | |
|--------------------|---|
| BAGEL_nonEssential | <i>Reference set of non essential genes</i> |
|--------------------|---|

Description

A list of reference non essential genes assembled from multiple RNAi studies used as classification template by the BAGEL algorithm to call gene depletion significance [1].

Usage

```
data(BAGEL_nonEssential)
```

Format

A vector of strings containing HGNC symbols of reference non essential genes.

References

[1] BAGEL: a computational framework for identifying essential genes from pooled library screens. Traver Hart and Jason Moffat. BMC Bioinformatics, 2016 vol. 17 p. 164.

See Also

[BAGEL_essential](#)

Examples

```
data(BAGEL_nonEssential)
head(BAGEL_nonEssential)
```

| | |
|----------------|---|
| CCLC.gisticCNA | <i>Genome-wide copy number data for 13 human cancer cell lines.</i> |
|----------------|---|

Description

Genome-wide Gistic [1] scores quantifying copy number status across a subset of the cell lines in [CL.subset](#) that are used to assess CRISPRcleaner results in [2].

Usage

```
data(CCLC.gisticCNA)
```

Format

A data frame with one observations per gene across 13 variables (one per cell line). Row names indicate HGNC gene symbols and column names indicate cell line COSMIC identifiers [3].

Source

This data frame has been derived from the tsv file downloadable at http://www.cbioportal.org/study?id=cellline_ccle_broad#summary. This has been obtained by processing Affymetrix SNP array data in the Cancer Cell Line Encyclopedia [4] repository (https://portals.broadinstitute.org/ccle_legacy/data/)

References

- [1] Mermel CH, Schumacher SE, Hill B, et al. *GISTIC2.0 facilitates sensitive and confident localization of the targets of focal somatic copy-number alteration in human cancers*. Genome Biol. 2011;12(4):R41. doi: 10.1186/gb-2011-12-4-r41.
- [2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>
- [2] Forbes SA, Beare D, Boutselakis H, et al. *COSMIC: somatic cancer genetics at high-resolution* Nucleic Acids Research, Volume 45, Issue D1, 4 January 2017, Pages D777-D783,
- [3] Barretina J, Caponigro G, Stransky N, et al. *The Cancer Cell Line Encyclopedia enables predictive modelling of anticancer drug sensitivity*. Nature. 2012 Mar 28;483(7391):603-7. doi: 10.1038/nature11003. Erratum in: Nature. 2012 Dec 13;492(7428):290.

Examples

```
data(CCLE.gisticCNA)
head(CCLE.gisticCNA)
```

| | |
|----------------------------|---|
| <code>ccr.cleanChrm</code> | <i>Identification and correction of genomic regions of equal log fold changes involving sgRNAs targeting a minimal number of genes within a given chromosome.</i> |
|----------------------------|---|

Description

This function applies a circular binary segmentation algorithm [1, 2] to genomic-sorted log fold changes of all the sgRNAs targeting genes on the same chromosome. This procedure yields a sets of genomic regions of estimated equal sgRNAs' log fold changes, significantly differing on average from adjacent regions. If some of these regions fulfill certain criteria (detailed below) then they are deemed as responding to CRISPR-Cas9 targeting in a gene independent manner, i.e. they might be biased by local feature of the DNA) and their pattern of log fold changes is mean centered [3].

Usage

```
ccr.cleanChrm(gwSortedFCs,CHR,display=TRUE,label='',
              saveTO=NULL,min.ngenes=3,ignoredGenes=NULL)
```

Arguments

| | |
|--------------|---|
| gwSortedFCs | <p>A data frame containing genome-wide genomic-sorted sgRNAs' log fold changes. This data frame must include one named row per each sgRNAs and the following columns/headers:</p> <ul style="list-style-type: none"> • CHR: the chromosome of the gene targeted by the sgRNA under consideration; • startp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration; • endp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration; • genes: the HGNC symbol of the gene targeted by the sgRNA under consideration; • avgFC: the log fold change of the sgRNA under consideration averaged across replicates; • BP: the genomic coordinate of the sgRNA defined as $\text{STARTpos} + (\text{ENDpos} - \text{STARTpos})/2$. <p>This can be generated using the ccr.logFCs2chromPos function, starting from a data frame containing sgRNAs' log fold changes generated by the ccr.NormfoldChanges function from raw sgRNAs' counts.</p> |
| CHR | Numerical value indicating the chromosome to analyse and correct. X and Y chromosome must be indicated with 23 and 24, respectively. |
| display | A logical value indicating whether genomic plots showing the results of the biased regions' identification and their log fold change correction should be generated or not. |
| label | A string indicating the experiment name, used in the main title of the plots and for the name of the folder where results are saved. |
| saveTO | If different from NULL then it will contain the path where pdf files with then genomic plots showing the results of the biased regions' identification (and their log fold change correction) will be saved (within a folder named as defined in the label parameter). |
| min.ngenes | A numerical value (>0) specifying the minimal number of different genes that the set of sgRNAs within a region of estimated equal log fold changes should target in order for that region to be corrected, i.e. mean centered. |
| ignoredGenes | A vector of strings containing HGNC symbols of genes that should not be considered when computing the minimal number of different genes targeted by the sgRNAs in the same identified region of estimated equal log fold changes. This vector could contain, for example, a priori known essential genes. This parameter should be set to NULL for a completely unsupervised correction. |

Value

A list containing two data frames. The first one (correctedFCs) contains a named row per each sgRNA and the following columns/header:

- CHR: the chromosome of the gene targeted by the sgRNA under consideration;
- startp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration;
- endp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;

- genes: the HGNC symbol of the gene targeted by the sgRNA under consideration;
- avgFC: the log fold change of the sgRNA averaged across replicates;
- correction: the type of correction: 1 = increased, -1 = decreased;
- correctedFC: the corrected log fold change of the sgRNA

The second one (regions) contains the identified region of estimated equal log fold changes (one region per row) and the following columns/headers:

- CHR: the chromosome of the region under consideration;
- startp: the genomic coordinate of the starting position of the region under consideration;
- endp: the genomic coordinate of the ending position of the region under consideration;
- n.sgRNAs: the number of sgRNAs targeting sequences in the region under consideration;
- avg.logFC: the average log fold change of the sgRNAs targeting the region;
- guideIdx: the indexes range of the sgRNAs targeting the region under consideration as they appear in the gwSortedFCs provided in input.

Author(s)

Francesco Iorio (iorio@ebi.ac.uk)

References

- [1] Olshen, A. B., Venkatraman, E. S., Lucito, R., Wigler, M. (2004). *Circular binary segmentation for the analysis of array-based DNA copy number data*. Biostatistics 5: 557-572.
- [2] Venkatraman, E. S., Olshen, A. B. (2007). *A faster circular binary segmentation algorithm for the analysis of array CGH data*. Bioinformatics 23: 657-63.
- [3] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

See Also

[ccr.logFCs2chromPos](#), [ccr.NormfoldChanges](#)

Examples

```
data(KY_Library_v1.0)

fn<-paste(system.file('extdata', package = 'CRISPRcleanR'),'HT-29_counts.tsv',sep='')
normANDfcs<-ccr.NormfoldChanges(fn,min_reads=30,EXPname='Example',
                                libraryAnnotation=KY_Library_v1.0)

gwSortedFCs<-ccr.logFCs2chromPos(normANDfcs$logFCs,KY_Library_v1.0)

chr8cleaned<-ccr.cleanChrm(gwSortedFCs,8,display=TRUE,label='HT-29',
                           min.ngenes=3)
```

| | |
|-------------------|---|
| ccr.correctCounts | <i>Correction of sgRNA treatment counts for gene independent responses to CRISPR-Cas9 targeting</i> |
|-------------------|---|

Description

This function applies an inverse transformation (described in ...) to CRISPRcleanR corrected sgRNAs' log fold changes and produces in output normalised corrected sgRNA counts (across treatments and control replicates), suitable for gene depletion/enrichment statistical testing via mean-variance modeling (for example through MAGeCK [1]*). *MAGeCK should be executed excluding initial normalisation, as the corrected sgRNA counts outputted by this function are already normalised.

Usage

```
ccr.correctCounts(CL,normalised_counts,
                  correctedFCs_and_segments,
                  libraryAnnotation,
                  minTargetedGenes=3,
                  OutDir='./',
                  ncontrols=1)
```

Arguments

- | | |
|---------------------------|---|
| CL | A string specifying the name of the experiment. This will be used to compose names of files and folde where results will be saved. |
| normalised_counts | A data frame containing normalised sgRNAs' read counts, which can be computed using the ccr.NormfoldChanges function from raw sgRNAs' counts. |
| correctedFCs_and_segments | sgRNAs log fold changes corrected for gene independent responses, generated with the function ccr.GWclean. |
| libraryAnnotation | <p>A data frame containing the sgRNAs' genome-wide annotations with at least a named row for each of the sgRNAs included in the foldchanges data frame provided in input. The following columns/headers should be present in this data frame (additional columns will be ignored):</p> <ul style="list-style-type: none"> • GENES: string vector containing the HGNC symbols of the genes targeted by the sgRNA under consideration; • EXONE: string vector containing the gene exon targeted by the sgRNA under consideration (these should include the prefix "ex" followed by the exone number); • CHRM: string vector the chromosome of the gene targeted by the sgRNA under consideration (X and Y chromosome should be specified as "X" and "Y"); • STRAND: string vector containing the strand targeted by the sgRNA under consideration ("+" or "-"); • STARTpop: numeric vector containing the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration; |

- ENDpos: numeric vector containing the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;

| | |
|------------------|--|
| minTargetedGenes | Minimanl number of different genes targeted by sgRNAs in a biased segment in order for the corresponding counts to be corrected (default = 3). |
| OutDir | Path of the folder where results and plots will be saved. |
| ncontrols | A numerical value indicating the number of control replicates (therefore columns to be considered as controls in the normalised counts). |

Value

A data frame with one entry per sgRNA and individual columns for the control/treatment samples included in the normalised count data object specified by the `normalised_counts` parameter, and containing sgRNA counts corrected for gene independent responses to CRISPR-Cas9 targeting and median-ratio normalised.

Author(s)

Francesco Iorio (fi9323@gmail.com)

References

[1] Li, W., Xu, H., Xiao, T., Cong, L., Love, M. I., Zhang, F., et al. (2014). MAGeCK enables robust identification of essential genes from genome-scale CRISPR/Cas9 knockout screens. *Genome Biology*, 15(12), 554. [2] Hart, T., & Moffat, J. (2016). BAGEL: a computational framework for identifying essential genes from pooled library screens. *BMC Bioinformatics*, 17(1), 164.

See Also

[ccr.NormfoldChanges](#), [ccr.GWclean](#)

Examples

```
## Loading sgRNA library annotation file
data(KY_Library_v1.0)

## Deriving the path of the file with the example dataset,
## from the mutagenesis of the EPLC-272H colorectal cancer cell line
fn<-paste(system.file('extdata', package = 'CRISPRcleanR'),
           '/EPLC-272H_counts.tsv', sep='')

## Loading, median-normalizing and computing fold-changes for the example dataset
normANDfcs<-ccr.NormfoldChanges(fn,min_reads=30,
                                EXPname='EPLC-272H',
                                libraryAnnotation = KY_Library_v1.0)

## Genome-sorting of the fold changes
gwSortedFCs<-ccr.logFCs2chromPos(normANDfcs$logFCs,KY_Library_v1.0)

## Identifying and correcting biased sgRNAs' fold changes
correctedFCs<-ccr.GWclean(gwSortedFCs,display=FALSE,label='EPLC-272H')

## correcting individual sgRNA treatment counts

correctedCounts<-ccr.correctCounts('EPLC-272H',normANDfcs$norm_counts,
```



```

        correctedFCs,
        KY_Library_v1.0,
        minTargetedGenes=3,
        OutDir='./')

head(correctedCounts)

```

| | |
|-----------------|------------------------------------|
| ccr.geneMeanFCs | <i>Gene level log fold changes</i> |
|-----------------|------------------------------------|

Description

This functions computes gene level log fold changes based on average log fold changes of targeting sgRNAs

Usage

```
ccr.geneMeanFCs(sgRNA_FCprofile, libraryAnnotation)
```

Arguments

sgRNA_FCprofile
A named numerical vector containing the sgRNAs' log fold-changes, with names corresponding to sgRNAs identifiers.

libraryAnnotation
A data frame containing the sgRNA library annotation (with same format of [KY_Library_v1.0](#)).

Value

A numerical vector containing gene average log fold-changes, with corresponding HGNC symbols as names.

Author(s)

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See Also

[KY_Library_v1.0](#)

Examples

```

## loading corrected sgRNAs log fold-changes and segment annotations for
## an example cell line (EPLC-272H)
data(EPLC.272HcorrectedFCs)

## loading sgRNA library annotation
data(KY_Library_v1.0)

## storing sgRNA log fold-changes in a named vector
FCs<-EPLC.272HcorrectedFCs$corrected_logFCs$avgFC

```

```
names(FCs)<-rownames(EPLC.272HcorrectedFCs$corrected_logFCs)

## computing gene level log fold-changes
geneFCs<-ccr.geneMeanFCs(FCs,KY_Library_v1.0)

head(geneFCs)
```

| | |
|------------------|-------------------------|
| ccr.genes2sgRNAs | <i>Targeting sgRNAs</i> |
|------------------|-------------------------|

Description

This function returns the set of sgRNAs targeting the set of genes provided in input, in a given pooled library.

Usage

```
ccr.genes2sgRNAs(libraryAnnotation,genes)
```

Arguments

| | |
|-------------------|--|
| libraryAnnotation | A data frame with a named row for each sgRNA with the same format of KY_Library_v1.0 |
| genes | A list of strings containing HGNC symbols |

Value

A list of strings containing the identifiers of the sgRNAs targeting the inputted set of genes

Author(s)

Francesco Iorio (fi9323@gmail.com)

See Also

[KY_Library_v1.0](#)

Examples

```
## Loading an sgRNA pooled library annotation
data(KY_Library_v1.0)
## Loading an example set of genes
data(BAGEL_essential)

ccr.genes2sgRNAs(KY_Library_v1.0,BAGEL_essential)
```

ccr.get.CCLEgisticSets

CCLE gistic score gene sets

Description

This function splits all the genes into 5 classes (-2, -1, 0, +1 and +2) based on the CNA Gistic [1] score observed in a given cell line.

Usage

```
ccr.get.CCLEgisticSets(cellLine,CCLE.gisticCNA=NULL)
```

Arguments

| | |
|----------------|--|
| cellLine | A string specifying the name of a cell line (or a COSMIC identifier [2]); |
| CCLE.gisticCNA | Genome-wide Gistic [1] scores quantifying copy number status across cell lines with the same format of CCLE.gisticCNA . If NULL then this function uses the CCLE.gisticCNA builtin data frame, containing data for 13 cell lines of the 15 used in [3] to assess the performances of CRISPRcleanR. |

Value

A named list of vectors with the following fields:

| | |
|-----|---|
| gm2 | A vector of strings containing identifiers of sgRNAs targeting genes whit a Gistic score = -2 in the cell line under consideration; |
| gm1 | A vector of strings containing identifiers of sgRNAs targeting genes whit a Gistic score = -1 in the cell line under consideration; |
| gz | A vector of strings containing identifiers of sgRNAs targeting genes whit a Gistic score = 0 in the cell line under consideration; |
| gp1 | A vector of strings containing identifiers of sgRNAs targeting genes whit a Gistic score = +1 in the cell line under consideration; |
| gp2 | A vector of strings containing identifiers of sgRNAs targeting genes whit a Gistic score = +2 in the cell line under consideration; |

Author(s)

Francesco Iorio (iorio@ebi.ac.uk)

References

- [1] Mermel CH, Schumacher SE, Hill B, et al. *GISTIC2.0 facilitates sensitive and confident localization of the targets of focal somatic copy-number alteration in human cancers*. Genome Biol. 2011;12(4):R41. doi: 10.1186/gb-2011-12-4-r41.
- [2] Forbes SA, Beare D, Boutselakis H, et al. *COSMIC: somatic cancer genetics at high-resolution* Nucleic Acids Research, Volume 45, Issue D1, 4 January 2017, Pages D777-D783,

[3] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

See Also

[ccr.get.gdsc1000.AMPgenes](#)

Examples

```
GS<-ccr.get.CCLEgisticSets('HT-29')

head(GS$gm2)
head(GS$gm1)
head(GS$gz)
head(GS$gp1)
head(GS$gp2)
```

```
ccr.get.gdsc1000.AMPgenes
```

Copy number amplified genes in a given cell line from the GDSC1000

Description

This function takes in input the name (or the COSMIC identifier [1]) of a cell line included in the GDSC1000 project [2] and it identifies the genes that are copy number amplified (according to a user defined minimal copy number value) in that cell line, using gene level copy number data from the Genomics of Drug Sensitivity in 1,000 Cancer Cell lines (GDSC1000) [2].

Usage

```
ccr.get.gdsc1000.AMPgenes(cellLine, minCN = 8, exact = FALSE,
                           GDSC.geneLevCNA=NULL)
```

Arguments

| | |
|-----------------|---|
| cellLine | A string specifying the name of a cell line used in (or a COSMIC identifier [1]); |
| minCN | Lower threshold for the minimum copy number of any genomic segment containing coding sequence of a gene in order for it to be considered as copy number amplified. |
| exact | If TRUE, then those genes for which any genomic segment containing coding sequence has a minimum copy number equal to minCN are considered as copy number amplified. |
| GDSC.geneLevCNA | Genome-wide copy number data with the same format of GDSC.geneLevCNA . This can be assembled from the xls sheet specified in the source section [a] (containing data for the GDSC1000 cell lines). If NULL, then this function uses the data in the built in GDSC.geneLevCNA data frame, containing data derived from [a] for 15 cell lines used in [3] to assess the performances of CRISPRcleanR. |

Value

A data frame, containing one row for each copy number amplified gene with the following columns:

| | |
|-------|---|
| Gene | HGNC symbol of the gene; |
| minCN | Minimum copy number of any genomic segment containing coding sequence of the gene in the cell line under consideration. |

Author(s)

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Source

[a] ftp://ftp.sanger.ac.uk/pub/project/cancerrxgene/releases/release-6.0/Gene_level_CN.xlsx.

References

- [1] Forbes SA, Beare D, Boutselakis H, et al. *COSMIC: somatic cancer genetics at high-resolution* Nucleic Acids Research, Volume 45, Issue D1, 4 January 2017, Pages D777-D783,
- [2] Iorio F, Knijnenburg TA, Vis DJ, Bignell GR, Menden MP, et al. *A landscape of pharmacogenomic interactions in cancer* Cell 2016 Jul 28;166(3):740-54
- [3] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

See Also

[ccr.get.CCLEgisticSets](#)

Examples

```
CNAgene<-
  ccr.get.gdsc1000.AMPgenes('HT-29')
head(CNAgene)
```

ccr.get.nonExpGenes *Non expressed genes in a given cell line*

Description

This function takes in input the name (or the COSMIC identifier [1]) of a cell line and it identifies genes that are not expressed (according to a user defined FPKM threshold) using a collection of RNAseq profile from [2].

Usage

```
ccr.get.nonExpGenes(cellLine, th = 0.05,
                    amplified = FALSE, minCN = 8,
                    RNAseq.fpkms=NULL)
```

Arguments

| | |
|--------------|---|
| cellLine | A string specifying the name of a cell line (or a COSMIC identifier [1]); |
| th | Minimum FPKM value for a gene to be considered as expressed; |
| amplified | A logic value specifying whether the selected not expressed genes should be also copy number amplified function; |
| minCN | If amplified = TRUE, this parameter defines a lower threshold for the minimum copy number of any genomic segment containing coding sequence of a gene in order for it to be considered as copy number amplified. |
| RNAseq.fpkms | Genome-wide substitute reads with fragments per kilobase of exon per million reads mapped (FPKM) across cell lines. These can be derived from a comprehensive collection of RNAseq profiles described in [2]. The format must be the same of the RNAseq.fpkms builtin data frame. If NULL then this function uses the RNAseq.fpkms builtin data fram containing data for 15 cell lines used in [3] to assess CRISPRcleaner results. |

Value

A vector of string containing the HGNC symbols of non expressed (optionally copy number amplified) genes in the cell line under consideration.

Author(s)

Francesco Iorio (fi9323@gmail.com)

References

- [1] Forbes SA, Beare D, Boutselakis H, et al. *COSMIC: somatic cancer genetics at high-resolution* Nucleic Acids Research, Volume 45, Issue D1, 4 January 2017, Pages D777-D783.
- [2] Garcia-Alonso L, Iorio F, Matchan A, et al. *Transcription factor activities enhance markers of drug response in cancer* doi: <https://doi.org/10.1101/129478>
- [3] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

See Also

[ccr.get.gdsc1000.AMPgenes](#)

Examples

```
ccr.get.nonExpGenes('HT-29', amplified = TRUE)
```

ccr.GWclean

Unsupervised identification and correction of gene independent cell responses to CRISPR-Cas9 targeting.

Description

This function takes in input a genome-wide essentiality profile derived from a CRISPR-Cas9 experiment employing a pooled library of single guide RNAs (sgRNAs) targeting protein coding genes, which are transfected in an *in vitro* model stably expressing Cas9. The essentiality profile quantifies the loss/gain-of-fitness caused by each sgRNA-targeting, and it is expressed as log fold changes (logFCs) between the abundance of the sgRNAs at an end point after cell purification and their abundance in the plasmid pool used for viral production, or at an initial time point, or in any other control condition. A circular binary segmentation algorithm [1, 2] is applied by this function to the genome-wide pattern of logFCs provided in input, in order to identify genomic regions including sgRNAs with sufficiently equal logFC (and mean logFC sufficiently different from background) and targeting a minimal number of different genes. Assuming that it is very unlikely to observe the same loss/gain-of-fitness effect when targeting a large number of contiguous genes, if certain user-defined condition (detailed below) are met then the logFCs of such regions are deemed as biased by some local feature of the involved genomic segment (which could be, for example, copy number amplified [3]), and they are corrected, i.e. mean centered [4].

Usage

```
ccr.GWclean(gwSortedFCs,label='',display=TRUE,
            saveT0=NULL,ignoredGenes=NULL,min.ngenes=3)
```

Arguments

- | | |
|-------------|---|
| gwSortedFCs | <p>A data frame containing genome-wide genomic-sorted sgRNAs' log fold changes. This data frame must include one named row per each sgRNA and the following columns/headers:</p> <ul style="list-style-type: none"> • CHR: the chromosome of the gene targeted by the sgRNA under consideration; • startp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration; • endp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration; • genes: the HGNC symbol of the gene targeted by the sgRNA under consideration; • avgFC: the log fold change of the sgRNA under consideration averaged across replicates; • BP: the genomic coordinate of the sgRNA defined as $\text{STARTpos} + (\text{ENDpos} - \text{STARTpos})/2$. |
|-------------|---|

This can be generated using the [ccr.logFCs2chromPos](#) function, starting from a data frame containing sgRNAs' log fold changes generated by the [ccr.NormfoldChanges](#) function (from raw sgRNAs' counts), from raw sgRNAs' counts.

| | |
|--------------|--|
| label | A string indicating the experiment name. This is used to compose the main title of the plots generated by this function and the name of the folder where the results are saved. |
| display | A logical value indicating whether genomic plots showing the results of the biased regions' identification and their log fold change correction should be generated or not. |
| saveTO | If different from NULL then this parameter will contain the path where pdf files with then genomic plots showing the results of the biased regions' identification (and their log fold change correction) will be saved (within a folder named as defined in the label parameter). |
| ignoredGenes | A vector of strings containing HGNC symbols of genes that should not be considered when computing the minimal number of different genes targeted by sgRNAs in the same identified region of estimated equal log fold changes. This could contain, for example, a-priori known essential genes. |
| min.ngenes | A numerical value (>0) specifying the minimal number of different genes that the set of sgRNAs within a region of estimated equal logFCs should target in order for their logFCs to be corrected, i.e. mean centered. |

Value

A list containing two data frames and a vector of strings. The first data frame (corrected_logFCs) contains a named row per each sgRNA and the following columns/header:

- CHR: the chromosome of the gene targeted by the sgRNA under consideration;
- startp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration;
- endp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;
- genes: the HGNC symbol of the gene targeted by the sgRNA under consideration;
- avgFC: the log fold change of the sgRNA averaged across replicates;
- correction: the type of correction: 1 = increased log fold change, -1 = decreased log fold change. 0 indicates no correction;
- correctedFC: the corrected log fold change of the sgRNA

The second data frame (segments) contains the identified region of estimated equal log fold changes (one region per row) and the following columns/headers:

- CHR: the chromosome of the gene targeted by the sgRNA under consideration;
- startp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration;
- endp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;
- genes: the HGNC symbol of the gene targeted by the sgRNA under consideration;
- avgFC: the log fold change of the sgRNA averaged across replicates;
- correction: the type of correction: 1 = increased log fold change, -1 = decreased log fold change. 0 indicates no correction;
- correctedFC: the corrected log fold change of the sgRNA

The second data frame (segments) contains the identified region of estimated equal log fold changes (one region per row) and the following columns/headers:

- CHR: the chromosome of the region under consideration;
- startp: the genomic coordinate of the starting position of the region under consideration;
- endp: the genomic coordinate of the ending position of the region under consideration;
- n.sgRNAs: the number of sgRNAs targeting sequences in the region under consideration;
- avg.logFC: the average log fold change of the sgRNAs in the region;
- guideIdx: the indexes range of the sgRNAs targeting the region under consideration as they appear in the gwSortedFCs provided in input.

The string of vectors (SORTED_sgRNAs) contains the sgRNAs' identifiers in the same order as they are reported in the gwSortedFCs input data frame, i.e. genome sorted.

Author(s)

Francesco Iorio (iorio@ebi.ac.uk)

References

- [1] Olshen, A. B., Venkatraman, E. S., Lucito, R., Wigler, M. (2004). *Circular binary segmentation for the analysis of array-based DNA copy number data*. Biostatistics 5: 557-572. \
- [2] Venkatraman, E. S., Olshen, A. B. (2007). *A faster circular binary segmentation algorithm for the analysis of array CGH data*. Bioinformatics 23: 657-63. \
- [3] Andrew J. Aguirre, Robin M. Meyers, Barbara A. Weir, Francisca Vazquez, Cheng-Zhong Zhang, Uri Ben-David, April Cook, Gavin Ha, William F. Harrington, Mihir B. Doshi, Maria Kost-Alimova, Stanley Gill, Han Xu, Levi D. Ali, Guozhi Jiang, Sasha Pantel, Yenarae Lee, Amy Goodale, Andrew D. Cherniack, Coyin Oh, Gregory Kryukov, Glenn S. Cowley, Levi A. Garraway, Kimberly Stegmaier, Charles W. Roberts, Todd R. Golub, Matthew Meyerson, David E. Root, Aviad Tsherniak and William C. Hahn. *Genomic copy number dictates a gene-independent cell response to CRISPR-Cas9 targeting*. Cancer Discov June 3 2016 DOI: 10.1158/2159-8290.CD-16-0154
- [4] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

See Also

[ccr.cleanChrm](#)

Examples

```
## Loading sgRNA library annotation file
data(KY_Library_v1.0)

## Deriving the path of the file with the example dataset,
## from the mutagenesis of the HT-29 colorectal cancer cell line
fn<-paste(system.file('extdata', package = 'CRISPRcleanR'), '/HT-29_counts.tsv', sep='')

## Loading, median-normalizing and computing fold-changes for the example dataset
normANDfcs<-ccr.NormfoldChanges(fn,min_reads=30,EXpname='HT-29',
```

```

libraryAnnotation = KY_Library_v1.0)

## Genome-sorting of the fold changes
gwSortedFCs<-ccr.logFCs2chromPos(normANDfcs$logFCs,KY_Library_v1.0)

## Identifying and correcting biased sgRNAs' fold changes
correctedFCs<-ccr.GWclean(gwSortedFCs,display=TRUE,label='HT-29')

## Visualising first five entries of the corrected fold changes
head(correctedFCs$corrected_logFCs)

```

ccr.logFCs2chromPos *Genomic sorting of sgRNAs' log fold changes.*

Description

This function maps genome-wide sgRNAs' log fold changes (averaged across replicates) on the genome and returns them sorted according to the position of their targeted region on the chromosomes.

Usage

```
ccr.logFCs2chromPos(foldchanges, libraryAnnotation)
```

Arguments

foldchanges A data frame containing genome-wide sgRNAs' log fold changes, one column per library transfection replicate, with first and second column containing the sgRNAs' identifiers and the HGNC symbols of the targeted genes, respectively. This can be generated from raw count files using the [ccr.NormfoldChanges](#) function.

libraryAnnotation A data frame containing the sgRNAs' genome-wide annotations with at least a named row for each of the sgRNAs included in the foldchanges data frame provided in input. The following columns/headers should be present in this data frame (additional columns will be ignored):

- **GENES**: string vector containing the HGNC symbols of the genes targeted by the sgRNA under consideration;
- **EXONE**: string vector containing the gene exon targeted by the sgRNA under consideration (these should include the prefix "ex" followed by the exon number);
- **CHRM**: string vector the chromosome of the gene targeted by the sgRNA under consideration (X and Y chromosome should be specified as "X" and "Y");
- **STRAND**: string vector containing the strand targeted by the sgRNA under consideration ("+" or "-");
- **STARTpop**: numeric vector containing the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration;
- **ENDpos**: numeric vector containing the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;

Additional columns can be optionally included and will be ignored by this function. The annotation for the genome-wide sgRNA library presented in [1] is included in the [KY_Library_v1.0](#) data object, formatted as described above.

Value

A data frame with a named row per each sgRNA and the following columns/headers:

- CHR: the chromosome where the gene targeted by the sgRNA under consideration resides;
- startp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration;
- endp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;
- avgFC: the log fold change of the sgRNA averaged across replicates;
- BP: the genomic coordinate of the sgRNA defined as $STARTpos + (ENDpos - STARTpos) / 2$.

Author(s)

Francesco Iorio (iorio@ebi.ac.uk)

References

[1] Tzelepis K, Koike-Yusa H, De Braekeleer E, Li Y, Metzakopian E, Dovey OM, Mupo A, Grinkevich V, Li M, Mazan M, Gozdecka M, Onishi S, Cooper J, Patel M, McKerrell T, Chen B, Domingues AF, Gallipoli P, Teichmann S, Ponstingl H, McDermott U, Saez-Rodriguez J, Huntly BJP, Iorio F, Pina C, Vassiliou GS, Yusa K. *A CRISPR dropout screen identifies genetic vulnerabilities and therapeutic targets in acute myeloid leukaemia*. Cell Reports 2016 Oct 18;17(4):1193-1205

See Also

[ccr.NormfoldChanges](#), [KY_Library_v1.0](#)

Examples

```
data(KY_Library_v1.0)
fn<-paste(system.file('extdata', package = 'CRISPRcleanR'), '/A2058_counts.tsv', sep='')
normANDfcs<-ccr.NormfoldChanges(fn,min_reads=30,
                                EXPname='Example',
                                libraryAnnotation=KY_Library_v1.0)
mappedLogFCs<-ccr.logFCs2chromPos(normANDfcs$logFCs,KY_Library_v1.0)
head(mappedLogFCs)
```

| | |
|---------------------|---|
| ccr.NormfoldChanges | <i>Median-ratio normalisation of sgRNA counts and fold change computation</i> |
|---------------------|---|

Description

This function median-ratio normalises [1,2] sgRNAs' counts stored in a tsv file whose path is provided in input, to adjust for the effect of library size and read count distributions. It computes log fold changes of transfected library replicates versus controls (typically the sgRNA counts in the plasmid). The output of this function is returned as a list, and it is also saved into two tsv files.

Usage

```
ccr.NormfoldChanges(filename, display=TRUE, saveToFig=FALSE,
                    outdir='.', min_reads=30, EXPname='',
                    libraryAnnotation, ncontrols=1)
```

Arguments

| | |
|-------------------|--|
| filename | <p>A string specifying the path of a tsv file containing the raw sgRNA counts. This must be a tab delimited file with one row per sgRNA and the following columns/headers:</p> <ul style="list-style-type: none"> • sgRNA: containing alphanumeric identifiers of the sgRNA under consideration; • gene: containing HGNC symbols of the genes targeted by the sgRNA under consideration; <p>followed by the columns containing the sgRNAs' counts for the controls and columns for library trasfected samples.</p> |
| display | A logic value specifying whether figures containing boxplots with the count values pre/post normalisation and log fold-changes should be visualised (TRUE, by default). |
| saveToFig | A logic value specifying whether figures containing boxplots with the count values pre/post normalisation and log fold-changes should be saved as pdf files (FALSE, by default). Setting this parameter to TRUE overrides the value of the display parameter. |
| outdir | Path of the directory where the normalised sgRNAs' counts and the log fold changes, as well as the pdf files (if the parameter saveToFig is set to TRUE), must be saved. |
| min_reads | This parameter defines a filter threshold value for sgRNAs, based on their average counts in the control sample. Specifically, it indicates the minimal number of counts that each individual sgRNA needs to have in the controls (on average) in order to be included in the output. |
| EXPname | A string specifying the name of the experiment. This will be used to compose main title of the generated figures and file names. |
| libraryAnnotation | A data frame containing the sgRNA annotations, with a named row for each sgRNA, and columns for targeted genes, genomic coordinates and possibly other informations. This should be formatted as the KY_Library_v1.0 data object containing the annotation of the sgRNA library presented in [3]. |
| ncontrols | A numerical value indicating the number of control replicates (therefore columns to be considered as control counts after the first two, in the inputted tsv file). |

Value

A list containing two data frames: for the normalised sgRNAs' counts (norm_counts) and the sgRNAs' log fold changes (logFCs) respectively. First two columns in these data frames contain sgRNAs' identifiers and HGNC symbols of targete gene, respectively.

Author(s)

Francesco Iorio (fi9323@gmail.com)

References

- [1] Wang T, Wei JJ, Sabatini DM, Lander ES. *Genetic screens in human cells using the CRISPR-Cas9 system*. Science. 2014, 343: 80-84.
- [2] Anders S, Huber W. *Differential expression analysis for sequence count data*. Genome Biol. 2010, 11: R106
- [3] Tzelepis K, Koike-Yusa H, De Braekeleer E, et al *A CRISPR dropout screen identifies genetic vulnerabilities and therapeutic targets in acute myeloid leukaemia*. Cell Reports 2016 Oct 18;17(4):1193-1205

See Also

[KY_Library_v1.0](#)

Examples

```
## loading sgRNA library annotation
data(KY_Library_v1.0)

## derive path for an example dataset
fn<-paste(system.file('extdata', package = 'CRISPRcleanR'), '/HT-29_counts.tsv', sep='')

## sgRNAs' normalisation and computation of log fold-changes
normANDfcs<-ccr.NormfoldChanges(fn,
                                min_reads=30,
                                EXPname='Example',
                                libraryAnnotation=KY_Library_v1.0)

## inspecting first 5 entries of the data frames containing the
## normalised counts and the log fold-changes
head(normANDfcs$norm_counts)
head(normANDfcs$logFCs)
```

ccr.PrecisionRecallCurve

*Classification performances of reference sets of genes (or sgRNAs)
based on depletion log fold-changes*

Description

This functions computes Precision/Recall (PR) curve, area under the PR curve and (optionally) Recall at fixed false discovery rate (and corresponding log fold change threshold) when classifying reference sets of genes (or sgRNAs) based on their depletion log fold-changes

Usage

```
ccr.PrecisionRecallCurve(FCsprofile,
                          positives,
                          negatives,
                          display = TRUE,
                          FDRth = NULL)
```

Arguments

| | |
|------------|--|
| FCsprofile | A numerical vector containing gene average depletion log fold changes (or sgRNAs' depletion log fold changes) with names corresponding to HGNC symbols (or sgRNAs' identifiers). |
| positives | A vector of strings containing a reference set of positive cases: HGNC symbols of essential genes or identifiers of their targeting sgRNAs. This must be a subset of FCsprofile names, disjointed from negatives. |
| negatives | A vector of strings containing a reference set of negative cases: HGNC symbols of essential genes or identifiers of their targeting sgRNAs. This must be a subset of FCsprofile names, disjointed from positives. |
| display | A logical parameter specifying if a plot containing the computed precision/recall curve with ROC indicators should be plotted (default = TRUE). |
| FDRth | If different from NULL, will be a numerical value ≥ 0 and ≤ 1 specifying the false discovery rate threshold at which fixed recall will be computed. In this case, if the display parameter is TRUE, an orizontal dashed line will be added to the plot at the resulting recall and its value will be visualised in the legend. |

Value

A list containint three numerical variable AUC, Recall, and sigthreshold indicating the area under the precision/recall curve and (if FDRth is not NULL) the recall at the specifying false discovery rate and the corresponding log fold change threshold (both equal to NULL, if FDRth is NULL), respectively.

Author(s)

Francesco Iorio (fi9323@gmail.com)

See Also

[BAGEL_essential](#), [BAGEL_nonEssential](#), [ccr.genes2sgRNAs](#), [ccr.VisDepAndSig](#)

Examples

```
## loading corrected sgRNAs log fold-changes and segment annotations for an example
## cell line (EPLC-272H)
data(EPLC.272HcorrectedFCs)

## loading reference sets of essential and non-essential genes
data(BAGEL_essential)
data(BAGEL_nonEssential)

## loading library annotation
data(KY_Library_v1.0)

## storing sgRNA log fold-changes in a named vector
FCs<-EPLC.272HcorrectedFCs$corrected_logFCs$avgFC
names(FCs)<-rownames(EPLC.272HcorrectedFCs$corrected_logFCs)

## deriving sgRNAs targeting essential and non-essential genes (respectively)
BAGEL_essential_sgRNAs<-ccr.genes2sgRNAs(KY_Library_v1.0,BAGEL_essential)
BAGEL_nonEssential_sgRNAs<-ccr.genes2sgRNAs(KY_Library_v1.0,BAGEL_nonEssential)
```

```
## computing classification performances at the sgRNA level
ccr.PrecisionRecallCurve(FCs,BAGEL_essential_sgRNAs,BAGEL_nonEssential_sgRNAs)

## computing gene level log fold-changes
geneFCs<-ccr.geneMeanFCs(FCs,KY_Library_v1.0)

## computing classification performances at the sgRNA level, with Recall at 5% FDR
ccr.PrecisionRecallCurve(geneFCs,BAGEL_essential,BAGEL_nonEssential,FDRth = 0.05)
```

| | |
|------------------|--|
| ccr.VisDepAndSig | <i>Depletion profile visualisation with genes signatures superimposed and recall</i> |
|------------------|--|

Description

This functions ranks the gene (or sgRNAs) log fold changes. Based on this it determines a log fold change threshold based on a user defined false discovery rate when classifying two gene (sgRNA) positive/negative references sets (typically core-fitness-essential and non-essential genes), and it computes the Recall (or True Positive Rate) of genes in other user defined sets at the determined threshold. It produces a plot where the log fold changes are visualised alongside the rank positions of the genes included in the inputted sets and, their recall and the determined FDR threshold.

Usage

```
ccr.VisDepAndSig(FCsprofile,SIGNATURES,TITLE='',
                 pIs=NULL,nIs=NULL,
                 th=0.05,plotFCprofile=TRUE)
```

Arguments

| | |
|---------------|--|
| FCsprofile | A numerical vector containing gene average depletion log fold changes (or sgRNAs' depletion log fold changes) with names corresponding to HGNC symbols (or sgRNAs' identifiers). |
| SIGNATURES | A named list of vectors containing HGNC gene symbols. Two of these lists are used as classification template (respectively for positive and negative cases) to determine a log fold-change threshold providing a user defined classification false discovery rate. |
| TITLE | A string specifying the title of the plot. |
| pIs | The index position of the signature that contains the positive cases of the classification template. |
| nIs | The index position of the signature that contains the negative cases of the classification template. |
| th | A numerical value specifying the desired classification false discovery rate (this must be a real number between 0 and 1). |
| plotFCprofile | A logic value specifying whether the log fold changes should be plotted. |

Value

A named numerical vector containing recall scores for all the inputted signatures at the computed false discovery rate threshold for log fold-changes.

CL.subset

*COSMIC identifiers of 15 immortalised human cancer cell lines***Description**

COSMIC identifiers [1] of 15 cell lines included in the GDSC1000 panel [2] that are used in [3] to assess CRISPRcleanR results.

Usage

```
data(CL.subset)
```

Format

A vector of strings.

References

[1] Forbes SA, Beare D, Boutselakis H, et al. *COSMIC: somatic cancer genetics at high-resolution* Nucleic Acids Research, Volume 45, Issue D1, 4 January 2017, Pages D777-D783,

[2] Iorio F, Knijnenburg TA, Vis DJ, Bignell GR, Menden MP, et al. *A landscape of pharmacogenomic interactions in cancer* Cell 2016 Jul 28;166(3):740-54

[3] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

Examples

```
data(CL.subset)

## Loading annotation for the GDSC1000 cell lines
data(GDSC.CL_annotation)

## Visualising annotation
GDSC.CL_annotation[CL.subset,]
```

EPLC.272HcorrectedFCs *CRISPRcleanR corrected data for an example cell line*

Description

This list contains corrected sgRNAs log fold-changes and segment annotations for an example cell line (EPLC-272H), obtained using the `ccr.GWclean` function, as detailed in its reference manual entry [ccr.GWclean](#).

Usage

```
data("EPLC.272HcorrectedFCs")
```

Format

A list containing two data frames and a vector of strings. The first data frame (corrected_logFCs) contains a named row per each sgRNA and the following columns/header:

- CHR: the chromosome of the gene targeted by the sgRNA under consideration;
- starttp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration;
- endtp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;
- genes: the HGNC symbol of the gene targeted by the sgRNA under consideration;
- avgFC: the log fold change of the sgRNA averaged across replicates;
- correction: the type of correction: 1 = increased log fold change, -1 = decreased log fold change. 0 indicates no correction;
- correctedFC: the corrected log fold change of the sgRNA

The second data frame (segments) contains the identified region of estimated equal log fold changes (one region per row) and the following columns/headers:

- CHR: the chromosome of the gene targeted by the sgRNA under consideration;
- starttp: the genomic coordinate of the starting position of the region targeted by the sgRNA under consideration;
- endtp: the genomic coordinate of the ending position of the region targeted by the sgRNA under consideration;
- genes: the HGNC symbol of the gene targeted by the sgRNA under consideration;
- avgFC: the log fold change of the sgRNA averaged across replicates;
- correction: the type of correction: 1 = increased log fold change, -1 = decreased log fold change. 0 indicates no correction;
- correctedFC: the corrected log fold change of the sgRNA

The second data frame (segments) contains the identified region of estimated equal log fold changes (one region per row) and the following columns/headers:

- CHR: the chromosome of the region under consideration;
- starttp: the genomic coordinate of the starting position of the region under consideration;
- endtp: the genomic coordinate of the ending position of the region under consideration;
- n. sgRNAs: the number of sgRNAs targeting sequences in the region under consideration;
- avg.logFC: the average log fold change of the sgRNAs in the region;
- guideIdx: the indexes range of the sgRNAs targeting the region under consideration as they appear in the gwSortedFCs provided in input.

The string of vectors (SORTED_sgRNAs) contains the sgRNAs' identifiers in the same order as they are reported in the gwSortedFCs data frame inputted to the ccr.GWclean function.

Examples

```
data(EPLC.272HcorrectedFCs)
head(EPLC.272HcorrectedFCs$corrected_logFCs)
head(EPLC.272HcorrectedFCs$segments)
head(EPLC.272HcorrectedFCs$SORTED_sgRNAs)
```

`EssGenes.DNA_REPLICATION_cons`*Core Fitness essential genes involved in DNA replication*

Description

List of core fitness essential genes involved in DNA replication assembled by merging together multiple DNA replication signatures from MSigDB [1] as detailed in [2].

Usage

```
data("EssGenes.DNA_REPLICATION_cons")
```

Format

A vector of strings containing HGNC symbols.

References

[1] Subramanian, A., Tamayo, P., Mootha, V. K., Mukherjee, S., Ebert, B. L., Gillette, M. A., et al. (2005). Gene set enrichment analysis: a knowledge-based approach for interpreting genome-wide expression profiles. *Proceedings of the National Academy of Sciences of the United States of America*, 102(43), 15545-15550. <http://doi.org/10.1073/pnas.0506580102>

[2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

Examples

```
data(EssGenes.DNA_REPLICATION_cons)
head(EssGenes.DNA_REPLICATION_cons)
```

`EssGenes.KEGG_rna_polymerase`*Core Fitness essential rna polymerase genes*

Description

List of core fitness essential rna polymerase genes downloaded from MSigDB [1].

Usage

```
data("EssGenes.KEGG_rna_polymerase")
```

Format

A vector of strings containing HGNC symbols.

References

- [1] Subramanian, A., Tamayo, P., Mootha, V. K., Mukherjee, S., Ebert, B. L., Gillette, M. A., et al. (2005). Gene set enrichment analysis: a knowledge-based approach for interpreting genome-wide expression profiles. *Proceedings of the National Academy of Sciences of the United States of America*, 102(43), 15545-15550. <http://doi.org/10.1073/pnas.0506580102>
- [2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

Examples

```
data(EssGenes.KEGG_rna_polymerase)
head(EssGenes.KEGG_rna_polymerase)
```

```
EssGenes.PROTEASOME_cons
```

Core Fitness essential proteasome genes

Description

List of core fitness essential proteasome genes assembled by merging together multiple DNA replication signatures from MSigDB [1] as detailed in [2].

Usage

```
data("EssGenes.PROTEASOME_cons")
```

Format

A vector of strings containing HGNC symbols.

References

- [1] Subramanian, A., Tamayo, P., Mootha, V. K., Mukherjee, S., Ebert, B. L., Gillette, M. A., et al. (2005). Gene set enrichment analysis: a knowledge-based approach for interpreting genome-wide expression profiles. *Proceedings of the National Academy of Sciences of the United States of America*, 102(43), 15545-15550. <http://doi.org/10.1073/pnas.0506580102>
- [2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

Examples

```
data(EssGenes.PROTEASOME_cons)
head(EssGenes.PROTEASOME_cons)
```

`EssGenes.ribosomalProteins`*Core Fitness essential genes coding for ribosomal proteins*

Description

List of core fitness essential coding for ribosomal proteins curated from [1].

Usage

```
data("EssGenes.KEGG_rna_polymerase")
```

Format

A vector of strings containing HGNC symbols.

References

- [1] Yoshihama, M. et al. The human ribosomal protein genes: sequencing and comparative analysis of 73 genes. *Genome Res.* 12, 379-390 (2002)
- [2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

Examples

```
data(EssGenes.ribosomalProteins)
head(EssGenes.ribosomalProteins)
```

`EssGenes.SPLICEOSOME_cons`*Core Fitness essential spliceosome genes*

Description

List of core fitness essential spliceosome genes assembled by merging together multiple DNA replication signatures from MSigDB [1] as detailed in [2].

Usage

```
data("EssGenes.SPLICEOSOME_cons")
```

Format

A vector of strings containing HGNC symbols.

References

- [1] Subramanian, A., Tamayo, P., Mootha, V. K., Mukherjee, S., Ebert, B. L., Gillette, M. A., et al. (2005). Gene set enrichment analysis: a knowledge-based approach for interpreting genome-wide expression profiles. *Proceedings of the National Academy of Sciences of the United States of America*, 102(43), 15545-15550. <http://doi.org/10.1073/pnas.0506580102>
- [2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

Examples

```
data(EssGenes.SPLICEOSOME_cons)
head(EssGenes.SPLICEOSOME_cons)
```

| | |
|--------------------|--|
| GDSC.CL_annotation | <i>Tissue type and other annotations for 1,001 human cancer cell lines</i> |
|--------------------|--|

Description

Tissue type and other annotations for 1,001 human cancer cell lines

Usage

```
data(GDSC.CL_annotation)
```

Format

A data frame with 1,001 observations of the following 7 variables.

CL.name Cell line name;
 COSMIC.ID Cosmic identifier of the cell line;
 GDSC.description_1 Tissue descriptor (Genomics of Drug Sensitivity in Cancer - Level 1);
 GDSC.description_2 Tissue descriptor (Genomics of Drug Sensitivity in Cancer - Level 2);
 'TCGA type' Manually curated matched TCGA cancer type;
 MMR Microsatellite instability status (MSI-S = Stable, MSI-L = Instable, MSI-H = highly-Instable).

Source

This data frame has been derived from the xls table available at http://www.cancerrxgene.org/gdsc1000/GDSC1000_WebResources/Data/suppData/TableS1E.xlsx.

References

- [1] Iorio F, Knijnenburg TA, Vis DJ, Bignell GR, Menden MP, et al. A landscape of pharmacogenomic interactions in cancer *Cell* 2016 Jul 28;166(3):740-54

Examples

```
data(GDSC.CL_annotation)
head(GDSC.CL_annotation)
```

GDSC.geneLevCNA

*Genome-wide copy number data for 15 human cancer cell lines.***Description**

Genome-wide copy number data derived from PICNIC analysis of Affymetrix SNP6 segmentation data (EGAS00001000978, part of the Genomics of Drug Sensitivity in 1,000 Cancer Cell Lines (GDSC1000) panel [1]) for 15 cell lines used in [2] to assess CRISPRcleanerR results.

Usage

```
data(GDSC.geneLevCNA)
```

Format

A data frame with HGNC gene symbols on the row cancer cell lines' cosmic identifiers on the columns. The entry in position i,j indicates the copy number status of gene i in cell line j .

Details

Each entry of the data frame is a string made of four comma separated pieces of data (n1 , n2 , n3 , n4), hyphen (-) is used when the corresponding data is unknown.

The four values indicate:

- n1: Maximum copy number of any genomic segment containing coding sequence of the gene (-1 indicates a value could not be assigned).
- n2: Minimum copy number of any genomic segment containing coding sequence of the gene (-1 indicates a value could not be assigned).
- n3: Zygoty - (H) if all segments containing gene sequence are heterozygous, (L) if any segment containing coding sequence has LOH, (0) if the complete coding sequence of the gene falls within a homozygous deletion.
- n4: Disruption (D) if the gene spans more than 1 genomic segment (-) if no disruption occurs.

Source

This data frame has been derived from the xls table available at ftp://ftp.sanger.ac.uk/pub/project/cancerrxgene/releases/release-6.0/Gene_level_CN.xlsx.

References

- [1] Iorio F, Knijnenburg TA, Vis DJ, Bignell GR, Menden MP, et al. *A landscape of pharmacogenomic interactions in cancer* Cell 2016 Jul 28;166(3):740-54
- [2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>

Examples

```
data(GDSC.geneLevCNA)
GDSC.geneLevCNA[1:10,1:10]
```

| | |
|-----------------|--|
| KY_Library_v1.0 | <i>sgRNAs' genome-wide annotation for the Sanger sgRNA pooled Library v1.0</i> |
|-----------------|--|

Description

A data frame with a named row for each sgRNA of the Sanger sgRNA pooled library presented in [1] including annotations such as targeted genes, and genomic coordinates.

Usage

```
data("KY_Library_v1.0")
```

Format

A a row named data frame with 90709 observations (one for each sgRNA) of the following 7 variables.

CODE alphanumeric identifier of the sgRNAs;

GENES targeted gene;

EXONE exon of the targeted genomic region (string with 'ex' prefix followed by the exon number);

CHRM chromosome of where the targeted region resides (string)

STRAND targeted DNA strand ('+' or '-')

STARTpos starting genomic coordinate of the targeted genomic region (numeric);

ENDpos ending genomic coordinate of the targeted genomic region (numeric).

References

[1] Tzelepis K, Koike-Yusa H, De Braekeleer E, Li Y, Metzakopian E, Dovey OM, Mupo A, Grinkevich V, Li M, Mazan M, Gozdecka M, Onishi S, Cooper J, Patel M, McKerrell T, Chen B, Domingues AF, Gallipoli P, Teichmann S, Ponstingl H, McDermott U, Saez-Rodriguez J, Huntly BJP, Iorio F, Pina C, Vassiliou GS, Yusa K. *A CRISPR dropout screen identifies genetic vulnerabilities and therapeutic targets in acute myeloid leukaemia*. Cell Reports 2016 Oct 18;17(4):1193-1205

Examples

```
data(KY_Library_v1.0)
head(KY_Library_v1.0)
```

| | |
|--------------|--|
| RNAseq.fpkms | <i>RNAseq derived genome-wide basal expression profiles for 15 cell lines.</i> |
|--------------|--|

Description

Genome-wide substitute reads with fragments per kilobase of exon per million reads mapped (FPKM) for the 15 cell lines specified in [CL.subset](#), derived from a comprehensive collection of RNAseq profiles described in [1] and used in [2] to assess CRISPRcleaner results.

Usage

```
data(RNAseq.fpkms)
```

Format

A data frame with one observations per gene and one variable per cell line. Row names indicates HGNC symbols and column names indicate cell line COSMIC identifiers [3].

References

- [1] Garcia-Alonso L, Iorio F, Matchan A, et al. *Transcription factor activities enhance markers of drug response in cancer* doi: <https://doi.org/10.1101/129478>
- [2] Iorio, F., Behan, F. M., Goncalves, E., Beaver, C., Ansari, R., Pooley, R., et al. (n.d.). Unsupervised correction of gene-independent cell responses to CRISPR-Cas9 targeting. <http://doi.org/10.1101/228189>
- [3] Forbes SA, Beare D, Boutselakis H, et al. *COSMIC: somatic cancer genetics at high-resolution* Nucleic Acids Research, Volume 45, Issue D1, 4 January 2017, Pages D777-D783,

See Also

[CL.subset](#)

Examples

```
data(RNAseq.fpkms)
head(RNAseq.fpkms)
```

Index

*Topic **Assessment and Visualisation**

ccr.PrecisionRecallCurve, [21](#)
ccr.VisDepAndSig, [23](#)

*Topic **analysis**

ccr.cleanChrm, [4](#)
ccr.correctCounts, [7](#)
ccr.GWclean, [15](#)
ccr.logFCs2chromPos, [18](#)
ccr.NormfoldChanges, [19](#)

*Topic **datasets**

BAGEL_essential, [2](#)
BAGEL_nonEssential, [3](#)
CCLE.gisticCNA, [3](#)
CL.subset, [25](#)
EPLC.272HcorrectedFCs, [25](#)
EssGenes.DNA_REPLICATION_cons, [27](#)
EssGenes.KEGG_rna_polymerase, [27](#)
EssGenes.PROTEASOME_cons, [28](#)
EssGenes.ribosomalProteins, [29](#)
EssGenes.SPLICEOSOME_cons, [29](#)
GDSC.CL_annotation, [30](#)
GDSC.geneLevCNA, [31](#)
KY_Library_v1.0, [32](#)
RNAseq.fpkms, [33](#)

*Topic **utils**

ccr.geneMeanFCs, [9](#)
ccr.genes2sgRNAs, [10](#)
ccr.get.CCLEgisticSets, [11](#)
ccr.get.gdsc1000.AMPgenes, [12](#)
ccr.get.nonExpGenes, [13](#)

BAGEL_essential, [2](#), [3](#), [22](#)
BAGEL_nonEssential, [2](#), [3](#), [22](#)

CCLE.gisticCNA, [3](#), [11](#)
ccr.cleanChrm, [4](#), [17](#)
ccr.correctCounts, [7](#)
ccr.geneMeanFCs, [9](#)
ccr.genes2sgRNAs, [10](#), [22](#)
ccr.get.CCLEgisticSets, [11](#), [13](#)
ccr.get.gdsc1000.AMPgenes, [12](#), [12](#), [14](#)
ccr.get.nonExpGenes, [13](#)
ccr.GWclean, [8](#), [15](#), [25](#)
ccr.logFCs2chromPos, [5](#), [6](#), [15](#), [18](#)

ccr.NormfoldChanges, [5](#), [6](#), [8](#), [15](#), [18](#), [19](#), [19](#)
ccr.PrecisionRecallCurve, [21](#), [24](#)
ccr.VisDepAndSig, [22](#), [23](#)
CL.subset, [3](#), [25](#), [33](#)

EPLC.272HcorrectedFCs, [25](#)
EssGenes.DNA_REPLICATION_cons, [27](#)
EssGenes.KEGG_rna_polymerase, [27](#)
EssGenes.PROTEASOME_cons, [28](#)
EssGenes.ribosomalProteins, [29](#)
EssGenes.SPLICEOSOME_cons, [29](#)

GDSC.CL_annotation, [30](#)
GDSC.geneLevCNA, [12](#), [31](#)

KY_Library_v1.0, [9](#), [10](#), [19–21](#), [32](#)

RNAseq.fpkms, [14](#), [33](#)