Package 'ASCAT'

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| Type Package | |
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| Title Allele-Specific Copy Number Analysis of Tumours | |
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| Description R package of ASCAT as published in http://www.ncbi.nlm.nih.gov/pubmed/20837533 | |
| Depends R (>= 2.13.0), RColorBrewer | |
| License GPL-3 | |
| LazyLoad yes | |
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| ascat.aspcf ascat.aspcf | _ |

Description

run ASPCF segmentation

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Usage

```
ascat.aspcf(ASCATobj, selectsamples = 1:length(ASCATobj$samples),
  ascat.gg = NULL, penalty = 25)
```

Arguments

ASCATobj an ASCAT object

 $selects ample \ \ a \ vector \ containing \ the \ sample \ number(s) \ to \ PCF. \ Default = all$

ascat.gg germline genotypes (NULL if germline data is available)

penalty penalty of introducing an additional ASPCF breakpoint (expert parameter, don't

adapt unless you know what you're doing)

Details

This function can be easily parallelised by controlling the selectsamples parameter it saves the results in LogR_PCFed[sample]_[segment].txt and BAF_PCFed[sample]_[segment].txt; if these files exist, the results are read from them.

Hence, after parallelisation, copy all the files into the current directory, and run this function to read in the results

Value

output: ascat data structure containing:

- 1. Tumor_LogR data matrix
- 2. Tumor_BAF data matrix
- 3. Tumor_LogR_segmented: matrix of LogR segmented values
- 4. Tumor_BAF_segmented: list of BAF segmented values; each element in the list is a matrix containing the segmented values for one sample (only for probes that are germline homozygous)
- 5. Germline_LogR data matrix
- 6. Germline_BAF data matrix
- 7. SNPpos: position of all SNPs
- 8. ch: a list containing vectors with the indices for each chromosome (e.g. Tumor_LogR[ch[[13]],] will output the Tumor_LogR data of chromosome 13
- 9. chr: a list containing vectors with the indices for each distinct part that can be segmented separately (e.g. chromosome arm, stretch of DNA between gaps in the array design)

ascat.GCcorrect ascat.GCcorrect

Description

Corrects logR of the tumour sample(s) with genomic GC content

Usage

```
ascat.GCcorrect(ASCATobj, GCcontentfile = NULL)
```

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Arguments

ASCATobj an ASCAT object

GCcontentfile File containing the GC content around every SNP for increasing window sizes

Details

Note that probes not present in the GCcontentfile will be lost from the results

Value

ASCAT object with corrected tumour logR

ascat.loadData

ascat.loadData

Description

Function to read in SNP array data

Usage

```
ascat.loadData(Tumor_LogR_file, Tumor_BAF_file, Germline_LogR_file = NULL,
  Germline_BAF_file = NULL, chrs = c(1:22, "X", "Y"), gender = NULL,
  sexchromosomes = c("X", "Y"))
```

Arguments

```
Tumor_LogR_file
```

file containing logR of tumour sample(s)

Tumor_BAF_file file containing BAF of tumour sample(s)

Germline_LogR_file

file containing logR of germline sample(s), NULL

Germline_BAF_file

file containing BAF of germline sample(s), NULL

chrs a vector containing the names for the chromosomes (e.g. c(1:22,"X"))

gender a vector of gender for each cases ("XX" or "XY"). Default = all female ("XX")

sexchromosomes a vector containing the names for the sex chromosomes

Details

germline data files can be NULL - in that case these are not read in

Value

ascat data structure containing:

- 1. Tumor_LogR data matrix
- 2. Tumor_BAF data matrix
- 3. Tumor_LogR_segmented: placeholder, NULL
- 4. Tumor_BAF_segmented: placeholder, NULL
- 5. Germline_LogR data matrix
- 6. Germline_BAF data matrix
- 7. SNPpos: position of all SNPs
- 8. ch: a list containing vectors with the indices for each chromosome (e.g. Tumor_LogR[ch[[13]],] will output the Tumor_LogR data of chromosome 13
- 9. chr: a list containing vectors with the indices for each distinct part that can be segmented separately (e.g. chromosome arm, stretch of DNA between gaps in the array design)
- 10. gender: a vector of gender for each cases ("XX" or "XY"). Default = NULL: all female ("XX")

ascat.plotAscatProfile

ascat.plotAscatProfile

Description

ascat.plotAscatProfile

Usage

ascat.plotAscatProfile(n1all, n2all, heteroprobes, ploidy, rho, goodnessOfFit, nonaberrant, y_limit, nAfull, ch, lrr, bafsegmented, chrs, textFlag)

Arguments

n1all copy number major allele n2all copy number minor allele

heteroprobes probes with heterozygous germline

ploidy ploidy of the sample rho purity of the sample goodnessOfFit estimated goodness of fit

nonaberrant

y_limit Optional parameter determining the size of the y axis in the nonrounded plot and

ASCAT profile. Default=5

nAfull copy number major allele

ch a list containing c vectors, where c is the number of chromosomes and every

vector contains all probe numbers per chromosome

1rr (unsegmented) log R, in genomic sequence (all probes), with probe IDs

bafsegmented B Allele Frequency, segmented, in genomic sequence (only probes heterozygous

in germline), with probe IDs

chrs a vector containing the names for the chromosomes (e.g. c(1:22,"X"))

textFlag Optional flag to add the positions of fragments located outside of the plotting

area to the plots. Default=F

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Value

plot showing the ASCAT profile of the sample

ascat.plotNonRounded ascat.plotNonRounded

Description

ascat.plotNonRounded

Usage

```
ascat.plotNonRounded(ploidy, rho, goodnessOfFit, nonaberrant, nAfull, nBfull,
  y_limit, ch, bafsegmented, lrr, chrs, textFlag)
```

Arguments

ploidy ploidy of the sample
rho purity of the sample

goodnessOfFit estimated goodness of fit

nonaberrant

nAfull copy number major allele nBfull copy number minor allele

y_limit Optional parameter determining the size of the y axis in the nonrounded plot and

ASCAT profile. Default=5

ch a list containing c vectors, where c is the number of chromosomes and every

vector contains all probe numbers per chromosome

bafsegmented B Allele Frequency, segmented, in genomic sequence (only probes heterozygous

in germline), with probe IDs

1rr (unsegmented) log R, in genomic sequence (all probes), with probe IDs

chrs a vector containing the names for the chromosomes (e.g. c(1:22,"X"))

textFlag Optional flag to add the positions of fragments located outside of the plotting

area to the plots. Default=F

Value

plot showing the nonrounded copy number profile

ascat.plotRawData

ascat.plotRawData

Description

Plots SNP array data

Usage

```
ascat.plotRawData(ASCATobj)
```

Arguments

ASCATobj

an ASCAT object (e.g. data structure from ascat.loadData)

Value

Produces png files showing the logR and BAF values for tumour and germline samples

```
ascat.plotSegmentedData
```

ascat.plot Segmented Data

Description

plots the SNP array data before and after segmentation

Usage

```
ascat.plotSegmentedData(ASCATobj)
```

Arguments

ASCATobj

an ASCAT object (e.g. from ascat.aspcf)

Value

png files showing raw and segmented tumour logR and BAF

```
as cat. predict Germline Genotypes \\ as cat. predict Germline Genotypes
```

Description

predicts the germline genotypes of samples for which no matched germline sample is available

Usage

```
ascat.predictGermlineGenotypes(ASCATobj, platform = "AffySNP6")
```

Arguments

ASCATobj an ASCAT object
platform used array platform

Details

Currently possible values for platform:

AffySNP6 (default)

Custom10k

Illumina109k

IlluminaCytoSNP

Illumina610k

Illumina660k

Illumina700k

Illumina1M

Illumina2.5M

IlluminaOmni5

Affy10k

Affy100k

Affy250k_sty

Affy250k_nsp

AffyOncoScan

Affy CytoScanHD

Value

predicted germline genotypes

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Description

ASCAT main function, calculating the allele-specific copy numbers

Usage

```
ascat.runAscat(ASCATobj, gamma = 0.55, pdfPlot = F, y_limit = 5,
  textFlag = F, rho_manual = NA, psi_manual = NA)
```

Arguments

| ASCATobj | an ASCAT object from ascat.aspcf |
|------------|---|
| gamma | technology parameter, compaction of Log R profiles (expected decrease in case of deletion in diploid sample, 100% aberrant cells; 1 in ideal case, 0.55 of Illumina $109K$ arrays) |
| pdfPlot | Optional flag if nonrounded plots and ASCAT profile in pdf format are desired. Default=F |
| y_limit | Optional parameter determining the size of the y axis in the nonrounded plot and ASCAT profile. Default=5 |
| textFlag | Optional flag to add the positions of fragments located outside of the plotting area to the plots. Default=F |
| rho_manual | optional argument to override ASCAT optimization and supply rho parameter (not recommended) |
| psi_manual | optional argument to override ASCAT optimization and supply psi parameter (not recommended) |

Details

Note: for copy number only probes, nA contains the copy number value and nB = 0.

Value

an ASCAT output object, containing:

- 1. nA: copy number of the A allele
- 2. nB: copy number of the B allele
- 3. aberrantcellfraction: the aberrant cell fraction of all arrays
- 4. ploidy: the ploidy of all arrays
- 5. failedarrays: arrays on which ASCAT analysis failed
- 6. nonaberrantarrays: arrays on which ASCAT analysis indicates that they show virtually no aberrations
- 7. segments: an array containing the copy number segments of each sample (not including failed arrays)
- 8. segments_raw: an array containing the copy number segments of each sample without any rounding applied

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Description

the ASCAT main function

Usage

```
runASCAT(lrr, baf, lrrsegmented, bafsegmented, gender, SNPpos, chromosomes,
  chrnames, sexchromosomes, failedqualitycheck = F, distancepng = NA,
  copynumberprofilespng = NA, nonroundedprofilepng = NA,
  aberrationreliabilitypng = NA, gamma = 0.55, rho_manual = NA,
  psi_manual = NA, pdfPlot = F, y_limit = 5, textFlag = F)
```

Arguments

1rr (unsegmented) log R, in genomic sequence (all probes), with probe IDs

baf (unsegmented) B Allele Frequency, in genomic sequence (all probes), with probe

IDs

1rrsegmented log R, segmented, in genomic sequence (all probes), with probe IDs

bafsegmented B Allele Frequency, segmented, in genomic sequence (only probes heterozygous

in germline), with probe IDs

SNPpos position of all SNPs

chromosomes a list containing c vectors, where c is the number of chromosomes and every

vector contains all probe numbers per chromosome

chrnames a vector containing the names for the chromosomes (e.g. c(1:22,"X"))

sexchromosomes a vector containing the names for the sex chromosomes

failedqualitycheck

did the sample fail any previous quality check or not?

distancepng if NA: distance is plotted, if filename is given, the plot is written to a .png file copynumberprofilespng

if NA: possible copy number profiles are plotted, if filename is given, the plot is written to a .png file

nonroundedprofilepng

if NA: copy number profile before rounding is plotted (total copy number as well as the copy number of the minor allele), if filename is given, the plot is written to a .png file

aberrationreliabilitypng

if NA: aberration reliability score is plotted, if filename is given, the plot is

written to a .png file

gamma technology parameter, compaction of Log R profiles (expected decrease in case

of deletion in diploid sample, 100% aberrant cells; 1 in ideal case, 0.55 of Illu-

mina 109K arrays)

rho_manual optional argument to override ASCAT optimization and supply rho parameter

(not recommended)

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| psi_manual | optional argument to override ASCAT optimization and supply psi parameter (not recommended) |
|------------|--|
| pdfPlot | Optional flag if nonrounded plots and ASCAT profile in pdf format are desired. Default=F |
| y_limit | Optional parameter determining the size of the y axis in the nonrounded plot and ASCAT profile. Default=5 |
| textFlag | Optional flag to add the positions of fragments located outside of the plotting area to the plots. Default=F |
| gender: | a vector of gender for each cases ("XX" or "XY"). Default = NULL: all female ("XX") |

Value

a list containing optimal purity and ploidy

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