

# Package ‘demuxSNP’

May 1, 2024

**Title** scRNAseq demultiplexing using cell hashing and SNPs

**Version** 1.2.0

## Description

This package assists in demultiplexing scRNAseq data using both cell hashing and SNPs data. The SNP profile of each group is learned using high confidence assignments from the cell hashing data.

Cells which cannot be assigned with high confidence from the cell hashing data are assigned to their most similar group based on their SNPs.

We also provide some helper function to optimise SNP selection, create training data and merge SNP data into the SingleCellExperiment framework.

**URL** <https://github.com/michaelplynych/demuxSNP>

**BugReports** <https://github.com/michaelplynych/demuxSNP/issues>

**License** GPL-3

**Encoding** UTF-8

**Roxygen** list(markdown = TRUE)

**RoxygenNote** 7.3.1

**Depends** R (>= 4.3.0), SingleCellExperiment, VariantAnnotation, ensemblDb

**Imports** MatrixGenerics, BiocGenerics, class, GenomeInfoDb, IRanges, Matrix, SummarizedExperiment, demuxmix, methods, KernelKnn, dplyr

**Suggests** knitr, rmarkdown, ComplexHeatmap, viridisLite, ggpubr, dittoSeq, EnsDb.Hsapiens.v86, BiocStyle, RefManageR, testthat (>= 3.0.0), Seurat

**biocViews** Classification, SingleCell

**VignetteBuilder** knitr

**LazyData** false

**Config/testthat/edition** 3

**git\_url** <https://git.bioconductor.org/packages/demuxSNP>

**git\_branch** RELEASE\_3\_19

**git\_last\_commit** 197fc63  
**git\_last\_commit\_date** 2024-04-30  
**Repository** Bioconductor 3.19  
**Date/Publication** 2024-04-30  
**Author** Michael Lynch [aut, cre] (<<https://orcid.org/0000-0001-9535-6461>>),  
Aedin Culhane [aut] (<<https://orcid.org/0000-0002-1395-9734>>)  
**Maintainer** Michael Lynch <michael.lynch@ul.ie>

Contents

add_snps . . . . .	2
common_genes . . . . .	3
commonvariants_1kgenomes_subset . . . . .	3
high_conf_calls . . . . .	4
multiplexed_scrnaseq_sce . . . . .	5
reassign . . . . .	5
reassign_balanced . . . . .	6
reassign_jaccard . . . . .	7
subset_vcf . . . . .	8
vartrix_consensus_snps . . . . .	9

<b>Index</b>	<b>10</b>
--------------	-----------

---

add_snps	<i>Add SNPs to SingleCellExperiment object</i>
----------	--

---

Description

Add SNPs to SingleCellExperiment object

Usage

add\_snps(sce, mat, thresh = 0.8)

Arguments

- sce                    object of class SingleCellExperiment
- mat                    object of class matrix, output from VarTrix in 'consensus' mode (default)
- thresh                threshold presence of SNP, defaults to 0.8

Value

Updated SingleCellExperiment object with snps in altExp slot

**Examples**

```
data(multiplexed_scrnaseq_sce, vartrix_consensus_snps)
multiplexed_scrnaseq_sce <- add_snps(sce = multiplexed_scrnaseq_sce,
mat = vartrix_consensus_snps,
thresh = 0.8)
```

---

common_genes	<i>Return a character vector of top n most frequent genes from a Single-CellExperiment object.</i>
--------------	--

---

**Description**

Returns a character vector of the top n most frequently expressed genes from the counts of the SingleCellExperiment object. Expression is based on having a count > 0 in a given cell.

**Usage**

```
common_genes(sce, n = 100)
```

**Arguments**

sce	a SingleCellExperiment object
n	number of genes to be returned. Defaults to n=100.

**Value**

character vector of n most frequently expressed genes.

**Examples**

```
data(multiplexed_scrnaseq_sce)
multiplexed_scrnaseq_sce <- common_genes(multiplexed_scrnaseq_sce)
```

---

commonvariants_1kgenomes_subset	<i>Sample vcf file</i>
---------------------------------	------------------------

---

**Description**

VCF file containing SNPs from a subset of the 1k Genomes common variants HG38 genome build.

**Usage**

```
data(commonvariants_1kgenomes_subset)
```

**Format**

An object of class CollapsedVCF with 2609 rows and 0 columns.

**Value**

commonvariants\_1kgenomes\_subset:  
An object of class CollapsedVcf

**Source**

[https://cellsnp-lite.readthedocs.io/en/latest/snp\\_list.html](https://cellsnp-lite.readthedocs.io/en/latest/snp_list.html)

---

high_conf_calls	<i>Run demuxmix to determine high-confidence calls</i>
-----------------	--

---

**Description**

Run demuxmix to determine high-confidence calls

**Usage**

```
high_conf_calls(sce, assay = "HTO", pacpt = 0.95)
```

**Arguments**

sce	Object of class SingleCellExperiment with HTO (or similar) altExp assay
assay	Name of altExp for cell hashing counts to be retrieved from
pacpt	acceptance probability for demuxmix model

**Value**

Updated SingleCellExperiment object with logical vector indicating training data, data to be classified (all cells) and assigned labels for all cells.

**Examples**

```
data(multiplexed_scrnaseq_sce)
multiplexed_scrnaseq_sce <- high_conf_calls(multiplexed_scrnaseq_sce)
```

---

multiplexed\_scrnaseq\_sce

*SingleCellExperiment object containing multiplexed RNA and HTO data from six biological smamples*

---

## Description

Example SingleCellExperiment object containing demultiplexed scRNAseq data from six donors, used throughout and built upon in demuxSNP workflow.

## Usage

```
data(multiplexed_scrnaseq_sce)
```

## Format

An object of class SingleCellExperiment with 259 rows and 2000 columns.

## Value

multiplexed\_scrnaseq\_sce:  
An object of class SingleCellExperiment

---

reassign

*Reassign cells using knn*

---

## Description

k-nearest neighbour classification of cells. Training data is intended to be labels of cells confidently called using cell hashing based methods and their corresponding SNPs. Prediction data can be remaining cells but can also include the training data. Doublets are simulated by randomly combining 'd' SNP profiles from each grouping combination.

## Usage

```
reassign(
  sce,
  k = 10,
  d = 10,
  train_cells = sce$train,
  predict_cells = sce$predict
)
```

**Arguments**

sce                    object of class SingleCellExperiment  
 k                     number of neighbours used in knn, defaults to 10  
 d                     number of doublets per group combination to simulate, defaults to 10  
 train\_cells         logical vector specifying which cells to use to train classifier  
 predict\_cells       logical vector specifying which cells to classify

**Value**

A SingleCellExperiment with updated group assignments called 'knn'

**Examples**

```
data(multiplexed_scrnaseq_sce, vartrix_consensus_snps)
multiplexed_scrnaseq_sce <- high_conf_calls(multiplexed_scrnaseq_sce)
multiplexed_scrnaseq_sce <- add_snps(sce = multiplexed_scrnaseq_sce,
  mat = vartrix_consensus_snps,
  thresh = 0.8)
multiplexed_scrnaseq_sce <- reassign(sce = multiplexed_scrnaseq_sce, k = 10)
```

---

reassign_balanced	<i>Reassign cells using balanced knn with jaccard distance</i>
-------------------	--

---

**Description**

k-nearest neighbour classification of cells. Training data is intended to be labels of cells confidently called using cell hashing based methods and their corresponding SNPs. Prediction data can be remaining cells but can also include the training data. Doublets are simulated by randomly combining 'd' SNP profiles from each grouping combination.

**Usage**

```
reassign_balanced(
  sce,
  k = 10,
  d = 10,
  train_cells = sce$train,
  predict_cells = sce$predict,
  n = NULL,
  nmin = 50
)
```

**Arguments**

sce	object of class SingleCellExperiment
k	number of neighbours used in knn, defaults to 10
d	number of doublets per group combination to simulate, defaults to 10
train_cells	logical vector specifying which cells to use to train classifier
predict_cells	logical vector specifying which cells to classify
n	number of cells per group (otherwise will be calculated from data)
nmin	min n per class (where available)

**Value**

A SingleCellExperiment with updated group assignments called 'knn'

**Examples**

```
data(multiplexed_scrnaseq_sce, vartrix_consensus_snps)
multiplexed_scrnaseq_sce <- high_conf_calls(multiplexed_scrnaseq_sce)
multiplexed_scrnaseq_sce <- add_snps(sce = multiplexed_scrnaseq_sce,
mat = vartrix_consensus_snps,
thresh = 0.8)
multiplexed_scrnaseq_sce <- reassign_balanced(sce = multiplexed_scrnaseq_sce, k = 10)
```

---

reassign_jaccard	<i>Reassign cells using knn with jaccard distance</i>
------------------	---

---

**Description**

k-nearest neighbour classification of cells. Training data is intended to be labels of cells confidently called using cell hashing based methods and their corresponding SNPs. Prediction data can be remaining cells but can also include the training data. Doublets are simulated by randomly combining 'd' SNP profiles from each grouping combination.

**Usage**

```
reassign_jaccard(
  sce,
  k = 10,
  d = 10,
  train_cells = sce$train,
  predict_cells = sce$predict
)
```

**Arguments**

sce	object of class SingleCellExperiment
k	number of neighbours used in knn, defaults to 10
d	number of doublets per group combination to simulate, defaults to 10
train_cells	logical vector specifying which cells to use to train classifier
predict_cells	logical vector specifying which cells to classify

**Value**

A SingleCellExperiment with updated group assignments called 'knn'

**Examples**

```
data(multiplexed_scrnaseq_sce, vartrix_consensus_snps)
multiplexed_scrnaseq_sce <- high_conf_calls(multiplexed_scrnaseq_sce)
multiplexed_scrnaseq_sce <- add_snps(sce = multiplexed_scrnaseq_sce,
mat = vartrix_consensus_snps,
thresh = 0.8)
multiplexed_scrnaseq_sce <- reassign(sce = multiplexed_scrnaseq_sce, k = 10)
```

---

subset\_vcf

---

*Subset common variants vcf file to only SNPs seen in 'top\_genes'*


---

**Description**

Subset common variants vcf file to only SNPs seen in 'top\_genes'

**Usage**

```
subset_vcf(vcf, top_genes, ensdb)
```

**Arguments**

vcf	object of class CollapsedVCF
top_genes	output from 'common_genes' function, alternatively character vector containing custom gene names.
ensdb	object of class EnsDb corresponding to organism, genome of data

**Value**

object of class CollapsedVCF containing subset of SNPs from supplied vcf seen in commonly expressed genes



**Examples**

```
data(multiplexed_scrnaseq_sce, commonvariants_1kgenomes_subset)
top_genes <- common_genes(multiplexed_scrnaseq_sce)
ensdb <- EnsDb.Hsapiens.v86::EnsDb.Hsapiens.v86
small_vcf <- subset_vcf(commonvariants_1kgenomes_subset, top_genes, ensdb)
```

---

vartrix\_consensus\_snps

*Sample VarTrix output*

---

**Description**

A sample output from VarTrix corresponding to the sce SingleCellExperiment objec for a subset of SNPs located in well observed genes.

**Usage**

```
data(vartrix_consensus_snps)
```

**Format**

An object of class matrix (inherits from array) with 2542 rows and 2000 columns.

**Value**

vartrix\_consensus\_snps:  
An object of class matrix

# Index

## \* **datasets**

commonvariants\_1kgenomes\_subset, [3](#)

multiplexed\_scrnaseq\_sce, [5](#)

vartrix\_consensus\_snps, [9](#)

add\_snps, [2](#)

common\_genes, [3](#)

commonvariants\_1kgenomes\_subset, [3](#)

high\_conf\_calls, [4](#)

multiplexed\_scrnaseq\_sce, [5](#)

reassign, [5](#)

reassign\_balanced, [6](#)

reassign\_jaccard, [7](#)

subset\_vcf, [8](#)

vartrix\_consensus\_snps, [9](#)