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🌐 FR-Match: cell type matching for scRNAseq data

📁 In 1 collection

DOI

<https://dx.doi.org/10.17504/protocols.io.bmyfk7tn>

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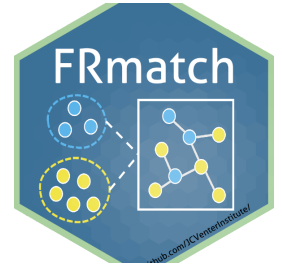
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Human Cell Atlas Metho...



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External link: <https://github.com/JCVenterInstitute/FRmatch>

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Protocol status: Working

We use this protocol and it's working

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Keywords: single cell RNA sequencing, cell types, data integration, cell type matching for scrnaseq data fr, cluster cell transcriptome integration across scrnaseq experiment, scrnaseq data fr, supervised cell phenotype matching strategy for cluster, supervised cell phenotype matching strategy, cell type matching, transcriptome integration, scrnaseq experiment, cluster cell, cluster, match,

Abstract

FR-Match is a supervised cell phenotype matching strategy for cluster-to-cluster cell transcriptome integration across scRNAseq experiments.

An R package and Shiny application are provided at <https://github.com/JCVenterInstitute/FRmatch>.

Before start

Software

R programming language

NAME

The R Foundation

DEVELOPER

[Comprehensive R Archive Network](https://comprehensive-r-network.org/)

REPOSITORY

<https://cran.r-project.org/>

SOURCE LINK

Require R Shiny package.



Launch Shiny app

- 1 Interactively explore and match scRNAseq cell type clusters with the seamless Shiny app. The Shiny app may serve as a quick start demo with pre-loaded datasets. 

Command

new command name

```
runShiny()
```

Data preparation and exploration

- 2 Use the built-in data preparation function to create data objects with required and optional input data elements. Create data objects for experiment 1 (E1) and experiment 2 (E2)

Command

new command name

```
dat_E1 <- make_data_object()  
dat_E2 <- make_data_object()
```

- 2.1 View comparative cell cluster sizes.



Command

new command name

```
plot_clusterSize(dat_E1, dat_E2)
```

2.2 View "barcode" plot for cluster of interest.

Command

new command name

```
plot_cluster_by_markers(dat_E1, cluster.name = "cluster_of_interest")
```

Run main algorithm

10m

3 Use wrapper function to perform bi-directional matching.



3.1 Map E1 to E2.

Command

new command name

```
rst12 <- FRmatch(sce.query = dat_E1, sce.ref = dat_E2)
```



3.2 Map E2 to E1.

Command

new command name

```
rst21 <- FRmatch(sce.query = dat_E2, sce.ref = dat_E1)
```

Combine and plot matching results

4 Combine the bi-directional matching results and plot.

Command

new command name

```
plot_bi_FRmatch(rst12, rst21)
```

Additional plots

5 Some optional plotting functions to help studying the matching results.



5.1 Plot one-directional matching results.

Command

new command name

```
plot_FRmatch(rst12)
```

Plot one-directional matching p-values.

Command

new command name

```
plot_FRmatch(rst12, type = "padj")
```

- 5.2 Minimum spanning tree (MST) plot. MST can be plotted by turning on the plot option in the test function.

Command

new command name

```
FR.test(samp1, samp2, plot.MST = TRUE)
```