Package 'flowCL'

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flowCL-package

Semantic labelling of flow cytometric cell populations.

Description

flowCL uses SPARQL to match a phenotype to a cell type from the cell ontology. If the match is not unique, then the best alternative is returned.

Details

Package: flowCL
Type: Package
License: Artistic-2.0

LazyLoad: yes

Depends: Rgraphviz, SPARQL

flowCL executes queries against the Cell Ontology (CL), available at http://cellontology.org. The CL file is hosted on a triplestore, i.e., a database for storage and retrieval of Resource Description Framework (RDF) triples. The SPARQL endpoint at http://cell.inference.me:7200/repositories/CL (old endpoint: http://cell.ctde.net:8080/openrdf-sesame/repositories/CL) is used to execute the SPARQL queries retrieving the correct matches from the CL. While other SPARQL endpoints can be used, users should be aware that in our case the CL file has been reasoned upon, and resulting extra inferred axioms have been added to the triplestore, providing a more complete result set.

Author(s)

Maintainer: Justin Meskas <jmeskas@bccrc.ca> Authors: Justin Meskas, Radina Droumeva

Examples

```
# Load a pre-loaded archive. Skipping this chuck will cause flowCL to
## slowly build a new one.
flowCL("archive")

# Simple two marker example
Res <- flowCL("CCR7+CD45RA+")
tmp <- Res$'CCR7+CD45RA+'
plot(tmp[[1]], nodeAttrs=tmp[[2]], edgeAttrs=tmp[[3]], attrs=tmp[[4]])
Res$Table

# Exact match example
Res <- flowCL("CD3-CD19-CD20-CD14+", Verbose = TRUE, OntolNamesTD = TRUE)
tmp <- Res$'CD3-CD19-CD20-CD14+'
plot(tmp[[1]], nodeAttrs=tmp[[2]], edgeAttrs=tmp[[3]], attrs=tmp[[4]])
Res$Table

# Cell Label Example
x <-"CD3-CD19-CD20-CD14+"</pre>
```

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```
Res <- flowCL(x)
Res$Cell_Label[[x]][[1]]

# As a secondary way to view the results,
## see "[current directory]/flowCL_results/".
# Figures created called tree_(phenotype).pdf give the cell hierarchy
## dependent on the markers in the phenotype.
# A list of results from Res$Table are stored in listPhenotypes.csv.</pre>
```

flowCL

Semantic labelling of flow cytometric cell populations.

Description

flowCL uses queries to match a phenotype to a cell type from the cell ontology. If the match is not unique, then the best alternative is returned.

Usage

```
flowCL ( MarkerList = "HIPC", ExpMrkrLst = NULL, Indices = NULL,
    Verbose = FALSE, KeepArch = TRUE, MaxHitsPht = 5,
    OntolNamesTD = FALSE, ResetArch = FALSE, VisualSkip = FALSE)
```

Arguments

MarkerList	A list of phenotypes to query the cell ontology with (Ex.	"CD3+CD4-CD8+"
Har Ker E15t	It not of phenotypes to query the een ontology with (Ex.	CD31CD1CD01

as a single element of the list). There is an option to use a preloaded list. This preloaded list can be used by inputting "HIPC" into MarkerList. This will first query all the individual markers, then will query all the common HIPC pheno-

types.

ExpMrkrLst A list of all of the phenotypes that were used in the experiment. This will be

used to inform the user that a certain marker could have been used to further identify the correct population. If ExpMrkLst is left blank, flowCL will define ExpMrkLst as each phenotype in MarkerList. If ExpMrkrLst has only one element, flowCL will define every ExpMrkrLst phenotype as being the same as the

one input.

Indices A vector of indices that dictate which of the MarkerList phenotypes will be

queried. If left blank, all phenotypes in the list will be queried.

Verbose A logical value that dictates if computational information is printed while the

code is running. The default is FALSE.

ResetArch A logical value that dictates if the archive folder, "flowCL_results", is deleted

before the queries take place. This will increase the simulation time, but should be done every once in a while to account for updates from the ontology. The

default is FALSE.

KeepArch A logical value that dictates if the archive folder, "flowCL_results", is deleted

after the queries take place. Set to FALSE to insure there is no unwanted files

being stored on the hard drive. The default is TRUE.

MaxHitsPht An integer for the maximum number of cell types that are returned per pheno-

type queried. The default is 5.

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 ${\tt OntolNamesTD} \qquad A \ logical \ value \ that \ controls \ if \ the \ phenotypes \ in \ the \ tree \ diagrams (TD) \ are \ short$

names (ex. CD4) or the long ontology names (ex. CD4 molecule). Short names are used if OntolNamesTD is FALSE, while long ontology names are used if

TRUE. The default is FALSE.

VisualSkip A logical value that controls if the visualization step is skipped or not. TRUE is

for skip, while FALSE is for no skip. The default is FALSE.

Details

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Value

A list containing N+5 elements. Where N is the number of phenotypes queried. Each of these N elements contains information for plotting the results. The other five elements show the cell labels (Cell_Labels), the matching markers in a list form (Marker_Groups) and a bracket form (Markers), ranking scores (Ranking) and a table (Table). The cell labels element lists the cell labels in order of highest score based on their ranking, which is in a form easily extracted and used by other N packages and functions. Marker_Groups and Markers list markers that were queried and that are part of a certain cell type. In Markers these markers are displayed in the form of N and N by N and N by the markers input for the query. B, C and D together are the markers that make up the definition of the particular cell type. C lists the markers that were pat of the experiment that were not part of the query, while D lists all other markers that make up the cell type that were not part of the experimental markers. A lists all the markers in the input for the query that were not required for that particular cell type. The table is a list of all the related information of each phenotype queried. This table is mainly for users to see the results in N.

Author(s)

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Examples

```
# Load a pre-loaded archive. Skipping this chuck will cause flowCL to
## slowly build a new one.
flowCL("archive")

# Simple two marker example
Res <- flowCL("CCR7+CD45RA+")
tmp <- Res$'CCR7+CD45RA+'
plot(tmp[[1]], nodeAttrs=tmp[[2]], edgeAttrs=tmp[[3]], attrs=tmp[[4]])
Res$Table

# Exact match example
Res <- flowCL("CCR7+CD45RA+CD8+", Verbose = TRUE, OntolNamesTD = TRUE)
tmp <- Res$'CCR7+CD45RA+CD8+'</pre>
```

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```
plot(tmp[[1]], nodeAttrs=tmp[[2]], edgeAttrs=tmp[[3]], attrs=tmp[[4]])
Res$Table

# Cell Label Example
x <-"CCR7+CD45RA+CD8+"
Res <- flowCL(x)
Res$Cell_Label[[x]][[1]]

# As a secondary way to view the results,
## see "[current directory]/flowCL_results/".
# Figures created called tree_(phenotype).pdf give the cell hierarchy
## dependent on the markers in the phenotype.
# A list of results from Res$Table are stored in listPhenotypes.csv.</pre>
```

flowCL_archive

A list of data used to build a local archive.

Description

A list of data used to build a local archive.

Usage

```
data(flowCL_archive)
```

Value

Loads the a pre-built archive.

Author(s)

Justin Meskas

```
test.flowCL.connection
```

A function to check that a connection exists between the user and the ontology server.

Description

A function to check that a connection exists between the user and the ontology server.

Usage

```
test.flowCL.connection()
```

Value

The date of the lastest update of the cell ontology is returned.

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Author(s)

Justin Meskas

Examples

test.flowCL.connection()

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