Alakazam: Reconstruction of Ig lineage trees

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Contents

Example data	1
Preprocess a clone	1
Run PHYLIP	2
Plotting of the lineage tree	3
Batch processing lineage trees	4

Reconstruction of an Ig lineage requires the following steps:

- 1. Load a Change-O tab-delimited database file and select a clone
- 2. Preprocess the clone to remove gap characters and duplicate sequences
- 3. Run PHYLIP, parse the output, and modify the tree topology

Example data

A small example Change-O database, ExampleDb, is included in the alakazam package. Lineage reconstruction requires the following fields (columns) to be present in the Change-O file:

- SEQUENCE ID
- SEQUENCE_IMGT
- CLONE
- GERMLINE_IMGT_D_MASK
- V_CALL
- J_CALL
- JUNCTION_LENGTH

```
# Load required packages
library(alakazam)
library(igraph)
library(dplyr)

# Select clone from example database
data(ExampleDb)
sub db <- subset(ExampleDb, CLONE == 3138)</pre>
```

Preprocess a clone

Before a lineage can be constructed the sequences must first be cleaned of gap (-, .) characters added by IMGT, duplicate sequences must be removed, and annotations must be combined for each cluster of duplicate sequences. Optionally, "ragged" ends of sequences, such as may occur from primer template switching, may also be cleaned by masking mismatched positions and the leading

and trailing ends of each sequence. The function makeChangeoClone is a wrapper function which combines these steps and returns a ChangeoClone object which may then be passed into the lineage reconstruction function.

Two arguments to makeChangeoClone control which annotations are retained following duplicate removal. Unique values appearing within columns given by the text_fields arguments will be concatenated into a single string delimited by a "," character. Values appearing within columns given by the num fields arguments will be summed.

```
# This example data set does not have ragged ends
# Preprocess clone without ragged end masking (default)
clone <- makeChangeoClone(sub_db, text_fields=c("SAMPLE", "ISOTYPE"),</pre>
                           num_fields="DUPCOUNT")
# Show combined annotations
clone@data[, c("SAMPLE", "ISOTYPE", "DUPCOUNT")]
##
     SAMPLE ISOTYPE DUPCOUNT
## 1
        +7d
                 IgA
## 2
        +7d
                 IgG
                            1
## 3
        +7d IgA, IgG
                           10
## 4
        +7d
                 IgG
                           36
## 5
        +7d
                           10
                 IgA
## 6
        +7d
                 IgG
                           13
```

Run PHYLIP

Lineage construction uses the dnapars (maximum parsimony) application of the PHYLIP package. The function buildPhylipLineage performs a number of steps to execute dnapars, parse its output, and modify the tree topology to meet the criteria of an Ig lineage. This function takes as input a ChangeoClone object output by makeChangeoClone and returns an igraph graph object. The igraph graph object will contain clone annotations as graph attributes, sequence annotations as vertex attributes, and mutations along edges as edge attributes.

The system call to dnapars requires a temporary folder to store input and output. This is created in the system temporary location (according to base::tempfile), and is not deleted by default (only because automatically deleting files is somewhat rude). In most cases, you will want to set rm_temp=TRUE to delete this folder.

```
## 1 3138
                        60 IGHV3-49 IGHJ5
# The vertices have sequence specific annotations
data.frame(SEQUENCE_ID=V(graph)$name,
           ISOTYPE=V(graph)$ISOTYPE,
           DUPCOUNT=V(graph)$DUPCOUNT)
##
        SEQUENCE ID ISOTYPE DUPCOUNT
## 1 GN5SHBT06HH3QD
                         IgA
                                   10
## 2 GN5SHBT08F45HV IgA,IgG
                                   10
## 3
           Germline
                        <NA>
                                   NA
## 4 GN5SHBT06IFVOR
                                   13
                         IgG
## 5 GN5SHBT08I3P11
                                   36
                         IgG
## 6 GN5SHBT01BXJY7
                                    1
                         IgG
```

IgA

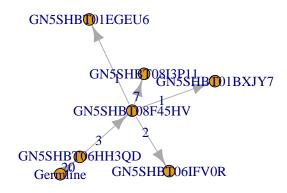
Plotting of the lineage tree

7 GN5SHBT01EGEU6

Plotting of a lineage tree may be done using the built-in functions of the igraph package. The default edge and vertex labels are edge weights and sequence identifiers, respectively.

1

```
# Plot graph with defaults
plot(graph)
```



The default layout and attributes are not very pretty. We can modify the graphical parameter in the usual igraph ways. A tree layout can be built using the layout_as_tree layout with assignment of the root position to the germline sequence, which is named "Germline" in the object returned by buildPhylipLineage.

Modify graph and plot attributes

```
V(graph)$color <- "steelblue"
V(graph)$color[V(graph)$name == "Germline"] <- "black"</pre>
V(graph)$color[grepl("Inferred", V(graph)$name)] <- "white"</pre>
V(graph)$label <- V(graph)$ISOTYPE</pre>
E(graph) $label <- ""
# Remove large default margins
par(mar=c(0, 0, 0, 0) + 0.1)
# Plot graph
plot(graph, layout=layout_as_tree, edge.arrow.mode=0, vertex.frame.color="black",
     vertex.label.color="black", vertex.size=40)
# Add legend
legend("topleft", c("Germline", "Inferred", "Sample"),
       fill=c("black", "white", "steelblue"), cex=0.75)
 ■ Germline
 □ Inferred
 Sample
                                        IgA,IgG
                        IgG
                                    IgG
                                                IgG
                                                           IgA
```

Which is much better.

Batch processing lineage trees

Multiple lineage trees may be generated at once, by splitting the Change-O data.frame on the clone column.