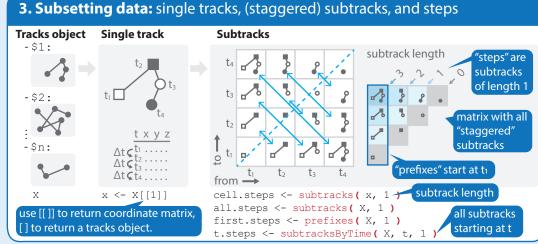
Analysing cell migration data in R CelltrackR cheat sheet

To analyze cell movement, we record a cell's coordinates in time-lapse videos to obtain a cell track. To facilitate the interpretation of tracking data, celltrackR implements a large variety of methods for the fast and flexible analysis of track data in R. Load data from a text file, get rid of artefacts and tracking errors by performing quality controls proposed in literature, and analyze any metric on the level of individual tracks, steps, or subtracks. CelltrackR supports angle analyses and allows rapid visualization, clustering, and simulation of tracks. Let's get started!



1. Loading & converting tracks Generate tracks object from a csv file: mydata.csv: - Scell1: txyz txyz t₁ cell1 t₁ . . . txyz tı tracks object contains a matrix for each cell read.tracks.csv(mydata.csv, id.column = 1, time.column = 2, pos.columns = 3:5) Concatenate two tracks objects: c(X1, X2) Convert between data structures: as.data.frame(X) dataframe tracks to dataframe ID txvz as.tracks(D) cel|1 t₁ dataframe to tracks cell2 t₁ as.list(X) cell2 t2 tracks to regular R list - \$cell1: wrapTrack(x) txyz wrap single track matrix tı into a track object Sort tracks by time-order: sort (X) Output of read. tracks.csv() t₈ . . . t₁ . . . t₃ . . . t₂ . . . and as.tracks.data.frame()

is time-ordered by default.

2. Quality control & preprocessing **Longer tracks** allow better inference of the cell's behavior, especially in cell-based analyses (box 4). hist(sapply(X, nrow)) max length distribution maxTrackLength (X) must return longest track (# steps) TRUE/FALSE filterTracks (function(x) nrow(x)>n, X) keep only tracks of at least n steps Filtering can cause bias. Consider a step-based analysis (box 4) instead of removing short tracks. Check for **unequal** Δt between steps, or gaps: position avdt <- timeStep(x); hist(sapply(</pre> subtracks(x, 1), duration) - avdt) Fix this issue automatically for all tracks in X with an irregular Δt above some threshold: or: "split" fix1 <- repairGaps(X, "interpolate")</pre> subsample every Adjust time resolution Δt : k-th timepoint subsample (x, k = 2) t₁... t₁... t3... interpolateTrack(x, dtvec) interpolate at times in dtvec Angle analyses (**box 6**) can help detect artifacts,

drift, and tracking errors (Beltman et al. 2009).

4. Analysis types: cell-based, step-based, and staggered metrics Track properties can be computed in a cell-based, step-based, or staggered fashion. For more information, please refer to (Beltman et al, 2009). Examples are shown for the analysis of speed, but can also be performed with other analysis measures (box 5). Step-based Cell-based Average speed over all steps, Find average speed of each pooled from all tracks together: individual cell (track): cell-based mean mean cell speed mean(sapply(X, speed) cells have equal weights; steps from short tracks weigh more Get instantaneous/"step" speed distribution for each cell (track): **T**cell 1:

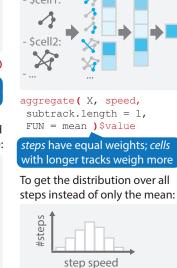
step speed

hist(sapply(

steps, speed)

steps <- subtracks (x, 1)

one cell >



hist(sapply(

steps, speed)

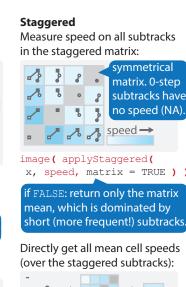
steps <- subtracks (X, 1)

all steps in

object X

sapply(X,

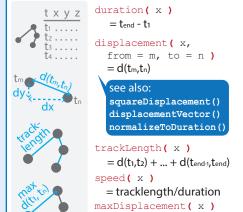
staggered (speed))





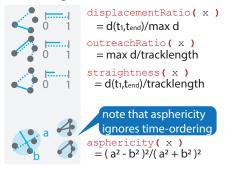
(see also ?TrackMeasures)

Speed and displacement

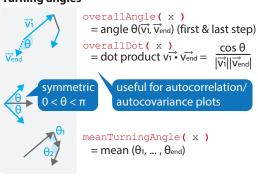


 $= \max d(t_1, t_n)$

Track straightness



Turning angles



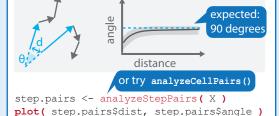
6. Angles & Directionality (see also ?AngleAnalysis)

Angles to a reference point, direction, or plane angleToPoint (x,p) = angle θ between first step and reference point distanceToPoint (x,p) = distance d between first step and reference point angleToDir (x, dvec) = angle θ between first step and reference direction angleToPlane (x,p1,p2,p3) = angle θ between first step and plane with points p1-p3 distanceToPlane (x,p1,p2,p3)

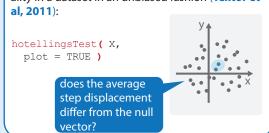
Angles between pairs of steps or tracks can help identify directional biases or artefacts (Beltman et al, 2009):

= distance d between first step

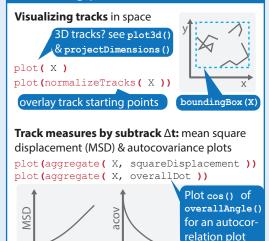
and plane with points p1-p3



Hotelling's test can help detect global directionality in a dataset in an unbiased fashion (Textor et al. 2011):



7. Visualization & Clustering: detecting patterns in track data



Θ

speed

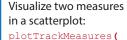
straightness

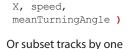
Other methods:

"UMAP"/"MDS"

Or: "kmeans'

Tracks in feature space:

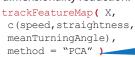




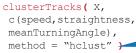
feature first: minv <- median (</pre>

```
minv <- median(
  sapply( X, speed ) )
fast <- selectTracks(
  X, speed, minv, Inf )</pre>
```

Or visualize higher dimensional feature sets with dimensionality reduction:



Cluster tracks by features:



8. Simulating tracks: Models & bootstrapping

Comparing observed data to idealized models is useful for interpretation. CelltrackR supports several methods for simulating tracks.

A **random walk** in dim dimensions:

```
brownianTrack( nsteps, dim, mean=c(0,0), sd=c(1,1)) non-zero for directional bias
```

A "**stop-and-go**" model designed for T cells (**Beauchemin et al, 2007**). Cells move at speed v.free for time t.free, and then pause for a time t.pause before changing direction (can be with directional persistence or directional bias):

```
beaucheminTrack( sim.time, delta.t,
  p.persist, p.bias, bias.dir, taxis.mode,
  t.free, v.free, t.pause )
```

unlike brownianTrack(), beaucheminTrack() has an explicit definition of time.

A **bootstrapped track** matches speeds and turning angles to those observed in data:

bootstrapTrack(nsteps, X)

Simulate multiple tracks at once:

```
simdata <- simulateTracks( 10,
  bootstrapTrack( nsteps, X ) )</pre>
```

or another simulation method

References

Beauchemin et al (2007). Characterizing T cell movement within lymph nodes in the absence of antigen. *Journal of Immunology*.

Beltman et al (2009). Analysing Immune cell migration. Nature Reviews Immunology.

Mokhtari et al (2013). Automated characterization and parameter-free classification of cell tracks based on local migration behavior. *PLoS ONE*.

Textor et al (2007). Defining the quantitative limits of intravital two-photon lymphocyte tracking. *PNAS*.



Learn more?

Check out the detailed examples in the package vignettes: browseVignettes(package = "celltrackR")

© Johannes Textor, Katharina Dannenberg, Jeffrey Berry, Gerhard Burger, Inge Wortel (2019). For the newest version, visit: https://github.com/ingewortel/celltrackR To cite celltrackR, please refer to: citation("celltrackR").