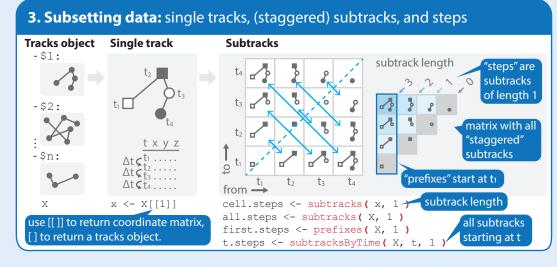
# Analysing cell migration data in R CelltrackR cheat sheet

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 analyse 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!

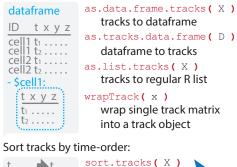


# 1. Loading & converting tracks

### Convert between data structures:

t<sub>8</sub> . . . t<sub>1</sub> . . .

t<sub>3</sub> . . . t<sub>2</sub> . . .



Output of read. tracks.csv()

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**).



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**  $\Delta t$  between steps, which can cause artefacts.



avdt <- timeStep( x ); hist( sapply(
 subtracks( x, 1 ), duration ) - avdt )</pre>

Fix this issue automatically for all tracks in X with an irregular  $\Delta t$  above some threshold: or: "split"

fix1 <- repairGaps( X, "interpolate")</pre>



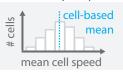
Angle analyses (**box 6**) can help detect artefacts, drift, and tracking errors (**Beltman et al, 2009**).

# **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**).

#### Cell-based

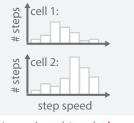
Find average speed of each individual cell (track):



mean( sapply( X, speed ) )

*cells* have equal weights; *steps* from short tracks weigh more

Get instantaneous/"step" speed distribution for each cell (track):



steps <- subtracks( x, 1 )
hist( sapply(
 steps, speed ) )
one cell x</pre>

### Step-based

Average speed over all steps, pooled from all tracks together:



aggregate( X, speed,
 subtrack.length = 1,
 FUN = mean )\$value

steps have equal weights; cells with longer tracks weigh more

To get the distribution over all steps instead of only the mean:



steps <- subtracks( X, 1 )
hist( sapply(
 steps, speed ) )
object X</pre>

### Staggered

Measure speed on all subtracks in the staggered matrix:



image( applyStaggered(
 x, speed, matrix = TRUE ) ;

if FALSE: return only the matrix mean, which is dominated by short (more frequent!) subtracks.

Directly get all mean cell speeds (over the staggered subtracks):

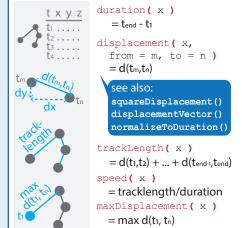


sapply( X,
 staggered( speed ) )

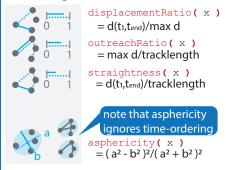


(see also ?TrackMeasures)

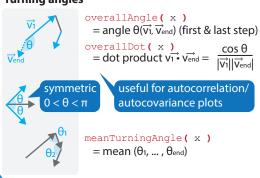
### **Speed and displacement**



### Track straightness



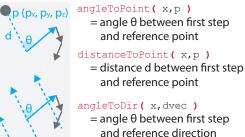
### **Turning angles**



# 6. Angles & Directionality

(see also ?AngleAnalysis)

### Angles to a reference point, direction, or plane

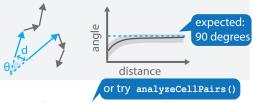


angleToPlane(x,p1,p2,p3)

= angle  $\theta$  between first step and plane with points p1-p3

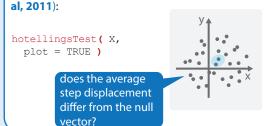
distanceToPlane(x,p1,p2,p3) = distance d between first step and plane with points p1-p3

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

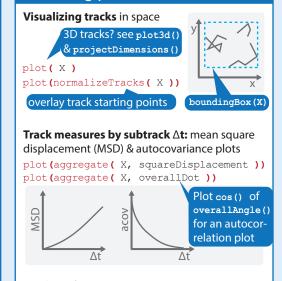


step.pairs <- analyzeStepPairs( X )</pre> plot( step.pairs\$dist, step.pairs\$angle )

# Hotelling's test can help detect global directionality in a dataset in an unbiased fashion (Textor et



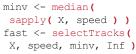
# 7. Visualization & Clustering: detecting patterns in track data



### Tracks in feature space: Visualize two measures in a scatterplot:

```
plotTrackMeasures(
X, speed,
meanTurningAngle )
```

### Or subset tracks by one feature first:



Or visualize higher dimensional feature sets with a clustering method of choice:

clusterTracks ( X, c (speed, straightness, meanTurningAngle), method = "PCA")

see also: getFeatureMatrix() to combine with another method

speed straightness

> PC<sub>1</sub> Other methods: "UMAP", "MDS" "hclust", or "kmeans"

# 8. Simulating tracks: Models & bootstrapping

Comparing observed data to some null model 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,</pre>
 bootstrapTrack( nsteps, X ) )
```

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