Figure1B_VideoReversals

MOD

2023-09-26

```
library(rhdf5)
library(tidyverse)
create_state_column <- function(df) {</pre>
 df %>%
   mutate(
     state = case_when(
       cumsum(!is.na(realSpeed) & realSpeed > 0) >= 1 ~ "post-reversal",
     TRUE ~ "pre-reversal"),
     censored = case_when(
       cumsum(!is.na(realSpeed) & realSpeed < 0 & state == "post-reversal") >= 1 ~ "censored",
       TRUE ~ "kept"
     ))
}
files <- fs::dir_ls(path='datasets/tracking', glob = "*cest-1.2*|*N2*")
mergedData <- map_df(files, read_csv, .id = "filename")</pre>
## Rows: 1246 Columns: 11
## -- Column specification -------
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm_index, timestamp, realSpeed, coord_x_midbody, coord_y_midbody,...
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
## Rows: 1155 Columns: 11
## -- Column specification ------
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm_index, timestamp, realSpeed, coord_x_midbody, coord_y_midbody,...
##
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
## Rows: 1063 Columns: 11
## -- Column specification ------
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm_index, timestamp, realSpeed, coord_x_midbody, coord_y_midbody,...
```

##

```
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
## Rows: 1120 Columns: 11
## -- Column specification ------
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm index, timestamp, realSpeed, coord x midbody, coord y midbody,...
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
## Rows: 1141 Columns: 11
## -- Column specification -----
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm_index, timestamp, realSpeed, coord_x_midbody, coord_y_midbody,...
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
## Rows: 1056 Columns: 11
## -- Column specification ------
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm_index, timestamp, realSpeed, coord_x_midbody, coord_y_midbody,...
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
## Rows: 1180 Columns: 11
## -- Column specification -----
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm_index, timestamp, realSpeed, coord_x_midbody, coord_y_midbody,...
##
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
## Rows: 1059 Columns: 11
## -- Column specification ------
## Delimiter: ","
## chr (3): genotype, condition, flag
## dbl (8): worm_index, timestamp, realSpeed, coord_x_midbody, coord_y_midbody,...
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
#calculate a running loess fit over the data, vary the span
mergedData <- mergedData |>
 filter(!(genotype == "cest-1.2" & worm_index == 561 & filename == "datasets/tracking/2022-07-12_cest-
 #make a unique wormID
 mutate(wormID = as.factor(as.numeric(as.factor(interaction(genotype, worm_index, filename))))) |>
 group_by(genotype, worm_index, filename) |>
 create_state_column() |>
 filter(censored == "kept") |>
 nest() |>
 mutate(loess_fit = purrr::map(data,
                              loess,
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formula = realSpeed ~ realTime,
                                span = .5),
         fitted = purrr::map(loess_fit, ~predict(.x, newdata = seq(1,15, by = 0.1))))
# get the x intercepts:
Xints <- mergedData |>
  unnest(data) |>
  filter(state != lag(state) | state != lead(state)) |>
group_by(worm_index, genotype, filename) |>
  nest() |>
  dplyr::mutate(
    # get slope and y-intercept to calculate x intercept
   m = purrr::map(data, lm,
                   formula = realSpeed ~ realTime),
   xint = purrr::map(m, function(x) {
      -x$coefficients[1]/x$coefficients[2]
   })) |>
  select(worm_index, genotype, xint) |>
  unnest(cols = c(xint))
## Adding missing grouping variables: 'filename'
plot <- mergedData |>
  unnest(data) |>
ggplot(aes(x = realTime, y = realSpeed)) +
 geom_hline(yintercept = 0, linetype = 2, color = "grey") +
  geom_line(aes(color = genotype, group = wormID), size = 0.5, alpha = 0.5) +
  geom_point(data = Xints, aes(x = xint, y = 0, color = genotype), alpha = 0.5) +
  geom_smooth(aes(color = genotype, group = genotype, span = 0.1)) +
  # scale color manual(values = c( "darkgoldenrod2", "black")) +
  scale color manual(values = c( "royalblue2", "black")) +
 coord_cartesian(xlim = c(0,7.5), ylim = c(-.4,.4))
## Warning: Using 'size' aesthetic for lines was deprecated in ggplot2 3.4.0.
## i Please use 'linewidth' instead.
## This warning is displayed once every 8 hours.
## Call 'lifecycle::last_lifecycle_warnings()' to see where this warning was
## generated.
## Warning in geom_smooth(aes(color = genotype, group = genotype, span = 0.1)):
## Ignoring unknown aesthetics: span
plot2 <- ggplot(filter(Xints, xint < 7.5), aes(x = xint, y = genotype, fill = genotype)) +</pre>
   ggdist::stat_halfeye(
    # adjust bandwidth
   adjust = 1,
   # move to the right
    justification = 0,
    # remove the slub interval
    \#.width = 0,
    #point colour = NA
  alpha = 0.5) +
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```
# scale_fill_manual(values = c( "darkgoldenrod2", "black")) +
    scale_fill_manual(values = c( "royalblue2", "black")) +
   theme_void()
 library(patchwork)
plot2 / plot + plot_layout(heights = c(1,2))
## 'geom_smooth()' using method = 'gam' and formula = 'y ~ s(x, bs = "cs")'
## Warning: Removed 1467 rows containing non-finite values ('stat_smooth()').
## Warning: Removed 430 rows containing missing values ('geom_line()').
## Warning: Removed 5 rows containing missing values ('geom_point()').
                                                                            genotype
                                                                                cest-1.2
                                                                                N2
   0.4
   0.2
realSpeed
                                                                            genotype
                                                                                cest-1.2
    0.0
                                                                                N2
  -0.2
  -0.4
                                                         6
                                   realTime
# first we want to get initial x/y positions of each animal post-reversal:
# start with midbody, may want to switch to head
Pos_start <- unnest(mergedData, data) |>
  group_by(wormID) |>
  summarize(X0 = first(coord_x_midbody), Y0 = first(coord_y_midbody))
mergedData <- unnest(mergedData, data) |>
```

full_join(Pos_start, by = join_by(wormID)) |>

```
# now use the XO, YO to calculate total (net) distance traveled at each time point
  mutate(distance = sqrt((X0 - coord_x_midbody)^2 + (Y0 - coord_y_midbody)^2))
#correct distance for pixelSize in 2022-07-12 data
mergedData <- mergedData |>
  mutate(
    distance = case_when(
      stringr::str_detect(filename, "2022-07-12") ~ distance * (9.7/11.1) / 10,
      TRUE ~ distance
    )
  )
# get max distance during reversal
maxD <- mergedData |>
  filter(state == "pre-reversal") |>
  group_by(wormID, genotype) |>
  summarise(maxD = max(distance), realTime = max(realTime))
## 'summarise()' has grouped output by 'wormID'. You can override using the
## '.groups' argument.
# now plot distance over time:
plot3 <- mergedData |>
    filter(state == "pre-reversal") |>
  ggplot() +
  geom_line(aes(x = realTime,
                y = distance,
                group = wormID,
                color = genotype), alpha = 0.5) +
  geom_point(data = maxD,
             aes(x = realTime, y = maxD, color = genotype), alpha = 0.5) +
  coord cartesian(xlim = c(0,7.5)) +
  scale_color_manual(values = c( "royalblue2", "black")) +
  guides(color = 'none')
plot4 <- ggplot(filter(maxD, realTime < 7.5),</pre>
                aes(y = maxD, x = genotype, fill = genotype)) +
   ggdist::stat_halfeye(
    # adjust bandwidth
    adjust = 1,
    # move to the right
    justification = 0,
    # remove the slub interval
    #.width = 0,
    \#point\_colour = NA
  alpha = 0.5) +
   # scale_fill_manual(values = c( "darkgoldenrod2", "black")) +
    scale fill manual(values = c( "royalblue2", "black")) +
   theme_void()
```

plot3 + plot4 + plot_layout(widths = c(5,2))

Warning: Removed 207 rows containing missing values ('geom_line()').
Warning: Removed 8 rows containing missing values ('geom_point()').
Warning: Removed 5 rows containing missing values ('stat_slabinterval()').

