

brachypoder

This is the vignette of **brachypoder**, an R package developed to help read and assemble the data simulated by the **brachypode** program.

To install the package from GitHub, use:

```
devtools::install_github("rscherrer/brachypoder")
```

To load the package, use:

```
library(brachypoder)
```

Some example data are provided with the package. To extract where they are saved, run:

```
root <- system.file("extdata", "sim-example", package = "brachypoder")
root
#> [1] "/home/raphael/R/x86_64-pc-linux-gnu-library/4.1/brachypoder/extdata/sim-example"
```

If we have a look at what is inside this folder,

```
list.files(root)
#> [1] "architecture.txt" "brachypode"      "demesizes.dat"   "gmon.out"
#> [5] "individuals.dat"  "parameters.txt"  "paramlog.txt"    "patchsizes.dat"
#> [9] "popsize.dat"      "time.dat"        "traitmeans.dat"
```

we see a few **.dat** files. These are the files that contain the data saved from one simulation. Different variables are saved in different files, e.g. **popsize.dat** contains the total population size at every time point saved. See the repository of `\textcolor{blue}{brachypode}` for details.

Each data file contains data with different resolutions. For example, **individuals.dat** contains five values per individual and covers all individuals of all the time points saved. The data files are saved in binary format, which is one-dimensional. This means that the arrays of saved values may have to be reshaped in order to be assembled into a data set that is workable within R. This is where the **read_data** function comes in.

The function **read_data** is the core function of the package. It takes the location of the simulation data as input, the variables to read, and rules that determine how each variable should be reshaped. For example,

```
read_data(root, "popsize")
#> # A tibble: 101 x 1
#>   popsize
#>   <dbl>
#> 1      10
#> 2     206
#> 3     250
#> 4     216
#> 5     220
#> 6     210
#> 7     230
#> 8     201
#> 9     206
#> 10    207
#> # ... with 91 more rows
```

reads the binary data in `popsiz.dat` back into numbers and places them into a one-column tibble. If we now run:

```
read_data(root, c("time", "popsiz"))
#> # A tibble: 101 x 2
#>   time popsiz
#>   <dbl> <dbl>
#> 1     0    10
#> 2   100   206
#> 3   200   250
#> 4   300   216
#> 5   400   220
#> 6   500   210
#> 7   600   230
#> 8   700   201
#> 9   800   206
#> 10  900   207
#> # ... with 91 more rows
```

we get multiple variables (`time.dat` and `popsiz.dat`) read and assembled together. Now, these two variables both have one value saved every saved time point, and so have the same number of values. When the variables to read have different dimensions (e.g. `patchsiz.dat` contains one value per patch, per site, per time point), we use the `ncols` argument:

```
read_data(root, c("time", "patchsiz"), ncols = c(1, 10))
#> # A tibble: 101 x 11
#>   time patchsiz1 patchsiz2 patchsiz3 patchsiz4 patchsiz5 patchsiz6
#>   <dbl>      <dbl>      <dbl>      <dbl>      <dbl>      <dbl>      <dbl>
#> 1     0         0        10         0         0         0         0
#> 2   100        16        47         34        31        36        21
#> 3   200        15        62         25        52        37        36
#> 4   300        10        57         35        29        34        26
#> 5   400        15        54         21        28        32        27
#> 6   500        14        62         25        41        15        33
#> 7   600        17        57         34        49        27        19
#> 8   700        16        43         34        37        33        16
#> 9   800         6        66         23        40        25        19
#> 10  900        12        53         34        42        24        16
#> # ... with 91 more rows, and 4 more variables: patchsiz7 <dbl>,
#> #   patchsiz8 <dbl>, patchsiz9 <dbl>, patchsiz10 <dbl>
```

which splits the `patchsiz` data into 10 columns, and so its number of rows becomes equal to the number of time points in `time` (the 1 tells the function not to split that column), with which they can be attached. The columns in the output data are named after their respective data file or origin, with numbers appended for each column.

In some cases some columns may need to be duplicated instead of split into multiple columns. Then, we use negative numbers in `ncols`. For example,

```
read_data(root, c("time", "patchsiz"), ncols = c(-10, 1))
#> # A tibble: 1,010 x 2
#>   time patchsiz
#>   <dbl>      <dbl>
#> 1     0         0
#> 2     0        10
#> 3     0         0
```

```
#> 4      0      0
#> 5      0      0
#> 6      0      0
#> 7      0      0
#> 8      0      0
#> 9      0      0
#> 10     0      0
#> # ... with 1,000 more rows
```

reads the same data as the previous chunk, but in a “longer” format, where each number of individuals is taken as an observation and therefore as a row. And so here, `patchsizes` was kept in a single column but each value in `time` was duplicated as many times as there are patches and sites (so 10 times).

Some use-cases may be more intricate. For example, reading `individuals.dat` and assigning each individual its own time point requires to know how many individuals there are at each time point. For this we provide a wrapper around `read_data` that reads `individuals.dat`, `time.dat` and `popsizes.dat` to perform this task:

```
read_individual_data(root)
#> # A tibble: 22,038 x 6
#>   time deme patch      x      y      z
#>   <dbl> <dbl> <dbl> <dbl> <dbl> <dbl>
#> 1     0     0     1     0     0     0
#> 2     0     0     1     0     0     0
#> 3     0     0     1     0     0     0
#> 4     0     0     1     0     0     0
#> 5     0     0     1     0     0     0
#> 6     0     0     1     0     0     0
#> 7     0     0     1     0     0     0
#> 8     0     0     1     0     0     0
#> 9     0     0     1     0     0     0
#> 10    0     0     1     0     0     0
#> # ... with 22,028 more rows
```

This data set contains all the relevant information for each individual saved in the simulation. Because this function has only one use-case (i.e. it is not designed to be flexible, unlike `read_data`) it gives the output tibble specific column names.

Same, we provide a function that directly reads and formats the data from the `traitmeans.dat` data set, which contains the mean trait value for each trait in each patch of each site, at each saved generation.

```
read_trait_mean_data(root)
#> # A tibble: 1,010 x 6
#>   time deme patch      x      y      z
#>   <dbl> <int> <int> <dbl> <dbl> <dbl>
#> 1     0     1     0     0     0     0
#> 2     0     1     1     0     0     0
#> 3     0     2     0     0     0     0
#> 4     0     2     1     0     0     0
#> 5     0     3     0     0     0     0
#> 6     0     3     1     0     0     0
#> 7     0     4     0     0     0     0
#> 8     0     4     1     0     0     0
#> 9     0     5     0     0     0     0
#> 10    0     5     1     0     0     0
#> # ... with 1,000 more rows
```

Besides the functions to read simulation data, one may want to read back the parameters that were used in a given simulation. To do this, use:

```
pars <- read_parameters(root)
pars[1:5] # just a few parameters
#> $type
#> [1] 1
#>
#> $popsize
#> [1] 10
#>
#> $pgood
#> [1] 5.0 0.8 0.6 0.5 0.3 0.1
#>
#> $maxgrowths
#> [1] 2 3
#>
#> $zwidths
#> [1] 1 2
```

Once the data are properly loaded and reshaped, they can be plotting using the usual tools available in R. Using `ggplot2`, for example:

```
library(tidyverse)
#> -- Attaching packages ----- tidyverse 1.3.1 --
#> v ggplot2 3.3.5      v purrr 0.3.4
#> v tibble 3.1.6       v dplyr 1.0.7
#> v tidyr 1.1.4        v stringr 1.4.0
#> v readr 2.1.1       v forcats 0.5.1
#> -- Conflicts ----- tidyverse_conflicts() --
#> x dplyr::filter() masks stats::filter()
#> x dplyr::lag()     masks stats::lag()

# Read the proportion of good patches
pgood <- read_parameters(root)$pgood
pgood <- pgood[-1]

# Read trait means per patch per deme
data <- read_trait_mean_data(root)
data$patch <- as.logical(data$patch)

# Relabel
data <- data %>% mutate(patch_lab = if_else(patch, "Facilitated", "Unfacilitated"))

# Add information about the coverage in good patches
data <- data %>%
  group_by(deme) %>%
  nest() %>%
  ungroup() %>%
  mutate(pgood = pgood) %>%
  unnest(data) %>%
  mutate(patch_size = if_else(patch, pgood, 1 - pgood))

# Plot trait z through time
data %>%
```

```
ggplot(aes(x = time, y = z, group = interaction(deme, patch), color = patch_lab, alpha = patch_size))
  geom_line() +
  xlab("Time (generations)") +
  ylab("Mean trait value") +
  labs(color = NULL, alpha = "% cover") +
  scale_color_manual(values = c("gray10", "gray60"))
```

