```
library(here)
## here() starts at /Users/ruanvanmazijk/Desktop/Cape-vs-SWA
library(tidyverse)
## -- Attaching packages --

    tidyverse

## v ggplot2 3.2.1
                    v purrr
                               0.3.3
## v tibble 2.1.3
                     v dplyr
                               0.8.3
## v tidyr
           1.0.0
                    v stringr 1.4.0
## v readr
            1.3.1
                     v forcats 0.4.0
## -- Conflicts ------ tidyverse_confl
## x dplyr::filter() masks stats::filter()
## x dplyr::lag()
                    masks stats::lag()
library(magrittr)
##
## Attaching package: 'magrittr'
## The following object is masked from 'package:purrr':
##
##
      set_names
## The following object is masked from 'package:tidyr':
##
##
      extract
library(raster)
## Loading required package: sp
##
## Attaching package: 'raster'
## The following object is masked from 'package:magrittr':
##
##
      extract
## The following object is masked from 'package:dplyr':
##
##
      select
## The following object is masked from 'package:tidyr':
##
##
      extract
library(rgdal)
## rgdal: version: 1.4-7, (SVN revision 845)
## Geospatial Data Abstraction Library extensions to R successfully loaded
## Loaded GDAL runtime: GDAL 2.4.2, released 2019/06/28
## Path to GDAL shared files: /Library/Frameworks/R.framework/Versions/3.6/Resources/library/rgdal/gda
## GDAL binary built with GEOS: FALSE
## Loaded PROJ.4 runtime: Rel. 5.2.0, September 15th, 2018, [PJ_VERSION: 520]
## Path to PROJ.4 shared files: /Library/Frameworks/R.framework/Versions/3.6/Resources/library/rgdal/p
```

Linking to sp version: 1.3-1

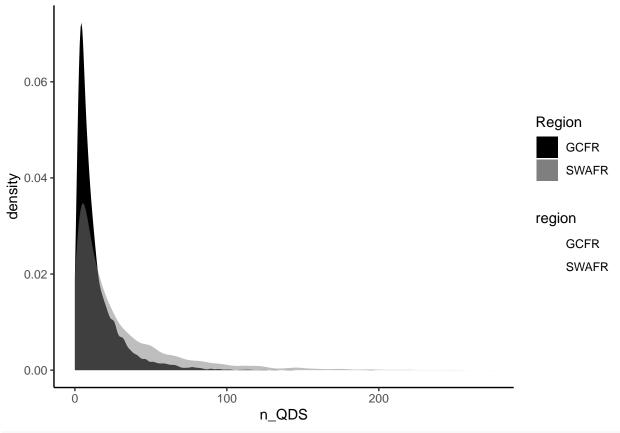
```
library(vegan)
## Loading required package: permute
## Loading required package: lattice
## This is vegan 2.5-6
data <- read_csv(here(</pre>
  "data/derived-data/May-2019",
  "data-QDS.csv"
))
## Parsed with column specification:
## cols(
##
     Elevation = col_double(),
     MAP = col_double(),
##
     PDQ = col_double(),
     Surface_T = col_double(),
##
    NDVI = col double(),
##
##
    CEC = col_double(),
##
     Clay = col_double(),
##
     Soil_C = col_double(),
##
    pH = col_double(),
##
     region = col_character(),
##
    PC1 = col_double(),
##
    lon = col_double(),
##
     lat = col_double(),
##
     QDS = col_character(),
##
     n_EDS_in_region = col_double(),
##
     n_collections = col_double(),
     QDS_richness = col_double()
## )
GCFR_species <- read_csv(here(</pre>
  "data/derived-data/May-2019",
  "GCFR-species.csv"
))
## Parsed with column specification:
## cols(
##
     species = col_character(),
     n_collections = col_double()
## )
SWAFR_species <- read_csv(here(
 "data/derived-data/May-2019",
  "SWAFR-species.csv"
))
## Parsed with column specification:
## cols(
     species = col_character(),
##
    n_collections = col_double()
## )
GCFR_box <- readOGR(here("data/derived-data/borders/GCFR_box"))</pre>
```

```
## OGR data source with driver: ESRI Shapefile
## Source: "/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/derived-data/borders/GCFR_box", layer: "value
## with 1 features
## It has 1 fields
SWAFR_box <- readOGR(here("data/derived-data/borders/SWAFR_box"))</pre>
## OGR data source with driver: ESRI Shapefile
## Source: "/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/derived-data/borders/SWAFR_box", layer: "valu
## with 1 features
## It has 1 fields
ZA_EDS <- readOGR(here("data/raw-data/QDGC/qdgc_zaf"), layer = "qdgc_03_zaf")
## OGR data source with driver: ESRI Shapefile
## Source: "/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/raw-data/QDGC/qdgc_zaf", layer: "qdgc_03_zaf"
## with 14144 features
## It has 4 fields
AU_EDS <- readOGR(here("data/raw-data/QDGC/qdgc_aus"), layer = "qdgc_03_aus")
## OGR data source with driver: ESRI Shapefile
## Source: "/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/raw-data/QDGC/qdgc_aus", layer: "qdgc_03_aus"
## with 89216 features
## It has 4 fields
GCFR_EDS <- crop(ZA_EDS, GCFR_box)</pre>
SWAFR_EDS <- crop(AU_EDS, SWAFR_box)
Larsen_grid <- rbind(GCFR_EDS, SWAFR_EDS)</pre>
Larsen_grid$edgc <- Larsen_grid$qdgc</pre>
Larsen_grid$qdgc <- str_remove(Larsen_grid$edgc, ".$")</pre>
Larsen_grid$hdgc <- str_remove(Larsen_grid$qdgc, ".$")</pre>
Larsen_grid$dgc <- str_remove(Larsen_grid$hdgc, ".$")</pre>
make_SpatialPointsDataFrame <- function(df) {</pre>
  SpatialPointsDataFrame(
                = df[, c("decimallongitude", "decimallatitude")],
    coords
               = df[, "species"],
    proj4string = crs(Larsen_grid)
  )
}
GCFR_species_occ <- make_SpatialPointsDataFrame(read_csv(here(</pre>
  "data/derived-data/flora",
  "GCFR clean flora 2017-09-14.csv"
)))
## Warning: Missing column names filled in: 'X1' [1]
## Parsed with column specification:
## cols(
    X1 = col_double(),
##
##
    family = col_character(),
    genus = col character(),
     species = col_character(),
##
##
     infraspecificepithet = col_character(),
     scientificname = col_character(),
```

```
##
     taxonrank = col_character(),
##
     decimallatitude = col_double(),
##
     decimallongitude = col_double(),
##
     coordinateuncertaintyinmeters = col_double(),
##
     coordinateprecision = col_logical()
## )
## Warning: 1 parsing failure.
                                         expected actual
## 392500 coordinateprecision 1/0/T/F/TRUE/FALSE
                                                     0.0 '/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/
SWAFR_species_occ <- make_SpatialPointsDataFrame(read_csv(here(</pre>
  "data/derived-data/flora",
  "SWAFR_clean_flora_2017-09-14.csv"
)))
## Warning: Missing column names filled in: 'X1' [1]
## Parsed with column specification:
## cols(
##
     X1 = col_double(),
##
     family = col_character(),
##
     genus = col_character(),
##
     species = col_character(),
##
     infraspecificepithet = col_character(),
##
     scientificname = col_character(),
##
     taxonrank = col_character(),
##
     decimallatitude = col_double(),
##
     decimallongitude = col_double(),
##
     coordinateuncertaintyinmeters = col_double(),
     coordinateprecision = col_logical()
##
## )
## Warning: 4 parsing failures.
                                         expected actual
## 72178 coordinateprecision 1/0/T/F/TRUE/FALSE
                                                    20.0 '/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/
                                                    20.0 '/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/
## 126045 coordinateprecision 1/0/T/F/TRUE/FALSE
## 126046 coordinateprecision 1/0/T/F/TRUE/FALSE
                                                     10.0 '/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/
## 225575 coordinateprecision 1/0/T/F/TRUE/FALSE
                                                     15.0 '/Users/ruanvanmazijk/Desktop/Cape-vs-SWA/data/
species_occ <- rbind(GCFR_species_occ, SWAFR_species_occ)</pre>
species_occ$EDS <- species_occ %over%</pre>
 Larsen_grid %>%
  pull(edgc)
species_occ@data$EDS %<>% as.character()
species_occ$QDS <- str_remove(species_occ$EDS, ".$")</pre>
species_occ$HDS <- str_remove(species_occ$QDS, ".$")</pre>
species_occ$DS <- str_remove(species_occ$HDS, ".$")</pre>
species_occ2 <- species_occ@data %>%
 mutate(
    lon = DS \%
     str_extract("E\\d\\d\\d") %>%
     str remove("E") %>%
      as.numeric(),
    region = ifelse(lon >= 112, "SWAFR", "GCFR")
```

```
) %>%
  split(.$region) %>%
  map(dplyr::select, -EDS, -lon, -region) %>%
  map(distinct)
GCFR_species %<>% filter(n_collections >= 5)
SWAFR_species %<>% filter(n_collections >= 5)
cells <- sort(unique(species_occ2$GCFR$QDS))</pre>
cells <- cells[cells %in% data$QDS]</pre>
species <- sort(unique(species_occ2$GCFR$species))</pre>
species <- species[species %in% GCFR_species$species]</pre>
n_cells <- length(cells)</pre>
n_species <- length(species)</pre>
GCFR_matrix <- matrix(nrow = n_cells, ncol = n_species)</pre>
rownames(GCFR_matrix) <- cells</pre>
colnames(GCFR_matrix) <- species</pre>
for (i in 1:nrow(GCFR_matrix)) {
  GCFR_matrix[i, ] <- species %in% species_occ2$GCFR$species[</pre>
    species_occ2$GCFR$QDS == cells[[i]]
}
cells <- sort(unique(species_occ2$SWAFR$QDS))</pre>
cells <- cells[cells %in% data$QDS]</pre>
species <- sort(unique(species occ2$SWAFR$species))</pre>
species <- species[species %in% SWAFR_species$species]</pre>
n_cells <- length(cells)</pre>
n_species <- length(species)</pre>
SWAFR_matrix <- matrix(nrow = n_cells, ncol = n_species)</pre>
rownames(SWAFR_matrix) <- cells</pre>
colnames(SWAFR_matrix) <- species</pre>
for (i in 1:nrow(SWAFR_matrix)) {
  SWAFR_matrix[i, ] <- species %in% species_occ2$SWAFR$species[</pre>
    species_occ2$SWAFR$QDS == cells[[i]]
}
GCFR_range_sizes <- GCFR_matrix %>%
  apply(2, sum) %>%
  sort(decreasing = TRUE) %>%
  {tibble(region = "GCFR", species = names(.), n_QDS = .)}
SWAFR_range_sizes <- SWAFR_matrix %>%
  apply(2, sum) %>%
  sort(decreasing = TRUE) %>%
  {tibble(region = "SWAFR", species = names(.), n_QDS = .)}
range sizes <-
  full_join(GCFR_range_sizes, SWAFR_range_sizes) %>%
  mutate(log10_n_QDS = log10(n_QDS))
## Joining, by = c("region", "species", "n_QDS")
ggplot(range sizes) +
  aes(n_QDS, fill = region, alpha = region) +
```

```
geom_density(colour = NA) +
scale_fill_manual(name = "Region", values = c("black", "grey50")) +
scale_alpha_manual(values = c(1, 0.5)) +
theme_classic()
```



```
ggplot(range_sizes[range_sizes$log10_n_QDS >= 0, ]) +
  aes(log10_n_QDS, fill = region, alpha = region) +
  geom_density(colour = NA) +
  scale_fill_manual(name = "Region", values = c("black", "grey50")) +
  scale_alpha_manual(values = c(1, 0.5)) +
  theme_classic()
```

```
0.75
                                                                               Region
                                                                                   GCFR
                                                                                   SWAFR
density density
                                                                               region
                                                                                   GCFR
                                                                                   SWAFR
  0.25
  0.00
         0.0
                     0.5
                                  1.0
                                               1.5
                                                            2.0
                                                                         2.5
                                  log10_n_QDS
t.test(
  log10_n_QDS ~ region,
  range_sizes[range_sizes$log10_n_QDS >= 0, ]
)
##
##
   Welch Two Sample t-test
## data: log10_n_QDS by region
## t = -30.579, df = 12721, p-value < 2.2e-16
\#\# alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -0.2500060 -0.2198856
## sample estimates:
    mean in group GCFR mean in group SWAFR
##
             0.9407647
                                  1.1757104
wilcox.test(
  n_QDS ~ region,
  range_sizes[range_sizes$log10_n_QDS >= 0, ]
)
##
##
   Wilcoxon rank sum test with continuity correction
## data: n_QDS by region
## W = 23322959, p-value < 2.2e-16
\#\# alternative hypothesis: true location shift is not equal to 0
```