Act 4 Field

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library(tidyverse) ## Warning: package 'tidyverse' was built under R version 4.0.5 ## -- Attaching packages ------ tidyverse 1.3.0 --## v ggplot2 3.3.3 v purrr 0.3.4 ## v tibble 3.1.0 v dplyr 1.0.5 v stringr 1.4.0 ## v tidyr 1.1.3 ## v readr 1.4.0 v forcats 0.5.1 ## Warning: package 'ggplot2' was built under R version 4.0.5 ## Warning: package 'tidyr' was built under R version 4.0.5 ## Warning: package 'readr' was built under R version 4.0.5 ## Warning: package 'forcats' was built under R version 4.0.5 ## -- Conflicts ----- tidyverse_conflicts() --## x dplyr::filter() masks stats::filter() ## x dplyr::lag() masks stats::lag() event <- read_csv("D:/kaya/Documents/2 SY 2021-2022/1st Semester 2021-2022/Bio 115 - Field Biology/Labo ## -- Column specification -------## cols(## .default = col_character(), ## year = col_double(), ## month = col_double(), day = col_logical(), ## decimalLongitude = col_double(), decimalLatitude = col_double(),

coordinateUncertaintyInMeters = col_double()

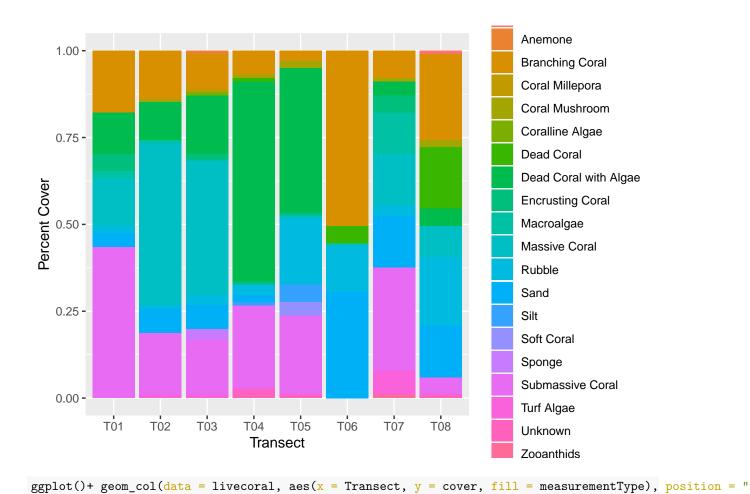
i Use 'spec()' for the full column specifications.

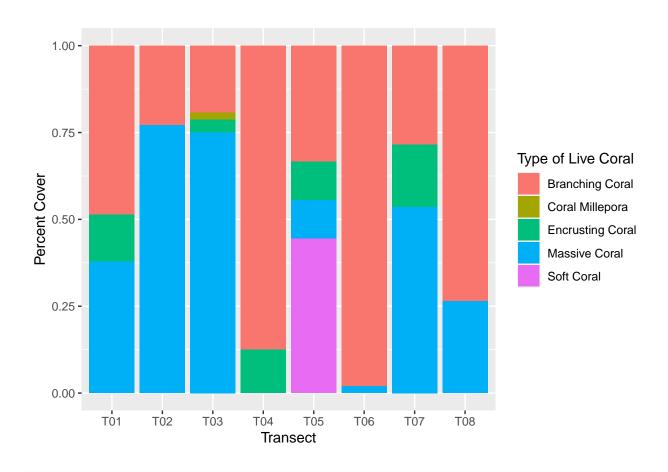
##

```
emof <- read_csv("D:/kaya/Documents/2 SY 2021-2022/1st Semester 2021-2022/Bio 115 - Field Biology/Labor
##
## -- Column specification ------
##
    eventID = col_character(),
    measurementID = col_character(),
##
##
    measurementType = col_character(),
##
    measurementValue = col_character(),
##
    measurementUnit = col_character(),
##
    measurementDeterminedDate = col_character(),
    measurementDeterminedBy = col_character(),
##
    measurementMethod = col_character(),
##
##
    measurementRemarks = col_character()
## )
occurrence <- read_csv("D:/kaya/Documents/2 SY 2021-2022/1st Semester 2021-2022/Bio 115 - Field Biology
## -- Column specification -------
    .default = col_character(),
##
    decimalLatitude = col_double(),
##
##
    decimalLongitude = col_double(),
    coordinateUncertaintyInMeters = col_double(),
##
    individualCount = col_double(),
##
    organismQuantity = col_double()
## )
## i Use 'spec()' for the full column specifications.
newoccurrence <- occurrence %>%
                 select(occurrenceID, organismQuantityType, scientificName, family, genus, order, indi
benthic <- emof %>%
       filter(str detect(measurementID, "benthic")) %>%
       filter(!str_detect(measurementID, "T09")) %>%
       filter(!str_detect(measurementID, "T10")) %>%
       select(measurementID, measurementType, measurementValue) %>%
 separate(col = measurementID, into = c("Location", "Transect", "Date", "Benthic", "Type"), sep = ":")
benthicfinal <- benthic %>%
           select(Transect, measurementType, measurementValue) %% mutate(cover = as.numeric(measureme
livecoral <- benthicfinal %>%
         filter(str_detect(measurementType, "Branching|Massive|Encrusting|Millepora| Submassive|Soft")
totalcover <-livecoral %>%
             spread(key = Transect, value = cover)
transect <- c("T01", "T02", "T03", "T04", "T05", "T06", "T07", "T08")
Totalcover \leftarrow c(0.3663, 0.6039, 0.5148, 0.0792, 0.0891, 0.5149, 0.2772, 0.3366)
```

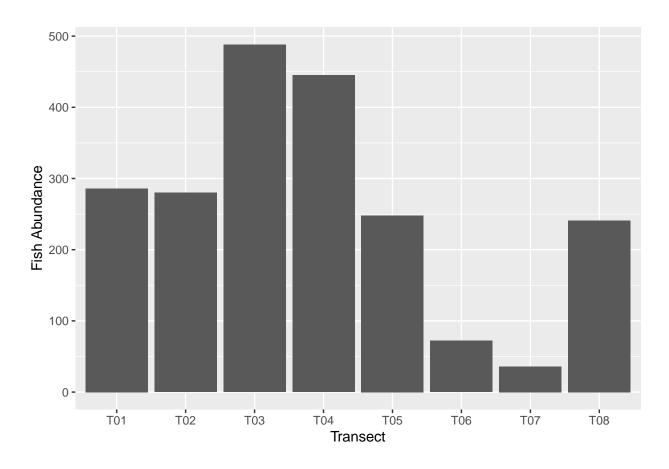
```
covertotal<-data.frame(transect, Totalcover)</pre>
fish <- newoccurrence %>%
      filter(str_detect(occurrenceID, "fish"))
fishfinal <- fish %>%
  select(occurrenceID, organismQuantityType, scientificName, individualCount) %>%
  separate(col = occurrenceID, into = c("Location", "Transect", "Date", "Benthic", "Species"), sep = ":
  select(Transect, Species, organismQuantityType, individualCount)
## Warning: Expected 5 pieces. Additional pieces discarded in 331 rows [1, 2, 3, 4,
## 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, ...].
fishcount <- fishfinal %>%
      spread(key = Transect, value = individualCount) %>%
      mutate_if(is.integer, as.numeric)
fishtotal <- data.frame(colSums(Filter(is.numeric,fishcount), na.rm = TRUE))</pre>
transects <- c("T01", "T02", "T03", "T04", "T05", "T06", "T07", "T08")
fishabundance <- c(286, 280, 488, 445, 248, 72, 36, 241)
totalfish <- data.frame(transects, fishabundance)</pre>
coralfish <- data.frame(totalfish, covertotal)</pre>
% cover per transect
```

 $ggplot() + geom_col(data = benthicfinal, aes(x = Transect, y = cover, fill = measurementType), position = fill = measurementType), position = fill = fill$

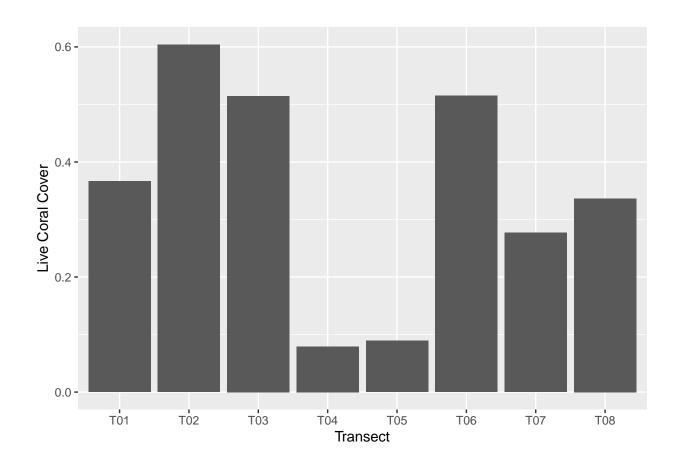




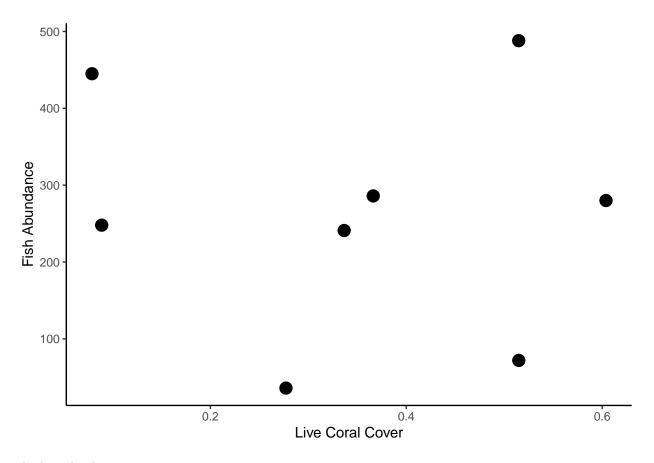
 $ggplot(data = totalfish, aes(x = transects, y = fishabundance)) + geom_col() + labs(x = "Transect", y = fishabundance)) + geom_col() + labs(x = "Transect", y = fishabundance))$



 $ggplot(data = covertotal, aes(x = transect, y = Totalcover)) + geom_col() + labs(x = "Transect", y = "Lotalcover")$

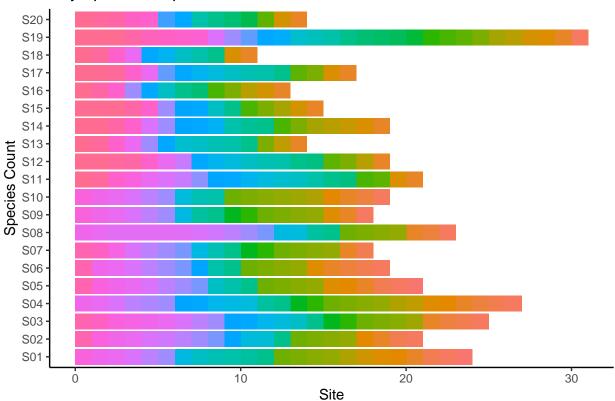


 $ggplot(data = coralfish, aes(x = Totalcover, y = fishabundance), fill = transect) + geom_point(size = 4)$



plankton biodiversity





```
phytoplanktonNew <- phytoplankton %>%
    select(Site,Species,individualCount) %>%
    group_by(Species, Site)

planktonwide <- phytoplanktonNew %>% pivot_wider(names_from = Species, values_from = individualCount, S

## Warning: Values are not uniquely identified; output will contain list-cols.

## * Use 'values_fn = list' to suppress this warning.

## * Use 'values_fn = length' to identify where the duplicates arise

## * Use 'values_fn = {summary_fun}' to summarise duplicates

planktonwide[planktonwide == "NULL"] <- NA

library(readxl)

## Warning: package 'readxl' was built under R version 4.0.5

library(vegan)

## Warning: package 'vegan' was built under R version 4.0.5

## Loading required package: permute</pre>
```

```
## Warning: package 'permute' was built under R version 4.0.5

## Loading required package: lattice

## This is vegan 2.5-7

Diversity <- read_excel("D:/kaya/Documents/2 SY 2021-2022/1st Semester 2021-2022/Bio 115 - Field Biolog sheet = "Sheet2")

diversity_results <- diversity(x = Diversity,index = "shannon")
spa <- specaccum(Diversity)

## Warning in cor(x > 0): the standard deviation is zero

plot(spa)
```

