# Homework 3 dada2 setup - Spring 2019 - ANSWER KEY

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### Document package installation

Use library() function to load each package to show that the packages are installed.

You have to use the BioConductor approach and tools to install the dada2 (which includes Shortread), phyloseq and DECIPHER. I installed my packages using a separate R script (see "packages\_install.R" stored in this Github repository). But here is the script code to install these packages.

```
# install BiocManager
if (!requireNamespace("BiocManager"))
  install.packages("BiocManager")
BiocManager::install()
# install dada2 - includes ShortRead pacakge
if (!requireNamespace("BiocManager", quietly = TRUE))
  install.packages("BiocManager")
BiocManager::install("dada2", version = "3.8")
# install phyloseq
if (!requireNamespace("BiocManager", quietly = TRUE))
  install.packages("BiocManager")
BiocManager::install("phyloseq", version = "3.8")
# install DECIPHER
if (!requireNamespace("BiocManager", quietly = TRUE))
  install.packages("BiocManager")
BiocManager::install("DECIPHER", version = "3.8")
ggplot2 can be installed from CRAN using the RStudio /Tools/Install Packages menu.
library(dada2)
## Loading required package: Rcpp
library(ShortRead)
## Loading required package: BiocGenerics
## Loading required package: parallel
```

```
##
## Attaching package: 'BiocGenerics'
## The following objects are masked from 'package:parallel':
##
##
       clusterApply, clusterApplyLB, clusterCall, clusterEvalQ,
##
       clusterExport, clusterMap, parApply, parCapply, parLapply,
##
       parLapplyLB, parRapply, parSapply, parSapplyLB
## The following objects are masked from 'package:stats':
##
##
       IQR, mad, sd, var, xtabs
## The following objects are masked from 'package:base':
##
##
       anyDuplicated, append, as.data.frame, basename, cbind,
       colMeans, colnames, colSums, dirname, do.call, duplicated,
##
##
       eval, evalq, Filter, Find, get, grep, grepl, intersect,
       is.unsorted, lapply, lengths, Map, mapply, match, mget, order,
##
##
       paste, pmax, pmax.int, pmin, pmin.int, Position, rank, rbind,
##
       Reduce, rowMeans, rownames, rowSums, sapply, setdiff, sort,
       table, tapply, union, unique, unsplit, which, which.max,
##
##
       which.min
## Loading required package: BiocParallel
## Loading required package: Biostrings
## Loading required package: S4Vectors
## Loading required package: stats4
##
## Attaching package: 'S4Vectors'
## The following object is masked from 'package:base':
##
##
       expand.grid
## Loading required package: IRanges
## Attaching package: 'IRanges'
## The following object is masked from 'package:grDevices':
##
##
       windows
## Loading required package: XVector
```

```
##
## Attaching package: 'Biostrings'
## The following object is masked from 'package:base':
##
##
       strsplit
## Loading required package: Rsamtools
## Loading required package: GenomeInfoDb
## Loading required package: GenomicRanges
## Loading required package: GenomicAlignments
## Loading required package: SummarizedExperiment
## Loading required package: Biobase
## Welcome to Bioconductor
##
##
       Vignettes contain introductory material; view with
       'browseVignettes()'. To cite Bioconductor, see
##
##
       'citation("Biobase")', and for packages 'citation("pkgname")'.
## Loading required package: DelayedArray
## Loading required package: matrixStats
##
## Attaching package: 'matrixStats'
## The following objects are masked from 'package:Biobase':
##
##
       anyMissing, rowMedians
##
## Attaching package: 'DelayedArray'
## The following objects are masked from 'package:matrixStats':
##
##
       colMaxs, colMins, colRanges, rowMaxs, rowMins, rowRanges
## The following object is masked from 'package:Biostrings':
##
##
       type
## The following objects are masked from 'package:base':
##
##
       aperm, apply
```

```
library(phyloseq)
##
## Attaching package: 'phyloseq'
## The following object is masked from 'package:SummarizedExperiment':
##
##
       distance
## The following object is masked from 'package:Biobase':
##
##
       sampleNames
## The following object is masked from 'package:GenomicRanges':
##
##
       distance
## The following object is masked from 'package: IRanges':
##
##
       distance
library(DECIPHER)
## Loading required package: RSQLite
## Warning: package 'RSQLite' was built under R version 3.5.3
library(ggplot2)
```

## Shows you've downloaded the fastq files

The dada2 tutorial explains this part at http://benjjneb.github.io/dada2/tutorial.html.

```
# CHANGE ME to the directory containing the fastq files after unzipping.
# path <- "~/MiSeq_SOP" # MAC path specification
path <- "./MiSeq_SOP" # PC path specification
list.files(path)</pre>
```

```
##
   [1] "F3D0_S188_L001_R1_001.fastq"
   [2] "F3D0_S188_L001_R2_001.fastq"
##
   [3] "F3D1_S189_L001_R1_001.fastq"
##
   [4] "F3D1_S189_L001_R2_001.fastq"
   [5] "F3D141_S207_L001_R1_001.fastq"
##
   [6] "F3D141_S207_L001_R2_001.fastq"
##
   [7] "F3D142_S208_L001_R1_001.fastq"
##
##
   [8] "F3D142_S208_L001_R2_001.fastq"
##
  [9] "F3D143_S209_L001_R1_001.fastq"
## [10] "F3D143_S209_L001_R2_001.fastq"
## [11] "F3D144_S210_L001_R1_001.fastq"
```

```
## [12] "F3D144_S210_L001_R2_001.fastq"
  [13] "F3D145_S211_L001_R1_001.fastq"
  [14] "F3D145 S211 L001 R2 001.fastq"
  [15] "F3D146_S212_L001_R1_001.fastq"
  [16] "F3D146_S212_L001_R2_001.fastq"
  [17] "F3D147 S213 L001 R1 001.fastq"
  [18] "F3D147 S213 L001 R2 001.fastq"
  [19] "F3D148_S214_L001_R1_001.fastq"
   [20] "F3D148_S214_L001_R2_001.fastq"
   [21] "F3D149_S215_L001_R1_001.fastq"
  [22] "F3D149_S215_L001_R2_001.fastq"
   [23] "F3D150_S216_L001_R1_001.fastq"
   [24] "F3D150_S216_L001_R2_001.fastq"
  [25] "F3D2_S190_L001_R1_001.fastq"
  [26] "F3D2_S190_L001_R2_001.fastq"
   [27] "F3D3_S191_L001_R1_001.fastq"
   [28] "F3D3_S191_L001_R2_001.fastq"
   [29] "F3D5 S193 L001 R1 001.fastg"
   [30] "F3D5_S193_L001_R2_001.fastq"
   [31] "F3D6_S194_L001_R1_001.fastq"
##
  [32] "F3D6_S194_L001_R2_001.fastq"
  [33] "F3D7_S195_L001_R1_001.fastq"
  [34] "F3D7_S195_L001_R2_001.fastq"
   [35] "F3D8_S196_L001_R1_001.fastq"
##
   [36] "F3D8_S196_L001_R2_001.fastq"
##
  [37] "F3D9_S197_L001_R1_001.fastq"
   [38] "F3D9_S197_L001_R2_001.fastq"
##
   [39] "HMP_MOCK.v35.fasta"
  [40] "Mock_S280_L001_R1_001.fastq"
  [41] "Mock_S280_L001_R2_001.fastq"
  [42] "mouse.dpw.metadata"
##
   [43] "mouse.time.design"
   [44] "silva_nr_v132_train_set.fa.gz"
   [45] "silva_species_assignment_v132.fa.gz"
   [46] "SILVA_SSU_r132_March2018.RData"
  [47] "stability.batch"
## [48] "stability.files"
```

### Get the Silva 132 training set and the Silva 132 species assignment zip files

see https://zenodo.org/record/1172783#.XJkdyKBKiUl

Put these 2 \*.gz files into the same directory as specified above.

## Get SILVA SSU r132 RData object

See http://www2.decipher.codes/Downloads.html - download the training file for Silva SSU r132. This is a RData file. Put in same directory as above.

List files again

```
path <- "./MiSeq_SOP"
list.files(path)</pre>
```

```
##
    [1] "F3D0 S188 L001 R1 001.fastg"
    [2] "F3D0_S188_L001_R2_001.fastq"
##
##
    [3] "F3D1 S189 L001 R1 001.fastq"
##
    [4] "F3D1_S189_L001_R2_001.fastq"
##
    [5] "F3D141_S207_L001_R1_001.fastq"
       "F3D141 S207 L001 R2 001.fastq"
##
       "F3D142 S208 L001 R1 001.fastq"
##
       "F3D142_S208_L001_R2_001.fastq"
##
    [8]
##
    [9]
       "F3D143_S209_L001_R1_001.fastq"
##
   [10] "F3D143_S209_L001_R2_001.fastq"
   [11] "F3D144_S210_L001_R1_001.fastq"
   [12] "F3D144_S210_L001_R2_001.fastq"
##
   [13]
       "F3D145_S211_L001_R1_001.fastq"
       "F3D145_S211_L001_R2_001.fastq"
   [15] "F3D146_S212_L001_R1_001.fastq"
   [16] "F3D146_S212_L001_R2_001.fastq"
   [17] "F3D147_S213_L001_R1_001.fastq"
##
   [18] "F3D147 S213 L001 R2 001.fastg"
   [19] "F3D148_S214_L001_R1_001.fastq"
##
   [20] "F3D148_S214_L001_R2_001.fastq"
##
  [21]
       "F3D149_S215_L001_R1_001.fastq"
       "F3D149 S215 L001 R2 001.fastq"
  [23] "F3D150_S216_L001_R1_001.fastq"
##
   [24] "F3D150 S216 L001 R2 001.fastq"
##
   [25] "F3D2 S190 L001 R1 001.fastq"
##
   [26] "F3D2_S190_L001_R2_001.fastq"
   [27] "F3D3_S191_L001_R1_001.fastq"
##
   [28]
       "F3D3_S191_L001_R2_001.fastq"
   [29]
       "F3D5_S193_L001_R1_001.fastq"
##
   [30]
       "F3D5_S193_L001_R2_001.fastq"
##
   [31]
        "F3D6_S194_L001_R1_001.fastq"
##
   [32]
        "F3D6_S194_L001_R2_001.fastq"
   [33]
       "F3D7_S195_L001_R1_001.fastq"
       "F3D7_S195_L001_R2_001.fastq"
##
   [34]
   [35]
        "F3D8_S196_L001_R1_001.fastq"
##
   [36]
       "F3D8_S196_L001_R2_001.fastq"
##
       "F3D9 S197 L001 R1 001.fastq"
  [38] "F3D9_S197_L001_R2_001.fastq"
##
   [39]
        "HMP MOCK.v35.fasta"
##
   [40]
       "Mock_S280_L001_R1_001.fastq"
##
       "Mock S280 L001 R2 001.fastq"
   [41]
   [42]
       "mouse.dpw.metadata"
##
##
   Γ431
       "mouse.time.design"
   [44]
       "silva_nr_v132_train_set.fa.gz"
##
       "silva_species_assignment_v132.fa.gz"
  [45]
        "SILVA_SSU_r132_March2018.RData"
   [46]
   [47]
       "stability.batch"
## [48] "stability.files"
```