

# Prokaryotic biomass and density database

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```
#ProkaBIO database ## Loading benthic prokaryote biomass dataset
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```

library(tidyverse)
library(ggplot2)
library(scales)
library(openxlsx)

setwd("C:/Users/Mama/Desktop/manuscript revised/Figures")
prokaryotic_biomass_data <- read.xlsx("ProkaBio database.xlsx") # read xlsx
file
head(prokaryotic_biomass_data)

##      Region      Ocean Latitude.(N) Longitude.(E)      Location
Depth.(m)
## 1 North Sea Atlantic Ocean          53        3.9 exact location
24
## 2 North Sea Atlantic Ocean          53        3.9 exact location
24
## 3 North Sea Atlantic Ocean          53        3.9 exact location
24
## 4 North Sea Atlantic Ocean          53        3.9 exact location
24
## 5 North Sea Atlantic Ocean          53        3.9 exact location
24
## 6 North Sea Atlantic Ocean          53        3.9 exact location
24
##            Depth.range upper.--lower.sediment.layers.(cm)
## 1 near-shore (0-50m)                0.0-0.3
## 2 near-shore (0-50m)                0.0-0.3
## 3 near-shore (0-50m)                0.0-0.3
## 4 near-shore (0-50m)                12.0-12.3
## 5 near-shore (0-50m)                3.0-3.3
## 6 near-shore (0-50m)                3.0-3.3
##      middle.depth.of.sampled.sediment.layer.(cm) sediment.type
## 1                               0.15      <2% mud
## 2                               0.15      <2% mud
## 3                               0.15      <2% mud
## 4                             12.20      <2% mud
## 5                               3.15      <2% mud
## 6                               3.15      <2% mud
##      median.sediment.grain.size.(µm) sediment.density.(g.cm-3) porosity
## 1                               225                  NA     <NA>

```

```

## 2 225 NA <NA>
## 3 225 NA <NA>
## 4 225 NA <NA>
## 5 225 NA <NA>
## 6 225 NA <NA>
## type.of.microorganisms
## 1 Prokaryote
## 2 Prokaryote
## 3 Prokaryote
## 4 Prokaryote
## 5 Prokaryote
## 6 Prokaryote
##
mean/.median.density.per.sediment.layer.as.presented.in.the.original.publication
## 1
54
## 2
52
## 3
32
## 4
50
## 5
21
## 6
29
## unit.of.mean/.median.as.presented.in.original.publication
## 1 µg C cm-3 wet sediment
## 2 µg C cm-3 wet sediment
## 3 µg C cm-3 wet sediment
## 4 µg C cm-3 wet sediment
## 5 µg C cm-3 wet sediment
## 6 µg C cm-3 wet sediment
## error.term.as.presented.in.original.publication
## 1 NA
## 2 NA
## 3 NA
## 4 NA
## 5 NA
## 6 NA
## type.of.error.term.as.presented.in.the.original.publication
## 1 <NA>
## 2 <NA>
## 3 <NA>
## 4 <NA>
## 5 <NA>
## 6 <NA>
## standardized.biomass.per.sediment.layer standardized.unit
## 1 5.4e-05 g C cm-3 wet sediment

```

```

## 2           5.2e-05 g C cm-3 wet sediment
## 3           3.2e-05 g C cm-3 wet sediment
## 4           5.0e-05 g C cm-3 wet sediment
## 5           2.1e-05 g C cm-3 wet sediment
## 6           2.9e-05 g C cm-3 wet sediment
## conversion.factor.to.convert.PD.to.PB/.ATP.to.PB
## 1           2.2*10^(-11) g C μm-3 biovolume
## 2           2.2*10^(-11) g C μm-3 biovolume
## 3           2.2*10^(-11) g C μm-3 biovolume
## 4           2.2*10^(-11) g C μm-3 biovolume
## 5           2.2*10^(-11) g C μm-3 biovolume
## 6           2.2*10^(-11) g C μm-3 biovolume
##
reference.for.conversion.factor
## 1 Bratbak, G., & Dundas, I. (1984). Bacterial dry matter content and
biomass estimations. Applied and Environmental Microbiology, 48(4), 755-757.
https://doi.org/10.1128/aem.48.4.755-757.1984
## 2 Bratbak, G., & Dundas, I. (1984). Bacterial dry matter content and
biomass estimations. Applied and Environmental Microbiology, 48(4), 755-757.
https://doi.org/10.1128/aem.48.4.755-757.1984
## 3 Bratbak, G., & Dundas, I. (1984). Bacterial dry matter content and
biomass estimations. Applied and Environmental Microbiology, 48(4), 755-757.
https://doi.org/10.1128/aem.48.4.755-757.1984
## 4 Bratbak, G., & Dundas, I. (1984). Bacterial dry matter content and
biomass estimations. Applied and Environmental Microbiology, 48(4), 755-757.
https://doi.org/10.1128/aem.48.4.755-757.1984
## 5 Bratbak, G., & Dundas, I. (1984). Bacterial dry matter content and
biomass estimations. Applied and Environmental Microbiology, 48(4), 755-757.
https://doi.org/10.1128/aem.48.4.755-757.1984
## 6 Bratbak, G., & Dundas, I. (1984). Bacterial dry matter content and
biomass estimations. Applied and Environmental Microbiology, 48(4), 755-757.
https://doi.org/10.1128/aem.48.4.755-757.1984
## integrated.sediment.layers.(lower.--.upper;.cm)
## 1           0.0-12.3
## 2           0.0-6.3
## 3           0.0-6.3
## 4           <NA>
## 5           <NA>
## 6           <NA>
## integrated.sediment.layer/.surface.sediment.layer
## 1           integrated sediment
## 2           integrated sediment
## 3           integrated sediment
## 4           <NA>
## 5           <NA>
## 6           <NA>
## integrated.sediment.depth.(cm)
## 1           12.3
## 2           6.3
## 3           6.3

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```

## 4 NA
## 5 NA
## 6 NA
## integrated.standardized.biomass.per.station/.sampling.event
## 1 0.000186
## 2 0.000091
## 3 0.000139
## 4 NA
## 5 NA
## 6 NA
## integrated.standardized.unit
## 1 g C cm-3 wet sediment
## 2 g C cm-3 wet sediment
## 3 g C cm-3 wet sediment
## 4 <NA>
## 5 <NA>
## 6 <NA>
##
method
## 1 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors
## 2 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors
## 3 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors
## 4 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors
## 5 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors
## 6 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors
## substrate.type
## 1 sediment
## 2 sediment
## 3 sediment
## 4 sediment
## 5 sediment
## 6 sediment
##
Reference
## 1 Osinga, R., Kop, A. J., Duineveld, G. C. A., Prins, R. A., & Van Duyl,
F. C. (1996). Benthic mineralization rates at two locations in the southern
North Sea. Journal of Sea Research, 36(3-4), 181-191.
https://doi.org/10.1016/S1385-1101\(96\)90788-1
## 2 Osinga, R., Kop, A. J., Duineveld, G. C. A., Prins, R. A., & Van Duyl,
F. C. (1996). Benthic mineralization rates at two locations in the southern
North Sea. Journal of Sea Research, 36(3-4), 181-191.
https://doi.org/10.1016/S1385-1101\(96\)90788-1
## 3 Osinga, R., Kop, A. J., Duineveld, G. C. A., Prins, R. A., & Van Duyl,
F. C. (1996). Benthic mineralization rates at two locations in the southern

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North Sea. Journal of Sea Research, 36(3-4), 181-191.
https://doi.org/10.1016/S1385-1101\(96\)90788-1
## 4 Osinga, R., Kop, A. J., Duineveld, G. C. A., Prins, R. A., & Van Duyl, F. C. (1996). Benthic mineralization rates at two locations in the southern North Sea. Journal of Sea Research, 36(3-4), 181-191.
https://doi.org/10.1016/S1385-1101\(96\)90788-1
## 5 Osinga, R., Kop, A. J., Duineveld, G. C. A., Prins, R. A., & Van Duyl, F. C. (1996). Benthic mineralization rates at two locations in the southern North Sea. Journal of Sea Research, 36(3-4), 181-191.
https://doi.org/10.1016/S1385-1101\(96\)90788-1
## 6 Osinga, R., Kop, A. J., Duineveld, G. C. A., Prins, R. A., & Van Duyl, F. C. (1996). Benthic mineralization rates at two locations in the southern North Sea. Journal of Sea Research, 36(3-4), 181-191.
https://doi.org/10.1016/S1385-1101\(96\)90788-1

prokaryotic_biomass_data <- prokaryotic_biomass_data[c(2:4, 6:7, 19:20, 28)]
cols <- c("Ocean", "Latitude", "Longitude", "Depth", "Depth_range", "Biomass", "Unit", "Method")
colnames(prokaryotic_biomass_data) <- cols

prokaryotic_biomass_corrected <-
prokaryotic_biomass_data[complete.cases(prokaryotic_biomass_data$Biomass), ]
#prokaryotic_biomass_corrected <- prokaryotic_biomass_data %>%
drop_na("Biomass")
head(prokaryotic_biomass_corrected)

##          Ocean Latitude Longitude Depth      Depth_range Biomass
## 1 Atlantic Ocean      53     3.9    24 near-shore (0-50m) 5.4e-05
## 2 Atlantic Ocean      53     3.9    24 near-shore (0-50m) 5.2e-05
## 3 Atlantic Ocean      53     3.9    24 near-shore (0-50m) 3.2e-05
## 4 Atlantic Ocean      53     3.9    24 near-shore (0-50m) 5.0e-05
## 5 Atlantic Ocean      53     3.9    24 near-shore (0-50m) 2.1e-05
## 6 Atlantic Ocean      53     3.9    24 near-shore (0-50m) 2.9e-05
##                  Unit
## 1 g C cm-3 wet sediment
## 2 g C cm-3 wet sediment
## 3 g C cm-3 wet sediment
## 4 g C cm-3 wet sediment
## 5 g C cm-3 wet sediment
## 6 g C cm-3 wet sediment
##
Method
## 1 PD measured by epifluorescence microscopy, PD converted to PB using conversion factors
## 2 PD measured by epifluorescence microscopy, PD converted to PB using conversion factors
## 3 PD measured by epifluorescence microscopy, PD converted to PB using conversion factors
## 4 PD measured by epifluorescence microscopy, PD converted to PB using conversion factors

```

```

## 5 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors
## 6 PD measured by epifluorescence microscopy, PD converted to PB using
conversion factors

```

## Preparing figure

```

knitr::opts_chunk$set(fig.width=13, fig.height=6.5)

#along a latitudinal gradient
lab1 <- c(expression(g~C~cm^-3~wet~sediment),
           expression(g~C~g~dry~sediment^-1),
           expression(g~C~g~wet~sediment^-1),
           expression(g~C~m^-2))

Fig1 <- ggplot(data = prokaryotic_biomass_corrected, aes(x = Biomass, y =
Latitude, color=Unit)) + geom_point(alpha = 0.3, size=2.5, shape=16) +
theme_classic() + scale_y_continuous(breaks = c(-90, -75, -60, -45, -30, -15,
0, 15, 30, 45, 60, 75, 90)) + ylab("° Latitude") + xlab("Prokaryotic
biomass") +
  theme(axis.text.y = element_text(vjust=0.5, size=15)) +
  theme(text = element_text(size=20)) + geom_hline(yintercept = 0,
linetype="dashed") + theme(legend.position = "bottom", legend.title =
element_blank()) + guides(col = guide_legend(nrow=1)) +
  theme(plot.margin=unit(c(0.5,0,0,0.5), "cm")) +
  scale_colour_manual(labels = lab1, values = c('red','orange', 'black',
'blue')) +
  scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x), labels =
trans_format("log10", math_format(10^.x)), position = "bottom", limits =
c(1e-4,1e4)) + annotation_logticks(sides = "b")

ggsave(file="prokaryotic biomass vs latitude.png", width=7,height=6.5)

## Warning: Removed 612 rows containing missing values (geom_point).

#along a depth gradient
lab1 <- c(expression(g~C~cm^-3~wet~sediment),
           expression(g~C~g~dry~sediment^-1),
           expression(g~C~g~wet~sediment^-1),
           expression(g~C~m^-2))

reverselog_trans <- function(base = exp(1)) {
  trans <- function(y) -log(y, base)
  inv <- function(y) base^(-y)
  trans_new(paste0("reverselog-", format(base)), trans, inv,
            log_breaks(base = base),
            domain = c(1e-100, Inf))
}

Fig2 <- ggplot(data = prokaryotic_biomass_corrected, aes(x = Biomass, y =
Depth, color=Unit)) + geom_point(alpha = 0.3, size=2.5, shape=16) +

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```

theme_classic() + scale_y_continuous(trans=reverselog_trans(10), breaks =
c(1, 10, 100, 1000, 10000))+ ylab("Depth (m)") + xlab("Prokaryotic biomass")
+
  theme(axis.text.y = element_text(vjust=0.5, size=15)) +
annotation_logticks(sides = "tl") + geom_hline(yintercept = 0.1,
linetype="dashed", col="white", alpha=0) +
  theme(text = element_text(size=20)) + guides(fill=guide_legend(nrow=1)) +
guides(col = guide_legend(nrow=2)) + theme(legend.position = "bottom",
legend.title = element_blank()) + theme(plot.margin=unit(c(0.5,0,0,0.5),
"cm")) +
  scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x),
labels = trans_format("log10", math_format(10^.x)), limits =
c(1e-4,1e4), position = "top") + scale_colour_manual(labels = lab1, values =
c('red','orange','black','blue'))
ggsave(file="prokaryotic biomass vs depth.png", width=7,height=6.5)

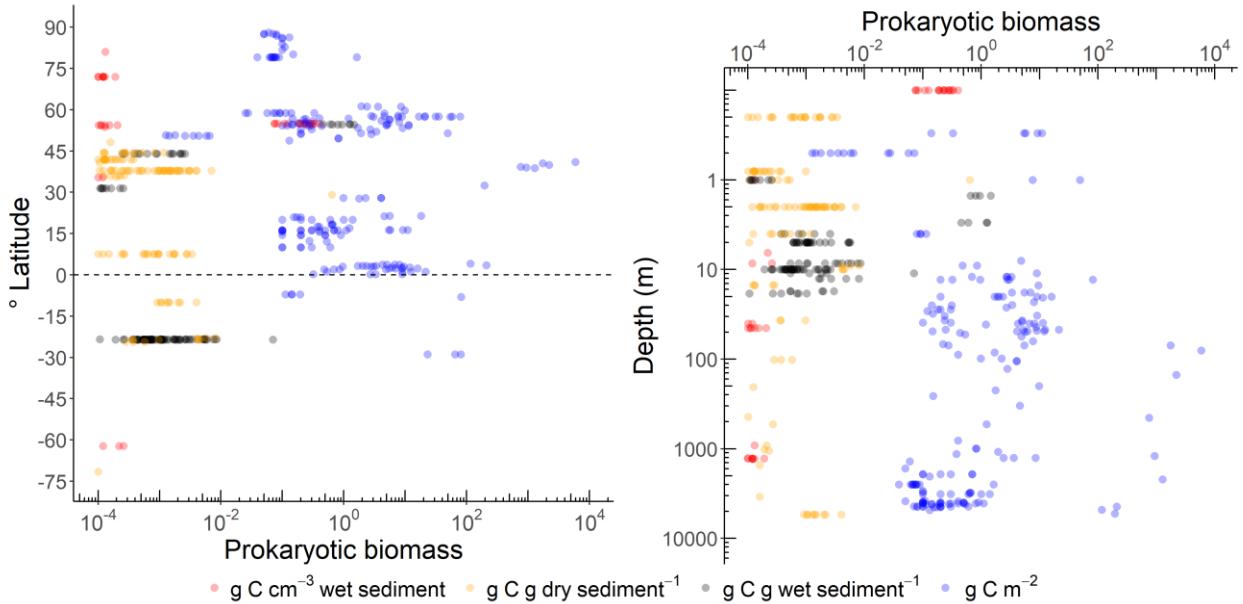
## Warning: Removed 629 rows containing missing values (geom_point).

ggpubr::ggarrange(Fig1, Fig2, common.legend = TRUE, legend = "bottom")

## Warning: Removed 612 rows containing missing values (geom_point).
## Removed 612 rows containing missing values (geom_point).

## Warning: Removed 629 rows containing missing values (geom_point).

```



```

setwd("C:/Users/Mama/Desktop/manuscript revised/Figures")
ggsave(file="prokaryotic biomass.png", width=13,height=6.5)

```

## Loading benthic prokaryote density dataset

```

library(tidyr)
library(ggplot2)

```

```

library(openxlsx)
library(dplyr)

##
## Attache Paket: 'dplyr'

## Die folgenden Objekte sind maskiert von 'package:stats':
##
##     filter, lag

## Die folgenden Objekte sind maskiert von 'package:base':
##
##     intersect, setdiff, setequal, union

setwd("C:/Users/Mama/Desktop/manuscript revised/Figures")
prokaryotic_density_data <- read.xlsx("ProkaDen database.xlsx") # read xlsx
file
head(prokaryotic_density_data)

##           Region          Ocean Latitude.(N) Longitude.(E)      Location
## 1 Adriatic Sea Mediterranean Sea    42.866     15.095 exact location
## 2 Adriatic Sea Mediterranean Sea    42.874     15.092 exact location
## 3 Adriatic Sea Mediterranean Sea    42.903     15.069 exact location
## 4 Adriatic Sea Mediterranean Sea    42.907     15.072 exact location
## 5 Adriatic Sea Mediterranean Sea    42.911     15.062 exact location
## 6 Adriatic Sea Mediterranean Sea    43.600     13.500 exact location
##   Depth.(m)      Depth.range upper.-
## 1.0 .lower.sediment.layers.(cm)
## 1.0 ## 1     189 continental shelf (>50-200m)             0.0-
## 1.0 ## 2     194 continental shelf (>50-200m)             0.0-
## 1.0 ## 3     190 continental shelf (>50-200m)             0.0-
## 1.0 ## 4     190 continental shelf (>50-200m)             0.0-
## 1.0 ## 5     197 continental shelf (>50-200m)             0.0-
## 1.0 ## 6       5 near-shore (0-50m)                   0.0-
##   middle.depth.of.sampled.sediment.layer.(cm) sediment.type
## 1.0 ## 1                           0.5 <NA>
## 1.0 ## 2                           0.5 <NA>
## 1.0 ## 3                           0.5 <NA>
## 1.0 ## 4                           0.5 <NA>
## 1.0 ## 5                           0.5 <NA>
## 1.0 ## 6                           0.5 <NA>
##   median.sediment.grain.size.(µm) sediment.density.(g.cm-3) porosity
## 1.0 ## 1                         NA NA
## 1.0 ## 2                         NA NA
## 1.0 ## 3                         NA NA

```

```

## 4 NA NA NA
## 5 NA NA NA
## 6 NA NA NA
## type.of.microorganisms
## 1 Prokaryote
## 2 Prokaryote
## 3 Prokaryote
## 4 Prokaryote
## 5 Prokaryote
## 6 Prokaryote
##
mean/.median.density.per.sediment.layer.as.presented.in.the.original.publication
## 1
3.35e+08
## 2
3.38e+08
## 3
1.18e+08
## 4
2.31e+08
## 5
1.43e+08
## 6
5.20e+08
## unit.of.mean/.median.as.presented.in.original.publication
## 1 cell g-1 dry sediment
## 2 cell g-1 dry sediment
## 3 cell g-1 dry sediment
## 4 cell g-1 dry sediment
## 5 cell g-1 dry sediment
## 6 cell g-1 dry sediment
## error.term.as.presented.in.original.publication
## 1 61674009
## 2 10572687
## 3 31718062
## 4 33480176
## 5 12334802
## 6 NA
## type.of.error.term.as.presented.in.the.original.publication
## 1 standard deviation
## 2 standard deviation
## 3 standard deviation
## 4 standard deviation
## 5 standard deviation
## 6 <NA>
## standardized.density.per.sediment.layer standardized.unit
## 1 3.35e+08 cell g-1 dry sediment
## 2 3.38e+08 cell g-1 dry sediment
## 3 1.18e+08 cell g-1 dry sediment

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```

## 4          2.31e+08 cell g-1 dry sediment
## 5          1.43e+08 cell g-1 dry sediment
## 6          5.20e+08 cell g-1 dry sediment
## integrated.sediment.layers.(lower.-.upper;.cm)
## 1          0.0-1.0
## 2          0.0-1.0
## 3          0.0-1.0
## 4          0.0-1.0
## 5          0.0-1.0
## 6          0.0-1.0
## integrated.sediment.layer/.surface.sediment.layer
## 1          surface sediment
## 2          surface sediment
## 3          surface sediment
## 4          surface sediment
## 5          surface sediment
## 6          surface sediment
## integrated.sediment.depth.(cm)
## 1          1
## 2          1
## 3          1
## 4          1
## 5          1
## 6          1
## integrated.standardized.density.per.station/.sampling.event
## 1          3.35e+08
## 2          3.38e+08
## 3          1.18e+08
## 4          2.31e+08
## 5          1.43e+08
## 6          5.20e+08
## integrated.standardized.unit
## 1          cell g-1 dry sediment
## 2          cell g-1 dry sediment
## 3          cell g-1 dry sediment
## 4          cell g-1 dry sediment
## 5          cell g-1 dry sediment
## 6          cell g-1 dry sediment
##                                     method substrate.type
## 1 epifluorescence microscopy (stain: acridine orange)    sediment
## 2 epifluorescence microscopy (stain: acridine orange)    sediment
## 3 epifluorescence microscopy (stain: acridine orange)    sediment
## 4 epifluorescence microscopy (stain: acridine orange)    sediment
## 5 epifluorescence microscopy (stain: acridine orange)    sediment
## 6 epifluorescence microscopy (stain: acridine orange)    sediment
##
Reference
## 1      Giovannelli, D., D'Errico, G., Fiorentino, F., Fattorini, D.,
Regoli, F., Angeletti, L., et al. (2016). Diversity and distribution of
prokaryotes within a shallow-water pockmark field. Frontiers in Microbiology,

```

```

7, 941. https://doi.org/10.3389/fmicb.2016.00941
## 2 Giovannelli, D., D'Errico, G., Fiorentino, F., Fattorini, D.,
Regoli, F., Angeletti, L., et al. (2016). Diversity and distribution of
prokaryotes within a shallow-water pockmark field. Frontiers in Microbiology,
7, 941. https://doi.org/10.3389/fmicb.2016.00941
## 3 Giovannelli, D., D'Errico, G., Fiorentino, F., Fattorini, D.,
Regoli, F., Angeletti, L., et al. (2016). Diversity and distribution of
prokaryotes within a shallow-water pockmark field. Frontiers in Microbiology,
7, 941. https://doi.org/10.3389/fmicb.2016.00941
## 4 Giovannelli, D., D'Errico, G., Fiorentino, F., Fattorini, D.,
Regoli, F., Angeletti, L., et al. (2016). Diversity and distribution of
prokaryotes within a shallow-water pockmark field. Frontiers in Microbiology,
7, 941. https://doi.org/10.3389/fmicb.2016.00941
## 5 Giovannelli, D., D'Errico, G., Fiorentino, F., Fattorini, D.,
Regoli, F., Angeletti, L., et al. (2016). Diversity and distribution of
prokaryotes within a shallow-water pockmark field. Frontiers in Microbiology,
7, 941. https://doi.org/10.3389/fmicb.2016.00941
## 6 Danovaro, R., Armeni, M., Dell'Anno, A., Fabiano, M., Manini, E.,
Marrale, D., et al. (2001). Small-scale distribution of bacteria, enzymatic
activities, and organic matter in coastal sediments. Microbial Ecology,
42(2), 177–185. https://doi.org/10.1007/s002480000109

prokaryotic_density_data <- prokaryotic_density_data[c(2:4, 6:7, 14, 19:20,
26)]
cols <- c("Ocean", "Latitude", "Longitude", "Depth", "Depth_range", "Type",
"Density", "Unit", "Method")
colnames(prokaryotic_density_data) <- cols

prokaryotic_density_corrected <- prokaryotic_density_data %>%
drop_na("Density")
head(prokaryotic_density_corrected)

##          Ocean Latitude Longitude Depth      Depth_range
## 1 Mediterranean Sea    42.866    15.095   189 continental shelf (>50-200m)
## 2 Mediterranean Sea    42.874    15.092   194 continental shelf (>50-200m)
## 3 Mediterranean Sea    42.903    15.069   190 continental shelf (>50-200m)
## 4 Mediterranean Sea    42.907    15.072   190 continental shelf (>50-200m)
## 5 Mediterranean Sea    42.911    15.062   197 continental shelf (>50-200m)
## 6 Mediterranean Sea    43.600    13.500      5             near-shore (0-50m)
##           Type Density          Unit
## 1 Prokaryote 3.35e+08 cell g-1 dry sediment
## 2 Prokaryote 3.38e+08 cell g-1 dry sediment
## 3 Prokaryote 1.18e+08 cell g-1 dry sediment
## 4 Prokaryote 2.31e+08 cell g-1 dry sediment
## 5 Prokaryote 1.43e+08 cell g-1 dry sediment
## 6 Prokaryote 5.20e+08 cell g-1 dry sediment
##           Method
## 1 epifluorescence microscopy (stain: acridine orange)
## 2 epifluorescence microscopy (stain: acridine orange)
## 3 epifluorescence microscopy (stain: acridine orange)

```

```

## 4 epifluorescence microscopy (stain: acridine orange)
## 5 epifluorescence microscopy (stain: acridine orange)
## 6 epifluorescence microscopy (stain: acridine orange)

##Divide datasets in Prokaryote, Bacteria, Archaea

Prokaryote_data <- prokaryotic_density_corrected %>% filter(Type ==
"Prokaryote")
Pro_data <- Prokaryote_data[!(Prokaryote_data$Density==0),]
head(Pro_data)

##          Ocean Latitude Longitude Depth             Depth_range
## 1 Mediterranean Sea    42.866    15.095   189 continental shelf (>50-200m)
## 2 Mediterranean Sea    42.874    15.092   194 continental shelf (>50-200m)
## 3 Mediterranean Sea    42.903    15.069   190 continental shelf (>50-200m)
## 4 Mediterranean Sea    42.907    15.072   190 continental shelf (>50-200m)
## 5 Mediterranean Sea    42.911    15.062   197 continental shelf (>50-200m)
## 6 Mediterranean Sea    43.600    13.500      5             near-shore (0-50m)
##          Type Density           Unit
## 1 Prokaryote 3.35e+08 cell g-1 dry sediment
## 2 Prokaryote 3.38e+08 cell g-1 dry sediment
## 3 Prokaryote 1.18e+08 cell g-1 dry sediment
## 4 Prokaryote 2.31e+08 cell g-1 dry sediment
## 5 Prokaryote 1.43e+08 cell g-1 dry sediment
## 6 Prokaryote 5.20e+08 cell g-1 dry sediment
##                  Method
## 1 epifluorescence microscopy (stain: acridine orange)
## 2 epifluorescence microscopy (stain: acridine orange)
## 3 epifluorescence microscopy (stain: acridine orange)
## 4 epifluorescence microscopy (stain: acridine orange)
## 5 epifluorescence microscopy (stain: acridine orange)
## 6 epifluorescence microscopy (stain: acridine orange)

Bacteria_data <- prokaryotic_density_corrected %>% filter(Type == "Bacteria")
head(Bacteria_data)

##          Ocean Latitude Longitude Depth
Depth_range
## 1 Mediterranean Sea    32.368    31.704   1158 continental slope (>200-
2000m)
## 2 Mediterranean Sea    32.369    31.711   1122 continental slope (>200-
2000m)
## 3 Mediterranean Sea    32.369    31.711   1122 continental slope (>200-
2000m)
## 4 Mediterranean Sea    32.368    31.704   1158 continental slope (>200-
2000m)
## 5 Mediterranean Sea    32.368    31.704   1158 continental slope (>200-
2000m)
## 6 Mediterranean Sea    32.369    31.711   1122 continental slope (>200-
2000m)
##          Type Density           Unit
##                  Method

```

```

## 1 Bacteria 1.12e+09 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 2 Bacteria 3.87e+09 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 3 Bacteria 8.90e+08 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 4 Bacteria 9.40e+08 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 5 Bacteria 7.00e+07 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 6 Bacteria 3.10e+08 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)

Bac_data <- Bacteria_data[!(Bacteria_data$Density==0),]
head(Bac_data)

##          Ocean Latitude Longitude Depth
Depth_range
## 1 Mediterranean Sea   32.368   31.704  1158 continental slope (>200-
2000m)
## 2 Mediterranean Sea   32.369   31.711  1122 continental slope (>200-
2000m)
## 3 Mediterranean Sea   32.369   31.711  1122 continental slope (>200-
2000m)
## 4 Mediterranean Sea   32.368   31.704  1158 continental slope (>200-
2000m)
## 5 Mediterranean Sea   32.368   31.704  1158 continental slope (>200-
2000m)
## 6 Mediterranean Sea   32.369   31.711  1122 continental slope (>200-
2000m)
##      Type Density           Unit           Method
## 1 Bacteria 1.12e+09 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 2 Bacteria 3.87e+09 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 3 Bacteria 8.90e+08 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 4 Bacteria 9.40e+08 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 5 Bacteria 7.00e+07 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)
## 6 Bacteria 3.10e+08 cell cm-3 wet sediment CARD-FISH (probe: EUB338I-III)

Archaea_data <- prokaryotic_density_corrected %>% filter(Type == "Archaea")
head(Archaea_data)

##          Ocean Latitude Longitude Depth
Depth_range
## 1 Mediterranean Sea   32.369   31.710  1120 continental slope (>200-
2000m)
## 2 Mediterranean Sea   32.368   31.704  1158 continental slope (>200-
2000m)
## 3 Mediterranean Sea   32.369   31.711  1122 continental slope (>200-
2000m)
## 4 Mediterranean Sea   32.368   31.704  1158 continental slope (>200-
2000m)
## 5 Mediterranean Sea   32.369   31.710  1120 continental slope (>200-
2000m)
## 6 Mediterranean Sea   32.368   31.704  1158 continental slope (>200-
2000m)
##      Type Density           Unit           Method
## 1 Archaea 5.0e+07 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)

```

```

## 2 Archaea 4.0e+06 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 3 Archaea 5.1e+08 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 4 Archaea 0.0e+00 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 5 Archaea 1.0e+07 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 6 Archaea 3.0e+06 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)

Arc_data <- Archaea_data[!(Archaea_data$Density==0),]
head(Arc_data)

##          Ocean Latitude Longitude Depth
Depth_range
## 1 Mediterranean Sea   32.369   31.710 1120 continental slope (>200-
2000m)
## 2 Mediterranean Sea   32.368   31.704 1158 continental slope (>200-
2000m)
## 3 Mediterranean Sea   32.369   31.711 1122 continental slope (>200-
2000m)
## 5 Mediterranean Sea   32.369   31.710 1120 continental slope (>200-
2000m)
## 6 Mediterranean Sea   32.368   31.704 1158 continental slope (>200-
2000m)
## 7 Mediterranean Sea   32.369   31.711 1122 continental slope (>200-
2000m)
##      Type Density           Unit           Method
## 1 Archaea 5.0e+07 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 2 Archaea 4.0e+06 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 3 Archaea 5.1e+08 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 5 Archaea 1.0e+07 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 6 Archaea 3.0e+06 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)
## 7 Archaea 1.7e+08 cell cm-3 wet sediment CARD-FISH (probe: ARCH915)

```

## Preparing figure: Microbial densities along a latitudinal gradient

```

#Plot for prokaryotic density
lab1 <- c(expression(cell~cm^-3~dry~sediment),
           expression(cell~cm^-3~wet~sediment),
           expression(cell~g^-1~dry~sediment),
           expression(cell~g^-1~dry~sulfide),
           expression(cell~g^-1~vent~precipitate),
           expression(cell~g^-1~wet~sediment),
           expression(cell~m^-2))

Fig3 <- ggplot(data = Pro_data, aes(x = Density, y = Latitude, color=Unit)) +
  geom_point(alpha = 0.3, size=2.5, shape=16) + theme_classic() +
  scale_y_continuous(breaks = c(-90, -75, -60, -45, -30, -15, 0, 15, 30, 45,
60, 75, 90)) + ylab("° Latitude") +
  theme(axis.text.y = element_text(vjust=0.5, size=15)) +
  theme(text = element_text(size=20)) + geom_hline(yintercept = 0,
linetype="dashed") +
  scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x),
               labels = trans_format("log10", math_format(10^.x))), limits =

```

```

c(1e4,1e16)) + theme(legend.position = "bottom", legend.title =
element_blank()) + guides(col = guide_legend(nrow=4)) +
theme(plot.margin=unit(c(0.5,0,0,0.5), "cm")) +
scale_colour_manual(labels = lab1, values = c('blue', 'red','orange',
'black', '#8918FA', '1CF339', '#1CD7F3')) + annotate(geom="text", x=1e14,
y=100, label="Prokaryotes", color="black", size=6) +
annotation_logticks(sides = "b")

ggsave(file="prokaryotic density vs latitude.png", width=6.5,height=6.5)

#Plot for Bacteria density
lab1 <- c(expression(cell~cm^-3~dry~sediment),
expression(cell~cm^-3~wet~sediment),
expression(cell~g^-1~dry~sediment),
expression(cell~g^-1~dry~sulfide),
expression(cell~g^-1~vent~precipitate),
expression(cell~g^-1~wet~sediment),
expression(cell~m^-2))

Fig4 <- ggplot(data = Bac_data, aes(x = Density, y = Latitude, color=Unit)) +
geom_point(alpha = 0.3, size=2.5, shape=16) + theme_classic() +
scale_y_continuous(breaks = c(-90, -75, -60, -45, -30, -15, 0, 15, 30, 45,
60, 75, 90)) + ylab("° Latitude") +
theme(axis.text.y = element_text(vjust=0.5, size=15)) +
theme(text = element_text(size=20)) + geom_hline(yintercept = 0,
linetype="dashed") +
scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x),
labels = trans_format("log10", math_format(10^.x)), limits =
c(1e4,1e16)) + theme(legend.position = "bottom", legend.title =
element_blank()) + guides(col = guide_legend(nrow=4)) +
theme(plot.margin=unit(c(0.5,0,0,0.5), "cm")) +
scale_colour_manual(labels = lab1, values = c('blue', 'red','orange',
'black', '#8918FA', '1CF339', '#1CD7F3')) + annotate(geom="text", x=1e14,
y=100, label="Bacteria", color="black", size=6) + annotation_logticks(sides =
"b")

ggsave(file="Bacteria density vs latitude.png", width=6.5,height=6.5)

#Plot for Archaea density
lab1 <- c(expression(cell~cm^-3~dry~sediment),
expression(cell~cm^-3~wet~sediment),
expression(cell~g^-1~dry~sediment),
expression(cell~g^-1~dry~sulfide),
expression(cell~g^-1~vent~precipitate),
expression(cell~g^-1~wet~sediment),
expression(cell~m^-2))

Fig5 <- ggplot(data = Arc_data, aes(x = Density, y = Latitude, color=Unit)) +
geom_point(alpha = 0.3, size=2.5, shape=16) + theme_classic() +
scale_y_continuous(breaks = c(-90, -75, -60, -45, -30, -15, 0, 15, 30, 45,

```

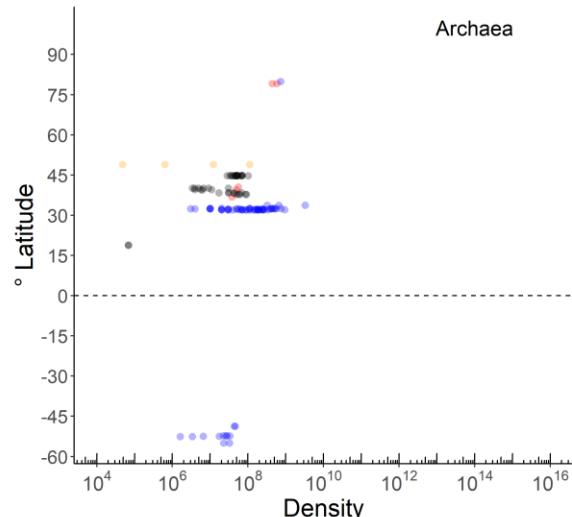
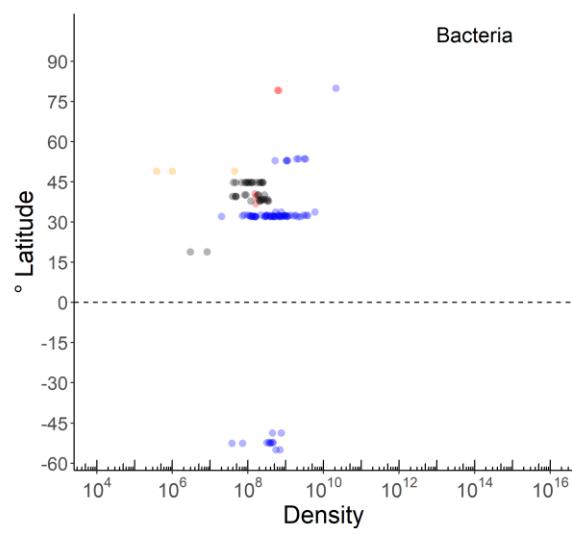
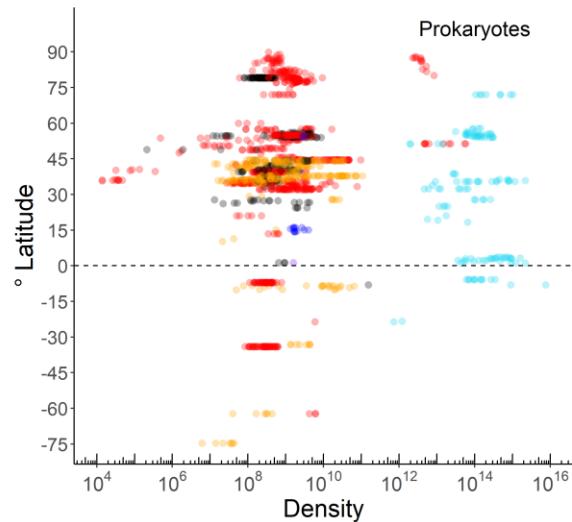
```

60, 75, 90)) + ylab("° Latitude") +
  theme(axis.text.y = element_text(vjust=0.5, size=15)) +
  theme(text = element_text(size=20)) + geom_hline(yintercept = 0,
linetype="dashed") +
  scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x),
                labels = trans_format("log10", math_format(10^.x)), limits =
c(1e4,1e16)) + theme(legend.position = "bottom", legend.title =
element_blank()) + guides(col = guide_legend(nrow=4)) +
  theme(plot.margin=unit(c(0.5,0,0,0.5), "cm")) +
  scale_colour_manual(labels = lab1, values = c('blue', 'red','orange',
'black', '#8918FA', '1CF339', '#1CD7F3')) + annotate(geom="text", x=1e14,
y=100, label="Archaea", color="black", size=6) + annotation_logticks(sides =
"b")

ggsave(file="Archaea density vs latitude.png", width=6.5,height=6.5)

ggpubr::ggarrange(Fig3, Fig4, Fig5, ncol=1, common.legend = TRUE, legend =
"bottom")

```



- cell  $\text{cm}^{-3}$  dry sediment
- cell  $\text{cm}^{-3}$  wet sediment
- cell  $\text{g}^{-1}$  dry sediment
- cell  $\text{g}^{-1}$  dry sulfide
- cell  $\text{g}^{-1}$  vent precipitate
- cell  $\text{g}^{-1}$  wet sediment
- cell  $\text{m}^{-2}$

```
setwd("C:/Users/Mama/Desktop/manuscript revised/Figures")
ggsave(file="microbial density vs latitude.png", width=6.5, height=19.5)
```

## Preparing figure: Microbial densities along a latitudinal gradient

```
library(ggplot2)
library(openxlsx)

#Plot for prokaryotic density
lab1 <- c(expression(cell~cm^-3~dry~sediment),
           expression(cell~cm^-3~wet~sediment),
           expression(cell~g~dry~sediment^-1),
           expression(cell~g~dry~sulfide^-1),
           expression(cell~g~wet~sediment^-1),
           expression(cell~m^-2))

reverselog_trans <- function(base = exp(1)) {
  trans <- function(y) -log(y, base)
  inv <- function(y) base^(-y)
  trans_new(paste0("reverselog-", format(base)), trans, inv,
            log_breaks(base = base),
            domain = c(1e-100, Inf))
}

Fig6 <- ggplot(data = Pro_data, aes(x = Density, y = Depth, color=Unit)) +
  geom_point(alpha = 0.3, size=2.5, shape=16) + theme_classic() +
  scale_y_continuous(trans=reverselog_trans(10), breaks = c(1, 10, 100, 1000,
  10000)) + ylab("Depth (m)") + xlab("Density") + theme(axis.text.y =
  element_text(vjust=0.5, size=15)) + annotation_logticks(sides = "tl") +
  geom_hline(yintercept = 10000, linetype="dashed", col="white", alpha=0) +
  geom_hline(yintercept = 0.1, linetype="dashed", col="white", alpha=0) +
  theme(text = element_text(size=20)) + guides(col = guide_legend(nrow=3)) +
  scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x), labels =
  trans_format("log10", math_format(10^.x)), position = "top", limits =
  c(1e4,1e16)) + theme(legend.position = "bottom", legend.title =
  element_blank()) + theme(plot.margin=unit(c(0.5,0,0,0.5), "cm")) +
  annotate(geom="text", x=1e14, y=0.2, label="Prokaryotes", color="black",
  size=6) + scale_colour_manual(labels = lab1, values = c('blue',
  'red','orange', 'black', '#8918FA', '1CF339', '#1CD7F3'))

ggsave(file="prokaryotic density vs depth.png", width=6.5, height=6.5)

## Warning: Removed 33 rows containing missing values (geom_point).

#Plot for Bacteria density
Fig7 <- ggplot(data = Bacteria_data, aes(x = Density, y = Depth, color=Unit)) +
  geom_point(alpha = 0.3, size=2.5, shape=16) + theme_classic() +
  scale_y_continuous(trans=reverselog_trans(10), breaks = c(1, 10, 100, 1000,
  10000)) + ylab("Depth (m)") + xlab("Density") + theme(axis.text.y =
  element_text(vjust=0.5, size=15)) + annotation_logticks(sides = "tl") +
  geom_hline(yintercept = 10000, linetype="dashed", col="white", alpha=0) +
```

```

geom_hline(yintercept = 0.1, linetype="dashed", col="white", alpha=0) +
theme(text = element_text(size=20)) + guides(col = guide_legend(nrow=3)) +
scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x), labels =
trans_format("log10", math_format(10^.x)), position = "top", limits =
c(1e4,1e16)) + theme(legend.position = "bottom", legend.title =
element_blank()) + theme(plot.margin=unit(c(0.5,0,0,0.5), "cm")) +
annotate(geom="text", x=1e14, y=0.2, label="Bacteria", color="black", size=6)
+ scale_colour_manual(labels = lab1, values = c('blue', 'red','orange',
'black', '#8918FA', '1CF339', '#1CD7F3'))

ggsave(file="Bacteria density vs depth.png", width=6.5,height=6.5)

## Warning: Transformation introduced infinite values in continuous x-axis

#Plot for Archaea density
Fig8 <- ggplot(data = Archaea_data, aes(x = Density, y = Depth, color=Unit))
+ geom_point(alpha = 0.3, size=2.5, shape=16) + theme_classic() +
scale_y_continuous(trans=reverselog_trans(10), breaks = c(1, 10, 100, 1000,
10000)) + ylab("Depth (m)") + xlab("Density") + theme(axis.text.y =
element_text(vjust=0.5, size=15)) + annotation_logticks(sides = "tl") +
geom_hline(yintercept = 10000, linetype="dashed", col="white", alpha=0) +
geom_hline(yintercept = 0.1, linetype="dashed", col="white", alpha=0) +
theme(text = element_text(size=20)) + guides(col = guide_legend(nrow=3)) +
scale_x_log10(breaks = trans_breaks("log10", function(x) 10^x), labels =
trans_format("log10", math_format(10^.x)), position = "top", limits =
c(1e4,1e16)) + theme(legend.position = "bottom", legend.title =
element_blank()) + theme(plot.margin=unit(c(0.5,0,0,0.5), "cm")) +
annotate(geom="text", x=1e14, y=0.2, label="Archaea", color="black", size=6)
+ scale_colour_manual(labels = lab1, values = c('blue', 'red','orange',
'black', '#8918FA', '1CF339', '#1CD7F3'))

ggsave(file="Archaea density vs depth.png", width=6.5,height=6.5)

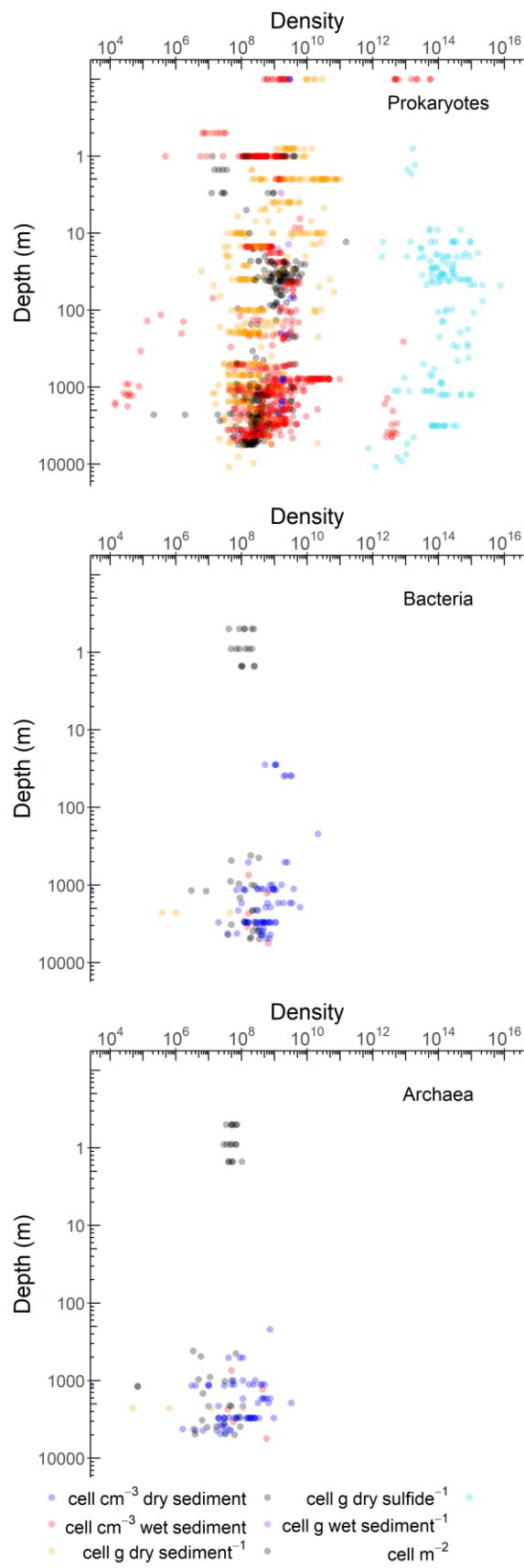
## Warning: Transformation introduced infinite values in continuous x-axis

ggpubr::ggarrange(Fig6, Fig7, Fig8, nrow=3, common.legend = TRUE, legend =
"bottom")

## Warning: Removed 33 rows containing missing values (geom_point).
## Removed 33 rows containing missing values (geom_point).

## Warning: Transformation introduced infinite values in continuous x-axis
## Transformation introduced infinite values in continuous x-axis

```



```
setwd("C:/Users/Mama/Desktop/manuscript revised/Figures")
ggsave(file="microbial density vs depth.png", width=6.5, height=19.5)
```