

all_strain_genome_stats

2023-07-12

Assembly stats for 13 *Emiliania huxleyi* genomes

```
genstats <- list()
gc <- list()
stats <- list()
strains <- as.character(read.csv('data/strains.csv',header=FALSE))
for (strain in strains){
  genstats[[strain]] <- read.table(paste0("data/2023-genome-stats/stats_after_pilon_round_2_decontam/",
                                           strain,".genstats.txt"),sep="\t",header=F)
  colnames(genstats[[strain]]) <- c('Contig_name','Avg_fold', 'Length', 'Ref_GC',
                                      'Covered_percent', 'Covered_bases','Plus_reads',
                                      'Minus_reads', 'Read_GC', 'Median_fold', 'Std_Dev')
  
  gc[[strain]] <- read.table(paste0("data/2023-genome-stats/stats_after_pilon_round_2_decontam/",
                                    strain,".gcscaffold.txt"),sep="\t",header=F)
  colnames(gc[[strain]]) <- c('Contig_name', 'Length', 'A', 'C', 'G', 'T', 'N',
                               'IUPAC','Other', 'GC')
  stats[[strain]] <- inner_join(genstats[[strain]],gc[[strain]],by="Contig_name")
}
```

Read in assembly stats data

```
n50s <- c()
lengths <- c()
contig_counts <- c()
mins <- c()
maxs <- c()
for (strain in strains){
  contig_lengths <- stats[[strain]]$Length.x
  contig_counts <- c(contig_counts,length(contig_lengths))
  total_assembled_length=sum(contig_lengths)
  contig_lengths <- sort(contig_lengths,decreasing=TRUE)

  sum <- 0
  for (length in contig_lengths){
    sum <- sum+length
    if (sum>=total_assembled_length/2){
      n50s <- c(n50s,as.numeric(length))
      lengths <- c(lengths,total_assembled_length)
    }
  }
}
```

```

    maxs <- c(maxs,contig_lengths[1])
    mins <- c(mins,contig_lengths[length(contig_lengths)])
    break
}
}

}

global_stats <- data.frame(n50s,lengths,contig_counts,mins,maxs)
rownames(global_stats) <- strains
colnames(global_stats) <- c('N50','Total assembled length','Contig count',
                            'Min contig length','Max contig length')

```

Calculate globals assembly stats including N50 and total assembled length

```

name_translation <- read.table('data/genomescope/illumina-run-conversions.txt',sep=' ')
temp <- name_translation$V2
names(temp) <- name_translation$V1
name_translation <- temp

```

```

global_stats$genome_haploid_length <- seq(1,nrow(global_stats))
global_stats$genome_unique_length <- seq(1,nrow(global_stats))
for (folder in list.dirs('data/genomescope/')){
  if (grepl('HA',folder)){
    key <- str_split(folder,'_',simplify=TRUE)[,1]
    key <- str_split(key,'/',simplify=TRUE)[,4]
    if (sum(grepl(name_translation[key],rownames(global_stats)))==1){
      temp <- read.csv(paste0(folder,"/summary.txt_fixed.csv"))
      global_stats[name_translation[key],'genome_haploid_length'] <-
        mean(temp[2,'min'],temp[2,'max'])
      global_stats[name_translation[key],'genome_unique_length'] <-
        mean(temp[4,'min'],temp[4,'max'])
    }
  }
}
#print(temp)
print(global_stats)

```

Add in estimated genome size stats, calculated using Genomescope

	N50	Total assembled length	Contig count	Min contig length
## CCMP371	167293	299709025	5373	671
## CCMP375	82247	247806427	8312	505
## CCMP377	70514	297357770	9460	493
## CCMP1280	104319	241936265	7168	510
## RCC874	3015506	152968590	396	522
## RCC914	63139	210245484	9395	508
## RCC1222	55476	179814701	10077	160
## RCC1239	364457	171886557	2195	506

	Max contig length	genome_haploid_length	genome_unique_length	
## RCC1256	43379	236745271	10971	393
## RCC3492	70125	211559349	9734	496
## RCC3963	146371	281844499	5463	544
## RCC6071	511805	200137934	1929	545
## RCC6856	78893	333916182	9838	528
## CCMP371	1476961	98806191	72440020	
## CCMP375	1947962	103218139	69423426	
## CCMP377	606376	138388017	76045285	
## CCMP1280	4348365	111401730	64523269	
## RCC874	7853977	118376237	91959391	
## RCC914	653764	86242315	53701350	
## RCC1222	1451258	107508536	66175986	
## RCC1239	1376033	116873978	83660661	
## RCC1256	579311	93127971	69271484	
## RCC3492	1245722	77328042	62355972	
## RCC3963	1638904	107212419	77275117	
## RCC6071	2885505	131960871	81986586	
## RCC6856	509900	123410186	76317769	

```
global_stats$Strain <- rownames(global_stats)
ggplot(global_stats,aes(x=N50,y=genome_haploid_length,label=Strain))+ 
  geom_point()+
  theme_bw()+
  scale_x_log10()+
  geom_smooth(method=lm,colour="black")+
  ylab("Haploid genome length")+
  xlab("Contig N50")+
  labs(caption=str_wrap("Figure 1. Relationship between predicted haploid genome
                        length and contig N50 for 13 Emiliania huxleyi
                        genomes.",75))+
  geom_text_repel(size=3)+
  theme(plot.caption = element_text(hjust = 0,size=12))
```

Genomescope predicted haploid genome length vs N50

```
## `geom_smooth()` using formula = 'y ~ x'

## Warning: The following aesthetics were dropped during statistical transformation: label
## i This can happen when ggplot fails to infer the correct grouping structure in
##   the data.
## i Did you forget to specify a 'group' aesthetic or to convert a numerical
##   variable into a factor?
```

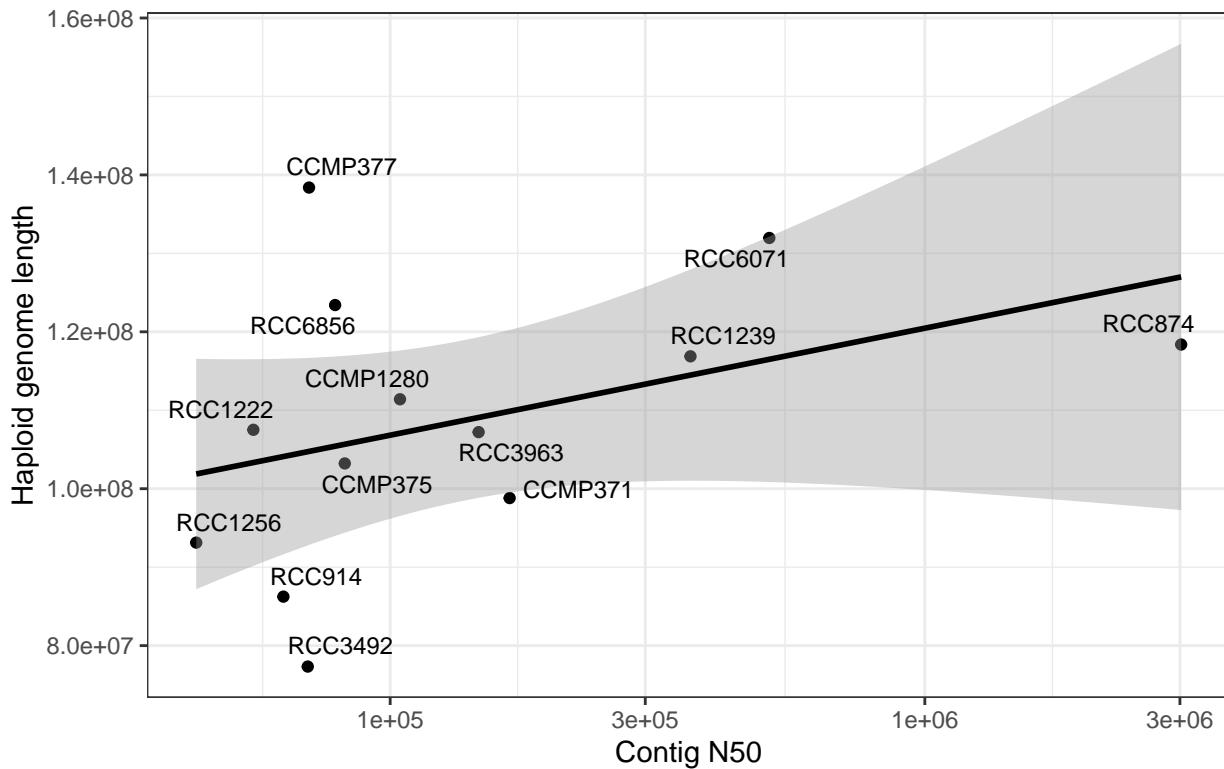


Figure 1. Relationship between predicted haploid genome length and contig N50 for 13 *Emiliania huxleyi* genomes.

```
ggsave("plots/haploid_genome_length_vs_N50.png")
```

```
## Saving 6.5 x 4.5 in image
## `geom_smooth()` using formula = 'y ~ x'

## Warning: The following aesthetics were dropped during statistical transformation: label
## i This can happen when ggplot fails to infer the correct grouping structure in
##   the data.
## i Did you forget to specify a 'group' aesthetic or to convert a numerical
##   variable into a factor?
```

```
global_stats$Strain <- rownames(global_stats)
ggplot(global_stats,aes(x=N50,y=genome_unique_length,label=Strain))+ 
  geom_point()+
  theme_bw()+
  scale_x_log10()+
  geom_smooth(method=lm,colour="black")+
  ylab("Unique genome length")+
  xlab("Contig N50")+
  labs(caption=str_wrap("Figure 2. Relationship between predicted unique genome
length and contig N50 for 13 Emiliania huxleyi
genomes.",75))+
```

```
geom_text_repel(size=3)+  
theme(plot.caption = element_text(hjust = 0, size=12))
```

Genomescope predicted unique genome length vs N50

```
## `geom_smooth()` using formula = 'y ~ x'  
  
## Warning: The following aesthetics were dropped during statistical transformation: label  
## i This can happen when ggplot fails to infer the correct grouping structure in  
##   the data.  
## i Did you forget to specify a 'group' aesthetic or to convert a numerical  
##   variable into a factor?
```

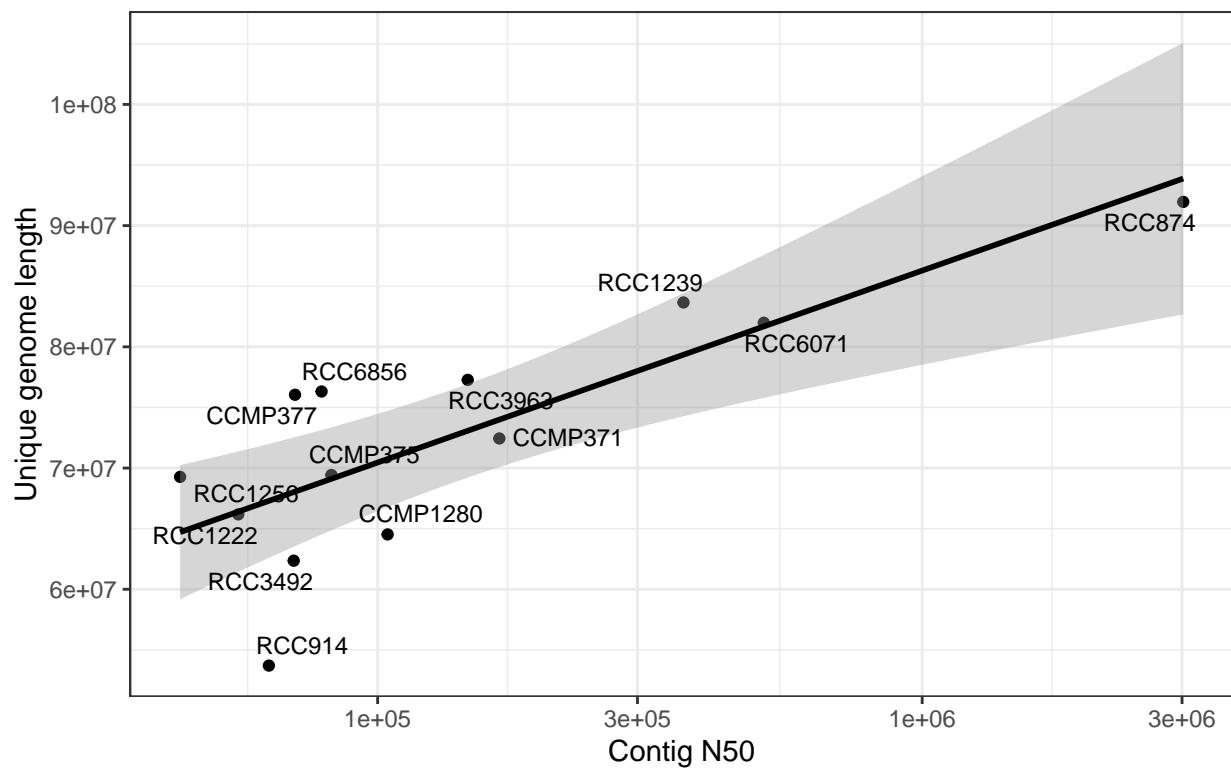


Figure 2. Relationship between predicted unique genome length and contig N50 for 13 *Emiliania huxleyi* genomes.

```
ggsave("plots/unique_genome_length_vs_N50.png")
```

```
## Saving 6.5 x 4.5 in image  
## `geom_smooth()` using formula = 'y ~ x'  
  
## Warning: The following aesthetics were dropped during statistical transformation: label  
## i This can happen when ggplot fails to infer the correct grouping structure in  
##   the data.  
## i Did you forget to specify a 'group' aesthetic or to convert a numerical  
##   variable into a factor?
```

```

stats_joined <- ldply(stats, rbind)
ggplot(stats_joined, aes(x=Length.x)) + geom_histogram() + scale_x_log10() +
  theme_bw() +
  xlab("Contig size") +
  ylab("Count") + facet_wrap(vars(.id)) +
  labs(caption = str_wrap("Figure 3. Contig length distribution for 13 Emiliania huxleyi genomes.", 75)) +
  theme(plot.caption = element_text(hjust = 0, size=12))

```

Contig length distribution for each strain

‘stat_bin()’ using ‘bins = 30’. Pick better value with ‘binwidth’.

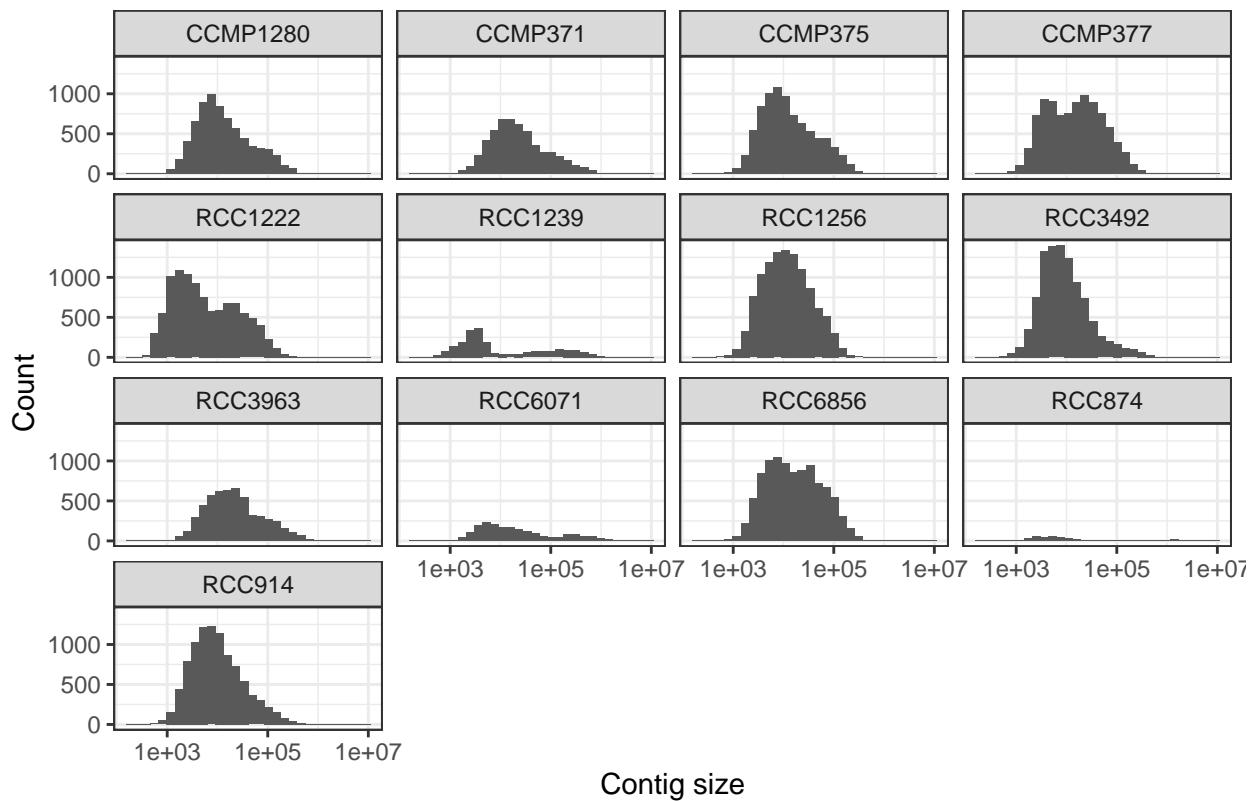


Figure 3. Contig length distribution for 13 *Emiliania huxleyi* genomes.

```
ggsave("plots/contig_length_distributions.png")
```

Saving 6.5 x 4.5 in image
‘stat_bin()’ using ‘bins = 30’. Pick better value with ‘binwidth’.

```

ggplot(data = stats_joined, aes(x = GC, y=Avg_fold)) +
  geom_point(size=0.5) + ylim(0, 300) + facet_wrap(vars(.id)) +

```

```

    theme_bw()+
  ylab("Contig GC%")+
  xlab("Average Fold Coverage")+
  labs(caption=str_wrap("Figure 4. Contig Fold coverage vs GC percentage for 13  
Emiliana huxleyi genomes.",75))+  
  theme(plot.caption = element_text(hjust = 0,size=12))

```

Scatterplots of fold coverage vs GC percentage (each point represents a contig)

Warning: Removed 150 rows containing missing values ('geom_point()').

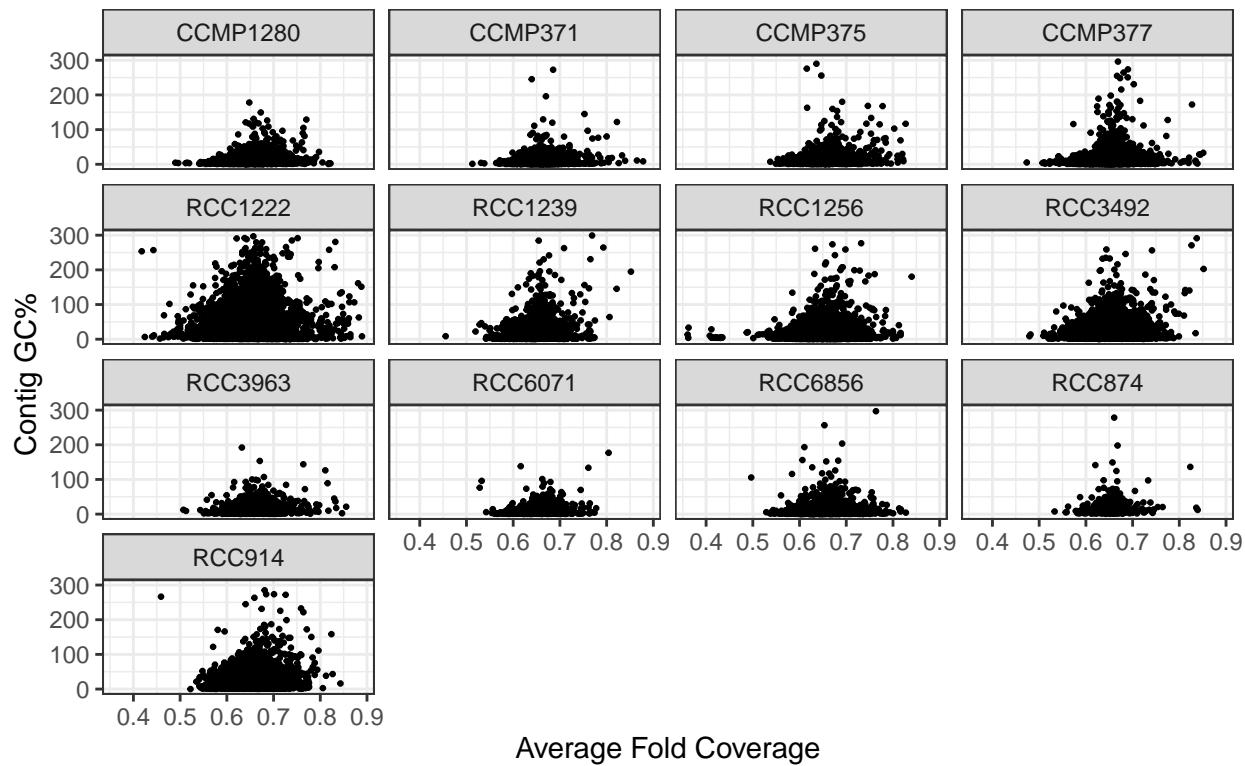


Figure 4. Contig Fold coverage vs GC percentage for 13 Emiliana huxleyi genomes.

```
ggsave("plots/coverage_vs_GC.png")
```

Saving 6.5 x 4.5 in image

Warning: Removed 150 rows containing missing values ('geom_point()').

```

global_stats$total_assembled_length <- global_stats[, "Total assembled length"]
ggplot(global_stats,aes(x=total_assembled_length,y=genome_unique_length,label=Strain))+  
  geom_point()+
  theme_bw()

```

```

geom_smooth(method=lm, colour="black")+
ylab("Unique genome length")+
xlab("Total assembled length")+
labs(caption=str_wrap("Figure 5. Relationship between predicted unique genome length and total assembled length for 13 Emiliania huxleyi genomes.", 75))+
  geom_text_repel(size=3)+
theme(plot.caption = element_text(hjust = 0, size=12))

```

Genomescope predicted unique genome length vs assembled length

```

## `geom_smooth()` using formula = 'y ~ x'

## Warning: The following aesthetics were dropped during statistical transformation: label
## i This can happen when ggplot fails to infer the correct grouping structure in
##   the data.
## i Did you forget to specify a 'group' aesthetic or to convert a numerical
##   variable into a factor?

```

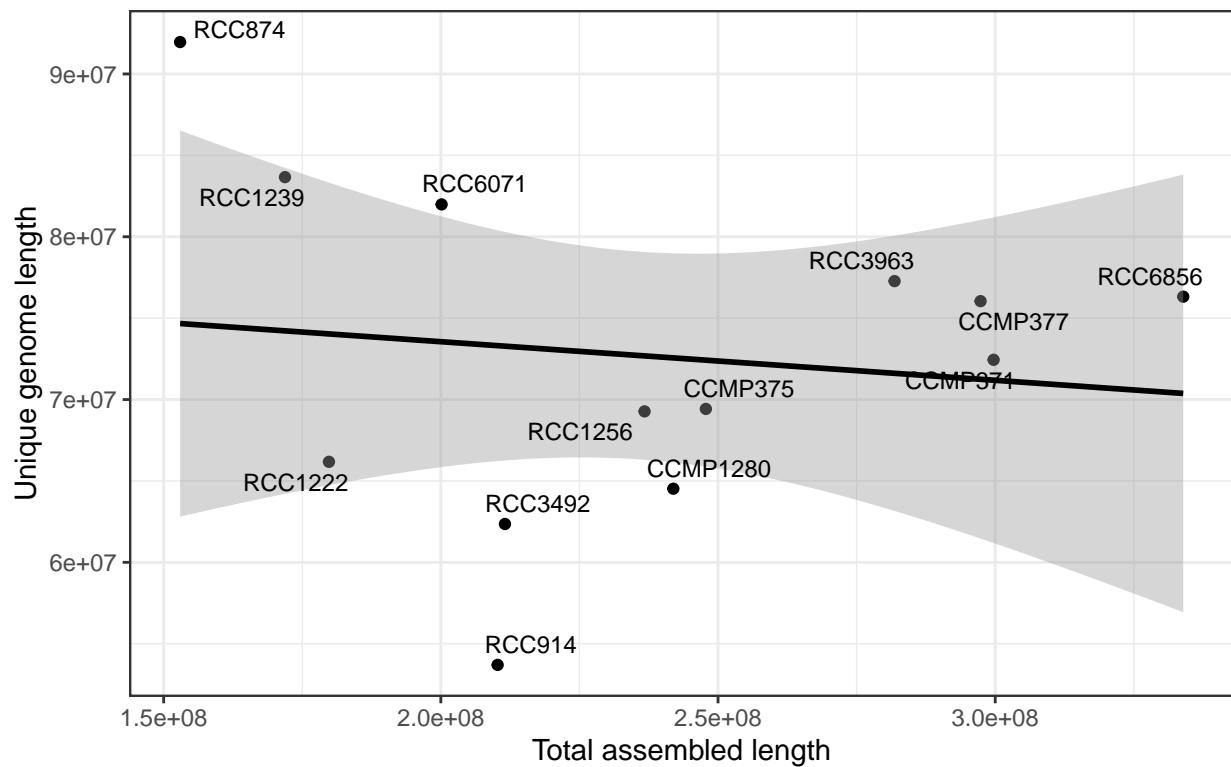


Figure 5. Relationship between predicted unique genome length and total assembled length for 13 *Emiliania huxleyi* genomes.

```
ggsave("plots/unique_genome_length_vs_total_assembled_length.png")
```

```

## Saving 6.5 x 4.5 in image
## `geom_smooth()` using formula = 'y ~ x'

```

```

## Warning: The following aesthetics were dropped during statistical transformation: label
## i This can happen when ggplot fails to infer the correct grouping structure in
##   the data.
## i Did you forget to specify a 'group' aesthetic or to convert a numerical
##   variable into a factor?

ggplot(global_stats,aes(x=total_assembled_length,y=genome_haploid_length,label=Strain))+  

  geom_point()+
  theme_bw()+
  geom_smooth(method=lm,colour="black")+
  ylab("Haploid genome length")+
  xlab("Total assembled length")+
  labs(caption=str_wrap("Figure 6. Relationship between predicted haploid genome  
length and total assembled length for 13 Emiliania huxleyi  
genomes.",75))+
  geom_text_repel(size=3)+
  theme(plot.caption = element_text(hjust = 0,size=12))

## `geom_smooth()` using formula = 'y ~ x'

## Warning: The following aesthetics were dropped during statistical transformation: label
## i This can happen when ggplot fails to infer the correct grouping structure in
##   the data.
## i Did you forget to specify a 'group' aesthetic or to convert a numerical
##   variable into a factor?

```

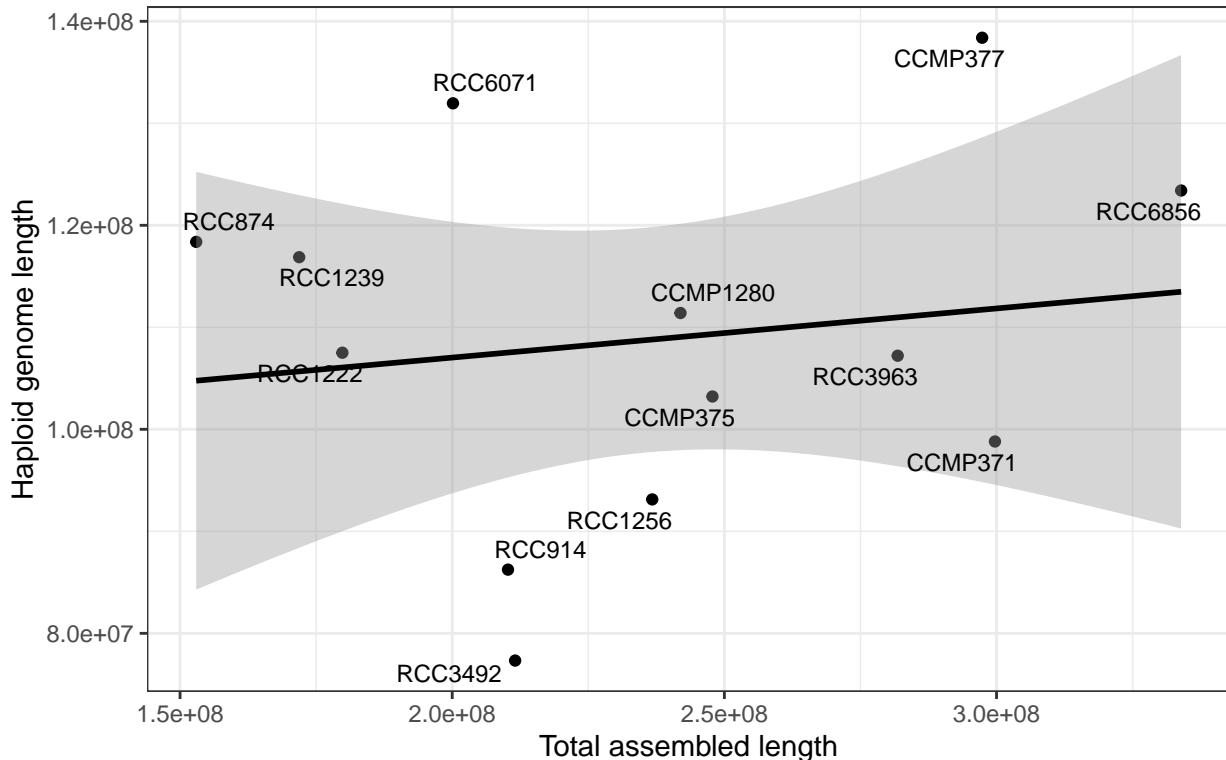


Figure 6. Relationship between predicted haploid genome length and total assembled length for 13 *Emiliania huxleyi* genomes.

```
ggsave("plots/haploid_genome_length_vs_total_assembled_length.png")

## Saving 6.5 x 4.5 in image
## `geom_smooth()` using formula = 'y ~ x'

## Warning: The following aesthetics were dropped during statistical transformation: label
## i This can happen when ggplot fails to infer the correct grouping structure in
##   the data.
## i Did you forget to specify a 'group' aesthetic or to convert a numerical
##   variable into a factor?
```