## VC\_Bench\_Poster\_Figures

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### A BUNCH OF PREPROCESSING

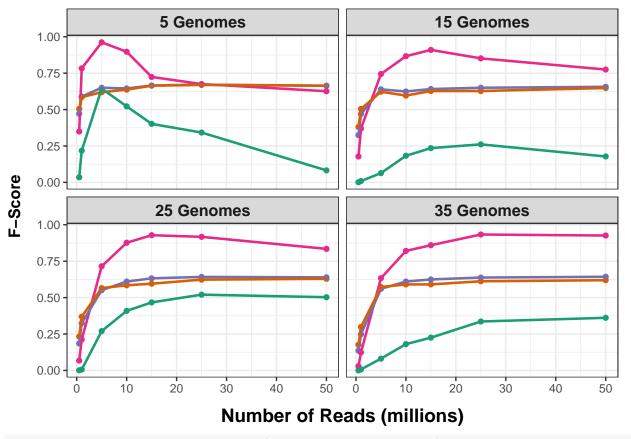
```
library(tidyverse)
## -- Attaching packages ----
                                                     ----- tidyverse 1.3.0 --
                    v purrr
## v ggplot2 3.3.3
                               0.3.4
## v tibble 3.1.0 v dplyr
                              1.0.5
## v tidyr
           1.1.3
                    v stringr 1.4.0
                     v forcats 0.5.1
## v readr
            1.4.0
## -- Conflicts ----- tidyverse_conflicts() --
## x dplyr::filter() masks stats::filter()
## x dplyr::lag()
                    masks stats::lag()
library(RColorBrewer)
library(reshape2)
##
## Attaching package: 'reshape2'
## The following object is masked from 'package:tidyr':
##
##
      smiths
library(scales)
##
## Attaching package: 'scales'
## The following object is masked from 'package:purrr':
##
##
## The following object is masked from 'package:readr':
##
bench_df<- read.csv("~/Desktop/NoyesLab/All_Benchmarks.csv")</pre>
\#bench\_df
###Dataset Fixing
bench_df$Dataset_num<-ifelse(bench_df$Dataset=="M0.5",0.5,
                        ifelse(bench_df$Dataset=="M1",1,
                               ifelse(bench_df$Dataset=="M5",5,
```

```
ifelse(bench_df$Dataset=="M10",10,
                                              ifelse(bench_df$Dataset=="M15",15,
                                                     ifelse(bench_df$Dataset=="M25",25,50))))))
bench_df$Dataset_f<-factor(bench_df$Dataset, levels=c("M0.5","M1","M5","M10","M15","M25","M50"))
###Subset Fixing
bench_df$Subset_num<-ifelse(bench_df$Subset=="S5",5,
                            ifelse(bench df$Subset=="S15",15,
                                   ifelse(bench df$Subset=="S25",25,35)))
bench_df$Subset<-ifelse(bench_df$Subset=="S5","5 Genomes",
                        ifelse(bench_df$Subset=="S15","15 Genomes",
                               ifelse(bench_df$Subset=="S25","25 Genomes","35 Genomes")))
bench_df$Subset_f<-factor(bench_df$Subset, levels=c("5 Genomes","15 Genomes","25 Genomes","35 Genomes")
###Variant Caller Fixing
bench_df$VCaller<-ifelse(bench_df$VCaller=="FB_Out", "FreeBayes",
                         ifelse(bench_df$VCaller=="GATK_Out", "GATK",
                                ifelse(bench_df$VCaller=="Disco_Out","DiscoSNP++","MetaSNV")))
bench_df$VCaller_f<-factor(bench_df$VCaller,levels=c("MetaSNV","GATK","FreeBayes","DiscoSNP++"))
cbbPalette<-rev(brewer.pal(n=4,name = "Dark2"))
```

#### Making color palettes

```
cbbPalette<-rev(brewer.pal(n=4,name = "Dark2"))
VCPalette<-rev(brewer.pal(n=7,name = "Dark2"))
VC_light_palette<-rev(brewer.pal(n=7,name = "BuGn"))</pre>
```

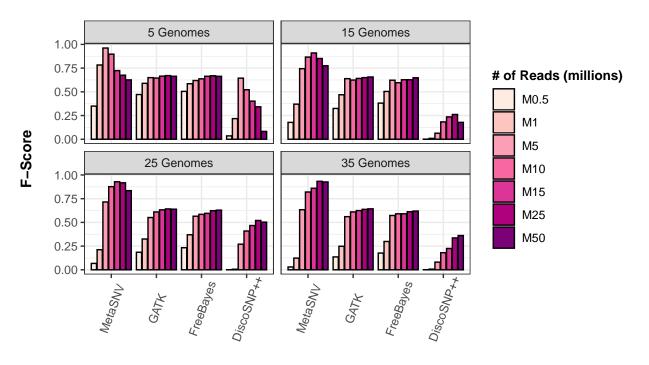
### SUBSET AS A FACET



#legend.title = element\_text(size = 10, face = "bold"),legend.position = "bottom")
###MUST INCLUDE

#### 1 A

```
###Reasonably good
#1B
ggplot(bench_df,aes(x=VCaller_f,y=F.score,fill=Dataset_f,group=Dataset_f)) +
    geom_col(colour="black",position="dodge2") + facet_wrap(~Subset_f,nrow=2) +
    ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
    xlab("Variant Caller") + ylab("F-Score") + labs(fill="# of Reads (millions)") +
    scale_fill_brewer(palette = "RdPu") + theme_bw() +
    theme(plot.title = element_text(hjust=0.5,size=13,face="bold",margin=margin(0,0,15,0)),
        axis.title.x = element_text(face="bold",margin=margin(15,0,0,0)),
        axis.title.y = element_text(face="bold",margin=margin(0,15,0,0)),
        legend.title = element_text(size = 10, face = "bold"))
```

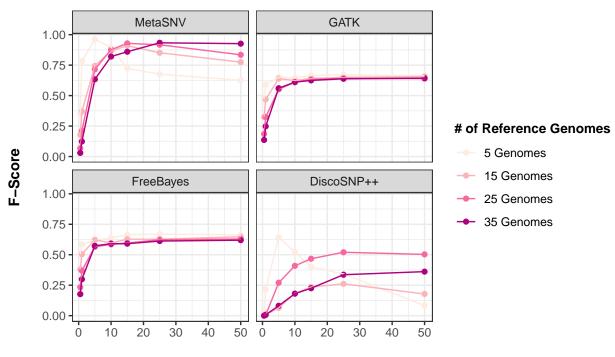


**Variant Caller** 

1B

## VARIANT CALLER AS A FACET

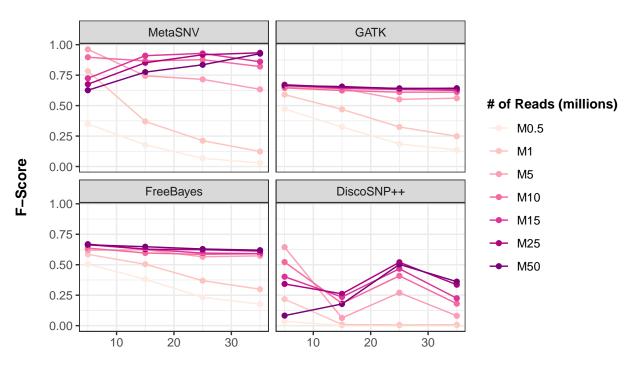
```
#####CONSIDER USING EITHER LINE GRAPHS
###This one is BETTER
#2A
ggplot(bench_df,aes(x=Dataset_num,y=F.score,colour=Subset_f,group=Subset_f)) +
    geom_point() + geom_line() + facet_wrap(~VCaller_f,nrow=2) +
    ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
    xlab("Number of Reads (millions)") + ylab("F-Score") + labs(col="# of Reference Genomes") +
    scale_color_brewer(palette = "RdPu") + theme_bw() +
    theme(plot.title = element_text(hjust=0.5,size=13,face="bold",margin=margin(0,0,15,0)),
        axis.title.x = element_text(face="bold",margin=margin(0,15,0,0)),
        axis.title.y = element_text(face="bold",margin=margin(0,15,0,0)),
        legend.title = element_text(size = 10, face = "bold"))
```



## **Number of Reads (millions)**

### 2A

```
###This is also good
#2B
ggplot(bench_df,aes(x=Subset_num,y=F.score,colour=Dataset_f,group=Dataset_f)) +
    geom_point() + geom_line() + facet_wrap(~VCaller_f,nrow=2) +
    ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
    xlab("Number of Reference Genomes") + ylab("F-Score") + labs(col="# of Reads (millions)") +
    scale_color_brewer(palette = "RdPu") + theme_bw() +
    theme(plot.title = element_text(hjust=0.5,size=13,face="bold",margin=margin(0,0,15,0)),
        axis.title.x = element_text(face="bold",margin=margin(15,0,0,0)),
        axis.title.y = element_text(face="bold",margin=margin(0,15,0,0)),
        legend.title = element_text(size = 10, face = "bold"))
```

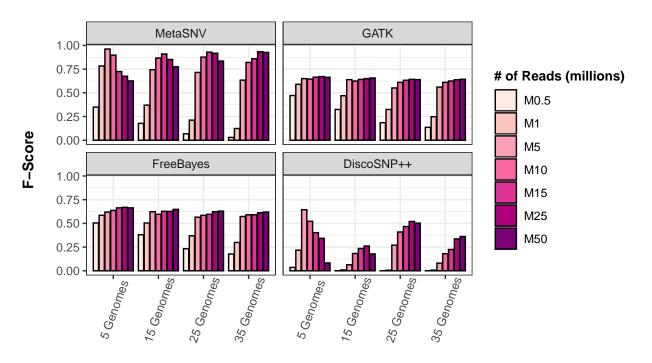


**Number of Reference Genomes** 

### 2B

```
#Barplot version is less good but viable
#2C

ggplot(bench_df,aes(x=Subset_f,y=F.score,fill=Dataset_f,group=Dataset_f)) +
    geom_col(colour="black",position="dodge2") + facet_wrap(~VCaller_f,nrow=2) +
    ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
    xlab("Number of Reference Genomes") + ylab("F-Score") + labs(fill="# of Reads (millions)") +
    scale_fill_brewer(palette = "RdPu") + theme_bw() +
    theme(plot.title = element_text(hjust=0.5,size=13,face="bold",margin=margin(0,0,15,0)),
        axis.title.x = element_text(face="bold",margin=margin(15,0,0,0)),
        axis.title.y = element_text(size=9,angle=70,vjust = 0.6),
        axis.title.y = element_text(face="bold",margin=margin(0,15,0,0)),
        legend.title = element_text(size = 10, face = "bold"))
```

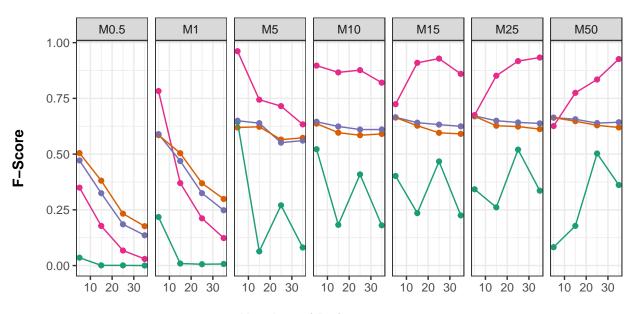


**Number of Reference Genomes** 

2C

## DATASET AS A FACET

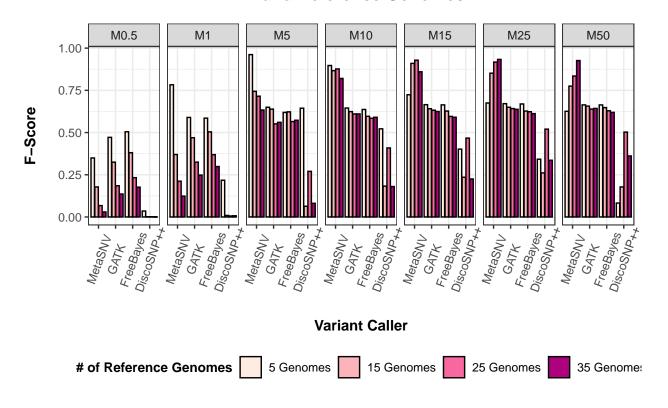
```
###INCLUDE THIS ONE IF POSSIBLE -- GOOD STORY HERE
#3A
ggplot(bench_df,aes(x=Subset_num,y=F.score,colour=VCaller_f,group=VCaller_f)) +
    geom_point() + geom_line() + facet_wrap(~Dataset_f,nrow=1) +
    ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
    xlab("Number of Reference Genomes") + ylab("F-Score") + labs(col="Variant Caller") +
    scale_color_manual(values=cbbPalette) + theme_bw() +
    theme(plot.title = element_text(hjust=0.5,size=13,face="bold",margin=margin(0,0,15,0)),
        axis.title.x = element_text(face="bold",margin=margin(0,15,0,0)),
        axis.title.y = element_text(face="bold",margin=margin(0,15,0,0)),
        legend.title = element_text(size = 10, face = "bold"), legend.position = "bottom")
```



### **Number of Reference Genomes**

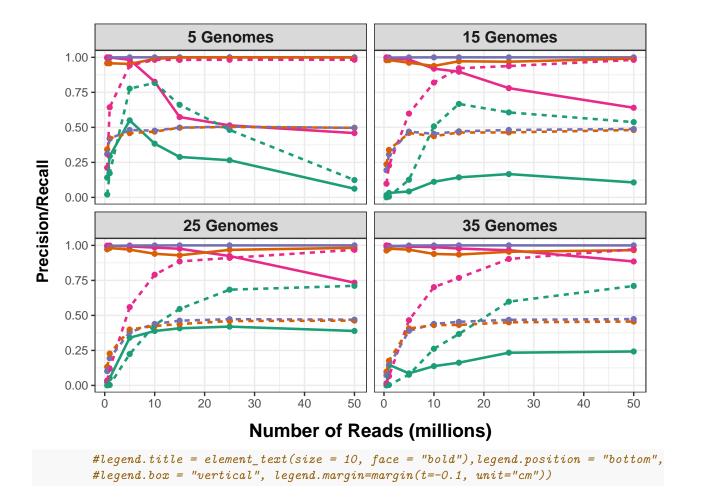
```
Variant Caller → MetaSNV → GATK → FreeBayes → DiscoSNP++
```

### **3A**

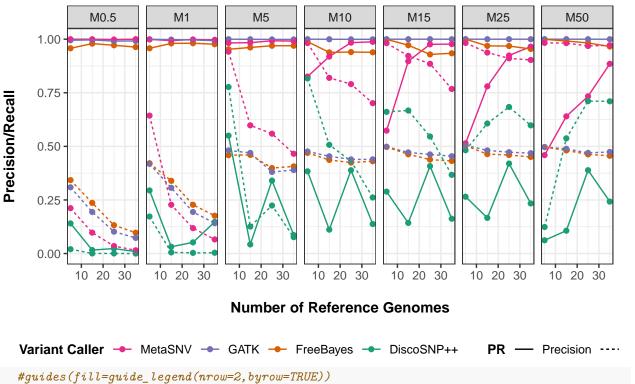


3B

## Figures using Precision/Recall

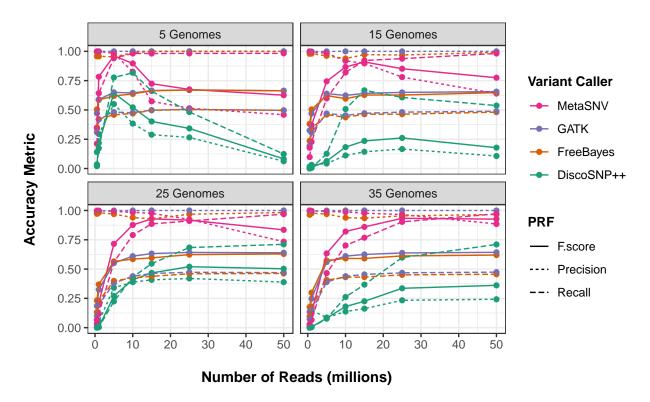


### 1PR



#guiues() itt-guiue\_tegenu(niiow=2,0giow=11tt

## 3PR



1PRF

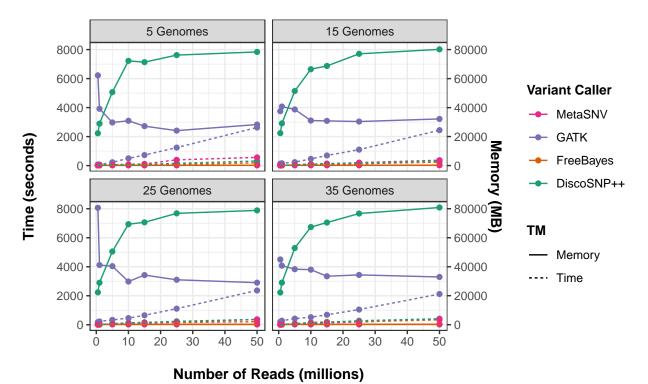
## Time and Memory Benchmarks

These ones aren't that good

```
tm_df<-gather(bench_df,"TM","TM_value",10:11)

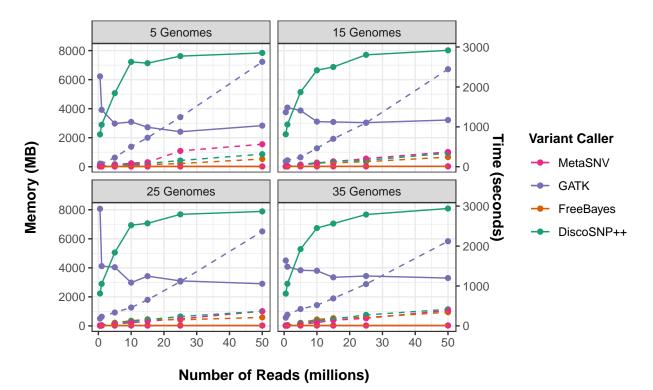
#Scaled axis, unscaled data
ggplot(tm_df,aes(x=Dataset_num,y=TM_value,colour=VCaller_f,group=interaction(VCaller_f,TM))) +
    geom_point() + geom_line(aes(linetype=TM)) + facet_wrap(~Subset_f,nrow=2) +
    ggtitle("Time & Memory Benchmarks by Number of Reads\n and Reference Genomes") +
    xlab("Number of Reads (millions)") + ylab("Time (seconds)") + labs(col="Variant Caller") +
    scale_y_continuous(name="Time (seconds)", sec.axis = sec_axis(~ 10*., name="Memory (MB)")) +
    scale_color_manual(values=cbbPalette) + theme_bw() +
    theme(plot.title = element_text(hjust=0.5,size=13,face="bold",margin=margin(0,0,15,0)),
        axis.title.x = element_text(face="bold",margin=margin(0,15,0,0)),
        axis.title.y = element_text(face="bold",margin=margin(0,15,0,0)),
        legend.title = element_text(size = 10, face = "bold")</pre>
```

## Time & Memory Benchmarks by Number of Reads and Reference Genomes



### TM1

# Time & Memory Benchmarks by Number of Reads and Reference Genomes

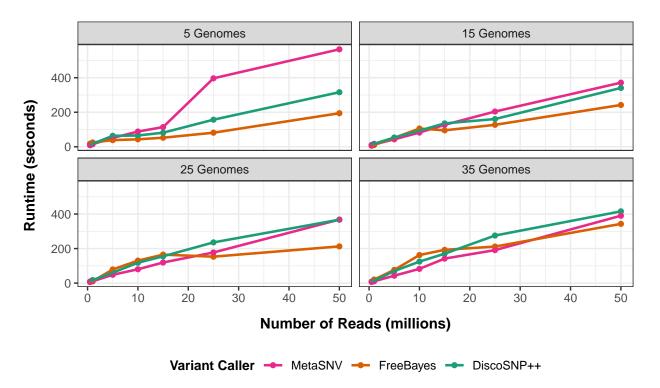


TM2

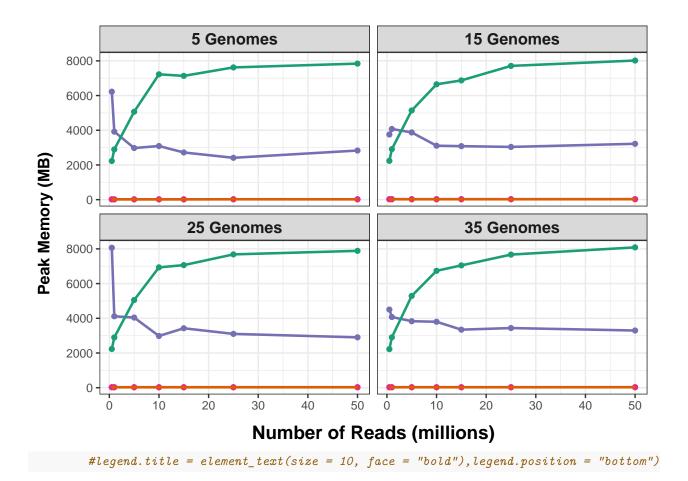
\

## These are Fab

# Runtime by Number of Reads and Reference Genomes



### T1



### M1

## Genome Heatmap

```
five<-read.csv("~/Desktop/NoyesLab/SNPSubs/SNPLog_subset5.csv")
fifteen<-read.csv("~/Desktop/NoyesLab/SNPSubs/SNPLog_subset15.csv")
twentyfive<-read.csv("~/Desktop/NoyesLab/SNPSubs/SNPLog_subset25.csv")
thirtyfive<-read.csv("~/Desktop/NoyesLab/SNPSubs/SNPLog_subset25.csv")

#Heat_df<-data.frame(Five=unique(five$CHROM),Fifteen=unique(fifteen$CHROM),TwentyFive=unique(fifteen$CHROM)

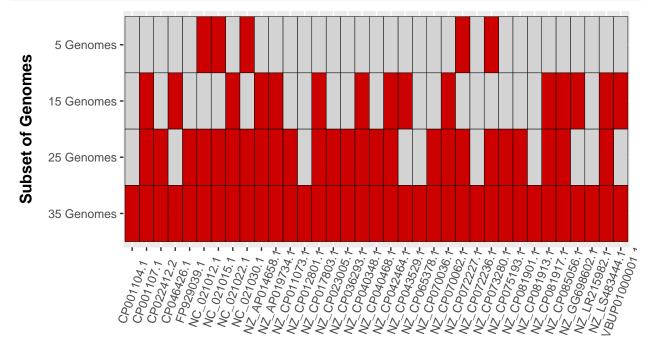
Heat_df<-data.frame(Full=unique(thirtyfive$CHROM))

Heat_df$"5 Genomes"<-ifelse(Heat_df$Full %in% unique(five$CHROM),1,0)
Heat_df$"15 Genomes"<-ifelse(Heat_df$Full %in% unique(fifteen$CHROM),1,0)
Heat_df$"25 Genomes"<-ifelse(Heat_df$Full %in% unique(twentyfive$CHROM),1,0)
Heat_df$"35 Genomes"<-ifelse(Heat_df$Full %in% unique(thirtyfive$CHROM),1,0)

Heat_gather<-gather(Heat_df, "Genome_num", "Genome_ext",2:5)

Heat_gather$Genome_num_f<-factor(Heat_gather$Genome_num,levels=c("35 Genomes","25 Genomes","15 Genomes"
```

```
ggplot(Heat_gather,aes(x=Full,y=Genome_num_f)) + geom_tile(col="black",aes(fill=factor(Genome_ext))) +
    scale_fill_manual(labels=c("Excluded", "Included"),values=c("0"="light grey","1"="red3")) + labs(fill=
    xlab("Reference Genomes") + ylab("Subset of Genomes") +
    theme(plot.title = element_text(hjust=0.5,size=13,face="bold",margin=margin(0,0,15,0)),
        axis.title.x = element_text(size=12,face="bold",margin=margin(10,0,0,0)),
        axis.text.x = element_text(size=9,angle=70,vjust = 0.6),
        axis.title.y = element_text(size=12,face="bold",margin=margin(0,10,0,0)),
        legend.title = element_text(size = 12, face = "bold"), legend.text = element_text(size=12), leg
```



#### **Reference Genomes**

