

VC_Bench_Poster_Figures

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A BUNCH OF PREPROCESSING

```
library(tidyverse)

## -- 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 tidyr 1.1.3        v stringr 1.4.0
## v readr 1.4.0        v forcats 0.5.1

## -- 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

bench_df<- read.csv("~/Desktop/NoyesLab/Benchmark.csv")
#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"))

```

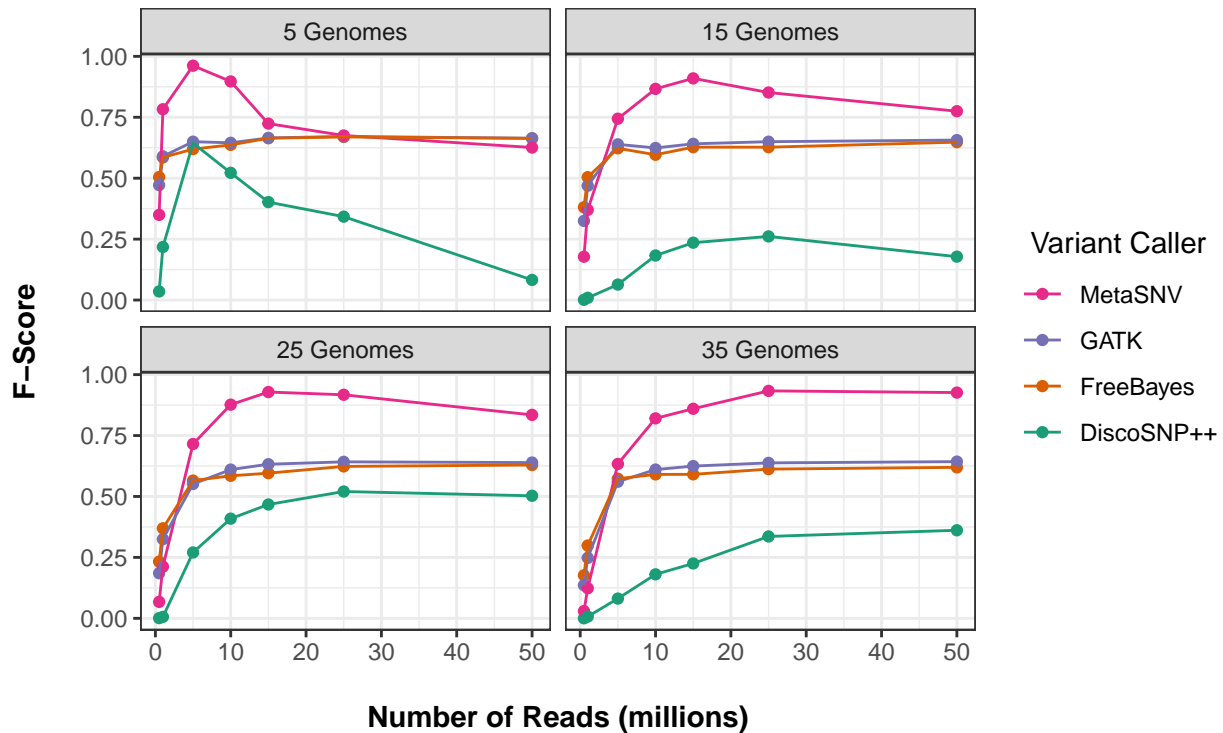
SUBSET AS A FACET

```

\\
###MUST INCLUDE
#1A
ggplot(bench_df, aes(x=Dataset_num, y=F.score, colour=VCaller_f, group=VCaller_f)) +
  geom_point() + geom_line() + facet_wrap(~Subset_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="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(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") +
        annotate("text", x=30, y = 0.5, label="HELLO WORLD", vjust=1, hjust=1))

```

Variant Caller Accuracy by Number of Reads and Reference Genomes

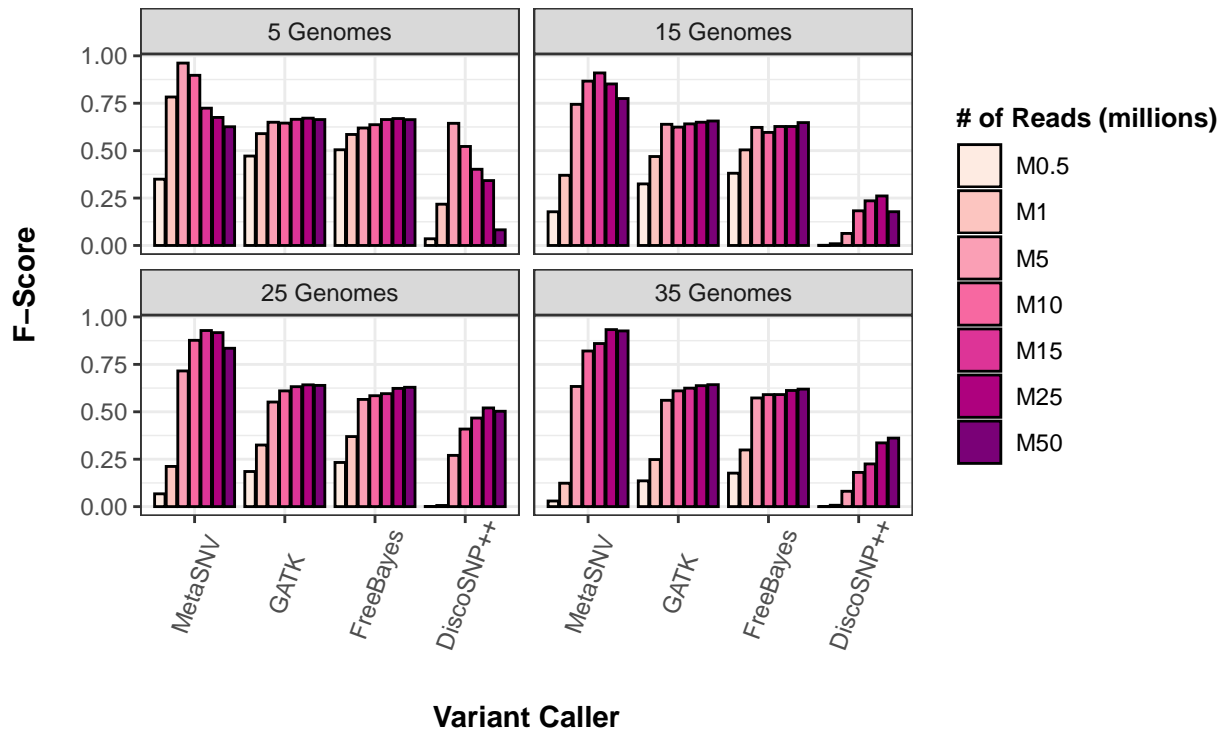


###MUST INCLUDE

1A

```
###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.text.x = 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"))
```

Variant Caller Accuracy by Number of Reads and Reference Genomes



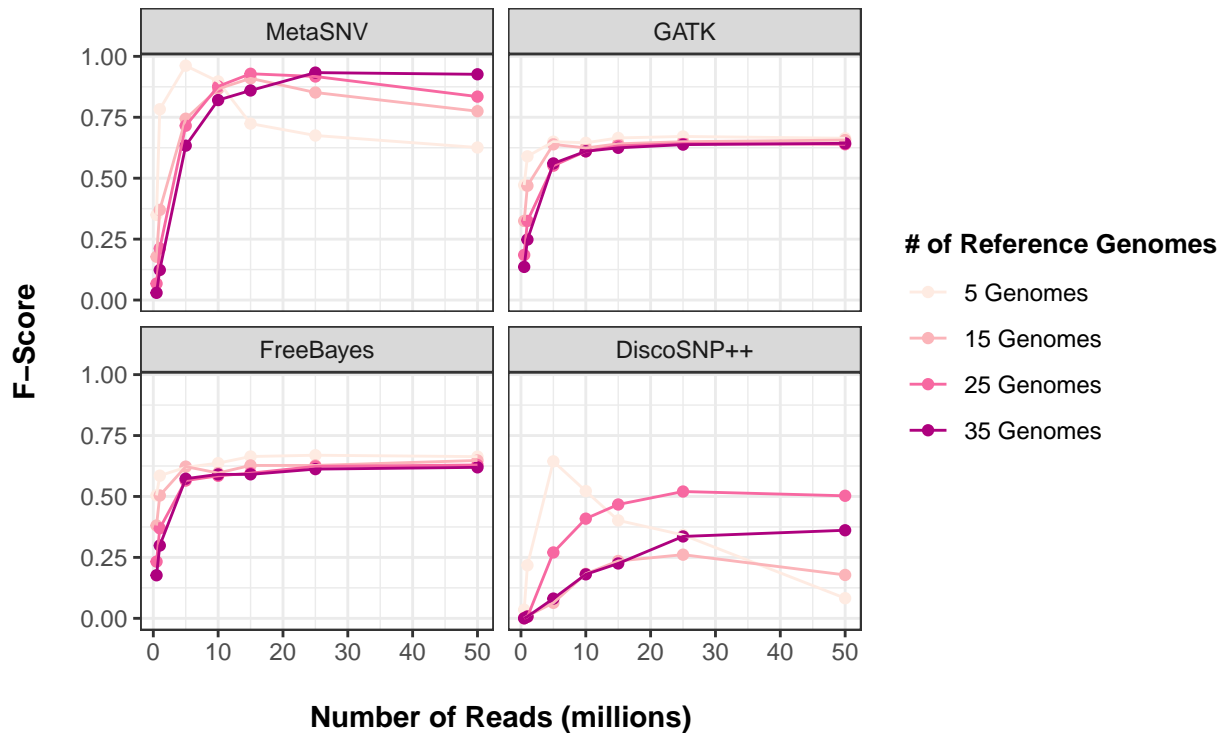
1B

VARIANT CALLER AS A FACET

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```
#####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(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"))
```

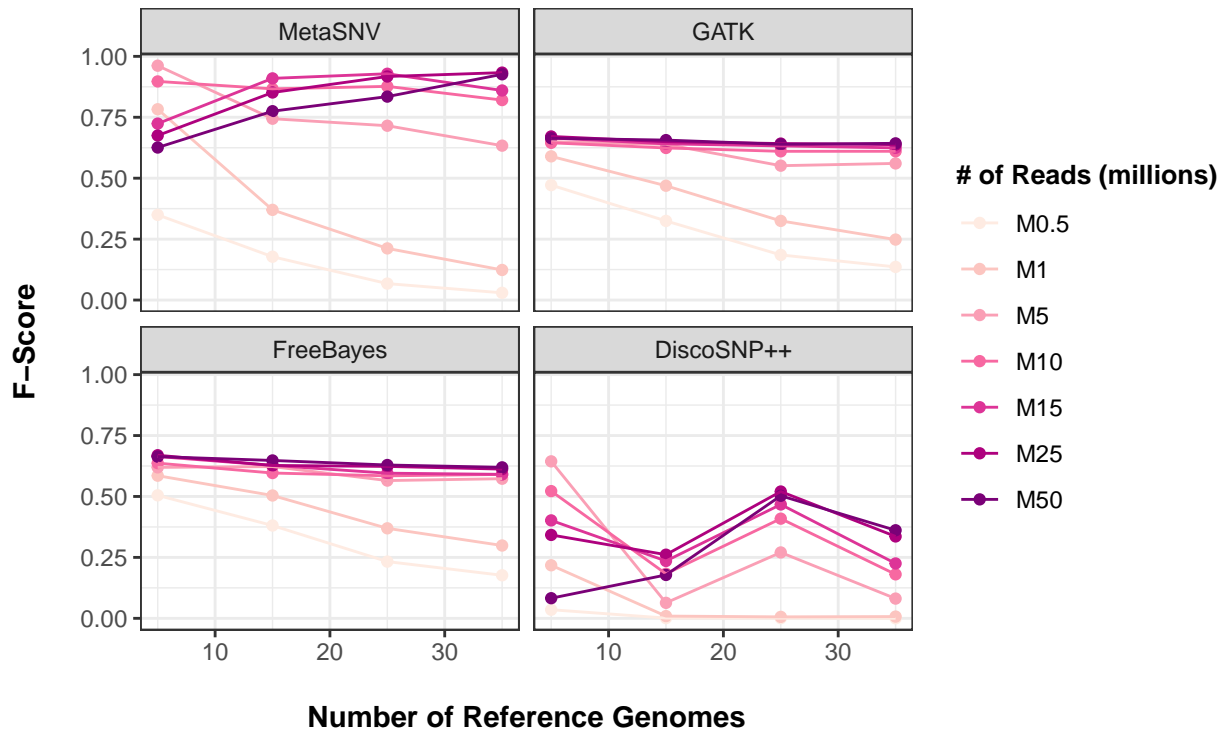
Variant Caller Accuracy by Number of Reads and Reference Genomes



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"))
```

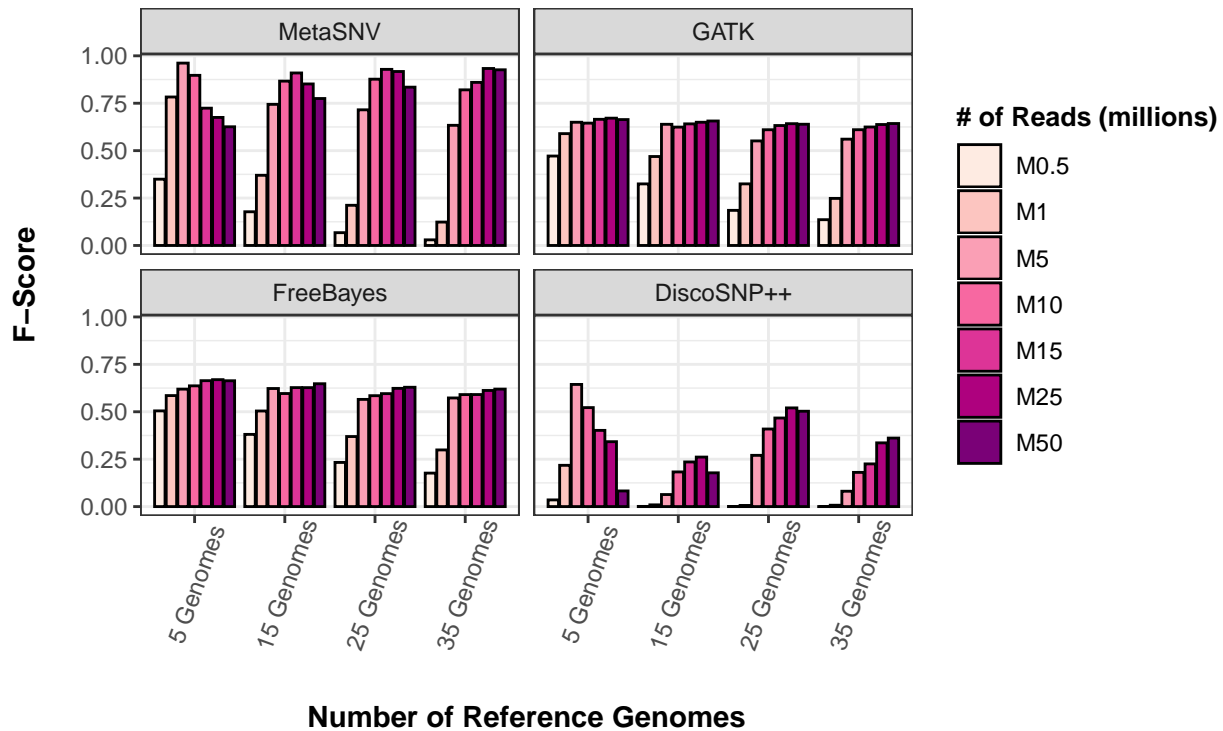
Variant Caller Accuracy by Number of Reads and Reference Genomes



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.text.x = 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"))
```

Variant Caller Accuracy by Number of Reads and Reference Genomes



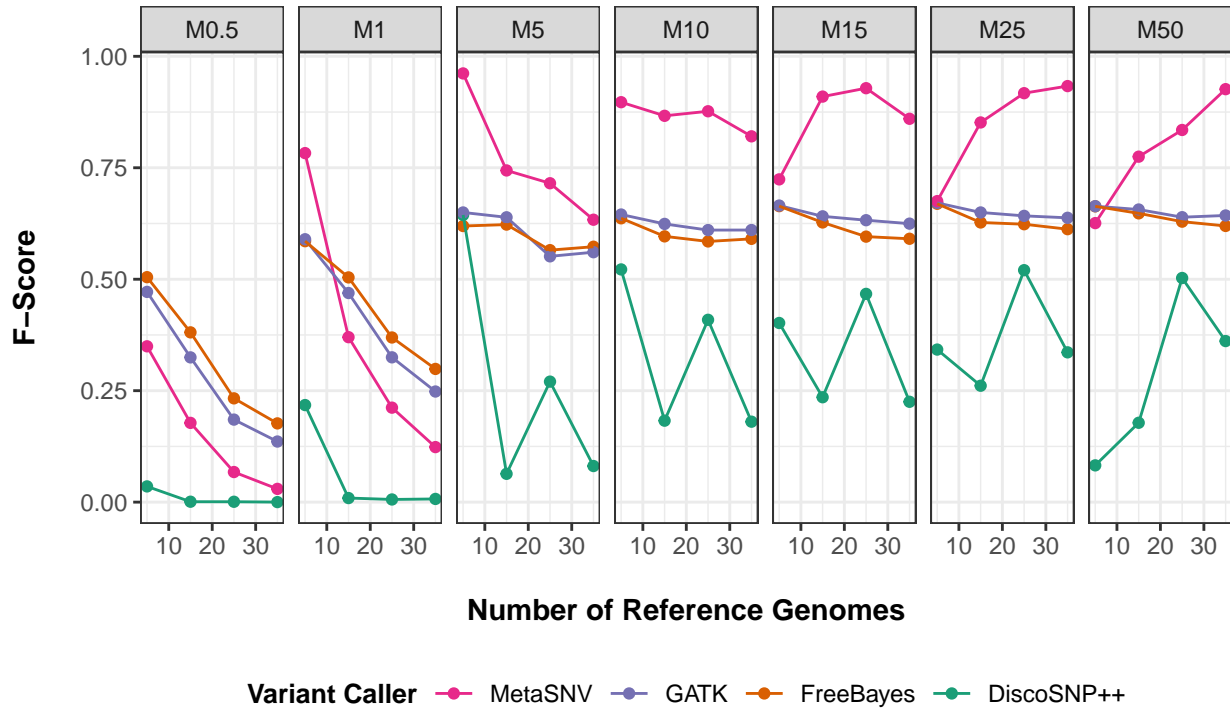
2C

DATASET AS A FACET

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```
###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(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"), legend.position = "bottom")
```

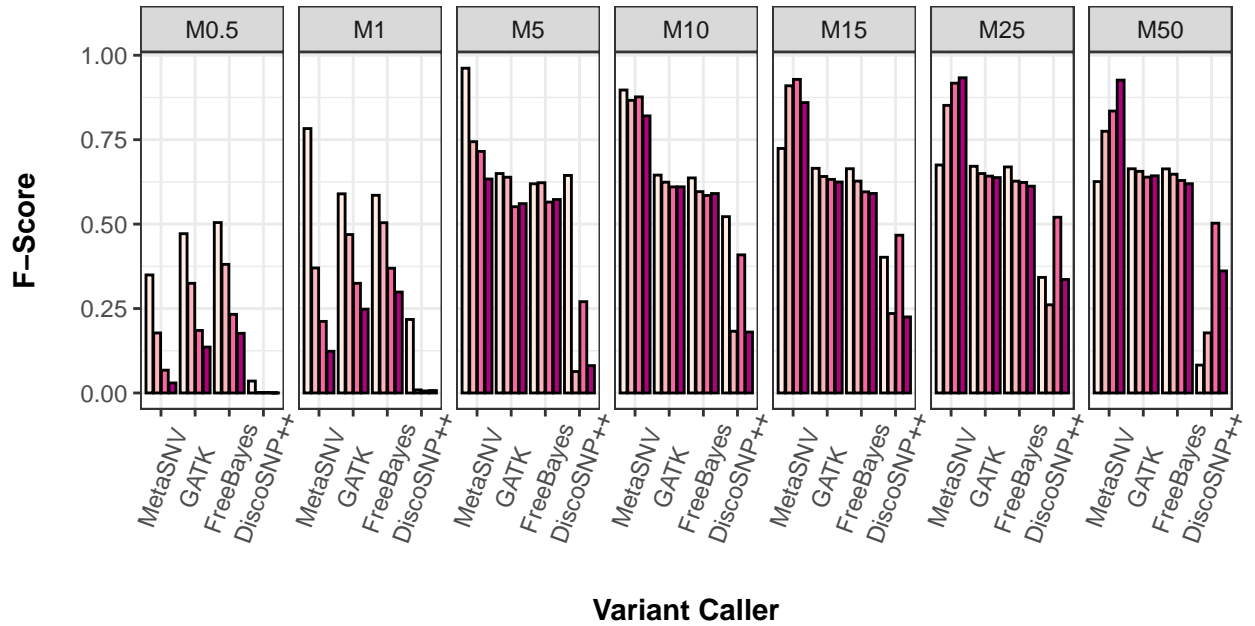
Variant Caller Accuracy by Number of Reads and Reference Genomes



3A

```
#3B
ggplot(bench_df, aes(x=VCaller_f, y=F.score, fill=Subset_f, group=Subset_f)) +
  geom_col(colour="black", position="dodge2") + facet_wrap(~Dataset_f, nrow=1) +
  ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
  xlab("Variant Caller") + ylab("F-Score") + labs(fill="# of Reference Genomes") +
  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.text.x = 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"), legend.position = "bottom")
```


Variant Caller Accuracy by Number of Reads and Reference Genomes



of Reference Genomes 5 Genomes 15 Genomes 25 Genomes 35 Genomes

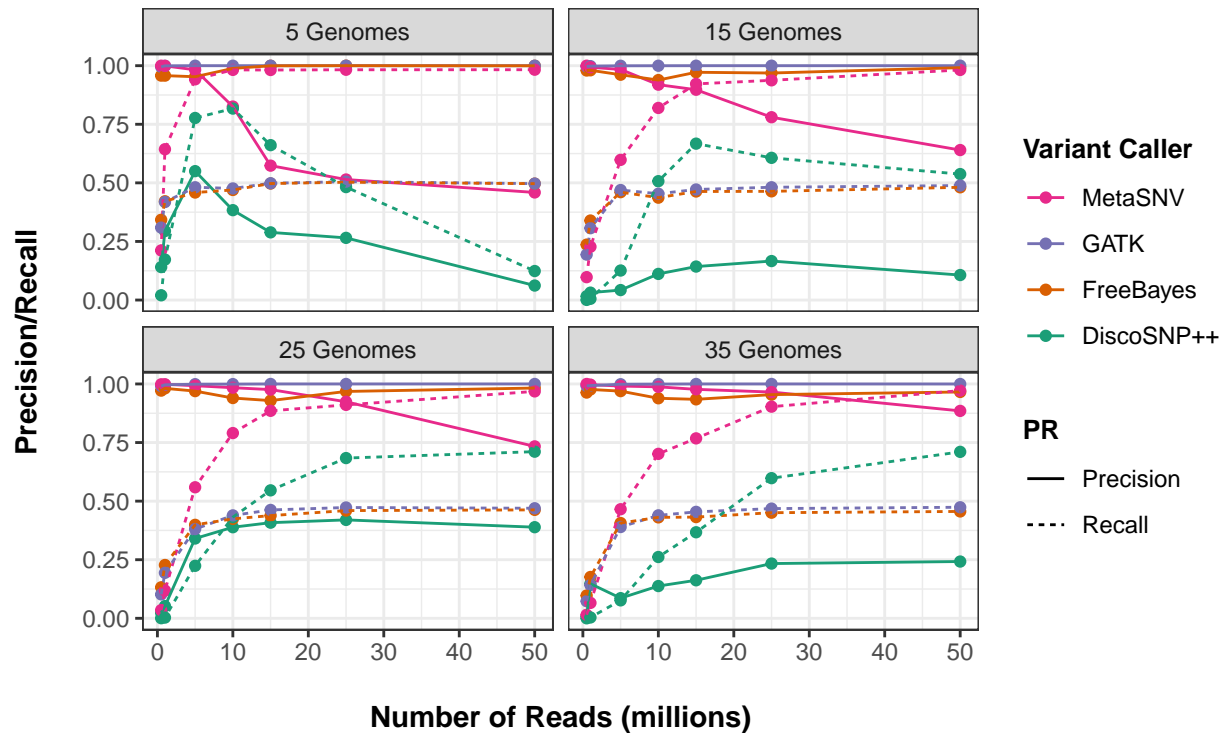
3B

Figures using Precision/Recall

```
bench_melt<-gather(bench_df,"PR","PR_value",7:8)

ggplot(bench_melt,aes(x=Dataset_num,y=PR_value,colour=VCaller_f,group=interaction(VCaller_f,PR))) +
  geom_point() + geom_line(aes(linetype=PR)) + facet_wrap(~Subset_f,nrow=2) +
  ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
  xlab("Number of Reads (millions)") + ylab("Precision/Recall") + 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(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") )
```

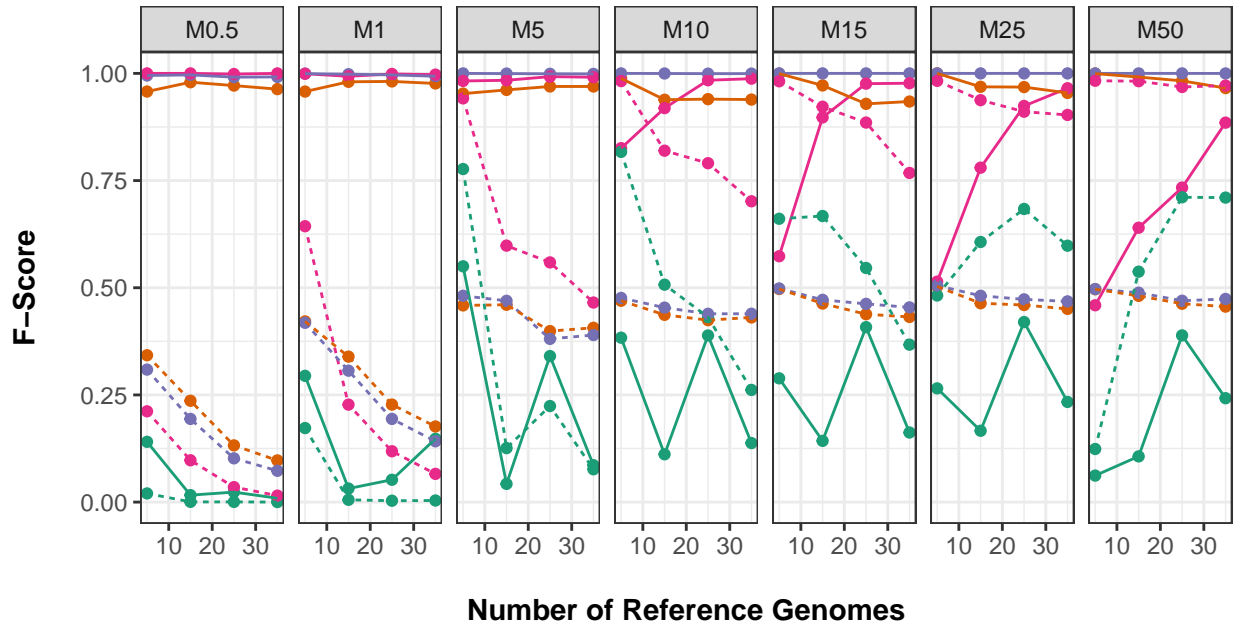
Variant Caller Accuracy by Number of Reads and Reference Genomes



1PR

```
ggplot(bench_melt, aes(x=Subset_num, y=PR_value, colour=VCaller_f, group=interaction(VCaller_f, PR))) +
  geom_point() + geom_line(aes(linetype=PR)) + 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(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"), legend.position = "bottom") #+
```

Variant Caller Accuracy by Number of Reads and Reference Genomes



Variant Caller MetaSNV GATK FreeBayes DiscoSNP++ PR — Precision - - - Recall

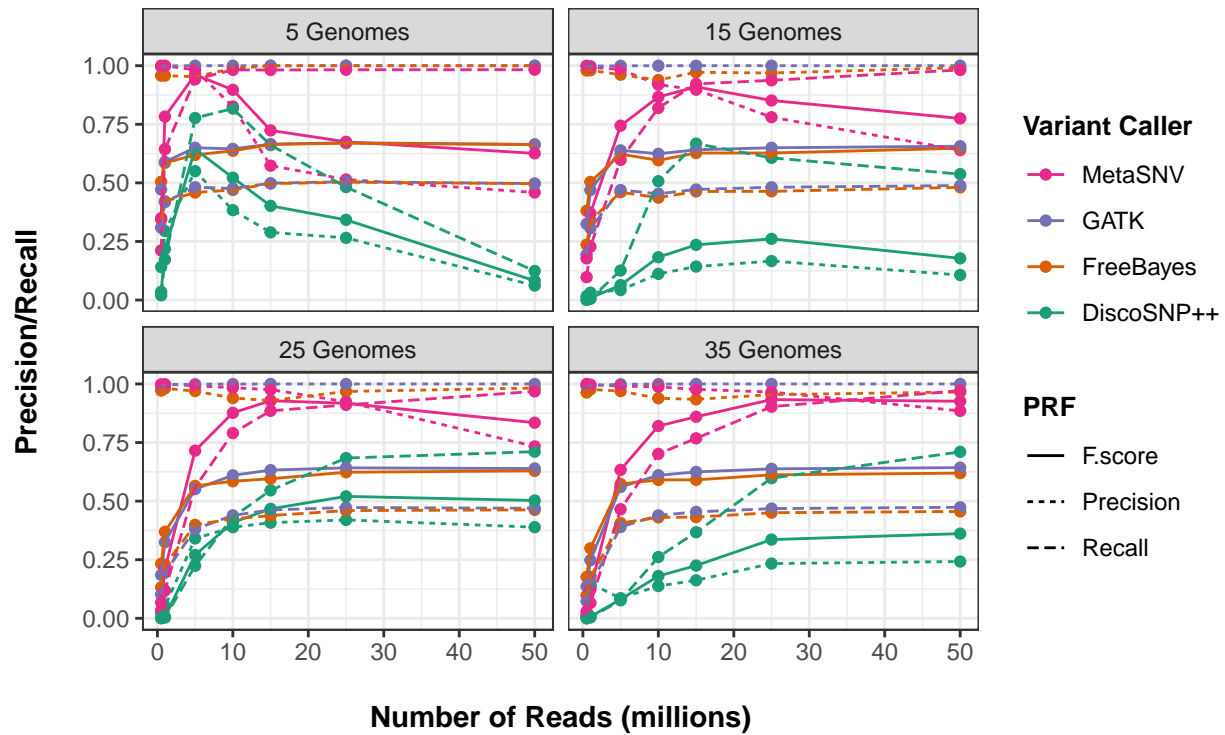
```
#guides(fill=guide_legend(nrow=2,byrow=TRUE))
```

3PR

```
bench_super_melt<-gather(bench_df, "PRF", "PRF_value", 7:9)

ggplot(bench_super_melt, aes(x=Dataset_num, y=PRF_value, colour=VCaller_f, group=interaction(VCaller_f, PRF))) +
  geom_point() + geom_line(aes(linetype=PRF)) + facet_wrap(~Subset_f, nrow=2) +
  ggtitle("Variant Caller Accuracy by Number of Reads\n and Reference Genomes") +
  xlab("Number of Reads (millions)") + ylab("Precision/Recall") + 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(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 Accuracy by Number of Reads and Reference Genomes



1PRF