

ClinVar Report

James Diao

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Contents

1	Collect and Merge ClinVar Data	2
1.1	Import ClinVar VCF	2
1.2	Merge ClinVar with 1000 Genomes and ExAC	2
2	Summary Statistics	3
2.1	Fraction of Individuals with Pathogenic Non-Reference Sites	3
3	Penetrance Estimates	4
3.1	Max/Min Penetrance as a Function of $P(D)$ and $P(V D)$	4
3.2	Penetrance Estimates by Ancestry	5
3.3	Empirical CDFs for All Penetrance Plots	7
3.4	Comparing Mean Penetrance between ExAC and 1000 Genomes	7

Sourcing ClinVar input from: clinvar_2016-01-04.vcf

Sending output to: Report_2016-01-04.pdf

1 Collect and Merge ClinVar Data

1.1 Import ClinVar VCF

```
## Processed ClinVar data frame 94971 x 14 (selected rows/columns):
```

1.2 Merge ClinVar with 1000 Genomes and ExAC

```
## Breakdown of ClinVar Variants
```

Subset_ClinVar	Number_of_Variants
Total ClinVar	94971
LP/P-ClinVar	29342
LP/P-ClinVar & ACMG	5923
LP/P-ClinVar & ACMG & ExAC	938
LP/P-ClinVar & ACMG & 1000 Genomes	165

```
## Breakdown of ACMG-1000 Genomes Variants
```

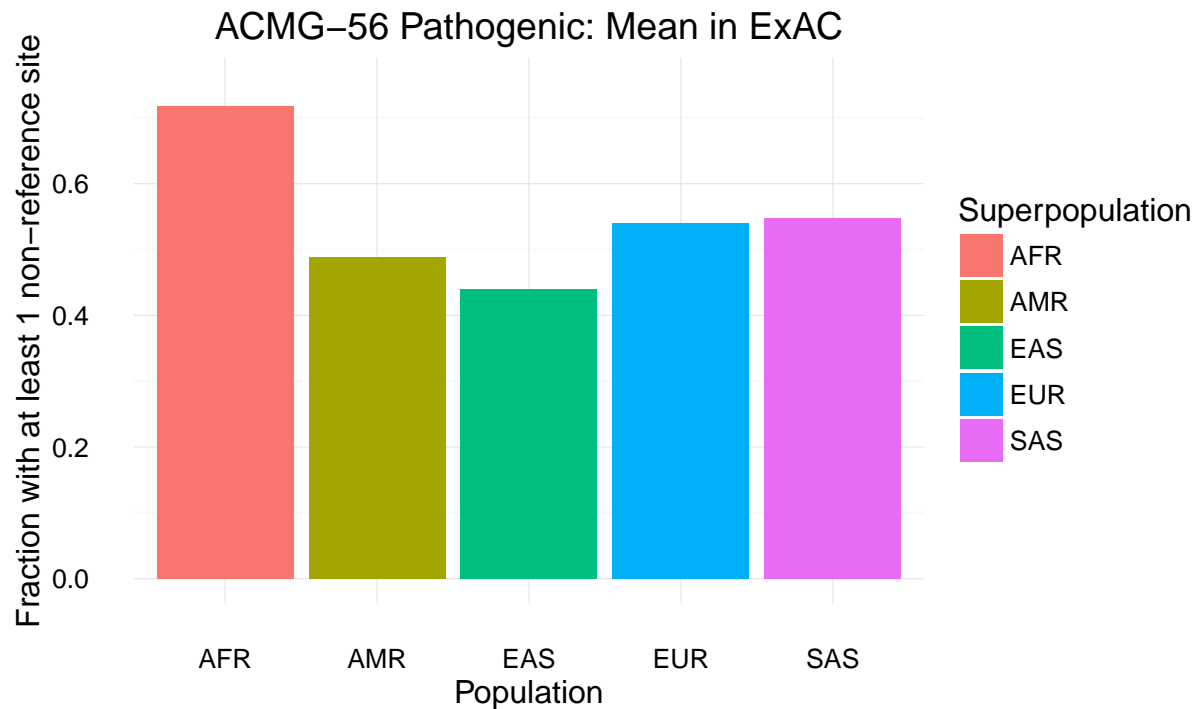
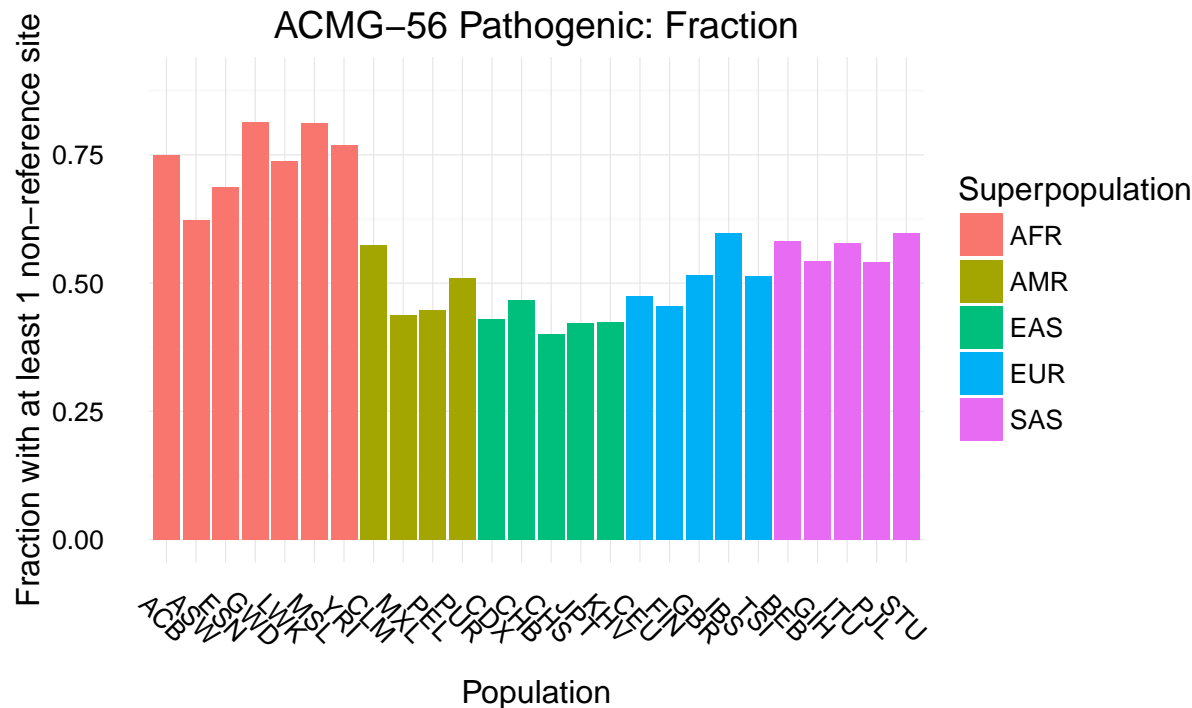
Subset_1000_Genomes	Number_of_Variants
Total 1000_Genomes & ACMG	139335
1000_Genomes & ACMG & ClinVar	3840
1000_Genomes & ACMG & LP/P-ClinVar	165

```
## Breakdown of ACMG-ExAC Variants
```

Subset_ExAC	Number_of_Variants
Total ExAC & ACMG	58873
ExAC & ACMG & ClinVar	7777
ExAC & ACMG & LP/P-ClinVar	938

2 Summary Statistics

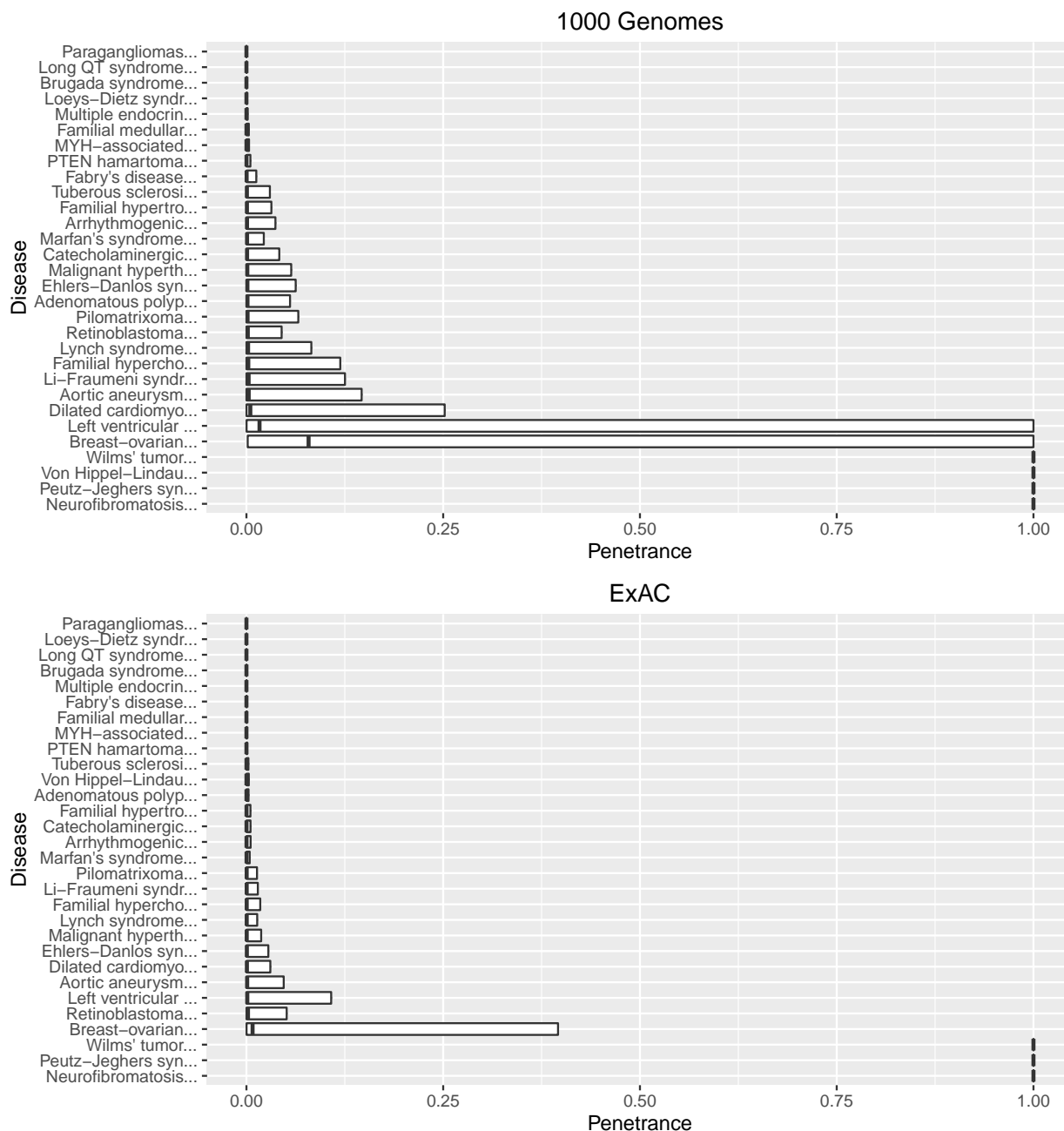
2.1 Fraction of Individuals with Pathogenic Non-Reference Sites



3 Penetrance Estimates

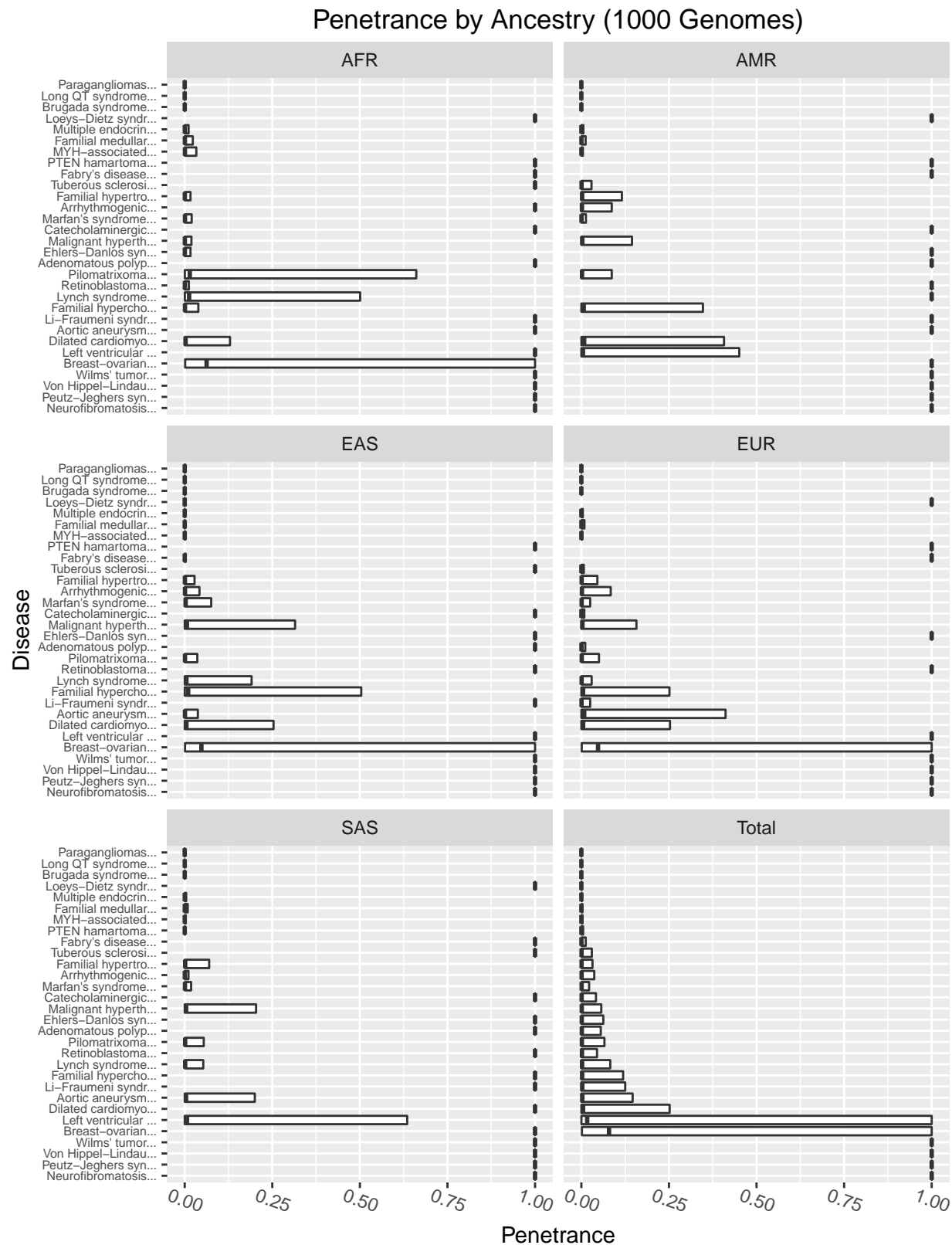
3.1 Max/Min Penetrance as a Function of $P(D)$ and $P(V|D)$

The left end of the boxplot indicates $P(D)$ AND $P(V|D)$ = lower value,
the bold line in the middle indicates $P(D)$ AND $P(V|D)$ = geometric_mean(values),
the right end of the boxplot indicates $P(D)$ AND $P(V|D)$ = upper value.

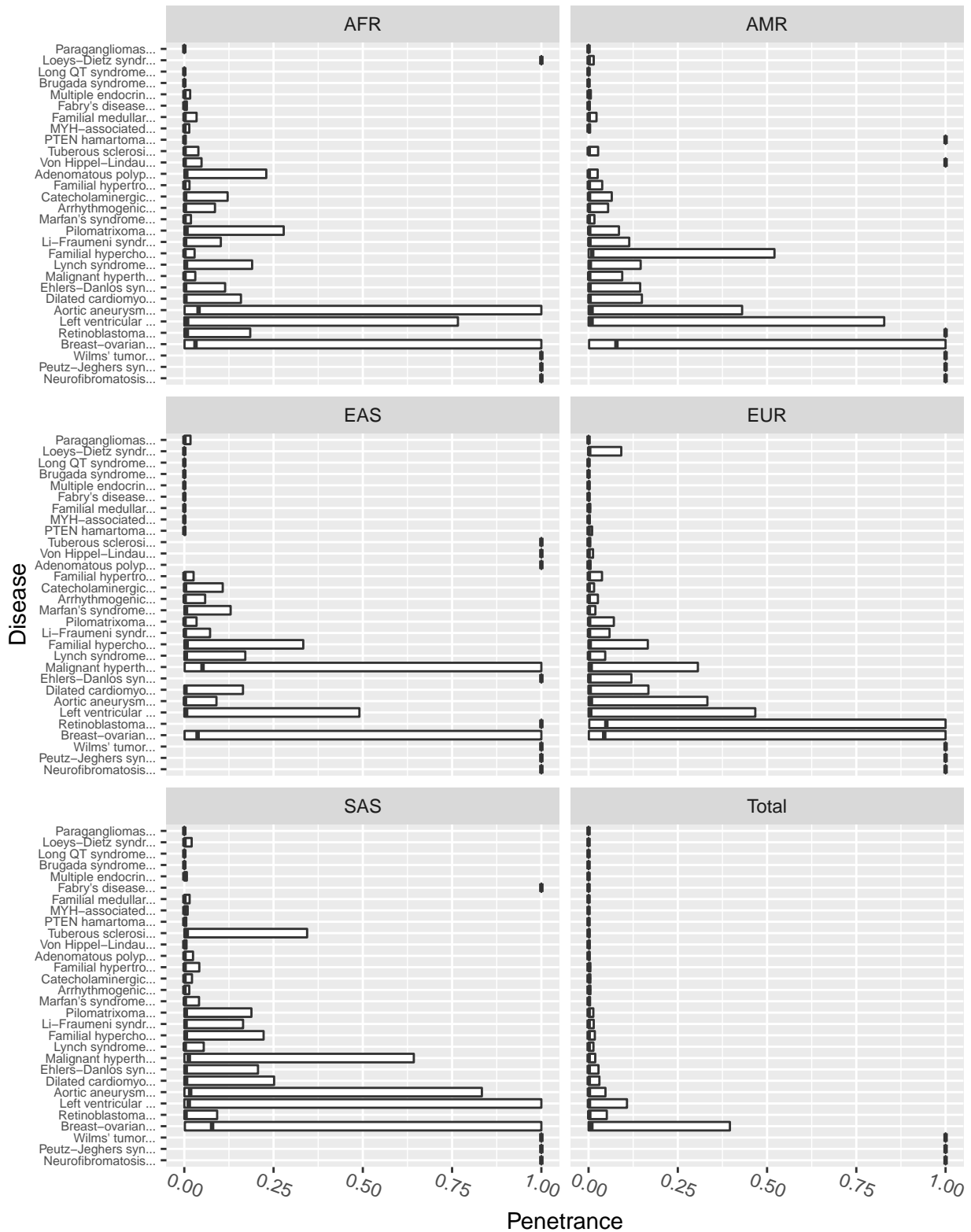


Note: Prevalence ranges of 5x were assumed for all point estimates of prevalence.
For example: a point estimate of 0.022 would be given the range 0.01-0.05.

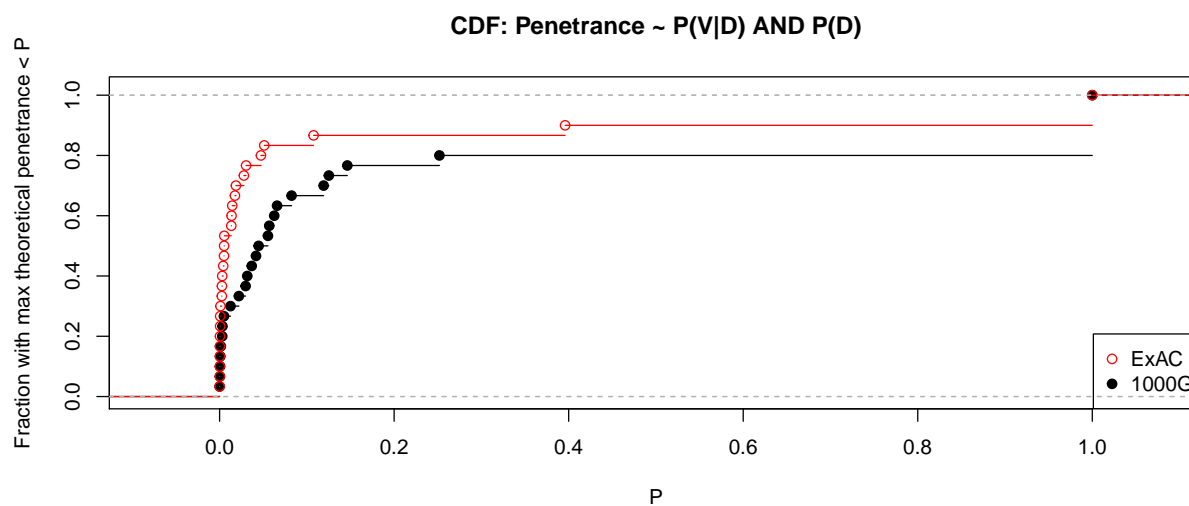
3.2 Penetrance Estimates by Ancestry



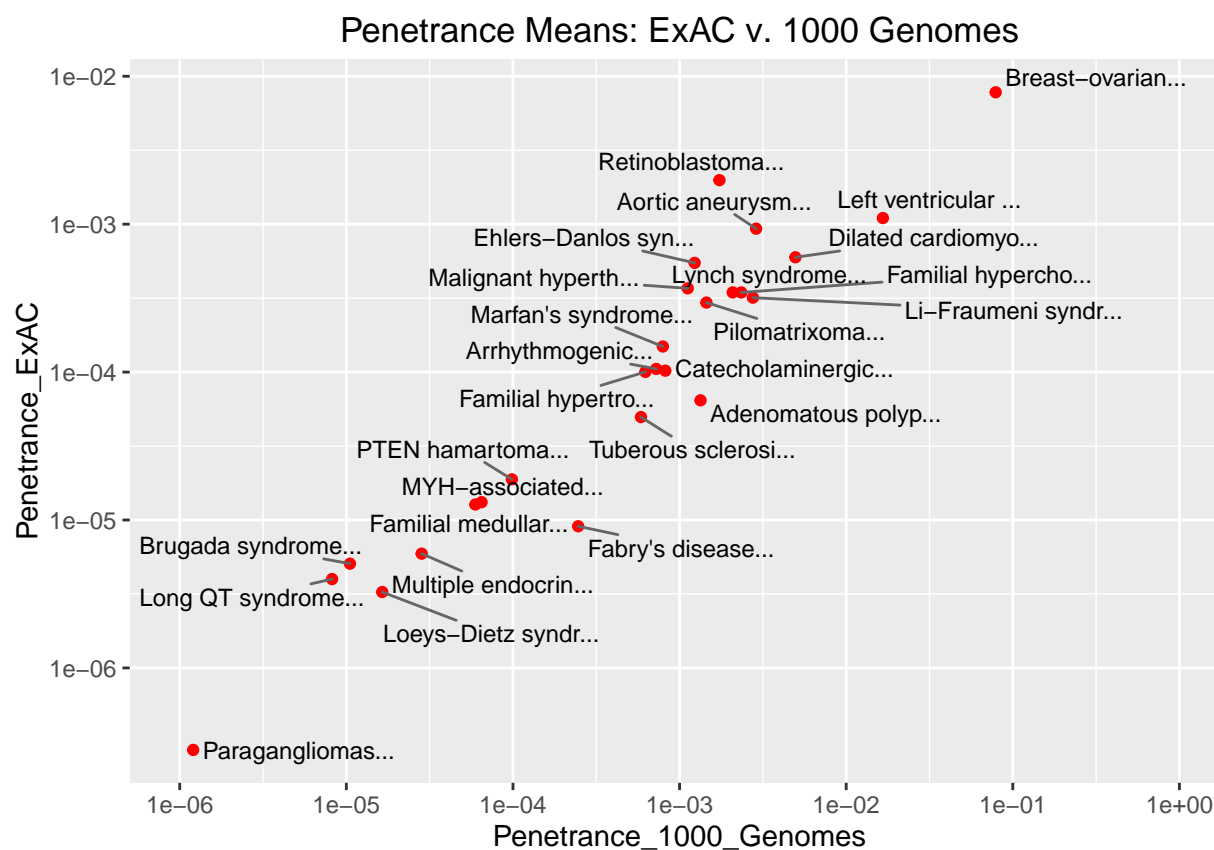
Penetrance by Ancestry (ExAC)



3.3 Empirical CDFs for All Penetrance Plots



3.4 Comparing Mean Penetrance between ExAC and 1000 Genomes



The Pearson correlation is 0.97.

Max penetrance values computed using 1000 Genomes are 9.8-fold larger than those computed using ExAC.