

ClinVar Report

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Sourcing ClinVar input from: clinvar_2015-11-02.vcf

Sending output to: Report_2015-11-02.pdf

1 Collect and Merge ClinVar Data

1.1 Import ClinVar VCF

Processed ClinVar data frame 93163 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	93163
LP/P-ClinVar	28627
LP/P-ClinVar & ACMG	5965
LP/P-ClinVar & ACMG & ExAC	985
LP/P-ClinVar & ACMG & 1000 Genomes	185

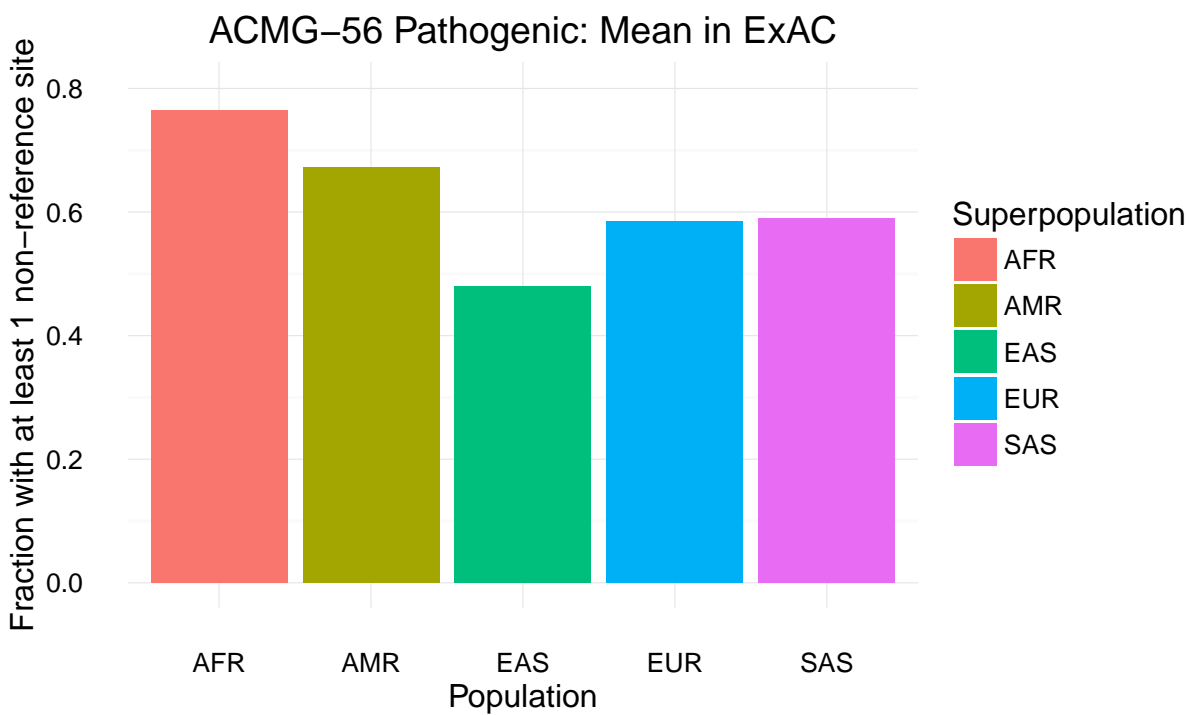
Breakdown of ACMG-1000 Genomes Variants

Subset_1000_Genomes	Number_of_Variants
Total 1000_Genomes & ACMG	139335
1000_Genomes & ACMG & ClinVar	3807
1000_Genomes & ACMG & LP/P-ClinVar	185

Breakdown of ACMG-ExAC Variants

Subset_ExAC	Number_of_Variants
Total ExAC & ACMG	58873
ExAC & ACMG & ClinVar	7714
ExAC & ACMG & LP/P-ClinVar	985

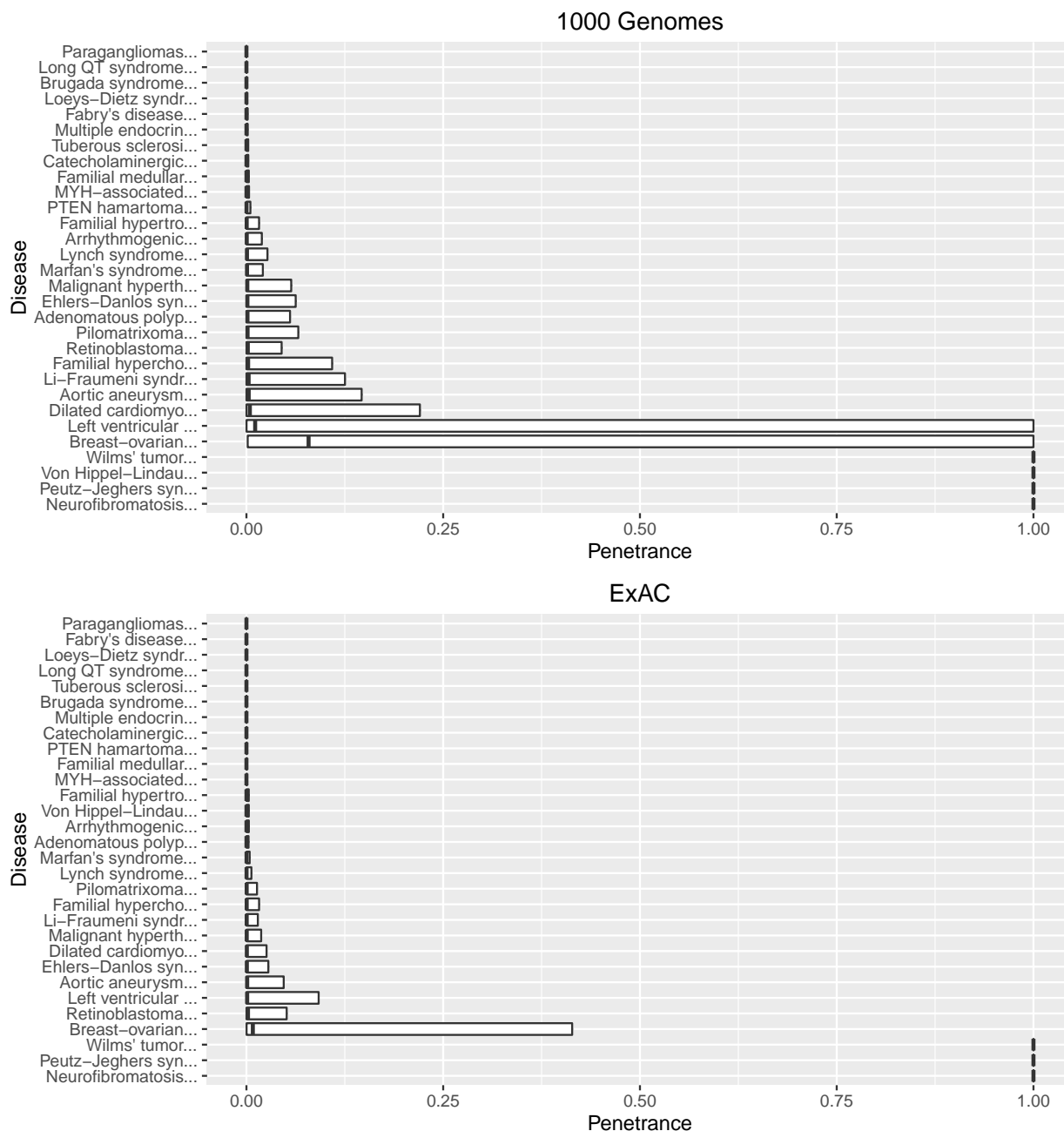
2.1 Fraction of Individuals with Pathogenic Non-Reference Sites



3 Penetrance Estimates

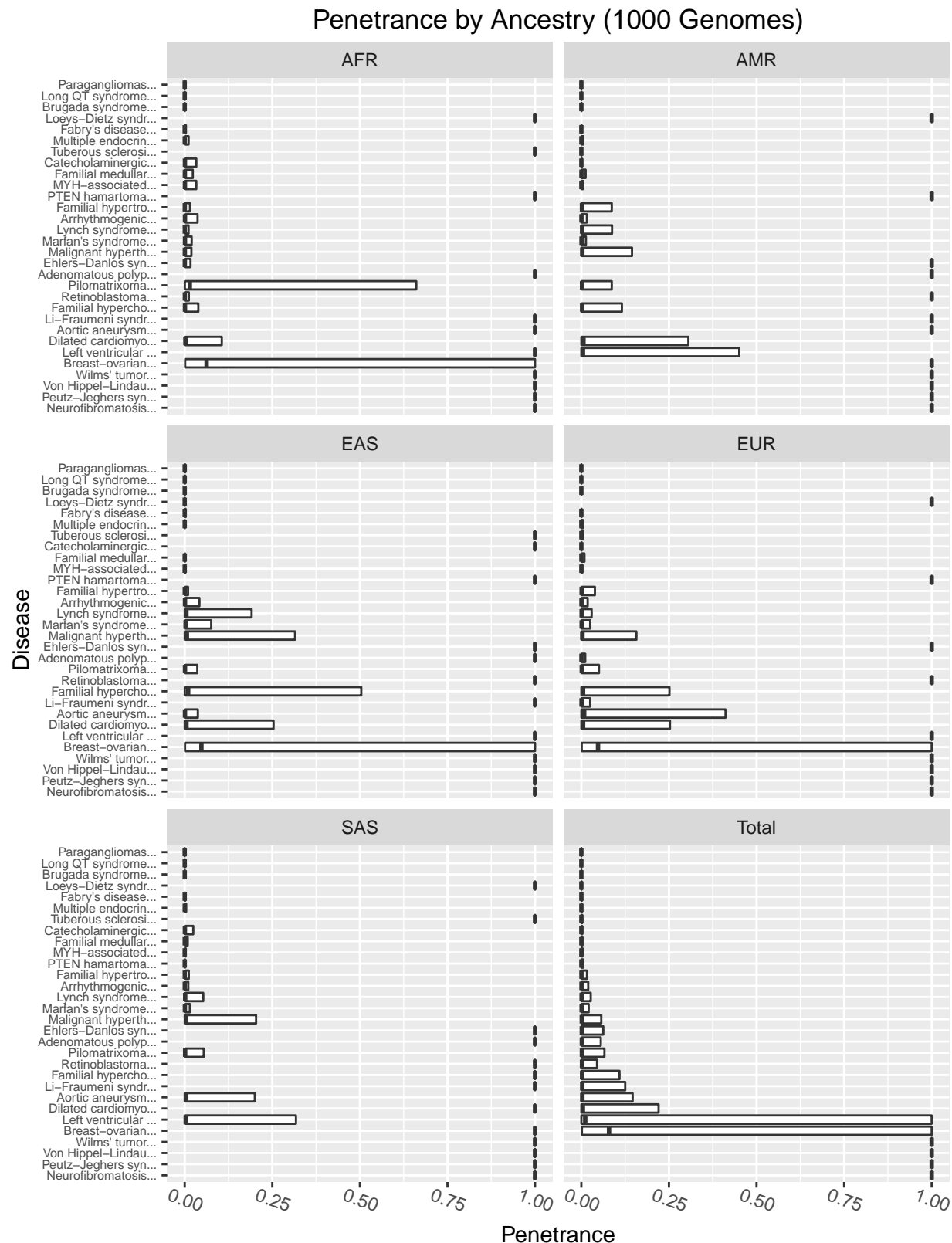
3.1 Max/Min Penetrance as a Function of $P(D)$ and $P(V|D)$

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.

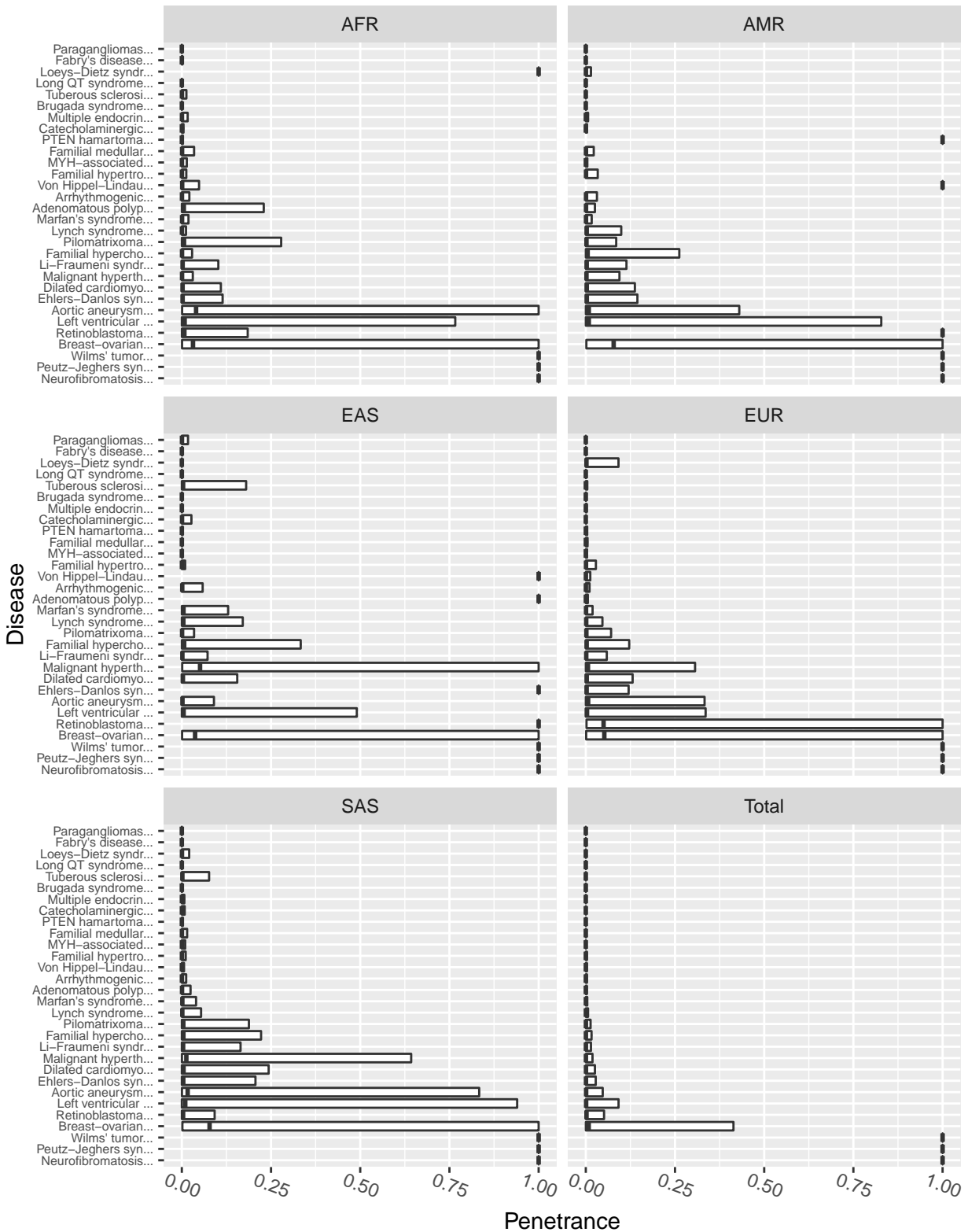


Note: Prevalence ranges of 5x were assumed for all point estimates of prevalence.
For example: a point estimate of 0.022 would be given the range 0.01-0.05.

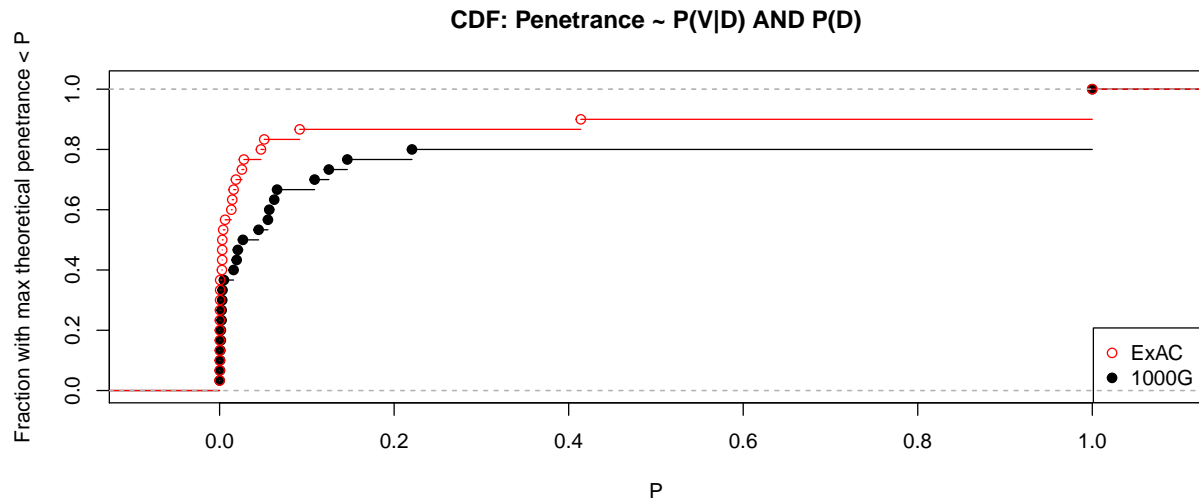
3.2 Penetrance Estimates by Ancestry



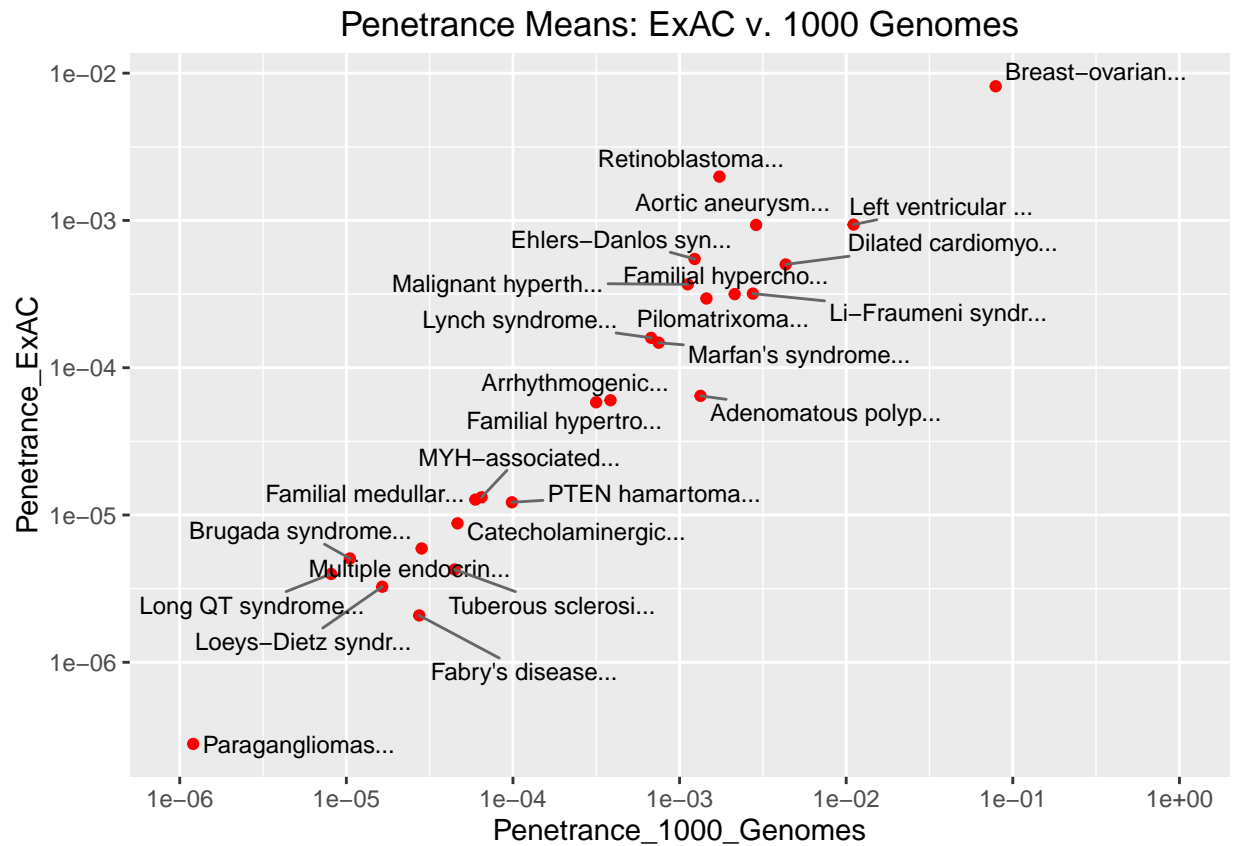
Penetrance by Ancestry (ExAC)



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 9.3-fold larger than those computed using ExAC.