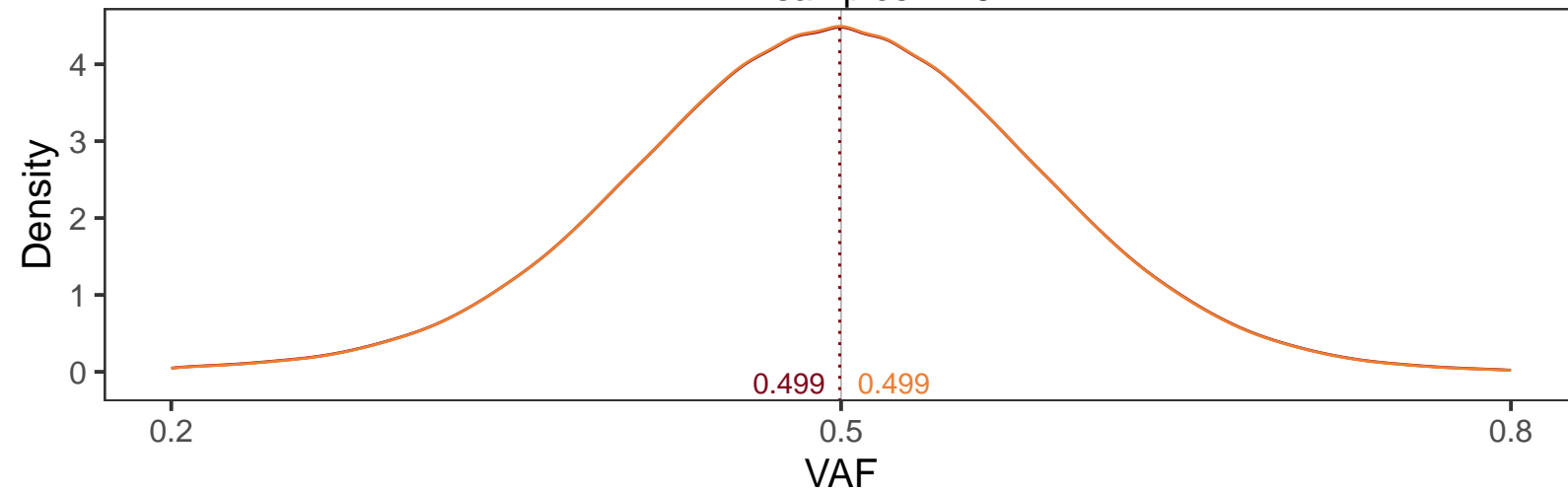


HD 1kGP WGS (30x)

N samples = 25

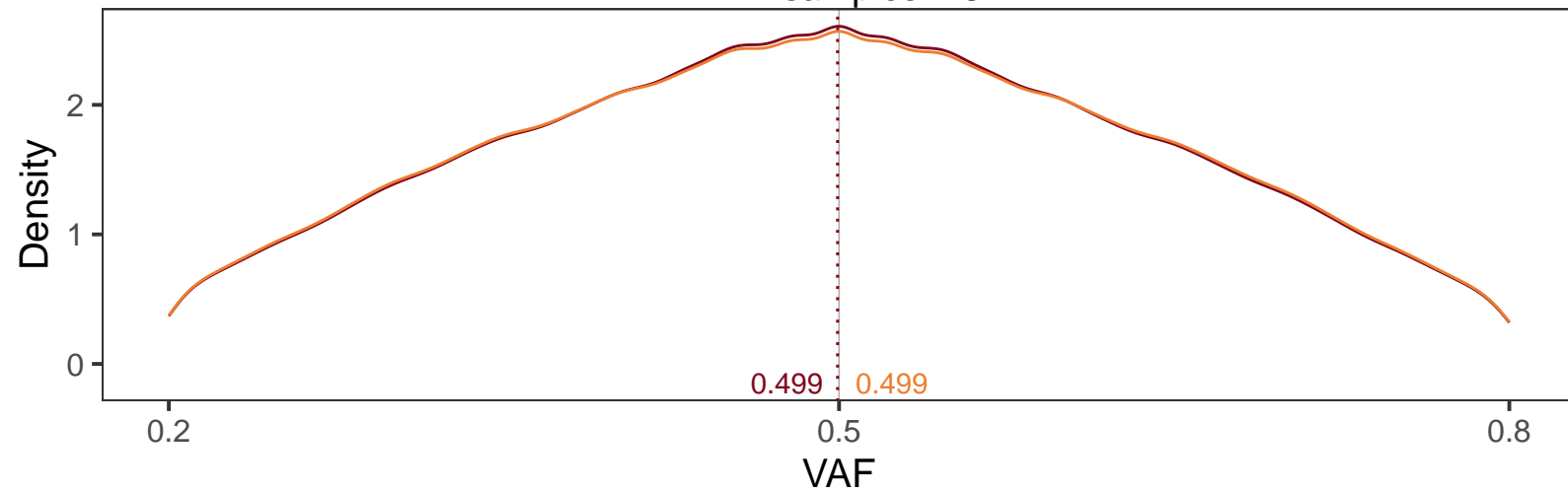


N SNPs = 43531.71*1e3



Tumor CCLE WGS (30x) – wt regions

N samples = 8



N SNPs = 8064.08*1e3



Assembly version

hg38 T2T-CHM13

Allele swapping

VAF coherent

VAF incoherent