

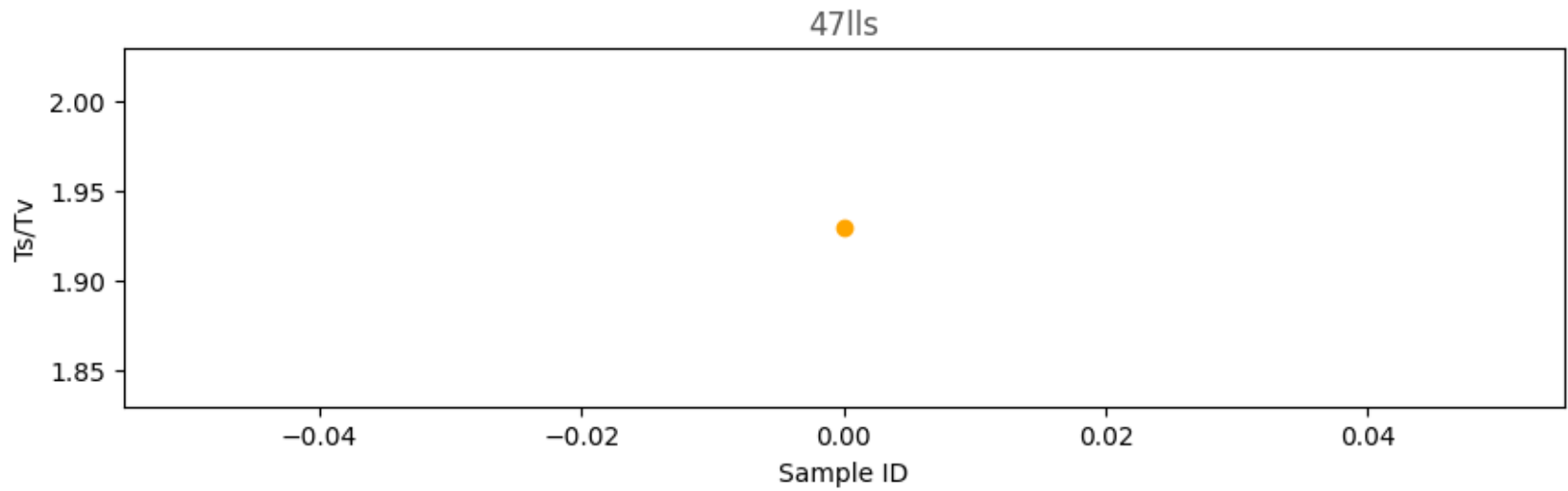
Summary Numbers

| Callset | SNPs | | | indels | | MNPs | others |
|----------------------------------|-----------|-------|-----------|---------|------|------|--------|
| | n | ts/tv | (1st ALT) | n | frm* | | |
| 47lls | 4,086,961 | 1.93 | 1.93 | 954,287 | – | 0 | 0 |
| * frameshift ratio: out/(out+in) | | | | | | | |

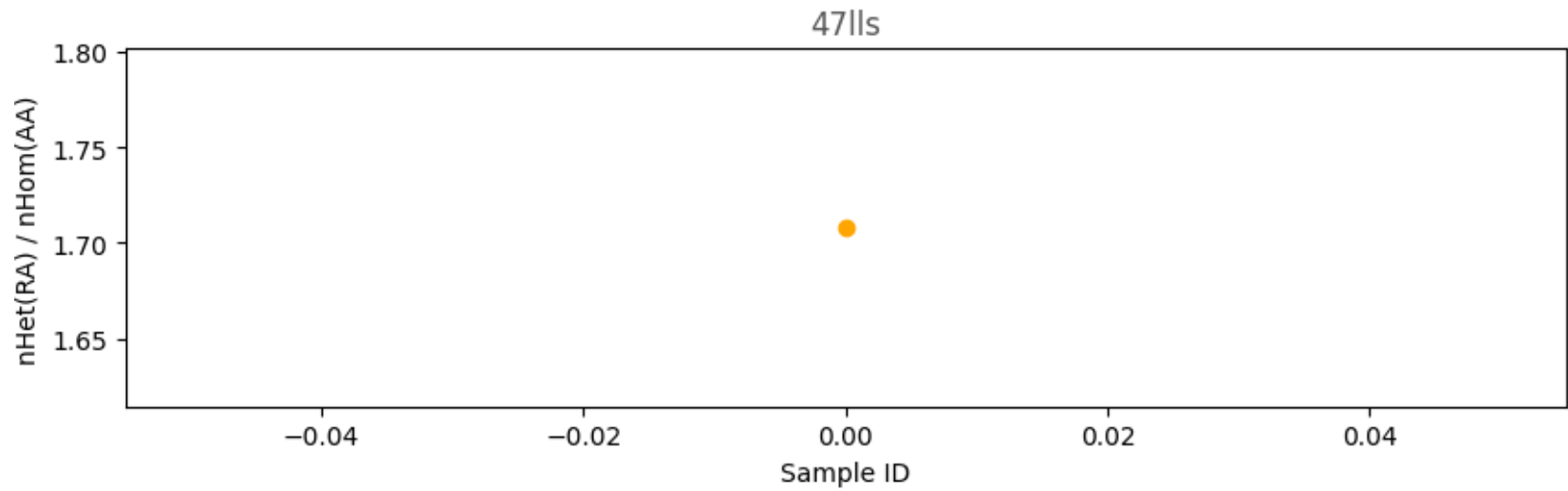
| Callset | singletons (AC=1) | | | multiallelic | |
|---------|-------------------|-------|--------|--------------|-------|
| | SNPs | ts/tv | indels | sites | SNPs |
| 47lls | 63.1% | 1.90 | 68.1% | 101,068 | 2,060 |

- 47lls .. /ngc/projects2/gm/data/archive/2022/variants/snv/47llsherf-103910771476-Normal_Blood_noinfo-WGS_v1_IlluminaDNAPCRFree_RHGM01558-220916_A01411 BHYNLVDSX3-EXT_LAB
KA_NGCWGS-NGCWGS05359_snv_germline_raw.haplotype_caller.vcf.gz

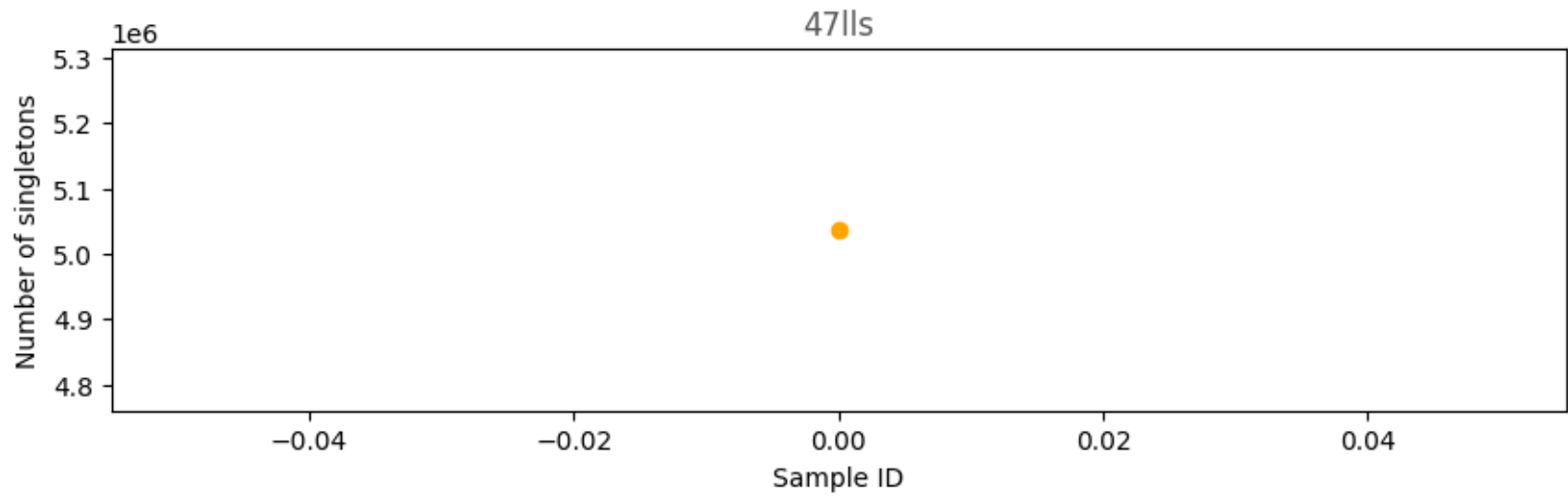
Ts/Tv by sample



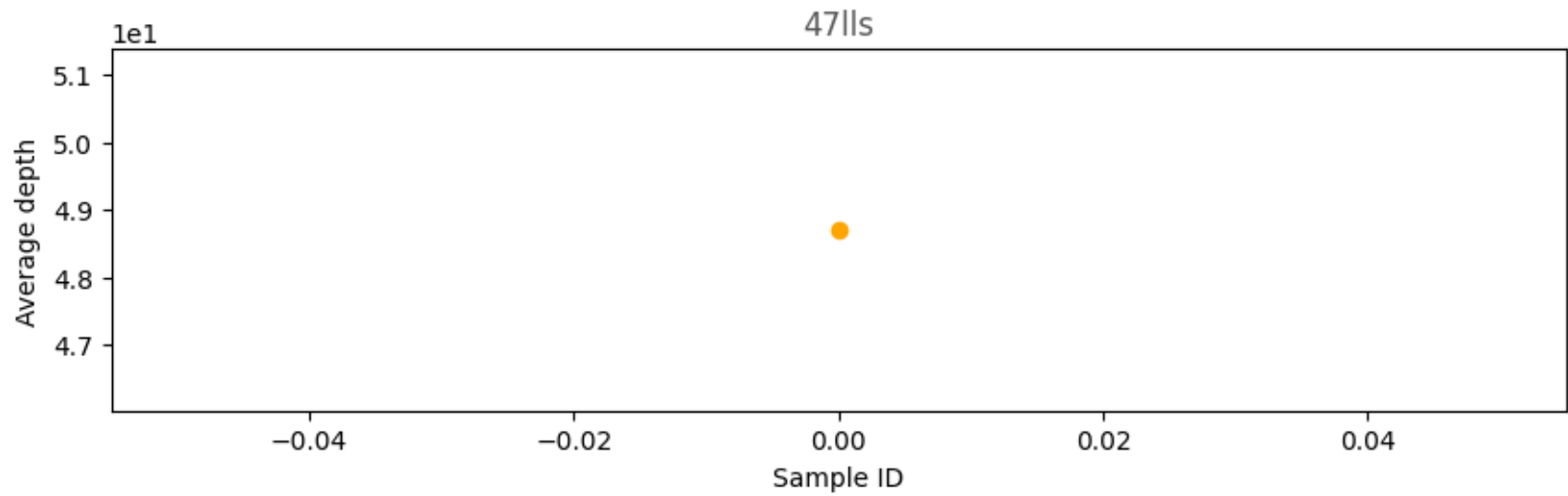
Hets vs non-ref Homs by sample



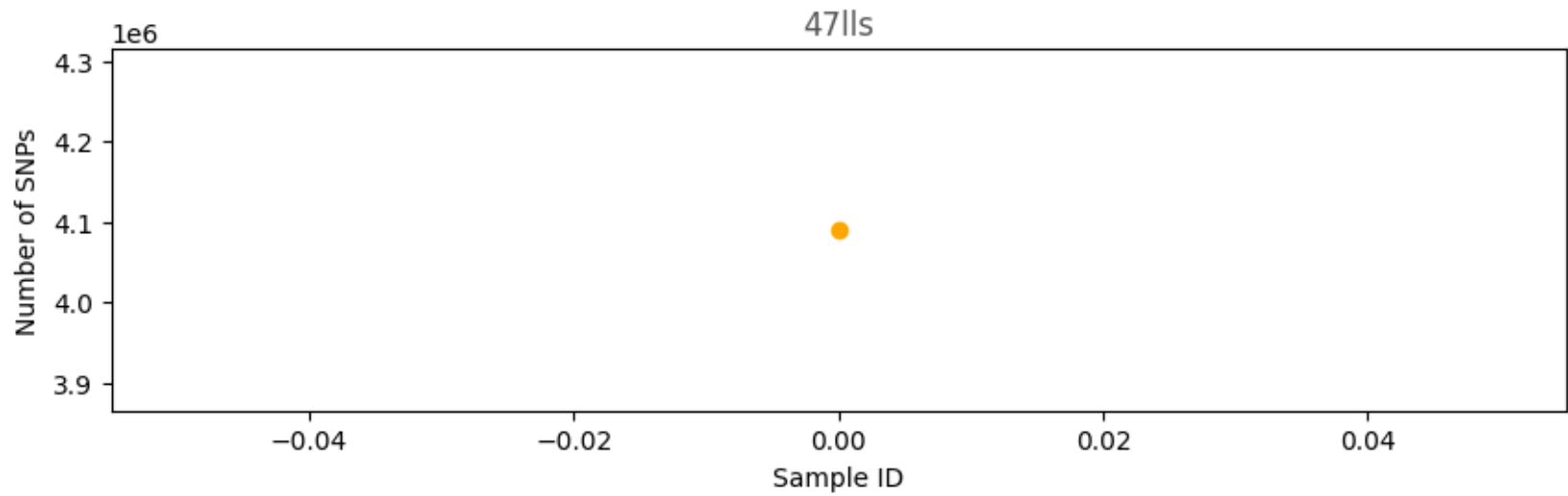
Singletons by sample (hets and homs)



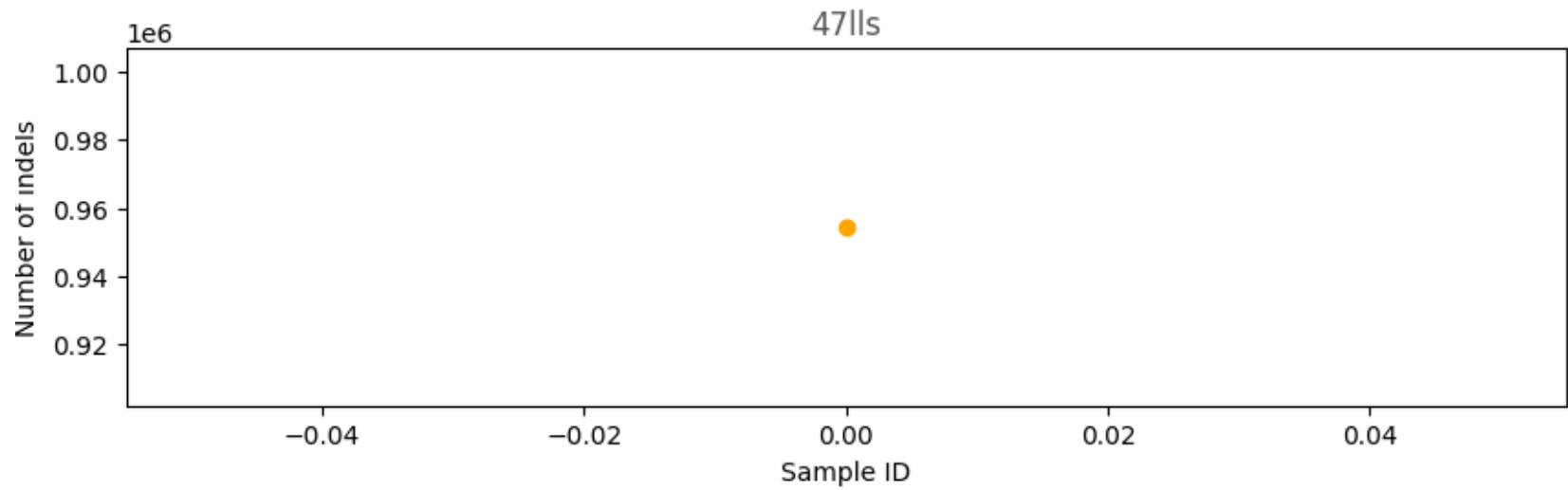
Average depth by sample



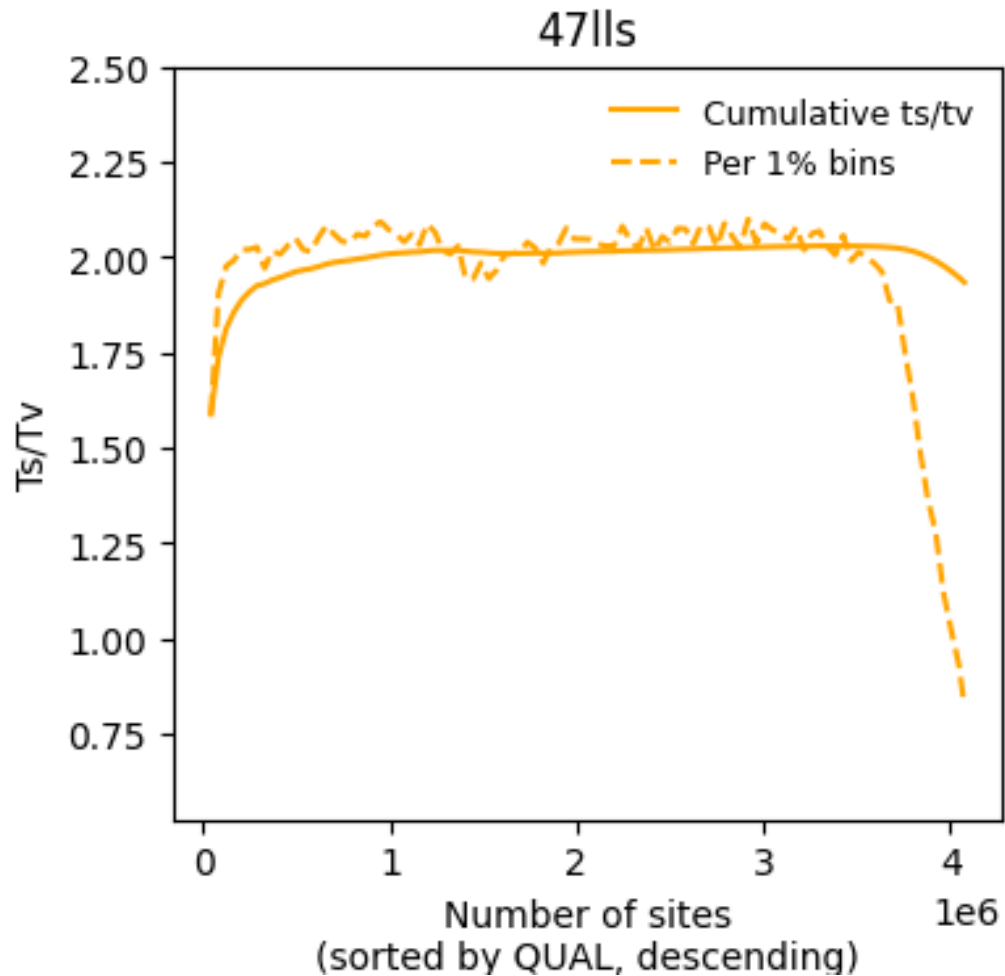
Number of SNPs by sample



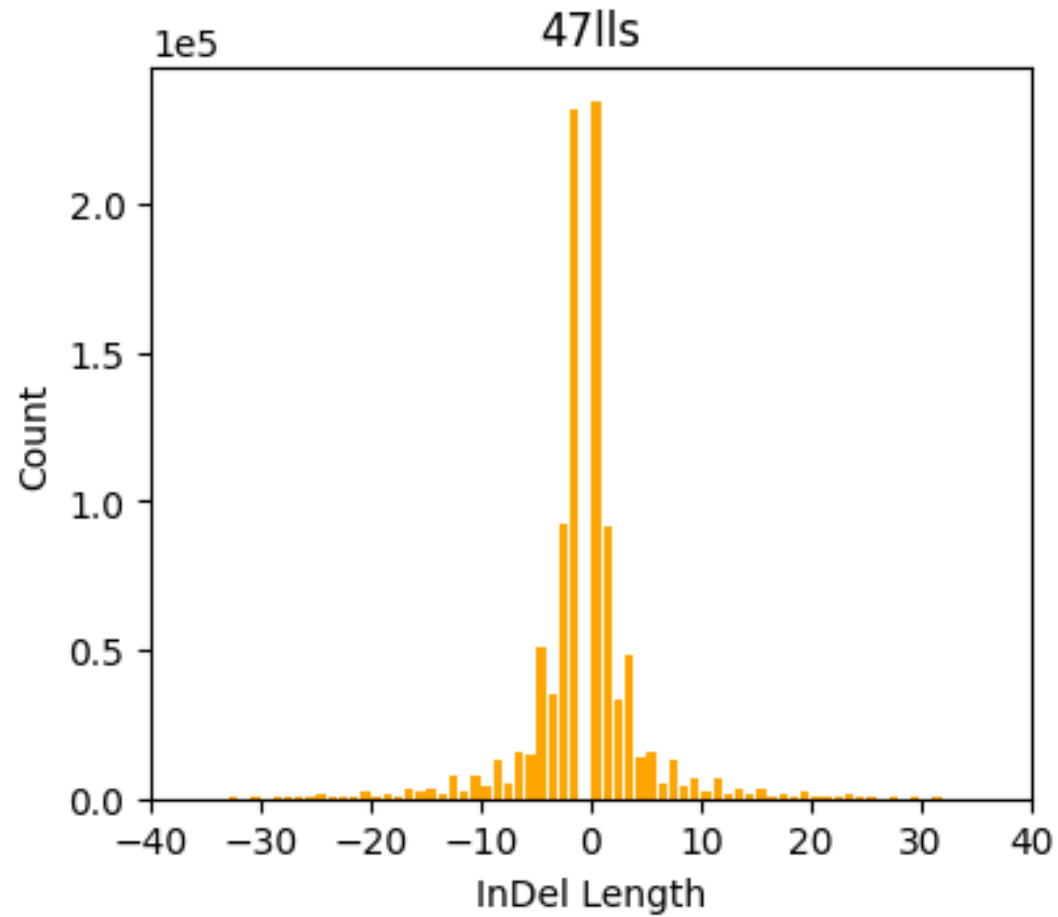
Number of indels by sample



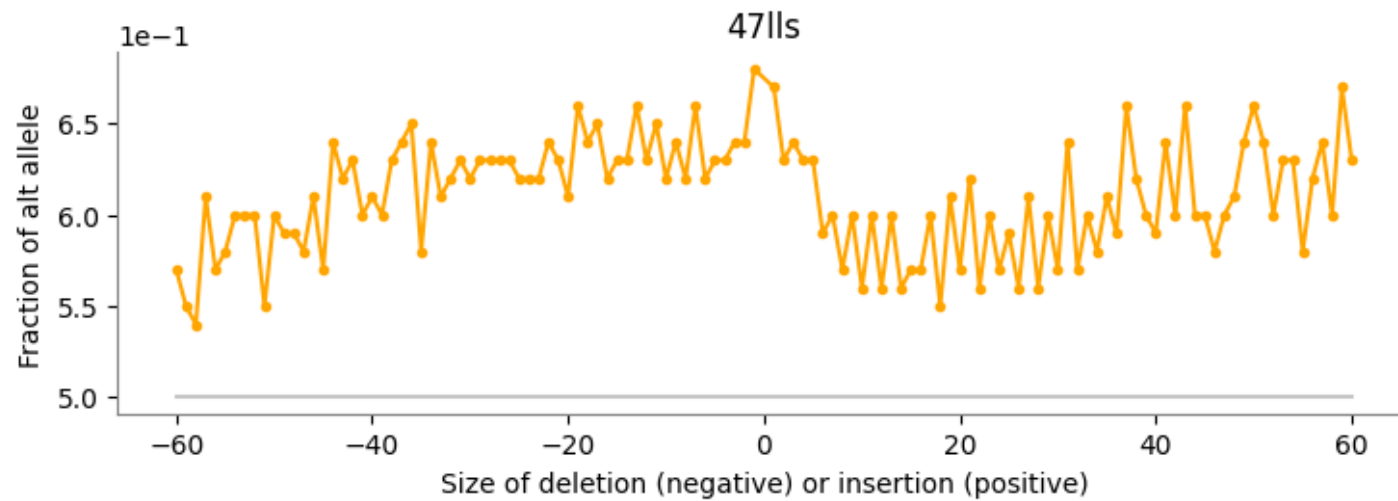
Ts/Tv stratified by QUAL



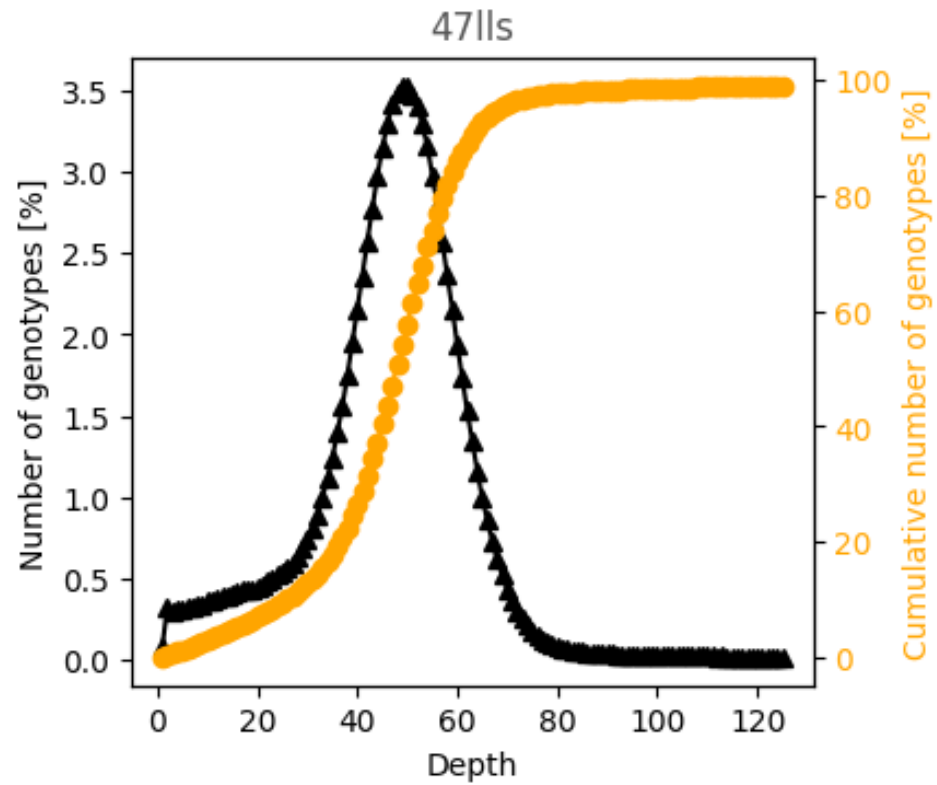
Indel distribution



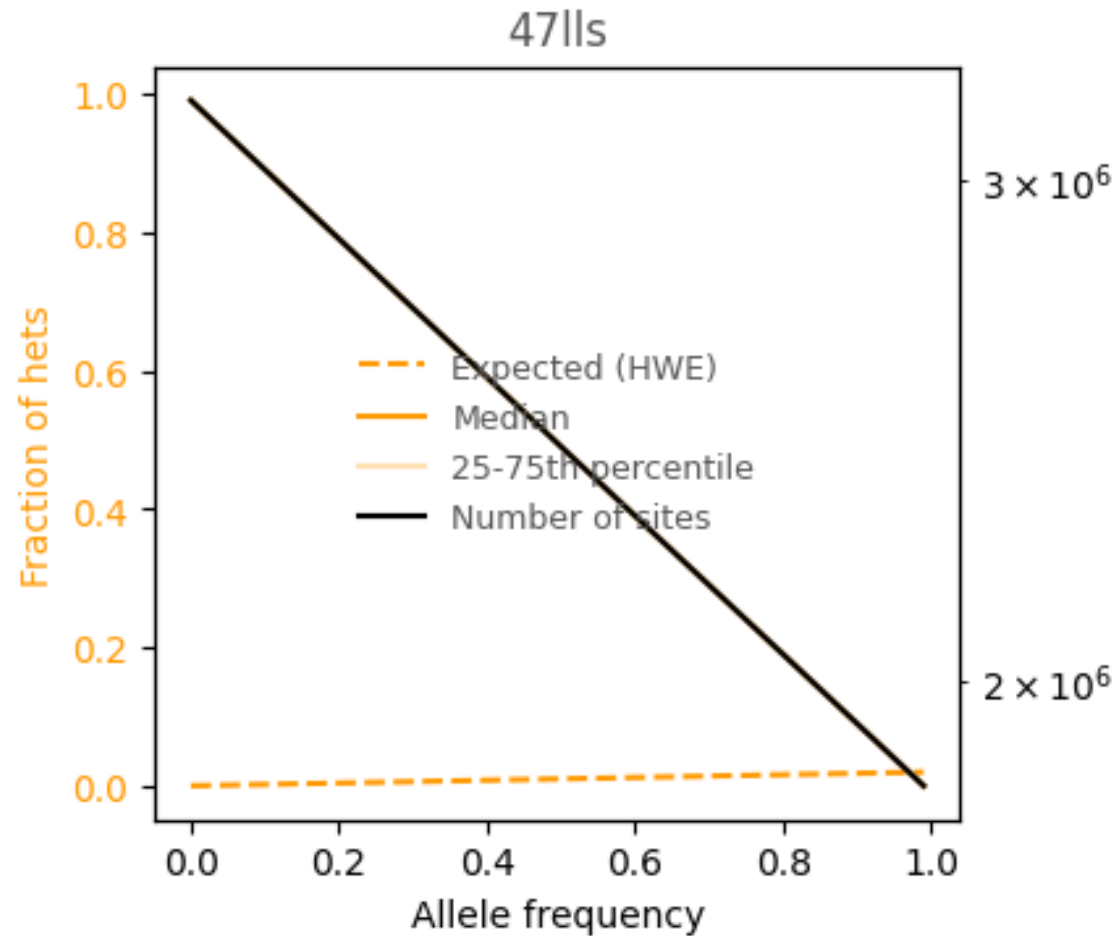
Fraction of alternate indel allele



Depth distribution



Number of HETs by AF



Substitution types

