

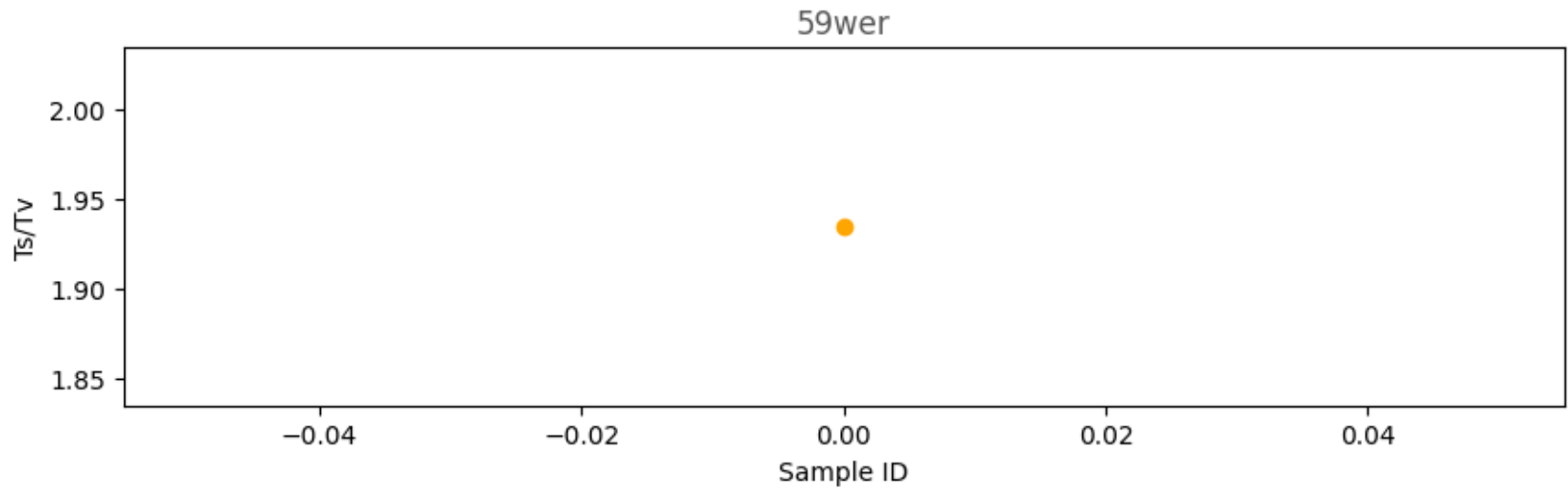
# Summary Numbers

| Callset                          | SNPs      |       |           | indels  |      | MNPs | others |
|----------------------------------|-----------|-------|-----------|---------|------|------|--------|
|                                  | n         | ts/tv | (1st ALT) | n       | frm* |      |        |
| 59wer                            | 4,073,330 | 1.93  | 1.94      | 939,829 | –    | 0    | 0      |
| * frameshift ratio: out/(out+in) |           |       |           |         |      |      |        |

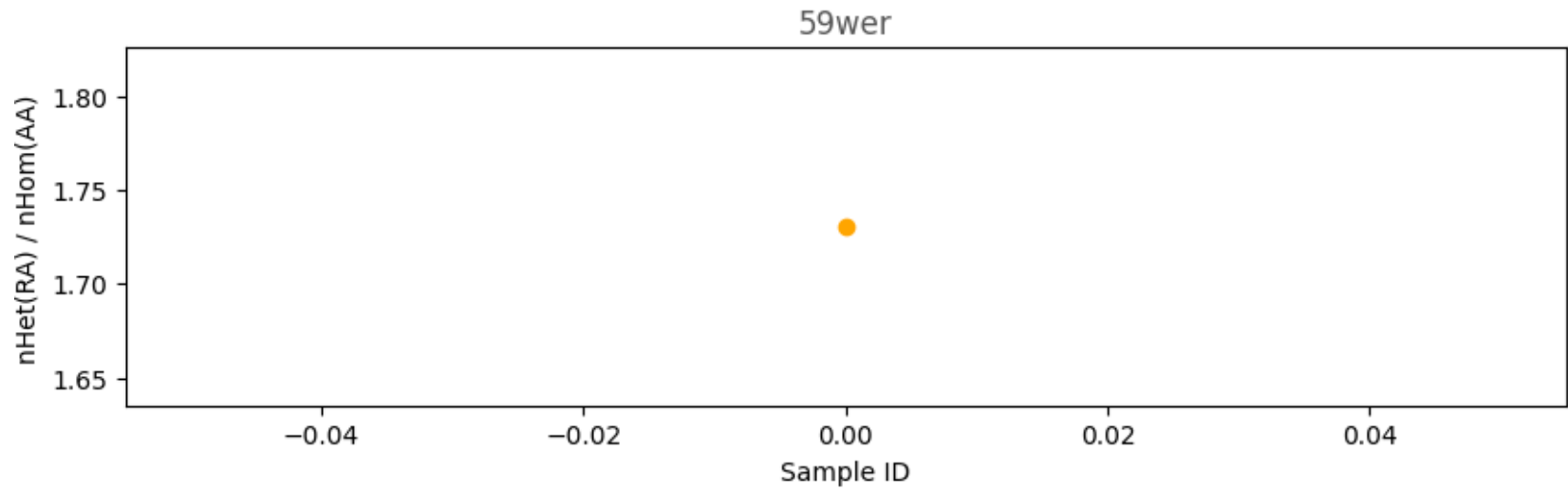
| Callset | singletons (AC=1) |       |        | multiallelic |       |
|---------|-------------------|-------|--------|--------------|-------|
|         | SNPs              | ts/tv | indels | sites        | SNPs  |
| 59wer   | 63.4%             | 1.91  | 67.8%  | 93,861       | 2,047 |

- 59wer .. /ngc/projects2/gm/data/archive/2022/variants/snv/59weribaf-103833689964-Normal\_Blood\_noinfo-WGS\_v1\_IlluminaDNAPCRFree\_X-220202\_A01176\_AH52HTDMXY-RHGM\_LABKA\_WGS-WGS03783\_snv\_germline\_raw.haplotype\_caller.vcf.gz

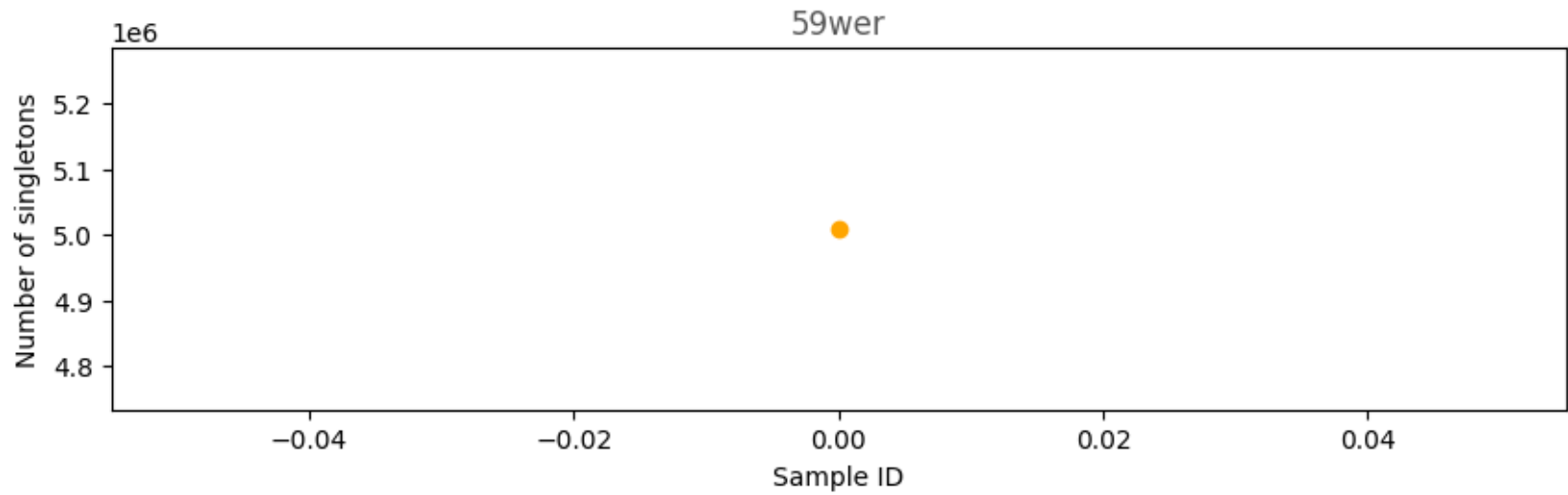
# Ts/Tv by sample



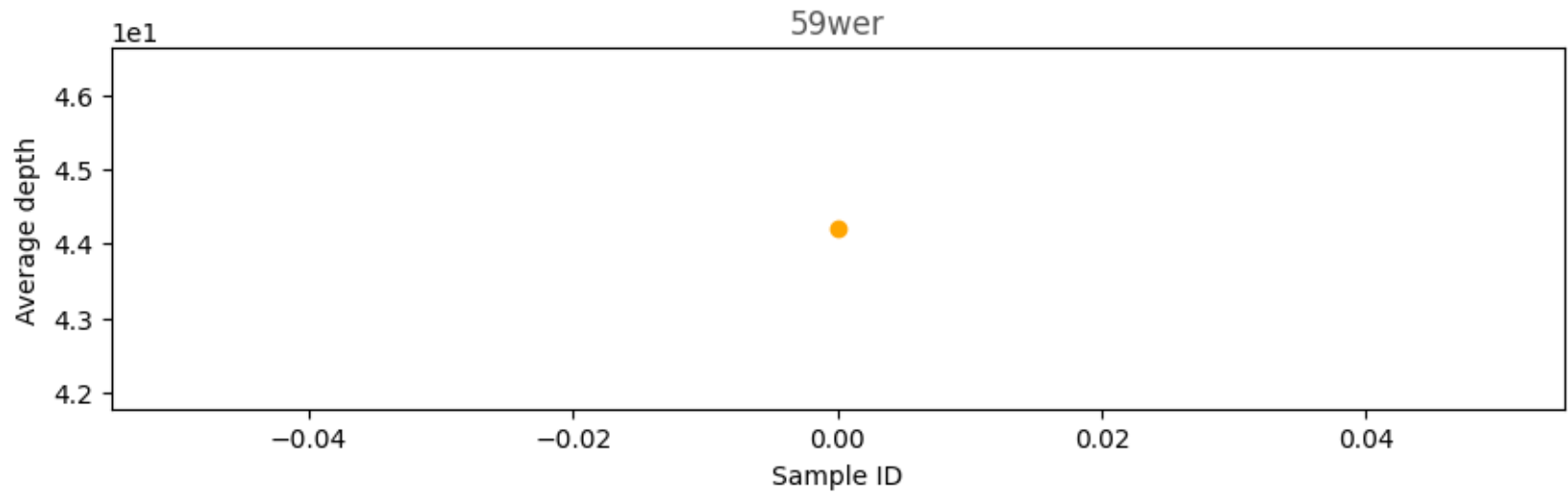
# Hets vs non-ref Homs by sample



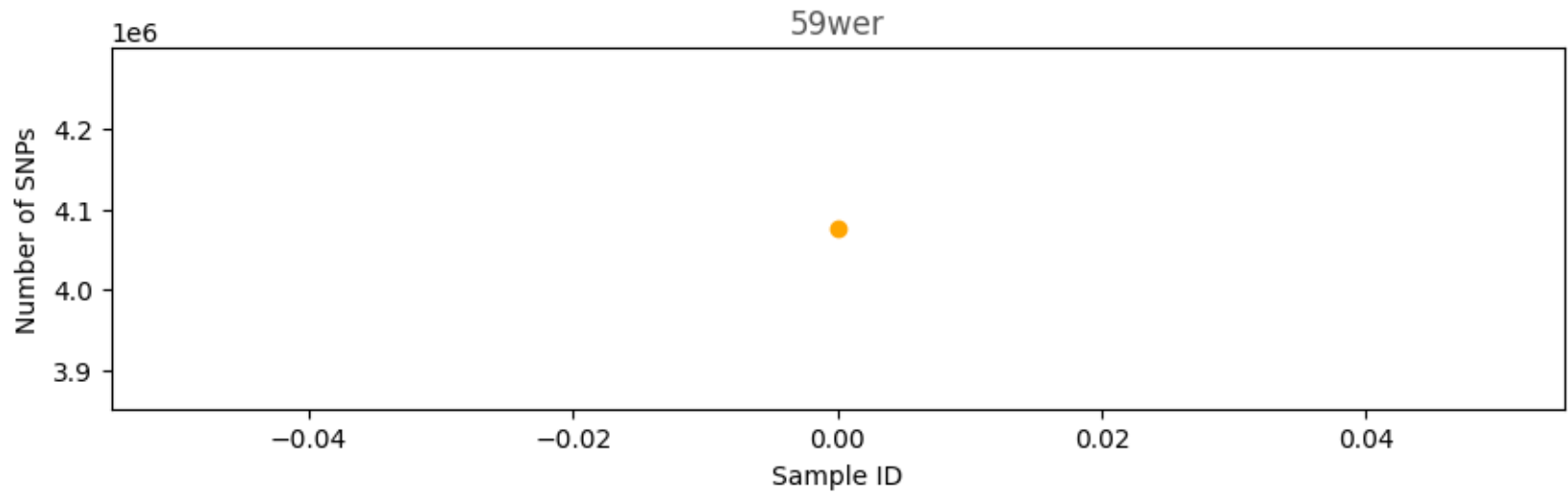
# Singletons by sample (hets and homs)



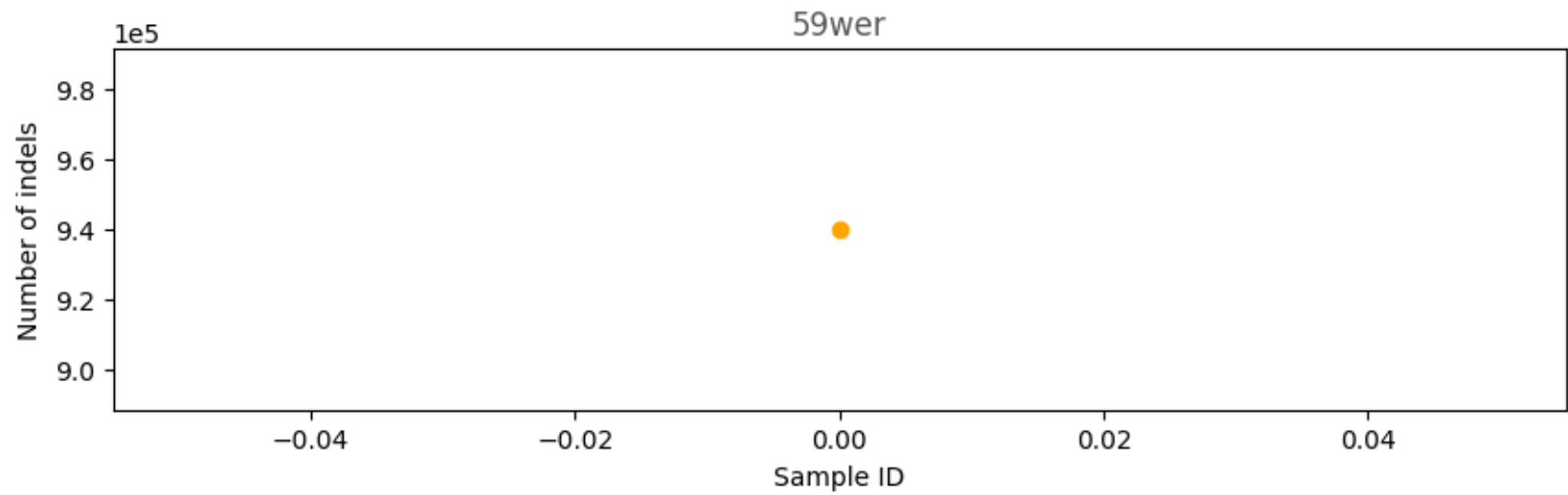
# Average depth by sample



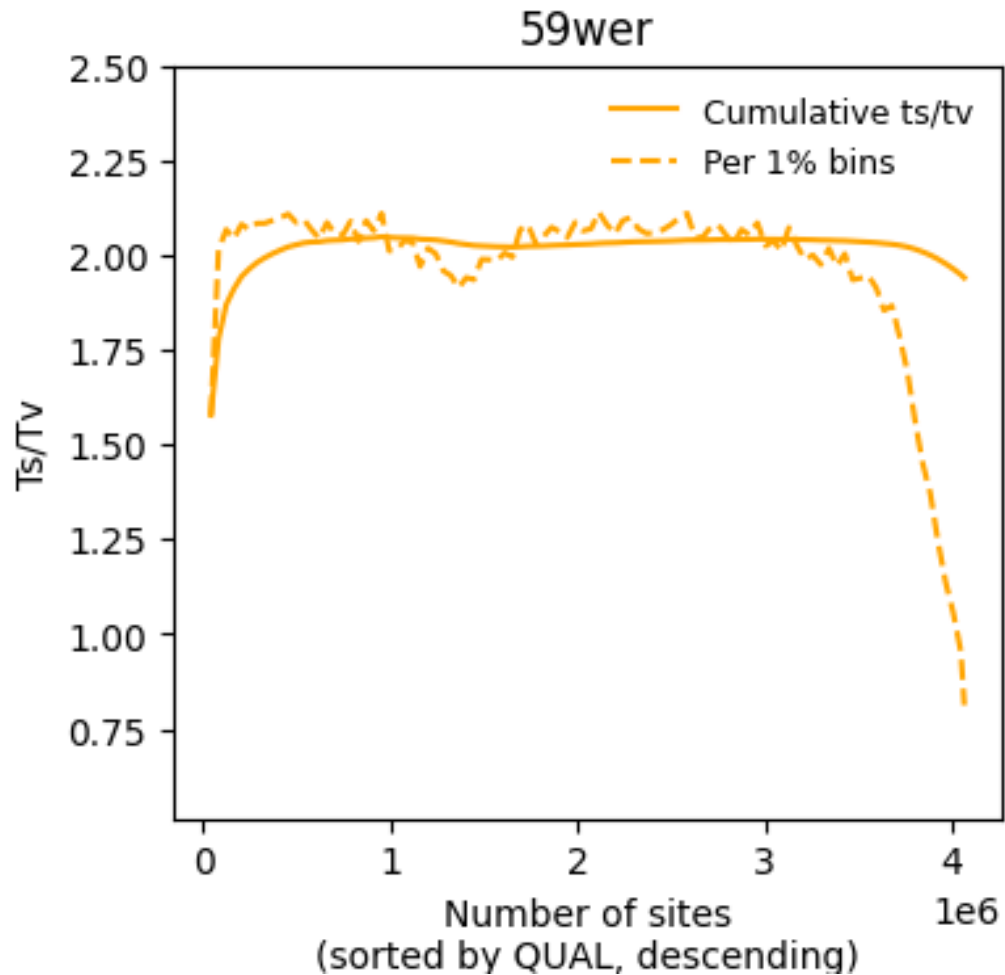
# Number of SNPs by sample



# Number of indels by sample

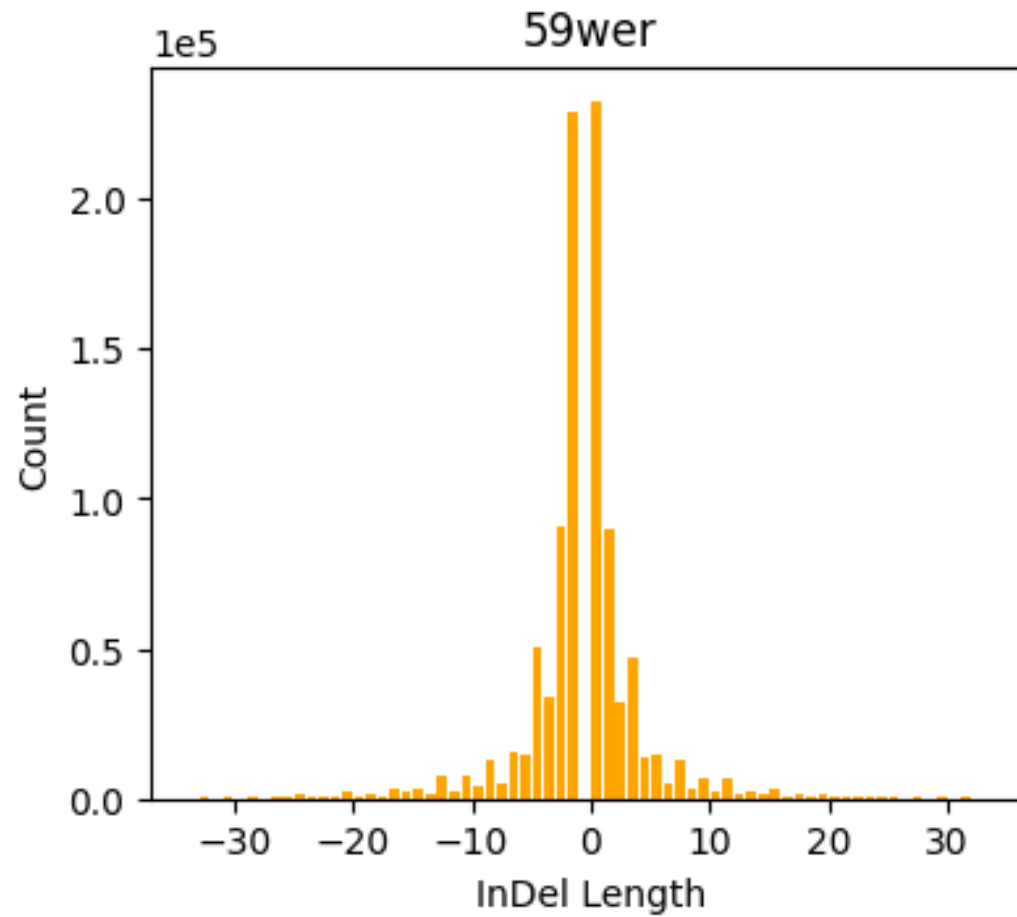


# Ts/Tv stratified by QUAL

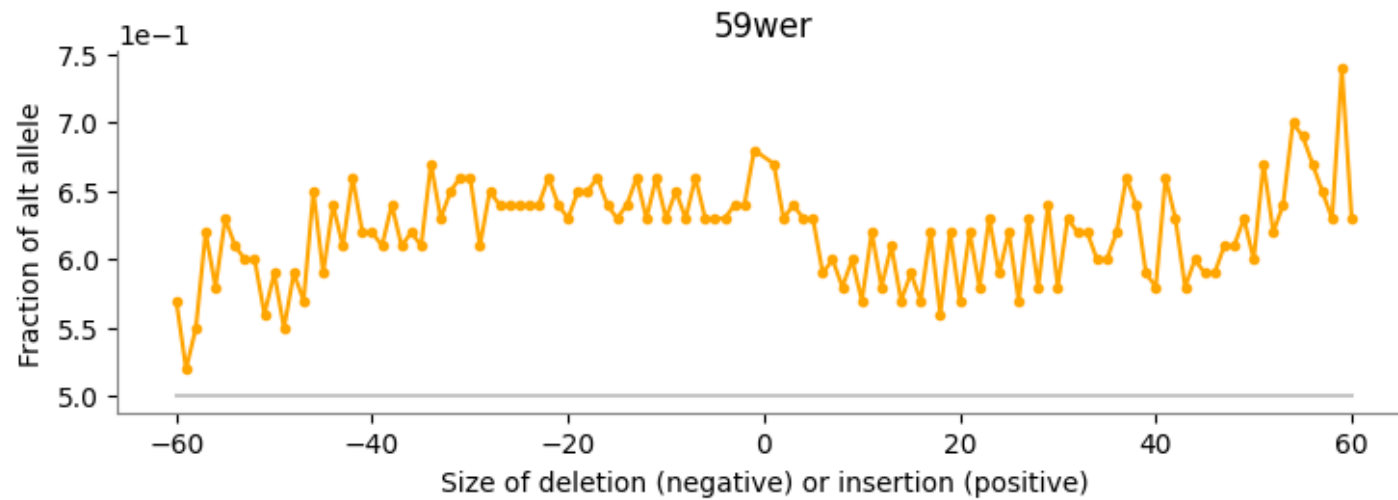




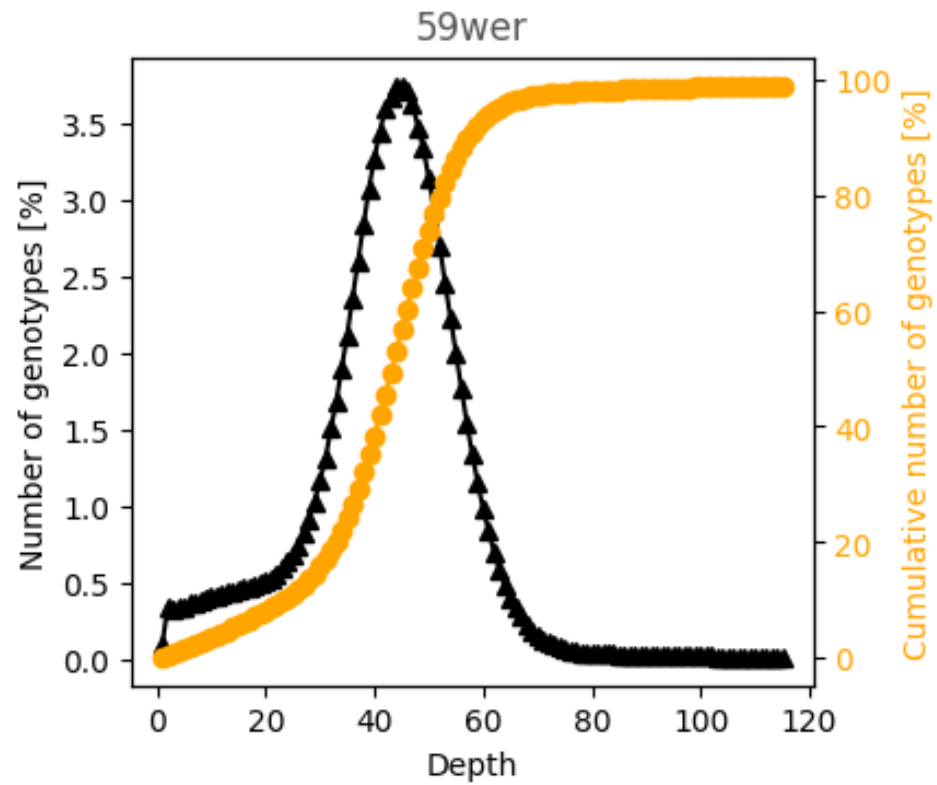
# Indel distribution



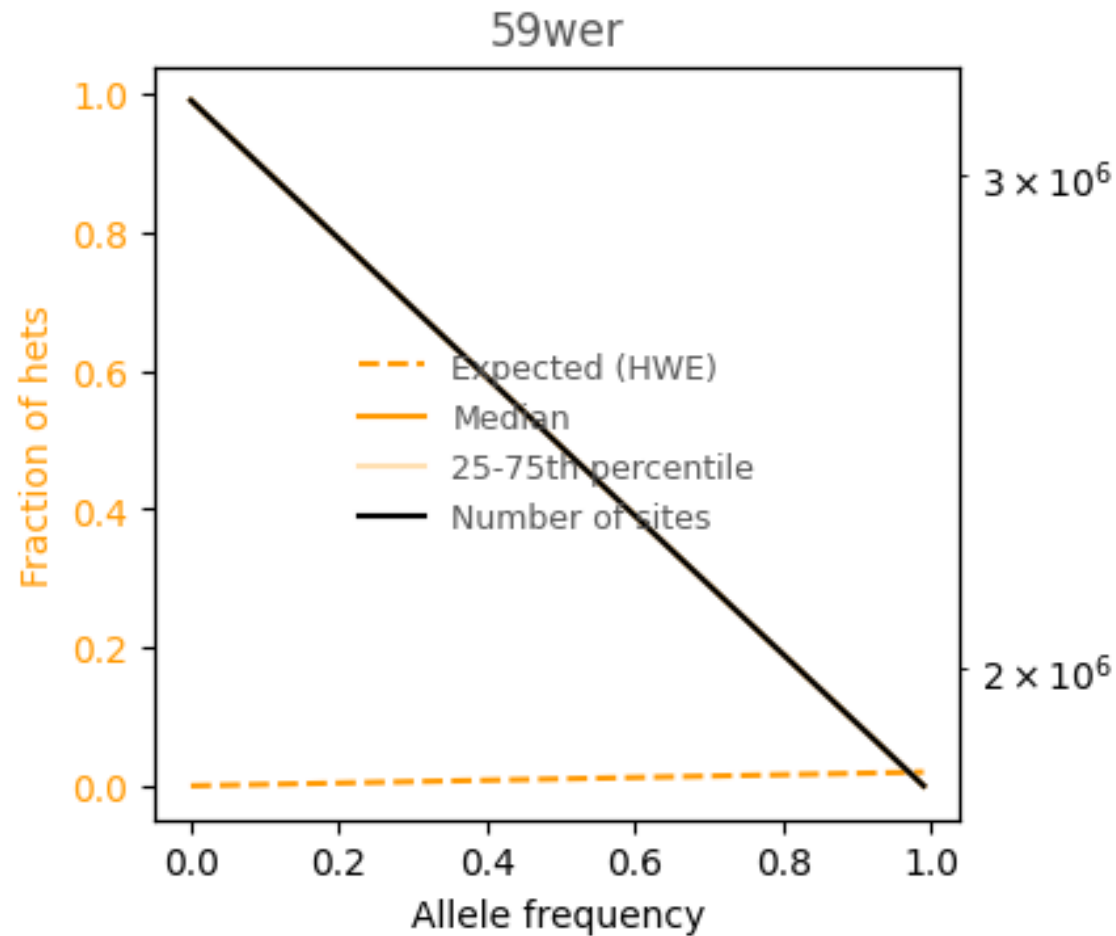
# Fraction of alternate indel allele



# Depth distribution



# Number of HETs by AF



# Substitution types

