

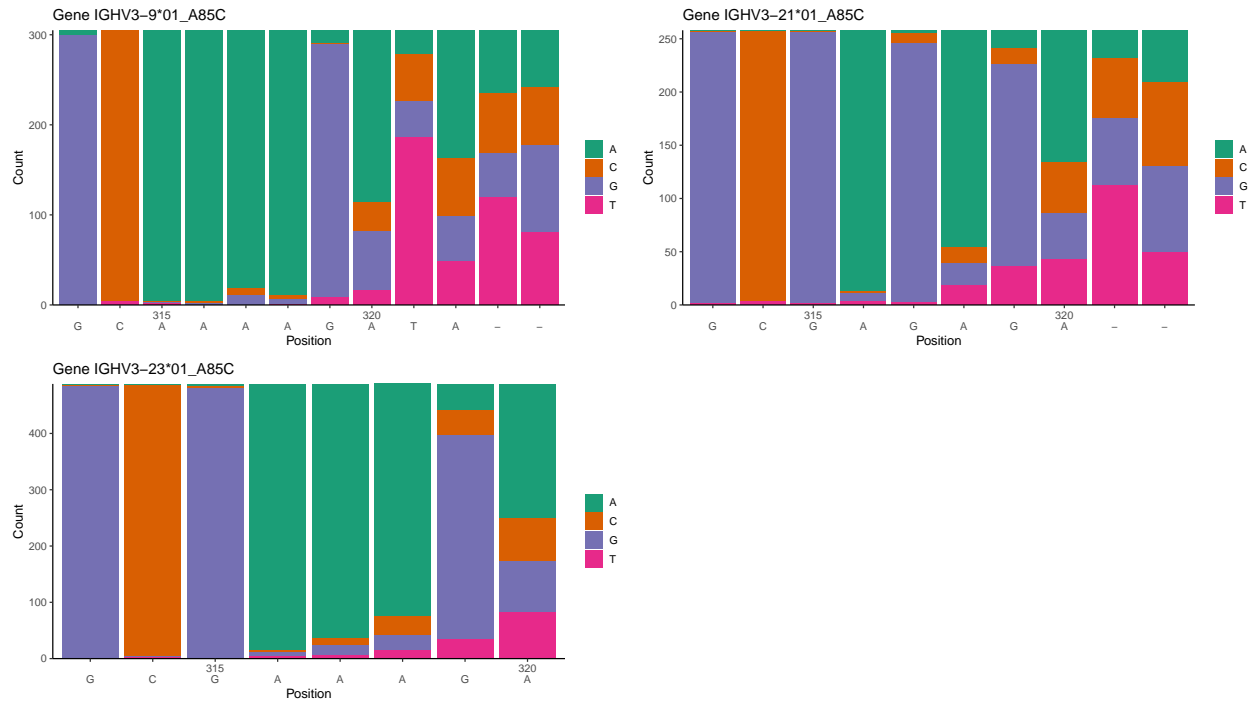
OGRDBstats Report

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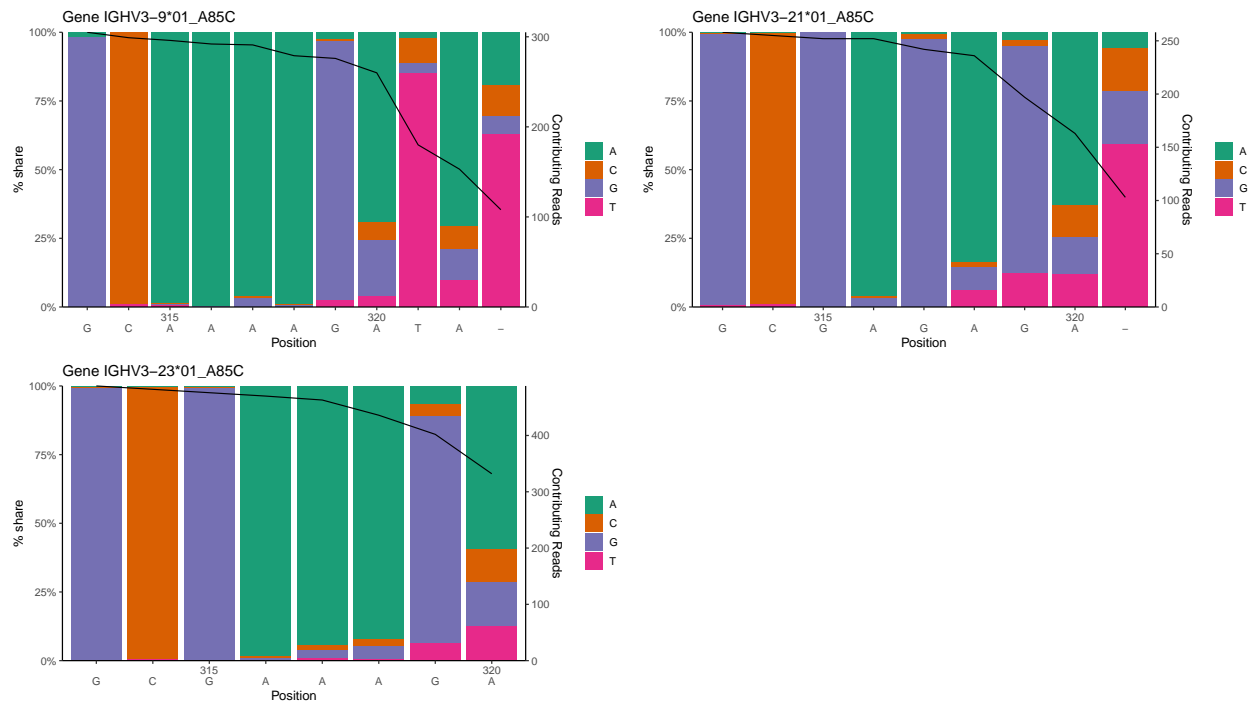
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1 Novel sequence analysis

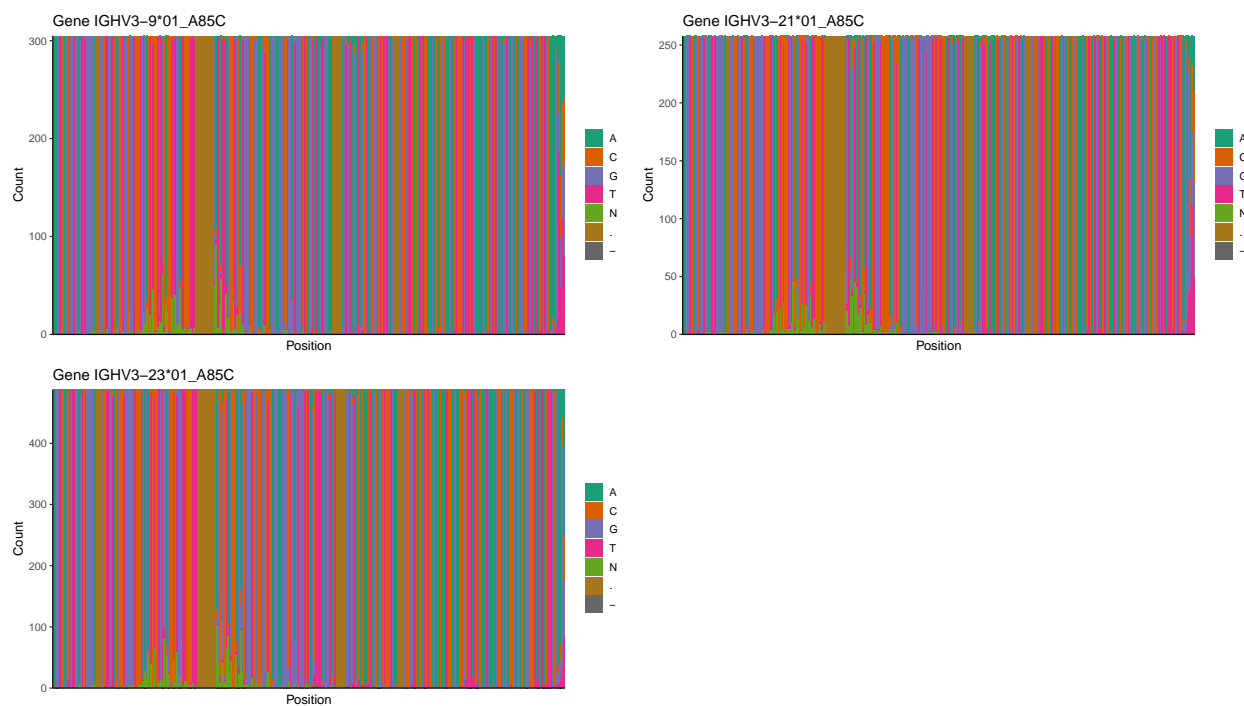
1.1 End-nucleotide composition



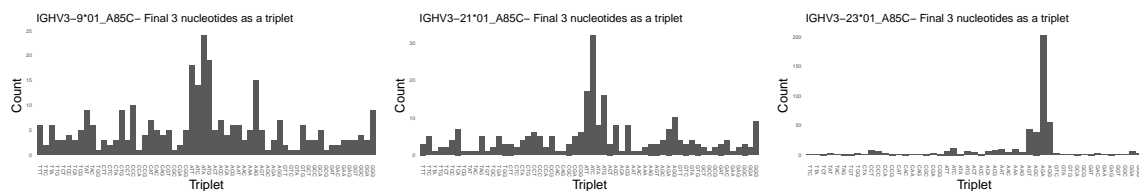
1.2 Per-nucleotide consensus where previous nucleotides match the consensus



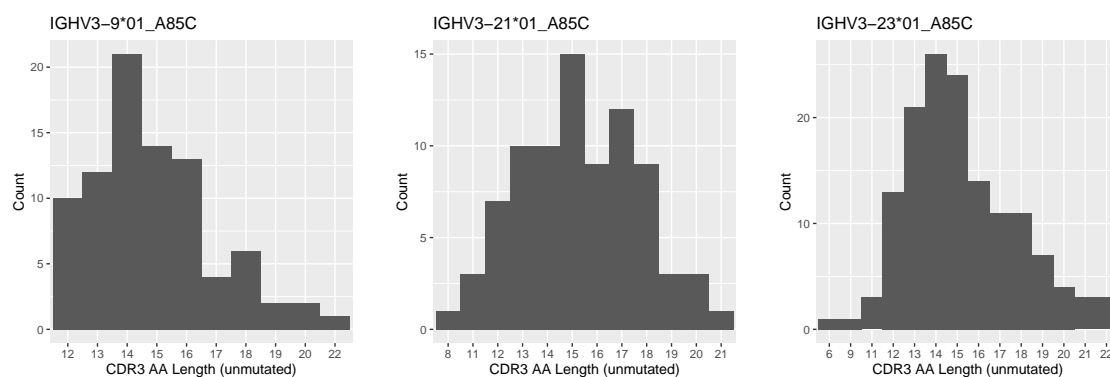
1.3 Whole-sequence composition of each assigned read



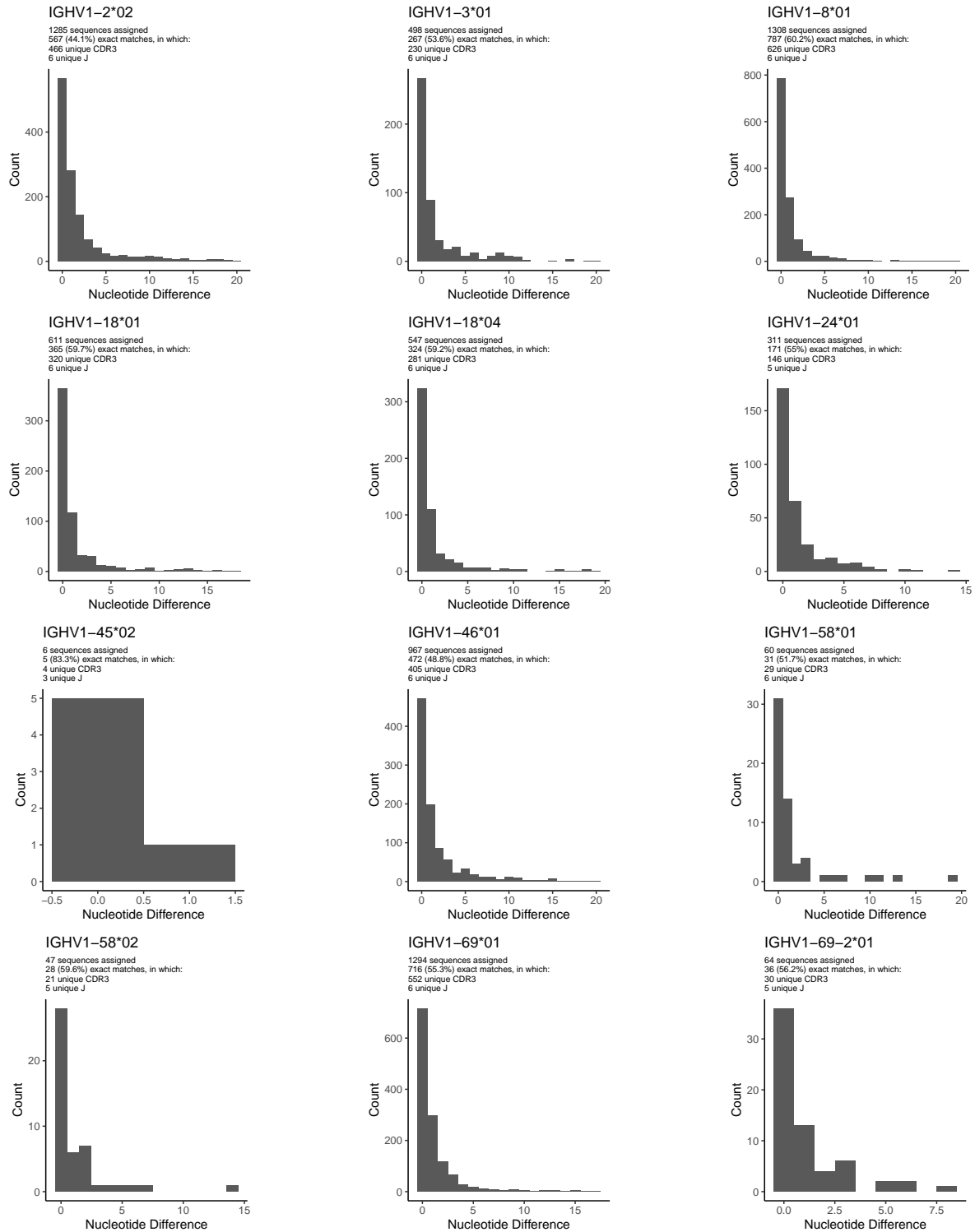
1.4 Final three nucleotides: frequency of each observed triplet

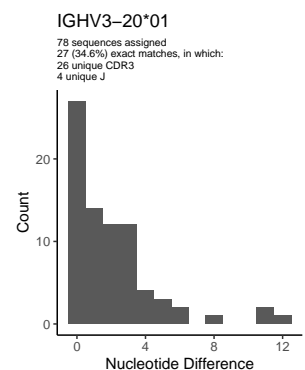
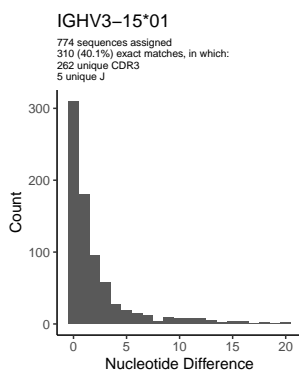
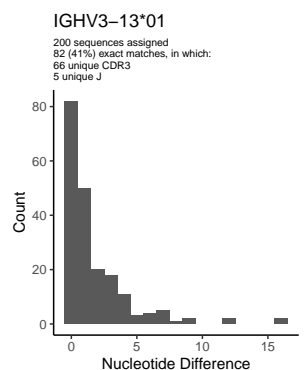
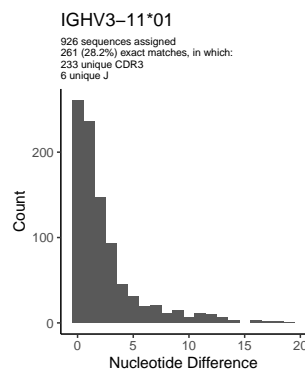
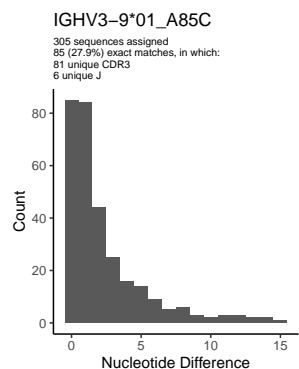
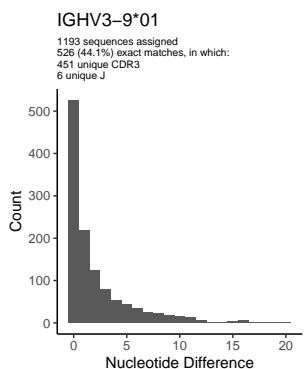
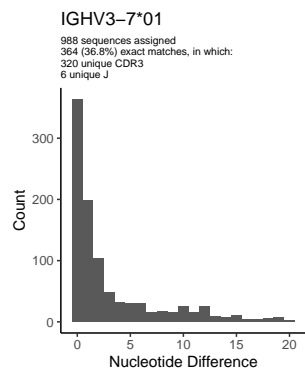
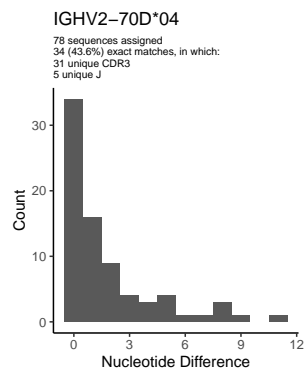
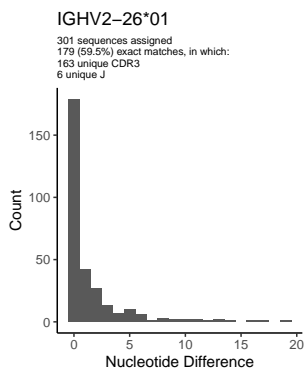
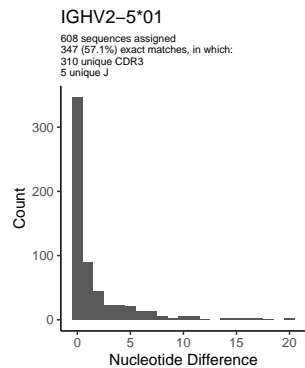
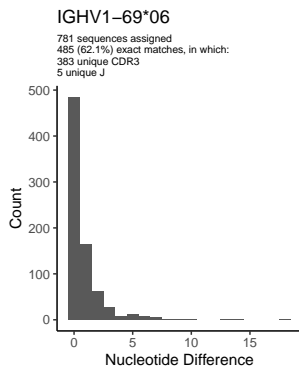
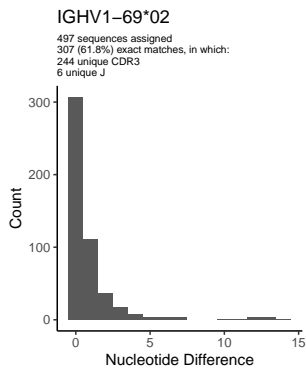


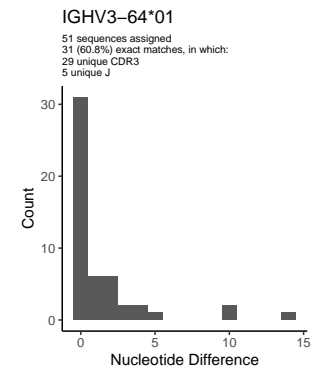
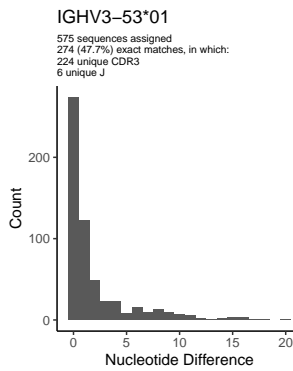
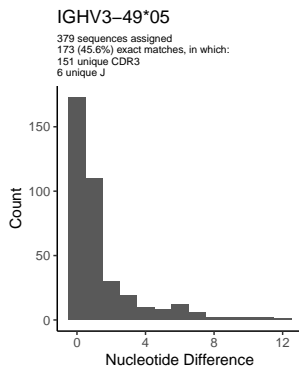
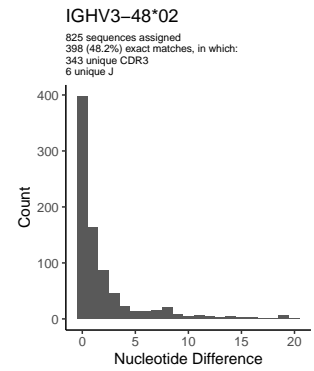
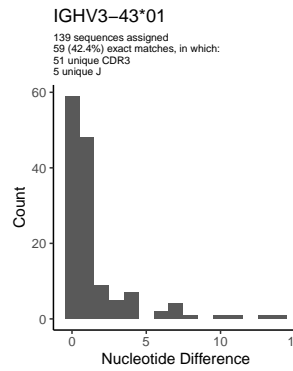
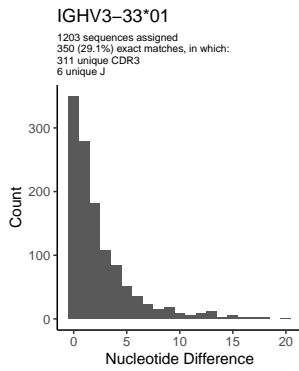
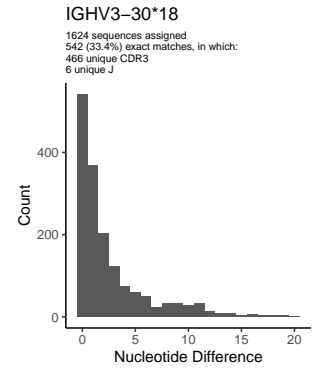
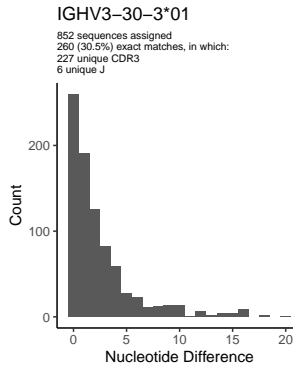
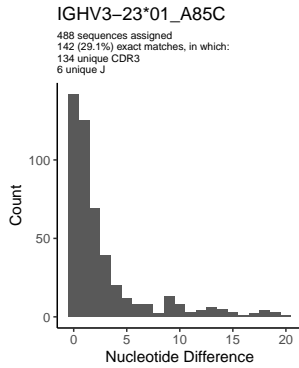
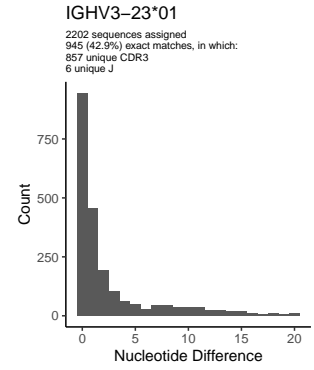
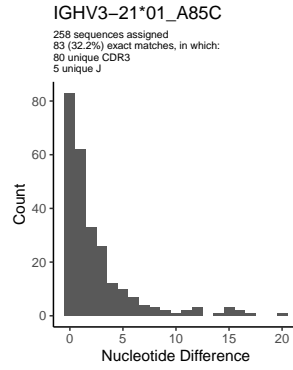
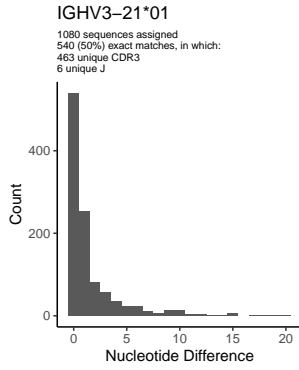
1.5 CDR3 length distribution, in assignments to novel alleles

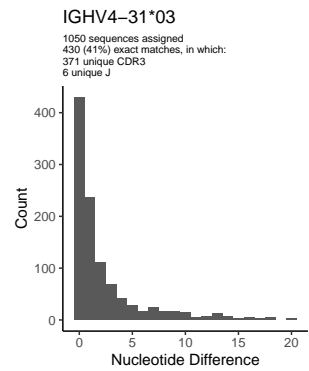
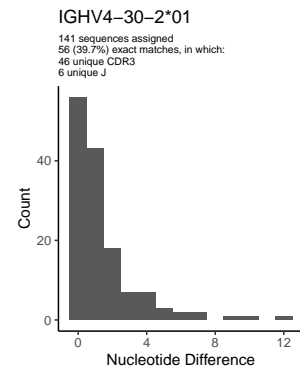
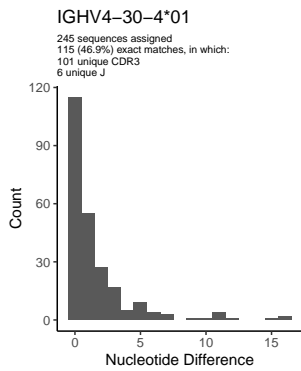
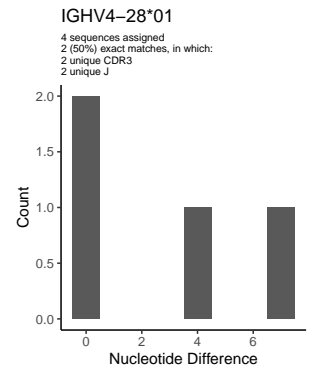
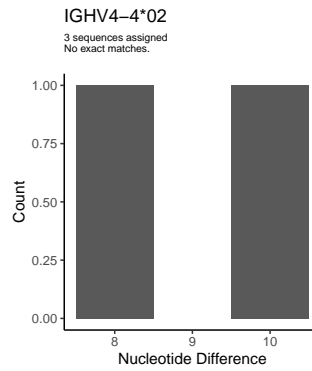
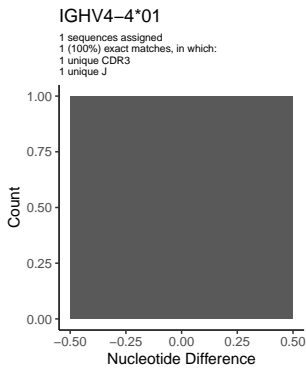
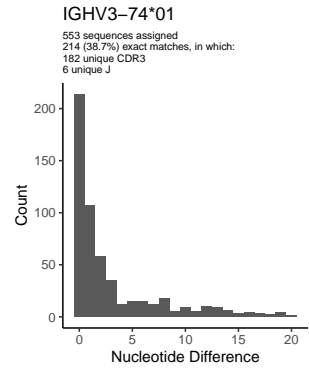
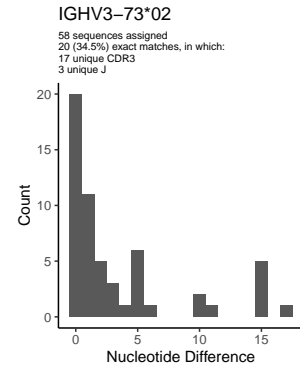
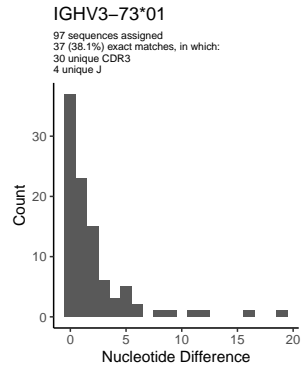
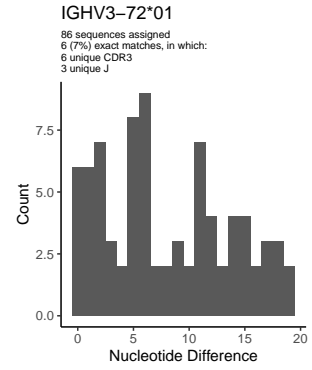
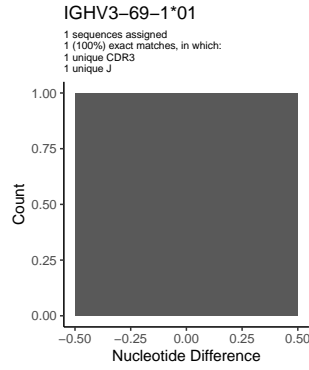
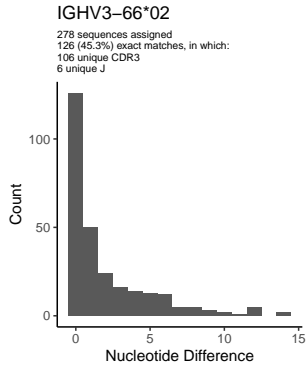


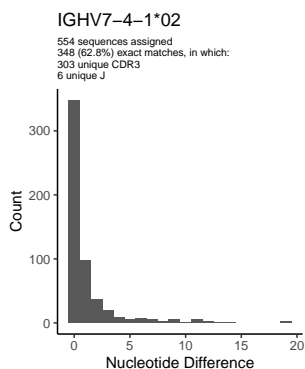
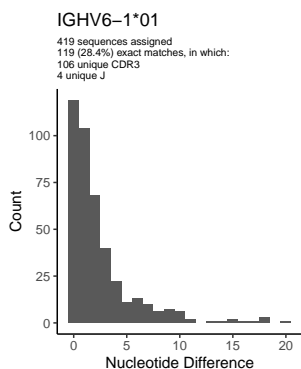
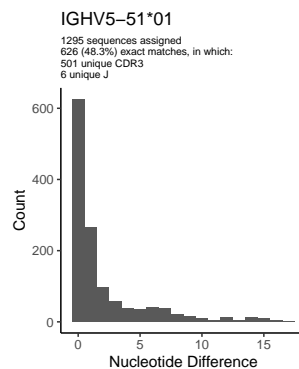
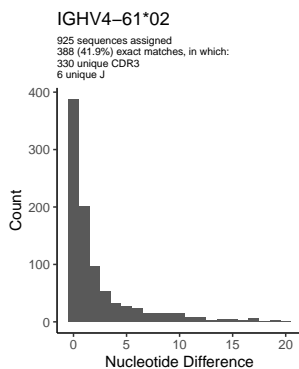
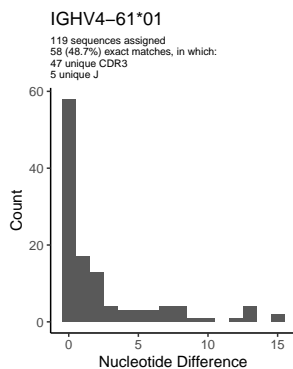
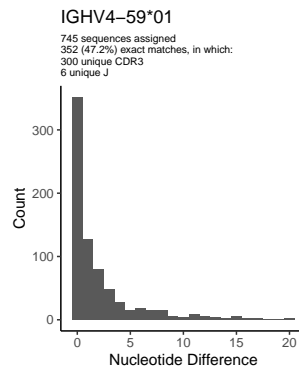
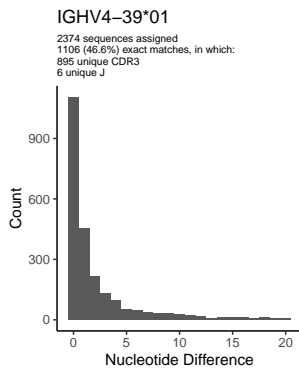
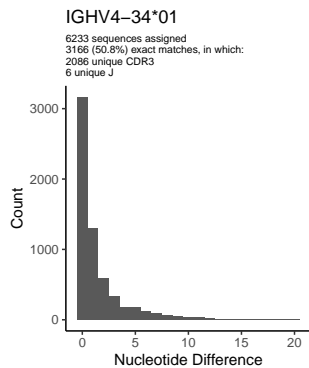
2 Variation from germline, in assignments to each allele



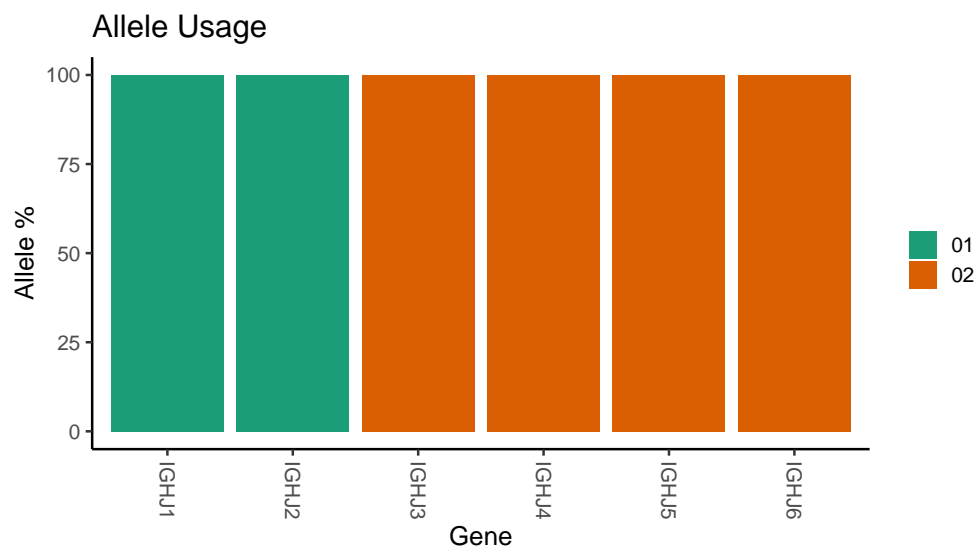








3 Allele usage in potential haplotype anchor genes



4 Haplotype plots

5 Configuration settings

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##  
## Germline reference file: /work/jenkins/PRJEB26509/S3/S3/3/3/c5/3e70dbc1464751fb42edc164455501/persona  
##  
## Novel allele file: /work/jenkins/PRJEB26509/S3/S3/3/3/c5/3e70dbc1464751fb42edc164455501/novel_sequen  
##  
## Species: Homosapiens  
##  
## Chain: IGHV  
##  
## Segment: V  
##  
## warning: non codon aligned gaps were found in novel sequence IGHV3 9 01_A85C  
##  
##  
## warning: non codon aligned gaps were found in novel sequence IGHV3 21 01_A85C
```