

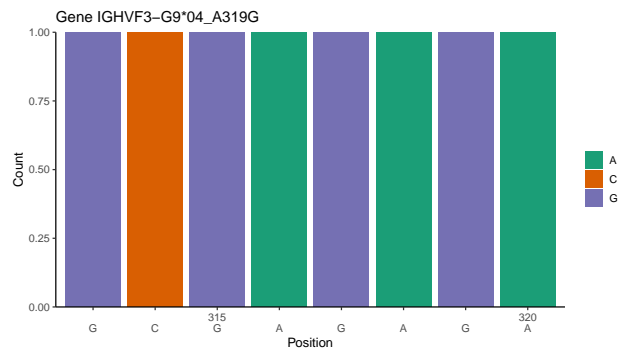
OGRDBstats Report

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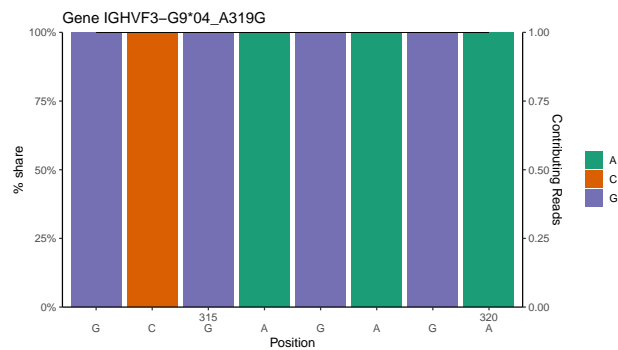
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1 Novel sequence analysis

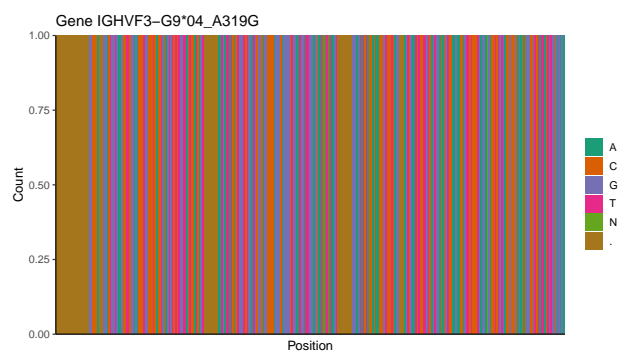
1.1 End-nucleotide composition



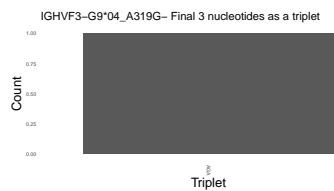
1.2 Per-nucleotide consensus where previous nucleotides match the consensus



1.3 Whole-sequence composition of each assigned read

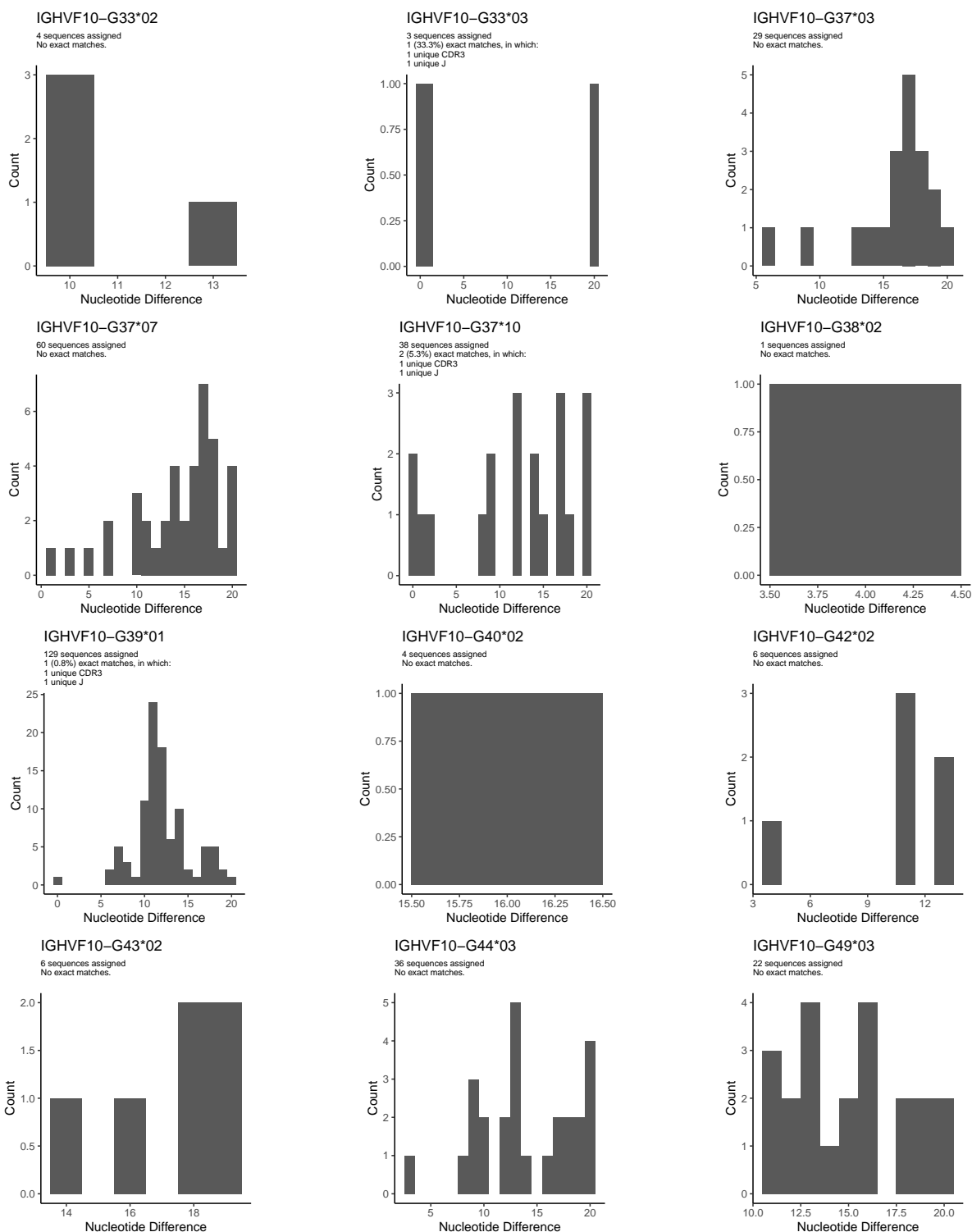


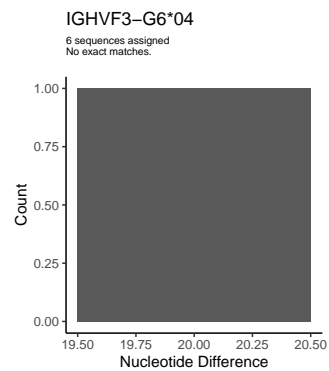
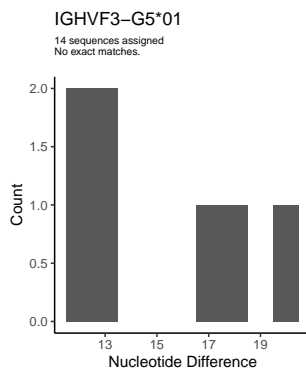
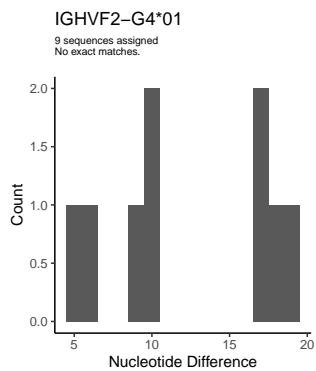
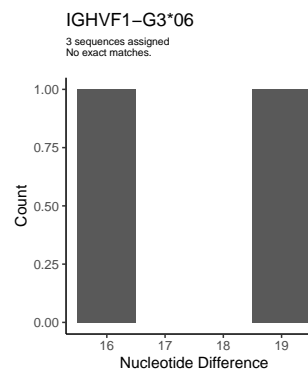
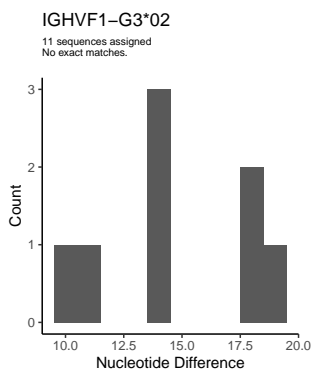
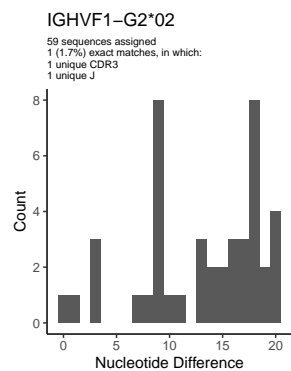
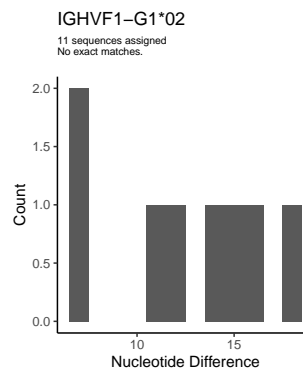
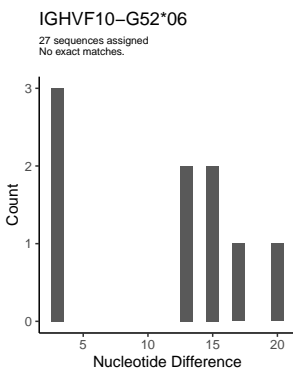
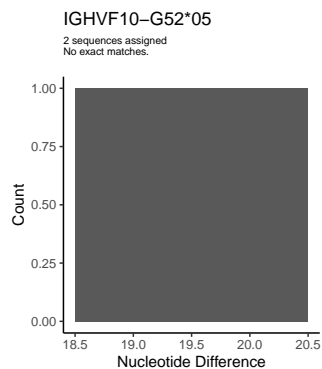
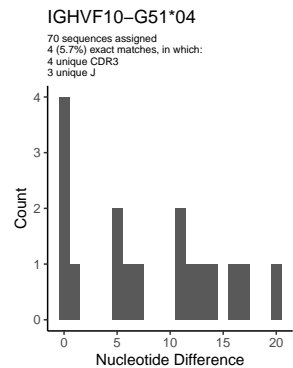
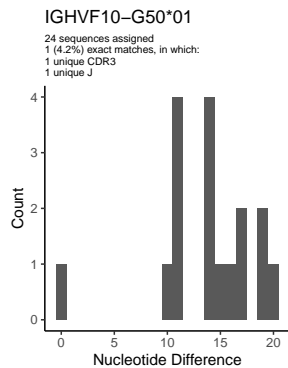
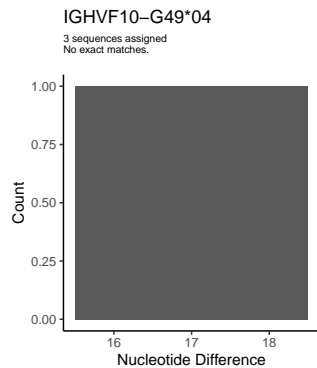
1.4 Final three nucleotides: frequency of each observed triplet

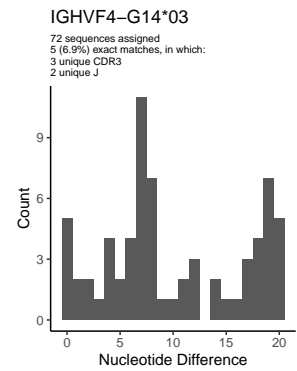
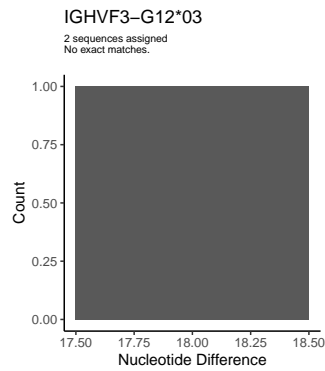
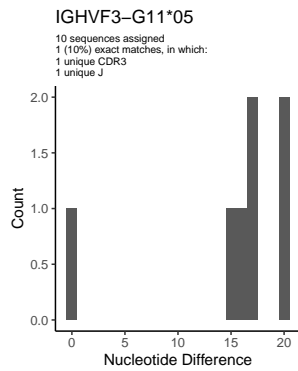
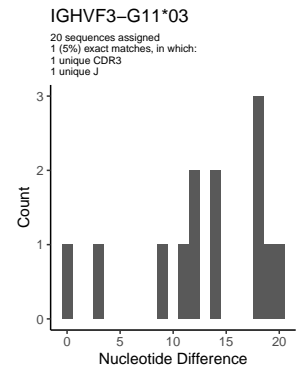
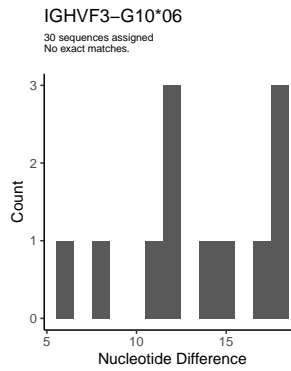
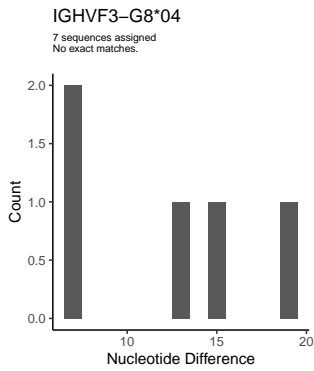
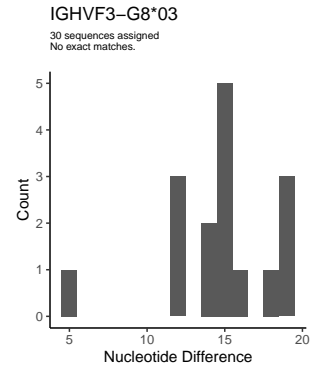
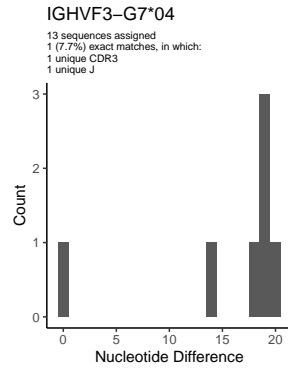
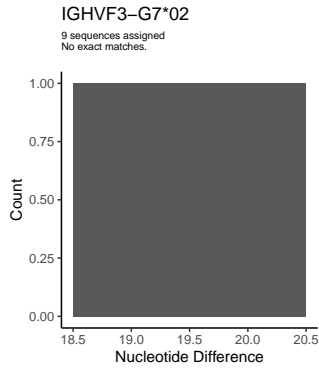
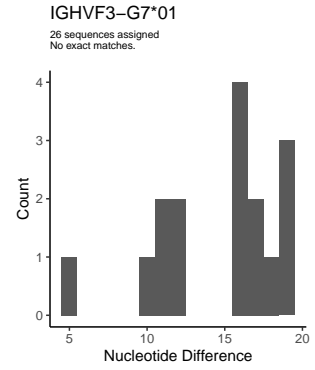
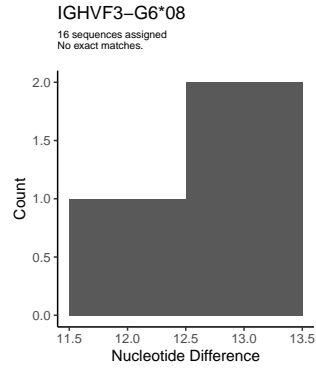
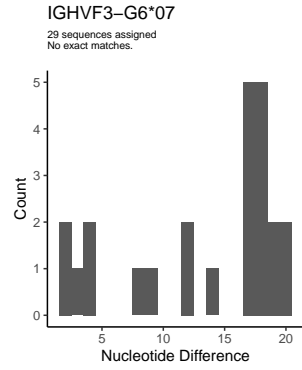


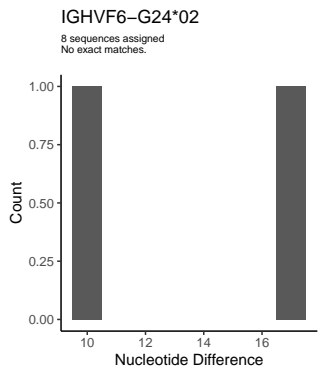
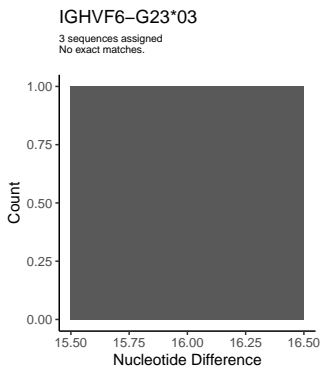
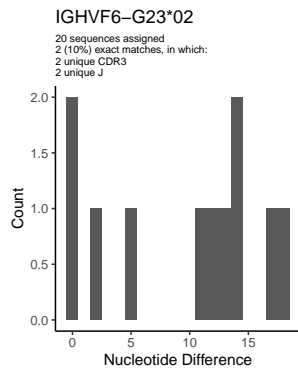
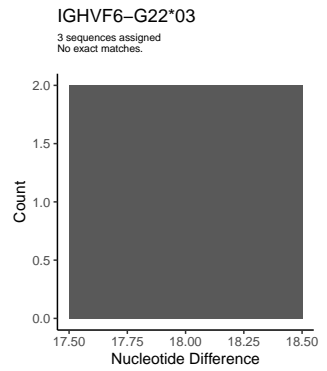
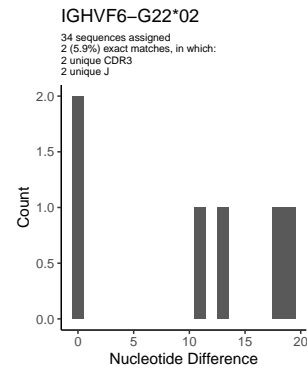
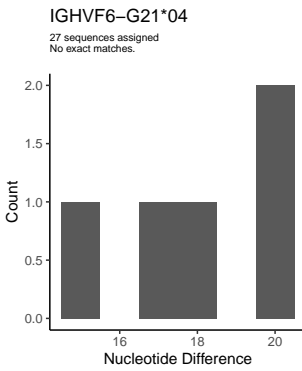
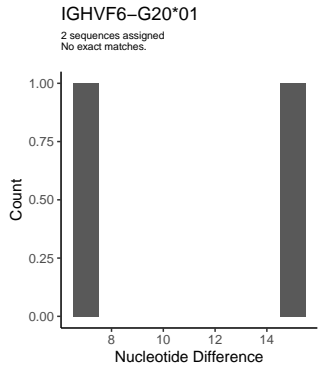
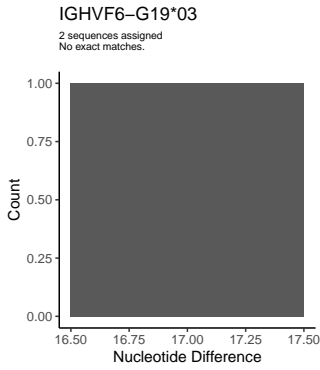
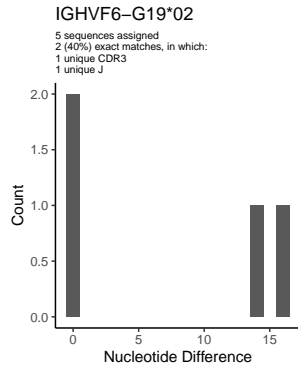
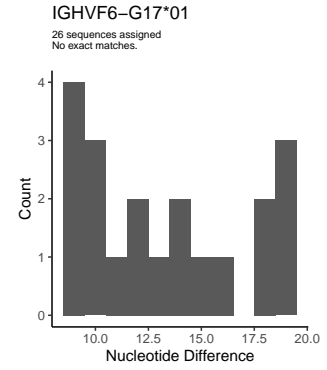
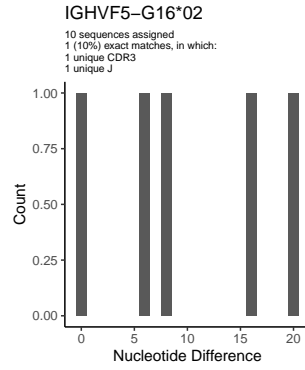
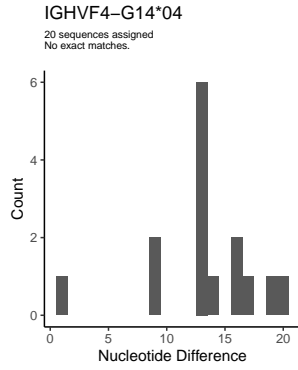
1.5 CDR3 length distribution, in assignments to novel alleles

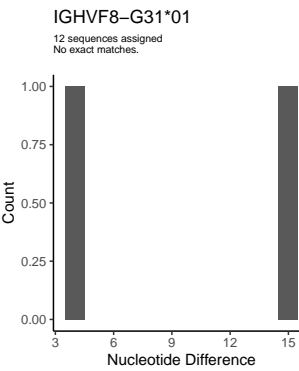
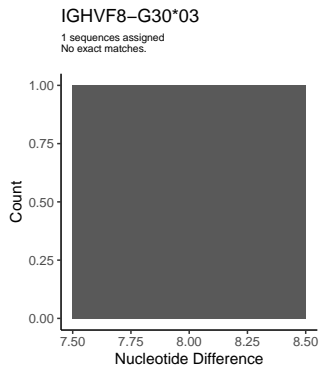
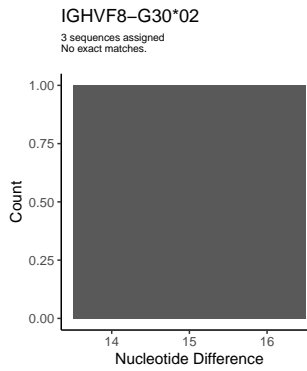
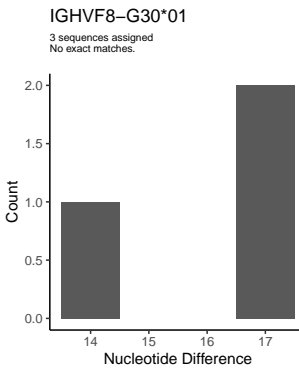
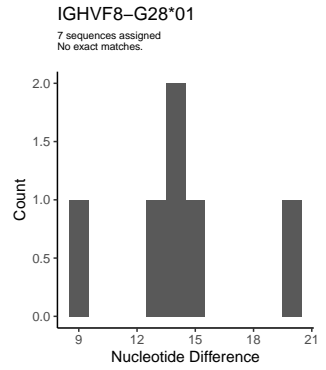
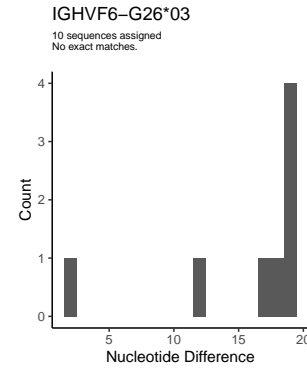
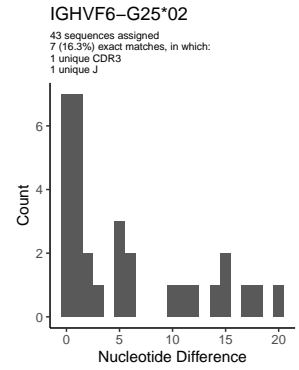
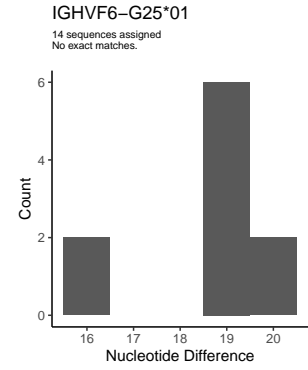
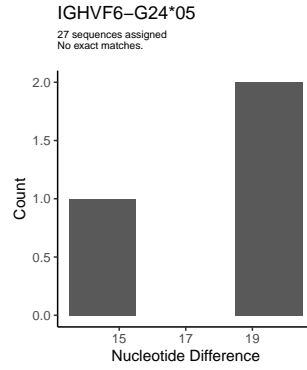
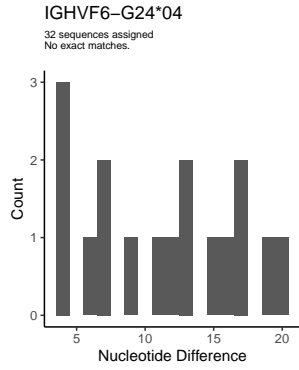
2 Variation from germline, in assignments to each allele











3 Allele usage in potential haplotype anchor genes



4 Haplotype plots

5 Configuration settings

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##
## Germline reference file: /misc/work/jenkins/PRJNA248475/M2/M2_lesion/M2_lesion/M2_lesion/d9/b063d9b2
##
## Novel allele file: /misc/work/jenkins/PRJNA248475/M2/M2_lesion/M2_lesion/M2_lesion/d9/b063d9b25adff8
##
## Species: Homosapiens
##
## Chain: IGHV
##
## Segment: V
```