The top CNV component has the followinf trio-states. Where '1' indicates a deletion and order is F, M, O.

```
000 001 010 011 100 101 110 111
326 3 14 35 14 32 1 20
```

## Methods

## Cleft Data Description

- Performed 284 tests. Bonferroni significant locus has width 4.7 kB.
- PennCNV joint HMM
- european, MAD < 0.3, non-WGA, aux  $\neq 1$
- coverage > 10
- 13140 hemi/homozygous deletions identified in 445 trios
- 4288 CNV components

```
Common (> 0.01): 954
Rare: 3334
```

• Construct trio-states for all CNV components

recall that we use indicator variable for hemi/homozygous deletions

ullet must be at least 5 informative mating pairs

01x and 10x

• count transmissions and perform binom.test (See "trans.tab")

GRanges with 284 ranges and 0 metadata columns:

```
      seqnames
      ranges
      strand

      <Rle>
      <IRanges>
      <Rle>

      comp120
      chr1
      [103969301, 103977611]
      *

      comp121
      chr1
      [103977612, 103983731]
      *

      comp122
      chr1
      [103983732, 103984810]
      *

      comp123
      chr1
      [103984811, 103988830]
      *

      comp124
      chr1
      [103988831, 103989217]
      *
```

```
. . .
                . . .
  comp3942
                       [20418019, 20422199]
              chr19
  comp3943
              chr19
                       [20422200, 20493452]
                       [20493453, 20507068]
  comp3944
              chr19
  comp4116
              chr20
                       [52081230, 52081774]
              chr20
                       [52081775, 52088118]
  comp4117
  seqlengths:
                   chr1 random
                                         chr2 ...
            chr1
                                                            chrY
                                                                           chrM
                                                        57772954
       247249719
                       1663265
                                    242951149 ...
                                                                          16571
> est.list.pitt <- lapply(binom.list.pitt, FUN = trioClasses:::get.est)
> ci.list.pitt <- lapply(binom.list.pitt, FUN = trioClasses:::get.ci)
> testable.pitt <- !is.na(ci.list.pitt)</pre>
> ci.mat.pitt <- matrix(unlist(ci.list.pitt[testable.pitt]),</pre>
     nrow = sum(testable.pitt), ncol = 2, byrow = TRUE)
> est.vec.pitt <- as(est.list.pitt, "numeric")[testable.pitt]
> cnv.pitt.obj$cmp.gr[testable.pitt]
GRanges with 354 ranges and 0 metadata columns:
           segnames
                                     ranges strand
              <Rle>
                                  <IRanges> <Rle>
   comp117
               chr1 [103977612, 103983731]
               chr1 [103983732, 103989830]
   comp118
               chr1 [103989831, 103990016]
   comp119
               chr1 [103990017, 103991756]
   comp120
   comp121
               chr1 [103991757, 103999836]
  comp3774
                       [52081775, 52088118]
              chr20
                       [14163409, 14164422]
  comp3831
              chr21
  comp3832
              chr21
                       [14164423, 14166029]
                       [14166030, 14166821]
  comp3833
              chr21
              chr21
                       [14166822, 14169701]
  comp3834
  seqlengths:
            chr1
                   chr1_random
                                         chr2 ...
                                                            chrY
                                                                           chrM
```

242951149 ...

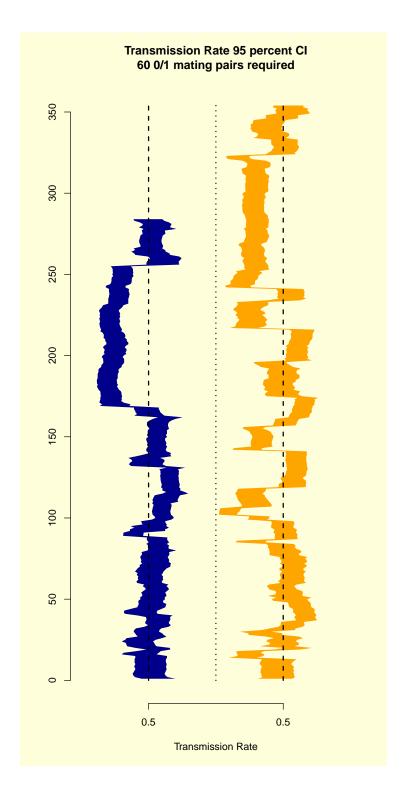
57772954

16571

The CNV components with significant p-values (Bonferroni) are given below.

1663265

247249719



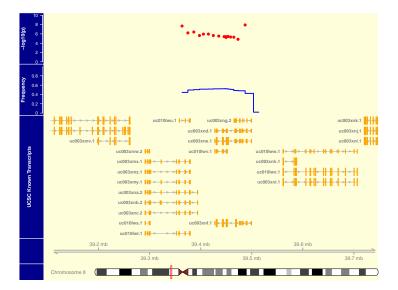


Figure 1: Cleft Trios — Red:  $-\log_{10} p$ -values for each CNV component that has at least **five** "0/1" mating pairs. Null hypothesis is transmission rate of  $\frac{1}{2}$ , and alternative hypothesis is "one-sided." Blue: The frequency of CNV component in the parents. Yellow: Gene tracks, labeled by transcript.

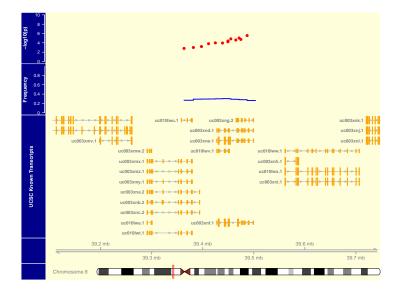


Figure 2: Control Trios — Red:  $-\log_{10} p$ -values for each CNV component that has at least **five** "0/1" mating pairs. Null hypothesis is transmission rate of  $\frac{1}{2}$ , and alternative hypothesis is "one-sided." Blue: The frequency of CNV component in the parents. Yellow: Gene tracks, labeled by transcript.



Figure 3: UCSC Genome Browser at chromosome 8 locus.

```
GRanges with 17 ranges and 0 metadata columns:
           seqnames
```

```
ranges strand
            <Rle>
                                <IRanges>
                                            <Rle>
comp1883
             chr7 [141435189, 141439888]
comp1994
             chr8 [ 39356825,
                                39370186]
comp1995
             chr8 [ 39370187,
                                39379683]
comp1996
             chr8 [ 39379684,
                                39393417]
comp1997
             chr8 [ 39393418,
                                39401865]
              . . .
comp2005
             chr8
                     [39450168, 39457081]
comp2006
             chr8
                     [39457082, 39460087]
                     [39460088, 39469612]
comp2007
             chr8
comp2008
             chr8
                     [39469613, 39476658]
comp2009
             chr8
                     [39476659, 39497557]
seqlengths:
                 chr1_random
                                        chr2 ...
          chr1
     247249719
                      1663265
                                  242951149 ...
                                                      57772954
```

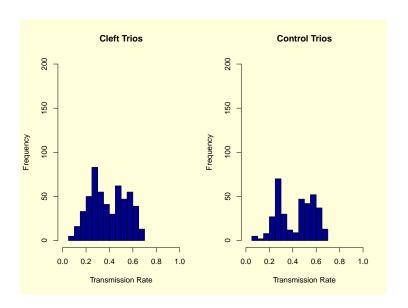
All CNV components are contiguous and the total width is 4.7 kB.

```
function (object)
{
    if (sum(object[c("101", "011", "100", "010")], na.rm = TRUE) >=
        60) {
        T \leftarrow sum(object[c("101", "011")], na.rm = TRUE)
        U <- sum(object[c("100", "010")], na.rm = TRUE)</pre>
        return(binom.test(x = T, n = T + U, p = 0.5, alternative = "greater")$p.value)
    }
    else {
        return(NA)
    }
}
```

chrY

chrM

16571



```
<environment: namespace:trioClasses>
> gr.deletion.pitt <- gr.pitt[values(gr.pitt)$numsnp >= 10 &
     values(gr.pitt)$cn %in% 0:1]
> sum(countOverlaps(gr.deletion.pitt, reduce(locus)))
[1] 649
> mean(est.vec[subjectHits(findOverlaps(reduce(locus), cnv.beaty.obj$cmp.gr))])
[1] NA
> mean(est.vec.pitt[subjectHits(findOverlaps(reduce(locus),
     cnv.pitt.obj$cmp.gr))])
[1] NA
> mean(-log10(trans.vec[subjectHits(findOverlaps(reduce(locus),
     cnv.beaty.obj$cmp.gr))]))
[1] 5.784253
> mean(-log10(trans.vec.pitt[subjectHits(findOverlaps(reduce(locus),
     cnv.pitt.obj$cmp.gr))]))
[1] 3.907709
```

> freq.pitt.vec[subjectHits(findOverlaps(reduce(locus), cnv.pitt.obj\$cmp.gr))]

```
      comp1670
      comp1820
      comp1821
      comp1822
      comp1823
      comp1824
      comp1825

      0.1914062
      0.2675781
      0.2910156
      0.2929688
      0.2949219
      0.2988281
      0.3027344

      comp1826
      comp1827
      comp1828
      comp1829
      comp1830
      comp1831
      comp1832

      0.3027344
      0.3007812
      0.2890625
      0.2871094
      0.2792969
      0.2792969
      0.2617188
```

> freq.vec[subjectHits(findOverlaps(reduce(locus), cnv.beaty.obj\$cmp.gr))]

 comp1883
 comp1994
 comp1995
 comp1996
 comp1997
 comp1998
 comp1999

 0.1534155
 0.4445689
 0.4960806
 0.5039194
 0.5139978
 0.5139978
 0.5162374

 comp2000
 comp2001
 comp2002
 comp2003
 comp2004
 comp2005
 comp2006

 0.5207167
 0.5218365
 0.5229563
 0.5251960
 0.5218365
 0.5139978
 0.5083987

 comp2007
 comp2008
 comp2009

 0.4848824
 0.4792833
 0.4232923