

# PepBed examples

November 18, 2017

importing bigbed files

```
library(PepBed)

# path to bigbed file
bigbedpath <- '/home/enrique/temp/pride_cluster_peptides_9606_Human.pogo.bb'

# convert bigbed to bed file (output bed file in the same directory)
bigBed2bed(inputFile = bigbedpath, compress = FALSE)

# getting basic information (output description file in the same directory)
getBigBedInfo(inputFile = bigbedpath)

# getting field names if available
fieldNames <- getBigBedFieldNames(inputFile = bigbedpath, only.names = TRUE)

print(fieldNames)

## [1] "chrom"      "chromStart" "chromEnd"    "name"        "score"
## [6] "strand"     "thickStart"  "thickEnd"    "reserved"     "blockCount"
## [11] "blockSizes" "chromStarts"
```

parsing Bed file

```
# path to bed file
bedpath <- '/home/enrique/temp/pride_cluster_peptides_9606_Human.bed'

# import bed file as dataframe
df <- readBedFile(inputFile = bedpath)

# set column name to bed file
names(df) <- fieldNames

# convert dataframe to GRanges
granges_peptide <- buildGRangesFromData(data = df, chrColName = "chrom",
                                          startColName = "chromStart",
                                          endColName = "chromEnd")
```

some basic stats

```
# getting number of features(peptides) by chromosome  
counts <- countsByChromosome(gr = granges_peptide)  
print(counts)
```

##	chromosome	countFeatures
## 1	chr1	9736
## 2	chr2	3688
## 3	chr3	5991
## 4	chr4	6191
## 5	chr5	1183
## 6	chr6	3822
## 7	chr7	2704
## 8	chr8	3524
## 9	chr9	6678
## 10	chr10	1176
## 11	chr11	5536
## 12	chr12	7490
## 13	chr13	2264
## 14	chr14	867
## 15	chr15	2236
## 16	chr16	5200
## 17	chr17	3505
## 18	chr18	4013
## 19	chr19	6463
## 20	chr20	4121
## 21	chr21	3064
## 22	chr22	3841
## 23	chrX	19
## 24	chrY	3138
## 25	chrM	131

```

# removing duplicated entries from original granges_peptide
gr_unique <- getUniqueFeatures(granges_peptide, colFeatures = 'name')

# getting unique number of features(peptides) by chromosome
counts_unique <- countsByChromosome(gr = gr_unique)
print(counts_unique)

```

```

##      chromosome countFeatures
## 1         chr1         8801
## 2         chr2         3379
## 3         chr3         5341
## 4         chr4         4985
## 5         chr5         1031
## 6         chr6         3123
## 7         chr7         2336
## 8         chr8         2942
## 9         chr9         5491
## 10        chr10        1016
## 11        chr11        4666
## 12        chr12        6073
## 13        chr13        1935
## 14        chr14          666
## 15        chr15        1742
## 16        chr16        4622
## 17        chr17        3145
## 18        chr18        3385
## 19        chr19        3759
## 20        chr20        3338
## 21        chr21        2543
## 22        chr22        3224
## 23         chrX          19
## 24         chrY        2388
## 25         chrM          8

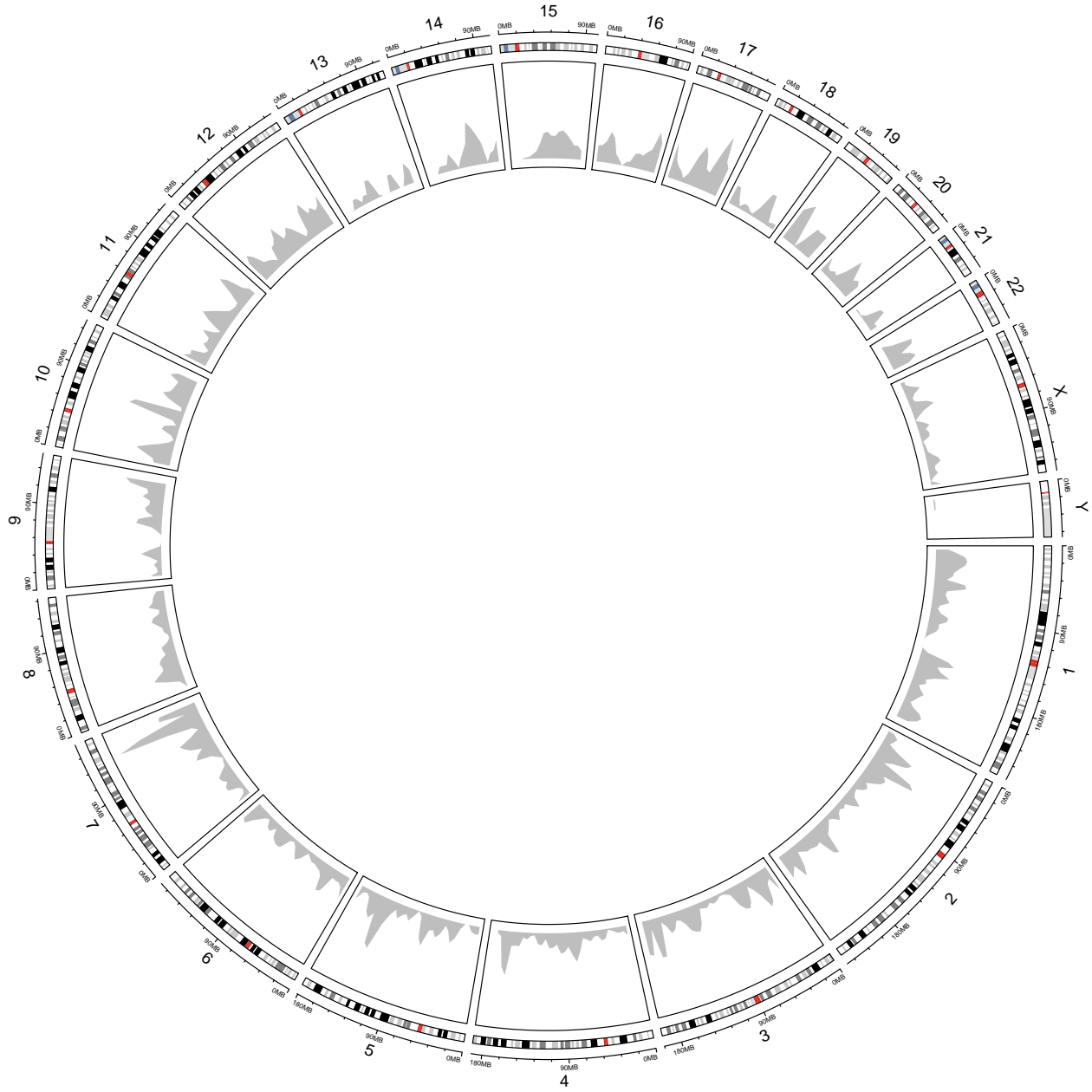
```

```
## compute coverage of query (peptide evidences) on subject (transcripts) by chromosome
data("protein_coding_transcript") # load protein coding transcript as GRanges object
coverage <- computeCoverageByChromosome(query = granges_peptide, subject = transcript)
print(coverage)
```

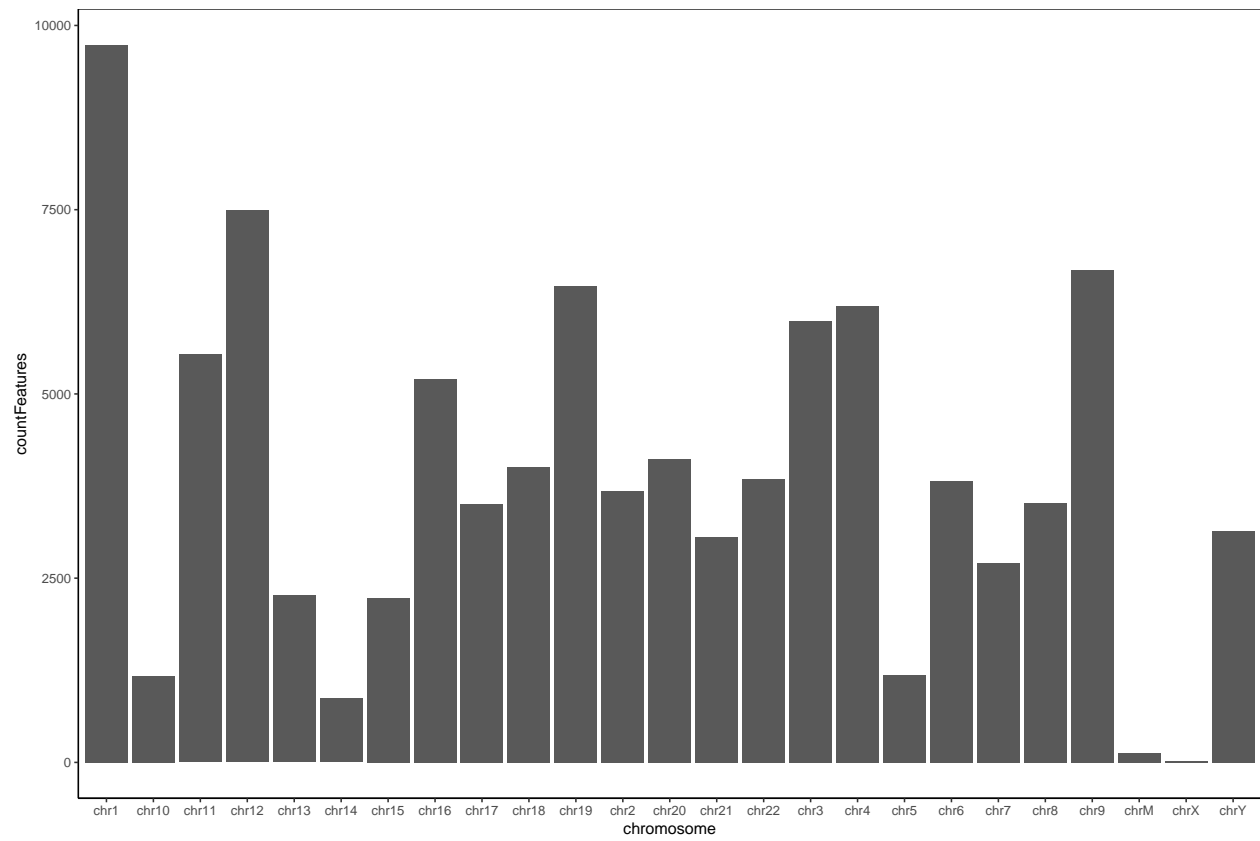
```
##      Chromosome Coverage
## 1         chr1    50.781
## 2         chr2    50.613
## 3         chr3    46.526
## 4         chr4    51.239
## 5         chr5    47.049
## 6         chr6    51.290
## 7         chr7    58.682
## 8         chr8    52.876
## 9         chr9    48.021
## 10        chr10   44.897
## 11        chr11   46.371
## 12        chr12   43.799
## 13        chr13   31.403
## 14        chr14   49.515
## 15        chr15   46.962
## 16        chr16   47.596
## 17        chr17   51.952
## 18        chr18   51.426
## 19        chr19   48.749
## 20        chr20   49.875
## 21        chr21   34.799
## 22        chr22   47.774
## 23         chrX   42.346
## 24         chrY   22.274
## 25         chrM    5.882
```

## plotting

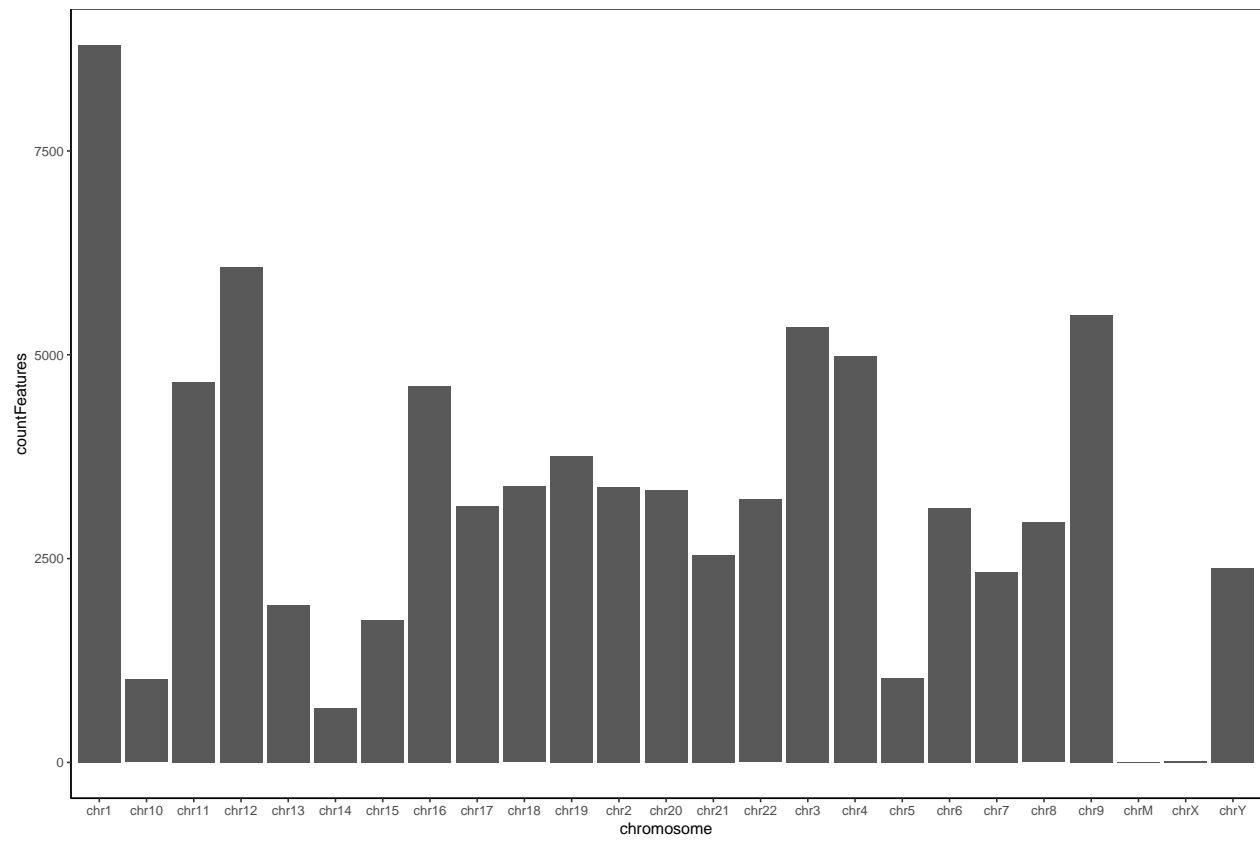
- The distribution of features (peptides) by chromosome



- barplot with counts by chromosome



- barplot with unique counts by chromosome



- barplot with coverage by chromosome

