

Compute coverage

```
bedtools_genomecov("-i a.bam -bga")
```

```
genome <- Seqinfo(genome = NA_character_)  
ga_a <- import("a.bam", genome = genome)  
cov <- coverage(granges(ga_a))  
cov_gr <- GRanges(cov)
```

Select zero runs

```
subset(score > 0)
```

Find intersection with regions

```
R_bedtools_intersect(cov_gr, "b.bed")
```

```
genome <- Seqinfo(genome = NA_character_)  
gr_a <- cov_gr  
gr_b <- import("b.bed", genome = genome)  
pairs <- findOverlapPairs(gr_a, gr_b,  
                           ignore.strand = TRUE)  
pintersect(pairs, ignore.strand = TRUE)
```