## exomePeak2

Here we use exomePeak2 to conduct reference based quantification and differential analysis.

## 1. Download and Convert Basic Site Information

Download single base RNA modification annotation of m6A on human genome from m6A-Atlas database, which should be tabular data in a txt file. Convert tabular data to genomic ranges by:

```
library(readr)
library(GenomicRanges)
# Downloaded from m6A-Atlas database
my.file="/path/to/home/tangyujiao/big/download/m6A_H.sapiens_basical_information
.txt"
# Load txt file
m6A_basic_info <- read_table2(my.file, col_names = FALSE)</pre>
m6A_basic_info <- m6A_basic_info[, c(1:11)]</pre>
colNames<- c("ID","chr","start","end", "num1","strand", "LOC", "ENS","RNA",
"Gene", "seq")
colnames(m6A_basic_info) <- colNames</pre>
# Concert to Grange object
mod_annot <- makeGRangesFromDataFrame(m6A_basic_info,</pre>
                          keep.extra.columns=FALSE,
                          ignore.strand=FALSE,
                          seginfo=NULL,
                          seqnames.field="chr",
                          start.field="start",
                          end.field="end",
                          strand.field="strand",
                          starts.in.df.are.Obased=FALSE)
# Save Grange to rds
saveRDS(mod_annot, "/path/to/mod_annot.rds")
```

## 2. Reference-based Analysis

```
library(exomePeak2)
set.seed(1)
root = "/path/to/homo_result"
setwd(root)

f1 = file.path(root, "SRR5978834_sorted.bam")
f2 = file.path(root, "SRR5978835_sorted.bam")
f3 = file.path(root, "SRR5978836_sorted.bam")
IP_BAM = c(f1,f2,f3)

f1 = file.path(root, "SRR5978827_sorted.bam")
f2 = file.path(root, "SRR5978828_sorted.bam")
f3 = file.path(root, "SRR5978829_sorted.bam")
```

An output folder named <code>exomePeak2\_output</code> will be created in the working directory containing: