

# Assign taxonomy to gene calls using Centrifuge

James Thornton Jr

## Abstract

Uses a custom Centrifuge pipeline to assign taxonomy to gene calls.

**Citation:** James Thornton Jr Assign taxonomy to gene calls using Centrifuge. **protocols.io**

dx.doi.org/10.17504/protocols.io.kpfcvjn

**Published:** 07 Nov 2017

## Protocol

### Step 1.

Log into the HPC

```
cmd COMMAND
$ ssh hpc
$ ocelote
```

### Step 2.

Move into your class directory.

```
cmd COMMAND
$ cd /rsgroups/bh_class/username
```

### Step 3.

Clone the Centrifuge github repository.

```
cmd COMMAND
$ git clone git@github.com:jetjr/Centrifuge.git
```

### Step 4.

Move into the Centrifuge directory.

```
cmd COMMAND
$ cd Centrifuge
```

## Dependencies

### Step 5.

This program uses R packages that must be installed prior to launching the job. Load the R module.

```
cmd COMMAND
$ module load unsupported
$ module load markb/R/3.1.1
```

## Dependencies

### Step 6.

Launch R.

```
cmd COMMAND  
$ R
```

## Dependencies

### Step 7.

Get the "optparse" package.

```
cmd COMMAND  
> install.packages("optparse", repos="http://R-Forge.R-project.org")
```

## Dependencies

### Step 8.

Get ggplot2 and plyr packages. You may be prompted to select a mirror. Any US server will work.

```
cmd COMMAND  
> install.packages("ggplot2")  
> install.packages("plyr")
```

#### 📌 NOTES

**James Thornton Jr** 07 Nov 2017

If you receive an error when installing the dependencies, continue with the protocol.

## Dependencies

### Step 9.

Quit the R session. Do not save workspace image.

```
cmd COMMAND  
> q()  
> Save workspace image? [y/n/c]: n
```

### Step 10.

Edit the config.sh file to include the correct variable declarations. The following steps will detail how the config.sh file should be edited.

```
cmd COMMAND  
$ nano config.sh
```

## CENT\_DB

### Step 11.

```
export CENT_DB="/rsgrps/bh_class/b_compressed+h+v/b_compressed+h+v"
```

## FASTA\_DIR

### Step 12.

```
export FASTA_DIR="/rsgrps/bh_class/username/prodigal"
```

## 🔗 NOTES

**James Thornton Jr** 07 Nov 2017

FASTA\_DIR should point to the directory containing your nucleotides.fna file generated from step 2 and transferred to the anvio-genes directory.

### TYPE

#### Step 13.

```
export TYPE="single"
```

### FILE\_EXT

#### Step 14.

```
export FILE_EXT="fna"
```

### REPORT\_DIR

#### Step 15.

```
export REPORT_DIR="/rsgrps/bh_class/username/taxonomy"
```

## 🔗 NOTES

**James Thornton Jr** 07 Nov 2017

The program will create this directory for you. Make sure to replace username.

### PLOT\_OUT

#### Step 16.

```
export PLOT_OUT='/rsgrps/bh_class/username/taxonomy/'
```

## 🔗 NOTES

**James Thornton Jr** 07 Nov 2017

Same as REPORT\_DIR but make sure to include the trailing / as stated in the config.sh file.

### PLOT\_FILE and PLOT\_TITLE

#### Step 17.

These should be named according to what sample you're working with. For example, ocean data may name these:

```
export PLOT_FILE='ocean_depth'
```

```
export PLOT_TITLE='ocean_depth'
```

## 📌 NOTES

**James Thornton Jr** 07 Nov 2017

PLOT FILE will be the file name of the bubble plot that is generated.

PLOT TITLE will be the title found on the actual plot.

## FILE\_TYPE

### Step 18.

```
export FILE_TYPE="f"
```

## 📌 NOTES

**James Thornton Jr** 07 Nov 2017

The nucleotides.fna file is in FASTA format.

## EXCLUDE

### Step 19.

The exclude parameter can be left blank.

```
export EXCLUDE=""
```

### Step 20.

Save and quit config.sh

### Step 21.

Submit the job using the submit script found in the Centrifuge directory.

```
cmd COMMAND  
$ ./submit.sh
```

### Step 22.

Status of the job can be determined by the following command:

```
cmd COMMAND  
$ stat -u username
```