

Mac version

Using ReadLine, Boost, HDF5, GSL

mothur v.1.47.0

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by

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<http://www.mothur.org>

When using, please cite:

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Type 'help()' for information on the commands that are available

For questions and analysis support, please visit our forum at <https://forum.mothur.org>

Type 'quit()' to exit program

[NOTE]: Setting random seed to 19760620.

Interactive Mode

```
mothur > make.contigs(file=stability.files)
```

Using 8 processors.

Unable to open F3D144_S120_L001_R1_001.fastq. Trying MOTHUR_FILES

directory mothur/F3D144_S120_L001_R1_001.fastq.

Unable to open mothur/F3D144_S120_L001_R1_001.fastq. Trying mothur's executable directory mothur/F3D144_S120_L001_R1_001.fastq.

Unable to open mothur/F3D144_S120_L001_R1_001.fastq.

[WARNING]: can't find mothur/F3D144_S120_L001_R1_001.fastq, ignoring pair.

```
>>>>> Processing file pair F3D0_S188_L001_R1_001.fastq -
```

```
F3D0_S188_L001_R2_001.fastq (files 1 of 3) <<<<<
```

```
Making contigs...
```

```
Done.
```

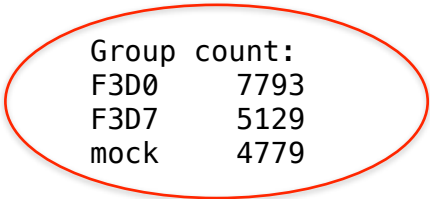
It took 1 secs to assemble 7793 reads.

```
>>>> Processing file pair F3D7_S195_L001_R1_001.fastq -
F3D7_S195_L001_R2_001.fastq (files 2 of 3) <<<<
Making contigs...
Done.
```

It took 1 secs to assemble 5129 reads.

```
>>>> Processing file pair Mock_S280_L001_R1_001.fastq -
Mock_S280_L001_R2_001.fastq (files 3 of 3) <<<<
Making contigs...
Done.
```

It took 0 secs to assemble 4779 reads.



```
Group count:
F3D0      7793
F3D7      5129
mock      4779
```

Total of all groups is 17701

It took 3 secs to process 17701 sequences.

Output File Names:
stability.trim.contigs.fasta
stability.scrap.contigs.fasta
stability.contigs_report
stability.contigs.count_table

```
mothur > summary.seqs(fasta=stability.trim.contigs.fasta)
```

Using 8 processors.

	Start	End	NBases	Ambigs	Polymer	NumSeqs
Minimum: 1	249	249	0	3	1	
2.5%-tile:	1	252	252	0	4	443
25%-tile:	1	252	252	0	4	4426
Median: 1	253	253	0	4	8851	
75%-tile:	1	253	253	0	5	13276
97.5%-tile:	1	254	254	6	6	17259
Maximum: 1	502	502	249	243	17701	
Mean: 1	252	252	0	4		
# of Seqs:	17701					

It took 0 secs to summarize 17701 sequences.

Output File Names:
stability.trim.contigs.summary

```
mothur >  
[ERROR]: You are missing (  
[ERROR]: Invalid.
```

```
mothur >  
[ERROR]: You are missing (  
[ERROR]: Invalid.
```

```
mothur > screen.seqs(fasta=stability.trim.contigs.fasta,  
groups=stability.contigs.count_table, maxambig=0, maxlength=275)  
[WARNING]: groups is not a valid parameter, ignoring.  
The valid parameters are: fasta, contigsreport, alignreport, summary,  
name, count, group, qfile, taxonomy, start, end, maxambig, maxhomop,  
minlength, maxlength, processors, criteria, optimize, seed, inputdir,  
outputdir, minoverlap, ostart, oend, mismatches, maxn, minscore,  
maxinsert, and minsim.
```

Using 8 processors.

It took 0 secs to screen 17701 sequences, removed 2565.

Output File Names:
stability.trim.contigs.good.fasta
stability.trim.contigs.bad.accnos

It took 0 secs to screen 17701 sequences.

```
mothur > get.current()
```

Current RAM usage: 0.133652 Gigabytes. Total Ram: 8 Gigabytes.

Current files saved by mothur:
accnos=stability.trim.contigs.bad.accnos
fasta=stability.trim.contigs.good.fasta
contigsreport=stability.contigs_report
count=stability.contigs.count_table
processors=8
summary=stability.trim.contigs.summary

Current default directories saved by mothur:
mothur/

Current working directory: /Users/natalieburkhard/bio-490/
independentStudy/Lab10/

Output File Names:
current_files.summary

```
mothur > unique.seqs(fasta=stability.trim.contigs.good.fasta)
15136    2698
```

Output File Names:
stability.trim.contigs.good.unique.fasta
stability.trim.contigs.good.count_table

```
mothur > count.seqs(name=stability.trim.contigs.good.names)
Unable to open stability.trim.contigs.good.names. Trying MOTHUR_FILES
directory mothur/stability.trim.contigs.good.names.
Unable to open mothur/stability.trim.contigs.good.names. Trying
mothur's executable directory mothur/
stability.trim.contigs.good.names.
Unable to open mothur/stability.trim.contigs.good.names.
Unable to open stability.trim.contigs.good.names
You have no current namefile or sharedfile and the name or shared
parameter is required, unless inflating or deflating an existing count
file.
[ERROR]: did not complete count.seqs.
```

```
mothur > count.seqs(name=stability.trim.contigs.good.count_table)
```

It took 0 secs to create a table for 2699 sequences.

Total number of sequences: 2699

Output File Names:
stability.trim.contigs.good.count_table

```
mothur > summary.seqs(count=stability.trim.contigs.good.count_table)
Using stability.trim.contigs.good.unique.fasta as input file for the
fasta parameter.
```

```
Using 8 processors.
[ERROR]: Your count file contains 2699 unique sequences, but your
fasta file contains 2698. File mismatch detected, quitting command.
```

```
mothur > unique.seqs(fasta=stability.trim.contigs.good.unique.fasta)
2698    2698
```

Output File Names:

```
stability.trim.contigs.good.unique.unique.fasta
stability.trim.contigs.good.unique.count_table
```

```
mothur > count.seqs(name=stability.trim.contigs.good.names)
Unable to open stability.trim.contigs.good.names. Trying MOTHUR_FILES
directory mothur/stability.trim.contigs.good.names.
Unable to open mothur/stability.trim.contigs.good.names. Trying
mothur's executable directory mothur/
stability.trim.contigs.good.names.
Unable to open mothur/stability.trim.contigs.good.names.
Unable to open stability.trim.contigs.good.names
Using stability.trim.contigs.good.count_table as input file for the
name parameter.
[ERROR]: did not complete count.seqs.
```

```
mothur > count.seqs(name=stability.trim.contigs.good.unique.fasta)
```

It took 0 secs to create a table for 4047 sequences.

Total number of sequences: 4047

Output File Names:

```
stability.trim.contigs.good.unique.count_table
```

```
mothur > summary.seqs(count=stability.trim.contigs.good.count_table)
Using stability.trim.contigs.good.unique.unique.fasta as input file
for the fasta parameter.
```

Using 8 processors.

[ERROR]: Your count file contains 2699 unique sequences, but your
fasta file contains 2698. File mismatch detected, quitting command.

```
mothur > summary.seqs(count=stability.trim.contigs.good.unique.fasta)
Using stability.trim.contigs.good.unique.unique.fasta as input file
for the fasta parameter.
```

Using 8 processors.

[ERROR]: 'M00967_43_000000000-A3JHG_1_2111_6887_10490' is not in your
name or count file, please correct.

```
mothur >
unique.seqs(fasta=stability.trim.contigs.good.unique.count_table)
1350      1350
```

Output File Names:

```
stability.trim.contigs.good.unique.unique.count_table
stability.trim.contigs.good.unique.count_table
```

```
mothur >
count.seqs(name=stability.trim.contigs.good.unique.count_table)
```

It took 0 secs to create a table for 1351 sequences.

Total number of sequences: 1351

Output File Names:
stability.trim.contigs.good.unique.count_table

```
mothur >
summary.seqs(count=stability.trim.contigs.good.unique.count_table)
Using stability.trim.contigs.good.unique.unique.count_table as input
file for the fasta parameter.
```

```
Using 8 processors.
[ERROR]: Your count file contains 1351 unique sequences, but your
fasta file contains 1350. File mismatch detected, quitting command.
```

```
mothur > quit
Logfile : mothur.1647403825.logfile
```

```

*****
*****
*****
Detected 10 [ERROR] messages, please review.
*****
*****
*****

```

[illegible]