

## QIIME2-DEBLUR PROCEDURE TO MAKE OTU TABLES

### Start qiime2

source activate qiime2-2018.6

### Go to qiime2 folder

cd qiime2

### Make directory to work in

mkdir q2\_"todaysdate"

cd q2\_"todaysdate"

### Make manifest file which shows path to data files

sample-id,absolute-filepath,direction  
sample-1,\$HOME/path/"name".fastq,forward  
sample-2,\$HOME/path/"name".fastq,forward  
sample-1,\$HOME/path/"name".fastq,reverse  
sample-2,\$HOME/path/"name".fastq,reverse

### Import files

qiime tools import --type 'SampleData[PairedEndSequencesWithQuality]' --input-path manifest --output-path paired-end-demux.qza --source-format PairedEndFastqManifestPhred33

### Denoise with Deblur

<https://docs.qiime2.org/2018.6/tutorials/read-joining/>

#### *Join pairs*

qiime vsearch join-pairs --i-demultiplexed-seqs paired-end-demux.qza --o-joined-sequences demux-joined.qza  
*\*Can vary the --p-maxdiffs parameter (default is 10)*

#### *View summary*

qiime demux summarize --i-data demux-joined.qza --o-visualization demux-joined.qzv

#### *Quality filter*

qiime quality-filter q-score-joined --i-demux demux-joined.qza --o-filtered-sequences demux-joined-filtered.qza --o-filter-stats demux-joined-filter-stats.qza

#### *Deblur*

qiime deblur denoise-16S --i-demultiplexed-seqs demux-joined-filtered.qza --p-trim-length 250 --p-sample-stats --o-representative-sequences rep-seqs.qza --o-table table.qza --o-stats deblur-stats.qza

### Export table

mkdir exported  
qiime tools export table.qza --output-dir exported  
qiime tools export rep-seqs.qza --output-dir exported

### Convert biom to tab file

biom convert -i exported/feature-table.biom -o table.txt --to-tsv