

Qiime

- Software is a series of python scripts
- Modular, and incorporates other software and algorithms as dependencies (e.g. BLAST, SWARM, uclust, cd-hit, fastq-join, seqprep, mothur, ChimeraSlayer, uchime, etc.)
 - High number of dependencies makes a native install of qiime....difficult
 - Easy alternatives for installation and use:
 - MacQiime
 - Qiime Virtual box
 - Qiime Amazon Web Services Virtual Machine

Qiime Documentation

 Each script has its own page on the qiime website that lists all of its options



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filter_distance_matrix.py —
Filter a distance matrix to
contain only a specified set of
samples.

Next topic

filter_otus_by_sample.py — Filter OTU mapping file and sequences by SampleIDs

Quick search

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Enter search terms or a module, class or function name.

filter_fasta.py - This script can be applied to remove sequences from a fasta or fastq file based on input criteria.

Description:

Usage: filter fasta.py [options]

Input Arguments:

[REQUIRED]

-f, --input_fasta_fp

Path to the input fasta file

-o, --output_fasta_fp

The output fasta filepath

[OPTIONAL]

-m, --otu map

An OTU map where sequences ids are those which should be retained.

-s, --seq id fp

A list of sequence identifiers (or tab-delimited lines with a seq identifier in the first field) which should be retained.

-b, --biom fp

A biom file where otu identifiers should be retained.

-a, --subject fasta fp

All giime commands have the same format

Script_name.py |--option myvalue -o myvalue

- name of the script you want to call
- usually intuitive eg. join_paired_ends.py, pick otus.py
- must always be in the first position

All giime commands have the same format

Script_name.py |--option myvalue -f myvalue

- name of the script you want to call
- usually intuitive eg. join_paired_ends.py, pick otus.py
- must always be in the first position

All giime commands have the same format

Script_name.py |--option|myvalue -f myvalue

- parameter for the script
- always preceded by ---
- usually intuitive eg.
- --max ambig or --fasta
- can have many options in any order

- value for the option
- separated from the option name by a space
- can be a filename, an algorithm name, a numeric value, or a TRUE/FALSE

All qiime commands have the same format

Script_name.py --option myvalue -f myvalue

- Qiime also uses shorthand to save you typing
- shorthand preceded by –
- here, -f would be the same as -fasta

Qiime Input Files

- 1. Sequences! (fastq, fasta + qual, or fasta)
- 2. Mapping File
 - Specifies metadata about samples
 - Includes list of barcodes, primers, etc.
- 3. Parameter File
 - Specifies your parameters for multiple

Qiime Input Files: Mapping File

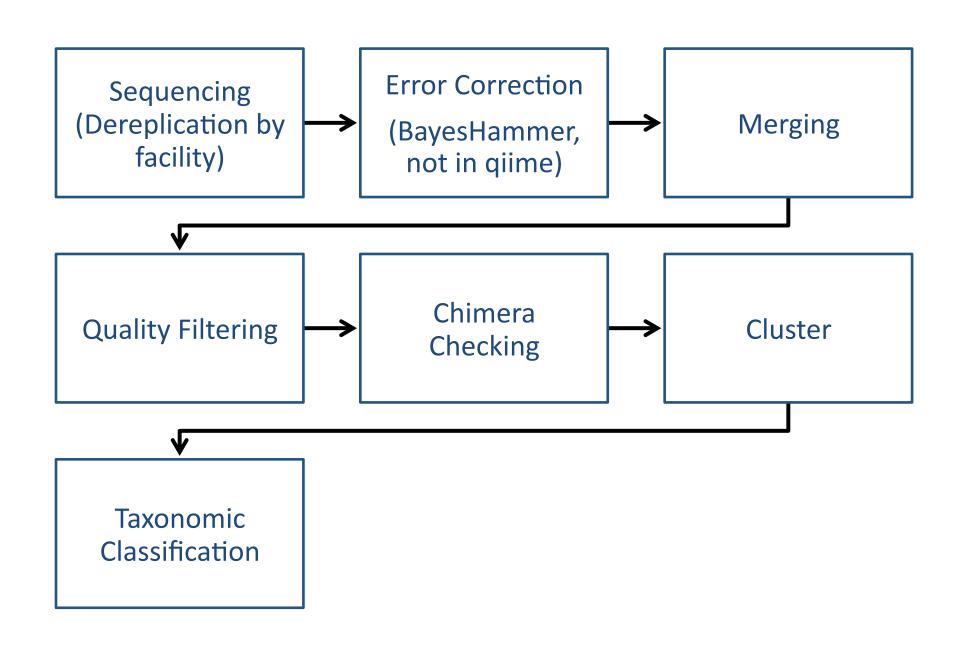
- Tab delimited text file
- Header line is very important
 - First field MUST BE #SampleID, second MUST BE BarcodeSequence, third MUST BE LinkerPrimerSequence, and last MUST BE Description

#SampleI	D BarcodeSequ	uence LinkerPrimerSequence	e Tre	atment DOB	Description
PC.354	AGCACGAGCCTA	YATGCTGCCTCCCGTAGGAGT	Control	20061218	Control_mouse_I.D354
PC.355	AACTCGTCGATG	YATGCTGCCTCCCGTAGGAGT	Control	20061218	Control_mouse_I.D355
PC.356	ACAGACCACTCA	YATGCTGCCTCCCGTAGGAGT	Control	20061126	Control_mouse_I.D356
PC.481	ACCAGCGACTAG	YATGCTGCCTCCCGTAGGAGT	Control	20070314	Control_mouse_I.D481
PC.593	AGCAGCACTTGT	YATGCTGCCTCCCGTAGGAGT	Control	20071210	Control_mouse_I.D593
PC.607	AACTGTGCGTAC	YATGCTGCCTCCCGTAGGAGT	Fast	20071112	Fasting_mouse_I.D607
PC.634	ACAGAGTCGGCT	YATGCTGCCTCCCGTAGGAGT	Fast	20080116	Fasting_mouse_I.D634
PC.635	ACCGCAGAGTCA	YATGCTGCCTCCCGTAGGAGT	Fast	20080116	Fasting_mouse_I.D635
PC.636	ACGGTGAGTGTC	YATGCTGCCTCCCGTAGGAGT	Fast	20080116	Fasting_mouse_I.D636

Qiime Input Files: Parameter File

- Parameter files are used for qiime scripts that are 'wrappers'
 - Wrappers run several scripts one after the another, or run the same script multiple times
- Text file where each line specifies a command, option, and value
- Consistent format script_name:parameter_name value

```
join_paired_ends:pe_join_method fastq-join
join_paired_ends:min_overlap 10
join_paired_ends:perc_max_diff 90
split_libraries_fastq:max_bad_run_length 3
split_libraries_fastq:min_per_read_length_fraction 0.75
split_libraries_fastq:sequence_max_n 0
split_libraries_fastq:phred_quality_threshold 30
```



Qiime Python Scripts Used in our Workflow

- multiple_join_paired_ends.py (merge sequences)
- multiple_split_libraries_fastq.py (quality filter)
- identify_chimeric_seqs.py (chimera check)
- filter_fasta.py (remove chimeras)
- pick_otus.py, pick_open_reference_otus.py (cluster)
- pick_rep_set.py (get representative sequences)
- assign_taxonomy.py (taxonomic assignment)
- make_otu_table.py (build OTU table)
- core_diversity_analyses.py (fun with data!)