AMOEBAE documentation

Lael D. Barlow

Contents

1 Introduction

1.1 What is AMOEBAE?

Analysis of MOlecular Evolution with BAtch Entry (AMOEBAE) is a bioinformatics software toolkit composed primarily of scripts written in the Python3 language. AMOEBAE scripts use existing Python packages including Biopython (Cock *et al.*, 2009), the Environment for Tree Exploration (ETE3) (Huerta-Cepas *et al.*, 2016), pandas, and Matplotlib (Hunter, 2007) for setting up, running, and summarizing analyses of molecular evolution using bioinformatics software packages including MUSCLE (Edgar, 2004), BLAST+ (Camacho *et al.*, 2009), HMMer3 (Eddy, 1998), and IQ-Tree (Nguyen *et al.*, 2015). Applications include identifying and classifying predicted peptide sequences according to their evolutionary relationships with homologues. All dependencies are freely available, and AMOEBAE code is open-source (see subsection 1.8) and available on GitHub (`https://github.com/laelbarlow/amoebae`).

1.2 Why use AMOEBAE?

Webservices such as those provided by NCBI (`https://blast.ncbi.nlm.nih.gov/Blast.cgi`) (Camacho *et al.*, 2009) provide a means to investigate the evolution of one or a few genes via similarity searching, and automated pipelines such as orthoMCL (Li, 2003) attempt to rapidly perform orthology prediction for all genes in several genomes. AMOEBAE addresses the problem mid-scale analyses which are too cumbersome to be done via webservices and yet requiring a level of detail and flexibility not offered by automated pipelines. AMOEBAE may be useful for analyzing the distribution of orthologues of up to perhaps 30 genes/proteins among a sampling of no more than approximately 100 eukaryotic genomes. However, you may need to carefully define the scope of your analysis depending on what additional steps you may find necessary beyond those that may be performed using AMOEBAE (30 queries and 100 genomes may in fact be unmanageable). AMOEBAE provides many options which can be tailored to the specific genes/proteins being analyzed, and allow analyses using complex sets of customized criteria to be reproduced more practically.

1.3 Key features

The core functionality is to run sequence similarity searches with multiple algorithms, multiple queries, and multiple databases simultaneously in parallel and to facilitate efficient and highly customizable implementation of reciprocal-best-hit search strategies. The output includes detailed summaries of results in the form of a spreadsheet and plots.

A particular advantage of AMOEBAE over other tools is the functionality for parsing results of TBLASTN (searching in nucleotide sequences with peptide sequence queries) search results. This allows rapid identification of High-scoring Segment Pair (HSP) clusters at separate gene loci (identified according to user-defined criteria), automatic checking of those loci

against information in genome annotation files, and systematic use of Exonerate (Slater and Birney, 2005) where possible for obtaining better exon predictions.

1.4   User support

For specific issues with the code, please use the issue tracker on the GitHub webpage here: `https://github.com/laelbarlow/amoebae/issues`.

If you have general questions regarding AMOEBAE, please email the author at lael (at) ualberta.ca.

1.5   Documentation

This document provides an overview of AMOEBAE and describes the functionality of the various commands/scripts. For a tutorial which includes a working example of a similarity search analysis run using AMOEBAE, see the Jupyter Notebook: amoebae/notebooks/similarity_search_tutorial_2.ipynb. For code documentation, please see the html file(s), which can be opened with your web browser: `amoebae/doc/code_documentation/html/index.html`.

1.6   How to cite AMOEBAE

Please cite the GitHub webpage `https://github.com/laelbarlow/amoebae` (or alternative permanent repositories if relevant). Also, the first publication to make use of a version of AMOEBAE was an analysis of Adaptor Protein subunits in embryophytes by Larson *et al.* (2019).

Also, you may wish to cite the software packages which are key dependencies of AMOEBAE, since AMOEBAE would not work without these (see subsection 2.2).

1.7   Acknowledgments

Also, help with testing AMOEBAE has been kindly provided by Raegan T. Larson, Shweta V. Pipalya, Kira More, and Kristýna Záhonová.

1.8 License

2 How to start using AMOEBAE

2.1 System requirements

Please note that the commands shown likely only work on macOS or Linux operating systems (you may have trouble running AMOEBAE directly on Windows).

2.2 Dependencies

All dependencies are free and open-source, and can be automatically installed in a virtual environment (see subsection 2.3).

These are the main depencencies of AMOEBAE:

- Python3 (the Anaconda distribution works well).

- Biopython, a Python package for bioinformatics (Cock *et al.*, 2009).

- The Environment for Tree Exploration 3 (ETE3), a Python package for working with phylogenetic trees (Huerta-Cepas *et al.*, 2016).

- Matplotlib, a Python package for generating plots (Hunter, 2007).

- (gffutils).

- NCBI BLAST+, a software package for sequence similarity searching (Camacho *et al.*, 2009).

- HMMer3, a software package for profile sequence similarity searching (Eddy, 1998).

- MUSCLE, for multiple sequence alignment (Edgar, 2004).

- IQ-TREE, for phylogenetic analysis (Nguyen *et al.*, 2015).

2.3 Setting up an environment for AMOEBAE using Singularity

Follow the steps below to set up AMOEBAE on your personal computer. This setup process should take approximately 20 minutes to complete. Instructions for setting up AMOBEAE on a remote server will soon be added as well.

1. Ensure that Git is installed on your computer If you do not already have git installed, then your computer will prompt you with instructions for how to install it when you type git into the command line. If you have a newer version of macOS it may prompt you to install developer tools, which may take up a considerable amount of storage space. Documentation for Git is available here: `https://git-scm.com/doc`. You can check which version you have (or whether it is installed at all) by running the command below. Please note: Here ">>>" is used to indicate that the following text in the line is to be entered in you terminal command prompt.

   ```
   >>> git --version
   ```

2. Clone the AMOEBAE repository using Git. If you simply download the code from GitHub, instead of cloning the repository, then AMOEBAE cannot record specifically what version of the code you use, and will not run properly. Make sure to use the appropriate directory path (the path shown is just an example). Also, replace the path shown below with the path to the directory on your system where you wish to put the main AMOEBAE directory.

   ```
   >>> cd /path/to/directory/where/you/keep/files
   >>> git clone https://github.com/laelbarlow/amoebae.git
   ```

3. Set up AMOEBAE. This performs several steps including checking for whether singularity is installed and attempting to use VirtualBox and Vagrant to run Singularity in a pre-built Ubuntu virtual machine with Singularity installed. This is because Singularity does not run on MacOS (or Windows), and installation of Singularity on Linux is complex, as several dependencies are required. This script downloads a pre-built singularity container, which was built using the singularity.recipe file, and provided on the Singularity Library (`https://cloud.sylabs.io/library/_container/5e8ca8fff0f8eb90a8a7b60d`).

   ```
   >>> cd amoebae
   >>> bash setup.sh
   ```

4. If you are setting up AMOEBAE on a high performance computing cluster, then you will not be able to install Singularity yourself, and may need to use specific procedures to load Singularity prior to use.

2.4 Running AMOEBAE using Jupyter notebooks

1. After setting up AMOEBAE according to the instructions above, the easiest way to start running analyses using AMOEBAE is via the tutorials, which are in the form of Jupyter notebooks (`https://jupyter.org/`). These Jupyter notebooks can be run using the installation of Jupyter in the Singularity container, and can be accessed using your browser (on a personal computer). To start a Jupyter server, run the bash script as indicated below (assuming your current working directory is the main amoebae directory that you cloned with Git).

   ```
   >>> bash singularity_jupyter.sh
   ```

2. Copy and past the resulting URL (the one at the bottom of the output) into the address bar of your web browser (either Firefox, Chrome, or Safari will work). This will open Jupyter to the notebooks subdirectory, which contains several tutorial and example notebooks (.ipynb files). These files are the files on your regular (host) filesystem, as the amoebae directory is synced with the Singularity container. Thus changes to files will persist after you shut down the Jupyter server and the Singularity container. Documentation on Jupyter is available here: `https://jupyter-notebook.readthedocs.io/en/stable/`.

3. Click on one of the tutorial files (.ipynb). These Jupyter notebooks include information on how to use them once opened. The first tutorial (amoebae_tutorial_1.ipynb) provides a simple example of similarity searching with BLASTP using a Jupyter notebook. The second tutorial (amoebae_tutorial_2.ipynb) provides an example using most of the similarity searching functionality that AMOEBAE provides.

4. To shut down the Jupyter server, click the logout button in the jupyter browser tab(s), and then go to the terminal window that you used to startup the Jupyter server, and press CTRL-C to kill the Jupyter kernel. This will close the Jupyter notebooks, but the analysis output files will remain, because they are saved to your amoebae/notebooks folder which is on your host machine and accessed from within the container.

5. Working with the Jupyter notebooks interactively in this manner on high-performance computing clusters is likely possible, but inconvenient, and procedures will vary. Also, running the tutorial notebooks would require access to the internet from compute nodes (as opposed to login nodes) which may not be supported. Therefore, it is recommended that you run the tutorials on a personal laptop/desktop computer if possible. To run your own notebooks on a cluster, you will need to write a job submission script that will be specific to the cluster, the job schedule it uses, and your account details. Please refer to documentation provided by your system administrators for this. For an example script that writes a script for running a notebook as a job to a SLURM job

scheduler see https://github.com/laelbarlow/amoebae/blob/master/notebooks/ write_notebook_slurm_script.sh.

2.5 Running AMOEBAE via the command line

1. The easiest way to access AMOEBAE dependencies via the command line is to use the bash script provided. If you are running singularity in a virtual machine (*e.g.*, on MacOS), then only one shell session may be opened at once (and these cannot be opened at the same time as the singularity_jupyter.sh script is running Singularity in a virtual machine). Running the script as indicated below will open a shell session in the Singularity container, with the amoebae directory being the only one accessible. Also, the amoebae executable script is added to the $PATH in the container, so you can run amoebae commands from any directory.

```
>>> bash singularity_shell.sh
```

2. You may find it useful to explore and test the environment using the following commands.

   - Print the paths included in the $PATH variable in the container.
     ```
     >>> tr ':' '\n' <<< "$PATH"
     ```

   - Check the location of the amoebae executable being run from within the container.
     ```
     >>> command -v amoebae
     ```

   - Check that the amoebae executable script can be run (print the help message).
     ```
     >>> amoebae -h
     ```

   - Check that all modules can be imported in all python files in the AMOEBAE code.
     ```
     >>> amoebae check_imports
     ```

   - Check that key dependencies such as BLASTP can be accessed (they are installed in the Singularity container).
     ```
     >>> amoebae check_depend
     ```

3. Again, running AMOEBAE commands on high-performance computing clusters will require you to write custom job submission scripts. Please refer to documentation provided by your system administrator(s) regarding details specific to your cluster, including the job scheduler used. Also, refer to the Singularity documentation for formulating Singularity commands (https://sylabs.io/docs/).

3 Command reference

Documentation for each AMOEBAE command and the various options may be accessed from the command line via the "-h" options. The following command reference information is the output of running amoebae (and each command) with the "-h" option.

3.1 amoebae

```
usage: amoebae <command> [<args>]

Commands for setting up data structure:
    mkdatadir       Make a directory with subdirectories and CSV files for
                    storing sequence data, etc.

Commands for similarity searching:
    setup_hmmdb     Construct an HMM database (with hmmpress).
    add_to_dbs      Format and add a file to a formatted directory.
    list_dbs        Print a list of all usable database files in the database
                    directory as defined in the settings file.
    add_to_queries  Add a query file to a formatted directory.
    list_queries    Print a list of all usable query files in the query
                    directory as defined in the settings file.
    get_redun_hits  Run searches with queries to find redundant hits in
                    databases (for interpreting results).
    setup_fwd_srch  Make directory in which to perform forward searches.
    run_fwd_srch    Perform searches with given queries into given dbs.
    sum_fwd_srch    Append information about forward searches to csv summary
                    file (this is used to organize reverse searches).
    setup_rev_srch  Make a directory in which to perform reverse searches.
    run_rev_srch    Perform searches with given forward search hits into given db.
    sum_rev_srch    Append information about reverse searches to csv summary
                    file.
    interp_srchs    Interpret search results based on summary.
    find_redun_seqs Identify sequences likely encoded on redundant loci
                    predicted for the same species.
    plot            Plot search results.

Commands for phylogenetic analysis using a reference tree:
    add_to_models   Add an alignment, tree, substitution model, names of
                    clade-defining sequences to a directory with other models.
    list_models     Print a list of all usable model/reference tree names in
                    the models directory as defined in the settings file.
    get_alt_topos   Take a tree and make copies with every alternative
                    topology for the branches connecting the clades of
                    interest.

Commands for phylogenetic analysis without a reference tree:
```

```
prune               Identify sequences in a tree, and remove them from a
                    given alignment for further phylogenetic analysis.
auto_prune          Automatically identify sequences in a tree, and remove
                    them from a given alignment for further phylogenetic
                    analysis.
reduce_tree         Remove terminal nodes from a given tree if there are
                    not any sequences with the same name in a given multiple
                    sequence alignment file.
constrain_mb        Add constraint commands to MrBayes input file based on a
                    given tree topology.
visualize_tree      Parse phylogenetic analysis output files for a single
                    alignment in a given directory, and write human-readable
                    tree figures to PDF files.
replace_seqs        Replace sequences in an alignment with their top hits in a
                    given fasta file (useful if genomes or taxon selection has
                    been updated).

Miscellaneous commands:
csv_to_fasta        Generate a fasta file from sequences detailed in a
                    spreadsheet of similarity search results.
check_depend        Check that all the dependencies are properly installed and
                    useable.
check_imports       Check that all the import statements used in the AMOEBAE
                    repository run without error.

positional arguments:
  command     Specify one of the functionalities of amoebae.

optional arguments:
  -h, --help  show this help message and exit

Copyright 2018 Lael D. Barlow Licensed under the Apache License, Version 2.0
(the "License"); you may not use this file except in compliance with the
License. You may obtain a copy of the License at
http://www.apache.org/licenses/LICENSE-2.0 Unless required by applicable law
or agreed to in writing, software distributed under the License is distributed
on an "AS IS" BASIS, WITHOUT WARRANTIES OR CONDITIONS OF ANY KIND, either
express or implied. See the License for the specific language governing
permissions and limitations under the License.
```

3.2   amoebae mkdatadir

```
usage: amoebae [-h] new_dir_path

Make a directory with subdirectories and CSV files for storing sequence data,
etc.

positional arguments:
  new_dir_path  Specify the full file path that you want the new directory to
```

```
1              have.

2

3 optional arguments:
4   -h, --help    show this help message and exit
```

3.3   amoebae setup_hmmdb

```
usage: amoebae [-h] indirpath


Construct an HMM database (with hmmpress). This is for later sorting of given
sequences into categories based on which HMM the score highest against.


positional arguments:
   indirpath   Path to directory containing amino acid sequence alignment
               file(s) to be constructed into an HMM database using hmmpress
               from the HMMer3 software package.


optional arguments:
   -h, --help  show this help message and exit
```

3.4   amoebae add_to_dbs

```
usage: amoebae [-h] [--split_char SPLIT_CHAR] [--split_pos SPLIT_POS]
               [--skip_header_reformat] [--auto_extract_accs]
               new_file


Format and add a file to a formatted directory.


positional arguments:
   new_file                Can be a fasta file (prot or nucl) or HMM databases,
                           generated using the hmmpress program in the HMMer
                           software package. Or a GFF3 annotation file.


optional arguments:
   -h, --help              show this help message and exit
   --split_char SPLIT_CHAR
                           Character to split the header string on for extracting
                           the accession. (default: )
   --split_pos SPLIT_POS
                           Position that the accession will be in after
                           splitting. (default: 0)
   --skip_header_reformat
                           Skip reformatting of header lines in input fasta file.
                           (default: False)
   --auto_extract_accs     Automatically identify accessions/IDs in sequence
                           headers (overrides split_char and split_pos options
                           above). (default: False)
```

3.5 amoebae list_dbs

```
usage: amoebae [-h]

Print a list of all usable query files in the query directory as defined in
the settings file.

optional arguments:
  -h, --help  show this help message and exit
```

3.6 amoebae add_to_queries

```
usage: amoebae [-h] query_file

Add a query file to a formatted directory. This command adds a given sequence
file to the directory with the path that you have specified in the settings.py
file, and appends a corresponding line to the CSV file that you specified
(e.g., '0_query_info.csv') to indicate the query title, etc.

positional arguments:
  query_file  Path to a sequence file in FASTA format that can be used as a
              similarity search query file. Or path to a directory containing
              only files for addition to the queries. Note: By default, the
              portion of the input filename preceding the first underscore
              character will be recorded as the "query title", the remaining
              substring preceding the second underscore character will be
              recorded as the taxon (e.g., "Hsapiens"), and the rest of the
              filename preceding the filename extension will be recorded as
              the sequence ID. So the filename might look like this:
              "QUERYTITLE_HSAPIENS_SEQUENCEID.fa". However, the relevant
              information can be revised in the "Queries/0_query_info.csv"
              file afterward if necessary.

optional arguments:
  -h, --help  show this help message and exit
```

3.7 amoebae list_queries

```
usage: amoebae [-h]

Print a list of all usable query files in the query directory as defined in
the settings file.

optional arguments:
  -h, --help  show this help message and exit
```

3.8 amoebae get_redun_hits

10

```
 1  usage: amoebae [-h] [--csv_file CSV_FILE] [--query_name QUERY_NAME]
 2                 [--query_list_file QUERY_LIST_FILE] [--db_name DB_NAME]
 3                 [--db_list_file DB_LIST_FILE] [--query_title QUERY_TITLE]
 4                 [--outdir OUTDIR]
 5                 [--blast_report_evalue_cutoff BLAST_REPORT_EVALUE_CUTOFF]
 6                 [--blast_max_target_seqs BLAST_MAX_TARGET_SEQS]
 7                 [--hmmer_report_evalue_cutoff HMMER_REPORT_EVALUE_CUTOFF]
 8                 [--hmmer_report_score_cutoff HMMER_REPORT_SCORE_CUTOFF]
 9                 [--num_threads_similarity_searching NUM_THREADS_SIMILARITY_SEARCHING
10      ]
11                 srch_dir
12
13  Run searches with queries to find redundant hits in databases (for
14  interpreting results).
15
16  positional arguments:
17    srch_dir              Path to directory that will contain output directory
18                         as a subdirectory.
19
20  optional arguments:
21    -h, --help           show this help message and exit
22    --csv_file CSV_FILE   Path to spreadsheet to append summary of result to for
23                         manual annotation. (default: None)
24    --query_name QUERY_NAME
25                         Query filename to use (not full path). (default: None)
26    --query_list_file QUERY_LIST_FILE
27                         Path to file containing a list of query files to use,
28                         if no query_name is specified (or all queries by
29                         default). (default: None)
30    --db_name DB_NAME    Name of database file in the database directory in
31                         which to do searches (not full path). (default: None)
32    --db_list_file DB_LIST_FILE
33                         Path to file containing a list of database files to
34                         use (if no db_name specified). (default: None)
35    --query_title QUERY_TITLE
36                         Name to be assigned to hits in databases that may be
37                         considered redundant with a search query to which the
38                         same title is assigned, otherwise it is taken from the
39                         query info spreadsheet specified in the settings.py
40                         file ('query_info_csv'). (default: None)
41    --outdir OUTDIR      Path to directory to write search results to.
42                         (default: None)
43    --blast_report_evalue_cutoff BLAST_REPORT_EVALUE_CUTOFF
44                         Maximum E-value for reporting BLAST hits. (default:
45                         0.05)
46    --blast_max_target_seqs BLAST_MAX_TARGET_SEQS
47                         Maximum BLAST target sequences to consider. (default:
48                         500)
49    --hmmer_report_evalue_cutoff HMMER_REPORT_EVALUE_CUTOFF
```

```
                        Maximum E-value for reporting HMMer hits. (default:
                        0.05)
  --hmmer_report_score_cutoff HMMER_REPORT_SCORE_CUTOFF
                        Minimum sequence score for reporting HMMer hits.
                        (default: 5)
  --num_threads_similarity_searching NUM_THREADS_SIMILARITY_SEARCHING
                        Number of threads to use for running searches.
                        (default: 4)

Recommendation: For most analyses, use the --query_name option and the
--db_name option, and run the get_redun_hits command for each query
separately. Otherwise, there will be redundant information in the output
spreadsheet(s).
```

3.9 amoebae setup_fwd_srch

```
usage: amoebae [-h] [--outdir OUTDIR] srch_dir query_list_file db_list_file

Make a directory in which to write output files from similarity searches.

positional arguments:
  srch_dir         Path to directory that will contain output directory as a
                   subdirectory.
  query_list_file  Path to file with list of queries to search with.
  db_list_file     Path to file with list of databases to search with.

optional arguments:
  -h, --help       show this help message and exit
  --outdir OUTDIR  Path to directory to put search results into (so that this
                   step can be piped together with other commands). (default:
                   None)

Note: Use the bash script to run forward searches on a remote server.
```

3.10 amoebae run_fwd_srch

```
usage: amoebae [-h] [--blast_report_evalue_cutoff BLAST_REPORT_EVALUE_CUTOFF]
               [--blast_max_target_seqs BLAST_MAX_TARGET_SEQS]
               [--hmmer_report_evalue_cutoff HMMER_REPORT_EVALUE_CUTOFF]
               [--hmmer_report_score_cutoff HMMER_REPORT_SCORE_CUTOFF]
               [--num_threads_similarity_searching NUM_THREADS_SIMILARITY_SEARCHING
    ]
               fwd_srch_dir

Perform searches with original queries into subject databases.

positional arguments:
  fwd_srch_dir          Path to directory that will contain forward search
```

```
 1                           output files.
 2
 3  optional arguments:
 4    -h, --help              show this help message and exit
 5    --blast_report_evalue_cutoff BLAST_REPORT_EVALUE_CUTOFF
 6                            Maximum E-value for reporting BLAST hits. (default:
 7                            0.05)
 8    --blast_max_target_seqs BLAST_MAX_TARGET_SEQS
 9                            Maximum BLAST target sequences to consider. (default:
10                            500)
11    --hmmer_report_evalue_cutoff HMMER_REPORT_EVALUE_CUTOFF
12                            Maximum E-value for reporting HMMer hits. (default:
13                            0.05)
14    --hmmer_report_score_cutoff HMMER_REPORT_SCORE_CUTOFF
15                            Minimum sequence score for reporting HMMer hits.
16                            (default: 5)
17    --num_threads_similarity_searching NUM_THREADS_SIMILARITY_SEARCHING
18                            Number of threads to use for running searches.
19                            (default: 4)
```

3.11   amoebae sum_fwd_srch

```
21  usage: amoebae [-h] [--max_evalue MAX_EVALUE]
22                 [--max_gap_between_tblastn_hsps MAX_GAP_BETWEEN_TBLASTN_HSPS]
23                 [--do_not_use_exonerate]
24                 [--exonerate_score_threshold EXONERATE_SCORE_THRESHOLD]
25                 [--max_hits_to_sum MAX_HITS_TO_SUM]
26                 fwd_srch_out csv_file
27
28  Append information about forward searches to csv summary file (this is used to
29  organize reverse searches). For TBLASTN searches (protein queries, nucleotide
30  target sequences), HSPs are clustered into groups that are close enough within
31  the target sequence to potentially represent exons from the same coding
32  sequence. The nucleotide subsequences in which these clusters of HSPs are
33  found are then analyzed using exonerate to identify and translate potential
34  exons, in "protein2genome" mode, because exonerate, unlike TBLASTN, attempts
35  to identify exon boundaries, yielding translations that are less likely to
36  include translations of non-coding regions outside exons (which might include
37  apparent stop codons).
38
39  positional arguments:
40    fwd_srch_out            Path to directory where forward search results were
41                            written.
42    csv_file                Path to summary spreadsheet (CSV) file, which may
43                            already contain search summaries, or may not exist
44                            yet.
45
46  optional arguments:
47    -h, --help              show this help message and exit
```

```
  1    --max_evalue MAX_EVALUE
  2                          Maximum E-value threshold for reporting forward search
  3                          hits. (default: 0.0005)
  4    --max_gap_between_tblastn_hsps MAX_GAP_BETWEEN_TBLASTN_HSPS
  5                          Maximum number of nucleotide bases between TBLASTN
  6                          HSPs to be considered part of the same gene locus.
  7                          This is important, because it will be assumed that HSP
  8                          separated by more than this number of nucleotide bases
  9                          are not part of the same gene or TBLASTN "hit".
 10                          (default: 10000)
 11    --do_not_use_exonerate
 12                          Override the default use of exonerate to identify
 13                          coding sequences and translations, and just use
 14                          TBLASTN instead. This option is provided because
 15                          concatenated TBLASTN HSPs may be more inclusive of
 16                          sequences within the target sequence, and the results
 17                          of TBLASTN and exonerate may need to be compared.
 18                          Also, note that HSPs identified by TBLASTN but for
 19                          which exonerate yields no alignments will be ignored
 20                          if exonerate is used. (default: False)
 21    --exonerate_score_threshold EXONERATE_SCORE_THRESHOLD
 22                          Set score threshold to be applied when running
 23                          exonerate on nucleotide sequences identified by
 24                          TBLASTN. The default for setting of exonerate is 100,
 25                          but a lower score is set as default here, because
 26                          otherwise exonerate cannot identify some of the
 27                          seqeunces identified by TBLASTN. This option is only
 28                          relevant if using exonerate. (default: 10)
 29    --max_hits_to_sum MAX_HITS_TO_SUM
 30                          Maximum number of forward search hits to list in the
 31                          summary spreadsheet. If zero, then reverse searches
 32                          will be performed for all hits. (default: 0)
```

3.12   amoebae setup_rev_srch

```
usage: amoebae [-h] [--outdir OUTDIR] [--aasubseq] [--nafullseq]
               srch_dir csv_file databases

Make directory in which to write results of reverse searches.

positional arguments:
  srch_dir        Path to directory that will contain output directory as a
                  subdirectory.
  csv_file        Path to summary spreadsheet (CSV) file, which contains a
                  summary of forward search(es).
  databases       Database filename (in database directory) or path to file
                  with list of database filenames. Note that filenames are
                  needed, not file paths.
```

```
1  optional arguments:
2    -h, --help        show this help message and exit
3    --outdir OUTDIR   Path to directory to put search results into (so that this
4                      step can be piped together with other commands). (default:
5                      None)
6    --aasubseq        Use only the portion of each (amino acid) forward hit
7                      sequence that aligns to the original query used (top HSP
8                      subject sequence). This is default for nucleotide hits.
9                      (default: False)
10   --nafullseq       Use the full (nucleic acid) forward hit sequence. This is
11                     default for amino acid hits. (default: False)
```

3.13   amoebae run_rev_srch

```
13 usage: amoebae [-h] [--blast_report_evalue_cutoff BLAST_REPORT_EVALUE_CUTOFF]
14                [--blast_max_target_seqs BLAST_MAX_TARGET_SEQS]
15                [--hmmer_report_evalue_cutoff HMMER_REPORT_EVALUE_CUTOFF]
16                [--hmmer_report_score_cutoff HMMER_REPORT_SCORE_CUTOFF]
17                [--num_threads_similarity_searching NUM_THREADS_SIMILARITY_SEARCHING
18     ]
19                rev_srch_dir
20
21 Perform searches with forward search hit sequences as queries into the
22 original query databases.
23
24 positional arguments:
25   rev_srch_dir        Path to directory that will contain output of
26                       searches.
27
28 optional arguments:
29   -h, --help          show this help message and exit
30   --blast_report_evalue_cutoff BLAST_REPORT_EVALUE_CUTOFF
31                       Maximum E-value for reporting BLAST hits. (default:
32                       0.05)
33   --blast_max_target_seqs BLAST_MAX_TARGET_SEQS
34                       Maximum BLAST target sequences to consider. (default:
35                       500)
36   --hmmer_report_evalue_cutoff HMMER_REPORT_EVALUE_CUTOFF
37                       Maximum E-value for reporting HMMer hits. (default:
38                       0.05)
39   --hmmer_report_score_cutoff HMMER_REPORT_SCORE_CUTOFF
40                       Minimum sequence score for reporting HMMer hits.
41                       (default: 5)
42   --num_threads_similarity_searching NUM_THREADS_SIMILARITY_SEARCHING
43                       Number of threads to use for running searches.
44                       (default: 4)
```

3.14 amoebae sum_rev_srch

```
usage: amoebae [-h] [--redun_hit_csv REDUN_HIT_CSV]
               [--min_evaldiff MIN_EVALDIFF] [--aasubseq] [--nafullseq]
               [--max_rev_srchs MAX_REV_SRCHS]
               csv_file rev_srch_out
```

Append information about reverse searches to csv summary file. Use information from redundant hit csv file to interpret results.

```
positional arguments:
  csv_file              Path to summary spreadsheet (CSV) file, which may
                        already contain reverse search summaries.
  rev_srch_out          Path to directory where reverse search results were
                        written.

optional arguments:
  -h, --help            show this help message and exit
  --redun_hit_csv REDUN_HIT_CSV
                        Path to spreadsheet (CSV) file, which specifies which
                        hits are redundant positive hits for a given query
                        (query title) in a given database. If this is not
                        provided, then it is assumed that any and all reverse
                        search hits are equivalent to/redundant with the
                        original query. (default: None)
  --min_evaldiff MIN_EVALDIFF
                        Minimum difference in E-value order of magnitude
                        between top reverse search hit and first reverse
                        search hit that is not redundant with the original
                        query. (default: 5)
  --aasubseq            Use only the portion of each (amino acid) forward hit
                        sequence that aligns to the original query used (top
                        HSP subject sequence). This is default for nucleotide
                        hits. Must be selected if selected when the
                        setup_rev_srch command was run. (default: False)
  --nafullseq           Use the full (nucleic acid) forward hit sequence. This
                        is default for amino acid hits. Must be selected if
                        selected when the setup_rev_srch command was run.
                        (default: False)
  --max_rev_srchs MAX_REV_SRCHS
                        Maximum number of forward search hits to perform
                        reverse searches for per query database. If zero, then
                        reverse searches will be performed for all hits.
                        (default: 0)
```

3.15 amoebae interp_srchs

```
usage: amoebae [-h] [--fwd_only] [--fwd_evalue_cutoff FWD_EVALUE_CUTOFF]
               [--rev_evalue_cutoff REV_EVALUE_CUTOFF]
```

```
1              [--hmmer_cutoff HMMER_CUTOFF] [--redun_hits]
2              [--out_csv_path OUT_CSV_PATH]
3              csv_file
4
5 Interpret search results based on final summary, which provides a basis for
6 further analyses of positive hits.
7
8 positional arguments:
9   csv_file             Path to spreadsheet with forward and reverse search
10                       results.
11
12 optional arguments:
13   -h, --help           show this help message and exit
14   --fwd_only           Interpret forward searches based on score (HMMer)
15                       cutoff. (default: False)
16   --fwd_evalue_cutoff FWD_EVALUE_CUTOFF
17                       Specify an (more stringent) E-value cutoff for forward
18                       search results. (default: None)
19   --rev_evalue_cutoff REV_EVALUE_CUTOFF
20                       Specify an (more stringent) E-value cutoff for reverse
21                       search results. (default: None)
22   --hmmer_cutoff HMMER_CUTOFF
23                       Specify a score that hits must exceed to be included.
24                       (default: 20)
25   --redun_hits         Interpret which hits are redundant in output of
26                       get_redun_hits command. (default: False)
27   --out_csv_path OUT_CSV_PATH
28                       Optionally specify an output file path, so that this
29                       command can be piped together with others. (default:
30                       None)
```

3.16   amoebae find_redun_seqs

```
32 usage: amoebae [-h] [--out_csv_path OUT_CSV_PATH]
33              [--remove_tblastn_hits_at_annotated_loci]
34              [--just_look_for_genes_in_gff3] [--ignore_gff3]
35              [--allow_internal_stops ALLOW_INTERNAL_STOPS]
36              [--min_length MIN_LENGTH]
37              [--min_percent_length MIN_PERCENT_LENGTH]
38              [--min_percent_query_cover MIN_PERCENT_QUERY_COVER]
39              [--overlap_required] [--max_percent_ident MAX_PERCENT_IDENT]
40              [--min_alig_res_in_overlap MIN_ALIG_RES_IN_OVERLAP]
41              [--min_ident_res_in_overlap MIN_IDENT_RES_IN_OVERLAP]
42              [--min_sim_res_in_overlap MIN_SIM_RES_IN_OVERLAP]
43              [--min_ident_span_len MIN_IDENT_SPAN_LEN]
44              [--min_sim_span_len MIN_SIM_SPAN_LEN]
45              [--min_percent_ident_in_overlap MIN_PERCENT_IDENT_IN_OVERLAP]
46              [--min_percent_sim_in_overlap MIN_PERCENT_SIM_IN_OVERLAP]
47              [--min_percent_overlap MIN_PERCENT_OVERLAP] [--add_ali_col]
```

```
1             csv_file

2

3  Identify hit sequences likely encoded by the same gene loci in the genome of a
4  given species, or otherwise not representing paralogous genes. Criteria are
5  applied in this order: 1. Peptide hits with the same ID as higher-ranking hits
6  for the same query (query title) are excluded. 2. Nucleotide hits for the same
7  loci as peptide sequence hits are excluded. 3. Sequences with internal stop
8  codons are excluded, as these are potentially pseudogenes. 4. Sequences are
9  excluded if they do not meet several minimum length criteria: Absolute minimum
10 length (in amino acids) and percent query cover. 5. Sequences are excluded if
11 they do not overlap to a specified degree with all included higher-ranking
12 hits for the same query (query title) in sequence data for the same
13 species/genome. This is determined by algorithmically comparing pairs of
14 sequences aligned to a reference alignment of homologues, and several minimum
15 measures of alignment overlap may be specified. 6. Secondary hit sequences are
16 excluded if they do not meet a specified maximum percent identity threshold.
17 Highly identical sequences may result from false segmental duplications in the
18 genome assembly, may represent alleles, etc. Note: Applying these criteria
19 requires a column to be manually added to the input csv file prior to running
20 with the header "Alignment for sequence comparison" and filled with the
21 appropriate alignment name to use (one for each query title, as listed in the
22 "Query title" column). Alternatively, you can run this command with the
23 --add_ali_col option to automatically identify appropriate alignments among
24 your aligned FASTA queries used for running HMMer searches.

25

26 positional arguments:
27   csv_file              Path to spreadsheet with interpreted search results
28                         outputted by the interp_srchs command.

29

30 optional arguments:
31   -h, --help            show this help message and exit
32   --out_csv_path OUT_CSV_PATH
33                         Optionally specify an output file path, so that this
34                         command can be piped together with others. (default:
35                         None)
36   --remove_tblastn_hits_at_annotated_loci
37                         Ignore tblastn hits that overlap with any previously
38                         annotated loci. The rationale for this would be that
39                         the corresponding protein sequences should have been
40                         retrieved if the tblastn hit were a true positive
41                         anyway. If this option is not specified, then
42                         sequences will still be excluded if they specifically
43                         correspond to the same loci as do higher-ranking hits.
44                         (default: False)
45   --just_look_for_genes_in_gff3
46                         When looking for records in GFF3 annotation files that
47                         overlap with subsequences identified by similarity
48                         searching (TBLASTN), ignore records that are not
49                         explicitly "gene" (for example, "CDS", "mRNA", and
```

```
                           "exon"). This option should probably not be selected,
                           because in some GFF3 annotation files do not include
                           "gene" records, but do include predicted coding
                           sequences for genes. (default: False)
  --ignore_gff3            Disregard any information regarding redundancy of
                           identified nucleotide sequences with identified
                           protein sequences that may be found in GFF3 annotation
                           files. (default: False)
  --allow_internal_stops ALLOW_INTERNAL_STOPS
                           Include sequences that have internal stop codons
                           (anywhere other than the N-terminal position).
                           (default: True)
  --min_length MIN_LENGTH
                           Absolute minimum length (in AA) of a hit sequence to
                           be considered a potential distinct paralogue.
                           (default: 55)
  --min_percent_length MIN_PERCENT_LENGTH
                           Minimum length (in AA) of a hit sequence as a
                           percentage of query length for the hit to be
                           considered a potential distinct paralogue. (default:
                           15)
  --min_percent_query_cover MIN_PERCENT_QUERY_COVER
                           Minimum number of residues aligning with the original
                           query as a percentage of the original query sequence
                           length. (default: 0)
  --overlap_required       True if hits must overlap with a higher-ranking hit to
                           be considered potential unique paralogues. (default:
                           False)
  --max_percent_ident MAX_PERCENT_IDENT
                           Maximum percent identity (among aligning residues) for
                           evaluating whether two sequences are redundant or not
                           (secondary hits showing a percent identity with a
                           higher-ranking hit exceeding this value will be
                           excluded). (default: 98.0)
  --min_alig_res_in_overlap MIN_ALIG_RES_IN_OVERLAP
                           Minimum number of residues which must align for two
                           sequences to be considered as potentially distinct
                           hits. This is only relevant if the overlap_required
                           option is specified. (default: 20)
  --min_ident_res_in_overlap MIN_IDENT_RES_IN_OVERLAP
                           Minimum number of aligning residues which must be
                           identical for two sequences to be considered as
                           potentially distinct hits. This is only relevant if
                           the overlap_required option is specified. (default:
                           10)
  --min_sim_res_in_overlap MIN_SIM_RES_IN_OVERLAP
                           Minimum number of aligning residues which must be
                           similar for two sequences to be considered as
                           potentially distinct hits. This is only relevant if
```

```
                        the overlap_required option is specified. (default:
                        15)
--min_ident_span_len MIN_IDENT_SPAN_LEN
                        Minimum number of aligning residues which are
                        identical that must exist in at least one continuous
                        span for two sequences to be considered as potentially
                        distinct hits (not counting positions where both
                        sequences have gaps). This is only relevant if the
                        overlap_required option is specified. (default: 0)
--min_sim_span_len MIN_SIM_SPAN_LEN
                        Minimum number of aligning residues which are similar
                        (or identical) that must exist in at least one
                        continuous span for two sequences to be considered as
                        potentially distinct hits (not counting positions
                        where both sequences have gaps). This is only relevant
                        if the overlap_required option is specified. (default:
                        0)
--min_percent_ident_in_overlap MIN_PERCENT_IDENT_IN_OVERLAP
                        Minimum percent identity between the two sequences of
                        interest in the alignment.This is only relevant if the
                        overlap_required option is specified. (default: 0)
--min_percent_sim_in_overlap MIN_PERCENT_SIM_IN_OVERLAP
                        Minimum percent similarity (including identity)
                        between the two sequences of interest in the
                        alignment.This is only relevant if the
                        overlap_required option is specified. (default: 0)
--min_percent_overlap MIN_PERCENT_OVERLAP
                        Minimum number of aligning residues between the two
                        sequences of interest as a percentage of the length of
                        the second sequence (the last sequence in the
                        alignment), not including gaps, for the two sequences
                        to be considered as potentially distinct hits. This is
                        only relevant if the overlap_required option is
                        specified. (default: 0)
--add_ali_col         Add a column to the csv file listing which alignment
                        file in the queries directory to use for comparing
                        sequences. Aligned FASTA queries are selected that
                        match the query titles of the original queries used to
                        retrieve each of the relevant hits listed in the csv
                        file. No other options need to be specified in this
                        case. (default: False)
```

3.17   amoebae plot

```
usage: amoebae [-h] [--csv_file2 CSV_FILE2] [--complex_info COMPLEX_INFO]
               [--row_order ROW_ORDER] [--out_pdf OUT_PDF]
               csv_file
```

Plot results of similarity search and sequence classification analyses. The

```
1   outputs are PDF files.
2
3   positional arguments:
4     csv_file             Path to a spreadsheet with the relevant results to be
5                          plotted. This can be either a CSV file output of the
6                          sum_rev_srch command or from the find_redun_seqs
7                          command. If the output of the sum_rev_srch command is
8                          used, however, redundant hits will be counted (e.g.,
9                          BLASTP and TBLASTN hits corresponding to the same or
10                         highly identical genomic loci).
11
12  optional arguments:
13    -h, --help           show this help message and exit
14    --csv_file2 CSV_FILE2
15                         Path to a second spreadsheet with relevant results to
16                         be compared to the first and plotted. (default: None)
17    --complex_info COMPLEX_INFO
18                         Path to file that specifies which query titles
19                         represent components of which protein complexes (or
20                         otherwise grouped proteins). (default: None)
21    --row_order ROW_ORDER
22                         Path to file that specifies the order in which data
23                         for each species will be displayed. (default: None)
24    --out_pdf OUT_PDF    Path to output pdf file. (default: None)
```

3.18   amoebae add_to_models

```
26  usage: amoebae [-h]
27                 model_name alignment tree_topology subs_model type_seqs taxon
28
29  Add a phylogenetic model for relationships between members of a gene family
30  (sequence_data matrix, data type, tree topology, type sequence defining each
31  clade of interest, and substitution model) to a directory for use in
32  classifying sequence (via the 'phylo_class' command.
33
34  positional arguments:
35    model_name     An arbitrary name for the model (which will refer to the
36                   alignment, tree, substitution model, etc. collectively).
37    alignment      A multiple amino acid sequence alignment in nexus format.
38    tree_topology  Text file containing a tree (identified previously using
39                   MrBayes, etc) containing the names of all the sequences in
40                   the alignment, in newick format.
41    subs_model     The name of the substitution model used to recover the
42                   provided topology (chosen with ModelFinder or similar
43                   software).
44    type_seqs      Names of sequences (sequence headers) that are to be used to
45                   define clades of interest. A csv file with seq names in one
46                   column and clade names in the next column.
47    taxon          Taxonomic group represented in the model (e.g., "Eukaryotes",
```

21

```
1                    or "Amorphea").
2
3  optional arguments:
4    -h, --help      show this help message and exit
```

3.19   amoebae list_models

```
6  usage: amoebae [-h]
7
8  Print a list of all usable model/reference tree names in the models directory
9  as defined in the settings file.
10
11  optional arguments:
12    -h, --help  show this help message and exit
```

3.20   amoebae get_alt_topos

```
14  usage: amoebae [-h] [--polytomy] [--not_polytomy_clades]
15                 [--keep_original_backbone] [--iqtree_au_test]
16                 model_name out_dir_path
17
18  Take a tree and make copies with every alternative topology for the branches
19  connecting the clades of interest. Output as additional models in the Models
20  directory.
21
22  positional arguments:
23    model_name              Name of model/backbone tree to modify (other info
24                            provided in the model info csv file).
25    out_dir_path            Path to directory in which output directory will be
26                            written.
27
28  optional arguments:
29    -h, --help              show this help message and exit
30    --polytomy              Just make one big polytomy connecting the clades of
31                            interest intead of making alternative bifurcating
32                            trees. (default: False)
33    --not_polytomy_clades
34                            Do not make subtrees/clades of interest polytomies in
35                            output topologies. (default: False)
36    --keep_original_backbone
37                            Keep the original backbone topology instead of
38                            generating a polytomy or alternative resolved
39                            topologies. (default: False)
40    --iqtree_au_test        Test all the relevant alternative topologies against
41                            each other using Approximately Unbiased (AU) test with
42                            IQ-tree. (default: False)
```

3.21 amoebae prune

```
usage: amoebae [-h] [--include_seqs] [--output_file OUTPUT_FILE]
               tree_file alignment name_replace_table

Identify sequences in a tree, and remove them from a given alignment for
further phylogenetic analysis.

positional arguments:
  tree_file             Tree in newick format (coded names, because ETE3
                        cannot parse taxon names with space characters without
                        quotation marks around them).
  alignment             Dataset used to make the tree (nexus alignment)
                        (original alignment with original taxon names either
                        trimmed or untrimmed).
  name_replace_table    File for decoding names in input tree file.

optional arguments:
  -h, --help            show this help message and exit
  --include_seqs        Include only listed sequences/nodes instead of
                        removing them. (default: False)
  --output_file OUTPUT_FILE
                        Path to output file. (default: None)
```

3.22 amoebae auto_prune

```
usage: amoebae [-h]
               [--max_bl_iqr_above_third_quartile MAX_BL_IQR_ABOVE_THIRD_QUARTILE]
               [--remove_redun_seqs REMOVE_REDUN_SEQS]
               [--remove_redun_seqs_threshold REMOVE_REDUN_SEQS_THRESHOLD]
               [--output_file OUTPUT_FILE]
               in_dir

Automatically identify sequences in a tree, and remove them from a given
alignment for further phylogenetic analysis.

positional arguments:
  in_dir                Path to directory that contains the phylogenetic
                        analysis output files (sequence name conversion table
                        file and original nexus alignment file can be in the
                        parent directory to this directory as long as their
                        names are mostly identical.

optional arguments:
  -h, --help            show this help message and exit
  --max_bl_iqr_above_third_quartile MAX_BL_IQR_ABOVE_THIRD_QUARTILE
                        Inclusion threshold for number of interquartile ranges
                        above the third quartile of terminal branch lengths
                        the length of a terminal branch can be before it is
```

```
                            considered an outlier (length is total distance from
                            root node after rooting on midpoint, or the longest
                            terminal branch on either side of the midpoint).
                            (default: 1.5)
  --remove_redun_seqs REMOVE_REDUN_SEQS
                            Remove taxonomically redundant sequences (longest
                            branch of two sister branches when both are sequences
                            from the same species. (default: True)
  --remove_redun_seqs_threshold REMOVE_REDUN_SEQS_THRESHOLD
                            Minimum support required to consider one of two sister
                            branches/sequences taxonomically redundant. Note: only
                            used if the remove_redun_seqs option is specified.
                            (default: 0.95)
  --output_file OUTPUT_FILE
                            Path to output file. (default: None)
```

3.23   amoebae reduce_tree

```
usage: amoebae [-h] [--output_file OUTPUT_FILE] alignment tree_file
```

Remove terminal nodes from a given tree if there are not any sequences with
the same name in a given alignment.

```
positional arguments:
  alignment             Alignment in nexus format with sequences representing
                        a subset of those represented in the input tree.
  tree_file             Tree in newick format.

optional arguments:
  -h, --help            show this help message and exit
  --output_file OUTPUT_FILE
                        Path to output file. (default: None)
```

3.24   amoebae constrain_mb

```
usage: amoebae [-h] [--out_alignment OUT_ALIGNMENT] alignment tree
```

Add constraint commands to MrBayes input file.

```
positional arguments:
  alignment             Nexus alignment for input to Mrbayes (without any
                        constraint commands).
  tree                  Tree in newick format with same taxon names as in
                        alignment. To be used as a topology constraint (all
                        nodes).

optional arguments:
  -h, --help            show this help message and exit
```

```
1  --out_alignment OUT_ALIGNMENT
2                         Path to nexus alignment for input to Mrbayes with
3                         constraints added. (default: None)
```

3.25   amoebae visualize_tree

```
5  usage: amoebae [-h] [--root_taxon ROOT_TAXON] [--highlight_paralogues]
6                 [--add_clade_names_from_file]
7                 input_directory method
8
9  Parse phylogenetic analysis output files in a given directory, and write
10 human-readable tree figures to PDF files.
11
12 positional arguments:
13   input_directory      Path to directory containing input files (must contain
14                        a .table file for decoding taxon names.
15   method               Name of tree searching program used. Either iqtree,
16                        raxml, or mrbayes accepted.
17
18 optional arguments:
19   -h, --help           show this help message and exit
20   --root_taxon ROOT_TAXON
21                        Name of species to root on (e.g.,
22                        "Klebsormidium_nitens").
23   --highlight_paralogues
24                        Highlight clades that contain paralogues found in at
25                        least one other clade in the tree.
26   --add_clade_names_from_file
27                        Use a file in the parent directory with clade names
28                        corresponding to representative sequences to add clade
29                        names to all the taxon names in the output trees.
```

3.26   amoebae replace_seqs

```
31 usage: amoebae [-h] [--fasta_file FASTA_FILE] alignment
32
33 Replace sequences in an alignment the full-length sequences from the relevant
34 file(s) in the Genomes directory, or with their top hits in a given fasta
35 file. And, align, mask, and trim the identified sequences to the input
36 alignment
37
38 positional arguments:
39   alignment            Path to multiple sequence alignment file in nexus
40                        format (trimmed alignment).
41
42 optional arguments:
43   -h, --help           show this help message and exit
44   --fasta_file FASTA_FILE
```

```
                        Path to file containing sequences with which to
                        replace sequences in the alignment. If this option is
                        not specified, then full-length sequences will be
                        retrieved from files in the Genomes directory.
```

3.27   amoebae csv_to_fasta

```
usage: amoebae [-h] [--output_dir OUTPUT_DIR] [--abbrev] [--paralogue_names]
               [--only_descr] [--subseq] [--all_hits] [--split_by_query_title]
               [--split_by_top_rev_srch_hit SPLIT_BY_TOP_REV_SRCH_HIT]
               csv_file

Extract sequences described in a spreadsheet output by AMOEBAE, and write to a
file in FASTA format.

positional arguments:
  csv_file              Path to csv file listing sequences.

optional arguments:
  -h, --help            show this help message and exit
  --output_dir OUTPUT_DIR
                        Path for output directory to contain FASTA files.
                        (default: None)
  --abbrev              Add species name instead of sequence description from
                        fasta header. Applicable when the output file is to be
                        used for alignment and phylogenetic analysis.
                        (default: False)
  --paralogue_names     Use species name, query title, and paralogue number
                        instead of sequence description from fasta header.
                        Applicable when the output file is to be used for
                        alignment and phylogenetic analysis. Does not work if
                        the abbrev option is specified. (default: False)
  --only_descr          Use the description but not the ID as the new fasta
                        sequence header. Does not work if the abbrev option is
                        specified. (default: False)
  --subseq              Write subsequences that aligned to forward search
                        query, instead of the full sequences. (default: False)
  --all_hits            Write all forward hits listed in the input csv file.
                        (default: False)
  --split_by_query_title
                        Write sequences to files according to the query title
                        of the query which retrieved them in a similarity
                        search. (default: False)
  --split_by_top_rev_srch_hit SPLIT_BY_TOP_REV_SRCH_HIT
                        Write sequences to files according to the top hit that
                        they retrieve in a reverse search, for each sequence
                        that meets the reverse search criteria. (Provide the
                        reverse search identifier, eg,
                        "rev_srch_20180924122402-1") (default: None)
```

3.28 amoebae check_depend

```
usage: amoebae [-h]

Check that all the dependencies (other than python modules) are properly
installed and useable.

optional arguments:
  -h, --help  show this help message and exit
```

3.29 amoebae check_imports

```
usage: amoebae [-h]

Check that all the import statements used in the AMOEBAE repository run
without error.

optional arguments:
  -h, --help  show this help message and exit
```

4 Miscellaneous scripts

Several scripts of less general applicablity than the amoebae commands descibed above are included in the AMOEBAE toolkit. See the amoebae/misc_scripts directory (`https://github.com/laelbarlow/amoebae/tree/master/misc_scripts`). Most scripts have information regarding usage in the files themselves. More detailed information regarding some of these scripts may be added to this documentation in the future.

5 References

Camacho, C., Coulouris, G., Avagyan, V., Ma, N., Papadopoulos, J., Bealer, K., and Madden, T.L. (2009). BLAST+: Architecture and applications. *BMC Bioinformatics*, 10(1):421. doi:10.1186/1471-2105-10-421.

Cock, P.J.A., Antao, T., Chang, J.T., Chapman, B.A., Cox, C.J., Dalke, A., Friedberg, I., Hamelryck, T., Kauff, F., Wilczynski, B., and de Hoon, M.J.L. (2009). Biopython: Freely available Python tools for computational molecular biology and bioinformatics. *Bioinformatics*, 25(11):1422–1423. doi:10.1093/bioinformatics/btp163.

Eddy, S.R. (1998). Profile hidden Markov models. *Bioinformatics*, 14(9):755–763. doi:10.1093/bioinformatics/14.9.755.

Edgar, R.C. (2004). MUSCLE: Multiple sequence alignment with high accuracy and high throughput. *Nucleic Acids Research*, 32(5):1792–1797. doi:10.1093/nar/gkh340.

Huerta-Cepas, J., Serra, F., and Bork, P. (2016). ETE 3: Reconstruction, Analysis, and Visualization of Phylogenomic Data. *Molecular Biology and Evolution*, 33(6):1635–1638. doi:10.1093/molbev/msw046.

Hunter, J.D. (2007). Matplotlib: A 2D Graphics Environment. *Computing in Science Engineering*, 9(3):90–95. doi:10.1109/MCSE.2007.55.

Larson, R.T., Dacks, J.B., and Barlow, L.D. (2019). Recent gene duplications dominate evolutionary dynamics of adaptor protein complex subunits in embryophytes. *Traffic*, page tra.12698. doi:10.1111/tra.12698.

Li, L. (2003). OrthoMCL: Identification of Ortholog Groups for Eukaryotic Genomes. *Genome Research*, 13(9):2178–2189. doi:10.1101/gr.1224503.

Nguyen, L.T., Schmidt, H.A., von Haeseler, A., and Minh, B.Q. (2015). IQ-TREE: A Fast and Effective Stochastic Algorithm for Estimating Maximum-Likelihood Phylogenies. *Molecular Biology and Evolution*, 32(1):268–274. doi:10.1093/molbev/msu300.

Slater, G. and Birney, E. (2005). [No title found]. *BMC Bioinformatics*, 6(1):31. doi:10.1186/1471-2105-6-31.