

# VIBRANT

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## Virus Identification By iteRative ANnoTation

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## Current Version

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VIBRANT v1.2.1

## Citation

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If you find VIBRANT useful please consider citing our manuscript in [Microbiome](#):

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## Content Addition (June 22 2020):

- The script `simplify_faa-ffn.py` was added to the `scripts/` folder. This script can be used to remove excess annotation information from the definition lines of VIBRANT phage protein (.faa) and gene (.ffn) output files. Specifically, this converts protein/gene outputs from VIBRANT's tab separated format to the standard format generated by other software (e.g., Prodigal). An example implementation of this script would be for using VIBRANT phage protein results for vConTACT2 taxonomy. Usage of this script: `python3 simplify_faa-ffn.py example.phages_combined.faa` . Replace the example .faa file with any VIBRANT output .faa or .ffn file of your choice.

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## Updates for v1.2.1 (Mar 13 2020):

### Summary:

- Note: predictions are identical to v1.2.0 and all data are 100% compatible (i.e., no change to true or false positive rate).
- Note: new version push to bioconda install in order to fix bug effecting some users (see issue below).
- Issue: resolved `name 'pfam_name' is not defined` issue from v1.2.0 causing some parallel runs to exit and data to be lost. This issue can also be identified by the presence of a file beginning with `temp2_` .
- Update: new output `phages_circular.fna` containing circular predicted viral sequences.
- Update: new output `integrated_prophage_coordinates.tsv` containing the scaffold coordinate locations of excised proviruses.
- Update: new figure `figure_PCA.pdf` displaying a summary PCA plot of predicted viruses (qualities, lifestyles, linear/circular, sizes and relationships).
- Update: new output `figure_PCA.tsv` containing the (x,y) coordinates of each predicted virus on the PCA plot.
- Update: new output `summary_normalized.tsv` is the normalized (to number of ORFs) version of `summary_results.tsv` and the information read into the neural network for classification. This is also the file used to generate the PCA plot.
- Update: new outputs `log_run.log` and `log_annotation.log` . The file `log_run.log` represents the original log file and contains summary information from the run (command, runtime, date, virus identifications, etc.). Errors that cause VIBRANT to exit will be written to `log_run.log` or `log_annotation.log` depending on the origin of the error.

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## Updates for v1.2.0 (Feb 9 2020):

### Summary:

- Update: pre-filtration of scaffolds based on strand switching was completely removed. This modification was found to increase virus identification with a negligible effect on the rate of false positive discovery.
- Note: two of the HMM databases are no longer required (Pfam-A\_phage and Pfam-A\_plasmid). You may want to remove these. There is no need to re-download or re-compile the other databases, though you can delete all databases and start over.
- Note: results of virus identification may increase compared to previous versions.

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### Updates for v1.1.0 (Feb 7 2020):

### Summary:

- Issue: incorrect storage of strand switching information caused inconsistent results and potential decrease in virus recovery. Issue is now resolved.
- Update: modification of `hmmsearch` command to increase speed if using a newer version (version  $\geq$  3.2.1).
- Update: addition of `-folder` flag and compression of databases (`-d`) and files flags (`-m`).
- Update: modification of `VIBRANT_setup.py` messages and removal of `VIBRANT_test_setup.py`.
- Issue: annotations with PF12761.7 caused issues.
- Note: there is no need to re-download or re-compile the databases.
- Note: results of virus identification may vary compared to v1.0.1.
- Note: please verify Scikit-Learn and numpy versions. Incorrect versions may cause inconsistent and invalid results.

### Explanations:

- Fixed a bug that was causing strand switching information to be stored with the wrong scaffolds. This caused scaffolds to undergo potentially incorrect pre-filtering (see flowchart and methods). Due to this issue, viral scaffolds may have been filtered out accidentally. This also led to inconsistent results between identical runs of VIBRANT because the same scaffold was given various strand switching information depending on the run. Both issues of incorrect strand switching and inconsistent results have been resolved.
- The usage of `hmmsearch --cpu` changed with newer versions of HMMER (version  $\geq$  3.2.1). For older versions the command used by VIBRANT is `--cpu 1` whereas newer versions `--cpu 0` is used. VIBRANT will automatically detect your version and use the appropriate command. This increases speed by optimizing cpu usage.
- The new `-folder` flag allows for the designation of an output directory to deposit the final VIBRANT output as well as all temporary files. When invoked this flag will create the new directory if it does not exist or add to the directory if it already exists. This may especially be useful when running VIBRANT on a shared computing cluster. The new `-d` and `-m` flags are useful if you have moved the location of the `databases/` and/or `files/` directories from their default location. Previously this required several flags that have now been condensed to two.
- The new `VIBRANT_setup.py` script generates additional messages for version control and it now does not require `VIBRANT_test_setup.py`. There is no need to re-run this script if you have ran it with a previous version. If you want to test dependencies/versions use `VIBRANT_setup.py -test`. It is suggested to run `VIBRANT_setup.py -test` to quickly verify correct setup.
- PF12761.7 was not given a name in `VIBRANT_names.tsv` and would cause errors if a protein was annotated with PF12761.7. This issue has been resolved.

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### Updates for v1.0.1:

#### Summary:

- Issue: some extracted proviruses were given the wrong genome sequence. Issue is now fixed.
- Update: minimum sequence length is now 1000bp which greatly increases virus identifications from some metagenomes.
- Update: metrics for quality analysis of scaffolds has been changed.
- Note: there is no need to re-download or re-compile the databases.

#### Explanations:

- Fixed a bug that was causing issues with extracting the correct genomic region of an integrated lysogenic virus (provirus). Briefly, the issue was causing some extracted proviruses (those with "`_fragment_#`" in the name) to have either the wrong genomic sequence or a sequence length of zero. The respective proteins and virus identification metrics for these scaffolds remains identical.
- The minimum scaffold length is now set to 1000bp (previous 3000bp). This greatly increases total virus identification within metagenomes that contain many sequences less than 3kb. This has no effect on false discovery since the minimum open reading frame requirement is still 4. For example, a 1kb and 3kb scaffold each encoding 4 open reading frames will be considered identically since neither sequence length nor sequence features impact virus identification.
- The quality analysis of scaffolds was updated. The category *fragment quality draft* was removed. The categories are now *complete circular*, *high*, *medium* and *low quality draft*. The update was made to better represent the completeness of

*Caudovirales* scaffolds which are likely to be the most abundant viral population in a metagenome. Baseline tests indicate the new metrics linearly represent the completeness of *Caudovirales* genomes.

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## Program Description

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VIBRANT is a tool for automated recovery and annotation of bacterial and archaeal viruses, determination of genome completeness, and characterization of viral community function from metagenomic assemblies. VIBRANT uses neural networks of protein annotation signatures and genomic features to maximize identification of highly diverse partial or complete viral genomes as well as excise integrated proviruses.

- Uses neural network machine learning of protein annotation signatures
- Assigns novel 'v-score' for determining the virus-like nature of all annotations
- Determines genome completeness
- Characterizes viral community function by metabolic analysis
- Identifies auxiliary metabolic genes (AMGs)
- Excises integrated viral genomes from host scaffolds
- Performs well in diverse environments
- Recovers novel and abundant viral genomes
- Built for dsDNA, ssDNA and RNA viruses

VIBRANT uses three databases for identifying viruses and characterizing virome metabolic potential:

- KEGG (March release): <https://www.genome.jp/kegg/> (FTP: <ftp://ftp.genome.jp/pub/db/kofam/archives/2019-03-20/>)
- Pfam (v32): <https://pfam.xfam.org> (FTP: <ftp://ftp.ebi.ac.uk/pub/databases/Pfam/releases/Pfam32.0/>)
- VOG (release 94): <http://vogdb.org/> (FTP: <http://files.share.csb.univie.ac.at/vog/vog94/>)

## Requirements

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*System Requirements:* VIBRANT has been tested and successfully run on Mac, Linux and Ubuntu systems.

*Program Dependencies:* Python3, Prodigal, HMMER3, gzip, tar, wget (see section below)

*Python Dependencies:* BioPython, Pandas, Matplotlib, Seaborn, Numpy, Scikit-learn, Pickle (see section below)

Don't worry, these should all be easy and quick to install if you do not already have the requirements satisfied. Suggested methods of installation are with [1] pip (pip3) (<https://pypi.org/project/pip/>), [2] conda (<https://anaconda.org/anaconda/conda>), [3] homebrew (<https://brew.sh/>) or [4] apt (you may need to use 'apt-get' or 'sudo') (<https://help.ubuntu.com/its/serverguide/apt.html>). The method will depend on your operating system and setup.

### Program Dependencies: Installation

Please ensure the following programs are installed and in your machine's PATH. Note: most downloads will automatically place these programs in your PATH.

#### Programs:

1. Python3: <https://www.python.org> (version  $\geq 3.5$ )
2. Prodigal: <https://github.com/hyatt/Prodigal>
3. HMMER3: <https://github.com/EddyRivasLab/hmmer>
4. gzip: <http://www.gzip.org/>
5. tar: <https://www.gnu.org/software/tar/>
6. wget: <https://www.gnu.org/software/wget/>

#### Example Installations:

1. Python3: see Python webpage. You can check your current version using `python --version`. Required version  $\geq 3.5$ .

2. Prodigal: `conda install -c bioconda prodigal` or `brew install prodigal` or `apt-get install prodigal` or GitHub clone
3. HMMER3: `conda install -c bioconda hmmer` or `brew install hmmer` or `apt install hmmer` or GitHub clone
4. gzip: you likely already have this installed
5. tar: you likely already have this installed
6. wget: you likely already have this installed or `brew install wget` or `apt install wget` or `pip install wget`

## Python3 Dependencies: Installation

There are several Python3 dependencies that must be installed as well. You may already have some of these installed.

### Packages

1. BioPython: <https://biopython.org/wiki/Download>
2. Pandas: <https://pandas.pydata.org/pandas-docs/stable/install.html>
3. Matplotlib: <https://matplotlib.org/>
4. Seaborn: <https://seaborn.pydata.org/>
5. Numpy: <https://numpy.org/> (version  $\geq 1.17.0$ )
6. Scikit-learn: <https://scikit-learn.org/stable/> (version  $= 0.21.3$ )
7. Pickle: <https://docs.python.org/3/library/pickle.html>

### Example Installations:

1. BioPython: `pip install biopython` or `apt-get install python-biopython` or `conda install -c conda-forge biopython`
2. Pandas: `pip install pandas` or `conda install -c anaconda pandas` or `apt-get install python-pip`
3. Matplotlib: `pip install matplotlib` or `conda install matplotlib` or `apt-get install python-matplotlib`
4. Seaborn: `pip install seaborn` or `conda install seaborn`
5. Numpy: `pip install numpy` (`pip install --upgrade numpy==1.17.0`) or `conda install -c anaconda numpy` or `apt-get install python-numpy`
6. Scikit-learn: `pip install --upgrade scikit-learn==0.21.3` or `conda install scikit-learn`
7. Pickle: this should already come with Python3. If you have issues try `pip install pickle-mixin`

## Running VIBRANT

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VIBRANT is built for efficiently running on metagenomes but can also run on individual or small groups of genomes. Each scaffold is considered individually, so results will *not* vary whether the scaffold is run as part of a metagenome or by itself.

### Quick Start

There are two different routes to downloading VIBRANT: [Anaconda](#) install or [GitHub](#) clone/download. Alternatively you can use the [CyVerse Discovery Environment](#) open source web server.

### CyVerse (currently running v1.0.1)

- VIBRANT app: <https://de.cyverse.org/de/?type=apps&app-id=c2864d3c-fd03-11e9-9cf4-008cfa5ae621&system-id=de>
- Information: <https://wiki.cyverse.org/wiki/display/DEapps/VIBRANT-1.0.1>

### Anaconda (currently running v1.2.0)

- 1) Install dependencies. See *Requirements* section above.
- 2) Install directly to \$PATH using bioconda.

```
conda install -c bioconda vibrant==1.2.0
```

- 3) Download and setup databases. This will take some time due to file sizes, but it only needs to be run once. This step requires ~20GB of temporary storage space and ~11GB of final storage space. To do this, run `download-db.sh` which should be in your

system's \$PATH.

```
download-db.sh
```

## GitHub (currently running v1.2.1)

*Note:* if at any time you are given a "permission denied" error you can run `chmod 777 <file_name>` or `chmod -R 777 <folder_name>`. Simply replace `<file_name>` or `<folder_name>` with the file/folder that you would like to add permissions to.

1) Install dependencies. See *Requirements* section above.

2) Download VIBRANT using git clone or download zip file. *Note:* if you download the zip file you will have the parent folder `VIBRANT-master` instead of `VIBRANT`.

```
git clone https://github.com/AnantharamanLab/VIBRANT
```

3) You may want to add permissions to all files.

```
chmod -R 777 VIBRANT
```

4) Move parent folder ( `VIBRANT` ) to desired location. VIBRANT will function no matter where the parent folder is located or moved to, but hierarchy of files and folders within the parent folder must remain constant.

5) Move into databases folder for setup.

```
cd VIBRANT/databases
```

6) Download and setup databases. This will take some time due to file sizes, but it only needs to be run once. This step requires 20GB of temporary storage space and ~11.2GB of final storage space. Run the following:

```
python3 VIBRANT_setup.py or ./VIBRANT_setup.py
```

7) This step is included in step 6. This can be used if you have already completed step 6 and want to re-test the setup.

VIBRANT can automatically verify that downloads, setup and installation of dependencies was completed properly. You may choose to skip this, but it's very quick. This flag can be called as many times as necessary to run verifications.

```
python3 VIBRANT_setup.py -test or ./VIBRANT_setup.py -test
```

## Testing VIBRANT

*Note:* VIBRANT does not write to standard out (command prompt screen) while running or when it finishes (i.e., not verbose). However, VIBRANT will write to standard out in the event that it encounters an error, such as incorrect use of optional arguments or incorrect input file format. See step 11 below for an example.

*Anaconda users:* these example datasets are not made available through Anaconda install. Steps 8-11 may be skipped or the files may be downloaded from the [GitHub site](#).

*Optional:* run VIBRANT on test datasets (steps 8-11). *Note:* you may want to try using the `-t` flag to increase VIBRANT's speed. See *Arguments and Flags* section for details.

8) Test out a small dataset of mixed viral and non-viral scaffolds in nucleotide format. See

`example_output/VIBRANT_mixed_example` for how the results should look.

```
python3 ../VIBRANT_run.py -i mixed_example.fasta from within the example_data/ folder
```

or

```
python3 VIBRANT_run.py -i example_data/mixed_example.fasta from within the parent folder
```

9) Test out a single viral scaffold in protein format.

```
python3 ../VIBRANT_run.py -i Microviridae_MH552510.2.faa -f prot from within the example_data/ folder
```

10) Test out a set of non-viral scaffolds in nucleotide format.

```
python3 ../VIBRANT_run.py -i no_phages.fna from within the example_data/ folder
```

11) Test out a scaffold in nucleotide format that does not meet the minimum requirement of 1kb in length. VIBRANT will exit because the input sequence is not 1kb. *Note:* Exiting only occurs if *all* the scaffolds do not meet the minimum requirements.

```
python3 ../VIBRANT_run.py -i short_scaffold.fsa from within the example_data/ folder
```

## Arguments and Flags

The only required flag is the input ( `-i` ) FASTA file which can either be in nucleotide or protein format. The file extension (e.g., fasta, fna, faa, fsa) does not matter. For nucleotide input, the definition lines can be in any format. The only exception is that the phrase "fragment" should not appear in the definition line because this term is used by VIBRANT during analysis and output parsing. For protein input, proteins must be grouped by genome and in descending order by protein number. Also, the definition lines for proteins must be in Prodigal format. Please see `example_data/Microviridae_MH552510.2.faa` for an example of Prodigal format definition lines. The required items are the scaffold name, protein number, start site, end site and strand. It is suggested to use nucleotide input unless inputting proteins will fit best with your current/downstream analyses.

VIBRANT comes with a couple of very simple optional arguments. At any point you can see the help menu with `VIBRANT_run.py -h`.

#### *Common optional arguments*

- `-t` : increase the number of VIBRANT parallel runs (similar to threads). The integer entered here will have no impact on results but may impact runtime. To parallelize VIBRANT, the `-t` flag will indicate how many separate files to split the input file into; these will all be run in tandem. For example, an input file with 10 scaffolds that has invoked `-t 5` will run VIBRANT 5 separate times simultaneously with each run consisting of 2 scaffolds each. Default = `1`.
- `-f` : identify either nucleotide or protein input. This flag is only required when inputting proteins ( `-f prot` ) but can be added with any nucleotide input ( `-f nucl` ). Default = `nucl`.
- `-folder` : specify the directory in which to deposit VIBRANT's output folder and temporary files. If the specified directory doesn't exist then it will be created. Default = your current working directory.

#### *Uncommon optional arguments*

- `-l` : increase the minimum scaffold length requirement. The minimum is 1000 basepairs ( `-l 1000` ). Default = `1000`. For example, if `-l 5000` is invoked VIBRANT will only consider scaffolds greater than or equal to 5000bp.
- `-o` : increase the minimum number of open readings frames (ORFs, or proteins) per scaffold requirement. The minimum is 4 ORFs ( `-o 4` ). Default = `4`. For example, if `-o 8` is invoked VIBRANT will only consider scaffolds encoding at least 8 proteins.
- `-virome` : This flag should be used cautiously. This will edit VIBRANT's sensitivity if the input dataset is a virome and not mixed metagenome. That is, if you expect the vast majority of your input scaffolds to be viruses then `-virome` can be used to remove obvious non-viral scaffolds. This will have no effect on runtime. Default = off.
- `-no_plot` : This flag can be used to skip generation of the graphical representations of the VIBRANT's results. You may wish to use this flag if you simply have no use for the output figures or if for some reason this function of VIBRANT is causing the program to break. This will have little effect on runtime. Default = off.

#### *Likely unused optional arguments*

- `-d` : specify the location of the `databases/` directory if moved from its default location.
- `-m` : specify the location of the `files/` directory if moved from its default location.
- For `-d` and `-m` please specify the full path to the new location of the files. For example, `-d new_location/databases/`.

## Output Explanations

VIBRANT outputs a lot of files and folders. Please see `output_explanations.pdf` within the parent folder for information regarding each file/folder that is generated by VIBRANT. Each file and folder will have a prefix or suffix respective to the name of the input file.

*Note:* some scaffolds will be proviruses that have been extracted from a host scaffold. These viral sequences will be given a new name. Specifically, the term "`_fragment_#`" will be appended to the name to indicate that it is a fragment of a larger scaffold. The number associated with the fragment is, for the most part, arbitrary.

## Useful Outputs

- FASTA file of identified virus genomes: `VIBRANT_phages_<input_file>/<input_file>.phages_combined.fna`

- List of identified virus genomes: `VIBRANT_phages_<input_file>/<input_file>.phages_combined.txt`
- GenBank file of identified virus genomes (if `-f nucl`):  
`VIBRANT_phages_<input_file>/<input_file>.phages_combined.gbk`

Note: integrated viruses that have been excised from a scaffold will have 'fragment\_#' appended to the genome and protein names.

## General Overview

Briefly, the folder `VIBRANT_phages_<input_file>` will contain FASTA, GenBank and list files for the identified viruses; the folder `VIBRANT_results_<input_file>` will contain annotation, metabolic and summary information for the identified viruses; the folders `VIBRANT_HMM_tables_parsed_<input_file>` and `VIBRANT_HMM_tables_unformatted_<input_file>` will contain raw HMM tables used for analyses; the folder `VIBRANT_figures_<input_file>` will contain summary figures for the dataset and identified viruses; the file `VIBRANT_log_<input_file>` will contain the log summary and run information. All outputs will be contained within a folder named `VIBRANT_<input_file>`.

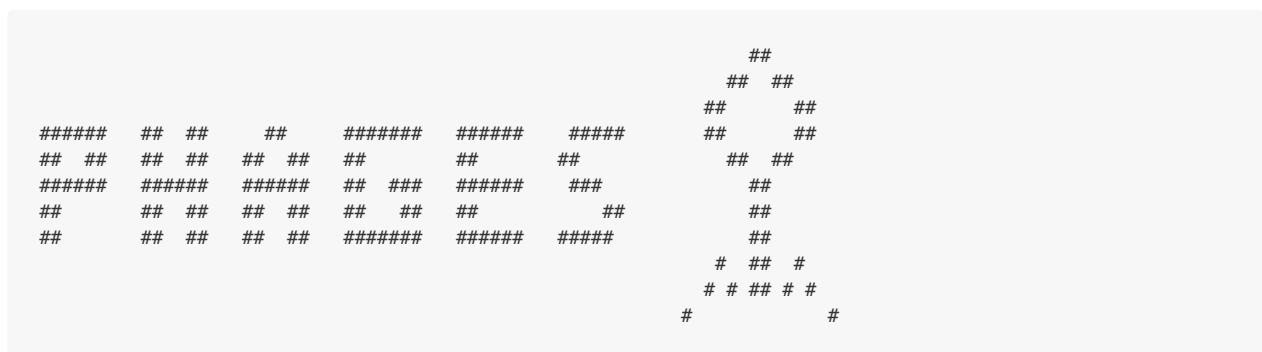
## VIBRANT Files and Folders

VIBRANT comes with several folders, files and scripts that are used during analysis. You will not need to interact with any of these, but knowing what they contain may be useful. The folder `databases` contains `VIBRANT_setup.py` that is only used during initial setup of VIBRANT. There is also a folder `profile_names` that contains text files with lists of all HMM profiles used per database. The folder `example_data` has a standard setup for testing VIBRANT and validating the correct outputs. The folder `files` contains several very useful documents: `VIBRANT_AMGs.tsv` - list of all KEGG KOs designated as auxiliary metabolic genes (AMGs); `VIBRANT_categories.tsv` - list of accession numbers for VOG, KEGG and Pfam with their respective "v-score" which is an essential metric used to identify viruses; `VIBRANT_KEGG_pathways_summary.tsv` - list of KEGG map pathways, and their associated metabolic and KO information which is used to summarize virome metabolism; `VIBRANT_machine_model.sav` - the neural network model used for virus classification; `VIBRANT_names.tsv` - list of names associated with each VOG, KEGG and Pfam accession number. The folder `scripts` contains three auxiliary scripts used to run VIBRANT. The script `VIBRANT_run.py` within the parent folder is actually a wrapper script to facilitate parallelization of VIBRANT and perform final metric analyses, whereas the script `VIBRANT_annotation.py` is what does the real virus identification. The two scripts `VIBRANT_extract_nucleotide.py` and `VIBRANT_extract_protein.py` are used when splitting the input file for parallelization. A fourth script `simplify_faa-ffn.py` can be used to remove all annotation information from the definition lines of protein (.faa) or gene (.ffn) output files.

## Contact

Please contact Kristopher Kieft (kieft@wisc.edu or GitHub Issues) with any questions, concerns or comments.

Thank you for using VIBRANT!



## Copyright



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