KLTPicker 2.0 – user guide (Python version)

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1 System requirements

Recommended Environments:

The package has been tested on Ubuntu 16.04 and Windows 10. It should probably work on other versions of Windows and Linux but has not been tested on them yet. Similarly, for macOS.

Python 3.6.0+ is required.

The package makes use of the pyfftw package, which in turn uses the FFTW library. Before installing KLTPicker make sure you have the FFTW library installed on your system: <http://www.fftw.org/fftw3_doc/Installation-and-Customization.html#Installation-and-Customization>

For optional GPU support, the package requires:

* NVIDIA CUDA GPU with the Compute Capability 3.0 or larger
* CUDA Toolkit: v9.0 / v9.2 / v10.0 / v10.1 / v10.2 / v11.0

2 Installing KLTPicker

2.1 Install KLTPicker via pip:

We recommend installing KLTPicker via pip.

For installation **without** GPU support run:

$ pip install kltpicker

For installation **with** GPU support (provided that your system satisfies the above requirements) run:

$ pip install kltpicker[gpu]

2.2 Install KLTPicker from source

The tarball of the source tree is available via:

$ pip download kltpicker

You can install KLTPicker from the tarball:

$ pip install kltpicker-x.x.x.tar.gz

You can also install the development version of KLTPicker from a cloned Git repository:

$ git clone https://github.com/ShkolniskyLab/kltpicker.git

$ cd kltpicker

$ pip install .

2.3 Uninstall KLTPicker

Use pip to uninstall KLTPicker:

$ pip uninstall kltpicker

2.4 Upgrade KLTPicker

Just use pip with -U option:

$ pip install -U kltpicker

3 Using the KLT picker

**Note**: after installing the KLT Picker, you need to open a new terminal and run the KLT Picker from there.

3.1 Interactive mode

To run the script in interactive mode simply run:

$ kltpicker

The following questions will appear, one by one

1. **Enter full path of micrographs MRC files**:

Type the full path to the directory which contains the micrograph MRC files.

1. **Enter the particle size in pixels**:

Type the particle diameter in pixels (more precisely, the diameter of the extracted box).

1. **Enter full path of output directory**:

Type the full path where the particles coordinate files will be saved.

If the output directory specified already contains coordinate files for some (but not all) of the micrographs in the input directory, the following question will appear:

* 1. **The directory you specified contains coordinate files for some of the micrographs in the input directory. Run only on micrographs which have no coordinate file? (Y/N):**

[Y] Type Y to pick particles only from micrographs for which no coordinate file exists in the specified output directory.

[N] If you type N, the program will abort.

1. **Pick all particles?** **(Y/N):**

[Y] Type Y to pick all particles using the optimal threshold derived on the paper.

[N] If you type N, then the following question will appear:

* 1. **How many particles to pick:**

Type the number of particles to pick in each micrograph.

1. **Pick noise images? (Y/N):**

[N] Type N if you don’t want to pick noise images.

[Y] If you type Y then the following question will appear:

* 1. **How many noise images to pick**?

Type the number of noise images to pick in each micrograph.

1. **Do you want to use ASOCEM for contamination removal? (Y/N):**

[N] Type N if you don’t want touse ASOCEM for contamination removal.

[Y] If you type Y then the following question will appear:

* 1. **Do you want to save ASOCEM masks? (Y/N)**?

Type Y to save for each micrograph a binary mask (.mrc) indicates the contaminated regions.

* 1. **Do you want to change ASOCEM default parameters (Y/N)?**

[N] Type N if you don’t want to change ASOCEM default parameters.

[Y] If you type Y then the following question will appear:

* + 1. **Enter ASOCEM downsample image size (should be a positive number)**Type the image size after downsampling (600 is the default parameter).
    2. **Enter ASOCEM covariance area size (should be a positive odd number)**Type covariance area sizeafter downsampling (5 is the default parameter).

1. **Display detailed progress? (Y/N):**

[N] Type N to display a simple progress-bar.

[Y] Type Y to display during runtime the number of particles and noise images picked from each micrograph.

1. **Enter maximum number of concurrent processes (-1 to let the program decide):**

Specify the maximum number of processes you want to allow the program to run in parallel. Enter -1 to let the program set the limit.

If you have installed the version that supports GPU, the following question will appear:

1. **Do you want to use the GPU? (Y/N)?**

[N] Type N to not use GPUs found on your system (CPU only).

[Y] Type Y to use the GPUs found on your system. If you type Y the following question will appear:

* 1. **Which GPUs would you like to use? (Valid indices: 0,...,7. Enter -1 to use all):**

Specify the indices of the GPUs on your system which you want to use, separated by whitespaces or commas. Type -1 to use all available GPUs.

3.2 Using flags to pass arguments

You can also run the program by passing arguments to the program using flags. To view a help message and the available flags run:

$ kltpicker -h

Usage:

$ kltpicker -i INPUT\_DIR -o OUTPUT\_DIR -s PARTICLE\_SIZE -p NUM\_PARTICLES -n NUM\_NOISE -a -–save-asocem –asocem-downsample ASOCEM\_DOWNSAMPLE –-asocem-area ASOCEM-AREA -v --verbose --max-processes MAX\_PROCESSES --gpus GPUS --no-gpu --only-do-unfinished

In red – flags that are only available in the version with GPU support.

Explanation of the arguments and flags:

|  |  |  |  |
| --- | --- | --- | --- |
| **Flag** | **Argument** | **Default** | **Optional** |
| -i, --input-dir | Full path of input directory. | (N/A) | No |
| -o, --output-dir | Full path of output directory. | (N/A) | No |
| -s, --particle-size | Expected size of particles in pixels. | (N/A) | No |
| -p, --num-particles | Number of particles to pick per micrograph.  Enter -1 to pick all particles. | -1 | Yes |
| -n, --num-noise | Number of noise images to pick per micrograph. | 0 | Yes |
| -a, --use-asocem | Choose to use ASOCEM to remove contaminations. | True | Yes |
| --save-asocem | Choose to save ASOCEM .mrc. | False | Yes |
| --asocem-downsample ASOCEM\_DOWNSAMPLE | Use to set ASOCEM downsample size. | 600 | Yes |
| --asocem-area ASOCEM\_AREA | Use to set ASOCEM covariance area size after downsampling. | 5 | Yes |
| -v, --verbose | Verbose. Choose this to display number of particles and noise images picked from each micrograph during runtime. Otherwise, you get a simple progress bar. | False | Yes |
| --max-processes | Limit the number of concurrent processes to run.  Enter -1 to let the program choose. | -1 | Yes |
| --only-do-unfinished | Only run on micrographs for which there are no coordinate files in the output directory. | False | Yes |
| --no-gpu | Don't use GPUs. | False | Yes |
| --gpus | Indices of GPUs to be used, **separated by whitespaces.**  Valid indices: zero through one less than the number of GPUs on the system.  Enter -1 to use all available GPUS. | -1 | Yes |

The KLT picker will then start running, while displaying progress notifications. The outputs are the coordinate files (box and star) and the contamination binary masks written to the output directory.

4 Examples

In this section we demonstrate the execution of the program on 5 micrographs of the EMPIAR-10028 data set (Plasmodium Falciparum 80S ribosome) [2] from the EMPIAR repository [1].

First, download the micrographs manually from <https://www.ebi.ac.uk/pdbe/emdb/empiar/entry/10028/>

For the sake of the example, assume you placed the downloaded files in a directory named /home/user/example/micrographs

Then proceed to run the program in interactive mode or by passing arguments using flags.

4.1 Interactive mode

In this section we demonstrate the execution of the program in interactive mode.

$ kltpicker

Enter full path of micrographs MRC files: /home/user/example/micrographs

Found 5 MRC files.

Enter the particle size in pixels: 300

Enter full path of output directory: /home/user/example/results

Pick all particles? (Y/N): y

Pick noise images? (Y/N): n

Do you want to use ASOCEM for contamination removal? (Y/N):y

Do you want to save ASOCEM masks? (Y/N):Y

Do you want to change ASOCEM default parameters? (Y/N):N

Display detailed progress? (Y/N): n

Enter maximum number of concurrent processes (-1 to let the program decide): -1

Use GPU? (Y/N): y

Which GPUs would you like to use?

(Valid indices: 0,...,7. Enter -1 to use all): 0 1 2 3

Running on 5 files.

Using GPUs 0, 1, 2, 3.

Preprocessing (usually takes up to 1 minute)...

Preprocess finished. Picking particles...

[Elapsed Time: 0:00:53] |#############################| (100%)

Finished successfully!

Picked \*\*\* particles and 0 noise images out of 5 micrographs.

4.2 Using flags to pass arguments

In this section we demonstrate the execution of the program using flags to pass arguments.

The most basic way to run the program:

$ kltpicker -i /home/user/example/micrographs -o /home/user/example/results -s 300

If you want detailed progress output, run:

$ kltpicker -i /home/user/example/micrographs -o /home/user/example/results -s 300 -v

If you don’t want to use the GPU, run:

$ kltpicker -i /home/user/example/micrographs -o /home/user/example/results -s 300 --no-gpu

To specify additional optional parameters such as number of noise images to pick, the maximum number of processes to run, using ASOCEM for contamination detection, or the GPUs you want to use:

$ /data/amitay/cmpKltAsocemItai/micsForTesting/10061/ -o /data/amitay/cmpKltAsocemItai/resultsPython/10061/ -s 720 -n 30 -a --save-asocem --asocem-downsample 500 --asocem-area 3 -–max-processes 16 –-gpus 0 1 2 5 7

etc.

4.3 Displaying the results in EMAN and RELION

In order to display the picking results in EMAN [3], open a terminal in the output directory /home/user/example/results and create a new directory named eman, using the command:

$ mkdir eman

Change directory to eman and enter the following commands one by one:

$ e2rawdata.py /home/user/example/micrographs/\*.mrc --invert --edgenorm --xraypixel --ctfest --apix=1.0 --voltage=200.0 --cs=2.0 --ac=10.0 --threads=4 --defocusmin=0.6 --defocusmax=4.0

$ e2import.py

/home/user/example/results/pickedParticlesParticleSize300/box/\*.box -–import\_boxes –-box\_type=boxes

$ e2boxer.py --allmicrographs --boxsize=300 --ptclsize=300 --apix=1.0 –-no\_ctf --gui --threads=4

In order to display the picking results of the first micrograph in RELION [4], open the terminal in the output directory /home/user/example/results and create a new directory named relion, using the command:

$ mkdir relion

Change directory to relion and enter the following command:

$ relion\_display --i /home/user/example/micrographs/001.mrc –-cords /home/user/example/results/pickedParticlesParticleSize300/star/001.star --scale 0.15 –-particle\_radius 150 --angpix 1 --lowpass 20 -–pick

5 Citation

If you use the KLT picker, please cite  
**Eldar A, Landa B, Shkolnisky Y. KLT picker: Particle picking using data-driven optimal templates. J Struct Biol. 2020 May;210(2) 107473.**

A preprint is available at <https://arxiv.org/abs/1912.06500>.

6 References

[1] Iudin, A., Korir, P., Salavert-Torres, J., Kleywegt, G., and Patwardhan, A. (2016). EMPIAR: A public archive for raw electron microscopy image data. Nature Methods, 13.

[2] Wong, Wilson and Bai, Xiao-chen and Brown, Alan and Fernandez, Israel S and Hanssen, Eric and Condron, Melanie and Tan, Yan Hong and Baum, Jake and Scheres, Sjors H W. (2014). Cryo-EM structure of the Plasmodium falciparum 80S ribosome bound to the anti-protozoan drug emetine. Wong et al. eLife ,3,e03080.

[3] G. Tang, L. Peng, P.R. Baldwin, D.S. Mann, W. Jiang, I. Rees & S.J. Ludtke. (2007). EMAN2: an extensible image processing suite for electron microscopy. Journal of structural biology, 157, 38-46.

[4] Scheres, Sjors HW. (2015). Semi-automated selection of cryo-EM particles in RELION 1.3. Journal of structural biology, 128, 114-122.

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