

# MSMuTect, Version 0.5 manual

## Installation methods:

### PIP

pip3 install msmutect (not on pypi yet; this will not work yet)

### Github

**# cython must be preinstalled; if it is not, run: pip3 install Cython**

git clone [https://github.com/MaruvkaLab/MSMuTect\\_0.5](https://github.com/MaruvkaLab/MSMuTect_0.5)

cd [MSMuTect\\_0.5](#)

sh rename.sh

python3 setup.py install

## Defintions:

**Histogram** - a histogram is a structure containing indel information about a microsatellite locus. For instance, if the reference motif repeat count is 5, but some cells have a motif-length deletion, a histogram could be: 5\_10, 4\_6. This means there were 10 reads with a motif repeat length of 5, and 4 reads with a motif repeat length of 6

**Alleles** - Alleles are the alleles called by MSMuTect's rigorous statistical noise model for each microsatellite locus. The output file contains both the called alleles, their projected frequencies, and the extent to which the call precisely models the data (log likelihood).

**Mutations** - Mutations are the mutations called by MSMuTect, by analyzing a normal and tumor sample from the same patient. It uses a rigorous noise model to weed out noisy loci, only analyzing loci that are good candidates for a mutation. The final step in the mutation calling process is a Fisher Exact Test for the histograms of the normal and tumor file for a locus. If the p value of this test is under the threshold, the locus is called a mutation. The p value is also written to the output file (note that the majority of microsatellite loci will not have a Fisher Exact test performed since they do not pass the noise model).

## Flags:

Run msmutect --help to see help message.

All flags

- T TUMOR\_FILE, --tumor\_file TUMOR\_FILE  
Tumor BAM file
- N NORMAL\_FILE, --normal\_file NORMAL\_FILE  
Non-tumor BAM file
- S SINGLE\_FILE, --single\_file SINGLE\_FILE  
Analyze a single file for histogram and/or alleles
- I LOCI\_FILE, --loci\_file LOCI\_FILE  
File of loci to be processed and included in the output
- O OUTPUT\_PREFIX, --output\_prefix OUTPUT\_PREFIX  
prefix for all output files
- c CORES, --cores CORES

Number of cores to run MSMuTect on  
 -b BATCH\_START, --batch\_start BATCH\_START  
 1-indexed number locus to begin analyzing at (Inclusive)  
 -e BATCH\_END, --batch\_end BATCH\_END  
 1-indexed number locus to stop analyzing at (Inclusive)  
 -H, --histogram Output a Histogram File  
 -A, --allele Output allele file  
 -m, --mutation Output mutation file  
 -F FLANKING, --flanking FLANKING  
 Length of flanking on both sides of an accepted read  
 -f, --force overwrite pre-existing files

MSMuTect can run for pairs of files (ie. a regular and a tumor), or call histograms/alleles for a single file. The -S flag denotes a single file, and -T and -N flags denote Tumor and Normal files, respectively. The -H flag will generate a histogram, the -A flag will generate allele calls, and the -m flag will generate mutation calls (for pairs of files only). MSMuTect will always show preceding steps. In other words, the -m flag will output a file with alleles and histograms, and the alleles flag will output a file with histograms as well. In addition, by default, MSMuTect will do more work, not less. So, if you give MSMuTect a single file, it will call alleles and histograms for it, and only by passing the -H flag (and not the -A flag) will it only call histograms.

Some examples:

msmutect -l generic\_loci.phobos -S myBam.bam -O myout [-A]

Will generate a file containing the alleles and histograms for myBam.bam called myout.all.tsv.

The -A argument is unnecessary, calling alleles is the default behavior.

msmutect -l generic\_loci.phobos -S myBam.bam -H -O myout

Will generate a file holding only histograms called myout.hist.tsv. It's first columns will be identical to myout.all.tsv

For pairs of files, by default MSMuTect will only analyze loci that are considered candidates for a mutation after analysis of the Normal file. So, for instance,

msmutect -l generic\_loci.phobos -N normalBam.bam -T tumorBam.bam -O myout [-m]

Will output a file with mutations and some non mutations called myout.partial.mut.tsv

If you would like to see full data for a pair, for all loci, simply feed the -A or -H flags, along with the -m flag

msmutect -l generic\_loci.phobos -N normalBam.bam -T tumorBam.bam -O myout -m -A

Will output 3 files:

Myout.tumor.all.tsv

Myout.normal.all.tsv

And

Myout.full.mut.tsv

This will incur a significant performance penalty

All files will record the mutation call for convenience

Please note, the -S and -T/-N flags cannot be used in tandem. -S is for single files. -T and -N is for running pairs

## Support:

For support, please email [k.avraham@technion.ac.il](mailto:k.avraham@technion.ac.il) or [yosi.maruvka@bfe.technion.ac.il](mailto:yosi.maruvka@bfe.technion.ac.il)  
The code is hosted at Github at [https://github.com/MaruvkaLab/MSMuTect\\_0.5](https://github.com/MaruvkaLab/MSMuTect_0.5) . If there is some feature you would like added to make MSMuTect work for you, please do not hesitate to email us and we will consider adding it. The code is licensed under an MIT license