
jbiot Documentation

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CHAPTER 1

Introduction

fastq is used to clean raw fastq data...

CHAPTER 2

Authors

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CHAPTER 3

Status

Note: not reviewed yet.

CHAPTER 4

Installation

use git to clone code:

```
git clone git@192.168.1.251:/home/git/fastqtools.git
```

Attention: if you want to run fastqtools on local server without docker , try to add config.py.

CHAPTER 5

Usage

just type command:

```
/path/to/fastqClean.py -h  
/path/to/fastqStat.py -h  
/path/to/fastqSplit.py -h
```

developments followed by Dcer rules, script will need a yaml file, which shoud contain following key and values

5.1 must_args

- args1 desc of args2
- args2 desc of args2

5.2 optional args

- args3 desc of args3

here is a sample yaml file:

```
args1: value of args1  
args2: value of args2
```


CHAPTER 6

RUN

6.1 cli way

copy and paste to your input yaml file and call script:

```
/path/of/ctpips.py -c your.yml
```

6.2 serer way

send request to jbios with /start/ctpips/:

```
req = requests.get("http://<server>:port/ctpips/", data=json.dumps(indict))
```

for jbios detail information check api documentation [here](#)

Code

7.1 latest

7.1.1 fastqtools package

Subpackages

`fastqtools.fastqReader package`

Submodules

`fastqtools.fastqReader.fastqReader module`

```
class fastqtools.fastqReader.fastqReader(fq1,fq2)
```

```
    next()
```

```
class fastqtools.fastqReader.fastqReader.fq(id,seq,flag,qual)
```

`fastqtools.fastqReader.fastqWriter module`

```
fastqtools.fastqReader.fastqWriter.fastqWriter(read,prefix)
    write reads to fastqs
```

Module contents

fastqtools.fastqSplit package

Submodules

fastqtools.fastqSplit.fastqMerge module

```
fastqtools.fastqSplit.fastqMerge.fastqMerge(fqs, prefix)
```

fastqtools.fastqSplit.fastqSplit module

```
fastqtools.fastqSplit.fastqSplit.fastqSplit(fq1, fq2, splitNum, prefix)  
split Fastq in to small ones to accelerate downstreaming analysis...
```

fastqtools.fastqSplit.getfileNum module

```
fastqtools.fastqSplit.getfileNum.getfileNum(afile)
```

Module contents

fastqtools.fastqStat package

Submodules

fastqtools.fastqStat.fastqStat module

```
fastqtools.fastqStat.fastqStat.code2score(code)  
fastqtools.fastqStat.fastqStat.fastqStat(fq1, fq2, prefix)  
fastqtools.fastqStat.fastqStat.qual20(scores)  
fastqtools.fastqStat.fastqStat.score2code(score)
```

fastqtools.fastqStat.getfileNum module

```
fastqtools.fastqStat.getfileNum.getfileNum(afile)
```

Module contents

fastqtools.readprocess package

Submodules

fastqtools.readprocess.readprocess module

```
class fastqtools.readprocess.readprocess.readprocess (read)
```

```
    autoadaptremove (flag)
    length (lenMin)
    nbase (percent)
    qual (q, percent)
    trim (head1, tail1, head2, tail2)
    umi (umis)
```

fastqtools.readprocess.tools module

```
fastqtools.readprocess.tools.autocutadaptor (seq1, seq2)
```

```
fastqtools.readprocess.tools.checkN (seq, percent)
```

```
fastqtools.readprocess.tools.checking_adaptor (seq1, seq2, common, threshold=0.95)
```

```
fastqtools.readprocess.tools.checkqual (qual, q_thread, percent)
```

```
fastqtools.readprocess.tools.checkumi (seq, umis)
```

```
fastqtools.readprocess.tools.code2score (code)
```

```
fastqtools.readprocess.tools.diffstr (str1, str2)
```

```
class fastqtools.readprocess.tools.dnaseq
```

```
    static complement (seq)
```

```
    static reverse (seq)
```

```
fastqtools.readprocess.tools.enlong_and_find_common (seq1, seq2, seed_choosed, mismatchMax=5)
```

return best common sequences.

```
fastqtools.readprocess.tools.score2code (score)
```

```
fastqtools.readprocess.tools.seeding (seq1, seq2, seed_len=10, seed_max=3, seed_step=1)
```

find seed candidates return seeds and locus.

Module contents

Module contents

7.1.2 setup module

Setup file for jbiot.

```
setup.find_bins()  
setup.readfile(filename)  
setup.setup_package()
```

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