

# Package ‘qckit’

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**Type** Package

**Title** What the Package Does (Title Case)

**Version** 0.1.0

**Author** Who wrote it

**Maintainer** The package maintainer <yourself@somewhere.net>

**Description** More about what it does (maybe more than one line)

Use four spaces when indenting paragraphs within the Description.

**License** What license is it under?

**Encoding** UTF-8

**LazyData** false

**Suggests** testthat

**RoxygenNote** 6.0.1.9000

**LinkingTo** Rcpp

**Imports** magrittr,  
ggplot2,  
dplyr,  
seqTools,  
RSQLite,  
zlibbioc,  
Rcpp

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basic_stat	<i>Generate the data frame that includes percentiles of quality score per position</i>
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**Description**

Generate the data frame that includes percentiles of quality score per position

**Usage**

basic\_stat(infile)

**Arguments**

infile            the object that is the dataframe of the mean, median and quantiles of the FASTQ file from basic statistics function

**Value**

boxplot of per position quality score distribution

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calc_over_rep_seq	<i>calculate Over Rep seqs</i>
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**Description**

Description

**Usage**

calc\_over\_rep\_seq(infile, out\_prefix, min\_size = 5L, buffer\_size = 1000000L)

**Arguments**

infile            A string giving the path for the fastqfile  
out\_prefix        A string giving the prefix to be used for outputs  
min\_size          An int for thhresholding over representation  
buffer\_size        An int for the number of lines to keep in memory

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dimensions	<i>Extract the dimensions for Fastq file</i>
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**Description**

ncolumnuse seqTool to extract the dimensions of a Fastq G zipped file

**Usage**

```
dimensions(fseq, selection)
```

**Arguments**

fseq	an object that is the read result of the seq.read function
selection	"reads" for number of reads/rows, "positions" for number of positions/columns

**Value**

a numeric value of the number of reads or the number of positions

---

GC_content	<i>Extract GC content separately and calculate GC content percentage for each sequence read</i>
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**Description**

Extract GC content separately and calculate GC content percentage for each sequence read

**Usage**

```
GC_content(infile)
```

**Arguments**

infile	the object that is the path to the FASTQ file
--------	---

**Value**

plot of GC content

---

GC_content_plot	<i>Generate GC content plot from the GC content</i>
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**Description**

Generate GC content plot from the GC content

**Usage**

```
GC_content_plot(nc, gc_df, writefile = FALSE, prefix = "")
```

**Arguments**

nc	the object that is the number of positions of the FASTQ files
gc_df	the object that is the GC content vectors generated from GC content function
writefile	the object indicating intent to save the plot as pdf file, set default as FALSE
prefix	the prefix for the output file of the plot

**Value**

a ggplot of the GC content across all positions

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gc_per_read	<i>calculate GC percent per read</i>
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**Description**

Description

**Usage**

```
gc_per_read(infile)
```

**Arguments**

infile	A string giving the path for the fastqfile
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hello	<i>Hello, World!</i>
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**Description**

Prints 'Hello, world!'.

**Usage**

```
hello()
```

**Examples**

```
hello()
```

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kmer	<i>Extract kmers and kmer counts from FASTQ file to a data frame</i>
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**Description**

Extract kmers and kmer counts from FASTQ file to a data frame

**Usage**

```
kmer(name, kcount, writefile = FALSE)
```

**Arguments**

name	the object that is the path to gzipped FASTQ file
kcount	the object that is the length of kmer that is in interest
writefile	the boolean object that asks whether to write output as csv file

**Value**

data frame of kmer and corresponding kmer count of the length of choice

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overrepresented_sequence	<i>Sort all sequences per read by count along with a density plot of all counts with top 5 repeated sequences marked</i>
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**Description**

Sort all sequences per read by count along with a density plot of all counts with top 5 repeated sequences marked

**Usage**

```
overrepresented_sequence(infile, prefix, nr)
```

**Arguments**

infile	the object that is the path to gzipped FASTQ file
prefix	the prefix to name the output file
nr	the number of reads of the FASTQ file

**Value**

table of sequences sorted by count  
density plot of sequence length with top 5 marked by rugs, saved as PDF file

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overrep_kmer	<i>Generate overrepresented kmers from all kmer counts results</i>
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**Description**

Generate overrepresented kmers from all kmer counts results

**Usage**

```
overrep_kmer(path, k, nc, nr)
```

**Arguments**

path	the path to the gz file
k	the length of the sequence looking for
nc	number of positions
nr	number of reads

**Value**

the index of reads that has overrepresented kmers

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overrep_plot	<i>Plot the top 5 sequences</i>
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**Description**

Plot the top 5 sequences

**Usage**

```
overrep_plot(overrep_order, prefix)
```

**Arguments**

overrep_order	the table that sorts the sequence content and corresponding counts in descending order
prefix	the prefix to the file saved

**Value**

plot of the top 5 overrepresented sequences

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plotall	<i>Plot all plots about the FASTQ file and save them in a pdf as comprehensive report</i>
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**Description**

Plot all plots about the FASTQ file and save them in a pdf as comprehensive report

**Usage**

```
plotall(name, nr, nc, basic_stat, fseq)
```

**Arguments**

name	the object that is the path to the gzipped FASTQ file
nr	the number of reads in the dataset
nc	the number of positions in the dataset
basic_stat	the dataframe of the basic stat informations
fseq	the object that is the seqTools processed data

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plotSeqContent	<i>Plot the per position nucleotide content</i> plotSeqContent
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**Description**

Plot the per position nucleotide content plotSeqContent

**Usage**

```
plotSeqContent(fseq, nr, nc)
```

**Arguments**

nr	the number of reads of the FASTQ file, acquired through previous functions
nc	the number of positions of the FASTQ file, acquired through previous functions
name	the object that is the path to the gzipped FASTQ file

**Value**

ggplot line plot of all nucleotide content inclding A, T, G, C and N

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plot_perseq_quality	<i>plot the mean quality score per read in histograms</i>
plot_perseq_quality	

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**Description**

plot the mean quality score per read in histograms plot\_perseq\_quality

**Usage**

```
plot_perseq_quality(infile)
```

**Arguments**

infile            the object that is the path to the file that

**Value**

plot of mean quality score per read

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plot_quality_score	<i>Generate a boxplot of the per position quality score from basic statistics results</i>
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**Description**

Generate a boxplot of the per position quality score from basic statistics results

**Usage**

```
plot_quality_score(basic_statistics)
```

**Arguments**

basic\_stat        the object that is the dataframe of the mean, median and quantiles of the FASTQ file from basic statistics function

**Value**

boxplot of per position quality score distribution



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plot_sequence_length	<i>extract the sequence length per read and plot corresponding bar plot</i>
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**Description**

extract the sequence length per read and plot corresponding bar plot

**Usage**

```
plot_sequence_length(fseq)
```

**Arguments**

fseq	the object that is the seqTools processed result
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**Value**

the plot of the sequence distribution among all reads

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process_fastq	<i>calculate Over Rep seqs</i>
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**Description**

Description

**Usage**

```
process_fastq(infile, out_prefix, buffer_size)
```

**Arguments**

infile	A string giving the path for the fastqfile
out_prefix	A string giving the prefix to be used for outputs
buffer_size	An int for the number of lines to keep in memory

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qual_score_per_read	<i>calculate mean quality per read</i>
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**Description**

Description

**Usage**

```
qual_score_per_read(infile)
```

**Arguments**

infile	A string giving the path for the fastqfile
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sequence_content	<i>Extract nucleotide sequence content per position from fastq file</i>
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**Description**

Extract nucleotide sequence content per position from fastq file

**Usage**

```
sequence_content(fseq, content)
```

**Arguments**

fseq	an object that is the read result from seq.read function
content	an object of string type that specifies the content in question, "A", "T", "G", "C", "N"(either capital or lower case)

**Value**

the per position

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