# Package 'qPCRtools'

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Version 0.1.1		
Title Tools for qPC	R	
Description A set of	of tools for qPCR data process.	
URL https://githu	ub.com/lixiang117423/qPCRtools	
BugReports https	://github.com/lixiang117423/qPCRtools/issues	
${\bf License}\ {\rm MIT}+{\rm file}$	LICENSE	
	ta.table, dplyr, ggplot2, ggpmisc, magrittr, lxl, reshape2, sjmisc, stringr, xlsx	
RoxygenNote 7.2.0		
NeedsCompilation	no	
Author Xiang LI [c	re, aut]	
Maintainer Xiang	m LI < lixiang 117423 @gmail.com >	
$ \begin{array}{c} {\rm CalCurve} & . \\ {\rm CalRTable} & . \\ \\ {\bf Index} \end{array}$		1 2 <b>3</b>
CalCurve	Standard Curve Calculation.	_
Description		
Standard Curve	Calculation.	
Arguments		
cq.table	The data frame of the position and Cq value.	
concen.table	The data frame of the position and concentration.	
highest.concen	The highest concentration.	
lowest.concen	The lowest concentration.	
dilution	Dilution factor of cDNA template.	
by.mean	Calculation by mean Cq value or not.	

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#### Value

A list.

#### Author(s)

Xiang LI ¡lixiang117423@gmail.com;

#### Examples

```
df.1.path <- system.file("examples", "calsc.cq.txt", package = "qPCRtools")
df.2.path <- system.file("examples", "calsc.info.txt", package = "qPCRtools")
df.1 <- data.table::fread(df.1.path)
df.2 <- data.table::fread(df.2.path)
CalCurve(
    cq.table = df.1,
    concen.table = df.2,
    lowest.concen = 4,
    highest.concen = 4096,
    dilu = 4,
    by = "mean"
) -> p

p[["table"]]
p[["figure"]]
```

CalRTable

Calculate volume.

## Description

Calculate RNA and other reagent volume required for reverse transcription.

#### **Arguments**

data A data frame contained the sample names and the concentration value.

The default unit of concentration is ng/uL.

template A data frame contained the information of reverse transcription.

RNA weight RNA weight required for reverse transcription. Default is 1 ug.

## Value

A data frame.

#### Author(s)

Xiang LI ¡lixiang117423@gmail.com¿

#### Examples

```
df.1.path <- system.file("examples", "crtv.data.txt", package = "qPCRtools")
df.2.path <- system.file("examples", "template", package = "qPCRtools")
df.1 <- data.table::fread(df.1.path)
df.2 <- data.table::fread(df.2.path)
result <- CalRTable(data = df.1, template = df.2, RNA.weight = 2)
head(result)</pre>
```

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