

# Package ‘qPCRtools’

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**Version** 0.1.1

**Title** Tools for qPCR

**Description** A set of tools for qPCR data process.

**URL** <https://github.com/lixiang117423/qPCRtools>

**BugReports** <https://github.com/lixiang117423/qPCRtools/issues>

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**Imports** broom, data.table, dplyr, ggplot2, ggpmisc, ggthemes,  
magrittr, multcomp, readxl, reshape2, rstatix, sjmisc, stringr,  
tibble, tidyr, xlsx

**RoxygenNote** 7.2.0

**NeedsCompilation** no

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CalCurve	<i>Standard Curve Calculation.</i>
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## 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.

**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"]]
```

---

CalExp2ddCt

*Calculate expression using standard curve.*

---

**Description**

Calculate expression using standard curve.

**Arguments**

cq.table	The data frame of the position and cq value.
design.table	The data frame of the position and corresponding information.
correction	Correct expression value by reference gene.
ref.gene	The name of reference gene.
ref.group	The name of reference group.
stat.method	Statistical method.
fig.type	Calculation by mean cq value or not.
fig.ncol	Number of columes of figure.

**Author(s)**

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**Examples**

```
df1.path = system.file("examples", "ddct.cq.txt", package = "qPCRtools")
df2.path = system.file("examples", "ddct.design.txt", package = "qPCRtools")

cq.table = data.table::fread(df1.path)
design.table = data.table::fread(df2.path)

CalExp2ddCt(cq.table,
            design.table,
            ref.gene = "OsUBQ",
            ref.group = "CK",
            stat.method = "t.test",
            fig.type = "box",
            fig.ncol = NULL) -> res

res[["table"]]
res[["figure"]]
```

---

CalExpCurve

*Calculate expression using standard curve.*

---

**Description**

Calculate expression using standard curve.

**Arguments**

<code>cq.table</code>	The data frame of the position and Cq value.
<code>design.table</code>	The data frame of the position and corresponding information.
<code>correction</code>	Correct expression value by reference gene.
<code>ref.gene</code>	The name of reference gene.
<code>stat.method</code>	Statistical method.
<code>ref.group</code>	The name of reference group.
<code>fig.type</code>	Calculation by mean Cq value or not.
<code>fig.ncol</code>	Number of columns of figure.

**Author(s)**

Xiang LI jlixiang117423@gmail.com;

## Examples

```
df1.path = system.file("examples", "cal.exp.curve.cq.txt", package = "qPCRtools")
df2.path = system.file("examples", "cal.expre.curve.sdc.txt", package = "qPCRtools")
df3.path = system.file("examples", "cal.exp.curve.design.txt", package = "qPCRtools")

cq.table = data.table::fread(df1.path)
curve.table = data.table::fread(df2.path)
design.table = data.table::fread(df3.path)

CalExpCurve(
  cq.table,
  curve.table,
  design.table,
  correction = TRUE,
  ref.gene = "OsUBQ",
  stat.method = "t.test",
  ref.group = "CK",
  fig.type = "box",
  fig.ncol = NULL) -> res

res[["table"]]
res[["figure"]]
```

---

CalExpRqPCR

*Calculate expression using standard curve.*

---

## Description

Calculate expression using standard curve.

## Arguments

<code>cq.table</code>	The data frame of the position and cq value.
<code>design.table</code>	The data frame of the position and corresponding information.
<code>correction</code>	Correct expression value by reference gene.
<code>ref.gene</code>	The name of reference gene.
<code>ref.group</code>	The name of reference group.
<code>stat.method</code>	Statistical method.
<code>fig.type</code>	Calculation by mean cq value or not.
<code>fig.ncol</code>	Number of columen of figure.

## Author(s)

Xiang LI [jlixiang117423@gmail.com](mailto:jlixiang117423@gmail.com),

**Examples**

```
df1.path <- system.file("examples", "cal.expre.rqpcr.cq.txt", package = "qPCRtools")
df2.path <- system.file("examples", "cal.expre.rqpcr.design.txt", package = "qPCRtools")

cq.table <- data.table::fread(df1.path, header = TRUE)
design.table <- data.table::fread(df2.path, header = TRUE)

CalExpRqPCR(cq.table,
            design.table,
            ref.gene = NULL,
            ref.group = "CK",
            stat.method = "t.test",
            fig.type = "box",
            fig.ncol = NULL
            ) -> res

res[["table"]]
res[["figure"]]
```

---

CalRTable

*Calculate volume.*


---

**Description**

Calculate RNA and other reagent volume required for reverse transcription.

**Arguments**

<b>data</b>	A data.frame contained the sample names and the concentration value. The default unit of concentration is ng/uL.
<b>template</b>	A data.frame contained the information of reverse transcription.
<b>RNA.weight</b>	RNA weight required for reverse transcription. Default is 1 ug.

**Value**

A data frame.

**Author(s)**

Xiang LI jlixiang117423@gmail.com

**Examples**

```
df1.path <- system.file("examples", "crtv.data.txt", package = "qPCRtools")
df2.path <- system.file("examples", "crtv.template.txt", package = "qPCRtools")
df1 <- data.table::fread(df1.path)
df2 <- data.table::fread(df2.path)
result <- CalRTable(data = df1, template = df2, RNA.weight = 2)
head(result)
```

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