

# Package ‘lmQCM’

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**Type** Package

**Title** An Algorithm for Gene Co-Expression Analysis

**Version** 0.2.1

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**Description**

Implementation based on Zhang, Jie & Huang, Kun (2014) <doi:10.4137/CIN.S14021> Normalized lmQCM: An Algorithm for Detecting Weak Quasi-Cliques in Weighted Graph with Applications in Gene Co-Expression Module Discovery in Cancers. Cancer informatics, 13, CIN-S14021.

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**Encoding** UTF-8

**Depends** genefilter, Biobase, progress, stats, methods

**Suggests** devtools, roxygen2

**LazyData** true

**RoxygenNote** 6.1.0

**URL** <http://github.com/huangzhii/lmQCM>

**BugReports** <http://github.com/huangzhii/lmQCM/issues>

**NeedsCompilation** no

**Repository** CRAN

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fastFilter	<i>fastFilter: Subroutine for filtering expression matrix</i>
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**Description**

Author: Zhi Huang

**Usage**

```
fastFilter(RNA, lowest_percentile_mean = 0.2,  
           lowest_percentile_variance = 0.2, var.func = "var")
```

**Arguments**

RNA	an expression matrix (rows: genes; columns: samples)
lowest_percentile_mean	a float value range 0-1
lowest_percentile_variance	a float value range 0-1
var.func	specify variance function

**Value**

An filtered expression matrix

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lmQCM	<i>lmQCM: Main Routine for Gene Co-expression Analysis</i>
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**Description**

Author: Zhi Huang

**Usage**

```
lmQCM(data_in, gamma = 0.55, t = 1, lambda = 1, beta = 0.4,  
       minClusterSize = 10, CCmethod = "pearson", normalization = F)
```

**Arguments**

data_in	real-valued expression matrix with rownames indicating gene ID or gene symbol
gamma	gamma value (default = 0.55)
t	t value (default = 1)
lambda	lambda value (default = 1)
beta	beta value (default = 0.4)
minClusterSize	minimum length of cluster to retain (default = 10)
CCmethod	Methods for correlation coefficient calculation (default = "pearson"). Users can also pick "spearman".
normalization	Determine if normalization is needed on massive correlation coefficient matrix.

**Value**

QCMObject - An S4 Class with lmQCM results

**Examples**

```
library(lmQCM)
library(Biobase)
data(sample.ExpressionSet)
data = assayData(sample.ExpressionSet)$exprs
data = fastFilter(data, 0.2, 0.2)
lmQCM(data)
```

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localMaximumQCM

*localMaximumQCM: Subroutine for Creating Gene Clusters*

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**Description**

Author: Zhi Huang

**Usage**

```
localMaximumQCM(cMatrix, gamma = 0.55, t = 1, lambda = 1)
```

**Arguments**

cMatrix	a correlation matrix
gamma	gamma value (default = 0.55)
t	t value (default = 1)
lambda	lambda value (default = 1)

**Value**

An unmerged clusters group 'C'

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`merging_lmQCM`*merging\_lmQCM: Subroutine for Merging Gene Clusters*

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**Description**

Author: Zhi Huang

**Usage**`merging_lmQCM(C, beta = 0.4, minClusterSize = 10)`**Arguments**

<code>C</code>	Resulting clusters
<code>beta</code>	beta value (default = 0.4)
<code>minClusterSize</code>	minimum length of cluster to retain (default = 10)

**Value**

mergedCluster - An merged clusters group

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