

# HTqPCR - high-throughput qPCR analysis in R and Bioconductor

Heidi Dvinge

November 1, 2011

## 1 Introduction

The package *HTqPCR* is designed for the analysis of cycle threshold (Ct) values from quantitative real-time PCR data. The main areas of functionality comprise data import, quality assessment, normalisation, data visualisation, and testing for statistical significance in Ct values between different features (genes, miRNAs).

Example data is included from TaqMan Low Density Arrays (TLDA), a proprietary format of Applied Biosystems, Inc. However, most functions can be applied to any kind of qPCR data, regardless of the nature of the experimental samples and whether genes or miRNAs were measured. Section 13 gives some examples of how to handle other input data, including data from a non-well based microfluidic system (Dynamic Arrays from Fluidigm Corporation).

```
> library("HTqPCR")
```

The package employs functions from other packages of the Bioconductor project ([Gentleman et al., 2004](#)). Dependencies include *Biobase*, *RColorBrewer*, *limma*, *statmod*, *affy* and *gplots*.

### Examples from the vignette

This vignette was developed in Sweave, so the embedded R code was compiled when the PDF was generated, and its output produced the results and plots that appear throughout the document. The following commands will extract all of the code from this file:

```
> all.R.commands <- system.file("doc", "HTqPCR.Rnw",  
+   package = "HTqPCR")  
> Stangle(all.R.commands)
```

This will create a file called HTqPCR.R in your current working directory, and this file can then either be sourced directly, or the commands run individually.

### General workflow

The main functions and their use are outlined in Figure 1. Note that the QC plotting functions can be used both before and after normalisation, in order to examine the quality of the data or look for particular trends.

For the full set of available functions type:

```
> ls("package:HTqPCR")
```

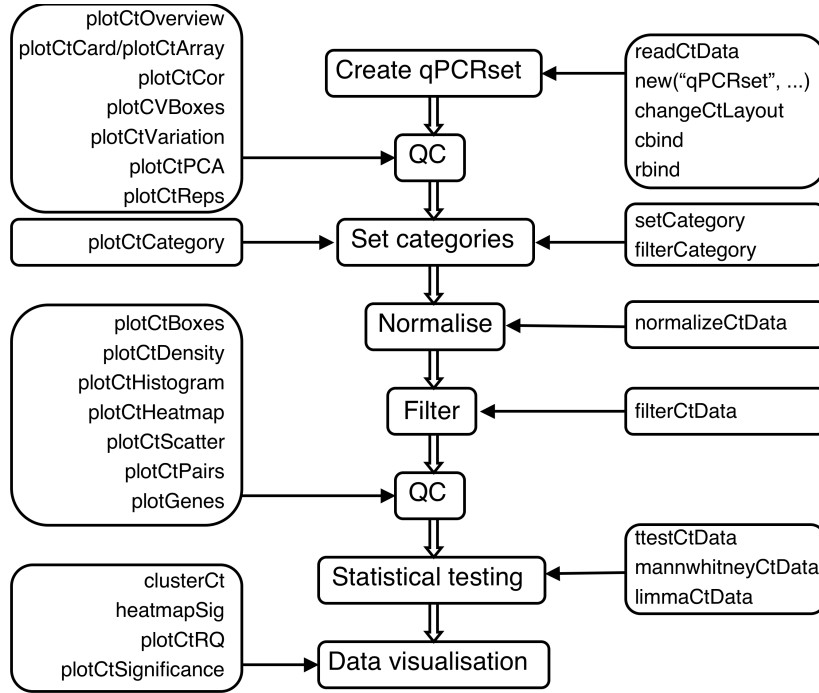


Figure 1: Workflow in *HTqPCR* analysis of qPCR data. Centre column: The main procedural steps in a typical qPCR analysis; Left: examples of visualisation functions; Right: data analysis functions.

[1] "cbind.qPCRset"	"changeCtLayout"
[3] "clusterCt"	"featureCategory"
[5] "featureCategory<-"	"featureClass"
[7] "featureClass<-"	"featurePos"
[9] "featurePos<-"	"featureType"
[11] "featureType<-"	"filterCategory"
[13] "filterCtData"	"flag"
[15] "flag<-"	"getCt"
[17] "getCtHistory"	"heatmapSig"
[19] "limmaCtData"	"mannwhitneyCtData"
[21] "n.samples"	"n.wells"
[23] "normalizeCtData"	"plotCVBoxes"
[25] "plotCtArray"	"plotCtBoxes"
[27] "plotCtCard"	"plotCtCategory"
[29] "plotCtCor"	"plotCtDensity"
[31] "plotCtHeatmap"	"plotCtHistogram"
[33] "plotCtLines"	"plotCtOverview"
[35] "plotCtPCA"	"plotCtPairs"
[37] "plotCtRQ"	"plotCtReps"
[39] "plotCtScatter"	"plotCtSignificance"
[41] "plotCtVariation"	"rbind.qPCRset"
[43] "readCtData"	"setCategory"
[45] "setCt<-"	"summary"
[47] "ttestCtData"	

## Getting help

Please send questions about *HTqPCR* to the Bioconductor mailing list.

See <http://www.bioconductor.org/docs/mailList.html>

## 2 *qPCRset* objects

The data is stored in an object of class *qPCRset*, which is similar to the *ExpressionSet* from package *Biobase* used for handling microarray data. The format is the same for raw and normalized data, and depending on how much information is available about the input data, the object can contain the following slots:

**featureNames** Object of class *character* giving the names of the features (genes, miRNAs) used in the experiment.

**sampleNames** Object of class *character* containing the sample names of the individual experiments.

**exprs** Object of class *matrix* containing the Ct values.

**flag** Object of class *data.frame* containing the flag for each Ct value, as supplied by the input files. These are typically set during the calculation of Ct values, and indicate whether the results are flagged as e.g. “Passed” or “Flagged”.

**featureType** Object of class *factor* representing the different types of features on the card, such as endogenous controls and target genes.

**featurePos** Object of class *character* representing the location “well” of a gene in case TLDA cards are used, or some other method containing a defined spatial layout of features.

**featureClass** Object of class *factor* with some meta-data about the genes, for example if it is a transcription factor, kinase, marker for different types of cancers or similar. This is typically set by the user.

**featureCategory** Object of class *data.frame* representing the quality of the measurement for each Ct value, such as “OK”, “Undetermined” or “Unreliable”. These can be set using the function **setCategories** depending on a number of parameters, such as how the Ct values are flagged, upper and lower limits of Ct values and variations between technical and biological replicates of the same feature.

**history** Object of class *data.frame* indicating what operations have been performed on the *qPCRset* object, and what the parameters were. Automatically set when any of the functions on the upper right hand side of figure 1 are called (**readCtData**, **setCategory**, **filterCategory**, **normalizeCtData**, **filterCtData**, **changeCtLayout**, **rbind**, **cbind**).

Generally, information can be handled in the *qPCRset* using the same kind of functions as for *ExpressionSet*, such as **exprs**, **featureNames** and **featureCategory** for extracting the data, and **exprs<-**, **featureNames<-** and **featureCategory<-** for replacing or modifying values.

The use of **exprs** might not be intuitive to users who are not used to dealing with microarray data, and hence *ExpressionSet*. The functions **getCt** and **setCt<-** that perform the same

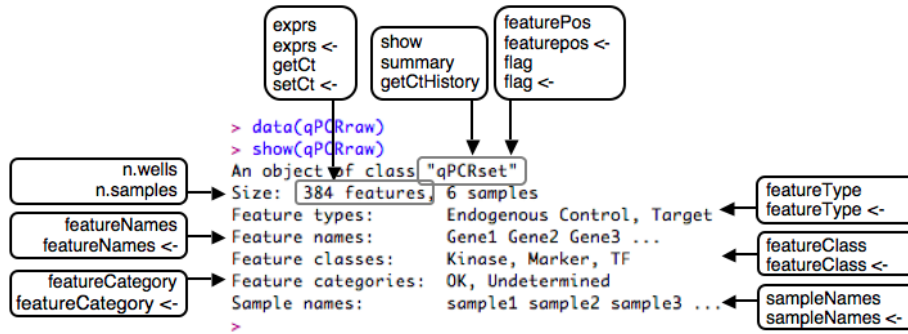


Figure 2: An example of a *qPCRset* object, and some of the functions that can be used to display and/or alter different aspects of the object, i.e. the accessor and replacement functions.

operations as `exprs` and `exprs<-` are therefore also included. For the sake of consistency, `exprs` will be used throughout this vignette for accessing the Ct values, but it can be replaced by `getCt` in all examples.

`qPCRset` objects can also be combined or reformatted in various ways (see section 12).

Two `qPCRset` test objects are included in the package: one containing raw data, and the other containing processed values. An example is shown in figure 2, along with some of the functions that can typically be used for manipulating *qPCRset* objects.

```

> data(qPCRraw)
> data(qPCRpros)
> class(qPCRraw)
[1] "qPCRset"
attr(,"package")
[1] ".GlobalEnv"

```

### 3 Reading in the raw data

The standard input consists of tab-delimited text files containing the Ct values for a range of genes. Additional information, such as type of gene (e.g. target, endogenous control) or groupings of genes into separate classes (e.g. markers, kinases) can also be read in, or supplied later.

The package comes with example input files (from Applied Biosystem's TLDA cards), along with a text file listing sample file names and biological conditions.

```
> path <- system.file("exData", package = "HTqPCR")
> head(read.delim(file.path(path, "files.txt")))

      File Treatment
1 sample1.txt   Control
2 sample2.txt LongStarve
3 sample3.txt LongStarve
4 sample4.txt   Control
5 sample5.txt   Starve
6 sample6.txt   Starve
```

The data consist of 192 features represented twice on the array and labelled "Gene1", "Gene2", etc. There are three different conditions, "Control", "Starve" and "LongStarve", each having 2 replicates.

The input data consists of tab-delimited text files (one per sample); however, the format is likely to vary depending on the specific platform on which the data were obtained (e.g., TLDA cards, 96-well plates, or some other format). The only requirement is that columns containing the Ct values and feature names are present.

```
> files <- read.delim(file.path(path, "files.txt"))
> raw <- readCtData(files = files$File, path = path)
```

The *qPCRset* object looks like:

```
> show(raw)
An object of class "qPCRset"
Size: 384 features, 6 samples
Feature types:          Endogenous Control, Target
Feature names:          Gene1 Gene2 Gene3 ...
Feature classes:
Feature categories:      OK, Undetermined
Sample names:            sample1 sample2 sample3 ...
```

NB: This section only deals with data presented in general data format. For notes regarding other types of input data, see section 13. This section also briefly deals with other types of qPCR results besides Ct data, notably the Cp values reported by the LightCycler System from Roche.

## 4 Data visualisation

### 4.1 Overview of Ct values across all groups

To get a general overview of the data the (average) Ct values for a set of features across all samples or different condition groups can be displayed. In principle, all features in a sample might be chosen, but to make it less cluttered Figure 3 displays only the first 10 features. The top plot was made using just the Ct values, and shows the 95% confidence interval across replicates within and between samples. The bottom plot represents the same values but relative to a chosen calibrator sample, here the “Control”. Confidence intervals can also be added to the relative plot, in which case these will be calculated for all values compared to the average of the calibrator sample per gene.

```
> g <- featureNames(raw)[1:10]
> plotCtOverview(raw, genes = g, xlim = c(0, 50), groups = files$Treatment,
+   conf.int = TRUE, ylim = c(0, 55))

> plotCtOverview(raw, genes = g, xlim = c(0, 50), groups = files$Treatment,
+   calibrator = "Control")
```

### 4.2 Spatial layout

When the features are organised in a particular spatial pattern, such as the 96- or 384-well plates, it is possible to plot the Ct values or other characteristics of the features using this layout. Figure 4 shows an example of the Ct values, as well as the location of different classes of features (using random examples here), across all the wells of a TLDA microfluidic card.

```
> plotCtCard(raw, col.range = c(10, 35), well.size = 2.6)

> featureClass(raw) <- factor(c("Marker", "TF", "Kinase")[sample(c(1,
+   1, 2, 2, 1, 3), 384, replace = TRUE)])
> plotCtCard(raw, plot = "class", well.size = 2.6)
```

### 4.3 Comparison of duplicated features within samples

When a sample contains duplicate measurements for some or all features, the Ct values of these duplicates can be plotted against each other to measure accordance between duplicates. In Figure 5 the duplicates in sample2 are plotted against each other, and those where the Ct values differ more than 20% from the average of a given feature are marked.

```
> plotCtReps(qPCRraw, card = 2, percent = 20)
```

Replicates differing > 20% on card 2:

	rep1	rep2
Gene135	40.000000	29.90044
Gene14	40.000000	27.69185
Gene37	7.408248	40.00000
Gene73	40.000000	23.11949
Gene84	21.673946	40.00000
Gene86	20.389658	40.00000
Gene9	22.494440	31.39404
Gene95	15.916160	40.00000

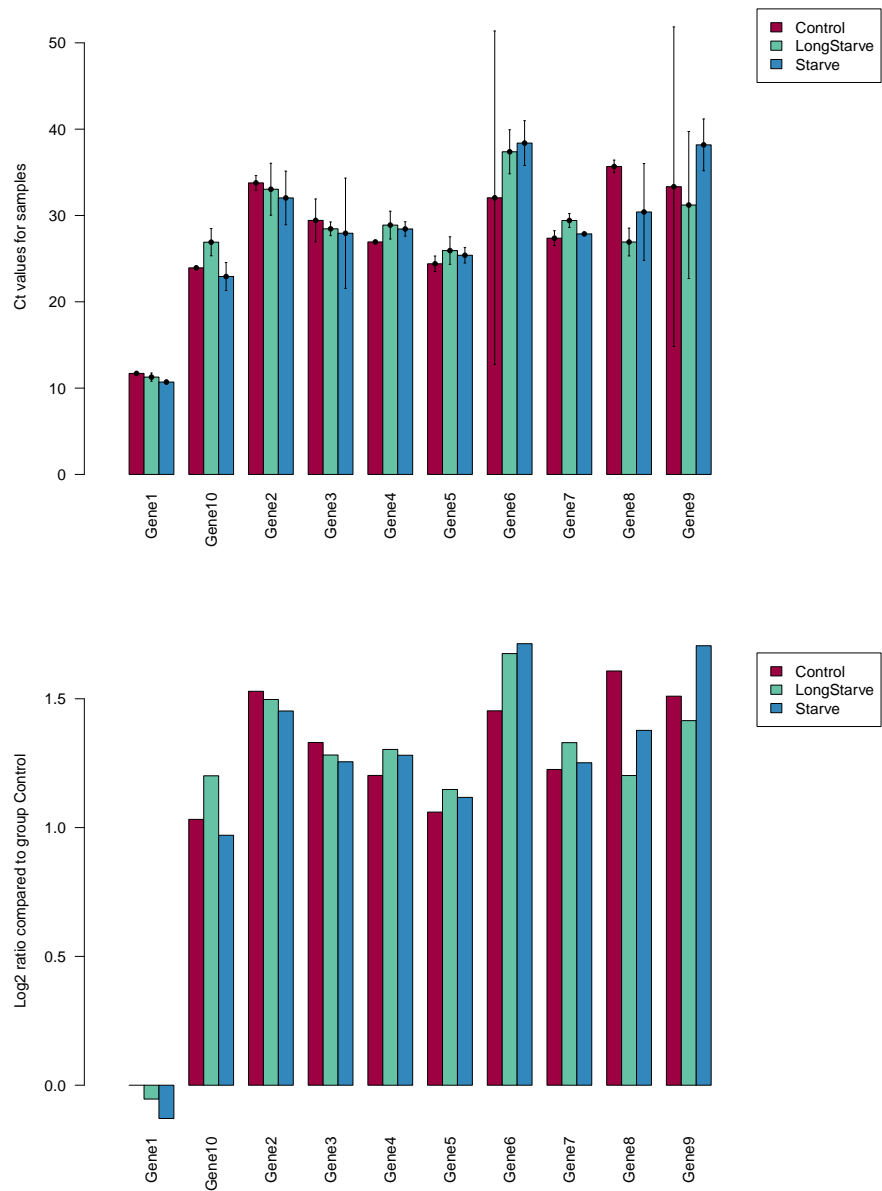


Figure 3: Overview of Ct values for the raw data.

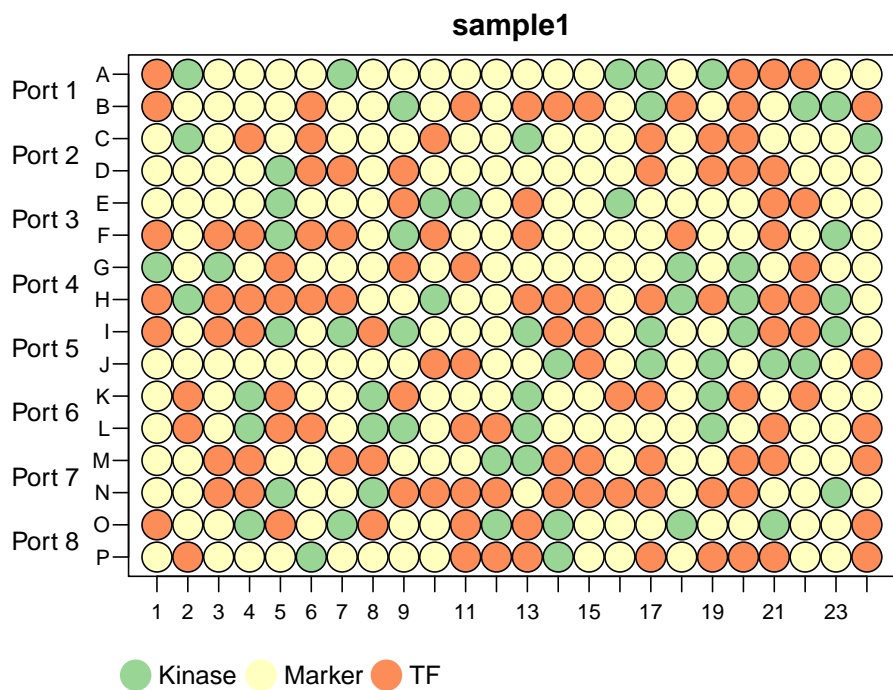
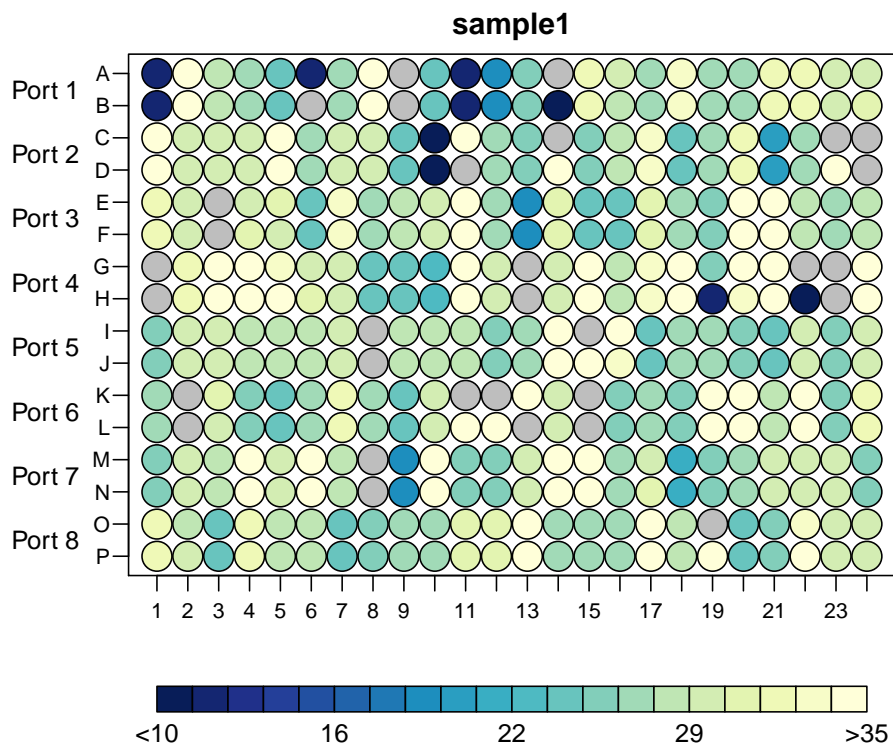


Figure 4: Ct values for the first sample (top)<sup>8</sup>, and the location of different feature classes (bottom). Ct values are visualised using colour intensity, and grey circles are features that were marked “undetermined” in the input file.

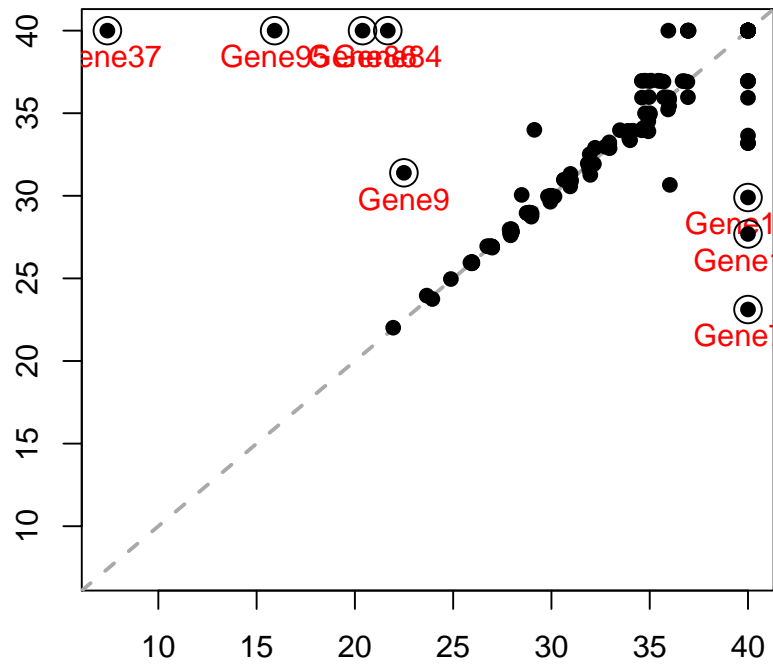


Figure 5: Concordance between duplicated Ct values in sample 2, marking features differing >20% from their mean.

Differences will often arise due to one of the duplicates marked as “Undetermined”, thus contributing to an artificially high Ct value, but other known cases exist as well.

#### 4.4 Variation within and across samples

In some cases more than two replicates are present, either within each qPCR card (feature replicates) or across cards (replicated samples). Assessing the variation within replicates can indicate whether some samples or individual features are less reliable, or if perhaps an entire qPCR card shows high variation between replicate features and needs to be discarded. `plotCtVariation` generates a boxplot with all the variation values, either across genes or with each samples. That way the general distribution of variation or standard deviation values can be compared quickly (Figure 6). For illustration, we use a data set where one of the samples is highly variable.

```
> raw.mix <- raw
> exprs(raw.mix)[, 6] <- sample(exprs(raw[, 6]))
> plotCtVariation(raw.mix, variation = "sd", log = TRUE,
+   main = "SD of replicated features", col = "lightgrey")
```

If it looks like there’s an unacceptable (or interesting) difference in the variation, this can be further investigated using the parameter `type="detail"`. This will generate multiple sub-plots, containing a single scatterplot of variation versus mean for each gene or sample (Figure 6). That way individual outliers can be identified, or whole samples removed by examining the resulting variation in more detail.

```
> raw.variation <- plotCtVariation(raw.mix, type = "detail",
+   add.featurenames = TRUE, pch = " ", cex = 1.2)
```

```
> names(raw.variation)
[1] "Var" "Mean"
> head(raw.variation[["Var"]][, 1:4])
  Group.1   sample1   sample2   sample3
1   Gene1 6.603053e-02 3.367143e-02 4.340261e-02
2  Gene10 6.087458e-05 2.864660e-04 1.869050e-03
3 Gene100 1.232561e-05 7.789010e-04 4.452426e-04
4 Gene101 5.859356e-03 8.787857e-02 7.988004e-06
5 Gene102 6.571738e-03 7.440192e-02 2.768647e-03
6 Gene103 0.000000e+00 2.544768e-06 7.899120e-04
> head(raw.variation[["Mean"]][, 1:4])
  Group.1 sample1 sample2 sample3
1   Gene1 11.70414 11.78938 10.76248
2  Gene10 23.93387 25.92385 27.88952
3 Gene100 27.93282 31.92726 30.92547
4 Gene101 27.88928 34.71769 28.93203
5 Gene102 28.91345 30.76310 28.90544
6 Gene103 40.00000 30.92895 29.90892
> apply(raw.variation[["Var"]][, 3:7], 2, summary)
      sample2   sample3   sample4   sample5 sample6
Min.    0.000e+00 0.000e+00 0.000e+00 0.000e+00  0.000
1st Qu. 3.674e-04 7.037e-04 3.074e-04 3.466e-04  2.186
```

```

Median  4.805e-03 5.066e-03 6.370e-03 6.630e-03 13.020
Mean    8.667e+00 4.440e+00 7.155e+00 6.887e+00 34.910
3rd Qu. 6.222e-02 3.446e-02 3.648e-02 3.863e-02 47.740
Max.    5.311e+02 5.716e+02 4.675e+02 3.328e+02 421.700
> colSums(raw.variation[["Var"]][, 3:7] > 20)
sample2 sample3 sample4 sample5 sample6
      11       3       7       6      82

```

In the example in this section a lot of features from sample 6 have intra-replicate variation above an arbitrary threshold selected based on Figure 6, and the mean and median values are much higher than for the remaining samples. Sample 1 is excluded due to the page width.

Variation across Ct values is discussed further in the following section regarding filtering.

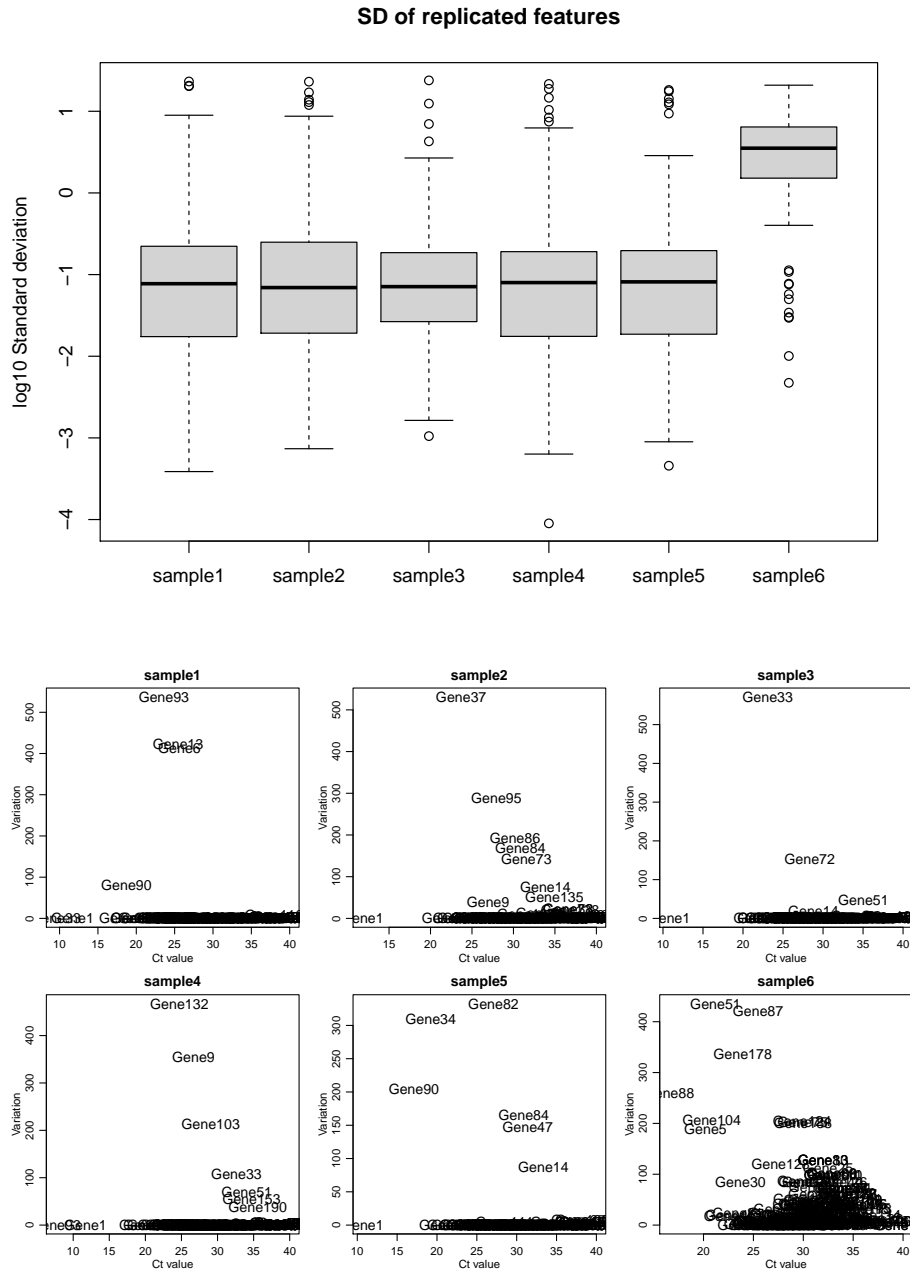


Figure 6: (Top) summary of standard deviation between replicated features within each of the six samples. (Bottom) Variation versus mean for replicated features.

## 5 Feature categories and filtering

Each Ct values in HTqPCR has an associated feature category. This is an important component to indicate the reliability of the qPCR data. Aside from the “OK” indicator, there are two other categories: “Undetermined” is used to flag Ct values above a user-selected threshold, and “Unreliable” indicates Ct values that are either so low as to be estimated by the user to be problematic, or that arise from deviation between individual Ct values across replicates. By default, only Ct values labelled as “undetermined” in the input data files are placed into the “Undetermined” category, and the rest are classified as “OK”. However, either before or after normalisation these categories can be altered depending on various criteria.

**Range of Ct values** Some Ct values might be too high or low to be considered a reliable measure of gene expression in the sample, and should therefore not be marked as “OK”.

**Flags** Depending on the qPCR input the values might have associated flags, such as “Passed” or “Failed”, which are used for assigning categories.

**Biological and technical replicates** If features are present multiple times within each sample, or if samples are repeated in the form of technical or biological replicates, then these values can be compared. Ct values lying outside a user-selected confidence interval (90% by default) will be marked as “Unreliable”.

```
> raw.cat <- setCategory(raw, groups = files$Treatment,
+   quantile = 0.8)
```

Categories after Ct.max and Ct.min filtering:

	sample1	sample2	sample3	sample4	sample5	sample6
OK	313	264	327	295	296	286
Undetermined	68	119	56	86	86	96
Unreliable	3	1	1	3	2	2

Categories after standard deviation filtering:

	sample1	sample2	sample3	sample4	sample5	sample6
OK	301	254	319	274	277	275
Undetermined	68	119	56	86	86	96
Unreliable	15	11	9	24	21	13

A summary plot for the sample categories is depicted in Figure 7. The result can be stratified by `featureType` or `featureClass`, for example to determine whether one class of features performed better or worse than others.

```
> plotCtCategory(raw.cat)
```

```
> plotCtCategory(raw.cat, stratify = "class")
```

The results can also be shown per feature rather than averaged across samples (Figure 8).

```
> plotCtCategory(raw.cat, by.feature = TRUE, cexRow = 0.1)
```

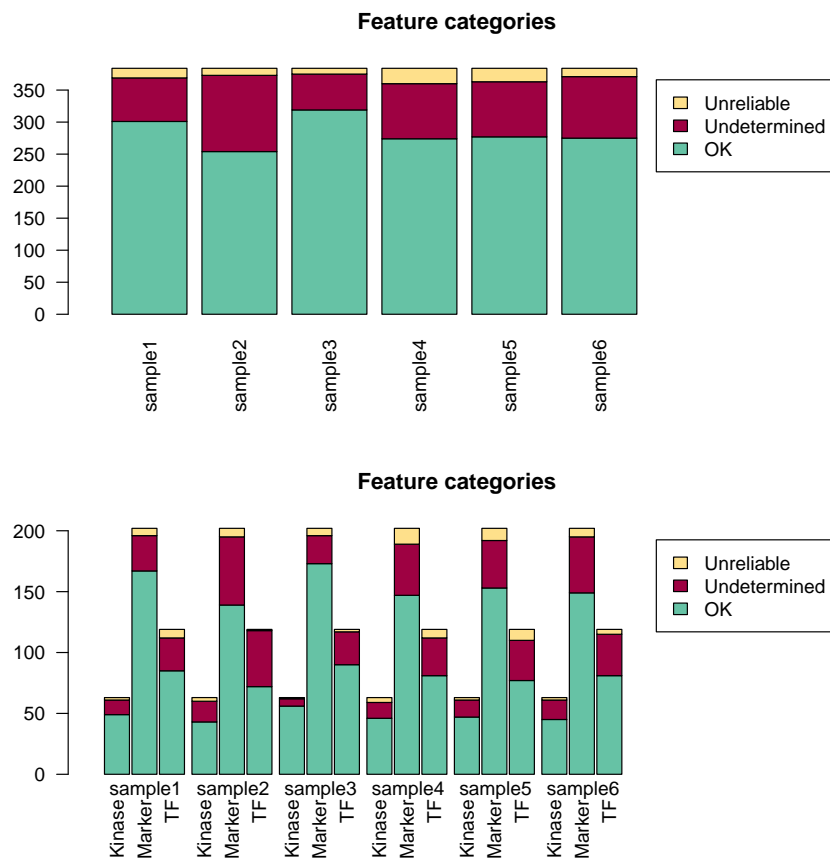


Figure 7: Summary of the categories, either for each sample individually or stratified by feature class.

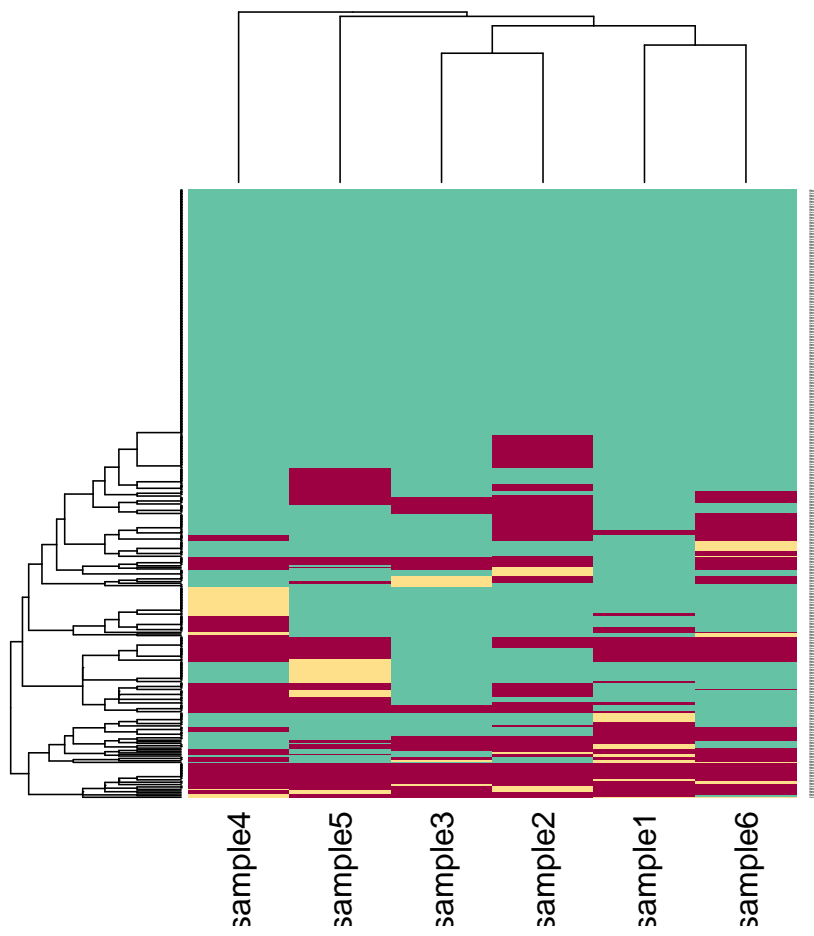


Figure 8: Summary of the categories, clustered across features.

If one doesn't want to include unreliable or undetermined data in part of the analysis, these Ct values can be set to NA using `filterCategory`. However, the presence of NAs could make the tests for differential expression less robust. When testing for differential expression the result will come with an associated category ("OK" or "Unreliable") that can instead be used to assess the quality of the results. For the final results both "Undetermined" and "Unreliable" are pooled together as being "Unreliable". However, the label for each feature can either be set according to whether half or more of the samples are unreliable, or whether only a single non-"OK" category is present, depending on the level of stringency the user wishes to enforce.

## 6 Normalisation

Five different normalisation methods are currently implemented in *HTqPCR*. Three of these (`scale.rankinvariant`, `deltaCt` and `geometric.mean`) will scale each individual sample by a given value, whereas the remaining two will change the distribution of Ct values.

**quantile** Will make the distribution of Ct values more or less identical across samples.

**norm.rankinvariant** Computes all rank-invariant sets of features between pairwise comparisons of each sample against a reference, such as a pseudo-mean. The rank-invariant features are used as a reference for generating a smoothing curve, which is then applied to the entire sample.

**scale.rankinvariant** Also computes the pairwise rank-invariant features, but then takes only the features found in a certain number of samples, and used the average Ct value of those as a scaling factor for correcting all Ct values.

**deltaCt** Calculates the standard deltaCt values, i.e. subtracts the mean of the chosen controls from all other values in the feature set.

**geometric.mean** Calculates the average Ct value for each sample, and scales all Ct values according to the ratio of these mean Ct values across samples. There are some indications that this is beneficial for e.g. miRNA studies ([Mestdagh et al., 2009](#)).

For the rank-invariant normalisation and geometric mean methods, Ct values above a given threshold can be excluded from the calculation of a scaling factor or normalisation curve. This is useful so that a high proportion of "Undetermined" Ct values (assigned a value of 40 by default) in a given sample doesn't bias the normalisation of the remaining features. In the example dataset, Gene1 and Gene60 correspond to 18S RNA and GADPH, and are used as endogenous controls. Normalisation methods can be run as follows:

```
> q.norm <- normalizeCtData(raw.cat, norm = "quantile")
> sr.norm <- normalizeCtData(raw.cat, norm = "scale.rank")

Scaling Ct values
  Using rank invariant genes: Gene1 Gene29
  Scaling factors: 1.00 1.06 1.00 1.03 1.00 1.00

> nr.norm <- normalizeCtData(raw.cat, norm = "norm.rank")

Normalizing Ct values
  Using rank invariant genes:
  sample1: 71 rank invariant genes
  sample2: 43 rank invariant genes
  sample3: 41 rank invariant genes
```

```

    sample4: 76 rank invariant genes
    sample5: 22 rank invariant genes
    sample6: 62 rank invariant genes
> d.norm <- normalizeCtData(raw.cat, norm = "deltaCt",
+   deltaCt.genes = c("Gene1", "Gene60"))
Calculating deltaCt values
    Using control gene(s): Gene1 Gene60
    Card 1:      Mean=14.45      Stdev=4.25
    Card 2:      Mean=15.19      Stdev=5.27
    Card 3:      Mean=14.5       Stdev=5.79
    Card 4:      Mean=14.79      Stdev=4.79
    Card 5:      Mean=14.07      Stdev=5.32
    Card 6:      Mean=13.82      Stdev=4.75
> g.norm <- normalizeCtData(raw.cat, norm = "geometric.mean")
Scaling Ct values
    Using geometric mean within each sample
    Scaling factors: 1.00 1.06 1.05 1.02 1.04 1.02

```

Comparing the raw and normalised values gives an idea of how much correction has been performed (Figure 9), as shown below for the `q.norm` object. Note that the scale on the y-axis varies.

```

> plot(exprs(raw), exprs(q.norm), pch = 20, main = "Quantile normalisation",
+   col = rep(brewer.pal(6, "Spectral"), each = 384))

```

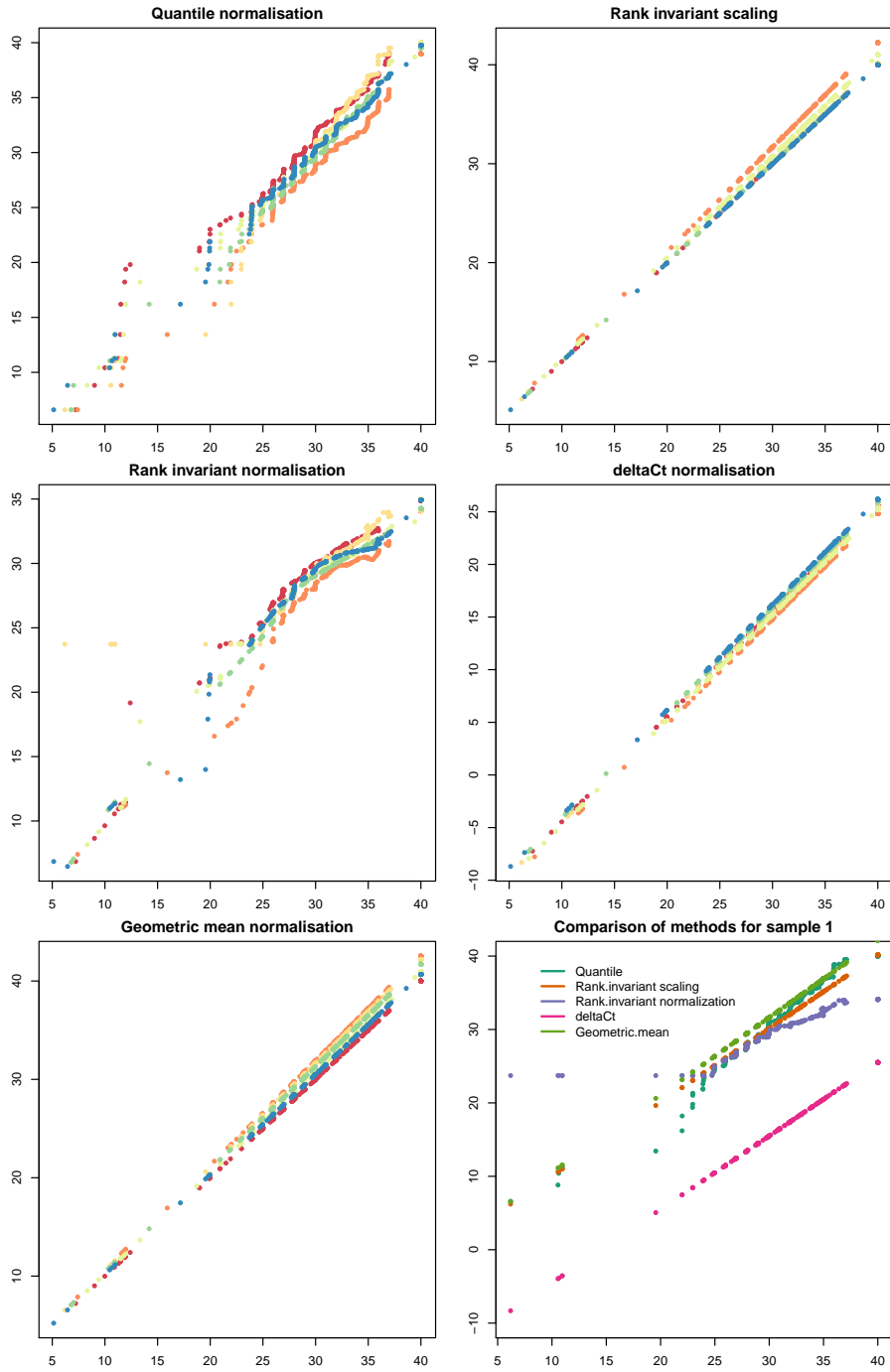


Figure 9: Normalized versus raw data, using a separate colour for each sample. The raw data is plotted along the x-axis and the normalised along y. The last plot is a comparison between normalization methods for the third sample, still with the raw Ct values along the x-axis.

## 7 Filtering and subsetting the data

At any point during the analysis it's possible to filter out both individual features or groups of features that are either deemed to be of low quality, or not of interest for a particular aspect of the analysis. This can be done using any of the feature characteristics that are included in the `featureNames`, `featureType`, `featureClass` and/or `featureCategory` slots of the data object. Likewise, the `qPCRset` object can be turned into smaller subsets, for example if only a particular class of features are to be used, or some samples should be excluded.

Simple subsetting can be done using the standard `[,]` notation of R, for example:

```
> nr.norm[1:10, ]
An object of class "qPCRset"
Size: 10 features, 6 samples
Feature types:      Endogenous Control, Target
Feature names:      Gene1 Gene2 Gene3 ...
Feature classes:    Kinase, Marker, TF
Feature categories: OK, Unreliable, Undetermined
Sample names:      sample1 sample2 sample3 ...

> nr.norm[, c(1, 3, 5)]
An object of class "qPCRset"
Size: 384 features, 3 samples
Feature types:      Endogenous Control, Target
Feature names:      Gene1 Gene2 Gene3 ...
Feature classes:    Kinase, Marker, TF
Feature categories: OK, Unreliable, Undetermined
Sample names:      sample1 sample3 sample5 ...
```

Filtering is done by specifying the components to remove, either by just using a single criteria, or by combining multiple filters:

```
> qFilt <- filterCtData(nr.norm, remove.type = "Endogenous Control")
Removed 4 'Endogenous Control' features based on featureType(q).

> qFilt <- filterCtData(nr.norm, remove.name = c("Gene1",
+       "Gene20", "Gene30"))
Removed 8 'Gene1/Gene20/Gene30' features based on featureNames(q).

> qFilt <- filterCtData(nr.norm, remove.class = "Kinase")
Removed 63 'Kinase' features based on featureClass(q).

> qFilt <- filterCtData(nr.norm, remove.type = c("Endogenous Control"),
+       remove.name = c("Gene1", "Gene20", "Gene30"))
Removed 4 'Endogenous Control' features based on featureType(q).
Removed 4 'Gene1/Gene20/Gene30' features based on featureNames(q).
```

The data can also be adjusted according to feature categories. With `filterCategory` mentioned previously it's possible to replace certain Ct values with NA, but one might want to completely exclude features where a certain number of the Ct values are for example unreliable.

```
> qFilt <- filterCtData(nr.norm, remove.category = "Undetermined")
Removed 64 features with >3 'Undetermined' based on featureCategory(q).
```

```
> qFilt <- filterCtData(nr.norm, remove.category = "Undetermined",
+   n.category = 5)
Removed 10 features with >5 'Undetermined' based on featureCategory(q).
```

Another typical filtering step would be to remove features showing little or no variation across samples, prior to testing for statistical significance of genes between samples. Features with relatively constant Ct levels are less likely to be differentially expressed, so including them in the downstream analysis would cause some loss of power when adjusting the p-values for multiple testing (the feature-by-feature hypothesis testing). Variation across samples can be assessed using for example the interquartile range (IQR) values for each feature (Figure 10).

```
> iqr.values <- apply(exprs(nr.norm), 1, IQR)
> hist(iqr.values, n = 20, main = "", xlab = "IQR across samples")
> abline(v = 1.5, col = 2)
```

All features with IQR below a certain threshold can then be filtered out.

```
> qFilt <- filterCtData(nr.norm, remove.IQR = 1.5)
Removed 212 features with IQR <1.5 based on exprs(q).
```

Note that filtering prior to normalisation can affect the outcome of the normalisation procedure. In some cases this might be desirable, for example if a particular feature class are heavily biasing the results so it's preferable to split the `qPCRset` object into smaller data sets. However, in other cases it might for example make it difficult to identify a sufficient number of rank invariant features for `thenorm.rankinvariant` and `scale.rankinvariant` methods. Whether to perform filtering, and if so then during what step of the analysis, depends on the genes and biological samples being analysed, as well as the quality of the data. It's therefore advisable to perform a detailed quality assessment and data comparison, as mentioned in the next section.

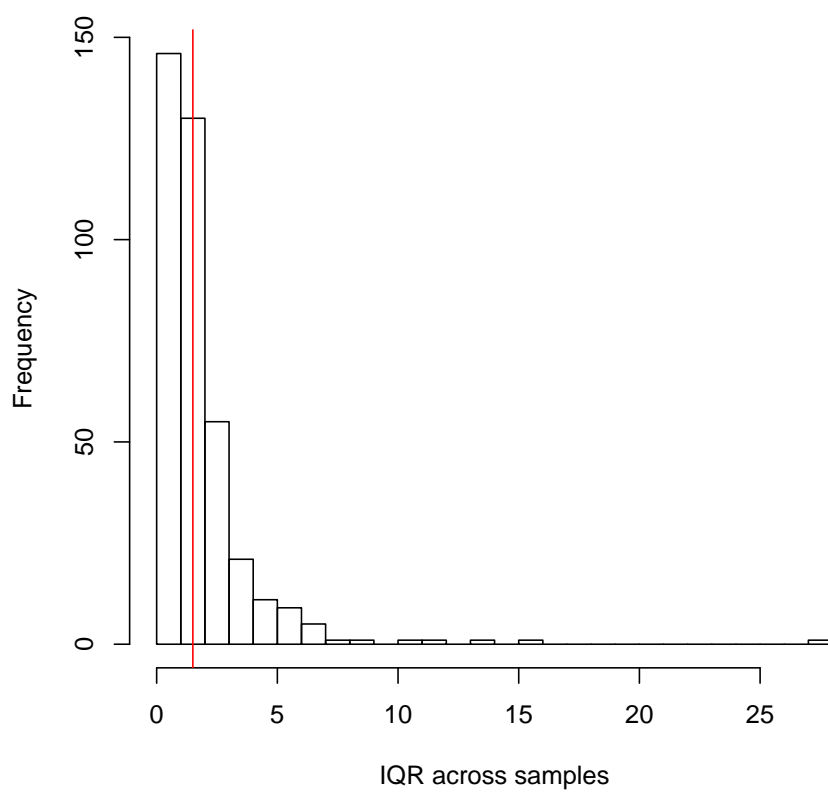


Figure 10: Histogram of the IQR values for all features across the samples, including the cut-off used in the filtering example.

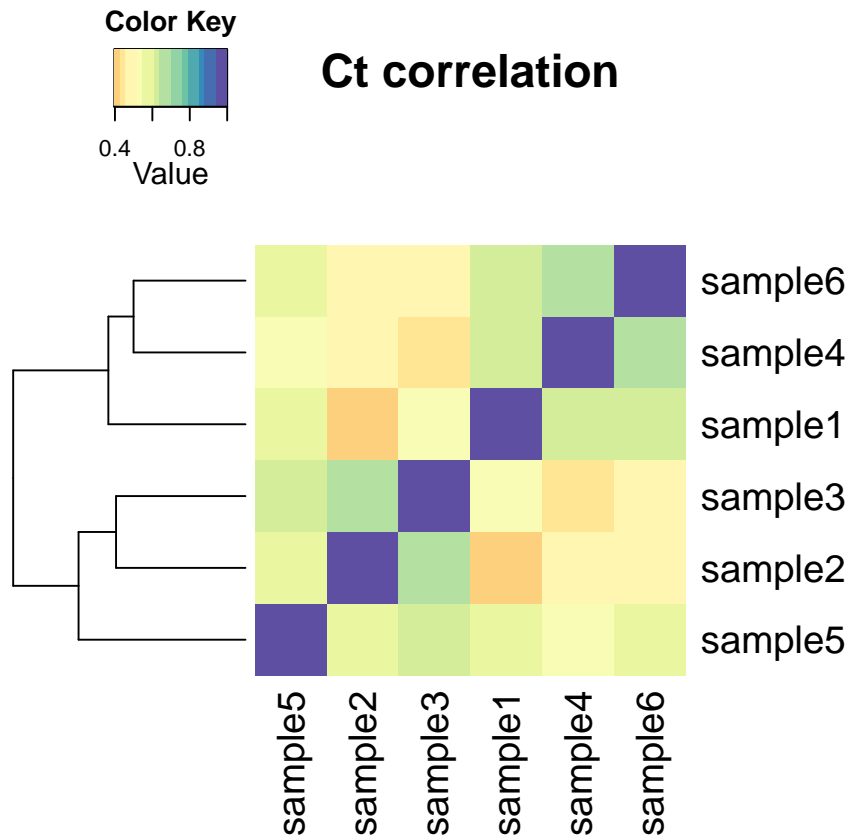


Figure 11: Correlation between raw Ct values.

## 8 Quality assessment

### 8.1 Correlation between samples

The overall correlation between different samples can be displayed visually, such as shown for the raw data in Figure 11.

```
> plotCtCor(raw, main = "Ct correlation")
```

### 8.2 Distribution of Ct values

It may be of interest to examine the general distribution of data both before and after normalisation. A simple summary of the data can be obtained using `summary` as shown below.

```
> summary(raw)
```

	sample1	sample2	sample3	sample4	sample5	sample6
Min.	" 7.218"	" 7.408"	" 6.19"	" 6.853"	" 6.787"	" 5.133"
1st Qu.	"26.738"	"28.855"	"27.90"	"26.964"	"27.913"	"27.514"
Median	"28.937"	"30.994"	"29.92"	"29.943"	"30.778"	"29.931"
Mean	"29.542"	"32.190"	"30.35"	"30.590"	"30.995"	"30.663"

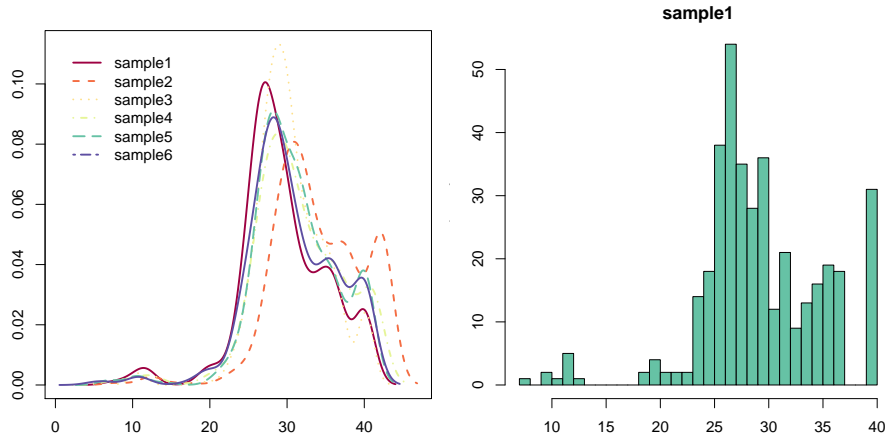


Figure 12: Distribution of Ct values for the individual samples, either using the density of all arrays (left) or a histogram of a single sample (right), after scale rank-invariant normalisation.

```
3rd Qu. "33.323" "35.985" "32.98" "34.694" "34.702" "35.046"
Max.    "40.000" "40.000" "40.00" "40.000" "40.000" "40.000"
```

However, figures are often more informative. To that end, the range of Ct values can be illustrated using histograms or with the density distribution, as shown in Figure 12.

```
> plotCtDensity(sr.norm)
```

```
> plotCtHistogram(sr.norm)
```

Plotting the densities of the different normalisation methods lends insight into how they differ (Figure 13).

Ct values can also be displayed in boxplots, either with one box per sample or stratified by different attributes of the features, such as `featureClass` or `featureType` (Fig. 14).

```
> plotCtBoxes(sr.norm, stratify = "class")
```

### 8.3 Comparison of Ct values for two samples

It is often of interest to directly compare Ct values between two samples. In Figure 15, two examples are shown for the rank-invariant normalised data: one for different biological samples, and one for replicates.

```
> plotCtScatter(sr.norm, cards = c(1, 2), col = "type",
+   diag = TRUE)

> plotCtScatter(sr.norm, cards = c(1, 4), col = "class",
+   diag = TRUE)
```

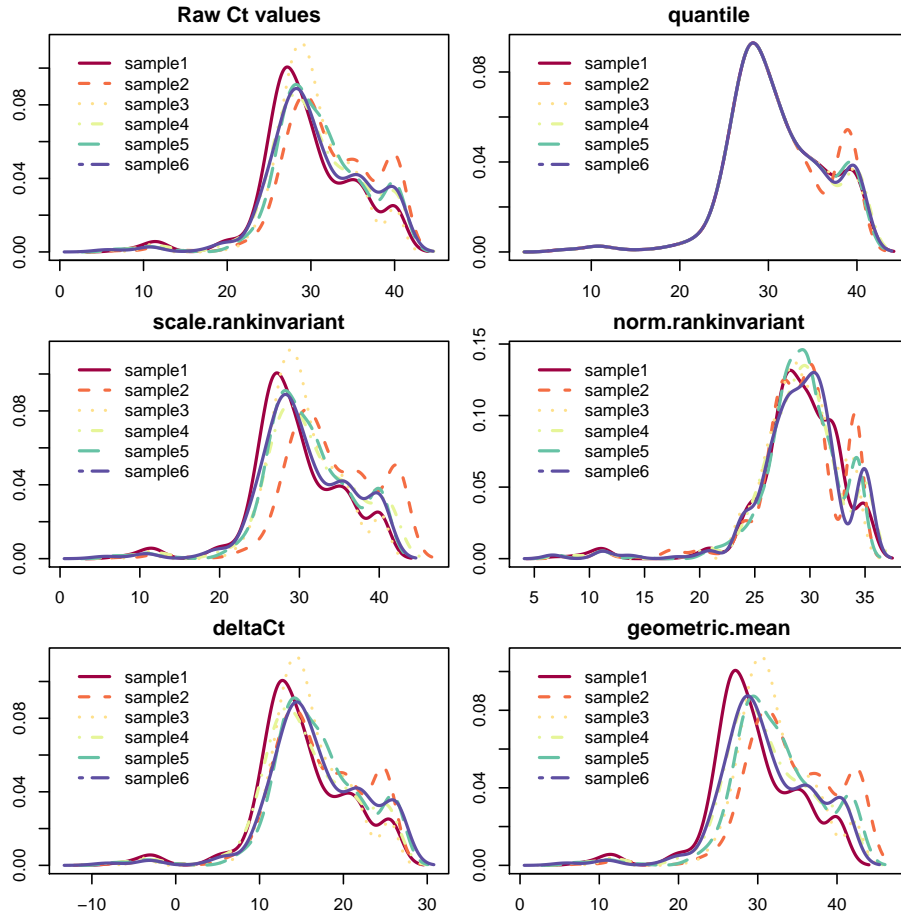


Figure 13: Densities of Ct values for all samples before and after each of the normalisation methods. The peak at the high end originates from features with “Undetermined” Ct values, which are assigned the Ct value 40 by default.

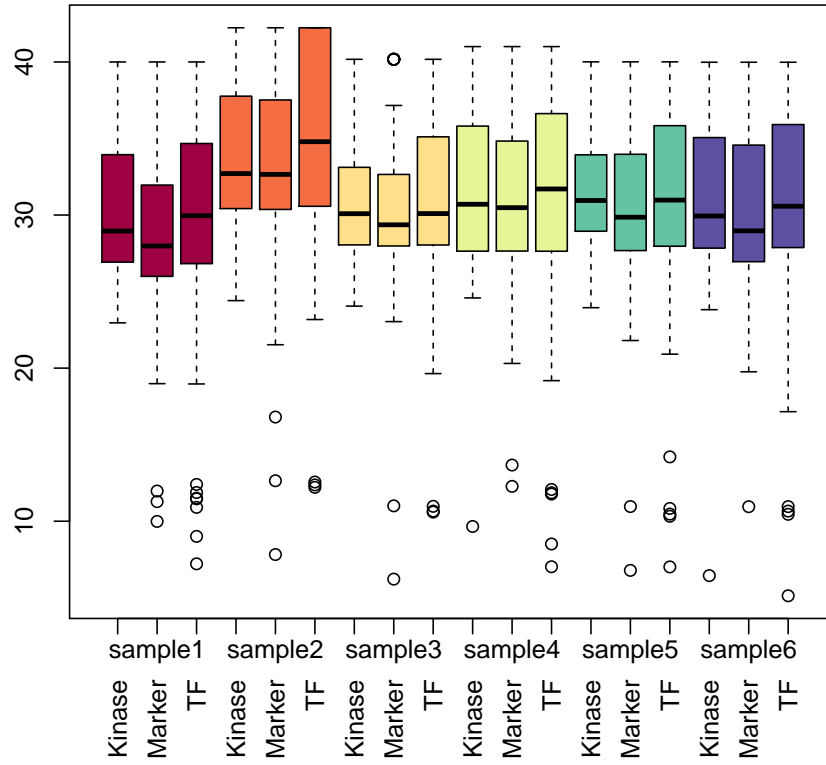


Figure 14: Boxplot of Ct values across all samples, stratified by feature classes.

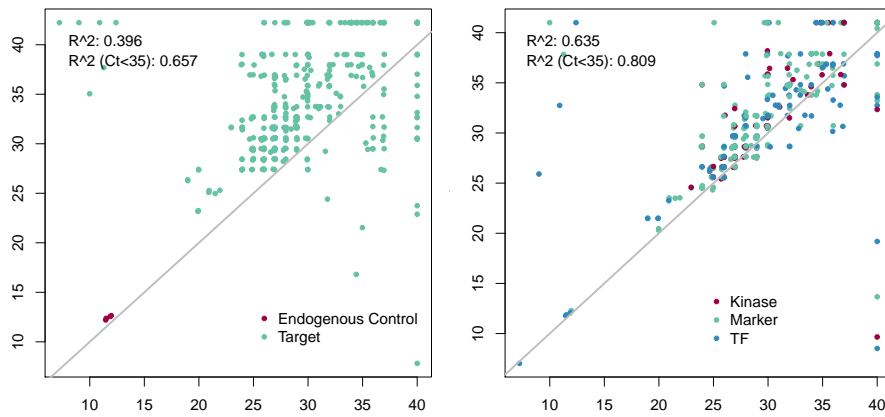


Figure 15: Scatter plot of Ct values in different samples, with points marked either by featureType (left) or featureClass (right) and the diagonal through  $x = y$  marked with a grey line.

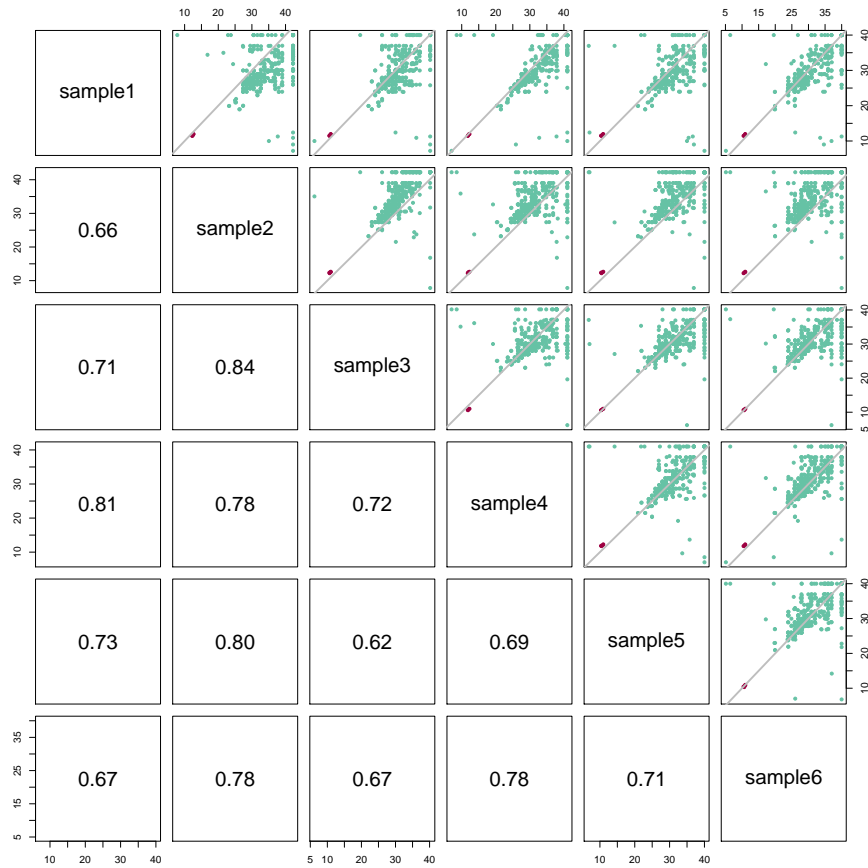


Figure 16: Scatterplot for all pairwise comparisons between samples, with spots marked depending on `featureType`, i.e. whether they represent endogenous controls or targets.

## 8.4 Scatter across all samples

It is also possible to generate a scatterplot of Ct values between more than the two samples shown above. In Figure 16 all pairwise comparisons are shown, along with their correlation when all Ct values <35 are removed.

```
> plotCtPairs(sr.norm, col = "type", diag = TRUE)
```

## 8.5 Ct heatmaps

Heatmaps provide a convenient way to visualise clustering of features and samples at the same time, and show the levels of Ct values (Figure 17). The heatmaps can be based on either Pearson correlation coefficients or Euclidean distance clustering. Euclidean-based heatmaps will focus on the magnitude of Ct values, whereas Pearson clusters the samples based on similarities between the Ct profiles.

```
> plotCtHeatmap(raw, gene.names = "", dist = "euclidean")
```

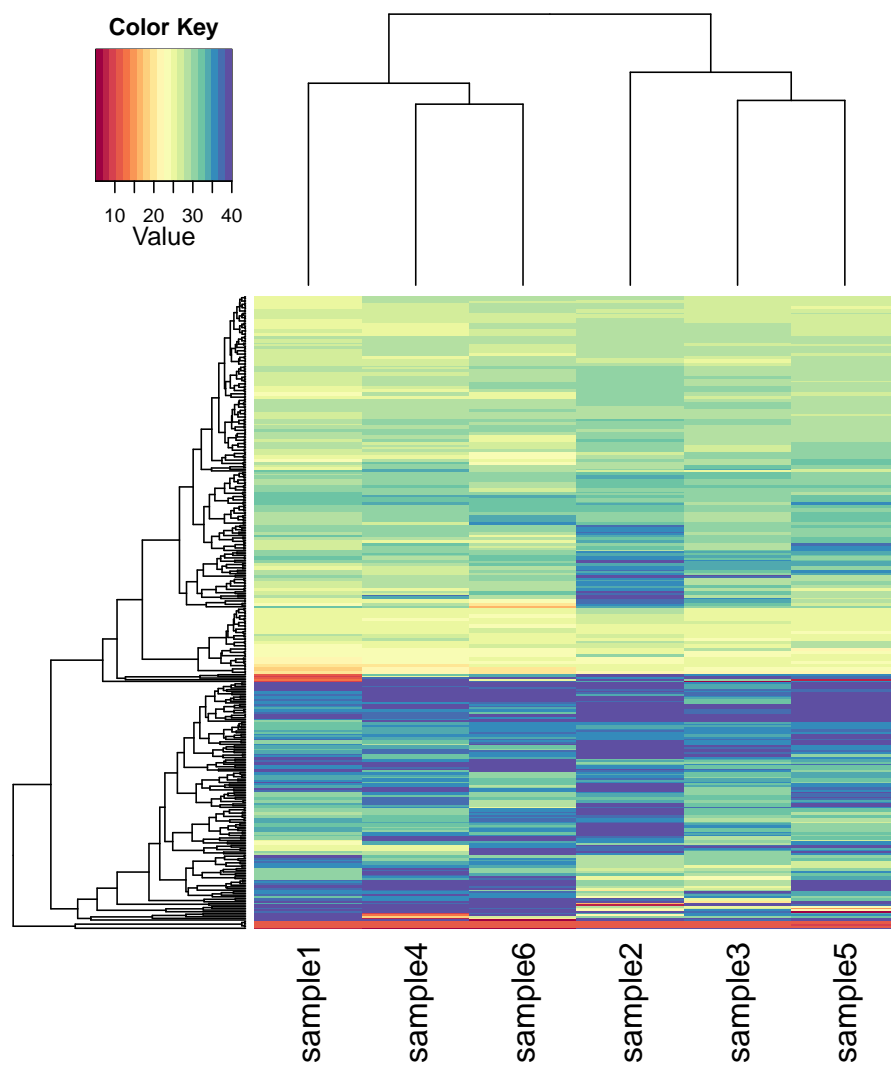


Figure 17: Heatmap for all samples and genes, based on the Euclidean distance between Ct values.

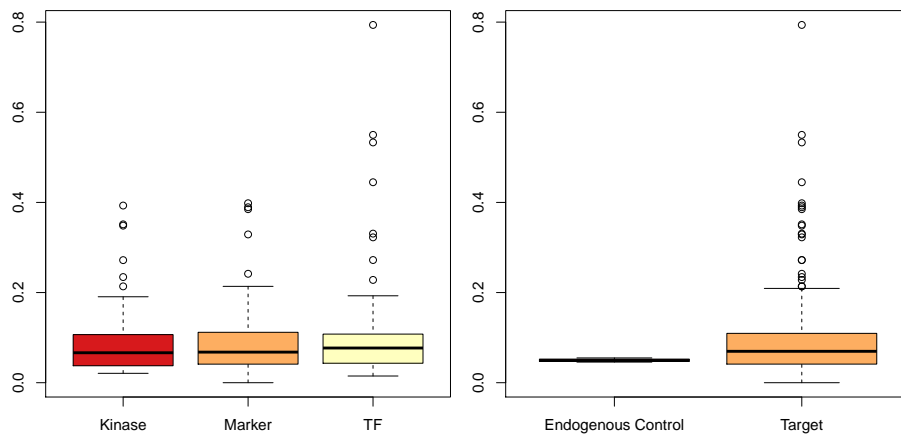


Figure 18: Coefficients of variation for each feature across all samples.

## 8.6 Coefficients of variation

The coefficients of variation (CV) can be calculated for each feature across all samples. Stratifying the CV values by `featureType` or `featureClass` can help to determine whether one class of features is more variable than another (Figure 18). For the example data feature classes have been assigned randomly, and the CVs are therefore similar, whereas for the feature types there's a clear difference between controls and targets.

```
> plotCVBoxes(qPCRraw, stratify = "class")
> plotCVBoxes(qPCRraw, stratify = "type")
```

## 9 Clustering

At the moment there are two default methods present in *HTqPCR* for clustering; hierarchical clustering and principal components analysis (PCA).

### 9.1 Hierarchical clustering

Both features and samples can be subjected to hierarchical clustering using either Euclidean or Pearson correlation distances, to display similarities and differences within groups of data. Individual subclusters can be selected, either using pre-defined criteria such as number of clusters, or interactively by the user. The content of each cluster is then saved to a list, to allow these features to be extracted from the full data set if desired.

An example of a clustering of samples is shown in Figure 19. In Figure 20 these data are clustered by features, and the main subclusters are marked.

```
> clusterCt(sr.norm, type = "samples")

> cluster.list <- clusterCt(sr.norm, type = "genes",
+   n.cluster = 6, cex = 0.5)

> c6 <- cluster.list[[6]]
> print(c6)
```

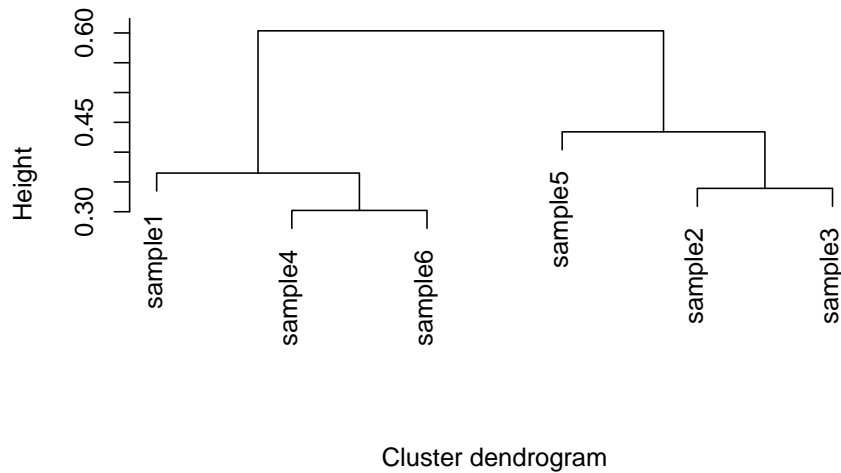


Figure 19: Hierarchical clustering of samples.

```

      Gene9  Gene13  Gene46  Gene50  Gene75  Gene93  Gene75  Gene103
           9      14      71      99     148     166     172     200
Gene132  Gene151  Gene151
      277      296      320
> show(sr.norm[c6, ])
An object of class "qPCRset"
Size:  11 features, 6 samples
Feature types:                Endogenous Control, Target
Feature names:                Gene9 Gene13 Gene46 ...
Feature classes:              Kinase, Marker, TF
Feature categories:           Undetermined, Unreliable, OK
Sample names:                 sample1 sample2 sample3 ...

```

## 9.2 Principal components analysis

PCA is performed across the selected features and samples (observations and variables), and can be visualized either in a biplot, or showing just the clustering of the samples (Figure 21).

```

> plotCtPCA(qPCRraw)
> plotCtPCA(qPCRraw, features = FALSE)

```

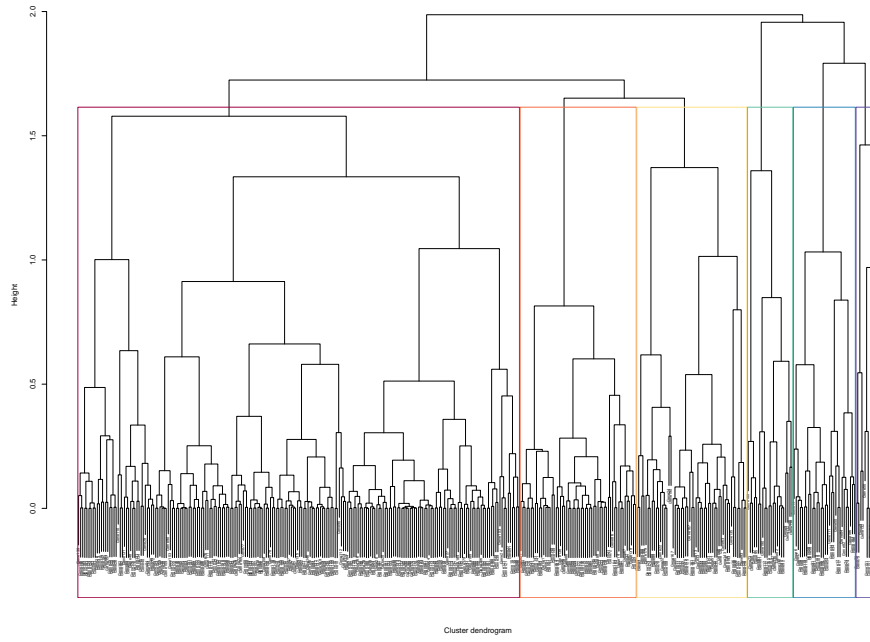


Figure 20: Hierarchical clustering of features, with subclusters marked.

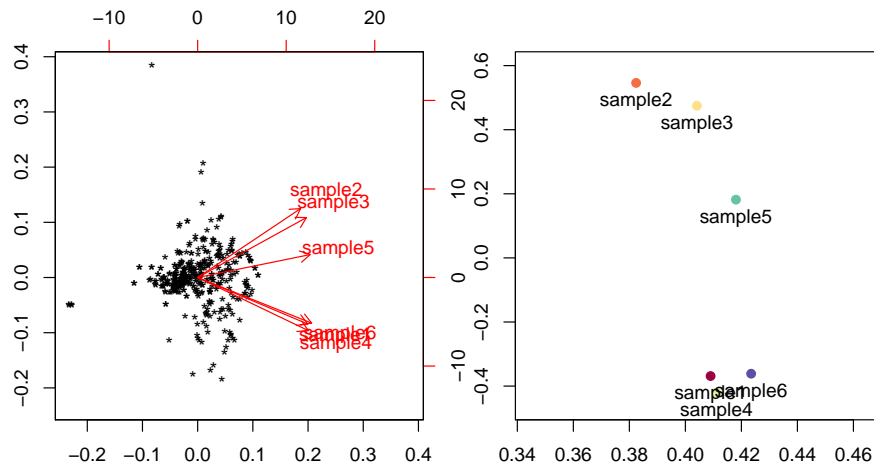


Figure 21: Left: a biplot including all features, with samples represented by vectors. Right: the same plot, including only the samples.

## 10 Differential expression

At this stage multiple filterings might have been performed on the data set(s). To remind yourself about those, you can use the `getCtHistory` function on the *qPCRset* object.

```
> getCtHistory(sr.norm)

                                history
1      readCtData(files = files$File, path = path)
2 setCategory(q = raw, quantile = 0.8, groups = files$Treatment)
3      normalizeCtData(q = raw.cat, norm = "scale.rank")
> getCtHistory(qFilt)

                                history
1      readCtData(files = files$File, path = path)
2 setCategory(q = raw, quantile = 0.8, groups = files$Treatment)
3      normalizeCtData(q = raw.cat, norm = "norm.rank")
4      filterCtData(q = nr.norm, remove.IQR = 1.5)
```

In general there are three approaches in *HTqPCR* for testing the significance of differences in Ct values between samples.

**t-test** Performing a standard t-test between two sample groups. This function will incorporate information about replicates to calculate t and p-values. This is a fairly simple approach, typically used when comparing a single treatment and control sample, and multiple pair-wise tests can be carried out one-by-one by the user.

**Mann-Whitney** This is a non-parametric test, also known as a two sample Wilcoxon test. Similar to the t-test, multiple pair-wise tests will have to be carried out one by one if more than two types of samples are present. This is a rank-based test that doesn't make any assumptions about the population distribution of Ct values.

**limma** Using a wrapper for functions from the package *limma* (Smyth, 2005) to calculate more sophisticated t and p-values for any number of groups present across the experiment. This is more flexible in terms of what types of comparisons can be made, but the users need to familiarise themselves with the *limma* conventions for specifying what contrasts are of interest.

Examples of how to use each of these are given in the next sections. In all cases the output is similar; a data frame containing the test statistics for each feature, along with fold change and information about whether the Ct values are “OK” or “Unreliable”. This result can be written to a file using standard functions such as `write.table`.

### 10.1 Two sample types - t-test

This section shows how to compare two samples, e.g. the control and long starvation samples from the example data. A subset of the *qPCRset* data is created to encompass only these samples, and a t-test is then performed.

```
> qDE.ttest <- ttestCtData(sr.norm[, 1:4], groups = files$Treatment[1:4],
+   calibrator = "Control")
> head(qDE.ttest)
```

	genes	feature.pos	t.test	p.value	adj.p.value
2	Gene10	A10;B10	-13.705640	9.812948e-06	0.001874273
22	Gene118	I23;J23	-10.407281	8.661980e-05	0.008272191
36	Gene130	K11;L11	7.644065	3.156334e-04	0.019863762
164	Gene74	G3;H3	7.201124	4.768225e-04	0.019863762
30	Gene125	K6;L6	-7.203614	5.199938e-04	0.019863762
33	Gene128	K9;L9	-10.357650	6.826374e-04	0.020708859

	ddCt	FC	meanCalibrator	meanTarget
2	3.460492	9.084231e-02	24.23106	27.69156
22	8.982157	1.977430e-03	26.70531	35.68746
36	-9.433460	6.914400e+02	39.74414	30.31068
164	-5.178293	3.620942e+01	35.40699	30.22869
30	7.052337	7.534164e-03	28.28445	35.33679
33	5.210558	2.700635e-02	25.61542	30.82598

	categoryCalibrator	categoryTarget
2	Undetermined	OK
22	OK	Undetermined
36	Undetermined	OK
164	Undetermined	OK
30	OK	OK
33	OK	OK

## 10.2 Two sample types - Mann-Whitney

When only two samples are of interest, these can also be compared using a Mann-Whitney test. As in the section above, the function is demonstrated using the control and long starvation samples from the example data. A subset of the `qPCRset` data is created to encompass only these samples, and a Mann-Whitney test is performed.

```
> qDE.mwtest <- mannwhitneyCtData(sr.norm[, 1:4], groups = files$Treatment[1:4],
+   calibrator = "Control")
> head(qDE.mwtest)
```

	genes	feature.pos	MW.value	p.value	adj.p.value
26	Gene121	K2;L2	W = 0	0.02746864	0.07842053
134	Gene47	C24;D24	W = 16	0.02842954	0.07842053
157	Gene68	E21;F21	W = 0	0.02842954	0.07842053
185	Gene93	G22;H22	W = 0	0.02842954	0.07842053
36	Gene130	K11;L11	W = 16	0.02940105	0.07842053
54	Gene147	M4;N4	W = 0	0.02940105	0.07842053

	ddCt	FC	meanCalibrator	meanTarget
26	2.316925	2.006947e-01	38.88749	41.20441
134	-12.205480	4.722973e+03	40.50267	28.29719
157	6.547033	1.069416e-02	34.65738	41.20441
185	25.515391	2.084980e-08	15.68902	41.20441
36	-9.433460	6.914400e+02	39.74414	30.31068
54	3.765620	7.352505e-02	35.92123	39.68685

	categoryCalibrator	categoryTarget
26	Undetermined	Undetermined
134	Undetermined	OK
157	Undetermined	Undetermined

185	Undetermined	Undetermined
36	Undetermined	OK
54	Undetermined	Undetermined

### 10.3 Multiple sample types - limma

In this example all three types of treatment are compared, as well as the control against both starvation samples combined. The data is sorted by feature names, to make easier use of replicated features.

```
> design <- model.matrix(~0 + files$Treatment)
> colnames(design) <- c("Control", "LongStarve", "Starve")
> print(design)
      Control LongStarve Starve
1           1           0      0
2           0           1      0
3           0           1      0
4           1           0      0
5           0           0      1
6           0           0      1
attr(,"assign")
[1] 1 1 1
attr(,"contrasts")
attr(,"contrasts")$`files$Treatment`
[1] "contr.treatment"

> contrasts <- makeContrasts(LongStarve - Control,
+   LongStarve - Starve, Starve - Control, (Starve +
+   LongStarve)/2 - Control, levels = design)
> colnames(contrasts) <- c("LS-C", "LS-S", "S-C", "bothS-C")
> print(contrasts)
              Contrasts
Levels      LS-C LS-S S-C bothS-C
Control     -1    0 -1    -1.0
LongStarve   1    1  0     0.5
Starve       0   -1  1     0.5

> sr.norm2 <- sr.norm[order(featureNames(sr.norm)),
+   ]
> qDE.limma <- limmaCtData(sr.norm2, design = design,
+   contrasts = contrasts, ndups = 2, spacing = 1)
```

The result is a list with one component per comparison. Each component is similar to the result from using `ttestCtData`.

```
> class(qDE.limma)
[1] "list"
> names(qDE.limma)
[1] "LS-C"    "LS-S"    "S-C"     "bothS-C" "Summary"
> head(qDE.limma[["LS-C"]])
```

	genes	feature.pos	t.test	p.value	adj.p.value
14	Gene11	A12;B12	7.603024	1.999665e-05	0.003839357
50	Gene142	K23;L23	6.271023	9.883760e-05	0.009488409
3	Gene10	A10;B10	5.076932	5.031756e-04	0.024152428
119	Gene32	C9;D9	5.217172	4.113509e-04	0.024152428
23	Gene118	I23;J23	4.812232	7.417071e-04	0.028481551
133	Gene45	C22;D22	4.195002	1.905657e-03	0.060981013

	ddCt	FC	meanTarget	meanCalibrator
14	6.051791	0.0150740287	26.23181	20.18002
50	10.752992	0.0005794645	36.48566	25.73267
3	3.460492	0.0908423086	27.69156	24.23106
119	5.645620	0.0199755612	31.37832	25.73270
23	8.982157	0.0019774303	35.68746	26.70531
133	6.108469	0.0144933090	34.42031	28.31184

	categoryTarget	categoryCalibrator
14	OK	Undetermined
50	Undetermined	OK
3	OK	Undetermined
119	OK	OK
23	Undetermined	OK
133	OK	OK

Furthermore, there is a “Summary” component at the end where each feature is denoted with -1, 0 or 1 to respectively indicate down-regulation, no change, or up-regulation in each of the comparisons.

```
> qDE.limma[["Summary"]][21:30, ]
```

	Contrasts			
	LS-C	LS-S	S-C	bothS-C
Gene116	0	0	0	0
Gene117	0	0	0	0
Gene118	1	0	0	0
Gene119	0	0	0	0
Gene12	0	0	0	0
Gene120	0	0	0	0
Gene121	0	0	0	0
Gene122	0	0	0	0
Gene123	0	0	0	0
Gene124	0	0	0	0

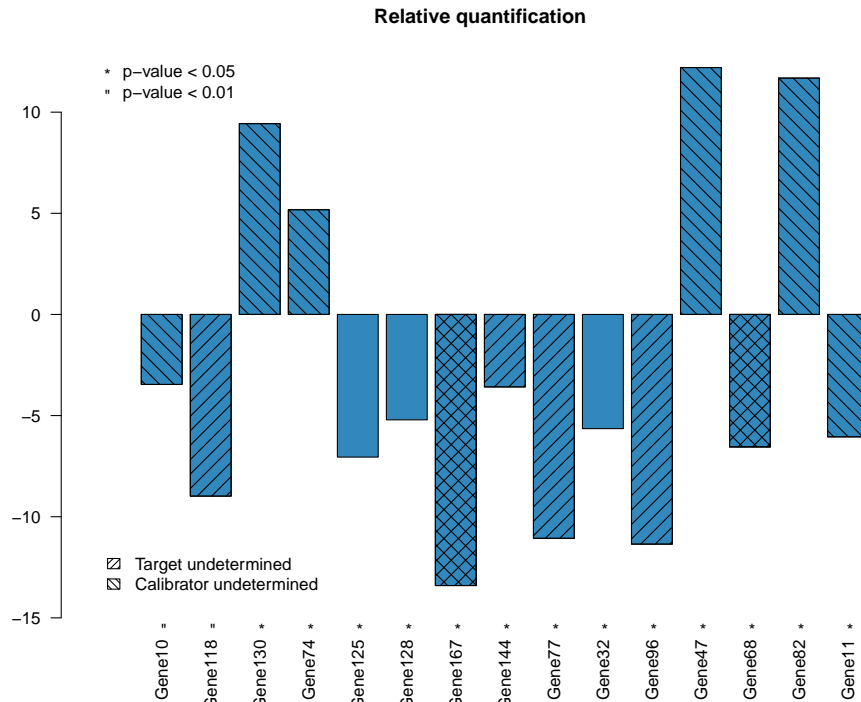


Figure 22: Relative quantification, using the top 15 features from `ttestCtData`.

## 11 Displaying the results

The results can be visualised using the generic `plotCtOverview` shown in Figure 3. However, *HTqPCR* also contains more specialised functions, for example to include information about whether differences are significant or not.

### 11.1 Fold changes: Relative quantification

The relative Ct levels between two groups can be plotted with the function `plotCtRQ`. Below are two examples: one the result of `ttestCtData` where the top 15 genes are selected (figure 22), and another from the first comparison in `limmaCtData` where all genes below a certain p-value are depicted (Figure 23).

```
> plotCtRQ(qDE.ttest, genes = 1:15)

> plotCtRQ(qDE.limma, p.val = 0.085, transform = "log10",
+         col = "#9E0142")
```

The hatching on the bars indicates whether the target and calibrator Ct samples were unreliable, but this also depend on whether the parameter `stringent=TRUE` or `stringent=FALSE` when testing for differential expression. See the help functions for details (`?ttestCtData` and `?limmaCtData`).

### 11.2 Fold changes: Detailed visualisation

In some cases it will be beneficial to more closely examine individual Ct data points from the fold changes, partly to look at the data dispersion, and partly to determine which of

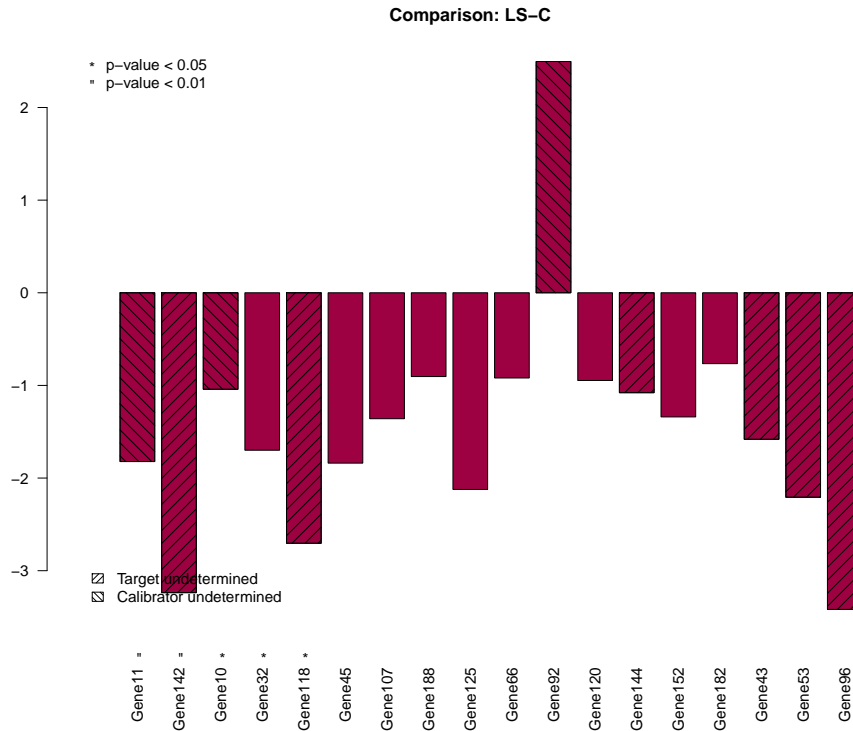


Figure 23: Relative quantification, using all features with p-value<0.085 from limmaCt-Data.

these values are in the “OK” versus “Unreliable”/“Undetermined” category. The function `plotCtSignificance` will take the result of `ttestCtData` or `limmaCtData`, along with the input data to these functions, and display a combined barplot showing the individual data points and marking those comparisons with a significant p-value.

```
> plotCtSignificance(qDE.limma, q = sr.norm, groups = files$Treatment,
+   target = "LongStarve", calibrator = "Control",
+   genes = featureNames(sr.norm)[11:20], un.col = "#3288BD",
+   jitter = 0.2)
```

### 11.3 Heatmap across comparisons

When multiple conditions are compared with `limmaCtData`, the fold changes from all comparisons can be compared to see if features cluster together in groups (Figure 25).

```
> heatmapSig(qDE.limma, dist = "euclidean")
```

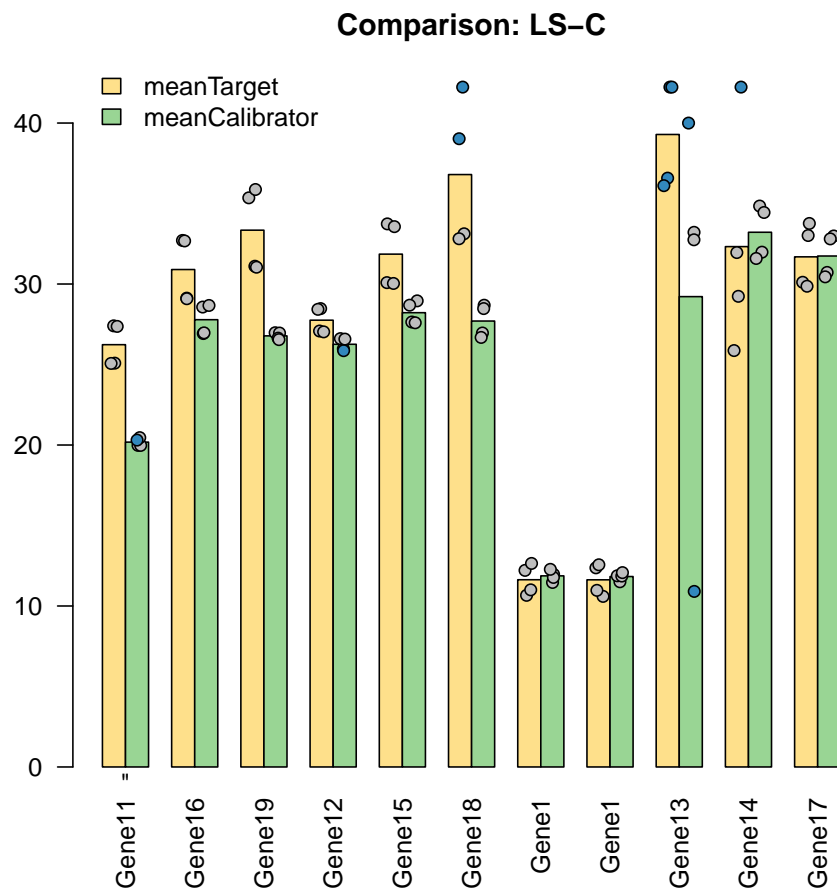


Figure 24: Ten genes from the data set, with the average Ct in two groups plotted along with the individual values. Points marked in blue are “Unreliable” or “Undetermined” whereas grey spots are “OK”.

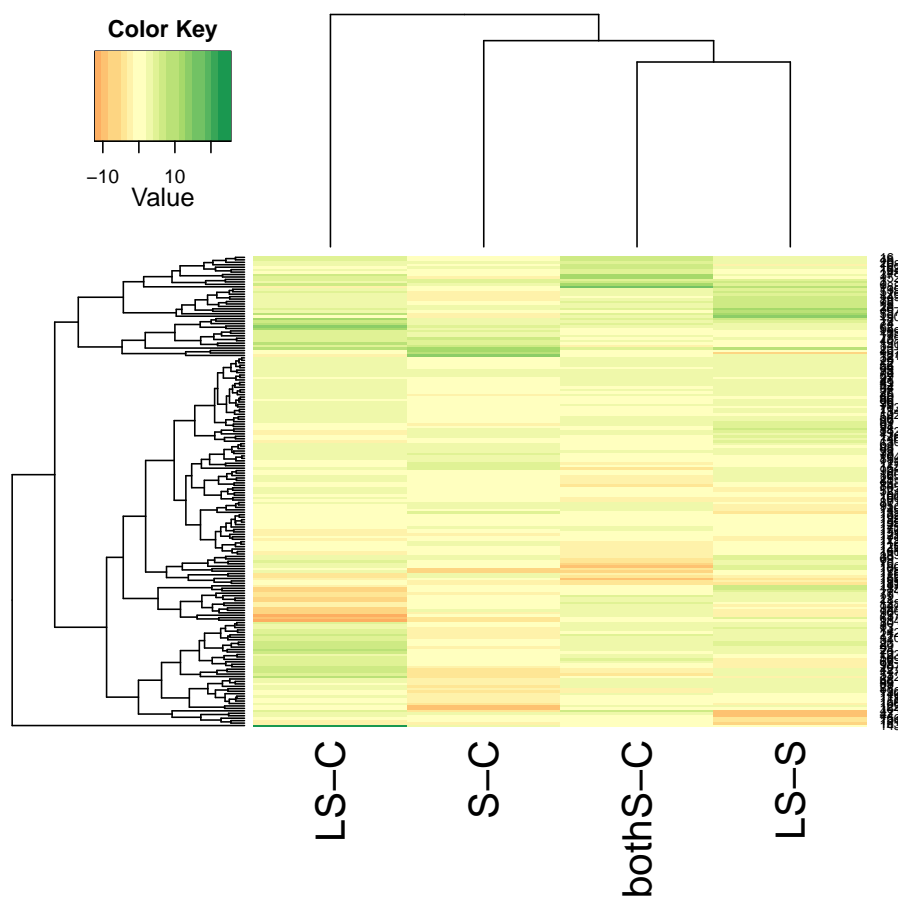


Figure 25: Fold changes across all comparisons, clustered based on Euclidean distance.

## 12 Manipulating qPCRset objects

Depending on the design of the qPCR card and how the data is to be analysed, it will sometimes be necessary to manipulate the `qPCRset` objects in different ways, such as by combining several objects or altering the layout of features x samples.

### 12.1 Multiple samples present on each plate

The result from each qPCR run of a given card typically gets presented together, such as in a file with 384 lines, one per feature, for 384 well plates. However, some cards may contain multiple samples, such as commercial cards that are designed to be loaded with two separate samples and then include 192 individual features.

Per default, `readCtData` reads each card into a *qPCRset* object as consisting of a single sample, and hence one column in the Ct data matrix. When this is not the case, the data can subsequently be split into the correct features x samples (rows x columns) dimensions using the function `changeCtLayout`. The parameter `sample.order` is a vector, that for each feature in the *qPCRset* indicates what sample it actually belongs to, and reformats all the information in the *qPCRset* (`exprs`, `featureCategory`, `flag` etc.) accordingly. The actual biological samples are likely to differ on each card, so `sample.order` merely indicates the *location* of different samples among the features present in the input data.

```
> sample2.order <- rep(c("subSampleA", "subSampleB"),
+   each = 192)
> sample4.order <- rep(c("subA", "subB", "subC", "subD"),
+   each = 96)
> qPCRnew2 <- changeCtLayout(sr.norm, sample.order = sample2.order)
> show(qPCRnew2)

An object of class "qPCRset"
Size: 192 features, 12 samples
Feature types:      Endogenous Control, Target
Feature names:      Gene1 Gene2 Gene3 ...
Feature classes:    Kinase, Marker, TF
Feature categories: OK, Unreliable, Undetermined
Sample names:      subSampleA:sample1 subSampleA:sample2 subSampleA:sample3 .
> qPCRnew4 <- changeCtLayout(sr.norm, sample.order = sample4.order)
> show(qPCRnew4)

An object of class "qPCRset"
Size: 96 features, 24 samples
Feature types:      Endogenous Control, Target
Feature names:      Gene1 Gene2 Gene3 ...
Feature classes:    Kinase, Marker, TF
Feature categories: OK, Unreliable, Undetermined
Sample names:      subA:sample1 subA:sample2 subA:sample3 ...
```

As with the other functions that manipulate the Ct data, the operation is stored in the history slot of the *qPCRset* for future reference.

```
> getCtHistory(qPCRnew4)

                                history
1      readCtData(files = files$File, path = path)
2 setCategory(q = raw, quantile = 0.8, groups = files$Treatment)
```

```

3           normalizeCtData(q = raw.cat, norm = "scale.rank")
4 changeCtLayout(q = sr.norm, sample.order = sample4.order)

```

## 12.2 Combining multiple qPCRset objects

In some cases it might be desirable to merge multiple qPCRset objects, that have been read into R or processed individually. The *HTqPCR* package contains two functions for combining multiple qPCRset objects into one, by either adding columns (samples) or rows (features). This can be done for either identical samples across multiple different cards (such as a 384 well plate), or if more samples have been run on cards with the same layout.

**cbind** combines data assuming that all experiments have been carried out on identical cards, i.e. that **featureNames**, **featureType**, **featurePos** and **featureClass** is identical across all the qPCRset objects. The number of features in each object must be identical, but number of samples can vary.

**rbind** combines data assuming that the same samples have been analysed using different qPCR cards. The number of samples in each object must be identical, but the number of features can vary.

Both these functions should be used with some care; consider e.g. whether to normalize before or after joining the samples, and what method to use.

In the examples here objects with different normalisation are combined, although in a real study the qPCRset objects would typically contain different data.

```

> q.comb <- cbind(q.norm[, 1:3], sr.norm[, 4], nr.norm[,
+               c(1, 5, 6)])
> q.comb
An object of class "qPCRset"
Size: 384 features, 7 samples
Feature types:           Endogenous Control, Target
Feature names:           Gene1 Gene2 Gene3 ...
Feature classes:         Kinase, Marker, TF
Feature categories:      OK, Unreliable, Undetermined
Sample names:            sample1 sample2 sample3 ...
> q.comb2 <- rbind(q.norm, sr.norm[1:4, ], nr.norm)
> q.comb2
An object of class "qPCRset"
Size: 772 features, 6 samples
Feature types:           Endogenous Control, Target
Feature names:           Gene1 Gene2 Gene3 ...
Feature classes:         Kinase, Marker, TF
Feature categories:      OK, Unreliable, Undetermined
Sample names:            sample1 sample2 sample3 ...

```

As with other functions where the qPCRset object is being manipulated, the information is stored in the **history** slot.

```

> getCtHistory(q.comb)

```

```

                                                                    history
1          qPCRset1: readCtData(files = files$File, path = path)
2  qPCRset1: setCategory(q = raw, quantile = 0.8, groups = files$Treatment)
3          qPCRset1: normalizeCtData(q = raw.cat, norm = "quantile")
4          qPCRset2: readCtData(files = files$File, path = path)
5  qPCRset2: setCategory(q = raw, quantile = 0.8, groups = files$Treatment)
6          qPCRset2: normalizeCtData(q = raw.cat, norm = "scale.rank")
7          qPCRset3: readCtData(files = files$File, path = path)
8  qPCRset3: setCategory(q = raw, quantile = 0.8, groups = files$Treatment)
9          qPCRset3: normalizeCtData(q = raw.cat, norm = "norm.rank")
10         cbind(deparse.level, ..1, ..2, ..3)

```

## 13 How to handle different input data

General information about how to read qPCR data into a *qPCRset* object is presented in section 3. Below, some more specific cases are illustrated.

### 13.1 Sequence Detection Systems format

The qPCR data might come from Sequence Detection Systems (SDS) software, in which case each file has a header containing some generic information about the initial Ct detection. This header varies in length depending on how many files were analysed simultaneously, and an example is shown below.

```
> path <- system.file("exData", package = "HTqPCR")
> cat(paste(readLines(file.path(path, "SDS_sample.txt"),
+   n = 19), "\n"))
```

```
SDS 2.3 RQ Results          1.2
Filename      Testscreen analys all.sdm
Assay Type      RQ Study
EmbeddedFile    FileA
Run DateTime    Fri May 15 17:10:28 BST 2009
Operator
ThermalCycleParams
EmbeddedFile    FileB
Run DateTime    Sat May 16 10:36:09 BST 2009
Operator
ThermalCycleParams
EmbeddedFile    FileC
Run DateTime    Sun May 17 13:21:05 BST 2009
Operator
ThermalCycleParams
```

#	Plate	Pos	Flag	Sample	Detector	Task
1	Control	A1	Passed	Sample01	Gene1	Endogenous
2	Control	A2	Passed	Sample01	Gene2	Target

Only the first 7 columns are shown, since the file shown here contains >30 columns (of which many are empty). All columns for the first 20 lines can be seen in an R terminal with the command:

```
> readLines(file.path(path, "SDS_sample.txt"), n = 20)
```

For these files the parameter `SDS=TRUE` can be set in `readCtData`. The first 100 lines of each file will be scanned, and all lines preceding the actual data will be skipped (in this case 17), even when the length of the header varies between files.

### 13.2 Non-Ct data

Some qPCR systems, such as the LightCycler from Roche, don't provide the results as Ct values, but instead as crossing points (Cp). Ct values are measured at the exponential phase of amplification by drawing a line parallel to the x-axis of the real-time fluorescence intensity curve (fit point method), whereas Cp (second derivative method) calculates the fractional cycle where the second derivative of the real-time fluorescence intensity curve reaches the maximum value (Luu-The et al., 2005).

As long as all samples and features are quantified using the same method, it shouldn't matter whether Ct or Cp values are being used to test for significant differences between samples. The analysis of e.g. LightCycler Cp data can therefore proceed as outlined in this vignette. One thing to bear in mind though, is that for Ct values a value of 40 generally means NA, and above 35 is considered unreliable, however these numbers will be different for Cp. When using the filtering methods to set results as being "OK", "Unreliable" and "Undetermined", the parameters in e.g. `setCategory(..., Ct.max = 35, Ct.min = 10, ...)` and `readCtData(..., na.value, ...)` will need to be adjusted accordingly.

### 13.3 Microfluidic devices

In addition to multi-well microtitre plates, other assay formats have also emerged for performing high-throughput qPCR analysis. These include for example the 48.48 and 96.96 Dynamic Array from Fluidigm Corporation. For the 48.48 assay, 48 individual samples are loaded onto a plate along with qPCR primers for 48 genes. Using microfluidic channels all possible sample  $\times$  primer reactions are assayed in a combinatorial manner using 2,304 individual reaction chambers, to generate e.g. 48 real-time curves for each of 48 samples. Results are then reported in a single file with 2304 rows of data.

*HTqPCR* includes a comma-separated file containing example data from a Fluidigm 48.48 array. Since the data format differs slightly from some other qPCR platforms, it'll currently need to be transformed into a `qPCRset` object in one of two ways. First, `qPCRset` can be constructed indirectly, by reading the data into a data frame, creating a 48x48 matrix manually, and generating a new `qPCRset`. The file was inspected first, and turned out to have an 11 line header plus a header line, which have to be skipped when reading the data into *HTqPCR*.

```
> exPath <- system.file("exData", package = "HTqPCR")
> exFiles <- "fluidigm_sample.csv"
> temp <- read.delim(file.path(exPath, exFiles), skip = 11,
+   sep = ",", colClasses = "character")
> n <- 48
> mat <- matrix(as.numeric(temp$Value), ncol = n, nrow = n,
+   byrow = FALSE)
> mat[mat > 40] <- NA
> raw <- new("qPCRset", exprs = mat, sampleNames = paste("S",
+   1:n, sep = ""), featureNames = paste("A", 1:n,
+   sep = ""), featureCategory = as.data.frame(array("OK",
+   c(n, n))))
```

Alternatively, the data can be read directly into a `qPCRset` in a 1x2304 format using `readCtData`, and then re-formatted into 48x48. This will automatically read in additional information such as feature names, positions and categories (Failed/Passed) if available.

```
> temp <- readCtData(exFiles, path = exPath, n.features = 48 *
+   48, flag = 9, feature = 5, type = 6, Ct = 7,
+   position = 1, skip = 12, sep = ",")
> raw <- changeCtLayout(temp, sample.order = rep(1:48,
+   each = 48))
```

Information from multiple microfluidic devices can then subsequently be combined using `cbind` and `rbind`. If array-specific effects are likely to be present, a useful normalisation strategy might be to combine the data into a `qPCRset` object containing one column for

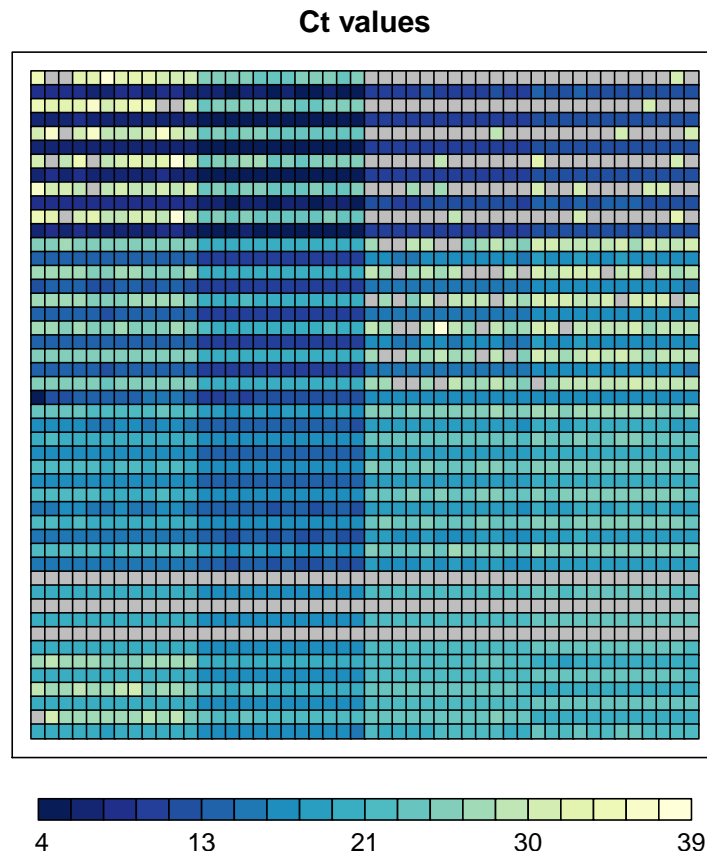


Figure 26: Ct values for a test Fluidigm 48.48 Array. 48 samples are loaded into rows, and 48 PCR primers into columns, resulting in 2,304 combinatorial qPCR reactions. Four individual genes were added in 12 replicates each into the 48 columns to assess technical variability. Grey corresponds to “NA”.

each array, and 48x48 or 96x96 samples. That way overall differences between the arrays can be removed, and afterwards the data can be split with `changeCtLayout` to generate one column per individual sample rather than one column per array.

The assay readouts are Ct values, which can be analysis using HTqPCR in a similar fashion to Ct values from other technologies, both regarding the statistical analysis and data visualisation. The function `plotCtArray` can be used for displaying the layout of the qPCR results instead of `plotCtCard` (Figure 26).

```
> plotCtArray(raw)
```

Functions specifically designed for such qPCR platforms are still on a test stage in *HTqPCR*, and will be expanded depending on demand.

## 14 Concluding remarks

Use the function `news` to see what bug fixes and updates have been incorporated into *HTqPCR*. For example:

```
> news(Version > 1.3, package = "HTqPCR")
```

Changes in version 1.7:

### SIGNIFICANT USER-VISIBLE CHANGES

- o

### NEW FEATURES

- o Added "geometric.mean" as a normalisation method.

### BUG FIXES

- o Check for all-NA features in `normalizeCtData` rank-invariant methods.
- o Modified `plotCtCategory` to handle cases with only 1 sample.
- o Modified `plotCtCategory` to make colours more consistent.
- o Fix in `deltaCt` for normalization with NAs
- o Added warning in `readCtData` if there are NAs in input

Changes in version 1.5:

### SIGNIFICANT USER-VISIBLE CHANGES

- o Made small adjustments to `readCtData` to make it more suitable for data from the Fluidigm platform.

### NEW FEATURES

- o Added function `plotCtVariation` for displaying variation across replicated features or samples.
- o Added function `plotCtArray` for visualising data from non-well plates (e.g. the Fluidigm platform)

### BUG FIXES

- o Fixed bug in `readCtData` causing repeats of `featureType` and `featurePos` when `n.data>1`.
- o Corrected bugs for subsetting when `flag()` or `featureCategory()` are missing.

- o Modified "history" slot for manually created qPCRset objects.

This vignette was generated using:

- R version 2.14.0 (2011-10-31), i386-pc-mingw32
- Locale: LC\_COLLATE=C, LC\_CTYPE=English\_United States.1252, LC\_MONETARY=English\_United States.1252, LC\_NUMERIC=C, LC\_TIME=English\_United States.1252
- Base packages: base, datasets, grDevices, graphics, methods, stats, utils
- Other packages: Biobase 2.14.0, HTqPCR 1.8.0, RColorBrewer 1.0-5, limma 3.10.0, statmod 1.4.13
- Loaded via a namespace (and not attached): BiocInstaller 1.2.0, affy 1.32.0, affyio 1.22.0, gdata 2.8.2, gplots 2.10.1, gtools 2.6.2, preprocessCore 1.16.0, tools 2.14.0, zlibbioc 1.0.0

## References

- R. C. Gentleman, V. J. Carey, D. M. Bates, B. Bolstad, M. Dettling, S. Dudoit, B. Ellis, L. Gautier, Y. Ge, J. Gentry, K. Hornik, T. Hothorn, W. Huber, S. Iacus, R. Irizarry, F. Leisch, C. Li, M. Maechler, A. J. Rossini, G. Sawitzki, C. Smith, G. Smyth, L. Tierney, J. Y. H. Yang, and J. Zhang. Bioconductor: Open software development for computational biology and bioinformatics. *Genome Biology*, 5:R80, 2004. [1](#)
- V. Luu-The, N. Paquet, E. Calvo, and J. Cumps. Improved real-time rt-pcr method for high-throughput measurements using second derivative calculation and double correction. *Biotechniques*, 38(2):287–93, 2005. [42](#)
- P. Mestdagh, P. V. Vlierberghe, A. D. Weer, D. Muth, F. Westermann, F. Speleman, and J. Vandesompele. A novel and universal method for microrna rt-qpcr data normalization. *Genome Biol.*, 10(6):R64, 2009. [16](#)
- G. K. Smyth. Limma: linear models for microarray data. In R. Gentleman, V. Carey, S. Dudoit, and W. H. R. Irizarry, editors, *Bioinformatics and Computational Biology Solutions using R and Bioconductor*, pages 397–420. Springer, New York, 2005. [31](#)