

RReportGenerator : Automatic Reports from Routine Statistical Analysis using R

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<http://www-bio3d-igbmc.u-strasbg.fr/~wraff/>

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Automatic Data Analysis



: Powerful Means for Statistical Analysis

- ☺ vast collection of additional modules (CRAN, Bioconductor)
- ☹ command-line structure : “inaccessible” for Biologists,...

Our Motivation : Make R available in convenient way for

- automatic routine use
- via graphical user interface (GUI)
- allowing to access prewritten R-code
- creating reports in pdf-format

RReportGenerator : Principle

GUI managing tasks of :

- **selecting the data to be treated,**
- **calling command-line R,**
- **passing Sweave code**
(customize by substituting specific terms)
- **file-name & location of report**
- Customizing options : preferred search paths, ...

RReportGenerator

Scenario 1

Input Data 2

Output 3

3b

RReportGenerator : statistical analysis with automatic reports

Main Items

Analysis Scenario File : **Library** **Browse ...**

Infos

Data Input File : **Browse ...**

Output Folder : **Browse ...**

Report File Name : **.pdf** **.dvi (+eps)**

Optional Items

Supplemental Data Input File : **Browse ...**

Supplemental Data Output File : **Keep the .tex file**

Reset **Generate Report**

WWW Scenario Library

WWW Scenario Library :

- RRG_test1.Rnw
- RRG_test2.Rnw
- automAffyQC1.Rnw
- automAffyQC2.Rnw
- combineMAIAdata1.Rnw
- segmentCGH3.Rnw
- segmentCGH4.Rnw
- summarizeTCA_fromExcel.Rnw
- summarizeTCA_fromTxt.Rnw

Dismiss **Infos** **OK**

Execution finished:
the following files have been created :
D://projects/TCA_140307.pdf
D://projects/TCA_140307.txt

www accessible collection of updated scenarios

RReportGenerator

Scenario 1

Input Data 2

Output 3

www accessible collection of updated scenarios

4 Generate Report

data for export



Sweave & L^AT_EX

final .pdf report

temporary report (.tex)



Sample Code : Sweave

```
%@RRG_INFO
% Read a tab-delimited file & draw a scatter plot.
%@RRG_INFO_END

\documentclass[a4paper]{article}
\title{Minimal test scenario for RReportGenerator}
\usepackage{a4wide,Sweave}
\begin{document}
\maketitle

The file selected as 'Data Input File' is read :
<<chunk_read, echo=FALSE, print=TRUE>>=
  mydata <- read.table("<DATA_IN_FILE>")
  t.test(mydata[,1],mydata[,2],paired=TRUE)
@

<<chunk_plot1, echo=FALSE, fig=TRUE>>=
  plot(mydata[,2]~mydata[,1])
@

% If a name for 'Supplemental Data Output File' was given ...
<<chunk_save, echo=F, print=F, results=hide>>=
  write.table(summary(mydata), "<DATA_OUT_FILE>", row.names=F)
@
\end{document}
```

RReportGenerator Applications

- **Automatic analyses** for *routine applications* in
 - **Transcriptomics** : Quality control, data normalization
 - **Comparative Genomics Hybridization (CGH)**
Counseling to amplified/lost regions
 - **Transfected cell array (TCA)** :
Graphical and statistical summary

Examples : Transcription Profiling (Affymetrix CQ)

Affymetrix Batch Quality Control using RReportGenerator and R

August 1, 2008

This document was generated by Analysis Type File 'automAffyQC1.Rnw', Version: 1.2.1, a protocol for automated QC analysis for Affymetrix expression array data (using R version 2.7.0 on a x86-64-unknown-linux-gnu system).

The aim of this report is to provide information on multiple QC aspects for a set of Affymetrix arrays. The interpretation of QC parameters and QC plots should be done with care since this may lead to delicate decisions. For further information and details about the various plots shown in this report please look at the references section. This analysis protocol was written by wolfgang.raffelsberger (at) igbmc.fr, LGBI, IGBMC (Strasbourg, France).

QC analysis of 16 cel-files of type 'HG-U133A' with 22283 probesets each, found in path :

```
[1] "/genomics/g6/CELS/cancer/calvano2005/A_t0_02"
```

page 3

Both plots give a resume of the original PM values, either as boxplot or as (kernel) density estimate for the signal distribution (similar to Bioconductor-packages simpleaffy and affyQCReport).

page 4

This figure shows the QC plot from the simpleaffy package. For this purpose the 3' to 5' ratio for spiked-in and control genes (triangles for b-Actin and squares for GAPDH) are considered. The filled (blue) dot with the vertical line (heading to 0) represents the scaling factor. Finally, the percentage of 'present calls' and value of average background are shown in red on the left side. Note that the order of arrays is inverted (compared to other plots in this report).

page 5

This pair of graphs shows the RLE (top) and NUSE (bottom) plots from the affyPLM package. The RLE compares for each probeset (across all arrays) its expression value against its median across all arrays. The NUSE plot shows the standard errors for each probe-set (variability intra), standardized across all arrays.

page 6

This page shows false color images for the RLE residuals of all probes (from the affyPLM package, with red intensities corresponding to positive residuals and blue to negative residuals). Note, that smaller deviating zones have little impact on the overall results. For further information, examples and details see e.g. chapter by Bolstad in book by Gentleman *et al.*

page 7

MA plots (from affyPLM package) of each array against a synthetic median array (constructed from probe-wise medians of all arrays in the current project). The red line represents a lowest fit to the scatter plot and could be helpful in indicating non-linear relationships.

page 8

In the top part the RNA degradation plot shows the average intensity with respect to the sorted 5' to 3' position of the probes in the target-sequence. Depending on the type of microarray specific patterns can be observed (see also references Bolstad, Gentleman *et al.*). The lower figure shows a density estimate for the signal distribution of data resulting from RMA.

page 9

Similarity between RMA summarized samples measured as Euclidean distance : Heatmap of distance values.

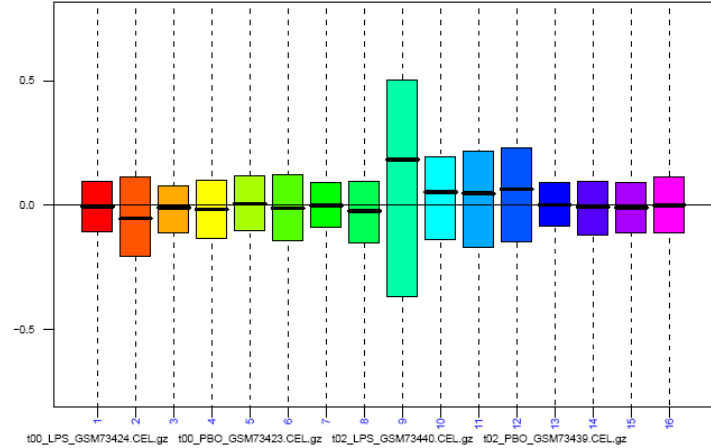
page 10

Principal components of RMA summarized samples. The main graphic shows the values of the first and second principal component for each of the samples. The smaller plots show the 2nd vs. 3rd as well as the 1st vs. 3rd component, the bargraph at the bottom indicates the amount of total variance captured by the individual principal components.

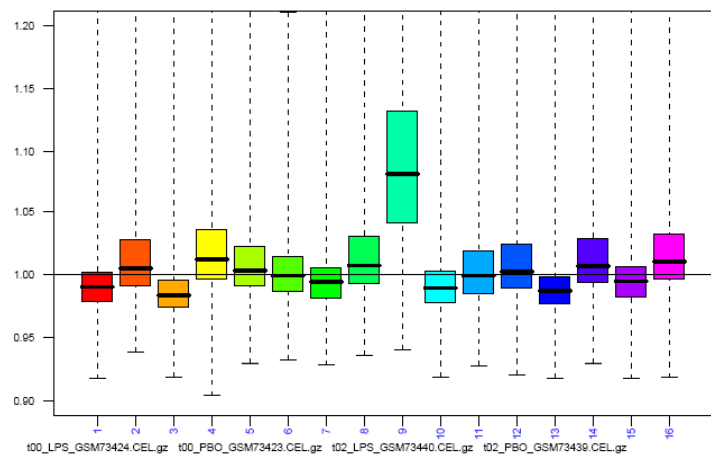
page 11 : References

Relative Log Expression (RLE) values

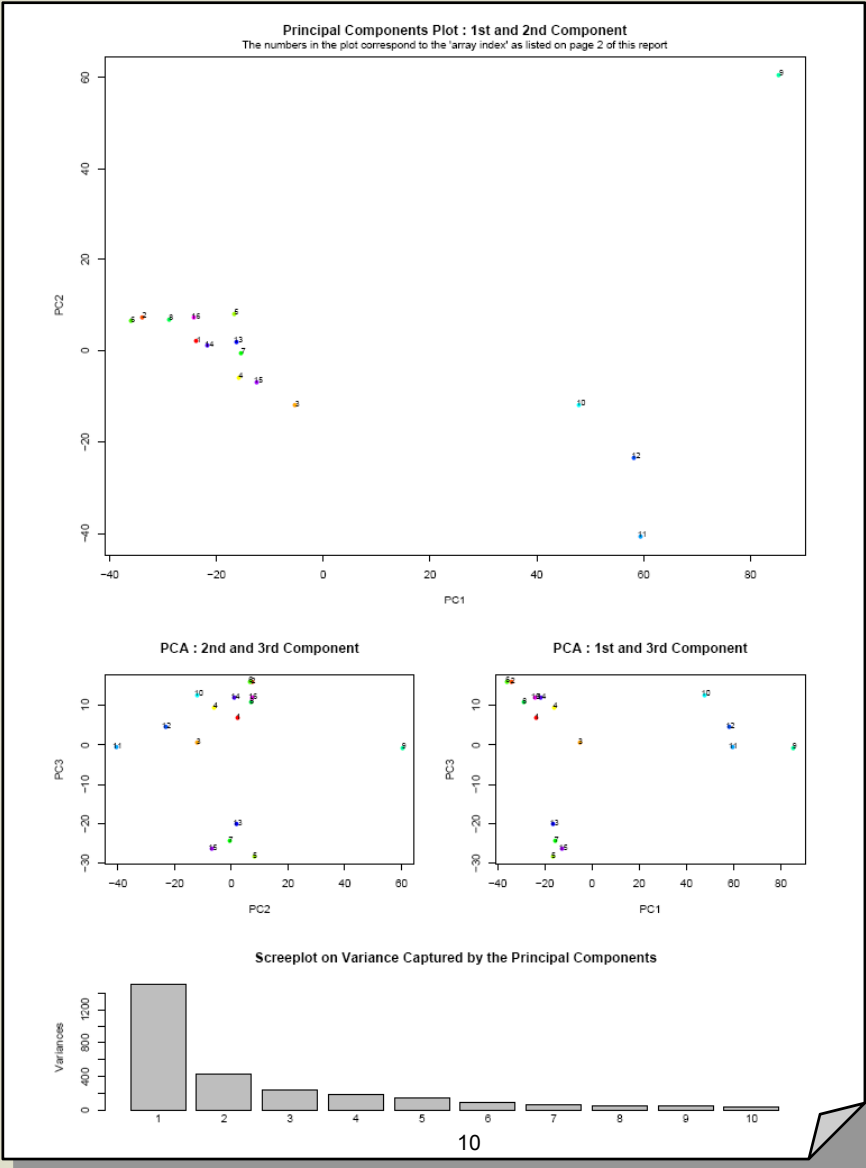
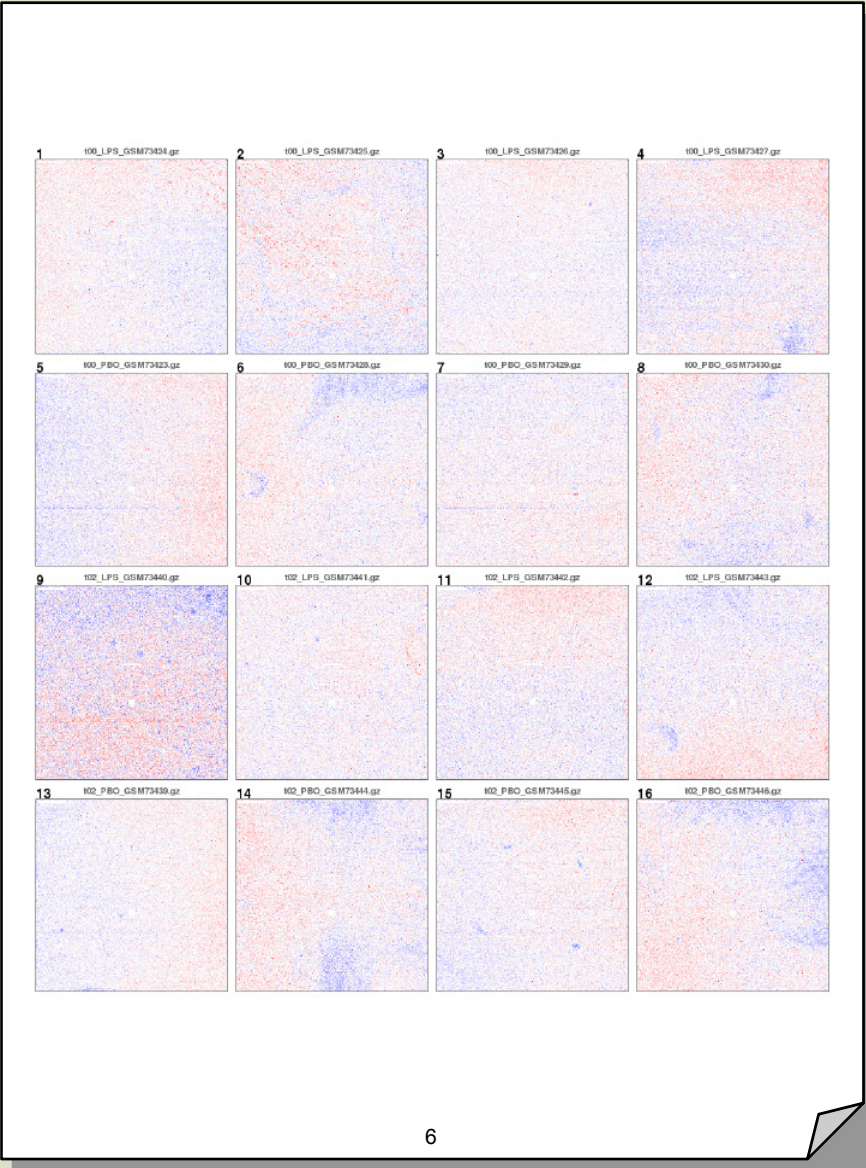
data from: /genomics/g6/CELS/cancer/calvano2005/A_t0_02



Normalized Unscaled Standard Errors (NUSE) values



Examples : Transcription Profiling (Affymetrix QC)



RReportGenerator Availability

Windows & Linux versions

coming with installer, user-manual and basic set of analysis scenarios under free GNU public license at :

<http://www-bio3d-igbmc.u-strasbg.fr/~wraff>

RReportGenerator Version 1.3.3

- **Paper Format** : Automatic switch between US and A4 paper format
- Windows : **Customize memory allocation of R**
- **Updated Analysis Scenarios**

More information :

RReportGenerator : Automatic reports from routine statistical analysis using R.
Raffelsberger W, Krause Y, Mouliner L, Kieffer D, Morand AL, Brino L, Poch O
Bioinformatics 2008, 24(2):276-278

Or : http://alnitak.u-strasbg.fr/wikili/index.php/RReportGenerator_English

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all users of RReportGenerator for feed-back!



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