

# Bioconductor mAPKL Package

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## Contents

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<b>1</b>	<b>Introduction</b>	<b>1</b>
<b>2</b>	<b>Identification of differentially expressed genes</b>	<b>2</b>
2.1	Breast cancer data . . . . .	2
2.2	Data normalization and transformation . . . . .	3
2.3	mAPKL gene selection . . . . .	5
2.4	Building and evaluating classification models . . . . .	5
<b>3</b>	<b>Advanced usage of the package</b>	<b>7</b>
3.1	Annotation analysis . . . . .	7
3.2	Network characteristics . . . . .	8
<b>4</b>	<b>Reporting</b>	<b>10</b>
<b>5</b>	<b>Session info</b>	<b>11</b>
<b>6</b>	<b>Reference</b>	<b>11</b>

## 1 Introduction

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The mAPKL bioconductor R package implements a hybrid gene selection method, which is based on the hypothesis that among the statistically significant genes in a ranked list, there should be clusters of

genes that share similar biological functions related to the investigated disease. Thus, instead of keeping a number of  $N$  top ranked genes, it would be more appropriate to define and keep a number of gene cluster exemplars.

The proposed methodology combines filtering and cluster analysis to select a small yet highly discriminatory set of non-redundant genes. Regarding the filtering step, a multiple hypothesis testing approach from *multtest* r-package (maxT) is employed to rank the genes of the training set according to their differential expression. Then the top  $N$  genes (e.g.  $N = 200$ ) are reserved for cluster analysis. First the index of Krzanowski and Lai as included in the *ClusterSim* r-package is applied on the disease samples of the training set to determine the number of clusters. The Krzanowski and Lai index is defined by  $DIFF(k) = (k-1)^{\frac{2}{p}}W_{k-1} - k^{\frac{2}{p}}W_k$  when choosing the number of clusters ( $k$ ) to maximize the quantity  $KL(k) = \left| \frac{DIFF(k)}{DIFF(k+1)} \right|$ . The  $W_k$  denotes the within-cluster sum of squared errors.

Finally, cluster analysis is performed with the aid of Affinity Propagation (AP) clustering algorithm, which detects  $n$  ( $n = k$  the Krzanowski and Lai index) clusters among the top  $N$  genes, the so called exemplars. Those  $n$  exemplars are expected to form a classifier that shall discriminate between normal and disease samples (Sakellariou et al. 2012, *BMC Bioinformatics* **13**:270). This package implements the pre-mentioned methodology through a core function, the *mAPKL*. In the upcoming sections follows a guidance of the included functions and its functionality through differential expression analysis scenarios on a breast cancer dataset (GSE5764) which is part of the *mAPKLData* package.

## 2 Identification of differentially expressed genes

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### 2.1 Breast cancer data

Throughout this tutorial we utilized a publicly available breast cancer dataset comprised of 30 samples, where 20 of them represent normal cases and the remaining 10 samples stand for tumor cases. We first load the package and then the breast cancer data. Then with the aid of the *sampling* function we create a separate training and validation sets where 60% of the samples will be used for training and the rest 40% of the samples will be used for evaluation purposes.

```
library(mAPKL)
library(mAPKLData)
data(mAPKLData)
varLabels(mAPKLData)
breast <- sampling(Data=mAPKLData, valPercent=40, classLabels="type", seed=135)
```

The *sampling* function returns an S3 class (breast) with two eSet class objects that nest the relevant training and validation samples. Those two objects are used throughout the rest of the analysis.

```
breast
## $trainData
```

```
## ExpressionSet (storageMode: lockedEnvironment)
## assayData: 54675 features, 18 samples
##   element names: exprs
## protocolData: none
## phenoData
##   sampleNames: GSM134584 GSM134690 ... GSM134695 (18 total)
##   varLabels: title type
##   varMetadata: labelDescription
## featureData: none
## experimentData: use 'experimentData(object)'
##   pubMedIds: 17389037
## Annotation: hgu133plus2
##
## $testData
## ExpressionSet (storageMode: lockedEnvironment)
## assayData: 54675 features, 12 samples
##   element names: exprs
## protocolData: none
## phenoData
##   sampleNames: GSM134691 GSM134588 ... GSM134692 (12 total)
##   varLabels: title type
##   varMetadata: labelDescription
## featureData: none
## experimentData: use 'experimentData(object)'
##   pubMedIds: 17389037
## Annotation: hgu133plus2
```

## 2.2 Data normalization and transformation

We perform normalization to the expression values through the *preprocess* function.

```
normTrainData <- preprocess(breast$trainData)
normTestData <- preprocess(breast$testData)
```

The *preprocess* function produces a list of several available normalization and transformation options. Besides density plots per method are produced and saved to current working directory to assist the user to decide upon which method to select before proceeding to mAPKL analysis.

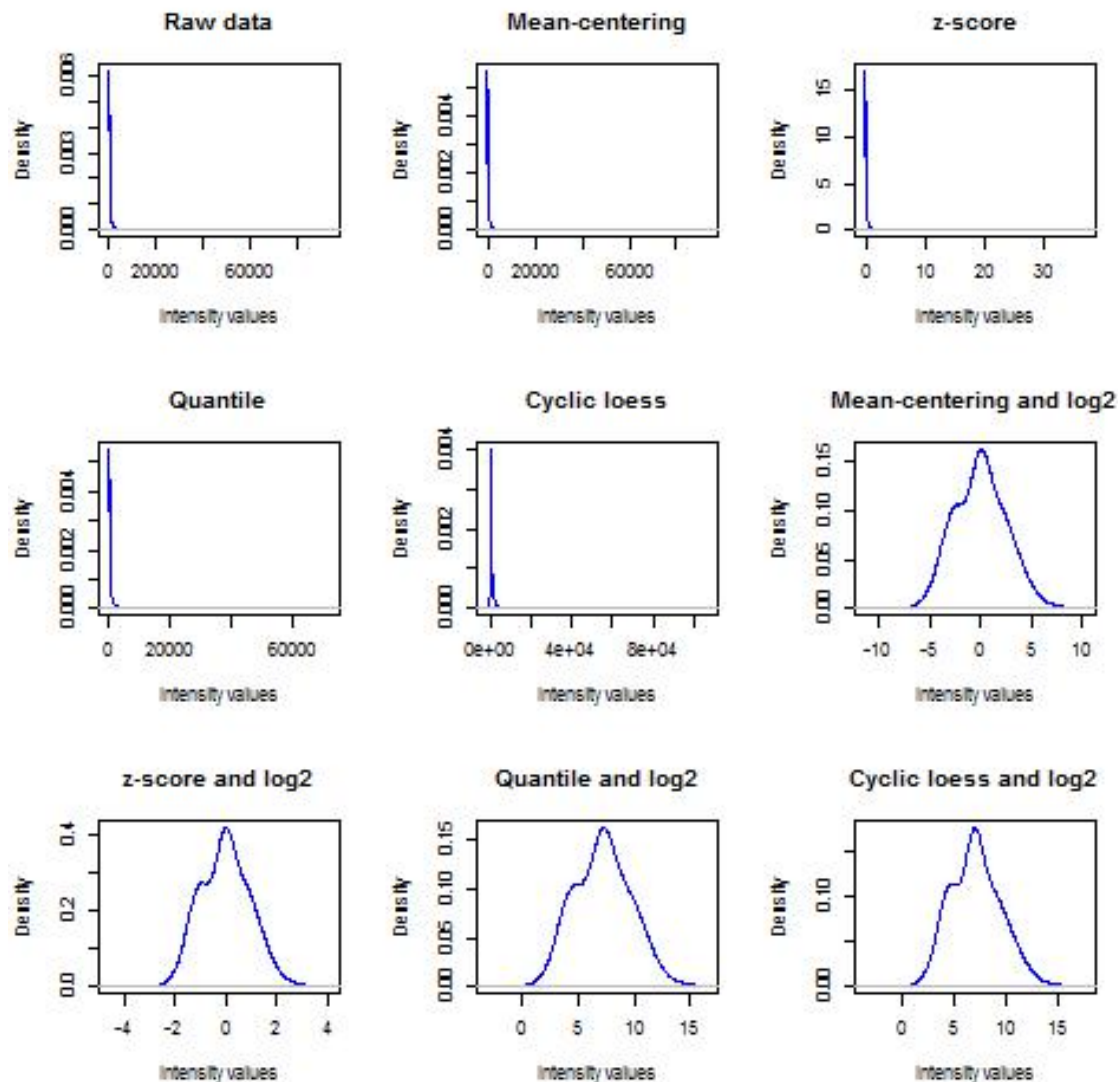


Figure 1: Density plots of normalized intensity values

```
attributes(normTrainData)
## $names
## [1] "rawdata"      "mc.normdata"  "z.normdata"   "q.normdata"   "cl.normdata"
## [6] "mcL2.normdata" "zL2.normdata" "qL2.normdata" "clL2.normdata"
```

The following graph presents the density plots of 8 possible normalization process with or without log2 transformation. The *preprocess* function applies all of them and it is up to the user, which one will engage for the rest of the analysis. In brief, the available approaches are mean-centering, z-score, quantile, and cyclic loess. During this case study we will proceed with the expression values following log2 transformation and cyclic loess normalization.

## 2.3 mAPKL gene selection

In this example we employ the expression values of log2 transformation and cyclic loess normalization to proceed with the *mAPKL* analysis.

```

exprs(breast$strainData) <- normTrainData$cLL2.normdata
exprs(breast$testData) <- normTestData$cLL2.normdata
out.cLL2 <- mAPKL(trObj = breast$strainData, classLabels = "type",
  valObj = breast$testData, dataType = 7)

## b=10 b=20 b=30 b=40 b=50 b=60 b=70 b=80 b=90 b=100
## b=110 b=120 b=130 b=140 b=150 b=160 b=170 b=180 b=190 b=200
## b=210 b=220 b=230 b=240 b=250 b=260 b=270 b=280 b=290 b=300
## b=310 b=320 b=330 b=340 b=350 b=360 b=370 b=380 b=390 b=400
## b=410 b=420 b=430 b=440 b=450 b=460 b=470 b=480 b=490 b=500
## b=510 b=520 b=530 b=540 b=550 b=560 b=570 b=580 b=590 b=600
## b=610 b=620 b=630 b=640 b=650 b=660 b=670 b=680 b=690 b=700
## b=710 b=720 b=730 b=740 b=750 b=760 b=770 b=780 b=790 b=800
## b=810 b=820 b=830 b=840 b=850 b=860 b=870 b=880 b=890 b=900
## b=910 b=920 b=930 b=940 b=950 b=960 b=970 b=980 b=990 b=1000

## Please wait! The (KL) cluster indexing may take several minutes...
## Asking for 15 number of clusters
## fc according to limma

```

## 2.4 Building and evaluating classification models

After having get the exemplars from *mAPKL* analysis we build an SVM classifier to test their discriminatory performance. Regarding the SVM setup, we utilize a linear kernel for which the cost attribute is inferred by the `tune.svm` function. however, the user may freely use another kernel and a different Cross Validation approach than 5-folds.

```

clasPred <- classification(out.cLL2@exemplTrain, "type", out.cLL2@exemplTest)

## The training set has 12 Negative and 6 Positive samples. Using k-fold=5 C-V
## ##### THE BEST PARAMETERS TUNING STAGE #####
## ##### THE TRAINING STAGE #####

##
## Call:
## svm.default(x = train.mtx, y = lbls, scale = FALSE, type = "C-classification",
##   kernel = "linear", gamma = best_gamma, cost = best_cost, cross = k_fold)
##

```

```

##
## Parameters:
##   SVM-Type: C-classification
##   SVM-Kernel: linear
##     cost: 2
##     gamma: 0.125
##
## Number of Support Vectors: 5
## ##### THE PREDICTION STAGE #####
##
##           Test Labels Prediction Labels
## GSM134691           0           0
## GSM134588           0           0
## GSM134688           0           0
## GSM134694           0           1
## GSM134697           0           0
## GSM134700           0           0
## GSM134687           0           0
## GSM134709           0           0
## GSM134710           1           1
## GSM134698           1           1
## GSM134689           1           1
## GSM134692           1           1
##
## Negative samples: 8
## Positive samples: 4
## TN=7
## FP=1
## TP=4
## FN=0
## AUC=0.94
## Accuracy=92.00
## MCC=0.84
## Specificity=0.88
## Sensitivity=1.00

```

The output of the *classification* inform us about the SVM set up, the number of Support Vectors and finally show the the predicted labels along with the initial. In this example there is a validation set different from the training set and therefore we may use the respective labels to obtain the performance characteristics. The relevant function *metrics* called inside the *classification* function, calculates five key measures: the Area Under the ROC curve AUC, the classification accuracy, the Matthews correlation

coefficient MCC classification measure, the degree of true negative's identification Specificity, and finally the degree of true positive's identification Sensitivity.

## 3 Advanced usage of the package

---

### 3.1 Annotation analysis

For each contemporary chip technology, there is a relevant annotation file, in which the the user may drag several *genome oriented* information. Regarding the breast cancer microarray data, the gene expression values were stored on Affumetrix gene chips. Using the *annotate* function, the user may obtain several info related to probe id, gene symbol, Entrez id, ensembl id, and chromosomal location.

```
gene.info <- annotate(out.cll2@exemplars, "hgu133plus2.db")
gene.info@results
```

##	PROBEID	SYMBOL	ENTREZID	ENSEMBL	MAP
## 1	215717_s_at	FBN2	2201	ENSG00000138829	5q23-q31
## 2	1561358_at	TXLNA	200081	ENSG00000084652	1p35.1
## 3	222752_s_at	TMEM206	55248	ENSG00000065600	1q32.3
## 4	233922_at	<NA>	<NA>	<NA>	<NA>
## 5	218871_x_at	CSGALNACT2	55454	ENSG00000169826	10q11.21
## 6	33323_r_at	SFN	2810	ENSG00000175793	1p36.11
## 7	244311_at	<NA>	<NA>	<NA>	<NA>
## 8	220932_at	<NA>	<NA>	<NA>	<NA>
## 9	205508_at	SCN1B	6324	ENSG00000105711	19q13.1
## 10	209596_at	MXRA5	25878	ENSG00000101825	Xp22.33
## 11	215180_at	<NA>	<NA>	<NA>	<NA>
## 12	1560638_a_at	<NA>	<NA>	<NA>	<NA>
## 13	201852_x_at	COL3A1	1281	ENSG00000168542	2q31
## 14	229947_at	PI15	51050	ENSG00000137558	8q21.11
## 15	221731_x_at	VCAN	1462	ENSG00000038427	5q14.3

We may exploit the output of the *annotate* function to extent our analysis. For instance, we may perform *pathway analysis* on the exemplars. For this purpose we will utilize the *reactome.db* package.

```
library("reactome.db")
# We first remove the 'NA' entries
genes <- gene.info@entrezId[!is.na(gene.info@entrezId)]
# Then we map the Entrez ID to Reactome pathway identifiers
qExtID2PathID <- mapIds(reactome.db, genes, "PATHID", "ENTREZID",
```

```

multiVals = "list")
notNA.idx <- unlist(lapply(qExtID2PathID, function(i) !all(is.na(i))))
qExtID2PathID <- qExtID2PathID[notNA.idx]
pathID <- as.character(qExtID2PathID[[1]])
# Finally we map Reactome pathway identifiers to pathway
# names
pathName <- mapIds(reactome.db, pathID, "PATHNAME", "PATHID")
pathName

##                                1474228
## "Homo sapiens: Degradation of the extracellular matrix"
##                                1474244
##      "Homo sapiens: Extracellular matrix organization"
##                                1566948
##      "Homo sapiens: Elastic fibre formation"
##                                2129379
## "Homo sapiens: Molecules associated with elastic fibres"

```

### 3.2 Network characteristics

Regarding the network characteristics, we compute through the *netwAttr* function three different types of centralities (degree, closeness, betweenness) and a measure for clustering coefficient called transitivity. The degree centrality of a node refer to the number of connections or edges of that node to other nodes. The closeness centrality describes the reciprocal accumulated shortest length distance from a node to all other connected nodes. The betweenness centrality depicts the number of times a node intervenes along the shortest path of two other nodes. Transitivity measures the degree of nodes to create clusters within a network. For all four network measures we provide both global and local values. Furthermore, we compose an edge list (Node1-Node2-weight) based on the  $N$  top ranked genes. We may exploit that measures to depict the exemplars' network characteristics

```

net.attr <- netwAttr(out.c1L2)
wDegreeL <- net.attr@degree$WdegreeL[out.c1L2@exemplars]
wClosenessL <- net.attr@closeness$WclosenessL[out.c1L2@exemplars]
wBetweennessL <- net.attr@betweenness$WbetweennessL[out.c1L2@exemplars]
wTransitivityL <- net.attr@transitivity$WtransitivityL[out.c1L2@exemplars]

Global.val <- c(net.attr@degree$WdegreeG, net.attr@closeness$WclosenessG,
               net.attr@betweenness$WbetweennessG, net.attr@transitivity$WtransitivityG)

```



```
Global.val <- round(Global.val, 2)
exempl.netattr <- rbind(wDegreeL, wClosenessL, wBetweenessL, wTransitivityL)
```

```
netAttr <- cbind(Global.val, exempl.netattr)
```

```
netAttr <- t(netAttr)
```

```
netAttr
```

##	wDegreeL	wClosenessL	wBetweenessL	wTransitivityL
## Global.val	330.18	0.93	741.81	0.57
## 215717_s_at	308.35	1.25	886.00	0.14
## 1561358_at	346.92	1.34	1141.00	0.14
## 222752_s_at	327.89	0.65	0.00	0.14
## 233922_at	317.58	0.79	2.00	0.15
## 218871_x_at	293.73	0.53	768.00	0.14
## 33323_r_at	338.19	0.27	0.00	0.13
## 244311_at	294.80	0.63	0.00	0.15
## 220932_at	359.10	0.66	0.00	0.14
## 205508_at	309.07	0.89	4.00	0.14
## 209596_at	345.13	1.34	278.00	0.14
## 215180_at	333.37	1.37	1440.00	0.14
## 1560638_a_at	368.23	1.38	4615.00	0.14
## 201852_x_at	353.34	0.93	24.67	0.15
## 229947_at	317.11	1.19	496.00	0.15
## 221731_x_at	331.01	0.61	14.00	0.15

and identify potential hubs.

```
# For local degree > global + standard deviation
```

```
sdev <- sd(net.attr@degree$WdegreeL)
```

```
msd <- net.attr@degree$WdegreeG + sdev
```

```
hubs <- wDegreeL[which(wDegreeL > msd)]
```

```
hubs
```

```
## 220932_at 1560638_a_at
```

```
## 359.10 368.23
```

Finally, we may plot the network for those nodes that their local weighted degree is greater than Global weighted degree plus 2 times the standard deviation. We set this rule for both significance and illustration purposes (that edge list has dimension 604 x 3).

```
sdev <- sd(net.attr@degree$WdegreeL)
```

```
ms2d <- net.attr@degree$WdegreeG + 2 * sdev
```



## 5 Session info

---

```

sessionInfo()

## R version 3.2.0 (2015-04-16)
## Platform: x86_64-unknown-linux-gnu (64-bit)
## Running under: Ubuntu 14.04.2 LTS
##
## locale:
##  [1] LC_CTYPE=en_US.UTF-8      LC_NUMERIC=C              LC_TIME=en_US.UTF-8
##  [4] LC_COLLATE=C              LC_MONETARY=en_US.UTF-8  LC_MESSAGES=en_US.UTF-8
##  [7] LC_PAPER=en_US.UTF-8     LC_NAME=C                 LC_ADDRESS=C
## [10] LC_TELEPHONE=C           LC_MEASUREMENT=en_US.UTF-8 LC_IDENTIFICATION=C
##
## attached base packages:
## [1] stats4      parallel  stats      graphics  grDevices  utils      datasets  methods
## [9] base
##
## other attached packages:
##  [1] igraph_0.7.1          reactome.db_1.52.1      hgu133plus2.db_3.1.2  org.Hs.eg.db_3.1.2
##  [5] AnnotationDbi_1.30.0  GenomeInfoDb_1.4.0     IRanges_2.2.0         S4Vectors_0.6.0
##  [9] mAPKLData_0.99.1      mAPKL_1.0.0            RSQlite_1.0.0         DBI_0.3.1
## [13] Biobase_2.28.0        BiocGenerics_0.14.0    knitr_1.9
##
## loaded via a namespace (and not attached):
##  [1] Rcpp_0.11.5          cluster_2.0.1          splines_3.2.0         MASS_7.3-40
##  [5] lattice_0.20-31     stringr_0.6.2         highr_0.4.1           tools_3.2.0
##  [9] grid_3.2.0          parmigene_1.0.2       e1071_1.6-4          class_7.3-12
## [13] multtest_2.24.0     survival_2.38-1       ade4_1.7-2           rgl_0.95.1201
## [17] Matrix_1.2-0        apcluster_1.4.1       formatR_1.1          clusterSim_0.44-2
## [21] evaluate_0.6        limma_3.24.0          R2HTML_2.3.1         BiocStyle_1.6.0

```

## 6 Reference

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Sakellariou, A., D. Sanoudou, and G. Spyrou. "Combining Multiple Hypothesis Testing and Affinity Propagation Clustering Leads to Accurate, Robust and Sample Size Independent Classification on Gene Expression Data." *BMC Bioinformatics* 13 (2012): 270.

<http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE5764>

### mAP-KL Analysis Report

Sat Dec 06 20:54:13 2014

#### Data Samples

Training Set	Class Labels	Validation Set	Class Labels
GSM134584	0	GSM134691	0
GSM134690	0	GSM134588	0
GSM134693	0	GSM134688	0
GSM134696	0	GSM134694	0
GSM134586	0	GSM134697	0
GSM134589	0	GSM134700	0
GSM134699	0	GSM134687	0
GSM134702	0	GSM134709	0
GSM134705	0	GSM134710	1
GSM134708	0	GSM134698	1
GSM134703	0	GSM134689	1
GSM134706	0	GSM134692	1
GSM134701	1		
GSM134704	1		
GSM134587	1		
GSM134591	1		
GSM134707	1		
GSM134695	1		

#### Exemplars

Exemplars	Adj.p-value	P-value	FC	wl.degree	wl.closeness	wl.betweenness	wl.transitivity
215717_s_at	0.885	0.001	2.252	308.35	1.25	886	0.14
1561358_at	0.928	0.001	1.246	346.92	1.34	1141	0.14
222752_s_at	0.947	0.001	0.88	327.89	0.65	0	0.14
233922_at	0.952	0.001	-1.102	317.58	0.79	2	0.15
218871_x_at	0.987	0.001	0.901	293.73	0.53	768	0.14
33323_r_at	0.997	0.001	-0.701	338.19	0.27	0	0.13
244311_at	0.999	0.003	-1.355	294.8	0.63	0	0.15
220932_at	0.999	0.002	-1.069	359.1	0.66	0	0.14
205508_at	1	0.001	1.349	309.07	0.89	4	0.14
209596_at	1	0.002	1.662	345.13	1.34	278	0.14
215180_at	1	0.003	-1.888	333.37	1.37	1440	0.14
1560638_a_at	1	0.001	-2.336	368.23	1.38	4615	0.14
201852_x_at	1	0.002	2.239	353.34	0.93	24.67	0.15
229947_at	1	0.003	-4.004	317.11	1.19	496	0.15
221731_x_at	1	0.001	2.046	331.01	0.61	14	0.15

#### Classification Performance

(Hold-out Validation)

AUC	Accuracy	MCC	Specificity	Sensitivity
0.94	92.00	0.84	0.88	1.00

#### Genome Annotation

PROBE ID	SYMBOL	ENTREZ ID	ENSEMBL ID	MAP
215717_s_at	FBN2	2201	ENSG00000138829	5q23-q31
1561358_at	TXLNA	200081	ENSG00000084652	1p35.1
222752_s_at	THM206	55248	ENSG00000065600	1q32.3
233922_at	NA	NA	NA	NA
218871_x_at	CSGALNACT2	55454	ENSG00000169826	10q11.21
33323_r_at	SFN	2810	ENSG00000175793	1p36.11
244311_at	NA	NA	NA	NA
220932_at	NA	NA	NA	NA
205508_at	SCN1B	6324	ENSG00000105711	19q13.1
209596_at	MXRA5	25878	ENSG00000101825	xp22.33
215180_at	NA	NA	NA	NA
1560638_a_at	NA	NA	NA	NA
201852_x_at	COL3A1	1281	ENSG00000168542	2q31
229947_at	PI15	51050	ENSG00000137558	8q21.11
221731_x_at	VCAN	1462	ENSG00000038427	5q14.3

Figure 3: mAPKL analysis report