

PCOT2: Principal Coordinates and Hotelling's T^2 for the analysis of microarray data

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1 Overview

`pcot2` is an R-package for the analysis of groups of genes in microarray experiments. It utilizes inter-gene correlation information to detect significant alterations in the activities of gene sets. Incorporating additional (usually functional) information into the data analysis process allows gene interactions to be investigated in a statistical framework. One of the reasons that gene set analysis is becoming important is that it is suitable for detecting small coordinated changes in expression of groups of genes which are functionally related, which may not be considered significant in a single gene analysis. This vignette gives a tutorial-style introduction to the functions in the `pcot2` package. These functions are used for testing and visualizing changes in expression activity for groups of genes.

2 Example: ALL/AML data

In this example the ALL/AML leukemia data set of Golub *et al.*(1999) is used to illustrate the functionality of the `pcot2` package. This data set contains 38 bone marrow samples obtained from adult leukemia patients, 11 relating to acute myeloid leukemia (AML, class 1) and 27 relating to acute lymphoblastic leukemia (ALL, class 0). Gene expression levels were measured using Affymetrix high density oligonucleotide arrays containing 6817 human genes, of which 3051 genes were considered suitable for analysis by Golub et al.(1999) after pre-processing. This data set is available as part of the `multtest` package and gene sets are defined as KEGG pathways using the `hu6800` annotation package. Both packages can be downloaded from www.bioconductor.org.

```
> library(pcot2)
> library(multtest)
> library(hu6800)
> set.seed(1234567)
```

3 The `pcot2` function

The `pcot2` function implements the PCOT2 testing method, which is a two-stage permutation-based approach for testing changes in activity in pre-specified

gene sets. The function requires at least three inputs: gene expression data, sample class labels, and a gene category indicator matrix. The gene expression data should be in the form of a matrix with no missing values. Data pre-processing (e.g. normalization) must therefore take place before running the PCOT2 analysis.

```
> data(golub)
> rownames(golub) <- golub.gnames[, 3]
> colnames(golub) <- golub.cl
```

The class labels represent two distinct experimental conditions (e.g., AML and ALL).

```
> golub.cl
[1] 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
```

The gene category indicator matrix is designed to indicate presence or absence of genes in the pre-defined gene categories (e.g., gene pathways). The indicator matrix contains rows representing gene identifiers for genes present in the expression data, and columns representing pre-defined group names. The values 1 or 0 indicate the presence or absence of a gene in a particular group.

In this example, the `hu6800` annotation package is used to define the KEGG (<http://www.genome.jp/kegg/pathway.html>) pathways for all of 3051 genes in the data. The `getImat` function is used to generate an indicator matrix which includes 65 KEGG pathways containing at least 10 of the total 3051 genes.

```
> KEGG.list <- as.list(hu6800PATH)
> imat <- getImat(golub, KEGG.list, ms = 10)
> colnames(imat) <- paste("KEGG", colnames(imat), sep = "")
> dim(imat)
[1] 3051 109
```

Permutations are used to produce *p*-values based on the null distribution of the T^2 statistic. By default `pcot2` will automatically run 1000 permutations. In order to minimize the time taken to build this vignette, only 10 permutations have been performed.

```
> results <- pcot2(golub, golub.cl, imat, iter = 10)
```

Comparison: 0-1

The output from the `pcot2` function can contain information on either all pathways or just significantly differentially expressed pathways, based on the value of α used in the function, where α determines the significance threshold for the permutation *p*-values. For each KEGG pathway, the number of genes in the pathway is listed, along with Hotelling's T^2 statistic. These are followed by parametric *p*-values for the test statistic, both raw and adjusted. The last two columns provide raw and adjusted permutation-based *p*-values. The default adjustment method is the false discovery rate controlling method of Benjamini and Yekutieli (2001).

```

> results$res.sig

[1] Num          T2          P.nor        P.adj        P.permu      P.permu.adj
<0 rows> (or 0-length row.names)

> results$res.all

    Num          T2          P.nor        P.adj        P.permu      P.permu.adj
KEGG03050  10 48.62835408 3.189816e-07 9.555251e-06  0.1  0.5805784
KEGG00980  13 69.18821215 7.093654e-09 5.824626e-07  0.1  0.5805784
KEGG04010  97 37.55521790 3.711571e-06 4.961185e-05  0.1  0.5805784
KEGG04210  44 27.29935197 5.138148e-05 5.469013e-04  0.1  0.5805784
KEGG04620  52 45.79163870 5.792791e-07 1.204330e-05  0.1  0.5805784
KEGG04660  43 33.33813073 1.042993e-05 1.275498e-04  0.1  0.5805784
KEGG04662  39 46.75747370 4.717016e-07 1.203039e-05  0.1  0.5805784
KEGG04920  30 57.38563184 5.693675e-08 3.636188e-06  0.1  0.5805784
KEGG05120  38 71.94302882 4.512199e-09 5.186977e-07  0.1  0.5805784
KEGG05212  48 30.39925365 2.225552e-05 2.664972e-04  0.1  0.5805784
KEGG05215  49 53.56622252 1.182413e-07 5.614801e-06  0.1  0.5805784
KEGG05220  52 40.10938567 2.042326e-06 3.260758e-05  0.1  0.5805784
KEGG05221  38 42.96538031 1.071892e-06 1.987400e-05  0.1  0.5805784
KEGG05222  54 44.61526063 7.464344e-07 1.479414e-05  0.1  0.5805784
KEGG04510  87 49.07535509 2.908893e-07 9.288624e-06  0.1  0.5805784
KEGG04512  32 50.31731778 2.257251e-07 8.108789e-06  0.1  0.5805784
KEGG04640  70 115.40054196 1.210776e-11 6.959209e-09  0.1  0.5805784
KEGG04810  90 48.42804428 3.324880e-07 9.555251e-06  0.1  0.5805784
KEGG01032  10 16.63239211 1.298268e-03 9.949453e-03  0.1  0.5805784
KEGG04060  86 50.94081419 1.990119e-07 7.625772e-06  0.1  0.5805784
KEGG04612  51 45.73226060 5.866882e-07 1.204330e-05  0.1  0.5805784
KEGG04630  56 38.32636360 3.092383e-06 4.426758e-05  0.1  0.5805784
KEGG04650  68 45.97734942 5.567382e-07 1.204330e-05  0.1  0.5805784
KEGG04670  57 40.31362320 1.948756e-06 3.200262e-05  0.1  0.5805784
KEGG04720  38 14.22610932 2.944604e-03 2.054732e-02  0.1  0.5805784
KEGG01510  27 14.13381648 3.040922e-03 2.080761e-02  0.1  0.5805784
KEGG05040  21 13.80932761 3.406880e-03 2.303743e-02  0.1  0.5805784
KEGG04020  62 42.29976180 1.243043e-06 2.232709e-05  0.1  0.5805784
KEGG00230  52 17.72039780 9.075225e-04 7.346748e-03  0.1  0.5805784
KEGG00240  31 53.31591846 1.241763e-07 5.614801e-06  0.1  0.5805784
KEGG05010  16 5.72814930 7.547169e-02 4.337907e-01  0.1  0.5805784
KEGG04742  10 9.16510728 1.889037e-02 1.193150e-01  0.1  0.5805784
KEGG00500  18 18.28339353 7.561735e-04 6.208969e-03  0.1  0.5805784
KEGG04080  58 43.84884711 8.822515e-07 1.690313e-05  0.1  0.5805784
KEGG04530  40 33.48419039 1.005284e-05 1.256108e-04  0.1  0.5805784
KEGG04012  39 21.80887165 2.514144e-04 2.330744e-03  0.1  0.5805784
KEGG04730  36 37.99949685 3.340337e-06 4.571273e-05  0.1  0.5805784
KEGG04910  59 26.20096780 6.979813e-05 7.038255e-04  0.1  0.5805784
KEGG05210  43 27.62201433 4.700716e-05 5.097817e-04  0.1  0.5805784
KEGG05213  29 26.29243943 6.802606e-05 6.982057e-04  0.1  0.5805784
KEGG05214  41 20.52326018 3.726642e-04 3.295342e-03  0.1  0.5805784
KEGG05218  32 19.21207000 5.619507e-04 4.749911e-03  0.1  0.5805784
KEGG05219  25 54.45386664 9.949904e-08 5.614801e-06  0.1  0.5805784

```

KEGG05223	32	17.10738266	1.109387e-03	8.616828e-03	0.1	0.5805784
KEGG00190	42	14.20955560	2.961639e-03	2.054732e-02	0.1	0.5805784
KEGG00010	36	8.52085605	2.429027e-02	1.501224e-01	0.1	0.5805784
KEGG00030	15	13.50674640	3.790243e-03	2.533172e-02	0.1	0.5805784
KEGG00051	17	26.65539600	6.144979e-05	6.421756e-04	0.1	0.5805784
KEGG00710	12	6.02236856	6.673974e-02	3.874766e-01	0.1	0.5805784
KEGG04514	72	25.59171339	8.291894e-05	8.077889e-04	0.1	0.5805784
KEGG04350	28	20.14928563	4.185715e-04	3.645204e-03	0.1	0.5805784
KEGG04520	36	23.80529823	1.387873e-04	1.329519e-03	0.1	0.5805784
KEGG04310	42	37.04027266	4.197126e-06	5.360874e-05	0.1	0.5805784
KEGG05216	22	29.27179542	3.003285e-05	3.452412e-04	0.1	0.5805784
KEGG04610	15	73.36386723	3.589229e-09	5.186977e-07	0.1	0.5805784
KEGG00760	10	42.06352176	1.310543e-06	2.282619e-05	0.1	0.5805784
KEGG04540	41	10.89127317	9.799036e-03	6.290863e-02	0.1	0.5805784
KEGG04912	38	15.91910929	1.648417e-03	1.214699e-02	0.1	0.5805784
KEGG00590	19	47.57667675	3.970128e-07	1.086629e-05	0.1	0.5805784
KEGG04916	33	16.52924108	1.343613e-03	1.016147e-02	0.1	0.5805784
KEGG04370	36	30.08550891	2.417879e-05	2.836185e-04	0.1	0.5805784
KEGG04664	38	61.37981160	2.735820e-08	1.965593e-06	0.1	0.5805784
KEGG04340	11	6.07312845	6.534459e-02	3.832478e-01	0.1	0.5805784
KEGG00071	19	39.18341951	2.530215e-06	3.827101e-05	0.1	0.5805784
KEGG00280	21	40.94467897	1.687207e-06	2.852237e-05	0.1	0.5805784
KEGG00310	14	28.90293286	3.316409e-05	3.737611e-04	0.1	0.5805784
KEGG00380	21	88.10612309	3.925660e-10	1.128181e-07	0.1	0.5805784
KEGG00410	13	46.66122632	4.814060e-07	1.203039e-05	0.1	0.5805784
KEGG00640	16	49.10831719	2.889241e-07	9.288624e-06	0.1	0.5805784
KEGG00650	16	15.98099220	1.614410e-03	1.205089e-02	0.1	0.5805784
KEGG00562	14	19.02129910	5.970409e-04	4.973373e-03	0.1	0.5805784
KEGG04070	31	25.77606406	7.869364e-05	7.798441e-04	0.1	0.5805784
KEGG05030	15	28.15020247	4.067506e-05	4.495945e-04	0.1	0.5805784
KEGG00350	13	5.31695549	8.975286e-02	5.107672e-01	0.1	0.5805784
KEGG00561	16	69.24258213	7.029794e-09	5.824626e-07	0.1	0.5805784
KEGG00564	10	45.87159715	5.694580e-07	1.204330e-05	0.1	0.5805784
KEGG04940	34	7.77927983	3.259065e-02	1.992789e-01	0.1	0.5805784
KEGG03320	20	53.20149317	1.269935e-07	5.614801e-06	0.1	0.5805784
KEGG00620	16	21.66912268	2.622921e-04	2.392989e-03	0.1	0.5805784
KEGG00860	15	51.68661364	1.713804e-07	7.036053e-06	0.1	0.5805784
KEGG04360	33	39.55981347	2.318480e-06	3.601619e-05	0.1	0.5805784
KEGG04740	10	14.88788894	2.341753e-03	1.703767e-02	0.1	0.5805784
KEGG04930	18	17.46695844	9.858206e-04	7.761955e-03	0.1	0.5805784
KEGG04110	51	46.18411534	5.327283e-07	1.204330e-05	0.1	0.5805784
KEGG00020	12	12.20751287	6.036512e-03	3.942752e-02	0.1	0.5805784
KEGG00330	13	17.47113357	9.844744e-04	7.761955e-03	0.1	0.5805784
KEGG00220	12	38.23761530	3.157719e-06	4.426758e-05	0.1	0.5805784
KEGG00260	12	9.01420916	2.002974e-02	1.251364e-01	0.1	0.5805784
KEGG00360	11	38.952477906	2.670169e-06	3.935231e-05	0.1	0.5805784
KEGG00252	15	20.66838185	3.563113e-04	3.199968e-03	0.1	0.5805784
KEGG00970	16	23.40339175	1.561698e-04	1.471510e-03	0.1	0.5805784
KEGG04115	24	37.09912863	4.138379e-06	5.360874e-05	0.1	0.5805784
KEGG04330	15	14.41382000	2.758517e-03	1.981900e-02	0.1	0.5805784

KEGG00052	15	19.84974037	4.596460e-04	3.943164e-03	0.1	0.5805784
KEGG00480	11	72.73975947	3.967321e-09	5.186977e-07	0.1	0.5805784
KEGG05050	11	7.68091632	3.389911e-02	2.050977e-01	0.1	0.5805784
KEGG05110	15	13.11246540	4.359517e-03	2.880151e-02	0.1	0.5805784
KEGG04150	17	10.87724970	9.850462e-03	6.290863e-02	0.1	0.5805784
KEGG05060	11	14.20423831	2.967133e-03	2.054732e-02	0.1	0.5805784
KEGG05211	34	3.47752979	1.991449e-01	1.000000e+00	0.2	1.0000000
KEGG05130	26	4.37723967	1.342465e-01	7.491378e-01	0.2	1.0000000
KEGG05131	26	4.37723967	1.342465e-01	7.491378e-01	0.2	1.0000000
KEGG00251	13	6.42852173	5.640018e-02	3.376800e-01	0.2	1.0000000
KEGG04120	12	6.18238811	6.244511e-02	3.700179e-01	0.2	1.0000000
KEGG01430	35	2.74775939	2.760440e-01	1.000000e+00	0.4	1.0000000
KEGG04320	10	1.20129027	5.630263e-01	1.000000e+00	0.7	1.0000000
KEGG00530	10	0.37998588	8.321461e-01	1.000000e+00	0.8	1.0000000
KEGG00510	12	0.07751927	9.630572e-01	1.000000e+00	0.9	1.0000000
KEGG01030	18	0.15071488	9.294974e-01	1.000000e+00	1.0	1.0000000

In the `pcot2` function, the T^2 statistic can be calculated in two ways, using either a pooled estimate of correlation for the two classes (default) or an unpooled estimate. And users can set `var.equal=F` if the correlation structure is assumed to differ across the two classes.

In the first step of the PCOT2 analysis, the dimensionality of the gene expression data is reduced via principal coordinates. The default dimensionality in the `pcot2` function is set as `ncomp=2`. In the second step of the PCOT2 analysis, the distances between the transformed groups are calculated via euclidean distances by default. Other distances (e.g., correlation or Spearman distances) can also be used by defining `dist.method` in the function. A permutation p -value for each category is calculated by re-arranging the sample labels. The permutations can also be performed by permuting rows (genes), using `permu='ByRow'`.

Table 1 lists computation times (in minutes) required to run 1000 permutations of the `pcot2` function on the AML/ALL data under various parameter configurations. The two machines used were a 3.2GHz Pentium 4 with 1Gb RAM running Microsoft Windows XP and R 2.1.0 (PC), and a 1.70GHz Pentium M with 256Mb of RAM running Fedora Core 3 and R 2.2.0 (Unix).

Table 1: *Computation times (minutes, 1000 permutations)*

Changes	PC machine	UNIX machine
default setting	5.6	6.8
<code>var.equal=F</code>	5.5	6.8
<code>comp=8</code>	6	7.6
<code>dist.method="euclidean"</code>	4.8	6
<code>permu="ByRow"</code>	5.6	6.8

4 The corplot and corplot2 functions

The `corplot` and `corplot2` functions enable visualization of both correlation and gene expression information for a particular gene category, in particular the groups identified as being differentially expressed. The plot produced by the

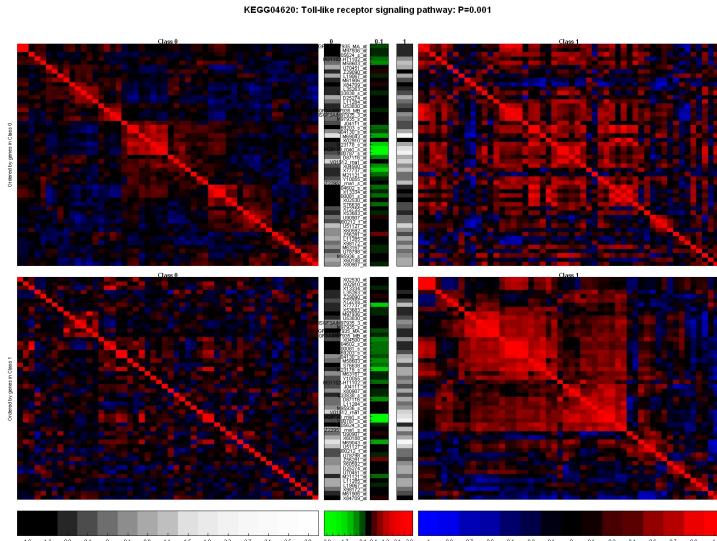


Figure 1: KEGG04620

`corplot` function displays the pooled correlation calculated from the two classes, while the `corplot2` function produces a plot based on unpooled correlation. Gene names can be added to the plot using `add.name=T` (default). The font size can be changed by setting the `font.size` argument. The `main` option specifies the title of the plot.

```
> sel <- c("04620", "04120")
> pvalue <- c(0.001, 0.72)
> library(KEGG)
> pname <- unlist(mget(sel, env = KEGGPATHID2NAME))
> main <- paste("KEGG", sel, ":", pname, ":", "P=", pvalue, sep = "")
> for (i in 1:length(sel)) {
+   fname <- paste("corplot2-KEGG", sel[i], ".jpg", sep = "")
+   jpeg(fname, width = 1600, height = 1200, quality = 100)
+   selgene <- rownames(imat)[imat[, match(paste("KEGG", sel,
+     sep = "")), colnames(imat))] == 1]
+   corplot2(golub, selgene, golub.cl, main = main[i])
+   dev.off()
+ }
```

The argument `inputP` allows users to input the p -values of individual genes calculated using other approaches, such as the limma package (Smyth *et al.*, 2004), allowing the results from both per-gene and per-pathway analysis to be printed on a single plot. To allow users to identify genes from in correlation image plots, the argument `gene.locator=T` allows the selection of interesting (e.g., highly correlated and differential expressed between two classes) genes by clicking beginning and end points on the main diagonal of the image plots. This prints the identifiers for the selected genes. Further details of this functionality are provided in the `HowToUseGeneLocator.pdf` document. The usage of `corplot2` is similar to that for the `corplot` function.

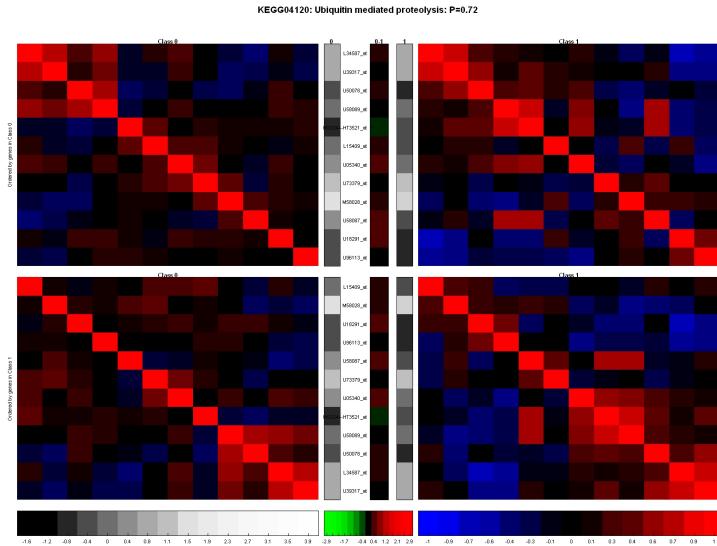


Figure 2: KEGG04120

5 The aveProbes function

In Affymetrix gene expression data, a unique gene can often link to multiple probe sets, with such genes then having a greater influence on the pathway analysis (particularly if the gene is differentially expressed). In order to solve this problem, the `aveProbe` function is provided to change the multiple probe data to the unique gene data by taking the median of the probe values. This function can be used to transform both expression data and the indicator matrix by providing a vector of unique gene identifiers.

```
> pathlist <- as.list(hu6800PATH)
> pathlist <- pathlist[match(rownames(golub), names(pathlist))]
> ids <- unlist(mget(names(pathlist)), env = hu6800SYMBOL))
> newdata <- aveProbe(x = golub, ids = ids)$newx
> output <- aveProbe(x = golub, imat = imat, ids = ids)
> newdata <- output$newx
> newimat <- output$newimat
> newimat <- newimat[, apply(newimat, 2, sum) >= 10]
> dim(newdata)

[1] 2763 38

> dim(newimat)

[1] 2763 103
```

After the multiple probe data set has been changed to the unique gene symbol data, further analysis such as testing and visualizing pathways can be done on the new data set.

References

- [1] Benjamini,B.Y. and Yekutieli,D. (2001) The control of the false discovery rate in multiple testing under dependency. *The Annals of Statistics*, **29**, 1165-1188.
- [2] Gentleman,R.C., Carey,V.J., Bates,D.M., Bolstad,B., Dettling,M., Du-doit,S., Ellis,B., Gautier,L., Ge,Y., Gentry,J. *et al.* (2004) Bioconductor: open software development for computational biology and bioinformatics. *Genome Biology*, **5**, R80.
- [3] Golub,T.R., Slonim,D.K., Tamayo,P., Huard,C., Gaasenbeek,M., Mesirov,J.P., Coller,H., Loh,M.L., Downing,J.R., Caligiuri,M.A. *et al.* (1999) Molecular Classification of Cancer: Class Discovery and Class Prediction by Gene Expression Monitoring, *Science*, **286**, 531-537.
- [4] Smyth,G.K. (2004) Linear models and empirical Bayes methods for assessing differential expression in microarray experiments. *Statistical Applications in Genetics and Molecular Biology*, **3**, No.1, Article 3.