Additional exercesis R bioinfo programming. Master ISCIII. 2014-2015. Second set.

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1 Recommendations

2 Purpose of the exercises

The purpose of this exercise is to get used to obtaining summary statistics from data sets and from subpieces of those data sets.

3 General hints

You will use the leukemia data (including the data, the class and the sex objects we used in the exercise for figures). Here are those instructions again:

3.1 The data

• Read the data "leukemia.data.txt". Note that there are row names that we do not want to be part of the data itself (look at the help for "read.table", the option "row.names"). Call this "leuk.dat" These data are based on the famous leukemia data from Golub et al. I've modified a few things below: the gene names and the sex of the patients are invented.

- That will be a data frame. For most figures your life will be simpler if you convert it into a matrix, using data.matrix. Call this data matrix "leuk.dat.m". (Why is it a good idea to convert this into a matrix? Try doing the figures using the original data frame.)
- Read the data "leukemia.class.txt". Use scan for this (do you know why?). Since these are labels, use what = "" in scan. Convert the classes into a factor. Call this "leuk.class".
- Create a vector for the sex of the patients. The patients are Male and Female, alternating, and staring with Male. Call this factor "sex".

4 Tables and cross-tabs

Obtain a simple table of how many males and females are there. Use table on the "sex" object:

```
## sex
## Female Male
## 19 19
```

Now, a table of how many of each sex are there in each class, or viceversa. This requires you to pass two arguments to table

```
## leuk.class
## sex ALL AML
## Female 13 6
## Male 14 5
```

(if your table looks transposed with respect to the above ... well, change the order of arguments). Now, do the same as above, but with a formula interface, which some times is more useful. Use the xtabs function

```
## leuk.class
## sex ALL AML
## Female 13 6
## Male 14 5
```

Finally, rerun the code from xtabs and assign that to an object, call it "mytable" and convert mytable to a data frame using as .data.frame and print it:

```
## sex leuk.class Freq
## 1 Female ALL 13
## 2 Male ALL 14
## 3 Female AML 6
## 4 Male AML 5
```

Notice that this is just a very simple but extremely powerful way of getting frequencies and saving them as a data frame for further analysis!! With the output from the tables we could do a chi-square. Or we could use the data frame for some other statistical analysis (e.g., logistic regressions, etc). Oh and for real we do not need to save all those intermediate objects.

5 Two subsetting operations

Since getting access to selected parts of matrices, data frames, etc, is so important, two quick exercises. First, obtain the mean of all the genes for the third subject, but only for those genes with a p-value less than 0.01. Or, to put it another way, you want to see the average (mean) expression of the genes in subject 3, but only for the genes that have a p-value < 0.01:

```
## [1] 0.01816
```

And now the median of the second gene, but ONLY for males

```
## [1] -1.045
```

6 Gene summaries by condition and sex

The next exercises all involve summaries of specific genes. Remember that genes were originally the rows of the data set. Since we will be using aggregate and similar, you will want to have genes as columns. How? Doing a transposition:

```
leuk.dat.t <- t(leuk.dat.m)</pre>
```

Make sure you understand what happened. For instance, do:

```
dim(leuk.dat.t)
## [1] 38 3051
```

and now check the first few columns (you do not want to use head, because it would show all 3051 columns)

```
leuk.dat.t[1:5, 1:6]

## G1 G2 G3 G4 G5 G6

## V2 -1.458 -0.75161 0.45695 3.1353 2.766 2.6434

## V3 -1.394 -1.26278 -0.09654 0.2142 -1.270 1.0142

## V4 -1.428 -0.09052 0.90325 2.0875 1.604 1.7048

## V5 -1.407 -0.99596 -0.07194 2.2347 1.532 1.6384

## V6 -1.427 -1.24245 0.03232 0.9381 1.637 -0.3608
```

6.1 aggregate: The median of three genes by condition

Use aggregate to obtain the median of genes in positions 1, 2124, and 2600 (HK-1, PTEN, and the other gene from the figures in the exercises from last week)

```
## type G1 G2124 G2600
## 1 ALL -1.361 -0.2167 -0.1101
## 2 AML -1.230 1.5763 1.3253
```

6.2 aggregate: The median of three genes by condition and sex

Like the previous exercise, but you want to obtain those summaries by condition and sex:

```
## type sex G1 G2124 G2600

## 1 ALL Female -1.368 0.07727 -0.00771

## 2 AML Female -1.320 1.71842 1.13218

## 3 ALL Male -1.321 -0.46043 -0.16511

## 4 AML Male -1.089 1.45710 1.55580
```

6.3 aggregate: The median of all the genes by condition and sex

This is like the previous one, but for **every** gene. So you do not want to print it. Assign the result of aggregate to an object, lets call it "all.median".

Now, show all the rows and the first 10 columns

```
## type sex G1 G2 G3 G4 G5 G6 G7

## 1 ALL Female -1.368 -1.0321 0.23381 0.6418 0.2285 0.9687 3.288

## 2 AML Female -1.320 -0.9223 0.14707 0.3602 -0.5765 -0.1641 3.126

## 3 ALL Male -1.321 -1.0346 0.26632 2.2069 1.7145 1.6147 3.087

## 4 AML Male -1.089 -1.0890 0.04609 1.3030 1.0160 0.5127 3.279

## G8

## 1 3.040

## 2 2.901

## 3 2.985

## 4 2.961
```

and the dimensions of "all.median"

```
dim(all.median)
## [1] 4 3053
```

6.4 aggregate: The mean and standard deviations of three genes by condition and sex

This is like the above ones, but we do not want just one statistic (the median); here we want the mean and standard deviation for all our three genes (1, 2124, 2600). And, if possible, with clear column names, so you will have to be explicit, when you pass a function to aggregate, about the names of the components you return. (In other words, DO NOT assign the return value of aggregate and change the column names; you have to play around with the names in the return object from the function you pass to aggregate)

```
## type
        sex G1.mean G1.sd G2124.mean G2124.sd G2600.mean
## 1 ALL Female -1.3180 0.1567 -0.1633 0.5816 -0.23135
## 2 AML Female -1.1570 0.3504
                              1.6656 0.3161 1.30764
## 3 ALL Male -1.2283 0.4254
                               -0.4169 0.8363
                                               -0.09673
## 4 AML Male -0.3260 1.1930
                               1.4919 0.4386
                                               1.42469
  G2600.sd
##
## 1 0.63892
## 2 0.50587
## 3 0.78109
## 4 0.40976
```

6.5 by and aggregate: those three genes again, but now use "summary"

As it says. First, do not use aggregate now, but use by, and the function you want to have applied to each subset is summary, which will give you different summary statistics.

```
## type: ALL
## sex: Female
##
       G1
                     G2124
                                     G2600
## Min.
        :-1.61 Min. :-1.3683 Min. :-1.4009
  1st Qu.:-1.40 1st Qu.:-0.4010
                                  1st Qu.:-0.6645
##
## Median :-1.37
                 Median : 0.0773
                                  Median :-0.0077
##
  Mean :-1.32
                Mean :-0.1633 Mean :-0.2313
##
  3rd Qu.:-1.22
                 3rd Qu.: 0.2502
                                  3rd Qu.: 0.1770
## Max. :-1.03 Max. : 0.6263 Max. : 0.5734
```

```
## type: AML
## sex: Female
             G2124 G2600
   G1
##
  Min. :-1.444 Min. :1.23 Min. :0.816
##
##
  1st Qu.:-1.410 1st Qu.:1.46 1st Qu.:0.929
##
  Median :-1.320 Median :1.72 Median :1.132
##
  Mean :-1.157 Mean :1.67 Mean :1.308
##
  3rd Qu.:-0.894 3rd Qu.:1.81
                           3rd Qu.:1.728
  Max. :-0.665 Max. :2.11 Max. :1.977
##
## -----
## type: ALL
## sex: Male
           G2124
##
  G1
                                G2600
## Min. :-1.568 Min. :-1.476 Min. :-1.2305
  1st Ou.:-1.427 1st Ou.:-1.225 1st Ou.:-0.5857
##
  Median :-1.321 Median :-0.460 Median :-0.1651
##
  Mean :-1.228 Mean :-0.417 Mean :-0.0967
##
  3rd Qu.:-1.234 3rd Qu.: 0.392
##
                              3rd Qu.: 0.3977
##
  Max. : 0.176 Max. : 0.579 Max. : 1.4982
## -----
## type: AML
## sex: Male
            G2124
##
                               G2600
      G1
  Min. :-1.262 Min. :1.05 Min. :0.814
##
  1st Qu.:-1.230 1st Qu.:1.19 1st Qu.:1.218
##
  Median :-1.089 Median :1.46
                           Median :1.556
##
## Mean :-0.326 Mean :1.49 Mean :1.425
## 3rd Qu.: 0.849 3rd Qu.:1.58 3rd Qu.:1.737
## Max. : 1.101 Max. :2.18 Max. :1.800
```

(Of course, if you wanted, you could do that to all of the genes in the array) Now, use aggregate:

```
## type sex G1.Min. G1.1st Qu. G1.Median G1.Mean G1.3rd Qu.
## 1 ALL Female -1.610 -1.400 -1.370 -1.320 -1.220
## 2 AML Female -1.440
                        -1.410
                                -1.320 -1.160
## 3 ALL Male -1.570 -1.430 -1.320 -1.230 
## 4 AML Male -1.260 -1.230 -1.090 -0.326
                                                -1.230
                                                 0.849
  G1.Max. G2124.Min. G2124.1st Qu. G2124.Median G2124.Mean
##
## 1 -1..
## 2 -0.665 1.2500
-1.4800
## 1 -1.030 -1.3700 -0.4010 0.0773 -0.1630
                                    1.7200
             1.2300 1.4600
-1.4800 -1.2300
                                              1.6700
                                   1.7200 1.6700
-0.4600 -0.4170
## 4 1.100 1.0500 1.1900 1.4600
                                             1.4900
## G2124.3rd Ou. G2124.Max. G2600.Min. G2600.1st Ou. G2600.Median
## 1
        2.1100 0.81600
                                      0.92900
                                                 1.13000
## 2
         1.8100
## 3
                   0.5790
                           -1.23000
                                      -0.58600
         0.3920
                                                 -0.16500
    1.5800 2.1800 0.81400
                                      1.22000
## 4
                                                 1.56000
## G2600.Mean G2600.3rd Qu. G2600.Max.
## 1 -0.23100 0.17700 0.57300
## 2
     1.31000
                 1.73000 1.98000
                0.39800 1.50000
## 3 -0.09670
## 4 1.42000 1.74000 1.80000
```

Please, notice the differences in output. They are different kinds of objects, something you can see if you assign the output to an object and do, for instance

class(objeto)