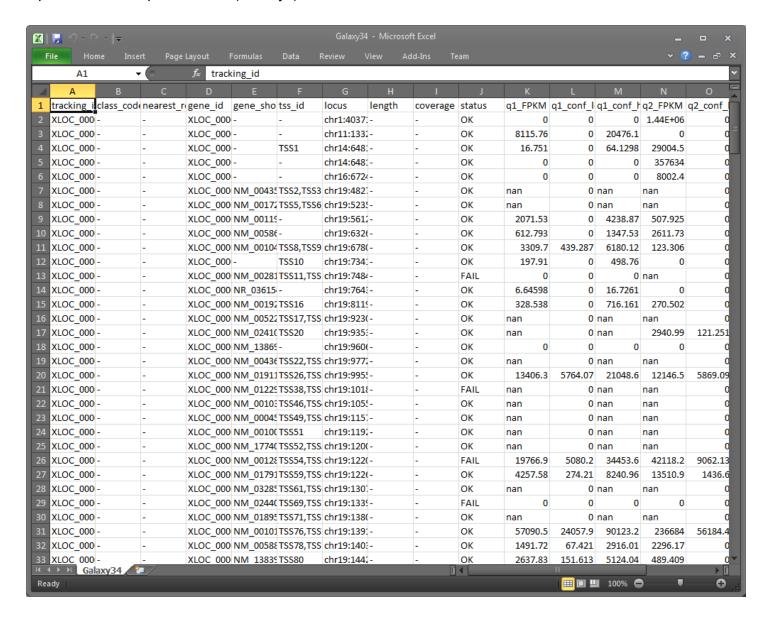
A preview of the input text data (3 arrays) in Excel.

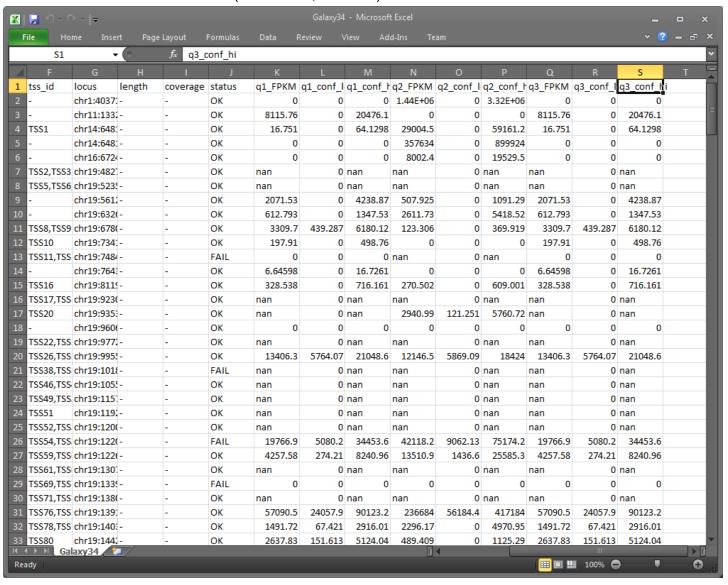


### The data header looks like

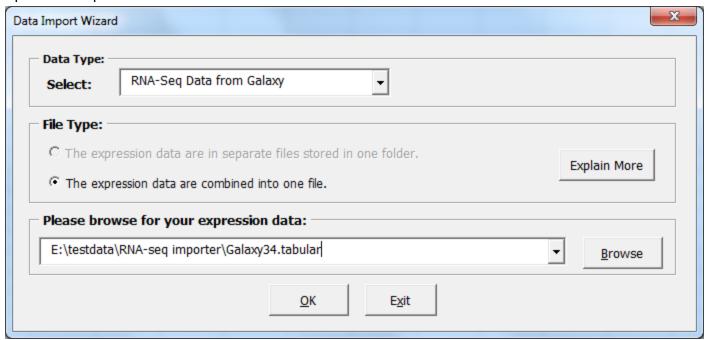
```
> x <- scan("clipboard", "")</pre>
Read 19 items
> x
 [1] "tracking id"
                        "class code"
                                            "nearest ref id"
                                                               "gene id"
 [5] "gene short name" "tss id"
                                            "locus"
                                                               "length"
                                            "q1 FPKM"
 [9] "coverage"
                        "status"
                                                               "q1 conf lo"
                                            "q2 conf lo"
[13] "q1 conf hi"
                        "q2 FPKM"
                                                               "q2 conf hi"
[17] "q3 FPKM"
                        "q3 conf lo"
                                            "q3 conf hi"
```

Each array occupies 3 columns in the input file although only FPKM column will be used.

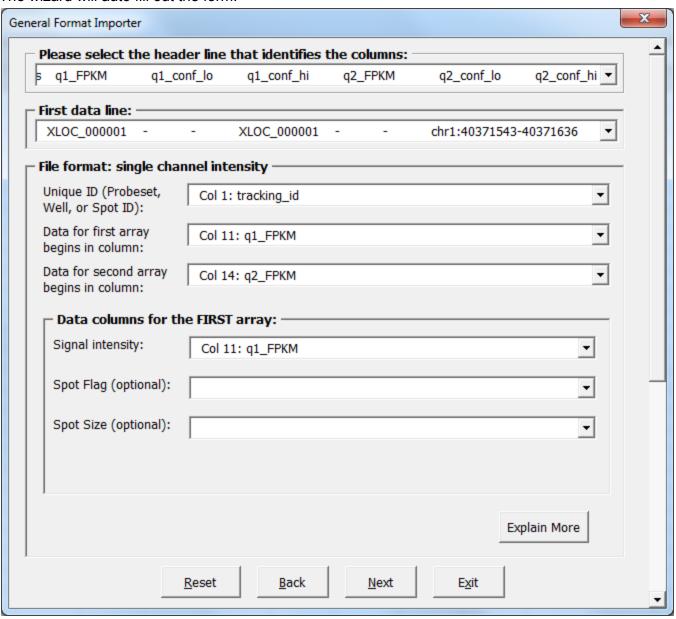
## A close look at the FPKM columns (Columns K, N and Q)

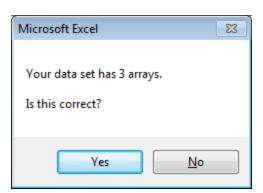


# Open data import wizard

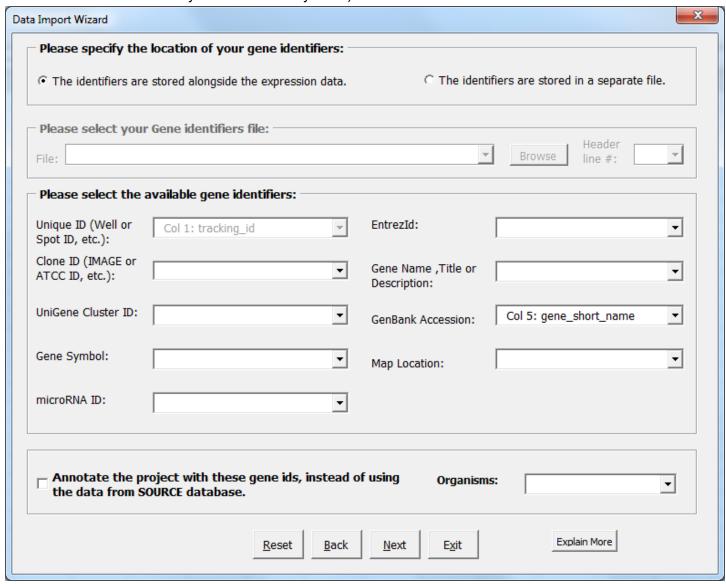


The wizard will auto fill out the form.

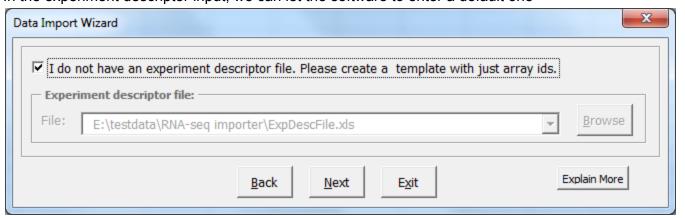




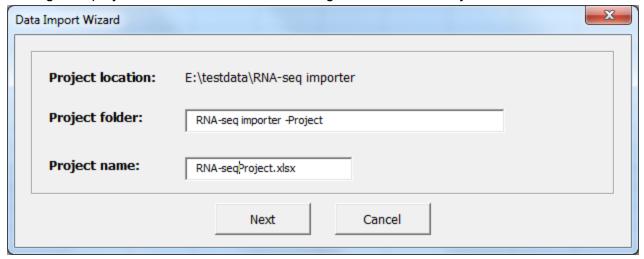
Select 'Col 5: gene\_short\_name' from the dropdown list of 'GenBank Accession' (RefSeq Accession) for annotation use (if I did not select any gene identifiers, the dataset will not be able to run annotation though we can still run most of analyses in BRB-ArrayTools).



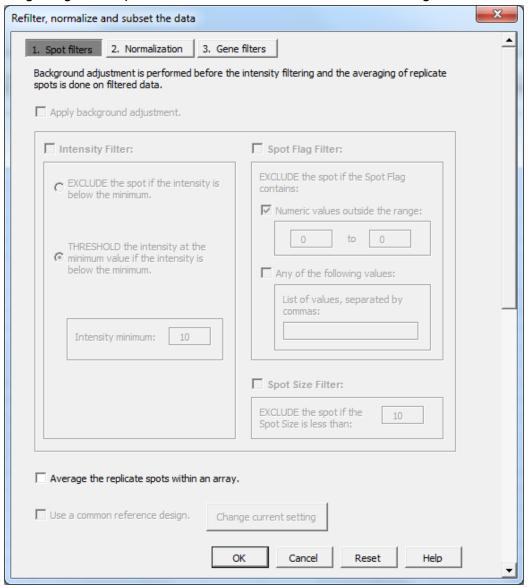
In the experiment descriptor input, we can let the software to enter a default one



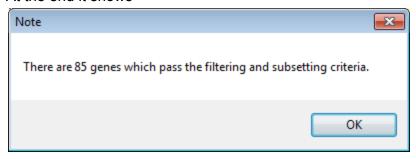
I change the project name from its default although it is not necessary



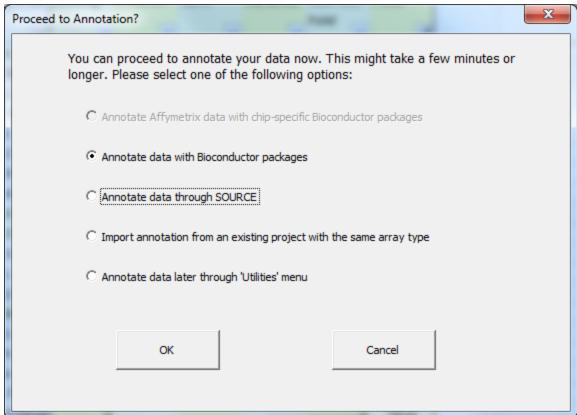
Program goes to spot filter, normalization and subset data dialog. Just click 'OK'



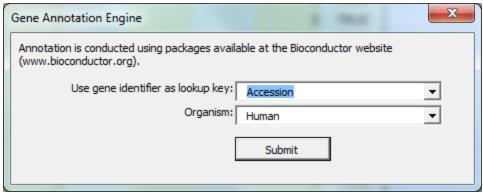
### At the end it shows



It will prepare to run annotation.

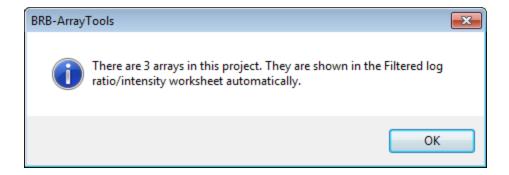


Since my data has only accession inform, this gene identifier is the only option as lookup key.

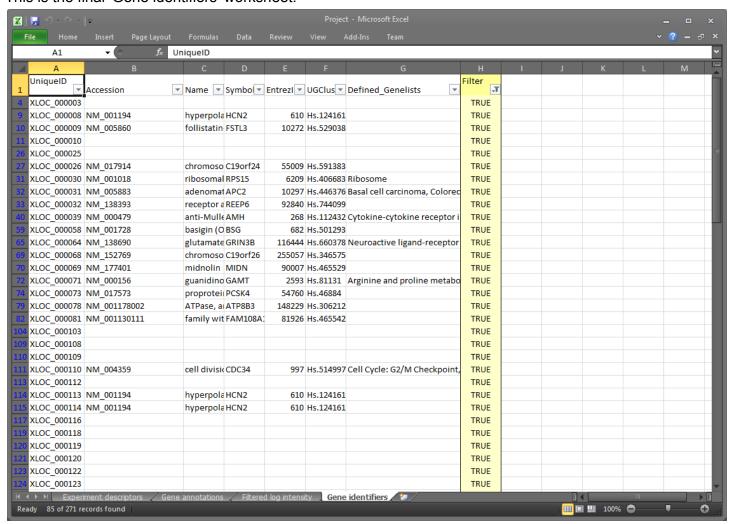


2 Tk windows will pop up to show the progress .... Be patient.

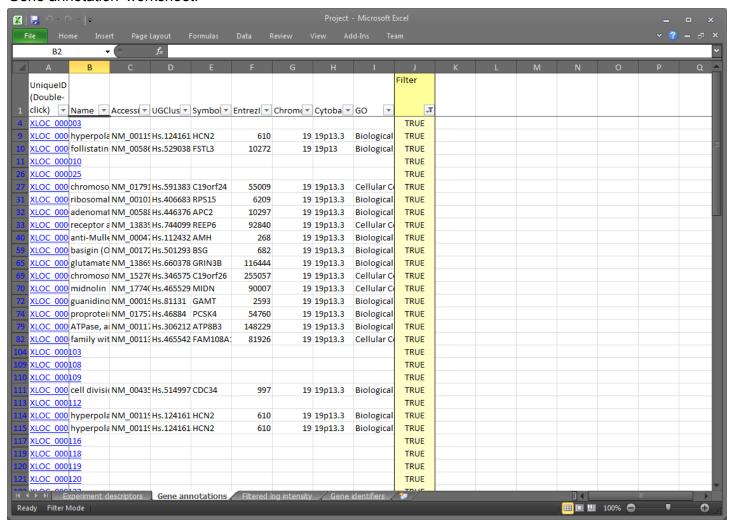




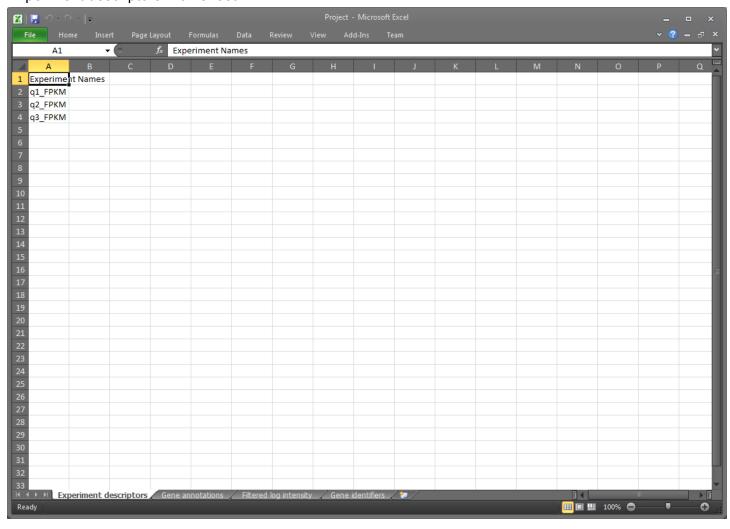
This is the final 'Gene identifiers' worksheet.



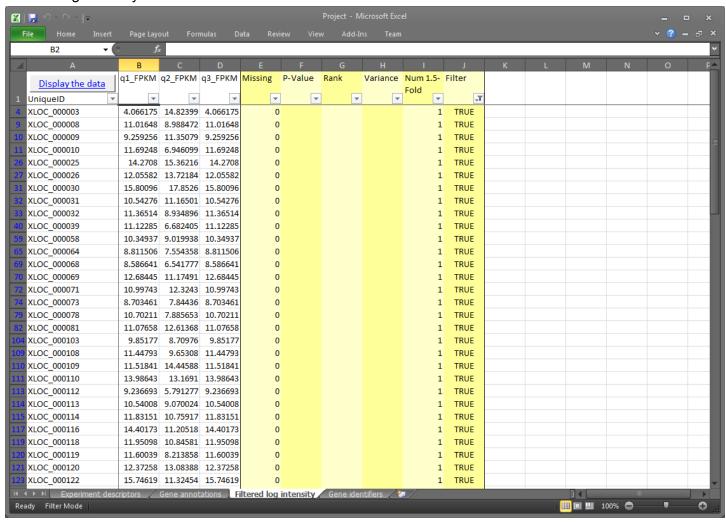
#### 'Gene annotation' worksheet.



# 'Experiment descriptors' worksheet



## 'Filtered log intensity' worksheet



## DONE!

We can double check the log intensity value. For example, UniqueID=XLOC\_000003 shall have log2(16.751)=4.066175.