



## ExpressionFileCreator Documentation

**Description:** Creates a RES or GCT file from a set of Affymetrix CEL files  
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### Summary

The ExpressionFileCreator module creates an expression dataset from a set of individual Affymetrix CEL files. The conversion is done using the standard Affymetrix probe modeling algorithm MAS5, the RMA algorithm, the GCRMA algorithm, or the dChip algorithm. The result is a matrix containing one intensity value per probe set, in the GCT or RES file format described at [http://www.broad.mit.edu/cancer/software/genepattern/tutorial/gp\\_fileformats.html](http://www.broad.mit.edu/cancer/software/genepattern/tutorial/gp_fileformats.html).

Samples can be annotated by specifying a clm file. A clm file allows you to change the name of the samples in the expression matrix, reorder the columns, select a subset of the scans in the input zip file, and create a class label file in the cls format, also described on the web page above.

By default, sample names are taken from the CEL file names contained in the zip file. A clm file allows you to specify the sample names explicitly. Additionally, the columns in the expression matrix are reordered so that they are in the same order as the scan names appear in the clm file. For example, the input zip file contains the files scan1.cel, scan2.cel, and scan3.cel. The clm file could contain the following text:

```
scan3      sample3      tumor
scan1      sample1      tumor
scan2      sample2      normal
```

The column names in the expression matrix would be: sample3, sample1, sample2. Additionally, only scan names in the clm file will be used to construct the GCT or RES file; scans not present in the clm file will be ignored.

### Notes:

- The MAS5 and dChip algorithms are based on their Bioconductor implementations. Therefore the results obtained from these algorithms will differ slightly from their official implementations.

### References:

1. Affymetrix. Affymetrix Microarray Suite User Guide. Affymetrix, Santa Clara, CA, version 5 edition, 2001.
2. Rafael A. Irizarry, Bridget Hobbs, Francois Collin, Yasmin D. Beazer-Barclay, Kristen J. Antonellis, Uwe Scherf, and Terence P. Speed. Exploration, normalization, and summaries of high density oligonucleotide array probe level data. Biostatistics, 2003b. To appear.
3. Li, C. and Wong, W.H. (2001) Genome Biology 2, 1–11.
4. Li, C. and Wong, W.H. (2001) Proc. Natl. Acad. Sci USA 98, 31–36.

### Parameters:

input file	A zip file of CEL files
method	The method to use
quantile normalization	(GCRMA and RMA only) Whether to normalize data using quantile normalization

# GenePattern

background correct	(RMA only) Whether to background correct using RMA background correction
compute present absent calls	Whether to compute Present/Absent calls
normalization method	(MAS5 only) The normalization method to apply after expression values are computed. The column having the median of the means is used as the reference unless the parameter value to scale to is given.
value to scale to	(median/mean scaling only) The value to scale to.
clm file	A tab-delimited text file containing one scan, sample, and class per line
annotate probes	Whether to annotate probes with the gene symbol and description
cdf file	Custom CDF file. Leave blank to use default internally provided CDF file.
output file	The base name of the output file

## Output Files:

1. gct or res file
2. cls file if clm file is supplied

## Platform dependencies:

**Module type:** Preprocess & Utilities  
**CPU type:** any  
**OS:** any  
**Language:** R