

# **GS01 0163**

## **Analysis of Microarray Data**

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## Lecture 2: The Basics of dChip

- So, why are we here?
- Getting the stuff required
- Using dChip
  - Loading Data
  - Looking at Data
  - Normalizing Data
  - Model Fitting
  - Exporting Results
- The Real World...

## So, why are we here?

We want to learn about dChip.

The freeware package dChip has become quite widely used for the analysis of Affymetrix gene chip data. We're going to look at using it now.

The main web page for dChip is

<http://biosun1.harvard.edu/complab/dchip/>

where you can download the software, get links to some publicly available data, and browse through the online manual.

Much of this lecture will follow the manual, and the associated "Short tutorial" and "Lab" with my editorial comments.

## Step 1: Get dChip

This step is fairly trivial; simply download the latest version (dchip2004.exe as of July 13, 2005) and put the application somewhere (eg, D:Program Files/dChip2004/). We keep this application on a shared drive at

`/data/bioinfo/affymetrix/00 Affymetrix Info/DChip Files`

The entire application is a bit over 1M in size. At present, dChip only runs on Windows platforms. Some success has been reported using windows emulators on the Mac, but there is a performance hit.

## A Biological Example

There is a genetic translocation that occurs in ALL, associated with a mixed-lineage leukemia gene (MLL). Patients with this translocation have noticeably worse outcomes. It is thought that this translocation may make the disease qualitatively different, and somewhat closer to AML. If the disease is different, we may want to adjust the therapy as well.

Using Affymetrix gene chips, can we identify differences between ALL, MLL, and AML?

## Step 2: Get CEL Files

The Lab page supplies a link to the example data CEL files on leukemia (ALL,MLL,AML) from Dana Farber.

```
http://www.broad.mit.edu/cgi-bin/cancer/  
publications/pub_paper.cgi?mode=view&paper_id=63
```

The CEL files are available as gzipped tar files, which WinZip should be able to uncompress. There are 6 CEL file collections at this site, each about 35-41M in size, or 100-127M in size when uncompressed. These files contain about 10-12 CEL files each.

The suffixes on these files should be .tar.gz, but for some reason they are tar.tar. This latter suffix needs to be changed so that the file type will be recognized.

## Step 2: Get CEL Files (cont)

If you are working with CEL files stored in more than one location, it is often useful to assemble a “data list file” specifying the locations of the files. This file should be a text file (and end in .txt). Every row should contain either a specific file name or a directory. An example from the manual:

```
E:\Affy data\dan\CA-H.cel  
E:\Affy data\dan\CA-HR.cel  
E:\Affy data\dan\zugen  
E:\Affy data\dan\PC-C.cel
```

Here, the AML samples were run later, so we put them in a different directory.

## Step 2A: Digression

The Dana Farber web site also supplies the quantifications that they used in their analyses, as

`expression_data.txt`

or

`expression_data_plus_APcalls.txt`

These data were initially quantified using MAS4.0 (AvDiff). We prefer to work with the CEL files as raw data and to construct our own quantifications.



## **Step 3: Get Explanatory Files**

Also at the above site, there are files describing the sample-to-chip mapping in more detail:

scaling\_factors\_and\_fig\_key.txt

and a link to the paper that appeared in Nature Genetics describing the biological context of the problem.

## Step 4: Find the CDF file

This requires that we know what type of Affy chip was used. In this case (according to the paper), the chips were U95A.

For this example, a compressed version of the CDF file can be downloaded from the dChip site; more generally, we have a collection of CDF files for the chip types we use in

`data/bioinfo/affymetrix/00 Affymetrix Info/CDF Files`

A warning – the `cdf` extension is also used for “channel files” by Microsoft, so don’t worry if you see a weird icon.

## Step 4A: Digression

Actually, the CDF file for these chips is a bit tricky.

There is a set of U95 chips, U95A,U95B,..,U95E that contain probes for all genes in the genome. The probes were assembled using the 95th build of the Unigene database to define what a “gene” was. However, while these chips surveyed the genome, most of the probes corresponding to “interesting” genes were put on the A chip, so most people just bought those as opposed to the set.

Soon after the U95A release, some mistakes were noted in the probe design, and Affy released the U95Av2, which is the type we have encountered more frequently here at MDA.

Can you tell them apart?

## Step 5: Get the Gene Info file(s)

Every chip type has a fixed set of probesets printed on it, but the probeset identifiers are typically not enough to suggest anything (1389\_at?). We need more context – is there a common name for the associated gene? Which chromosome is it on, and where? Is the gene known or thought to be part of a functional family (eg, cytoskeleton)? Are there IDs that can let us look up more information in national databases?

The above information for each chip type has been collected and assembled into GeneInfo files available at the dChip website.

*These files are tab-delimited text files, but they've had an xls extension placed on them so that Excel is the default program for opening them.*

These info files can change over time!

## Step 5: Get the Gene Info file(s) (cont)

Actually, when we download the zip file from the dChip web site, we get 3 files:

HG-U95Av2 gene info2.xls

HG-U95Av2 gene info2 Gene Ontology.xls

HG-U95Av2 gene info2 Protein Domain.xls

We're going to look at each of these in turn, but I want to quickly note that these files are for the U95Av2 chip, as opposed to the U95A chip. In terms of the probesets that were used, the overlap is so large (12600 of 12625) that working with these should be fine.

These files are on our system in

/data/bioinfo/affymetrix/00 Affymetrix Info/DChip Files

## HG-U95Av2 gene info2.xls

The first few entries:

```

Probe Set Name : Identifier : LocusLink :
Name : Gene Ontology.xls : Protein Domain.xls :
Pathway : Chromosome : Description
1000_at : X60188 : 5595 :
mitogen-activated protein kinase 3 : |7165|
7154|6935|42330|9605|6928|8151|4707|4702|4674|
4672|16301|3824|16773|16772|16740|5057|4871| :
|2290|719|3527| : : |16|16p|16p12| :
X60188 /FEATURE=mRNA /DEFINITION=HSERK1 Human ERK1
mRNA for protein serine/threonine kinase
1001_at : X60957 : 7075 :
tyrosine kinase with immunoglobulin and epidermal
growth factor homology domains : |7498|9888|

```

# HG-U95Av2 gene info2 Gene Ontology.xls

Term ID	Term Name	Frequency	
3	reproduction	101	
18	regulation of DNA recombination	9	
41	transition metal transport	16	
67	DNA replication and chromosome cycle		103
70	mitotic chromosome segregation	7	
72	M-phase specific microtubule process		8
74	regulation of cell cycle	330	
75	cell cycle checkpoint	35	
76	DNA replication checkpoint	8	

# HG-U95Av2 gene info2 Protein Domain.xls

Term ID	Term Name	Frequency
1	Kringle 16	
2	Cdc20/Fizzy	4
3	Retinoid X receptor	15
4	Saposin type B	5
5	Helix-turn-helix, AraC type	11
6	Vertebrate metallothionein, family 1	6
7	Tubby	7
8	C2 domain	84
10	Cysteine proteases inhibitor	18

“what ghastly names they all have...” E. J. (Ernest John) Moncrieff



## Step 6: Get the Sample Info file

Most of the files that we have worked with so far have described properties associated with a given chip type, not with the samples we have used. We can also supply and use sample-specific information in a tab-delimited text file. The first few entries here:

scan name	sample_name	type
CL2001011101AA	ALL_1	A
CL2001011104AA	ALL_2	A
CL2001011105AA	ALL_3	A
CL2001011108AA	ALL_4	A
CL2001011109AA	ALL_5	A
CL2001011111AA	ALL_6	A
CL2001011112AA	ALL_7	A
CL2001011116AA	ALL_8	A
CL2001011113AA	ALL_9	A

## Step 6: Get the Sample Info file (cont)

The header row and the first two columns are required, but any columns beyond that are at our discretion. By default, column values are treated as factors, but adding the string “(numeric)” to a column name will override this.

What else could we have included?

- Presence/absence of other translocations
- train/test status
- specimen type (diagnostic, relapse)
- run date...

## **Step 7: write a README file**

Strictly speaking, this is not mentioned in the Lab or Tutorial, but I'll put it here, right before actually running the program.

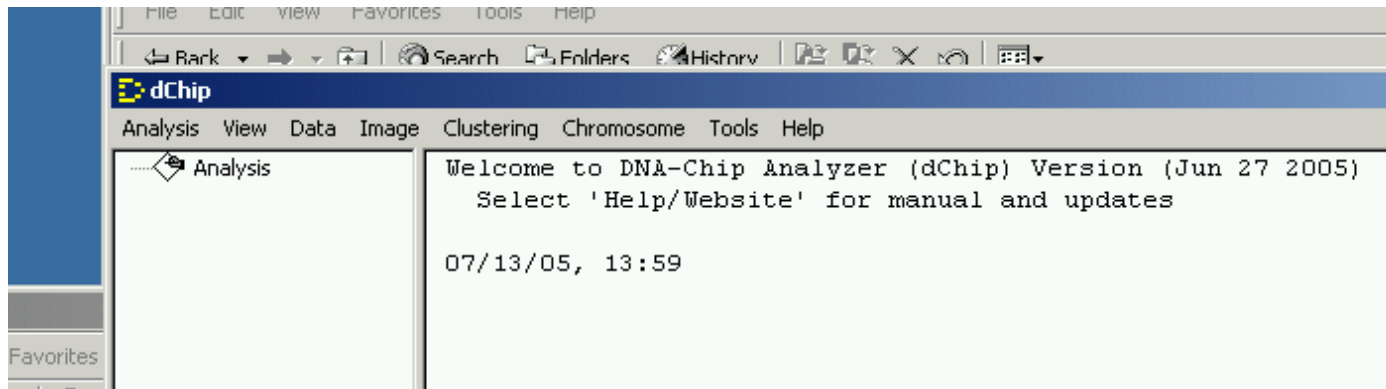
What is the biological question you are seeking to address?

What contrasts of data samples will allow you to address this?

Sending a brief description of this type off to the investigator before running the analysis can save some time...

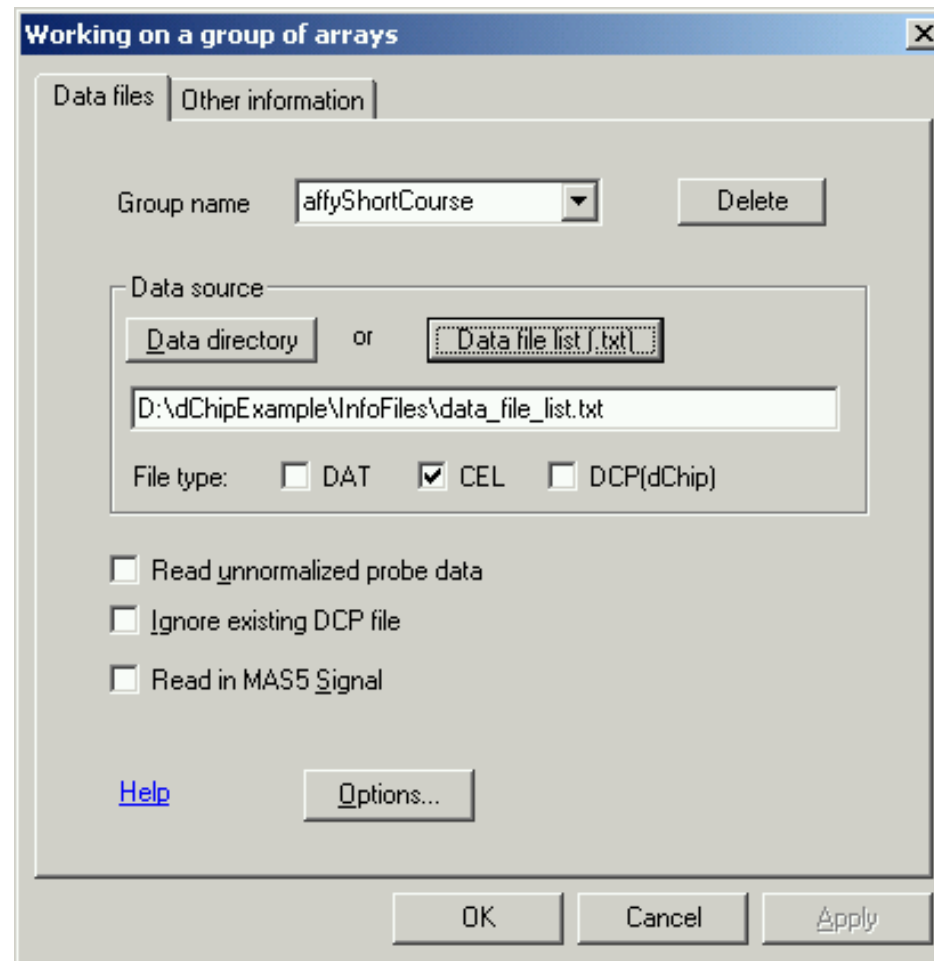
## Step 8: run dChip

Nice, friendly, unexciting...



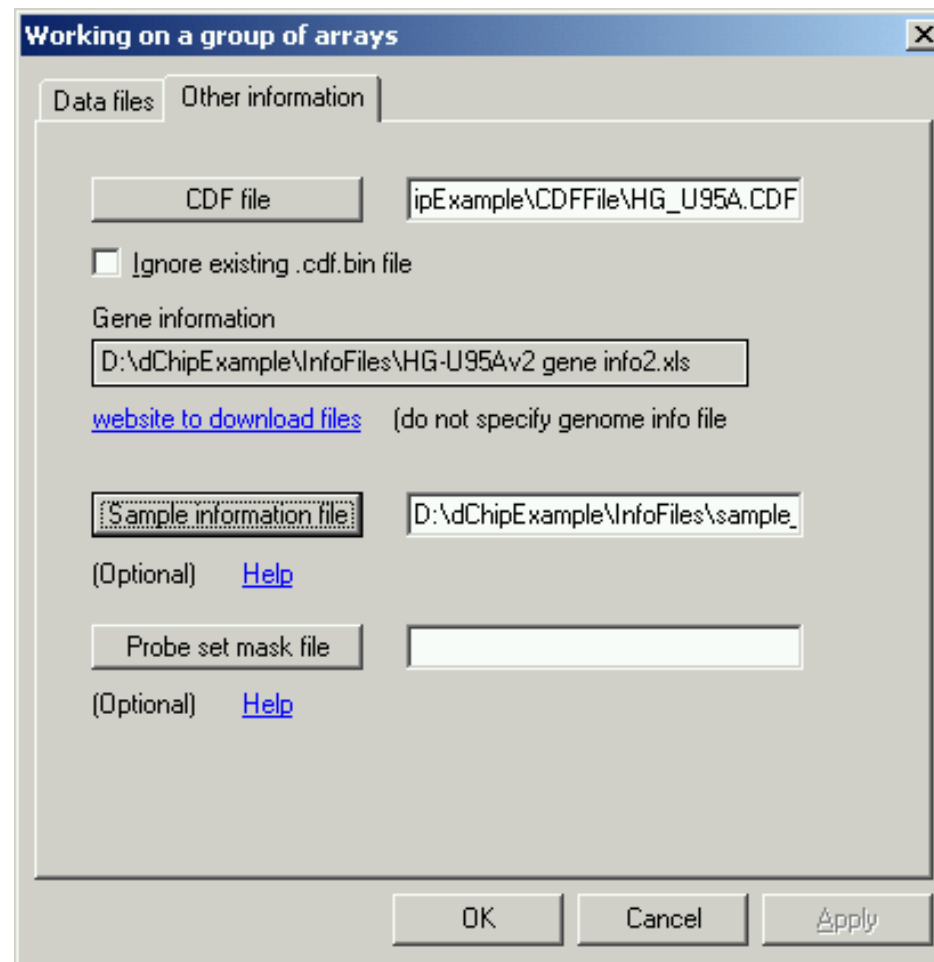
Now, we need to tell it where to find the data for analysis. Go to Analysis/Open Group.

# Finding files, part 1



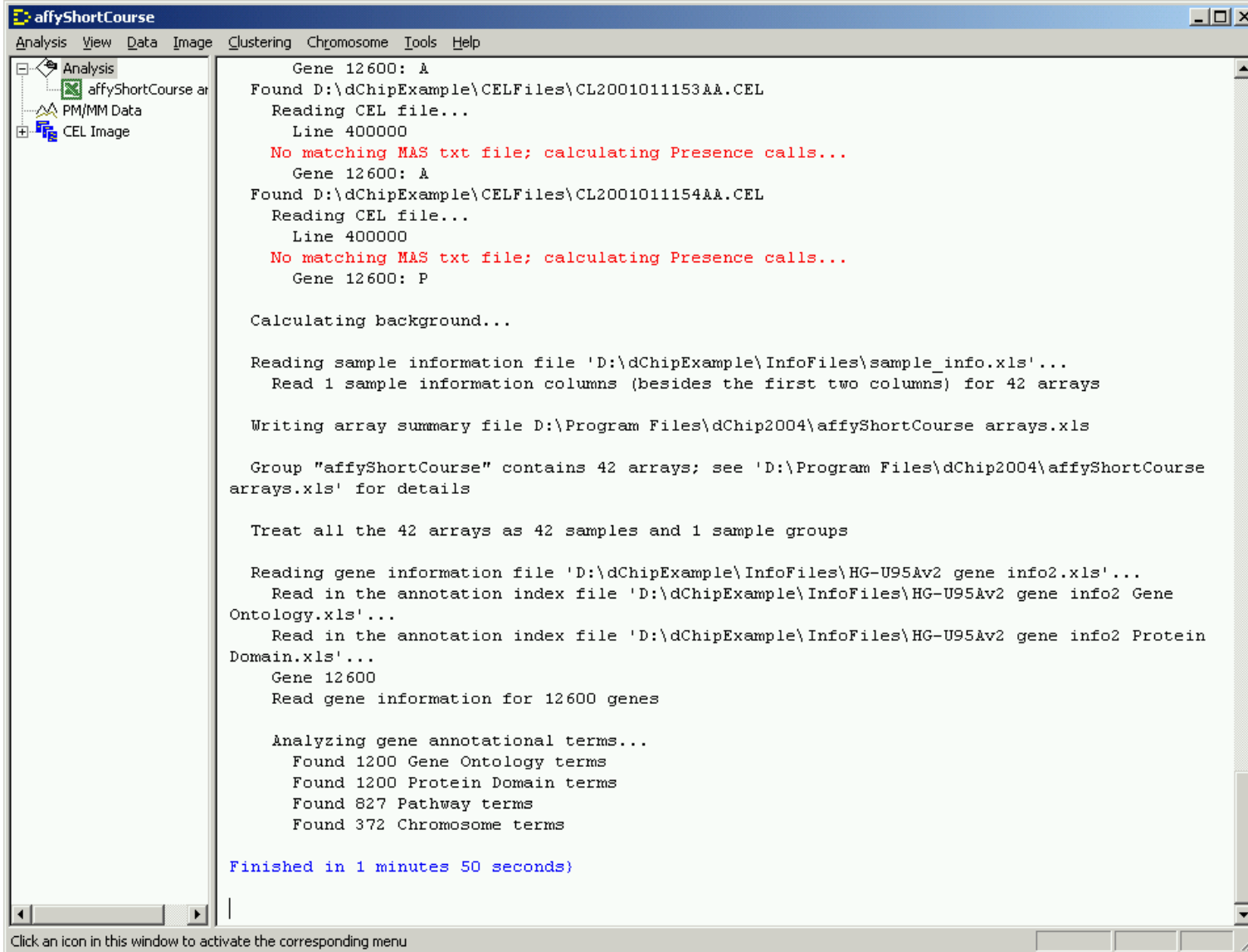
assign a group name, locate data files

## Finding files, part 2



locate CDF, gene info, and sample info files. Under Options, we can set the working directory where results should be stored.

# Reading files



The screenshot shows the 'affyShortCourse' application window. The title bar reads 'affyShortCourse'. The menu bar includes 'Analysis', 'View', 'Data', 'Image', 'Clustering', 'Chromosome', 'Tools', and 'Help'. The left sidebar contains a tree view with 'Analysis' expanded, showing sub-items: 'affyShortCourse ar', 'PM/MM Data', and 'CEL Image'. The main window displays a log of operations:

```
Gene 12600: A
Found D:\dChipExample\CELFiles\CL2001011153AA.CEL
Reading CEL file...
Line 400000
No matching MAS txt file; calculating Presence calls...
Gene 12600: A
Found D:\dChipExample\CELFiles\CL2001011154AA.CEL
Reading CEL file...
Line 400000
No matching MAS txt file; calculating Presence calls...
Gene 12600: P

Calculating background...

Reading sample information file 'D:\dChipExample\InfoFiles\sample_info.xls'...
Read 1 sample information columns (besides the first two columns) for 42 arrays

Writing array summary file D:\Program Files\dChip2004\affyShortCourse arrays.xls

Group "affyShortCourse" contains 42 arrays; see 'D:\Program Files\dChip2004\affyShortCourse
arrays.xls' for details

Treat all the 42 arrays as 42 samples and 1 sample groups

Reading gene information file 'D:\dChipExample\InfoFiles\HG-U95Av2 gene info2.xls'...
Read in the annotation index file 'D:\dChipExample\InfoFiles\HG-U95Av2 gene info2 Gene
Ontology.xls'...
Read in the annotation index file 'D:\dChipExample\InfoFiles\HG-U95Av2 gene info2 Protein
Domain.xls'...
Gene 12600
Read gene information for 12600 genes

Analyzing gene annotational terms...
Found 1200 Gene Ontology terms
Found 1200 Protein Domain terms
Found 827 Pathway terms
Found 372 Chromosome terms

Finished in 1 minutes 50 seconds)
```

At the bottom of the window, there is a status bar with the text: 'Click an icon in this window to activate the corresponding menu'.

## What has been wrought?

For each CEL file, a binary “dcp” file has been produced:

CL2001011101AA.CEL	10,393	KB
CL2001011101AA.dcp	1,764	KB
CL2001011102AA.CEL	10,324	KB
CL2001011102AA.dcp	1,764	KB

$$(2 * 640^2) * 2 = 1638400$$

Keep the means as 16-bit integers, and allocate space for 2 CEL equivalents in each dcp file – 1 for the raw data, and 1 for the processed data.

This saves space, and uses an intelligent data structure.



## What has been wrought?

A binary version of the CDF file has been produced for quicker processing.

HG_U95A.CDF	29,814	KB
HG_U95A.CDF.bin	5,492	KB

## What has been wrought?

3 interim files have been produced:

dChip.ini

affyShortCourse.ini

affyShortCourse arrays.xls

The first two are configuration files, and are stored with the exe file. The last summarizes some aspects of the files examined, and is stored in the working directory.

# The dChip.ini file

## dChip.ini

CDF\_FILE=

READ\_DAT=0

READ\_CEL=1

READ\_DCP=0

DATA\_PATH=D:\Program Files\dChip2004

WORKING\_DIR=D:\Program Files\dChip2004

GOSURFER\_DIR=D:\Program Files\dChip2004

USE\_UNNORM=0

MAS5\_SIGNAL=0

# The affyShortCourse.ini file

## affyShortCourse.ini

CDF\_FILE=D:\dChipExample\CDFFile\HG\_U95A.CDF

READ\_DAT=0

READ\_CEL=1

READ\_DCP=0

DATA\_PATH=D:\dChipExample\InfoFiles\data\_file\_list.txt

WORKING\_DIR=D:\Program Files\dChip2004

GOSURFER\_DIR=D:\Program Files\dChip2004

USE\_UNNORM=0

MAS5\_SIGNAL=0

# The affyShortCourse arrays.xls file

## affyShortCourse arrays.xls

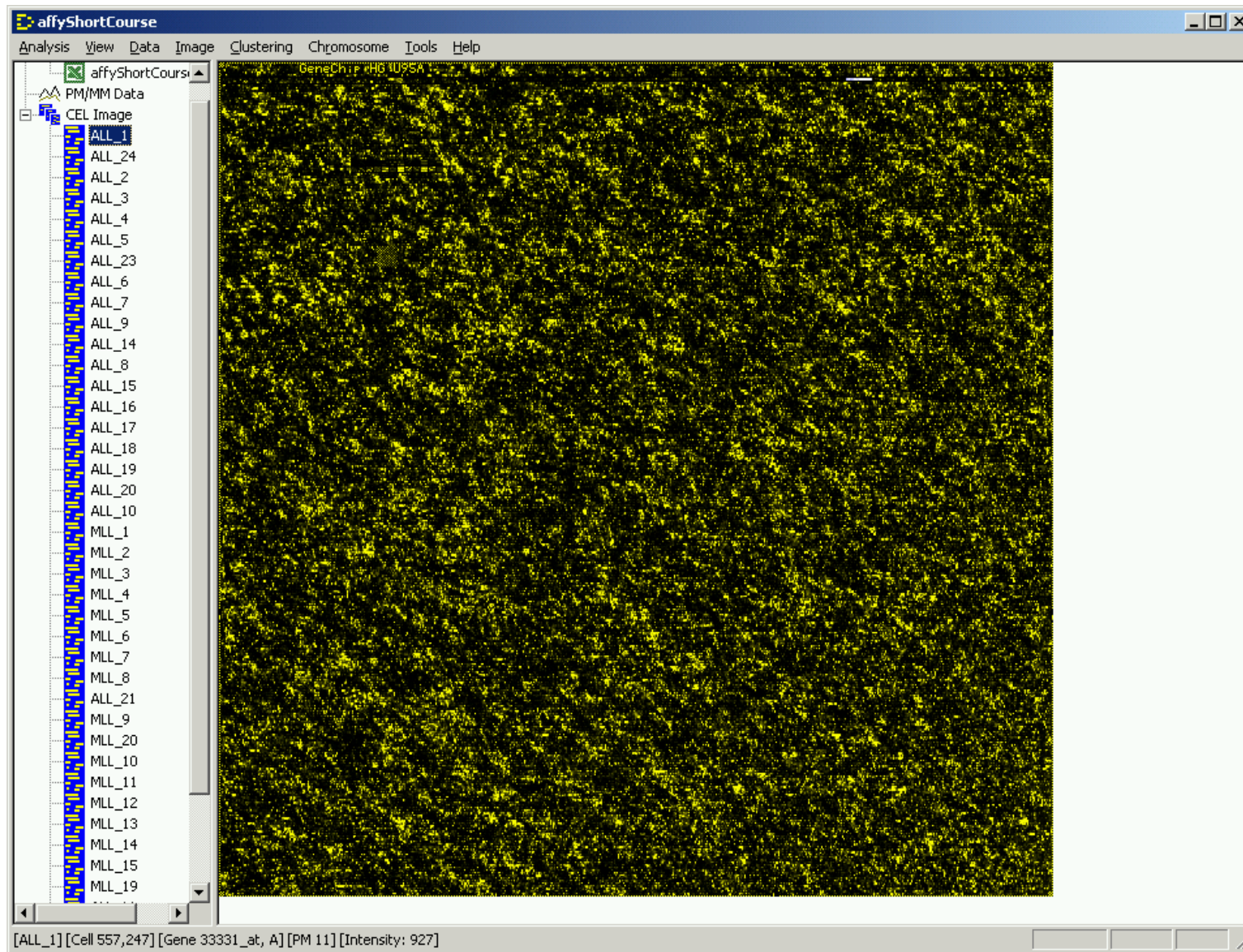
```
Number : Array : File Name : Median Intensity  
(unnormalized) : P call %  
1 : ALL_1 : D:\dChipExample\CELFiles\  
CL2001011101AA.CEL : 1497 : 48.2  
2 : ALL_24 : D:\dChipExample\CELFiles\  
CL2001011102AA.CEL : 1196 : 38.3  
3 : ALL_2 : D:\dChipExample\CELFiles\  
CL2001011104AA.CEL : 1778 : 49.5  
4 : ALL_3 : D:\dChipExample\CELFiles\  
CL2001011105AA.CEL : 1097 : 36.8
```

## Look at the Chips

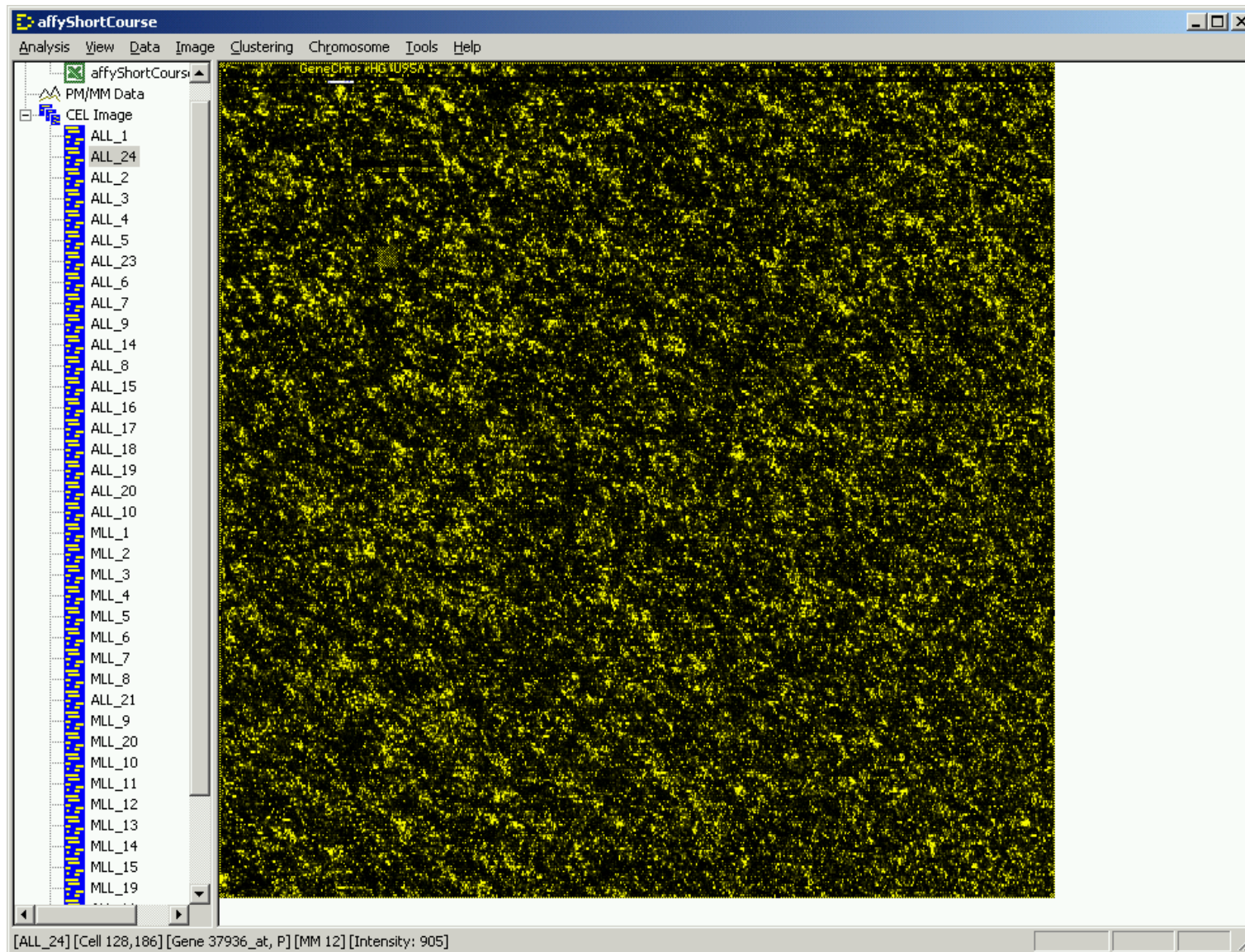
The “Short Tutorial” next suggests going to “View/CEL Image” to look at the data. Unfortunately, this is for an earlier version of dChip, as this pulldown option no longer exists.

So, we click on the “CEL Image” icon at the left of the display and cycle through. If you click on one of the file names, the up and down arrows will let you cycle through them, or Page Up/Page Down also works. The display range covers from the 1st percentile (black) to the 95th (bright yellow).

# Look at the First Chip: ALL\_1



# Look at the Second Chip: ALL\_24





## Zoom In

If you click on a part of the image, you select the corresponding probe set. The arrow keys will let you zoom in on the image to look at that spots more closely.

Down arrow: zoom in lots

Up arrow: zoom out lots

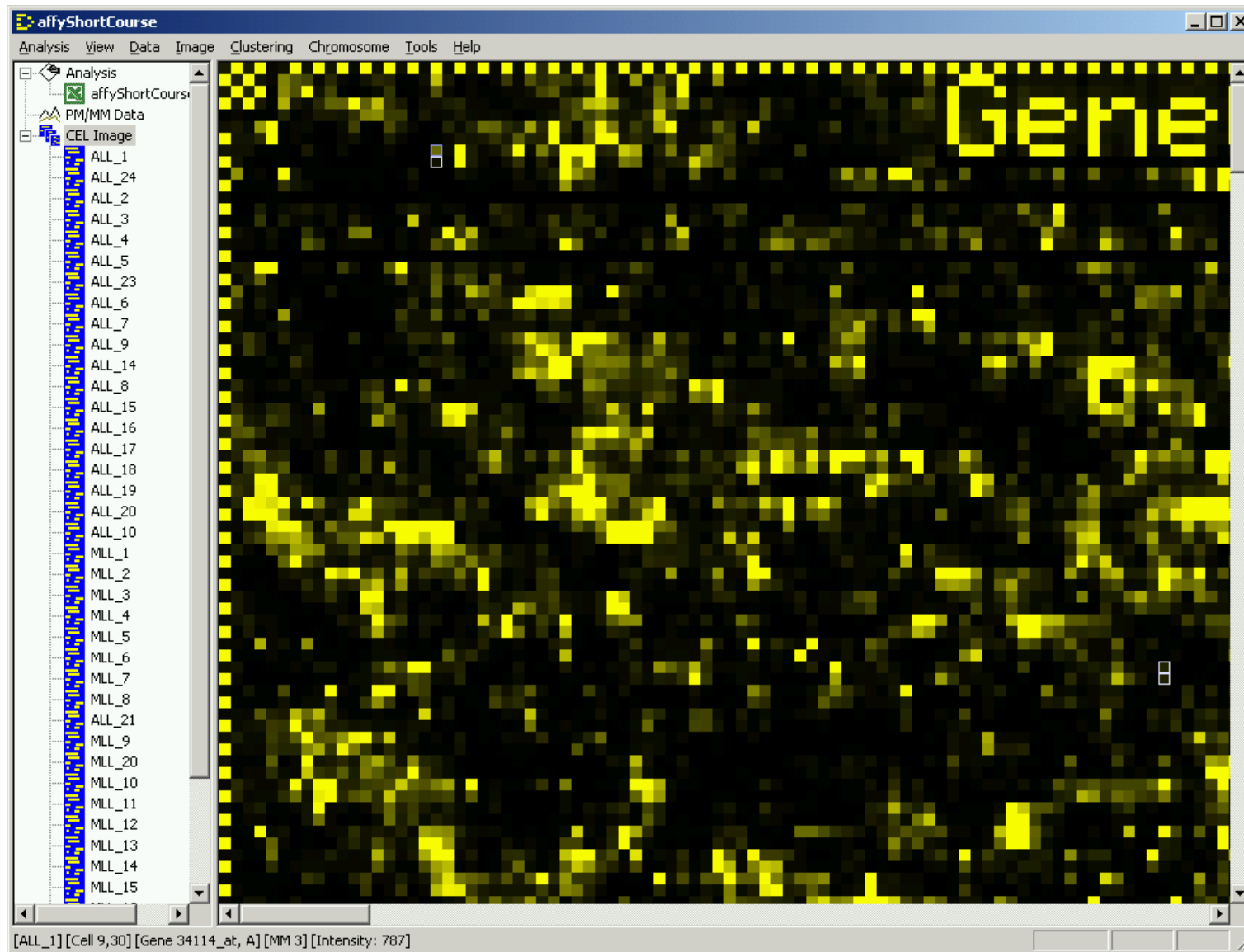
Right arrow: zoom in a little

Left arrow: zoom out a little

Scrollbars move about

Page Up and Page Down cycle you through the set of chips.

# Zoom In: ALL\_1



## Normalize the data

go to Analysis/Normalize

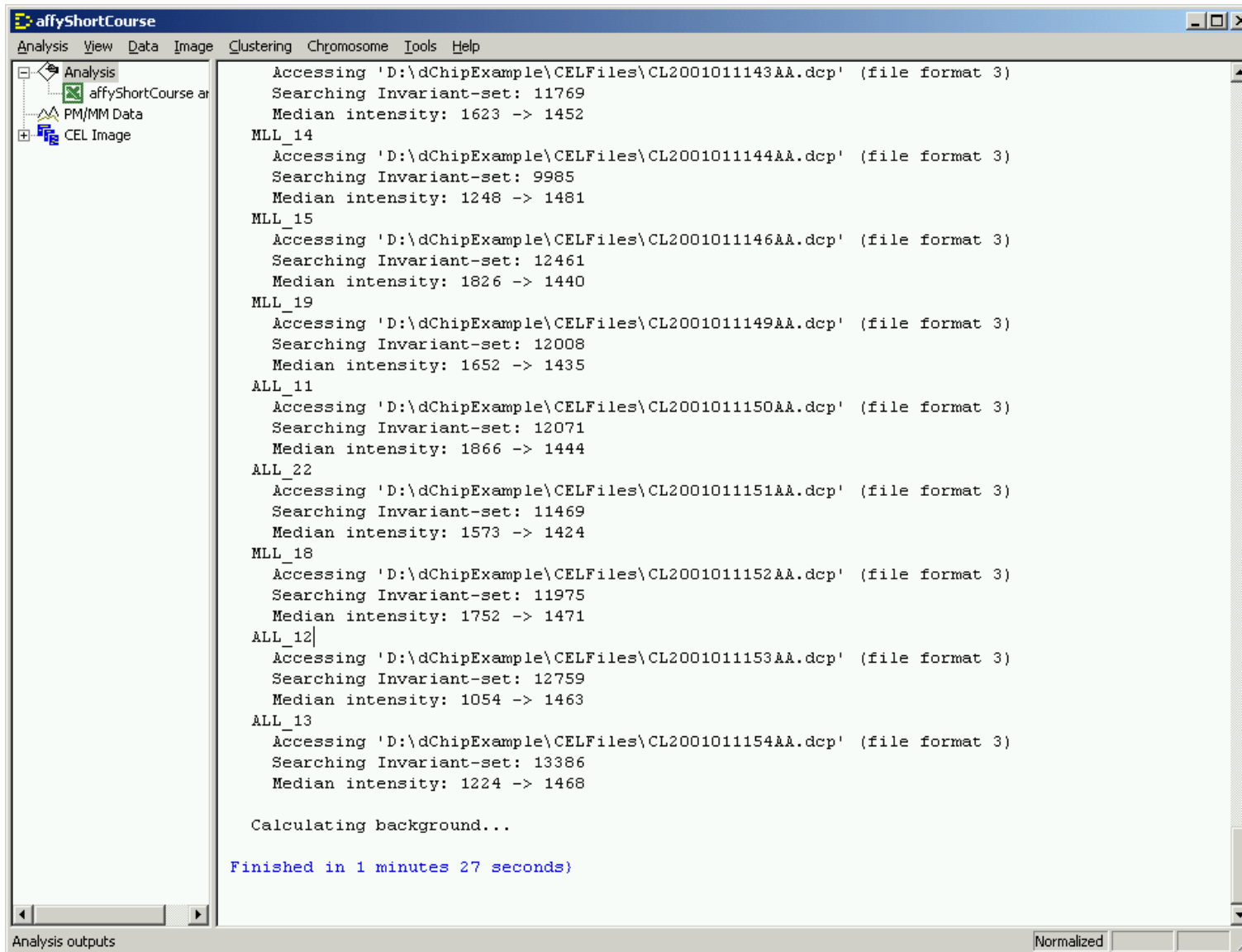
dChip will pick one array in the set to normalize all of the others to; by default it will choose the array with the median overall feature intensity.

(This can make a difference. Trying it with at least two different chips is recommended.)

For each chip, dChip then calculates an “invariant set” of features whose ranks do not change a great deal, and uses those to define a normalization curve.

Functionally, this often works like quantile normalization to the target chip.

# Is it normalized?



The screenshot shows the affyShortCourse application window. The left sidebar contains a tree view with 'Analysis' selected, showing sub-items for 'affyShortCourse ar', 'PM/MM Data', and 'CEL Image'. The main window displays a list of analysis outputs for various CEL files, including MLL\_14, MLL\_15, MLL\_19, ALL\_11, ALL\_22, MLL\_18, ALL\_12, and ALL\_13. Each entry shows the file path, the number of invariant genes searched, and the median intensity before and after normalization. The process concludes with 'Calculating background...' and 'Finished in 1 minutes 27 seconds'. A 'Normalized' checkbox is visible at the bottom right of the window.

```
Analysis View Data Image Clustering Chromosome Tools Help
Analysis
  affyShortCourse ar
  PM/MM Data
  CEL Image

Accessing 'D:\dChipExample\CELFiles\CL2001011143AA.dcp' (file format 3)
Searching Invariant-set: 11769
Median intensity: 1623 -> 1452
MLL_14
Accessing 'D:\dChipExample\CELFiles\CL2001011144AA.dcp' (file format 3)
Searching Invariant-set: 9985
Median intensity: 1248 -> 1481
MLL_15
Accessing 'D:\dChipExample\CELFiles\CL2001011146AA.dcp' (file format 3)
Searching Invariant-set: 12461
Median intensity: 1826 -> 1440
MLL_19
Accessing 'D:\dChipExample\CELFiles\CL2001011149AA.dcp' (file format 3)
Searching Invariant-set: 12008
Median intensity: 1652 -> 1435
ALL_11
Accessing 'D:\dChipExample\CELFiles\CL2001011150AA.dcp' (file format 3)
Searching Invariant-set: 12071
Median intensity: 1866 -> 1444
ALL_22
Accessing 'D:\dChipExample\CELFiles\CL2001011151AA.dcp' (file format 3)
Searching Invariant-set: 11469
Median intensity: 1573 -> 1424
MLL_18
Accessing 'D:\dChipExample\CELFiles\CL2001011152AA.dcp' (file format 3)
Searching Invariant-set: 11975
Median intensity: 1752 -> 1471
ALL_12
Accessing 'D:\dChipExample\CELFiles\CL2001011153AA.dcp' (file format 3)
Searching Invariant-set: 12759
Median intensity: 1054 -> 1463
ALL_13
Accessing 'D:\dChipExample\CELFiles\CL2001011154AA.dcp' (file format 3)
Searching Invariant-set: 13386
Median intensity: 1224 -> 1468

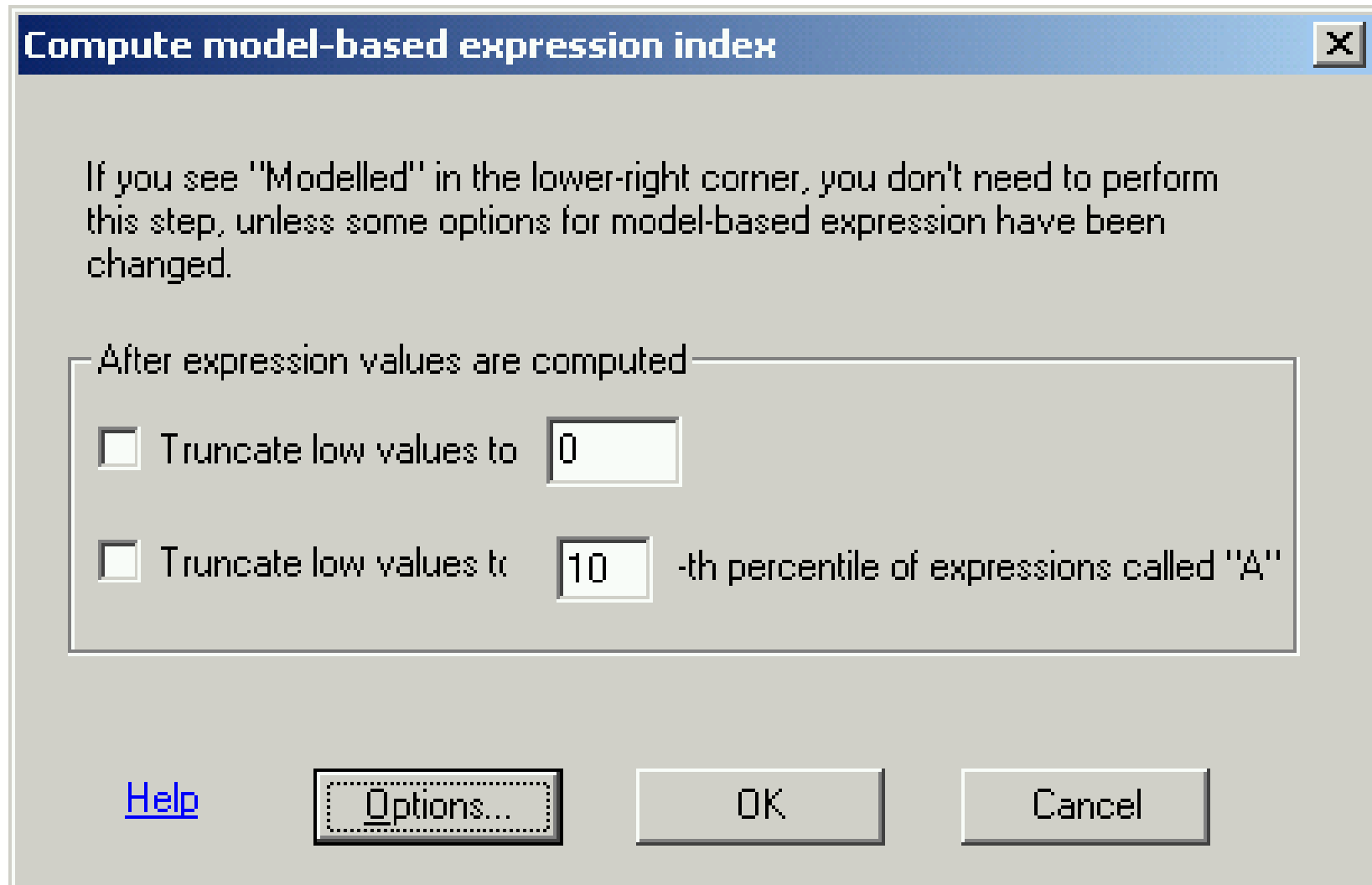
Calculating background...

Finished in 1 minutes 27 seconds)

Analysis outputs Normalized
```

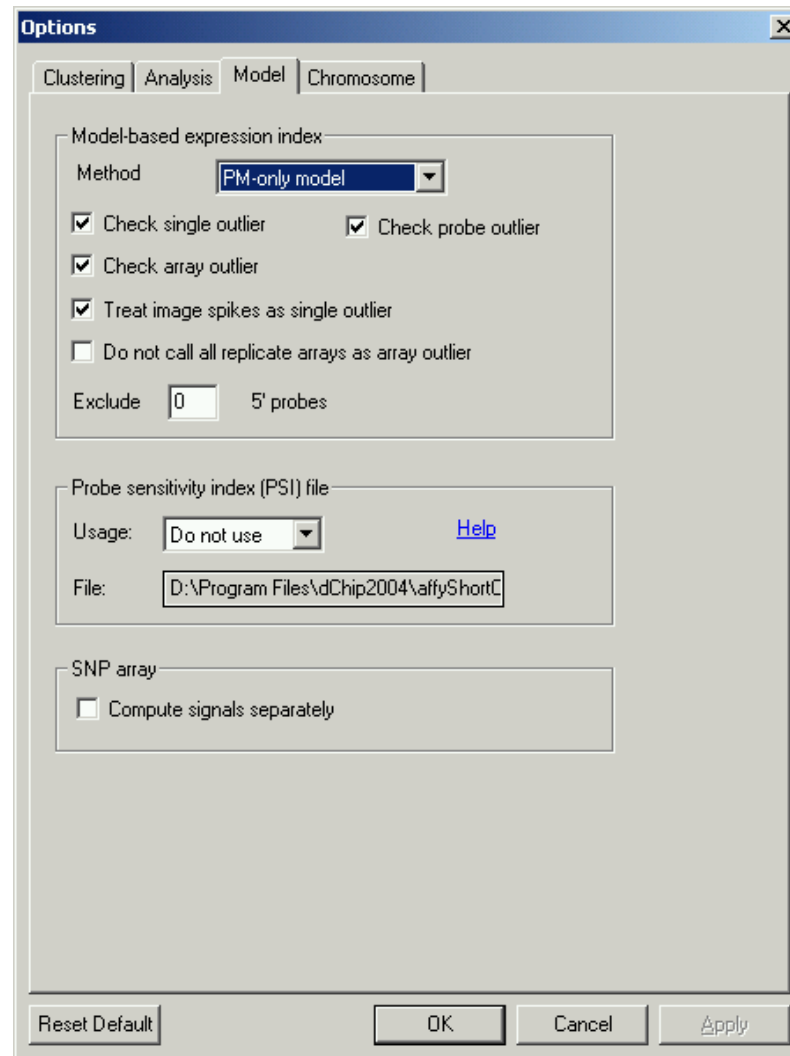
# Fit the Model 1

go to Analysis/Model-based Expression

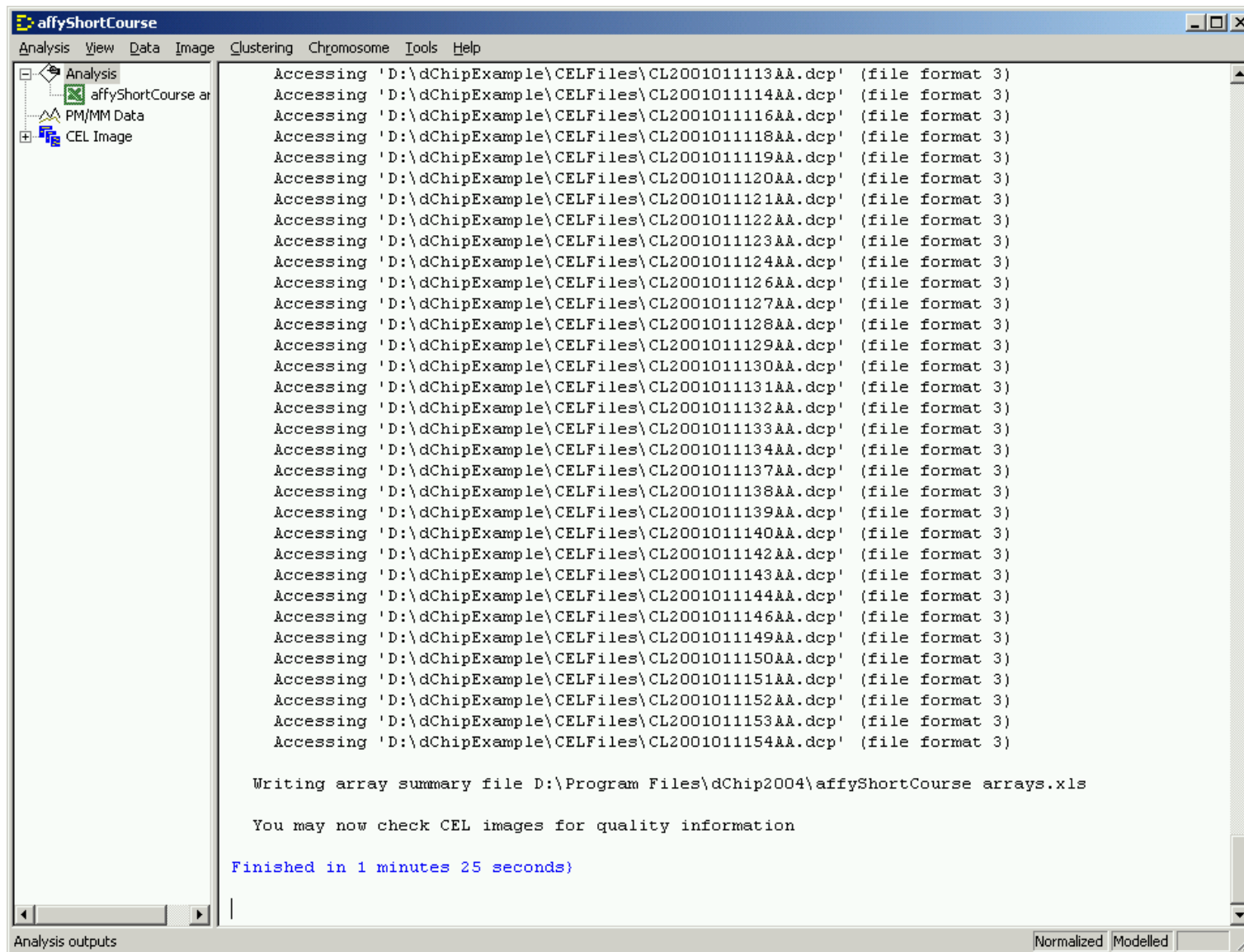


# Fit the Model 2

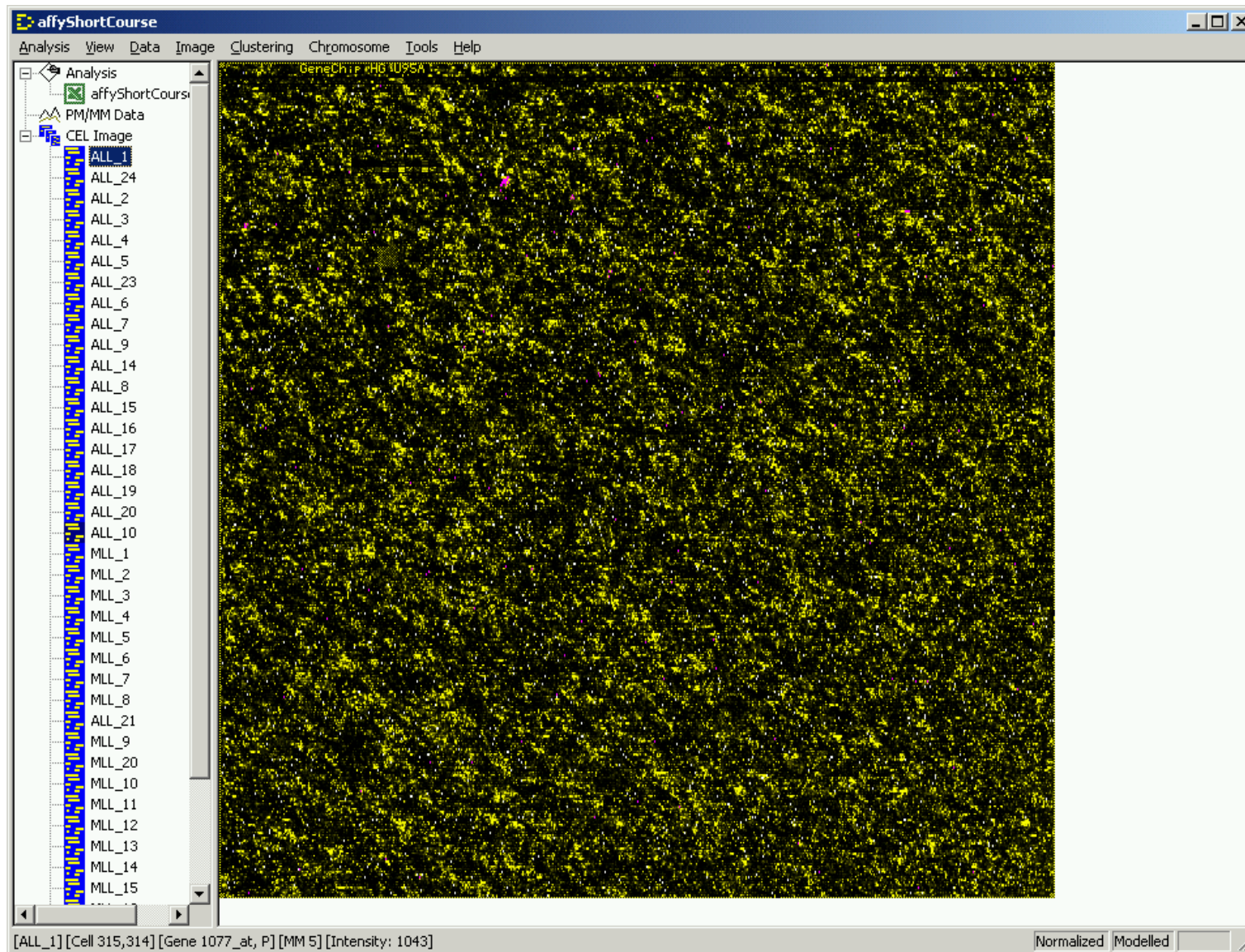
Choose “Options” and select the PM-only model



# Fit the Model 3

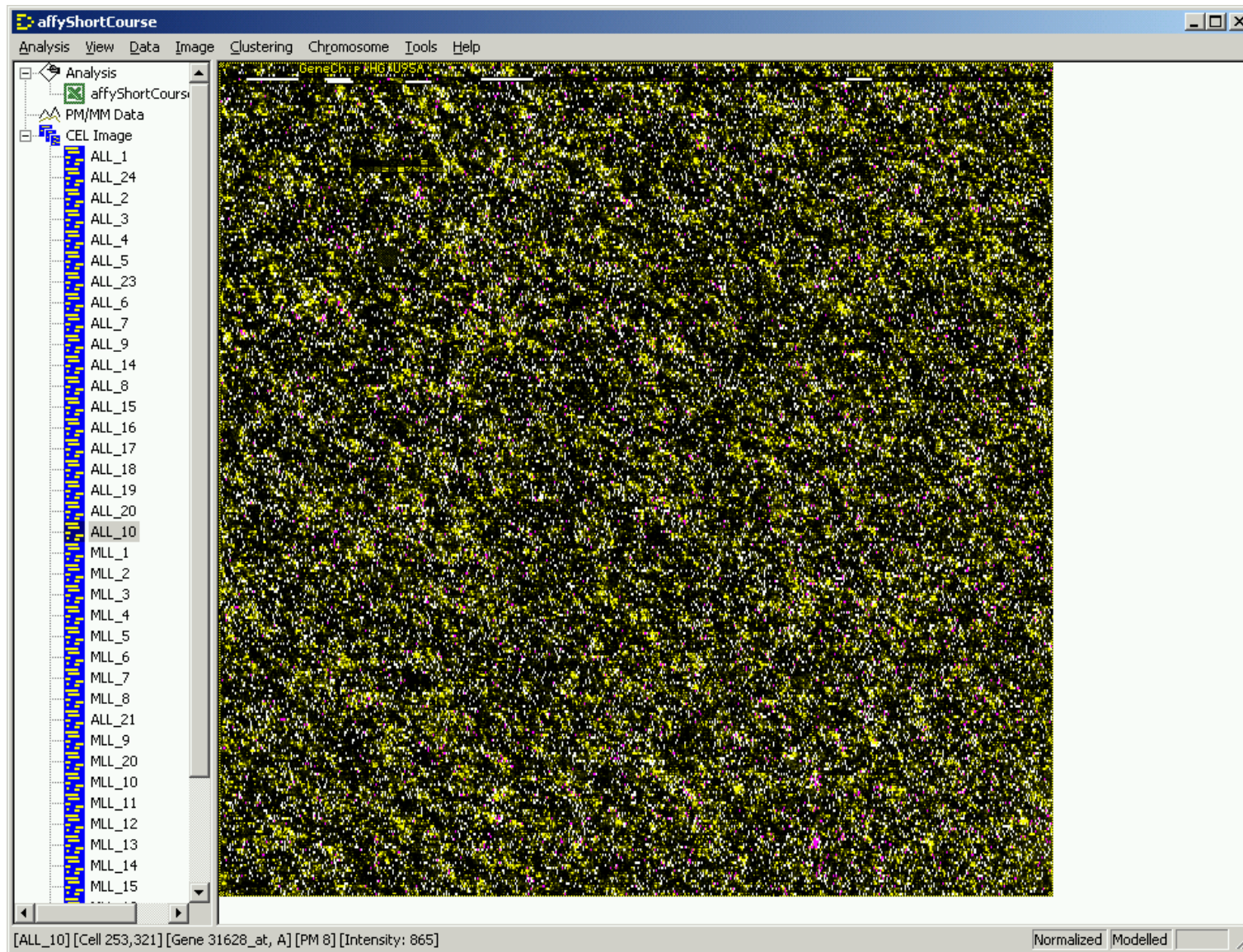


# Look at the Chips, with Cues

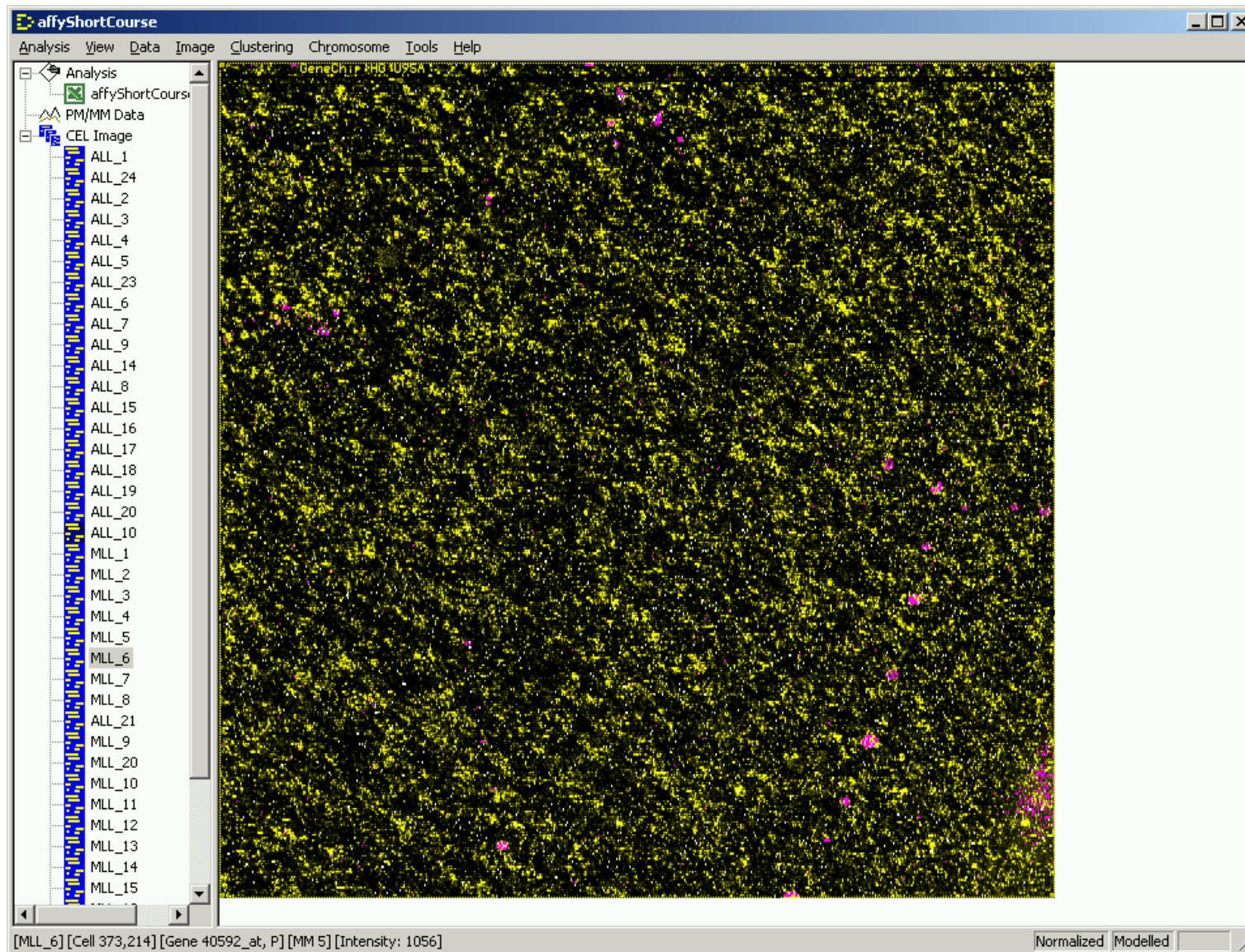




# Look at the Chips, with Cues



# Look at the Chips, with Cues



## Residual Checking is Useful

Hitting the “o” key toggles the display of outliers, which can let us look at the values underneath to see if we can spot what the model is picking up.

The file

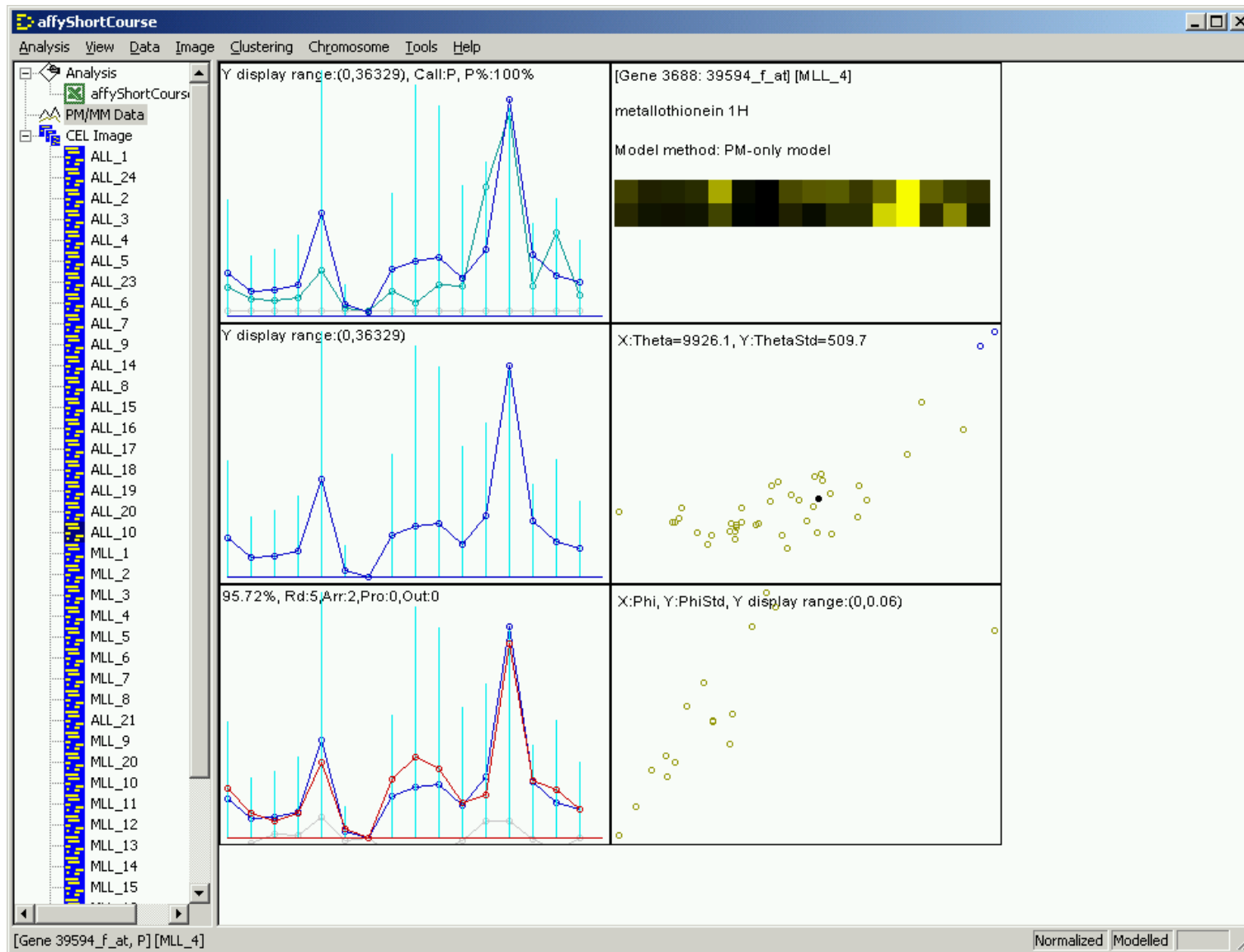
`affyShortCourse.xls`

has been updated in the model-fitting process to record the number of “array outliers” (high standard errors, in white) and “single outliers” (discounted measurements, in purple). Model fitting is performed in a robust fashion.



So, what does a probeset look like?

# Look at a Probeset



## Look at a Probeset

Various panels show

- The PM/MM values for this probeset in this array
- A heatmap view of the same thing
- The target PM-MM values or PM-BG values in this array
- The MBEI values, plotted against their standard errors
- The target values, fitted values, and residuals
- The probe sensitivity indices, plotted against their standard errors

outliers are indicated with colored dots.

## Look at a Probeset

Cycling through the different chips can be accomplished using the Page Up or Page Down keys. The arrow keys zoom in and out as before, but this feature is less useful here.

Holding down the Page Down key produces an animation effect, which can also be achieved using Data/Animate.

The samples are sorted in order of increasing MBEI values, so cycling through produces a differential effect.

For the sample in question, there were 2 array outliers, 0 probe outliers, and 0 single outliers. The model explained 95.72% of the variation, and iterative fitting took 5 rounds.



So, which probesets are “interesting”?

## Find Interesting Genes

Go to Analysis/Compare Samples

Choose the groups using “Select by Category”; this exploits the information that we supplied in the Sample Info file.

One group is “Baseline”, the other “Experiment”

Filter using the lower bound of fold change

Filter on absolute differences

# Find Interesting Genes: Panel 1

**Compare Samples**

Compare samples | Combine comparisons

Baseline (B)

- MLL\_12 [33]
- MLL\_13 [34]
- MLL\_14 [35]
- MLL\_15 [36]
- MLL\_19 [37]
- ALL\_11 [38]
- ALL\_22 [39]
- MLL\_18 [40]
- ALL\_12 [41]

Experiment (E)

- MLL\_10 [31]
- MLL\_11 [32]
- MLL\_12 [33]
- MLL\_13 [34]
- MLL\_14 [35]
- MLL\_15 [36]
- MLL\_19 [37]
- ALL\_11 [38]
- ALL\_22 [39]

Select by category

Select by category

Comparison criteria

(1)   $E / B > 1.2$   or  $B / E > 1.2$   
 Use lower 90% confidence bound of fold change

(2)   $E - B > 100$   or  $B - E > 100$   
For logged data, use (2) instead of (1) for fold

(3)  (P value for testing  $E = B$ )  $\leq 0.05$

(4)  P call of B  $\geq 20\%$  and P call of E  $\geq 20\%$

(5)  (P value for paired t-test)  $\leq 0.05$

[Help](#)

OK Cancel Apply



## **Find Interesting Genes**

Look at “Combine Comparisons”

See where the comparison results will be sent

Estimate FDR using permutations

# Find Interesting Genes: Panel 2

**Compare Samples** [X]

Compare samples | Combine comparisons

Combine type

And  And not  Or  Or not

Combine	Baseline	Experiment	E/...	or B/E>	U...	E-B>	or B
{	1,2,3,4,...	20,21,22,23,...	1,...	1.200	L...	10...	100.
}							

Compare on gene list:

Compare result file

Output all genes

Permute samples to assess False Discovery Rate (FDR):  times

[Help](#)

# Find Interesting Genes – Voila!

affyShortCourse

Analysis View Data Image Clustering Chromosome Tools Help

Analysis

- affyShortCourse
- affyShortCourse
- PM/MM Data
- CEL Image
  - ALL\_1
  - ALL\_24
  - ALL\_2
  - ALL\_3
  - ALL\_4
  - ALL\_5
  - ALL\_23
  - ALL\_6
  - ALL\_7
  - ALL\_9
  - ALL\_14
  - ALL\_8
  - ALL\_15
  - ALL\_16
  - ALL\_17
  - ALL\_18
  - ALL\_19
  - ALL\_20
  - ALL\_10
  - MLL\_1
  - MLL\_2
  - MLL\_3
  - MLL\_4
  - MLL\_5
  - MLL\_6
  - MLL\_7
  - MLL\_8
  - ALL\_21
  - MLL\_9
  - MLL\_20
  - MLL\_10
  - MLL\_11
  - MLL\_12
  - MLL\_13
  - MLL\_14

Permutation 34: [14,21,4,34,24,30,20,10,36,22,9,32,2,33,41,13,15,40,42,28,25,18,23,3 vs. 35,11,27,16,12,7,31,19,37,38,17,1,39,26,29,6,8,5], genes obtained: 24

Permutation 35: [22,25,40,20,8,37,30,5,36,19,35,26,10,9,2,21,42,13,32,16,14,28,31,6 vs. 1,17,39,24,7,18,3,34,12,41,27,15,11,23,33,4,29,38], genes obtained: 14

Permutation 36: [38,9,27,23,34,8,18,22,4,1,2,3,25,13,14,42,16,17,11,6,15,5,24,7 vs. 41,29,12,26,21,33,19,40,39,30,35,10,28,37,36,32,20,31], genes obtained: 85

Permutation 37: [18,31,1,16,24,39,22,21,6,42,29,32,20,27,33,40,23,11,2,8,36,19,38,3 vs. 28,17,15,5,13,35,9,10,30,41,26,4,12,7,37,25,14,34], genes obtained: 14

Permutation 38: [32,14,22,9,8,38,23,18,1,37,40,36,5,30,26,34,31,25,2,20,28,21,3,7 vs. 11,42,39,13,10,19,24,12,16,4,41,6,35,29,33,17,15,27], genes obtained: 47

Permutation 39: [37,20,41,11,34,33,15,2,27,1,39,16,36,35,25,22,42,7,10,17,40,14,18,29 vs. 6,26,23,12,5,24,38,3,28,32,4,30,9,13,31,8,21,19], genes obtained: 2

Permutation 40: [38,37,12,40,35,20,42,27,16,4,22,17,18,8,24,28,10,5,25,39,41,34,32,3 vs. 13,26,29,6,15,9,33,7,31,1,21,11,2,14,30,19,36,23], genes obtained: 8

Permutation 41: [5,14,3,7,41,36,19,26,40,42,35,4,38,2,29,1,6,18,12,10,34,22,15,16 vs. 37,27,21,39,32,9,17,28,25,31,23,20,8,30,33,11,24,13], genes obtained: 14

Permutation 42: [27,6,18,25,15,9,22,7,40,3,11,19,24,36,29,32,33,41,12,23,42,30,37,26 vs. 34,39,1,20,21,35,10,8,5,31,28,13,16,2,14,38,4,17], genes obtained: 57

Permutation 43: [13,2,21,11,5,41,14,7,34,8,20,36,18,19,33,35,10,22,16,12,24,17,29,42 vs. 23,32,40,15,31,38,6,25,3,28,30,26,27,9,39,4,1,37], genes obtained: 15

Permutation 44: [32,4,5,8,42,38,10,26,18,12,2,22,27,40,24,7,41,34,13,30,1,14,29,19 vs. 39,37,3,28,20,31,36,33,16,17,6,21,23,15,35,9,25,11], genes obtained: 2

Permutation 45: [36,14,4,10,27,31,21,13,9,8,17,24,3,32,7,42,37,18,20,41,23,29,30,15 vs. 35,26,38,16,40,25,11,5,6,22,19,33,28,39,34,1,12,2], genes obtained: 1

Permutation 46: [22,7,35,11,8,21,36,12,9,27,23,34,28,3,18,2,40,32,14,6,10,19,26,39 vs. 41,13,1,4,38,25,16,31,24,29,37,17,20,15,5,42,33,30], genes obtained: 7

Permutation 47: [42,13,24,26,28,38,20,12,23,22,7,14,29,27,33,34,6,41,9,19,1,16,30,21 vs. 15,32,40,35,5,3,37,11,25,8,2,4,10,36,17,39,18,31], genes obtained: 9

Permutation 48: [24,2,3,7,13,42,29,23,35,12,19,40,10,37,16,22,33,25,31,11,36,41,21,38 vs. 30,34,9,32,4,20,15,17,14,1,39,18,5,28,27,8,6,26], genes obtained: 5

Permutation 49: [14,3,34,5,30,6,40,16,29,31,17,39,12,26,8,37,4,1,33,2,38,22,24,9 vs. 15,28,19,21,10,42,20,13,27,36,25,32,41,11,7,35,18,23], genes obtained: 45

Permutation 50: [14,19,42,16,2,35,31,26,36,21,11,13,7,1,28,38,32,9,10,37,39,4,29,22 vs. 18,24,34,15,3,20,17,8,30,40,6,25,41,23,12,33,27,5], genes obtained: 18

Obtained 610 genes; False Discovery Rate (Number) of 50 permutations, Median: 2.3% (14), 90th percentile: 11.5% (70)

Finished)

Analysis outputs

Normalized Modelled

# Find Interesting Genes

Results are exported to

affyShortCourse compare result.xls

```
[ COMPARE_CRITERIA_V2 ]  
$NUM_OPTION_LINE=5  
$ARRAY_LIST_FILE=  
$COMPARE_ON_GENE_LIST=  
$COMPARE_ON_USE_LIST=1  
$AVERAGE_USING_STANDARD_ERROR=Yes  
$OMIT_AFFY_CONTROL_GENE=Yes  
$NUM_CRITERION=1
```

## More compare result.xls (1)

```

$Parenthesis : Combine : Baseline : Experiment :
  E/B> : or B/E> : Use : E-B> : or B-E>
  P value <= : P call % of B >= :
  and P call % of E >= : % Pair P value <=
No : and : 1,2,3,4,5,6,7,8,9,10,11,12,13,14,15,
  16,17,18,19,28,38,39,41,42 :
  20,21,22,23,24,25,26,27,29,30,31,32,33,34,35,
  36,37,40 :
  1.200 : 1.200 : Lower Bound : 100.000 : 100.000
NA : NA : NA : NA

```

## More compare result.xls (2)

[COMPARE\_RESULT]

```
probe set : gene : Accession : LocusLink
Description : ALL_1 ALL_24 ALL_2 ... :
baseline mean : baseline mean's SE :
MLL_1 MLL_2 MLL_3 MLL_4 ... :
experiment mean : experiment mean's SE :
fold change : lower bound of FC : upper bound
of FC : difference of means : filtered
```

## More compare result.xls (3)

```
31407_at : protease, serine, 7 (enterokinase) :  
: U09860 : 5651 :  
Cluster Incl. U09860:Human enterokinase mRNA,  
complete cds /cds=(40,3099) /gb=U09860 /gi=  
746412 /ug=Hs.158333 /len=3696 :  
950.09 155.37 285.85 76.51 412.50 ... :  
267.28 64.25 :  
105.22 67.40 151.85 111.09 106.78 ... :  
125.50 8.56 :  
-2.13 : -1.28 : -3.04 : -141.78 : *
```

# Find This Gene

in Probeset View, use View/Find Gene

**Find probe set or gene** [X]

Change one of the following to search:

Probe set number:

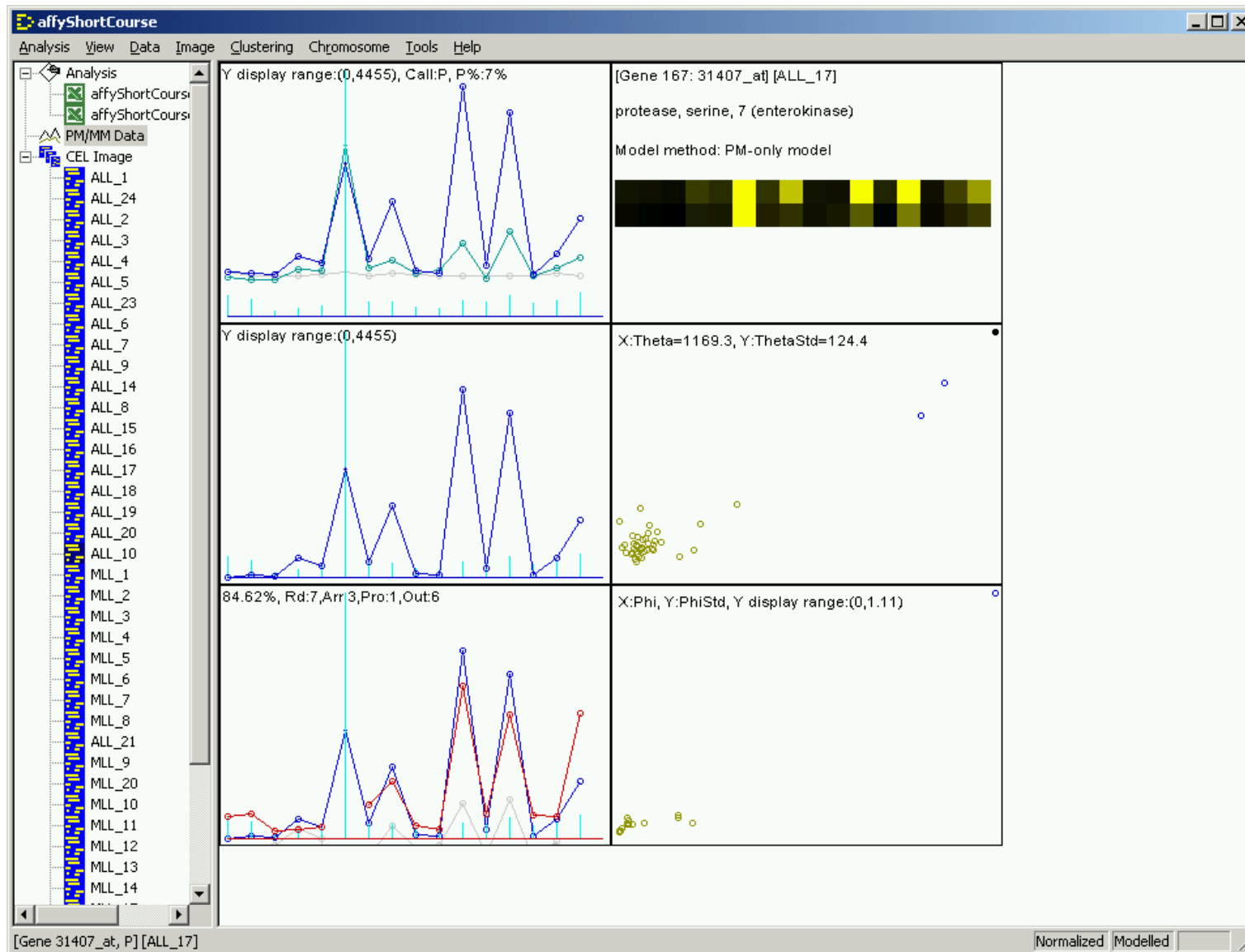
Probe set name:

Gene keywords:

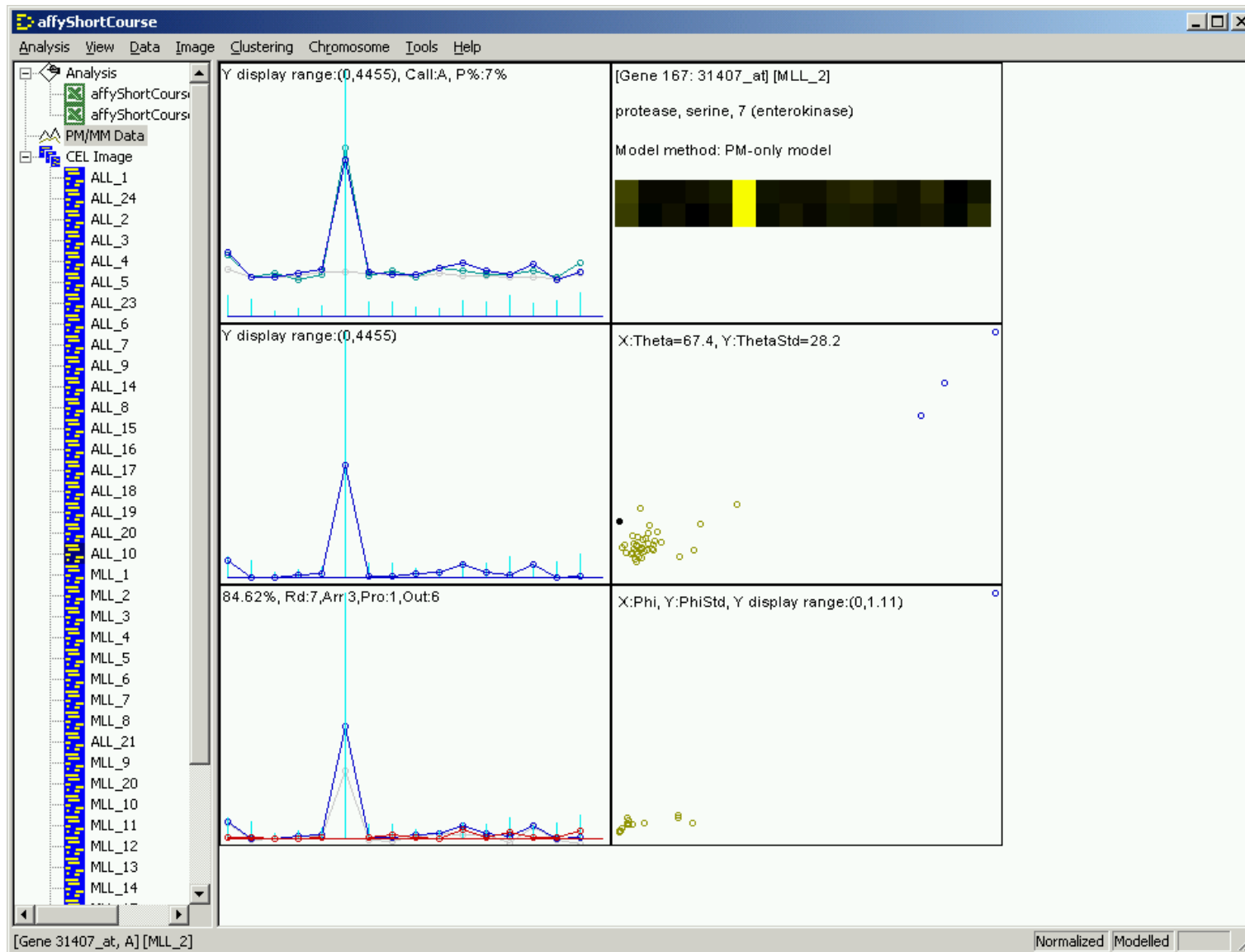
(Can use wildcards, e.g. "receptor \* kinase" matches with "receptor tyrosine kinase", and both "receptor [1-9]" and "receptor ?" match with "receptor 4" )



# Find This Gene: ALL\_17, High End

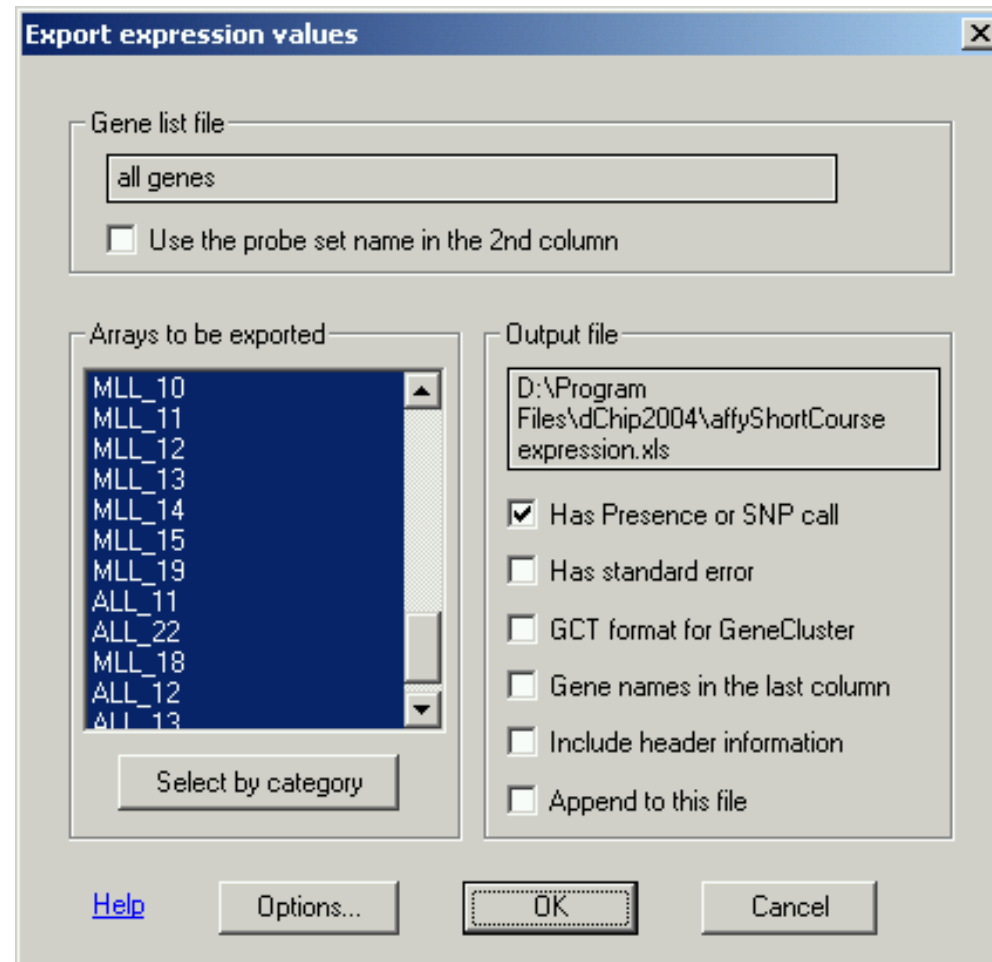


# Find This Gene: MLL\_17, Low End



# Other Exports: Expression Results

Tools/Export Expression Value...



turn off Presence call option

## Export all Expression Results (2)

produces affyShortCourse expression.xls

probe set	gene	Accession			LocusLink	
Description		ALL_1	ALL_24		ALL_2	
ALL_3	ALL_4	ALL_5	ALL_23	ALL_6	ALL_7	
...						
AFFX-MurIL2_at	M16762	M16762	Mouse interleukin 2 (IL-2) ge			
M16762		M16762	Mouse interleukin 2 (IL-2) ge			
1324.22	1766.49	1562.23	1739.9	1486.82		
1624.63	1759.31	1763.18	1558.21	1555.06		
...						
AFFX-MurIL10_at	interleukin 10	M37897	16153			
M37897	Mouse interleukin 10 mRNA, complete cds					
917.868	1360.26	1067.69	1380.64	1037.5	1074.34	
1294.49	1109.37	1181.09	1090.53	1121.5		

## Other Exports: Probe Results

Tools/Export Probe Set...

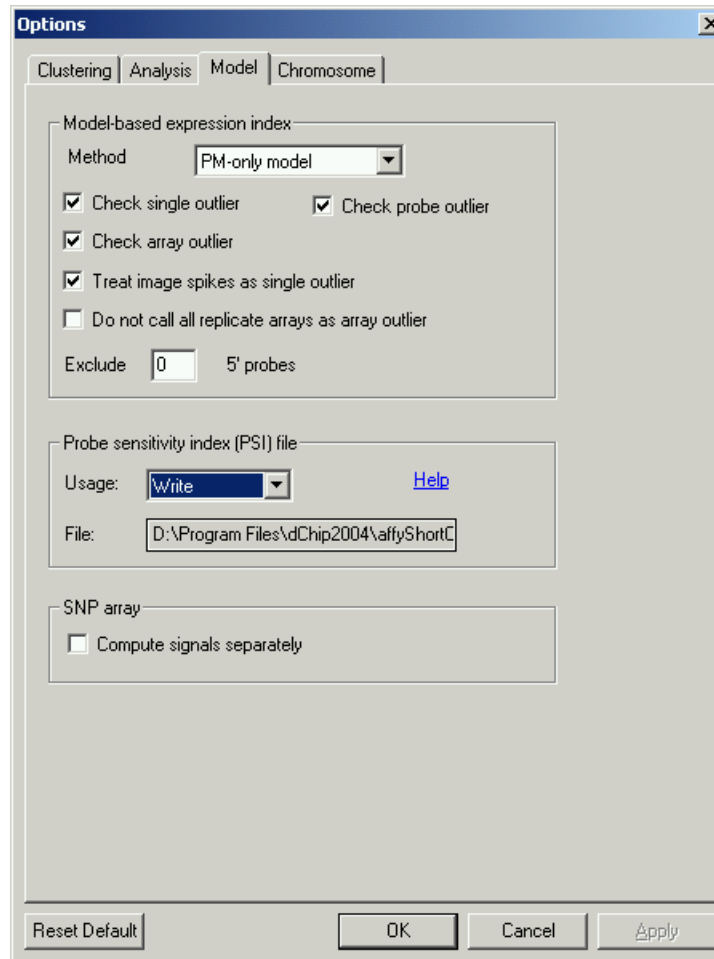
produces

affyShortCourse 31407\_at probe data.xls

Probeset	Probe	Array	PM-bkgd	MM-bkgd
MBEI	PSI			
31407_at	0	0	121	-59
950.092	0.339297			
31407_at	0	1	154	-35
155.365	0.339297			
31407_at	0	2	83	-44
285.847	0.339297			
31407_at	0	3	-96	-73
76.511	0.339297			

# Other Exports: PSIs

Keep the PSIs? Analysis/Model-based Expression, Options,  
Usage: Write



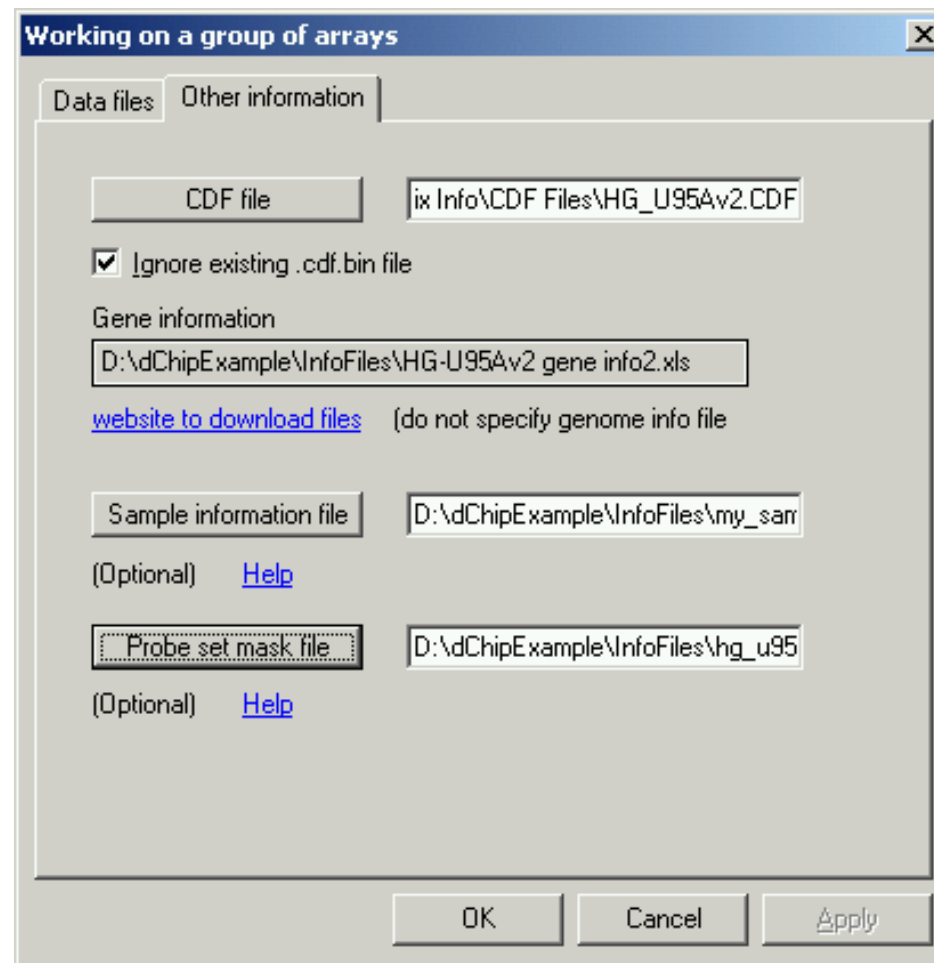
## So, Did We Find What They Did?

Well...■

It turns out that half of the chips used were U95A, and the rest (including all of the AML samples) were U95Av2. By default, dChip does not combine results from different chip types.

However, since the difference is not large (25 probesets out of 12625), we can mask the ones that don't overlap and get it to fit anyway.

# Combine the Chip Types



the mask file is from the dChip web site, and we use the U95Av2 CDF file.



# Do We Find What They Did Now?

Well...■

It turns out that the paper reported gene names and gene symbols, but did not specify the Affymetrix probe ids. Unfortunately, some of the annotation has changed over time.

If we look for

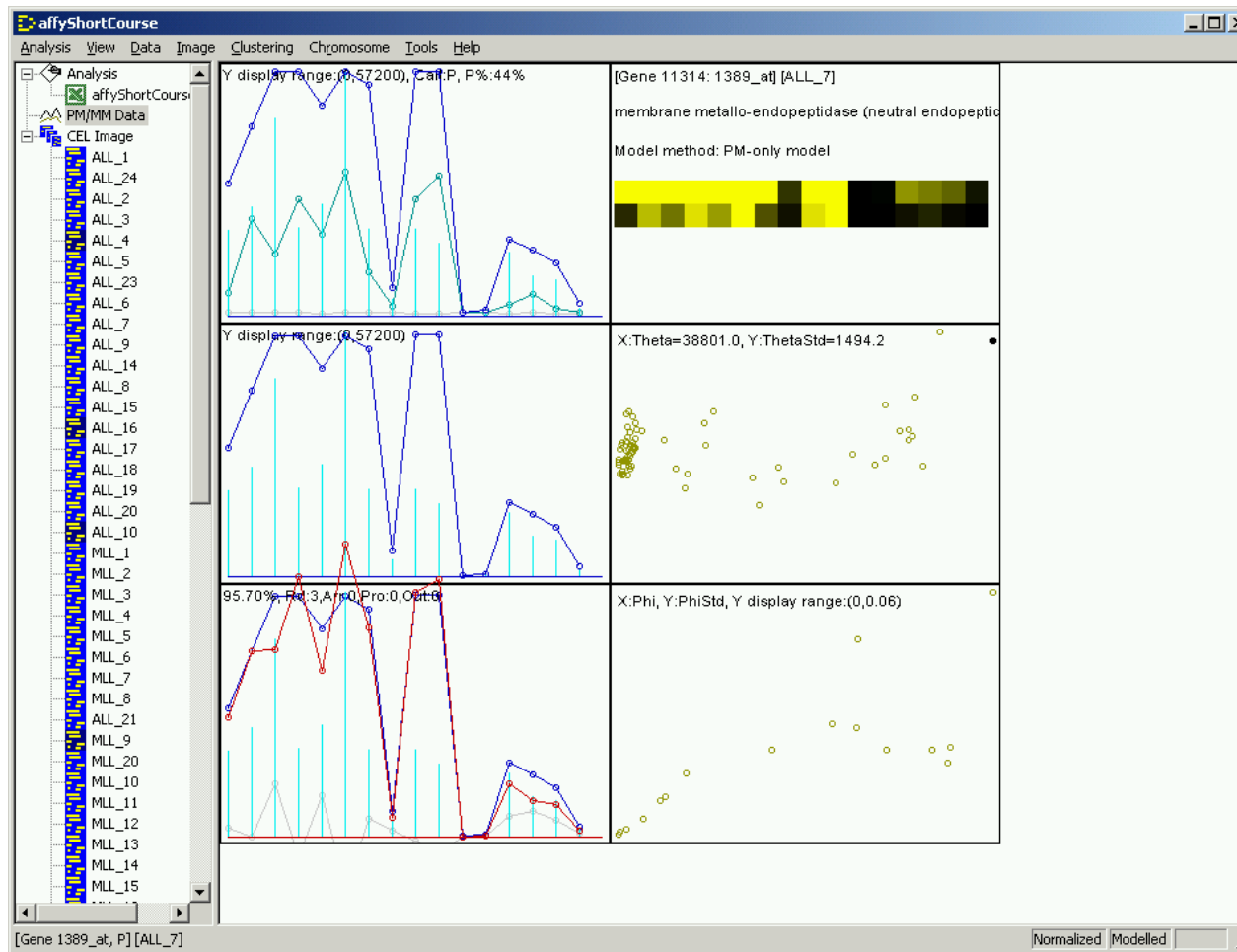
J03779 (gene accession number), aka  
CD10 (gene symbol)

in the expression tables supplied with the paper, it's not there.

■ But if we look in the gene info files supplied with dChip, it *is* there (it's 1389\_at).

# And?

FC: -3.97, CI: (-3.38,-4.6), Diff: -17310. Different!



# Summary

We know what files to track down

We know how to load them in for processing

We know how to normalize and fit models

We know how to export results

We've seen how finicky indexing can be.



And we struck biology!

Thus endeth the lesson...