

Quality Control & Pre-processing Evaluation  
of  
E-MEXP-1956.raw.1\_0  
**REPORT**

Array names and grouping

ArrayDataFile	SourceName	FactorValue
08-F2C14.CEL	Array1	Group1
12-F3C18.CEL	Array2	Group1
11-F3C15.CEL	Array3	Group1
07-F2C03.CEL	Array4	Group1
10-F6C04.CEL	Array5	Group1
09-F6C01.CEL	Array6	Group1

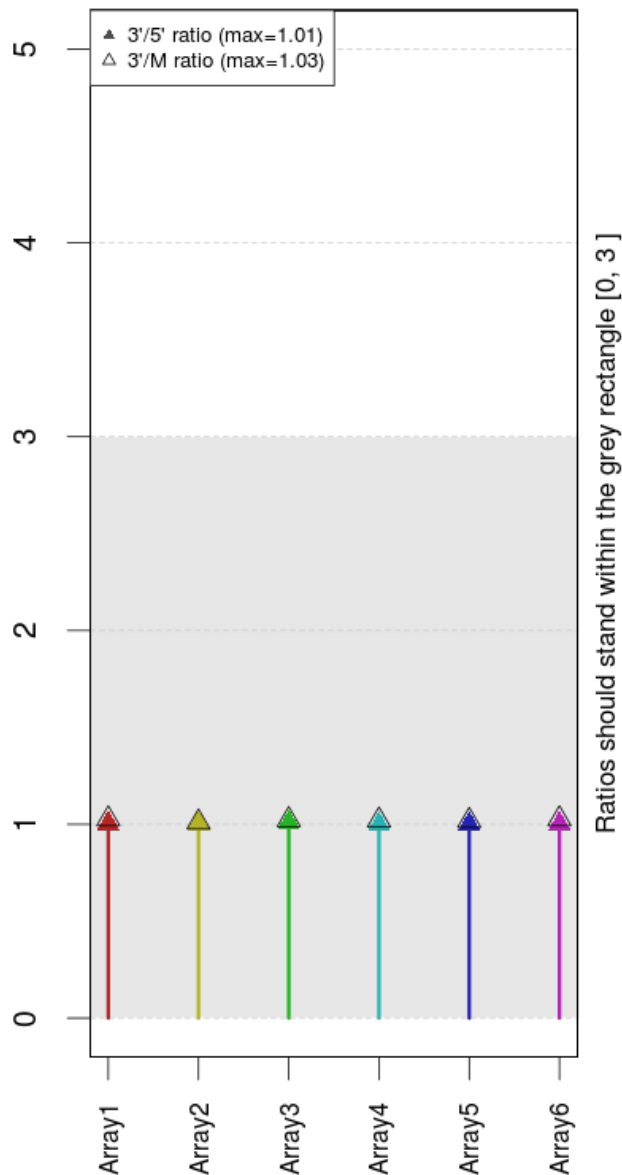
# Summary of raw data quality indicators

blue = "within" / red = "out of" recommended cut-off

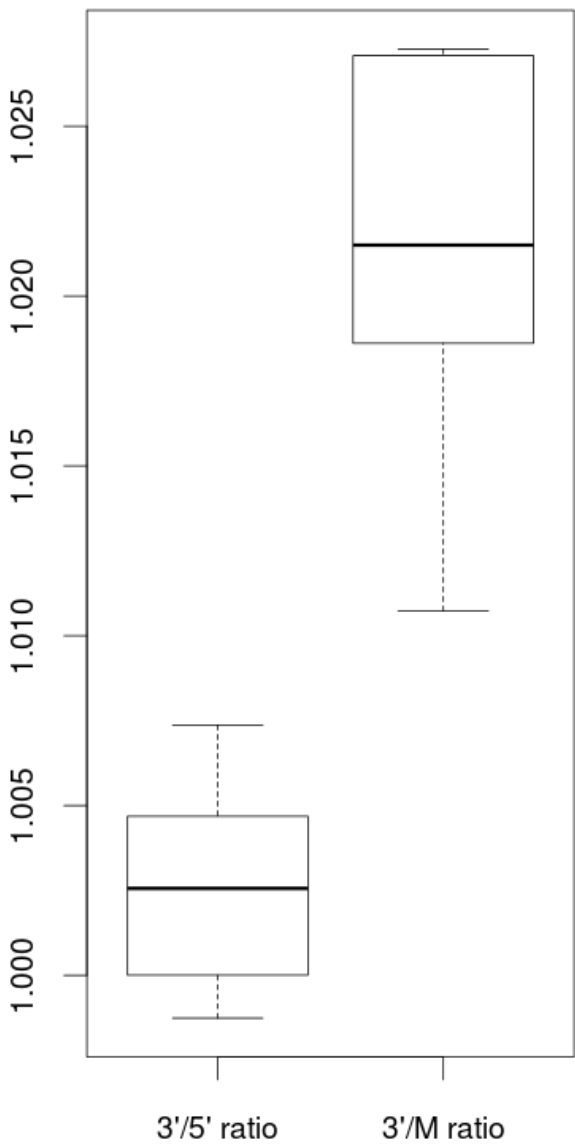
	3'/5' beta-actin (cutoff=3)	3'/5' GAPDH (cutoff= 1.25)	Hybridization BioB<BioC<BioD<CreX	Hybridization BioB=Present	Percent Present spread<= 10%	Background spread<=20%	Log Scale Factor spread<=3
Array1	1	1.02	T	P	41 %	34	0.81
Array2	1	1.02	T	P	39 %	34	1
Array3	1.01	1.02	T	P	36 %	35	1.45
Array4	1	1.02	T	P	39 %	39	1.18
Array5	1	1.02	T	P	38 %	39	1.07
Array6	1	1.02	T	P	39 %	34	1

# RNA degradation of beta-actin

3'/5' and 3'/M ratios



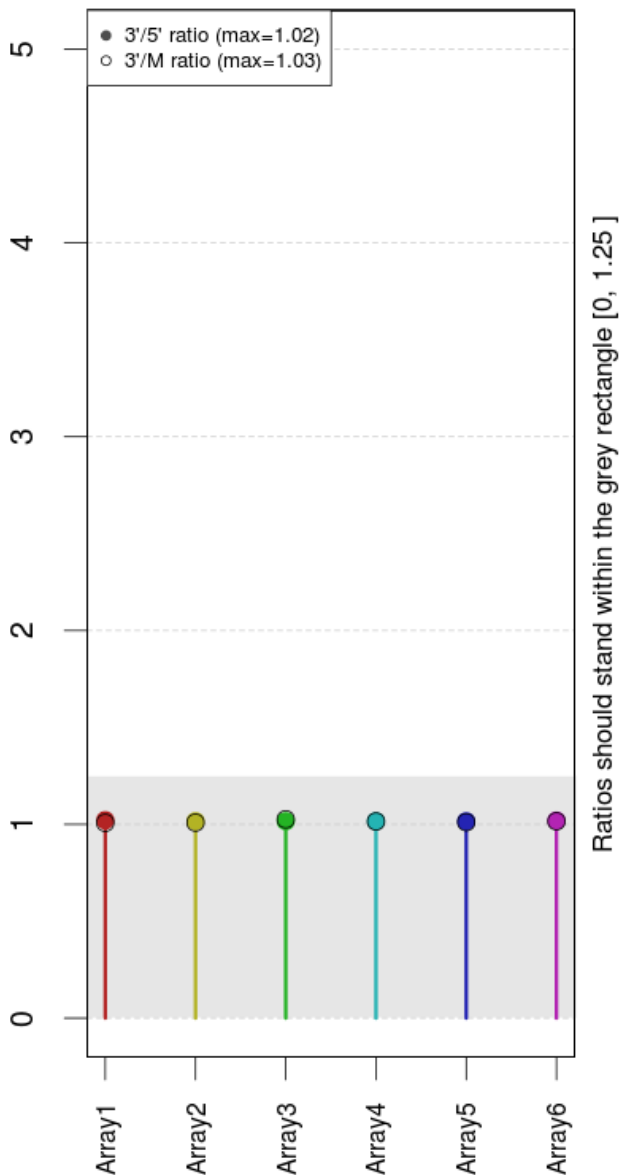
# Boxplot of beta-actin ratios



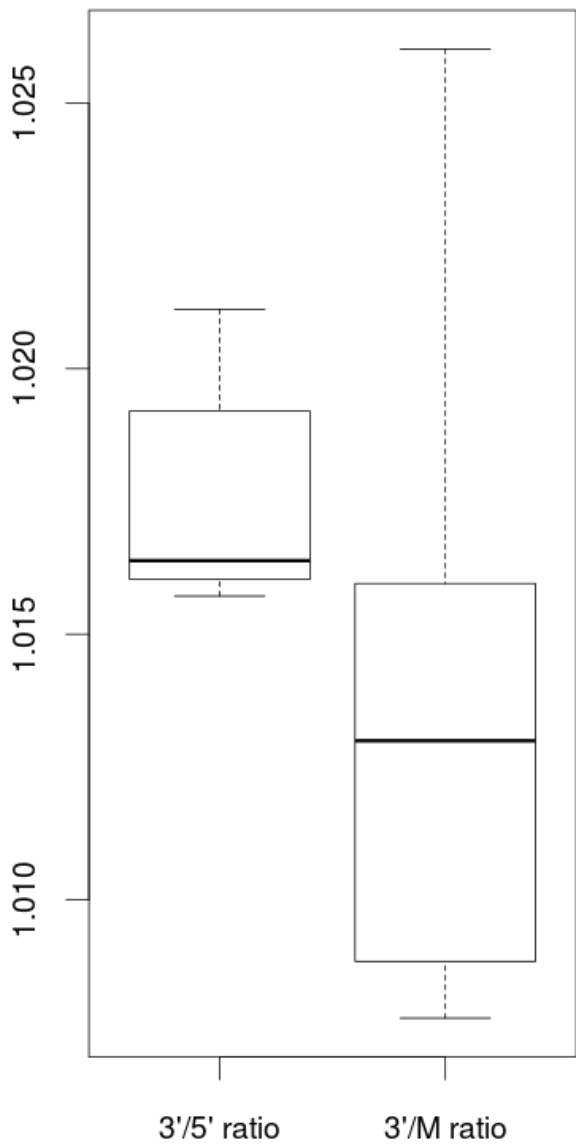
beta-actin QC: OK (all 3'/5' ratios < 3)

# RNA degradation of GAPDH

3'/5' and 3'/M ratios

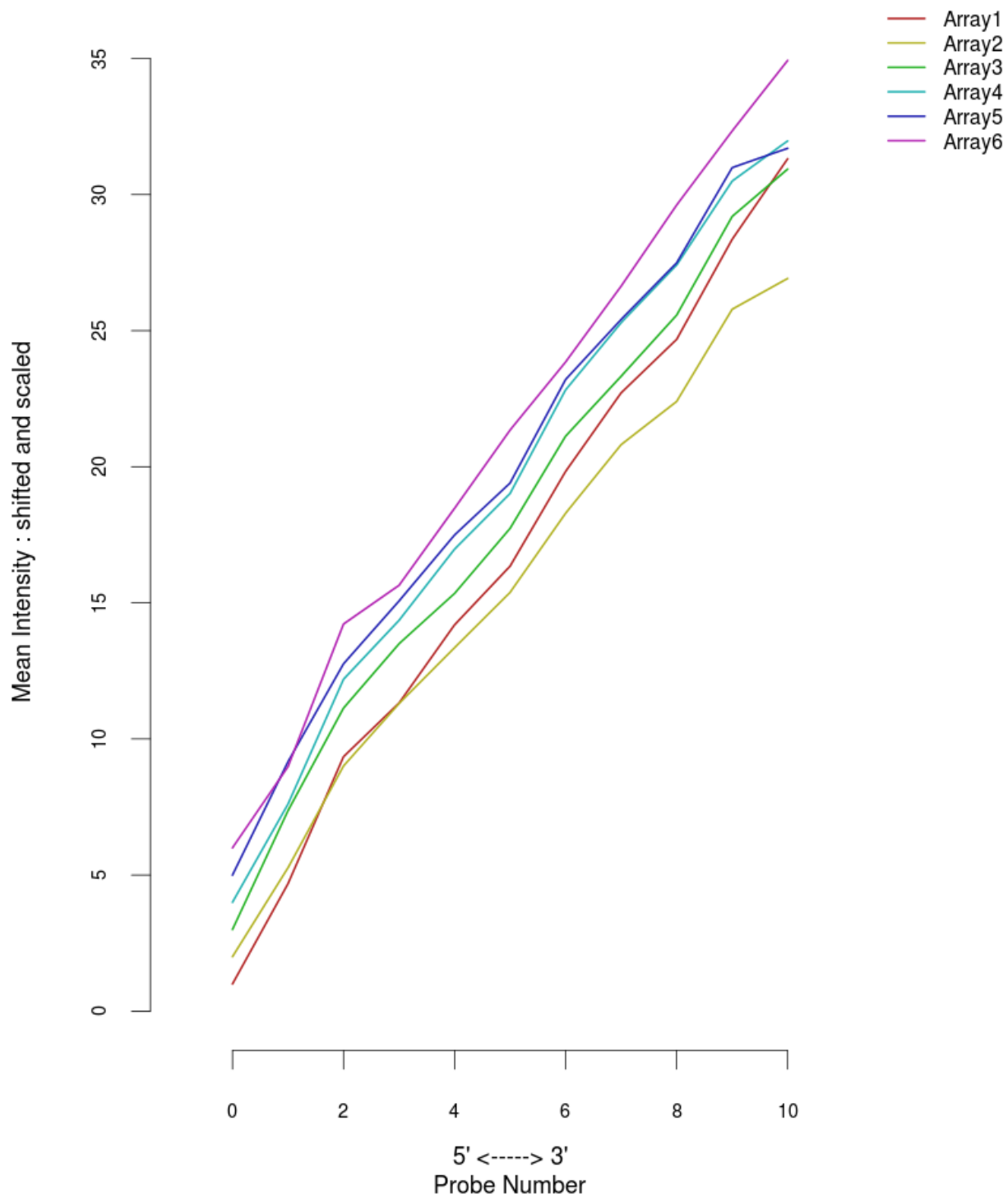


# Boxplot of GAPDH ratios

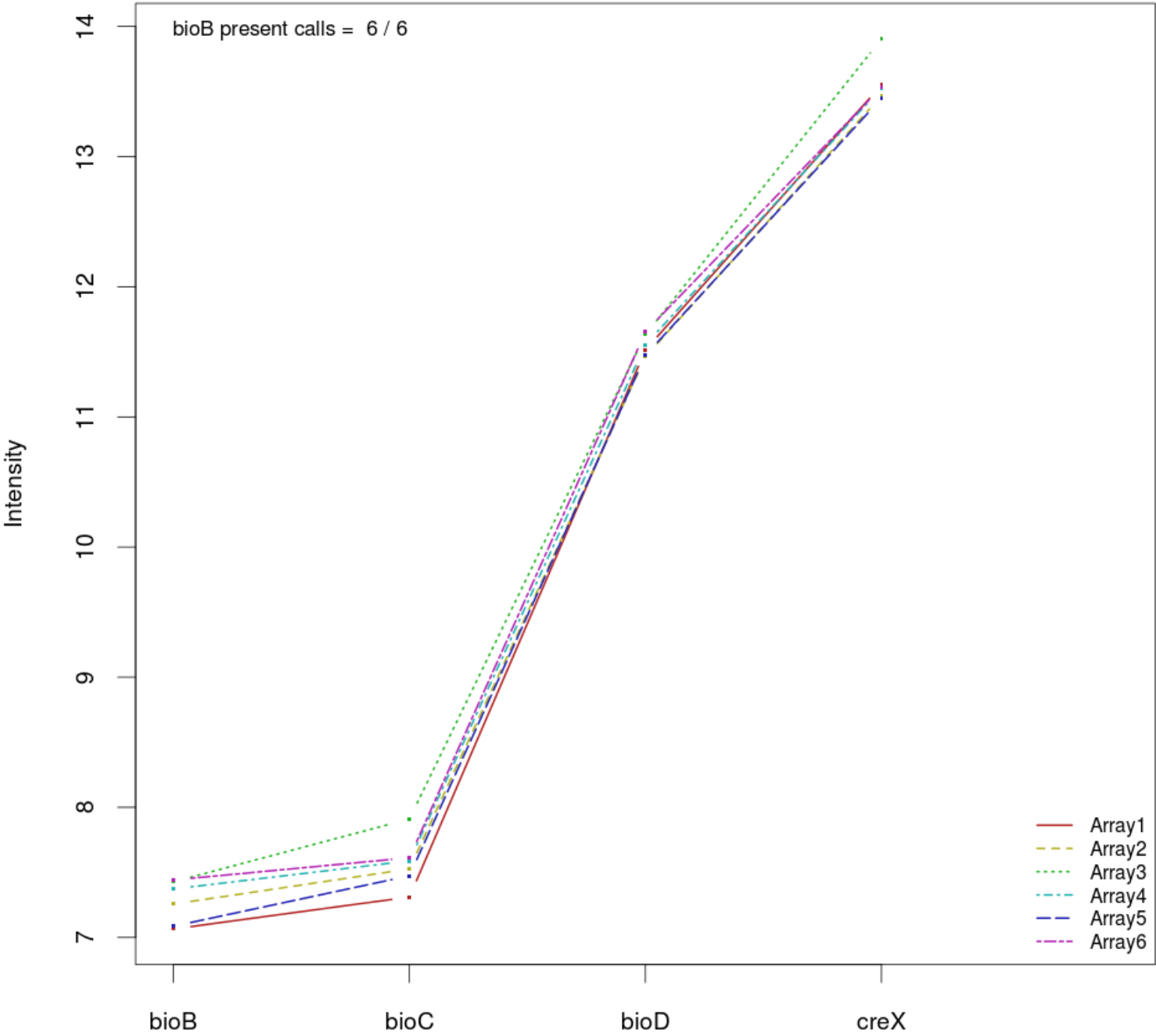


GAPDH QC: OK (all 3'/5' ratios < 1.25)

# RNA degradation plot



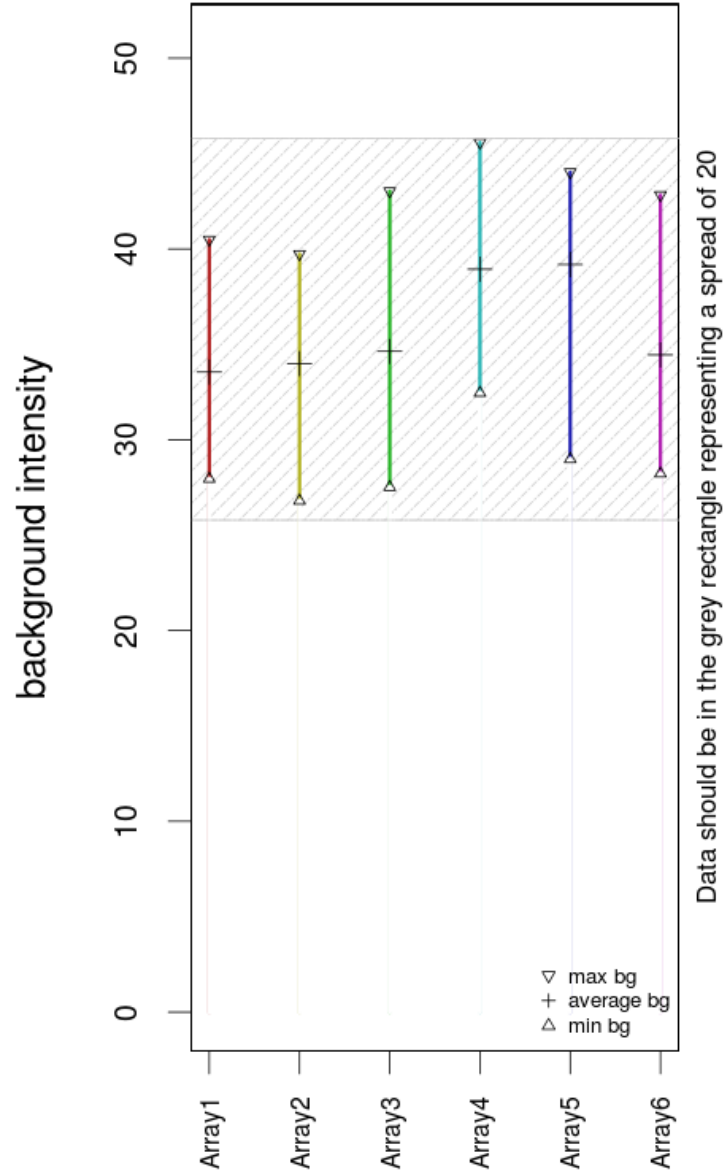
# Spike-in Hybridization controls intensities and calls



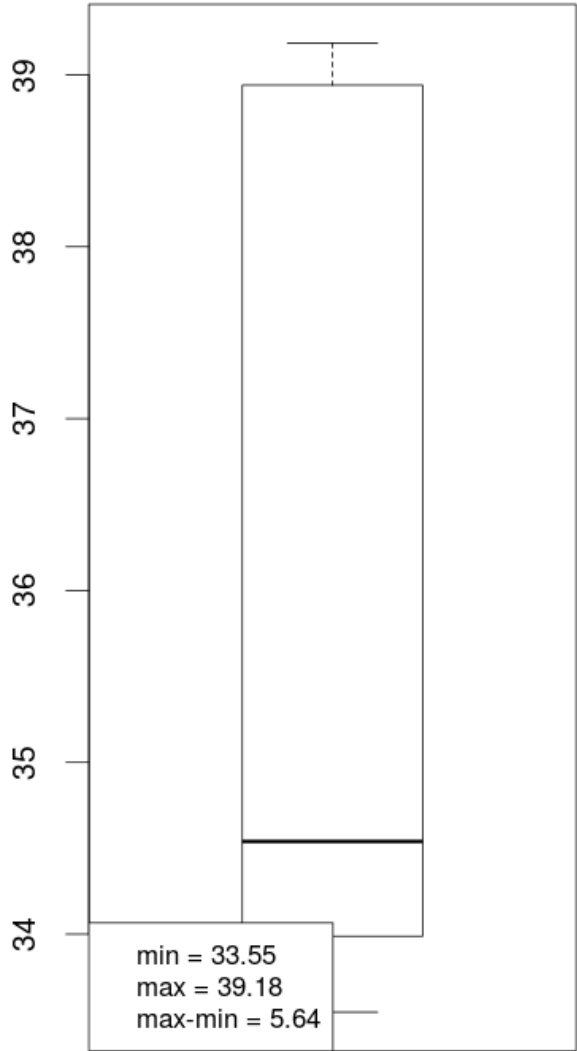
Intensities: OK (bioB < bioC < bioD < creX for all arrays)

BioB Present calls: OK (indeed all bioB are called present)

Plot of background intensity



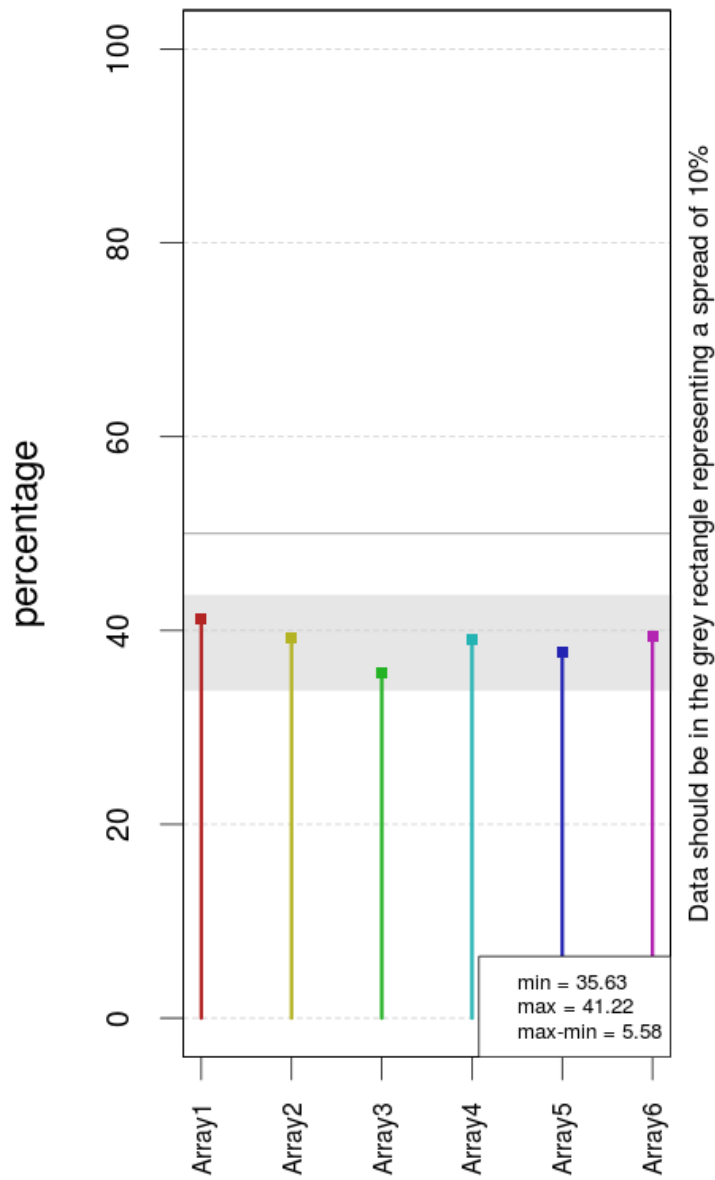
Average background intensity



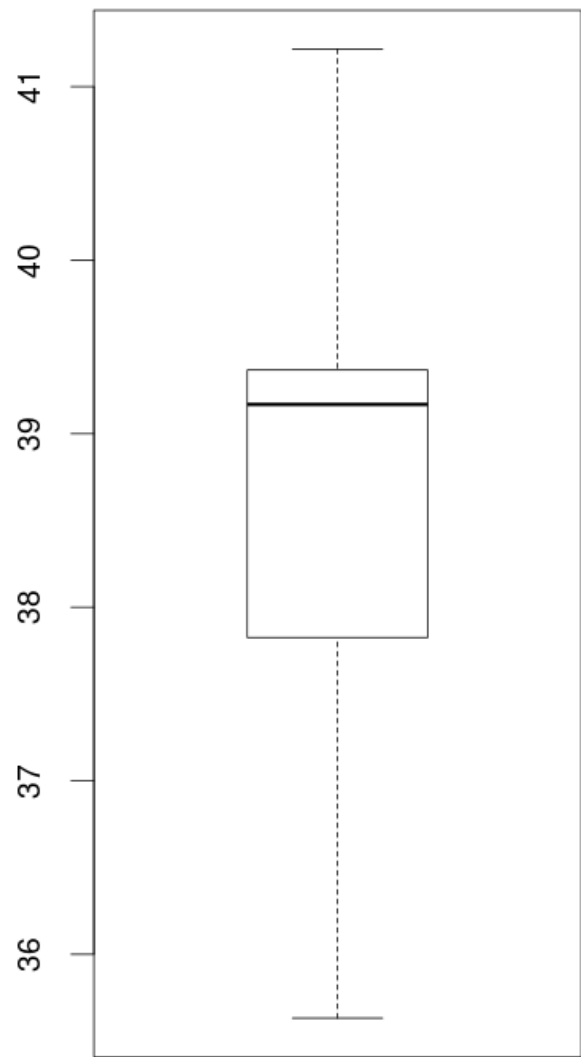
Background QC: OK (spread <= 20)



Plot of percent present

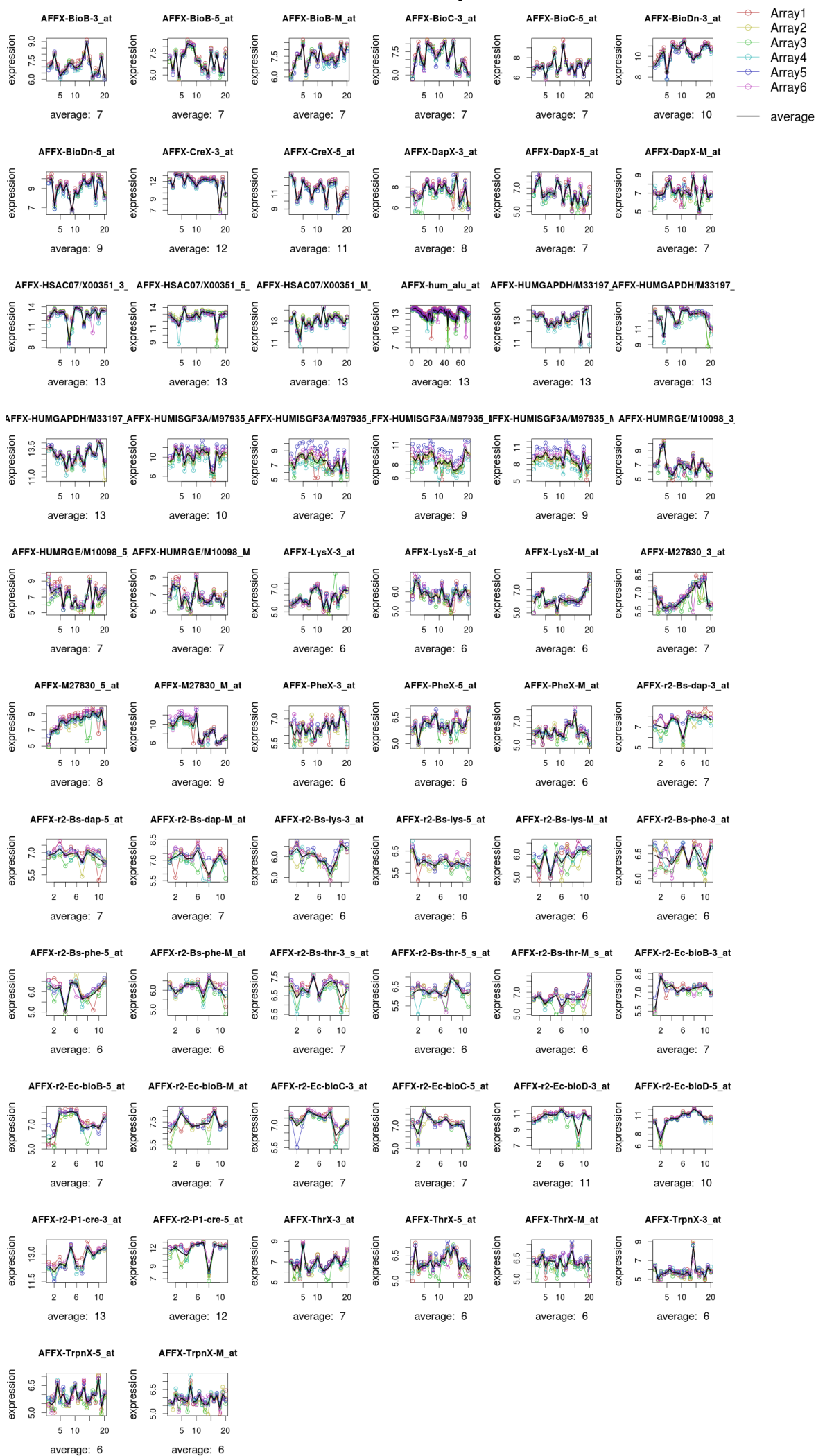


Boxplot of percent present

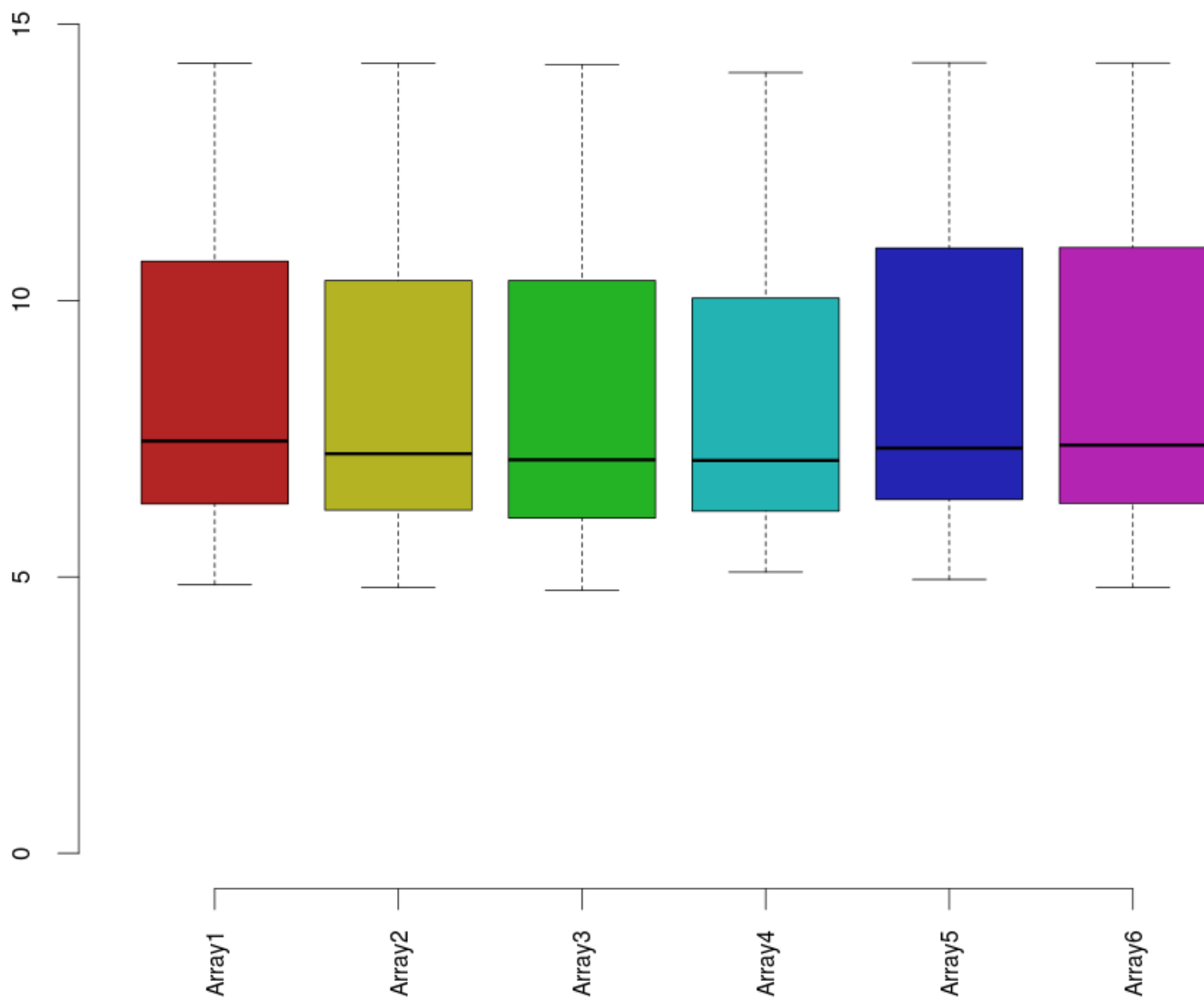


Percent present QC: OK (spread <= 10%)

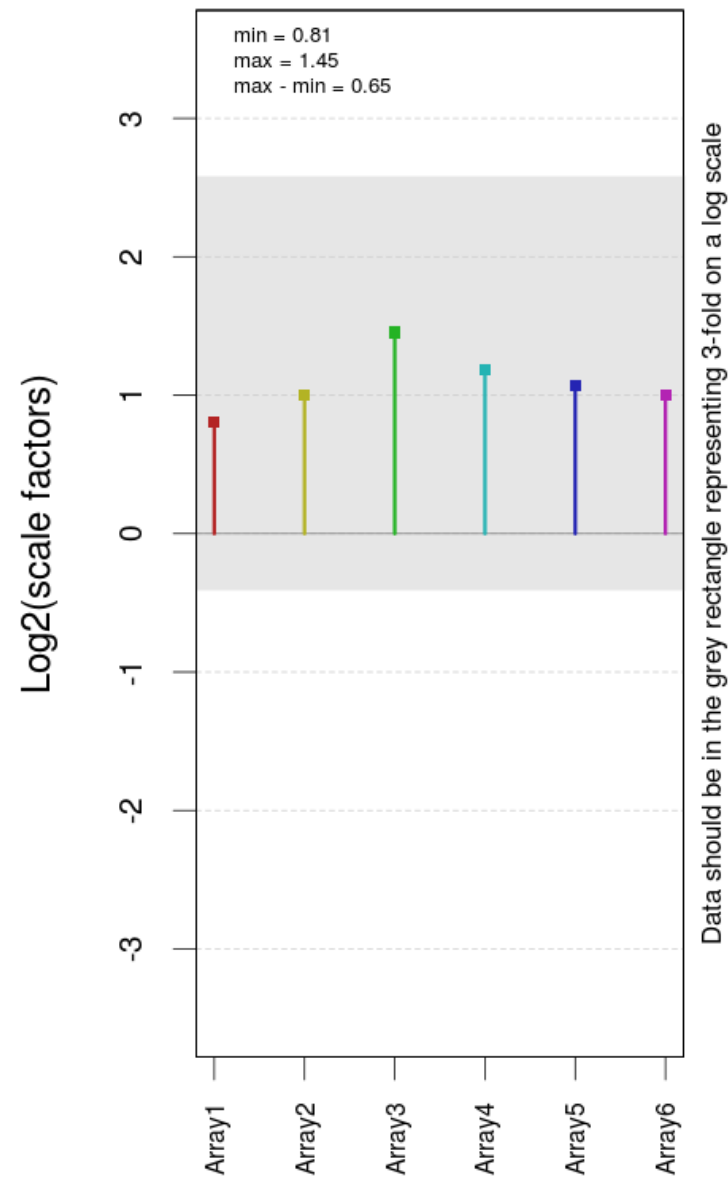
# affx control profiles



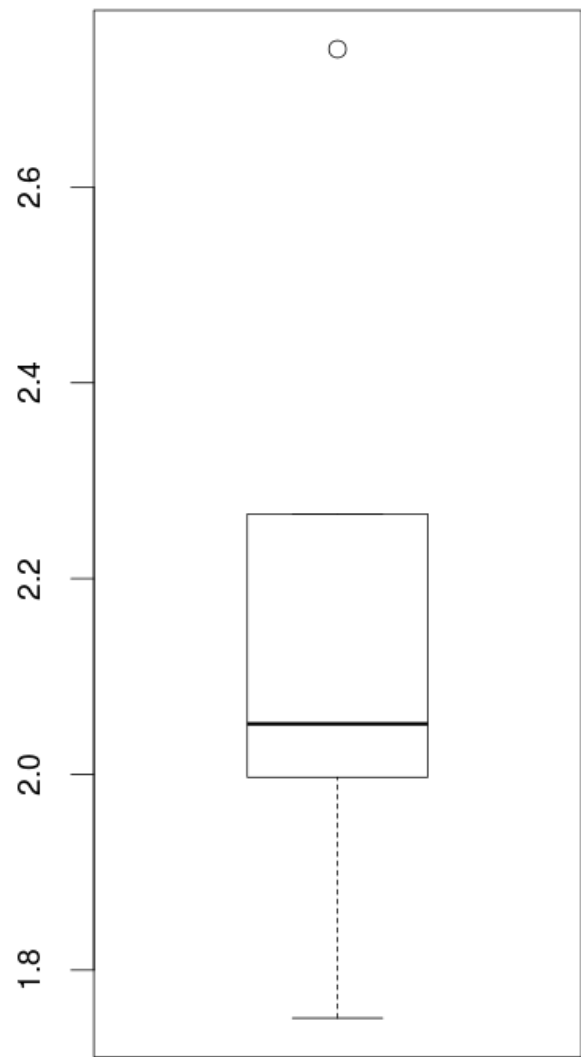
# affx controls



Plot of Log scale factors



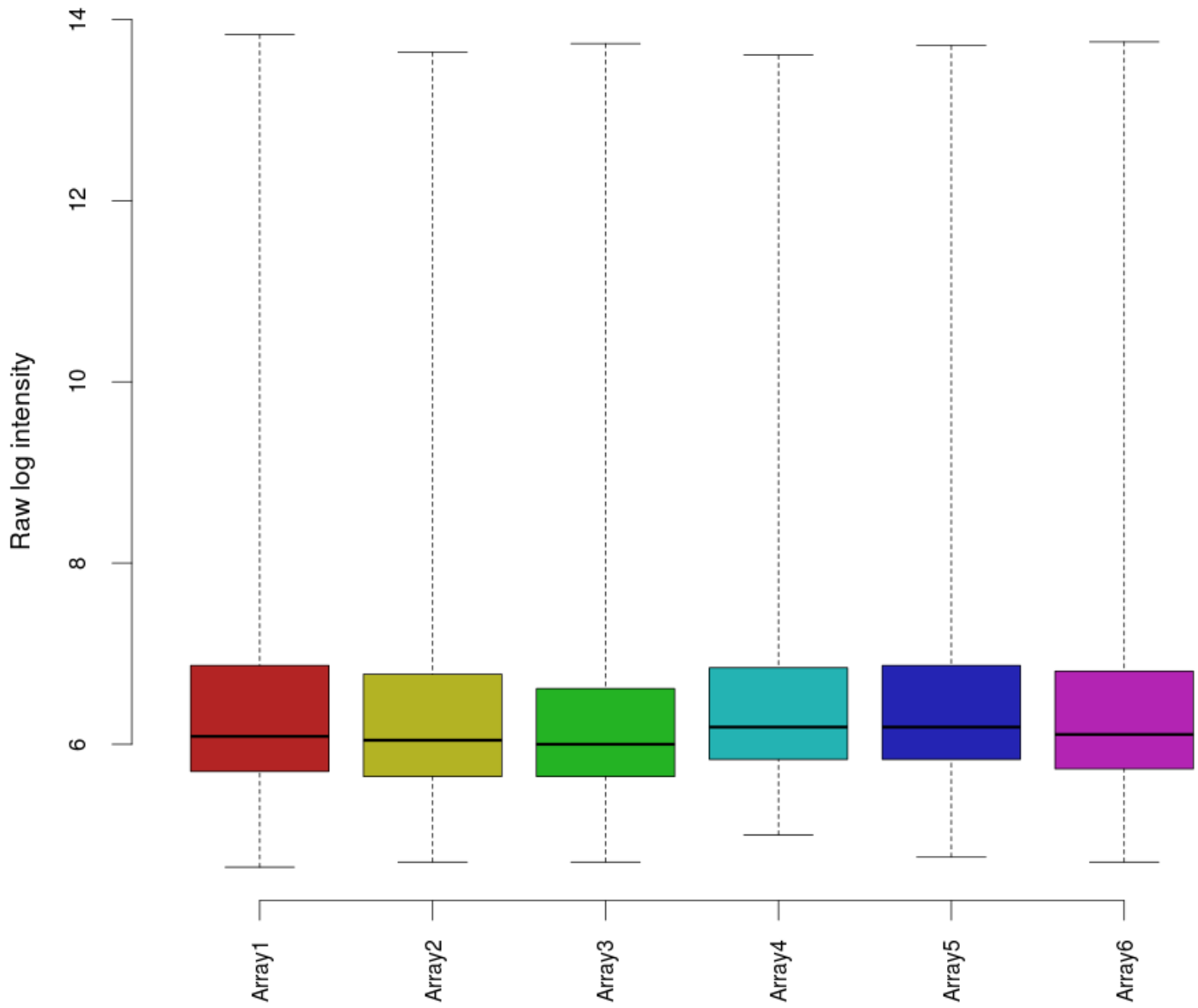
Boxplot of scale factors  
(natural scale)



Scale factors QC: OK (spread < 3-fold)

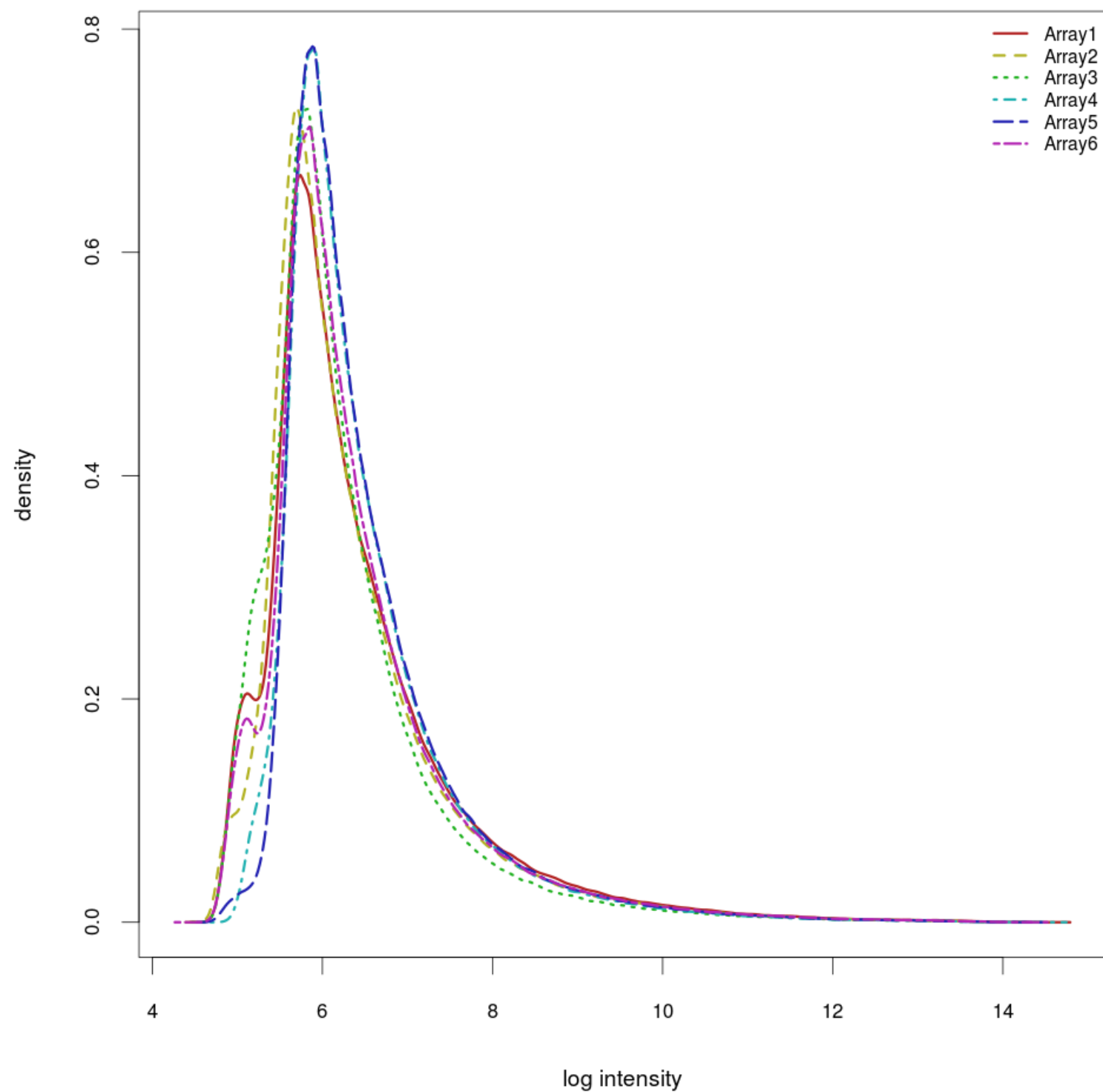
# Boxplot of raw intensities

Distributions should be comparable between arrays



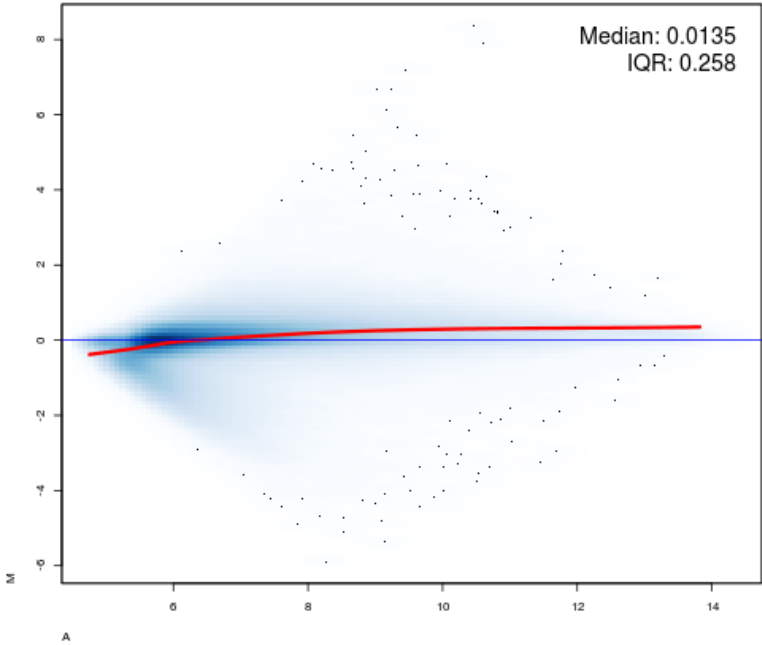
# Density histogram of raw intensities

Curves should be comparable between arrays

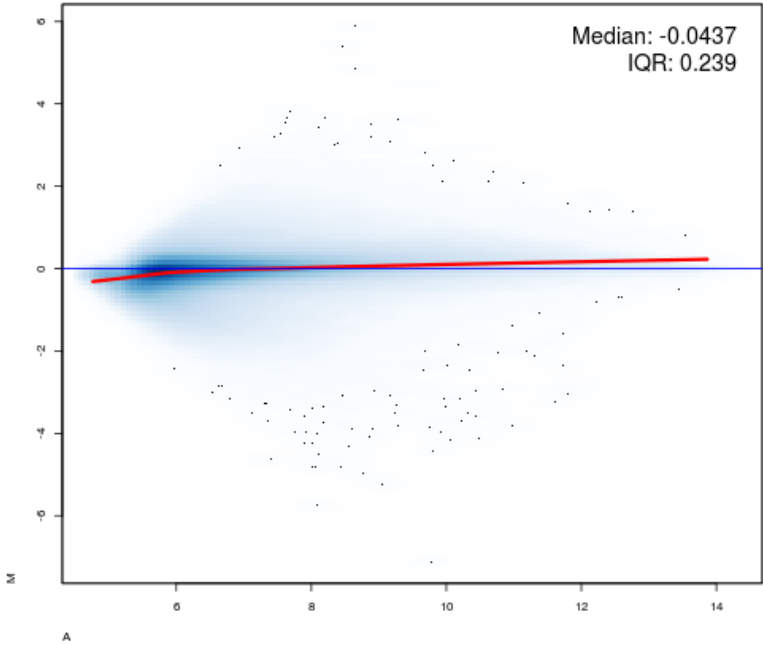


# MA plots of raw data

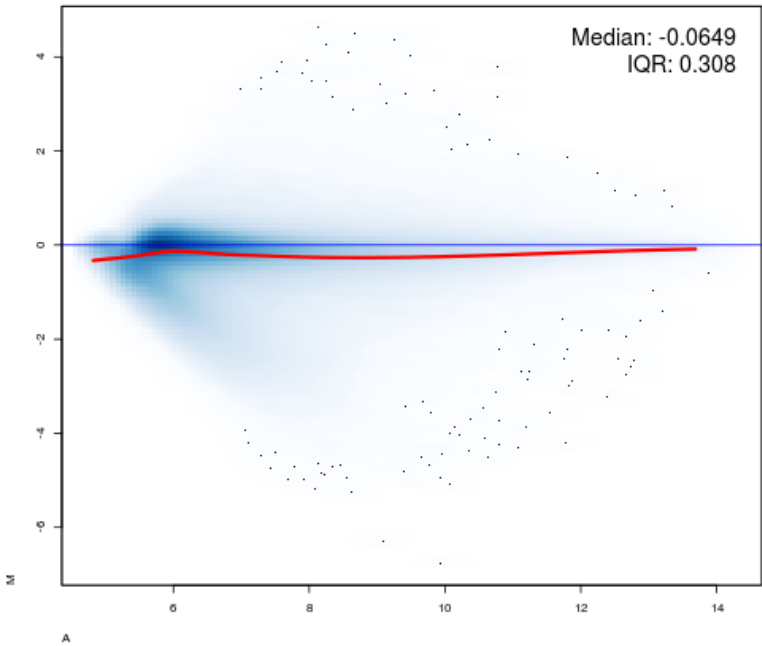
Array1 vs pseudo-median reference chip



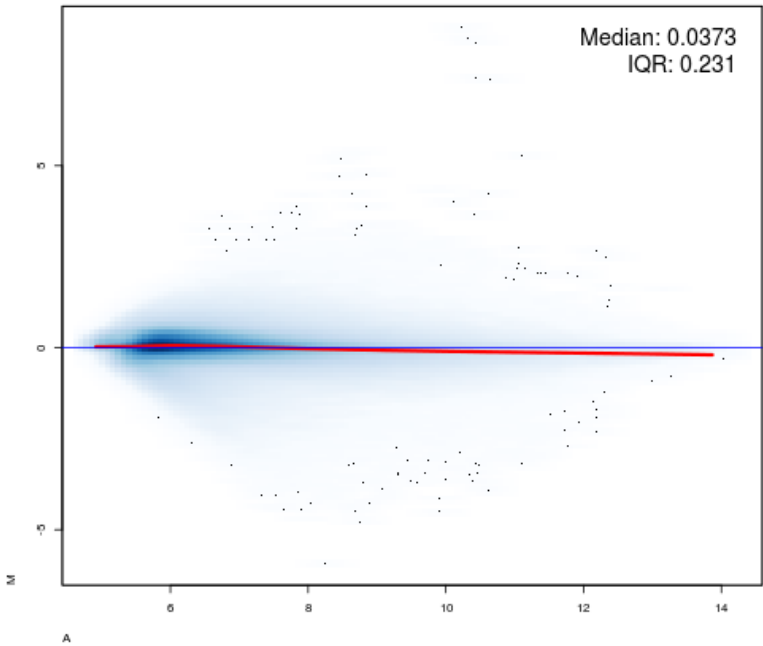
Array2 vs pseudo-median reference chip



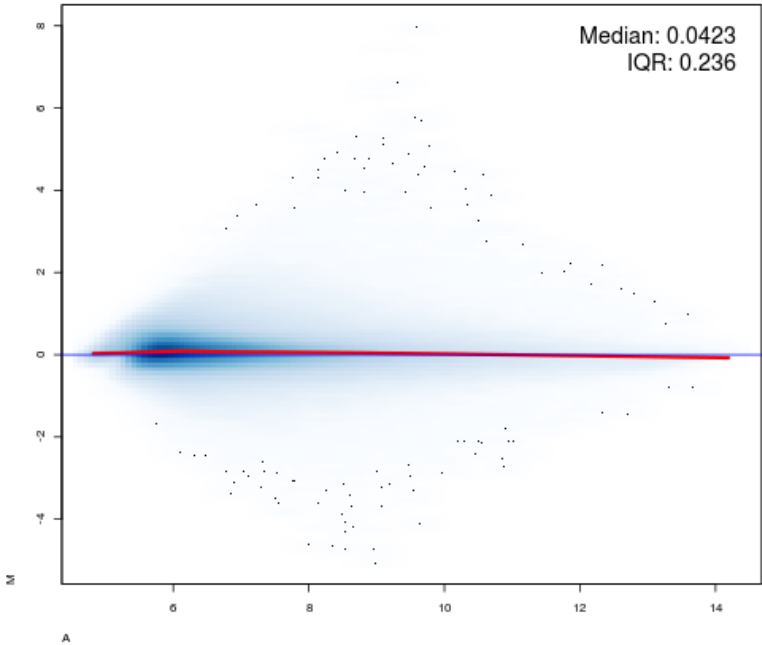
Array3 vs pseudo-median reference chip



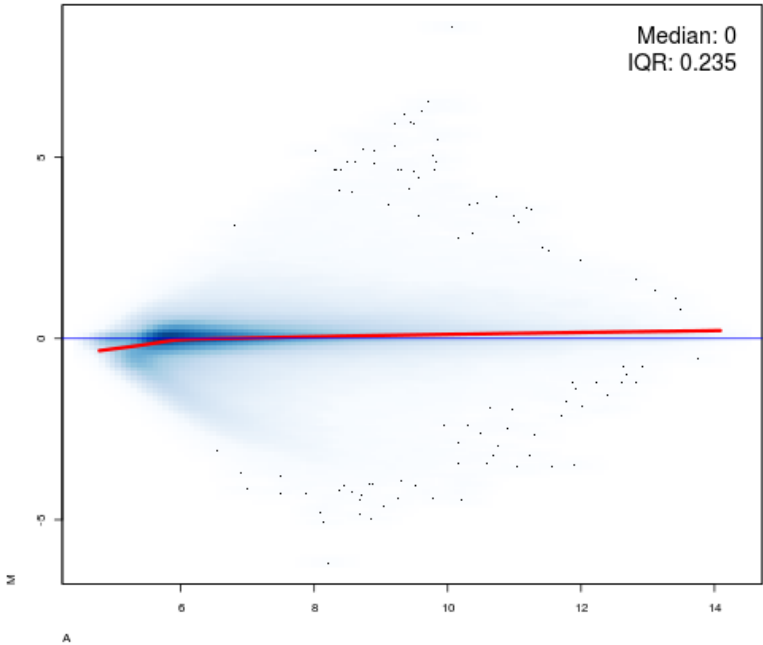
Array4 vs pseudo-median reference chip



Array5 vs pseudo-median reference chip



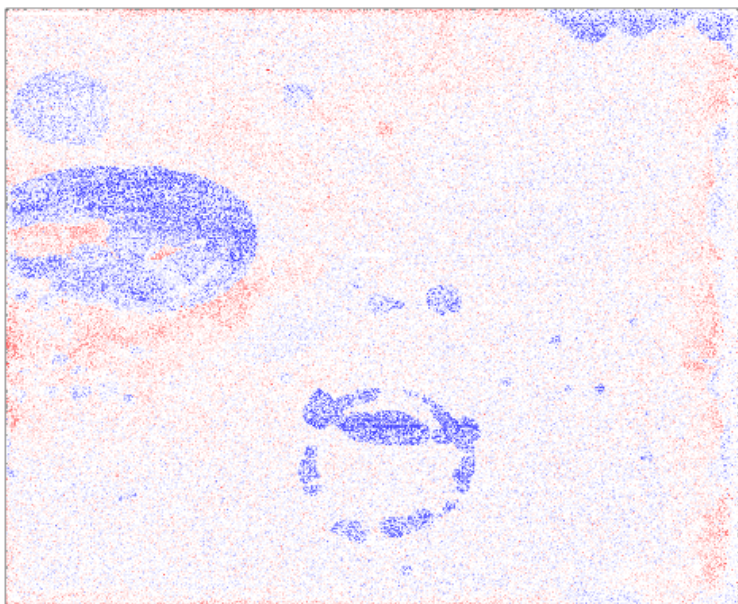
Array6 vs pseudo-median reference chip



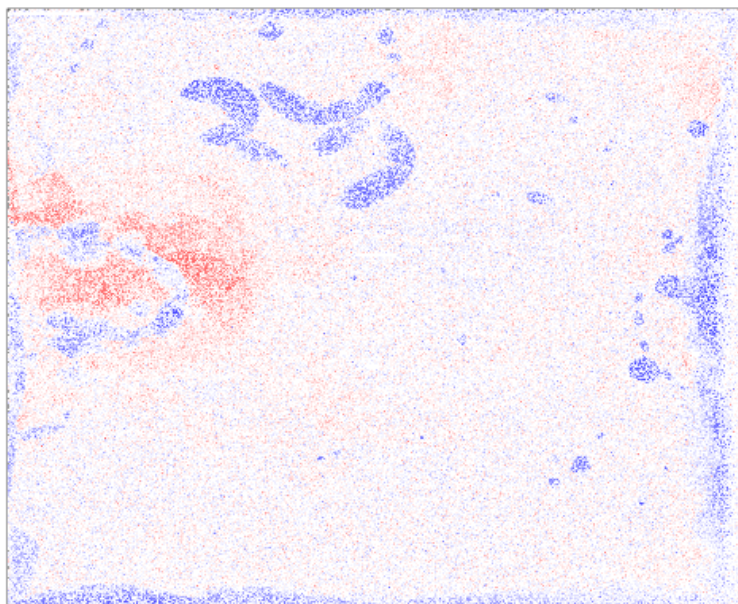


## 2D virtual PLM image for model characteristic: resides

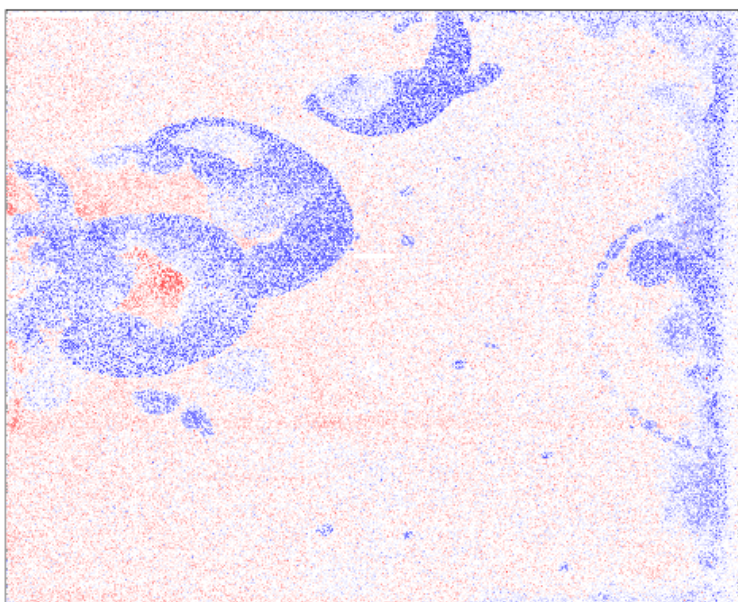
Array1



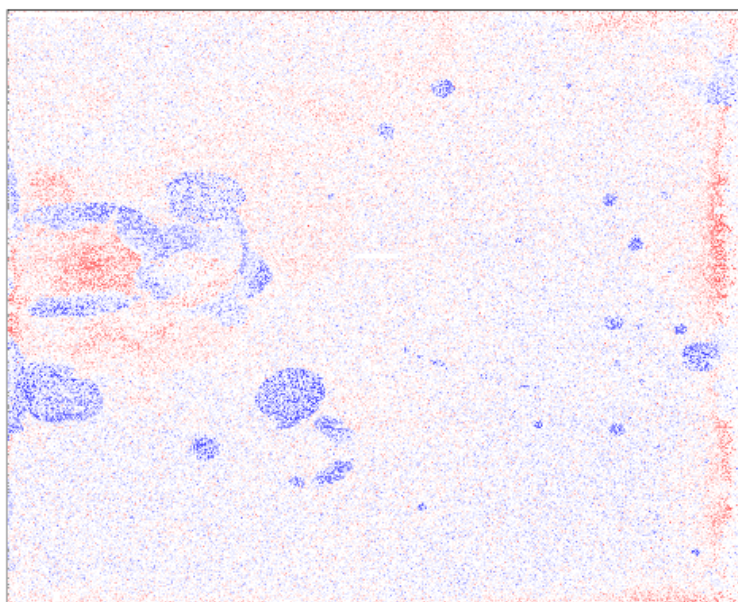
Array2



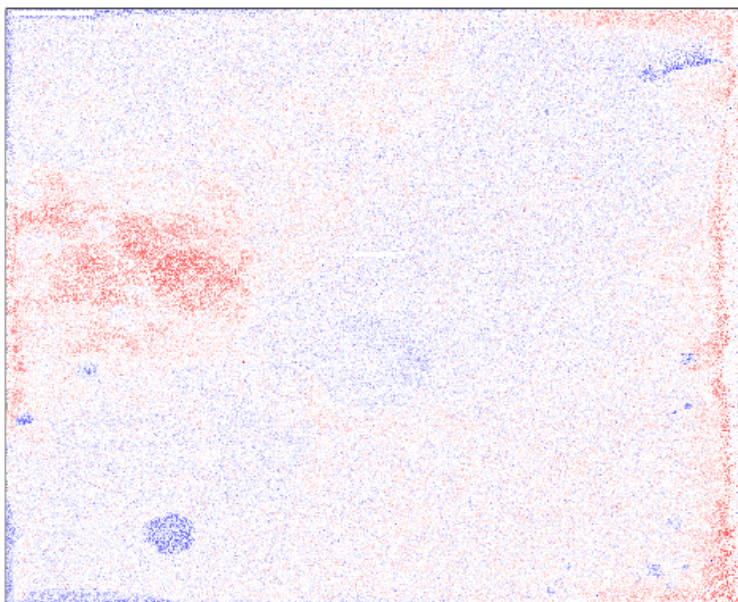
Array3



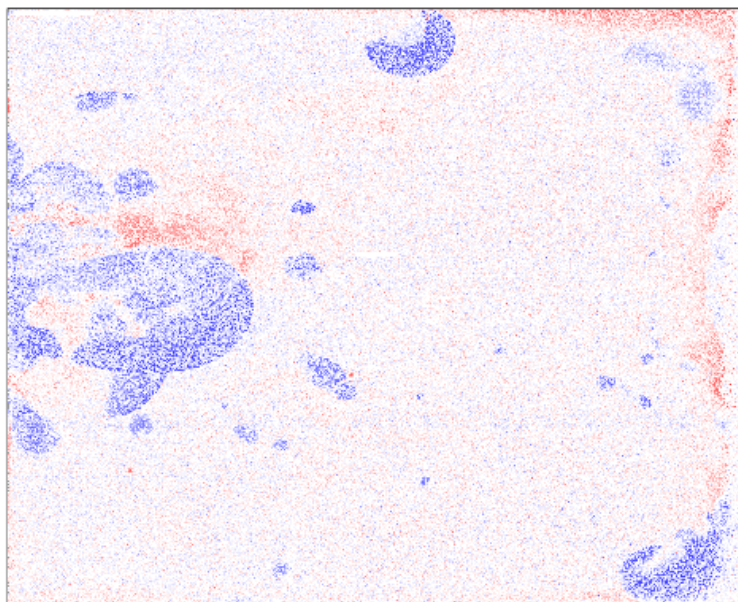
Array4



Array5



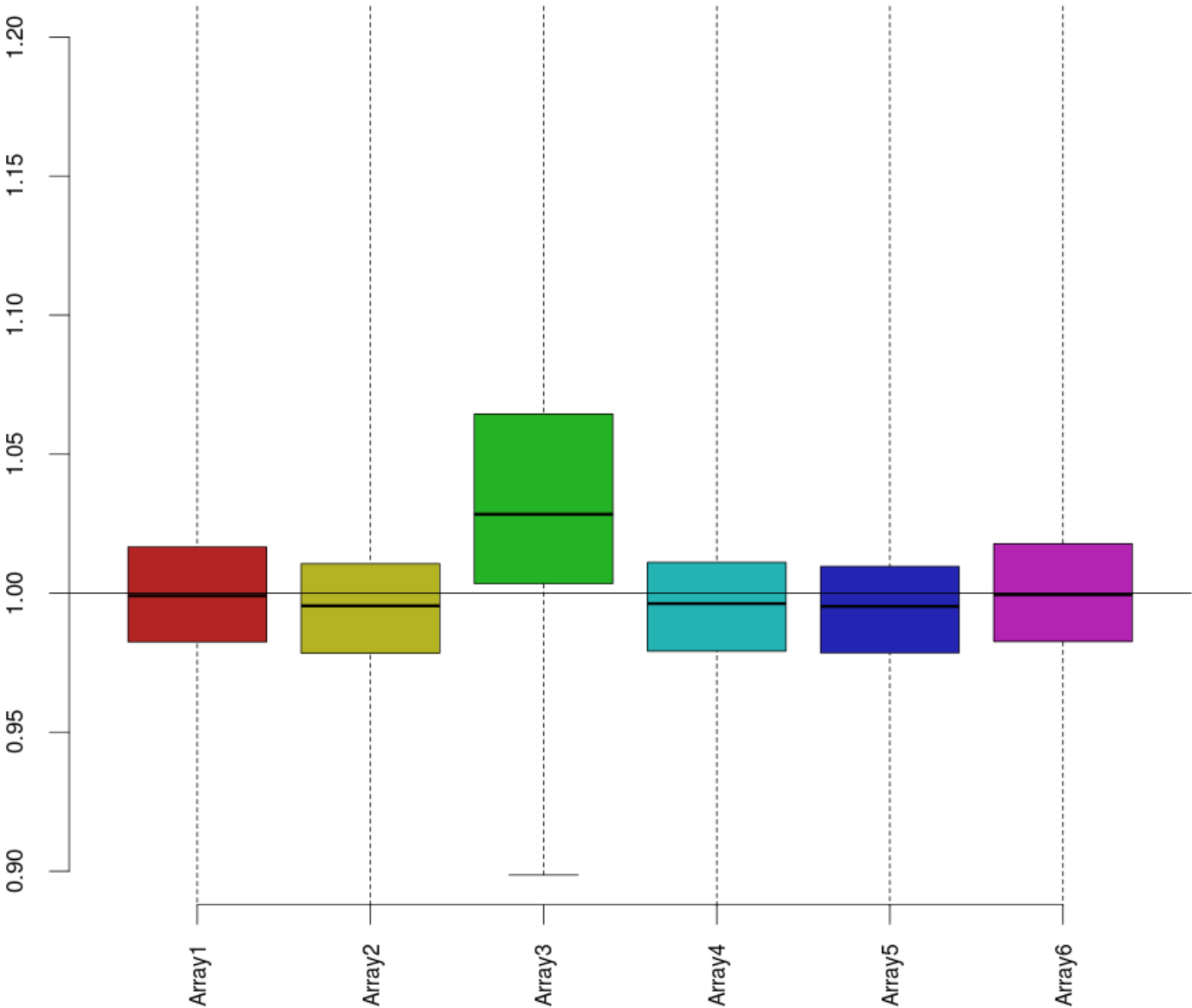
Array6





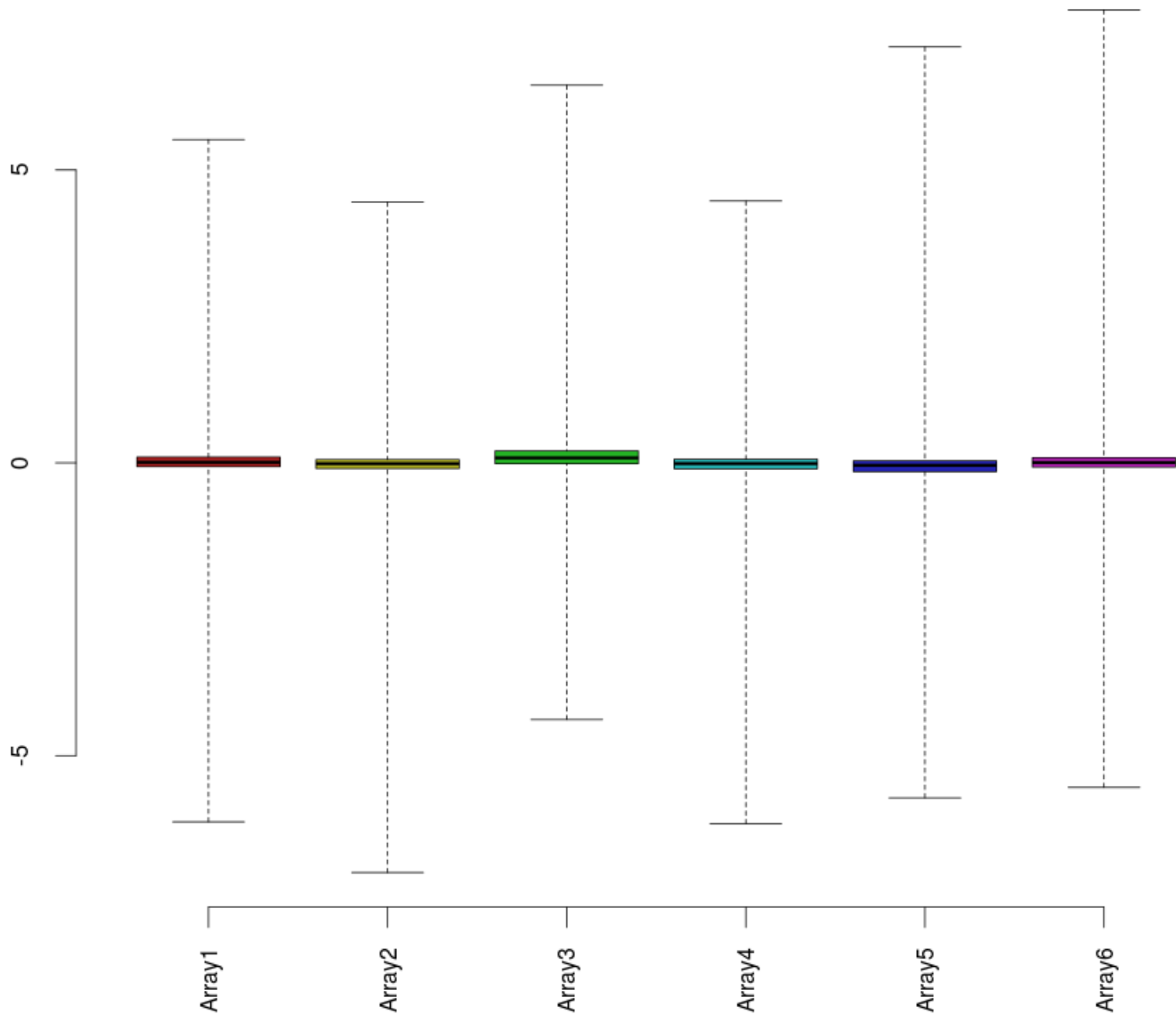
# Normalized Unscaled Standard Errors (NUSE)

NUSE median value should be < 1.1

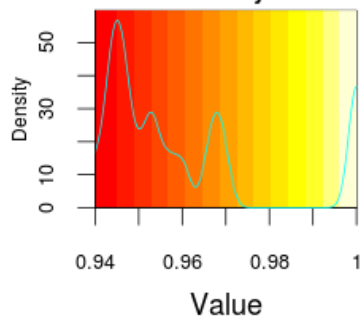


# Relative Log Expression (RLE)

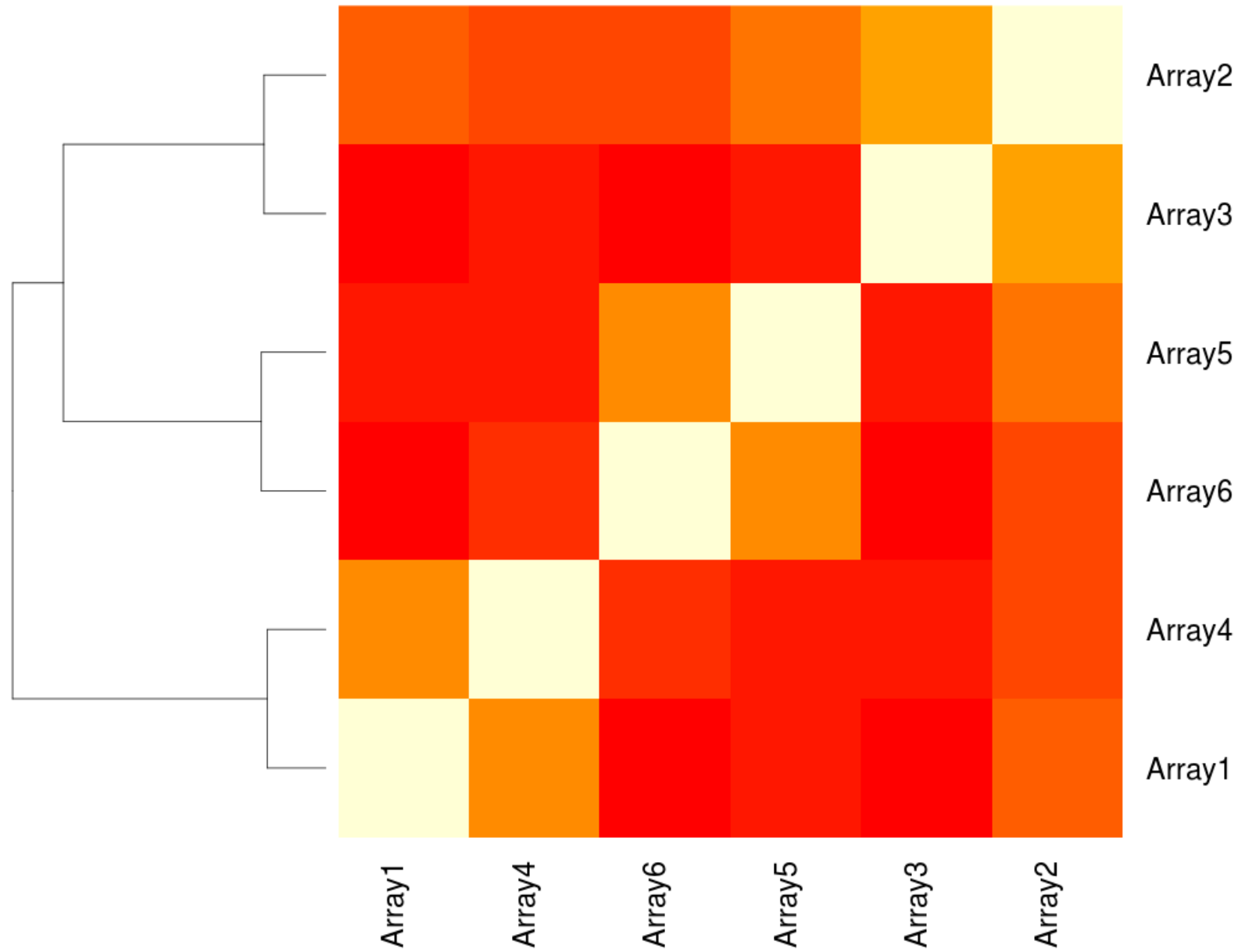
RLE distributions should be centered around 0



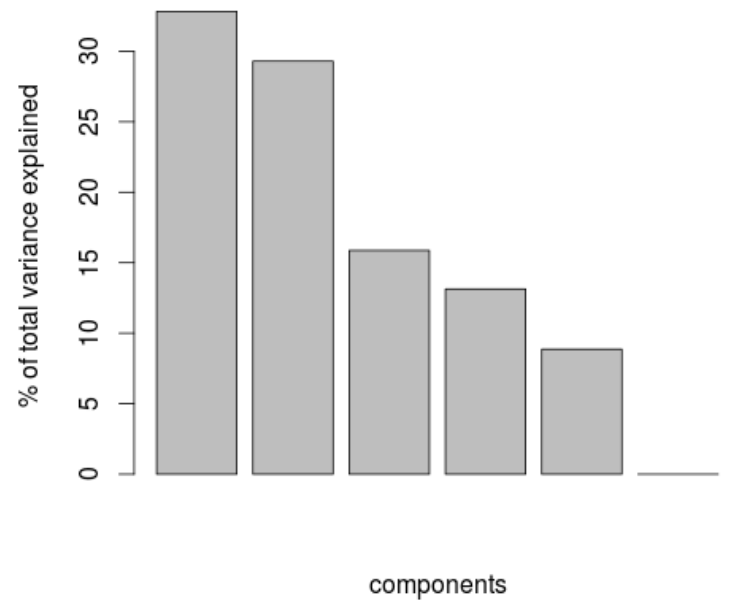
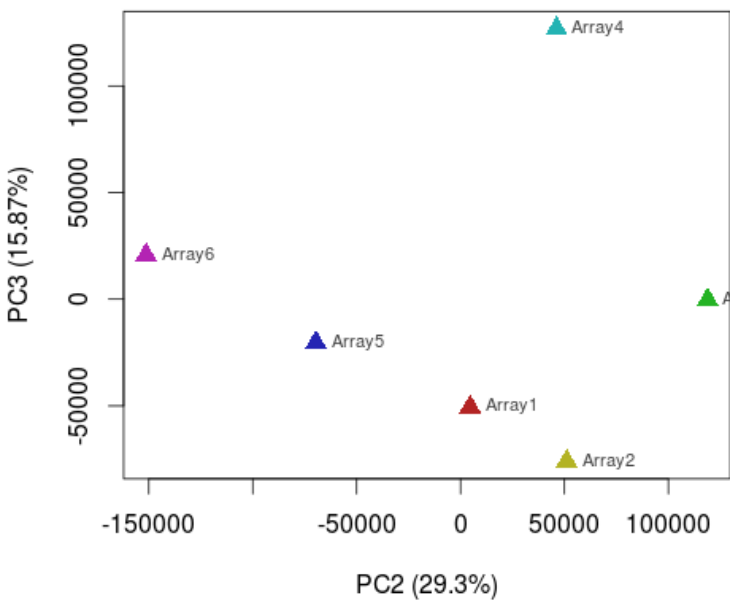
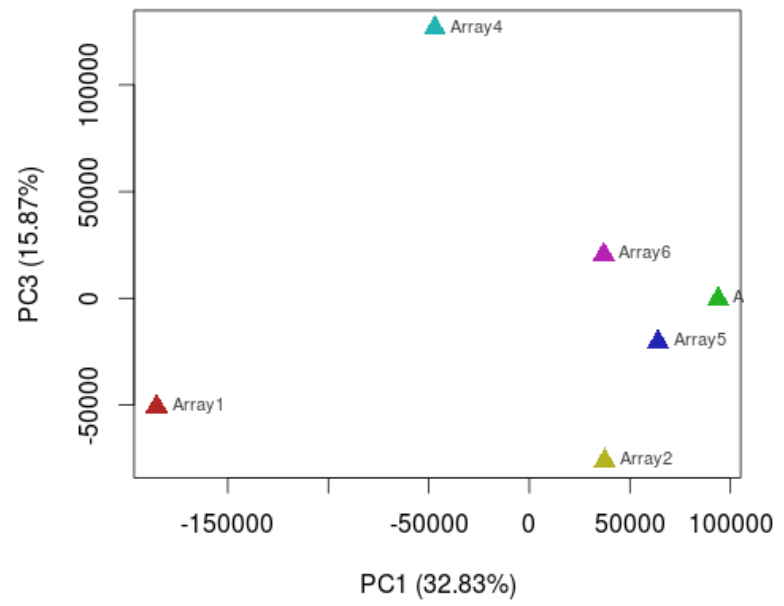
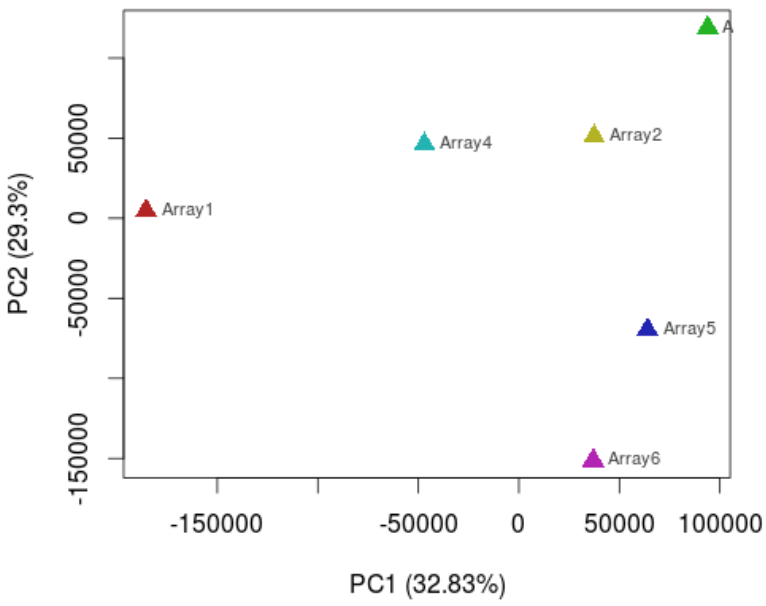
Color Key  
and Density Plot



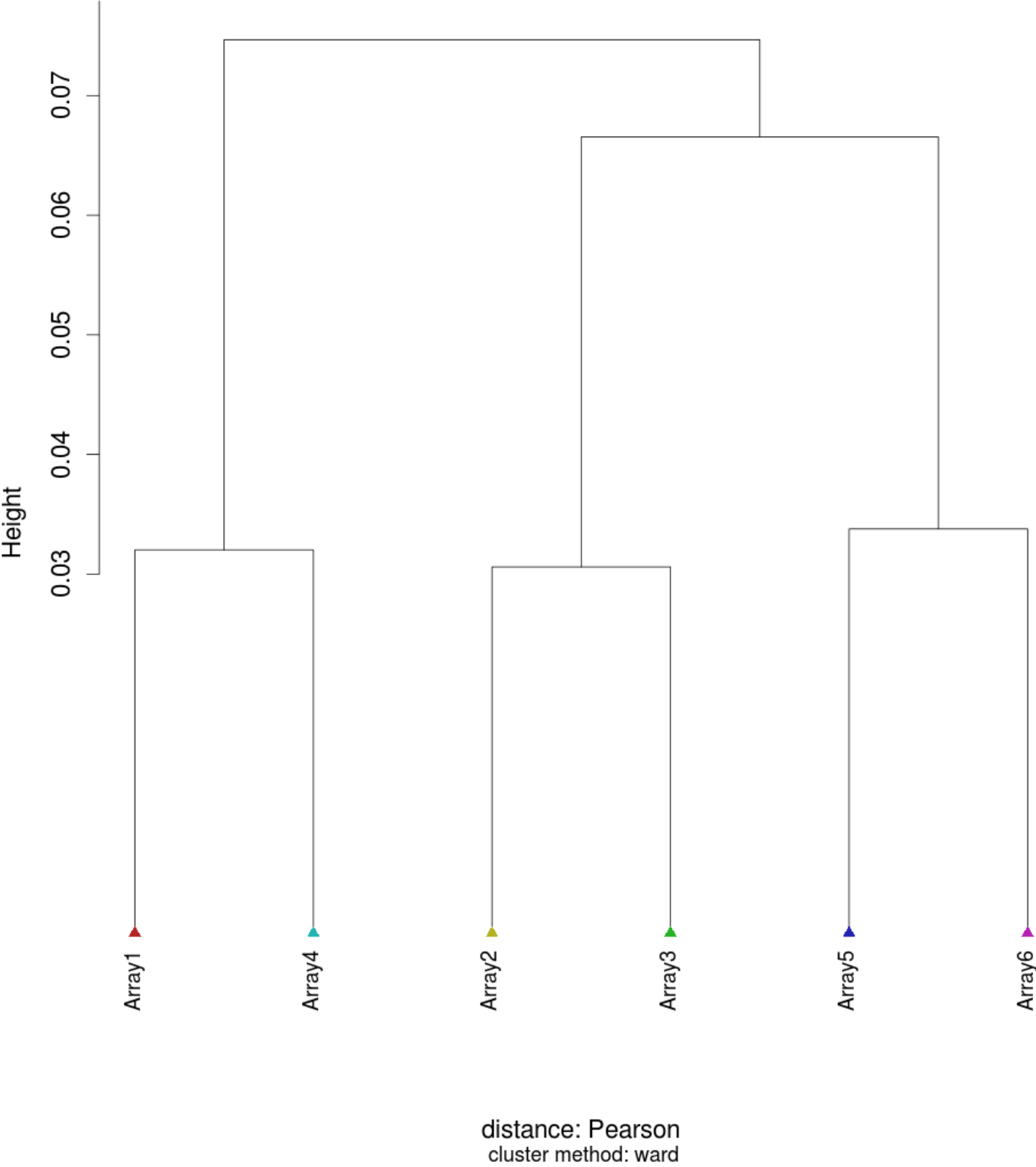
Raw data correlation plot  
correlation method: pearson  
cluster method: ward



# PCA analysis of Raw data



Cluster dendrogram of raw data



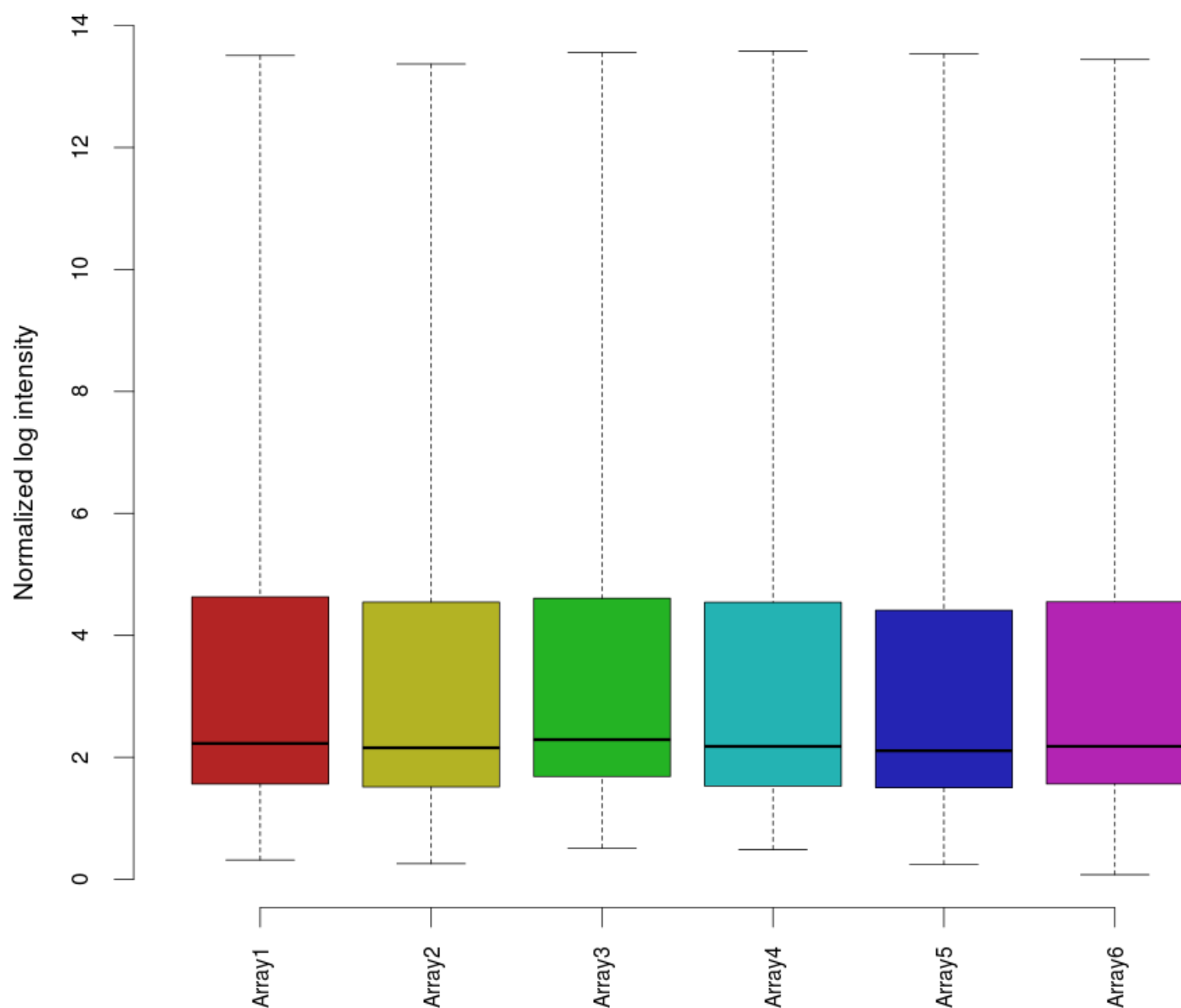
## Pre-processing of Raw Data

Method: GCRMA

Annotation: hgu133plus2\_Hs\_ENSG

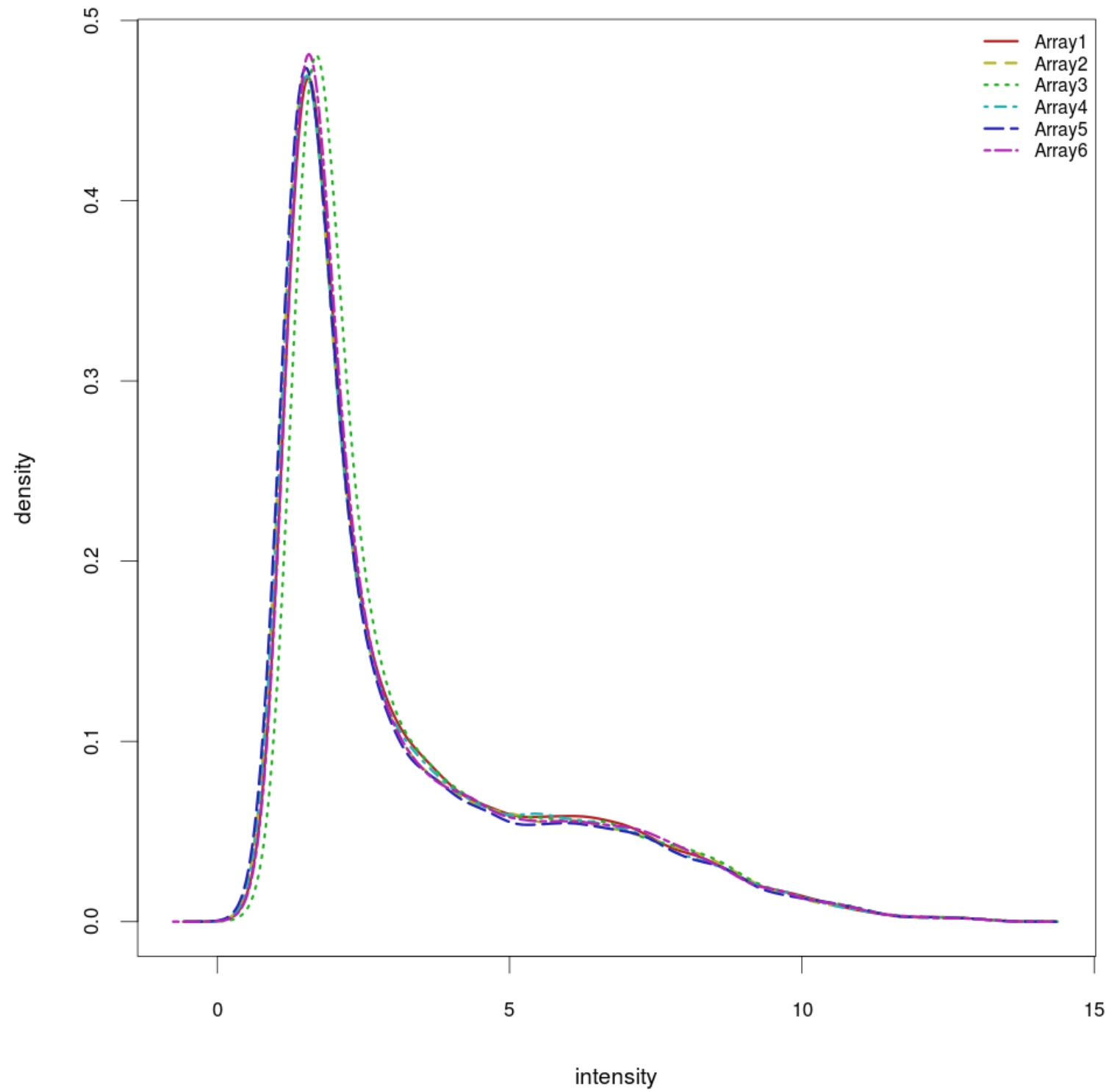
# Boxplot after GCRMA

Distributions should be comparable between arrays



# Density histogram after GCRMA

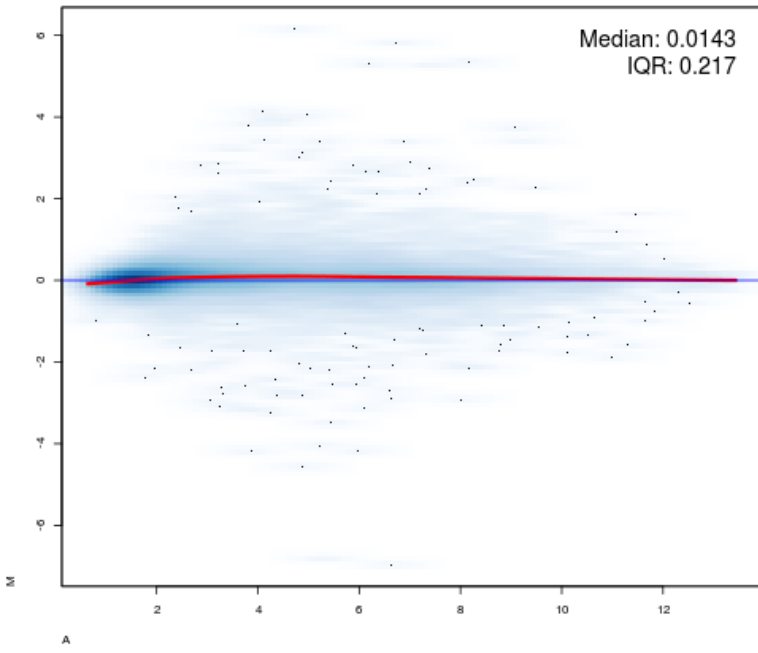
Curves should be comparable between arrays



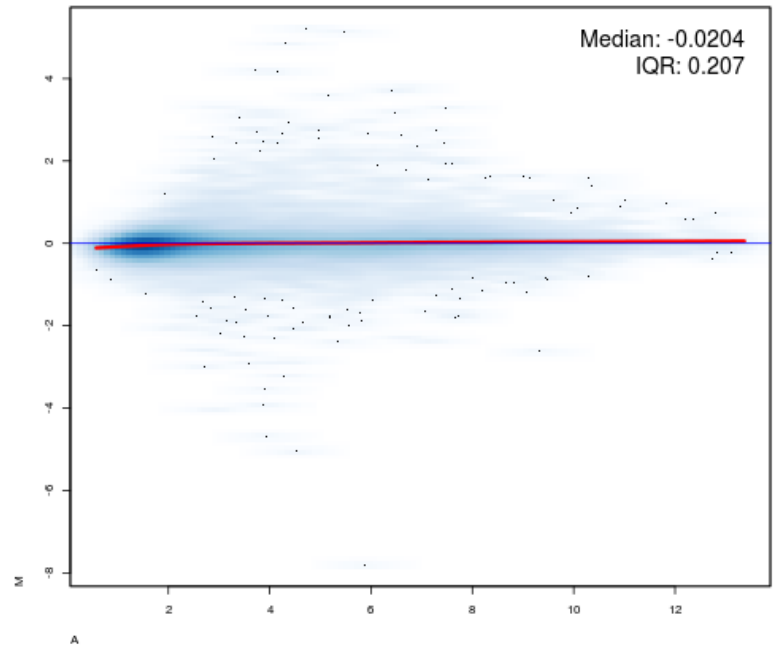


# MA plots afterGCRMANormalization

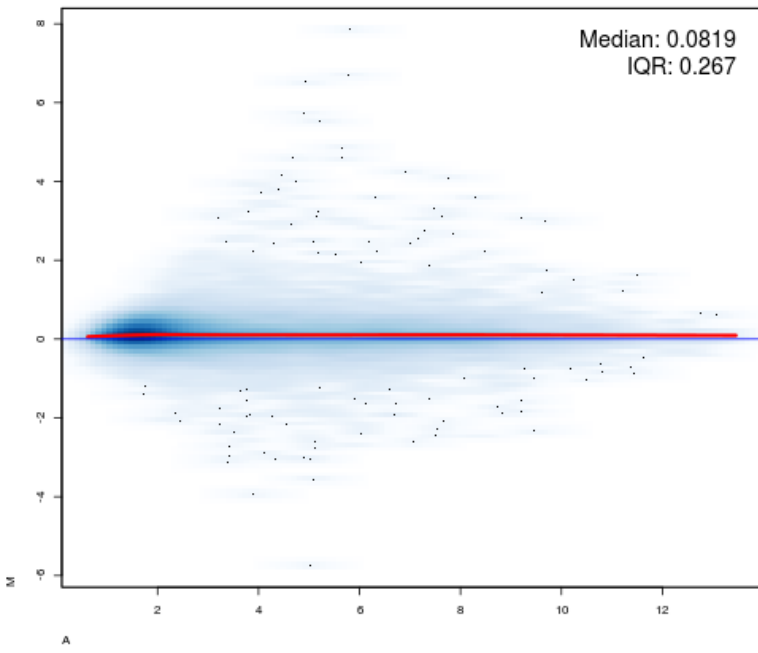
Array1 vs pseudo-median reference chip



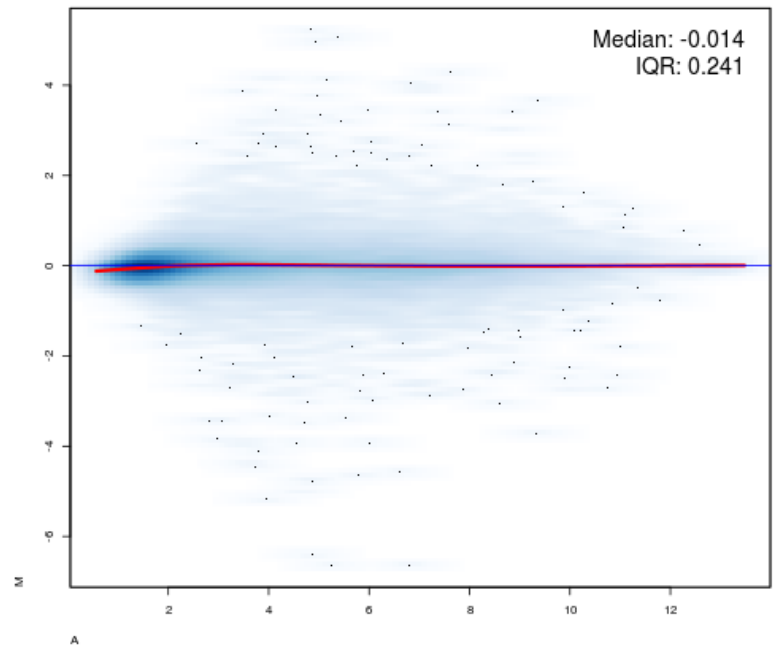
Array2 vs pseudo-median reference chip



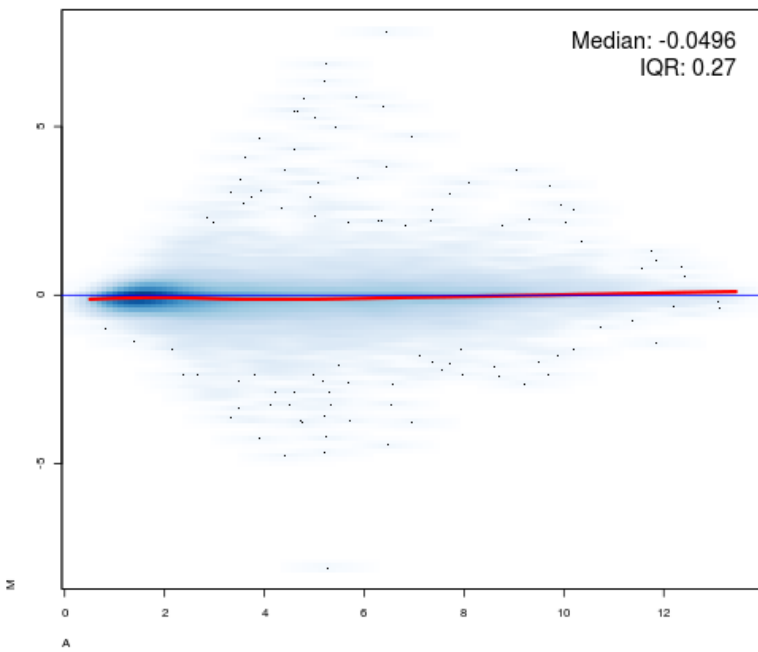
Array3 vs pseudo-median reference chip



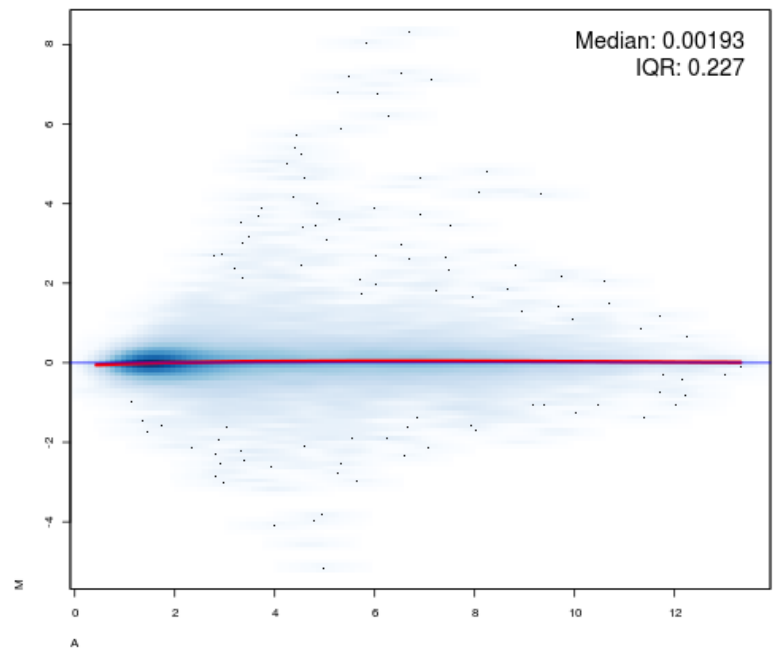
Array4 vs pseudo-median reference chip



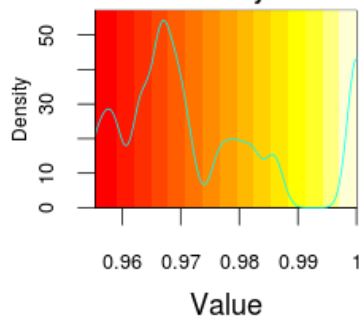
Array5 vs pseudo-median reference chip



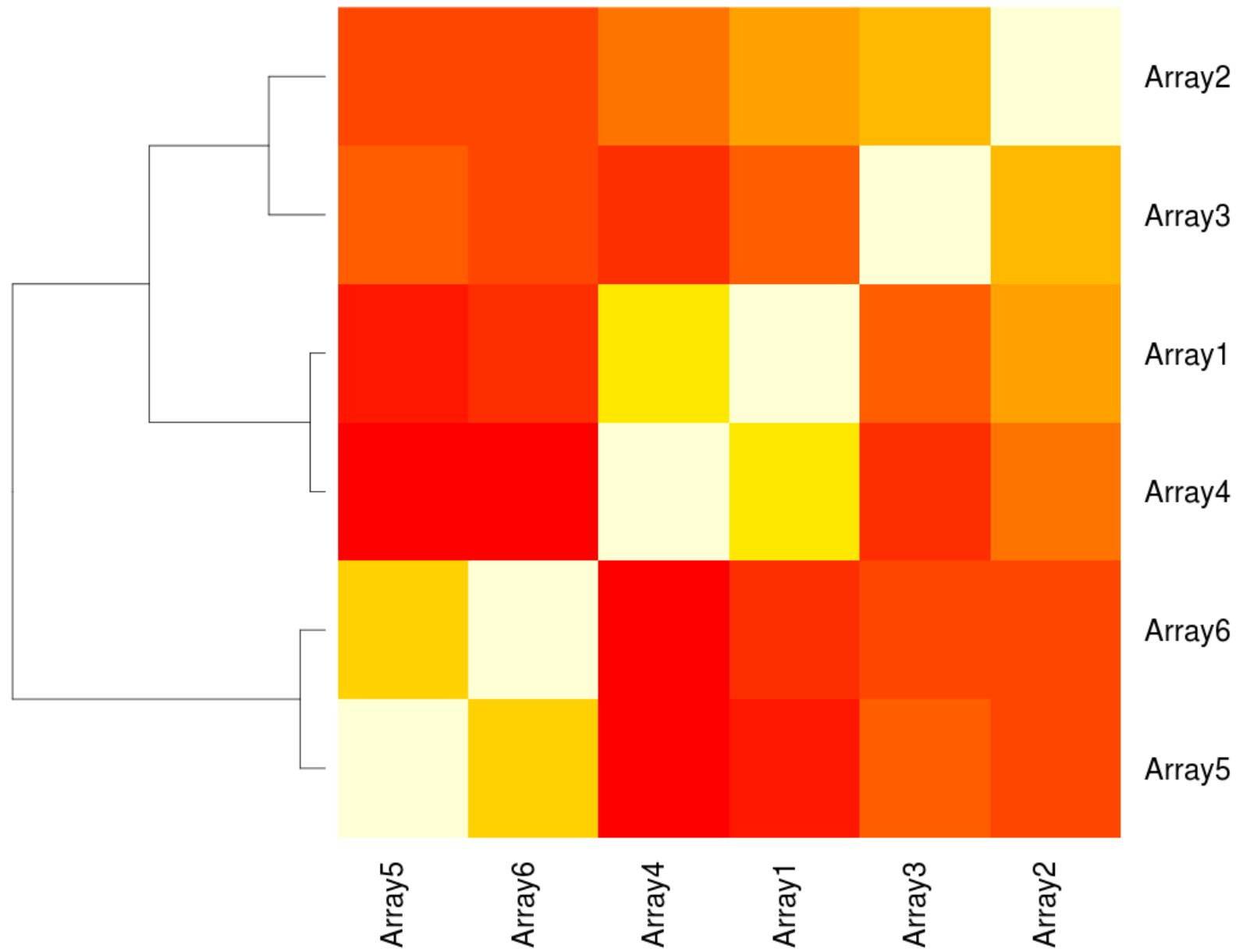
Array6 vs pseudo-median reference chip



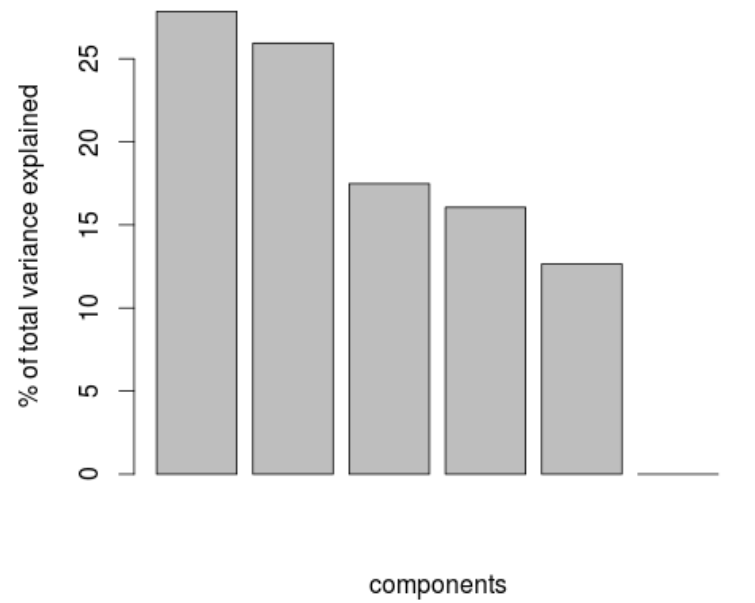
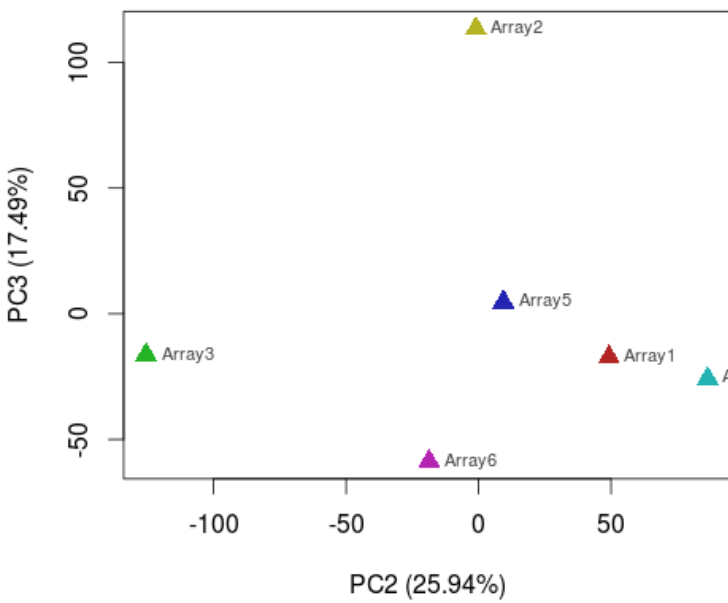
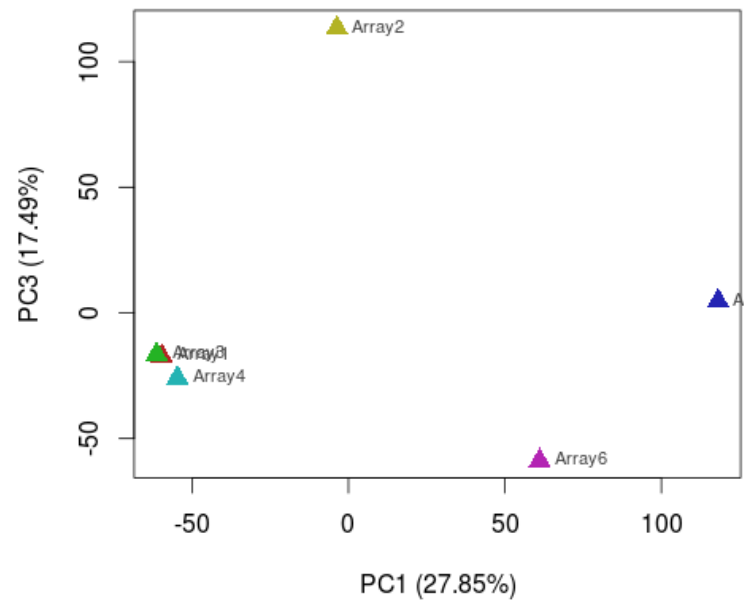
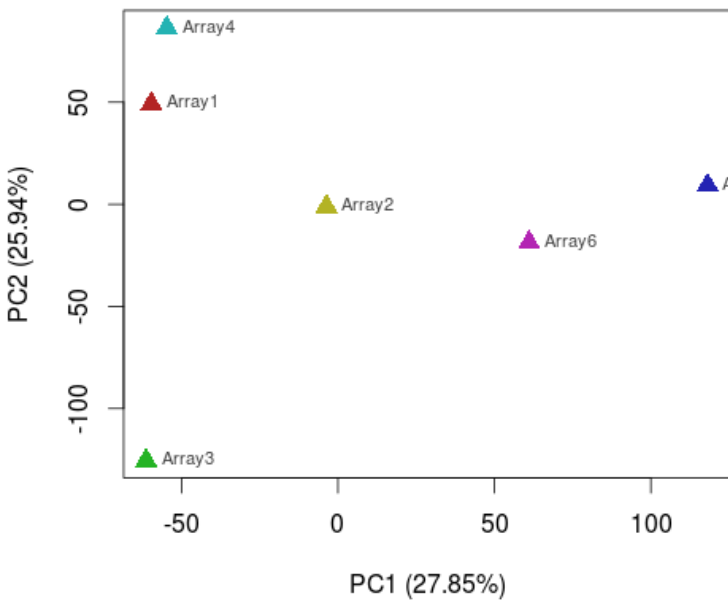
Color Key  
and Density Plot



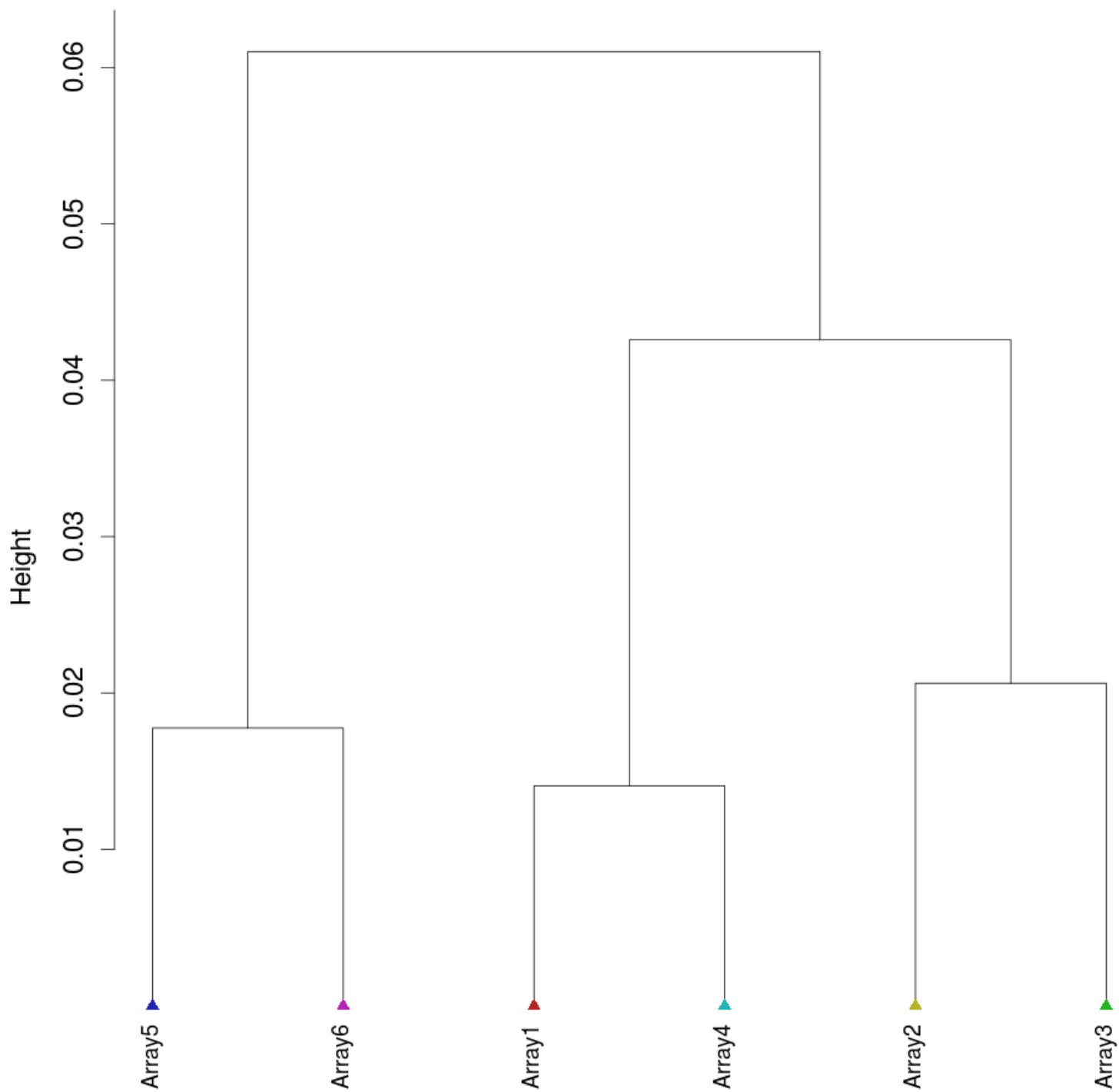
Array correlation plot  
after GCRMA normalization  
correlation method: pearson  
cluster method: ward



# PCA analysis after GCRMA normalization



Cluster dendrogram of GCRMA normalized data



distance: Pearson  
cluster method: ward