

Quality Control & Pre-processing Evaluation
of
E-GEOD-4600.raw.1_0
REPORT

Array names and grouping

ArrayDataFile	SourceName	FactorValue
GSM102842.CEL	Array1	Group1
GSM102878.CEL	Array2	Group2
GSM102883.CEL	Array3	Group3
GSM102825.CEL	Array4	Group1
GSM102879.CEL	Array5	Group2
GSM102880.CEL	Array6	Group2
GSM102881.CEL	Array7	Group3
GSM102876.CEL	Array8	Group4
GSM102877.CEL	Array9	Group4
GSM102875.CEL	Array10	Group4
GSM102882.CEL	Array11	Group3
GSM102870.CEL	Array12	Group1

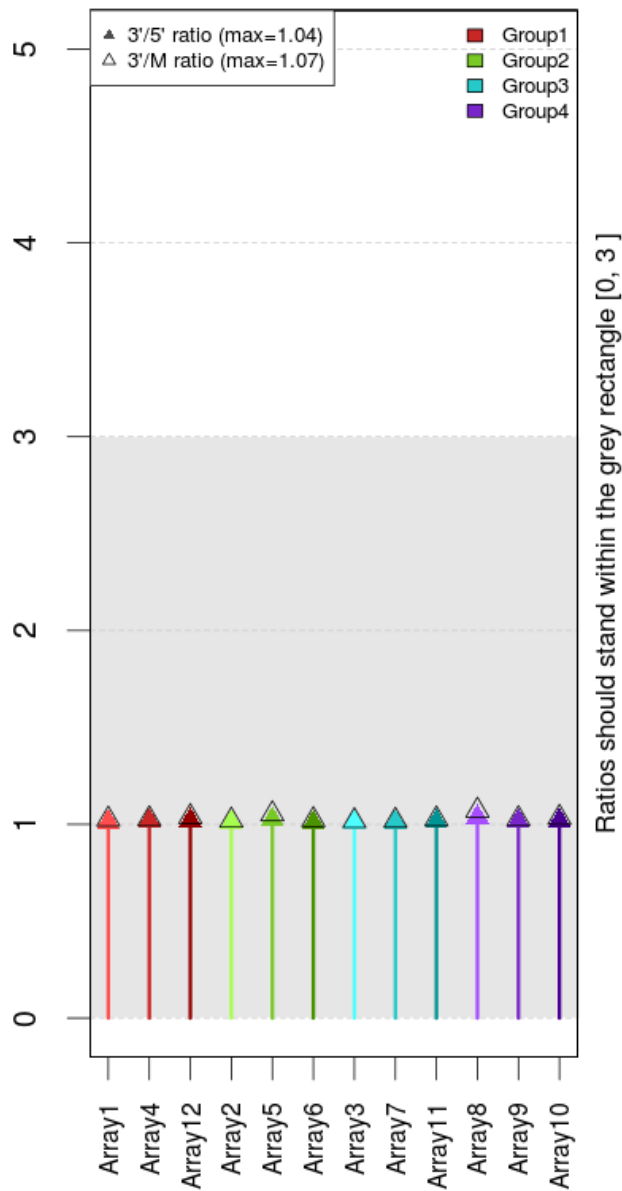
Summary of raw data quality indicators

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

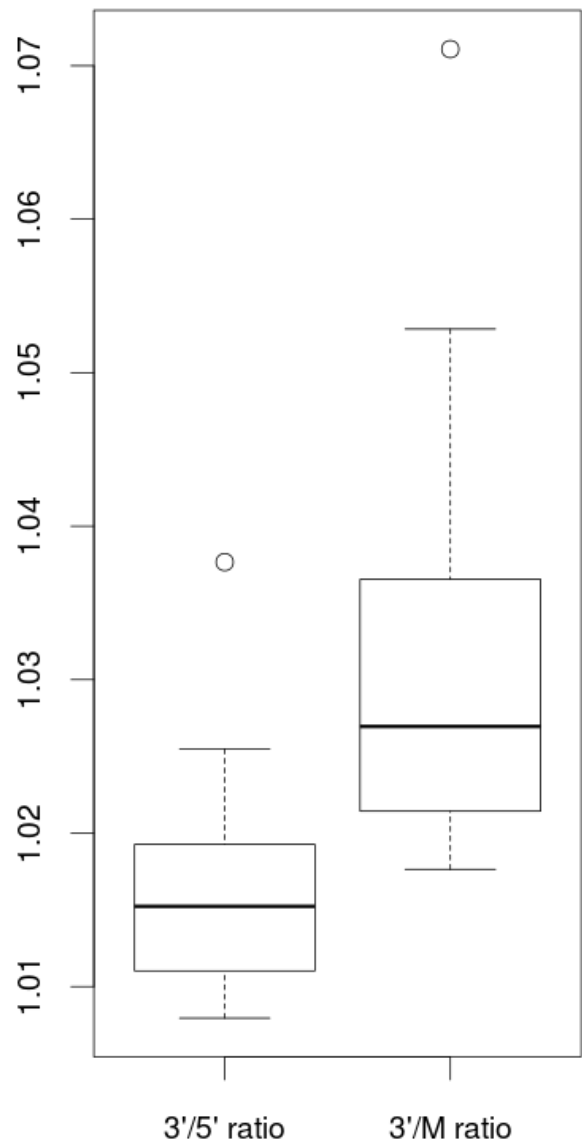
	3'/5' beta-actin (cutoff=3)	3'/5' GAPDH (cutoff= 1.25)	Hybridization BioB<BioC<BioD<CreX	Hybridization BioB=Present	Percent Present spread<= 10%	Background spread<=20%	Log Scale Factor spread<=3
Array1	1.01	1.02	T	P	30 %	83	0.9
Array4	1.02	1.02	T	P	33 %	117	0.66
Array12	1.02	1.02	T	P	33 %	74	0.75
Array2	1.01	1.01	T	P	31 %	85	1.26
Array5	1.03	1.03	T	P	32 %	68	1
Array6	1.01	1.02	T	P	34 %	66	0.91
Array3	1.01	1.02	T	P	34 %	76	0.67
Array7	1.01	1.02	T	P	29 %	121	1.14
Array11	1.01	1.02	T	P	33 %	67	0.97
Array8	1.04	1.03	T	P	34 %	72	0.64
Array9	1.02	1.02	T	P	34 %	81	0.63
Array10	1.02	1.01	T	P	34 %	87	0.94

RNA degradation of beta-actin

3'/5' and 3'/M ratios



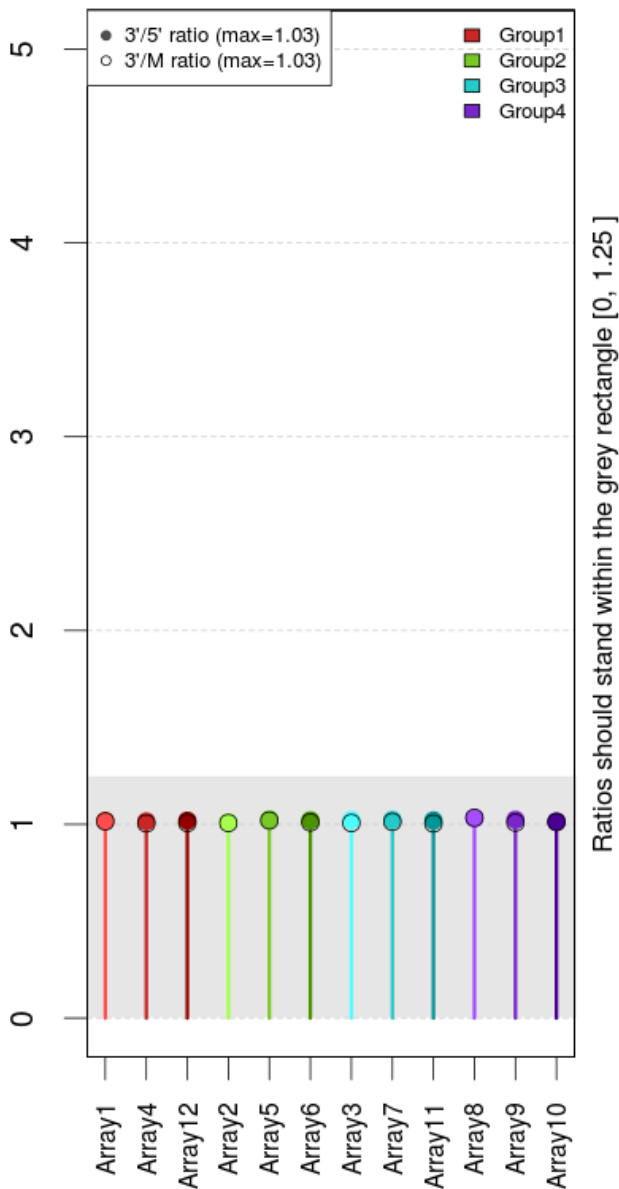
Boxplot of beta-actin ratios



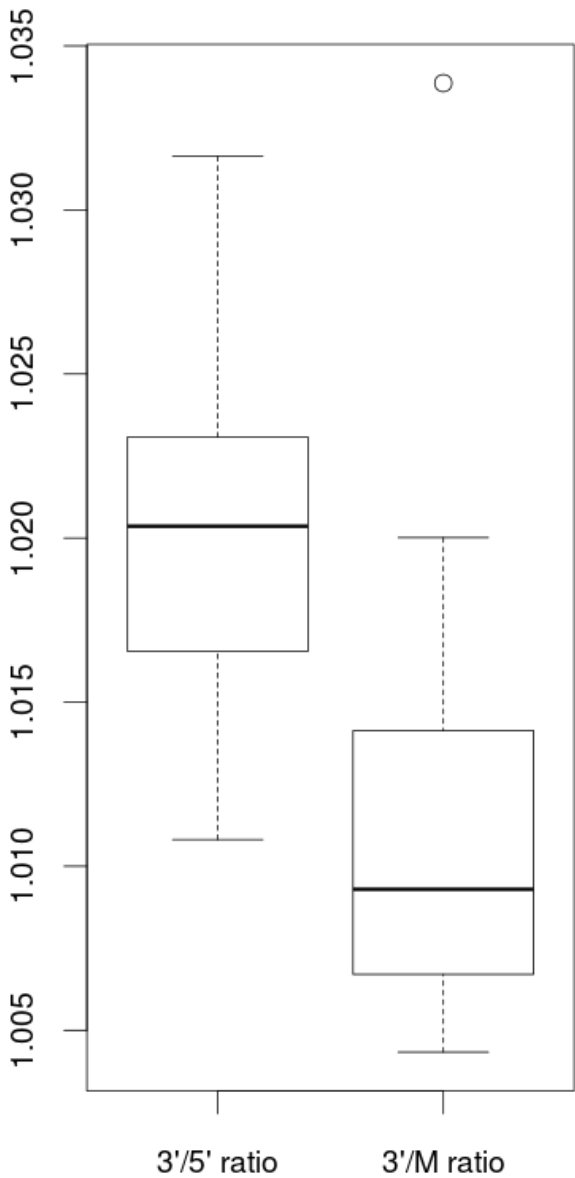
beta-actin QC: OK (all 3'/5' ratios < 3)

RNA degradation of GAPDH

3'/5' and 3'/M ratios

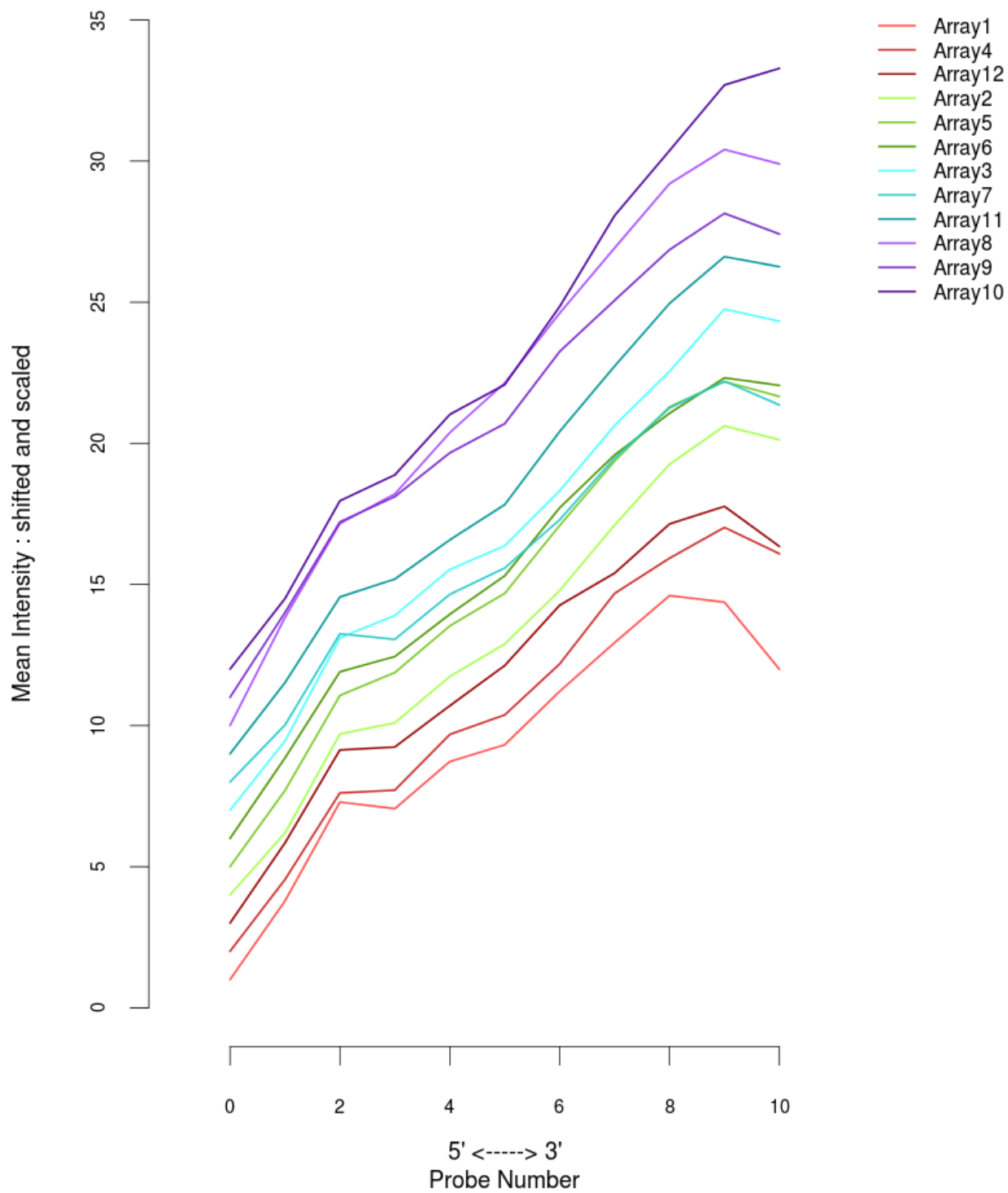


Boxplot of GAPDH ratios

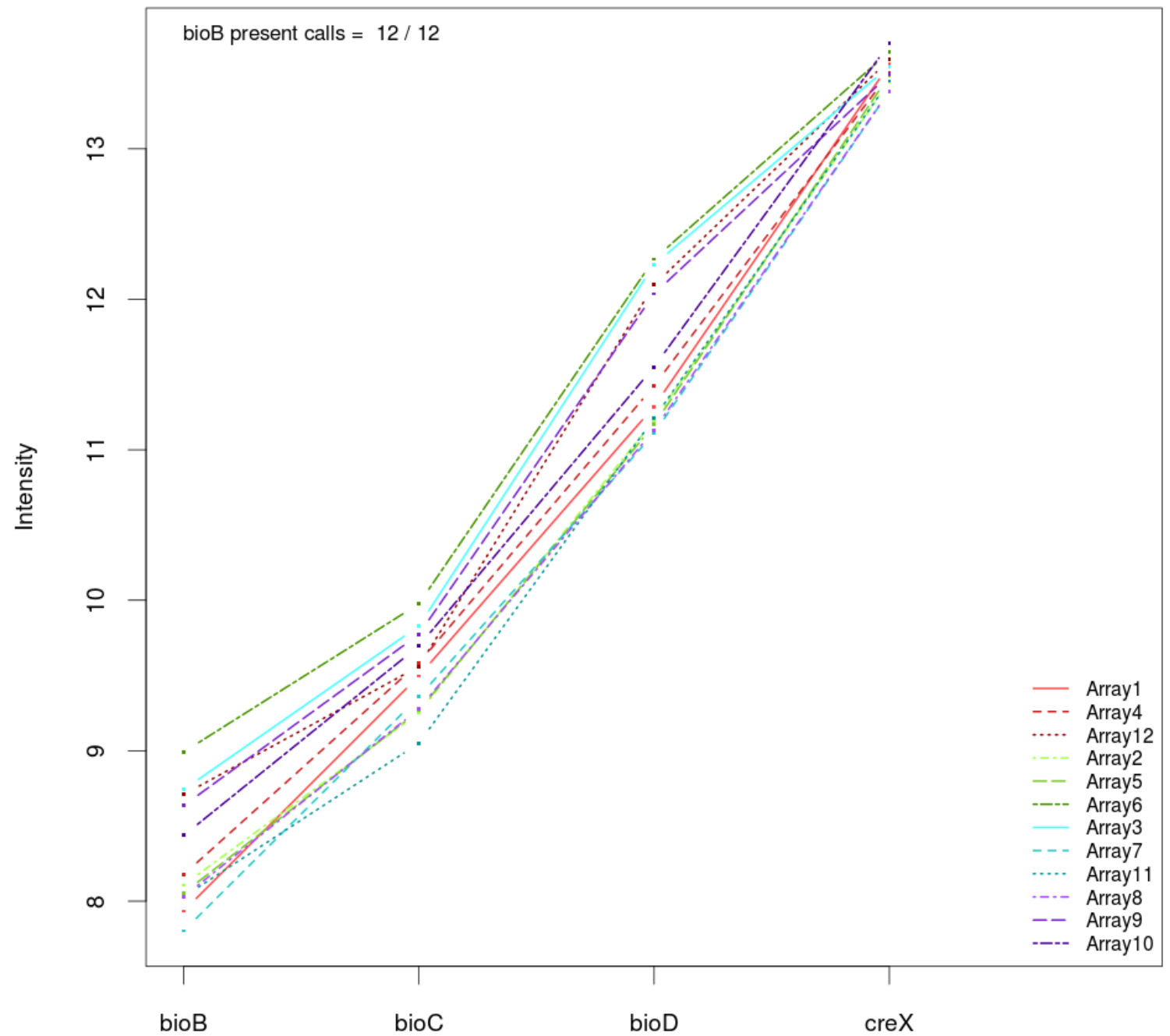


GAPDH QC: OK (all 3'/5' ratios < 1.25)

RNA degradation plot

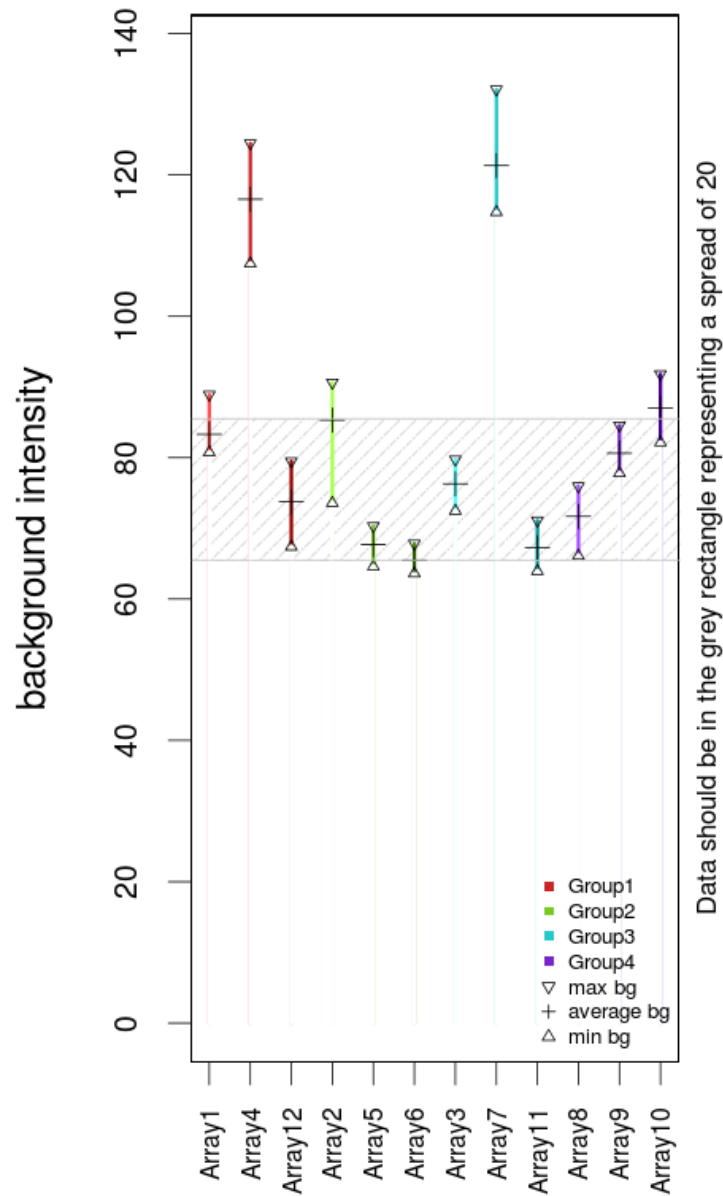


Spike-in Hybridization controls intensities and calls

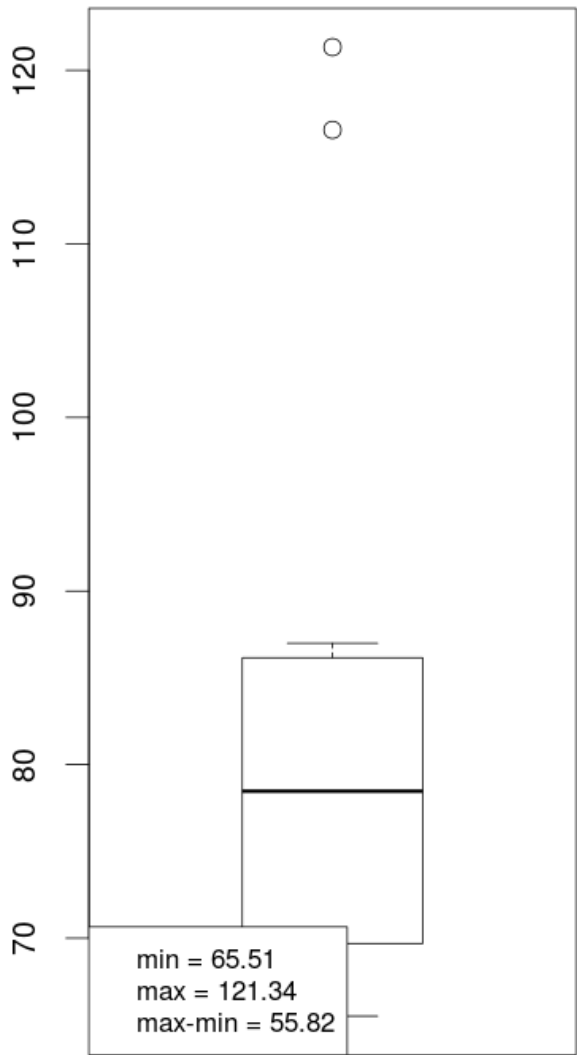


Intensities: OK (bioB < bioC < bioD < creX for all arrays)
BioB Present calls: OK (indeed all bioB are called present)

Plot of background intensity

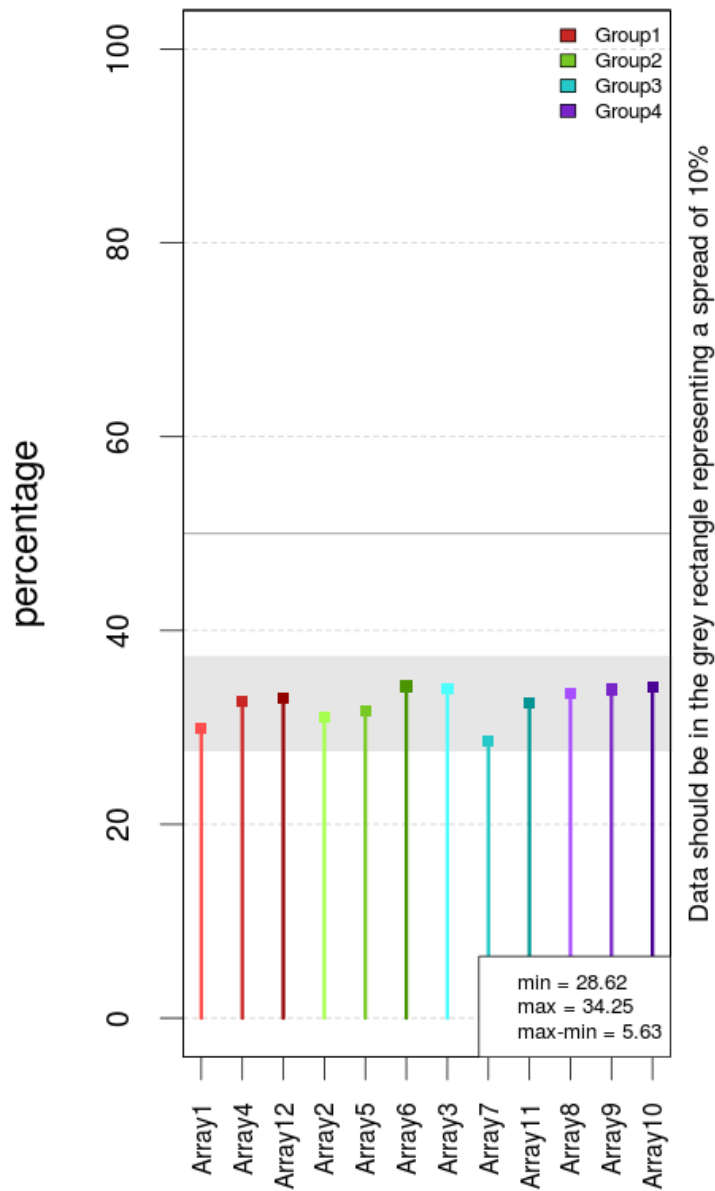


Average background intensity

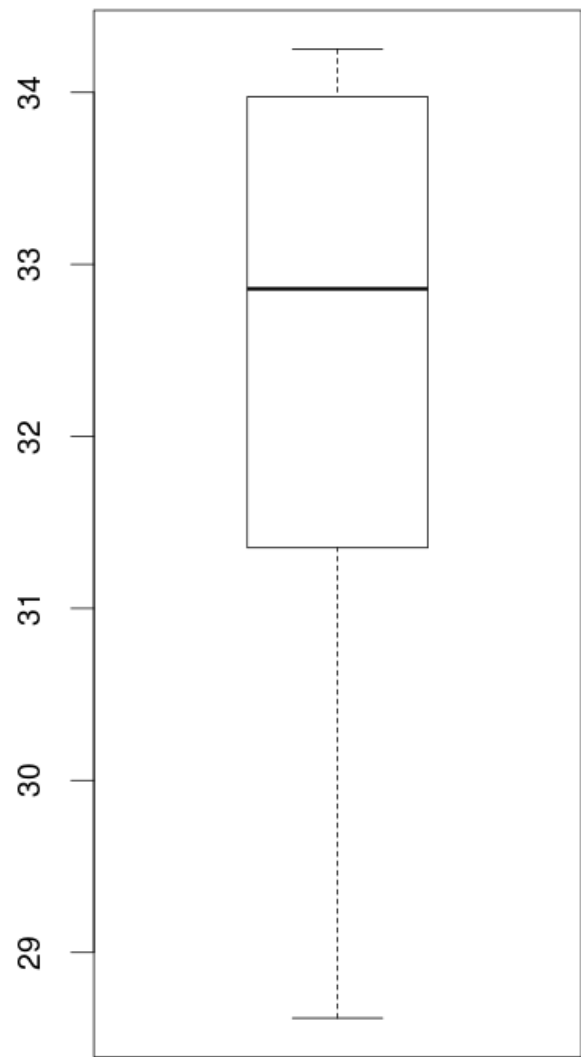


Background QC: not OK (spread > 20)

Plot of percent present

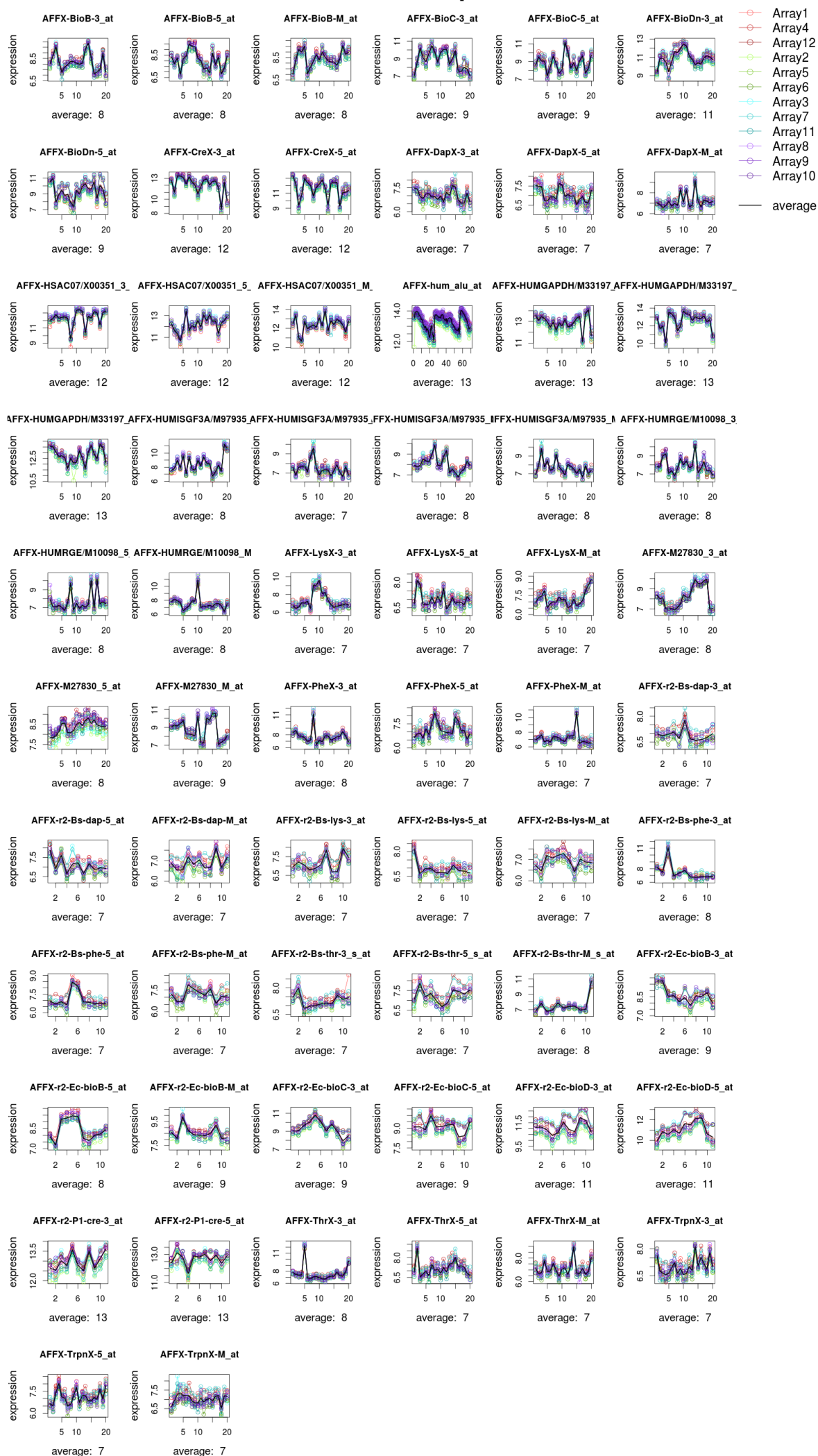


Boxplot of percent present

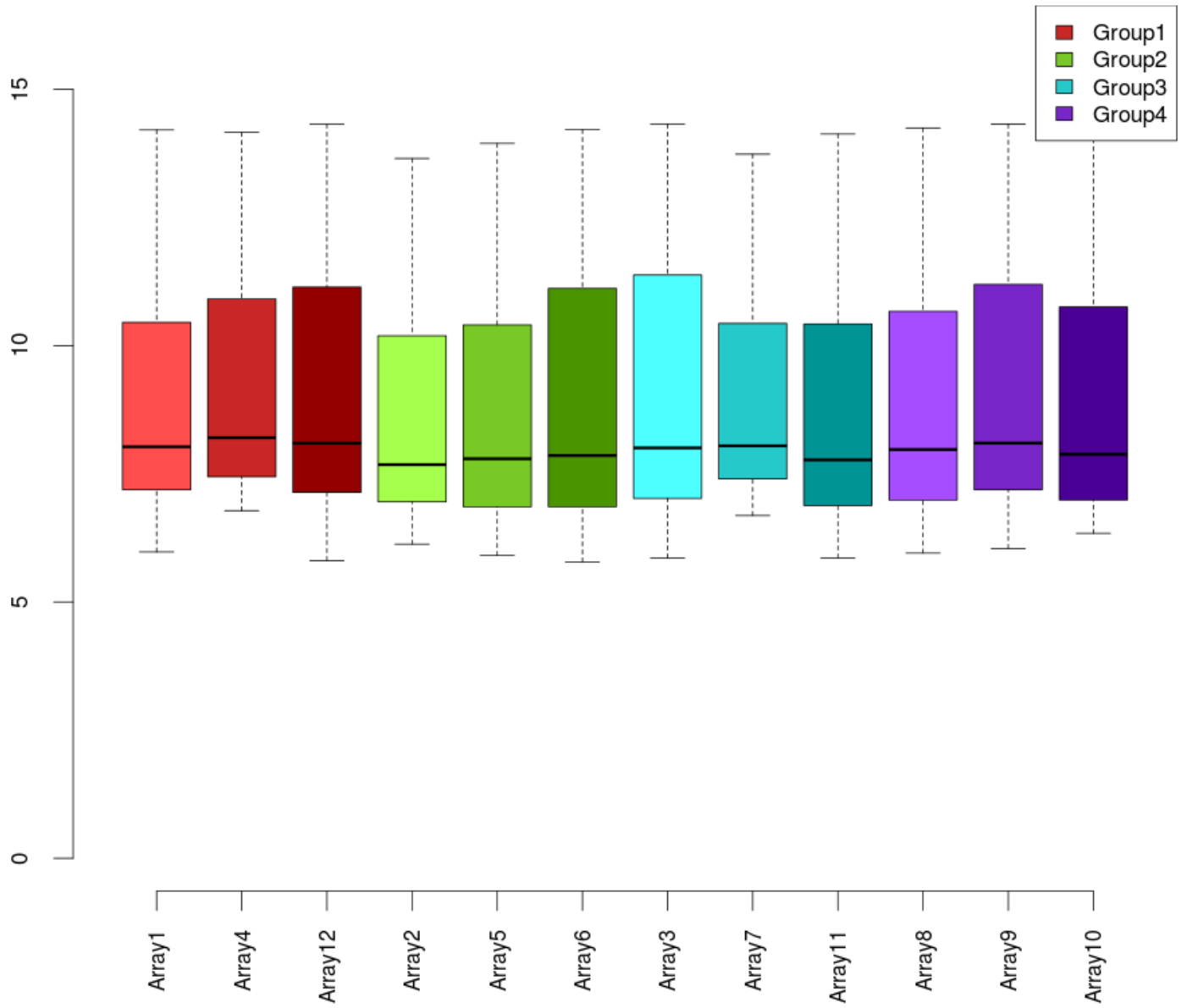


Percent present QC: OK (spread <= 10%)

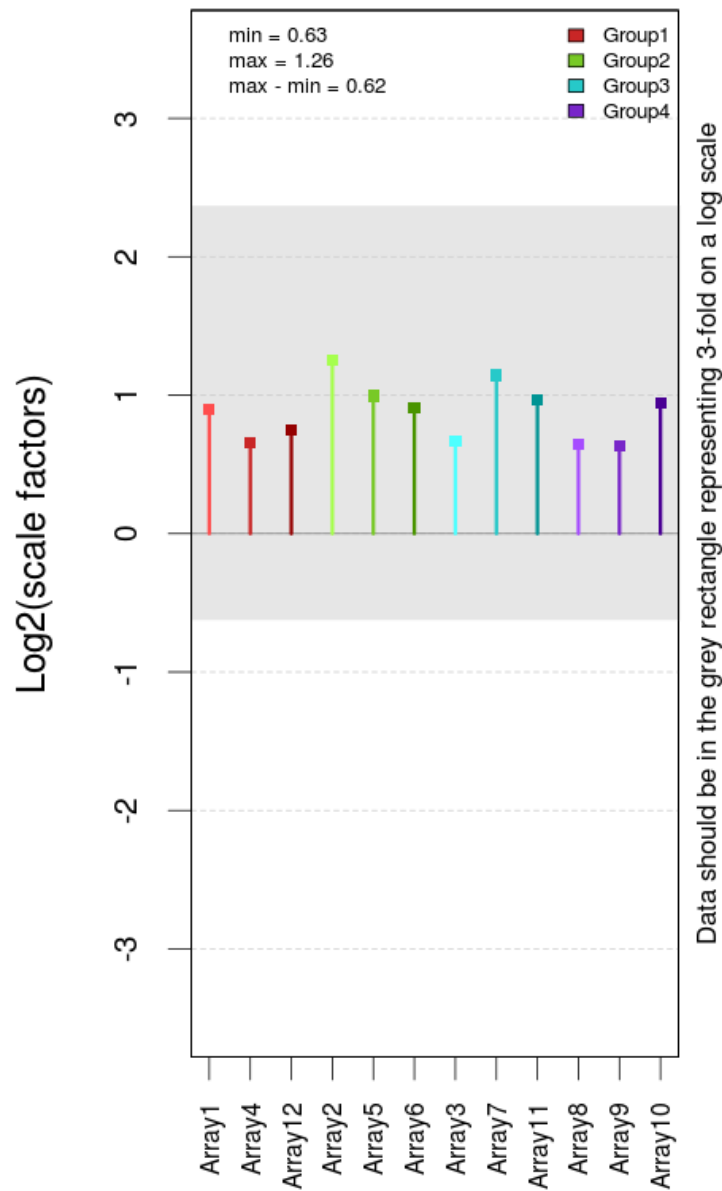
affx control profiles



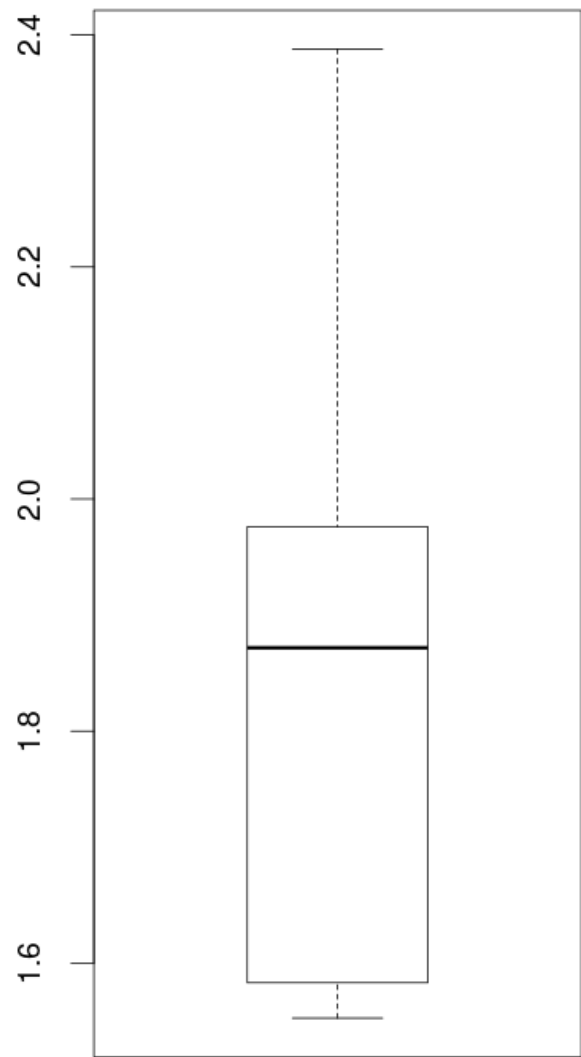
affx controls



Plot of Log scale factors



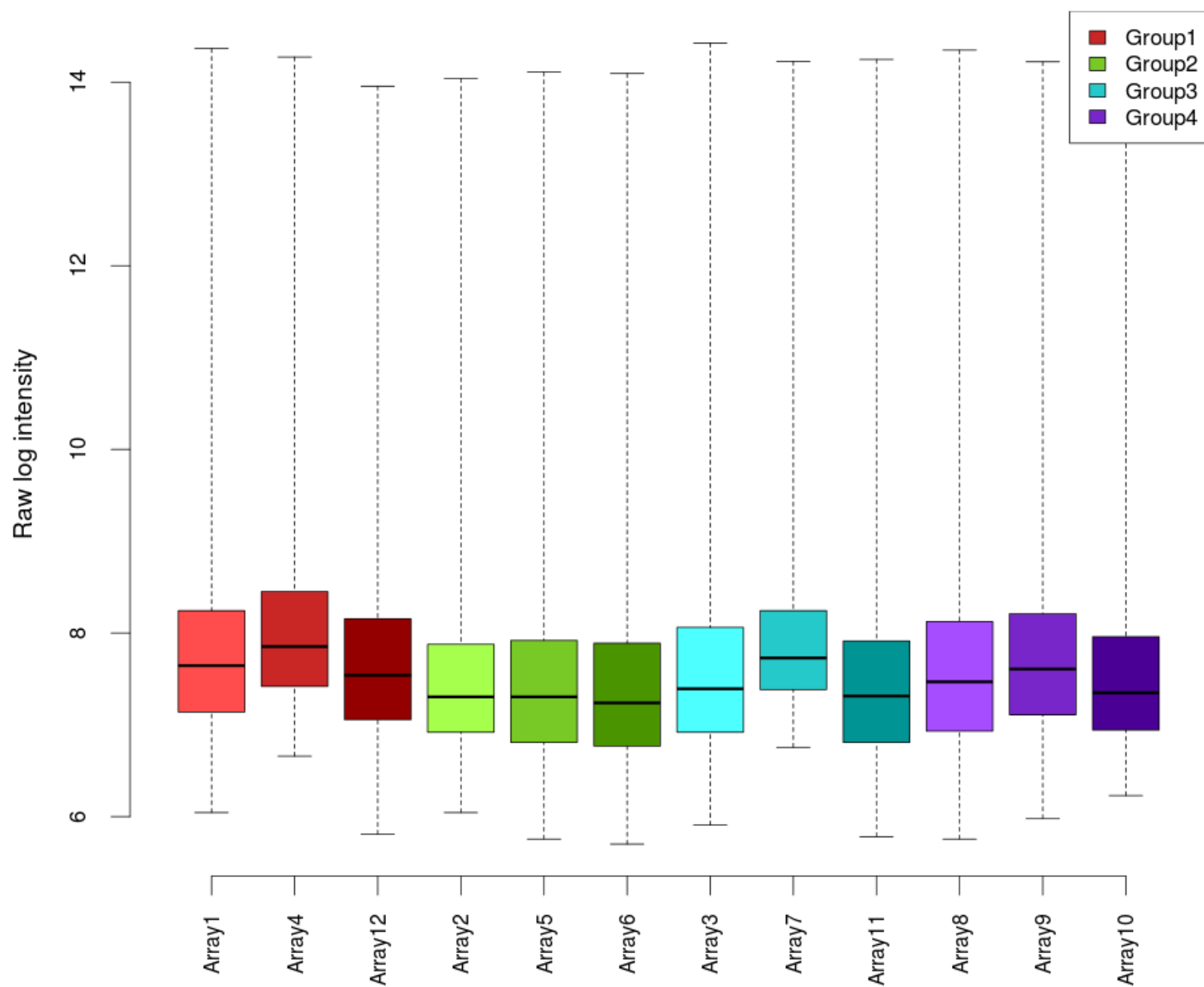
Boxplot of scale factors
(natural scale)



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

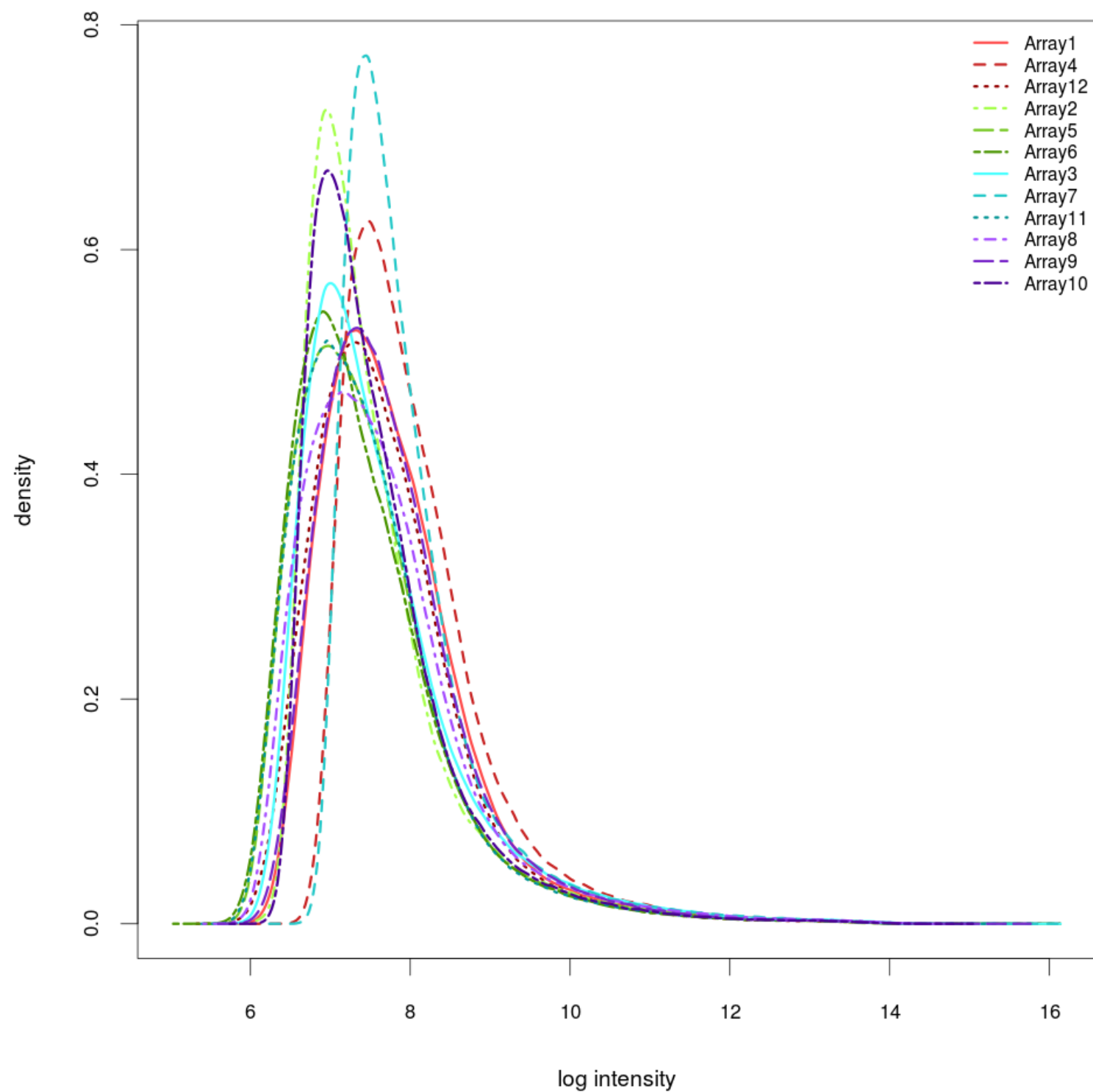
Boxplot of raw intensities

Distributions should be comparable between arrays



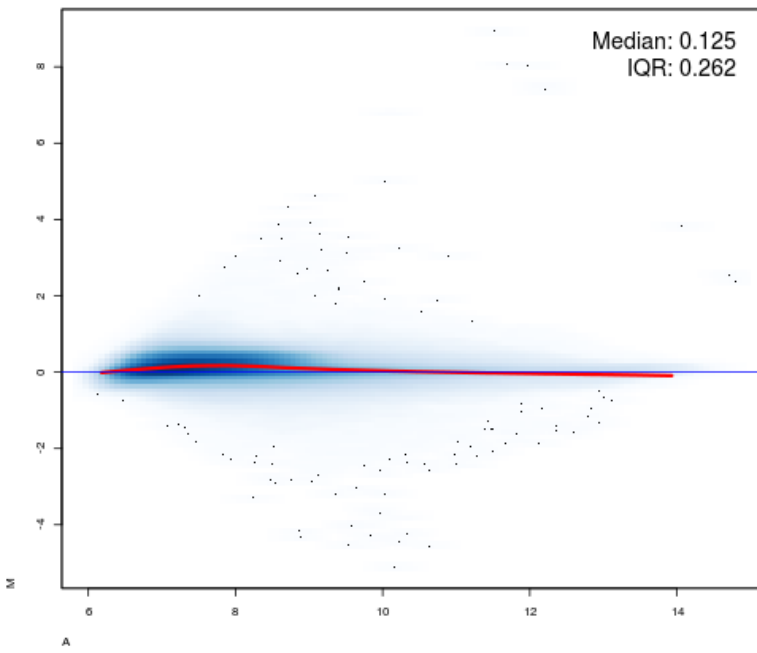
Density histogram of raw intensities

Curves should be comparable between arrays

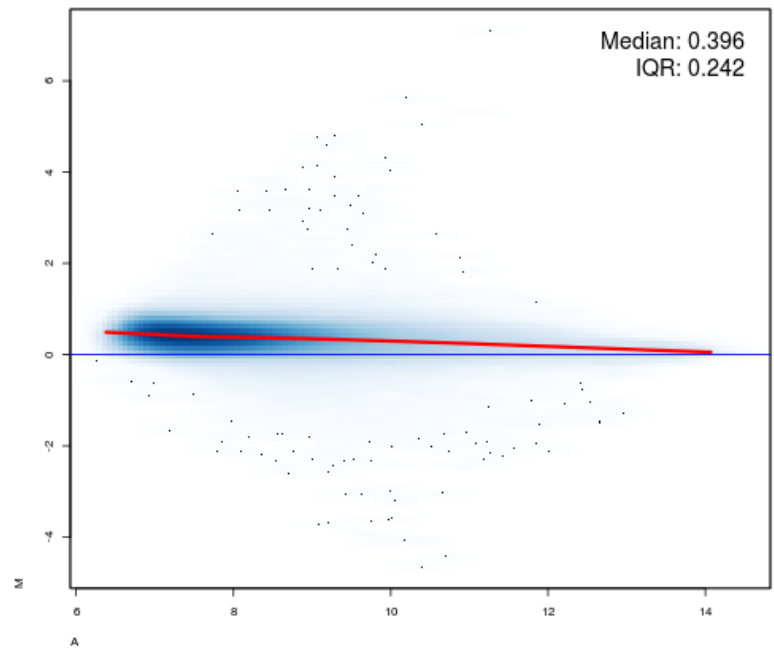


MA plots of raw data 1 / 2

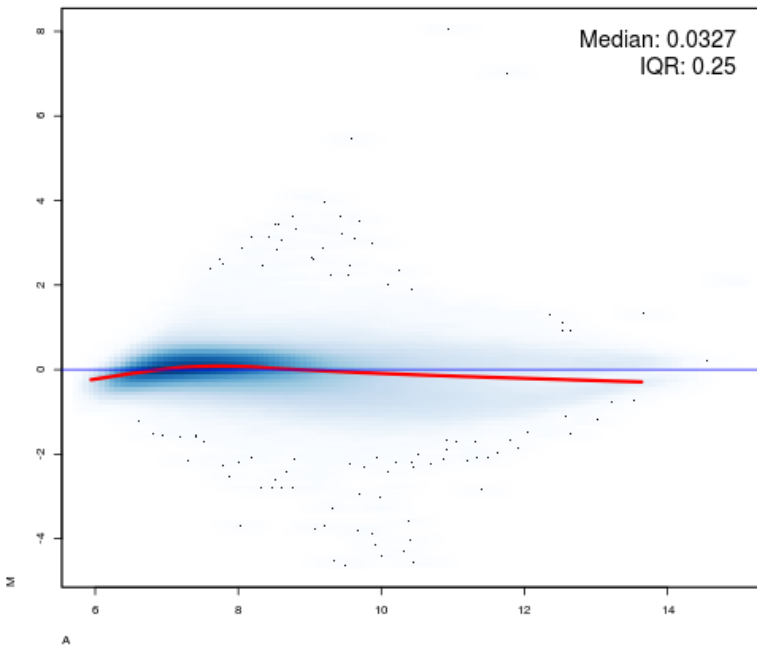
Array1 vs pseudo-median reference chip



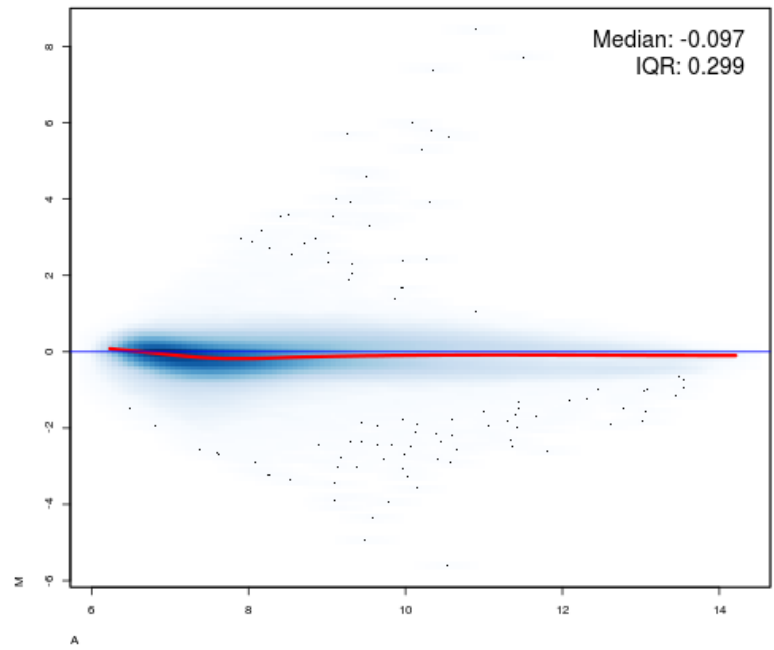
Array4 vs pseudo-median reference chip



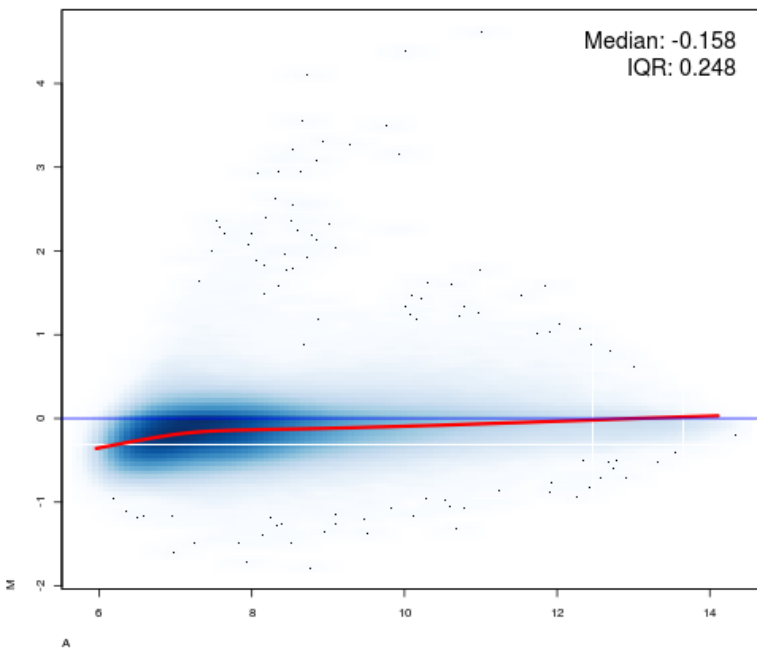
Array12 vs pseudo-median reference chip



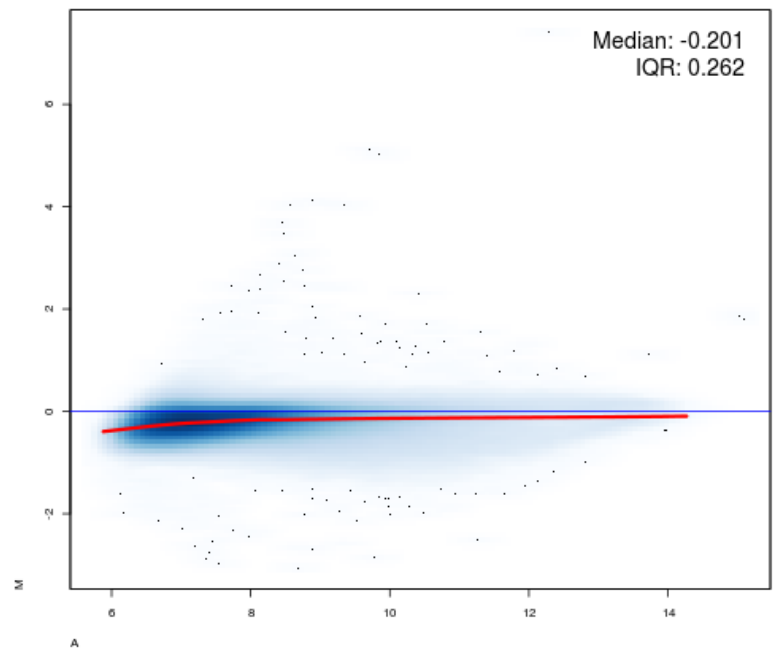
Array2 vs pseudo-median reference chip



Array5 vs pseudo-median reference chip

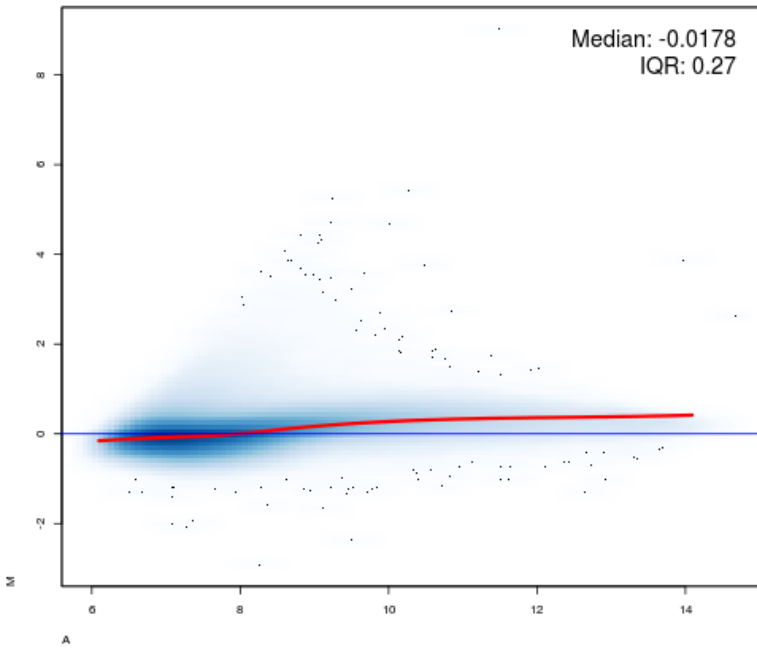


Array6 vs pseudo-median reference chip

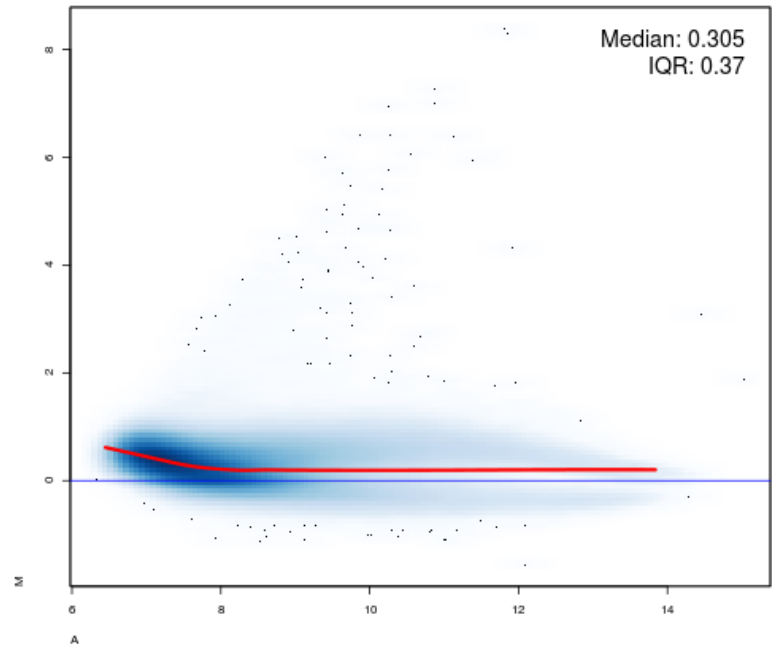


MA plots of raw data 2 / 2

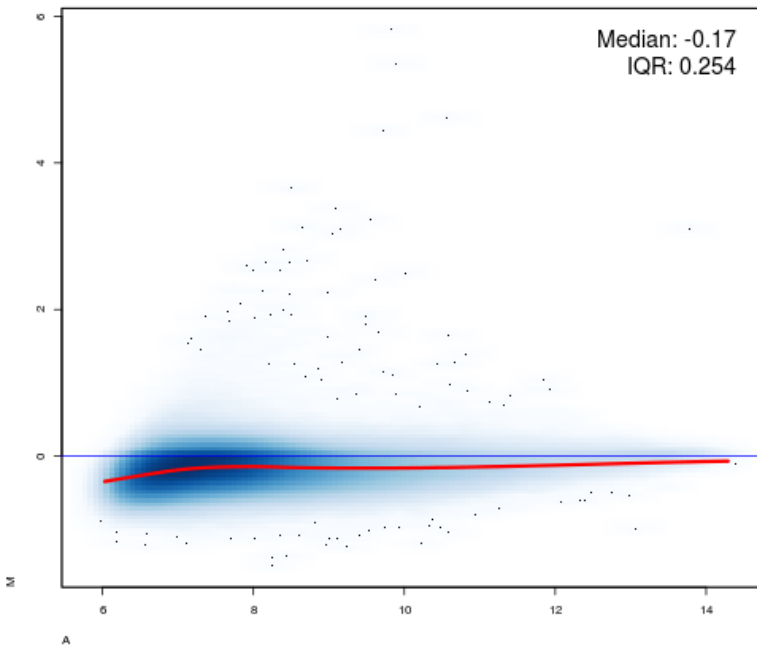
Array3 vs pseudo-median reference chip



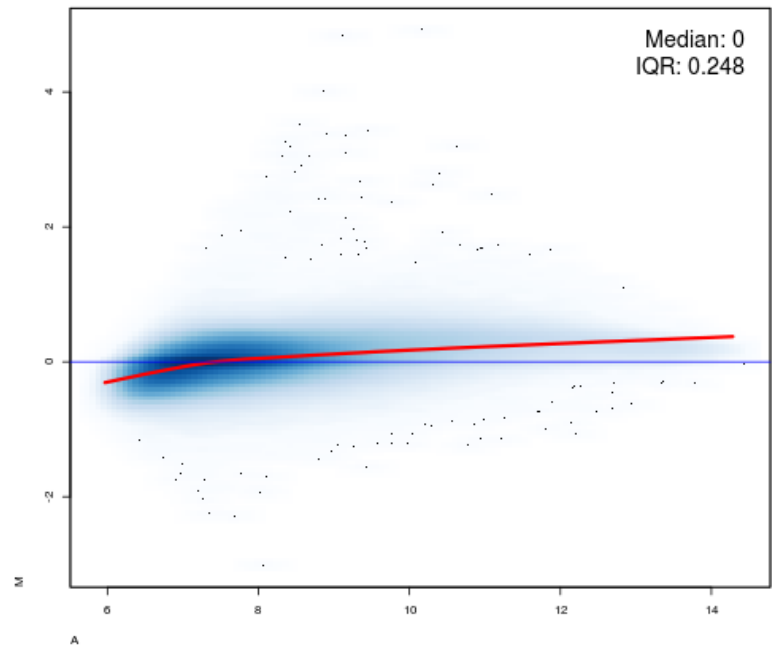
Array7 vs pseudo-median reference chip



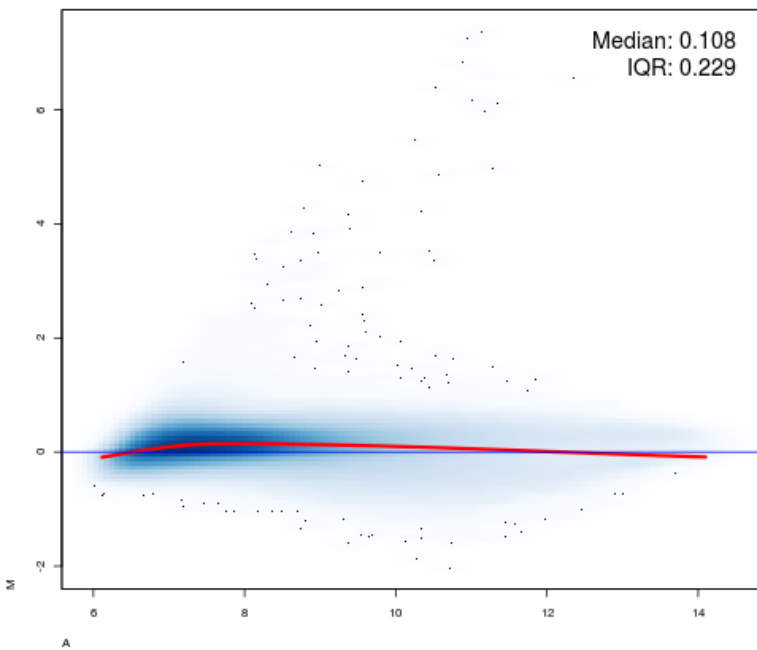
Array11 vs pseudo-median reference chip



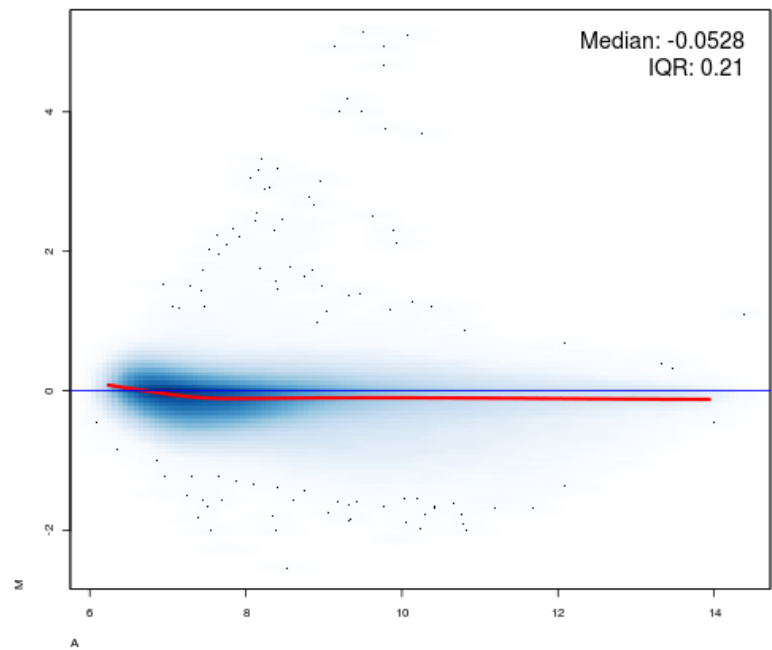
Array8 vs pseudo-median reference chip



Array9 vs pseudo-median reference chip

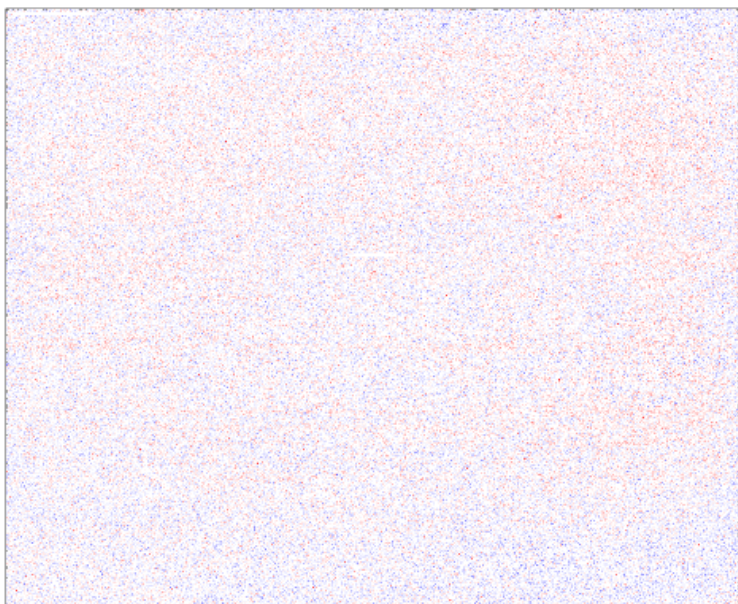


Array10 vs pseudo-median reference chip

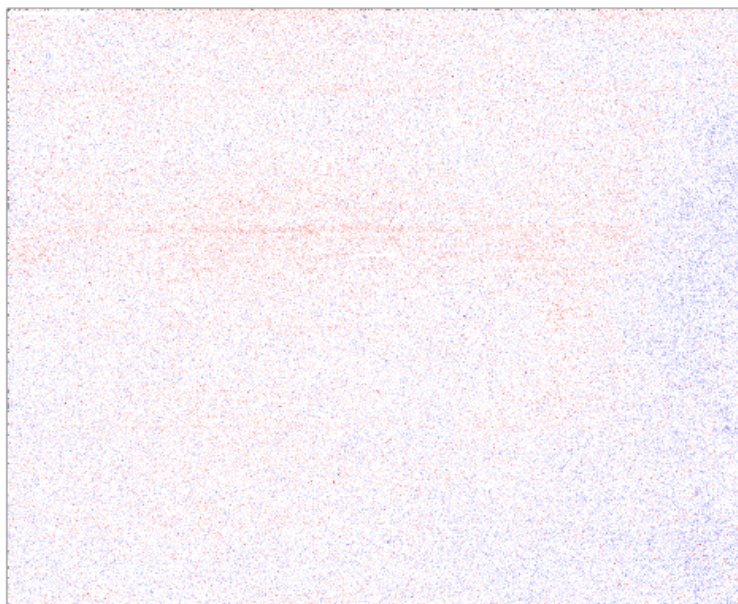


2D virtual PLM image for model characteristic: resid 1 / 2

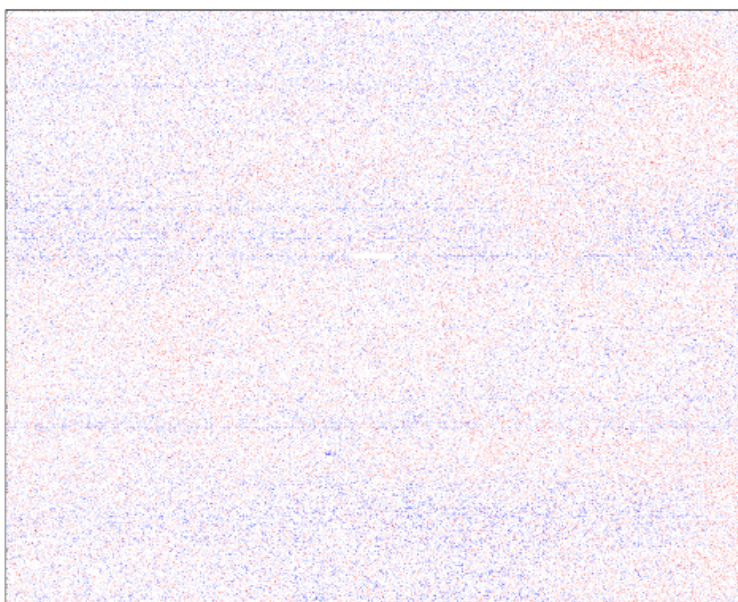
Array1



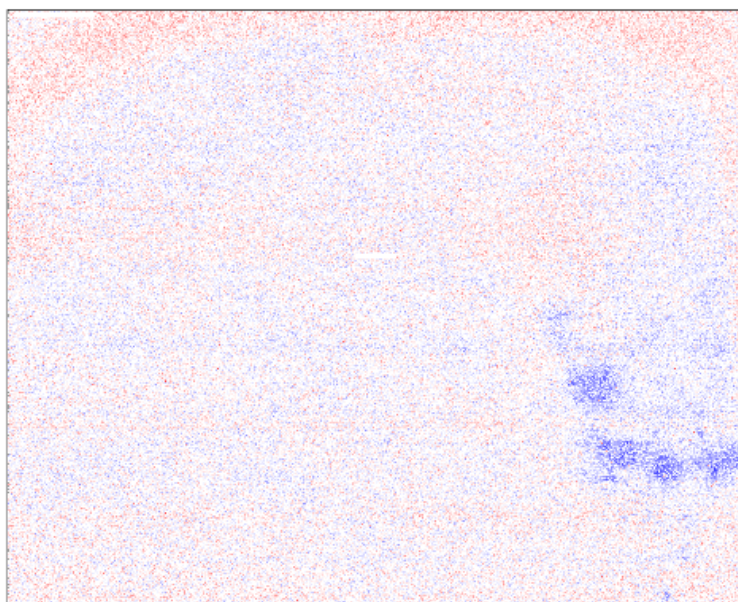
Array4



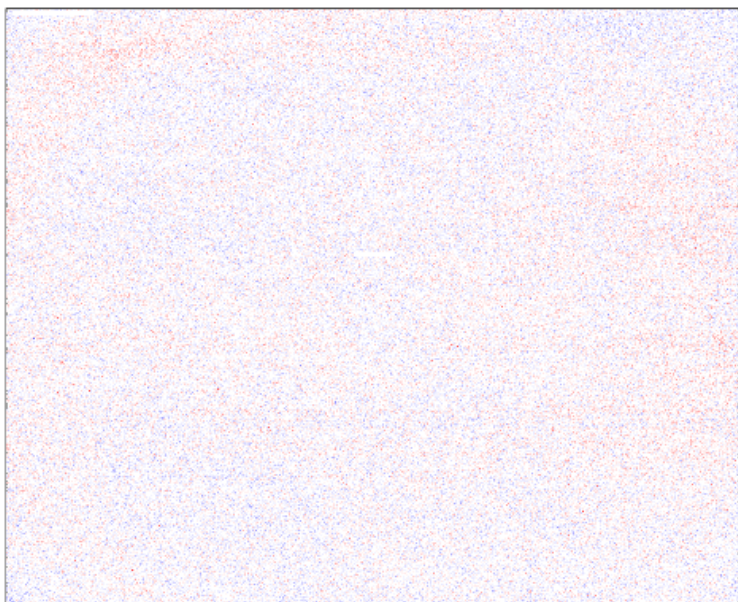
Array12



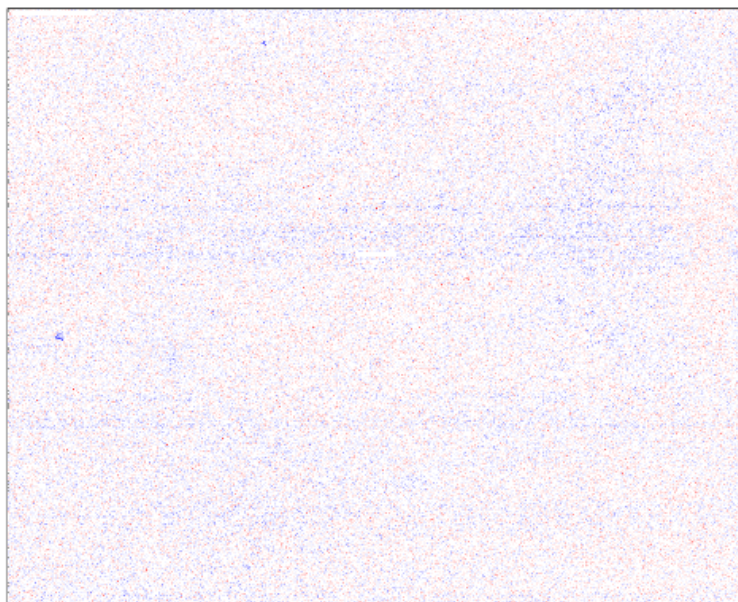
Array2



Array5

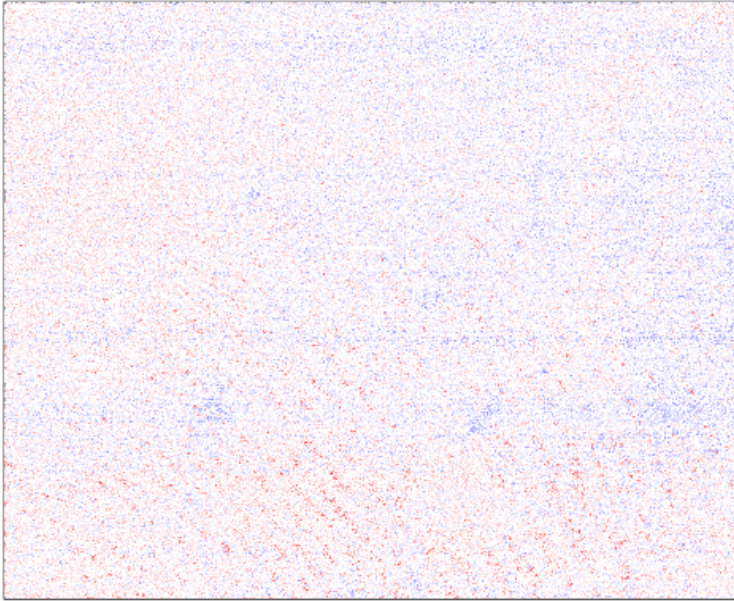


Array6

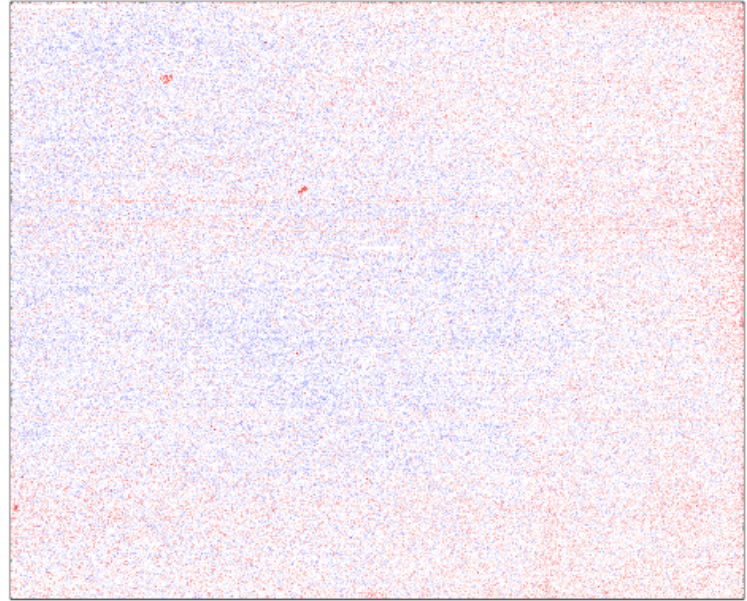


2D virtual PLM image for model characteristic: resids 2 / 2

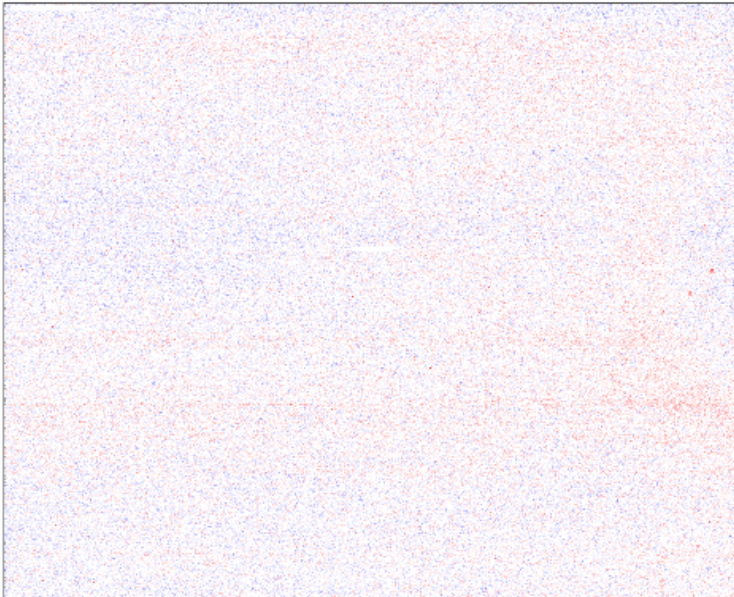
Array3



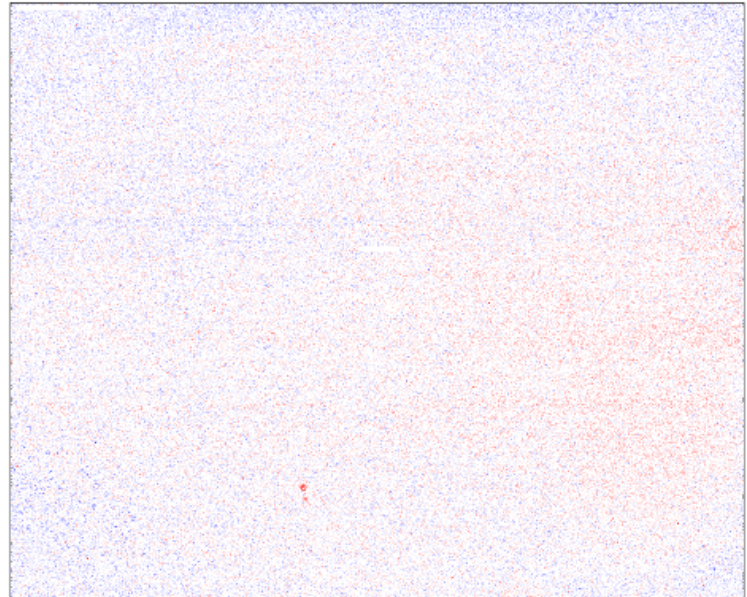
Array7



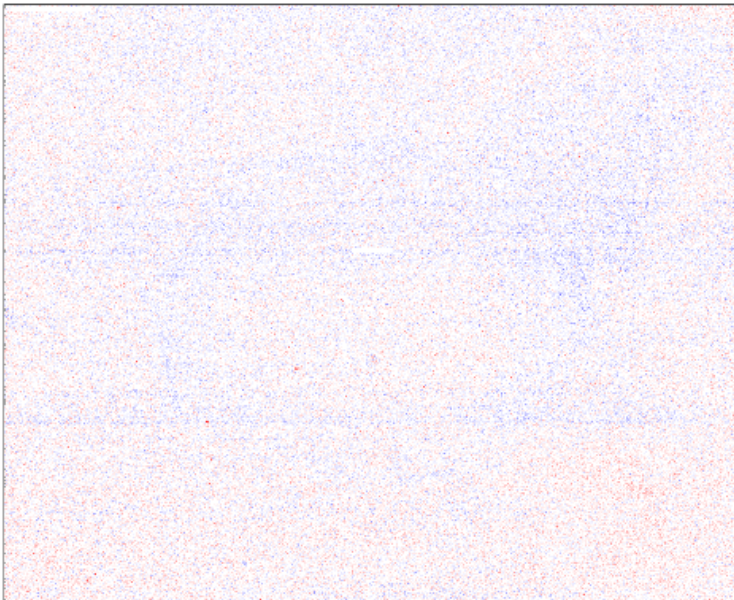
Array11



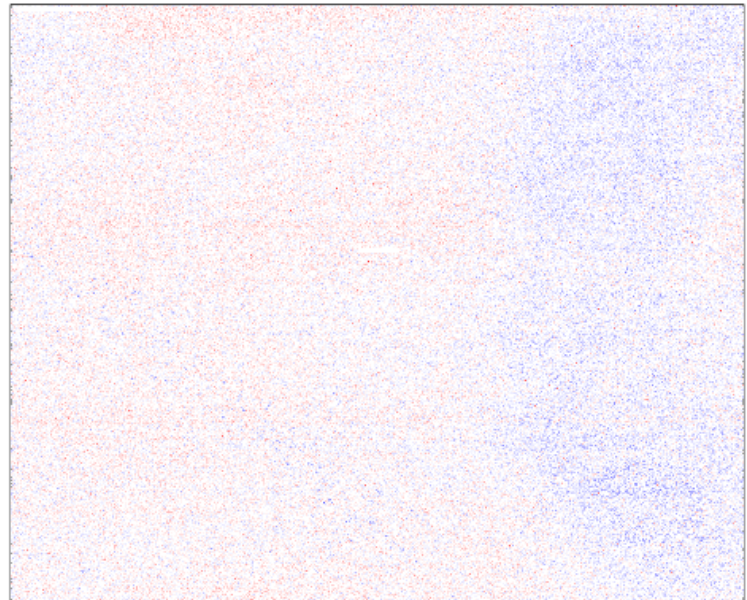
Array8



Array9

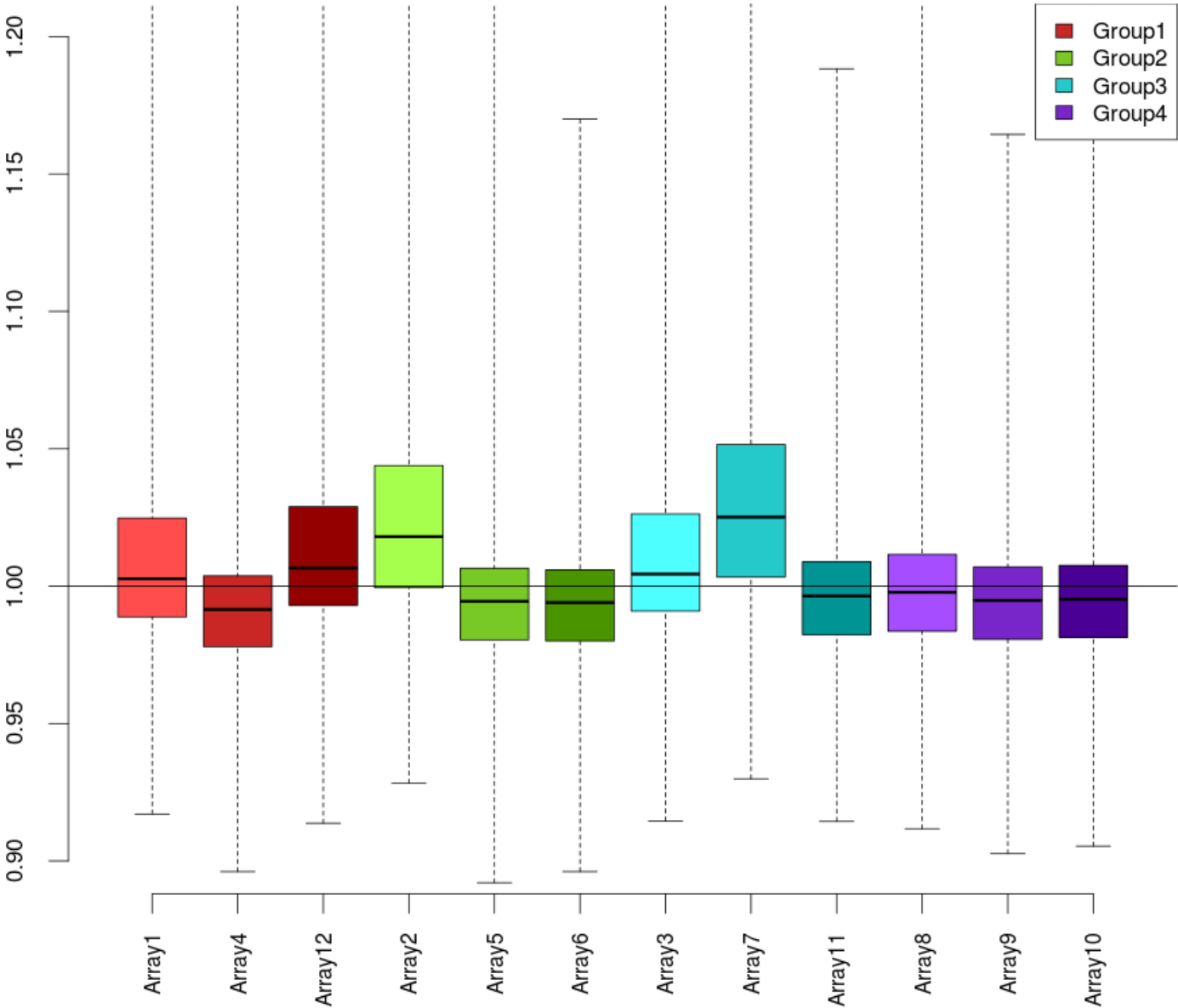


Array10



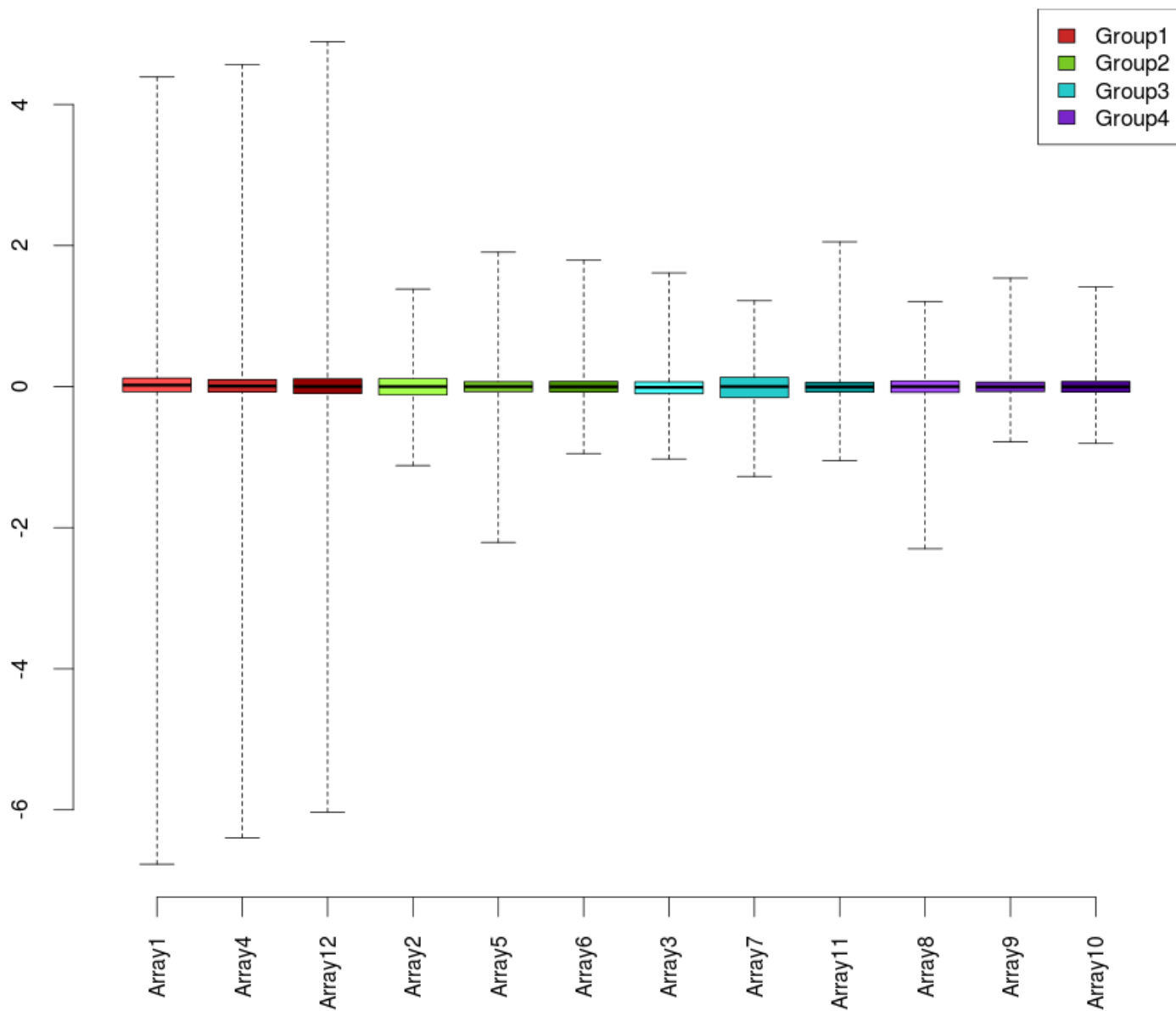
Normalized Unscaled Standard Errors (NUSE)

NUSE median value should be < 1.1

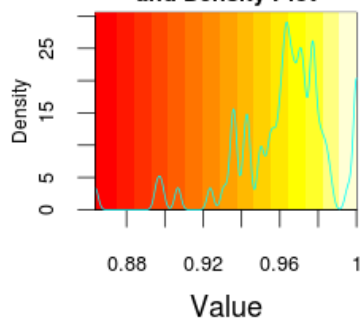


Relative Log Expression (RLE)

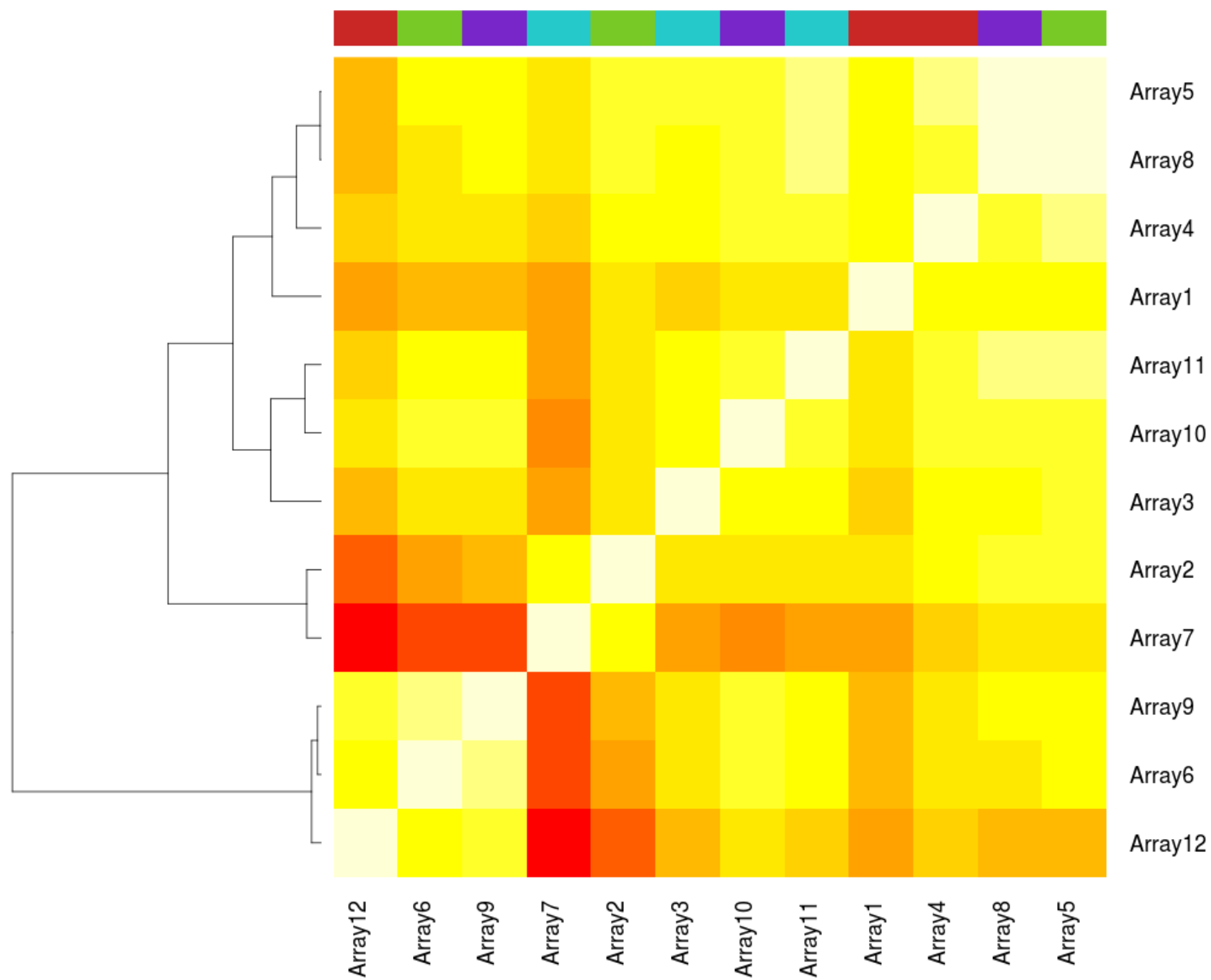
RLE distributions should be centered around 0



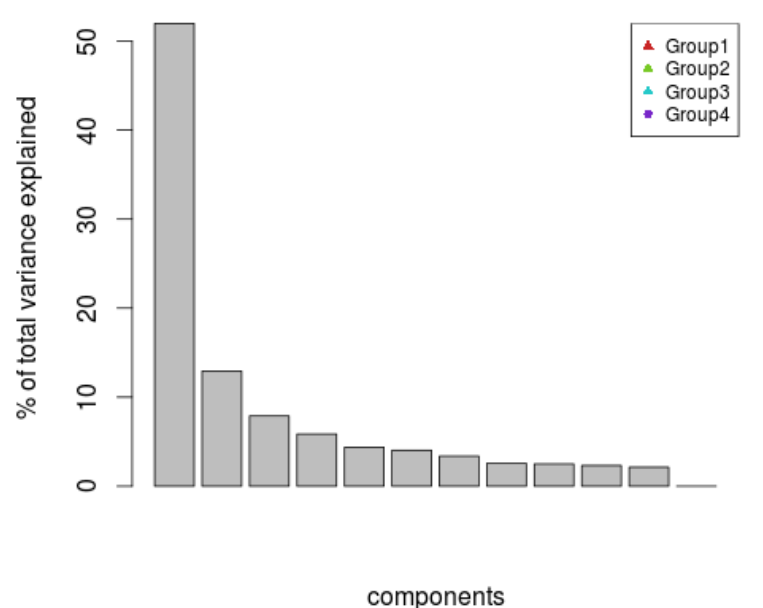
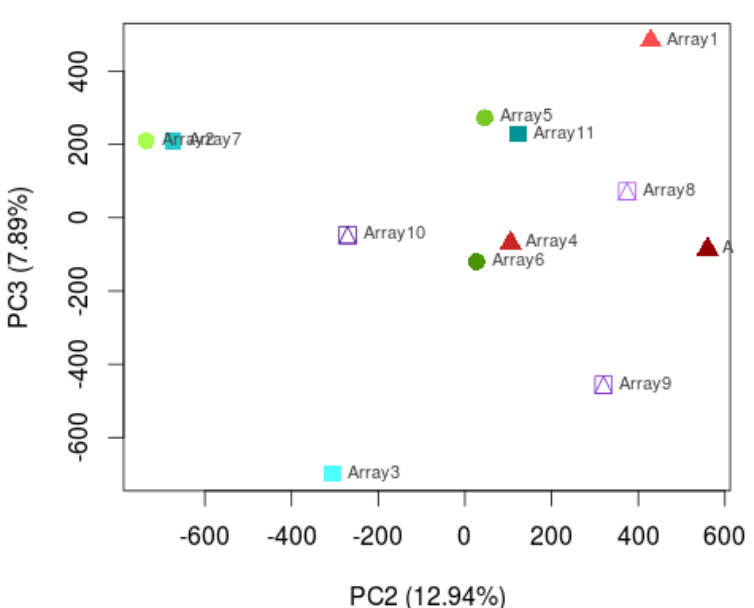
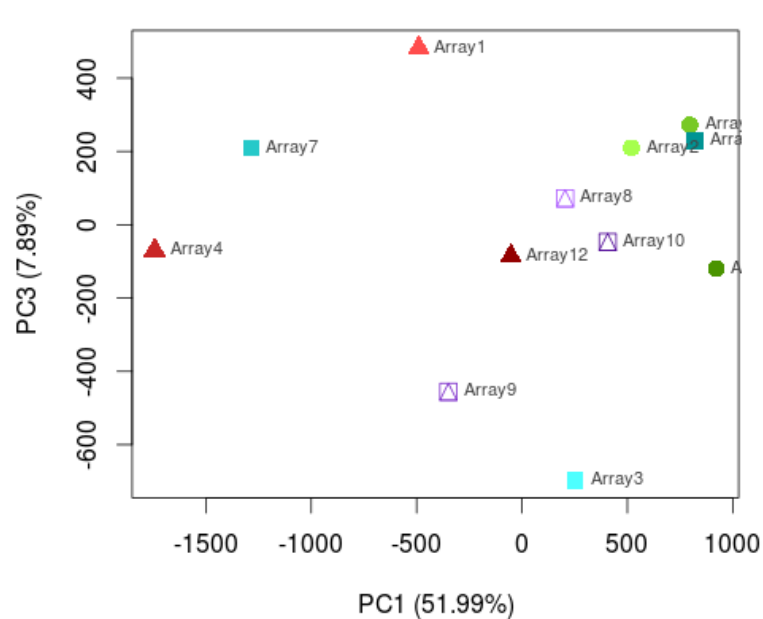
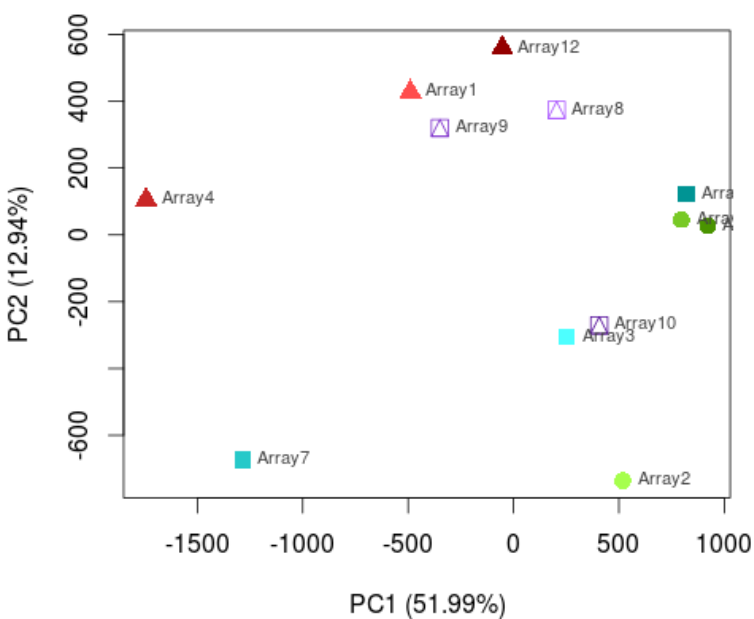
Color Key
and Density Plot



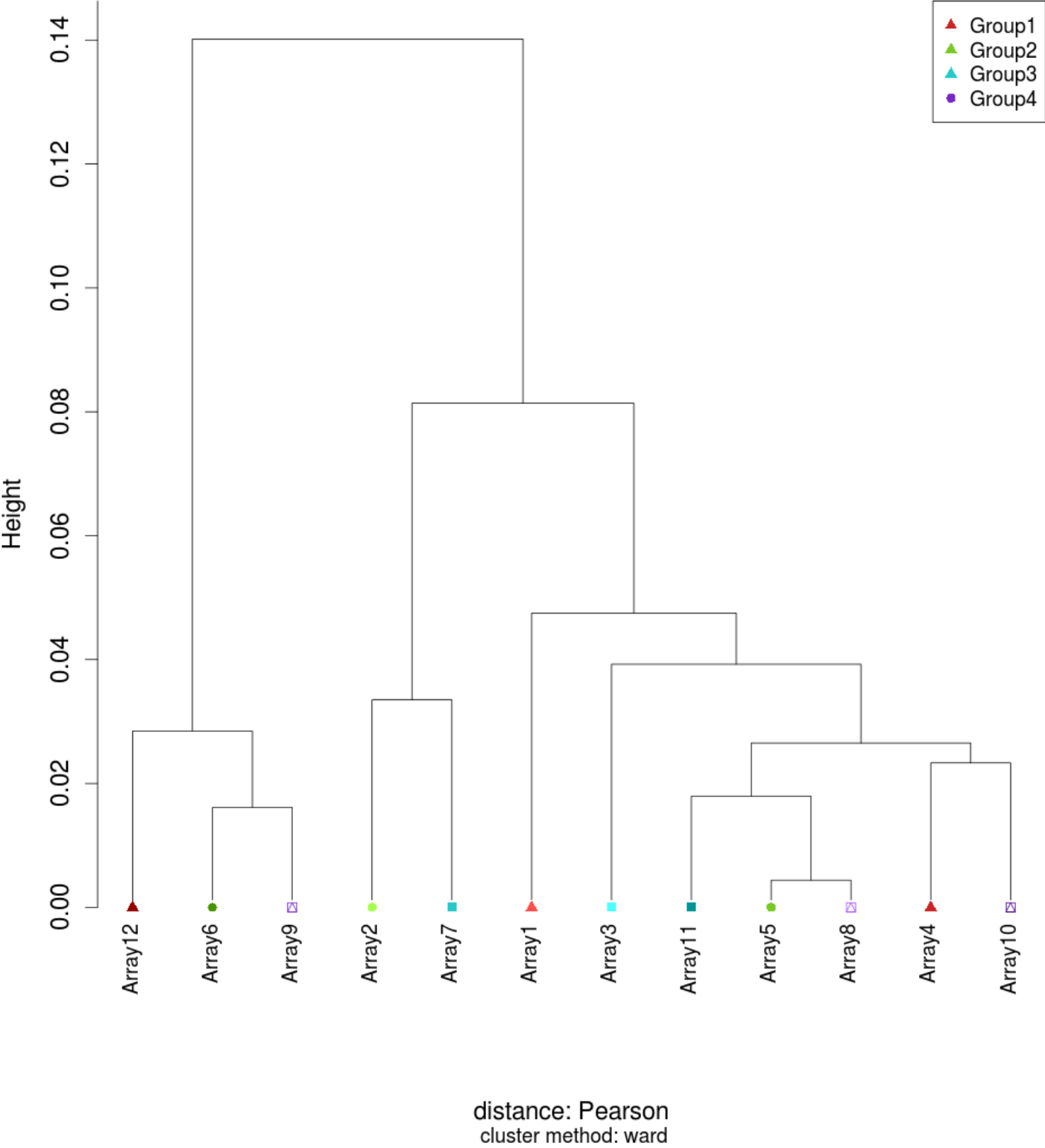
Raw data correlation plot
correlation method: pearson
cluster method: ward



PCA analysis of Raw data



Cluster dendrogram of raw data



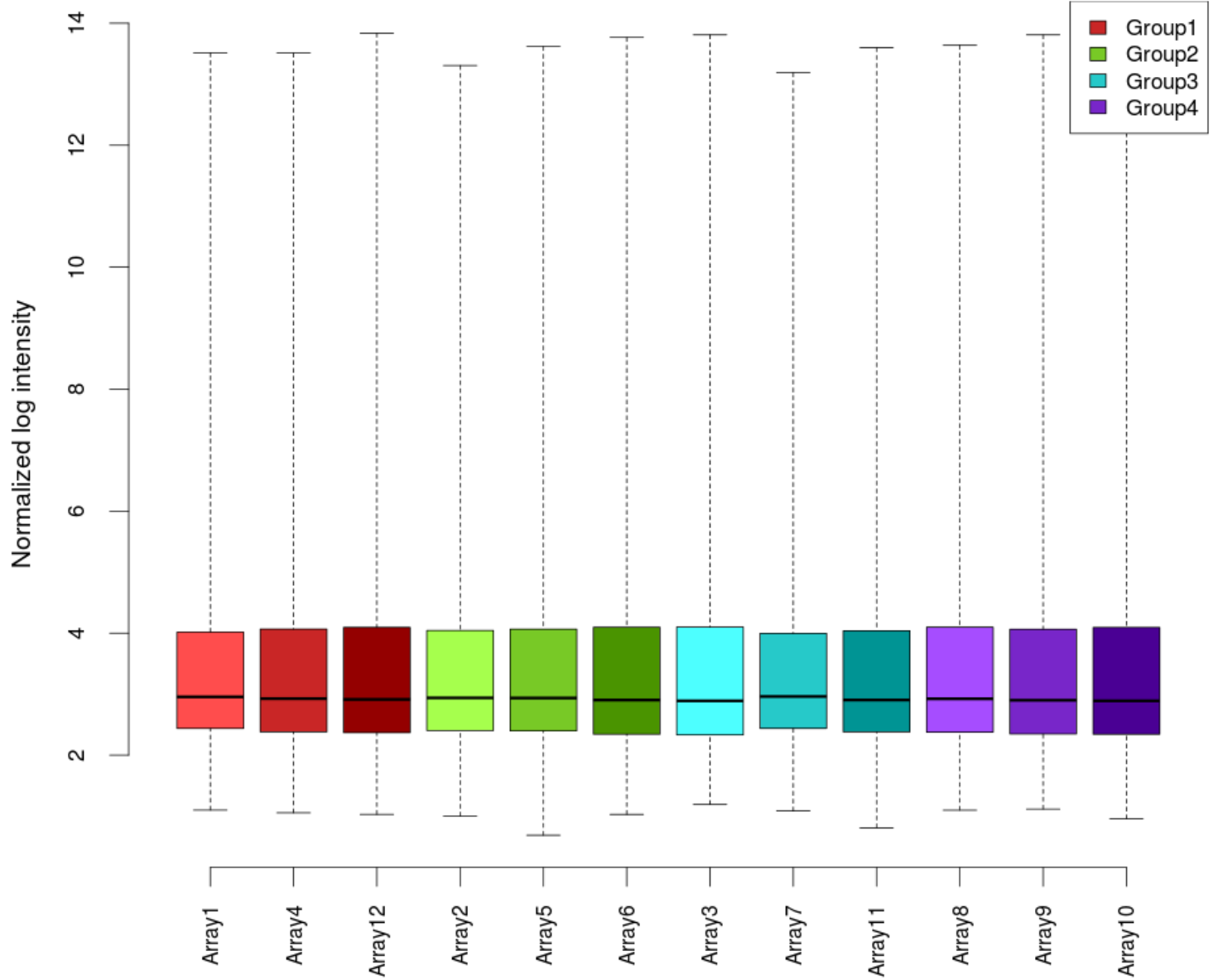
Pre-processing of Raw Data

Method: GCRMA

Annotation: hgu133plus2_Hs_ENSG

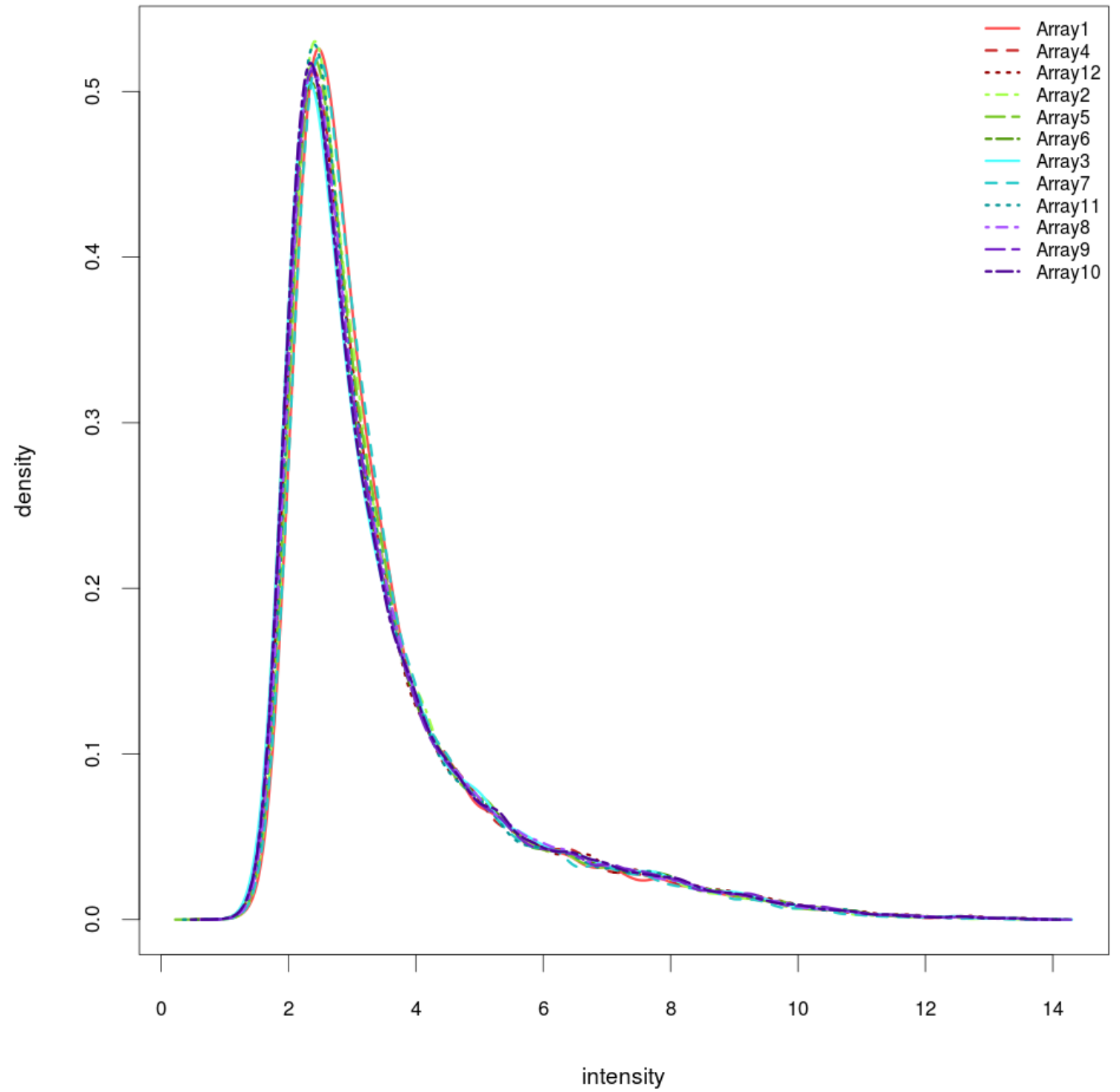
Boxplot after GCRMA

Distributions should be comparable between arrays



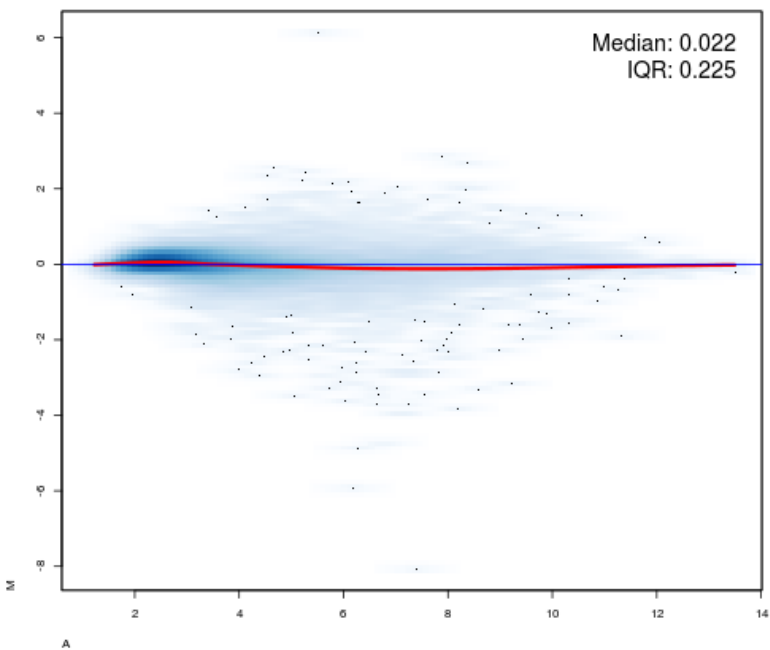
Density histogram after GCRMA

Curves should be comparable between arrays

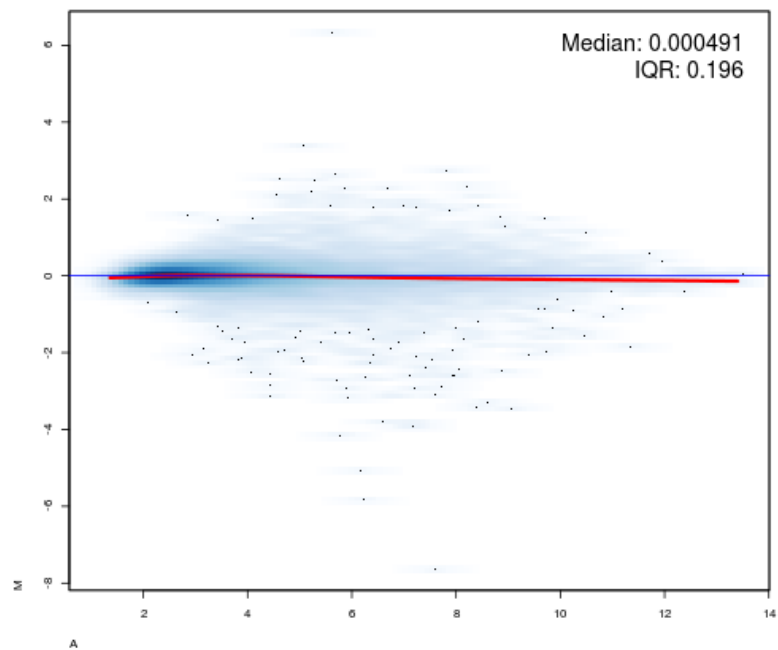


MA plots after GCRMA normalization 1 / 2

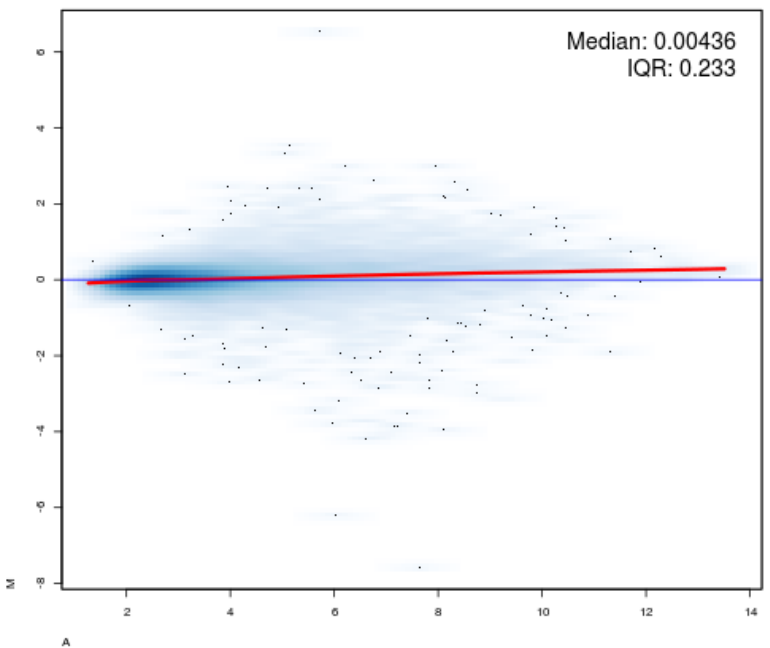
Array1 vs pseudo-median reference chip



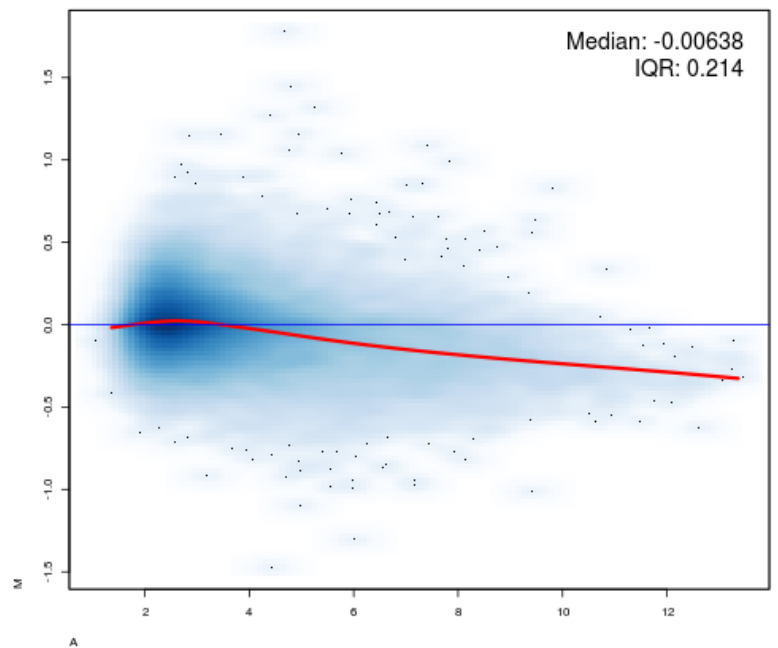
Array4 vs pseudo-median reference chip



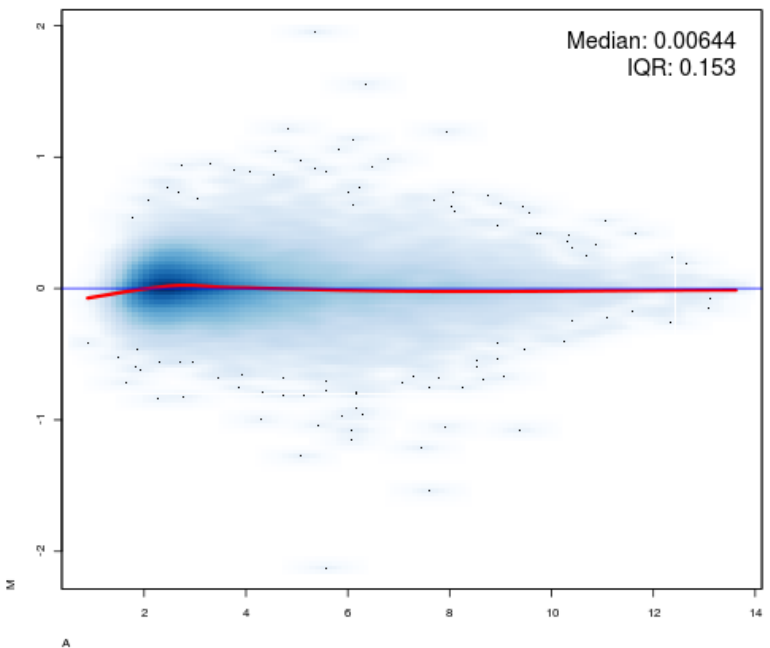
Array12 vs pseudo-median reference chip



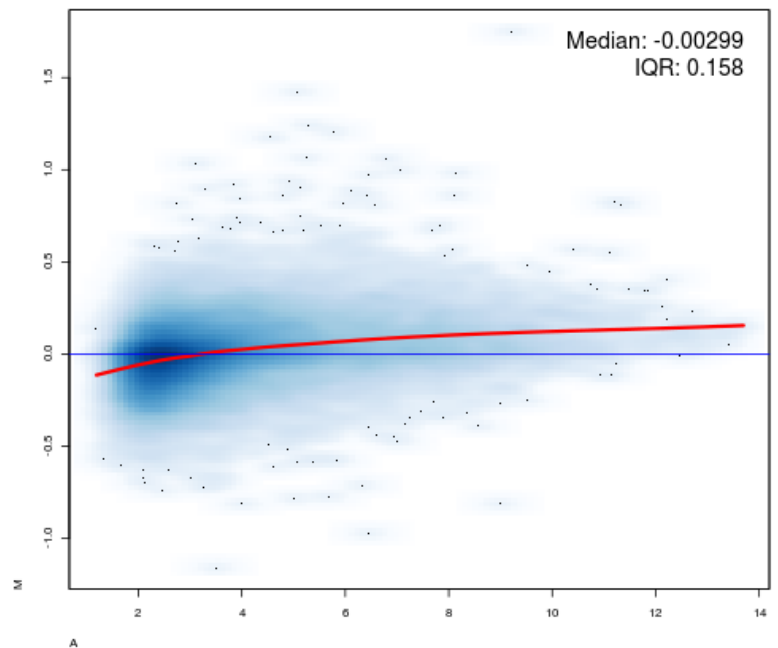
Array2 vs pseudo-median reference chip



Array5 vs pseudo-median reference chip

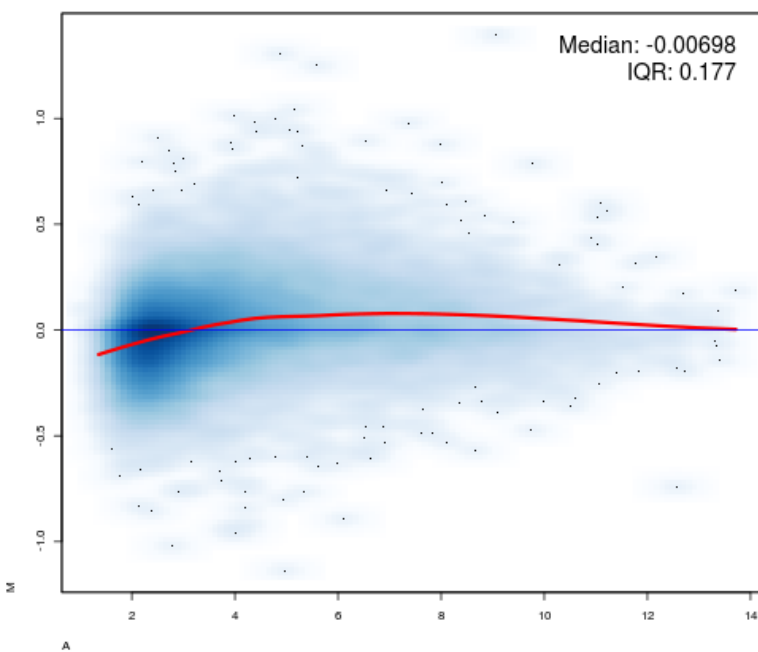


Array6 vs pseudo-median reference chip

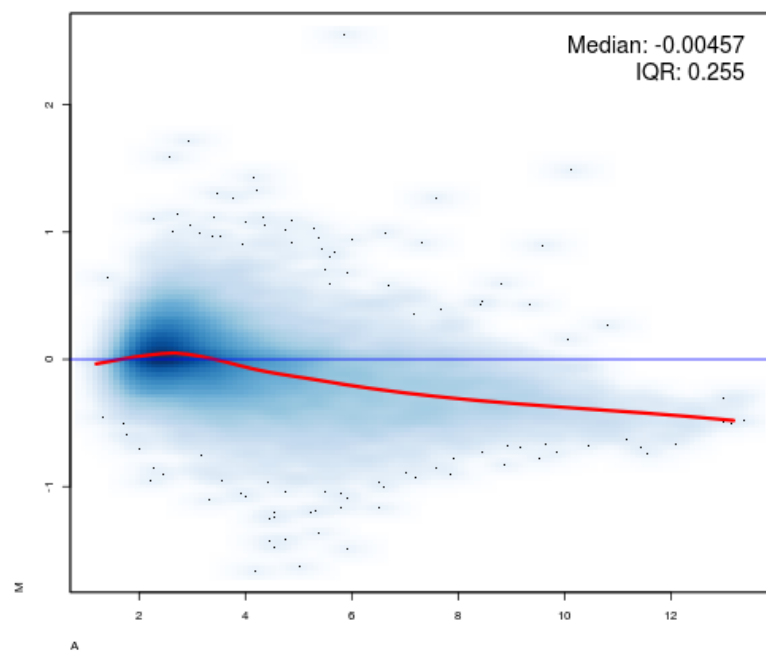


MA plots after GCRMA normalization 2 / 2

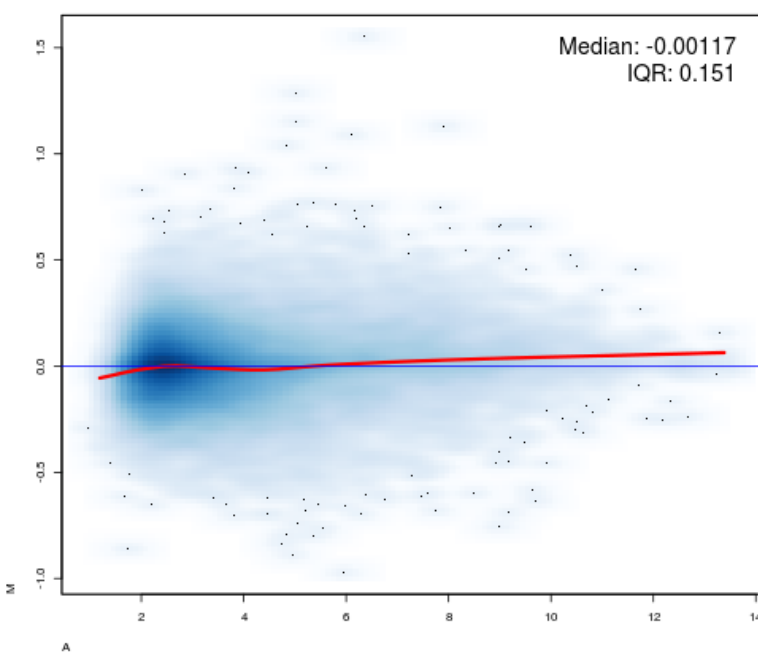
Array3 vs pseudo-median reference chip



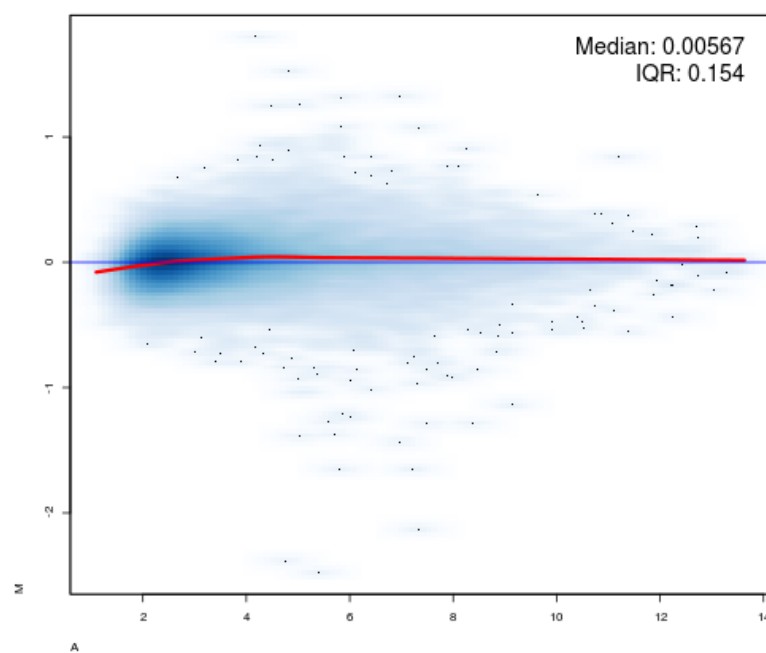
Array7 vs pseudo-median reference chip



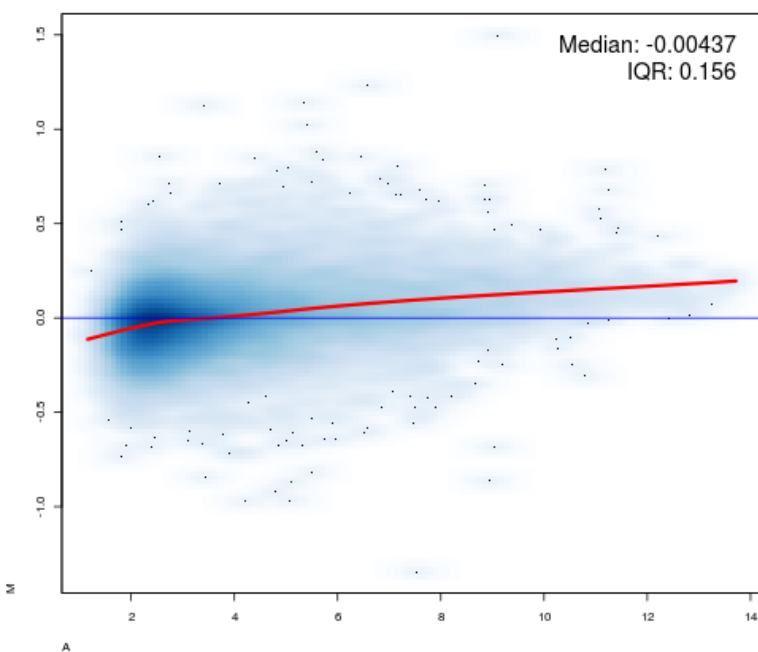
Array11 vs pseudo-median reference chip



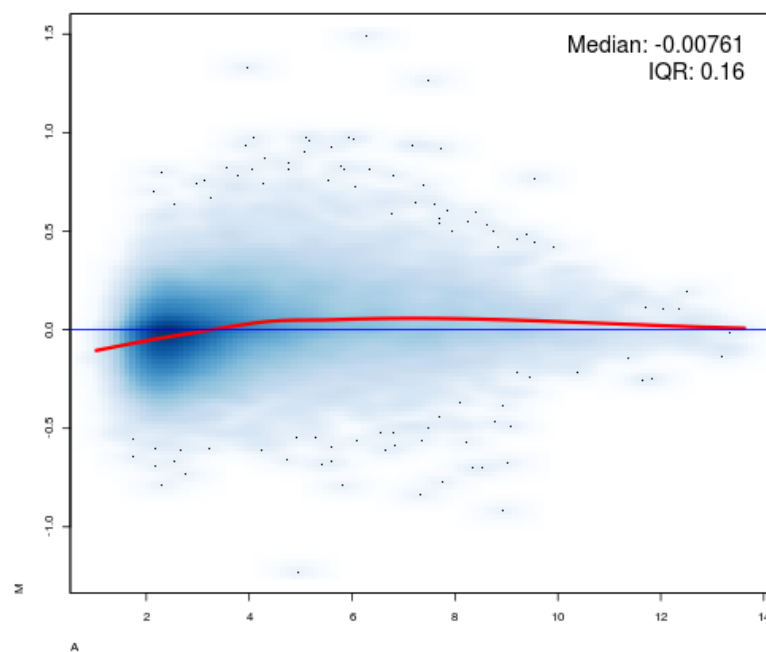
Array8 vs pseudo-median reference chip



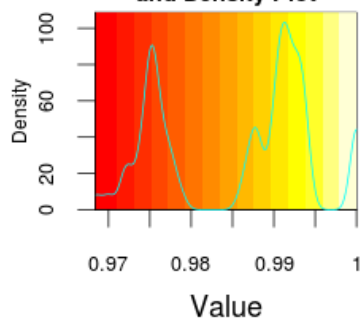
Array9 vs pseudo-median reference chip



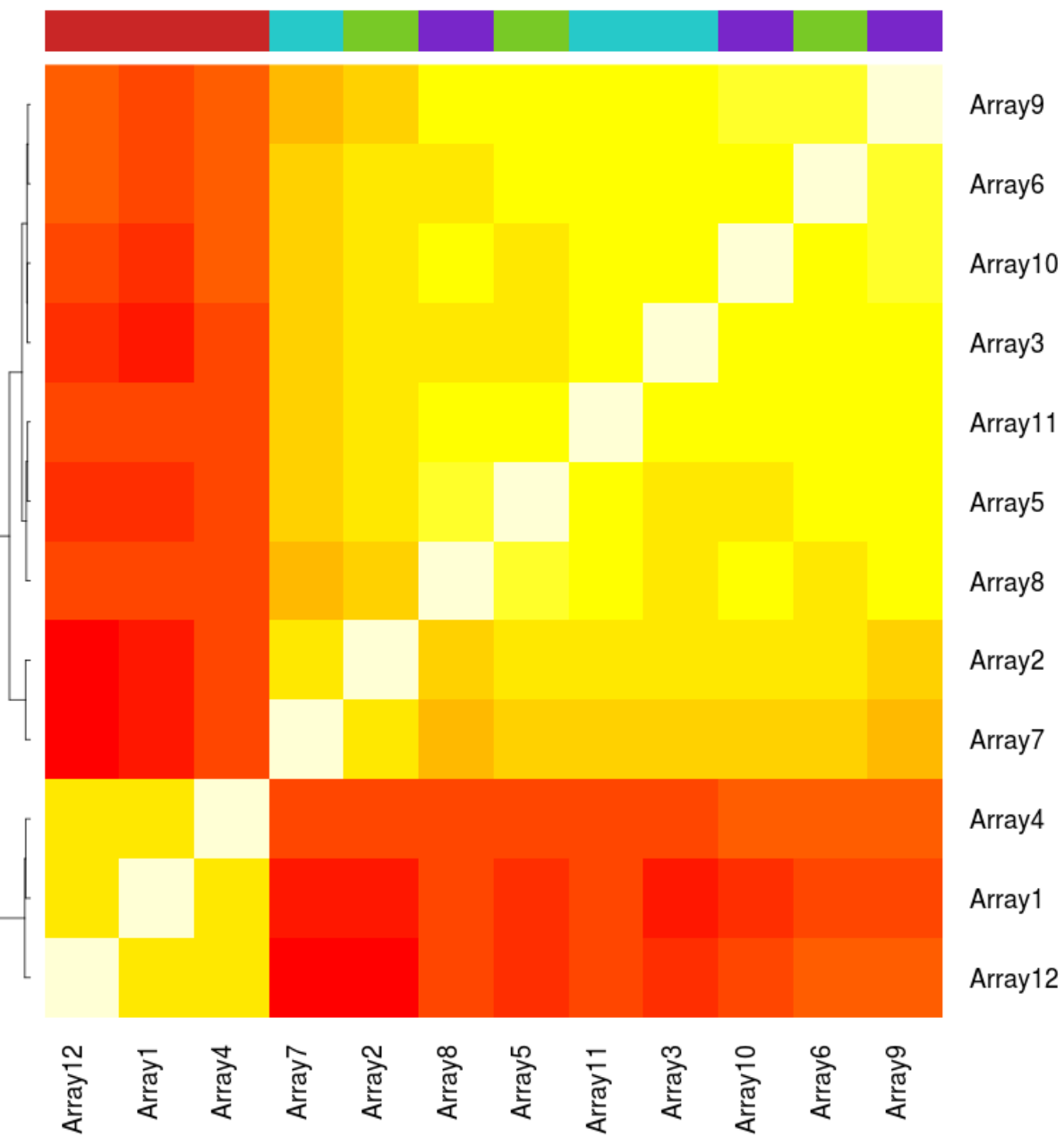
Array10 vs pseudo-median reference chip



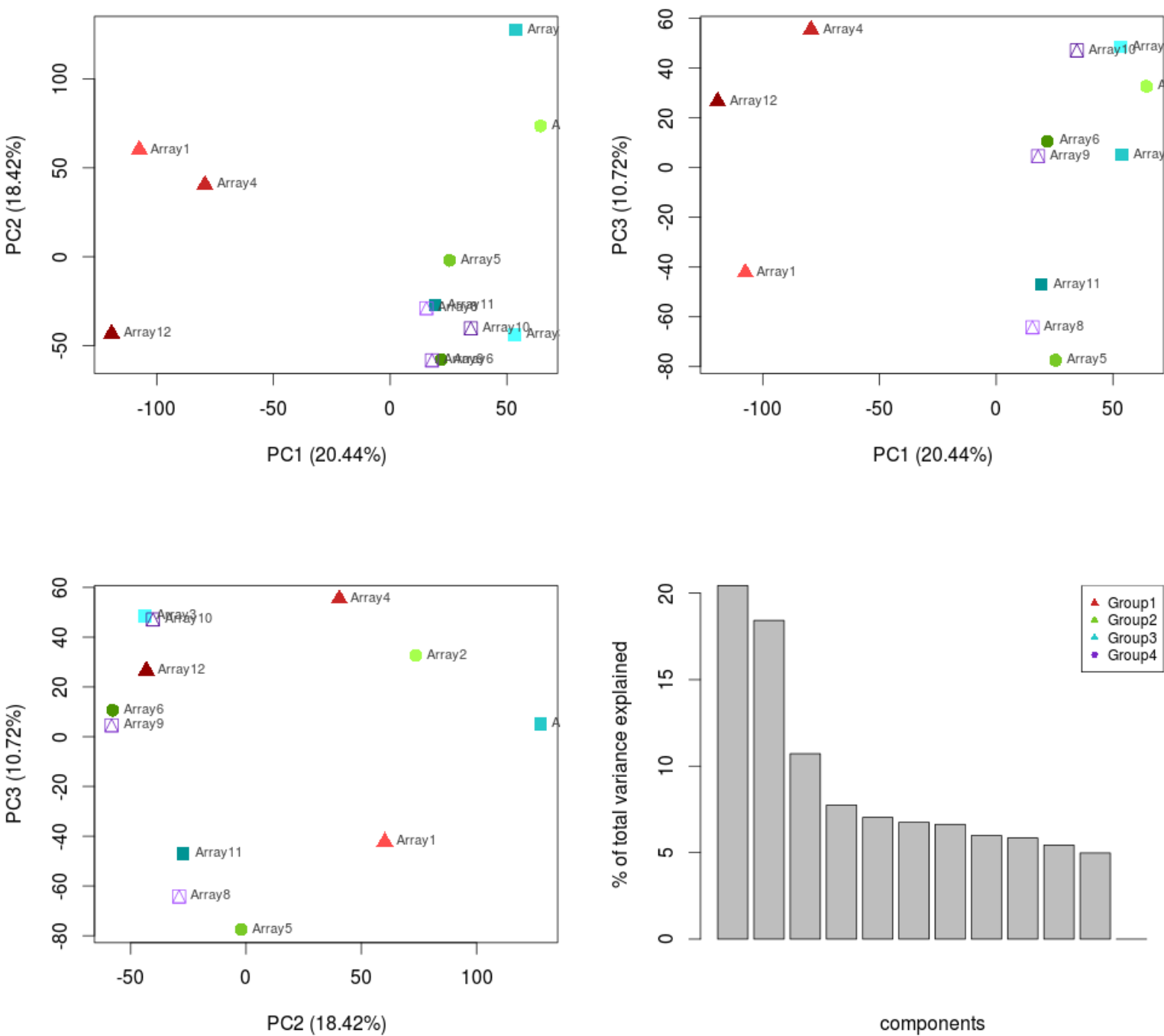
Color Key
and Density Plot



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



PCA analysis after GCRMA normalization



Cluster dendrogram of GCRMA normalized data

