

#### Array names and grouping

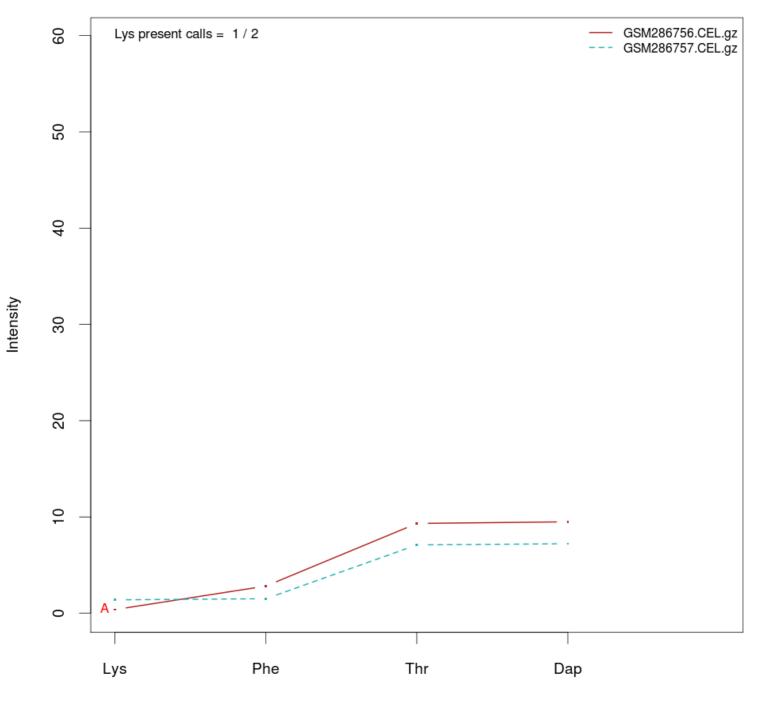
ArtayDataFlie	SourceName	Factor/Value
GSM288766.CEL.gz	GSV256758 CEL gz	1
GSM288767.CEL.gz	GSV296757.CEL.gz	1

#### Array names and grouping

ArrayDataFile	SourceName	FactorValue	
GSM286756.CEL.gz	GSM286756.CEL.gz	1	
GSM286757.CEL.gz	GSM286757.CEL.gz	1	



## Spike-in Sample Prep controls intensities and calls



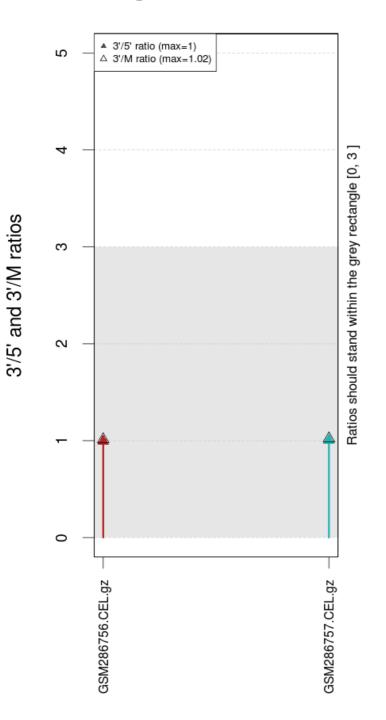
Intensity: OK (Lys < Phe < Thr < Dap for all arrays)
Lys Present calls: 1 Lys not called present

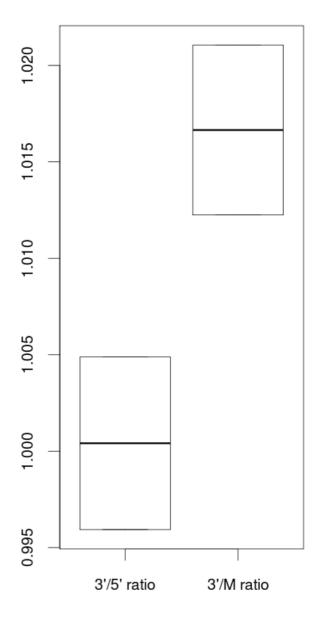
## Summary of raw data quality indicators



## RNA degradation of beta-actin

#### **Boxplot of beta-actin ratios**

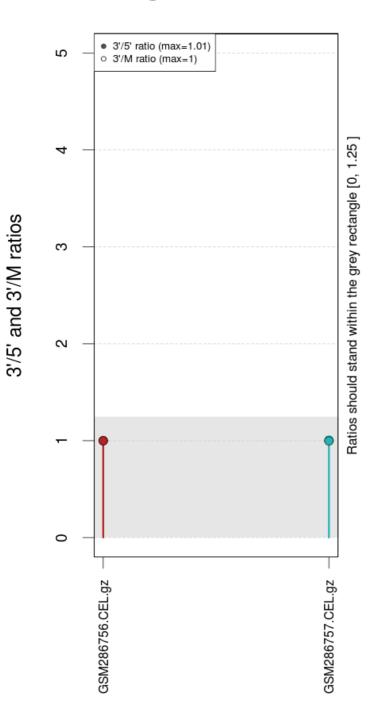


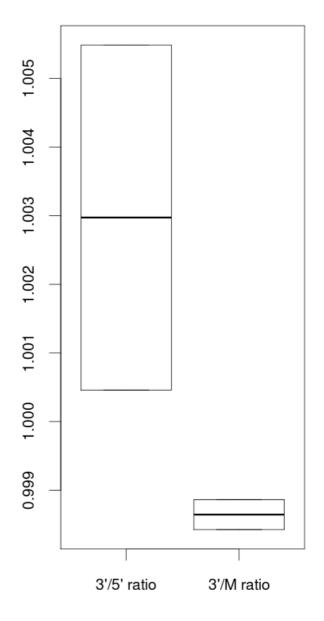


beta-actin QC: OK (all 3'/5' ratios < 3)

#### **RNA degradation of GAPDH**

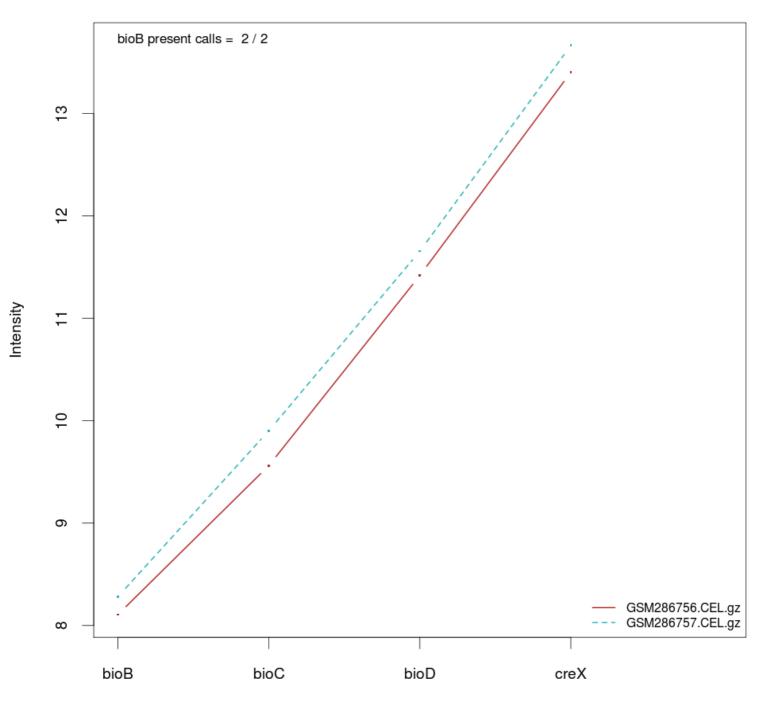
#### **Boxplot of GAPDH ratios**





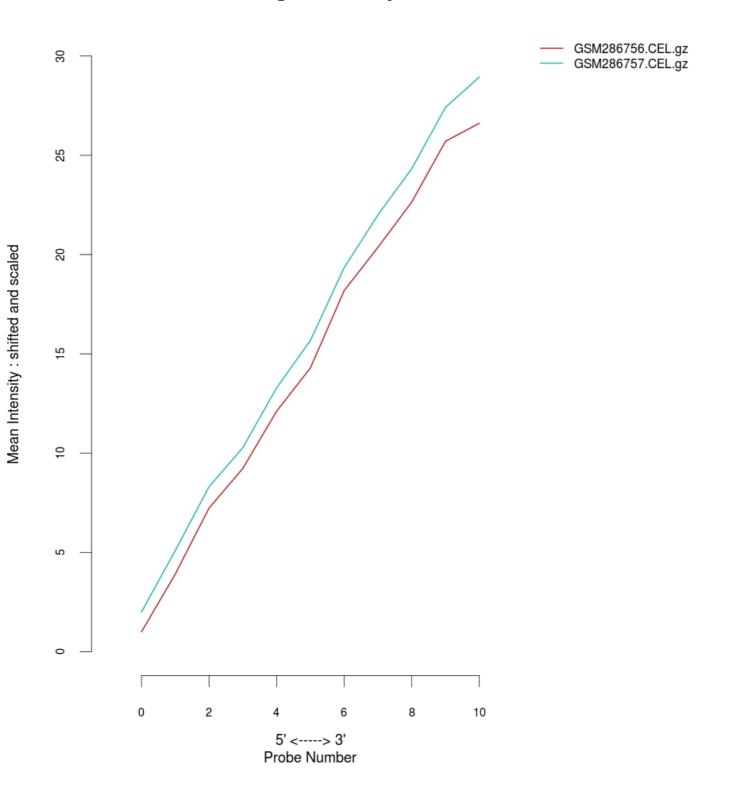
GAPDH QC: OK (all 3'/5' ratios < 1.25)

## 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)

# **RNA** degradation plot



# Plot of percent present

100

80

9

40

20

0

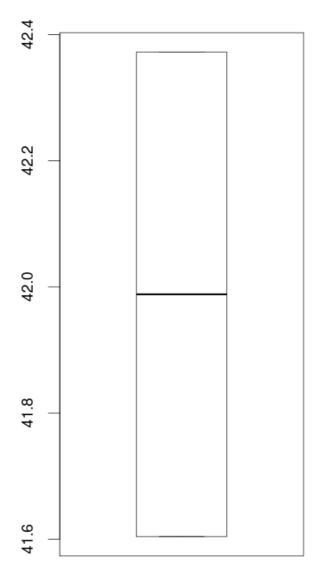
GSM286756.CEL.gz

Data should be in the grey rectangle representing a spread of 10%

min = 41.6 max = 42.37 max-min = 0.77

GSM286757.CEL.gz

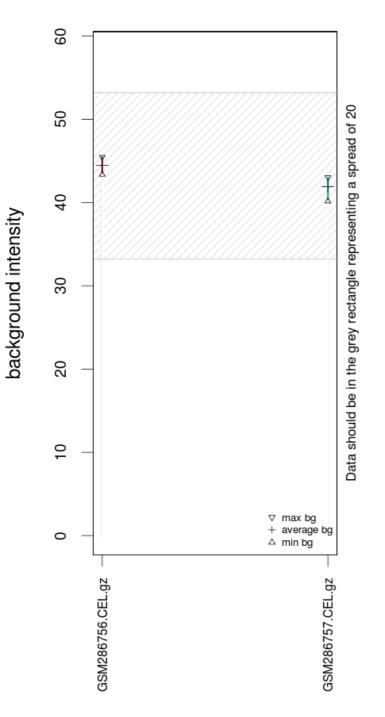
## **Boxplot of percent present**

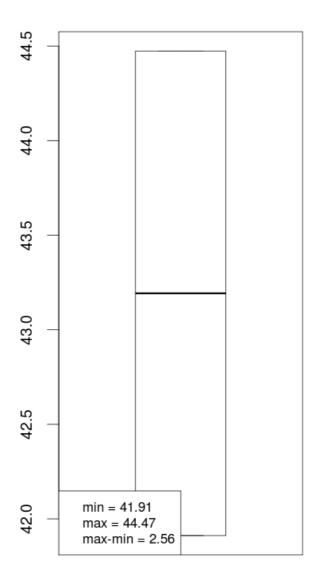


Percent present QC: OK (spread <= 10%)

## Plot of background intensity

## itensity Average background intensity



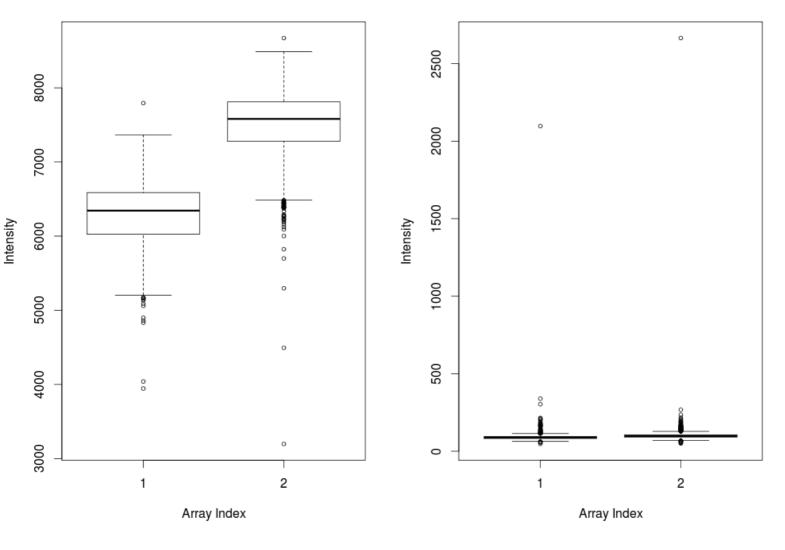


Background QC: OK (spread <= 20)

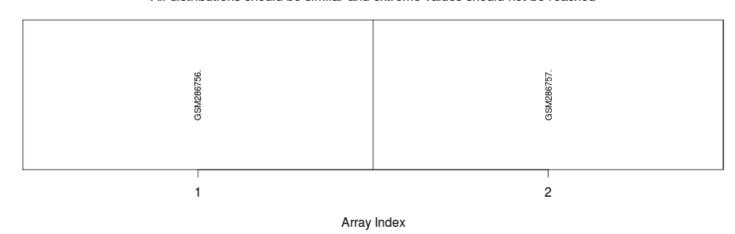


Sample Name

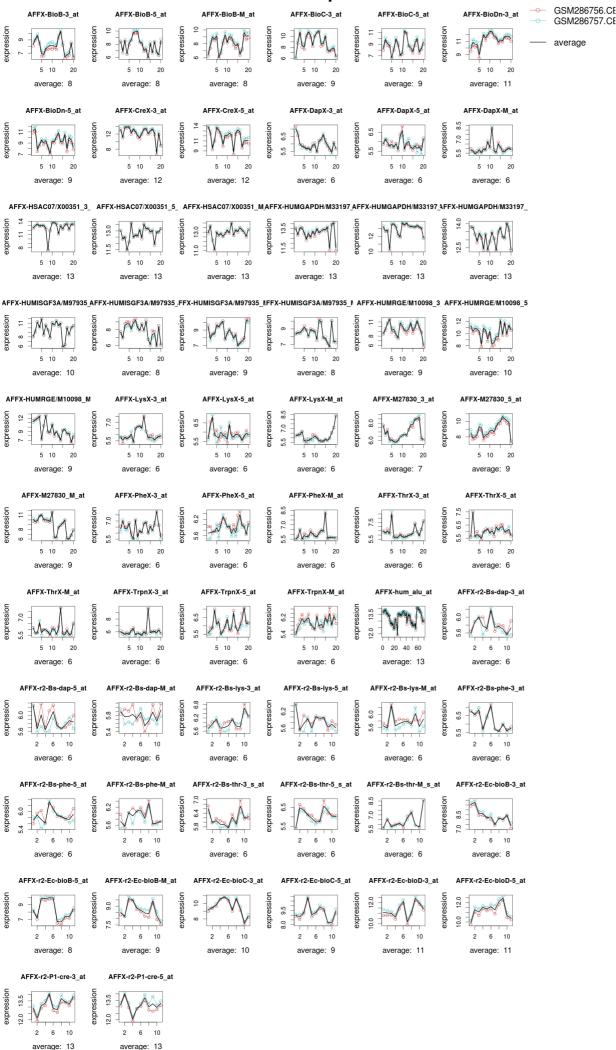
#### **Negative Border Elements**



All distributions should be similar and extreme values should not be reached

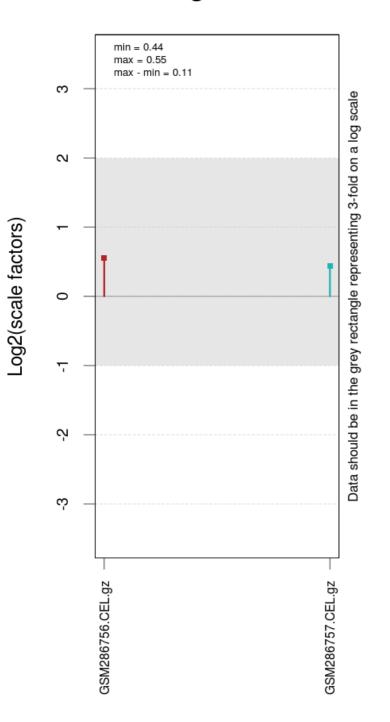


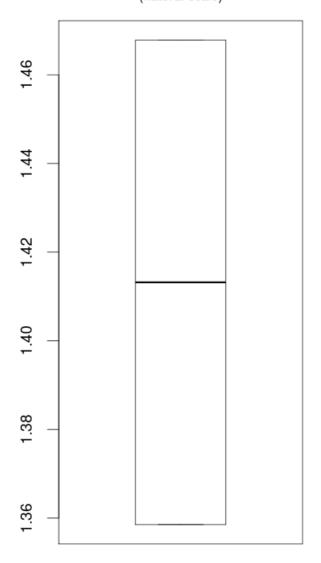
#### affx control profiles



## Plot of Log scale factors

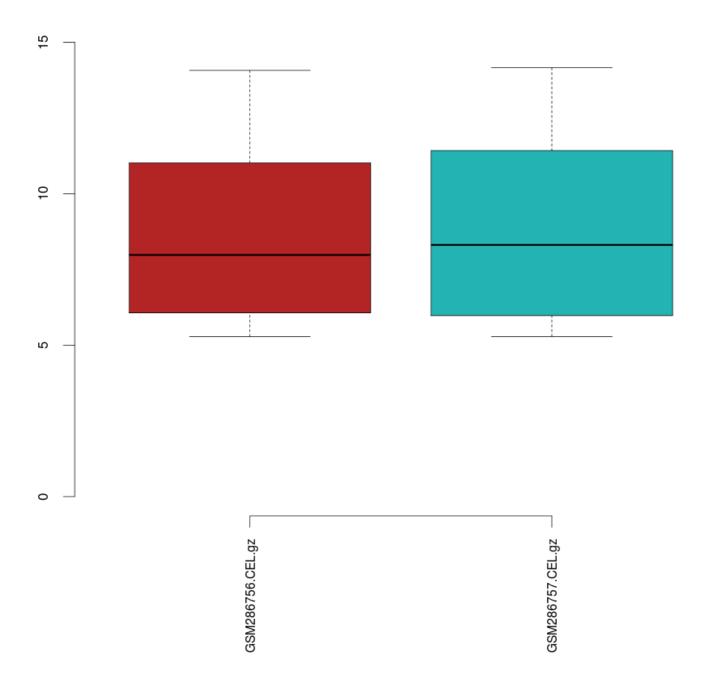
# Boxplot of scale factors (natural scale)



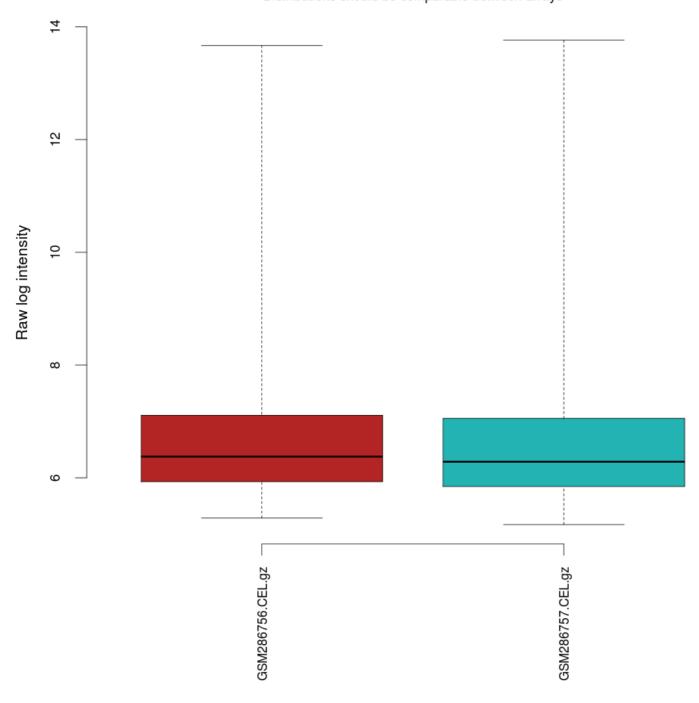


Scale factors QC: OK (spread < 3-fold)

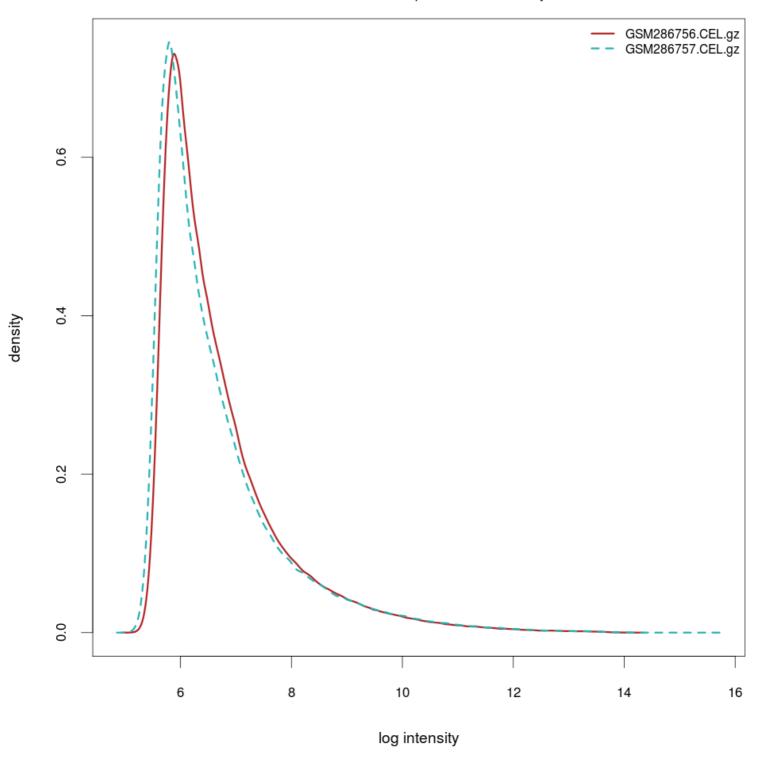
# affx controls



# Boxplot of raw intensities Distributions should be comparable between arrays



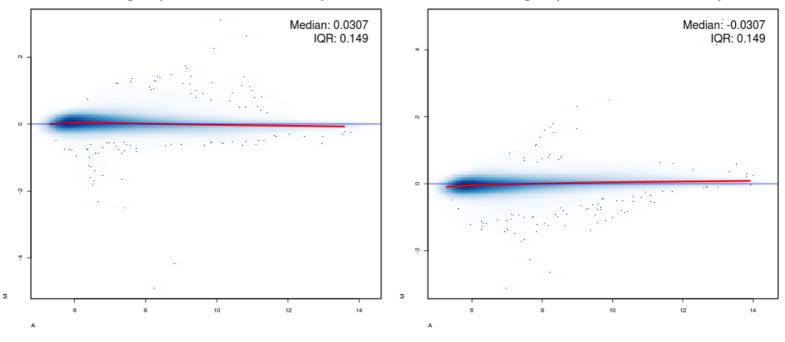
# Density histogram of raw intensities Curves should be comparable between arrays



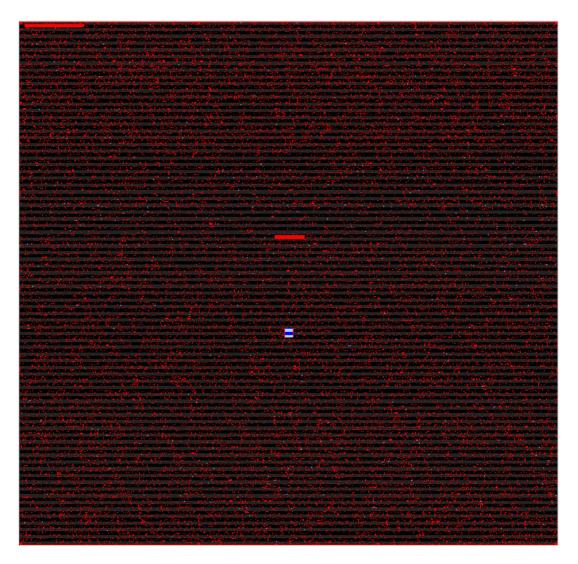
#### MA plots of raw data



#### GSM286757.CEL.gz vs pseudo-median reference chip



## Array reference layout

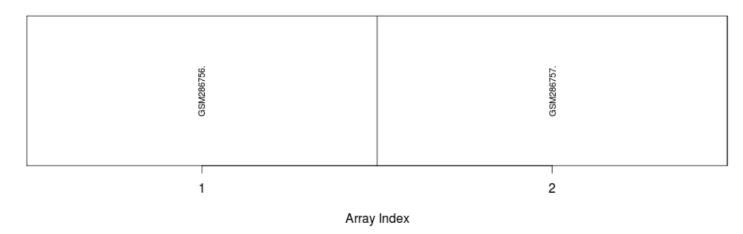


black/gray: regular match/mismatch probe blue/light blue: control match/mismatch probe red: unannotated probe (control region)

X Center of Intensity position

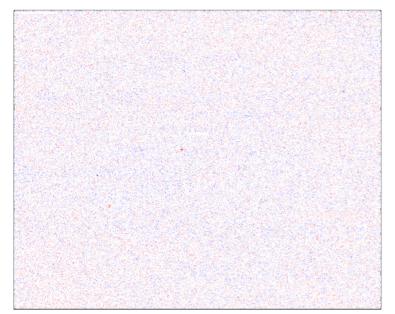
X Center of Intensity position

Sample Name



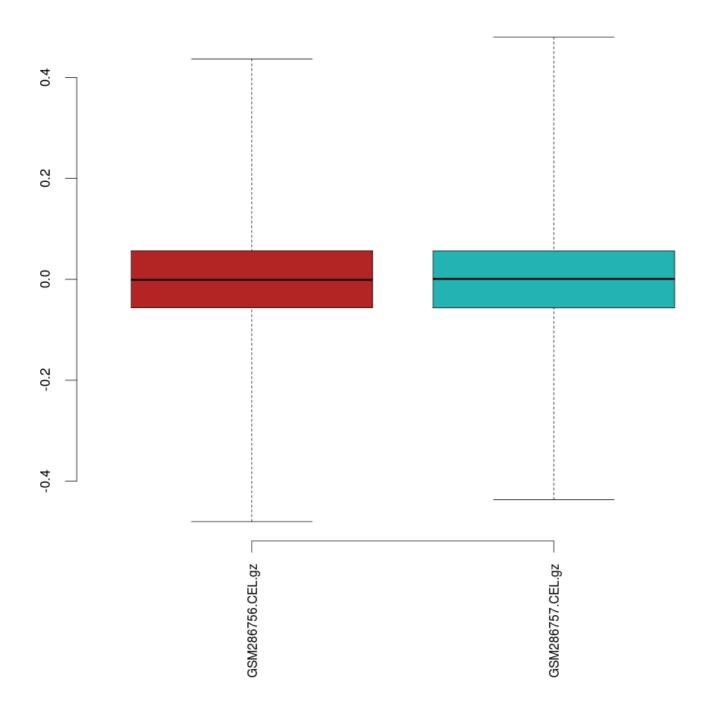
# 2D virtual PLM image for model characteristic: resids

GSM286756.CEL.gz GSM286757.CEL.gz





# Relative Log Expression (RLE) RLE distributions should be centered around 0



#### Array-Array Intensity Correlation

