

The ENmix User's Guide

Analyzing Illumina HumanMethylation450 BeadChip

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1 Introduction

The ENmix package provides tools for data preprocessing of the Illumina HumanMethylation450 array data to improve data quality, including functions for data quality control, background correction and inter-array normalization. In addition, the package also provides parallel computing wrappers for BMIQ probe design type bias correction and ComBat batch effect correction.

The Illumina Infinium HumanMethylation450 BeadChip has a complicated design. The array uses bisulfite converted DNA to estimate methylated (M) and unmethylated (U) allele intensity at individual CpG site. Two different assay chemistries are employed. The Infinium I assay is used for 28% (135, 476) of the CpGs on array and has 2 bead types for each CpG locus: one for the methylated and one for the unmethylated alleles. Signal intensities for both alleles at a locus are scanned on the same color channel (Cy3 green for some loci and Cy5 red for others). For a given type I bead, the intensity of the unused color channel has been proposed as a means to estimate background, and termed the out-of-band (oob) intensity. The Infinium II assay is used for 72% (350, 036) of the CpGs on the array and uses a single bead type per CpG. It utilizes two different colors to represent the two different alleles. These are assessed via single base extension with guanine (labeled with Cy3) for methylated, or adenine (labeled with Cy5) for unmethylated alleles. The HumanMethylation450K Beadchip has 850 internal control probes to monitor experimental procedures at different steps, including 613 negative control probes to measure background intensity and 16 probes to monitor bisulfite conversion efficiency.

2 Citation

If you are using ENmix package, please cite the following publications:

- The package:
Zongli Xu, Liang Niu, Leping Li and Jack A. Taylor ENmix: a novel background correction method for Illumina HumanMethylation450 BeadChip, Under review.
- Function `normalize.quantile.450k`:
Pidsley R, Y Wong CC, Volta M, Lunnon K, Mill J, Schalkwyk LC. A data-driven approach to preprocessing Illumina 450K methylation array data, BMC Genomics, 2013, 14:293
- Function `bmiq.mc`:
Teschendorff AE, Marabita F, Lechner M, Bartlett T, Tegner J, Gomez-Cabrero D, Beck S. A beta-mixture quantile normalization method for correcting probe design bias in Illumina Infinium 450 k DNA methylation data, Bioinformatics, 2013, 29(2):189-96
- Function `Combat.mc`:
Johnson WE, Li C, Rabinovic A. Adjusting batch effects in microarray expression data using empirical Bayes methods, Biostatistics, 2007, 8(1):118-27

Thanks for your help!

3 Setting up the data

The first step is to import array raw data files (*.idat) using functions provided in R package `minfi` to create an object of `RGChannelSetExtended`.

```
> library(ENmix)
> require(minfi)
> #see minfi user guide for the format of sample_sheet.txt file
> targets <- read.table("./sample_sheet.txt", header=T)
> rgSet <- read.450k.exp( targets = targets, extended = TRUE)
> # or read in all idat files under a directory
> rgSet <- read.450k.exp(base = "path_to_directory_idat_files",
+ targets = NULL, extended = TRUE, recursive=TRUE)
```

When methylation IDAT raw data files are not available, such as in most publically available datasets, users can use methylated (M) and unmethylated (U) intensity data to create an object of `MethylSet`.

```
> M<-matrix_for_methylated_intensity
> U<-matrix_for_unmethylated_intensity
> pheno<-as.data.frame(cbind(colnames(M), colnames(M)))
> names(pheno)<-c("Basename", "filenames")
> rownames(pheno)<-pheno$Basename
> pheno<-AnnotatedDataFrame(data=pheno)
> anno<-c("IlluminaHumanMethylation450k", "ilmn12.hg19")
```

```
> names(anno) <- c("array", "annotation")
> mdat <- MethySet(Meth = M, Unmeth = U, annotation=anno,
+ phenoData=pheno)
```

As an example for testing, users can use IDAT files installed in R data package `minfiData` to create an object of `RGChannelSetExtended`.

```
> library(ENmix)
> require(minfi)
> require(minfiData)
> sheet <- read.450k.sheet(file.path(find.package("minfiData"),
+ "extdata"), pattern = "csv$")
> rgSet <- read.450k.exp(targets = sheet, extended = TRUE)
```

4 Quality Control

4.0.1 Internal control probes

Illumina 450k chip incorporated 15 different types of internal controls (total of 848 probes). The function `plotCtrl` can plot intensity values for each type of the controls to evaluate data quality or the performance of specific steps in the process flow. See Illumina Infinium HD Methylation Assay for detailed description on how to interpret these control figures. Here is a list of the control types:

Control types	Number of probes
Sample-Independent Controls	
STAINING	4
EXTENSION	4
HYBRIDIZATION	3
TARGET REMOVAL	2
RESTORATION	1
Sample-Dependent Controls	
BISULFITE CONVERSION I	12
BISULFITE CONVERSION II	4
SPECIFICITY I	12
SPECIFICITY II	3
NON-POLYMORPHIC	4
NORM_A	32
NORM_C	61
NORM_G	32
NORM_T	61
NEGATIVE	613

```
> plotCtrl(rgSet)
```

4.0.2 Filtering out low quality samples and probes

Data quality measures, including detection P values, number of bead for each methylation reads and average intensities for bisulfite conversion probes can be extracted using function `QCinfo` from an object of `RGChannelSetExtended`. The information can be used for identifying low quality data points. Samples or probes with large percentage of low quality data can be excluded using function `QCfilter`. Appropriate thresholds (`samplethre`, `CpGthre` and `bisulthre`) can be explored from figures and data outputted by function `QCinfo`. Users can also specify a list of samples or CpGs to be filtered out by using options `outid` and `outCpG`. Density plot of total intensity (methylated intensity + unmethylated intensity) density plot of methylation beta values before and after filtering can be produced using option `plot=TRUE`.

```
> qc<-QCinfo(rgSet)
> mraw <- preprocessRaw(rgSet)
> mraw<-QCfilter(mraw, qcinfo=qc, samplethre = 0.01, CpGthre = 0.05
+               ,bisulthre=5000,plot=TRUE, outid=NULL, outCpG=NULL)
```

5 Background correction

Function `preprocessENmix` incorporated a model based background correction method ENmix, which models methylation signal intensities with a flexible exponential-normal mixture distribution, together with a truncated normal distribution to model background noise.

```
> mdat<-preprocessENmix(rgSet, bgParaEst="oob", nCores=6)
> mdat<-QCfilter(mdat, qcinfo=qc, samplethre = 0.01, CpGthre = 0.05
+ ,bisulthre=5000,plot=TRUE, outid=NULL, outCpG=NULL)
```

6 Inter-array normalization

Function `normalize.quantile.450k` can be used to perform quantile normalization on methylation intensity value

```
> mdat<-normalize.quantile.450k(mdat, method="quantile1")
```

7 Probe type bias correction

Function `bmiq.mc` is a multi-core parallel computing wrapper for the `BMIQ` function in R package `watermelon`.

```
> beta<-bmiq.mc(mdat, nCores=6)
```

8 Principal component regression analysis plot

Principal component regression can be used to explore methylation data variance structure (or source of variance) and identifying possible confounding variables. Principal components are derived using singular value decomposition method in standardized (for each probe) beta value matrix.

```
> cov<-data.frame(group=pData(mdat)$Sample_Group,
+ , slide=factor(pData(mdat)$Slide))
> pcrplot(beta, cov, npc=6)
```

9 Batch effect correction

Function `ComBat.mc` is a multi-core parallel computing wrapper for the `ComBat` function in R package `sva`.

```
> batch<-factor(pData(mdat)$Slide)
> betaC<-ComBat.mc(beta, batch, nCores=6, mod=NULL)
```

10 Multimodal CpGs

Function `nmode.mc` uses an empirical approach to identify multimodal distributed CpGs. When measured in a population of people the majority of CpGs on the Illumina HumanMethylation450 BeadChip have unimodal distributions of DNA methylation values with relatively small between-person variation. However, some CpGs (typically 10,000 often seemingly the result of SNPs in the probe region) may have multimodal distributions of methylation values with sizeable differences between modes and greater between-person variation. These multimodal distributed data are usually caused by SNP effect, problematic probe design or some other unknown artifacts instead of actual methylation level and thus should be excluded from DNA methylation analysis. Although researchers have often excluded CpGs based on SNP annotation information, we found that this approach alone may exclude many well-distributed (unimodal) CpGs, while still failing to identify other multi-modal CpGs. We developed an empirical approach to identify CpGs that are not uni-modally distributed, so that researchers can make more informed decisions about whether to exclude them in their particular study populations and analyses.

```
> nmode<- nmode.mc(beta, minN = 3, modedist=0.2, nCores = 5)
```

To illustrate the function, we have applied the approach in two different datasets: Sister study data (SS, n=200, Harlid et al, Plos one, 2015) and Norway Facial Clefts Study (NCL, n=891, Markunas et al. Environ Health Perspect. 2014) datasets. To simplify our illustration, we excluded 52,238 CpGs on X or Y chromosome, and non-specific probes annotated by Price et al (Price et al, Epigenetics Chromatin, 2013).

From the following summary table we can see this approach can correctly identifies all 65 known SNP probes as not unimodal in both datasets. More than 90% of CpGs with an annotated SNP in probe region are uni-modal in both populations and could be kept in data analysis with exclusion of a few outliers. For CpGs with any SNP at CpG target site, more than 80% are unimodal, and even 50% of CpGs with an annotated common (minor allele frequency > 0.05) SNP at target site have unimodal distributions.

Table 1. Number of unimodal and multimodal CpGs in two independent datasets.

	Unimodal in NCL				Concordance	% unimodal	
	Yes		No			NCL	SS
	Unimodal in SS		Unimodal in SS				
	Yes	No	Yes	No			
SNP probe	0	0	0	65	1.00	0	0
SNP in target site with MAF 0.05	2785	493	315	2584	0.87	0.531	0.502
SNP in target site#	14131	810	409	2844	0.93	0.821	0.799
SNP in probe#	109480	2140	676	3227	0.98	0.966	0.954
No SNP in probe#	314502	2264	443	542	0.99	0.997	0.991

*Based on 1000 genome project data for European population

based on annotation by Price et al, Epigenetics Chromatin, 2013

11 Example Analysis

Working with IDAT files

```
> library(ENmix)
> #read in raw intensity data
> sheet <- read.450k.sheet(file.path(find.package("minfiData"),
+   "extdata"), pattern = "csv$")
> rgSet <- read.450k.exp(targets = sheet, extended = TRUE)
> #Control plots
> plotCtrl(rgSet)
> #QC info
> qc<-QCinfo(rgSet)
> #Search for multimodal CpGs
> #sample size in this example data is too small for this purpose!
> #should not use beta matrix after ComBat analysis for this purpose!
> mraw <- preprocessRaw(rgSet)
> beta<-getBeta(mraw, "Illumina")
> nmode<-nmode.mc(beta, minN = 3, modedist=0.2, nCores = 6)
> #Frequency polygon plot to examining beta value distribution
> multifreqpoly(beta)
> #user specified CpG list to be excluded
> outCpG = names(nmode)[nmode>1]
> #exclude non-specific binding probes or probes affected by SNP et al.
> #outCpG = unique(c(outCpG,non-specific bind probes,snp probes,...))
> #filter out low quality samples and probes
```

```

> mraw<-QCfilter(mraw, qcinfo=qc, samplethre = 0.01, CpGthre = 0.05
+      ,bisulthre=5000,plot=TRUE, outid=NULL, outCpG=outCpG)
> cglisr=rownames(mraw)
> idlist=colnames(mraw)
> exCpG=rownames(beta)[!(rownames(beta) %in% cglisr)]
> #background correction
> mdat<-preprocessENmix(rgSet, bgParaEst="oob", dyeCorr=TRUE, exCpG=exCpG,
> mdat=mdat[,colnames(mdat) %in% idlist]
> #between-array normalization
> mdat<-normalize.quantile.450k(mdat, method="quantile1")
> #probe type bias correction
> beta<-bmiq.mc(mdat, nCores=6)
> # Principal component regression analysis plot
> cov<-data.frame(group=pData(mdat)$Sample_Group,
+      slide=factor(pData(mdat)$Slide))
> pcrplot(beta, cov, npc=6)
> #batch correction
> batch<-factor(pData(mdat)$Slide)
> betaC<-ComBat.mc(beta, batch, nCores=6, mod=NULL)

```

12 SessionInfo

- R version 2.14.1 (2011-12-22), x86_64-unknown-linux-gnu
- Locale: LC_CTYPE=en_US.UTF-8, LC_NUMERIC=C, LC_TIME=en_US.UTF-8, LC_COLLATE=en_US.UTF-8, LC_MONETARY=en_US.UTF-8, LC_MESSAGES=en_US.UTF-8, LC_PAPER=C, LC_NAME=C, LC_ADDRESS=C, LC_TELEPHONE=C, LC_MEASUREMENT=en_US.UTF-8, LC_IDENTIFICATION=C
- Base packages: base, datasets, graphics, grDevices, methods, stats, utils
- Loaded via a namespace (and not attached): tools 2.14.1

13 References

Zongli Xu, Liang Niu, Leping Li and Jack A. Taylor, ENmix: a novel background correction method for Illumina HumanMethylation450 BeadChip, Under review

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DNA Methylation microarrays. *Bioinformatics*, 30(10), pp. 1363-1369.

Pidsley, R., CC, Y.W., Volta, M., Lunnon, K., Mill, J. and Schalkwyk, L.C. (2013) A data-driven approach to preprocessing Illumina 450K methylation array data. *BMC genomics*, 14, 293.

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