

# iChip: A Package for Analyzing Multi-platform ChIP-chip data with Various Sample Sizes

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## Contents

<b>1</b>	<b>Introduction</b>	<b>1</b>
<b>2</b>	<b>Agilent and Affymetrix ChIP-chip Data</b>	<b>2</b>
<b>3</b>	<b>Example1 — Analyzing the Agilent Promoter Array Data</b>	<b>2</b>
<b>4</b>	<b>Example2 — Analyzing the Affymetrix Tiling Array Data</b>	<b>8</b>
<b>5</b>	<b>Tips</b>	<b>13</b>

## 1 Introduction

This package implements the models proposed by Mo and Liang (2010a, b) for ChIP-chip data analysis. The package can be used to analyze the ChIP-chip data from multiple platforms (e.g. Affymetrix, Agilent, and NimbleGen) with various genomic resolutions and various sample sizes. Mo and Liang (2010a,b) proposed Bayesian Hierarchical models to model the ChIP-chip data in which the spatial dependency of the data is modeled through ferromagnetic high-order or standard Ising models. Briefly, without loss of generality, the proposed methods let each probe be associated with a binary latent variable  $X_i \in (0, 1)$ , where  $i$  denotes the ID for the probe, and  $X_i = 1$  denotes that the probe is an enriched probe, and 0 otherwise. In the first stage, conditioning on the latent variable, the probe enrichment measurements for each state (0 or 1) are modeled by normal distributions. Here, the probe enrichment measurement could be any appropriate measurement for comparison of IP-enriched and control samples. For example, the measurement could be a log2 ratio of the IP-enriched and control samples for a single replicate, or a summary statistic such as t-like statistic or mean difference for multiple replicates. In the second stage, the latent variable is modeled by ferromagnetic Ising models. The Gibbs sampler and Metropolis

algorithm are used to simulate from the posterior distributions of the model parameters, and the posterior probabilities for the probes in the enriched state ( $X_i = 1$ ) is used for statistical inference. A probe with a high posterior probability in the enriched state will provide strong evidence that the probe is an enriched probe. For further details, we refer the user to Mo and Liang's papers. If you use this software to analyze your data, we will appreciate it if you can cite our papers.

## 2 Agilent and Affymetrix ChIP-chip Data

A subset of the Oct4 (Boyer et al., 2005) and the p53 (Cawley et al, 2004) data are used for the purpose of illustration. The average genomic resolutions for the Oct4 and p53 data are about 280 bps and 35 bps, respectively. Both the Oct4 and p53 data have been log2 transformed and quantile-normalized. Note iChip software doesn't provide functions for data normalization. The users should normalize their data before using iChip software. For one-color and two-color data, one can use the quantile method (e.g., see the function *normalize.quantiles()* in the **affy** package). For two-color data, one can also use the *loess* method (e.g., see the function *normalizeWithinArrays()* in the **limma** package). The full Oct4 data can be obtained from

```
http://jura.wi.mit.edu/young\_public/hESregulation/Data\_download.html
```

The full p53 data can be obtained from

```
http://www.gingeras.org/affy\_archive\_data/publication/tfbs/
```

## 3 Example1 — Analyzing the Agilent Promoter Array Data

Let's start analyzing the low resolution Oct4 data. Firstly, we need to calculate the enrichment measurement for each probe. Although the enrichment measurement could be any appropriate measurement for comparison of IP-enriched and control samples, we suggest using the empirical Bayes t-statistic for multiple replicates, which can be easily calculated using the **limma** package (Smyth, 2004). Here, we call the empirical Bayes t-statistic limma t-statistic. For the users who are not familiar with limma t-statistic, we provide a wrapper function **lmtstat** for the calculation.

There are two replicates for the Oct4 data. The enriched DNA was labeled with Cy5 (red) dye and the control DNA was labeled with Cy3 (green) dye.

```
> library(iChip)
> data(oct4)
> head(oct4, n = 3L)

  chr position    green1    green2     red1     red2
1 20      70312  6.969102  6.847819  6.808445  7.063581
2 20      70601  6.625190  6.176981  6.996920  6.391692
3 20      70873 10.334613 11.072903  9.521095 10.785880
```

To use the iChip1 and iChip2 function, the data must be sorted, firstly by chromosome then by genomic position. It may be a good habit to sort the data at the beginning, although function **lmtstat** doesn't require the data be sorted.

```
> oct4 = oct4[order(oct4[, 1], oct4[, 2]), ]
```

Calculate the enrichment measurements — two-sample limma t-statistics.

```
> oct4lmt = lmtstat(oct4[, 5:6], oct4[, 3:4])
```

Here, we treat the IP-enriched and control data as independent data although both the IP-enriched and control samples were hybridized to the same array. This is because the quantile-normalization method was applied to the oct4 data. If the data are normalized using *loess* method, the resulting data are in log ratio format (e.g.,  $\log_2(\text{IP-enriched}/\text{control})$ ). In this case, one can calculate the paired limma t-statistics. Suppose a matrix called `log2ratio` are the loess-normalized data, where each column corresponds to a sample, the paired limma t-statistics can be calculated using **lmtstat(log2ratio)**.

Prepare the data for iChip2 function.

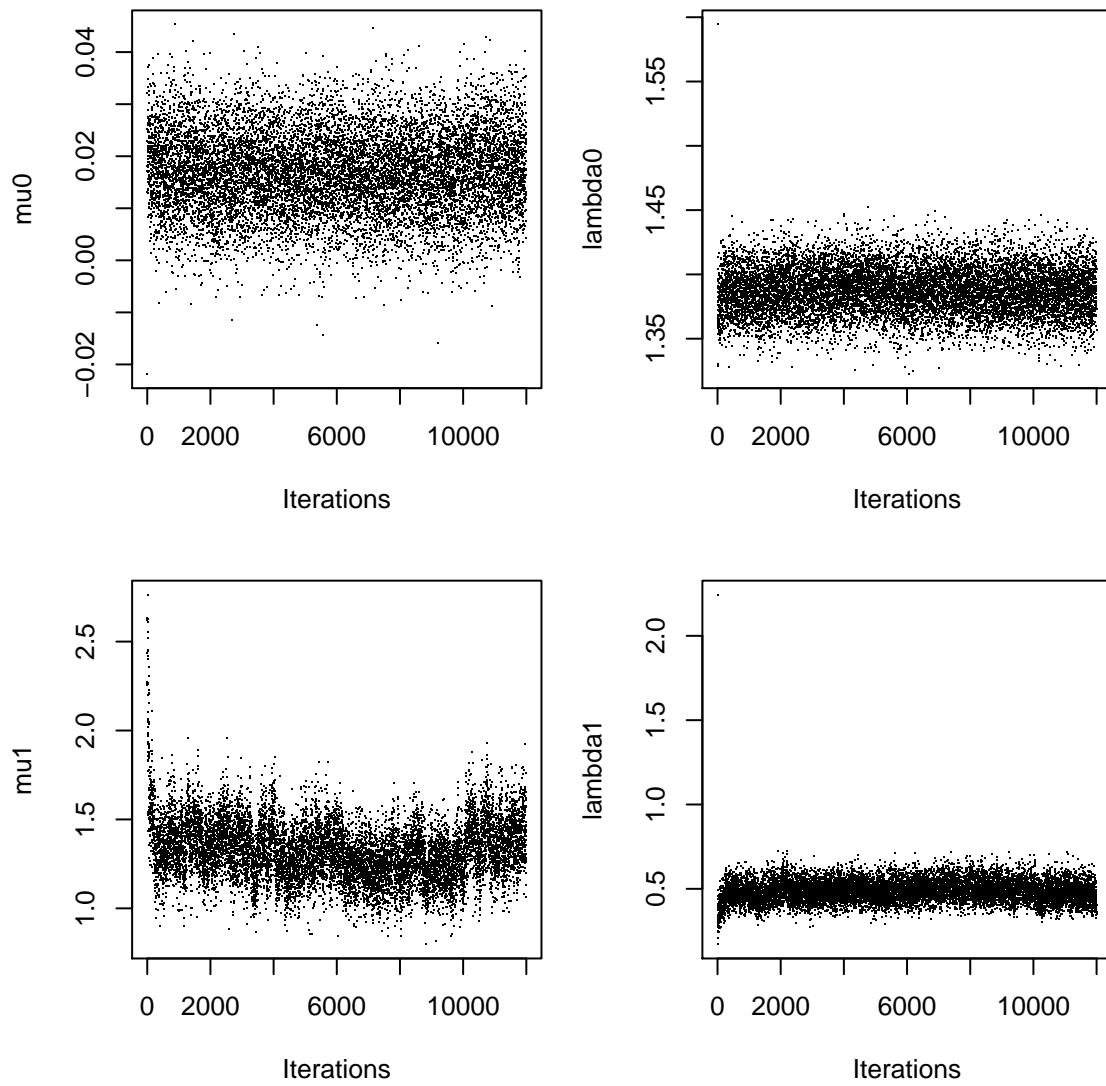
```
> oct4Y = cbind(oct4[, 1], oct4lmt)
```

Apply the second-order Ising model to the ChIP-chip data by setting `winsize = 2`. According to our experience, a balance between high sensitivity and low FDR can be achieved when `winsize = 2`. The critical value of the second-order Ising model is about 1.0. For low resolution data, the value of beta could be around the critical value. In general, increasing beta value will lead to less enriched regions, which amounts to setting a stringent criterion for detecting enriched regions.

```
> set.seed(777)
> oct4res2 = iChip2(Y = oct4Y, burnin = 2000, sampling = 10000,
+   winsize = 2, sdcut = 2, beta = 1.25, verbose = FALSE)
```

Plot the model parameters to see whether they converge. In general, the model has converged when the parameters fluctuate around the modes of their distributions. If there is an obvious trend(e.g. continuous increase or decrease), one should increase the number of iterations in the burn-in phase. If this doesn't work, one can try to adjust the parameter **beta** to see how it affect the results.

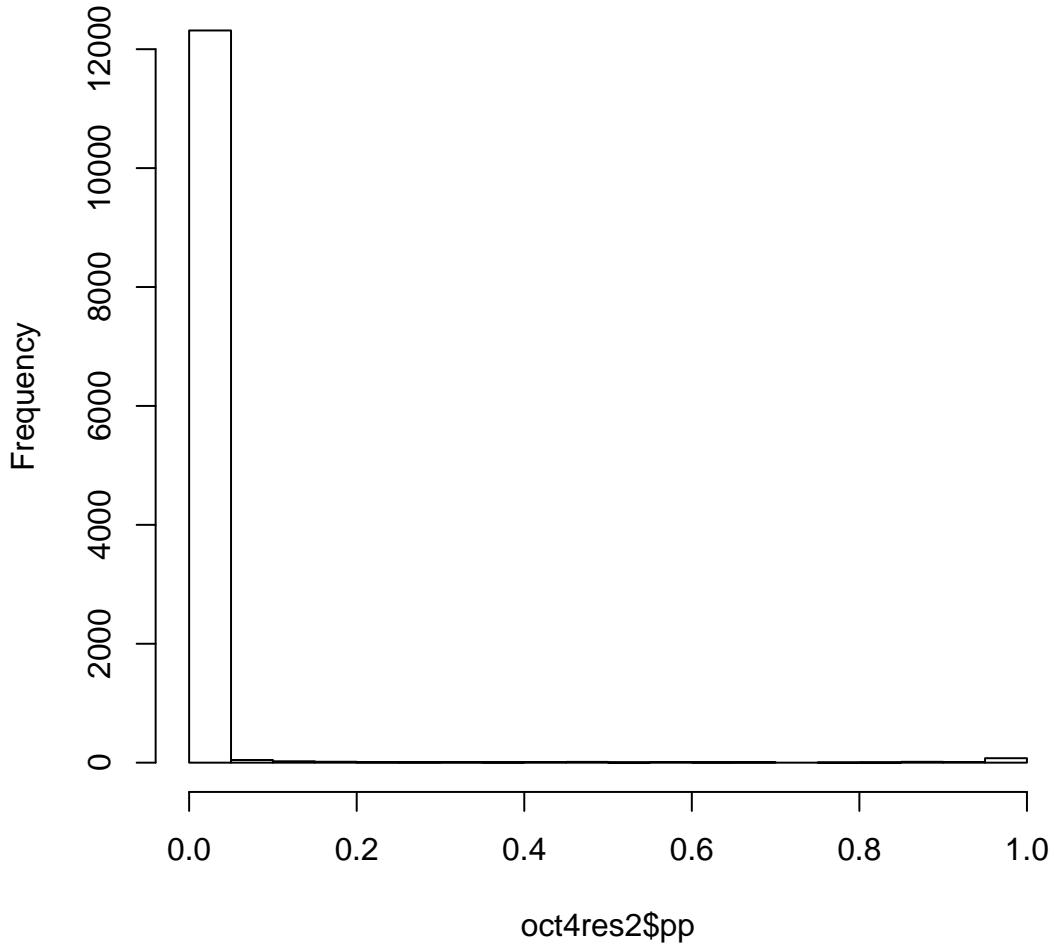
```
> par(mfrow = c(2, 2), mar = c(4.1, 4.1, 2, 1))
> plot(oct4res2$mu0, pch = ".", xlab = "Iterations", ylab = "mu0")
> plot(oct4res2$lambda0, pch = ".", xlab = "Iterations", ylab = "lambda0")
> plot(oct4res2$mu1, pch = ".", xlab = "Iterations", ylab = "mu1")
> plot(oct4res2$lambda1, pch = ".", xlab = "Iterations", ylab = "lambda1")
```



The histogram of the posterior probabilities should be dichotomized, either 0 or 1. For transcription factor binding site studies, the histogram should be dominated by 0.

```
> hist(oct4res2$pp)
```

### Histogram of oct4res2\$pp



Call the enriched regions detected by iChip2 using a posterior probability (pp) cutoff of 0.9.

```
> reg1 = enrichreg(pos = oct4[, 1:2], enrich = oct4lmt, pp = oct4res2$pp,
+       cutoff = 0.9, method = "ppcut", maxgap = 500)
> print(reg1)
```

	chr	gstart	gend	rstart	rend	peakpos	meanpp	maxpp	nprobe
1	20	3944132	3946241	1415	1427	3946061	0.96	1	13
2	20	20291072	20291658	3293	3295	20291658	0.96	1	3
3	20	20292352	20294499	3296	3304	20293941	1.00	1	9
4	20	21441187	21450238	3441	3477	21445231	0.99	1	37
5	20	22519126	22519690	3545	3547	22519406	1.00	1	3
6	20	28137489	28138889	4307	4312	28137489	1.00	1	6

```

7 20 34633143 34633770    6132 6134 34633506    0.98    1     3
8 20 44034352 44034352    8313 8313 44034352    1.00    1     1
9 20 54633181 54635934    9703 9713 54633459    0.99    1    11

```

Call the enriched regions detected by iChip2 using a FDR cutoff of 0.01. The FDR is calculated using a direct posterior probability approach (Newton et al., 2004).

```

> reg2 = enrichreg(pos = oct4[, 1:2], enrich = oct4lmt, pp = oct4res2$pp,
+   cutoff = 0.01, method = "fdrcut", maxgap = 500)
> print(reg2)

```

	chr	gstart	gend	rstart	rend	peakpos	meanpp	maxpp	nprobe
1	20	3944132	3946241	1415	1427	3946061	0.96	1	13
2	20	20291344	20291658	3294	3295	20291658	0.98	1	2
3	20	20292352	20294499	3296	3304	20293941	1.00	1	9
4	20	21441187	21449717	3441	3475	21445231	1.00	1	35
5	20	22519126	22519690	3545	3547	22519406	1.00	1	3
6	20	28137489	28138889	4307	4312	28137489	1.00	1	6
7	20	34633143	34633770	6132	6134	34633506	0.98	1	3
8	20	44034352	44034352	8313	8313	44034352	1.00	1	1
9	20	54633181	54635934	9703	9713	54633459	0.99	1	11

BED file can be easily created using the output from function **enrichreg**, which can be used for motif discovery and visualized in the UCSC genome browser. For example,

```

> bed1 = data.frame(chr = paste("chr", reg2[, 1], sep = ""), reg2[, 2:3])
> print(bed1[1:3, ])

```

	chr	gstart	gend
1	chr20	3944132	3946241
2	chr20	20291344	20291658
3	chr20	20292352	20294499

Alternatively, one may create a BED file using the peak position of the enriched regions. For example,

```

> bed2 = data.frame(chr = paste("chr", reg2[, 1], sep = ""), gstart = reg2[, 6] - 100, gend = reg2[, 6] + 100)
> print(bed2[1:3, ])

```

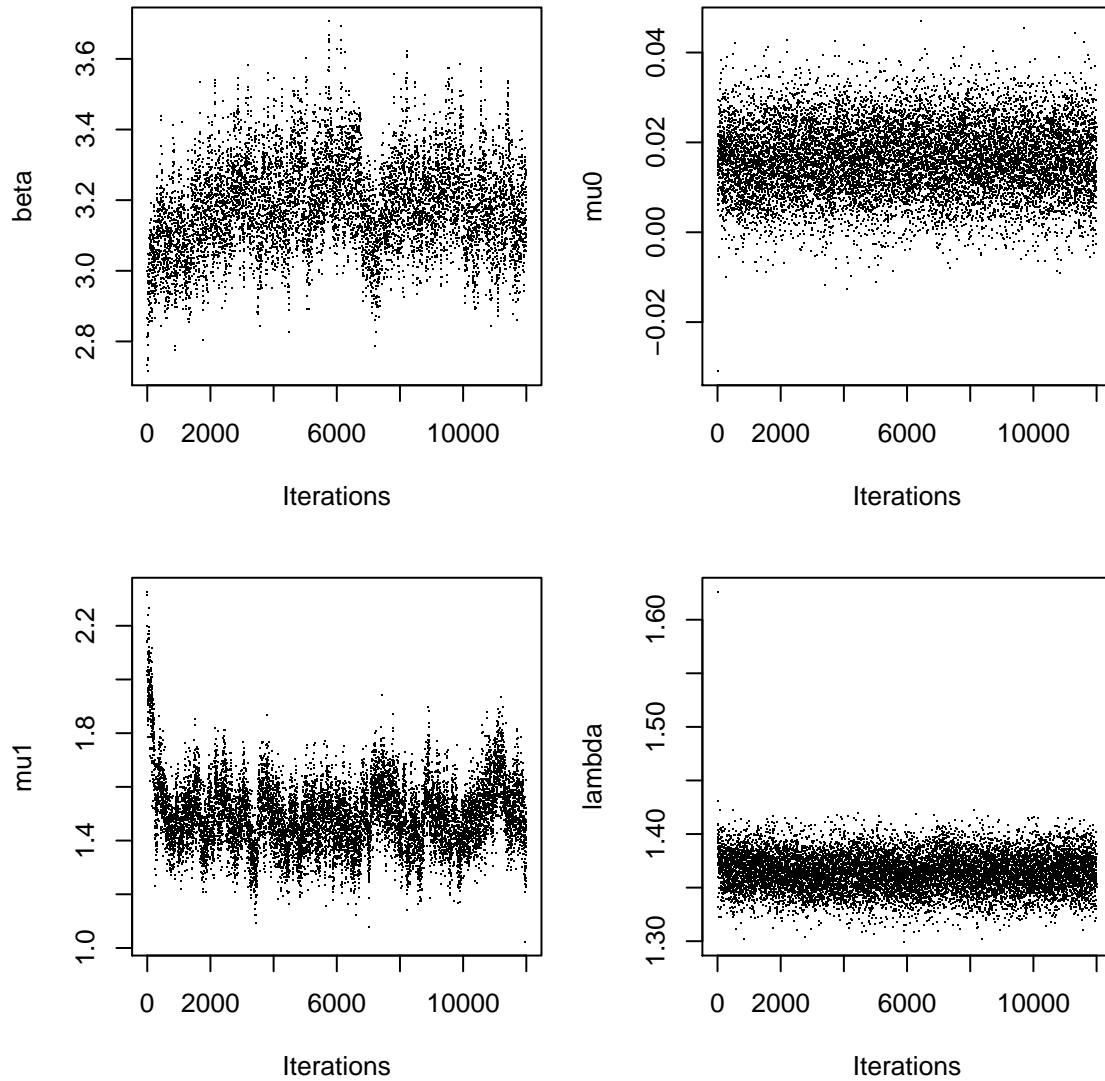
	chr	gstart	gend
1	chr20	3945961	3946161
2	chr20	20291558	20291758
3	chr20	20293841	20294041

Model the oct4 data using the first-order Ising model.

```
> oct4res1 = iChip1(enrich = oct4lmt, burnin = 2000, sampling = 10000,
+      sdcut = 2, beta0 = 3, minbeta = 0, maxbeta = 100, normsd = 0.1,
+      verbose = FALSE)
```

Plot the model parameters to see whether they converge.

```
> par(mfrow = c(2, 2), mar = c(4.1, 4.1, 2, 1))
> plot(oct4res1$beta, pch = ".", xlab = "Iterations", ylab = "beta")
> plot(oct4res1$mu0, pch = ".", xlab = "Iterations", ylab = "mu0")
> plot(oct4res1$mu1, pch = ".", xlab = "Iterations", ylab = "mu1")
> plot(oct4res1$lambda, pch = ".", xlab = "Iterations", ylab = "lambda")
```



Call the enriched regions detected by iChip1.

```
> enrichreg(pos = oct4[, 1:2], enrich = oct4lmt, pp = oct4res1$pp,
+           cutoff = 0.9, method = "ppcut", maxgap = 500)
```

	chr	gstart	gend	rstart	rend	peakpos	meanpp	maxpp	nprobe
1	20	3944132	3946241	1415	1427	3946061	1.00	1	13
2	20	20291072	20291658	3293	3295	20291658	0.96	1	3
3	20	20292352	20294805	3296	3305	20293941	0.99	1	10
4	20	21441187	21449717	3441	3475	21445231	0.99	1	35
5	20	22519126	22519690	3545	3547	22519406	1.00	1	3
6	20	28137489	28138889	4307	4312	28137489	0.99	1	6
7	20	34632735	34633770	6131	6134	34633506	0.97	1	4
8	20	54633181	54636203	9703	9714	54633459	0.99	1	12

```
> enrichreg(pos = oct4[, 1:2], enrich = oct4lmt, pp = oct4res1$pp,
+           cutoff = 0.01, method = "fdrcut", maxgap = 500)
```

	chr	gstart	gend	rstart	rend	peakpos	meanpp	maxpp	nprobe
1	20	3944132	3946241	1415	1427	3946061	1.00	1	13
2	20	20291072	20291658	3293	3295	20291658	0.96	1	3
3	20	20292352	20294805	3296	3305	20293941	0.99	1	10
4	20	21441187	21449717	3441	3475	21445231	0.99	1	35
5	20	22519126	22519690	3545	3547	22519406	1.00	1	3
6	20	28137489	28138889	4307	4312	28137489	0.99	1	6
7	20	34632735	34633770	6131	6134	34633506	0.97	1	4
8	20	54633181	54636203	9703	9714	54633459	0.99	1	12

## 4 Example2 — Analyzing the Affymetrix Tiling Array Data

Now, let's analyze the high resolution p53 data.

```
> data(p53)
> head(p53, n = 3L)

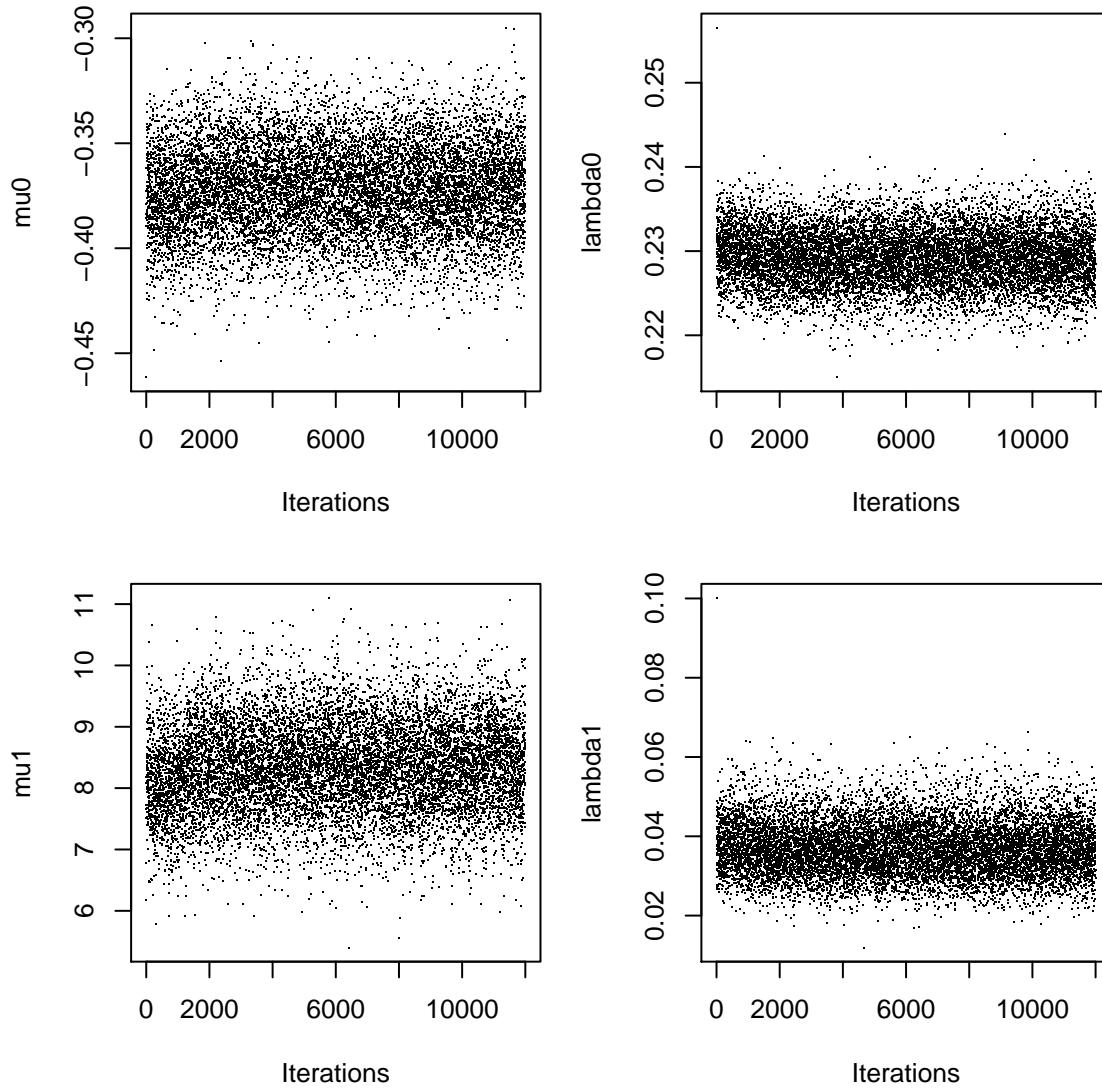
      chr position      CON      CON      CON      CON      CON      CON
783581  22 27980300 9.576077 10.90728  9.585894 11.14744 10.23070 11.01191
783582  22 27980347 9.941713 10.66333 10.031774 10.86761 10.23361 10.81113
783583  22 27980372 9.932955 10.63290  9.995038 10.42966 10.02872 10.39427
          IP      IP      IP      IP      IP      IP
783581  9.70315 10.64889 10.53961  9.376407  9.869731 10.89024
783582 10.32478 10.39944 10.39757 10.602358 10.544956 10.24796
783583 10.27690 10.28804 10.27836 10.244013  9.961192 10.18641

> p53 = p53[order(p53[, 1], p53[, 2]), ]
> p53lmt = lmtstat(p53[, 9:14], p53[, 3:8])
> p53Y = cbind(p53[, 1], p53lmt)
```

For high resolution data, beta could be set to a relatively large value (e.g. from 2-4). In general, increasing beta value will lead to less enriched regions, which amounts to setting a stringent criterion for detecting enriched regions.

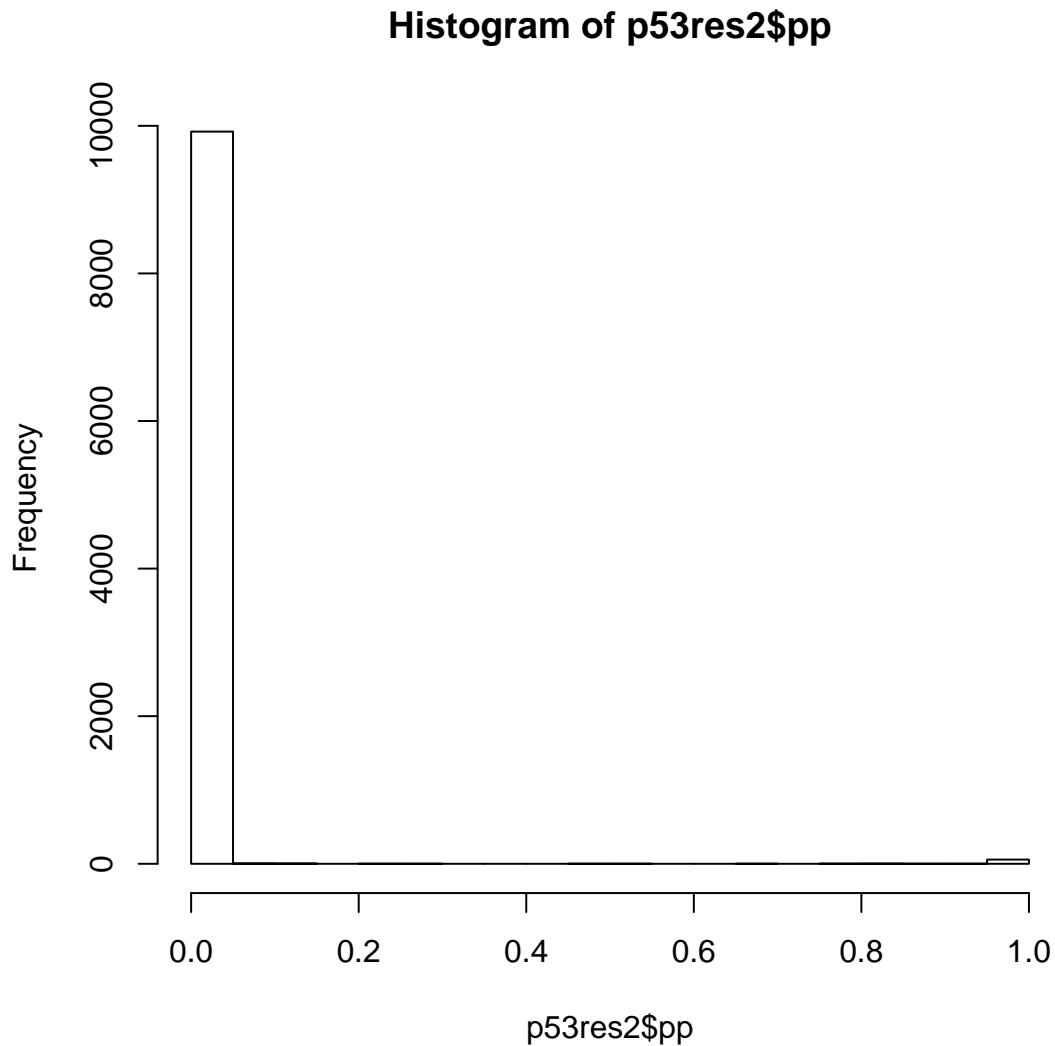
```
> p53res2 = iChip2(Y = p53Y, burnin = 2000, sampling = 10000, winsize = 2,
+      sdcut = 2, beta = 2.5)

> par(mfrow = c(2, 2), mar = c(4.1, 4.1, 2, 1))
> plot(p53res2$mu0, pch = ".", xlab = "Iterations", ylab = "mu0")
> plot(p53res2$lambda0, pch = ".", xlab = "Iterations", ylab = "lambda0")
> plot(p53res2$mu1, pch = ".", xlab = "Iterations", ylab = "mu1")
> plot(p53res2$lambda1, pch = ".", xlab = "Iterations", ylab = "lambda1")
```



The histogram of the posterior probabilities should be dichotomized, either 0 or 1. For transcription factor binding site studies, the histogram should be dominated by 0.

```
> hist(p53res2$pp)
```



```
> enrichreg(pos = p53[, 1:2], enrich = p53lmt, pp = p53res2$pp,  
+           cutoff = 0.9, method = "ppcut", maxgap = 500)
```

chr	gstart	gend	rstart	rend	peakpos	meanpp	maxpp	nprobe	
1	22	28211540	28211540	2991	2991	28211540	0.96	0.96	1
2	22	28269526	28270158	3705	3724	28269751	1.00	1.00	20
3	22	28345939	28346341	4831	4842	28346226	1.00	1.00	12
4	22	28380484	28380676	5440	5446	28380676	1.00	1.00	7
5	22	28775272	28775790	9878	9895	28775550	1.00	1.00	18

```

> enrichreg(pos = p53[, 1:2], enrich = p53lmt, pp = p53res2$pp,
+           cutoff = 0.01, method = "fdrcut", maxgap = 500)

  chr    gstart      gend rstart rend peakpos meanpp maxpp nprobe
1 22 28211540 28211540    2991 2991 28211540    0.96  0.96      1
2 22 28269497 28270158    3704 3724 28269751    0.99  1.00     21
3 22 28345939 28346341    4831 4842 28346226    1.00  1.00     12
4 22 28380484 28380725    5440 5447 28380676    0.98  1.00      8
5 22 28775272 28775790    9878 9895 28775550    1.00  1.00     18

```

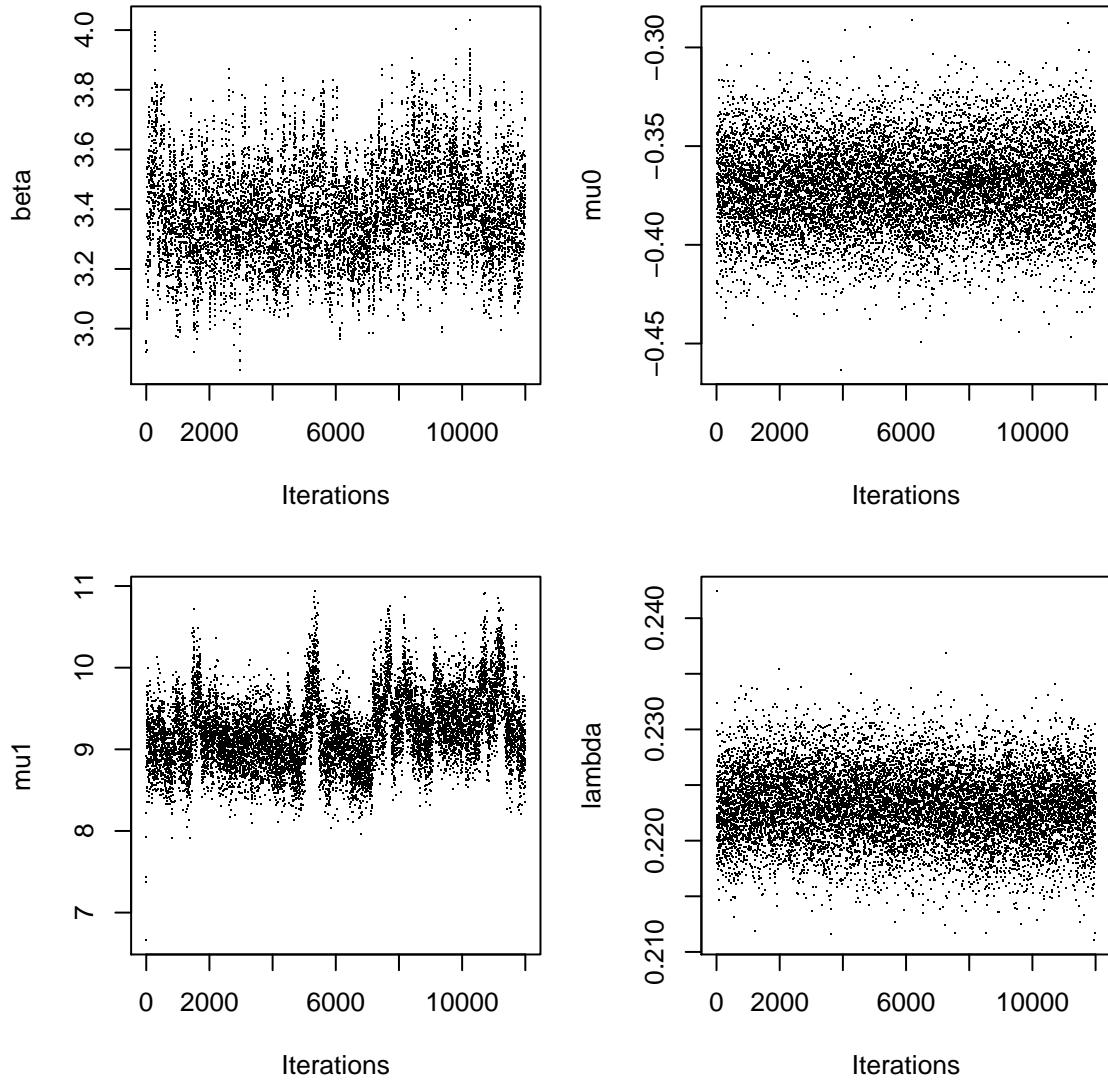
Model the p53 data using the first-order Ising model.

```

> p53res1 = iChip1(enrich = p53lmt, burnin = 2000, sampling = 10000,
+                   sdcut = 2, beta0 = 3, minbeta = 0, maxbeta = 100, normsd = 0.1,
+                   verbose = FALSE)

> par(mfrow = c(2, 2), mar = c(4.1, 4.1, 2, 1))
> plot(p53res1$beta, pch = ".", xlab = "Iterations", ylab = "beta")
> plot(p53res1$mu0, pch = ".", xlab = "Iterations", ylab = "mu0")
> plot(p53res1$mu1, pch = ".", xlab = "Iterations", ylab = "mu1")
> plot(p53res1$lambda, pch = ".", xlab = "Iterations", ylab = "lambda")

```



```
> enrichreg(pos = p53[, 1:2], enrich = p53lmt, pp = p53res1$pp,
+           cutoff = 0.9, method = "ppcut", maxgap = 500)
```

	chr	gstart	gend	rstart	rend	peakpos	meanpp	maxpp	nprobe
1	22	28211540	28211540	2991	2991	28211540	1.00	1	1
2	22	28269526	28270058	3705	3721	28269751	0.97	1	17
3	22	28345939	28346341	4831	4842	28346226	1.00	1	12
4	22	28380574	28380676	5443	5446	28380676	1.00	1	4
5	22	28775272	28775790	9878	9895	28775550	0.99	1	18

```
> enrichreg(pos = p53[, 1:2], enrich = p53lmt, pp = p53res1$pp,
+           cutoff = 0.01, method = "fdrcut", maxgap = 500)
```

	chr	gstart	gend	rstart	rend	peakpos	meanpp	maxpp	nprobe
1	22	28211540	28211540	2991	2991	28211540	1.00	1	1
2	22	28269526	28270058	3705	3721	28269751	0.97	1	17
3	22	28345939	28346341	4831	4842	28346226	1.00	1	12
4	22	28380546	28380676	5442	5446	28380676	0.97	1	5
5	22	28775272	28775790	9878	9895	28775550	0.99	1	18

## 5 Tips

What happens when there is no enriched region? Suppose the data are just random noises.

```
> randomY = cbind(p53[, 1], rnorm(10000, 0, 1))
> randomres2 = iChip2(Y = randomY, burnin = 2000, sampling = 10000,
+   winsize = 2, sdcut = 2, beta = 2.5, verbose = FALSE)
```

Warning: all probes are in the same state at the last MCMC iteration.

NO enriched region is found!

```
> table(randomres2$pp)
```

```
1
10000
```

In this case, all the probes are only in one state. Since there are not enriched probe, the mean and variance become  $-\infty$  or  $\infty$ . In the MCMC simulations, we relabel the outputs according to the constraint  $\mu_0 < \mu_1$ , where  $\mu_0$  and  $\mu_1$  are the population means for the non-enriched and enriched probes, respectively (For details, see Mo and Liang, 2010a). That is, when  $\mu_0 > \mu_1$ ,  $\mu_0$  will be treated as the population mean of the enriched probes. As a result, no matter  $\mu_1 = \infty$  or  $\mu_1 = -\infty$ , the posterior probabilities of probes are all 1s or close to 1. Therefore, when this happens, it means there are not enriched region.

In addition, for transcription factor binding site studies, if the posterior probabilities are not dichotomized (major 0, minor 1), it suggests that the Ising model is not in the super-paramagnetic phase. Only the super-paramagnetic phase reflects the binding events on the chromosomes. Therefore, the users should increase the value of beta to let the phase transition occur so that the Ising model reach the super-paramagnetic phase.

The states of probes produced by iChip1 are also in the same state if there is not enriched region.

```
> randomres1 = iChip1(enrich = randomY[, 2], burnin = 2000, sampling = 10000,
+   sdcut = 2, beta0 = 3, minbeta = 0, maxbeta = 100, normsd = 0.1,
+   verbose = FALSE)
```

Warning: all probes are in the same state at the last MCMC iteration.

NO enriched region is found!

```
> table(randomres1$pp)
```

```
0  
10000
```

Although the above two examples only show the analysis for the data on a single chromosome, one can use iChip2 and iChip1 functions to analyze data with multiple chromosomes. Although a probe may not physically interact with its adjacent probes (e.g., the last probe of a chromosome and the first probe of the next chromosome, and two probes in the same chromosome that are adjacent but separated by a long genomic distance), in practice, it should be acceptable to consider the interactions between these adjacent and boundary probes. There are two reasons for this argument. Firstly, the number of these probes is quite small, compared to all the probes in the tiling arrays. Secondly, these boundary probes have a very high probability to be non-binding probes, thus it should be reasonable to consider the interactions between them. If we let these boundary and adjacent probes interact with each other, it has little effect on the results, but significantly simplify the algorithms for modeling the ChIP-chip data.

Finally, it should be noted that when the data are very noisy, the posterior mean of beta will be relatively small (e.g., around 1) when the iChip1 method is used. In this case, the probes are not dominated by non-binding probes. The user should increase the value of parameter **minbeta** or use the **iChip2** method for modeling. Both iChip1 and iChip2 functions run reasonably fast. It costs about 15-20 minutes for a data set with 500,000 probes for running 15,000 iterations on a 64-bit Linux machine with 2.4 GHZ CPU. If the total number of probes is relatively small, say, a half million, one may analyze the data in a single run. If the total number of probes are large, say, several millions, one may perform parallel computation. For example, one can divide the data to several pieces and run them simultaneously.

```
> sessionInfo()  
  
R version 2.10.1 (2009-12-14)  
x86_64-pc-linux-gnu  
  
locale:  
[1] LC_CTYPE=en_US.UTF-8      LC_NUMERIC=C  
[3] LC_TIME=en_US.UTF-8       LC_COLLATE=en_US.UTF-8  
[5] LC_MONETARY=C            LC_MESSAGES=en_US.UTF-8  
[7] LC_PAPER=en_US.UTF-8     LC_NAME=C  
[9] LC_ADDRESS=C              LC_TELEPHONE=C  
[11] LC_MEASUREMENT=en_US.UTF-8 LC_IDENTIFICATION=C  
  
attached base packages:  
[1] tools      stats      graphics   grDevices  utils      datasets  methods  
[8] base
```

```
other attached packages:  
[1] iChip_0.99.3 limma_3.2.1
```

## References

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- Newton, M., Noueiry, A., Sarkar, D., Ahlquist, P. (2004). Detecting differential gene expression with a semiparametric hierarchical mixture method. *Biostatistics* **5**, 155-176.
- Smyth, G. (2004). Linear models and empirical Bayes methods for assessing differential expression in microarray experiments. *Statistical Applications in Genetics and Molecular Biology*, **3**, Iss. 1, Article 3.