

# *msap* (v. 0.1-2) - User's Guide

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## 1 Introduction

*msap* provides a deep analysis of epigenetic variation starting from a binary data matrix indicating the presence or absence of EcoRI-HpaII and EcoRI-MspI fragments, typical of MSAP technique. After compare the data from both enzyme combinations, the program determines if each fragment is susceptible of methylation (representative of epigenetic variation) or if there is no evidence of methylation (representative of genetic variation). Different analyses of the variation and differentiation (genetic and epigenetic) among user-defined groups of samples are then performed, as well as the classification of the methylation occurrences in those groups. A comprehensive report of the analyses and several useful plots could help researchers to asses the epigenetic variation in their experiments using MSAP. Standard AFLP data is also suitable to be analyzed. All analyses follow a band-based strategy (Bonin et al. , 2007). There are several examples in the literature of MSAP experiments using some of the analyses provided by *msap* (see Herrera and Bazaga , 2010; Moran and Perez-Figueroa , 2011; Chwedorzewska and Bednarek , 2012; Gupta et al. , 2012; Rodriguez et al. , 2012) so the package could be useful in those kind of approaches.

The package is intended to be easy to use even for those people non-familiar to the R environment. Advanced users could take advantage of available source code to adapt *msap* for more complex analyses.

## 2 R basics. All you need to know about R to run *msap*

The only knowledge required for installing and running *msap* is about how to open an R session in your computer. R is a statistical programming language that provides many built in functions for performing statistical analysis and is also flexible to allow users to write their own functions.

R can be downloaded and installed for free from the website <http://cran.r-project.org> where detailed instructions for installing R on any operating system are provided. Accessing R is different for every operating system. For windows users, simply double click the R icon that is created after installation. For Mac

users, you can double click the R icon under your Applications menu. On Linux, in the terminal window simply type R at the command prompt and R will be opened within the terminal window.

When you open R, no matter the operating system you are using, you will see the command prompt symbol `>` which simply means that R is waiting for you to give it a command. To quit R, simply type `q()` in the command prompt and R will ask you if you want to save the workspace before quitting. And that's all.

### 3 Installing *msap*

You can install *msap* automatically from a R session. To install the last stable version from CRAN:

```
> install.packages("msap")
```

To get the last daily development version from R-Forge:

```
> install.packages("msap", repos="http://R-Forge.R-project.org")
```

The above instructions should install *msap* and all required dependencies.

### 4 Preparation of data

In order to use *msap* for analyzing your results from a MSAP experiment, you need to provide a data file with a binary matrix (1/0) indicating the presence or absence of EcoRI-HpaII and EcoRI-MspI fragments in a bunch of samples of two or more populations/groups. Data file should be a .csv file with markers as columns and two rows by sample, one for each isoschizomer reaction. File could be edited in the a spreadsheet of your choice (see Figure 1) and then saved as csv (with ',' as field separator). The final text file should look like Figure 2 if opened in a text editor.

The first row should include the markers name/references. The first column should provide the label for the group where the sample is included, with the aim to make comparisons between different groups. Second column is reserved for an arbitrary label (i.e. to name the sample). Third column should identify the isoschizomer with 'HPA' or 'MSP'. If you want to analyze a standard AFLP dataset the datafile format is the same, but the program will ignore content of third column and treat all rows as independent samples.

### 5 Running *msap*

We start by loading the *msap* package into an R session.

```
> library(msap)
```

	A	B	C	D	E	F	G	H	I	J	K	L	M	N	O	P	Q	R	S	T
1				m1	m2	m3	m4	m5	m6	m7	m8	m9	m10	m11	m12	m13	m14	m15	m16	m17
2	Pop1	a1	HPA	0	0	0	0	1	0	0	0	1	1	0	0	0	0	0	0	0
3	Pop1	a1	MSP	0	0	0	0	1	0	0	0	1	1	1	0	0	0	0	0	1
4	Pop1	a2	HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0
5	Pop1	a2	MSP	0	0	1	0	0	1	1	1	1	1	1	1	0	0	1	0	1
6	Pop1	a3	HPA	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0
7	Pop1	a4	MSP	1	1	0	1	0	0	0	0	1	1	1	0	0	0	0	0	1
8	Pop1	a5	HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0
9	Pop1	a5	MSP	0	0	1	0	0	0	1	0	1	1	1	1	0	0	1	0	1
10	Pop1	a6	HPA	1	1	0	1	0	0	0	0	1	1	1	0	0	0	0	0	1
11	Pop1	a6	MSP	0	0	0	0	0	0	0	1	1	1	1	1	0	0	1	0	0
12	Pop1	a7	HPA	0	1	0	1	0	0	0	0	1	1	1	1	0	0	1	0	0
13	Pop1	a7	MSP	0	0	1	0	0	0	0	0	1	1	0	0	0	0	0	0	0
14	Pop2	b1	HPA	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0
15	Pop2	b1	MSP	0	0	0	0	1	0	0	0	1	1	0	1	0	0	0	0	0
16	Pop2	b2	HPA	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0
17	Pop2	b2	MSP	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0
18	Pop2	b3	HPA	0	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0
19	Pop2	b3	MSP	0	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0
20	Pop2	b4	HPA	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0
21	Pop2	b4	MSP	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
22	Pop2	b5	HPA	0	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0
23	Pop2	b5	MSP	0	0	0	0	0	0	0	0	0	1	1	0	0	0	1	0	0
24	Pop2	b6	HPA	0	1	0	0	0	0	0	0	1	1	1	1	0	0	0	0	0
25	Pop2	b6	MSP	0	1	0	0	1	1	0	1	1	0	1	0	0	1	0	0	0
26	Pop3	c1	HPA	0	1	0	1	1	0	0	1	1	1	0	0	0	1	0	0	1
27	Pop3	c1	MSP	0	0	1	1	0	0	0	0	1	1	1	1	0	0	0	0	0
28	Pop3	c2	HPA	0	1	0	1	1	0	0	1	1	1	0	0	0	0	0	1	0
29	Pop3	c2	MSP	0	0	0	1	1	0	0	1	1	1	1	1	0	0	1	0	1
30	Pop3	c3	HPA	1	1	0	1	1	0	0	1	1	1	1	0	0	0	1	0	1

Figure 1: Data format as seen in a spreadsheet for edition

	m1	m2	m3	m4	m5	m6	m7	m8	m9	m10	m11	m12	m13	m14	m15	m16	m17	m18	m19	m20	m
Pop1,a1,HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	1	1	0	0
Pop1,a1,MSP	0	0	0	0	1	1	0	0	1	1	1	0	0	0	0	0	1	0	0	0	0
Pop1,a2,HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	1	1	0	0
Pop1,a2,MSP	0	0	0	1	0	0	1	1	1	1	1	0	0	1	0	1	0	0	1	0	0
Pop1,a3,HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	1	1	0	0
Pop1,a4,MSP	1	1	0	1	0	0	0	1	1	1	0	0	0	0	0	0	1	1	0	0	0
Pop1,a5,HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	1	1	0	0
Pop1,a5,MSP	0	0	0	1	0	0	0	1	0	1	1	0	0	1	0	1	0	0	1	0	0
Pop1,a6,HPA	1	1	0	1	0	0	0	1	1	1	0	0	0	0	0	0	1	0	1	0	0
Pop1,a6,MSP	0	0	0	0	0	0	0	1	1	1	1	0	0	1	0	0	1	0	0	0	0
Pop1,a7,HPA	0	1	0	1	0	0	0	1	1	1	1	0	0	1	0	0	0	1	0	0	0
Pop1,a7,MSP	0	0	1	0	0	0	0	1	1	0	0	0	0	0	0	0	0	1	0	1	0
Pop2,b1,HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	1	1	0	0
Pop2,b1,MSP	0	0	0	0	1	0	0	0	1	1	0	1	0	0	0	0	0	0	1	1	0
Pop2,b2,HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	1	1	0
Pop2,b2,MSP	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	1	1	0
Pop2,b3,HPA	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0	0	0	1	1	0
Pop2,b3,MSP	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0	0	0	1	1	0
Pop2,b4,HPA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	1	1	0
Pop2,b4,MSP	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0
Pop2,b5,HPA	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0	0	0	0	0	0
Pop2,b5,MSP	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0	0	0	1	0	0
Pop2,b6,HPA	0	1	0	0	0	0	0	1	1	1	0	0	0	0	0	0	0	1	0	0	0
Pop2,b6,MSP	0	1	0	0	1	1	0	1	1	0	1	0	0	0	0	0	0	1	0	1	1

Figure 2: Final data format in the .csv file

It is highly recommended to change the working directory to that where datafile is located. Windows users can use the menu item 'File>Change dir' and choose the appropriate folder. To change the working directory within an R console run the command `setwd(dir)` where *dir* is the absolute path to the directory. The output files created by *msap* will be save in that working directory.

Once we are in the rigth working directory with an appropriate data file, we can run all analyses of *msap* with a single command (change "example.csv" by the name of your datafile, keeping the quotes, and change "Example" by custom name to identify your data):

```
> msap("example.csv", name="Example")
```

On execution, *msap* will run all analyses and will show an on screen text report with the results:

```
msap - Statistical analysis for Methilation-Sensitive Amplification Polimorphism data

Reading example.csv
Number of loci: 701
Number of samples/individuals: 38
Number of groups/populations: 4
Number of Methylation-Susceptible Loci (MSL): 494
Number of No Methylated Loci (NML): 207

Number of polymorphic MSL: 257 ( 52 % of total MSL)
Number of polymorphic NML: 49 ( 24 % of total NML)

Shannon's Diversity Index
MSL: I = 0.5836821 (SD: 0.1035363 )
NML: I = 0.3299215 (SD: 0.1078034 )
Wilcoxon rank sum test with continuity correction : W = 11710 ( P < 0.0001 )
```

```
*****
```

```
Analysis of MSL
```

```
Report of methylation levels
```

	pop1
HPA+/MSP+ (Unmethyalted)	0.1917
HPA+/MSP- (Hemimethylated)	0.1706
HPA-/MSP+ (Internal cytosine methylation)	0.2156
HPA-/MSP- (Full methylation or absence of target)	0.4221
	pop2
HPA+/MSP+ (Unmethyalted)	0.1688
HPA+/MSP- (Hemimethylated)	0.1432
HPA-/MSP+ (Internal cytosine methylation)	0.1430

HPA-/MSP- (Full methylation or absence of target) 0.5450  
 pop3  
 HPA+/MSP+ (Unmethylated) 0.1601  
 HPA+/MSP- (Hemimethylated) 0.1599  
 HPA-/MSP+ (Internal cytosine methylation) 0.2298  
 HPA-/MSP- (Full methylation or absence of target) 0.4502  
 pop4  
 HPA+/MSP+ (Unmethylated) 0.1670  
 HPA+/MSP- (Hemimethylated) 0.1281  
 HPA-/MSP+ (Internal cytosine methylation) 0.1589  
 HPA-/MSP- (Full methylation or absence of target) 0.5460

#### Performing AMOVA

AMOVA TABLE	d.f.	SSD	MSD	Variance
among groups	3	1.332	0.4439	0.03146
within groups	34	4.961	0.1459	0.1459
Total	37	6.293	0.1701	

Phi\_ST = 0.1773 (P<0.0001)

#### Pairwise Phi\_ST

```

-----
pop1 - pop2 : 0.2971          (P<0.0001)
pop1 - pop3 : 0.05566        (P= 0.024 )
pop1 - pop4 : 0.2341          (P<0.0001)
pop2 - pop3 : 0.203           (P<0.0001)
pop2 - pop4 : 0.09294        (P= 0.0006999 )
pop3 - pop4 : 0.1518          (P<0.0001)

```

\*\*\*\*\*

#### Analysis of NML

#### Performing AMOVA

AMOVA TABLE	d.f.	SSD	MSD	Variance
among groups	3	0.3587	0.1196	0.006628
within groups	34	1.931	0.05679	0.05679
Total	37	2.29	0.06188	

Phi\_ST = 0.1045 (P<0.0001)

#### Pairwise Phi\_ST

```

-----
pop1 - pop2 : 0.2213          (P<0.0001)
pop1 - pop3 : 0.02315        (P= 0.119 )
pop1 - pop4 : 0.1757          (P<0.0001)

```

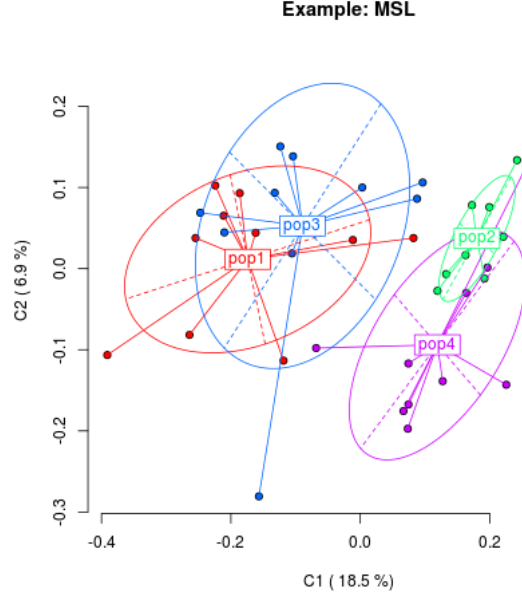


Figure 3: Representation of Principal Coordinate Analysis (PCoA) for epigenetic (MSL) differentiation between groups. The first two coordinates (C1 and C2) are shown with the percentage of variance explained by them. Different point types represent individuals from different groups. Group labels show the centroid for the points cloud in each group. Ellipses represent the average dispersion of those points around their centre. The long axis of the ellipse shows the direction of maximum dispersion and the short axis, the direction of minimum dispersion.

pop2	-	pop3	:	0.08653	(P= 0.0013 )
pop2	-	pop4	:	0.0206	(P= 0.07849 )
pop3	-	pop4	:	0.07359	(P= 0.009799 )

In addition to the above text report, *msap* produces some exploratory figures that are directly saved into .png files:

- A plot of Principal Coordinate Analysis (PCoA), see Figure 3, showing the first two axes. They are saved as '<name>-MSL.png' and '<name>-NML.png' for MSL and NML respectively, where <name> represents the name passed as argument in the calling to *msap* function.
- A neighbor-joining tree of all samples (see Figure 4) saved as '<name>-MSL-NJ.png' and '<name>-NML-NJ.png' for MSL and NML respectively.

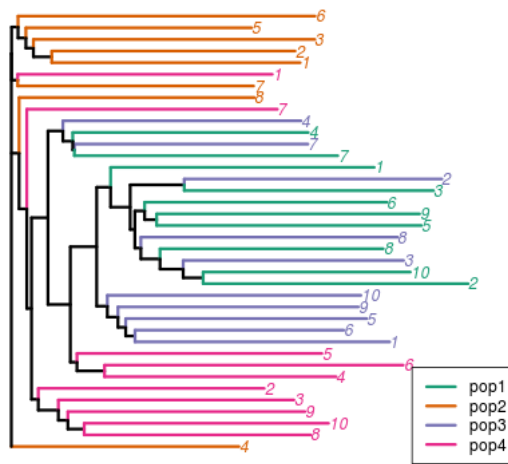


Figure 4: Neighbor-Joining tree of all samples (numbered labels at the tips) for epigenetic (MSL) distances. Colors represent different groups/populations.

## 5.1 Further options

In the previous section, the basic use of *msap* was described. However, it is possible to set some different options in the program if passed as arguments to the `msap()` function.

Here is the full usage of `msap()` function including all the arguments and their default values (if applicable). Except for the 'datafile' that is required, all other arguments are optional.

```
msap(datafile, name=datafile, uninformative=TRUE,  
nDec=4, meth=TRUE, rm.redundant=TRUE,  
rm.monomorphic=TRUE, do.pcoa=TRUE, do.shannon=TRUE,  
do.amova=TRUE, do.pairwisePhiST=TRUE, do.cluster=TRUE,  
use.groups=NULL)
```

**datafile** String containing the url of the csv file with the data. Required.

**name** a name for the dataset to be included in the output files. By default, the name of the given datafile is used.

**uninformative** A logical value determining how to deal with HPA-/MSP- pattern. 'FALSE' assumes that HPA-/MSP- (no band for both isoschizomers) pattern represents full methylation of cytosines in the target, while 'TRUE' (default value) consider that pattern as uninformative as could be caused by a missing target (mutation). See 'Details' below.

**nDec** number of digits of precision for floating point output.

**meth** Logical value switching between MSAP ('TRUE') and standard AFLP ('FALSE') analysis. The difference lies in that for AFLP (`meth=FALSE`) the 'enzyme' column is ignored and every row in data represent an independent sample, without combination of data.

**rm.redundant** Not implemented yet.

**rm.monomorphic** Logical value switching between the removal ('TRUE', by default) of monomorphic fragments (defined as those with only one state or just one occurrence of the second state across the whole dataset) after data transformation.

**do.pcoa** Option switcher for doing a Principal Coordinate Analysis for variation between groups. TRUE by default.

**do.shannon** Option switcher for Shannon's Diversity Index comparison between MSL and NML.

**do.amova** Option switcher for doing an AMOVA for differentiation between groups. TRUE by default.



**do.pairwisePhiST** Logical value switching between the calculation of the pairwise  $\Phi_{st}$  between pairs of groups/populations ('TRUE' by default) or skip it ('FALSE').

**do.cluster** Calculates and plots a Neighbour-Joining tree ('TRUE' by default) or skip it ('FALSE').

**use.groups** Gives the groups/populations/treatments of the datafile to be analysed. By default all groups are considered into the analysis. To provide a subset of the groups a vector should be passed with the names of groups to be included. For example, in a datafile with 5 groups (Control, pop1, pop2, pop3 and pop4) we are interested only in Control and pops 1 and 3. Then, *msap* should be called with 'use.groups=c('Control','pop1','pop3')'.

## 5.2 What if I need another kind of analysis for my MSAP data?

The analyses currently provided by *msap* are limited but that does not mean that further analysis could be added in the future. I try to keep the package updated to allow exploratory assays of epigenetic diversity and differentiation, focused on the field of evolutionary ecology. If you are trying to analyse your MSAP data and options in *msap* do not fit your requirements, then you have three alternatives to get your tasks done by *msap*:

- If you have programming skills or experience with R, then you can get the source code of *msap* and adapt it to your needs. That is the main advantage of open source software!
- If you have programming skills or experience with R, and want to collaborate with me to expand *msap*, then do not hesitate to contact me for joining to the development team.
- If you are not familiar with R or do not feel confident to make code yourself, then use the support tracker to request new features in *msap*.

## References

- Bonin, A., Ehrlich, D., and Manel, S. (2007). Statistical analysis of amplified fragment length polymorphism data: a toolbox for molecular ecologists and evolutionists. *Molecular ecology*, 16(18), 3737-58.
- Chwedorzewska, K. J., and Bednarek, P. T. (2012). Genetic and epigenetic variation in a cosmopolitan grass *Poa annua* from Antarctic and Polish populations. *Polish Polar Research*, 33(1), 63-80.
- Gupta, V., Bijo, J., Kumar, M., Reddy, C. R. K., and Jha, B. (2012). Detection of Epigenetic Variations in the Protoplast-Derived Germings of *Ulva*

- reticulata Using Methylation Sensitive Amplification Polymorphism (MSAP). *Marine biotechnology* DOI: 10.1007/s10126-012-9434-7
- Herrera, C. M., and Bazaga, P. (2010). Epigenetic differentiation and relationship to adaptive genetic divergence in discrete populations of the violet *Viola cazorlensis*. *The New phytologist*, 187(3), 867-76.
- Moran, P., and Perez-Figueroa, A. (2011). Methylation changes associated with early maturation stages in the Atlantic salmon. *BMC genetics*, 12(1), 86.
- Rodriguez, C. M., Moran, P., Lago, F., Beckmann, M., and Consuegra, S. (2012). Detection and quantification of tissue of origin in salmon and veal products using methylation sensitive AFLPs, *Food Chemistry* 131, 1493-1498.

## 6 Session Info

This document was created using the following:

```
> sessionInfo()

R version 2.15.0 (2012-03-30)
Platform: i686-pc-linux-gnu (32-bit)

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=en_US.UTF-8  LC_MESSAGES=en_US.UTF-8
 [7] LC_PAPER=C               LC_NAME=C
 [9] LC_ADDRESS=C             LC_TELEPHONE=C
[11] LC_MEASUREMENT=en_US.UTF-8 LC_IDENTIFICATION=C

attached base packages:
[1] grid      stats      graphics  grDevices utils
[6] datasets  methods  base

other attached packages:
[1] msap_0.1-2      cba_0.2-9      proxy_0.4-7
[4] pegas_0.4-2     adegenet_1.3-4 MASS_7.3-16
[7] ape_3.0-3       socrime_1.2.8  ade4_1.4-17

loaded via a namespace (and not attached):
[1] gee_4.13-18     lattice_0.20-6 Matrix_1.0-6
[4] nlme_3.1-103    tools_2.15.0
```