

This guide helps you get started with rSalvador. For more advanced usage, you should consult the accompanying technical manual or use the inbuilt help facility. To get inbuilt help for a function, e.g., `newton.LD`, type the R command `?newton.LD`.

## 1 Installing rSalvador on the Windows Platform

- Download and install R 3.0.1 or higher version on your computer.
- Make a new folder as your working folder. We recommend naming your working folder `c:/rsalvador`. Copy the rSalvador distribution file `rsalvador_1.4.zip` into that new folder.
- Invoke R. rSalvador requires the R package `hypergeo`, which you can install by the R command

```
install.packages('hypergeo')
```

- rSalvador also requires the R package `gdata`, which in turn requires the `perl` software. However, most Windows computers do not have the software `perl` pre-installed. We recommend Strawberry Perl, which is available on the Internet for free download. Choose the stable version to download and accept all defaults during installation. After installing `perl`, install the R package `gdata` with the R command:

```
install.packages('gdata')
```

- Users not interested in the Excel file import capability can skip the installation of both `gdata` and `perl`. In this case, you can ignore the harmless warning messages about the unavailability of `perl` and `gdata` that you may see each time you load rSalvador.
- Use the R command `setwd('c:/rsalvador')` to set the new folder as your working directory. If your working folder has a different name, replace `'c:/rsalvador'` with the actual name. You can verify your working directory with the R command `getwd()`. You can check whether the rSalvador distribution file `rsalvador_1.4.zip` is present in your working folder by executing `list.files()`. Finally, install rSalvador as follows:

```
install.packages('rsalvador_1.4.zip')
```

## 2 Installation on the Linux Platform

To begin with, install R and the two R packages (`hypergeo`, `gdata`) as described in the above section. Now, follow the following five steps:

1. Make an installation directory and copy the installation file `rSalvadorV1.4.tar` into the installation directory. For example:

```
mkdir myrsalvador
```

2. Untar the installation file that is already in your installation directory (e.g., `~/myrsalvador`):  

```
tar xvf rSalvadorV1.4.tar
```

3. A sub-directory `rsalvador` is now created under the installation directory (e.g., `~/myrsalvador`), but continue to stay in the installation directory. From the installation directory, perform the following checking process (optional but recommended):

```
R CMD check rsalvador
```

4. If no error messages are generated (except for a few benign warning messages), you are ready to build `rSalvador`:

```
R CMD build rsalvador
```

5. Finally, install `rSalvador` by executing

```
R CMD INSTALL rsalvador
```

## 3 Staring `rSalvador`

Each time you invoke an R session, you need to load `rSalvador` with the R command `library(rsalvador)`.

## 4 Your First `rSalvador` Calculations

Here we use the `Demerec` data to demonstrate the basic capabilities of `rSalvador`. To view this data set, type `demerec.data`.

- to find the maximum likelihood estimate of  $m$ :

```
newton.LD(demerec.data)
```

- to find the 95% confidence interval for  $m$ :

```
confint.LD(demerec.data)
```

- To view the iteration details:

```
newton.LD(demerec.data, show.iter=TRUE)
```

- to display the log-likelihood function:

```
plot.likelihood.LD(demerec.data)
```

## 5 Importing and Exporting Data

There are three ways to transfer data into rSalvador.

- creating a sequence of numbers within R:

```
y=c(0, 16, 20, 2, 2, 56, 3, 361, 9)
```

- importing data from a text file. This data file should have just one column of numbers. It can have one or more memo lines. (Use built-in help for details.)

```
y=import.text.data('example1.txt')
```

- Typically, you may have saved your data in an Excel spreadsheet file, like the accompanying example file `example2.xlsx`. (See built-in help for details.) To import data from that file, type

```
y=import.excel.data('example2.xlsx')
```

- Now you can repeat the calculations in the above section by replacing `demerec.data` with the new data variable `y`.
- Occasionally it may be desirable to save your data (say in `y`) into a plain text file for future use. This can be done by

```
export.text.data('mytest.txt', y)
```

To read this data file back into rSalvador for analysis:

```
import.text.data('mytest.txt')
```

## 6 Adjusting for Plating efficiency

We use data from experiment #16 of Luria and Delbruck as an example. This experiment has a plating efficiency of 0.4. You can view the data by typing `luria.16.data` and learn more by typing `?luria.16.data`.

- to find the maximum likelihood estimate of  $m$ :

```
newton.LD.plating(luria.16.data,e=0.4)
```

- to find the 95% confidence interval for  $m$ :

```
confint.LD.plating(luria.16.data,e=0.4)
```

- To view the iteration details:

```
confint.LD.plating(luria.16.data,e=0.4,show.iter=TRUE)
```

- to view the log-likelihood function graphically:

```
plot.likelihood.LD.plating(luria.16.data,e=0.4)
```

## 7 Adjusting for Relative Fitness

For the sake of illustration, assume that from a fitness assay the experimentalist learns that the relative fitness is  $w = 0.21$  for the Demerec data. To get a maximum likelihood estimate of  $m$ , execute the R command

```
newton.MK(demerec.data,w=0.21)
```

The output is  $\hat{m} = 25.86$ . To set up a 95% CI for  $m$ , execute

```
confint.MK(demerec.data,w=0.21)
```

which yields a 95% CI for  $m$ : (22.98, 28.81).

## 8 Comparison of mutation rates

Two mutation rates can be compared by checking whether the two 84% confidence intervals overlap.

```
confint.LD.plating(crane.data[[1]],e=0.1,alpha=0.16)/3.6e9
confint.LD.plating(crane.data[[2]],e=0.1,alpha=0.16)/3.9e9
```

The two confidence intervals for the two mutation rates overlap, and hence the difference is not significant. If terminal cell population sizes are the same in the two experiments, `compare.LD` performs a likelihood ratio test. In the Newcombe experiments, the difference in  $N_t$  between Experiments F and H are small, so a likelihood ratio test is possible.

```
compare.LD(newcombe.data[[6]],newcombe.data[[8]])
```

The likelihood ratio test statistic is  $\Lambda = 0.0119$  and the  $p$ -value is 0.9130.

If final cell population sizes differ noticeably, a more flexible approach is to conduct a likelihood ratio test (LRT) that accounts for the difference in final cell population size. `rSalvador` provides

three functions for that purpose, namely, `LRT.LD`, `LTR.LD.plating` and `LRT.MK`. All these functions require a value of  $R$ , the ratio of the two cell population sizes. The function names are intended as a mnemonic aid, e.g., `LRT.LD.plating` indicates that the underlying distribution is the usual Luria-Delbrück distribution with plating efficiency less than perfect. Take the Werngren-Hoffner data for example, Here we have plating efficiency  $\epsilon = 0.4$ . Immediately prior to plating, the first experiment had final cell density  $2.3 \times 10^8$  per mL, and the second experiment had final cell density  $0.5 \times 10^8$  per mL. Therefore,  $R=0.5/2.3$ . To compare mutation rates in these two experiments, one executes the following

```
R=0.5/2.3
LRT.LD.plating(wh.data[[1]],wh.data[[2]],R=R,e1=0.4,e2=0.4)
```

which gives  $\Lambda = 15.16$  and  $p = 9.86 \times 10^{-5}$ .

## 9 Accounting for variation in $N_t$

The final cell count  $N_t$  may vary considerably if the initial inoculum  $N_0$  is too small. The gamma mixture model can be used to reduce the potential bias caused by excessive variation in  $N_t$ . An estimate of the coefficient of variation for  $N_t$  is needed.

```
newton.B0(demerec.data,cv=0.2)
```

To get a confidence interval, execute:

```
confint.B0(demerec.data,cv=0.2)
```