The "mc2d" package.

Régis POUILLOT, Marie-Laure DELIGNETTE-MULLER & Jean-Baptiste DENIS

November 7, 2008

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0.1

This documentation is intended for readers with:

- A medium level in R. Please refer to the Manual "An Introduction to R" available with R distribution if needed;
- Some knowledge about Monte-Carlo simulations (its basic principles and its utility) and about Quantitative Risk Assessment (QRA).

This documentation will not described all arguments of the functions. The reference remains the documentation attached to the package.

1 Introduction

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1.1 What is mc2d?

"mc2d" means Two-Dimensional Monte-Carlo ("Monte-Carlo à Deux Dimensions"). This package :

- provides additional distributions;
- provides tools to build One-Dimensional and Two-Dimensional Monte-Carlo Simulations;
- provides tools to study One-Dimensional and Two-Dimensional Monte-Carlo Simulations.

In a previous version, some tools to fit parametric distributions to data were included. These functions being useful for other purposes, they have been placed in a specific package called specdist. These packages are available at the URL http://www.inra.fr/unites/miaj/public/matrisq/jbdenis/outils/welcome.html, and soon on the CRAN: http://lib.stat.cmu.edu/R/CRAN/.

mc2d was built for QRA in the Food Safety domain but it might be used for other domains.

1.2 What is Two-Dimensional Monte-Carlo Simulation (briefly)?

The following text and Figure 1 are adapted from [3] and [4] where this method was used. The major reference for Two-Dimensional Monte-Carlo simulations remains [1].

According to international recommendations, a QRA should reflect the "variability" of the risk and estimate the "uncertainty" of the risk estimate. The "variability" represents the temporal, geographical and/or individual heterogeneity of the risk for a given population. The "uncertainty" is understood as the lack of perfect knowledge of the QRA model structure and parameters.

In order to estimate the natural "variability" of the risk, a Monte-Carlo simulation approach may be useful: the empirical distribution of the risk within the population may be estimated from the mathematical combination of distributions reflecting the variability of parameters in the population.

In order to estimate the "uncertainty" of the risk estimates issued from data uncertainty, a two-dimensional (or second-order) Monte-Carlo simulation was proposed [1]. A two-dimensional Monte-Carlo simulation is a Monte-Carlo simulation where the alea reflectiong "variability" and the alea reflecting "uncertainty" are transfered separately in the simulation, so that "variability" and "uncertainty" of the output may be estinated separately. It may be described as following (see Figure 1):

- 1. The parameters of the model should be divided in three categories: the parameters whose alea reflects "variability only", hereinafter denoted as "variable parameters", the parameters whose alea reflects "uncertainty only", denoted as "uncertain parameters" and the parameters whose alea reflects uncertainty and variability. For this latter category, a hierarchical structure, using "hyper-parameters", should be specified: if a parameter is uncertain and variable, one should be able to specify an empirical or parametric distribution reflecting variability only conditionally on other parameters known with uncertainty. As an example, one should be able to set a structure as $X \mid a, b \sim N(a, b)$, this normal distribution reflecting variability of x conditionally to x and x with, e.g., x and x unif x and x and x unif x uncertainty around the parameters x and x and x unif x and x unif x
- 2. A set of uncertain parameters are randomly sampled from their respective distributions;
- 3. The QRA is performed using a classical (one-dimensional) Monte-Carlo simulation of size N_v , conditionally to these uncertain parameters considered as fixed. This QRA takes into account the variability of all variable parameters, and leads to an empirical density function reflecting the variability of exposure/risk among the population conditionally to the uncertain parameters. Various statistics (e.g. the mean, the standard deviation, some percentiles) of the resulting empirical density are evaluated and stored:
- 4. Steps 2) and 3) are performed a large number of time $(N_u \text{ times})$, leading to $N_u \text{ set of statistics}$;
- 5. As output, the 50th percentile (median) of each statistic is used to establish an estimate of this statistic; the 2.5^{th} and 97.5^{th} percentiles of each statistic are used to establish a 95% credible interval (CI95) of this statistic. The median of the N_u estimated values for each of the 101 estimated percentiles allows us to represent a "variability cumulative distribution" using a graph. This curve is surrounded by the 2.5th and 97.5th percentiles obtained from the N_u estimates of each of the 101 percentiles.

"mc2d" is a set of R functions that will help to develop such two-dimensional Monte-Carlo simulations. The main difference with the procedure described above is that mc2d will use arrays of (at least) two dimensions to derive the results: the first dimension will reflect variability, the second will reflect uncertainty. This document will not develop further the method, but the practical use of mc2d, based on a fictive example.

1.3 A basic example

Quantitative Risk Assessment: Escherichia coli O157:H7 infection linked to the consumption of frozen ground beef in <3 year old children.

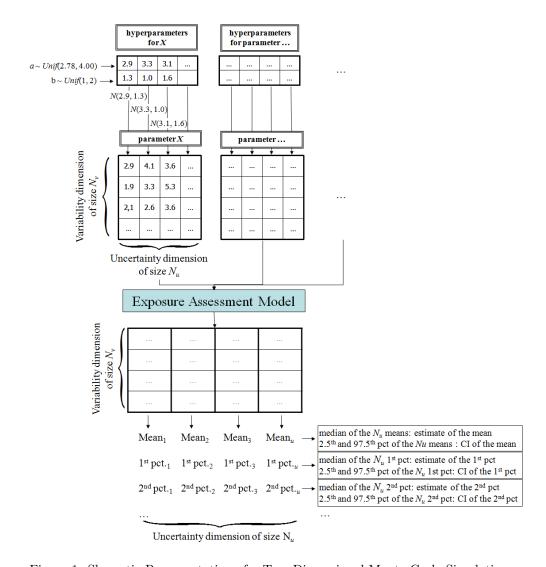


Figure 1: Shematic Representation of a Two-Dimensional Monte-Carlo Simulation.

- We assume that, in a given batch of ground beef, $E.\ coli$ O157:H7 are randomly distributed with a mean concentration of c=10 bacteria (cfu) per gram of product;
- We assume that no bacterial growth occurs, since the product is kept frozen until cooked, just before consumption;
- 2.7% of consumers cook their beef "rare", 37.3% "medium" and 60.0% "well cooked";
- The following inactivation i is associated to these cooking practices:
 - No inactivation for "rare" cooking;
 - 1/5 surviving bacteria for a "medium" cooking;
 - -1/50 surviving bacteria for a "well done" cooking.
- The variability distribution of steak serving sizes s for <3 year children was estimated in a consumption survey. The "best fit" was obtained using a gamma distribution with parameters: shape = 3.93, rate = 0.0806.
- The dose-response relationship, describing the probability of illness P according to the dose is a one hit model. The probability of illness per hit r is assumed to be constant and r = 0.001.

The question is: "What is the distribution of the risk of illness in the population that consumed the contaminated lot?"

This distribution will be estimated using Monte-Carlo simulations performed using R with the "mc2d" package. First, the model will be developed in a one dimensional framework. Then, including some uncertainties in the model, it will be derived in a two dimensional framework.

1.3.1 One Dimensional Monte-Carlo Simulation

In a first step, we assume that no uncertainty exists in our model. All distributions reflects variability. The model may be written as:

```
c = 10.
i \sim emp(\{1, 1/5, 1/50\}, \{0.027, 0.373, 0.600\})
s \sim gamma(3.93, 0.0806)
n \sim Poisson(c \times i \times s)
P = 1 - (1 - 0.001)^{n}
```

where emp(X, P) is the empirical distribution where each value X_i is associated to the probability P_i . We will use a "classical" one dimensional Monte-Carlo simulation, with 1000 iterations. Using the "mc2d" package, the model may be written as:

```
> dose <- mcstoc(rpois, lambda = expo)</pre>
> r <- 0.001
> risk <- 1 - (1 - r)^dose
> EC1 <- mc(cook, serving, expo, dose, risk)
> print(EC1)
     node
             mode nsv nsu nva variate min
                                               mean median
                                                                max Nas type
     cook numeric 1000
                                     1 0.02
                                            0.1165 0.0200
                                                              1.000
                                                                      0
1
                         1
                             1
                                                                           V
2 serving numeric 1000
                             1
                                     1 5.17 48.4451 44.0195 219.976
                                                                           V
     expo numeric 1000
                                     1 1.03 56.2452 14.1530 935.189
                         1
                             1
4
     dose numeric 1000
                        1
                             1
                                     1 0.00 56.0520 15.0000 938.000
                                                                           V
     risk numeric 1000
                             1
                                     1 0.00 0.0507 0.0149
                                                                           V
  outm
1 each
2 each
3 each
4 each
5 each
> summary(EC1)
cook:
               sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
NoUnc 0.116 0.176 0.02 0.02 0.02 0.02 0.2
serving:
             sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
NoUnc 48.4 24.3 5.17 14.5 29.8 44 62.6
                                          103 220 1000
expo:
             sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
NoUnc 56.2 96.8 1.03 3.5 8.11 14.2 79.1
                                           229 935 1000
dose :
             sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
NoUnc
                             15 79
                                       226 938 1000
        56 96.3
                 0
                       2
                           7
risk:
                                  25%
                                         50%
                                               75% 97.5%
                 sd Min 2.5%
                                                           Max nsv Na's
        mean
                      0 0.002 0.00698 0.0149 0.076 0.203 0.609 1000
NoUnc 0.0507 0.0755
```

This One-Dimensional Monte-Carlo simulation provides an estimates of the mean risk (around 5%), as well as some quantiles of the risk (2.5% of the population has a risk of illness greater than 20.3%).

1.3.2 Two dimensional Monte-Carlo Simulation

Assume now that:

• The mean concentration of bacteria in the batch is not known with certainty. It was an estimate. Microbiologists think that the uncertainty around this estimate might be reflected through a normal distribution with parameters $\mu = 10$ and $\sigma = 2$;

• Epidemiological studies suggest that the r parameter is known with uncertainty. The uncertainty around the mean value 0.001 may be reflected through a uniform distribution bounded within 0.0005 and 0.0015.

The model could then be written as:

```
c \sim N(10,2)

i \sim emp(\{1,1/5,1/50\},\{0.027,0.373,0.600\})

s \sim gamma(3.93,0.0806)

n \sim Poisson(c \times i \times s)

r \sim Unif(0.0005,0.0015)

P = 1 - (1 - r)^n
```

Nevertheless, the distributions of r and c do not reflect the same kind of alea then do the distributions of i and s. r and c are uncertain, while i and s are variable. n, as a function of c, i and s will be variable and uncertain.

We will use a two dimensional Monte-Carlo simulation, with 1000 iterations in the variability dimension and 100 iterations in the uncertainty dimension. Using the "mc2d" package, the model may be written as:

```
> ndunc(100)
[1] 100
> conc <- mcstoc(rnorm, type = "U", mean = 10, sd = 2)</pre>
> cook <- mcstoc(rempiricalD, type = "V", values = c(1, 1/5, 1/50),
     prob = c(0.027, 0.373, 0.6))
> serving <- mcstoc(rgamma, type = "V", shape = 3.93, rate = 0.0806)
> expo <- conc * cook * serving
> dose <- mcstoc(rpois, type = "VU", lambda = expo)</pre>
> r < -mcstoc(runif, type = "U", min = 5e-04, max = 0.0015)
> risk <- 1 - (1 - r)^dose
> EC2 <- mc(conc, cook, serving, expo, dose, r, risk)
> EC2
     node
             mode nsv nsu nva variate
                                             min
                                                            median
                                                                         max Nas
                                                     mean
1
     conc numeric
                     1 100
                                      1 5.557709 9.94e+00 9.72e+00 1.75e+01
                                      1 0.020000 1.07e-01 2.00e-02 1.00e+00
     cook numeric 1000
                             1
                                      1 2.665858 4.97e+01 4.50e+01 1.61e+02
3 serving numeric 1000
                             1
                                                                               0
     expo numeric 1000 100
                                      1 0.705346 5.31e+01 1.37e+01 1.68e+03
                                                                               0
                            1
5
     dose numeric 1000 100
                                      1 0.000000 5.31e+01 1.40e+01 1.71e+03
                                                                               0
6
        r numeric
                     1 100
                                      1 0.000514 9.62e-04 9.02e-04 1.48e-03
                                                                               0
                             1
     risk numeric 1000 100
                                      1 0.000000 4.55e-02 1.36e-02 8.40e-01
  type outm
    U each
1
2
     V each
3
    V each
4
   VU each
    VU each
    U each
6
    VU each
```

```
conc :
       NoVar
       9.72
median
mean
        9.94
2.5%
        5.96
97.5% 14.46
cook:
               sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
NoUnc 0.107 0.166 0.02 0.02 0.02 0.02 0.2
                                                   1 1000
serving:
             sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
NoUnc 49.7 24.9 2.67 13.6 31 45 64.2
                                         110 161 1000
expo:
       mean
               sd
                    Min 2.5%
                               25%
                                     50%
                                           75% 97.5%
                                                      Max
                                                           nsv Na's
median 51.9
            94.2 1.234 3.06
                              7.87 13.58
                                          71.8
                                                  240
                                                       938 1000
                                                                   0
            96.3 1.261 3.12
                              8.04 13.89
                                          73.4
                                                  245
                                                       959 1000
                                                                   0
mean
2.5%
       31.8 57.8 0.756 1.87
                              4.82
                                   8.33
                                          44.0
                                                      575 1000
                                                                   0
                                                  147
97.5% 77.2 140.2 1.836 4.55 11.71 20.21 106.8
                                                  357 1396 1000
                                                                   0
dose :
               sd Min 2.5%
                              25% 50%
                                         75% 97.5%
       mean
                                                    Max
                                                          nsv Na's
median 51.9
             94.7 0.00
                          2
                             7.00 14.0
                                        71.8
                                                242
                                                     958 1000
                                                                 0
                             7.53 14.2
                                                                 0
       53.1 96.7 0.04
                          2
                                        73.4
                                                245
                                                    964 1000
mean
                          1 4.47 9.0
2.5%
       31.7 57.8 0.00
                                        43.5
                                                146
                                                    573 1000
                                                                 0
97.5% 77.6 140.7 1.00
                          3 11.00 20.5 107.8
                                                355 1379 1000
                                                                 0
r:
          NoVar
median 0.000902
mean
       0.000962
2.5%
       0.000525
97.5% 0.001459
risk:
         mean
                  sd
                          Min
                                  2.5%
                                            25%
                                                    50%
                                                           75% 97.5%
median 0.0445 0.0703 0.00e+00 0.001713 0.00687 0.01298 0.0645 0.2027 0.589 1000
       0.0455 0.0706 3.88e-05 0.001902 0.00717 0.01347 0.0674 0.2061 0.582 1000
       0.0191 0.0324 0.00e+00 0.000583 0.00282 0.00538 0.0271 0.0841 0.290 1000
2.5%
       0.0730 0.1057 7.08e-04 0.004115 0.01226 0.02242 0.1116 0.3259 0.788 1000
97.5%
       Na's
median
          0
          0
mean
2.5%
          0
97.5%
```

Note that the syntax is similar. Nevertheless, for each distribution, a "type" argument is provided, indicating if the parameter distribution reflects uncertainty (type="U"), variability (type="V"), or both (type="VU").

The summary now provides estimates of the variability distribution (in row) but with a measure of their uncertainty, linked to the uncertainty around conc and r. The estimate of the mean risk is now known with uncertainty. The median of the 100 simulations lead to a "best estimate" of 0.0445, with a "credible interval" of [0.191, 0.0730].

2 Basic Principles and Functions

A classical session of R using "mc2d" is as following:

- From data, expert knowledge, *etc.* empirical or parametric distributions are chosen for each "parents" parameters. For data fitting, the "specdist" package is recommended;
- For each parameter, an mcnode object is built (key functions: mcdata, mcstoc);
- Various mcnode objects are grouped in a mc object (key function: mc).
- The mc object is studied through summaries, graphs, sensitivity analysis (key functions: summary.mc, plot.mc, tornado, tornadounc).

2.1 Before All

The "mc2d" library should be loaded during your R session ("library(mc2d)").

The default size of the Monte-Carlo Simulation should be defined using the ndvar() function (dimension of variability) and the ndunc() function (dimension of uncertainty).

2.2 The mcnode Object as an Elementary Object.

2.2.1 mcnode Object Structure

An mcnode object is the basic element of an mc object. It is an array of dimension $(nsv \times nsu \times nvariates)$ where nsv is the dimension of variability, nsu is the dimension of uncertainty and nvariates is the number of variates of the mcnode¹. Four types of mcnode exist:

- "V" mcnode, for "Variability", are arrays of dimension $(nsv \times 1 \times nvariates)$. The alea in the data reflects variability of the parameter;
- "U" mcnode, for "Uncertainty", are arrays of dimension $(1 \times nsu \times nvariates)$. The alea in the data reflects uncertainty of the parameter.
- "VU" mcnode, for "Variability and Uncertainty", are arrays of dimension (nsv × nsu × nvariates). The alea in the data reflects separated variability (in the first dimension) and uncertainty (in the second dimension) of the parameter.
- Additionally, "0" mcnode are defined for some use. "0" stand for "Neither Variability or Uncertainty". They are arrays of dimension (1 × 1 × nvariates). No alea is considered for these nodes. "0" mcnode are not necessary in the univariate context (use scalar instead) but are useful to build multivariate nodes (See section 3).

There are 5 ways to build a mcnode object:

- 1. The mcstoc function builds mcnode from random generating functions;
- 2. The mcdata function fills mcnode from data sets;
- 3. mcnode are built directly from operations on mcnode objects;
- 4. mcprobtree is a special function that builds an mcnode from various mcnode using a probability tree;
- 5. Some functions, as "==" or ">", is.na, is.finite build an mcnode when applied to an mcnode.

¹In this section, we will only consider mcnode with nvariates = 1.

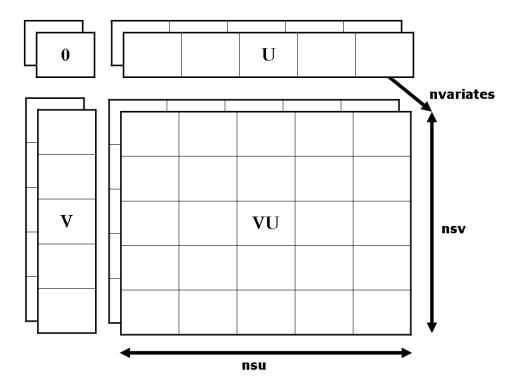


Figure 2: Structure of the various mcnode objects.

2.2.2 The mcstoc function

The mcstoc function is written as 2 :

mcstoc(func=runif, type=c("V", "U", "VU", "0"), ..., nsv=ndvar(), nsu=ndunc(),
nvariates=1, outm="each", nsample="n", seed=NULL, rtrunc=FALSE, linf=-Inf, lsup=Inf,
lhs=FALSE)

- func is a function providing random data or its name as character. The table 1 provides available distributions from the stats and the mc2d libraries that can be used in mcstoc;
- type is the type of requested mcnode. By default, mcstoc builds a "V" mcnode;
- ... are the arguments to be passed to the function func, with the exception of the argument providing the size of the sample. This latter is calculated by the function according to func, type, nsv, nsu and nvariates. If the name of the argument specifying the size of the sample is not n (e.g. functions rhyper and rwilcox, see table 1), the name of this parameter should be provided in the nsample argument. Note that all following arguments should be named;
- nsv and nsu are the number of values needed in the variability and the uncertainty dimension, respectively. By default, these values are the one provided by ndvar() and ndunc(), respectively;
- nvariates is the desired number of variates in the mcnode;
- outm is the default output for multivariate nodes;
- seed optionally specify a seed for the random generator;

²as classically in R, most of arguments have logical default values and will be infrequently modified.

Table 1: Available distributions

Package	Distribution	function	Parameter n	Other Parameters	trunc	lhs
stats	beta	rbeta	n	shape1, shape2, ncp	Y	Y
	binomial	rbinom	n	size, prob	Y	Y
	Cauchy	rcauchy	n	location, scale	Y	Y
	chi-squared	rchisq	n	df, ncp	Y	Y
	exponential	rexp	n	rate	Y	Y
	F	rf	n	df1, df2, ncp	Y	Y
	gamma	rgamma	n	shape, rate (or scale)	Y	Y
	geometric	rgeom	n	prob	Y	Y
	hypergeometric	rhyper	nn	m, n, k	Y	Y
	lognormal	rlnorm	n	meanlog, sdlog	Y	Y
	logistic	rlogis	n	location, scale	Y	Y
	negative binomial	rnbinom	n	size, prob (or mu)	Y	Y
	normal	rnorm	n	mean, sd	Y	Y
	Poisson	rpois	n	lambda	Y	Y
	Student's t	rt	n	df, ncp	Y	Y
	uniform	runif	n	min, max	Y	Y
	Weibull	rweibull	n	shape, scale	Y	Y
	Wilcoxon	rwilcox	nn	m,n	Y	Y
mc2d	Bernoulli	rbern	n	prob	Y	Y
	empirical discrete	rempiricalD	n	values, prob	Y	Y
	PERT	rpert	n	min, mode, max, shape	Y	Y
	triangular	rtriang	n	min, mode, max	Y	Y
	generalised beta	rbetagen	n	shape1,shape2,min,max,ncp	Y	Y
	multinomial	rmultinomial	n	n, size, prob	N	N
	Dirichlet	rdirichlet	n	alpha	N	N
	multinormal	rmultinormal	n	mean, sigma	N	N

- rtrunc allows to truncate the distribution between linf and lsup. This function is not valid for all distributions (see table 1). See the rtrunc function help for further details;
- 1hs allows to sample the node in a Latin Hypercube Sampling framework. This function is not valid for all distributions (see table 1). See the 1hs function help for further details.

In our basic example, mcstoc was used to specify conc (a normal distribution), cook (an empirical discrete distribution), serving (a gamma distribution), and dose (a Poisson distribution). Note that the argument lambda of the Poisson distribution (node dose) is an mcnode.

```
> conc <- mcstoc(rnorm, type = "U", mean = 10, sd = 2)
> cook <- mcstoc(rempiricalD, type = "V", values = c(1, 1/5, 1/50),
+ prob = c(0.027, 0.373, 0.6))
> serving <- mcstoc(rgamma, type = "V", shape = 3.93, rate = 0.0806)
> ...
> dose <- mcstoc(rpois, type = "VU", lambda = expo)
> r <- mcstoc(runif, type = "U", min = 5e-04, max = 0.0015)
> ...
```

A normal distribution with parameters mean = 2, sd = 3, truncated on [1.5, 2] with a Latin Hypercube Sampling could be written³:

³Note that the mean and the standard deviation of the Gaussian distribution are not kept due to the truncation.

For your use in mcstoc, additionnal distributions have been implemented: the Bernoulli distribution (rbern), the empirical discrete distribution (rempiricalD), the PERT distribution (rpert)[5], the triangular distribution (rtriang), the Dirichlet distribution (rdirichlet) and the multinormal distribution (rmultinormal). The multinomial distribution has been adapted (vectorized): rmultinomial (library mc2d) should be used in place of rmultinom (library stats). The empirical discrete (e.g. for bootstrap), the Dirichlet, the multinomial and the multinormal may be used with uncertain and/or variable parameters using multivariate nodes. See section 3.

2.2.3 The mcdata function

Another way to build mcnode object is via the mcdata function, when the data are available.

```
mcdata(data, type=c("V", "U", "VU", "0"), nsv=ndvar(), nsu=ndunc(), nvariates=1,
outm="each")
```

See the documentation associated to this function to see the size/mode of data that can be used to specify an mcnode. The following example place a TRUE value in a "U" node in half of the simulations:

```
> nu <- ndunc()
> tmp <- (1:nu) > (nu/2)
> mcdata(tmp, type = "U")

node    mode nsv nsu nva variate min mean median max Nas type outm
1    x logical 1 100 1 1 0 0.5 0.5 1 0 U each
```

2.2.4 Operations on mcnode

mcnodes are automatically built using operations on mcnode. Rules are built to transfer coherently uncertainty and variability within the model. Logically, the rules are as following (illustrated here with a "+")⁴:

```
• "0" + "0" = "0":
```

- "V" + "U" = "VU": the "U" menode is recycled by row, the "V" menode is recycled classically by column;
- "V" + "VU" = "VU": the "V" mcnode is recycled classically by column;

⁴These rules are not classical R rules of recycling.

```
• "U" + "U" = "U";
```

- "U" + "VU" = "VU": the "U" mcnode is recycled by row;
- "VU" + "VU" = "VU"

Thus, in our example:

```
> ...
> expo <- conc * cook * serving
> ...
> risk <- 1 - (1 - r)^dose</pre>
```

expo is function of a "U" and two "V" mcnode: it is a "VU" mcnode with the variability dimension in row and the uncertainty dimension in column. risk is a function of a "U" and a "VU" node: it is a "VU" node.

2.2.5 The mcprobtree function

The mcprobtree function should be used if a "probability tree" is needed to build an mcnode. Assume that the distribution reflecting the uncertainty on conc was not sure, and that the microbiologists suggest that they are 75% confident that $conc \sim N(10,2)$ but that they are 25% confident that $conc \sim U(8,12)$. This could be written using mcprobtree as⁵:

```
> conc1 <- mcstoc(rnorm, type = "U", mean = 10, sd = 2)
> conc2 <- mcstoc(runif, type = "U", min = 8, max = 12)
> whichdist <- mcstoc(rbern, type = "U", prob = 0.25)
> concbis <- mcprobtree(whichdist, list("0" = conc1, "1" = conc2),
+ type = "U")</pre>
```

mcprobtree could also be used to provide a mixture distribution in the variability dimension.

2.2.6 Other functions to build mcnode

The functions "==", "<", "<=", ">=", ">=", ">", provides an mcnode when applied on a mcnode.

Special functions is.na(x), is.nan(x), is.finite(x), is.infinite(x) are implemented to test if any values are NA (missing data), NaN ("Not A Number"), finite or not.

```
> cook < 1
          mode nsv nsu nva variate min mean median max Nas type outm
 node
     x logical 1000
                                      0 0.975
                                  1
> tmp <- log(mcstoc(runif, min = -1, max = 1))</pre>
> tmp
 node
          mode nsv nsu nva variate
                                      min mean median
                                                             max Nas type outm
     x numeric 1000
                                  1 -8.19 -1.03 -0.699 -0.00167 512
                      1
                          1
> is.na(tmp)
 node
          mode nsv nsu nva variate min mean median max Nas type outm
                                       0 0.512
                                                                 V each
     x logical 1000
                                   1
                                                        1
```

⁵an alternative for whichdist could be whichdist <- mcstoc(rempiricalD, type="U", values=c(0,1), prob=c(75,25))

2.2.7 Building correlation between mcnode

Structural links between set of parameters may be very important in QRA. In mc2d, a correlation structure (in the sense of Spearman) implying 2 or more nodes may be built with the cornode function. This function use the Iman & Conover method [2]. Assume that a study suggests that people that eat their ground beef "rare" eat bigger serving sizes. We could build this relation using:

```
> cornode(cook, serving, target = 0.5, result = TRUE)
output Rank Correlation per variates
variates: 1
[1] 1.0000000 0.3796997 0.3796997 1.0000000
$cook
 node
          mode nsv nsu nva variate min mean median max Nas type outm
    x numeric 1000
                                  1 0.02 0.107
                                                 0.02
$serving
 node
          mode nsv nsu nva variate min mean median max Nas type outm
    x numeric 1000
                      1
                          1
                                  1 2.67 49.7
                                                  45 161
                                                                 V each
```

Note that the resulting correlation (around 0.4) is obviously an approximation in this case, where a discrete distribution (cook: 3 categories) is correlated to a continuous distribution (serving).

It is possible to create such correlation between "V" nodes, between "U" nodes, between "VU" nodes or between one "V" node and some "VU" nodes.

The use of a multinormal distribution (rmultinormal) is another way to create such relationship between nodes.

2.3 The mc Object

Once your mcnode objects are built, one should group them in a single object to study the Monte-Carlo results. The "mc" object is a list of mcnode. There are three ways to build a mc object: using the mc function, the evalment function or within the evalment function.

2.3.1 The mc Function

mc(..., name=NULL, devname=FALSE)

```
... are mcnode or mc objects to be gathered in a mc object. mc value is an mc object with specific methods, e.g. print or summary. In our example, we used:
```

```
> ...
> EC2 <- mc(conc, cook, serving, expo, dose, r, risk)
> print(EC2)
> summary(EC2)
```

2.3.2 The mcmodel and the evalmemod Functions

A model may be written in one step using mcmodel (just a wrap of your model in a function), and then evaluated using evalmcmod. These functions may be used once your model is correct and tested using a small number of iterations. For our example:

```
> modelEC3 <- mcmodel({</pre>
      conc <- mcstoc(rnorm, type = "U", mean = 10, sd = 2)</pre>
      cook \leftarrow mcstoc(rempiricalD, type = "V", values = c(1, 1/5,
           1/50), prob = c(0.027, 0.373, 0.6))
      serving <- mcstoc(rgamma, type = "V", shape = 3.93, rate = 0.0806)
      r \leftarrow mcstoc(runif, type = "U", min = 5e-04, max = 0.0015)
      expo <- conc * cook * serving
      dose <- mcstoc(rpois, type = "VU", lambda = expo)</pre>
      risk \leftarrow 1 - (1 - r)^dose
      mc(conc, cook, serving, expo, dose, r, risk)
+ })
> modelEC3
expression({
    conc <- mcstoc(rnorm, type = "U", mean = 10, sd = 2)</pre>
    cook <- mcstoc(rempiricalD, type = "V", values = c(1, 1/5,</pre>
        1/50), prob = c(0.027, 0.373, 0.6))
    serving <- mcstoc(rgamma, type = "V", shape = 3.93, rate = 0.0806)
    r \leftarrow mcstoc(runif, type = "U", min = 5e-04, max = 0.0015)
    expo <- conc * cook * serving
    dose <- mcstoc(rpois, type = "VU", lambda = expo)</pre>
    risk \leftarrow 1 - (1 - r)^dose
    mc(conc, cook, serving, expo, dose, r, risk)
})
attr(,"class")
[1] "mcmodel"
```

Note that:

- the model is wrapped between "{" and "}";
- any (valid) R code may be placed in the model⁶;
- The model should end by an mc() function.

The model is then evaluated using the evalmcmod function:

```
evalmcmod(expr, nsv=ndvar(), nsu=ndunc(), seed=NULL)
```

The interest lay in the possibility to re-run the model with various dimensions or random seeds in one line.

```
> EC3 <- evalmcmod(modelEC3, nsv = 100, nsu = 10, seed = 666)
> EC4 <- evalmcmod(modelEC3, nsv = 100, nsu = 1000, seed = 666)
```

2.3.3 The mcmodelcut and the evalmccut Functions

If you want to evaluate a high dimension model, R may reach its memory limit. evaluates a 2-dimensional Monte-Carlo model (written with the mcmodelcut function) using a loop, calculates and stores statistics in the uncertainty dimension for further analysis. Readers should refer to the corresponding documentation for further details. Our example would be written as:

 $^{^6}$ If needed, it is possible to make reference to the simulation dimensions using ndvar() and/or ndunc().

```
> modEC4 <- mcmodelcut({</pre>
      {
          cook <- mcstoc(rempiricalD, type = "V", values = c(0,</pre>
+
               1/5, 1/50), prob = c(0.027, 0.373, 0.6))
          serving <- mcstoc(rgamma, type = "V", shape = 3.93, rate = 0.0806)
          conc <- mcstoc(rnorm, type = "U", mean = 10, sd = 2)</pre>
          r \leftarrow mcstoc(runif, type = "U", min = 5e-04, max = 0.0015)
      }
      {
          expo <- conc * cook * serving
          dose <- mcstoc(rpois, type = "VU", lambda = expo)</pre>
          risk <- 1 - (1 - r)^dose
          res <- mc(zero, conc, cook, serving, expo, dose, r, risk)
      }
      ſ
          list(sum = summary(res), plot = plot(res, draw = FALSE),
              minmax = lapply(res, range), tor = tornado(res),
               et = sapply(res, sd))
      }
+ })
> evalmccut(modEC4, nsv = 10001, nsu = 101, seed = 666, progress.bar = TRUE)
```

Note that the use of a tornado function in the model should be avoided since it slows considerably the evalment function. The tornado function will be rewritten in the near future.

2.4 Studying an mc Object

As a reminder, the print function provides a very basic summary of the mc object. It has a digits argument (default: 3). Obviously, other more informative functions are provided in the mc2d package.

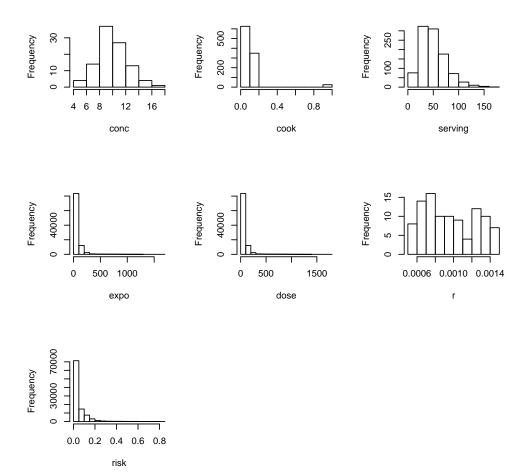
2.4.1 The summary Function

The summary function provide statistics on the mc object:

```
summary(object, probs=c(0,0.025,0.25,0.5,0.75,0.975,1), lim=c(0.025,0.975), ...)
```

The mean, the standard deviation and the quantiles provided in the probs arguments are evaluated on the variability dimension. Then, the median and the quantiles provided in the lim argument are evaluated on these statistics. Of course, these arguments should be changed if other quantiles are needed.

Figure 3: Function hist.



2.4.2 The hist Function

The hist provides an histogram of the different mcnode of the mc object (cf. Figure 3).

In the current version, uncertainty and variability distributions are collapsed. The histogram might be meaningless.

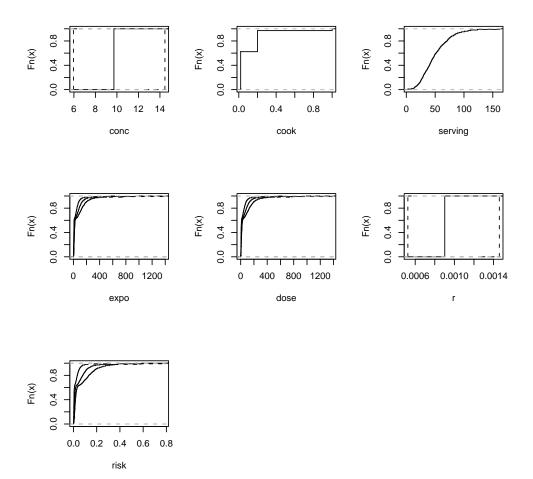
> hist(EC2)

2.4.3 The plot function

The plot function provides a graph of the empirical distribution function of the estimate (mean or median) of the quantiles.

```
plot (x, prec = 0.01, stat = c("median", "mean"), \lim = c(0.025, 0.975), na.rm = TRUE, griddim = NULL, xlab = NULL, ylab = "Fn(x)", main = "", draw = TRUE, ...)
```

Figure 4: plot Function.



For our example, see the Figure 4 as a default graph.

> plot(EC2)

Note that mcnode objects have the same methods print, summary, plot, and hist.

2.4.4 The tornado function

The tornado function builds the Spearman (default) rank correlation between nodes of the mc object.

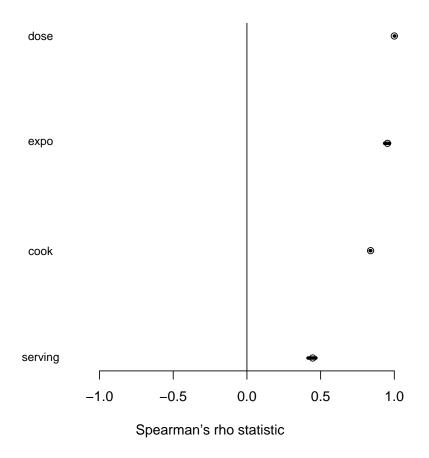
 $\label{tornado} \ \ (x, output=length(x), use="all.obs", method=c("spearman", "kendall", "pearson"), \\ \ \ lim=c(0.025, 0.975))$

where output is the mcnode (name or rank) of the output (default: the last mcnode). Missing data are treated using the use arguments (see the reference documentation). tornado creates a tornado object with a plot method (cf. Figure 5).

> torEC2 <- tornado(EC2)
> plot(torEC2)

18

Figure 5: plot.tornado Function .



2.4.5 The tornadounc function

The tornadounc explore the impact of the uncertainty on the uncertainty of an output. It builds the Spearman (default) rank correlation between statistics of the mc object calculated in the variability dimension.

```
tornadounc(mc,output = length(mc), quant=c(0.5,0.75,0.975), use = "all.obs",
method=c("spearman", "kendall", "pearson"), ...)
```

The quant argument indicates which quantiles should be used in the variability dimension. tornadounc creates a tornadounc object with a plot method

The output shows the impact of the uncertain nodes (type "U" nodes) and some statistics (mean, median and, here, the 99thpercentile) calculated on the variability dimension (type "V" and type "VU" nodes) on some statistics of the output.

2.5 Other Functions and mc Objects

mc objects are simply lists of three dimensional arrays; within each arrays, values in a given columns represent variability of the parameter.

Knowing the structure of the mc and the structure of the mcnode objects, it is direct to apply any R function to these objects. The "\$" function is helpful to extract an mcnode from an mc object, the unmc function removes all attributes, classes, and dimension equal to one, providing a list of vectors, matrices and/or arrays.

Here is a (silly) example building a linear model (in fact ndunc() linear models) between the risk and the dose within each uncertainty dimension and studying some statistics on the coefficients. This example is here only to show you that the whole power of R is available for your analysis.

```
> tmp <- unmc(EC2, drop = TRUE)
> dimu <- ncol(tmp$risk)</pre>
> coef <- sapply(1:dimu, function(x) lm(tmp$risk[, x] ~ tmp$dose[,
      x])$coef)
> apply(coef, 1, summary)
        (Intercept) tmp$dose[, x]
          0.0007991
                         0.0004028
Min.
1st Qu.
          0.0038060
                         0.0005948
          0.0064130
                         0.0007084
Median
          0.0072600
                         0.0007334
Mean
          0.0092290
                         0.0008837
3rd Qu.
          0.0206100
                         0.0011200
Max.
```

3 Multivariate Nodes

The dimension nvariates is the third dimension of the mcnode. One can use mc2d ignoring it. Nevertheless, its use is mandatory to deal with some multivariate distributions, and it may be useful in some circonstances. Building multivariate nodes is direct. We just have to say that the following code:

```
> mcstoc(runif, nvariates = 3, min = c(1, 2, 3), max = 4)
```

will logically not provide a nodes with 3 variates, each having a different limit. The recycling rule tells you that c(1, 2, 3) will be used in the first dimension, i.e. the variability dimension. Use instead:

```
> \lim <- mcdata(c(1, 2, 3), type = "0", nvariates = 3)
> mcstoc(runif, nvariates = 3, min = lim, max = 4)
 node
               nsv nsu nva variate min mean median max Nas type outm
                                                                    V each
1
     x numeric 1000
                       1
                           3
                                    1 1.00 2.54
                                                   2.58
                                                          4
                                                               0
2
     x numeric 1000
                       1
                           3
                                    2 2.00 3.00
                                                   3.00
                                                          4
                                                               0
                                                                    V each
3
     x numeric 1000
                           3
                                    3 3.00 3.52
                                                   3.52
                                                          4
                                                               0
                                                                    V each
```

3.1 Multivariate Nodes for Multivariate Distributions

The basic usage of multivariate nodes (and the reason why it has been implemented) is for multivariate distributions such as the dirichlet distribution, the multinomial distribution, the multinormal distribution and, possibly, the empirical distribution

As an example, assume that 3-member families buy 500 g of ground beef. The proportion of steak eaten by the baby, his older brother and his mom follow a Dirichlet (uncertainty) distribution of parameter $\alpha = (2, 3, 5)$. You want to derive the distribution (variability) of steak eaten by 500 babies issued from these 500 families.

```
(p <- mcstoc(rdirichlet, type = "U", nsu = 100, nvariates = 3,
      alpha = c(2, 3, 5))
  node
          mode nsv nsu nva variate
                                       min mean median
                                                              Nas type outm
                 1 100
                         3
                                  1 0.0164 0.194
                                                  0.168 0.647
                                                                      U each
1
     x numeric
2
     x numeric
                 1 100
                          3
                                  2 0.0389 0.299
                                                  0.283 0.728
                                                                      U each
3
     x numeric
                 1 100
                         3
                                  3 0.1968 0.507 0.512 0.846
                                                                      U each
> s <- mcstoc(rmultinomial, type = "VU", nsv = 500, nsu = 100,
      nvariates = 3, size = 500, prob = p)
> summary(s)
node:
[[1]]
                           2.5%
                                   25%
                                         50%
                                               75% 97.5%
                                                            Max nsv Na's
                sd
                                 78.9
median
                    57.00
                                              89.5 99.8 109.5 500
        84.2
              8.15
                           68.2
                                        84.2
                                                                       0
              8.08
                    73.20
                           81.7
                                  91.6
                                        97.0 102.4 113.1 122.5 500
                                                                       0
mean
2.5%
        12.1 3.42
                     2.95
                             5.9
                                   9.9
                                        11.9 14.5 18.9 23.3 500
                                                                       0
97.5%
       248.9 11.25 214.92 227.0 241.5 249.0 256.0 270.9 285.8 500
[[2]]
                     Min 2.5%
                                  25%
                                        50%
                                              75% 97.5%
                                                          Max nsv Na's
        mean
                sd
median 141.9 10.06 114.0 122.7 135.5 141.5 148.5 161.5 171.0 500
```

```
149.4 9.56 121.3 131.0 142.9 149.4 155.8 168.2 179.2 500
                                                                    0
mean
2.5%
        24.3 4.77 11.4 15.1 20.9 24.4 27.4 33.9 39.3 500
                                                                    0
97.5%
      331.3 11.35 295.3 310.2 324.6 331.7 338.2 351.4 363.1 500
                                                                    0
[[3]]
       mean
               sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
        256 10.78 222 235.0 249 256 264
                                          277 290 500
median
        253 10.65 221 232.8 246 253 261
                                          274 286 500
                                                         0
mean
2.5%
        114 8.83 86 96.4 108 114 121
                                          132 143 500
                                                         0
97.5%
        380 11.59 350 360.0 374 380 386
                                          398 412 500
                                                         0
```

Assume that each member of these families eat a "normal" distribution (variability) of steak with mean 100, 150 and 250 g. There is a positive correlation between the serving of the children, and a negative one with the one of the mother. You want to derive the distribution (variability) of steak eaten by 500 babies.

```
> (x <- mcstoc(rmultinormal, type = "V", nvariates = 3, mean = c(100,
      150, 250), sigma = c(10, 2, -5, 2, 10, -5, -5, -5, 10)))
 node
                                      min mean median max Nas type outm
          mode nsv nsu nva variate
                                     90.7
                                            99.8
1
     x numeric 1000
                      1
                          3
                                   1
                                                    100 109
                                                              0
                                                                    V each
2
     x numeric 1000
                      1
                          3
                                  2 140.1 149.9
                                                    150 160
                                                              0
                                                                    V each
     x numeric 1000
                          3
                                  3 239.5 250.2
                                                    250 261
                                                                    V each
> cor(x[, 1, ])
           [,1]
                      [,2]
                                  [,3]
     1.0000000
                0.1964287 -0.4623170
[2,] 0.1964287 1.0000000 -0.5011393
[3,] -0.4623170 -0.5011393 1.0000000
```

In this example, mean could be variable or uncertain, as well as sigma⁷. You could have used, for an uncertain mean.

The correlation is preserved, but the mean of each categories is known with uncertainty.

Multivariate nodes may finally be useful to derive non parametric bootstrap. Assume that, from a study, you obtained 6 individuals that eat 100 g, 12 individuals that eat 150 g, 6 individuals that eat 170 g and 6 individuals that eat 200 g of ground beef. You want to derive a non parametric bootstrap to derive uncertainty [1], and then pick in the empirical distribution.

⁷Caution: the use of a varying sigma would be very slow.

```
> (x <- mcstoc(rempiricalD, type = "U", outm = c("min", "mean",
      "max"), nvariates = 30, values = c(100, 150, 170, 200), prob = c(6, 100, 100)
      12, 6, 6)))
 node
          mode nsv nsu nva variate min mean median max Nas type outm
1
     x numeric
                 1 100
                        30
                                NA 100
                                         100
                                                100 100
                                                          0
                                                                U min
2
                 1 100
                        30
                                NA 139
                                         154
                                                154 169
                                                          0
                                                                U mean
     x numeric
                 1 100 30
                                NA 200 200
                                                200 200
                                                          0
                                                                U max
     x numeric
> mcstoc(rempiricalD, type = "VU", values = x)
  node
          mode nsv nsu nva variate min mean median max Nas type outm
     x numeric 1000 100
                                   1 100 154
                                                 150 200
                                                               VU each
```

Printing the statistics of the 30 variates of x has no interest. Instead, we use the "outm" option which allows to specify which output we want ("none" for none, "each", the default, for a series of statistics for each variates, or, as in the example, a vector of functions that are applied over all the 30 variates).

3.2 Multivariate Nodes as a "Third Dimension" for Multiple Options in a Model

The recycling rules in mc2d regarding the nvariate dimension is as following: the recycling will be done from nvariates=1 to nvariates=n with n > 1. This allows to use the multivariates nodes as a third dimension, in case you want to test various alternatives.

Assume as in section 2.2.5 that the distribution reflecting the uncertainty on conc was not sure, and that the microbiologists suggest that $conc \sim N(10,2)$ is possible, but that $conc \sim U(8,12)$ is also possible. We can i) build a "bivariate" node reflecting these two options; ii) transfer these options until the final risk estimate. We obtain a bivariate node for the risk, one using the first hypothesis, the second the second hypothesis.

```
> conc1 <- mcstoc(rnorm, type = "U", mean = 10, sd = 2)</pre>
> conc2 <- mcstoc(runif, type = "U", min = 8, max = 12)</pre>
> conc <- mcdata(c(conc1, conc2), type = "U", nvariates = 2)</pre>
> cook <- mcstoc(rempiricalD, type = "V", values = c(1, 1/5, 1/50),
      prob = c(0.027, 0.373, 0.6))
> serving <- mcstoc(rgamma, type = "V", shape = 3.93, rate = 0.0806)
> expo <- conc * cook * serving
> dose <- mcstoc(rpois, type = "VU", nvariates = 2, lambda = expo)</pre>
> r <- mcstoc(runif, type = "U", min = 5e-04, max = 0.0015)
> risk <- 1 - (1 - r)^dose
> EC5 <- mc(conc, cook, serving, expo, dose, r, risk)
> summary(EC5)
conc :
\lceil \lceil 1 \rceil \rceil
       NoVar
median 10.3
        10.2
mean
2.5%
         5.4
97.5%
        14.0
[[2]]
       NoVar
```

```
median 10.14
mean 10.10
       8.08
2.5%
97.5% 11.81
cook:
              sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
      mean
NoUnc 0.117 0.176 0.02 0.02 0.02 0.02 0.2 1 1 1000
serving:
           sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
NoUnc 49.7 25.5 4.47 14.0 31.8 44.5 62.2 109 169 1000
expo:
[[1]]
      mean
              sd
                  Min 2.5%
                            25%
                                  50%
                                        75% 97.5% Max nsv Na's
median 61.2 113.8 1.298 3.36 8.57 15.07 83.7
                                              326 1374 1000
mean 60.4 112.3 1.281 3.32 8.46 14.88 82.6
                                              322 1356 1000
      32.0 59.5 0.679 1.76 4.48 7.88 43.7
                                              170 719 1000
2.5%
97.5% 83.0 154.2 1.760 4.56 11.63 20.44 113.4
                                             442 1863 1000
[[2]]
      mean sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
median 60.1 112 1.28 3.30 8.42 14.8 82.2
                                      320 1350 1000
mean 59.9 111 1.27 3.29 8.39 14.8 81.9 319 1345 1000
2.5% 47.9 89 1.02 2.63 6.71 11.8 65.5 255 1076 1000
97.5% 70.0 130 1.49 3.85 9.81 17.2 95.7 373 1573 1000
dose :
[[1]]
      mean
              sd Min 2.5%
                          25% 50%
                                     75% 97.5% Max nsv Na's
median 61.3 113.5 0.00 2.00 8.00 16.0 82.5
                                          325 1373 1000
mean 60.4 112.6 0.09 2.33 8.11 15.8 81.3
                                          322 1361 1000
                                                            0
      32.2 60.3 0.00 1.00 4.00 9.0 43.6
2.5%
                                           169 700 1000
                                                            0
97.5% 82.9 155.5 1.00 4.00 11.00 21.3 111.5
                                           436 1892 1000
[[2]]
      mean sd Min 2.5%
                         25% 50% 75% 97.5% Max nsv Na's
median 60.3 112 0.00 2.00 8.00 16.0 81.0 324 1340 1000
mean 59.9 112 0.06 2.34 8.05 15.7 80.4 322 1346 1000
     48.1 90 0.00 1.00 6.00 13.0 64.5 262 1069 1000
2.5%
                                                         0
97.5% 69.9 130 1.00 3.00 10.00 19.0 95.0 384 1596 1000
         NoVar
median 0.000921
```

0

mean 0.000950 2.5% 0.000518

97.5% 0.001455

risk:

```
[[1]]
                                   2.5%
                                            25%
                                                    50%
                                                            75% 97.5%
                  sd
                           Min
                                                                        Max nsv
         mean
median 0.0479 0.0761 0.00e+00 0.002156 0.00714 0.01360 0.0688 0.253 0.704 1000
       0.0504 0.0781 8.98e-05 0.002178 0.00763 0.01476 0.0736 0.257 0.694 1000
       0.0262 0.0443 0.00e+00 0.000641 0.00342 0.00746 0.0366 0.139 0.457 1000
       0.0825 0.1184 1.25e-03 0.004039 0.01291 0.02508 0.1246 0.405 0.899 1000
97.5%
       Na's
median
          0
          0
mean
2.5%
          0
97.5%
          0
[[2]]
                                   2.5%
                                            25%
                                                    50%
                                                            75% 97.5%
                  sd
                           Min
                                                                        Max nsv
         mean
median 0.0496 0.0783 0.00e+00 0.002163 0.00732 0.01436 0.0708 0.256 0.710 1000
       0.0506 0.0784 6.48e-05 0.002238 0.00766 0.01486 0.0737 0.260 0.697 1000
mean
2.5%
       0.0268 0.0455 0.00e+00 0.000754 0.00379 0.00753 0.0375 0.139 0.478 1000
97.5%
       0.0800 0.1153 1.07e-03 0.004172 0.01262 0.02446 0.1208 0.407 0.889 1000
       Na's
          0
median
          0
mean
          0
2.5%
97.5%
          0
```

(Do not forget to transfer the number of variates you want in mcstoc... (see the definition of dose). mc2d can not guess...)

3.3 Multivariate Nodes as a "Third Dimension" for Multiple Vectors/Contaminants

The recycling rules in mc2d also allows to use the multivariate nodes as a third dimension for multiple vectors/Contaminants.

Assume in our ground beef example that we have two contaminants: one has a mean concentration that follows an uncertainty distribution $conc \sim N(10,2)$, the second one $conc \sim N(14,2)$. We can i) build a "bivariate" node reflecting these two concentrations⁸; ii) transfer these options until the final dose; iii) sum the dose over the variates (using mcapply). The behavior of contaminants is transferred in the model.

```
> mconc <- mcdata(c(10, 14), type = "0", nvariates = 2)
> conc <- mcstoc(rnorm, nvariates = 2, type = "U", mean = mconc,
+ sd = 2)
> cook <- mcstoc(rempiricalD, type = "V", values = c(1, 1/5, 1/50),
+ prob = c(0.027, 0.373, 0.6))
> serving <- mcstoc(rgamma, type = "V", shape = 3.93, rate = 0.0806)
> expo <- conc * cook * serving
> dose <- mcstoc(rpois, type = "VU", nvariates = 2, lambda = expo)
> dosetot <- mcapply(dose, margin = "variates", fun = sum)
> r <- mcstoc(runif, type = "U", min = 5e-04, max = 0.0015)
> risk <- 1 - (1 - r)^dosetot
> EC6 <- mc(conc, cook, serving, expo, dose, dosetot, r, risk)
> summary(EC6)
```

⁸Note that we could simulate a correlation between both contaminants using a multinormal distribution.

```
conc :
[[1]]
      NoVar
median 10.06
      10.17
mean
2.5%
        6.47
97.5% 14.18
[[2]]
      NoVar
median 14.0
mean
        14.2
2.5%
        10.2
97.5%
       18.3
cook:
              sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
      mean
NoUnc 0.116 0.173 0.02 0.02 0.02 0.02 0.2
                                            1 1 1000
serving:
            sd Min 2.5% 25% 50% 75% 97.5% Max nsv Na's
     mean
NoUnc 48.1 23.9 6.19 12.4 30.0 45.5 61.4
                                         102 171 1000
expo:
[[1]]
                   Min 2.5%
                                    50%
                                          75% 97.5% Max nsv Na's
      mean
              sd
                              25%
median 55.3 96.1 1.246 3.01 8.28 15.05 81.3
                                                239 1017 1000
      55.9 97.1 1.260 3.04 8.37 15.21 82.1
                                                241 1028 1000
                                                                0
mean
2.5%
      35.6 61.7 0.801 1.94 5.32 9.68 52.2
                                                154 654 1000
                                                                0
97.5% 77.9 135.3 1.755 4.24 11.66 21.20 114.5
                                                337 1432 1000
[[2]]
               sd Min 2.5%
                              25% 50% 75% 97.5% Max nsv Na's
        mean
median 77.0 133.6 1.73 4.19 11.51 20.9 113
                                             332 1414 1000
        77.9 135.2 1.75 4.24 11.65 21.2 114
                                             336 1431 1000
                                                             0
        55.8 96.9 1.26 3.04 8.35 15.2 82
                                             241 1025 1000
2.5%
                                                             0
97.5% 100.5 174.5 2.26 5.47 15.03 27.3 148
                                             434 1847 1000
dose :
[[1]]
      mean
              sd Min 2.5% 25% 50%
                                      75% 97.5% Max nsv Na's
median 55.3 96.2 0.00 2.00 8.0 15.5 81.0
                                             236 1010 1000
      55.9 97.3 0.04 2.02 7.9 15.7 81.1
                                             239 1026 1000
                                                             0
mean
       35.7 62.1 0.00 1.00 5.0 10.0 51.5
2.5%
                                             157 636 1000
                                                             0
97.5% 77.7 134.8 1.00 3.00 11.0 21.5 113.3
                                             337 1426 1000
                                                             0
[[2]]
        mean sd Min 2.5% 25% 50%
                                      75% 97.5% Max nsv Na's
median 76.9 134 0.00 3.0 11.0 21.5 113.0
                                            330 1420 1000
        77.9 135 0.14 3.2 11.2 21.6 113.4
                                            335 1431 1000
                                                            Λ
mean
2.5%
        55.6 96 0.00 2.0 8.0 15.5 80.1
                                            235 981 1000
                                                            0
```

97.5% 100.3 174 1.00 4.0 14.0 27.0 146.2

434 1842 1000

0

```
dosetot:
       mean
            sd Min 2.5% 25%
                                50% 75% 97.5%
                                               Max nsv Na's
       132 229 1.00 6.00 19.0 36.0 194
                                           567 2426 1000
median
        134 232 0.98 6.24 19.4 36.4 195
                                           576 2457 1000
                                                            0
mean
2.5%
        113 196 0.00 5.00 16.4 31.0 164
                                           484 2049 1000
                                                            0
        167 289 2.52 8.51 24.0 45.0 242
                                           730 3063 1000
97.5%
                                                            0
r :
          NoVar
median 0.001018
       0.000996
mean
2.5%
       0.000521
97.5% 0.001449
risk:
                  sd
                          Min
                                  2.5%
                                           25%
                                                  50%
                                                         75% 97.5%
                                                                     Max
         mean
median 0.1045 0.1354 0.000858 0.00586 0.01843 0.0354 0.1710 0.432 0.910 1000
       0.1062 0.1352 0.000975 0.00621 0.01916 0.0356 0.1755 0.428 0.888 1000
       0.0614 0.0885 0.000000 0.00307 0.00997 0.0186 0.0973 0.261 0.719 1000
97.5%
      0.1633 0.1866 0.002889 0.01047 0.03259 0.0588 0.2806 0.623 0.983 1000
       Na's
          0
median
mean
          0
          0
2.5%
97.5%
          0
```

As a conclusion, this "third" dimension is highly flexible...

4 Another Example

As a Conclusion

We think and hope that "mc2d" could help risk assessors to build or study their models, and that it may help developing the use of "Two-dimensional" simulation. Nevertheless, "mc2d" is currently under development and

CHECK CAREFULLY YOUR MODEL AND RESULTS TO TRACK THE BUGS.

Please refer any commentary or bugs to rpouillot@yahoo.fr.

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