

Multi-Arm Multi-Stage Trials: Developing the R package "MAMS"

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INTRODUCTION

Multi-Arm Multi-Stage (MAMS) trials are innovative designs that allow the comparison of multiple treatments against a common control across several stages. These trials incorporate adaptations such as stopping for futility or efficacy at either the arm or trial level, enhancing efficiency in identifying promising interventions. Despite the existence of some specialised MAMS software, these tools are often limited in scope and usability. Our project addresses this gap by developing the MAMS R package, which is designed to be flexible, user-friendly, and accessible to users with minimal programming experience.

METHODS

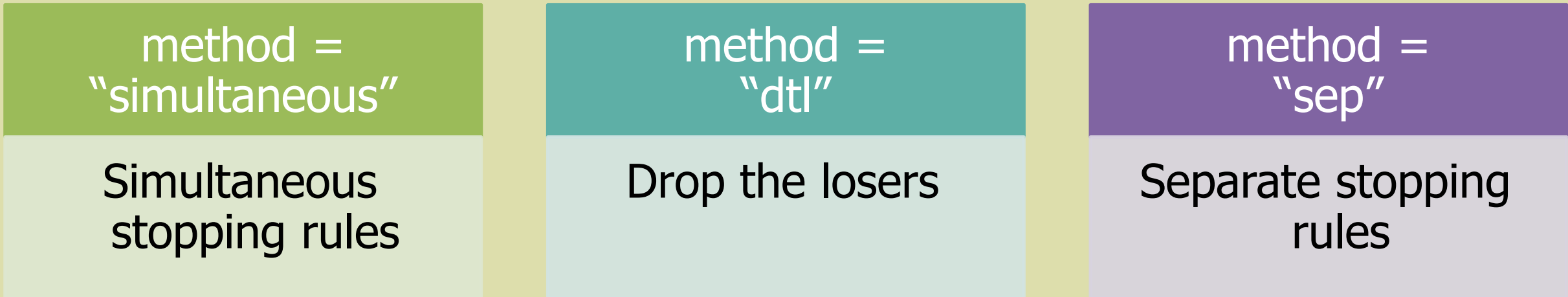
The MAMS R package employs a modular approach, enabling easy integration of various methods for trial adaptation. Currently, the package implements three key methods: Simultaneous stopping, Drop-the-losers design, and separate stopping rule for platform trials.

The modular design ensures that new methods can be seamlessly incorporated into the package as they are developed. This approach enhances the package's adaptability and broadens its applicability to a wide range of MAMS (platform) trial designs.

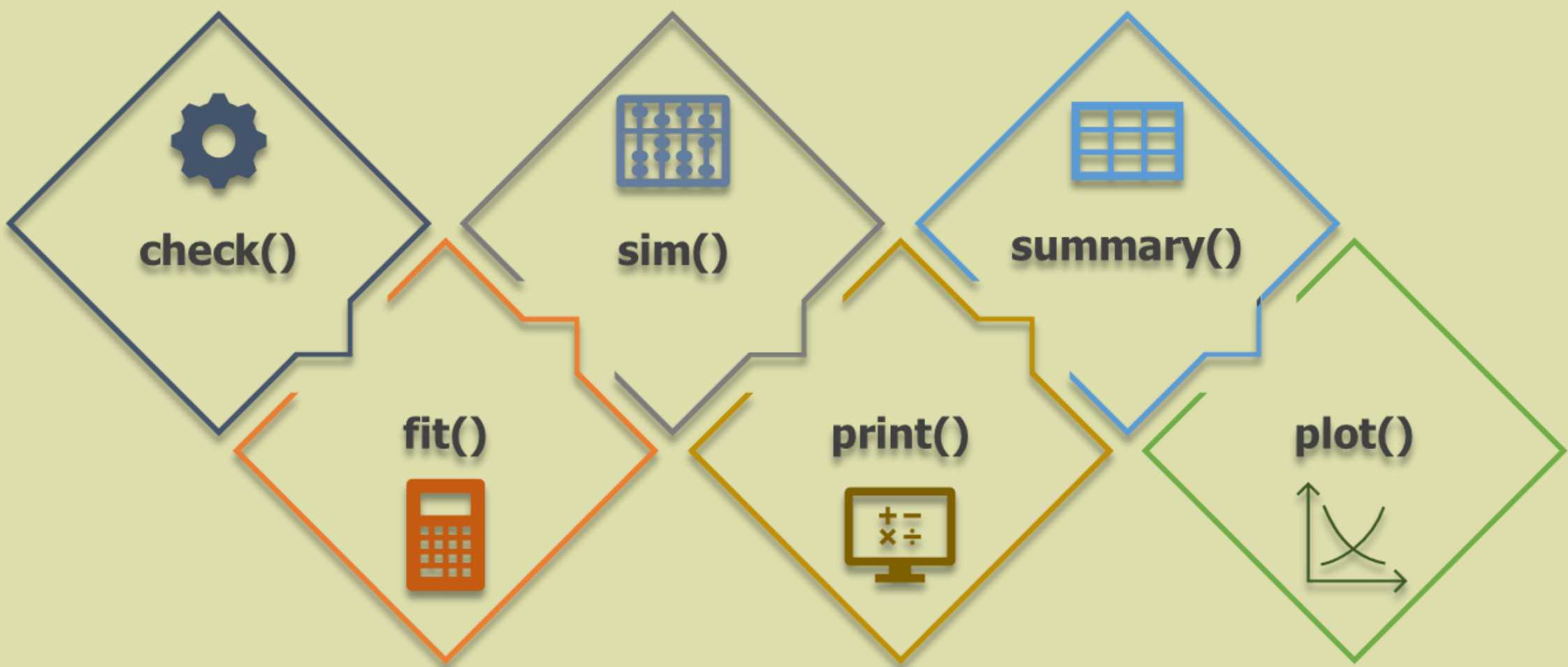
STRUCTURE

Main MAMS package function

mams(method = "module_name", ...)



MAMS module



EXAMPLE APPLICATION

R call: `> mams(method = "dtl")`

Design	Function call	Description
Investigation of 4 experimental treatments in 2 stages with normal endpoints.	K = 4 J = 2	Number of experimental treatments Number of stages
One-sided familywise error rate of 0.05 and desired power of 0.9.	alpha = 0.05 power = 0.9	One-sided familywise error rate Desired power
Allocation ratios: r = 1:2 for experimental treatments,	r = 1:2	Vector of allocation ratios
r0 = 1:2 for control.	r0 = 1:2	Vector ratio on control
Effect size parameters	p = 0.75	Interesting treatment effect on probability scale
	p0 = 0.5	Uninteresting treatment effect on probability scale
	delta = NULL	Interesting treatment effect on traditional scale
	delta0 = NULL	Uninteresting treatment effect on traditional scale
Fixed lower boundary with shape "obf" for upper boundary.	sd = NULL	Standard deviation
	ushape = obf	Shape of upper boundary
	lshape = fixed	Shape of lower boundary
	ufix = NULL	Fixed upper boundary
Sample Size and Computational Settings	lfix = 0	Fixed lower boundary
	nstart = 1	Starting point for sample size
	nstop = NULL	Stopping point for sample size
	sample.size = TRUE	Find sample size
Simulations using 50,000 iterations, considering H0 = True.	Q = 20	Number of quadrature points
	nsim = 50000	Number of simulations
	H0 = TRUE	Consider case with all effect sizes set to 0

OUTPUT

```
-- MAMS design -----
-- Design characteristics --

* Normally distributed endpoint
* Drop the losers
* 2 stages
* 4 treatment arms
* 5% overall type I error
* 90% power of detecting Treatment 1 as the best arm
* Assumed effect sizes per treatment arm:

      | Under H1 | Under H0
abbr | cohen.d | prob.scale | cohen.d | prob.scale
Treatment 1 T1 | 0.954 | 0.75 | 0 | 0.5
Treatment 2 T2 | 0.000 | 0.50 | 0 | 0.5
Treatment 3 T3 | 0.000 | 0.50 | 0 | 0.5
Treatment 4 T4 | 0.000 | 0.50 | 0 | 0.5

-- Arms allocation per stage --

      Stage 1 | Stage 2
Control      1 | 1
Treatment    4 | 1

-- Limits --

      Stage 1 | Stage 2 | shape
Upper bounds  NA | 2.055 | dtl
Lower bounds  NA | 2.055 | dtl
```

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-- Sample sizes --

      Cumulative | Expected ($)
      Stage 1 | Stage 2 | Under H1 | Under H0
      low | mid | high | low | mid | high
Control      13 | 26 | 26 | 26 | 26 | 26
Treatment 1   13 | 26 | 26 | 13 | 16 | 26
Treatment 2   13 | 26 | 13 | 13 | 16 | 26
Treatment 3   13 | 26 | 13 | 13 | 16 | 26
Treatment 4   13 | 26 | 13 | 13 | 16 | 26
TOTAL†        65 | 91 | 91 | 91 | 91 | 91
† Max cumulative size per arm
‡ Based on arms allocation at each stage
-- Futility cumulated probabilities ($) --
      Under H1 | Under H0
      Stage 1 | Stage 2 | Stage 1 | Stage 2
Treatment 1    0 | 0.077 | 0 | 0.241
Treatment 2    0 | 0.005 | 0 | 0.238
Treatment 3    0 | 0.006 | 0 | 0.235
Treatment 4    0 | 0.006 | 0 | 0.236
ANY            0 | 0.095 | 0 | 0.950
ALL            0 | 0.000 | 0 | 0.000
-- Efficacy cumulated probabilities ($) --
      Under H1 | Under H0
      Stage 1 | Stage 2 | Stage 1 | Stage 2
Treatment 1    0 | 0.902 | 0 | 0.013
Treatment 2    0 | 0.001 | 0 | 0.013
Treatment 3    0 | 0.001 | 0 | 0.012
Treatment 4    0 | 0.001 | 0 | 0.012
ANY            0 | 0.905 | 0 | 0.050
T1 IS BEST     0 | 0.902 | 0 | 0.013
ALL            0 | 0.000 | 0 | 0.000

* Estimated T1 related power ($) = 90.218%, [89.956, 90.478] 95% CI
* Estimated overall type I error ($) = 5.012%, [4.822, 5.204] 95% CI

($) Operating characteristics estimated by a simulation
considering 50000 Monte Carlo samples
```

REFERENCES

Jaki, T., Pallmann, P., & Magirr, D. (2019). The R Package MAMS for Designing Multi-Arm Multi-Stage Clinical Trials. *Journal of Statistical Software*, 88(4), 1–25. doi: 10.18637/jss.v088.i04

Wason J, Stallard N, Bowden J, Jennison C. A multi-stage drop-the-losers design for multi-arm clinical trials. *Statistical Methods in Medical Research*. 2017;26(1):508-524. doi:10.1177/0962280214550759

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