Dose-Finding Clinical Trials in Stan

Kristian Brock, StanCon Helsinki, 30-Aug-18



Not too toxic...but High enough to be effective



Not too toxic...but High enough to be effective



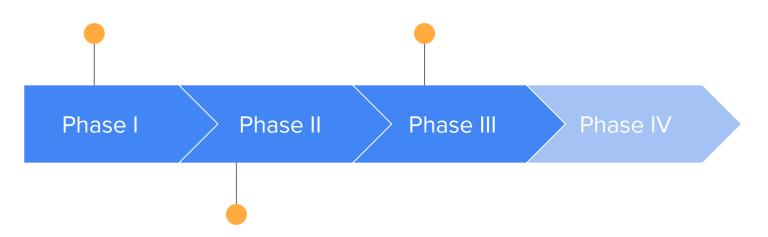
Is it any good?

Does it do something desirable?

Not too toxic...but High enough to be effective

Pivotal study

The challenger vs the standard of care



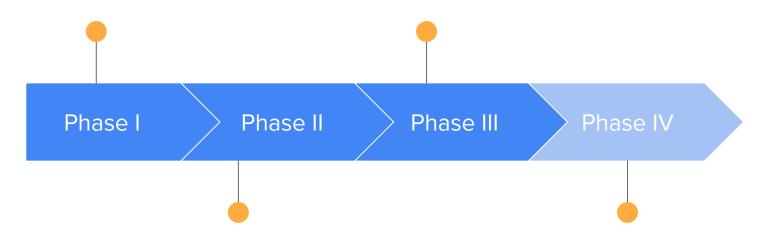
Is it any good?

Does it do something desirable?

Not too toxic...but High enough to be effective

Pivotal study

The challenger vs the standard of care



Is it any good?

Does it do something desirable?

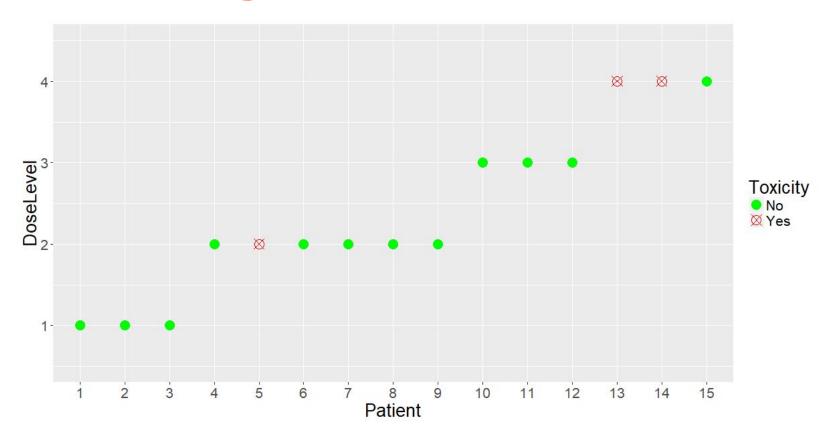
Post-marketing

We'll use it...

But let's keep an eye on it

Dose-finding by toxicity

Dose Finding and the 3+3 Method



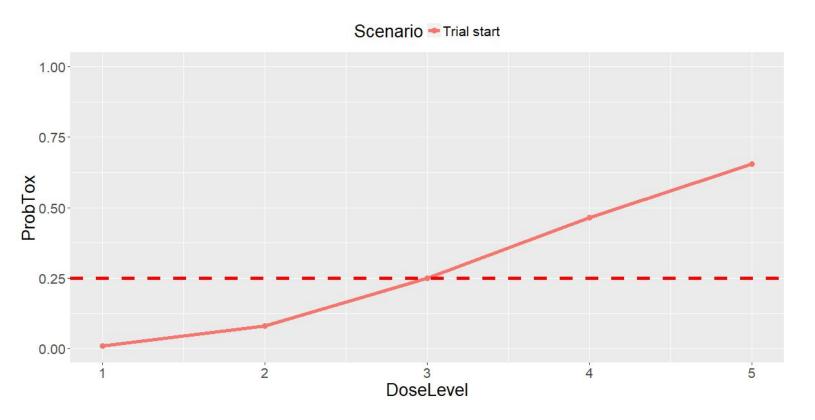
Surely folk use statistical models these days?

- Chiuzan et al. (2017) conducted systematic review of dose-finding methods
- Cancer trials published between 2008 and 2014



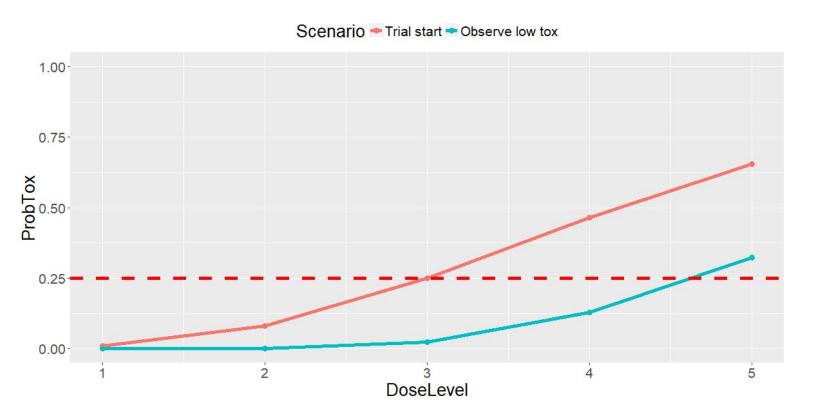
"adoption of [model-based] designs continues to remain low"

Continual Reassessment Method (CRM)



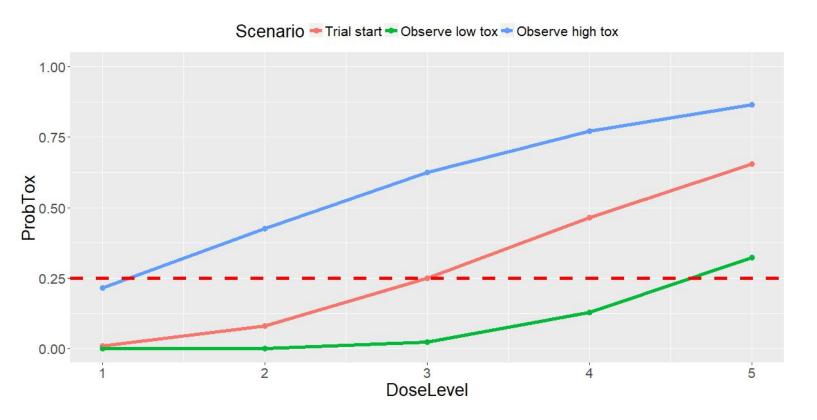
O'Quigley, Pepe, and Fisher (1990). Continual Reassessment Method: A Practical Design for Phase 1 Clinical Trials in Cancer. Biometrics. https://doi.org/10.2307/2531628.

Continual Reassessment Method (CRM)



O'Quigley, Pepe, and Fisher (1990). Continual Reassessment Method: A Practical Design for Phase 1 Clinical Trials in Cancer. Biometrics. https://doi.org/10.2307/2531628.

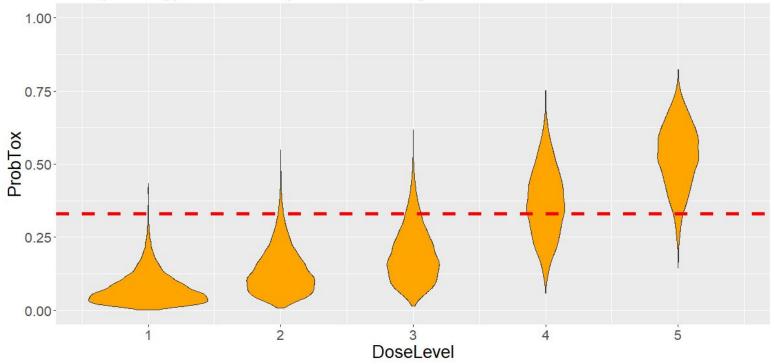
Continual Reassessment Method (CRM)



O'Quigley, Pepe, and Fisher (1990). Continual Reassessment Method: A Practical Design for Phase 1 Clinical Trials in Cancer. Biometrics. https://doi.org/10.2307/2531628.

Levy et al. - ssHHT in leukaemia

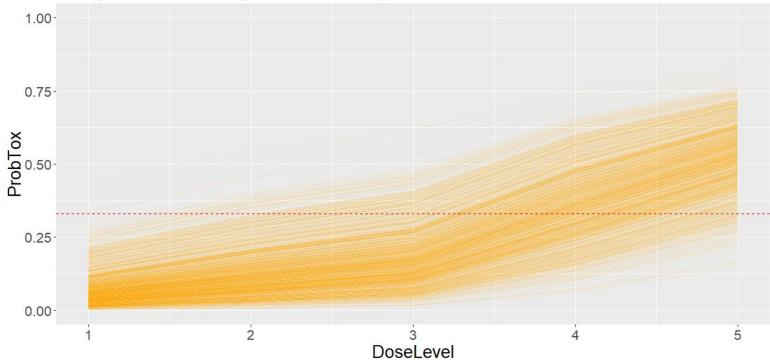
Prob(Toxicity) after all 18 patients in Levy, et al.



Levy, et al. (2006) A Phase I Dose-Finding and Pharmacokinetic Study of Subcutaneous Semisynthetic Homoharringtonine (ssHHT) in Patients with Advanced Acute Myeloid Leukaemia. BJC. https://doi.org/10.1038/sj.bjc.6603265.

Levy et al. - ssHHT in leukaemia

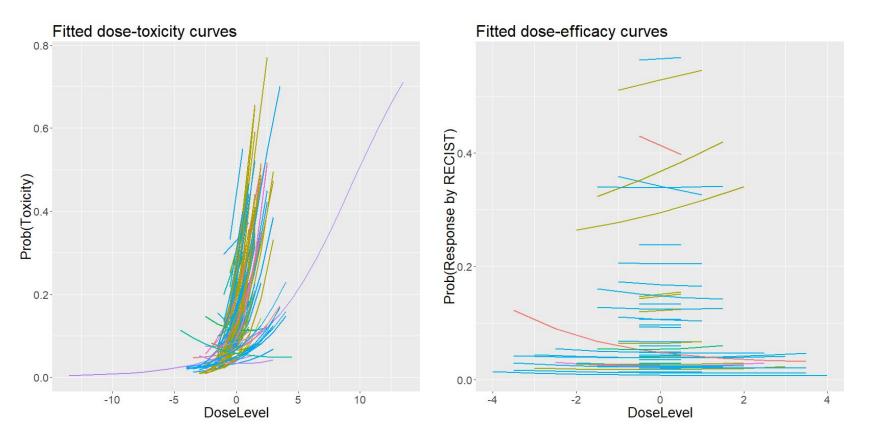
Sampled dose-toxicity curves in Levy, et al.



Levy, et al.(2006) A Phase I Dose-Finding and Pharmacokinetic Study of Subcutaneous Semisynthetic Homoharringtonine (ssHHT) in Patients with Advanced Acute Myeloid Leukaemia. BJC. https://doi.org/10.1038/sj.bjc.6603265.

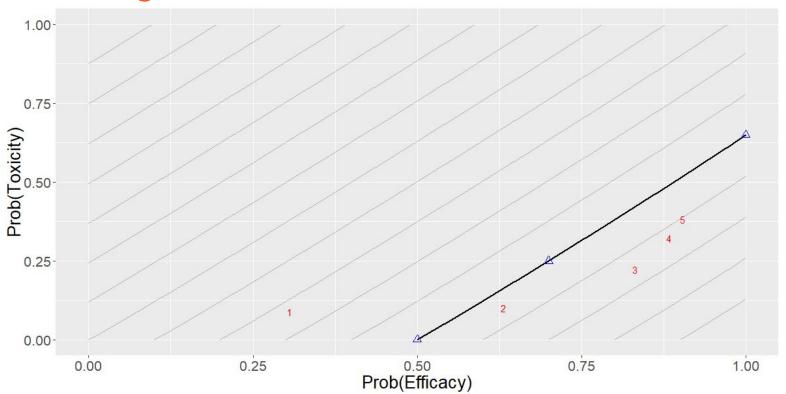
Dose-finding by toxicity & efficacy

Dose-toxicity & efficacy in modern therapies



Brock, et al. (unpublished). Meta-analysis of the dose-toxicity and dose-efficacy curves in the manuscripts in the Chiuzan (2017) review paper.

EffTox - dose-finding based on efficacy & toxicity trade-offs



Thall, PF, and JD Cook. (2004) Dose-Finding Based on Efficacy-Toxicity Trade-Offs. Biometrics 60, no. 3 (2004): 684–93.

Posterior beliefs on attractiveness

Prob("dose in column is more attractive than dose in row")

Dose	1	2	3	4	5
1		0.92	0.86	0.82	0.8
2			0.75	0.68	0.64
3				0.58	0.54
4					0.52
5					

trialr - clinical trial designs in R & Stan

```
# Install
    devtools::install github('brockk/trialr')
    install.packages('trialr')
    # CRM
    levv ← '1NNN 3NNT 4NNT 4NNN 4NTN 4TNT'
    skeleton \leftarrow c(0.05, 0.10, 0.15, 0.33, 0.5)
    target ← 0.33
    levy mod ← stan crm(levy, skeleton = skeleton, target = target,
12
                         model = 'logistic gamma', a0 = 4,
13
                         beta shape = 1, beta inverse scale = 1,
                         seed = 123, control = list(adapt_delta = 0.95))
14
15
    # FffTnx
    outcomes ← '1NEN 2NBE'
    mod2 ← stan_efftox_demo(outcomes, seed = 123)
    # That is short-hand for
    mod2 ← stan efftox(outcomes,
21
                        real doses = c(1.0, 2.0, 4.0, 6.6, 10.0),
22
                        efficacy hurdle = 0.5, toxicity hurdle = 0.3,
23
                        p = 0.1, p t = 0.1,
                        eff0 = 0.5, tox1 = 0.65,
24
25
                        eff star = 0.7, tox star = 0.25.
                        alpha mean = -7.9593, alpha_sd = 3.5487,
26
27
                        beta_mean = 1.5482, beta_sd = 3.5018,
28
                        gamma_mean = 0.7367, gamma_sd = 2.5423,
29
                        zeta mean = 3.4181, zeta sd = 2.4406,
                        eta mean = 0, eta sd = 0.2,
31
                        psi mean = 0, psi sd = 1,
                        seed = 123)
```

Future work:

- Add more trial designs :-)
- Plumb it to work with tidybayes
- More visualisation via bayesplot
- Scrutinise fit by shinystan, etc
- Automated documents via Flexdashboards?

The End

Kristian Brock

Senior Biostatistician
Cancer Research UK Clinical Trials Unit
University of Birmingham
k.brock@bham.ac.uk or
kristian.brock@gmail.com





