RQ: Real-time evidence collection in data streams Updates april 2021

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 - Prof. Floor Scheepers (UMCU)
 - Karin Hagoort (UMCU)
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 - Roel van Est (Parnassia)



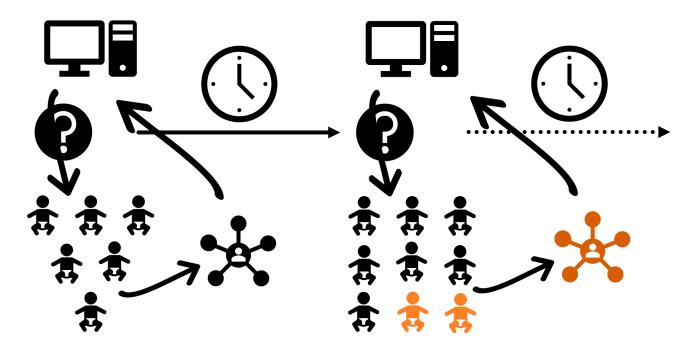






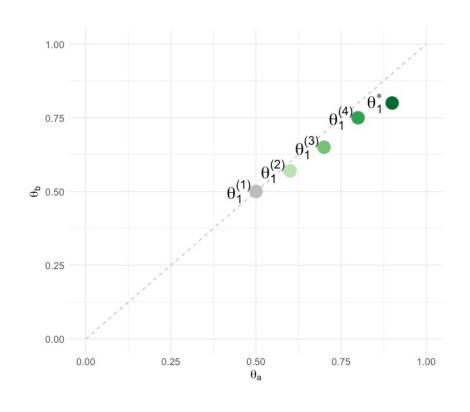
Real-time analysis: safe statistics

Collect and adjust inference about evidence for treatment strategies in real time



Learn from data for adjusting evidence collection process

- With safe statistics can collect "statistically sound" evidence for A/B testing (e.g. treatment recommendations!) in real-time
- Learning approach: take safe test for next data block that is optimal relative to estimate $\hat{\theta}_1$ based on data seen so far
- Technical detail: take $\hat{\theta}_1$ as posterior mean for θ_1 with Beta prior on parameter space



Learn from data for adjusting evidence collection process

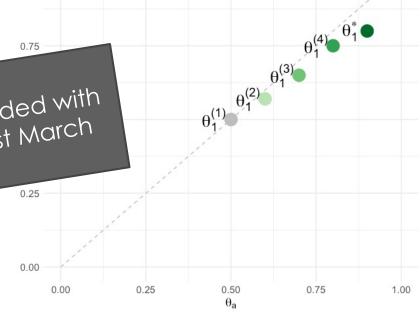
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With safe statistics can collect "statistically sound" evidence for A/B testing (e.g. treatment recommendations!) in real-time

Learning as

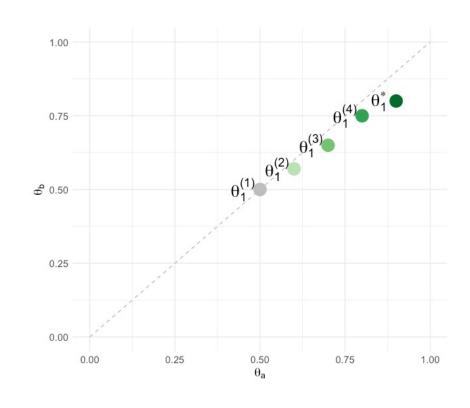
Work partly from master thesis; awarded with
the VVSOR Jan Hemelrijk award last March

posterior mean for θ_1 with Beta prior on parameter space



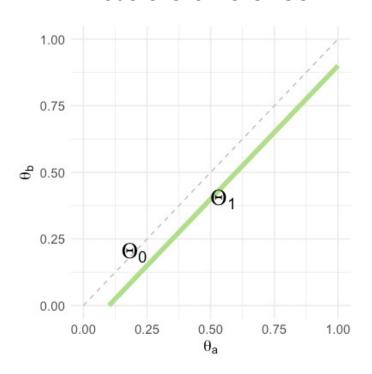
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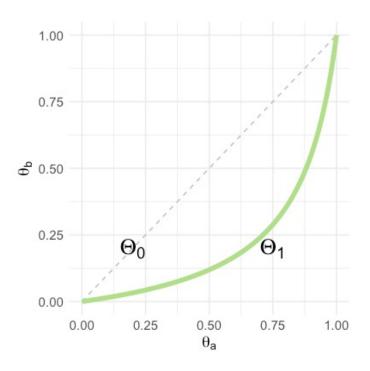


Can also add expert/ prior knowledge: restrict options!

Absolute difference



Odds ratio/ relative risk

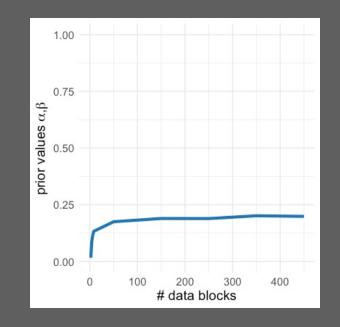


Learn from data for evidence collect

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Progress since last meeting: Beta priors that are "optimal" w.r.t.

REGRET and data collected





Real-life example: SWEPIS¹ perinatal death study

- Comparing perinatal death in labour induction at 41 or 42 weeks
- "All significance tests were two sided at the 0.05 level."

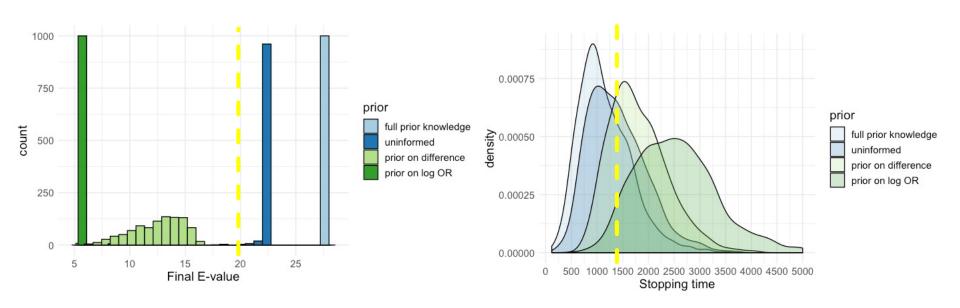
After observing approx. 1380 births in each group:

"On 2 October 2018 the Data and Safety Monitoring Board strongly recommended the SWEPIS steering committee to stop the study owing to a statistically significant higher perinatal mortality in the expectant management group. Although perinatal mortality was a secondary outcome, it was not considered ethical to continue the study. No perinatal deaths occurred in the early induction group but six occurred in the expectant management group (five stillbirths and one early neonatal death; P=0.03)."

Safe testing applied to collect evidence in the SWEPIS scenario

Optionally: use knowledge from previous studies in prior of safe test

- Mean perinatal death rate at 41 weeks: 0.0001
- Difference risk between 42 and 41 weeks: 0.00318



Application to use case: evaluating the usefullness of a recommender system for treatment of depression

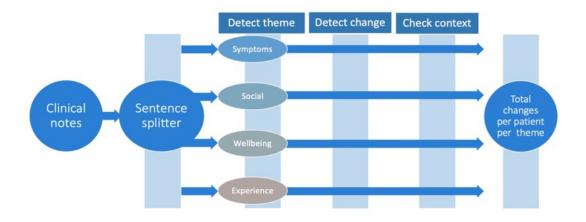
Extracting information on the outcome of treatment trajectories from electronic health records in psychiatry

R.J. Turner^{1,2}, F. Coenen², K. Hagoort², F.E. Scheepers², P.D. Grünwald¹, and A. Härmä³

¹CWI, Amsterdam, NL

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³Philips Research, Eindhoven, NL



Application to use case: evaluating the usefullness of a recommender system for treatment of depression

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NLP part submitted to OCUPAI'21 conference
Full draft will be submitted to JAMA Psychiatry

Experience

Detect change
Check context

Total changes per patient per theme

Application to use case: evaluating the usefullness of a recommender system for treatment of depression

- Which types of recommendations assist clinicians the best?
- Plan: offer, in (micro-)randomized format, different forms of recommendations to clinicians based on the four outcome measures extracted from free text
 - ECT
 - Antidepressants
- Continuously analyze results with safe tests

ARTICLE

Open Access

How machine-learning recommendations influence clinician treatment selections: the example of the antidepressant selection

Maia Jacobs o¹, Melanie F. Pradier¹, Thomas H. McCoy Jr.^{2,3}, Roy H. Perlis^{2,3}, Finale Doshi-Velez¹ and Krzysztof Z. Gajos o¹