

Incentivizing Participation in Randomized Clinical Trials (RCTs)

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Incentivize participation in RCTs

Use **information asymmetry** rather than money.

RCT as a bandit algorithm with T rounds and n arms:

- in each round t , a new patient arrives
- algorithm chooses distribution p_t over arms, draws an arm from p_t , observes outcome $\omega_t \in \Omega$
- patients do not observe previous arms & outcomes, RCT commits to the algorithm (both are **realistic!**)

Goal 1: incentivize each patient to participate (**new!**)

Goal 2: counterfactual estimates (**standard**)

Model: patients and their incentives

Heterogenous agents with subjective utilities:

- public type θ_{pub} : medical history, affects outcomes;
- private type θ_{pri} : subjective utilities over the outcomes, strategically reported by the patient

Patients' beliefs (*not reality!*):

- type $(\theta_{\text{pub}}, \theta_{\text{pri}})$ is drawn from fixed distribution F_{type} ;
- outcome is drawn from fixed distribution $\psi^*(\text{arm}, \theta_{\text{pub}})$;
- ψ^* is **state**, initially drawn from known prior \mathcal{P} .

The algorithm must incentivize each patient to:

- report θ_{pri} (Bayesian Incentive-Compatibility, **BIC**)
- participate (Bayesian Individual Rationality, **BIR**)

Model: statistics (standard for RCTs)

- initially, an **adversary** chooses, for each patient t : public & private types, outcome $\omega_{a,t} \in \Omega$ for each arm a
- each outcome $\omega \in \Omega$ assigned a known **score** $\text{score}(\omega) \in [0, 1]$
- **average score** over a subset S of rounds:

$$f_{\text{adv}}(\text{arm } a) := \frac{1}{|S|} \sum_{t \in S} \text{score}(\omega_{a,t}).$$

- initially, algorithm chooses $S \subset T$; after round T , it computes estimators $\hat{f}(\cdot)$ for $f_{\text{adv}}(\cdot)$, for all arms.

- Objective: maximal mean-squared error

$$\text{MaxMSE} = \max_{\text{all arms } a, \text{ all adversaries}} \mathbb{E} [(\hat{f}(a) - f_{\text{adv}}(a))^2],$$

where $\mathbb{E}[\cdot]$ is over the randomness in the algorithm.

- Standard IPS estimators suffice for our algorithm. Then:

$$\text{MaxMSE} \leq |S|^{-2} \cdot \max_{\text{arms } a} \sum_{t \in S} 1/p_t(a).$$

- Incentives aside, **uniform randomization** is best.

Benchmark: what if state ψ^* is known?

- focus on a **single round** of the algorithm
 - policy class Π : input patient's type $(\theta_{\text{pub}}, \theta_{\text{pri}})$ and state ψ , output distribution over arms, satisfy BIC and BIR.
 - **Informally**: policy $\pi \in \Pi$ and state ψ determine **MaxMSE**;
- $$\text{bench}(\mathcal{P}, F_{\text{type}}) := \inf_{\pi \in \Pi} \sup_{\psi \in \text{support}(\mathcal{P})} \text{MaxMSE}(\pi, \psi).$$
- homogeneous patients \Rightarrow dependence on F_{type} is not needed.

Our focus: "two-stage algorithms"

- initially, algorithm chooses $T_0 < T/2$.
- **warm-up stage**: first T_0 rounds, collects "warm-up data",
- **main stage**: the remaining rounds $S = [T] \setminus [T_0]$.
- main-stage data ignored in the main stage; "warm-up data" ignored by the estimators $\hat{f}(\cdot)$.
- **Realistic!** all main-stage patients can be treated simultaneously; adapting to data is challenging in practice

Our results (homogeneous patients)

Our algorithm: if "warm-up data" contains $\geq N_{\mathcal{P}}$ samples from each arm, with $N_{\mathcal{P}}$ determined by prior \mathcal{P} (but constant in T)

$$\text{MaxMSE} \leq 2 \cdot \text{bench}(\mathcal{P}) / (T - T_0)$$

- "main stage" approximates **bench**(\mathcal{P}) given "warm-up data"
- "warm-up data" collected via techniques from "incentivized exploration", with T_0 determined by \mathcal{P} (and constant in T)

Impossibility: for any BIR mechanism there is an adversary s.t.

$$\text{MaxMSE} \geq \text{constant} \cdot \text{bench}(\mathcal{P}) / T.$$

Heterogeneous patients: similar results, more complex to state.

Working paper at arXiv:2202.06191

Comparison to the work on "incentivized exploration": ask us!