

Bayesian inference of causal effects with incorrectly measured interference network

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Introduction

- Interference occurs when the potential outcomes of a unit depend on treatments assigned to other units.
- The interference structure can be represented by a network. Nodes are units and edges indicate whether interference is possible between pairs of units.
- An observed network is often assumed to correctly specify the interference structure, and analysis is conditioned on it [4].
- However, accurately measuring the interference network is challenging.
- A misspecified network leads to biased estimation [5].
- Observing an incorrect network may result from:
 - Constraints in data acquisition methods (e.g., measurement errors, edge censoring).
 - Sampling a sub-network from the population network.

Setup and assumptions

- $\mathbf{Z} \in \{0, 1\}^n$ binary treatments; $Y_i(\mathbf{z})$ potential outcomes; \mathbf{Y} observed outcomes; \mathbf{X} covariates; \mathbf{A} observed network.
- Interference via true network \mathbf{A}^* . Assumed to be undirected and unweighted.
- A.1** (Consistency) If $\mathbf{Z} = \mathbf{z}$, then $Y_i = Y_i(\mathbf{z})$.
- Exposure mapping $f(\mathbf{Z}_{-i}, \mathbf{A}_i^*)$ with image space $\mathcal{C} \subseteq \mathbb{R}$.
- A.2** (Neighborhood interference) For any \mathbf{z}, \mathbf{z}' , if $z_i = z'_i$ and $f(\mathbf{z}_{-i}, \mathbf{A}_i^*) = f(\mathbf{z}'_{-i}, \mathbf{A}_i^*)$, then $Y_i(\mathbf{z}) = Y_i(\mathbf{z}')$ w.p.1.
- A.3** (Positivity) $0 < \Pr(\mathbf{Z}_i = z, f(\mathbf{Z}_{-i}, \mathbf{A}_i^*) = c | \mathbf{X} = \mathbf{x}) < 1$, $\forall z \in \{0, 1\}, c \in \mathcal{C}, \forall \mathbf{x}$.

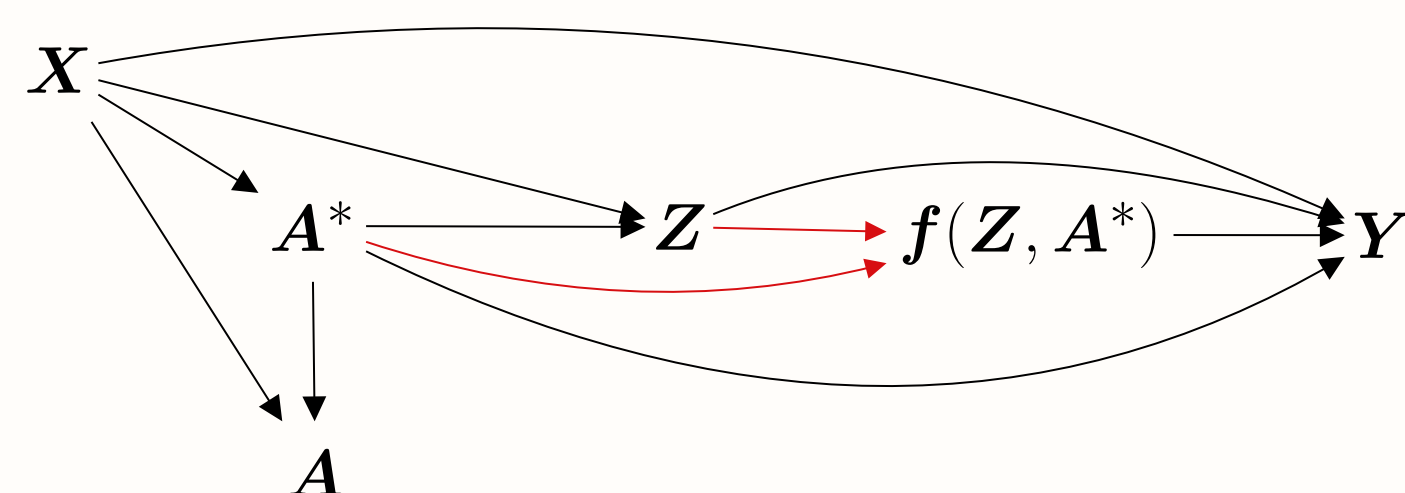


Figure 1. DAG representing the assumed causal structure. Red arrows are deterministic.

Estimands and identification

$\tilde{Y}_i(z, c), z \in \{0, 1\}, c \in \mathcal{C}$ potential outcomes expressed in term of exposure values. From A.2, $\tilde{Y}_i(z, c)$ is equivalent to all $Y_i(\mathbf{z})$ with the same effective treatments.

Causal estimands

Comparisons of the following.

- Exposure levels. $\mathbb{E}[\tilde{Y}_i(z, c)]$.
- Treatment intervention. $\mathbb{E}[Y_i(\mathbf{z})]$.

The first estimand reveals how changes in treatment and exposure values influence outcomes, while the second depicts the effect of setting \mathbf{Z} to \mathbf{z} . Alternatively, conditional estimands, e.g., $\mathbb{E}[Y_i(\mathbf{z}) | \mathbf{A}^*]$, can be used instead.

Identification

- $\mathbb{E}[\tilde{Y}_i(z, c)] = \mathbb{E}_{\mathbf{X}} \mathbb{E}_{\mathbf{A}^* | \mathbf{X}} \mathbb{E}[Y_i | \mathbf{Z}_i = z, f(\mathbf{Z}_{-i}, \mathbf{A}_i^*) = c, \mathbf{A}^*, \mathbf{X}]$.
- $\mathbb{E}[Y_i(\mathbf{z})] = \mathbb{E}_{\mathbf{X}} \mathbb{E}_{\mathbf{A}^* | \mathbf{X}} \mathbb{E}[Y_i | \mathbf{Z} = \mathbf{z}, \mathbf{A}^*, \mathbf{X}]$

- Both requires obtaining $\mathbf{A}^* | \mathbf{X}$ distribution.
- Can be extended to stochastic and network interventions [4].

Generative model and posterior distribution

- True network generation $p(\mathbf{A}^* | \mathbf{X}, \theta)$.
- Observed network (network misspecification model) $p(\mathbf{A} | \mathbf{A}^*, \mathbf{X}, \gamma)$.
- Outcome model $p(\mathbf{Y} | \mathbf{Z}, \mathbf{X}, \mathbf{A}^*, \eta)$.
- Possible to augment outcome model with propensity scores.
- Parameters space can be finite or infinite.
- Assume prior independence $\pi(\eta, \theta, \gamma) = \pi(\eta)\pi(\theta)\pi(\gamma)$.
- Denote observed data by $\mathbf{O} = (\mathbf{Y}, \mathbf{Z}, \mathbf{X}, \mathbf{A})$.

Posterior distribution

$$\pi(\eta, \theta, \gamma, \mathbf{A}^* | \mathbf{O}) \propto p(\mathbf{Y} | \mathbf{Z}, \mathbf{X}, \mathbf{A}^*, \eta) \pi(\eta) \cdot p(\mathbf{A} | \mathbf{X}, \mathbf{A}^*, \gamma) \pi(\gamma) \cdot p(\mathbf{A}^* | \mathbf{X}, \theta) \pi(\theta)$$

Examples of network misspecification models

Measurement error

$p(\mathbf{A} | \mathbf{A}^*, \mathbf{X}, \gamma)$ can be differential or non-differential measurement error model. For instance,

- Random noise.** Edges in \mathbf{A} are observed with true positive rate $1 - \gamma_1$ and false positive rate γ_0 .
- Censoring.** Edges between units with degrees larger than a censoring threshold are missing w.p. γ .

Sampled network

- Study on a sample from a population $n < N$.
- \mathbf{A} is obtained via a network sampling procedure, such as random node, egocentric, or link-tracing sampling.
- Write $\mathbf{A}_o, \mathbf{A}_m$, as the observed and missing parts, respectively.
- Posterior can be written as

$$\pi(\eta, \theta, \mathbf{A}_m | \mathbf{Y}, \mathbf{Z}, \mathbf{X}, \mathbf{A}_o) \propto \pi(\eta) p(\mathbf{Y} | \mathbf{Z}, \mathbf{X}, \mathbf{A}_o, \mathbf{A}_m, \eta) \cdot \pi(\theta) p(\mathbf{A}_o, \mathbf{A}_m | \mathbf{X}, \theta)$$
- In this scenario, further restrictions are required:
 - Missingness mechanism of the network sampling design (*ignorability*).
 - Interference between recruited and non-recruited units. RCTs are possible with further restrictions on exposure mappings. Observational studies are tricky.
 - Projective network models [2].

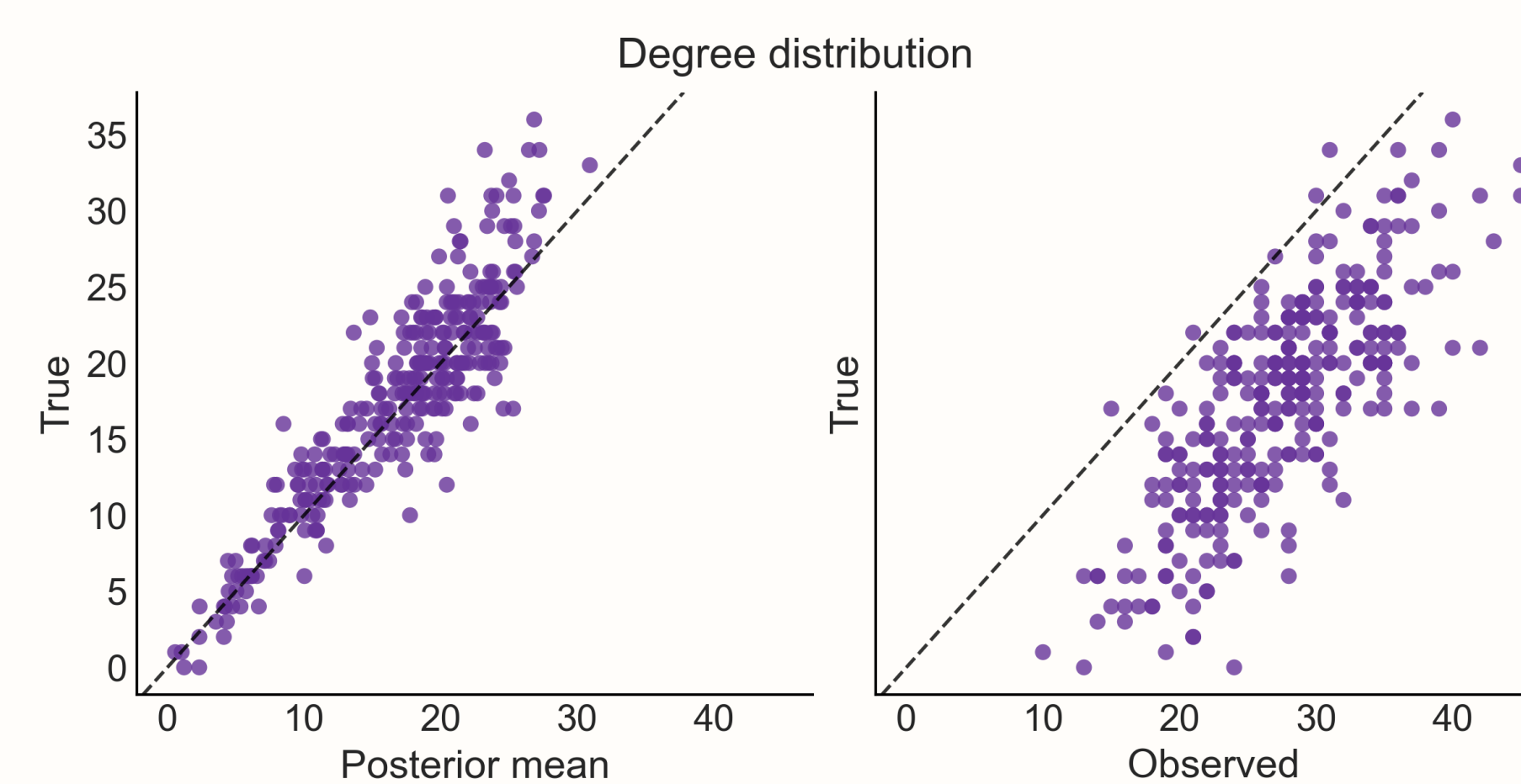


Figure 2. Reconstructing \mathbf{A}^* from \mathbf{A} . Degrees $d_i = \sum_{j \neq i} A_{ij}$ in the observed network (right) and the posterior mean degrees (left) versus the true degrees.

Sampling from the posterior

- The full posterior is a mixed space of continuous (η, θ, γ) and discrete (\mathbf{A}^*) latent variables. The discrete space has $\mathcal{O}(2^{n^2})$ terms.
- MCMC methods such as MH or modified HMC/NUTS [6, 7] do not scale well. Marginalizing over \mathbf{A}^* is problematic since Y_i depends on \mathbf{A}_i^* .

Bayesian modularization

The posterior can be written as a composition of modules

$$\pi(\eta, \theta, \gamma, \mathbf{A}^* | \mathbf{O}) \propto \underbrace{\pi(\eta | \mathbf{Y}, \mathbf{Z}, \mathbf{X}, \mathbf{A}^*)}_{\text{Outcome module}} \underbrace{\pi(\theta, \gamma, \mathbf{A}^* | \mathbf{X}, \mathbf{A})}_{\text{Network module}} \underbrace{p(\mathbf{Y} | \mathbf{Z}, \mathbf{X}, \mathbf{A}^*)}_{\text{Feedback term}}$$

Consequently, sampling from the ‘cut’ posterior [1, 3]

$$\pi_{\text{cut}}(\eta, \theta, \gamma, \mathbf{A}^* | \mathbf{O}) \propto \pi(\eta | \mathbf{Y}, \mathbf{Z}, \mathbf{X}, \mathbf{A}^*) \pi(\theta, \gamma, \mathbf{A}^* | \mathbf{X}, \mathbf{A}),$$

is attractive since

$$\pi(\theta, \gamma, \mathbf{A}^* | \mathbf{X}, \mathbf{A}) = \pi(\mathbf{A}^* | \mathbf{X}, \mathbf{A}, \theta, \gamma) \sum_{\mathbf{A}^*} \pi(\theta, \gamma, \mathbf{A}^* | \mathbf{X}, \mathbf{A}),$$

can be simplified tremendously. Sampling from the ‘cut’ posterior by [1, 3]:

- Generate (θ_m, γ_m) samples. Then, sample multiple \mathbf{A}_m^* . For each \mathbf{A}_m^* , sample η from the outcome module. Networks can be sampled via either
 - “Three-stage”**. For each (θ_m, γ_m) sample one network from $\mathbf{A}_m^* \sim \pi(\mathbf{A}^* | \mathbf{X}, \mathbf{A}, \theta_m, \gamma_m)$.
 - “Two-stage”**. Compute $\mathbb{E}[\theta, \gamma | \cdot]$ and sample $\mathbf{A}_m^* \sim \pi(\mathbf{A}^* | \mathbf{X}, \mathbf{A}, \mathbb{E}[\theta, \gamma | \cdot])$.
- “Plug-in”**. Sample multiple \mathbf{A}^* , estimate sufficient statistics of $p(\mathbf{Y} | \cdot)$ (e.g., exposure values), and sample η from the outcome module.

Numerical illustration

- Outcome model $Y_i = \eta_0 + \eta_1 Z_i + \eta_2 \sum_{j \neq i} Z_j A_{ij}^* + \eta_3 X_i + \varepsilon_i$
- Network generation $\text{logit}(\Pr(A_{ij}^* = 1)) = \theta_0 + \theta_1 |X_i - X_j|$.
- Observed network from a random noise measurement error model.
 - Implemented in the probabilistic programming language NumPyro. Accelerated cut-posterior sampling with JAX JIT compilation.
 - MCMC sampling via NUTS.

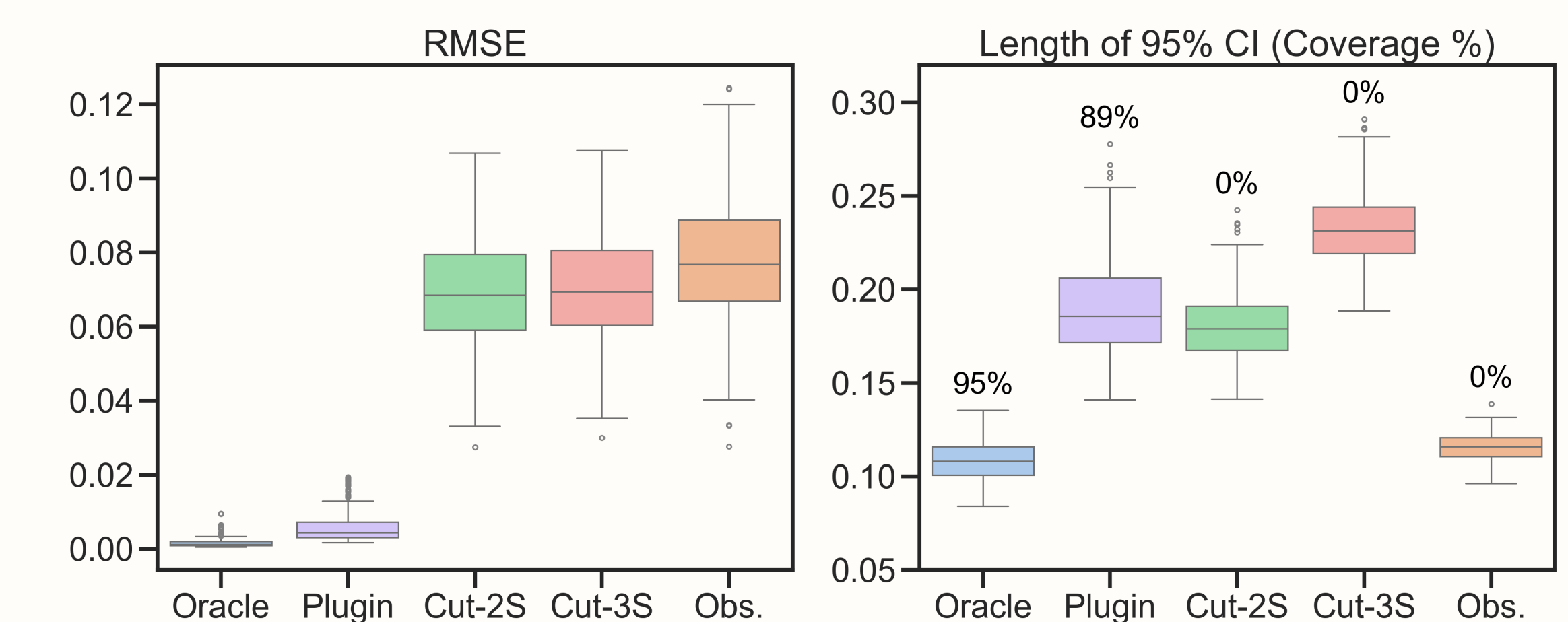


Figure 3. Distribution of η_2 RMSE and 95% credible intervals (coverage) for $n = 300$ and 300 replications. ‘Oracle’ and ‘Obs.’ display results using \mathbf{A}^*, \mathbf{A} , respectively. ‘Cut-2S’ is “Two-stage” sampling, and similarly for “Cut-3S”.

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