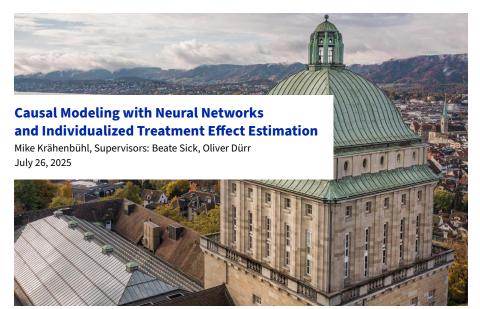
Master Program in Biostatistics www.biostat.uzh.ch Master Exam



Background

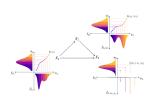
Supervisors:

- Beate Sick, UZH
- Oliver Dürr, HTWG Konstanz

Paper "Interpretable Neural Causal Models with TRAM-DAGs" (Sick and Dürr, 2025):

- Framework to model causal relationships
- Based on transformation models
- Rely on (deep) neural networks
- Compromise between interpretability and flexibility

They showed on synthetic data, that TRAM-DAGs can be fitted on observational data and tackle causal queries on all three levels of Pearl's causal hierarchy.



Research Questions

In this presentation:

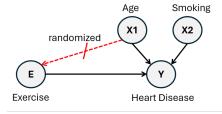
- 1. TRAM-DAGs
 - How do they work?
- 2. Individualized Treatment Effect (ITE) estimation
 - Does it work on real data (International Stroke Trial)?
 - When and why does ITE estimation fail (simulation)?
 - How to estimate ITEs with TRAM-DAGs in a complicated graph (simulation)?



TRAM-DAGs: Motivation

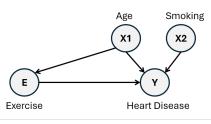
Randomized Controlled Trial:

- Gold standard for estimating causal effect
- Solves problem of confounding



Observational Data:

- Real world, potential confounding
- We assume no unobserved confounding



TRAM-DAGs: Motivation

Pearl's causal hierarchy (Pearl, 2009)

Observational: $P(Y = 1 \mid E = 1)$

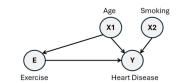
"Probability of heart disease given that the person exercises"

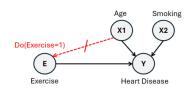
Interventional: $P(Y = 1 \mid do(E = 1))$

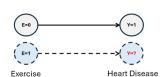
"Probability of heart disease if we made people start exercising"

Counterfactual: $P(Y_{(E=1)} = 1 \mid E = 0, Y = 1)$

"Would someone who does not exercise and has heart disease still have it if they had exercised?"



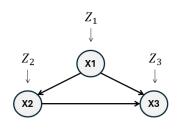




Structural Causal Model: Describes the causal mechanism and probabilistic uncertainty (Pearl, 2009)

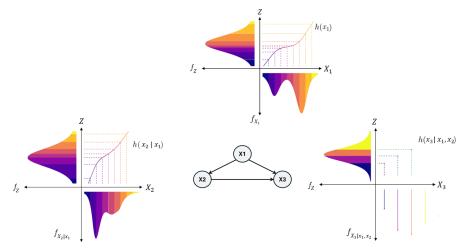
- $-X_i$ = observed variable
- $-Z_i$ = noise distribution
- f_i = deterministic function: $X_i = f_i(Z_i, pa_i)$

 \rightarrow We want a model that estimates $X_i = f_i(Z_i, pa_i)$ in a flexible and interpretable way!



$$Z \sim F_{Z_1}$$
, $Z_2 \sim F_{Z_2}$, $Z_3 \sim F_{Z_3}$
 $X_1 = f_1(Z_1)$
 $X_2 = f_2(Z_2, X_1)$
 $X_3 = f_3(Z_3, X_1, X_2)$

Proposed framework: TRAM-DAGs (Sick and Dürr, 2025)



Transformation Models: Flexible distributional regression method (Hothorn et al., 2014)

Continuous $Y \in \mathbb{R}$:

$$F_{Y|\mathbf{X}=\mathbf{x}}(y) = F_{Z}(h(y) + \mathbf{x}^{\top}\boldsymbol{\beta})$$

Discrete $Y \in \{y_1, y_2, \dots, y_K\}$:

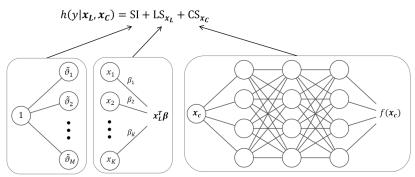
$$P(Y \le y_k \mid \mathbf{X} = \mathbf{x}) = F_Z(\vartheta_k + \mathbf{x}^{\top}\boldsymbol{\beta}), \quad k = 1, 2, \dots, K - 1$$

- $-F_Z$: CDF of the latent distribution (e.g. standard logistic)
- h: Transformation function, monotonically increasing
- x: Predictors

Extended to Deep TRAMs (Sick et al., 2021)

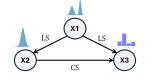
- Customizable transformation model using neural networks (NNs)
- Minimizing negative log-likelihood (NLL) via NN optimization

Effects of predictors: LS (Linear Shift), CS (Complex Shift), CI (Complex Intercept)



Setup:

- We have:
 - Observational data (simulated)
 - Predefined DAG
- We want:
 - Estimate $Z_i = h_i(X_i \mid pa(X_i))$ of each variable i
 - Sample from conditional distributions for causal queries with structural equations $X_i = h_i^{-1}(Z_i \mid pa(X_i))$



$$X_1 \sim F_Z(h(x_1))$$

 $X_2 \sim F_Z(h(x_2) + LS_{x_1})$
 $X_3 \sim F_Z(h(x_3) + LS_{x_1} + CS_{x_2})$

Data-generating process (DGP):

 X_1 : Continuous, bimodal. *Source node* (independent).



 X_2 : Continuous. Depends on X_1 (linear):

$$\frac{\beta_{12} = 2}{h(X_2 \mid X_1) = h_I(X_2) + \beta_{12}X_1}$$



 X_3 : Ordinal. Depends on X_1 (linear) and X_2 (complex):

$$\beta_{13} = 0.2$$
, $f(X_2) = 0.5 \cdot \exp(X_2)$, $\vartheta_k \in \{-2, 0.42, 1.02\}$

$$h(X_{3,k} \mid X_1, X_2) = \vartheta_k + \beta_{13}X_1 + f(X_2)$$



Construct Model: Modular Neural Network

Inputs: Observations + assumed structure

Outputs:

- Simple Intercepts (SI): ϑ
- Linear Shifts (LS): $\beta_{12}X_1$, $\beta_{13}X_2$
- Complex Shift (CS): $f(X_2)$

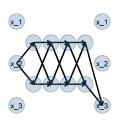
Assemble transformation functions:

$$h(X_{i} \mid pa(X_{i})) = SI + LS + CS$$

$$h(X_{1}) = h_{I}(X_{1})$$

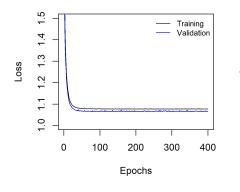
$$h(X_{2} \mid X_{1}) = h_{I}(X_{2}) + \beta_{12}X_{1}$$

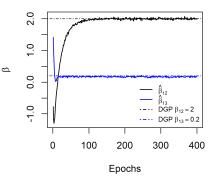
$$h(X_{3} \mid X_{1}, X_{2}) = \vartheta_{k} + \beta_{13}X_{1} + f(X_{2})$$



 CS_{X_2} on X_3

Model fitting: 20,000 training samples, 400 epochs





Sampling from the Fitted TRAM-DAG

Nodes $X_i, i \in \{1, 2, 3\}$:

— Sample latent value:

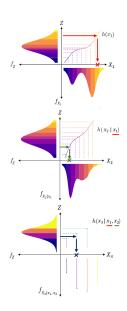
$$z_i \sim F_{Z_i}$$
 (e.g., rlogis() in R)

- Determine x_i such that:
 - **If** X_i **is continuous:** Solve for x_i using numerical root-finding:

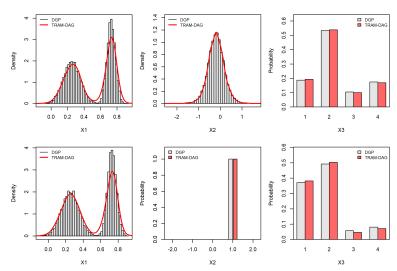
$$h(x_i \mid pa(x_i)) - z_i = 0$$

If X_i is ordinal: find the smallest category
 x_i such that

$$x_i = \max(\{0\} \cup \{x : z_i > h(x \mid pa(x_i))\}) + 1$$

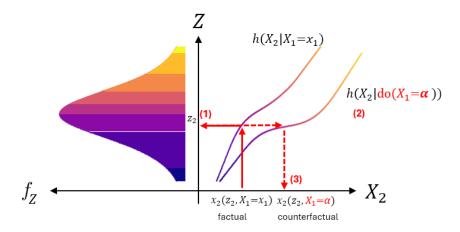


Sampled observational and interventional distributions:



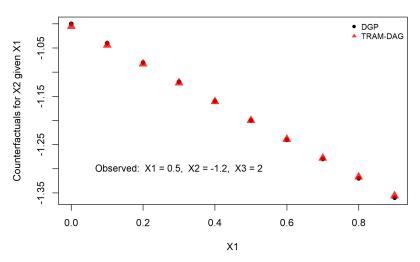
Experiment 1: TRAM-DAGs (simulation)

How to determine a counterfactual value for X_2 , given some observation?



Experiment 1: TRAM-DAGs (simulation)

Counterfactuals: Counterfactual value of X_2 under varying X_1



Experiment 1: TRAM-DAGs (simulation)

Discussion: With TRAM-DAGs we can

- estimate the functional form of the edges in the DAG
- customize flexibility and interpretability (SI/CI, LS, CS)
- sample from the fitted model (observational/interventional)
- estimate counterfactuals

Individualized Treatment Effects

(ITEs)

Individualized Treatment Effect (ITE): Motivation

Why ITE?

- RCTs estimate the Average Treatment Effect (ATE)
- Individuals may respond differently based on covariates
- Important for personalized medicine, targeted marketing, etc.
- Heterogeneity mostly driven by treatment-covariate interactions

Definition: Difference in potential outcomes (Rubin, 2005)

$$Y_i(1) - Y_i(0)$$

where $Y_i(1)$: outcome if treated, $Y_i(0)$: if not treated

Fundamental problem: We never observe both $Y_i(1)$ and $Y_i(0)$ for the same individual (Holland, 1986)

From Unobservable to Estimable ITE

Goal: Estimate the *individualized treatment effect (ITE)* from observed data (Hoogland et al., 2021).

$$\begin{aligned} \mathsf{ITE}(\mathbf{x}_i) &= \mathbb{E}[Y_i(1) - Y_i(0) \mid \mathbf{X} = \mathbf{x}_i] \\ &= \mathbb{E}[Y_i(1) \mid T = 1, \mathbf{X} = \mathbf{x}_i] - \mathbb{E}[Y_i(0) \mid T = 0, \mathbf{X} = \mathbf{x}_i] \\ & \textit{(by ignorability/exchangeability: no unmeasured confounding)} \\ &= \mathbb{E}[Y_i \mid T = 1, \mathbf{X} = \mathbf{x}_i] - \mathbb{E}[Y_i \mid T = 0, \mathbf{X} = \mathbf{x}_i] \\ & \textit{(by consistency: observed = potential outcome, e.g. correct label)} \end{aligned}$$

Further assumptions:

- Positivity: every individual could receive either treatment (e.g. no deterministic assignment)
- No interference: one person's treatment doesn't affect another's outcome

Individualized Treatment Effect (ITE): Models

How did we estimate the potential outcomes $\mathbb{E}[Y_i \mid T = t, \mathbf{X} = \mathbf{x}_i]$?

— T-learner:

- 1. Fit two separate models on treated and control groups
- 2. Predict $\mathbb{E}[Y_i \mid \mathbf{X} = \mathbf{x}_i]$ from each model
- Logistic regression / Random forest (with hyperparameter tuning)

— S-learner:

- 1. Fit one model on all data with treatment as a feature
- 2. Predict $\mathbb{E}[Y_i \mid do(T=t), \mathbf{X} = \mathbf{x}_i]$ by setting T=0 and T=1
- TRAM-DAGs (SCM, flexible, interactions, generative)

Experiment 2: ITE on International Stroke Trial (IST)

Background/Motivation: Chen et al. (2025) showed that results of models used for ITE estimation did not generalize to the test set.

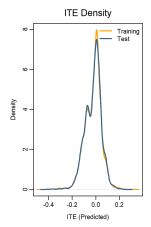
International Stroke Trial (IST):

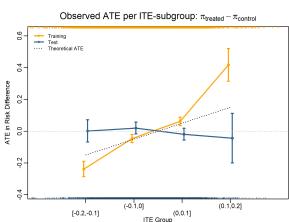
- Large RCT on stroke patients (19,435 patients, 21 baseline covariates)
- Evaluated the effects of aspirin on stroke patients
- Binary treatment and outcome

Research question: Do we reach similar conclusion as Chen et al. (2025) when estimating ITEs with T-learners (logistic regression, tuned random forest) and S-learner (TRAM-DAGs) on IST dataset.

Experiment 2: ITE on International Stroke Trial (IST)

Results: with T-learner tuned random forest (comets package (Kook, 2024)):





Experiment 2: ITE on International Stroke Trial (IST)

Discussion:

- We obtained similar results as Chen et al. (2025)
- Some models suggest a range of ITEs, but these ITEs do not generalize to the test set (no effect)
- We do not know why, since ground truth is unknown

Experiment 3: ITE model robustness in RCTs (simulation)

Motivation: ITE estimation failed on the real-world RCT of the International Stroke Trial (IST). We want to know why.

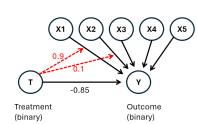
Research question: What factors contribute to the failure of ITE estimation in causal models?

Setup:

- Simulate different RCT scenarios to understand when ITE estimation fails
- Apply simple model (logistic regression; matching DGP) and non-parametric model (tuned random forest)

Setup:

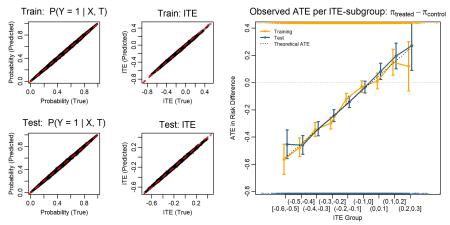
- -n = 20,000
- $T \sim \text{Bernoulli}(0.5)$
- $\mathbf{X} = (X_1, \dots, X_5)^{\top} \sim \mathcal{N}(\mathbf{0}, \Sigma)$
- $\mathbf{X}_{\mathbf{TX}} = (X_1, X_2)^{\top}$ interaction



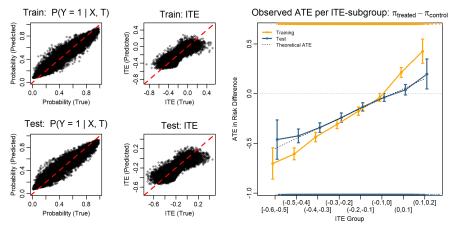
Outcome model:

$$\mathbb{P}(Y = 1 \mid \mathbf{X}, T) = \mathsf{logit}^{-1} \left(\beta_0 + \beta_T T + \boldsymbol{\beta}_X^\top \mathbf{X} + \underline{T} \cdot \boldsymbol{\beta}_{TX}^\top \mathbf{X}_{\mathsf{TX}}\right)$$

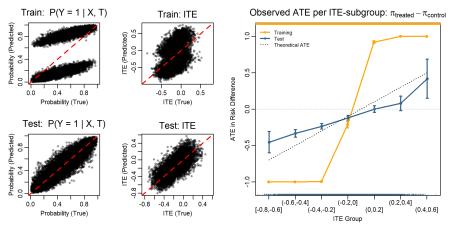
Results with T-learner logistic regression (glm):



Results with T-learner tuned random forest (comets package):



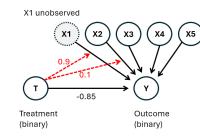
Results with (untuned) T-learner random forest (randomForest package):



Simulation Case 2: Unobserved Interaction

Setup:

- n = 20,000
- − T ~ Bernoulli(0.5)
- $\mathbf{X} = (X_1, \dots, X_5)^{\top} \sim \mathcal{N}(\mathbf{0}, \Sigma)$
- $\mathbf{X}_{\mathbf{TX}} = (X_1, X_2)^{\top}$ interaction



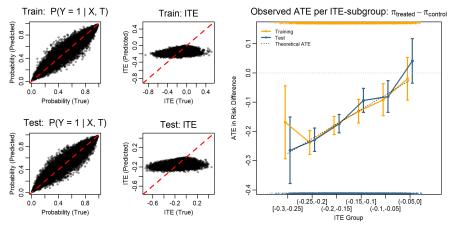
Outcome model:

$$\mathbb{P}(\textit{Y} = 1 \mid \textbf{X}, \textit{T}) = \mathsf{logit}^{-1} \left(\beta_{0} + \beta_{\textit{T}}\textit{T} + \boldsymbol{\beta}_{\textit{X}}^{\top}\textbf{X} + \boldsymbol{T} \cdot \boldsymbol{\beta}_{\textit{TX}}^{\top}\textbf{X}_{\textbf{TX}}\right)$$

Note: Same DGP, but X_1 is not observed!

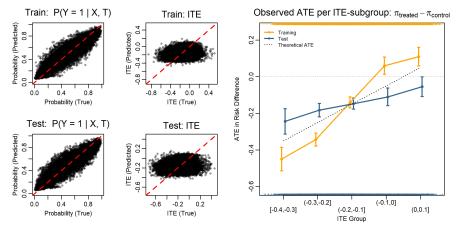
Simulation Case 2: Unobserved Interaction

Results with T-learner logistic regression (glm):



Simulation Case 2: Unobserved Interaction

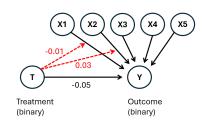
Results with T-learner tuned random forest (comets package):



Simulation Case 3: Fully Observed, Small Effects

Setup:

- n = 20,000
- $T \sim Bernoulli(0.5)$
- $\mathbf{X} = (X_1, \dots, X_5)^{\top} \sim \mathcal{N}(\mathbf{0}, \Sigma)$
- $\mathbf{X}_{\mathbf{TX}} = (X_1, X_2)^{\top}$ interaction



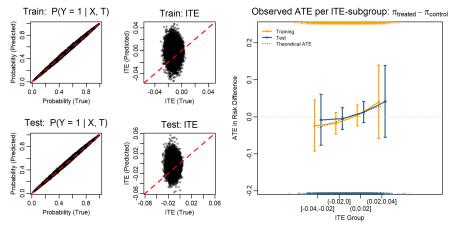
Outcome model:

$$\mathbb{P}(Y = 1 \mid \mathbf{X}, T) = \mathsf{logit}^{-1} \left(\beta_0 + \beta_T T + \boldsymbol{\beta}_X^\top \mathbf{X} + \underline{T} \cdot \boldsymbol{\beta}_{TX}^\top \mathbf{X}_{\mathsf{TX}} \right)$$

Note: Same DGP, but weak treatment effects!

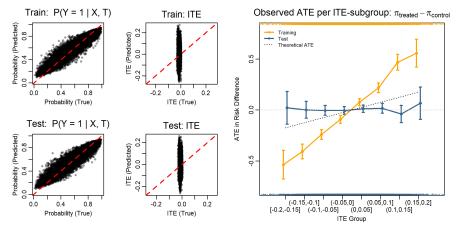
Simulation Case 3: Fully Observed, Small Effects

Results with T-learner logistic regression (glm):



Simulation Case 3: Fully Observed, Small Effects

Results with T-learner Random Forest (comets package):



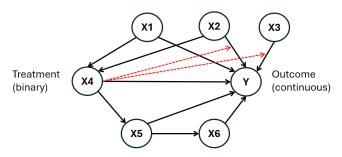
Experiment 3: ITE model robustness in RCTs (simulation)

Key Insights:

- Calibration and tuning of models are crucial for reliable ITE estimation
- Ignorability alone may not guarantee unbiased ITEs if important effect modifiers are unobserved
- Low true heterogeneity may be mistaken for model failure

These factors may explain the limited ITE performance in the IST dataset.

TRAM-DAGs: Example for ITE estimation

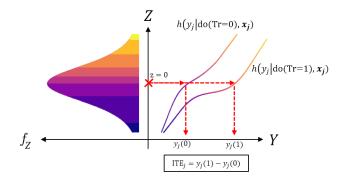


DGP:

- $X5 = h_5^{-1} (\epsilon 0.8 X4)$ → (depends on treatment)
- $X6 = h_6^{-1}(\epsilon + 0.5X5) \rightarrow \text{(depends on treatment through X5)}$
- $Y = h_7^{-1} (\epsilon \beta_1 X 1 \beta_2 X 2 \beta_3 X 3 \beta_4 X 4 \beta_5 X 5 \beta_6 X 6 Tr \cdot (\beta_{2Tr} X 2 + \beta_{3Tr} X 3))$

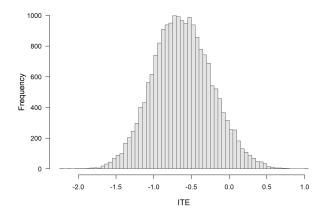
TRAM-DAGs: Example for ITE estimation

$$\mathsf{ITE} = \mathsf{median}(Y \mid \mathsf{do}(T=1), X) - \mathsf{median}(Y \mid \mathsf{do}(T=0), X)$$



TRAM-DAGs: Example for ITE estimation

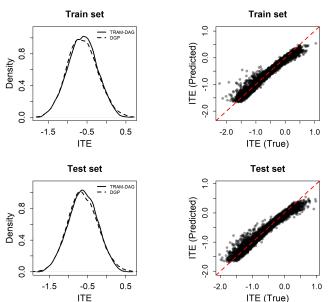
$$ITE = median(Y \mid do(T = 1), X) - median(Y \mid do(T = 0), X)$$



TRAM-DAGs: Estimate Potential Outcomes

- 1. Estimate each $h_i(X_i \mid pa(X_i))$ fully flexible (deep-NN / complex intercept)
- 2. Take the train set or a test set
- 3. $Z_i = h(X_i \mid pa(X_i))$ gives us the (observed) latent variable for each X_i
- 4. Determine counterfactuals for X5 and X6 with the (observed) latent variables *Z*_i
- 5. Determine medians of potential outcomes Y(1) and Y(0)
- 6. ITE = median($Y(1) \mid X_{tx}$) median($Y(0) \mid X_{ct}$)

TRAM-DAGs: Example for ITE estimation (Results)



Key Findings

Findings: TRAM-DAGs

- Customizable; accurately recovers causal relationships in known DAG; allows sampling of L1-L3
- Can model interactions between variables

Findings: Individualized treatment effects (ITE)

- Calibration is important for ITE prediction
- Missing effect modifiers (or weak heterogeneity) are problematic
- TRAM-DAGs yield unbiased ITEs when DAG is correct and heterogeneity exists

Outlook

Limitations

- Simulations may not reflect real-world complexity
- TRAM-DAGs are computationally expensive (long training time)
- TRAM-DAGs require correct model specification for interpretability
- ITE estimation used median QTE for continuous outcomes

Recommendations

- Apply TRAM-DAGs to real-world datasets, including semi-structured data
- Investigate ITE estimation under unobserved effect modifiers

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