

Doubly-Robust Functional Average Treatment Effect Estimation

Reading Group Presentation

Lorenzo Testa, Tobia Boschi, Francesca Chiaromonte, Edward H. Kennedy, Matthew Reimherr

March 28, 2025

Presented by Yuankang Zhao

- **Motivation:** Understanding causal effects in observational studies with functional data.
- Many traditional methods target scalar outcomes; however, many applications (e.g., longitudinal medical data) require handling functional outcomes.
- The paper addresses the estimation of the Functional Average Treatment Effect (FATE) over a continuous domain.

- **Challenges:**

- Functional data is infinite-dimensional and exhibits complex dependence across the domain.
- Standard causal inference methods (e.g., AIPW) are designed for scalar outcomes.

- **Existing Methods:**

- Function-on-scalar regression models are often non-robust.
- Some approaches (e.g., Liu et al. (2024)) require strong parametric assumptions.

Introduction: Need for a New Approach

- **DR-FoS:** A novel estimator that extends double robustness to the functional data setting.
- **Key Idea:** Combine outcome regression and propensity score weighting to achieve consistency if either model is correctly specified.
- **Additional Feature:** Uses cross-fitting and provides simultaneous confidence bands over the continuous domain.

Definition of FATE

- The target parameter is the **Functional Average Treatment Effect (FATE)**:

$$\beta(t) = E \left[Y^{(1)}(t) - Y^{(0)}(t) \right]$$

- Here, $Y^{(a)}(t)$ denotes the potential outcome function at time t under treatment $a \in \{0, 1\}$.
- The estimation target is a function over a continuous domain (e.g., time interval $T = [0, 1]$).
- **Identifiability Assumption**
- **Consistency** The potential outcome of a treatment is the same regardless of the mechanism by which the treatment is administered; that is, $Y^{(a)}(t)$ if $A = a$ for any $t \in T$.
- **No unmeasured confounding.** $Y^{(a)}(t) \perp\!\!\!\perp A \mid X$ for any $t \in T$.
- **Positivity.** $0 < \mathbb{P}[A = 1 \mid X] < 1$ for any $t \in T$.

Estimation: Model Setup

- Two main components are modeled:

- **Propensity Score:**

$$\pi^{(a)}(X) = P(A = a \mid X)$$

- **Outcome Regression:**

$$\mu^{(a)}(X, t) = E[Y^{(a)}(t) \mid X, A = a]$$

- Both models can be estimated using flexible methods (e.g., logistic regression, nonparametric regression).
- **Double Robustness:** The estimator remains consistent if either the propensity model or the outcome model is correctly specified.

- **Corrected Regression Function:**

$$\gamma^{(a)}(D, t) = \begin{cases} \mu^{(a)}(X, t) + \frac{Y(t) - \mu^{(a)}(X, t)}{\pi^{(a)}(X)} & \text{if } A = a, \\ \mu^{(a)}(X, t) & \text{if } A \neq a. \end{cases}$$

where $D = (A, X, Y)$

- Then FATE could be rewritten as:

$$\beta(t) = E \left[\gamma^{(1)}(D, t) - \gamma^{(0)}(D, t) \right]$$

Estimation: The DR-FoS Estimator

Proof:

$$\begin{aligned} E \left[\gamma^{(1)}(D, t) \right] &= E \left[\mu^{(1)}(X, t) + \frac{A(Y(t) - \mu^{(1)}(X, t))}{\pi^{(1)}(X)} \right] \\ &= E \left[\mu^{(1)}(X, t) \right] + E \left[\frac{A(Y(t) - \mu^{(1)}(X, t))}{\pi^{(1)}(X)} \right] \\ &= E \left[E[Y(t)|X, A = 1] \right] + E \left[E \left[\frac{A(Y(t) - \mu^{(1)}(X, t))}{\pi^{(1)}(X)} \middle| X \right] \right] \\ &= E \left[E[Y(t)|X, A = 1] \right] \\ &= E[Y^{(1)}(t)] \end{aligned}$$

Estimation: The DR-FoS Estimator

The DR-FoS estimator is then given by:

$$\begin{aligned}\hat{\beta}_{\text{DR-FoS}}(t) &= P_n \left[\hat{\gamma}^{(1)}(D, t) - \hat{\gamma}^{(0)}(D, t) \right] \\&= \frac{1}{n} \sum_{i=1}^n \left[\hat{\gamma}^{(1)}(D_i, t) - \hat{\gamma}^{(0)}(D_i, t) \right] \\&= \frac{1}{n} \sum_{i=1}^n \left[\left(\hat{\mu}^{(1)}(X_i, t) + \frac{A_i(Y_i(t) - \hat{\mu}^{(1)}(X_i, t))}{\hat{\pi}^{(1)}(X_i)} \right) \right. \\&\quad \left. - \left(\hat{\mu}^{(0)}(X_i, t) + \frac{(1 - A_i)(Y_i(t) - \hat{\mu}^{(0)}(X_i, t))}{1 - \hat{\pi}^{(1)}(X_i)} \right) \right]\end{aligned}$$

Cross-Fitting Procedure for DR-FoS Estimation

Step 1: Data Partitioning

Randomly partition dataset into J folds:

$$\{D_1, D_2, \dots, D_n\} D_i = (A_i, X_i, Y_i(t))$$

Step 2: Training-Prediction Cycle (for each fold j)

For each fold $j = 1, \dots, J$:

- ① Train models using all data except fold j :
 - Estimate propensity score: $\hat{\pi}^{(a), [-j]}(X)$
 - Estimate outcome regression: $\hat{\mu}^{(a), [-j]}(X, t)$
- ② Compute DR-FoS estimator on fold j :

$$\hat{\beta}_{\text{DR-FoS}}^{[j]}(t) = \frac{1}{n_j} \sum_{i \in \text{Fold } j} \left[\hat{\gamma}^{(1), [-j]}(D_i, t) - \hat{\gamma}^{(0), [-j]}(D_i, t) \right]$$

Step 3: Averaging Over Folds

$$\hat{\beta}_{\text{DR-FoS}}(t) = \frac{1}{J} \sum_{j=1}^J \hat{\beta}_{\text{DR-FoS}}^{[j]}(t)$$

Inference: Asymptotic Properties

- Under regularity conditions, the estimator converges in distribution:

$$\sqrt{n}(\hat{\beta}_{\text{DR-FoS}}(t) - \beta(t)) \Rightarrow \mathcal{GP}(0, \Sigma(s, t))$$

- \mathcal{GP} denotes a Gaussian process, ensuring pointwise asymptotic normality.
- The covariance function is given by:

$$\Sigma(s, t) = E[\phi(D; s)\phi(D; t)],$$

where $\phi(D; t) = \gamma^{(1)}(D, t) - \gamma^{(0)}(D, t) - \beta(t)$.

Inference: Constructing Simultaneous Confidence Bands

- Two approaches are discussed:
 - ① **Critical Value Function Approach:** Uses an estimated critical value $u_{\alpha/2}^*$ to form bands:

$$C_{\alpha}(t) = \hat{\beta}_{\text{DR-FoS}}(t) \pm \frac{u_{\alpha/2}^* \hat{\sigma}(t)}{\sqrt{n}}.$$

- ② **Parametric Bootstrap:** Repeatedly sample from the estimated Gaussian process to obtain empirical quantiles, forming the band:

$$C_{\alpha}(t) = [\hat{\beta}_{\alpha/2}(t), \hat{\beta}_{1-\alpha/2}(t)].$$

- These bands provide simultaneous coverage over the entire function domain.

- **Data Generating Process:**

- True outcome model:

$$\mu^{(a)}(t) = a\beta(t) + \rho(t),$$

where $\beta(t)$ and $\rho(t)$ are drawn from a Gaussian process with Matern covariance.

- Observations:

$$Y_i^{(a)}(t) = \mu^{(a)}(t) + \epsilon_i(t),$$

and

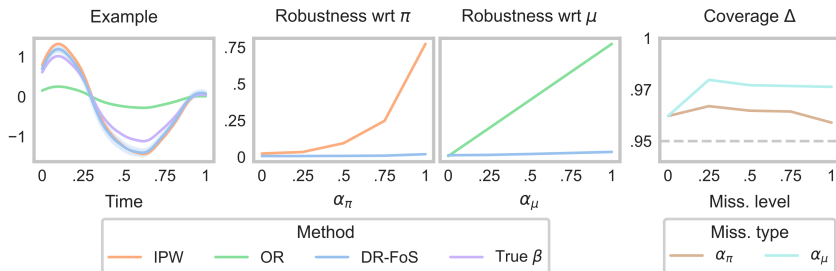
$$Y_i(t) = A_i Y_i^{(1)}(t) + (1 - A_i) Y_i^{(0)}(t).$$

- **Controlling Model Misspecification:**

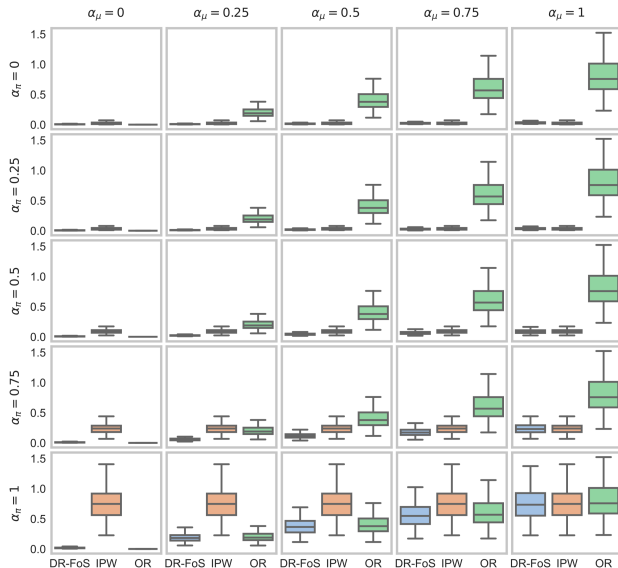
- α_π controls the misspecification of the propensity score model.
- α_μ controls the misspecification of the outcome regression model.

Simulation Study: Results and Implications

- Compared methods: DR-FoS, Outcome Regression (OR), and Inverse Probability Weighting (IPW) via MSE and Coverage probability of the simultaneous confidence bands.
- Key findings:
 - When one model is misspecified (either $\alpha_\pi > 0$ or $\alpha_\mu > 0$), DR-FoS remains robust.
 - When both models are misspecified, DR-FoS outperforms the other methods.



Simulation Study: Results and Implications



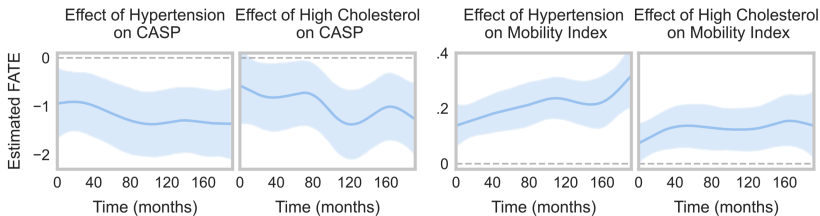
SHARE Application: Dataset and Variables

- **Dataset:** Survey of Health, Ageing and Retirement in Europe (SHARE).
- **Participants:** 1518 subjects with at least 7 out of 8 survey waves.
- **Key Variables:**
 - **Treatments:** High blood pressure and high cholesterol.
 - **Outcomes:** Functional variables such as the Mobility Index and CASP (quality of life measure).
 - **Covariates:** Age, education, numeracy score, gender, smoking status, childhood vaccinations.
- **Preprocessing:**
 - Time-varying outcomes are smoothed using cubic B-splines.
 - Functional representations are constructed to align data across survey waves.
- **Estimation Setup:**
 - Logistic regression for the propensity score.
 - Function-on-scalar least squares for the outcome regression.
 - 5-fold cross-fitting is employed.
 - Parametric bootstrap is used to build simultaneous confidence bands.

SHARE Application: Main Findings and Implications

Findings:

- Both high blood pressure and high cholesterol have a significant negative impact on quality of life (CASP) over time.
- Effects on the Mobility Index indicate a deterioration in mobility (a higher index implies worse mobility).
- The adverse effects become more pronounced with increasing age.



Implications:

- The DR-FoS method demonstrates practical utility in real-world, complex longitudinal data.
- These insights can inform healthcare policy and personalized treatment planning.

Summary

- Proposed a **Doubly-Robust Functional Average Treatment Effect** (DR-FoS) estimator, and established rigorous asymptotic theory to make inference.
- Demonstrated excellent performance in simulation studies, and applied DR-FoS to real longitudinal data (SHARE study).

Concerns and Future Directions:

- **Parametric Assumptions and Robustness:**
 - Current simulations assume Gaussian processes. How robust is the DR-FoS estimator to deviations from such assumptions (e.g., non-Gaussian errors)?
 - Need to further assess robustness in diverse practical data-generating scenarios.
- **Limited Exploration of High-dimensional Covariates:**
 - Analyses involved relatively low-dimensional covariates. High-dimensional settings could challenge estimation accuracy.
 - Future work should explore regularization or dimensionality reduction approaches suitable for high-dimensional covariates.