#### **Covariate Balancing Propensity Score**

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#### Joint work with Marc Ratkovic and Christian Fong

- Main Paper: Imai, K. and Ratkovic, M. (2013). "Covariate Balancing Propensity Score" *Journal of the Royal Statistical Society, Series B (Methodological)*, Forthcoming.
- Software: Ratkovic, M., K. Imai, C. Fong. (2013). CBPS: R Package for Covariate Balancing Propensity Score available for download at the CRAN

These and other related materials available at http://imai.princeton.edu

- Causal inference is a central goal of scientific research
- Randomized experiments are not always possible
   ⇒ Causal inference in observational studies
- Experiments often lack external validity
   Need to generalize experimental results
- Importance of statistical methods to adjust for confounding factors

## Overview of the Talk

#### Review: Propensity score

- propensity score is a covariate balancing score
- matching and weighting methods
- Problem: Propensity score tautology
  - sensitivity to model misspecification
  - adhoc specification searches
- Solution: Covariate balancing propensity score (CBPS)
  - Estimate propensity score so that covariate balance is optimized
- Evidence: Reanalysis of two prominent critiques
  - Improved performance of propensity score weighting and matching
- Software: R package CBPS
- Extension: General Treatment Regimes

# **Propensity Score**

#### • Setup:

- $T_i \in \{0, 1\}$ : binary treatment
- X<sub>i</sub>: pre-treatment covariates
- $(Y_i(1), Y_i(0))$ : potential outcomes
- $Y_i = Y_i(T_i)$ : observed outcomes
- Definition: conditional probability of treatment assignment

$$\pi(X_i) = \Pr(T_i = 1 \mid X_i)$$

• Balancing property (without assumption):

$$T_i \perp\!\!\!\perp X_i \mid \pi(X_i)$$

• Assumptions:



$$0 < \pi(X_i) < 1$$

Our Control Control

$$\{Y_i(1), Y_i(0)\} \perp T_i \mid X_i$$

• Propensity score as a dimension reduction tool:

 $\{Y_i(1), Y_i(0)\} \perp T_i \mid \pi(X_i)$ 

# Matching and Weighting via Propensity Score

- Propensity score reduces the dimension of covariates
- But, propensity score must be estimated (more on this later)
- Once estimated, simple nonparametric adjustments are possible
- Matching
- Subclassification
- Weighting (Horvitz-Thompson estimator):

$$\frac{1}{n}\sum_{i=1}^{n}\left\{\frac{T_{i}Y_{i}}{\hat{\pi}(X_{i})}-\frac{(1-T_{i})Y_{i}}{1-\hat{\pi}(X_{i})}\right\}$$

often, weights are normalized

• Doubly-robust estimators (Robins et al.):

$$\frac{1}{n}\sum_{i=1}^{n}\left[\left\{\hat{\mu}(1,X_{i})+\frac{T_{i}(Y_{i}-\hat{\mu}(1,X_{i}))}{\hat{\pi}(X_{i})}\right\}-\left\{\hat{\mu}(0,X_{i})+\frac{(1-T_{i})(Y_{i}-\hat{\mu}(0,X_{i}))}{1-\hat{\pi}(X_{i})}\right\}\right]$$

• They have become standard tools for applied researchers

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- Propensity score is unknown
- Dimension reduction is purely theoretical: must model T<sub>i</sub> given X<sub>i</sub>
- Diagnostics: covariate balance checking
- In practice, adhoc specification searches are conducted
- Model misspecification is always possible
- Theory (Rubin *et al.*): ellipsoidal covariate distributions
   ⇒ equal percent bias reduction
- Skewed covariates are common in applied settings
- Propensity score methods can be sensitive to misspecification

# Kang and Schafer (2007, Statistical Science)

• Simulation study: the deteriorating performance of propensity score weighting methods when the model is misspecified

• Setup:

- 4 covariates X<sub>i</sub><sup>\*</sup>: all are *i.i.d.* standard normal
- Outcome model: linear model
- Propensity score model: logistic model with linear predictors
- Misspecification induced by measurement error:

• 
$$X_{i1} = \exp(X_{i1}^*/2)$$

• 
$$X_{i2} = X_{i2}^* / (1 + \exp(X_{1i}^*) + 10)$$

• 
$$X_{i3} = (X_{i1}^* X_{i3}^* / 25 + 0.6)^3$$

• 
$$X_{i4} = (X_{i1}^* + X_{i4}^* + 20)^2$$

- Weighting estimators to be evaluated:
  - Horvitz-Thompson
  - Inverse-probability weighting with normalized weights
  - Weighted least squares regression
  - Doubly-robust least squares regression

# Weighting Estimators Do Fine If the Model is Correct

		Bi	as	RM	RMSE		
Sample size	Estimator	GLM	True	GLM	True		
(1) Both mode	els correct						
	HT	0.33	1.19	12.61	23.93		
n - 200	IPW	-0.13	-0.13	3.98	5.03		
n = 200	WLS	-0.04	-0.04	2.58	2.58		
	DR	-0.04	-0.04	2.58	2.58		
	HT	0.01	-0.18	4.92	10.47		
n = 1000	IPW	0.01	-0.05	1.75	2.22		
<i>II</i> = 1000	WLS	0.01	0.01	1.14	1.14		
	DR	0.01	0.01	1.14	1.14		
(2) Propensity	y score mode	el correct					
- 000	HT	-0.05	-0.14	14.39	24.28		
	IPW	-0.13	-0.18	4.08	4.97		
11 = 200	WLS	0.04	0.04	2.51	2.51		
	DR	0.04	0.04	2.51	2.51		
<i>n</i> = 1000	HT	-0.02	0.29	4.85	10.62		
	IPW	0.02	-0.03	1.75	2.27		
	WLS	0.04	0.04	1.14	1.14		
	DR	0.04	0.04	1.14	1.14		

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# Weighting Estimators are Sensitive to Misspecification

		Bia	as	RMS	RMSE		
Sample size	Estimator	GLM	True	GLM	True		
(3) Outcome	model corre	ct					
	HT	24.25	-0.18	194.58	23.24		
n = 200	IPW	1.70	-0.26	9.75	4.93		
11 = 200	WLS	-2.29	0.41	4.03	3.31		
	DR	-0.08	-0.10	2.67	2.58		
	HT	41.14	-0.23	238.14	10.42		
n = 1000	IPW	4.93	-0.02	11.44	2.21		
<i>II</i> = 1000	WLS	-2.94	0.20	3.29	1.47		
	DR	0.02	0.01	1.89	1.13		
(4) Both mod	els incorrect	t					
	HT	30.32	-0.38	266.30	23.86		
n 000	IPW	1.93	-0.09	10.50	5.08		
n = 200	WLS	-2.13	0.55	3.87	3.29		
	DR	-7.46	0.37	50.30	3.74		
<i>n</i> = 1000	HT	101.47	0.01	2371.18	10.53		
	IPW	5.16	0.02	12.71	2.25		
	WLS	-2.95	0.37	3.30	1.47		
	DR	-48.66	0.08	1370.91	1.81		

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**Covariate Balancing Propensity Score** 

- LaLonde (1986; Amer. Econ. Rev.):
  - Randomized evaluation of a job training program
  - Replace experimental control group with another non-treated group
  - Current Population Survey and Panel Study for Income Dynamics
  - Many evaluation estimators didn't recover experimental benchmark
- Dehejia and Wahba (1999; J. of Amer. Stat. Assoc.):
  - Apply propensity score matching
  - Estimates are close to the experimental benchmark
- Smith and Todd (2005):
  - Dehejia & Wahba (DW)'s results are sensitive to model specification
  - They are also sensitive to the selection of comparison sample

# Propensity Score Matching Fails Miserably

- One of the most difficult scenarios identified by Smith and Todd:
  - LaLonde experimental sample rather than DW sample
  - Experimental estimate: \$886 (s.e. = 488)
  - PSID sample rather than CPS sample
- Evaluation bias:
  - Conditional probability of being in the experimental sample
  - Comparison between experimental control group and PSID sample
  - "True" estimate = 0
  - Logistic regression for propensity score
  - One-to-one nearest neighbor matching with replacement

Propensity score model	Estimates
Linear	-835
	(886)
Quadratic	-1620
	(1003)
Smith and Todd (2005)	-1910
	(1004)

## Covariate Balancing Propensity Score

- Idea: Estimate the propensity score such that covariate balance is optimized
- Covariate balancing condition:

$$\mathbb{E}\left\{\frac{T_i\widetilde{X}_i}{\pi_{\beta}(X_i)}-\frac{(1-T_i)\widetilde{X}_i}{1-\pi_{\beta}(X_i)}\right\} = 0$$

where  $\widetilde{X}_i = f(X_i)$  is any vector-valued function

• Score condition from maximum likelihood:

$$\mathbb{E}\left\{\frac{T_i\pi'_{\beta}(X_i)}{\pi_{\beta}(X_i)}-\frac{(1-T_i)\pi'_{\beta}(X_i)}{1-\pi_{\beta}(X_i)}\right\} = 0$$

# Weighting to Balance Covariates

• Balancing condition: 
$$\mathbb{E}\left\{\frac{T_iX_i}{\pi_{\beta}(X_i)} - \frac{(1-T_i)X_i}{1-\pi_{\beta}(X_i)}\right\} = 0$$



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Covariate Balancing Propensity Score

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# Generalized Method of Moments (GMM) Framework

- Just-identified CBPS: covariate balancing conditions alone
- Over-identified CBPS: combine them with score conditions
- GMM (Hansen 1982):

$$\hat{eta}_{\mathrm{GMM}} = \operatorname*{argmin}_{eta \in \Theta} ar{g}_eta(T,X)^ op \Sigma_eta(T,X)^{-1}ar{g}_eta(T,X)$$

where

$$\bar{g}_{\beta}(T,X) = \frac{1}{N} \sum_{i=1}^{N} \underbrace{\left(\begin{array}{c} \text{score condition} \\ \text{balancing condition} \end{array}\right)}_{g_{\beta}(T_i,X_i)}$$

 $\bullet\,$  "Continuous updating" GMM estimator for  $\Sigma\,$ 

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# Revisiting Kang and Schafer (2007)

		Bias				RN	ISE		
	Estimator	GLM	CBPS1	CBPS2	True	GLM	CBPS1	CBPS2	True
(1) Both r	nodels cor	rect							
	HT	0.33	2.06	-4.74	1.19	12.61	4.68	9.33	23.93
n — 200	IPW	-0.13	0.05	-1.12	-0.13	3.98	3.22	3.50	5.03
11 = 200	WLS	-0.04	-0.04	-0.04	-0.04	2.58	2.58	2.58	2.58
	DR	-0.04	-0.04	-0.04	-0.04	2.58	2.58	2.58	2.58
	HT	0.01	0.44	-1.59	-0.18	4.92	1.76	4.18	10.47
n - 1000	IPW	0.01	0.03	-0.32	-0.05	1.75	1.44	1.60	2.22
<i>II</i> = 1000	WLS	0.01	0.01	0.01	0.01	1.14	1.14	1.14	1.14
	DR	0.01	0.01	0.01	0.01	1.14	1.14	1.14	1.14
(2) Prope	nsity score	e model	correct						
	HT	-0.05	1.99	-4.94	-0.14	14.39	4.57	9.39	24.28
n — 200	IPW	-0.13	0.02	-1.13	-0.18	4.08	3.22	3.55	4.97
11 = 200	WLS	0.04	0.04	0.04	0.04	2.51	2.51	2.51	2.51
	DR	0.04	0.04	0.04	0.04	2.51	2.51	2.52	2.51
	HT	-0.02	0.44	-1.67	0.29	4.85	1.77	4.22	10.62
n 1000	IPW	0.02	0.05	-0.31	-0.03	1.75	1.45	1.61	2.27
11 - 1000	WLS	0.04	0.04	0.04	0.04	1.14	1.14	1.14	1.14
	DR	0.04	0.04	0.04	0.04	1.14	1.14	1.14	1.14

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## **CBPS Makes Weighting Methods Work Better**

		Bias					RMS	E	
	Estimator	GLM	CBPS1	CBPS2	True	GLM	CBPS1	CBPS2	True
(3) Outco	me model	correct							
	HT	24.25	1.09	-5.42	-0.18	194.58	5.04	10.71	23.24
n 200	IPW	1.70	-1.37	-2.84	-0.26	9.75	3.42	4.74	4.93
11 = 200	WLS	-2.29	-2.37	-2.19	0.41	4.03	4.06	3.96	3.31
	DR	-0.08	-0.10	-0.10	-0.10	2.67	2.58	2.58	2.58
	HT	41.14	-2.02	2.08	-0.23	238.14	2.97	6.65	10.42
n 1000	IPW	4.93	-1.39	-0.82	-0.02	11.44	2.01	2.26	2.21
n = 1000	WLS	-2.94	-2.99	-2.95	0.20	3.29	3.37	3.33	1.47
	DR	0.02	0.01	0.01	0.01	1.89	1.13	1.13	1.13
(4) Both I	models inc	correct							
	HT	30.32	1.27	-5.31	-0.38	266.30	5.20	10.62	23.86
n 000	IPW	1.93	-1.26	-2.77	-0.09	10.50	3.37	4.67	5.08
n = 200	WLS	-2.13	-2.20	-2.04	0.55	3.87	3.91	3.81	3.29
	DR	-7.46	-2.59	-2.13	0.37	50.30	4.27	3.99	3.74
	HT	101.47	-2.05	1.90	0.01	2371.18	3.02	6.75	10.53
n 1000	IPW	5.16	-1.44	-0.92	0.02	12.71	2.06	2.39	2.25
n = 1000	WLS	-2.95	-3.01	-2.98	0.19	3.30	3.40	3.36	1.47
	DR	-48.66	-3.59	-3.79	0.08	1370.91	4.02	4.25	1.81

## **CBPS Sacrifices Likelihood for Better Balance**



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Covariate Balancing Propensity Score

# Revisiting Smith and Todd (2005)

- Evaluation bias: "true" bias = 0
- CBPS improves propensity score matching across specifications and matching methods
- However, specification test rejects the null

	1-to-1	Nearest Ne	ighbor	Optimal 1-to-N Nearest Neighbor			
Specification	GLM	CBPS1	CBPS2	GLM	CBPS1	CBPS2	
Linear	-1209.15	-654.79	-505.15	-1209.15	-654.79	-130.84	
	(1426.44)	(1247.55)	(1335.47)	(1426.44)	(1247.55)	(1335.47)	
Quadratic	-1439.14	-955.30	-216.73	-1234.33	-175.92	-658.61	
	(1299.05)	(1496.27)	(1285.28)	(1074.88)	(943.34)	(1041.47)	
Smith & Todd	-1437.69	-820.89	-640.99	-1229.81	-826.53	-464.06	
	(1256.84)	(1229.63)	(1757.09)	(1044.15)	(1179.73)	(1130.73)	

## Comparison with the Experimental Benchmark

- LaLonde, Dehejia and Wahba, and others did this comparison
- Experimental estimate: \$866 (s.e. = 488)
- LaLonde+PSID pose a challenge: e.g., GenMatch -571 (1108)

	1-to-1	Nearest Ne	ighbor	Optimal 1-to-N Nearest Neighbor			
Specification	GLM	CBPS1	CBPS2	GLM	CBPS1	CBPS2	
Linear	-304.92	423.30	183.67	-211.07	423.30	138.20	
	(1437.02)	(1295.19)	(1240.79)	(1201.49)	(1110.26)	(1161.91)	
Quadratic	-922.16	239.46	1093.13	-715.54	307.51	185.57	
	(1382.38)	(1284.13)	(1567.33)	(1145.82)	(1158.06)	(1247.99)	
Smith & Todd	-734.49	-269.07	423.76	-439.54	-617.68	690.09	
	(1424.57)	(1711.66)	(1404.15)	(1259.28)	(1438.86)	(1288.68)	

#### Software: R Package CBPS

```
## upload the package
library("CBPS")
## load the LaLonde data
data(LaLonde)
## Estimate ATT weights via CBPS
fit <- CBPS(treat \sim age + educ + re75 + re74 +
                     I(re75==0) + I(re74==0),
            data = LaLonde, ATT = TRUE)
summary(fit)
## matching via MatchIt
library (MatchIt)
## one to one nearest neighbor with replacement
m.out <- matchit(treat ~ 1, distance = fitted(fit),
                 method = "nearest", data = LaLonde,
                 replace = TRUE)
summary(m.out)
```

## Extensions to Other Causal Inference Settings

- Propensity score methods are widely applicable
- This means that CBPS is also widely applicable
- Non-binary treatment regimes
- Imai, K. and van Dyk, D. (2004). "Causal Inference with General Treatment Regimes: Generalizing the Propensity Score" *Journal of the American Statistical Association*
- Challenge: many treatment groups ⇒ covariate balance checking is difficult
- Estimate the generalized propensity score such that covariate is balanced across *all* treatment groups

## **Multi-valued Categorical Treatment**

• Propensity score for each value:

$$\pi_{\beta}(t, X_i) = \Pr(T_i = t \mid X_i)$$

- Commonly used model: multinomial logistic regression
- CBPS: balance covariates across all groups

$$\mathbb{E}\left\{\frac{\mathbf{1}\{T_i=t\}X_i}{\pi_{\beta}(t,X_i)}\right\} = \mathbb{E}\left\{\frac{\mathbf{1}\{T_i=t'\}X_i}{\pi_{\beta}(t',X_i)}\right\}$$

- Orthogonalize the conditions when the number of groups is 2<sup>J</sup>
- Estimation of ATE: weighting or multi-dimensional matching/subclassification

## **Continuous and Other Treatments**

• Generalized propensity score:

$$\pi_{\beta}(t,X_i) = p(T_i = t \mid X_i)$$

- Propensity function:  $\psi_{\beta}(X_i)$  where  $p_{\psi}(T_i = t \mid X_i)$
- Commonly used models: linear regression, GLMs

$$\pi_{\beta}(t,X_{i}) = \frac{1}{\sqrt{2\pi\sigma^{2}}} \exp\left\{-\frac{1}{2\sigma^{2}}(t-X_{i}^{\top}\beta)^{2}\right\}, \quad \psi_{\beta}(X_{i}) = X_{i}^{\top}\beta$$

- CBPS: balance covariates across discretized treatment categories
- Estimation of causal effects:
  - subclassification on propensity function (Imai and van Dyk)
  - subclassification on treatment (Zhao, van Dyk, and Imai)
  - smooth coefficient model (Zhao, van Dyk, and Imai)

- Covariate balancing propensity score:
  - simultaneously optimizes prediction of treatment assignment and covariate balance under the GMM framework
  - is robust to model misspecification
  - improves propensity score weighting and matching methods

#### • Extensions:

- Non-binary treatment regimes
- Dynamic treatment regimes in longitudinal analysis
- Generalizing experimental estimates
- Generalizing instrumental variable estimates
- Weighting methods for causal mediation analysis
- Model and confounder selection in a high-dimensional setting