

Estimating Causal Effects of Credit Decisions Using Propensity Score Methodologies

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Agenda



- » Objective and Background
- » Case Studies:
 - » Risk Based Pricing
 - » Credit Line Increase
- » Discussion

Objective



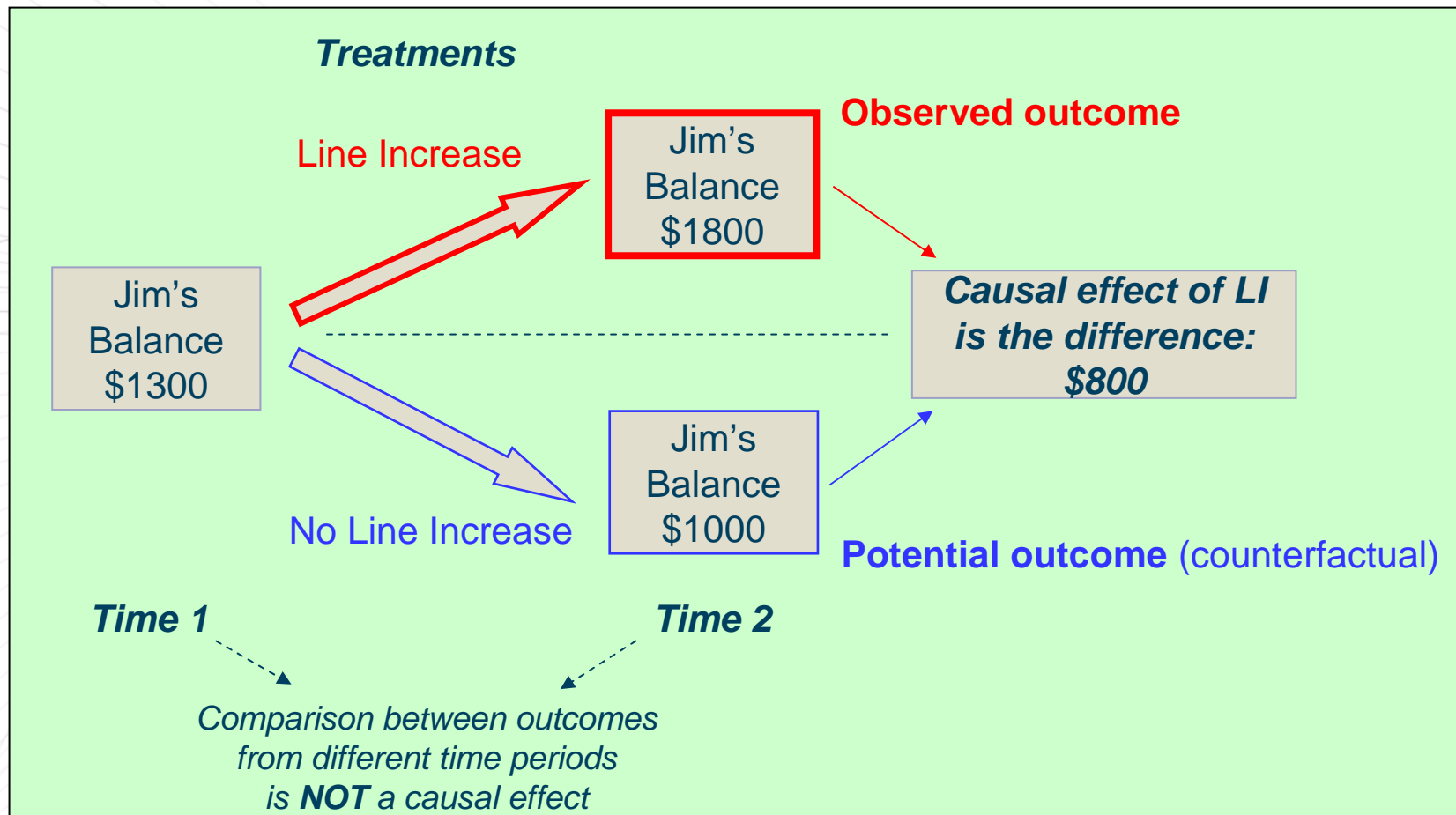
» Lenders and marketers want to determine the best credit decisions, offers, customer treatments

For this, they would like to understand the *causal effects* of their actions on future outcomes

Causal effect: “what will happen if...?”

» Rubin’s Causal Model

- » Causal effects are a comparison of outcomes from two or more alternative treatments, with only one of the outcomes observed



Responses to causal effect estimation problem



- » Gold standard
 - » Randomized experiments generate maximally informative data to estimate causal effects of interest

Example*

	Nonsmokers	Cigarette Smokers	Cigar and Pipe Smokers
Mortality Rates per 1000 person-years	13.5	13.5	17.4

- » Treatments (self-selected, not randomly assigned)
 - » Nonsmoking
 - » Cigarette smoking
 - » Cigar and pipe smoking
- » Naïve conclusion
 - » Cigarette smoking is good for your health, especially relative to cigar and pipe smoking
 - ...may be misleading

*Adapted from Rubin and Cochran

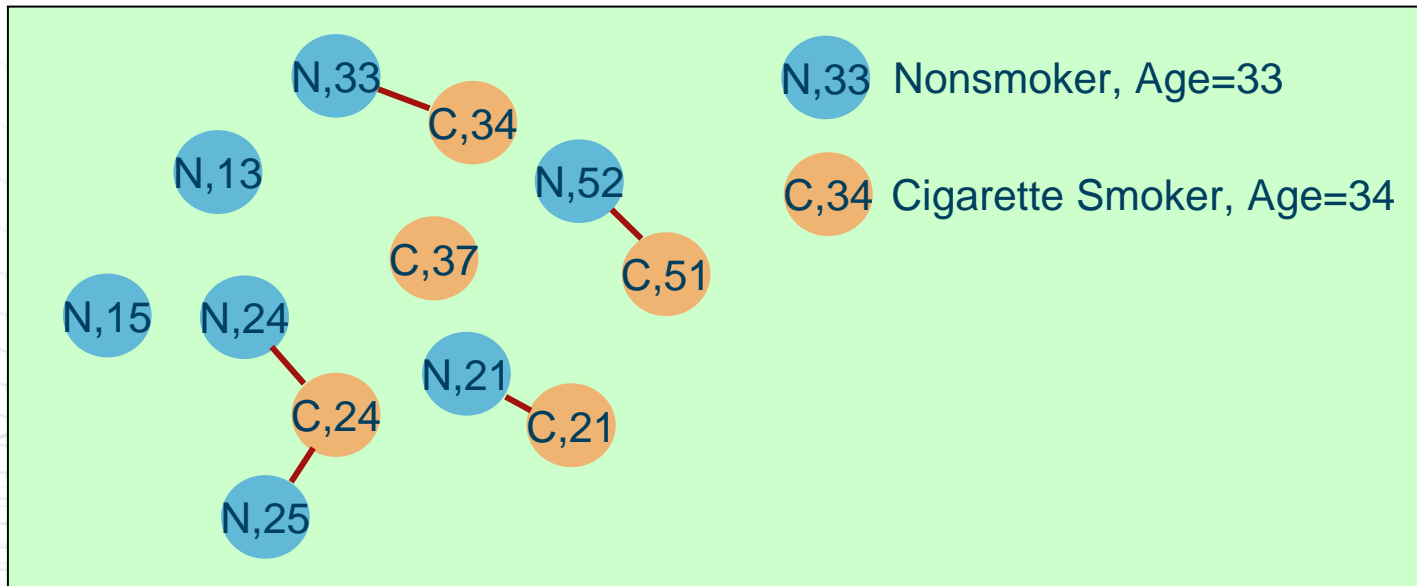
Age is a confounding variable



	Nonsmokers	Cigarette Smokers	Cigar and Pipe Smokers
Mortality Rates per 1000 person-years	13.5	13.5	17.4
Average Age	57.0	53.2	59.7

- » Age is correlated with mortality and also with smoking habits
 - » Conclusions about effects of smoking should be adjusted for age
- » Adjustment approaches include
 - » Regression analysis
 - » Subclassification
 - » Matched sampling

Matched sampling on Age*



- » For each nonsmoker, find a cigarette smoker of similar age if possible. Analogous for cigar and pipe smokers
- » Age will be balanced between the matched samples
- » Compare mortality rates based on matched samples only
 - » Some samples may be discarded for estimation

* Many variants of matching, see e.g. D. Rubin's book, "*Matched Sampling for Causal Effects*"

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Effect of risk-based pricing on Take Rate



Risk Premium	Base Rate	Base + 3%	Base + 5%
Take Rate (TR)	68%	37%	44%

- » Naïve conclusions:
 - » Changing from (Base+3%) to (Base+5%) increases TR by 7%
...doesn't make sense, assuming consumers behave rational
 - » Changing from Base Rate to (Base+3%) decreases TR by 31%
...makes more sense, but estimate may nevertheless be biased

APR offer was determined in two steps:

1. Decide on Base Rate (not discussed here)
2. Decide on Risk Premium to add to Base Rate

Should adjust for many covariates



Risk Premium	Base Rate	Base + 3%	Base + 5%
Take Rate (TR)	68%	37%	44%
Application Score (mean)	172	140	134
Applicant Age (mean)	47	42	37
Channel: Supermarket	30%	36%	52%
Internet	16%	20%	1%
Marketing	54%	44%	47%
Residence: Owns	86%	68%	51%
Rents	11%	29%	45%
Other	3%	3%	4%
.. many other covariates

Data spanning across multiple marketing channel

Different marketing channels used different risk-based pricing strategies

Outline of process steps

- I. Develop *propensity score(s)* between alternative treatments
 - » Will simplify matching when many covariates need adjustment
- II. Generate *matched samples* based on propensity scores
 - » Matched data will resemble data from randomized experiments
 - » Useful for descriptive analysis and as basis for regression estimation
- III. Estimate *causal effects* from matched samples
 - » Will use regression estimation techniques

Benefits

- » Reduced risk of confounding causal effect estimates
- » Clearer appreciation of what information is in the data and what isn't
- » More objective assessment of causal relationships

Step I: Develop propensity score(s)



- » Propensity score models probability of individuals receiving treatment (as opposed to control), given covariates

$$\Pr\{T = 1 | X\} = p(X)$$

where :

$T : \{0,1\}$ treatment indicator (0 : control, 1 : treatment)

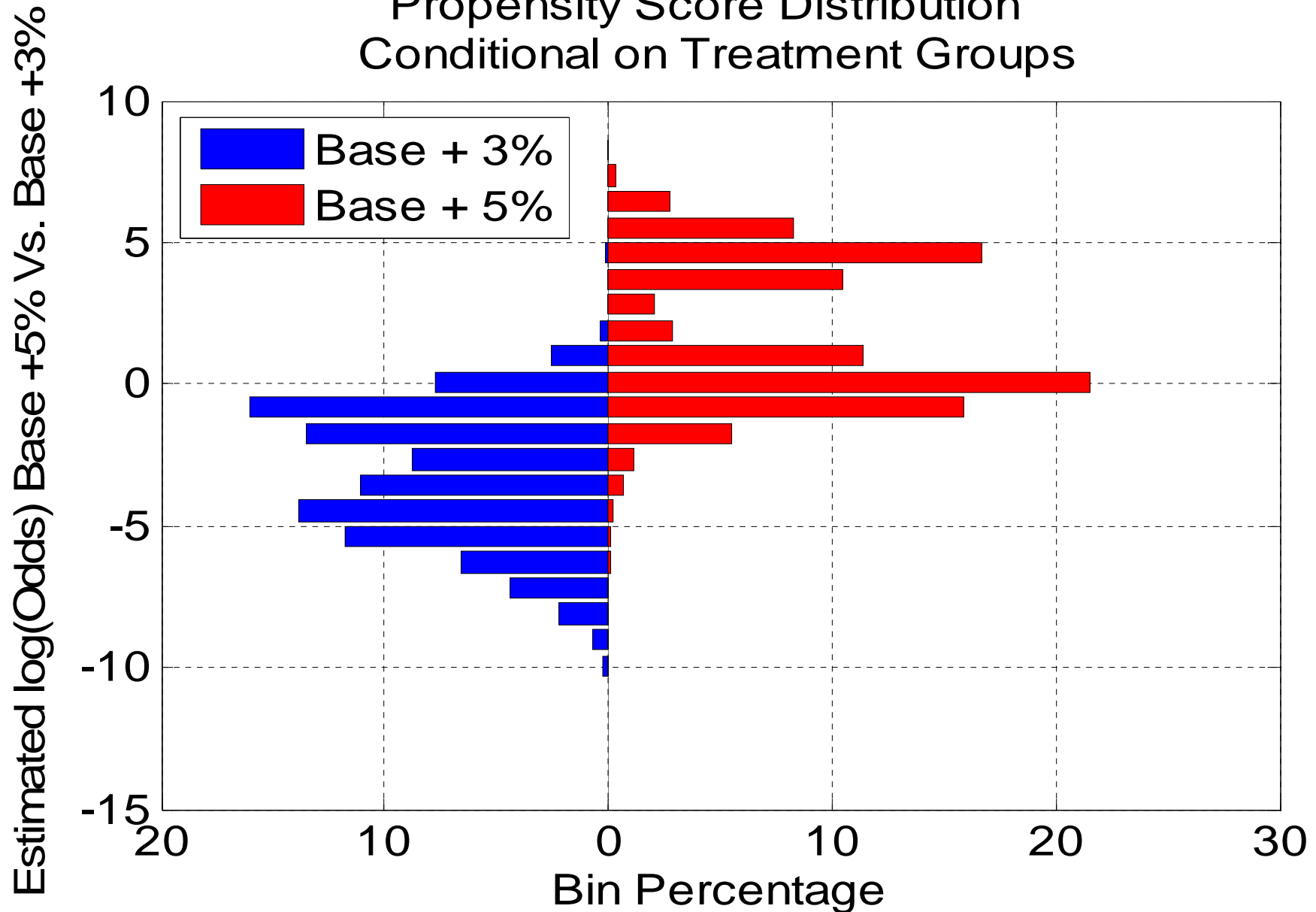
X : Covariates

- » Score is developed from observed data
- » May estimate using logistic regression, scorecard or other generalized additive models, random forests, neural nets, etc.
- » Develop multiple propensity scores for various treatment dichotomies, such as:
 - » Control = Base Rate Control = Base + 3%
 - » Treatment = Base + 3% Treatment = Base + 5% etc.

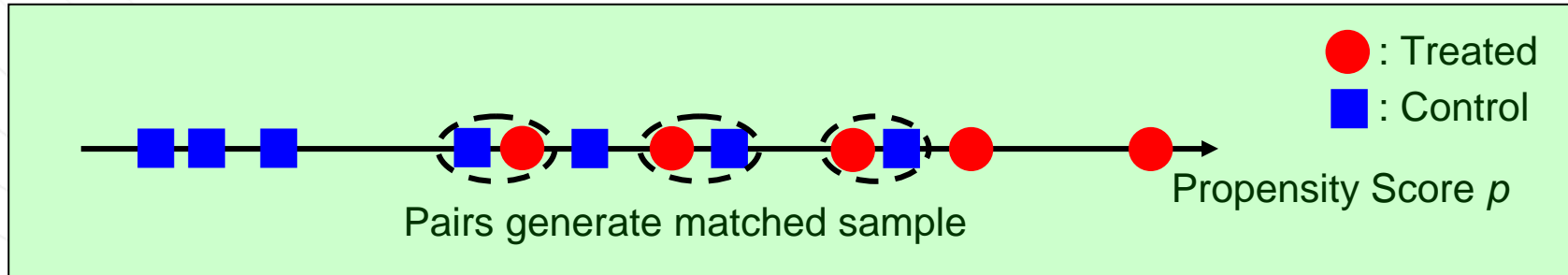
Limited overlap between Base+3% and Base+5%



Propensity Score Distribution Conditional on Treatment Groups



Step II: Matching on propensity score



- » Propensity score simplifies matching, by reducing all covariates to be matched, to a single covariate (p) to be matched
- » If the propensity score is reasonably developed, then, within any stratum of p that contains treated and controls, treated individuals are similar to controls in terms of their covariate distributions
- » Practical assessment of estimated propensity score
 - » Do the matching
 - » Covariates should be well balanced on the matched sample
 - » Diagnosis of balance is important. Seek to improve propensity score until satisfied with balance, *before* proceeding with estimation step

Risk-based pricing example before and after matching



Observed sample
N=11,837
Unbalanced covariates

Matched sample
N=3,404
Balanced covariates

APR Premium (treatment)	Base + 3%	Base + 5%	Base + 3%	Base + 5%
Application Score	140	134	137	137
Applicant Age	42	37	40	39
Channel: Supermarket	36%	52%	50%	51%
Internet	20%	1%	1%	1%
Marketing	44%	47%	49%	48%
.. many other covariates

- » Matched sample balances all covariates going into propensity score
 - » (Almost) as if data came from a randomized experiment

Step III: Regression analysis on matched sample

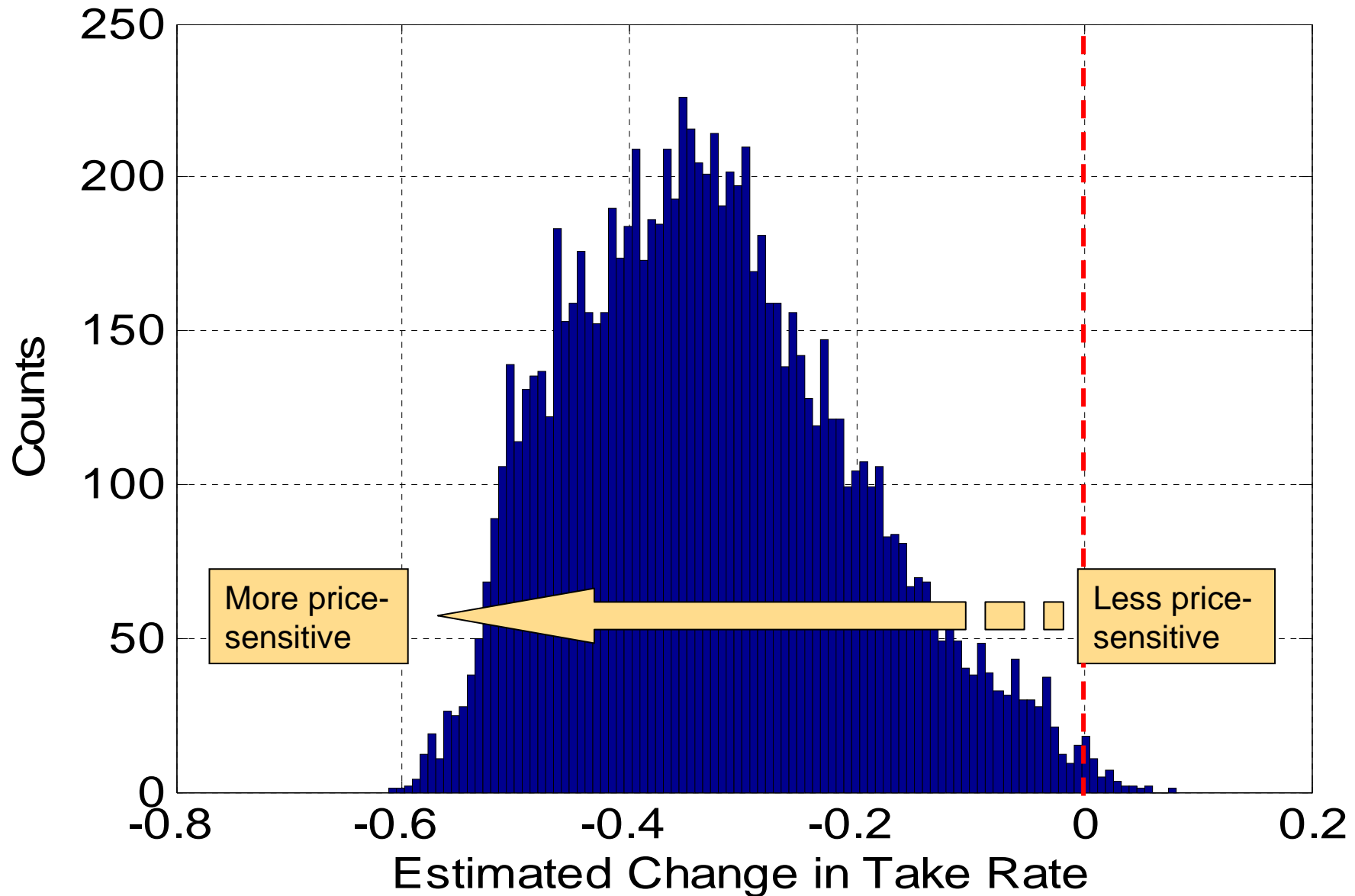


- » Include treatment variable, along with covariates, in regression model (GLM, GAM, Scorecard, NN, ...) to predict outcomes
 - » Specify model depending on nature of data and analysis goals
- » Models with additive treatment effect:
 - » May improve estimate of average treatment effect
 - » By further adjusting for covariate distributions
- » Testing for treatment x covariate interactions:
 - » Do treatment effects differ between individuals?
 - » Infer segment-level or individual-level treatment effects

Effects of changing treatment from Base \rightarrow Base + 3% are heterogeneously distributed



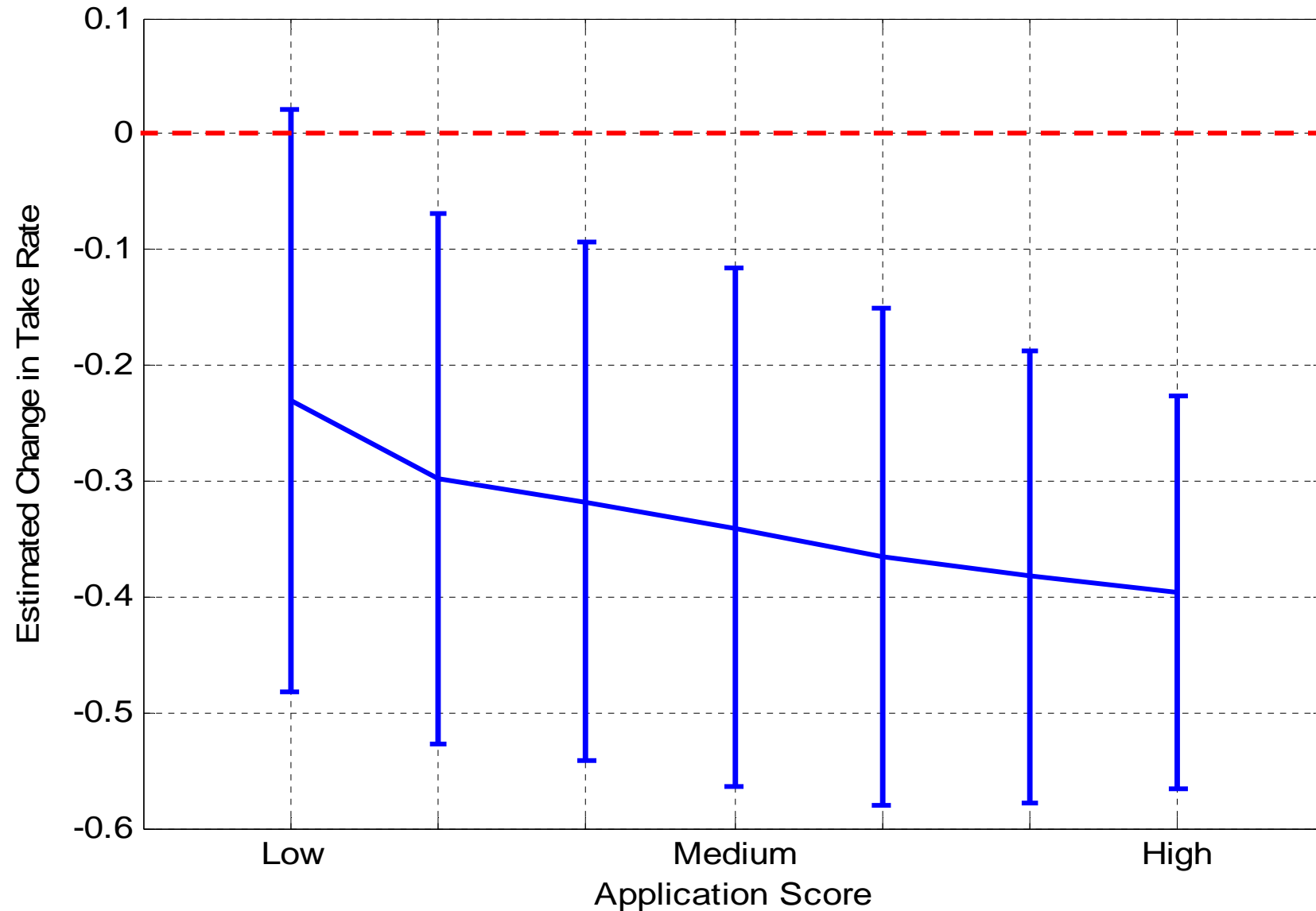
Effect Distribution in Matched Sample



Price sensitivity (Base \rightarrow Base + 3%) By Risk band



Mean Effects and 95% Prediction Intervals by Application Score Band



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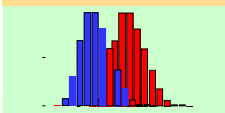
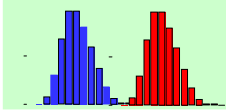


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Line increases in a credit card portfolio



- » Mostly deterministic line increase strategy (very limited testing)
 - » Increase amounts in multiples of \$500
- » Analyzed all treatment-control pairs and generated matched samples, where possible
 - » In tendency, overlap decreased with distance between treatments (with some exceptions). Example:

Control [\$]	Treatment [\$]	Overlap	P'score distributions
1,000	1,500	23%	
1,000	2,000	26%	
1,000	2,500	13%	
1,000	3,000	5%	
1,000	3,500	0%	

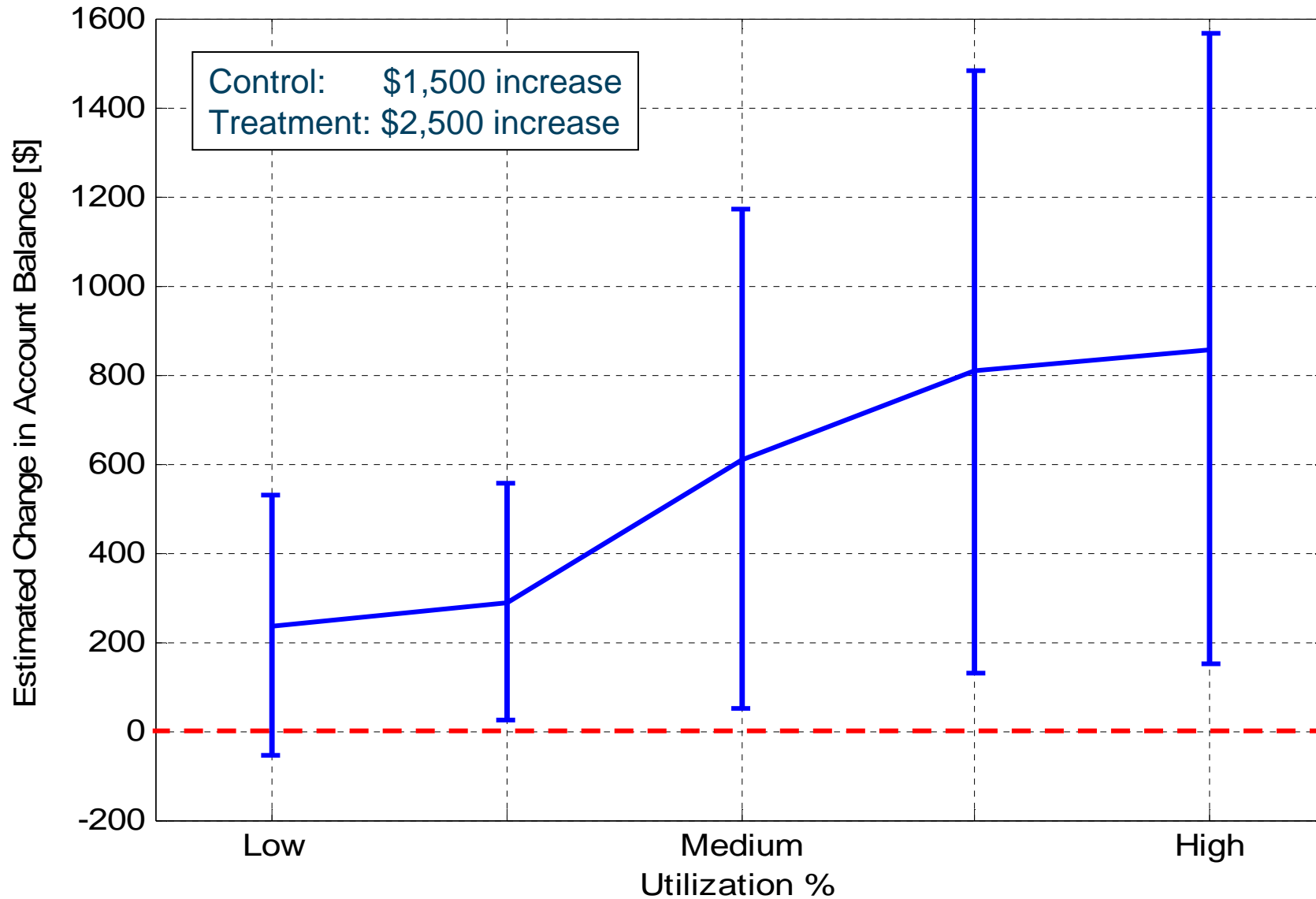
- » Some comparisons struggled with too few matched samples. More aggressive testing would have been a boon for analysis
- » But decent overlap across various treatment-control pairs, even with limited testing, is encouraging news for causal modeling

Effect of line increase on account balance

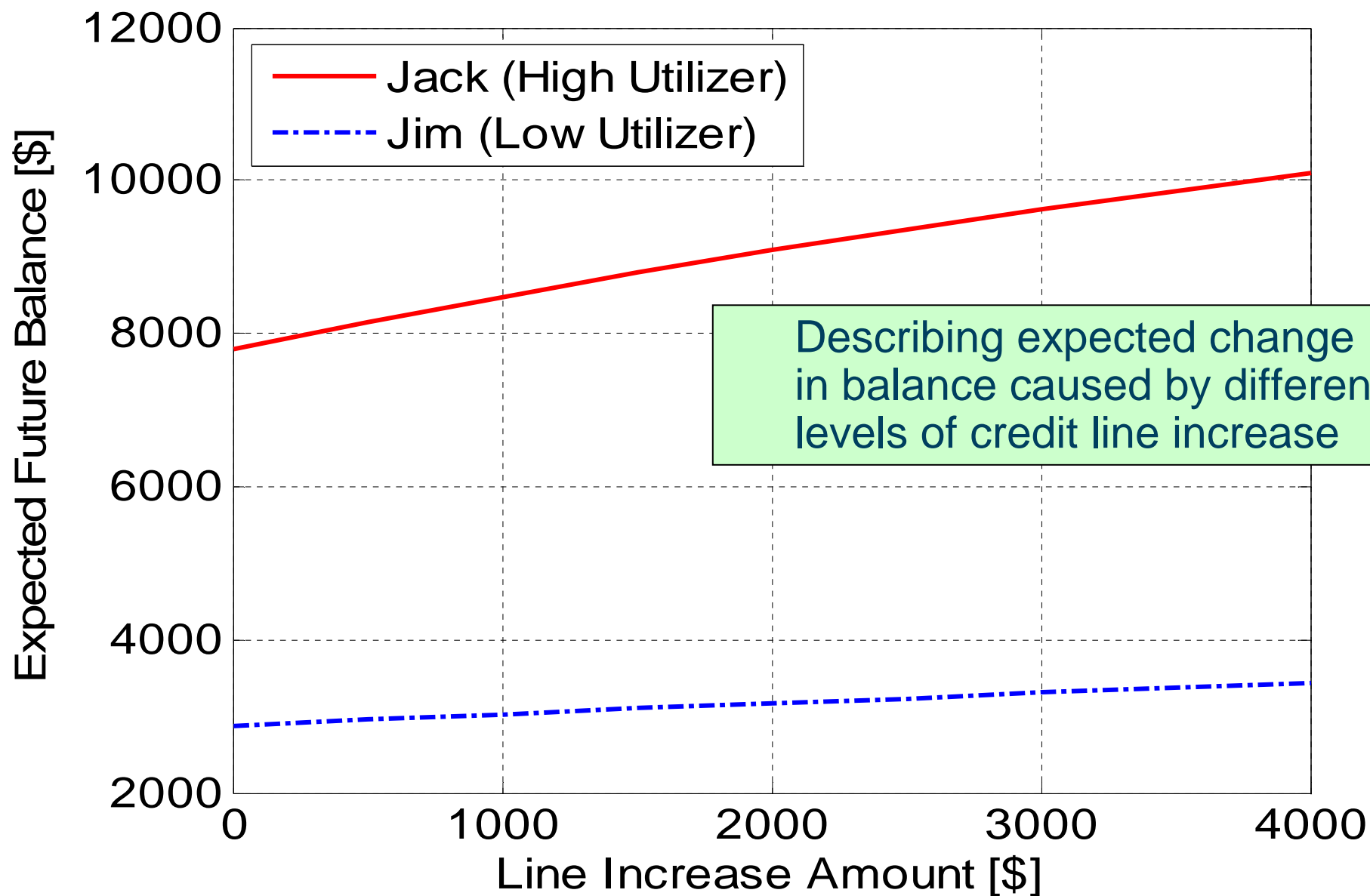
By Utilization band



Mean Effects and 95% Prediction Intervals by Utilization %



Credit line increase-response curves



- » Selection bias in business-as-usual data poses analytic challenges for estimating causal effects
- » Direct regression analysis of BAU data can yield unreliable results
- » Benefit of matching prior to regression analysis
- » No miracle cure exists if important confounders are not collected

- » Mainstays of analytic success continue to include:
 - » Purposeful design of experiments
 - » Exhaustive collection of covariates associated with treatments or outcomes
 - » Sniffer dog capabilities to sense lurking variables
 - » Incorporation of domain expertise into the modeling process

Landmark paper on properties and virtues of the propensity score:

“The Central Role of the Propensity Score in Observational Studies for Causal Effects”, by Paul R. Rosenbaum and Donald B. Rubin, *Biometrika*, Vol. 70, No. 1. (Apr., 1983), pp. 41-55.

Assembly of many papers from introductory to advanced, and good advice:

“*Matched Sampling for Causal Effects*”, by Donald B. Rubin, Cambridge University Press, 2006.

For a leisurely introduction into propensity score ideas:

“Estimating Causal Effects from Large Data Sets Using Propensity Scores”, by Donald B. Rubin, *Annals of Internal Medicine*, Volume 127(8S), October 1997, pp 757-763.

THANK YOU

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Relative distribution analysis over time

How treatment affects account balance distribution



Reference distribution (matched controls): No increase
Comparison distribution (matched treated): \$1,000 increase

For each quantile of the reference distribution, *relative density* indicates enrichment (if >1), or depletion (if <1) of comparison distribution in this quantile.

