# Clever Randomization and Ensembling Strategies for Accommodating Multiple Data Pathologies in Conjoint Studies

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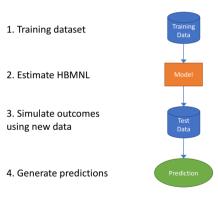
Marketing Science

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## Conjoint Analysis in Marketing

- Products are defined by attributes, each with a number of levels
- Individuals choose from among a number of alternatives
- We infer respondent-level preferences for each attribute level
- We make use of preference estimates to predict demand for potential products compared with the competition
- ► These **market simulators** can inform new product development, pricing, product line optimization, and go-no-go decision-making

## Current Approach to Conjoint Prediction



#### **Data Pathologies**

**Pathology:** Structural and functional **deviations from the normal** that constitute disease or characterize a particular disease

**Data Pathology:** Any **respondent-level behavior** that deviates from the **normative model** of consumer choice, thus impeding accurate out-of-sample prediction

#### Normative Model of Consumer Choice

For a given individual and choice task:

$$y \in \{1, 2, \cdots, J\}$$
 $U_j = V_j + \varepsilon_j$ 
 $V_j = \beta_1 x_{1j} + \beta_2 x_{2j} + \beta_k x_{kj}$ 
 $\varepsilon_j \sim \mathsf{iid} \; \mathsf{Gumbel}$ 

Across respondents:

$$eta \sim N\left(ar{eta}, \Sigma
ight)$$

#### Models of Data Pathologies

- ▶ Many innovations in conjoint focus on addressing specific data pathologies
- These models fit better and provide marginal improvements in predictive fit
- Although prospectively useful, these models are rarely used in practice

## Examples of Data Pathologies

- Attribute Non-Attendance
- Screening Rules
- ► Poor Respondent Quality
- ► Respondent Fatigue
- ► Alternative Decision Rules (e.g., Regret Minimization)

#### Difficulties in Practice

- ► They are theoretically and computationally complex (i.e., difficult to understand and time-consuming to estimate and simulate)
- We don't have high-quality commercial software that can be used to fit these models
- Each model deals with a single data pathology
  - A priori how do I know which pathology will prove problematic (i.e., which model should I fit)?
  - What if multiple pathologies are present in a single dataset?

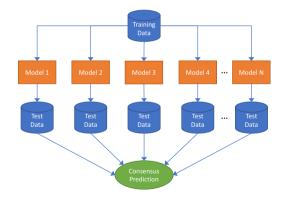
## Ensemble Approaches to Prediction

- With a few exceptions, most conjoint projects are still conducted using a single model and dataset for both inference and simulation
- ► However, **ensemble approaches** currently dominate the world of competitive prediction (e.g., Kaggle, Netflix Prize, etc.)

#### An Ensemble Approach to Prediction



- 2. Estimate different types of models
- 3. Simulate outcomes on new data
- 4. Generate a consensus prediction



#### Why Do Ensembles Perform So Well?

- They strike an optimal trade-off on the bias/variance frontier
  - Wisdom of the Crowd
- ▶ They allow for efficient exploration of the model space
  - All models are wrong, but some models are useful
- ► They provide a hedge against data conditions that are pathological with respect to a single model (our hypothesis)

## Our Proposed Approach

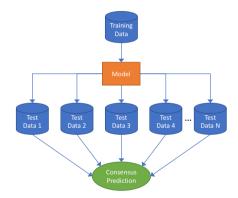
- Create an ensemble where diversity is created through various forms of (clever)
   randomization
- Induce diversity in the ensemble by thinking about how specific forms of randomization are related to specific data pathologies
- ► Generate a consensus prediction from the ensemble

Note: Our focus in this study is to improve **out-of-sample predictive validity**!

## An Ensemble Approach to Conjoint Prediction

1. Training dataset

- 2. Estimate HBMNL
- 3. Simulate outcomes using random versions of holdout data
- 4. Generate a consensus prediction



# Pathology 1: Attribute Non-Attendance (ANA)

**Attribute Non-Attendance**: Respondents ignore subsets of attributes when making decisions (i.e.,  $\beta = 0$  for all levels of the attribute)

**Randomization Strategy for the Ensemble**: Create an ensemble where we randomly set the coefficients for a full attribute to 0

#### Ensemble Algorithm for Attribute Non-Attendance

- 1. Estimate an HB MNL on training data
- 2. Loop over respondent-level estimates  $(\hat{\beta}_h)$ 
  - randomly select an attribute
  - with probability heta set all coefficients for that attribute to 0
- 3. Predict first choices (e.g., max utility) for each choice set in the out-of-sample holdout (test) data
- 4. Repeat 2 and 3 many times
- 5. Generate a consensus (e.g., most commonly selected) prediction

## Pathology 2: Screening Behavior

**Screening Behavior**: Respondents use certain attribute levels to screen out alternatives from consideration (i.e.,  $\beta \approx -\infty$  for levels being screened on)

Randomization Strategy for the Ensemble: Create an ensemble where we randomly set the coefficients for attribute levels to  $\approx -\infty$ 

## Ensemble Algorithm for Screening Behavior

- 1. Estimate HB MNL on training data
- 2. Loop over respondent-level estimates  $(\hat{\beta}_h)$ 
  - randomly select an attribute level
  - with probability  $\psi$  set the coefficient for that attribute level to  $\approx -\infty$
- 3. Predict first choices (e.g., max utility) for each choice set in the validation data
- 4. Repeat 2 and 3 many times
- 5. Generate a consensus (e.g., most commonly selected) prediction

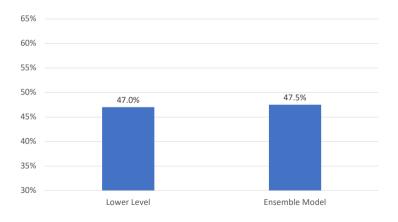
# Joint Ensemble Algorithm for Attribute Non-Attendance and Screening

- 1. Estimate HB MNL on training data
- 2. Loop over respondent-level estimates  $(\hat{\beta}_h)$ 
  - randomly select an attribute level
  - with probability  $\psi$  set the coefficient for that attribute level to  $\approx -\infty$
  - randomly select an attribute
  - with probability heta set all coefficients for that attribute to 0
- 3. Predict first choices (e.g., max utility) for each choice set in the validation data
- 4. Repeat 2 and 3 many times
- 5. Generate a consensus (e.g., most commonly selected) prediction

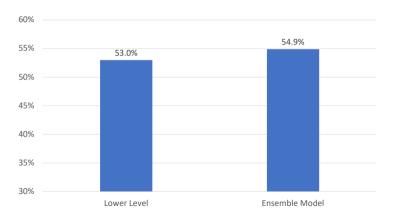
## Simulation Study

- ► Simulate data with the presence or absence of pathologies: attribute non-attendance and/or screening
- ▶ 500 respondents; 12 choice tasks; 6 attributes
- Generate prediction from standard HBMNL results (lower level) and the ANA, Screening, or joint ANA + Screening Ensemble

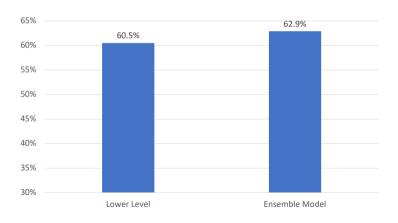
# Simulated Data: No ANA + No Screening



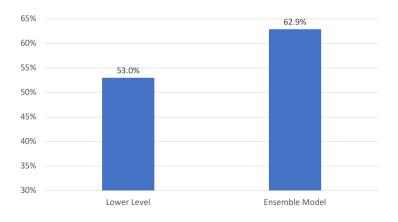
# Simulated Data: ANA without Screening



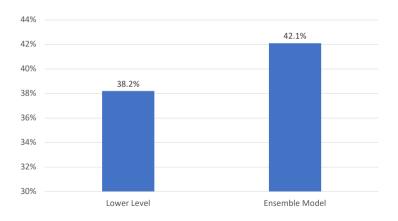
# Simulated Data: Screening without ANA



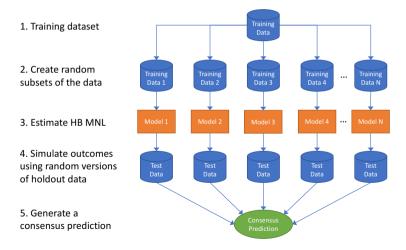
# Simulated Data: ANA + Screening



#### Performance on Interior Paint Data



#### Another Ensemble Approach to Conjoint Prediction



#### Next Steps

- Utilize computational advantages of Hamiltonian Monte Carlo via Stan
- Explore clever randomization strategies on test data
- Include more pathologies
- ▶ Better aggregation of prediction (e.g., model stacking via a meta-learner rather than consensus predictions)

#### Conclusion

- Cleverly designed ensembles of models appear to help hedge against data pathologies and improve prediction
- ► The ensemble approach is especially powerful in the presence of multiple data pathologies
- ► Ensembles for inference: Can we learn about the severity of a particular pathology by examining the relative fit of ensemble members as part of a meta-learner?

#### Thank You!

Questions? Contact me at marc.dotson@byu.edu.