Image-based Treatment Effect Heterogeneity

Abstract

Randomized controlled trials (RCTs) are considered the gold standard for estimating the average treatment effect (ATE) of interventions. One important use of RCTs is to study the causes of global poverty—a subject explicitly cited in the 2019 *Sveriges Riksbank Prize in Economic Sciences in Memory of Alfred Nobel* awarded to Duflo, Banerjee, and Kremer "for their experimental approach to alleviating global poverty." Because the ATE is a population summary, anti-poverty experiments often seek to unpack the treatment effect variation around the ATE by conditioning (CATE) on tabular variables such as age and ethnicity that were measured during the RCT data collection. Although such variables are key to unpacking CATE, using only such variables may fail to capture historical, geographical, or neighborhood-specific contributors to effect variation, as tabular RCT data are often only observed near the time of the experiment. In global poverty research, when the geographical location of the experiment units is approximately known, satellite imagery can provide a window into such historical and geographical factors important for understanding heterogeneity. However, there is no causal inference method that specifically enables applied researchers to analyze CATE from images. In this paper, using a deep probabilistic modeling framework, we develop such a method that estimates latent clusters of images by identifying images with similar treatment effects distributions. Our model also emphasizes an image sensitivity factor that quantifies the importance of image segments in contributing to the mean effect cluster prediction, obtained via Monte Carlo using the approximate posterior distribution over the clustering. We compare the proposed methods against alternatives in simulation; additionally, we show how the model works in an actual RCT, estimating the effects of an anti-poverty intervention in northern Uganda and obtaining a posterior predictive distribution over treatment effects for the rest of the country were no experimental data was collected. We make code for all modeling strategies available in an open-source software package and discuss their applicability in other domains—such as the biomedical sciences—where image data is also prevalent.

Keywords: Causal inference; Treatment effect heterogeneity; Remove sensing; Image data; Probabilistic reasoning

1. Introduction

Field experiments in the social and health sciences help us understand the effects of interventions in the natural habitat where people live (Banerjee et al., 2011). Their primary goal is often to identify the overall effect of a treatment T_i on an outcome Y_i, marginalizing over units ($i \in \mathcal{I}$) in the sample population, and thereby, calculating the average treatment effect (ATE). By collecting tabular characteristics, $\mathbf{X}_i$, such as age and sex, investigators can unpack results by sub-populations, estimating the Conditional Average Treatment Effects (CATEs) (Künzel et al., 2019; Balgi et al., 2022). However, the features in $\mathbf{X}_i$ are often measured only at baseline, just before the experiment is initiated. Thus, $\mathbf{X}_i$ rarely contain information on an experimental unit's historical characteristics, including its past neighborhood-level features and geographical context, that may be useful in identifying sub-populations that react differently to the same treatment (Kino et al., 2021).

When experimental units i are associated with a particular location, satellite images, $\mathbf{M}_i$, can fill in this gap, providing pieces of the necessary information about the otherwise missing historical, neighborhood, and geographical contexts (Burke et al., 2021; Daoud et al., 2021).

Indeed, in contrast to covariates measured during experiments, satellite imagery is collected passively from space for every geographic context on earth, and thus (except for clouds covering the line sight of a satellite) there is no systematic missingness in the data source. Moreover, these data have been collected for parts of the world since the CORONA intelligence satellites of the 1950s and for the entire world since the start of NASA's and the US Geological Survey's Landsat mission in the 1970s. The Landsat data are publicly available with a revisiting time of on average 16 days. Therefore, by combining satellite imagery with experimental data, researchers can investigate not only the historical and geographical roots of effect variation, but they can also predict how an intervention will likely impact places outside the scope of the original study where no researcher-collected covariate data is available. That is, predictive distributions over treatment effects can be estimated not only for the experimental context but also investigated in places not originally conceived of during the experiment—where no tabular background covariates were measured. In this way, remote sensing data has the potential to increase the applicability of ideas in casual transportability (Pearl and Bareinboim, 2022).

However, despite the potential offered by images for causal inference, as evident by the growing literature (Castro et al., 2020; Chalupka et al., 2016b,a, 2015; Schölkopf et al., 2021; Yi et al., 2020; Ding et al., 2021; Paciorek, 2010; Kaddour et al., 2021b; Louizos et al., 2017; Pawlowski et al., 2020; Lopez-Paz et al., 2017), it remains unclear how researchers should use these $\mathbf{M}_i$'s for CATE analysis. One key challenge is that $\mathbf{M}_i$ is high-dimensional and rarely annotated. A CATE analysis using $\mathbf{M}_i$ but relying on tabular methods, such as a linear model or the generalized random forest Athey et al. (2019), would likely lead to poor interpretability or struggle to find heterogeneity within the high-dimensional image objects.

Thus, there is a major research need to form a causal inference framework for CATE analysis in images. Equipped with such a method, applied researchers would not only be enabled to start incorporating satellite images in future RCTs, but also re-analyze data from past experiments that have already been performed, with the goal of deepening understanding using satellite image data, potentially uncovering missed yet significant CATE–informing policymaking.

In this article, we develop machine-learning models which identify image-based CATE in RCT data, by analyzing satellite imagery collected independently of the experiments. The model we develop is a probabilistic model, employing Bayesian convolutional neural network arms (CNNs) with categorical gates that allow us to directly model mixtures of image clusters with similar effects. Our models estimate treatment effects for all units, conditional on treatment status, the images $\mathbf{M}_i$, and, if desired, accounting for available $\mathbf{X}_i$ through orthogonalization. By residualizing, our model will identify what additional CATE that stems from $\mathbf{M}_i$, separately from $\mathbf{X}_i$, for enhanced heterogeneity analysis. The models construct image-type clusters that probabilistically categorize units based on their CATE similarity.

In the following sections, we develop our methods, show some of their properties analytically, and explore others in simulated experiments. We demonstrate the usefulness of our methods by replicating the results of an RCT study in Uganda—the Youth Opportunities Program (YOP) (Blattman et al., 2014). This study, conducted in 2008, was designed to help the poor break unemployment cycles by financially assisting their artisans and business activity. The government solicited young adults to participate in YOP, asking them to form teams and compose a business

plan. After screening these teams and their plans, the government randomly assigned a one-time grant worth about \$7,500 to each team. This amount equals approximately their joint annual income. Most of the applicants were young, rural farmers having low educational attainment ($\sim$ eighth grade), earning less than \$1 per day, and working less than twelve hours per week. In many RCTs like YOP, the researchers collected a set of baseline covariates, but none of them explicitly capture historical neighborhood or geographical characteristics. Our replication analysis demonstrates the benefits, and limitations, of using $\mathbf{M}_i$ for CATE, as a complement to the tabular version.

While our contribution focuses on the use of satellite images in global poverty research, our methods are designed such that they generalize to other forms of RCTs where complementary image data is available. In the last few decades, there has been a rapid increase in the availability of imaging technologies. Most notably, these technologies are readily available in biomedical fields in the form of X-ray, positron emission, MRI, and ultrasound modalities. These images data streams are likely useful not only for estimating ATE (Castro et al., 2020; Lopez-Paz et al., 2017; Chalupka et al., 2016b), but also for evaluating CATE. Nonetheless, more research is required to systematically evaluate the uses of our modeling approach for these domains.

2. Background and Related Work

Conditional Average Treatment Effects with Tabular Data   Let $Y_i(t)$ denote the potential outcome (Rubin, 2005) of an intervention $t \in \{0, 1\}$ for a unit of study i. For example, $Y_i(1)$ may represent the poverty level in household i following an aid intervention, and $Y_i(0)$ is the level without intervention.

We may define the unit-level treatment effect as $\tau_i = Y_i(1) - Y_i(0)$. When τ_i is greater than 0, the unit's outcome is greater under treatment than otherwise. The quantity τ_i cannot be exactly identified without strong assumptions (Pearl, 2009). Because a unit can only receive a single treatment at a given time, only either of the potential outcomes $Y_i(1)$ or $Y_i(0)$ are observed, and thus, the counterfactual remains unobserved. Assuming consistency—that is, units comply with their treatment assignment—the observed outcome can be written as, $Y_i = Y_i(T_i) = Y_i(1)T_i + Y_i(0)(1 - T_i)$, where T_i denotes the (random) treatment status of i (Miguel and Robins, 2020). The ATE captures the population effect by averaging over all unit-level effects:

$$\text{Average Treatment Effect (ATE):} \quad \overline{\tau} = \mathbb{E}[\tau_i] = \mathbb{E}[Y_i(1) - Y_i(0)].$$

The ATE is useful as it marginalizes over the heterogeneity present in a population to form an overall assessment of an experiment. In an RCT, due to treatment randomization, ATE can be estimated non-parametrically by the difference between treatment and control outcomes (Rubin, 2005). Despite the importance of aggregate quantities such as the ATE, it is useful to disaggregate average effects using based on contextual information. Such a disaggregation is critical for not only scientific understanding but also for policy and (e.g., by personalizing treatments (Greenland et al., 2020; Balgi et al., 2022). This disaggregation can be a function of any type of general pre-treatment data variable, $\mathbf{G}_i$:

$$\text{Conditional Average Treatment Effect (CATE):} \quad \tau(\mathbf{g}) = \mathbb{E}[Y_i(1) - Y_i(0) \mid \mathbf{G}_i = \mathbf{g}],$$

The literature has primarily focused on conditioning sets that contain tabular data, $\mathbf{X}_i$:

$$\text{Tabular Conditional Average Treatment Effect:} \quad \tau(\mathbf{x}) = \mathbb{E}[Y_i(1) - Y_i(0) \mid \mathbf{X}_i = \mathbf{x}],$$

where $\mathbf{x} \in \mathbb{R}^D$ denotes the vector of D pre-treatment covariates (Athey and Imbens, 2016; Athey et al., 2019; Ding et al., 2016; Imai and Ratkovic, 2013; Shalit et al., 2017; Zhao et al., 2017; Luedtke and van der Laan, 2016; Nie and Wager, 2021). For example, the generalized random forest is one such machine-learning method (Athey et al., 2019), and it has proven useful in a variety of applied settings (Shiba et al., 2021; Daoud and Johansson, 2019). However, these methods are tailored for annotated tabular data and images are high-dimensional, often non-annotated. These non-annotated image features consist of image bands and pixels that may jointly induce effect heterogeneity. Thus, more research is required to improve the ability of investigators to understand CATE in the context of unstructured high-dimensional image data.

Causal Inference with Image Data While most causal-inference studies use tabular data, there is an increasing realization that image data provides a creative yet useful way to conduct causal inference (Castro et al., 2020; Ramachandra, 2019; Daoud and Dubhashi, 2020). To this end, there is a growing methodological literature investigating how images should be integrated to identify and estimate ATEs in the observational setting (Kallus, 2020; Kaddour et al., 2021a; Pawlowski et al., 2020; Jerzak et al., 2022). Yet these approaches tend to mainly treat images as proxies, for inclusion in the adjustment set, thereby securing causal identification; these approaches are not tailored for CATE analysis in images. Hence, little is known about how to use images for CATE.

Like any tabular information in $\mathbf{X}_i$, images as a whole or through its segments may be associated with treatment effect heterogeneity. In the observational setting, images $\mathbf{M}_i$ could be part of both the conditioning and effect modification sets. In the RCT setting, $\mathbf{M}_i$ is not a confounder (and thus, unnecessary to identify ATE), since the treatment was randomized, yet it may still provide significant information for CATE (as discussed in §1). In both settings, one target estimand is the Image CATE,

$$\textit{Image CATE:} \quad \tau(\mathbf{m}) = \mathbb{E}[Y_i(1) - Y_i(0) \mid \mathbf{M}_i = \mathbf{m}],$$

where $\mathbf{m} \in \mathbb{R}^{W \times H \times D}$ is an image, obtained before treatment assignment, of width W, height H, and with D data channels. These data channels often correspond to reflectance information from various bands. While image and tabular CATEs are both special cases of the general CATE, there are conceptual differences. First, by their nature, images are unstructured, high-dimensional objects. In RCT applications, every image will likely be unique as image effects act much as individual-level treatment effects, where summarizing the quantity of interest becomes increasingly difficult with a growing number of units—aggravating the curse of dimensionality. Second, as images are unstructured, it is unclear how to interpret the act of conditioning on an image. We need a conceptual language and modeling strategy for characterizing proximity between images in the space of conditional effects. The rest of our article contributes to establishing this conceptual language, focusing for the sake of clarity on the RCT setting, and thus, leaving image-CATE in observational design for a future study.

3. Modeling Causal Effect Heterogeneity in Images

In experimental settings, CATEs may be estimated readily by function approximation. Perhaps the simplest approach is to use a parameterized function, $f_{\hat{Y}_t}(\mathbf{m})$, to predict observations of the potential outcome $Y(t)$ of each intervention—a so-called *T-learner* (Künzel et al., 2019). The CATE may then be estimated as $\hat{\tau}(\mathbf{m}) = f_{\hat{Y}_1}(\mathbf{m}) - f_{\hat{Y}_0}(\mathbf{m})$. Shalit et al. (2017) found that learning a shared representation Φ used to predict both treatment outcomes improved prediction quality and named this approach TARNet.

Given a CATE model, we can partition units by their predicted causal effect, creating a clustering of inputs *post hoc*. Let $f_{\text{Cluster}}(\hat{\tau}(\mathbf{m})) \in \{1, 2, ..., K\}$ denote a cluster labeling function determined by the output of $\hat{\tau}$, which partitions the space of effect sizes and, as a result, the space of images. C may be constructed by uniform binning of $\hat{\tau}$, quantile binning, or, as in our experiments, by k-means clustering. Then, the CATE with respect to the post-hoc clustering labels is

$$\textit{Prediction Cluster CATE:} \quad \tau(c) = \mathbb{E}[Y_i(1) - Y_i(0) \mid f_{\text{Cluster}}(\hat{\tau}(\mathbf{M}_i)) = c],$$

In our experiments, we use the post-hoc clustering of TARNet's $\hat{\tau}(\mathbf{M}_i)$ estimates as an entry point of comparison, as the approach is a standard model for high-dimensional causal estimation.

A drawback of a post-hoc model is that it compounds approximations to arrive at a discrete representation of treatment effect heterogeneity. If two similar images yield very different predictions due to misestimation in either the model for $Y_i(0)$ or for $Y_i(1)$, they are likely to be placed in different clusters. We would prefer to cluster images in a way that smoothly best approximates the Image CATE in a single model. Moreover, it is difficult to quantify the aggregation of uncertainty in post-hoc clustering because it may be computationally prohibitive to propagate gradients through both the outcome and subsequent clustering model—gradients that are useful in understanding which parts of the image contribute to the heterogeneity (a topic explored in §3.2).

3.1. Probabilistic Effect Clustering Based on Image Type

We now introduce a set of model strategies for directly targeting CATE clusters in images. There are many possible approaches to this task; here, we seek to develop a modeling strategy for summarizing CATE that is driven by image patterns that can be succinctly summarized for the purposes of interpretability. To the best of our knowledge, this is the first work to use images to estimate CATE.

We seek to fulfill the following criteria: (1) *model potential outcomes and treatment effects flexibly (e.g., allowing for non-linearities)* (2) *identify interpretable image clusters with similar in-cluster effect sizes, and different cross-cluster effects*, and (3) *allow for the modeling of uncertainty regarding the image clusters and treatment effects*.

Our target quantity of interest will be

$$\textit{Image-Type CATE:} \quad \tau(z) = \mathbb{E}[Y_i(1) - Y_i(0) \mid Z_i = z],$$

where $Z_i \in \{1, 2, ..., K\}$ denotes the effect cluster of image $\mathbf{M}_i$, where there are K total clusters. We will search for assignments Z_i of clusters to images that best explain treatment effect variation. A somewhat similar existing analog to this quantity is the mixture-of-experts approach for CATEs in the conjoint setting using linear models with interactions (Goplerud et al., 2022).

Because the treatment effects are decomposed by image type, there are only K quantities needed to effectively summarize the heterogeneity attributable to images. Since human working memory can track between 3 to 5 distinct chunks at a time (Cowan, 2010), this probabilistic summary of the complex heterogeneity process stemming from images can in principle be communicated to human stakeholders, as long as the number of image-effect types is reasonably small. This possibility of specifying the number of clusters in our model is important because it may facilitate discussions regarding fairness in future treatment-targeting decisions. With this substantive motivation in mind, we now discuss how we fulfill the modeling criteria for capturing treatment effect heterogeneity with images. We also note that all our modeling arms will have a probabilistic component in order to fulfill the third criterion of allowing uncertainty estimation.

Probabilistic Estimation of the Baseline Outcome   We satisfy the first criterion by allowing the baseline potential outcome, $Y_i(0)$, to be estimated flexibly as a function of the image. In particular, we let the mean of the baseline potential outcome, conditional on the image, be parameterized by a Bayesian convolutional neural network,

$$\mathbb{E}[Y_i(0) \mid \mathbf{M}_i = \mathbf{m}] = \{\mu_{Y_i(0)} \mid \mathbf{M}_i = \mathbf{m}\} \sim \text{Bayesian CNN}(\mathbf{m}), \tag{1}$$

where convolutional and dense parameters are not deterministic but instead defined according to a distribution. This model for the baseline mean is listed as (2) in Schema 1.

(1a) Effect Mixture Based on Image Type   Having defined the baseline, we next turn our attention to the modeling component that targets the Image-Type CATE estimand. In this component, we first compute image type probabilities $\mathbf{P}_i$ using another Bayesian CNN. This arm looks for image features that predict the mean treatment effect associated with the image. This mean treatment effect model can take on two forms—one more interpretable and the other more flexible.

(1b.i) Image-Type Effect Cluster Model   In the first, conditional on the *image type $Z = z$*, mean treatment effects are drawn from a Normal with a mean $\mu_{\tau,z}$ and variance $\sigma_{\tau,z}^2$ indexed to that image type. We do not assume that there is a single treatment effect per image type, but instead that there is a specific *distribution* over treatment effects by image type. The cluster effect means and variances offer a complete summarization of this distribution.

(1b.ii) Image-Type Differential Effect Model   The first effect mixture model could be useful to investigators because conditional effects can be succinctly summarized with one or two parameters, but in some circumstances, investigators may want a more flexible model for the heterogeneity structure. In that case, the distribution of treatment effects given the image type z is parameterized by a Bayesian CNN arm indexed to z:

$$\{\mu_{\tau_i} \mid \mathbf{M}_i = \mathbf{m}, \ Z_i = z\} \sim \text{Bayesian CNN}_z(\mathbf{m})$$

Here, the image type acts as a stochastic gate that determines which image patterns will be used in predicting the treatment effect means.

Overall, the probabilistic generative model for image-based CATE is defined as follows:

Schema 1: The Image Effect Cluster Model. See Figure A.1 for visualization.

(1a) *Generating the Image Type*

$$\{\mathbf{P}_i \mid \mathbf{M}_i = \mathbf{m}\} \sim \text{Bayesian CNN}(\mathbf{m})$$
$$\{Z_i \mid \mathbf{P}_i = \mathbf{p}\} \sim \text{Categorical}(\mathbf{p})$$
$$\downarrow$$

(1b) *Generating the Treatment Effect Distributions*

i. Image-Type Effect Cluster Model: $\{\mu_{\tau_i} \mid Z_i = z\} \sim \mathcal{N}(\mu_{\tau,z}, \ \sigma_{\tau,z}^2)$

ii. Image-Type Differential Effect Model: $\{\mu_{\tau_i} \mid \mathbf{M}_i = \mathbf{m}, \ Z_i = z\} \sim \text{Bayesian CNN}_z(\mathbf{m})$
$$\downarrow$$

(2) *Generating the Baseline Outcome, $Y_i(0)$*   (3) *Generating the Outcome under Treatment, $Y_i(1)$*

$\{\mu_{Y_i(0)} \mid \mathbf{M}_i = \mathbf{m}\} \sim \text{Bayesian CNN}(\mathbf{m})$ $\rightarrow$ $\{Y_i(1) \mid \mu_{Y_i(0)}, \mu_{\tau_i}, \sigma_{1,z}^2\} \sim \mathcal{N}(\mu_{Y_i(0)} + \mu_{\tau_i}, \ \sigma_{1,z}^2)$

$\{Y_i(0) \mid \mu_i(0), \sigma_{0,z}^2\} \sim \mathcal{N}(\mu_{Y_i(0)}, \ \sigma_{0,z}^2)$

By the structure of the model outlined in Figure A.1, we have conditional mean and variance (see A.1.1.2),

$$\tau(z) = \mathbb{E}[Y_i(1) - Y_i(0) \mid Z_i = z] = \mu_{\tau,z}, \quad \text{Var}(Y_i(1) - Y_i(0) \mid Z_i = z) = \sigma_{0,z}^2 + \sigma_{1,z}^2 + \sigma_{\tau,z}^2.$$

Besides the improved interpretability from summarizing image-derived effect heterogeneity in K discrete clusters, there are other advantages to our modeling strategy. First, as it is probabilistic, we can explore uncertainty not only in the image cluster effects by also in the cluster assignment probabilities. Second, the cluster decomposition also may facilitate scientific inquiry: an image type serves as a generalization tool for reasoning across images, facilitating theorizing about the causal mechanisms at play. Third, we can readily compute the gradients of the expected cluster probabilities with respect to the image in order to identify *how* the image affects the typology, a matter we explore in §3.2.[1]

Using the model outlined in Figure A.1, we attempt learn the joint distribution of the model parameters $\boldsymbol{\Theta}$ and the image clustering, $\mathbf{Z}$, given the observed dataset, $\mathbf{D} = \{Y_i(T_i),\ \mathbf{M}_i,\ T_i\}_{i=1}^n$:

$$\textit{Target Posterior:} \quad p(\mathbf{Z},\ \boldsymbol{\Theta} \mid \mathbf{D}), \tag{2}$$

where $\mathbf{Z} = \{Z_i\}_{i=1}^n$. For details of how we model the uncertainties, see A.1.1. To estimate the posterior in (2), we maximize the Evidence Lower Bound (ELBO) (Ranganath et al., 2014),

$$\underset{q(\mathbf{Z},\ \boldsymbol{\Theta})}{\text{maximize}} \quad \mathbb{E}_{q(\mathbf{Z},\ \boldsymbol{\Theta})}\left[\log\left(p(\mathbf{Y}(\mathbf{T}) \mid \mathbf{Z},\ \boldsymbol{\Theta},\ \mathbf{M},\ \mathbf{T})\right)\right] \ - \ D_{\text{KL}}\left(q(\mathbf{Z},\ \boldsymbol{\Theta}) \ || \ p(\mathbf{Z},\ \boldsymbol{\Theta})\right).$$

We solve the problem approximately using stochastic gradient descent with gradients passing through discrete sampling nodes using re-parameterization techniques (Parmas and Sugiyama, 2021). An additional benefit of the approach proposed here, as opposed to post-hoc clustering, is that uncertainty of the variational clustering model can be further quantified under model misspecification using M-estimation theory (Westling and McCormick, 2019). The choice of priors affects finite sample performance; when possible, we specify priors using observable marginal information (e.g., prior means for cluster effects are set to the differences-in-means ATE estimator value).

3.2. Determination of Salience Regions in Posterior Mean Probabilities

One benefit of the proposed probabilistic image-type heterogeneity decomposition is that we can examine the model in order to assess how image information translates into the predicted effect cluster. In particular, we can take the derivative of the posterior mean cluster probabilities with respect to pixel (w, h):

$$s_{whk}^{\text{Direction}} = \sum_{c=1}^{C} \frac{\partial\, \mathbb{E}\left[\Pr(Z_i = k \mid \mathbf{M}_i = \mathbf{m})\right]}{\partial\, m_{whc}},$$

where $c \in \{1, ..., C\}$ denotes the channel (band) dimension. The quantity, s_{whk} is a scalar summary of how changing pixel (w, h) would induce changes in the predicted effect cluster probability. Because the modeling strategy is probabilistic, the salience must average over the randomness in the predicted cluster probabilities, hence the expectation on the inside of the derivative. This expectation is approximated via Monte Carlo. Positive values of this quantity would indicate that increasing the pixel intensity at (w, h) would increase the probability of cluster k; with the same logic, negative values indicate that increasing pixel intensity at (w, h) would decrease the probability of cluster k.

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1. The approach outlined here can be readily adapted to outcomes having non-Normally distributed outcomes by altering the observed data likelihoods.

Because directional salience may not be interpretable when increasing the pixel intensity is not itself interpretable, we can also consider an approach based on magnitudes, sidestepping the potentially complex interpretation of pixel intensity in different bands:

$$s_{whk}^{\text{Magnitude}} = \sqrt{\sum_{c=1}^{C} \left(\frac{\partial \, \mathbb{E}\left[\Pr(Z_i = k \mid \mathbf{M}_i = \mathbf{m}) \right]}{\partial \, m_{whc}} \right)^2}.$$

Large values of $s_{whk}^{\text{Magnitude}}$ reveal locations in the image that, if changed, would lead to large changes in cluster k probabilities. This measure is agnostic about whether those changes would be associated with increases or decreases in those expected probabilities. This salience information is difficult to compute for post-hoc clustering methods, as the clustering of the $\hat{\tau}_i$'s needs to solve a second optimization problem (as in k-means) through which gradients with respect to the initial outcome model(s) may not be efficiently traced. Moreover, in contrast to the post-hoc approach, the salience measure here incorporates uncertainty in the cluster prediction itself, since the salience maps are generated after integrating over the uncertainty in the probabilities.

3.3. Policy Action Using the Image-based Heterogeneity Model

A major motivation for considering satellite-image-based CATEs is that we can readily generate predictive distributions over treatment effects for contexts outside the experimental setting and where no tabular covariates were measured by researchers—a possibility that may meaningfully expand the reach of causal analyses. In this context, for a new out-of-sample point not from the original dataset, $i^{\text{Out}} \in \mathcal{I}(\text{Out})$, we form a predictive distribution over image treatment effects using

$$p(\tau_{i^{\text{Out}}} \mid \mathbf{M}_{i^{\text{Out}}}, \mathbf{D}) = \sum_{z=1}^{K} \int p(\tau_i \mid Z_{i^{\text{Out}}} = z; \, \boldsymbol{\Theta} = \boldsymbol{\theta}) \cdot p(Z_{i^{\text{Out}}} = z \mid \mathbf{M}_{i^{\text{Out}}}; \, \boldsymbol{\Theta} = \boldsymbol{\theta}) \cdot p(\boldsymbol{\Theta} = \boldsymbol{\theta} \mid \mathbf{D}) \, \mathrm{d}\boldsymbol{\theta}$$

We can use the predictive distribution over treatment effects to improve treatment targeting for out-of-sample individuals. With a fixed treatment budget of size n_1^O for the new dataset of size $n^O = |\mathcal{I}(\text{Out})|$, this policy can be written as $\boldsymbol{\Pi}(\{\mathbf{M}_{i^{\text{Out}}}\}_{\mathcal{I}(\text{Out})}) \rightarrow \{0, 1\}^{n^O}$. There are many approaches to this problem, and we refer readers to the relevant literature (Hitsch and Misra, 2018).

3.4. Image and Tabular Conditional Information

3.4.1. ANALYZING BOTH IMAGE AND TABULAR HETEROGENEITY

Tabular information can be readily incorporated into this modeling pipeline. For example, tabular covariates can be appended to the input of the dense part of the cluster type and baseline outcome models. The resulting treatment effect heterogeneity clusters are therefore conditional on both individual-level and also image-context-level information. This image and tabular data integration can be useful when investigators are focused on understanding the holistic heterogeneity dynamic in an experimental context, integrating both individual- and neighborhood-level information.

3.4.2. DISTINGUISHING IMAGE FROM TABULAR HETEROGENEITY

In some contexts—like when tabular covariates are not measured or when we aim to generate predictions for observations having no measured tabular covariates—it may be useful to perform

image-type effect clustering directly using images alone. However, when other tabular covariates are measured for the experimental sample, researchers may seek to understand the heterogeneity stemming from image information that is not redundant when compared with tabular covariates.

For example, information about economic class is embedded in remote sensing image information about neighborhoods in Africa and beyond, where the urban poor are often concentrated in dense city centers (with little green space) while the affluent often live in green-space-rich areas on the outskirts of cities (Venter et al., 2020). We might wonder about the remaining heterogeneity after accounting for the heterogeneity due to key tabular variables such as race. We can therefore perform the image clustering on the orthogonalized outcomes: $Y_i(t)^\perp = Y_i(t) - \widehat{\mathbb{E}}[Y_i(t)|\mathbf{X}_i]$. The resulting clusters can then be interpreted as image effect types after accounting for the additive heterogeneity from measured tabular data. This image-specific decomposition can be useful when geographic or neighborhood information is the focus of study.

4. Treatment Effect Cluster Recovery in Simulation

Design We now explore the dynamics of the proposed methods in simulation, where true treatment effects are known. We generate image-based treatment effect heterogeneity using,

$$H_i = \text{GN}(\max(f_l(\mathbf{M}_i))), \qquad H_i^+ = \underbrace{\lfloor \min\{H_i\}_{i=1}^n \rfloor}_{\text{Ensures } \tau_i > 0} + \underbrace{\text{sign}(H_i) \cdot |H_i|^{1/\gamma}}_{\text{Generates bimodality as } \gamma \to \infty} \qquad (3)$$

where $f_l(\cdot)$ denotes the application of a $l \times l$ filter to the image, $\max(\cdot)$ denotes the global maximum operation across the image, and $\text{GN}(\cdot)$ denotes a global normalization function scaling the H_i values to have mean 0 and variance 1 across the image pool. The specific transformation generating H_i^+ is selected (1) to ensure all the treatment effects are in the same direction (i.e., all positive) and (2) to generate heterogeneity in the effect distribution, with greater bimodality in the treatment response as $\gamma \to \infty$. We let $\gamma = 2$. We define the treatment and outcome:

$$T_i \sim \text{Binomial}(0.5), \quad Y_i = T_i H_i^+ + \epsilon_i,$$

with $\epsilon_i \sim \mathcal{N}(0, \nu \cdot \text{Var}(H_i^+))$. The value of ν controls the extent to which the image is predictive of the outcomes, where smaller values indicate a stronger image heterogeneity signal. To explore the effect of the signal-to-noise ratio, we vary $\nu \in \{0.01, 0.1, 1\}$.

The filter used in the convolution function of Equation 3 is visualized in Figure A.2, along with high and low responders from the set of images used in the simulation that we take from Landsat mosaics of sub-Saharan Africa. We have some degree of model misspecification in our estimation model, as the way the data is generated is distinct from the estimation model; given this misspecification, we will examine the degree to which our model will recover key properties of the image-based causal heterogeneity.

Cluster Recovery Measure We compare the estimated effect clustering with an oracle baseline from the true, in practice unknown, τ_i's. That is, we first compute the oracle k-means clustering:

$$\tau(z)_{\text{Oracle}} = z^{\text{th}} \text{ center from the oracle } k\text{-means applied to the true (in practice unknown) } \tau_i\text{'s}$$

The clustering quality measure then compares the oracle with estimated cluster means:

$$\text{Cluster Recovery } R^2 \;\; = \;\; 1 - \frac{\sum_{z=1}^{K} \min_{z'} (\widehat{\tau}(z) - \tau(z')_{\text{Oracle}})^2}{\sum_{z''=1}^{K} (\tau(z'')_{\text{Oracle}} - \overline{\tau}_{\text{Oracle}})^2},$$

where $\overline{\tau}_{\text{Oracle}}$ denotes the mean across the oracle cluster centers. This measure is equivalent to the R^2 in predicting the oracle cluster means from the estimated ones, where the ordering of the clusters has been arranged so that each oracle center is compared to its nearest estimated counterpart.

Simulation Results We see in the left panel of Figure 1 one representative posterior distribution over $Y_i(1) - Y_i(0)$ given estimated cluster information. We find that the estimated clusters capture the bimodality present in the true distribution of τ_i's. The right panel shows how the cluster recovery measure for the Image-Type Differential Effects Model and TARNet post-hoc clustering are similar in the low residual variance setting. In the high residual variance setting, the TARNet clustering struggles somewhat in recovering the oracle cluster centers. In contrast, the parsimonious Image-Type Effect Clustering Model performs best at recovering the clustering of the treatment effects across the noise range.

Figure 1: *Left:* Capturing the treatment effect heterogeneity with our Image-Type Effect Model. *Right:* Comparing models as we vary the signal-to-noise ratio.

5. Application to an Anti-Poverty Experiment in Uganda

Data In our application, we explore the effects of the anti-poverty experiment performed in Uganda and described in §1. The treatment variable is the random assignment of small teams to receive grants for business ventures. The outcome variable is an aggregate summary of skilled labor (see A.1.3.1) measured at the end of the experiment (two years after treatment assignment). De-identified outcome and treatment data were given voluntarily by subjects and are available under CC0 1.0 license. Longitude/latitude information about respondents' villages is found using OpenStreetMap.

Pre-treatment image data is taken from Landsat. We use the Orthorectified ETM+ pan-sharpened data product, processed to contain minimal cloud cover. Reflectance is measured in the green, near-infrared, and short-wave infrared bands. These bands are useful in capturing information about peak vegetation, water content, and thermal dynamics, in addition to structural land features. In contrast to RBG-only images' resolution of 30 meters, these ETM+ mosaics have resolution of 14.25 meters.

Empirical Results Due to space constraints, we focus only on results from the Image-Type Effect Cluster Model (details in §A.1.5). We set the cluster number to 2 after finding that cluster probabilities become highly correlated with additional clusters. The top three rows in the left panel of Figure 2 show results for the highest images having the highest posterior mean probabilities for cluster 1. For each image, this figure visualizes the salience measures defined in §3.2. The bottom three rows show results for the highest posterior mean cluster 2 images. The treatment effect for

cluster 1 is substantially different than for cluster 2. Visually, we see that smaller treatment effects exist for places with harsher terrain and less developed transportation networks, hampering economic growth. These low responders are found in the harsh mountainous northern part of Uganda. This is logical, as skilled labor tends to thrive in areas that are not isolated geographically (Ashraf and Galor, 2011).

We show in the right panel of Figure 2 how the results of the experiment may be generalized to the entire country of Uganda, assuming no systematic bias in places chosen conditional on the image information. In particular, we show the posterior predictive mean cluster 2 probability for the entire country. This kind of analysis can provide policymakers with potentially useful information for how to improve the targeting of treatments in the future across larger geographic contexts.

The Appendix contains supplementary analyses for this application. For example, we show in Figure A.5 the correlation between the estimated Image CATEs and Tabular CATEs using various conditioning sets, as well as, in Table A.1, between the cluster probabilities and other individual-level covariates. Moreover, we show in Figure A.4 that the places having the most uncertainty in the cluster probabilities (estimated by the posterior standard deviation). In addition, we orthogonalize the potential outcomes using tabular information (see A.1.4), and the results remain similar: the correlation between raw and non-orthogonalized cluster probabilities is 0.85. Because our results remain similar after orthogonalization, the Ugandan satellite images supply independent CATE information that the original experiments could not assess.

6. Discussion and Conclusion

Applied researchers and policymakers use RCTs to estimate population-wide effects (ATE) and sub-population effects (CATE), using tabular data collected at baseline, often near-time to when the RCT is launched. However, these near-time variables tend to miss historical characteristics of the experimental units, such as their neighborhood and geographical characteristics, that are important for CATE. While such historical data are often expensive to collect or unavailable, satellite images are a data stream that captures such characteristics in an unstructured form. As no CATE method exists explicitly for image analysis, this paper presents principles and modeling strategies for analyzing image-based CATE using probabilistic image-type models. We derive some properties of this model and perform approximate inference using variational methods. Dynamics are explored via simulation, and an anti-poverty field experiment from Uganda was analyzed, where we seem to find interesting heterogeneity dynamics.

Our approach has other limitations, which form an incentive for future research. First, our models estimate heterogeneity clusters at the image level, but not explicitly for smaller segments of an image. Having such within-image heterogeneity segmentation would further improve interpretability of what exactly in the image is generating heterogeneity. Second, our methods estimate heterogeneity with respect to a fixed baseline (i.e. the control intervention). While the choice of baseline is clear in most settings in RCTs, in others, it may not be. In unclear cases, investigators may need to run the models using different baselines and compare results. Third, our model is tailored for RCTs (i.e., assuming unconfoundedness); more research is required to adapt it for observational settings. Using experimental data, effect estimates are confounding-free by design; heterogeneity can be studied independently of identification. Observational data are more plentiful but require adjustment (Rosenbaum et al., 2010).

Figure 2: *Left, top 3 rows:* High probability cluster 1 images. *Left, bottom 3 rows:* High probability cluster 2 images. "Salience Magnitude" and "Direction" are defined in §3.2. *Right, top:* A land use map of Uganda. *Right, center:* Posterior predictive mean cluster 2 probabilities for the entire country. Black circles represent observed sample points. *Right, bottom.* Posterior standard deviation of the cluster 2 probabilities.

Finally, our focus on *CATE for images* opens exciting possibilities. It not only encourages others to start incorporating satellite images in their planned experiments but also reanalyze past experiments, potentially unraveling previously undetected yet significant CATE. As demonstrated in our analysis of the Ugandan anti-poverty experiment, our method identifies heterogeneity not initially detected by incorporating informative satellite data. Thus, our image-based methods have the potential to contribute to policy by complementing traditional RCT heterogeneity analysis based on tabular $\mathbf{X}_i$—not to mention analyses in other fields such as agriculture, disaster relief, climate science, and medicine where image data is also prevalent. □

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