Degree of Interference: A General Framework For Causal Inference Under Unknown Interference

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Abstract

One core assumption typically adopted for valid causal inference is that of no interference between experimental units, i.e., the outcome of an experimental unit is unaffected by the treatments assigned to other experimental units. This assumption can be violated in real-life experiments, which significantly complicates the task of causal inference as one must disentangle direct treatment effects from “spillover” effects. Current methodologies are lacking, as they cannot handle arbitrary, unknown interference structures to permit inference on causal estimands. We present a general framework to address the limitations of existing approaches. Our framework is based on the new concept of the “degree of interference” (DoI). The DoI is a unit-level latent variable that captures the latent structure of interference. We also develop a data augmentation algorithm that adopts a blocked Gibbs sampler and Bayesian nonparametric methodology to perform inferences on the estimands under our framework. We illustrate the DoI concept and properties of our Bayesian methodology via extensive simulation studies and an analysis of a randomized experiment investigating the impact of a cash transfer program for which interference is a critical concern. Ultimately, our framework enables us to infer causal effects without strong structural assumptions on interference.

Keywords: Rubin Causal Model, Interference, Spillover effect, Stable Unit Treatment Value Assumption, Bayesian Nonparametric, Dependent Dirichlet Process mixture

1 Introduction

The two components of the Stable Unit Treatment Value Assumption (SUTVA) are fundamental for causal inference for a treatment effect. The first component of SUTVA is that...
there are well-defined treatments for each experimental unit that yield well-defined outcomes, and the second is that no interference exists between units. The first assumption is typically satisfied by the design of an experiment, whereas the second relates to the complicated phenomenon of interference and may not be true in certain experiments. Interference is said to arise when a particular experimental unit’s outcome is a function of both their assigned treatment as well as the treatments assigned to other experimental units. For example, interference can naturally arise in an experiment on the efficacy of a vaccine for a contagious disease, as one subject’s health outcome can depend on whether their close contacts received the vaccine. The existence of interference complicates causal inferences because one must disentangle direct treatment effects from the spillover effects of the treatments assigned to other units.

This paper proposes a Bayesian semiparametric methodology for causal inference under unknown interference. The existing methods for causal inference under interference, as we discuss in §2, either assume the interference structure or assess the robustness of inference when it is misspecified. In the latter case, the inference is generally sensitive to misspecification. In our proposed approach, a unit’s interference structure is modeled as a latent function of the unit’s features and other units’ treatments. This latent function identifies the direct and spillover effect. We develop a Bayesian nonparametric prior for this latent function and a fast data augmentation algorithm that adopts a blocked Gibbs sampler to draw posterior inferences for the causal effects. The code implementing our method is available in the online supplement.

We call the unit-level latent functions “degree of interference.” One of our main contributions is to formalize the notion of the degree of interference. Using this notion, we develop a flexible modeling strategy that avoids strong assumptions on the interference structure. In the next section, we review existing methods and highlight our contributions in their context.

2 Current Literature on Causal Inference Under Interference

The first foundational work towards causal inference under interference was by Hudgens and Halloran (2008), who proposed a design-based approach to identify and estimate direct and spillover effects in a two-stage hierarchical experimental design. The first stage of the design randomly assigns clusters of units to one of two treatment assignment strategies, and then the second stage assigns treatments within the clusters based on the selected strategies. Their methods assume that the interference is (1) partial, i.e., operates only within clusters, and (2) stratified, i.e., is specified via the proportion of treated units within clusters. Several important works have built upon the above framework (Liu and Hudgens, 2014; Kang and Imbens, 2016; Basse and Feller, 2018; Imai et al., 2021; Ohnishi and Sabbaghi, 2022). However, in practice, the validity of the assumptions of partial interference, which requires multiple independent clusters, and stratified interference, which is typically a simplification for the unknown interference structure, may be questionable (Hudgens and Halloran, 2008, p. 836). When these assumptions fail, the above methodologies provide biased inferences. In contrast, we utilize a Bayesian nonparametric prior for the “degree of interference”, specifically, the Dependent Dirichlet process mixture prior to flexibly model the unknown interference structure. These priors can data-adaptively separate units into several clusters based on how they interfere with one another, thus accommodating a stratified nature of the
Aronow and Samii (2017) introduced the idea of exposure mapping to move beyond the partial and stratified interference assumptions. Under this framework, a mapping is specified that relates the vector of treatment assignments for the other experimental units to a finite set of exposures. Then, an experimental unit’s outcome is specified as a function of its own treatment label and this exposure. In other words, an exposure map is supposed to capture the full nature of interference of a unit from all other units. Once an exposure map is well-specified, causal effects are defined in terms of comparisons of outcomes under different exposures. A line of research exists that uses exposure maps to provide causal inference under interference. Notably, Toulis and Kao (2013) developed randomization-based and Bayesian model-based inferential approaches for “peer influence effects” defined by fixing the specific number of neighbors who receive treatment. Basse et al. (2019) developed a framework for constructing valid randomization tests for general exposure contrasts. Other research works have addressed interference by specifying their interference mapping based on peers’ treatment status and attributes of a given network (Li et al., 2019; Zigler and Papadogeorgou, 2021; Forastiere et al., 2021; Papadogeorgou and Samanta, 2023).

As Basse and Airoldi (2018) mention, causal inference under interference is impossible without assumptions about the interference structure. Thus, it is natural to use exposure maps specifying the structure. However, current causal inferential frameworks and methodologies based on exposure maps are limited in several respects. First, they require that the interference structure be fully characterized as a finite set of exposures \textit{a priori} so that each exposure corresponds to a unique potential outcome. In practice, satisfying this requirement may be unrealistic because the interference structure is usually unknown and can be very complicated to understand. Second, as observed by Sävje (2023), a misspecified exposure mapping can lead to imprecise variance estimation and, thus, incorrect inference. Partial knowledge of the interference structure can mitigate this problem. Leung (2021) proposed a flexible model that permits treatments that are assigned to units “farther” from a particular unit to have non-vanishing, but smaller, effects on the unit’s response. However, this is not very helpful when we are unaware of this structure, which is often the case in practice. Our proposal of the “degree of interference” generalizes the exposure map but does not require precise knowledge of the interference structure; rather learns a flexible model for the interference given the data using a fast Gibbs sampler.

Third, the exposure mapping framework simultaneously defines both the interference structure that determines a single unique potential outcome and exposure effects that determine causal quantities. Since exposure mapping is typically problem specific, exposure effects generally do not hold any meaning outside of the problem instance, even when for the same treatment-outcome pair. In contrast, we define new causal estimands that are not bonded with the interference structure. Our estimands are generalizations of those suggested by Sävje et al. (2021) as natural causal estimands for direct and spillover effects under interference. Finally, our approach allows for continuous treatment factors and factors with many discrete levels, while most existing frameworks lead to overly complicated formulations of causal inference under interference under these general treatment factors.

The rest of the paper is organized as follows. We proceed in §3 to review the Rubin Causal Model and propose new spillover effect estimands under interference. §4 contains our new framework based on the “degree of interference” concept and our corresponding data augmentation algorithm for inferring causal estimands under interference. The frequentist properties of our
method are demonstrated via extensive simulation studies in §5. In §6 we demonstrate our framework using a case study on cash transfer programs in Colombia. §7 concludes the paper.

3 Background

3.1 The Rubin Causal Model and Interference Structures

Adopting the Rubin Causal Model (Rubin, 1974), consider $N$ experimental units, indexed by $i = 1, \ldots, N$, that correspond to physical objects at a particular point in time. Each experimental unit $i$ has an observed set of pre-treatment covariates $X_i \in \mathcal{X}$, with $\mathcal{X}$ denoting the covariate space; typically; $\mathcal{X} = \mathbb{R}^d$ for $d$ covariates. Treatment assignments are contained in the $N$-dimensional vector $Z = (Z_1, \ldots, Z_N)^T$, where $Z_i \in \Omega$, the treatment assignment for unit $i$, can be binary, categorial or continuous, and $\Omega$ is the set of treatment levels. So, the vector $Z \in \Omega^N$. For example, $\Omega = \mathbb{R}$ denotes a continuous treatment factor with “0” denoting the specified “control” level, and $\Omega = \{0, 1, \ldots, K\}$ denotes a categorical treatment with $K + 1$ levels and with “0” again denoting control. Let $Z_{-i} = (Z_1, \ldots, Z_{i-1}, Z_{i+1}, \ldots, Z_N)^T$ be the sub-vector of $Z$ with the $i$th element removed; so, $Z_{-i} \in \Omega^{N-1}$.

Next, consider the potential outcomes for unit $i$. To facilitate the exposition and indicate how, in the general setting of interference, unit $i$’s potential outcomes depend on multiple entries in $Z$, we write the potential outcomes for unit $i$ as $Y_i(Z) \equiv Y_i(Z_i, Z_{-i})$. We let $Y = \{Y_i(z) : i = 1, \ldots, N, z \in \Omega^N\}$ denote the set of all potential outcomes. The observed outcomes are contained in vector $Y_{\text{obs}} = (Y_{\text{obs}}^1, \ldots, Y_{\text{obs}}^N)^T$ where $Y_{\text{obs}}^i = \sum_{z \in \Omega^N} Y_i(z)I(Z = z)$ for each $i$. Thus, only one outcome is observable to us among the many potential outcomes of unit $i$.

The probability distribution of $Z$ given covariates $X_1, \ldots, X_N$ is known to us. We assume without loss of generality that the support of the probability distribution is equal to $\Omega^N$.

Since the experimental units interfere, as is common in the literature, we assume throughout that they belong to a single, known network that governs their interference structure. Let $A$ denote the $N \times N$ matrix that captures how the units are related to one another. The specification of $A$ can vary depending on how one wishes to describe interference between units. For example, to describe the interference structure based just on unit adjacencies, $A$ can be defined simply via the adjacency matrix of the units. If instead one wishes to use the shortest spatiotemporal path between units to describe their interference structure, then $A$ is defined as the distance matrix of the units, with element $(i, j)$ of $A$ being the shortest distance between units $i$ and $j$. Later, in our definition of the degree of interference, $A$ becomes an argument of its functional form.

3.2 Causal Estimands

Throughout, we consider the finite-population perspective, where we view the $N$ units are given, and an external, known randomization process assigns treatment to these units. All potential outcomes are defined as fixed but mostly missing for the unrealized treatment assignment.

We define two types of causal estimands—the first type is assignment-conditional effects, and the second is expected effects marginalized over a treatment assignment. Assignment-conditional causal estimands are defined by conditioning on a specific treatment assignment vector. We consider two effects for this type of estimand: average treatment and spillover effects.
Following Sävje et al. (2021), who used similar estimands in the case of a binary treatment, the unit-level assignment-conditional treatment effect for unit $i$ is the contrast between its two potential outcomes corresponding to $Z_i = 0$ and $Z_i = z$ when the treatment assignments $Z_{-i}$ for the other experimental units are held fixed at $z_{N-1} \in \Omega^{N-1}$. Denote this effect by $\tau^\tr_i(Z_i = z, Z_{-i} = z_{N-1}) = Y_i(Z_i = z, Z_{-i} = z_{N-1}) - Y_i(Z_i = 0, Z_{-i} = z_{N-1})$. The average of the $\tau^\tr_i(Z_i = z, Z_{-i} = z_{N-1})$ across the $N$ experimental units corresponds to a finite-population average treatment effect.

**Definition 1** (Assignment-Conditional Average Treatment Effect). An assignment-conditional average treatment effect (A-CATE) comparing a pair of treatment, a $z \in \Omega$ and vector $z' \in \Omega^N$ is

$$
\tau_{A-CATE}(z, z') = \frac{1}{N} \sum_{i=1}^{N} \tau^\tr_i(Z_i = z, Z_{-i} = z'_{-i}),
$$

where $z'_{-i}$ is an $(N - 1)$-dimensional sub-vector of $z'$ with element $i$ removed.

A-CATE is a natural extension of the estimand of Sävje et al. (2021) for the cases of categorical or continuous treatments. This effect quantifies the average change in the outcome as one unit’s treatment changes from 0 to $z$ when the treatments for all other units are fixed at $z'_{-i}$.

We define our unit-level spillover effect that is inspired by the A-CATE. Specifically, for unit $i$ with treatment $z$ and for two treatment assignment vectors $z'_{N-1}, z^*_N \in \Omega^{N-1}$ for the other units, the unit-level spillover effect is defined as $\tau^\sp_i(z, z'_{N-1}, z^*_N) = Y_i(Z_i = z, Z_{-i} = z'_{N-1}) - Y_i(Z_i = z, Z_{-i} = z^*_N)$.

**Definition 2.** An assignment-conditional average spillover effect (A-CASE) under a treatment $z \in \Omega$ for unit $i$ and treatment vectors $z'_N, z^*_N \in \Omega^N$

$$
\tau_{A-CASE}(z, z'_N, z^*_N) = \frac{1}{N} \sum_{i=1}^{N} \tau^\sp_i(z, z'_i, z^*_i),
$$

where $z'_i$ and $z^*_i$ are $(N - 1)$-dimensional sub-vectors of $z'$ and $z^*$ with the $i$th elements removed.

Our definition of the A-CASE corresponds to the average comparison of the units’ outcomes for a single treatment under two different cases, corresponding to the two different treatment vectors considered for the other units. Of special interest is the case with $z = 0, z^*_N = (0, \ldots, 0)^T$, and $z'$ set at some other vector, where A-CASE captures a pure spillover effect.

Our second type of causal estimands is a set of three expected effects, which marginalize the assignment-conditional causal estimands over a probability distribution on the assignment vectors, i.e., the treatment assignment mechanism. Expected effects are useful in practice because the number of assignment-conditional effects grows rapidly as a function of $N$, whereas the number of expected effects is generally smaller.

**Definition 3** (Expected Average Treatment and Spillover). Let $\mathbb{E}_\pi(h(Z))$ denote the expectation of a function $h(Z)$ of the assignment vector over the known probability mass/density function $\pi(\mathbf{z}) = p(\mathbf{z} \mid \mathbf{X}_1, \ldots, \mathbf{X}_N)$. Then for each assignment-conditional average effect, we define the corresponding expected average effect based on the expectation of the assignment-conditional average effect over the assignment mechanism: $\tau_{E-ATE}(z; \pi) = \mathbb{E}_\pi\{\tau_{A-CATE}(z, Z)\}$ and $\tau_{E-ASE}(z; \pi) = \mathbb{E}_\pi\{\tau_{A-CASE}(z, Z, 0)\}$.
Our version of this estimand, $\tau_{E-ATE}$, is a natural extension of the estimand of Sävje et al. (2021) for categorical treatment factors with multiple levels and continuous factors. Our definitions of $\tau_{E-ASE}$ is motivated in a similar manner as that of $\tau_{E-ATE}$. As noted by Sävje et al. (2021), the E-ATE—similarly, E-ASE—captures the expected average effect of changing the treatment of a single unit in the “current” experiment.

4 The Degree of Interference Framework and Bayesian Methodology

4.1 Definition and Assumptions for the Degree of Interference

The Degree of Interference (DoI) is defined in terms of the characteristics of a unit and the treatments for all other units. Our DoI framework addresses a limitation of exposure maps that, while an exposure map needs to be pre-specified, the DoI is an unknown latent function that represents the underlying interference structure between units, which is inferred from the data.

We let characteristics $T_i \in \mathcal{T}$ for unit $i$ contain, its covariates, $X_i$, covariates for units other than $i$, $X_{-i}$, and the adjacent network information of the other units. In all that follows, we let $\mathcal{T} = \mathcal{X} \times \mathcal{X}^{N-1} \times \mathcal{A}$, where $\mathcal{A}$ denotes the set of matrices of networks, i.e., set of $\mathbf{A}$s.

Definition 4. The DoI for unit $i$ is a random function $G_i : \mathcal{T} \times \Omega^{N-1} \to \mathbb{R}$ that maps the unit’s characteristics, residing in $\mathcal{T}$, and treatments for all other units to a number.

The DoI satisfies the following consistency assumption which says that unit $i$’s potential outcomes are identified from its own treatment and DoI.

Assumption 1. (Consistency of DoI) For all $i \in \{1, ..., N\}$, $T_i \in \mathcal{T}$, $z \in \Omega$, and $z_{N-1}, z_{N-1}' \in \Omega^{N-1}$ in which $G_i(T_i, z_{N-1}) \overset{D}{=} G_i(T_i, z_{N-1}')$, where $\overset{D}{=}$ denotes equivalence in the marginal prior distributions of $G_i(T_i, z_{N-1})$ and $G_i(T_i, z_{N-1}')$, we have $Y_i(z, z_{N-1}) = Y_i(z, z_{N-1}')$.

Remark (Randomness of DoI). Assumption 1 is critical, as it implies that the DoI is effectively a latent representation of the interference structure for unit $i$ with characteristics $T_i$ when other units are assigned to $z_{N-1}$. The distributional equivalence allows for additional flexibility of the DoI; it indicates the unknown uncertainty about the interference structure. Under the finite-population perspective, the treatment assignment is usually the only source of randomness for observables. Therefore, the existing design-based approach under the finite-population perspective (e.g., partial interference, stratified interference and exposure maps) inevitably defines the interference structure a priori even when there is limited knowledge about it. Under our framework, by contrast, the interference structure is an “unobserved” quantity, representing uncertainty for analysts. We incorporate the knowledge about interference and the associated uncertainty into our model-based inference developed in the following sections. More specifically, we use a Bayesian approach to infer unknown DoIs, requiring us to specify their prior distributions. From the Bayesian perspective, the randomness of the DoI can be viewed as the representation of prior knowledge and uncertainty about the interference structure. This uncertainty should be distinguished from the ones on the underlying network between units (Egami, 2021). We place priors on functional spaces, which are flexible enough to recover the latent DoIs that satisfy Assumption 1.
This assumption ultimately corresponds to equivalence in the potential outcomes according to DoIs that have the same marginal prior distributions. More formally, for all $i \in \{1, ..., N\}$, $z \in \Omega$, $z_{N-1} \in \Omega^{N-1}$, and $T_i \in \mathcal{T}$, the equivalencies of the potential outcomes for unit $i$ with $Y_i(z, z_{N-1})$ are based on $z$ and the marginal prior distribution on $G_i \equiv G(T_i, z_{N-1})$. We represent its conditional cumulative distribution function $F_{G_i(T_i, z_{N-1})}$. Hence, as a consequence of Assumption 1, we consider “auxiliary” potential outcomes $\tilde{Y}_i(z, g)$, where $\tilde{Y}_i(z, G_i)$ and $\tilde{Y}_i(z, G'_i)$ have same distributions for any $G_i, G'_i$ that have the same marginal prior distributions. In contrast to the original potential outcome, the auxiliary potential outcomes have randomness induced by $G_i$. Thus, we write the original potential outcome using the auxiliary potential outcomes as:

$$ Y_i(z, z_{N-1}) = \mathbb{E}_{G(T_i, z_{N-1})}\{\tilde{Y}_i(z, G_i)\} = \int_{\mathbb{R}} \tilde{Y}_i(z, g) dF_{G_i(T_i, z_{N-1})}(g), \quad (1) $$

Equation (1) says that the uncertainty about $G_i$ is incorporated by marginalizing the auxiliary potential outcomes over $G(T_i, z_{N-1})$ for the original potential outcomes with $Z_{-i} = z_{N-1}$. Also, it implies an inferential approach for $Y_i(z, z_{N-1})$ via the Monte Carlo simulation: $Y_i(z, z_{N-1}) \approx \sum_{m=1}^M \tilde{Y}_i(z, g_m)/M$ where $g_m \sim F_{G_i(T_i, z_{N-1})}$ for $m = 1, \ldots, M$. The Monte Carlo procedure is introduced in §4.4. Finally, the unit-level assignment-conditional treatment effect is

$$ \tau_i^{tr}(Z_i = z, Z_{-i} = z_{N-1}) = \int_{\mathbb{R}} \{\tilde{Y}_i(z, g) - \tilde{Y}_i(0, g)\} dF_{G_i(T_i, z_{N-1})}(g). \quad (2) $$

Unit-level assignment conditional spillover effects can also be expressed in terms of the distribution of the DoI. Equation (2) further clarifies the conceptual role of the DoI that it is effectively a low-dimensional compression of interference, and the direct effect is defined as the comparison of two auxiliary potential outcomes with different treatment assignments under the same DoI.

DoI, along with Assumption 1, generalizes exposure mappings (Aronow and Samii 2017). In particular, if $G_i$ is a prespecified, known function that maps to a finite set then the DoI is analogous to exposure mappings. For example, the exposure mapping for stratified interference (Hudgens and Halloran 2008) would correspond to prespecifying $G_i$ as the number of treated units in the neighborhood of unit $i$. However, in contrast to existing approaches based on exposure maps, the DoI is not prespecified but is instead inferred from data via our Bayesian methodology in §4.2 and §4.3. Inferring the DoI is advantageous in principle, as imposing a misspecified interference structure and failing to modify it could yield biased inferences (Savje 2023).

Additionally, exposure mapping plays a dual role in the existing literature—pre-specified exposure maps define the direct and spillover effect, and they are used to impose assumptions on the interference structure. The DoI framework addresses an open question posed by Savje (2023) regarding estimators that fully separate the two roles of exposure maps. Our inferential approach can incorporate knowledge about the interference structure in estimating an exposure effect without necessarily changing the effect being estimated. This is because, first, our causal estimands are generally defined in terms of treatment assignments and are not dependent on exposures. Second, we incorporate knowledge about the interference structure by specifying priors on the DoI that can capture their complexities and uncertainties and yield valid posterior distributions for the finite-population causal estimands. The Bayesian methodology propagates the uncertainties about interference structures using an imputation-based approach outlined in §4.2.

Finally, the DoI framework allows causal inferences for experiments that satisfy the following new version of the unconfoundedness assumption.
Assumption 2. A treatment assignment is unconfounded under the DoI framework if the probability mass/density function in the treatment assignment mechanism does not depend on \( Y \) or the \( G_i(T_i, z_{-i}) \) conditional on the \( T_i \), i.e.,
\[
p(z|T_1, \ldots, T_N, Y, G_1(T_1, z_{-1}), \ldots, G_N(T_N, z_{-N})) = p(z|T_1, \ldots, T_N), \text{ for all } T_1, \ldots, T_N \in \mathcal{T}, z \in \Omega^N \text{ and } G_1(T_1, z_{-1}), \ldots, G_N(T_N, z_{-N}).
\]

While this assumption is less restrictive than a randomized treatment assignment, this assumption may not be valid in observational studies, e.g., when units receive treatments so as to minimize the DoI. In our case study in Assumption 2 is valid due to the design of the experiment.

4.2 Overview of the Bayesian Methodology

Bayesian inference is one of the modes of inference within the potential outcome framework, where we consider the missing potential outcomes as unobserved random variables, no different from unknown parameters. Specifically, Bayesian inference involves specifying a model for all random variables, including the potential outcomes, treatment, and covariates. Based on this model, one can infer causal estimates from the joint posterior distributions of the parameters and the missing potential outcomes, conditional on the observed data (Rubin 1978).

The Bayesian approach not only provides a principled framework for analyzing causal studies but also provides a refined map of identifiability, clarifying what can be learned when causal estimates are not fully identifiable but instead weakly identifiable (i.e., when the likelihood functions of parameters and causal estimates have substantial regions of flatness) (Imbens and Rubin 1997). In particular, issues of identification differ from those in the frequentist paradigm because if prior distributions are proper, then posterior distributions are always proper. Weak identifiability is reflected in the flatness of the posterior distribution and can be evaluated quantitatively (Gustafson 2009). Li et al. (2022) give further comprehensive reviews of Bayesian causal inference.

Our Bayesian methodology involves multiple imputations of the DoI for missing potential outcomes to derive the posterior distributions of the finite-population causal estimands. To formally describe this, let \( \tau \) denote one of the causal estimands from \( \tau \). We let \( T \) the \( N \times P \) matrix of characteristics for all units (where the columns correspond to the concatenation of covariates and adjacent network information), \( Z \) the \( N \times 1 \) vector of assigned treatments, \( Y^{obs} \) the vector of observed outcomes, and \( Y^{mis} \) the vector of all missing potential outcomes. The \( Y^{mis} \) is determined according to the treatment assignment, with each missing potential outcome for unit \( i \) being either \( Y_i(z'_i, z_{-i}) \) or \( Y_i(z, z'_{-i}) \) with \( z'_i \) denoting an alternate treatment for unit \( i \) and \( z'_{-i} \) denoting an alternate treatment vector for all units excluding unit \( i \). In these cases, the DoI for unit \( i \) would be either \( G_i(T_i, z_{-i}) \) or \( G_i(T_i, z'_{-i}) \) respectively. We let \( G^o = (G_1(T_1, Z_{-1}), \ldots, G_N(T_N, Z_{-N}))^T \) and \( G^u(z') = (G_1(T_1, z'_{-1}), \ldots, G_N(T_N, z'_{-N}))^T \) contain the DoIs for the realized and an unrealized treatment assignment vector \( z' \), respectively. None of the entries in \( G^o \) or the \( G^u(z') \) are observable. In accordance with the matrix \( Y^{mis} \), we generally define the matrix \( G^u \) as the column-wise concatenation of all the \( G^u(z') \) for treatment assignment vectors \( z' \) that were not realized.

Standard types of point estimates of estimand \( \tau \) are obtained via means, medians, or modes of the posterior distribution \( p(\tau | T, Z, Y^{obs}) \), and standard interval estimates are obtained via the central credible intervals or highest posterior density intervals. As causal estimands are functions of observed and missing potential outcomes, we recognize that we can calculate this posterior
distribution by integrating out $Y^{\text{mis}}$, $G^o$, and $G^u$ according to

$$
\int p(\tau \mid T, Z, Y^{\text{obs}}, G^o, G^u, Y^{\text{mis}}) p(G^o, G^u, Y^{\text{mis}} \mid T, Z, Y^{\text{obs}}) \, dG^o \, dG^u \, dY^{\text{mis}}. \tag{3}
$$

In order to sample from this posterior, we first derive the posterior distribution of the DoI and missing outcomes conditional on the observed data. Then, we calculate the posterior distribution of the estimand conditional on the observed data and imputations of the DoI and missing outcomes drawn from their posterior distributions. We perform all imputations and posterior calculations via Markov Chain Monte Carlo technique.

Four inputs are necessary to derive the posterior distributions of the causal estimands. (i) The first is knowledge of the assignment mechanism, as encoded in the probability mass/density function $p(Z \mid T)$. As we consider designed experiments, this probability mass/density function is known. (ii) The second is the prior distribution for the DoIs conditional on $T$ and $Z$, which we denote by the joint probability density function $p(G^o, G^u \mid T, Z, \phi)$. We assume that $G^o$ and $G^u$ are a priori independent of each other given $T, Z$. (iii) The third input is a model for the potential outcomes conditional on $T, Z, G^o, G^u$, which we represent via a joint probability density/mass function $p(Y_{i, obs}^{\text{mis}} \mid T, Z, G^o, G^u, \theta)$ with parameter vector $\theta$. (iv) The final input is the prior distribution for $\phi$ and $\theta$, denoted by $p(\phi, \theta)$. We assume that $\phi$ and $\theta$ are distinct and do not share any parameters. We describe our priors for the DoI, potential outcomes and the parameters $(\phi, \theta)$ in §4.3.

Given those four inputs, we first express the complete data likelihood function under Assumption 2 and the assumption of independence in the statistical model according to

$$
L(\theta, \phi \mid Y^{\text{obs}}_{i, Y^{\text{mis}}}, G^o, G^u, T, Z) = p(Y^{\text{obs}}_{i, Y^{\text{mis}}}, G^o, G^u \mid T, Z, \theta, \phi)
$$

$$
= p(Y^{\text{obs}}_{i, Y^{\text{mis}}}, G^o, G^u \mid T, \theta, \phi) = \prod_{i=1}^{N} p(Y^{\text{obs}}_{i, Y^{\text{mis}}}, G^o_{i, G^u_{i}}, T, \theta) \, p(G^o_{i, G^u_{i}} \mid T, \phi).
$$

The posterior distribution of the model parameters $p(\phi, \theta \mid Y^{\text{obs}}_{i, Z})$ is thus derived as

$$
p(\phi, \theta \mid Y^{\text{obs}}_{i, Z}) \propto p(\phi, \theta) \int \prod_{i=1}^{N} p(Y^{\text{obs}}_{i, Y^{\text{mis}}}_{i}, G^o_{i, G^u_{i}}, T, \theta) \, p(G^o_{i, G^u_{i}} \mid T, \phi) \, dG^o_{i} \, dG^u_{i} \, dY^{\text{mis}}_{i}.
$$

The posterior distribution of the model parameters based on the complete data likelihood function is derived in a straightforward manner as

$$
p(\phi, \theta \mid Y^{\text{obs}}_{i, Y^{\text{mis}}}, G^o, G^u, T, Z) \propto p(\phi, \theta) \prod_{i=1}^{N} p(Y^{\text{obs}}_{i, Y^{\text{mis}}}_{i}, G^o_{i, G^u_{i}}, T, \theta) \, p(G^o_{i, G^u_{i}} \mid T, \phi).
$$

Similarly, the posterior of the missing outcomes and DoIs conditional on the observed data and model parameters is

$$
\prod_{i=1}^{N} p\left\{ (Y^{\text{mis}}_{i} \mid Y^{\text{obs}}_{i}, G^o_{i, G^u_{i}}, T, \theta) \, p(G^o_{i, G^u_{i}} \mid Y^{\text{obs}}_{i}, T, \phi) \right\}.
$$

We use the Gibbs sampler to obtain posterior draws for the imputations and thereby derive the posterior distributions of the causal estimands using (Imbens and Rubin, 1997). Specifically, in each iteration of the sampler we alternate between drawing from the conditional posterior distributions of $\phi, \theta, G^o, G^u$, and $Y^{\text{mis}}$ given the other variables, respectively, and then use the draws of $Y^{\text{mis}}$ to obtain draws of the causal estimands from their respective posterior distributions. Details on the Gibbs sampler are in §4.4.
4.3 Bayesian Semiparametric Models

The models in our Bayesian approach consist of a combination of parametric and nonparametric specifications. We adopt a parametric model for the $Y$-model, with the particular form specified based on the specific setting under consideration. We shall discuss the specification of the $Y$-model later in our simulation studies and case study. In those studies, we set $\theta = (\beta, \lambda)^T$, where $\beta$ contain regression coefficients associated with the covariates and $\lambda$ is a scale parameter in the regression. The prior for $(\beta, \lambda)$ is the Normal-Inverse-Gamma distributions, $N(0, \sigma^2)IG(a_1, b_1)$ with $\sigma^2$, $a_1$ and $b_1$ being the variance, shape and scale parameters respectively.

The interference structure captured by the DoI is generally complex, hence unlikely to be adequately represented by parametric models. As such, we utilize a Bayesian nonparametric prior for the $G$-model, specifically, the Dependent Dirichlet process mixture prior (DDPM). As indicated by equation (2), the estimation of the causal estimands is reduced to estimation of the probability density function of the DDPM.

Our DDPM prior for the probability density function of the DoI, and this further justifies our use of the DDPM as it is a natural choice for density estimation under the Bayesian paradigm. In addition to its flexibility, the clustering property of the DDPM, in terms of its potential ability to automatically separate units into several clusters based on how they interfere with one another, provides another advantage that motivates our adoption of this nonparametric model. Quintana et al. (2022) provides a comprehensive review of the DDPM.

A random probability measure $H$ drawn from a Dirichlet Process, $\text{DP}(\alpha, H_0)$, with concentration parameter $\alpha > 0$ and base probability measure $H_0$ over a measurable space $(\Phi, B)$ is such that for any finite partition $(B_1, \ldots, B_k)$ of $\Phi$, $H(B_1), \ldots, H(B_k) \sim \text{Dir}(\alpha H_0(B_1), \ldots, \alpha H_0(B_k))$, where $\text{Dir}(\alpha_1, \ldots, \alpha_k)$ denotes the Dirichlet distribution with parameters $\alpha_1, \ldots, \alpha_k > 0$ (Ferguson 1974). The concentration parameter $\alpha$ determines the number of clusters in the Dirichlet Process. Our DDPM prior for the probability density function of the $G_i$ is specified as

$$(G_1, \ldots, G_N) | H_{T,Z} \overset{\text{ind}}{\sim} p(G | H_{T,Z}),$$

where $p(G | H_{T,Z}) = \int \eta(G | \phi) dH_{T,Z}(\phi)$ with $\eta(\cdot | \phi)$ being a continuous density function for every $\phi \in \Phi$, and $H_{T,Z} \sim \text{DP}(\alpha, H_0)$ for some $\alpha > 0$ and a base measure $H_0$ (Maceachern 2000), where $\overset{\text{ind}}{\sim}$ indicates independently distributed variables. The subscript of $H_{T,Z}$ indicates the dependence of the measure on $T$ and $Z$. As shown below, we include covariate information in the DDPM by allowing the locations in the stick-breaking construction of the $\text{DP}(\alpha, H_0)$ to depend on covariates. We consider using common weights across the values of DoI, so that our DDPM is a “single-weights” DDPM, for its simplicity and ease of implementation (which follows as it ignores the dependence of the weights on covariates). The representation of the stick-breaking process is

$$H_{T,Z} = \sum_{k=1}^{\infty} w_k \delta_{\phi_k(T,Z)}(\cdot), \quad w_k = v_k \prod_{i<k}(1 - v_i),$$

and $v_i \overset{\text{ind}}{\sim} \text{Beta}(1, \alpha)$. Ultimately, our $G$-model is

$$p(G_i | T_i, Z_{-i}, \Phi) \propto \sum_{k=1}^{\infty} w_k (\sigma_k^2)^{-1/2} \exp \left( -\frac{1}{2\sigma_k^2} (G_i - \mu(T_i, Z_{-i}, \gamma_k))^2 \right)$$

where the atoms $\phi_k = (\gamma_k, \sigma_k^2)$ and the weights $w_k$ are nonparametrically specified via the $\text{DP}(\alpha, H_0)$. This is an infinite mixture of Normal distributions with $\mu(T_i, Z_{-i}, \gamma_k)$ being the location parameter of each component that is a function of the covariates of unit $i$, the $i$th row of $A$, and treatment assignments excluding that of unit $i$. 

10
Finally, we specify the priors for $\alpha$ and $H_0$. We use the Gamma($a, b$) prior for $\alpha$ with shape $a$ and scale $b$ [Escobar and West (1995)], and obtain posterior draws of $\alpha$ via a Metropolis-Hastings step in our Gibbs sampler. Next, we take $H_0$ as a Normal-Inverse-Gamma conjugate $N(\mu_0, \sigma^2_0)I G(a_0, b_0)$ with $a_0$ and $b_0$ being the shape and scale parameter respectively. Our simulation studies and case study illustrate how the values for all the hyperparameters are set.

4.4 Details on the Gibbs Sampler

A discussion on efficient MCMC algorithms for DDPM models is provided by Escobar (1994). Our $Y$-model is generally specified by $\theta$ as $p(Y_i(z, g_i) | T_i, G_i = g_i, \theta)$. For our DDPM, we utilize an approximated blocked Gibbs sampler based on a truncation of the stick-breaking representation of the DP proposed by Ishwaran and Zarepour (2000). This algorithm proceeds by first selecting a conservative upper bound on the number $K$ of clusters. Let $C_i \in \{1, ..., K\}$ denote the latent class indicators for unit $i$. We specify a Multinomial distribution $C_i \sim MN(w)$ on $C_i$, where $w = (w_1, ..., w_K)^T$ contains the weights from the DDPM. Conditional on $C_i = k$, equation [4] is simplified to a single Normal component. Ishwaran and James (2001) demonstrated that an accurate approximation to the exact DP is obtained as long as $K$ is sufficiently large. To ensure this, we ran several MCMC iterations with different values of $K$ and increased $K$ after an iteration if all clusters were occupied. We terminated this process when the number of occupied clusters was less than $K$. Our entire algorithm proceeds as follows; throughout, ‘| − ’ indicates conditional on all others.

1. Given $\theta$, $\phi$ and $C = (C_1, ..., C_N)$, draw each $G_i^o$ from $p(G_i^o | − ) \propto p(Y_i^{obs} | T_i, G_i^o, \theta) p(G_i^o | T_i, Z_{−i}, \phi)$.

2. Given $\phi$, $w$ and $G_i^o$, draw each $C_i$ from $p(C_i = k | − ) \propto w_k p(G_i^o | T_i, Z_{−i}, \phi_k)$. 

3. Let $w_{K}^i = 1$. Given $\alpha$, $C$, draw $w_k^i$ for $k \in \{1, ..., K - 1\}$ from $w_k^i \sim \text{Beta}(1 + \sum_{i : C_i = k} 1, \alpha + \sum_{i : C_i > k} 1)$. Then, update $w_k = w_k^i \prod_{j \neq k} (1 - w_j^i)$ for $k = 1, \ldots, K$.

4. Given $C$ and $w'$, draw $\alpha$ from $p(\alpha | − ) \propto p(\alpha) \prod_{k = 1}^{K} f(w_k' | 1 + \sum_{i : C_i = k} 1, \alpha + \sum_{i : C_i > k} 1)$ where $f$ is the pdf of $w_k'$, the beta distribution.

5. Given $C$ and $G^o$, draw $\phi_k$ from $p(\phi_k | − ) \propto H_0(\phi_k) \prod_{i : C_i = k} p(G_i^o | T_i, Z_{−i}, \phi_k)$.

6. Given $\phi$, $C$ and $G^o$, draw $\theta$ from $p(\theta | − ) \propto p(\theta) \prod_{i = 1}^{N} p(Y_i^{obs} | T_i, G_i^o, \theta)$.

The imputations of the $Y_i^{mis}$ are generated from the $Y$-model using the $G_i^o$ in Step 1 and the $\theta$ in Step 6. Specifically, for $i = 1, \ldots, N$:

7. Given $G_i^o$ in Step 1 and $\theta$ in Step 6, draw from $Y_i(z', z_{−i}) \sim p(Y_i | Z_i = z', T_i, G_i^o, \theta)$.

To obtain posterior draws of the spillover effect estimands, we impute the $Y_i(z, z_{−i})$ by using the posterior draws of $G_i^u = G(T_i, z')$ and $\theta$.

8. Given $w$ in step 3, draw $C_i^u$ from $C_i^u \sim MN(w)$.

9. Given $C_i^u$ and $\phi$, draw $G_i^u$ from $G_i^u \sim N(\mu(T_i, z_{−i}^u, \gamma C_i^u), \sigma^2_{G_i^u})$.

10. Given $G_i^u$ in step 9 and $\theta$ in step 6, draw from $Y_i(z', z_{−i}^u) \sim p(Y_i | Z_i = z, T_i, G_i^u, \theta)$.

This Gibbs sampler ultimately enables us to impute all of the missing potential outcomes that define the assignment-conditional effects, and yields approximations for the expected effects. Additional technical details regarding these points and each step are provided in the supplement.
5 Simulation Studies

We perform simulation studies to evaluate the performance of our Bayesian methodology under our DoI framework with respect to that of the Horvitz-Thompson (HT) estimator in the case of a Bernoulli trial. The HT estimator was selected as the comparison method in our evaluation because Säve et al. (2021) and Li and Wager (2022) have demonstrated that it is a consistent estimator for the E-ATE under a Bernoulli trial under interference and that it possesses desirable large sample properties.

To compare these methods, we assess the bias, mean square error (MSE), and coverage of a procedure based on $M$ simulated datasets by calculating $\sum_{n=1}^{N_{\text{sim}}} (\tau - \hat{\tau}_n)/N_{\text{sim}}$, $\sum_{n=1}^{N_{\text{sim}}} (\tau - \hat{\tau}_n)^2 / N_{\text{sim}}$, and $\sum_{n=1}^{N_{\text{sim}}} 1 (\hat{\tau}_n \leq \tau \leq \hat{\tau}_n)/N_{\text{sim}}$, respectively, where $\tau$ denotes the true causal estimand, and $\hat{\tau}_n$, $\hat{\tau}_n^l$, and $\hat{\tau}_n^u$ are the estimates of the causal estimand, 2.5% lower endpoint of the interval estimator, and 95% upper endpoint of the interval estimator in dataset $n = 1, \ldots, N_{\text{sim}}$. Here, we take the posterior mean of the causal estimand as our Bayesian point estimator and the 95% central credible interval as the interval estimator. For the HT estimator under the Bernoulli trial, Säve et al. (2021) gave the variance estimator and confidence interval. As the true value of E-ATE is hard to obtain in closed form, we approximated it in our simulation study via the Monte Carlo average of 1000 draws for each generated dataset and graph.

We consider five different data-generating mechanisms and two different underlying networks of experimental units. First, the five data-generating mechanisms that we consider are:

1. $Y_i = X_i^T \beta + Z_i \tau + \psi_1 \sum_{j \in N_i} Z_j A_{ij} / (|N_i| + 1) + \epsilon_i,$
2. $Y_i = X_i^T \beta + Z_i \tau + \psi_1 \sum_{j \in N_i} P_j Z_j A_{ij} / (|N_i| + 1) + \epsilon_i,$
3. $Y_i = \left( X_i^T \beta + \psi_1 \sum_{j \in N_i} P_j Z_j A_{ij} / (|N_i| + 1) \right) \exp \left( \psi_2 Z_i \sum_{j \in N_i} P_j Z_j A_{ij} / (|N_i| + 1) \right) + \epsilon_i,$
4. $Y_i = \left( X_i^T \beta + Z_i \tau + \psi_1 \sum_{j \in N_i} P_j Z_j A_{ij} / (|N_i| + 1) \right) \exp \left( \psi_2 Z_i \sum_{j \in N_i} P_j Z_j A_{ij} / (|N_i| + 1) \right) + \epsilon_i,$
5. $Y_i = \left( X_i^T \beta + Z_i \tau + \psi_1 \sum_{j \in N_i} P_j Z_j A_{ij} / (|N_i| + 1) \right) \cos \left( \pi \psi_2 \sum_{j \in N_i} P_j Z_j A_{ij} / (|N_i| + 1) \right) + \epsilon_i,$

where $\beta = (-1, 1.5)^T$, $\tau = 5$, $\psi_1 = 2$, $\psi_2 = 0.2$, $X_i \in \mathbb{R}^2$ has first entry $X_{i,1} \equiv 1$ and second entry $X_{i,2} \sim N(0, 1)$, $Z_i \sim \text{Bernoulli}(0.5)$, $\epsilon_i \sim N(0, 1)$ is an additive error term, $A_{ij}$ is the $(i, j)$ entry of the adjacency matrix, $N_i$ is the set of neighboring units for unit $i$, and $P_i$ denotes a measure computed by the PageRank algorithm (Page et al. 1998). The $P_i$ is considered as a proxy for the “importance” of unit $i$ on the network. Scenario 1 corresponds to the stratified interference structure. Including $P_i$ in Scenarios 2–5 creates more complicated interference structures because $P_i$ is computed based on the number of incoming relationships and the importance of the corresponding source nodes. Furthermore, the treatment effect in Scenario 3 has a nonlinear multiplicative form in that the treatment variable $Z_i$ appears in the exponential function. Additionally, Scenarios 4 and 5 have additive and multiplicative treatment and spillover effects.

To see the performance of the methodologies under different underlying networks, we use two different random graph models to generate the underlying networks: the Erdős-Rényi and Barabási–Albert graphs (Frieze and Karoński 2016). In all five scenarios, we simulate $N_{\text{sim}} = 200$ datasets for each sample size of $N = 300, 500, 1000, 2000, 3000$. We ran the MCMC algorithm for 2000 iterations after a burn-in of 2000. We chose the iteration numbers after experimentation verifying they deliver stable results over multiple runs.
Figure 1: MSE of the HT estimator and our proposed Bayesian methodology for the Erdös-Rényi graphs ER(\(N,p\)) with different sparsity parameters \(p\).

Figure 2: MSE of the HT estimator and our proposed Bayesian methodology for the Barabási–Albert graphs BA(\(N,n_0,k\)) with different sparsity parameters \((n_0,k)\); \(n_0\) is the initial number of nodes and \(k\) is the number of edges a new node attaches to existing nodes.

The \(Y\)-model is specified by \(\theta = (\beta, \lambda)^T\) as: \(Y_i(z) \sim N(X_i^T\beta_z + G_i(T_i, Z_{-i}), \lambda_z)\). In the \(G\)-model we set \(\mu(T_i, Z_{-i}, \gamma_k) = \gamma_{k,1} \sum_{j \in N_i} Z_j A_{ij} + \gamma_{k,2} \sum_{j \in N_i} Z_j P_{ij} A_{ij}\). The prior distributions for the parameters in this case are \(\beta_z \sim N((0,0)^T, 10^2 I_2)\), where \(I_2\) is the 2 \(\times\) 2 identity matrix, \(\lambda_z \sim IG(0.1, 0.1)\), \(\gamma_{k,1}, \gamma_{k,2} \sim N(0, 10^2)\) and \(\sigma_k^2 \sim IG(0.1, 0.1)\). For scenarios 3–5, the inference model is severely misspecified in that the data-generating models are multiplicative of treatment and spillover effect terms (exp and cos), whereas the inferential model is additive. These scenarios assess the robustness of our approach to model misspecification.

Figures 1 and 2 summarize the MSE of the two methods. These figures show that the proposed approach outperforms the HT estimator for all scenarios across both networks. The MSE converges at the same rate across all scenarios, indicating that the proposed approach is as efficient as the frequentist estimator. The proposed approach works consistently well even under model misspecifications, i.e., Scenarios 3–5, demonstrating the efficiency and robustness of the DoI-based approach. Further evaluation of bias and coverage for the treatment effects and all metrics for the spillover effects are provided in the Supplementary Material in tabular forms. Those results also demonstrate the efficiency of the proposed method under interference.

6 Case Study: Cash Transfer Programs in Colombia

The methodology developed in §4 is applied to a randomized experiment investigating the impact of a conditional cash transfer program, treatment, on students’ attendance rates (Barrera-Osorio et al., 2011). The study was conducted in two regions of Bogota, Colombia: San Cristobal and
Suba. Households with school children, ranging from one to five children per household, were recruited for the experiment. Within each household, school children were randomly assigned to either enroll in the cash transfer program or not, by a stratified randomization approach based on locality (San Cristobal/Suba), school type (public/private), gender, and grade level.

Multiple children within a household had the potential to receive the treatment, and within each stratum, all children had an equal chance of being treated. Additionally, given the known treatment assignment, Barrera-Osorio et al. (2011) examined both the direct effect of the program and the possible spillover effect on siblings within the same household. Enrolled students were eligible for cash subsidies if they attended school at least 80% of the time in a given month. Enrolling one student in the program could influence the attendance rate of their sibling(s) in the same household, either positively or negatively. Our objective is to employ the proposed methodology to analyze these direct and spillover effects in the two regions of Bogota, Colombia.

Formally, for student $i$, we denote $Y_i \in \{0, 1\}$ where $Y_i = 1$ if the student is eligible for cash subsidies, i.e., their attendance rate is over 80%. We also let $Z_i$ denote a treatment indicator that equals 1 if student $i$ was enrolled in the program, and 0 otherwise. We consider the interference occurs within the same household and thus denote $A_{i,j}$ as the adjacency that equals 1 if student $i$ and $j$ are in the same household (siblings), and 0 otherwise. $N_i$ denotes the number of siblings of unit $i$. Additionally, $X_i$ represents the pre-treatment covariates, including the student’s age, grade level, gender, household head’s age, single-parent household indicator, household size, household’s poverty score, household’s income status, locality, and the number of students in the household who participated in the lottery. We establish three cluster types based on the two regions and household sizes: households in Suba with two students, households in San Cristobal with two students, and households in San Cristobal with three or more students. Overall, our analysis comprises 1012 households containing 2135 students.

We adopt the original treatment randomization probability as the treatment allocation strategy, indicated by the treatment allocation probabilities $\pi = (\pi_{SC}, \pi_{Su}) = (0.628, 0.449)$ for the two localities San Cristobal and Suba respectively. The primary estimands of interest are E-ATE $\tau_{E-ATE}(1; \pi)$ and E-ASE $\tau_{E-ASE}(0; \pi)$ for the current treatment allocation strategy $\pi$, see §3.2. In the given context, the spillover effect measures the discrepancy in the attendance rate of an unenrolled student under the allocation strategy $\pi$ when none of their siblings is enrolled.

We use a probit model for the $Y$-model: $Y_i(z) \sim \text{Bernoulli (}\Psi(X_i^T \beta + G_i(T_i, Z_{-i})))$, where $\Psi(\cdot)$ denotes the standard normal cumulative distribution function. The details of the Gibbs sampler for the binary outcome are provided in the Supplementary Material. We here customize the components of $G_i$ to reflect our knowledge about the interference: $\mu(T_i, Z_{-i}, \gamma_k) = \gamma_{k,1} \sum_{j \in N_i} Z_j A_{ij} / |N_i| + \gamma_{k,2} \left( \text{Grade}_i - \sum_{j \in N_i} \text{Grade}_j Z_j A_{ij} / |N_i| \right)$, where Grade denotes the grade of student $i$. The intuition of this specification is that students are more encouraged to enroll in the program if more or fewer siblings are treated, and they are likely influenced if their siblings are close in grade. We use the same weakly informative priors, $\beta_\cdot \sim N((0, 0)^T, 10^2 I_2)$, where $I_2$ is the $2 \times 2$ identity matrix, $\gamma_{k,1}, \gamma_{k,2} \sim N(0, 10^2)$ and $\sigma_k^2 \sim IG(0.01, 0.01)$. We present a sensitivity analysis with respect to different prior specifications in the following section.

Table II presents the estimates of E-ATE and E-ASE under the current treatment allocation strategy $\pi$. We also present the conditional E-ASE for those with 1 or 2 siblings and those whose siblings’ treated proportion is 33%, 50%, or 66%. We observe that the lower bound of E-ATE is positive and the upper bound of E-ASE is negative, suggesting the positive direct treatment effect and
Table 1: Causal estimands and their estimates of posterior means, standard deviation, percentiles, and interval length under the current treatment allocation strategy $\pi$. The superscript $nb,k$ denotes the spillover effects on students with $k$ siblings. The superscript $rt,p$ denotes the spillover effects on students for whom $p\%$ of their siblings are treated. E-ATE, expected average treatment effects; E-ASE, expected average spillover effects.

| Estimands  | Mean   | SD    | 2.5%     | Median | 97.5%  | Length |
|------------|--------|-------|----------|--------|--------|--------|
| E-ATE$_{\pi}$ | 0.011  | 0.007 | -0.003   | 0.011  | 0.025  | 0.028  |
| E-ASE$_{\pi}$ | -0.003 | 0.006 | -0.014   | -0.003 | 0.009  | 0.023  |
| E-ASE$_{nb,1}^{rt}$ | -0.003 | 0.006 | -0.014   | -0.003 | 0.01   | 0.024  |
| E-ASE$_{nb,2}^{rt}$ | 0.0   | 0.017 | -0.033   | 0.0    | 0.036  | 0.069  |
| E-ASE$_{rt,\pi}^{p}$ | -0.001 | 0.026 | -0.053   | 0.0    | 0.051  | 0.105  |
| E-ASE$_{rt,0.50}^{p}$ | -0.0  | 0.008 | -0.016   | -0.001 | 0.016  | 0.032  |
| E-ASE$_{rt,0.66}^{p}$ | 0.004 | 0.029 | -0.051   | 0.0    | 0.061  | 0.113  |

Table 2: The spillover effects and their estimates of posterior means, standard deviation, percentiles, and interval length under different treatment allocation strategies $\pi'$. E-ASE$_{p}$ denotes expected average spillover effects under the treatment allocation strategy $\pi' = (p, p)$.

| Estimands  | Mean   | SD    | 2.5%     | Median | 97.5%  | Length |
|------------|--------|-------|----------|--------|--------|--------|
| E-ASE$_{\pi}$ | -0.003 | 0.006 | -0.014   | -0.003 | 0.009  | 0.023  |
| E-ASE$_{0.1}^{rt}$ | -0.011 | 0.008 | -0.029   | -0.011 | 0.004  | 0.033  |
| E-ASE$_{0.3}^{rt}$ | -0.009 | 0.007 | -0.023   | -0.008 | 0.005  | 0.029  |
| E-ASE$_{0.5}^{rt}$ | -0.006 | 0.007 | -0.019   | -0.007 | 0.006  | 0.025  |
| E-ASE$_{0.7}^{rt}$ | -0.004 | 0.006 | -0.017   | -0.004 | 0.008  | 0.025  |
| E-ASE$_{0.9}^{rt}$ | -0.002 | 0.006 | -0.015   | -0.002 | 0.01   | 0.024  |

negative spillover effects. The spillover effects tend to be slightly larger as the number of siblings or the proportion of treated siblings increases. Table 2 presents the estimates of the E-ASE under different treatment allocation strategies $\pi' \in \{(0.1, 0.1), (0.3, 0.3), (0.5, 0.5), (0.7, 0.7), (0.9, 0.9)\}$, $\tau_{E-ASE}(0; \pi')$. We observe that the effect is always negative. As the proportion of treated units increases, the effects slightly get larger.

Finally, when performing Bayesian analyses with weakly identifiable models, it is important to investigate the robustness of the results with respect to the prior specifications so as to make inferences more reliable. Considering different prior specifications, we examined the robustness of our analyses; the results are provided in the online supplement.

7 Concluding Remarks

This article proposed our new notion of the Degree of Interference (DoI) that flexibly handles unknown interference. We presented several new types of causal estimands and developed a fast data augmentation based Bayesian semiparametric methodology for estimating them. In simulation studies across different scenarios with complicated interference structures our approach exhibited better finite-sample performance than that of the conventional HT estimator in terms of MSE. Our two case studies demonstrate that our DoI framework and Bayesian methodology yield more insights and knowledge on interference in a straightforward manner. A possible future
direction to build upon this framework would be to extend our framework to observational studies where the treatment assignment mechanisms take unknown functional forms.

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A Detailed Steps of Gibbs Sampler

In this section we present the detailed steps of the Gibbs sampler developed in §4.4. First, our implementation of the Gibbs sampler is outlined below.

1. Initialize parameters $\phi^{(0)}, \theta^{(0)}$.
2. Repeat the following steps. For $t = 0, 1, \ldots$
   
   (a) Draw $G^{o,(t+1)} \sim p \left( G \mid T, Y^{\text{obs}}, \phi^{(t)}, \theta^{(t)} \right)$.
   
   (b) Draw $\phi^{(t+1)} \sim p \left( \phi, \theta \mid T, Y^{\text{obs}}, G^{o,(t+1)} \right)$.
   
   (c) Draw $G^{u,(t+1)} \sim p \left( G \mid T, Y^{\text{obs}}, G^{o,(t+1)}, \phi^{(t+1)}, \theta^{(t+1)} \right)$.
   
   (d) Draw $Y^{\text{mis},(t+1)} \sim p \left( Y^{\text{mis}} \mid T, Y^{\text{obs}}, G^{o,(t+1)}, G^{u,(t+1)}, \phi^{(t+1)}, \theta^{(t+1)} \right)$.
   
   (e) Calculate the causal estimand based on the observed and imputed data.

The detailed steps are provided below.

1. For the units with $C_i = k$ and $Z_i = z$, draw $G^o_i$ from $N\left( \{ \lambda_i \gamma_0 + \gamma_k \sum_{j \in N_i} Z_j A_{ij} \} + (Y_i - X_i^T \beta_i) \sigma_k^2, (\lambda_z + \sigma_k^2), \lambda_z \sigma_k^2 / (\lambda_z + \sigma_k^2) \right)$.
2. For each unit, draw $C_i$ from $\text{Pr}(C_i = k | \cdot) = \frac{w_k N(G_i | \gamma_0 + \gamma_k \sum_{j \in N_i} Z_j A_{ij}, \sigma_k^2)}{\sum_{k=1}^K w_k N(G_i | \gamma_0 + \gamma_k \sum_{j \in N_i} Z_j A_{ij}, \sigma_k^2)}$
3. Update $w_k$'s as Step 3 in §4.4.
4. We adopt the Metropolis Hasting step with a proposal draw $\alpha^* \sim \text{Gamma}(1, 1)$. Given $w'$ and $C$, we accept the draw $\alpha^*$ with probability $r$ where

$$r = \min \left\{ 1, \frac{p(\alpha^* | 1, 1) \prod_{k=1}^K f \left( w'_k \mid 1 + \sum_{i:C_i = k} 1, \alpha^* + \sum_{i:C_i > k} 1 \right)}{p(\alpha | 1, 1) \prod_{k=1}^K f \left( w'_k \mid 1 + \sum_{i:C_i = k} 1, \alpha + \sum_{i:C_i > k} 1 \right)} \right\}.$$ 

We reject the draw $\alpha^*$ with probability $1 - r$ and accept the draw $\alpha$, obtained in the previous iteration. Note that $p(\alpha \mid a, b)$ denotes the prior distribution of $\alpha$, which we specify by the Gamma distribution with parameters $a = 1, b = 1$ and $f$ is specified in Step 4 in §4.4.

5. (a) If $N_k = \sum_{i=1}^N \mathbf{1}(C_i = k) > 0$, draw $\sigma^2_k$ from $\text{IG}(2 + 0.5N_k, 1 + 0.5s_k)$ where $s_k = \sum_{i:C_i = k} (G^o_i - \gamma_0 - \gamma_k \sum_{j \in N_i} Z_j A_{ij})^2$. If $N_k = 0$, then draw $\sigma^2_k$ from the prior $\text{IG}(2, 1)$.
   
   (b) If $N_k > 0$, draw $\gamma_k$ from $\text{N}((M_k^T M_k + \sigma^2_k / 10 \times I_2)^{-1} M_k^T G_k^o, \sigma^2_k (M_k^T M_k + \sigma^2_k / 10 \times I_2)^{-1})$ where $G_k^o$ is a subvector of $G^o$ for units with $C_i = k$ and $M_k$ is an $N_k \times 2$ matrix with the row vector $(1, \sum_{j \in N_i} Z_j A_{ij})$ for units with $C_i = k$. If $N_k = 0$, then draw from $\text{N}((0, 0)^T, \sigma^2_k / 10 \times I_2)$.
6. (a) Draw $\lambda_z$ from $\text{IG}(2 + N_z / 2, 1 + 0.5 \sum_{i:Z_i = z} (Y_i - X_i^T \beta_z - G_z^o)^2)$.
   
   (b) Draw $\beta_z$ from $\text{N}(X_z^T X_z + \lambda_z / 10 \times I_2)^{-1} X_z^T (Y_z - G_z^o), \lambda_z (X_z^T X_z + \lambda_z / 10 \times I_2)^{-1})$, where $G_z^o$, $X_z$, and $Y_z$ are respectively a submatrix and a subvector of $G^o$, $X$ and $Y$ for units with $Z_i = z$.

7-10. Follow the procedures in §4.4 for assignment-conditional effects or the steps provided in the next section for the expected effects, depending on which of the effects (conditional effects or expected effects) one is interested in.
A.1 Estimation of Expected Effects

We present a methodology to infer the expected effects. It suffices to derive the procedures for estimating $\mathbb{E}_{Z_{-i} \sim \pi}[Y_i(z, Z_{-i})]$ where $\pi$ is the marginalized distribution of $Z_{-i}$, which depends on the experimental design. For notational convenience, the outer integral below is taken with respect to the counting measure $\pi$ for the distribution of $Z_{-i}$.

\[
\mathbb{E}_{Z_{-i} \sim \pi}[Y_i(z, Z_{-i})] = \mathbb{E}_{Z_{-i} \sim \pi} \left[ Y_i(z, Z_{-i}) \int_g dF_{G_i|Z_{-i}=z_{-i}}(g) \right] = \mathbb{E}_{Z_{-i} \sim \pi} \left[ \int_g Y_i(z, Z_{-i}) dF_{G_i|Z_{-i}=z_{-i}}(g) \right] = \int_{\Omega^{N-1}} \int_g Y_i(z, g) dF_{G_i|Z_{-i}=z_{-i}}(g) d\pi(z_{-i}) \approx \frac{1}{M} \sum_{m=1}^M Y_i(z, g_i^{(m)})
\]

$g_i^{(m)}$ is obtained by the following two steps. For $m = 1, \ldots, M$, (1) $z^m \sim \pi$, and then (2) $g_i^{(m)} \sim F_{G_i|Z_{-i}=z^m_i}(g)$ where $z_{-i}^m$ is a subvector of $z^m$ with the $i$-th entry removed. We infer the expected effects by following the same steps from step 1 to step 6 in §4.4 and then we proceed as:

7. Draw $z^* \sim \pi$.
8. Given $w$ in step 3, draw $C_i^*$ from $C_i^* \sim \text{MN}(w)$.
9. (a) Given $C_i^*$, $\gamma$, and $\sigma^2$, draw $G_i^*$ from $G_i^* \sim N(\mu(T_i, z_{-i}^*, \gamma(C_i^*)), \sigma^2_{C_i^*})$.
    (b) Given $C_i^*$, $\gamma$, and $\sigma^2$, draw $G_i^0$ from $G_i^0 \sim N(\mu(T_i, 0^{N-1}, \gamma(C_i^*)), \sigma^2_{C_i^*})$.
10. (a) Given $G_i^*$ in step 9(a) and $\theta$ in step 6, draw from $Y_i(z, z_{-i}^*) \sim \Pr(Y_i|Z_i = z, T_i, G_i^*, \theta)$ for $z = 0, 1$.
     (b) Given $G_i^0$ in step 9(b) and $\theta$ in step 6, draw from $Y_i(0, 0^{N-1}) \sim \Pr(Y_i|Z_i = 0, T_i, G_i^0, \theta)$.

A.2 Extension to Binary Outcomes

When we adopt a Probit model as in our illustrative analyses in §6 we draw the sample of $\beta_z$ as follows. Given $G_i^0 = g_i$ and $\beta_z$ in the previous iteration, we first draw an auxiliary variable $v_i$ for unit $i$ with $Z_i = z$ from

\[
v_i \sim \begin{cases} 
\text{TN}(X_i \beta_z + g_i, 1, 0, \infty) & \text{if } Y_i = 1, \\
\text{TN}(X_i \beta_z + g_i, 1, -\infty, 0) & \text{if } Y_i = 0,
\end{cases}
\]

where $\text{TN}(\mu, \sigma^2, l, u)$ denotes the truncated normal distribution with the mean, variance, lower bound, and upper bound parameters. Then, we draw from $\beta_z \sim N(M, V)$, where $V = (H_0^{-1} + X_i^T X_z)^{-1}$, $M = V(H_0^{-1} h_0 + X_i^T v_z)$ with $v_z$ being a subvector of $v = (v_1, \ldots, v_N)$ for units with $Z_i = z$. The prior distribution of $\beta_z$ is specified here by $\beta_z \sim N(h_0, H_0)$.

B Simulation Results

This section presents the simulation evaluation not presented in §5; bias and coverage for E-ATE, and bias, MSE, and coverage for E-ASE. We consider the same data-generating mechanisms as in §5; Table 3-7 present the results.
Table 3: Evaluation metrics of simulations studies (Scenario 1). The data-generating process (DGP) and the underlying network between units are provided in §5.

| DGP   | Network (parameters) | N  | HT (E-ATE) Bias | MSE | Coverage | DoI (E-ATE) Bias | MSE | Coverage | DoI (E-ASE) Bias | MSE | Coverage |
|-------|----------------------|----|----------------|-----|----------|-----------------|-----|----------|-----------------|-----|----------|
| ER(p = 0.01) |                        |    | -0.018 | 0.1104 | 0.985 | -0.0146 | 0.0129 | 0.95 | -0.0011 | 0.0071 | 0.99 |
|        | ER(p = 0.03) |                        |    | -0.0121 | 0.0679 | 0.985 | -0.0059 | 0.0069 | 0.945 | 0.0104 | 0.0098 | 0.965 |
|        | 1000 | 0.0053 | 0.0337 | 0.99 | 0.0021 | 0.0046 | 0.96 | 0.0018 | 0.0097 | 0.95 |
|        | ER(p = 0.05) |                        |    | -0.0137 | 0.0341 | 0.99 | -0.0083 | 0.0044 | 0.975 | -0.0021 | 0.0409 | 0.83 |
|        | 1000 | 0.0122 | 0.1354 | 0.98 | -0.0047 | 0.0141 | 0.97 | 0.0137 | 0.0538 | 0.925 |
|        | BA(n0 = 10, k = 3) |                        |    | 0.0315 | 0.114 | 0.99 | 0.0036 | 0.0118 | 0.975 | -0.0035 | 0.0156 | 0.985 |
|        | 1000 | 0.0051 | 0.0732 | 0.98 | -0.006 | 0.0087 | 0.95 | 0.0101 | 0.0084 | 0.99 |
|        | BA(n0 = 10, k = 5) |                        |    | -0.0005 | 0.0325 | 0.995 | -0.0011 | 0.004 | 0.97 | 0.0054 | 0.0036 | 0.99 |
|        | 1000 | -0.0428 | 0.1237 | 0.975 | -0.0175 | 0.0125 | 0.975 | 0.0121 | 0.0269 | 0.97 |
|        | BA(n0 = 30, k = 3) |                        |    | 0.0001 | 0.0695 | 0.99 | -0.0072 | 0.0085 | 0.95 | -0.0064 | 0.0075 | 0.965 |
|        | 1000 | 0.0026 | 0.0337 | 0.985 | -0.0002 | 0.004 | 0.97 | 0.0026 | 0.0042 | 0.985 |
|        | BA(n0 = 30, k = 5) |                        |    | -0.0040 | 0.1212 | 0.98 | -0.0168 | 0.0122 | 0.97 | -0.003 | 0.0171 | 0.98 |
|        | 1000 | -0.0068 | 0.0631 | 1.0 | -0.0032 | 0.0077 | 0.975 | 0.005 | 0.0124 | 0.99 |
|        | 1000 | -0.0046 | 0.0372 | 0.99 | 0.0012 | 0.0036 | 0.975 | 0.0006 | 0.0074 | 0.975 |

For E-ATE, the DoI approach outperforms the HT estimator in terms of MSE and bias for almost all scenarios of both networks even under severe model misspecifications. The MSE decreases as N increases, and the coverage probability is well-calibrated with 95% probability. These observations demonstrate the efficiency and robustness of our DoI approach. The HT estimator sometimes exhibits large bias when N is small, whereas the Bayesian methodology consistently works well. Both methodologies exhibit smaller bias as N increases. They are well-calibrated in that they yield intervals with nearly 95% coverage across nearly all conditions for the E-ATE, indicating the robustness of our methodologies for mispecifications.

Another advantage of the DoI methodology is that it provides a principled estimation procedure for the spillover effects. Overall, the DoI methodology yields good bias, MSE, and coverage for the E-ASE across most scenarios. Only when the data-generating process is scenario 4, it sometimes exhibits slightly larger bias and smaller coverages, especially when the model is severely misspecified (Scenario 4, ER(p = 0.03, 0.05)). This phenomenon can be explained in terms of the connectedness of the underlying graph. The Erdős-Rényi (ER) graph is known to be almost surely connected when $p > \frac{(1+\epsilon)\log N}{N}$ for any $\epsilon > 0$, and disconnected otherwise (Frieze and Karoński, 2016). Therefore, when $p = 0.05$ and $N = 1000$, the ER graph is likely to be fully connected, hence the influence of misspecification inflates and dominates the flexibility of our models; that is, in the case of a misspecified model, the Bayesian posterior credible intervals could be discrepant with the confidence interval. In contrast, the BA graph usually yields disconnected graphs from its construction, leading to the good performance of our methodology for the E-ASE under the BA graphs. All scenarios exhibit good performance when the underlying graph is the Barabási–Albert (BA) graph. This observation should have minimal impact on our empirical analyses because any of our datasets in the empirical analyses do not have fully connected graphs as the underlying structure, and have multiple independent connected subgraphs in it.
| Scenario 2 | Network (parameters) | N   | Bias  | MSE  | Coverage | Bias  | MSE  | Coverage |
|------------|----------------------|-----|-------|------|----------|-------|------|----------|
| ER(p = 0.01) |                      | 50  | -0.0116 | 0.0542 | 0.995 | -0.0066 | 0.0091 | 0.95 | 0.0139 | 0.0103 | 0.96 |
|             |                      | 1000 | 0.005  | 0.0273 | 0.995 | 0.0018 | 0.0046 | 0.96 | -0.0038 | 0.0092 | 0.935 |
| ER(p = 0.03) |                      | 300  | -0.0332 | 0.0868 | 1.0 | 0.0007 | 0.0122 | 0.975 | -0.0047 | 0.0289 | 0.94 |
|             |                      | 500  | 0.0022 | 0.0667 | 0.995 | -0.0121 | 0.0077 | 0.95 | -0.0247 | 0.0371 | 0.905 |
| ER(p = 0.05) |                      | 1000 | -0.0116 | 0.0285 | 1.0 | -0.0082 | 0.0041 | 0.965 | 0.003 | 0.0306 | 0.965 |
| BA(n_0 = 10, k = 3) |              | 300  | 0.0291 | 0.0827 | 0.995 | 0.0044 | 0.0119 | 0.985 | 0.0021 | 0.0176 | 0.955 |
|             |                      | 500  | -0.0034 | 0.0524 | 0.995 | -0.0059 | 0.0065 | 0.94 | 0.0142 | 0.0101 | 0.97 |
| BA(n_0 = 10, k = 5) |              | 1000 | -0.0062 | 0.0212 | 1.0 | -0.001 | 0.0042 | 0.94 | 0.0055 | 0.0045 | 0.975 |
| BA(n_0 = 30, k = 3) |             | 300  | 0.0399 | 0.0824 | 0.995 | 0.003 | 0.0117 | 0.985 | 0.0063 | 0.0111 | 0.985 |
|             |                      | 500  | -0.0015 | 0.0523 | 0.995 | -0.0077 | 0.0084 | 0.955 | -0.0103 | 0.0081 | 0.95 |
| BA(n_0 = 30, k = 5) |             | 1000 | -0.0055 | 0.0222 | 1.0 | -0.0006 | 0.004 | 0.97 | 0.0038 | 0.0047 | 0.955 |
| Table 5: Evaluation metrics of simulations studies (Scenario 3). |

| Scenario 3 | Network (parameters) | N   | Bias  | MSE  | Coverage | Bias  | MSE  | Coverage | Bias  | MSE  | Coverage |
|------------|----------------------|-----|-------|------|----------|-------|------|----------|-------|------|----------|
| ER(p = 0.01) |                      | 300  | -0.0267 | 0.0509 | 0.965 | -0.0133 | 0.013 | 0.955 | -0.006 | 0.0086 | 0.97 |
|             |                      | 500  | 0.0146 | 0.0321 | 0.925 | -0.0057 | 0.0088 | 0.94 | 0.0081 | 0.0103 | 0.96 |
| ER(p = 0.03) |                      | 300  | -0.0246 | 0.0452 | 0.965 | 0.0016 | 0.0122 | 0.975 | -0.0086 | 0.0301 | 0.95 |
|             |                      | 500  | -0.0084 | 0.0309 | 0.925 | -0.0131 | 0.0078 | 0.955 | -0.014 | 0.0345 | 0.915 |
| ER(p = 0.05) |                      | 1000 | 0.0028 | 0.013 | 0.96 | -0.0088 | 0.0041 | 0.965 | 0.0143 | 0.0413 | 0.86 |
| BA(n_0 = 10, k = 3) |              | 300  | -0.0091 | 0.0543 | 0.92 | -0.045 | 0.0145 | 0.97 | 0.0241 | 0.0591 | 0.905 |
|             |                      | 500  | 0.0055 | 0.0305 | 0.945 | 0.0036 | 0.0083 | 0.975 | -0.0066 | 0.0684 | 0.89 |
| BA(n_0 = 10, k = 5) |              | 1000 | 0.0015 | 0.0152 | 0.95 | 0.0005 | 0.0039 | 0.955 | -0.0166 | 0.0644 | 0.82 |
| BA(n_0 = 30, k = 3) |             | 300  | 0.0242 | 0.0674 | 0.95 | 0.005 | 0.012 | 0.975 | -0.041 | 0.0172 | 0.97 |
|             |                      | 500  | 0.0018 | 0.0342 | 0.94 | -0.0054 | 0.0085 | 0.945 | 0.0088 | 0.0098 | 0.97 |
| BA(n_0 = 30, k = 5) |             | 1000 | 0.0102 | 0.0188 | 0.95 | -0.0004 | 0.004 | 0.965 | 0.0011 | 0.0043 | 0.97 |
| Table 4: Evaluation metrics of simulations studies (Scenario 2). |
### Table 6: Evaluation metrics of simulations studies (Scenario 4).

| DGP          | Network (parameters) | N  | HT (E-ATE) | DoI (E-ATE) | DoI (E-ASE) |
|--------------|----------------------|----|------------|-------------|-------------|
| Scenario 4   | ER($p = 0.01$)       | 300| -0.0201   | 0.0921      | 0.995       |
|              |                      | 500| -0.0119   | 0.0579      | 0.99        |
|              |                      | 1000| 0.0051   | 0.0295      | 1.0         |
|              | ER($p = 0.03$)       | 300| -0.0336   | 0.0972      | 1.0         |
|              |                      | 500| 0.0032   | 0.0733      | 0.995       |
|              |                      | 1000| -0.0123  | 0.0317      | 0.995       |
|              | ER($p = 0.05$)       | 300| -0.0123   | 0.1243      | 0.98        |
|              |                      | 500| -0.0184   | 0.0679      | 1.0         |
|              | BA($n_0 = 10, k = 3$) | 300| 0.0296    | 0.0857      | 0.99        |
|              |                      | 500| -0.0035   | 0.0554      | 0.99        |
|              | BA($n_0 = 10, k = 5$) | 1000| -0.0061  | 0.0217      | 1.0         |
|              |                      | 1000| -0.003  | 0.0456      | 1.0         |
|              | BA($n_0 = 30, k = 3$) | 300| 0.032    | 0.0855      | 0.99        |
|              |                      | 500| -0.0011   | 0.0538      | 0.995       |
|              | BA($n_0 = 30, k = 5$) | 1000| -0.0004  | 0.0227      | 1.0         |

### Table 7: Evaluation metrics of simulations studies (Scenario 5).

| DGP          | Network (parameters) | N  | HT (E-ATE) | DoI (E-ATE) | DoI (E-ASE) |
|--------------|----------------------|----|------------|-------------|-------------|
| Scenario 5   | ER($p = 0.01$)       | 300| -0.0194   | 0.0851      | 0.995       |
|              |                      | 500| -0.0116   | 0.0532      | 0.99        |
|              |                      | 1000| 0.0042  | 0.0266      | 0.995       |
|              | ER($p = 0.03$)       | 300| -0.0332   | 0.0859      | 1.0         |
|              |                      | 500| 0.0018   | 0.0644      | 0.99        |
|              |                      | 1000| -0.0114  | 0.0275      | 1.0         |
|              | ER($p = 0.05$)       | 300| -0.0123   | 0.1094      | 0.98        |
|              |                      | 500| -0.0173   | 0.0588      | 1.0         |
|              |                      | 1000| 0.0032  | 0.0352      | 0.995       |
|              | BA($n_0 = 10, k = 3$) | 300| 0.0291    | 0.0821      | 0.99        |
|              |                      | 500| -0.0034   | 0.0521      | 0.99        |
|              |                      | 1000| -0.0062  | 0.0212      | 1.0         |
|              | BA($n_0 = 10, k = 5$) | 300| -0.0388   | 0.0884      | 0.98        |
|              |                      | 500| -0.0029   | 0.0437      | 1.0         |
|              |                      | 1000| -0.0201  | 0.0211      | 1.0         |
|              | BA($n_0 = 30, k = 3$) | 300| 0.0305    | 0.0817      | 0.99        |
|              |                      | 500| -0.0016   | 0.052       | 0.995       |
|              |                      | 1000| -0.0005  | 0.0222      | 1.0         |
|              | BA($n_0 = 30, k = 5$) | 300| -0.0368   | 0.0874      | 0.98        |
|              |                      | 500| -0.0038   | 0.0434      | 1.0         |
|              |                      | 1000| -0.0028  | 0.0252      | 0.99        |
Table 8: Sensitivity Analysis for different prior specifications (Prior 1).

| Estimands | Mean   | SD     | 2.5%  | Median | 97.5% | Length |
|-----------|--------|--------|-------|--------|-------|--------|
| E-ATEπ   | 0.011  | 0.007  | -0.002| 0.011  | 0.026 | 0.028  |
| E-ASEπ   | -0.004 | 0.006  | -0.016| -0.004 | 0.008 | 0.024  |
| E-ASEπb,1| -0.004 | 0.006  | -0.016| -0.004 | 0.009 | 0.026  |
| E-ASEπb,2| -0.002 | 0.018  | -0.036| -0.004 | 0.033 | 0.069  |
| E-ASEπr,33| -0.003 | 0.027  | -0.055| 0.0     | 0.05  | 0.105  |
| E-ASEπr,50| -0.003 | 0.008  | -0.019| -0.004 | 0.013 | 0.033  |
| E-ASEπr,66| -0.0  | 0.03   | -0.058| 0.0     | 0.06  | 0.118  |

Table 9: Sensitivity Analysis for different prior specifications (Prior 2).

| Estimands | Mean   | SD     | 2.5%  | Median | 97.5% | Length |
|-----------|--------|--------|-------|--------|-------|--------|
| E-ATEπ   | 0.01   | 0.006  | -0.002| 0.009  | 0.023 | 0.025  |
| E-ASEπ   | -0.0   | 0.006  | -0.011| -0.0   | 0.012 | 0.023  |
| E-ASEπb,1| -0.0   | 0.006  | -0.012| 0.0    | 0.013 | 0.024  |
| E-ASEπb,2| 0.002  | 0.017  | -0.033| 0.0    | 0.036 | 0.069  |
| E-ASEπr,33| 0.002  | 0.026  | -0.048| 0.0    | 0.054 | 0.101  |
| E-ASEπr,50| 0.0    | 0.008  | -0.015| 0.0    | 0.016 | 0.031  |
| E-ASEπr,66| 0.003  | 0.03   | -0.052| 0.0    | 0.065 | 0.117  |

C Sensitivity Analyses for the Case Study in the Main Text

In this section, we present sensitivity analyses with different prior specifications: Table 9 and 11 report the results with the following priors:

Prior 1: \[ \beta_2 \sim N((0, 0)^T, 20^2I_2), \gamma_{k,1}, \gamma_{k,2} \sim N(0, 20^2), \sigma_k^2 \sim IG(0.1, 0.1). \]

Prior 2: \[ \beta_2 \sim N((0, 0)^T, 5^2I_2), \gamma_{k,1}, \gamma_{k,2} \sim N(0, 5^2), \sigma_k^2 \sim IG(0.1, 0.1). \]

Prior 3: \[ \beta_2 \sim N((0, 0)^T, 10^2I_2), \gamma_{k,1}, \gamma_{k,2} \sim N(0, 10^2), \sigma_k^2 \sim IG(0.01, 0.01). \]

Prior 4: \[ \beta_2 \sim N((0, 0)^T, 10^2I_2), \gamma_{k,1}, \gamma_{k,2} \sim N(0, 10^2), \sigma_k^2 \sim IG(1.0, 1.0). \]

Compared with Table 11, the results change only slightly, validating the robustness of our analyses.

Table 10: Sensitivity Analysis for different prior specifications (Prior 3).

| Estimands | Mean   | SD     | 2.5%  | Median | 97.5% | Length |
|-----------|--------|--------|-------|--------|-------|--------|
| E-ATEπ   | 0.011  | 0.007  | -0.002| 0.011  | 0.026 | 0.028  |
| E-ASEπ   | -0.003 | 0.006  | -0.015| -0.003 | 0.01  | 0.025  |
| E-ASEπb,1| -0.003 | 0.007  | -0.015| -0.003 | 0.01  | 0.025  |
| E-ASEπb,2| -0.0   | 0.018  | -0.033| 0.0    | 0.036 | 0.069  |
| E-ASEπr,33| -0.001 | 0.026  | -0.052| 0.0    | 0.051 | 0.103  |
| E-ASEπr,50| -0.002 | 0.008  | -0.017| -0.002 | 0.015 | 0.032  |
| E-ASEπr,66| 0.001  | 0.03   | -0.056| 0.0    | 0.06  | 0.116  |
Table 11: Sensitivity Analysis for different prior specifications (Prior 4).

| Estimands | Mean  | SD    | 2.5% | Median | 97.5% | Length |
|-----------|-------|-------|------|--------|-------|--------|
| E-ATE$\pi$ | 0.012 | 0.007 | −0.001 | 0.011 | 0.027 | 0.028  |
| E-ASE$\pi$ | −0.003 | 0.006 | −0.015 | −0.003 | 0.008 | 0.024  |
| E-ASE$^{ab,1}_\pi$ | −0.003 | 0.006 | −0.016 | −0.003 | 0.009 | 0.025  |
| E-ASE$^{ab,2}_\pi$ | −0.002 | 0.018 | −0.036 | −0.004 | 0.033 | 0.069  |
| E-ASE$^{rt,33}_\pi$ | −0.002 | 0.027 | −0.053 | 0.0 | 0.052 | 0.104  |
| E-ASE$^{rt,50}_\pi$ | −0.003 | 0.008 | −0.019 | −0.004 | 0.013 | 0.032  |
| E-ASE$^{rt,66}_\pi$ | −0.001 | 0.029 | −0.058 | 0.0 | 0.039 | 0.117  |

Figure 3: Experimental design (left) and observed deformation (right). Dashed lines on the left figure represent assigned compensations.

D Case Study: Inference on the Degree of Interference in an Experiment on Stereolithography

Additive manufacturing (AM), or three-dimensional (3D) printing, refers to a manufacturing technology in which material is deposited and solidified additively based on a computer-aided design (CAD) model to manufacture the corresponding physical product. A significant challenge with AM technologies is dimensional accuracy control of a printed product. Shape deviations inevitably arise in additively manufactured products due to the physics associated with the rapid heating and cooling in any AM process. Several studies have been performed to investigate different strategies for dimensional accuracy control. One strategy is to implement a compensation plan for shape deviations, which corresponds to a modification of the CAD model such that when the modified model is manufactured the corresponding physical product should exhibit reduced shape deviations compared to the original printed product (Huang et al., 2015).

Sabbaghi et al. (2014) studied shape deviations in a stereolithography process. Their experiment considered in-plane deviations for the top layers of four circular cylinders with negligible heights. The cylinder was of nominal radius 0.5”. The objective of their experiment was to determine whether interference would arise in the shape deviations for the points on the cylinders under discretized compensation plans. The experimental units in this context are the individual
points on the top boundary of a printed product. The covariates for each unit correspond to the location of the unit on a cylinder under the polar coordinate system (i.e., the angle $\psi_i \in (0, 2\pi)$ in radians). The treatment is a continuous treatment factor, corresponding to the amount of compensation (i.e., the addition or subtraction of material) applied to an experimental unit. Each of the potential outcomes $Y_i(Z)$ for unit $i$ is defined as the difference between the potential and nominal radius at $\psi_i$ under compensation plan $Z$ applied to the product containing unit $i$. The experimental design corresponds to a restricted Latin square design on a cylinder. The cylinder was divided into 16 equal-sized sections of $\pi/8$ radians, and one of four levels of the continuous treatment factor was applied to each, with the levels set based on the size of the cylinder. The compensations for the 0.5” cylinder were $-0.004”$, $0.004”$, and $0.008”$. The left panel of Figure 3 summarizes the shape deviation data of all the measured points.

We proceed to apply our DoI framework to analyze the $N = 6159$ datapoints from the cylinder with nominal radius $r = 0.5”$. This application is challenging in the sense that we are interested in the unit-level effects. We first estimate the unit-level assignment-conditional total effects. These are similar in spirit to the effective treatments considered by Sabbaghi et al. (2014). However, a major difference is that Sabbaghi et al. (2014) defined the estimand in terms of the parameters of their Bayesian model whereas we define the estimand according to the finite-sample perspective, i.e., in terms of observed and missing potential outcomes such as $\tau_{i,\text{tot}}(z) = Y_i(z_i, z_{-i}) - Y_i(0, 0)$. The left panel of Figure 4 summarizes the inferences on the total effects obtained from our methodology. These inferences match the effective treatments presented in Figure 5 of Sabbaghi et al. (2014). Considering the fact that their analysis was based on parametric Bayesian models tailored to this particular AM data set, we conclude that our DoI framework and Bayesian approach successfully captures the effective treatment of compensation as the total effect of compensation in a conceptually more straightforward and general manner. In addition, our DDPM is much more flexible than the parametric models considered by Sabbaghi et al. (2014), with the flexibility arising from the infinite mixture of kernels with atoms that depend on $A$. Instead of the common choice of the adjacency matrix, we employed an inverse-distance matrix that was defined as

$$A_{ij} = \begin{cases} \frac{1}{|\psi_i - \psi_j|} & \text{if } |\psi_i - \psi_j| \leq \frac{\pi}{8} \\ 0 & \text{otherwise.} \end{cases}$$

The set of neighboring units for unit $i$ is defined as $N_i = \{ j : A_{ij} \neq 0 \}$. Our potential outcome model was specified according to $Y_i(z) = X_i^T \beta_Z + G_i(T_i, Z_{-i})$, with $\mu(X_i, A_i, Z_{-i}, \gamma_k) = \gamma_k \sum_{j \in N_i} Z_j A_{ij}$. Following Sabbaghi et al. (2014) we set $X_i = (1, \cos \psi_i)^T$ and specify our prior distributions for the model parameters as $\beta_{Z_i} \sim N((0,0)^T, 10I_2)$, where $I_2$ is the $2 \times 2$ identity matrix, for $Z_i \in \{-0.004”, 0, 0.004”, 0.008”\}$, $\gamma_k \sim N(0, 10), \sigma_k^2 \sim IG(1, 10^{-7}), \sigma^2 \sim IG(1, 10^{-7})$.

In addition to the difference in perspectives, our approach possesses additional advantages. First and foremost, we need only use a single data set to estimate the total effect, whereas Sabbaghi et al. (2014) used two distinct sets of experimental data to assess which units on the cylinders have negligible interference. In particular, their first data set consisted of control data from previously manufactured cylinders for which every unit received zero compensation. They computed the posterior distribution of the model parameters and formed the posterior predictive distribution of the potential outcomes based on these control data. After performing these inferences, they then
used the data from the compensation experiment to assess interference in terms of how the units’ effective treatments differed from that to which they were physically assigned. This approach is similar to that of Hudgens and Halloran (2008) in the sense that they prepared multiple clusters of units by their designs and assessed interference by comparing the outcomes in different clusters. In our approach, we only need the data from the compensated products, and not control data on products that did not have a compensation plan, to infer the effects of interference.

As discussed in §3.2, the unit-level spillover effect is \( \tau_{i}^{sp}(Z_i, Z_{-i}, Z'_{-i}) = Y_i(Z_i, Z_{-i}) - Y_i(Z_i, Z'_{-i}) \) for two different assignment vectors \( Z \) and \( Z' \) with \( Z_i = Z'_i \). Following the definition of negligible interference by Sabbaghi et al. (2014, p. 1403, 1405), we consider spillover effects in which, for a given \( Z \), we set \( Z' = Z_i1_{N-1} \), where \( 1_{N-1} \) is the \((N-1)\)-dimensional vector of ones. Our unit-level spillover estimand in this case is formally expressed as \( \tau_{i}^{sp}(Z_i, Z_{-i}, Z'_{-i}) = Y_i(Z_i, Z_{-i}) - Y_i(Z_i, Z_i1_{N-1}) \). The right panel of Figure 4 presents the spillover effect of all units as inferred from our Bayesian method. The central 95% posterior intervals for significantly positive (negative) units are always positive (negative), and never contain zero, which indicates that these units receive nonzero spillover effects. We observe that our Bayesian method indicates that approximately 90% of units, primarily in the central regions of the 16 sections of the cylinders, exhibit negligible interference. This result is consistent with that obtained by Sabbaghi et al. (2014), and was obtained in a more general and straightforward manner compared to Sabbaghi et al. (2014).

Another advantage of our Bayesian method compared to the work of Sabbaghi et al. (2014) is the clustering property of the DDPM. Figure 5 summarizes the DoI of all experimental units as inferred from our model, with different colors representing different clusters of units. The clusters of units are determined by the posterior modes of the latent class indicators \( C_i \). Different clusters exhibit different interference structures, which are governed by the atoms of the DDPM model. As seen in this figure, most of the units in the middle of the sections belong to one major cluster (Cluster 6, indicated as green). Units on the edges of the sections belong to other minor clusters and exhibit more significant interference, which corresponds to the fact that in this experiment these units should have different interference structures compared to that of the major cluster.

Each atom of the DDPM model has its own covariate-dependent location parameter, so that each component corresponds to a single parametric model with a different location parameter. As we obtain multiple clusters, it is clear that modeling the interference structure with a single parametric model, as was done by Sabbaghi et al. (2014), is not appropriate. Our methodology automatically captures the interference structure and separates the units into different clusters based on the interference effects that they receive. Of the total \( N = 6159 \) units, 4458 (or approx-
Figure 5: Summary of the DoIs for all points on the 0.5" cylinder.

approximately 70%) belong to the major cluster. Clusters 3, 4 and 8 appear only on the edge of the sections in which the absolute difference in their compensation level versus that of the neighboring section is 0.008". As such, units in clusters 3, 4 and 8 are highly affected by other units through interference. Finally, we see that the cluster membership changes depending on the distance of a unit from the edge of the section, which gradates the interference structures within sections.