On randomization-based causal inference for matched-pair factorial designs

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Abstract

Under the potential outcomes framework, we introduce matched-pair factorial designs, and propose the matched-pair estimator of the factorial effects. We also calculate the randomization-based covariance matrix of the matched-pair estimator, and provide the “Neymanian” estimator of the covariance matrix.

Keywords: Experimental design; factorial effect; precision; potential outcome.

1. INTRODUCTION

Randomization is widely regarded as the gold standard of causal inference (Rubin 2008). Under the potential outcomes framework (Neyman 1923; Rubin 1974), for a two-level factor, we define the causal effect as the linear contrast of the potential outcomes under treatment and control. To investigate multiple factors simultaneously, $2^K$ factorial designs (Fisher 1935; Yates 1937) can be employed. Randomization-based casual inference for factorial designs has deep roots in the experimental design literature (e.g., Kempthrone 1952), and was recently presented using the language of potential outcomes (Dasgupta et al. 2015; Mukerjee et al. 2016).

Pair-matching (Cochran 1953), as a special form of stratification, has been widely adopted by researchers and practitioners (e.g., Grossarth-Maticke and Ziegler 2008). For treatment-control

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studies (i.e., $2^1$ factorial designs), pair-matching has been extensively investigated by the causal inference community (Rosenbaum 2002; Imai 2008; Imai et al. 2009; Ding 2016; Fogarty 2016a,b). Unfortunately, similar discussion appears to be missing for general factorial designs. In this paper, we fill this theoretical gap by extending Imai (2008)’s analysis to matched-pair factorial designs. We restrict the experimental units to be a fixed finite population, for a two-fold reason. First, as shown in Imai (2008), it is straightforward to generalize the finite-population analyses to infinite populations. Second, for some practical examples, it might be unreasonable to view the experimental units as a random sample from an infinite population.

The paper proceeds as follows. Section 2 reviews the randomization-based causal inference framework for completely randomized factorial designs. Section 3 introduces matched-pair factorial designs, proposes the matched-pair estimator for the factorial effects, calculates its covariance matrix and the corresponding estimator. Section 4 briefly discusses the precision gains by pair-matching in factorial designs, and concludes.

2. CAUSAL INFERENCE FOR COMPLETELY RANDOMIZED FACTORIAL DESIGNS

To ensure self-containment, we first review the randomization-based causal inference framework for completely randomized factorial designs. Although most materials are adapted from Dasgupta et al. (2015) and Lu (2016a,b), some are refined for better clarity. For more detailed discussions on factorial designs, see, e.g., Wu and Hamada (2009).

2.1. Factorial designs

A $2^K$ factorial design consists of $K$ two-level (coded $-1$ and $+1$) factors. We represent it by the corresponding model matrix (Wu and Hamada 2009), a $2^K \times 2^K$ matrix $H_K = (h_0, \ldots, h_{2^K-1})$ that can be constructed as follows:

1. Let $h_0 = 1_{2^K}$;

2. For $k = 1, \ldots, K$, construct $h_k$ by letting its first $2^{K-k}$ entries be $-1$, the next $2^{K-k}$ entries be $+1$, and repeating $2^{k-1}$ times;
3. If $K \geq 2$, order all subsets of $\{1, \ldots, K\}$ with at least two elements, first by cardinality and then lexicography. For $k = 1, \ldots, 2^K - K - 1$, let $\sigma_k$ be the $k$th subset and $h_{K+k} = \prod_{l \in \sigma_k} h_l$, where “$\prod$” stands for entry-wise product.

The use of the constructed $H_K$ is two-fold:

1. $h_0$ corresponds to the null effect; $h_1$ to $h_K$ correspond to the main effects of the $K$ factors; $h_{K+1}$ to $h_{K+(K/2)}$ correspond to the two-way interactions; \ldots; $h_{2K-1}$ corresponds to the $K$-way interaction;

2. The $j$th row of $(h_1, \ldots, h_K)$ corresponds to the $j$th treatment combination $z_j$.

For $j = 1, \ldots, 2^K$, let $\lambda_j$ denote the $j$th row of $H_K$.

**Example 1.** For $2^2$ factorial designs, the model matrix is:

$$H_2 = \begin{pmatrix}
    h_0 & h_1 & h_2 & h_3 \\
    \lambda_1 & +1 & -1 & -1 & +1 \\
    \lambda_2 & +1 & -1 & +1 & -1 \\
    \lambda_3 & +1 & +1 & -1 & -1 \\
    \lambda_4 & +1 & +1 & +1 & +1
\end{pmatrix}.$$  

The four treatment combinations are $z_1 = (-1, -1)$, $z_2 = (-1, +1)$, $z_3 = (+1, -1)$ and $z_4 = (+1, +1)$. We represent the main effects of factors 1 and 2 by $h_1 = (-1, -1, +1, +1)'$ and $h_2 = (-1, +1, -1, +1)'$ respectively, and the two-way interaction by $h_3 = (+1, -1, -1, +1)'$.

### 2.2. Randomization-based causal inference

We consider a $2^K$ factorial design with $N = 2^K r$ units. By invoking the Stable Unit Treatment Value Assumption [Rubin 1980], for $i = 1, \ldots, N$ and $l = 1, \ldots, 2^K$, let the potential outcome of unit $i$ under $z_l$ be $Y_i(z_l)$, the average potential outcome for $z_l$ be $\bar{Y}(z_l) = N^{-1} \sum_{i=1}^{N} Y_i(z_l)$, and $Y_i = \{Y_i(z_1), \ldots, Y_i(z_{2^K})\}'$. Define the individual and population-level factorial effect vectors as

$$\tau_i = \frac{1}{2^{K-1}} H_K' Y_i \quad (i = 1, \ldots, N); \quad \tau = \frac{1}{N} \sum_{i=1}^{N} \tau_i,$$

(1)
respectively. Our interest lies in $\tau$.

We denote the treatment assignment mechanism by

$$W_i(z_l) = \begin{cases} 
1, & \text{if unit } i \text{ is assigned treatment } z_l, \\
0, & \text{otherwise.}
\end{cases}$$

($i = 1,\ldots,N; l = 1,\ldots,2^K$).

We impose the following restrictions on the treatment assignment mechanism:

$$\sum_{l=1}^{2^K} W_i(z_l) = 1 \quad (i = 1,\ldots,N); \quad \sum_{i=1}^{N} W_i(z_l) = r \quad (l = 1,\ldots,2^K).$$

In other words, we assign $r$ units to each treatment, and one treatment to each unit. Therefore, the observed outcome of unit $i$ is $Y^\text{obs}_i = \sum_{l=1}^{2^K} W_i(z_l)Y_i(z_l)$, and the average observed outcome for treatment $z_l$ is $\bar{Y}^\text{obs}(z_l) = r^{-1} \sum_{i=1}^{N} W_i(z_l)Y_i(z_l)$. Under complete randomization, Dasgupta et al. (2015) estimated $\tau$ by

$$\hat{\tau}_C = 2^{-(K-1)} H_K \tilde{Y}^\text{obs}, \quad \tilde{Y}^\text{obs} = \{\tilde{Y}^\text{obs}(z_1),\ldots,\tilde{Y}^\text{obs}(z_{2^K})\}'.$$

The sole source of randomness of $\hat{\tau}_C$ is the treatment assignment. Dasgupta et al. (2015) and Liu (2016b) derived the covariance matrix of this estimator, and the “Neymanian” estimator of the covariance matrix. We summarize their main results in the following lemmas.

**Lemma 1.** $\hat{\tau}_C$ is unbiased, and its covariance matrix is

$$\text{Cov}(\hat{\tau}_C) = \frac{1}{2^{2(K-1)}r} \sum_{l=1}^{2^K} \lambda_l \chi_l \left[ \frac{1}{N-1} \sum_{i=1}^{N} \{Y_i(z_l) - \bar{Y}(z_l)\}^2 - \frac{1}{N(N-1)} \sum_{i=1}^{N} (\tau_i - \bar{\tau})(\tau_i - \bar{\tau})' \right] s^2(z_l).$$

Moreover, the “Neymanian” estimator of the covariance matrix is

$$\widehat{\text{Cov}}(\hat{\tau}_C) = \frac{1}{2^{2(K-1)}r} \sum_{l=1}^{2^K} \lambda_l \chi_l \left[ \frac{1}{r-1} \sum_{i=1}^{N} W_i(z_l) \{Y^\text{obs}_i - \bar{Y}^\text{obs}(z_l)\}^2 \right] s^2(z_l),$$

whose bias is $\sum_{i=1}^{N} (\tau_i - \bar{\tau})(\tau_i - \bar{\tau})'/(N^2 - N)$. 

The covariance matrix estimator $\hat{\Cov}(\hat{\tau}_C)$ is “conservative,” because its diagonal entries, i.e., the variance estimators of the components of $\hat{\tau}_C$, have non-negative biases.

3. CAUSAL INFERENCE FOR MATCHED-PAIR RANDOMIZED FACTORIAL DESIGNS

3.1. Matched-pair designs and causal parameters

As pointed out by Imai (2008), the key idea behind matched-pair designs is that “experimental units are paired based on their pre-treatment characteristics and the randomization of treatment is subsequently conducted within each matched pair.” To apply this idea to factorial designs, we group the $N$ experimental units into $r$ “pairs” of $2^K$ units, and within each pair randomly assign one unit to each treatment. Let $\psi_j$ be the set of indices of the units in pair $j$, such that

$$|\psi_j| = 2^K \quad (j = 1, \ldots, r); \quad \psi_j \cap \psi_{j'} = \emptyset \quad (\forall j \neq j'); \quad \bigcup_{j=1}^r \psi_j = \{1, \ldots, N\}.$$  

For pair $j$, denote the average outcomes for treatment $z_l$ as $\bar{Y}_j(z_l) = 2^{-K} \sum_{i \in \psi_j} Y_i(z_l)$, and $\bar{Y}_j = \{\bar{Y}_j(z_1), \ldots, \bar{Y}_j(z_{2^K})\}'$, and the factorial effect vector as $\tau_j = 2^{-(K-1)} H_K' \bar{Y}_j$. It is apparent

$$\frac{1}{r} \sum_{j=1}^r \bar{Y}_j(z_l) = \bar{Y}(z_l) \quad (l = 1, \ldots, 2^K); \quad \frac{1}{r} \sum_{j=1}^r \tau_j = \tau.$$

Within each pair, we randomly assign one unit to each treatment. Let the observed outcome of treatment $z_l$ in pair $j$ be $Y_{j}^\text{obs}(z_l) = \sum_{i \in \psi_j} Y_i(z_l)W_i(z_l)$, and $Y_j^\text{obs} = \{Y_j^\text{obs}(z_1), \ldots, Y_j^\text{obs}(z_{2^K})\}'$. We estimate $\tau_j$ by $\hat{\tau}_j = 2^{-(K-1)} H_K' Y_j^\text{obs}$. The matched-pair estimator for $\tau$ is

$$\hat{\tau}_M = \frac{1}{r} \sum_{j=1}^r \hat{\tau}_j.$$  \hspace{1cm} (3)

3.2. Randomization-based inference

We now present the main results of this paper.
Proposition 1. \( \hat{\tau}_M \) is an unbiased estimator of \( \tau \), and its covariance matrix is

\[
\text{Cov}(\hat{\tau}_M) = \frac{1}{2^{2(K-1)r^2}} \sum_{l=1}^{2^K} \lambda'_l \lambda_l \Delta_l - \frac{1}{2^K (2^K - 1)r^2} \Sigma,
\]

where

\[
\Delta_l = \frac{1}{2^K - 1} \left[ (N - 1)S^2(z_l) - 2^K \sum_{j=1}^r \{ \bar{Y}_j(z_l) - \bar{Y}(z_l) \}^2 \right] (l = 1, \ldots, 2^K),
\]

and

\[
\Sigma = \sum_{i=1}^N (\tau_i - \tau)(\tau_i - \tau)' - 2^K \sum_{j=1}^r (\tau_j - \tau)(\tau_j - \tau)'.
\]

Proof. To prove the first part, note that \( \hat{\tau}_j \) is an unbiased estimator of \( \tau_j \), for \( j = 1, \ldots, r \). This fact combined with (3) completes the proof.

To prove the second part, let \( W_j = \{W_i(z_l)\}_{i \in \psi_j, l=1,\ldots,2^K} \) denote the treatment assignment for pair \( j \). By definition, \( W_j \)'s are independently and identically distributed, implying the (joint) independence of \( \hat{\tau}_j \)'s. Consequently, we can treat each pair as a completely randomized factorial design with \( 2^K \) units. Therefore by Lemma [4],

\[
\text{Cov}(\hat{\tau}_j) = \frac{1}{2^{2(K-1)r^2}} \sum_{l=1}^{2^K} \lambda'_l \lambda_l \frac{1}{2^K - 1} \sum_{i \in \psi_j} (Y_i(z_l) - \bar{Y}_j(z_l))^2 - \frac{1}{2^K (2^K - 1)r^2} \sum_{i \in \psi_j} (\tau_i - \tau_j)(\tau_i - \tau_j)'.
\]

This implies that

\[
\text{Cov}(\hat{\tau}_M) = \frac{1}{r^2} \sum_{j=1}^r \text{Cov}(\hat{\tau}_j) = \frac{1}{2^{2(K-1)r^2}} \sum_{l=1}^{2^K} \lambda'_l \lambda_l \sum_{j=1}^r S^2_j(z_l) - \frac{1}{2^K (2^K - 1)r^2} \sum_{j=1}^r \sum_{i \in \psi_j} (\tau_i - \tau_j)(\tau_i - \tau_j)'.
\]

To prove the equivalence between (4) and (5), simply note that

\[
(2^K - 1) \sum_{j=1}^r S^2_j(z_l) + 2^K \sum_{j=1}^r \{ \bar{Y}_j(z_l) - \bar{Y}(z_l) \}^2 = (N - 1)S^2(z_l)
\]
and
\[ \sum_{j=1}^{r} \sum_{i \in \psi_j} (\tau_i - \tau_j)(\tau_i - \tau_j)' + 2^K \sum_{j=1}^{r} (\tau_j - \tau)(\tau_j - \tau)' = \sum_{i=1}^{N} (\tau_i - \tau)(\tau_i - \tau)' . \]

The proof is complete. \( \square \)

We discuss a special case before moving forward. When \( K = 1 \), we have the classic treatment-control studies, and label the treatment and control as +1 and −1, respectively. We are interested in the difference-in-mean estimator
\[ \hat{\tau}_{MP} = \frac{1}{r} \sum_{j=1}^{r} \{ Y_{j1}^{\text{obs}}(+1) - Y_{j2}^{\text{obs}}(-1) \} . \]

Denote \( \psi_j = \{ j_1, j_2 \} \). Imai (2008) (p. 4861, Eq. (8)) derived the variance of \( \hat{\tau}_{MP} \) as
\[ \text{Var}(\hat{\tau}_{MP}) = \frac{1}{4r^2} \sum_{j=1}^{r} \{ Y_{j1}(+1) - Y_{j2}(-1) - Y_{j2}(+1) + Y_{j1}(-1) \}^2 . \] (6)

As a validity check, Proposition 1 reduces to (6) when \( K = 1 \). We leave the proof to the readers.

We discuss the estimation of \( \text{Cov}(\hat{\tau}_{M}) \), because Lemma 1 does not apply for matched-pair factorial designs. Inspired by Imai (2008), we propose the following estimator:
\[ \hat{\text{Cov}}(\hat{\tau}_{M}) = \frac{1}{r(r-1)} \sum_{j=1}^{r} (\hat{\tau}_j - \hat{\tau}_{M})(\hat{\tau}_j - \hat{\tau}_{M})' . \] (7)

**Proposition 2.** The bias of the covariance estimator in (7) is
\[ E \{ \hat{\text{Cov}}(\hat{\tau}_{M}) \} - \text{Cov}(\hat{\tau}_{M}) = \frac{1}{r(r-1)} \sum_{j=1}^{r} (\tau_j - \tau)(\tau_j - \tau)' . \]

**Proof.** The proof is a basic maneuver of the expectation and covariance operators. First, by (3) and the joint independence of \( \hat{\tau}_j \)'s,
\[ \text{Cov}(\hat{\tau}_{M}) = r^{-2} \sum_{j=1}^{r} \text{Cov}(\hat{\tau}_j) . \]
Therefore by (7),

$$r(r - 1)E \left\{ \hat{\text{Cov}}(\hat{\tau}_M) \right\} = \sum_{j=1}^{r} E(\hat{\tau}_j \hat{\tau}_j') - rE(\hat{\tau}_M \hat{\tau}_M')$$

$$= \sum_{j=1}^{r} \text{Cov}(\hat{\tau}_j) + \sum_{j=1}^{r} \tau_j \tau_j' - r\text{Cov}(\hat{\tau}_M) - r\tau \tau'$$

$$= r(r - 1)\text{Cov}(\hat{\tau}_M) + \sum_{j=1}^{r} (\tau_j - \tau)(\tau_j - \tau)' .$$

Proposition 2 implies that the estimator of Cov(\(\hat{\tau}_M\)) is also “conservative.” We leave it to the readers to prove that for treatment-control studies, Proposition 2 reduces to the corresponding results in Imai (2008) (p. 4862, Prop. 2, Part 1).

4. DISCUSSIONS AND CONCLUDING REMARKS

For treatment-control studies, Imai (2008) compared the variance formulas for the complete-randomization and matched-pair estimators, and derived the condition under which pair-matching leads to precision gains. For general factorial designs, analogous comparisons can be made between (2) and (4). However, to our best knowledge, intuitive closed-form expressions might not be available without additional assumptions on the potential outcomes.

There are multiple future directions based on our current work. First, we may compare the precisions of the complete-randomization and matched-pair estimators under certain mild restrictions on the potential outcomes. Second, it is possible to unify the randomization-based and regression-based inference frameworks, as pointed out by Samii and Aronow (2012) and Lu (2016b). Third, additional pre-treatment covariates may shed light on the pair-matching mechanism, and help sharpen our current analysis.

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