

1 **Nonparametric Methods for the Nondecreasing**
2 **Ordered Hypothesis in a Mixed Design**

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ABSTRACT

Aims: Introduce two new test statistics in testing for the nondecreasing alternative in a mixed design consisting of a Completely Randomized Portion and a Randomized Complete Block Portion.

Study design: Simulation study comparing four test statistics for the nondecreasing alternative in a mixed design consisting of a CRD and an RCBD portion. The test statistics included two new test statistics and two existing test statistics. Random samples were taken from three different types of underlying distributions. Different percentages of the CRD portion will be considered as well as different sample sizes. Powers were estimated based on a variety location parameter shifts. Three, four, and five populations were considered.

Place and Duration of Study: The simulation study took place on the campus of North Dakota State University during the calendar year 2019.

Methodology: Levels of significance for each of the three types of underlying distributions, when the RCBD portion was larger than the CRD portion, when the CRD portion was larger than the RCBD portion, and when the CRD portion was equal to the RCBD portion, and when the number of populations were 3, 4, and 5.

Results: Regardless of the underlying population types, the proposed test statistics did better than the existing test statistics when the difference between the last two parameters is large. This was true for 3, 4, and 5 populations.

Conclusion: When the differences between the last two parameters is large, the two new test statistics performed better. Otherwise, the existing test statistics are better. In both cases, it is better to use the combined test statistic that first standardizes the individual test statistics for the CRD and RCBD portions before adding them together.

22

23 *Keywords:* nonparametric; order-restricted inference; JonckheereTerpstra -;
24 completely randomized design; randomized complete block design; mixed design.

25 1. INTRODUCTION

26

27 In some experimental studies, researchers may wish to test the null hypothesis that there
28 are no differences among treatment effects against an ordered alternative hypothesis.
29 Researchers may be able to assume the treatments follow an a priori ordering, if they are
30 different. The scope of this paper focusses on the nondecreasing ordered alternative
31 hypothesis. That is,

32 $H_0: \mu_1 = \mu_2 = \dots = \mu_k$ versus $H_1: \mu_1 \leq \mu_2 \leq \dots \leq \mu_k$

33 where at least one inequality is strict and μ_i denotes the location parameter of the i^{th}
34 treatment. It is also possible that researchers may start out conducting an experiment using
35 a randomized complete block design, but realize later that the design becomes too
36 expensive and too hard to continue without several observations missing on treatments
37 within a block (14,16). For instance, let us suppose a large company is interested in
38 controlling the increasing cost of insurance of its employees. Hence, the company introduces
39 a wellness program in an attempt to improve the overall health by reducing the average
40 cholesterol level (*LDL*) of its employees. To test the competence of the program, the

41 company starts measuring the cholesterol levels of random employees' samples – after their
 42 consent – three times in two years: at the beginning of the program and two times, annually,
 43 afterward (13,15). However, because the company is large and has an annual turnover of
 44 nearly 18% in its employees, many observations become obsolete because their donor
 45 employees left the company before completing them; therefore, their cholesterol level cannot
 46 be obtained anymore. To counter this problem, the company decides to discard
 47 observations that were incomplete for at least one year and perform a test using only a
 48 randomized complete block design (*RCBD*) to get accurate results on cholesterol levels over
 49 two years.

50
 51 The company realizes that it loses a lot of data by sticking to a randomized complete block
 52 design (*RCBD*) which, in turn, hampers its effort to enhance its employees' general health.
 53 Thus, the company comes up with an idea to take advantage of the leftover observations
 54 that do not constitute a complete block and collect more observations over the next two
 55 years using a completely randomized design (*CRD*). Therefore, statistical test is required
 56 which combines the observations from the randomized complete block with those from the
 57 completely randomized design.

58
 59 Page (1963)[1] proposed a nonparametric procedure that is applicable for testing
 60 nondecreasing ordered alternative when the data fit the two-way analysis of variance
 61 structure. (Daniel, 1990) [2] mentioned several assumptions for the validity of this test,
 62 including the independency of blocks, and no interaction between the blocks and the
 63 treatments. The test statistic is defined as

$$L = \sum_{j=1}^k jR_j \quad (1)$$

64 where R_j is the j^{th} treatment rank sum, based on the within block ranks of the original
 65 observations. Under H_0 , the statistic L has an asymptotic normal distribution with mean and
 66 variance, $bk(k+1)^2/4$, $b(k^3-k)^2/144(k-1)$, respectively. The standardized version of
 67 the statistic L is defined as

$$Z_p = \frac{L - [bk(k+1)^2/4]}{\sqrt{b(k^3-k)^2/144(k-1)}} \quad (2)$$

68 where b denotes the number of the blocks and k denotes the number of treatments. Under
 69 H_0 , the statistic Z_p follows an asymptotic standard normal distribution (i.e., $N(0,1)$) and so the
 70 standard normal table can be used to get the critical values.

71 Jonckheere-Terpstra test (referred to as *JT*) is a widely known nonparametric test for the
 72 nondecreasing ordered alternatives in the k -sample case when the design is a completely
 73 randomized. The test was proposed independently by Terpstra (1952) [3] and Jonckheere
 74 (1954) [4]. For this test, the samples must be independent, and each sample is assumed to
 75 be drawn from a continuous population in which the distribution is the same for each
 76 population and may only differ in the location parameters. The test statistic is based on the
 77 summation of $k(k-1)/2$ Mann-Whitney statistics. Namely, it can be expressed as

$$JT = \sum_{i=1}^{k-1} \sum_{j=i+1}^k U_{ij} \quad (3)$$

78 where U_{ij} is the U-statistics of Mann-Whitney and is defined as the number of pairs of
 79 observations (X_{ia}, X_{ib}) in which X_{ia} is less than X_{ib} , once more, X_{ia} is the a^{th} observation in
 80 i^{th} treatment sample, $a = 1, 2, \dots, n_i$ and X_{ib} is the b^{th} observation in j^{th} treatment sample,
 81 $b = 1, 2, \dots, n_j$. Under the null hypothesis, H_0 , the *JT* statistic follows an asymptotic standard
 82 normal distribution with mean

$$E_0(JT) = \frac{N^2 - \sum_{i=1}^k n_i^2}{4} \quad (4)$$

83 and variance

$$Var_0(JT) = \frac{N^2(2N + 3) - \sum_{i=1}^k n_i^2(2n_i + 3)}{72} \quad (5)$$

84 where $N = \sum_{i=1}^k n_i$, and n_i denotes the sample size of the i^{th} treatment. However, the
 85 asymptotic normality of JT depends on the number of samples. Jonckheere (1954) [4]
 86 mentioned that the normality approximation might be inaccurate if only one n_i tends to
 87 infinity as N increases. Therefore, to achieve the normality approximation, at least two
 88 samples increase as N goes to infinity. Therefore, the standardized version of the test
 89 statistic, JT , is defined as

$$Z_{JT} = \frac{JT - E_0(JT)}{\sqrt{Var_0(JT)}} \quad (6)$$

90

91 Tryon and Hettmansperger(1973) [5] introduced a modified version of the JT test, MJT . Both
 92 tests, JT and MJT , are dealing with the nondecreasing ordered alternatives. However, the
 93 MJT test assigns more weights for each Mann-Whitney statistic based on the distance
 94 between the i^{th} and j^{th} populations. Thus, if the distance between the two populations is
 95 considered to be large, more amount of weight will be assigned to each Mann-Whitney
 96 statistic. The test statistic can be written as follows:

$$MJT = \sum_{i=1}^{k-1} \sum_{j=i+1}^k (j-i)U_{ij} \quad (7)$$

97

98 Under H_0 , the asymptotic distribution for the test statistic follows a normal distribution with
 99 mean

$$E_0(MJT) = \sum_{i=1}^{k-1} \sum_{j=i+1}^k (j-i) \frac{n_i n_j}{2} \quad (8)$$

100 and variance

$$\begin{aligned} Var_0(MJT) &= Var \left\{ \sum_{i=1}^{k-1} \sum_{j=i+1}^k (j-i)U_{ij} \right\} \\ &= \sum_{i=1}^{k-1} \sum_{j=i+1}^k (j-i)^2 Var(U_{ij}) + 2 \sum_{i=1}^{k-1} \sum_{j=i+1}^k \sum_{s=1}^{k-1} \sum_{t=s+1}^k (j-i)(t-s) Cov(U_{ij}, U_{st}) \end{aligned} \quad (9)$$

101 where the values of the variance and the covariance terms can be defined as in Hollander
 102 and Wolfe (1999) [6] where

$$Var(U_{ij}) = \frac{n_i n_j (n_i + n_j + 1)}{12} \quad \text{for } 1 \leq i \leq j \leq k \quad (10)$$

103

$$Cov(U_{ij}, U_{pq}) = \begin{cases} \frac{n_i n_j n_q}{12} & \text{if } 1 \leq i, p \leq j \leq k, j \neq q \\ -\frac{n_i n_j n_p}{12} & \text{if } 1 \leq p < j \leq k, i = q \\ -\frac{n_i n_j n_q}{12} & \text{if } 1 \leq i < j < q \leq k, j = p \\ \frac{n_i n_j n_p}{12} & \text{if } 1 \leq i, p \leq j \leq k, i \neq p \\ 0 & \text{if } i, j, p, q \text{ are different} \end{cases} \quad (11)$$

104

105 Neuhäuser *et al.* (1998) [7] illustrated that in situations when the sample sizes are relatively
106 small, the modified Jonckheere-Terpstra (*MJT*) has higher power than the original
107 Jonckheere-Terpstra (*JT*).

108

109 Terpstra and Magel (2003) [8] introduced a new nonparametric test for testing the
110 nondecreasing ordered alternative that does not depend on pairwise information. Instead, it
111 depends on the information that is obtained across all samples at the same time. The test
112 statistic is given as follows

$$TM = \sum_{i_1=1}^{n_i} \dots \sum_{i_k=1}^{n_k} I(X_{1i_1} \leq X_{2i_2} \leq \dots \leq X_{ki_k}) \quad (12)$$

113 where $I(X_{1i_1} \leq X_{2i_2} \leq \dots \leq X_{ki_k})$ is the indicator function that is equal to one only when there
114 is at least one strict inequality and zero otherwise. Terpstra and Magel (2003) [8] compared
115 their test to the *JT*, and the *MJT* tests. The results indicated that the proposed test has fairly
116 higher power when the priori ordering is correct. However, if it is the other way around, the
117 proposed test can have smaller power than the *JT*, and the *MJT* tests.

118

119 Ferdhiana *et al.* (2008) [9] proposed a test that is similar to the test proposed by Terpstra and
120 Magel (2003) [8], with the exception of function. Ferdhiana *et al.* (2008) [9] used the Kendall's
121 Tau correlation coefficient instead of the indicator function used in the *TM* test.

122

123 Moreover, another nonparametric test is given by Terpstra *et al.* (2011) [10] that is analogous
124 test to the *TM*; however, in this test the indicator function in Equation (12) was replaced by
125 Spearman's correlation coefficient. The result of both tests indicate that the proposed test
126 has higher power than the *JT*, the *MJT* and the *TM* when the sample sizes are small with
127 large shift between the two adjacent location parameters.

128

129 Magele *et al.* (2009 [11]) developed two tests for the nondecreasing ordered alternatives in
130 mixed design. Part of the design is considered a data from a randomized complete block
131 design (RCBD) and the other part is considered a data from a complete randomized design
132 (CRD). The tests in [11] are a mixture of Page's test and Jonckheere-Terpstra test. The first
133 test in [11], can be written as follows:

$$C_1 = \frac{Z_p + Z_{JT}}{\sqrt{2}} \quad (13)$$

134

135 Under H_0 , C_1 follows a standard normal distribution since the standardized version of Page's
136 test, and the standardized version of Jonckheere-Terpstra, Z_p, Z_{JT} , respectively, follow a
137 standard normal distribution. Thus, the null hypothesis will be rejected when $C_1 \geq Z_\alpha$
where Z_α is the upper α quantile of the standard normal distribution.

138

139 The second test in Magel et al. (2009) [11] is based on the idea that the Page's test and
 140 Jonckheere-Terpstratest were added together first and then standardized. That is,

$$C_2 = \frac{L + JT - E(0)}{\sqrt{Var(0)}} \quad (14)$$

141 where

$$E(0) = \frac{bk(k+1)^2 + (N^2 - \sum_{i=1}^k n_i^2)}{4} \quad (15)$$

142 and

$$Var(0) = \frac{b(k^3 - k)^2}{144(k-1)} + \frac{N^2(2N+3) - \sum_{i=1}^k n_i^2(2n_i+3)}{72} \quad (16)$$

143 It can be noted that the mean and variance are the sum of Page's test and the Jonckheere-

144 Terpstra test. More specifically, $\frac{bk(k+1)^2}{4}$ denotes the mean of Page's test and $\frac{(N^2 - \sum_{i=1}^k n_i^2)}{4}$

145 denotes the mean of the Jonckheere-Terpstra. Similarly, $\frac{b(k^3 - k)^2}{144(k-1)}$ and $\frac{N^2(2N+3) - \sum_{i=1}^k n_i^2(2n_i+3)}{72}$

146 denote the variance of Page test (L) and the variance of the Jonckheere-Terpstra test (JT),
 147 respectively. The null hypothesis would again be rejected if $C_2 \geq Z_{\alpha}$.

148

149 2. PROPOSED TESTS AND SIMULATION STUDY

150

151 The Jonckheere-Terpstra (JT) test and some modifications of the JT test already exist for the
 152 completely randomized design (CRD). However, we need to propose new versions of the JT
 153 test for the randomized complete block design ($RCBD$) portion of the design.

154

155 2.1. First proposed method

156

157 In this proposed method, we are applying the idea discussed by Tryon and Hettmansperger
 158 (1973) [5] where the distance between the two populations is multiplied by the i^{th} population.
 159 That is,

$$NMJT = \sum_{i=1}^{k-1} \sum_{j=i+1}^k i(j-i)U_{ij} \quad (17)$$

160

161 Since we are dealing with a mixed design, the standardized version of the test that will be
 162 applied to the CRD portion is written as

$$Z_{NMJT} = \frac{NMJT - E(NMJT)}{\sqrt{Var(NMJT)}} \quad (18)$$

163 where $NMJT$ is defined as the unstandardized version of the new modified Jonckheere-
 164 Terpstra test with mean

$$E(NMJT) = \sum_{i=1}^{k-1} \sum_{j=i+1}^k i(j-i) \frac{n_i n_j}{2} \quad (19)$$

165 and variance

$$\begin{aligned} Var(NMJT) &= Var\left(\sum_{i=1}^{k-1} \sum_{j=i+1}^k i(j-i)U_{ij}\right) \\ &= \sum_{i=1}^{k-1} \sum_{j=i+1}^k i^2(j-i)^2 Var(U_{ij}) + 2 \sum_{i=1}^{k-1} \sum_{j=i+1}^k \sum_{s=1}^{k-1} \sum_{t=s+1}^k (i.s)(j-i)(t-s) Cov(U_{ij}, U_{st}) \end{aligned} \quad (20)$$

166 Moreover, the values of the variance and the covariance terms can be obtained using
 167 Equation (10) and (11). Under H_0 , as $\min(n_1, n_2, \dots, n_k)$ tends to infinity,

$$Z_{NMJT} \xrightarrow{D} N(0,1)$$

168

169 As for the *RCBD* portion, a test is designed by applying the *NMJT* test to each block (i.e.,
 170 $NMJT_1, NMJT_2, \dots, NMJT_b$; $l = 1, 2, \dots, b$) where $NMJT_1$ denotes the new version of the
 171 modified Jonckheere-Terpstra test for the first block, $NMJT_2$ denotes the new version of the
 172 modified Jonckheere-Terpstra test for the second block, and so on. Then, we sum up all the
 173 *NMJT* tests together to form the *BNMJT* test. That is,

$$BNMJT = \sum_{l=1}^b NMJT_l \quad (21)$$

174 where b is the number of blocks. Corresponding to equation (19) and since we are
 175 considering *one* observation per block-treatment, the mean of *BNMJT* test can then be
 176 written as

$$E(BNMJT) = \sum_{l=1}^b E(NMJT_l) = \sum_{l=1}^b \left(\sum_{i=1}^{k-1} \sum_{j=i+1}^k \frac{i(j-i)}{2} \right) \quad (22)$$

177 In like manner, the variance of *BNMJT* test can be defined as

$$\begin{aligned} Var(BNMJT) &= \sum_{l=1}^b Var(NMJT_l) \\ &= \sum_{l=1}^b \left(3 \sum_{i=1}^{k-1} \sum_{j=i+1}^k \frac{i^2(j-i)^2}{12} + 2 \sum_{i=1}^{k-1} \sum_{j=i+1}^k \sum_{s=1}^{k-1} \sum_{t=s+1}^k (i,s)(j-i)(t-s) Cov(U_{ij}, U_{st}) \right) \end{aligned} \quad (23)$$

178 where b is the number of blocks and $NMJT_l$ is the new (multiplied) modified Jonckheere-
 179 Terpstra test for the l^{th} block. Moreover, the covariance term can be obtained from Equation
 180 (11). The standardized version of *BNMJT* test is written as

$$Z_{BNMJT} = \frac{BNMJT - E(BNMJT)}{\sqrt{Var(BNMJT)}} \quad (24)$$

181 where *BNMJT* is defined as the unstandardized version of the summation of the new modified
 182 Jonckheere-Terpstra test for the entire blocks. Under H_0 , as $\min(n_1, n_2, \dots, n_k)$ tends to infinity,

$$Z_{BNMJT} \xrightarrow{D} N(0,1)$$

183

184 Therefore, the first proposed method for the nondecreasing ordered alternatives in a mixed
 185 design is written as follows:

$$T_1 = \frac{Z_{NMJT} + Z_{BNMJT}}{\sqrt{2}} \quad (25)$$

186 Here, we added the standardized version of *NMJT* and *BNMJT* together first, and then we
 187 standardized the two tests by subtracting the means and divided by the standard deviations.
 188 Under H_0 , for large sample sizes, this method will follow an asymptotic normal distribution
 189 since it is a combination of two tests which follow a standard normal distribution. Thus, the
 190 null hypothesis, H_0 , will be rejected when $T_1 \geq Z_\alpha$ where Z_α is the upper α quantile of the
 191 standard normal distribution.

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196 **2.2. Second proposed method**

197

198 The second method we are proposing for the nondecreasing ordered alternatives in a mixed
 199 design is designed as follows:

$$T_2 = \frac{T_2^* - [E(NMJT) + E(BNMJT)]}{\sqrt{V(NMJT) + V(BNMJT)}} \quad (26)$$

200 where

$$T_2^* = NMJT + BNMJT = \sum_{i=1}^{k-1} \sum_{j=i+1}^k i(j-i)U_{ij} + \sum_{l=1}^b NMJT_l \quad (27)$$

201 Here, k is the number of treatments, and b is the number of blocks. $NMJT$ is the new modified
 202 Jonckheere-Terpstra test for the l^{th} block. Under H_0 , for large sample sizes, this test will also
 203 follow an asymptotic normal distribution. The null hypothesis, H_0 , will be rejected when
 204 $T_2 \geq Z_\alpha$.

205
 206 It is noted that versions of the Jonckheere-Terpstra test are being developed for other types
 207 of designs such as a two-stage nested design [12].
 208

209 2.3. Simulation Study

210
 211 The aim of this section is to describe in detail the Monte Carlo simulation study that is used to
 212 investigate the performance of the proposed tests and to compare them to each other and
 213 with the tests proposed by Magelet *al.* (2009) [11]. The performance of the tests are evaluated
 214 by the estimated powers and whether or not they maintain the stated level of significance
 215 (α). The power of a test can be defined as the probability of rejecting a false H_0 . Likewise,
 216 the level of significance (α) is defined as the probability of rejecting a true H_0 .

217
 218 In this study, three underlying distributions are used including, the standard normal
 219 distribution, the exponential distribution with mean one, and t-distribution with $DF = 3$. For
 220 each distribution, we consider three scenarios of proportions of the number of blocks in the
 221 *RCBD* portion to the sample size in the *CRD* portion, namely, assuming that the portion of
 222 the number of blocks in *RCBD* is *larger*, *equal*, and *smaller* than the portion of the sample
 223 size in the *CRD*. However, regarding the *CRD* portion, we consider cases where the sample
 224 sizes are equal.

225
 226 Based on 5000 iterations, the study is conducted for each combination of distributions with
 227 nondecreasing location parameters at 0.05 level of significance. The level of significance is
 228 estimated for each test by generating 5,000 sets of samples from the populations when the
 229 null hypothesis is true (i.e., the location parameter arrangements are the same for all
 230 treatments) and counting the number of times the null hypothesis is rejected, dividing by the
 231 number of iterations. Similarly, the power for each test is estimated by generating 5,000 sets
 232 of samples from the populations when the alternative hypothesis is true (i.e., the location
 233 parameter arrangements are different for at least one treatment) and count the number of
 234 times the null hypothesis is rejected, divided by the number of iterations. All the simulations
 235 are performed using the statistical program SAS9.4.

236
 237 Moreover, powers are estimated based on a variety of location parameter arrangements. In
 238 particular, we considered cases when there is an equal distance between the parameters;
 239 cases where the distances between the parameters are distinct; cases where the some
 240 parameters are equal and the rest were different; cases where the distance between the
 241 some parameters is twice as large as the distance of others; and cases where the distance
 242 between the first two parameters = chosen to be the same as the distance between the

243 last parameters. The location parameters are denoted by $\mu_1, \mu_2, \mu_3, \mu_4,$ and μ_5 for treatment
 244 one, two, three, four, and five, respectively.

245
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247 3. RESULTS AND DISCUSSION

248

249 The results of the simulation study play a significant role in evaluating the performance of the
 250 proposed tests compared to each other and to those proposed by Magelet *al.* (2009). Thus,
 251 the aim of this section is to introduce the results of the simulation study described in the
 252 previous section. The proposed tests are designed to analyze data in a mixed design of a
 253 CRD and a RCBD.

254

255 In each table, the results of the simulation study are defined based on number of treatments
 256 (k), the distribution used to simulate the data, the sample size in the CRD portion, and the
 257 number of blocks in the RCBD portion. Besides, a variety of location parameter
 258 arrangements, shifts, are considered on $k = 3, 4,$ and 5 treatments. The estimated level of
 259 significance (α) and the estimated power for each test are given so that the first row of each
 260 table represents the estimated α -level; however, the rest of rows represent the estimated
 261 powers.

262

263 Results as to which test did better in particular situation are consistent regardless of the
 264 number of treatments (namely, 3, 4, or 5) and the different distributions considered. Results
 265 are shown in Tables 1 and 2 for cases when we have four treatments ($k = 4$), and the
 266 number of blocks in the randomized complete block design (RCBD) portion is larger than the
 267 sample sizes in the completely randomized design (CRD) portion. These results are for the
 268 standard normal and standard exponential distributions. It can be seen that the first
 269 proposed test (T_1)
 270

271 Table 1. Percentage of Rejection for $k = 4$; Exponential Distribution: Block = 16 and $n = 8$

μ_1	μ_2	μ_3	μ_4	C_1	C_2	T_1	T_2
0	0	0	0	5.46	4.96	5.32	5.04
0	0.1	0.2	0.3	47.44	38.36	43.56	29.48
0	0	0.25	0.25	47.22	37.30	44.64	30.54
0	0.125	0.25	0.25	40.18	32.42	31.44	22.04
0	0	0	0.5	67.54	56.12	81.80	60.38
0.05	0.1	0.3	0.5	74.88	64.76	76.68	56.02
0	0.25	0.5	0.5	80.66	71.60	69.94	50.62
0	0.5	0.5	1	98.84	96.24	99.08	90.26
0.1	0.2	0.6	1	99.42	97.64	99.50	94.22
0.25	0.25	0.5	0.5	47.22	37.30	43.46	29.76
0	0.1	0.3	0.7	94.54	88.52	96.24	83.78
0	0.05	0.15	0.35	55.40	44.96	57.38	38.74
0	0.15	0.2	0.5	74.84	64.26	76.14	54.54
0	0	0.05	0.3	42.26	33.54	48.24	33.06
0	0	0.1	0.6	85.60	75.74	92.66	74.76

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274 Table 2. Percentage of Rejection for $k = 4$; Normal Distribution: Block = 32 and $n = 8$

μ_1	μ_2	μ_3	μ_4	C_1	C_2	T_1	T_2
0	0	0	0	4.78	5.32	5.00	4.86
0	0.1	0.2	0.3	33.84	28.40	31.24	20.68
0	0	0.25	0.25	33.58	28.18	31.88	21.08
0	0.125	0.25	0.25	28.84	24.18	22.54	15.86
0	0	0	0.5	55.48	47.06	68.00	45.72
0.05	0.1	0.3	0.5	59.14	49.76	59.34	38.16
0	0.25	0.5	0.5	67.10	58.12	55.18	36.58
0	0.5	0.5	1	97.14	93.06	95.26	77.86
0.1	0.2	0.6	1	97.80	94.56	98.12	86.34
0.25	0.25	0.5	0.5	33.58	28.18	31.70	21.60
0	0.1	0.3	0.7	85.56	77.40	89.26	67.12
0	0.05	0.15	0.35	40.08	33.68	42.32	27.36
0	0.15	0.2	0.5	59.00	50.32	58.60	39.26
0	0	0.05	0.3	31.28	26.88	35.98	24.52
0	0	0.1	0.6	72.80	63.10	83.18	58.68

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has higher powers in cases where the arrangements of location parameters follow the pattern that the first two parameters are equal and the last two parameters are different such as (0, 0, 0.05, 0.3), cases where the arrangements of location parameters following the pattern that distance between the first two parameter is equal to the distance between the last two parameters such as (0, 0.5, 0.5, 1), and lastly cases where the arrangements of location parameters follow the pattern that the distance between the fourth and third parameters is twice as large as the distance between the third and the second, while the distance between the third and the second parameters is twice as large as the distance between the second and the first such as (0, 0.1, 0.3, 0.7).

Tables 3 and 4 show the results of the simulation study for cases when $k = 4$, and the number of blocks in the *RCBD* portion is equal relative to the sample sizes in the *CRD* portion. When one of the parameters is quite a bit larger relative to the other parameters, the first proposed test (T_1) has the highest powers. As an example of this, consider the case where the first two parameters are equal and the last two are distinct such as (0, 0, 0.1, 0.6). When the parameters are equally spaced, or the last two parameters are equal to each other, C_1 has the highest powers. The test C_1 was one of the tests proposed by Magel et al. [11] and given in equation (13).

305 Table 3. Percentage of Rejection for $k = 4$; Exponential Distribution: Block = 10 and $n = 10$

μ_1	μ_2	μ_3	μ_4	C_1	C_2	T_1	T_2
0	0	0	0	5.26	5.00	4.88	5.04
0	0.1	0.2	0.3	42.36	35.24	38.36	28.80
0	0	0.25	0.25	41.70	35.80	40.60	31.66
0	0.125	0.25	0.25	36.12	31.04	29.50	23.26
0	0	0	0.5	62.86	52.34	76.82	60.74
0.05	0.1	0.3	0.5	69.52	60.74	72.60	57.36
0	0.25	0.5	0.5	76.84	67.42	66.26	51.62
0	0.5	0.5	1	98.08	94.84	97.32	90.84
0.1	0.2	0.6	1	98.76	96.44	99.20	95.48
0.25	0.25	0.5	0.5	41.64	34.92	40.44	31.12
0	0.1	0.3	0.7	92.10	85.46	94.96	84.32
0	0.05	0.15	0.35	51.64	43.26	53.28	40.66
0	0.15	0.2	0.5	69.10	59.52	70.60	54.58
0	0	0.05	0.3	39.72	31.58	46.56	33.36
0	0	0.1	0.6	80.86	70.32	89.16	74.68

306 Table 4. Percentage of Rejection for $k = 4$; Normal Distribution: Block = 20 and $n = 20$

μ_1	μ_2	μ_3	μ_4	C_1	C_2	T_1	T_2
0	0	0	0	4.46	5.08	4.68	5.16
0	0.1	0.2	0.3	35.22	27.38	32.92	24.40
0	0	0.25	0.25	37.24	27.72	34.82	25.12
0	0.125	0.25	0.25	31.24	24.10	25.74	19.04
0	0	0	0.5	61.36	46.78	73.12	53.98
0.05	0.1	0.3	0.5	63.34	49.48	64.60	47.76
0	0.25	0.5	0.5	73.80	56.50	60.36	42.82
0	0.5	0.5	1	98.78	93.06	97.50	87.28
0.1	0.2	0.6	1	99.10	94.78	99.40	94.24
0.25	0.25	0.5	0.5	36.48	28.28	34.80	25.48
0	0.1	0.3	0.7	90.90	78.30	92.10	77.94
0	0.05	0.15	0.35	44.28	34.06	46.22	33.30
0	0.15	0.2	0.5	63.84	49.10	65.02	47.02
0	0	0.05	0.3	34.18	25.80	40.46	28.96
0	0	0.1	0.6	79.74	64.04	87.98	70.48

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308 Tables 5 and 6 show the results of the proposed tests and the tests proposed by Magel et
 309 al.[11] in terms of the estimated level of significance and the estimated powers for the
 310 normal, exponential, and student's t distributions for four treatments ($k = 4$) when the
 311 proportion of the *RCBD* portion is *smaller*, than the *CRD* portion. Results were the same as

312 when the CRD and RCBD portions were equal as to which test performed better. when the
 313 RCBD portion was $\frac{1}{2}$ the CRD portion (Table 5).

314 Table 5. Percentage of Rejection for $k = 4$; Exponential Distribution: Block = 8 and $n = 16$

μ_1	μ_2	μ_3	μ_4	C_1	C_2	T_1	T_2
0	0	0	0	5.00	4.72	4.68	4.50
0	0.1	0.2	0.3	49.48	44.84	46.14	39.62
0	0	0.25	0.25	47.62	42.34	44.96	39.12
0	0.125	0.25	0.25	40.94	36.76	32.36	27.86
0	0	0	0.5	69.84	62.88	83.86	75.62
0.05	0.1	0.3	0.5	77.68	70.96	79.70	70.98
0	0.25	0.5	0.5	83.06	76.72	72.40	64.06
0	0.5	0.5	1	99.40	98.22	99.08	96.88
0.1	0.2	0.6	1	99.58	98.90	99.88	99.14
0.25	0.25	0.5	0.5	46.10	41.28	43.28	37.70
0	0.1	0.3	0.7	95.84	92.14	97.44	93.56
0	0.05	0.15	0.35	56.50	50.82	59.92	50.76
0	0.15	0.2	0.5	76.32	70.08	78.42	69.04
0	0	0.05	0.3	43.66	37.30	51.80	42.40
0	0	0.1	0.6	86.58	81.58	93.80	88.44

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 316 When the proportion of the number of blocks in the RCBD is *one-eighth* the sample size in
 317 the CRD portion (Table 6), the proposed test T_2 tends to have higher estimated powers than
 318 the others under the following location parameters arrangements: (0, 0, 0, 0.5), (0.05, 0.1,
 319 0.3, 0.5), (0, 0.1, 0.3, 0.7), (0, 0.05, 0.15, 0.35) and (0, 0, 0.05, 0.3). Note that in this case, it
 320 is the second versions of the test statistics that perform better.

321 Table 6. Percentage of Rejection for $k = 4$; Normal Distribution: Block = 4 and $n = 32$

μ_1	μ_2	μ_3	μ_4	C_1	C_2	T_1	T_2
0	0	0	0	4.76	4.84	5.26	5.16
0	0.1	0.2	0.3	31.10	34.60	29.88	32.32
0	0	0.25	0.25	31.20	34.04	29.90	32.24
0	0.125	0.25	0.25	25.58	27.62	21.04	23.26
0	0	0	0.5	52.04	56.78	63.36	68.78
0.05	0.1	0.3	0.5	54.36	59.78	55.38	61.34
0	0.25	0.5	0.5	64.84	69.18	52.80	56.84
0	0.5	0.5	1	96.50	97.80	93.68	96.46
0.1	0.2	0.6	1	97.08	98.46	97.34	98.60
0.25	0.25	0.5	0.5	30.42	34.58	29.54	32.88
0	0.1	0.3	0.7	84.74	88.54	86.80	90.86
0	0.05	0.15	0.35	36.52	41.76	39.52	43.92
0	0.15	0.2	0.5	54.80	61.02	55.30	61.34
0	0	0.05	0.3	29.58	31.78	35.20	37.54
0	0	0.1	0.6	69.60	75.26	78.96	83.08

323 Table 7. Percentage of Rejection for $k = 3$; Normal Distribution: Block = 16 and $n = 4$

μ_1	μ_2	μ_3	C_1	C_2	T_1	T_2
0	0	0	4.88	4.98	4.74	4.36
0	0	0.5	36.52	35.58	41.28	33.70
0	0.5	0.5	36.32	35.68	27.48	23.18
0.05	0.25	0.5	31.92	30.96	32.14	26.18
0	0.3	0.5	37.10	36.14	33.76	27.90
0	0	1	80.92	78.96	88.84	80.40
0	1	1	80.86	78.92	66.56	56.16
0	0.5	1	81.86	80.32	80.62	70.96
0.5	0.5	1	36.52	35.58	41.70	34.44
0.5	1	1	36.22	35.68	29.34	23.98
0.1	0.5	1	74.12	72.38	75.36	64.90
0.1	0.3	0.7	46.30	44.92	47.60	38.78
0	0.25	0.5	36.88	35.94	35.32	29.52
0.2	0.5	0.8	46.12	44.96	43.98	36.38
0	0.1	0.8	65.68	63.34	71.64	61.24

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325 In Tables 7 and 8, the RCBD portion is larger than the CRD portion. The results in this case
 326 are the same as the results when the RCBD portion is equal to the CRD portion. The first
 327 versions of the test statistics have larger powers. The first newly proposed test statistic has
 328 larger powers when there is a relatively large difference between the last two location
 329 parameters. When the parameters are equally spaced or fairly close to each other, C_1 (one of
 330 the tests in Magel et al. [12]) has higher powers. In Table 7, $k=3$ and in Table 8, $k=5$. The
 331 number of populations did not affect the result as to which test statistic had higher powers,
 332 but the spacing between the last two parameters did.

333 Table 8. Percentage of Rejection for $k = 5$; Normal Distribution: Block = 16 and $n = 8$

μ_1	μ_2	μ_3	μ_4	μ_5	C_1	C_2	T_1	T_2
0	0	0	0	0	5.38	4.86	5.06	4.78
0.05	0.15	0.25	0.35	0.45	40.76	35.06	37.22	26.12
0	0.025	0.075	0.175	0.375	34.90	30.04	39.42	27.02
0	0	0	0	0.5	39.86	33.24	51.70	35.72
0	0	0.125	0.25	0.25	27.70	23.74	25.48	19.02
0	0.05	0.05	0.3	0.3	32.62	27.78	31.02	21.82
0.05	0.2	0.3	0.4	0.5	45.74	39.62	41.16	28.78
0	0	0	0.25	0.5	53.34	46.82	62.08	43.60
0	0	0	0.35	0.35	42.90	37.00	46.80	32.04
0	0	0.25	0.25	0.5	53.88	46.86	53.66	36.66
0	0	0	0.1	0.3	25.42	22.06	30.28	20.30
0	0	0	0.2	0.7	71.18	63.20	81.62	61.48
0	0.1	0.1	0.6	0.6	76.70	69.10	76.76	56.04

0	0.1	0.3	0.4	0.4	45.82	39.80	38.80	25.88
0	0.05	0.2	0.4	0.4	48.64	42.20	43.76	29.74

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4. CONCLUSION

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In this paper, novel nonparametric methods for the nondecreasing ordered alternative are proposed for a mixed design consisting of a combination of a CRD and *RCBD*. Three cases were considered where the number of blocks were proportional to the sample sizes. In particular, the number of blocks in the *RCBD* portion are *larger*, *equal*, and *smaller* than the sample sizes in the *CRD* portion. In either case, from the findings of the simulation study, it was shown that both the proposed tests appear to maintain their type one errors. This was also true of the tests proposed in Magel et al. [11].

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Moreover, the estimated powers for the method formed by standardized last idea (T_2) are less than the method formed by standardized first idea (T_1) under all distributions considered for all cases when the *RCBD* portion is at least $\frac{1}{2}$ or greater of the *CRD* portion. When the number of blocks in the *RCBD* portion are *one-eighth* the sample sizes in the *CRD* portion, we found (T_2) had higher powers than (T_1).

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The simulation study has also shown that the estimated power for the proposed test (T_1) is better than the test statistics proposed by Magelet *al.* [11] under the nondecreasing ordered alternative as long as a large jump is present between the last two adjacent location parameters such as (0, 0, 0.1, 0.6) and (0, 0, 0, 0.5) when the *RCBD* portion was $\frac{1}{2}$ or greater of the *CRD* portion. If the *RCBD* portion was only $\frac{1}{8}$ that of the *CRD* portion, then T_2 had the largest powers of all the test statistics. If there was not a large jump between the last two parameters, C_1 had the largest powers when the *RCBD* portion was at least $\frac{1}{2}$ of the *CRD* portion. If the *RCBD* portion was only $\frac{1}{8}$ the *CRD* portion under this circumstance, then C_2 had the larger powers.

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