

Data-driven treatment selection for seamless phase II/III trials incorporating early-outcome data

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Seamless phase II/III clinical trials are conducted in two stages with treatment selection at the first stage. In the first stage, patients are randomized to a control or one of $k > 1$ experimental treatments. At the end of this stage, interim data are analysed, and a decision is made concerning which experimental treatment should continue to the second stage. If the primary endpoint is observable only after some period of follow-up, at the interim analysis data may be available on some early outcome on a larger number of patients than those for whom the primary endpoint is available. These early endpoint data can thus be used for treatment selection. For two previously proposed approaches, the power has been shown to be greater for one or other method depending on the true treatment effects and correlations. We propose a new approach that builds on the previously proposed approaches and uses data available at the interim analysis to estimate these parameters and then, on the basis of these estimates, chooses the treatment selection method with the highest probability of correctly selecting the most effective treatment. This method is shown to perform well compared with the two previously described methods for a wide range of true parameter values. In most cases, the performance of the new method is either similar to or, in some cases, better than either of the two previously proposed methods. © 2014 The Authors. *Pharmaceutical Statistics* published by John Wiley & Sons Ltd.

Keywords: adaptive seamless design; multi-arm multi-stage trial; surrogate endpoint

1. INTRODUCTION

Drug development is very expensive and risky with many compounds failing in late development phases. Adaptive designs have been recognized as a way to improve efficiency of drug development by industry and regulators alike [1–3]. Of particular interest are designs combining aspects of the clinical development process into one single study that would have traditionally been assessed in separate trials and phases, for instance, adaptive seamless phase II/III designs [4–7]. Seamless phase II/III clinical trials are conducted in two stages. In the first stage, patients are randomized to control or some number $k > 1$ experimental treatments. At the end of this stage, interim data are analysed, and a decision is made concerning which experimental treatment should continue, along with the control, to the second stage. If the primary endpoint is observable only after some period of follow-up, at the interim analysis, data may be available on some early outcome on a larger number of patients than those for whom the primary endpoint is available. These early endpoint data can thus be used for guiding the choice of treatments to continue. It has been demonstrated, for a range of settings, that adaptive trial designs are generally more efficient, and thereby quicker due to improved resource management, than conventional programmes with phase II studies for treatment selection and phase III studies for confirmation of the efficacy of the selected treatments [8].

Two different procedures for incorporating early endpoint data in a treatment selection design have been proposed by Stallard [9] and Friede *et al.* [10]; the focus in this paper is on treatment selection, general principles and methods for hypothesis testing in adaptive seamless designs are discussed in detail elsewhere [5,8,10], as are alternative methods in the setting where early endpoint data are available at interim analysis [11,12]. Friede *et al.* propose a method of treatment selection using early endpoint data only. In contrast, Stallard uses any available final (primary) endpoint data in addition to early endpoint data for treatment selection at interim. Although both approaches differ in the way in which data from the two stages are combined, they have both been shown to control the type I error rate. In this paper,

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we propose a new data-driven methodology to pick the optimal approach to treatment selection, which uses data available at an interim analysis to estimate treatment effects and correlations between endpoints and then, on the basis of these estimates, chooses the treatment selection method with the highest probability of selecting the most effective treatment. This new data-driven method is compared with the more established data-driven methods of Friede *et al.* and Stallard for a wide range of true parameter values. We focus here on using interim data available at an early point in a trial to select the method to undertake treatment selection. If, however, this was not possible, and it was necessary to decide on the appropriate method at the planning stage, then one would have little option but to rely on for instance other (external) sources of information or data (possibly from a pilot study) to augment ones beliefs about the likely effect sizes and associations between early and final endpoints. Section 2 describes a randomized trial assessing the efficacy of a novel compound in patients with primary hypertension, which provides a motivating example of how early outcome data might be used for decision-making at an interim analysis of a seamless phase II/III trial. Appropriate notation is established, and treatment selection for the two previously proposed methods is described in detail in Section 3. The new data-driven selection rule is developed in Section 4. Section 5 provides a worked example of how selection probabilities are calculated using data from the motivating example. In Section 6, numerical evaluation of the new data-driven selection rule is undertaken in a simulation study, and the performance of the new method is compared with the two previously proposed methods. The paper concludes with a discussion in Section 7.

2. MOTIVATING EXAMPLE

Calhoun *et al.* [13] report a double-blind, placebo-controlled, randomized trial assessing the efficacy and safety of a novel compound in patients with primary hypertension. The primary efficacy endpoint was the change in clinic diastolic blood pressure (DBP; mmHg) from baseline to the end of the 8-week double-blind treatment period. A total of $n=524$ patients were randomized to receive either one of four dose regimens of the new anti-hypertensive, an active control (AC) or placebo for 8 weeks. The standard deviation (SD) of the differences from baseline was assumed to be 10 mmHg in the sample size calculation. Additional readouts of the primary endpoints were available at weeks 1, 2 and 4. The primary comparisons were with placebo (and not with the AC) using a multiple-contrast test [14] to control the familywise type I error rate at 0.025 one-sided. Figure 1 gives the differences in diastolic blood pressure (DBP) between the dose regimens and AC with placebo over the time course of the 8 weeks of double-blind treatment. The differences of dose regimens 3 (DR3) and the AC to placebo at the end of the 8-week treatment period were found to be statistically significant. These two treatments also showed the biggest reductions in DBP at weeks 2 and 4 in comparison to placebo. Here, we investigate the possibility that a single interim analysis be undertaken at some point before the trial endpoint had been observed on all patients, but where, dependent on timing, a single early endpoint were available on a group of patients, and additionally some primary efficacy endpoint data were also available from a sub-group of these patients at interim. We would expect both that differences in DBP at different time points for the same patient would be correlated and that treatment effects on the earlier outcomes might

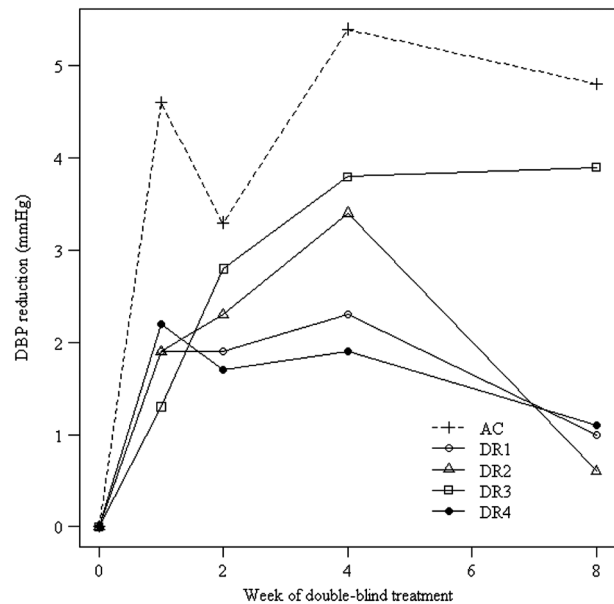


Figure 1. Differences in diastolic blood pressure (DBP) between active treatments (i.e. dose regimens (DR) 1–4 and active control (AC)) and placebo over the time course of the 8 weeks of double-blind treatment.

indicate the ordering of treatment effects for the final outcome. This suggests that we might be able to use the early time point data in the treatment selection at an interim analysis. The patients were recruited into the trial over a period of about seven months. Since the trial included a two-week washout and a two-week single-blind run-in period, at an interim analysis half way through recruitment, say after 4 months, only those patients recruited during the first month would have had completed their 8-week double-blind treatment period whereas the other patients could have only contributed 1-, 2- or 4-week measurements to the interim analysis. In what follows, we will explore selection rules for adaptive treatment selection at interim incorporating incomplete patient follow-up data.

3. PHASE II/III CLINICAL TRIALS WITH EARLY OUTCOME DATA

3.1. A flexible hypothesis testing method for phase II/III clinical trials with early outcome data

Following, e.g. Stallard [9], we envisage a two-stage clinical trial, in which in the first stage, patients are randomized to the control treatment T_0 or to one of k experimental treatments, $T_i, i = 1, \dots, k$, with one of these experimental treatments, T_i , selected to continue, along with the control treatment, T_0 , to the second stage.

We suppose that, at the end of the first stage, data are available on the primary, final outcome for n_1 patients in each treatment group, but that in addition, early outcome data are available for some larger number of patients, N_1 per group. At the end of the second stage, final outcome data will be available for all n_2 patients receiving treatments T_i and T_0 .

Denote by X_{ij} and Y_{ij} respectively the early and final outcome data from patient j in group $i, i = 0, \dots, k, j = 1, \dots, n_2$. The two endpoints for each patient are assumed to follow a bivariate

normal distribution with

$$\begin{pmatrix} X_{ij} \\ Y_{ij} \end{pmatrix} \sim N \left(\begin{pmatrix} \mu_{b_i} \\ \mu_{B_i} \end{pmatrix}, \begin{pmatrix} \sigma_0^2 & \rho_w \sigma_0 \sigma \\ \rho_w \sigma_0 \sigma & \sigma^2 \end{pmatrix} \right). \quad (1)$$

Given the mean values, individual patients are assumed to be independent so that $cov(X_{ij}, X_{i'j'}) = 0$, $cov(Y_{ij}, Y_{i'j'}) = 0$ and $cov(X_{ij}, Y_{i'j'}) = 0$ for $i \neq i'$ or $j \neq j'$.

The parameters of interest are the treatment effects on the final outcome for each treatment relative to the control, $\mu_{B_i} - \mu_{B_0}$, which will be denoted by θ_i , $i = 1, \dots, k$, with H_{0i} denoting the null hypotheses $\theta_i = 0$, which will be tested against the one-sided alternative hypotheses $H_{1i} : \theta_i > 0$. The familywise type I error rate may be controlled in the strong sense by application of the closed testing procedure [15] and combination test [16,17]. Intersection hypotheses $H_S = \bigcap_{i \in S} H_{0i}$, where $S \subseteq \{1, \dots, k\}$, are tested using data from each of the two stages and combined using a combination test. Individual elementary hypotheses H_{0i} are rejected at level α if and only if all intersection hypotheses with index sets that include i are rejected at individual local test level α [8]. Stagewise p -values, $p_{S,1}$ and $p_{S,2}$, are combined using the weighted inverse normal method, with (pre-specified) weights (w_1 and w_2) proportional to the planned sample size at each stage, with H_S rejected at level α if $C(p_{S,1}, p_{S,2}) \leq \alpha$ where $C(p_{S,1}, p_{S,2}) = 1 - \Phi(w_1 \Phi^{-1}(1 - p_{S,1}) + w_2 \Phi^{-1}(1 - p_{S,2}))$ [18].

For σ assumed known, a test of H_{0i} may be based on standardized test statistics using final endpoint data from stages one and two given by

$$Z_{i,1} = \frac{\sum_{j=1}^{N_1} (Y_{ij} - Y_{0j})}{\sqrt{2N_1\sigma^2}}$$

and

$$Z_{i,2} = \frac{\sum_{j=N_1+1}^{n_2} (Y_{ij} - Y_{0j})}{\sqrt{2(n_2 - N_1)\sigma^2}}.$$

Note that, in order to ensure independence between test statistics from the two stages, $Z_{i,1}$ uses data from all patients for whom early endpoint data are available in stage one and $Z_{i,2}$ uses data from only those patients recruited in the second stage. From equation (1), Z_{ij} are normally distributed, with $E(Z_{i,1}) = \theta_i \sqrt{N_1 / (2\sigma^2)}$, $E(Z_{i,2}) = \theta_i \sqrt{(n_2 - N_1) / (2\sigma^2)}$, $var(Z_{ij}) = 1$, $cov(Z_{ij}, Z_{i'j'}) = 0$ for $j \neq j'$ and $cov(Z_{ij}, Z_{i'j}) = 1/2$ for $i \neq i'$. The p -values for a test of each of the intersection hypotheses H_S at each stage may be obtained by a Dunnett test [19] that uses the test statistic $Z^{max} = \max_{i \in S} Z_{ij}$. The second stage p -value for a test of H_S , $p_{S,2}$, is set to p_l , the p -value for selected treatment T_l , if $l \in S$ and to 1 if $l \notin S$, as in the conservative approach suggested by Posch et al. [20].

This approach to hypothesis testing in the setting of treatment selection with early endpoint data was proposed by Friede et al. [10] who considered specifically the setting where early endpoint data only were available for decision making (treatment selection) at stage one. Although they also propose a treatment selection rule for use in this setting, the analysis approach controls the familywise type I error rate in the strong sense for any treatment selection method based on the data observed at the end of stage one. An alternative treatment selection method for a setting in which a combination of final and early endpoint data are available from stage one was proposed by Stallard [9]. The treatment selection methods of Friede et al. and Stallard are described in detail in the succeeding text.

3.2. Treatment selection method of Friede et al.

Friede et al. [10] proposed a method for selection of the treatment T_l that should continue to the second stage. In the setting they envisage, early endpoint data only are available at the time when the treatment selection is made, so that the selection does not use any final endpoint data. The method can, however, be applied when some final endpoint data are available, as discussed by Kunz et al. [21].

Let

$$Z_i^* = \frac{\sum_{j=1}^{N_1} (X_{ij} - X_{0j})}{\sqrt{2N_1\sigma_0^2}}$$

be the standardized test statistic based on the early endpoint data from stage one. From equation (1), Z_i^* are normally distributed with $E(Z_{i,1}) = (\mu_{b_i} - \mu_{b_0}) \sqrt{N_1 / (2\sigma_0^2)}$, $var(Z_{i,1}) = 1$ and $cov(Z_{i,1}, Z_{i',1}) = 1/2, i \neq i'$.

Treatment T_l is then selected where $l = \arg \max \{Z_i^*\}$. The probability that, e.g. treatment 1 is selected is therefore given by

$$\begin{aligned} &Pr(\text{select treatment 1 using Friede et al. method}) = \\ &Pr(Z_1^* = \max \{Z_i^*\}) = Pr(Z_2^* - Z_1^* < 0, \dots, Z_k^* - Z_1^* < 0). \end{aligned}$$

This may be calculated using the joint distribution of $Z_2^* - Z_1^*, \dots, Z_k^* - Z_1^*$ given by

$$\begin{pmatrix} Z_2^* - Z_1^* \\ \vdots \\ Z_k^* - Z_1^* \end{pmatrix} \sim N \left(\begin{pmatrix} \frac{\mu_{b_2} - \mu_{b_1}}{\sigma_0} \sqrt{\frac{N_1}{2}} \\ \vdots \\ \frac{\mu_{b_k} - \mu_{b_1}}{\sigma_0} \sqrt{\frac{N_1}{2}} \end{pmatrix}, \begin{pmatrix} 1 & \frac{1}{2} & \dots & \frac{1}{2} \\ \frac{1}{2} & 1 & \dots & \vdots \\ \vdots & \vdots & \ddots & \vdots \\ \frac{1}{2} & \dots & \frac{1}{2} & 1 \end{pmatrix} \right).$$

Denoting by $F(\mathbf{x}, \mu, \Sigma)$, the cumulative distribution function of a multivariate normal with mean μ and variance-covariance matrix Σ evaluated at \mathbf{x} , we have

$$\begin{aligned} &Pr(\text{select treatment 1 using Friede et al. method}) = \\ &F \left(\mathbf{0}, \begin{pmatrix} \frac{\mu_{b_2} - \mu_{b_1}}{\sigma_0} \sqrt{\frac{N_1}{2}} \\ \vdots \\ \frac{\mu_{b_k} - \mu_{b_1}}{\sigma_0} \sqrt{\frac{N_1}{2}} \end{pmatrix}, \begin{pmatrix} 1 & \frac{1}{2} & \dots & \frac{1}{2} \\ \frac{1}{2} & 1 & \dots & \vdots \\ \vdots & \vdots & \ddots & \vdots \\ \frac{1}{2} & \dots & \frac{1}{2} & 1 \end{pmatrix} \right), \quad (2) \end{aligned}$$

where here $\mathbf{0}$ denotes the zero-vector with $k - 1$ elements. Similar joint distributions enable calculation of the probabilities that other treatments are selected.

3.3. Treatment selection method of Stallard

Stallard [9] proposed a method for selection of the treatment T_l that should continue to the second stage using a combination of early and final endpoint data.

Let S_j denote the standardized score statistic for θ_j given all data available at the end of stage one. For known σ , σ_0 and ρ_w , S_j is given by

$$S_j = \frac{\sum_{j=1}^{n_1} (Y_{ij} - Y_{0j} - \rho_w \frac{\sigma}{\sigma_0} (X_{ij} - X_{0j} - \bar{X}_i + \bar{X}_0)) / n_1}{\sigma \sqrt{2/N_1^*}}$$

where $\bar{X}_i = \sum_{j=1}^{N_1} X_{ij}$ and $N_1^* = n_1 N_1 / (N_1 - \rho_w^2 (N_1 - n_1))$. With σ , σ_0 and ρ_w unknown, S_j may be estimated using the double

regression method of Engel and Walstra [22]; Galbraith and Marschner [23] provide an analogous expression to the above in the setting of a series of interim analyses at which the two groups are compared, with the number of patients for whom short-term and long-term data are available increasing through the duration of the trial. Stallard and Kunz *et al.* show that S_1, \dots, S_k follow a multivariate normal distribution with $E(S_i) = \theta_i \sqrt{N_1^* / (2\sigma^2)}$, $\text{var}(S_i) = 1$ and $\text{cov}(S_i, S_{i'}) = 1/2, i \neq i'$.

Stallard proposes selecting treatment T_l where $l = \arg \max\{S_i\}$. The probability that, e.g. treatment 1 is selected is therefore given by

$$\begin{aligned} \Pr(\text{select treatment 1 using Stallard method}) &= \\ \Pr(S_1 = \max\{S_i\}) &= \Pr(S_2 - S_1 < 0, \dots, S_k - S_1 < 0). \end{aligned}$$

This may be calculated using the joint distribution of $S_2 - S_1, \dots, S_k - S_1$ given by

$$\begin{pmatrix} S_2 - S_1 \\ \vdots \\ S_k - S_1 \end{pmatrix} \sim N \left(\begin{pmatrix} \frac{\mu_{B_2} - \mu_{B_1}}{\sigma} \sqrt{\frac{N_1^*}{2}} \\ \vdots \\ \frac{\mu_{B_k} - \mu_{B_1}}{\sigma} \sqrt{\frac{N_1^*}{2}} \end{pmatrix}, \begin{pmatrix} 1 & \frac{1}{2} & \dots & \frac{1}{2} \\ \frac{1}{2} & 1 & \ddots & \vdots \\ \vdots & \ddots & \ddots & \frac{1}{2} \\ \frac{1}{2} & \dots & \frac{1}{2} & 1 \end{pmatrix} \right),$$

so that

$$\Pr(\text{select treatment 1 using Stallard method}) = F \left(\mathbf{0}, \begin{pmatrix} \frac{\mu_{B_2} - \mu_{B_1}}{\sigma} \sqrt{\frac{N_1^*}{2}} \\ \vdots \\ \frac{\mu_{B_k} - \mu_{B_1}}{\sigma} \sqrt{\frac{N_1^*}{2}} \end{pmatrix}, \begin{pmatrix} 1 & \frac{1}{2} & \dots & \frac{1}{2} \\ \frac{1}{2} & 1 & \ddots & \vdots \\ \vdots & \ddots & \ddots & \frac{1}{2} \\ \frac{1}{2} & \dots & \frac{1}{2} & 1 \end{pmatrix} \right). \quad (3)$$

As in the preceding text, similar joint distributions enable calculation of the probabilities that other treatments are selected.

3.4. Practical considerations

Successful application of the methods of Friede *et al.* [10] and Stallard [9], described in Sections 3.2 and 3.3, depends in part at least on the adequacy of the early endpoint as a surrogate for the final endpoint. In a trial setting where a group of experimental treatments (i.e. two or more) are compared with a control the concept of group-level surrogacy corresponds to correlation between the group means for the early (surrogate) and final endpoints, and analogously individual level surrogacy corresponds to within-group correlation. For a fully validated surrogate endpoint, we require both individual and group-level surrogacy. For the methods of both Friede *et al.* and Stallard, the use of an early endpoint for decision making improves the power of the test. However, the properties of the two methods are very different. For the method of Stallard, the gain in power comes from the within-group correlation between the early (surrogate) and final outcomes (ρ_w); i.e. from the presence of individual-level surrogacy. Whereas for the method of Friede *et al.* unless there is a large within-group correlation, the gain in power arises from the correlation between treatment effects for the early and final outcomes; i.e. from the presence of group-level surrogacy. The latter property implies that in some settings, the Friede *et al.* method

could perform badly. For instance, consider a setting where the μ_{b_i} 's were widely spaced, but the μ_{B_i} 's were close together. The method of Friede *et al.* will pick a treatment based on early outcome data μ_{b_i} with high probability, but because of the close spacing of the μ_{B_i} 's, it is likely that the selected treatment may not perform equally well based on the final outcome. Clearly, the Friede *et al.* methodology is only really sensible if the early end-point data alone provide useful information for treatment selection; if it leads one to pick the *wrong* treatment(s), it is not sensible. The Stallard method may do better in such settings, dependent on the magnitude of ρ_w . So we would advise that before either of these methods are used, some thought is given to the issues discussed here, and the likely nature and magnitude of all associations between early and final outcomes be considered during the trial planning stage. Kunz *et al.* [21] provide a comparison of the treatment selection rules proposed by Stallard and Friede *et al.* They show that the properties of the methods depend on the true unknown treatment effects and correlations between the endpoints, and that neither the Stallard method nor the Friede *et al.* method is always preferable in terms of selection probability or power. The new data-driven selection rule discussed in Section 4 describes how one might select which of the two methods to use, but if one had strong beliefs or data to support one or other methodology above the other at the planning stage of a trial, then clearly the most sensible option would be to proceed with that methodology.

4. A NEW DATA-DRIVEN SELECTION RULE

In this section, we introduce a new treatment selection method based on the methods proposed by Friede *et al.* and Stallard. The idea is that following observation of the data at the end of stage one, both the methods of Friede *et al.* and Stallard will be applied and compared. If both methods agree on which treatment should proceed along with the control to the second stage, the treatment indicated by both methods will be selected. If the methods indicate that different treatments should proceed, the stage one data will be used to assess which of the methods is likely to be correct, as explained in more detail in the remainder of this section, and the treatment indicated by this method will be selected.

The aim is to provide a method that has high power, where this is defined to be the probability of correctly selecting the most effective treatment and rejecting the null hypothesis corresponding to this treatment having effect no greater than the control treatment. This probability is bounded above by the probability that the most effective treatment is selected on the basis of the data observed at stage one. In principle, it may be possible to compute the power analytically; however, it is very hard compared with computation of the selection probabilities from (2) and (3); our proposed method for choosing between the treatment selection rules is based on the latter method.

Given observed stage one data, estimates of the effects, μ_b and μ_B , variances, σ_0 and σ , and the correlation, ρ_w , can be obtained. These will be denoted by $\hat{\mu}_b$, $\hat{\mu}_B$, $\hat{\sigma}_0$, $\hat{\sigma}$ and $\hat{\rho}_w$, respectively.

On the basis of these estimates, we can calculate the estimated probability of selection of each treatment for each of the two methods given by (2) and (3), with the estimated probability of, e.g. selection of treatment T_1 given by

$$F \left(\mathbf{0}, \begin{pmatrix} \frac{\hat{\mu}_{B_2} - \hat{\mu}_{B_1}}{\hat{\sigma}_0} \sqrt{\frac{N_1}{2}} \\ \vdots \\ \frac{\hat{\mu}_{B_k} - \hat{\mu}_{B_1}}{\hat{\sigma}_0} \sqrt{\frac{N_1}{2}} \end{pmatrix}, \begin{pmatrix} 1 & \frac{1}{2} & \dots & \frac{1}{2} \\ \frac{1}{2} & 1 & \ddots & \vdots \\ \vdots & \ddots & \ddots & \frac{1}{2} \\ \frac{1}{2} & \dots & \frac{1}{2} & 1 \end{pmatrix} \right), \quad (4)$$

for the Friede *et al.* selection method and

$$F \left(\mathbf{0}, \begin{pmatrix} \frac{\hat{\mu}_{B_2} - \hat{\mu}_{B_1}}{\hat{\sigma}} \sqrt{\frac{\hat{N}_1^*}{2}} \\ \vdots \\ \frac{\hat{\mu}_{B_k} - \hat{\mu}_{B_1}}{\hat{\sigma}} \sqrt{\frac{\hat{N}_1^*}{2}} \end{pmatrix}, \begin{pmatrix} 1 & \frac{1}{2} & \dots & \frac{1}{2} \\ \frac{1}{2} & 1 & \ddots & \vdots \\ \vdots & \ddots & \ddots & \frac{1}{2} \\ \frac{1}{2} & \dots & \frac{1}{2} & 1 \end{pmatrix} \right) \quad (5)$$

for the Stallard selection method, where $\hat{N}_1^* = n_1 N_1 / (N_1 - \hat{\rho}_w^2 (N_1 - n_1))$.

These estimated selection probabilities give an indication of the preference of each treatment based on each selection method. For the Friede *et al.* and Stallard selection methods, the treatments with the largest estimated selection probability is that with the largest value of Z_i^* and S_i , respectively, and so will be the treatments selected. When the two methods indicate selection of different treatments, we therefore propose to select the method for which the estimated selection probability for the selected treatment is largest. With the treatment selected in this way, we propose to test the hypotheses $H_{0,i}, i = 1, \dots, k$ using the method as described in Section 3.1. As Friede *et al.* show that this controls the familywise type I error rate in the strong sense for any treatment selection method based on data from patients enrolled in stage one of the trial, the familywise type I error rate is strongly controlled.

5. EXAMPLE

The motivating example provides an informative setting in which to demonstrate how selection probabilities are calculated for the Stallard and Friede *et al.* methods. Let us consider the possibility that an interim analysis was undertaken where early outcome data, DBP at the 4-week assessment time point, were available from N_1 patients, and 8-week assessment data, the primary efficacy endpoint, were available from n_1 patients. For the selected motivating example, we would expect there to be strong within group correlations (ρ_w) between early (4-week) and final (8-week) outcome data. Also, available data (Figure 1) suggest strong correlations between treatment group effects, albeit the sample of available treatment group means is small. Clearly, in this setting, the methods of both Friede *et al.* and Stallard have potential application, so it is of interest to understand under what conditions one or other method performs the better. Depending on the timing of the interim analysis, we investigate two options; (i) where 4 week data were available from half of the target population ($N_1 = 45$), and 8 week data were also available from a small proportion ($n_1 = 10$) of these patients, and (ii) where the interim analysis was undertaken at an earlier time point where $N_1 = 25$ and $n_1 = 5$. The AC treatment is always selected in addition to the placebo. The parameters of interest, the effects for the four test treatments, on the final outcome relative to the placebo $\mu_{B_i} - \mu_{B_0}$ for $i = 1, \dots, 4$, are estimated from Figure 1 to be approximately 1.0, 0.6, 3.9 and 1.1 mmHg for treatments DR1, DR2, DR3 and DR4, respectively. Similarly, the estimates of early outcome (4 week)

treatment effects relative to the placebo $\mu_{B_i} - \mu_{B_0}$ are estimated from Figure 1 to be approximately 2.3, 3.4, 3.8 and 1.9 mmHg. Given these estimates for $\hat{\mu}_{B_1}, \dots, \hat{\mu}_{B_4}$, $\hat{\mu}_{B_1}, \dots, \hat{\mu}_{B_4}$, estimates for early and final outcome standard deviations of $\hat{\sigma} = \hat{\sigma}_0 = 10$ and correlation parameter $\hat{\rho}_w = 0.9$, treatment selection probabilities for the Stallard and Friede *et al.* methods can be calculated using expressions (4) and (5). These can be evaluated simply using the `pvmnorm` function in the `mvtvnorm` package [24] in R [25], to give treatment selection probabilities for setting (i) of (0.102, 0.072, 0.716, 0.110) and (0.118, 0.337, 0.468, 0.076) for the Stallard and Friede *et al.* methods respectively and for setting (ii) treatment selection probabilities of (0.143, 0.113, 0.592, 0.152) and (0.150, 0.324, 0.426, 0.110) for the two methods, all for treatments DR1, DR2, DR3 and DR4 respectively. Both methods would select treatment DR3 with highest probability, in both settings, with setting (ii), where fewer patients were available, having lower selection probabilities for the DR3 group. Treatment group DR2 illustrates where the methods can differ. For the Stallard method the selection probability for this group is much smaller than for the Friede *et al.* method, due to the relatively poor performance of this treatment group at the 8-week assessment, from which data are not used by the Friede *et al.* method for determining selection probabilities. In general, the two methods may give different selection probabilities, the properties of these procedures together with the data-driven method are explored in more detail in a simulation study.

6. NUMERICAL EVALUATION OF THE NEW DATA-DRIVEN SELECTION RULE

6.1. Selection probability and power

In general, there are two different ways to define the selection probability: (i) the probability to select any effective treatment and (ii) the probability to select the most effective treatment. Throughout this paper, we use the latter definition. In order to be consistent with this definition, we also define the power as the probability to reject the null hypothesis belonging to the most effective treatment. Furthermore, we will, without loss of generality, focus on T_1 and report the probability of selecting T_1 as well as rejecting the corresponding null hypothesis, H_{01} .

In this section, we report the results of a simulation study to explore the properties of the new procedure. Results are shown for a trial with three experimental groups, that is $k = 3$, compared with a control group, T_0 , with $n_1 = 4$, $N_1 = 32$ and $n_2 = 64$. For treatment T_1 , we assume final endpoint effects, μ_{B_1} , from -0.5 to 1 in steps of 0.1 and early endpoint effects, μ_{b_1} , of $-0.2, 0$, and 0.2 , and we set $\mu_{B_0} = \mu_{b_0} = 0$, $\mu_{B_2} = \frac{1}{2}\mu_{B_1}$, $\mu_{B_3} = \frac{1}{4}\mu_{B_1}$, $\mu_{b_2} = \frac{1}{2}\mu_{b_1}$ and $\mu_{b_3} = \frac{1}{4}\mu_{b_1}$ to give early and final endpoint effects of a half and a quarter of the size of the effect for T_1 . Note that even for $\mu_{B_1} < 0$, the probability to select T_1 is reported although, in this case, this is the worst-performing treatment. We investigate correlation ρ_w values of $-0.9, -0.5, 0, 0.5$ and 0.9 . Results are based on 10,000 simulations for each scenario considered. Supplementary material describes the results of additional simulation studies, covering a wide range of alternative parameter settings.

Simulation results are shown in Figure 2. The panels on the left-hand side of the figure show the probability of selecting treatment T_1 for the Stallard (black dashed line), the Friede *et al.* (black dash-dotted line) and the data-driven method (black solid line). The panels on the right-hand side show the probability of both selecting treatment T_1 and rejecting H_{01} at the $\alpha = 0.025$ level,

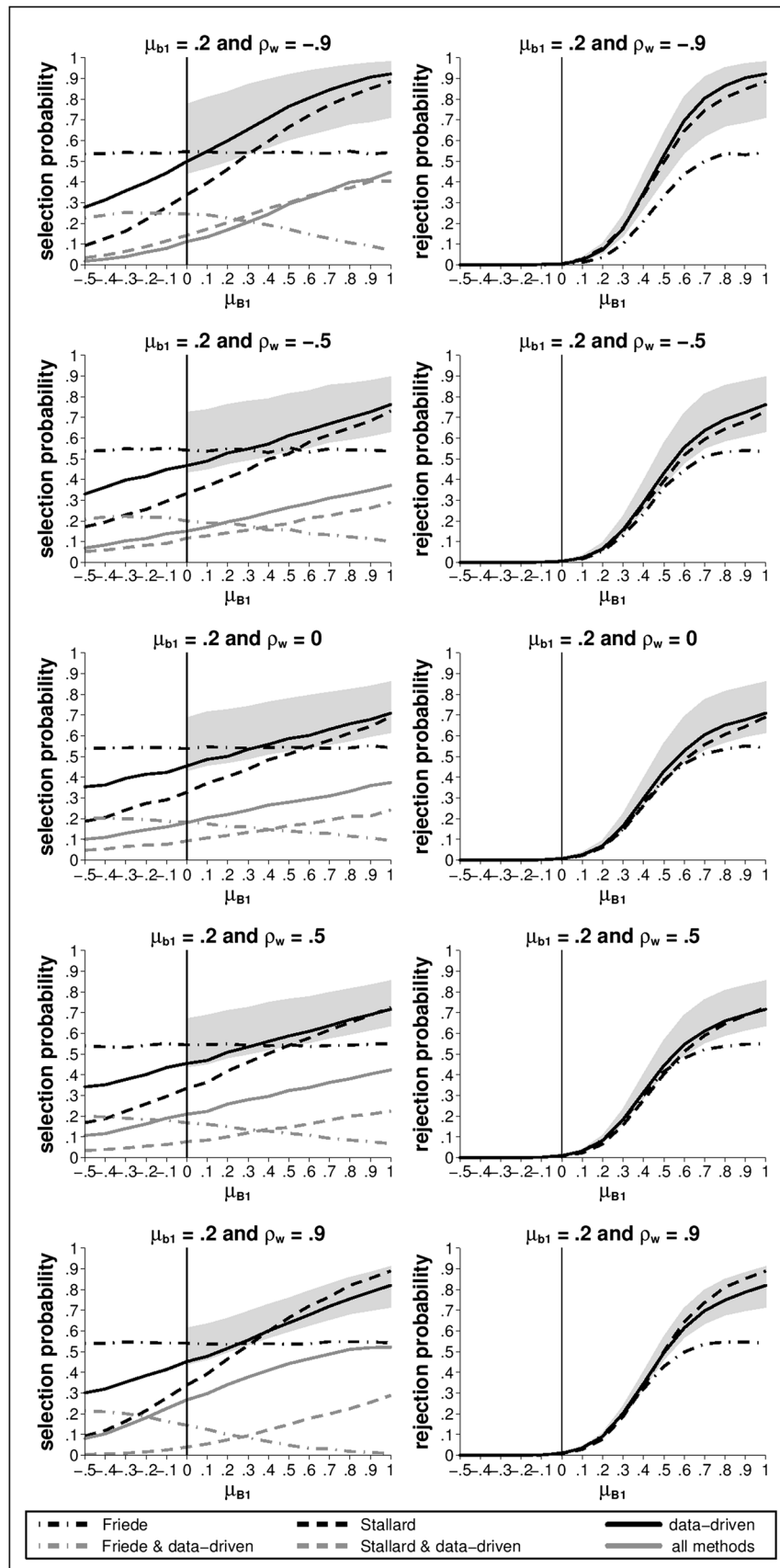


Figure 2. Selection (left) and rejection probabilities (right) for T_1 , as a function of final endpoint effect μ_{B1} , using the Stallard [9], Friede et al. [10] and data-driven methods for T_1 early endpoint effect $\mu_{b1} = 0.2$ and correlation between endpoints within each group $\rho_w = \{-0.9, -0.5, 0, 0.5, 0.9\}$; $\mu_{B2} = \frac{1}{2}\mu_{B1}$, $\mu_{B3} = \frac{1}{4}\mu_{B1}$, $\mu_{b2} = 0.1$ and $\mu_{b3} = 0.05$.

which is the power as defined in the preceding text, for the three methods using the same line types.

For the Friede *et al.* method, the selection is based on early endpoint data only. The selection probability for this method thus does not depend on μ_{B_1} or ρ_w . This is in contrast to the selection probability for the Stallard method, which increases with increasing μ_{B_1} and, to a lesser extent, with the magnitude of ρ_w .

For ρ_w positive, for μ_{B_1} larger than about 0.3, the selection probability for the data-driven method is mostly similar to that for the Stallard method, i.e. the better of the two methods under these scenarios. For μ_{B_1} less than 0.3, the selection probability for the new method is between that for the two previous methods. For ρ_w less than zero, for μ_{B_1} above about 0.3, the new method leads to selection of treatment T_1 with higher probability than either the Stallard or Friede *et al.* methods.

Under the scenarios considered, the power is generally higher for the Stallard selection method than for the Friede *et al.* method. The power for the new method is generally similar to that for the Stallard method for larger ρ_w and greater than that for the Stallard method for smaller or negative ρ_w .

In order to evaluate the performance of the data-driven method, two boundaries can be defined against which the data-driven method is then compared. The data-driven method can only select a treatment group that was selected by at least one of the Stallard and the Friede *et al.* method. An upper bound for the selection probability for the new method is thus the probability that at least one of the Stallard and Friede *et al.* methods selects treatment T_1 . This boundary is only meaningful if T_1 is the most effective treatment, that is if $\mu_{B_1} > 0$. An alternative to choosing between the treatments selected by the Stallard and the Friede *et al.* methods using the data-driven rule introduced in this paper is to randomly select between the treatments recommended by the two methods when they disagree. The selection probability for such an approach would be the average of that for the Stallard and Friede *et al.* methods. Although not a lower bound for the selection probability, it is desirable that the new method should perform at least as well as this approach. The grey-shaded area in Figure 2 marks the range of probabilities bounded by these two values. Ideally, the results for the data-driven method should be between the boundaries and as close as possible to the upper one. For the selection probability, we see that the data-driven method starts closer to the lower boundary for smaller values of μ_{B_1} but comes closer to the upper boundary for higher values of μ_{B_1} . A range of power values defined similarly is shown on the plots on the right-hand side of Figure 2. The power for the data-driven method is quite close to the upper boundary for all scenarios considered.

To understand the results for the data-driven method in more detail, the selection probability can be split up in three different categories as illustrated by the grey lines on Figure 2: T_1 was selected by the data-driven method because (1) the Stallard method but not the Friede *et al.* method selected T_1 (grey dashed line), (2) the Friede *et al.* but not the Stallard method selected T_1 (grey dash dotted line), (3) both the Friede *et al.* and the Stallard method selected T_1 (grey solid line). Note that the probabilities for category (3) can be calculated using equation (6), given in the succeeding text and that the results for the three categories sum to the probabilities for the data-driven method. As explained above, it is more likely that both the Stallard and the Friede *et al.* methods select T_1 if the correlation is positive than if the correlation is negative. For example, for $\mu_{B_1} = 0$, the probability that

all three methods select T_1 is just about 10% for $\rho_w = -0.9$ but about 26% for $\rho_w = +0.9$. On the other hand, the Stallard and the data-driven method selected T_1 in about 14% of the cases for $\rho_w = -0.9$ but in only 4% of the cases for $\rho_w = +0.9$. Similarly, in about 25% of the cases, the Friede *et al.* and the data-driven method selected T_1 if the correlation is negative, but in only 15% of the cases if the correlation is positive.

6.2. Effect of ρ_w

It is interesting to note that although the selection probabilities and power when using the Stallard or Friede *et al.* selection method is the same for ρ_w of given magnitude, irrespective of the sign, the properties of the new data-driven selection method do depend on the sign of ρ_w . The selection probabilities for the Friede *et al.* method and Stallard method are given by expressions such as equations (4) and (5), respectively. The first of these does not depend on ρ_w , and the second depends on ρ_w only through N_1^* , which depends on ρ_w^2 , but not on the sign of ρ_w . As the new selection method is based on the treatments indicated by both the Stallard method and the Friede *et al.* method, it is important to note that the treatments selected by the two methods are not independent, because they both depend on the early outcome data $X_{ij}, i = 0, \dots, k, j = 1, \dots, n_1$. The probability that both methods lead to selection of treatment T_1 is obtained from the joint distribution of $Z_2^* - Z_1^*, \dots, Z_k^* - Z_1^*, S_2 - S_1, \dots, S_k - S_1$, which does depend on the sign of ρ_w , and is given by

$$\begin{pmatrix} Z_2^* - Z_1^* \\ \vdots \\ Z_k^* - Z_1^* \\ S_2 - S_1 \\ \vdots \\ S_k - S_1 \end{pmatrix} \sim N \left(\begin{pmatrix} \frac{\mu_{b_2} - \mu_{b_1}}{\sigma_0} \sqrt{\frac{N_1}{2}} \\ \vdots \\ \frac{\mu_{b_k} - \mu_{b_1}}{\sigma_0} \sqrt{\frac{N_1}{2}} \\ \frac{\mu_{B_2} - \mu_{B_1}}{\sigma_0} \sqrt{\frac{N_1^*}{2}} \\ \vdots \\ \frac{\mu_{B_k} - \mu_{B_1}}{\sigma_0} \sqrt{\frac{N_1^*}{2}} \end{pmatrix}, \Sigma \right) \quad (6)$$

where

$$\Sigma = \begin{pmatrix} 1 & \rho_w \sqrt{\frac{N_1^*}{N_1}} \\ \rho_w \sqrt{\frac{N_1^*}{N_1}} & 1 \end{pmatrix} \otimes \begin{pmatrix} 1 & \frac{1}{2} & \dots & \frac{1}{2} \\ \frac{1}{2} & 1 & \ddots & \vdots \\ \vdots & \ddots & \ddots & \frac{1}{2} \\ \frac{1}{2} & \dots & \frac{1}{2} & 1 \end{pmatrix}$$

Joint distributions enabling calculation of the joint probability of selection of other treatments by the Stallard and Friede *et al.* methods are given by similar expressions.

The effect of the dependence on the sign of ρ_w is illustrated in Table I, which gives the probabilities for each of the four cases that T_1 is selected by (i) the Stallard method but not the Friede *et al.* method, (ii) the Friede *et al.* method but not the Stallard method, (iii) both methods, (iv) neither of the methods, together with the marginal distributions based on equations (2), (3) and (6). The first two sub-tables give probabilities for $\rho_w = \pm 0.9$. For $\rho_w = -0.9$, T_1 is selected by both methods with probability 0.21 compared with 0.39 for $\rho_w = 0.9$. On the other hand, both methods fail to select T_1 with probability only 0.13 for $\rho_w = -0.9$ but with probability 0.31 for $\rho_w = 0.9$. Conversely, for $\rho_w = -0.9$, T_1 is selected by at least one of the two methods with probability 0.87 while for $\rho_w = 0.9$, T_1 is selected by at least one of the

Table I. Probability of selecting T_1 for the Stallard and Friede *et al.* methods for (a) $\mu_{B_1} = 0.3, \mu_{B_2} = 0.15, \mu_{B_3} = 0.075, \mu_{b_1} = 0.2, \mu_{b_2} = 0.1$ and $\mu_{b_3} = 0.05$, and (b) $\mu_{B_1} = \mu_{B_2} = \mu_{B_3} = \mu_{b_1} = \mu_{b_2} = \mu_{b_3} = 0$ based on equations (2), (3) and (6).

		(a)						(b)					
		$\rho_w = -0.9$			$\rho_w = +0.9$			$\rho_w = -1$			$\rho_w = +1$		
		Stallard			Stallard			Stallard			Stallard		
		No	Yes	Total	No	Yes	Total	No	Yes	Total	No	Yes	Total
Friede	No	0.13	0.33	0.46	0.31	0.15	0.46	0.33	0.33	0.67	0.67	0.00	0.67
	Yes	0.33	0.21	0.54	0.15	0.39	0.54	0.33	0.00	0.33	0.00	0.33	0.33
Total		0.46	0.54	1.00	0.46	0.54	1.00	0.67	0.33	1.00	0.67	0.33	1.00

two methods with probability 0.69. As the data-driven method can only select a treatment group that was selected by at least one of the Stallard and the Friede *et al.* method, the differences between the joint distributions for different signs of the correlation lead to the different results for the data-driven method that can be seen in Figure 2. An even more extreme example is given in the last two subtables in Table I, when $\rho_w = \pm 1$. In this case, when $\rho_w = -1$, either the Stallard method or the Friede *et al.* method is correct but never both of them, while for $\rho_w = 1$, the two methods always agree, so that the probability of at least one method selecting treatment T_1 changes from 0.33 to 0.67 depending on the sign of ρ_w .

7. CONCLUSION

When a clinical trial with a primary endpoint that is observed only after a relatively long follow-up period includes interim analyses, it is desirable, if possible, to base these analyses at least in part on more rapidly observable short-term endpoint data. In general, any data available at the interim analysis could be used for treatment selection, as long as the type I error rate is controlled. Recently, Jenkins *et al.* [26] proposed a design for subgroup selection at interim using correlated survival outcomes. In this paper, we have considered two-stage adaptive seamless phase II/III clinical trials in which interim analysis data are used to decide which of a number of experimental treatments should continue along with the control treatment to the second stage. Two methods have previously been proposed for the use of short-term endpoint data in this setting [9,10]. A comparison of the properties of these methods [21] has indicated that neither method is uniformly more powerful than the other, but that each can be more powerful depending on the (unknown) true values of the treatment effects on long-term and short-term endpoints and the correlation between the endpoints.

The method of analysis proposed by Friede *et al.* allows considerable flexibility in terms of the choice of the rule used to select the most promising treatment based on the interim data observed. In particular, the selection rule itself may be chosen in a data-dependent way based on data from patients recruited in stage one of the trial whilst maintaining control of the familywise type I error rate in the strong sense. This is the starting point for the method proposed in this paper, in which, when the Stallard and Friede *et al.* methods would lead to selection of different treatments, the method that maximizes the selection probability based on parameter values estimated from the observed data is used. The focus here has been on the development of a novel methodological approach, so for reasons of

conciseness, we have not discussed some of the more practical issues concerning implementation, for instance recruitment patterns, whether recruitment should continue seamlessly and what triggers the interim analysis. These are all key issues, amongst many others, that would need to be considered when deciding whether the methods described here might work in any given setting; a fuller discussion of these issues is provided elsewhere [1,27]. Mainly, for reasons of clarity of exposition, we have assumed fixed-effects models throughout this paper. Random-effects models for the methods of Stallard and Friede *et al.* have been discussed elsewhere [21], and also developed for the new data-driven approach of Section 4, but as the conclusions and simulation results were not changed qualitatively, the simpler fixed-effects approach only is presented here. We have restricted discussion in this paper to the most simple setting of a two-stage design where a single treatment is selected at an interim analysis, using a single early endpoint. Future work will investigate how this might be generalized to designs where more than one early endpoint were available for treatment selection, more than one treatment were selected at interim analysis and multiple (more than two) stages were planned.

The comparison in the preceding text, together with additional simulation results available as supplementary material, indicate that the new data-driven method has power higher than the average of the two existing methods, which is higher than a strategy that would pick one or other of the methods at random when they disagree, for nearly all parameter values considered. The only exceptions are when the treatment effect on the short-term endpoint for treatment T_1 , μ_{b_1} , is less than 0 and the treatment effect for T_1 on the long-term endpoint, μ_{B_1} , is close to 0, or when $\mu_{b_1} \approx 0$ and $\rho_w \approx 1$. In many cases, the new method is similar in performance to the most powerful of the other two methods. In some cases, the new method is better than either of the two existing methods, e.g. for large μ_{B_1} , if ρ_w is close to 0 and $\mu_{b_1} > 0$, for nearly all values of $\mu_{B_1} > 0$, if ρ_w is close to -1 and $\mu_{b_1} > 0$ and for some μ_{B_1} if ρ_w is close to 1 and $\mu_{b_1} > 0$.

The choice of whether or not to use the new data-driven method depends on the level of confidence in predictions of μ_{b_1} , μ_{B_1} , σ_0 , σ and ρ_w at the planning stage. If reliable estimates are available, it is possible to work out which method will give the best results based on equations (2), (3), and (6) and simulation studies for the power as reported above. Depending on the parameter estimates, as indicated above, the preferred method could be one of the existing methods or the new method. If there is uncertainty regarding parameter values, the new data-driven method would be a good choice. In summary, the numerical evaluations of the new data-driven methodology, presented in

Section 6 and the supplementary material, show that the power of the data-driven method was close to the upper boundary of all the methods considered; i.e. it always performed relatively well. Also, the new data-driven method was always similar in performance to the more powerful of the two previously described methods. That is, it was either marginally more or less powerful, across all the scenarios tested, than the most powerful of either the Stallard or the Friede *et al.* methods. More specifically, when we assumed positive study findings for both early and final outcomes, the new method generally performed better than either the Stallard or the Friede *et al.* methods. Therefore, given that *a priori* we would generally not know which of the two previously described methods will perform the better, the new data-driven method is a good choice, as it will perform nearly as well or often better than either of the other methods.

The new data-driven method described here provides a straightforward and appealing way of choosing between two methods that have been suggested elsewhere for treatment selection in the chosen setting, where up to now there was no clear guidance. Our simulations have shown that this new method generally performs well. The new method just represents another way of using the data available at the interim analysis to select a treatment, and we accept that this is somewhat arbitrary. An interesting area for future work would be to attempt to make this decision in an optimal way, perhaps using prior knowledge of the expected magnitude of effects and associations between the endpoints in a Bayesian setting.

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REFERENCES

- [1] Gallo P, Chuang-Stein C, Dragalin V, Gaydos B, Krams M, Pinheiro J. Adaptive designs in clinical drug development - an executive summary of the pharma working group. *Journal of Biopharmaceutical Statistics* 2006; **16**:275–283.
- [2] Food and Drug Administration (FDA). Innovation or stagnation? challenge and opportunities on the critical path to new medical products, 2012. Available at: <http://www.fda.gov/ScienceResearch/SpecialTopics/CriticalPathInitiative/CriticalPathOpportunitiesReports/ucm077262.htm> (accessed November 28, 2012).
- [3] Phillips AJ, Keene ON. Adaptive designs for pivotal trials: discussion points from the psi adaptive design expert group. *Pharmaceutical Statistics* 2006; **5**(1):61–66.
- [4] Stallard N, Friede T. A group-sequential design for clinical trials with treatment selection. *Statistics in Medicine* 2008; **27**:6209–6227.
- [5] Bretz F, Koenig F, Brannath W, Glimm E, Posch M. Tutorial in biostatistics: adaptive designs for confirmatory clinical trials. *Statistics in Medicine* 2009; **28**:1181–1217.
- [6] Di Scala L, Glimm E. Time-to-event analysis with treatment arm selection at interim. *Statistics in Medicine* 2011; **30**:3067–3081.
- [7] Brannath W, Koenig F, Bauer P. Multiplicity and flexibility in clinical trials. *Pharmaceutical Statistics* 2007; **6**(3):205–216.

- [8] Bretz F, Schmidli H, König F, Racine A, Maurer W. Confirmatory seamless phase II/III clinical trials with hypotheses selection at interim: general concepts. *Biometrical Journal* 2006; **48**:623–634.
- [9] Stallard N. A confirmatory seamless phase II/III clinical trial design incorporating short-term endpoint information. *Statistics in Medicine* 2010; **29**:959–971.
- [10] Friede T, Parsons N, Stallard N, Todd S, Valdés-Márquez E, Chataway J, Nicholas R. Designing a seamless phase II/III clinical trial using early outcomes for treatment selection: an application in multiple sclerosis. *Statistics in Medicine* 2011; **30**:1528–1540.
- [11] Liu Q, Pledger GW. Phase 2 and 3 combination designs to accelerate drug development. *Journal of the American Statistical Association* 2005; **100**:493–502.
- [12] Todd S, Stallard N. A new clinical trial design combining phases II and III: Sequential designs with treatment selection and a change of endpoint. *Drug Information Journal* 2005; **39**:109–118.
- [13] Calhoun DA, White WB, Krum H, Guo W, Bermann G, Trapani A, Lefkowitz MP, Menard J. Results of a randomized, double-blind, placebo- and active-controlled phase 2 trial. *Circulation* 2011; **124**:1945–1955.
- [14] Bretz F, Pinheiro JC, Branson M. Combining multiple comparisons and modeling techniques in dose-response studies. *Biometrics* 2005; **61**:738–748.
- [15] Marcus R, Peritz E, Gabriel KR. On closed testing procedures with special reference to ordered analysis of variance. *Biometrika* 1976; **63**(3):655–660.
- [16] Bauer P, Köhne K. Evaluation of experiments with adaptive interim analyses. *Biometrics* 1994; **50**(4):1029–1041.
- [17] Brannath W, Posch M, Bauer P. Recursive combination tests. *Journal of the American Statistical Association* 2002; **97**:236–244.
- [18] Lehman W, Wassmer G. Adaptive sample size calculations in group sequential trials. *Biometrics* 1999; **55**(4):1286–1290.
- [19] Dunnett CW. A multiple comparison procedure for comparing several treatments with a control. *Journal of the American Statistical Association* 1955; **50**:1096–1121.
- [20] Posch M, König F, Branson M, Brannath W, Dunger-Baldauf C, Bauer P. Testing and estimation in flexible group sequential designs with adaptive treatment selection. *Statistics in Medicine* 2005; **24**:3697–3714.
- [21] Kunz CU, Friede T, Parsons N, Todd S, Stallard N. A comparison of methods for treatment selection in seamless phase II/III clinical trials incorporating information on short-term endpoints. *Journal of Biopharmaceutical Statistics*. DOI: 10.1080/10543406.2013.840646.
- [22] Engel B, Walstra P. Increasing precision or reducing expense in regression experiments by using information from a concomitant variable. *Biometrics* 1991; **47**(1):13–20.
- [23] Galbraith S, Marschner IC. Interim analysis of continuous long-term endpoints in clinical trials with longitudinal outcomes. *Statistics in Medicine* 2003; **22**:1787–1805.
- [24] Genz A, Bretz F, Miwa T, Mi X, Leisch F, Scheipl F, Bornkamp B, Hothorn T. Package ‘mvtnorm’, 2012. Available at: <http://CRAN.R-project.org> (accessed 16.04.2014), R package version 0.9-9992.
- [25] R Development Core Team. *R: A language and Environment for Statistical Computing*. R Foundation for Statistical Computing: Vienna, Austria, 2011. Available at: <http://www.R-project.org> (accessed 16.04.2014), ISBN 3-900051-07-0.
- [26] Jenkins M, Stone A, Jennison C. An adaptive seamless phase II/III design for oncology trials with subpopulation selection using correlated survival endpoints. *Pharmaceutical Statistics* 2011; **10**(4):347–356.
- [27] Maca J, Bhattacharya S, Dragalin V, Gallo P, Krams M. Adaptive seamless phase II/III designs – background, operational aspects, and examples. *Drug Information Journal* 2006; **40**:463–473.

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