A Bayesian Adaptive Design with Biomarkers for Targeted Therapies

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Summary. Targeted therapies have become increasingly important for the treatment of various diseases. Biomarkers are a critical component of a targeted therapy as they can be used to identify patients who are more likely to benefit from a treatment. Targeted therapies, however, have created major challenges in the design, conduct and analysis of clinical trials. In traditional clinical trials, treatment effects for various biomarkers are typically evaluated in an exploratory fashion and only limited information about the predictive values of biomarkers are provided. New study designs are thus required which effectively evaluate both the diagnostic and the therapeutic implication of biomarkers. The Bayesian approach provides a useful framework for optimizing the clinical trial design by directly integrating information about biomarkers and clinical outcomes as they become available. We propose a Bayesian covariate-adjusted response-adaptive (BCARA) randomization design which utilizes individual biomarker profiles and patient’s clinical outcomes as they become available during the course of the trial, to assign the most efficacious treatment to individual patients. Predictive biomarker subgroups are determined adaptively using a partial least squares regression approach. A series of simulation studies were conducted to examine the operating characteristics of the proposed study design. The simulation studies show that the proposed
design efficiently identifies patients who benefit most from a targeted therapy and that there are substantial savings in the sample size requirements when compared to alternative designs. We conclude that the proposed design may serve a useful role in the early efficacy phase of targeted therapy development.

**Key words:** Bayesian logistic regression, personalized medicine, partial least squares regression.
1. INTRODUCTION

With the availability of new genomic and proteomic technologies, biomarkers have become increasingly important in both drug discovery and development. A biomarker is defined as a characteristic that is objectively measured or evaluated as an indicator of a biological or pharmacological response to a therapeutic intervention [1]. Biomarkers have the potential to allow for the selection of patients who are more likely to benefit from a targeted therapy. For example in cancer clinical trials, biomarkers can serve as a molecular indication of drug efficacy or toxicity. This allows individual patients to be treated based on the molecular determinants of their tumor cells. Biomarkers can be used to stratify patients, to make diagnosis, and to guide treatment [2]. The development of biomarkers for diagnosis and prognosis of personalized therapies allows the targeting of individualized treatments to patients most likely to benefit. New molecular profiling technologies which allow comprehensive analysis of single nucleotide polymorphisms (SNPs), gene expression, and protein profiles have resulted in a vast increase in the data available for biomarkers development [3].

At the design stage of the early clinical trial phases, the treatment effect sizes for the general patient population and the biomarker sub-populations are typically unknown. In traditional early efficacy phase clinical trials, patients are treated with the new drug and outcomes in patients with a positive biomarker are compared to outcomes in patients with a negative biomarker in an exploratory fashion. However, these trials provide only limited information about the predictive value of a biomarker, especially in situations where the prevalence of a positive biomarker in patients is small.

Several design strategies have been proposed for the prospective validation of biomarkers in the phase III setting [4,5,6,7]. Sargent *et al.* [6] divided clinical trial designs for predictive marker validation into two classes: Biomarker-by-treatment interaction design and biomarker-based-strategy design. In the biomarker-by-treatment interaction design, patients
are stratified according to their biomarker status, i.e., positive versus negative biomarker. Patients in each stratum are randomized to receive either the experimental arm or control arm. In the biomarker-based-strategy design, patients are randomized into a biomarker based-strategy arm and a non-biomarker based strategy arm. In the biomarker-based-strategy arm, patients with a positive biomarker are assigned to the experimental arm while patients with a negative biomarker are assigned to the control arm. In the non-biomarker based strategy arm, all patients are assigned to the control arm. In a modified version of the biomarker-based-strategy design, patients in the non-biomarker based strategy arm undergo a second randomization which assigns them to the experimental or control arm. This modification allows clarification of whether any finding regarding the efficacy of the biomarker directed approach to therapy is due to a true effect of the biomarker status or to an improved regimen regardless of biomarker status [6]. Moreover, this design may also allow a retrospective assessment of an alternative classification for the biomarker. As pointed out by Sargent et al. [6], the choice of the design for any particular trial depends on the nature of the conclusion that one wishes to draw and the strength of the evidence desired at the trial’s conclusion. The advantage of the biomarker-based strategy design over the biomarker-by-treatment interaction design is that it allows an assessment of the prognostic value of the biomarker. However, biomarker-based-strategy design generally requires a larger sample size than the biomarker-by-treatment interaction design.

Simon et al. [8] evaluated the relative efficiency of a targeted versus an untargeted clinical trial design for a randomized clinical trial design comparing a new treatment to a control. The untargeted trial design is the standard approach where patients are randomized into the treatment or control arm regardless of biomarker status. In the targeted trial design, only those patients who are predicted to respond, based on their biomarker profile, are randomized into the treatment or control arm. For example, the prediction of whether a patient will
respond to a treatment may be based on an assay that measures expression levels, or a multivariate gene expression model derived from transcript profiling [9,10,11]. Simon et al. [8] compared the two designs with regard to the number of patients required to achieve a fixed statistical power for detecting treatment effects. They showed that a targeted clinical trial design often requires fewer randomized patients than the untargeted design. The degree of reduction in sample size depends on the availability of the biomarker for identifying patients who will benefit from the new treatment and the prevalence of such patients. When the new treatment benefits only a subset of patients and those patients can be accurately identified, then the targeted design requires fewer patients than the untargeted design. However, targeted designs may lose some relative efficiency when there is a partial treatment effect for biomarker negative patients, possibly because of multiple potential mechanisms of action [8]. Therefore, while it is clear that a biomarker-based strategy can increase the efficiency of the trial, much depends on the performance of the diagnostics and the size of the treatment effect for target negative patients.

An adaptive design is a design that allows modifications of some aspects of the trial after its initiation without undermining the validity and integrity of the trial. It provides a mechanism for incorporating biomarker information during clinical trials. Adaptive methods for clinical trials have been studied extensively by many authors [12,13,14,15,16]. For example in a response adaptive clinical trial, patient outcomes can be used as they become available to adjust the allocation of future patients. This allows one to improve expected patient outcomes during the experiment, while still being able to reach good statistical decisions in a timely fashion. As we enter the era of personalized medicine, the role of adaptive designs in drug discovery and development has become increasingly important. Zhou et al. [17] recently proposed a Bayesian adaptive design for targeted therapy using pre-defined biomarker profile groups.
In this article, we propose a Bayesian covariate-adjusted response-adaptive randomization design for targeted therapies which utilizes individual patient’s biomarker profiles and clinical outcomes as they become available and which identifies subgroups of patients who respond best to a targeted therapy. Predictive biomarker subgroups are determined adaptively using a partial least squares logistic regression approach. The remainder of this article is organized as follows. In Section 2, the study design and computational aspects for implementing the design are described. The operating characteristics based on simulation studies are examined in Section 3. Finally, a brief summary is given in Section 4.

2. METHODS

2.1 A Clinical Trial with Biomarkers Evaluations

In the following, we assume a clinical trial with \( J \) treatments where the primary outcome is a dichotomous response variable which can be measured within a short time period. For example in cancer clinical trials, pathologic response, defined as the absence of residual tumors, can be measured via biopsy within one month after administrating treatment. It is assumed that measurements from a set of \( K \) biomarkers are available where the predictive or prognostic status of each biomarker can be unknown. Let \( Y_{tj} \) denote the response for the \( t \)th subject in treatment arm \( j \) \((j = 1, \ldots, J)\) with

\[
Y_{tj} = \begin{cases} 
1 & \text{if subject } t \text{ in treatment arm } j \text{ has a response,} \\
0 & \text{else.} 
\end{cases}
\]

Furthermore, let \( \mathbf{x}_t = (x_{1t}, \ldots, x_{Kt})^T \) denote a set of \( K \) biomarkers for subject \( t \). It is assumed each subject’s biomarker assessment is performed at or before study entry and that each biomarker is measured on a quantitative scale. For example, pharmacogenomic biomarkers are often measured quantitatively using immunohistochemistry staining with a scale ranging from with 0-3. The main objective is to develop an adaptive clinical trial design which evaluates the therapeutic intervention of targeted therapy, identifies subset of subjects who respond better to a targeted therapy, and optimize the treatment allocation by
randomizing more subjects to the superior treatment arm based on each subject’s individual biomarker profile.

2.2 Bayesian Logistic Regression Model for Predicting Clinical Outcome

We use a Bayesian logistic regression approach to predict response to a treatment of a newly accrued subject. The Bayesian approach provides a powerful framework for optimizing the clinical trial design by integrating information about biomarkers and clinical outcomes as they become available during the trial. Bayesian logistic regression is a natural choice to this problem as it can select predictive biomarkers among a large numbers of biomarkers without loss of model performance. For subject \( t \) in treatment arm \( j \), the probability of a response is modeled as

\[
\Pr(y_{tj} = 1|x_t) = \psi(\theta_j^T x_t) = \exp(\theta_j^T x_t) / (1 + \exp(\theta_j^T x_t))
\]

where \( \theta_j = (\theta_{j1}, \cdots, \theta_{jK}) \) has the prior distribution \( N(\mu_j, \Sigma_j) \), i.e., a multivariate normal distribution with mean \( \mu_j \) and covariance matrix \( \Sigma_j \). Holmes and Held [18] proposed an auxiliary variable approach to generate samples from the posterior distribution. Let

\[
y_{tj} = \begin{cases} 
1 & \text{if } z_{tj} > 0, \\
0 & \text{if } z_{tj} \leq 0,
\end{cases}
\]

with \( z_{tj} = \theta_j^T x_t + \epsilon_t, \epsilon_t \sim N(0, \lambda_t) \), \( \lambda_t = (2\omega_t)^2 \) where \( \omega_t \) has a Kolmogorov-Smirnov distribution [19]. The error term \( \epsilon_t \) has a scale mixture of normal form with a marginal logistic distribution.

For subjects \( 1, \cdots, i \) let \( X^{(i)} = (x_1^T, \cdots, x_i^T) \), \( \lambda^{(i)} = (\lambda_1, \cdots, \lambda_i)^T \), \( z^{(i)} = (z_{1j}, \cdots, z_{ij})^T \) and \( y^{(i)} = (y_{1j}, \cdots, y_{ij})^T \). The conditional distribution of \( \theta_j | z_j, \lambda^{(i)}, y^{(i)} \) has a normal distribution with mean \( B_j^{(i)} \) and covariance matrix \( \Xi_j^{(i)} \) where

\[
B_j^{(i)} = \left( \Sigma_j^{-1} + X^{(i)T}(\text{diag}(\lambda^{(i)}))^{-1}X^{(i)} \right) \left( \Sigma_j^{-1} \mu_j + X^{(i)T}(\text{diag}(\lambda^{(i)}))^{-1}z^{(i)} \right),
\]

\[
\Xi_j^{(i)} = \left( \Sigma_j^{-1} + X^{(i)T}(\text{diag}(\lambda^{(i)}))^{-1}X^{(i)} \right).
\]
A straightforward Gibbs sampling strategy can be implemented where $z_{tj}$ is sampled from a truncated normal distribution with

$$
z_{tj} | \theta_j, x_t, y_{tj}, \lambda_t \propto \begin{cases} 
N(\theta_j^T x_t, \lambda_t) I(z_{tj} > 0) & \text{if } y_{tj} = 1, \\
N(\theta_j^T x_t, \lambda_t) I(z_{tj} \leq 0) & \text{if } y_{tj} = 0,
\end{cases}
$$

for $t = 1, \cdots, i$. The conditional distribution $p(\lambda_t | z_{tj}, \theta_j)$ does not have a closed form. However, samples can be conveniently generated from this conditional distribution using standard rejection sampling [20]. Holmes and Held [18] proposed an efficient block Gibbs sampling strategy using iterative updates, i.e., $z_j^{(i)}, \lambda^{(i)} | \theta_j$ followed by $\theta_j | z_j, \lambda^{(i)}$. In this approach, $z_{tj} | \theta_j, x_t, y_{tj}$ follows a truncated logistic distribution with mean $\theta_j^T x_t$ and scale 1, i.e.,

$$
z_{tj} | \theta_j, x_t, y_{tj}, \lambda_t = o \begin{cases} 
\psi(\theta_j^T x_t) I(z_{tj} > 0) & \text{if } y_{tj} = 1, \\
\psi(\theta_j^T x_t) I(z_{tj} \leq 0) & \text{if } y_{tj} = 0.
\end{cases}
$$

The Gibbs sampling strategy described above can be conveniently implemented to generate samples from the posterior distribution $\theta_j | X^{(i)}, z_j^{(i)}$.

### 2.3 Bayesian Covariate-Adjusted Response-Adaptive Randomization

Response-adaptive randomization is a randomization technique in which the allocation of patients to the study arms is based on the responses of the previous outcomes. Response adaptive designs have been extensively studied, see, e.g., [12,21,22,23]. A covariate-adjusted response-adaptive randomization design can be used to incorporate biomarker information, as they become available during the conduct of the clinical trial, into a decision making process to evaluate both the diagnostic and the therapeutic intervention using biomarkers. One challenge involves the problem of how to incorporate information from multiple, possibly correlated, biomarkers into the design. The Bayesian paradigm may provide an ideal solution to both the diagnostic and the therapeutic intervention using multiple biomarkers with a covariate-adjusted response-adaptive randomization design. A Bayesian covariate-adjusted response-adaptive (BCARA) randomization can be implemented using the following steps.
For subject $i + 1$, we compute $E(y_{i+1,j}|x_{i+1};\theta_j) = \Pr(y_{i+1,j} = 1|x_{i+1};\theta_j)$ with respect to the posterior distribution $\pi(\theta_j|X^{(i)}, y^{(i)}_j)$, using the Gibbs sampling strategy described in Section 2.2. Furthermore, we compute
\[
p_j(y_{i+1,j}|x_{i+1}) = \Pr\left(\bigcap_{j' \neq j} E(y_{i+1,j'}|x_{i+1};\theta_{j'}) > E(y_{i+1,j'}|x_{i+1};\theta_{j'})|X^{(i)}, y^{(i)}_j\right)
\]
for each $j = 1, \cdots, J$. The probabilities in (1) can be used to calculate the randomization allocation probabilities, i.e., the probabilities of assigning subject $i + 1$ to treatment arms $j = 1, \cdots, J$. Specifically, the randomization rate for subject $i + 1$ and treatment $j$ is be computed as
\[
RR_{i+1,j} = \frac{(p_j(y_{i+1,j}|x_{i+1}))^c}{\sum_{j'=1}^{J} (p_{j'}(y_{i+1,j'}|x_{i+1}))^c},
\]
for a constant $c > 0$. Following the recommendations from Thall and Wathen [24] for Bayesian response adaptive designs we set $c = (i + 1)/(2N_{\text{max}})$ in (2) where $N_{\text{max}}$ is the trial’s maximum sample size. Before the BCARA randomization is implemented, a run-in phase is required. During the run-in phase, the first $n^*$ subjects are randomized to the treatment arms using a standard randomization procedure. A minimum of $n^* = J \times (K + 1)$ subjects is required for the run-in phase.

2.4 Adaptive Determination of Predictive Biomarker Profile Groups

A major objective of the study design is to identify subgroups of subject who respond better to a targeted therapy. We use a partial least squares regression (PLSR) approach to identify predictive biomarker subgroups. PLSR has become a popular dimension reduction tool that is based on a latent variables approach [25,26]. Contrary to principal component analysis, which attempts to find linear combinations of the predictors that explain most of the variation in these predictors using a small number of components, PLSR takes into account information about both the predictors and outcome variable in the definition of
scores and loadings. Specifically, PLSR finds linear combinations of the predictors that, in addition to the maximal variance constraint, also best explain the response. Originally, PLSR methods were developed for continuous response variables. Bastien et al [27] extended PLSR to partial least squares generalized linear regression models.

We use a partial least squares logistic regression (PLSLR) approach to classify subjects into biomarker profile groups which predict the clinical outcome. The biomarker profile groups are determined iteratively, i.e., after each new subject has entered the study. Let \( \mathbf{V}^{(i)} = (\mathbf{V}_1, \ldots, \mathbf{V}_i)^T \) where \( \mathbf{V}_t = (v_{t1}, \ldots, v_{t,J-1})^T \) denotes the vector of a dummy variables, i.e., \( v_{tj} = 1 \) if subject \( t \) (\( t = 1, \ldots, i \)) is assigned to treatment \( j \) (\( j = 1, \ldots, J - 1 \)) and zero otherwise. Furthermore, let \( \mathbf{W}^{(i)} \) denote the \( i \times (K + J - 1) \) matrix which consists of the standardized columns of \( \mathbf{X}^{(i)} \) and \( \mathbf{V}^{(i)} \) and let \( \mathbf{y}^{(i)} = (y^{(i)}_1, \ldots, y^{(i)}_J) \) denote the vector of responses for patients 1,..,i. The PLSLR model of \( \mathbf{y}^{(i)} \) on \( \mathbf{W}^{(i)} \) with \( m \) components can be written as

\[
g(\eta^{(i)}) = \sum_{h=1}^{m} b_h^{(i)} = \sum_{h=1}^{m} c_h \left( \sum_{l=1}^{K+J-1} \beta_h^* w_l^{(i)} \right), \tag{3}
\]

where \( c_h \) and \( \beta_h^* \) are the parameters of the PLSLR model, \( g(\cdot) \) is the canonical link function for the binomial distribution and \( m \leq K + J - 1 \).

The PLS components \( b_h^{(i)} \) of (3) can be computed iteratively, using the steps shown below. The computation of the first PLS component \( b_1^{(i)} \) is performed as follows:

1. Compute the regression coefficients \( a_{1l} \) of the logistic regression models of \( \mathbf{y}^{(i)} \) on \( \mathbf{w}_l^{(i)} \) for each \( l = 1, \ldots, K + J - 1 \).

2. Compute \( \zeta_1 = (a_{11}, \ldots, a_{1,K+J-1})^T / \sum_{l=1}^{K+J-1} a_{1l}^2 \).

3. Compute the first PLS component \( b_1^{(i)} = \mathbf{W}^{(i)} \zeta_1 / \zeta_1^T \zeta_1 \).

The \( h \)th PLS component is computed using the following steps:
1. Fit the logistic regression model of $y^{(i)}$ on $w_l^{(i)}$ and $b_1^{(i)}, \ldots, b_{h-1}^{(i)}$ for each $l = 1, \cdots, K+J-1$ and compute the regression coefficient $a_{hl}$, which corresponds to the predictor $w_l^{(i)}$, for each logistic regression model.

2. Compute $\zeta_h = (a_{h1}, \cdots, a_{hK+J-1})^T / \sum_{l=1}^{K+J-1} a_{hl}^2$.

3. Compute the residual matrix $R_{h-1}^{(i)}$ of the linear regression model of $W^{(i)}$ on $b_1^{(i)}, \cdots, b_{h-1}^{(i)}$.

4. Compute the $h$th PLS component $b_h^{(i)} = R_{h-1}^{(i)} \zeta_h / \zeta_h^T \zeta_h$.

The number $m$ of PLS components to be retained can be chosen by evaluating the predictive power of the current PLSLR model after each iteration using a cross-validation procedure [27]. Specifically, the algorithm is terminated if the predictive power of the PLSLR model does not increase after adding a new PLS component.

The number of biomarker subgroups can be determined after the run-in phase of the trial when the first $n^*$ subjects are randomized to the treatment arms using a standard equal randomization procedure. In the simplest scenario, subjects can be divided into two biomarker profile groups. For example, if after the run-in phase of the trial it is concluded that the PLSLR model with the first PLS component $b_1^{(i)}$ has a sufficiently high predictive power, the biomarker subgroups can be defined as follows:

$$
\delta_t^{(i)} = \begin{cases} 
1 & \text{if } b_1^{(i)} > 0, \text{ i.e., subject } t \ (t = 1, \cdots, i) \text{ has a positive biomarker profile,} \\
0 & \text{if } b_1^{(i)} \leq 0, \text{ i.e., subject } t \ (t = 1, \cdots, i) \text{ has a negative biomarker profile.}
\end{cases}
$$

A larger number of biomarker profile groups can be defined in a similar fashion.

2.5 Stopping Rule

It is assumed that $g = 1, \cdots, G$ different biomarker profile groups have been identified after the run-in phase. Let

$$
p_{jg}^{(i)} = \Pr \left( \bigcap_{j' \neq j} \mathbb{E}(\sum_{t=1}^i y_{t,j'}|x_t; \theta_{j'}) > \mathbb{E}(y_{t,j'}|x_t; \theta_{j'}) | \delta_1^{(i)} = g, \cdots, \delta_i^{(i)} = g \right),
$$
which can be conveniently computed using Gibbs sampling techniques describes in Section 2.2. The following stopping rule will be employed after the \( i \)th subject has been enrolled and evaluated for response:

1. If \( \max_{j=1,\ldots,J} p_{jg}^{(i)} > \delta \), for a pre-specified stopping criterion \( \delta \), e.g., 0.99 or 0.999, the trial is stopped after subject \( i \) has been evaluated for response. In this case, treatment \( j^* = \arg\max_{j=1,\ldots,J} p_{jg}^{(i)} \) for the biomarker profile group \( g \) is declared as superior over all other treatment-biomarker group combinations. Consequently, the treatment recommendation is that all subjects who are classified in biomarker profile group \( g \) according the final PLSLR model, should be treated with treatment \( j^* \).

2. If a pre-specified maximum sample size \( N_{max} \) is reached and \( p_{jg}^{(N_{max})} \leq \delta \) for all \( g = 1, \ldots, G \) and \( j = 1, \ldots, J \) then the trial is declared as inconclusive.

3. **SIMULATION STUDIES**

3.1 *Simulation Study I*

We performed a series of simulation studies to evaluate the operating characteristics of the proposed BCARA randomization design where subject’s memberships in biomarker subgroups were determined adaptively using the PLSLR approach. For each simulated hypothetical clinical trial, the basic scenario was a three-arm study with treatments A, B, and C. A set of \( K = 5 \) biomarker measurements were generated from a multivariate normal distribution with means 0, variances 1 and correlations \( \rho = 0.50 \). For the targeted therapy (treatment A), various levels of predictive power of the five biomarkers were assumed. Specifically, three different scenarios were considered. In the first scenario, a weak association between each of the five biomarkers and treatment response to the targeted therapy, i.e., treatment A, was assumed with a polyserial correlation of \( \rho_{ps} = 0.1 \). In the second scenario, a moderately strong association between each of the five biomarkers and treatment response was assumed with a polyserial correlation of \( \rho_{ps} = 0.5 \). In the last scenario, a strong
association was assumed with a polyserial correlation of $\rho_{ps} = 0.8$. Furthermore, it was assumed that the biomarkers were uncorrelated with treatment response for treatment B and C. The overall response rates for the targeted therapy were assumed to be 0.5, 0.6, and 0.7. The overall response rates for treatments B and C were assumed to be 0.4. The stopping criterion parameter for terminating the trial was fixed at 0.99 and 0.999, respectively. A maximum sample size of $N_{\text{max}} = 200$ was used for each hypothetical trial. The run-in phase, during which subjects were randomized to one of the three treatment arms using equal randomization, consisted of the first 24 subjects. Afterward, the BCARA randomization design described in Section 2.3 was used to randomize subjects into the three treatment arms where the prior distribution for $\theta_j$ was assumed to be normally distributed with mean zero and $\Sigma_j = 100 \mathbf{I}_5$ for $j = 1, 2, 3$. Biomarker groups were determined using the PLSLR approach as described in Section 2.4. Subjects were adaptively classified in biomarker positive or biomarker negative groups according the sign of first PLS component, i.e., subjects were classified as biomarker positive if $b_{it}^{(i)} > 0$ for $t = 1, \ldots, i$ and $i = n^* + 1, \ldots, N_{\text{max}}$ and as biomarker negative otherwise. Each scenario was simulated $M = 500$ times. The results are summarized in Table 1.

For the first scenario with $\rho_{ps} = 0.1$ and $\delta = 0.99$, the mean total sample size ranged from 86-145. As expected, more subjects from the biomarker positive group were randomized to the targeted treatment arm, i.e., 51% of the subjects with a positive biomarker profile were randomized to the targeted therapy arm when the overall response rate was 50%, and 46% of the subjects with a positive biomarker profile were randomized to that arm when the overall response rate was 70%. When the overall response of the targeted therapy was 50%, the probability that the targeted therapy in subjects with a positive biomarker profile was
declared as superior over all other treatment-biomarker group combinations was 25%. There was a 51% probability that the maximum sample size $N_{max} = 200$ was reached without declaring one of the treatment-biomarker groups as superior when the overall response rate of the targeted therapy was 50%. However, this probability decreased to 29% and 20% when the overall response rate of the targeted therapy increased to 60% and 70%, respectively. Changing the stopping criterion parameter $\delta$ from 0.99 to 0.999 resulted in a substantial increase in the mean total sample size. For example, when the overall response rate of the targeted therapy was 50%, the mean total sample size increased from 145 to 190. Furthermore, changing the stopping criterion parameter $\delta$ from 0.99 to 0.999 resulted in an increase in the probability of an inconclusive result. Under the second and third scenarios, i.e., assuming a moderately strong and strong association between biomarkers and response to the targeted therapy, the mean total sample sizes were substantially less. For example, assuming a strong association between biomarkers and treatment response and using a stopping criterion parameter $delta = 0.99$, the mean total sample sizes ranged from 66 to 86. There were no changes in the treatment allocations. However, there was a substantial increase in the sensitivity for the third scenario when compared to the first scenario. Specifically, assuming an overall response rate for the targeted therapy of 50% and a stopping criterion of $\delta = 0.99$, the probability of declaring the targeted therapy as superior in subjects with a positive biomarker profile increased from 25% under the first scenario to 57% under the second scenario and to 82% under the third scenario. Assuming a stopping criterion parameter of $\delta = 0.999$ and a strong association between biomarkers and treatment response for the targeted therapy, the probability of an inconclusive result increased from 5% to 17% when the overall response rate for the targeted therapy was 50%.
3.2 Simulation Study II

In a second simulation study, the operating characteristic of the proposed BCARA randomization design where biomarker subgroups are determined using the PLSLR approach was compared to the operating characteristic of a marker-by-treatment interaction design [6] with a Bayesian response adaptive randomization [24]. In the following, we will denote the first design as BCARA-PLSLR and the second design as MBTID-BRA. The response rate of a targeted therapy (treatment A) was compared to the response rate of an untargeted therapy (treatment B). The response rate of the untargeted therapy was 0.5 while the overall response rates for the targeted therapy were 0.6, 0.7, and 0.8. A set of \( K = 3 \) correlated (\( \rho = 0.3 \)) biomarker measurements were generated from a multivariate multinomial distribution with a scale ranging from 0-3. This a standard scale used for quantifying molecular markers based on immunohistochemistry (IHC) staining with 0 (none), 1 (low), 2 (medium), or 3 (high). The frequency for each of the 4 categories was assumed to be 25%. Various levels of predictive power of the three biomarkers on the probability of response for the targeted therapy were assumed, with polyserial correlations ranging from 0.1-0.8. Furthermore, it was assumed that there were no associations between biomarker measurements and response for the untargeted therapy. The maximum sample size for each trial was \( N_{\text{max}} = 200 \). Hypothetical trials were simulated \( M = 500 \) times. For the BCARA-PLSLR design, equal randomization was used for the first \( n^* = 10 \) subjects during the run-in phase. Biomarker subgroups were determined based on the sign of the first PLS component. The stopping criterion parameter \( \delta \) was fixed at 0.99. For the MBTID-BRA, a Beta(1,1) distribution was assumed as a prior distribution for the success probabilities for both treatment A and treatment B. The biomarker groups are defined as positive if at least one of the three biomarker has a value of 2 or higher and as negative if all values are 0 or 1, i.e., “none” or “low” category on the standard IHC scale. The randomization ratios for this design were
computed in the same fashion as in the BCARA-PLSLR design, i.e., subject $i + 1$ was randomized to treatment A with probability $RR^{(g)}_{i+1,A} = (\frac{p_A^{(g)}}{N_{\text{max}}})^c / \left( (\frac{p_A^{(g)}}{N_{\text{max}}})^c + (1 - (\frac{p_A^{(g)}}{N_{\text{max}}}))^c \right)$, where $c = (i + 1)/2N_{\text{max}}$ and $p_A^{(g)}$ denotes the probability that the response rate in treatment A is greater than in treatment B given the observed responses in group $g$ for subjects 1, ..., $i$. The same stopping rule as for the BCARA-PLSLR design was employed to terminate the trial with $\delta = 0.99$. The results of this simulation study are summarized in Table 2.

[Table 2 about here.]

As expected, the operating characteristics of the BCARA-PLSLR and the MBTID-BRA design were similar when the biomarkers are non-predictive and when the difference in response rates between the two arms is small. For example, in the scenario where the overall response rate of treatment A is 0.6 and the polyserial correlations between the three biomarkers and response are 0.1, the mean total sample size for the BCARA-PLSLR design was 92 while that for the MBTID-BRA design was 82. Under both designs, the probability that treatment A was declared as superior in the biomarker positive group was approximately 50%. There were substantial differences in the operating characteristics between the two designs when the levels of predictive power varied between the three biomarkers. For example, when the overall response rate for the targeted therapy was 0.7 and the polyserial correlations for the three biomarkers were 0.5, 0.1, and 0.1, respectively, the mean total sample size for the MBTID-BRA design was 76 while that for the BCARA-PLSLR design was 61. The probability that the targeted therapy was declared as superior in the biomarker positive group over all other treatment-biomarker group combinations was 0.69 for the BCARA-PLSLR design and 0.50 for the MBTID-BRA design. When the overall response rate for the targeted therapy was 0.8 and there were substantial differences in the level of predictive power between the three biomarkers, with polyserial correlations of 0.8, 0.1, and 0.1, the mean total
sample size for the MBTID-BRA design was 80 with a 38% probability that the targeted therapy was declared as superior in the biomarker positive group was 38% while the mean total sample size for the BCARA-PLSLR design was only 30 with a 90% probability that the targeted therapy was declared as superior in the biomarker positive group. In summary, the simulation studies show that the proposed BCARA-PLSLR design effectively identifies subjects who benefit most from a targeted therapy and that there may be substantial savings in the sample size requirements when compared to the MBTID-BRA design.

4. DISCUSSION

In this paper, we proposed a Bayesian covariate-adjusted response-adaptive design that utilizes information about individual subject’s biomarker profiles and clinical outcomes as they become available during the conduct of the trial. A computationally efficient algorithm using block Gibbs sampling techniques is implemented to predict patient’s responses to treatment and to compute allocation probabilities adaptively. Predictive biomarker subgroups are also determined adaptively using a PLSLR approach. Simulation studies were conducted to examine the operating characteristics. The simulation studies showed that the proposed design effectively identifies subjects who benefit most from a targeted therapy and that there may be substantial savings in the sample size requirements when compared to alternative designs.

The proposed combination of an adaptive design strategy, Bayesian approach, and biomarker classification is not meant to replace the traditional paradigm for drug development. We conclude, however, that it may serve a useful role in the early efficacy phase of targeted therapy development.

5. ACKNOWLEDGMENTS

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NIH/NCI grant P30 CA14520.
REFERENCES


Table 1
Operating characteristics of BCARA randomization design where biomarker subgroups are determined adaptively using the PLSLR approach.

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<th>ρ_A^2</th>
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<th>RR_A-</th>
<th>δ</th>
<th>N^S</th>
<th>P_A+^P</th>
<th>N_A^P</th>
<th>Prob_A+^P</th>
<th>Prob_I^P</th>
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1 Overall response rate of targeted therapy (treatment A)
2 Polyserial correlation between response and individual biomarkers for treatment A
3 Response rate of treatment A in biomarker positive group
4 Response rate of treatment A in biomarker negative group
5 Mean total sample size across 500 replications
6 Proportion of subjects with positive biomarker profile randomized to treatment A
7 Mean number of subjects randomized to treatment A
8 Probability of declaring treatment A in subjects with positive biomarker profile superior
9 Probability of inconclusive result
### Table 2
Comparison of operating characteristics between proposed BCARA-PLSLR design and MBTID-BRA design.

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1. Overall response rate of targeted therapy (treatment A)
2. Polyserial correlations between response and biomarkers 1, 2, and 3 for treatment A
3. Mean total sample size across 500 replications
4. Probability of declaring treatment A in subjects with positive biomarker profile superior
5. Probability of declaring treatment A in subjects with negative biomarker profile superior
6. Probability of declaring treatment B in subjects with positive biomarker profile superior
7. Probability of declaring treatment B in subjects with negative biomarker profile superior
8. Probability of inconclusive result
9. Bayesian covariate-adjusted response-adaptive randomization design with PLSR to determine predictive biomarker subgroups
10. Marker-by-treatment interaction design using a Bayesian response adaptive randomization