# Fully Bayesian Optimal Experimental Design: A Review

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## Abstract

Bayesian experimental design is a fast growing area of research with many real-world applications. As computational power has increased over the years, so has the development of simulation-based design methods, which involve a number of algorithms, such as Markov chain Monte Carlo, sequential Monte Carlo and approximate Bayes methods, and which have enabled more complex design problems to be solved. This paper provides an overview of the literature on Bayesian experimental design that uses a decision-theoretic approach. The Bayesian framework provides a unified approach for incorporating prior information and/or uncertainties regarding the statistical model with a utility function which describes the experimental aims. In this paper, we provide a general overview on the concepts involved in Bayesian experimental design, and focus on describing some of the more commonly-used Bayesian utility functions and methods for their estimation, as well as a number of algorithms that are used to search over the design space to find the optimal Bayesian design. We also provide some examples from the literature of real-world applications and discuss future directions for Bayesian experimental design.

KEYWORDS: Bayesian optimal design; Decision theory; Utility function; Stochastic optimisation; Posterior distribution approximation.

# 1 Introduction

## 1.1 Background

Statistical experimental design provides rules for the allocation of resources in an information gathering exercise in which there is variability that is not under control of the experimenter. Experimental design has very broad applications across the natural, medical and social sciences, as well as engineering, business and finance. Experimental design reflects the purpose of the experiment. Prior to the commencement of an experiment, experimental design often requires choices to be made regarding which treatments to study and how these treatments will be

defined or administered (e.g., amount, timing, frequency), and the proportion of observations to allocate to each treatment. Experimental design can also require choices of blocking factors, randomisation methods and sample size to be made. The experimental units must also be clearly defined prior to the commencement of the experiment, along with the time period over which the experiment is to be performed. Due to costs, ethics and other constraints on time, efficient use of resources is highly critical.

Experimental designs incorporate features into studies with the aim to control systematic error (bias), reduce random variations, increase precision of parameter estimates (or some measure of interest), make predictions about future observations, or discriminate between competing models. Essentially, non-optimal designs require more resources to make inferences on the features of interest with the same level of reward that an optimal design would. Experimental design problems are commonly viewed as optimisation problems, and optimal experimental designs may be used to achieve the experimental goals more rapidly and hence reduce experimental costs.

Experimental design has been widely developed within the classical framework, in both theory and practice (e.g., Atkinson and Donev [1992]). In the classical framework, optimal experimental designs are commonly derived using optimality criteria that are based on the expected Fisher information matrix (e.g., Fedorov [1972], Pukelsheim and Torsney [1991], Atkinson and Donev [1992]).

Classical experimental design is well suited to linear or linearised models. For nonlinear models, designs are dependent on the values which are chosen for the model parameters. Often, the aim of experimental design is to precisely estimate model parameters. Therefore, selection of the parameter values from which to construct the design is highly important and use of unsuitable parameter values may result in sub-optimal designs. Only locally optimal designs can be obtained in the classical framework for nonlinear models. Several studies have incorporated probability distributions on the model parameters and averaged local design criteria over the distributions so that the designs obtained may be robust to the initial choice of the parameter values (e.g., Pronzato and Walter [1985], D'Argenio [1990]). These probability distributions are known as *prior distributions* and can incorporate information from previous studies, expert elicited data or subjective beliefs of the experimenters. Similar methods are also used for situations in which there is model uncertainty.

It is a common misconception in the experimental design literature that designs which have arisen from averaging classical design criteria over prior distributions are termed "Bayesian designs". We propose that to qualify as a "fully Bayesian design", one must obtain the design by using a design criterion that is a functional of the posterior distribution. Designs which have arisen from averaging the classical design criteria over the parameter space are termed "pseudo-Bayesian", "on average" or "robust" designs (Pronzato and Walter [1985], Federov and Hackl [1997]).

Bayesian methodologies for optimal experimental design have become more prominent in the literature (e.g., Müller [1999], Han and Chaloner [2004], Amzal et al. [2006], Müller et al. [2006], Cook et al. [2008], Huan and

Marzouk [2013]). One advantage of using a Bayesian design criterion is that a single design point can be used, and the prior distribution is updated by the single observation. Lindley (1972) presents a decision theoretic approach to experimental design, upon which Bayesian experimental design is based. Bayesian optimal design involves defining a design criterion, or a utility function $U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})$, that describes the worth (based on the experimental aims) of choosing the design $\mathbf{d}$ from the design space $\mathbf{D}$ yielding data $\mathbf{y}$, with model parameter values $\boldsymbol{\theta}$. A probabilistic model, $p(\boldsymbol{\theta}, \mathbf{y}|\mathbf{d})$, is also required. This consists of a likelihood $p(\mathbf{y}|\mathbf{d}, \boldsymbol{\theta})$ for observing a new set of measurements $\mathbf{y}$ at the design points $\mathbf{d}$, given parameter values $\boldsymbol{\theta}$, and a prior distribution $p(\boldsymbol{\theta})$ for the parameters $\boldsymbol{\theta}$. The prior distribution is usually assumed to be independent of the design $\mathbf{d}$.

The Bayesian optimal design, $\mathbf{d}^*$, maximises the expected utility function $U(\mathbf{d})$ over the design space $\mathbf{D}$ with respect to the future data $\mathbf{y}$ and model parameters $\boldsymbol{\theta}$:

$$
\begin{aligned}
\mathbf{d}^* &= \arg\max_{\mathbf{d} \in \mathbf{D}} \ E\{U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})\} \\
&= \arg\max_{\mathbf{d} \in \mathbf{D}} \ \int_{\mathbf{Y}} \int_{\boldsymbol{\Theta}} U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y}) p(\boldsymbol{\theta}, \mathbf{y}|\mathbf{d}) d\boldsymbol{\theta} d\mathbf{y} \\
&= \arg\max_{\mathbf{d} \in \mathbf{D}} \ \int_{\mathbf{Y}} \left\{ \int_{\boldsymbol{\Theta}} U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y}) p(\boldsymbol{\theta}|\mathbf{d}, \mathbf{y}) d\boldsymbol{\theta} \right\} p(\mathbf{y}|\mathbf{d}) d\mathbf{y}.
\end{aligned}
\tag{1}
$$

Thus, the optimal design (given the observed data), maximises the posterior expected utility (defined in {} in equation (1)). Unless the likelihood and prior are specifically chosen to enable analytic evaluation of the integration problem, equation (1) does not usually have a closed form solution. Therefore, numerical approximations or stochastic solution methods are required to solve the maximisation and integration problem.

Due to the computational challenges of performing the integration and maximisation of equation (1), the use of standard optimisation algorithms, such as the Newton-Raphson method, to find the optimal design is inappropriate. A number of stochastic algorithms have been proposed in the literature to approximate the maximisation and integration problem of equation (1). These include: prior simulation (Müller [1999]); smoothing of Monte Carlo simulations (Müller [1999]); gridding methods which involve numerical quadrature or Laplace approximations to perform backward induction (Brockwell and Kadane [2003]); Markov chain Monte Carlo simulation in an augmented probability model (Müller [1999]); and sequential Monte Carlo methods (Kück et al. [2006], Amzal et al. [2006]). These algorithms will be discussed further in Sections 5 and 6.

## 1.2 Brief History of the Bayesian Design Literature

A broad range of literature exists on optimal experimental design. This article aims to review those papers which present solutions to fully Bayesian experimental design problems.

Early work on Bayesian decision-theory includes Lindley (1968, 1972), which notes that the design of an experiment should depend on the experimental objectives (e.g., precise estimation of certain parameters, predic-

tion of future responses). Other work includes Chaloner (1984), who further developed Bayesian optimal design theory in a linear regression context and explicitly describes how the prior causes a difference between the classical and Bayesian optimal designs. DasGupta and Studden (1991) gave a structured formulation to demonstrate the sensitivity of designs to the priors and presented designs that were robust to the prior specification. Other notable works on Bayesian designs for linear regression models include Pilz (1991) and El-Krunz and Studden (1991). Simulation-based design methods have frequently been used more recently (e.g., Clyde et al. [1996], Bielza et al. [1999], Müller [1999], Stroud et al. [2001], Amzal et al. [2006], Müller et al. [2006], Cook et al. [2008], Cavagnaro et al. [2010]) in which Markov chain Monte Carlo and sequential Monte Carlo algorithms are utilised to solve complex optimal Bayesian design problems (e.g., designing for nonlinear models). Sequential, or adaptive designs, have become increasingly popular in the Bayesian design literature as they provide flexible and efficient designs. Rather than using the same design throughout the experimental process, as in *static* design problems, the design which maximises the expected utility is chosen at each stage of experimentation, based on the outcomes of previous experiments. Recent developments in static and sequential designs will be discussed further in Sections 5 and 6.

There are already several notable review papers on Bayesian experimental design. DasGupta (1995) presents a review of both classical and Bayesian experimental design, with a focus on designing for linear models. Atkinson (1996) review classical and pseudo-Bayesian optimal design for linear and nonlinear models. Verdinelli (1992) and Chaloner and Verdinelli (1995) present a comprehensive review on Bayesian experimental design, for both linear and nonlinear models. Müller (1999) provides an overview of simulation-based methods in optimal design. Clyde (2001) presents a broad review on several of the key concepts involved in Bayesian experimental design, such as, choice of utility functions; prior elicitation; and methods for calculating the expected utility.

## 1.3   Contribution and Outline

There has been a lack in review papers on fully Bayesian experimental design since the early 2000s. These earlier review papers have often been written from a rather mathematical view point, and have often focused on defining Bayesian design criteria and their relationship to classical design criteria. In the past two decades there has been a substantial increase in computational power and, along with it, the use of Bayesian methodologies for optimal design. At the present time, we have been unable to find any recent review articles which discuss the various algorithms that are used in the Bayesian design literature to solve optimal design problems. Designs for complex models have also received little attention in Bayesian experimental design literature reviews. This article is concerned with reviewing the computational methods that have been used to find fully Bayesian experimental designs and aims to address the aspects of Bayesian experimental design which have received little or no emphasis in previous review papers. This article is aimed at readers with some understanding of Bayesian methods, but

not necessarily with knowledge of experimental design.

In Section 2 we describe how the prior distribution has been elicited in previous Bayesian experimental design studies. Section 3 discusses methods for posterior distribution approximation for use in Bayesian utility functions. In Section 4 we discuss some of the more commonly-used Bayesian utility functions, along with the methods that have been used for their estimation. Sections 5 and 6 provide an overview of the optimisation algorithms that have been used to search for static and sequential Bayesian experimental designs, respectively. We provide some real-world examples of Bayesian experimental designs in Section 7, discuss future directions of Bayesian experimental design in Section 8 and provide a conclusion in Section 9.

## 2   The Prior Distribution

The Bayesian design framework (as well as Bayesian analysis) requires the elicitation of a prior distribution for the statistical model(s). The requirement of having to specify a prior distribution, upon which the conclusions depend, has caused many practitioners to be reluctant to use Bayesian experimental design methods. It is very important to check the sensitivity of the optimal design to the specification of the prior distribution (see, for example, DasGupta and Studden [1991]).

A number of studies (e.g., Clyde et al. [1996], Stroud et al. [2001], Ryan et al. [2014a]) have used historical data from previous experiments to construct a prior distribution for the design of future experiments. Toman and Gastwirth (1994) suggest the use of results from a pilot study to specify the prior distribution. Tsai and Chaloner (2002) used information from over 50 clinical experts to elicit prior distributions for their design problem. Kadane (1996) discusses a number of the practical issues that occur in subjective elicitation for clinical trials.

Several studies have considered the problem where the prior used for the design phase is different from the prior used for analysis (e.g., Etzione and Kadane [1993], Han and Chaloner [2004]). For example, a noninformative prior may be used for analysis to mimic a classical analysis, but all available prior information may be used in the design process so that an informative prior may be used.

## 3   Estimation of the Posterior Distribution

Bayesian utility functions are based on the posterior distribution and generally assume that a Bayesian analysis will be performed on any data that are generated from the experimental design. In general, the posterior distribution does not have a closed form expression, and numerical methods are required to sample from or approximate the posterior distribution. Generally, thousands of posterior distributions need to be considered, since each possible future data set that is drawn from the prior predictive distribution requires calculations of the posterior distribution. For this to be computationally feasible, rapid methods for obtaining the posterior

distributions for many datasets that are drawn from the same model and prior are required.

## 3.1 Markov Chain Monte Carlo

Markov chain Monte Carlo (MCMC) has often been used to approximate the posterior distribution for Bayesian utility function calculations (e.g., Wakefield [1994], Palmer and Müller [1998], Han and Chaloner [2004]). Although MCMC is often appropriate and useful for Bayesian data analysis, it can be too computationally intensive to perform MCMC to approximate the posterior distribution for each of the thousands of iterations required in the Bayesian experimental design algorithms.

## 3.2 Importance Sampling

Importance sampling is a popular method for approximating target distributions of interest, from which it may be difficult to sample (Geweke [1989]). Importance sampling involves choosing an *importance distribution* $g(\cdot)$, from which it is easy to sample, and then appropriately weighting the samples that have been drawn from the importance distribution to account for the discrepancy between $g(\cdot)$ and the target distribution. In the Bayesian design context, the target distribution is the posterior $p(\boldsymbol{\theta}|\mathbf{d}, \mathbf{y})$. Weighted samples $\{\boldsymbol{\theta}_k, W_k\}_{k=1}^{N_p}$ are produced, where $N_p$ is the number of particles used to approximate the posterior; $w(\boldsymbol{\theta}) = \frac{p(\mathbf{y}|\mathbf{d}, \boldsymbol{\theta})p(\boldsymbol{\theta})}{g(\boldsymbol{\theta})}$ are the importance weights; and $W_k \propto w(\boldsymbol{\theta}_k), k = 1, ..., N_p$ are the normalised importance weights, $\sum_{k=1}^{N_p} W_k = 1$. The target and importance distributions should have the same support. To measure the efficiency of importance sampling, the effective sample size (ESS) is used and can be approximated via

$$ESS = \frac{1}{\sum_{k=1}^{N_p} W_k^2}, 1 \le ESS \le N_p.$$

Importance sampling is a very useful method for approximating the posterior distribution in Bayesian experimental design since the importance samples only need to be drawn once (unlike MCMC) and can then be re-weighted in each iteration of the optimisation algorithm according to the current design and data. The ability to re-use the importance samples offers substantial computational savings.

Importance sampling from the prior distribution has commonly been used in Bayesian experimental design to approximate the posterior distribution (e.g., Cook et al. [2008], McGree et al. [2012c], Ryan et al. [2014a,c]). This reduces the importance weights to the likelihood function. However, this is usually inefficient for large amounts of data since the posterior distribution is very different from the prior (e.g., Bengtsson et al. [2008], Ryan et al. [2014a,c]). Importance sampling from the prior may also not be useful for diffuse priors (Chopin [2002]) or when the model parameter is high dimensional.

Ryan et al. (2014a) used Laplace approximations (to the posterior) to form the importance distribution for importance sampling, and found that this approach corrects for some non-normality that is not accommodated

by the Laplace approximation, and can also be used when large amounts of data are involved in the design problem since fewer particles are required in the importance sampling to obtain a reasonable ESS.

The use of adaptive importance sampling (e.g., Kinas [1996], Pennanen and Koivu [2006]) is largely unexplored for approximating the posterior distribution in Bayesian experimental design problems.

## 3.3 Deterministic Approximations

Laplace approximations (or Gaussian approximations) and numerical quadrature provide fast methods for obtaining approximations to the posterior distribution in Bayesian design problems (e.g., Lewi et al. [2009], Cavagnaro et al. [2010], Bornkamp et al. [2011], Long et al. [2013], Ryan et al. [2014a]). These methods are particularly useful when large amounts of data are involved. However, their suitability depends on whether it is reasonable to assume that the posterior distribution is well approximated by a multivariate normal distribution and they also suffer from the curse of dimensionality. To overcome the issue of dimensionality, Long et al. (2013) use polynomial-based sparse quadrature for the integration over the prior distribution.

Integrated nested Laplace approximation (INLA) is a relatively new method for rapidly approximating posterior distributions (see Rue et al. [2009]). INLA generally is a significantly faster alternative to MCMC and importance sampling for approximating the posterior. To date, INLA has mostly been used for approximate posterior inference for models in which the posterior marginals are not available in closed form due to non-Gaussian response variables, such as latent Gaussian Markov random field (GMRF) models (e.g., Rue et al. [2009]) with non-Gaussian observations. INLA enables fast Bayesian inference by using accurate approximations to the marginal posterior density for the hyperparameters and the posterior marginal densities for the latent variables. The use of INLA in the context of Bayesian experimental design is currently unexplored.

Variational Bayesian (VB) methods facilitate approximate inference for intractable posteriors (or other target densities) and provide an alternative to other approaches for approximate Bayesian inference such as MCMC and Laplace approximations. VB can also be used to determine a lower bound for the evidence for use in model selection problems. The VB approach is fast and deterministic, and involves approximating the intractable target densities, e.g., $p(\boldsymbol{\theta}|\mathbf{y})$, by a factored form $q(\boldsymbol{\theta}) = q_1(\boldsymbol{\theta}_1) \times .... \times q_r(\boldsymbol{\theta}_r)$, for which $q(\boldsymbol{\theta})$ is more tractable than $p(\boldsymbol{\theta}|\mathbf{y})$. The $q_j$ are found iteratively from Gibbs sampling like expectations (see, for example, Ormerod and Wand [2010]). An issue is the factorization for the variational approximation $q(\cdot)$. Variational approximations have commonly been used for Bayesian inference (e.g., Ormerod and Wand [2010]), but have not yet been used in a Bayesian experimental design context. These methods could provide a fast alternative for approximating the posterior for use in Bayesian utility function calculation. However, the error of the VB approximation is generally unknown and can be substantial (e.g., Rue et al. [2009]).

## 3.4 Approximate Bayesian Computation

Approximate Bayesian computation (ABC) is a likelihood-free method that is used to approximate the posterior distribution in situations where the likelihood function is intractable, but simulation from the likelihood is relatively straightforward. ABC has commonly been used to perform inference (e.g., Drovandi and Pettitt [2011], Drovandi et al. [2011], Sisson and Fan [2011]). One of the most common ABC algorithms is ABC rejection (see Beaumont et al. [2002]). ABC rejection prevents one from having to evaluate the likelihood by instead drawing many parameter values from the prior, and simulating data from the model, conditional on those parameter values. Only those parameters that generate simulated data that are close in some sense to the observed data are kept. The efficiency of this method is dependent on how close the posterior distribution is to the prior.

Drovandi and Pettitt (2013) and Hainy et al. (2013) used ABC rejection in the Bayesian experimental design context to approximate the posterior distributions (for Bayesian utility function calculation) for models with computationally intractable likelihoods. The ABC posterior is given by:

$$p(\boldsymbol{\theta}|\mathbf{d},\mathbf{y},\epsilon) = \int_{\mathbf{x}} p(\mathbf{x}|\mathbf{d},\boldsymbol{\theta})p(\boldsymbol{\theta})1(\rho(\mathbf{y},\mathbf{x}) \leq \epsilon)d\mathbf{x},$$

where $\mathbf{y}$ represents the 'observed data' (that is generated from the model at each iteration of the optimisation (e.g., MCMC) algorithm); $\mathbf{x}$ is simulated data; $1(\cdot)$ is an indicator function; $\rho(\cdot,\cdot)$ is function that measures the discrepancy between the observed and simulated data; and $\epsilon$ is a tolerance threshold that controls the error of the approximation. The discrepancy function typically compares summary statistics of the observed and simulated data. However, Drovandi and Pettitt (2013) only considered low dimensional designs and so they were able to compare the observed and simulated data directly. ABC rejection is very useful since the ABC data, i.e., the $\mathbf{x}$ values, as well as the model parameters $\boldsymbol{\theta}$, only need to be simulated once and can be re-used at each iteration of the optimisation algorithm (much in the same spirit as importance sampling) for comparison to the observed data, $\mathbf{y}$. This offers substantial computational savings.

# 4 Bayesian Utility Functions and Methods for Their Estimation

It is highly important that the utility function incorporates the experimental aims and is specific to the application of interest. For instance, designs which efficiently estimate the model parameters may not be useful for prediction of future outcomes. Several approaches have been suggested in the literature to assist in the elicitation of the utility function (see Spiegelhalter et al. [1996], Wolfson et al. [1996]). In practice, the utility function is often not specified as a single function, due to the difficulty of combining competing goals, and instead a set of possible utility functions is used. Christen *et al.* (2004) formally acknowledged the fact that the decision maker may be unwilling or unable to specify a unique utility function by considering a set of possible utility functions.

Sensitivity analyses to misspecifications in the utility function have been proposed (see Rios Insua and Ruggeri [2000] for a review). In this section we will discuss some of the more commonly used Bayesian utility functions, as well as methods for their estimation based on the approximation to the posterior. One of the most commonly used and versatile Bayesian design criteria is the mutual information, which is based on entropy, and has been used for designing for efficient parameter estimation (Bernardo [1979], Ryan [2003], Paninski [2005]), as well as minimising prediction uncertainty (Liepe et al. [2013]), and model discrimination (Box and Hill [1967], Ng and Chick [2004], Cavagnaro et al. [2010], Drovandi et al. [2014]). For discussion of other Bayesian utility functions, see Chaloner and Verdinelli (1995).

For normal linear models, analytical expressions for equation (1) can be obtained for many Bayesian utilities, provided the model dimension and decision space is small (e.g., Borth [1975], Chaloner and Verdinelli [1995], Ng and Chick [2004]). For nonlinear design problems, one cannot usually obtain an analytical expression, and the integrals in equation (1) can instead be approximated by Monte Carlo methods (e.g., Palmer and Müller [1998], Cook et al. [2008], Ryan et al. [2014a]), Laplace approximations (e.g., Lewi et al. [2009], Ryan et al. [2014a]), or numerical quadrature (e.g., Cavagnaro et al. [2010]).

## 4.1 Parameter Estimation Utility Functions

Precise parameter estimation is a common goal of experimental design and many different utility functions have been used to achieve this purpose. Bayesian utility functions that design for precise parameter estimation are discussed below.

### 4.1.1 Information-based Utilities

When interest lies in estimating some function of $\boldsymbol{\theta}$, say $\phi(\boldsymbol{\theta})$, the mutual information between $\phi(\boldsymbol{\theta})$ and the data $\mathbf{y}$, conditional on the design $\mathbf{d}$, may be given by:

$$I(\phi(\boldsymbol{\theta}); \mathbf{y}|\mathbf{d}) = U(\mathbf{d}) = \int_{\phi(\boldsymbol{\theta})} \int_{\mathbf{Y}} p(\phi(\boldsymbol{\theta}), \mathbf{y}|\mathbf{d}) \bigg[ \log p(\phi(\boldsymbol{\theta}), \mathbf{y}|\mathbf{d}) - \log p(\mathbf{y}|\mathbf{d}) - \log p(\phi(\boldsymbol{\theta})) \bigg] d\mathbf{y} d\phi(\boldsymbol{\theta}). \qquad (2)$$

The optimal design that maximises the utility function is the one that yields the largest information gain, on average, about $\phi(\boldsymbol{\theta})$ upon observation of the data.

Another commonly-used Bayesian design criterion is the Kullback-Leibler divergence (KLD) (Kullback and Leibler [1951]) between the prior and posterior distributions, which is given by:

$$\begin{aligned} U(\mathbf{d}, \mathbf{y}) &= E_{\phi(\boldsymbol{\theta})|\mathbf{d},\mathbf{y}}(\log p(\phi(\boldsymbol{\theta})|\mathbf{d}, \mathbf{y}) - \log p(\phi(\boldsymbol{\theta}))) \\ &= \int_{\phi(\boldsymbol{\theta})} p(\phi(\boldsymbol{\theta})|\mathbf{d}, \mathbf{y}) \log p(\mathbf{y}|\mathbf{d}, \phi(\boldsymbol{\theta})) d\phi(\boldsymbol{\theta}) - \log p(\mathbf{y}|\mathbf{d}). \end{aligned} \qquad (3)$$

Lindley (1956) suggested that this utility should be used if one is interested in maximising the expected information gain on the model parameters (or functions of) due to performing an experiment at design points **d**. Mathematically, the mutual information is the KLD between the joint distribution $p(\boldsymbol{\theta}, \mathbf{y}|\mathbf{d})$ and product of marginal distributions of $\boldsymbol{\theta}$ and $\mathbf{y}$ (Borth [1975]). This criterion is equivalent to the classical $D$-optimal criterion when designing for normal linear models with a normal prior distribution for the model parameter (see Chaloner and Verdinelli [1995] and Verdinelli [2000] for further details).

Ryan (2003) used mutual information to find static designs for efficient parameter estimation. Kim et al. (2013) used the mutual information utility to find sequential designs to efficiently estimate parameters, which was of the form:

$$U(\mathbf{d}_{(t)}) = \int_{\boldsymbol{\Theta}} \int_{\mathbf{Y}} \left[\log\left(\frac{p(\boldsymbol{\theta}|\mathbf{d}_{(t)}, \mathbf{y}_{(1:t)})}{p(\boldsymbol{\theta}|\mathbf{y}_{(1:t-1)})}\right)\right] p(\mathbf{y}_{(t)}|\mathbf{d}_{(t)}, \boldsymbol{\theta}) p(\boldsymbol{\theta}|\mathbf{y}_{(1:t-1)}) d\mathbf{y}_{(t)} d\boldsymbol{\theta},$$

where $\mathbf{y}_{(1:t)}$ are the data that were observed from the 1st to the $t$-th trial, $\mathbf{y}_{(t)}$ are the data that were observed at the current, $t$-th trial, using design $\mathbf{d}_t$, $\mathbf{y}_{(1:t-1)}$ are the data that were measured from the 1st to the $(t-1)$-th trials using the designs $\mathbf{d}_{(1:t-1)}$. Paninski (2005) proved that under acceptably weak modelling conditions, utility functions based on mutual information can choose designs that lead to consistent and efficient parameter estimates in the adaptive design framework.

Despite the theoretical appeal, mutual information is computationally complex, due to the difficulty in calculating the evidence $p(\mathbf{y}|\mathbf{d})$ in equation (2). Therefore, many design problems have been restricted to special cases, such as designing for parameter estimation of linear gaussian models (e.g., Lewi et al. [2009]) or binary models (e.g., Kujala and Lukka [2006]) in which the evidence can be computed analytically. Conjugate priors have been used to obtain analytic results (e.g., Borth [1975]) and numerical quadrature has also been used (e.g., Cavagnaro et al. [2010]). Drovandi et al. (2013) used sequential Monte Carlo algorithms (which are described in more detail in Section 5) for both posterior and evidence approximation so that the mutual information could be calculated for sequential design problems for parameter estimation. Ryan et al. (2014c) used importance sampling to calculate the KLD between the prior and posterior distributions for static design problems, but found this to be computationally intensive. Huan and Marzouk (2012, 2013) used polynomial chaos approximations and nested Monte Carlo integration (Ryan [2003]) to estimate the KLD between the prior and posterior distributions for static design problems for parameter estimation.

### 4.1.2  Scalar Functions of the Posterior Covariance Matrix

The inverse of the determinant of the posterior covariance matrix is a useful utility function if one is interested in maximising the (joint) posterior precision of all (or a subset) of the model parameters $\boldsymbol{\theta}$ (e.g., Drovandi et al. [2013], Ryan et al. [2014c]) or a function of the model parameters $\phi(\boldsymbol{\theta})$ (e.g., Stroud et al. [2001], Drovandi et al.

[2013], Ryan et al. [2014a]). This utility is also known as the 'Bayesian D-posterior precision' and is given by:

$$U(\mathbf{d}, \mathbf{y}) = \frac{1}{\det(cov(\phi(\boldsymbol{\theta})|\mathbf{d}, \mathbf{y}))}.$$

If one were interested in maximising the precision of the marginal posterior distributions of the model parameters, then one should use the trace instead of the determinant to obtain the Bayesian A-posterior precision. The Bayesian D-posterior precision is much less computationally intensive to estimate than equation (2). However, if the posterior distribution is multi-modal, then use of the Bayesian D-posterior precision utility may be inappropriate and one should instead use equation (2) as the utility function.

The posterior variance-covariance matrix can easily be obtained from the weighted posterior samples that are obtained from importance sampling (e.g., Stroud et al. [2001], McGree et al. [2012c], Ryan et al. [2014a]), ABC rejection (e.g., Drovandi and Pettitt [2013]) and via sequential Monte Carlo (Drovandi et al. [2013]). The posterior variance-covariance matrix is also easily obtained when one uses numerical quadrature or Laplace approximations to the posterior distribution.

### 4.1.3 Quadratic Loss

When one is interested in obtaining a point estimate of the parameters, or linear combinations of them, a quadratic loss function may provide a suitable utility function:

$$U(\mathbf{d}, \mathbf{y}) = -\int_{\phi(\boldsymbol{\theta})} (\phi(\boldsymbol{\theta}) - \widehat{\phi(\boldsymbol{\theta})})^T \mathbf{A} (\phi(\boldsymbol{\theta}) - \widehat{\phi(\boldsymbol{\theta})}) p(\phi(\boldsymbol{\theta})|\mathbf{d}, \mathbf{y}) d\phi(\boldsymbol{\theta}),$$

where $\mathbf{A}$ is a symmetric non-negative definite matrix (e.g., Chaloner [1984], Chaloner and Verdinelli [1995], Han and Chaloner [2004]) and $\widehat{\phi(\boldsymbol{\theta})}$ is some estimate (e.g., the mean) of $p(\phi(\boldsymbol{\theta})|\mathbf{d}, \mathbf{y})$. Once the posterior distribution has been approximated, it is quite straightforward to estimate this utility. For normal linear models, when one is interested in point estimates of parameters, this utility is the Bayesian equivalent of the classical $A$-optimal criterion. When one is interested in linear combinations of the parameters, this utility is the Bayesian equivalent of the classical $c$-optimal criterion (for normal linear models).

## 4.2 Utilities for Model Discrimination

Model discrimination is an important experimental design problem which has generated a substantial amount of research (see, for example, Box and Hill [1967], Hill et al. [1968], Borth [1975], Cavagnaro et al. [2010], Drovandi et al. [2014]). Much of the design literature has focused on producing designs that offer efficient and precise parameter estimates. However, these designs can perform poorly on model discrimination problems (see, for example Atkinson [2008], Waterhouse et al. [2009]).

Mutual information has commonly been used as the utility function in the Bayesian design literature to design for model discrimination (e.g., Box and Hill [1967], Borth [1975], Ng and Chick [2004], Cavagnaro et al. [2010], Drovandi et al. [2014], McGree et al. [2012b]). The optimal design $\mathbf{d}$ is the one that maximises the mutual information between the (random variable) model indicator, $m$, and the future observation $\mathbf{y}$ (see, for example, Cavagnaro et al. [2010]). Drovandi et al. (2014) give an expression of this utility to design for model discrimination for discrete data, and McGree et al. (2012b) provide an expression for continuous data. Both Drovandi et al. (2014) and McGree et al. (2012b) used sequential Monte Carlo methods to approximate the necessary quantities so that mutual information could be used to obtain sequential designs for model discrimination.

Roth (1965) proposed a model discrimination utility that is known as 'total separation', and selects design points that yield the largest differences between the posterior predictive means of rival models. This is achieved by maximising a weighted sum (over all of the potential models) of the product of the absolute differences between the posterior predicted mean responses from all rival models and the given ('true') model. Total separation has recently been used by Masoumi et al. (2013) and McGree et al. (2012b) to design for model discrimination. The total separation utility can be approximated quite easily once the posterior predictive distribution has been found (see, for example McGree et al. [2012b]). This utility does not account for the variance of the predicted responses (Hill [1978]), which is problematic if the competing models differ in their error structures (e.g., additive vs. multiplicative error) (McGree et al. [2012b]).

Both mutual information and total separation do not rely on the assumption of a particular model being true (unlike many of the classical design criteria), but require the experimenter to define a set of rival models with prior probability of being true. That is, these utilities use the $M$-closed approach of Bernardo and Smith (2000, chapter 6).

Vanlier et al. (2014) proposed a model discrimination utility that is based on a k-nearest neighbour estimate of the Jensen Shannon divergence (which is the averaged KLD between the probability densities and their mixture) between the multivariate predictive densities of competing models. They showed that their utility is monotonically related to the expected change in the Bayes Factor in favour of the model that generated the data. MCMC was used to sample from the posterior distributions and the predictive distributions were sampled using these posterior distribution values and by adding noise generated by the error model. This was found to be computationally intensive, especially for their application which involved nonlinear models of biochemical reaction networks.

## 4.3   Utilities for Prediction of Future Observations

If one is interested in choosing $\mathbf{d}$ to predict $\mathbf{y}_{n+1}$ from $\mathbf{y} = (\mathbf{y}_1, ..., \mathbf{y}_n)$, then the expected gain in Shannon information for a future observation, $\mathbf{y}_{n+1}$, from the prior predictive distribution to the posterior predictive

distribution can be used as the utility function:

$$U(\mathbf{d}_{(n+1)}, \mathbf{y}) = \int_{\boldsymbol{\theta}} \int_{\mathbf{Y}_{n+1}} p(\mathbf{y}_{n+1}|\mathbf{d}_{n+1}, \mathbf{y}_{1:n}, \boldsymbol{\theta}) \log p(\mathbf{y}_{n+1}|\mathbf{d}_{n+1}, \mathbf{y}_{1:n}, \boldsymbol{\theta}) d\mathbf{y}_{n+1} d\boldsymbol{\theta} - \log p(\mathbf{y}_{1:n}|\mathbf{d}_{1:n}),$$

(e.g., Chaloner and Verdinelli [1995] and references therein). This is equivalent to the mutual information between the future observation $\mathbf{y}_{n+1}$ and the previous observations $\mathbf{y}_{1:n}$, conditional on the future designs $\mathbf{d}_{n+1}$ and previous designs $\mathbf{d}_{1:n}$. For normal linear models, this criterion is related to the classical $c$-optimal criterion (which maximises the precision of estimates of linear combinations of model parameters for linear models). Leipe et al. (2013) used mutual information to minimise prediction uncertainty in sequential systems biology experiments. Zidek et al. (2000) used maximum entropy to obtain designs that maximised information about expected responses for air quality monitoring sites.

Geostatistical design problems often use utilities that are functions of the prediction variance. For example, Diggle and Lophaven (2006) propose a Bayesian design criterion that chooses a set of sampling locations to enable efficient spatial prediction by minimising the expectation of the spatially averaged prediction variance (with respect to the marginal distribution of the data).

If one is interested in minimising the variance of the expected response, then one could use the utility function developed by Solonen et al. (2012) which places the next design point where the prior variance of the mean response is largest. The utility is calculated by bringing in the observations one-at-a-time and is given by:

$$U(\mathbf{d}, \mathbf{y}) = \prod_{k=1}^{K} (\sigma^2 + Var_{\boldsymbol{\theta}|\mathbf{y}_{1:(k-1)}}(m_k(\boldsymbol{\theta}))), \tag{4}$$

where $m_k(\boldsymbol{\theta}) = E(y_k|d_k, \boldsymbol{\theta})$ and $K$ is the number of observations. The expression $Var_{\boldsymbol{\theta}|\mathbf{y}_{1:(k-1)}}(m_k(\boldsymbol{\theta}))$ gives the variance of the mean response at $d_k$, given measurements $\mathbf{y}_{1:(k-1)}$ at points $\mathbf{d}_{1:(k-1)}$. The utility at $d_k$ is evaluated using a weighted variance, where each simulated response is weighted based on the likelihood of previous simulated measurements, $p(\mathbf{y}_{1:(k-1)}|\mathbf{d}_{1:(k-1)}, \boldsymbol{\theta})$.

Solonen et al. (2012) advocate the use of this utility function to design for parameter estimation since it is easier to compute than information-based utility functions (equation (2)). Solonen et al.'s (2012) utility function assumes a constant variance. Ryan et al. (2014c) present a generalised version of this utility function which may be used when the error structure of a model has a non-constant variance.

## 4.4 Utilities for Several Design Objectives

Researchers often have several competing goals for an experiment, rather than one single goal, and so these competing design objectives can be incorporated into one or several utility functions. One approach to dealing with competing design objectives is to weight each design criterion and search for the design that optimises the

weighted average of these criteria. This is known as a compound or weighted design problem (e.g., Dette [1990]). Clyde and Chaloner (1996) discuss compound design criteria and present an equivalence theorem for Bayesian constrained design problems. DasGupta et al. (1992) gave examples of compromise designs in which one is interested in finding a design that is highly efficient for several design problems.

Borth (1975) extends the mutual information utility proposed by Box and Hill (1967) so that fully Bayesian designs could be obtained for the dual goals of model discrimination and parameter estimation. This utility is known as "Total entropy". This dual design problem has been investigated in a number of classical design papers through use of compound criteria such as $D|T$- and $T|D-$optimality and hybrid DT-optimality (e.g., Atkinson [2008], Tommasi [2009], Waterhouse et al. [2009]), but is largely unexplored in the Bayesian design literature.

Chaloner and Verdinelli (1995) discuss several Bayesian utility functions that may be used for the dual purpose of maximising the expected value of the response and the expected information gain, and utilities which may be used to design for parameter estimation and prediction.

McGree et al. (2012c) considered compound utility functions in the context of Bayesian adaptive designs for dose-finding studies for the dual design objectives of estimating the maximum tolerated dose and addressing the safety of the study subjects. A number of different estimation utilities were used, and the utility functions only allowed doses to be available for selection if the 95th percentile of the posterior predictive probability of toxicity was less than some pre-specified tolerance level. Drovandi et al. (2013) developed a hybrid utility function for an adaptive dose-finding study to obtain robust estimates of the target stimulus-response curve in the presence of model and parameter uncertainty.

A number of studies have had the dual objectives of designing for parameter estimation or prediction accuracy and to minimise study costs (or inconvenience to study subjects). Stroud et al. (2001) used utility functions which designed for the precise estimation of parameters of interest, as well as minimising inconvenience to study subjects by penalising samples that were collected after a certain time period. Palmer and Müller (1998) searched for the optimal sampling times for stem cell collections in cancer patients, to minimise the expected loss function over the posterior predictive distribution for a new patient. Their utility function also included a penalty for failing to collect a certain target number of stem cells and a cost penalty for each sampling time scheduled.

# 5 Static Design Search Algorithms

Static design problems assume that the same design will be used throughout the experimental process, regardless of the incoming information that may be collected from the experiment. Static designs are useful when data are collected in a batch, according to a fixed protocol. A number of different algorithms have been used to solve Bayesian static design problems and they will be discussed below.

## 5.1 MCMC Algorithms

A number of stochastic algorithms have been proposed in the literature to approximate the maximisation and integration problem of equation (1) for static design problems. These include: prior simulation (Müller [1999]); smoothing of Monte Carlo simulations (Müller [1999]); Markov chain Monte Carlo (MCMC) simulation in an augmented probability model (Müller [1999]); and sequential Monte Carlo methods (Kück et al. [2006]).

### 5.1.1 Monte Carlo Integration

In many situations, one can simulate values of $(\boldsymbol{\theta}_i, \mathbf{y}_i)$ (for $i = 1, ..., M$) from $p(\boldsymbol{\theta}, \mathbf{y}|\mathbf{d})$ and the utility function can be estimated using these values. The integral is approximated by using:

$$\hat{U}(\mathbf{d}) = \frac{1}{M} \sum_{i=1}^{M} U(\mathbf{d}, \boldsymbol{\theta}_i, \mathbf{y}_i). \tag{5}$$

The optimal design, $\mathbf{d}^* = \arg \max \hat{U}(\mathbf{d})$, can then be found by using a suitable maximisation method to search over the estimates, $\hat{U}(\mathbf{d})$ (see Müller [1999]). This approach has commonly been used in the literature (e.g., Wakefield [1994], Carlin et al. [1998], Palmer and Müller [1998]) and is useful when a discrete set of possible designs that are of low dimension are used.

Müller and Parmigiani (1995) use a similar approach to equation (5), in which stochastic optimisation is performed by fitting curves to the Monte Carlo samples. First, they simulate draws from $(\boldsymbol{\theta}, \mathbf{y})$ and evaluate the observed utilities. Then, a smooth curve is fitted through these simulated points, which serves as an estimate of the expected utility surface. The optimal design can then be found deterministically. Kuo et al. (1999) also used these curve fitting methods for solving design problems of low dimension.

Straightforward Monte Carlo integration over $(\boldsymbol{\theta}, \mathbf{y})$ for each design $\mathbf{d}$ may be computationally intensive for design problems involving a large number of design variables or model parameters, since a large value of $M$ is required to obtain an estimate of $U(\mathbf{d})$ with high accuracy.

### 5.1.2 MCMC Simulation in an Augmented Probability Model

Alternatively, Clyde *et al.* (1996), Bielza *et al.* (1999) and Müller (1999) solved the optimal design problem by treating the expected utility as an unnormalised marginal probability density function. This was achieved by placing a joint distribution on $(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})$ to form an augmented probability model $h(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})$, which is given by:

$$h(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y}) \propto U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})p(\boldsymbol{\theta}, \mathbf{y}|\mathbf{d}),$$

where it was assumed that $U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})$ satisfies the appropriate conditions for $h(\cdot)$ to be positive and integrable over $(\mathbf{D}, \boldsymbol{\Theta}, \mathbf{Y})$. The probability distribution $h(\cdot)$ is defined such that the marginal distribution in $\mathbf{d}$ is proportional

to the expected utility, i.e.,

$$h(\mathbf{d}) \propto \int \int U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})p(\boldsymbol{\theta}, \mathbf{y}|\mathbf{d})d\boldsymbol{\theta}d\mathbf{y}$$

$$= U(\mathbf{d}).$$

It is assumed that the design space $\mathbf{D}$ is bounded and that the utility $U(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})$ is non-negative and bounded. One can then use a Metropolis-Hastings (MH) MCMC scheme to simulate from $h(\cdot)$ and select random draws from the design space that are proportional to the utility that is attached to the design. The MH MCMC algorithm focuses on sampling designs in areas of high expected utility and discourages sampling in areas of low expected utility (see Müller [1999]). The sample of simulated $\mathbf{d}$ may be used to provide an estimate of $h(\mathbf{d})$ and the joint mode of $h(\mathbf{d})$, $\mathbf{d}^*$, corresponds to the optimal design.

We note that the joint mode of $h(\mathbf{d})$ needs to be found rather than the marginal modes for each element of $\mathbf{d}$ as the latter may be very different from the former. Cook et al. (2008) and Drovandi and Pettitt (2014) propose methods for searching for the multivariate mode of the multivariate normal kernel smoothing density estimates of the design variables. However, for design problems that involve a large number of design points ($dim(\mathbf{d}) \geq 4$), the problem of finding the multivariate mode is more difficult than finding marginal modes and one may need to use dimension reduction techniques, such as those that Ryan et al. (2014c) propose. However, dimension reduction techniques may not always be appropriate and further research is needed into the problem of finding the multivariate mode for a large number of design variables.

### 5.1.3 Simulated Annealing-type Approach

In many design problems the shape of the expected utility surface can be very flat around its mode and prohibitively large simulation sample sizes may be required to estimate the mode. To overcome this problem, one can use an approach that is similar to simulated annealing (see Van Laarhoven and Aarts [1987]) in which the expected utility surfaces are replaced by a more peaked surface. This does not change the solution of the optimal design problem. The target function $h(\mathbf{d})$ is replaced with $h_J(\mathbf{d})$ where $J$ is an integer, usually large (say 20 or higher). The joint augmented distribution to simulate from is:

$$h_J(\mathbf{d}, \boldsymbol{\theta}_1, ..., \boldsymbol{\theta}_J, \mathbf{y}_1, ..., \mathbf{y}_J) \propto \prod_{j=1}^{J} U(\mathbf{d}, \boldsymbol{\theta}_j, \mathbf{y}_j)p(\boldsymbol{\theta}_j, \mathbf{y}_j|\mathbf{d}). \qquad (6)$$

For each $\mathbf{d}$, one simulates $J$ experiments $(\boldsymbol{\theta}_j, \mathbf{y}_j), j = 1, ..., J$, independently from $p(\boldsymbol{\theta}, \mathbf{y}|\mathbf{d})$ and considers the product of the calculated utilities. The product of the calculated utilities (rather than the sum) is used to ensure that $h_J(\mathbf{d}) \propto U^J(\mathbf{d})$.

This approach has been very popular in the literature (e.g., Bielza et al. [1999], Müller [1999], Stroud et al.

[2001], Cook et al. [2008], Ryan et al. [2014c]) and uses similar ideas to simulated annealing (see Van Laarhoven and Aarts [1987]) where $T = 1/J$ may be interpreted as the 'annealing temperature'. As $T \to 0$, the original target function is replaced with a point mass at the mode (Müller [1999]). As $J$ increases, the utility surface will become more peaked and simulations will cluster more tightly around the mode. However, increasing $J$ obviously increases the number of required computations. An annealing schedule is not required, i.e., the same value of $J$ may be used for all simulations. However, this is not efficient for high dimensional problems (see Amzal et al. [2006]) and a 'cooling' schedule may be required where $J$ increases to $+\infty$. Müller et al. (2004) recommend that $J$ should be gradually increased as the algorithm progresses so that the search will not become trapped in a local mode for situations where several modes exist. In Müller et al.'s (2004) approach, the algorithm initially explores the entire design space, but as the $J$ value increases, the MCMC draws focus around one of the highest modes.

Whilst the algorithm presented by Müller (1999) has "theoretically appealing" properties, it has been found to have slow convergence in practice, particularly for situations where there are a large number of design variables for which this algorithm becomes inefficient (Stroud et al. [2001], Amzal et al. [2006]). Use of this algorithm has therefore mostly been restricted to up to four design variables (e.g., Bielza et al. [1999], Müller [1999], Stroud et al. [2001], Cook et al. [2008]) and further research is required for searching for solutions to high dimensional design problems.

## 5.2   SMC Algorithms

Sequential Monte Carlo (SMC) algorithms, also known as "particle filters", use a population of particles to approximate a distribution and move through a smooth sequence of connected target distributions using resampling and diversification of particles until the final target distribution is reached (see Chopin [2002], Del Moral et al. [2006]). SMC combined with Markov and MCMC kernels provides a powerful and efficient computational approach for approximating target distributions. SMC has only recently been applied to static design problems (see Amzal et al. [2006], Kück et al. [2006]).

SMC methods can be useful for sampling from target distributions that change over time. This also includes the target distribution $h_J(\mathbf{d}, \boldsymbol{\theta}_{1:J}, \mathbf{y}_{1:J})$ (Müller et al. [2004]) in which $J$ increases over time. For nonlinear and high dimensional design problems, Amzal et al. (2006) extended the approach of Müller (1999) and Müller et al. (2004) through the use of particle methods, which are similar to particle filters (e.g., Doucet et al. [2001], Chopin [2002]) and population Monte Carlo simulations (e.g., Cappé et al. [2004]). This involves the simulation of $N_p$ 'parallel' Markov chains from the target distribution $h_J(\cdot)$, which are known as an 'interacting particles system'. At each iteration of the algorithm, an approximate weighted sample is generated from $h_J(\mathbf{d}, \boldsymbol{\theta}_1, ..., \boldsymbol{\theta}_J, \mathbf{y}_1, ..., \mathbf{y}_J)$ via importance sampling (Geweke [1989]) and a selection procedure, such as 'sampling with replacement' (e.g.,

Chopin [2002]), is then used to duplicate those particles that occur near the modes of the target distribution, whilst eliminating those that fall further away. An independent Markov step (Chopin [2002]) could also be added to the algorithm to avoid degeneracy problems and enrich the sample. Amzal et al. (2006) suggest that the proposal distribution for the design points should have a fairly large variance to enable the detection of other modes.

Amzal et al. (2006) also propose a "resampling-Markov algorithm" in which the importance sampling step is only implemented at initialisation of the algorithm. It is assumed that at time $t-1$, a sample that approximates $h_{J(t-1)}$ is available. If $J(t) > J(t-1)$, then additional values for each of the particles $\{\boldsymbol{\theta}_{j,k}^{(t-1)}, \mathbf{y}_{j,k}^{(t-1)}\}_{k=1}^{N_p}$, $j = J(t-1)+1, ..., J(t)$, are drawn from the posterior for $t-1$ and weights are computed that are proportional to the product of the utilities of the newly sampled values. The resampling and Markov steps are then implemented.

Kück et al. (2006) generalise the approach of Müller et al. (2004) to non integer annealing steps and used SMC samplers (similar to Del Moral et al. [2006], Johansen et al. [2006]) to search for designs which maximise the KLD between the prior predictive and posterior predictive distributions. In their application the model parameters $\boldsymbol{\theta}$ could be analytically integrated out of the expected utility which simplified the problem. A sequence of target distributions is generated artificially by incrementally "powering up" some measure of the utility for each particle. They define the following sequence of artificial target distributions:

$$h_{J(t),\nu_t}(\mathbf{d}, \boldsymbol{\theta}_{1:J(t)}, \mathbf{y}_{1:J(t)}) \propto \left( \prod_{j=1}^{J(t)-1} U(\mathbf{d}, \boldsymbol{\theta}_j, \mathbf{y}_j)p(\mathbf{y}_j|\mathbf{d}, \boldsymbol{\theta}_j)p(\boldsymbol{\theta}_j) \right) \left\{ U(\mathbf{d}, \boldsymbol{\theta}_{J(t)}, \mathbf{y}_{J(t)})p(\mathbf{y}_{J(t)}|\mathbf{d}, \boldsymbol{\theta}_{J(t)})p(\boldsymbol{\theta}_{J(t)}) \right\}^{\nu_t}.$$

The inverse annealing temperature, $J(t)$, is assumed to have an integer value. Kück et al. (2006) assumed that $J(t)$ could increase by at most one per iteration. Kück et al. (2006) also used the variable $\nu_t \in [0, 1]$ to enable smoother non-integer increases of the inverse annealing temperature. If $\nu_t = 1$, then the target density given in equation (6) is obtained and the dimension of the target distribution increases by $J(t) = J(t-1) + 1$, and $\nu_t$ is set back to zero. In all other instances, $J(t) = J(t-1)$. The choice of how to increase $J(t)$ is important, since large increments could result in degeneracy of the particles, and small increments are computationally inefficient. McGree et al. (2012a) propose to choose the increment to maintain a specific level of efficiency (based on the ESS) in the sample.

At time $t-1$, the particle set $\{\mathbf{d}_k^{(t-1)}, W_k^{(t-1)}\}_{k=1}^{N_p}$ provides an approximation for $h_{J(t-1)}$. A re-weight step is then implemented in the SMC algorithm via importance sampling to update the weighted particle set to approximate $h_{J(t)}$. Particles with a higher utility are given more weight than those with a lower utility. As $J$ increases, the target distribution becomes more peaked around the mode. Resampling and mutation steps are also used to avoid denegeracy in the particle set.

Kück et al.'s (2006) approach was found to behave well when exploring multi-modal target distributions.

## 5.3 Other Stochastic Approximation Algorithms

Huan and Marzouk (2013) used simultaneous perturbation stochastic approximation (SPSA) (Spall [1998]) and Nelder-Mead nonlinear simplex (NMNS) (Nelder and Mead [1965]) algorithms to perform stochastic optimisation for nonlinear and computationally intensive models. SPSA is a stochastic approximation method that is similar in nature to a steepdest-descent method that uses a finite difference estimate of the gradient. However, SPSA only uses two random perturbations to estimate the gradient, regardless of the dimension of the problem. Whilst the finite differences stochastic approximation (FDSA) algorithm only perturbs in one direction at a time, the SPSA algorithm perturbs in all directions at once. In SPSA, the error in the estimation of the gradient is "averaged out" over a large number of iterations (Spall [1998]) and the algorithm has a similar convergence rate to FDSA. SPSA has a global convergence property that relies on the existence of a non-negligable noise level in the objective function and the finite-difference-like perturbations (Maryak and Chin [2004]). However, high noise levels can cause slow convergence or can cause the algorithm to become stuck in local optima. SPSA is suitable for large-scale population models.

The NMNS algorithm has commonly been used for deterministic optimisation of nonlinear functions. It is a well-studied numerical method that is useful for problems in which gradients may be unknown. The NMNS algorithm is useful when dealing with noisy objective functions since it only requires a relative ordering of the function values, rather than the magnitudes of the differences (as when estimating gradients). NMNS is less sensitive than SPSA to the noise level, but can converge to non-stationary points. Huan and Marzouk (2013) found that the NMNS algorithm performed better than SPSA overall, in terms of the asymptotic distribution of the design variables and how quickly convergence was achieved.

Huan and Marzouk (2012) used the Robbins-Munro (RM) (Robbins and Monro [1951]) stochastic approximation, and compared it to a sample average approximation combined with the Broyden-Fletcher-Goldfarb-Shanno method (SAA-BFGS) to solve the optimal design problem for partial differential equations. The RM algorithm is one of the oldest stochastic approximation methods. It uses an iterative update that is similar to steepest descent, but uses stochastic gradient information. Sampling average approximation (SAA) algorithms reduce a stochastic optimisation problem to a deterministic one. For instance, in the optimal experimental design framework, we may define the problem to be solved as:

$$\mathbf{d}^* = \arg\max_{\mathbf{d} \in \mathbf{D}}\{U(\mathbf{d})\} = E_W[\hat{U}(\mathbf{d}, W)],$$

where $\mathbf{d}$ is the design variable, $W$ is the "noise" random variable, and $\hat{U}(\mathbf{d}, W)$ is an unbiased estimate of the objective function, $U(\mathbf{d})$ (e.g., KLD between the prior and posterior distributions). SAA approximates this

optimisation problem using

$$\hat{\mathbf{d}}_s = \arg\max_{\mathbf{d} \in \mathbf{D}}\{\hat{U}_M(\mathbf{d}, w_s) \equiv \frac{1}{M}\sum_{i=1}^{M}\hat{U}(\mathbf{d}, w_i)\},$$

where $\hat{\mathbf{d}}_s$ and $\hat{U}_M(\mathbf{d}, w_s)$ are the optimal design and utility function values under a particular set of $M$ realisations of $W$, where $w_s \equiv \{w_i\}_{i=1}^{M}$. The same set of realisation of $W$ is used for different values of $\mathbf{d}$ throughout the optimisation process, which makes the maximisation problem deterministic. The Broyden-Fletcher-Goldfarb-Shanno (BFGS) method (Nocedal and Wright [2006]), which is a deterministic quasi-Newton method, was used to find $\hat{\mathbf{d}}_s$ as an approximation to $\mathbf{d}^*$.

Huan and Marzouk (2012) used infinitesimal perturbation analysis (Ho and Cao [1983]) to construct an unbiased estimator of the gradient of the KLD for use in the RM algorithm. A polynomial chaos approximation of the forward model was also used to speed up computation of the utility function and gradient evaluations. Huan and Marzouk (2012) found that, although SAA-BFGS generally required fewer iterations, each iteration had a longer run time than a step of RM. As the evaluation of the utility function becomes more expensive, RM may be the more suitable of the two methods. RM was also found to outperform SAA-BFGS in terms of the size of the mean square error (between the "true" optimal value of the KLD and the value of the KLD for the current iteration), for a given computational effort.

# 6 Sequential Design Search Algorithms

Decisions are often made in stages, with additional data being observed between the decisions. For example, in dose-finding trials, dose allocation decisions are often made after previous cohorts have been administered the treatment so that future cohorts may be given doses that are closer to the maximum tolerated dose. Whitehead and Brunier (1995) and Whitehead and Williamson (1998) implement a Bayesian $m$-step look-ahead procedure to find the optimal treatment dose to administer to the next $m$ patients in a dose-finding study. Sequential design problems are those that involve an alternating sequence of decisions and observations. The Bayesian paradigm is extremely useful for sequential design problems since the posterior can be used as the prior distribution for the next experiment.

## 6.1 Backwards Induction

Although many approaches to solving sequential design problems use a myopic approach, which involves looking ahead only to the next observation (e.g., Cavagnaro et al. [2010], Drovandi et al. [2014], McGree et al. [2012b]), in general, this is not optimal, and one should instead look ahead to all future observations in the experiment (Borth [1975]). To achieve this, the computationally intensive *backward induction* method should be used (see, for example, DeGroot [1970], Berger [1985], Bernardo and Smith [2000] for a description) which considers all

future observations. Backward induction is also known as stochastic dynamic programming (e.g., Ross [1983]).

Early work in this area was restricted to simple model settings, such as one-sided tests of a univariate parameter (Berry and Ho [1988]), and binary outcome settings (Lewis and Berry [1994]). These approaches typically used only two or three backwards steps (interim looks at the data). Carlin et al. (1998) extend these approaches by including a forward sampling algorithm that can be used to find the optimal stopping boundaries in clinical trials and eases the computational burdens associated with backward induction. However, Carlin et al. (1998) used a univariate normal likelihood, assumed that the standard deviations were known at each step, and considered a maximum of 4 backwards steps.

Brockwell and Kadane (2003) proposed a gridding method which approximates the expected loss function (utility function) at each decision time, and consists of a function of certain summary statistics (low dimensional) of the posterior distribution of the parameter of interest. Their approach is similar to that of Berry et al. (2000). Brockwell and Kadane (2003) use a one-step-ahead forward simulation procedure to evaluate the expected utilities and focus on problems related to parameter estimation. Müller et al. (2006) also use a similar approach to Brockwell and Kadane (2003) which involves forward simulation to approximate the utility functions and constrain the action space to circumvent the problem of an increasing number of possible trajectories in the backward induction steps. Rossell et al. (2007) extend the approaches of Carlin et al. (1998), Brockwell and Kadane (2003), and Müller et al. (2006), in which they compute a summary statistic when new data are observed and use decision boundaries that partition the sample space. Once the summary statistic falls in the stopping region, the experiment is terminated. Thus the sequential problem is reduced to the problem of finding optimal stopping boundaries, and the choice of these boundaries accounts for all future data. Rossell and Müller (2013) extend these ideas to high-dimensional data by assuming that the data are suitably pre-processed.

## 6.2 MCMC Algorithms

McGree et al. (2012c) used MCMC methods (MH algorithms) to sample from the posterior distribution to find adaptive designs for a dose-finding study. Bayesian compound utility functions were used to find the dose for the next subject for the dual purposes of estimating the maximum tolerated dose (MTD) and addressing safety issues of toxicity. To estimate the utility functions, importance sampling was used in which the posterior distribution of the parameters (using the observations up to the $i - 1$th subject) $p(\boldsymbol{\theta}|\mathbf{y}_{(1:i-1)})$ was used as the importance distribution, and the target distribution was $p(\boldsymbol{\theta}|\mathbf{y}_{(1:i)})$, where $\mathbf{y}_i$ is the new data point given by dose $D$. McGree et al.'s (2012c) algorithm involved a form of self-tuning in that the proposal distribution for the model parameters $\boldsymbol{\theta}$ was based on a bivariate normal distribution in which the mean and variance were obtained from a maximum likelihood fit to the current data. Each time a new dose was selected, the proposal distribution was updated.

### 6.2.1 SMC Algorithms

SMC provides a natural framework for sequential design problems and has been used for parameter estimation design problems (e.g., Drovandi et al. [2013]), and model discrimination design problems (see Cavagnaro et al. [2010], Drovandi et al. [2014]). Its design applications are diverse and include computer experiments (e.g., Loeppky et al. [2010]), astrophysics (e.g., Loredo [2004]), cognitive science (e.g., Cavagnaro et al. [2010]), neurophysiology experiments (e.g., Lewi et al. [2009]), clinical trials (e.g., Liu et al. [2009]) and bioassays (e.g.,Tian and Wang [2009]).

SMC algorithms have commonly been used to design for model discrimination for Bayesian sequential design problems (e.g., Cavagnaro et al. [2010], Drovandi et al. [2014], McGree et al. [2012b]). Designs that efficiently and precisely estimate model parameters usually perform poorly on model discrimination problems (e.g., Atkinson [2008]).

Cavagnaro et al. (2010) use a similar approach to Amzal et al. (2006) in which an SMC algorithm was implemented to design optimally for model discrimination in the context of memory retention models. A simulated annealing effect (Müller [1999]) was used in which the utility function was incrementally "powered up". Cavagnaro et al.'s (2010) SMC algorithm designs for experiments one-observation-at-a-time, using the posterior distribution that is based on all of the data that has been observed thus far. Whilst these myopic approaches are sub-optimal, they are necessary in many applications of Bayesian design of experiments due to computational complexity of the backwards induction algorithm (Section 6.1).

Drovandi et al. (2014) present an SMC algorithm to sequentially design experiments one-at-a-time in the presence of model uncertainty for discrete data. McGree et al. (2012b) extended this approach for continuous data. In these works, an SMC algorithm is run in parallel for each of the competing models and the results are combined to compute the utility function in the presence of model uncertainty. This algorithm avoids between-model or cross dimensional proposals. The SMC algorithm produces an approximation to the evidence (the marginal likelihood of the data given a particular model) as a by-product (Del Moral et al. [2006]), which is used to compute the posterior model probabilities and to estimate the utility function. This avoids the need to use computationally intensive numerical integration techniques, such as quadrature (e.g., Cavagnaro et al. [2010]) to obtain an estimate of the evidence. Once the posterior model probabilities are computed, model discrimination utility functions, that are derived from information theory, such as the entropy of model probabilities (Box and Hill [1967], Borth [1975]) can be evaluated. The design $d$ that is chosen is the one that maximises the mutual information between the model indicator, $m$, and the predicted observation (Cavagnaro et al. [2010]). Little problem specific tuning is required for this algorithm and it is much less computationally intensive than approaches that rely on MCMC for posterior simulation in sequential design contexts (e.g., McGree et al. [2012c], Section 6.2).

In both Drovandi et al. (2014) and McGree et al.'s (2012b) work, only a discrete design space was considered and no optimisation algorithm was implemented. To reduce the computational requirements, the utility was evaluated for all possible choices of design, and the design which maximised the utility was chosen. For high dimensional design problems or those with continuous support, optimisation routines such as the exchange algorithm (Meyer and Nachtsheim [1995]) or simulated annealing (Corana et al. [1987]) may be required. Alternatively, one could incorporate the simulation-based algorithms of Müller (1999) or Amzal et al. (2006) to search over the design space.

# 7   Applications

We will now highlight some of the key areas that Bayesian experimental design is being applied to. Please note that this is not a comprehensive review on the applications of Bayesian experimental design, but rather an overview of some of the key papers in the literature.

## 7.1   Clinical Trial Design

There is a wealth of literature on Bayesian designs for clinical trial studies, with many practical developments in both sequential and static frameworks. The requirement of a decision-theoretic approach for the design of clinical trials was recognised as early as Anscombe (1968). Clinical trial design typically involves making decisions prior to the commencement of the experiment in relation to the drug dosage (e.g., dose level, timing of doses, number of doses, number of subjects to assign to each dose level), and for pharmacokinetic (what the subjects' body does to the treatment) and pharmacodynamic (what the treatment does to the subjects' body) studies, sampling times (e.g., number of samples to take, timing of the samples, assignment of subjects to sampling schedules).

Spiegelhalter et al. (2004) and Berry (2006) provide a general overview on how Bayesian methods can be used for inference and experimental design for clinical trial design. Berry (1993), Spiegelhalter et al. (1994), Kadane (1996), and Stangl and Berry (1998) discuss a number of important issues present in the use of Bayesian methods for the design and analysis of clinical trials, such as: ethics, prior elicitation, randomisation, treatment allocation, utilities, and decision making. However, the use of fully Bayesian designs for clinical trial design remains mostly theoretical and their use in practice is still uncommon. This is most likely due to the difficulties associated with prior elicitation for complex models and selection of a utility function, as well as the computational difficulties of optimising the expected utility function.

Decisions are often made sequentially in clinical trials as information is gathered on the experimental process from previous study subjects and enables the investigators to modify the experiment according to the accumulated information. Modifications to sequential clinical trial designs include adaptively assigning study subjects to treatments that have a higher performance or that will give more information about the experimental aims;

adding or deleting treatment arms; termination of the trial; and incorporation of more study subjects if the experimental aims have not been satisfied. Bornkamp et al. (2011), Müller et al. (2006), Wathen and Thall (2008), and Dragalin et al. (2010) present Bayesian sequential designs for clinical trials.

Dose-finding studies are concerned with determining the effect of different doses of the treatment on the response of interest and are required to ensure that marketed drug doses are safe and efficacious. One of the earliest works on Bayesian adaptive design for dose-finding studies is that by Whitehead and Brunier (1995), in which they obtain priors through elicited data and select treatment doses based on the gain in statistical information about an estimate.

Bornkamp et al. (2011) determine adaptive designs in a dose-finding study so that the minimum effective dose (MED), i.e., the smallest dose that achieves a clinically beneficial response over the placebo response, can be precisely estimated (using the approaches described in Bornkamp et al. [2007]). The dose-response relationship of a treatment is often unknown prior to the study. To account for model uncertainty, Bornkamp et al. (2011) average the design criterion (the posterior variance of the MED), conditional on model $m$, with respect to the model probabilities (see also, Dette et al. [2008]). This produces designs that are robust to model uncertainty. Bornkamp et al. (2011) also use a Bayesian shrinkage approach to stabilise the parameter estimates during the sequential updates of the parameter estimates and model probability.

Müller et al. (2006) use a grid-based backwards induction approach to make decisions about adaptive dose allocation, optimal stopping of a trial and the optimal decision upon stopping to enable optimal learning about the dose-response curve. Wathen and Thall (2008) describe an approach to find a group sequential design that maintains a targeted false-positive rate and power, under a wide range of true event time distributions for right-censored data in a phase III clinical trial. At each interim analysis, Wathen and Thall's (2008) procedure adaptively chooses the most likely model (based on the posterior probability) for the hazard function, using Bayesian model selection, and then they apply the decision bounds that are optimal for the chosen model. Their focus is on two-sided tests in two-arm trials. Dragalin et al. (2010) conduct an extensive simulation study that compares five different adaptive dose-finding designs. These designs differed in the number of doses, the number of interim analyses, and the number of patients allocated to each design, and were derived under different experimental objectives.

Stroud et al. (2001) use the MH MCMC algorithm of Müller (1999) to determine the optimal blood sampling times for the next patient to precisely estimate pharmacokinetic parameters of interest (subject to a cost penalty) for the anticancer agent, paclitaxel. The priors were obtained by fitting nonlinear mixed effects models to existing data. Ryan et al. (2014b) present fully Bayesian static designs for a horse population pharmacokinetic study. The design problem was to determine the optimal urine sampling times, as well as the number of subjects and samples per subject to obtain precise posterior distributions of the population parameters (subject to a cost

constraint). These designs were also obtained using an adaption of the MH MCMC algorithm of Müller (1999).

Other recent examples of Bayesian clinical trial design include Christen et al. (2004), Dragalin et al. (2007), Miller et al. (2007), Ding et al. (2008), Drovandi et al. (2013), and McGree et al. (2012c).

## 7.2 Cognitive Science

Optimal experimental design methods are also commonly used in the field of cognitive science. For example, Kujala and Lukka (2006) and Lesmes et al. (2006) use Bayesian sequential designs to estimate psychometric functions using utility functions based on maximum entropy. Lewi et al. (2009) present a sequential design framework that searches for the optimal design for a neurophysiology experiment that maximises the mutual information between the prior and posterior distribution for a generalised linear model. To facilitate estimation of the mutual information, a Gaussian approximation to the posterior was used. Myung and Pitt (2009) search for optimal static designs for model discrimination for memory retention and categorisation examples, using the approach of Amzal et al. (2006). Cavagnaro et al. (2010) use a similar approach to Myung and Pitt (2009), but instead search for sequential designs for model discrimination for a memory retention example. Kim et al. (2013) extend the approach of Cavagnaro et al. (2010) to find sequential designs for a population study of visual perception. Zhang and Lee (2010) find sequential designs to discriminate amongst competing models for a two arm bandit problem for human choice behaviour.

## 7.3 Natural Sciences

Loredo (2004) found sequential Bayesian designs for astrophysics experiments to detect extrasolar planets, using the maximum expected Shannon information gain on the posterior parameter estimates for the utility function. Huan and Marzouk (2013) use polynomial chaos approximations and nested Monte Carlo integration (Ryan [2003]) to estimate the KLD between the prior and posterior distributions to find static designs which enable inference about parameters for chemical kinetic models for combustion. Solonen et al. (2012) derive optimal static designs for an exothermic example. Cook et al. (2008) used Bayesian simulation-based strategies similar to Müller (1999) to determine observation times for botanical epidemic experiments that were governed by nonlinear stochastic processes.

# 8  Directions for Future Research

We believe the future of Bayesian experimental design lies in: (1) developing and implementing fast methods for approximating the posterior distribution for use in Bayesian utility functions, and fast computation of the Bayesian utility functions, as these are the most computationally intensive components of Bayesian experimental

| Search Algorithm framework | Method for approx. posterior | Example(s) |
|---|---|---|
| **Static designs** | | |
| | | |
| MCMC | Laplace approximation | Ryan et al. [2014a] |
| MCMC | Importance sampling | Cook et al. [2008], Ryan et al. [2014a,c] |
| MCMC | ABC | Drovandi and Pettitt [2013], Hainy et al. [2013] |
| MCMC | MCMC | Clyde et al. [1996] |
| Monte Carlo | MCMC | Han and Chaloner [2004] |
| SMC | Importance sampling | Amzal et al. [2006] |
| SPSA and NMNS | Polynomial chaos approximations and nested Monte Carlo integration | Huan and Marzouk [2013] |
| RM stochastic approximation | Nested Monte Carlo integration | Huan and Marzouk [2012] |
| SAA-BFGS | Nested Monte Carlo integration | Huan and Marzouk [2012] |
| **Sequential designs** | | |
| | | |
| Discrete search | Laplace approximation | Lewi et al. [2009] |
| SMC | Numerical quadrature | Cavagnaro et al. [2010] |
| Discrete search | SMC / importance sampling | Drovandi et al. [2013] |
| MCMC | Importance sampling | Stroud et al. [2001], McGree et al. [2012c] |
| Monte Carlo | MCMC | Wakefield [1994], Palmer and Müller [1998] |

Table 1: Summary of methods used to approximate the posterior distributions for Bayesian utility function estimation and for optimisation over $(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})$.

design; and (2) finding solutions to complex Bayesian experimental design problems, such as problems in which the likelihood is intractable or computationally prohibitive to calculate, or problems with a large number of design points.

## 8.1 Fast Algorithms for Bayesian Experimental Design

In Table 1 we provide a summary of the methods which have previously been used to approximate the posteriors for Bayesian utility functions, along with the search algorithms in which they are embedded.

MCMC and importance sampling have been found to be computationally intensive to perform at each iteration of the optimisation algorithm that searches over the space $(\mathbf{d}, \boldsymbol{\theta}, \mathbf{y})$, due to the large number of samples that are required to ensure that the Bayesian utility is well estimated. In particular, importance sampling from the prior performs poorly when large amounts of data are involved due to a low ESS (Ryan et al. [2014a]). Adaptive importance sampling (e.g., Kinas [1996], Pennanen and Koivu [2006]) may provide a faster method for approximating the posterior distributions, but is yet to be explored for Bayesian experimental design.

Laplace approximations and numerical quadrature have been found to be fast alternatives for approximating the posterior distribution in Bayesian design, and can be used when large amounts of data are involved, but rely on the assumption that the posterior distribution follows a multivariate normal distribution and also suffer from the curse of dimensionality. INLA can also provide a fast method for approximating the posterior distribution, but

has not been used for Bayesian experimental design. VB methods are a fast method for facilitating approximate inference for intractable posterior distributions, but are yet to be used in a Bayesian experimental design context.

Drovandi and Pettitt (2013) and Hainy et al. (2013) have explored the use of ABC rejection (see Beaumont et al. [2002]) within an MCMC framework to approximate the posterior distributions for Bayesian utility functions for design problems in which the likelihood function is intractable. Further use of ABC methods for posterior distribution approximation should be explored in the experimental design context.

A few studies have investigated the use of SMC for approximating the necessary quantities for Bayesian utility functions (e.g., Drovandi et al. [2013]), but its use has been limited. Future studies should focus on extending previous approaches to allow for more complicated design problems. SMC with a Liu West filter (Liu and West [2001]) could offer a fast method for posterior approximation for Bayesian design problems.

Computational burden is a major obstacle in all Bayesian design problems for complex models and must be overcome so that designs can be obtained efficiently and in real time, and to broaden the applicability of Bayesian design methodology by making it more accessible to practitioners, scientists and industry. This may be achieved through algorithmic developments and the exploitation of current parallel computing technology (such as graphics processing units or GPUs). Indeed, new parallel architectures are becoming increasingly available to individual researchers, and will have a significant impact on Bayesian experimental design. In order to take advantage of this increased power, computational problems and approaches should be adapted from the current serial processing paradigm to one that optimises algorithms for parallel processing. To our knowledge, there is no published, peer reviewed research on the use of GPUs in the derivation of a Bayesian experimental design.

## 8.2   Finding Optimal Designs for Complex Models

The future of Bayesian experimental design also lies in solving complex or nonstandard problems, such as problems in which the likelihood is intractable or computationally prohibitive to evaluate, problems where the observed data likelihood cannot be evaluated analytically, or problems with a large number of design points. Whilst sophisticated inference techniques are available for Bayesian data analysis for complex data models, corresponding methodology for deriving Bayesian experimental designs is severely lacking, and it is important that the methods for inference are complemented with appropriate experimental design methodologies that enable more informative data to be collected in a more timely manner. Use of parallel computing technology may be required to ease the computational burden of finding optimal Bayesian experimental designs for complex models (such as mixed effects models).

Fully Bayesian experimental designs for nonlinear mixed effects models are largely unexplored. Most of the current work has focused on evaluating Bayesian utility functions for a fixed set of discrete designs (e.g., Han and Chaloner [2004], Palmer and Müller [1998]) and selecting the design that produces the highest utility value

(i.e., no search over a continuous design space is performed). Ryan et al. (2014b) extend this by searching over a continuous design space to determine (near) optimal sampling times for a horse population pharmacokinetic study. Kim et al. (2013) find optimal sequential designs for population studies. Further work on using SMC algorithms (Chopin [2002]) to search for optimal designs for mixed effects models in the presence of model uncertainty is currently being conducted, so that solutions to real-world design problems can be found. The main difficulty in finding solutions to experimental design problems in which the data is modelled by mixed effects models is that the observed data likelihood is unavailable in closed form for all but the simplest examples.

## 8.3 Finding Optimal Designs for a Large Number of Design Variables

Better search algorithms are also required to find static designs. Many of the search algorithms for obtaining optimal designs (e.g., Müller [1999], Amzal et al. [2006]) are restricted to a small number of design variables ($\leq 4$), as these algorithms are computationally prohibitive for a large number of design variables (e.g., Bielza et al. [1999], Müller [1999], Stroud et al. [2001], Cook et al. [2008]). MCMC algorithms are good at estimating the marginal distribution of random variables, but experimental design requires the joint distribution, and in particular the joint mode of the design variables, which is quite difficult to find and estimate.

Ryan et al. (2014c) propose the use of lower dimensional parameterisations to enable near optimal designs to be found for problems that require a large number of design points. The lower dimensional parameterisations consist of a few design variables, which are optimised, and are then input into various functions to generate multiple design points. This was found to have substantial computational savings, and it was much easier to obtain the multivariate mode for a few design variables than for a large number of design variables. However, designs found using this method are not optimal but *near* optimal, which is a compromise of the computational savings achieved. The approach is only useful for design variables (e.g., sampling times/locations) that require multiple measures to be taken at specific points that are separated from one another in the design space. This approach does not overcome the problem of having a large number of different types of design variables (e.g., temperatures, pressures), and further research needs to be conducted for solving this design problem.

## 9 Conclusion

Bayesian experimental design is a fast growing area of research with many exciting recent developments. The Bayesian approach to experimental design offers many advantages over frequentist approaches, the most notable of which is the ability to optimise design criteria that are functions of the posterior distribution and can easily be tailored to the experimenters' design objectives. Bayesian design criteria are optimised often with the assumption that Bayesian inference will be performed on the data that is obtained from the experimental design. Bayesian frameworks also provide a formal approach for incorporating parameter uncertainties and prior information into

the design process via prior distributions, and provide a unified approach for joining these quantities with the model and design criterion. Another advantage of using a Bayesian design criterion is that a single design point can be used, and the prior distribution is updated by the single observation in a sequential manner. The prior information is not "thrown away" in fully Bayesian experimental design, as it is in pseudo-Bayesian design.

Whilst several review papers on Bayesian experimental design have been written, there is a lack of recent Bayesian experimental design papers that reflect the computational advancements that have occurred in recent times. In this article we have reviewed the computational methods that have been used to approximate the posterior distribution for Bayesian utility functions, along with methods for calculating the Bayesian utility functions (once the posterior has been approximated) and the search algorithms that have been used for finding the optimal designs. We have also highlighted some numerical methods and stochastic algorithms that have previously been used to perform Bayesian inference, but have not been used in the design context, and may provide fast alternatives for finding Bayesian designs.

It is our opinion that the future of Bayesian experimental design lies in the development and implementation of rapid methods for approximating the Bayesian utility functions, since this is the most computationally intensive component of the Bayesian experimental design process. We also believe that the future of Bayesian experimental design lies in finding solutions to complex or nonstandard design problems, such as problems in which the likelihood is intractable or computationally prohibitive to evaluate, problems where the observed data likelihood cannot be evaluated analytically, or problems with a large number of design points or design variables. Solutions to these difficult problems can only be achieved through algorithmic developments and the exploitation of current parallel computing technology.

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# References

B. Amzal, F. Bois, E. Parent, and C. P. Robert. Bayesian-optimal design via interacting particle systems. *Journal of the American Statistical Association*, 101(474):773–785, 2006.

F. Anscombe. Sequential medical trials. *Journal of the American Statistical Association*, 58:365–383, 1968.

A. C. Atkinson. Dt-optimum designs for model discrimination and parameter estimation. *Journal of Statistical Planning and Inference*, 138:56–64, 2008.

A. C. Atkinson and A. N. Donev. *Optimum Experimental Designs*. Oxford University Press, New York, 1992.

M. A. Beaumont, W. Zhang, and D. J. Balding. Approximate Bayesian computation in population genetics. *Genetics*, 162(4):2025–2035, 2002.

T. Bengtsson, P. Bickel, and B. Li. Curse-of-dimensionality revisited: Collapse of the particle filter in very large scale systems. In *Probability and Statistics: Essays in Honor of David A. Freedman*. Institute of Mathematical Statistics, 2008.

J. O. Berger. *Statistical Decision Theory and Bayesian Analysis*. Springer-Verlag, New York, 2nd edition, 1985.

J. M. Bernardo. Expected information as expected utility. *Annals of Statistics*, 7(3):686–690, 1979.

J. M. Bernardo and A. F. M. Smith. *Bayesian Theory*. John Wiley & Sons, 2nd edition, 2000.

D. Berry. A case for Bayesianism in clinical trials (with discussion). *Statistics in Medicine*, 12:1377–1404, 1993.

D. Berry. Bayesian clinical trials. *Nature Reviews Drug Discovery*, 5:27–36, 2006.

D. Berry, P. Müller, A. Grieve, M. Smith, T. Parke, R. Blazek, N. Mitchard, and M. Krams. *Case Studies in Bayesian Statistics*, chapter Adaptive bayesian designs for dose-ranging drug trials. Springer, 2000.

D. A. Berry and C.-H. Ho. One-sided sequential stopping boundaries for clinical trials: A decision-theoretic approach. *Biometrics*, 44:219–227, 1988.

C. Bielza, P. Müller, and D. R. Insua. Decision analysis by augmented probability simulation. *Management Science*, 45(7):995–1007, 1999.

B. Bornkamp, F. Bretz, A. Dmitrienko, G. Enas, B. Gaydos, C. Hsu, F. König, M. Krams, Q. Liu, B. Neuenschwander, T. Parke, and J. Pinheiro. Innovative approaches for designing and analyzing adaptive dose-ranging trials (with discussion). *Journal of Biopharmaceutical Statistics*, 17:965–995, 2007.

B. Bornkamp, F. Bretz, H. Dette, and J. Pinheiro. Response-adaptive dose-finding under model uncertainty. *Annals of Applied Statistics*, 5(2B):1611–1631, 2011.

D. M. Borth. A total entropy criterion for the dual problem of model discrimination and parameter estimation. *Journal of the Royal Statistical Society: Series B (Methodological)*, 37:77–87, 1975.

G. E. P. Box and W. J. Hill. Discrimination among mechanistic models. *Technometrics*, 9:57–71, 1967.

A. E. Brockwell and J. B. Kadane. A gridding method for Bayesian sequential decision problems. *Journal of Computational and Graphical Statistics*, 12(3):566–584, 2003.

O. Cappé, A. Guillin, J.-M. Marin, and C. Robert. Population Monte Carlo. *Journal of Computational and Graphical Statistics*, 13:907–929, 2004.

B. Carlin, J. Kadane, and A. Gelfand. Approaches for optimal sequential decision analysis in clinical trials. *Biometrics*, 54(3):964–975, 1998.

D. R. Cavagnaro, J. I. Myung, M. A. Pitt, and J. V. Kujala. Adaptive design optimization: A mutual information-based approach to model discrimination in cognitive science. *Neural Computation*, 22(4):887–905, 2010.

K. Chaloner. Optimal bayesian experimental designs for linear models. *Annals of Statistics*, 12:283–300, 1984.

K. Chaloner. An approach to design for generalised linear models. In *Proceedings of the Workshop on Model-oriented data analysis, Wartburg. Lecture Notes in Economics and Mathematical Systems.*, Berlin, 1987. Springer.

K. Chaloner and I. Verdinelli. Bayesian experimental design: A review. *Statistical Science*, 10:273–304, 1995.

N. Chopin. A sequential particle filter method for static models. *Biometrika*, 89(3):539–552, 2002.

J. Christen, P. Müller, K. Wathen, and J. Wolf. Bayesian randomized clinical trials: A decision-theoretic sequential design. *Canadian Journal of Statistics*, 32(4):387–402, 2004.

M. Clyde and K. Chaloner. The equivalence of constrained and weighted designs in multiple objective design problems. *Journal of the American Statistical Association*, 91:1236–1244, 1996.

M. A. Clyde, P. Müller, and G. Parmigiani. Exploring expected utility surfaces by Markov Chains. Technical report, Duke University, 1996.

A. Cook, G. Gibson, and C. Gilligan. Optimal observation times in experimental epidemic processes. *Biometrics*, 64(3):860–868, 2008.

A. Corana, M. Marchesi, C. Martini, and S. Ridella. Minimizing multimodal functions of continuous variables with the simulated annealing algorithm. *ACM Transactions on Mathematical Software*, 13:262–280, 1987.

D. D'Argenio. Incorporating prior parameter uncertainty in the design of sampling schedules for pharmacokinetic parameter estimation experiments. *Mathematical Biosciences*, 99(1):105–118, 1990.

A. DasGupta and W. Studden. Robust bayes designs in normal linear models. *Annals of Statistics*, 19:1244–1256, 1991.

A. DasGupta, S. Mukhopadhyay, and W. Studden. Compromise designs in heteroscedastic linear models. *Journal of Statistical Planning and Inference*, 32:363–384, 1992.

M. H. DeGroot. *Optimal Statistical Decisions*. McGrawHill, New York, 1970.

P. Del Moral, A. Doucet, and A. Jasra. Sequential Monte Carlo samplers. *Journal of the Royal Statistical Society: Series B (Statistical Methodology)*, 68(3):411–436, 2006.

H. Dette. A generalization of D-and D1-optimal designs in polynomial regression. *The Annals of Statistics*, 18: 1784–1804, 1990.

H. Dette, F. Bretz, A. Pepelyshev, and J. Pinheiro. Optimal designs for dose finding studies. *Journal of the American Statistical Association*, 103:1225–1237, 2008.

P. Diggle and S. Lophaven. Bayesian geostatistical design. *Scandinavian Journal of Statistics*, 33(1):53–64, 2006.

M. Ding, G. L. Rosner, and P. Müller. Bayesian optimal design for phase II screening trials. *Biometrics*, 64: 886–894, 2008.

A. Doucet, N. de Freitas, and N. Gordon. *Sequential Monte Carlo Methods in Practice*. Springer-Verlag, New York, 2001.

V. Dragalin, F. Hsuan, and S. Padmanabhan. Adaptive designs for dose-finding studies based on sigmoid Emax model. *Journal of Biopharmaceutical Statistics*, 17:1051–1070, 2007.

V. Dragalin, B. Bornkamp, F. Bretz, F. Miller, S. Padmanabhan, N. Patel, I. Perevozskaya, J. Pinheiro, and J. Smith. A simulation study to compare new adaptive doseranging designs. *Statistics in Biopharmaceutical Research*, 2:487–512, 2010.

C. Drovandi, J. McGree, and A. Pettitt. A sequential Monte Carlo algorithm to incorporate model uncertainty in Bayesian sequential design. *Journal of Computational and Graphical Statistics*, 23(1):3–24, 2014.

C. C. Drovandi and A. N. Pettitt. Estimation of parameters for macroparasite population evolution using approximate Bayesian computation. *Biometrics*, 67(1):225–233, 2011.

C. C. Drovandi and A. N. Pettitt. Bayesian experimental design for models with intractable likelihoods. *Biometrics*, 69(4):937–948, 2013.

C. C. Drovandi, A. N. Pettitt, and M. J. Faddy. Approximate Bayesian computation using indirect inference. *Journal of the Royal Statistical Society: Series C (Applied Statistics)*, 60(3):503–524, 2011.

C. C. Drovandi, J. M. McGree, and A. N. Pettitt. Sequential Monte Carlo for Bayesian sequential design. *Computational Statistics and Data Analysis*, 57(1):320 – 335, 2013.

S. El-Krunz and W. Studden. Bayesian optimal designs for linear regression models. *Annals of Statistics*, 19: 2183–2208, 1991.

R. Etzione and J. B. Kadane. Optimal experimental design for another's analysis. *Journal of the American Statistical Association*, 88:1401–1411, 1993.

V. Federov and P. Hackl. *Model-oriented Design of Experiments*. Springer-Verlag, Berlin, 1997.

V. V. Fedorov. *Theory of Optimal Experiments*. Academic Press, New York, 1972.

J. Geweke. Bayesian inference in Econometric models using Monte Carlo integration. *Econometrica*, 57(6): 1317–1339, 1989.

M. Hainy, W. Müller, and H. Wagner. Likelihood-free simulation-based optimal design. Technical report, Johannes Kepler University, Linz, 2013.

C. Han and K. Chaloner. Bayesian experimental design for nonlinear mixed-effects models with application to HIV dynamics. *Biometrics*, 60:25–33, 2004.

W. Hill, W. Hunter, and D. Wichern. A joint design criterion for the dual problem of model discrimination and parameter estimation. *Technometrics*, 10(1):145–160, 1968.

W. J. Hill. A review of experimental design procedures for regression model discrimination. *Technometerics*, 20: 15–21, 1978.

Y. C. Ho and X. Cao. Perturbation analysis and optimization of queueing networks. *Journal of Optimization Theory and Applications*, 40:559–582, 1983.

X. Huan and Y. M. Marzouk. Gradient-based stochastic optimization methods in bayesian experimental design. Technical report, Massachusetts Institute of Technology, Cambridge, 2012.

X. Huan and Y. M. Marzouk. Simulation-based optimal Bayesian experimental design for nonlinear systems. *Journal of Computational Physics*, 232(1):288–317, 2013.

A. Johansen, A. Doucet, and M. Davy. Maximum likelihood parameter estimation for maximum likelihood models using sequential Monte Carlo methods. In *ICASSP*, 2006.

J. B. Kadane. *Bayesian methods and ethics in clinical trial design*. John Wiley & Sons, 1996.

J. Kiefer. Optimum experimental designs. *Journal of the Royal Statistical Society. Series B*, 21:272–304, 1959.

J. Kiefer. Optimum designs in regression problems. II. *Annals of Mathematical Statistics*, 32(2):298–325, 1961.

J. Kiefer. General equivalence theory for optimum designs (approximate theory). *Annals of Statistics*, 2(5): 849–1063, 1974.

J. Kiefer and J. Wolfowitz. Optimum designs on regression problems. *Annals of Mathematical Statistics*, 30: 271–94, 1959.

J. Kiefer and J. Wolfowitz. The equivalence of two extremum problems. *Canadian Journal of Mathematics*, 14: 363–366, 1960.

J. Kiefer and J. Wolfowitz. On a theorem of Hoel and Levine on extrapolation designs. *Annals of Mathematical Statistics*, 36(6):1627–1655, 1965.

P. Kinas. Bayesian fishery stock assessment and decision making using adaptive importance sampling. *Canadian Journal of Fisheries and Aquatic Sciences*, 53:414–423, 1996.

H. Kück, N. de Freitas, and A. Doucet. Smc samplers for Bayesian optimal nonlinear design. Technical report, University of British Columbia, 2006.

J. V. Kujala and T. J. Lukka. Bayesian adaptive estimation: The next dimension. *Journal of Mathematical Psychology*, 50(4):369–389, 2006.

S. Kullback and R. A. Leibler. On information and sufficiency. *The Annals of Mathematical Statistics*, 22(1): 79–86, 1951.

L. Kuo, R. Soyer, and F. Wang. *Bayesian Statistics VI*, chapter Optimal design for quantal bioassay via Monte Carlo methods, pages 795–802. Oxford University Press, New York, 1999.

L. Lesmes, S.-T. Jeon, Z.-L. Lu, and B. Dosher. Bayesian adaptive estimation of threshold versus contrast external noise functions: The quick TvC method. *Vision Research*, 46:3160–3176, 2006.

J. Lewi, R. Butera, and L. Paninski. Sequential optimal design of neurophysiology experiments. *Neural Computation*, 21:619–687, 2009.

R. Lewis and D. A. Berry. Group sequential clinical trials: A classical evaluation of Bayesian decision-theoretic designs. *Journal of the American Statistical Association*, 89:1528–1534, 1994.

J. Liepe, S. Filippi, M. Komorowski, and M. P. H. Stumpf. Maximising the information content of experiments in systems biology. *PLoS Computational Biology*, 9(1):e1002888. doi:10.1371/journal.pcbi.1002888, 2013.

D. Lindley. On a measure of the information provided by an experiment. *Annals of Mathematical Statistics*, 27: 986–1005, 1956.

D. Lindley. The choice of variables in multiple regression. *Journal of the Royal Statistical Society Series B*, 30: 31–53, 1968.

D. Lindley. *Bayesian Statistics - A Review.* SIAM, Philadelphia, 1972.

G. Liu, W. F. Rosenberger, and L. M. Haines. Sequential designs for ordinal phase I clinical trials. *Biometrical Journal*, 51(2):335–347, 2009.

J. Liu and M. West. *Sequential Monte Carlo Methods in Practice*, chapter 10: Combined Parameter and State Estimation in Simulation-Based Filtering. Springer Verlag New York, 2001.

J. Loeppky, L. Moore, and B. Williams. Batch sequential designs for computer experiments. *Journal of Statistical Planning and Inference*, 140:1452–1464, 2010.

Q. Long, M. Scavino, R. Tempone, and S. Wang. Fast estimation of expected information gains for Bayesian experimental designs based on Laplace approximations. *Computer Methods in Applied Mechanics and Engineering*, 259:24–39, 2013.

J. López-Fidalgo, C. Tommani, and P. Trandafir. An optimal experimental design criterion for discriminating between non-normal models. *Journal of the Royal Statistical Society:Series B (Statistical Methodology)*, 69: 231–242, 2007.

T. Loredo. Bayesian adaptive exploration. In *Bayesian Inference and Maximum Entropy Methods in Science and Engineering: 23rd International Workshop on Bayesian Inference and Maximum Entropy Methods in Science and Engineering*, pages 330–346, 2004.

J. Maryak and D. Chin. Global random optimization by simultaneous perturbation stochastic approximation. *Johns Hopkins APL Technical Digest*, 25(2):91–100, 2004.

T. A. Masoumi, S.and Duever and P. M. Reilly. Sequential Markov chain Monte Carlo (MCMC) model discrimination. *The Canadian Journal of Chemical Engineering*, 91(5):862–869, 2013.

J. McGree, C. C. Drovandi, and A. N. Pettitt. A sequential Monte Carlo approach to derive sampling times and windows for population pharmacokinetic studies. *Journal of Pharmacokinetics and Pharmacodynamics*, 39(5): 519–526, 2012a.

J. McGree, C. C. Drovandi, and A. N. Pettitt. A sequential Monte Carlo approach to the sequential design for discriminating between rival continuous data models. Technical report, Queensland University of Technology, 2012b.

J. McGree, C. C. Drovandi, H. Thompson, J. Eccleston, S. Duffull, K. Mengersen, A. N. Pettitt, and T. Goggin. Adaptive Bayesian compound designs for dose finding studies. *Journal of Statistical Planning and Inference*, 142(6):1480–1492, 2012c.

R. Meyer and C. Nachtsheim. The coordinate-exchange algorithm for constructing exact optimal experimental designs. *Technometerics*, 37(1):60–69, 1995.

F. Miller, O. Guilbaud, and H. Dette. Optimal designs for estimating the interesting part of a dose-effect curve. *Journal of Biopharmaceutical Statistics*, 17(6):1097–1115, 2007.

P. Müller. Simulation-based optimal design. *Bayesian Statistics*, 6:459–474, 1999.

P. Müller and G. Parmigiani. Optimal design via curve fitting of monte carlo experiments. *Journal of the American Sta*, 90(432):1322–1330, 1995.

P. Müller, B. Sansó, and M. De Iorio. Optimal Bayesian design by inhomogeneous Markov chain simulation. *Journal of the American Statistical Association*, 99(467):788–798, 2004.

P. Müller, D. A. Berry, A. P. Grieve, and M. Krams. A Bayesian decision-theoretic dose-finding trial. *Decision Analysis*, 3(4):197–207, Dec. 2006.

J. A. Nelder and R. Mead. A simplex method for function minimization. *The Computer Journal*, 7(4):308–313, 1965.

S. H. Ng and S. E. Chick. Design of follow-up experiments for improving model discrimination and parameter estimation. *Naval Research Logistics*, 51:1129–1148, 2004.

J. Nocedal and S. J. Wright. *Numerical Optimization*. Springer, 2nd edition, 2006.

J. Ormerod and M. Wand. Explaining variational approximations. *American Statistical Association*, 64(2): 140–153, 2010.

J. Palmer and P. Müller. Bayesian optimal design in population models for haematologic data. *Statistics in Medicine*, 17:1613–1622, 1998.

L. Paninski. Asymptotic theory of information-theoretic experimental design. *Neural Computation*, 17:1480–1507, 2005.

T. Pennanen and M. Koivu. An adaptive importance sampling technique. In H. Niederreiter and D. Talay, editors, *Monte Carlo and Quasi-Monte Carlo Methods 2004*, pages 443–455. Springer Berlin, Heidelberg, 2006.

J. Pilz. *Bayesian estimation and experimental design in linear regregression models (2nd ed)*. Wiley, New York, 1991.

L. Pronzato and E. Walter. Robust experiment design via stochastic approximation. *Mathematical Biosciences*, 75(1):103–120, 1985.

F. Pukelsheim. *Optimal Design of Experiments*. Wiley, New York, 1993.

F. Pukelsheim and B. Torsney. Optimal weights for experimental designs on linearly independent support points. *The Annals of Statistics*, 19(3):1614–1625, 1991.

D. Rios Insua and F. Ruggeri. *Robust Bayesian Analysis*. Springer Verlag, New York, 2000.

H. Robbins and S. Monro. A stochastic approximation method. *The Annals of Mathematical Statistics*, 22(3): 400–407, 1951.

S. Ross. *Introduction to stochastic dynamic programming*. Academic Press, 1983.

D. Rossell and P. Müller. Sequential stopping for high-throughput experiments. *Biostatistics*, 14(1):75–86, 2013.

D. Rossell, P. Müller, and G. L. Rosner. Screening designs for drug development. *Biostatistics*, 8(3):595–608, 2007.

P. Roth. *Design of Experiments for Discrimination Among Rival Models*. PhD thesis, Princeton University, New Jersey, USA., 1965.

H. Rue, S. Martino, and N. Chopin. Approximate Bayesian inference for latent Gaussian models using integrated nested Laplace approximations (with discussion). *Journal of the Royal Statistical Society, Series B*, 71(2): 319–392., 2009.

E. Ryan, C. C. Drovandi, and A. N. Pettitt. Fully Bayesian experimental design for pharmacokinetic studies. Technical report, Queensland University of Techonology, 2014a.

E. Ryan, C. C. Drovandi, and A. N. Pettitt. Simulation-based fully bayesian experimental design for mixed effects models. Technical report, Queensland University of Technology, 2014b.

E. Ryan, C. C. Drovandi, M. Thompson, and A. N. Pettitt. Towards Bayesian experimental design for nonlinear models that require a large number of sampling times. *Computational Statistics and Data Analysis*, 70:45–60, 2014c.

K. Ryan. Estimating expected information gains for experimental designs with application to the random fatigue-limit model. *Journal of Computational and Graphical Statistics*, 12:585–603, 2003.

C. Shannon. A mathematical theory of communication. *Bell System Technical Journal*, 27:379–423, 623–656, 1948.

S. D. Silvey. *Optimal design.* Chapman and Hall, London, 1980.

S. A. Sisson and Y. Fan. *MCMC handbook*, chapter Likelihood-free Markov chain Monte Carlo, pages 313–335. Chapman & Hall., 2011.

A. Solonen, H. Haario, and M. Laine. Simulation-based optimal design using a response variance criterion. *Journal of Computational and Graphical Statistics*, 21(1):234–252, 2012.

J. C. Spall. An overview of the simultaneous perturbation method for efficient optimization. *Johns Hopkins APL Technical Digest*, 19(4):482–492, 1998.

D. Spiegelhalter, L. Freedman, and M. Parmar. Bayesian approaches to randomize trials. In D. Berry and D. Stangl, editors, *Bayesian Biostatistics*, pages 67–108. Dekker, New York, 1996.

D. J. Spiegelhalter. Incorporating Bayesian ideas into health-care evaluation. *Statistical Science*, 19(1):156–174, 2004.

D. J. Spiegelhalter, L. S. Freedman, and M. K. B. Parmar. Bayesian approaches to randomized trials. *Journal of the Royal Statistical Society. Series A (Statistics in Society)*, 157(3):357–416, 1994.

D. Stangl and D. Berry. Bayesian statistics in medicine: Where are we and where should we be going? *The Indian Journal of Statistics*, 60:176–195, 1998.

J. Stroud, P. Müller, and G. Rosner. Optimal sampling times in population pharmacokinetic studies. *Journal of the Royal Statistical Society: Series C (Applied Statistics)*, 50(3):345–359, 2001.

Y. Tian and D. Wang. Sequential bayesian design for estimation of EDp. In *The 2nd International Conference on Biomedical Engineering and Informatics, 2009. BMEI'09*, 2009.

B. Toman and J. Gastwirth. Efficienct robust experimental design and estimation using a data-based prior. *Statistical Sinica*, 4:603–615, 1994.

C. Tommasi. Optimal designs for both model discrimination and parameter estimation. *Journal of Statistical Planning and Inference*, 139:4123–4132, 2009.

C. Tsai and K. Chaloner. *Case studies in Bayesian Statistics 5*, chapter Using Prior opinions to examine sample size in a clinical trial: two examples, pages 409–423. Springer-Verlag, New York, 2002.

P. Van Laarhoven and E. Aarts. *Simulated Annealing: Theory and Applications.* Reider, Dordrecht, 1987.

J. Vanlier, C. Tiemann, P. Hilbers, and N. van Riel. Optimal experimental design for model selection in biochemical networks. *BMC Systems Biology*, 8:20, 2014.

I. Verdinelli. Bayesian design for the normal linear model with unknown error variance. *Biometrika*, 87:222–227, 2000.

J. Wakefield. An expected loss approach to the design of dosage regimens via sampling-based methods. *Journal of the Royal Statistical Society. Series D (The Statistician)*, 43(1):13–29, 1994.

T. H. Waterhouse, J. A. Eccleston, and S. B. Duffull. Optimal design criteria for discrimination and estimation in nonlinear models. *Journal of Biopharmaceutical Statistics*, 19:386–402, 2009.

J. Wathen and P. Thall. Bayesian adaptive model selection for optimizing group sequential clinical trials. *Statistics in Medicine*, 27:5586–5604, 2008.

J. Whitehead and H. Brunier. Bayesian decision procedures for dose determining experiments. *Statistics in Medicine*, 14:885–893, 1995.

J. Whitehead and D. Williamson. Bayesian decision procedures based on logisic regression models for dose-finding studies. *Journal of Biopharmaceutical Statistics*, 8:445–467, 1998.

L. Wolfson, J. Kadane, and M. Small. Expected utility as a policy making tool: an environmental health example. In D. Berry and D. Stangl, editors, *Bayesian Biostatistics*, pages 261–277. Dekker, New York, 1996.

S. Zhang and M. Lee. Optimal experimental design for a class of bandit problems. *Journal of Mathematical Psychology*, 54(6):499–508, 2010.

J. Zidek, W. Sun, and N. Le. Designing and integrating composite networks for monitoring multivariate Gaussian pollution fields. *Applied Statistics*, 49:63–79, 2000.