A Bayesian approach to targeted experiment design

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ABSTRACT

Motivation: Systems biology employs mathematical modelling to further our understanding of biochemical pathways. Since the amount of experimental data on which the models are parameterized is often limited, these models exhibit large uncertainty in both parameters and predictions. Statistical methods can be used to select experiments that will reduce such uncertainty in an optimal manner. However, existing methods for optimal experiment design (OED) rely on assumptions that are inappropriate when data are scarce or considering model complexity.

Results: We have developed a novel method to perform OED for models that cope with large parameter uncertainty. We employ a Bayesian approach involving importance sampling of the posterior predictive distribution to predict the efficacy of a new measurement at reducing the uncertainty of a selected prediction. We demonstrate the method by applying it to a case where we show that specific combinations of experiments result in more precise predictions.

Availability and implementation: Source code is available at: http://bmi.bmt.tue.nl/sysbio/software/pua.html

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1 INTRODUCTION

Computational models can be used to predict (un)measured behaviour or system responses and formalize hypotheses in a testable manner. To be able to make predictions parameters are required. Despite the development of new quantitative experimental techniques, data are often relatively scarce. Consequently the modeller is faced with a situation where large regions of parameter space can describe the measured data to an acceptable degree (Höfer et al. 2008). This is not a problem when the predictions required for testing the hypothesis (which we shall refer to as predictions of interest) are well constrained (Kreutz et al. 2006). However, when data are scarce or considering model complexity the model complexity or the model is strongly non-linear, such methods are not appropriate (Kreutz et al. 2006). This makes investigating the role of parameter uncertainty in OED a relevant topic to explore.

We propose a method for experimental design that overcomes these issues by adopting a probabilistic approach which incorporates prediction uncertainty. Our method enables the modeller to target experimental efforts in order to selectively reduce the uncertainty of predictions of interest. Using our approach, multiple experiments can be designed simultaneously revealing potential benefits that might arise from specific combinations of experiments.

We focus on biochemical networks that can be modelled using a system of ordinary differential equations. These models comprise of equations of the form:

\[ \dot{x}(t) = f(x(t), u(t), p) \]  

\[ \tilde{y}(t) = g(x(t), \tilde{q}) + \xi(t) \]  

\[ \tilde{q}(0) = \tilde{q} \]  

In order to perform inference and experiment design an error model is required. For ease of notation we shall demonstrate our method using a Gaussian error model. If we consider \( \tilde{y}(t) \) state variables \( \tilde{q}(t) \) (constant in time), inputs \( \tilde{u}(t) \) and state variables \( \tilde{x}(t) \). Given a set of parameters, inputs and initial conditions \( \tilde{q}(0) \) these equations can subsequently be simulated. Measurements \( \tilde{y}(t) \) are performed on a subset and/or a combination of the total number of states in the model.

Measurements are hampered by measurement noise \( \xi \) while many techniques used in biology (e.g. western blotting) necessitate the use of scaling and offset parameters \( \tilde{q} \) (Kreutz et al. 2006). We define \( \tilde{q} \) as \( \tilde{q} = [\tilde{p}, \tilde{q}_0] \), which lists all the required variables to simulate the model.

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Equation (3) multiplied with the prior to a unit area. Here \( P(y|\theta) \) corresponds to the probability of observing dataset \( y \) given the true parameters \( \theta \). Both computational as well as methodological advances have made Markov Chain Monte Carlo (MCMC) an attractive option for obtaining samples from such a distribution. 

Given a sample of the posterior parameter distribution, predictions can be made by simulating the model for each of the parameter sets. The distribution of such predictions shall be referred to as the posterior predictive distribution (PPD) and reflects their uncertainty. Since all of these predictions are linked via the parameter distributions, the relations between the different predictions can be exploited for experimental design. By considering the effects of a new measurement on the PPD, it is possible to predict how useful an experimental design would be. This approach consists of a number of steps. First, we briefly mention how to compute the PPD. Subsequently we detail how to compute the efficacy of the new measurement. In a third step this measure is used for experimental design. We conclude by demonstrating the method by applying it to a case study.

2 APPROACH

In order to overcome the limitations of existing OED methods, a sampling-based approach for experimental design is proposed. This approach consists of four consecutive steps which we shall outline below.

2.1 Step 1. Computation of the posterior parameter distribution

The first step in the analysis is computing the posterior parameter distribution of the model based on the available data:

\[
P(\theta|y) = \frac{P(y|\theta)P(\theta)}{\int P(y|\theta)P(\theta)d\theta}
\]

Here probability reflects a degree of belief and prior knowledge regarding the parameters is included in the form of priors, and \( P(y|\theta) \) denotes the conditional probability of the data given the model parameters. The unnormalized form of this function is often referred to as the likelihood function. Furthermore, \( P(\theta) \) refers to the prior distribution of the parameters.

In order to sample from the posterior distribution we employ a MCMC method known as the Metropolis–Hastings algorithm. This algorithm performs a random walk through parameter space where each subsequent step is based on a proposal distribution (centred on the current step) and an acceptance criterion based on the proposal and probability densities at the sampled points. In brief, after an initial burn-in period (which is discarded), MCMC methods generate samples from probability distributions whose posterior densities are known up to a normalizing factor. The Metropolis–Hastings algorithm proceeds as follows:

1. Generate a sample \( \theta_{n+1} \) by sampling from a proposal distribution based on the current state \( \theta_n \).
2. Compute the likelihood of the data \( L(y|\theta_{n+1}) \) and calculate \( P(\theta_{n+1}|y) = L(y|\theta_{n+1})P(\theta_{n+1}) / P(\theta_n) \), where \( P(\theta_{n+1}) \) refers to the prior density of \( \theta_{n+1} \).
3. Draw a random number \( y \) from a uniform distribution between 0 and 1 and accept the new step if \( y < \min\left\{1, \frac{L(\theta_{n+1}|y)P(\theta_{n+1})}{L(\theta_n|y)P(\theta_n)}\right\} \).

Here \( (\theta_n → \theta_{n+1}) \) refers to the proposal density from current parameter set \( \theta_n \) to \( \theta_{n+1} \). The ratio of \( \theta_n \) ensures detailed balance, a sufficient condition for the Markov chain to converge to the equilibrium distribution. This can be found in the Supplementary Materials.

It corrects for sampling biases resulting from non-symmetric proposal distributions and is defined as the ratio between the proposal densities associated with going from \( n \) to \( n+1 \) and \( n+1 \) to \( n \). We employ an adaptive Gaussian proposal distribution whose covariance matrix is based on a quadratic approximation to the posterior probability at the current sample point. Further details regarding the implementation can be found in the Supplementary Materials.

2.2 Step 2. Determine PPDs for all candidate experiments

A PPD is a distribution of predictions conditioned on the available data as shown in Equation (3). A PPD is obtained by simulating the model (including the addition of measurement noise) for a sample of parameter sets from the posterior parameter distribution. We simulate a PPD for each candidate experiment. These PPDs link the parameters to the predictions and via the parameters also link predictions (across different experiments) to each other. The model and data constrain the dynamics of the system and hereby implicitly impose non-trivial relations between the different predictions. Therefore, the observables of candidate experiments are related to our prediction of interest. The next step is to exploit the relations within these distributions for experimental design.

\[
P(y|\theta_{n+1}) = \frac{P(y|\theta_{n+1})P(\theta_{n+1})}{\int P(y|\theta)P(\theta)d\theta}
\]

2.3 Step 3. Predict EVR based on PPDs and measurement accuracies

To be able to perform experiment design a measure of expected measurement efficacy is required. For this purpose, we introduce the expected variance reduction (EVR). Consider an independent new measurement of a specific parameter \( G \) which reflects a certain degree of uncertainty associated with the new experiment. If this new experiment were to be performed and a value is obtained, then the subsequent step would be to incorporate the new data point (and its associated error model) in the likelihood function and re-perform the MCMC. This new data would subsequently constrain the posterior parameter distribution, hence also affecting the prediction of interest (which cannot be measured directly). This process is illustrated in Figure 4.

The outcome of the experiment is known only after the experiment has been performed. Therefore the measured value in the error model is not known a priori. However, the PPD provides us with a predicted distribution of this value (shown in grey in Fig 1) which reflects the uncertainty associated with this value. Samples from the PPD can subsequently be substituted as ‘true’ values in the error model. By repeating this process for every R-th point of our MCMC chain and averaging the result, we weight by the probability distribution of this predicted value. Considering that a single MCMC is often already computationally demanding, such a nested MCMC is likely not tractable. Here we propose an alternative approach. Consider the normalized densities \( P(y|\theta), P(y_2|\theta) \) and \( P(y_1|\theta) \), respectively, corresponding to the density model of the data used to determine the initial posterior distribution, the density model for the new data point, and the parameter prior. Assuming that the new data point is independent of the existing data points we can state that \( P(y|\theta, y_1, y_2) \propto P(y|\theta)P(y_1|\theta)P(y_2|\theta) \) in order to obtain the following equation for the new normalized posterior. In this equation, \( Z_1 \) and \( Z_2 \) denote the normalization constants of the old and new posterior, respectively.

\[
P(\theta|y, y_1, y_2) = \frac{P(y_1|\theta)P(y_2|\theta)}{P(y|\theta)P(y_1|\theta)P(y_2|\theta)} \frac{P(y|\theta)P(y_1|\theta)P(y_2|\theta)}{P(y|\theta)P(y_1|\theta)P(y_2|\theta)}
\]

\[
Z_1 = \frac{P(y_1|\theta)P(y_2|\theta)}{P(y|\theta)P(y_1|\theta)P(y_2|\theta)}
\]

\[
Z_2 = \frac{P(y_1|\theta)P(y_2|\theta)}{P(y|\theta)P(y_1|\theta)P(y_2|\theta)}
\]

This relation between the two posteriors can be exploited in order to compute expected values by re-weighting samples from the old posterior.
appropriately. Rather than running a new MCMC for every sample, we can use self normalized importance sampling on the predictions of the output in order to compute expected values. This is shown in Equation (11), where \( E(z|y) \) is the expected value of \( z \) given \( y \).

\[
E(z|y) = \int P(z|y) \, d\theta
= \int P(z|\theta) P(\theta|y) \, d\theta
= \sum_{i=1}^Q \frac{P(z_i|\theta_i)}{\sum_{j=1}^Q P(z_j|\theta_j)} \theta_i
\]

As mentioned before, the value of \( y \) is not known a priori. Therefore, we subsequently compute such an expected value for each parameter set in the PPD with \( y \) set to the predicted output value from the original PPD. Hence this provides a distribution of expected values considering possible outcomes of the experiment. The mean of these expected values then provides us with a prediction of the quantity of interest. The entire approach can then be succinctly summarized in Equation (12). Here the expected value of \( z \) is computed, \( G \) corresponds to the error model and \( \theta \) refers to the \( q \)th parameter vector of the chain. Assuming a Gaussian error model with SD \( \sigma \) for a new measurement or the output \( y \), probability model \( G \) is given by Equation (13). Note that both input as well as output can be any quantity of interest (prediction or parameter) indicating the flexibility of the approach.

\[
E(z|y) = \frac{1}{T \times Q} \sum_{i=1}^T \sum_{j=1}^Q G(t(\theta_t), \tilde{y}, \tilde{y}_j) \, \theta_t
\]

Since the variance of a variable of interest can be computed by Equation (14), we can use the aforementioned method to estimate this quantity. The variance reduction can then be computed as shown in Equation (15), where \( \sigma_{\text{new}}^2 \) corresponds to the posterior variance without the new measurement and \( \sigma_{\text{old}}^2 \) corresponds to the expected posterior variance with the new measurements taken into account. In other words, one obtains the mean variance reduction considering the prediction uncertainty. The variance reduction computed by this sampling method is referred to as the variance reduction (SVR).

\[
\text{Var}(z) = E(z^2) - [E(z)]^2
\]

\[
\text{Var}R = 1 - E\left( \frac{\sigma_{\text{old}}^2}{\sigma_{\text{new}}^2} \right)
\]

### 2.4 Step 4. Determine measurement points for optimal variance reduction

The probability density model can be obtained by multiplying the error models for each candidate measurement. Subsequently, the space of all candidate measurements is sampled using Monte Carlo sampling. The efficacy of a specific combination of measurements is evaluated by computing the variance reduction, which is defined as Equation (16). During this sampling stage, additional constraints which arise because of practical considerations can be imposed on the experimental design (simply by rejecting such samples). An example of this could be the inability to measure certain states simultaneously. The optimal experiment is then obtained by determining the combination of measurements that yields the maximal predicted variance reduction.

### 3 COMPUTATIONAL METHODS

All of the discussed algorithms were implemented in Matlab (Natick, MA, USA). Numerical integration of the differential equations was performed with compiled MEX files using numerical integrators from the SUNDIALS CVode package (Lawrence Livermore National Laboratory, Livermore, CA, USA). Absolute and relative tolerances were set to \( 10^{-10} \) and \( 10^{-13} \), respectively. The Gaussian proposal distribution for the MCMC was based on an approximation to the Hessian computed using a Jacobian obtained.
concentration of phosphorylated STAT in the cytoplasm (\(x_2+2x_3\)) were measured. Note that the step from \(x_4\) back to \(x_1\) is associated with a delay. Using finite differencing (\(H=J^TJ\)). All available priors were subsequently included in the Hessian approximation. After convergence, the chain was thinned to 10,000 samples. The SVR was computed in parallel using OpenCL on the GPU using a compiled MEX file.

4 RESULTS

To illustrate our method, we apply it to a model of the JAK-STAT signalling pathway [Raue et al. 2009; Toni et al. 2009]. The model is based on a number of hypothesized steps (see Figure 2). The first reaction describes the activation of the erythropoietin receptor which subsequently phosphorylates cytoplasmic STAT (\(x_1\)). Phosphorylated STAT (\(x_2\)) dimerises (\(x_3\)) and is imported into the nucleus (\(x_4\)). Here dissociation and dephosphorylation occurs which are associated with a time delay. Similar to the implementation given in the original paper, the driving input function was approximated by a spline interpolant, while the delay was approximated using a linear chain approximation \((x_5, \ldots, x_{13})\).

\[
\begin{align*}
x_1 & = \frac{V_{	ext{cyto}}}{V_{	ext{nucl}}} (p_4 x_{13}) - p_1 x_{12} x_4 \\
x_2 & = p_1 x_{12} x_4 - 2p_2 x_2 x_3 \\
x_3 & = p_2 x_2 - p_3 x_4 \\
x_4 & = \frac{V_{	ext{nucl}}}{V_{	ext{cyto}}} (p_3 x_3) - p_4 x_4 \\
x_5 & = p_4 x_4 - p_4 x_5 \\
x_6 & = p_4 x_5 - p_4 x_6 \\
x_7 & = p_4 x_6 - p_4 x_7 \\
x_8 & = p_4 x_7 - p_4 x_8 \\
x_9 & = p_4 x_8 - p_4 x_9 \\
x_{10} & = p_4 x_9 - p_4 x_{10} \\
x_{11} & = p_4 x_{10} - p_4 x_{11} \\
x_{12} & = p_4 x_{11} - p_4 x_{12} \\
x_{13} & = p_4 x_{12} - p_4 x_{13} \\
x_4 & = p_4 x_{13} - p_4 x_4 \\
\end{align*}
\]

In order to infer the posterior distribution data from the paper by Swamye et al. [2003; http://webber.physik.uni-freiburg.de/~jeti/\(\text{PNAS_Swamye_Data/dataset 1}\)] was used. Measured quantities were the total concentration of STAT (\(x_1 + x_2 + 2x_3\)) and the total concentration of phosphorylated STAT in the cytoplasm (\(x_2 + 2x_3\)), both reported in arbitrary units (which necessitates two scaling parameters). The initial cytoplasmic concentration of STAT is unknown while all other forms of STAT are assumed zero at the start of the simulation. Given the data, not all parameters are identifiable [Raue et al. 2009]. We used uniform priors in logspace for the kinetic parameters and a Gaussian (\(\mu = 200 \text{nM}, \sigma = 20 \text{nM}\)) for the initial condition. Parameter two was bounded between ranges, since this parameter was non-identifiable from the data [Raue et al. 2009]. We simulated two chains starting at different initial values up to one million parameter sets and assessed convergence by visually inspecting differences between batches of samples. The uncertainty in model parameters propagates as an uncertainty in the predicted responses of the state variables. PPDs were simulated for all states as well as the summations of states already measured. To simulate the PPDs the chain of parameter sets was thinned to 10,000 samples using equidistant thinning. Since the error model in this case is additive Gaussian noise, there is no need to explicitly simulate measurement noise. This can be taken into account by multiplying the SD of the measurement by \(\sqrt{2}\) (see Supplementary Materials for more information). An example is shown in Figure 3, revealing the relation between two predictions at different time points. For a complete overview of the PPDs for all states, see the Supplementary Materials. The relation between the PPDs of different states was explored. This relation between two states at the indicated time points is shown in more detail in both scatter plot and 2D histogram form in Figure 4. The former shows the actual samples from the PPD for one point in time. Here each dot represents a simulated value for one parameter set from the MCMC chain. As shown in the figure, these different states are often non-linearly related at specific points in time. The associated 2D histogram corresponds to the same information interpreted as probability density. Considering state 3 as observable and state 4 as prediction, while assuming a measurement accuracy of \(\sigma = 10/\sqrt{2}\) for \(x_3\), it can be observed that a significant decrease in variance can be attained during the rise of state 3. Measuring state 3 at the peak value however, results in a smaller variance reduction. A few things can be observed. In order for the measurement to be useful, there should be a correlation between the measurement and the prediction of interest. Additionally, the uncertainty in both should be large enough. Since all predictions of state 3 start with an initial condition of zero, this implies that the uncertainty at this point is low. Therefore, an additional measurement at \(t=0\) would not yield appreciable variance reduction which is also reflected by the fact that the SVR starts at a value of zero.

In order to demonstrate the flexibility of our method it was decided to perform OED for a quantity that depends on the model predictions in a highly non-linear fashion, namely the time to peak for the concentration of dimerized STAT in the nucleus (state 4). The time to peak was computed for the state 4 prediction for each parameter set from the posterior parameter distribution. We assumed that all states except state 4 are measurable with an accuracy of \(\sigma = 10/\sqrt{2}\). As potential measurements we also included the two sums of states as measured in earlier experiments. The experiment space was sampled using a Monte Carlo approach, uniformly sampling the experiment space.

This sampling is shown in Figure 4 where the SVR is shown for several combinations of two measurements. In this figure each axis corresponds to a potential measurement. Different model outputs (potential measurements) are separated using grid lines, while...
reductions are attained but the timing accuracy required for a given experiment could consider measuring state three and one instead. Here smaller reductions are obtained by measuring state one at an early and late time point. This result underlines the benefit of being able to combine multiple measurements in the OED. Furthermore, the analysis reveals that the largest reduction in variance could be obtained by measuring state one at an early and late time point. This result underlines the benefit of being able to combine multiple measurements in the OED. Interestingly, performing the experimental design for two measurements revealed that the largest reduction in variance could be attained but the timing accuracy required for a reasonable reduction is less stringent. Additionally, we investigated how the bounds of the priors on the non-identifiable kinetic rates affected our experimental design by widening them. This revealed that the EVRs obtained when measuring state 2 or 3 in combination with state 1 were more robust (for more information see the Supplementary Materials).

Since both error models in this case are Gaussian, the same analysis can be performed using the LVR (which for T = 1000 samples is about 100-fold faster). The resulting sampling is shown in Figure 5. Qualitatively, the results agree well with those in Figure 4, revealing its applicability as an initial sampling step. Information gained from an initial LVR sweep can subsequently be used to sample only relevant regions of the experiment design space. Another example can be found in the Supplementary Materials.

### 5 DISCUSSION AND CONCLUDING REMARKS

In this article, we have outlined a flexible method to perform experimental design. Here a Gaussian probability density function was used to model the uncertainties. Note, however, that our method is not restricted to such error models. In statistical parameter inference it is important to determine which error model to use for each experiment as this will define the appropriate likelihood function. If the likelihood function cannot be computed explicitly then approximate Bayesian methods can provide a solution.

In the OED the timing of the new measurement is assumed instantaneous (infinitely accurate). It remains an open but relevant challenge to incorporate temporal inaccuracies in the current framework. It is expected that when timing is more error prone and explicitly accounted for, experiments that are only effective during brief time intervals will be marked as less beneficial for variance reduction.

In our method, we base the experimental design on the expected value of a distribution of variance reductions. However, since the entire distribution of possible variance reductions has been computed one could also consider incorporating information regarding the accuracy of this estimate into the selection process. Finding a sensible trade off between EVR and its inaccuracy considering the prediction uncertainty remains an open topic for further research.

In order to obtain the posterior distribution, the parameters are required to be either identifiable or restricted by means of
a finite prior distribution. Even for a small model, identifying a sufficient amount of data, the posterior distribution is relatively insensitive to the assumed priors. It is important to vary the priors or determine the effect of a measurement on the assumed prior. Note that the latter strongly depends on the initial prior, which should be chosen sufficiently wide to cover all potential parameter regimes. The number of samples required to order to get a reliable estimate is highly problem specific. Convergence is hard to assess and only non-convergent runs of the MCMC can be diagnosed (Calderhead and Girolami, 2009). Once convergence has been attained, one should verify that the model sufficiently describes the acquired data as EVRs will be based on model predictions.

The method is not limited to a specific family of distributions for the parameters and model predictions. However, strong distributions (such as high variance logarithmic distributions) can be problematic. The reason for this is that in such cases estimates from a small sample of the tail are quite unreliable and give a poor description of the distribution. Therefore it is important to posteriori visually inspect the distribution at the determined optimal. In the case of heavy tailed distributions, it is beneficial to perform a transformation of the PPD before the experimental design.

Consider performing a new measurement as illustrated in Figure 4. The estimation of the measurement efficacy multiplying samples of the old posterior with new weights in order to keep track of the quality of the estimation. When the before and after a new experiment is very different, many sample weights will be very low and a large fraction of the samples will contribute only negligibly to the estimation of the new variance. It follows that such an estimate will be poor. We monitor this degeneracy by estimating the effective sample size (ESS) below (Del Moral et al., 2009).

\[
\text{ESS} = \frac{\sum_{k=1}^{N} G(t, \tilde{u}(t), \tilde{y}_{k}, \theta)}{\sum_{k=1}^{N} G(t, \tilde{u}(t), \tilde{y}_{k}, \theta)^2}
\]

We compute a distribution of ESS values (one for each incorporated sample) which we characterize by its median value. This ESS gives a measure for the quality of the samples. In the case that the importance sampling distribution agrees well with the posterior, it should scale linearly with the number of included samples. When the values for the ESS are very low, the variance reduction can be inaccurate and it also implies that such a measurement would be very informative on an inferential perspective. This stems from the fact that the probability distribution would be much narrower. In such a case, it would be beneficial to perform the experiment and subsequently redo the MCMC step in such cases (for more information see the Supplementary Materials).

Obtaining the PPDs as well as performing the experiment is computationally expensive. For the former, model time is a primary concern which can be significantly reduced by using compiled simulation code (see COPASI (Hoops et al., 2006); ABC-SysBio (Lluch et al., 2010); Potters Wheel and Timme). Additionally, posteriori visual inspection of the LVR and Calderhead design part, contribution of each experiment into the estimation of the posterior distribution is limited. Therefore it is important that the importance sampling distribution agrees well with the posterior, it should scale linearly with the number of included samples. When the values for the ESS are very low, the variance reduction can be inaccurate and it also implies that such a measurement would be very informative on an inferential perspective. This stems from the fact that the probability distribution would be much narrower. In such a case, it would be beneficial to perform the experiment and subsequently redo the MCMC step in such cases (for more information see the Supplementary Materials).

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