

Efficient Likelihood Approximation via Gaussian Processes

With an Application to a *P. Vivax* Malaria Model

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Malaria

- ▶ 600,000 deaths/year, 75% children under 5
- ▶ Two main species *P. vivax* and *P. falciparum*
- ▶ *P. falciparum* main cause of death, but *P. vivax* traditionally underestimated.
- ▶ As *P. falciparum* decrease, *P. vivax* cases increase

P. vivax has Dormant Stage

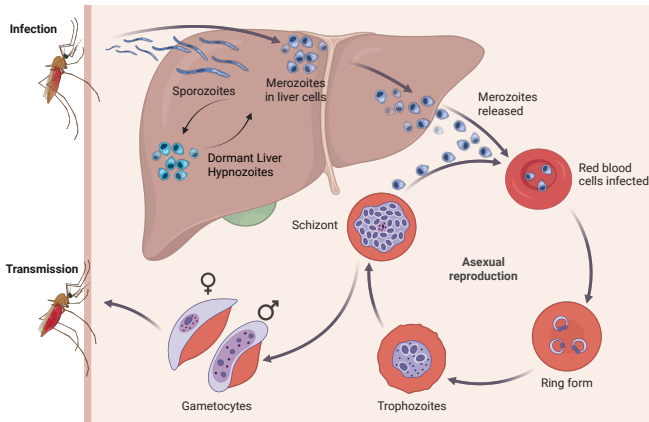


Figure: *P. vivax* lifecycle. Created with BioRender.com

Vivax Model - Champagne et. al

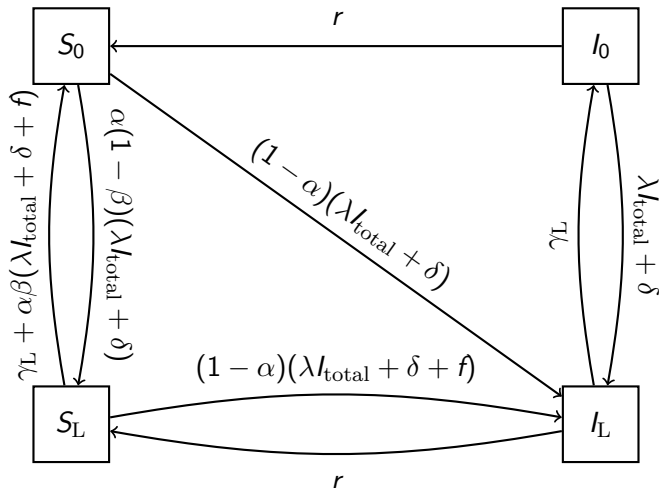


Figure: Champagne et al. 2022 *P. vivax* model

Champagne Model Parameters

- ▶ α : proportion of those infected who clear blood stage infections through treatment
- ▶ β : proportion of those cleared of blood stage infection who are also cleared of liver stage parasites (radical cure)
- ▶ λ : rate of infection
- ▶ γ_L : rate of liver stage disease clearance
- ▶ f : rate of relapse
- ▶ r : rate of blood stage clearance
- ▶ $\delta = 0$ importation rate (fixed)

Ordinary Differential Equations - Champagne et. al

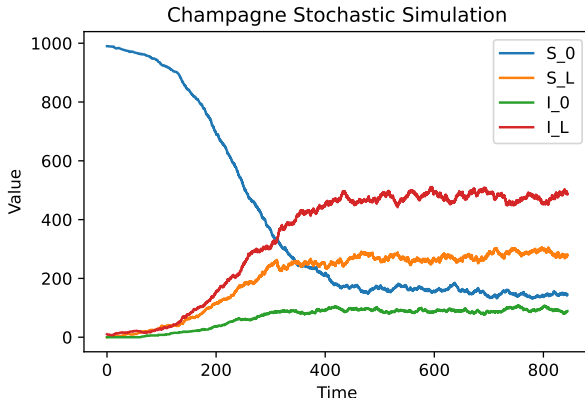
$$\begin{aligned}\frac{dI_L}{dt} = & (1 - \alpha)(\lambda I_{\text{total}} + \delta)(S_0 + S_L) + (\lambda I_{\text{total}} + \delta)I_0 \\ & + (1 - \alpha)fS_L - \gamma_L I_L - rI_L\end{aligned}$$

$$\frac{dI_0}{dt} = -(\lambda I_{\text{total}} + \delta)I_0 + \gamma_L I_L - rI_0$$

$$\begin{aligned}\frac{dS_L}{dt} = & -(1 - \alpha(1 - \beta))(\lambda I_{\text{total}} + \delta + f)S_L \\ & + \alpha(1 - \beta)(\lambda I_{\text{total}} + \delta)S_0 - \gamma_L S_L + rI_L\end{aligned}$$

$$\begin{aligned}\frac{dS_0}{dt} = & -(1 - \alpha\beta)(\lambda I_{\text{total}} + \delta)S_0 + (\lambda I_{\text{total}} + \delta)\alpha\beta S_L \\ & + \alpha\beta fS_L + \gamma_L S_L + rI_0\end{aligned}$$

Example Simulation



The Problem

- ▶ How to parametrise such a model.

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- ▶ Simulations take long time (and models get a lot more complicated)

Notation

- ▶ θ vector of parameters - e.g. $[\alpha, \beta, \gamma_L, \lambda, f, r]^T$
- ▶ \mathbf{Y}_{obs} : a (summary) vector of observed data e.g. (weekly) incidence, prevalence, (monthly) hospitalisations

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- ▶ \mathbf{Y}_{θ} : a random vector of model statistics for given θ .

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- ▶ There would be an explicit form for the likelihood:

$$\mathcal{L}(\theta|\mathbf{Y}_{\text{obs}}) := \Pr(\mathbf{Y}_{\text{obs}}|\theta)$$

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- ▶ Off to the pub

Or not...

- ▶ Explicit likelihoods often don't exist/are intractible
 - ▶ Champagne model
 - ▶ Agent based models.

A Standard Bayesian Solution

- ▶ Approximate Bayesian Computation (ABC)
 1. Sample θ_i from prior
 2. Run model and observe \mathbf{Y}_{θ_i}
 3. Accept or reject θ_i run based on how well \mathbf{Y}_{θ_i} 'matches' \mathbf{Y}_{obs} .

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2. Rescale \mathbf{Y} s, and use discrepancy function $D : \mathbb{R}^d \times \mathbb{R}^d \rightarrow \mathbb{R}$
e.g. p -norm

$$D(\mathbf{Y}_{\theta_i}, \mathbf{Y}_{\text{obs}}) := \left(\sum_{j=1}^d |\{\mathbf{Y}_{\theta_i}\}_j - \{\mathbf{Y}_{\text{obs}}\}_j|^p \right)^{1/p}$$

Discrepancy Function

$\mathcal{D}(\theta) := D(\mathbf{Y}_\theta, \mathbf{Y}_{\text{obs}})$ how 'close' our model is to the observed data using parameters θ

1. Sample θ_i from prior
2. Run model
3. Accept θ_i if $\mathcal{D}(\theta_i) < \varepsilon$.

Overall Idea of my Research

- ▶ ABC fixes one problem but leaves another:
 - ▶ Don't need $\mathcal{L}(\theta)$.
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- ▶ ABC fixes one problem but leaves another:
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- ▶ $\mathcal{D}(\theta), \mathcal{D}(\theta')$ will be highly correlated when θ is near θ' .
 - ▶ Gaussian Processes

Gaussian Process Setup

A common assumption is that

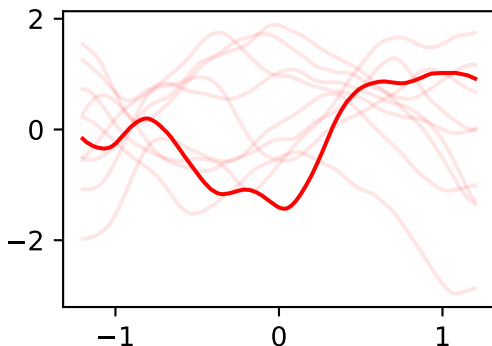
$$\begin{bmatrix} \mathbb{E}(\mathcal{D}(\boldsymbol{\theta}_1)) \\ \vdots \\ \mathbb{E}(\mathcal{D}(\boldsymbol{\theta}_n)) \end{bmatrix} \sim \text{MVN}(\mathbf{0}, \mathbf{K})$$

where $\mathbf{K}_{ij} = k(\boldsymbol{\theta}_i, \boldsymbol{\theta}_j)$ for some covariance kernel k that decays to 0 as $\boldsymbol{\theta}_i$ is further away than $\boldsymbol{\theta}_j$.

Gaussian Processes on \mathbb{R}^d

Definition (Gaussian Process)

A collection of random variables $\{f(\mathbf{x})\}_{\mathbf{x} \in \mathbb{R}^d}$ is a Gaussian process if all finite dimensional distributions are multivariate normal distributed.



Equivalent GP definition

Definition (Gaussian Process)

A collection of random variables $\{f(\mathbf{x})\}_{\mathbf{x} \in \mathbb{R}^d}$ is a Gaussian process if there is a function $m : \mathcal{X} \rightarrow \mathbb{R}$ and covariance kernel $k : \mathcal{Y} \times \mathcal{Y} \rightarrow \mathbb{R}$ such that for all $\{\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_n\}$,

$$\begin{bmatrix} f(\mathbf{x}_1) \\ f(\mathbf{x}_2) \\ \vdots \\ f(\mathbf{x}_n) \end{bmatrix} \sim \text{MVN} \left(\begin{bmatrix} m(\mathbf{x}_1) \\ m(\mathbf{x}_2) \\ \vdots \\ m(\mathbf{x}_n) \end{bmatrix}, \mathbf{K} \right)$$

where $\mathbf{K}_{ij} := k(\mathbf{x}_i, \mathbf{x}_j)$.

k Determines Smoothness

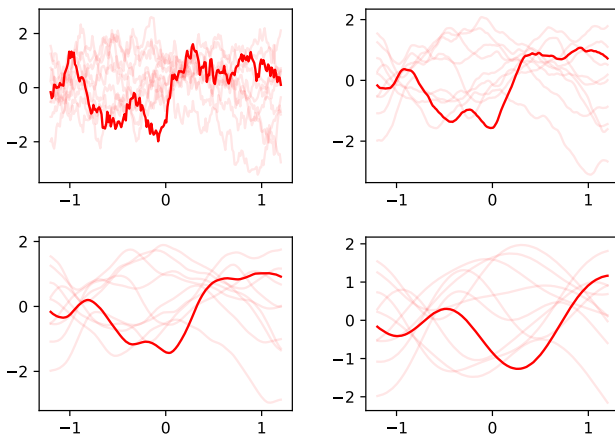


Figure: Matérn 1/2, 3/2, 5/2, and squared exponential kernels.

Gaussian Process Continuity

- ▶ Induce continuity by forcing $k(\mathbf{x}, \mathbf{x}') \rightarrow \text{Var}(f(\mathbf{x}))$ (hence $\text{Cor}(f(\mathbf{x}), f(\mathbf{x}')) \rightarrow 1$) as $\mathbf{x} \rightarrow \mathbf{x}'$.

Common Covariance Kernels

- ▶ Matérn Kernel with hyperparameter $\nu : \lfloor \nu \rfloor$ times mean square differentiable.
- ▶ $\nu \rightarrow \infty$ - infinitely mean square differentiable squared exponential covariance kernel (strong assumption)

$$k(x, x') = \sigma_k^2 \exp\left(-\frac{\|x - x'\|^2}{2\ell^2}\right)$$

k Determines Class of Functions

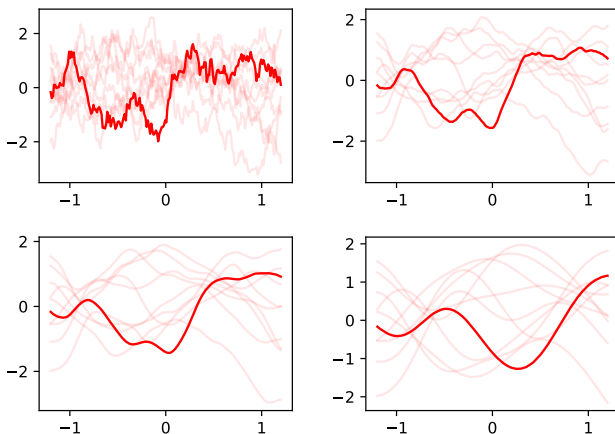
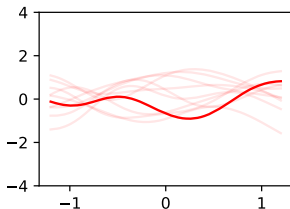


Figure: Matérn 1/2, 3/2, 5/2, and squared exponential kernels.

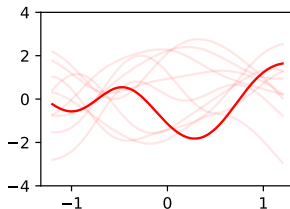
Kernel Hyperparameters

- ▶ Matérn and squared exponential kernel can both be written in the form $k(\mathbf{x}, \mathbf{x}') = \sigma_k^2 \kappa(\|\mathbf{x}, \mathbf{x}'\|/\ell)$
- ▶ $1/\ell$ rate of covariance decay
- ▶ $\sigma_k = \text{Var}(f(\mathbf{x}))$

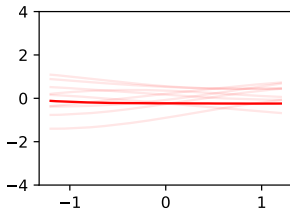
Kernel Hyperparameters



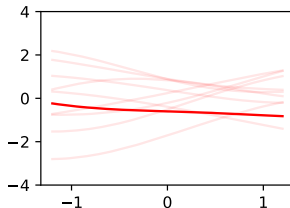
(a) $\ell = 1/2$ $\sigma_k^2 = 1/2$



(b) $\ell = 1/2$ $\sigma_k^2 = 2$



(c) $\ell = 2$ $\sigma_k^2 = 1/2$



(d) $\ell = 2$ $\sigma_k^2 = 2$

Discrepancy Function Context

- ▶ Long term play: use a Gaussian process as a surrogate model for $\mathbb{E}[\mathcal{D}(\boldsymbol{\theta})]$

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- ▶ Long term play: use a Gaussian process as a surrogate model for $\mathbb{E}[\mathcal{D}(\boldsymbol{\theta})]$
- ▶ What if we have observations already?

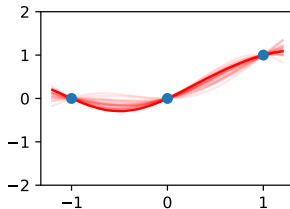
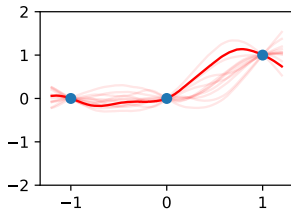
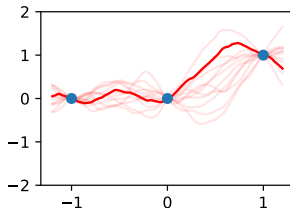
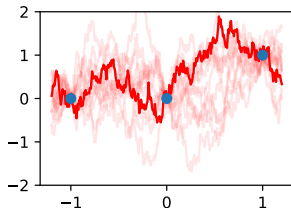
Gaussian Process Regression

$$\begin{bmatrix} f(\mathbf{x}) \\ f(\mathbf{x}_*) \end{bmatrix} \sim \text{MVN} \left(\begin{bmatrix} m(\mathbf{x}) \\ m(\mathbf{x}_*) \end{bmatrix}, \begin{bmatrix} K & K_* \\ K_*^T & K_{**} \end{bmatrix} \right)$$

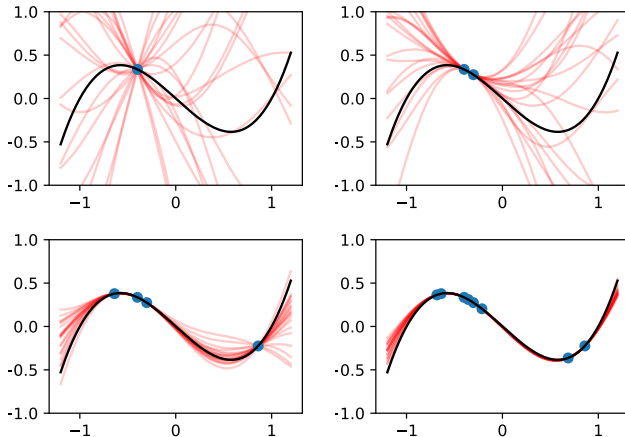
implies

$$f(\mathbf{x})|f(\mathbf{x}_*) \sim \text{MVN} \left(m(\mathbf{x}) + K_* K_{**}^{-1} (f(\mathbf{x}_*) - m(\mathbf{x}_*)), K - K_* K_{**}^{-1} K_*^T \right).$$

Conditioning Gaussian Processes



GP regression on $x(x-1)(x+1)$

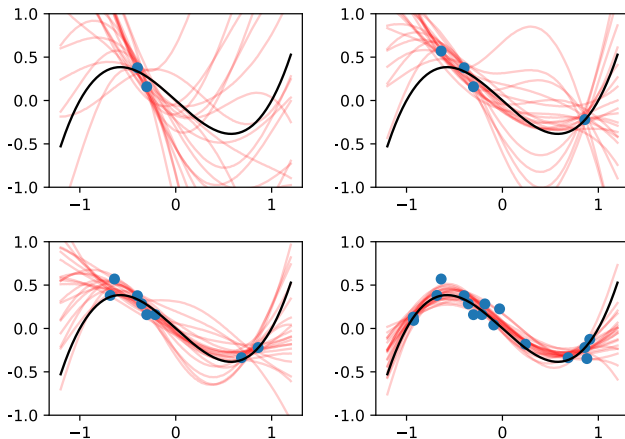


Normal observation noise

If observations actually $f(\mathbf{x}_i) + \varepsilon_i$, with $\varepsilon_i \sim N(0, \sigma_o^2)$ i.i.d., then

$$\begin{bmatrix} f(\mathbf{x}_1) + \varepsilon_1 \\ \vdots \\ f(\mathbf{x}_n) + \varepsilon_n \end{bmatrix} \sim \text{MVN} \left(\begin{bmatrix} m(\mathbf{x}_1) \\ \vdots \\ m(\mathbf{x}_n) \end{bmatrix}, \mathbf{K} + \sigma_o^2 \mathbf{I}_n \right)$$

GP regression on $x(x-1)(x+1) + \varepsilon$, $\varepsilon \sim N(0, \sigma_o^2)$



Key Assumptions

- ▶ $\mathcal{D}(\boldsymbol{\theta}) \stackrel{d}{\approx} \mathcal{D}(\boldsymbol{\theta}')$ for $\boldsymbol{\theta}, \boldsymbol{\theta}'$ close.
- ▶ $\mathcal{D}(\boldsymbol{\theta})$ approximately distributed $N(\cdot, \sigma_0^2)$ with σ_0^2 independent of $\boldsymbol{\theta}$.
- ▶ $\mathbb{E}[\mathcal{D}(\boldsymbol{\theta})]$ can be well approximated by a Gaussian process.

Key Idea

Approximate $\mathcal{D}(\boldsymbol{\theta})$ with $\mathcal{D}_{GP}(\boldsymbol{\theta})$, a Gaussian process with observation noise.

1. Sample θ_i from prior
2. Run model
3. Accept θ_i if $\mathcal{D}(\theta_i) < \varepsilon$.

Approximate ABC...??

1. Sample θ_i from prior
2. Run model
3. Accept θ_i if $\mathcal{D}_{\mathcal{GP}}(\theta_i) < \varepsilon$.

Synthetic Likelihood

The probability of drawing and accepting θ under the ABC is

$$\Pr(\mathcal{D}_{\mathcal{GP}}(\theta) < \varepsilon) \Pr(\theta)$$

and hence we consider $L(\theta|\mathbf{Y}_{\text{obs}}) := \Pr(\mathcal{D}_{\mathcal{GP}}(\theta) < \varepsilon)$ a synthetic likelihood - an approximation of the true likelihood $\mathcal{L}(\theta|\mathbf{Y}_{\text{obs}})$

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 - ▶ $\mathcal{D}(\boldsymbol{\theta})$ approximately distributed $LN(\cdot, \sigma_0^2)$ with σ_0^2 independent of $\boldsymbol{\theta}$.
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 - ▶ $\mathbb{E}[\ln \mathcal{D}(\boldsymbol{\theta})]$ can be well approximated by a Gaussian process.
- ▶

$$\Pr(\mathcal{D}(\boldsymbol{\theta}_i) < \varepsilon) \approx \Pr(d_{\mathcal{GP}}(\boldsymbol{\theta}_i) < \ln \varepsilon)$$

Where to sample $\mathcal{D}(\theta)$

- ▶ To generate a reliable $\mathcal{D}_{\mathcal{GP}}$, we need to sample widely
- ▶ Generating $\mathcal{D}(\theta)$ still costly...
- ▶ Therefore sample where:

Where to sample $\mathcal{D}(\theta)$

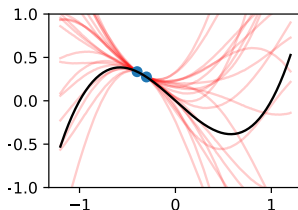
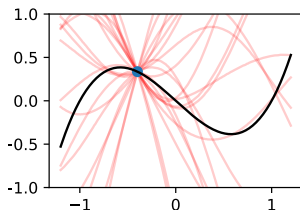
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Bayesian Acquisition Functions

- Quantify where we expect $\mathcal{D}(\theta)$ small and highly unknown with Bayesian acquisition function A , and choose $\arg \min_{\theta} A(\theta)$

Bayesian Acquisition Functions

- ▶ Gutmann and Cor 2016 use lower confidence bound

$$A_{\text{LCB}}(\boldsymbol{\theta}) := \mu(\boldsymbol{\theta}) - \eta_t \sqrt{v(\boldsymbol{\theta})}$$

- ▶ $\mu(\boldsymbol{\theta})$, $v(\boldsymbol{\theta})$ are posterior mean and variance of $D_{\mathcal{GP}}(\boldsymbol{\theta})$
- ▶ η_t a slowly increasing function in t , the number of previous samples

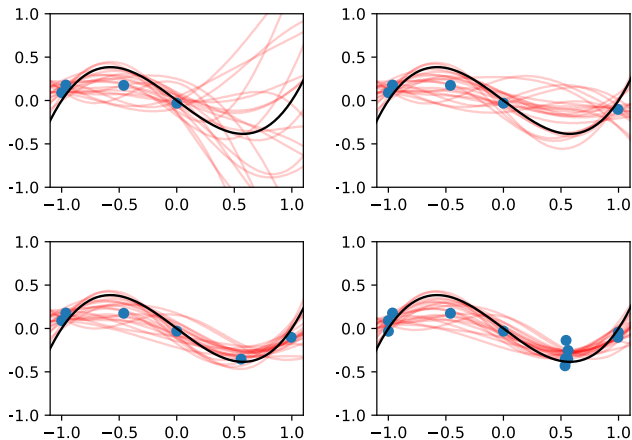
Bayesian Acquisition Functions

- ▶ Expected information

$$A_{\text{EI}}(\boldsymbol{\theta}) := \mathbb{E}[\min(\mu_{\min} - \mathcal{D}_{\mathcal{GP}}(\boldsymbol{\theta}), 0)]$$

- ▶ $\mu_{\min} := \min_{\boldsymbol{\theta}} \mu(\boldsymbol{\theta})$

Lower Confidence Bound



Vivax Model - Champagne et. al

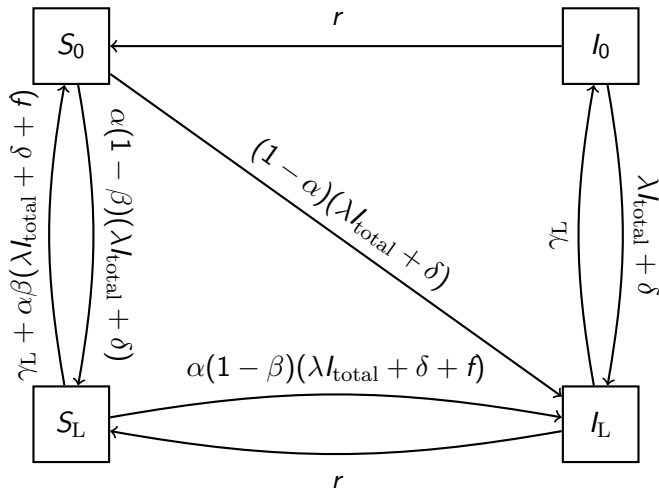


Figure: Champagne et al. 2022 *P. vivax* model

Champagne Model Parameters

- ▶ α : proportion of those infected but cleared of blood stage infections (through treatment)
- ▶ β : a further proportion that are also cleared of liver stage parasites, given that they were also cleared of blood stage infection (radical cure)
- ▶ λ : the rate of infection
- ▶ γ_L : rate of clearance of liver stage disease
- ▶ f : rate of relapse
- ▶ r : rate of blood stage clearance

'Observed' Data

- ▶ 'Observed' data from one simulation of 10 initial infections, in a population of 1000 people using the parameters reported in Champagne et al. 2022.
- ▶ $\mathbf{Y}_{\text{obs}} := \{w_{\text{obs}}, p_{\text{obs}}, m_{\text{obs}}\}$
 - ▶ w_{obs} : weekly incidence around (stochastic) equilibrium
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 - ▶ m_{obs} : incidence in the first month of the epidemic
- ▶ $\mathcal{D}(\alpha, \beta, \gamma_L, \lambda, f, r)$ is the L_2 norm of the relative differences

$$\sqrt{\left(\frac{p - p_{\text{obs}}}{p_{\text{obs}}}\right)^2 + \left(\frac{m - m_{\text{obs}}}{m_{\text{obs}}}\right)^2 + \left(\frac{w - w_{\text{obs}}}{w_{\text{obs}}}\right)^2}$$

GP choices

- ▶ $\mathcal{D}(\alpha, \beta, \gamma_L, \lambda, f, r)$ is the L_2 norm of the relative differences

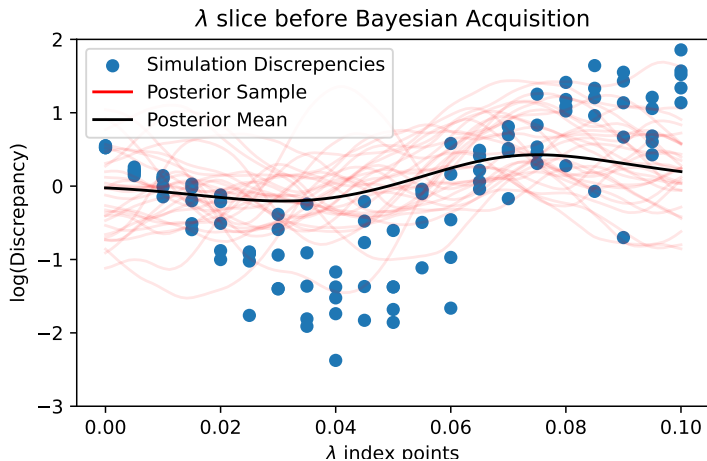
$$\sqrt{\left(\frac{p - p_{\text{obs}}}{p_{\text{obs}}}\right)^2 + \left(\frac{m - m_{\text{obs}}}{m_{\text{obs}}}\right)^2 + \left(\frac{w - w_{\text{obs}}}{w_{\text{obs}}}\right)^2}$$

- ▶ \mathcal{GP} choices
 - ▶ Modelled In \mathcal{D} as a Gaussian process
 - ▶ Matern kernel with $\nu = 5/2$
 - ▶ $\ell, \sigma_k^2, \sigma_o^2$ selected by leave one out cross validation.

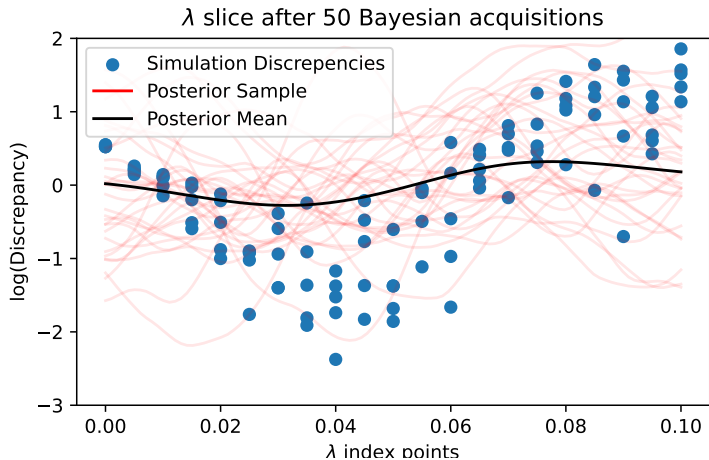
How did it go?



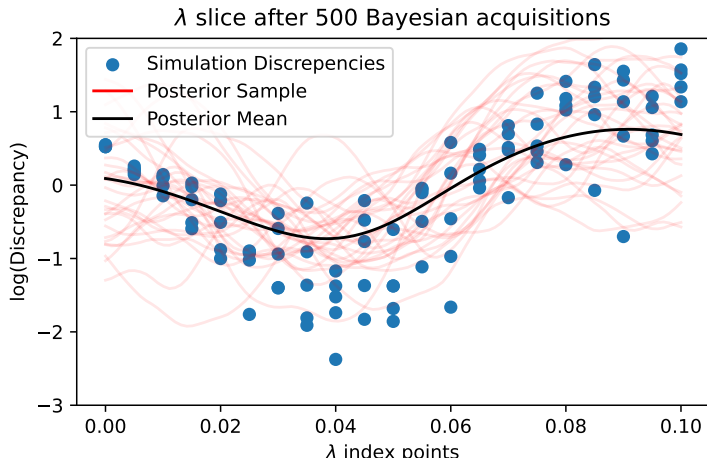
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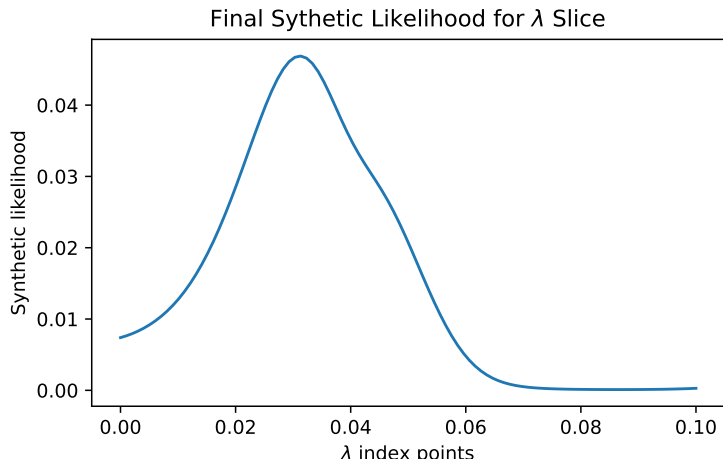
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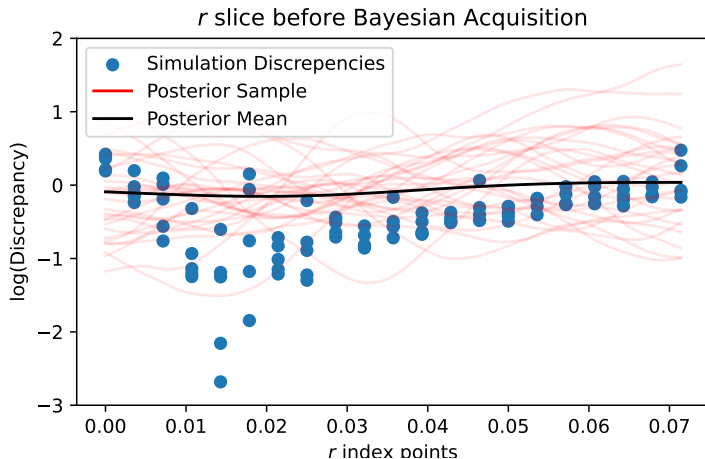
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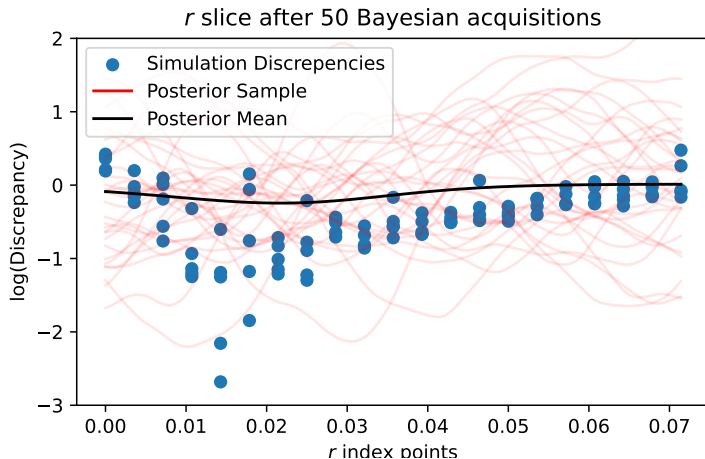
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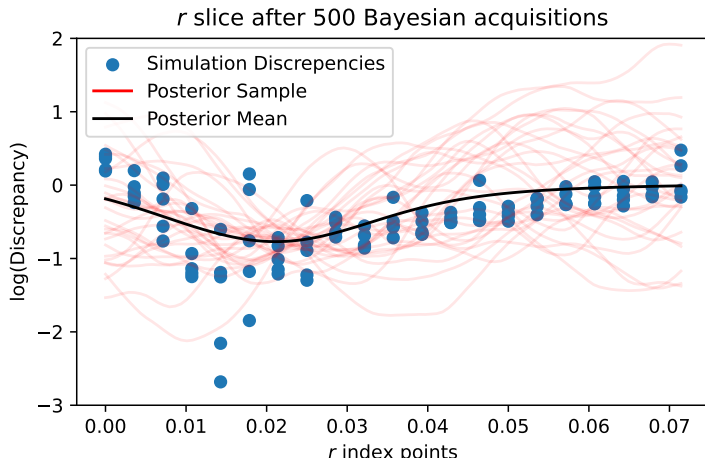
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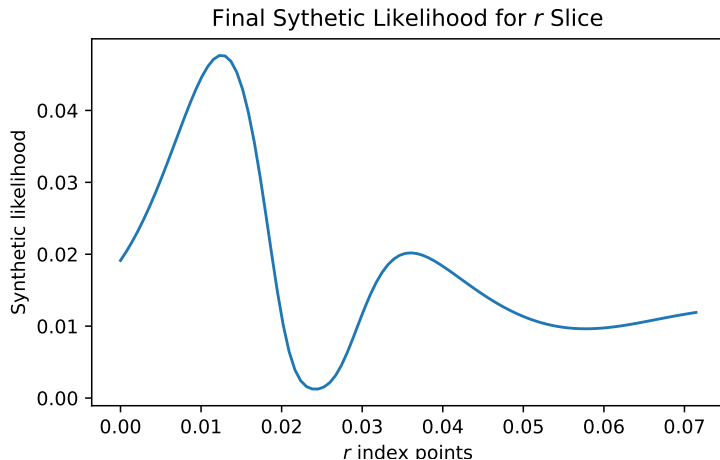
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How did it go?



Discussion

- ▶ Observation variance is considered constant across the GP (or log GP)
 - ▶ Particularly a problem at the threshold
- ▶ Assumes that normal/log-normal distribution approximates $\mathcal{D}(\theta)$
- ▶ Jumps where there is threshold/bifurcation behaviour
 - ▶ Student t -Process?

Thanks to

- ▶ Eamon Conway
- ▶ Jennifer Flegg



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