Bayesian analysis of growth curves using mixed models defined by stochastic differential equations

Sophie Donnet¹
Ceremade, Université Dauphine, France
*email: sophie.donnet@ceremade.dauphine.fr

and

Jean-Louis Foulley²
INRA-GABI, PSGen, CR Jouy, France
*email: jean-louis.foulley@jouy.inra.fr

and

Adeline Samson³
Laboratoire MAP5, Université Paris Descartes, France
*email: adeline.samson@parisdescartes.fr

Summary: Growth curve data consist of repeated measurements of a continuous growth process over time in a population of individuals. These data are classically analyzed by nonlinear mixed models. However, the standard growth functions used in this context prescribe monotone increasing growth and can fail to model unexpected changes in growth rates. We propose to model these variations using stochastic differential equations (SDEs) that are deduced from the standard deterministic growth function by adding random variations to the growth dynamics. A Bayesian inference of the parameters of these SDE mixed models is developed. In the case when the SDE has an explicit solution, we describe an easily implemented Gibbs algorithm. When the conditional distribution of the diffusion process has no explicit form, we propose to approximate it using the Euler-Maruyama scheme. Finally, we suggest to validate the SDE approach via criteria based on the predictive posterior distribution. We illustrate the efficiency of our method using the Gompertz function to model data on chicken growth, the modeling being improved by the SDE approach.
Key words: Bayesian estimation; Euler-Maruyama scheme; Gompertz model; Growth curves; Mixed models; Predictive posterior distribution; Stochastic differential equation
1. Introduction

Growth curve data consist of repeated measurements of a continuous growth process over time among a population of individuals. In agronomy, growth data allow differentiating animal or vegetal phenotypes by characterizing the dynamics of the underlying biological process. In gynecology or pediatrics, height and weight of fetus/children are regularly recorded to control their development. The parametric statistical approach commonly used to analyze these longitudinal data is mixed model methodology (Huggins and Loesch, 1998). The regression function of this mixed model is classically a parametric growth function, such as the Gompertz, logistic, Richards or Weibull functions (Zimmerman and Núñez-Antón, 2001) which prescribe monotone increasing growth, whatever the parameter values. These models have proved their efficiency in animal genetics (Hou et al., 2005; Jaffrézic et al., 2006, e.g.) and in pediatrics (Spyrides et al., 2008, e.g.). However, as pointed out by Davidian and Giltinan (2003), the used function may not capture the exact process, as responses for some individuals may display some local fluctuations such as weight decreases or growth slow down. These phenomena are not due to error measurements but are induced by an underlying biological process that is still unknown today. In animal genetics, a wrong modeling of these curves could affect the genetic analysis. In fetal growth, the detection of growth slow down is a crucial indicator of fetal development problems.

The aim of this paper is to model these variations in growth rate using a stochastic differential equation (SDE) whose solution is the regression term of the mixed model. More precisely, each growth function is defined as the solution of an ordinary differential equation (ODE); this ODE models the growth rate. We suggest to add a random perturbation to the ODE, resulting in an SDE. Thus, the growth rate varies randomly around the mean dynamics. In this paper, we propose and study Bayesian estimators for mixed models defined by SDEs. Estimator properties are illustrated for simulated and real animal growth data.
Parametric estimation by maximum likelihood of SDE with random parameters (without measurement noise) has been studied by Ditlevsen and De Gaetano (2005). However, estimation of SDE mixed models (including the measurement noise modeling) has received little attention. Overgaard et al. (2005) and Tornoe et al. (2005) proposed estimators based on an extended Kalman filter, but the algorithm convergence was not proved. Donnet and Samson (2008) proposed an EM-based estimator and proved the convergence of their algorithm. Whereas the Bayesian point of view is widely used on standard growth curves, Bayesian estimation of SDE mixed models has not been much investigated. Cano et al. (2006) computed the posterior distribution by approximating the diffusion process by an Euler scheme. Oravecz et al. (in press) studied the Bayesian estimation of an Ornstein-Uhlenbeck process with random parameters. In this paper, we propose either to use a judicious transformation of the SDE to compute the exact conditional distribution of the diffusion process, or, if it is not possible, to approximate the diffusion by the Euler-Maruyama scheme. Then we propose a Gibbs algorithm to simulate the exact or the approximate posterior distributions. In the case of approximation by the Euler scheme, we control the error induced by this scheme on the posterior distributions. Finally, we adapt the computation of the posterior predictive distributions to validate the SDE mixed model (Meng, 1994).

Section 2 presents the classical nonlinear mixed model and the mixed model defined by SDEs. We discuss the choice of the volatility term in the SDEs. In Section 3, we suggest some prior specifications and posterior computation. We also present the Euler-Maruyama scheme. Section 4 shows how to validate the SDE mixed model using posterior predictive distributions. In Section 5, the theory is illustrated with the particular case of the Gompertz function applied on chicken growth data.
2. Models and notations

2.1 Nonlinear mixed models

Let \( y = (y_i)_{1 \leq i \leq n} = (y_{ij})_{1 \leq i \leq n, 1 \leq j \leq n_i} \) denote the data, where \( y_{ij} \) is the noisy measurement of the observed biological process for individual \( i \) at time \( t_{ij} \), for \( i = 1, \ldots, n, j = 0, \ldots, n_i \).

In classical mixed models, the process is modeled by a deterministic function, depending on individual random parameters. Formally, the classical nonlinear mixed model is defined as:

\[
\begin{align*}
y_{ij} &= f(\phi_i, t_{ij}) + \epsilon_{ij}, \quad \epsilon_{ij} \sim \text{i.i.d. } \mathcal{N}(0, \sigma^2) \\
\phi_i &\sim \mathcal{N}(\mu, \Omega)
\end{align*}
\]

with \( f \) being a parametric deterministic function and \( \phi = (\phi_i)_{1 \leq i \leq n} \) the \( p \)-vectors of individual parameter vectors. The \( \phi_i \) are assumed to be independently and identically normally distributed with expectation \( \mu \) and variance \( \Omega \). The \( \epsilon_{ij} \) are the residual errors, assumed to be independently and identically normally distributed with null mean and variance \( \sigma^2 \).

For growth curve data, \( f \) is classically one of the four most famous parametric functions modeling growth curves, namely the logistic, the Gompertz, the Richards and the Weibull functions. Each of them can be written as the solution of an ordinary differential equation (ODE) describing the evolution of growth rate, which are respectively:

\[
\begin{align*}
f'(t) &= Cf(t) \left[ 1 - \frac{1}{A} f(t) \right], \quad f(0) = \frac{A}{1+B} \quad \text{(Logistic)} \\
f'(t) &= BCe^{-Ct} f(t), \quad f(0) = Ae^{-B} \quad \text{(Gompertz)} \\
f'(t) &= \frac{BCDe^{-Ct}}{1 + Be^{-Ct}} f(t), \quad f(0) = \frac{A}{(1+B)\sigma} \quad \text{(Richards)} \\
f'(t) &= DCl^{D-1}(A - f(t)), \quad f(0) = A - B \quad \text{(Weibull)}
\end{align*}
\]

where \( A, B, C, D \) are non-negative parameters. \( A \) is the upper asymptote, \( C \) and \( D \) are growth rate parameters. All four models prescribe monotone increasing curves. More generally, if \( \phi \) denotes the parameter vector (either \( (A, B, C) \), \( (A, B, C, D) \) or a well-chosen parametrization), \( f \) is the solution of the following ODE:
\[
\frac{\partial f(\phi, t)}{\partial t} = F(f, t, \phi), \quad f(\phi, 0) = f_0(\phi)
\]  

(6)

2.2 Nonlinear mixed models defined by stochastic differential equations

The classical nonlinear mixed model is extended by replacing the regression function by a stochastic process. We propose to introduce a stochastic term in the ODE (6) to take into account individuals whose growth curve suffers from an unexpected growth rate change. Growth curve is thus described by a random process, denoted \( Z_t \), solution of the SDE:

\[
dZ_t = F(Z_t, t, \phi)dt + \Gamma(Z_t, \phi, \gamma^2)dW_t, \quad Z(t = 0) = Z_0(\phi)
\]  

(7)

where \( W_t \) is a Brownian motion. \( \Gamma(Z_t, \phi, \gamma^2) \) is the volatility function depending on the unknown parameter \( \gamma^2 \). The nonlinear mixed model defined by an SDE is thus:

\[
y_{ij} = Z_{t_{ij}}(\phi_i) + \varepsilon_{ij}, \quad \varepsilon_{ij} \sim_{i.i.d.} \mathcal{N}(0, \sigma^2)
\]

\[
dZ_t(\phi_i) = F(Z_t, t, \phi_i)dt + \Gamma(Z_t, \phi, \gamma^2)dW_t
\]

(8)

\[
\phi_i \sim \mathcal{N}(\mu, \Omega)
\]

In model (8), three fundamentally different noises are distinguished: the inter-subject variability \( \Omega \), the dynamic noise \( \gamma^2 \), reflecting the random fluctuations around the corresponding theoretical dynamic model, and the measurement noise \( \sigma^2 \) representing the uncorrelated part of the residual variability associated with assay or sampling errors.

Many types of volatility functions can be proposed to extend an ODE into an SDE (e.g. constant, square root or polynomial volatility). This choice depends on several considerations. If the observed biological process is non-negative, a volatility function ensuring the positivity of \( Z_t \) will be chosen. If biological reasons imply that a model parameter fluctuates along the experiment record, the volatility can be derived by adding a random perturbation to this parameter. If heteroscedastic variances have been used in an ODE modeling approach, a polynomial volatility can be chosen. Finally, algorithmic and computational constraints have to be considered: an SDE with explicit solution implies a simpler estimation scheme.
leading to good estimation properties (convergence of the algorithm to the true posterior
distribution) whereas an SDE without explicit solution implies additional computational
difficulties (use of an approximation scheme). As an example, we propose to use an affine
volatility function \( \Gamma(Z_t, \phi, \gamma^2) = \gamma Z_t \), for the logistic (2), Gompertz (3) and Richards (4)
models: the process \((Z_t)\) is then a log-Gaussian process (see Section 5.2 for more details).

3. Bayesian estimation

3.1 Prior specification

The Bayesian approach consists in the evaluation of the posterior distribution of the popu-
lation parameters \((\mu, \Omega), \sigma^2\) and the volatility \(\gamma^2\) for the SDE model. The first step is thus
the choice of the prior distributions. Usual diffuse prior distributions can be chosen but the
resulting posterior distributions may not be proper. Therefore, we suggest to use standard
prior distributions proposed, among others, by De la Cruz-Mesia and Marshall (2006) for
expectation or variance parameters in hierarchical models:

\[
\begin{align*}
\mu_k &\sim \mathcal{N}(m_{k}^{\text{prior}}, v_{k}^{\text{prior}}), \quad k = 1, \ldots, p \\
\Omega^{-1} &\sim W(R, p + 1), \quad 1/\sigma^2 \sim \Gamma(\alpha_{\sigma}^{\text{prior}}, \beta_{\sigma}^{\text{prior}})
\end{align*}
\]  

(9)

where \(W\) and \(\Gamma\) are respectively the Wishart and Gamma disributions. The \(\gamma^2\) parameter
controls the variance of the random perturbation. Many prior distributions can be considered
such as uniform, inverse-Gamma or Jeffreys. A sensitivity analysis is performed on the real
data set (see Section 5.5). In pratice the specification of hyperparameters \(m_{k}^{\text{prior}}, v_{k}^{\text{prior}}, R, \alpha_{\sigma}^{\text{prior}}, \beta_{\sigma}^{\text{prior}}\)
may be difficult. Therefore we choose the values of hyperparameters to obtain non-
informative priors.

3.2 Posterior computation

Since models (1) and (8) are non-linear, posterior distributions are not explicit and iterative
estimation procedures have to be used. For the ODE model (1), Gibbs sampling algorithms
Biometrics, –

including the sampling of the auxiliary random variables \( \phi_i \) under their conditional distributions have been proposed in the literature (Carlin and Louis, 2000, e.g.). These algorithms do not present any particular difficulties and are not detailed here. For the SDE model (8), we propose to use a Gibbs algorithm, including the sampling of the auxiliary random variables \( \phi_i \) and the vectors \( Z_i \) of realizations of process \( (Z_t) \) for each individual at each observation time. Let \( Z = (Z_1, \ldots, Z_n) \in \mathbb{R}^{(n_1+1)\times\cdots\times(n_n+1)} \) denote the vector of the \( n \) realizations. Hence the Gibbs sampling algorithm for the SDE model is outlined as follows:

• **Step 1:** initialize the iteration counter of the chain \( k = 1 \) and start with initial values \( \sigma^{-2}(0), \gamma^2(0), \mu(0), \phi(0), Z(0) \).

• **Step 2:** obtain \( \sigma^{-2(k)}, \gamma^2(k), \mu(k), \phi(k), Z(k) \) from \( \sigma^{-2(k-1)}, \gamma^2(k-1), \mu(k-1), \phi(k-1), Z(k-1) \) through successive generations of

1. \( Z^{(k)} \sim p(Z|\phi^{(k-1)}, \gamma^{-2(k-1)}, \sigma^{-2(k-1)}, y) \)
2. \( \phi^{(k)} \sim p(\phi|\sigma^{-2(k-1)}, \gamma^{-2(k-1)}, \mu(k-1), \Omega(k-1), Z^{(k)}, y_0) \) where \( y_0 = (y_{i0})_{i=1\ldots n} \)
3. \( \mu^{(k)} \sim p(\mu|\phi^{(k)}) \) and \( \Omega^{(k)} \sim p(\Omega|\phi^{(k)}) \)
4. \( \sigma^{-2(k)} \sim p(\sigma^{-2}|Z^{(k)}, \phi^{(k)}, y) \) and \( \gamma^{-2(k)} \sim p(\gamma^{-2}|Z^{(k)}, \phi^{(k)}) \)

• **Step 3:** change \( k \) to \( k + 1 \) and return to **Step 2** until convergence is reached.

Some conditional distributions are explicit. A Gamma prior distribution on \( \sigma^{-2} \) implies that \( p(\sigma^{-2}|Z^{(k)}, \phi^{(k)}, y) \) is a Gamma density. The prior distribution of \( p(\phi|\mu, \Omega) \) being Gaussian, the conditional distribution of \( \mu \) is Gaussian and the conditional distribution of \( \Omega \) is inverse Wishart. The conditional distributions on \( \phi, Z \) and \( \gamma^2 \) depend on the specific form of the SDE and are detailed in the particular example of the Gompertz model in Section 5. Depending on the model complexity, we may have to resort to Metropolis-Hastings algorithms. Moreover, for SDEs without explicit solution, the conditional distribution on \( Z \) has generally no closed form. In this case, we suggest to approximate the diffusion by the Euler-Maruyama scheme, which leads to Gaussian approximations of the transition densities. We then introduce an
approximate statistical model on which the posterior distributions are computed. Details are
given thereafter.

3.3 Posterior distribution using Euler-Maruyama approximation

The Euler-Maruyama scheme is presented for subject $i$. If the time intervals between the
observation instants are too great to obtain a good approximation of the transition density,
a natural approach is to introduce a set of auxiliary latent data points between every pair of
observations, as first proposed by Pedersen (1995). Let $t_{i0} = \tau_0 < \ldots < \tau_m < \ldots < \tau_{Mi} = t_{i,n_i}$
denote the equally spaced discretization of the time interval $[t_{i0}, t_{i,n_i}]$ and $h$ be the step size
of the discretization. We assume that, for all $j = 0 \ldots n_i$, there exists an integer $m_j$ verifying
$t_{ij} = \tau_{mj}$ ($m_0 = 0$ by definition). Using the approximated diffusion process from the Euler-
Maruyama scheme of step size $h$, an approximate statistical model is defined as:

$$y_{ij} = \hat{Z}_{mj}(\phi_i) + \varepsilon_{ij}, \quad \varepsilon_{ij} \sim_{i.i.d.} \mathcal{N}(0, \sigma^2)$$

$$\hat{Z}_m(\phi_i) = \hat{Z}_{m-1}(\phi_i) + h F(\hat{Z}_{m-1}, \tau_{m-1}, \phi_i) + \Gamma(\hat{Z}_{m-1}, \phi_i, \gamma^2) \sqrt{h} \xi_m, 1 \leq m \leq M_i,$$

$$\xi_m \sim_{i.i.d} \mathcal{N}(0, 1), \quad \phi_i \sim_{i.i.d} \mathcal{N}(\mu, \Omega)$$

For model (10), the conditional distribution of the approximate diffusion $\hat{Z}^h$ is Gaussian,
allowing to implement the previously presented Gibbs algorithm. The convergence of this
Gibbs algorithm is ensured by classical results (Carlin and Louis, 2000). However, this
Gibbs algorithm is performed on the approximate model (10), and computes the posterior
distribution $p^h(\theta|y)$ of model (10), with $\theta = (\mu, \Omega, \sigma^2, \gamma^2)$, instead of the original posterior
distribution $p(\theta|y)$. But, the error induced by the Euler scheme on the posterior distributions
can be controlled, as shown in the Supplementary materials.

4. Model validation

The goal in model checking is to monitor the quality of the proposed model, i.e. to determine
whether the observed data are representative of the type of data we might expect under this
Biometrics, –

model. Posterior predictive checks set this up by generating replicated data sets from the estimated posterior predictive distribution. These replicated data sets are then compared with the observed data. The function used to compare observed and replicated datasets is the discrepancy function; it depends on data and model parameters and is denoted $T(y, \eta)$, $\eta$ being used as generic notation for a function of the model parameters. It quantifies incompatibility of the model with the observed data. In our case, we consider for $T$ the $\chi^2$ discrepancy function, computed for the whole population at each age as:

$$T_j(y, \eta) = \sum_i \frac{(y_{ij} - \eta_{ij})^2}{\text{Var}(y_{ij} - \eta_{ij})}.$$ 

For the observation at time $t_{ij}$, we choose $\eta_{ij} = f(\phi_i, t_{ij})$ for the ODE model and $\eta_{ij} = Z_{ij}(\phi_i)$ for the SDE model. Consequently, for both models, $\text{Var}(y_{ij} - \eta_{ij}) = \sigma^2$.

We aim at comparing the posterior distribution $p(T_j(y, \eta) | y)$ of the observed data $y$ with the posterior distribution $p(T_j(y_{rep}, \eta) | y)$ where $y_{rep}$ denotes the replicated data drawn from the posterior predictive distribution $p(y_{rep} | y)$. A short version of that posterior predictive distribution is the posterior predictive $p$-value:

$$p_{pp,j} = P \left[ T_j(y_{rep}, \eta) \geq T_j(y, \eta) | y \right] = \int P \left[ T_j(y_{rep}, \eta) > T_j(y, \eta) | y, \eta \right] p(\eta | y) d\eta \quad (11)$$

Since this quantity has no closed form, the idea is to approximate it by the Monte Carlo method. For each estimated model (ODE and SDE), the Gibbs algorithm used to estimate the posterior distribution provides a set of vectors $\eta^l (l = 1 \ldots L)$ drawn from the posterior distribution $p(\eta | y)$. For each of this draw, a replicated data set $y_{rep}^l$ is simulated from the posterior predictive distribution of the data $p(y_{rep}^l | y)$. Finally, the posterior predictive $p$-value (11) is estimated by the Monte Carlo method as

$$\frac{1}{L} \sum_{l=1}^L 1_{T_j(y_{rep}^l, \eta^l) > T_j(y, \eta^l)}.$$ 

5. An example: chicken growth modeling with the Gompertz function

We focus on the modeling of chicken growth. Data $y$ are noisy weight measurements of $n = 50$ chickens at weeks $t = 0, 4, 6, 8, 12, 16, 20, 24, 28, 32, 36, 40$ after birth: see the
corresponding curves on Figure 1. Such a data set has been previously analyzed by Mignon-Grasteau et al. (1999), Jaffrézic et al. (2006) and Meza et al. (2007), who concluded that, among the standard growth models, the monotonic mixed Gompertz model is the most appropriate one. This model is adapted to the most subjects, however it fails to model the unexpected variations of growth rate for some individuals (see Figure 1).

5.1 The classical Gompertz nonlinear mixed model

Jaffrézic and Foulley (2006) underline that a heteroscedastic error model is required to obtain satisfactory results. For simplicity’s sake, we consider modeling the logarithm of the data \( y \) by adding an additive measurement error with a constant variance:

\[
\begin{align*}
\log y_{ij} &= \log A_i - B_i e^{-C_i t_{ij}} + \varepsilon_{ij}, \varepsilon_{ij} \sim \text{i.i.d.} N(0, \sigma^2), \forall i = 1, \ldots, n, j = 0, \ldots, n_i \\
\phi_i &= (\log A_i, B_i, \log C_i) \sim \text{i.i.d.} N(\mu, \Omega), \forall i = 1, \ldots, n
\end{align*}
\]

We use the log-parametrization for parameters \( A_i \) and \( C_i \). This parametrization has two advantages: it simplifies the computation of the posterior distributions and it ensures the positivity of the parameters. We set \( \mu = (\log(a), b, \log(c)) \).

5.2 Extension to the Gompertz stochastic nonlinear mixed model

We now deduce the SDE model from the Gompertz equation (3). Given the heteroscedasticity of the process, the volatility function is set to be equal to \( \Gamma(Z_t, \phi, \gamma^2) = \gamma Z_t \):

\[
dZ_t = B C e^{-C t} Z_t dt + \gamma Z_t dW_t, \quad Z_0 = A e^{-B}
\]

We use the log-parametrization for parameters \( A_i \) and \( C_i \). This parametrization has two advantages: it simplifies the computation of the posterior distributions and it ensures the positivity of the parameters. We set \( \mu = (\log(a), b, \log(c)) \).

This means that the standard error of the random perturbations of the growth rate is proportional to weight. This choice of volatility has two main advantages. First, SDE (13) has an explicit solution. Indeed, set \( X_t = \log(Z_t) \). By the Itô’s formula, for \( h > 0 \), the conditional distribution of \( X_{t+h} \) given \( (X_s)_{s \leq t} \) is:

\[
X_{t+h} | (X_s)_{s \leq t} \sim N(X_t - Be^{-Ct}(e^{-Ch} - 1) - \frac{1}{2} \gamma^2 h, \gamma^2 h), \quad X_0 = \log(A) - B
\]
Thus, ∀t > 0, we have $Z_t = A e^{-B e^{-C_i t} e^{-\frac{1}{2} \gamma^2 t + \eta_t}} = f(t) e^{-\frac{1}{2} \gamma^2 t + \eta_t}$ with $\eta_t \sim \mathcal{N}(0, \gamma^2 t)$ and $Z_0 = A e^{-B}$. As a consequence, $Z_t$ is a multiplicative random perturbation of the solution of the Gompertz model. Second, due to the assumption of the non-negativity of $A$, $Z_t$ is almost surely non-negative, which is a natural constraint to model weight records.

We then discretize the SDE. The discrete realization ($X_{t,i}$) of the SDE is Markovian:

$$X_{t,i} | X_{t,i-1} \sim \mathcal{N} \left( X_{t,i-1} - B_i e^{-C_i t_i} (e^{-C_i (t_i - t_{i-1})} - 1) - \frac{1}{2} \gamma^2 (t_i - t_{i-1}), \gamma^2 (t_i - t_{i-1}) \right)$$

with $X_{t,0} = \log(A_i) - B_i$. The SDE model (8) on the logarithm of data is thus defined as:

$$\begin{align*}
\left( \log y_{i0}, \log y_{i1}, \ldots, \log y_{in} \right)' &= \left( \log(A_i) - B_i, X_{t,i1}, \ldots, X_{t,im} \right)' + \varepsilon_i, \quad \varepsilon_i \sim \text{i.i.d. } \mathcal{N}(0, \sigma^2 I_{n+1}) \\
\left( X_{t,i1}, \ldots, X_{t,im} \right)' &= \left( \log(A_i) - B_i e^{-C_i t_i}, \ldots, \log(A_i) - B_i e^{-C_i t_m} \right)' - \gamma^2 (t_i, \ldots, t_m)' + \eta_i \\
\eta_i &\sim \text{i.i.d. } \mathcal{N}(0, \gamma^2 T_i), \quad T_i = (\min(t_{ij}, t_{ij}'))_{1 \leq j,j' \leq n} \\
(\log A_i, B_i, \log C_i) &\sim \text{i.i.d. } \mathcal{N}(\mu, \Omega)
\end{align*}$$

(14)

5.3 Posterior computation and inference in the Gompertz model

Conditional distribution computation for the ODE mixed model is standard. We detail the computation under the SDE mixed model. Let $m_a^{\text{prior}}, m_b^{\text{prior}}, m_c^{\text{prior}}, \gamma_a^{\text{prior}}, \gamma_b^{\text{prior}}, \gamma_c^{\text{prior}}$ denote the prior parameters of the 3 components of $\mu$. The conditional distribution of $X_i = (X_{ij})_{1 \leq j \leq n_i}$ given $(\phi_i, \gamma^{-2}, y_i, \sigma^2)$ is Gaussian with mean $m_i^{\text{post}}$ and variance $V_i^{\text{post}}$:

$$V_i^{\text{post}} = (\sigma^2 I_{n_i-1} + \gamma^2 T_i^{-1})^{-1}, \quad m_i^{\text{post}} = V_i^{\text{post}} \left[ \sigma^2 (\log y_{i1} \ldots \log y_{in})' + \gamma^2 T_i^{-1} u_{X_i} \right]$$

with $u_{X_i} = \log A_i - B_i \left( e^{-C_i t_{i1}} \ldots e^{-C_i t_{in}} \right)' - \frac{1}{2} \gamma^2 (t_{i1} \ldots t_{in})'$. Let $\omega_{\log A}, \omega_B, \omega_{\log C}$ denote the diagonal elements of $\Omega$. Let $\Omega_{k,(j,j')}$ denote the two-vector composed of the elements on the $k$-th row and $(j, j')$ columns of $\Omega$ and $\Omega_{(j,j'),(j,j')}$ the two-symmetric-matrix composed of the elements on the $(j, j')$-th rows and $(j, j')$-th columns of $\Omega$. Set the $(n_i + 1) \times (n_i + 1)$-matrix:

$$G_i = \begin{pmatrix}
\sigma^2 & 0 \\
0 & \gamma^2 T_i
\end{pmatrix}$$

(15)
The conditional distributions for the individual parameters \(\log A_i\) and \(B_i\) are \(\mathcal{N}(m_{A_i}^{post}, V_{A_i}^{post})\) and \(\mathcal{N}(m_{B_i}^{post}, V_{B_i}^{post})\), respectively, with:

\[
\begin{align*}
V_{A_i}^{post} &= \left[ (1 \ldots 1)G_i^{-1}(1 \ldots 1)' + \frac{1}{\omega_{\log A | B, \log C}^2} \right]^{-1} \\
m_{A_i}^{post} &= V_{A_i}^{post} \left[ (1 \ldots 1)G_i^{-1}u_{A_i} + \frac{\mu_{\log A | B, \log C}}{\omega_{\log A | B, \log C}^2} \right] \\
V_{B_i}^{post} &= \left[ (e^{-C_i t_0} \ldots e^{-C_i t_{m_i}})G_i^{-1}(e^{-C_i t_0} \ldots e^{-C_i t_{m_i}})' + \frac{1}{\omega_B^2 | \log A, \log C} \right]^{-1} \\
m_{B_i}^{post} &= V_{B_i}^{post} \left[ (e^{-C_i t_0} \ldots e^{-C_i t_{m_i}})G_i^{-1}u_{B_i} + \frac{\mu_B | \log A, \log C}{\omega_B^2 | \log A, \log C} \right]
\end{align*}
\]

where

\[
\begin{align*}
&u_{A_i} = (\log y_{i0}, X_{i1} \ldots X_{i m_i})' + B_i (e^{-C_i t_0} \ldots e^{-C_i t_{m_i}})' - \frac{1}{2} \gamma^2 (t_{i0} \ldots t_{im_i})' \\
&\omega_{\log A | B, \log C}^2 = \omega_{\log A}^2 - \Omega_{\log A, (B, \log C)}^2 \omega_{(B, \log C), (B, \log C)}^{-1} \omega_{(B, \log C), (B, \log C)} \omega_{B, (B, \log C)} \\
&\mu_{\log A | B, \log C} = \log a + \Omega_{\log A, (B, \log C)}^{-1} \Omega_{(B, \log C), (B, \log C)} ( (B_i, \log C_i)' - (b, \log c)' ) \\
&u_{B_i} = (\log y_{i0}, X_{i1} \ldots X_{i m_i})' + \log A_i - \frac{1}{2} \gamma^2 (t_{i0} \ldots t_{im_i}) \\
&\omega_B^2 | \log A, \log C = \omega_B^2 - \Omega_B \omega_{(B, \log A, \log C)}^{-1} \omega_{(B, \log A, \log C), (B, \log A, \log C)} \omega_B \omega_{(B, \log A, \log C)} \\
&\mu_B | \log A, \log C = b + \Omega_B \omega_{(B, \log A, \log C)}^{-1} \omega_{(B, \log A, \log C)} ( (\log A_i, \log C_i)' - (\log a, \log c)' )
\end{align*}
\]

The conditional distributions of \(\log(a)\) and \(b\) are Gaussian with parameters:

\[
\begin{align*}
V_a^{post} &= \left[ n \omega_{\log A}^{-2} + (v_a^{prior})^{-1} \right]^{-1} \quad \text{and} \quad m_a^{post} = V_a^{post} \left[ \omega_{\log A}^{-2} \frac{1}{n} \sum_{i=1}^n \log A_i + m_a^{prior} \right] \\
V_b^{post} &= \left[ n \omega_B^{-2} + (v_b^{prior})^{-1} \right]^{-1} \quad \text{and} \quad m_b^{post} = V_b^{post} \left[ \omega_B^{-2} \frac{1}{n} \sum_{i=1}^n B_i + m_b^{prior} \right]
\end{align*}
\]

The conditional distribution of \(\Omega^{-1}\) is \(W(R + (\phi - \mu)(\phi - \mu)', n + p + 1)\) where \(\phi - \mu = [(\phi_1 - \mu) \ldots (\phi_n - \mu)] \in \mathbb{R}^{3 \times n}\). The conditional distribution of \(\sigma^2\) is \(\Gamma(\alpha^{post}_{\sigma}, \beta^{post}_{\sigma})\) with:

\[
\alpha^{post}_{\sigma} = \alpha^{prior}_{\sigma} + \sum_{i=1}^n \frac{n_i}{2} + 1 \quad \text{and} \quad \beta^{post}_{\sigma} = \left[ \frac{1}{\beta^{prior}_{\sigma}} + \frac{1}{2} \sum_{i=1,j=0}^{n,n} (\log y_{ij} - X_{ij})^2 \right]^{-1}
\]

The posterior distributions of \(\log C_i\), \(\log c\) and \(\gamma^2\) have no explicit form and we use the Metropolis-Hastings random-walks.
The Gibbs algorithm convergence is ensured by the classical convergence theorem proposed by Carlin and Louis (2000), the convergence of the Metropolis-Hastings algorithm is ensured by the theorem proposed by Mengersen and Tweedie (1996). The implementations of the ODE and SDE approaches have the same level of complexity. The convergence and the stability of the MCMC algorithms produced by these two algorithms are equivalent.

5.4 Simulations

We simulate datasets mimicking chicken growth with \( n = 50 \) individuals and \( n_i = 9 \) measurements obtained every 5 weeks after birth. The population parameters are \( \log(a) = \log(3000) \), \( b = 5 \), \( \log(c) = \log(14) \), \( \Omega \) is assumed diagonal with diagonal elements equal to 100 and \( \sigma^{-2} = 5 \). A 100 datasets are simulated via the mixed model defined by the Gompertz model (12) and a 100 datasets with the mixed model defined by the Gompertz SDE (14), with \( \gamma^2 = 1 \). We estimate all the parameters under the ODE mixed model (12) and the SDE mixed model (14), successively. The two algorithms take 96s and 206s on a dataset with a Intel Core2 Duo CPU (2.4 GHz), respectively. Estimates are obtained as the expectation of the parameter posterior distribution. Means and standard errors computed for each parameter on the 100 estimation results obtained with both algorithms are presented in Table 1.

[Table 1 about here.]

When data are simulated under the ODE model, estimates obtained with the Bayesian ODE algorithm are very satisfactory. Those obtained by the Bayesian SDE algorithm are also satisfactory although the bias for the variance parameter \( \omega_{lnA}^{-2} \) is larger. Note that, as expected, the estimation of the volatility parameter \( \gamma^2 \) is rather low (0.19). When data are simulated under the SDE model, estimates obtained with the Bayesian SDE model are very satisfactory, with small bias and standard error. On the contrary, the estimates obtained with the Bayesian ODE algorithm have larger bias, including the parameters of fixed effects. The parameter \( \omega_{logA}^{-2} \) is very badly estimated (8.63 to be compared to the true value 100).
5.5 Application on chicken growth data

In this section, we apply the proposed models on real data of chicken growth. The ODE and SDE models (12) and (14) are used to model the logarithm of the data. The influence of the prior choices of the ODE model has been validated in Jaffrézic et al. (2006). For the SDE model, the influence of the prior choice for the parameter $\gamma^2$ is studied using the Deviance Information Criterion (DIC), which combines two terms, one for "goodness of fit" and the other for penalizing the complexity of the model (Spiegelhalter et al. (2002)). Only differences in DIC are meaningful. A inverse Gamma prior on $\gamma^2$, a Jeffreys prior on $\gamma^{-2}$, a log normal prior on $\log(\gamma)$, a uniform prior on $\gamma^2$ have been tested leading to DIC variations from the uniform prior choice equal to 24, 1.5, 1.4 and 0 respectively. In addition, the influence on the posterior distributions was very light. Results are presented with a uniform prior on $\gamma^2$. Posterior expectations of the parameters are presented in Table 2. Diagnostic tools to validate the models are applied to both ODE and SDE models: Figure 2 presents the posterior predictive distributions and p-values of both models computed for each time point. The estimate of the volatility parameter $\gamma^2$ is strictly positive and its credibility interval does not include 0: this means that the dynamical process that most likely represents the growth is a stochastic process with non-negligible noise. Furthermore, the diagnostic tools show a clear improvement from the ODE model to SDE model for the whole population, both at early and late ages. Here, the reduction in DIC from the ODE to the SDE models is equal to 393, which clearly indicates the better predictive ability of the SDE model. Figure 3 reports, for four subjects, the observed weights, the ODE prediction, the empirical mean of the last 1000 simulated trajectories of the SDE (14) generated during the Gibbs algorithm, their empirical 95% confidence limits (from the 2.5th percentile to the 97.5th percentile) and one simulated trajectory. Subjects 4 and 13 are examples of subjects with no growth slow down. Both ODE and SDE models satisfactorily fit the observations. Subject 14 has a
small observed weight decrease. For subject 1, the weight decrease is more important. For both subjects, the ODE model fails to capture this phenomenon while the SDE model does. Furthermore, the SDE model provides different estimates for the individual parameters. For example for subject 1, the individual parameter $A_1$ (adult weight) is estimated at 3.922 kg and 3.484 kg by the ODE and SDE models, respectively.

6. Conclusion and discussion

We propose a Bayesian approach to nonlinear mixed models defined by stochastic differential equations. These models are an alternative to classical nonlinear mixed models whose deterministic regression function is too restrictive to model some unexplained biological processes such as growth rate changes. On the presented data set, the introduction of this SDE model leads to a clear validation of the model (Figure 2) which was not the case in the standard model, justifying the introduction of the new stochastic component. Moreover, the modelling of these fluctuations has a non-neglectible impact on the estimation of the parameters. This might have a great influence on the conclusions about genetic specifications.

We consider SDEs defined with a general volatility function. As discussed in Section 2, the study context can induce a natural choice of this volatility function (non-negativity of the process, fluctuation of a parameter, heteroscedastic variances,...). This volatility function choice has obviously some consequences on the complexity of the Bayesian posterior computation. As detailed in Section 3, a volatility function which induces an explicit solution of the SDE will imply a very easy implementation of the algorithm. The example detailed in
this work belongs to this case. On the contrary, when the SDE has no explicit distribution, we propose to use the Euler-Maruyama scheme to approximate the diffusion: the conditional distribution is then Gaussian, implying an easy Bayesian implementation. We control the error induced by this Euler approximate scheme on the posterior distribution. In this context, auxiliary latent points are introduced to obtain a better approximation of the diffusion. The choice of the discrete grids \((\tau_0, \ldots, \tau_{M_t})\) is complex and has been evoked by Pedersen (1995) and Donnet and Samson (2008). In conclusion, generally speaking, there is no limitations to the choice of the volatility function of the SDE approach.

Our model differs from mixed models with continuous time autoregressive measurement errors, as proposed by De la Cruz-Mesia and Marshall (2006) or others. These authors assume that measurement errors have an auto-regressive structure \((Y_{t_i} = f(t_{ij}, \phi_i) + \varepsilon_{t_{ij}} \text{ and } d\varepsilon_t = -a\varepsilon_t dt + \sigma dW_t)\). We assume that the auto-regressive structure observed in residuals of classical nonlinear mixed models comes from a model failure: the regression function is too restrictive and rigid to model random variations of the biological process. Therefore, in our model, it is the regression process that has an auto-regressive structure, while the observation measurements are assumed to be independant. These two models are different. De la Cruz-Mesia and Marshall (2006) consider a stationnary CAR process, implying that the process of the observations has a homoscedastic variance. On the contrary, we do not assume any stationnarity for the process \((Z(t))\). Moreover, heteroscedastic error model can be considered with our approach, depending on the choice of the volatility function.

The proposed model should prove to be useful for other applications in which deterministic models are too restrictive to take into account different sources of variation that exist in real life. For example, Picchini et al. (2006) propose a stochastic differential equation to model glucose/insulin dynamics, where sources of variability are various (anxiety, rest, etc). The extension of this work to mixed models using our approach should be of great interest.
An interesting area for future research is the development of model selection tools in this context. Indeed, the analysis of covariate effects and the comparison between the ODE and the SDE models require specific selection tools. Indeed, the method of pseudo-priors proposed by Carlin and Chib (1995) and developed by others, which is very sensitive to the choice of priors and pseudo-priors, would be difficult to use in practice in our context. Bayes factors are complex to compute in these models but could be an interesting alternative. Finally, the extension of this work to multidimensional SDEs would be of great interest in several biological applications.

7. Supplementary Materials

Web Appendix referenced in Section 3 is available under the Paper Information link at the Biometrics website http://www.biometrics.tibs.org.

Acknowledgements

The authors are grateful to Dr F. Jaffrézic for helpful discussions on mixed model and to Mrs Brand Williams and Vincent-Arnaud for their English revision.

References


*Received. Revised.*

*Accepted.*
Figure 1. Growth curves of the 50 chickens and mean growth curve in dashed bold line.
Figure 2. Posterior predictive distributions for the ODE and SDE models on chicken growth data. Posterior predictive p-values at times (0, 4, 6, 8, 12, 16, 20, 24, 28, 32, 36, 40) for the ODE (resp. SDE) mixed models are equal to (0.55, 0.00, 0.86, 0.80, 0.08, 0.48, 0.90, 0.73, 0.02, 0.99, 0.91, 0.46) (resp (0.49, 0.23, 0.40, 0.45, 0.43, 0.60, 0.64, 0.60, 0.43, 0.57, 0.64, 0.56)).
Figure 3. Observations (circles), predictions obtained with the ODE mixed model (long dashed line), mean SDE prediction (smooth solid line), 95% credibility interval obtained with the SDE mixed model (dotted line) and one SDE realization (solid line), for subjects 1, 4, 13 and 14.
Mean estimates and standard errors in brackets obtained from the ODE and the SDE mixed models on 100 datasets simulated with the ODE or the SDE mixed model.

<table>
<thead>
<tr>
<th>Simulation model</th>
<th>true</th>
<th>ODE ($\gamma^2 = 0$)</th>
<th>SDE ($\gamma^2 = 1$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Estimation model</td>
<td>value</td>
<td>ODE</td>
<td>SDE</td>
</tr>
<tr>
<td>$\mu_{\ln A}$</td>
<td>8.01</td>
<td>8.00 (0.04)</td>
<td>8.03 (0.06)</td>
</tr>
<tr>
<td>$\mu_B$</td>
<td>5.00</td>
<td>4.99 (0.08)</td>
<td>5.02 (0.08)</td>
</tr>
<tr>
<td>$\mu_{\ln C}$</td>
<td>2.64</td>
<td>2.64 (0.04)</td>
<td>2.63 (0.04)</td>
</tr>
<tr>
<td>$\omega_{\ln A}^2$</td>
<td>100.00</td>
<td>122.24 (39.95)</td>
<td>160.84 (27.84)</td>
</tr>
<tr>
<td>$\omega_B^2$</td>
<td>100.00</td>
<td>106.70 (22.17)</td>
<td>103.16 (23.74)</td>
</tr>
<tr>
<td>$\omega_{\ln C}^2$</td>
<td>100.00</td>
<td>126.27 (45.69)</td>
<td>131.03 (55.02)</td>
</tr>
<tr>
<td>$\gamma^2$</td>
<td>- (-)</td>
<td>0.19 (0.02)</td>
<td>- (-)</td>
</tr>
<tr>
<td>$\sigma^{-2}$</td>
<td>5.00</td>
<td>5.05 (0.39)</td>
<td>5.35 (0.43)</td>
</tr>
</tbody>
</table>
### Table 2

*Posterior distributions for the ODE and SDE models on chicken growth data: mean estimated parameters and their 95% credibility intervals (95% CI).*

<table>
<thead>
<tr>
<th></th>
<th>ODE</th>
<th>SDE</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>mean</td>
<td>95% CI</td>
</tr>
<tr>
<td>log $a$</td>
<td>7.77</td>
<td>[7.70; 7.84]</td>
</tr>
<tr>
<td>$b$</td>
<td>4.17</td>
<td>[4.11; 4.23]</td>
</tr>
<tr>
<td>log $c$</td>
<td>2.75</td>
<td>[2.70; 2.81]</td>
</tr>
<tr>
<td>$\Omega^{-1}_{\log a,\log a}$</td>
<td>117.30</td>
<td>[66.53; 190.90]</td>
</tr>
<tr>
<td>$\Omega^{-1}_{\log a,b}$</td>
<td>-128.50</td>
<td>[-217.90; -68.32]</td>
</tr>
<tr>
<td>$\Omega^{-1}_{\log a,\log c}$</td>
<td>-4.57</td>
<td>[-29.53; 16.81]</td>
</tr>
<tr>
<td>$\Omega^{-1}_{b,b}$</td>
<td>172.10</td>
<td>[94.21; 287.90]</td>
</tr>
<tr>
<td>$\Omega^{-1}_{b,\log c}$</td>
<td>22.64</td>
<td>[-4.73; 57.69]</td>
</tr>
<tr>
<td>$\Omega^{-1}_{\log c,\log c}$</td>
<td>36.68</td>
<td>[23.04; 54.99]</td>
</tr>
<tr>
<td>$\sigma^{-2}$</td>
<td>225.5</td>
<td>[197.40; 255.50]</td>
</tr>
<tr>
<td>$\gamma^2$</td>
<td>0.09</td>
<td>[0.07; 0.12]</td>
</tr>
</tbody>
</table>