

Comparison of Multi-EKF and MHE for PD parameter Estimation

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1 Introduction

This report details the pipeline for the comparison of the two methods to identify PD parameter from Propofol and Remifentanil using BIS as the measured signal. In order to have a fair comparison, this is done on simulated data, where the true parameters are known.

This report is organized as follow, Section 2 explain the method to generate the simulated data. Section 3 explain the details of the two proposed methods and the metrics used to evaluate the performances. The results are presented in section 4 and some conclusion are exposed in section 5.

2 Data generation

2.1 Drug model

In order to simulate the effect of Propofol and Remifentanil on the depth of hypnosis the standard pharmacokinetic-pharmacodynamic (PK-PD) structure is used. The PK part models the distribution of drugs in the body using a compartment-model resulting in two decoupled LTI system, one for each drug. The outputs of this models are two effect-site drug concentrations which can be linked to the hypnotic effect by the PD part using a surface model response to model the synergic effect between both drugs. The final, model can be expressed by the following equation:

$$\begin{aligned}\dot{x}(t) &= Ax(t) + Bu(t) \\ y(t) &= f(x(t)) + w(t)\end{aligned}\tag{1}$$

where $x(t) \in \mathbb{R}^8$ represent the concentration of drugs inside the different compartments, particularly $x_4(t)$ and $x_8(t)$ are respectively the Propofol and Remifentanil effect-site concentration. $u(t) = (u_p(t) \ u_r(t))^{\top}$ are the Propofol and Remifentanil drug rates injected into the patient blood. f is the surface model function that can be describe by the following equation:

$$f(x(t)) = BIS_0 + E_{max} \frac{I(t)^{\gamma}}{1 + I(t)^{\gamma}}\tag{2}$$

where BIS_0 is the initial BIS, E_{max} the maximum effect of combined drugs, γ the slope coefficient of the Hill curve and $U(t)$ the interaction term defined by:

$$I(t) = \frac{x_4(t)}{C_{50p}} + \frac{x_8(t)}{C_{50r}}; \quad (3)$$

C_{50p} and C_{50r} are the Propofol and Remifentanil half-effect concentrations for BIS (*i.e.* the concentrations to achieve half the effect of the drugs).

Finally $w(t)$ is the measurement noise. In the paper, a white noise filtered by a second order low pass filter with a cut-off frequency of 0.03 Hz is used to replicate real condition.

In the simulation, the parameters of [1] and [2] are used respectively for Propofol and Remifentanil PK model. For the PD model the parameters from [3] are implemented. To perform simulation as close to the reality as possible uncertainties are added to the parameters. Particularly, each parameter is following a log-normal distribution (parameters of the distribution are specified in the previously cited paper) and a realization of the distribution is used for each patient. Simulation are done using the Python Anesthesia Simulator [?].

2.2 Drug rates tunning

During the total intravenous anesthesia, Propofol and Remifentanil are dosed using Target Concentration Infusion (TCI) pumps. Those pumps include the nominal PK models, the anesthetist can select a concentration target and the pumps will compute and inject the drug to reach the target thanks to the model in an open loop matter. At the end the anesthetist is tuning the concentration target to reach the desired level of hypnosis.

In order to recreate this scheme of control, TCI control algorithm [4] have been reproduced. To reproduce the comportment of the anesthesiologist the following rules have been coded:

- At the beginning the target concentration are set to $u_p = 4\mu\text{g}/\text{mL}$ and $u_r = 4\text{ng}/\text{mL}$.
- A control period is randomly draw for each patient and each drugs between 3 and 7 minutes. At each end of the period, the concentration target of the given drugs is updated as follow:
 - If $55 < BIS$ the target is increased by 0.5.
 - If $40 < BIS$ the target is decreased by 0.5.

In addition for Propofol only:

- If $BIS > 60$ the target is increased by 1.
- If $BIS < 40$ the target is decreased by 1.

An example of the resulting control is exposed Fig 1.

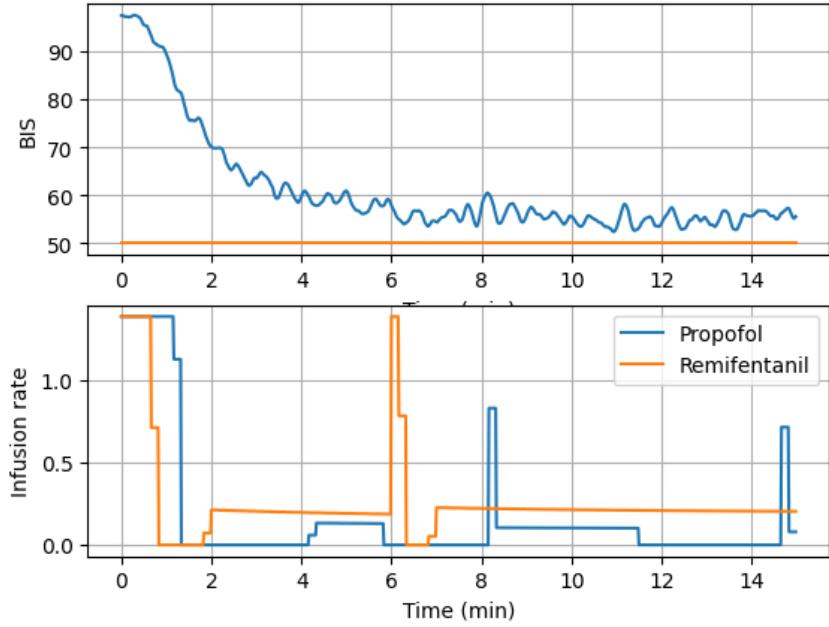


Figure 1: Results of the control algorithm for a single random patient.

2.3 Characteristics of the database

The final databased includes induction (15 minutes) simulation files for 1000 different patients with a sampling time of one second. Patients characteristic have been randomly chosen using uniform distribution ($\text{age} \in [18, 70]$, $\text{height} \in [150, 190]$, $\text{weight} \in [50, 100]$, and gender $\in \{0, 1\}$). Each files includes all the signals from the simulator, particularly the BIS values and the drug rates over time are available. An additional file includes the parameters of each patient. Figures 2-4 expose the results of the simulations.

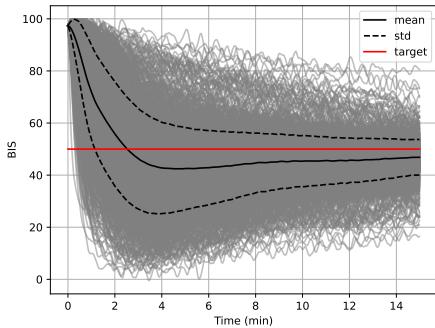


Figure 2: BIS trajectory for all the simu-

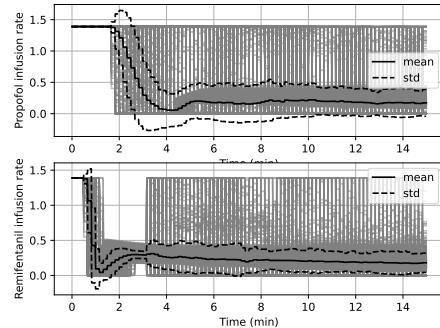


Figure 3: Drug input rates for all the simulations.

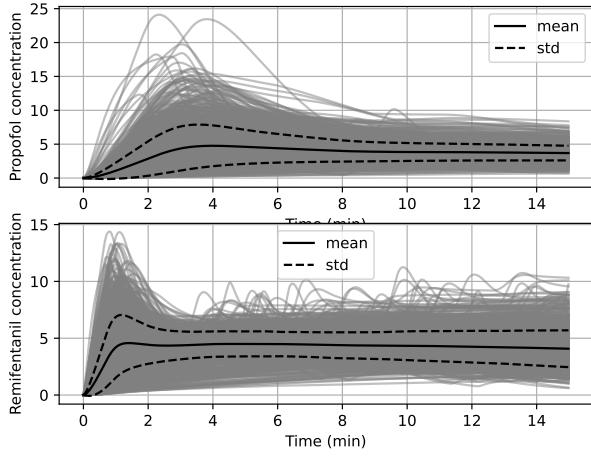


Figure 4: Drug effect-site concentration for all the simulations.

3 Methods

In this section, different methods used to estimate the unknown parameters of the PD models are detailed. BIS_0 can be measured at before the induction of anesthesia and E_{max} is usually set equal to BIS_0 . Thus the remaining parameters are C_{50p} , C_{50r} , and γ . in this section, $\theta = (C_{50p} \ C_{50r} \ \gamma)$ is used to describe the vector of unknown parameters.

The Multi Extended Kalman Filter (MEKF) method select the best vector among a grid in space of the parameters. This discrete choice allow a fast convergence but less precision at the end. The Moving Horizon Estimation (MHE) method uses an extended state formulation to estimate the vector of parameters along with the state in a continuous manner. Thus the method could identify more precisely the parameters but is also more subject to noise and could be slower than MEKF.

3.1 Multi Extended Kalman Filter

In order to identify the PD parameters, the MEKF method uses a set of EKF, one for every realization of the vector selected within a grid in the space of the parameters. The grid is designed to reasonably represent the variability of the parameter vector. Next, a vector is chosen using a model-matching criterion.

EKF is a state estimation method that relies on the linearization of a non-linear model. If we consider the model given in (1) with the non-linear function f parametrized by θ , the estimator using the parameter vector θ_i is given by:

$$\begin{aligned}
H_i(k) &= \frac{\partial f(x, \theta_i)}{\partial x} \Big|_{x=\hat{x}_i(k_{|k-1})} \\
K_i(k) &= P_i(k_{|k-1}) H_i^\top(k) (H_i(k) P_i(k_{|k-1}) H_i^\top(k) + R_2)^{-1} \\
\hat{x}_i(k_{|k}) &= \hat{x}_i(k_{|k-1}) + K_i(k)(y(k) - f(\hat{x}_i(k_{|k-1}), \theta_i)) \\
P_i(k_{|k}) &= P_i(k_{|k-1}) - K_i(k) H_i(k) P_i(k_{|k-1}) \\
\hat{x}_i(k+1_{|k}) &= A\hat{x}_i(k_{|k}) + Bu(k) \\
P_i(k+1_{|k}) &= AP_i(k_{|k})A^\top + R_1
\end{aligned}$$

Here the notation $X(k_{1|k_2})$ represents the value of variable X computed at time step k_1 based on the knowledge available at k_2 . The estimated state vector is \hat{x} and P is the covariance matrix. R_1 and R_2 are two constant matrices used to respectively characterize the process uncertainties and the measurements noise.

3.2 Moving Horizon Estimation

3.3 Metrics for the comparison

4 Results

5 conclusion

References

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