Supplementary material: Errors-in-variables Modeling of Personalized Treatment-Response Trajectories

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I. Specification of model prior distributions

Response function:

$$\tilde{\beta}_l \sim Normal(mu = 0, sd = 5)$$
 (1)

$$\tilde{\beta}_h \sim Normal(mu = 0, sd = 5)$$
 (2)

$$\Sigma_l \sim HalfNormal(sd = 1) \tag{3}$$

$$\Sigma_h \sim HalfNormal(sd = 0.5)$$
 (4)

$$\beta_n^l \sim Normal(mu = \tilde{\beta}_l, sd = \Sigma_l)$$
 (5)

$$\beta_n^h \sim Normal(mu = \tilde{\beta}_h, sd = \Sigma_h)$$
 (6)

where the dimension of covariates P is equal to 1 for notation simplicity. For P > 1, an independence among different dimensions is assumed.

Counterfactual trend:

$$ls_n \sim HalfNormal(sd = 10) + 10 \tag{7}$$

$$nu_n \sim HalfNormal(sd = 10)$$
 (8)

$$c_n \sim HalfNormal(sd = 10)$$
 (9)

$$K1_n = nu_n * ExpQuad(\cdot|ls_n) \tag{10}$$

$$K2_n = Constant(\cdot|c_n) \tag{11}$$

$$K_n = K1_n + K2_n \tag{12}$$

where $ExpQuad(\cdot)$ and $Constant(\cdot)$ are kernel functions. Time is measured in minutes, and the length-scale ls_n is bounded from below by 10 minutes to increase stability of the estimation (measurements are taken at an interval of 15 minutes in our real-world data set).

Mearurement models:

$$d_n \sim Normal(mu = 0, sd = 10) \tag{13}$$

$$\sigma_n^t \sim HalfNormal(sd = 10) \tag{14}$$

$$t_{nm}^* \sim d_n + \sigma_n^t * Normal(mu = 0, sd = 1)$$
(15)

$$\delta_{nm} \sim Lognormal(mu = 0, sd = 0.05) \tag{16}$$

II. DETAILS OF THE FIRST SIMULATION EXPERIMENT

In total 300 markers consisting of a zero trend and 5 evenly distributed treatments are generated, where 2-dimension input is generated randomly from Uniform(0.3, 1). A perturbation rate sampled from $N(1, 0.2^2)$ are multiplied by each input. The

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groundtruth coefficients for the height of treatment response are [5, 3] and [15, 2] for length-scale. The result is shown in Figure 1.

III. DERIVATION FOR THE MARGINAL INCREMENT OF TREATMENT RESPONSE AREA BY ONE COVARIATE

For simplicity, we focus on a single individual and drop the unnecessary indexing in the notation. The area A is proportional to length-scale l and height h of the response. Hence

$$A = \lambda h l \tag{17}$$

for some constant λ (knowing the shape of the response, solving for λ analytically is straightforward). Denote the amount of one covarate, e.g. sugar, in the *m*th meal by x_{mi} where $i \in$ {1, 2, ..., *P*}. Now the length-scale *l* depends on *x* through

$$l_m(x_{mi}) = g(y_m^l) = g(\beta_i^l x_{mi} + c_m^l),$$
(18)

where g is the *softplus* function and c_m^l comprises the other parts of the linear predictor that do not depend on the sugar x_{mi} . Similarly, the height h depends on x through

$$h_m(x_{mi}) = g(y_m^l) = g(\beta_i^h x_{mi} + c_m^h).$$
(19)

We want to know how area A_m changes if we change the amount of sugar x_{mi} by one unit.

$$\frac{dA_m}{dx_{mi}} = \lambda \frac{dl_m}{dx_{mi}} h_m + \lambda l_m \frac{dh_m}{dx_{mi}}$$
(20)

$$= \lambda \frac{dl_m}{dy_m^l} \frac{dy_m^l}{dx_{mi}} h_m + \lambda \frac{dh_m}{dy_m^h} \frac{dy_m^h}{dx_{mi}} l_m$$
(21)

$$= \lambda (1 + e^{-y_m^l})^{-1} \beta_i^l h_m + \lambda (1 + e^{-y_m^h})^{-1} \beta_i^h l_m.$$
(22)

By replacing x_m with an average meal, we have

$$\frac{dA}{dx_{i}} = \lambda (1 + e^{-\bar{y^{i}}})^{-1} \beta_{i}^{l} \bar{h} + \lambda (1 + e^{-\bar{y^{h}}})^{-1} \beta_{i}^{h} \bar{l}$$
(23)

$$= \lambda (1 + e^{-(\beta^{l})^{T} \bar{\mathbf{x}}})^{-1} \beta_{l}^{l} \bar{h} + \lambda (1 + e^{-(\beta^{h})^{T} \bar{\mathbf{x}}})^{-1} \beta_{l}^{h} \bar{l}.$$
(24)

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Fig. 1: EIV results in the first simulation experiment. The left plot shows how the perturbations are restored; the middle compares the coefficients estimated with and without EIV; the last one exhibits overall fittings of EIV and baseline models. Red stars and dots stand for groundtruth values and observed points respectively.



Fig. 2: Results of the simulation experiment in Section IV-A of the main text for the second patient.



Fig. 3: Replication of the results of the simulation experiment in Section IV-A of the main text, initialized with a different seed. Results for both patients are shown.



Fig. 4: Posterior uncertainty in the personalized regression coefficients, estimated with different models from the real-world data. \mathcal{M}_{ind} model is excluded because of its much wider and distracting uncertainty.



Fig. 5: Histogram of personalized coefficients for different covariates, estimated with the real-world data. Red line and the surrounding region coloured red show mean +/- one SD.



Fig. 6: Visualization of the observed and estimated trajectories for all patients.