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Non-fragile state estimation for reaction-diffusion genetic regulatory networks with mode-dependent timevarying delays

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Abstract

This paper investigates the problem of non-fragile state estimation for a class of reaction-diffusion genetic regulatory networks with mode-dependent time-varying delays and Markovian jump parameters. First, the Markov chain with partially unknown probabilities is used in this paper to describe the switching between system modes, which can make the model more generalizable. Moreover, considering the possible gain variations, we design a non-fragile state estimator that makes the estimation performance non-fragile to gain variations, thus guaranteeing the estimation performance. Sufficient conditions that ensure the asymptotic stability of the estimation error can be derived by using the Lyapunov stabilization theory and several inequality treatments. Finally, a simulation example is presented to demonstrate the effectiveness of the proposed estimator design scheme.

Keywords: Non-fragile state estimation, reaction-diffusion genetic regulatory networks, mode-dependent time-varying delays, partially unknown probabilities

1. INTRODUCTION

Recently, genetic regulatory networks (GRNs) have gained significant interest in the fields of biology, mathematics, and medicine [1-3]. This is because mathematical models of GRNs can describe the complex dynamic

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behaviors and mutual regulation of mRNAs and proteins within genes during transcription and translation. The models describing GRNs are mainly classified into two types: Boolean models^[4,5] and differential equation models^[6,7]. In the Boolean model, the state of the gene is viewed as either on or off. The differential equation model can describe the regulatory relationship between mRNAs and proteins in detail and accurately. Therefore, differential equation models are more suitable for modeling GRNs than Boolean network models.

The diffusion phenomenon is widespread; e.g., molecules propagate in the air, and mRNAs and proteins diffuse orderly within genes. However, in^[8,9], the authors default that mRNA and protein concentrations are uniformly distributed in space, which does not consider the existence of diffusion phenomena, making the constructed model inaccurate. Therefore, this paper introduces the reaction-diffusion term in the GRN model. During the modeling process, in addition to taking the diffusion phenomenon into account, it is crucial not to ignore the significance of time delay. This is because delay phenomena are present in the process of gene expression and demand attention. Based on the above analysis, we incorporate time delays in reaction-diffusion GRNs (RDGRNs)^[10-13]. In^[13], Zhang *et al.* constructed RDGRNs with constant delays. However, since constant delays cannot accurately describe the gene regulation process, many scholars investigate GRNs with time-varying delays ^[14-16]. Furthermore, compared with common time-varying delays^[17], mode-dependent time-varying delays can reduce the conservativeness of the stability criterion. Thus, Tian *et al.* in^[18] focused on the stability analysis of Markovian jump neural networks with mode-dependent time-varying delays. To the best of our knowledge, there has been no relevant research on RDGRNs with mode-dependent time-varying delays, which is one of the motivations for this paper.

For RDGRNs with time-varying delays, the mode of the system may switch when there are mutations in the gene sequence or changes in the external environment, and the probabilities of switching are described by the Markov chain^[19–21]. Thus, the investigations of RDGRNs with Markovian jump parameters and time-varying delays have received much attention^[22,23]. For example, in^[23], Zou *et al*. studied the robust stochastic stability of delayed RDGRNs with Markovian jump parameters. However, it is known that the switching probability of the system is difficult to obtain. Yet, in^[24] and^[25], the authors have assumed that the switching probabilities are completely known, which is not realistic. Therefore, it is worthwhile for Markovian jump RDGRNs with partially unknown transfer probabilities^[26–28] to be further scrutinized.

So far, for some applications of RDGRNs, it is a prerequisite to know the exact concentrations of mRNA and protein within genes. Nevertheless, due to the specificity of RDGRNs and the cost issue, the mRNA and protein concentrations are often only partially available, so it is extremely investigative to utilize the partially available state information to estimate the system state. In research on state estimation of RDGRNs^[29–31], there are defaults that the estimator parameters can be executed exactly. However, in general, the parameters may be perturbed to some extent due to factors such as component aging and internal noise, leading to system vulnerability. Therefore, some scholars have begun to focus on the non-fragile state estimation and synchronization of GRNs, and several results have been achieved^[32–34]. For example, Li *et al.* in^[32] investigated the issue of non-fragile state estimation for delayed GRNs. Nevertheless, the design of non-fragile state estimators for RDGRNs has not been investigated, which is another research motivation for this paper.

In light of the above discussion, this work studies the non-fragile state estimation for Markovian jump RDGRNs with partially unknown transfer probabilities and mode-dependent time-varying delays. The main innovations of this paper are as follows: (1) The reaction-diffusion term, mode-dependent time-varying delays, and Markovian jump parameters are considered in the model, which makes this model investigated in this paper more generalized compared to the existing studies on GRNs^[8]; (2) The partially unknown transfer probabilities^[26] are used in this work, which is more realistic compared to Markovian switching, where the transfer probabilities^[22] are fully known; and (3) The estimation performance of conventional state estimators^[29] is

affected when facing cases such as component aging and internal noise, whereas the non-fragile state estimator designed in this paper for RDGRNs makes the estimation performance non-fragile to gain variations.

Notations: $\Re^{n \times n}$ denote the space of $n \times n$ real matrices, $\operatorname{col}(\cdot)$ denotes the column vector, $\operatorname{diag}(\cdot)$ indicates the diagonal matrix, and $\operatorname{sym}\{\mathbb{M}\} = \mathbb{M} + \mathbb{M}^T$; $(g_{hq})_{n \times n}$ represents the $n \times n$ matrix consisting of the elements g_{hq} ; Ω denotes the set $\{v_1 \le y \le v_2\}$. In addition, this paper uses the following abbreviations: $\tilde{m}_h \triangleq \tilde{m}_h(x, y)$, $\tilde{p} \triangleq \tilde{p}_h(x, y), \bar{m} \triangleq \tilde{m}(x, y), \bar{p} \triangleq \bar{p}(x, y), L_m \triangleq L_m(x, y), L_p \triangleq L_p(x, y), \hat{m} \triangleq \hat{m}(x, y), and \hat{p} \triangleq \hat{p}(x, y)$.

2. NETWORK MODEL AND PRELIMINARIES

Consider the Markovian jump RDGRNs with mode-dependent time-varying delays as follows:

$$\begin{cases} \frac{\partial \tilde{m}_{h}}{\partial x} = e_{h} \frac{\partial^{2} \tilde{m}_{h}}{\partial y^{2}} - b_{h,\varepsilon(x)} \tilde{m}_{h} + \sum_{q=1}^{n} g_{hq,\varepsilon(x)} \xi_{q} \left(\tilde{p}_{q} \left(x - \alpha_{\varepsilon(x)}(x), y \right) \right) + \delta_{h}, \\ \frac{\partial \tilde{p}_{h}}{\partial x} = \bar{e}_{h} \frac{\partial^{2} \tilde{p}_{h}}{\partial y^{2}} - \bar{b}_{h,\varepsilon(x)} \tilde{p}_{h} + \bar{g}_{h,\varepsilon(x)} \tilde{m}_{h} \left(x - \beta_{\varepsilon(x)}(x), y \right), \end{cases}$$
(1)

where the border and initial conditions are given as follows:

$$\tilde{m}_h = 0, \ y \in \partial\Omega, \ x \in [-\beta, +\infty), \ \tilde{m}_h = \varphi(x, y), \ y \in \Omega, \ x \in [-\beta, 0],$$

$$\tilde{p}_h = 0, \ y \in \partial\Omega, \ x \in [-\alpha, +\infty), \ \tilde{p}_h = \theta(x, y), \ y \in \Omega, \ x \in [-\alpha, 0],$$

$$(2)$$

in which h = 1, 2, ..., n; $\partial \Omega$ indicates the boundary of Ω , $\varphi(x, y)$ and $\theta(x, y)$ denote the initial conditions of \tilde{m}_h and \tilde{p}_h , respectively. \tilde{m}_h and \tilde{p}_h are the mRNA and protein concentrations; x and $y \in \Omega$ denote time and space position, respectively; $\xi_q(\tilde{p}_q)$ denotes the monotonic feedback regulation function, which is described in the form $\xi_q(\tilde{p}_q) = \frac{(\frac{\tilde{p}_q}{\varrho_q})^{H_q}}{1+(\frac{\tilde{p}_q}{\varrho_q})^{H_q}}$, where H_q is the Hill coefficient representing the degree of cooperativity, ϱ_q is a positive constant, and q = 1, ..., n; the meaning and form of $\xi_q(\tilde{p}_q)$ are explained in detail by ^[35-37], so they are omitted here; $\delta_h = \sum_{q \in \mu_h} g^*_{hq, \varepsilon(x)}$, and μ_h is the set containing all the repressors of gene h; $\alpha_{\varepsilon(x)}(x)$ and $\beta_{\varepsilon(x)}(x)$ are time-varying delays that satisfy the following conditions

$$0 \le \alpha_{\varepsilon(x)}(x) \le \alpha_{\varepsilon(x)}, \ \dot{\alpha}_{\varepsilon(x)}(x) \le \psi_{1,\varepsilon(x)} < 1, \ 0 \le \beta_{\varepsilon(x)}(x) \le \beta_{\varepsilon(x)}, \ \dot{\beta}_{\varepsilon(x)}(x) \le \psi_{2,\varepsilon(x)} < 1,$$
(3)

with known scalars $\alpha_{\varepsilon(x)}$, $\beta_{\varepsilon(x)}$, $\psi_{1,\varepsilon(x)}$, and $\psi_{2,\varepsilon(x)}$. e_h and \bar{e}_h represent the diffusion rate; $b_{h,\varepsilon(x)}$ and $\bar{b}_{h,\varepsilon(x)}$ denote the mRNA and protein degradation rate, respectively; $\bar{g}_{h,\varepsilon(x)}$ indicates the translation rate; $G_{\varepsilon(x)} = (g_{hq,\varepsilon(x)})_{n\times n}$ with

$$g_{hq,\varepsilon(x)} = \begin{cases} g_{hq,\varepsilon(x)}^*, & \text{if } q \text{ is an activator of gene } h, \\ 0, & \text{if there is no link between gene } q \text{ and } h, \\ -g_{hq,\varepsilon(x)}^*, & \text{if } q \text{ is a repressor of gene } h, \end{cases}$$

denotes the coupling matrix, and $g^*_{hq,\varepsilon(x)}$ is the dimensionless transcriptional rate.

Define $\varepsilon(x) \in \mathbb{U} \triangleq \{1, 2, \dots, U\} (t \ge 0)$ to be a Markov chain, the corresponding transfer probability matrix $\prod \triangleq (\pi_{ab})_{U \times U}$ is given as

$$\operatorname{Prob}\{\varepsilon(t+\Delta t) = b \mid \varepsilon(x) = a\} = \begin{cases} \pi_{ab}\Delta t + 0(\Delta t), & a \neq b, \\ 1 + \pi_{ab}\Delta t + 0(\Delta t), & a = b, \end{cases}$$

where $\Delta t > 0$, $\lim_{\Delta t \to 0} \frac{0(\Delta t)}{\Delta t} = 0$, π_{ab} is the transfer probability from mode *a* to *b*, and $\pi_{aa} = -\sum_{a=1, a \neq b}^{U} \pi_{ab}$. Moreover, the transfer matrix with partially unknown probabilities can be expressed as

$$\begin{bmatrix} \pi_{11} & ? & \dots & ? \\ \pi_{21} & ? & \dots & \pi_{2U} \\ \vdots & \vdots & \ddots & \vdots \\ \pi_{U1} & ? & \dots & \pi_{UU} \end{bmatrix}$$
(4)

where "?" represents the unknown transfer probabilities. Define $\Re_r^a \triangleq \{b : \pi_{ab} \text{ is known}\}, \Re_{ur}^a \triangleq \{b : \pi_{ab} \text{ is unknown}\}$, and $\Re \triangleq \Re_r^a + \Re_{ur}^a$.

Let $\tilde{m}^*(y) = \operatorname{col}\left(\tilde{m}_1^*(y), \tilde{m}_2^*(y), \dots, \tilde{m}_n^*(y)\right)$ and $\tilde{p}^*(y) = \operatorname{col}\left(\tilde{p}_1^*(y), \tilde{p}_2^*(y), \dots, \tilde{p}_n^*(y)\right)$ be the unique equilibrium point of the system (1), that is

$$\begin{cases} 0 = e_h \frac{\partial^2 \tilde{m}_h^*(y)}{\partial y^2} - b_{h,\varepsilon(x)} \tilde{m}_h^*(y) + \sum_{q=1}^n g_{hq,\varepsilon(x)} \xi_q(\tilde{p}_q^*(y)) + \delta_h, \\ 0 = \bar{e}_h \frac{\partial^2 \tilde{p}_h^*(y)}{\partial y^2} - \bar{b}_{h,\varepsilon(x)} \tilde{p}_h^*(y) + \bar{g}_{h,\varepsilon(x)} \tilde{m}_h^*(y). \end{cases}$$

Define $\bar{m}_h = \tilde{m}_h - \tilde{m}_h^*(y)$ and $\bar{p}_h = \tilde{p}_h - \tilde{p}_h^*(y)$, the system (1) is converted into

$$\begin{cases} \frac{\partial \bar{m}}{\partial x} = E \frac{\partial^2 \bar{m}}{\partial y^2} - B_{\varepsilon(x)} \bar{m} + G_{\varepsilon(x)} f(\bar{p}(x - \alpha_{\varepsilon(x)}(x), y)), \\ \frac{\partial \bar{p}}{\partial x} = \bar{E} \frac{\partial^2 \bar{p}}{\partial y^2} - \bar{B}_{\varepsilon(x)} \bar{p} + \bar{G}_{\varepsilon(x)} \bar{m}(x - \beta_{\varepsilon(x)}(x), y), \end{cases}$$
(5)

where

$$\begin{split} \bar{m} &= \operatorname{col}(\bar{m}_1, \bar{m}_2, \dots, \bar{m}_n), \ E = \operatorname{diag}(e_1, e_2, \dots, e_n), \ B_{\varepsilon(x)} = \operatorname{diag}(b_{1,\varepsilon(x)}, b_{2,\varepsilon(x)}, \dots, b_{n,\varepsilon(x)}), \\ \bar{p} &= \operatorname{col}(\bar{p}_1, \bar{p}_2, \dots, \bar{p}_n), \ \bar{E} = \operatorname{diag}(\bar{e}_1, \bar{e}_2, \dots, \bar{e}_n), \ \bar{B}_{\varepsilon(x)} = \operatorname{diag}(\bar{b}_{1,\varepsilon(x)}, \bar{b}_{2,\varepsilon(x)}, \dots, \bar{b}_{n,\varepsilon(x)}), \\ \bar{G}_{\varepsilon(x)} &= \operatorname{diag}(\bar{g}_{1,\varepsilon(x)}, \dots, \bar{g}_{n,\varepsilon(x)}), \ f_h(\bar{p}_h(x - \alpha_{\varepsilon(x)}(x), y)) = \xi_h(\bar{p}_h(x - \alpha_{\varepsilon(x)}(x), y) + \tilde{p}_h^*(y)) - \xi_h(\tilde{p}_h^*(y)), \\ f(\bar{p}(x - \alpha_{\varepsilon(x)}(x), y)) &= \operatorname{col}(f_1(\bar{p}_1(x - \alpha_{\varepsilon(x)}(x), y)), f_2(\bar{p}_2(x - \alpha_{\varepsilon(x)}(x), y)), \dots, f_n(\bar{p}_n(x - \alpha_{\varepsilon(x)}(x), y))). \end{split}$$

Additionally, the measured outputs of (5) are

$$L_m = N\bar{m}, \ L_p = \bar{N}\bar{p}, \tag{6}$$

where $N \in \mathfrak{R}^{n \times n}$ and $\overline{N} \in \mathfrak{R}^{n \times n}$ are known matrices; L_m and L_p represent the measured outputs of mRNA and protein, respectively.

To estimate the states of (5) by using the measured outputs in (6), one can construct the estimator as follows:

$$\begin{cases} \frac{\partial \hat{m}}{\partial x} = E \frac{\partial^2 \hat{m}}{\partial y^2} - B_{\varepsilon(x)} \hat{m} + G_{\varepsilon(x)} f(\hat{p}(x - \alpha_{\varepsilon(x)}(x), y)) + (K_{1,\varepsilon(x)} + \Delta K_{1,\varepsilon(x)})(L_m - N\hat{m}), \\ \frac{\partial \hat{p}}{\partial x} = \bar{E} \frac{\partial^2 \hat{p}}{\partial y^2} - \bar{B}_{\varepsilon(x)} \hat{p} + \bar{G}_{\varepsilon(x)} \hat{m}(x - \beta_{\varepsilon(x)}(x), y) + (K_{2,\varepsilon(x)} + \Delta K_{2,\varepsilon(x)})(L_p - \bar{N}\hat{p}), \end{cases}$$
(7)

where \hat{m} and \hat{p} represent the estimations of \bar{m} and \bar{p} , $K_{1,\varepsilon(x)}$ and $K_{2,\varepsilon(x)}$ are the estimator gain matrices, and $\Delta K_{1,\varepsilon(x)}$ and $\Delta K_{2,\varepsilon(x)}$ denote the gain perturbation, which takes the following form:

$$\Delta K_{1,\varepsilon(x)} = A_{1,\varepsilon(x)} \Lambda_{1,\varepsilon(x)}(x) C_{1,\varepsilon(x)}, \ \Delta K_{2,\varepsilon(x)} = A_{2,\varepsilon(x)} \Lambda_{2,\varepsilon(x)}(x) C_{2,\varepsilon(x)}, \tag{8}$$

where $A_{1,\varepsilon(x)}$, $C_{1,\varepsilon(x)}$, $A_{2,\varepsilon(x)}$, and $C_{2,\varepsilon(x)}$ stand for the given matrices; $\Lambda_{1,\varepsilon(x)}(x)$ and $\Lambda_{2,\varepsilon(x)}(x)$ are unknown matrices satisfying $\Lambda_{1,\varepsilon(x)}^{T}(x)\Lambda_{1,\varepsilon(x)}(x) \leq I$ and $\Lambda_{2,\varepsilon(x)}^{T}(x)\Lambda_{2,\varepsilon(x)}(x) \leq I$, respectively.

Set $\varepsilon(x) = a$, we denote $B_a \triangleq B_{\varepsilon(x)}$, $G_a \triangleq G_{\varepsilon(x)}$, $\alpha_a \triangleq \alpha_{\varepsilon(x)}$, $\bar{B}_a \triangleq \bar{B}_{\varepsilon(x)}$, $\bar{G}_a \triangleq \bar{G}_{\varepsilon(x)}$, $\beta_a \triangleq \beta_{\varepsilon(x)}$, $K_{da} \triangleq K_{d,\varepsilon(x)}$, $\Delta K_{da} \triangleq \Delta K_{d,\varepsilon(x)}$, $A_{da} \triangleq A_{d,\varepsilon(x)}$, $\Lambda_{da} \triangleq \Lambda_{d,\varepsilon(x)}$, $C_{da} \triangleq C_{d,\varepsilon(x)}$, (d = 1, 2). Define $\phi_m = \bar{m} - \hat{m}$ and $\phi_p = \bar{p} - \hat{p}$ to be the error state vectors; based on (5)–(7), the following error system is derived,

$$\begin{cases} \frac{\partial \phi_m}{\partial x} = E \frac{\partial^2 \phi_m}{\partial y^2} - B_a \phi_m + G_a f(\phi_p(x - \alpha_a(x), y)) - (K_{1a} + \Delta K_{1a}) W \phi_m, \\ \frac{\partial \phi_p}{\partial x} = \bar{E} \frac{\partial^2 \phi_p}{\partial y^2} - \bar{B}_a \phi_p + \bar{G}_a \phi_m(x - \beta_a(x), y) - (K_{2a} + \Delta K_{2a}) \bar{W} \phi_p, \end{cases}$$
(9)

where $f(\phi_p(x - \alpha_a(x), y)) = f(\bar{p}(x - \alpha_a(x), y)) - f(\hat{p}(x - \alpha_a(x), y)), \phi_m(x - \beta_a(x), y) = \bar{m}(x - \beta_a(x), y) - \hat{m}(x - \beta_a(x), y), \Delta K_{1a} = A_{1a}\Lambda_{1a}(x)C_{1a}, \text{ and } \Delta K_{2a} = A_{2a}\Lambda_{2a}(x)C_{2a}.$ Because $f_h(\cdot)$ is a monotonically increasing function, which satisfies the following inequality

$$0 \le \frac{J_h(z)}{z} \le \vartheta_h, \ \forall z \ne 0; \ h = 1, \dots, n,$$
(10)

where ϑ_h is a known positive real number.

The following lemmas are important tools for obtaining the principal results of this article.

Lemma 1.^[38] Assume $\hbar \ge 0 \in \Re^{n \times n}$ and the function $\ell \in \mathcal{H}_l^n([r_1, r_2])$ satisfying $\ell(r_1) = 0$ or $\ell(r_2) = 0$, then

$$\int_{r_1}^{r_2} \ell^T(y) \hbar \ell(y) dy \le \frac{4(r_2 - r_1)^2}{\pi^2} \int_{r_1}^{r_2} \dot{\ell}(y)^T \hbar \dot{\ell}(y) dy.$$

Further, if $\ell(r_1) = \ell(r_2) = 0$, one has

$$\int_{r_1}^{r_2} \ell^T(y) \hbar \ell(y) dy \le \frac{(r_2 - r_1)^2}{\pi^2} \int_{r_1}^{r_2} \dot{\ell}(y)^T \hbar \dot{\ell}(y) dy.$$

Lemma 2.^[39] With real matrices H and J of appropriate dimensions, the inequality satisfies

$$HJ^T + JH^T \le \gamma^{-1}HH^T + \gamma JJ^T,$$

where γ is a positive scalar.

Lemma 3.^[40] The existence of matrices $\Psi = \Psi^{T}$, *H* and *J* of appropriate dimensions, then

$$\Psi + H\Lambda(x)J + H^{\mathrm{T}}\Lambda^{\mathrm{T}}(x)J^{\mathrm{T}} < 0,$$

for all satisfying $\Lambda^{T}(x)\Lambda(x) \leq I$, the sufficient condition for $\Lambda(x)$ to hold is the existence of a positive constant γ such that the following equation holds:

$$\Psi + \gamma^{-1} H H^{\mathrm{T}} + \gamma J^{\mathrm{T}} J < 0.$$

3. MAIN RESULTS

For this section, establishing new criteria by choosing appropriate Lyapunov-Krasovskii functionals (LKFs), which assures that system (9) is asymptotically stable.

Theorem 1. Given constants ψ_{1a} , ψ_{2a} , ϖ , matrix \mathbb{N} , and estimator gain matrices K_{1a} and K_{2a} , the system (9) is asymptotically stable if there exist positive matrices P_{la} , F_l , Q_l , (l = 1, 2) with suitable dimensions, symmetric matrices $W_a \in \mathfrak{R}^{n \times n}$ and $\overline{W}_a \in \mathfrak{R}^{n \times n}$, and the diagonal matrix $R_1 > 0 \in \mathfrak{R}^{n \times n}$, which makes the below linear matrix inequalities (LMIs) feasible:

$$\Psi_1 + \Psi_2 + \Psi_3 \le 0, \tag{11}$$

$$P_{1b} + W_a \le 0, \ P_{2b} + \bar{W}_a \le 0, \ \forall b \in \Re^a_{wr}, \ a \ne b,$$
 (12)

where

$$\begin{split} \Psi_{1} = & \operatorname{sym}\{\zeta_{1}^{T}P_{1a}\zeta_{6} + \zeta_{2}^{T}P_{2a}\zeta_{7}\} + \zeta_{1}^{T}\sum_{b=\Re_{r}^{a}}\pi_{ab}(P_{1b}+W_{a})\zeta_{1} + \zeta_{2}^{T}\sum_{b=\Re_{r}^{a}}\pi_{ab}(P_{2b}+\bar{W}_{a})\zeta_{2} + \zeta_{1}^{T}F_{1}\zeta_{1} + \zeta_{2}^{T}F_{2}\zeta_{2} \\ &-(1-\psi_{2a})\zeta_{3}^{T}F_{1}\zeta_{3} - (1-\psi_{1a})\zeta_{4}^{T}F_{2}\zeta_{4} + \operatorname{sym}\{\zeta_{5}^{T}R_{1}(\mathbb{N}\zeta_{4}-\zeta_{5})\}, \\ \Psi_{2} = & \operatorname{sym}\{\zeta_{1}^{T}Q_{1}B_{a}\zeta_{1}\} - \operatorname{sym}\{\zeta_{1}^{T}\frac{\pi^{2}}{\varpi^{2}}Q_{1}E\zeta_{1}\} - \operatorname{sym}\{\zeta_{1}^{T}Q_{1}B_{a}\zeta_{6}\} + \operatorname{sym}\{\zeta_{1}^{T}Q_{1}G_{a}\zeta_{5}\} - \operatorname{sym}\{\zeta_{1}^{T}Q_{1}\zeta_{6}\} \\ &+ \operatorname{sym}\{\zeta_{5}^{T}Q_{1}G_{a}\zeta_{6}\} - \operatorname{sym}\{\zeta_{6}^{T}Q_{1}\zeta_{6}\} - \operatorname{sym}\{\zeta_{1}^{T}Q_{1}K_{1a}N\zeta_{1}\} - \operatorname{sym}\{\zeta_{1}^{T}Q_{1}\Delta K_{1a}N\zeta_{1}\} - \operatorname{sym}\{\zeta_{1}^{T}Q_{1}\Delta K_{1a}N\zeta_{6}\}, \\ \Psi_{3} = & -\operatorname{sym}\{\zeta_{2}^{T}Q_{2}\bar{B}_{a}\zeta_{2}\} - \operatorname{sym}\{\zeta_{2}^{T}\frac{\pi^{2}}{\varpi^{2}}Q_{2}\bar{E}\zeta_{2}\} - \operatorname{sym}\{\zeta_{2}^{T}Q_{2}\bar{B}_{a}\zeta_{7}\} + \operatorname{sym}\{\zeta_{2}^{T}Q_{2}\bar{G}_{a}\zeta_{3}\} - \operatorname{sym}\{\zeta_{2}^{T}Q_{2}\zeta_{7}\} \\ &+ \operatorname{sym}\{\zeta_{3}^{T}Q_{2}\bar{G}_{a}\zeta_{7}\} - \operatorname{sym}\{\zeta_{7}^{T}Q_{2}\zeta_{7}\} - \operatorname{sym}\{\zeta_{2}^{T}Q_{2}K_{2a}\bar{N}\zeta_{2}\} - \operatorname{sym}\{\zeta_{2}^{T}Q_{2}\Delta K_{2a}\bar{N}\zeta_{2}\} - \operatorname{sym}\{\zeta_{2}^{T}Q_{2}\Delta K_{2a}\bar{N}\zeta_{7}\} \\ &- \operatorname{sym}\{\zeta_{2}^{T}Q_{2}\Delta K_{2a}\bar{N}\zeta_{7}\}, \end{split}$$

where $\psi_{1a} = \psi_{1,\varepsilon(x)}, \psi_{2a} = \psi_{2,\varepsilon(x)}, \mathbb{N} = \text{diag}(\mathbb{N}_1, \mathbb{N}_2, \dots, \mathbb{N}_n), \varpi = \nu_2 - \nu_1, \text{ and } \zeta_{\upsilon} = [0_{n,(\upsilon-1)n} I_n 0_{n,(7-\upsilon)n}], (\upsilon = 1, \dots, 7).$

Proof. Construct the LKFs for the system (9) as follows:

$$V(x) = V_1(x) + V_2(x) + V_3(x),$$
(13)

where

$$\begin{split} V_1(x) &= \int_{\Omega} \phi_m^T P_{1a} \phi_m dy + \int_{\Omega} \phi_p^T P_{2a} \phi_p dy, \\ V_2(x) &= \int_{\Omega} \frac{\partial \phi_m^T}{\partial y} Q_1 E \frac{\partial \phi_m}{\partial y} dy + \int_{\Omega} \frac{\partial \phi_p^T}{\partial y} Q_2 \bar{E} \frac{\partial \phi_p}{\partial y} dy, \\ V_3(x) &= \int_{\Omega} \int_{x-\beta_a(x)}^x \phi_m^T(z, y) F_1 \phi_m(z, y) dz dy + \int_{\Omega} \int_{x-\alpha_a(x)}^x \phi_p^T(z, y) F_2 \phi_p(z, y) dz dy. \end{split}$$

The time derivative of V(x) is calculated as follows:

$$\dot{V}_1(x) = 2\int_{\Omega} \phi_m^T P_{1a} \frac{\partial \phi_m}{\partial x} dy + \int_{\Omega} \sum_{b=1}^U \pi_{ab} \phi_m^T P_{1b} \phi_m dy + 2\int_{\Omega} \phi_p^T P_{2a} \frac{\partial \phi_p}{\partial x} dy + \int_{\Omega} \sum_{b=1}^U \pi_{ab} \phi_p^T P_{2b} \phi_p dy, \quad (14)$$

$$\begin{split} \dot{V}_{2}(x) &= 2 \int_{\Omega} \frac{\partial^{2} \phi_{m}^{T}}{\partial y \partial x} Q_{1} E \frac{\partial \phi_{m}}{\partial y} dy + 2 \int_{\Omega} \frac{\partial^{2} \phi_{p}^{T}}{\partial y \partial x} Q_{2} \bar{E} \frac{\partial \phi_{p}}{\partial y} dy, \end{split}$$
(15)
$$\dot{V}_{3}(x) &= \int_{\Omega} \phi_{m}^{T} F_{1} \phi_{m} dy - (1 - \dot{\beta}_{a}(x)) \int_{\Omega} \phi_{m}^{T} (x - \beta_{a}(x), y) F_{1} \phi_{m} (x - \beta_{a}(x), y) dy \\ &+ \int_{\Omega} \phi_{p}^{T} F_{2} \phi_{p} dy - (1 - \dot{\alpha}_{a}(x)) \int_{\Omega} \phi_{p}^{T} (x - \alpha_{a}(x), y) F_{2} \phi_{p} (x - \alpha_{a}(x), y) dy \\ &\leq \int_{\Omega} \phi_{m}^{T} F_{1} \phi_{m} dy - (1 - \psi_{2a}) \int_{\Omega} \phi_{m}^{T} (x - \beta_{a}(x), y) F_{1} \phi_{m} (x - \beta_{a}(x), y) dy \\ &+ \int_{\Omega} \phi_{p}^{T} F_{2} \phi_{p} dy - (1 - \psi_{1a}) \int_{\Omega} \phi_{p}^{T} (x - \alpha_{a}(x), y) F_{2} \phi_{p} (x - \alpha_{a}(x), y) dy. \end{split}$$
(16)

According to (9), the following is obviously true

$$0 = 2 \int_{\Omega} \left(\phi_m + \frac{\partial \phi_m}{\partial x} \right)^T Q_1 \left[-\frac{\partial \phi_m}{\partial x} + E \frac{\partial^2 \phi_m}{\partial y^2} - B_a \phi_m + G_a f(\phi_p(x - \alpha_a(x), y)) - (K_{1a} + \Delta K_{1a}) N \phi_m \right] dy,$$

$$0 = 2 \int_{\Omega} \left(\phi_p + \frac{\partial \phi_p}{\partial x} \right)^T Q_2 \left[-\frac{\partial \phi_p}{\partial x} + \bar{E} \frac{\partial^2 \phi_p}{\partial y^2} - \bar{B}_a \phi_p + \bar{G}_a \phi_m(x - \beta_a(x), y) - (K_{2a} + \Delta K_{2a}) \bar{N} \phi_p \right] dy.$$
(17)

According to the definition of the transfer probability matrix (4), for symmetric matrices $W_a \in \mathbb{R}^{n \times n}$ and $\overline{W}_a \in \mathbb{R}^{n \times n}$, one gets

$$0 = \sum_{b=1}^{U} \pi_{ab} \phi_m^T W_a \phi_m, \ 0 = \sum_{b=1}^{U} \pi_{ab} \phi_p^T \bar{W}_a \phi_p, \ \forall a \in U.$$
(18)

Based on (17) and the boundary conditions (2), using the integration by parts, one can get

$$2\int_{\Omega}\phi_m^T Q_1 E \frac{\partial^2 \phi_m}{\partial y^2} dy = -2\int_{\Omega} \frac{\partial \phi_m^T}{\partial y} Q_1 E \frac{\partial \phi_m}{\partial y} dy,$$
(19)

and

$$2\int_{\Omega}\phi_p^T Q_2 \bar{E}\frac{\partial^2 \phi_p}{\partial y^2} dy = -2\int_{\Omega}\frac{\partial \phi_p^T}{\partial y} Q_2 \bar{E}\frac{\partial \phi_p}{\partial y} dy.$$
(20)

Following (19) and (20) and applying Lemma 1, we can derive that

$$-2\int_{\Omega}\frac{\partial\phi_m^T}{\partial y}Q_1E\frac{\partial\phi_m}{\partial y} + \frac{\partial\phi_p^T}{\partial y}Q_2\bar{E}\frac{\partial\phi_p}{\partial y}dy \le -2\frac{\pi^2}{\varpi^2}\int_{\Omega}\phi_m^TQ_1E\phi_m + \phi_p^TQ_2\bar{E}\phi_pdy.$$
(21)

From (17) and the boundary conditions (2), we have that

$$2\int_{\Omega} \frac{\partial \phi_m^T}{\partial x} Q_1 E \frac{\partial^2 \phi_m}{\partial y^2} dy = -2\int_{\Omega} \frac{\partial \phi_m^T}{\partial y} Q_1 E \frac{\partial^2 \phi_m}{\partial y \partial x} dy, \tag{22}$$

and

$$2\int_{\Omega} \frac{\partial \phi_p^T}{\partial x} Q_2 \bar{E} \frac{\partial^2 \phi_p}{\partial y^2} dy = -2\int_{\Omega} \frac{\partial \phi_p^T}{\partial y} Q_2 \bar{E} \frac{\partial^2 \phi_p}{\partial y \partial x} dy.$$
(23)

Based on (10), for arbitrary diagonal matrix $R_1 > 0$, one obtains

$$0 \le 2f^{T}(\phi_{p}(x - \alpha_{a}(x), y))R_{1}(\mathbb{N}\phi_{p} - f(\phi_{p}(x - \alpha_{a}(x), y))).$$
(24)

By combining (14)-(24), it can be proved that

$$\dot{V}(x) \leq \int_{\Omega} \ell^{T}(x, y) (\Psi_{1} + \Psi_{2} + \Psi_{3}) \ell(x, y) dy + \int_{\Omega} \sum_{b = \Re_{ur}^{a}} \pi_{ab} \phi_{m}^{T} (P_{1b} + W_{a}) \phi_{m} + \sum_{b = \Re_{ur}^{a}} \pi_{ab} \phi_{p}^{T} (P_{2b} + \bar{W}_{a}) \phi_{p} dy,$$
(25)

where $\ell(x, y) = \operatorname{col}\left(\phi_m, \phi_p, \phi_m(x - \beta_a(x), y), \phi_p(x - \alpha_a(x), y), f(\phi_p(x - \alpha_a(x), y)), \frac{\partial \phi_m}{\partial x}, \frac{\partial \phi_p}{\partial x}\right)$. Therefore, combining the above analysis, one can conclude that the error system (9) is asymptotically stable. The proof is complete.

Having performed the stability analysis in Theorem 1, we will deal with the design of a non-fragile estimator. The following theorem gives a solution for designing a non-fragile estimator.

Theorem 2. Given positive constants ψ_{1a} , ψ_{2a} , ϖ , γ_1 , γ_2 , and matrix \mathbb{N} , the error system (9) is asymptotically stable if there exist positive matrices P_{la} , F_l , Q_l , Γ_{la} , (l = 1, 2) with suitable dimensions, symmetric matrices $W_a \in \mathbb{R}^{n \times n}$ and $\overline{W}_a \in \mathbb{R}^{n \times n}$, and the diagonal matrix $R_1 > 0 \in \mathbb{R}^{n \times n}$, which makes the below LMIs feasible:

$$\begin{bmatrix} \hat{\Psi} & \gamma_1 \mathbb{A}_{1a} & \mathbb{C}_{1a}^T & \gamma_2 \mathbb{A}_{2a} & \mathbb{C}_{2a}^T \\ * & -\gamma_1 I & 0 & 0 & 0 \\ * & * & -\gamma_1 I & 0 & 0 \\ * & * & * & -\gamma_2 I & 0 \\ * & * & * & * & -\gamma_2 I \end{bmatrix} \leq 0,$$
(26)

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$$\begin{split} \mathbb{A}_{2a} &= \begin{bmatrix} 0 & -A_{2a}^{T} Q_{2}^{T} & \underbrace{0 & 0 & \dots & 0}_{5 \text{ times}} \end{bmatrix}^{T}, \mathbb{C}_{2a} = \begin{bmatrix} 0 & C_{2a}\bar{N} & \underbrace{0 & 0 & \dots & 0}_{4 \text{ times}} & C_{2a}\bar{N} \end{bmatrix}, \text{ and} \\ \Psi_{1} &= \text{sym}\{\zeta_{1}^{T} P_{1a}\zeta_{6} + \zeta_{2}^{T} P_{2a}\zeta_{7}\} + \zeta_{1}^{T} \sum_{b=\Re_{r}^{\alpha}} \pi_{ab}(P_{1b} + W_{a})\zeta_{1} + \zeta_{2}^{T} \sum_{b=\Re_{r}^{\alpha}} \pi_{ab}(P_{2b} + \bar{W}_{a})\zeta_{2} + \zeta_{1}^{T} F_{1}\zeta_{1} + \zeta_{2}^{T} F_{2}\zeta_{2} \\ &- (1 - \psi_{2a})\zeta_{3}^{T} F_{1}\zeta_{3} - (1 - \psi_{1a})\zeta_{4}^{T} F_{2}\zeta_{4} + \text{sym}\{\zeta_{5}^{T} R_{1}(\mathbb{N}\zeta_{4} - \zeta_{5})\}, \\ \hat{\Psi}_{1} &= - \text{sym}\{\zeta_{1}^{T} Q_{1} B_{a}\zeta_{1}\} - \text{sym}\{\zeta_{1}^{T} \frac{\pi^{2}}{\varpi^{2}} Q_{1} E\zeta_{1}\} - \text{sym}\{\zeta_{1}^{T} Q_{1} B_{a}\zeta_{6}\} + \text{sym}\{\zeta_{1}^{T} Q_{1} G_{a}\zeta_{5}\} - \text{sym}\{\zeta_{1}^{T} Q_{1}\zeta_{6}\} \\ &+ \text{sym}\{\zeta_{5}^{T} Q_{1} G_{a}\zeta_{6}\} - \text{sym}\{\zeta_{6}^{T} Q_{1}\zeta_{6}\} - \text{sym}\{\zeta_{1}^{T} \Gamma_{1a} N\zeta_{1}\} - \text{sym}\{\zeta_{1}^{T} \Gamma_{1a} N\zeta_{6}\}, \\ \hat{\Psi}_{2} &= - \text{sym}\{\zeta_{2}^{T} Q_{2} \bar{B}_{a}\zeta_{2}\} - \text{sym}\{\zeta_{2}^{T} \frac{\pi^{2}}{\varpi^{2}} Q_{2} \bar{E}\zeta_{2}\} - \text{sym}\{\zeta_{2}^{T} Q_{2} \bar{B}_{a}\zeta_{7}\} + \text{sym}\{\zeta_{2}^{T} Q_{2} \bar{G}_{a}\zeta_{3}\} - \text{sym}\{\zeta_{2}^{T} Q_{2}\zeta_{7}\} \\ &+ \text{sym}\{\zeta_{3}^{T} Q_{2} \bar{G}_{a}\zeta_{7}\} - \text{sym}\{\zeta_{7}^{T} Q_{2}\zeta_{7}\} - \text{sym}\{\zeta_{2}^{T} \Gamma_{2a} \bar{N}\zeta_{2}\} - \text{sym}\{\zeta_{2}^{T} \Gamma_{2a} \bar{N}\zeta_{7}\}. \end{split}$$

Furthermore, the other parameters are consistent with Theorem 1, and the estimator gain matrices are obtained by $K_{1a} = \bar{Q}_1^{-1} \Gamma_{1a}$ and $K_{2a} = \bar{Q}_2^{-1} \Gamma_{2a}$.

Proof. Based on (8) and the proof procedure of Theorem 1, one obtains that

$$\Theta = \hat{\Psi} + \mathbb{A}_{1a}\Lambda_{1a}\mathbb{C}_{1a} + \mathbb{C}_{1a}^{\mathrm{T}}\Lambda_{1a}^{\mathrm{T}}\mathbb{A}_{1a}^{\mathrm{T}} + \mathbb{A}_{2a}\Lambda_{2a}\mathbb{C}_{2a} + \mathbb{C}_{2a}^{\mathrm{T}}\Lambda_{2a}^{\mathrm{T}}\mathbb{A}_{2a}^{\mathrm{T}} \le 0.$$
(27)

Since $\Lambda_{1a}^{\mathrm{T}}(x)\Lambda_{1a}(x) \leq I$ and $\Lambda_{2a}^{\mathrm{T}}(x)\Lambda_{2a}(x) \leq I$, by using Lemma 2, one obtains

$$\mathbb{A}_{1a}\Lambda_{1a}\mathbb{C}_{1a} + \mathbb{C}_{1a}^{\mathrm{T}}\Lambda_{1a}^{\mathrm{T}}\mathbb{A}_{1a}^{\mathrm{T}} \leq \gamma_{1}\mathbb{A}_{1a}\mathbb{A}_{1a}^{\mathrm{T}} + \gamma_{1}^{-1}\mathbb{C}_{1a}^{\mathrm{T}}\mathbb{C}_{1a},$$

$$(28)$$

and

$$\mathbb{A}_{2a}\Lambda_{2a}\mathbb{C}_{2a} + \mathbb{C}_{2a}^{\mathrm{T}}\Lambda_{2a}^{\mathrm{T}}\mathbb{A}_{2a}^{\mathrm{T}} \leq \gamma_{2}\mathbb{A}_{2a}\mathbb{A}_{2a}^{\mathrm{T}} + \gamma_{2}^{-1}\mathbb{C}_{2a}^{\mathrm{T}}\mathbb{C}_{2a}.$$
(29)

According to (27), (28), and (29), by using Lemma 3, we can get

$$\Theta \leq \hat{\Psi} + \gamma_1 \mathbb{A}_{1a} \mathbb{A}_{1a}^{\mathrm{T}} + \gamma_1^{-1} \mathbb{C}_{1a} \mathbb{C}_{1a}^{\mathrm{T}} + \gamma_2 \mathbb{A}_{2a} \mathbb{A}_{2a}^{\mathrm{T}} + \gamma_2^{-1} \mathbb{C}_{2a} \mathbb{C}_{2a}^{\mathrm{T}} \leq 0,$$
(30)

where \mathbb{A}_{1a} , \mathbb{A}_{2a} , \mathbb{C}_{1a} , and \mathbb{C}_{2a} are already given in the statement of Theorem 2.

By combining (14)-(24) and (27)-(30), it can be proved that

$$\dot{V}(x) \leq \int_{\Omega} \ell^{T}(x, y) (\hat{\Psi} + \gamma_{1} \mathbb{A}_{1a} \mathbb{A}_{1a}^{T} + \gamma_{1}^{-1} \mathbb{C}_{1a} \mathbb{C}_{1a}^{T} + \gamma_{2} \mathbb{A}_{2a} \mathbb{A}_{2a}^{T} + \gamma_{2}^{-1} \mathbb{C}_{2a} \mathbb{C}_{2a}^{T}) \ell(x, y) dy + \int_{\Omega} \sum_{b = \Re_{ur}^{a}} \pi_{ab} \phi_{m}^{T}(P_{1b} + W_{a}) \phi_{m} + \sum_{b = \Re_{ur}^{a}} \pi_{ab} \phi_{p}^{T}(P_{2b} + \bar{W}_{a}) \phi_{p} dy,$$
(31)

Table 1. The mode-dependent time-varying delays and related parameters

	$\alpha_a(x)$	$\beta_a(x)$	α_a	β_a	ψ_{1a}	ψ_{2a}
<i>a</i> = 1	$0.3 + 0.3 \sin(x)$	$0.25 + 0.25 \sin(x)$	0.6	0.5	0.3	0.25
<i>a</i> = 2	$0.4 + 0.4 \sin(x)$	$0.35+0.35\sin(x)$	0.8	0.7	0.4	0.35
<i>a</i> = 3	$0.5+0.5\sin(x)$	$0.45 + 0.45\sin(x)$	1	0.9	0.5	0.45

where $\ell(x, y) = \operatorname{col}\left(\phi_m, \phi_p, \phi_m(x - \beta_a(x), y), \phi_p(x - \alpha_a(x), y), f(\phi_p(x - \alpha_a(x), y)), \frac{\partial \phi_m}{\partial x}, \frac{\partial \phi_p}{\partial x}\right)$, we employ Schur's complement, which can convert the inequality (30) into (26).

Combining the above analysis, we can conclude that the system (9) is asymptotically stable when the inequalities (12) and (26) hold. The proof is complete.

4. SIMULATION EXAMPLE

In this section, a numerical simulation is provided to illustrate the validity of the theoretical results in this paper. The corresponding parameters of the system (1) are selected as:

$$\begin{split} B_1 &= \begin{bmatrix} 0.74 & 0 & 0 \\ 0 & 0.72 & 0 \\ 0 & 0 & 0.65 \end{bmatrix}, B_2 &= \begin{bmatrix} 0.69 & 0 & 0 \\ 0 & 0.75 & 0 \\ 0 & 0 & 0.71 \end{bmatrix}, B_3 &= \begin{bmatrix} 0.71 & 0 & 0 \\ 0 & 0.74 & 0 \\ 0 & 0 & 0.67 \end{bmatrix}, N = \begin{bmatrix} 1.0 & 0 & 0 \\ 0 & 0.8 & 0 \\ 0 & 0 & 1.1 \end{bmatrix}, \\ \bar{B}_1 &= \begin{bmatrix} 0.75 & 0 & 0 \\ 0 & 0.69 & 0 \\ 0 & 0 & 0.62 \end{bmatrix}, \bar{B}_2 &= \begin{bmatrix} 0.77 & 0 & 0 \\ 0 & 0.71 & 0 \\ 0 & 0 & 0.68 \end{bmatrix}, \bar{B}_3 &= \begin{bmatrix} 0.72 & 0 & 0 \\ 0 & 0.76 & 0 \\ 0 & 0 & 0.72 \end{bmatrix}, \bar{N} = \begin{bmatrix} 1.1 & 0 & 0 \\ 0 & 1.3 & 0 \\ 0 & 0 & 0.9 \end{bmatrix}, \\ G_1 &= \begin{bmatrix} 0.49 & -0.29 & 0.43 \\ -0.52 & 0.91 & 0.63 \\ 0.51 & 0.34 & -0.62 \end{bmatrix}, G_2 &= \begin{bmatrix} 0.53 & -0.34 & 0.42 \\ -0.52 & 0.94 & 0.61 \\ 0.53 & 0.31 & -0.63 \end{bmatrix}, G_3 &= \begin{bmatrix} 0.55 & -0.36 & 0.41 \\ -0.53 & 0.97 & 0.67 \\ 0.56 & 0.37 & -0.68 \end{bmatrix}, \\ \bar{G}_1 &= \begin{bmatrix} 0.7 & 0 & 0 \\ 0 & 0.7 & 0 \\ 0 & 0 & 0.6 \end{bmatrix}, \bar{G}_2 &= \begin{bmatrix} 0.6 & 0 & 0 \\ 0 & 0.6 & 0 \\ 0 & 0 & 0.6 \end{bmatrix}, \bar{G}_3 &= \begin{bmatrix} 0.5 & 0 & 0 \\ 0 & 0.55 & 0 \\ 0 & 0 & 0.5 \end{bmatrix}, E &= \begin{bmatrix} 0.15 & 0 & 0 \\ 0 & 0.16 & 0 \\ 0 & 0 & 0.13 \end{bmatrix}, \\ \bar{E} &= \begin{bmatrix} 0.2 & 0 & 0 \\ 0 & 0.21 & 0 \\ 0 & 0 & 0.19 \end{bmatrix}. \end{split}$$

The selection of the transfer matrix with partially unknown probabilities is shown below:

$$\Pi = \begin{bmatrix} -0.8 & 0.5 & 0.3 \\ ? & -1.0 & ? \\ 0.8 & ? & ? \end{bmatrix};$$

based on the above transfer matrix, the process of switching between modes is shown in Figure 1.

The mode-dependent time-varying delays and related parameters are shown in Table 1.



Figure 1. Switching Modes of the system.

The parameters of the estimator gain perturbation are chosen as:

$$\begin{split} A_{11} &= \begin{bmatrix} 0.25 & 0 & 0 \\ 0 & 0.16 & 0 \\ 0 & 0 & 0.19 \end{bmatrix}, A_{12} = \begin{bmatrix} 0.31 & 0 & 0 \\ 0 & 0.18 & 0 \\ 0 & 0 & 0.23 \end{bmatrix}, A_{13} = \begin{bmatrix} 0.35 & 0 & 0 \\ 0 & 0.25 & 0 \\ 0 & 0 & 0.25 \end{bmatrix}, \\ A_{21} &= \begin{bmatrix} 0.33 & 0 & 0 \\ 0 & 0.26 & 0 \\ 0 & 0 & 0.18 \end{bmatrix}, A_{22} = \begin{bmatrix} 0.28 & 0 & 0 \\ 0 & 0.22 & 0 \\ 0 & 0 & 0.17 \end{bmatrix}, A_{23} = \begin{bmatrix} 0.26 & 0 & 0 \\ 0 & 0.19 & 0 \\ 0 & 0 & 0.23 \end{bmatrix}, \\ C_{11} &= \begin{bmatrix} 0.55 & 0 & 0 \\ 0 & 0.45 & 0 \\ 0 & 0 & 0.45 \end{bmatrix}, C_{12} = \begin{bmatrix} 0.35 & 0 & 0 \\ 0 & 0.31 & 0 \\ 0 & 0 & 0.31 \end{bmatrix}, C_{13} = \begin{bmatrix} 0.42 & 0 & 0 \\ 0 & 0.33 & 0 \\ 0 & 0 & 0.35 \end{bmatrix}, \\ C_{21} &= \begin{bmatrix} 0.45 & 0 & 0 \\ 0 & 0.35 & 0 \\ 0 & 0 & 0.35 \end{bmatrix}, C_{22} = \begin{bmatrix} 0.38 & 0 & 0 \\ 0 & 0.32 & 0 \\ 0 & 0 & 0.35 \end{bmatrix}, C_{23} = \begin{bmatrix} 0.51 & 0 & 0 \\ 0 & 0.43 & 0 \\ 0 & 0 & 0.45 \end{bmatrix}, \\ \Lambda_{1a}(x) = \Lambda_{2a}(x) = \begin{bmatrix} \sin x & 0 & 0 \\ 0 & \sin x & 0 \\ 0 & 0 & \sin x \end{bmatrix}, a = 1, 2, 3. \end{split}$$

Let $\gamma_1 = \gamma_2 = 0.5$, $\varpi = 1$, $\mathbb{N} = 0.65I_3$, and regulation function $f(\epsilon) = \frac{\epsilon^2}{1+\epsilon^2}$, by solving the LMIs in Theorem 2, it can prove that the LMIs (12) and (26) of this paper are feasible; the estimator gain matrices are derived:

$$\begin{split} K_{11} &= \begin{bmatrix} 0.9024 & -0.4021 & -0.1306 \\ -0.3080 & 2.1696 & -0.2721 \\ -0.1623 & -0.3567 & 1.1212 \end{bmatrix}, \ K_{21} = \begin{bmatrix} 0.0671 & -9.0706 & -5.7745 \\ -8.5686 & 0.0830 & -4.1928 \\ -5.7549 & -5.9144 & 0.0639 \end{bmatrix}, \\ K_{12} &= \begin{bmatrix} 0.9036 & -0.4430 & -0.1300 \\ -0.2972 & 2.2427 & -0.2630 \\ -0.1473 & -0.3743 & 1.0542 \end{bmatrix}, \ K_{22} = \begin{bmatrix} 0.0703 & -7.1541 & -3.5354 \\ -7.5068 & 0.1065 & -3.5237 \\ -3.7345 & -5.1003 & 0.1021 \end{bmatrix}, \\ K_{13} &= \begin{bmatrix} 0.5541 & -0.4042 & -0.1278 \\ -0.2766 & 1.6326 & -0.2426 \\ -0.1444 & -0.3375 & 0.7699 \end{bmatrix}, \ K_{23} = \begin{bmatrix} 0.0110 & -6.1614 & -4.1572 \\ -6.2536 & 0.0032 & -1.0914 \\ -5.7068 & -2.1649 & -0.0302 \end{bmatrix}. \end{split}$$

Based on the obtained estimator gain matrix, Figures 2-5 illustrate the simulation results we obtained. Figures 2 and 3 demonstrate the estimation errors for mRNA and protein, respectively. To observe the trend of



Figure 2. The evolution of mRNA estimation errors.



Figure 3. The evolution of protein estimation errors.

the estimation errors more clearly, we take the spatial point at y = 0.2 as the example and display the trajectory plots of the estimation errors of mRNA and protein in Figures 4 and 5, respectively. The above simulation results show that the estimation errors are asymptotically stable, thus proving the effectiveness of the non-fragile state estimator designed in this paper.



Figure 4. The trajectories of mRNA estimation errors ϕ_{mh_1} h = 1, 2, 3.



Figure 5. The trajectories of protein estimation errors ϕ_{ph} , h = 1, 2, 3.

5. CONCLUSIONS

This paper has investigated the non-fragile state estimation for Markovian jump RDGRNs with mode-dependent time-varying delays. Moreover, the switching topology of the system satisfies the Markov chain with partial unknown transfer probabilities. By utilizing the suitable LKFs, sufficient conditions that guarantee the asymptotic stability of the error system were established. Ultimately, a numerical example was used to demonstrate the validity of the designed non-fragile state estimator. Nonetheless, some limitations of this paper still exist. For example, the measurement methods adopted in this paper are costly and challenging to implement. Therefore, we will employ point measurements to reduce the measurement cost in future work.

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REFERENCES

- Pratap A, Raja R, Agarwal RP, Alzabut J, Niezabitowski M, Hincal E. Further results on asymptotic and finite-time stability analysis of fractional-order time-delayed genetic regulatory networks. *Neurocomputing* 2022;475:26-37. DOI
- Yu T, Zhao Y, Wang J, Liu J. Event-triggered sliding mode control for switched genetic regulatory networks with persistent dwell time. Nonlinear Anal Hybrid Syst 2022;44:101135. DOI
- 3. Yu T, Liu J, Zeng Y, Zhang X, Zeng Q, Wu L. Stability analysis of genetic regulatory networks with switching parameters and time delays. *IEEE Trans Neural Netw Learn Syst* 2017;29:3047-58. DOI
- Chaves M, Albert R, Sontag E. Robustness and fragility of Boolean models for genetic regulatory networks. J Theor Biol 2005;235:431-49. DOI
- 5. Li R, Yang M, Chu T. State feedback stabilization for Boolean control networks. IEEE Trans Automat Contr 2013;58:1853-7. DOI
- 6. Ali MS, Gunasekaran N, Ahn CK, Shi P. Sampled-data stabilization for fuzzy genetic regulatory networks with leakage delays. *IEEE/ACM Trans Comput Biol Bioinf* 2016;15:271-85. DOI
- Zhang X, Fan X, Wu L. Reduced-and full-order observers for delayed genetic regulatory networks. *IEEE Trans Cybern* 2017;48:1989-2000. DOI
- 8. Liu C, Wang X, Xue Y. Global exponential stability analysis of discrete-time genetic regulatory networks with time-varying discrete delays and unbounded distributed delays. *Neurocomputing* 2020;372:100-8. DOI
- 9. Yue D, Guan Z, Li J, Liu F, Xiao J, Ling G. Stability and bifurcation of delay-coupled genetic regulatory networks with hub structure. J Franklin Inst 2019;356:2847-69. DOI
- 10. Wang W, Dong, Y, Zhong S, Shi K, Liu F. Secondary delay-partition approach to finite-time stability analysis of delayed genetic regulatory networks with reaction-diffusion terms. *Neurocomputing* 2019;359:368-83. DOI
- 11. Song X, Li X, Ahn CK, Song S. Space-dividing-based cluster synchronization of reaction-diffusion genetic regulatory networks via intermittent control. *IEEE Trans NanoBiosci* 2021;21:55-64. DOI
- 12. Qin Y, Wang J, Chen X, Shi K, Shen H. Anti-disturbance synchronization of fuzzy genetic regulatory networks with reaction-diffusion. *J Franklin Inst* 2022;359:3733-48. DOI
- 13. Zhang Y, Liu H, Yan F, Zhou J. Oscillatory behaviors in genetic regulatory networks mediated by microRNA with time delays and reaction-diffusion terms. *IEEE Trans NanoBiosci* 2017;16:166-76. DOI
- 14. Xue Y, Zhang L, Zhang X. Reachable set estimation for genetic regulatory networks with time-varying delays and bounded disturbances. *Neurocomputing* 2020;403:203-10. DOI
- 15. Zou C, Wei X, Zhang Q, Zhou C. Passivity of reaction-diffusion genetic regulatory networks with time-varying delays. *Neural Process Lett* 2018;47:1115-32. DOI
- 16. Xue Y, Liu C, Zhang X. State bounding description and reachable set estimation for discrete-time genetic regulatory networks with time-varying delays and bounded disturbances. *IEEE Trans Syst Man Cybern Syst* 2022;52:6652-61 . DOI
- 17. Sun J, Wang H. Stability analysis for highly nonlinear switched stochastic systems with time-varying delays. *Complex Eng Syst* 2022;2:17. DOI
- 18. Tian J, Li Y, Zhao J, Zhong S. Delay-dependent stochastic stability criteria for Markovian jumping neural networks with mode-dependent time-varying delays and partially known transition rates. *Appl Math Comput* 2012;218:5769-81. DOI
- Hua M, Zheng D, Deng F, Fei J, Cheng P, Dai X. H_{oc} filtering for nonhomogeneous Markovian jump repeated scalar nonlinear systems with multiplicative noises and partially mode-dependent characterization. *IEEE Trans Syst Man Cybern Syst* 2021;51:3180-92. DOI
- 20. Hua M, Qian Y, Deng F, Fei J, Cheng P, Chen H. Filtering for discrete-Time Takagi-Sugeno fuzzy nonhomogeneous Markov jump systems with quantization effects. *IEEE Trans Cybern* 2022;52:982-95. DOI
- 21. Zhuang G, Ma Q, Zhang B, Xu S, Xia J. Admissibility and stabilization of stochastic singular Markovian jump systems with time delays.

Syst Control Lett 2018;114:1-10. DOI

- 22. Ma Y, Zhang Q, Li X. Dissipative control of Markovian jumping genetic regulatory networks with time-varying delays and reactiondiffusion driven by fractional Brownian motion. *Differ Equ Dyn Syst* 2020;28:841-64. DOI
- 23. Zou C, Wang X. Robust stability of delayed Markovian switching genetic regulatory networks with reaction-diffusion terms. *Comput Math Appl* 2020;79:1150-64. DOI
- 24. Chen G, Xia J, Park JH, Shen H, Zhuang G. Sampled-data synchronization of stochastic Markovian jump neural networks with timevarying delay. *IEEE Trans Neural Netw Learn Syst* 2022;33:3829-41. DOI
- 25. Shen H, Huo S, Yan H, Park JH, Sreeram V. Distributed dissipative state estimation for Markov jump genetic regulatory networks subject to round-robin scheduling. *IEEE Trans Neural Netw Learn Syst* 2019;31:762-71. DOI
- 26. Tao J, Xiao Z, Li Z, et al. Dynamic event-triggered state estimation for Markov jump neural networks with partially unknown probabilities. *IEEE Trans Neural Netw Learn Syst* 2021;33:7438-47. DOI
- 27. Tan Y, Liu Q, Liu J, Xie X, Fei S. Observer-based security control for interconnected semi-Markovian jump systems with unknown transition probabilities. *IEEE Trans Cybern* 2021;52:9013-25. DOI
- Qiu L, Zhang B, Xu G, Pan J, Yao F. Mixed H₂/H_∞ control of Markovian jump time-delay systems with uncertain transition probabilities. Inf Sci 2016;373:539-56. DOI
- 29. Song X, Wang M, Song S, Ahn CK. Sampled-data state estimation of reaction diffusion genetic regulatory networks via space-dividing approaches. *IEEE/ACM Trans Comput Biol Bioinf* 2019;18:718-30. DOI
- Zou C, Zhou C, Zhang Q, He X, Huang C. State estimation for delayed genetic regulatory networks with reaction diffusion terms and Markovian jump. *Complex Intell Syst* 2023;9:5297-311. DOI
- Sun L, Wang J, Chen X, Shi K, Shen H. H₁₀ fuzzy state estimation for delayed genetic regulatory networks with random gain fluctuations and reaction-diffusion. J Franklin Inst 2021;358:8694–714. DOI
- Li J, Dong H, Liu H, Han F. Sampled-data non-fragile state estimation for delayed genetic regulatory networks under stochastically switching sampling periods. *Neurocomputing* 2021;463:168–76. DOI
- Wan X, Xu L, Fang H, Ling G. Robust non-fragile H_∞ state estimation for discrete-time genetic regulatory networks with Markov jump delays and uncertain transition probabilities. *Neurocomputing* 2015;154:162-73. DOI
- 34. Ali MS, Agalya R, Hong KS. Non-fragile synchronization of genetic regulatory networks with randomly occurring controller gain fluctuation. *Chinese J Phys* 2019;62:132–43. DOI
- 35. Wu FX. Delay-independent stability of genetic regulatory networks. IEEE Trans Neural Netw 2011;22:1685-93. DOI
- 36. Polynikis A, Hogan SJ, Di Bernardo M. Comparing different ODE modelling approaches for gene regulatory networks. *J Theor Biol* 2009;261:511-30. DOI
- 37. Lipan O, Ferwerda C. Hill functions for stochastic gene regulatory networks from master equations with split nodes and time-scale separation. *Phys Rev E* 2018;97:022413. DOI
- Wang JW, Wu HN. Some extended Wirtinger's inequalities and distributed proportional-spatial integral control of distributed parameter systems with multi-time delays. J Franklin Inst 2015;352:4423-45. DOI
- 39. Zhou K, Khargonekar PP. Robust stabilization of linear systems with norm-bounded time-varying uncertainty. *Syst Control Lett* 1988;10:17-20. DOI
- 40. Petersen IR, Hollot CV. A riccati equation approach to the stabilization of uncertain linear systems. Automatica 1986;22:397-411. DOI