

**DNN PROJECTIONAL OBSERVER FOR  
ADVANCED OZONATION SYSTEMS OF  
COMPLEX CONTAMINANTS MIXTURES**

**Olga Andrianova \* Tatyana Poznyak \*\*  
Alexander Poznyak \*\*\* Isaac Chairez \*\*\*\***

*\* Higher School of Economics and ICS RAS, Russia*

*\*\* SEPI, ESQIE, Instituto Politecnico Nacional, Mexico*

*\*\*\* Automatic Control Department, CINVESTAV-IPN,  
Mexico*

*\*\*\*\* Bioprocesses Department, UPIBI, Instituto Politecnico  
Nacional, Mexico*

Abstract: The aim of this study is to provide a class of state observers, based on differential neural networks, to approximate a class of advanced oxidation systems, based on the application of ozone high oxidant power and catalyst (the named catalytic ozonation). The study considers the design of a state observer for uncertain systems with the restrictions of the ozonation system, including the positivity of the states, as well as the control action. The observer includes a projection operator which is motivated by the state constraints. The learning laws of the proposed differential neural networks are obtained using a class of controlled state restricted Lyapunov functions. The detailed stability analysis proves the input to state stability with respect to the modeling error, as well as the bounded uncertainties of the ozonation system. The experimental confirmation of the state estimation is also presented. The experimental case considers the ozonation of a toxic organic contaminant (terephthalic acid) which is a regular pollutant of the plastic industry wastewater.

Keywords: Catalytic ozonation; Adaptive state observer, Differential neural network; State restrictions; Projection operators.

## INTRODUCTION

There is a wide variety of inorganic materials which have been tested as catalysts in heterogeneous catalytic ozonation systems. Such material include Titanium oxide ( $TiO_2$ ), supported metals ( $Ni$ ,  $Mo$ ,  $Ce$ , etc), and unsupported or supported metal oxides, which are regular catalysts. In the last few decades, the catalytic ozonation reaction has been explained by two main reaction mechanisms (Rodríguez *et al.* 2013). The first of these mechanisms assumes that organics, reacting with ozone, are adsorbed on the catalyst surface and

then, decomposed by either ozone or the produced hydroxyl radicals (Guan *et al.* 2013). The second of the proposed mechanisms supposes that both ozone and organic compounds are simultaneously adsorbed on the catalyst surface, and then the organic derivative products (formed by the redox process) are oxidized by both ozone and the produced hydroxyl radicals (Nawrocki and Kasprzyk-Hordern 2010). In consequence, the catalytic ozonation is considered as a non selective process; however, the chemical structure of the initial compound is a main factor which may

define its interactions with the catalyst along the ozonation. With the aim of suggesting a suitable method to apply the catalytic ozonation system in real treatment plants, it is necessary to optimize the use of the catalyst (Wang and Bai 2017).

One of the possible options to justify the application of the catalytic ozonation is supporting the catalyst on solid inert support forming thin films (Li *et al.* 2003). These catalytic films do not require filtering the liquid and may preserve the catalyst in the reactor and promote its re-use for several reaction rounds. Catalytic ozonation can be considered as a promising method to decompose toxic organics, which cannot be decomposed by other methods. However, the enhancement of the catalytic ozonation efficiency requires the application of diverse strategies, including the adjustment of the operating conditions for the ozonation system. For exerting such optimization, it may be necessary to use a suitable model of the catalytic ozonation system. The development of such a model is a complex task considering the high number of reactions between the organic compounds, ozone, the catalyst and the formed byproducts. An alternative is using adaptive approximate representations of the ozonation model.

Differential neural networks (DNNs) have proven to be efficient approximative representations of systems with uncertain models (Poznyak *et al.* 2001, Wu 1998). DNNs, also known as Auto Associative or Feedback Networks, define a class of artificial neural networks, where the connections between units and output form a directed cycle. This creates an internal state of the network, yielding the exhibition of dynamic temporal-dependent behavior (Chairez 2009). DNNs can use their internal memory to process arbitrary sequences of inputs. In DNN, the signal travels in both forward and backward directions, by introducing loops in the network architecture or topology. The potential application of DNN, to obtain approximate models of the catalytic ozonation, must consider several challenging aspects, such as the limited available information, to obtain the approximate model, the positive nature of all the states, which describe the catalytic ozonation system, and the large number of states needed to obtain a usable DNN-based model of the system.

The main contributions of this study are: a) the development of an approximate model for a class of catalytic ozonation systems, considering that the catalyst is supported in solid supports and forming thin films, b) the development of a software sensor, to approximate the catalytic ozonation dynamic behavior, considering the positiveness of all the states, and c) an experimental confirmation of the software sensor application to recover the approximate model of the catalytic

ozonation of 4-chlorophenol (4-*CPh*) and 4-phenolsulfonic acid (4-*PSA*) using  $CeO_2$  as the catalyst. The rest of this manuscript is organized as follows. Section II presents the development of an approximate model of the catalytic ozonation system including the design of the DNN based state observer. Section III describes the design of the learning law which is applicable to the class of developed software sensors. Section IV details the application of the DNN state observer to reproduce the dynamics of a catalytic ozonation of 4-*CPh* as well as 4-*PSA*. Section V closes the study with some final remarks.

## 1. ADAPTIVE STATE OBSERVER FOR OZONATION SYSTEMS

This section describes the design of the suggested state observer. The system to be approximated is described by the vector of variables  $x = [c^g, Q, OH, c_1, c_{11}, \dots, c_{1n}]$ , where the sense of all states is defined as follows: the catalytic ozonation model may include two sections: a) the first one formed by a basic nominal model, that contains the mass-transfer of ozone and its reaction with contaminants, and b) an uncertain section that includes all the non-modeled reactions taking place. The ozone dissolved in the aqueous phase ( $Q$ , *mole*) is transferred from the gaseous phase (with a given ozone concentration  $c^g$ ), obeying the double layer theory. This  $Q$  is consumed by the reaction with the contaminants, as well as the corresponding by-products and it is also used to produce hydroxyl radicals. The initial contaminant concentration is represented by  $c_1$ , *mole* ·  $L^{-1}$  and its byproducts concentrations are given by  $(c_{1i}, \text{mole} \cdot L^{-1}, i \in [1, N])$ . All these organics are decomposed by the reaction with the dissolved ozone as well as with the hydroxyl radicals ( $OH$ , *mole* ·  $L^{-1}$ ). These radicals are produced from the self-decomposition of dissolved ozone, as well as by the action of the catalyst.

With the aim of providing a suitable approximate model of the catalytic ozonation system, let's propose a general class of dynamic systems with uncertain dynamics that can represent the time variation of  $x$ .

### 1.1 Uncertain dynamic system

Assume that the catalytic ozonation system can be represented by an uncertain non-negative ODE given by:

$$\frac{d}{dt}x(t) = f(x(t), u(t), t), \quad x(0) = x_0, \quad \|x_0\| < \infty. \quad (1)$$

The output  $y = c^g$  is described by

$$y(t) = Cx(t) + \nu(t), \quad (2)$$

then  $C = [1 \ 0 \ \dots \ 0]$ . In view of the nature of the variables in  $x$ , the state has non-negative bounded components

$$0 \leq x_i \leq x_i^+, \quad i = 1, \dots, n, \quad (x^+)^2 := \sum_{i=1}^n (x_i^+)^2. \quad (3)$$

The variable  $u(t) \in U_{adm} \subset \mathbb{R}^m$ ,  $m < n$  may describe a possible control action, which must satisfy boundedness conditions on state components (3). Variable  $\nu(t) \in \mathbb{R}^g$  defines bounded noises affecting the output.

### 1.2 Main assumptions

The principal assumptions considered in this study are:

- (1) The dimension of the state  $n$  is known.
- (2) The admissible controls  $u(t)$  and noises  $\nu(t)$  are within the following bounds (the bounds  $u_+$  and  $\nu_+$  are known), i.e.,

$$U_{adm} = \{u : \|u\|^2 \leq u_+^2 < \infty\} \quad (4)$$

and

$$\|\nu(t)\| \leq \nu_+ < \infty. \quad (5)$$

- (3) The mapping  $f : \mathbb{R}^n \times \mathbb{R}^m \times \mathbb{R} \rightarrow \mathbb{R}^n$  is unknown, but locally Lipschitz.
- (4) The upper bounds  $x_i^+$  in (3) are known and constant.
- (5) There is a Hurwitz matrix  $A \in \mathbb{R}^{n \times n}$  (maybe unknown) such that  $\forall x \in \mathbb{R}^n$ , all  $u(t) \in U_{adm}$  and all  $t \geq 0$  the following property holds (satisfying (3)):

$$\|\delta_A f(x, u, t)\|^2 \leq \theta, \quad (6)$$

where

$$\delta_A f(x, u, t) := f(x, u, t) - Ax. \quad (7)$$

## 2. DNN OBSERVER FOR POSITIVE UNCERTAIN SYSTEMS

Consider the DNN observer (DNNO) satisfying the following structure:<sup>1</sup>

$$\left. \begin{aligned} \frac{d}{dt} \bar{x}(t) &= A\hat{x}(t) + W_1(t)\Psi_1(\hat{x}(t)) + \\ &W_2(t)\Psi_2(\hat{x}(t))u(t) + L[y(t) - C\hat{x}(t)], \\ \hat{x}(t) &= \Pi_X \{\bar{x}(t)\}. \end{aligned} \right\} \quad (8)$$

Here  $\Pi_X \{\bar{x}\}$  defines the projection operator as

$$\begin{aligned} \Pi_X \{\bar{x}\} &= \{\text{sat}_1(\bar{x}_1), \dots, \text{sat}_n(\bar{x}_n)\}, \\ \text{sat}_i(\bar{x}_i) &= \begin{cases} 0 & \text{if } \bar{x}_i < 0 \\ \bar{x}_i & \text{if } \bar{x}_i \in [0, x_i^+] \\ x_i^+ & \text{if } \bar{x}_i > x_i^+. \end{cases} \end{aligned} \quad (9)$$

The vector  $\hat{x}(t) \in \mathbb{R}^n$  is the state estimation vector,  $\bar{x}(t) \in \mathbb{R}^n$  is an intermediate auxiliary variable,  $W_1(t) \in \mathbb{R}^{n \times l}$  and  $W_2(t) \in \mathbb{R}^{n \times s}$  are the time varying weighting matrices which must be adjusted during the training procedure,  $\Psi_1(x) \in \mathbb{R}^l$  and  $\Psi_2(x) \in \mathbb{R}^{s \times m}$  are given activation vectorial and matrix functions. These activations have sigmoidal components, which satisfy the following structure:

$$\left. \begin{aligned} \Psi_{1,r}(x) &= \frac{a_r}{1 + b_r e^{(-c_r^\top x)}} \\ \Psi_{2,rp}(x) &= \frac{a_{rp}}{1 + b_{rp} e^{(-c_{rp}^\top x)}} \\ a_r, a_{rp}, b_r, b_{rp} &> 0, \\ c_r \in \mathbb{R}, c_{rp} \in \mathbb{R}^n \end{aligned} \right\} \quad (10)$$

Evidently,  $\Psi_1(x)$  and  $\Psi_2(x)$  are bounded:

$$\|\Psi_1(x)\| \leq L_{\Psi_1}^+, \quad \|\Psi_2(x)u\| \leq Lu_+.$$

Notice that, in view of the projection operator property (9), vectors  $\hat{x}(t)$  and  $\bar{x}(t)$  satisfy the inequality

$$\|\hat{x}(t) - x(t)\| \leq \|\bar{x}(t) - x(t)\|. \quad (11)$$

## 3. LEARNING LAWS

Assume that the right-hand side of the system with uncertain dynamics (1) can be represented as

$$\frac{d}{dt}x(t) = Ax(t) + \hat{W}_1\Psi_1(x(t)) + \hat{W}_2\Psi_2(x(t))u(t),$$

let's consider the following learning laws for the DNNO weights on-line adjustment:

$$\begin{aligned} \dot{W}_1(t) &= -\frac{\alpha}{2}\tilde{W}_1(t) - \\ &\frac{1}{k_{w,1}}P(C^\top C + \delta I)^{-1}C^\top e\Psi_1^\top(\hat{x}(t)) \\ &- \frac{1}{2k_{w,1}}P\Lambda_w^{-1}P\tilde{W}_1(t)\Psi_1(\hat{x}(t))\Psi_1^\top(\hat{x}(t)) \\ \dot{W}_2(t) &= -\frac{\alpha}{2}\tilde{W}_2(t) - \\ &\frac{1}{k_{w,2}}P(C^\top C + \delta I)^{-1}C^\top e(t)u^\top(t)\Psi_2^\top(\hat{x}(t)) \\ &- \frac{1}{2k_{w,2}}P\Lambda_w^{-1}P\tilde{W}_2(t)\Psi_2(\hat{x}(t))u(t)u^\top(t)\Psi_2^\top(\hat{x}(t)). \end{aligned} \quad (12)$$

Here  $\tilde{W}_i(t) := W_i(t) - \hat{W}_i$  ( $i = 1, 2$ );  $\alpha, k_{w,1}, k_{w,2}, \delta$  are positive parameters,  $\Lambda_w \in \mathbb{R}^{n \times n}$  is any positive definite matrix, and

$$e(t) := C\bar{x}(t) - y(t). \quad (13)$$

Matrices  $\hat{W}_1, \hat{W}_2, L, P$ , as well as the parameters  $\alpha, \delta$ , participating in (12), will be obtained using

<sup>1</sup> The similar DNNO structure without the projection component was suggested in (Poznyak *et al.* 2001) and (Poznyak *et al.* 2019).

an optimization procedure based on the theory of matrix inequalities.

#### 4. MAIN RESULT ON THE QUALITY OF THE SUGGESTED PROJECTIONAL DNNO

*Theorem 1.* Consider the projectional DNNO (8) supplied by Learning Laws (12), where all matrix, vector and scalar parameters are fixed. If, for the given DNNO, there exist positive definite (symmetric) matrices  $P \in \mathbb{R}^{n \times n}$ ,  $L \in \mathbb{R}^{n \times q}$ ,  $A \in \mathbb{R}^{n \times n}$ ,  $\hat{W}_1 \in \mathbb{R}^{n \times l}$ ,  $\hat{W}_2 \in \mathbb{R}^{n \times s}$ , and a positive scalar parameter  $\varepsilon$  such that the following matrix inequality holds:

$$H \left( P, L, A, \hat{W}_1, \hat{W}_2 \mid \alpha, \varepsilon, \delta \right) := \begin{pmatrix} H_{11} & -P \\ -P & -\varepsilon I_{n \times n} \end{pmatrix} < 0, \\ H_{11} = P \left( A - LC + \frac{\alpha}{2} I \right) + \left( A - LC + \frac{\alpha}{2} I \right)^\top P + \\ P \left[ (A - LC) \Lambda_x^{-1} (A - LC)^\top + L \Lambda_L^{-1} L^\top \right] P + \\ P \left( \hat{W}_1 \Lambda_{\hat{w}_1}^{-1} \hat{W}_1^\top + \hat{W}_2 \Lambda_{\hat{w}_2}^{-1} \hat{W}_2^\top \right) P + \\ 2 \|\Lambda_x\| I_{n \times n} + \delta^2 (C^\top C + \delta I)^{-1} \Lambda_w (C^\top C + \delta I)^{-1}, \quad (14)$$

then all possible trajectories of the state estimation error  $\hat{\Delta}(t) := \hat{x}(t) - x(t)$  asymptotically arrive (approach) to the centered attractive ellipsoid  $\mathcal{E}(P_{attr})$  with the ellipsoidal matrix  $P_{attr} \in \mathbb{R}^{n \times n}$ , fulfilling

$$\limsup_{t \rightarrow \infty} \hat{\Delta}^\top(t) P_{attr} \hat{\Delta}(t) \leq 1 \quad (15)$$

where

$$P_{attr} = \frac{\alpha}{\beta} P \quad (16)$$

with

$$\beta = 2 \|\Lambda_x\| \left( \sum_{i=1}^n (x_i^+)^2 + (x^+)^2 \right) + \|\Lambda_L\| \nu_+^2 + \varepsilon \theta + \\ \|\Lambda_{\hat{w}_1}\| (L_{\Psi_1}^+)^2 + \|\Lambda_{\hat{w}_2}\| (L_{\Psi_2}^+ u_+)^2 + \\ \left\| C (C^\top C + \delta I)^{-1} \Lambda_w (C^\top C + \delta I)^{-1} C^\top \right\| (\nu_+)^2. \quad (17)$$

Here matrices  $\Lambda_x \in \mathbb{R}^{n \times n}$ ,  $\Lambda_L \in \mathbb{R}^{q \times q}$ ,  $\Lambda_w \in \mathbb{R}^{n \times n}$ ,  $\Lambda_{\hat{w}_1} \in \mathbb{R}^{l \times l}$ ,  $\Lambda_{\hat{w}_2} \in \mathbb{R}^{s \times s}$  are known. A brief summary of the proof of this theorem is given in Appendix.

## 5. EXPERIMENTAL EVALUATIONS

### 5.1 Materials

The compounds  $Ce(NO_3)_3 \times 6H_2O$  (99%), 4-CPH (99%) and 4-PSA (99%) were used without further purification (Sigma-Aldrich). All the other reagents were analytical grade (J. T. Baker). The fixed initial concentration of all the contaminants (4-CPH and 4-PSA) was  $100 \text{ mgL}^{-1}$  and initial pH was fixed to 3.5. The pH parameter was not controlled during the ozonation.

### 5.2 Preparation of CeO<sub>2</sub> film

The selected catalyst ( $CeO_2$ ) was deposited over clean glass slides as films. The supported catalyst, namely  $CeO_2(f)$ , was synthesized using an ultrasonic spray pyrolysis technique. The procedure initiated with a solution of cerium nitrate ( $0.6 \text{ gL}^{-1}$ ) which operated as precursor. The films of  $CeO_2(f)$  were made over clean glass slides (LAUKA,  $26.4 \times 76.2 \text{ mm}$ ).

### 5.3 Catalyst evaluation on contaminant decomposition

Ozonation experiments for compounds 4-CPH and 4-PSA were carried out in a glass reactor (500 mL) with a porous filter at the bottom, which served as the ozone diffuser to the liquid phase. Ozone was produced using an HTU500G generator fed with ultra-dried oxygen. The  $O_2 - O_3$  gas mixture flow ( $0.5 \text{ Lmin}^{-1}$ ) was limited by a mechanical rotameter. The input ozone concentration was ( $25 \text{ mgL}^{-1}$ ). The output ozone concentration was measured with a BMT 964 BT analyzer. In the experiments with the catalyst, the glass slides, coated with the  $CeO_2$  film (an average of 1.64 mg in each film), were fixed in the reactor using a teflon support.

### 5.4 Analytical methods

The 4-CPH and 4-PSA concentrations were determined by high performance liquid chromatography (HPLC, Perkin Elmer, Flexar) with an array-diode detector. The C18 column (Nova Pack C18) was used in an isocratic mode with a mobile phase of water: acetonitrile (50 : 50 and 90 : 10 for 4-CPH and 4-PSA, respectively) at  $0.5 \text{ mL} \times \text{min}^{-1}$ . In the case of oxalic acid, prevail organic acid column was used with  $KH_2PO_4$  buffer solution at pH 2.5 ( $1.0 \text{ mL min}^{-1}$ ). All the experiments were done in triplicate.

### 5.5 Estimation results

The observer was applied considering that the ozone concentration at the reactor's output was the on-line available measure. The reconstructed information included the variation of the main contaminant. The application of the observer can be extended to include the variation of the byproducts' concentrations formed during ozonation, the dissolved oxygen in the aqueous phase in the reactor, as well as the concentration of hydroxyl radicals. In this study, we presented the reconstruction of the contaminants and their byproducts concentrations (of those identified

compounds), which were considered the most valuable information for the practitioners.

The parameters used for the projected DNN-observer evaluation were

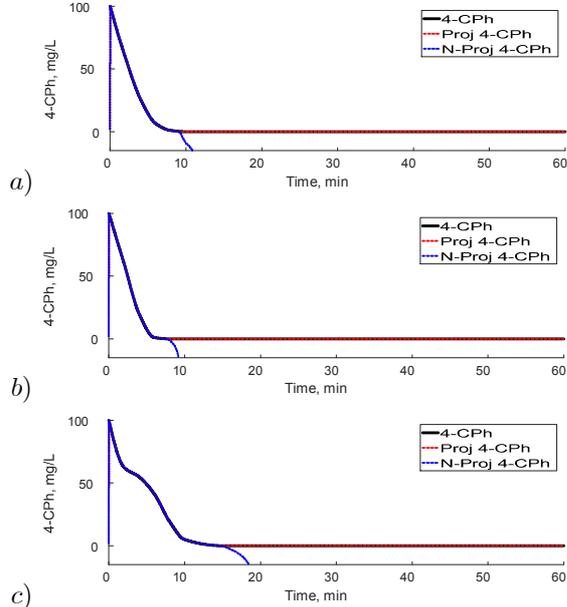
$$A = \text{diag} \{ [-0.25 \ -0.34 \ -1.24 \ -1.26] \},$$

$$C = [1 \ 0 \ 0 \ 0]^T$$

$$P = \begin{bmatrix} 9.2 & 1.2 & -2.5 & -2.9 \\ 1.2 & 18.7 & 0.5 & 0.9 \\ -2.5 & 0.5 & 25.2 & -0.7 \\ -2.9 & 0.9 & -0.7 & 34.6 \end{bmatrix}$$

$$L = [26.1 \ 15.7 \ 22.4 \ 18.2].$$

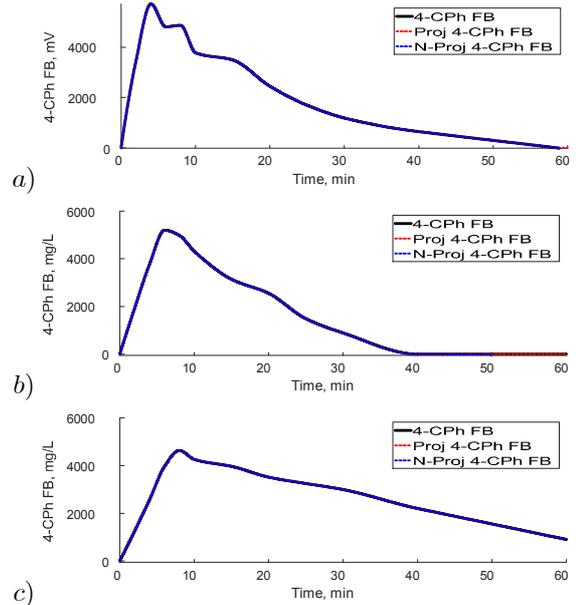
For calculation of the matrix  $P$  the following parameters  $\alpha = 0.25$  and  $\beta = 1.62 \times 10^{-2}$  were used. *Figure 1* depicts the comparison of the 4-CP and 4-PSA concentrations obtained with the conventional, suspended and films-based catalytic ozonations, its non-projected estimation by the DNNO ( $\hat{x}$ ) as well as the projected state ( $\hat{x}$ ) are presented. This comparison does have a significant difference between the projected and non-projected state because the non-projected form yields the incorrect estimation of a negative concentration for the main contaminant.



*Figure 1.* Comparison of the estimated concentrations of 4-CP estimated with the projected and non-projected observer.

a) Conventional, b)  $CeO_2(s)$ , b)  $CeO_2(f)$

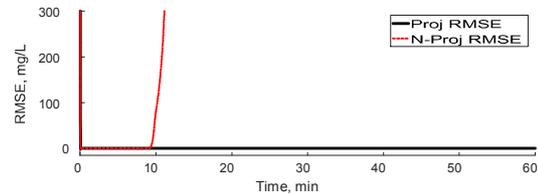
The comparison of the byproduct's concentration formation and decomposition confirms the benefits of introducing the projection operator, because the estimated concentration is forced to be non-negative ( $\hat{x}$ ) while the non-projected state detours to negative values (*Figure 2*).



*Figure 2.* Comparison of the estimated concentrations of the first byproduct obtained from the 4-CP estimated with the projected and non-projected observer.

a) Conventional, b)  $CeO_2(s)$ , b)  $CeO_2(f)$

The estimation of the root mean square error (RMSE) confirmed the significant differences of introducing the projection operator over the DNN observer. The magnitude of the RMSE for the non-projected case was 2 orders larger than the case for the projected case (*Figure 3*).



*Figure 3.* Comparison of the estimated RMSE for the projected and non-projected observers.

## 6. CONCLUSIONS

This study considers the design of a new learning law for DNN, based on the application of a new variant of a projectional algorithm to estimate the states of a complex system, describing the ozonation of the toxic contaminant under the presence of the catalyst in two diverse variants: suspended and supported on thin films. The explicit idea of how to implement the projectional method in the continuous-time NN for estimating the variables of a positive system is described. The strict analysis for obtaining the upper bound of the method and to justify the asymptotic convergence of the estimation error is provided. The proposed method opens new researching opportunities to design a wide diversity of NN with a clear motivation on real plants.

REFERENCES

- Chairez, Isaac (2009). Wavelet differential neural network observer. *IEEE Transactions on Neural Networks* **20**(9), 1439–1449.
- Guan, Ying-Hong, Jun Ma, Yue-Ming Ren, Yu-Lei Liu, Jia-Yue Xiao, Ling-qiang Lin and Chen Zhang (2013). Efficient degradation of atrazine by magnetic porous copper ferrite catalyzed peroxymonosulfate oxidation via the formation of hydroxyl and sulfate radicals. *Water research* **47**(14), 5431–5438.
- Li, Laisheng, Wanpeng Zhu, Pengyi Zhang, Zhongyin Chen and Wenya Han (2003). Photocatalytic oxidation and ozonation of catechol over carbon-black-modified nano-tio2 thin films supported on al sheet. *Water Research* **37**(15), 3646–3651.
- Nawrocki, Jacek and Barbara Kasprzyk-Hordern (2010). The efficiency and mechanisms of catalytic ozonation. *Applied Catalysis B: Environmental* **99**(1-2), 27–42.
- Poznyak, A., E. Sanchez and W. Yu (2001). *Differential Neural Networks for Robust Nonlinear Control (Identification, State Estimation and Trajectory Tracking)*. World Scientific.
- Poznyak, AS (2009). *Advanced Mathematical Tools for Automatic Control Engineers. Vol. I. Deterministic Systems..* Springer.
- Poznyak, T., I. Chairez and A. Poznyak (2019). *Ozonation and Biodegradation in Environmental Engineering: Dynamic Neural Network Approach*. 1st ed.. Elsevier.
- Rodríguez, Julia L, Tatiana Poznyak, Miguel A Valenzuela, Hugo Tiznado and Isaac Chairez (2013). Surface interactions and mechanistic studies of 2, 4-dichlorophenoxyacetic acid degradation by catalytic ozonation in presence of ni/tio2. *Chemical engineering journal* **222**, 426–434.
- Wang, Jianlong and Zhiyong Bai (2017). Fe-based catalysts for heterogeneous catalytic ozonation of emerging contaminants in water and wastewater. *Chemical Engineering Journal* **312**, 79–98.
- Wu, Jianhong (1998). Symmetric functional differential equations and neural networks with memory. *Transactions of the American mathematical society* **350**(12), 4799–4838.

APPENDIX

**Proof.** Consider the Lyapunov-like function

$$V_P(t) = \|\bar{x}(t) - x(t)\|_P^2 + \sum_{i=1}^2 k_{w,i} \text{tr} \left\{ \tilde{W}_i^T(t) \tilde{W}_i(t) \right\} \quad (18)$$

and define the vectors  $\Delta := \bar{x}(t) - x(t)$  and  $e(t) = C\bar{x}(t) - y(t)$  for which we have

$$\begin{aligned} e(t) &= C\bar{x}(t) - y(t) = C\Delta(t) - \nu(t), \\ \Delta(t) &= (C^T C + \delta I)^{-1} C^T e(t) + w(t), \\ w(t) &:= (C^T C + \delta I)^{-1} [\delta I \Delta + C^T \nu(t)]. \end{aligned} \quad (19)$$

Differentiation of (18) on the trajectories of the system gives

$$\begin{aligned} \dot{V}_P(t) &= S(t) + S_1(t) + S_{11}(t) + \\ &+ 2[\bar{x}(t) - x(t)]^T P[(A - LC)(\bar{x}(t) - x(t))] \\ &- 2[\bar{x}(t) - x(t)]^T P \delta_A f(x, u, t) \end{aligned}$$

where

$$\begin{aligned} S(t) &:= 2 \sum_{i=1}^2 k_{w,i} \text{tr} \left\{ \tilde{W}_i^T(t) \dot{W}_i(t) \right\} + \\ &2 \left[ e^T C (C^T C + \delta I)^{-1} + w^T(t) \right] P W_1(t) \Psi_1(\hat{x}(t)), \\ &2 \left[ e^T C (C^T C + \delta I)^{-1} + w^T(t) \right] P W_2(t) \Psi_2(\hat{x}(t)) u(t) \\ S_1(t) &:= 2[\bar{x}(t) - x(t)]^T P (A - LC) (\hat{x}(t) - \bar{x}(t)), \\ S_{11}(t) &:= 2[\bar{x}(t) - x(t)]^T P L \nu(t). \end{aligned}$$

In view of the  $\Lambda$ -inequality (see Lemma 12.1 in (Poznyak 2009)) the following upper estimates for the derivative of the Lyapunov-like function holds:

$$\begin{aligned} \dot{V}_P(t) &\leq \varpi^T \tilde{H}(P, L, A | \alpha, \varepsilon) \varpi - \\ &\alpha \|\bar{x}(t) - x(t)\|_P^2 + S(t) + \beta_2, \\ \beta_1 &:= 2 \|\Lambda_x\| \left( \sum_{i=1}^n (x_i^+)^2 + (x^+)^2 \right) + \\ \|\Lambda_L\| \nu_+^2, \quad \beta_2 &:= \beta_1 + \varepsilon \theta, \\ \varpi &= \begin{bmatrix} \bar{x}(t) - x(t) \\ \delta_A f(x, u, t) \end{bmatrix} \end{aligned}$$

with  $\tilde{H}(P, L, A | \alpha, \varepsilon) := \begin{pmatrix} \tilde{H}_{11} & -P \\ -P & -\varepsilon I_{n \times n} \end{pmatrix}$ . Here

$$\begin{aligned} \tilde{H}_{11} &= P(A - LC) + (A - LC)^T P + \\ &P(A - LC) \Lambda_x^{-1} (A - LC)^T P + \alpha P + \\ &2 \|\Lambda_x\| I_{n \times n} + P L \Lambda_L^{-1} L^T P \end{aligned}$$

. In view of Learning Laws (12)

$$\begin{aligned} &\text{tr} \left\{ 2 \tilde{W}_1^T(t) P (C^T C + \delta I)^{-1} C^T e \Psi_1^T(\hat{x}(t)) \right\} + \\ &\text{tr} \left\{ 2 \tilde{W}_2^T(t) P (C^T C + \delta I)^{-1} C^T e u^T(t) \Psi_2^T(\hat{x}(t)) \right\} + \\ &\text{tr} \left\{ \tilde{W}_1^T(t) P \Lambda_w^{-1} P \tilde{W}_1(t) \Psi_1(\hat{x}(t)) \Psi_1^T(\hat{x}(t)) \right\} + \\ &\text{tr} \left\{ \tilde{W}_2^T(t) P \Lambda_w^{-1} P \tilde{W}_2(t) \Psi_2(\hat{x}(t)) u(t) \cdot \right. \\ &\left. u^T(t) \Psi_2^T(\hat{x}(t)) \right\} + 2 \sum_{i=1}^2 k_{w,i} \text{tr} \left\{ \tilde{W}_i^T(t) \dot{W}_i(t) \right\} + \\ &\alpha \sum_{i=1}^2 k_{w,i} \text{tr} \left\{ \tilde{W}_i^T(t) \tilde{W}_i(t) \right\} = 0 \end{aligned}$$

and in view of the property (14), one gets  $\dot{V}_P(t) \leq -\alpha V_P(t) + \beta$  implying

$$V_P(t) \leq \left[ V_P(0) + \frac{\beta}{\alpha} \right] e^{-\alpha t} + \frac{\beta}{\alpha}.$$

By (11),

$$\begin{aligned} \limsup_{t \rightarrow \infty} \|\hat{x}(t) - x(t)\|_P^2 &\leq \limsup_{t \rightarrow \infty} \|\bar{x}(t) - x(t)\|_P^2 \\ &\leq \limsup_{t \rightarrow \infty} V_P(t) \leq \frac{\beta}{\alpha} \end{aligned}$$

which is equivalent to (15). ■