

**Effects of stochastic time-delayed feedback on a dynamical system modeling a chemical oscillator**

Héctor O. González Ochoa\* and Gualberto Solís Perales

*Departamento de Electrónica, Universidad de Guadalajara. Av. Revolución 1500, 44430, Guadalajara Jal, México*

Irving R. Epstein

*Department of Chemistry, Brandeis University, Waltham, Massachusetts 02454-9110, USA*

Ricardo Femat

*Instituto Potosino de Investigación Científica y Tecnológica A.C., San Luis Potosí, México*

(Received 10 January 2018; published 23 May 2018)

We examine how stochastic time-delayed negative feedback affects the dynamical behavior of a model oscillatory reaction. We apply constant and stochastic time-delayed negative feedbacks to a point Field-Körös-Noyes photosensitive oscillator and compare their effects. Negative feedback is applied in the form of simulated inhibitory electromagnetic radiation with an intensity proportional to the concentration of oxidized light-sensitive catalyst in the oscillator. We first characterize the system under nondelayed inhibitory feedback; then we explore and compare the effects of constant (deterministic) versus stochastic time-delayed feedback. We find that the oscillatory amplitude, frequency, and waveform are essentially preserved when low-dispersion stochastic delayed feedback is used, whereas small but measurable changes appear when a large dispersion is applied.

DOI: [10.1103/PhysRevE.97.052214](https://doi.org/10.1103/PhysRevE.97.052214)**I. INTRODUCTION**

Oscillatory chemical reactions show complex nonlinear phenomena with a rich diversity of patterns, many resembling those found in biological processes. For instance, striking similarities can be noted in wave patterns displayed by the Belousov-Zhabotinsky (BZ) reaction [1–5], starving amoeba *Dictyostelium discoideum* [6], and calcium waves in *Xenopus laevis* oocytes [7,8]. Turing was the first to propose reaction-diffusion as a possible underlying mechanism in the process of morphogenesis [9]. The structures predicted by Turing have been experimentally observed in both continuous [10] and cell-compartmentalized chemical systems [11]. Chemical oscillators are reaction-diffusion systems, which have been proposed and used as model systems for elucidating complex biomimetic dynamics [11–14]. The BZ reaction is the prototypical chemical oscillator in such investigations. The reaction, which occurs in acidic aqueous solution, consists of the oscillatory oxidation by bromate of an organic substrate, usually malonic acid, catalyzed by metal ions or metal complexes. A photosensitive version of the BZ reaction, which uses the light-sensitive complex  $\text{Ru}(\text{bpy})_3^+$  as the metal catalyst, can be modulated by the application of an inhibiting electromagnetic field [15,16]. Field, Körös, and Noyes performed, on the grounds of chemical kinetics, a detailed analysis of the BZ reaction and developed a mechanism, the Field-Körös-Noyes (FKN) mechanism, to account for its oscillatory behavior [17]. The FKN mechanism identifies 11 principal reactions and 12 chemical species and successfully reproduces the behavior of

the BZ reaction [18]. Later, the model was extended to include photoinhibition [16,19,20].

The application of feedback to a nonlinear dynamical system can induce complex behavior. Feedback can also be used as a control mechanism to drive the system toward a desired dynamics. Different feedback strategies have been explored. Time-delayed feedback has demonstrated its efficacy in controlling chaotic systems [21,22] since this approach was introduced by Ott and coworkers [23] and later by Pyragas [24]. Additionally, delayed feedback has been used to suppress relaxation oscillations [25], synchronize ensembles of coupled oscillators [26], and investigate emergent dynamics in mechanical [27] and optoelectronic [28] systems. This technique has also been investigated in chemical and biological systems in both pointlike and spatially extended diffusive systems. Time-delayed feedback was applied to control unstable orbits in the oscillatory BZ reaction [29]. A rich diversity of spatiotemporal patterns has been investigated by the use of differential-difference equations in a one-species reaction-diffusion system with delay [30]. Control of spatiotemporal chaotic patterns has been attained in the reaction-diffusion Gray-Scott two-species model by means of time-delayed feedback [31]. This strategy can be harnessed to induce pattern transitions [32], spiral waves and their modulation [33], and complex dynamics of localized structures [34] and Turing patterns [35] in reaction-diffusion systems. The emergence of spatial patterns triggered by a time delay via a Hopf bifurcation is observed in a plankton prey-predator model [36]. The combination of noise, inherent to any physical system, with time delay in the feedback applied to oscillatory systems can induce undesirable oscillations [25,37], a phenomenon observed in different systems [38–40]. The introduction of a second feedback loop with delay has been applied to stabilize the system or suppress the undesirable dynamics [41–43].

\*hector.tnh@gmail.com

<sup>1</sup> $\text{Ru}(\text{bpy})_3$  tris(bipyridine)ruthenium(II) is a coordination complex that can be used to trigger photooxidation or photoreduction.

The application of global feedback has been investigated with oscillatory systems [44], coupled oscillator networks [45,46], and spatially distributed reaction-diffusion systems [47,48]. The combination of time-delayed with global feedback has been explored in chemical systems [49] and oscillator arrays [50]. Double delayed feedback [51], event-triggered feedback [52], and variable delay feedback [53] have also been explored. The above investigations have employed negative feedback, positive feedback, or both.

Time delays can arise from finite time propagation and processing of signals. A reasonable assumption in systems with time-delayed feedback is that in real processes time delay will not be deterministic, but in general will undergo stochastic fluctuations as a consequence of uncontrolled variables or perturbations. This is particularly important in synchronization processes when the intended synchronizing elements do not share a unique time delay in their feedback. Nevertheless, in the active and diverse field of research on feedback, to our knowledge there have been no previous investigations of the role of feedback with stochastic time delay and its effects on system dynamics. We investigate here the behavior of a model chemical oscillator subjected to a time-delayed negative feedback where the delay time is stochastic with a known probability distribution, and we compare our results with the effects of a deterministic time delay. We perform numerical experiments on the photosensitive FKN mechanism augmented with inhibitory electromagnetic radiation to study the effects of applying a negative feedback (NFB). The light intensity is set proportional to the instantaneous concentration of photosensitive catalyst in the reaction, and we choose the normal and Gumbel distributions for the stochastic time delay.

## II. SIMULATION

We investigate the effect of time-delayed feedback on the dynamics of the FKN mechanism with photoinhibition. NFB is applied as an inhibitory electromagnetic radiation field on a point FKN oscillator. An experimental realization approximating a point FKN oscillator is a small drop [54,55] or a continuously stirred tank reactor containing the BZ reaction [56]. The complete model used in the simulations is presented in Eqs. (1)–(10), which include time-delayed feedback by photoinhibition:

$$\dot{x}_1 = -k_1 x_1 x_2 + k_2 x_2 - 2k_3 x_1^2 - k_4 x_1 + k_r x_6^2 + k_{red} x_6 x_7, \quad (1)$$

$$\dot{x}_2 = -k_1 x_1 x_2 - k_2 x_2 - k_5 x_2 x_4 + k_6 x_5 + k_7 x_5 + k_9 x_3 + \phi, \quad (2)$$

$$\dot{x}_3 = k_{red} x_6 x_7 - k_9 x_3 - k_{10} x_3 + \phi, \quad (3)$$

$$\dot{x}_4 = 2k_1 x_1 x_2 + k_2 x_2 + k_3 x_1^2 - k_5 x_2 x_4 + k_6 x_5 - k_8 x_4, \quad (4)$$

$$\dot{x}_5 = k_5 x_2 x_4 - k_6 x_5 - k_7 x_5, \quad (5)$$

$$\dot{x}_6 = 2k_4 x_1 - 2k_r x_6^2 - k_{red} x_6 x_7, \quad (6)$$

$$\dot{x}_7 = -k_{red} x_6 x_7 + k_9 x_3 + k_{10} x_3 - \phi, \quad (7)$$

$$\phi(x, t) = k(I) x_7 b / (b_C + b), \quad (8)$$

$$k(I) = K_1 I, \quad (9)$$

$$I = K_2 x_3(t - \tau). \quad (10)$$

The ordinary differential Eqs. (1)–(7), excluding the function  $\phi$ , are generated from the original FKN mechanism under mass action kinetics and constitute an adequate model of the BZ reaction. Here,  $x_1 = [\text{HBrO}_2]$ ,  $x_2 = [\text{Br}^-]$ ,  $x_3 = [\text{oxidized catalyst}]$ ,  $x_4 = [\text{HOBr}]$ ,  $x_5 = [\text{Br}_2]$ ,  $x_6 = [\text{BrO}_2]$ , and  $x_7 = [\text{reduced catalyst}]$ . Additionally,  $x_3 + x_7 = c_0$ , where  $c_0$  is the total concentration of the catalyst either in reduced or oxidized form, which remains constant over time. The catalyst,  $\text{Ru}(\text{bpy})_3$ , which has reduced  $[\text{Ru}(\text{bpy})_3]^{2+}$  and oxidized  $[\text{Ru}(\text{bpy})_3]^{3+}$  forms, is used in the photosensitive version of the BZ reaction. When there is no illumination, an idealized point BZ reactor, or alternatively the point FKN mechanism, behaves as a nonlinear oscillator. The photosensitive BZ reaction is inhibited partially or totally by electromagnetic radiation, which is accounted for by the presence of the function  $\phi$  [defined in Eq. (8)] in Eqs. (2), (3), and (7) of the photosensitive FKN mechanism. Photoinhibition is mediated by bromide ion when the photosensitive catalyst  $\text{Ru}(\text{bpy})_3$  is subjected to electromagnetic radiation ( $\lambda = 450 \text{ nm}$ ) [16]. Here,  $b = [\text{BrCH}(\text{COOH})_2]$ ,  $b_C = 0.05 \text{ M}$ , and  $k(I)$  is the reaction rate constant for light-mediated production of  $[\text{Ru}(\text{bpy})_3]^{2+*}$  (photo-activated  $[\text{Ru}(\text{bpy})_3]^{2+}$ ), which in turn promotes the production of bromide (see Refs. [19,20] and [16] for details).

We perform numerical experiments to simulate the application of an inhibiting radiation beam of intensity  $I$  to the FKN oscillator to build a NFB signal. We assume that  $k(I)$  is proportional to the inhibitory radiation intensity  $I$  according to Eq. (9). Additionally, in Eq. (10) we define  $I$  to be proportional to the concentration of oxidized catalyst ( $x_3 = [\text{Ru}(\text{bpy})_3]^{3+}$ ) at  $t - \tau$ , where  $t$  is time and  $\tau$  a time delay.  $K_1$  and  $K_2$  are appropriate constants for dimensional consistency. Thus, if  $x_3$  (the oxidized catalyst concentration) increases, the inhibitory radiation illuminating the oscillator rises after a time delay  $\tau$ , which in turn increases the concentration of bromide, thereby incrementing the inhibition on the reaction. The net result is that the radiation inhibition  $\phi$  is dependent upon the time-delayed concentration of the species  $x_3$ . We can tune the total and average radiation intensity by varying  $I$ .

We explore different scenarios as described below by applying feedback without time delay ( $\tau = 0$ ) and with a time delay  $\tau$ . When  $\tau > 0$ , we examine the delay effects and discover the preservation of key aspects of the FKN dynamics in the presence of feedback with constant (therefore deterministic) and with stochastic time delay. The constant time delay calculation is straightforward. We apply stochastic time delay as follows. At each solution point while solving the system of differential Eqs. (1)–(7), we control, through Eq. (10), the time delay  $\tau$  stochastically, choosing it from a given probability distribution. Hence,  $\tau$  is time-dependent. For stochastic time delay, we investigate two probability distributions, the normal distribution and the type I extreme value distribution (Gumbel distribution) of the minimum [57]. In the first case,  $\tau$  is a random number taken from a normal distribution with known mean  $\mu$  and standard deviation  $\sigma$ . In the second case,  $\tau$  is a random number taken from a Gumbel distribution with known location parameter  $\alpha$  and scale parameter  $\beta$  according to the expression.

$$P(t - \tau) = \beta^{-1} \exp \left[ \frac{t - \tau - \alpha}{\beta} - \exp \left( \frac{t - \tau - \alpha}{\beta} \right) \right]. \quad (11)$$

For each distribution we analyze six families, varying the width of the distribution by choosing six values of the standard deviation ( $\sigma$ ) for the normal distribution and six values of the scale parameter ( $\beta$ ) for the Gumbel distribution. For simplicity in exposition, we define the nominal time delay  $\tau_N$  such that: (i)  $\tau = \tau_N$  when  $\tau$  is constant, (ii)  $\mu = t - \tau_N$  when  $\tau$  is stochastic and normally distributed, and (iii)  $\alpha = t - \tau_N$  when  $\tau$  is stochastic and follows a Gumbel distribution. As a result, the distribution from which stochastic time delay,  $\tau$ , is obtained, is constantly translated to the right at the same speed as  $t$  increases, i.e., as each new solution point is calculated. The distribution is centered ( $\mu$  for the normal and  $\alpha$  for the Gumbel are located) behind  $t$  an amount  $\tau_N$ . Feedback is applied only when  $t \geq 397 + 2\tau_N$  s, and  $\tau_N$  is at least five times larger than the width of the distribution ( $\tau_N \geq 5\sigma, 5\beta$ ). As a consequence, only the tails of the distribution reach  $\tau < 0$  and  $\tau > t$  and the probability to obtain these values of  $\tau$  is zero in our simulations.

The system of differential Eqs. (1)–(7) is numerically integrated with MATLAB variable order, multistep solver ode15s, which is based on the numerical differentiation formulas and optionally the backward differentiation formulas, methods particularly suitable to solve stiff problems [58–60].

### III. RESULTS AND DISCUSSION

First, we explore the role played by the average intensity level of the applied inhibiting radiation intensity,  $I$ . Due to the fundamental role played by the oxidized photosensitive catalyst  $x_3$ , and since all seven FKN variables have the same oscillation frequency, we use  $x_3$  to characterize the system dynamics. Figure 1(a) shows a comparison of the behavior of  $x_3$  versus

time in the free system, i.e., when there is no inhibitory radiation, and when inhibitory radiation is present. At  $I = 0.01$ , the oscillation period  $T$  is increased and the peak amplitude  $x_{3,p}$  slightly decreased. Oscillations are suppressed at a critical average radiation intensity  $I_C = 0.011$ . Below this critical intensity, the main consequences of increasing inhibitory radiation intensity are the lengthening of the oscillation period and a slight decrease of the peak amplitude, with no significant effect on the shape of the curve. Figures 1(b)–1(d) illustrate these effects over a range of intensities. In Figs. 1(b), 1(c) and 1(d),  $t_0$  is the time at which the radiation feedback is turned on. The graphs show no difference when inhibitory radiation is turned on at time  $t_0 = 0$ , the time at which the simulation is initiated, or when the radiation is turned on at  $t_0 = 397$  s. This result is unexpected, because transient behavior occurs during the first cycle of  $x_3$ . We selected the time at the second minimum of  $x_3$  ( $t = 397$  s) because at this point regular periodic behavior is already established in the free system [Fig. 1(a),  $I = 0$  curve]. Even though there is no dependence on the time at which radiation inhibition is turned on, henceforth unless specified otherwise, we turn on inhibitory radiation at  $t_0 = 397 + 2\tau$  s.

Next, we introduce a time delay  $\tau$  in the application of the feedback and examine its effects. First we consider constant, i.e., deterministic time delay. Figure 2 shows  $x_3$  versus time at the critical inhibiting radiation for different values of  $\tau$ . As previously noted, oscillations are suppressed at the critical radiation intensity in the absence of time delay. Interestingly, the system recovers its oscillatory dynamics if the inhibiting critical radiation is applied with a time delay. For  $\tau \geq 2$  s, system oscillations are recovered with no significant change in the  $x_3$  peak amplitude and time profile.

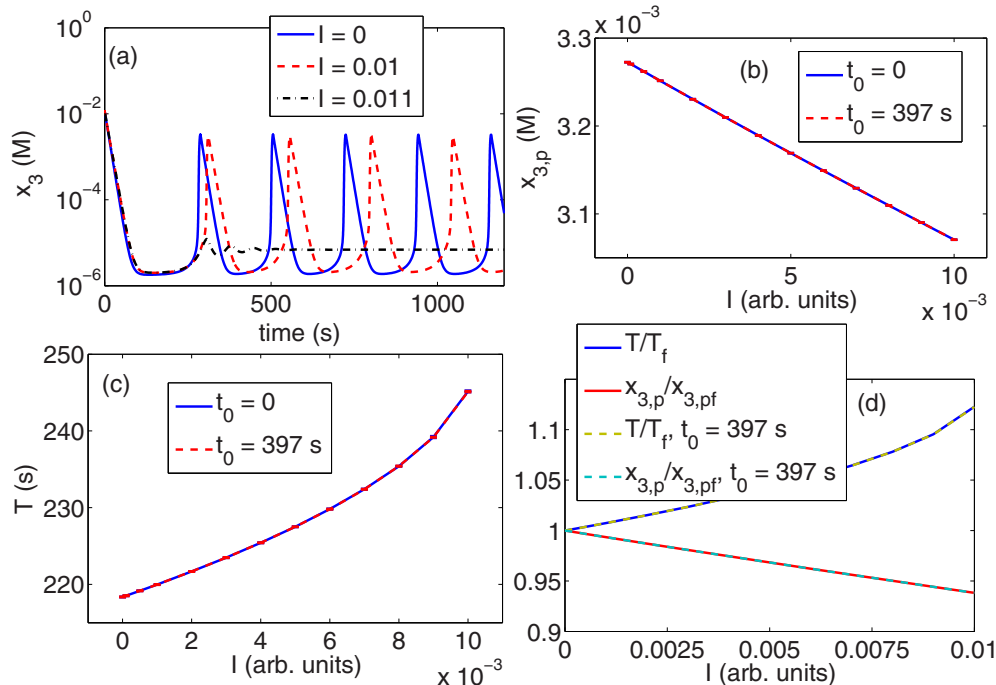


FIG. 1. Effects of radiation intensity on system dynamics. (a) Plot of  $x_3$  time dependence for increased values of  $I$ . Critical radiation intensity  $I_C = 0.011$  suppresses oscillatory dynamics. (b) Peak amplitude  $x_{3,p}$  decreases linearly and (c) oscillation period  $T$  increases as inhibitory radiation increases. (d)  $T$  and  $x_{3,p}$  time profiles normalized to free system (no-inhibition) values. Values of maximum amplitude and period are means averaged over several oscillatory cycles. Error bars (standard deviation) are barely noticeable in plots (b) and (c).

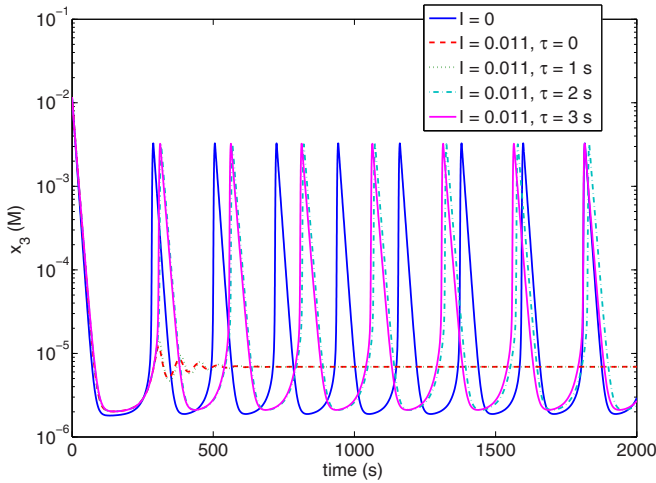


FIG. 2. Time dependence of  $x_3$  at critical radiation intensity for increasing time-delayed feedback. The introduction of  $\tau \geq 2$  s in feedback, results in system recovery of oscillatory dynamics at critical radiation intensity. The scale is the same for all plots.

We now consider how the system dynamics is modified when inhibitory feedback is applied at a subcritical radiation intensity with constant time delay. Figure 3 shows nine plots of  $x_3$  versus time at  $I = 0.01$  for increasing values of  $\tau$ . These plots correspond to the different characteristic time profiles exhibited by  $x_3$  as a function of time-delayed feedback. The time window for each plot was selected after the transients died out and the system exhibited regular periodic behavior.

The figure shows significant changes in the  $x_3$  time profile as  $\tau$  increases. At  $\tau = 40, 80, 110, 120, 200,$  and  $240$  s, we observe more complex time profiles, which contain 2, 3, 4, 3, 2, and 3 peaks per cycle, respectively. Interestingly, at  $\tau = 250$  s,

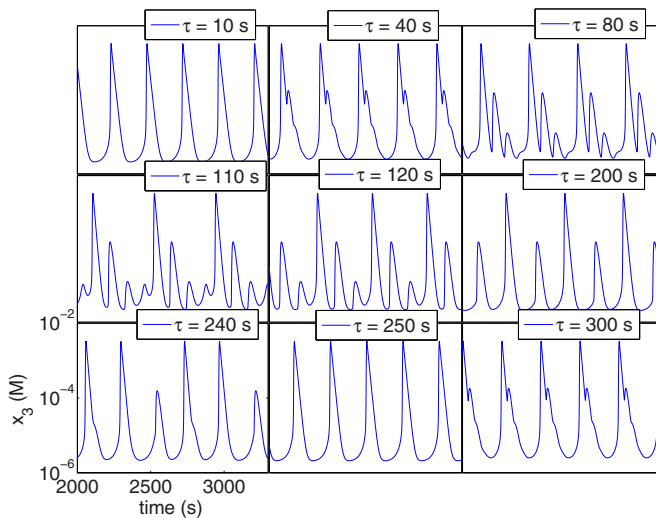


FIG. 3. Plot of  $x_3$  vs. time as a function of increasing constant time delay (shown in the top right side in seconds). As time delay increases new peaks per cycle emerge, differentiate, and disappear. Note that free system dynamics appears to be recovered at  $\tau = 250$  s, compare with  $\tau = 10$  s and with Fig. 1(a) curve  $I = 0$ . Also note that system behaves similarly at  $\tau = 40$  s and  $\tau = 300$  s.

the waveform of the system at  $\tau \leq 10$  s is recovered (which essentially is the same of the free system), with one peak per cycle. The oscillation period of 245 s is the same as that at  $\tau = 10$  s (Fig. 9). The last plot at  $\tau = 300$  s resembles that at  $\tau = 40$  s with a similar period. This could suggest that system behavior is periodic in  $\tau$  with a period of about 250 s. See discussion of Fig. 9 on this issue.

Finally, we investigated the role of negative feedback with stochastic time delay at a subcritical radiation intensity ( $I = 0.01$ ). Stochastic time delays with normal and Gumbel probability distributions were used. In choosing the normal and Gumbel distributions, one of our purposes was to compare the effects of stochastic time delay from a symmetric and a nonsymmetric distribution. The dispersions applied were  $\sigma = 0.25, 1, 2, 4, 10,$  and  $20$  s for the normal distribution and  $\beta = 0.25, 1, 2, 4, 10,$  and  $20$  s for the Gumbel distribution. Figure 4 shows plots of  $x_3$  in the real [Figs. 4(a) and 4(b)] and Fourier [Fig. 4(c)] spaces at  $\tau_N = 100$  s for the 13 conditions of time-delayed feedback investigated: one constant and the 12 stochastic just described. Again, the frame has been chosen after transients, once periodic behavior is established. In real space, all plots exhibit strikingly similar dynamics. We observe good matching between plots for dispersions of 4 s and below for both distributions. At large dispersion, 10 and 20 s [Fig. 4(b)], the only important difference between plots is in frequency and/or phase. The first peaks, which occur at the same location in Fourier space in most of the plots, correspond to the major frequency component. The slight frequency difference at low dispersion ( $\leq 4$  s), noticeable in real space, is not evidenced in the peaks location in Fourier space, due to the large scale of the plot. However, at large dispersion ( $\geq 10$  s) the frequency is significantly larger, as evidenced by the shift of the main peaks in the Fourier transform. Additionally, a large set of lower amplitude frequency components is apparent in the Fourier domain. Figure 5 shows an analogous plot at  $\tau_N = 200$  s. Again, there is a remarkable match of dynamic behavior in real space [Figs. 5(a) and 5(b)] for all the conditions investigated. The most significant differences occur in phase for large dispersions [Fig. 5(b)]. Primary and several secondary frequency components exhibit impressive agreement as shown in plots of the Fourier domain [Fig. 5(c)], for all conditions.

The same trend of closely matching dynamics for the entire set of time-delayed feedback conditions investigated is observed for the complete interval of nominal time delay explored ( $10 \text{ s} \leq \tau_N \leq 300 \text{ s}$ ). Time profile, frequency, and mainly phase exhibit growing differences as the dispersion in time delay increases. The similarity is particularly noticeable at low time-delay dispersion ( $\leq 4$  s) with the striking exceptions of  $\tau_N = 110$  and  $240$  s. Figure 6 shows the case of  $\tau_N = 110$  s. In real space [Fig. 6(a)], only plots 1, 2, 3, and 8 (constant,  $\sigma = 0.25$  s,  $\sigma = 1$  s, and  $\beta = 0.25$  s) exhibit near-perfect agreement in every aspect of their dynamics. These plots present four maxima (one global and three local) per oscillation cycle in contrast with three or two for the remaining conditions with periodic behavior. The transition from four to three maxima per oscillation occurs at  $\sigma = 2$  s in the normal distribution (plot 4). This plot appears to be aperiodic with an apparently random peak sequence 4-3-3-4-3. An analogous

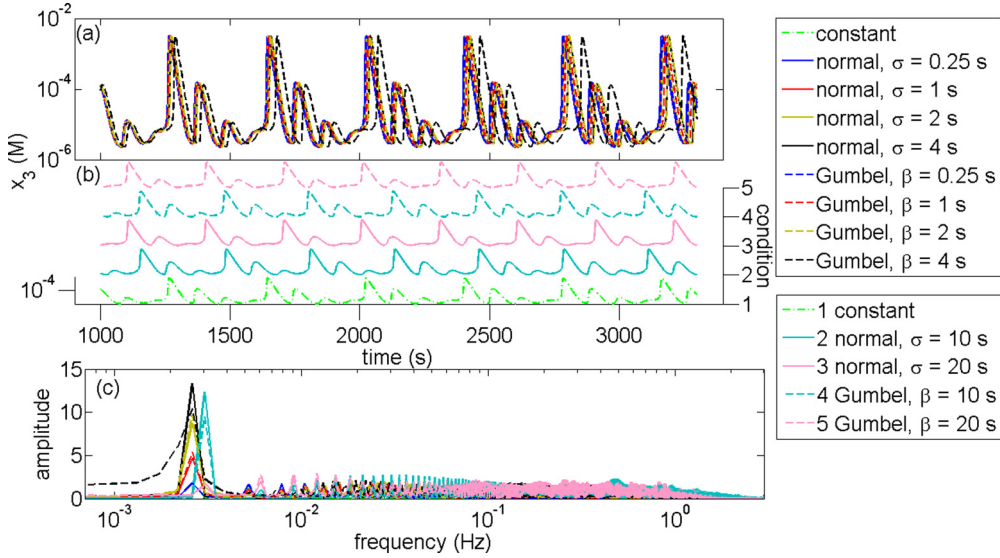


FIG. 4. Real (a, b) and Fourier (c) space plots of  $x_3$  for all time-delay conditions investigated at  $\tau_N = 100$  s. Real-space profiles exhibit closely similar dynamic behavior for all time delay conditions. Fourier analysis shows the concordance of the main and some secondary frequency components roughly divided into small and large dispersions. Plots of  $x_3$  at small (a) and large (b) dispersion exhibit essentially equal amplitude.

transition in the number of peaks per oscillation with aperiodic behavior is observed in plot 9 (Gumbel distribution  $\beta = 1$  s), which has the peak sequence 3-4-4-4-3. In both cases (normal and Gumbel distributions), two different runs with identical simulation conditions but different random sequences of time delays generated different peak sequences. One outcome of the increase in time delay dispersion is the suppression of secondary peaks in the real-space dynamics and a decrease in oscillation period. This is revealed in Fourier space [Fig. 6(b)], where plots for  $\sigma \geq 4$  s and  $\beta \geq 2$  s show the main frequency component peaks shifted to the right.

To have a quantitative measure of the match between  $x_3$  waveforms subjected to feedback with stochastic and constant time delays independent of frequency and phase, we define the

quantity  $q_i$  as

$$q_i = \frac{\int_0^1 |v_i - u| dt}{\int_0^1 u dt}, \quad i = 1, 2, \dots, 12, \quad (12)$$

where  $u(t) = x_3(Tt + t_M)$  under constant time delay and  $v_i(t) = x_{3i}(T_i t + t_{Mi})$  under stochastic time delay (the index  $i$  specifies which form of stochastic time delay is used, see Fig. 7),  $T(T_i)$  is the oscillation period at constant(stochastic) time delay, and  $t_M(t_{Mi})$  is the time of the penultimate peak when time delay is constant(stochastic). Equation (12) integrates the absolute difference between frequency-normalized states  $x_3$  under stochastic and constant time delay, over one oscillation cycle (the last simulated), with their phases matched, and

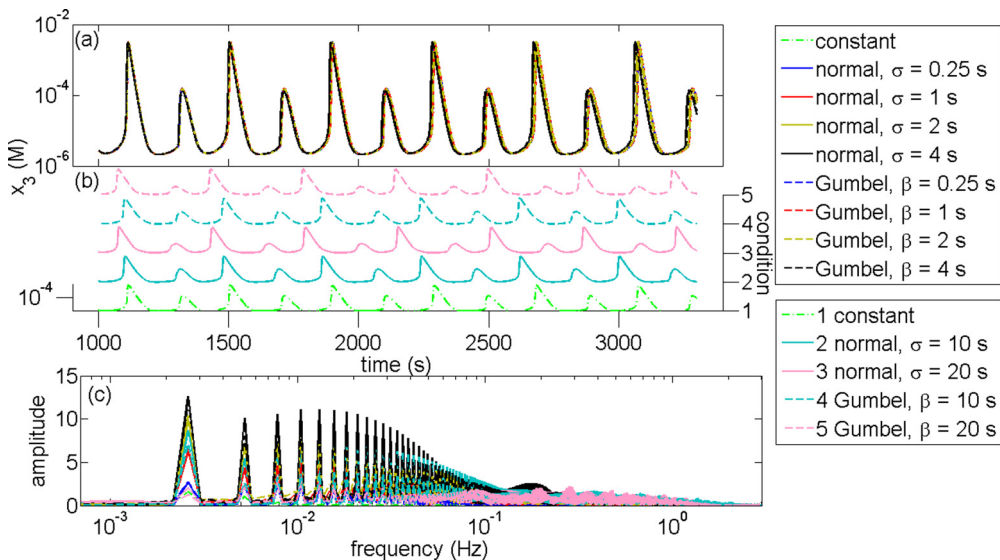


FIG. 5. Real (a, b) and Fourier (c) domains of  $x_3$  at  $\tau_N = 200$  s for all time delay conditions explored. As in the previous figure, the agreement between all plots is remarkable. Plots of  $x_3$  at small (a) and large (b) dispersion exhibit essentially equal amplitude.

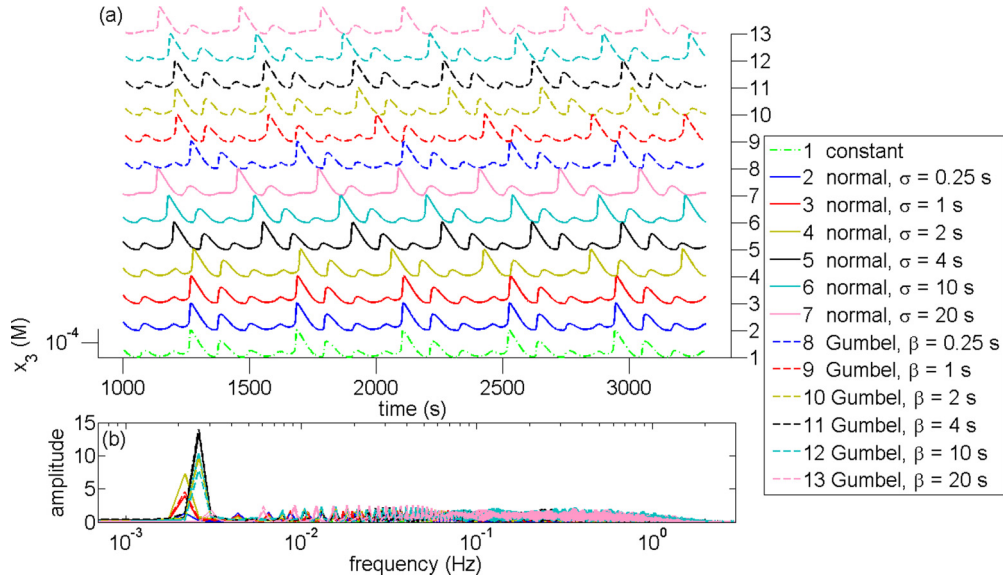


FIG. 6. Plots of  $x_3$  in real (a) and Fourier (b) spaces for all conditions of time delay investigated at  $\tau_N = 110$  s. Periodicity is preserved in curves 1, 2, 3, 5, and 7. Plots 1, 2, 3, and 5 still exhibit a close match while plot 7 shows a larger frequency, evidenced in the Fourier domain, where the frequency component peaks are shifted to the right. Plots 4 and 6 appear to be aperiodic functions, though they still exhibit main frequency components very similar to those of the periodic plots.

normalizes this integrated difference to the one-cycle integral of  $u$ .

Figure 7 shows  $q_i$  as a function of nominal time delay for all conditions of stochastic time-delayed feedback investigated. Remarkably, there is only one condition of nominal time delay,  $\tau_N = 240$  s, at which  $q_i$  is larger than 30%. This occurs solely for the largest dispersions in time delay explored ( $\geq 10$  s). For discussion purposes and based on the figure, we define low ( $\leq 4$  s) and high ( $\geq 10$  s) time-delay dispersions for the behavior of  $q_i$ . At low dispersion  $q_i$  is always less than 2% over the entire

range of nominal time delay, with two exceptions  $\tau_N = 110$  and 240 s. The case of  $\tau_N = 110$  s has been discussed in Fig. 6. A similar situation occurs at  $\tau_N = 240$  s. Even in these cases,  $q_i$  is close to 10%. At large dispersion,  $q_i$  increases moderately, remaining below 30% with the exception mentioned above. The largest  $q_i$  values occur at the largest dispersion explored, 20 s, and the Gumbel distribution generally exhibits larger  $q_i$  than the normal distribution. There are three intervals that contain large values of  $q_i$ :  $\tau_N = 100$ –110 s,  $\tau_N = 130$ –180 s and  $\tau_N = 240$  s.

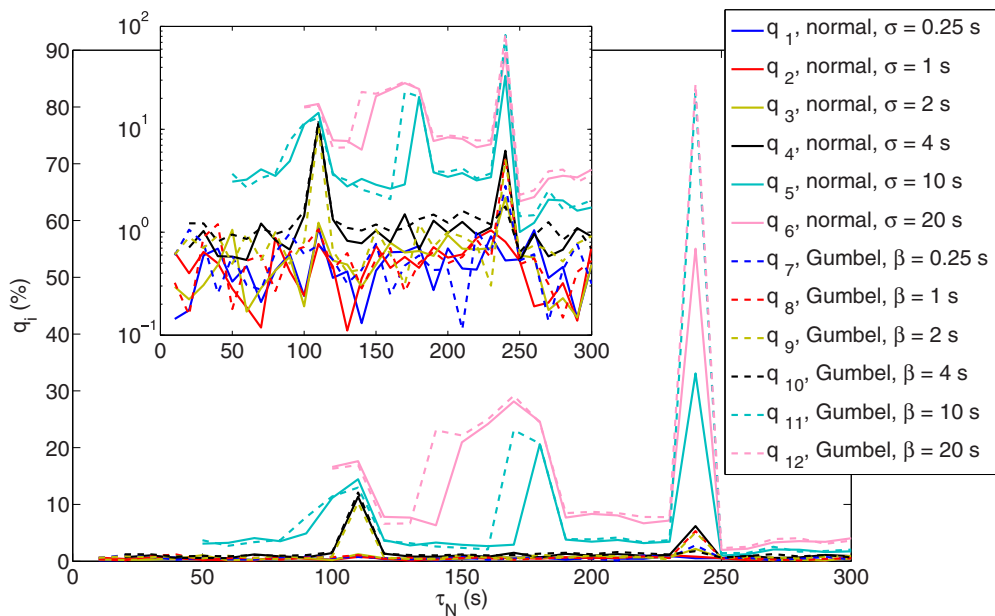


FIG. 7. Plot of  $q_i$  vs.  $\tau_N$  shows that for all conditions of time delay studied  $q_i$  remains below 30% for nearly all  $\tau_N$ , except near  $\tau_N = 240$  s. The inset shows a logarithmic plot of the same data.

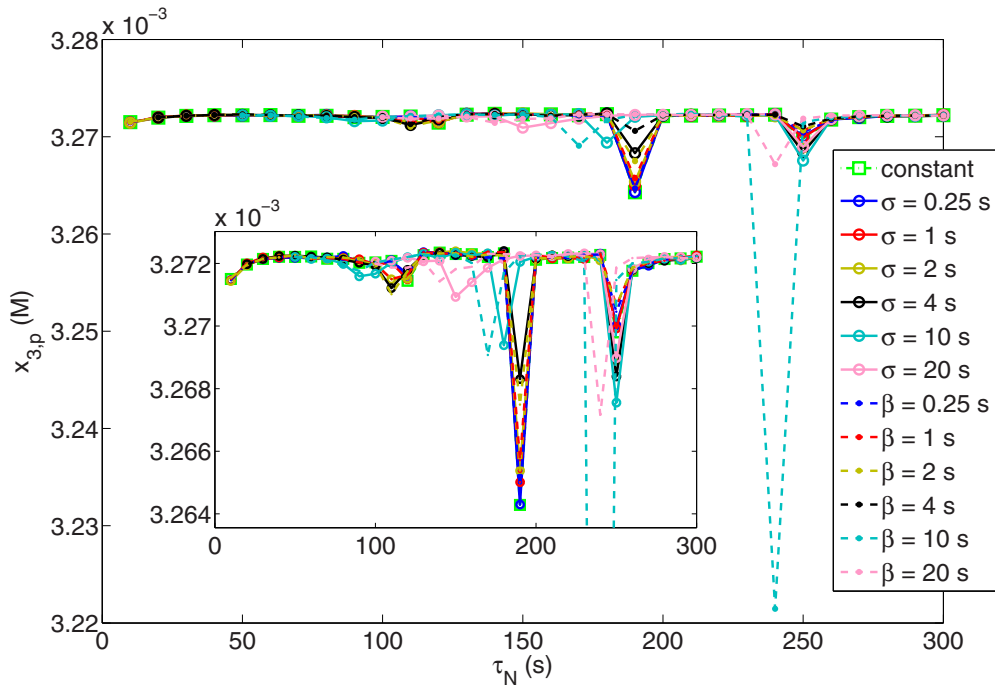


FIG. 8. Plot of  $x_3$  peak amplitude,  $x_{3,p}$ , as a function of  $\tau_N$ . The plot shows a remarkable agreement between all time-delayed feedback conditions. A logarithmic plot of these data is shown in the inset.

Finally, we present results for the signal  $x_3$  amplitude (maxima) and period (frequency). Figure 8 shows a plot of signal amplitude,  $x_{3,p}$ , versus nominal time delay. The plot compares constant and all stochastic time-delayed feedback conditions. Besides small local differences, there is a surprising match for the entire set of curves. The largest mismatch, which occurs at  $\tau_N = 240$  s for  $\beta = 10$  s, is less than 1.6% relative to constant time delay. There are three intervals where noticeable decrements in peak amplitude occur (including conditions at

constant and small dispersion stochastic time delay), at or around  $\tau_N = 110$  s,  $\tau_N = 140-190$  s, and  $\tau_N = 240-250$  s. In all three cases there is a decrease in the number of maxima per oscillation cycle of the state  $x_3$  (see Fig. 3). The number of peaks per cycle decreases from four to three peaks at  $\tau_N \simeq 110$  s, from three to two at  $\tau_N = 190$  s, and from three to one at  $\tau_N = 250$  s (in this case the number of peaks increases from two to three at  $\tau_N = 240$  s). Notice that secondary (local) maxima fade or disappear as the dispersion increases, as seen

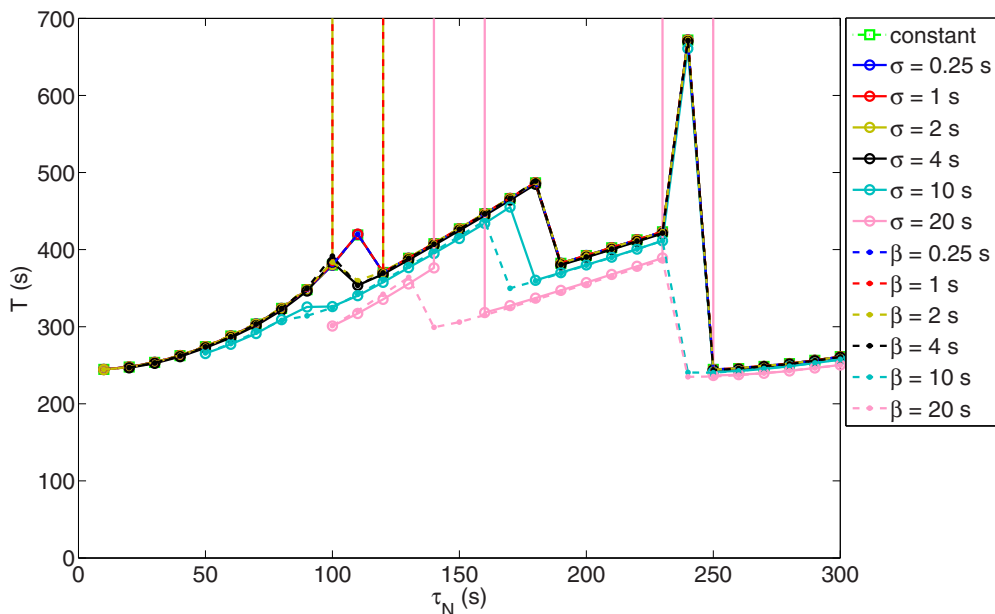


FIG. 9. Oscillation period  $T$  as a function of  $\tau_N$ . As in Fig. 8, the curves exhibit a remarkable match for all conditions of time-delayed feedback applied.

in Figs. 4 and 6. We emphasize that the intervals mentioned roughly correspond to the peaks in Fig. 7.

Figure 9 shows the oscillation period as a function of the nominal time delay. The signal exhibits four continuous regions of increasing period as  $\tau_N$  increases. In the boundary between these regions there are intervals (three) which suggest the existence of a discontinuity. These intervals loosely coincide with (a) the regions where the  $x_3$  amplitude and number of peaks per cycle decrease and with (b) the peaks of Figure 7, i.e., at  $\tau_N \simeq 110$  s, 140–190 s, and 240 s. Two regions of period increase exhibit nonlinear behavior (first and fourth regions from left to right) and two exhibit linear behavior (regions two and three). We find four cases at which state  $x_3$  does not exhibit periodic behavior, marked with infinite period in the Figure (vertical lines). The cases are: (1)  $\sigma = 2$  s,  $\tau_N = 110$  s; (2)  $\beta = 1$  s,  $\tau_N = 110$  s; (3)  $\sigma = 20$  s,  $\tau_N = 150$  s; and (4)  $\sigma = 20$  s,  $\tau_N = 240$  s. These cases lie within the intervals described. As discussed in Fig. 3, the recovering of the oscillation period and waveform at  $\tau_N = 250$  s suggests that system behavior is periodic in time delay  $\tau$ . However, we note in Fig. 9 that the first region of the plots ( $10 \text{ s} \leq \tau_N \leq 60 \text{ s}$ ) grows faster and has a larger first derivative than the last region ( $250 \text{ s} \leq \tau_N \leq 300 \text{ s}$ ).

#### IV. CONCLUSION

We have investigated the effect of time-delayed negative feedback on a point photosensitive FKN oscillator by means of simulated inhibitory radiation. We find that increasing the radiation intensity with no time delay lengthens the oscillatory period and lowers the peak amplitude of the oxidized photosensitive catalyst concentration profile. At a critical intensity of illumination, oscillatory dynamics is suppressed. These observations can be attributed to the production of bromide, which in turn inhibits the oxidation of  $[\text{Ru}(\text{bpy})_3]^{2+}$ . If the concentration of the inhibitor increases, the time needed for its concentration to decrease to values compatible with oxidation transitions will also increase; thus, the refractory time and therefore the oscillation period should also increase. Similarly, at increased inhibitor concentrations, the maximum concentration of oxidized catalyst should decrease, as the available activator is partially depleted relative to the inhibition-free case. We also observe that the response of the system is the

same whether nondelayed feedback is initiated during transient behavior or once periodic behavior is established, and that introduction of delayed feedback allows recovery of oscillatory behavior in the presence of a radiation level that suppresses oscillation in the absence of delay.

Our main objective is to investigate the effects of incorporating a stochastic time delay in the feedback. We compare the response of the FKN oscillator under constant versus stochastic time-delayed feedback. Investigation with stochastic time delay included two probability distributions, a symmetric normal and a nonsymmetric Gumbel, with six values of parameter dispersion for each. The FKN model exhibits a surprising amplitude and frequency match for low dispersion ( $\leq 4$  s) in the probability distribution of the time-delayed feedback over a wide interval of time-delay values, which spanned more than 1.3 times the natural oscillation period of the free system ( $10 \text{ s} \leq \tau_N \leq 300 \text{ s}$ ). Additionally, there is a striking agreement in the time profiles for the situations described. The amplitude and period of  $x_3$  decrease or show a significant curvature change in three intervals of the nominal time delay: (1)  $\tau_N = 90$ –120 s, (2)  $\tau_N = 140$ –190 s, and (3)  $\tau_N = 240$ –250 s. The time delay (nominal) at which the transition occurs shifts toward lower values as the dispersion in the probability distribution of stochastic time delay increases. These regions closely coincide with those at which  $q_i$  exhibits bumps or peaks, i.e., when the waveform of  $x_3$  under stochastic time delay feedback diverges most from that under constant delay. In these parameter regions, one secondary peak in the oscillation cycle of  $x_3$  fades and disappears, and the signal exhibits aperiodic behavior in some cases when time delay is stochastic.

Experimental implementation of this investigation will be carried out in future work. This can be accomplished with droplets of the BZ reaction prepared in the oscillatory regime and subjected to an inhibitory light beam [11,61]. A PC connected to a CCD video camera can evaluate, from droplet image brightness, the concentration of oxidized catalyst in the drops and tune accordingly the beam intensity with an appropriate constant or stochastic time delay.

#### ACKNOWLEDGMENT

I.R.E. acknowledges support from the U.S. National Science Foundation through Grant No. CHE-1362477.

- 
- [1] B. P. Belousov, in *Collection of Short Papers on Radiation Medicine* (Medgiz, Moscow, 1959).
  - [2] A. M. Zhabotinsky, Dokl. Akad. Nauk SSSR **157**, 392 (1964).
  - [3] A. N. Zaikin and A. M. Zhabotinsky, *Nature* **225**, 535 (1970).
  - [4] A. T. Winfree, *Science* **175**, 634 (1972).
  - [5] Edited by R. Kapral and K. Showalter, *Chemical Waves and Patterns* (Kluwer, Dordrecht, 1995).
  - [6] I. Prigogine, *From Being to Becoming: Time and Complexity in the Physical Sciences* (W. H. Freeman and Company, New York, 1980).
  - [7] J. Lechleiter, S. Girard, E. Peralta, and D. Clapham, *Science* **252**, 123 (1991).
  - [8] S. Girard, A. Lückhoff, J. Lechleiter, J. Sneyd, and D. Clapham, *Biophys. J.* **61**, 509 (1992).
  - [9] A. M. Turing, *Philos. T. Roy. Soc.* **237**, 37 (1952).
  - [10] V. Castets, E. Dulos, J. Boissonade, and P. De Kepper, *Phys. Rev. Lett.* **64**, 2953 (1990).
  - [11] N. Tompkins, N. Li, C. Girabawe, M. Heymann, G. B. Ermentrout, I. R. Epstein, and S. Fraden, *Proc. Natl. Acad. Sci. USA* **111**, 4397 (2014).
  - [12] A. F. Taylor, M. R. Tinsley, F. Wang, Z. Huang, and K. Showalter, *Science* **323**, 614 (2009).
  - [13] H. Fukuda, N. Tamari, H. Morimura, and S. Kai, *J. Phys. Chem. A* **109**, 11250 (2005).



- [14] B. T. Ginn, B. Steinbock, M. Kahveci, and O. Steinbock, *J. Phys. Chem. A* **108**, 1325 (2004).
- [15] L. Kuhnert, K. I. Agladze, and V. I. Krinsky, *Nature* **337**, 244 (1989).
- [16] S. Kádár, T. Amemiya, and K. Showalter, *J. Phys. Chem. A* **101**, 8200 (1997).
- [17] R. J. Field, E. Körös, and R. M. Noyes, *J. Am. Chem. Soc.* **94**, 8649 (1972).
- [18] P. Ruoff, *J. Phys. Chem.* **88**, 2851 (1984).
- [19] V. K. Vanag, A. M. Zhabotinsky, and I. R. Epstein, *J. Phys. Chem. A* **104**, 8207 (2000).
- [20] V. K. Vanag and I. R. Epstein, *J. Chem. Phys.* **131**, 104512 (2009).
- [21] K. Pyragas and A. Tamaševičius, *Phys. Lett. A* **180**, 99 (1993).
- [22] K. Pyragas, *Phil. Trans. R. Soc. A* **364**, 2309 (2006).
- [23] E. Ott, C. Grebogi, and J. A. Yorke, *Phys. Rev. Lett.* **64**, 1196 (1990).
- [24] K. Pyragas, *Phys. Lett. A* **170**, 421 (1992).
- [25] V. Flunkert and E. Schöll, *Phys. Rev. E* **76**, 066202 (2007).
- [26] M. G. Rosenblum and A. S. Pikovsky, *Phys. Rev. Lett.* **92**, 114102 (2004).
- [27] S. A. Campbell, J. Blair, T. Ohira, and J. Milton, *Chaos* **5**, 640 (1995).
- [28] F. Lin and J. Liu, *IEEE J. Quantum Electron.* **39**, 562 (2003).
- [29] F. W. Schneider, R. Blittersdorf, A. Förster, T. Hauck, D. Lebender, and J. Müller, *J. Phys. Chem.* **97**, 12244 (1993).
- [30] M. Bestehorn, E. V. Grigorieva, and S. A. Kaschenko, *Phys. Rev. E* **70**, 026202 (2004).
- [31] Y. N. Kyrychko, K. Blyuss, S. Hogan, and E. Schöll, *Chaos* **19**, 043126 (2009).
- [32] Q. Li and H. Hu, *J. Chem. Phys.* **127**, 154510 (2007).
- [33] H. Hu, X. Li, Z. Fang, X. Fu, L. Ji, and Q. Li, *Chem. Phys.* **371**, 60 (2010).
- [34] S. V. Gurevich, *Phys. Rev. E* **87**, 052922 (2013).
- [35] R. Iri, Y. Tonosaki, K. Shitara, and T. Ohta, *J. Phys. Soc. Jpn.* **83**, 024001 (2014).
- [36] C. Tian and L. Zhang, *Phys. Rev. E* **88**, 012713 (2013).
- [37] D. Goldobin, M. G. Rosenblum, and A. Pikovsky, *Phys. Rev. E* **67**, 061119 (2003).
- [38] D. Arsenijević, M. Kleinert, and D. Bimberg, *Appl. Phys. Lett.* **103**, 231101 (2013).
- [39] M. C. Soriano, J. Garca-Ojalvo, C. R. Mirasso, and I. Fischer, *Rev. Mod. Phys.* **85**, 421 (2013).
- [40] R. Vicente, L. L. Gollo, C. R. Mirasso, I. Fischer, and P. Gordon, *Proc. Natl. Acad. Sci. USA* **105**, 17157 (2008).
- [41] C. Masoller, *Phys. Rev. Lett.* **88**, 034102 (2002).
- [42] N. B. Janson, A. G. Balanov, and E. Schöll, *Phys. Rev. Lett.* **93**, 010601 (2004).
- [43] L. Jaurigue, E. Schöll, and K. Lüdge, *Phys. Rev. Lett.* **117**, 154101 (2016).
- [44] Y. Kanevsky and A. A. Nepomnyashchy, *Phys. Rev. E* **76**, 066305 (2007).
- [45] B. Lindner, B. Doiron, and A. Longtin, *Phys. Rev. E* **72**, 061919 (2005).
- [46] S. Gil and A. S. Mikhailov, *Phys. Rev. E* **79**, 026219 (2009).
- [47] V. K. Vanag, A. M. Zhabotinsky, and I. R. Epstein, *J. Phys. Chem. A* **104**, 11566 (2000).
- [48] V. K. Vanag, L. Yang, M. Dolnik, A. M. Zhabotinsky, and I. R. Epstein, *Nature* **406**, 389 (2000).
- [49] A. S. Mikhailov and K. Showalter, *Phys. Rep.* **425**, 79 (2006).
- [50] W. Wang, I. Z. Kiss, and J. Hudson, *Ind. Eng. Chem. Res.* **41**, 330 (2002).
- [51] S. Yanchuk and G. Giacomelli, *Phys. Rev. Lett.* **112**, 174103 (2014).
- [52] J. A. Kromer, B. Lindner, and L. Schimansky-Geier, *Phys. Rev. E* **89**, 032138 (2014).
- [53] A. Gjurchinovski and V. Urumov, *Phys. Rev. E* **81**, 016209 (2010).
- [54] M. Toiya, V. K. Vanag, and I. R. Epstein, *Angew. Chem. Int. Ed.* **47**, 7753 (2008).
- [55] M. Toiya, H. O. González-Ochoa, V. K. Vanag, S. Fraden, and I. R. Epstein, *J. Phys. Chem. Lett.* **1**, 1241 (2010).
- [56] A. N. Zaikin and A. M. Zhabotinsky, in *Biological and Biochemical Oscillators*, edited by B. Chance, E. K. Pye, A. K. Ghosh, and B. Hess (Academic Press, New York, 1973).
- [57] R. D. Reiss and M. Thomas, *Statistical Analysis of Extreme Values*, Handbook on Statistical Distributions for Experimentalists (Birkhäuser Verlag, University of Stockholm, 2007).
- [58] L. F. Shampine and M. W. Reichelt, *SIAM J. Sci. Comput.* **18**, 1 (1997).
- [59] L. F. Shampine, M. W. Reichelt, and J. A. Kierzenka, *SIAM Rev.* **41**, 538 (1999).
- [60] R. W. Klopfenstein, *RCA Rev.* **32**, 447 (1971).
- [61] J. Delgado, N. Li, M. Leda, H. O. González-Ochoa, S. Fraden, and I. R. Epstein, *Soft Matter* **7**, 3155 (2011).