Thin liquid film as an optical nonlinear-nonlocal medium and memory element in integrated optofluidic reservoir computer

Chengkuan Gao, Prabhav Gaur, Shimon Rubin,* and Yeshaiahu Fainman
University of California, San Diego, Department of Electrical and Computer Engineering, La Jolla, California, United States

Abstract. Understanding light–matter interaction lies at the core of our ability to harness physical effects and to translate them into new capabilities realized in modern integrated photonics platforms. Here, we present the design and characterization of optofluidic components in an integrated photonics platform and computationally predict a series of physical effects that rely on thermocapillary-driven interaction between waveguide modes and topography changes of optically thin liquid dielectric film. Our results indicate that this coupling introduces substantial self-induced phase change and transmittance change in a single channel waveguide, transmittance through the Bragg grating waveguide, and nonlocal interaction between adjacent waveguides. We then employ the self-induced effects together with the inherent built-in finite relaxation time of the liquid film, to demonstrate that the light-driven deformation can serve as a reservoir computer capable of performing digital and analog tasks, where the gas–liquid interface operates both as a nonlinear actuator and as an optical memory element.

Keywords: light–liquid interaction; optofluidics; nanophotonics; silicon photonics; reservoir computing; thermocapillary effect.

© The Authors. Published by SPIE and CLP under a Creative Commons Attribution 4.0 International License. Distribution or reproduction of this work in whole or in part requires full attribution of the original publication, including its DOI.
[DOI: 10.1117/1.AP.4.4.046005]

1 Introduction

Light–matter interaction resides at the core of advancing our understanding of both light and matter properties. The emergence of robust-integrated photonic platforms over the last two decades, characterized by miniaturized cross-section and higher refractive index contrast, was leveraged to enhance the optical intensity, thus achieving nonlinear response of various matter degrees of freedom. More recently, nonlinear integrated photonics platforms1 demonstrated promising capabilities to conduct basic scientific research in technologically attractive applications, such as frequency conversion and third-harmonic generation,2 supercontinuum generation,3 emerging quantum photonics applications,4 and are considered as an attractive platform for future nonconventional computation architectures.3 In particular, efficiency of neuro systems to process computational tasks, which are challenging to traditional Turing–von-Neumann machines6 due to sequential “line-by-line” operation, inspired the development of machine learning-based neuromorphic computing (NC) and recurrent neural network (RNN) computational models7,8 as well as its subset called reservoir computing (RC).9,10 Similarly to RNN, the recurrent signal in RC constitutes a memory capable of participating in parallel computation; however, in contrast to RNN, where computationally complex algorithms are required to tune internal weights in the network, in RC, the dynamics occurs in a fixed recurrent network, and only external weights are digitally tuned, thus leading to a significant reduction of the training computation time and the size of the required memory. Consequently, various physical systems can, in principle, serve as powerful RC platforms,11 where the complex and nonlinear signal output produced by the physical system is collected and used for supervised learning to obtain the desired set of weight coefficients during the digital learning stage. Key properties of physical RC systems should satisfy sufficiently large reservoir dimensionality, allowing us to separate features of distinct states according to their dynamics, short-term (fading) memory, and a balance between sensitivity and separability. Among the different proposed physical
mechanisms, such as wave propagation, optical-based systems are particularly attractive due to their inherent parallelism, speed-of-light data propagation and processing, and relatively low energy consumption, leading to both on-chip and free-space realizations.

In this work, we employ the recently theoretically proposed light–liquid nonlinear and nonlocal interaction mechanism, where localized light-induced heating invokes thermocapillary (TC) driven deformation of an optically thin gas–liquid interface, to predict and characterize a set of new nonlinear–nonlocal effects in integrated components and then leverage these effects to achieve RC capabilities. Figure 1 presents the schematic illustration of the typical setup, where the optical mode, which propagates in the waveguide (WG) covered with a thin liquid film, invokes heating in a small gold patch on the top facet of the WG due to optical absorption, leading in turn to the TC effect and liquid deformation. In case the liquid film is sufficiently thin, liquid deformation modifies the overlap of the evanescent tail with the gas phase above the liquid film, leading to self-induced change of optical mode properties, such as phase, intensity, or coupling coefficient to another WG, which we consider below. For numerical simulations, we define silicon (Si) or silicon-nitride (SiN) channel WGs designed to carry a single mode at a wavelength 1550 nm (we consider TM polarization), which is buried in a silica (SiO₂) layer to provide a flat surface for more straightforward numerical implementation.

Our extensive three-dimensional (3D) multiphysics numerical analysis, which takes into account the full dynamics of various degrees of freedom, allows quantitative analysis beyond previous studies, which considered limiting assumptions such as small liquid deformation and effective two-dimensional (2D) approximation schemes of surface plasmon polaritons and slab WG modes. In particular, our simulations include intricate coupling, where optical dissipation in the gold patch leads to Joule heat generation and its transport to the gas–liquid interface; the latter triggers surface tension gradients along the gas–liquid interface, which, in turn, invoke TC flow accompanied by the liquid’s thickness change above the active WG [Fig. 1(b)] (in our terminology active/passive WG corresponds to presence/absence of gold patch). For the simplest case of a single active WG, the corresponding deformation of the dielectric cladding leads to a self-induced change of the accumulated optical phase, thus constituting a two-way interaction, where the optical mode affects itself through the liquid, provided the gas–liquid interface overlaps with the evanescent tail. Furthermore, a substantial in-plane spatial scale of liquid’s indentation allows us to achieve the so-called optical nonlocality, where the active optical mode affects the passive optical mode, even if the passive WG is displaced from the region of maximal optical intensity. We then employ the self-induced nonlinear phase change in a single WG or alternatively the nonlocal coupling change between adjacent WGs as a nonlinear actuator that provides

Fig. 1 Schematic description of the key components in the integrated optofluidic system under study describing the underlying mechanism of the light–liquid interaction and the memory used for RC. (a) 3D perspective presenting a box-shaped liquid cell of length $L = 10 \, \mu m$ bracketed by vertical silicone walls, and an active Si channel WG on a silica substrate covered with a gold patch of dimensions $\ell_x \times \ell_y \times k$ enabling light-induced heating and subsequent heat transport to the gas–liquid interface, which in turn triggers surface tension gradients, represented by blue arrows in (b), leading to the TC effect and liquid film thinning. The latter leads to a self-induced phase change and/or to transmittance change in the single WG setup and also to a nonlocal effect, where the phase in the adjacent passive WG is modified due to liquid deformation. Darker colors on the liquid’s surface in (a) correspond to higher temperature, whereas the dark red lines denote equal height levels. A sequence of high and low optical power pulses correspond to logic “0” and “1,” respectively. The finite relaxation time of the gas–liquid interface allows us to employ it as a short-term memory, where the $(n-1)$th pulse of power $P(n-1)$ induces $h_{z(n-1)}$ liquid thickness affecting the subsequent $n$th pulse of power $P(n)$. (b) 2D normal section, consisting of Si channel WGs of width $w = 500 \, nm$ and height $u = 220 \, nm$ buried in depth $b = 80 \, nm$ under silica top surface and hosting a gold patch of thickness $k = 20 \, nm$ and in-plane dimensions $\ell_x = \ell_y = 500 \, nm$; the initial liquid depth is $h_0 = 500 \, nm$. 

Gao et al.: Thin liquid film as an optical nonlinear-nonlocal medium...
internal feedback and also exhibits memory capability due to finite relaxation time of the gas–liquid interface, allowing us to perform digital and analog RC tasks.

2 Methods

2.1 Details of Multiphysics Simulation

To capture the complex coupling between light propagation, fluid dynamics, heat transport, and surface tension effects, we employed finite-element-based commercial software COMSOL Multiphysics\textsuperscript{287} with Wave Optics, heat transfer, and CFD module, where changes of thin liquid film geometry are simulated by the moving mesh method. Since electromagnetic dynamics occurs on much shorter time scales compared to other processes we consider, the optical simulation is performed in the frequency domain, whereas all other processes are simulated in the time domain. For the WG and thinner gold patch, we use a free tetrahedral mesh resulting in $6 \times 10^5$ mesh elements and $4 \times 10^6$ degrees of freedom. These memory demanding simulations are performed on a constructed dedicated server with 16 cores and 512 GB memory; for instance, the single self-induced phase change simulation presented in Fig. 2 takes more than 24 h simulation time. The governing equations, matching and boundary conditions, as well as key parameters used for the simulation, are provided in the Supplementary Material. In all simulations, we assume the initial liquid surface is flat and forms a contact angle with a value of 90 deg with the vertical silica boundary walls.

2.2 Details of RC Simulation

Construction of the computational scheme for optofluidic RC involved three main steps. In the first step, we employed $\sim 100$ 3D simulations, each providing gas–liquid interface deformation under an equivalent heat source (see Fig. S4a in the Supplementary Material) for a variety of different developed temperatures in the gold patch and initial configurations (see Fig. S3 in the Supplementary Material). In the second step, we extracted the corresponding data describing evolution of $h_c(t)$ and performed linear fitting to derive the following reduced phenomenological one-dimensional (1D) model for $h_c(t)$:

$$h_c(t) = \alpha(P) \cdot h_c(t) + \beta(P),$$

with power dependent coefficients $\alpha(P)$, $\beta(P)$ (see Fig. S5b in the Supplementary Material). While the first step enabled us to generate a sufficiently large dataset in a reasonable time, the second step allowed us to reduce the computation time of the RC algorithm shortly described below. Finally, we implemented the RC scheme\textsuperscript{28} using MATLAB (R2019b, Mathworks, Natick, Massachusetts)\textsuperscript{28} to simulate the optical power at the detectors in Mach–Zehnder interferometer (MZI) circuit, described in Fig. 5(a), as a function of the exciting signal. In particular, we employed the Euler method to integrate Eq. (1) with a time step of 0.1 ms and employed the relation in Fig. S5c in the Supplementary Material to derive the corresponding phase shift $\Delta \phi_{TC}$ and the accompanying optical signal at the detectors. To implement RC by employing our system, we first obtain a training matrix, without tuning the internal system properties, and then use this matrix to perform the test stage, which performs relevant computations. In particular, to accomplish the training stage, we first collected the output data in $D_{1,2}$ and arranged it in an $N_{pw} \times 2$ dimensional matrix, where $N$ is the number of bits used for training, and $p_w$ is the excitation time (which affects the number of rows in the output matrix). For instance, in our XOR simulation for training steps, we used $p_w = 25$, $N = 1000$. We then rearrange the output matrix in matrix $X = (M, N)$, where $M = 2p_w$ and each column in the matrix $X$ represents data from time step $i$ ($i = 1, \ldots, N$), and use it to solve the following linear equation for the weight matrix $W_{\text{out}}$:

$$Y = X \cdot W_{\text{out}},$$

where $Y$ is a predetermined desired output. Here, $W_{\text{out}}$ is an $M \times 1$ dimensional vector that is determined by employing Tikhonov (ridge) regression, which is expected to be executed digitally in future experimental realizations, via MATLAB’s built-in “ridge” function, which minimizes the following expression:

$$|Y - Y_p|^2 + \beta|W_{\text{out}}|^2,$$

where $\beta$ serves as a regularization parameter, and $[\ldots]$ denotes the Euclidean norm. Note that the first term in Eq. (3) penalizes large differences between the output vector $Y$ and the desired output $Y_p$, whereas the second term penalizes large weight values, which facilitates better performance.\textsuperscript{28} The corresponding closed form for $W_{\text{out}}$ is then given as

$$W_{\text{out}} = (X^T X + \beta I_N)^{-1} X^T Y,$$

where non-zero $\beta$ multiplies a unity matrix $I_N$ of dimension $N$, thus adding nonzero values to the diagonal of matrix $X^T \cdot X$ and potentially regularizing it. To implement the test stage, we feed the previously unseen driving sequence, generate new design matrix $X_{\text{test}}$, and operate on it with the previously derived $W_{\text{out}}$ via

$$Y_{\text{test}} = X_{\text{test}} W_{\text{out}}.$$
3 Results

3.1 Nonlinear Self-Induced Phase Change

Consider a single Si channel WG with integrated gold patch, covered with thin liquid film shown in Fig. 2(a). Following the mechanism described above, the gold patch serves as an optical heater, which induces TC-driven flows and deformation of the liquid dielectric film. In our simulations, we set liquid properties close to silicone oil, which is an optically transparent and highly nonvolatile liquid, thus appropriate for future experimental realizations; see Table S1 in the Supplementary Material for simulation parameters.

Figure 2(a) presents simulation results of the self-induced phase change effect due to the TM mode propagating in a single active WG covered with a thin liquid dielectric film of the initial thickness of 500 nm. In particular, Fig. 2(a) presents the corresponding deformation and temperature field of the gas–liquid interface at time $t = 20$ ms after the 0.1 mW continuous wave (CW) mode began to propagate in the WG, whereas Fig. 2(b) presents the gas–liquid deformation as a feedback response (circular arrow), allowing us to store information and affect subsequent optical pulses at the same compact spatial region without the need of dedicated large footprint optical feedback elements, e.g., optical rings. The colormap describes electric field intensity at $t = 20$ ms and thin liquid-deformation along the $x$–$z$ plane, where $x$ is the propagation direction; note the effect of light reflection from the gold patch leading to distortion of the incoming mode. Figure 2(b) presents the electric field intensity at two different times along the planes $x = -4.5 \mu$m, $x = 0 \mu$m, and $x = 4.5 \mu$m, where $x = 0 \mu$m is the center of the gold patch. Specifically, the images at the first row indicate that at $t = 0$ ms, i.e., prior to liquid film deformation, the incident mode experiences reduction in intensity but preserves its TM polarization. However, once the liquid film begins to deform, the second row indicates that after 20 ms the incident mode changes its shape due to superposition with the reflected wave from the gold patch, and the transmitted mode is no longer top-bottom symmetric. Figures 2(d)–2(g) present the self-induced phase change, deformation of the thin liquid film above

Fig. 2 Numerical multiphysics simulation results presenting self-induced phase change in a single active WG covered with a thin liquid film. (a) Deformed gas–liquid interface under 0.1-mW CW light-induced TC effect at time $t = 20$ ms (where $t = 0$ s is the time moment when the optical mode began to propagate) and (b) graphic representation of the internal optical-generated feedback (circular arrow) due to deformation of the gas–liquid interface, allowing us to invoke transient dynamics serving as memory storing the previous optical pulse and using the dynamics to affect the deformed profile and the optical pulse at the next time moments at the same actuation region. The colormap describes electric field magnitude along the $x$–$z$ plane at time moment $t = 20$ ms; the gold patch resides on the top facet of the WG, and its dimensions along the $x$-axis are bracketed by the white dashed lines. (c) Electrical field magnitude along the normal section $y$–$z$ at $x = -4.5 \mu$m, $x = 0 \mu$m, and $x = 4.5 \mu$m corresponding to left, central, and right columns, respectively, under 0.1 mW incident intensity; first and second rows correspond to mode profiles at initial time $t = 0$ ms and $t = 20$ ms, respectively. (d) The corresponding phase change as a function of different optical powers in the WG due to (e) liquid film thickness change, where $h_L$ is liquid thickness above the center of the gold patch. (f) The underlying change of the average temperature of the gold patch, $T_{C}$; (g) the corresponding transmittance as a function of time. Key parameters: initial temperature 20°C; wavelength of TM mode is 1550 nm; refractive index of liquid is 1.44 (same as silica substrate); geometric WG parameters are presented in Fig. 1.
the gold patch center \( h_c(t) \) (i.e., at region of minimal thickness just above the center of the gold patch), average temperature of the gold patch \( T_c(t) \), and the corresponding transmittance as a function of time, respectively, under various optical powers. Naturally, increasingly higher power levels and accompanying temperature gradients lead to more significant \( h_c(t) \) and the corresponding self-induced phase change effects. Interestingly, at optical powers of 0.1, 0.07, and 0.05 mW, the corresponding liquid film deformation also induces prominent transmittance change, leading to a power reduction by approximately a factor of 2/3, which we attribute to impedance mismatch created by the liquid deformation. Notably, owing to the relatively low power levels required to activate TC-driven film thinning as well as the large refractive index contrast across the gas–liquid interface, sufficiently thin liquid film with high overlap of the evanescent optical tail with the gas phase supports a few orders of magnitude higher self-induced phase change compared with that of the thermo-optical (TO) effect. In particular, Fig. S1 in the Supplementary Material indicates that the system described in Fig. 2(a) admits approximately three orders of magnitude higher self-induced phase change compared with a similar system, where instead of liquid film one employs a dielectric solid material which does not support the TC effect. It is worth mentioning that due to the moving mesh method employed in the multiphysics software,\(^{32}\) the simulation terminates once the gas–liquid interface reaches the few elements thick film, as measured from the bottom of the liquid cell, leading to a phase change of \( \sim 0.3\pi \) rad. In future experimental realizations, which are not subject to such limitation, we expect to obtain even higher values of self-induced phase change and nonlocality scales, as well as self-induced transmittance/reflection modulation, shortly described below. Furthermore, given the fact that liquid deformation is mainly concentrated around the compact gold patch, based on our study, the size of the liquid cell can be, in principle, reduced to 2 \( \mu m \), whereas higher values of self-induced phase change can be achieved by placing several gold (or other metal) patches along the WG.

3.2 Nonlocal Effect

Consider the optofluidic components schematically presented in Fig. 3 and analyze the effect of the self-induced TC-driven deformation on modes in the two WGs as a function of their distance \( d \). In particular, we consider close and far separation regimes characterized by \( d < l_e \) and \( d > l_e \), respectively, where \( l_e \) is the corresponding evanescent scale of the TM mode. To investigate the close separation regime, we focus on the directional coupler geometry\(^{27}\) with minimal distance \( d = 400 \text{ nm} \) presented in Figs. 3(a) and 3(b), whereas in the far separation regime we consider parallel WGs geometry presented in Figs. 3(d) and 3(e). Figure 3(c) shows the simulation results of the TM mode injected only into the active WG, indicating a decrease of the coupling coefficient as a function of liquid deformation, leading to increased (decreased) transmittance of the TM mode through the passive (active) WG as a function of time. In case the distance between the WGs is several microns \( (d = 1, 2, 3 \mu m) \), we can increase the nonlocal effect of the active WG on the passive WG by increasing the transverse dimension of the gold patch \( \epsilon_c \), which facilitates more efficient heat transport and enables liquid film deformation profile, which extends toward the passive WG. This approach also allows us to alleviate the simulation limitation taking place due to its termination once the liquid film reaches a thickness of a few elements. Figure 3(d) presents 3D thin liquid film deformation and the underlying temperature map for the \( d = 2 \mu m \) case, whereas Fig. 3(e) presents a normal cross section with the computational result of the corresponding modal structure. In particular, the latter presents numerical results at time \( t = 23 \text{ ms} \) of two different simulations when the light was injected into the active or passive WGs and indicates that the gold patch does not guide surface plasmon polaritons from one WG to another. Figure 3(f) presents the associated phase change in the passive WG as a function of time for three different separations, naturally indicating smaller values of phase shift for increasingly larger separations. Note that in contrast to the close separation regime, an initially low power (which is not sufficient to deform the liquid film) optical TM mode was also injected into the passive WG to analyze its evolution under the effect of the higher power mode in the active WG. Notably, the maximal separation between the WGs is much higher than the corresponding evanescent scale and also significantly exceeds other known optical nonlocality scales stemming from alternative optical nonlinearities, such as photorefraction,\(^{33}\) TO effect,\(^{34,35}\) and long range molecular interaction between liquid crystals molecules.\(^{36,37}\)

3.3 Self-Induced Transmittance and Reflection in Channel Bragg Waveguides

Consider the SiN Bragg WG on silica substrate presented in Fig. 4, where some of the ribs are covered with a gold patch of thickness \( k \). Here, we employ SiN rather than Si because the former has a lower refractive index (2 at wavelength 1550 nm) leading to larger mode volume and thus to higher sensitivity of perturbations in the geometry of the gas–liquid interface. First, we design the periodic WG to admit a stop-band at a wavelength 1550 nm when it is covered with a thin liquid film of thickness 1 \( \mu m \) (as measured from silica substrate) and admit non-zero transmittance without the thin liquid film. The evanescent penetration of the incident light a few periods into the SiN Bragg WG, which is inherent to stop-band conditions, triggers heating of the gold patches close to the input facet (left), which in turn triggers TC-driven deformation of the liquid film. As liquid thickness becomes thinner near the input facet, the designed propagation conditions facilitate deeper penetration of the optical mode into the periodic structure, thus leading to film thinning above more distant regions from the input region, which in turn amplifies the process. Indeed, Fig. 4(c) presents an initially flat gas–liquid interface at \( t = 0 \text{ ms} \), liquid deformation with indentation closer to the input facet at \( t = 8 \text{ ms} \), and wider indentation over the central part of the periodic structure at the later time of \( t = 16 \text{ ms} \), leading to almost minimally thin liquid film above all WG ribs. Similarly, a periodic structure with slightly different period can be designed to support propagation once the WG is covered with liquid film and a stop band once the liquid film is replaced by air. Following the same arguments as above, self-induced liquid thinning over the periodic structure is expected to shift the propagation conditions and lead to lower transmittance, as indeed described by the blue curve in Fig. 4(d). Interestingly, the presented effects are reminiscent of electromagnetically induced transparency, which relies on a very different phenomenon, where quantum interference allows the propagation of light through an otherwise opaque atomic medium.\(^{38}\)
3.4 Reservoir Computing of Digital and Analog Tasks

Consider Fig. 5(a) describing an MZI circuit with two directional couplers and an optofluidic cell in one of the arms, which supports the nonlinear self-induced phase change effect described above, and the circuit described by Fig. 5(c) with a liquid cell placed in the coupling region, which affects the transmittance via self-induced coupling change (nonlocal effect). In both cases, the self-induced phase/coupling change due to liquid deformation invoked by the active WG is translated into intensity changes in the output arms $D_1$ and $D_2$. To allow sufficiently large actuation of the liquid, the corresponding actuation time scale ($\tau_l$) must be smaller than pulse width ($\tau_w$), whereas to ensure memory, i.e., that the liquid deformation imprinted by the $(n-1)$th bit will affect the phase/coupling change of the $n$th bit, the distance between subsequent pulses ($\tau_f$) must be smaller than $\tau_l$. Both conditions can be summarized as

$$\tau_f < \tau_l < \tau_w;$$

(7)

and these ensure that the optically imprinted light pulse in the gas–liquid interface at time moment $n-1$ affects the liquid through the subsequent pulse at time moment $n$. Since, for an optical signal of given power and operation time, thinner film is expected to introduce more significant nonlinear response (unless it is so thin that the optical signal leads to saturation due to very fast liquid depletion above the gold patch), we choose liquid thickness in the region between 0 and $0.5 \mu m$. Below, we employ the nonlinear self-induced phase change and the self-induced coupling change (nonlinear effect) between adjacent WGs and the accompanying memory effect to perform both digital and analog tasks.

3.4.1 Digital task

Consider the delayed XOR operation, where an arbitrary input time series $\{x_n\}_{n=1}^N$ $(n = 1, 2, \cdots, N)$ of zeros and ones is mapped to output sequence $\{y_n\}_{n=1+\delta}^N$ according to

$$\{x_n\}_{n=1}^N \rightarrow \{y_n\}_{n=1+\delta}^N; \quad y_n = x_{n-\delta} \oplus x_n,$$

(8)
where $\oplus$ is the addition modulo two (i.e., XOR operation), and $\delta > 0$ is an integer encoding the delay. In particular, $\delta = 1$ corresponds to the simplest case of smallest delay, where the XOR operation is performed on adjacent bits, whereas $\delta = 2$ corresponds to the case where it is performed on bits that are separated by one bit. The corresponding dynamics of the thin liquid film, applicable irrespective of the nonlocal or nonlinear circuit in Fig. 5, admits the following evolution equation explicitly:

$$h_c(n) = f[h_c(n-1), P(n)],$$

(9)

where the state of the liquid at discrete time moment $n$ is determined by the result of the nonlinear saturation function $f$, which depends on an incident power $P(n)$ at the same time $n$ and liquid thickness at preceding time $n-1$ (see the Supplementary Material for derivation). This dependence describes the memory effect of the liquid film, where the finite (typically ms) relaxation time imprinted in the gas–liquid interface at time $n-1$ interacts with a subsequent pulse at time $n$.

To implement XOR operation, we can either employ the (nonlinear) self-induced phase change or the (nonlinear–nonlocal) self-induced coupling change with circuits Figs. 5(a) and 5(c), respectively. The input series $\{x_n\}_{n=1}^N$ is encoded as a sequence of square pulses of power level $P_n = P$ or $2P$ depending on the value of logic “0” or “1,” as described by Fig. 5(b); the corresponding power level operates during a time $\tau_w$ and is followed by relaxation time $\tau_r$. Figures 5(d) and 5(e) present the $P-\tau_r$ performance diagram with different colors encoding the corresponding prediction error, for the case of nonlinear circuit and nonlocal circuit described by Figs. 5(a) and 5(c).

---

**Fig. 4** Numerical simulation results of the self-induced transmittance and reflection effects. (a) The underlying 3D geometric setup of the periodic SiN Bragg WG on silica substrate and (b) the corresponding 2D normal cross section in the $x$–$z$ plane. Here, $a = 0.7 \mu m$, $b = 0.4 \mu m$, $c = r = 0.2775 \mu m$, $k = 20$ nm; number of ribs without gold patch $n = 4$ and with gold patch $m = 15$ for optimized performance. (c) Colormap of the corresponding power-flow along the WG as a function of time. At successively later time moments, liquid film over the periodic structure becomes increasingly thinner, thus facilitating optical propagation through the periodic structure designed to admit a stop band in the presence of liquid film. (d) Transmittance (black curve) as a function of time presenting also the transmittance values for thin liquid topography presented for the corresponding time moments in (c). The blue curve presents the decreasing function of transmittance for the complementary case of self-induced reflection, where a similar structure increasingly rejects light as liquid thickness approaches zero (see the Supplementary Material for more details as well as the video file showing the corresponding 3D liquid deformation as a function of time) (Video 1, mp4, 6.40 MB [URL: https://doi.org/10.1117/1.AP.4.4.046005.1]).
respectively. Note that the performance diagram of the nonlocal circuit presents the (white) region of vanishing error, which has a larger area compared with the corresponding area in the performance diagram of the nonlinear circuit. Furthermore, the lowest power value of the white region in the nonlinear case is \( \sim 40 \mu W \) with a lower value around \( \sim 10 \mu W \) for the nonlocal case. Higher performance of the nonlocal circuit stems from higher sensitivity of the intensity of the transmitted signal with respect to changes of the liquid thickness, compared with the nonlinear case. Specifically, sensitivity estimates can be determined by taking the derivative of the MZI signal due to a phase shift \( \varphi \), given by \( d \cos^2(\varphi/2)/dh_c = (1/2) \cdot \sin(\varphi) d\varphi/dh_c \), where the derivative \( d\varphi/dh_c \) can be estimated from Fig. S5c in the Supplementary Material; similarly, Fig. S7 in the Supplementary Material provides an estimate for the typical slope/sensitivity in the nonlocal case.

**Fig. 5** Simulation results presenting RC computing of the XOR task by employing self-induced phase change (nonlinear) and self-induced coupling change (nonlocal) effects. (a) Schematic diagram presenting the MZI circuit with two linear couplers, where the liquid cell introduces self-induced phase change in one of the arms measured by a pair of detectors D1,2, (b) encoding scheme of “0” (“1”), which employs a square wave modulation with actuation period \( \tau_w \) of power \( 2P \) (P), followed by relaxation period \( \tau_r \), and (c) nonlocal circuit, which employs self-induced coupling change and identical encoding. (d) Test results presenting performance error of the XOR task for delay parameter \( \delta = 1 \) as a function of \( P_{\text{max}} \) and relaxation time \( \tau_r \) in nonlinear circuit (a), and colormap (e) performing an identical task with nonlocal circuit (c) demonstrating the enhanced area of vanishing error. (f) Test results of the XOR task with \( \delta = 2 \) using nonlocal circuit (c) showing minimal error \( \sim 11\% \). (g) Dynamics evolution of the optical signal in detector D1 (blue) and liquid thickness \( h_c \) (red) due to the actuation signal (green) with \( \tau_w = 25 \text{ ms} \), \( \tau_r = 5 \text{ ms} \), and \( P = 0.033 \text{ mW} \). (h) Folded dynamics of optical intensity in the nonlocal circuit and (i) nonlocal circuit showing enhanced separability of the different regimes.
prominent memory of only one time backward in time. The underlying dynamics of the thin liquid film was obtained by constructing a reduced 1D model, which was obtained from analyzing numerous 3D multiphysics simulations results, where the thin liquid film was subjected to various initial conditions and actuation powers. In our reduced model, the transient dynamics of the thin liquid film is subject to the first-order ordinary differential equation with power dependent coefficients, allowing a simple numerical scheme to predict the accompanying phase/coupling change and the resultant power levels at the detectors (see Sec. 2 and the Supplementary Material). In particular, Figs. S3, S5a, and S5c in the Supplementary Material present \( h_1(t) \) as a function of time for several different initial thicknesses and input powers, the relation between \( h_1(t) \) and \( h_2(t) \) and the one-to-one relation between \( h_1(t) \) and the induced phase shift, respectively, whereas Fig. S7 in the Supplementary Material presents transmittance values as a function of optical power in the active WG. Figure 5(g) presents the underlying dynamics of the thin liquid film (red curve) and the output signal (blue curve) in the D1 detector in the nonlocal circuit [Fig. 5(c)], under an exciting signal (green curve). To visually analyze the different patterns of detected power, we construct folded dynamics diagrams, where the optical signals at different time intervals of length \( T = \tau_e + \tau_w \) are plotted on top of each other. Figures 5(h) and 5(i) present folded dynamics graphs for the nonlinear and nonlocal circuits between times \( T \) to \( 200T \), respectively, where the signal in the nonlocal circuit shows enhanced separability compared to the nonlinear case performed under similar conditions \( \tau_e = 25 \text{ ms}, \tau_w = 5 \text{ ms}, \) and \( P = 33 \mu \text{W} \). Interestingly, Figs. 5(h) and 5(i) do not show dependence on initial conditions, which is in concert with the graph in Fig. S6a in the Supplementary Material, demonstrating that the system practically “forgets” the initial conditions after \( 5T \) and, furthermore, showing the emergence of patterns that are characteristic of particular dynamics and actuation signals. For instance, Fig. 5(i) presents four main curves with an internal substructure, corresponding to “1” or “0,” which was preceded by “1” or by “0.” These represent the four dominant combinations, which allow an efficient solution of the delayed (\( \delta = 1 \)) XOR task, whereas the internal structure represents next order memory effects, which allows us to some extent to solve the \( \delta = 2 \) XOR task. More formally, since the dynamics of our system converges to states that do not depend on the initial conditions, it satisfies the so-called common-signal-induced synchronization (see Ref. 40 and references within), also known as echo state,\(^4\) which is a necessary property enabling a dynamical system to serve as an RC. To determine the required optical energy \( E \) needed for training, we assume that the number of “0” and “1” pulses is equal, leading to \( E = (NP/2 + NP)\tau_w \); for instance, \( N = 1000, P = 10 \mu \text{W}, \) and \( \tau_w = 20 \text{ ms} \) yields \( E = 0.1 \text{ J} \). Similarly, the energy during the test stage \( E_{\text{test}} \), required for a solution of a string of length \( N_{\text{test}} \), is given by \( E_{\text{test}} = (N_{\text{test}}P/2 + N_{\text{test}}P)\tau_w \). The case with \( N_{\text{test}} = 20 \) and similar parameters used during the training stage discussed above yields \( E_{\text{test}} = 6 \mu \text{J} \).

### 3.4.2 Analog task

Next, we consider the classification task of hand-written “zero” and “one” digits; Fig. 6(a) presents a sample of 20 images from Modified National Institute of Standards and Technology (MNIST) database,\(^4\), each of the size 28 × 28 pixels. To encode each image as a signal, without filtering or other preprocessing procedures, we partitioned the image into a set of horizontal rows or columns, where each pixel is represented by a 1-ms long constant optical power proportional to the brightness value of the corresponding pixel. Specifically, the pixels are encoded according to power values \( cP_{\text{max}} \), where \( P_{\text{max}} \) is the power needed to encode the brightest pixel of unity brightness (\( c = 1 \)), whereas \( c = 0 \) corresponds to dark pixels that carry no features, and other values \( 0 < c < 1 \) correspond to pixels of intermediate brightness. Figure 6(b) presents an example of a hand-written “one” digit used for classification, where the 28 ms long signal encoding the highlighted red horizontal row is given by Fig. 6(c) with left/right pixels corresponding to early/late times. To perform image classification, we employ either the nonlinear or nonlocal circuits presented in Figs. 5(a) and 5(c), respectively, where the top and the bottom arms receive in parallel the 1-ms long bits encoding the \( i \)th and the following (\( i + 1 \))th rows, respectively, thus generating more correlations between different lines and leading to higher accuracy compared to feeding just one arm at a time. Figure 6(d) presents the classification error
during the training and testing stages, with training values naturally admitting lower values. Interestingly, both errors admit fairly low values below 0.5% across a wide range of power values \( P_{\text{max}} \) spanning from 0.01 to 0.1 mW. The total energy needed to feed one image is determined by \( 28 \times 28 \times 1 \text{ ms} \times P_{\text{max}} \times \tau \), where \( \tau \) is the average brightness level of the image laying between 0 < \( \tau \) < 1 and typically admitting value \( \sim 0.15 \). The corresponding energy for the training/testing stage is achieved by multiplying the one image energy above by the total number of corresponding images (see Sec. 2); for \( P_{\text{max}} = 10 \ \mu \text{W} \), the corresponding average power level is given by \( \tau P_{\text{max}} = 1.5 \ \mu \text{W} \). Importantly, the single liquid cell circuits in Figs. 5(a) and 5(c) can be extended to form a more complex network. For instance, Fig. S9 in the Supplementary Material presents a network consisting of three inputs, two liquid cells, and two detectors. While it does not present enhancement in accuracy, it allows us to reduce the computation time by a factor of two without increasing the total required energy. Analyzing networks with more inputs, allowing us to achieve optimization of the handwritten digits recognition task and further reduction in operation time is beyond the scope of this work.

To learn more about the role of the reservoir in our modality, we also performed linear regression analysis, where the output signal is made equal to the input signal, which is followed by ridge regression, allowing us to obtain the corresponding weights that perform the corresponding task with corresponding error given by 0.61%. Interestingly, employing a reservoir presented in Fig. 5(a) but without liquid response yields similar 0.61% performance for “zero/one” digits classification, which is around 4.3 times higher compared to the highest 0.14% reservoir performance for this task. It is worth mentioning that the performance described in Fig. 5(d) and Fig. S8 in the Supplementary Material is not a prominent function of the optical power and using larger power does not lead to significantly enhanced performance. While in this work the input data used for the analog task were not subject to any preprocessing, we expect that common methods such as edge detection should improve the accuracy.

4 Discussion

To summarize, we believe that the family of physical effects predicted to take place due to nonlinear–nonlocal interaction between thin liquid film and channel WG modes in integrated chip-scale photonics platforms, and the emergent concept suggested to employ these effects to realize optofluidic RNN supporting RC applications, will stimulate both pure light–matter interaction studies and development of RNN architectures/ schemes. In particular, our 3D simulation results, which take into account optical propagation with losses, heat transport, and fluid dynamics as well as the underlying surface tension physics, indicate that the magnitude of the self-induced phase change due to the TC effect in the liquid film is approximately three orders of magnitude higher compared to the more common heat-based TO effect. Therefore, the predicted effects of self-induced coupling change and self-induced bandstructure and transmittance change can stimulate experimental and theoretical studies validating the predictions and exploring bandstructure tuning in photonic crystals and their topological properties, as well as studying nonlocal interaction between a large number of WGs, even if their separation exceeds the optical evanescent scales. While in our work, we employed a gold patch due to its optical absorption properties, studying metal structures designed to support resonant absorption due to surface plasmon polariton oscillations is another intriguing possibility capable of inducing and modifying resonant absorption in channel WGs, as we briefly mention in Fig. S11 in the Supplementary Material. From the ML-based perspective, the employed self-induced phase/coupling change allows us to achieve nonlocal–nonlinear interaction and memory at the same compact actuation region and perform the digital XOR and analog handwritten images classification tasks, which can lead to new RNN elements and architecture that take advantage of the built-in internal feedback capability. Notably, our work demonstrates for the first time that controlled deformation of the gas–liquid interface on the submicron scale can operate as an RNN, which can be extended to a network with a larger number of liquid cells (or a larger number of WGs in a single liquid cell) and further provides design principles of compact optofluidic and complementary metal–oxide–semiconductor compatible optofluidic RC systems, which are capable of inducing additional nonlinearity beyond the commonly employed square-law detection, without utilizing optical resonant structures, and furthermore operates under a wide range of conditions that do not require phase matching conditions. In particular, our design of an optofluidic-based RC system is capable of leading to 4–5 orders of magnitude reduction in size compared to previous liquid-based RC systems and also to a few orders of magnitude faster computation compared to the surface waves dynamics and reaction-diffusion processes, which were employed in Refs. 44 and 45, respectively. While the presented optofluidic approach for RC requires more energy compared to the present state-of-the-art approaches such as the opto-electronic based method, offering \( \sim 10 \text{ GHz} \) modulation rate with hundreds of fJ per bit \(^4\) (see Table S2 in the Supplementary Material), we should keep in mind that the electro-optic approach requires an external modulation unit, whereas in the optofluidic approach the modulation is internal due to the specificity of the gas–liquid interaction. In particular, internal feedback allows us to invoke nonlinearities and memory effects in the interaction region and thus translates into a very compact computation region without feedback loops, which are required in systems that employ external feedback. It is not clear how to achieve a self-induced refractive index response of the order \( O(10^{-1}) \) with the electro-optic effect, though self-induced modulation can be achieved by employing two-photon absorption and other nonlinearities \(^3\) using significantly higher power levels compared with state-of-the-art externally modulated microring resonator devices.\(^46\) Furthermore, we should keep in mind that neural brain activity also occurs on a ms time scale and is based on several ionic transport processes, which also take place in a liquid environment. Therefore, even if our study in its current version cannot suggest computational capabilities comparable to that of a human brain, the general strategy of combining several different physical processes rather than relying on a small number of physical effects may prove to be a useful strategy to achieve more efficient computational approaches. It is worth mentioning that while phase-change materials, where optically induced phase change between crystalline and amorphous phases invokes self-induced transmittance change, were recently employed for NC applications,\(^47,48\) the corresponding material phase change is not able to nonlocally affect nearby WGs. Given the fact that in this work we demonstrated that nonlocality enhances performance of both digital and analog tasks, we believe that our study will motivate future studies with more
efficient optofluidic-based architectures that employ nonlocality effects and, in parallel, also stimulate material science-oriented studies on how nonlocality can be harnessed in more conventional phase-change materials. It is worth noting that we performed preparatory experiments demonstrating controlled deposition of nonvolatile liquid into etched trenches above a silicon channel WG, bringing future experimental realization of the predicted effects within a closer reach (see Fig. S10 in the Supplementary Material).

Acknowledgments

The authors have no relevant financial interests in the article and no other potential conflicts of interest to disclose. The authors cordially thank the Defense Advanced Research Projects Agency (DARPA), Nature as Computer (NAC) program management team, for stimulating discussions during the program. This work was supported by the DARPA Defense Sciences Office NAC (HR00112090009) and NLM Programs, the Office of Naval Research (ONR); the National Science Foundation (NSF), grants CBET-1704085, DMR-1707641, NSF ECCS-180789, NSF ECCS-1901884, NSF ECCS-2023730; the Army Research Office (ARO); the San Diego Nanotechnology Infrastructure (SDNI), supported by the NSF National Nanotechnology Coordinated Infrastructure (grant ECCS-2025752); the Quantum Materials for Energy Efficient Neuromorphic Computing – an Energy Frontier Research Center, funded by the U.S. Department of Energy (DOE) Office of Science, Basic Energy Sciences, under award #DE-SC0019273; and the Cymer Corporation.

References

1. M. Borghi et al., “Nonlinear silicon photonics,” J. Opt. 19(9), 093002 (2017).
2. B. Corcoran et al., “Green light emission in silicon through slow-light enhanced third-harmonic generation in photonic-crystal waveguides,” Nat. Photonics 3(4), 206–210 (2009).
3. A. L. Gaeta, M. Lipson, and T. J. Kippenberg, “Photonic-chip-based frequency combs,” Nat. Photonics 13(3), 158–169 (2019).
4. J. W. Silverstone et al., “On-chip quantum interference between silicon photon-pair sources,” Nat. Photonics 8(2), 104–108 (2014).
5. G. Van der Sande, D. Brunner, and M. C. Soriano, “Advances in photonic reservoir computing,” Nanophotonics 6(3), 561–576 (2017).
6. J. Hasler and H. B. Marr, “Finding a roadmap to achieve large neuromorphic hardware systems,” Front. Neurosci. 7, 118 (2013).
7. H. Jaeger, “The echo state approach to analysing and training recurrent neural networks,” in Photonic Reservoir Computing, Vol. 4(4), 337 (1871).
8. J. R. A. Pearson, “On convection cells induced by surface tension,” J. Fluid Mech. 4(5), 489–500 (1958).
9. V. G. Levich, Physicochemical Hydrodynamics, Prentice Hall (1962).
10. H. M. J. M. Wedershoven et al., “Infrared laser induced rupture of thin liquid films on stationary substrates,” Appl. Phys. Lett. 104, 054101 (2014).
11. S. Rubin, B. Hong, and Y. Fainman, “Nonlinear, tunable, and active optical metasurface with liquid film,” Light: Sci. Appl. 8, 177 (2019).
12. www.comsol.com.
13. M. Lukoševičius, A practical guide to applying echo state networks, in Neural Networks: Tricks of the Trade, G. Montavon et al., Eds., pp. 659–686, Springer, Berlin, Heidelberg (2012).
14. The MathWorks Inc., Natick, MATLAB version 9.7.0 (R2019b), Massachusetts.
15. S. Chatterjee and A. S. Hadi, Regression Analysis by Example, John Wiley & Sons (2013).
16. C. Mesaritakis, V. Papatxiarihs, and D. Syrvidis, “Micro ring resonators as building blocks for an all-optical high-speed reservoir-computing bit-pattern-recognition system,” J. Opt. Soc. Am. B 30(11), 3048–3055 (2013).
17. G. Lifante, Integrated Photonics: Fundamentals, John Wiley and Sons, Chichester (2003).
18. G. C. Duric, Jr. et al., “Observation of self-trapping of an optical beam due to the photorefractive effect,” Phys. Rev. Lett. 71(4), 533–536 (1993).
19. C. Rotschild et al., “Solitons in nonlinear media with an infinite range of nonlocality: first observation of coherent elliptic solitons and of vortex-ring solitons,” Phys. Rev. Lett. 95(21), 215904 (2005).
20. J. Komma et al., “Thermo-optic coefficient of silicon at 1550 nm and cryogenic temperatures,” Appl. Phys. Lett. 101(4), 041905 (2012).
21. D. W. McLaughlin et al., “A paraxial model for optical self-focusing in a nematic liquid crystal,” Physica D 88(1), 55–81 (1995).
22. C. Conti, M. Peccianti, and G. Assanto, “Route to nonlocality and observation of accessible solitons,” Phys. Rev. Lett. 91(7), 073901 (2003).
23. G. Assanto and M. Peccianti, “Spatial solitons in nematic liquid crystals,” IEEE J. Quantum Electron. 39(1), 13–21 (2003).
24. M. Inubushi et al., “On the characteristics and structures of dynamical systems suitable for reservoir computing,” in Reservoir Computing: Theory, Physical Implementations, and Applications,
K. Nakajima and I. Fischer, Eds., pp. 97–116, Nature Computing Series, Springer, Singapore (2020).
41. Y. LeCun, “The MNIST database of handwritten digits,” http://yann.lecun.com/exdb/mnist/ (1998).
42. L. Lu, J. D. Joannopoulos, and M. Soljačić, “Topological photonics,” Nat. Photonics 8(11), 821–829 (2014).
43. T. Ozawa et al., “Topological photonics,” Rev. Mod. Phys. 91(1), 015006 (2019).
44. C. Fernando and S. Sojakka, “Pattern recognition in a bucket,” Lect. Notes Comput. Sci. 2801, 588–597 (2003).
45. A. Adamatzky and B. D. L. Costello, “Experimental logical gates in a reaction-diffusion medium: the XOR gate and beyond,” Phys. Rev. E 66(4), 046112 (2002).
46. J. Sun et al., “A 128 Gb/s PAM4 silicon microring modulator with integrated thermo-optic resonance tuning,” IEEE J. Lightwave Technol. 37(1), 110–115 (2019).
47. J. Feldmann et al., “All-optical spiking neurosynaptic networks with self-learning capabilities,” Nature 569(7755), 208–214 (2019).
48. C. Rìos et al., “Integrated all-photonic non-volatile multi-level memory,” Nat. Photonics 9(11), 725–732 (2015).
49. M. Wuttig, H. Bhaskaran, and T. Taubner, “Phase-change materials for non-volatile photonic applications,” Nat. Photonics 11(8), 465–476 (2017).

Chengkuan Gao received his BSc degree in physics from Wuhan University in 2017 and his MSc Eng. in materials science from University of Pennsylvania in 2019. He is currently a PhD student in the Ultrafast & Nanoscale Optics (UNO) Group, majoring in photonics at the Department of Electrical and Computer Engineering (ECE) at University of California, San Diego.

Prabhav Gaur received his BTech degree in electrical engineering from the Indian Institute of Technology Kanpur in 2019 and his MS degree in electrical and computer engineering (photons option) from University of California, San Diego (UCSD) in 2021. Currently, he is pursuing his PhD in electrical and computer engineering (photons option) from the UCSD, and his research interest includes optical signal processing and optical metrology.

Shimon Rubin received his BSc and MSc physics degrees from Technion – Israel Institute of Technology in 2001 and 2004, respectively, and his PhD physics degree from Ben-Gurion University of the Negev in 2012. Between 2013 and 2016 he served as a postdoctoral fellow in the Department of Mechanical Engineering at Technion, and in 2017 he joined as a postdoctoral researcher and later as assistant project scientist in ECE at UCSD.

Yeshaiahu Fainman received his MSc and PhD degrees from Technion, Israel, in 1979 and 1983, respectively. He joined the ECE Department at UCSD in July 1990, following a faculty appointment at the University of Michigan, and now serves as a Cymer chair and distinguished professor, directing the Ultrafast and Nanoscale Optics Group in research of nanophotonics. He has contributed over 280 manuscripts and 450 conference proceedings and is a fellow of OSA, IEEE, and SPIE, and a recipient of numerous prizes and awards.