



Nanoscale

Pathway of Transient Electronics towards Connected Biomedical Applications

Journal:	<i>Nanoscale</i>
Manuscript ID	NR-MRV-10-2022-006068.R1
Article Type:	Minireview
Date Submitted by the Author:	19-Dec-2022
Complete List of Authors:	Dutta, Ankan; The Pennsylvania State University, Engineering Science and Mechanics Cheng, Huanyu; Pennsylvania State University, Engineering Science and Mechanics

SCHOLARONE™
Manuscripts

Pathway of Transient Electronics towards Connected Biomedical Applications

Ankan Dutta¹, Huanyu Cheng^{1*}

¹Department of Engineering Science and Mechanics, The Pennsylvania State University, University Park, 16802, USA

*To whom correspondence should be addressed: E-mail: huanyu.cheng@psu.edu

Abstract

Transient electronic devices have shown promising implications in hardware security and medical implants with diagnosing therapeutics capabilities since their inception. The control of the device transience allows the device to “dissolve at will” after its functional operation, leading to the development of on-demand transient electronics. This review discusses the recent developments and advantages of triggering strategies (e.g., electrical, thermal, ultrasound, and optical) for controlling the degradation of on-demand transient electronics. We also summarize bioresorbable sensors for medical diagnoses, including representative applications in electrophysiology and neurochemical sensing. Along with the profound advancement in medical diagnosis, the commencement of therapeutic systems such as electrical stimulation and drug delivery for the biomedical or medical implant community has also been discussed. However, implementing a transient electronic system in real healthcare infrastructure is still in its infancy. Many critical challenges still need to be addressed, including strategies to decouple multimodal sensing signals, dissolution selectivity in the presence of multiple stimuli, and a complete sensing-stimulation closed-loop system. Therefore, the review future discusses opportunities in transient decoupling sensors and robust transient devices that are selective to a particular stimulus and act as hardware-based passwords. Recent advancement in closed-loop controller-enabled electronics has also been analyzed for future opportunities of using data-driven Artificial Intelligence-powered controllers in fully closed-loop transient systems.

Keywords: transient electronics, medical implants, closed-loop system, electrical stimulation and drug delivery, medical diagnosis

1. Introduction

Bioelectronics has been an integrated part of medical implants such as artificial pacemakers. However, surgical interventions are often performed to remove the implants to avoid adverse long-term consequences after the functional operation of the implants. The recent development of medical implants with on-demand transience has largely overcome post-surgical complications. Therefore, transient electronics are widely employed in medical implants for biomedical diagnosis^{1–12}. Besides sensing and diagnosis, bioresorbable electronics have also been implemented for biomedical therapeutics^{13–21}. With the need to use advanced transient systems in real-world healthcare infrastructure, control of their operational time becomes essential. Wirelessly triggered transience using an external stimulus such as electrical, optical, or ultrasound has recently been leveraged to dissolve the implant at a controlled rate and prescribed time, thereby controlling the implant's operational time^{22–28} (**Figure 1a**). Moreover, introducing an intelligent closed-loop controller is imperative to elevate the transient system to an advanced intelligent system for minimal clinician intervention on large-scale applications^{17,18,29–32}. As a first step to address this need, closed-loop systems with bioresorbable sensors and actuators have been recently developed to sense and actuate on demand³³. However, gaps and possible directions in truly data-driven intelligent, robust transient electronic systems are yet to be reported.

Traditional implants are fabricated using non-bioresorbable materials showing near-constant performance until their infinite operational time (**Figure 1b-i**), requiring secondary surgery. On the other hand, transient electronics leverages bioresorbable substrates such as poly lactic-co-glycolic acid (PLGA)^{34–46} and metals such as magnesium and zinc^{47–51} fabricated with standard or unconventional techniques (e.g., light-induced zinc mass transfer)⁵² to enable transience. The material selection, design, and external stimulus mode are crucial to determining its operational time ranging from seconds to months. However, passive transient systems have a destined degradation period, leaving no control over their operational time trajectory. On the contrary, active or on-demand transient systems could use an external triggering stimulus to initiate the onset of degradation^{22–27,34,35} (**Figure 1b-ii**). The control over the onset of transience allows the device to “dissolve at will” after its designed usage in both the inpatient and outpatient settings. The new dimension in the home monitoring/treatment reduces the hospital burden^{24,30,33} (**Figure 1c**).

This review discusses the advancement in different strategies and their mechanisms underlying the dissolution dynamics of on-demand transient electronics. Along with the advancement in the various diagnostic applications enabled by transient electronics, the review also discusses a few initial steps toward transient therapeutics such as electrical stimulation and drug delivery. By incorporating transient therapeutics with sensing capabilities, transient closed-

loop systems and their possible advancement to provide autonomous biomedical therapy from recent studies are also reviewed. Lastly, limitations and challenges have been discussed, which may provide tangible pathways to implement fully robust closed-loop transient electronics in practical healthcare infrastructure.

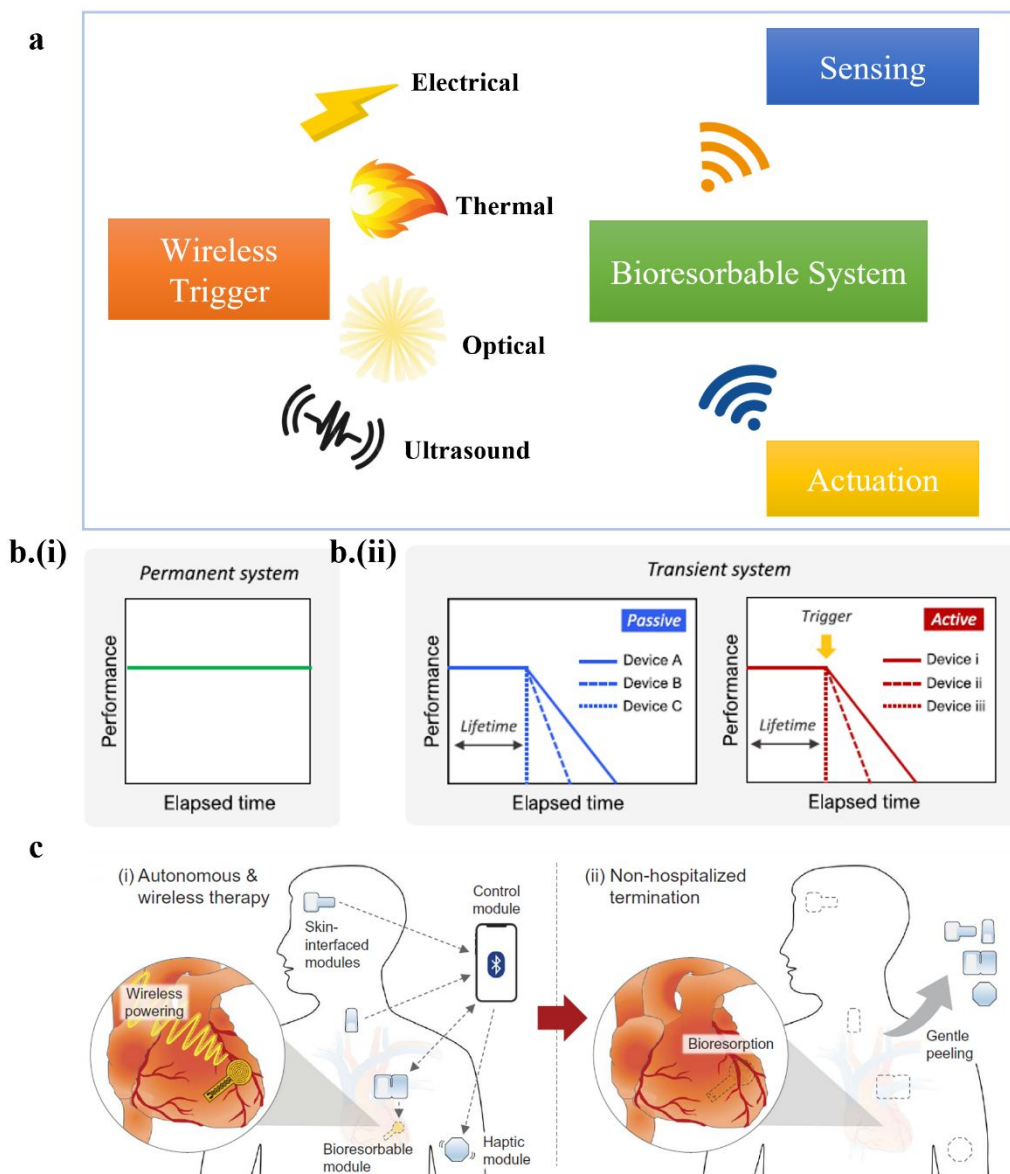


Figure 1: (a) Schematic showing on-demand transient electronics in biomedical sensing and actuation. (b) Comparison in the strategies between (i) permanent system and (ii) passive or active transient electronics. This figure has been reproduced from ref.²⁴ with permission from Elsevier, copyright 2020. (c) Advantage of active transient electronics in a clinical setup. This figure has been reproduced from ref.³³ with permission from The American Association for the Advancement of Science, copyright 2022.

2. Strategies of Wirelessly Triggered Transient Electronics

Degradation of materials commences immediately with the introduction of water or biofluids such as phosphate-buffered saline (PBS) solution in passive transient systems, leading to its destined operational failure^{24,34,35,38,39,42,47,53,54}. In contrast, materials degrade only in the presence of an external stimulus in active transient systems, enabling control over the operation timeline of the device^{14,22–25,27,55,56}. The transience onset and rate can be conveniently controlled by the device structure (e.g., thickness) and constituent materials, as well as the degradation mechanism of stimuli-responsive material, to modulate the transient process and operation timeline^{53,54}. The dissociation of one or various bioresorbable components within the device can initiate the degradation process. On the other hand, the device can also disintegrate due to the complete degradation of the substrate supporting the device components²⁴. The material dissociation can be triggered by various external stimuli, ranging from etchants and temperature to ultrasound. However, triggering stimuli such as water or solvents/etchants applied physically in contact with the dissociating material are challenging to use in a wirelessly controlled setting. On the other hand, light³ or ultrasound²³ can be applied externally and controlled wirelessly to trigger or initiate the dissociation of the material in the surrounding solution. The following subsections focus on the degradation mechanism and strategies exploited to trigger device disintegration wirelessly by external stimuli.

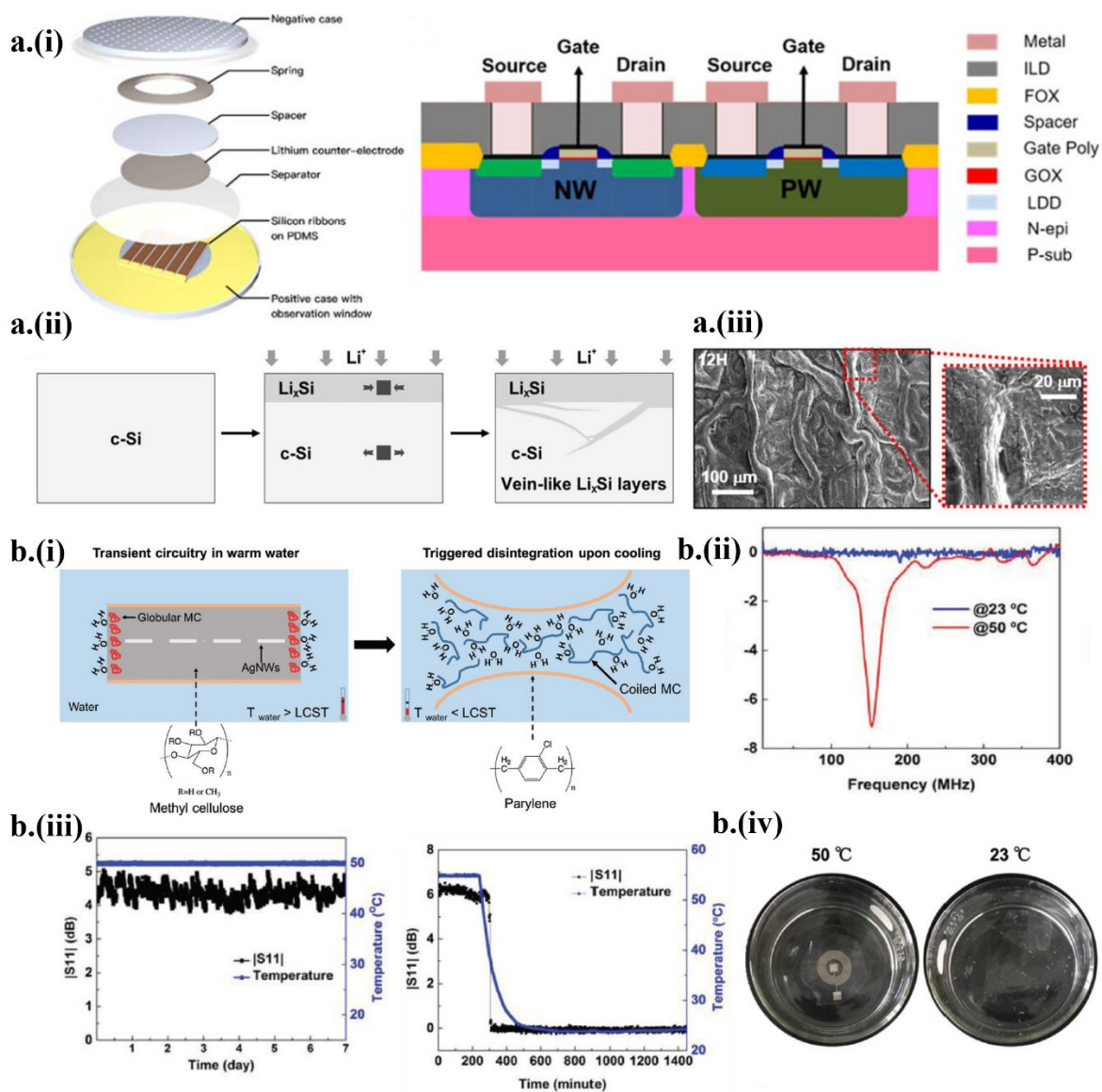


Figure 2. Strategies of wirelessly triggered on-demand transient electronics by (a) electrical or (b) temperature stimuli. (a) (i) Electrically triggered on-demand transient electronics based on Si nanomembrane. (ii) The morphological transformation of Li_xSi layers upon lithiation. (iii) Morphology of the Si side of the MOSFETs after 12 h of lithiation. This figure has been reproduced from ref.²⁷ with permission from IOP Publishing, Ltd., copyright 2019. (b) Triggered by the temperature change, (i) the methylcellulose substrate dissolves when the solution temperature drops below the polymer's lower critical solution temperature (LCST). (ii-iii) Measured S_{11} return loss in warm and cold water and (iv) demonstration of transience in cold water. This figure has been reproduced from ref.⁵⁵ with permission from John Wiley and Sons, copyright 2019.

2.1 Design strategy and mechanism of electrically triggered transient electronics

Electrical stimuli initiate fast and controllable triggering events in transient systems. Typically, electrical energy can be provided wirelessly to either generate an electrical spark to release corrosive agents or enable electrochemical reaction-assisted triggering of the device disintegration into the surrounding fluid²⁴. Though the onset of triggering is controllable, the prior does not provide any control over the dynamic releasing process of corrosive agents, resulting in a destined disintegration of the device. In contrast, the disintegration due to electrochemical reaction is controllable as it depends on the generated bias to facilitate chemical kinetics^{14,53,54,57}. Understanding the transience of silicon is critical to advance transient circuitries by leveraging the existing silicon-based electronics. However, accelerating the dissolution of Si nanomembranes for fast transient Si circuits is a challenge. With the application of a constant current density of $400 \mu\text{A cm}^{-2}$ in galvanostatic mode using a metal-oxide-semiconductor field-effect transistor (MOSFET), lithiation was performed to provide a fast yet controllable on-demand disintegration of Si⁵⁷ (**Figure 2a-i**). Inhomogeneous diffusion of lithium (Li) into Si formed a vein-like Li_xSi network and increased its volume by $\sim 280\%$, which induced microcracks to accelerate the degradation of Si (**Figure 2a-ii, iii**).

2.2 Design strategy and mechanism of thermally triggered transient electronics

It is intuitive to utilize water as a solvent to dissolve the substrate and as a reactant to initiate hydrolysis of the device's constitutive materials in water-triggered transient electronics^{22,35,55}. However, the water or other etchants often need to be in contact with the device to initiate hydrolysis and are limited in the application of passive triggering transient electronics. In order to improve wireless controllability, it is imperative to combine external parameters such as the temperature of the water as wirelessly triggered stimuli to control the triggering events. As a representative example, the hydrophilicity of a thermoresponsive polymer such as methylcellulose (MC) with a lower critical solution temperature (LCST) of 50°C was leveraged in a transient radio frequency (RF) antenna⁵⁵ (**Figure 2b-i**). For temperatures above LCST, the hydrogen bonding between water molecules and polar groups of the polymer prevents random mixing, decreasing the entropy and thus favoring unspontaneous mixing due to an increase in Gibb's free energy, which induces the hydrophobic nature of MC. At a temperature below LCST, the MC polymer becomes hydrophilic with the decreased formation enthalpy due to the formation of more hydrogen bonds, thus favoring spontaneous mixing. As a result, the antenna showed excellent mechanical and electrical stability in a warm PBS with a temperature higher than LCST (**Figure 2b-ii**), but it

degenerated and dissolved when the solution cooled down to be lower than LCST (**Figure 2b-iii**), resulting in physical disappearance of the antenna (**Figure 2b-iv**).

2.3 Design strategy and mechanism of ultrasonically triggered transient electronics

The rise of transient implantable devices has spurred the development of various transient powering sources. The implants are generally powered by batteries, taking the bulk of the device's weight. In addition, the transient battery is still in its infancy, suffering from the trade-off between the electrochemical performance and transient behavior^{58,59}. Therefore, wireless powering systems such as ultrasound become attractive for transient implantable devices^{23,60}. However, the tissue attenuation often results in drastically reduced power transmitted to the implantable. In a recent demonstration²³, ultrasound-based acoustic energy transmission was harnessed both as wireless powering and a triggering mechanism to power implantable triboelectric nanogenerator (TENG) (**Figure 3a**). The TENG was designed with a poly(3-hydroxybutyrate-co-3-hydroxyvalerate)/polyethylene glycol (PHBV/PEG) composite membrane on top of an Mg electrode layer sandwiched between two PHBV layers (**Figure 3a-i**). The implantable TENG showed a stable energy generation of 338.7 mV root mean square (RMS) using a 0.5 W cm⁻², 20 kHz ultrasound probe (**Figure 3a-ii**). After 60 mins of operation, an increased acoustic power density of 3 W cm⁻² from the ultrasound probe triggered the onset of the transient process to result in a sudden reduction in the generated voltage (**Figure 3a-iii**). Thus, controlling the ultrasound intensity allowed the TENG to power (<1 W cm⁻²) or initiate degradation (>3 W cm⁻²) (**Figure 3a-iv**).

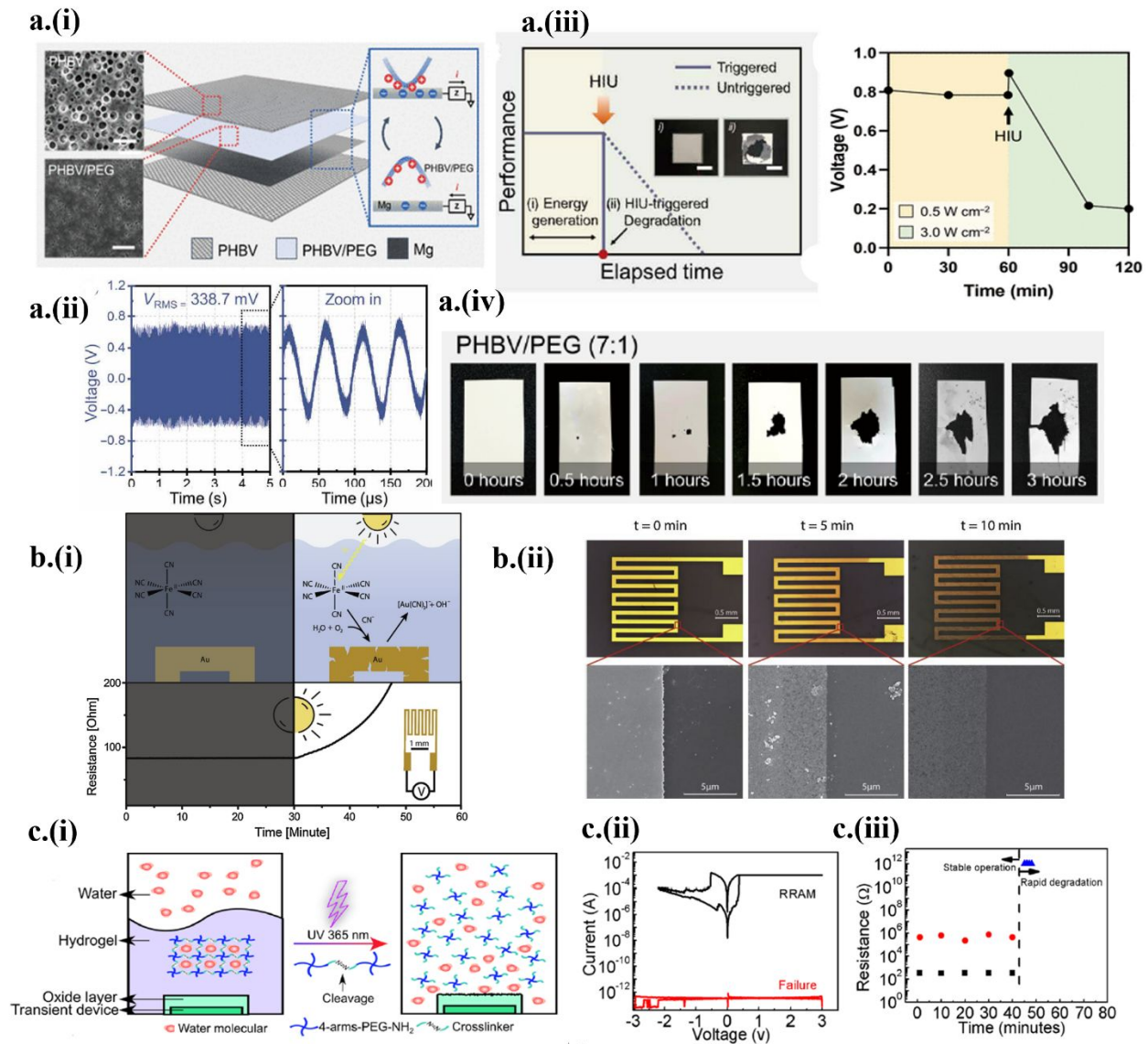


Figure 3: Wirelessly triggered on-demand transient electronics by ultrasound and light. (a) (i) Working principle of ultrasound-triggered TENG based on fully bioresorbable PHBV, PHBV/PEG, and Mg, with field-effect scanning electron microscopy (FE-SEM) images of PHBV and PHBV/PEG shown in the inset (scale 20 μm). (ii) Generated voltage from the TENG in *ex vivo* experiment. (iii) Voltage generated before and after the ultrasound-powered triggering event. (iv) Demonstration of different degradation stages during the triggering event. This figure has been reproduced under the terms of the Creative Commons CC BY NC License.²³ Copyright 2022, The American Association for the Advancement of Science. (b) (i) Schematic showing the release of photo-initiated cyanide and gold cyanidation, which increases the resistance after UV triggered event. (ii) Demonstration of transience and SEM image at 0, 5, and 10 min. This figure has been reproduced from ref.⁵⁶ with permission from Elsevier, copyright 2019. (c) (i) Encapsulation design of hydrogel-oxide bilayer for controlled UV-triggered degradation. (ii) Electrical characterization of RRAM before and after the triggering

event (failure). (iii) Investigation of degradation of RRAM without a light trigger. This figure has been reproduced from ref.³ with permission from American Chemical Society, copyright 2018.

2.4 Design strategy and mechanism of optically triggered transient electronics

Besides ultrasound, photo-triggerable materials (e.g., photo-acid generators or photo-induced phase transition materials) have also been recently leveraged in on-demand transient electronics systems^{3,22,35,56,61}. For instance, light-induced free cyanide species produced from the solar simulator oxidized the Au surface⁵⁶ (**Figure 3b-i**). The light-activated Au surface degradation increased with the time of light exposure, as observed in the increased resistance (**Figure 3b-ii**). In another work, UV (365 nm, 300 mW cm⁻²) acted as a triggering external stimulus for a light-responsive hydrogel-oxide bilayer to undergo a gel-to-sol phase transition³ (**Figure 3c-i**). The hydrogel layer acted as a transparent encapsulation of the Mg-based Resistive Random Access Memory (RRAM) device, where resistive switching between ON (conducting) and OFF (insulating) (ratio of >10²) was achieved using oxygen vacancies-based conductive filaments (**Figure 3c-ii**). The triggering of UV collapsed the hydrogel network, allowing water to pass and dissolve Mg-based RRAM (**Figure 3c-iii**). The advent of transient memory devices could add an extra dimension to hardware security and zero electronics waste.

3. Implantable Transient Electronics in Diagnostic Applications

By avoiding secondary surgical complications, the implantable devices with the transient nature have unique application opportunities in biomedical diagnostics^{22,35,61–63}, including the sensors to monitor the intracranial pressure^{1,34,62} and dopamine^{5,34,63}. The ideal bioresorbable implants must maintain a user-defined operational time with unchanged sensing performance for diagnosis. However, it is challenging to maintain stable sensing performance while controlling the degradation characteristics in a transient system^{24,34,63,64}. A few representative transient-based diagnostic devices from recent studies are discussed in the following.

3.1 Transient electrodes for electrophysiology and temperature/pressure sensors

As an essential biomarker for neuronal disorders (e.g., epilepsy and Alzheimer's disease) and diagnosing irregular cardiac rhythms^{1,5,6,35,62}, electrophysiology from transient transistor-based neural electrode arrays can provide high-resolution spatiotemporal brain mapping. Passive transient systems typically exhibit degraded electrical performance with gradually increased bulk resistance or contact impedance over time^{6,47,48,65}. To address this challenge, an

electrocochleography (ECoG) array combined with an intracranial pressure sensor was fabricated with a Mo electrode embedded between the poly (L-lactic acid)/polycaprolactone (PLLA/PCL) substrate and encapsulation, connected to six metal contacts in the connector pad¹ (**Figure 4a-i**). Penicillin given to a rat induced seizures to drastically increase the amplitude of the measured ECoG signals to 1430 μV within a narrow frequency range of 3-13 Hz, compared to the ECoG data of the rat in its nonrapid eye movement (NREM) sleep state (**Figure 4a-ii**). After stable operation with an impedance of 10.4 k Ω for 80 h, degradation of the ECoG electrode in PBS buffer (37°C, pH 7.4) signaled metal hydrolysis, drastically increasing the impedance¹ (**Figure 4a-iii**). The localized hydrolysis initiates from Mg contacts with a weak adhesion to the polymer. The hydrolysis follows the single-layer reactive diffusion model⁴⁷, increasing the impedance as a function of reduced thickness.

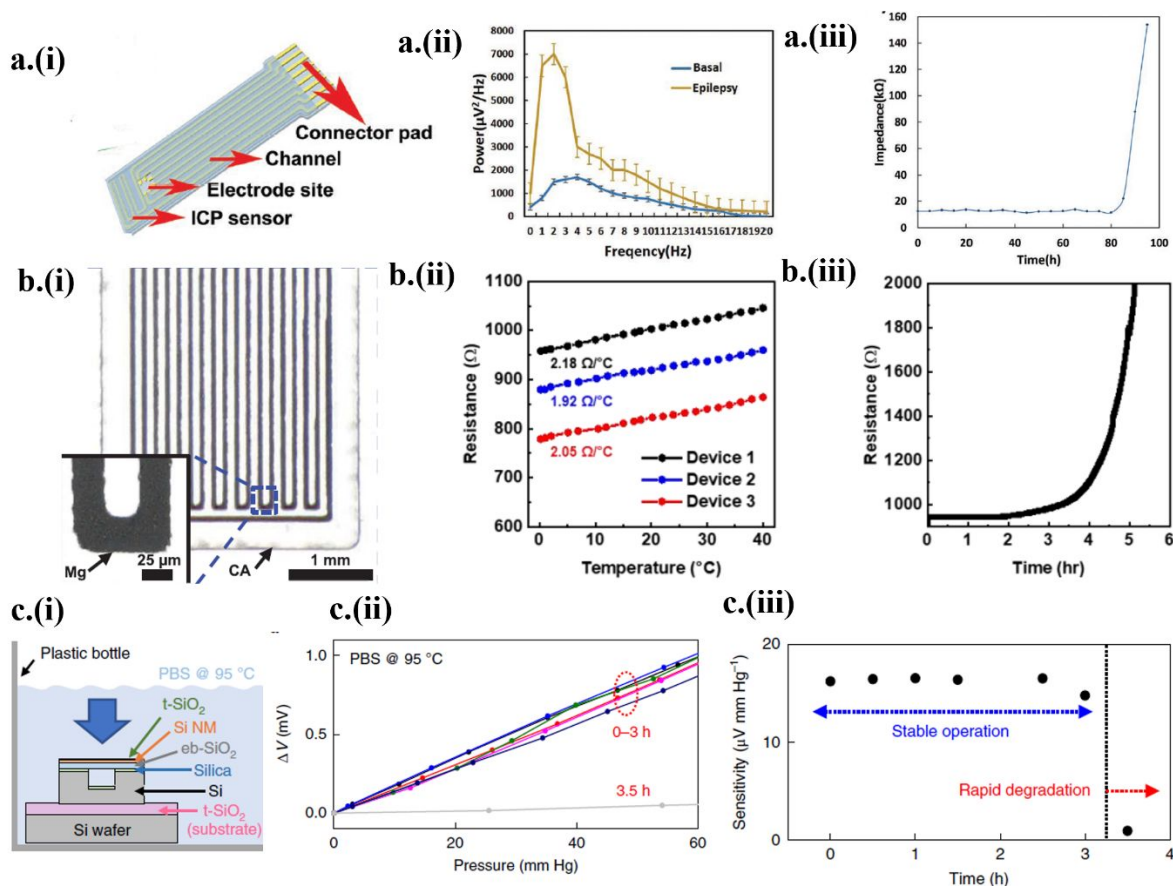


Figure 4. Transient electronics in diagnostic applications. (a) Measurements of electrophysiological signals by the ECoG device (i) designed with an intracortical pressure sensor. (ii) Power spectrum of ECoG data before and during the onset of epilepsy. (iii) Increase in impedance after the onset of degradation. This figure has been reproduced from ref.¹ with permission from John Wiley and Sons, copyright 2019. (b) (i) Design and (ii) calibration curve of Mg-based temperature sensor, with (iii) increased resistance after the initiation of the

degradation process. This figure has been reproduced from ref.²¹ with permission from The American Association for the Advancement of Science, copyright 2022. (c) (i) Design of the *in vitro* dissolution experiment with the eb-SiO₂/SiNM/t-SiO₂-based pressure sensor immersed in PBS solution. (ii) Linear voltage response with the pressure and (iii) decreased sensitivity after 3 hr due to the onset of degradation. This figure has been reproduced from ref.⁶⁶ with permission from Springer Nature, copyright 2018.

As a thermistor, an Mg-based temperature sensor with a width of 25 μm and length of 72 μm (**Figure 4b-i**) showed a sensitivity of $2.05 \pm 0.13 \text{ } \Omega/\text{ }^\circ\text{C}$ from 0 to 40°C (**Figure 4b-ii**)²¹. The encapsulation layers of POC and SiO₂ provide a stable operational time of 2 hr, followed by a degradation in PBS (pH 7.4, 37°C)²¹. The onset of hydrolysis of Mg conductors decreases its thickness following the single-layer reactive diffusion model, increasing the resistance (**Figure 4b-iii**). With comparable sensitivity to conventional, commercialized pressure sensors, the transient pressure sensors have been used to detect intracranial, intraocular, and blood pressure^{1,22,24,35,66,67}. Despite good sensitivity, transient pressure sensors often lack stability over time. Efforts to overcome this challenge led to the recent development⁶⁶ of a pressure sensor based on Si nanomembranes (NMs) sandwiched between a thermally grown Si dioxide (t-SiO₂) encapsulation layer and a thick electron-beam evaporated SiO₂ (eb-SiO₂). The thick eb-SiO₂ improved the response of the strain gauges to pressure due to the increased distance between the neutral mechanical plane and the Si NMs. With the accelerated dissolution testing setup in PBS (pH 7.4, 95°C) (**Figure 4c-i**), the complete dissociation of the encapsulating t-SiO₂ layer (~10 nm) was observed to take place within 5 hr, with relatively stable performance for 3 hr (**Figure 4c-ii, iii**). The dissolution of the t-SiO₂ encapsulation with a rate of a few hundred nanometers per day eventually leads to rapid degradation of Si NMs and bioresorbable metals due to hydrolysis. The time of stable operation could also be increased with the increased thickness of the t-SiO₂ layer.

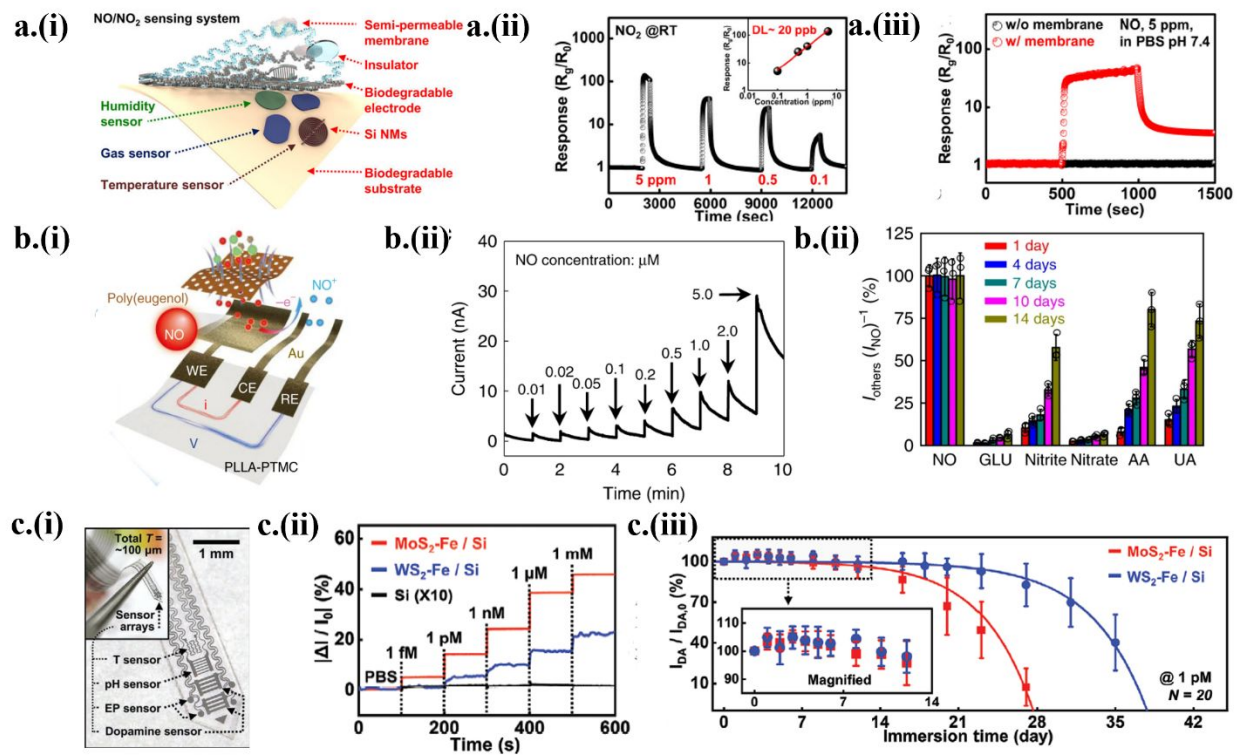


Figure 5. Transient electronics to monitor NO_x and dopamine. (a) (i) Design of the NO_x gas integrated with temperature and humidity sensors. (ii) Linear dynamic response to various NO₂ concentrations at room temperature with a linear fit to show the estimation of the limit of detection of 20 ppb. (iii) Dynamic response of the gas sensor with and without semipermeable to NO in the PBS solution. This figure has been reproduced under the terms of the Creative Commons Attribution 4.0 International License. ⁴ Copyright 2020, Springer Nature. (b) (i) Design of the electrochemical NO sensor with Au nanomembrane electrodes on PLLA-PTMC. (ii) Dynamic electrochemical response of the sensor to varying NO concentrations (bias voltage of 0.8 V). (iii) Stability of the NO sensor selectivity showing a decrease after 7 days. This figure has been reproduced under the terms of the Creative Commons Attribution 4.0 International License. ² Copyright 2020, Springer Nature. (c) (i) Integration of dopamine sensor with electrophysiology, temperature, and pH sensors in a neural probe. (ii) Relative current change of the neurochemical sensor to various dopamine concentrations. (iii) Change of response to 1 pM dopamine in a buffer solution for 5 weeks. This figure has been reproduced from ref.⁵ with permission from John Wiley and Sons, copyright 2022.

3.2 Nitric oxide and dopamine Sensors

As one of the most important gaseous biomarkers for environmental and health monitoring, nitric oxide (NO) is produced in the human body to maintain homeostatic equilibrium and provide nutrients to the tissues ^{4,68}. With n-doped monocrystalline silicon nanomembranes (~100 nm) and Mg (~300 nm) electrodes encapsulated by a semipermeable membrane (**Figure 5a-i**), a

transient NO_x sensor showed a sensitivity of 13600% to 5 ppm NO₂ at room temperature and fast response/recovery (30/60 s)⁴. The linear fit between the response and gas concentration gave a limit of detection (LOD) of ~20 ppb at room temperature (**Figure 5a-ii**). The semipermeable membrane encapsulation also allowed the gas sensor to show good stability and respond to NO_x in PBS solution (pH 7.4), whereas the transient gas sensor without any encapsulation dissolved in the solution with no signal (**Figure 5a-iii**). Along with the degradation of the PLGA substrate, all the constituent components such as Mg electrodes and Si NMs also get dissolved via hydrolysis, leaving nontoxic end products and byproducts.

Although the chemiresistive transient gas sensors may detect NO_x as it adsorbs on the surface to oxidize Si NMs, more accurate real-time NO_x monitoring from the solution relies on the use of chemical sensors. A transient electrochemical NO sensor was developed with Au nanomembrane electrodes and a selective membrane of poly(eugenol) film on a bioresorbable poly L-lactic acid/poly (trimethylene carbonate) (PLLA/PTMC) copolymer substrate² (**Figure 5b-i**). By using chronoamperometry, the electrochemical sensor demonstrated linear current response with a high sensitivity of 5.29 nA μM⁻¹ (in 0-5 μM NO) and 4.17 nA μM⁻¹ (in 5-100 μM NO), as well as a LOD of 3.97 nM (**Figure 5b-ii**). The electrochemical NO sensor showed a good selectivity with an almost linear current response to NO concentration for 7 days, but the sensitivity decreased up to 1.5 times on the 14th day along with the selectivity (**Figure 5b-iii**). The excellent stability over 7 days followed by the gradual decrease in the sensitivity is attributed to the stable Au nanomembrane and slow degradation rate of poly(eugenol) film and the copolymer substrate. The continuous and wireless detection of higher NO in the articular cavity of New Zealand White rabbits with anterior cruciate ligament (ACL) rupture injuries was observed to associate with the onset of osteoarthritis (OA) at the later stage⁶⁸.

The electrochemical sensors can also be exploited to detect dopamine⁵ for neural and cardiac health monitoring, predicting diseases ranging from Parkinson's and Alzheimer's and cardiovascular diseases to cancer. For instance, a bioresorbable, wireless neurochemical system was developed based on the heterostructure of 2D molybdenum/tungsten disulfide (MoS₂/WS₂) nanosheets and catalytic iron nanoparticles (Fe NPs) on Si NMs⁵ (**Figure 5c-i**). The resulting electrochemical sensor showed a linear response across the dopamine concentration from 10⁻¹⁵ M to 10⁻³ M. Compared with Fe-adsorbed Si, the 2D transition metal dichalcogenides (TMDs) in MoS₂-Fe/Si and WS₂-Fe/Si acted as a mediator between Si electrodes and catalyst Fe NPs⁵ (**Figure 5c-ii**). The excellent durability and robustness of the sensor allowed it to detect 1 pM

dopamine over 40 days. The transient electrochemical sensor could retain up to 90% of the initial value over 27 days for WS₂-Fe/Si and 18 days for MoS₂-Fe/Si⁵ (**Figure 5c-iii**). The relatively faster dissolution rate of MoS₂ can be attributed to the thermodynamically less stable MoS₂ in forming sulfate ions, favoring the forward chemical kinetics. Taken together with a wireless communication system connected to the brain-integrated probe, the simultaneously monitored dopamine and peripheral neurophysiological signals (e.g., pH, temperature, and electrophysiology) established the platform for *in vivo* studies of neurodegenerative diseases.

4. Implantable Transient Electronics in Therapeutic Applications

Along with sensing biomarkers, transient electronics can also deliver stimuli to modulate the biological activity or trigger drug delivery at specific target locations for therapeutics^{14,24,33,35,62}. Explored in neurotechnology, pancreatic and cardiac systems^{13,14,62}, these implantable stimulator or drug delivery systems allow the operator to control transience onset on demand.

4.1 Transient electronics-based drug-delivery system

The implantable drug delivery system is still in its infancy before realizing its full potential to deliver a drug at a prescribed rate and time to targeted locations for avoiding off-target side effects^{14,33}. An ideal drug delivery system should integrate wireless control with zero leakage for avoiding unwanted delivery sites. As a first step to achieving this deterministic control over the delivery process, a programmable, bioresorbable, wireless drug delivery system based on transient Polybutanedithiol 1,3,5-triallyl-1,3,5-triazine-2,4,6(1H,3H,5H)-trione pentaonic anhydride (PBTPA) reservoirs with Mg active valves was developed¹⁴ (**Figure 6a-i**). The wireless power harvester consisted of an Mg-based RF coil with a resonant frequency of ~5 MHz and a rectifier circuit using a Si NM diode and Mg/SiO₂/Mg capacitor¹⁴ (**Figure 6a-ii**). The generated overpotential between the anode gate and cathode from the wireless power harvester accelerated the Faradaic reaction at the anodic end, opening the anode gate to release the drug from the reservoir¹⁴ (**Figure 6a-iii**). The increased bias decreased the time delay of the triggering onset for the drug release (**Figure 6a-iv**). After the triggering onset, the diffusion-assisted drug release increased sharply and proceeded toward saturation, marking the equilibrium in drug mixing¹⁴ (**Figure 6a-v**). Sequential wireless insulin triggering into the blood at different resonant frequencies in response to increased glucose (simulated from injection into the blood) demonstrated robust control of the blood glucose level¹⁴ (**Figure 6a-vi**).

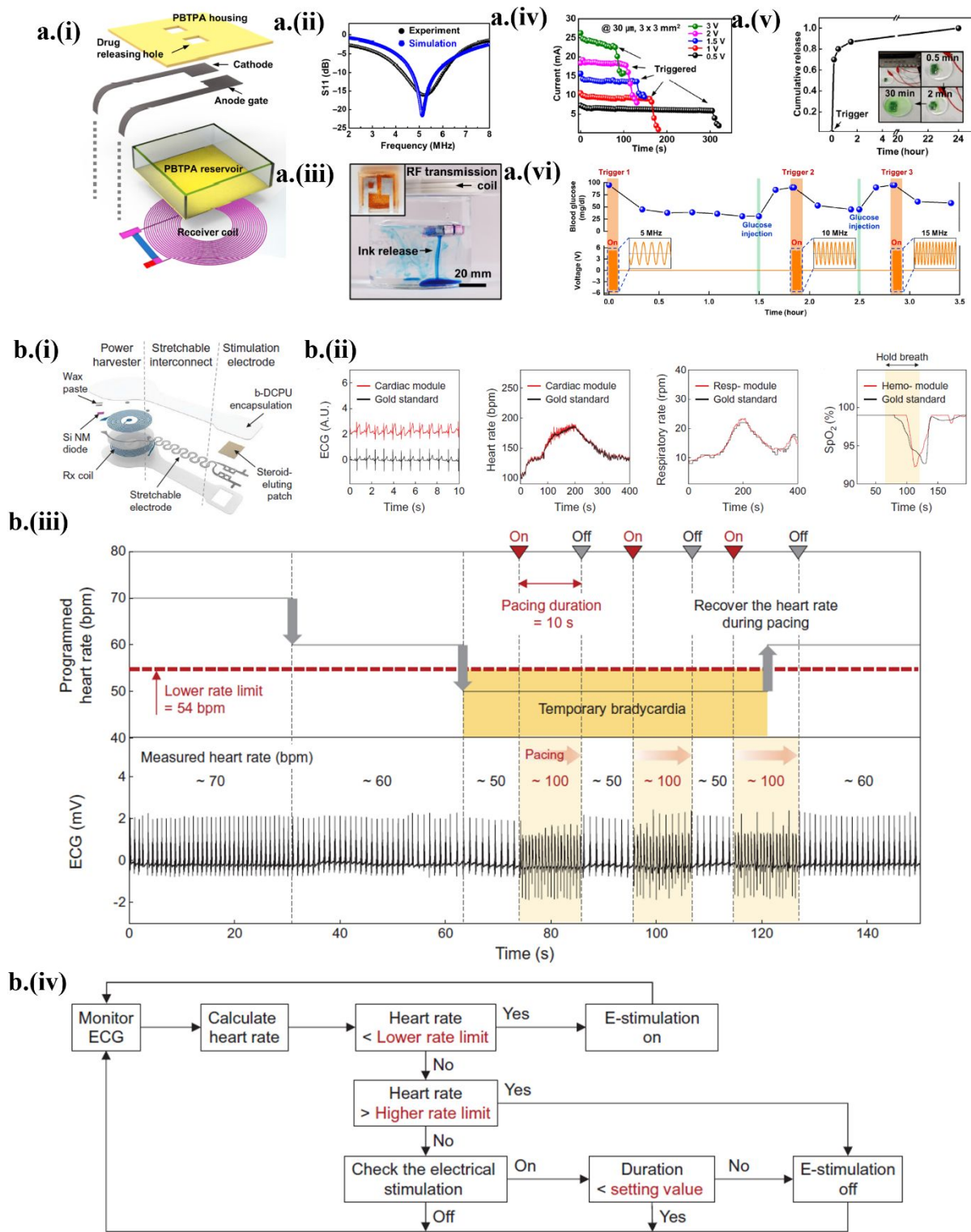


Figure 6. Transient actuating system in drug delivery and electrical stimulation. **(a)** (i) Design of the drug delivery system with an electrical triggering unit and bioresorbable reservoir. (ii) The return loss S_{11} of the Mg RF coil. (iii) Wireless controlled release of blue dye during degradation of the sensor in PBS solution. (iv) Voltage-dependent triggering event for a fixed Mg gate size. (v) Cumulative drug release after the triggered opening of the reservoir gate. (vi) Demonstration of insulin delivery to maintain the blood glucose level. This figure has been reproduced under the terms of the Creative Commons CC BY NC License.¹⁴ Copyright 2020, The American Association for the Advancement of Science. **(b)** (i) Bioresorbable stimulator with integrated sensing and actuating platform. (ii) Comparison of ECG, heart and respiratory rate, and SpO₂ bioresorbable sensor with the commercially available sensors. (iii) Triggering mechanism to maintain the average heart rate of 60 bpm based on a condition-based algorithm. (iv) Flowchart showing the condition-based algorithm to stimulate in the closed-loop system. This figure has been reproduced from ref.³³ with permission from The American Association for the Advancement of Science, copyright 2022.

4.2 Electrical stimulation with transient electrodes

The commercially available electrical stimulators such as the responsive neurostimulator (RNS) have been used in clinical setups to help nerve recovery and gain stability in neuronal disorders such as epilepsy and Alzheimer's disease^{13,33,69}. To avoid post-surgical complications from the commercially rigid and non-bioresorbable stimulators^{32,70}, a transient stimulator with sensors to detect biomarker signals in a closed-loop system allowed automatic modulation of the biological activity by stimulating based on the detected signals³². The Si NM-based PIN diode was used to rectify the voltage generated in the Mo-based receiver coil, which generated stimulation through Mo-based electrodes³³ (**Figure 6b-i**). The slow dissolution rate of the bioresorbable Mo conductor (tested in PBS at 95°C) enables a long-term (>1 month) operational lifetime. The holistic sensing system comprised skin-interfaced ECG electrodes, a microcontroller unit to calculate heart rate and respiratory rate, and a hemodynamic module to collect the SpO₂ data³³ (**Figure 6b-ii**). The wirelessly transferred sensing data to the cloud were used to process and optimize stimulation parameters according to a threshold-based algorithm. The closed-loop system automatically initiated pacing of 100 beats per minute (bpm) when the heart rate fell below 54 bpm and ceased on-demand stimulation to maintain an average heart rate of 60 bpm³³ (**Figure 6b-iii**) using a condition-based algorithm (**Figure 6b-iv**).

5. Conclusion and Future Directions

Since the initial demonstration of transient devices a little more than ten years ago, different bioresorbable materials and unconventional fabrication approaches have been explored to enable

controlled degradation or disintegration of the device at a prescribed rate or time. After the investigations of strategies for controlling the on-demand degradation of the device, stimulating devices have further been integrated with sensors to yield a transient closed-loop system that performs the stimulation according to condition-based algorithm³³. However, the if-else algorithms result in inefficient and impotent control logic. A pre-defined threshold-based decision-making controller almost always operates in a limited scope in practical settings, as it is unaware of past medical conditions and present environmental scenarios. The closed-loop system based on a more practical and Artificial Intelligence (AI)-powered controller is thereby imperative³². Lessons can be learned from other biomedical closed-loop systems. For instance, an implantable pulse generator could carry out epidural electrical stimulation (EES) of the spinal cord with a spatiotemporally adaptive trigger controller⁷¹. The sensing system consisted of the electromyography (EMG) response of the leg muscle and the whole-body ground reaction. With an option of pre-programmed open-loop or closed-loop, the wireless communication module carried out the EES with dynamic control over EES parameters⁷¹ (**Figure 7a-i**). Only with the introduction of closed-loop spatiotemporal EES, the participants could voluntarily walk, and reintroducing EES after a stoppage resumed the participant's locomotion⁷¹ (**Figure 7a-ii**). Similar to spinal cord stimulation, closed-loop deep brain stimulation (DBS) or neuromodulation was recently reported as well. With measured local field potential (LFP) to automatically detect pain in a closed-loop system, a brain-machine interface was set up to optogenetically or electrically perform DBS⁶⁹ (**Figure 7b-i**). The multiregional neural interface from the anterior cingulate cortex (ACC) and somatosensory cortex (S1) could detect and treat acute evoked and chronic pain. By computing frequency-dependent LFP power features, a real-time neural decoder based on a Markovian-driven state space model (unsupervised learning) was designed to decode the pain onset⁶⁹ (**Figure 7b-ii**), which applied the DBS to reduce the acute thermal pain⁶⁹ (**Figure 7b-iii**). Along with intelligent closed loop-based stimulation, a protocol of sequential searching for biomarker signals and optimizing stimulation parameters has also been effective in complicated, biological systems. As an example, biomarker-driven closed-loop DBS was carried out to treat the major depressive disorder (MDD)³². Unlike Parkinson's disease or epilepsy, MDD requires personalized neural circuit targeting, which results in the patient's own physiological biomarkers-based trigger or stimulation. The stimulus-response mapping was initially performed for the emotion neural circuit using stereoelectroencephalography (SEEG) electrodes at important regions such as the amygdala, hippocampus, and ventral capsule/ventral striatum (VC/VS). The SEEG

spectral activity identified amygdala gamma power as a sufficient biomarker. With the right VC/VS as the selected stimulation site, the correlation between the stimulation parameters and the amygdala gamma power was analyzed³² (**Figure 7c-i**). According to the correlation, the FDA-approved NeuroPace RNS system was used with two separate sensing and stimulation leads simultaneously placed in the amygdala and VC/VS region to stimulate with the optimized parameters³² (**Figure 7c-ii**). The implementation of closed-loop neuromodulation resulted in significant improvement in the symptom severity according to the visual analog scale (VAS) scales³² (**Figure 7c-iii**).

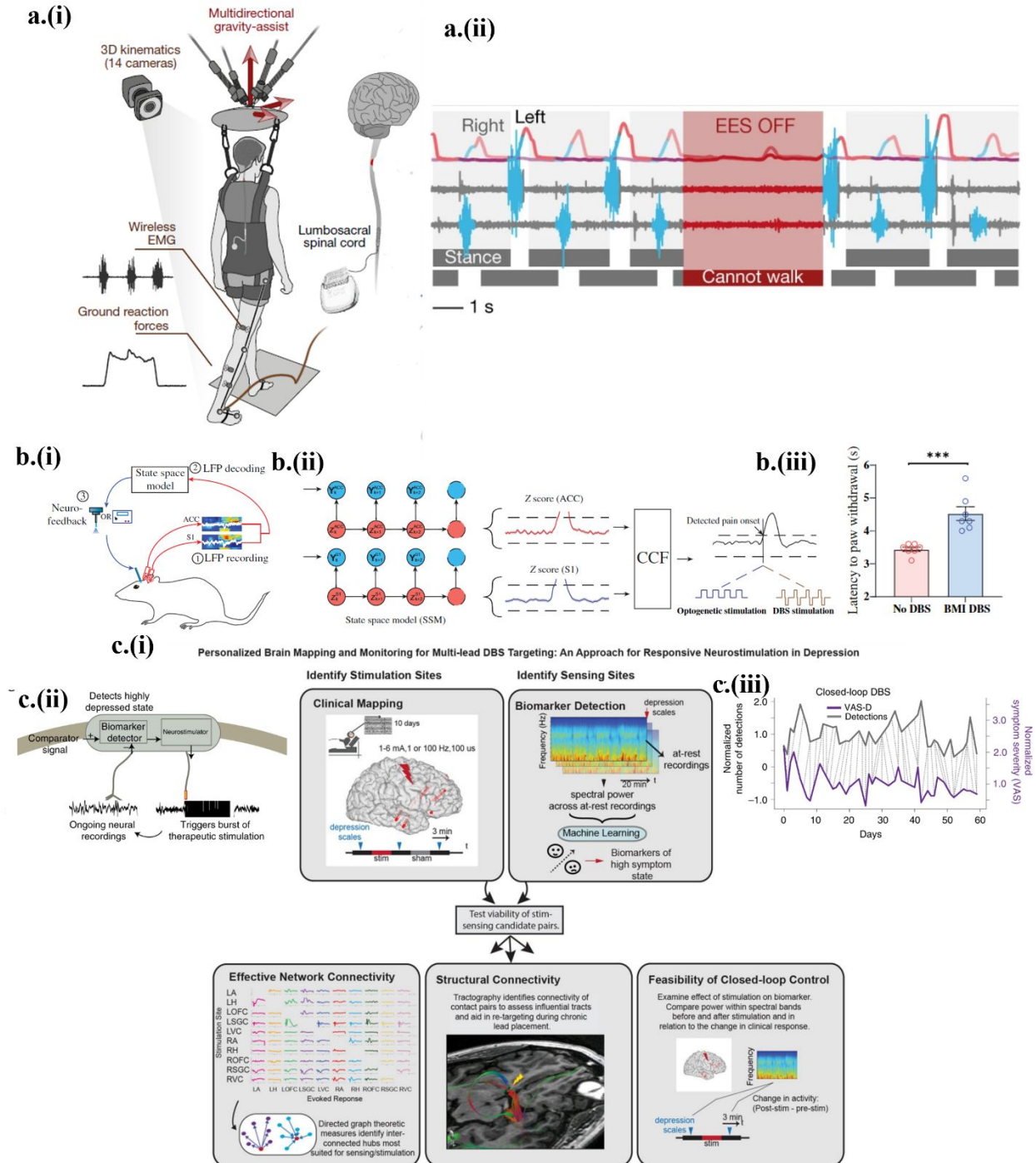


Figure 7. On-demand closed-loop transient systems. (a) Epidural electrical stimulation (EES) of the spinal cord with (i) the control module to (ii) demonstrate spatiotemporal EES for voluntary walking. This figure has been reproduced from ref. ⁷¹ with permission from Springer Nature, copyright 2018. (b) Closed-loop brain-machine interface with (i) the control module based on the detection of low field potential for deep brain stimulation

(DBS). (ii) Representation of the Markovian-driven state space model to detect pain onset. (iii) Demonstration of improved latency with the brain-machine interface-based DBS. This figure has been reproduced from ref.⁶⁹ with permission from The American Association for the Advancement of Science, copyright 2022. (c) Closed-Loop neuromodulation relies on (i) the sequential algorithm optimizing the location of the biomarker, validating the network and structural connectivity and the stimulation parameters. (ii) The NeuroPace RNS system with biomarker detection and stimulation. (iii) Aligning biomarker detection with VAS rating using dynamic time wrapping for demonstrated improvement due to biomarker-driver stimulation. This figure has been reproduced from ref.³² with permission from Springer Nature, copyright 2021.

In addition to the closed-loop system or system-level integration of transient electronics, there are still a few fundamental challenges that need to be addressed. For example, multiple sensing units (e.g., temperature, pH, pressure) in the transient device may affect each other, so the decoupling of the sensing data would need to be performed⁷². Furthermore, with the advent of on-demand transience, the selectivity of multimodal transient electronic devices in the presence of more than one external (active and passive) stimuli is often ignored. For example, transient devices that disintegrate faster in electrical (active) and comparatively slower in optical (passive) stimuli are expected to dissociate at an unexpected rate in the presence of both stimuli. Moreover, it becomes difficult to control the influence of passive stimuli on the disintegration of the device, presenting challenges to its use and robustness in real-world applications. An intuitive strategy to solve this issue is to incorporate multilayer encapsulations with materials having complementary dissociation-selectivity towards stimuli. For example, materials that dissociate in electrical and water stimuli and materials with electrical and optical stimuli-specific dissolution may form a multi-layer encapsulation to implement transience that can only be triggered by the electrical stimulus, increasing the selectivity towards electrical triggered transience. Employing specific sequence-based external stimuli can result in a complete dissociation of the transient device with composite encapsulations⁶⁴. The sequence of stimuli-selective materials in the composite forms an intelligent hardware-based password to trigger dissolution rather than a destined degradation, adding an extra dimension to the security. The ultimate practical, robust transient electronics system hinges on the mature developments in closed-loop therapeutics, decoupling mechanism, and dissolution selectivity discussed above. Although considerable advancements have been made, the field of intelligent transient electronics systems is still in its infancy. However, the introduction of intelligent, robust transient electronics systems will open up opportunities from hardware security to biomedical diagnosis and therapeutics in the real world.

Acknowledgments

HC acknowledges the support provided by NIH (Award Nos. R21EB030140, U01DA056242, and R61HL154215), NSF (Grant No. ECCS-1933072), and Penn State University.

Declaration of Competing Interest

The authors declare that there are no competing financial interests that appear to influence this work.

References:

- 1 K. Xu, S. Li, S. Dong, S. Zhang, G. Pan, G. Wang, L. Shi, W. Guo, C. Yu and J. Luo, *Adv Healthc Mater*, 2019, **8**, 1801649.
- 2 R. Li, H. Qi, Y. Ma, Y. Deng, S. Liu, Y. Jie, J. Jing, J. He, X. Zhang, L. Wheatley, C. Huang, X. Sheng, M. Zhang and L. Yin, *Nat Commun*, 2020, **11**, 3207.
- 3 S. Zhong, X. Ji, L. Song, Y. Zhang and R. Zhao, *ACS Appl Mater Interfaces*, 2018, **10**, 36171–36176.
- 4 G. J. Ko, S. D. Han, J. K. Kim, J. Zhu, W. B. Han, J. Chung, S. M. Yang, H. Cheng, D. H. Kim, C. Y. Kang and S. W. Hwang, *NPG Asia Mater*, 2020, **12**, 71.
- 5 S. M. Yang, J. H. Shim, H. U. Cho, T. M. Jang, G. J. Ko, J. Shim, T. H. Kim, J. Zhu, S. Park, Y. S. Kim, S. Y. Joung, J. C. Choe, J. W. Shin, J. H. Lee, Y. M. Kang, H. Cheng, Y. Jung, C. H. Lee, D. P. Jang and S. W. Hwang, *Advanced Materials*, 2022, **34**, 2108203.
- 6 S. K. Kang, R. K. J. Murphy, S. W. Hwang, S. M. Lee, D. v. Harburg, N. A. Krueger, J. Shin, P. Gamble, H. Cheng, S. Yu, Z. Liu, J. G. McCall, M. Stephen, H. Ying, J. Kim, G. Park, R. C. Webb, C. H. Lee, S. Chung, D. S. Wie, A. D. Gujar, B. Vemulapalli, A. H. Kim, K. M. Lee, J. Cheng, Y. Huang, S. H. Lee, P. v. Braun, W. Z. Ray and J. A. Rogers, *Nature*, 2016, **530**, 71–76.
- 7 C. M. Boutry, Y. Kaizawa, B. C. Schroeder, A. Chortos, A. Legrand, Z. Wang, J. Chang, P. Fox and Z. Bao, *Nat. Electron.*, 2018, **1**, 314–321.
- 8 H. Lopez Hernandez, S. K. Kang, O. P. Lee, S. W. Hwang, J. A. Kaitz, B. Inci, C. W. Park, S. Chung, N. R. Sottos, J. S. Moore, J. A. Rogers and S. R. White, *Advanced Materials*, 2014, **26**, 7637–7642.
- 9 T. Lei, M. Guan, J. Liu, H. C. Lin, R. Pfattner, L. Shaw, A. F. McGuire, T. C. Huang, L. Shao, K. T. Cheng, J. B. H. Tok and Z. Bao, *Proc Natl Acad Sci U S A*, 2017, **114**, 5107–5112.
- 10 P. S. Liu and H. F. Tse, *J. Arrhythmia*, 2013, **29**, 314–319.
- 11 R. J. Vetter, J. C. Williams, J. F. Hetke, E. A. Nunamaker and D. R. Kipke, *IEEE Trans. Biomed. Eng.*, 2004, **51**, 896–904.
- 12 H. U. Chung, A. Y. Rwei, A. Hourlier-Fargette, S. Xu, K. H. Lee, E. C. Dunne, Z. Xie, C. Liu, A. Carlini, D. H. Kim, D. Ryu, E. Kulikova, J. Cao, I. C. Odland, K. B. Fields, B. Hopkins, A. Banks, C. Ogle, D. Grande, J. bin Park, J. Kim, M. Irie, H. Jang, J. H. Lee, Y. Park, J. Kim, H. H. Jo, H. Hahm, R. Avila, Y. Xu, M. Namkoong, J. W. Kwak, E. Suen, M. A. Paulus, R. J. Kim, B. v. Parsons, K. A. Human, S. S. Kim, M. Patel, W. Reuther, H. S. Kim, S. H. Lee, J. D. Leedle, Y. Yun, S. Rigali, T. Son, I. Jung, H. Arafa, V. R. Soundararajan, A. Ollech, A. Shukla, A. Bradley, M. Schau, C. M. Rand, L. E. Marsillio, Z. L. Harris, Y. Huang, A. Hamvas, A. S. Paller, D. E. Weese-Mayer, J. Y. Lee and J. A. Rogers, *Nat Med*, 2020, **26**, 418–429.
- 13 G. Lee, E. Ray, H. J. Yoon, S. Genovese, Y. S. Choi, M. K. Lee, S. Şahin, Y. Yan, H. Y. Ahn, A. J. Bandodkar, J. Kim, M. Park, H. Ryu, S. S. Kwak, Y. H. Jung, A. Odabas, U. Khandpur, W. Z. Ray, M. R. MacEwan and J. A. Rogers, *Sci Adv*, 2022, **8**, eabp9169.
- 14 J. Koo, S. B. Kim, Y. S. Choi, Z. Xie, A. J. Bandodkar, J. Khalifeh, Y. Yan, H. Kim, M. K. Pezhouh, K. Doty, G. Lee, Y. Y. Chen, S. M. Lee, D. D’Andrea, K. Jung, K. H. Lee, K. Li, S. Jo, H. Wang, J. H. Kim, J. Kim, S. G. Choi, W. J. Jang, Y. S. Oh, I. Park, S. S. Kwak, J. H. Park, D. Hong, X. Feng, C. H. Lee, A. Banks, C. Leal, H. M. Lee, Y. Huang, C. K. Franz, W. Z. Ray, M. MacEwan, S. K. Kang and J. A. Rogers, *Sci Adv*, 2020, **6**, 20.
- 15 M. Prakasam, J. Locs, K. Salma-Ancane, D. Loca, A. Largeteau and L. Berzina-Cimdina, *J Funct*

- Biomater*, 2017, **8**, 44.
- 16 J. Koo, M. R. MacEwan, S. K. Kang, S. M. Won, M. Stephen, P. Gamble, Z. Xie, Y. Yan, Y. Y. Chen, J. Shin, N. Birenbaum, S. Chung, S. B. Kim, J. Khalifeh, D. v. Harburg, K. Bean, M. Paskett, J. Kim, Z. S. Zohny, S. M. Lee, R. Zhang, K. Luo, B. Ji, A. Banks, H. M. Lee, Y. Huang, W. Z. Ray and J. A. Rogers, *Nat Med*, 2018, **24**, 1830–1836.
- 17 C. Li, C. Guo, V. Fitzpatrick, A. Ibrahim, M. J. Zwierstra, P. Hanna, A. Lechtig, A. Nazarian, S. J. Lin and D. L. Kaplan, *Nat. Rev. Mater.*, 2020, **5**, 61–81.
- 18 A. D. Mickle, S. M. Won, K. N. Noh, J. Yoon, K. W. Meacham, Y. Xue, L. A. McIlvried, B. A. Copits, V. K. Samineni, K. E. Crawford, D. H. Kim, P. Srivastava, B. H. Kim, S. Min, Y. Shiuan, Y. Yun, M. A. Payne, J. Zhang, H. Jang, Y. Li, H. H. Lai, Y. Huang, S. il Park, R. W. Gereau and J. A. Rogers, *Nature*, 2019, **565**, 361–365.
- 19 Y. S. Choi, R. T. Yin, A. Pfenniger, J. Koo, R. Avila, K. Benjamin Lee, S. W. Chen, G. Lee, G. Li, Y. Qiao, A. Murillo-Berlioz, A. Kiss, S. Han, S. M. Lee, C. Li, Z. Xie, Y. Y. Chen, A. Burrell, B. Geist, H. Jeong, J. Kim, H. J. Yoon, A. Banks, S. K. Kang, Z. J. Zhang, C. R. Haney, A. V. Sahakian, D. Johnson, T. Efimova, Y. Huang, G. D. Trachiotis, B. P. Knight, R. K. Arora, I. R. Efimov and J. A. Rogers, *Nat Biotechnol*, 2021, **39**, 1228–1238.
- 20 Y. S. Choi, Y. Y. Hsueh, J. Koo, Q. Yang, R. Avila, B. Hu, Z. Xie, G. Lee, Z. Ning, C. Liu, Y. Xu, Y. J. Lee, W. Zhao, J. Fang, Y. Deng, S. M. Lee, A. Vázquez-Guardado, I. Stepien, Y. Yan, J. W. Song, C. Haney, Y. S. Oh, W. Liu, H. J. Yun, A. Banks, M. R. MacEwan, G. A. Ameer, W. Z. Ray, Y. Huang, T. Xie, C. K. Franz, S. Li and J. A. Rogers, *Nat Commun*, 2020, **11**, 5990.
- 21 J. T. Reeder, Z. Xie, Q. Yang, M. H. Seo, Y. Yan, Y. Deng, K. R. Jinkins, S. R. Krishnan, C. Liu, S. McKay, E. Patnaude, A. Johnson, Z. Zhao, M. J. Kim, Y. Xu, I. Huang, R. Avila, C. Felicelli, E. Ray, X. Guo, W. Z. Ray, Y. Huang, M. R. MacEwan and J. A. Rogers, *Science (1979)*, 2022, **377**, 109–115.
- 22 C. You, H. Zhao, Q. Guo and Y. Mei, *MRS Bull*, 2020, **45**, 129–134.
- 23 D. M. Lee, N. Rubab, I. Hyun, W. Kang, Y. J. Kim, M. Kang, B. O. Choi and S. W. Kim, *Sci Adv*, 2022, **8**, 8423.
- 24 G. Lee, Y. S. Choi, H. J. Yoon and J. A. Rogers, *Matter*, 2020, **3**, 1031–1052.
- 25 J. W. Shin, J. Chan Choe, J. H. Lee, W. B. Han, T. M. Jang, G. J. Ko, S. M. Yang, Y. G. Kim, J. Joo, B. H. Lim, E. Park and S. W. Hwang, *ACS Nano*, 2021, **15**, 19310–19320.
- 26 J. Ausra, M. Madrid, R. T. Yin, J. Hanna, S. Arnott, J. A. Brennan, R. Peralta, D. Clausen, J. A. Bakall, I. R. Efimov and P. Gutruf, *Sci Adv*, 2022, **8**, eabq7469.
- 27 Y. Chen, H. Wang, Y. Zhang, R. Li, C. Chen, H. Zhang, S. Tang, S. Liu, X. Chen, H. Wu, R. Lv, X. Sheng, P. Zhang, S. Wang and L. Yin, *Nanotechnology*, 2019, **30**, 394002.
- 28 C. M. Boutry, H. Chandralim, P. Streit, M. Schinhammer, A. C. Hänzi and C. Hierold, *Philos. Trans. R. Soc. A*, 2012, **370**, 2418–2432.
- 29 R. Proietti, G. Manzoni, L. di Biase, G. Castelnuovo, L. Lombardi, C. Fundarò, N. Vegliante, G. Pietrabissa, P. Santangeli, R. A. Canby, A. Sagone, M. Viecca and A. Natale, *PACE - Pacing and Clinical Electrophysiology*, 2012, **35**, 990–998.
- 30 C. G. Kanakry, *Sci Transl Med*, 2020, **12**, eabb2779.
- 31 R. Hovorka, *Diabetic Medicine*, 2006, **23**, 1–12.
- 32 K. W. Scangos, A. N. Khambhati, P. M. Daly, G. S. Makhoul, L. P. Sugrue, H. Zamanian, T. X. Liu, V.

- R. Rao, K. K. Sellers, H. E. Dawes, P. A. Starr, A. D. Krystal and E. F. Chang, *Nat Med*, 2021, **27**, 1696–1700.
- 33 Y. S. Choi, H. Jeong, R. T. Yin, R. Avila, A. Pfenniger, J. Yoo, J. Y. Lee, A. Tzavelis, Y. J. Lee, S. W. Chen, H. S. Knight, S. Kim, H. Y. Ahn, G. Wickerson, A. Vázquez-Guardado, E. Higbee-Dempsey, B. A. Russo, M. A. Napolitano, T. J. Holleran, L. A. Razzak, A. N. Miniovich, G. Lee, B. Geist, B. Kim, S. Han, J. A. Brennan, K. Aras, S. S. Kwak, J. Kim, E. A. Waters, X. Yang, A. Burrell, K. S. Chun, C. Liu, C. Wu, A. Y. Rwei, A. N. Spann, A. Banks, D. Johnson, Z. J. Zhang, C. R. Haney, S. H. Jin, A. V. Sahakian, Y. Huang, G. D. Trachiotis, B. P. Knight, R. K. Arora, I. R. Efimov and J. A. Rogers, *Science (1979)*, 2022, **376**, 1006–1012.
- 34 R. Li, L. Wang, D. Kong and L. Yin, *Bioact Mater*, 2017, **3**, 322–333.
- 35 J. S. Shim, J. A. Rogers and S. K. Kang, *Materials Science and Engineering: R: Reports*, 2021, **145**, 100624.
- 36 L. Yin, X. Huang, H. Xu, Y. Zhang, J. Lam, J. Cheng and J. A. Rogers, *Advanced Materials*, 2014, **26**, 3879–3884.
- 37 Y. H. Jung, H. Zhang, S. Gong and Z. Ma, *Semicond. Sci. Technol.*, 2017, **32**, 063002.
- 38 Q. Yang, S. Lee, Y. Xue, Y. Yan, T. L. Liu, S. K. Kang, Y. J. Lee, S. H. Lee, M. H. Seo, D. Lu, J. Koo, M. R. MacEwan, R. T. Yin, W. Z. Ray, Y. Huang and J. A. Rogers, *Adv. Funct. Mater.*, 2020, **30**, 1910718.
- 39 S. W. Hwang, X. Huang, J. H. Seo, J. K. Song, S. Kim, S. Hage-Ali, H. J. Chung, H. Tao, F. G. Omenetto, Z. Ma and J. A. Rogers, *Advanced Materials*, 2013, **25**, 3526–3531.
- 40 X. Huang, Y. Liu, S. W. Hwang, S. K. Kang, D. Patnaik, J. F. Cortes and J. A. Rogers, *Advanced Materials*, 2014, **26**, 7371–7377.
- 41 Y. Choi, J. Koo and J. A. Rogers, *MRS Bull*, 2020, **45**, 103–112.
- 42 S. W. Hwang, D. H. Kim, H. Tao, T. il Kim, S. Kim, K. J. Yu, B. Panilaitis, J. W. Jeong, J. K. Song, F. G. Omenetto and J. A. Rogers, *Adv Funct Mater*, 2013, **23**, 4087–4093.
- 43 J.-K. Chang, *Proc. Natl Acad. Sci. USA*, 2017, **114**, E5522–E5529.
- 44 Q. Yang, T. Wei, R. T. Yin, M. Wu, Y. Xu, J. Koo, Y. S. Choi, Z. Xie, S. W. Chen, I. Kandela, S. Yao, Y. Deng, R. Avila, T. L. Liu, W. Bai, Y. Yang, M. Han, Q. Zhang, C. R. Haney, K. Benjamin Lee, K. Aras, T. Wang, M. H. Seo, H. Luan, S. M. Lee, A. Brikha, N. Ghoreishi-Haack, L. Tran, I. Stepien, F. Aird, E. A. Waters, X. Yu, A. Banks, G. D. Trachiotis, J. M. Torkelson, Y. Huang, Y. Kozorovitskiy, I. R. Efimov and J. A. Rogers, *Nature Materials 2021 20:11*, 2021, **20**, 1559–1570.
- 45 Y. S. Choi, J. Koo, Y. J. Lee, G. Lee, R. Avila, H. Ying, J. Reeder, L. Hambitzer, K. Im, J. Kim, K. M. Lee, J. Cheng, Y. Huang, S. K. Kang and J. A. Rogers, *Adv Funct Mater*, 2020, **30**, 2000941.
- 46 S. W. Hwang, J. K. Song, X. Huang, H. Cheng, S. K. Kang, B. H. Kim, J. H. Kim, S. Yu, Y. Huang and J. A. Rogers, *Advanced Materials*, 2014, **26**, 3905–3911.
- 47 L. Yin, H. Cheng, S. Mao, R. Haasch, Y. Liu, X. Xie, S. W. Hwang, H. Jain, S. K. Kang, Y. Su, R. Li, Y. Huang and J. A. Rogers, *Adv Funct Mater*, 2014, **24**, 645–658.
- 48 H. Ryu, M. H. Seo and J. A. Rogers, *Adv Healthc Mater*, 2021, **10**, 2002236.
- 49 S. K. Kang, G. Park, K. Kim, S. W. Hwang, H. Cheng, J. Shin, S. Chung, M. Kim, L. Yin, J. C. Lee, K. M. Lee and J. A. Rogers, *ACS Appl Mater Interfaces*, 2015, **7**, 9297–9305.
- 50 L. Teng, S. Ye, S. Handschuh-Wang, X. Zhou, T. Gan, X. Zhou, L. Teng, S. C. Ye, S. Handschuh-Wang,

- X. H. Zhou, T. S. Gan and X. C. Zhou, *Adv Funct Mater*, 2019, **29**, 1808739.
- 51 Q. Guo, J. Koo, Z. Xie, R. Avila, X. Yu, X. Ning, H. Zhang, X. Liang, S. B. Kim, Y. Yan, M. R. MacEwan, H. M. Lee, A. Song, Z. Di, Y. Huang, Y. Mei and J. A. Rogers, *Adv Funct Mater*, 2019, **29**, 1905451.
- 52 N. Yi, Y. Gao, A. lo Verso, J. Zhu, D. Erdely, C. Xue, R. Lavelle and H. Cheng, *Materials Today*, 2021, **50**, 24–34.
- 53 R. Li, H. Cheng, Y. Su, S. W. Hwang, L. Yin, H. Tao, M. A. Brenckle, D. H. Kim, F. G. Omenetto, J. A. Rogers and Y. Huang, *Adv Funct Mater*, 2013, **23**, 3106–3114.
- 54 S. K. Kang, S. W. Hwang, H. Cheng, S. Yu, B. H. Kim, J. H. Kim, Y. Huang and J. A. Rogers, *Adv Funct Mater*, 2014, **24**, 4427–4434.
- 55 X. Zhang, C. M. Weber and L. M. Bellan, *Adv Mater Technol*, 2019, **4**, 1900528.
- 56 W. D. Chen, S. K. Kang, W. J. Stark, J. A. Rogers and R. N. Grass, *Sens Actuators B Chem*, 2019, **282**, 52–59.
- 57 Y. Chen, H. Wang, Y. Zhang, R. Li, C. Chen, H. Zhang, S. Tang, S. Liu, X. Chen, H. Wu, R. Lv, X. Sheng, P. Zhang, S. Wang and L. Yin, *Nanotechnology*, 2019, **30**, 394002.
- 58 L. A. Wehner, N. Mittal, T. Liu and M. Niederberger, *ACS Cent Sci*, 2021, **7**, 231–244.
- 59 C. Dagdeviren, S. W. Hwang, Y. Su, S. Kim, H. Cheng, O. Gur, R. Haney, F. G. Omenetto, Y. Huang and J. A. Rogers, *Small*, 2013, **9**, 3398–3404.
- 60 R. Hinchet, H. J. Yoon, H. Ryu, M. K. Kim, E. K. Choi, D. S. Kim and S. W. Kim, *Science (1979)*, 2019, **365**, 491–494.
- 61 R. Jamshidi, M. Taghavimehr, Y. Chen, N. Hashemi and R. Montazami, *Adv Sustain Syst*, 2022, **6**, 2100057.
- 62 A. Fanelli and D. Ghezzi, *Curr Opin Biotechnol*, 2021, **72**, 22–28.
- 63 W. B. Han, H. Lee, J.-W. Shin, S.-W. Hwang, W. B. Han, J. H. Lee, J.-W. Shin and S.-W. Hwang, *Advanced Materials*, 2020, **32**, 2002211.
- 64 S. Wei, J. Jiang, L. Sun, J. Li, T. H. Tao, Z. Zhou, S. Wei, J. Jiang, L. Sun, T. H. Tao, Z. Zhou and J. Li, *Advanced Materials*, 2022, **34**, 2201035.
- 65 Q. Yang, T. L. Liu, Y. Xue, H. Wang, Y. Xu, B. Emon, M. Wu, C. Rountree, T. Wei, I. Kandela, C. R. Haney, A. Brikha, I. Stepien, J. Hornick, R. A. Sponenburg, C. Cheng, L. Ladehoff, Y. Chen, Z. Hu, C. Wu, M. Han, J. M. Torkelson, Y. Kozorovitskiy, M. T. A. Saif, Y. Huang, J. K. Chang and J. A. Rogers, *Nature Electronics 2022 5:8*, 2022, **5**, 526–538.
- 66 J. Shin, Y. Yan, W. Bai, Y. Xue, P. Gamble, L. Tian, I. Kandela, C. R. Haney, W. Spees, Y. Lee, M. Choi, J. Ko, H. Ryu, J. K. Chang, M. Pezhouh, S. K. Kang, S. M. Won, K. J. Yu, J. Zhao, Y. K. Lee, M. R. MacEwan, S. K. Song, Y. Huang, W. Z. Ray and J. A. Rogers, *Nat Biomed Eng*, 2019, **3**, 37–46.
- 67 D. Lu, Y. Yan, Y. Deng, Q. Yang, J. Zhao, M. H. Seo, W. Bai, M. R. MacEwan, Y. Huang, W. Z. Ray and J. A. Rogers, *Adv Funct Mater*, 2020, **30**, 2003754.
- 68 Y. Deng, H. Qi, Y. Ma, S. Liu, M. Zhao, Z. Guo, Y. Jie, R. Zheng, J. Jing, K. Chen, H. Ding, G. Lv, K. Zhang, R. Li, H. Cheng, L. Zhao, X. Sheng, M. Zhang and L. Yin, *Proceedings of the National Academy of Sciences*, 2022, **119**, e2208060119.
- 69 G. Sun, F. Zeng, M. McCartin, Q. Zhang, H. Xu, Y. Liu, Z. S. Chen and J. Wang, *Sci Transl Med*, 2022, **14**, eabm5868.

- 70 G. K. Bergey, M. J. Morrell, E. M. Mizrahi, A. Goldman, D. King-Stephens, D. Nair, S. Srinivasan, B. Jobst, R. E. Gross, D. C. Shields, G. Barkley, V. Salanova, P. Olejniczak, A. Cole, S. S. Cash, K. Noe, R. Wharen, G. Worrell, A. M. Murro, J. Edwards, M. Duchowny, D. Spencer, M. Smith, E. Geller, R. Gwinn, C. Skidmore, S. Eisenschenk, M. Berg, C. Heck, P. van Ness, N. Fountain, P. Rutecki, A. Massey, C. O'Donovan, D. Labar, R. B. Duckrow, L. J. Hirsch, T. Courtney, F. T. Sun and C. G. Seale, *Neurology*, 2015, **84**, 810–817.
- 71 F. B. Wagner, J. B. Mignardot, C. G. le Goff-Mignardot, R. Demesmaeker, S. Komi, M. Capogrosso, A. Rowald, I. Seáñez, M. Caban, E. Pirondini, M. Vat, L. A. McCracken, R. Heimgartner, I. Fodor, A. Watrin, P. Seguin, E. Paoles, K. van den Keybus, G. Eberle, B. Schurch, E. Pralong, F. Becce, J. Prior, N. Buse, R. Buschman, E. Neufeld, N. Kuster, S. Carda, J. von Zitzewitz, V. Delattre, T. Denison, H. Lambert, K. Minassian, J. Bloch and G. Courtine, *Nature* 2018 563:7729, 2018, **563**, 65–71.
- 72 R. Yang, W. Zhang, N. Tiwari, H. Yan, T. Li and H. Cheng, *Advanced Science*, 2022, **9**, 2202470.