

## SENSORS

# Biofuel-powered soft electronic skin with multiplexed and wireless sensing for human-machine interfaces

You Yu<sup>1</sup>, Joanna Nassar<sup>1</sup>, Changhao Xu<sup>1</sup>, Jihong Min<sup>1</sup>, Yiran Yang<sup>1</sup>, Adam Dai<sup>2</sup>, Rohan Doshi<sup>2</sup>, Adrian Huang<sup>3</sup>, Yu Song<sup>1</sup>, Rachel Gehlhar<sup>4</sup>, Aaron D. Ames<sup>4</sup>, Wei Gao<sup>1\*</sup>

Existing electronic skin (e-skin) sensing platforms are equipped to monitor physical parameters using power from batteries or near-field communication. For e-skins to be applied in the next generation of robotics and medical devices, they must operate wirelessly and be self-powered. However, despite recent efforts to harvest energy from the human body, self-powered e-skin with the ability to perform biosensing with Bluetooth communication are limited because of lack of a continuous energy source and limited power efficiency. Here, we report a flexible and fully perspiration-powered integrated electronic skin (PPES) for multiplexed metabolic sensing *in situ*. The battery-free e-skin contains multimodal sensors and highly efficient lactate biofuel cells that use a unique integration of zero- to three-dimensional nanomaterials to achieve high power intensity and long-term stability. The PPES delivered a record-breaking power density of 3.5 milliwatt-centimeter<sup>-2</sup> for biofuel cells in untreated human body fluids (human sweat) and displayed a very stable performance during a 60-hour continuous operation. It selectively monitored key metabolic analytes (e.g., urea, NH<sub>4</sub><sup>+</sup>, glucose, and pH) and the skin temperature during prolonged physical activities and wirelessly transmitted the data to the user interface using Bluetooth. The PPES was also able to monitor muscle contraction and work as a human-machine interface for human-prostheses walking.

## INTRODUCTION

Recent advances in robotics have enabled soft electronic devices at different scales with excellent biocompatibility and mechanical properties; these advances have rendered novel robotic functionalities suitable for various medical applications, such as diagnosis and drug delivery, soft surgery tools, human-machine interaction (HMI), wearable computing, health monitoring, assistive robotics, and prosthesis (1–6). Electronic skin (e-skin) can have similar characteristics to human skin, such as mechanical durability and stretchability and the ability to measure various sensations such as temperature and pressure (7–11). Moreover, e-skin can be augmented with capabilities beyond those of the normal human skin by incorporating advanced bioelectronics materials and devices. For example, the compliance of soft e-skin enables wearable sensing of biochemicals from the environment and the human body, showing great potential for wearable personalized health monitoring at molecular levels (12–16).

Sophisticated tasks in practical robotic and biomedical applications—including multiplexed sensing, on-demand actuation, and wireless data transmission during prolonged operation—set high standards for energy consumption (17). Batteries have been used as the primary power source of the majority of wireless e-skin systems; however, existing ones suffer from inadequate long-term continuous usability, especially when electricity is not readily available (18–20). Soft e-skins that can directly harvest energy from other accessible sources—including human motion, body heat, and solar light—are limited by their low-power density. Thus, they fail to power signal-processing circuitry and wireless data transmission (e.g., Bluetooth or radiofre-

quency communication) (21, 22). Battery-free e-skins using near-field communication (NFC) for wireless power and data transmission with the user interface have been developed recently; however, these approaches are limited by the required proximity for data processing and readout (23, 24).

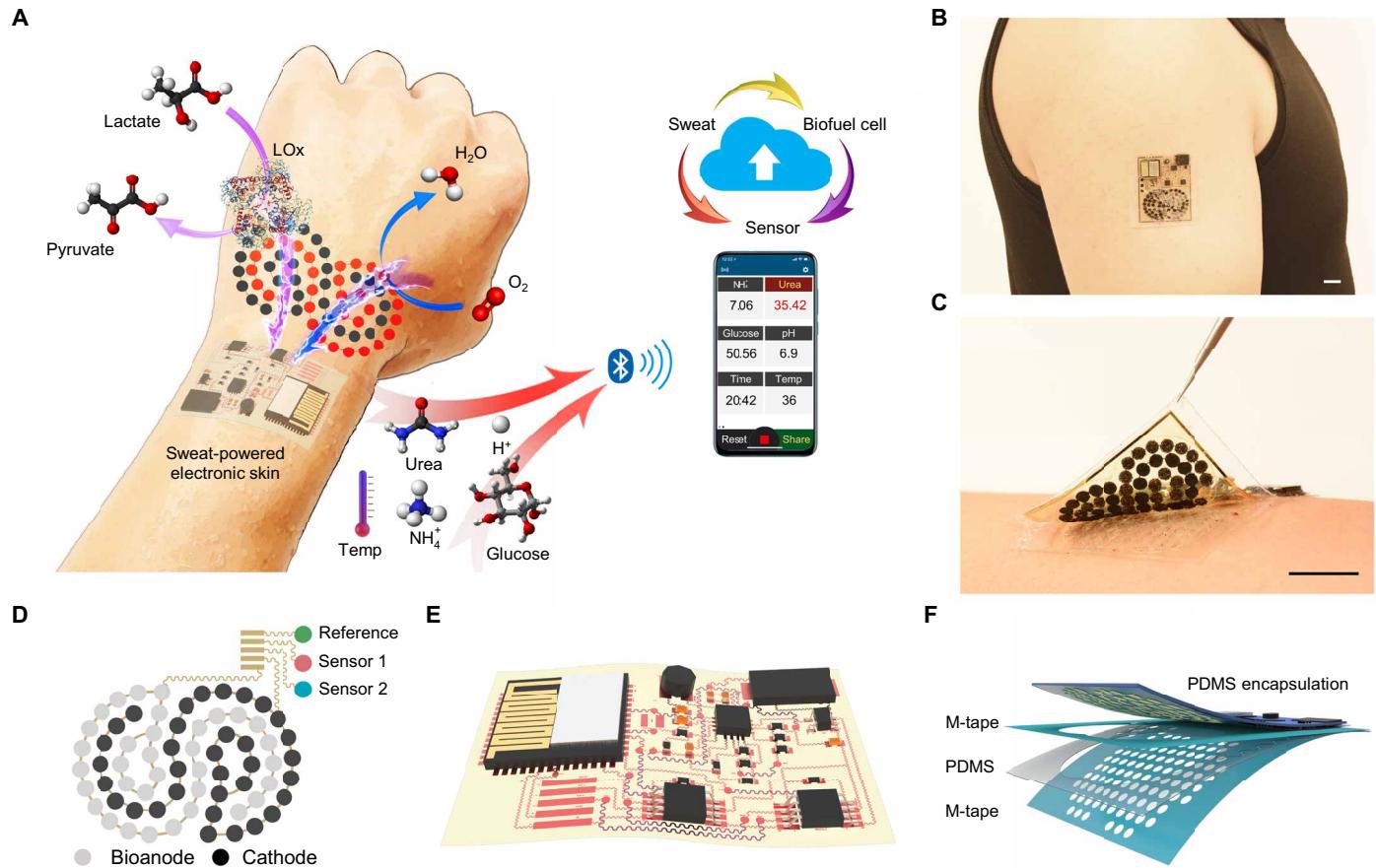
Human biofluids, such as human sweat, could serve as an ideal and sustainable bioenergy source for powering future e-skin devices (25, 26). Moreover, they contain a wealth of chemicals that can reflect an individual's health status (27–29). Recent advances in wearable sweat analysis have enabled a number of fundamental and clinical applications, such as noninvasive metabolic monitoring and disease diagnosis (30–32).

Here, we report a battery-free, fully perspiration-powered electronic skin (PPES) that harvests energy from human sweat through lactate biofuel cells (BFCs), performs continuous multiplexed monitoring of key metabolic biomarkers (e.g., glucose, urea, NH<sub>4</sub><sup>+</sup>, and pH), and wirelessly transmits the personalized information to a user interface via Bluetooth low energy (BLE) (Fig. 1A). Built on an ultra-soft polymeric substrate, the PPES can comply with the skin's modulus of elasticity and conformally laminate on different body parts for accurate biosensing (Fig. 1, B and C). Using cross-dimensional nanomaterial integration, the highly efficient and stable nanoengineered BFC array was able to achieve a record-breaking and sustainable high-power density, enabling continuous self-powered health monitoring. On this basis, we report the use of BFCs to fully power an e-skin with both multiplexed sensing and wireless data-transmission capabilities. The PPES was successfully validated *in vivo* in a cycling human trial, and its use toward noninvasive metabolic monitoring was further evaluated in humans involving dietary and nutrition challenges. Moreover, we demonstrate the use of the PPES as a human-machine interface for assistive robotic control: Integrated with strain sensors, the self-powered e-skin could wirelessly transmit signals to a user interface to direct a human operator to control a robotic prosthesis. This technology may substantially advance both self-powered electronic skin and personalized health care.

Copyright © 2020  
The Authors, some  
rights reserved;  
exclusive licensee  
American Association  
for the Advancement  
of Science. No claim  
to original U.S.  
Government Works

<sup>1</sup>Andrew and Peggy Cherng Department of Medical Engineering, California Institute of Technology, Pasadena, CA 91125, USA. <sup>2</sup>Department of Electrical Engineering, California Institute of Technology, Pasadena, CA 91125, USA. <sup>3</sup>Division of Chemistry and Chemical Engineering, California Institute of Technology, Pasadena, CA 91125, USA. <sup>4</sup>Department of Mechanical and Civil Engineering, California Institute of Technology, Pasadena, CA 91125, USA.

\*Corresponding author. Email: weigao@caltech.edu



**Fig. 1. Perspiration-powered soft electronic skin (e-skin) for multiplexed wireless sensing.** (A) Schematic of a battery-free, biofuel-powered e-skin that efficiently harvests energy from the human body, performs multiplexed biosensing, and wirelessly transmits data to a mobile user interface through Bluetooth. (B and C) Photographs of a PPES on a healthy individual's arm. Scale bars, 1 cm. (D and E) Schematic illustrations of the flexible BFC-biosensor patch (D) and the soft electronic-skin interface (E). (F) System-level packaging and encapsulation of the PPES for efficient on-body biofluid sampling. M-tape, medical tape.

## RESULTS

### Design of the soft PPES for multiplexed sensing

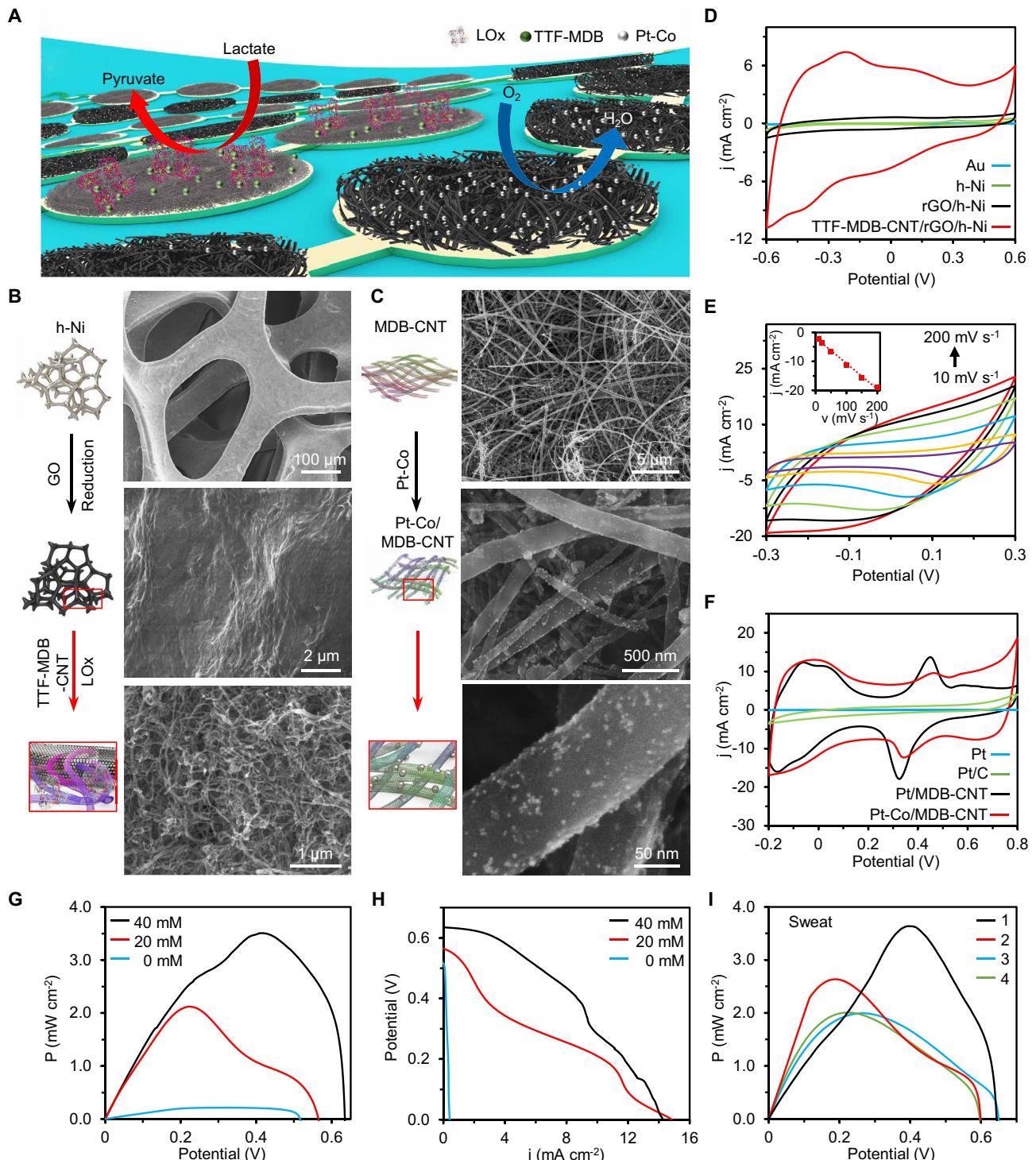
The PPES consists of two main parts: (i) A nanoengineered flexible electrochemical patch contains a BFC array and a biosensor array for energy harvesting and molecular analysis in human sweat; biosensing films and biocatalytic nanomaterials are immobilized on serpentine-connected electrode arrays (Fig. 1D). (ii) A flexible electronic patch consolidates the rigid electronics on an ultrathin polyimide (PI) substrate through flexible interconnects for power management, signal processing, and wireless transmission (Fig. 1E). A skin-interfaced microfluidic module was integrated into the PPES to achieve efficient fresh sweat sampling for stable BFC operation and accurate sweat analysis with high temporal resolution (Fig. 1F). The independent inlet-outlet design of the microfluidics for BFCs and sensors further minimized the potential influences of the BFC by-products on the sensing accuracy. The electronic components and interconnects of the PPES were encapsulated with polydimethyl siloxane (PDMS) to avoid sweat/electronic contact (Fig. 1F).

### Design and characterization of nanoengineered BFCs for highly efficient energy harvesting

BFCs, using the enzymes as biocatalysts to transform the bioenergy into the electricity, are attractive power sources for future wearable and implantable electronics (33, 34). Among bioenergy resources,

lactate, the main metabolic product of both muscle and brain exertion, is present in sweat at tens of millimolar levels and ideally suitable for powering future skin-interfaced electronic devices (35, 36). However, reports of BFCs-powered wearable sensing remain very limited. The main bottlenecks/challenges for the practical use of current BFCs to power e-skin are (i) limited power density harvested from the human body (table S1) and (ii) poor BFC stability and short lifetime. Here, we used a BFC configuration that consists of lactate oxidase (LOx) immobilized bioanodes to catalyze the lactic acid to pyruvate and Pt alloy nanoparticle-decorated cathodes that reduce oxygen to water (Fig. 2A). Such redox reactions on BFC electrodes yield a stable current to power the electrical loads. Monolithic integration of zero-dimensional (0D) to 3D nanomaterials was used on the BFC electrodes to obtain optimal energy harvesting performance (Fig. 2, B and C).

To prepare the bioanode, we sequentially modified hierarchical Ni (h-Ni) microstructures, reduced graphene oxide (rGO) films, and Meldola's blue-tetrathiafulvalene-modified carbon nanotubes (MDB-TTF-CNT) on an Au electrode array (Fig. 2B). The extremely high current density and stability of the MDB-TTF-CNT/rGO/h-Ni bioanode illustrated in Fig. 2D and fig. S1 can be attributed to (i) the high electrochemically active surface area (ECSA) that is increased by 3000 times after nanomaterial modification (Fig. 2E, fig. S2, and Materials and Methods); (ii) the π-π interaction between the CNTs and rGO that enhances the electron-transfer rate between LOx and



**Fig. 2. Flexible nanoengineered BFC array for efficient energy harvesting.** (A) Schematic of a soft BFC array consisting of Lox-modified bioanodes and Pt alloy nanoparticle-modified cathodes. (B and C) Schematic and SEM images showing modification process of bioanode assembled with porous h-Ni, rGO, CNTs, and Lox (B), and cathode assembled with CNT film and Pt alloy nanoparticles (C). (D) CVs of the Au, h-Ni, rGO/h-Ni, and MDB-CNT/rGO/h-Ni electrodes.  $j$ , current density. Scan rate, 50 mV s<sup>-1</sup>. (E) CVs of the MDB-CNT/rGO/h-Ni with scan rates of 10, 20, 50, 100, 150, and 200 mV s<sup>-1</sup>. Inset: Calibration curve of the current density at  $\sim$ 0.1 V versus scan rate ( $v$ ). (F) CVs of the Pt-Co/MDB-CNT, Pt/MDB-CNT, commercial Pt/C, and bulk Pt electrodes recorded in  $N_2$ -purged 0.1 M  $H_2SO_4$  solutions. Pt/C, platinum carbon black. Scan rate, 100 mV s<sup>-1</sup>. (G and H) Power density ( $P$ ) (G) and polar curves (H) of BFCs recorded in 0 to 40 mM lactate. Experiments in (D) to (H) were repeated three times independently with similar results. (I) Power density curves of BFCs in sweat samples from four healthy humans. Reference electrodes in (D) to (I), Ag/AgCl.

electrodes; (iii) the TTF-MDB redox mediator that decreases the overpotential of the lactate oxidation reaction (text S1).

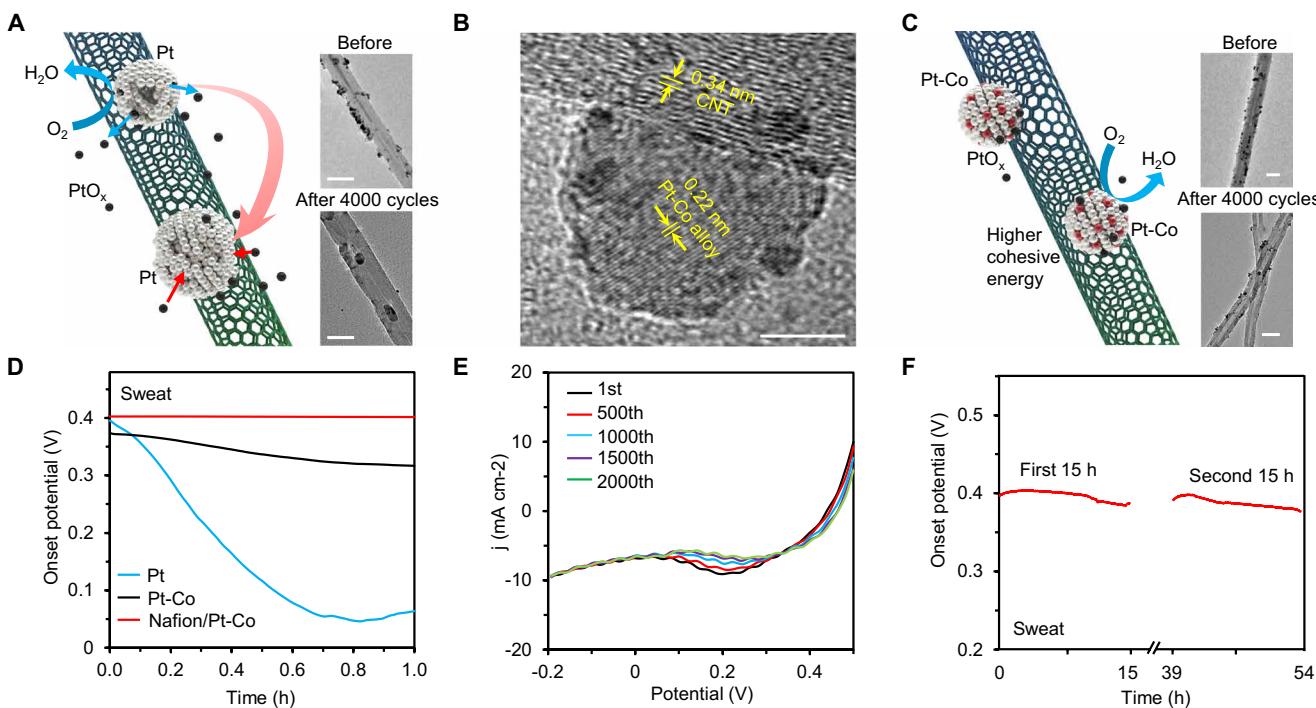
On the other hand, Pt-based nanoparticles were decorated on an MDB-modified CNT network (MDB-CNT) through electroless plating to form the BFC cathode (Fig. 2C). Note that the MDB modification was crucial to achieve uniform nanoparticle distribution with controlled sizes (figs. S3 and S4). Compared with that of the conventional bulk Pt electrode, Pt and Pt-Co nanoparticle-coated MDB-CNT electrodes showed higher ECSA of 170 and 210 times, respectively (Fig. 2F, fig. S5, and Materials and Methods) (37).

The assembled BFC array showed excellent performance, with an open-circuit potential (OCP) at ~0.6 V and maximum power outputs of ~2.0 and ~3.5 mW cm<sup>-2</sup> in 20 and 40 mM lactate solutions, respectively (Fig. 2, G and H). Note that the maximum power densities shifted as the lactate concentrations changed because of the varied redox reaction of MDB at different pHs (fig. S6) (38). When operating in human sweat samples, the BFCs could provide a power density as high as 3.6 mW cm<sup>-2</sup> (Fig. 2I). This is, to the best of our knowledge, the highest power density for BFCs in untreated human body fluids (table S1).

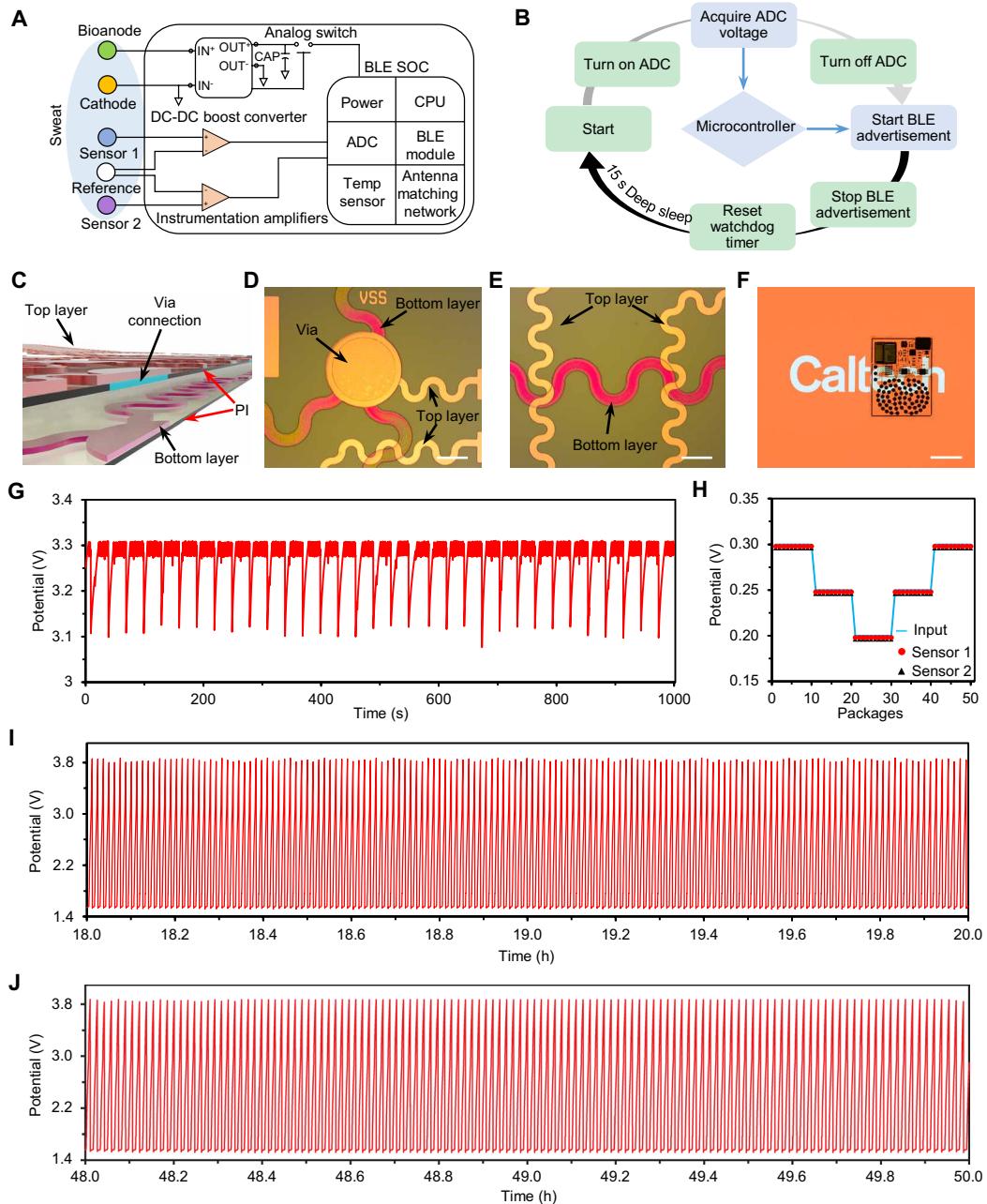
### Characterization of Pt-Co nanoparticle-decorated BFC cathode for enhanced stability

Enzymatic BFCs usually suffer from poor long-term stability, primarily due to the limited stability of their cathode. Among commonly used cathode materials, Prussian blue, Ag<sub>2</sub>O, and MnO<sub>2</sub> are consum-

able and cannot be recovered in situ (39, 40). Pt could be easily fouled during the practical uses due to the formation of the —OH groups on the electrode surface (Fig. 3A and fig. S7) (41); the dissociated oxidized Pt from the nanoparticles tends to redeposit on the larger particles via the Ostwald ripening mechanism (the average cohesive energy of Pt atoms in smaller nanoparticles is smaller than that in larger ones) (42). To enhance the long-term stability of the Pt particles, transition-metal dopants (i.e., Co) are introduced through electroless codeposition (43, 44). The resulting Pt/Co alloy nanoparticles were characterized using transmission electron microscopy (TEM) (Fig. 3B) and energy-dispersive x-ray spectroscopy (fig. S8). The Pt/Co nanoparticles showed an average size of ~10 to 20 nm and an edge lattice spacing of 0.22 nm (Fig. 3B). The Co dopants could enhance the cohesive energy and thus stabilize the nanoparticles, leading to substantially reduced biofouling in body fluids and higher onset potential for oxygen reduction (45). In sweat samples, the Pt-Co/CNT showed a relatively stable onset potential compared with that of Pt/CNT (Fig. 3D). To further improve the long-term stability of the cathode in biofluids, we modified a permselective Nafion layer onto the Pt-Co/CNT (Fig. 3D). The Nafion/Pt-Co/CNT cathode showed stable performance over 2000 cycles of cyclic voltammetry (CV) scans (Fig. 3E) and a negligible fluctuation in the onset potential in a sweat sample for more than 30 hours (Fig. 3F). In addition to high stability, Pt-Co/MDB-CNT cathodes also showed the highest oxygen reduction performance compared with those modified with Pt or other Pt alloys (fig. S5), attributed to the Sabatier principle (46).



**Fig. 3. Characterization of role of Pt alloy nanoparticles in the BFC stability.** (A) Schematic illustration showing that the Pt nanoparticles (decorated on CNTs) merge into larger ones during the prolonged catalysis. Inset: TEM images of the Pt/MDB-CNT before and after 4000 CV cycles between −0.2 and 0.5 V. Scale bars, 50 nm. (B) TEM image showing the crystalline structure of a Pt-Co nanoparticle. Scale bar, 5 nm. (C) Schematic illustration of the Pt-Co alloy nanoparticles on CNT, which maintain the uniform size owing to their higher surface-cohesive energy level. Inset: TEM images of the Pt-Co/MDB-CNT before and after 4000 cycles between −0.2 and 0.5 V. Scale bars, 50 nm. (D) The onset potentials of Pt/MDB-CNT-, Pt-Co/MDB-CNT-, and Nafion/Pt-Co/MDB-CNT-modified cathodes measured in sweat more than a 1-hour period. (E) Repetitive LSVs of a Nafion/Pt-Co/MDB-CNT-modified cathode obtained during 2000 CV cycles between −0.2 and 0.5 V. (F) Long-term stability test of a Nafion/Pt-Co/MDB-CNT cathode in sweat more than 30 hours. Reference electrodes in (A) and (C to F), Ag/AgCl. Experiments in (D) to (F) were repeated three times independently with similar results.



**Fig. 4. System-level integration and evaluation of the PPES.** (A) Schematic diagram of the PPES system including the BFC array, sensor array, boost converter, instrumentation amplifiers, and BLE module. SOC, system-on-chip; CPU, central processing unit. (B) Operation flow of the energy control and data-transmission processes. (C to E) Schematic (C) and microscopic images (D and E) showing the via structure and soft interconnects of the e-skin system. Scale bars, 3  $\mu$ m. (F) A photographic image of a fully integrated PPES. Scale bar, 2 cm. (G) Real-time potential of the capacitor measured during continuous operation of a PPES in 20 mM lactate. (H) Characterization of BFC-powered multiplexed sensing with varied electrical inputs in 20 mM lactate. (I and J) Long-term stability of the capacitor (220  $\mu$ F)-charging process with BFCs through the DC-DC converter more than 50 hours in 20 mM lactate. Potential dynamics from 18 to 20 hours and from 48 to 50 hours are illustrated in (I) and (J), respectively. Experiments in (G) to (J) were repeated three times independently with similar results.

### System-level integration of the PPES for energy management and signal processing

The fully integrated PPES consists of a nanoengineered BFC array (total bioanode area, 1  $\text{cm}^2$ ), a boost converter, a biosensor array, instrumentation amplifiers, and a programmable system on chip (PSoC) module (integrated with a BLE module, a microcontroller, and a temperature sensor) (Fig. 4A and figs. S9 and S10). The DC-DC

boost converter amplifies the signal potential with a small power loss (~20%) (fig. S9). The output signal (3.3 V) continuously charges a capacitor (660  $\mu$ F) that temporarily stores the energy and powers biosensors and other electronic components. The BLE module runs in bursts of activity, periodically waking up from deep sleep mode to acquire measurements with the embedded successive-approximation analog-to-digital converter then wirelessly broadcasting the data to

the user interfaces (Fig. 4B). BLE advertising is chosen here owing to the small size of the data packets and the low-power consumption.

The PPES is patterned on an ultrasoft PI substrate through a standard micro/nanofabrication process to comply with the skin's modulus of elasticity for accurate wearable biosensing (figs. S11 to S14). The fabricated via connection and serpentine-shaped interconnects in the PPES are illustrated in Fig. 4 (C to E). To minimize the overall patch size, reduce the strain, and achieve uniform strain distribution of the patch during mechanical deformation, we designed the whole BFC electrode array as a serpentine structure (fig. S15). The fully encapsulated PPES shows excellent transparency, mechanical flexibility, and stability (Fig. 4F and fig. S16).

In deep sleep mode, the whole PPES system draws a total current of  $\sim 100 \mu\text{A}$  at 3.3 V, primarily from the two instrumental amplifiers (fig. S17A). The PPES is programmed to wake up periodically from deep sleep for  $\sim 10$  ms for acquiring and sending the sensing data with an average consumption of 9.35 mA (fig. S17B). As a result, the capacitor discharges when the PPES wakes up and is recharged within a few seconds by the BFCs. In 20 mM lactate, dynamic changes in the potential of the capacitor on the PPES are shown in Fig. 4G. Wirelessly received BLE data in the user interface showed good agreement with the dual-channel sensor inputs (Fig. 4H). Our results demonstrate that, by using a smaller capacitor (220 or 400  $\mu\text{F}$ ), the whole PPES can be continuously powered in lactate solutions (5 to 20 mM) without the need of deep sleep mode (fig. S18). Excellent long-term stability of the BFC-based electrical charging/discharging process was demonstrated by the continuous charging activity for  $\sim 60$  hours, as shown in Fig. 4 (I and J) (and fig. S19): A capacitor was charged from 1.5 to 3.8 V continuously and repeatedly, and the charging periods could remain stable when fresh lactate fuel was supplied (fig. S19).

### Characterization of the PPES for multiplexed biosensing

The PPES holds great promise for multiplexed sensing, and a particularly attractive one is wearable sweat analysis for personalized health monitoring. Important sweat biomarkers—such as urea, glucose, pH, and  $\text{NH}_4^+$ —contain meaningful information about an individual's physiological status. Despite high interest, only a limited number of sweat analytes can be accurately monitored by currently reported wearable sensors (12, 13, 47). Given the complicated sweat secretion process and the electrochemical sensing mechanisms, multiplexed sensing is often essential to achieve an accurate assessment of the specific analyte. As an example, two sweat biosensor arrays (urea and  $\text{NH}_4^+$  sensor array and glucose and pH sensor array) were developed and evaluated toward metabolic monitoring.

The  $\text{NH}_4^+$  and urea sensor array was designed on the soft electrochemical patch based on the  $\text{NH}_4^+$  ion-selective electrodes (ISEs) (Fig. 5A). Compared with the  $\text{NH}_4^+$  sensor, the urea sensor contains an additional enzymatic layer where urease converts urea to carbon dioxide and ammonia; the increased  $\text{NH}_4^+$  product reflects the urea level. Figure 5 (B and C) shows the potentiometric responses of the urea and  $\text{NH}_4^+$  sensors, measured in 40 to 2.5 mM  $\text{NH}_4^+$  solutions and 40 to 2.5 mM urea solutions, respectively. A linear relationship between potential output and logarithmic concentrations of the target analytes was obtained, with near-Nernstian sensitivities of 60.3 and 60.0 mV per decade of concentration for  $\text{NH}_4^+$  and urea sensors, respectively. The sensors have good selectivity over common analytes in human sweat (fig. S20). The dependence of urea and  $\text{NH}_4^+$  concentrations on the sensor response is illustrated in Fig. 5D and fig. S21. Considering that  $\text{NH}_4^+$  level has a significant influence on urea sen-

sor reading, it is essential to simultaneously monitor both urea and  $\text{NH}_4^+$  with real-time calibration for accurate sweat analysis.

The glucose and pH sensor array was prepared using a similar potentiometric sensing approach (Fig. 5E). A sandwich structure, Nafion/chitosan (CS)-glucose oxidase (GOx)/Nafion, was coated on the platinum-deposited electrode to form a highly sensitive and selective glucose sensor; an electropolymerized polyaniline film serves as the hydrogen ion-selective film for pH sensing. Figure 5 (F and G) illustrates the responses of the glucose and pH sensors in 40 to 200  $\mu\text{M}$  glucose and pH 4 to 8 solutions, respectively. A linear response between the potential output of glucose sensor and glucose concentrations (in a physiologically relevant range of 0 to 150  $\mu\text{M}$ ) was obtained with a sensitivity of  $0.1 \text{ mV } \mu\text{M}^{-1}$ . A near-Nernstian sensitivity of 55.3 mV per pH was observed for the pH sensor. Considering that the glucose sensor response is heavily dependent on the solution pH (Fig. 5H), multiplexed glucose and pH sensing with real-time calibration is also crucial to obtain high sensing accuracy (fig. S22).

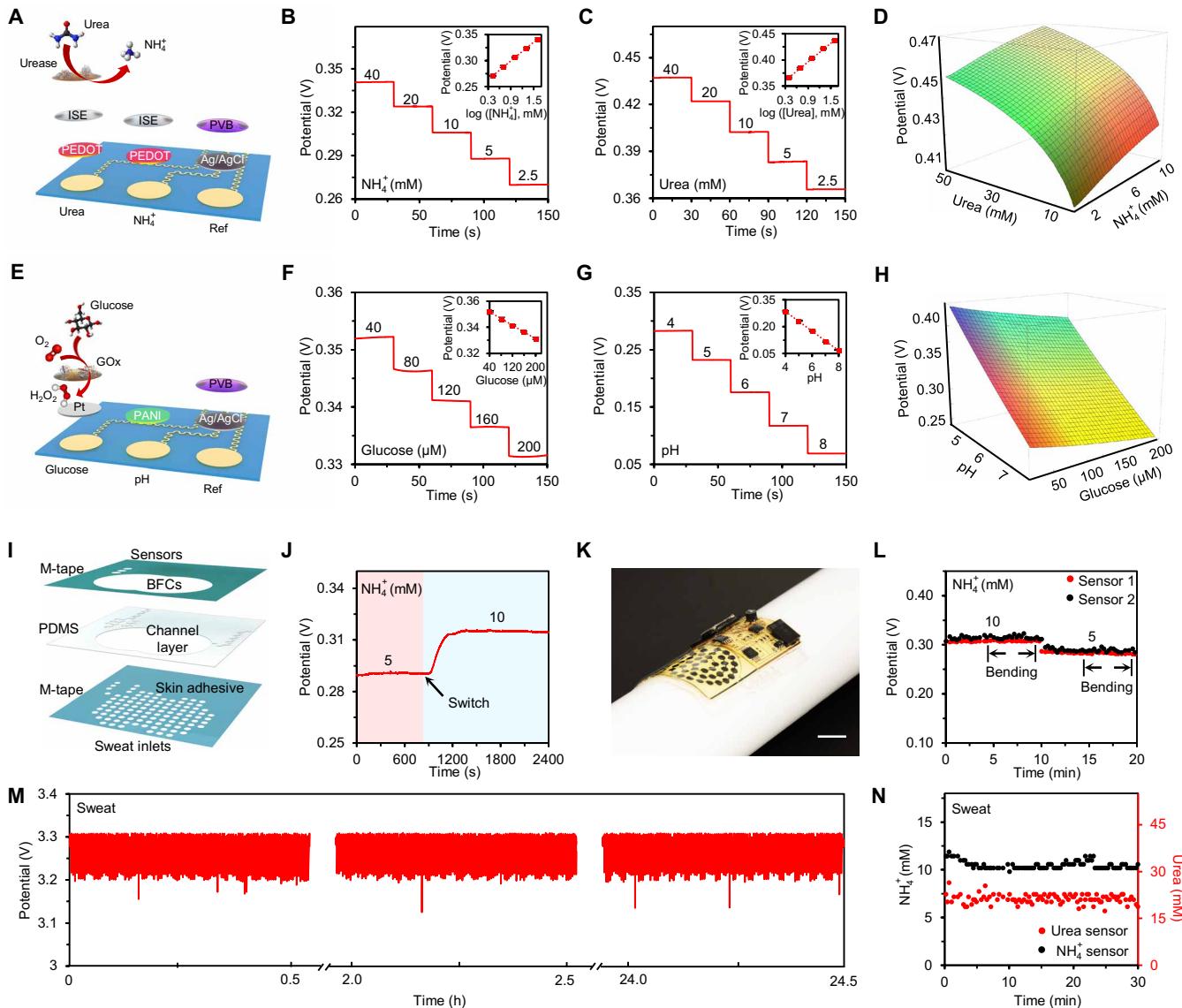
All the sensors showed excellent long-term electrochemical and mechanical stabilities during continuous operation, indicating their promise for wearable use (figs. S23 and S24). Considering that skin temperature has a direct influence on the enzymatic biosensors (glucose and urea sensors here as shown in fig. S25), the on-chip temperature sensor in the BLE module could provide the skin temperature information for real-time calibration.

For wearable on-body use, the integration of a microfluidic module could greatly enhance the sweat sampling process and lead to a higher temporal resolution for wearable sensing and more stable power output from BFCs (Fig. 5I). The laser-patterned microfluidics was assembled in a sandwich structure (M-tape/PDMS/M-tape) and contains two reservoirs to minimize the influence of the BFC by-products on the sensing accuracy (fig. S26). An in vitro flow test showed that when the  $\text{NH}_4^+$  level in the input solution is switched from 5 to 10 mM at a physiologically measured sweat rate of  $0.05 \text{ ml hour}^{-1}$ , it takes  $\sim 4$  min for the  $\text{NH}_4^+$  sensor to reach new stable reading (Fig. 5J), indicating the small time delay for the on-body continuous monitoring. The PPES is mechanically flexible and can conformally laminate on a curved substrate (Fig. 5K). Under mechanical deformation with a bending curvature of 1.5 cm in radius, the PPES maintained consistent sensor readings (Fig. 5L). During long-period operation in human sweat samples, the PPES demonstrated stable performance in both energy harvesting (Fig. 5M) and analyte monitoring (Fig. 5N).

### On-body validation and evaluation of the PPES toward wearable biosensing

On-body validation of the PPES was conducted on healthy humans toward continuous metabolic monitoring during a constant-load stationary biking exercise (Fig. 6A). Figure 6 (B and C) shows the data collected from the PPES for real-time monitoring of sweat urea,  $\text{NH}_4^+$ , glucose, and pH. The pH and  $\text{NH}_4^+$  levels measured here, along with the on-chip temperature sensor readings, were also used to calibrate the readings from the glucose and urea sensors, respectively. During the biking process, the urea and  $\text{NH}_4^+$  levels in sweat decreased rapidly and then stabilized over time (Fig. 6B). A similar trend was observed for sweat glucose, whereas a stable pH response throughout the exercise was obtained (Fig. 6C). The PPES showed good reusability, stability, and biocompatibility during long-term usage (fig. S27).

In addition to on-body validation, the use of the PPES in metabolic and nutritional management was evaluated through controlled

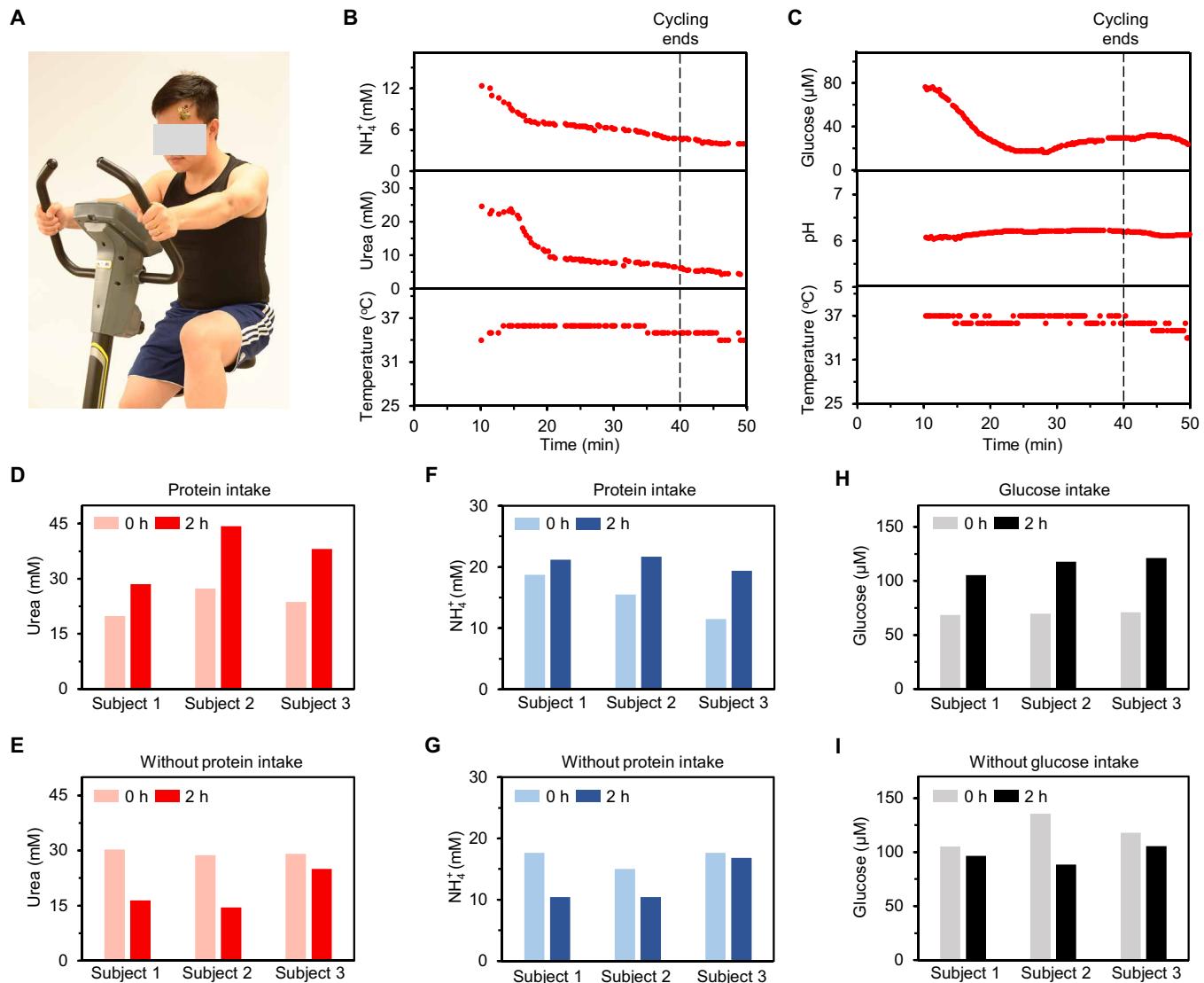


**Fig. 5. Characterization of the PPES for multiplexed biosensing.** (A) Schematic of a sensor array for simultaneous urea and  $\text{NH}_4^+$  monitoring. (B) and (C) OCP responses of a  $\text{NH}_4^+$  sensor (B) and a urea sensor (C) in standard analyte solutions. Insets: The corresponding calibration plots of the  $\text{NH}_4^+$  and urea sensors. Data recording was paused for 30 s for each solution change. Error bars represent the SDs from five sensors. (D) A color map showing the dependence of the urea sensor response on urea and  $\text{NH}_4^+$  concentrations. Red indicates higher  $\text{NH}_4^+$  level and green indicates higher urea level. (E) Schematic of a sensor array for simultaneous glucose and pH sensing. (F) and (G) OCP responses of a glucose sensor (F) and a pH sensor (G) in standard analyte solutions. Insets: The corresponding calibration plots of the glucose and pH sensors. Data recording was paused for 30 s for each solution change. Error bars represent the SDs from five sensors. (H) A color map showing the dependence of the glucose sensor on the glucose and pH levels. Red indicates higher pH value and green indicates higher glucose level. (I) Schematic of the microfluidic design for biofluid sampling. (J) Dynamic response of an  $\text{NH}_4^+$  sensor upon switching the inflow solutions at a flow rate of 0.05 ml hour<sup>-1</sup>. (K) A photograph of an integrated PPES under mechanical deformation (with a radius of bending curvature of 1.5 cm). (M) and (N) Potential of the capacitor (M) and responses of a urea and  $\text{NH}_4^+$  sensor array (N) when the PPES operates in a human sweat sample.

dietary challenges (Fig. 6, D to I). Figure 6 (D to G) demonstrates that, compared with the initial levels, sweat urea and  $\text{NH}_4^+$  levels measured 2 hours after a standardized protein intake significantly increased in all three individuals. In contrast, decreased trends were obtained during the 2-hour period from all the individuals without protein intake. In an oral glucose-tolerance test, sweat glucose levels increased markedly for all individuals after the glucose intake and decreased after 2 hours for individuals without intake. These data indicate that the PPES has a great potential in self-powered personalized physiological and metabolic monitoring.

### Evaluation of the PPES as a human-machine interface for robotic assistance

HMI has attracted substantial attention over the past decade owing to its high promise in real-world biomedical applications in rehabilitation. On skin, detection of strains induced by muscle contraction using soft strain sensors integrated in e-skin systems is one promising approach to enable dynamic HMI. When integrated with soft strain sensors, the skin-interfaced soft PPES could function as a human-machine interface toward robotic applications (Fig. 7A and figs. S28 and S29). The strain sensors were designed on the basis of



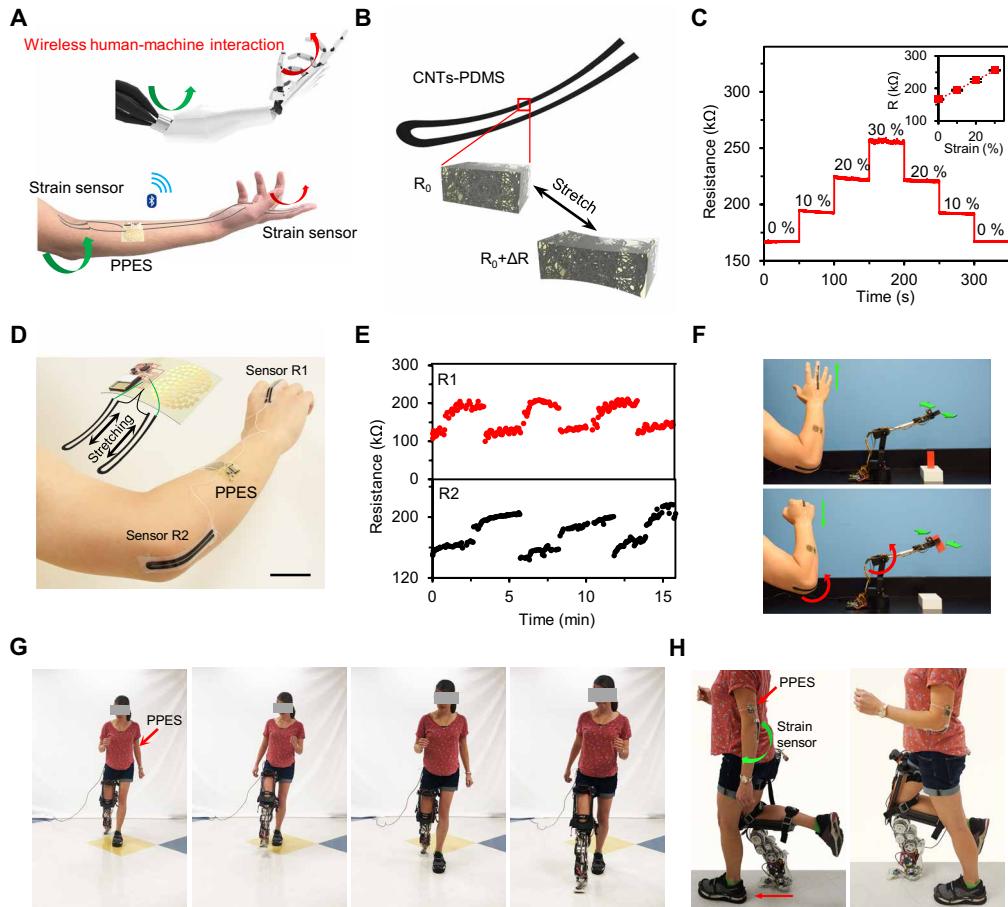
**Fig. 6. On-body evaluation of the PPES toward personalized metabolic monitoring.** (A) Photograph of an individual wearing a PPES during cycling exercise. (B) Real-time multiplexed urea and  $\text{NH}_4^+$  analysis using a PPES on an individual's forehead. (C) Real-time multiplexed glucose and pH monitoring using a PPES on an individual's forehead. (D to I) Evaluation of the PPES in dietary challenges: sweat urea levels with (D) and without (E) protein intake, sweat  $\text{NH}_4^+$  levels with (F) and without (G) protein intake, and sweat glucose levels with (H) and without (I) glucose intake.

CNTs/PDMS elastomer: The resistance of the sensors increased linearly with the applied strain (Fig. 7, B and C). Two strain sensors were placed on the hand and the elbow, respectively, and connected to the PPES (Fig. 7D); the bending of the finger and elbow could be monitored from the resistance change of the strain sensors (Fig. 7E). Each resistive-type strain sensor as part of a voltage divider consumed a total of  $\sim 5 \mu\text{A}$ . The battery-free e-skin placed on the sweaty arm was able to wirelessly control the motion of a robotic arm in real time: The robotic arm recognized the gestures of the human arm and then approached and grabbed the target object (Fig. 7F and movie S1). The PPES could also be used for robotic assistance in the rehabilitation settings. Figure 7 (G and H) (and corresponding movie S2) demonstrates that prosthesis walking control could be achieved with a strain sensor-integrated PPES to achieve assistive walking in real-world environments. By incorporating more physical sensors for electroencephalogram and electromyography recording along with the continuous

metabolic monitoring, the multimodal PPES could facilitate the design and optimization of novel prostheses that bring the human into the loop of prosthesis control to enable real-time user-specific responses to human intent and behavior.

## DISCUSSION

Here, we developed a flexible, fully integrated, and self-powered e-skin platform that can provide real-time, continuous, multiplexed sensing and wirelessly transmit the data to the user interface through Bluetooth communication. Our e-skin platform provided real-time chemical sensing of  $\text{NH}_4^+$ , urea, glucose, and pH, enhancing the capabilities of e-skin with multimodal and multianalyte detection. Moreover, the e-skin platform was able to monitor physical parameters—such as temperature, strain, and pressure—enabling self-powered wireless human-machine interaction for robotic applications such as powered



**Fig. 7. On-body evaluation of the PPES as a wireless human-machine interface for robotic assistance.** (A) Schematic illustration of the PPES for remote human-machine interaction. (B) Schematic of the CNTs-PDMS elastomer-based stain sensors. (C) Resistance response of a CNTs-PDMS strain sensor under different strains. (D) Photograph and schematic (inset) of the PPES integrated with strain sensors. (E) Real-time multidegree motion tracking using a PPES with the strain sensors on an individual's finger and elbow. (F) Time-lapse images of the wireless robotic arm control using a PPES. (G and H) Time-lapse images of front view (G) and side view (H) of the use of the PPES for robotic prosthesis control.

prostheses walking. The incorporation of multiplexed, multimodal chemical and physical sensing may expand the potential applications of e-skin for personalized health care.

For an e-skin to perform real-time and continuous sensing, on-board powering is necessary. Most e-skin platforms use the battery power supply, which could introduce constraints on the duration of use between charging and affect the long-lasting functionality when the electricity is not readily available. To circumvent the battery requirements, other wireless platforms have used battery-free data transmission/powering strategies, such as NFC, that require very small readout distance. The use of BLE transmission alleviates such constraints and offers a more realistic communication scheme for e-skin/wearable applications but would impose a higher power consumption. To accommodate the use of BLE, various battery-free powering strategies have been tested, such as piezo-based or biofuel-based energy harvesting. However, these strategies either have limited power densities for wearable or robotic use that fail to support the BLE functionality on a compact e-skin platform or suffer from short lifetime caused by biofouling.

The PPES proposed here has successfully resolved both challenges for on-board integration of BLE within the compact yet flexible

e-skin format. With a unique integration of 0D-3D nanomaterials on bioanodes and cathodes, the BFC designed here yields a record-breaking power density (as high as  $3.5 \text{ mW cm}^{-2}$  in human sweat) that could support sensing and BLE functionalities on a small e-skin patch. The use of carefully designed and selected Pt-Co alloy nanoparticles has greatly enhanced the long-term stability of the BFC in sweat and enabled the long-term continuous use on skin.

Moreover, the developed e-skin here is a fully integrated platform with enhanced wearability and sensing accuracy. The platform is ultrathin and transparent using electronics with low-power consumption and integrating electronics onto the soft substrate with a minimal mechanical mismatch, rendering a fully compliant e-skin for maximal comfort and wearability. The use of microfluidics significantly improves sweat refreshing and reduces interference between BFC and biosensors; a carefully studied correlation between different chemical targets yields calibrated sensor readouts that reflect higher accuracy. The system-level integration of electronics, microfluidics, and on-board calibration on a soft e-skin provides a much-enhanced accuracy of sensing results for on-body use. Moreover, the successful demonstration of the prosthetic control for robotic assistance using the PPES indicates

the promise of using such platform for human machine interaction and for design/optimization of next-generation prostheses. The development of such a multimodal, fully integrated, and self-powered platform enriches the functionalities and potentials of e-skins and opens the door to numerous robotic and wearable healthcare possibilities, such as personalized medicine.

## MATERIALS AND METHODS

### Materials

Graphite flake and silver nitrate were purchased from Alfa Aesar. Sodium nitrate, potassium permanganate, hydrogen peroxide, multiwalled CNTs, TTF, citric acid, chloroplatinic acid, sodium borohydride, ammonium chloride, nickel chloride, copper acetate, iron(III) chloride, copper acetate, zinc nitrate, Nafion perfluorinated resin solution (5%), sodium thiosulfate pentahydrate, sodium bisulfite, polyvinyl chloride (PVC), polyvinyl butyral (PVB), F127, nonactin, bis(2-ethylhexyl) sebacate (DOS), 3,4-ethylenedioxythiophene (EDOT), poly(sodium 4-styrenesulfonate) (PSS), CS, PDMS, aniline, and platinum carbon black (Pt/C) (0.5%) were purchased from Sigma-Aldrich. Sulfuric acid, hydrochloric acid, ascorbic acid, methanol, ethanol, acetone, tetrahydrofuran, sodium

chloride, disodium phosphate, urea, and dextrose ( $\text{D}$ -glucose) were purchased from Fisher Scientific. MDB was purchased from Combi-Blocks Inc. Lactate oxidase ( $106 \text{ U mg}^{-1}$ ) was purchased from Toyobo Co. The CNT film and Ni substrate (h-Ni) were purchased from NTL Inc. and MXBaocheng Products, respectively. Medical tapes were purchased from Adhesives Research. PIs (PI-2611 and PI-2610) were purchased from HD MicroSystems Inc. Silver conductive paint was purchased from Structure Probe, Inc. (SPI supplies).

### Preparation of the BFCs

To prepare the bioanodes, we first prepared GO suspension following the modified Hummer's method (48). Briefly, a mixture of 1 g of graphite flake and 23 ml of  $\text{H}_2\text{SO}_4$  was stirred more than 24 hours, and then 100 mg of  $\text{NaNO}_3$  was added into the mixture. Subsequently, 3 g of  $\text{KMnO}_4$  was added to the mixture below  $5^\circ\text{C}$  in the ice bath. After stirring at  $40^\circ\text{C}$  for another 30 min, 46 ml of  $\text{H}_2\text{O}$  was added at  $80^\circ\text{C}$ . Last, 140 ml of  $\text{H}_2\text{O}$  and 10 ml of  $\text{H}_2\text{O}_2$  (30%, w/v) were introduced into the mixture to complete the reaction. The GO was washed and filtered with 1 M HCl. The self-supported h-Ni was cut into 2-mm-diameter circles using a  $\text{CO}_2$  laser cutter and cleaned by ultrasonication in 4 M HCl for 30 min until the color changed from black to silver. After drying, the h-Ni substrates were immersed into a GO suspension with a concentration of  $2.0 \text{ mg ml}^{-1}$  in water for 1 hour. Then, the h-Ni substrates were transferred to 5 ml of ascorbic acid ( $10 \text{ mg ml}^{-1}$ ) overnight and heated at  $75^\circ\text{C}$  for 2 hours. After cooling down to room temperature, the rGO/h-Ni composite electrodes were rinsed with water. The free-standing CNTs were immersed into a 2 mM MDB solution and then heated to  $140^\circ\text{C}$  overnight, followed by rinsing with water for several times. The resulted MDB-CNTs were dropcasted onto the rGO/h-Ni electrode to achieve a higher ECSA. The MDB-CNTs/rGO/h-Ni composite was soaked into the 20 mM TTF ethanol/acetone (9:1, v/v) solution. Then, bioanodes were obtained by immersing TTF-MDB-CNTs/rGO/h-Ni composite into an LOx solution ( $20 \text{ mg ml}^{-1}$ ) for 2 hours and dried at  $4^\circ\text{C}$ . Two microliters of 0.5% Nafion perfluorinated resin solution was dropcasted on the LOx/TTF-MDB-CNTs/rGO/h-Ni bioanodes to protect the enzymes during the operation.

The BFC cathodes were prepared as follows: The CNT film was first laser cut into 2-mm-diameter disks. The CNT pieces were immersed into 2 mM MDB solution, heated to  $140^\circ\text{C}$  overnight, and then rinsed with water for several times; the MDB-CNT pieces were immersed in a 60 mM  $\text{H}_2\text{PtCl}_6$  solution with 20 mM doping metal ions (like Co, Ni, Cu, and Zn) and then immersed in a 0.1 M  $\text{NaBH}_4$  solution for seconds, followed by several times of water rinsing; 2  $\mu\text{l}$  of 0.5% Nafion perfluorinated resin solution was dropcasted onto the Pt- or Pt alloy-decorated MDB-CNT composite surfaces.

### Characterizations of the BFCs

For characterization, the scanning electron microscopy (SEM) images of the electrodes were obtained by a field-emission SEM (FEI Sirion). High-resolution TEM images were obtained by a TEM (Tecnai TF-20). CV and linear sweep voltammetry (LSV) analyses were performed through an electrochemical workstation (CHI 660E). The characterizations of BFCs were performed in McIlvaine buffer solutions (pH 6.0) unless otherwise noted. Ag/AgCl references electrodes were used in the BFC characterization. The Pt electrode was fabricated by photolithography (Microchemicals GmbH, AZ 9260) and electron beam (E-beam) evaporation of Cr/Pt (20/100 nm, at a speed of 0.2 and 0.5  $\text{\AA/s}$ , respectively), followed by lift-off in acetone.

The commercial Pt/C electrode was fabricated by drop-casting 5  $\mu\text{l}$  of Pt/C solution ( $2 \text{ mg ml}^{-1}$ ) and 2.5  $\mu\text{l}$  of Nafion (0.5%) on the Au electrode (3 mm in diameter). To quantify the ECSA of the bioanode, we tested the electrodes in McIlvaine buffer (pH 6.0) with different scan rates ranging from 10 to 200  $\text{mV s}^{-1}$ . The slope of the calibration curve (the current density at  $-0.1 \text{ V}$  versus scan rate) was used as an estimate of the ECSA of each anode. The adsorption/desorption of hydrogen on Pt (here we used integral area in the CV in the range of  $-0.2$  to  $0.1 \text{ V}$ ) was used to characterize the ECSA of the Pt-based BFC cathode. The polarization curves and power output were achieved by the LSV measurement at a scan rate of  $2 \text{ mV s}^{-1}$ .

### Fabrication of the soft PPES

The fabrication process of the PPES platform is illustrated in figs. S11 to S13. PI (PI-2611) was spin-coated on the silicon handling wafer with a speed of 2000 rpm for 30 s. Then, the PI was cured at  $350^\circ\text{C}$  for 1 hour with a ramping speed of  $4^\circ\text{C min}^{-1}$ . The resulted PI substrate thickness was about 9  $\mu\text{m}$ . Photolithography (Microchemicals GmbH, AZ 9260) was used to define the inner connection wires. The photoresist was spin-coated on the wafer at a speed of 2400 rpm for 30 s and measured to be around 10  $\mu\text{m}$  thick. For surface treatment, reactive ion etching (RIE) [Oxford Plasmalab, 100 inductively coupled plasma (ICP)/RIE, 80 standard cubic centimeter per minute (sccm) of  $\text{O}_2$ , 5 sccm of SF<sub>6</sub>, 70 W, 20 mtorr] was used for 2 min to enhance surface adhesion of PI layers. E-beam evaporation of Cu (1.5  $\mu\text{m}$  at a speed of  $2.5 \text{ \AA s}^{-1}$ ) was deposited on the PI, followed by lift-off in acetone for minutes. An insulating layer of PI (PI-2610) was spin-coated on the surface with a speed of 5000 rpm for 30 s and then cured at  $350^\circ\text{C}$  for 30 min with a ramping speed of  $4^\circ\text{C min}^{-1}$ . The resulted intermediate PI layer thickness was about 1  $\mu\text{m}$ . Another photolithography step was used to define via connections between Cu layers. The wafer was selectively dry-etched using ICP (Oxford Plasmalab, 100 ICP/RIE, 50 sccm of  $\text{O}_2$ , 150 W, 80 mtorr, 9 min) to form via pattern. Photolithography was used to define outer connection wires. The wafer was surface-treated with RIE using the same recipe described above before metal evaporation. E-beam evaporation of Cu (2.5  $\mu\text{m}$  at a speed of  $2.5 \text{ \AA s}^{-1}$ ) was performed, followed by lift-off in acetone. Another encapsulation layer of PI (1  $\mu\text{m}$  thick; PI-2610) was spin-coated, followed by fully curing. Photolithography was performed to define openings of sensors and BFC patterns, and then dry etching was performed using ICP (Oxford Plasmalab, 100 ICP/RIE, 50 sccm of  $\text{O}_2$ , 150 W, 80 mtorr, 9 min).

After wiring system patterning, PI was spin-coated on the silicon handling wafer with a thickness of 9  $\mu\text{m}$ . Photolithography (Microchemicals GmbH, AZ 9260) was used to define the shapes of BFCs and sensor arrays. The PI was then surface-treated with RIE to enhance surface adhesion (Oxford Plasmalab, 100 ICP/RIE, 80 sccm of  $\text{O}_2$ , 5 sccm of SF<sub>6</sub>, 70 W, 20 mtorr). E-beam evaporation of Cr/Au (20/100 nm at a speed of 0.2 and 0.5  $\text{\AA/s}$ , respectively) was performed, followed by lift-off in acetone. A thin layer of parylene (ParaTech LabTop 3000 Parylene coater) was deposited (1  $\mu\text{m}$ ), followed by photolithography and RIE (Oxford Plasmalab, 100 ICP/RIE, 30 sccm of  $\text{O}_2$ , 100 W, 50 mtorr, 3 min) to expose openings for further treatments.

### Assembly and characterization of the PPES

The packaging and assembly of the PPES are illustrated in fig. S14. First, the electronic system pattern was set on 100  $\mu\text{m}$  of PDMS and connected with the BFC and sensor array pattern by the conductive silver paint (Structure Probe Inc.). One-hundred micrometers of PDMS

was coated on the BFC and electronic system patterns. The bottom layer was cut as the BFC and sensor array pattern to expose the electrode area. After electrodes were modified on the BFC and sensor array, it was combined with the laser-cut microfluidic channels, which were assembled with two medical tape layers and a single PDMS layer in the middle.

The core of the sensor sampling and data-processing transmission system is the CYBLE-214009-00 BLE module. This microcontroller provided onboard BLE capability and 12-bit analog-to-digital converter (ADC) resolution and a minimal power consumption of  $1.3\text{ }\mu\text{A}$  in deep sleep. The OCP of BFCs could hardly reach 1 V, which was significantly lower than the regular electronics needed. Thus, a soft integrated electronic patch, which contained energy boost converter for increasing the applied voltage, multiplexed sensing channels, and Bluetooth broadcast, was combined with the wearable BFCs. The Texas Instruments BQ25504 boost converter formed the core of the circuitry used to harvest energy from the BFCs. The BQ25504 was set to draw current from the BFC at 75% of its OCP to maximize the power provided by the cell. The output of the BQ25504 was connected to a  $680\text{-}\mu\text{F}$  capacitor, which was set to charge to 3.3 V before triggering an analog switch to close, connecting the charged capacitor to the rest of the circuitry and powering the system. Then, the instrumentation amplifier and microcontroller recorded the response of sensor arrays and sent it to the Bluetooth module to broadcast, which were all powered under the boost voltage. The module would run in bursts of activity, periodically waking up to take ADC measurements of the sensors, convert the measurement into a voltage, and then wirelessly broadcast the data via BLE advertisement before re-entering deep sleep mode. The device was programmed through the Cypress programmable system-on-chip interface using the MiniProg3 SWD debugger and PSoC Creator 4.2 design environment. To reduce noise and eliminate the effects of impedance mismatching, instrumentation amplifiers were connected as intermediaries between the sensor electrode output and the microcontroller ADC input. The Texas Instruments INA333 was used as the instrumentation amplifier. The BLE packets broadcasted by the microcontroller were received on a computer with a CY5677 CySmart BLE 4.2 Universal Serial Bus (USB) dongle connected. The computer ran a scanner program written in C# with the CySmart 1.3 application programming interface (API), which continually searched for an advertising device that matched the BLE address of the Cyble module, and then decoded and stored any received data.

### Preparation and characterization of biosensors

To prepare the shared reference electrode of the potentiometric sensor array, we first electrodeposited Ag on the Au electrodes with a potentiostatic method ( $-0.25\text{ V}$  for 600 s) in the solution containing  $0.25\text{ M }$   $\text{AgNO}_3$ ,  $0.75\text{ M }$   $\text{Na}_2\text{S}_2\text{O}_3$ , and  $0.43\text{ M }$   $\text{NaHSO}_3$ ; the Ag/AgCl electrode was obtained by dropping the  $0.1\text{ M }$   $\text{FeCl}_3$  solution on top of the Ag surface for 60 s, and then the PVB reference cocktail was prepared by dissolving 79.1 mg of PVB, 50 mg of NaCl, 1 mg of F127, and 0.2 mg of multiwalled CNT in 1 ml of methanol; a  $6.6\text{ }\mu\text{l}$  of reference cocktail was modified on the Ag/AgCl electrode and left overnight.

The  $\text{NH}_4^+$  selective electrode was prepared as follows: 100 mg of the  $\text{NH}_4^+$  selective membrane cocktail consisting of 1%  $\text{NH}_4^+$  ionophore (nonactin), 33% PVC, and 66% DOS (w/w) was dissolved in  $660\text{ }\mu\text{l}$  of tetrahydrofuran. The membrane cocktail was stored at  $4^\circ\text{C}$ . A constant current of  $0.2\text{ mA cm}^{-2}$  was applied to electrodeposit the poly(EDOT) polystyrene sulfonate (PEDOT:PSS) membrane on the Au electrode in the solution containing  $0.01\text{ M }$  EDOT and  $0.1\text{ M }$

NaPSS to minimize the potential drift of the ISEs. Cocktail solution ( $6.6\text{ }\mu\text{l}$ ) was dropcasted over the PEDOT layer to create an  $\text{NH}_4^+$  selective membrane. The modified electrodes were left drying overnight. To prepare urea-sensing electrode, we dropcasted  $3.7\text{ }\mu\text{l}$  of urease solution ( $10\text{ mg ml}^{-1}$ ) onto the  $\text{NH}_4^+$  ISE four times, and then  $3.3\text{ }\mu\text{l}$  of 0.5% Nafion perfluorinated resin solution was dropped over the sensor area. The modified sensors were dried at  $4^\circ\text{C}$  overnight.

To prepare the glucose sensor, we used differential pulse amperometry (100 cycles in total) to electrodeposit the Pt on the Au electrode. A  $-0.4\text{-V}$  potential was applied for 1 s, and  $1.0\text{ V}$  was used as the cleaning voltage for 0.5 s. A 1% Nafion ( $1.1\text{ }\mu\text{l}$ ; prepared by dilution of Nafion perfluorinated resin solution in water) was dropped on the Pt surface, and  $2\text{ }\mu\text{l}$  of CS-GOx mixture (3:1, v/v) was modified on the electrode. The potentiometric glucose sensors were dried at  $4^\circ\text{C}$  overnight, and then another  $1.1\text{ }\mu\text{l}$  of 1% Nafion (1%) was dropped, covering the enzymes to form the sandwich structure. For the pH sensor, the polyaniline was electropolymerized on the Au electrodes in a solution containing  $0.1\text{ M }$  aniline and  $0.1\text{ M }$  HCl using CV from  $-0.2$  to  $1\text{ V}$  for 50 cycles at a scan rate of  $50\text{ mV s}^{-1}$ .

For the in vitro sensor characterizations, analyte solutions were prepared in McIlvaine buffer solutions (pH 6.0 for urea, glucose, and  $\text{NH}_4^+$ ). To obtain the best performance for long-term continuous measurements, we placed the  $\text{NH}_4^+$  ion-selective sensors and urea sensors in a solution containing  $0.1\text{ M }$   $\text{NH}_4\text{Cl}$  and  $20\text{ mM}$  urea for 1 hour before measurements. The glucose and pH sensors were placed in a solution containing  $100\text{ }\mu\text{M}$  glucose and  $\text{H}_2\text{O}$  for 1 hours, separately. This conditioning process greatly helps to minimize the potential drift.

### Preparation of the strain sensors

CNT-PDMS elastomer was produced by the solution-evaporation method. CNTs (7% w/w) were added to SYLGARD 184 silicone elastomer base and toluene mixture (1:4 v/v) at room temperature. Then, the mixture was poured into a culture dish. After toluene was evaporated, uncured CNT-PDMS was mixed with curing agent (10:1) and poured onto the mask made by tape and scraped flat with a glass slide. After the mask was removed, the CNT-PDMS was baked at  $80^\circ\text{C}$  for 1 hour. Then, uncured EcoFlex was spin-coated on it and cured at  $80^\circ\text{C}$  for 1 hour. The silver paste was used to link the pad on the patterned CNT-PDMS with thin wires. Another layer of EcoFlex was spin-coated for encapsulation and protection. The strain sensors were then connected to the PPES through external wires.

### On-body evaluation of the PPES for wearable sensing

The validation and evaluation of the lab-on-skin platform were performed using humans in compliance with a protocol (ID: 19-0892) that was approved by the Institutional Review Board (IRB) at California Institute of Technology (Caltech). The participating individuals (age range, 18 to 65) were recruited from the Caltech campus and the neighboring communities through advertisement by posted notices, word of mouth, and email distribution. All individuals gave written informed consent before participation in the study.

To validate the PPES, constant-load cycling exercise was conducted on three individuals. A stationary exercise bike (Kettler Axos Cycle M-LA) was used for cycling trials. The individuals reported to the laboratory with overnight fasting. The individuals' foreheads were cleaned with water and alcohol swabs before the PPES was placed. The individuals cycled at 60 rpm for 40 min, followed by a 10-min cooldown session. When the sweat filled the BFC reservoir, the PPES

started to transmit the sensing data to the user interface through the Bluetooth every 15 s, and the data were further converted to the concentration levels after temperature and sweat analyte calibrations. During biking, the sweat would continuously refill both the BFC and the sensor array reservoir. For each diary challenge study, after the first cycling trial, the individuals were given a standardized protein drink (30 g of protein) for urea and  $\text{NH}_4^+$  test and three energy bars (66 g of total sugars) for glucose, respectively. After 2 hours, the individuals cycled at 60 rpm for 40 min, followed by a 10-min cooldown session, and the PPES would monitor the sweat component change. The first set of data received by the PPES was plotted in Fig. 6 (D to I).

### Evaluation of the PPES as a human-machine interface for robotic assistance

For robotic arm control, two strain sensors were set on the hand and the elbow of a healthy individual with medical tape, respectively. The individual's arm was cleaned with water and alcohol swabs before the PPES was placed. The individual cycled at 60 rpm for 10 min. When the sweat filled the BFC reservoir, the PPES wirelessly transmitted the strain sensor data to the user interface, and the data were further used to control the robotic arm.

For the PPES-based prosthesis control study, a strain sensor was set on the elbow of a healthy individual with medical tape and connected with the PPES. When filled with 20 mM lactate, the PPES wirelessly transmitted the measured strain sensor data to the user interface. When the sensor reached a certain threshold, a human operator sent a computer command through USB to the custom-built powered transfemoral prosthesis, AMPRO3 (49), to trigger the start of one swing step. With a proportional derivative (PD) controller, the prosthesis tracked a time-based trajectory designed with the methods in (50). This research was approved by Caltech IRB with a protocol (ID: 16-0693) for human subject testing.

### SUPPLEMENTARY MATERIALS

[robotics.sciencemag.org/cgi/content/full/5/41/eaaz7946/DC1](http://robotics.sciencemag.org/cgi/content/full/5/41/eaaz7946/DC1)

Fig. S1. CVs of an MDB-CNT/rGO/h-Ni electrode.

Fig. S2. Characterization of the surface area of the BFC anodes.

Fig. S3. Role of MDB in the preparation of the Pt/CNT BFC cathodes.

Fig. S4. Characterization of the Pt/CNT and the Pt/MDB-CNT cathodes.

Fig. S5. Characterization of the bulk Pt, Pt/MDB-CNT, and Pt alloy/MDB-CNT cathodes.

Fig. S6. LSV characterization of the bioanode and the cathodes in lactate solutions.

Fig. S7. Characterization of the stability of the Pt/MDB-CNT cathode.

Fig. S8. Characterization of the Pt-Co alloy nanoparticles.

Fig. S9. Schematic diagram of the energy-harvesting circuit.

Fig. S10. Schematic illustration and component list of the electronic system of the PPES.

Fig. S11. Fabrication process of the electrochemical patch of the PPES.

Fig. S12. Exploded view of the multilayered electronic patch.

Fig. S13. Fabrication process of the flexible electronic patch of the PPES.

Fig. S14. Assembly and encapsulation of the PPES.

Fig. S15. Mechanical deformation study of the BFC patch.

Fig. S16. Mechanical deformation study of the electronic patch.

Fig. S17. Power consumption breakdown of the PPES.

Fig. S18. Continuous operation of the PPES with small capacitors in 5 to 20 mM lactate.

Fig. S19. Long-term stability of the BFCs to charge a capacitor.

Fig. S20. Selectivity of the sensor arrays.

Fig. S21. Interactive study of the urea/ $\text{NH}_4^+$  sensor array.

Fig. S22. Interactive study of the glucose/pH sensor array.

Fig. S23. Stability of the sensor arrays.

Fig. S24. Performance of the sensor arrays under mechanical deformation.

Fig. S25. Influence of the temperature on the performance of enzymatic sensors.

Fig. S26. Skin-interfaced microfluidic patch in the PPES.

Fig. S27. Investigation of the stability, reusability, and biocompatibility of the PPES.

Fig. S28. Operation flow of data transmission processes and human-machine interaction.

Fig. S29. Schematic illustration and component list of the strain sensor integrated PPES for human-machine interaction.

Text S1. Mechanism of TTF-MDB bimediator in BFC.

Table S1. List of BFCs for energy-harvesting in body fluids.

Movie S1. Biofuel-powered electronic skin for robotic arm control and cargo handling.

Movie S2. Biofuel-powered electronic skin for prosthesis control.

References (51–55)

### REFERENCES AND NOTES

1. G.-Z. Yang, J. Bellingham, P. E. Dupont, P. Fischer, L. Floridi, R. Full, N. Jacobstein, V. Kumar, M. McNutt, R. Merrifield, B. J. Nelson, B. Scassellati, M. Taddeo, R. Taylor, M. Veloso, Z. L. Wang, R. Wood, The grand challenges of *Science Robotics*. *Sci. Robot.* **3**, eaar7650 (2018).
2. T. Someya, M. Amagai, Toward a new generation of smart skins. *Nat. Biotechnol.* **37**, 382–388 (2019).
3. S. Xu, Y. Zhang, L. Jia, K. E. Mathewson, K.-I. Jang, J. Kim, H. Fu, X. Huang, P. Chava, R. Wang, S. Bhole, L. Wang, Y. J. Na, Y. Guan, M. Flavin, Z. Han, Y. Huang, J. A. Rogers, Soft microfluidic assemblies of sensors, circuits, and radios for the skin. *Science* **344**, 70–74 (2014).
4. S. I. Rich, R. J. Wood, C. Majidi, Untethered soft robotics. *Nat. Electron.* **1**, 102–112 (2018).
5. M. L. Hammock, A. Chortos, B. C.-K. Tee, J. B.-H. Tok, Z. Bao, 25th anniversary article: The evolution of electronic skin (e-skin): A brief history, design considerations, and recent progress. *Adv. Mater.* **25**, 5997–6038 (2013).
6. C. Laschi, B. Mazzolai, M. Cianchetti, Soft robotics: Technologies and systems pushing the boundaries of robot abilities. *Sci. Robot.* **1**, eah3690 (2016).
7. A. Chortos, J. Liu, Z. Bao, Pursuing prosthetic electronic skin. *Nat. Mater.* **15**, 937–950 (2016).
8. S. Gong, W. Schwalb, Y. Wang, Y. Chen, Y. Tang, J. Si, B. Shirinzadeh, W. Cheng, A wearable and highly sensitive pressure sensor with ultrathin gold nanowires. *Nat. Commun.* **5**, 3132–3140 (2014).
9. X. Wang, L. Dong, H. Zhang, R. Yu, C. Pan, Z. L. Wang, Recent progress in electronic skin. *Adv. Sci.* **2**, 1500169–1500190 (2015).
10. Y. Liu, J. Liu, S. Chen, T. Lei, Y. Kim, S. Niu, H. Wang, X. Wang, A. M. Foudeh, J. B. Tok, Z. Bao, Soft and elastic hydrogel-based microelectronics for localized low-voltage neuromodulation. *Nat. Biomed. Eng.* **3**, 58–68 (2019).
11. D. Son, J. Lee, S. Qiao, R. Ghaffari, J. Kim, J. E. Lee, C. Song, S. J. Kim, D. J. Lee, S. W. Jun, S. Yang, M. Park, J. Shin, K. Do, M. Lee, K. Kang, C. S. Hwang, N. Lu, T. Hyeon, D. H. Kim, Multifunctional wearable devices for diagnosis and therapy of movement disorders. *Nat. Nanotechnol.* **9**, 397–404 (2014).
12. W. Gao, S. Emaminejad, H. Y. Y. Nyein, S. Challa, K. Chen, A. Peck, H. M. Fahad, H. Ota, H. Shiraki, D. Kiriyama, D.-H. Lien, G. A. Brooks, R. W. Davis, A. Javey, Fully integrated wearable sensor arrays for multiplexed in situ perspiration analysis. *Nature* **529**, 509–514 (2016).
13. M. Baria, H. Y. Y. Nyein, A. Javey, Wearable sweat sensors. *Nat. Electron.* **1**, 160–171 (2018).
14. J. Kim, A. S. Campbell, B. E.-F. de Avila, J. Wang, Wearable biosensors for healthcare monitoring. *Nat. Biotechnol.* **37**, 389–406 (2019).
15. Y. Yang, W. Gao, Wearable and flexible electronics for continuous molecular monitoring. *Chem. Soc. Rev.* **48**, 1465–1491 (2019).
16. S. Nakata, M. Shiomi, Y. Fujita, T. Arie, S. Akita, K. Takei, A wearable pH sensor with high sensitivity based on a flexible charge-coupled device. *Nat. Electron.* **1**, 596–603 (2018).
17. S. Park, S. W. Heo, W. Lee, D. Inoue, Z. Jiang, K. Yu, H. Jinno, D. Hashizume, M. Sekino, T. Yokota, K. Fukuda, K. Tajima, T. Someya, Self-powered ultra-flexible electronics via nano-grating-patterned organic photovoltaics. *Nature* **561**, 516–521 (2018).
18. W. Zeng, L. Shu, Q. Li, S. Chen, F. Wang, X.-M. Tao, Fiber-based wearable electronics: A review of materials, fabrication, devices, and applications. *Adv. Mater.* **26**, 5310–5336 (2014).
19. A. M. Zamarayeva, A. E. Ostfeld, M. Wang, J. K. Duey, I. Deckman, B. P. Lechêne, G. Davies, D. A. Steingart, A. C. Arias, Flexible and stretchable power sources for wearable electronics. *Sci. Adv.* **3**, e1602051 (2017).
20. J. Li, J. Zhao, J. A. Rogers, Materials and designs for power supply systems in skin-interfaced electronics. *Acc. Chem. Res.* **52**, 53–62 (2019).
21. J. Chen, Y. Huang, N. Zhang, H. Zou, R. Liu, C. Tao, X. Fan, Z. L. Wang, Micro-cable structured textile for simultaneously harvesting solar and mechanical energy. *Nat. Energy* **1**, 16138–16146 (2016).
22. F. Suarez, A. Nozariasbmaz, D. Vashaei, M. C. Öztürk, Designing thermoelectric generators for self-powered wearable electronics. *Energ. Environ. Sci.* **9**, 2099–2113 (2016).
23. Z. Lou, L. Li, L. Wang, G. Shen, Recent progress of self-powered sensing systems for wearable electronics. *Small* **13**, 1701791–1701818 (2017).

24. A. J. Bandodkar, P. Gutruf, J. Choi, K. Lee, Y. Sekine, J. T. Reeder, W. J. Jeang, A. J. Aranyosi, S. P. Lee, J. B. Model, R. Ghaffari, C.-J. Su, J. P. Leshock, T. Ray, A. Verrillo, K. Thomas, V. Krishnamurthi, S. Han, J. Kim, S. Krishnan, T. Hang, J. A. Rogers, Battery-free, skin-interfaced microfluidic/electronic systems for simultaneous electrochemical, colorimetric, and volumetric analysis of sweat. *Sci. Adv.* **5**, eaav3294 (2019).
25. R. A. Escalona-Villalpando, E. Ortiz-Ortega, J. P. Bocanegra-Ugalde, S. D. Minteer, J. Ledesma-García, L. G. Arriaga, Clean energy from human sweat using an enzymatic patch. *J. Power Sources* **412**, 496–504 (2019).
26. A. J. Bandodkar, J.-M. You, N.-H. Kim, Y. Gu, R. Kumar, A. M. V. Mohan, J. Kurniawan, S. Imani, T. Nakagawa, B. Parish, M. Parthasarathy, P. P. Mercier, S. Xu, J. Wang, Soft, stretchable, high power density electronic skin-based biofuel cells for scavenging energy from human sweat. *Energ. Environ. Sci.* **10**, 1581–1589 (2017).
27. H. Guo, X. Pu, J. Chen, Y. Meng, M.-H. Yeh, G. Liu, Q. Tang, B. Chen, D. Liu, S. Qi, C. Wu, C. Hu, J. Wang, Z. L. Wang, A highly sensitive, self-powered triboelectric auditory sensor for social robotics and hearing aids. *Sci. Robot.* **3**, eaat2516 (2018).
28. J. Heikenfeld, A. Jajack, B. Feldman, S. W. Granger, S. Gaitonde, G. Begtrup, B. A. Katchman, Assessing analytes in biofluids for peripheral biochemical monitoring. *Nat. Biotechnol.* **37**, 407–419 (2019).
29. K. Suhre, S.-Y. Shin, A.-K. Petersen, R. P. Mohney, D. Meredith, B. Wägele, E. Altmaier; CARDIOGRAM, P. Deloukas, J. Erdmann, E. Grundberg, C. J. Hammond, M. H. de Angelis, G. Kastenmüller, A. Köttgen, F. Kronenberg, M. Mangino, C. Meisinger, T. Meitinger, H.-W. Mewes, M. V. Milburn, C. Prehn, J. Raffler, J. S. Ried, W. Römisich-Margl, N. J. Samani, K. S. Small, H. E. Wichmann, G. Zhai, T. Illig, T. D. Spector, J. Adamski, N. Soranzo, C. Gieger, Human metabolic individuality in biomedical and pharmaceutical research. *Nature* **477**, 54–60 (2011).
30. A. Koh, D. Kang, Y. Xue, S. Lee, R. M. Pielak, J. Kim, T. Hwang, S. Min, A. Banks, P. Bastien, M. C. Manco, L. Wang, K. R. Ammann, K.-I. Jang, P. Won, S. Han, R. Ghaffari, U. Paik, M. J. Slepian, G. Balooch, Y. Huang, J. A. Rogers, A soft, wearable microfluidic device for the capture, storage, and colorimetric sensing of sweat. *Sci. Transl. Med.* **8**, 366ra165 (2016).
31. B. Meng, W. Tang, Z.-h. Too, X. Zhang, M. Han, W. Liu, H. Zhang, A transparent single-friction-surface triboelectric generator and self-powered touch sensor. *Energ. Environ. Sci.* **6**, 3235–3240 (2013).
32. H. Lee, T. K. Choi, Y. B. Lee, H. R. Cho, R. Ghaffari, L. Wang, H. J. Choi, T. D. Chung, N. Lu, T. Hyeon, S. H. Choi, D. H. Kim, A graphene-based electrochemical device with thermoresponsive microneedles for diabetes monitoring and therapy. *Nat. Nanotechnol.* **11**, 566–572 (2016).
33. X. Xiao, H.-q. Xia, R. Wu, L. Bai, L. Yan, E. Magner, S. Cosnier, E. Lojou, Z. Zhu, A. Liu, Tackling the challenges of enzymatic (bio)fuel cells. *Chem. Rev.* **119**, 9509–9558 (2019).
34. W. Jia, G. Valdés-Ramirez, A. J. Bandodkar, J. R. Windmiller, J. Wang, Epidermal biofuel cells: Energy harvesting from human perspiration. *Angew. Chem. Int. Ed.* **52**, 7233–7236 (2013).
35. G. A. Dienel, L. Hertz, Glucose and lactate metabolism during brain activation. *J. Neurosci. Res.* **66**, 824–838 (2001).
36. L. F. Barros, Metabolic signaling by lactate in the brain. *Trends Neurosci.* **36**, 396–404 (2013).
37. I. T. McCrum, M. J. Janik, Deconvoluting cyclic voltammograms to accurately calculate Pt electrochemically active surface area. *J. Phys. Chem. C* **121**, 6237–6245 (2017).
38. C. U. Ferreira, Y. Gushikem, L. T. Kubota, Electrochemical properties of Meldola's Blue immobilized on silica-titania phosphate prepared by the sol-gel method. *J. Solid State Electrochem.* **4**, 298–303 (2000).
39. X. Xie, M. Ye, P.-C. Hsu, N. Liu, C. S. Criddle, Y. Cui, Microbial battery for efficient energy recovery. *Proc. Natl. Acad. Sci. U.S.A.* **110**, 15925–15930 (2013).
40. L. Bai, L. Jin, L. Han, S. Dong, Self-powered fluorescence controlled switch systems based on biofuel cells. *Energ. Environ. Sci.* **6**, 3015–3021 (2013).
41. W. Vogel, L. Lundquist, P. Ross, P. Stonehart, Reaction pathways and poisons—II. *Electrochim. Acta* **20**, 79–93 (1975).
42. I. E. L. Stephens, A. S. Bondarenko, U. Grønbjerg, J. Rossmeisl, I. Chorkendorff, Understanding the electrocatalysis of oxygen reduction on platinum and its alloys. *Energ. Environ. Sci.* **5**, 6744–6762 (2012).
43. Y. Nie, L. Li, Z. Wei, Recent advancements in Pt and Pt-free catalysts for oxygen reduction reaction. *Chem. Soc. Rev.* **44**, 2168–2201 (2015).
44. J. Greeley, I. E. L. Stephens, A. S. Bondarenko, T. P. Johansson, H. A. Hansen, T. F. Jaramillo, J. Rossmeisl, I. Chorkendorff, J. K. Nørskov, Alloys of platinum and early transition metals as oxygen reduction electrocatalysts. *Nat. Chem.* **1**, 552–556 (2009).
45. B.-W. Zhang, H.-L. Yang, Y.-X. Wang, S.-X. Dou, H.-K. Liu, A comprehensive review on controlling surface composition of Pt-based bimetallic electrocatalysts. *Adv. Energy Mater.* **8**, 1703597–1703614 (2018).
46. V. R. Stamenkovic, B. S. Mun, M. Arenz, K. J. J. Mayrhofer, C. A. Lucas, G. Wang, P. N. Ross, N. M. Markovic, Trends in electrocatalysis on extended and nanoscale Pt-bimetallic alloy surfaces. *Nat. Mater.* **6**, 241–247 (2007).
47. Y. Yang, Y. Song, X. Bo, J. Min, O. S. Pak, L. Zhu, M. Wang, J. Tu, A. Kogan, H. Zhang, T. K. Hsiao, Z. Li, W. Gao, A laser-engraved wearable sensor for sensitive detection of uric acid and tyrosine in sweat. *Nat. Biotechnol.* **38**, 217–224 (2020).
48. W. S. Hummers Jr., R. E. Offeman, Preparation of graphitic oxide. *J. Am. Chem. Soc.* **80**, 1339–1339 (1958).
49. H. Zhao, J. Horn, J. Reher, V. Paredes, A. D. Ames, First steps toward translating robotic walking to prostheses: A nonlinear optimization based control approach. *Auton. Robot.* **41**, 725–742 (2017).
50. H. Zhao, E. Ambrose, A. D. Ames, Preliminary results on energy efficient 3D prosthetic walking with a powered compliant transfemoral prosthesis, in *IEEE Int. Conf. Robot. Autom. (ICRA)* (2017), pp. 1140–1147.
51. P. Ó. Conghaile, M. Falk, D. MacAodha, M. E. Yakovleva, C. Gonau, C. K. Peterbauer, L. Gorton, S. Shleev, D. Leech, Fully enzymatic membraneless glucose/oxygen fuel cell that provides  $0.275 \text{ mA cm}^{-2}$  in 5 mM glucose, operates in human physiological solutions, and powers transmission of sensing data. *Anal. Chem.* **88**, 2156–2163 (2016).
52. R. C. Reid, S. D. Minteer, B. K. Gale, Contact lens biofuel cell tested in a synthetic tear solution. *Biosens. Bioelectron.* **68**, 142–148 (2015).
53. M. Falk, D. Pankratov, L. Lindh, T. Arnebrant, S. Shleev, Miniature direct electron transfer based enzymatic fuel cell operating in human sweat and saliva. *Fuel Cells* **14**, 1050–1056 (2014).
54. M. Falk, V. Andorlov, Z. Blum, J. Sotres, D. B. Suyatin, T. Ruzgas, T. Arnebrant, S. Shleev, Biofuel cell as a power source for electronic contact lenses. *Biosens. Bioelectron.* **37**, 38–45 (2012).
55. W. Jia, X. Wang, S. Imani, A. J. Bandodkar, J. Ramírez, P. P. Mercier, J. Wang, Wearable textile biofuel cells for powering electronics. *J. Mater. Chem. A* **2**, 18184–18189 (2014).

**Funding:** This work was supported by California Institute of Technology Startup Grant, the Rothenberg Innovation Initiative ( $\text{RI}^2$ ) program, the Carver Mead New Adventures Fund, and NIH (no. 5R21NR018271) (all to W.G.). We gratefully acknowledge critical support and infrastructure provided for this work by the Kavli Nanoscience Institute at Caltech.

**Author contributions:** W.G. and Y.Y. conceived the project. W.G. supervised the studies and Y.Y. led the experiments. J.N., C.X., and Y.Y. contributed to the system development, fabrication, and characterization; J.M., A.D., and R.D. contributed to the electronic circuit design and test. A.H. and Y.S. contributed to sensor preparation and characterization. R.G. and A.D.A. contributed to the prosthesis control study. Y. Yu, Y. Yang, and W.G. contributed the data analysis and co-wrote the paper. All authors provided feedback on the manuscript.

**Competing interests:** The authors declare that they have no competing financial interest.

**Data and materials availability:** All data needed to evaluate the conclusions in the paper are present in the paper or the Supplementary Materials.

Submitted 11 October 2019

Accepted 18 March 2020

Published 22 April 2020

10.1126/scirobotics.aaz7946

**Citation:** Y. Yu, J. Nassar, C. Xu, J. Min, Y. Yang, A. Dai, R. Doshi, A. Huang, Y. Song, R. Gehlhar, A. D. Ames, W. Gao, Biofuel-powered soft electronic skin with multiplexed and wireless sensing for human-machine interfaces. *Sci. Robot.* **5**, eaaz7946 (2020).

# Science Robotics

## Biofuel-powered soft electronic skin with multiplexed and wireless sensing for human-machine interfaces

You Yu, Joanna Nassar, Changhao Xu, Jihong Min, Yiran Yang, Adam Dai, Rohan Doshi, Adrian Huang, Yu Song, Rachel Gehlhar, Aaron D. Ames and Wei Gao

*Sci. Robotics* **5**, eaaz7946.  
DOI: 10.1126/scirobotics.aaz7946

### ARTICLE TOOLS

<http://robotics.sciencemag.org/content/5/41/eaaz7946>

### SUPPLEMENTARY MATERIALS

<http://robotics.sciencemag.org/content/suppl/2020/04/20/5.41.eaaz7946.DC1>

### RELATED CONTENT

<http://robotics.sciencemag.org/content/robotics/4/32/eaax2352.full>

### REFERENCES

This article cites 54 articles, 5 of which you can access for free  
<http://robotics.sciencemag.org/content/5/41/eaaz7946#BIBL>

### PERMISSIONS

<http://www.sciencemag.org/help/reprints-and-permissions>

Use of this article is subject to the [Terms of Service](#)

---

*Science Robotics* (ISSN 2470-9476) is published by the American Association for the Advancement of Science, 1200 New York Avenue NW, Washington, DC 20005. The title *Science Robotics* is a registered trademark of AAAS.

Copyright © 2020 The Authors, some rights reserved; exclusive licensee American Association for the Advancement of Science. No claim to original U.S. Government Works