

## REVIEW

[View Article Online](#)  
[View Journal](#) | [View Issue](#)Cite this: *Mater. Adv.*, 2023,  
4, 1444Received 11th July 2022,  
Accepted 26th September 2022

DOI: 10.1039/d2ma00818a

[rsc.li/materials-advances](https://rsc.li/materials-advances)

## Wearable strain sensors: state-of-the-art and future applications

Ashish Yadav, <sup>a</sup> Neha Yadav, <sup>a</sup> Yongling Wu, <sup>a</sup> Seeram RamaKrishna <sup>b</sup> and Zheng Hongyu <sup>a</sup>

Wearable strain sensors have drawn massive awareness in various studies and industrial fields. The growing suitability and adaptability of such systems have increased the demand for developing instruments with improved aspects and packaging required to investigate bio-integrated gadgets, electronic skins, wearable wellness care techniques, and smooth robotics. Despite the outstanding performance, there are enormous challenges in the production of significant fundamental enrichments to the latest optical, mechanical, electrical, and sensing modalities and developing these for precise detection needs. Given all these demanding challenges, state-of-the-art wearable strain sensor technology is evolving towards a roadmap for exploring next-level generation innovations and breakthroughs in nanotechnology, and flexible strain sensors incorporate the virtues of stretchability, high detection power, and several other features. Skin-attachable wearable devices are developing as a tremendously attractive area due to their capable applications in the fields of artificial intelligence, human-machine systems, and healthcare tools. To understand and explore wearable strain sensors, this brief review describes aspects of flexible wearable strain sensing systems along with their challenges and prospects.

## 1. Introduction

Wearable sensing technology<sup>1–5</sup> has attracted significant attention and has swiftly transformed from a scientific idea to an extensive collection of traditional consumer and healthcare goods. This rapid evolution of wearable sensors may be credited to

<sup>a</sup> Center for Advanced Laser Manufacturing (CALM), Shandong University of Technology, Zibo, 255000, P. R. China. E-mail: [ashish@sdut.edu.cn](mailto:ashish@sdut.edu.cn), [zhenghongyu@sdut.edu.cn](mailto:zhenghongyu@sdut.edu.cn)

<sup>b</sup> Nanoscience and Nanotechnology Initiative, National University of Singapore, 10 Kent Ridge Crescent, 119260, Singapore



Ashish Yadav

Ashish Yadav, PhD Assc. Professor from Shandong University of Technology, China. He obtained a master's degree in applied physics from India. He got a PhD degree from the department of Fisica from the University of Roma Tor Vergata, Italy in 2014. His work focuses on advanced self-assembly-based photonic crystals and plasmonic nanostructures. Presently he is working as an Associate Professor on fabrication methods and the develop-

ment of three-dimensionally ordered high-quality photonic crystal films and their application as optical materials for lasing and devices. He is working multi directions research e.g. wearable electronics, sensors, and energy harvesting devices also interested in laser spectroscopic techniques for low-threshold devices.



Neha Yadav

Dr Neha Yadav obtained her bachelor's degree in science and Master's degree in Computer application. Now she is working on smart materials and Artificial Intelligence.



numerous features, *e.g.*, comparatively low-cost prices and ergonomics implemented by advancements in the small sizes of electronic components, the vast demand and use of smart-phones and associated gadgets, a rising human yearning for health-related awareness, and the unachieved medical requirements to continuously obtain correct and precise medical quality data from the patients.<sup>6–9</sup> Despite the substantial preliminary success, a large amount of knowledge is still required to monitor exact information from the body. These requirements

are still only partially met by the detecting modalities used in contemporary wearables,<sup>10–12</sup> which are not specific. Besides, most wearable sensor products are still being manufactured based on outdated techniques, which have been accessible for decades. Even the most innovative wearables, such as continuous transdermal glucose monitoring equipment, leverage almost 30 years of advances in enzyme electrodes using simple and ultra-cheap finger-prick test strips of glucose.<sup>13</sup> Transcutaneous blood glucose monitoring may be the only prevalent wearable sensor that measures the persistent observation of underlying explicit disease or infection.<sup>13–16</sup>

Technology has been playing larger roles in our daily lives, which helps to meet some of the demands of modern living while also adding to others. To fulfill these needs, there are numerous opportunities in the textile business to increase the usefulness and performance of textiles. The emergence of intelligent nanotextiles<sup>17</sup> will transform how we dress, how our houses are furnished, and how industrial materials are utilised. Expectations for textile performance have increased due to the impending revolution, and there is a high demand for “smart materials” that are more aware of their surroundings. A wide variety of uses for technical and functional textiles can be found in fields ranging from the military and security to individualized healthcare, cleanliness, and entertainment.<sup>18,19</sup>

Doctors can diagnose the disease and monitor the wellness of the patients by using numerous outdated available diagnostic tools for nearly every analyte. Nevertheless, these tests require the drawing of blood from a patient's body because of their non-wearability and outdated bench-top assessment procedures.<sup>20–25</sup> The main question of whether wearable sensor technology can evolve for the precise and more specific



**Yongling WU**

*Professor WU is a distinguished professor in Shandong University of Technology (SDUT). She gained her B.Eng and M.Eng in Mechanical Engineering from Tsinghua University, China, and PhD in Materials Science and Engineering from Nanyang Technological University of Singapore. Prior to joining SDUT, she was a Senior Scientist in the Agency for Science, Technology and Research (A-STAR) of Singapore leading a research group in Surface Technology. Prof. WU won the National Technology Award - Singapore's highest technology award. Her current research interests include: nanomaterials and coatings, laser materials processing, surface science and engineering, functional materials and surfaces.*



**Seeram Ramakrishna**

*Professor Seeram Ramakrishna, FREng, Everest Chair is a world-renowned poly-disciplinary scholar at the National University of Singapore, NUS. He is an elected Fellow of UK Royal Academy of Engineering (FREng), American Association of Advancement of Science, AAAS, Singapore Academy of Engineering, and Indian National Academy of Engineering. He is also an elected Fellow of ASM International, ASME, and AIMBE (USA); IMechE and IoM3 (UK); ISTE (India); International Union of Biomaterials Science & Engineering (FBSE). His publications to date have received 177 H-index and 153,245 citations. He is named among the World's Most Influential Minds (Thomson Reuters) and the Top 1% Highly Cited Researchers in cross-field (Clarivate Analytics). Stanford University C-score ranks him among top six impactful researchers of the world in materials, biomedical engineering, and enabling & strategic technologies. He is the Director of Center for Nanotechnology and Sustainability.*



**Zheng Hongyu**

*Professor Zheng Hongyu is a distinguished professor in Shandong University of Technology (SDUT), and director of Centre for Advanced Laser Manufacturing (CALM). He gained his B.Eng in Mechanical Engineering from Tsinghua University in 1985, and Ph.D in laser materials processing from Imperial College London in 1990. Prior to joining SDUT, he was a principal scientist with the Agency for Science, Technology and Research (A-STAR) of Singapore. He is Fellow of the International Society of Nano-Manufacturing (ISNM), and has co-authored more than 300 scientific publications. His current research interests include ultrafast laser-matter interactions, two-photon polymerization, micro-/nano-processing mechanisms, in-process monitoring and data analytics.*



physiological phenomena, such as monitoring the health of a baby inside the mother's womb by the measurement of simple mechanical movements of the fetus, or the differentiation of a fatal seizure from just higher physical exertion,<sup>26–29</sup> or to warn a sports person or a worker about their dangerous dehydration because of over-exercising or overwork by continuous measurement of their health analysis data, still needs answers.<sup>9,20,30</sup>

Usually, wearable electronic gadgets are fabricated using wearable non-toxic materials, smart sensors, modules and linkages, actuators, power sources, control and processing units, a user interface, software, and sophisticated algorithms for data collecting and decision-making, which are all components of wireless communication systems. As a result, the system processes data from factors including body temperature, blood pressure, the patient's level of exertion, and the concentrations of gases, different ions, and proteins in circulation to monitor the patient's physiological information. The smart sensors will be utilized in wearable electronic devices, which are surrounded by conducting electrodes. However, these constituent materials must be lightweight, highly flexible, ultrathin, stretchable, and with low moduli. Various state-of-the-art applications of wearable strain sensors are illustrated in Fig. 1.

In this brief review, our focus will be on wearable technologies that can extract data from the interior of the body, but without the need to implant a sensor within the body. Typical environmental sensors and limb-movement accelerometers are not in the scope of this review. This report focuses on the new borderline of wearable techniques and terms traditionally utilized by analytical chemists. In brief, this report describes the categorization of wearable strain sensors,<sup>31–33</sup> the basic physics principles of the body-to-signal transduction method, the state of the art, unsettled

challenges, and finally, comments on the prospects. The authors believe that the outcomes of this mini-review will serve as a basic text for beginners in the field of wearable sensors, and it will also be helpful to experienced researchers in wearables by expanding the knowledge of challenging sensing modalities and the vital obstacles that are being faced by wearable sensors and devices.

## 2. Historical viewpoint

There is a long history of wearable strain sensors. Some are briefly described herein. Wearables were first realized in the Apollo Space Program during the early 1960s, when scientists were well aware of the physical extremes of humans on this mission. They recognized the need for some specific non-invasive sensors<sup>30</sup> to continuously monitor the health status of astronauts, as well as transmit real-time data back to earth. Wearable sensors fulfilled this job in a very efficient manner. These sensors included an electrocardiogram (Fig. 2(a)), and a heated thermistor to sense breathing by measuring the cooling due to inhaled and exhaled air movement from the mouth. Within very little time, pulse oximetry proved itself as an entirely safe and standard measurement technique during general anesthesia (Fig. 2(b)). There was also a biosensor probe for the precise measurement of glucose in the body (Fig. 2(c)).<sup>34</sup> Wireless electrocardiogram (EKG) (Fig. 2(d)) heart rate monitors started as standard equipment in 1977. Wearable monitors first gained popularity in the early 1980s and started being used by the common man in the late 1980s.

Leland Clark and Ann Lyons from the Cincinnati Children's hospital developed the first precise glucose enzyme sensing



Fig. 1 Potential state-of-the-art applications of wearable strain sensors.





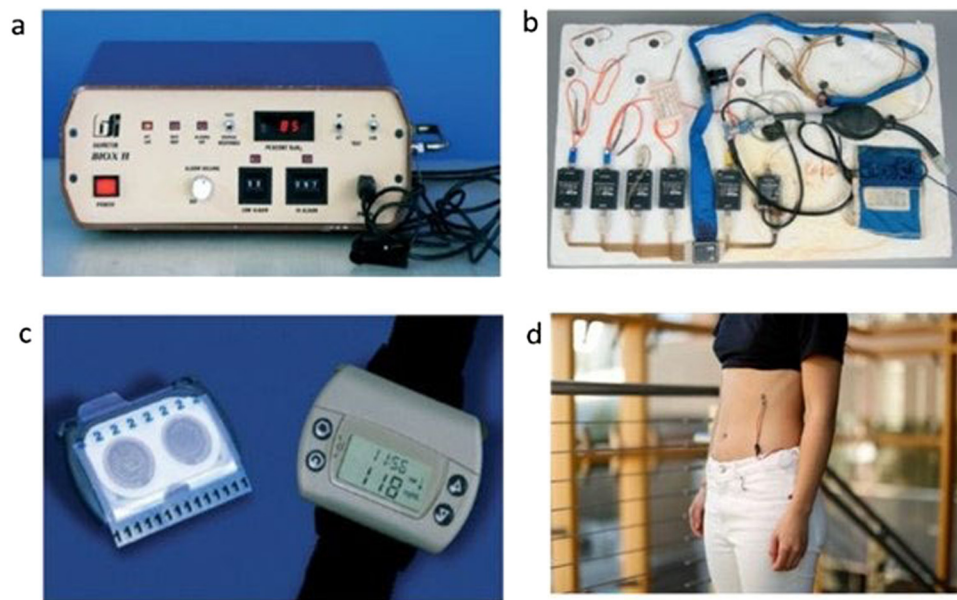


Fig. 2 Some historical glimpses of wearable sensors: (a) pulse oximetry for wearing on the fingertip, (b) wearable sensors used during the Apollo Space Program, (c) an early wearable digital glucose sensor for diabetic patients, and (d) 'Sport Tester PE2000' heart rate monitor by Polar. (Adapted and reprinted with permissions,<sup>35</sup> Copyright 2018, Royal Society of Chemistry).

electrode in 1962. However, it took a long time to realize the non-invasive wearable sensor. Electro-osmotic flow, also known as reverse iontophoresis, involves generating interstitial fluid through paracellular pathways so that the negatively charged plasma membranes can promote a moving electro-osmotic sheath of  $\text{Na}^+$  ions. Cygnus Inc. was granted FDA approval for its GlucoWatch, which was a significant novel noninvasive approach to diabetes monitoring since diabetes can be deadly if glucose monitoring is not performed accurately. However, to date, non-invasive wearable chemical sensors are still unavailable as popular products<sup>36</sup> (the broadly used transcutaneous glucose monitors are invasive and are not a part of this review).

Commercially available wearable sensors like watches by Fitbit and Apple and medical patches by Medtronic's SEEQ cardiac monitoring system are examples of the latest wearable strain sensors, which indicate the advancement of technology; the current wearable sensors consist of simple electrical and optical measurement components on the skin.

### 3. Wearable sensing mechanism

Inertial motion and plantar force sensors can empirically monitor the movement of the human body. The miniaturized size of these sensors makes them wearable. The measurement of linear acceleration, angular velocity, and the direction of human body movements allow inertial motion sensors to distinguish the postural sway of humans. Also, the measurement of the coefficient of performance (COP) path at the plantar surface of the foot and stance/swing time during walking enables plantar force sensors to perceive the postural sway and gait variability. Flexible and wearable electronics that can be implanted into clothing have recently drawn increasing interest.<sup>37,38</sup> In such a

flexible system, information on mechanical deformation has always been an important research topic.<sup>39–42</sup> The gauge factor of a strain sensor is low to facilitate a higher strain sensing range. Further, the mechanical properties of fundamentally fragile materials are incompatible with those of textiles for integrated systems.

Another approach is to fabricate sensing devices from soft and flexible materials. Composites of polymers and conductive fillers have been successfully demonstrated for this purpose. Such materials can sense mechanical strain by the electron tunneling effect between adjacent particles or by sensing the resistance change due to the opening and closing of micro-cracks by mechanical deformation.<sup>43–50</sup> These schemes have shown more extensive strain sensing ranges with similar gauge factors compared to rigid materials. Furthermore, they can be integrated into textiles with low manufacturing costs. The conductive composites are usually obtained by mixing an insulating polymer with carbon black powders or carbon nanotubes. In general, these flexible strain sensors are fabricated based on thin-film technologies with limited placements and configurations.

### 4. Classification of wearable sensors

In this section of the review article, we discuss optical, mechanical, electrical, and chemical sensors. The primary body-to-signal transduction method for each sensing modality presented is also explained in brief, and the real-life devices and their practical illustrations are discussed. Newly developed wearable physical sensors have enough stretchability and flexibility to resist the deformations caused by human activities.<sup>51–53</sup> For example, portable sensors measure physical factors in a patient's body condition, including the temperature of the body and skin, wrist, blood pressure, and skin fatigue. These aspects



critically indicate the complete health of a person. With this measuring capability, physical sensors can measure these physical parameters and can electronically monitor the human-machine interface, skin, or overall patient health.<sup>54–57</sup>

#### 4.1. Pressure sensors

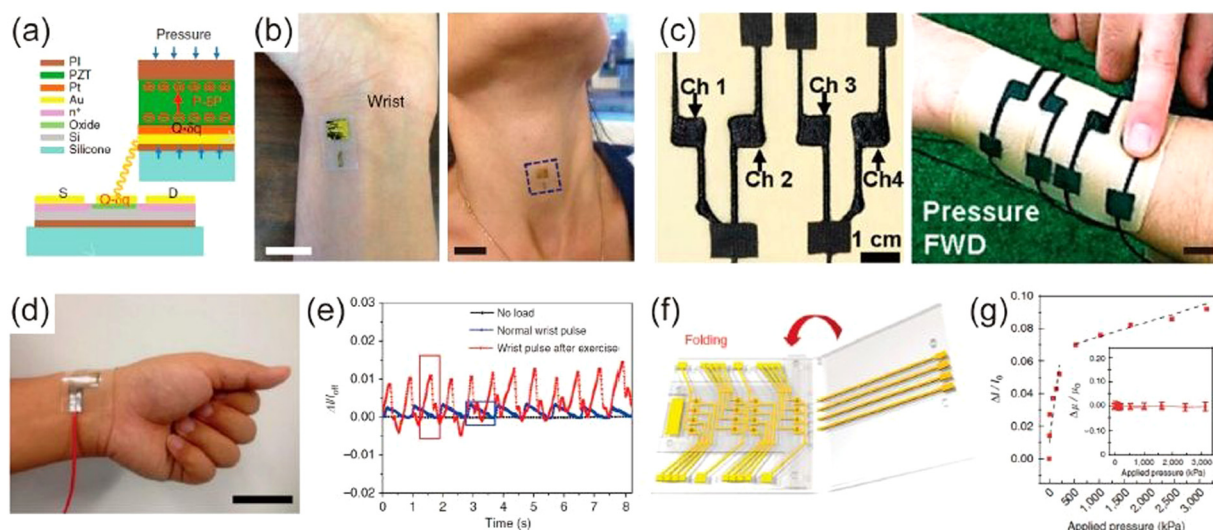
Pressure ranges vary differently from one part or target of the body to another to measure the parameters. For wearable pressure sensors, pressure ranges can be divided into three different categories, namely a range of low pressure (100 kPa),<sup>58</sup> which includes a person's weight or high-altitude atmospheric pressure.<sup>59</sup> Diseases related to the eyes, heart, damaged vocal cords, *etc.*, can be monitored by sensing the changes in some different types of pressure. The extensive studies on portable pressure sensors have opened a pathway for their utilization in personal health-related wellness and medical diagnostic instruments. Many sensing mechanisms, including piezoresistivity, piezoelectricity, and capacitance, can be used to convert physical stimuli into measurable electric signals. The basic principle of piezoelectric pressure sensors is founded on the piezoelectric effect, which is the change in the electrical charge that occurs in some specific solid materials as a result of the pressure applied on them. The following equations can indicate pressure-induced electricity:

$$D_i = \varepsilon_0 \varepsilon_{ij} \sigma E_j + d_{ij} \sigma_j \text{ and } \delta I = S_{ij} E \sigma_j + d_{Ij} E_j \quad (1)$$

where,  $S$ ,  $E$ ,  $\sigma$ ,  $\delta$ ,  $\varepsilon_0$ ,  $\varepsilon$ ,  $\sigma$ ,  $d$  and  $D$  denote the mechanical compliance matrix, electric field, the stress induced by applied pressure, the strain induced by applied pressure, the electrical permittivity of free space, the relative electrical permittivity, the

piezoelectric coefficient, and the electric displacement vector.  $i, j$ , and  $I, J$ , are different indices. The  $d_{31}$  and  $d_{33}$  are the transverse and longitudinal coefficients. Therefore, the piezoelectric effect is the induction of polarization in the dipole as a result of the pressure applied to the material. This polarization is proportional to the applied pressure. The piezoelectric pressure sensors are extensively used to detect different dynamic pressures, *e.g.*, sound vibrations, owing to their rapid response time and comparatively low energy consumption.  $\text{PbTiO}_3$ ,  $\text{BaTiO}_3$ , PVDF, and  $\text{P(VDF-TrFE)}$  are some of the appropriate materials used in the fabrication of piezoelectric pressure sensors. A representative material for a film-type sensor, known as P, <sup>60–64</sup> is used extensively to manufacture wearable pressure sensors. An ultrathin and ultra-high-quality PZT sheet is used as a component of a capacitor, and it connects to the gate electrodes of a FET (Fig. 3(a)). FETs intensify the piezoelectric output of the PZT and change it to an output current. This is a very thin and lightweight device with high sensitivity ( $\sim 0.005$  Pa) and fast response time (0.1 ms). It can be softly attached to the neck/throat or wrist to monitor blood pressure<sup>54,65</sup> (Fig. 3(b)).

The human skin can successfully adopt this wearable porous PSR pressure sensor for several applications, *e.g.*, man-machine interface, sanitary healthcare observations, and remotely controlling robots (Fig. 3(c)). Researchers have not only developed pressure sensors based on the microstructure and porous structures, but they have also successfully developed piezoresistive pressure sensors based on several other materials.<sup>66–68</sup> Gong *et al.* successfully fabricated a highly sensitive flexible pressure sensor by inserting gold nanowires (AuNWs) between two PDMS films.<sup>69</sup> The Au-NW-incorporated flexible pressure sensors



**Fig. 3** Wearable pressure sensing gadgets. (a) A schematic diagram of the cross-section of the pressure sensor and its contacts in an accompanying transistor. (b) The pressure sensor positioned on a wrist and neck to measure instantaneous variations in blood pressure<sup>70</sup> (reprinted with permission 2014, Copyright, Nature Publishing Group (NPG)). (c) Printed pressure sensor mounted on a commercially available elastomeric patch. The arrays of the sensor consist of four channels of pressure sensors<sup>71</sup> (reprinted with permission 2014, Copyright, John Wiley and Sons). (d) The skin-mounted sensor directly above the artery of the wrist (scale bar 3 cm). (e) The heartbeat is measured by sensing the physical force under healthy and exercising conditions<sup>72</sup> (reprinted with permission 2014, Copyright, Nature Publishing Group (NPG)). (f) A picture representation of pressure-sensitive graphene FETs. (g) A plot of changes in normalized drain current versus pressure applied on the device<sup>73</sup> (reprinted with permission 2014, Copyright, Nature Publishing Group (NPG)); (figure reproduced with permission,<sup>64</sup> Copyright 2017, from MDPI, Polymer).



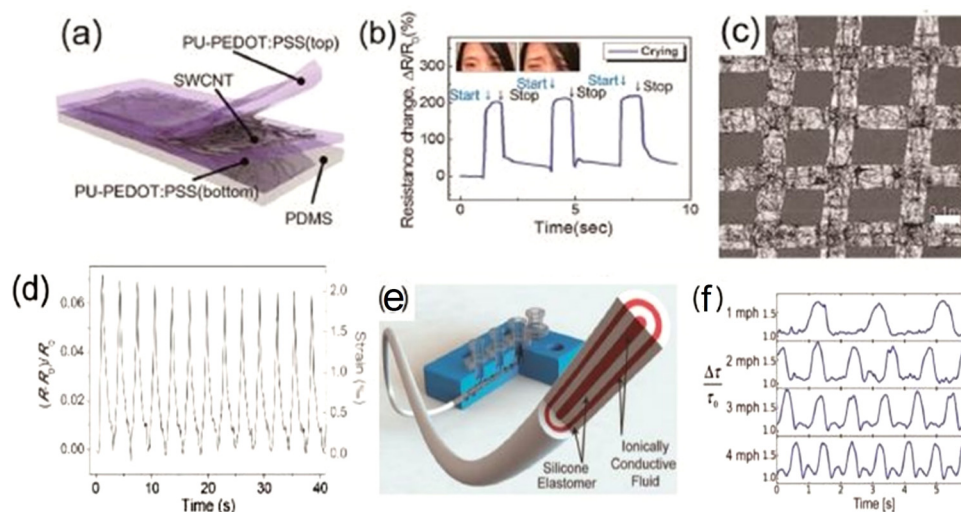
delivered real-time blood pressure monitoring with high precision and sensitivity, as shown in Fig. 3(d) and (e).

As is evident from the name, capacitors make up capacitive pressure sensors. Fundamentally, the magnitude of capacitance varies with the thickness of the layer of dielectric material. The capacitance  $C$  may be expressed by the relation  $C = \epsilon_0 \epsilon_r (A/d)$ , where  $A$  is the overlapping area between the two plates,  $d$  is the thickness of the dielectric material,  $\epsilon_0$  is the electrical permittivity of free space, and  $\epsilon_r$  is the relative static electrical permittivity of the dielectric. As an inference, the capacitance increases while externally applied pressure reduces the thickness. To alter the pressure-dependent thickness, materials with small modulus values, such as polyurethane (PU), PDMS, and eco-flex are generally used. Sun *et al.* fabricated a pressure sensor comprised of a stretchable and transparent dielectric material crammed in between two flexible ion conductors.<sup>31,68</sup> Owing to its transparency, stretchability, and better biocompatibility with the human body, this sensor can be applied in implantable or wearable electronic gadgets. Park *et al.* also developed a flexible and highly sensitive capacitive pressure sensor by raising an air gap between a single-walled carbon nanotube (SWCNT) film and a porous PDMS film.<sup>74</sup> Nevertheless, capacitive pressure sensors have certain restrictions, such as slow response time and low sensitivity due to the small modulus of the elastomer used in the fabrication. The use of air as a dielectric layer improves sensitivity. To resolve this issue, Zang *et al.* successfully demonstrated a flexible pressure sensor with remarkably high sensitivity in the low-pressure region, which was constructed on a suspended FET gate electrode.<sup>75</sup> It can be worn on the wrist and can monitor pulse waves. The pressure sensors described above are sensitive and can detect precise pressure intervals. However, a pressure sensor capable of detecting a wide range of pressures for diverse applications is

always needed. Very recently, Shin *et al.* also demonstrated an alternative approach to fabricating a pressure sensor array<sup>64</sup> filled with an air dielectric layer using folding panels (Fig. 3(f)). The sensor can be used successfully over a vast interval in tactile sensing, as shown in Fig. 3(g). All the results above have established that pressure sensing devices are arising as promising candidates for monitoring human movements, individual healthcare, and wellness.

#### 4.2. Strain sensors

While a person moves, small and large deformations occur. Wearable strain sensors must be able to precisely detect and observe the movements of the targeted organs of the human body continuously, whether this movement is generated from the faint vibration of the vocal cords or the activities of the joints; therefore, there is a need for flexible strain sensors. Because of the flexibility requirements, the strain sensors must consist of a flexible and stretchable material, and this differentiates them from conventional silicon-based strain sensors.<sup>76</sup> The constituent material of wearable strain sensors must be mechanically reliable, stretchable, sensitive, should have the property of hysteresis, and must produce a linear form of the output signal.<sup>77</sup> The classification of wearable strain sensors is according to the working mechanism used for sensing. Piezo-resistive strain sensors notice the deformations within the targeted body organ primarily by variations in the resistance. The following well-known equation may describe the change in the resistance of the materials used in piezoresistive strain sensors:  $R = \rho(l/A)$ , where  $\rho$  is resistivity,  $l$  is the length, and  $A$  is the area of the cross-section of the constituent material used in the piezoresistive sensor.  $A$  and  $l$  are both geometrical factors, and the piezoresistivity of the used materials induces changes in  $\rho$ . When a metal or semiconductor is deformed, its



**Fig. 4** Some examples of wearable strain sensors: (a) schematic cross-sectional diagram of the strain sensor comprised of 3-layers stacked in a nanohybrid structure. (b) Time-dependent resistance variations  $\Delta R/R_0$  of the sensors<sup>82</sup> (reprinted with permission, American Chemical Society, Copyright 2015). (c) Optical micrograph of a graphene woven fabric's relative change of resistance. (d) Relative resistance variation between 0% and 0.2% strain<sup>96</sup> (reprinted with permission, John Wiley & Sons, Copyright 2014). (e) Representation of the multicore-shell printing procedure for a fiber-type capacitive strain sensor. (f) Normalized decay-time response of the sensor (reprinted with permission John Wiley & Sons,<sup>87</sup> Copyright 2015).





interatomic spacing changes and as a result, its bandgap changes. This change is often expressed as the gauge factor,  $\Delta R/R$  of strain sensors. Consequently, because of substantial changes in resistivity, semiconductors have a high gauge factor. Carbon black, zinc oxide (ZnO), and CNT are some examples of semiconducting materials for use as piezoresistive strain sensors.<sup>63,78–81</sup> Roh *et al.* embedded single-walled carbon nanotubes in a conductive elastomer polyurethane-poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (PU-PEDOT:PSS) to fabricate the strain sensor<sup>82</sup> as shown in Fig. 4(a). This sensor exhibited a gauge factor of 62, stretchability of 100% strain, and 62% transmittance<sup>63,82</sup> in the visible range (Fig. 4(b)). Furthermore, deformations may lead to changes in the resistance (Fig. 4(c)) as a result of disconnections in the network geometry or cracks by the applied force or pressure. CNTs, graphene flakes/films, and metal nanowires (mNWs) are some examples (Fig. 4(d)) of materials possessing a conductive network. While these materials are stretched or bent under external force, there may be the sliding of atoms or molecules within the network, which produces a measurable change in resistance.<sup>69,83–86</sup> Frutiger *et al.* demonstrated a capacitive strain sensor<sup>87</sup> analogous to fiber using silicon elastomer and conductive ionic fluid (Fig. 4(g) and (h)). The strain can also be detected using the property of the piezoelectric effect of some explicit materials.<sup>88</sup> Various oxides have been used to develop a piezoelectric strain sensor.<sup>89–91</sup> Recently, a flexible piezoelectric sensor based on a ZnSnO<sub>3</sub> nano/microwire was fabricated successfully by Wu *et al.*; interestingly, the gauge factor of the sensor was 3740 at 0.35% strain.<sup>92</sup> There is no requirement for an external power source for piezoelectric strain sensors as they are mostly self-powered. However, the main challenges are still stretchability and flexibility because of the unsuitable mechanical properties of piezoelectric substances.<sup>91,93,94</sup>

#### 4.3. Piezoresistive sensors

Several transduction mechanisms with capacitance, piezoresistivity, piezoelectricity, *etc.*, are available for converting tactile stimuli into electrical signals.<sup>95,97–100</sup> Every transduction method has its characteristic features (Fig. 5), which are presented in the paragraph below. Accordingly,  $R$  is the resistance of the conductive material,  $A$  is the conductor's cross-sectional area,  $L$  is its length, and  $\rho$  is its resistivity. Wearable electronics frequently exploit the piezoresistive effect. The variation of resistance with the applied pressure and its transduction into electrical signals is the foundation principle of piezoresistive sensors (Fig. 5(a)). These sensors have been studied intensively due to the fact of their remarkably intense pixel density, comparatively simple device structure, and protected read-out mechanism algorithms.<sup>101</sup> The suitability of piezoresistive sensors over vast pressure ranges makes the precise measurement of large strains possible. The variation of contact resistance ( $R_c$ ) between two materials is governed by the applied forces ( $F$ ) and is the primary source of the change in the electrical signal.<sup>102</sup> The resistance of a pressure sensor changes as a consequence of the pressure applied to the device. The power law,  $R_c \sim F^{-1/2}$ , confirms the remarkably high sensitivity of piezoresistive sensors in low-pressure regimes and their relatively wide operating

ranges. Additionally, these devices have a fast response speed.<sup>103</sup>

The term capacitance characterizes the ability of a body to store electrical charge. An applied pressure produces a deflection in the plate, followed by a change in the capacitance of the used parallel-plate capacitor (Fig. 5(b)). These changes don't need to be linear and they may be of the order of several picofarads (pF). By changing the pressure-induced capacitance, the frequency of an oscillator can be controlled, or the coupling of an alternating current (AC) signal through a network can be varied. These sensors tend to exhibit low sensitivity due to the relatively small changes in the parallel plate capacitance.

The OFETs can be categorized among the most meticulously studied electronic components. A typical OFET structure comprises a semiconductor layer, a gate dielectric layer, source-drain electrodes, and a gate electrode. In a typical device structure, the gate dielectric layer is positioned between the gate electrode and the semiconductor layer. As a gate voltage is applied, the charge carriers gather at the interface of the dielectric/organic layer and establish a conductive channel and finally, current drifts from the source to drain electrodes while applying a source-drain bias.<sup>104–106</sup>

Piezoelectricity is also one of the frequently followed transduction methods to fabricate pressure sensors. Piezoelectricity denotes the generation of electrical charges in some specific solid materials as a result of the applied mechanical stress.<sup>107</sup> It is the presence of electric dipole moments in solids that contributes to the generation of the piezoelectric effect (Fig. 5(c)). Piezoelectric sensors can be used in a wide range of applications for the detection of dynamic pressures such as sound vibrations, owing to their remarkably high sensitivity and ultrafast response time. Piezoelectric sensors are promising for the development of ultra-low-power-consuming or even self-powered pressure sensors.<sup>14,108</sup>

#### 4.4. Wearable humidity sensors based on porous graphene:

One of the most important physiological markers that may be utilized to track a person's health and activity levels is respiration. The porous graphene network is flexible, light, and extremely conductive, thus making it a good candidate for a humidity sensor for respiration monitoring. Graphene oxide (GO), poly(3,4-ethylenedioxythiophene)-poly(styrene sulfonate) (PEDOT:PSS), and Ag colloids (AC) were employed to modify the porous graphene to improve the sensing performance. At various relative humidities (RH), the characteristics of porous-based graphene networks have been studied. The porous graphene sensors are capable of monitoring various breathing patterns, including mouth and nasal respiration, and normal and deep respiration. Additionally, the sensors record the signal changes before and after water consumption, demonstrating the capacity to monitor health processes. Furthermore, the humidity sensor showed the ability to detect physiological activities including skin moisture, and speaking and whistling rhythms, which could be promising for clinical respiration monitoring.<sup>109–111</sup>



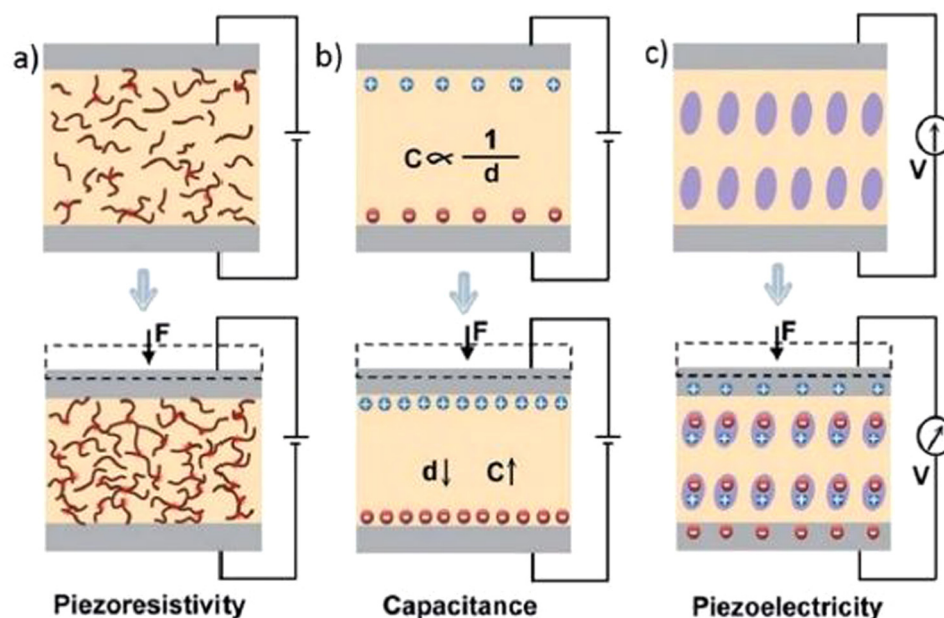


Fig. 5 Schematic illustrations<sup>109</sup> of transduction methods: (a) piezoresistivity, (b) capacitance, and (c) piezoelectricity. (Adapted and reprinted with permission, Copyright 2015, Royal Society of Chemistry).

## 5. Promising state-of-the-art strain sensing applications

Due to the excellent flexibility and fitting exhibited by flexible stress sensing materials, they have great potential in many applications, especially in the monitoring of stress and stress distribution on surfaces with three-dimensional irregular curved structures. The chemical fabrication process for flexible (humidity) stress sensors based on porous graphene network has been shown, step-by-step in Fig. 6(a)–(g). The process is initiated with a nickel foam template, then a graphene layer is prepared on the foam using CVD, followed by acidic chemical etching to obtain a porous network, chemical modification of this porous network, and finally transferring the as-obtained porous network to a flexible PET substrate. Flexible stress-sensing materials are commonly used in applications with different

types of stress regions.<sup>82,112–115</sup> The applications include ultra-low stress (<1 Pa) signal monitoring such as sound capture; weak stress (<1 kPa) signal acquisition, such as ultra-sensitive electronic skin, and touchpads; low stress (<10 kPa) signal detection, including motion monitoring in daily activities; and medium pressure (<100 kPa) signal acquisition, which includes plantar pressure collection.<sup>116–119</sup>

The current applications of wearable, flexible stress sensors are mainly concentrated in the fields of human physiological signal monitoring, motion and gesture recognition, flexible touch switches, electronic skin, and robots, which are introduced briefly below.

### 5.1. Human physiological signal monitoring

The flexible wearable stress sensors can realize real-time and long-term evaluation and feedback of individual physiological

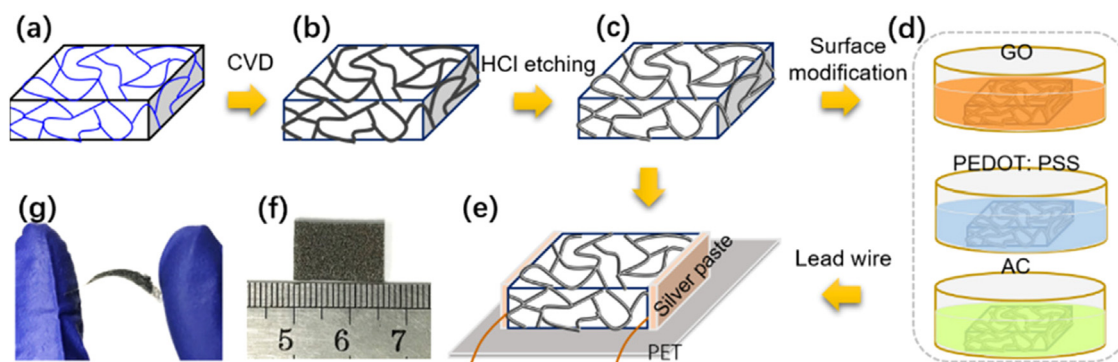
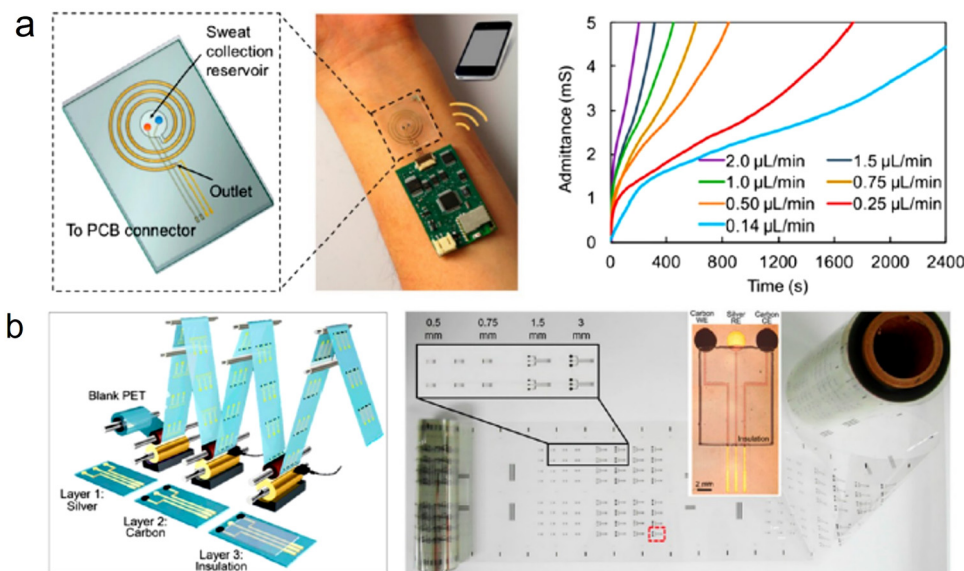


Fig. 6 Detailed step-by-step diagram of the fabrication process for devices. (A) A template made of nickel foam. (b) Using the CVD process, a graphene layer was created on the nickel foam. (c) A porous graphene network was created by etching the nickel skeleton with a solution of hydrochloric acid. (d) To obtain the surface-modified samples, the porous graphene networks were submerged in GO, PEDOT: PSS, and AC solutions. (e) Copper wires were led out after the modified samples were put into a flexible PET substrate. (f) The graphene/nickel foam image. (g) The image of the porous graphene network sensor demonstrates its ability to bend and stretch. Adapted and reprinted with permission,<sup>113</sup> Copyright 2018, Elsevier.







**Fig. 7** (a) Microfluidics-based sensing system for sweat sampling, sensing, and sweat rate analysis. Reprinted with permission, Copyright,<sup>109</sup> 2018 American Chemical Society. (b) Roll-to-roll gravure printing enabled mass production of high-performance flexible chemical sensors at low cost. Reproduced from ref.110 Copyright 2018 American Chemical Society.

data by avoiding the burden of heavy instruments and the storage of complicated cables. They can continuously perceive, acquire and transmit physiological signals such as the pulse and blood pressure of the human body without being affected by daily activities and can realize better personal medical treatment.<sup>76,77,120–123</sup> Currently, researchers have designed and developed a variety of flexible sweat sensors (Fig. 7) that monitor the pulse, blood pressure, and respiratory rate. Bao *et al.* designed a flexible stress sensor (Fig. 8(a)) with a micro-hair structure<sup>124</sup> that fits well on the surface of the human body and uses the micro-hair structure to adhere to the skin to monitor the depth. Wang *et al.* integrated the sensor with the breathing mask<sup>125</sup> to form a device that can monitor the respiratory rate of the human body and can accurately observe the breathing process in different states of the human body in real-time (Fig. 8(b)). To observe the pulsation of blood vessels, Lee *et al.* prepared a bending-insensitive stress sensor<sup>126–130</sup> based on graphene and carbon tube composites, which can continuously monitor the pressure on the artificial heart and blood vessel wall in real-time without being affected by activities such as human limb bending (Fig. 8(c)).

## 5.2. Motion and gesture recognition

Statistical data shows that when humans communicate, the amount of information transmitted by language is about 7%, and expressions and physical movements express about 50% of the messages. Therefore, facial expression recognition and gesture recognition have gradually become research hotspots.<sup>131</sup> A flexible and lightweight stress-strain sensor matrix can be placed on the face or different joints to analyze the expression and posture of the human body.<sup>132–134</sup> It is safe to wear without any harmful effects, and the cost-effective mass

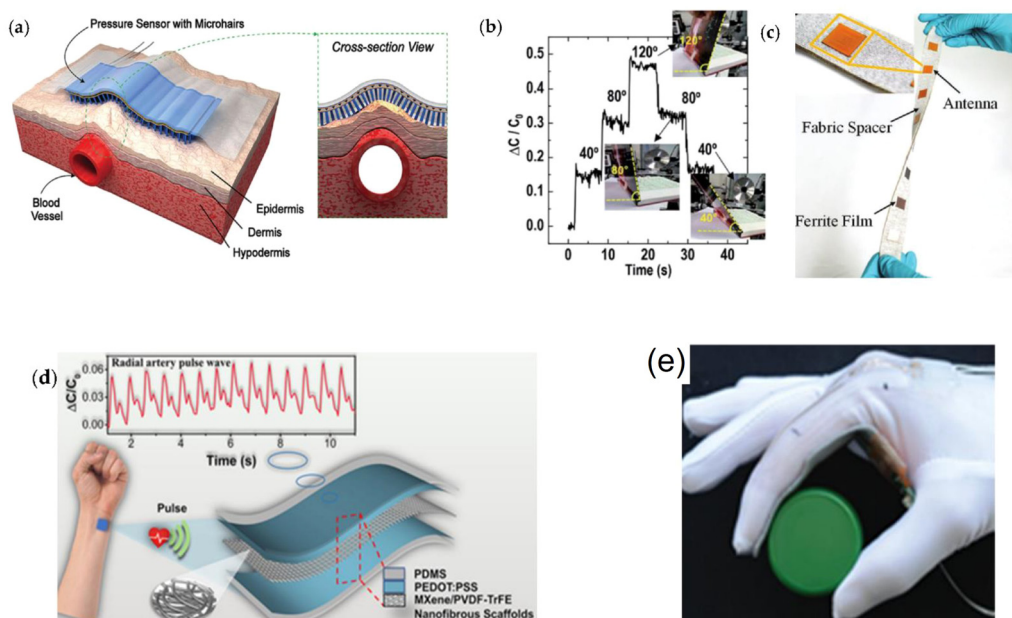
production of such devices is also possible. Su *et al.* laced multiple flexible stress sensors<sup>135</sup> on the forehead, eyebrows, nose, chin, and corners of the mouth for the multi-recognition of body expressions. Multi-channel linkage was used for smiles, laughter, surprise, sadness, fear, frustration, anger, and relaxation to identify the eight primary facial expressions. The same stress-strain sensor can also be integrated directly into the joints and muscles to detect movement of the joints or muscles<sup>136</sup> of the human body. The realization of the detection function of the joint activity of the human body can be applied in the medical field, such as in physical therapy rehabilitation, and prostheses.

Self-powered body motion skin sensors fabricated using triboelectric nanogenerator<sup>137</sup> (TENGs) can be categorized into several groups based on the triboelectric materials<sup>138</sup> used. The first type is built on materials that can be stretched, such as rubber and silicone elastomers. A stretchable rubber-based (SR-based) TENG was reported by Yin *et al.*,<sup>139</sup> which makes use of the triboelectricity between a stretchable rubber and an aluminum (Al) film, as shown in Fig. 9(a)–(c). The SR-based TENG operates on a novel theoretical foundation. It causes in-plane charge separation between rubber and Al instead of rubber shifting in position, which results in the potential difference between the Al electrode and the ground. Fig. 9(d)–(f) illustrates the soft, stretchable triboelectric band described by the author, which is made of a rubber tube filled with physiological saline.

## 5.3. Electronic skin and a flexible wearable touch switch

Electronic skin<sup>140–142</sup> is a flexible sensor device that mimics the human skin's perception and collection of external stimulus signals. Its main feature is the ability to monitor the stress





**Fig. 8** The applications of flexible stress sensors on (a) a pulse-detectable pressure sensor with a microhair structure (reprinted with permission from ref. 126. Copyright 2014 WILEY-VCH Verlag GmbH & Co. KGaA, Weinheim). (b) Capacitance response plot obtained from the wrinkled pressure sensor (reprinted with permission from ref. 127. Copyright 2017 Royal Society of Chemistry). (c) Textile-based flexible wireless pressure sensor (reprinted with permission from ref. 128. Copyright 2019 John Wiley and Sons). (d) Wearable capacitive pressure sensor based on MXene for reliable human physiological signals (reprinted with permission from ref. 129. Copyright 2020 American Chemical Society). (e) Textile glove with capacitive pressure sensors (reprinted with permission from ref. 130. Copyright 2015 John Wiley and Sons).

distribution at the pixel points in the 3D surface in real time.<sup>69,143–145</sup> Electronic skin was prepared by Liu *et al.*<sup>146</sup> using an elastic carbon sponge obtained by the carbonization of melamine, which utilizes inter-digitated electrodes to increase sensitivity. The ability of the material to contact the electrode<sup>147</sup> results in stress distribution and the magnitude is determined by scanning the rate of change of resistance at each pixel in the sensor dot matrix. The case shown is a schematic diagram for detecting the areas and sizes of different chess pieces.

Similar to electronic skin, after integration, the trajectory of the finger or other object moving on the sensor can be observed, and the stress between them can also be recorded by the sensor array so that it can be applied to the flexible touch-field,<sup>148–150</sup> (Fig. 10(a)). There are integrated sensor components with different shapes on the wristband, and wireless manipulation of the wrist band for household appliances.<sup>151</sup> They have been proven to be useful for Washable Electronic Textiles<sup>152</sup> as Self-Powered Touch/Gesture Tribo-Sensors for Intelligent Human–Machine Interactions (Fig. 10(b)). Flexible touch panels may also achieve the remote control of the turning on and off of household appliances such as lamps, fans, and microwave ovens *via* signal amplifiers (Fig. 10(c)), relays, wireless transmission (Fig. 10) and receiving devices.<sup>153</sup>

#### 5.4. Robots and intelligent interaction

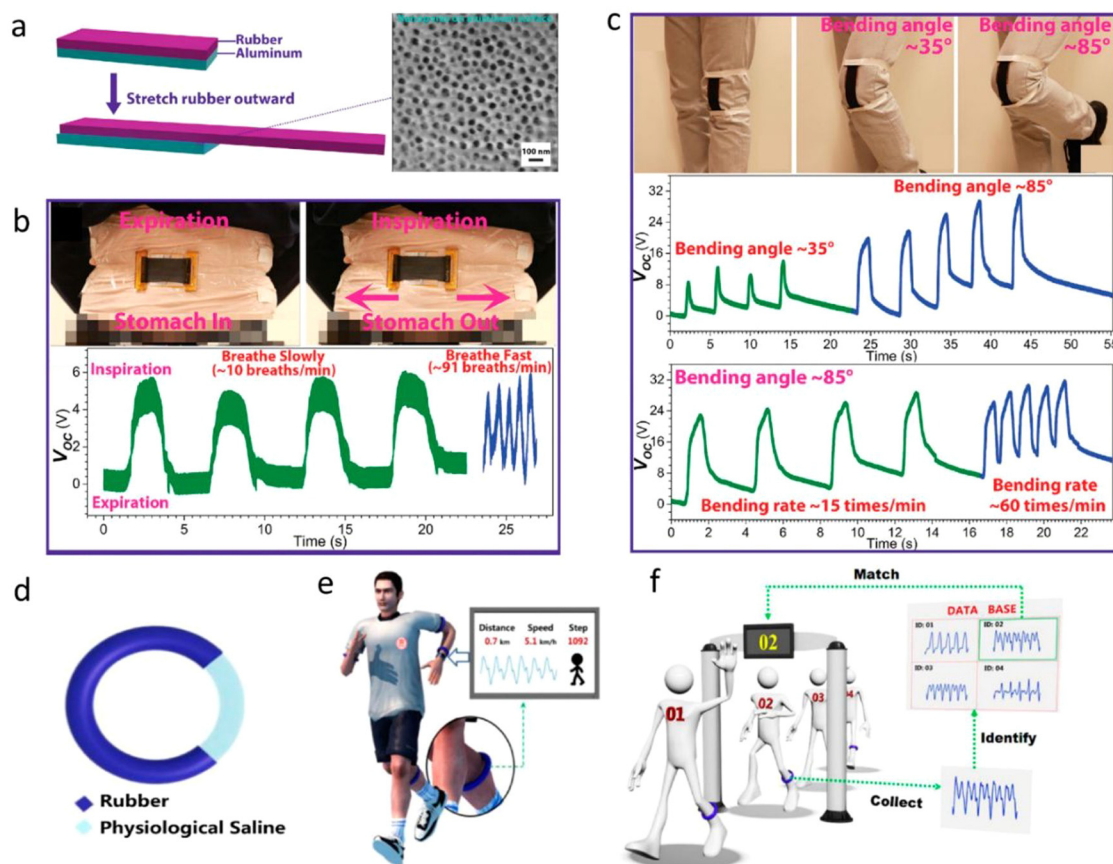
The stress sensor can accurately capture the fine movements of the human body or humanoid robots and can thus be applied to the field of robots and intelligent interactions.<sup>154–160</sup>

At present, the stress sensors used in robotics or intelligent interaction fields are mainly multi-channel work. Similar to expression recognition, the multi-channel sensors are used to monitor the stress and strain of different parts, and the action or posture of the object to be tested is judged in real-time.<sup>161–163</sup> Fig. 11(a)–(c) are the applications of robot motion recognition, somatosensory games, and wrist movement digitization.

## 6. Challenges in wearable strain sensing technology

There has been profound progress in the clever use of flexible and stretchable wearable sensors in this field throughout the years. The latest achievements and continuous advancement of the technology have proved the sensing conductors to be promising candidates in numerous profound applications to detect the motion of the human body by utilizing the simple principle of the variation of resistance in terms of a measurable electric current under ambient distortions, and thus an all-new sector of consumer health care has emerged. Numerous sensing devices concerning health-centered applications, which were just a theoretical possibility, have turned into reality. However, lots of significant challenges still exist on the long path towards precision and more diverse applications. For the further development of more innovative and versatile devices, a robust understanding of the transduction mechanism is a must because it is the most critical part of enhancing the overall





**Fig. 9** (a) Device structure of the stretchable rubber based on a triboelectric nanogenerator (TENG). (b) Images and electrical outputs of the TENG on the abdomen during expiration and inspiration. (c) Optical images of the device on the knee at different bending angles and voltage responses when bending the knee at different angles and different rates.<sup>139</sup> (Reprinted with permission Copyright 2015, John Wiley & Sons). (d) The typical device structure of the TENG band. (e) Schematic illustration of the skin TENG band for body motion detection.<sup>140</sup> (Reprinted with permission Copyright 2018, Elsevier). (f) Schematic illustration of the skin TENG band for identification. (Reprinted with permission Copyright 2018, Elsevier).<sup>140</sup>

detection precision and further enhancement of the gain factor of the transduced signal obtained as a result of deformations. Recently, the resistance and hence the current variation in strain sensors were proved to be produced due to numerous mechanisms, including piezoresistivity, the connection-disconnection mechanism, geometry effect, and tunneling effect. The scientific community needs to focus on further sharpening the knowledge and the understanding of robust mechanical properties to develop highly intelligent sensors for monitoring the specifically targeted organs of the human body; failure to do this could result in worse situations for individuals.

For the precise measurement of the data, a secure data read-write mechanism in severe circumstances is also of vital importance for real-world applications, along with a stable and diverse signal output.

## 7. Conclusions and outlooks

In this review, we have presented a compressive study on the recent progress of wearable strain sensors and advanced applications. The latest research outcomes of several papers in the field were considered to provide a glimpse of the state-of-the-art

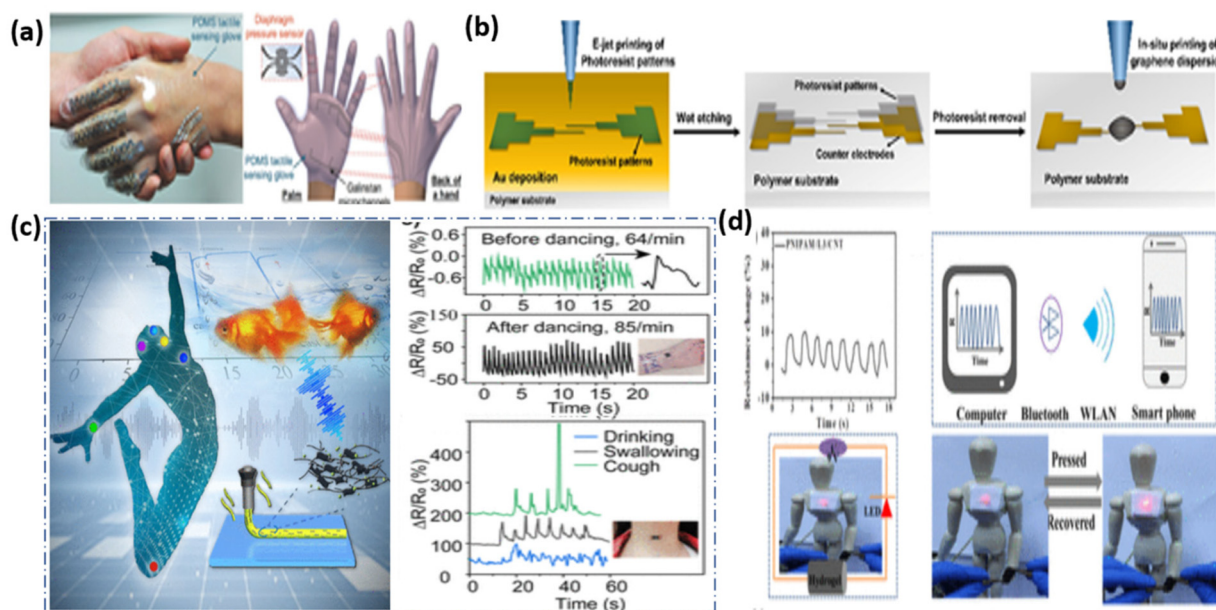
technology and to understand the physical phenomena behind the strain-responsive mechanisms of strain sensors at the microscale. Lots of conventional approaches, including several new ones such as disconnection between overlapped nanomaterials, crack proliferation in thin films, and the tunneling effect, have been used for the development of highly efficient flexible strain sensors.

After speedy progress on wearable strain sensors in the previous few decades, incredible advancements have been achieved, thus making the use of pressure sensors in real-life electronic applications possible. The thriving development of organic electronics inspired the unprecedented advancement of flexible pressure sensors with sensitive response proficiencies. Ultrasensitive, stretchable, and flexible pressure sensors are currently emerging as ideal candidates for wearable wellness monitoring applications. These accomplishments and potential applications have enabled flexible pressure sensors to be a crucial component of upcoming electronics.

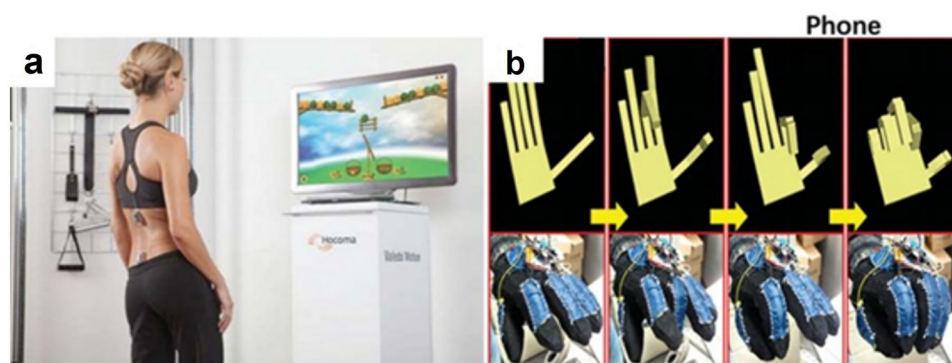
We strongly believe that stretchable, flexible strain sensors can serve as a vital component for several applications such as biomedicine, robotics, and entertainment sectors. However, high dynamic response with low overshoot and decay, a certain







**Fig. 10** (a) Photograph of a hand-shake greeting after wearing the PDMS tactile sensing gloves (left); representative diagram of the PDMS tactile sensing glove<sup>154</sup> (reprinted with permission Copyright 2017, John Wiley & Sons). (b) Representative diagram of the process of fabricating fully printed graphene photodetector gadgets using the mask-free direct-writing method<sup>155</sup> (reprinted with permission Copyright 2021, Elsevier). (c) Flexible composite pressure sensors for underwater monitoring (left); pulse wave curves and response signals under different conditions.<sup>156</sup> (Reprinted with permission Copyright 2022, American Chemical Society). (d) Photographs of the increased light intensity.<sup>157</sup> (Reprinted with permission Copyright 2019, American Chemical Society).



**Fig. 11** (a) Motion-sensing game with a wearable stress sensor system<sup>162</sup> (reprinted with permission Copyright 2012, BioMed Central). (b) The capture and digitalization of hand motion with a smart glove integrated with sensors.<sup>163</sup> (Reprinted with permission Copyright 2014, American Chemical Society).

degree of hysteresis, robust and flexible packaging, and conformal attachment to the targeted organs of the human body still need to be achieved.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

Each contributor thanks the appropriate government for providing the necessary resources and facilities. We gratefully

accept financial assistance from the Chinese National Natural Science Foundation.

## References

- 1 S. F. Clarke and J. R. Foster, *Br. J. Biomed. Sci.*, 2012, **69**, 83–93.
- 2 C. E. Sims and N. L. Allbritton, *Lab Chip*, 2007, **7**, 423–440.
- 3 T. Matsunaga, M. Hosokawa, A. Arakaki, T. Taguchi, T. Mori, T. Tanaka and H. Takeyama, *Anal. Chem.*, 2008, **80**, 5139–5145.
- 4 K. Klepárník and M. Horký, *Electrophoresis*, 2003, **24**, 3778–3783.



- 5 E. Proksch, J. M. Brandner and J.-M. Jensen, *Exp. Dermatol.*, 2008, **17**, 1063–1072.
- 6 X. Liu, C. Tang, X. Du, S. Xiong, S. Xi, Y. Liu, X. Shen, Q. Zheng, Z. Wang, Y. Wu, A. Horner and J.-K. Kim, *Mater. Horiz.*, 2017, **4**, 477–486.
- 7 Y. Wang, A. X. Wang, Y. Wang, M. K. Chyu and Q.-M. Wang, *Sens. Actuators, A*, 2013, **199**, 265–271.
- 8 M. Abkarian, M. Faivre and H. A. Stone, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**, 538–542.
- 9 M. L. Yarmush, A. Golberg, G. Serša, T. Kotnik and D. Miklavčič, *Annu. Rev. Biomed. Eng.*, 2014, **16**, 295–320.
- 10 H.-B. Yao, J. Ge, C.-F. Wang, X. Wang, W. Hu, Z.-J. Zheng, Y. Ni and S.-H. Yu, *Adv. Mater.*, 2013, **25**, 6692–6698.
- 11 L. Sheng, Y. Liang, L. Jiang, Q. Wang, T. Wei, L. Qu and Z. Fan, *Adv. Funct. Mater.*, 2015, **25**, 6545–6551.
- 12 Y. Wang, C. Xie, D. Liu, X. Huang, J. Huo and S. Wang, *ACS Appl. Mater. Interfaces*, 2016, **8**, 18652–18657.
- 13 A. El-Laboudi, N. S. Oliver, A. Cass and D. Johnston, *Diabetes Technol. Ther.*, 2013, **15**, 101–115.
- 14 Y. Zang, F. Zhang, C. Di and D. Zhu, *Mater. Horiz.*, 2015, **2**, 140–156.
- 15 S. Mitragotri, *Adv. Drug Delivery Rev.*, 2013, **65**, 100–103.
- 16 P. Garstecki, I. Gitlin, W. DiLuzio, G. M. Whitesides, E. Kumacheva and H. A. Stone, *Appl. Phys. Lett.*, 2004, **85**, 2649–2651.
- 17 S. Coyle, Y. Wu, K.-T. Lau, D. De Rossi, G. Wallace and D. Diamond, *MRS Bull.*, 2007, **32**, 434–442.
- 18 W. Zhong, H. Jiang, L. Yang, A. Yadav, X. Ding, Y. Chen, M. Li, G. Sun and D. Wang, *Polymers*, 2019, **11**, 1883.
- 19 W. Zhong, X. Ding, W. Li, C. Shen, A. Yadav, Y. Chen, M. Bao, H. Jiang and D. Wang, *Polymers*, 2019, **11**, 1289.
- 20 J. Heikenfeld, A. Jajack, J. Rogers, P. Gutruf, L. Tian, T. Pan, R. Li, M. Khine, J. Kim, J. Wang and J. Kim, *Lab Chip*, 2018, **18**, 217–248.
- 21 S. P. Nichols, A. Koh, W. L. Storm, J. H. Shin and M. H. Schoenfish, *Chem. Rev.*, 2013, **113**, 2528–2549.
- 22 X. Zhao, Q. Hua, R. Yu, Y. Zhang and C. Pan, *Adv. Electron. Mater.*, 2015, **1**, 1500142.
- 23 X. Liang and S. A. Boppart, *IEEE Trans. Biomed. Eng.*, 2010, **57**, 953–959.
- 24 M. Amjadi, Y. J. Yoon and I. Park, *Nanotechnology*, 2015, **26**, 375501.
- 25 X. Ye, Z. Yuan, H. Tai, W. Li, X. Du and Y. Jiang, *J. Mater. Chem. C*, 2017, **5**, 7746–7752.
- 26 T. Li, H. Luo, L. Qin, X. Wang, Z. Xiong, H. Ding, Y. Gu, Z. Liu and T. Zhang, *Small*, 2016, **12**, 5042–5048.
- 27 J. Wang, J. Jiu, M. Nogi, T. Sugahara, S. Nagao, H. Koga, P. He and K. Suganuma, *Nanoscale*, 2015, **7**, 2926–2932.
- 28 B. C.-K. Tee, A. Chortos, R. R. Dunn, G. Schwartz, E. Eason and Z. Bao, *Adv. Funct. Mater.*, 2014, **24**, 5427–5434.
- 29 G. Schwartz, B. C.-K. Tee, J. Mei, A. L. Appleton, D. H. Kim, H. Wang and Z. Bao, *Nat. Commun.*, 2013, **4**, 1859.
- 30 J. Levin and H. Maibach, *J. Controlled Release*, 2005, **103**, 291–299.
- 31 H. Sun, Y. Bu, H. Liu, J. Wang, W. Yang, Q. Li, Z. Guo, C. Liu and C. Shen, *Sci. Bull.*, 2022, **67**, 1669–1678.
- 32 D. Zhang, R. Yin, Y. Zheng, Q. Li, H. Liu, C. Liu and C. Shen, *Chem. Eng. J.*, 2022, **438**, 135587.
- 33 Y. Bu, T. Shen, W. Yang, S. Yang, Y. Zhao, H. Liu, Y. Zheng, C. Liu and C. Shen, *Sci. Bull.*, 2021, **66**, 1849–1857.
- 34 P. R. Miller, R. J. Narayan and R. Polsky, *J. Mater. Chem. B*, 2016, **4**, 1379–1383.
- 35 J. Heikenfeld, A. Jajack, J. Rogers, P. Gutruf, L. Tian, T. Pan, R. Li, M. Khine, J. Kim, J. Wang and J. Kim, *Lab Chip*, 2018, **18**, 217–248.
- 36 A. Abbosh, *Sensors*, 2019, **19**, 1662.
- 37 Q. Li, R. Yin, D. Zhang, H. Liu, X. Chen, Y. Zheng, Z. Guo, C. Liu and C. Shen, *J. Mater. Chem. A*, 2020, **8**, 21131–21141.
- 38 R. Yin, S. Yang, Q. Li, S. Zhang, H. Liu, J. Han, C. Liu and C. Shen, *Sci. Bull.*, 2020, **65**, 899–908.
- 39 D.-Y. Khang, H. Jiang, Y. Huang and J. A. Rogers, *Science*, 2006, **311**, 208–212.
- 40 D.-H. Kim, J.-H. Ahn, W. M. Choi, H.-S. Kim, T.-H. Kim, J. Song, Y. Y. Huang, Z. Liu, C. Lu and J. A. Rogers, *Science*, 2008, **320**, 507–511.
- 41 Y. Huang, W. Dong, C. Zhu and L. Xiao, *Complexity*, 2018, **2018**, e3016343.
- 42 R. Bogue, *Sens. Rev.*, 2015, **35**, 321–328.
- 43 A. Nocke, S. Richter, M. Wolf and G. Gerlach, *Procedia Chem.*, 2009, **1**, 1151–1154.
- 44 M. Farooq and E. Sazonov, in *Wearable Electronics Sensors: For Safe and Healthy Living*, ed. S. C. Mukhopadhyay, Springer International Publishing, Cham, 2015, pp.221–239.
- 45 M. Knite, V. Teteris, A. Kiploka and J. Kaupuzs, *Sens. Actuators, A*, 2004, **110**, 142–149.
- 46 T. Yamada, Y. Hayamizu, Y. Yamamoto, Y. Yomogida, A. Izadi-Najafabadi, D. N. Futaba and K. Hata, *Nat. Nanotechnol.*, 2011, **6**, 296–301.
- 47 D. J. Lipomi, M. Vosgueritchian, B. C.-K. Tee, S. L. Hellstrom, J. A. Lee, C. H. Fox and Z. Bao, *Nat. Nanotechnol.*, 2011, **6**, 788–792.
- 48 Y. Wang, H. Mi, Q. Zheng, H. Zhang, Z. Ma and S. Gong, *J. Mater. Chem. C*, 2016, **4**, 460–467.
- 49 Alamus, N. Hu, H. Fukunaga, S. Atobe, Y. Liu and J. Li, *Sensors*, 2011, **11**, 10691–10723.
- 50 C. Dagdeviren, Y. Su, P. Joe, R. Yona, Y. Liu, Y.-S. Kim, Y. Huang, A. R. Damadoran, J. Xia, L. W. Martin, Y. Huang and J. A. Rogers, *Nat. Commun.*, 2014, **5**, 4496.
- 51 R. C. Webb, A. P. Bonifas, A. Behnaz, Y. Zhang, K. J. Yu, H. Cheng, M. Shi, Z. Bian, Z. Liu, Y.-S. Kim, W.-H. Yeo, J. S. Park, J. Song, Y. Li, Y. Huang, A. M. Gorbach and J. A. Rogers, *Nat. Mater.*, 2013, **12**, 938–944.
- 52 M. Kaltenbrunner, T. Sekitani, J. Reeder, T. Yokota, K. Kuribara, T. Tokuhara, M. Drack, R. Schwödiauer, I. Graz, S. Bauer-Gogonea, S. Bauer and T. Someya, *Nature*, 2013, **499**, 458–463.
- 53 L. Y. Chen, B. C.-K. Tee, A. L. Chortos, G. Schwartz, V. Tse, D. J. Lipomi, H.-S. P. Wong, M. V. McConnell and Z. Bao, *Nat. Commun.*, 2014, **5**, 5028.
- 54 S. Jung, J. H. Kim, J. Kim, S. Choi, J. Lee, I. Park, T. Hyeon and D.-H. Kim, *Adv. Mater.*, 2014, **26**, 4825–4830.



- 55 W.-H. Yeo, Y.-S. Kim, J. Lee, A. Ameen, L. Shi, M. Li, S. Wang, R. Ma, S. H. Jin, Z. Kang, Y. Huang and J. A. Rogers, *Adv. Mater.*, 2013, **25**, 2773–2778.
- 56 J. Park, Y. Lee, J. Hong, Y. Lee, M. Ha, Y. Jung, H. Lim, S. Y. Kim and H. Ko, *ACS Nano*, 2014, **8**, 12020–12029.
- 57 S. C. B. Mannsfeld, B. C.-K. Tee, R. M. Stoltenberg, C. V. H.-H. Chen, S. Barman, B. V. O. Muir, A. N. Sokolov, C. Reese and Z. Bao, *Nat. Mater.*, 2010, **9**, 859–864.
- 58 Y. Wang, C. Zhu, R. Pfattner, H. Yan, L. Jin, S. Chen, F. Molina-Lopez, F. Lissel, J. Liu, N. I. Rabiah, Z. Chen, J. W. Chung, C. Linder, M. F. Toney, B. Murmann and Z. Bao, *Sci. Adv.*, 2017, **3**, e1602076.
- 59 X. Wang, T. Xu, S. Dong, S. Li, L. Yu, W. Guo, H. Jin, J. Luo, Z. Wu and J. M. King, *RSC Adv.*, 2017, **7**, 48461–48465.
- 60 S. Bauer and S. Bauer-Gogonea, in *Electromechanically Active Polymers: A Concise Reference*, ed. F. Carpi, Springer International Publishing, Cham, 2016, pp.1–15.
- 61 J. S. Lee, K.-Y. Shin, O. J. Cheong, J. H. Kim and J. Jang, *Sci. Rep.*, 2015, **5**, 7887.
- 62 S.-W. Kim, Y. Lee, J. Park, S. Kim, H. Chae, H. Ko and J. J. Kim, *Sensors*, 2018, **18**, 78.
- 63 D. Zaharie-Butucel, L. Digianantonio, C. Leordean, L. Ressler, S. Astilean and C. Farcau, *Carbon*, 2017, **113**, 361–370.
- 64 B. W. An, J. H. Shin, S.-Y. Kim, J. Kim, S. Ji, J. Park, Y. Lee, J. Jang, Y.-G. Park, E. Cho, S. Jo and J.-U. Park, *Polymers*, 2017, **9**, 303.
- 65 G. Yu, J. Hu, J. Tan, Y. Gao, Y. Lu and F. Xuan, *Nanotechnology*, 2018, **29**, 115502.
- 66 Y. Meng, H. Li, K. Wu, S. Zhang and L. Li, *Polymers*, 2018, **10**, 587.
- 67 D. Kang, P. V. Pikhitsa, Y. W. Choi, C. Lee, S. S. Shin, L. Piao, B. Park, K.-Y. Suh, T. Kim and M. Choi, *Nature*, 2014, **516**, 222–226.
- 68 J.-Y. Sun, C. Keplinger, G. M. Whitesides and Z. Suo, *Adv. Mater.*, 2014, **26**, 7608–7614.
- 69 S. Gong, D. T. H. Lai, B. Su, K. J. Si, Z. Ma, L. W. Yap, P. Guo and W. Cheng, *Adv. Electron. Mater.*, 2015, **1**, 1400063.
- 70 C. Dagdeviren, Y. Su, P. Joe, R. Yona, Y. Liu, Y.-S. Kim, Y. Huang, A. R. Damadoran, J. Xia, L. W. Martin, Y. Huang and J. A. Rogers, *Nat. Commun.*, 2014, **5**, 4496.
- 71 S. Jung, J. H. Kim, J. Kim, S. Choi, J. Lee, I. Park, T. Hyeon and D.-H. Kim, *Adv. Mater.*, 2014, **26**, 4825–4830.
- 72 S. Gong, W. Schwalb, Y. Wang, Y. Chen, Y. Tang, J. Si, B. Shirinzadeh and W. Cheng, *Nat. Commun.*, 2014, **5**, 3132.
- 73 S.-H. Shin, S. Ji, S. Choi, K.-H. Pyo, B. Wan An, J. Park, J. Kim, J.-Y. Kim, K.-S. Lee, S.-Y. Kwon, J. Heo, B.-G. Park and J.-U. Park, *Nat. Commun.*, 2017, **8**, 14950.
- 74 S. Park, H. Kim, M. Vosgueritchian, S. Cheon, H. Kim, J. H. Koo, T. R. Kim, S. Lee, G. Schwartz, H. Chang and Z. Bao, *Adv. Mater.*, 2014, **26**, 7324–7332.
- 75 Y. Zang, F. Zhang, D. Huang, X. Gao, C. Di and D. Zhu, *Nat. Commun.*, 2015, **6**, 6269.
- 76 T. Q. Trung and N.-E. Lee, *Adv. Mater.*, 2016, **28**, 4338–4372.
- 77 M. Amjadi, K.-U. Kyung, I. Park and M. Sitti, *Adv. Funct. Mater.*, 2016, **26**, 1678–1698.
- 78 N. Lu, C. Lu, S. Yang and J. Rogers, *Adv. Funct. Mater.*, 2012, **22**, 4044–4050.
- 79 C. Yan, J. Wang, W. Kang, M. Cui, X. Wang, C. Y. Foo, K. J. Chee and P. S. Lee, *Adv. Mater.*, 2014, **26**, 2022–2027.
- 80 T. Lee, W. Lee, S.-W. Kim, J. J. Kim and B.-S. Kim, *Adv. Funct. Mater.*, 2016, **26**, 6206–6214.
- 81 X. Xiao, L. Yuan, J. Zhong, T. Ding, Y. Liu, Z. Cai, Y. Rong, H. Han, J. Zhou and Z. L. Wang, *Adv. Mater.*, 2011, **23**, 5440–5444.
- 82 E. Roh, B.-U. Hwang, D. Kim, B.-Y. Kim and N.-E. Lee, *ACS Nano*, 2015, **9**, 6252–6261.
- 83 C. S. Boland, U. Khan, C. Backes, A. O'Neill, J. McCauley, S. Duane, R. Shanker, Y. Liu, I. Jurewicz, A. B. Dalton and J. N. Coleman, *ACS Nano*, 2014, **8**, 8819–8830.
- 84 Y. Wang, T. Yang, J. Lao, R. Zhang, Y. Zhang, M. Zhu, X. Li, X. Zang, K. Wang, W. Yu, H. Jin, L. Wang and H. Zhu, *Nano Res.*, 2015, **8**, 1627–1636.
- 85 B.-U. Hwang, J.-H. Lee, T. Q. Trung, E. Roh, D.-I. Kim, S.-W. Kim and N.-E. Lee, *ACS Nano*, 2015, **9**, 8801–8810.
- 86 B. Saha, S. Baek and J. Lee, *ACS Appl. Mater. Interfaces*, 2017, **9**, 4658–4666.
- 87 A. Frutiger, J. T. Muth, D. M. Vogt, Y. Mengüç, A. Campo, A. D. Valentine, C. J. Walsh and J. A. Lewis, *Adv. Mater.*, 2015, **27**, 2440–2446.
- 88 Y. Tomimatsu, H. Takahashi, T. Kobayashi, K. Matsumoto, I. Shimoyama, T. Itoh and R. Maeda, *J. Micromech. Microeng.*, 2013, **23**, 125023.
- 89 J. Lei, B. Yin, Y. Qiu, H. Zhang, Y. Chang, Y. Luo, Y. Zhao, J. Ji and L. Hu, *RSC Adv.*, 2015, **5**, 59458–59462.
- 90 G. Poulin-Vittrant, C. Oshman, C. Opoku, A. S. Dahiya, N. Camara, D. Alquier, L.-P. T. H. Hue and M. Lethiecq, *Phys. Procedia*, 2015, **70**, 909–913.
- 91 J. Zhong, Q. Zhong, Q. Hu, N. Wu, W. Li, B. Wang, B. Hu and J. Zhou, *Adv. Funct. Mater.*, 2015, **25**, 1798–1803.
- 92 J. M. Wu, C.-Y. Chen, Y. Zhang, K.-H. Chen, Y. Yang, Y. Hu, J.-H. He and Z. L. Wang, *ACS Nano*, 2012, **6**, 4369–4374.
- 93 Y. Wang, Y. Yu, X. Wei and F. Narita, *Adv. Mater. Technol.*, 2021, 2200318.
- 94 H. Høyer, M. Knaapila, J. Kjelstrup-Hansen and G. Helgesen, *J. Appl. Phys.*, 2012, **112**, 094324.
- 95 E. Roh, B.-U. Hwang, D. Kim, B.-Y. Kim and N.-E. Lee, *ACS Nano*, 2015, **9**, 6252–6261.
- 96 Y. Wang, L. Wang, T. Yang, X. Li, X. Zang, M. Zhu, K. Wang, D. Wu and H. Zhu, *Adv. Funct. Mater.*, 2014, **24**, 4666–4670.
- 97 W. Choi, J. Lee, Y. Kyoung Yoo, S. Kang, J. Kim and J. Hoon Lee, *Appl. Phys. Lett.*, 2014, **104**, 123701.
- 98 H. C. Lim, B. Schulkin, M. J. Pulickal, S. Liu, R. Petrova, G. Thomas, S. Wagner, K. Sidhu and J. F. Federici, *Sens. Actuators, A*, 2005, **119**, 332–335.
- 99 G. Darlinski, U. Böttger, R. Waser, H. Klauk, M. Halik, U. Zschieschang, G. Schmid and C. Dehm, *J. Appl. Phys.*, 2005, **97**, 093708.
- 100 J. Zhou, Y. Gu, P. Fei, W. Mai, Y. Gao, R. Yang, G. Bao and Z. L. Wang, *Nano Lett.*, 2008, **8**, 3035–3040.
- 101 L. Pan, A. Chortos, G. Yu, Y. Wang, S. Isaacson, R. Allen, Y. Shi, R. Dauskardt and Z. Bao, *Nat. Commun.*, 2014, **5**, 3002.





- 102 C.-L. Choong, M.-B. Shim, B.-S. Lee, S. Jeon, D.-S. Ko, T.-H. Kang, J. Bae, S. H. Lee, K.-E. Byun, J. Im, Y. J. Jeong, C. E. Park, J.-J. Park and U.-I. Chung, *Adv. Mater.*, 2014, **26**, 3451–3458.
- 103 M. L. Hammock, A. Chortos, B. C.-K. Tee, J. B.-H. Tok and Z. Bao, *Adv. Mater.*, 2013, **25**, 5997–6038.
- 104 Y. Guo, G. Yu and Y. Liu, *Adv. Mater.*, 2010, **22**, 4427–4447.
- 105 Y. Zang, F. Zhang, D. Huang, C. Di, Q. Meng, X. Gao and D. Zhu, *Adv. Mater.*, 2014, **26**, 2862–2867.
- 106 C. Di, Y. Liu, G. Yu and D. Zhu, *Acc. Chem. Res.*, 2009, **42**, 1573–1583.
- 107 K.-I. Park, J. H. Son, G.-T. Hwang, C. K. Jeong, J. Ryu, M. Koo, I. Choi, S. H. Lee, M. Byun, Z. L. Wang and K. J. Lee, *Adv. Mater.*, 2014, **26**, 2514–2520.
- 108 S. Xu, Y. Qin, C. Xu, Y. Wei, R. Yang and Z. L. Wang, *Nat. Nanotechnol.*, 2010, **5**, 366–373.
- 109 H. Y. Y. Nyein, L.-C. Tai, Q. P. Ngo, M. Chao, G. Zhang, W. Gao, M. Bariya, J. Bullock, H. Kim, H. M. Fahad and A. Javey, A Wearable Microfluidic Sweat Sensing Patch for Dynamic Sweat Secretion Analysis, *ACS Sens.*, 2018, **3**, 944–952.
- 110 M. Bariya, Z. Shahpar, H. Park, J. Sun, Y. Jung, W. Gao, H. N. N. Nyein, T. S. Liaw, L.-C. Tai, Q. P. Ngo, M. Chao, Y. Zhao, M. Hettick, G. Cho and A. Javey, Roll-to-Roll Gravure Printed Electrochemical Sensors for Wearable and Medical Devices, *ACS Nano*, 2018, **12**, 6978–6987.
- 111 A. dos Santos, E. Fortunato, R. Martins, H. Águas and R. Igreja, *Sensors*, 2020, **20**, 4407.
- 112 L. Lin, Y. Xie, S. Wang, W. Wu, S. Niu, X. Wen and Z. L. Wang, *ACS Nano*, 2013, **7**, 8266–8274.
- 113 X. He, Q. Liu, J. Wang and H. Chen, *Front. Mater. Sci.*, 2019, **13**, 305–313.
- 114 Y. Pang, J. Jian, T. Tu, Z. Yang, J. Ling, Y. Li, X. Wang, Y. Qiao, H. Tian, Y. Yang and T.-L. Ren, *Biosens. Bioelectron.*, 2018, **116**, 123–129.
- 115 L.-C. Tai, T. S. Liaw, Y. Lin, H. Y. Y. Nyein, M. Bariya, W. Ji, M. Hettick, C. Zhao, J. Zhao, L. Hou, Z. Yuan, Z. Fan and A. Javey, *Nano Lett.*, 2019, **19**, 6346–6351.
- 116 L. M. Castano and A. B. Flatau, *Smart Mater. Struct.*, 2014, **23**, 053001.
- 117 D. T.-P. Fong and Y.-Y. Chan, *Sensors*, 2010, **10**, 11556–11565.
- 118 S. C. Mukhopadhyay, *IEEE Sens. J.*, 2015, **15**, 1321–1330.
- 119 A. H. Abdul Razak, A. Zayegh, R. K. Begg and Y. Wahab, *Sensors*, 2012, **12**, 9884–9912.
- 120 J. R. Windmiller and J. Wang, *Electroanalysis*, 2013, **25**, 29–46.
- 121 X. Liu, B. Du, Y. Sun, M. Yu, Y. Yin, W. Tang, C. Chen, L. Sun, B. Yang, W. Cao and M. N. R. Ashfold, *ACS Appl. Mater. Interfaces*, 2016, **8**, 16379–16385.
- 122 Z. He, A. Elbaz, B. Gao, J. Zhang, E. Su and Z. Gu, *Adv. Healthcare Mater.*, 2018, **7**, 1701306.
- 123 Kenry, J. C. Yeo and C. T. Lim, *Microsyst. Nanoeng.*, 2016, **2**, 1–19.
- 124 T. Wang, H. Yang, D. Qi, Z. Liu, P. Cai, H. Zhang and X. Chen, *Small*, 2018, **14**, 1702933.
- 125 X. Wang, Y. Gu, Z. Xiong, Z. Cui and T. Zhang, *Adv. Mater.*, 2014, **26**, 1336–1342.
- 126 C. Pang, J. H. Koo, A. Nguyen, J. M. Caves, M. G. Kim, A. Chortos, K. Kim, P. J. Wang, J. B. Tok and Z. Bao, Highly skin-conformal microhairy sensor for pulse signal amplification, *Adv. Mater.*, 2015, **27**, 634–640.
- 127 S. Baek, H. Jang, S. Y. Kim, H. Jeong, S. Han, Y. Jang, D. H. Kim and H. S. Lee, Flexible piezocapacitive sensors based on wrinkled microstructures: Toward low-cost fabrication of pressure sensors over large areas, *RSC Adv.*, 2017, **7**, 39420–39426.
- 128 B. Nie, R. Huang, T. Yao, Y. Zhang, Y. Miao, C. Liu, J. Liu and X. Chen, Textile-Based Wireless Pressure Sensor Array for Human-Interactive Sensing, *Adv. Funct. Mater.*, 2019, **29**, 1808786.
- 129 S. Sharma, A. Chhetry, M. Sharifuzzaman, H. Yoon and J. Y. Park, Wearable Capacitive Pressure Sensor Based on MXene Composite Nanofibrous Scaffolds for Reliable Human Physiological Signal Acquisition, *ACS Appl. Mater. Interfaces*, 2020, **12**, 22212–22224.
- 130 A. P. Gerratt, H. O. Michaud and S. P. Lacour, Elastomeric Electronic Skin for Prosthetic Tactile Sensation, *Adv. Funct. Mater.*, 2015, **25**, 2287–2295.
- 131 X. Guo, Y. Huang, Y. Zhao, L. Mao, L. Gao, W. Pan, Y. Zhang and P. Liu, *Smart Mater. Struct.*, 2017, **26**, 095017.
- 132 K. Muldoon, Y. Song, Z. Ahmad, X. Chen and M.-W. Chang, *Micromachines*, 2022, **13**, 642.
- 133 G. Jia, H. Wang, L. Yan, X. Wang, R. Pei, T. Yan, Y. Zhao and X. Guo, *Environ. Sci. Technol.*, 2005, **39**, 1378–1383.
- 134 D. Kwon, T.-I. Lee, J. Shim, S. Ryu, M. S. Kim, S. Kim, T.-S. Kim and I. Park, *ACS Appl. Mater. Interfaces*, 2016, **8**, 16922–16931.
- 135 M. Su, F. Li, S. Chen, Z. Huang, M. Qin, W. Li, X. Zhang and Y. Song, *Adv. Mater.*, 2016, **28**, 1369–1374.
- 136 A. M. Almassri, W. Z. Wan Hasan, S. A. Ahmad, A. J. Ishak, A. M. Ghazali, D. N. Talib and C. Wada, *J. Sens.*, 2015, **2015**, e846487.
- 137 R. Cao, X. Pu, X. Du, W. Yang, J. Wang, H. Guo, S. Zhao, Z. Yuan, C. Zhang, C. Li and Z. L. Wang, *ACS Nano*, 2018, **12**, 5190–5196.
- 138 F. Rahimi Sardo, A. Rayegani, A. Matin Nazar, M. Balaghiinaloo, M. Saberian, S. A. H. Mohsan, M. H. Alsharif and H.-S. Cho, *Biosensors*, 2022, **12**, 697.
- 139 F. Yi, L. Lin, S. Niu, P. K. Yang, Z. Wang, J. Chen, Y. Zhou, Y. Zi, J. Wang and Q. Liao, *et al.*, *Adv. Funct. Mater.*, 2015, **25**, 3688–3696.
- 140 Y. Han, F. Yi, C. Jiang, K. Dai, Y. Xu, X. Wang and Z. You, *Nano Energy*, 2018, **56**, 516–523.
- 141 J. Rao, Z. Chen, D. Zhao, Y. Yin, X. Wang and F. Yi, *Sensors*, 2019, **19**, 2763.
- 142 X. Wang, L. Dong, H. Zhang, R. Yu, C. Pan and Z. L. Wang, *Adv. Sci.*, 2015, **2**, 1500169.
- 143 W. Liu, N. Liu, Y. Yue, J. Rao, C. Luo, H. Zhang, C. Yang, J. Su, Z. Liu and Y. Gao, *J. Mater. Chem. C*, 2018, **6**, 1451–1458.
- 144 C. G. Núñez, W. T. Navaraj, E. O. Polat and R. Dahiya, *Adv. Funct. Mater.*, 2017, **27**, 1606287.



- 145 S. Gong, D. T. H. Lai, Y. Wang, L. W. Yap, K. J. Si, Q. Shi, N. N. Jason, T. Sridhar, H. Uddin and W. Cheng, *ACS Appl. Mater. Interfaces*, 2015, **7**, 19700–19708.
- 146 Y. Liu, H. Wang, W. Zhao, M. Zhang, H. Qin and Y. Xie, *Sensors*, 2018, **18**, 645.
- 147 J. Kim, W. J. da Silva, A. R. bin Mohd Yusoff and J. Jang, *Sci. Rep.*, 2016, **6**, 19813.
- 148 Y. Li, B. Zhou, G. Zheng, X. Liu, T. Li, C. Yan, C. Cheng, K. Dai, C. Liu, C. Shen and Z. Guo, *J. Mater. Chem. C*, 2018, **6**, 2258–2269.
- 149 S. J. Lim, J. H. Bae, S. J. Jang, J. Y. Lim and J. H. Ko, *Fibers Polym.*, 2018, **19**, 2622–2630.
- 150 C. Nam and D. Shin, *Int. J. Distrib. Sens. Netw.*, 2018, **14**, 1550147718767794.
- 151 X. Liu, J. Miao, Q. Fan, W. Zhang, X. Zuo, M. Tian, S. Zhu, X. Zhang and L. Qu, *Adv. Fiber Mater.*, 2022, **4**, 361–389.
- 152 H. Liu, Q. Li, Y. Bu, N. Zhang, C. Wang, C. Pan, L. Mi, Z. Guo, C. Liu and C. Shen, *Nano Energy*, 2019, **66**, 104143.
- 153 H. Jin, S. Jung, J. Kim, S. Heo, J. Lim, W. Park, H. Y. Chu, F. Bien and K. Park, *Sci. Rep.*, 2017, **7**, 10854.
- 154 Y. Gao, H. Ota, E. W. Schaler, K. Chen, A. Zhao, W. Gao, H. M. Fahad, Y. Leng, A. Zheng, F. Xiong, C. Zhang, L.-C. Tai, P. Zhao, R. S. Fearing and A. Javey, *Adv. Mater.*, 2017, **29**, 1701985.
- 155 W. Zou, H. Yu, P. Zhou, Y. Zhong, Y. Wang and L. Liu, *Appl. Surf. Sci.*, 2021, **543**, 148800.
- 156 Y. Song, H. Dong, W. Liu, X. Fu, Z. Fu, P. Li, L. Chen, Z. Ahmad, J. Liu, X. Chen and M.-W. Chang, *ACS Appl. Polym. Mater.*, 2022, **4**, 868–878.
- 157 Z. Deng, T. Hu, Q. Lei, J. He, P. X. Ma and B. Guo, *ACS Appl. Mater. Interfaces*, 2019, **11**, 6796–6808.
- 158 S. Kagami, K. Nishiwaki, J. Kuffner, S. Thompson, J. Chestnutt, M. Stilman and P. Michel, in *Robotics Research*, ed. S. Thrun, R. Brooks and H. Durrant-Whyte, Springer, Berlin, Heidelberg, 2007, pp.103–117.
- 159 H. Morishita, R. Fukui and T. Sato, in *IEEE/RSJ International Conference on Intelligent Robots and Systems*, 2002, vol. 2, pp. 1246–1251 vol.2.
- 160 V. J. P. Amorim, M. C. Silva and R. A. R. Oliveira, *Sensors*, 2019, **19**, 1904.
- 161 Z. He, W. Chen, B. Liang, C. Liu, L. Yang, D. Lu, Z. Mo, H. Zhu, Z. Tang and X. Gui, *ACS Appl. Mater. Interfaces*, 2018, **10**, 12816–12823.
- 162 S. Patel, H. Park, P. Bonato, L. Chan and M. Rodgers, *J. Neuroeng. Rehabilitation*, 2012, **9**, 21.
- 163 M. Amjadi, A. Pichitpajongkit, S. Lee, S. Ryu and I. Park, *ACS Nano*, 2014, **8**, 5154–5163.

