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# **Review of pH sensing materials from macro- to nano-scale: recent developments and examples of seawater applications**

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## Abstract

Over the last decades, a large number of pH sensitive materials with new compositions and structures have been proposed. Solid state sensors based on organic, inorganic and composite materials are actively investigated, with an increasing interest in the performances offered by nano-scale materials. Our review provides a thorough, up-to-date knowledge of a wide range of pH measurement methods and related-sensing materials, first introducing well established materials and methods for pH sensing and then covering recent developments in inorganic, organic and nano-engineered devices. The main sensor parameters, including sensitivity, stability, response time and testing conditions are reported. Given the importance of pH sensing in environmental applications, in particular seawater monitoring, sensors tested in seawater are highlighted and discussed.

**Key words:** pH sensors; environmental monitoring; nanomaterials; water quality

## 1. Introduction

Due to the relevance of pH for many chemical and biochemical processes, pH measurements are routinely carried out in a very broad range of activities, from industrial processes to chemistry, medicine and environmental monitoring.

pH strongly affects environmental and biological processes. The availability of nutrients, the uptake of pollutants like heavy metals, the occurrence and distribution of microorganisms, the efficiency of enzymatic bioprocesses and metabolism, the occurrence of oxidative stress and its consequences on living organisms, are all pH-related phenomena (González Durán et al., 2018; Jin & Kirk, 2018; Kahn et al., 2017). Accurate quantification of pH is then vital for monitoring and protecting the health of our planet. In particular, pH is intimately linked with the dynamics of nutrients, contaminants, and trace metals in seawater and is entangled with the complex ocean carbonate system. As the pH of ocean surface decreases (-0.15 since pre-industrial times due to increasing dissolution of atmospheric CO<sub>2</sub>, Clarke et al., 2015), the delicate equilibria among chemical species in solution are perturbed, with effects on coastal biodiversity, ecosystem functioning (Lacoue-Labarthe et al., 2016) and the health of ecosystems worldwide (Kroeker et al., 2013; Somero et al., 2016). Continuous, accurate and punctual recording of seawater pH is needed to increase our understanding of the local and global pH dynamics and enable a better prediction of their effects (Bushinsky et al., 2019; Stow et al., 2009).

Ion sensitive glass electrodes are the most popular pH sensors, due to their reliability, affordability and fast (few s) response time. This includes environmental applications like seawater monitoring and most oceanic probes are equipped with this kind of pH sensors for routine pH recording. However, glass electrodes exhibit signal instability or drift and, therefore, require constant re-calibration: this operation can cause significant error, that may arise from the quality and handling of the calibration solutions (McLaughlin et al., 2017b). The need for an inner electrolyte solution, connecting the reference electrode with the sample solution through a liquid junction, can be another source of error as the potential that develops across the junction varies as a consequence of external factors like pressure. Finally, glass electrodes are brittle, need a storage solution and cannot be miniaturized.

For all these reasons, a number of alternative pH sensing devices have been proposed over the past decades. High precision measurements (up to 0.001 pH units) can be provided by spectrophotometric devices that are, however, much more expensive and complex than potentiometric sensors and have long sampling time (up to minutes). Solid state sensors can provide a cheap, robust and miniaturizable alternative for pH measurements (Korostynska et al., 2007), as demonstrated by the presence on the market of Ion Sensitive Field-Effect Transistors (ISFETs) based pH probes. These features can be exploited to realize sensing system with low cost, low power consumption and ease of operation (Radu et al., 2015). In the case of seawater monitoring, desired uncertainties for pH sensing have been specified as 0.02 for the study of short term, local variation and 0.003 for global, long term trends (Newton et al., 2015) and the quest for sensors with optimal field performances is

still open (Okazaki et al., 2017). A discussion of problems and techniques related to the measurement of pH in marine waters can be found in specialized papers (Byrne, 2014; Marion et al., 2011).

This review will discuss developments in the field of solid-state pH sensors, covering organic, inorganic and composite sensing materials and focusing on recent devices based on nanomaterials. Parameters like sensitivity, stability, robustness to interfering ions and response time of the sensors will be reported and organized in tables for a fast reference. Recent examples of pH sensors developed for seawater applications will be provided and critically reviewed at the end of each chapter. Providing a thorough, up-to-date knowledge of a wide range of pH measurement methods and related-sensing materials, our review may assist materials scientists, sensors developers and marine scientists interested in new pH sensing solutions.

## 2. Traditional methods and materials for pH measurement

The hydrogen ion is a ubiquitous species that plays a role in most chemical and biochemical reactions carried out in aqueous solutions. Firstly introduced by the Danish biochemist Søren Peter Lauritz Sørensen, pH is defined as the negative logarithm of  $H^+$  activity (Sørensen, 1909; Buck et al., 2002):

$$pH = -\log(a_{H^+}) \quad (1)$$

Due to the importance of this parameter for a wide range of applications, pH measurements are routinely performed in chemical, industrial, biological and medical practice. In the following sections, well established measurement techniques will be summarized, introducing some examples of seawater-designed devices.

### 2.1 Optical/spectrophotometric methods

A practical measurement of pH can be obtained using the so-called acid-base indicators, substances that change their color as a function of pH. In general, an indicator dye is an amphoteric compound with a dissociation constant that is close to the pH to be determined. The pH of the sample-indicator system can be expressed as a function of the dissociation constant of the indicator ( $pK$ ) and of the concentration of its protonated ( $HA$ ) and unprotonated ( $A^-$ ) form:

$$pH = pK + \log \frac{[A^-]}{[HA]} \quad (2)$$

As the two forms of the indicator in solution have different colors due to different light absorption, their concentration can be measured from their absorption spectra.

Based on this principle, spectrophotometric methods for pH measurement, reaching an accuracy as high as 0.001, have been developed using different indicators such as m-cresol purple, cresol red, bromocresol green, bromocresol purple and thymol blue (King & Kester, 1989; Millero et al., 2009). A schematic example of automated spectrophotometric pH system is reported in Figure 1. Once calibrated, these devices do not need to be recalibrated for use at sea. A description of a spectrophotometric pH sensor designed for in situ measurements can be found in Cullison Gray et al. (2011) and in Lai et al. (2018).

Recent technological developments of optical/spectrophotometric-based sensors represent a promising tool for monitoring the ocean carbonate system. In particular, pH sensors using spectrophotometric techniques are currently used for surface water measurements on research vessels and, similarly, optodes for pCO<sub>2</sub> measurements have been successfully tested in seawater for oceanographic applications (Rérolle et al., 2018; Staudinger et al., 2018; 2019 and references therein). Optical methods for pH detection will not be further discussed. A comprehensive review can be found in (Rérolle et al., 2012).

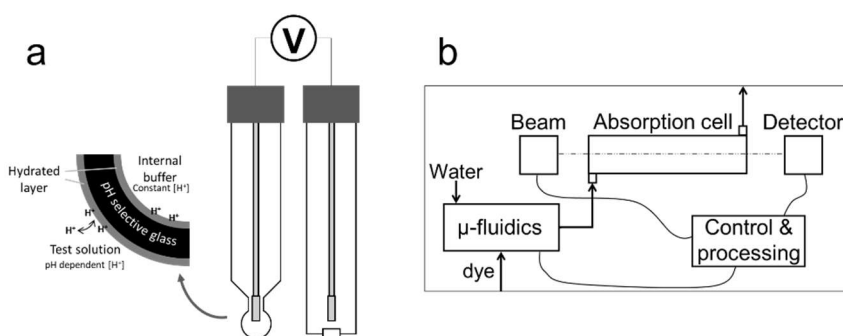
## 2.2 Electrochemical methods

Probably the most common techniques for pH sensing are based on the measurement of electrical parameters, such as conductivity or resistivity, impedance, potential. Conductometric devices correlate the change in conductivity/resistivity of an active material connecting two electrodes to the concentration of the analyte (H<sup>+</sup> for pH). Voltammetric devices measure the current flowing between the electrodes when the potential is swept in a defined manner; in this case, the pH measurement can be correlated to a peak potential of an electroactive compound (Dai et al., 2016).

Potentiometric sensors are the most used for routine pH determination. In principle, a potentiometric measurement consists of the measurement of the electromotive force (EMF) in an electrochemical cell, composed of a working electrode and a reference electrode. The pH of the sample is calculated comparing the EMF measured in the sample (E<sub>s</sub>) and in a standard buffer solution (E<sub>b</sub>) of known pH (pH<sub>b</sub>), following the Nernst equation:

$$pH = pH_b + \frac{(E_b - E_s)F}{RT \ln 10} \quad (3)$$

where R is the gas constant, F is the Faraday constant and T is the temperature (Rérolle et al., 2012).



**Figure 1.** Scheme of: a potentiometric pH sensor with glass ion sensitive electrode (a) and a spectrophotometric pH measurement device (b).

The most used working electrode for this application is made of a silver/silver chloride electrode embedded into a glass tube that ends in an ion selective glass membrane. On both sides of the glass membrane, a hydrated gel layer is formed with the aqueous solutions that are in contact with glass surfaces (Figure 1). The concentration of H<sup>+</sup> ions on the inner layer, containing a reference solution, is constant while on the outer layer it varies depending on pH. As a consequence, there is an exchange of alkaline ions between the outer layer and the glass membrane that changes the overall potential of

the membrane. The reference electrode is usually of the same type (Ag/AgCl), immersed into a KCl solution and can be included with the working electrode in a single device.

The glass electrode potentiometric equipment is relatively cheap and has been the only practical way to measure pH of seawater for many years. However, the glass electrodes must be handled with care due to the brittleness that is associated with glass, and properly stored in electrolyte solutions to prevent ions leaching from the glass membrane (if stored in deionized water) and to preserve the hydrated layer onto glass surface from drying out. They also have a limited shelf life due to the degradation of the glass membrane and need a regular calibration in seawater buffers, whose accurate preparation determines the accuracy of the measurement (McLaughlin et al., 2017b; Weldborg et al., 2009). The stability and pressure sensitivity of the “liquid junction”, the porous membrane that allows an ion flow to close the electrochemical cell, can also be an issue. In practice, electrode potential drift and experimental problems can limit the accuracy of potentiometric measurements to less than 0.01, with a drift of 0.02 pH/day (Rérolle et al., 2012).

### **3. Inorganic materials for solid state sensors**

The realization of a miniaturizable, stable and cheap pH sensor to substitute glass membrane based devices is still a challenge. A number of solid-state sensors have been proposed and some of them are already available on the market.

A common approach to solid state Nernstian pH sensors is based on the realization of Ion Sensitive Field Effect Transistors (ISFETs). ISFETs are traditional Metal Oxide Semiconductor Field Effect Transistors (MOSFET), where the gate electrode is modified (or substituted) by a thin layer of an insulating material ( $\text{Si}_3\text{N}_4$ ,  $\text{Al}_2\text{O}_3$ ,  $\text{Y}_2\text{O}_3$ ,  $\text{ZrO}_2$ ). The protonation/deprotonation process occurring on the insulator layer when in contact with water solutions of different pH determines the electrostatic field at the gate, controlling the current flowing into the FET (Bergveld, 2003). The circuit must be closed using a reference electrode connected to the source in lieu of the now removed gate (liquid gating).

ISFET pH sensors exploit a mature (more than 20 years) technology and have been used extensively for industrial, clinical and environmental pH monitoring as they offer a number of advantages, relative to glass electrodes. First, the sensor can be fabricated with conventional silicon based semiconductor technologies at reduced costs and ease of integration with electronic devices. Furthermore, it is small, resistant to mechanical shock and does not need a storage solution. Due to the different structure, the impedance of ISFET devices is lower with respect to glass electrodes, which has a beneficial effect on noise and stability. Commercially available sensors based on ISFET technology have been tested at sea with encouraging results and devices specifically designed for oceanographic research, mainly based on the Honeywell Durafet™ sensor, are currently used by research institutions (Johnson et al., 2016; Saba et al., 2019).

Despite the good performances of ISFET sensors, further refinements are required for their extended use in ocean acidification studies, concerning, as an example, the reliability of the reference electrode, long signal stabilization time and the stability of the sensor during long-term oceanic deployments (Martz et al., 2015; McLaughlin et al., 2017a; Rérolle et al., 2012).

A wide number of variations to the standard ISFET design have been proposed over the years. Some examples of the most advanced solutions will be reported here (see Table 1 for main parameters). A double gate architecture that can push sensitivity above the Nernst limit has been developed. As an example, a double gate ISFET based on ZnO was claimed, with a sensitivity as high as 2.25 V/pH (Spijkman et al., 2011a). The high sensitivity is generated by a capacitive coupling effect (Spijkman et al., 2011b) that, in this case, was maximized by applying an extremely thin passivation layer, a self assembled monolayer of octadecyl phosphonic acid. However, the device was tested only at the pH values of 6 and 8, and showed a large standard deviation in the measured potential. An alternative design to reduce noise and increase stability relies on the realization of an extended sensing layer, connected to the gate (extended-gate FET or EGFET; Pullano et al., 2018). Parizi et al. (2012) proposed a device that couples two EGFETs (n- and p- type) in parallel, matched to have the same transconductance to cancel a large part of the noise. This design allows the substitution of the external reference electrode (e.g. Ag/AgCl) with a simple, solid state pseudo-reference.

In a recent paper, Takechi et al. (2015) demonstrated a signal amplification effect similar to Spijkman (2011a) using an amorphous InGaZnO<sub>4</sub> (IGZO) layer as the bottom gate and a thin film of TaO<sub>x</sub> as an ion sensitive top gate. The resulting sensitivity is as high as 450 mV/pH but the resolution limit, calculated taking into account drift and hysteresis of the device, was estimated in 0.02 pH in a narrow range (pH 4 – 6). An optimization of the fabrication process led to a similar IGZO/Ta<sub>2</sub>O<sub>5</sub> based ISFET with a sensitivity of 402 mV/pH in the 4 – 9 pH range (Kumar et al., 2017). However, stability and drift problems still constitute a serious limit to the use of this kind of device in demanding applications (Pyo & Cho, 2017). Ta<sub>2</sub>O<sub>5</sub> has been investigated also for the realization of flexible extended gate electrodes, printed on plastics and coupled to a FET device (Wu et al., 2017). The sensitivity of this assembly was relatively low, 24 mV/pH, but good temporal stability (drift < 1% during tests) and repeatability were observed.

Recently, an interesting combination of organic semiconductor and SiO<sub>x</sub> thin layer was tested as gate in a dual-gate ISFET device, showing an improvement in response time and an amplification of the signal up to 10 times with respect to a bare SiO<sub>x</sub> layer (Pfattner et al., 2019). However, the stability of the response was not addressed.

In summary, ISFET devices take advantage of well-established semiconductor fabrication processes for the production and integration of pH sensors. It is a technology with a long history and a high maturity level, with at least one product dedicated to seawater application already on the market. In the quest for increased stability and accuracy, a number of improved designs have been proposed and tested at laboratory scale. Latest developments make use of nano-engineered active layers and electrodes and will be discussed in Section 5.

| Sensing material and setup | Testing range and media        | Sensitivity     | Stability       | Response time | Reference              |
|----------------------------|--------------------------------|-----------------|-----------------|---------------|------------------------|
| ZnO<br>Dual gate<br>ISFET  | 6 – 8<br>Commercial<br>buffers | Up to 2.25 V/pH | Low hysteresis. | n/a           | Spijkman et al., 2011a |

|                                                |                                                                                                   |                                                                          |                                                            |       |                       |
|------------------------------------------------|---------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------|------------------------------------------------------------|-------|-----------------------|
|                                                |                                                                                                   |                                                                          | High standard deviation on sensitivity estimation.         |       |                       |
| Al <sub>2</sub> O <sub>3</sub> EGFET           | 4 – 10                                                                                            | Up to 130 mV/pH                                                          | n/a                                                        | n/a   | Parizi et al., 2012   |
| TaO <sub>x</sub> Dual gate ISFET               | 4 – 6<br>McIlvaine buffer (Na <sub>2</sub> HPO <sub>4</sub> – citric acid)                        | 453 mV/pH<br>Resolution up to 0.02 pH                                    | Drift and hysteresis low with respect to the high response | n/a   | Takechi et al. 2015   |
| Ta <sub>2</sub> O <sub>5</sub> Dual gate ISFET | 4 – 9<br>Commercial pH buffers                                                                    | 402 mV/pH                                                                | Relatively stable after 1.5 years of storage               | n/a   | Kumar et al., 2017    |
| Ta <sub>2</sub> O <sub>5</sub> EGFET           | 1 – 13<br>Water + HCl/NaOH<br>Weak effect of monovalent cations                                   | 28 mV/pH                                                                 | <1% drift over 16 min                                      | <10 s | Wu et al., 2017       |
| SiO <sub>x</sub> ISFET                         | 2.4 – 11.7<br>PBS Response is not linear below pH 5.<br>Low effect of varying NaCl concentration. | Up to 14%/pH (drain current normalized to the reference value at pH 7.4) | n/a                                                        | Few s | Pfattner et al., 2019 |

**Table 1.** Main characteristics of ISFET based sensors

A second family of solid state probes for pH are electrodes based on oxides or metal/metal oxide couples, suitable for a potentiometric sensing setup. Metal/metal oxide pH sensors respond to pH due to an equilibrium involving the metal and its oxide where, in the metal oxide electrodes, the metal is not involved in the potential-determining reaction (Glab et al., 1989).

Due to their robustness, relatively easy miniaturization, fast response and good sensing performance, metal/metal oxide and metal oxide materials represent promising substitutes to glass electrodes. pH responsiveness has been observed in many semiconducting oxides, including Sb<sub>2</sub>O<sub>3</sub>, PtO<sub>2</sub>, OsO<sub>2</sub>, Ta<sub>2</sub>O<sub>5</sub>, TiO<sub>2</sub>, PdO, SnO<sub>2</sub>, ZrO<sub>2</sub>, PbO<sub>2</sub> and, notably, IrO<sub>2</sub> and RuO<sub>2</sub> (Hayat & Marty, 2014; Koncki & Mascini, 1997; Yao et al., 2001).

Antimony based electrodes have been among the first to be developed and proposed (Kinoshita et al., 1986). As the potential developed by antimony, in response to hydrogen ion activity, is to some degree sensitive to other dissolved anions, the use of a Nafion membrane to cover the electrode has been proposed, resulting in a response stable within 2 mV/pH over 1 month (Xu et al., 2016, 2018, see table 2).

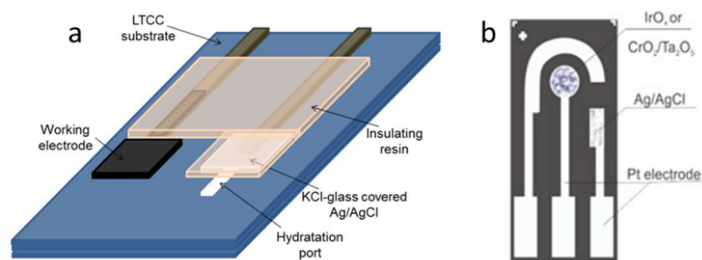
Ruthenium oxide is one of the most investigated oxides for pH sensors; its sensing mechanism is attributed to the presence of oxygen vacancies at the surface that lead to the formation of hydroxyl groups by dissociative adsorption of water, generating a pH sensitive layer (Trasatti, 1991). Thick films can be produced by screen printing and 3D structures can be built by the low temperature co-firing of ceramics (LTCC), both industrially scalable processes, showing very high sensitivity and robustness (Manjakkal et al., 2014, 2016; Figure 2). Thick films based on RuO<sub>2</sub> containing glass paste by screen printing and sintering; potential was measured against Ag/AgCl in the pH range 2-12 and a linear Nernstian behaviour was observed with a slope of 56 mV/pH. In a formulation with 30 wt% of titania, the sensitivity was maintained at 56.11 mV/pH with a good response time of about 15s and a

good 60 days stability. In a similar way, mixed  $\text{RuO}_2/\text{Ta}_2\text{O}_5$  based films were prepared by screen printing and sintering with glass forming oxides. In this case, the response was higher in acid to neutral environment (64.7 mV/pH from 2 to 8) than in basic conditions (43.1 mV/pH from 8 to 11) probably due to the effect of alkaline pH on the supporting glass paste (Manjakkal et al., 2016). Remarkably, in these examples the behaviour of  $\text{RuO}_2$  based sensors was not influenced by common anions. However, an influence of oxygen and redox agents has been observed in industrial applications of  $\text{RuO}_2$  based sensors. Recently, a double protective layer ( $\text{Ta}_2\text{O}_5$  thin film and Nafion membrane) has been introduced to mitigate the effect of interfering species (Lonsdale et al., 2018).

Some of the reported papers, in the quest for miniaturization and integration of their sensors, propose an integrated solid  $\text{Ag}/\text{AgCl}$  pseudo-reference electrode, to be fabricated into the same substrate as the sensing electrode. The need for a stable reference is, in fact, a key problem for the development of miniaturized solid pH sensors (Hu et al., 2015; Michalska, 2012). We can affirm that a robust and stable alternative to liquid or gel filled electrodes is not yet available, although a number of different designs for disposable and/or reusable solid pseudo-reference electrodes are available (Sophocleous & Atkinson, 2017). The use of modern fabrication technologies can lead to miniaturized multilayered electrodes with stability comparable to traditional ones (Moya et al., 2019).

Iridium oxide is also widely employed for pH sensing. It is usually indicated as  $\text{IrO}_x$  due to its complex stoichiometry, strongly influenced by synthesis conditions (Jang & Lee, 2020). An optimized electrodeposition, followed by an annealing procedure, was developed for the deposition of porous  $\text{IrO}_x$  films onto gold electrodes (Kim & Yang, 2014). The modified electrodes (Figure 2) showed a nearly perfect Nernstian response to pH changes, with a slope of 59 mV/pH very stable towards cyclic pH changes. Iridium and tantalum oxide thin films were deposited onto platinum electrodes by means of electro-deposition and e-beam sputtering respectively (Uria et al., 2016). Being designed for biological media, these devices were tested in a phosphate buffer saline solution (PBS) within a narrow pH range, resulting in a potentiometric response of 59.4 mV/pH for tantalum and 72 mV/pH for iridium oxide.

$\text{IrO}_x$  based sensors have good pH sensing performances but their response can be affected by reactions with oxidizing and reducing species dissolved in the test solution. A tantalum oxide layer deposited over  $\text{IrO}_x$  has been tested as a barrier layer, increasing the stability of the signal against oxygen (Kuo et al., 2014). Recently, a further refinement in the oxidation procedure of iridium wires led to the production of a remarkably stable sensor, with no need for barrier layers (Pan et al., 2018). This sensor was tested in the presence of a large set of anions and cations and in marine water, exhibiting stability and sensing performances in line with the glass electrode used as a reference. A very similar  $\text{Ir}/\text{Ir}(\text{OH})_x$  pH electrode has been recently fabricated and field tested in seawater, comparing the results with a commercial pH meter (Zhang et al., 2017). The solid state electrode showed good stability (137 days) and a precision comparable to the reference glass sensor, with a life span up to 5 months.



**Figure 2.** Assembly of: sintered  $\text{RuO}_2$  working electrode onto LTCC ceramic substrate (a) (Reprinted from Manjakkal et al. (2016), with permission from Elsevier); gold electrode modified by electrodeposited  $\text{IrO}_x$  (b) (Reprinted from Uria et al. (2016), with permission from Elsevier). Both solutions include a pseudo-reference solid  $\text{Ag/AgCl}$  electrode.

Other metal oxides used for pH measurements include  $\text{WO}_3$ ,  $\text{TiO}_2$ ,  $\text{ErO}_2$  and  $\text{MnO}_2$ . Manganese oxide was shown to exhibit a non-linear electrical response to pH (and a tendency to dissolve in acidic solutions) due to the chemical equilibrium among the oxide and the oxo-hydroxide species. A microelectrode was fabricated by coating  $\text{MnO}_2$  with a polymeric proton-conductive Nafion membrane, showing a linear response in the 4 – 12 pH range with a slope of 60 mV/pH (Cachet-Vivier et al., 2010). In a recent study, a tungsten bronze with a well-defined composition and crystal structure was produced by oxidation of tungsten wire and has been proposed as electrode material for potentiometric pH detection (Cisternas et al., 2017). The response of this material was found to be highly reproducible and stable (variations in the order of 0.3 mV) upon storage and continuous operation conditions. The use of multiple metals in a single device has been investigated by Sadig et al. (2018) that realized an iridium, ruthenium and titanium oxide based tri-oxide system. Though no details are given on the structure of the deposited oxide layer, the response recorded showed a linear potential/pH relation whose slope was stable within 0.3 mV over 120 days of testing. Finally, it is worth reporting on the design of a sensor based on solid metal rods, expressly developed for seawater monitoring (Brooke et al., 2016). To overcome the interferences of corrosion, surface reactions and fouling, 8 different metals were simultaneously used and their potential against a common zinc counter electrode was recorded continuously against pH, measured by a reference pH meter, allowing the calibration of the device through a self-learning neural network algorithm. After calibration, the device was able to reproduce actual pH values over 3 weeks of deployment.

| Sensing material and setup                               | Testing range and media                                                                                    | Sensitivity         | Stability                                                                | Response time | Reference              |
|----------------------------------------------------------|------------------------------------------------------------------------------------------------------------|---------------------|--------------------------------------------------------------------------|---------------|------------------------|
| $\text{Sb}_2\text{O}_3$ – Nafion membrane Potentiometric | 4 – 9<br>Commercial buffers                                                                                | $54.5 \pm 2$ mV/pH  | Stable within 2 mV/pH over 1 month (measurement repeated every week)     | 20 s          | Xu et al., 2018        |
| $\text{RuO}_2/\text{TiO}_2$ Potentiometric               | 2 – 12<br>HCl/NaOH solutions.<br>Interference of $\text{Li}^+$ , $\text{Na}^+$ and $\text{K}^+$ negligible | 56.11 mV/pH         | Storage in ambient condition up to 2 months with no change in properties | 15 s          | Manjakkal et al., 2014 |
| $\text{RuO}_2/\text{Ta}_2\text{O}_5$ Potentiometric      | 2 – 12<br>HCl/NaOH solutions and $\text{H}_3\text{BO}_3$ /citric                                           | 64.7 mV/pH (pH 2–8) | Storage in ambient conditions up to 2 months led to a                    | 15 s          | Manjakkal et al., 2016 |

|                                                                                                    |                                                                                                                                                          |                                                                       |                                                                                                 |                             |                                                |
|----------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------|-------------------------------------------------------------------------------------------------|-----------------------------|------------------------------------------------|
|                                                                                                    | acid/ $\text{Na}_3\text{PO}_4$ buffer. Interference of $\text{Li}^+$ , $\text{Na}^+$ and $\text{K}^+$ negligible                                         | 43.1 mV/pH (pH 8–11)                                                  | small reduction in sensitivity.                                                                 |                             |                                                |
| $\text{IrO}_x$ Potentiometric                                                                      | 2.4 – 11.6 Commercial buffers                                                                                                                            | 59.5 mV/pH                                                            | n/a                                                                                             | 2 s                         | Kim & Yang, 2014                               |
| $\text{Ta}_2\text{O}_5$ / $\text{IrO}_x$ Potentiometric                                            | 3 – 8 PBS acidified with $\text{HNO}_3$ Chloride ion concentration can influence reference stability.                                                    | 59.4 mV/pH ( $\text{Ta}_2\text{O}_5$ )<br>72 mV/pH ( $\text{IrO}_x$ ) | Stable after incubation in LB/ glucose for 24h.                                                 | Few s                       | Uria et al., 2016                              |
| $\text{IrO}_x$ Potentiometric                                                                      | 2 – 13 Britton – Robinson buffer Good selectivity against common cations                                                                                 | 59.5 mV/pH                                                            | Drift < 0.1 mV/h                                                                                | n/a                         | Kuo et al., 2014                               |
| $\text{Ir(OH)}_x$ carbonate oxidized Potentiometric                                                | 2 – 10 Commercial buffers <b>Tested in seawater (pH 7.9)</b> Negligible effect of common cations anions and $\text{O}_2$                                 | 56.8 – 57.6 mV/pH                                                     | No drift over 48 h at pH 6.                                                                     | 1 s                         | Pan et al., 2018                               |
| $\text{Ir(OH)}_x$ Potentiometric                                                                   | 4 – 9 Calibrated in commercial buffers <b>Tested in Dickinson seawater (pH 7.876) and in open sea</b>                                                    | 56.1 – 59.5 mV/pH                                                     | Negligible drift over 200s. Stable during 137 d of continuous recalibration in standard buffers | 5 s                         | Zhang et al., 2017                             |
| $\text{MnO}_2$ – Nafion membrane Potentiometric                                                    | 2 – 12 $\text{H}_2\text{SO}_4/\text{NaOH}$ solutions Interference by $\text{Fe}^{2+}$ ions.                                                              | $\approx 60$ mV/pH                                                    | n/a                                                                                             | 35 to 74 s                  | Cachet-Vivier et al., 2010                     |
| $\text{Na}_{0.75}\text{WO}_3$ Potentiometric                                                       | 1 – 10 Commercial buffers, $\text{KCl}/\text{HCl}$ solution (pH 1) High selectivity against $\text{Na}^+$ $\text{K}^+$ $\text{Mg}^{2+}$ $\text{Ca}^{2+}$ | $\approx 56$ mV/pH                                                    | Stable for storage in air up to 6 months and for repeated measurements over 1 w                 | 13 – 18 s (depending on pH) | Cisternas et al., 2015; Cisternas et al., 2017 |
| $\text{IrO}_2$ - $\text{RuO}_2$ - $\text{TiO}_2$ Potentiometric                                    | 1 – 13 Tris buffer Some influence of $\text{K}^+$ ions                                                                                                   | 59 mV/pH                                                              | Stable for 120 d                                                                                | 4 – 8 s                     | Sadig et al., 2018                             |
| Stainless Steel, Cu, WC, Brass, Ni, Al, Ti, Bronze Potentiometric vs a common Zn counter-electrode | <b>Tested in seawater</b>                                                                                                                                | Neural network calibration correlates potential readings with pH      | Signal degradation after 1 month of deployment                                                  | n/a                         | Brooke et al., 2016                            |

**Table 2.** Main characteristics of metal/metal oxide based sensors. In bold, sensors that have been tested in seawater.

Metal oxide based pH sensors are finding increasing popularity due to their ruggedness and relatively low price. However, most of the prospective applications are not demanding, in terms of accuracy and stability, like seawater monitoring.

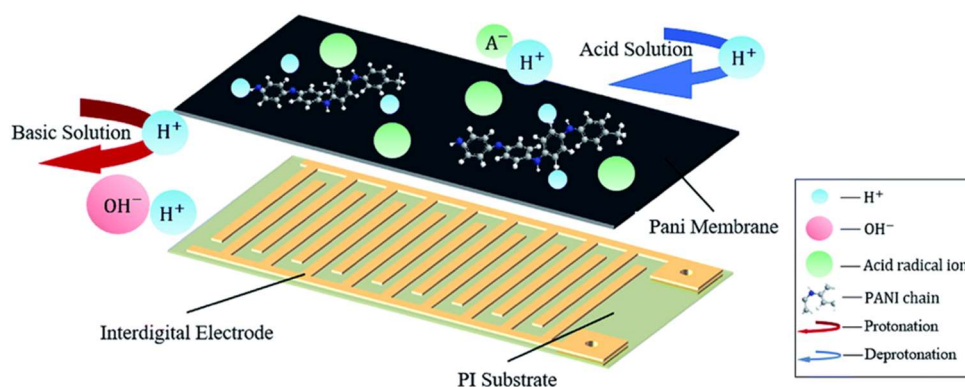
There are interesting examples of metal oxide sensors tested in seawater. Zhang et al. (2017) integrated four IrO<sub>x</sub> pH electrodes and one Ag/AgCl reference electrode in a self-made chemical sensor, and deployed it in a profile detection of nearly 70 m for a sea trial, near Newport Harbor, California. The pH value measured by the sensor was very close to the data given by a Sea-Bird 911 plus CTD, taken as a reference (maximum deviation 0.06 pH units), with the IrO<sub>x</sub> sensor showing a better data matching in the 0–40 m water depth range. The sensors were subjected to periodic calibrations for a 137 days period, showing a remarkable response stability. The authors contend that the high precision and accuracy of the sensor make it possible to use in the ocean observation field. Pan et al. (2018), fabricated an IrO<sub>x</sub> based electrode, whose response to pH was tested in various buffers and in seawater samples. Their sensors showed a good agreement (maximum deviation 0.04 pH units) with a commercial glass electrode in all testing conditions, showing a remarkable selectivity against interfering ions. No data is available for long term deployment in seawater. Furthermore, Brooke et al. (2016) described the simultaneous use of eight metal electrochemical cell for measuring ocean pH through a non-linear calibration algorithm obtained using a neural network self-learning approach. A prototype sensor was deployed in a seawater tank at the Seattle Aquarium for one month and, after the calibration period, was able to reproduce pH values within 0.02 pH units vs. the reference pH electrode for up to 3 weeks, before corrosion and fouling started to affect the response. The latest developments in the field of inorganic films, for both FET and potentiometric pH sensing devices, are directed towards the fabrication of nanostructured/multilayer electrodes with improved performances and reduced cost. These approaches will be treated in Section 5.

#### **4. Polymer-based pH sensors**

Polymer based materials, in particular conducting polymers, are finding ever increasing applications in the sensing field, due to their versatility, low cost and robustness (Adhikari & Majumdar, 2004). A general feature of conducting polymers is their “redox” activity and, as a consequence, the possibility to change their electrical behaviour (charge carrier density, band structure) through a doping-dedoping effect generated by the interaction with ions or small molecules (Adhikari & Majumdar, 2004; Culebras et al., 2014). These interactions constitute the basis for the use of conducting polymers for sensing (Gupta et al., 2004; Persaud & Pelosi, 1985). Huang et al. (1986) investigated in detail the effect of pH on conducting polymers, in particular polyaniline (PANI) and showed that the pH influences the redox processes of PANI in aqueous electrolytes. Since the first pioneering studies, the most investigated polymers in sensing have been polythiophene, polypyrrole (Ppy) and, notably, polyaniline and its derivatives, deposited or polymerized directly onto metal electrodes. Doped PANI can be produced by electrochemical polymerization of aniline in the presence of tetraphenyl borate (Pandey & Singh, 2001). The potentiometric measurement carried out in buffers and electrolytic solutions showed a linear potential/pH relationship, and a claimed stability of 6 months

(Table 3). However, a super-Nernstian response was observed, attributed to a non equilibrium protonation/deprotonation process. In a more complex design, a graphite lead was covered with in situ polymerized PANI (Gao & Song, 2009) and used for amperometric sensing of pH in the range 1.8 – 9.9. The voltammetric I/V curve shifted towards negative potential with increasing pH, showing a bilinear correlation and high reproducibility (0.5% error on repeated measurements). The higher slope recorded in the acidic range was attributed to multiple oxidation states possible for PANI. Recently, disposable and low cost sensors were realized by drop casting a PANI solution on carbon electrodes, printed on a paper substrate. Ag/AgCl solid pseudo-references were produced on the same substrate to fabricate an integrated device that showed a linear response to pH in the range 4 – 10, stable during 24h (Rahimi et al., 2016). Flexible interdigitated electrodes deposited on a polyimide film (Figure 3) have been covered by spin casting with a PANI film, doped with dodecyl benzene sulfonic acid (Li et al., 2020). The flexible sensors were calibrated in phosphate buffers and showed a linear potential response vs. pH up to pH 8.6.

Platinum electrodes, realized by photolithography, have been modified with polypyrrole and used by Lakard et al. (2007); the potentiometric response of these sensors was tested in the pH range 2 – 11, showing a nearly linear dependence of potential with pH. The sensitivity, however, showed a progressive decrease over 30 days of monitoring, attributed to the degradation of the silver pseudo-reference electrode. Ppy polymerized onto PEI modified electrodes showed improved stability, due to the adhesion granted by the imine layer (Segut et al., 2007). As a more recent example of a potentiometric sensor made by electropolymerization, it is worth mentioning the device proposed by (Li et al., 2011). By polymerization of bisphenol A (BPA) onto ITO glass, the authors developed an electrode that was tested in either potentiostatic or potentiometric setup, in a wide pH range (1 to 14) showing a sensitivity close to the Nernst limit and a reasonable stability of the response up to 12 days.



**Figure 3.** Schematic representation of an interdigitated gold electrode with deposited PANI sensing layer undergoing reversible protonation/deprotonation. Reproduced from Li et al. (2020) - Published by The Royal Society of Chemistry.

Recently, a non-conjugated, redox active polymer, poly(dopamine), demonstrated a linear correlation of the redox peak measured by voltammetry with pH. The polymer was deposited on a carbon electrode and tested in a wide pH range, in different buffers or saline solutions showing an excellent stability of the response (Amiri et al., 2016).

Combinations of conducting polymers with support polymers have been also realized by various methods, including the deposition of preformed polymer from solutions, reducing the cost of the assembly and overcoming the difficulties of electrodeposition. Gill et al. (2008) developed a composite conductimetric pH sensor mixing doped PANI particles with polyvinyl butyral and polypyrrole. The composite was deposited by screen printing on an interdigitated electrode and showed a linear response to pH in the range 2 – 8, but a response time of about 200 s. An analysis of the sensor response as a function of composition revealed that PANI is the active component while polypyrrole contributes to increase the system conductivity. As a development of this concept, a gel with similar composition was tested for the real time detection of pH in drinking water (Banna et al., 2014). Gold interdigitated electrodes were covered with the sensitive polymers and exposed to solutions in the pH range 6.5 – 9 showing a non-linear change in resistivity that was stable over 30 days of continuous exposure. The accuracy and resolution of this sensor were similar to commercial devices.

| Sensing material and setup                             | Testing range and media                                                                                                                | Sensitivity                                                            | Stability                                                                                                     | Response time        | Ref.                 |
|--------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------|----------------------|----------------------|
| PANI/<br>tetraphenylborate<br>Potentiometric           | 2 – 9<br>Tris-HCl buffer<br>Negligible effect of Na <sup>+</sup> , K <sup>+</sup> , Ca <sup>2+</sup>                                   | ≈ 86 mV/pH                                                             | Stable after 6 months storage                                                                                 | n/a                  | Pandey & Singh, 2001 |
| PANI<br>Amperometric                                   | 1.8 – 9.9<br>Britton-Robinson buffer                                                                                                   | 32.4 mA/pH (pH 1.8 – 5.5)<br>15.9 mA/pH (pH 5.5 – 9.9)                 | 0.5% error on consecutive measurements                                                                        | 5 s (85% of reading) | Gao & Song, 2009     |
| PANI<br>Potentiometric                                 | 4 – 10<br>Commercial buffers                                                                                                           | 50 mV/pH                                                               | Drift ≤ 0.01 pH/h during 24h                                                                                  | 12 s                 | Rahimi et al., 2016  |
| PANI<br>Potentiometric                                 | 5.45 – 8.62<br>Phosphate buffer                                                                                                        | 58.6 mV/pH<br>2.4% standard deviation                                  | Hysteresis < 12% of full scale                                                                                | 54 s                 | Li et al., 2020      |
| PEI / Ppy<br>Potentiometric                            | 4 – 9<br>Commercial buffers.<br>Interference of carbonate ions                                                                         | ≈ 50 mV/pH<br>Dependent on film structure                              | Slight decrease of sensitivity over 30 d                                                                      | <60 s                | Segut et al., 2007   |
| Poly(bisphenol A)<br>Potentiostatic/<br>Potentiometric | -1 – 15<br>50nM NaCl + HCl or NaOH.<br>No effect of Na <sup>+</sup> , K <sup>+</sup> , Cl <sup>-</sup> , SO <sub>4</sub> <sup>2-</sup> | 58.6 ± 1.4 mV/pH (Potentiostatic)<br>56.7 ± 1.6 mV/pH (Potentiometric) | Stable within ≈ 4% after 12 d of storage.                                                                     | 20 s                 | Li et al., 2011      |
| Poly(dopamine)<br>Potentiostatic                       | 1 – 12<br>Phosphate, acetate, carbonate, Britton-Robinson buffers.<br>HCl/KCl solution                                                 | 58.2 mV/pH                                                             | Stable within 0.8% for repeated measurement. Slight effect of buffer ionic strength, corrected by calibration | n/a                  | Amiri et al., 2016   |
| PANI/ Ppy in poly(vinyl butyral) gel<br>Conductometric | 6.4 – 9<br>Tap water                                                                                                                   | Non-linear resistivity/pH calibration curve.                           | Precision 0.07 pH units.<br>Stable for 30 d of continued use                                                  | n/a                  | Banna et al., 2014   |

**Table 3.** Main characteristics of polymer based sensors.

Polymer based pH sensors, mainly based on organic conductive polymers, have been known for a long time. Many different designs and compositions have been proposed, but their development has been limited up to now to lab scale studies. This fact can be due to the low compatibility of polymer processing conditions with the traditional electronic technologies that rely on inorganic semiconductors and oxides. Moreover, the relatively low stability of polymer electrical response may have contributed to the low diffusion of polymeric sensors for pH monitoring. Nevertheless, the popularity of polymer based sensors is now increasing, following the development of flexible, printable organic electronics, and polymers can be the ideal candidates for the fabrication of disposable devices with short service life. In the most recent researches, conductive polymers are combined with nanomaterials for enhanced sensitivity, response time and selectivity (Ates, 2013).

## **5. Nanomaterial-based sensors**

The continuous quest for high sensitivity, fast response time, flexibility and cost-effectiveness is the driving force for the research of new solutions and materials for sensing. The use of nanoscale materials, both organic and inorganic in the realization of sensing devices, has been recently proposed leading to very interesting improvements in sensor performances (Salavagione et al., 2014).

The first and more obvious consequence of the structuring at very small length scale, is the large increase in surface area. As the interactions with probe solutions are usually limited to the surface of the sensing material, this leads to an immediate increase in sensitivity that allows the design of miniaturized devices with weight, energy and cost savings.

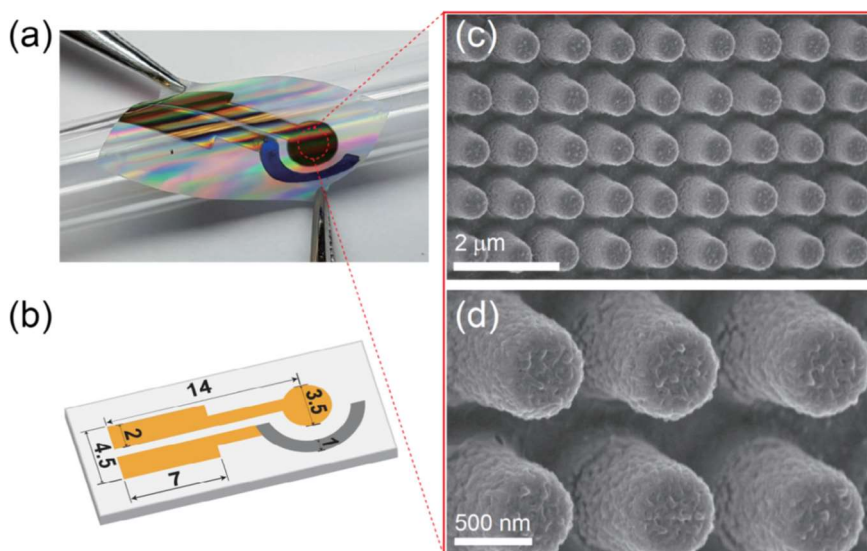
### **5.1 Organic and carbon-based nanomaterials**

One-dimensional nanomaterials based on conducting polymers can be fabricated using well-established wet chemical techniques and their properties can be easily tuned during synthesis or with a doping step. Nanotubes and nanowires with enhanced sensitivity toward various chemical/biological species are then ideal candidates for the design of new sensors (Bangar et al., 2010).

Nanowires fabricated with different methods have been proposed for the realization of pH sensing devices. Shirale et al. (2010) fabricated a FET sensor based on a single PPy nanowire for real-time pH monitoring and examined how the diameter of the nanowire affects the sensor performance. The sensor showed a linear correlation of the drain current with pH in the range 1 – 11 (Table 4). Doped Ppy nanowires were fabricated by electropolymerization on a gold substrate (Sulka et al., 2013). The gold/PPy electrode was then used as a potentiometric sensor in buffer solutions, in the pH range 2 – 12: it was shown that the oxidizing agent used for the polymerization influences the response, with  $\text{LiClO}_4$  giving the best sensitivity (49.3 mV/pH). Remarkably, the nanowires showed a ten-fold increase in sensitivity compared with thin films of Ppy prepared in the same conditions.

Recent developments in polymer based pH sensors are generally directed towards the realization of flexible devices, as an example, by printing carbon based electrodes onto plastic films and modifying them with active materials. PANI nanofibers directly polymerized at the surface of carbon electrodes supported on PET were tested at pH between 4 and 10 (Park et al., 2019). The Nernstian response

was observed with good repeatability (97.9%) and reasonable stability (drift of 3 mV/h over 15 h). An interesting flexible pH sensor was fabricated by soft-lithography templating of nanopillars on a polyurethane/acrylate layer followed by electrodeposition of polyaniline (Figure 4). A solid Ag/AgCl pseudo-reference electrode was deposited as a reference and the sensor was tested in the 2 – 12 pH range, showing a remarkably fast ( $\approx 1$  s) and accurate response (compared to a reference glass electrode) even in complex samples like juices and coffee (Yoon et al., 2017).



**Figure 4.** Templated nanopillars realized by soft lithography and flexible electrode assembly (Journal of Colloid and Interface Science 490 (2017) 53–58). Reprinted from Yoon et al. (2017), with permission from Elsevier.

Dodecyl benzene sulfonic acid doped PANI nanoparticles were conveniently incorporated into an epoxy resin to produce thin films for conductometric measurement of pH in soil (Patil et al., 2019). The films showed a high conductivity when loaded with 10 wt% of PANI and were tested in commercial buffers showing a good linearity of relative conductance vs. pH.

Carbon nanotubes (CNTs) and graphene (G) are among the most investigated nanomaterials for sensing applications, thanks to their unique chemical structure, very high conductivity, chemical stability and high surface area (Chen et al., 2011; Martin & Escarpa, 2014).

Ideal graphene (G) is a single layer of  $sp^2$  carbons arranged in a hexagonal structure extended in 2 dimensions (Li et al., 2009; Novoselov et al., 2012). Carbon nanotubes are tubular structures ideally formed by rolling up one (single-wall, SWCNT) or more (multi-wall, MWCNT) graphene sheets. The surface chemistry of carbon nanostructures can be tuned by the introduction of specific chemical groups, influencing their electronic and chemical behaviour (Ramanathan et al., 2008; Tasis et al., 2006). Graphene derivative materials known as graphene oxide (GO) and reduced graphene oxide (rGO) are interesting alternatives to graphene, showing higher reactivity at the expense of conductivity.

An interesting report on the correlation of CNT conductivity with pH was published in (Lei et al., 2012). The authors simply deposited a layer of multiwall CNTs onto filter paper and then showed a nice correlation of the system resistivity with pH of buffer solutions. Similarly, the pH response of graphene was observed on a simple resistive device, by deposition of exfoliated graphene onto a silicon wafer. Platinum electrodes were then deposited and the resistivity measured showed a linear correlation with

pH that was explained by an n- and p-doping effect induced by  $H^+$  and  $OH^-$  ions respectively (Lei et al., 2011). A number of studies show that the electrical response of graphene and CNTs exposed to aqueous electrolyte solutions depend on various interfering factors (pH, dissolved ions, substrate surface, Heller et al., 2010) and that the formation of charges at CNT or graphene surfaces is mainly driven by the presence of “defects” (Back & Shim, 2006), such as oxidized groups (Tan et al., 2013). This findings are in line with papers reporting a negligible sensitivity to pH for perfect, defect free graphene sheets (Fu et al., 2011). Summarizing, a consistent explanation of the pH response of carbon nanomaterials is still lacking.

Recently, ink-jet printing was used to deposit  $-COOH$  functionalized SWCNTs on glass and polymeric substrates, obtaining a potentiometric sensor. A linear response, with slope related to the number of layers, was recorded in the pH range 3 – 11 (Qin et al., 2016). Carbon nanotubes can also be integrated into traditional semiconductor-based electronics for the realization of transistor-like devices with sensing properties. An extended gate FET (EGFET) was realized with a CNT network (Chien et al., 2012) employed for both the contact electrode and the sensing membrane. The CNTs were first acid-oxidized and then irradiated with a laser beam to increase the defect concentration on their surface. This treatment resulted in a greater sensitivity (50.9 mV/pH) of the FET to pH and in a good linearity (Correlation coefficient  $R^2$ : 0.998) of the response.

Similarly, most graphene based sensors are, realized as transistors. Ohno et al. (2009) reported on the fabrication of a solution-gated FET (SGFET) made by a single layer of mechanically exfoliated graphene onto  $SiO_2$ /silicon substrate. The charge transport properties of the graphene layer depend on pH and a nearly linear correlation was found between the gate potential (measured at the Dirac point) and pH, with a sensitivity of approximately 30 mV/pH. For the same kind of device (Ohno et al., 2010), the authors analyzed the signal/noise parameters in a narrower pH range (5 – 8) and calculated a promising detection limit of 0.025. Using a different approach, few-layer graphene (thickness 1-2 or 3-4 layers) was grown epitaxially on silicon to realize a SGFET, tested in the pH range 2 – 12. Interestingly, a super-Nernstian sensitivity of 99 mV/pH was recorded, irrespective of the thickness (Ang et al., 2008). The authors performed impedance spectroscopy to rule out any external influence on the conduction behaviour of the device, demonstrating that only the adsorption of  $OH^- / H_3O^+$  species determines the properties.

One of the interesting advantages of carbon nanomaterials is the possibility to use conventional fabrication techniques to realize electronic devices and sensors on flexible substrates (Jung et al., 2014; Sharma & Ahn, 2013). Single wall nanotubes were employed for the fabrication of flexible FETs supported on polyethylene terephthalate (PET) films, using a layer-by-layer (LbL) approach. The film was obtained by LbL deposition of carboxylated SWCNT with two polyelectrolytes, to work as the gate electrode. The response of the FET was found to be dependent on pH, although in a non-linear way (Lee & Cui, 2010). Mailly-Giacchetti et al. (2013) transferred graphene layers, grown by CVD, onto poly(ethylene 2,6-naphthalenedicarboxylate) (PEN), silicon modified with octadecyltrichlorosilane (OTS) and  $SiO_2$ , to evaluate the influence of the substrate on sensing. Although the different devices showed different conductivities, the sensitivity to pH was around 22 mV/pH for all of them.

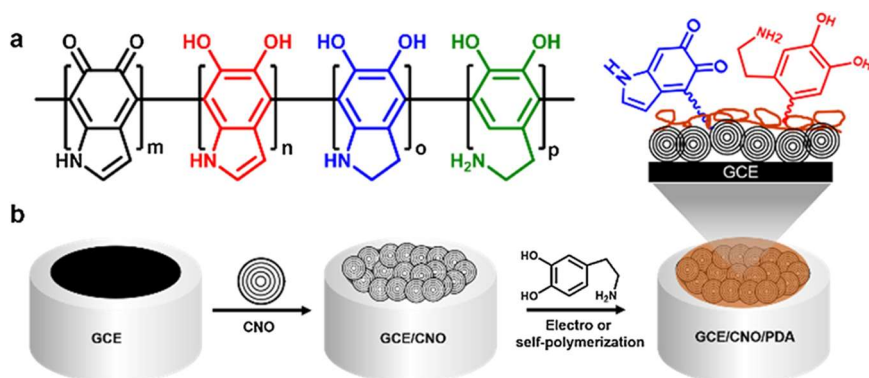
Some refinements in the design of graphene devices have been proposed to improve the sensing performances. A suspended graphene FET was fabricated with a claimed increase in the signal to noise ratio by 14 dB with respect to the same unsuspended device (Cheng et al., 2010). The increase in signal quality allowed measurements to be carried out with very low applied voltage, reducing the risk of interference from the testing solution (polarization, redox reactions). An interesting approach to increase the contact surface with the aqueous solution has been recently proposed by Ameri et al. (2016) with the realization of a high porosity graphene foam covered by a thin layer of  $\text{HfO}_2$ . This device was tested in Dulbecco phosphate buffer, showing a super Nernstian sensitivity of 71 mV/pH and a fast response. Finally, a solid gated G-FET designed to avoid the need for an external reference electrode has been reported. In this device, a layer of  $\text{HfO}_2$  is deposited between the graphene layer and a gold gate electrode (Zhu et al., 2015). The response of the FET was linear with pH in the range 5.3 – 9.1, with a sensitivity of 56.5 mV/pH.

GO and rGO materials can be used for the fabrication of membranes and networks, showing lower electrical properties compared to graphene but good pH sensitivity, probably due to the high concentration of oxidized groups (Sohn et al., 2013). A potentiometric GO based sensor has been realized for medical applications by printing the electrodes on a plastic substrate (Salvo et al., 2017) and calibrated in a buffer with isotonic salt concentration. The potential response to pH was linear and the sensors proved to be relatively stable for 1 week in serum. A sensor for seawater pH detection was derived from this system (Poma et al., 2019) and validated in high ionic strength buffers and real seawater. GO and rGO electrodes were coated with Nafion to increase stability and tested, with rGO (functionalized with 4-aminophenylacetic acid) showing the highest sensitivity. Stability was assessed for up to 8 days in seawater but the accuracy of the sensor was worse than the reference glass electrode.

Nanomaterials are often combined with polymeric substrates/matrices, for processing reasons and to enhance sensing performances by exploiting the synergism between the components. Synergistic effects can be observed in carbon nanomaterials combined with conducting polymers: the polymer increases the robustness and selectivity of the response; at the same time, the incorporation of nanoparticles improve the stability and the conductivity of polymers, enhancing the electric properties. Polyaniline is by far the most studied conducting polymer for the development of composites (Oueiny et al., 2014). Interesting results have been obtained by Loh et al. (2007), by combining CNTs with a conducting layer-by-layer thin film of PSS/PANI and employing the assembly in a resistive pH sensor. The electrical resistance showed a large shift upon pH change (pH 1 to 10), with a sensitivity of approximately 19.9  $\text{k}\Omega \text{ cm}^{-2}/\text{pH}$ . Similarly, Boeva et al. (2014) produced few-layer graphene and exfoliated (30 – 50 layers) graphite coated by PANI. The redox behaviour of these materials was investigated by cyclic voltammetry, showing that PANI nanocomposite preserve their electroactivity up to neutral pH due to interactions with the graphene, whereas neat PANI loses its conductivity above pH 3. A miniaturized pH meter based on amino-functionalized graphene/PANI nanocomposite was fabricated by electropolymerization on ITO/glass substrate and tested by voltammetric measurements in PBS buffer, resulting remarkably stable up to pH 11 (Su et al., 2016). Similarly, a good sensing performance was also recorded on polyaniline functionalized rGO, tested in both potentiometric and

resistive setup in the range 2 – 9. The PANI-rGO electrodes were coated with a Nafion film to decrease interference from other ions and tested in a L. Lactis fermentation reactor (Chinnathambi & Euverink, 2018). Recently, Grozdanov et al. (2018, 2019) have tested screen printed electrodes (SPE) modified with PANI/carbon nanotubes composites as pH nanosensors, in the frame of FP7 project COMMON SENSE (Cleary et al., 2014; Barton et al., 2016; Ribotti et al., 2015). Nanosensors were prepared by electropolymerization and exhibited a high value of conductivity, which was attributed to the synergistic effect of the conductive polymer and carbon nanostructure via  $\pi$ - $\pi$  stacking. Conductivity changes were measured at different pH (4 to 10) in commercial buffers, as well as in seawater samples showing a non linear response to pH. Similar electrodes were produced by Bao et al. (2019), who produced a PANI/MWCNT ink for screen printing of miniaturized working electrodes. Here, the response was measured by chronoamperometry, observing a linear relationship of potential vs. pH in the range 2 – 11; the role of nanotube/PANI interactions in the enhancement of the electric response was pointed out. Amperometric pH sensors were produced (Sha et al., 2017) by electropolymerization of well-ordered PANI chains on graphene-modified carbon electrodes, showing a nearly linear response to pH. The sensitivity was higher in alkaline solutions, which is rarely observed for PANI due to the dependence of electroactivity on acid doping.

A number of other polymers have been used for nanocomposite sensors fabrication. Gou et al. (2014). deposited a layer of oxidized SWCNTs between gold electrodes onto a silicon substrate, then poly(1-amino anthracene, PAA) was electropolymerized onto CNTs. The obtained device was tested in either liquid gated FET or conductometric configuration in various pH, showing high sensitivity and stability of the response over long time. The detection limit (resolution) is 0.04 pH. Recently, a biomimetic polymer, polydopamine (PDA), has also shown redox properties (Amiri et al., 2016) and, thanks to its excellent adhesion properties, has been used to modify nanostructured carbon electrodes (Figure 5; Zuaznabar-Gardona & Frago, 2018). PDA response was investigated by both cyclic voltammetry and potentiometry, showing a higher sensitivity when combined with carbon nanostructures, up to 53 mV/pH. The electrodes were stable for several months in water and only attacked by strong alkaline solutions. They were also tested in seawater showing a very good agreement with the reference pH meter.



**Figure 5.** General structure of polydopamine (a) and schematic fabrication of nano-onions (CNO) and polydopamine (PDA) deposition on glassy carbon electrodes (GCE) (b). Reprinted from Zuaznabar-Gardona and Frago (2018), with permission from Elsevier.

As a further example of polymer/nanoparticle synergism, it is worth mentioning the design proposed by Crespo et al. (2009). An acrylic ion selective membrane, doped to increase the selectivity towards  $H^+$  ions, was casted on a MWCNT-modified carbon electrode and tested for potentiometric pH measurements. Nanotubes here are introduced as a solid contact between the polymeric membrane, exhibiting pH dependent ionic conduction, and the working electrode. The result is a sensor with Nernstian response and good selectivity. The design of this kind of sensor has been refined over the years leading to the fabrication of a complete apparatus for field testing campaigns in freshwater (Athavale et al., 2017) and, notably, in seawater (Cuartero et al., 2017) showing performances comparable with commercial sensors.

| Sensing material and setup                           | Testing range and media                                                                                              | Sensitivity                                      | Stability                                          | Response time    | Reference                     |
|------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------|--------------------------------------------------|----------------------------------------------------|------------------|-------------------------------|
| Polypyrrole NW ISFET                                 | 1 – 11<br>Commercial buffers                                                                                         | 0.4/pH<br>(normalized drain current)             | n/a                                                | Few s            | Shirale et al., 2010          |
| Polypyrrole NW array Potentiometric                  | 2 – 12                                                                                                               | Up to 49.3 mV/pH                                 | Stable up to 50 d of storage                       | n/a              | Sulka et al., 2013            |
| PANI nanofibers Potentiometric                       | 3 – 10<br>Commercial buffers<br>Good selectivity against common cations                                              | 62.4 mV/pH                                       | Drift of 3 mV/h over 15h                           | 12.8 s           | Park et al., 2019             |
| PANI nanopillars Potentiometric                      | 2 – 12<br>Commercial buffers<br>Negligible effects of $Na^+$ , $K^+$ , $NH_4^+$ , $Ca^{2+}$ , and $Mg^{2+}$ at 10 mM | 60.3 mV/pH                                       | Low drift (0.64 mV/h) during a 15h test            | 5 s              | Yoon et al., 2017             |
| PANI particles in epoxy Resistive                    | 2.4 – 10<br>Commercial buffers                                                                                       | 957 $\mu$ S/pH                                   | Stable within 4% for 30 d (tested every 5 d)       | 5 – 30 s         | Patil et al., 2019            |
| SWCNT-COOH Potentiometric                            | 3 – 11<br>Britton-Robinson buffer                                                                                    | 48.1 $\pm$ 0.4 mV/pH                             | Response is reduced by 4% after 15 d of storage    | 7 s              | Qin et al., 2016              |
| MWCNT-COOH EGFET                                     | 3 – 13<br>PBS                                                                                                        | 50.9 mV/pH                                       | n/a                                                | n/a              | Chien et al., 2012            |
| Single graphene layer on $SiO_2$ Solution gated FET  | 4 – 8.2<br>10 mM phthalate, phosphate and borate buffers                                                             | Approx. 30 mV/pH<br>Resolution is 0.025 pH units | n/a                                                | n/a              | Ohno et al., 2009, 2010       |
| Single graphene layer on PEN Solution gated FET      | 4 – 9<br>Phosphate buffer adjusted with strong acids/alkali                                                          | 22 mV/pH                                         | Response decrease after exposure to acid solutions | $\approx$ 10 min | Maily-Giacchetti et al., 2013 |
| Graphene foam coated with $HfO_2$ Solution gated FET | 3 – 9<br>Dulbecco buffer adjusted with strong acids/alkali                                                           | 71 $\pm$ 7 mV/pH                                 | n/a                                                | $\leq$ 240 s     | Ameri et al., 2016            |
| Graphene + $HfO_2$ dielectric layer FET              | 5.3 – 9.2<br>Phosphate saline buffer                                                                                 | 56.5 mV/pH                                       | n/a                                                | < 60 s           | Zhu et al., 2015              |

|                                                              |                                                                                                                                                          |                                                                                                                     |                                                                                                             |         |                                                       |
|--------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------|---------|-------------------------------------------------------|
| rGO<br>Solution gated FET                                    | 6 – 9<br>Phosphate<br>buffer                                                                                                                             | 29.2 mV/pH                                                                                                          | n/a                                                                                                         | Few s   | Sohn et al.,<br>2013                                  |
| GO<br>Potentiometric                                         | 4 – 10<br>Citrate, borate,<br>phosphate<br>buffers<br>(isotonic)<br>Tested in<br>human serum                                                             | 40 ± 4<br>mV/pH                                                                                                     | Negligible drift<br>over 1 h in buffers.<br>Stable for 1 w in<br>serum                                      | n/a     | Salvo et al.,<br>2017                                 |
| Reduced GO + Nafion<br>membrane<br>Potentiometric            | 4 – 10<br>Citrate, borate,<br>phosphate<br><b>Tested in<br/>superficial<br/>seawater.</b>                                                                | 45 mV/pH                                                                                                            | Fluctuations<br>observed in the<br>response over 56h<br>in buffers. Signal<br>stable for 8 d in<br>seawater | n/a     | Poma et al.,<br>2019                                  |
| Graphene platelets<br>Resistive                              | 1 – 10<br>Commercial<br>buffers                                                                                                                          | 19.9 (k $\Omega$ /<br>cm <sup>2</sup> )/pH<br>Poor<br>linearity                                                     | n/a                                                                                                         | n/a     | Loh et al.,<br>2007                                   |
| Amino functionalized<br>graphene / PANI<br>Voltammetry       | 1 – 11<br>PBS                                                                                                                                            | 51.1 mV/pH                                                                                                          | Response<br>decrease after 1 w                                                                              | n/a     | Su et al., 2016                                       |
| rGO/PANI + Nafion<br>membrane<br>Potentiometric<br>Resistive | 2 – 9<br>Britton-<br>Robinson buffer<br>Also tested in a<br>bacterial<br>fermentation<br>broth                                                           | 55 mV/pH<br>1.71 $\Omega$ /pH                                                                                       | n/a                                                                                                         | n/a     | Chinnathambi<br>& Euverink,<br>2018                   |
| PANI-MWCNT<br>Chronoamperometry                              | 2 – 11<br>Water adjusted<br>with HCl/NaOH                                                                                                                | 20.6 mV/pH                                                                                                          | n/a                                                                                                         | Few s   | Bao et al.,<br>2019                                   |
| PANI on graphene-<br>carbon electrode<br>Amperometric        | 1 – 11<br>NaCl solution<br>adjusted with<br>HCl, H <sub>3</sub> PO <sub>4</sub> ,<br>NaOH                                                                | 50.17 $\mu$ A<br>/(pH cm <sup>2</sup> ) for<br>pH 1 – 5<br>139.2 $\mu$ A<br>/(pH cm <sup>2</sup> ) for<br>pH 7 – 11 | n/a                                                                                                         | ≈ 100 s | Sha et al.,<br>2017                                   |
| SWCNT – PAA<br>Solution gated FET /<br>Conductometric        | 2 – 12<br>Britton-<br>Robinson<br>buffers<br>Response to<br>Na <sup>+</sup> and Ca <sup>+</sup><br>negligible                                            | 0.073<br>mS/pH                                                                                                      | Stable for ≈ 2 h<br>(no drift). Same<br>sensitivity after<br>120 d of storage.                              | ≈ 60 s  | Gou et al.<br>2014                                    |
| Polydopamine/carbon<br>nano-onions<br>Potentiometric         | 1.5 – 10.5<br>Universal buffer<br><b>Tested in<br/>seawater (pH<br/>8.3)</b><br>Low<br>interference of<br>alkaline cations                               | 53 mV/pH                                                                                                            | Stable over 4 w                                                                                             | 15 s    | Zuaznabar-<br>Gardona &<br>Fragoso, 2018              |
| MWCNT solid contact<br>+ acrylic membrane<br>Potentiometric  | 3 – 10<br>Various buffers<br>Lake freshwater<br><b>Tested in<br/>seawater (pH<br/>7.9 – 8)</b><br>Selective<br>against alkali<br>cations and<br>sulfides | 58.8 ± 0.4<br>mV/pH                                                                                                 | Estimated drift 0.1<br>mV/h.                                                                                | 10 s    | Athavale et al.,<br>2017;<br>Cuartero et al.,<br>2017 |

**Table 4.** pH sensors based on nanostructured polymers, carbon nanomaterials and their combination. In bold, sensors that have been tested in seawater.

Three of the pH sensors listed in Table 4 have been recently tested in seawater, while most of the published works on sensors based on carbon-based nanomaterials only tested the sensors in buffered solutions. The first consists of a graphene-based pH sensor, part of an autonomous system for the remote monitoring of pH and temperature at sea (Poma et al., 2019); the pH measurement is performed through a potentiometric sensor with a wireless, smartphone-based real time acquisition system. The pH sensor was initially validated in the laboratory at controlled temperatures and in water previously collected at sea. Then it was left at sea with a sampling rate of one measurement per hour for 8 days. In both cases, a commercial glass electrode pH-meter was used as a reference device. Laboratory and on-field results have shown the great versatility of such a low cost system, providing pH values comparable with commercial sensors but with a lower energy consumption and a greater calibration stability. The second pH sensor is again potentiometric like the first one but based on polydopamine (PDA) films coated on a carbon nano-onion conductive surface. Also in this case, the new pH sensor was validated through comparison with a commercial combined glass pH electrode coupled to a pH meter from the same builder in water sampled at sea. They showed an excellent correspondence between these new PDA pH sensors and commercial ones with the advantages of an easy fabrication, an excellent reproducibility, a stability of the PDA coating in water over several months and the possibility of its integration into miniaturized devices. Both the potentiometric pH sensors described above must be tested in the field for longer times in order to verify stability and the long term effects e.g. of biofouling on system performances. The last example of a potentiometric sensor successfully tested in freshwater (Athavale et al., 2017) and in seawater (Cuartero et al., 2017), is based on an acrylic ion selective membrane with a carbon nanotube solid contact layer. For tests in seawater, the sensor was deployed in different coastal marine environments: Arcachon Bay on the Atlantic French coast for 14 hours, Genoa harbor on the Italian Mediterranean coast for 58 and 167 hours, and a mix sea-freshwater effluent, the Eyre River, in the Arcachon Bay during high tides for 14 hours. In all these tests the sensor showed good agreement with a reference glass electrode. This was particularly evident during the tests inside the harbour of Genoa where the sensor was compared with that mounted on a commercial multiparametric or Conductivity Temperature Depth (CTD) probe.

## **5.2 Semiconductor and metal/metal oxide nanomaterials**

The most traditional of semiconductor materials, silicon, has found new interesting applications in sensing with the development of Si nanowires (NW). Nanostructures with high packing density and tailored spacing can be fabricated by electron beam lithography on a silicon-on-insulator substrate with high accuracy and reproducibility (Bedner et al., 2013; Park et al., 2010). Choi et al. (2012) produced NWs on boron-doped silicon and deposited a protective layer of  $\text{Si}_3\text{N}_4$  to ensure better stability. The resistivity of the NW was measured as a function of pH and both short-time noise and long-term drift were measured (Choi et al., 2012). The pH sensitivity of this NW based device has been attributed to charge accumulation at the surface that induces a change in carrier density into the

high surface area wires, affecting conductivity. Recently Kim et al. (2014) produced As-doped suspended NWs by a lithographic approach. A linear correlation between normalized conductance and pH was found in the range 4 – 8, with a slope of 0.3 that is twice the slope of non-suspended nanowires (Table 5). The sensitivity was found to exceed the theoretical Nernst limit and, depending on the working current chosen, varied between 87 and 103 mV/pH (Salaün et al., 2014). This unexpected behaviour was observed also in double-gate NW transistors (Ahn et al., 2013) and rationalized taking into account the capacitance of the gates themselves (Knopfmacher et al., 2010). Finally, it is worth reporting a different application of Si NW, grown as a dense array to modify the gate of a FET device. The wires were sputtered with indium-gallium-zinc oxide (IGZO) resulting in a sensitivity of 50 mV/pH (Lin et al., 2013) at a working current of 200  $\mu$ A.

Metal oxide nanostructures can be produced by means of various fabrication techniques, exhibiting interesting electrochemical properties. An example is nanometric sulfated iron oxide (Alizadeh & Jamshidi, 2015). The particles produced by sol-gel were supported with a carbon paste and heat treated at 600°C to produce a regular crystal structure. With an optimized structure, the sensitivity was 57.5 mV/pH and a stability of 1 week was observed, providing the electrode is stored in water or immersed for a few hours in water after storage in dry conditions. Many other semiconducting metal oxides with very interesting properties can be shaped into nanometric wires, ribbons or tubes by different techniques and, interestingly, they can be easily integrated with well established silicon technologies. Titanium and zinc oxide nanotubes/wires are probably the most tested nanomaterials for pH sensing. Both materials show an amphoteric behaviour and can be used in both acidic and alkaline media; the active sites for sensing are oxygen vacancies found at the surface of the oxide structures. Titania nanotubes (NT) with lengths ranging from 33 to 800 nm were produced by anodization of a titanium electrode and embedded in PDMS for testing (Zhao et al., 2010). A nearly Nernstian behaviour was recorded for the nanotube modified electrodes, with best sensitivity and linearity obtained with amorphous titania. Materials prepared in different anodization conditions to produce a dense and thick nanotube layer onto titanium electrodes showed that the production parameters can affect the potentiometric response vs. pH (Albertin et al., 2013). To increase chemical stability, titania can be converted to nitride (TiN), producing a dense array of NTs onto platinum electrodes (Liu et al., 2016). TiN showed a higher pH sensitivity with respect to TiO<sub>2</sub>, excellent reproducibility and a good stability over 1 month of storage.

Fulati et al. (2009) produced zinc oxide nanotubes and wires, growing them onto gold substrates from a zinc nitrate solution. The response at different pH was measured showing a higher sensitivity for the NTs, explained in terms of higher surface area, and stability of the signal over several days. ZnO nanostructures are increasingly investigated for miniaturized devices (Kumar et al., 2019) and in particular for medical applications (Young & Tang, 2019). A linear response was recorded in the pH range 2 – 12 with aluminium-doped zinc oxide nanosheets (Tsai et al., 2019), tested as the gate layer of an ISFET.

A large number of nanostructured oxides have been exploited for pH sensing in different configurations, with a recent trend towards the realization of low cost, flexible devices. High

crystallinity tin oxide ( $\text{SnO}_2$ ) nanorods have been produced by a low temperature process onto conductive ITO glass (Li et al., 2012), and this layer was employed as a sensitive gate in an EGFET device. The sensitivity was increased with respect to thin film devices in both the linear and saturation regions of the transistor. Moreover, this device showed low hysteresis and no signal degradation during many hours of operation. Ruthenium oxide nanoparticles, deposited on a plastic supported electrode, have been tested as pH sensitive material in an EGFET configuration (Singh et al., 2019). The device showed a super-Nernstian behaviour, not usually observed for this oxide and a stabilization of the observed drift after 8 hours. Nanostructured platinum electrodes were realized by ink-jet printing onto a plastic substrate by Zea et al. (2019) and modified by electrodeposition of a thin  $\text{IrO}_x$  amorphous layer. The flexible devices were tested in the pH range 2 – 11 and aged in both dry and wet conditions over 1 year, showing excellent stability.

Tungsten oxide ( $\text{WO}_3$ ) is gaining increasing attention as a pH sensitive material. A  $\text{WO}_3$  layer was deposited by Zhang and Xu (2009) on a nanostructured electrode composed by aligned CNTs, obtaining a sort of nanopillar. Such modified electrodes showed a sensitivity of about 41 mV/pH, a low drift rate and a good stability after 1 month of storage. Tungsten oxide nanoparticles deposited onto a flexible, plastic supported electrode, showed a linear potential response to pH in the range 5 – 9 (Santos et al., 2014). A reduction of the sensitivity was however observed with continuous operation at different pH over  $\approx 1$ h. The same material was deposited onto glassy carbon to realize a sensor for voltammetric measurement of pH (Jamal et al., 2019), obtaining a high sensitivity (60 mV/pH) and linearity of the response. Drift was observed during the initial hours of sensor testing, but the signal stabilized thereafter remaining stable for up to 7 days. Recently, Choi et al. (2019), reported a new type of potentiometric pH sensor based on 1D tungsten oxide nanofibers with an amplified signal exceeding the Nernstian limit. Nanofibers with high porosity were synthesized and stabilized in a chloromethylated triptycene poly (ether sulfone) matrix, allowing a fast proton diffusion into the composite membrane. A high pH sensitivity of -377.5 mV/pH was obtained with the amplified sensor, linearity was acceptable in a narrow pH range (6.9 – 8.9). Testing in artificial seawater demonstrated a negligible effect of dissolved ions.

The advantages of nano-scale dimensions can also be exploited, in combination with organic support and/or ion-selective layers, for the realization of multicomponent sensing systems. An ISFET was realized by LbL deposition, using poly(diallyl dimethylammonium) (PDDA) and poly(styrene sulfonate) (PSS) embedding alternate layers of silica and  $\text{In}_2\text{O}_3$  nanoparticles (Liu & Cui, 2007). The semiconducting indium oxide granted a sufficient conductivity to the device while the protonation/deprotonation of  $\text{SiO}_2$  is responsible for pH sensing. A parabolic dependence of current vs. pH was recorded, with higher sensitivity in acid solutions. A development of this concept led to the realization of reliable and sensitive pH sensors based on the LbL assembly of iridium oxide nanoparticles and PDDA. The sensors produced showed a fast response and excellent reproducibility, by using a very low amount of iridium, paving the way for low cost, robust disposable sensors (Jović et al., 2018). Another example of synergistic combination of conducting polymer and nanoparticles was proposed by Kim et al. (2016), who developed a poly(terthiophene benzoic acid) (pTBA) / nanostructured  $\text{AuZnO}_x$  composite for disposable, solid state pH sensors. These devices were calibrated in the range

696 2 – 12 and showed fast response and stability when tested in biological samples. Lenar et al. (2019)  
697 recently proposed RuO<sub>2</sub> nanoparticles showing low resistivity, high stability and redox behaviour, as a  
698 solid contact layer between a carbon electrode and a modified PVC-based H<sup>+</sup> selective membrane.  
699 The assembly showed a fast, Nernstian response, largely due to the performance of oxide  
700 nanoparticles in synergy with the selectivity provided by the polymeric membrane.

| Sensing material and setup                                           | Testing range and media                                                                                                                                                                                                                                             | Sensitivity                                                                | Stability                                                                                                              | Response time | Ref.                      |
|----------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------|---------------|---------------------------|
| Si nanowire + Si <sub>3</sub> N <sub>4</sub> passivation layer ISFET | 4 – 9<br>HCl and KOH solutions                                                                                                                                                                                                                                      | 5.4%/pH                                                                    | Maximum drift of 1.68% at pH = 9                                                                                       | n/a           | Choi et al., 2012         |
| Si suspended nanowire ISFET                                          | 4 – 8<br>PBS buffer adjusted with HCl and NaOH                                                                                                                                                                                                                      | 0.3/pH<br>(expressed as relative conductance $\Delta G/G_0$ )              | n/a                                                                                                                    | n/a           | Kim et al., 2014          |
| Polycrystalline Si nanowire ISFET                                    | 4 – 9.2                                                                                                                                                                                                                                                             | Up to 103 mV/pH                                                            | n/a                                                                                                                    | n/a           | Salaün et al., 2014       |
| Si NW Double gated FET                                               | 4 – 10<br>PBS buffer                                                                                                                                                                                                                                                | 69 mV/pH                                                                   | Drift of 27 mV/h                                                                                                       | n/a           | Ahn et al., 2013          |
| Si NW sputtered with IGZO ISFET                                      | 2 – 10                                                                                                                                                                                                                                                              | 50 mV/pH                                                                   | n/a                                                                                                                    | Few seconds   | Lin et al., 2013          |
| Si QD EGFET                                                          | 2 – 12<br>Commercial buffers                                                                                                                                                                                                                                        | 108.3 mV/pH (linearity 98.97%)<br>2.65 $\mu A^{1/2}/pH$ (linearity 99.81%) | Hysteresis of 14 mV cycling pH from 4 to 10.                                                                           | ≈ 100 s       | Slewa et al., 2019        |
| Fe <sub>2</sub> O <sub>3</sub> nanoparticles Potentiometric          | 1.5 – 12.5<br>Negligible influence of common cations                                                                                                                                                                                                                | 57.5 mV/pH<br>Hysteresis effects ≤ 6%                                      | Stable for > 1 w of storage. Surface can be renewed by rubbing with paper. Reconditioning at pH=7 for few h is needed. | ≈ 10 s        | Alizadeh & Jamshidi, 2015 |
| TiO <sub>2</sub> nanotubes Potentiometric                            | 2 – 12<br>Britton-Robinson buffer Negligible interference of common ions (Na <sup>+</sup> , K <sup>+</sup> , Cl <sup>-</sup> , NO <sub>3</sub> <sup>-</sup> , SO <sub>4</sub> <sup>2-</sup> , F <sup>-</sup> , I <sup>-</sup> , Fe(CN) <sub>6</sub> <sup>4-</sup> ) | Up to 59 mV/pH (54 mV/pH before UV irradiation)                            | n/a                                                                                                                    |               | Zhao et al., 2010         |
| TiN nanotubes Potentiometric                                         | 2 – 11<br>Britton-Robinson buffer<br>Low effect of monovalent cations and F <sup>-</sup>                                                                                                                                                                            | 55.3 mV/pH                                                                 | Negligible drift over 200s. Stable after 1 month storage.                                                              | 4.4 s         | Liu et al., 2016          |
| ZnO nanotubes Potentiometric                                         | 4 – 12<br>Commercial buffers<br>Response influenced by CaCl <sub>2</sub>                                                                                                                                                                                            | 45.9 mV/pH                                                                 | Up to 5 d (tested at day 0, 2 ad 5).                                                                                   |               | Fulati et al., 2009       |
| Al-doped ZnO ISFET                                                   | 2 – 12<br>Commercial buffers                                                                                                                                                                                                                                        | ≈ 50 mV/pH                                                                 | Stable for 12 w at pH 2                                                                                                | 0.3 s         | Tsai et al., 2019         |

|                                                                       |                                                                                     |                                                                   |                                                                                  |                             |                     |
|-----------------------------------------------------------------------|-------------------------------------------------------------------------------------|-------------------------------------------------------------------|----------------------------------------------------------------------------------|-----------------------------|---------------------|
| SnO <sub>2</sub> nanorods EGFET                                       | 1 – 13<br>Commercial buffers                                                        | 55.2 mV/pH (linear regime)<br>0.86 $\mu$ A/pH (saturation regime) | Up to 6h continuous operation                                                    | n/a                         | Li et al., 2012     |
| RuO <sub>2</sub> nanomembrane EGFET                                   | 2 – 12<br>Commercial buffers<br>Low interference of mono and divalent cations       | 65.1 mV/pH (linear regime)<br>1.05 $\mu$ A/pH (saturation regime) | Drift of 2 mV/h, stabilizes after 8 h of immersion                               | n/a                         | Singh et al., 2019  |
| IrO <sub>x</sub> onto nanostructured Pt Potentiometric                | 2 – 11<br>KCl solution adjusted with strong acid/base                               | 70.9 mV/pH<br>Standard deviation < 1%.                            | Stable for 1 year, dry or immersed in PBS. Sensitivity stabilizes after 1 month. | 6 – 8 s                     | Zea et al., 2019    |
| WO <sub>3</sub> layer on CNTs Potentiometric                          | 2 – 12<br>Britton-Robinson buffer                                                   | 40.73 mV/pH                                                       | Standard error <1%<br>Stable after 1 month of storage                            | 30 s (pH 4)<br>90 s (pH 12) | Zhang & Xu, 2009    |
| WO <sub>3</sub> nanoparticles Potentiometric                          | 5 – 9<br>Commercial buffers                                                         | 56.7 $\pm$ 1.3 mV/pH                                              | Sensitivity reduction over time                                                  | 28 s                        | Santos et al., 2014 |
| WO <sub>3</sub> nanoparticles Voltammetry                             | 3 – 11<br>Phosphate buffer<br>Tested in vinegar                                     | 60.0 $\pm$ 0.01 mV/pH                                             | Average drift of 33 mV over 3 h. 95% sensitivity retained after 7 d of use.      | n/a                         | Jamal et al., 2019  |
| WO <sub>3</sub> nanofibers Potentiometric, amplified                  | 3 – 11<br>Commercial buffers<br><b>Tested in artificial seawater (pH 8.0 – 7.6)</b> | 38.9 mV/pH (amplified to 377.5 mV/pH)                             | n/a                                                                              | n/a                         | Choi et al., 2019   |
| IrO <sub>x</sub> / PDDA Potentiometric                                | 3 – 10<br>Commercial buffers                                                        | 59 mV/pH                                                          | n/a                                                                              | 3 s                         | Jović et al., 2018  |
| pTBA / AuZnOx Potentiometric                                          | 2 – 13<br>Commercial buffers<br>Also tested in saliva and urine samples             | 59.2 $\pm$ 0.5 mV/pH                                              | Stable for cyclic measurements (200 s). Good stability upon storage for 15 d     | 1 s                         | Kim et al., 2016    |
| RuO <sub>2</sub> np solid contact + PVC based membrane Potentiometric | 2 – 12<br>Tris buffer<br>Good selectivity against monovalent cations                | 59 mV/pH                                                          | Stable for 1 w of daily calibrations                                             | n/a                         | Lenar et al., 2019  |

**Table 5.** pH sensors based on semiconductor and metal/metal oxide nanomaterials. In bold, sensors that have been tested in seawater.

None of the potentiometric pH sensors listed in Table 5 was tested in seawater apart from the WO<sub>3</sub> nanofibers potentiometric amplified sensor realized by Choi et al. (2019). They tested such sensors in artificial seawater and calibrated the reading against a commercial pH meter, Due to the high sensitivity obtained through the amplification, the authors concluded that their new pH sensor is promising for portable and low-cost applications for the monitoring of seawater; however, stability and long term performances were not assessed.

## 6. Conclusions

711

712 pH is a key parameter in many chemical, biological and biogeochemical phenomena and is of  
713 particular interest in environmental monitoring. Ion sensitive glass electrodes are the most used  
714 sensors for pH measurements, but new solutions for the realization of robust, precise and affordable  
715 pH sensors are actively investigated.

716 In this review, we have presented the most recent developments in pH sensing materials, reporting  
717 sensor performances and main parameters. Solid state sensors based on inorganic materials, (metals  
718 of semiconductors), and carbon based materials (polymers and carbon particles) have been  
719 reviewed, revealing a general trend towards the realization of miniaturized, low cost/disposable  
720 sensors.

721 The development of nano-engineered materials and composites as active sensing elements has  
722 emerged as a promising strategy to improve sensitivity, response time, flexibility and ease of  
723 fabrication. Thin films and nanomaterials based on metal oxides provide good sensing performances  
724 and relatively good stability and can be easily integrated in potentiometric sensors or silicon-based  
725 FET devices. Examples of application of metal oxide pH sensors in different environments, including  
726 seawater, have been reported pointing out their robustness and flexibility.

727 Carbon nanoparticles, despite having attracted a large research effort, are not so stable in their  
728 response (sensitive to surface defects, functional groups and morphology), nor easy to produce and  
729 handle. Polymer-based sensors, finally, seems to be non-competitive in terms of precision and  
730 stability. However, the limitations shown by this class of materials can be overcome by properly  
731 combining them. In this respect, the synergy observed between polymeric components, and inorganic  
732 nanomaterials seems to be a key factor for the realization of robust and affordable sensors. Polymers  
733 can be used as efficient ion-selective or protective elements, to enhance the response of inorganic  
734 sensing elements and decrease the interference of dissolved ions. On the other hand, the response  
735 and the stability of pH sensitive polymers can be greatly improved by combining them with conductive  
736 and semiconductive nanomaterials, as shown for the most common electroactive polymer, PANI, and  
737 for polydopamine.

738 For each sensor class, results of testing in seawater, when available, have been reported and  
739 discussed. Only few new sensors have been designed for seawater, however, the examples reported  
740 show promising results, in terms of sensitivity, selectivity vs. interfering ions and stability. While some  
741 inorganic material (metal oxides) has shown good sensing performances at sea, among the devices  
742 based on polymers or carbon nanomaterials the only ones successfully tested in seawater are based  
743 on composite or multilayer structures. Design refinement and extensive field testing and validation are  
744 needed to assess the suitability of the sensors presented for seawater monitoring. Even if the  
745 possibility to replace well-established measurement technologies like glass electrodes and  
746 spectrophotometry looks, as yet, unrealistic, in the near future, robust, miniaturized, integrated arrays  
747 of solid state electrochemical pH sensors can represent a valuable alternative for specific  
748 applications.

749

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## References

- Adhikari, B., & Majumdar, S. (2004). Polymers in sensor applications. *Progress in Polymer Science*, 29(7), 699–766. doi: 10.1016/j.progpolymsci.2004.03.002
- Ahn, J.-H., Kim, J.-Y., Seol, M.-L., Baek, D.J., Guo, Z., Kim, C.-H., Choi, S.-J., & Choi, Y.-K. (2013). A pH sensor with a double-gate silicon nanowire field-effect transistor. *Applied Physics Letters*, 102, 083701. doi: 10.1063/1.4793655
- Albertin, K.F., Carreño, M.N.P., & Pereyra, I. (2013). Study of TiO<sub>2</sub> Nanotubes for Sensors and Integrated Devices. *Precision Instrument and Mechanology*, 2(3), 114-121.
- Alizadeh, T., & Jamshidi, F. (2015). Synthesis of nanosized sulfate-modified  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> and its use for the fabrication of all-solid-state carbon paste pH sensor. *Journal of Solid State Electrochemistry*, 19, 1053–1062. doi: 10.1007/s10008-014-2716-4
- Ameri, S.K., Singh, P.K., & Sonkusale, S.R. (2016). Three dimensional graphene transistor for ultra-sensitive pH sensing directly in biological media. *Analytica Chimica Acta*, 934, 212-217. doi: 10.1016/j.aca.2016.05.048
- Amiri, M., Amali, E., Nematollahzadeh, A., & Salehniya, H. (2016). Poly-dopamine films: Voltammetric sensor for pH monitoring. *Sensors and Actuators B: Chemical*, 228, 53–58. doi: 10.1016/j.snb.2016.01.012
- Ang, P.K., Chen, W., Wee, A.T.S. & Ping, K. (2008). Solution-Gated Epitaxial Graphene as pH Sensor. *Journal of the American Chemical Society*, 130, 14392–14393. doi: 10.1021/ja805090z
- Ates, M. (2013). A review study of (bio)sensor systems based on conducting polymers. *Materials Science and Engineering C*, 33(4), 1853–1859. doi: 10.1016/j.msec.2013.01.035
- Athavale, R., Dinkel, C., Wehrli, B., Bakker, E., Crespo, G.A., & Brand, A. (2017). Robust solid-contact ion selective electrodes for high-resolution in situ measurements in fresh water systems. *Environmental Science and Technology Letters*, 4(7), 286-291. doi: 10.1021/acs.estlett.7b00130
- Back, J.H., & Shim, M. (2006). pH-Dependent Electron-Transport Properties of Carbon Nanotubes. *Journal of Physical Chemistry B*, 110, 23736-23741. doi: 10.1021/jp063260x
- Bangar, M., Chen, W., Myung, N., & Mulchandani, A. (2010). Conducting polymer 1-dimensional nanostructures for FET sensors. *Thin Solid Films*, 519, 964-973. doi: 10.1016/j.tsf.2010.08.023
- Banna, M.H., Najjara, H., Sadiq, R., Imran, S.A., Rodriguez, M.J., & Hoorfar, M. (2014). Miniaturized water quality monitoring pH and conductivity sensors. *Sensors and Actuators B: Chemical*, 193, 434-441. doi: 10.1016/j.snb.2013.12.002
- Bao, Q., Yang, Z., Song, Y., Fan, M., Pan, P., Liu, J., Liao, Z., & Wei, J. (2019). Printed flexible bifunctional electrochemical urea-pH sensor based on multiwalled carbon nanotube/polyaniline electronic ink. *Journal of Materials Science: Materials in Electronics*, 30(2), 1751-1759. doi: 10.1007/s10854-018-0447-5

788 Barton, J., Begoña González García, M., Hernández Santos, D., Fanjul-Bolado, P., Ribotti, A.,  
789 McCaul, M., Diamond, D., & Magni P. (2016). Screen-printed electrodes for environmental  
790 monitoring of heavy metal ions: a review. *Microchimica Acta*, 183, 503-517. doi: 10.1007/s00604-  
791 015-1651-0

792 Bedner, K., Guzenko, V.A., Tarasov, A., Wipf, M., Stoop, R.L., Just, D., Rigante, S., Fu, W.,  
793 Minamisawa, R.A., David, C., & Calame, M. (2013). pH response of silicon nanowire sensors:  
794 impact of nanowire width and gate oxide. *Sensors and Materials*, 25(8), 567-576. doi:  
795 10.18494/SAM.2013.890

796 Bergveld, P. (2003). Thirty years of ISFETOLOGY: What happened in the past 30 years and what  
797 may happen in the next 30 years. *Sensors and Actuators B: Chemical*, 88(1), 1–20. doi:  
798 10.1016/S0925-4005(02)00301-5

799 Boeva, Z.A., Milakin, K.A., Pesonen, M., Ozerin, A.N., Sergeyev, V.G., & Lindfors, t. (2014).  
800 Dispersible composites of exfoliated graphite and polyaniline with improved electrochemical  
801 behaviour for solid-state chemical sensor applications. *RSC Advances*, 4, 46340. doi:  
802 10.1039/c4ra08362h

803 Brooke, M., Cole, E., Dale, J., Prasad, A., Quach, H., Bau, B., Nowacek, D., & Bhatt, E. (2016). An  
804 ocean sensor for measuring the seawater electrochemical response of 8 metals referenced to zinc,  
805 for determining ocean pH. *9th International Conference on Sensing Technology (ICST)*, Auckland  
806 (NZ), 8-11 December 2015, 147-150. doi: 10.1109/ICSensT.2015.7438381

807 Buck, R.P., Rondinini, S., Covington, A.K., Baucke, F.G.K., Brett, C.M.A., Camões, M.F., Milton,  
808 M.J.T., Mussini, T., Naumann, R., Pratt, K.W., & Spitzer, P. (2002). Measurement of pH. Definition,  
809 standards, and procedures. *Pure Appl. Chem.*, 74(11), 2169–2200. <http://hdl.handle.net/1808/8412>

810 Bushinsky, S.M., Takeshita, Y., & Williams, N.L. (2019). Observing Changes in Ocean Carbonate  
811 Chemistry: Our Autonomous Future. *Current Climate Change Reports*, 5(3), 207-220. doi:  
812 10.1007/s40641-019-00129-8

813 Byrne, R.H. (2014). Measuring Ocean Acidification: New Technology for a New Era of Ocean  
814 Chemistry. *Environmental Science and Technology*, 48(10), 5352-5360. doi: 10.1021/es405819p

815 Cachet-Vivier, C., Tribollet, B., Vivier, V. (2010). Cavity microelectrode for studying manganese  
816 dioxide powder as pH sensor. *Talanta*, 82(2), 555–559. doi: 10.1016/j.talanta.2010.05.006

817 Chen, X.M., Wu, G.H., Jiang, Y.Q., Wang, Y.R., & Chen, X. (2011). Graphene and graphene-based  
818 nanomaterials: the promising materials for bright future of electroanalytical chemistry. *Analyst*,  
819 136(22), 4631-40. doi: 10.1039/c1an15661f

820 Cheng, Z., Li, Q., Li, Z., Zhou, Q., & Fang, Y. (2010). Suspended Graphene Sensors with Improved  
821 Signal and Reduced Noise. *Nano Letters*, 10, 1864–1868. doi: 10.1021/nl100633g

822 Chien, Y.-S., Tsai, W.-L., Lee, I.-C., Chou, J.-C., & Cheng, H.-C. (2012). Novel Ph Sensor Of Egfets  
 823 With Laser-Irradiated Carbon-Nanotube Network. *IEEE Electron Device Letters*, 33(11), 1622-  
 824 1624. doi: 10.1109/LED.2012.2213794

825 Chinnathambi, S., & Euverink, G.J.W. (2018). Polyaniline functionalized electrochemically reduced  
 826 graphene oxide chemiresistive sensor to monitor the pH in real time during microbial  
 827 fermentations. *Sensors and Actuators B: Chemical*, 264, 38–44. doi: 10.1016/j.snb.2018.02.087

828 Choi, J.-S, Savagatrup, S., Kim, Y., Lang, J.H., & Swager, T.M. (2019). Precision pH sensor based on  
 829 WO<sub>3</sub> nanofiber-polymer composites and different amplification. *ACS Sensors*, 4(10), 2593-2598.  
 830 doi:10.1021/acssensors.9b01579

831 Choi, S., Park, I., Hao, Z., Holman, H.-Y. N., & Pisano, A.P. (2012). Quantitative studies of long-term  
 832 stable, top-down fabricated silicon nanowire pH sensors. *Applied Physics A*, 107, 421–428. doi:  
 833 10.1007/s00339-011-6754-9

834 Cisternas, R., Kahlert, H., Scholz, F., & Wulff, H. (2015). Direct contact tungsten bronze electrodes for  
 835 calibration-free potentiometric pH measurements. *Electrochemistry Communications*, 60, 17-20.  
 836 doi: 10.1016/j.elecom.2015.07.017

837 Cisternas, R., Ballesteros, L., Valenzuela, M.L., Kahlert, H., & Scholz, F. (2017). Decreasing the time  
 838 response of calibration-free pH sensors based on tungsten bronze nanocrystals. *Journal of*  
 839 *Electroanalytical Chemistry*, 801, 315–318. doi: 10.1016/j.jelechem.2017.08.005

840 Clarke, J.S., Achterberg, E.P., Rérolle, V.M.C., Abi Kaed Bey, S., Floquet, C.F.A., Mowlem, M.C.  
 841 (2015). Characterisation and deployment of an immobilised pH sensor spot towards surface ocean  
 842 pH measurements. *Analytica Chimica Acta*, 897, 69–80. doi: 10.1016/j.aca.2015.09.026

843 Cleary, J., McCaul, M., Diamond, D., García, M.B.G., Díez, C., Rovira, C., Challiss, M., Lassoued, Y.,  
 844 Ribotti, A., Sáez, J. (2014). COMMON SENSE: Cost-effective sensors, interoperable with  
 845 international existing ocean observing systems, to meet EU policies requirements. *2014 IEEE*  
 846 *Sensor Systems for a Changing Ocean (SSCO 2014)*, 1-7, doi: 10.1109/SSCO.2014.7000384

847 Crespo, G.A., Gugsa, D., Macho, S., & Rius, F.X. (2009). Solid-contact pH-selective electrode using  
 848 multi-walled carbon nanotubes. *Analytical and Bioanalytical Chemistry*, 395, 2371-2376. doi:  
 849 10.1007/s00216-009-3127-8

850 Cuartero, M., Pankratova, N., Cherubini, T., Crespo, G., Massa, F., Confalonieri, F., & Bakker, E.  
 851 (2017). In Situ Detection of Species Relevant to the Carbon Cycle in Seawater with Submersible  
 852 Potentiometric Probes. *Environmental Science & Technology Letters*, 4. doi:  
 853 10.1021/acs.estlett.7b00388

854 Culebras, M., Gómez, C.M., & Cantarero, A. (2014). Review on Polymers for Thermoelectric  
 855 Applications. *Materials*, 7(9), 6701-6732. doi: 10.3390/ma7096701

856 Cullison Gray, S.E., DeGrandpre, M.D., Moore, T.S., Martz, T.R., Friederich, G.E., & Johnson, K.S.  
857 (2011). Applications of in situ pH measurements for inorganic carbon calculations. *Marine*  
858 *Chemistry*, 125(1-4), 82–90. doi: 10.1016/j.marchem.2011.02.005

859 Dai, C., Chan, C.-W.I., Barrow, W., Smith, A., Song, P., Potier, F., Wadhawan, J.D., Fisher, A.C., &  
860 Lawrence, N.S. (2016). A Route to Unbuffered pH Monitoring: A Novel Electrochemical Approach.  
861 *Electrochimica Acta*, 190, 879-886. doi: 10.1016/j.electacta.2016.01.004

862 Fu, W., Nef, C., Knopfmacher, O., Tarasov, A., Weiss, M., Calame, M., & Schönenberger, C. (2011).  
863 Graphene Transistors Are Insensitive to pH Changes in Solution. *Nano Letters*, 11(9), 3597–3600.  
864 doi: 10.1021/nl201332c

865 Fulati, A., Usman Ali, S.M., Riaz, M., Amin, G., Nur, O., & Willander, M. (2009). Miniaturized pH  
866 Sensors Based on Zinc Oxide Nanotubes/Nanorods. *Sensors*, 9(11), 8911-8923. doi:  
867 10.3390/s91108911

868 Gao, W., & Song, J. (2009). Polyaniline Film Based Amperometric pH Sensor Using A Novel  
869 Electrochemical Measurement System. *Electroanalysis*, 21(8), 973-978. doi:  
870 10.1002/elan.200804500

871 Gill, E., Arshak, A., Arshak, K., & Korostynska, O. (2008). Conductometric pH sensor based on novel  
872 conducting polymer composite thick films. *Proceedings of the 2008 31st International Spring*  
873 *Seminar on Electronics Technology*, 478–483. doi: 10.1109/ISSE.2008.5276613

874 Glab, S., Hulanicki, A., Edwall, G., & Ingman F. (1989). Metal-Metal Oxide and Metal Oxide  
875 Electrodes as pH Sensors. *Critical Reviews in Analytical Chemistry*, 21(1), 29-47. doi:  
876 10.1080/10408348908048815

877 González Durán, E., Cuaya, M.P., Gutiérrez, M.V. & Ancona León, J. (2018). Effects of Temperature  
878 and pH on the Oxidative Stress of Benthic Marine Invertebrates. *Biology Bulletin*, 45, 610–616.  
879 <https://doi.org/10.1134/S1062359018660019>

880 Gou, P., Kraut, N.D., Feigel, I.M., Bai, H., Morgan, G.J., Chen, Y., Tang, Y., Bocan, K., Stachel, J.,  
881 Berger, L., Mickle, M., Sejdic, E., & Star, A., (2014). Carbon Nanotube Chemiresistor for Wireless  
882 pH Sensing. *Scientific Reports*, 4, 4468. doi: 10.1038/srep04468

883 Grozdanov, A., Petrovski, A., Paunovic, P., Dimitrov, T.A., & Avella M. (2018). MWCNT/Polyaniline  
884 nanocomposites used for pH nanosensors of marine waters. In M. Cocca, E. Di Pace, M. Errico,  
885 G. Gentile, A. Montarsolo, R. Mossotti (Eds.), *Proceedings of the International Conference on*  
886 *Microplastic Pollution in the Mediterranean Sea*, (pp. 231-238). Springer International Publishing.  
887 doi: 10.1007/978-3-319-71279-6\_32

888 Grozdanov, A., Petrovski, A., Avella, M., Paunovic, P., Errico, E.M., Avolio, R., Gentile, G., De Falco,  
889 F., & Dimitrov, T. A. (2019). Spectroscopic study of nanocomposite based on PANI and Carbon  
890 nanostructures for pH sensors. *Bulgarian Chemical Communications*, 51(D), 36-41.

891 Gupta, N., Sharma, S., Mir, I.A., & Kumar, D. (2006). Advances in sensors based on conducting  
 892 polymers. *Journal of Scientific and Industrial Research*, 65(7), 549-557.  
 893 <http://nopr.niscair.res.in/handle/123456789/4862>

894 Hayat, A., & Marty, J.L. (2014). Disposable screen printed electrochemical sensors: tools for  
 895 environmental monitoring. *Sensors*, 14(6), 10432-53. doi: 10.3390/s140610432

896 Heller, I., Chatoor, S., Mannik, J., Zevenbergen, M.A.G., Dekker, C., & Lemay, S.G. (2010). Influence  
 897 of Electrolyte Composition on Liquid-Gated Carbon Nanotube and Graphene Transistors. *Journal*  
 898 *of the American Chemical Society*, 132(48), 17149-17156. doi: 10.1021/ja104850n

899 Hu, J., Ho, K.T., Zou, X.U., Smyrl, W.H., Stein, A., & Bühlmann, P. (2015). All-Solid-State Reference  
 900 Electrodes Based on Colloid-Imprinted Mesoporous Carbon and their Application in Disposable  
 901 Paper-based Potentiometric Sensing Devices. *Analytical Chemistry*, 87(5), 2981–2987. doi:  
 902 10.1021/ac504556s

903 Huang, W.S., Humphrey, B.D., & MacDiarmid, A.G. (1986). Polyaniline, a novel conducting polymer.  
 904 Morphology and chemistry of its oxidation and reduction in aqueous electrolytes. *J. Chem. Soc.,*  
 905 *Faraday Trans. 1: Physical Chemistry in Condensed Phases*, 82(8), 2385-2400. doi:  
 906 10.1039/F19868202385

907 Jamal, M., Razeed, K.M., Shao, H., Islam, J., Akhter, I., Furukawa, H., & Khosla, A. (2019).  
 908 Development of Tungsten Oxide Nanoparticle Modified Carbon Fibre Cloth as Flexible pH Sensor.  
 909 *Scientific Reports*, 9(1), Article number 4659. doi: 10.1038/s41598-019-41331-w

910 Jang, H., & Lee, J. (2020). Iridium oxide fabrication and application: A review. *Journal of Energy*  
 911 *Chemistry*, 46, 152-172. doi: 10.1016/j.jechem.2019.10.026

912 Jin, Q. & Kirk, M.F. (2018). pH as a Primary Control in Environmental Microbiology: 1.  
 913 Thermodynamic Perspective. *Frontiers in Environmental Science*, 6:21. doi:  
 914 10.3389/fenvs.2018.00021

915 Johnson, K.S., Jannasch, H.W., Coletti, L.J., Elrod, V.A., Martz, T.R., Takeshita, Y., Carlson, R.J., &  
 916 Connery, J.C. (2016). Deep-Sea DuraFET: A Pressure Tolerant pH Sensor Designed for Global  
 917 Sensor Networks. *Analytical Chemistry*, 88(6), 3249-3256. doi: 10.1021/acs.analchem.5b04653

918 Jović, M., Hidalgo-Acosta, J.C., Lesch, A., Costa Bassetto, V., Smirnov, E., Cortés-Salazar, F., &  
 919 Girault, H.H. (2018). Large-scale layer-by-layer inkjet printing of flexible iridium-oxide based pH  
 920 Sensors. *Journal of Electroanalytical Chemistry*, 819, 384–390. doi:  
 921 10.1016/j.jelechem.2017.11.032

922 Jung, M.W., Myung, S., Song, W., Kang, M.A., Kim, S.H., Yang, C.S., Lee, S.S., Lim, J., Park, C.Y.,  
 923 Lee, J.O., & An, K.S. (2014). Novel fabrication of flexible graphene-based chemical sensors with  
 924 heaters using soft lithographic patterning method. *ACS Applied Materials & Interfaces*, 6(16),  
 925 13319-13323. doi: 10.1021/am502281t

926 Khan, M.I., Mukherjee, K., Shoukat, R. & Dong, H. (2017). A review on pH sensitive materials for  
 927 sensors and detection methods. *Microsystem Technologies*, 23, 4391–4404. doi: 10.1007/s00542-  
 928 017-3495-5

929 Kim, D.-M., Cho, S.J., Cho, C.-H., Kim, K.B., Kim, M.-Y., & Shim, Y.-B. (2016). Disposable all-solid-  
 930 state pH and glucose sensors based on conductive Polymer covered hierarchical AuZn oxide.  
 931 *Biosensors and Bioelectronics*, 79, 165–172. doi: 10.1016/j.bios.2015.12.002

932 Kim, K., Park, C., Rim, T., Meyyappan, M., & Lee, J.-S. (2014). Electrical and pH Sensing  
 933 Characteristics of Si Nanowire-Based Suspended FET Biosensors. *Proceedings of the 14th IEEE*  
 934 *International Conference on Nanotechnology Toronto, Canada, August 18-21*

935 Kim, T.Y., & Yang, S. (2014). Fabrication method and characterization of electrodeposited and heat-  
 936 treated iridium oxide films for pH sensing. *Sensors and Actuators B: Chemical*, 196, 31-38. doi:  
 937 10.1016/j.snb.2014.02.004

938 King, D.W., & Kester, D.R. (1989). Determination of seawater pH from 1.5 to 8.5 using colorimetric  
 939 indicator. *Marine Chemistry*, 26(1), 5-20. doi: 10.1016/0304-4203(89)90061-3

940 Kinoshita, E., Ingman, F., Edwalla, G., Thulina, S., & Glåb, S. (1986). Polycrystalline and  
 941 monocrystalline antimony, iridium and palladium as electrode material for pH-sensing electrodes.  
 942 *Talanta*, 33(2), 125-134. doi: 10.1016/0039-9140(86)80028-5

943 Knopfmacher, O., Tarasov, A., Fu, W., Wipf, M., Niesen, B., Calame, M., & Schönenberger, C. (2010).  
 944 Nernst Limit in Dual-Gated Si-Nanowire FET Sensors. *Nano Letters*, 10(6), 2268–2274. doi:  
 945 10.1021/nl100892y

946 Koncki, R., & Mascini, M. (1997). Screen-printed ruthenium dioxide electrodes for pH measurements.  
 947 *Analytica Chimica Acta*, 351(1–3), 143–149. doi: 10.1016/S0003-2670(97)00367-X

948 Korostynska, O., Arshak, K., Gill, E., & Arshak, A. (2007). Review on state-of-the-art in Polymer  
 949 Based pH sensors. *Sensors*, 7(12), 3027-3042. doi: 10.3390/s7123027

950 Kroeker, K.J., Kordas, R., Crim, R., Hendriks, I.E., Ramajo, L., Singh, G.S., Duarte, C.D., & Gattuso,  
 951 J.-P. (2013). Impacts of ocean acidification on marine organisms: quantifying sensitivities and  
 952 interaction with warming. *Global Change Biology*, 19(6), 1884-1896. doi: 10.1111/gcb.12179

953 Kumar, A., Kumar, N., Aniley, A.A., Fernandez, R.E., & Bhansali, S. (2019). Hydrothermal growth of  
 954 zinc oxide (ZnO) nanorods (NRs) on screen printed IDEs for ph measurement application. *Journal*  
 955 *of the Electrochemical Society*, 166, B3264-B3270. doi: 10.1149/2.0431909jes

956 Kumar, N., Sutradhar, M., Kumar, J., & Panda, S. (2017). Role of deposition and annealing of the top  
 957 gate dielectric in a-IGZO TFT-based dual-gate ion-sensitive field-effect transistors. *Semiconductor*  
 958 *Science and Technology*, 32, 035013. doi: 10.1088/1361-6641/aa5584

959 Kuo, L.-M., Chou, Y.-C., Chen, K.-N., Lu, C.-C., Chao, S. (2014). A precise pH microsensor using RF-  
 960 sputtering IrO<sub>2</sub> and Ta<sub>2</sub>O<sub>5</sub> films on Pt-electrode. *Sensors and Actuators B: Chemical*, 193, 687-  
 961 691. doi: 10.1016/j.snb.2013.11.109

962 Lacoue-Labarthe, T., Nunes, P.A.L.D., Ziveri, P., Cinar, M., Gazeau, F., Hilmi, N., Moschella, P., Safa,  
 963 A., Sauzade, D., & Turley, C. (2016). Impacts of ocean acidification in a warming Mediterranean  
 964 Sea: An overview. *Regional Studies in Marine Science*, 5, 1-11. doi: 10.1016/j.rsma.2015.12.005

965 Lai, C.-Z., DeGrandpre, M.D., & Darlington, R.C. (2018). Autonomous Optofluidic Chemical Analyzers  
 966 for Marine Applications: Insights from the Submersible Autonomous Moored Instruments (SAMI)  
 967 for pH and pCO<sub>2</sub>. *Frontiers in Marine Science*, 4, 438. doi: 10.3389/fmars.2017.00438

968 Lakard, B., Segut, O., Lakard, S., Herlem, G., & Gharbi, T. (2007). Potentiometric miniaturized pH  
 969 sensors based on polypyrrole films. *Sensors and Actuators B: Chemical*, 122, 101-108. doi:  
 970 10.1016/j.snb.2006.04.112

971 Lee, D., & Cui, T. (2010). Low-cost, transparent, and flexible single-walled carbon nanotube  
 972 nanocomposite based ion-sensitive field-effect transistors for pH/glucose sensing. *Biosensors and*  
 973 *Bioelectronics*, 25(10), 2259–2264. doi: 10.1016/j.bios.2010.03.003

974 Lei, K.F., Lee, K.-F., & Yang, S.-I (2012). Fabrication of carbon nanotube-based pH sensor for paper-  
 975 based microfluidics. *Microelectronic Engineering*, 100, 1–5. doi: 10.1016/j.mee.2012.07.113

976 Lenar, N., Paczosa-Bator, B., & Piech, R. (2019). Ruthenium dioxide nanoparticles as a high-capacity  
 977 transducer in solid-contact polymer membrane-based pH-selective electrodes. *Microchimica Acta*,  
 978 186(12), Article n. 777. doi: 10.1007/s00604-019-3830-x

979 Li, H.-H., Dai, W.-S., Chou, J.-C., & Cheng, H.-C. (2012). An Extended-Gate Field-Effect Transistor  
 980 With Low-Temperature Hydrothermally Synthesized SnO<sub>2</sub> Nanorods as pH Sensor. *IEEE Electron*  
 981 *Device Letters*, 33(10), 1495-1497. doi: 10.1109/LED.2012.2210274

982 Li, Q., Li, H., Zhang, J., & Xu, Z. (2011). A novel pH potentiometric sensor based on electrochemically  
 983 synthesized polybisphenol A films at an ITO electrode. *Sensors and Actuators B: Chemical*, 155,  
 984 730–736. doi: 10.1016/j.snb.2011.01.038

985 Li, X., Cai, W., An, J., Kim, S., Nah, J., Yang, D., Piner, R., Velamakanni, A., Jung, I., Tutuc, E.,  
 986 Banerjee, S.K., Colombo, L., & Ruoff, R.S. (2009). Large-area synthesis of high-quality and  
 987 uniform graphene films on copper foils. *Science*, 324(5932), 1312-1314. doi:  
 988 10.1126/science.1171245

989 Li, Y., Mao, Y., Xiao, C., Xu, X., & Li, X. (2020). Flexible pH sensor based on a conductive PANI  
 990 membrane for pH monitoring. *RSC Advances*, 10, 21-28. doi: 10.1039/C9RA09188B

991 Lin, J.-C., Huang, B.-R., & Yang, Y.-K. (2013). IGZO nanoparticle-modified silicon nanowires as  
 992 extended-gate field-effect transistor pH sensors. *Sensors and Actuators B: Chemical*, 184, 27–32.  
 993 doi: 10.1016/j.snb.2013.04.060

994 Liu, M., Ma, Y., Su, L., Chou, K.-C., & Hou, X. (2016). A titanium nitride nanotube array for  
 995 potentiometric sensing of pH. *Analyst*, 141(5), 1693-1699. doi: 10.1039/c5an02675j

996 Liu, Y., & Cui, T. (2007). Ion-sensitive field-effect transistor based pH sensors using nano self-  
 997 assembled polyelectrolyte/nanoparticle multilayer films. *Sensors and Actuators B: Chemical*, 123,  
 998 148–152. doi:10.1016/j.snb.2006.08.006

999 Loh, K.J., Lynch, J.P., & Kotov, N.A. (2007). Passive wireless strain and pH sensing using carbon  
 1000 nanotube-gold nanocomposite thin films. In M. Tomizuka, C.-B. Yun, V. Giurgiutiu (Eds.), *Sensors*  
 1001 and Smart Structures Technologies for Civil, Mechanical, and Aerospace Systems, *Processing of*  
 1002 *SPIE*, 6529, 652919-652931. doi: 10.1117/12.715826

1003 Lonsdale, W., Wajrak, M., & Alameh, K. (2018). Manufacture and application of RuO<sub>2</sub> solid-state  
 1004 metal-oxide pH sensor to common beverages. *Talanta*, 180, 277-281. doi:  
 1005 10.1016/j.talanta.2017.12.070

1006 Maily-Giacchetti, B., Hsu, A., Wang, H., Vinciguerra, V., Pappalardo, F., Occhipinti, L., Guidetti, E.,  
 1007 Coffa, S., Kong, J., & Palacios, T. (2013). pH sensing properties of graphene solution-gated field-  
 1008 effect transistors. *Journal of Applied Physics*, 114, 084505. doi: 10.1063/1.4819219

1009 Manjakkal, L., Cvejic, K., Kulawik, J., Zaraska, K., Szwagierczak, D., & Socha, R.P. (2014).  
 1010 Fabrication of thick film sensitive RuO<sub>2</sub>-TiO<sub>2</sub> and Ag/AgCl/KCl reference electrodes and their  
 1011 application for pH measurements. *Sensors and Actuators B: Chemical*, 204, 57–67.  
 1012 doi:10.1016/j.snb.2014.07.067

1013 Manjakkal, L., Synkiewicz, B., Zaraska, K., Cvejic, K., Kulawik, J., & Szwagierczak, D. (2016).  
 1014 Development and characterization of miniaturized LTCC pH sensors with RuO<sub>2</sub> based sensing  
 1015 electrodes. *Sensors and Actuators B: Chemical*, 223, 641–649. doi: 10.1016/j.snb.2015.09.135

1016 Marion, G. M., Millero, F. Camões, M.F., Spitzer, P., Feistel, R., & Chen, C.T.A. (2011). pH of  
 1017 seawater. *Marine Chemistry*, 126, 89-96. doi: 10.1016/j.marchem.2011.04.002

1018 Martin, A., & Escarpa, A. (2014). Graphene: The cutting-edge interaction between chemistry and  
 1019 electrochemistry. *TrAC Trends in Analytical Chemistry*, 56, 13-26. doi: 10.1016/j.trac.2013.12.008

1020 Martz, T., McLaughlin, K., Weisberg, S.B. (2015). Best Practices for autonomous measurement of  
 1021 seawater pH with the Honeywell Durafet pH sensor. *Technical Report 861. California Current*  
 1022 *Acidification Network. Santa Barbara, CA.*  
 1023 [http://ftp.sccwrp.org/pub/download/DOCUMENTS/TechnicalReports/861\\_CCAN\\_Durafet\\_Best\\_Pr](http://ftp.sccwrp.org/pub/download/DOCUMENTS/TechnicalReports/861_CCAN_Durafet_Best_Practices_Manual.pdf)  
 1024 [actices\\_Manual.pdf](http://ftp.sccwrp.org/pub/download/DOCUMENTS/TechnicalReports/861_CCAN_Durafet_Best_Practices_Manual.pdf)

1025 McLaughlin, K., Dickson, A., Weisberg, S.B., Coale, K., Elrod, V., Hunter, C., Johnson, K.S., Kram, S.,  
 1026 Kudela, R., Martz, T., Negrey, K., Passow, U., Shaughnessy, F., Smith, J.E., Tadesse, D.,  
 1027 Washburn, L., & Weis, K.R. (2017a). An evaluation of ISFET sensors for coastal pH monitoring  
 1028 applications. *Regional Studies in Marine Science*, 12, 11-18. doi: 10.1016/j.rsma.2017.02.008

1029 McLaughlin, K., Nezhlin, N.P., Weisberg, S.B., Dickson, A.G., Booth, J.A., Cash, C.L., Feit, A., Gully,  
 1030 J.R., Johnson, S., Latker, A., Mengel, M.J., Robertson, G.L., Steele, A., & Terriquez, L. (2017b).  
 1031 An evaluation of potentiometric pH sensors in coastal monitoring applications. *Limnology and*  
 1032 *Oceanography: Methods*, 15, 679-689. doi: 10.1002/lom3.10191

1033 Michalska, A. (2012). All-Solid-State Ion Selective and All-Solid-State Reference Electrodes.  
 1034 *Electroanalysis*, 24, 1253–1265. doi: 10.1002/elan.201200059

1035 Millero, F.J., DiTrollo, B., Suarez, A.F., & Lando, G. (2009). Spectroscopic measurements of the pH in  
 1036 NaCl brines. *Geochimica Cosmochimica Acta*, 73, 3109–3114. doi: 10.1016/j.gca.2009.01.037

1037 Moya, A., Pol, R., Martínez-Cuadrado, A., Villa, R., Gabriel, G., & Baeza, M. (2019). Stable Full-Inkjet-  
 1038 Printed Solid-State Ag/AgCl Reference Electrode. *Analytical Chemistry*, 91(24), 15539-15546. doi:  
 1039 10.1021/acs.analchem.9b03441

1040 Newton, J.A., Feeley, R.A., Jewett, E.B., Williamson, P., & Mathis, J. (2015). *Global Ocean*  
 1041 *Acidification Observing Network: Requirements and Governance Plan (2<sup>nd</sup> edition)*. GOA-ON.  
 1042 [http://www.goa-on.org/documents/general/GOA-ON\\_2nd\\_edition\\_final.pdf](http://www.goa-on.org/documents/general/GOA-ON_2nd_edition_final.pdf)

1043 Novoselov, K.S., Falko, V.I., Colombo, L., Gellert, P.R., Schwab, M.G., & Kim, K. (2012). A roadmap  
 1044 for graphene. *Nature*, 490(7419), 192-200. doi: 10.1038/nature11458

1045 Ohno, Y., Maehashi, K., Yamashiro, Y., & Matsumoto, K. (2009). Electrolyte-Gated Graphene Field-  
 1046 Effect Transistors for Detecting pH and Protein Adsorption. *Nano Letters*, 9(9), 3318-3322. doi:  
 1047 10.1021/nl901596m

1048 Ohno, Y., Maehashi, K., & Matsumoto, K. (2010). Chemical and biological sensing applications based  
 1049 on graphene field-effect transistors. *Biosensors and Bioelectronics*, 26, 1727–1730. doi:  
 1050 10.1016/j.bios.2010.08.001

1051 Okazaki, R.R., Sutton, A.J., Feely, R.A., Dickson, A.G., Alin, S.R., Sabine, C.L., Bunje, P.M.E., &  
 1052 Virmani, J.I. (2017). Evaluation of marine pH sensors under controlled and natural conditions for  
 1053 the Wendy Schmidt Ocean Health XPRIZE. *Limnology and Oceanography: Methods*, 15, 586–600.  
 1054 doi: 10.1002/lom3.10189

1055 Oueiny, C., Berlioz, S., & Perrin, F.-J. (2014). Carbon nanotube–polyaniline composites. *Progress in*  
 1056 *Polymer Science*, 39(4), 707–748. doi: 10.1016/j.progpolymsci.2013.08.00

1057 Pan, Y., Sun, Z., He, H., Li, Y., You, L., & Zheng, H. (2018). An improved method of preparing iridium  
 1058 oxide electrode based on carbonate-melt oxidation mechanism. *Sensors and Actuators B:*  
 1059 *Chemical*, 261, 316-324. doi: 10.1016/j.snb.2018.01.069

1060 Pandey, P.C., & Singh, G. (2001). Tetraphenylborate doped polyaniline based novel pH sensor and  
 1061 solid-state urea biosensor. *Talanta*, 55(4), 773–782. doi: 10.1016/S0039-9140(01)00505-7

1062 Parizi, K.B., Yeh, A.J., Poon, A.S.Y., & Wong, H.S.P. (2012). Exceeding Nernst limit (59mV/pH):  
 1063 CMOS-based pH sensor for autonomous applications. *2012 International Electron Devices*  
 1064 *Meeting*, 24.7.1 - 24.7.4. doi: 10.1109/IEDM.2012.6479098

1065 Park, H.J., Yoon, J.H., Lee, K.G., & Choi, B.G. (2019). Potentiometric performance of flexible pH  
 1066 sensor based on polyaniline nanofiber arrays. *Nano Convergence*, 6(1), Article n. 9. doi:  
 1067 10.1186/s40580-019-0179-0

1068 Park, I., Li, Z., Pisano, A.P., & Williams, R.S. (2010). Top-down fabricated silicon nanowire sensors  
1069 for real-time chemical detection. *Nanotechnology*, 1, 015501. doi: 10.1088/0957-4484/21/1/015501

1070 Patil, S., Ghadi, H., Ramgir, N., Adhikari, A., & Rao, V.R. (2019). Monitoring soil pH variation using  
1071 Polyaniline/SU-8 composite film based conductometric microsensor. *Sensors and Actuators B:  
1072 Chemical*, 286, 583-590. doi: 10.1016/j.snb.2019.02.016

1073 Persaud, K.C., & Pelosi, P. (1985). An approach to an artificial nose. *Transactions - American Society  
1074 for Artificial Internal Organs*, 31, 297–300.

1075 Pfattner, R., Foudeh, A.M., Chen, S., Niu, W., Matthews, J.R., He, M., & Bao, Z. (2019). Dual-Gate  
1076 Organic Field-Effect Transistor for pH Sensors with Tunable Sensitivity. *Advanced Electronic  
1077 Materials*, 5(1), Article n. 1800381. doi: 10.1002/aelm.201800381

1078 Poma, N., Vivaldi, F., Bonini, A., Carbonaro, N., Di Rienzo, F., Melai, B., Kirchhain, A., Salvo, P.,  
1079 Tognetti, A., & Di Francesco, F. (2019). Remote monitoring of seawater temperature and pH by  
1080 low cost sensors. *Microchemical Journal*, 148, 248-252. doi: 10.1016/j.microc.2019.05.001

1081 Pyo, J.-Y., & Cho, W.-J. (2017). High-performance SEGISFET pH Sensor using the structure of  
1082 double-gate a-IGZO TFTs with engineered gate oxides. *Semiconductor Science and Technology*,  
1083 32(3), Article N. 035015. doi: 10.1088/1361-6641/aa584b

1084 Qin, Y., Kwon, H.-J., Subrahmanyam, A., Howlader, M.M.R., Selvaganapathy, P.R., Adronov, A., &  
1085 Deen, M.J. (2016). Inkjet-printed bifunctional carbon nanotubes for pH sensing. *Materials Letters*,  
1086 176, 68–70. doi: 10.1016/j.matlet.2016.04.048

1087 Radu, A., Anastasova, S., Fay, C., Diamond, D., Bobacka, J. & Lewenstam, A. (2010). Low cost,  
1088 calibration-free sensors for in situ determination of natural water pollution. *SENSORS, 2010 IEEE*,  
1089 1487-1490. doi: 10.1109/ICSENS.2010.5690357

1090 Rahimi, R., Ochoa, M., Parupudi, T., Zhao, X., Yazdi, I.K., Dokmeci, M.R., Tamayol, A.,  
1091 Khademhosseini, A., & Ziaie, B. (2016). A low-cost flexible pH sensor array for wound  
1092 assessment. *Sensors and Actuators B: Chemical*, 229, 609–617. doi: 10.1016/j.snb.2015.12.082

1093 Ramanathan, T., Abdala, A.A., Stankovich, S., Dikin, D.A., Herrera-Alonso, M., Piner, R.D., Adamson,  
1094 D.H., Schniepp, H.C., Chen, X., Ruoff, R.S., Nguyen, S.T., Aksay, I.A., Prud'Homme, R.K., &  
1095 Brinson, L.C.(2008). Functionalized graphene sheets for polymer nanocomposites. *Nature  
1096 Nanotechnology*, 3(6), 327-331. doi: 10.1038/nnano.2008.96

1097 Rérolle, V.M.C., Floquet, C.F.A., Mowlem, M.C., Connelly, D.P., Achterberg, E.P., & Bellerby,  
1098 R.R.G.J. (2012). Seawater-pH measurements for ocean-acidification observations. *TrAC Trends in  
1099 Analytical Chemistry*, 40, 146-157. doi: 10.1016/j.trac.2012.07.016

1100 Rérolle, V.M.C., Achterberg, E.P., Ribas-Ribas, M., Kitidis, V., Brown, I., Bakker, D.C.E., Lee, G.A., &  
1101 Mowlem, M.C. (2018). High Resolution pH Measurements Using a Lab-on-Chip Sensor in Surface  
1102 Waters of Northwest European Shelf Seas. *Sensors*, 18(8), 2622. doi: 10.3390/s18082622

1103 Ribotti, A., Magni, P., Borghini, M., Schroeder, K., Barton, J., McCaul, M., Diamond, D., (2015), New  
 1104 cost-effective, interoperable sensors tested on existing ocean observing platforms in application of  
 1105 European directives: The COMMON SENSE European project. *Proceedings of the IEEE OCEANS*  
 1106 *2015 Conference*, 1-9. doi: 10.1109/OCEANS-Genova.2015.7271340

1107 Saba, G.K., Wright-Fairbanks, E., Chen, B., Cai, W.-J., Barnard, A.H., Jones, C.P., Branham, C.W.,  
 1108 Wang, K., & Miles, T. (2019). The Development and Validation of a Profiling Glider Deep ISFET-  
 1109 Based pH Sensor for High Resolution Observations of Coastal and Ocean Acidification. *Frontiers*  
 1110 *in Marine Science*, 6, Article n. 664. doi: 10.3389/fmars.2019.00664

1111 Sadig, H.R., Cheng, L., & Xiang, T. (2018). Using sol-gel supported by novel economic and  
 1112 environment-friendly spray-coating in the fabrication of nanostructure tri-system metal oxide-based  
 1113 pH sensor applications. *Journal of Electroanalytical Chemistry*, 827, 93-102. doi:  
 1114 10.1016/j.jelechem.2018.09.017

1115 Salaün, A-C., Pichon, L., & Wenga, G. (2014). Polysilicon nanowires FET as highly-sensitive pH-  
 1116 sensor: modeling and measurements. *Procedia Engineering*, 87, 911 – 914.  
 1117 doi:10.1016/j.proeng.2014.11.303

1118 Salavagione, H.J., Diez-Pascual, A.M., Lazaro, E., Vera, S., & Gomez-Fatou, M.A. (2014). Chemical  
 1119 sensors based on polymer composites with carbon nanotubes and graphene: the role of the  
 1120 polymer. *Journal of Materials Chemistry A*, 2, 14289. doi: 10.1039/C4TA02159B

1121 Salvo, P., Calisi, N., Melai, B., Cortigiani, B., Mannini, M., Caneschi, A., Lorenzetti, G., Paoletti, C.,  
 1122 Lomonaco, T., Paolicchi, A., Scataglini, I., Dini, V., Romanelli, M., Fuoco, R., & Di Francesco, F.  
 1123 (2017). Temperature and pH sensors based on graphenic materials. *Biosensors and*  
 1124 *Bioelectronics*, 91, 870-877. doi: 10.1016/j.bios.2017.01.062

1125 Santos, L., Neto, J.P., Crespo, A., Nunes, D., Costa, N., Fonseca, I.M., Barquinha, P., Pereira, L.,  
 1126 Silva, J., Martins, R., & Fortunato, E. (2014). WO<sub>3</sub> nanoparticle-based conformable pH sensor.  
 1127 *ACS Applied Materials and Interfaces*, 6(15), 12226-12234. doi: 10.1021/am501724h

1128 Segut, O., Lakard, B., Herlem, G., Rauch, J.-Y., Jeannot, J.-C., Robert, L., & Fahys, B. (2007).  
 1129 Development of miniaturized pH biosensors based on electrosynthesized polymer films. *Analytica*  
 1130 *Chimica Acta*, 597(2), 313-321. doi: 10.1016/j.aca.2007.06.053

1131 Sha, R., Komori, K., & Badhulika, S. (2017). Amperometric pH Sensor Based on Graphene-  
 1132 Polyaniline Composite. *IEEE Sensors Journal*, 17(16), Article n. 7959556, 5038-5043. doi:  
 1133 10.1109/JSEN.2017.2720634

1134 Sharma, B.K., & Ahn, J.-H. (2013). Graphene based field effect transistors: Efforts made towards  
 1135 flexible electronics. *Solid-State Electronics*, 89, 177–195. doi:10.1016/j.sse.2013.08.007

1136 Shirale, D.J., Bangar, M.A., Chen, W., Myung, N.V., & Mulchandani, A. (2010). Effect of aspect ratio  
 1137 (length:diameter) on a single polypyrrole nanowire FET device. *Journal of Physical Chemistry C*,  
 1138 114, 13375–13380. doi: 10.1021/jp104377e

1139 Singh, K., Lou, B.-S., Her, J.-L., Pang, S.-T., & Pan, T.-M. (2019). Super Nernstian pH response and  
 1140 enzyme-free detection of glucose using sol-gel derived RuO<sub>x</sub> on PET flexible-based extended-gate  
 1141 field-effect transistor. *Sensors and Actuators B: Chemical*, 298, Article n.126837. doi:  
 1142 10.1016/j.snb.2019.126837

1143 Sohn, I.-Y., Kim, D.-J., Jung, J.-H., Yoon, O.J., Thanh, T.N., Quang, T.T., & Lee, N.E. (2013). pH  
 1144 sensing characteristics and biosensing application of solution-gated reduced graphene oxide field-  
 1145 effect transistors. *Biosensors and Bioelectronics*, 45, 70–76. doi: 10.1016/j.bios.2013.01.051

1146 Somero, G.N., Beers, J.M., Chan, F., Hill, T.M., Klinger, T., & Litvin, S.Y. (2016). What changes in the  
 1147 carbonate system, oxygen, and temperature portend for the Northeastern Pacific Ocean: A  
 1148 physiological perspective. *BioScience*, 66, 14–26. doi: 10.1093/biosci/biv162

1149 Sophocleous, M., & Atkinson, J.K. (2017). A review of screen-printed silver/silver chloride (Ag/AgCl)  
 1150 reference electrodes potentially suitable for environmental potentiometric sensors. *Sensors and*  
 1151 *Actuators A: Physical*, 267, 106–120. doi: 10.1016/j.sna.2017.10.013

1152 Sørensen, S.P.L. (1909). Enzymstudien II: Über die Messung und die Bedeutung der  
 1153 Wasserstoffionenkonzentration bei enzymatischen Prozessen. *Biochemie Zeitung*, 21, 131–200.  
 1154 <https://d-nb.info/1125891521/34>

1155 Spijkman, M., Myny, K., Smits, E.C.P., Heremans, P., Blom, P.W.M., & de Leeuw, D.M. (2011a) Dual-  
 1156 Gate Thin-Film Transistors, Integrated Circuits and Sensors. *Advanced Materials*, 23(29), 3231-  
 1157 3242. doi: 10.1002/adma.201101493

1158 Spijkman, M., Smits, E., Cillessen, J.F.M., Biscarini, F., Blom, P.W.M., & de Leeuw, D.M. (2011b).  
 1159 Beyond the Nernst-limit with dual-gate ZnO ion-sensitive field-effect transistors. *Applied Physics*  
 1160 *Letters*, 98(4), 043502-043502-3. doi: 10.1063/1.3546169

1161 Staudinger, C., Strobl, M., Breininger, J., Klimant, I., & Borisov, S.M. (2019). Fast and stable optical  
 1162 pH sensor materials for oceanographic applications. *Sensors and Actuators B: Chemical*, 282,  
 1163 204–217. doi: 10.1016/j.snb.2018.11.048

1164 Staudinger, C., Strobl, M., Fischer, J.P., Thar, R., Mayr, T., Aigner, D., Müller, B.J., Müller, B., Lehner,  
 1165 P., Mistlberger, G., Fritzsche, E., Ehgartner, J., Zach, P.W., Clarke, J.S., Geißler, F., Mutzberg, A.,  
 1166 Müller, J.D., Achterberg, E.P., Borisov, S.M., & Klimant, I. (2018). A versatile optode system for  
 1167 oxygen, carbon dioxide, and pH measurements in seawater with integrated battery and logger.  
 1168 *Limnology and Oceanography: Methods*, 16(7), 459–473. doi: 10.1002/lom3.10260

1169 Stow, C.A., Jolliff, J., McGillicuddy Jr., D.J., Doney, S.C., Allen, J.I., Friedrichs, M.A.M., Rose, K.A., &  
 1170 Wallhead, P. (2009). Skill assessment for coupled biological/physical models of marine systems.  
 1171 *Journal of Marine Systems*, 76(1-2), 4–15. doi: 10.1016/j.jmarsys.2008.03.011

1172 Su, W., Xu, J., & Ding, X. (2016). An Electrochemical pH Sensor Based on the Amino-Functionalized  
 1173 Graphene and Polyaniline Composite Film. *IEEE Transactions on Nanobioscience*, 15, 8. doi:  
 1174 10.1109/TNB.2016.2625842

- 1175 Sulka, G.D., Hnida, K., & Brzózka, A. (2013). pH sensors based on polypyrrole nanowire arrays.  
1176 *Electrochimica Acta*, 104, 536– 541. doi: 10.1016/j.electacta.2012.12.064
- 1177 Takechi, K., Iwamatsu, S., Konno, S., Yahagi, T., Abe, Y., Katoh, M., & Tanabe, H. (2015). Gate-to-  
1178 source voltage response in high-sensitivity amorphous InGaZnO<sub>4</sub> thin-film transistor pH sensors.  
1179 *Japanese Journal of Applied Physics*, 54(7), 078004. doi: 10.7567/jjap.54.078004
- 1180 Tan, X., Chuang, H.-J., Lin, M.-W., Zhou, Z., & Cheng, M. M.-C. (2013). Edge Effects on the pH  
1181 Response of Graphene Nanoribbon Field Effect Transistors. *Journal of Physical Chemistry C*,  
1182 117(51), 27155–27160. doi: 10.1021/jp409116r
- 1183 Tasis, D., Tagmatarchis, N., Bianco, A., & Prato, M. (2006). Chemistry of carbon nanotubes. *Chemical*  
1184 *Reviews*, 106(3), 1105-1136. doi: 10.1021/cr050569o
- 1185 Trasatti, S. (1991). Physical electrochemistry of ceramic oxides. *Electrochimica Acta*, 36, 225–241.  
1186 doi: 10.1016/0013-4686(91)85244-2
- 1187 Tsai, Y.-T., Chang, S.-J., Ji, L.-W., Hsiao, Y.-J., & Tang, I.-T. (2019). Fast Detection and Flexible  
1188 Microfluidic pH Sensors Based on Al-Doped ZnO Nanosheets with a Novel Morphology. *ACS*  
1189 *Omega*, 4(22), 19847-19855. doi: 10.1021/acsomega.9b02778
- 1190 Uria, N., Abramova, N., Bratov, A., Muñoz-Pascual, F.-X., & Baldrich, E. (2016). Miniaturized metal  
1191 oxide pH sensors for bacteria detection. *Talanta*, 147, 364–369. doi: 10.1016/j.talanta.2015.10.011
- 1192 Weldborg, M., Turner, D.R., Anderson, L.G., & Dyrssen, D. (2009). Determination of pH. In K.  
1193 Grasshoff, K. Kremling, & M.Ehrhardt (Eds.), *Methods of Seawater Analysis*, John Wiley & Sons.
- 1194 Wu, Y.-C., Wu, S.-J., & Lin, C.-H. (2017). Mass-produced polyethylene-terephthalate film coated with  
1195 tantalum pentoxide for pH measurement under ISFET detection configuration. *Microsystem*  
1196 *Technologies*, 23(2), 293-298. doi: 10.1007/s00542-015-2474-y
- 1197 Xu, K., Zhang, X., Hou, K., Geng, M., & Zhao, L. (2016). The effects of antimony thin film thickness on  
1198 antimony pH electrode coated with nafion membrane. *Journal of the Electrochemical Society*,  
1199 163(8), B417-B421. doi: 10.1149/2.0191608jes
- 1200 Xu, K., Zhang, X., Chen, C., & Geng, M. (2018). Development and Performance of an All-Solid-Stated  
1201 pH Sensor Based on Modified Membranes. *International Journal of Electrochemical Science*, 13,  
1202 3080-3090. doi: 10.20964/2018.03.04
- 1203 Yao, S., Wang, M., & Madou, M. (2001). A pH Electrode Based on Melt-Oxidized Iridium Oxide.  
1204 *Journal of the Electrochemical Society*, 148(4), H29-H36. doi: 10.1149/1.1353582
- 1205 Yoon, J.H., Hong, S.B., Yun, S.-O., Lee, S.J., Lee, T.J., Lee, K.G., Choi, B.G. (2017). High  
1206 performance flexible pH sensor based on polyaniline nanopillar array electrode. *Journal of Colloid*  
1207 *and Interface Science*, 490, 53–58. doi: 10.1016/j.jcis.2016.11.033
- 1208 Young, S.-J., & Tang, W.-L. (2019). Wireless Zinc Oxide Based pH Sensor System. *Journal of the*  
1209 *Electrochemical Society*, 166(9), B3047-B3050. doi: 10.1149/2.0071909jes

1210 Zea, M., Moya, A., Fritsch, M., Ramon, E., Villa, R., & Gabriel, G. (2019). Enhanced Performance  
 1211 Stability of Iridium Oxide-Based pH Sensors Fabricated on Rough Inkjet-Printed Platinum. *ACS*  
 1212 *Applied Materials and Interfaces*, 11(16), 15160-15169. doi: 10.1021/acsami.9b03085

1213 Zhang, X., Ye, Y., Kan, Y., Huang, Y., Jia, J., Zhao, Y., Chen, C.-T.A., Qin, H. (2017). A new  
 1214 electroplated Ir/Ir(OH)<sub>x</sub> pH electrode and its application in the coastal areas of Newport Harbor,  
 1215 California. *Acta Oceanologica Sinica*, 36 (5), 99–104. doi: 10.1007/s13131-017-1064-5

1216 Zhang, W.-D., & Xu, B. (2009). A solid-state pH sensor based on WO<sub>3</sub>-modified vertically aligned  
 1217 multiwalled carbon nanotubes. *Electrochemistry Communications*, 11, 1038–1041.  
 1218 doi:10.1016/j.elecom.2009.03.006

1219 Zhao, R., Xu, M., Wang, J., & Chen, G. (2010). A pH sensor based on the TiO<sub>2</sub> nanotube array  
 1220 modified Ti electrode. *Electrochimica Acta*, 55 (20), 5647-5651.  
 1221 doi:10.1016/j.electacta.2010.04.102

1222 Zhao, W.-S., Fu, K., Wang, D.-W., Li, M., Wang, G., Yin, W.-Y. (2019). Mini-review: modeling and  
 1223 performance analysis of nanocarbon interconnects. *Applied Sciences*, 9 (11), 2174.  
 1224 doi:10.3390/app9112174

1225 Zhu, Y., Wang, C., Petrone, N., Yu, J., Nuckolls, C., Hone, J., & Lin, Q. (2015). A solid-gated  
 1226 graphene FET sensor for pH measurements. In *Proceedings of the IEEE International Conference*  
 1227 *on Micro Electro Mechanical Systems (MEMS)*. doi: 10.1109/MEMSYS.2015.7051097

1228 Zuaznabar-Gardona, J.C., & Frago, A. (2018). A wide-range solid state potentiometric pH sensor  
 1229 based on poly-dopamine coated carbon nano-onion electrodes. *Sensors and Actuators B:*  
 1230 *Chemical*, 273, 664-671. doi: 10.1016/j.snb.2018.06.103