Disposable Screen Printed Electrochemical Sensors: Tools for Environmental Monitoring

Akhtar Hayat, Jean-Louis Marty

Sensors **2014**, *14*, 10432-10453; doi:10.3390/s140610432

OPEN ACCESS

sensors

ISSN 1424-8220

www.mdpi.com/journal/sensors

Review

Disposable Screen Printed Electrochemical Sensors: Tools for Environmental Monitoring

Akhtar Hayat [1,2] and Jean Louis Marty [1,*]

[1] Images, Universite De Perpignan Via Domitia, 52 Avenue Paul Alduy, Perpignan Cedex 66860, France

[2] Interdisciplinary Research Centre in Biomedical Materials (IRCBM), COMSATS Institute of Information Technology (CIIT), Lahore 54000, Pakistan; E-Mail: akhtarloona@gmail.com

***** Author to whom correspondence should be addressed; E-Mail: jlmarty@univ-perp.fr; Tel.: +33-468-661-756.

Received: 9 March 2014; in revised form: 3 June 2014 / Accepted: 5 June 2014 / Published: 13 June 2014

Abstract: Screen printing technology is a widely used technique for the fabrication of electrochemical sensors. This methodology is likely to underpin the progressive drive towards miniaturized, sensitive and portable devices, and has already established its route from "lab-to-market" for a plethora of sensors. The application of these sensors for analysis of environmental samples has been the major focus of research in this field. As a consequence, this work will focus on recent important advances in the design and fabrication of disposable screen printed sensors for the electrochemical detection of environmental contaminants. Special emphasis is given on sensor fabrication methodology, operating details and performance characteristics for environmental applications.

Keywords: screen printed electrode; fabrication methods; sensors; electrochemical detection; environmental investigation

1. Introduction

A major part of analytical research activity is devoted to the development of new and robust methodologies. For example, new analytical tools are required for economical and real time monitoring of environmental pollutants, and for prevention of toxic materials in the environment. Progress in the field of analytical chemistry is aimed at bringing the analytical data close to the production

operations [1–3]. Such advances offer improved analytical methods with reduced environmental impact. A real time field detection system is highly desirable for continuous environmental monitoring to overcome the limitations such as sample collection and transport to a central laboratory, problems associated with commonly used methods for environmental pollutants. In this context, real time methods offer a rapid return of the chemical profile (alarm tools for sudden discharge) with minimized errors and costs as compared to the offsite laboratory-based analyses [1]. The development of portable approaches and devices with reduced sample volume is of considerable interest for both centralized and decentralized (field) analyses. This review paper highlights recent advances, primarily from the authors' laboratories, aimed at designing electrochemical systems for meeting the needs of analytical chemistry. Electrochemical devices offer unique properties to address the challenges of analytical chemistry. The advantages of electrochemical devices include possibility of miniaturization and portability, sensitivity, selectivity, a wide linear range, minimal space and power requirement and cost effective instrumentation. Devices based on the electrochemical detection are well established for many years. The past decades have seen enormous progress in electro-analytical chemistry with the development of ultra-microelectrodes, tailored interfaces, molecular devices and smart sensors. These developments have resulted in substantial popularity of electro-analyses and to their expansion into new phases and environments [4,5].

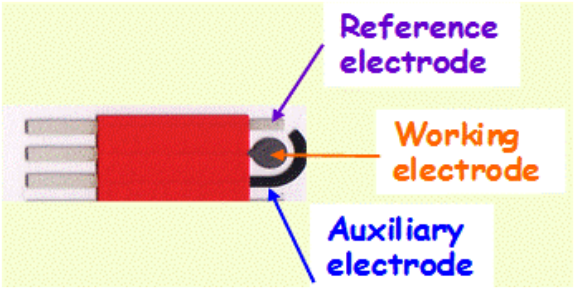
As we enter the 21 century, we do not want to rely on cumbersome electrochemical cells and bulky electrodes but rather would like to have fast, small, easy to use, portable, economical and disposable electrode systems. A vast array of electrodes for on site and *in situ* environmental monitoring has been developed during recent years [6–10]. Several representative examples, illustrating the scope, power, versatility and application of such miniaturized electrodes for environmental monitoring are described in the coming sections. This review paper as whole will focus on the trends in screen printed electrode design, screen printed electrode fabrication processes, types of screen printed electrodes and finally their environmental applications.

2. Towards Easy to Use, Disposable and Portable Screen Printed Electrodes

The elimination of bulky materials and instruments from the analytical protocol is a major thrust of analytical chemistry. The performance of analytical methods is directly related to the material of the working electrode. For many years, mercury was used as the most suitable electrode material due to its very attractive behavior and highly reproducible, renewable and smooth surface. These distinct properties of the mercury drop electrode led to the Nobel Prize in Chemistry awarded to Heyerovský in 1959. Both doping and hanging electrodes have been widely used in various polarographic and electrochemical techniques [1]. With the advancements in electro-analytical science, various non-mercury electrodes have also been examined. For example, bismuth and carbon electrodes started to be used in electro-analysis more than three decades ago due to their low background current, wide potential range, chemical inertness and suitability for various sensing and detection applications [11–13]. Recently, miniaturization of the solid electrodes was used to get several fundamental and practical advantages including such as a dramatic reduction in sample volume, portability and cost effectiveness. To address the needs of on-site analysis, it was necessary to move away from the commonly used cumbersome electrodes and cells. The exploitation of new fabrication techniques allows the replacement of traditional beaker type electrochemical cells and bulky electrodes with easy to use sensors. Fabrication

of printed devices on bendable substrates has enabled the development of a wide range of new electrode systems. Screen printing technology is a well established technique for the fabrication of economical, portable and disposable electrode systems [14,15]. The whole electrode system, including reference, counter and working electrodes can be printed on the same substrate surface [8] (Figure 1). One prominent commercialization of screen printed electrode is the glucose biosensor used for diabetes which represents a billion dollar per year global market [16–18]. Society is in constant state of growth and development, and it is evident that demands for sensing devices related to the environment will increase with the passage of time. In order to achieve this, accurate, portable and rapid devices are highly needed. Decentralized analyses are necessary and thus traditional analytical methods cannot cope with these requirements.

Figure 1. Design of a disposable and portable screen printed electrode (with reference, working and auxiliary electrodes on the same substrate) (IMAGES, Perpignan, France).



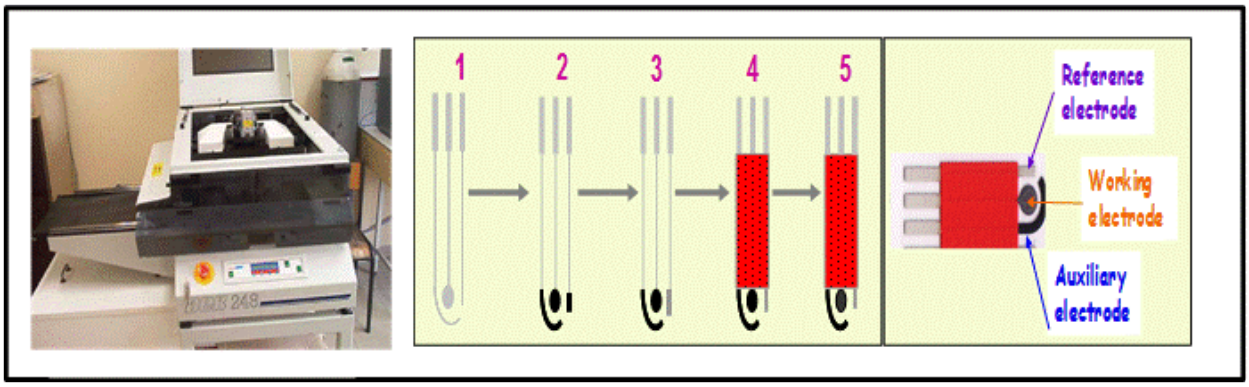
Screen printed electrodes not only address the issue of cost effectiveness but also satisfy the requirement of portability, a progress towards decentralized analysis. The adaptability of screen printed electrodes is of vital importance in the area of research, the ability to modify electrodes with ease through different inks commercially available for the reference, counter and working electrode, allows for highly specific and finally calibrated electrodes to be produced for specific target analytes [8,19,20]. Many kinds of screen printed electrode modifiers exist for environmental monitoring such as noble metals, inorganic nanomaterials, proteins, enzymes and DNA sequences [21]. Screen printed electrodes combine the properties of ease of use and portability with simple, inexpensive analytical methods [22,23]. Consequently, screen printed electrodes can be easily adapted to *in situ* environmental monitoring to achieve improved performance, as has been demonstrated over the past several years.

3. Working Principle of a Screen Printed Electrochemical Sensor

Screen printed methodology offers an attractive way to design new generation electrochemical sensors. Scientists from across different fields have shown their interest in designing low cost and reliable screen printed electrochemical sensors. A detailed description of screen printed electrode fabrication methodology was already reported in a review paper by Li *et al*. [9]. Briefly, a screen printed electrode comprises a chemically inert substrate on which three electrodes, including working electrode, reference electrode and counter electrode, are printed through screen printing methodology. The working electrode is the principal electrode on which electrochemical reactions are performed, while the reference electrode and counter electrode are used to complete the electronic circuit. Figure 2 represents the stepwise fabrication of a screen printed electrode. The chemical or biological event on

the screen printed electrode is converted into a detectable signal with the integration of a transducer element. Among the different transduction techniques, the electrochemical method of detection has attracted more interest for the design of low cost devices. Electrochemical methods of detection include amperometric (based on the current measurement), potentiometric (based on the voltage or potential differences) and conductometric (based on the conductivity or resistance). Among the electrochemical sensing techniques, amperometric detection is widely adopted due to its high sensitivity and applicability.

Figure 2. Fabrication of a three electrode system. Chemically inert substrate; screen printing of working and auxiliary electrode; screen printing of reference electrode; screen printing of protection paste; working electrode incubation with the analyte of interest (IMAGES, Perpignan, France).



The fabrication of an electrochemical screen printed sensor usually involves three steps: fabrication of the screen printed electrode, surface design of the screen printed electrode and subsequently utilization for a sensing application. The next section will focus on the fabrication strategies employed to design efficient electrochemical sensors in term of analytical characteristics.

4. Dynamics of Screen Printed Electrochemical Sensors

The inks used in screen printed electrode fabrication consist of particles, polymeric binder and other additives for improved dispersion, printing and adhesion process. The exact ink formulation and composition are patented by the respective companies, and are not disclosed to the users. The variation in the ink composition such as types, size or loading of particles strongly influence the electron transfer process and change the overall performance of the designed screen printed sensor [24–27]. However, screen printed electrodes surface can be very easily modified with a variety of materials and structurally related materials to compensate these limitations. Although these screen printed sensors have found widespread applications, the fundamental understanding of the electrochemical reactivity at the screen printed electrode is still rarely studied and addressed. In this regards, Sljukie *et al*. have shown that the performance of a macro-screen printed sensor can be improved by the use of ultrasound through increase in mass transport of the analyte and removal of surface active species. [28] Choudhry *et al*., have for the first time explored the fact that the electrode morphology can be changed dramatically by varying the concentration of the polymeric binder [29]. The same group has demonstrated that a bespoke screen printed electrode can be modified with electro-active palladium for the electrochemical oxidation of hydrazine where the unmodified electrode exhibited slow electron

transfer [30]. It has also been shown that non-linear diffusion over an electrode surface may affect the individual contributions of edge and basal plane materials, affecting the relative area of the electrode surfaces [31]. Recently, Choudhry and Banks have designed a screen printed electrode substrate modified with nickel nanoparticles supported on boron diamond. These nickel-modified electrodes were shown as a potential analytical tool for the detection of alcohols [32].

The other fundamental expect of understanding screen printed electrodes reactivity is to explore the creation of surface oxygen functionalities, and the use of mediators on the electrode surface [4]. The next section will highlight some of the important advances made on screen printed electrodes creating oxygen functionalities and edge plane like sites, along with some chemical modifications.

4.1. Preanodized Screen Printed Carbon Electrode

The research on several carbon materials, especially carbon nanotubes and graphene, has revolutionized the scope of screen printed electrodes in electro-analysis. Compton's group observed enhanced electrocatalytic properties of CNT towards several target analytes attributed to the defect/edge plane-like sites [33,34]. Zen *et al*. demonstrated the creation of defect/edge-like sites and oxygen functionalities on the screen printed electrode surface through a simple pre-anodization process. Prasad *et al*. performed a comparative study on the role of oxygen functionalities and edge plane sites created achieved through a pre-anodization process and oxygen plasma treated screen printed electrodes [4]. Consequently, the pre-anodized electrode showed better characteristics in terms of reduction in overpotential and separation of oxidation peaks for the detection of uric acid and dopamine compared to the oxygen plasma-treated screen printed electrodes [35,36]. Similarly, the versatility of the pre-anodized screen printed electrode has been demonstrated in different media for enzymeless detection, detection of poorly electro-active analytes and direct electron transfer-related researches [37,38].

4.2. Mediator Integrated Screen Printed Electrode

There are many target analytes which have no significant electroactivity or near impossible to get electrochemical signals. In such cases, the use of an electrocatalyst mediator and methodologies to improve the sensor performance are very common. The appropriate selection of the mediator can improve the selectivity and lower the working potential for electrocatalysis. Redox mediators such as metal/metal complexes and pure organic polymers can be used as electrode modifiers. The viable approaches to immobilize the mediator on the screen-printed electrode include drop casting, physical attachment, and covalent binding or mixing into carbon paste. The mediator mixing approach has found more applications, and has been the mainly used technique for many decades, since the pioneering work of Adams *et al*., Various types of mediators such as Meldola's Blue, Prussian Blue, crown ethers, cobalt phthalocyanine and nickel hexacyanoferrate have been successfully integrated into the screen printed ink to design sensors for many target analytes. Ionic liquids have been used in analytical chemistry and carbon composites due to their physiochemical properties and biocompatible nature [39,40], but they were never been used in screen printed electrodes fabrication until the notable work by Ping *et al*., who incorporated a variety of ionic liquids into screen printed electrodes [41]. This work was further extended by Ren *et al*. in the fabrication of DNA sensors to achieve nano-level sensitivity [42]. Recently, carbon nanotubes-mediated screen printed electrodes have been used to increase the electrochemically active area of screen printed electrodes,

subsequently employed in the detection of *p*-aminophenol [43]. This work has provided a base to use other carbon nanotubes such as single walled carbon nanotubes and multi-walled carbon nanotube derivatives in designing screen printed carbon electrodes, with the possibility of accessing mass produced and reproducible nanotube-modified screen printed electrodes.

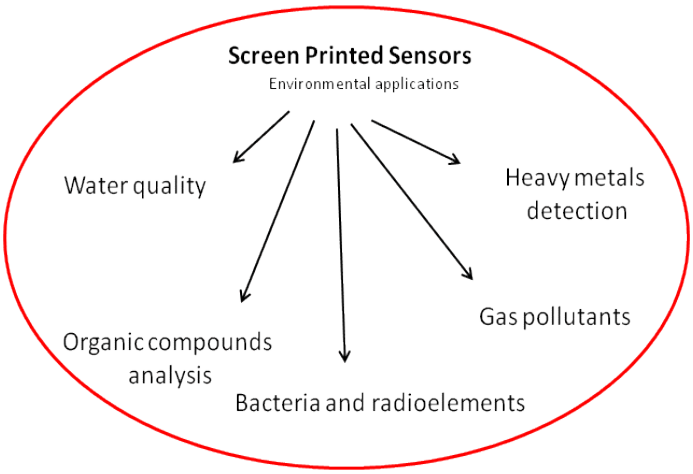
4.3. Metal Oxide Based Screen Printed Electrodes

Despite their various advantages, mediator approaches have the drawback of instability, and the fact they are not easily mass produced reproducibly, posing the problem of optimization of the sensors. As an alternative, metal oxides, including ruthenium oxide, copper oxide, nickel oxide, manganese oxide and bismuth oxide, have been used in the modification of electrodes [8,44], and the subsequently modified electrodes were used for many sensing applications. Among all the oxides, bismuth oxide is well documented to enhance the electro-analytical performance of the sensors, with reduced toxic effects. Since the pioneering work of Wang *et al.*, bismuth-modified electrodes have been extensively explored for diverse applications. Bismuth nanopowder immobilized with Nafion, and electrochemically oxidized bismuth oxides are normally used to obtain a uniform layer of bismuth on the electrode surface. The electrochemical deposition of the target species on the bismuth layer rather than the underlying graphite electrode results in the improved analytical performance of the sensors. Bismuth domains, as preferential nucleation sites, are distributed across the electrode surface producing their own diffusion zones and minimizing the effect of surface coverage which may result in an increased electron transfer resistance thus reducing the sensor sensitivity [45,46]. An advantageous approach for screen printed electrode modification is the one in which any metal oxide can be readily incorporated onto an electrode surface, allowing a true platform technology.

5. Screen Printed Sensors for Environmental Monitoring

Screen printed electrodes have been employed as a tool to design disposable and portable electrochemical sensors for environmental monitoring, such as water quality tests, organic compound analyses, heavy metals detection and gas pollutants (Figure 3).

Figure 3. Potential environmental applications of screen printed sensors.



5.1. Water Quality Tests

It is of vital importance to monitor source water and the aquatic systems that can be contaminated by industrial waste, sewage treatment plants and runoff from urban and agricultural lands. Water quality monitoring is mainly based on the measurement of the physical, chemical and bacteriological characteristics of water. The physical elements of analysis include monitoring of temperature, pH and conductivity, while chemical analyses measure oxygen, alkalinity, nitrogen and phosphorus compounds. Thousands of emerging contaminants may be present in water resources, including those used for drinking water production, therefore, the development of cost-effective devices for on-line and continuous monitoring of water quality is highly desirable. In this context, the use of screen printed electrodes to monitor pH changes has attracted great attention to replace the commonly used methods for testing pH changes. Koncki *et al*., designed a plastic, fully screen-printed, disposable pH sensor based on ruthenium dioxide by the application of a thick-film technology. The electrodes enable fast measurements, with good sensitivity in acidic and neutral solutions [47]. pH sensors with both printed reference and working electrodes on one substrate are reported in the literature [46], however this concept led to printing of the three electrode system for pH sensing with enhanced sensitivity as compared to the two electrode system [48]. Kampouris *et al*. designed a screen printed pH sensor by incorporating the pH sensitive phenanthraquinone moiety which undergoes a Nernstian potential shift with pH, and the pH insensitive dimethylferrocene one which acts as an internal reference. This generic approach offered a calibration-less and reproducible approach for portable pH measurements with the possibility of miniaturization, allowing incorporation into existing sensing devices [49]. Betelu *et al*., investigated the applicability of CeO_2-based screen printed electrodes for monitoring the pH of the COx pore water. However, this study was limited to the pH range between 5.5 and 13.2 [50]. Xiong *et al.* designed a calibration-less sensor based on nitrosophenyl-modified edge plane pyrolytic electrodes and screen printed electrodes to monitor pH changes [51]. Dissolved oxygen concentration is also another parameter that can be employed to test water quality. Zen *et al*., developed an efficient photocatalytic amperometric sensor for the determination of dissolved oxygen in phosphate buffer solution using a disposable copper-plated screen-printed carbon electrode. Real sample assays for groundwater and tap water were also consistent with those measured by a commercial dissolved oxygen meter [52]. Various modified screen printed sensors have been demonstrated as potential candidates to measure the chemical oxygen demand and biochemical chemical oxygen demand for various environmental studies [53].

Nitrate is also an important analyte for environmental and human health monitoring thus its detection and quantification is very important. In this regards, some screen printed electrodes have been designed and used to detect low levels of toxic ions. Moreover, microelectrodes in combination with screen printing technology have been employed to measure the nitrate level in water samples [54,55]. For example, Lin *et al*., fabricated poly (3,4-ethylenedioxythiophene) and PEDOT/multi-wall carbon nanotubes (PEDOT/MWCNTs)-modified screen-printed carbon electrodes (SPCEs) and studied their catalytic properties for nitrite measurement. The developed sensor was also applied to the determination of nitrite concentration in tap water samples [56]. Monchindu *et al*., electropolymerised aniline doped with polyvinyl sulphonate on screen printed carbon electrodes. The designed electrochemical sensors exhibited good analytical characteristics for nitrate detection [57]. Lin *et al*., investigated the oxidative electrochemistry of nitrite on a poly(3,4-ethylenedioxythiophene)/iron

phthalocyanine/multi-wall carbon nanotubes-modified screen-printed carbon electrode. The developed sensor was also applied for the determination of nitrite in tap water samples [58]. Metters *et al*. reported the fabrication of screen printed graphite micro-band electrodes which were electrochemically characterized and critically explored in electro-analytical applications for the sensing of nitrite [55] .Saljukis *et al*., fabricated manganese dioxide screen printed graphite electrodes for electro-analytical sensing purposes. The prepared sensors exhibited attractive performances as electrocatalysts for the sensing of nitrite ions with detection limits comparable or lower than those obtainable with other electrochemical sensors [59].

Similarly, the hydrophilic nature of phosphate ions makes them difficult to detect in water analysis. Ion selective electrodes, due to their ability to measure various species in turbid and colored medium, have been appeared as the prominent tools to measure phosphate ions for routine water sample analysis. Screen printed carbon paste and conventional PVC membrane electrodes have been integrated in ion selective sensors for phosphate ion analysis [60]. Some of the recently developed screen printed sensors for water quality are listed in Table 1.

Table 1. Some of the recently developed screen printed sensors for water quality tests.

Analyte	Modifier	Detection Method	Ref.
Liquids	Iridium and ruthenium oxide	pH sensor	[61]
Liquids	Phenanthraquinone moiety	pH sensor	[49]
Hydroxide ions	Nickel oxide bulk	pH sensor	[48]
Dissolved oxygen	CdS modified	Cathodic electrochemiluminescenc	[53]
Nitrite	Poly(dimethylsiloxane)	Amperometric detection	[54]
Nitrite	Shallow recessed unmodified	Amperometric detection	[25]
Phosphate	Bisthiourea ionophores	Amperometric detection	[60]
Nitrite	Carbon Black	Multi-electrochemical methods	[62]
Phosphate	Electrocatalyst cobalt phthalocyanine	Amperometric	[63]
Phosphate	Cobalt phthalocyanine	Amperometric	[64]
Nitrate	Modified screen printed electrodes	Electrochemical detection	[65]
Nitrate	polymer (poly(vinyl alcohol)) modified	Amperometric	[66]
Nitrate	commercial screen-printed electrochemical cell	Amperometric	[67]

5.2. Organic Compounds

Phenols are organic compounds broadly employed in the chemical, petrochemical, pharmaceutical, pesticide, pulp and paper, textile, metallurgic, resin and plastic, and pulp and paper industries. Phenol poisoning by skin absorption, inhalation of vapors or ingestion causes accumulation and damage to the brain, kidneys, liver, muscle, and eyes, as well as necrosis [21]. Therefore, detection of phenolic compounds and their derivatives is highly desirable to meet the corresponding environmental challenges. Even though the standardized methods are able to obtain accurate results for a wide range of phenolic compounds, conventional approaches are time-consuming and cost-intensive. Furthermore, they require large volumes of toxic organic solvents such as methylene chloride, acetone, and

methanol. Consequently, there is a demand for the development of reliable, portable, sensitive, simple and cost-effective methods for the fast detection of phenolic compounds. Electrochemical sensors based on screen printed electrodes have been used as low cost, simple, sensitive and disposable tools for *in situ* monitoring of phenolic compounds. The possibility of direct electrochemical oxidation of these phenolic compounds at the screen printed electrode facilitates their detection [68], Many modification strategies and immobilization methods have been reported in the literature to design innovative electrochemical sensors for monitoring phenolic compounds [69–72].

Pesticides are released intentionally into the environment, and through various processes can contaminate the environment. Although pesticides are associated with many health hazards, there is a lack of monitoring of these contaminants. Traditional chromatographic methods—high-performance liquid chromatography, capillary electrophoresis, and mass spectrometry—are effective for the analysis of pesticides in the environment, but have certain limitations such as complexity, time-consuming sample preparation, and the requirement of expensive apparatus and trained persons to operate them. Over the past decades, acetylcholinesterase (AChE) inhibition-based biosensors have emerged as simple, rapid, and ultra-sensitive tools for pesticide analysis in environmental monitoring, food safety, and quality control. These biosensors have the potential to complement or replace the classical analytical methods by simplifying or eliminating sample preparation and making field-testing easier and faster with a significant decrease in cost per analysis [73,74]. Based on the inhibition mechanism of the pesticide, various electrochemical biosensors based on screen printed electrodes have been constructed to analyse water and soil samples for the presence of pesticides. Prussian Blue, carbon nanotubes, cobalt phthalocyanine and conductive polymers have been successfully integrated as mediators in screen printed electrochemical biosensors for pesticide detection [75–78].

Despite the use of modern, less persistent agrochemicals, herbicide residues and herbicide metabolites in water are a serious environmental problem. Even when used appropriately, water soluble herbicides can be found in surface waters, ground waters, and tap water. For this reason, the monitoring of herbicides and herbicide metabolites is important to ensure the quality of water. Electrochemical immunosensors based on screen printed carbon electrodes are used for single shot determination of herbicides, eliminating the cleaning and reuse of components [79,80]. However, the immunoassays undergo some drawbacks such as the time consuming antibody production process and the possibility of cross-reactivity. Alternatively, photosynthetic electrochemical biosensors based on screen printed electrodes have been proposed, and successfully implemented for herbicide detection [81–83]. Polycyclic aromatic hydrocarbons (PAHs) are a large group of organic compounds with two or more fused aromatic rings. They have a relatively low solubility in water, but are highly lipophilic. Aromatic compounds can interact with graphite walls and thus stack onto carbon materials through non-covalent binding. After concentrating polyaromatic hydrocarbons on screen printed electrodes, an operating potential can be applied for the individual electrochemical detection of a specific aromatic compound [84–86]. Similarly, immunosensor approaches based on screen printed electrodes have also been reported in the literature for a mixture of individual polyaromatic hydrocarbon compounds. Nevertheless, the very similar structures of polyaromatic hydrocarbon compounds make difficult the production of a specific antibody for only one polyaromatic hydrocarbons. Future work may focus on the integration of various antibodies within a single screen printed sensor to get different signals and detailed information regarding polyaromatic hydrocarbon mixtures. Sensitive and decentralized analysis of antibiotic residues

in environmental samples is also high desirable. Tetracyclines are important classes of antibiotic that are detected by employing screen printed electrodes. Immunoassays in combination with nanoparticles on screen printed electrodes have shown great potential in the development of high sample throughput screening systems for antibiotics in environmental samples [87,88]. Table 2 provides examples of the some of the recently developed screen printed sensors for organic compounds (Table 2).

Table 2. Examples of the some of the recently developed screen printed sensors for organic compounds detection in environmental samples.

Analyte	Modifier	Detection Method	Ref.
Organophosphate	Poly(3,4-ethylenedioxythiophene) (PEDOT)	Amprometric	[76]
Organophosphate pesticides	Cobalt phthalocyanine	Chronoamperometry	[89]
Organophosphorus	Cysteamine self-assembled monolayer	Amprometric	[90]
Organophosphorus and Carbamate Pesticides	Unmodified	Amperometry, flow system	[91]
Aminophenol isomers	Untreated SPCE	Voltammetric	[21]
Organophosphorus Pesticide	Single-walled carbon nanotubes—Co phtalocyanine	Amperometry	[79]
Organophosphorus Pesticide Dichlofenthion	Nanometer-Sized Titania	Photoelectrochemical	[81]
Herbicide isoproturon	Unmodified	Amperometric	[92]
Herbicide	Magnetic nanoparticles	Amperometric	[83]
Picric acid and atrazine	Self-assembled monolayer	Photo-electrochemical	[93]
Chlorsulfuron	Gold (Au) metal ions	Stripping voltammetry	[80]
Phenol and catechol	Bismuth nanoparticles	Amperometric measurements	[94]
Phenol and pesticide	Iridium oxide nanoparticles	Electrochemical measurement	[95]
Phenol	Carbon Black Paste	Amperometric	[96]
Phenolic compounds	Nano-HA-chitosan nanocomposite-modified gold electrode	Amperometric	[97]
Phenolic compounds	Screen-printed PEDOT:PSS electrodes	Amperometric	[98]
Carbamate Insecticide	Prussian Blue-Multi-Walled Carbon Nanotubes	Amperometric	[99]

5.3. Heavy Metals

Due to the major negative impact of heavy metal ions toward human health and the environment, even at low concentrations, the development of simple, fast and not expensive detection methods for heavy metals is a major challenge for scientists. Among the different analytical methods for the analysis of heavy metal ions, the methods based on electrochemical sensors are widely applied for the detection of metals. Among toxic heavy metals, lead continues to be one of the most problematic. Despite considerable efforts to identify and eliminate Pb exposure sources, this metal still remains a significant health concern. Pb(II) is one of the heavy metals that has been detected with improved sensitivity by using modified carbon, bismuth, gold or other materials. These modifiers were integrated onto the surface of screen printed electrodes to make portable and disposable devices, improving their

suitability for on-site analysis [100–103]. However, such approaches exhibit shortcomings such as necessity of acid or alkaline working media. Alternatively, mercury has the potential to perform analysis over a wide range of pH, and can be used as a possible screen printed electrode modifier for trace metal detections. As a proof of concept, mercury-modified screen printed electrodes have been employed to detect low levels of Cd(II) in environmental samples [104,105].

Mercury ions, the most stable form of inorganic mercury, are highly toxic environmental pollutants and have serious medical effects. Therefore, it is highly desirable to develop sensitive methods for the detection of Hg^{2+}. Indeed, there have been numerous reports on optical Hg^{2+} detection by using Hg^{2+} sensitive fluorophores or chromophores, however, most of these fluorophores or chromophore-based Hg^{2+} sensors only work in organic media, which cannot be directly used to detect mercury contaminants in aqueous media. Bare gold or modified gold electrodes are normally used for the detection of Hg^{2+} due to their strong affinity for Hg^{2+} [106,107]. Commercial screen printed electrodes are reported for simple detection of Hg^{2+} in water samples. Nanostructured carbon black and screen printed electrodes modified with conducting polymer layers have also been designed for the trace level measurement of Hg^{2+} in water samples [108,109]. Arsenic is also a common compound found in drinking water, especially in some Asian countries. The toxicity of arsenic is greatly dependent on its oxidation state since As(III) is 50 times more toxic than arsenate due to its reactions with enzymes involved in human metabolism. Many detection methods have been developed for determination of such levels of arsenic. Among these methods, electrochemical methods provide accurate measurements of low concentrations of metal ions at ppb levels with rapid analysis times and low cost instrumentation. Screen printed electrodes modified with nanoparticles have been utilized to detect arsenic in water environments [110]. However, to avoid interferences from other metals, enzymatic biosensors based on screen printed electrodes for the measurement of arsenic in water samples have also been reported in the literature [111]. Designing screen printed sensors for simultaneous detection of various metals is also interesting for time and cost reasons. Screen printed electrodes modified with gold nanoparticles/gold films have also been reported for stripping voltammetric determination of mercury (II) and lead (II) [112]. Selected and recently developed screen printed sensors for heavy metals detection are listed in Table 3.

Table 3. Selected and recently developed sreen printed sensors for heavy metal detections.

Analyte	Modifier	Detection Method	Ref.
Pb^{2+} and Cd^{2+}	screen-printed antimony and tin	anodic stripping detection	[113]
Cu^{2+}	Macrocyclic Polyamine Modified Screen-Printed Electrodes	Square wave anodic stripping voltammetry	[114]
Cd^{2+}, Cu^{2+}	Diazonium modified electtrodes	Amperometric detection	[115]
Pb^{2+} and Cd^{2+}	Bismuth-coated	Stripping voltammetry	[116]
Pb^{2+}	Reduced graphene oxide	Square wave anodic stripping voltammetry	[117]
Zn^{2+}, Cd^{2+} and Pb^{2+}	Multiwalled carbon nanotubes	Differential pulse stripping voltammetry	[118]
Hg^{2+} and Pb^{2+}	Polypyrrole/carbonaceous nanospheres	Square wave anodic stripping voltammetry	[119]
Pb^{2+} and Cd^{2+}	Bismuth–carbon nanocomposites	Differential electrochemical methods	[120]

Table 3. *Cont.*

Analyte	Modifier	Detection Method	Ref.
Pb^{2+}	Bismuth-antimony film	Stripping voltammetric	[121]
Pb^{2+}	4-carboxyphenyl-grafted	Anodic Square Wave Voltammetry	[122]
As(III)	Gold electrode	Sequential injection/anodic stripping voltammetry	[123]
As(III)	Nanoparticles	Linear sweep voltammetric	[124]
As(III)	Modified screen printed electrodes	Amperometric	[111]
Cd^{2+}, Pb^{2+}, Cu^{2+} and Hg^{2+} ions	Heated graphitenanoparticle	Electrochemical stripping	[125]
Hg^{2+}	Gold nanoparticles-modified	Square wave anodic stripping voltammetry	[126]
Pb^{2+}, Cu^{2+} and Cd^{2+}	Mercury nano-droplets	Square wave anodic stripping voltammetry	[127]
Pb^{2+}	Paper disk impregnated	One-step electrochemical detection	[128]
Cd^{2+}	Nafion. Cd	Square Wave Anodic Stripping Voltammetry	[129]

5.4. Gas Pollutants

The air pollution caused by exhaust gases from automobiles has become a critical issue. In some regions, fossil fuel combustion is a problem as well. The principal gases that cause air pollution from automobiles are nitrogen oxide and carbon monoxide. Conventional and traditional methods to detect the levels of toxic gases include color reactions, chemiluminescence and IR absorption approaches. In comparison to these described methods, electrochemical gas sensor based on screen printed electrodes can provide low cost, easy to use and portable devices for environmental analysis.

Carbon monoxide is a colorless, odorless, tasteless and poisonous gas mainly produced by the combustion of fossil fuels. Bare gold, nanoparticles particles-modified carbon and SnO_2-modified carbon electrodes based on screen printing technology have been employed to detect the levels of this gas in environmental samples [130,131]. These devices have the potential to be used for *in situ* measurement and for continuous monitoring. Nitrogen oxide is a prominent air pollutant produced during high temperature combustion processes. The symptoms of nitrogen oxide poising appear several hours after its inhalation and require sensitive methods to detect it at low levels. Tin-doped and indium oxide thin films on screen printed electrodes have been used for the detection of nitrogen oxide in the air samples [132,133]. Volatile organic compounds including formaldehyde, acetone and methanol pose harmful effects to human health and contaminate the environment. Screen printed nanocomposite films integrated with multi-walled carbon nanotubes and silicon binders have been used to measure organic gases [134,135].

5.5. Other Environmental Pollutants

The presence of bacteria may also pose some enteric disease problems and indirectly results in economic losses. Enzyme-labeled and impedimetric immunoassays based on screen printed electrodes

have been developed for bacterial detection in river and tap water samples without pre-concentration steps [136]. Screen printed micro-system have been designed for pathogen detection that serve as both functional and structural components, to improve the simplicity of the fabrication steps [137]. Radio-elements are also considered as radiological and chemical toxic compounds, with their presence in the aquatic system needs to be monitored. Screen printing technology has efficiently contributed in the detection of radioelements, and screen printed sensors to monitor uranium have been reported in the literature [138,139].

6. Conclusions and Future Prospects

As discussed in this review paper, there have been many exciting developments in the use of screen printing to design new types of electrochemical sensors. The combination of modern electrochemical systems with screen printing technology along with breakthroughs in micro-electronoics and miniaturization permits the introduction of powerful and potential analytical tools for effective monitoring of environmental pollutants. Such real time on-site monitoring methodologies have successfully addressed the time constraints associated with classical laboratory analysis. With the passage of time, electrochemical devices are becoming more and more sophisticated and versatile while dramatically shrinking in size and weight. Screen printed methodologies offer the advantage of production of simple, economical, disposable, portable and mass produced devices suitable for on-site analysis of environmental pollutants. Disposable screen printed electrodes have extensively improved the sensitivity and selectivity of the analytical approaches, especially in the detection of certain environmental analytes that were difficult and challenging to measure with conventional and traditional techniques. The field of screen printed electrodes, however, continues to grow and find new application domains. It is expected that future work shall focus on the integration of nanomaterials in the screen printed electrodes to improve the electron transfer rates, thus enhancing the analytical performance of the sensors. Furthermore, microchip formats may find application to improve the miniaturization process to decrease the analysis time, sample volumes and reagent consumption and enhance portability and for on-site analysis.

Conflicts of Interest

The authors declare no conflict of interest.

References

1. Wang, J. Real-Time Electrochemical Monitoring: Toward Green Analytical Chemistry. *Acc. Chem. Res.* **2002**, *35*, 811–816.
2. Anastas, P.T. Green Chemistry and the Role of Analytical Methodology Development. *Crit. Rev. Anal. Chem.* **1999**, *29*, 167–175.
3. Workman, J.; Creasy, K.E.; Doherty, S.; Bond, L.; Koch, M.; Ullman, A.; Veltkamp, D.J. Process Analytical Chemistry. *Anal. Chem.* **2001**, *73*, 2705–2718.
4. Thiyagarajan, N.; Chang, J.-L.; Senthilkumar, K.; Zen, J.-M. Disposable electrochemical sensors: A mini review. *Electrochem. Commun.* **2014**, *38*, 86–90.

5. Honeychurch, K.C.; Hart, J.P. Screen-printed electrochemical sensors for monitoring metal pollutants. *TrAC, Trends Anal. Chem.* **2003**, *22*, 456–469.
6. Bakker, E.; Bühlmann, P.; Pretsch, E. Polymer Membrane Ion-Selective Electrodes—What are the Limits? *Electroanalysis* **1999**, *11*, 915–933.
7. Wang, J. Remote electrochemical sensors for monitoring inorganic and organic pollutants. *TrAC, Trends Anal. Chem.* **1997**, *16*, 84–88.
8. Metters, J.P.; Kadara, R.O.; Banks, C.E. New directions in screen printed electroanalytical sensors: An overview of recent developments. *Analyst* **2011**, *136*, 1067–1076.
9. Li, M.; Li, Y.-T.; Li, D.-W.; Long, Y.-T. Recent developments and applications of screen-printed electrodes in environmental assays—A review. *Anal. Chim. Acta* **2012**, *734*, 31–44.
10. Renedo, O.D.; Alonso-Lomillo, M.A.; Martínez, M.J.A. Recent developments in the field of screen-printed electrodes and their related applications. *Talanta* **2007**, *73*, 202–219.
11. Wang, J.; Lu, J.; Hocevar, S.B.; Farias, P.A.M.; Ogorevc, B. Bismuth-Coated Carbon Electrodes for Anodic Stripping Voltammetry. *Anal. Chem.* **2000**, *72*, 3218–3222.
12. Murray, R.W.; Ewing, A.G.; Durst, R.A. Chemically modified electrodes. Molecular design for electroanalysis. *Anal. Chem.* **1987**, *59*, 379A–390A.
13. Hayat, A.; Yang, C.; Rhouati, A.; Marty, J. Recent Advances and Achievements in Nanomaterial-Based, and Structure Switchable Aptasensing Platforms for Ochratoxin A Detection. *Sensors* **2013**, *13*, 15187–15208.
14. Hayat, A.; Andreescu, S.; Marty, J.-L. Design of PEG-aptamer two piece macromolecules as convenient and integrated sensing platform: Application to the label free detection of small size molecules. *Biosens. Bioelectron.* **2013**, *45*, 168–173.
15. Hayat, A.; Haider, W.; Rolland, M.; Marty, J.-L. Electrochemical grafting of long spacer arms of hexamethyldiamine on a screen printed carbon electrode surface: Application in target induced ochratoxin A electrochemical aptasensor. *Analyst* **2013**, *138*, 2951–2957.
16. Heller, A.; Feldman, B. Electrochemical Glucose Sensors and Their Applications in Diabetes Management. *Chem. Rev.* **2008**, *108*, 2482–2505.
17. Newman, J.D.; Turner, A.P.F. Home blood glucose biosensors: A commercial perspective. *Biosens. Bioelectron.* **2005**, *20*, 2435–2453.
18. Wilson, R.; Turner, A.P.F. Glucose oxidase: An ideal enzyme. *Biosens. Bioelectron.* **1992**, *7*, 165–185.
19. Hart, J.P.; Wring, S.A. Recent developments in the design and application of screen-printed electrochemical sensors for biomedical, environmental and industrial analyses. *TrAC, Trends Anal. Chem.* **1997**, *16*, 89–103.
20. Hayat, A.; Marty, J.-L.; Radi, A.-E. Novel Amperometric Hydrogen Peroxide Biosensor Based on Horseradish Peroxidase Azide Covalently Immobilized on Ethynyl-Modified Screen-Printed Carbon Electrode via Click Chemistry. *Electroanalysis* **2012**, *24*, 1446–1452.
21. Su, W.-Y.; Wang, S.-M.; Cheng, S.-H. Electrochemically pretreated screen-printed carbon electrodes for the simultaneous determination of aminophenol isomers. *J. Electroanal. Chem.* **2011**, *651*, 166–172.
22. Hayat, A.; Barthelmebs, L.; Marty, J.L. Electrochemical impedimetric immunosensor for the detection of okadaic acid in mussel sample. *Sens. Actuators B Chem.* **2012**, *171*, 810–815.

23. Hayat, A.; Sassolas, A.; Marty, J.L.; Radi, A. Highly sensitive ochratoxin A impedimetric aptasensor based on the immobilization of azido-aptamer onto electrografted binary film via click chemistry. *Talanta* **2013**, *103*, 14–19.
24. Fan, Z.; Ho, J.C.; Takahashi, T.; Yerushalmi, R.; Takei, K.; Ford, A.C.; Chueh, Y.-L.; Javey, A. Toward the Development of Printable Nanowire Electronics and Sensors. *Adv. Mater.* **2009**, *21*, 3730–3743.
25. Khairy, M.; Kadara, R.O.; Banks, C.E. Electroanalytical sensing of nitrite at shallow recessed screen printed microelectrode arrays. *Anal. Methods* **2010**, *2*, 851–854.
26. Kadara, R.O.; Jenkinson, N.; Banks, C.E. Screen printed recessed microelectrode arrays. *Sens. Actuators B* **2009**, *142*, 342–346.
27. Fanjul-Bolado, P.; Hernández-Santos, D.; Lamas-Ardisana, P.J.; Martín-Pernía, A.; Costa-García, A. Electrochemical characterization of screen-printed and conventional carbon paste electrodes. *Electrochim. Acta* **2008**, *53*, 3635–3642.
28. Šljukić, B.; Malakhova, N.A.; Brainina, K.Z.; Banks, C.E.; Compton, R.G. Screen Printed Electrodes and Screen Printed Modified Electrodes Benefit from Insonation. *Electroanalysis* **2006**, *18*, 928–930.
29. Choudhry, N.A.; Kampouris, D.K.; Kadara, R.O.; Jenkinson, N.; Banks, C.E. Next generation screen printed electrochemical platforms: Non-enzymatic sensing of carbohydrates using copper (ii) oxide screen printed electrodes. *Anal. Methods* **2009**, *1*, 183–187.
30. Choudhry, N.A.; Kadara, R.O.; Jenkinson, N.; Banks, C.E. Screen printed electrodes provide micro-domain sites for fabricating disposable electro-catalytic ensembles. *Electrochem. Commun.* **2010**, *12*, 406–409.
31. Davies, T.J.; Hyde, M.E.; Compton, R.G. Nanotrench Arrays Reveal Insight into Graphite Electrochemistry. *Angew. Chem.* **2005**, *117*, 5251–5256.
32. Banks, C.E.; Compton, R.G. New electrodes for old: From carbon nanotubes to edge plane pyrolytic graphite. *Analyst* **2006**, *131*, 15–21.
33. Banks, C.E.; Crossley, A.; Salter, C.; Wilkins, S.J.; Compton, R.G. Carbon Nanotubes Contain Metal Impurities Which Are Responsible for the "Electrocatalysis" Seen at Some Nanotube-Modified Electrodes. *Angew. Chem. Int. Ed.* **2006**, *45*, 2533–2537.
34. Shi, K.; Shiu, K.-K. Determination of Uric Acid at Electrochemically Activated Glassy Carbon Electrode. *Electroanalysis* **2001**, *13*, 1319–1325.
35. Chen, J.C.; Chung, H.H.; Hsu, C.T.; Tsai, D.M.; Kumar, A.S.; Zen, J.M. A disposable single-use electrochemical sensor for the detection of uric acid in human whole blood. *Sens. Actuators B* **2005**, *110*, 364–369.
36. Tsai, T.-H.; Thiagarajan, S.; Chen, S.-M. Detection of Melamine in Milk Powder and Human Urine. *J. Agric. Food Chem.* **2010**, *58*, 4537–4544.
37. Yang, T.-H.; Hung, C.-L.; Ke, J.-H.; Zen, J.-M. An electrochemically preanodized screen-printed carbon electrode for achieving direct electron transfer to glucose oxidase. *Electrochem. Commun.* **2008**, *10*, 1094–1097.
38. Zen, J.-M.; SenthillKumar, A.; Tsai, D.-M. Recent Updates of Chemically Modified Electrodes in Analytical Chemistry. *Electroanalysis* **2003**, *15*, 1073–1087.

39. Kozub, B.R.; Compton, R.G. Voltammetric studies of the redox mediator, cobalt phthalocyanine, with regard to its claimed electrocatalytic properties. *Sens. Actuators B* **2010**, *147*, 350–358.
40. Ping, J.; Wu, J.; Ying, Y. Development of an ionic liquid modified screen-printed graphite electrode and its sensing in determination of dopamine. *Electrochem. Commun.* **2010**, *12*, 1738–1741.
41. Ren, R.; Leng, C.; Zhang, S. A chronocoulometric DNA sensor based on screen-printed electrode doped with ionic liquid and polyaniline nanotubes. *Biosens. Bioelectron.* **2010**, *25*, 2089–2094.
42. Lamas-Ardisana, P.J.; Queipo, P.; Fanjul-Bolado, P.; Costa-García, A. Multiwalled carbon nanotube modified screen-printed electrodes for the detection of p-aminophenol: Optimisation and application in alkaline phosphatase-based assays. *Anal. Chim. Acta* **2008**, *615*, 30–38.
43. Wang, J.; Naser, N.; Angnes, L.; Wu, H.; Chen, L. Metal-dispersed carbon paste electrodes. *Anal. Chem.* **1992**, *64*, 1285–1288.
44. Wang, J. Stripping Analysis at Bismuth Electrodes: A Review. *Electroanalysis* **2005**, *17*, 1341–1346.
45. Economou, A. Bismuth-film electrodes: Recent developments and potentialities for electroanalysis. *TrAC, Trends Anal. Chem.* **2005**, *24*, 334–340.
46. Müller, A.; Brinz, T.; Simon, U. Preparation and Measurement of Combinatorial Screen Printed Libraries for the Electrochemical Analysis of Liquids. *J. Comb. Chem.* **2008**, *11*, 138–142.
47. Koncki, R.; Mascini, M. Screen-printed ruthenium dioxide electrodes for pH measurements. *Anal. Chim. Acta* **1997**, *351*, 143–149.
48. Hallam, P.M.; Kampouris, D.K.; Kadara, R.O.; Jenkinson, N.; Banks, C.E. Nickel oxide screen printed electrodes for the sensing of hydroxide ions in aqueous solutions. *Anal. Methods* **2010**, *2*, 1152–1155.
49. Kampouris, D.K.; Kadara, R.O.; Jenkinson, N.; Banks, C.E. Screen printed electrochemical platforms for pH sensing. *Anal. Methods* **2009**, *1*, 25–28.
50. Betelu, S.; Polychronopoulou, K.; Rebholz, C.; Ignatiadis, I. Novel CeO_2-based screen-printed potentiometric electrodes for pH monitoring. *Talanta* **2011**, *87*, 126–135.
51. Xiong, L.; Batchelor-McAuley, C.; Compton, R.G. Calibrationless pH sensors based on nitrosophenyl and ferrocenyl co-modified screen printed electrodes. *Sens. Actuators B* **2011**, *159*, 251–255.
52. Zen, J.-M.; Song, Y.-S.; Chung, H.-H.; Hsu, C.-T.; Senthil Kumar, A. Photoelectrochemical Oxygen Sensor Using Copper-Plated Screen-Printed Carbon Electrodes. *Anal. Chem.* **2002**, *74*, 6126–6130.
53. Zheng, R.-J.; Fang, Y.-M.; Qin, S.-F.; Song, J.; Wu, A.-H.; Sun, J.-J. A dissolved oxygen sensor based on hot electron induced cathodic electrochemiluminescence at a disposable CdS modified screen-printed carbon electrode. *Sens. Actuators B* **2011**, *157*, 488–493.
54. Chang, J.-L.; Zen, J.-M. A poly(dimethylsiloxane)-based electrochemical cell coupled with disposable screen printed edge band ultramicroelectrodes for use in flow injection analysis. *Electrochem. Commun.* **2007**, *9*, 2744–2750.
55. Metters, J.P.; Kadara, R.O.; Banks, C.E. Electroanalytical properties of screen printed graphite microband electrodes. *Sens. Actuators B* **2012**, *169*, 136–143.
56. Lin, C.-Y.; Vasantha, V.S.; Ho, K.-C. Detection of nitrite using poly(3,4-ethylenedioxythiophene) modified SPCEs. *Sens. Actuators B* **2009**, *140*, 51–57.

57. Muchindu, M.; Waryo, T.; Arotiba, O.; Kazimierska, E.; Morrin, A.; Killard, A.J.; Smyth, M.R.; Jahed, N.; Kgarebe, B.; Baker, P.G.L.; *et al*. Electrochemical nitrite nanosensor developed with amine- and sulphate-functionalised polystyrene latex beads self-assembled on polyaniline. *Electrochimica Acta* **2010**, *55*, 4274–4280.
58. Lin, C.-Y.; Balamurugan, A.; Lai, Y.-H.; Ho, K.-C. A novel poly(3,4-ethylenedioxythiophene)/iron phthalocyanine/multi-wall carbon nanotubes nanocomposite with high electrocatalytic activity for nitrite oxidation. *Talanta* **2010**, *82*, 1905–1911.
59. Sljukic, B.R.; Kadara, R.O.; Banks, C.E. Disposable manganese oxide screen printed electrodes for electroanalytical sensing. *Anal. Methods* **2011**, *3*, 105–109.
60. Khaled, E.; Hassan, H.N.A.; Girgis, A.; Metelka, R. Construction of novel simple phosphate screen-printed and carbon paste ion-selective electrodes. *Talanta* **2008**, *77*, 737–743.
61. Muller, A.; Brinz, T.; Simon, U. Preparation and measurement of combinatorial screen printed libraries for the electrochemical analysis of liquids. *J. Comb. Chem.* **2009**, *11*, 138–42.
62. Malha, S.I.R.; Mandli, J.; Ourari, A.; Amine, A. Carbon Black-Modified Electrodes as Sensitive Tools for the Electrochemical Detection of Nitrite and Nitrate. *Electroanalysis* **2013**, *25*, 2289–2297.
63. Gilbert, L.; Jenkins, A.T.; Browning, S.; Hart, J.P. Development of an amperometric assay for phosphate ions in urine based on a chemically modified screen-printed carbon electrode. *Anal. Biochem.* **2009**, *393*, 242–247.
64. Gilbert, L.; Jenkins, A.T.A.; Browning, S.; Hart, J.P. Development of an amperometric, screen-printed, single-enzyme phosphate ion biosensor and its application to the analysis of biomedical and environmental samples. *Sens. Actuators B* **2011**, *160*, 1322–1327.
65. Karousos, N.; Chong, L.C.; Ewen, C.; Livingstone, C.; Davis, J. Evaluation of a multifunctional indicator for the electroanalytical determination of nitrite. *Electrochim. Acta* **2005**, *50*, 1879–1884.
66. Quan, D.; Shim, J.H.; Kim, J.D.; Park, H.S.; Cha, G.S.; Nam, H. Electrochemical Determination of Nitrate with Nitrate Reductase-Immobilized Electrodes under Ambient Air. *Anal. Chem.* **2005**, *77*, 4467–4473.
67. Plumeré, N.; Henig, J.; Campbell, W.H. Enzyme-Catalyzed O_2 Removal System for Electrochemical Analysis under Ambient Air: Application in an Amperometric Nitrate Biosensor. *Anal. Chem*. **2012**, *84*, 2141–2146.
68. Brugnera, M.F.; Trindade, M.A.G.; Zanoni, M.V.B. Detection of Bisphenol A on a Screen-Printed Carbon Electrode in CTAB Micellar Medium. *Anal. Lett.* **2010**, *43*, 2823–2836.
69. Li, D.; Li, D.-W.; Fossey, J.S.; Long, Y.-T. Portable Surface-Enhanced Raman Scattering Sensor for Rapid Detection of Aniline and Phenol Derivatives by On-Site Electrostatic Preconcentration. *Anal. Chem.* **2010**, *82*, 9299–9305.
70. Song, W.; Li, D.-W.; Li, Y.-T.; Li, Y.; Long, Y.-T. Disposable biosensor based on graphene oxide conjugated with tyrosinase assembled gold nanoparticles. *Biosens. Bioelectron.* **2011**, *26*, 3181–3186.
71. Alkasir, R.S.J.; Ganesana, M.; Won, Y.-H.; Stanciu, L.; Andreescu, S. Enzyme functionalized nanoparticles for electrochemical biosensors: A comparative study with applications for the detection of bisphenol A. *Biosens. Bioelectron.* **2010**, *26*, 43–49.

72. Ibarra-Escutia, P.; Gómez, J.J.; Calas-Blanchard, C.; Marty, J.L.; Ramírez-Silva, M.T. Amperometric biosensor based on a high resolution photopolymer deposited onto a screen-printed electrode for phenolic compounds monitoring in tea infusions. *Talanta* **2010**, *81*, 1636–1642.
73. Ben Oujji, N.; Bakas, I.; Istamboulié, G.; Ait-Ichou, I.; Ait-Addi, E.; Rouillon, R.; Noguer, T. Acetylcholinesterase Immobilized on Magnetic Beads for Pesticides Detection: Application to Olive Oil Analysis. *Sensors* **2012**, *12*, 7893–7904.
74. Gan, N.; Yang, X.; Xie, D.; Wu, Y.; Wen, W. A Disposable Organophosphorus Pesticides Enzyme Biosensor Based on Magnetic Composite Nano-Particles Modified Screen Printed Carbon Electrode. *Sensors* **2010**, *10*, 625–638.
75. Istamboulie, G.; Sikora, T.; Jubete, E.; Ochoteco, E.; Marty, J.-L.; Noguer, T. Screen-printed poly(3,4-ethylenedioxythiophene) (PEDOT): A new electrochemical mediator for acetylcholinesterase-based biosensors. *Talanta* **2010**, *82*, 957–961.
76. Won, Y.-H.; Jang, H.S.; Kim, S.M.; Stach, E.; Ganesana, M.; Andreescu, S.; Stanciu, L.A. Biomagnetic Glasses: Preparation, Characterization, and Biosensor Applications. *Langmuir* **2009**, *26*, 4320–4326.
77. Arduini, F.; Ricci, F.; Tuta, C.S.; Moscone, D.; Amine, A.; Palleschi, G. Detection of carbamic and organophosphorous pesticides in water samples using a cholinesterase biosensor based on Prussian Blue-modified screen-printed electrode. *Anal. Chim. Acta* **2006**, *580*, 155–162.
78. Kumar, J.; D'Souza, S.F. Microbial biosensor for detection of methyl parathion using screen printed carbon electrode and cyclic voltammetry. *Biosens. Bioelectron.* **2011**, *26*, 4289–4293.
79. Baskeyfield, D.E.H.; Davis, F.; Magan, N.; Tothill, I.E. A membrane-based immunosensor for the analysis of the herbicide isoproturon. *Anal. Chim. Acta* **2011**, *699*, 223–231.
80. Nangia, Y.; Bhalla, V.; Kumar, B.; Suri, C.R. Electrochemical stripping voltammetry of gold ions for development of ultra-sensitive immunoassay for chlorsulfuron. *Electrochem. Commun.* **2012**, *14*, 51–54.
81. Bhalla, V.; Zazubovich, V. Self-assembly and sensor response of photosynthetic reaction centers on screen-printed electrodes. *Anal. Chim. Acta* **2011**, *707*, 184–190.
82. Shitanda, I.; Takamatsu, S.; Watanabe, K.; Itagaki, M. Amperometric screen-printed algal biosensor with flow injection analysis system for detection of environmental toxic compounds. *Electrochim. Acta* **2009**, *54*, 4933–4936.
83. Zamaleeva, A.I.; Sharipova, I.R.; Shamagsumova, R.V.; Ivanov, A.N.; Evtugyn, G.A.; Ishmuchametova, D.G.; Fakhrullin, R.F. A whole-cell amperometric herbicide biosensor based on magnetically functionalised microalgae and screen-printed electrodes. *Anal. Methods* **2011**, *3*, 509–513.
84. Leyton, P.; Gómez-Jeria, J.S.; Sanchez-Cortes, S.; Domingo, C.; Campos-Vallette, M. Carbon Nanotube Bundles as Molecular Assemblies for the Detection of Polycyclic Aromatic Hydrocarbons: Surface-Enhanced Resonance Raman Spectroscopy and Theoretical Studies. *J. Phys. Chem. B* **2006**, *110*, 6470–6474.
85. Honeychurch, K.C.; Hart, J.P.; Kirsch, N. Voltammetric, chromatographic and mass spectral elucidation of the redox reactions of 1-hydroxypyrene occurring at a screen-printed carbon electrode. *Electrochim. Acta* **2004**, *49*, 1141–1149.

86. Fähnrich, K.A.; Pravda, M.; Guilbault, G.G. Disposable amperometric immunosensor for the detection of polycyclic aromatic hydrocarbons (PAHs) using screen-printed electrodes. *Biosens. Bioelectron.* **2003**, *18*, 73–82.
87. Masawat, P.; Slater, J.M. The determination of tetracycline residues in food using a disposable screen-printed gold electrode (SPGE). *Sens. Actuators B* **2007**, *124*, 127–132.
88. Centi, S.; Stoica, A.I.; Laschi, S.; Mascini, M. Development of an Electrochemical Immunoassay Based on the Use of an Eight-Electrodes Screen-Printed Array Coupled with Magnetic Beads for the Detection of Antimicrobial Sulfonamides in Honey. *Electroanalysis* **2010**, *22*, 1881–1888.
89. Crew, A.; Lonsdale, D.; Byrd, N.; Pittson, R.; Hart, J.P. A screen-printed, amperometric biosensor array incorporated into a novel automated system for the simultaneous determination of organophosphate pesticides. *Biosens. Bioelectron.* **2011**, *26*, 2847–2851.
90. Arduini, F.; Guidone, S.; Amine, A.; Palleschi, G.; Moscone, D. Acetylcholinesterase biosensor based on self-assembled monolayer-modified gold-screen printed electrodes for organophosphorus insecticide detection. *Sens. Actuators B* **2013**, *179*, 201–208.
91. Alonso, G.A.; Muñoz, R.; Marty, J.-L. Automatic Electronic Tongue for On-Line Detection and Quantification of Organophosphorus and Carbamate Pesticides Using Enzymatic Screen Printed Biosensors. *Anal. Lett.* **2012**, *46*, 1743–1757.
92. Ivanov, A.N.; Younusov, R.R.; Evtugyn, G.A.; Arduini, F.; Moscone, D.; Palleschi, G. Acetylcholinesterase biosensor based on single-walled carbon nanotubes—Co phtalocyanine for organophosphorus pesticides detection. *Talanta* **2011**, *85*, 216–221.
93. Li, H.; Li, J.; Yang, Z.; Xu, Q.; Hu, X. A novel photoelectrochemical sensor for the organophosphorus pesticide dichlofenthion based on nanometer-sized titania coupled with a screen-printed electrode. *Anal. Chem.* **2011**, *83*, 5290–5295.
94. Mayorga-Martinez, C.C.; Cadevall, M.; Guix, M.; Ros, J.; Merkoci, A. Bismuth nanoparticles for phenolic compounds biosensing application. *Biosens. Bioelectron.* **2013**, *40*, 57–62.
95. Mayorga, C.; Pino, F.; Kurbanoglua, S.; Rivas, L.; Ozkan, S.A.; Merkoci, A. Iridium oxide nanoparticles induced dual catalytic/inhibition based detection of phenol and pesticide compounds. *J. Mater. Chem. B* **2014**, *2*, 2233–2239.
96. Nadifiyine, S.; Haddam, M.; Mandli, J.; Chadel, S.; Blanchard, C.C.; Marty, J.L.; Amine, A. Amperometric Biosensor Based on Tyrosinase Immobilized on to a Carbon Black Paste Electrode for Phenol Determination in Olive Oil. *Anal. Lett.* **2013**, *46*, 2705–2726.
97. Lu, L.; Zhang, L.; Zhang, X.; Huan, S.; Shen, G.; Yu, R. A novel tyrosinase biosensor based on hydroxyapatite–chitosan nanocomposite for the detection of phenolic compounds. *Anal. Chim. Acta* **2010**, *665*, 146–151.
98. Moczko, E.; Istamboulie, G.; Calas-Blanchard, C.; Rouillon, R.; Noguer, T. Biosensor employing screen-printed PEDOT:PSS for sensitive detection of phenolic compounds in water. *J. Polym. Sci. Part A* **2012**, *50*, 2286–2292.
99. Chai, Y.; Niu, X.; Chen, C.; Zhao, H.; Lan, M. Carbamate Insecticide Sensing Based on Acetylcholinesterase/Prussian Blue-Multi-Walled Carbon Nanotubes/Screen-Printed Electrodes. *Anal. Lett*. **2013**, *46*, 803–817.
100. Arduini, F.; Calvo, J.Q.; Palleschi, G.; Moscone, D.; Amine, A. Bismuth-modified electrodes for lead detection. *TrAC, Trends Anal. Chem.* **2010**, *29*, 1295–1304.

101. Krystofova, O.; Trnkova, L.; Adam, V.; Zehnalek, J.; Hubalek, J.; Babula, P.; Kizek, R. Electrochemical Microsensors for the Detection of Cadmium(II) and Lead(II) Ions in Plants. *Sensors* **2010**, *10*, 5308–5328.
102. Honeychurch, K.C.; Al-Berezanchi, S.; Hart, J.P. The voltammetric behaviour of lead at a microband screen-printed carbon electrode and its determination in acetate leachates from glazed ceramic plates. *Talanta* **2011**, *84*, 717–723.
103. Mazumdar, D.; Liu, J.; Lu, G.; Zhou, J.; Lu, Y. Easy-to-use dipstick tests for detection of lead in paints using non-cross-linked gold nanoparticle-DNAzyme conjugates. *Chem. Commun.* **2010**, *46*, 1416–1418.
104. Zaouak, O.; Authier, L.; Cugnet, C.; Castetbon, A.; Potin-Gautier, M. Electroanalytical Device for Cadmium Speciation in Waters. Part 1: Development and Characterization of a Reliable Screen-Printed Sensor. *Electroanalysis* **2010**, *22*, 1151–1158.
105. Betelu, S.; Parat, C.; Petrucciani, N.; Castetbon, A.; Authier, L.; Potin-Gautier, M. Semicontinuous Monitoring of Cadmium and Lead with a Screen-Printed Sensor Modified by a Membrane. *Electroanalysis* **2007**, *19*, 399–402.
106. Giacomino, A.; Abollino, O.; Malandrino, M.; Mentasti, E. Parameters affecting the determination of mercury by anodic stripping voltammetry using a gold electrode. *Talanta* **2008**, *75*, 266–273.
107. Gong, J.; Zhou, T.; Song, D.; Zhang, L.; Hu, X. Stripping Voltammetric Detection of Mercury(II) Based on a Bimetallic Au−Pt Inorganic−Organic Hybrid Nanocomposite Modified Glassy Carbon Electrode. *Anal. Chem.* **2009**, *82*, 567–573.
108. Bernalte, E.; Sánchez, C.M.; Gil, E.P. Determination of mercury in ambient water samples by anodic stripping voltammetry on screen-printed gold electrodes. *Anal. Chim. Acta* **2011**, *689*, 60–64.
109. Somerset, V.S.; Hernandez, L.H.; Iwuoha, E.I. Stripping voltammetric measurement of trace metal ions using screen-printed carbon and modified carbon paste electrodes on river water from the Eerste-Kuils River System. *J. Environ. Sci. Health Part A* **2010**, *46*, 17–32.
110. Sanllorente-Méndez, S.; Domínguez-Renedo, O.; Arcos-Martínez, M.J. Determination of Arsenic(III) Using Platinum Nanoparticle-Modified Screen-Printed Carbon-Based Electrodes. *Electroanalysis* **2009**, *21*, 635–639.
111. Sanllorente-Méndez, S.; Domínguez-Renedo, O.; Arcos-Martínez, M.J. Immobilization of Acetylcholinesterase on Screen-Printed Electrodes. Application to the Determination of Arsenic(III). *Sensors* **2010**, *10*, 2119–2128.
112. Mandil, A.; Idrissi, L.; Amine, A. Stripping voltammetric determination of mercury(II) and lead(II) using screen-printed electrodes modified with gold films, and metal ion preconcentration with thiol-modified magnetic particles. *Microchim. Acta* **2010**, *170*, 299–305.
113. Maczuga, M.; Economou, A.; Bobrowski, A.; Prodromidis, M.I. Novel screen-printed antimony and tin voltammetric sensors for anodic stripping detection of Pb(II) and Cd(II). *Electrochim. Acta* **2013**, *114*, 758–765.
114. Andreuccetti, C.; Bettazzi, F.; Giorgi, C.; Laschi, S.; Marrazza, G.; Mascini, M.; Palchetti, I. Macrocyclic Polyamine Modified Screen-Printed Electrodes for Copper(II) Detection. In *Sensors*; Baldini, F., D'Amico, A., Di Natale, C., Siciliano, P., Seeber, R., De Stefano, L., Bizzarri, R., Andò, B., Eds.; Springer: New York, NY, USA, 2014; Volume 162, pp. 471–474.

115. Bouden, S.; Bellakhal, N.; Chaussé, A.; Vautrin-Ul, C. Performances of carbon-based screen-printed electrodes modified by diazonium salts with various carboxylic functions for trace metal sensors. *Electrochem. Commun.* **2014**, *41*, 68–71.
116. Chen, C.; Niu, X.; Chai, Y.; Zhao, H.; Lan, M. Bismuth-based porous screen-printed carbon electrode with enhanced sensitivity for trace heavy metal detection by stripping voltammetry. *Sens. Actuators B* **2013**, *178*, 339–342.
117. Jian, J.-M.; Liu, Y.-Y.; Zhang, Y.-L.; Guo, X.-S.; Cai, Q. Fast and Sensitive Detection of Pb2+ in Foods Using Disposable Screen-Printed Electrode Modified by Reduced Graphene Oxide. *Sensors* **2013**, *13*, 13063–13075.
118. Fu, L.; Li, X.; Yu, J.; Ye, J. Facile and Simultaneous Stripping Determination of Zinc, Cadmium and Lead on Disposable Multiwalled Carbon Nanotubes Modified Screen-Printed Electrode. *Electroanalysis* **2013**, *25*, 567–572.
119. Wei, Y.; Yang, R.; Liu, J.-H.; Huang, X.-J. Selective detection toward Hg(II) and Pb(II) using polypyrrole/carbonaceous nanospheres modified screen-printed electrode. *Electrochim. Acta* **2013**, *105*, 218–223.
120. Gich, M.; Fernandez-Sanchez, C.; Cotet, L.C.; Niu, P.; Roig, A. Facile synthesis of porous bismuth-carbon nanocomposites for the sensitive detection of heavy metals. *J. Mater. Chem. A* **2013**, *1*, 11410–11418.
121. Chen, C.; Niu, X.; Chai, Y.; Zhao, H.; Lan, M.; Zhu, Y.; Wei, G. Determination of Lead(II) Using Screen-Printed Bismuth-Antimony Film Electrode. *Electroanalysis* **2013**, *25*, 1446–1452.
122. Bouden, S.; Chaussé, A.; Dorbes, S.; El Tall, O.; Bellakhal, N.; Dachraoui, M.; Vautrin-Ul, C. Trace lead analysis based on carbon-screen-printed-electrodes modified via 4-carboxy-phenyl diazonium salt electroreduction. *Talanta* **2013**, *106*, 414–421.
123. Punrat, E.; Chuanuwatanakul, S.; Kaneta, T.; Motomizu, S.; Chailapakul, O. Method development for the determination of arsenic by sequential injection/anodic stripping voltammetry using long-lasting gold-modified screen-printed carbon electrode. *Talanta* **2013**, *116*, 1018–1025.
124. Khairy, M.; Kampouris, D.K.; Kadara, R.O.; Banks, C.E. Gold Nanoparticle Modified Screen Printed Electrodes for the Trace Sensing of Arsenic(III) in the Presence of Copper(II). *Electroanalysis* **2010**, *22*, 2496–2501.
125. Aragay, G.; Pons, J.; Merkoci, A. Enhanced electrochemical detection of heavy metals at heated graphite nanoparticle-based screen-printed electrodes. *J. Mater. Chem.* **2011**, *21*, 4326–4331.
126. Bernalte, E.; Marín Sánchez, C.; Pinilla Gil, E. Gold nanoparticles-modified screen-printed carbon electrodes for anodic stripping voltammetric determination of mercury in ambient water samples. *Sens. Actuators B* **2012**, *161*, 669–674.
127. Song, W.; Zhang, L.; Shi, L.; Li, D.-W.; Li, Y.; Long, Y.-T. Simultaneous determination of cadmium(II), lead(II) and copper(II) by using a screen-printed electrode modified with mercury nano-droplets. *Microchim. Acta* **2010**, *169*, 321–326.
128. Fang, H.-L.; Zheng, H.-X.; Ou, M.-Y.; Meng, Q.; Fan, D.-H.; Wang, W. One-step sensing lead in surface waters with screen printed electrode. *Sens. Actuators B* **2011**, *153*, 369–372.

129. Henríquez, C.; Laglera, L.M.; Alpizar, M.J.; Calvo, J.; Arduini, F.; Cerdà, V. Cadmium determination in natural water samples with an automatic multisyringe flow injection system coupled to a flow-through screen printed electrode. *Talanta* **2012**, *96*, 140–146.
130. Morata, A.; Viricelle, J.P.; Tarancón, A.; Dezanneau, G.; Pijolat, C.; Peiro, F.; Morante, J.R. Development and characterisation of a screen-printed mixed potential gas sensor. *Sens. Actuators B* **2008**, *130*, 561–566.
131. Chou, C.-H.; Chang, J.-L.; Zen, J.-M. Effective analysis of gaseous formaldehyde based on a platinum-deposited screen-printed edge band ultramicroelectrode coated with Nafion as solid polymer electrolyte. *Sens. Actuators B* **2010**, *147*, 669–675.
132. Fergus, J.W. Materials for high temperature electrochemical NOx gas sensors. *Sens. Actuators B* **2007**, *121*, 652–663.
133. Mbarek, H.; Saadoun, M.; Bessaïs, B. Porous screen printed indium tin oxide (ITO) for NOx gas sensing. *Phys. Status Solidi C* **2007**, *4*, 1903–1907.
134. Khadayate, R.S.; Sali, J.V.; Patil, P.P. Acetone vapor sensing properties of screen printed WO_3 thick films. *Talanta* **2007**, *72*, 1077–1081.
135. Cantalini, C.; Valentini, L.; Lozzi, L.; Armentano, I.; Kenny, J.M.; Santucci, S. NO_2 gas sensitivity of carbon nanotubes obtained by plasma enhanced chemical vapor deposition. *Sens. Actuators B* **2003**, *93*, 333–337.
136. Lin, Y.-H.; Chen, S.-H.; Chuang, Y.-C.; Lu, Y.-C.; Shen, T.Y.; Chang, C.A.; Lin, C.-S. Disposable amperometric immunosensing strips fabricated by Au nanoparticles-modified screen-printed carbon electrodes for the detection of foodborne pathogen *Escherichia coli* O157:H7. *Biosens. Bioelectron.* **2008**, *23*, 1832–1837.
137. Mata, D.; Bejarano, D.; Botero, M.L.; Lozano, P.; Constantí, M.; Katakis, I. Screen-printed integrated microsystem for the electrochemical detection of pathogens. *Electrochim. Acta* **2010**, *55*, 4261–4266.
138. Kostaki, V.T.; Florou, A.B.; Prodromidis, M.I. Electrochemically induced chemical sensor properties in graphite screen-printed electrodes: The case of a chemical sensor for uranium. *Electrochim. Acta* **2011**, *56*, 8857–8860.
139. Betelu, S.; Vautrin-Ul, C.; Ly, J.; Chaussé, A. Screen-printed electrografted electrode for trace uranium analysis. *Talanta* **2009**, *80*, 372–376.