REVIEW

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A review of biosensor for environmental monitoring: principle, application, and corresponding achievement of sustainable development goals

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ABSTRACT

Human health/socioeconomic development is closely correlated to environmental pollution, highlighting the need to monitor contaminants in the real environment with reliable devices such as biosensors. Recently, variety of biosensors gained high attention and employed as *in-situ* application, in real-time, and cost-effective analytical tools for healthy environment. For continuous environmental monitoring, it is necessary for portable, cost-effective, quick, and flexible biosensing devices. These benefits of the biosensor strategy are related to the Sustainable Development Goals (SDGs) established by the United Nations (UN), especially with reference to clean water and sources of energy. However, the relationship between SDGs and biosensor application for environmental monitoring is not well understood. In addition, some limitations and challenges might hinder the biosensor application on environmental monitoring. Herein, we reviewed the different types of biosensors, principle and applications, and their correlation with SDG 6, 12, 13, 14, and 15 as a reference for related authorities and administrators to consider. In this review, biosensors for different pollutants such as heavy metals and organics were documented. The present study highlights the application of biosensor for achieving SDGs. Current advantages and future research aspects are summarized in this paper.

Abbreviations: ATP: Adenosine triphosphate; BOD: Biological oxygen demand; COD: Chemical oxygen demand; Cu-TCPP: Cu-porphyrin; DNA: Deoxyribonucleic acid; EDCs: Endocrine disrupting chemicals; EPA: U.S. Environmental Protection Agency; Fc-HPNs: Ferrocene (Fc)-based hollow polymeric nanospheres; $Fe_3O_4@3D$ -GO: $Fe_3O_4@$ three-dimensional graphene oxide; GC: Gas chromatography; GCE: Glassy carbon electrode; GFP: Green fluorescent protein; GHGs: Greenhouse gases; HPLC: High performance liquid chromatography; ICP-MS: Inductively coupled plasma mass spectrometry; ITO: Indium tin oxide; LAS: Linear alkylbenzene sulfonate; LIG: Laser-induced graphene; LOD: Limit of detection; ME: Magnetoelastic; MFC: Microbial fuel cell; MIP: Molecular imprinting polymers; MWCNT: Multi-walled carbon nanotube; MXC: Microbial electrochemical cell-based; NA: Nucleic acid; OBP: Odorant binding protein; OPs: Organophosphorus; PAHs: Polycyclic aromatic hydrocarbons; PBBs: Polybrominated biphenyls; PBDEs: Polybrominated diphenyl ethers; PCBs: Polychlorinated biphenyls; PGE: Polycrystalline gold electrode; photoMFC: photosynthetic MFC; POPs: Persistent organic pollutants; rGO: Reduced graphene oxide; RNA: Ribonucleic acid; SDGs: Sustainable Development Goals; SERS: Surface enhancement Raman spectrum; SPGE: Screen-printed gold electrode; SPR: Surface plasmon resonance; SWCNTs: single-walled carbon nanotubes; TCPP: Tetrakis (4-carboxyphenyl) porphyrin; TIRF: Total internal reflection fluorescence; TIRF: Total internal reflection fluorescence; TOL: Toluene-catabolic; TPHs: Total petroleum hydrocarbons; UN: United Nations; VOCs: Volatile organic compounds



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Highlights

- Biosensors are robust in both specific and total responses to environmental pollutants
- Promising prospects for achieving on-site monitoring and real-time environmental data
- Biosensors illustrate a reliable, simple, effective, and fast method for monitoring pollution
- Biosensors consume less energy and leave a smaller carbon footprint
- High performing biosensors make a solid contribution to SDGs

1. Introduction

Detection and monitoring of pollutants in the environment is crucial for assessing the harmful effects of potential toxicants to people, flora, and fauna [1-3]. Chemical analysis using instruments such as high performance liquid chromatography (HPLC), gas chromatography (GC), and inductively coupled plasma-mass spectrometry (ICP-MS) has been traditionally used for monitoring environmental water or soil samples for their sensitivity and accuracy [4]. To further understand the bioavailability of environmental pollutants, biosensors using biological sensor to detect such pollutants have been invented, and they have features of low-cost, energy-saving, and feasibility for real-time in situ monitoring [4,5]. Different types of biosensors were successfully applied for specific or nonspecific detection of heavy metals [6,7] and organic pollutants [1,8]. In groundwater and river water samples, the bioavailable and toxic fractions of metals and organic pollutants such as Cd and toluene could be successfully evaluated using biosensors [9,10]. Moreover, bioavailability of the pollutants such as Hg and phenanthrene was revealed by the use of biosensors [11,12], suggesting the useful application of biosensors on bioavailability assessment of pollutants in the environmental samples.

Biosensor is an analytical strategy for pollutants based on biotechnology, and consists of elements for signal transducer producing detectable or quantifiable signals when sensing pollutants [6,13]. Types of biosensor include cell-free-based and whole-cell-based, nonspecific and specific ones according to the biological elements and sensing ability [14]. The

bioavailability and toxic effects can be established by whole-cell-based biosensor compared with a cell-free one [14,15]. For example, microorganisms possess various responsive mechanisms to cope with environmental stress, organic pollutants, and heavy metals, which are associated with different regulatory genes and proteins. The Cd detecting protein cadC regulator could be used in biosensor for Cd detection [16]. In addition, the benzene metabolizing regulatory protein would be useful in recognizing benzene-like organic pollutants [17]. Those regulatory systems interacting with environmental pollutants are the key factors for distinguishing specific pollutants from others [16,17]. However, pollutants with similar structure might hamper the selectivity of biosensors, such as the benzene regulatory protein targeting toluene, ethylbenzene, toluene, and xylene [17]. Therefore, the choice of regulatory systems or recognizing elements would greatly affect the selectivity of specific biosensors. Currently, biosensors targeting heavy metals, pesticides, or pharmaceuticals show the advantages of biosensor such as portability, ease of use, and saving time [18,19]. Recent studies have highlighted the advances made in sensitivity, stability, selectivity, and their contribution to environmental monitoring [1,8].

It has been shown that in clinical laboratories, the consumption of electricity, water, and gas as well as the waste produced are the key factors generating large carbon footprints [20]. Energy use was also the major contributor to greenhouse gases (GHGs) emissions [21,22]. It is suggested that potential carbon reduction could be achieved by partially using biosensors in pollutants analysis due to their low energy requirements. One review has suggested that the contribution of using biosensor for human health on SDGs [23]. However, discussions about the application of biosensor on environmental monitoring and its relationship between sustainable development goals (SDGs) are scarce. Biosensors technology is anticipated to be powerful tool for monitoring the environmental pollutants for its advantages on sustainable development. In this article, different types of biosensors, principles of biosensor construction as well as the application and performance are reviewed. The relationship between biosensors and SDGs is also noted. This review will also help to achieve the SDGs since biosensors play a significant role to achieve the SDG 6, 12, 13, 14, and 15.



Figure 1. General mechanisms of biosensor for pollutants detection. The basic construction of biosensor contains sensing elements and signaling elements. The biological-sensing elements including DNA (aptamer), protein (enzyme), antibody, and whole-cell (bacteria) are able to recognize the environmental pollutants such as heavy metals and organic pollutants. The signaling elements would be triggered by sensing elements and produce the different signals such as fluorescence, luminescence, color, pH change, or electricity that could be measured or detected by the operators.



Figure 2. The types of biosensors and the applications on SDGs achievements. The different environmental pollutants including heavy metals and organic pollutants could trigger the sensing, transducing, and signaling of biosensors. The different types of biosensors include cell-free biosensor, nonspecific whole-cell biosensor, and specific whole-cell biosensor, which is categorized mainly by their sensing elements and selectivity. Those biosensors detecting the environmental pollutants could help achieving SDG 6, 12, 13, 14, 15.

2. Types of biosensors

2.1 Cell-free biosensors

The general mechanism of biosensors for environmental monitoring is displayed in Figure 1. The different types of biosensors and the application on SDGs achievement were shown in Figure 2. The sensing elements of cell-free biosensors include deoxyribonucleic acid (DNA), protein, and aptamer. DNA-based biosensors could be used for monitoring heavy metals such as arsenic and lead made possible by changes in the specific DNA structure and oxidative damage [14,24]. Sulfur on the protein structure reacts easily with toxicants like arsenic and the resulting inhibition of enzyme activities can help detect toxic metals like arsenic [25,26]. Other enzyme such as acetylcholinesterase could also be used in inhibition-based biosensor, which is contributed from the binding of organic P pesticides with acetylcholinesterase inhibiting the further signal producing from the reaction of acetylcholinesterase [27].

However, the response of DNA and protein would involve other toxic substances and might lack selectivity to specific pollutants. To enhance the selectivity of target chemicals, aptamer, an artificial single-strand DNA or ribonucleic acid (RNA), was developed for specific pollutants detection based on selective bonding [28]. Currently, aptamer-based biosensors coupled with nanomaterials have been employed in several studies for detecting heavy metals and organic pollutants such as pesticides [29].

2.2 Nonspecific whole-cell biosensors

Nonspecific whole-cell biosensors are constructed mainly by stress responsive genetic regulation, such as heat shock and DNA damage responses [6]. Information about bioavailability and toxicity can be revealed by nonspecific cell-based biosensors. The damage done to protein will trigger heat shock stressrelated genes, which could be used as a sensing element of nonspecific cell-based biosensors for pollutants that damage protein [30-32]. The expression of SOS response genes enhanced by DNA damage could be incorporated in nonspecific sensing elements [33-36]. Indigenous bacteria and their activity as far as environmental stress is concerned could also serve as a nonspecific whole-cell biosensor. For example, sulfur oxidizing and iron oxidizing abilities in certain types of bacteria would be inhibited by toxic pollutants and could be used for acid samples [37-39]. As well, indigenous bacterium isolated from soil environments was successfully engineered with green fluorescent protein (GFP) signaling gene, which could help assess bioavailable heavy metals in the soils by the GFP signals [40]. The nonspecific biosensor based on the stress response provides an early warning of hazard presence in samples that could be harmful for microorganisms despite the lack of selectivity.

2.3 Specific whole-cell biosensors

Specific whole-cell biosensors are mostly based on the metabolism or detoxification genes in microorganisms [8,41]. For organic pollutants, toluene and xylene can be metabolized by toluene-catabolic (TOL) plasmid in bacteria harboring those regulatory genes [42]. The *xylR* and *xylS* genes on the TOL plasmid can serve in biosensors specifically designed for benzenerelated compounds [43-45]. Other operons such as nah, alkBAC, and DntR were applied in a specific cell-based biosensor for naphthalene, linear alkanes, and pesticides [46-49]. For heavy metals, the sensing elements of specific cell-based biosensor belong to resistance genes instead of metabolized genes, which are located in the plasmid in microorganisms that could survive in toxic metal-rich environments [50]. Expression of the resistance genes including redox transformation, active transport, and intracellular/extracellular precipitation is regulated by intracellular metal level [51]. Both toxic metals and excessive essential metals could trigger the response of resistance or detoxification genes [52,53]. The arsR or arsD in ars operon for arsenic [14], the cadC/A in cad operon for cadmium [54], the merR in mer operon for mercury [6], and the chrB in chr operon for chromium were used in specific cell-based biosensors [55]. Cell-based biosensors for cadmium and mercury were able to detect other metal ions, such as copper, zinc, or lead, which reduces their selectivity [6,54]. zntA and copA promoter/operator evaluated the bioavailability of Zn and Cu [56]. zntA accounts for the removal of intracellular Zn and responds to Cd and Pb [57,58]. copA is responsible for the concentration of Cu and Ag in cells [59]. Regulatory proteins of *zntA* and *copA* belong to the MerR homologue, thereby making them responsible for Hg [60,61]. The cnrYXH gene was used to detect the bioavailability of Co and Ni in soil [62]. In addition, the Pb(II)-binding regulator PbrR in MerR family has the ability to recognize lead after being engineered in wholecell biosensor [63]. The characteristics of biosensors of different types are summarized in Table 1. In addition, the advantages of biosensors on SDGs achievement are displayed in Figure 3.

3. Principle of whole-cell biosensor construction

3.1 Toxicity-based nonspecific biosensor

The mechanisms of nonspecific biosensors based on toxicity include Microtox and adenosine triphosphate (ATP)-bioluminescence system, bacterial

Table 1. Characteristics of various type biosensors.

		Response		
Biosensors	Selectivity	time	Sensitivity	Bioavailability
Cell-free biosensor	High	Fast	High	No
Nonspecific whole- cell biosensor	Low	Medium	Medium	Yes
Specific whole-cell biosensor	High	Medium	Medium	Yes

functions such as iron and sulfur oxidizing, and DNA damage or heat shock protein-related genes. The Microtox test has been devised using the bacterium Vibrio fischeri whose bioluminescence would diminish due to the toxicity and disrupted metabolism [64,65]. The mechanisms of bioluminescence-based inhibitory biosensors might be associated with the ATP reduction resulted from stress responsive mechanisms or cell death resulted from excessive environmental stress, which further decreased the bioluminescence by lower ATP and cell viability [17,66]. It is worth noting that the genes of bioluminescence can be genetically engineered in other labcultivable bacteria such as E. coli. Similarly, the constitutively expressed bioluminescence of these toxicity-based nonspecific biosensors would diminish upon the exposure of toxicants, thereby making them capable of quantifying the general toxicity equivalent of multiple pollutants [66]. As well, the ATP level reflected by bioluminescence could serve as an indicator of toxicity and metabolic disruption.

The modified Saccharomyces cerevisiae with substrate luciferin could serve as a reporter system based on ATP-bioluminescence for detecting nonspecific toxicity [67]. The metabolic cost might increase for stress response when the bacterial functions are inhibited by environmental pollutants, thereby reducing the amount of products of iron and sulfur oxidizing [37-39]. For example, the pH decrease and conductivity increase resulted from sulfur oxidation was inhibited by the toxicants [37], which might be due to the bacterial death or disrupted microbial functions. Based on the inhibition level, toxicity could be quantified by using iron and sulfur oxidizing bacteria [37-39]. The recA and its promoter responsible for DNA repair and maintenance were widely used in DNA damagesensing biosensors for their response to genocompounds toxic and pharmaceuticals [34,68,69]. Recent studies have applied both recA-based and metabolic inhibition-based bioluminescence bacterial biosensor for pollutants such as toxic metals in urban rivers and seawater [70,71]. Similarly, the stress-inducible heat shock genes like hsp70 and its promoter were used in biosensors for detecting nonspecific toxicity since the heat shock protein systems will be triggered not only by heat shock but also other environmental stressors, and then induce the downstream fused signals [72,73].



Figure 3. The advantages of biosensors on SDGs achievements. Several advantages of biosensors include low-cost, ease of use, saving energy, smaller carbon footprints, nonuse of hazardous materials, and minimal sample pretreatment compared with traditional physicochemical methods. These advantages are helpful to achieving SDGs such as SDG 6, 12, 13, 14, and 15, which relate to clean water, responsible consumption and production, climate action, and marine and terrestrial life.

The metabolic inhibition-based bioluminescence bacterial biosensors have been developed and utilized in several pollutants detection [74]. Different signaling elements producing bioluminescence such as luciferase were used in the construction of bacterial biosensor, or naturally presented in bacteria that could be directly used in sensing [74]. The luciferase genes from bacteria and firefly have been suggested to be highly sensitive and broadly used as a reporter gene. The *luxCDABE* from bacterial operon is conservative in many bacteria species [75]. The luxAB is enough for producing the luminescence, while using the whole *luxCDABE* operon could reduce the amount of additional substrates that is needed for luminescence reaction [15]. The protein from *luxAB* is responsible for catalyzing long-chain aldehydes shown in the reaction below:

 $FMNH_2 + R - CHO + O_2 \rightarrow FMN + H_2O + RCOOH + hv (490 nm)$

The *lucFF* gene from the firefly requires luciferin, O_2 , and ATP for bioluminescence [76]:

Firefly luciferin + O_2 + ATP \rightarrow oxyluciferin + AMP + P_i + hv (550–575 nm)

The bioluminescence would diminish in the presence of environmental pollutants due to the inhibited bioluminescence reaction. The ATPdependent bioluminescence could be used in developing the metabolic inhibition-based biosensors for it can reflect the ATP decrease caused by environmental toxicants [67,74]. On the other hand, the energy-consuming *lux* operon reaction in Vibro spp. could also reflect the metabolic state of the microorganisms [66]. Therefore, the bioluminescence biosensors with *lux* operon such as Microtox (Vibrio fischeri) were applied in assessing the toxicity of environmental pollutants in river and sediment samples [71,77,78]. In addition to Vibro spp., the recombinant Escherichia coli harboring the *lux* operon could successfully detect the environmental pollutants, suggesting the feasibility of *lux* operon engineered in other bacterial species that could be easily cultured [8,79].

3.3 Microbial fuel cell-based biosensors

Microbial fuel cell (MFC)-based biosensors with single- or dual-chamber design were developed based on the oxidation of organic compounds and the corresponding electrical currents [80,81]. The dual chambers of MFC-based biosensors consist of an anaerobic anodic chamber where the oxidation happens and aerobic cathodic chamber, transferring electrons through circuits and protons through the ion-exchange membrane from anode to cathode [82]. MFC-based biosensors can detect the water quality and wastewater parameters such as biological oxygen demand (BOD) and toxins [81]. Compared with nonspecific whole-cell biosensors, MFC-based biosensors save more energy and are portable; they produce the electrical current and do not need signal transducers [82]. It has been reported that heavy metals such as copper and arsenic in wastewater were detected by MFCbased biosensor based on the toxicity associated inhibition [83]. The bacteria strains for organics degradation could be mixed culture from activated sludge, and the toxicants in wastewater could be sensed by the inhibited degradation [82]. In addition to pollutant-degrading bacteria, the electroactive bacteria such as Geobacter and Shewanella were shown to facilitate the current production in MFCbased biosensor [84]. Other bacterial strains in MFC-based biosensors for sensing specific pollutants were also reported in recent studies. For example, the linear alkylbenzene sulfonate could be simultaneously quantified and degraded dominantly by Pseudomonas species, generating electrical power using a MFC-based biosensor [85]. To increase the selectivity for specific detection, Cr(VI)-reducing bacterium Exiguobacterium aestuarii YC211 was cultured in a MFC-based biosensor system [86]. It can detect in situ Cr(VI) in industrial wastewater via the electrical currents generated from the reduction of Cr(VI) to Cr(III) [86], which suggests the possibility of specific detection and indigenous bacterium application in MFCbased biosensors. In addition, the genetically engineered E. coli was developed in MFC-based biosensors for detecting Zn(II) and Cu(II), which is accomplished by the recombining metal-sensing promoter, and the synthesis genes of riboflavin and porin [87,88]. The riboflavin and porin are responsible for facilitating electron transfer and improving cell membrane permeability, respectively [87,88].

3.4 Genetic modification using detoxification *and metabolism genes*

To detect specific pollutants, genetic methods using detoxification and metabolism genes to differentiate environmental pollutants have been developed. For organic compounds, the metabolic and utilization mechanisms and their regulatory genes were mainly used [8]. Recent research has combined genes and hosts from different bacterial species and used AtzR regulatory protein to detect cyanuric acid [89]. Biodegradable organic pollutants in the whole-cell biosensors trigger metabolic mechanisms and the downstream utilizing proteins or signals such as bioluminescence [8,90]. Subsequently, it is difficult to detect biodegradable organic pollutants since the metabolism mechanisms are unclear in the microorganisms. However, recent research has developed a whole-cell biosensor to detect persistent polybrominated diphenyl ethers (PBDEs) using genome-wide screening. It can find regulatory networks in the microorganisms [91], shedding light on the application of biosensors on organic pollutants that can only be degraded with difficulty.

For heavy metals, the resistance and detoxification mechanisms were applied in the sensing elements of specific whole-cell biosensors [92]. The regulatory systems consisted of regulatory protein that could bind metals and promoter regions that trigger downstream detoxification or signaling responses [92,93]. For example, arsR and the promoter region from Geobacter sulfurreducens could be genetically combined with a fluorescence reporter, usable for arsenic screening with the detection limit down to the ppb level [41]. Basically, the detoxification and metabolism operons were activated when the toxicants permeated in the cells, but it has been suggested that the two-component regulatory systems could be incorporated in bacterial biosensors for detecting extracellular pollutants [94]. The regulatory membrane protein has the ability to bind pollutants and then trigger downstream phosphorylation of the other regulator proteins that activate the signal transduction [94].

4. Application of biosensor to environmental monitoring

4.1 Heavy metals monitoring

Heavy metals pollution is usually discharged as a result of anthropogenic and industrial activities (e.g., refineries, metal works, mining, cement factories, smelting plants, etc.), and their pollution endangers human health and the environment [95-97]. Elements such as Fe, Cu, Mn, and Zn, then highly toxic Pb, Cr, Cd, As, Hg, etc, are very resistant to biodegradation. Heavy metals are transported into the environment, especially water sources and easily absorbed by living organisms. Due to their pronounced toxicity, these heavy metals are important for monitoring the environment (water, wastewater, air, solid waste, and organisms) [95,97,98]. Monitoring water contamination is essential for environmental conservation and the prevention of illnesses. Due to the accumulation in the environment (e.g., water, soil, sediment) and wildlife such as plants, animals over a long period of time, heavy metals pose a serious danger. Several methods have been devised to detect their concentrations and presence in the environmental matrix. Especially, biosensors can detect the levels of heavy metals and determine how much pollution they cause. Biosensors can easily detect the presence of heavy metals in order to regulate and manage water safety and quality.

The DNA probe is used as the element recognition to detect heavy metal elements/ions and is based on the principles as follows. The first is the formation of a stable DNA-duplex due to the selective binding of specific DNA bases with heavy metal ions; for the second, heavy metal ions aid in the breaking DNAzymes (deoxyribozymes), and finally, the guanine-rich probe switches to a stable G-quadruplex structure [99,100]. Similar to the design for heavy metals detection in environmental monitoring, microbial whole-cell biosensors detect concentrations based on the genetic elements that respond to chemical species. Their effectiveness is determined by the regulatory protein types connected with these promoters, as well as the reporter genes for transcriptional pollutants control [101,102]. A reporter gene in living cells (i.e., microorganisms) is employed as a sensor to convert its biological response into detectable physicochemical signals.

The whole-cell biosensors illustrate a prospective technique for the detection/monitoring of heavy metals and are critical to selectivity and sensitivity [70,100,103]. Using biosensors for continuous and/or individual detection and measurements normally depends on the biologically active element types. Also, biosensors (i.e., organisms) can be integrated into pollutants exposure and used as a method for predicting chemical pollution in the environment. The literature review demonstrated that Shen et al. [104] designed a toxicity MFC system for detecting Cu concentrations of 5–7 mg L^{-1} in a quick speedy response, and Wu et al. [105] also found at a level of 12 mg L^{-1} in their study. Recently, by using the decrease in the cell voltage, the MFC biosensor could monitor arsenic $(0.5-5.0 \text{ mg } \text{L}^{-1})$ and copper $(1.0-10 \text{ mg L}^{-1})$ with added Cu/As to the anolyte solution [83]. Wu et al. [86] inoculated a facultative anaerobe bacterium into an MFC to investigate an in-situ Cr⁶⁺ detection sensor. In their research, Wu et al. [105] developed a new sediment MFC-based toxic sensor for online and *in-situ* Cu²⁺ monitoring by using bacteria species such as Clostridium and Geobacter. A similar study was conducted using Pseudomonas and Geobacter for in-situ Cr⁶⁺ detection in real-time from industrial wastewater [106]. Researchers have recently become interested in electro-active biofilms because of their potential applications in environmental monitoring as amperometric biosensors, i.e., in-situ toxicant detection. For the assessment of heavy metal contamination in tap water, an O₂ reducing microbial cathode-based MFC biosensor was designed, with detection limits in the 1.0 to 10 mg L^{-1} range for ions, including Cr^{6+} , Hg²⁺ and Pb^{2+} [107]. As a result, biosensors can accurately reflect the harmful effects of numerous contaminants in the environment. This understanding provides strong technical solutions for online, direct pollutant detection (i.e., heavy metals) and the establishment of an early warning sensor system. Table 2 summarizes the materials and performance of biosensors for heavy metals monitoring.

Several biosensors based on electrochemical, colorimetric, and fluorescence measurements were successfully devised for heavy metal detection [114–116]. It has been suggested that the nanomaterials such as metal oxide and nanostructured carbon could be effectively used in biosensors development [5,117-119]. In addition, several nanomaterials with heterostructure such as a-Fe₂ O₃-g-C₃N₄, V₂O₅ g-C₃N₄, and CuS-WuO₃ were developed, indicating the potential use in electrochemical biosensors [120-122]. For electrochemical sensors, nano-scale structures are attractive materials [123], and their huge specific surface can enhance enzyme, signaling molecule, and catalyst immobilization. Applying electrochemical DNA biosensors for heavy metals detection includes implementing core-shell nanoparticles, nicking enzyme-assisted amplification, and nanocomposites modification [124,125]. Apart from these, applications of nanomaterials based on electrochemical nucleic acid (NA) biosensors for quantitative and qualitative analysis/monitoring of environmental pollutants, e.g., heavy metals have been discussed [126,127]. Table 2 demonstrates how biosensors were classified depending on the recognition component that was used for detecting heavy metals. For example, the bioavailable copper ions in synthetic samples were monitored by biosensor used as an optical transducer and immobilized engineered bacterium Alcaligenes eutrophus (AE1239), with the limit of detection (LOD) being 1 µM [128]. DNA-based electrodes biosensor is an excellent candidate for Hg detection, with a LOD of 3 fM [127]. Another DNA biosensor using electrode modification with Fe₃O₄ @3D-GO reached an excellent LOD for Ag⁺ ion detection, which was equal to 2.0 pM and in the wide range of 0.01–100 nM [129]. A self-cleaning electrochemical biosensor based on DNA nanomotors, novel super G-quadruplex (G4), and 2D Cu-porphyrin (Cu-TCPP) metal-organic nanofilms was designed for the cyclic detection of Pb^2 ⁺ ions, which the LOD was equal to 1.7 nM, and the linear range of 5 nM-5 μ M [123]. The synergistic effects of G4-hemin DNAzymes and Cu-

	Refs.	[83] [101]		[106]	[108]	[109]	[130]	[66]	100 5		[40]	[127]	[12]	[70]		[110]		[105]	[111]	[112]	101	[/8]	[103]	[123]		[113]		[138]	[102]		i-walled
	Remarks	MFC as a heavy metal biosensor A simple, cheap and sensitive		Real-time in-situ detection	Environmentally friendly method	High-sensitive	Based on 8–17 DNAzyme	Providing a platform for heavy	metal detection	Lost-effective	A green fluorescent protein	A time and cost-effective	Respond at low levels	Rapid detection, sensitive	biosensor	Cost-effective		A cost-effective	A simple, facile, and cost-effective	Good sensitivity, selectivity, simulicity, and ranidity	אוווידיר, מוש ומאושיניע אווידיריידי ידי-י	High sensitivity	Highly selective and sensitive	recognition Hich sensitive and selective		High sensitivity		A satisfactory recovery rate, high sensitive	On-site detection, saving time and	costs	eoxyribonucleic acid, MWCNT: Mult
Response	times	5 min N/A		18.3 min	N/A	5 min	N/A	N/A		N/A	4–7 h	N/A	30 min	A quick		Short time		N/A	N/A	N/A	V 1 V	N/A	N/A	N/A		10-	20 min	N/A	45 min		astic, DNA: De
	Detection limits	0.5–10 mg L ^{–1} 2 mg L ^{–1}	ו ע ו	0.2–0.7 mg L ^{–1}	200 mg L ⁻¹	0.23–0.33 µM	10 µM	0.0001-300 nM	- 	л Бш 07-с	1.42×10^{-4} - 3.16 × 10 ⁻⁴ mg L ⁻¹	$1 \times 10^{-6} \text{nM}$	Low	0.02–1.0 mg L ⁻¹)	0.0125-200 mg L ⁻¹	I	0.5–12.5 mg L ^{–1}	60.7 nM	$8.4 imes10^{-4}$ nM		0.9/–14.54 mg L	5×10^{-5} –100 nM	Mn 7.1		1.25 mM		19.93 nM	5 uM	-	oxide, ME: Magnetoel
	Environmentalmatrix	Municipal wastewater Water containing heavy	metal ions	Industrial wastewater	Artificial water	Heavy metal ions solutions	Lead ions solutions	Environmental matrices		ingustrial wastewater	Soilenvironment	Water samples	River water	Seawater		Water, wastewater,	sewage sludge	Artificial solution	Water, serum samples	Aqueous solution		Aquatic environments	Tap, river, lake,	wastewater, etc. Artificial solution		Artificial solution		Artificial solution	Artificially	contaminated soil	electrode, ITO: Indium tin o
es/Materials	Cathode	Carbon fiber Granhite felts		Graphite felt	Graphitepaper			nagnetic beads, etc.		urapnite sneet	i film of sol-gel)	/CNT	P1 Tox2			Graphite, carbon	cloth, platinum	Carbon cloth	-Rad)	, SWCNT substrate and		scentbacterium <i>Vibrio sp.</i>	DNAzymes	DNAzvmes		invertase, mutarotase,		resis, oligonucleotides	sor system		SPGE: Screen-printed gold
Electrode	Anode	Carbon fiber Graphite felts		Carbon cloth	Graphitefelt	ME ribbon	A silicon membrane	Gold, SPGE, PGE, ITO, n		urapnite sneet	Sol-gel matrices (a thin	Gold nanoparticles, MM	Acinetobacter baylyi AD	A. baylyi Tox2biosensor	×	Graphite, carbon cloth,	stainless steel	Carbon brush	Micro-spin column (Bio	Magnetic nanoparticles		A novel marine lumine: <i>MM1</i>	Nanomaterial-assisted [Cu-TCPP nanofilm and		Three-enzyme system (glucose oxidase)	Agarose gel electropho DNA	A das reporting biosen	-	stalline gold electrode, S
	Configuration of biosensors	Double chamber MFC A dual-chamber self-powered	MFC	Sediment MFC	Doublechamber MFC	Magnetoelastic (ME) biosensor	Micromechanicalbiosensor	DNA-based electrochemical	biosensors	keusaple single-cnampered cylindrical MFC	Whole-cell bacterial biosensor	DNA-based electrodes	Bacterial molecular biosensors	A bacterial whole-cell	biosensor	MFC-based biosensors		A single-chamber MFC	Light-up biosensor	SERS-based biosensor		biosensor cells	DNAzyme-based biosensors	A self-cleaning	electrochemical biosensor	Novel conductometric	biosensor	A novel signal amplification biosensor	Whole-cell biosensors		licrobial fuel cell, PGE: Polycrys
	Heavy metals	Cu, As, Cr, Hg Ph. etc	- 21 - 22 -	Ľ	Hg	Pb, Cd, Cu	Pb	Hg, Ag, Cu, Pb		ru, rr, zn, NI	Cd, Cu,Zn	Ha	Se, Sn, Cr	Ha, Zn, Cu, Cd		Cu, Hg, Zn, Fe,	Cr, Cd	Cu	Pb	Hg		Pb, NI, Cd, etc.	Hg	Ph	2	Hg, Ag		Cd	Ha	'n	Remarks: MFC: N

Table 2. Biosensors for heavy metal monitoring with detection limits and response time.

TCPP, which has efficient and catalytic H_2O_2 reduction, resulting in better performance of electrochemical biosensors [123].

DNA biosensors focusing on the recent design of selective and sensitive detection of heavy metal elements/ions by electrochemical transduction are depicted in Table 2. DNA biosensors used electrochemical transduction to provide a suitable platform for early monitoring/detection of heavy metal ions. Many effective DNA biosensors based on various electrochemical redox indicators have been applied for Ag^+ , Hg^{2+} , Pb^{2+} ions detection bacterial А [99]. whole-cell biosensor (Acinetobacter baylyi Tox2) was designed to detect heavy metal cytotoxicity in the polluted seawater sources [70]. Consequently, A. baylyi Tox2 exhibits excellent application as a sensitive and quick biosensor for investigating cytotoxicity in the marine environment. Many detection methods have been assessed using the 8-17 DNAzyme specific for lead ions [115,130,131]. Shen et al. [132] used DNA-Au bio-barcode based on a signal amplification assay for Pb²⁺ ion detection. Thus, biosensors have attracted much interest for biomonitoring purposes, especially of heavy metal ions.

Results found that Hg²⁺ concentrations analyzed by biosensor could be confirmed by standard methods (e.g., Inductively coupled plasma mass spectrometry, ICP-MS), showing that the biosensor was reliable and relevant for Hg^{2+} ion detection in real samples [133]. For instance, autocatalytic DNA circuit based on exonuclease III and G-quadruplex DNAzyme for mercury ions detection achieved high selectivity and sensitivity. For an LOD, it was 10 fM and the linear range was from 10 fM to 100 nM [134]. Shi et al. [133] used a biosensing system for detecting Hg^{2+} in tap and lake water samples with a 3D graphene/ gold electrode and a reporter probe attached to Au nanoparticles. An excellent widely linear range from 0.1 fM to 0.1 µM was reached. In contrast, Hg²⁺, Pb²⁺ and Cd²⁺ levels are strictly monitored by the European Union [103] due to these substances' highly toxic, bioaccumulative properties, and their impact on human health and the environment [133]. According to the U.S. Environmental Protection Agency (EPA) regulations, the maximum levels of heavy metals such as Hg²⁺, Pb²⁺, and Cd²⁺ in drinking water are 10 nM, 72 nM and 45 nM, respectively, so these described biosensors could effectively monitor their concentrations [103,116]. The LOD of these heavy metals are detected by biosensors lower than the toxicity safety levelfor example, in drinking water to monitor the quality. Shown here is the critical role of advanced biosensors for heavy metals detection/monitoring to meet requirements, such as monitoring Hg^{2+} concentration in drinking water (i.e., level of 10 nM), following the U.S. EPA standard [134].

Several studies have used eukaryotic microorganisms to investigate whole-cell biosensors for environmental pollution monitoring of heavy metals in aquatic habitats or soils [135]. Yeasts, ciliated protozoa, and microalgae are generally selected as the three main eukaryotic taxonomic groups. For example, a new conductometric biosensor based on alkaline phosphatase activity was developed utilizing immobilized whole-cell microalgae to detect cadmium ions in aquatic environments and habitats [136]. A novel whole-cell microbial biosensor was invented for rapid onsite detection related to Hg contaminated soil with gas signals [102]. This technique could detect bioavailable mercury levels within 45 min and a range from 5 to 500 µM, effectively showing how much pollution there was in the soil. Guo et al. [137] illustrated that a fluorescent wholecell biosensor could detect mercury (Hg²⁺) contamination in cosmetics with the detection range from 50 nM to 10 µM with incubating time for two hours. It means that biosensors can be used for many purposes and in many industries.

Biosensors could detect and provide information quickly about the toxic pollutants and contamination zone, which is necessary for good environmental management and monitoring. Another advantage of biosensors over traditional analytic techniques is related to their mobility, making possible the measurement of in-situ pollutant levels without added chemical agents and sample preparation. Biosensors can detect and conduct single measurements or continuous realtime monitoring during analysis processes. DNA biosensors based on electrochemical transduction, constitute an sensitive technique and affordable method for detection and monitoring of heavy metal elements [99,138]. Biological sensors are promising in heavy metal ions detection, especially DNA. DNA and DNAzymes are biodegradable compositions, exhibit high selectivity, the advantage of a portable analytical device and in-vitro technique, as well. Based on the highly selectivity of DNAzymes (as a biological recognition element), electrochemical biosensors can determine heavy metal compounds. DNA is easy to synthesize, low-cost, stable at room or higher temperatures, and is an ideal material for biosensors [134]. It shows novel advantages with excellent potential in toxicity determination, i.e., sensitive, costeffective devices, rapid response to the toxin and short life cycle [71,139,140].

The excellent sensitivity of DNA-based or DNAzyme-based biosensors on heavy metals was indicated by its low LOD (below nM), especially those integrating nanomaterials (Table 2). However, the cell-free biosensor could not provide the information of bioavailable metals in the environment. In addition, the high cost and requirements of advanced technology of nanomaterials might impede the production and application of these cell-free biosensors in developing countries. On the other hand, although the whole-cell or MFCbased biosensors could reveal the bioavailable heavy metals in the samples with lower cost, the sensitivity of them are mostly lower than DNA-based biosensors, limiting their commercial feasibility.

4.2 Organic pollutants monitoring

Anthropogenic activities ensure that the natural environment is contaminated by organic pollutants. A wide range of contaminants originates from different industrial, household and agricultural activities. Agricultural waste organic herbicides and pesticides contain toxic compounds mostly found in wastewater, and the compounds are widely used to remove weeds, pests and unwanted vegetation. Industrial wastewater contains organic matter carrying various hydrocarbons, chlorine compounds, aromatic substances and surfactants. Persistent organic pollutants (POPs) including polychlorinated biphenyls (PCBs), polybrominated diphenyl ethers (PBDEs), polybrominated biphenyls (PBBs), phthalate esters (PAEs), and other hazardous pollutants exist in wastewater [141–143]. industrial Household

wastes also contribute to organic pollutants in wastewater such as phenyl ether which is found in everyday household products such as soaps, deodorants, plastics and cosmetics [144]. Monitoring the particular organic matter in wastewater is an important aspect of human health and the environment, particularly wastewater treatment and water reclamation processes. The application of biosensor for detecting organic contaminants in wastewater achieves the fastest and most accurate results compared to other traditional methods [145]. Biosensor application can be used for environmental organic pollutants including endocrine disrupting chemicals (EDCs), petroleum PCBs, POPs, total hydrocarbons polycyclic aromatic hydrocarbons (TPHs), (PAHs) and volatile organic compounds (VOCs) (see Table 3).

Advances in the field of biotechnology have enabled biological materials to function as receptors, so these bioreceptors can analyze VOCs with high selectivity and sensitivity [161]. Some applications can detect VOCs based on protein and peptides biosensors, such as sensitive olfactory biosensors having a remarkable performance for high selectivity and sensitivity at very low concentrations by using odorant-binding protein (OBP) operated as a sensing component [162]. In addition, the nonspecific biosensor based on microbial electrochemical cell-based (MXC) structure can be successfully used to monitor PCBs and PAHs such as toluene [163]. For specific targeting of benzene, phenol, toluene, and other related pollutants, engineered whole-cell biosensors based on the metabolic genes as sensing elements were developed [164,165]. Other substances including pesticides and EDCs could be detected in river water and wastewater by biosensors especially nanomaterialsbased ones, suggesting their feasibility in real environmental samples [166].

Currently, the successful application on VOCs and PAHs such as alkanes, toluene, naphthalene in wastewater, seawater, and soil using whole-cell luminescence biosensor has been reported [8]. However, limited information about the key sensing element based on the metabolic mechanisms of microorganisms constrain the development of biosensors for specific organic pollutants such as emerging contaminants [91]. Recent research has

Connguration of bio Bisphenol A Classification Sensing element Si Bisphenol A EDC Enzyme biosensor Tyr- mm Bisphenol A EDC Enzyme biosensor Tyr- mm Bisphenol A EDC EDC Enzyme biosensor Tyr- mm Bisphenol A EDC Sol-gel films Tyr/ bisphenol Tyr/ bisphenol Bisphenol A EDC Sol-gel films Tyr/ bisphenol Tyr/ bisphenol Bisphenol A EDC Sol-gel films Tyr/ bisphenol Tyr/ bisphenol Tyr/ bisphenol Phenol EDC Nonvjphenol EDC Whole-cell (Acinetobacter Ligh Pricosan EDC Mole-cell (Acinetobacter Ligh Sutificacion Piot Presticide EDC Mole-cell (Acinetobacter Ligh Sutificacion Piot Presticide EDC Mole-cell (Acinetobacter Ligh Piot Piot Presticide Presticide Problement inporting Piot Piot Piot ActazineBisphenol Herbic	Contiguration of biosensorsg elementg elementrosinase)asensorTyr - AuNPs/GraphitesensorTyr - AuNPs/Graphiteoxideoxideoxideoxideoxideoxideoxideoxideoxideoxideoxideoxideoxideoxideoxidefluorimetryrosinase)AmperometryfibodyFluorimetryfluorimetrycosinaseibodyFluorimetryrosinaserosinasefibodyfluorimetryrosinasefluorimetryrosinasei (MIP)resonance (SPR)ribodiesfluorescence (TIRF)igG antibodyFluorimetry	Detection limit 0.01 nM 0.01 nM 1 nM 0.014 µg L ⁻¹ 66 nM 4.4 mg L ⁻¹ 4.06 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg 1.5.2 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg 2.5 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg 0.2 ng L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg 0.2 ng L ⁻¹ 0.017 µg L ⁻¹ 0.016 µg L ⁻¹ 0.016 µg L ⁻¹	Remarks Promising reliable Excellent performanceLong-term stabilityPromising on-site analysis Satisfactory repeatabilityGood agreement High sensitivity Good precision FastGood sensitivity FastGood sensitivity Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	Refs. [146] [147] [148] [149] [149] [150] [151] [151] [152] [153] [153]
Pollutants Classification Sensing element Si Bisphenol A EDC Enzyme (tyrosinase) Amp Bisphenol A EDC Enzyme biosensor Tyr/- Bisphenol A EDC EDC Enzyme biosensor Tyr/- Bisphenol A EDC Labeled antibody Fluo Bisphenol A EDC Sol-gel films Tyr/- Bisphenol A EDC Sol-gel films Tyr/- Bisphenol A EDC Sol-gel films Tyr/- Estradiol Nov/lohenol Nov/lohenol Nov/lohenol Phenol Nov/lohenol EDC Whole-cell (Acinetobacter Ligh Protone EDC Whole-cell (Acinetobacter Ligh Surf. Protonal EDC Whole-cell (Acinetobacter Dight No Phenol EDC Whole-cell (Acinetobacter Dight No Protonal EDC Whole-cell (Acinetobacter Dight No Protonal EDC Molecular imprinting Surf.	g element Signaling element rosinase) Amperometry/MWCNT rosinase) Amperometry/MWCNT ssensor Tyr - AuNPs/Graphite oxide Fluorimetry tibody Fluorimetry no Nafion/graphite rosinase) Amperometry fuorimetry - Nafion/graphite rosinase) Amperometry fibody Fluorimetry fibody Fluorimetry rosinase Light emission/ fibody Fluorimetry fibodis Light emission/ fibodies Total internal reflection fibodies Fluorimetry gG antibody Fluorimetry	Detection limit 0.01 nM 1 nM 0.014 µg L ⁻¹ 66 nM 4.4 mg L ⁻¹ 4.06 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg L ⁻¹ 5.2 mg L ⁻¹ 0.2 ng L ⁻¹ 0.2 ng L ⁻¹ 0.017 µg L ⁻¹ 0.017 µg L ⁻¹	Remarks Promising reliable Excellent performanceLong-term stabilityPromising on-site analysis Satisfactory repeatabilityGood agreement High sensitivity Good precision Fast, sensitivity Good precision Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	Refs. [146] [147] [148] [149] [150] [151] [151] [153] [153]
Bisphenol A EDC Enzyme (tyrosinase) Amp Bisphenol A EDC EDC Enzyme biosensor 7yr- Bisphenol A EDC Labeled antibody Fluo Bisphenol A EDC Sol-gel films -1 Bisphenol A EDC Sol-gel films -1 Extrogen EDC Sol-gel films -1 Estradiol Phenol Nonylphenol Amp Disphenol A EDC Sol-gel films -1 Estradiol Nonylphenol EDC Sol-gel films Nonylphenol EDC Nolecular imprinting -1 Phenol None-cell (Acinetobacter bi Nonylphenol EDC Whole-cell (Acinetobacter bi Phenol EDC Nolecular imprinting Surf. Phenol EDC Nolecular imprinting Surf. Propanil Herbicide Labeled antibody Fluo Acetamiprid Pesticide Labeled antibody Fluo Acetamiprid Pesticide Labeled antibody Fluo Acetamiprid Pesticide Pesticide Acetamipody Fluo Acetamiprid Pesticide Pesticide Polyclonal IgG antibody No <th>rosinase) Amperometry/MWCNT modified Jyr – AuNPs/Graphite oxide Fluorimetry - Nafion/graphite rosinase) Amperometry - Nafion/graphite Amperometry fluorimetry fluorimetry ison/ bioluminescence mprinting Surface plasmon i (MIP) Total internal reflection fluorescence (TIRF) gG antibody Fluorimetry</th> <th>0.01 nM 1 nM 0.014 µg L⁻¹ 66 nM 4.4 mg L⁻¹4.06 mg L⁻¹0.94 mg L⁻¹2.2 mg L⁻¹5.2 mg L⁻¹0.94 mg L⁻¹2.2 mg 2.5 mg L⁻¹0.94 mg L⁻¹2.2 mg 0.2 ng L⁻¹0.017 µg L⁻¹ 0.017 µg L⁻¹</th> <th>Promising reliable Excellent performanceLong-term stabilityPromising on-site analysis Satisfactory repeatabilityGood agreement High sensitivity Good precision Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor</th> <th>[146] [147] [148] [149] [150] [151] [151] [153]</th>	rosinase) Amperometry/MWCNT modified Jyr – AuNPs/Graphite oxide Fluorimetry - Nafion/graphite rosinase) Amperometry - Nafion/graphite Amperometry fluorimetry fluorimetry ison/ bioluminescence mprinting Surface plasmon i (MIP) Total internal reflection fluorescence (TIRF) gG antibody Fluorimetry	0.01 nM 1 nM 0.014 µg L ⁻¹ 66 nM 4.4 mg L ⁻¹ 4.06 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg L ⁻¹ 5.2 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg 2.5 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg 0.2 ng L ⁻¹ 0.017 µg L ⁻¹ 0.017 µg L ⁻¹	Promising reliable Excellent performanceLong-term stabilityPromising on-site analysis Satisfactory repeatabilityGood agreement High sensitivity Good precision Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	[146] [147] [148] [149] [150] [151] [151] [153]
Bisphenol A EDC Enzyme biosensor Tyr/- Bisphenol A EDC Labeled antibody Fluo Bisphenol A EDC Sol-gel films -1 Estrogen EDCs Sol-gel films -1 Estrogen EDCs Sol-gel films -1 Estrogen EDCs EDCs Enzyme (tyrosinase) Amp Estrone EDC Labeled antibody Fluo Nonylphenol Bisphenol A EDC Labeled antibody Fluo Phenol EDC Whole-cell (Acinetobacter Ligh Nonscheide EDC Whole-cell (Acinetobacter Iigh Phenol EDC Molecular imprinting Suff Propanil Herbicide Labeled antibody Fluo Acetamiprid Pesticide Polyclonal IgG antibody Fluo Acetamiprid Pesticide Polyclonal IgG antibody Flor	ssensor Tyr – AuNPs/Graphite etibody Fluorimetry as Tyr/TiO_z-MWCNTs-PDDA - Nafion/graphite rosinase) Amperometry Amperometry fluorimetry fluorimetry bioluminescence mprinting Surface plasmon resonance (SPR) tibodies fluorescence (TIRF) gG antibody Fluorimetry	1 nM 0.014 µg L ⁻¹ 66 nM 4.4 mg L ⁻¹ 4.06 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg L ⁻¹ 5.2 mg L ⁻¹ 0.2 ng L ⁻¹ 0.2 ng L ⁻¹ 0.2 ng L ⁻¹ 0.017 µg L ⁻¹	Excellent performanceLong-term stabilityPromising on-site analysis Satisfactory repeatabilityGood agreement High sensitivity Good precision FastGood sensitivity FastGood sensitivity Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	[147] [148] [149] [150] [151] [151] [153]
Bisphenol A EDC Labeled antibody Fluo Bisphenol A EDC Sol-gel films Tyr/1-1-1 Estrogen EDCs Enzyme (tyrosinase) Amp Estrodiol Nonylphenol Enzyme (tyrosinase) Amp Denol Nonylphenol Enzyme (tyrosinase) Amp Estrodiol Nonylphenol Enzyme (tyrosinase) Amp Phenol Nonylphenol Enzyme (tyrosinase) Amp Pisphenol EDCs Enzyme (tyrosinase) Amp Pisobenol A EDC Nole-cell (Acinetobacter Ligh Nonylphenol EDC Whole-cell (Acinetobacter Ligh Pisobenol A EDC Nole-cell (Acinetobacter Ligh Picolan EDC Nole-cell (Acinetobacter Ligh Picolan EDC Nole-cell (Acinetobacter Ligh AtrazineBisphenol Herbicide Polyclonal IgG antibody Fluo Actamiprid Pesticide Polyclonal IgG antibody Fluo Actamiprid Pesticide Monoclonal antibody Nu Actamiprid Pesticide Monoclonal antibody Nu	tibody Fluorimetry Tyr/TiO ₂ -MWCNTs-PDDA - Nafion/graphite rosinase) Amperometry tibody Fluorimetry (<i>Acinetobacter</i> Light emission/ bioluminescence mprinting Surface plasmon resonance (SPR) tibodies Fluorimetry gG antibody Fluorimetry	0.014 µg L ⁻¹ 66 nM 4.4 mg L ⁻¹ 4.06 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg L ⁻¹ 5.2 mg L ⁻¹ 0.2 ng L ⁻¹ 0.2 ng L ⁻¹ 0.017 µg L ⁻¹ 0.017 µg L ⁻¹	Satisfactory repeatabilityGood agreement High sensitivity Good precision FastGood sensitivity Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	[148] [149] [150] [151] [151] [144] [153]
Bisphenol A EDC Sol-gel films Tyr/l Estrogen EDCs Enzyme (tyrosinase) Amp Estradiol Nonylphenol Amp Enzyme (tyrosinase) Amp Phenol Nonylphenol Nonylphenol Amp Enzyme (tyrosinase) Amp Phenol Nonylphenol EDC Enzyme (tyrosinase) Amp Pisphenol Nonylphenol Nonylphenol Amp Enzyme (tyrosinase) Amp Pisphenol Nonylphenol EDC Labeled antibody Fluo Pisphenol EDC Whole-cell (Acinetobacter Ligh Picolan EDC Nhole-cell (Acinetobacter Ligh Picolan EDC Nhole-cell (Acinetobacter Nin Picolan EDC Nhole-cell (Acinetobacter Ligh AtrazineBisphenol Herbicide Labeled antibodies Tota Actamiprid Herbicide Polyclonal IgG antibody Ferricide Actamiprid Pesticide Monoclonal antibody Surf.	rs Tyr/TiO ₂ -MWCNTs-PDDA - Nafion/graphite rosinase) Amperometry Amperometry (<i>Acinetobacter</i> Light emission/ bioluminescence mprinting Surface plasmon resonance (SPR) tibodies fluorescence (TIRF) igG antibody Fluorimetry	66 nM 4.4 mg L ⁻¹ 4.06 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg L ⁻¹ 5.2 mg L ⁻¹ 0.2 ng L ⁻¹ 0.2 ng L ⁻¹ 2.5 mg L ⁻¹ 0.017 μg L ⁻¹ 0.002 μg L ⁻¹ 0.016 μg L ⁻¹	High sensitivity Good precision FastGood sensitivity Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	[149] [150] [151] [152] [144] [153]
Estrogen EDCs Enzyme (tyrosinase) Amp Estradiol Phenol Nonylphenol A Estrone EDC Labeled antibody Fluo Bisphenol A Estrone EDC Whole-cell (<i>Acinetobacter</i> Ligh sp. DF4) bi Triclosan EDC Molecular imprinting Surfi polymers (MIP) re AtrazineBisphenol Herbicide Labeled antibodies Tota AtrazineBisphenol Herbicide Polyclonal IgG antibody Fluo Propanil Herbicide Polyclonal IgG antibody Fluo Carbaryl Pesticide Monoclonal antibody Surfi	rosinase) Amperometry tibody Fluorimetry (<i>Acinetobacter</i> Light emission/ bioluminescence mprinting Surface plasmon : (MIP) resonance (SPR) ribodies fluorescence (TIRF) igG antibody Fluorimetry	4.4 mg L ⁻¹ 4.06 mg L ⁻¹ 0.94 mg L ⁻¹ 2.2 mg L ⁻¹ 5.2 mg L ⁻¹ 0.2 ng L ⁻¹ 2.5 mg L ⁻¹ 0.017 μg L ⁻¹ 0.002 μg L ⁻¹ 0.016 μg L ⁻¹	FastGood sensitivity Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	[150] [151] [144] [153]
Estrone EDC Labeled antibody Fluo Phenol EDC Whole-cell (<i>Acinetobacter</i> Ligh Triclosan EDC Whole-cell (<i>Acinetobacter</i> bi Triclosan EDC Molecular imprinting Surft polymers (MIP) re AtrazineBisphenol Herbicide Labeled antibodies flu Propanil Herbicide Polyclonal IgG antibody Fluo Acetamiprid Pesticide Electrochemical DNA Ferr Actararyl Pesticide Monoclonal antibody Surft	tibody Fluorimetry (<i>Acinetobacter</i> Light emission/ bioluminescence mprinting Surface plasmon : (MIP) resonance (SPR) tibodies Total internal reflection fluorescence (TIRF) igG antibody Fluorimetry	0.2 ng L ⁻¹ 2.5 mg L ⁻¹ 0.017 µg L ⁻¹ 0.002 µg L ⁻¹ 0.016 µg L ⁻¹ 0.016 µg L ⁻¹	Fast, sensitive, and cost-effective detectionA powerful tool in aquatic analytics Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	[151] [152] [144] [153]
Phenol EDC Whole-cell (Acinetobacter Lightsplan Triclosan EDC sp. DF4) bi Triclosan EDC Molecular imprinting Surfa AtrazineBisphenol Herbicide Labeled antibodies Tota AfstroneIsoproturon EDCEDCHerbicide Polyclonal IgG antibody Fluo Propanil Herbicide Electrochemical DNA Ferranci Acetamiprid Pesticide Electrochemical DNA Hi Carbaryl Pesticide Monoclonal antibody Surfa	(Acinetobacter Light emission/ bioluminescence mprinting Surface plasmon : (MIP) resonance (SPR) tibodies Total internal reflection fluorescence (TIRF) igG antibody Fluorimetry	2.5 mg L ⁻¹ 0.017 μg L ⁻¹ 0.002 μg L ⁻¹ 0.005 μg L ⁻¹ 0.016 μg L ⁻¹	Response to phenol-contaminated soils Excellent performanceFast, sensitive and simple nanosensor	[152] [144] [153]
Triclosan EDC Molecular imprinting Surfa- triclosan EDC Molecular imprinting Surfa- AtrazineBisphenol Herbicide Labeled antibodies Tota AEstroneIsoproturon EDCEDCHerbicide Polyclonal IgG antibody Fluo Propanil Herbicide Electrochemical DNA Ferr Acetamiprid Pesticide Electrochemical DNA Ferr Acetamiprid Pesticide Monoclonal antibody Surfi	mprinting Surface plasmon (MIP) resonance (SPR) tibodies Total internal reflection fluorescence (TIRF) igG antibody Fluorimetry	0.002 µg L ⁻¹ 0.005 µg L ⁻¹ 0.019 µg L ⁻¹ 0.016 µg L ⁻¹	Excellent performanceFast, sensitive and simple nanosensor	[144] [153]
AtrazineBisphenol Herbicide Labeled antibodies Total AEstronelsoproturon EDCEDCHerbicide Polyclonal IgG antibody Fluo Propanil Herbicide Polyclonal IgG antibody Fluo Acetamiprid Pesticide Electrochemical DNA Ferrini Acetamiprid Pesticide Monoclonal antibody Surfini Acetamiprid Pesticide Monoclonal antibody Surfini	tibodies Total internal reflection fluorescence (TIRF) gG antibody Fluorimetry	0.002 µg L ⁻¹ 0.005 µg L ⁻¹ 0.019 µg L ⁻¹ 0.016 µg L ⁻¹		[153]
Propanil Herbicide Polyclonal IgG antibody Fluo Acetamiprid Pesticide Electrochemical DNA Ferr Aptasensor hc Carbaryl Pesticide Monoclonal antibody Surfi	igG antibody Fluorimetry		Low-costEarly warning application	
Acetamiprid Pesticide Electrochemical DNA Ferra Aptasensor hc nä Carbaryl Pesticide Monoclonal antibody Surf.		0.6 ng L ⁻¹	Fast, sensitive, and cost-effective detection	[154]
Carbaryl Pesticide Monoclonal antibody Surfa	nical DNA Ferrocene (Fc)- based or hollow polymeric nanospheres (Fc- HPNs)	$3.3 \times 10^{-7} \text{nM}$	Excellent sensitivity	[155]
	l antibody Surface plasmon resonance (SPR)	1.38 μg L ⁻¹	Useful and portable for on-line monitoring	[156]
Urganopnosphorus (UPs) Pesticide A portable LIG-based Minc. electrochemical ai biosensor ar	LIG-based MnO ₂ -bridged enzyme- remical aided signal amplification	1.2 µg L ⁻¹	Good performance for pesticideApplications in the environment and food safety fields	[157]
Pyrethroid Pesticide Whole-cell and antibody Calo	and antibody Calorimetry	3 µg L ^{_1}	Simple and portable for on-site colorimetric applicationFunctional at least 90 days after lyophilization	[169]
NaphthalenePhenanthrene PAHs DNA/Sol-gel array Fluo	el array Fluorimetry	0–10 mg L ^{–1}	Effective PAHs detection in water, and biological samples	[158]
Polybrominated diphenyl POP Whole-cell (<i>Sphingobium</i> Bioluethers (PBDEs) xenophagum)	(<i>Sphingobium</i> Bioluminescence <i>jum</i>)	10 nM	Novel PBDE-specific sensing element	[16]
PCB-28PCB-101 PCB Enzyme (peroxidase)/ Amp immobilized polyaniline modified Pt electrode	eroxidase)/ Amperometry zed polyaniline Pt electrode	0.016 µg L ⁻¹ 0.019 µg L ⁻¹	Useful in POPs detection in landfill leachates	[159]
TPH TPH Antibody and enzyme Imm	nd enzyme Immunoassay	25 mg kg^{-1}	Quick on-site useDetect TPH in soil and water	[160]

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Product name	Analyte	Sensing element	Signaling element	Manufacturer (country)
Microtox	Toxicity	Whole-cell	Bioluminescence	Azur Environmental (US)
LUMIStox	Toxicity	Whole-cell	Bioluminescence	Hach (US)
ToxAlert	Toxicity	Whole-cell	Bioluminescence	Merck (Germany)
Catalytic DNA Sensors	Metals	DNAzyme	Fluorescence	Quasar Instruments (US)
PETRO RISc Soil Test	TPHs	Enzyme and antibody	Color	Ensys Energy Systems (US)
EnvironGard Petroleum Fuels in Soil	TPHs	Enzyme and antibody	Color	Merck Millipore (Germany)

Table 4. Commercial biosensors and the product details.

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highlighted the importance of high-throughput sequencing and synthetic biology tools on biosensors for organic pollutants since these techniques provide more genetic information about sensing elements [91,167]. Additionally, the usage of two different fluorescence as signaling elements enables the biosensor to detect different types of hydrocarbon simultaneously [165]. For simple and rapid on-site use, the colorimetric biosensors for pyrethroid insecticide and halogenated hydrocarbons performed well in testing samples [168,169]. So it can be states that a biosensor for detecting organic pollutants in the environment is now more efficient and feasible in rapid on-site screening. It complements traditional chemical analysis for large-scale monitoring.

Nowadays widespread industrial spillage is the major problem causing environmental degradation or destruction. For unique applications, biosensors can be personalized such as TPHs screening kit which is a favored tool for monitoring analytes on site in different media (soil, water and vapor) [170]. The immunoassay-based TPHs kit founded on antigen-antibody and enzyme is a biosensor due to its biological sensing elements. It has been suggested that TPHs immunoassay is one of the most widely used methods to quickly detect petroleum compounds especially in soil samples [171,172]. The low-cost and quick on-site utility are the main strengths of using the TPHs immunoassay kit [172]. However, the drawbacks of interference of media and nonspecific target were reported in TPHs assay kit, which is similar to other biosensors [171,172]. For this reason, the TPHs immunoassay could be used as large-scale screening method prior to precise quantification of TPHs using instruments such as GC. It could reduce the amount of energy consumed and carbon footprint as required by the SDGs.

An overview of the current achievements of biosensors and those devised to monitor organic pollutants was done in this section. Monitoring of organic contaminants plays an important role in the protection of human health and the ecosystem. By applying a biosensor, the rapid detection and screening of organic contaminants can be achieved. These biosensors are quite inexpensive, portable, available for on-site use, and no waste is generated during analysis. However, there are some limitations of these biosensors such as sensitivity and selectivity of the target pollutants, limiting applicability. Some challenges need to be overcome for biosensors including interference of humidity, effect of pH on sensitivity and selectivity, enhancing the sustainability of sensing components of biosensors, and ensuring the adaptation of nanomaterials, which can improve the performance of biosensors. More knowledge of metabolic genes for POPs is needed for further improvement of sensing elements. Thus, current biosensors technology could be utilized in monitoring EDCs, PCBs, POPs, TPHs, PAHs, and VOCs, while those difficult-to-degrade organic pollutants might not be efficiently monitored.

Similar to the biosensor detecting heavy metals, the cell-free biosensors using enzyme, DNA, and aptamer showed high sensitivity to organic pollutants including EDCs, pesticides, and PCBs (Table 3). However, the lack of bioavailability data and higher requirements of nanomaterials are also the limitations of cellfree biosensors. The whole-cell biosensors detecting organic pollutants are relatively few, which is due to the low biodegradability of the organic pollutants that hinders their development. In addition, most of the whole-cell biosensors have higher LOD than cell-free biosensors, showing the lower sensitivity (Table 3). Currently, the commercial whole-cell biosensors are mainly nonspecific and toxicitybased with the bioluminescence signals, whereas the biosensors specific to heavy metals and organic pollutants (TPHs) are all cell-free biosensors with DNAzyme, enzyme, and antibody (Table 4).

In short, establishing monitoring activities based on biosensors can be applied for the significant recognition/detection of heavy metals, chemicals, organic microorganisms, etc. Biosensors are robust environmental monitoring solutions with important novel characteristics and advantages [173]. Biosensors are early detection techniques with much promise in achieving on-site monitoring and continuous real-time environmental data. Wastewater and water monitoring biosensors are an innovative approach for the ultimate goal of on-site monitoring within the framework of the UN's Sustainable Development Goals (SDGs).

4.3 Wastewater quality monitoring and environmental viability

Wastewater discharge is the one of the source of environmental pollution, and the wastewater quality monitoring could be also achieved by the biosensors [145]. The biosensors especially MFCbased were applied in wastewater monitoring globally. In Asia Pacific countries, it has been reported that Au³⁺ ions in wastewater from Taiwan could be detected by engineered bacteria [174]. In addition, The MFC-based BOD biosensor was used in monitoring swine wastewater in Japan, which minimizes the energy cost for aeration [175]. The multiple heavy metals and phenols in real wastewater samples from China were able to be detected by MFC-based biosensors [176]. The immobilized engineered bacteria were used in screening PAHs in industrial wastewater in India [177]. The toxic metals (Cr⁶⁺ and Fe³⁺), NO₃⁻, and sodium acetate in the wastewater from Connecticut (USA) could be assessed by the batch mode MFC-based biosensors [178]. Similarly, the brewery wastewater obtained from Canada containing COD and others such as NH4⁺ was shown to successfully monitored by MFC-based biosensors [179]. The MFC-

based biosensors were also used in measuring the BOD in rice washed wastewater in Ecuador [180]. In South Africa, it has been shown that the influents and effluents from wastewater treatment plants were assessed by nonspecific whole-cell bioluminescence biosensors [181]. In Europe countries, the MFC-based biosensors were applied in detecting the BOD in domestic and brewery wastewaters in Hungary [182]. The COD in domestic wastewater from Spain discharged into constructed wetlands could be monitored by MFCbased biosensors [183]. Taken together, the MFCbased biosensors are broadly applied in wastewater quality monitoring due to its strengths of saving energy and continuous monitoring, which could correlated to the SDGs 6 (clean water and sanitation). However, further studies on detecting specific pollutants such as toxic metals in industrial wastewater are required for better wastewater quality. The performance and application of biosensors in real environmental samples such as river, wastewater, and soil samples were suggested in recent studies (Tables 2 and 3). For better commercial and environmental feasibility, more research about the real environmental sample tests is needed.

5. Biosensor for Sustainable Development Goals (SDGs)

Biosensors can detect impurities in the air, soil or water and in this way lead to finding the main sources of pollutants by implementing activities that mitigate impurities and help realize the SDGs, especially SDG 6 (clean water and sanitation), 12 (responsible consumption and production), 13 (climate action), 14 (life below water), and 15 (life on land). In developing countries, the environmental pollution might raise the health concern such as As contamination in groundwater, drinking water, and irrigation water, highlighting the needs for screening and monitoring the contaminants [184]. Compared with traditional environmental analysis methods, biosensors for environmental monitoring possess characteristics such as low-cost, minimal technical expertise and sample pretreatment, and feasibility for on-site use, saving energy and nonuse of hazardous materials [167]. For instance, the color-based

SDGs	Contributions of biosensor application
SDG 6: clean water and sanitation	• Large screening of environmental pollutants in waters for reducing the pollution in drinking water and aquatic ecosystemsImproving the access to safe and affordable drinking water due to ease of use and portability
SDG 12: responsible consumption and production	• Reducing the hazardous chemi- cals and wastes in sample pre- treatment and instrumental analysisBetter management of production and consumption in environmental monitoring to prevent the adverse impacts on human health and the environment
SDG 13: climate action	• Reducing the energy and corre- sponding carbon footprint from traditional chemical analysis
SDG 14: life below water	• Help managing the pollution into marine ecosystems
SDG 15: life on land	• Help managing the pollution into terrestrial ecosystems

 Table 5. The relationship of SDGs and biosensor application.

arsenic biosensor on paper strip can be easily used in drinking water by naked eyes without complicate instruments [14]. Therefore, applying the biosensors with such advantages on environmental monitoring could help achieving SDG 6 (clean water and sanitation) by accomplishing large screening of pollutants for clean water and improving the access to affordable drinking especially developing water in countries (Table 5). Furthermore, the SDG 6 (clean water and sanitation) and its target 6.3 (water quality and wastewater) could be achieved by using biosensors in simultaneous monitoring of multiple toxic metals in polluted rivers before human health is affected [71]. Water quality monitoring such as BOD, COD, and toxicants using the MFCbased biosensors and the toxicity-based nonspecific biosensors reflects not only the toxicity in drinking water but also the extent of eutrophication [23,81]. Targeted here are both SDGs 6 and 14 (life below water) which refers to detecting the potential pollution in freshwater and marine ecosystems as well as drinking water. For terrestrial ecosystems this refers to SDG 15 (life on land), and the soil contaminants-detecting biosensor could be applied to screen potential soil pollution, which can guide what happens in terrestrial ecosystems [185,186].

Furthermore, the results of biosensors provide more information about bioavailability and toxicity, which can complement traditional chemical analysis methods. From the sustainability point of view, biosensors' deployment reduces the amount of hazardous chemicals that are used in operational and sample pretreatment processes, which is also related to SDG targets 6.3 and 12.4 (reducing the hazardous waste produced) [167]. Portability and on-site monitoring feasibility using the biosensors reduce the carbon footprint which is a consequence of transportation. In addition, the large-scale screening of biosensor enviranalysis helps onmental with precise measurements and saves electricity and energy [167]. It has been reported that chemicals and used laboratories energy in make did a substantial contribution to the carbon footprint and green GHGs [20-22]. A summary of the relationship between SDGs and biosensor application is shown in Table 5. The complementary use of biosensors and traditional chemical analysis methods could realize SDG 13 (climate action) by reducing energy consumption and the carbon footprint.

6. Challenges of biosensor application on SDGs achievement

To provide reliable environmental monitoring results with low carbon footprint and less hazardous waste for achieving SDGs, biosensors need to be more feasible for quick-screening and onsite usage. The progress of biosensors which involves better selectivity, limits of detection and sensitivity is improving in the way that contaminants in the environments are detected, promoting a clean and green environment, while the challenges of sustainability, portability, and reusability still remain. Researchers want to resolve these challenges to enhance the biosensor application at advanced levels [187]. Currently, several advanced nanomaterials have been applied in the electrodes of cell-free and whole-cell biosensor to improve pollutants detection [188]. For cell-free biosensor, aptamer coupled with nanomaterials biosensors were commonly used for their good selectivity and reliability [29]. However, the cost for nanomaterials and aptamer-based biosensor production means they are still not commercially feasible, limiting their environmental application. The energy and carbon footprint from manufacturing these biosensors with nanomaterials and high performance might compromise their contribution to the SDGs. The possible risk of environmental nanomaterial use is still unclear so further information about the safety of nanomaterial-based biosensor and lowering the costs and energy consumption is needed. Recent studies mostly focused on sensing and signaling elements optimization, whereas the practicality tests using real wastewater samples are still rarely undertaken. More studies using biosensors in real environmental samples to assess the influence of other interfering substances such as organic matter chelation are warranted.

7. Conclusions and future prospects

Recognizing the growing call for more environmentally, economically, and socially responsible societies, emerging remediation technologies and governing strategies are being developed in alignment to the sustainable future. Biosensors using biological sensing elements coupled with signaling elements can be used to detect impurities in environmental samples so that the responsible sources of pollutants are known. By implementing remedial activities to remove harmful impurities from the sources, the UNSDG goals can be achieved. Despite the potential contribution to SDG achievements, few research reported the importance of biosensor application on SDGs, and the correlation of environmental monitoring using biosensors and SDGs. Herein, this article reviewed the current progress being made in biosensors for the purposes of environmental monitoring and their contribution to achieving the UNSDGs, including clean water and sanitation (SDG 6), responsible consumption and production (SDG 12), climate action (SDG 13), saving life below water (SDG 14), and saving life on land (SDG 15). While bio-sensing technologies have

advanced significantly in the past decades, further research on how to improve the sensitivity, stability, safety, and portability is warranted for environmental monitoring biosensors and their future applications.

Cell-free biosensors are powerful in terms of their quick and specific response to environmental pollutants, whereas a whole-cell biosensor can provide additional information about the bioavailability and toxicity that cannot be analyzed by cell-free biosensors. The results from nonspecific whole-cell biosensors directly reflect the total impact of pollutants on the test microorganisms, which is mostly based on the inhibition of microbial functions. However, nonspecificity is the main weakness when designing a specific targeting policy. On the other hand, the specific whole-cell biosensor can provide good information about the target pollutants, yet the sensitivity and stability still need to be improved for feasible commercial application. Recently, genetic methods have been developed in whole-cell biosensor to improve its sensing and signaling elements for better sensitivity. However, research into environmental sample tests, storage time such as shelf life, and safety of genetically modified biosensors is needed.

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Ethical statement

No ethics approval was required for this study as it involved no human participants or animals.

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