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## Expanding the circularity of plastic and biochar materials by developing alternative low environmental footprint sensors

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Flexible screen-printing technology combined with the use of a nano/material coating for improving electrode functionalities boosted the manufacturing of highly sensitive electrochemical sensors addressing the need for fast and easy-to-handle tests in different application fields. However, due to the large-scale production and disposable and single-use nature of these devices, their environmental footprint should be taken into careful consideration. Herein, the innovative reuse of post-consumer polyethylene terephthalate (PET) plastics as an alternative substrate coupled with biochar as an environmentally friendly and cost-effective modifier is described as a sustainable alternative for the production of robust electrochemical sensors. The good printability of reused plastics with graphite inks despite the chemical heterogeneity, different crystallinity, and surface roughness was demonstrated using atomic force microscopy and attenuated total reflection Fourier transform infrared spectroscopy. Functionalization with brewers' spent grain biochar enabled the fabrication of highly performing electrochemical sensors for nitrite detection in water having a limit of detection and a limit of quantification of 3.3 nM and 10.3 nM, respectively, with a linear range spanning from 0.01 to 500  $\mu$ M, and good reproducibility (RSD% 8%). The innovative intervention of the biochar-multilayer system markedly enhanced the electron transfer process at the electrode interface while simultaneously serving as an absorptive material for the investigated analyte. This work lays a foundation for repurposing end-of-life plastics for the electronics industry and presents a customizable reuse strategy aimed to keep the value of plastics in the economy and reduce waste and leakage into the natural environment.

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## Introduction

Polyethylene terephthalate (PET) is a thermoplastic polyester used worldwide to produce commodities in different application fields, mostly in food packaging and beverage sectors, owing to its lightweight, safety, transparency, flexibility, chemical inertia, durability, and thermal- and water-resistance.<sup>1,2</sup> PET polymers are also durable and have a slow degradation rate which makes them persisting materials, estimated to have a lifespan ranging from decades to hundreds of years.<sup>3,4</sup> Due to massive production, PET waste accumulation is un-

avoidable. In 2021, PET waste accounted for approximately 12% of global solid waste. Like other plastics, after leakage to different ecosystems, PET can break down into micro- and nano-plastics and pose grave risks to human health and wildlife.<sup>5</sup> PET is also one of the most recyclable and recycled materials whose physical-chemical properties enable multiple regeneration cycles or manufacturing of novel fibres, film sheets, and again novel food and beverage containers.<sup>6,7</sup> However, recycling or repurposing of damaged PET is not always appropriate for the food industry, thus the management of PET waste is directed toward other manufacturing end-users.<sup>8</sup> As a possible new industrial target, we envisage the electrochemical sensor industry, which has experienced phenomenal growth over the past few decades in response to the rising demand for innovative, smart, and disposable sensing devices for diagnostics, pharmaceuticals, agri-food, and environmental monitoring.<sup>9</sup> Electrochemical sensors are analytical devices that detect, quantify, and monitor target analytes even in complex matrices. The fundamental sensor

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design consists of a receptor, which binds the target analyte and produces a physical or chemical signal, and a transducer, which converts the analyte–receptor interaction into readable output signals.<sup>10</sup> Properly connecting the receptor–transducer coupling provides selectivity and sensitivity; nevertheless, various nano/materials, conducting polymers, and composites are frequently employed to functionalize the electrode transducer and enhance the analytical performance of the sensor.<sup>11,12</sup> Plastics are substituting glass, metals, and ceramics as substrates in screen-printing technology and for the development of wearable stretchy sensors, lab-on-a-chip, and point-of-care diagnostic tools.<sup>9</sup> The use of plastic materials in electrochemistry permits a greater degree of functional integration, including the introduction of wireless transmission modules, control and data acquisition instrumentation, and in-built power units, among others, and provides advantages over metallic and ceramic materials, including mild synthetic conditions, scalable and large-area processing, low operating temperature, and biocompatibility. However, the adoption of flexible sensors increases plastic demand and consumption, and the exploitation of high-footprint materials improving the performance of the electrodes increases waste.<sup>13</sup> In recent years, biochar has emerged as a valuable and effective electrode material and modifier, owing to its electrochemical properties combined with eco-friendly, economical, and renewable characteristics.<sup>14–16</sup> Biochar's electrochemical properties (pseudo-capacitance, electron conductivity, and double-layer capacitance) result from its heterogeneous composition, which consists of an organic aromatic matrix with different levels of oxidation and inorganic constituents whose composition varies depending on the original feedstock and synthesis process.<sup>17</sup>

In this paper, we propose repurposing post-consumer PET plastics for manufacturing robust and sustainable electrochemical sensors aiming to reduce the use of raw materials in the electronics sector while expanding the circularity of biomass derived from brewers' spent grain biochar. Towards this purpose, we developed highly sensitive biochar-based sensors for the detection of nitrite in water samples, by exploiting both the conventional Autostat™ polyester and post-consumer PET supports for the fabrication of screen-printed electrodes, namely SPEs and PET\_SPEs, respectively. We provided evidence on the usefulness and value of PET\_SPE exploitation and demonstrated the capability of multi-layered biochar to enhance the electrochemical performance of both types of electrodes, namely Bio-SPE and Bio-PET\_SPEs, respectively, after functionalization. The developed sensors demonstrated higher analytical performance, lower costs, and sustainability compared to previously realized devices.

## Results and discussion

### Morphological characterization

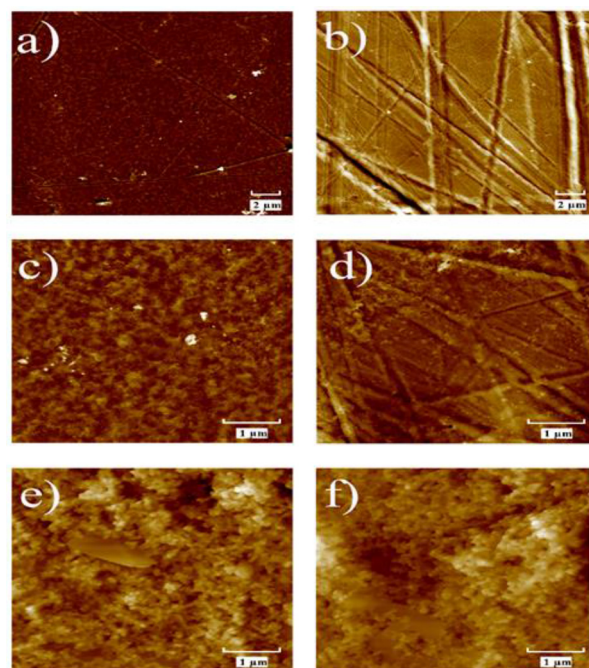
The surface morphology of plastic substrates and graphite-based screen-printed electrodes was investigated by atomic force microscopy (AFM) before and after graphite deposition.

The difference in the height of the morphological characteristics of plastic-based screen-printed electrodes after biochar modification (Bio-PET\_SPE) larger than 1  $\mu\text{m}$  made AFM not an appropriate investigation technique for this sample (see Bio-PET\_SPEs scanning electron microscopy images reported in our previous work<sup>19</sup>).

Fig. 1 shows the AFM images of the Autostat™ polyester and post-consumer PET substrates before and after graphite electrode deposition. 20  $\mu\text{m} \times 20 \mu\text{m}$  and 5  $\mu\text{m} \times 5 \mu\text{m}$  AFM images of the Autostat™ polyester substrate (Fig. 1a and c) show a porous morphology. AFM scans recorded in different locations of the same sample point to the similar morphology and value of surface roughness across the surface. In contrast, the surface morphology of the post-consumer PET substrate shows long and deep scratches and grooves (Fig. 1b and d), unevenly distributed on the sample surface, resulting in increased surface roughness and different roughness measured in different areas of the sample surface.

Table 1 reports the average root mean squared (RMS) roughness values and the related standard deviation calculated from AFM scans recorded in 3–5 different areas of the four measured samples. The standard deviation values provide a measure of the lateral uniformity of the sample.

Representative surface morphology of the graphite screen-printed electrodes is shown in Fig. 1e and f for both SPE and PET-SPE samples. Both AFM images show a small grain morphology and have height change across the surface of several



**Fig. 1** Surface morphology of plastics substrate and graphite-coated screen-printed electrodes. Representative 20  $\mu\text{m} \times 20 \mu\text{m}$  AFM images: (a) Autostat™ polyester, z scale = 100 nm; (b) post-consumer PET, z scale = 100 nm. Representative 5  $\mu\text{m} \times 5 \mu\text{m}$  AFM images: (c) Autostat™ polyester, z scale = 80 nm; (d) post-consumer PET, z scale = 50 nm; (e) SPE, z scale = 300 nm; (f) PET-SPE, z scale = 500 nm.



**Table 1** RMS roughness values calculated from  $5\ \mu\text{m} \times 5\ \mu\text{m}$  and  $20\ \mu\text{m} \times 20\ \mu\text{m}$  AFM scans and averaged over 3–5 images. The standard deviation values provide a measure of the lateral uniformity of the sample

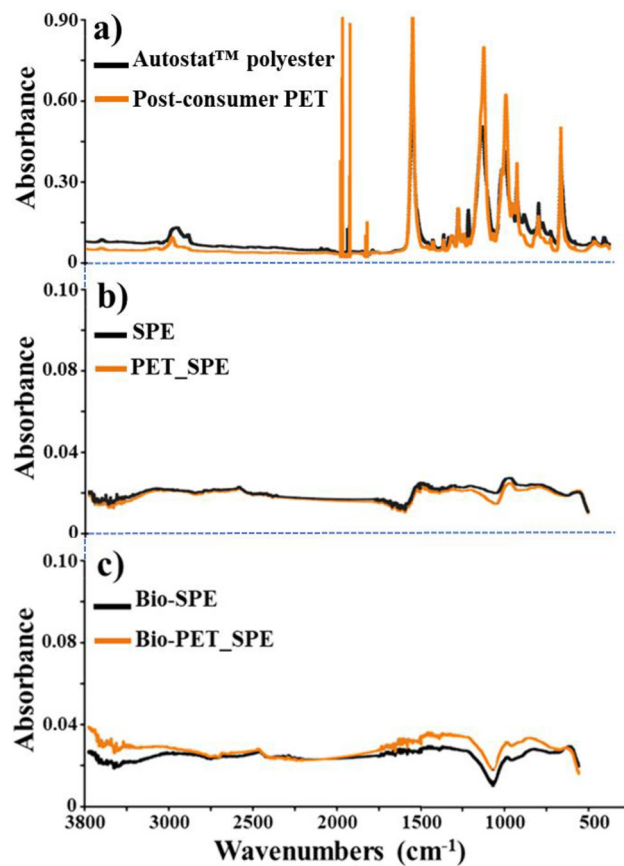
	Autostat <sup>TM</sup> polyester substrate		Post-consumer PET substrate	
	RMS (nm)	Stand. dev. (nm)	RMS (nm)	Stand. dev. (nm)
<b>20 <math>\mu\text{m} \times 20\ \mu\text{m}</math></b>				
Bare substrate	6.5	0.5	23.6	17.9
<b>5 <math>\mu\text{m} \times 5\ \mu\text{m}</math></b>				
Graphite-screen-printed electrodes	5.2	0.3	12.9	11.6
	82.6	22.4	68.5	12.6

hundredth nm ( $z$  scale of the AFM images) resulting in RMS roughness greater than that of the bare substrate samples and quite different in different surface areas (greater standard deviation, see Table 1). Also, the large height changes hindered recording of high-quality, reliable AFM scans larger than  $5\ \mu\text{m} \times 5\ \mu\text{m}$ .

However, and noteworthily, graphite screen-printed on post-consumer PET is smoother than the one printed on Autostat<sup>TM</sup> polyester as shown by both the lower value of the average RMS roughness and the decreased standard deviation (see Table 1).

Attenuated total reflection Fourier transform infrared spectroscopy (ATR-FTIR) was performed to characterize the chemical composition and polymer crystallinity of the screen-printed electrodes and plastic substrates (Fig. 2).

Autostat<sup>TM</sup> polyester and post-consumer PET samples displayed sharp peaks at  $\sim 1715\ \text{cm}^{-1}$  and  $\sim 725\ \text{cm}^{-1}$ , attributed to C=O stretching and C=O out-of-plane bending vibrations, respectively (see Fig. 2a). Several ring stretching and bending vibrations provided bands at  $\sim 1410\ \text{cm}^{-1}$ ,  $\sim 1020\ \text{cm}^{-1}$ , and  $\sim 872\ \text{cm}^{-1}$ . Moreover, both samples showed broad peaks at  $\sim 1240\ \text{cm}^{-1}$  and  $\sim 1100\ \text{cm}^{-1}$ , attributed to C(=O)–O stretching, ring-ester CC stretching, C=O in-plane bending and C–O stretching vibrations, respectively (see Fig. 2a).<sup>20–22</sup> The bands at  $1470$ ,  $1340$ ,  $1120$ ,  $970$ , and  $849\ \text{cm}^{-1}$  are markers of the crystalline structure of PET; their intensities appear higher in the spectrum of Autostat<sup>TM</sup> polyester, compared to that of post-consumer PET. However, bands at  $1370$ ,  $1044$ , and  $898\ \text{cm}^{-1}$ , characterizing the amorphous confirmation of samples, show a higher intensity for post-consumer PET material. Furthermore, FTIR spectra performed exactly on the Autostat<sup>TM</sup> polyester electrode (SPE) and post-consumer PET electrode (PET\_SPEs) are nearly superimposable (Fig. 2b), demonstrating that the printed graphite layer, which is at least  $2\ \mu\text{m}$  thick, makes the surfaces extremely similar despite that the pristine plastic substrates were largely dissimilar as also observed in the AFM images (Fig. 1c–f). These results revealed the functional equivalence of post-consumer PET and Autostat<sup>TM</sup> polyester for our screen-printing technology, leading us to proceed with the subsequent steps of the sensor assembly, *i.e.*, the layer-by-layer deposition of biochar to construct the Bio-



**Fig. 2** ATR-FTIR characterization: (a) Autostat<sup>TM</sup> polyester and post-consumer PET, (b) Autostat<sup>TM</sup> polyester electrode (SPE) and post-consumer PET electrode (PET\_SPE), (c) biochar-modified electrodes: SPE and Bio-PET\_SPE. Spectra are reported separately for clarity.

PET\_SPEs. FTIR analyses of Bio-PET\_SPEs (Fig. 2c) provided further support for the feasibility of using post-consumer PET as a substrate in electrode printing.

### Electrochemical characterization

An in-depth study was hence performed to verify the electrochemical behaviour of the developed electrodes for the fabrication of reliable sensors. To that end, the electrochemical performances of SPEs and PET\_SPEs before and after biochar modification were compared. Initially, the background current was measured in 50 mM KCl solution for each electrode. The following currents were registered:  $29 \pm 4$ ,  $34 \pm 5$ ,  $5.2 \pm 0.3$ , and  $6.4 \pm 0.4\ \text{nA}$  for SPEs, PET\_SPEs, Bio-SPEs and Bio-PET\_SPEs, respectively ( $n = 10$  electrodes for each electrode type). However, the following signal-to-noise ratios (S/N) corresponding to the current measured in the presence (10 mM  $[\text{Fe}(\text{CN})_6]^{3-/4-}$  in 100 mM KCl) and absence (only 100 mM KCl) of the redox probe were obtained: 230, 226, 476, and 473 for SPEs, PET\_SPEs, Bio-SPEs and Bio-PET\_SPEs, respectively. As previously observed for SPEs,<sup>12</sup> PET\_SPEs also showed a 5-fold decrease of the capacitive current associated with a 2-fold increase of the signal-to-noise ratio (S/N) when modified with biochar. Thus, both unmodified and biochar-modified electro-



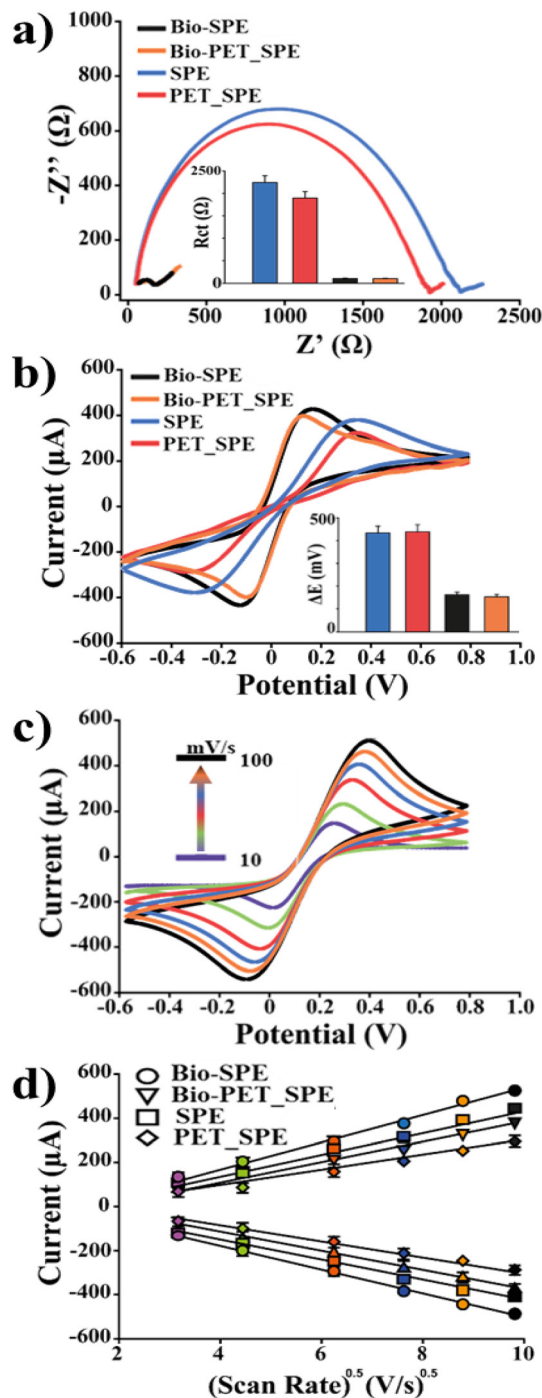
des demonstrated the applicability of post-consumer recycled plastic as a substrate material for the fabrication of trustworthy electrochemical devices. Impedance electrochemical spectroscopy (EIS) and cyclic voltammetry (CV) were used as complementary tools to investigate the electron transfer and diffusivity processes of the electrode/electrolyte interface in 10 mM  $[\text{Fe}(\text{CN})_6]^{3-/4-}$  in 100 mM KCl solution (see Fig. 3). From the respective Nyquist plots and voltammograms, reported in Fig. 3a and b, it is possible to observe that the modification with biochar produced a dramatic improvement of the sluggish surface kinetics typical of bare electrodes (both SPEs and PET\_SPEs) as demonstrated in our previous works.<sup>12</sup>

This is evidenced by the decrease in the generally high charge transfer resistance ( $R_{ct}$ , diameter of the semicircle) of bare platforms compared to that of biochar-modified SPEs and PET\_SPEs. Specifically, the calculated  $R_{ct}$  values for bare SPEs and PET\_SPEs were  $1.8 \pm 0.2 \Omega$  and  $1.9 \pm 0.2 \Omega$ , respectively, and achieved 10-fold lower values after the modification with biochar (*i.e.*,  $0.15 \pm 0.01 \Omega$  and  $0.17 \pm 0.01 \Omega$ , for Bio-SPE and Bio-PET\_SPEs, respectively). The CV study also found this enhancement in the electron transfer kinetics of the redox process between the electrode/electrolyte interface attributable to biochar modification. This is demonstrated by the ideal trend of the anodic and cathodic peak ratio ( $I_{pa}/I_c = 1$ ), the decrease of  $\Delta E$  (from 0.35 to 0.16 and 0.40 to 0.18 V for SPEs and PET\_SPEs, respectively), and the correlated  $k^0$  and  $k^0$ , calculated using Randles's and Marcus's theories (reported in Theoretical methods section),<sup>11,23</sup> respectively. The correlated results are shown in Table 2.

A study of the diffusivity process occurring at the electrode interface was carried out. In Fig. 3c the experiment performed using Bio-PET\_SPEs is reported as an example. By examining the effect of the scan rate on the current peaks, it is possible to observe that the faster the scan rates, the smaller the size of the diffusion layer and, as a result, the higher the peak current. The diffusion coefficient ( $D_0$ ) was calculated as the average of the anodic and cathodic coefficients ( $D_{Ox}$  and  $D_{Red}$ ) using the Randles-Sevcik equation (eqn (4), reported in the Theoretical methods section).

As a result, the diffusivity of bare SPEs and PET\_SPEs comparable to the one reported in the literature by Konopka and McDuffie was assessed.<sup>24</sup> Analogously, a similar planar diffusion-controlled process in the oxidation/reduction reactions of the selected redox probe was found for Bio-SPEs and Bio-PET\_SPEs. Subsequently,  $I_{pa}$  and  $I_{pc}$  were plotted as a function of the scan rate ( $\nu^{0.5}$ ). Fig. 3d clearly shows the direct proportionality between the peak current and the square root of the scan rate according to the Randles-Sevcik equation.

The estimated slopes ( $\mu\text{A} (\text{mV s}^{-1})^{-0.5}$ ) of the linear correlation of  $I_{pa}$  and  $I_{pc}$  are as follows: 183.4 ( $R^2 = 0.999$ ) and  $-183.6$  ( $R^2 = 0.999$ ), 148.1 ( $R^2 = 0.998$ ) and  $-148.2$  ( $R^2 = 0.999$ ), 157.7 ( $R^2 = 0.997$ ) and  $-158.2$  ( $R^2 = 0.995$ ), and 132.1 ( $R^2 = 0.997$ ) and  $-130.8$  ( $R^2 = 0.997$ ), corresponding, to the SPE, PET\_SPEs, Bio-SPE, and Bio-PET\_SPEs. Table 3 provides a summary of the characterization of the analytical performances of the four different electrodes.



**Fig. 3** Electrochemical characterization of the electron transfer and diffusivity process. In Fig. (a) and (b): improved conductivity observed by characterizing impedimetrically and voltammetrically an Autostat™ polyester electrode (SPE), post-consumer PET electrode (PET\_SPE) and Biochar modified electrodes (Bio-SPE and Bio-PET\_SPE). Fig. (c) depicts the scan rate analysis conducted on Bio-PET SPE, whereas Fig. (d) illustrates the corresponding calibration plot for the square root of the scan rate vs. the oxidation current peak.

An in-depth investigation in terms of LOD, LOQ, and reproducibility of the developed sensors was performed by using ferro-ferricyanide, ascorbic acid (AA), uric acid (UA) and hydro-



**Table 2** Comparison of the electrochemical parameters of SPE and PET\_SPE, before and after biochar modification

	CV			EIS	
	$I_{pa}/I_{pc}$	$D_0 \times 10^{-6} \text{ (cm}^2 \text{ s}^{-1}\text{)}$	$k^0 \times 10^{-3} \text{ (cm s}^{-1}\text{)}$	$W \text{ (K}\Omega\text{)}$	$k^{0'} \times 10^{-5} \text{ (cm s}^{-1}\text{)}$
SPE	$0.8 \pm 0.1$	$1.4 \pm 0.3$	—	$0.5 \pm 0.1$	$1.6 \pm 0.3$
PET_SPE	$1.3 \pm 0.1$	$1.2 \pm 0.2$	—	$0.5 \pm 0.1$	$2.3 \pm 0.2$
Bio-SPE	$1.1 \pm 0.1$	$2.1 \pm 0.2$	$2.4 \pm 0.2$	$0.6 \pm 0.1$	$7.4 \pm 0.3$
Bio-PET_SPE	$1.1 \pm 0.1$	$2.4 \pm 0.2$	$2.1 \pm 0.2$	$0.6 \pm 0.1$	$7.4 \pm 0.3$

**Table 3** Square wave voltammetry (SWV) analysis: limit of detection (LOD), limit of quantification (LOQ), sensitivity and inter-electrode reproducibility calculated for potassium ferricyanide, ascorbic acid, uric acid and hydroquinone for SPE, PET\_SPE, Bio-SPE and Bio-PET\_SPE

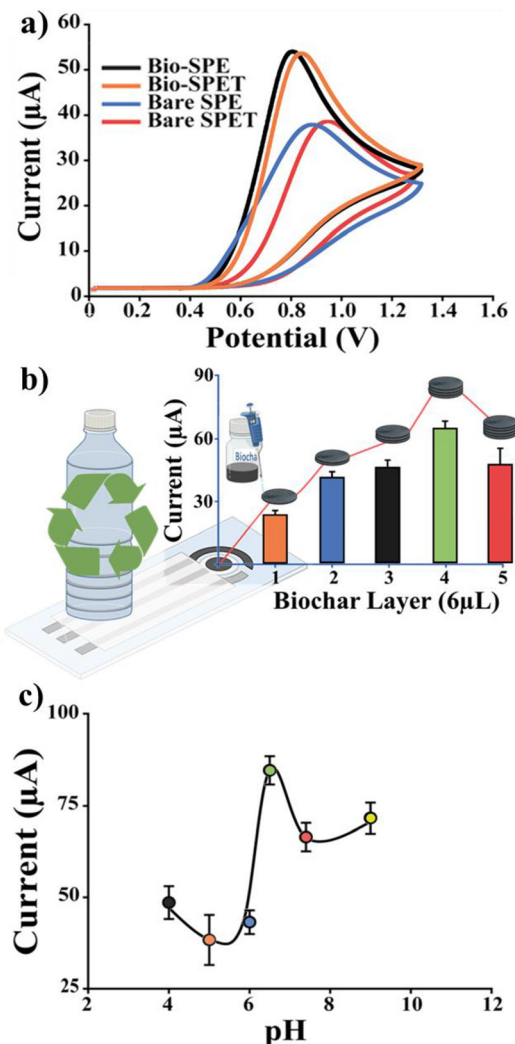
	SPE	PET_SPE	Bio-SPE	Bio-PET_SPE
<b>Ferricyanide</b>				
LOD ( $\mu\text{M}$ )	9.2	10.4	3.4	4.2
LOQ ( $\mu\text{M}$ )	28	30.2	10.5	12.1
Sensitivity ( $\text{mA Mcm}^2\text{)}^{-1}$	40	40.7	20.2	21.2
Reproducibility (RSD%)	9	10	4	5
<b>Uric acid</b>				
LOD ( $\mu\text{M}$ )	7.4	6.7	2.1	2.3
LOQ ( $\mu\text{M}$ )	22.3	20.7	7.3	8.7
Sensitivity ( $\text{mA Mcm}^2\text{)}^{-1}$	49.4	36.8	26.3	27.1
Reproducibility (RSD%)	9	10	5	5
<b>Ascorbic acid</b>				
LOD ( $\mu\text{M}$ )	3.2	4.7	0.5	0.8
LOQ ( $\mu\text{M}$ )	9.7	14.3	1.8	3.5
Sensitivity ( $\text{mA Mcm}^2\text{)}^{-1}$	21.3	12.7	5.9	6.9
Reproducibility (RSD%)	9	10	5	5
<b>Hydroquinone</b>				
LOD ( $\mu\text{M}$ )	9.2	10.5	2.1	1.9
LOQ ( $\mu\text{M}$ )	28	31.7	10.2	8.9
Sensitivity ( $\text{mA Mcm}^2\text{)}^{-1}$	40	42.2	19.4	16.8
Reproducibility (RSD%)	9	8	6	5

quinone (HQ) as electroactive probes and performing square wave voltammetry (SWV) as the analytical technique. The reproducibility calculated for six different electrodes of each substrate showed comparable results between bare and biochar-modified electrodes: RSD% of 10, 11, 5 and 5% for SPEs, PET\_SPEs, Bio-SPEs and Bio-PET\_SPEs. These results demonstrate that conventional SPEs and PET\_SPEs share the same analytical performance, as evidenced by the identical LOD, LOQ, and reproducibility values observed by analysing the different probes. This behaviour is also observed for biochar-modified electrodes, confirming the analytical robustness of PET\_SPEs and their suitability for sensing applications.

#### Application of PET\_SPEs to the detection of nitrite in a real sample

Biochar-modified PET\_SPEs were used to develop an electrochemical sensor for quantifying nitrites in drinking water; their levels are regulated by government directives due to their potential toxicity for environmental and human health.<sup>25,26</sup> Regarding this, we designed a biochar-multilayer system capable of adsorbing the analyte, which is then detected using SWV voltammetry.

A careful optimization of parameters such as the deposited layers of biochar, the effect of pH on nitrite oxidation conditions, and the nitrite adsorption time on the carbonaceous materials layer was explored, and the corresponding results are shown in Fig. 4. Considering the complexity of nitrite reduction reaction, which may involve interferences such as



**Fig. 4** Optimization of nitrite oxidation using Bio-PET\_SPE: (a) voltammograms obtained analysing a constant concentration of  $\text{NO}_2^-$  ( $100 \mu\text{M}$ ) using SPEs, PET\_SPE, Bio-SPE and Bio-PET\_SPE; (b) SWV current as a function of the number of Biochar layers deposited at constant nitrite concentration; (c) currents as a function of pH.



nitrate ions and reduction of molecular oxygen, oxidation is the most efficient method for electrochemically detecting nitrite. Initially, the behaviour of nitrite was investigated by CV measurements across all our platforms (SPEs, PET-SPEs, Bio-SPEs and Bio-PET\_SPE).

In Fig. 4a, the voltammograms obtained analysing a constant concentration of  $\text{NO}_2^-$  ( $100 \mu\text{M}$ ) using SPEs, PET-SPEs, Bio-SPEs and Bio-PET\_SPE are reported. According to Fig. 4a, biochar-modified platforms exhibited well-defined current peaks ( $0.75$  and  $0.81$  V for Bio-SPE and Bio-PET\_SPE, respectively) accompanied by a shift to a lower oxidizing potential compared to unmodified screen-printed electrodes ( $0.91$  and  $0.99$  for SPE and PET\_SPE, respectively). This can be attributed to the enhancing effect of biochar, which promotes the oxidation of  $\text{NO}_2^-$  in two ways: by exhibiting high electrocatalytic activity because of the presence of carbonyl and carboxyl groups and by increasing the surface area of the working electrode (WE). Specifically, to maximize the latter effect, biochar was deposited in multiple layers.

Fig. 4(b) and (c) show the effect of multiple layers of biochar and pH optimization on the detection of nitrite. SWV was performed using a constant nitrite concentration ( $100 \mu\text{M}$ ) on each biochar deposited layer. As anticipated, the greater the number of layers, the larger the electrochemically active surface area ( $A$ ,  $0.12$ ,  $0.25$ ,  $0.47$ ,  $0.63$ ,  $0.78 \text{ cm}^2$ , for bare, 1, 2, 3 and 4 layers, respectively), and the higher the measured current. However, for more than 5 layers (not reported here), a decrease in the peak current was observed compared to 4-layer biochar-modified PET\_SPEs (4-Bio-PET\_SPEs). This is because the thick layer of biochar significantly reduces nitrite's diffusivity, thereby nullifying the effect of the increased surface area ( $A$ ). Considering the best results obtained with 4-Bio-PET\_SPEs, this platform was applied to explore the effect of pH on  $\text{NO}_2^-$  oxidation. As depicted in Fig. 4c, the maximum peak current was observed at pH 6.5; consequently, this value was chosen for further research. In contrast, the peak current decreases at pH values below or above 6.5. A possible explanation is that at low pH levels (below 4) protonation of nitrite occurs, forming nitrous acid thereby affecting its redox activity.<sup>6</sup> A more basic pH, on the other hand, affects the stability of the silver reference electrode due to the formation of a passivating oxide layer.

At this point, the sorption capacity of biochar was determined by measuring the current as a function of time (0, 5, 10, and 20 minutes) while utilising the same nitrite concentration ( $100 \text{ M}$ ). Fig. 5 illustrates the relative results.

Fig. 5a depicts the biochar's negligible sorption effect on the analyte as a function of time, despite its well-documented sorption properties over N-based compounds. Typically, the biochar coatings used for sorption are thicker, so despite the four layers of biochar deposited in this case, the area is too small to observe this effect as a function of a short timeframe. After the oxidation conditions of nitrite were optimized, the analytical performances of Bio-PET\_SPE as  $\text{NO}_2^-$  sensors were investigated. 4-Bio-PET\_SPEs were tested at different concentrations of nitrite in PBS pH 6.5 (from 0 to 5 mM) using SWV

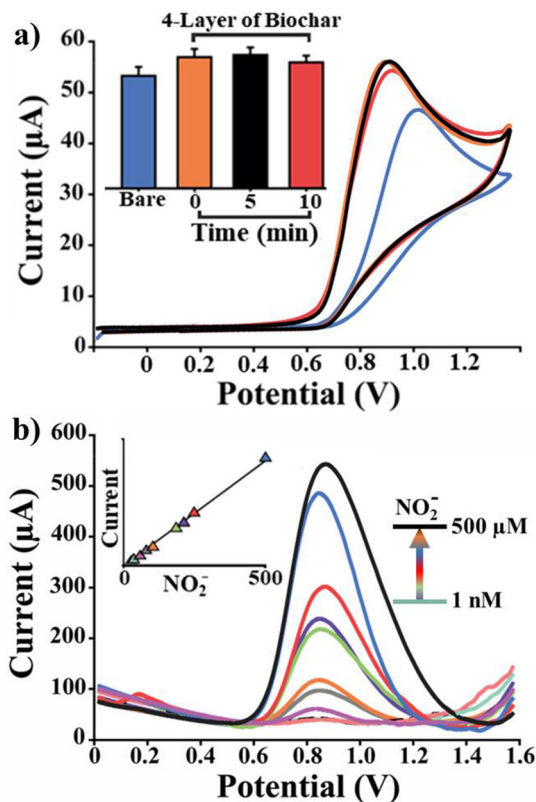


Fig. 5 Analytical performances of Bio-PET\_SPE based  $\text{NO}_2^-$  sensor at different nitrite concentrations from 0 to 500  $\mu\text{M}$  in 0.01 M PBS (pH 7.4) by performing CV (a) and SWV (b) as electrochemical techniques. In the inset of Fig. (a) biochar's sorption effect on the analyte as a function of time is reported.

as the analytical technique. The following analytical parameters were derived from the voltammograms shown in Fig. 5b and the relative calibration line (slope = 0.012,  $R^2 = 0.998$ ): LOD 2.91 nM, LOQ 10.2 nM, linear range from 0.01 to 500  $\mu\text{M}$ , and sensitivity  $0.174 \mu\text{A} \mu\text{M}^{-1} \text{ cm}^2$ . 4-Bio-SPEs analyzed under identical conditions exhibited nearly identical performances (LOD 2.6 nM, LOQ 10.3 nM, linear range 0.01 to 500  $\mu\text{M}$ , and sensitivity  $0.167 \mu\text{A} \text{ M}^{-1} \text{ cm}^2$ ), demonstrating once again the remarkable comparability of the performances of the Autostat<sup>TM</sup> polyester electrode (SPE) and post-consumer PET electrode (PET\_SPEs), and the viability of reusing PET. Furthermore, nitrite concentrations were measured using mineral water (zero nitrite certified samples) for real sample analysis by the standard addition method in the range of 0.1–100  $\mu\text{M}$  nitrite.

Extremely promising results were observed in terms of sensitivity LOD of 3.0 and 3.3 nM, reproducibility RSD = 7 and 8% for SPE and PET\_SPE-based sensors, respectively, and recovery. Data reported in Table 4 highlight the reliability of Bio-PET\_SPEs to detect nitrite as indicated by the recovery percentage ranging from 90 to 99%. In addition to the similarity of the electrochemical performance between SPEs and PET\_SPE-based sensors, good reproducibility (always < 10%) was observed on using even nanomolar concentrations of  $\text{NO}_2^-$ .



**Table 4** Determination of NO<sub>2</sub><sup>-</sup> concentrations in mineral water

	Concentration of NO <sub>2</sub> <sup>-</sup>		
	Spiked (C <sub>S</sub> , μM)	Recovered (C - C <sub>0</sub> , μM)	Recovery % (n = 6)
Bio-SPE	0.1	0.92 ± 0.05	92
	10	9.4 ± 0.4	94
	100	101 ± 5	101
Bio-PET_SPE	0.1	0.90 ± 0.09	90
	10	9.3 ± 0.6	92
	100	99 ± 5	99

### Comparison of the performance and cost-benefit analysis

The performance of the biochar-based sensor under study was compared to earlier NO<sub>2</sub><sup>-</sup> sensors reported in the literature. According to Table 5, both SPE and PET\_SPE-based sensors exhibited a significantly lower LOD and wider linear range when modified with a biochar-multilayer system. It is essential to highlight the analytical performance of the proposed electrodes, especially in light of the simplicity of their construction and the predicted costs, which are roughly ten times lower than those of their competitors. Furthermore, it should be noted that this work addresses the challenge outlined in the European Union's "European Strategy for Plastics in a Circular Economy" starting that in 2030, all plastic packaging placed on the EU market will be either reusable or recyclable at a low cost.<sup>18,27</sup> Indeed, this method proposes the sustainable and cost-effective reuse of post-consumer plastics for technological applications beyond their initial discarding application.

**Table 5** Comparison of 4-Bio-PET\_SPE performance to earlier nitrite sensors reported in the literature<sup>28–36</sup>

Device	LOD (nM)	Linear range (μM)	Matrix	Ref.
SPCE/AuNPs-PEI	2.5	0.01–4.0	Water	28
SPCE/GR-CD	260	0.7–2150	Water	8
SPCE/NBC180	2080	100–400	Water/wine	9
LIG/f-MWCNT-AuNPs electrode	900	10–140	Tap water	10
GCE/GO-CS-AuNPs	300	0.9–18.9	PBS buffer	11
GCE/MoS <sub>2</sub> -AuNPs	90	10–2100	River and drinking water	12
Bare-GCE	400	0.5–1000	Lake water	13
Ur-Pd/SWCNT electrode	250	2–238	Deionised water	14
CPE/f-MWCNT/1,8-DAN	75	0.3 to 65	Tap water	36
4-Bio-SPE	3	0.01 to 500	Mineral water	This work
4-Bio-PET_SPEs	3.3	0.01 to 500		

Acronyms: screen-printed carbon electrode (SPCE), gold nanoparticles (AuNPs), polyethyleneimine (PEI), graphite (GR), β-cyclodextrin (CD), nanobiochar obtained by hydrothermal carbonization 180 °C (NBC180), laser-induced graphene (LIG), COOH functionalized multiwalled carbon nanotubes (f-MWCNT), glassy carbon electrode (GCE), graphene oxide (GO), chitosan (CS), molybdenum disulfide (MoS<sub>2</sub>), urchin-like palladium nanostructures (Ur-Pd), single-walled carbon nanotubes (SWNT), poly 1,8-diaminonaphthalene (1,8-DAN).

## Conclusions

Tackling the plastic pollution problem involves every part of the plastic value chain – from sourcing to end-of-life, and direct reuse of post-consumer products could be the quickest and most sustainable pathway towards plastic circularity.

The method developed in this work is shown to be an easy, sustainable, and low-cost alternative for SPE construction by repurposing post-consumer PET to realize environmentally sustainable sensors with robust analytical performances. The peculiarity of the proposed method lies in the combined use of inexpensive PET beverage packaging as the substrate material for printing of serigraphic electrodes and biochar, which has the unique ability to function both as an absorbent substrate and as an enhancing material of the sluggish electron transfer kinetics of graphite-based electrodes.

Herein, we realized innovative, low-environmental footprint PET-biochar-based sensors having high sensitivity for detection of nitrite in water samples. The analytical parameters of Bio-PET-SPE sensors indicated sensitivity in the nanomolar range (LOD = 3 nM), high reproducibility (RSD% < 10%) and recovery in real sample analysis (always ≥ 90%), proving the viability of these devices for water monitoring.

These results showed for the first time that a biochar multi-layer functionalization strategy can be adopted to achieve highly improved performance in terms of sensitivity compared to previously reported sensors.

Currently, we are developing an enzyme-free sensor based on PET\_SPEs functionalised with metal-incorporating biochar to simulate nitrite oxidoreductase activity while improving the selectivity of our devices. Combined with a microfluidic system, these sensors hold the promise to be effective devices for real-time and on-site water monitoring.

## Experimental section

### Materials

All commercially available compounds were of analytical grade. Biochar was produced by pyrolytic micro-gasification (*T* = 400 °C) in an Elsa D17 micro pyrolytic reactor (Bluecomb Ltd., Udine, Italy).<sup>19</sup> Commercially available Autostat™ (HT5) polyester was purchased from Sirpi S.r.l (Milan, Italy), whereas post-consumer PET substrates are from water bottles and food packaging. Sigma-Aldrich (Steinheim, Germany) supplied sodium nitrite, hydroquinone, L-ascorbic acid, uric acid, and potassium chloride. Ferrocyanide and potassium ferricyanide were acquired from Fluka Chemie, Sigma-Aldrich (Buchs, Switzerland). The buffer solutions were 0.05 M phosphate buffered saline (PBS) + 0.1 M KCl with pH 7.4 and 0.05 M carbonate buffer (CB) with a pH of 9.0.

### Apparatus and experimental

All AFM images of the graphite-based screen-printed electrodes (SPE and PET\_SPEs) before and after biochar modification (only Bio-PET\_SPEs) were recorded in air in Scanasyst@ mode



using a Multimode 8 microscope (BRUKER, USA) equipped with silicon nitride probes (SCANASYST AIR, BRUKER, USA) having a nominal curvature radius of 2 nm and an elastic constant of  $0.4 \text{ N m}^{-1}$ .  $512 \times 512$  points AFM scans were collected from 3–5 randomly chosen locations of the samples and with different scan sizes in order to check the lateral uniformity of the surface morphology. Dimensions of the scans varied between  $2 \mu\text{m} \times 2 \mu\text{m}$  and  $20 \mu\text{m} \times 20 \mu\text{m}$ . Root mean square (RMS) roughness values were calculated using Nanoscope Analysis software (BRUKER, USA).

FTIR absorption spectra of electrodes were acquired with a Thermo-Scientific instrument (model Is50) (Thermo Scientific Inc., Madison WI USA) in attenuated total reflectance (ATR) mode using a single reflection diamond cell. Spectra were recorded from  $4000$  to  $525 \text{ cm}^{-1}$ , averaging over 32 scans with a resolution of  $2 \text{ cm}^{-1}$ . All experiments were performed in triplicate, yielding consistent and reproducible results.

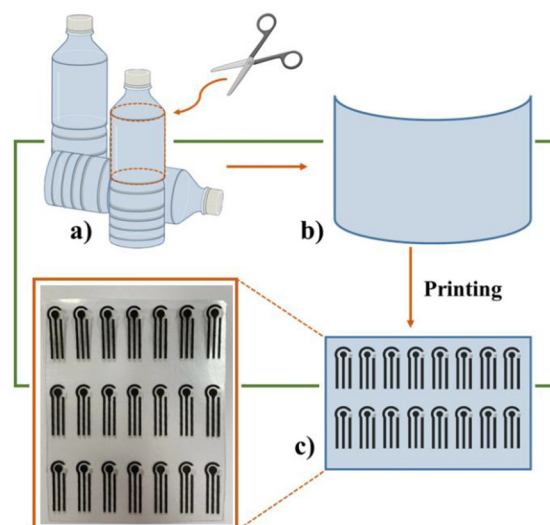
Amperometry, CV, SWV and EIS analyses were performed using a PalmSens Instrument (PalmSens, Netherlands). A Hielscher UP200St-ultrasonic transducer was used for the preparation of biochar dispersions. Chronoamperometry ( $0.4 \text{ V}$ ,  $150 \text{ s}$ ) was performed to measure the background current in  $50 \text{ mM KCl}$  solution by examining six different electrodes ( $n = 6$ ) for each type of substrate.

### Preparation of PET\_SPEs

SPEs and PET\_SPEs were manufactured in-house with a 245 DEK high-performance, multi-purpose, precision screen-printing machine. Specifically, both the working electrode (WE, geometric area  $\sim 0.07 \text{ cm}^2$ ) and counter electrodes (CEs) were printed on graphite-based ink (Elettrodag 421 at  $80 \text{ }^\circ\text{C}$ ) while silver ink was used to print the reference electrode (RE at  $80 \text{ }^\circ\text{C}$ ). The only distinction between SPEs and PET\_SPEs is the used polyester.<sup>36</sup> In detail, a small portion of used PET that is in good condition overall is identified. This material (flat or made flat by the application of physical weight) is initially rinsed with tap water and soap to remove any remaining dust and coarse dirt. After that, it is rinsed with a 4 : 1 (v/v) ethanol/distilled water solution. This process removes any dirt or grease and prepares the surface for printing. To maximize the number of electrodes that can be printed on the portion of the recycled PET, the substrate is then cut to the appropriate dimensions and utilized in the fabrication of PET\_SPEs. In Scheme 1 a representation of the PET\_SPE fabrication is reported.

### Preparation of Bio-PET\_SPEs

SPEs and PET\_SPEs were amperometrically pre-treated (a steady anodic potential of  $+1.7 \text{ V}$  was applied for  $180 \text{ s}$ ) in  $0.05 \text{ M}$  phosphate buffer  $+0.1 \text{ M KCl}$ , pH 7.4 solution. Then, a rinsing step ( $100 \mu\text{L}$ , three times) was carried out to remove any eventual salt residues, and  $6 \mu\text{L}$  of a biochar dispersion ( $1 \text{ mg mL}^{-1}$  in 1 : 3 v/v ethanol-water solution), prepared using an ultrasonic transducer ( $200 \text{ W}$ ,  $26 \text{ kHz}$ ,  $30 \text{ minutes}$ ), was deposited by drop-casting onto the SPE's and PET\_SPE's WE. The deposited solution is dried by heating the electrodes for  $15 \text{ minutes}$  at  $38 \text{ }^\circ\text{C}$ . To prepare multiple biochar-deposited



**Scheme 1** Fabrication of PET\_SPEs: (a) and (b) plastic bottles as starting and printing substrate materials, (c) the final PET-based electrodes.

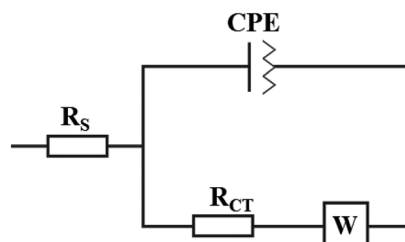
films, the final operation is repeated as many times as the desired number of coatings.

### Theoretical methods

The impedimetric experimental data were fitted using a common Randles circuit (see Scheme 2). This equivalent circuit, one of the simplest possible models for describing electrochemical interface processes, consists of an active electrolyte resistance  $R_s$  in series with parallel combination of the double-layer capacitance constant phase element (CPE). In addition to the previously mentioned components, the charge transfer resistance ( $R_{ct}$ ) and Warburg impedance are also essential ( $W$ ).<sup>15</sup>

The electronic transfer process was studied following the redox process of the reversible couple ferro-ferricyanide:  $[\text{Fe}(\text{CN})_6]^{3-} + 1e^- \rightleftharpoons [\text{Fe}(\text{CN})_6]^{4-}$ . The heterogeneous electron transfer constant was calculated voltammetrically ( $k^0$ , Marcus' theory, eqn (1)) and impedimetrically ( $k^{0r}$ , Randles's theory, eqn (2)).  $k^0$  and  $k^{0r}$  were calculated using the following equations:<sup>15,16</sup>

$$k^0 = \varphi \sqrt{\frac{D_0 \pi \nu n F}{RT} \left( \frac{D_R}{D_O} \right)^\alpha} \quad (1)$$



**Scheme 2** Equivalent circuit of Randles.





$$\varphi = \frac{(-0.6288 + 0.0021 \cdot \Delta E)}{(1 - 0.0170 \cdot \Delta E)} \quad (2)$$

$$k^{o'} = \frac{(RT)}{(n^2 F^2 A C R_{ct})} \quad (3)$$

where  $D_O$  and  $D_R$  are the diffusion coefficients for ferricyanide ( $D_O$ ) and ferrocyanide ( $D_R$ ),  $\nu$  is the scan rate ( $V s^{-1}$ ),  $n$  is the number of electrons involved in the process,  $T$  is the temperature (K),  $F$  is the Faraday constant ( $mol^{-1}$ ),  $R$  is the universal gas constant ( $J K^{-1} mol^{-1}$ ) and  $\alpha$  is the dimensional transfer coefficient. However, the parameter  $\varphi$  is based on the potential difference between the anodic and cathodic peaks ( $\Delta E$ ).

The Randles-Sevcik equation (eqn (4)) was utilized to examine the electrochemical behavior of several electrodes:

$$I_p = (0.4463)nFAC\sqrt{\frac{nF\nu D_O}{RT}} \quad (4)$$

where  $I_p$  is the current peak,  $n$  is the number of electrons exchanged,  $F$  is the Faraday constant ( $mol^{-1}$ ),  $R$  is the universal gas constant ( $J K^{-1} mol^{-1}$ ),  $A$  is the area of the electrode surface ( $cm^2$ ),  $C$  is the analyte concentration ( $mol cm^{-3}$ ),  $D_O$  is the diffusion coefficient ( $cm^2 s^{-1}$ ), and  $\nu$  is the scan rate ( $mV s^{-1}$ ).

## Author contributions

Rocco Cancelliere: conceptualization, data curation, formal analysis, investigation, methodology, project administration, supervision, validation, visualization, writing – original draft, and writing – review and editing. Giuseppina Rea: methodology, investigation, formal analysis, supervision, writing – original draft, and writing – review and editing. Leonardo Severini: investigation and formal analysis. Luciana Cerri: methodology, investigation and data curation. Gabriella Leo: methodology, investigation, formal analysis, and data curation. Elisa Paialunga: visualization and data curation. Pietro Mantegazza: investigation. Claudia Mazzuca: investigation, formal analysis, and data curation. Laura Micheli: methodology, project administration, funding acquisition, resources, software, supervision, and writing – review and editing.

## Conflicts of interest

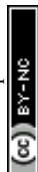
There are no conflicts to declare.

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## References

- P. Sarda, J. C. Hanan, J. G. Lawrence and M. Allahkarami, *J. Polym. Sci.*, 2022, **60**, 7–31.
- P. Benyathiar, P. Kumar, G. Carpenter, J. Brace and D. K. Mishra, *Polymers*, 2022, **14**, 2366.
- C. Ioakeimidis, K. N. Fotopoulou, H. K. Karapanagioti, M. Geraga, C. Zeri, E. Papathanassiou, F. Galgani and G. Papatheodorou, *Sci. Rep.*, 2016, **6**, 23501.
- K. Zhang, A. H. Hamidian, A. Tubić, Y. Zhang, J. K. H. Fang, C. Wu and P. K. S. Lam, *Environ. Pollut.*, 2021, **274**, 116554.
- M. Arhant, M. Le Gall, P.-Y. Le Gac and P. Davies, *Polym. Degrad. Stab.*, 2019, **161**, 175–182.
- F. Awaja and D. Pavel, *Eur. Polym. J.*, 2005, **41**, 1453–1477.
- P. Benyathiar, P. Kumar, G. Carpenter, J. Brace and D. K. Mishra, *Polymers*, 2022, **14**, 2366.
- K. Ragaert, L. Delva and K. Van Geem, *Waste Manage.*, 2017, **69**, 24–58.
- C. Dincer, R. Bruch, E. Costa-Rama, M. T. Fernández-Abedul, A. Merkoçi, A. Manz, G. A. Urban and F. Güder, *Adv. Mater.*, 2019, 1806739.
- J. Wang, *Analytical Electrochemistry*, Wiley, 1st edn, 2006.
- R. Cancelliere, A. D. Tinno, A. Cataldo, S. Bellucci and L. Micheli, *Biosensors*, 2021, **12**, 2.
- A. Di Tinno, R. Cancelliere, P. Mantegazza, A. Cataldo, A. Paddubskaya, L. Ferrigno, P. Kuzhir, S. Maksimenko, M. Shuba, A. Maffucci, S. Bellucci and L. Micheli, *Nanomaterials*, 2022, **12**, 1779.
- A. Ahamed, L. Ge, K. Zhao, A. Veksha, J. Bobacka and G. Lisak, *Chemosphere*, 2021, **278**, 130462.
- R. Cancelliere, M. Cianciaruso, K. Carbone and L. Micheli, *Chemosensors*, 2022, **10**, 344.
- J. Wang and S. Wang, *J. Cleaner Prod.*, 2019, **227**, 1002–1022.
- M. Chu, W. Tian, J. Zhao, M. Zou, Z. Lu, D. Zhang and J. Jiang, *Chemosphere*, 2022, **307**, 136024.
- F. J. Chacón, M. L. Cayuela, A. Roig and M. A. Sánchez-Monedero, *Rev. Environ. Sci. Biotechnol.*, 2017, **16**, 695–715.
- European Commission and Directorate-General for Communication, Circular economy action plan: for a cleaner and more competitive Europe, Publications Office of the European Union, 2020.
- R. Cancelliere, K. Carbone, M. Pagano, I. Cacciotti and L. Micheli, *Biosensors*, 2019, **9**, 139.
- K. C. Cole, J. Guèvremont, A. Ajji and M. M. Dumoulin, *Appl. Spectrosc.*, 1994, **48**, 1513–1521.
- S. K. Bahl, D. D. Cornell, F. J. Boerio and G. E. McGraw, *J. Polym. Sci., Polym. Lett. Ed.*, 1974, **12**, 13–19.
- F. J. Boerio, S. K. Bahl and G. E. McGraw, *J. Polym. Sci., Polym. Phys. Ed.*, 1976, **14**, 1029–1046.
- R. Cancelliere, D. Albano, B. Brugnoli, K. Buonasera, G. Leo, A. Margonelli and G. Rea, *Appl. Surf. Sci.*, 2021, **567**, 150791.
- L. Celio, M. Ottaviani, R. Cancelliere, A. Di Tinno, P. Panjan, A. M. Sesay and L. Micheli, *Front. Chem.*, 2021, **9**, 626630.



- 25 Z. Bahadoran, P. Mirmiran, A. Ghasemi, A. Kabir, F. Azizi and F. Hadaegh, *Nitric Oxide*, 2015, **47**, 65–76.
- 26 A. Ferlazzo, V. Bressi, C. Espro, D. Iannazzo, E. Piperopoulos and G. Neri, *J. Electroanal. Chem.*, 2023, **928**, 117071.
- 27 C. Matthews, F. Moran and A. K. Jaiswal, *J. Cleaner Prod.*, 2021, **283**, 125263.
- 28 M. Talbi, A. Al-Hamry, P. R. Teixeira, L. G. Paterno, M. B. Ali and O. Kanoun, *Chemosensors*, 2022, **10**, 40.
- 29 S. Palanisamy, B. Thirumalraj and S.-M. Chen, *J. Electroanal. Chem.*, 2016, **760**, 97–104.
- 30 A. Ferlazzo, V. Bressi, C. Espro, D. Iannazzo, E. Piperopoulos and G. Neri, *J. Electroanal. Chem.*, 2023, **928**, 117071.
- 31 S. Nasraoui, A. Al-Hamry, P. R. Teixeira, S. Ameer, L. G. Paterno, M. Ben Ali and O. Kanoun, *J. Electroanal. Chem.*, 2021, **880**, 114893.
- 32 R. Mo, X. Wang, Q. Yuan, X. Yan, T. Su, Y. Feng, L. Lv, C. Zhou, P. Hong, S. Sun, Z. Wang and C. Li, *Sensors*, 2018, **18**, 1986.
- 33 X. Li, N. Zou, Z. Wang, Y. Sun, H. Li, C. Gao, T. Wang and X. Wang, *Chem. Pap.*, 2020, **74**, 441–449.
- 34 S. Yallappa, M. Shivakumar, K. L. Nagashree, M. S. Dharmaprakash, A. Vinu and G. Hegde, *J. Electrochem. Soc.*, 2018, **165**, H614–H619.
- 35 B. R. Kozub, N. V. Rees and R. G. Compton, *Sens. Actuators, B*, 2010, **143**, 539–546.
- 36 O. Salhi, T. Ez-zine, L. Oularbi and M. El Rhazi, *Front. Chem.*, 2022, **10**, 870393.

