

Comparative study of ZnO and ZnFe<sub>2</sub>O<sub>4</sub> micro and nanoparticle-based screen-printed electrodes in pH sensing

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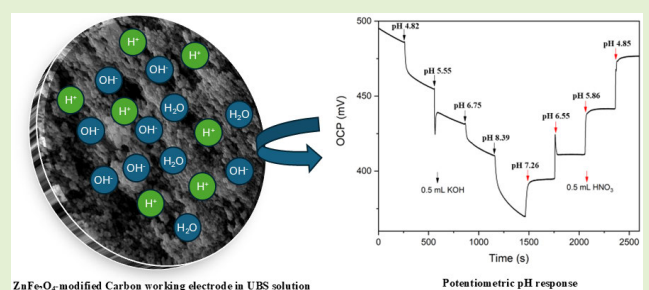
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# Comparative Study of ZnO and ZnFe<sub>2</sub>O<sub>4</sub> Microparticle and Nanoparticle-Based Screen-Printed Electrodes in pH Sensing

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**Abstract**—This work presents the application of zinc oxide (ZnO) and zinc ferrite (ZnFe<sub>2</sub>O<sub>4</sub>) for electrochemical pH sensing. ZnO and ZnFe<sub>2</sub>O<sub>4</sub> are synthesized by auto-combustion synthesis method. Field emission scanning electron microscopic (FESEM) images revealed that ZnO particles have pyramid- and spherical-shaped morphology with micrometer dimensions, while ZnFe<sub>2</sub>O<sub>4</sub> particles have spherical shape at the nanoscale. The surface-modified screen-printed electrodes with ZnO and ZnFe<sub>2</sub>O<sub>4</sub> particles are initially characterized by the ferri/ferrocyanide redox couple. Significant improvement in sensitivity (bare carbon:  $6.3 \pm 0.4 \mu\text{A}/\text{mM}$ , ZnO:  $8.5 \pm 0.3 \mu\text{A}/\text{mM}$ , ZnFe<sub>2</sub>O<sub>4</sub>:  $8.9 \pm 0.5 \mu\text{A}/\text{mM}$ ) and rate constant (bare carbon:  $10 \pm 1 \text{ms}^{-1}$ , ZnO:  $46 \pm 4 \text{ms}^{-1}$ , ZnFe<sub>2</sub>O<sub>4</sub>:  $42 \pm 3 \text{ms}^{-1}$ ) is observed with the surface-modified sensors. Chronopotentiometric pH response of the sensors showed hysteresis behavior with pH loop. No interference effects are observed, and the pH sensitivity of the bare carbon sensor ( $23.9 \pm 1.4 \text{mV}/\text{pH}$ ) is increased by the introduction of ZnO ( $38.1 \pm 1.3 \text{mV}/\text{pH}$ ) and ZnFe<sub>2</sub>O<sub>4</sub> ( $37.2 \pm 1.1 \text{mV}/\text{pH}$ ) particles. Stability of the pH response is discussed, and ways for its improvement are proposed.

**Index Terms**—Microparticles, nanoparticles, potentiometric pH sensor, sensor stability, zinc ferrite (ZnFe<sub>2</sub>O<sub>4</sub>), zinc oxide (ZnO).



## I. INTRODUCTION

ELECTROCHEMICAL pH sensing is valued for its high sensitivity, rapid response, and adaptability to a wide range of environments, making it a versatile solution in fields such as clinical diagnostics [1], environmental monitoring [2], water quality assessment [3], [4], and industrial process control [5]. These sensors operate based on the electrochemical interaction between a pH-sensitive material and hydrogen ions (H<sup>+</sup>) in a solution, allowing for real-time, in situ measurements of acidity, neutrality, or alkalinity of a solution [6].

Electrochemical pH sensors typically work through potentiometric, amperometric, or conductometric approaches. The most common type is the potentiometric pH sensor, which measures the potential difference between a pH-sensitive

working electrode (WE) (such as glass electrodes or metal oxides) and a reference electrode (RE) (typically Ag/AgCl) immersed in the solution. The potential generated at the WE is proportional to the activity of H<sup>+</sup> ions, which are related to the solution's pH value through Nernst equation [6], [7], [8].

Metal oxide nanomaterials have emerged as highly effective materials for electrochemical pH sensing due to their excellent electrochemical properties, stability, and sensitivity to hydrogen ions (H<sup>+</sup>) in solutions. Their unique structural, electrical, and chemical characteristics make them suitable for a wide range of pH sensing applications [1], [6], [9], [10]. In environmental applications, titanium dioxide (TiO<sub>2</sub>) [11] and zinc oxide (ZnO) [3], [12] nanomaterials have been employed in pH sensors to monitor water quality due to their wide bandgap [13], [14]. These sensors can detect shifts in pH caused by contaminants in rivers, lakes, or oceans, providing crucial data for water management. Nickel oxide (NiO) [15] and copper oxide (CuO) [6], [9], [16] nanoparticles are also commonly used in soil pH sensors due to their robust sensing capabilities in harsh environments.

In clinical diagnostics, monitoring pH is vital for tracking physiological parameters such as blood acidity (pH of blood)

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or the pH of interstitial fluids, which are critical in conditions like acidosis or alkalosis [1]. Iridium oxide ( $\text{IrO}_x$ ) [8], [17] and ruthenium oxide ( $\text{RuO}_2$ ) [1], [6], [10], [18], [19] are used to develop pH sensors with super Nernstian behavior that can be integrated into biomedical devices, including wearable and implantable sensors. These materials exhibit fast response times and stability under physiological conditions. Tin oxide ( $\text{SnO}_2$ ) [6], [18], [20] nanomaterials are also explored for implantable pH sensors due to their nontoxic nature and high sensitivity to pH changes, which are essential features for detecting pH changes in tissue environments during medical treatments. Tungsten oxide ( $\text{WO}_3$ ) is also used to develop pH sensors for clinical diagnostics [21], [22].

Spinel oxides are another class of multitransition metal oxides such as zinc ferrite ( $\text{ZnFe}_2\text{O}_4$ ) and nickel ferrite ( $\text{NiFe}_2\text{O}_4$ ). The spinels possess a crystal structure of the form  $\text{AB}_2\text{O}_4$  (A—metal cation with “+2” oxidation state and B—metal cation with “+3” oxidation state) with excellent chemical stability and electrical neutrality [23]. Several spinel oxide nanomaterials are used as sensing materials in electrochemical sensing applications for drug detection [24]. An improvement of the electrochemical sensing response is possible due to the tunability of their crystal structure. Their composition can be easily modified maintaining the chemical stability of the structure [25], [26], [27], [28], [29]. The spinel can be normal, inverse, or mixed with different cations occupying different sites in the crystal structure and leading to different electrochemical performance due to the change in their semiconducting nature either from  $p$  to  $n$  or from  $n$ - to  $p$ -type [25], [26]. The magnetic nature (ferromagnetic and antiferromagnetic behavior leading to electron scattering effects and charge trapping) of the spinel materials also plays an important role in the conductivity of spinel oxides [30] which in turn can affect the electrochemical sensing performance. This class of spinel nanomaterials is not explored as pH sensing electrode materials in the literature as far as our knowledge is concerned. Therefore, in this work, we investigate the capability of  $\text{ZnFe}_2\text{O}_4$  normal spinel material to act as an electrode sensing material of pH.

$\text{ZnO}$  and  $\text{ZnFe}_2\text{O}_4$  particles are synthesized and characterized using different techniques. Commercial screen-printed electrodes (SPEs) are used as an electrochemical sensing cell and the surface of the WE is modified with the newly synthesized materials. The electrochemical behavior of the new sensors is first assessed using the well-known ferri/ferrocyanide probe molecule, then their performance as pH sensors is assessed. Different performance characteristics of pH sensors such as hysteresis, sensor stability, sensitivity, and interference effects are presented and discussed.

## II. MATERIALS AND METHODS

### A. Chemicals

Zinc nitrate ( $\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ ), iron nitrate ( $\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ ),  $\text{C}(\text{NH}_2)_2\text{O}$  (Urea), Butanol ( $\text{C}_4\text{H}_{10}\text{O}$ ), potassium hexacyanoferrate(II) ( $\text{K}_4\text{Fe}(\text{CN})_6$ ), potassium hexacyanoferrate(III) ( $\text{K}_3\text{Fe}(\text{CN})_6$ ), acetic acid ( $\text{CH}_3\text{COOH}$ ), potassium nitrate ( $\text{KNO}_3$ ), boric acid ( $\text{H}_3\text{BO}_3$ ), phosphoric acid ( $\text{H}_3\text{PO}_4$ ), potassium hydroxide (KOH), and nitric acid

( $\text{HNO}_3$ ) are purchased from Sigma Aldrich and used without any further purification.

### B. Material Synthesis

A simple and cost-effective auto-combustion synthesis is used to synthesize nanomaterials based on our previous work [25]. Synthesis of  $\text{ZnFe}_2\text{O}_4$  nanoparticles is reported in our previous work [25], [31].  $\text{ZnO}$  is synthesized by using only  $\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$  following the same process as  $\text{ZnFe}_2\text{O}_4$  synthesis.

### C. Preparation of Universal Buffer Solution (UBS)

UBS covering the pH range from 2 to 12 is prepared to characterize electrochemical sensors. A measure of 0.3 mL (0.01 M) of  $\text{CH}_3\text{COOH}$ , 5.05 g (0.1 M) of  $\text{KNO}_3$ , 0.31 g (0.01 M) of  $\text{H}_3\text{BO}_3$ , and 0.58 mL (0.01 M) of  $\text{H}_3\text{PO}_4$  are mixed with 0.5 L of deionized (DI) water in a glass container. The mixture is kept in an ultrasonic bath for 5 min to dissolve all the chemicals and obtain a final homogeneous solution.

### D. Apparatus

Field emission scanning electron microscopic (FESEM) images are collected utilizing a Zeis SupraTM 50 (Oberkochen, Germany) to examine the nanomaterial's morphology. The crystal structure of the materials is determined by X-ray powder diffraction (XRD) using a Panalytical Empyrean diffractometer (Malvern Panalytical, Malvern, U.K.). The XRD analysis of the powder samples is performed at room temperature using a Bragg–Brentano setup with  $\text{Cu K}\alpha$  ( $\lambda = 1.5418 \text{ \AA}$ ) radiation at 40 kV and a current of 40 mA. The analysis is carried out using a time step of 60 s and a step size of  $0.013^\circ/\text{s}$ , covering a  $2\theta$  range from  $10^\circ$  to  $70^\circ$ . Palmsens potentiostat with PSTrace 5.9 is used to perform cyclic voltammetry (CV) in solutions containing ferri/ferrocyanide. Metrohm Autolab with Nova 2.1 is used for electrochemical impedance spectroscopic (EIS) studies and chronopotentiometric pH sensing.

### E. Electrochemical Sensors

Commercial screen-printed carbon WE (area  $0.12 \text{ cm}^2$ ) and carbon counter electrode (CE) on a ceramic substrate along with an external double junction  $\text{Ag}/\text{AgCl}$  RE are used as electrodes for ferri/ferrocyanide sensing. The same method as reported in the literature [25] is adopted to prepare the nanomaterial solutions and the surface of the carbon WE is modified using the drop-casting technique as reported previously [25], [31], [32].

### F. Electrochemical Measurements

CV is used to characterize the sensors performance for the detection of ferri/ferrocyanide redox couple with a potential window ranging from  $-0.4$  to  $0.8 \text{ V}$  at a scan rate of  $100 \text{ mV/s}$  in phosphate buffer (PB) at pH 7. Scan rate analysis is performed by changing the scan rate from  $25 \text{ mV/s}$  to  $125 \text{ mV/s}$  in steps of  $25 \text{ mV/s}$ . Calibration of the sensors is conducted by recording cyclic voltammograms in the concentration range

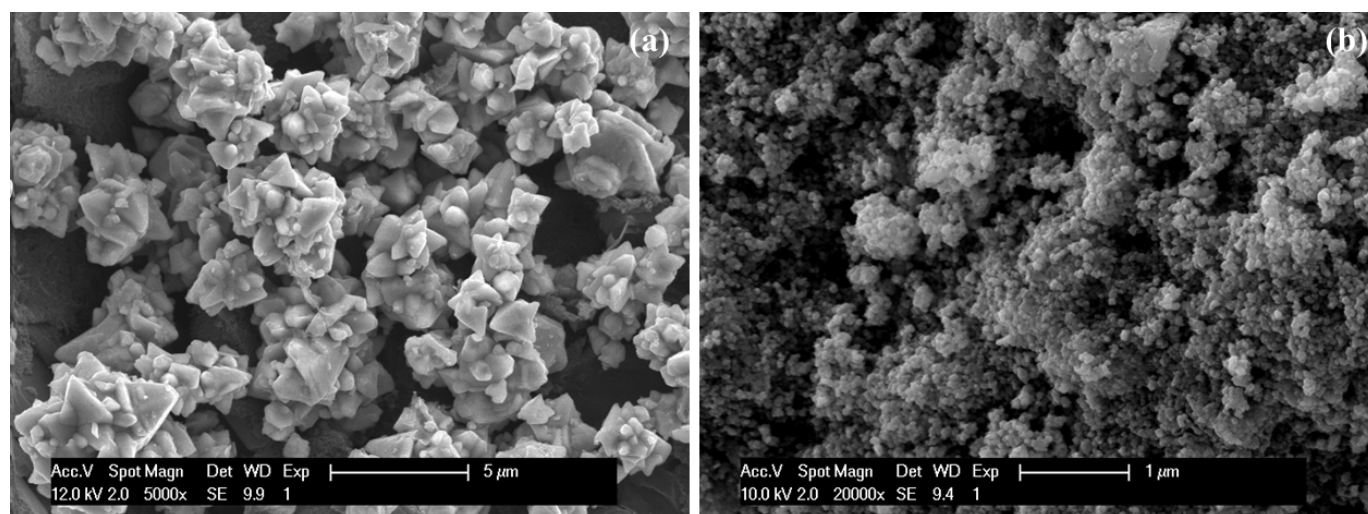


Fig. 1. FESEM images of (a) ZnO with 5k $\times$  and (b) ZnFe<sub>2</sub>O<sub>4</sub> with 20k $\times$  magnification, respectively.

164 from 1 to 8 mM of ferri/ferrocyanide redox couple. EIS  
 165 is performed in the frequency from 0.1 to 100 kHz with  
 166 10 points/decade at an amplitude of 5 mV. Chronopotentiometric  
 167 experiments are conducted for pH initially in the full  
 168 loop of 2-12-2 and in the physiologically relevant pH range  
 169 of 5-8.5. The RE and CE probes of the Autolab are coupled  
 170 together, and the solution is continuously stirred at 500 r/min.  
 171 Open-circuit potential (OCP) is recorded with time and the  
 172 average of the last 60 s of the response of OCP before any  
 173 pH change in solution is plotted against the pH value for the  
 174 calibration. The slope of the plot is reported as the sensitivity  
 175 of the sensors. Interference measurements are conducted using  
 176 30 mM KCl and 5 mM NaCl to verify the robustness of the  
 177 sensor toward interfering ions.

### 178 III. RESULTS AND DISCUSSION

#### 179 A. Material Characterization

180 Fig. 1 shows the FESEM images of ZnO and ZnFe<sub>2</sub>O<sub>4</sub>  
 181 particles-modified carbon surface. ZnO particles have a pyramidal  
 182 shape at microscale, while some smaller particles exhibit  
 183 a different morphology like spherical one. ZnFe<sub>2</sub>O<sub>4</sub> particles  
 184 show spherical-shaped morphology. A certain amount of  
 185 aggregation is observed in both materials that are possibly  
 186 due to the adopted synthesis method [25], [33] as it provides  
 187 a weak control over the particle growth kinetics.

188 The XRD patterns of ZnFe<sub>2</sub>O<sub>4</sub> and ZnO are shown in  
 189 Fig. 2. For ZnFe<sub>2</sub>O<sub>4</sub>, the experimental pattern fully matches  
 190 with the normal spinel Franklinite (ZnFe<sub>2</sub>O<sub>4</sub>, reference code  
 191 01-089-4926). Some low-intensity peaks are noticed in the  
 192  $2\theta$  range between 30° and 37°: these peaks are compatible  
 193 with the most intense peaks of ZnO (reference code 01-080-0075)  
 194 and iron oxide (Fe<sub>2</sub>O<sub>3</sub>, reference code 01-085-0987) in accordance  
 195 with previous studies [25], [26]. For ZnO, the XRD pattern  
 196 neatly matches with the ZnO one (reference code 01-080-0075).  
 197 The low-intensity peak at low angles is attributed to graphite,  
 198 which might be present as an impurity due to the graphite reactor  
 199 used for the material synthesis. Crystallite size of the materials  
 200 is calculated using Scherrer's method [25], [26]. ZnFe<sub>2</sub>O<sub>4</sub> has  
 201 an average crystallite size of 35.2  $\pm$  3.3 nm, while ZnO has an  
 202 average

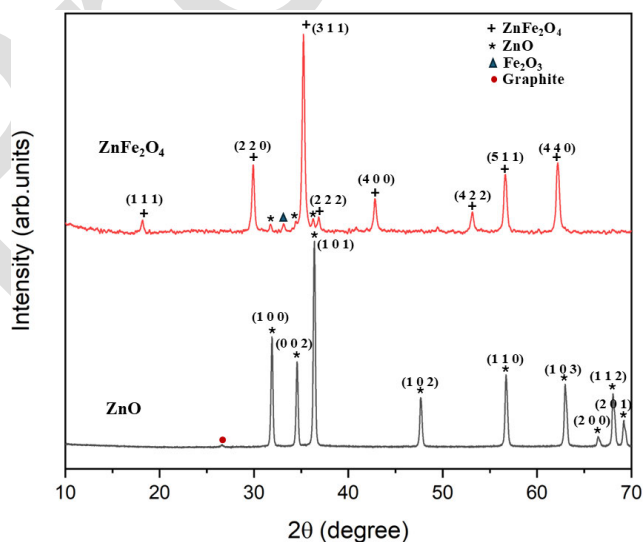


Fig. 2. XRD patterns of ZnFe<sub>2</sub>O<sub>4</sub> and ZnO particles synthesized by auto-combustion.

203 crystallite size of 45.7  $\pm$  1.3 nm. In the case of ZnO-particles,  
 204 particles are formed by abnormal asymmetric growth leading  
 205 to the micrometer dimensions of particles, as evidenced in  
 206 Fig. 1.

#### 207 B. Electrochemical Characterization

208 Fig. 3 shows the cyclic voltammograms of the bare,  
 209 ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO sensors characterized in 4 mM  
 210 ferri/ferrocyanide redox couple in 0.1 M PB at pH 7 at  
 211 a scan rate of 100 mV/s. The introduction of nanostructured/  
 212 microstructured materials has enhanced the performance  
 213 of screen-printed carbon electrodes. The oxidation peak current  
 214 ( $I_{pa}$ ) is increased [ $\sim$ 14  $\mu$ A:  $I_{pa}$  = 44.1  $\pm$  0.6  $\mu$ A  
 215 (ZnFe<sub>2</sub>O<sub>4</sub>), 44.3  $\pm$  0.4  $\mu$ A (ZnO)] and the anodic peak potential  
 216 (oxidation peak potential:  $E_{pa}$ ) is decreased [ $\sim$ 70 mV:  
 217  $E_{pa}$  = 413  $\pm$  3 (ZnFe<sub>2</sub>O<sub>4</sub>), 407  $\pm$  3 mV (ZnO)] compared to  
 218 bare carbon sensor ( $I_{pa}$  = 30.6  $\pm$  1.4  $\mu$ A,  $E_{pa}$  = 483  $\pm$  7 mV).

219 Cyclic voltammograms obtained at various scan rates are  
 220 depicted in Fig. S1 for bare carbon, ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO  
 221 sensors. The anodic ( $I_{pa}$ ) and cathodic peak currents ( $I_{pc}$ )

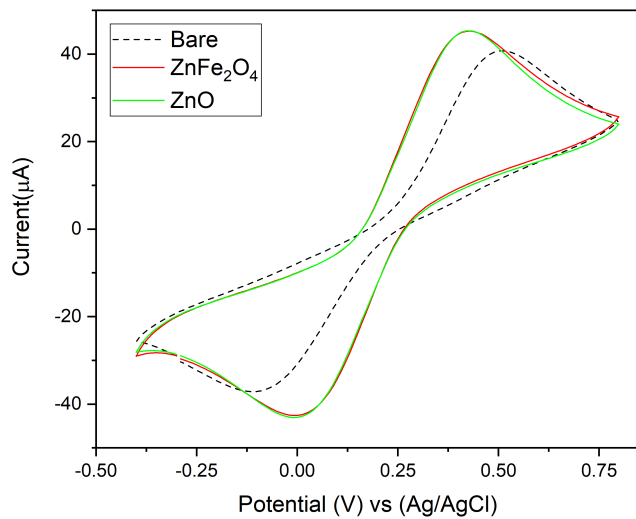


Fig. 3. Cyclic voltammograms of different sensors at a scan rate of 100 mV/s. (Electrolytic solution: 4 mM ferri/ferrocyanide in PB (0.1 M) at pH 7).

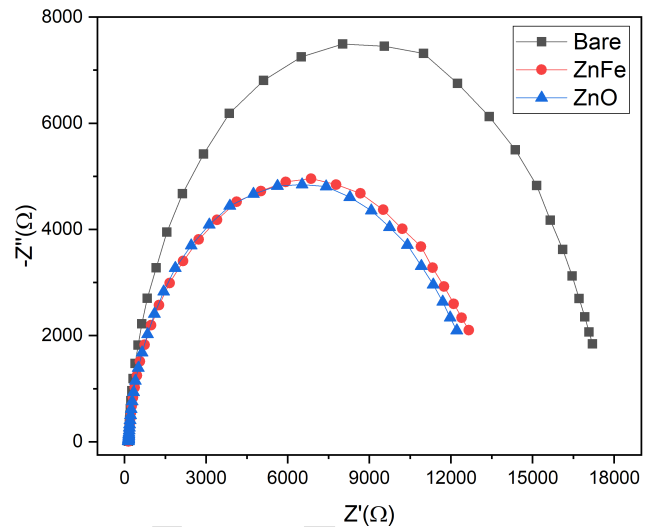


Fig. 4. Nyquist plots of different sensors in 4 mM ferri/ferrocyanide in PB (0.1 M) at pH 7. Frequency range is 0.5–100 kHz with 10 points per decade at an amplitude of 5 mV.

TABLE I

ELECTROACTIVE SURFACE AREA AND INTERFACE IMPEDANCES IN CHARACTERIZING FERRI/FERROCYANIDE REDOX COUPLE ON BARE CARBON AND SURFACE-MODIFIED SENSORS

| Sensor                           | $A_{RS}$ (cm <sup>2</sup> ) | $R_s$ (Ω) | $R_p$ (kΩ) |
|----------------------------------|-----------------------------|-----------|------------|
| Bare                             | $0.27 \pm 0.05$             | 166       | 17.2       |
| ZnFe <sub>2</sub> O <sub>4</sub> | $0.49 \pm 0.03$             | 146       | 13.4       |
| ZnO                              | $0.49 \pm 0.02$             | 139       | 11.8       |

increased with the increase in scan rate and varied linearly with the square root of the scan rate ( $\sqrt{\nu}$ ) in the range 25–125 mV/s, as shown in Fig. S2. The peak currents linear regression equations with their regression coefficients, confirming the high level of linearity, are listed in Table S1. Absolute redox current peak values are plotted in Fig. S3 to show the differences between the oxidation and reduction peak currents. The redox peak current ratios with scan rate are reported in Table S2, and the ratio  $I_{pa}/I_{pc}$  is  $\approx 1$ . Electroactive surface area ( $A_{RS}$ ) of the sensors is calculated using Randles–Sevcik equation [26] and reported in Table I. It is observed that the electroactive surface area of the sensors is increased due to the presence of nanoparticle/microparticle of ZnFe<sub>2</sub>O<sub>4</sub>/ZnO on the carbon surface compared to the bare carbon sensor. Both ZnFe<sub>2</sub>O<sub>4</sub> and ZnO-based sensors have a similar electroactive surface area indicating the same loading of the nanoparticle/microparticle on the carbon surface that participates in the electrochemical reaction with ferri/ferrocyanide redox couple.

The peak-to-peak separation ( $\Delta E_p = E_{pa} - E_{pc}$ , where  $E_{pc}$  is the cathodic peak potential or reduction peak potential) lowered by 150 mV compared to bare carbon sensor due to the presence of either ZnO or ZnFe<sub>2</sub>O<sub>4</sub> particles. The ZnO-based sensor showed the lowest  $\Delta E_p$  of  $387 \pm 9$  mV, while the ZnFe<sub>2</sub>O<sub>4</sub> sensor and bare carbon sensor exhibited a  $\Delta E_p$  of  $397 \pm 7$  and  $547 \pm 9$  mV, respectively. As the electron transfer kinetics are directly related to  $\Delta E_p$ , the

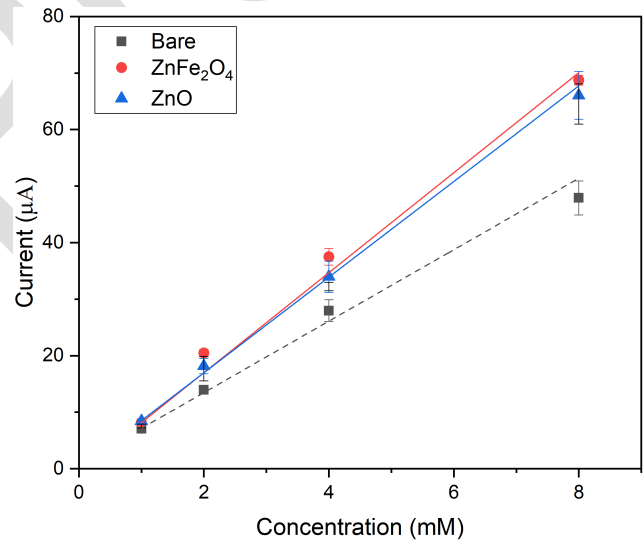


Fig. 5. Calibration of sensors for concentrations of ferri/ferrocyanide redox couple in PB (0.1 M) up to 8 mM.

lower  $\Delta E_p$  indicates faster electrochemical reactions with high reversibility. In Figs. S4 and S5, the redox peak positions  $E_{pa}$  and  $E_{pc}$ , and  $\Delta E_p$  are plotted and a linear variation against  $\ln(\nu)$  is observed. The linear regression equations with their coefficients are reported in Tables S3 and S4, respectively. Half-potential ( $E_{1/2} = (E_{pa} + E_{pc})/2$ ) is shown in Fig. S6 and has almost a constant relation with  $\ln(\nu)$  is observed. The redox peak current ratio,  $\Delta E_p$ , and  $E_{1/2}$  suggest that the interface is a quasi-reversible electrochemical interface from Randles–Sevcik theory [34], [35]. Following the equations from the Laviron model [25], [36], [37], the electron transfer rate coefficient “ $\alpha$ ” and kinetic rate constant “ $k_s$ ” using  $\Delta E_p$  ( $\nu = 100$  mV/s) and  $n = 1$ , are calculated and presented in Table II.

EIS is performed to study the impedance properties of the electrochemical interface between 4 mM ferri/ferrocyanide redox couple and different sensors. Nyquist plots of bare, ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO-based sensors are reported in Fig. 4. The Nyquist profiles of the sensors show only semicircles

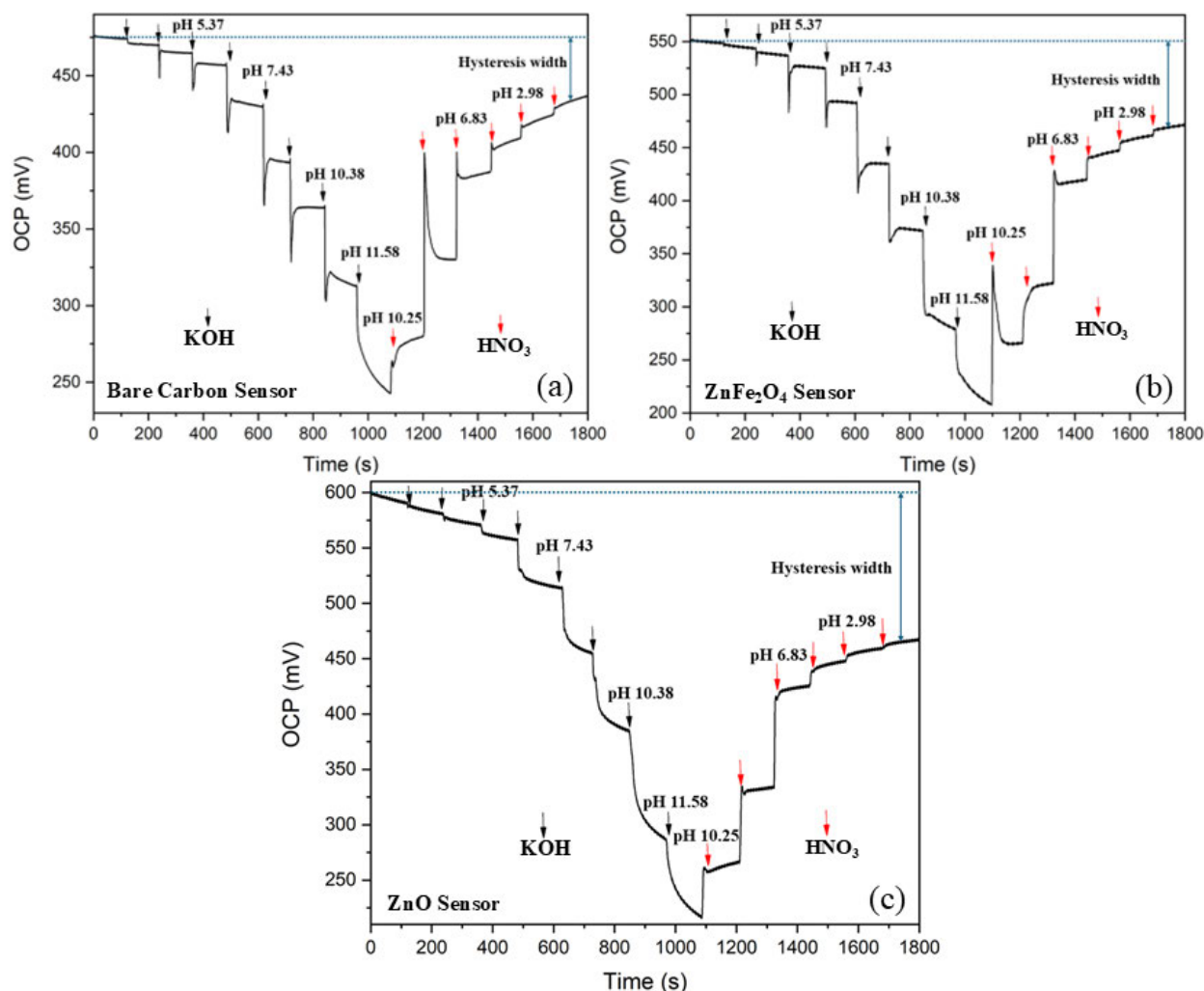


Fig. 6. Chronopotentiometric response of (a) bare, (b) ZnFe<sub>2</sub>O<sub>4</sub>, and (c) ZnO pH sensors showing the hysteresis effect in pH loop 2-12-2.

TABLE II  
ELECTROCHEMICAL PARAMETERS IN CHARACTERIZING  
FERRI/FERROCYANIDE REDOX COUPLE ON BARE  
CARBON AND SURFACE-MODIFIED SENSORS

| Sensor                           | $\alpha$      | $\Delta E_p$ (mV) | $k$ (ms <sup>-1</sup> ) |
|----------------------------------|---------------|-------------------|-------------------------|
| Bare                             | 0.47 ± 0.01   | 547 ± 9           | 9.97 ± 0.78             |
| ZnFe <sub>2</sub> O <sub>4</sub> | 0.49 ± 0.01   | 397 ± 7           | 41.8 ± 2.6              |
| ZnO                              | 0.496 ± 0.002 | 387 ± 9           | 46 ± 4                  |

268 associated with charge transfer kinetic limitations. Therefore,  
269 Randles equivalent circuit model with a series resistance  
270 ( $R_s$ ), charge transfer resistance ( $R_p$ ), and a constant phase  
271 element (CPE) are used to fit the obtained Nyquist plots  
272 of bare, ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO-modified sensors without any  
273 Warburg impedance (diffusion-controlled) as shown in Fig. S7.  
274 From the circuit fitting, series (solution) resistance ( $R_s$ ) and  
275 charge transfer resistance ( $R_p$ ) of the sensors are extracted  
276 and reported in Table I. The charge transfer resistance of the  
277 ZnFe<sub>2</sub>O<sub>4</sub> and ZnO-based sensors is significantly lower than  
278 the bare carbon sensor supporting the higher electron transfer  
279 rate constant. ZnO-based sensor showed 1.6 k $\Omega$  lower charge

280 transfer resistance with similar CPE parameters compared to  
281 ZnFe<sub>2</sub>O<sub>4</sub>-based sensor supporting its higher electron transfer  
282 rate (higher “ $k$ ”).

283 The sensors are calibrated by taking the average response  
284 of three sensors of each type, and the calibration plots are  
285 shown in Fig. 5. Sensitivity is computed with reference to the  
286 oxidation peaks of the ferri/ferrocyanide redox couple in the  
287 concentration range 1–8 mM. The bare carbon sensor has a  
288 sensitivity of 6.3 ± 0.4  $\mu$ A/mM, while the ZnFe<sub>2</sub>O<sub>4</sub>-based  
289 sensor and the ZnO-based sensor have sensitivities of 8.9 ±  
290 0.5 and 8.5 ± 0.3  $\mu$ A/mM, respectively. Though there are  
291 significant differences in crystallite size and morphology, the  
292 materials have shown similar electrochemical performance ( $k$   
293 and sensitivity) which might be due to their similar electroac-  
294 tive surface area and charge transfer resistance.

295 Fig. 6 shows the chronopotentiometric response of bare  
296 carbon, ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO-based sensors in the full pH  
297 loop of 2-12-2 (with 15 different pH values reported in the  
298 supporting information Table S5). The response at each pH is  
299 measured for two minutes (total time span of 30 min), and  
300 it is observed that ZnO has the higher initial OCP (lowest  
301 pH values). Moreover, there is huge shift in OCP after pH  
302 10 for all sensors. In the reverse pH from basic to acidic  
303 region, the OCP is similar in the surface-modified sensors

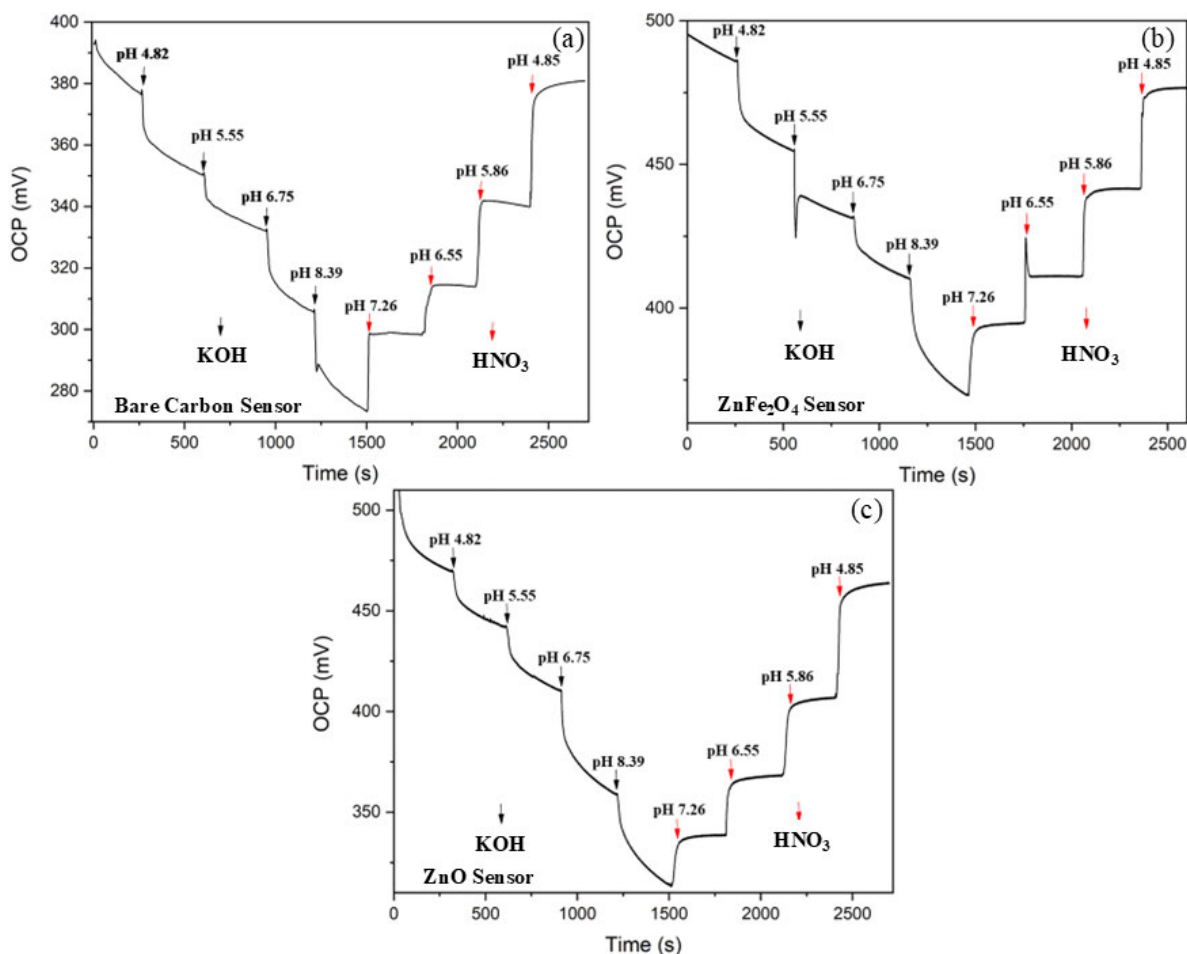


Fig. 7. Chronopotentiometric response of (a) bare, (b)  $\text{ZnFe}_2\text{O}_4$ , and (c)  $\text{ZnO}$  pH sensors with reduced hysteresis effect in pH loop 4.5–8.5–4.5.

304 compared to the initial OCP from acidic to basic region. 305 These differences in response leads to a phenomenon called 306 “hysteresis” in pH sensors especially in the case of metal 307 oxide-based pH sensors [6]. The potential difference from 308 initial response to the final response in the complete pH loop 309 is called the “hysteresis width” as indicated in Fig. 6. The 310  $\text{ZnO}$ -based sensor has the highest hysteresis width of 133 mV, 311 bare carbon sensor has the lowest width of 39 mV, and the 312  $\text{ZnFe}_2\text{O}_4$ -based sensor has a hysteresis width of 81 mV. The 313 hysteresis width defines the functionality of a pH sensor and 314 the pH range in which the sensor can be utilized. The lower 315 the hysteresis width, the better the performance of the sensor 316 in that pH range. As reported before, the hysteresis behavior 317 depends on the pH looping (pH range in which the sensor is 318 operated) [38]. Since our main interest is in the physiological 319 pH range, the sensors are characterized in the pH loop 320 4.5–8.5–4.5 (with nine different pH values with a total time 321 span of 45 min) as shown in Fig. 7. It is clearly observed that 322 the hysteresis behavior is reduced to 3, 18, and 44 mV for 323 bare carbon,  $\text{ZnFe}_2\text{O}_4$ , and  $\text{ZnO}$ -based sensors, respectively. 324 This is significantly lower compared to the previous pH loop 325 proving that the hysteresis behavior is pH loop-dependent.

326 Stability of the pH sensors is another important performance 327 factor. As is clear from Figs. 6 and 7, the potentiometric 328 response is not reaching a steady-state value when the pH is 329 changing from lower to higher values by the addition of  $\text{KOH}$

330 solution. To understand the sensors’ stability, the response 331 is recorded at different pH values for longer times (10 and 332 20 min after each change in pH value) as presented in Fig. 8. 333 A drift over time is still observed without reaching any steady- 334 state value. The response starts to stabilize immediately after 335 the addition of  $\text{HNO}_3$  solution into the solution under test. 336 To understand this effect, it is important to consider how 337 a pH sensor works. When the sensor is immersed into the 338 test solution, the hydroxide ( $\text{OH}^-$ ) ions get attracted by the 339 electrode surface due to the dissociative adsorption of water. 340 This adsorption leads to a potential difference between the 341 WE and the RE. Due to the addition of more  $\text{KOH}$  solution 342 into the test UBS solution, the sensor’s surface is enriched 343 with more  $\text{OH}^-$  ions, leading to a difference in the potential 344 between WE and RE. When the  $\text{HNO}_3$  solution is added, 345 the response starts to stabilize immediately as we introduce 346  $\text{H}^+$  ions. This is due to the faster diffusion of  $\text{H}^+$  ions (and 347 so faster protonation of the electrode surface) compared to 348 the process of interaction of  $\text{OH}^-$  ions with the electrode 349 surface [6]. To improve the stability of the sensors, instead 350 of using  $\text{KOH}$  as the base stock solution, bicarbonates such as 351  $\text{KHCO}_3$  or  $\text{NaHCO}_3$  can be employed. This might improve the 352 stability of the potential response as well as the pH sensitivity 353 of the sensors. When the pH of the solution under investigation 354 is controlled with  $\text{KOH}$  and  $\text{HNO}_3$ , the sensors are calibrated 355 in the pH range mentioned above to compare the OCP. When

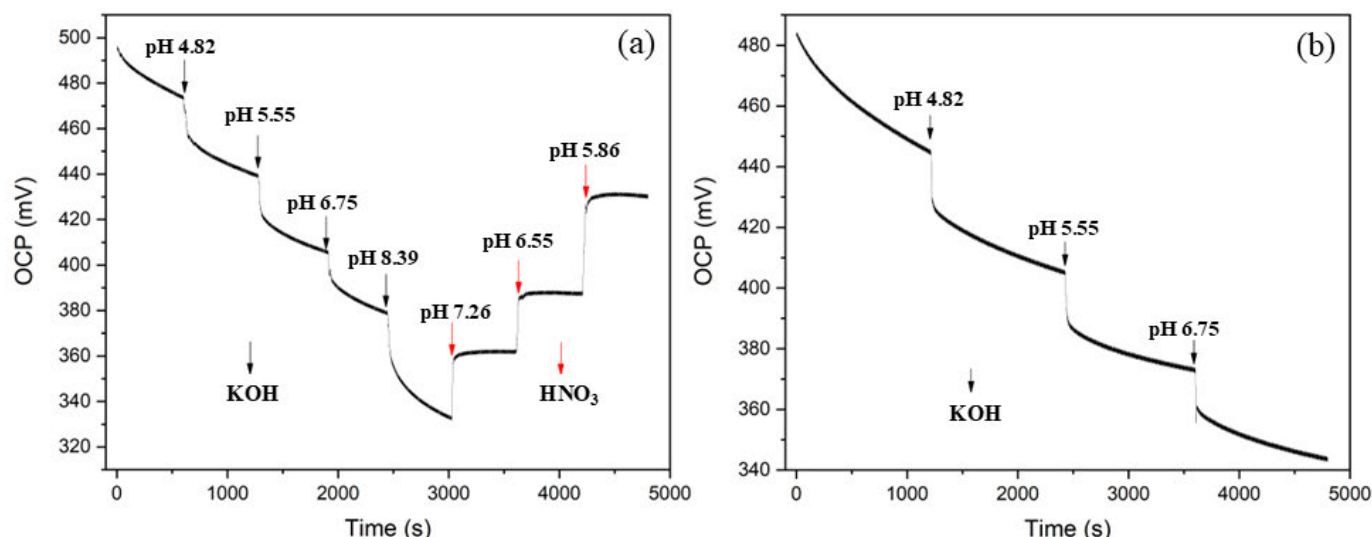


Fig. 8. Stability tests: chronopotentiometric response of ZnFe<sub>2</sub>O<sub>4</sub> pH sensor with longer stabilization times [(a) 10 and (b) 20 minutes of waiting time in between subsequent pH changes] after each pH change in the range between 4.5 and 8.5.

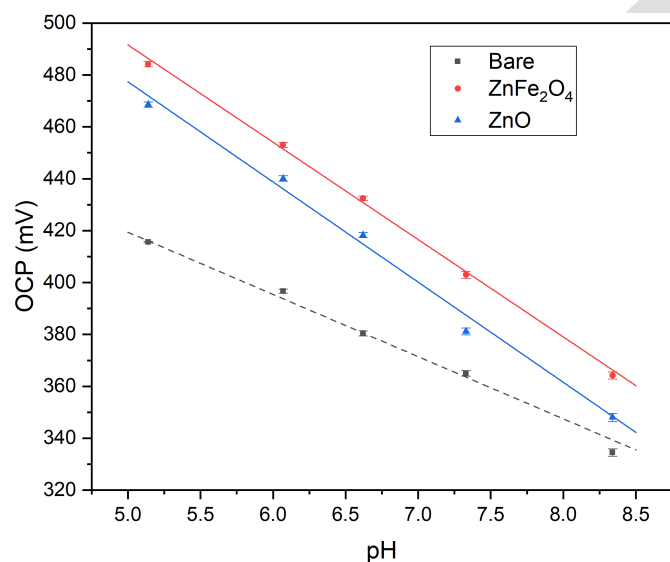


Fig. 9. Calibration of bare carbon, ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO pH sensors in the pH between 5 and 8.5.

HNO<sub>3</sub> is used after KOH in continuous measurements, the sensitivity of the sensors is higher, according to the calibration plots shown in Fig. S8. However, this is unreliable because the OCP response is unstable in the first case when KOH is used, which affects the OCP levels in the second case when HNO<sub>3</sub> is used for pH control. The response time of the sensors is difficult to investigate due to unstable response in the case of KOH; therefore, the response time is reported for the second case when HNO<sub>3</sub> is used for pH control. The bare, ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO-based sensors showed response times ( $T_{95}$ ) of 5, 80, and 95 s, respectively. These times are just indicative but not reliable due to unstable continuous OCP measurements with KOH for pH control.

To assess the performance of the electrodes as pH sensors, three freshly prepared electrodes of each type are taken, and the potential response is measured for three minutes as reported in Fig. S9. The average potential response in the last

60 s before changing the pH is considered to calibrate the sensors in the pH range between 5 and 8.5. pH calibration of the sensors is presented in Fig. 9. Bare carbon sensor, ZnFe<sub>2</sub>O<sub>4</sub>, and ZnO-based sensors have shown sensitivities of  $23.9 \pm 1.4$ ,  $37.3 \pm 1.1$ , and  $38.1 \pm 1.3$  mV/pH, respectively. The presence of either ZnFe<sub>2</sub>O<sub>4</sub> or ZnO significantly improved the pH sensing performance of bare carbon sensor, resulting in an increase of sensitivity of around 60%. pH sensing response is usually analyzed with the Nernst equation [6] as written below

$$E = E_0 - \frac{2.303RT}{nF} \text{pH} \quad (1)$$

where  $E$  is the Nernst potential,  $E_0$  is the standard surface potential,  $R$  is the universal gas constant,  $T$  is the temperature, and " $n$ " is the number of electrons transferred in the reaction. At atmospheric conditions with  $T = 25$  °C for one electron process (i.e.,  $n = 1$ ), the slope of the plot ( $E$  versus pH) is 59 mV/pH. This response is usually termed as Nernstian response for pH. Though for our sensors the response is not Nernstian, we observed a significant increase with respect to bare sensor, proving the potential of ZnFe<sub>2</sub>O<sub>4</sub> or ZnO in pH sensing applications.

The ZnO particles have a pyramid shape where the sides and long edges of the particles are electroactive, whereas ZnFe<sub>2</sub>O<sub>4</sub> has (close to) uniform spherical-shaped particles. Another difference is in the crystal structure which is hexagonal for ZnO microparticles, while ZnFe<sub>2</sub>O<sub>4</sub> particles exhibit a complex normal spinel where the metal centers of Fe and Zn play an important role in the electrochemical activity of the sensors [24], [39]. ZnO is a wide bandgap material (either p- or n-type semiconductor) with a bandgap around 3.2 eV [40], whereas ZnFe<sub>2</sub>O<sub>4</sub> (n-type semiconductor) has a smaller bandgap of about 2.5 eV [25], [26]. Even though they have these differences, we have observed that a comparable electroactive surface area (from ferri/ferrocyanide sensing), charge transfer resistance [kinetic limited reactions without diffusion limitations from Nyquist plots (see Fig. 4)], electron transfer rate constant (" $k$ ") and sensitivity in pH sensing for

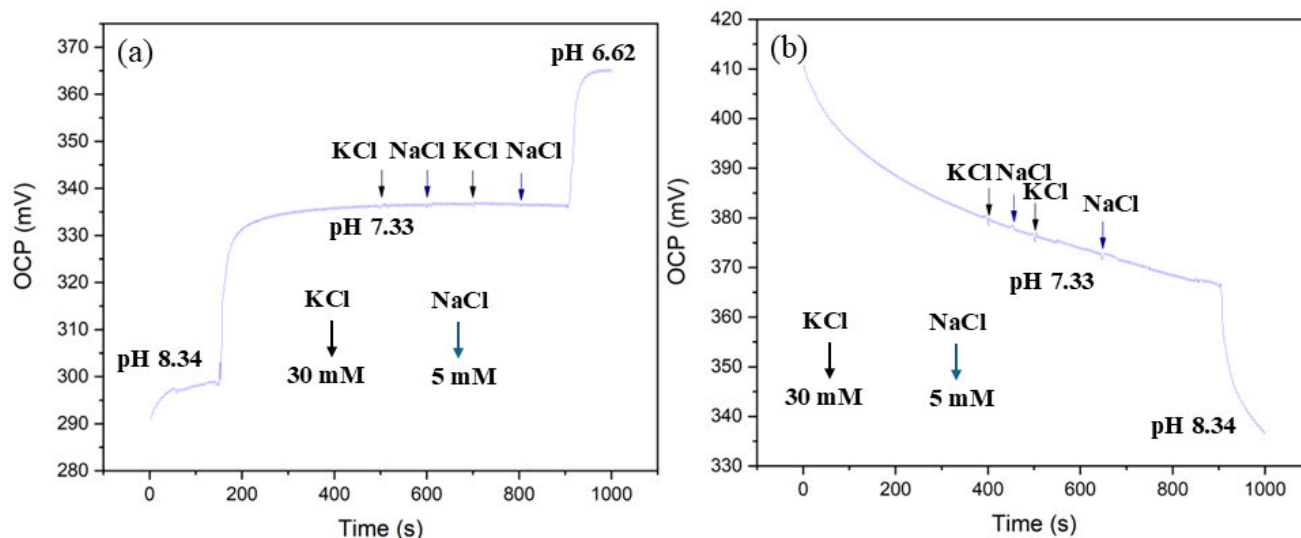


Fig. 10. Interference tests for ZnFe<sub>2</sub>O<sub>4</sub> pH sensor by adding 30 mM KCl and 5 mM NaCl in solution at neutral pH. The tests are performed by changing the pH from (a) basic to neutral pH and (b) neutral to basic pH.

our ZnFe<sub>2</sub>O<sub>4</sub> and ZnO -based sensors. ZnO-based sensor has shown higher hysteresis width compared to the other two types of sensors in this work. For the selection of the electrode type, it is crucial to consider eventual metal oxide dissolution effects. The dissolution depends on the metal solubility in the solution and the loading of the nanoparticle/microparticle on the surface [41]. ZnO can dissociate into Zn(II) and Zn(OH)<sup>+</sup> ions in aqueous solutions with alkaline or neutral pH conditions [40], [41], [42], [43]. On the other hand, there is no evidence of dissolution of ZnFe<sub>2</sub>O<sub>4</sub> reported so far in the literature. Thus, interference tests are performed with the ZnFe<sub>2</sub>O<sub>4</sub>-based sensor by using relevant interferent species, namely KCl and NaCl, at the physiological pH of 7.33 (close to the blood pH of 7.4). The measurement graph is shown in Fig. 10. During the measurement, 30 mM KCl is added followed by the successive addition of 5 mM NaCl. There is no significant effect on the potential response of the sensor proving an excellent pH sensing selectivity to interfering ions. In this work, we used the simple auto-combustion synthesis method to produce ZnO and ZnFe<sub>2</sub>O<sub>4</sub> particles based on our previous work [25], [26]. The electrochemical pH sensor response of the SPEs is improved by modification with the synthesized micromaterial/nanomaterial. The performance of the sensors can be further improved to have more control over the particle growth kinetics and the material structural properties. For instance, using appropriately selected surfactants during the synthesis can reduce particle aggregation. This would lead to an improvement in particle activity when incorporated on the electrode surface.

#### IV. CONCLUSION

In this work, ZnO and ZnFe<sub>2</sub>O<sub>4</sub> particles are successfully produced using a simple and cost-effective auto-combustion synthesis method. The materials are characterized by FESEM and XRD to understand their morphological and structural features. Commercial SPEs were modified with the developed particles and studied as new electrochemical sensors. First, the sensors are characterized using ferri/ferrocyanide redox

couple as a probe molecule *via* CV. Significant enhancement in the performance of bare carbon sensor is achieved in terms of sensitivity and rate constant using ZnO or ZnFe<sub>2</sub>O<sub>4</sub> particles. Successively, the electrodes' performance as pH sensor is studied by means of chronopotentiometric measurements. The hysteresis behavior of the sensors is improved by using the physiological pH loop (4.5–8.5–4.5). We observed that the stability of the pH sensors is affected by the addition of OH<sup>-</sup> ions in the test solution. Because of this, we propose the use of alternative stock solutions for pH change to obtain a stable steady-state signal upon the increase in the pH. Overall, the pH sensitivity of bare carbon sensor is increased (>55%) by using two different surface-modified sensors (ZnO or ZnFe<sub>2</sub>O<sub>4</sub>). Due to potential dissolution issues of the ZnO particles when in water-based solution and reduced hysteresis in pH sensing, interference studies are conducted at ZnFe<sub>2</sub>O<sub>4</sub>-modified electrodes and shown that the sensor is silent to potentially relevant interfering ions.

In this work, we choose to study ZnFe<sub>2</sub>O<sub>4</sub> due to its chemical stability, easy processing, low-cost synthesis, high surface area at nanoscale, easily tunable structure, and its n-type semiconducting and ferromagnetic nature with normal spinel structure. For continuous monitoring, no dissolution is expected due to its highly stable closely packed face-centered cubic structure which is crucial in pH sensing. Additionally, by incorporating surfactants in the synthesis process (hydrothermal or co-precipitation or sol-gel method), agglomeration can be avoided leveraging nanoscale properties which can lead to improved performance in the future work.

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materials and microfabrication techniques for implantable, bioresorbable electrochemical sensors.

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