


Cite this: *RSC Adv.*, 2024, 14, 24661

PDA@ α -ZrP/SiO₂ incorporated phosphate coating with enhanced corrosion resistance and friction resistance

Yubin Tian,^a Wei Li,^a Qiaoxin Yang,^a Yuan Gong,^a Wenfei Yue,^a Yujing Ou,^a Chunlei Li^{*a} and Xinxin Sheng^{ID *bc}

In this study, α -ZrP and SiO₂ composite nanomaterials were used to as phosphating accelerators. Experiments show that 2D nanosheets and 0D nanoparticles modified by PDA (PDA@ α -ZrP/SiO₂) play a synergistic role in effectively increasing the number of phosphate crystals and refining the crystal's size, thereby forming dense and uniform phosphate coatings. The friction resistance and corrosion resistance of phosphate coatings are simultaneously enhanced. Especially when the PDA@ α -ZrP/SiO₂ addition amount is 0.55 g L⁻¹, the coating porosity of phosphate coating drops from 64.24% to 4.38%. The friction resistance coefficient drops from 0.32 to 0.02 and the polarization resistance increased from 1381 Ω cm² to 20 520 Ω cm².

Received 29th May 2024
Accepted 29th July 2024

DOI: 10.1039/d4ra03958k

rsc.li/rsc-advances

1. Introduction

Traditional phosphating technology is widely used in metal surface treatment industries, including the machinery industry, automobile industry, aviation industry and other fields.^{1–4} The main function of the phosphate chemical conversion coatings is to protect the base metal materials from corrosion damage. Phosphate coatings can also play a role in lubrication and friction reduction in metal cold plastic processing technology. Although traditional chemical phosphating process technology is constantly being optimized and adjusted, it cannot fundamentally solve the problems of phosphating coatings in industrial applications. Traditional high-temperature phosphating has changed to low-temperature phosphating due to problems such as acid mist and high energy consumption. However, in low-temperature phosphating, due to the low free acidity, temperature and H⁺ concentration, cathodic polarization is easily caused, making the phosphate coating forming reactions difficult to proceed. Therefore, a certain phosphating accelerator needs to be introduced.^{5,6} Traditional phosphating accelerators are mainly oxidation accelerators, including nitrite, nitrate, chlorate, *etc.*^{7,8} On the one hand, the accelerator solves the cathodic polarization process; on the other hand, it oxidizes ferrous ions into ferric ions, which effectively prevents the

enrichment of ferrous ions. However, these traditional phosphating accelerators, which are currently the most widely used and effective, are carcinogenic, harmful to human health and environment. This is not in line with the concepts and trends of green production and sustainable development.^{9,10} Moreover, for low-temperature phosphating, improper oxidant concentration results in the generation of a large amount of phosphate sediment and contamination of the bath liquid. Phosphate sediment contains high phosphorus content and strong acidity, and is clearly classified as toxic waste. This is an urgent problem that needs to be solved in the phosphating treatment industry. An effective solution is to seek effectively and environmentally friendly green phosphate accelerators to replace traditional oxidizing accelerators.^{11–13}

The nanomaterials have received widespread research attention in phosphating technology, due to the large specific surface area and excellent physical and chemical properties. Some research works have confirmed that nanomaterials, including 2D and 0D nanomaterials, can promote the formation of phosphate coatings and have the potential to replace traditional phosphate accelerators.^{14,15} M. Tamilselvi *et al.* introduced SiO₂ into the zinc phosphating baths and confirmed that SiO₂ has the ability to reduce the activation energy of the phosphating process and increase the nucleation sites.¹⁶ Other works have also confirmed that nanoparticles such as nano ZnO, TiO₂ and ZrO₂ have the ability to promote phosphate coating formation and give phosphate coatings better performances including better corrosion resistance, coating weight, and greater substrate coverage *etc.*^{17–19} In addition to 0D nanoparticles, 2D nanosheet materials have also been proven to be used to new phosphate accelerators. Xie *et al.* first explored the possibility of using graphene oxide (GO) as the new

^aCollege of Petrochemical Technology, Lanzhou University of Technology, Lanzhou 730050, PR China. E-mail: licl@lut.edu.cn

^bDepartment of Polymeric Materials and Engineering, School of Materials and Energy, Guangdong University of Technology, Guangzhou, 510006, PR China. E-mail: xinxin.sheng@gdut.edu.cn; cexxsheng@gmail.com

^cGuangdong Provincial Key Laboratory of Functional Soft Condensed Matter, Guangdong University of Technology, Guangzhou 510006, PR China



environmentally friendly phosphate accelerator.¹⁴ Experimental results confirmed that in the initial stage of phosphate coating formation, the 2D GO sheets were conducive to capturing metal ions and thereby promoting the nucleation of phosphate crystals. When the number of phosphate crystals increases, the grain size is refined, which is beneficial to the formation of phosphate coatings with small porosity. At the same time, it is crucial for blocking corrosive media. Boron nitride nanosheets (BNs) has also been shown to increase the nucleation sites of phosphate crystals and thereby improve surface coverage of metal substrates.²⁰ In our previous works, dopamine-modified MXene (MXene@PDA) played an active role in the chemical phosphating process.⁹ In another work, stearic acid (SA)-modified MXene not only promoted phosphating, but also imparted hydrophobic properties to phosphate coatings, further enhancing the corrosion resistance.⁶ Other 2D nanosheet materials such as molybdenum disulfide (MoS₂), blank phosphorus, α -zirconium phosphate (α -ZrP), *etc.*^{21,22} have also been proven to have great potential as phosphate accelerators. Nanomaterials are expected to break the key bottlenecks faced by traditional phosphating.

α -ZrP, a new multifunctional 2D layered materials, is a strong solid acid with excellent ion exchange, selective adsorption and catalytic properties.²³ α -ZrP exchanges with metal ions and has great dispersion stability in acid solution, which has a positive effect on promoting the process of phosphating reactions and increasing the phosphating rate.⁴ Polydopamine (PDA) is formed by the self-polymerization of catechol natural substance dopamine. It contains a large number of phenolic hydroxyl and amino functional groups and has strong adhesion.²⁴ Modifying nanomaterials with PDA enhances the dispersion of nanomaterials. In addition, PDA can also adsorb metal ions such as Fe³⁺, Zn²⁺, Cu²⁺, *etc.* through surface functional groups, which also has positive effect on accelerating the phosphating process.^{25,26}

Herein, we modified α -ZrP and SiO₂ with PDA (PDA@ α -ZrP/SiO₂) and explored the potential of 2D/0D nanocomposites as phosphating accelerator. 2D/0D nanocomposites were designed to effectively combine the advantages of two separate components.^{27,28} The physicochemical properties of phosphate crystals were assessed through a series of different characterization techniques. The formation process of phosphate coatings was monitored by scanning electron microscopy (SEM). The corrosion resistance and friction resistance of the phosphate coatings were also evaluated. The results demonstrated that PDA@ α -ZrP/SiO₂ nanocomposite phosphate coatings presented better performances, which also imply that 2D/0D nanocomposites play a synergistic effect to achieve the more positive promotion effect on phosphating.

2. Materials and methods

2.1 Materials

All chemical reagents used were of analytical reagent. Zn(H₂-PO₄)₂·2H₂O, Zn(NO₃)₂·6H₂O and NaCl were purchased from Sinopharm Chemical Reagent Co., Ltd.

Tris, SiO₂ (20 ± 5 nm) and PDA were provided from Shanghai Macklin Biochemical Technology Co., Ltd. α -ZrP was produced by Fujian Runsen New Materials Co., Ltd. H₃PO₄ and NaOH were acquired from Guangzhou Chemical Reagent Factory. The Q235 steel samples (50 mm × 50 mm × 1 mm) with C = 0.22%, Si = 0.35%, S = 0.05%, P = 0.05%, Mn = 1.40% and Fe = balance (wt%) elemental composition were supplied from Shenzhen Oudifu Co., Ltd.

2.2 Fabrication of PDA@ α -ZrP/SiO₂ nanocomposite

First, 0.20 g Tris was dissolved in 200 mL water to prepare a Tris buffer solution, and then 0.20 g α -ZrP and 0.20 g SiO₂ were added to the Tris buffer solution for sonication.

Secondly, 0.20 g dopamine was directly added to aforementioned Tris buffer and stirred magnetically at room temperature for 24 h. Finally, after centrifugation and washing with deionized water, the PDA@ α -ZrP/SiO₂ was freeze-dried for subsequent use.

2.3 Fabrication of chemical conversion phosphate coatings

The metal substrates were pre-treated before phosphating. SiC-600 grits papers were used to polish and remove rust, and then 10% NaOH aqueous solution was used to degrease the metal substrate at 50 °C for 10 min. The formula of phosphating bath was based on our previous work.²⁹ Different concentrations of PDA@ α -ZrP/SiO₂ (0.25 g L⁻¹, 0.55 g L⁻¹, 0.85 g L⁻¹) were introduced into the phosphating bath and ultrasonicated at room temperature for 30 min to obtain the uniform phosphating solution. The pretreated Q235 substrates were placed in the prepared phosphating bath and phosphating at 50 °C for 30 min. The phosphate samples were rinsed with deionized water and dried with cold air for subsequent tests.

2.4 Corrosion property test

The corrosion resistance of phosphate coatings was evaluated by EIS and potentiodynamic polarization tests, performed on the CHI-760E electrochemical workstation (Chenhua, China). A standard three-electrode system was used, with the saturated Ag/AgCl electrode, graphite electrode and phosphate coating used as the reference electrode, counter electrode and working electrode respectively. The samples were tested with 3.5 wt% NaCl solution as the electrolyte. Before EIS test, the open circuit potential (OCP) was performed for 120 s to obtain a stable OCP. And then EIS was performed at the OCP under an alternation current with an amplitude of 10 mV in the frequency range from 100 kHz to 10 mHz. The data obtained from EIS test were analyzed using ZSimpWin software (EChem Software, Q5 Ann Arbor, MI, USA) for equivalent circuit fitting. The potentiodynamic polarization curve test was performed at a scan rate of 1 mV s⁻¹ from the cathode to anode direction (from OCP – 300 mV to OCP + 300 mV). In addition, the corrosion resistance of phosphate coatings was also evaluated through the copper sulfate drop test. All tests were performed at least three times to ensure data reproducibility.



2.5 Tribological test

The tribological properties of phosphate coatings were assessed utilizing a block-on-ring wear tester (MRH-3, Jinan Yihua Tribology Testing Technology Co., Ltd, China).

The tribological tests were performed for 30 min at 25 °C and 40–70% relative humidity. The rotational speed is 100 rpm and the load is 100 N for the tribological tests. The friction resistance tests were all dry friction. Each experiment was performed three times to ensure the accuracy of experimental data.

2.6 Characterizations

The microscopic morphologies of PDA@ α -ZrP/SiO₂ and phosphate crystals were characterized by the scanning electron microscope (SEM, Zeiss Ultra plus, Germany). Fourier transform infrared (FT-IR) and UV-vis were employed to analyze the chemical structure of 2D/0D nanocomposites. X-ray diffraction (XRD, Bruker D8 ADVANCE, Germany) was performed using a CuK radiation source to analyze the phase composition of nanocomposite and phosphate coatings. The phosphate coating thickness was measured using the coating thickness instrument (BGD 542/1 thickness meter, China). The phosphate coating weight test method refers to our previous work.⁴

3. Results and discussion

3.1 Characterization of PDA@ α -ZrP/SiO₂ nanocomposite

FT-IR was performed to characterize the chemical composition of SiO₂, α -ZrP and PDA@ α -ZrP/SiO₂, and the results were shown in Fig. 1a. As to the curve of SiO₂, the strong and broad absorption peak at the 1118 cm⁻¹ position is attributed to the asymmetric stretching vibration absorption peak of Si–O–Si of SiO₂. The peak around 800 cm⁻¹ is the symmetric stretching vibration of Si–O–Si.³⁰ The peak at 1612 cm⁻¹ is the bending vibration peak of O–H. The character peak at 3432 cm⁻¹ could be assigned to the stretching vibration of –OH. The characteristic bands of α -ZrP are seen at 1045 cm⁻¹, which is attributed to the stretching vibration of P–O.³¹ And the peaks at 1619 cm⁻¹ and around 3417 cm⁻¹ are the symmetric stretching vibration peak of O–H bonds of the water molecules. In the PDA@ α -ZrP/SiO₂ spectrum, the peaks located at 1608 cm⁻¹ and 1492 cm⁻¹ are due to the C=C stretching vibration of the benzene ring of PDA. Around 1207 cm⁻¹ is the peak of C–O–H. In addition, the characteristic peaks of SiO₂ and α -ZrP coexist, which also confirms that PDA effectively modified and connected SiO₂ and α -ZrP.

In order to further verify that PDA successfully modified the SiO₂ and α -ZrP nanocomposites, XRD was used to detect

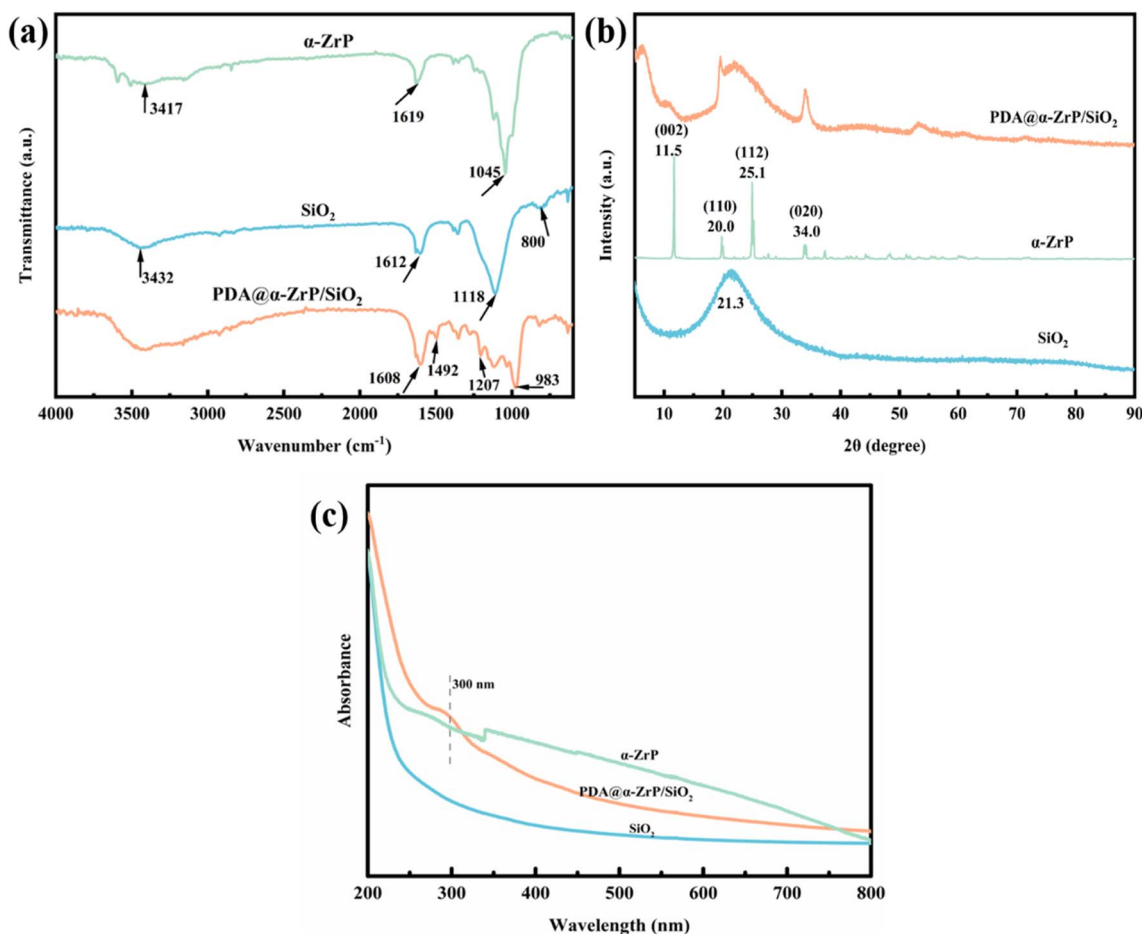


Fig. 1 FT-IR spectra (a), XRD spectra (b) and UV-vis spectra (c) for SiO₂, α -ZrP and PDA@ α -ZrP/SiO₂.

changes in phase composition (Fig. 1b). For SiO_2 , a broad and strong peak around 21.3° degree can be seen, which corresponds to the amorphous phase of SiO_2 . For $\alpha\text{-ZrP}$, $\alpha\text{-ZrP}$ with obvious diffraction peaks has high crystallinity. The peaks at 12° , 20° , 25° and 35° respectively correspond to the (002), (110), (112) and (020) crystal planes of $\alpha\text{-ZrP}$.³¹ When PDA was polymerized on the surface of SiO_2 and $\alpha\text{-ZrP}$, both the amorphous peak of SiO_2 and the crystalline peak of $\alpha\text{-ZrP}$ can be vaguely found in $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$. It confirms that the modification of PDA effectively combined the 2D $\alpha\text{-ZrP}$ and 0D SiO_2 .

From the UV-vis result of $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$ (Fig. 1c), it can be found that a peak attributed to PDA appears at 300 nm, which further proves that the modification of PDA was successful.

The morphologies of SiO_2 , $\alpha\text{-ZrP}$ and $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$ were observed with TEM and SEM, as shown in Fig. 2. Fig. 2a shows the TEM image of SiO_2 , and the size of SiO_2 particles used is 20 ± 5 nm. Fig. 2b shows the lamellar $\alpha\text{-ZrP}$. And Fig. 2c is $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$ nanocomposite. It can be clearly seen that SiO_2 and $\alpha\text{-ZrP}$ can be combined together through PDA modification. The surface roughness of the 2D/0D nanocomposite significantly increases due to the polymerization of dopamine on the surface of $\alpha\text{-ZrP}$ and SiO_2 . Elemental analysis on $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$ was further conducted using EDS-mapping (Fig. 2d), which confirmed that there are Zr, P, Si, P, O and N elements. It also confirms the successful preparation of 2D/0D nanocomposites.

3.2 Characterization of phosphate coatings

The growth process and morphology of phosphate crystals were observed through SEM. Fig. 3a1–d1 shows the growth state of phosphate crystals on the surface of metal substrate when phosphate coatings were formed for 1 min. It can be found that at 1 min, only a few phosphate crystals appears of the blank

phosphate coating on the surface of metal substrate. On the contrary, after incorporation $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$, a lot of phosphate crystals appears, which fully confirms that the 2D/0D composite nanomaterials can promote phosphating. When the phosphate coatings were formed for 30 min as shown in Fig. 3a2–d2, the phosphate crystals size of blank phosphate coating is larger, while the phosphate grains of nanocomposite phosphate coatings are obviously refined. In addition, the pure phosphate crystals have a tendency to grow vertically upwards from the substrate. In contrast, this upward growth tendency of phosphate crystals is inhibited when the promoter was introduced. Moreover, the nanocomposite phosphate coatings are more uniform and dense, which is extremely crucial for blocking corrosive media.

XRD analysis on the phase composition of phosphate coatings was further conducted, and the results are presented in Fig. 4. It is obvious that phosphate coatings comprise two crystal forms, $\text{Zn}_3(\text{PO}_4)_2 \cdot 4\text{H}_2\text{O}$ (hopeite, JCPD file # 37-0465) and $\text{Zn}_2\text{Fe}(\text{PO}_4)_2 \cdot 4\text{H}_2\text{O}$ (phosphophyllite, JCPD file # 29-1427).³²

In addition, we tested the coating thickness and coating weight (Fig. 5). The experimental results found that the blank phosphate coating has the largest coating thickness and the smallest coating weight. The incorporation of 2D/0D nanomaterials reduced the coating thickness but increased the weight of the phosphate coating. The statement appears paradoxical at first glance, yet the results have consistently been verified. The underlying explanation for these findings resides in the mechanism by which the integration of the nanocomposite fosters a heightened number of initial nucleation points for phosphate crystal formation. As more phosphate crystals grew, they would come into contact with each other in a shorter period of time and would restrict each other's continued growth. It would maintain a uniform coating

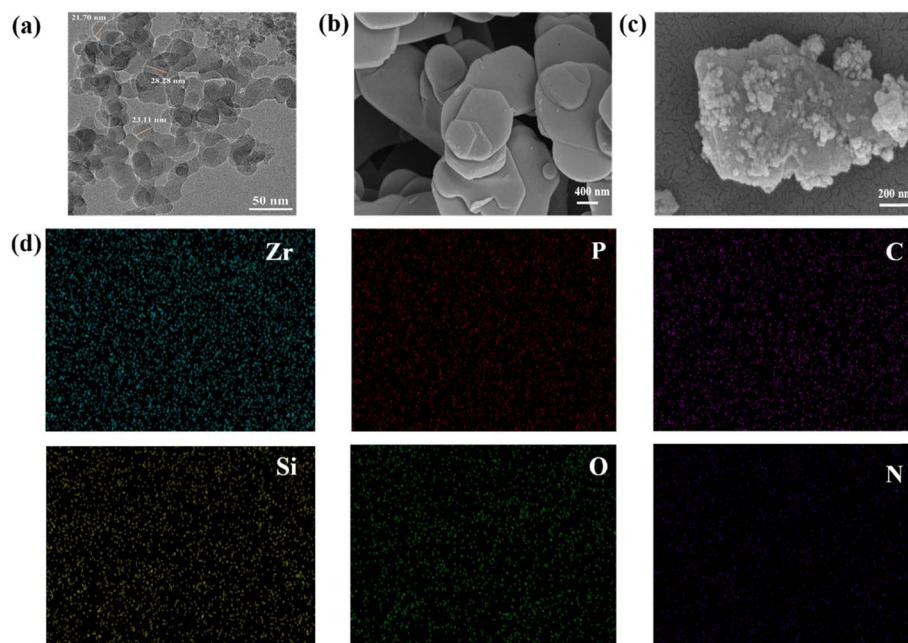


Fig. 2 TEM image of SiO_2 (a), SEM images of $\alpha\text{-ZrP}$ (b), $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$ (c) and EDS mapping of $\text{PDA}@ \alpha\text{-ZrP}/\text{SiO}_2$ (d).



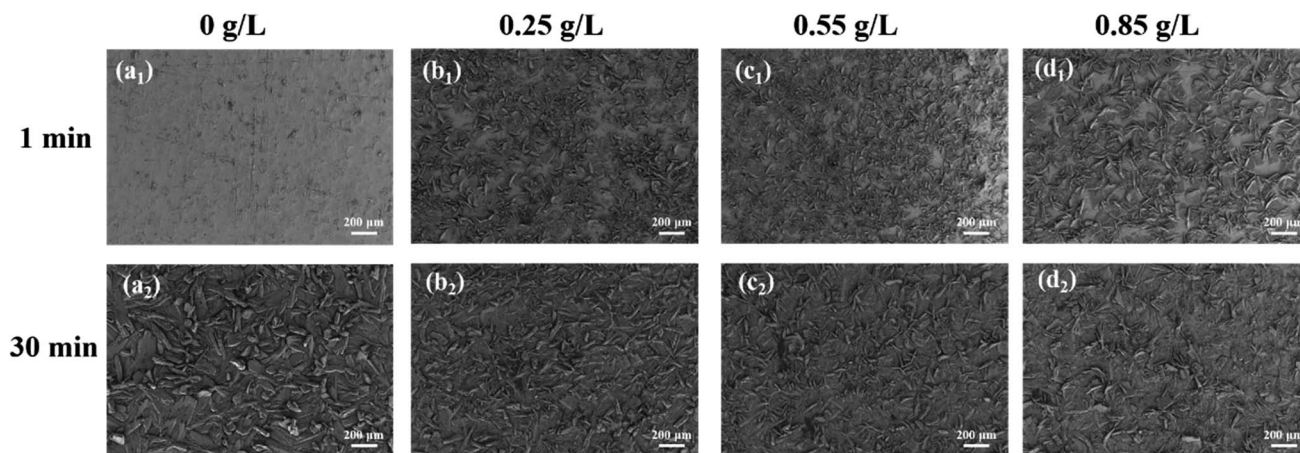


Fig. 3 SEM images of phosphate coatings with different PDA@α-ZrP/SiO₂ concentration as a function of time.

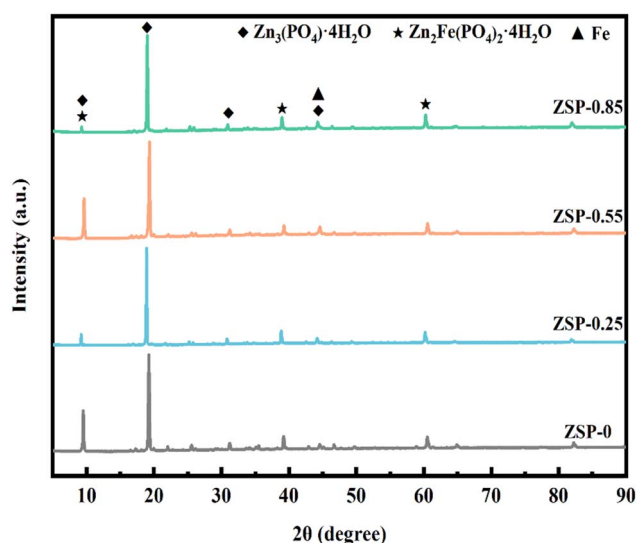


Fig. 4 XRD results of phosphate coatings.

thickness and limit the upward growth of the phosphate crystals. On the contrary, it can be found from SEM that the size of phosphate crystals is generally larger in the blank phosphate coating, and the growth of phosphate crystals was not restricted. Therefore, the upward growth orientation of phosphate crystals of blank phosphate coating resulted in a greater coating thickness. The difference in coating weight is that the nanocomposite phosphate coatings are denser and have the higher coverage of metal substrate, so they have a relatively higher coating weight.

3.3 Tribological properties of phosphate coatings

In cold plastic processing, the phosphate coatings need to have great friction resistance when used as lubricating coatings. The friction resistance of the 2D/0D nanocomposite phosphate coatings was evaluated, and Fig. 6 illustrates the coefficient of friction for phosphate coatings after the dry friction test conducted under a load of 100 N and at a speed of 100 revolutions per minute. Due to the special high-dispersion microporous structure, certain hardness, shock absorption and other

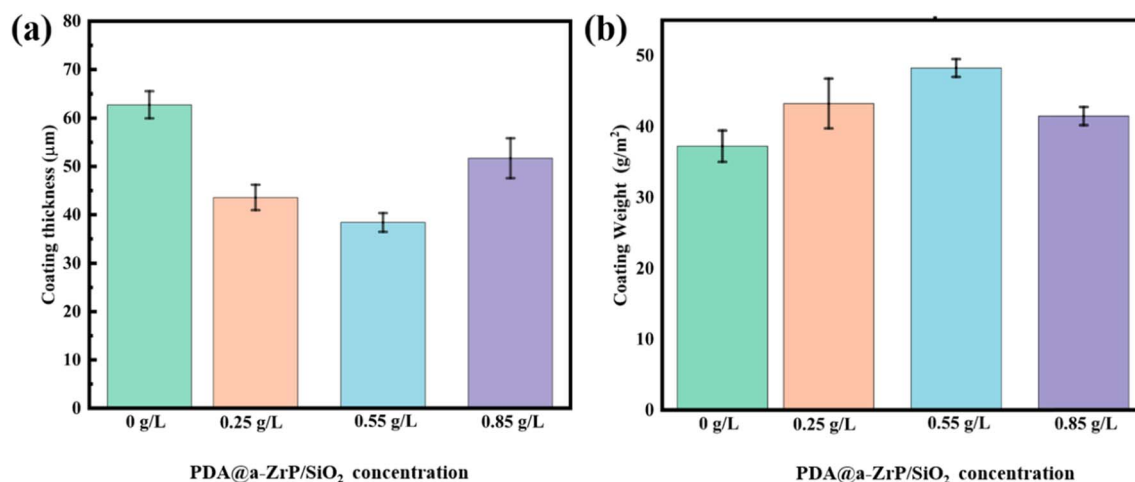


Fig. 5 Phosphate coatings thickness (a) and weight (b) with different PDA@α-ZrP/SiO₂ concentration.

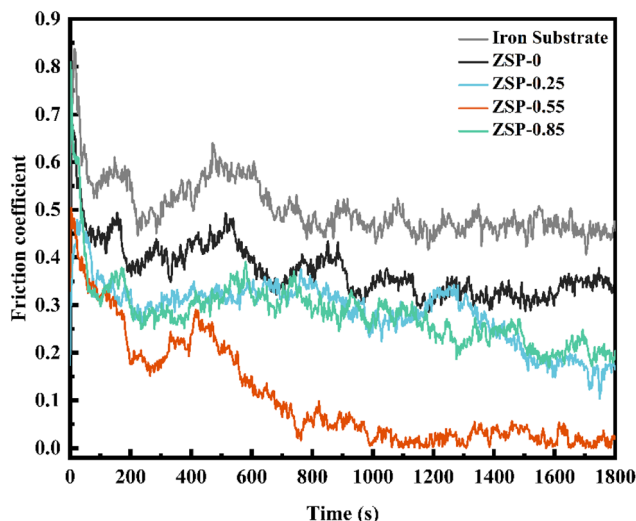


Fig. 6 Friction coefficients curves of phosphate coatings under 100 rpm and a fixed load of 100 N for 30 min at ambient temperature.

characteristics, the phosphate coatings effectively reduce the friction coefficient of friction pair surface. The coefficient of friction for pure metallic substrates is approximately 0.46. When 0.55 g L^{-1} PDA@ α -ZrP/SiO₂ was incorporated, the friction resistance coefficient of phosphate coating is reduced to 0.02. Smaller friction coefficient means better friction resistance.^{33–36}

The 2D α -ZrP nanosheets have large specific surface area, smooth surface atoms, low shear strength, excellent mechanical properties, and have low shear stress when slip occurs, which greatly enhances the friction resistance of phosphate coatings. In addition, the addition of 0D SiO₂ can also enhance the hardness and friction resistance of phosphate coatings. It can be found from Fig. 3 that after the introduction of PDA@ α -ZrP/SiO₂, the surface of phosphate coatings become more dense, uniform and fine, which also reduces mechanical movement resistance and noise, and improves the friction resistance of the phosphate coatings. The enhancement of friction resistance is expected to broaden the application fields of phosphate coatings.

3.4 Electrochemical corrosion behavior of phosphate coatings

To assess the corrosion resistance of phosphate coatings, the potentiodynamic polarization curve test was conducted, and the results are depicted in Fig. 7. From Fig. 7, it is evident that the composite phosphate coatings, after incorporating the nanocomposite, exhibits a smaller current density and a more positive potential compared to the blank phosphate coating. Tafel extrapolation method was applied to analyze the test data to obtain parameters including corrosion potential (E_{corr}), corrosion current (I_{corr}), cathode and anode slope (β_a and β_b), polarization resistance (R_p) and corrosion rate (CR). The corresponding data are presented in Table 1. The corrosion resistance of phosphate coating can be intuitively judged by the values of R_p and CR.³⁷ R_p of blank phosphate coating is $1381 \Omega \text{ cm}^2$, and the CR is 3.72 mpy. The R_p of the 2D/0D nanocomposite phosphate coatings is greatly improved, and the CR is significantly reduced. Especially when the introduction amount of PDA@ α -ZrP/SiO₂ is 0.55 g L^{-1} , the R_p rose to $20\,520 \Omega \text{ cm}^2$ and CR decreased to 0.21 mpy. As shown in the SEM results, the introduction of nanocomposites effectively increased the number of nucleated phosphate crystals, refined the grain size, and improved the coverage of phosphate coatings on the metal substrate. A denser film layer has the better shielding effect on corrosive media, which is the fundamental reason why CR decreases.

The porosity of the phosphate coatings was also analyzed and calculated. The following formula is used to calculate the porosity of phosphate coatings.³⁸

$$p = \frac{R_{\text{ps}}}{R_{\text{ps}}} 10^{-\frac{\Delta E_{\text{corr}}}{\beta_a}}$$

Among them, P , R_{ps} and R_p respectively represent the porosity of phosphate coating, the polarization resistance of bare metal substrate and phosphate coating. ΔE_{corr} is the

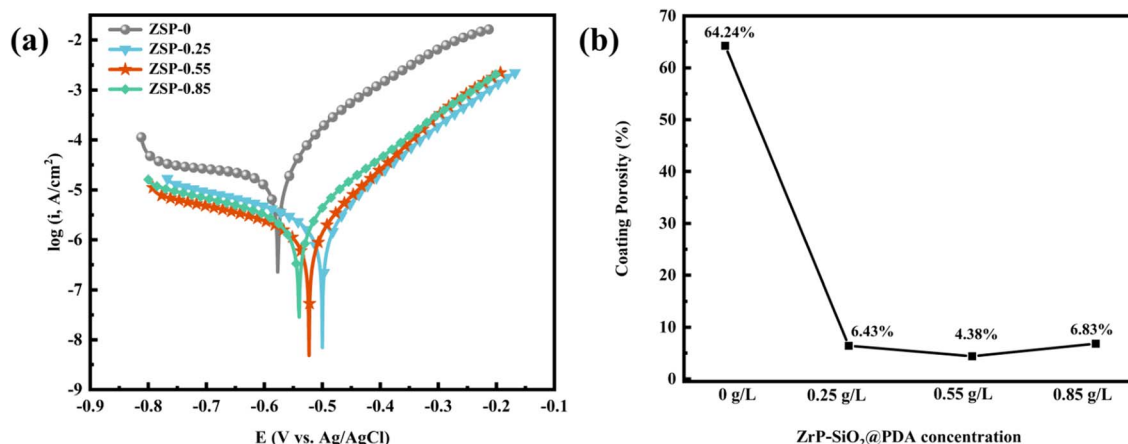


Fig. 7 Potentiodynamic polarization curves (a) and coating porosity (b) of phosphate coatings with different content of PDA@ α -ZrP/SiO₂.



Table 1 Tafel extrapolation results obtained from Fig. 7a

Specimens	E_{corr} (V)	I_{corr} (A cm^{-2})	R_p ($\Omega \text{ cm}^2$)	β_a (mV dec^{-1})	β_c (mV dec^{-1})	CR (mpy)
ZSP-0	−0.55	2.55×10^{-5}	1381	10.78	−1.58	3.72
ZSP-0.25	−0.47	2.10×10^{-6}	14 042	10.65	−4.11	0.31
ZSP-0.55	−0.49	1.45×10^{-6}	20 520	10.81	−3.86	0.21
ZSP-0.85	−0.50	2.75×10^{-6}	13 184	9.13	−2.88	0.40

Table 2 Tafel extrapolation results of bare metal substrate

Specimens	E_{corr} (V)	I_{corr} (A cm^{-2})	R_p ($\Omega \text{ cm}^2$)
Bare metal substrate	−0.62	2.53×10^{-5}	874

corrosion potential difference between the bare metal substrate and phosphate coating sample and β_a represents the anode slope. The test data of the bare metal substrate are listed in Table 2, where E_{corr} is −0.62 V and the R_p is $874 \Omega \text{ cm}^2$. The coating porosity of the phosphate coatings was calculated through the above formula and the results were plotted in Fig. 7b. The porosity of the blank phosphate coating is 64.24%. When the PDA@ α -ZrP/SiO₂ was introduced, the porosity of the phosphate coatings drops below 10%. The reduction in porosity effectively enhances the corrosion resistance of phosphate coatings.

EIS, as a rapid and minimally invasive inspection method, was conducted to evaluate the corrosion resistance of

phosphate coatings.^{39–41} The Nyquist and Bode plots of ZSP-0, ZSP-0.25, ZSP-0.55, and ZSP-0.85 are illustrated in Fig. 8. The Nyquist plot reveals a direct correlation with the corrosion resistance of phosphate coatings. Specifically, ZSP-0.55 displays the largest arc radius among all samples, indicating its superior corrosion resistance (Fig. 8a). Furthermore, the impedance modulus ($|Z|_{0.01 \text{ Hz}}$) at 0.01 Hz frequency is used to analyze the barrier properties of the phosphate coatings. The higher $|Z|_{0.01 \text{ Hz}}$ value indicates that the phosphate coating has better blocking effect on corrosive media.^{42,43} In Fig. 8b, $|Z|_{0.01 \text{ Hz}}$ values of phosphate coatings decrease in the following order: ZSP-0.55 > ZSP-0.25 > ZSP-0.85 > ZSP-0, revealing that the incorporation of PDA@ α -ZrP/SiO₂ positively impacted the corrosion resistance of phosphate coating. Nevertheless, ZSP-0.85 sample exhibited a lower $|Z|_{0.01 \text{ Hz}}$ value compared to that of the ZSP-0.55 sample, which may be that when a large amount of PDA@ α -ZrP/SiO₂ was introduced, PDA@ α -ZrP/SiO₂ would occupy the nucleation position of phosphate crystals and thus affect the film-forming effect of phosphate coatings. In Fig. 8c, compared with the

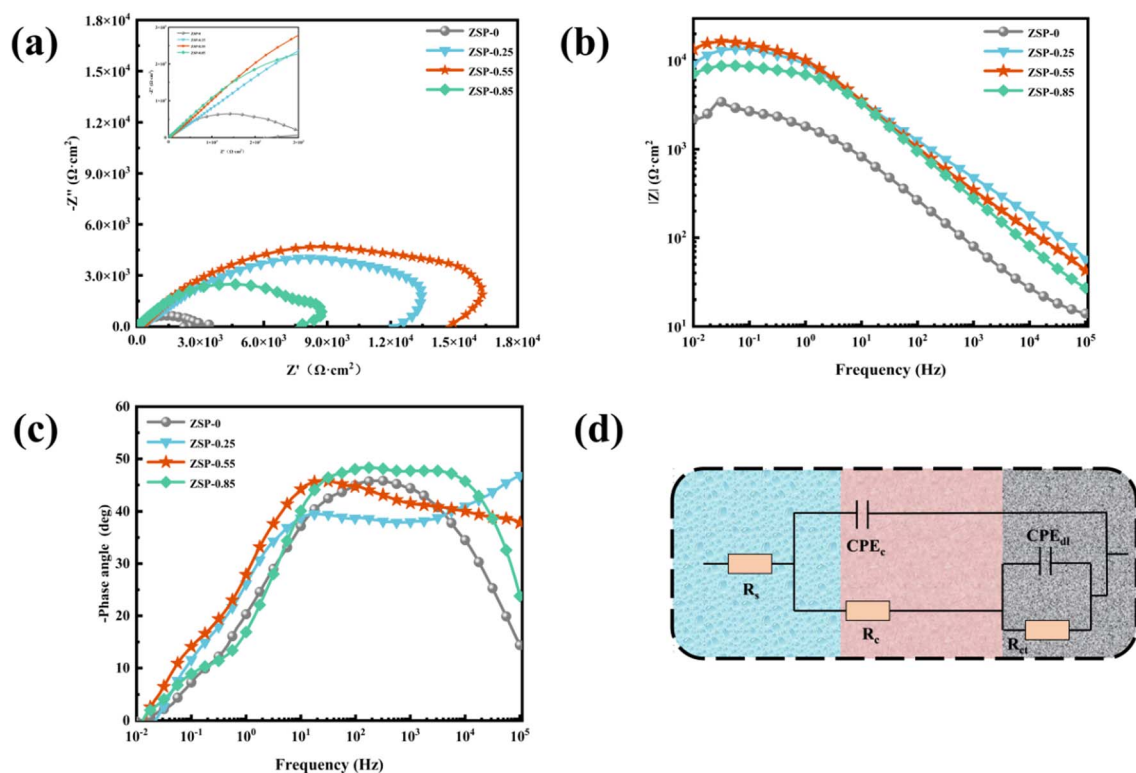


Fig. 8 EIS Nyquist plots (a), EIS Bode plots of different phosphate coatings (b and c), and the equivalent electrical circuit (d).



Table 3 Fitting values obtained from Fig. 8 using equivalent electrical circuit

Specimens	R_s (Ω cm ²)	CPE _c		R_c (Ω cm ²)	CPE _{dl}		R_{ct} (Ω cm ²)
		Y (Ω^{-1} cm ⁻² s ^{<i>n</i>})	<i>n</i>		Y (Ω^{-1} cm ⁻² s ^{<i>n</i>})	<i>n</i>	
ZSP-0	34	4.9×10^{-5}	0.64	612	1.4×10^{-5}	0.70	5029
ZSP-0.25	35	7.7×10^{-6}	0.67	1097	8.0×10^{-6}	0.52	54 840
ZSP-0.55	34	4.0×10^{-6}	0.57	1881	2.4×10^{-6}	0.64	57 890
ZSP-0.85	34	4.5×10^{-6}	0.61	1235	1.6×10^{-6}	0.61	28 320

blank phosphate coating, the nanocomposite phosphate coatings have the higher and wider phase angle, which also confirms that PDA@ α -ZrP/SiO₂ has great promotion effect on the phosphating process.

To delve deeper into the corrosion resistance of phosphate coatings, a comprehensive analysis of EIS data was undertaken utilizing the equivalent circuit illustrated in Fig. 8d. The electrochemical parameters are documented in Table 3. R_s , R_c , R_{ct} , CPE_c and CPE_{dl} symbolize the solution resistance, coating pore resistance, charge transfer resistance, coating capacitance and double-layer capacitance.⁴⁴ High R_c and low CPE_c usually indicate reduced permeability to H₂O and O₂. Remarkably, the ZSP-0.55 sample has the peak R_c value among the tested samples, affirming its supreme corrosion resistance. Additionally, the R_{ct} value of ZSP-0.55 is significantly higher than that of ZSP-0, underscoring a dramatic boost in corrosion inhibition. This pronounced improvement is attributed to the small porosity of ZSP-0.55.

The copper sulfate dropping test was further carried out to test the corrosion resistance of phosphate coatings. The principle of the copper sulfate dropping test is that a substitution reaction occurs between CuSO₄ and Fe in the metal substrate, resulting in discoloration. The length of discoloration time is used to determine the porosity and ability to block corrosive media of phosphate coatings. The results are shown in Fig. 9.

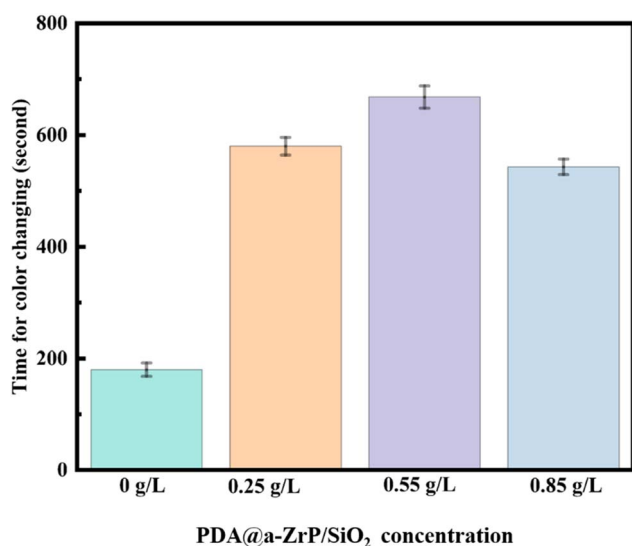
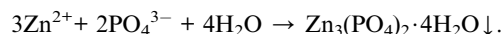
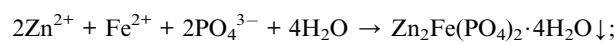
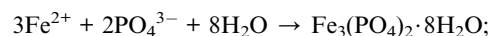
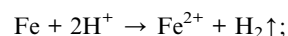


Fig. 9 The time for color changing of samples in copper sulfate dropping experiments.

Discoloration time of the blank phosphate coating is only 180 s. When the 2D/0D nanocomposite material was introduced, the discoloration time becomes significantly longer, which is consistent with the results of electrochemical corrosion resistance test. Discoloration time of 0.55 g L⁻¹ ZSP sample is 668 s, which also proves that the 2D/0D nanocomposite effectively reduced the coating porosity and enhanced the shielding effect of phosphate coating on corrosive media.

From the study of mechanism of traditional phosphating, the reactions that occur during phosphating process are as follows:



When Fe²⁺, PO₄³⁻ and Zn²⁺ ions reach saturation, phosphate coating forming reactions occur, that is, insoluble phosphate crystals are deposited on the surface of metal substrate. Therefore, it is hoped that the ions participating in reactions can quickly reach saturation on surface of metal substrate. α -ZrP has large specific surface area and negative charge density, and can be absorbed on the surface of metal substrate as the new cathode to accelerate hydrogen evolution reaction. In addition, it has strong adsorption capacity for Fe²⁺ and Zn²⁺, which makes the ions concentration reach saturation faster, thereby promoting the production of more phosphate crystals. The addition of SiO₂ also changes the initial potential of the interface between substrate surface and phosphating solution, reduces the activation energy of phosphating process, and increases the phosphating nucleation sites. 2D α -ZrP and 0D SiO₂ synergistically promote the phosphating process, forming the uniform and dense phosphate coatings.

In addition, PDA uses its own adhesion to wrap the nanoparticles, so that the nanoparticles show strong stability in the phosphating solution. Moreover, the catechol is strong ligand for metal ions including Fe²⁺ and Zn²⁺, which also allows the metal ions at the interface between substrate and phosphating solution to reach saturation faster and generate more phosphate crystals. The corrosion resistance and friction resistance of the more uniform and dense phosphate coating are enhanced.



4. Conclusions

In this study, α -ZrP and SiO₂ nanocomposites modified with PDA were introduced into the phosphating solution and used as new phosphate accelerators. SEM results confirms that PDA@ α -ZrP/SiO₂ effectively increased the number of phosphating nucleation in the initial stage of phosphate coating formation and further improved the coverage of phosphate coating on the metal substrate. It not only improves the friction resistance, but also enhances the corrosion resistance of phosphate coating. The friction coefficient of phosphate coating was reduced from 0.32 to about 0.02. In the corrosion resistance test, R_p of blank phosphate coating is 1381 Ω cm², while R_p of ZSP-0.55 could be increased to 20 520 Ω cm². And the coating porosity of phosphate coatings dropped from 64.24% to 4.38% via the incorporation of PDA@ α -ZrP/SiO₂. 2D/0D nanocomposites provide feasible solution to promote phosphating and reduce phosphating deposits.

Data availability

The authors confirm that the data supporting the findings of this study are available within the article.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

This work was supported by the Hongliu Outstanding Young Talent Support Program of Lanzhou University of Technology (Grant No. 062313); the Centrally Guided Local Science and Technology Development Fund Project (Grant No. 22ZY1QA011); and the Lanzhou Talent Innovation and Entrepreneurship Project (Grant No. 2023-RC-46).

References

- 1 A. H. Riyas, C. V. Geethanjali, S. Arathy, A. Anil and S. M. A. Shibli, *Appl. Surf. Sci.*, 2022, **593**, 153370.
- 2 W. Wang, Y. Tian, H. Shen and X. Zhang, *J. Alloys Compd.*, 2023, **955**, 170247.
- 3 J. Wu, Y. Chen, L. Zhang and X. Sheng, *J. Ind. Eng. Chem.*, 2024, **129**, 424–434.
- 4 Y. Tian, H. Huang, H. Wang, Y. Xie, X. Sheng, L. Zhong and X. Zhang, *J. Alloys Compd.*, 2020, **831**, 154906.
- 5 S. Silva-Fernández, B. Díaz, I. Feijoo, X. R. Nóvoa and C. Pérez, *Electrochim. Acta*, 2023, **457**, 142510.
- 6 X. Huang, Q. Weng, Y. Chen, L. Zhang and X. Sheng, *Surf. Interfaces*, 2024, **45**, 103911.
- 7 R. Ramanauskas, O. Girčienė, L. Gudavičiūtė and A. Selskis, *Appl. Surf. Sci.*, 2015, **327**, 131–139.
- 8 S. Silva-Fernández, B. Díaz, R. Figueroa, X. R. Nóvoa and C. Pérez, *Surf. Coat. Technol.*, 2024, **476**, 130260.
- 9 X. He, J. Wu, Y. Chen, L. Zhang and X. Sheng, *Appl. Surf. Sci.*, 2022, **603**, 154455.

- 10 H. Yue, D. Wang, J. Zhu and L. Li, *ACS Sustain. Chem. Eng.*, 2017, **5**, 3019–3026.
- 11 M. A. Hafeez, A. Farooq, A. Zang, A. Saleem and K. M. Deen, *J. Coat. Technol. Res.*, 2020, **17**, 827–849.
- 12 E. Rumyantsev, V. Rumyantseva and V. Konovalova, *Coatings*, 2022, **14**, 498.
- 13 D. P. Burduhos-Nergis, A. V. Sandu, D. D. Burduhos-Nergis, P. Vizureanu and C. Bejinariu, *Arch. Metall. Mater.*, 2023, **68**, 1029–1034.
- 14 Y. Xie, M. Chen, D. Xie, L. Zhong and X. Zhang, *Corros. Sci.*, 2017, **128**, 1–8.
- 15 H. Khosravi, R. Naderi and B. Ramezanzadeh, *Mater. Today Chem.*, 2023, **27**, 101282.
- 16 M. Tamilselvi, P. Kamaraj, M. Arthanareeswari, S. Devikala and J. A. Selvi, *Appl. Surf. Sci.*, 2015, **332**, 12–21.
- 17 M. Tamilselvi, P. Kamaraj, M. Arthanareeswari and S. Devikala, *Appl. Surf. Sci.*, 2015, **327**, 218–225.
- 18 M. Arthanareeswari, P. Kamaraj, M. Tamilselvi and S. Devikala, *Mater. Today: Proc.*, 2018, **5**, 9012–9025.
- 19 M. Tamilselvi, P. Kamaraj, M. Arthanareeswari, S. Devikala, J. Arockiaselvi and T. Pushpamalini, *Mater. Today: Proc.*, 2018, **5**, 8880–8888.
- 20 H. Huang, H. Wang, Y. Xie, D. Dong, X. Jiang and X. Zhang, *Surf. Coat. Technol.*, 2019, **374**, 935–943.
- 21 S. Hu, M. Muhammad, M. Wang, R. Ma, A. Du, Y. Fan, X. Cao and X. Zhao, *Mater. Lett.*, 2020, **265**, 127256.
- 22 M. Wang, R. Ma, A. Du, S. Hu, M. Muhammad, X. Cao, Y. Fan, X. Zhao and J. Wu, *Mater. Chem. Phys.*, 2020, **250**, 123056.
- 23 X. He, S. Li, J. Wu, Y. Chen, L. Zhang and X. Sheng, *Ind. Eng. Chem. Res.*, 2022, **61**, 12576–12589.
- 24 Y. Ma, H. Huang, H. Zhou, M. Graham, J. Smith, X. Sheng, Y. Chen, L. Zhang, X. Zhang, E. Shchukina and D. Shchukin, *J. Mater. Sci. Technol.*, 2021, **95**, 95–104.
- 25 H. Jiang, J. Li, Y. Xie, H. Guo, M. He, X. Shi, Y. Mei, X. Sheng and D. Xie, *J. Mater. Sci. Technol.*, 2025, **209**, 207–218.
- 26 Y. Fan, J. Wu, L. Zhang and X. Sheng, *Ind. Eng. Chem. Res.*, 2024, **63**, 10637–10650.
- 27 X. He, J. Wu, S. Li, Y. Chen, L. Zhang and X. Sheng, *Prog. Org. Coat.*, 2022, **171**, 107042.
- 28 J. Li, F. Meng, L. Liu, Y. Cui, R. Liu, H. Zheng and F. Wang, *Corros. Commun.*, 2022, **5**, 62–72.
- 29 Y. Tian, W. Qiu, Y. Xie, H. Huang, J. Hu, L. Zhong, X. Jiang and X. Zhang, *J. Electrochem. Soc.*, 2020, **167**, 101505.
- 30 B. Ramezanzadeh, Z. Haeri and M. Ramezanzadeh, *Chem. Eng. J.*, 2016, **303**, 511–528.
- 31 Y. Zhao, Y. He, S. Yan, C. Li, H. Li, W. Chen, J. Yan, G. Wu and X. Yuan, *Colloids Surf., A*, 2023, **656**, 130472.
- 32 R. Amini and A. A. Sarabi, *Appl. Surf. Sci.*, 2011, **257**, 7134–7139.
- 33 X. He, Q. Weng, R. Guo, L. Zhang, X. Sheng and D. Xie, *J. Mater. Sci. Technol.*, 2024, **201**, 197–209.
- 34 X. Chang, X. Chen, Q. Zhang, Y. Lei, D. Wang, J. Li and S. Sun, *Corros. Commun.*, 2021, **4**, 1–11.
- 35 Q. Weng, L. Zhang, X. Sheng and D. Xie, *J. Mater. Sci. Technol.*, 2024, DOI: [10.1016/j.jmst.2024.07.023](https://doi.org/10.1016/j.jmst.2024.07.023).



- 36 A. Kumar, H. Rachwani and M. Kumar Singh, *Proc. Inst. Mech. Eng., Part J.*, 2024, **238**, 662–675.
- 37 Y. Tian, L. Zhong, X. Sheng and X. Zhang, *Adv. Compos. Hybrid Mater.*, 2022, **5**, 1922–1938.
- 38 B. Elsener, A. Rota and H. Böhni, *Mater. Sci. Forum*, 1991, **44–45**, 29–38.
- 39 P. Zhou, L. Yang, Y. Hou, G. Duan, B. Yu, X. Li, Y. Zhai, B. Zhang, T. Zhang and F. Wang, *Corros. Commun.*, 2021, **1**, 47–57.
- 40 B. Rebbah, A. El Haib, S. Lahmady, I. Forsal, M. Gouygou, S. Mallet-ladeira, A. Medaghri-alaoui, E. M. Rakib and A. Hannioui, *RSC Adv.*, 2024, **14**, 14152–14160.
- 41 Q. A. Yousif, Z. Fadel, A. M. Abuelela, E. H. Alosaimi, S. Melhi and M. A. Bedair, *RSC Adv.*, 2023, **13**, 13094–13119.
- 42 P. Li, Z. Shao, W. Fu, W. Ma, K. Yang, H. Zhou and M. Gao, *Corros. Commun.*, 2023, **9**, 44–56.
- 43 Z. Huang, Q. Yong and Z.-H. Xie, *Corros. Commun.*, 2023, **10**, 38–47.
- 44 M. Cui, D. I. Njoku, B. Li, L. Yang, Z. Wang, B. Hou and Y. Li, *Corros. Commun.*, 2021, **1**, 1–9.

