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Electrochemical corrosion performance and microhardness of the electrodeposited Ni-CeO₂ and Ni-ZrO₂ nanocomposite coatings on the AA2219 aluminum alloy

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Although Ni-based composite coatings have been widely studied, a systematic and direct comparison of the CeO₂- and ZrO₂-reinforced Ni nanocomposite coatings on AA2219 remains limited. Nickel-based nanocomposite coatings are widely used to enhance the surface hardness and corrosion resistance of the AA2219 aluminium alloy in chloride-containing environments. In this work, to increase the durability of the AA2219 aluminium alloy, nickel nanocomposite coatings reinforced with varying contents of CeO₂ and ZrO₂ nanoparticles were fabricated on the alloy using composite electroplating. The results demonstrate that the uniform nanoparticle dispersion, particle chemistry, and optimized concentration (3.5 wt%) significantly enhance microhardness from 422 HV for pure Ni to 1089 HV for Ni-ZrO₂ and 1055 HV for Ni-CeO₂. Electrochemical corrosion resistance results for Ni-CeO₂ show a significant reduction in corrosion current density (I_{corr}) to 1.08 μA compared with 14.7 μA for Ni coatings, and charge-transfer resistance (R_{ct}) of 25.1 k Ω compared with 0.386 k Ω for Ni coatings, corresponding to a significant enhancement and confirming the strong barrier effect of nanoparticle dispersion. Microstructural characterization and nanoparticle distribution were examined using transmission electron microscopy (TEM), scanning electron microscopy (SEM), and energy-dispersive X-ray spectroscopy (EDS), revealing pronounced grain refinement induced by both CeO₂ and ZrO₂ nanoparticles. The key scientific contribution of this study lies in the direct, mechanism-based comparison of CeO₂- and ZrO₂-reinforced Ni coatings deposited under identical conditions, providing new insights into their respective strengthening and anticorrosion mechanisms. These results advance the current understanding of Ni-based composite coatings and provide a useful framework for designing high-performance protective coatings for aluminium alloys for future aerospace coating applications.

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1 Introduction

Aluminium alloy AA2219 is a heat-treatable wrought alloy widely used in demanding engineering applications in the marine and aerospace industries due to its high strength-to-weight ratio, excellent weldability, and high fracture toughness. It exhibits

reliable load-bearing performance over a wide temperature range.^{1–3} However, Al alloys are highly susceptible to galvanic and atmospheric corrosion because of the galvanic couple formation between CuAl₂ intermetallic precipitates and the aluminium matrix. These precipitates tend to form during thermal processing and preferentially segregate along grain boundaries, which significantly accelerates corrosion, particularly in a chloride-rich environment.⁴ Although a high copper content (≈ 5.8 – 6.8 wt%) is essential for achieving superior mechanical strength and thermal stability, it simultaneously increases the corrosion susceptibility of the alloy.^{5–7} To address these challenges and extend the service life while enhancing mechanical performance and corrosion resistance, surface modifications have attracted considerable research attention.^{2,8} Material degradation is typically initiated at the surface. Electrodeposited metal–ceramic composite coatings are promising candidates for commercial applications in harsh and chemically aggressive environments. Nickel-based coatings, in

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particular, are widely employed due to their good corrosion resistance, strong adhesion to substrates, high hardness, and long-term reliability across various engineering fields.^{9–11} The Ni-electroplated coatings with the incorporation of ceramic or second-phase reinforcement particles, such as TiO₂, SiC, CeO₂, Al₂O₃, and ZrO₂, have been shown to further enhance the mechanical properties and corrosion resistance. In particular, the dispersion of CeO₂ nanoparticles in Ni matrices has been reported to significantly increase microhardness (up to ~824 HV) compared with pure Ni, while also promoting grain refinement and dispersion strengthening, thereby improving corrosion protection through the formation of an effective passive barrier.¹² M. Safavi *et al.*¹⁰ reported that the metal composite coatings exhibit significantly improved mechanical properties, wear resistance, and corrosion performance. The Ni-P coatings reinforced with CeO₂ on a mild steel substrate show significant improvements in microhardness and corrosion resistance compared to those without the CeO₂ dispersion Ni-P coatings.¹³ It has been reported that the corrosion and wear resistance of AA6060 alloy are significantly enhanced by the incorporation of ceria (CeO₂) nanoparticles into electroplated Ni composite coatings. The resulting composite coating exhibited a microhardness of approximately 560 HV, representing a substantial improvement compared with ~421 HV for pure Ni coatings. However, this strengthening effect is non-linear, as excessive nanoparticle content can lead to particle agglomeration, resulting in a rough and porous coating morphology that degrades mechanical integrity and overall coating performance.¹⁴ Most recently, Farough *et al.*¹⁵ successfully electrodeposited pure Co and Co/CeO₂ composite coatings onto carbon steel substrates and systematically investigated the effects of CeO₂ concentration, current density, and coating thickness on corrosion behaviour, mechanical properties, and wear resistance. Potentiodynamic polarization (PDP) results indicated that both pure Co and Co/CeO₂ coatings exhibited less noble corrosion potentials compared with Ti-6Al-4V, while no pitting corrosion was observed. Electrochemical impedance spectroscopy (EIS) measurements further revealed a significant increase in total polarization resistance, reaching approximately 1540 Ω cm² for pure Co and 1777 Ω cm² for the Co/CeO₂ composite coating, confirming enhanced corrosion protection. Moreover, the incorporation of CeO₂ nanoparticles markedly improved the wear resistance of the composite coating compared with plain Co coatings and the uncoated carbon steel substrate. Zirconia nanoparticles with their well-dispersed characteristics enhance microhardness and corrosion resistance by promoting microstructural refinement and passive barrier effect, which is a more uniform and impermeable layer.¹⁶ A Ni-P coating with ZrO₂ nanoparticle reinforcement on a copper substrate shows a significant increase in microhardness, which they correlated to the dispersion strengthening mechanism (Orowan strengthening) imparted by the hard ZrO₂ particles, which explores the electrodeposition route for producing Ni-ZrO₂ coatings.¹⁷ Pulse electrodeposited Ni-P-ZrO₂ and Ni-W/ZrO₂ nanocomposite coatings, in which ZrO₂ nanoparticles have also been incorporated into PEO coatings on Ti-6Al-4V substrates, influence corrosion, antibacterial properties,

and microstructure, also with decreased grain diameters, from 0.55 μm for pure Ni to 0.31 μm, and a significantly increased hardness, from 325 HV to 401 HV.¹⁸ Numerous studies have reported the incorporation of zirconia and ceria nanoparticles using various approaches to improve wear and corrosion resistance and mechanical properties. The existing literature lacks detailed studies to address the systematic and direct comparison of the dispersion of individual CeO₂ and ZrO₂ nanoparticles in Ni electroplating under the same deposition conditions, which produce galvanic cells that weaken the Ni coating. This work shows that the active redox nature of ceria provides excellent long-term protection and reduces galvanic drive more efficiently than the physical barrier provided by zirconia. In this study, the individual role of CeO₂ and ZrO₂ nanoparticles on grain refinement, improved hardness and corrosion of Al alloy are investigated. In addition, a significant improvement in microhardness was achieved due to controlled nanoparticle dispersion and the formation of a nanocrystalline Ni film, which therefore provides new insights into the comparative analysis of CeO₂ and ZrO₂ in Ni coatings, offering a simple and cost-effective approach to fabricate a predictive model to protect AA2219 alloy in chloride-rich environments Table 1.

2 Experimental and characterization

2.1 Materials

Pure aluminium alloy AA2219 samples, with wt% compositions, as shown in Table 2, and dimensions of 15 × 10 × 2 mm³ were cut from the plate with purity >99%. A pure Ni plate was used as the anode material. The anode had a rectangular shape with dimensions of 50 × 30 × 5 mm³. ZrO₂ and CeO₂ nanoparticles were directly received from Macklin Company, Shanghai, China, as reported in our previous work. Analytical-grade chemicals, for the electroplating solution, including nickel sulfate hexahydrate (NiSO₄·6H₂O), boric acid (H₃BO₃), trisodium citrate dihydrate (C₆H₅Na₃O₇·2H₂O), and sodium chloride (NaCl), were purchased from Macklin Biochemical Co., Ltd, Shanghai, China.

2.2 Fabrication of nanocomposite coatings on AA2219 alloy

Pure aluminum alloy AA2219 samples with dimensions of 15 × 10 × 2 mm³ were cut from the plate with purity >99%, polished

Table 1 Electrodeposition parameters for the Ni-based nanocomposite coatings

Parameter	Value/description
Substrate (cathode)	AA2219 alloy
Anode	Pure Ni plate
Anode dimension	50 × 30 × 5 mm ³
Inter-electrode gap	~2 to 3 cm
pH	5.5–6.0
Current density	1.5 A dm ⁻²
Bath temperature	40 °C
Deposition time	50–60 minutes
Magnetic stirrer	200 rpm



Table 2 Composition of the aluminum alloy (AA2219)

Aluminium (Al)	Copper (Cu)	Manganese (Mn)	Silicon (Si)	Titanium (Ti)	Vanadium(v)	Zinc (Zn)	Iron (Fe)
91.5 to 93.8 wt%	5.8 to 6.8 wt%	0.2 to 0.4 wt%	0.2 wt% max	0.02 to 0.10 wt%	0.05 to 0.15 wt%	0.03 wt% max	0.05 to 0.15 wt%

with SiC paper, starting from 320 to 1200 grit, and cleaned ultrasonically with ethanol and acetone. The procedure for the surface treatment of an aluminium alloy prior to Ni electroplating involves several key steps to ensure a uniform and strongly adherent coating.^{19,20} The primary challenge is overcoming the natural surface oxide film on aluminium, which prevents proper plating. The sample is immersed in an alkaline solution. After alkaline cleaning, the specimen is rinsed and then immersed in a nitric acid solution (*e.g.*, 65% HNO₃ for 5 seconds) to neutralize any remaining alkaline residues and to activate the surface. The double zincate process deposits a thin, uniform, and adherent layer of zinc onto the aluminium surface, which acts as a conductive layer and provides a compatible base for the subsequent nickel plating. The specimen is first immersed in a zincate bath solution for 45 s, rinsed and again immersed in the zincate bath solution for 15 s at room temperature, then dried by using a mechanical drier. After the zincate treatment, a thin copper layer is electroplated on the specimen in a copper bath at room temperature without magnetic stirring. The deposition time is 25 min with the same parameters as Ni electroplating, as shown in Table 1. The sample is again dried using a mechanical drier, and subsequently, the sample is ready for electrodeposition.²¹ Nanocomposite Ni films were electrodeposited on an AA2219 substrate at a current density of 1.5 A dm⁻² using a bath containing 150 g l⁻¹ NiSO₄·6H₂O, 35 g l⁻¹ H₃BO₃, 120 g l⁻¹ C₆-H₅Na₃O₇·2H₂O, and 12 g l⁻¹ NaCl. ZrO₂ and CeO₂ nanoparticles were added separately to the bath at concentrations ranging from 5 to 10 g l⁻¹. Then, samples with different wt% of Ni-ZrO₂ or Ni-CeO₂ nanocomposite are prepared by the same procedure.²²⁻²⁴ The weight percentages of dispersed oxide particles in the Ni films were determined by EDS using the INCA software package attached to the SEM instrument. Fig. 1 shows a schematic of composite coatings deposited on the AA2219 alloy. The bath was maintained at 40 °C with a pH of 5.5–6.0. These nanoparticles

were ultrasonically dispersed for 30 min at 35 kHz before electrodeposition,²⁵ and the solution was continuously stirred at 200 rpm using a magnetic stirrer to prevent nanoparticle agglomeration. The interelectrode gap was ~2 to 3 cm.²⁶⁻³⁰ The electroplated samples were washed with acetone/ethanol, distilled water, and dried with a mechanical drier. The complete list of electrodeposition parameters for the Ni-based nanocomposite coatings is shown in Table 1.

2.3 Characterization

Field-emission scanning electron microscopy (FE-SEM, Inspect F50, Holliboro) was used to analyze the surface morphology of the as-prepared nanocomposite coatings. SEM/EDS results confirmed the presence of nanoparticles and quantified the weight percentage of dispersoids after electroplating. Transmission electron microscopy (TEM, JEM-2100F, JEOL, Japan) was used to confirm the size of the as-received nanoparticles. A Vickers microhardness tester (HMV Shimadzu company) was employed to measure microhardness of the layers. The measurements were performed at a load of 10 N for 10 s. The average of five measurements is presented. The electrochemical corrosion experiment was conducted using a Gamry electrochemical workstation, where the nickel-coated sample served as the working electrode, Ag/AgCl as the reference electrode, and a platinum mesh as the counter electrode. Tafel curves and impedance data of the coatings were obtained. The interelectrode gap was maintained at approximately 5–10 mm, and the immersed surface area in the solution was about 1 cm². The size of the platinum mesh electrode was 1 cm. Corrosion tests were conducted at room temperature using a working electrode with an area of 1 cm² immersed in a 3.5 wt% sodium chloride (NaCl) solution at room temperature in a 60 mL solution with natural aeration. The sample was initially placed in the solution prior to the test to allow the open-circuit potential to stabilize before measurement. Tafel experiments were performed with a 4 mV s⁻¹ scan rate, and electrochemical impedance spectroscopy (EIS) measurements were used to analyze the EIS at a frequency range of 0.1 Hz to 100 kHz. Tafel and EIS results of the Ni nanocomposite samples were analyzed by the Gamry Echem analyst software to determine the corrosion parameters. The coating thickness was experimentally confirmed in the laboratory using a LANDTEK digital meter, showing a thickness of approximately 20 μm. To confirm the accuracy of the results, the experiment was repeated three times.

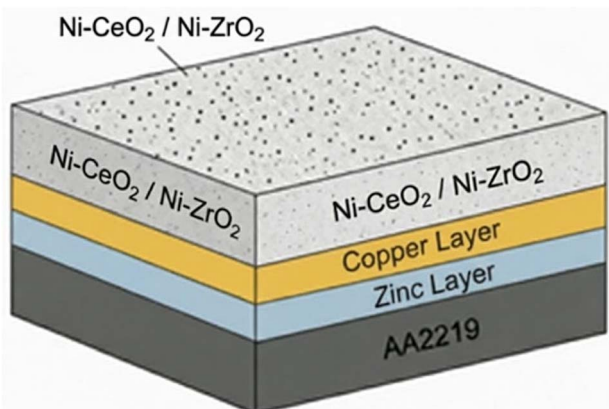


Fig. 1 Schematic of the Ni-coated samples with CeO₂ and ZrO₂ nanoparticles on the AA2219 alloy.

3 Results and discussion

3.1 TEM results of the as-received nanoparticles and as-fabricated nanocomposite coatings

Fig. 2 displays the as-received CeO₂ and ZrO₂ nanoparticles and their size distributions, along with the microstructure and grain



size of the Ni nanocomposite film with dispersed ZrO_2 nanoparticles. TEM results confirm the morphologies of the as-received CeO_2 and ZrO_2 nanoparticles, as shown in Fig. 2(a) and (c), respectively. The TEM analysis displays that ZrO_2 nanoparticles exhibit smaller and more homogeneous particle sizes, whereas CeO_2 nanoparticles appear as irregular and partially agglomerated clusters. Furthermore, the average particle sizes of CeO_2 and ZrO_2 nanoparticles were calculated to be approximately 31 nm and 26 nm, respectively, using the Nano measurer software, as shown in Fig. 2(b) and (d). Fig. 2(e) shows the surface microstructure of the as-prepared Ni nanocomposite film obtained from TEM analysis. The surface microstructure confirms that no cracks, cavities, or pores are present in the coatings. However, some twin structures were observed within the Ni film grains. The results also demonstrate

a uniform distribution of ZrO_2 nanoparticles throughout the nanocomposite Ni film, as indicated by the yellow arrows as shown in Fig. 2(e). In addition, localized agglomeration of ZrO_2 nanoparticles can be observed in certain areas, as shown in Fig. 2(e). To further confirm the grain size of the nanocomposite Ni film with dispersed ZrO_2 nanoparticles, the average grain size was measured to be approximately 50 nm using the Nano measurer software, as shown in Fig. 2(f). In a previous work by our group,³¹ it was reported that Ni films with dispersed CeO_2 nanoparticles exhibited an average grain size of approximately 61 nm, whereas pure Ni films showed an average grain size of around 91 nm, which is significantly larger than that of Ni films with CeO_2 and ZrO_2 nanoparticle dispersions. More recently, the authors reported²⁹ that pure Ni films exhibited an average grain size of approximately 100 nm. The dispersion of CeO_2 and

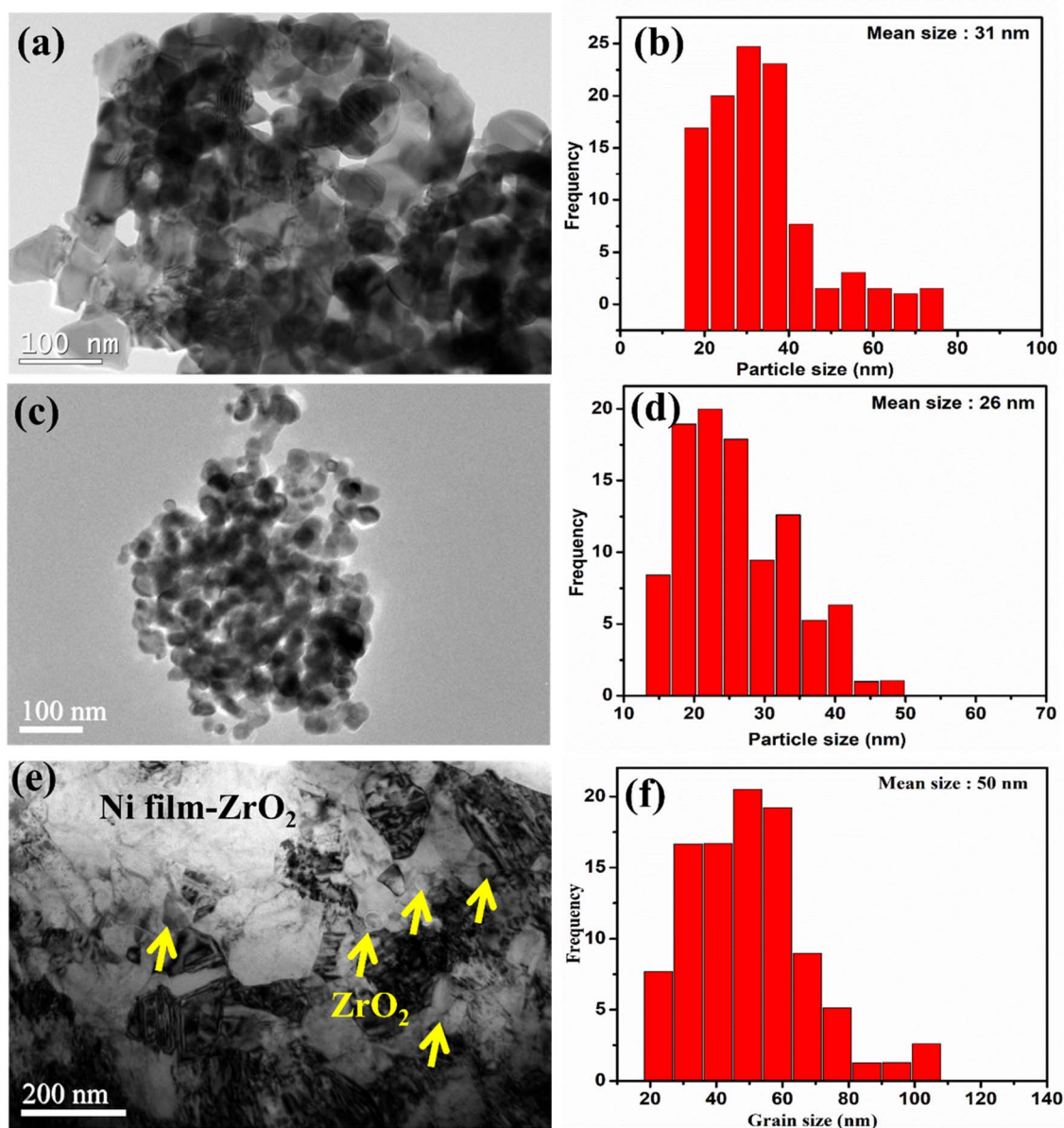


Fig. 2 TEM images of the as-received CeO_2 and ZrO_2 nanoparticles (a and c), respectively, and their corresponding particle sizes (b and d). (e) TEM image of the Ni film with dispersed ZrO_2 nanoparticles and (f) the corresponding grain size distributions.



ZrO₂ nanoparticles leads to pronounced grain refinement. This grain refinement is attributed to the freshly co-deposited nanoparticles, which suppress the growth of Ni grains along their preferred growth directions and/or promote the nucleation of new grains.

3.2 SEM/EDS surface morphologies of as-prepared Ni film with dispersed ZrO₂ nanoparticles

Fig. 3 displays the surface SEM/EDS results of the as-prepared Ni nanocomposite films with various concentrations of dispersed ZrO₂ nanoparticles. The surface microstructure of the Ni film with dispersed ZrO₂ with a low content of approximately 1 wt% deposited on the Al alloy is shown in Fig. 3(a). The corresponding high-magnification image clearly reveals the dispersion of ZrO₂ nanoparticles, as indicated by the yellow arrows in Fig. 3(b).

The EDS analysis taken from the surface region marked by the purple square in Fig. 3(a), which confirms the presence of approximately 1 wt% ZrO₂ nanoparticles, is shown in the EDS spectrum in Fig. 3(c). The elemental composition was

quantified using the INCA software attached to the SEM/EDS system. Multiple samples were prepared and examined by SEM/EDS, and appropriate samples were selected for further experiments. The particle concentration was increased to 3.5 wt%, and the resultant surface microstructure of the Ni nanocomposite film is shown in Fig. 3(d). The high-magnification image (Fig. 3(e)) confirms the uniform dispersion of ZrO₂ nanoparticles, as indicated by yellow arrows. The increased nanoparticle content is clearly reflected in the surface microstructure. The corresponding SEM/EDS spectrum further confirms the wt% of ZrO₂ nanoparticles in the nanocomposite film, as shown in Fig. 3(f). The ZrO₂ nanoparticle content was further increased from 3.5 wt% to 5.6 wt%, and the resultant surface morphologies and EDS results are presented in Fig. 3(g-i). The surface microstructure clearly indicates a higher concentration of nanoparticles, as highlighted by yellow arrows in Fig. 3(h). At this higher wt%, a small degree of particle agglomeration can also be observed. The EDS spectrum confirms the elemental wt% in the nanocomposite film, as shown in Fig. 3(i). The results indicate the successful and uniform dispersion of ZrO₂ nanoparticles at various

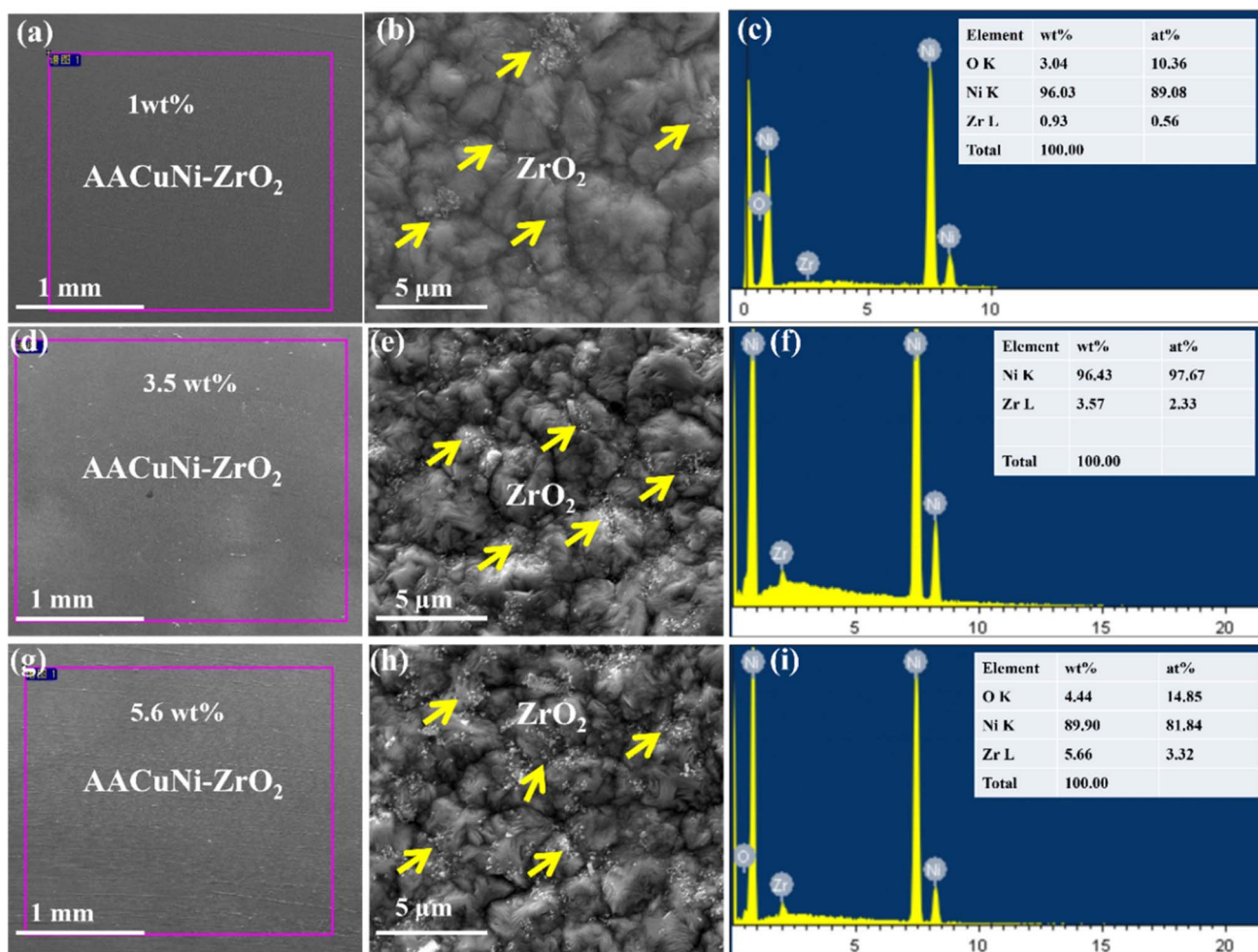


Fig. 3 Surface morphologies and the corresponding EDS results of as-prepared Ni-ZrO₂ nanocomposite coatings deposited on the AA2219 alloy: (a, d, and g) SEM images under low magnification and (b, e, and h) SEM images under high magnification of ZrO₂ particle concentrations of 1 wt%, 3.5 wt%, and 5.6 wt%, respectively; and (c, f, and i) the corresponding EDS spectra.

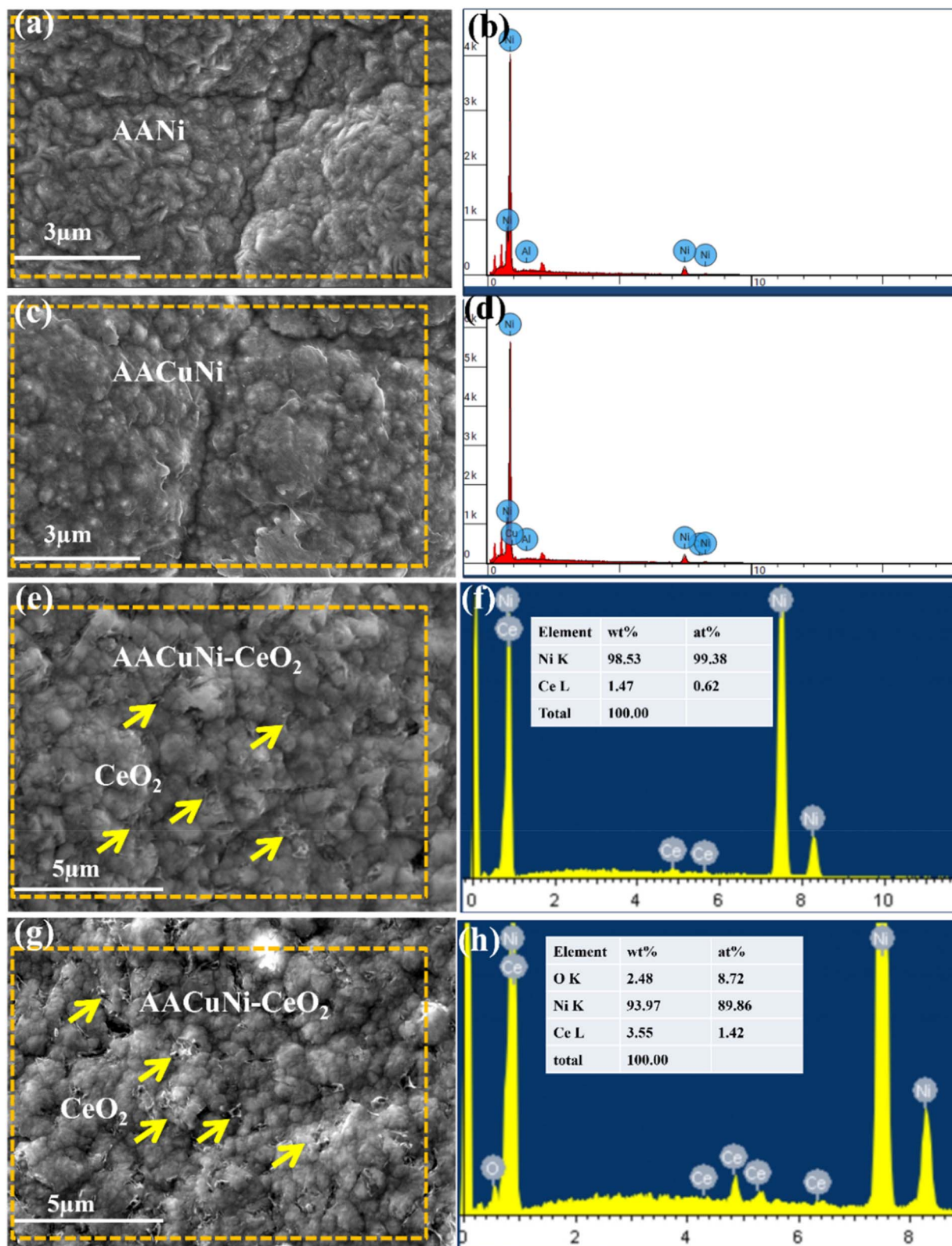


Fig. 4 Surface microstructures and the corresponding EDS results of the as-prepared coatings on the Al alloy. Surface morphologies of (a) AA Ni coating, (c) AACuNi, and (e and g) AACuNi-CeO₂ nanoparticle-dispersed coatings. (b), (d), (f) and (h) present the corresponding EDS spectra, confirming the elemental composition of the coatings.



concentrations (1 wt%, 3.5 wt%, and 5.6 wt%) in the nanocrystalline Ni film, without cavities, pores, or cracks.

3.3 Microstructures and the corresponding EDS results of the as-prepared Ni, Cu–Ni, and CeO₂-dispersed coatings on Al alloy

Fig. 4 illustrates the surface microstructures and the corresponding EDS results of coatings prepared on the Al alloy using composite electroplating. The AANi and AANiCu samples shown in Fig. 4(a–c) present clear surface morphologies in the magnified images. The corresponding EDS results are shown in Fig. 4(b–d). Fig. 4(e–g) show the surface morphologies of AACuNi coatings with dispersed CeO₂ nanoparticles at concentrations of 1.4 wt% and 3.5 wt%, respectively. The corresponding SEM/EDS analyses confirm the nanoparticle concentrations, as shown in Fig. 4(f–h). The surface microstructures clearly evidence the presence of CeO₂ nanoparticles, as indicated by yellow arrows in Fig. 4. It is also evident from the surface morphologies that the coatings with the CeO₂ nanoparticle dispersion exhibit a more compact structure with finer grain morphologies compared to the coatings without the nanoparticle dispersion. This observation is consistent with the TEM results shown in Fig. 2(e), where a smaller grain size of approximately 50 nm is observed. In our previous work, the grain size of CeO₂-dispersed coatings was reported to be around 61 nm, compared to approximately 100 nm for the pure Ni film reported in our recent study.^{29,32}

Generally, it is anticipated that combining a dense nickel matrix with particle reinforcement (ZrO₂ or CeO₂) will increase corrosion resistance primarily through enhancing route flexibility for corrosive species, introducing chemically stable inclusions that prevent ion transport and fine grains promoting uniform passive-film formation. The presence and uniform dispersion of Ce, Zr, and O elements inside the Ni matrix are confirmed by the SEM/EDS spectra of Ni-CeO₂ and Ni-ZrO₂ coatings, as shown in Fig. 3 and 4. In addition to improving the coating compactness, the addition of CeO₂ and ZrO₂ offers unique corrosion protective mechanisms, as CeO₂ provides active corrosion inhibition through Ce³⁺/Ce⁴⁺ redox transitions and as an oxide film promoter. Meanwhile, ZrO₂ acts as a stable, inert barrier phase that prevents corrosive species from diffusing. Therefore, when compared to pure Ni coatings, the results demonstrate that both Ni-CeO₂ and Ni-ZrO₂ composite coatings exhibit better structural integrity and a higher potential for corrosion resistance. The nanoparticle dispersion improves enhanced grain refinement, which increases the surface to volume ratio. The resulting top passive layer of NiO and Ni(OH)₂ (ref. 33 and 34) forms a smooth surface suitable for various industrial applications. The crack-free coatings prevent the penetration of corrosive ions in capillaries/pores/cracks, which enhances the anticorrosive protection performance of materials.

3.4 Micro-hardness

Surface modification by nickel electrodeposition provides significant enhancement in mechanical properties, particularly

microhardness, for demanding structural applications to improve the performance, service life, and corrosion resistance of components. These nanocomposite coatings offer superior corrosion resistance and excellent mechanical performance through the incorporation of hard ceramic nanoparticles.³⁵ In this study, we focus on the effects of incorporating different wt% zirconia (ZrO₂) and ceria (CeO₂) nanoparticles in nickel electroplating on AA2219 substrates. Microhardness of the Ni nanocomposite coating depends on various factors like strengthening the interface between Ni nanoparticles, grain refinement, nanoparticle distribution, reinforcement effect, and substrate/coating adhesion. These hard, non-deformable nanoparticles are uniformly distributed within the FCC Ni matrix, forming a compact coating (see Fig. 2(e), 3, and 4). Their presence affect the microstructure of the Ni coatings, which restrict grain growth and lead to grain refinement, as confirmed by TEM (Fig. 2(e)). This refinement significantly enhances hardness *via* the Hall–Petch effect. Both these nanoparticles act as nucleation sites during coating and provide a more compact and nanoscale structure,^{36,37} thus blocking the dislocation motion and plastic deformation in the nickel matrix, which is explained by the Orowan strengthening process and the primary agents of plastic deformation in crystalline materials have to bend around unsharable nanoparticles.³⁸ A uniform dispersion is achieved by controlled magnetic stirring and ultrasonic agitation. Grain refinement enhances the hardness of nanocomposite coatings. High current density decreases coating adhesion; therefore, the adhesion is the highest at 1.5 A dm⁻², as confirmed in the literature.^{29,39} During electrodeposition, a nanoparticle content of 3.5 wt% showed a significant improvement in mechanical properties, hardness and corrosion performance of coatings. Zirconia nanoparticles are inert, hard particles that promote a more compact coating structure. The resulting grain refinement, as shown in Fig. 2 based on the TEM results, leads to changes in surface morphology and increases surface density, which is one of the reasons for the higher

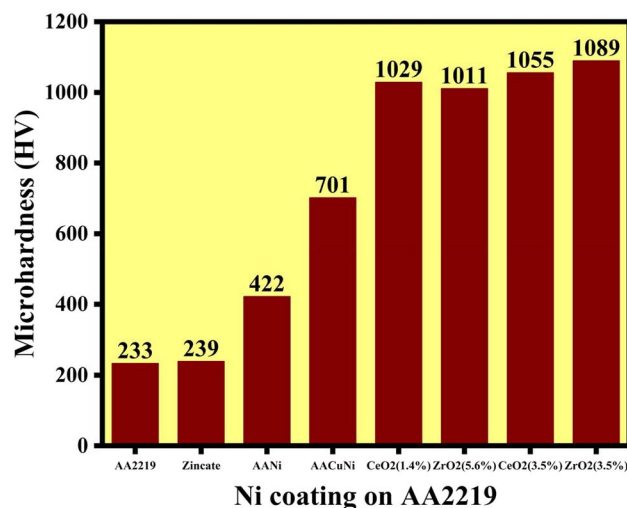


Fig. 5 Microhardness plot of different wt% of CeO₂ and ZrO₂ nanoparticles in Ni coatings on the AA2219 aluminum alloy.

hardness of the zirconia-containing coatings compared with ceria-containing coatings. In this study, the electrodeposition of ZrO₂ led to a substantial increase in the hardness of the Ni coating, from 422 ± 8 HV to 1089 ± 9 HV, demonstrating a remarkable enhancement in the hardness of Ni electrodeposited on the aluminum alloy, as shown in Fig. 5. Similarly, the incorporation of CeO₂ also resulted in a significant hardness improvement, reaching 1055 ± 11 HV, which is only slightly lower than that achieved with the dispersion of ZrO₂ nanoparticles. This optimum nanoparticle content (~3.5 wt%) shows a beneficial effect and prevents the nanoparticles from agglomeration, which leads to the degradation in mechanical properties at higher and much lower loadings, as shown in Fig. 5. When used individually, both ceramic nanoparticles produced pronounced hardness enhancement, along with improved corrosion resistance. In contrast, most reported studies show comparatively lower improvements in hardness, even with the addition of other hardening elements such as Co or W.⁴⁰ The improved adhesion of successive coatings is achieved through the double zincating (DZ) process, which includes nitric acid etching and generates a micro/nanoporous Al surface with finer Zn nucleation, as reported previously (Table 3). The electroplated Cu interlayer promotes the formation of CuO/ZnO phases that suppress Al₂O₃ formation and act as a compact metallic barrier, thereby reducing charge transfer, ion transport, and oxygen permeability.⁴¹ Although the protective Cu oxide layer may partially dissolve under high chloride conditions, the Cu interlayer still enhances hardness, provides a crack-barrier effect, and improves long-term mechanical stability when the Ni layer is damaged. However, excessive nanoparticle agglomeration can degrade coating integrity and reduce the hardness of coatings.^{42,43}

3.5 Electrochemical characterization

Electrochemistry plays important role in various fields, including batteries, water splitting, corrosion studies and fuel cells. Material durability and anticorrosion properties are determined by oxidation-reduction redox reactions on the metal

surface.⁴⁴ Electrochemical behavior of Ni nanocomposite coatings, without and with nanoparticles (ceria and zirconia) on AA2219, is investigated in a 3.5% NaCl solution to study the corrosion resistance under corrosive and harsh environment conditions.

3.5.1 Potentiodynamic polarization. The Tafel extrapolation method was utilized to examine the electrochemical measurements, as shown in Table 4. AA2219 is inherently prone to corrosion, and the electrodeposition of nickel nanocomposites provides microstructural refinement and densification, establishing a robust physical barrier that significantly enhances corrosion resistance. The corrosion potential (E_{corr}) represents the equilibrium potential at which the material neither corrodes nor passivates.⁴⁵ The Ni-CeO₂ coating exhibits a significant increase in corrosion potential ($E_{\text{corr}} = -133$ mV) compared to pure Ni ($E_{\text{corr}} = -509$ mV) and Ni-ZrO₂ ($E_{\text{corr}} = -368$ mV) coatings, shifting to a more positive value, as shown in Fig. 6(a). The observed shift signifies a reduced thermodynamic susceptibility to corrosion, while the intermediate copper layer significantly enhances the overall corrosion resistance. All the coated samples demonstrate high corrosion resistance because of low corrosion current density (I_{corr}), which reflects the rate of the electrochemical reaction due to the addition of ceria nanoparticles, reducing I_{corr} from 14.70 μA to 1.80 μA. The R_p is inversely proportional to I_{corr} , and a higher R_p indicates better anticorrosion properties.⁴⁶ The Ni-CeO₂ sample demonstrated enhanced protective performance, characterized by the lowest I_{corr} and highest polarization resistance. As shown in Fig. 6(a), Ni-CeO₂ and Ni-ZrO₂ exhibit superior resistance to localized corrosion, characterized by high pitting potentials and stable I_{corr} .⁴⁷ A passive film, as shown in Fig. 7(e-h), enhances the stability of the passive film and provides a robust barrier against chloride corrosive species, thereby minimizing the susceptibility to pitting corrosion; the homogeneous distribution of nanoparticles mitigates microstructural defects and obstructs aggressive ion transport, facilitating stable passivation and superior corrosion resistance.⁴⁸ The optimum concentration of 3.5 wt% for both nanoparticles prevents agglomeration, resulting in an efficient crystalline structure and

Table 3 Microhardness values of different wt% of the CeO₂ and ZrO₂ nanoparticles in Ni coating on the AA2219 aluminum alloy

Sample	AA2219	Zincate	AANi	AACuNi	CeO ₂ (1.4%)	ZrO ₂ (5.6%)	CeO ₂ (3.5%)	ZrO ₂ (3.5%)
Hardness (HV)	233 ± 4	239 ± 4	422 ± 8	701 ± 12	1029 ± 16	1011 ± 14	1055 ± 11	1089 ± 9

Table 4 Tafel parameters obtained by the fitting of the Tafel curves of the Ni nanocomposite samples

Sample	Beta A mV per decade	Beta C mV per decade	I_{corr} (μA)	E_{corr} (mV)	Corrosion rate (mpy)	Resistance polarization (KΩ)	Corrosion efficiency (η_p %)	Chi squared (χ^2)
AA bare	74.8	209	200.0	-991.0	57.46	0.12		0.12
Double zincate	94.9	253.7	165.0	-1080	14.21	0.18	17.5	0.1
AACuNi	166.5	209.1	14.70	-509.0	7.2	2.73	92.65	0.05
AACuNi-ZrO ₂ (3.5 wt%)	222.0	639.6	5.600	-368.0	2.228	13.05	97.2	0.01
AACuNi-ZrO ₂ (5.6 wt%)	80.0	87.7	26.50	-432.0	10.56	1.29	86.75	0.026
AACuNi-CeO ₂ (1.4 wt%)	555	250.6	3.27	-275	1.303	1.88	98.3	0.049
AACuNi-CeO ₂ (3.5 wt%)	471.2	272.1	1.08	-133.0	0.431	69.35	99.4	0.08



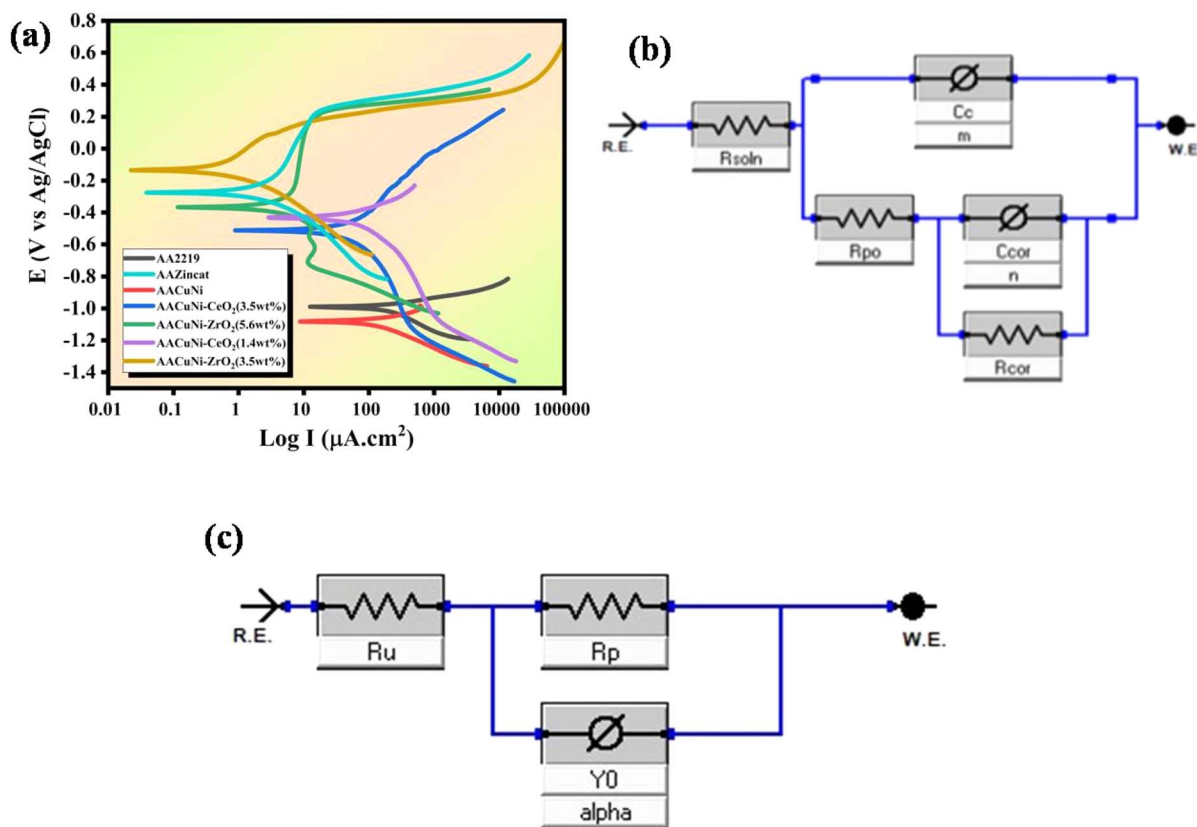


Fig. 6 (a) Tafel curves of the Ni-coated samples with different wt% of the CeO_2 and ZrO_2 nanoparticles. (b) Equivalent reape2cpe circuit model. (c) Equivalent cpe circuit model.

superior corrosion resistance, as verified by hardness and EIS studies. While the Ni- CeO_2 and Ni- ZrO_2 coatings with 1.4 wt% and 5.6 wt% nanoparticles, respectively, show lower corrosion current density, but lower than that of 3.5 wt% due to the agglomeration of high concentration nanoparticles, indicating good anti-corrosion characteristics in a 3.5 wt% NaCl solution at 25 °C.

3.5.2 Surface microstructure analysis after corrosion in 3.5 wt% NaCl solution. Fig. 7 shows the surface morphology of Ni films without and with nanoparticle dispersion after corrosion on an Al alloy in a 3.5 wt% NaCl environment. The AANi coatings exhibit severe corrosion, characterized by corrosion pits, microcracks (indicated by yellow arrows), and a non-uniform corrosion product layer, as shown in Fig. 7(a). Furthermore, it is confirmed that the formed oxide or hydroxide film is loosely adhered to the coating surface. The magnified image more clearly shows the significant growth of corrosion products and microcracks, along with the formation of an oxide film (NiO) layer, as presented in Fig. 7(b). This behavior is attributed to galvanic coupling between Ni and Al and the easy penetration of chloride ions through microcracks and pores, leading to unstable Ni passive films for Ni coatings on Al substrates in NaCl environments. The introduction of a Cu interlayer significantly improves corrosion resistance⁴⁹ by reducing the galvanic mismatch between Al and Ni and enhancing the formation of a more adherent corrosion film, as shown in Fig. 7(c), in comparison with the coating without a Cu

layer (Fig. 7(a)). However, cracks and corrosion product layers are still observed in the chloride environment. In such conditions, Cu oxides/hydroxides may partially dissolve, allowing localized corrosion to initiate at coating defects in the 3.5 wt% NaCl solution.

The Ni film with dispersed CeO_2 nanoparticles shows further improvement. CeO_2 nanoparticles provides active corrosion inhibition through the $\text{Ce}^{3+}/\text{Ce}^{4+}$ redox couple, promoting re-passivation and suppressing cathodic oxygen reduction.^{50,51} The surface passivation layer becomes smoother and more adherent, with only a few localized pits, as indicated by the yellow arrow in Fig. 7(e). These minor localized pits may be associated with nanoparticle agglomeration, which can be observed more clearly in the magnified image shown in Fig. 7(f). The AACuNi- ZrO_2 coatings exhibit the best corrosion resistance, characterized by a more compact and adherent oxide layer (Fig. 7(g)) and suppressed crack formation. ZrO_2 is chemically inert and does not participate in redox reactions; instead, it provides a strong physical barrier protection. The dispersion of ZrO_2 nanoparticles causes significant grain refinement of the Ni matrix, which enhances passivity, reduces defect size, and promotes the formation of a uniform and stable NiO/Ni(OH)₂ film. Fine grains also shorten diffusion paths for re-passivation, effectively preventing chloride penetration into the substrate and contributing to the formation of a thicker and more protective surface layer, as clearly observed in the magnified



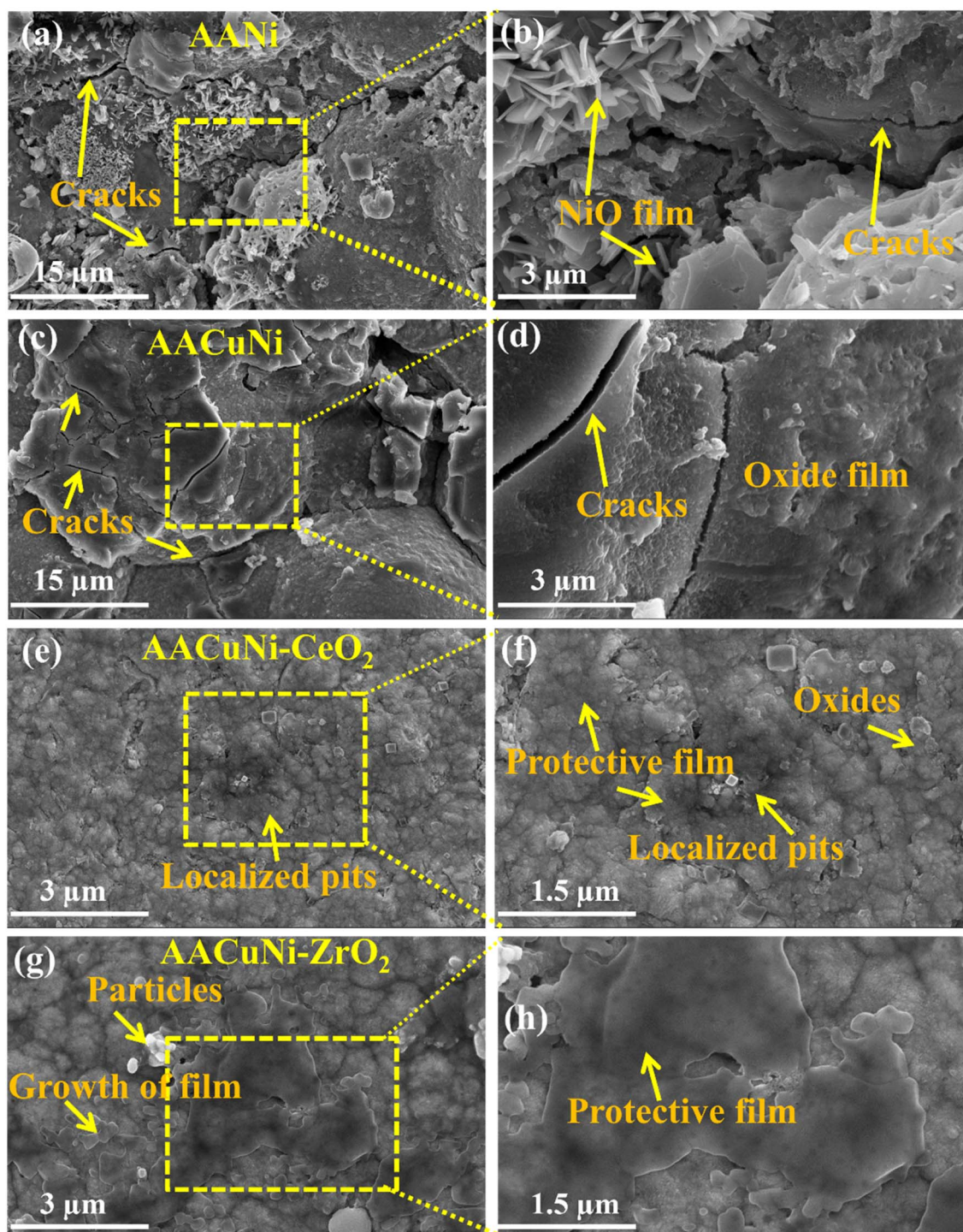


Fig. 7 (a), (c), (e), and (g) SEM surface morphologies of the Ni film without and with dispersed CeO_2 and ZrO_2 nanoparticles on the AA2219 alloy, after corrosion in 3.5% NaCl solution. (b), (d), (f), and (h) Corresponding magnified images.

image (Fig. 7(h)) and presented in Fig. 9, as a schematic of the corrosion mechanism.

3.5.3 Electrochemical impedance spectroscopy (EIS). Electrochemical impedance spectroscopy (EIS) is used to evaluate corrosion resistance by analyzing the coating–electrolyte interface. The key parameters include R_s (electrolyte resistance), R_{ct} (charge transfer resistance), R_c (coating resistance), C_{dl} (double

layer capacitance), and C_c (coating capacitance), which reflects the dielectric properties of the coating layer. Higher R_{ct} suggests slow corrosion, while lower C_c indicates a coating with low water absorption, dielectric constant, and high barrier characteristics. For marine and aerospace corrosion resistance, immersion tests in 3.5 wt% NaCl are utilized. Impedance modulus indicates



corrosion resistance; larger semicircles or longer arcs signify high corrosion resistance.

An equivalent circuit fitting model (reap2cpe) for Ni films with and without the nanoparticles, as shown in Fig. 6(b), and an equivalent circuit fitting model (cpe) for the bare sample, as shown in Fig. 6(c), were used to analyze the EIS data. Also, the data is well-fitted, as shown in Fig. 8. As demonstrated in Fig. 6(a), adding CeO₂ to the protective coating significantly increases the corrosion resistance. The formation of a passive stable layer on the Ni coating, where ceria works similarly to what is stated in the Tafel plot, thereby limiting anodic dissolution and Cl⁻ ion entry, is one of the microstructural and morphological factors responsible for the improvement in stability.^{45,50,52} FESEM images verify that ceria and zirconia nanoparticles seal surface pores, forming a homogeneous, dense structure that prevents corrosive agents and enhances the anticorrosive properties of the coating. Table 5 shows that Ni coating has a film resistance (R_{po}) of 0.3 KΩ. With the increase of both ceria and zirconia contents, R_{po} increases from 0.301 KΩ to 5.03 KΩ, demonstrating that ceria and zirconia nanoparticles create a compact, uniform layer. This dense coating prevents

water, oxygen, and corrosive ions from entering the metal substrate. The R_{po} values are confirmed by comparable results for R_{cor} , which also increased from 0.3 KΩ to 25.1 KΩ, while the C_{dl} and C_c values decreased. Ni-CeO₂ and Ni-ZrO₂ with the optimal concentration of 3.5 wt% improve the coating's anti-corrosive capabilities. Ceria nanoparticles provide a strong nickel oxide layer that prevents Cl⁻ ions.⁵³ The n and m values are close to 0.9, showing a highly uniform coating, which is verified by the FESEM images. The Bode plot shows the phase angle (ϕ) and impedance magnitude (Z) in relation to the frequency of the AC signal (f), as shown in Fig. 8(c) and (d).

A strong correlation exists between corrosion resistance and the height of the curve (Z) across the frequency range. This plateau value is proportional to polarization resistance (R_p), which is inversely related to the corrosion rate. At mid-frequency, the line slope is related to the capacitive behaviour of the protective layer.⁵⁴ Higher impedance, especially at low frequencies, indicated improved corrosion resistance. The zincate curve, which is the least resistant to corrosion, has the lowest impedance at low frequencies, whereas the Ni-CeO₂ curve has the highest impedance compared to Ni-ZrO₂. As

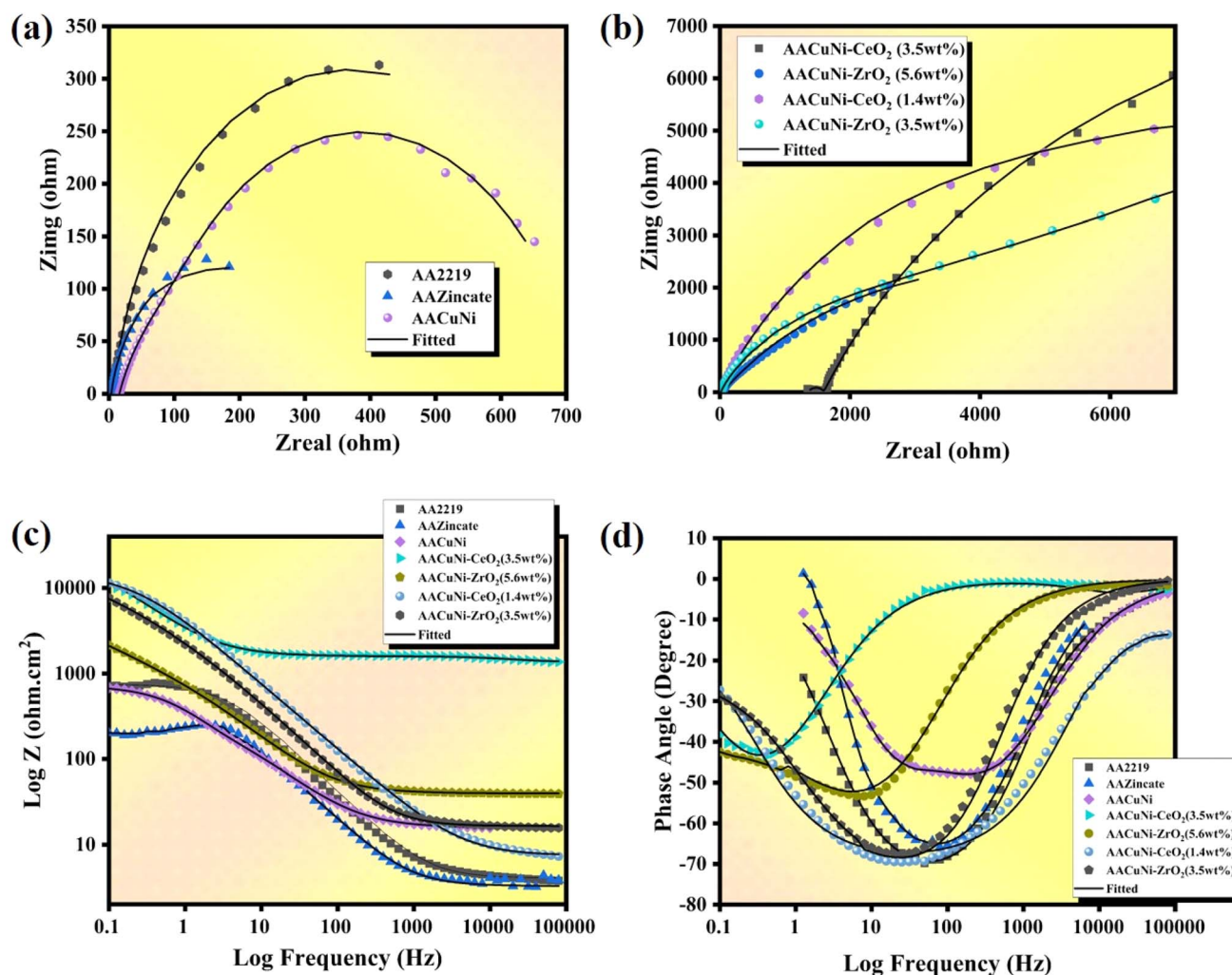


Fig. 8 (a) Nyquist plot of the Ni film without nanoparticles. (b) Nyquist plot of Ni-CeO₂ and Ni-ZrO₂ with different wt%. (c) Impedance plot of the Ni nanocomposite. (d) Phase angle plot of the Ni Nanocomposite.



Table 5 EIS parameters obtained by the fitting of EIS curves of the Ni nanocomposite samples

Sample	R_{soln} (k Ω)	R_{cor} (k Ω)	R_{po} (k Ω)	C_{cor} (S s n_1)	n	C_c (S s n_2)	m	Thickness (mm)	Goodness of fit (χ^2)
AA bare	4.575		0.555	175.5×10^{-6}	0.95			2 ± 0.2	0.008
AA zincate	3.353	0.164	0.301	4.001×10^{-6}	0.8	206.5×10^{-6}	0.856	28 ± 0.4	0.004
AACuNi	15.76	0.386	0.328	263.2×10^{-6}	0.9	381.3×10^{-6}	0.755	53 ± 0.5	0.00025
AACuNi-ZrO $_2$ (3.5 wt%)	16.20	19.1	2.72	105.2×10^{-6}	0.5	70.34×10^{-6}	0.833	22 ± 0.24	0.012
AACuNi-CeO $_2$ (1.4 wt%)	7.45	20.8	4.1	42.5×10^{-6}	0.8	52.16×10^{-6}	0.775	21 ± 0.12	0.004
AACuNi-ZrO $_2$ (5.6 wt%)	19.1	10.13	1.1	565.5×10^{-6}	0.52	237.1×10^{-6}	0.768	26 ± 0.2	0.001
AACuNi-CeO $_2$ (3.5 wt%)	15.2	25.1	5.03	38.47×10^{-6}	0.88	84.73×10^{-6}	0.723	24 ± 0.1	0.00056

Table 6 Comparative electrochemical performance of Ni-based nanocomposite coatings

Coating system	Substrate	Electrolyte	$R_{\text{ct}}/R_{\text{cor}}$ (k Ω)	$R_{\text{po}}/\text{film}$ resistance (k Ω)	I_{corr} (μA)	Year
Ni-ceramic nanocomposite	Al alloy	3.5 wt% NaCl	8–15	N/R	3–10	2023
Polymer/ceramic composite	Steel	NaCl	5–12	1–3	5–20	2023
Ni-based nanocomposite	Al alloy	3.5 wt% NaCl	10–18	2–4	2–6	2023
Ni-CeO $_2$ composite	Steel	NaCl	12–20	3–4	2–5	2024
Oxide-reinforced metal coating	Al alloy	NaCl	6–14	1–3	4–12	2024
AACuNi-ZrO $_2$ (3.5 wt%)	AA2219	3.5 wt% NaCl	19.1	2.72	5.60	This work
AACuNi-CeO $_2$ (3.5 wt%)	AA2219	3.5 wt% NaCl	25.1	5.03	1.08	This work

shown in Fig. 8(d), the Ni nanocomposite curves exhibit a more complex pattern, suggesting two overlapping time constants, and the downward shifts in phase angle indicate coating degradation due to the penetration of corrosive species.⁵⁵ Ni-CeO $_2$ with 3.5 wt% nanoparticles exhibits a lower and broader phase angle peak around -50° at low frequencies. A large peak at a low frequency with a minimal phase shift typically signifies a highly stable barrier layer. The peak shifts in Ni-CeO $_2$ to lower frequencies suggest an increased corrosion resistance. This EIS result confirms that both CeO $_2$ and ZrO $_2$ (3.5 wt%), along with the Ni coating, exhibit superior anticorrosive performance on AA2219, resembling the Tafel results.

Table 6 systematically compares the electrochemical performance parameters, including charge-transfer resistance ($R_{\text{ct}}/R_{\text{cor}}$), coating resistance (R_{po}), corrosion current density (I_{corr}), and test environment, of Ni-CeO $_2$ and Ni-ZrO $_2$ nanocomposite coatings in the present work with the previously reported nanocomposite coatings from the literature.^{56,57} The comparison clearly demonstrates that AACuNi-CeO $_2$ and AACuNi-ZrO $_2$ (3.5 wt%) coatings developed in this work exhibit high charge-transfer resistance and coating resistance among the Ni-based nanocomposite coatings evaluated in a 3.5 wt% NaCl solution, as discussed in detail above and in the discussion section. The high charge-transfer and coating resistance values indicate the formation of a highly effective barrier layer, resulting in superior corrosion protection.

4 Discussion

4.1 Effect of uniform nanoparticle dispersion and optimized wt% on the corrosion resistance of nanocomposite films

The present work clearly shows that the optimal concentration with uniform dispersion of CeO $_2$ and ZrO $_2$ nanoparticles

(around 3.5 wt%) significantly enhances the hardness and corrosion resistance of the Ni coating on AA2219 alloy. The uniform dispersion of nanoparticles is important because homogeneous distribution of CeO $_2$ and ZrO $_2$ nanoparticles hinders the movement of dislocations through the Orowan mechanism.⁵⁸ The SEM/EDS results confirmed the uniform distribution of both nanoparticles within the Ni matrix without causing agglomeration, porosity, cracks, or cavities. These findings were further confirmed by TEM investigations. Furthermore, uniform dispersion results in a more compact and dense coating, which effectively inhibits the pathways for electrolyte (Cl $^-$) penetration.⁵⁹ This observation is also supported by electrochemical spectroscopy. The Tafel plot indicates a significant decrease in the corrosion current density (I_{corr}), and a positive change in the corrosion potential (E_{corr}) of both Ni-CeO $_2$ and Ni-ZrO $_2$ wt% coatings compared to the Ni coating without the dispersion of nanoparticles. Specifically, the Ni-CeO $_2$ coating with 3.5 wt% content exhibits the lowest I_{corr} and the highest polarization resistance, yielding the most effective corrosion resistance. The EIS results, in which a drastic rise in the resistance of charge transfer (R_{cor}) and coating resistance (R_{po}) accompanied by a fall in coating capacitance, prove the barrier layer formation as being quite an effective anticorrosive layer. CeO $_2$ nanoparticles prevent corrosion through a reversible Ce $^{3+}/\text{Ce}^{2+}$ redox pair.⁵⁰ As a result of the dissolution of Ce $^{3+}$ ions, CeO $_2$ can be partially reduced to Ce $_2\text{O}_3$. When these cerium species migrate to the substrate surface, they are re-oxidized to Ce $^{4+}$ due to their strong affinity for oxygen. This process promotes the formation of a stable cerium oxide layer on the surface, which acts as a protective barrier and enhances the anticorrosion performance of the coating.⁶⁰ The schematic of this mechanism is shown in Fig. 9. The reversible Ce $^{3+}/\text{Ce}^{4+}$ redox couple enables rapid re-passivation and self-healing



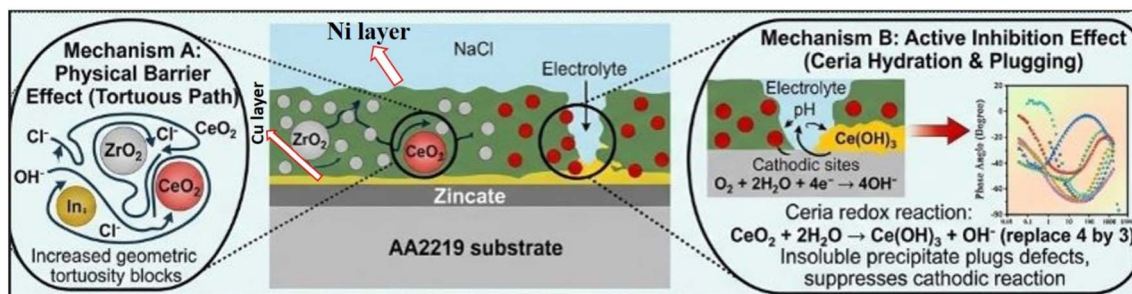


Fig. 9 Schematic of the corrosion mechanism of the Ni coatings with dispersed ZrO_2 and CeO_2 nanoparticles on the AA2219 alloy.

behavior.⁵⁰ Due to this redox activity, ceria offers more effective corrosion inhibition than zirconia. ZrO_2 nanoparticles provide a high anti-corrosivity because of blocking the pathways of diffusion of chloride ions, as well as retaining the passive $\text{NiO}/\text{Ni}(\text{OH})_2$ surface, and make it a chemically inert and physically strong barrier phase. The post-corrosion SEM images provide clear evidence that both nanoparticle-reinforced coatings exhibit significantly enhanced surface protection and fewer corrosion pits and microcracks compared to the pure Ni coating.

4.2 Grain refinement enhanced corrosion resistance and hardness

The CeO_2 and ZrO_2 nanoparticle dispersions act as heterogeneous nucleation sites in electrodeposition, thereby leading to increased coating density, as well as significantly enhanced grain refinement. This grain refinement reduces internal defects and microcracks that normally improve the corrosion resistance. By successfully blocking the diffusion pathways of aggressive species like Cl^- ions, the nanoparticles increase the complexity of corrosion pathways and slow down electrochemical reactions at the coating electrolyte interface.⁵⁰ The TEM analysis shows that upon adding CeO_2 and ZrO_2 nanoparticles, the grain size was significantly refined up to 50 nm, compared to the Ni coating without nanoparticles. This grain refinement is attributed to the pinning effect of the co-deposited nanoparticles, which facilitates the nucleation of new grains because they do not allow grains to grow preferentially.^{61,62} Grain refinement is a mechanical strengthening process that raises microhardness by means of the Hall–Petch strengthening process, in which the fine grains improve the grain boundary density and prevent dislocation motion. Hence, the hardness was significantly increased from 422 HV of pure Ni coating to 1089 HV and 1055 HV of Ni-ZrO_2 and Ni-CeO_2 coatings, respectively.⁶³ The optimal nanoparticle content of 3.5 wt% is a tradeoff between these positive effects and the prevention of agglomeration, which is found to reduce both mechanical integrity and corrosion resistance at higher loadings. This refined grain structure aids in the development of a more uniform, dense and adherent passive layer. Fine grains increase the number of grain boundaries and the grain boundary forms the fast moving diffusion channels of the formation and repair of passive oxide layers. As a result, the $\text{NiO}/\text{Ni}(\text{OH})_2$ film formed on the nanocrystalline coatings is

more uniform and compact. This is clearly reflected not only in the increased impedance at low frequencies in the EIS results, but also in the post-corrosion SEM images, where the nanoparticle-reinforced surfaces appear smoother and more continuous. These findings demonstrate that a controlled dispersion with an optimized concentration of approximately 3.5 wt% CeO_2 and ZrO_2 nanoparticles enables the formation of a dense, nanocrystalline Ni-based coating with enhanced mechanical strength and corrosion resistance on the AA2219 alloy. These improvements are directly associated with the grain refinement down to ~ 50 nm, dispersion strengthening, and the formation of a stable passive film in both Ni-CeO_2 and Ni-ZrO_2 coatings. Moreover, the Ni-CeO_2 coating provides additional active corrosion inhibition due to the redox behavior of ceria, resulting in superior corrosion resistance. These findings offer a clear mechanism-based strategy for designing high-performance Ni-based nanocomposite coatings for aerospace and marine applications.

5 Conclusions

In this study, Ni-CeO_2 and Ni-ZrO_2 nanocomposite coatings were successfully fabricated on AA219 aluminium alloy *via* composite electrodeposition. The key outcomes of this study are as follows.

SEM and TEM analyses confirmed that an optimal concentration (3.5 wt%) and well-controlled homogeneous dispersion of CeO_2 and ZrO_2 nanoparticles yielded dense, uniform coatings, free from pores and cracks. This optimum introduction of nanoparticles resulted in pronounced grain refinement, reducing the Ni grain to about 50 nm, which led to a substantial enhancement in surface hardness up to 1089 HV (158%) and 1055 HV (150%) for Ni-ZrO_2 and Ni-CeO_2 coatings, respectively, compared with pure Ni coating. Electrochemical measurements further demonstrated a remarkable improvement in corrosion resistance, with potentiodynamic polarization results showing a 92.6% reduction in corrosion current density for Ni-CeO_2 while Ni-ZrO_2 exhibited a 61.9% reduction. Consistently, EIS analysis revealed a 65-fold increase in R_{ct} for Ni-CeO_2 , confirming the strong barrier effect provided by the uniformly dispersed nanoparticles.



5.1 Suggestion and future research direction

(1) In future work, the co-deposition of both ceria and zirconia results in high hardness and active inhibition.

(2) Investigating the long-term stability of coatings and different electrolyte concentrations will further validate for different harsh environments.

(3) In electrochemical measurement, the rate of $\text{Ce}(\text{OH})_3$ precipitation provides deep insights into active protection.

Conflicts of interest

The authors declare that there are no conflicts of interest regarding the publication of this paper. All authors have contributed to this work according to the academic and research standards, and there are no competing interests, financial or otherwise, that could have influenced the outcomes of this study.

Data availability

The data supporting this article have been included as part of the manuscript.

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References

- H. Tao, *et al.*, Effect of heat treatment on the microstructure and corrosion performance of 2219 Al alloy cold sprayed coatings, *Corros. Sci.*, 2024, **241**, 112526.
- G. V. A. S. Mideena and A. K. Jhab, Corrosion behavior of aluminium alloy Aa2219– T87 welded plates in sea water, *Indian J. Sci. Technol.*, 2012, **5**(11), 3578–3583.
- Q. Qiao, *et al.*, In-situ monitoring of additive friction stir deposition of AA6061: Effect of rotation speed on the microstructure and mechanical properties, *Mater. Sci. Eng., A*, 2024, **902**, 146620.
- Y. Cui, *et al.*, Grain boundary segregation engineering in high-entropy multiphase alloys for overall water splitting at ultra-high current density, *Appl. Catal. B Environ.*, 2025, 126035.
- M. Margono, *et al.*, Enhanced corrosion resistance of aluminum 6061 alloy using Ti-based thin films and plasma nitriding, *JCIS Open*, 2025, **18**, 100139.
- K. Srinivasa Rao and K. Prasad Rao, Corrosion resistance of AA2219 aluminium alloy: electrochemical polarisation and impedance study, *Mater. Sci. Technol.*, 2006, **22**(1), 97–104.
- Z. Yuan, *et al.*, Microstructure and corrosion behavior of Al-Cu intermetallics in laminated sheets, *Intermetallics*, 2025, **187**, 109022.
- C. S. Witharamage, *et al.*, Corrosion-resistant metallic coatings for aluminum alloys by cold spray, *Corros. Sci.*, 2022, **209**, 110720.
- C. V. Rao, G. M. Reddy and K. S. Rao, Microstructure and pitting corrosion resistance of AA2219 Al-Cu alloy friction stir welds—effect of tool profile, *Def. Technol.*, 2015, **11**(2), 123–131.
- M. S. Safavi and F. C. Walsh, The electrodeposition of particle-metal matrix composite coatings to meet tribological and corrosion challenges, *Adv. Colloid Interface Sci.*, 2025, 103584.
- Z. Liu, *et al.*, Microstructure and cavitation erosion resistance of arc ion plating NiCrAlY coating on the 304L stainless steel, *Tribol. Int.*, 2022, **173**, 107618.
- S. Aruna, V. W. Grips and K. Rajam, Ni-based electrodeposited composite coating exhibiting improved microhardness, corrosion and wear resistance properties, *J. Alloys Compd.*, 2009, **468**(1–2), 546–552.
- J. N. Balaraju, T. S. N. Sankara Narayanan and S. K. Seshadri, Electroless Ni-P composite coatings, *J. Appl. Electrochem.*, 2003, **33**(9), 807–816.
- B. Zhang, *et al.*, Effect of CeO₂ Content on Microstructure and Wear Resistance of Laser-Cladded Ni-Based Composite Coating, *Lubricants*, 2024, **12**, 227.
- F. S. Var, *et al.*, Response surface methodology optimization of electroplating parameters for Co/CeO₂ composite coatings with enhanced corrosion and wear resistance, *J. Mater. Res. Technol.*, 2025, **38**, 4049–4063.
- Y. Wang, *et al.*, The microstructure and mechanical properties of Ni-Co-ZrO₂ nanocomposite coatings, *Int. J. Mod. Phys. B*, 2017, **31**(16–19), 1744022.
- A. Laszczyńska, *et al.*, Electrodeposition and characterization of Ni-Mo-ZrO₂ composite coatings, *Appl. Surf. Sci.*, 2016, **369**, 224–231.
- H. Yan, *et al.*, Study on Properties of Ni-ZrO₂ Nanocomposite Coatings Prepared by Pulsed Electrocasting Under the Coupling Effect of Ultrasonic and Magnetic Field, *JOM*, 2025, **77**, 6781–6791.
- A. Khan, *et al.*, Thermally grown oxide formation on Ni₂Al₃ aluminide coating: The effect of nanocrystalline nickel film on oxide scale adhesion, *Vacuum*, 2022, **197**, 110843.
- Y. Hu, *et al.*, Study on the wettability, intermetallic compound growth, voids formation and mechanical properties of Cu/Sn joints with changes in substrate roughness for electronic packaging, *J. Mater. Sci.: Mater. Electron.*, 2025, **36**(14), 851.
- A. Khiabani, Z.-S. Seyedraoufi and M. H. Sohi, Effect of heat treatment on surface characteristics and corrosion behavior of nickel plated AA2024 aluminum alloy, *Surf. Topography: Metrol. Prop.*, 2021, **9**(3), 035048.
- A. Khan, *et al.*, A Comparative Electrochemical Study of Pt and Ni-Oxide Cathodes: Performance and Economic Viability for Scale-Up Microbial Fuel Cells, *Catalysts*, 2025, **15**(12), 1153.
- A. Ullah, *et al.*, Temperature effect on early oxidation behavior of NiCoCrAlY coatings: Microstructure and phase transformation, *Acta Metall. Sin.*, 2022, **35**(6), 975–984.



- 24 A. Khan, *et al.*, Role of TiO₂ nanoparticles in accelerating α -Al₂O₃ formation and enhancing the oxidation resistance of Ni₂Al₃ coatings, *Ceram. Int.*, 2026, **52**, 8864–8878.
- 25 P. Zhang, *et al.*, Study on the influence of multi-energy field combined surface modification on the subsurface microstructure evolution of AA7075 aluminum alloy, *Mater. Charact.*, 2025, **219**, 114644.
- 26 A. Khan, Z. Dong and X. Peng, Accelerated phase transformation of thermally grown alumina on Ni₂Al₃: Effect of dispersion of hcp-oxides with various content and particle size and chemistry, *Surf. Coat. Technol.*, 2020, **394**, 125861.
- 27 A. Khan, *et al.*, Effect of Cr nanoparticle dispersions with various contents on the oxidation and phase transformation of alumina scale formation on Ni₂Al₃ coating, *Surf. Coat. Technol.*, 2022, **438**, 128397.
- 28 A. Khan, *et al.*, Effect of Cr₂O₃ nanoparticle dispersions on oxidation kinetics and phase transformation of thermally grown alumina on a nickel aluminide coating, *Corros. Sci.*, 2019, **150**, 91–99.
- 29 A. Khan, *et al.*, The effect of grain refinement on the oxidation and phase transformation of alumina scale on Ni₂Al₃ coating, *Intermetallics*, 2022, **146**, 107571.
- 30 A. Khan, *et al.*, ZrO₂-nanoparticle assisted phase transformation and oxidation kinetics of thermally grown alumina on nickel aluminide coatings, *Surf. Coat. Technol.*, 2023, **470**, 129852.
- 31 C. Xu, X. Peng and F. Wang, Cyclic oxidation of an ultrafine-grained and CeO₂-dispersed δ -Ni₂Al₃ coating, *Corros. Sci.*, 2010, **52**(3), 740–747.
- 32 A. Ullah, *et al.*, Effect of vacuum annealing on initial oxidation behavior and alumina transition of NiCoCrAlY coatings, *Surf. Coat. Technol.*, 2020, **404**, 126441.
- 33 X. Wei, *et al.*, Development of growth and thermal stresses in NiO scale on nanocrystalline Ni without and with dispersion of CeO₂ nanoparticles, *Corros. Sci.*, 2017, **118**, 60–68.
- 34 J. Peng, *et al.*, Investigation of process and properties of Cu-Mn-Al alloy cladding deposited on 27SiMn steel via cold metal transfer, *Crystals*, 2025, **15**(10), 858.
- 35 D. G. Pradeep, *et al.*, Study on scratch behavior of Ni-Al₂O₃ coating composition on Al-2219 substrate by electro deposited technique, *Mater. Today: Proc.*, 2021, **46**, 8716–8722.
- 36 I. A. Shozib, *et al.*, Modelling and optimization of microhardness of electroless Ni-P-TiO₂ composite coating based on machine learning approaches and RSM, *J. Mater. Res. Technol.*, 2021, **12**, 1010–1025.
- 37 J. Peng, *et al.*, Effect of the activator B (OCH₃)₃ on the microstructure and mechanical properties of Cu-Mn-Al alloy coating via CMT cladding, *Crystals*, 2025, **15**(10), 881.
- 38 Z. Zhang and D. L. Chen, Consideration of Orowan strengthening effect in particulate-reinforced metal matrix nanocomposites: A model for predicting their yield strength, *Scr. Mater.*, 2006, **54**(7), 1321–1326.
- 39 D. Pradeep, *et al.*, Study on scratch behavior of Ni-Al₂O₃ coating composition on Al-2219 substrate by electro deposited technique, *Mater. Today: Proc.*, 2021, **46**, 8716–8722.
- 40 A. John, A. Saeed and Z. A. Khan, Effect of Alloying and Reinforcing Nanocomposites on the Mechanical, Tribological, and Wettability Properties of Pulse-Electrodeposited Ni Coatings, *Micromachines*, 2025, **16**, 175.
- 41 L. M. Muresan, Nanocomposite coatings for anti-corrosion properties of metallic substrates, *Materials*, 2023, **16**(14), 5092.
- 42 S. Bamberg, *et al.*, The effect of nickel on the strain evolution in chemical copper films, *Thin Solid Films*, 2012, **520**(23), 6935–6941.
- 43 W. Kou, *et al.*, Effect of cold metal transfer mode and arc oscillation on the microstructure and properties of additive-manufactured 2319 aluminum alloy, *Mater. Sci. Technol.*, 2025, 02670836251383115.
- 44 W. Kou, *et al.*, Effects of arc oscillation on interface behavior and properties of additive-manufactured 2319/5356 dissimilar aluminum alloy, *Mater. Today Commun.*, 2025, **47**, 113068.
- 45 Q. Huang, *et al.*, The effect of ZnAl-LDHs-CO₃ on the corrosion behaviour of Zn-5Al alloys in 3.5 wt.% NaCl solution, *Corros. Sci.*, 2021, **179**, 109165.
- 46 D. Dwivedi, K. Lepková and T. Becker, Carbon steel corrosion: a review of key surface properties and characterization methods, *RSC Adv.*, 2017, **7**(8), 4580–4610.
- 47 P. Zhang, *et al.*, Characterization of surface integrity of 7075-T6 aluminum alloy subjected to microbiologically induced corrosion during high-speed machining, *J. Alloys Compd.*, 2024, **1008**, 176843.
- 48 Q. Li, *et al.*, Effect of temperature on the corrosion behavior and corrosion resistance of copper-aluminum laminated composite plate, *Materials*, 2022, **15**(4), 1621.
- 49 J. Du, *et al.*, Influence of a Nb interlayer on the oxidation behavior of a Cr coating deposited on Inconel 718 alloy substrate, *J. Mater. Res. Technol.*, 2025, 9283–9297.
- 50 S. Chen, *et al.*, Microstructure regulation and tribological response mechanism of β -Si₃N₄-modified 304 stainless-steel coatings, *J. Mater. Eng. Perform.*, 2026, 12966.
- 51 S. Ding, *et al.*, Phosphate-Induced Corrosion of Nickel Alloys in Supercritical Water: Mechanistic Insights into Passivation and Accelerated Corrosion at Elevated Temperatures, *J. Supercrit. Fluids*, 2026, 106901.
- 52 Z. Li, *et al.*, Microstructural evolution and corrosion resistance of additively manufactured Ti-6Al-4V alloy annular shaped components using multistage heat treatment, *Mater. Chem. Phys.*, 2025, 131414.
- 53 R. Zhu, *et al.*, Effect of CeO₂ on microstructure and properties of Ni-Co-based coatings, *J. Mater. Res. Technol.*, 2023, **26**, 7329–7339.
- 54 C. Magaña-Zavala, *et al.*, Electrochemical impedance spectroscopy (EIS) modelling of different behaviours of Ni and Ni oxide thin films for corrosion prevention in sour media, *Anti-Corros. Methods Mater.*, 2010, **57**(3), 118–125.
- 55 Z. Bai, *et al.*, A stable anticorrosion coating with multifunctional linkage against seawater corrosion, *Composites, Part B*, 2023, **259**, 110733.



- 56 S.-R. Zhao, *et al.*, Stepwise-Induced Synthesis and Excellent Corrosion Protection of Ce/Eu Codoped ZnO Solid Solution Materials, *ACS Appl. Mater. Interfaces*, 2024, **16**(37), 49533–49543.
- 57 J. R. Xavier and S. P. Vinodhini, Multifunctional nanocomposite coating for aluminium alloy: Corrosion resistance, flame retardancy, and mechanical enhancement for automotive components, *Mater. Chem. Phys.*, 2024, **323**, 129628.
- 58 Z. Zhang and D. Chen, Consideration of Orowan strengthening effect in particulate-reinforced metal matrix nanocomposites: A model for predicting their yield strength, *Scr. Mater.*, 2006, **54**(7), 1321–1326.
- 59 F. Walsh, *et al.*, Electrochemical characterisation of the porosity and corrosion resistance of electrochemically deposited metal coatings, *Surf. Coat. Technol.*, 2008, **202**(21), 5092–5102.
- 60 X. Wang, *et al.*, The reactive element effect of ceria particle dispersion on alumina growth: A model based on microstructural observations, *Sci. Rep.*, 2016, **6**(1), 29593.
- 61 J. Li, *et al.*, Structure and properties of super-hard (MoSiTiVZr) N high-entropy nitride coatings regulated by substrate bias, *Surf. Coat. Technol.*, 2025, **497**, 131788.
- 62 W. Hou, *et al.*, Improvements in surface integrity, corrosion and wear resistances of TA2 titanium alloy via cool-assisted ball burnishing, *Tribol. Int.*, 2025, 111566.
- 63 M. Alizadeh and A. Cheshmpish, Electrodeposition of Ni-Mo/Al₂O₃ nano-composite coatings at various deposition current densities, *Appl. Surf. Sci.*, 2019, **466**, 433–440.

