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# Enhancing Wear Resistance of A390 Aluminum Alloy: A Comprehensive Evaluation of Thermal Sprayed WC, CrC, and Al<sub>2</sub>O<sub>3</sub> Coatings

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**Abstract:** This study comparatively analyzed the wear characteristics and adhesion properties of 86WC–10Co–4Cr (WC) coatings deposited using the high velocity oxygen fuel process and 75Cr<sub>3</sub>C<sub>2</sub>–25NiCr (CrC) and Al<sub>2</sub>O<sub>3</sub>–3TiO<sub>2</sub> (Al<sub>2</sub>O<sub>3</sub>) coatings deposited using the atmospheric plasma spray process on an A390 aluminum alloy substrate. The adhesion strength and wear test results demonstrated that the WC coating exhibited superior wear resistance. In contrast, the CrC and Al<sub>2</sub>O<sub>3</sub> coatings showed lower adhesion properties and unstable frictional variations due to a higher number of defects compared to the WC coating. The WC coating layer, protected by WC particles, exhibited minimal damage and a low wear rate, followed by CrC and Al<sub>2</sub>O<sub>3</sub>. Ultimately, WC coating is highlighted as the optimal choice to enhance the wear resistance of A390 aluminum alloy.

**Keywords:** aluminum alloy; thermal spray coatings; wear resistance; adhesion strength; microstructure



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## 1. Introduction

Aluminum alloys are widely used in the automotive, marine, and aerospace industries due to their high strength, excellent thermal conductivity, and corrosion resistance [1,2]. These properties make aluminum alloys ideal for manufacturing lightweight and fuel-efficient vehicles, as well as for constructing durable marine structures and high-performance aerospace components [3–6]. The inherent light weight of aluminum, combined with its ability to resist corrosion, significantly enhances the lifespan and performance of parts exposed to harsh environments [1–6].

However, the inherently low hardness and poor wear resistance of aluminum alloys restrict their use in extreme environments characterized by high temperatures and friction. For instance, in automotive engines and aerospace turbines, components are subjected to severe mechanical stresses and thermal loads that can lead to rapid wear and tear. This necessitates frequent maintenance and part replacements, thereby increasing operational costs and downtime. Therefore, enhancing the wear resistance of aluminum alloys through surface treatments is essential for their use in high-performance components [4,5].

One of the most effective methods to improve surface properties is surface coating using materials with excellent mechanical properties [7,8]. Thermal spray methods such as flame spray, high velocity oxygen fuel (HVOF), atmospheric plasma arc spray (APS), and wire arc spray (WAS) are primarily used for superior surface protection [8,9]. These methods are applied across various industries by selecting materials that suit the characteristics of each process. Among them, APS and HVOF have been the most widely used due to their ability to generate dense coating layers using plasma at 20,000 degrees Celsius or high-temperature supersonic gas streams. These processes allow for the use of metals and ceramics, providing a wide range of material choices, low porosity, high wear resistance, and strong adhesion [10–16].

APS and HVOF processes typically employ oxides or carbides due to their high melting points. Oxides such as  $\text{Al}_2\text{O}_3$  and carbides like WC and  $\text{Cr}_3\text{C}_2$  are combined with metals like Co, Ni, and Cr to protect the substrate surface. These materials are favored in APS and HVOF coatings due to their low oxygen content in the coating layer, short reaction times resulting in low porosity, and excellent adhesion properties. Consequently, they are widely used in components requiring high wear resistance and corrosion resistance [5,17,18].

Despite ongoing efforts, the inherent limitations of aluminum alloys due to their low surface hardness and wear resistance persist [19,20]. Existing studies have attempted to address these issues through various surface treatment technologies, but most have failed to fully elucidate the wear resistance mechanisms of oxide and carbide coatings. For instance, while methods such as laser cladding can enhance surface hardness and wear resistance, they face issues such as limited application range and difficulty in maintaining consistent quality [21].

To date, there has been little research on the wear resistance mechanisms of ceramic coatings on aluminum surfaces. Therefore, this study comparatively analyzed the wear characteristics of 86WC–10Co–4Cr coatings applied using the HVOF process and 75 $\text{Cr}_3\text{C}_2$ –25NiCr and  $\text{Al}_2\text{O}_3$ –3 $\text{TiO}_2$  coatings applied using the APS process. This study presents coating materials and processes with the potential to significantly enhance the wear resistance of aluminum alloys, providing valuable insights for the development of high-performance components in various industrial applications and demonstrating the potential to improve the durability of the alloys.

## 2. Materials and Methods

The aluminum alloy (A390) substrate was prepared by depositing all coating materials following a solution heat treatment at 525 °C for 4 h and subsequent aging at 190 °C for 6 h. The composition of the alloy was Si (17.0 wt.%), Cu (4.5 wt.%), Mg (0.55 wt.%), Fe (<0.5 wt.%), Ti (<0.2 wt.%), Mn (<0.1 wt.%), Zn (<0.1 wt.%), with the balance being Al.

The coating materials used in the experiment were 86WC–10Co–4Cr, 75 $\text{Cr}_3\text{C}_2$ –25NiCr, and  $\text{Al}_2\text{O}_3$ –3 $\text{TiO}_2$ . The 86WC–10Co–4Cr material was deposited using the high velocity oxygen fuel (HVOF) process, while the 75 $\text{Cr}_3\text{C}_2$ –25NiCr and  $\text{Al}_2\text{O}_3$ –3 $\text{TiO}_2$  materials were deposited using the atmospheric plasma spray (APS) process. The variables for each coating process are listed in Table 1. For convenience, the materials will be referred to as WC, CrC, and  $\text{Al}_2\text{O}_3$ , respectively, throughout this paper. Prior to coating application, the substrate underwent cleaning and blasting for surface roughening. Subsequently, the  $\text{Al}_2\text{O}_3$  material received a bond coating via a buffer layer of Ni–5Al to prevent delamination from the substrate, followed by thermal spraying. The WC and CrC materials were directly thermal sprayed without any bond coating. To ensure a uniform layer, the substrate was rotated during the thermal spraying process, maintaining a consistent thickness.

**Table 1.** Process parameters of spray coating.

Spray Parameters	Unit		
Coating materials	86WC–10Co–4Cr	$\text{Cr}_3\text{C}_2$ –25NiCr	$\text{Al}_2\text{O}_3$ –3 $\text{TiO}_2$
Spray type	HVOF	APS	APS
Buffer layer	-	Ni–5Al	-
Voltage	36 V	60 V	60 V
Current	600 mA	600 mA	600 mA
Air pressure	40 psi	105 psi	105 psi
Spray Distance	8 inch	9 inch	9 inch
Gun spraying angle	90	90	90
Gun/feed/spray rate (g/min)	25	80	80
Thickness	150 $\mu\text{m}$	150 $\mu\text{m}$	100 $\mu\text{m}$
Buffer layer	-	-	Ni-5Al (50 $\mu\text{m}$ )

Specimens were cut and polished from #400 to #2400 grit, followed by polishing with 9, 6, 3, 1, and 0.04  $\mu\text{m}$  diamond suspensions. Microstructural analyses of the cross-sections

and surfaces of the coating layers were conducted using a field emission scanning electron microscope (FE-SEM; CLARA, Tescan, Brno, Czech Republic) and an energy dispersive spectrometer (EDS, EDAX, Pleasanton, CA, USA) with an acceleration voltage of 15.0 kV. The area fraction of cross-sectional defects was measured through image analysis. The adhesion strength between the substrate and coating layers was evaluated according to ASTM C633, with three tests conducted per specimen [22]. The upper cylinder and coating layer were fixed with adhesive, and a biaxial load was applied perpendicularly to the surface to determine the bonding strength. To assess the mechanical properties of each coating layer, a micro Vickers hardness test according to ASTM E384 [23] and a ball-on-disc test according to ASTM G99 [24] were performed. These tests were conducted using an HM-122 hardness test machine (Akashi Co., Tokyo, Japan) with a 0.1 kgf load applied for 10 s. Measurements were taken perpendicular to the coating layers and the substrate. Wear tests were conducted using a room-temperature abrasion tester (R&B, Daejeon, Republic of Korea) under the following conditions: load of 30 N, 100 RPM, 25 °C, and a radius of gyration of 11.5 mm, with Al<sub>2</sub>O<sub>3</sub> as the counterpart material. Each test was repeated three times per sample, and the wear rate (cm<sup>3</sup>/N·m) was calculated from the weight loss after the tests using Equation (1):

$$W = V / (L \times S) \quad (1)$$

where *V* is the wear loss volume (cm<sup>3</sup>) calculated from the cross-sectional area of the wear track, *L* is the load (N), and *S* is the sliding distance (m). Post-test wear track analysis was performed using FE-SEM and a high-resolution 3D microscope (VHX-7000, KEYENCE, Osaka, Japan) to analyze the wear mechanisms.

### 3. Results

Figure 1 shows the cross-sectional microstructure and EDS mapping results of the coating layers and substrates for different materials. In Figure 1a, the WC coating layer exhibits a uniformly distributed Co–Cr binder matrix (grey areas) with rounded WC particles (light grey areas), as revealed by quantitative EDS analysis. The coating layer displays very low porosity and high density due to the high impact velocity of the coating particles. The small WC particles within the coating layer were formed by the fracture of relatively faster and larger particles impacting the substrate during spraying [22]. The CrC coating in Figure 1b displays the typical lamellar structure of plasma-sprayed coatings, consisting of a Ni–Cr binder matrix (grey areas) and Cr carbide (light grey areas). It shows a lamellar structure parallel to the substrate, including porosity and splats. This lamellar structure is a characteristic microstructural feature of arc spray coatings, occurring when the coating material particles do not completely melt before application to the substrate or react with oxygen in the atmosphere to form oxides that mix with unmelted particles [22]. The Al<sub>2</sub>O<sub>3</sub> coating in Figure 1c consists of the coating material and a Ni–5Al buffer layer that includes compounds formed between the coating material and NiAl metal, along with the substrate. The buffer layer shows splat boundaries, and microcracks were observed at the bonding interface with the substrate.

Based on the cross-sectional images, the interfacial cross-sectional defect (porosity and cracks) area fraction analysis using Image J software (<https://imagej.net/ij/>) showed defect area fractions of 0.101 for WC, 0.255 for CrC, and 0.351 for Al<sub>2</sub>O<sub>3</sub>. The WC coating layer exhibited relatively fewer interfacial defects compared to the other coating layers, which is expected to result in the highest bonding strength. This is followed by CrC and then Al<sub>2</sub>O<sub>3</sub> in terms of expected bonding strength.

The actual bonding strength results between the substrate and coating materials are shown in Figure 2. The WC coating, which exhibited the fewest defects at the interface, demonstrated the highest bonding strength at 72.90 MPa. This was followed by CrC at 34.94 MPa and Al<sub>2</sub>O<sub>3</sub> at 17.92 MPa. These results are consistent with the interfacial defect area fraction analysis. The lower bonding strengths of CrC and Al<sub>2</sub>O<sub>3</sub> compared to WC are attributed to the presence of microcracks and splat boundaries in the coating layers and at the interface between the coating materials and the substrate. Lower bonding strength

in a wear environment can lead to reduced durability of the coating, resulting in the separation of the coating layer from the substrate when subjected to forces exceeding the bonding strength. This, in turn, can compromise the protection of the substrate and cause deformation. Therefore, WC is deemed the most suitable coating material for protecting the A390 substrate.

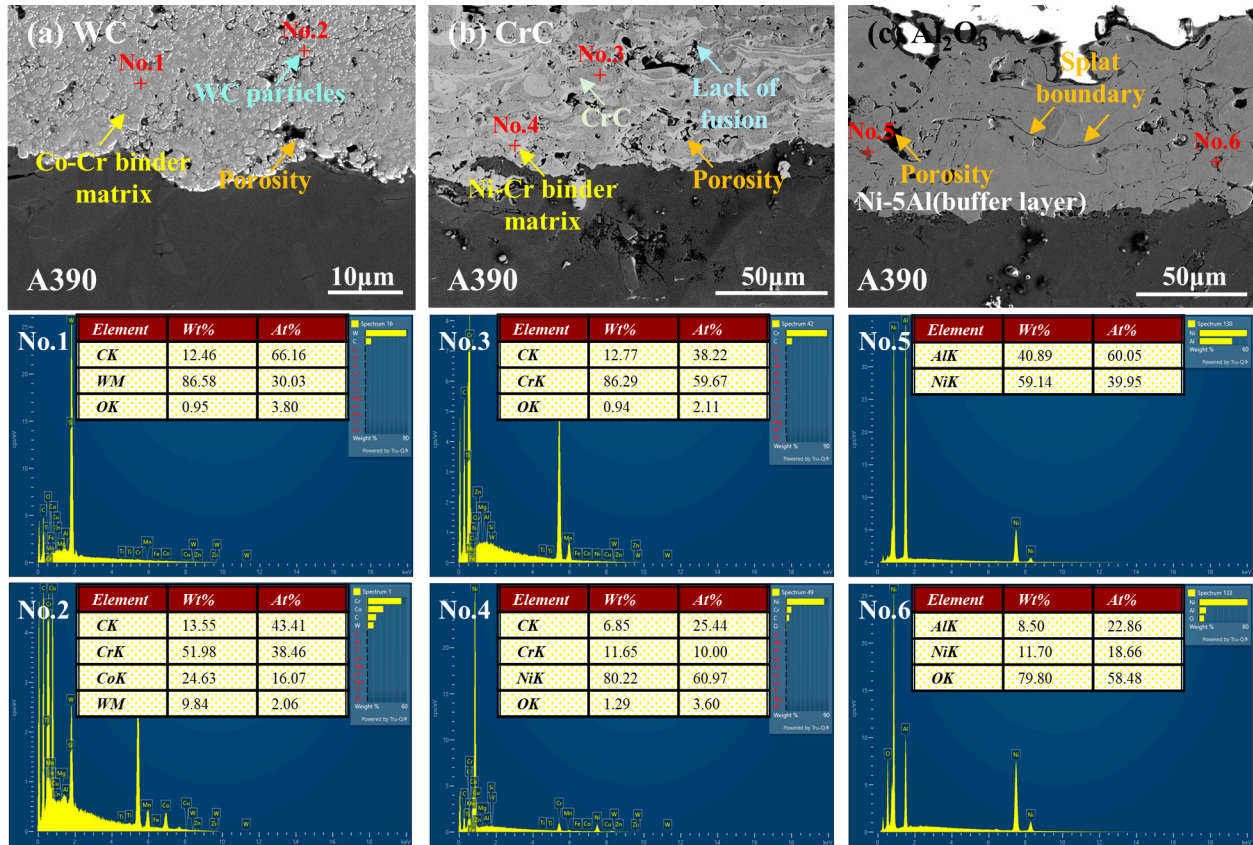


Figure 1. Cross-sectional SEM images of the coatings on A390 substrate: (a) WC, (b) CrC, and (c) Al<sub>2</sub>O<sub>3</sub> coatings, along with EDS qualitative analysis results of the points marked with red crosses.

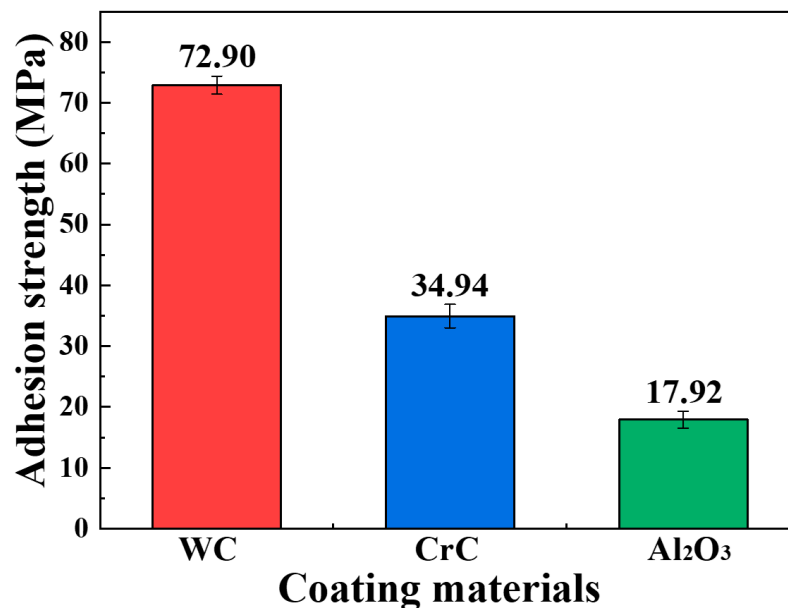
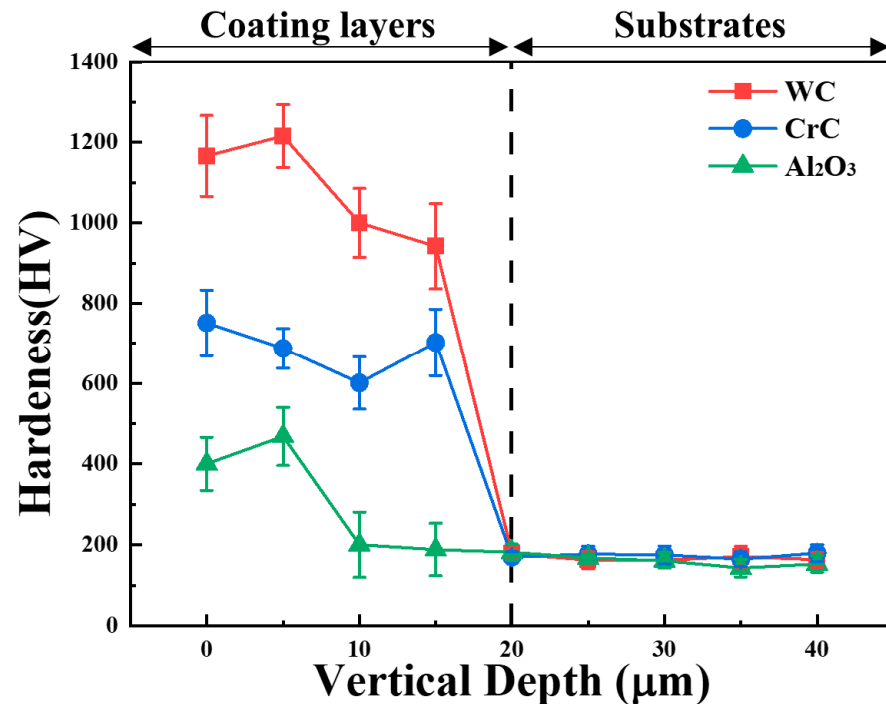


Figure 2. Adhesion strength between the substrate and the coating layers.



Figure 3 shows the micro Vickers hardness values measured in the depth direction from the surface of the coating materials to the substrate. The average hardness values of the WC, CrC, and Al<sub>2</sub>O<sub>3</sub> coating layers were 1081.39 HV, 685.40 HV, and 314.31 HV, respectively. The higher hardness value observed in the WC coating layer compared to other coating materials is attributed to the high-hardness tungsten carbide particles found in Figure 1. The Al<sub>2</sub>O<sub>3</sub> coating layer exhibits an average micro hardness of 194.26 HV in the buffer layer, similar to the substrate. All coating materials were uniformly applied to a certain extent, overcoming the characteristic uneven thickness of thermal spray coatings, and no deformation of the substrate due to exposure to high temperatures during the process was observed.



**Figure 3.** Vertical distribution of micro Vickers hardness from the top surface of the coating layers to the substrate.

Figure 4a,b show the wear rate based on the volume loss and the varying friction coefficient of the specimens after the wear test. The wear test results indicate that the wear rate follows the same order as the hardness results, with Al<sub>2</sub>O<sub>3</sub>, CrC, and WC showing decreasing wear rates, respectively. The variations in the friction coefficient graph are attributed to the increase (decrease) in the friction coefficient caused by wear debris (porosity). As shown in area fraction analysis, the WC material, which has fewer defects, exhibits uniform variations of microhardness. The average friction coefficients for the WC, CrC, and Al<sub>2</sub>O<sub>3</sub> materials are 0.548, 0.710, and 0.742, respectively. The friction coefficients of the WC and CrC materials stabilize over time, indicating increasing friction resistance [25]. The CrC and Al<sub>2</sub>O<sub>3</sub> materials exhibit rapid fluctuations compared to WC due to the significant number of cross-sectional defects, such as pores and cracks, which trap wear debris generated during the wear process and affect subsequent wear cycles. Notably, Al<sub>2</sub>O<sub>3</sub> shows a continuously increasing variation in the friction coefficient.

Figure 5a,b show the 3D and 2D profiles of the wear track width and depth. The depth and width of the tracks increase in the order of WC, CrC, and Al<sub>2</sub>O<sub>3</sub>, correlating with the increase in wear loss rate. The 3D profile results in Figure 5a reveal that the surface of the CrC and Al<sub>2</sub>O<sub>3</sub> coatings was indented by the counter material, whereas the WC surface appeared to be healed and convex. The analysis of the wear track depth (D) and width (W) using the 3D profiles in Figure 5b shows that the WC, CrC, and Al<sub>2</sub>O<sub>3</sub> materials have

wear track depths and widths of 7.7  $\mu\text{m}$  (D) and 626.6  $\mu\text{m}$  (W), 42.4  $\mu\text{m}$  (D) and 2120.6  $\mu\text{m}$  (W), and 24.0  $\mu\text{m}$  (D) and 1472.1  $\mu\text{m}$  (W), respectively. The WC material exhibits the least damaged wear track, suggesting it is the most effective coating layer for protecting the substrate.

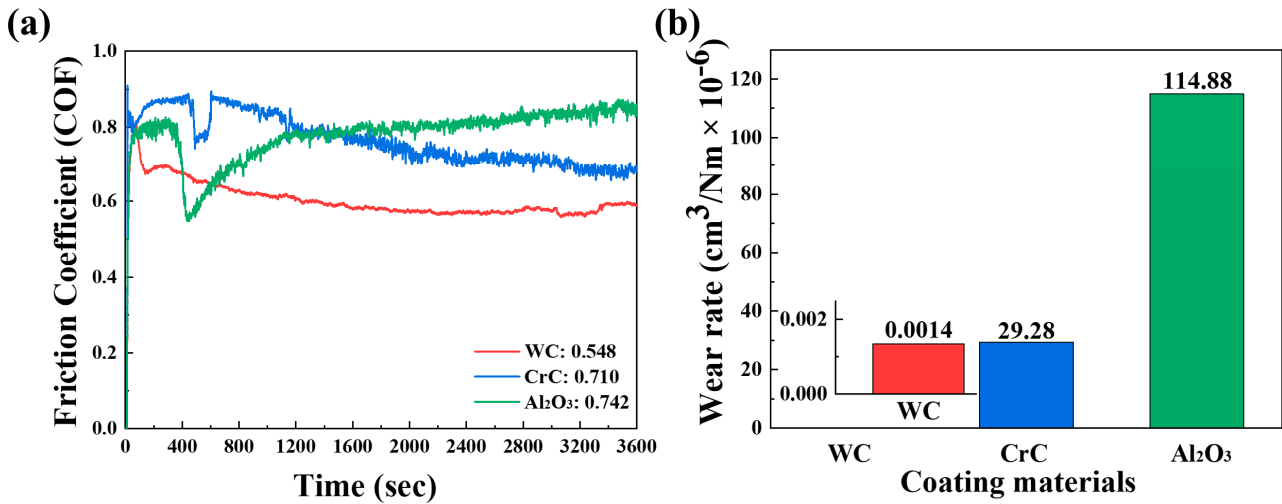


Figure 4. (a) Friction coefficient and (b) wear rate of the coating layers.

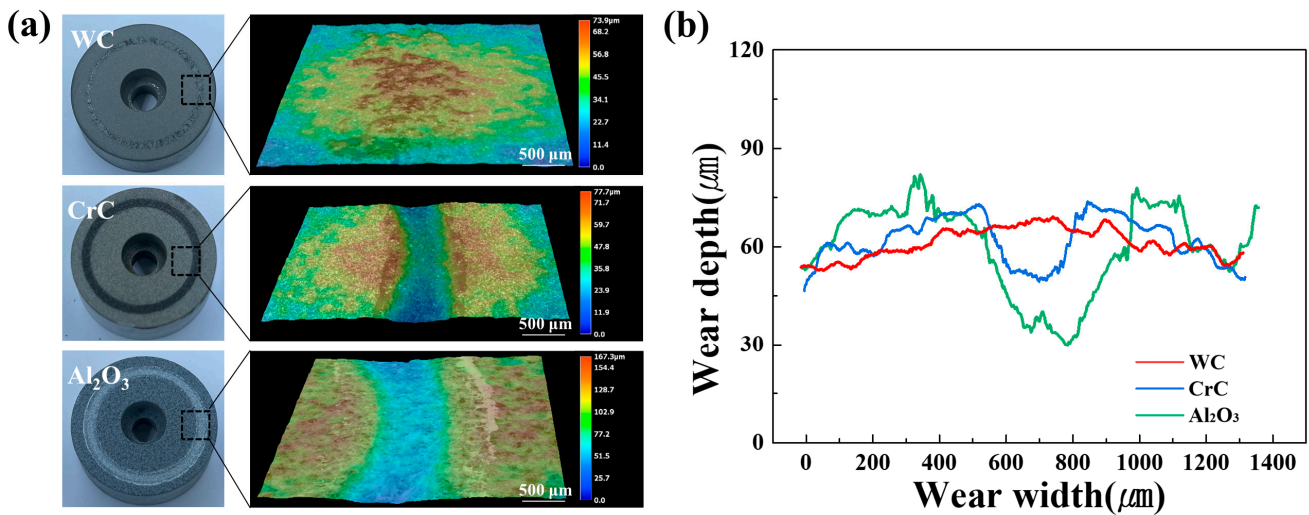
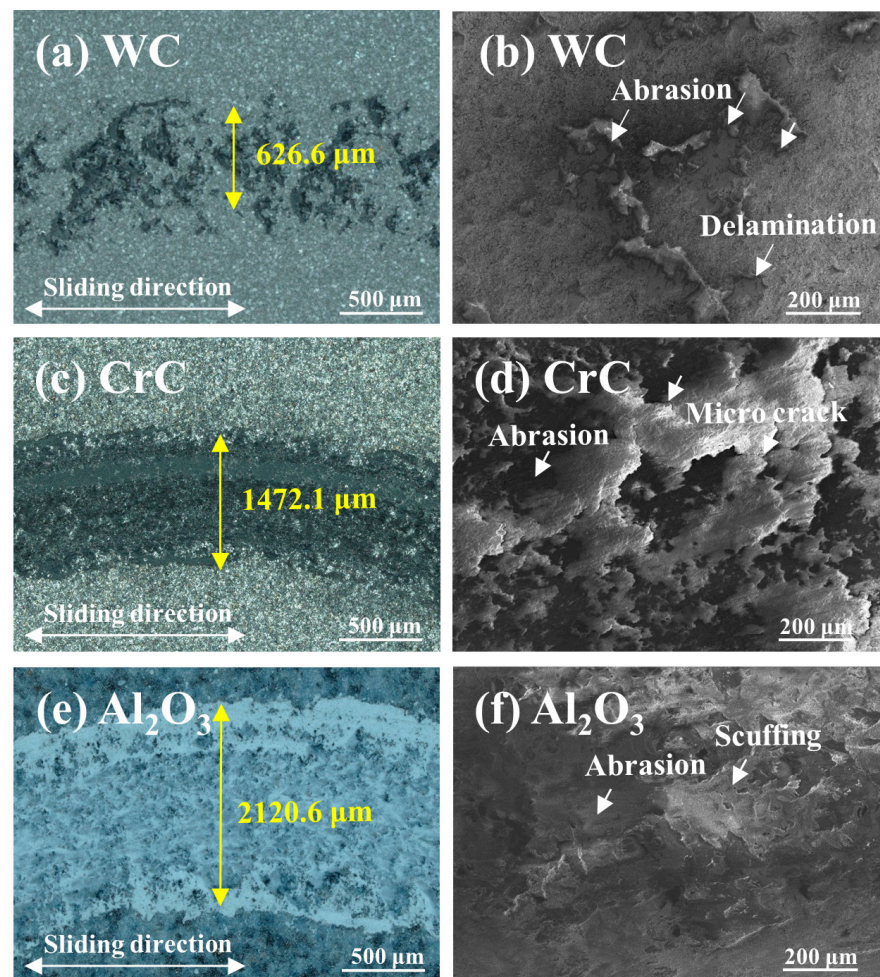
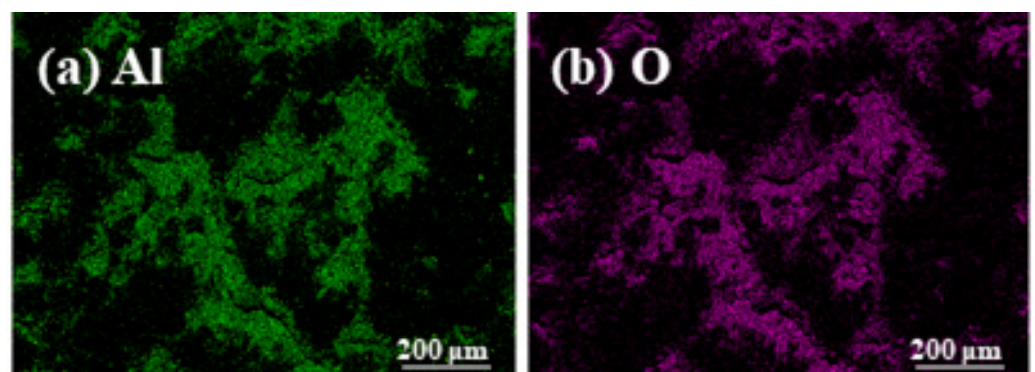


Figure 5. (a) 3D profiles and (b) 2D profiles showing the width and depth of the worn track on the coating layers.

To analyze the wear modes of each coating material, the microstructure of the wear track surface and EDS mapping results are shown in Figures 6 and 7. Figure 6a–c, respectively, display the wear track surfaces of WC, CrC, and Al<sub>2</sub>O<sub>3</sub>. The wear marks on the protruded WC coating layer indicate adhesive wear behavior, where material from the counter face adheres to the coating layer. The microstructure shows evidence of abrasion and microcracks, and the EDS results in Figure 7 depict the presence of (a) Al and (b) O, which are components of the counter material. Typical features of abrasive wear, such as abrasion and microcracks, were observed. In the CrC coating, abrasion, microcracks, and craters were also identified, while the Al<sub>2</sub>O<sub>3</sub> coating exhibited scuffing in the wear track due to the propagation of spalling pit cracks under repeated loading. The wear marks along the grooves in the Al<sub>2</sub>O<sub>3</sub> and CrC wear tracks suggest the occurrence of adhesive wear in the coating layers. It is evident that both abrasive and adhesive wear modes occurred in all coating layers.



**Figure 6.** Wear track microstructures of WC, CrC, and Al<sub>2</sub>O<sub>3</sub> analyzed by (a,c,e) 3D optical microscopy and (b,d,f) SEM.



**Figure 7.** The element mapping of (a) Al and (b) O detected in Figure 6b.

#### 4. Conclusions

This study confirms that the WC coating is the most effective option for enhancing the wear resistance of the A390 aluminum alloy, providing superior adhesion strength, hardness, and wear resistance. While the CrC and Al<sub>2</sub>O<sub>3</sub> coatings showed higher defect densities and wear rates, the WC coating's dense microstructure and minimal defects contribute significantly to its robust performance. Overall, the findings underscore the WC coating's potential to markedly enhance the durability of the A390 alloy in various industrial settings, offering a clear advantage over other tested coatings.



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