



# Friction and Wear Properties of CrAl-Based Coatings for Nuclear Fuel Cladding

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Friction and wear performance is one of the key mechanical properties of accident tolerant fuel cladding coatings. In this study, reciprocating sliding wear tests had performed on two types of CrAl and CrAlN coatings with two different Al content ratios and Zr-4. The coefficient of friction, wear depth, and abrasion loss were measured and compared. The results indicated that the CrAl-based coatings improve the wear behavior significantly and nitrogen has an obvious improvement on the wear resistance of the coating. The friction and wear performance was also studied in a water environment. The results show that the presence of water degrade the wear performance of Zr-4 and CrAl coatings but ameliorates the friction and wear performance of CrAlN coatings. The feasibility of depositing ATF coating on conventional Zr-4 substrates to mitigate the influence of grid-to-rod fretting was demonstrated.

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### Specialty section:

This article was submitted to  
Nuclear Energy,  
a section of the journal  
Frontiers in Energy Research

**Received:** 29 October 2020

**Accepted:** 18 January 2021

**Published:** 23 February 2021

### Citation:

Ma B, Luo B, Wang Z, Meng C and  
He X (2021) Friction and Wear  
Properties of CrAl-Based Coatings for  
Nuclear Fuel Cladding.  
Front. Energy Res. 9:622708.  
doi: 10.3389/fenrg.2021.622708

**Keywords:** accident-tolerant fuel, coating, chromium-aluminum coatings, friction and wear behavior, reciprocating wear

## INTRODUCTION

Zr alloys are widely used in the nuclear area because of its low neutron absorbing cross-section, good mechanical properties, and high corrosion resistance in reactor water (Choudhuri et al., 2012; Cekić et al., 2013; Kuprin et al., 2015). However, in the Fukushima nuclear accident, under the loss of coolant accident condition, zirconium reacts violently with water to produce a large amount of hydrogen, which causes the explosion and the radiation leak (Buessler et al., 2011; Blandford and Ahn, 2012; Buessler et al., 2012). These risks show that it is essential to improve the fuel safety and continually study the fuel degradation phenomena (Kurata et al., 2018). Hence, the researchers propose a new fuel system named accident-tolerant fuel (ATF). The ATF can be defined as a new system that increases the accident tolerance of loss of coolant accident condition over a long period of time while maintaining or improving fuel performance during normal operation condition (Zinkle et al., 2014; Kim et al., 2016; Terrani, 2018). One of the currently recognized method of the investigation of ATF is to deposit a coating on the Zr-4 cladding. A number of studies have exposed that ceramic coatings have excellent resistance to oxidation at high temperature, such as silicon carbide coating (Kato et al., 2014; Deck et al., 2015; Stone et al., 2015), TiAlCrN (Ma et al., 2019), and CrN (Meng et al., 2019), thus avoiding the oxidation of Zr-4 at high temperature. The other is to use the metallic coatings. For example, FeCrAl coatings, as Fe-based alloys have a high oxidation resistance (Zinkle et al., 2014; Pint et al., 2015; Massey et al., 2016). Cr coating is also used, because it has excellent corrosion resistance under severe condition on Zircoloy (Park et al., 2015; Brachet et al., 2016; Tang et al., 2017; He et al., 2019). Nowadays, CrAl-based coatings have attracted remarkable attention due to their excellent oxidation resistance. Zhong et al. (Zhong et al., 2018) performed high-temperature steam exposure to assess the oxidation behavior of CrAl coatings. It was demonstrated

that CrAl coatings with higher Al composition has lower weight gain in high temperature steam. Chen et al. (Chen et al., 2016) introduced TiN and ZrN insertion layer into CrAlN coating to study its mechanical and thermal properties. The results exhibit that CrAlN/TiN coating have a higher hardness than CrAlN/ZrN coating and TiN insertion layer have an improvement on the thermal stability of CrAlN coatings. Li et al. (Li et al., 2014) deposited CrAlN coatings with Zr alloying (Zr contents from 0 to 29.5 at.%) by d.c. reactive magnetron sputtering and investigated their mechanical properties, thermal stability, and oxidation resistance. They suggested that the coating hardness is improved by alloying of low contents of Zr. Zhang et al. (Zhang et al., 2019) used high current pulsed electron beam to deposit a CrAl coating on Al to study its microstructure and properties. They found that the microhardness and corrosion resistance were ameliorated because of the presence of Cr element.

Most published work mainly focused on the oxidation behavior and corrosion behavior of CrAl-based coatings nowadays. However, there are few studies on the friction and wear behavior of this coating. Furthermore, the influence of the water environment on the friction and wear behavior of this coating has not been investigated in details. In fact, in addition to oxidation performance, the mechanical properties of ATF also play an important role. In nuclear reactor components, such as fuel assemblies, flow induced vibrations can cause severe fretting wear (Rubiolo, 2006; Rubiolo and Young, 2009). Fretting is a low amplitude oscillatory motion between contacting bodies which leads to wear and fatigue damage (Neu, 2011). Many parameters such as amplitude, normal force, and coefficient of friction affect the type of wear and fatigue damage (Waterhouse, 1992). The grid-to-rod fretting (GTRF) is the most common cause of the fuel failure, and fuel rod fretting is a significant issue for designers of pressurized water reactor fuels (Edsinger et al., 2009). It is generally acknowledged that the turbulence from flow induced vibrations will cause the GTRF at the contact point between the grid and the fuel rod (Kim et al., 2016). According to the past research, the mechanism of GTRF degradation depends extremely on the grid-to-rod gap as a function of burnup, in particular under condition of high flow rates which is the vibration caused by the excessive flow (Kim and Lee, 2003; Kovács et al., 2009, Kovács et al., 2013; Jiang et al., 2016). Though the reliability of nuclear fuels can be increased by the new design features to strengthen the wear resistance of spacer grids with the same Zr alloy, the irradiation causes the fuel rod to elongate in the axial direction, and the thermal creep caused by the pressure difference between the inside and outside of the cladding results in radial contraction. These two factors lead to a gradual increasement of the grid-to-rod gap. It is excessively difficult to avoid this phenomenon under condition of operating temperature and high neutron irradiation. As a result, the wear behavior becomes the important parameters of the ATF and the tribological properties of new ATF material is not clearly understood. Therefore, it is worthy to study on the friction and wear behavior of new ATF materials.

In this work, CrAl-based coatings were deposited on Zr-4 by multi-arc ion plating. The GTRF situation is characterized by the

**TABLE 1** | Concentration of Cr, Al, and N in the CrAl-based coatings and mechanical properties for ATF candidates and Zr-4.

Material	Target Cr:Al ratio (at. %)	Element proportions (at.%)			Hardness [HV]
		Cr	Al	N	
Zr-4	—	—	—	—	234
CrAl-1	30:70	32.1	67.9	—	380
CrAl-2	50:50	45.8	54.2	—	390
CrAlN-1	30:70	17.4	36.7	46.0	1,020
CrAlN-2	50:50	35.1	25.6	39.4	1,080

friction and wear of the coating with a zirconia ball under different conditions, i.e., unlubricated condition and water lubrication condition. The results show that CrAlN coatings' friction and wear performance is more predominant than that of CrAl coatings, and the CrAlN coating with lower Al composition have the best friction and wear performance.

## EXPERIMENT PROCEDURE

In this study, four ATF candidates (two CrAl coatings and two CrAlN coatings with different Al content the detail is exhibited in **Table 1**) and a conventional Zr-4 alloy with a length of 25 mm, a width of 15 mm and a thickness of 2 mm were prepared and tested. The coatings are deposited on the Zr-4 alloy substrate by multi-arc ion plating. The deposition time of CrAl coatings is 2 h and 6 h, and for CrAlN coatings, the deposition time is 2 h. The hardness of Zr-4 and four CrAl-based coatings were measured by Vickers hardness tester and the results are shown in **Table 1**. The friction and wear tests were in progress in a reciprocating sliding tester at room temperature (27°C) in unlubricated (air) condition and in water lubrication. For the water lubrication condition, the specimens were placed in a tank filled with deionized water. The effects of water fluidity, temperature, and hydrochemistry on the results were not taken into account. The schematic diagram of test machine is shown in **Figure 1**. Each specimen is subjected to a friction test with a zirconia ball. The reciprocating stroke is fixed at 10 mm, the reciprocating frequency is 13 Hz, and the friction time is 3 min and 10 min. Since the friction and wear properties of coatings cannot be clearly compared with either too large or too small loads, the contact normal forces of 1 and 5 N are applied under the experiment condition. During the experiment, the coefficient of friction (COF) was continuously monitored in order to observe the relationship between the change of COF curve and the wear amount of each sample. After the wear test, the worn surface of each sample is observed by the surface profilometer to measure the wear depth and the amount of friction wear. The amount of wear volume was detected using a non-contact three-dimensional laser interferometer.

To characterize the coatings, X-ray diffraction (XRD) was conducted to determine the phase composition of samples. The energy-dispersive X-ray spectroscopy (EDS) profiles were performed to study the distribution of Cr, Al, and N elements. The scanning electron microscopy (SEM) was applied to assess

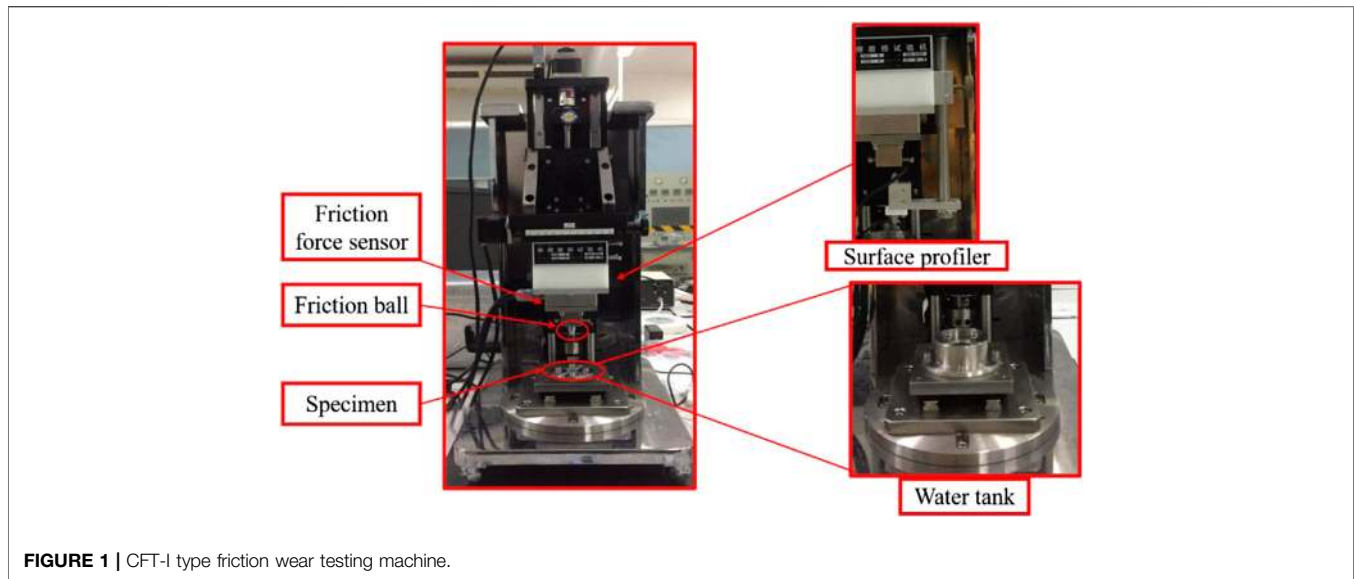


FIGURE 1 | CFT-I type friction wear testing machine.

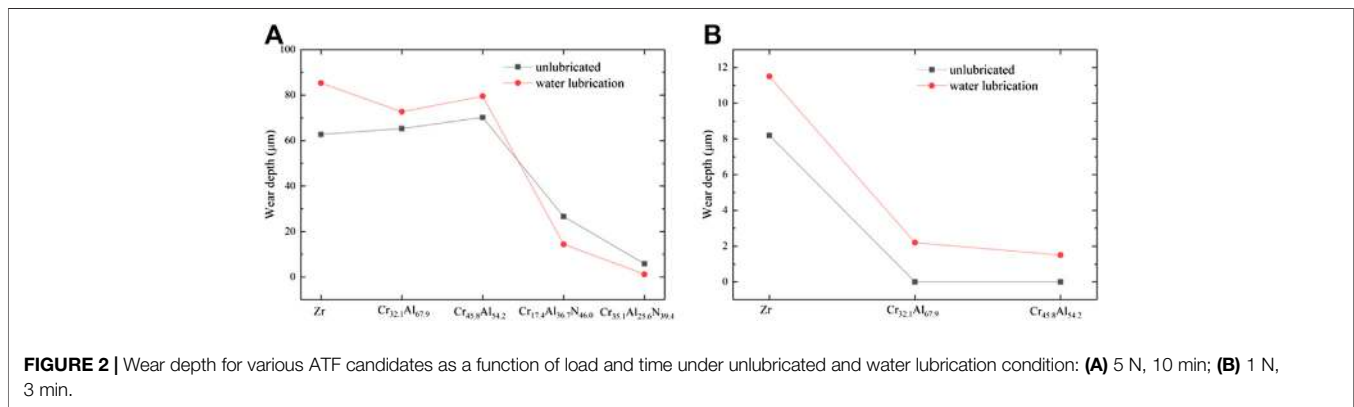


FIGURE 2 | Wear depth for various ATF candidates as a function of load and time under unlubricated and water lubrication condition: (A) 5 N, 10 min; (B) 1 N, 3 min.

the coatings thickness and adhesion with substrate. Meanwhile, nanoindentation test was performed to assess the mechanical properties of coatings.

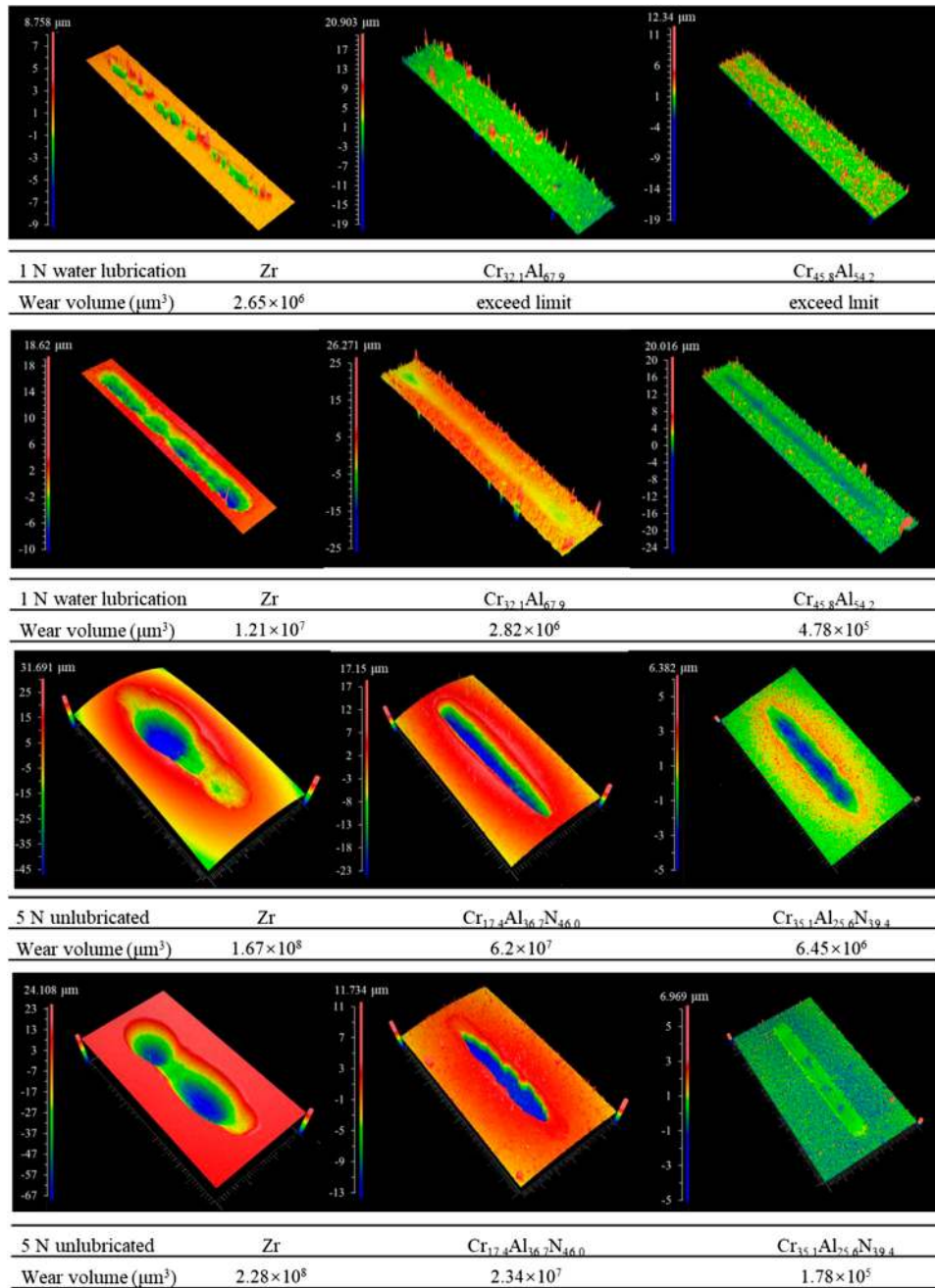
## RESULTS AND DISCUSSION

### Wear Behavior

After the reciprocating sliding friction test, the wear depth was obtained from the bottom sample. Figure 2 depicts the measurements of the maximum wear depth for various ATF coating under different loads in unlubricated and water lubrication condition, the experiment time is 10 and 3 min. The difference in wear depth of different samples is considered as the difference in the friction and wear performance of the samples. It can be found that the wear depth under different loads is quite different, which indicates that different loads have different effects on the damage of the coating. Under a load of 5 N, it will be found that Zr, Cr<sub>32.1</sub>Al<sub>67.9</sub>, and Cr<sub>45.8</sub>Al<sub>54.2</sub> have similar wear depths. Under a load of 1 N, the wear depth of the CrAl coating will be smaller than that of Zr.

This shows that when the load is small, the CrAl coating has a protective effect on the Zr substrate. However, the coating was destroyed by relatively large load, which will lead to the disappearance of the protective effect of the CrAl coating on the Zr substrate. Furthermore, it can be found that under the load of 5 N, the CrAlN coatings have a more significant protective effect on the Zr substrate compared to the CrAl coatings. At the same time, the wear performance of the Cr<sub>35.1</sub>Al<sub>25.6</sub>N<sub>39.4</sub> coating is better than that of the Cr<sub>17.4</sub>Al<sub>36.7</sub>N<sub>46.0</sub> coating due to the smaller wear depth.

The presence of water also has influence on wear performance, in addition, the impact on CrAl coatings and CrAlN coatings is distinct. For Zr, Cr<sub>32.1</sub>Al<sub>67.9</sub>, and Cr<sub>45.8</sub>Al<sub>54.2</sub> coatings, under 1 and 5 N loads, the wear depths under water lubrication are more significant than that under unlubricated condition. For Cr<sub>17.4</sub>Al<sub>36.7</sub>N<sub>46.0</sub> and Cr<sub>35.1</sub>Al<sub>25.6</sub>N<sub>39.4</sub>, the wear depths under water lubrication are less than that under unlubricated condition. It is considered that the wear behavior of CrAl coatings and CrAlN coatings in water environment is different (He et al., 2020). For the CrAl coatings and Zr-4, the wear behavior is abrasive wear, during this process, there will be



**FIGURE 3** | Results of noncontact three-dimensional optical profilometry under different conditions.

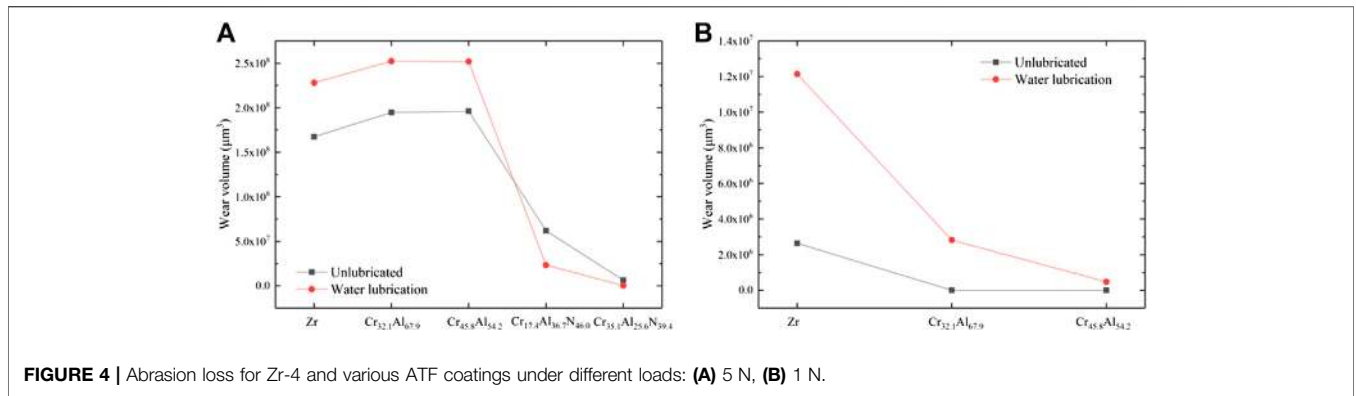
debris attached to the scratched surface, which lead to the increasement of wear depth. For the CrAlN coatings, the wear behavior is adhesive wear, the lubrication during friction reduce the degree of wear.

In order to assess the lubricant effect to the wear of various coatings more accurately, for the CrAl coatings, a 3-minute friction and wear under water lubrication and unlubricated condition was carried out under a load of 1 N. Zr-4 was also tested as a control group under the same condition. For the

CrAlN coatings, a 10-minute friction and wear test under water lubrication and unlubricated condition was conducted, identically, Zr-4 had the experiment under the same condition. After the experiment, the scratches were measured using noncontact three-dimensional optical profilometry.

According to **Figure 3**, the wear shapes of various ATF candidate coatings and Zr-4 are substantially similar under the presence or absence of lubrication condition. The wear shape of the Zr-4 and the CrAl coatings is largely different from the wear





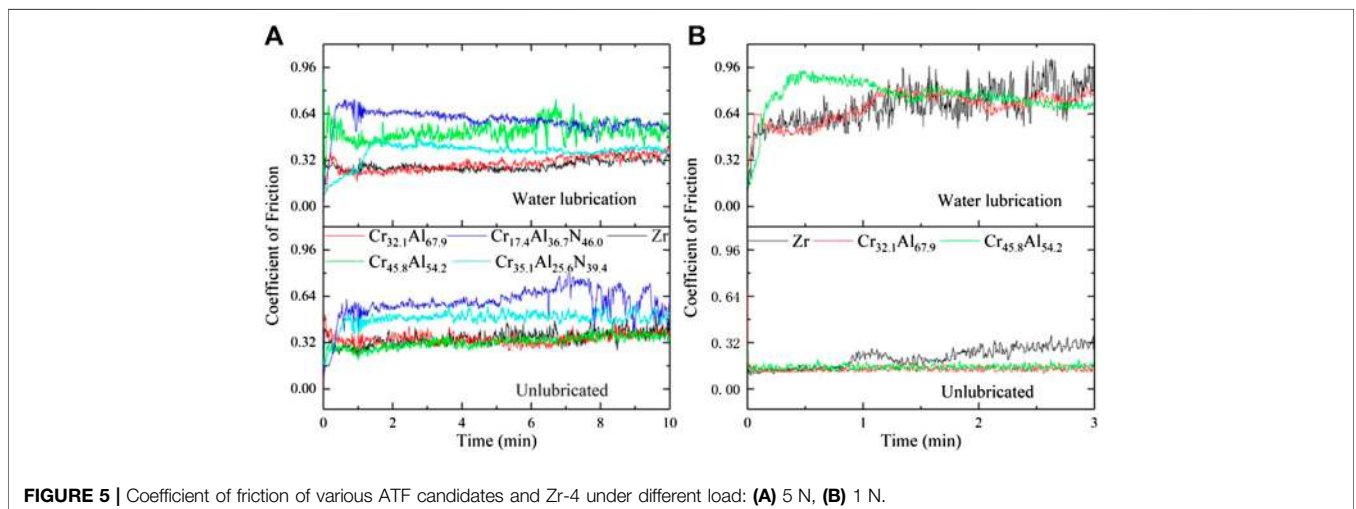
shape of the CrAlN coatings, which can be rationalized by the degree of damage of the coatings. As can be seen from the results of the noncontact three-dimensional optical profilometry, the surface of the coating is relatively rough. The shallower the scratch, the rougher the surface is observed. For the  $Cr_{32.1}Al_{67.9}$  and  $Cr_{45.8}Al_{54.2}$  coatings under unlubricated condition, the wear depth is too small, which has exceeded the lower limit of the noncontact three-dimensional optical profilometry measurement. Compared with unlubricated condition, the uncoated Zr-4,  $Cr_{32.1}Al_{67.9}$ , and  $Cr_{45.8}Al_{54.2}$  coatings wear more under water lubrication condition. Meanwhile the wear depth of CrAlN coatings under water lubrication condition are less than that of unlubricated condition, which is consistent with the results obtained by the scratch measurement instrument. Under the same condition, it can be realized that the Zr-4 with coatings have less wear debris than uncoated Zr-4. Thus, the CrAl-based coatings has a protective effect on the substrate and enhances the wear resistance.

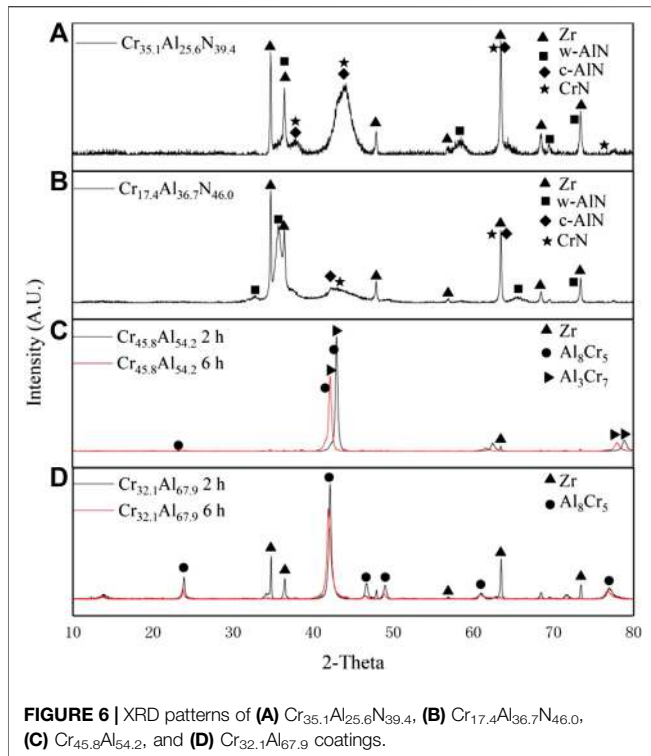
As can be seen from the results of the noncontact three-dimensional optical profilometry, the surface of the coating is relatively rough. When the wear depth is smaller, the surface roughness observed in the noncontact three-dimensional optical

profilometry is more obvious, because the sample surface has an unevenness of about  $0.5\ \mu m$ , which is not negligible in the noncontact three-dimensional optical profilometry result when the wear depth is small. Whereas the wear depth under 5 N is deep, and the unevenness of the surface is negligible, thus this difference is produced in the picture results of the noncontact three-dimensional optical profilometry.

The abrasion loss of Zr-4 and two CrAl coatings increased under the condition of water lubrication. According to the results of the noncontact three-dimensional optical profilometry, water lubrication has influence on the scratch shape, hence, the main reason for this result was the change of wear depth. This is consistent with previous results. **Figure 4** reveals the wear volume of each ATF coatings after experiment with different conditions. The abrasion loss of CrAlN coating has little change, among which the abrasion loss of  $Cr_{35.1}Al_{25.6}N_{39.4}$  has almost no change, which indicates that it has good wear resistance under both conditions.

Under a load of 1 N, the abrasion loss of the CrAl coating is much less than that of Zr under unlubricated and water lubrication conditions, and shows a good protective effect on the Zr substrate, which is consistent with the previous results.





## Coefficient of Friction

Figure 5 shows the friction coefficient for different ATF candidates and Zr-4 under water lubrication and unlubricated condition with different loads. The friction coefficient curve showed fluctuations under water lubrication and unlubricated conditions, which lasted until the end of tests. It can be rationalized that under unlubricated condition, since there are worn debris detached from the contact surface and remains on the surface, the worn surface becomes rougher, and then small worn debris may gather in an appropriate position on the irregular worn surface, which may form a worn debris layer.

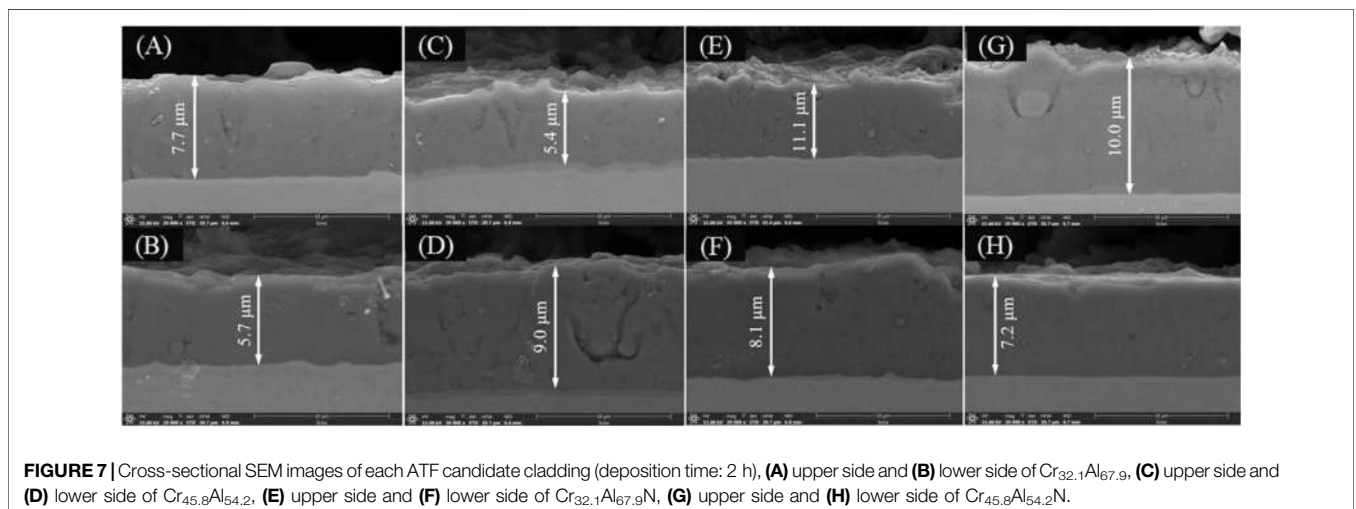
Therefore, the irregularity and fluctuation of the friction coefficient curve are closely related to the fracture of the worn debris layer and the interaction between the worn debris and the contact surface. It is possible that the frictional debris layer is difficult to be formed under the condition of a small load due to water lubrication. As a result, the fluctuation of the friction coefficient curve is caused by the damage of a surface coating and the periodic accumulation and damage of debris layer on the coating surface.

Under a load of 1 N without water lubrication, the friction coefficient of  $\text{Cr}_{32.1}\text{Al}_{67.9}$  and  $\text{Cr}_{45.8}\text{Al}_{54.2}$  is always around 0.15, indicating that the surface of the CrAl-coating was not severely damaged during the experiment, which is consistent with the previous results of wear depth and wear volume as shown in Figures 2, 3, respectively. Under water lubrication, the friction coefficient is quite different from that under the condition without water lubrication. It can be observed that when other conditions are the same, the coefficient of friction under water lubrication is more significant than that in air condition, which may be related to the viscosity of water. The fluctuation of the friction coefficient curve also reflects a certain change for the coating surface.

Under a load of 5 N, it is noticed that the friction coefficient curves of CrAl coating and CrAlN coating are distinct. This is due to the fact that the CrAl coating is a metallic coating and the CrAlN coating is a ceramic coating and their surface properties are different. Meanwhile, the coefficient of friction does not increase significantly under water conditions compared to 1 N with the same condition.

## X-Ray Diffraction Analysis

XRD patterns of  $\text{Cr}_{32.1}\text{Al}_{67.9}$  and  $\text{Cr}_{45.8}\text{Al}_{54.2}$  coatings with different deposition time and  $\text{Cr}_{17.4}\text{Al}_{36.7}\text{N}_{46.0}$ ,  $\text{Cr}_{35.1}\text{Al}_{25.6}\text{N}_{39.4}$  coatings with 2 h deposition are presented in Figure 6. According to Figures 6A,B, the c-AlN, w-AlN and CrN pattern can be detected in the results of  $\text{Cr}_{35.1}\text{Al}_{25.6}\text{N}_{39.4}$  and  $\text{Cr}_{17.4}\text{Al}_{36.7}\text{N}_{46.0}$  coating. The  $\text{Cr}_{35.1}\text{Al}_{25.6}\text{N}_{39.4}$  and  $\text{Cr}_{17.4}\text{Al}_{36.7}\text{N}_{46.0}$  coating have the same composition, nevertheless, their peak distributions are



**TABLE 2** | Thickness of various ATF coatings with 2 h deposition.

Materials	Thickness of the upper side ( $\mu\text{m}$ )	Thickness of the lower side ( $\mu\text{m}$ )	Average thickness of two sides ( $\mu\text{m}$ )
$\text{Cr}_{32.1}\text{Al}_{67.9}$	7.7	5.7	6.7
$\text{Cr}_{45.8}\text{Al}_{54.2}$	5.4	9.0	7.2
$\text{Cr}_{17.4}\text{Al}_{36.7}\text{N}_{46.0}$	11.1	8.1	9.6
$\text{Cr}_{35.1}\text{Al}_{25.6}\text{N}_{39.4}$	10.0	7.2	8.6

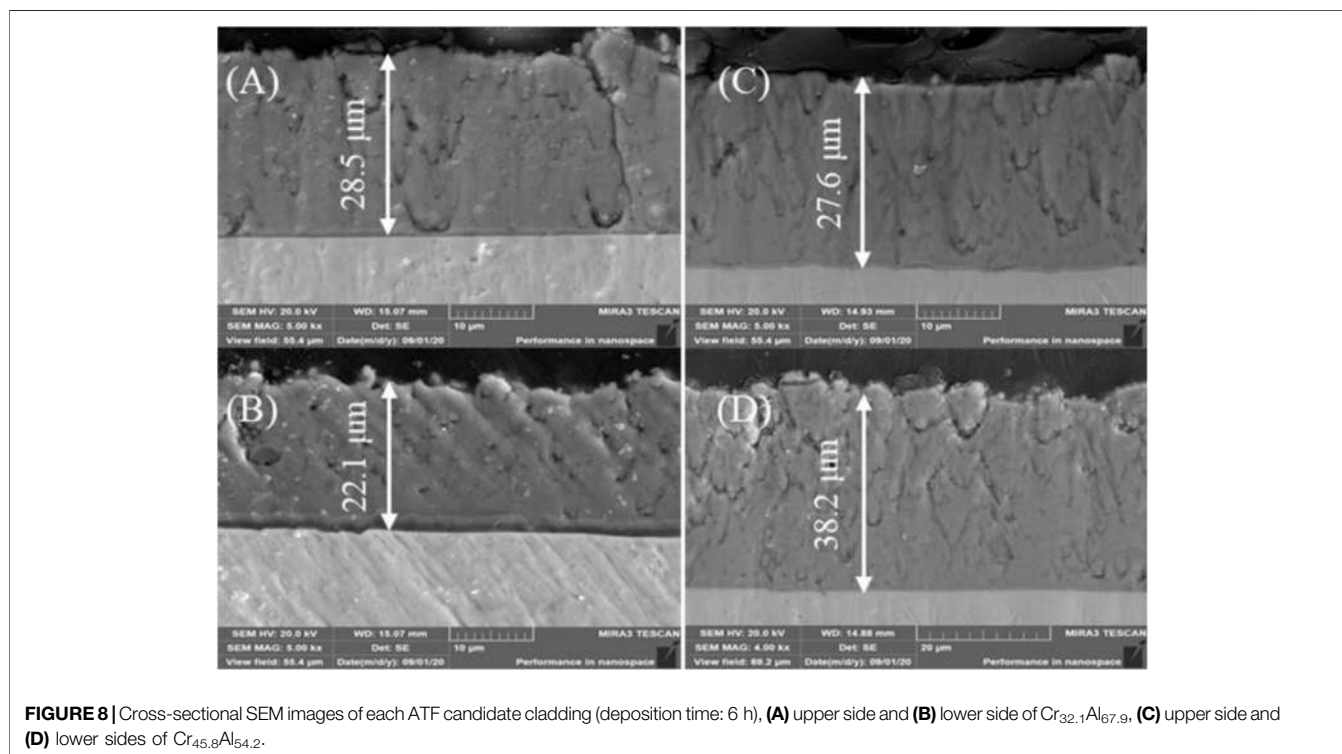
quite different. Strong  $\text{Al}_8\text{Cr}_5$  peaks can be observed on both  $\text{Cr}_{32.1}\text{Al}_{67.9}$  and  $\text{Cr}_{45.8}\text{Al}_{54.2}$  coatings, meanwhile  $\text{Al}_3\text{Cr}_7$  peak is only observed on  $\text{Cr}_{45.8}\text{Al}_{54.2}$  coating. Compared with the results of  $\text{Cr}_{32.1}\text{Al}_{67.9}$  and  $\text{Cr}_{45.8}\text{Al}_{54.2}$  with 2 h deposition, the XRD results indicate that CrAl coatings with 6 h deposition have the same composition, however, the coatings with 6 h deposition have weaker peak of Zr. The XRD results above indicate that Cr, Al and N elements are well involved in the crystal structure in the coatings, and the crystal structure will change with the change of element type and content. In addition, with the extension of deposition time, the diffraction signal of CrAl coating also increases, which is the result of the increase of coating thickness.

## Scanning Electron Microscopy and Energy-Dispersive X-Ray Spectroscopy Analyses

Figure 7 shows the results of SEM for four ATF candidate coatings. The measured thickness of each ATF candidate

coating is shown in the Table 2 according to the results in Figure 7. The thickness of each coating does not exceed  $10\ \mu\text{m}$ , the hardness of the CrAl coating is larger than that of Zr-4, meanwhile, the wear depth of CrAl coatings with 2 h deposition time is pretty close to that of Zr-4, which is due to the coating is relatively thin. When the coating is destroyed, its protective effect on the substrate disappears. The thickness of  $\text{Cr}_{32.1}\text{Al}_{67.9}$  and  $\text{Cr}_{45.8}\text{Al}_{54.2}$  coating with 2 h deposition time are thinner than that of  $\text{Cr}_{17.4}\text{Al}_{36.7}\text{N}_{46.0}$   $\text{Cr}_{35.1}\text{Al}_{25.6}\text{N}_{39.4}$ . It can be rationalized by the material compatibility of different coatings, which means the presence of nitrogen accelerates the deposition rate of particles, resulting in a better adhesion for CrAlN with Zr-4. Figure 8 exhibits the SEM result of CrAl coatings with 6 h deposition. Table 3 reveals the thickness of CrAl coatings with 6 h deposition time. As the deposition time increases, the coating remains highly dense, the thickness of the coating increases, and the wear depth does not exceed the thickness of the coating. This also explains the protective effect of the CrAl coating on the Zr substrate, that is, it can significantly enhance the wear resistance before the coating is severely damaged.

The distribution of various elements on the coating with 2 h deposition is shown in the EDS results in Figure 9. The coating and the Zr-4 substrate have a distinct boundary. The substrate does not contain Cr, Al, and N elements, and the coating does not contain Zr elements, which means that the coating does not affect the properties of the Zr substrate, this also proves that it is feasible to prevent GTRF by physical vapor deposition. When compare the result of Figures 9A–D, it can be seen that the content of Cr and Al in the coating is significantly less in the CrAlN coatings. This is consistent with the conclusion that the presence of



**FIGURE 8** | Cross-sectional SEM images of each ATF candidate cladding (deposition time: 6 h), (A) upper side and (B) lower side of  $\text{Cr}_{32.1}\text{Al}_{67.9}$ , (C) upper side and (D) lower sides of  $\text{Cr}_{45.8}\text{Al}_{54.2}$ .

**TABLE 3** | Thickness of various ATF coatings with 6 h deposition.

Materials	Thickness of the upper side (μm)	Thickness of the lower side (μm)	Average thickness of two sides (μm)
Cr <sub>32.1</sub> Al <sub>67.9</sub>	28.5	22.1	25.3
Cr <sub>45.8</sub> Al <sub>54.2</sub>	27.6	38.2	32.9

nitrogen changes the properties of the coating. **Figure 10** reveals the distribution of elements on CrAl coatings with 6 h deposition. The distribution of coating elements did not change, indicating that the coating structure would not be affected by the deposition time.

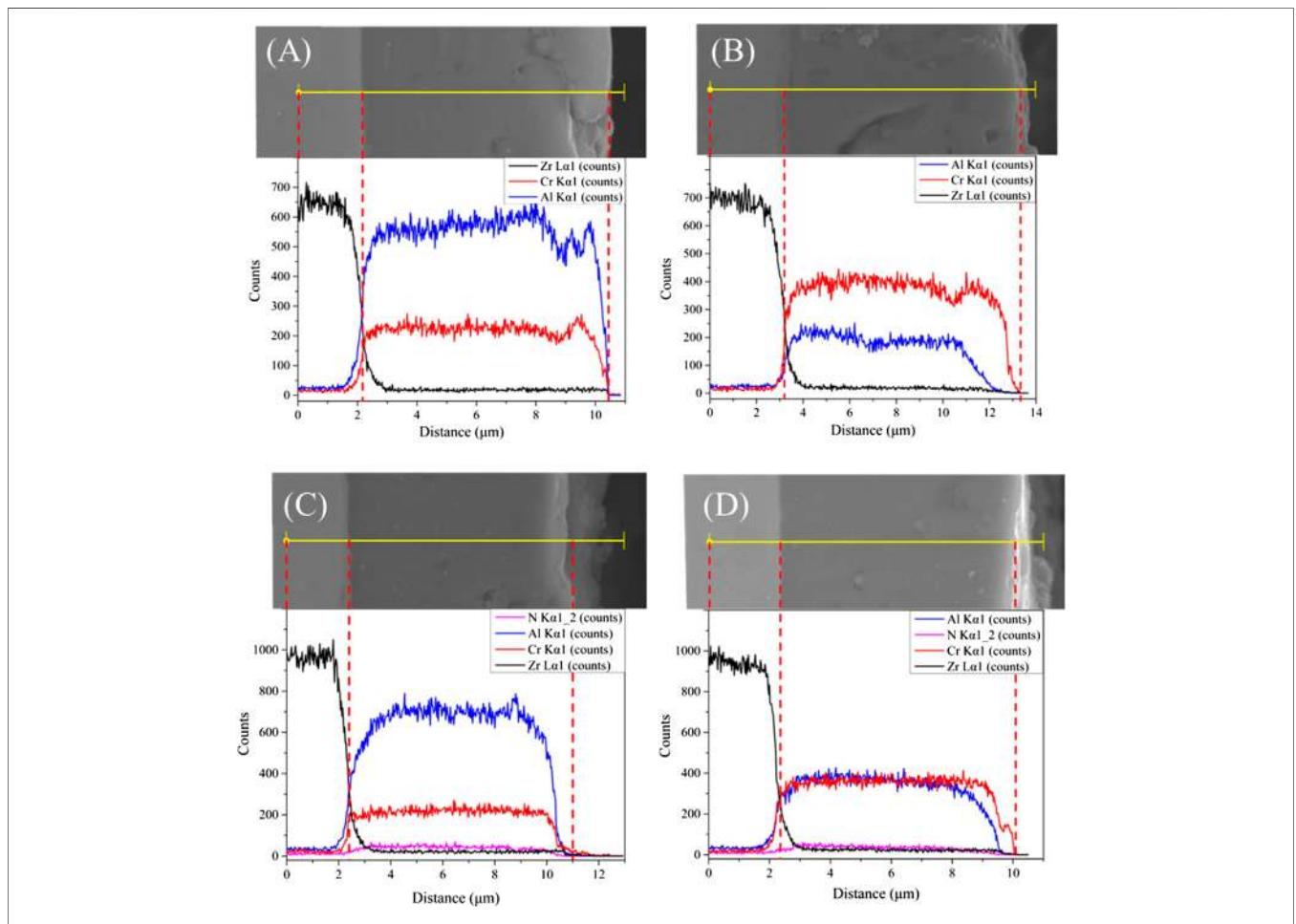
### Young's Modulus

The results of nanoindentation test are presented in **Figure 11**. Young's modulus of CrAl coatings are close to that of Zr-4, while the Young's modulus for CrAlN coatings are much higher than that of Zr-4. Considering the Young's modulus and hardness of the samples, the indentation depth of 3,000 nm and 2000 nm for CrAl and CrAlN coatings was chosen, respectively. The Young's

modulus of Cr<sub>35.1</sub>Al<sub>25.6</sub>N<sub>39.4</sub> coating is the highest, followed by Cr<sub>17.4</sub>Al<sub>36.7</sub>N<sub>46.0</sub> coating. The larger Young's modulus means the smaller elastic deformation under the same stress, which is also consistent with the results of previous friction and wear tests. The larger the Young's modulus is, the substrate will have more efficient protection. However, if the difference of Young's modulus between coating and substrate is huge (e.g., Cr<sub>35.1</sub>Al<sub>25.6</sub>N<sub>39.4</sub> and Cr<sub>17.4</sub>Al<sub>36.7</sub>N<sub>46.0</sub> coatings), a superior adhesion between coating and Zr-4 is required. The results of SEM and friction experiment show that the CrAlN coatings have an excellent adhesion, hence, they have a significant protective effect. Meanwhile, for CrAl and CrAlN coatings, the Young's modulus of the coating with high content is larger. This is consistent with the results of hardness and wear behavior of coatings.

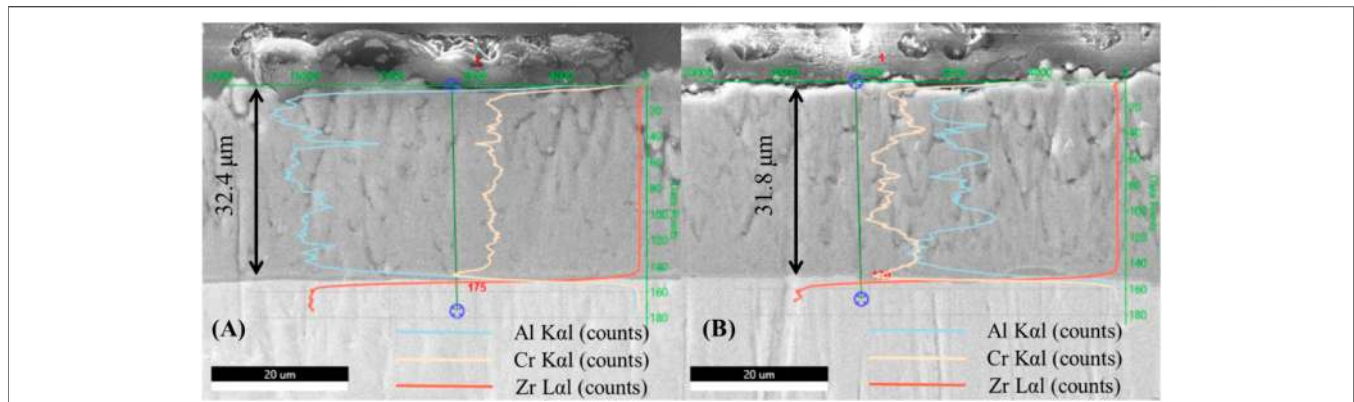
### SUMMARY AND CONCLUSION

Various types of accident-tolerant fuel coatings are examined to improve the resistance to grid-to-rod

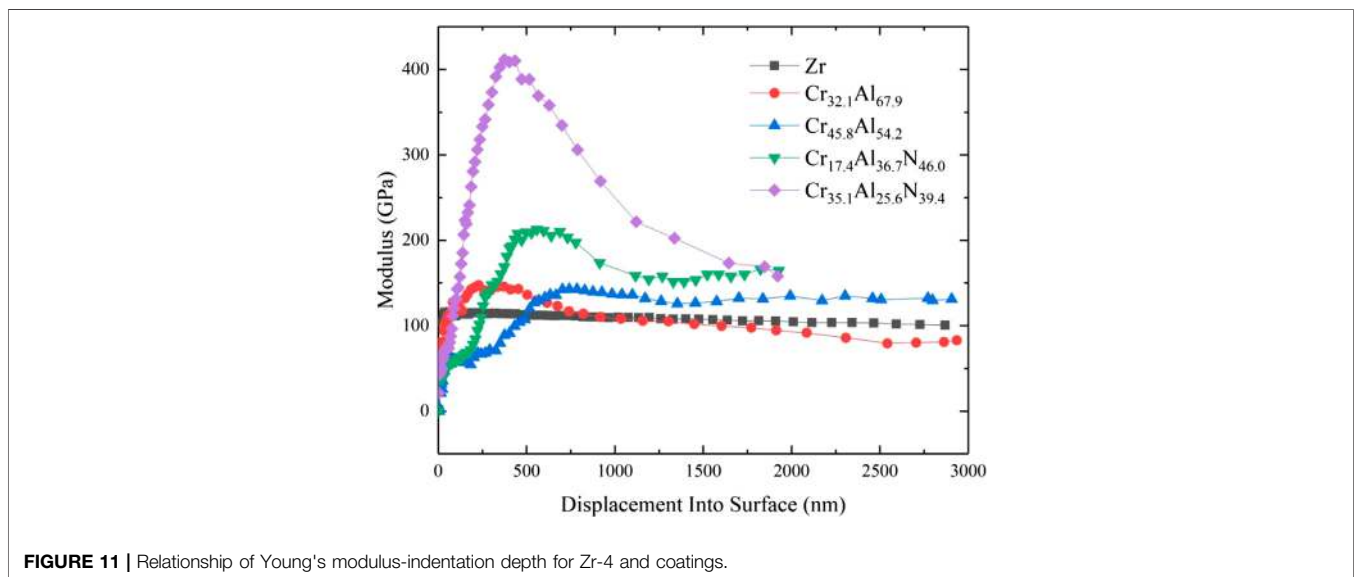


**FIGURE 9** | EDS results of cross-section of each ATF candidate cladding with 2 h deposition: **(A)** Cr<sub>32.1</sub>Al<sub>67.9</sub>, **(B)** Cr<sub>45.8</sub>Al<sub>54.2</sub>, **(C)** Cr<sub>32.1</sub>Al<sub>67.9</sub>N, and **(D)** Cr<sub>45.8</sub>Al<sub>54.2</sub>N.





**FIGURE 10** | EDS results of cross-section of each ATF candidate cladding with 6 h deposition: **(A)**  $\text{Cr}_{32.1}\text{Al}_{67.9}$ , **(B)**  $\text{Cr}_{45.8}\text{Al}_{54.2}$ .



**FIGURE 11** | Relationship of Young's modulus-indentation depth for Zr-4 and coatings.

fretting degradation. It was found that CrAl-based coatings have a pretty protect effect to Zr-4 substrate, which improve the resistance to GTRF degradation. Furthermore, CrAlN coatings have a better protective effect than CrAl coatings. When the coating is severely damaged, this protective effect of CrAl coatings will disappear. Under water lubrication, the CrAl coatings' resistance decrease, whereas the CrAlN coatings' resistance improve. The presence of nitrogen significantly affects coating properties, including coating hardness, abrasion resistance, and coating deposition rate.

Therefore, in order to improve the resistance to grid-to-rod fretting degradation, it is suggested to use coatings with higher hardness (e.g., CrAlN coatings) to deposit on current Zr-4, and the coating should be deposited thicker. Deposit coatings on Zr-4 is a feasible method to improve the resistance to GTRF degradation.

## DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

## AUTHOR CONTRIBUTIONS

BM: Original draft preparation BL: Friction and wear experiment and characterization ZW: Article polishing CM: Coating preparation XH: Guide the whole research idea.

## FUNDING

This work was supported by the Fundamental Research Funds for the Central Universities (19lgpy299).

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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