Research Article

Investigation on Microblasting Applied to CrN Coatings

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A microblasting treatment carried out on CrN coated samples was studied to investigate the induced effect on corrosion and wear resistance. CrN coating was deposited through Cathodic Arc Evaporation technique on quenched and tempered steel. The properties of the coating were studied by hardness measurements, scratch, potentiodynamic, and pin-on-disk tests. The results show that microblasting reduces the corrosion resistance while improving the wear behavior.

1. Introduction

Physical Vapor Deposition (PVD) coatings are widely used to increase the wear and corrosion resistance of many substrates [1, 2] and for different applications [3, 4]. As known, in fact, the presence of a ceramic layer increases the hardness of the surface and reduces the friction coefficient [2]. Additionally, due to the chemical stability of the deposited material, the corrosion of the substrate is strongly reduced [1, 2].

Different authors have already demonstrated that the corrosion resistance of a coated part is influenced by the presence of defects formed during the deposition process, such as pores or droplets [5–7]. It follows that the removal of coating defects, without affecting other coating properties, can be an interesting method to enhance corrosion resistance [1].

The microblasting of the film surface has been proven to remove droplets from Cathodic Arc Evaporation (CAE) coating surface, without reducing the adhesion or the mechanical properties of the coating itself [8]. This technique is commercially used to clean the surface of each layer during multilayer depositions, by removing some defects that have a detrimental effect on layers adhesion.

Authors also stated that microblasting, performed with the correct parameters, extends the life of cutting tools in the case of TiAlN coated cemented carbide inserts [9–12]. Additionally, a study on CrN deposited on steel demonstrates that the fatigue resistance is enhanced after the mechanical surface treatment, due to the increase in the residual compressive stress of the substrate skin [13]. As known, in fact, the fatigue cracks in PVD coated parts nucleate at the interface between coating and substrate [14, 15].

To the authors’ knowledge, no research works correlating microblasting with corrosion and wear resistance of CAE CrN coatings have been published. Therefore, the aim of this paper is to investigate these properties on a 4 μm CrN coating, deposited on quenched and tempered steel and subjected to microblasting. A first set of tests was performed to identify the correct microblasting parameters. The properties of the coating after microblasting were then assessed by means of hardness measurements, scratch, potentiodynamic, and pin-on-disk tests. The results were also compared to those obtained for the as-deposited state.

2. Materials and Methods

The specimens were machined from quenched and tempered steel, whose hardness is 400 HB. In Table 1 the chemical composition of the steel is reported.

Wear and corrosion test samples were cut as disks of 40 mm and 18 mm in diameter, respectively. In both cases, the flat surfaces were polished up to mirror finishing.

CrN deposition was carried out with a Metaplast system, working in a nitrogen atmosphere with a pressure of 1.4 × 10⁻² bar. Cathode power and bias were set at 60 Ah...
and 80 V, respectively. Before the deposition, argon etching was performed with a bias of 200 V. The obtained coating thickness was 4 µm, as resulting by Calotest measurements.

After the coating deposition, dry microblasting (MB) was carried out with silica microspheres, having a diameter ranging between 70 and 100 µm and a mean hardness of 47 HRC. The nozzle angle was set at 90° with respect to the coating surface. Three blasting pressures were used (2, 3, and 6 bar) with the aim of identifying the maximum pressure able to modify the surface without causing damage of the coating.

In order to understand the effect of the microblasting on tribological and corrosion resistance of the investigated coating, three groups of samples were considered:

(i) bare steel (named BS);
(ii) coated samples without microblasting treatment (named CrN);
(iii) coated samples with microblasting treatment (named CrN + MB).

A mechanical characterization of the coating was performed on the samples with and without microblasting. Nanoindentation and elastic modulus were obtained through a Table Top Nanoindentation Tester (TTX-NHT) of CSM Instruments. The maximum load (50 mN) was chosen in order to maintain the penetration depth lower than 10% of the coating thickness, thus excluding any substrate interferences [16, 17]. In order to obtain a statistical value, at least 20 indentations were carried out on each sample.

 Scratch tests were done with a Revetest by CSM Instruments equipped with a Rockwell indenter with a tip radius of 200 µm. Two critical loads were defined to evaluate the coating adhesion: LCl, corresponding to the detection of the first lateral crack (cohesive failure), and LC2, which is related to the observation of complete substrate exposure (adhesive failure).

Substrate residual stresses were measured by means of X-Stress Analyzer Rigaku Strainflex with CrKα radiation. The stress values were calculated applying the well-known $d$ - sin $^2 \Psi$ method, which considers the 2θ shift of steel (211) peak [18].

Scanning Electron Microscope (SEM) analyses of the coating surfaces were performed with a LEO EVO 40, equipped with an Energy Dispersive Spectroscopy probe (EDS).

Corrosion tests were made in 3.5% NaCl solution with AMEL 7050 potentiostat, considering a potential between −0.8 and +1.5 V, in order to observe both Tafel region and passivation/pitting phenomena. Three tests for each sample configuration were executed. Corrosion current density ($i_0$) and free corrosion potential ($E_{corr}$) were extrapolated by considering the tangent at both cathodic and anodic branches of the potentiodynamic curves. Coating porosity was also calculated according to the method reported in [19].

Finally, the tribological behavior of the samples was studied with a pin-on-disk tribometer (CSM instrument), using alumina balls of 6 mm in diameter as counterparts. The sliding distance and the track radius were set, respectively, at 4 cm/s and 3 mm.

Short tests were carried out with a fixed sliding distance of 100 m and changing the applied loads between 1 and 19 N (limits of the tribometer), in order to define a proper load for the long run test performed until the coating failure occurred. In fact, no precise indications about the load to be used are present in ASTM G 99.

The worn area was monitored during the wear tests without dismounting the sample. In particular, the test was paused every 1000 m and the track profile was acquired, using a stylus profilometer (Tribotechnique) fixed on the tribometer. The worn area was calculated as the average of the measurements in five different positions of each track.

The profile measurements were executed with a tip radius of 5 µm and an applied load of 1 mN. Track length was set at 48 mm. The instrument was equipped with a Diasoft standard software by Mountains Technologies for the direct worn area calculation.

3. Results and Discussion

3.1. Steel Substrate Residual Stress. Figure 1 reports the residual stresses of the steel substrate measured on the samples.

The BS samples show a high compressive stress due to the finishing operations (grinding and polishing). After deposition, the compression decreases probably because of stress relief induced by the high deposition temperature (450°C). Concerning the CrN + MB samples, the microblasting treatment significantly increases the residual stress of the steel substrate, as a function of the applied pressure. Hence, after microblasting, even with the lower pressure, the compressive stress is higher than that of the BS. This effect has been already investigated by the authors in [8].

### Table 1: Chemical composition (wt%) of the steel substrate.

<table>
<thead>
<tr>
<th>C</th>
<th>S</th>
<th>P</th>
<th>Mn</th>
<th>Cr</th>
<th>Ni</th>
<th>Mo</th>
<th>Cu</th>
<th>Si</th>
<th>V</th>
<th>Al</th>
<th>Fe</th>
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<tbody>
<tr>
<td>0.283</td>
<td>0.002</td>
<td>0.008</td>
<td>0.623</td>
<td>0.850</td>
<td>3.065</td>
<td>0.506</td>
<td>0.167</td>
<td>0.312</td>
<td>0.093</td>
<td>0.023</td>
<td>Bal.</td>
</tr>
</tbody>
</table>

**Figure 1:** Residual stress measurements in the steel substrate.
3.2. Morphological Analysis of the Coatings. In Figure 2, the images of the coatings collected by SEM are shown. CrN samples in as-deposited state are characterized by a high presence of droplets, whose size ranges from 1 to tens of micron (Figure 2(a)). The microblasting treatment seems to be effective in removing the larger droplets (Figure 2(b)). It can be also seen that microblasting performed at higher pressure can locally damage the coating, causing cracks and, in some cases, also detachments (Figures 2(c) and 2(d)). It follows that the proper microblasting pressure has to be defined for each specific coating and substrate, in order to have a good compromise between residual compressive stress and coating soundness.

On the base of these observations, subsequent mechanical, corrosion, and wear investigations were performed only on CrN and CrN + MB with a pressure of 2 bar. This pressure, in fact, guarantees the integrity of the PVD coating, contrary to 3 and 6 bar.

3.3. Hardness and Adhesion. Table 2 shows the HV hardness of the coatings obtained by nanoindentation tests and the scratch tests results.

<table>
<thead>
<tr>
<th></th>
<th>CrN</th>
<th>CrN + MB (2 bar)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HV</td>
<td>2009 ± 95</td>
<td>2004 ± 110</td>
</tr>
<tr>
<td>LC1</td>
<td>14.9 ± 1.5</td>
<td>14.4 ± 1.5</td>
</tr>
<tr>
<td>LC2</td>
<td>50.6 ± 1.3</td>
<td>49.9 ± 0.9</td>
</tr>
</tbody>
</table>

No significant variations either in the coating hardness or in the critical loads can be detected as a consequence of the microblasting. These results confirm the data reported by the authors elsewhere on different CrN coatings [8].

It can be concluded that the microblasting treatment at 2 bar is effective in increasing the residual stress of the substrate and in cleaning the CrN surface from the droplets, without appreciably affecting the hardness and adhesion of the coating.

3.4. Corrosion Resistance. Figure 3 shows the potentiodynamic curves obtained for the BS, CrN, and CrN + MB 2 bar samples. As expected, the coated samples revealed a better corrosion behavior. Ceramic coatings, in fact, act as a barrier against aggressive environments, because of their higher nobility than metals, reducing the corroilable areas to those regions where the substrate is exposed (e.g., porosity) [20].

Comparing the two coating conditions, it can be observed that the corrosion current density is lower for the CrN + MB sample of about 15% (see Table 3). It follows that a lower corrosion occurs on the microblasted sample. The corrosion current density is measured during the tests as sum of the effective corrosion of the substrate and that of the droplets, which are known to act as preferential corrosion sites (Figure 4). Therefore, the coating with less droplets (CrN + MB) exhibits a lower corrosion current density.
Table 3: Corrosion tests results.

<table>
<thead>
<tr>
<th></th>
<th>Corrosion current density $i_0$ [A/cm$^2$]</th>
<th>Free corrosion potential $E_0$ [V]</th>
<th>Coating porosity $P$ [%]</th>
<th>Passivation current density $i_p$ [A/cm$^2$]</th>
<th>Rupture potential $E_t$ [V]</th>
</tr>
</thead>
<tbody>
<tr>
<td>BS</td>
<td>$5.8 \times 10^{-6}$</td>
<td>$-0.59$</td>
<td>1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CrN</td>
<td>$1.8 \times 10^{-7}$</td>
<td>$-0.39$</td>
<td>$2.6 \times 10^{-3}$</td>
<td>$7.9 \times 10^{-6}$</td>
<td>0.55</td>
</tr>
<tr>
<td>CrN + MB</td>
<td>$1.5 \times 10^{-7}$</td>
<td>$-0.36$</td>
<td>$2.5 \times 10^{-5}$</td>
<td>$4.9 \times 10^{-5}$</td>
<td>0.37</td>
</tr>
</tbody>
</table>

Moreover, Cr droplets are less noble than CrN; thus, the presence of such defects explains the reduction of the free corrosion potential ($E_0$).

In Table 3, the calculated porosity level ($P$ [%]) is also reported, which represents the fraction of substrate area exposed to the environment. For both CrN and CrN + MB the calculated $P$ [%] results are similar, further confirming that the difference in their corrosion behavior is only related to the different amount of droplets.

Additionally, because the porosity level does not increase in the CrN + MB coating, it can be concluded that no damage was introduced by using the chosen microblasting parameters, as already assessed by the SEM analyses (Figure 2).

3.5. Wear Resistance. Figure 5 shows the friction coefficient obtained during the wear tests as a function of the applied load. The reported values were obtained as the mean value of the friction coefficient recorded during each test after the run-in (about 1 meter). In both the CrN and CrN + MB samples, increasing the applied load the friction coefficient decreases until it reaches stable values around 0.3, in agreement with the literature [21]. The wear mechanism is abrasive for both coatings.

To understand the behavior of the friction coefficient, SEM images of the wear tracks were collected for each test condition. After the wear test was performed at 1N, the droplets are still present and almost undamaged inside the wear track of the CrN samples (Figure 6(a)), notwithstanding the fact that they are softer than the ceramic coating. By increasing the load, as, for instance, at 5N, the droplets appear flattened on the scar surface (Figure 6(b)). For higher loads, no more droplets can be found on the wear path, as already detectable at 9N (Figure 6(c)).
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Figure 6: SEM images of the wear scar of samples CrN at 1, 5, and 9 N, respectively (a, b, c), and of samples CrN + MB at 1, 5, and 9 N, respectively (d, e, f). Sliding distance of 100 m.

In the case of CrN + MB, also at lower loads the wear tracks appear already flat and without droplets (Figures 6(d)–6(f)), thanks to the cleaning effect of the microblasting. In fact, stable friction coefficient values were reached at lower loads than for the CrN samples.

The long run wear tests were done at 9 N, which is a load able to guarantee the achievement of a stable friction coefficient in both CrN and CrN + MB.

Figure 7 reports an example of the friction coefficient measured during the whole tests duration.

It can be clearly seen that the coefficients are similar for both CrN and CrN + MB. In particular, the coefficients rapidly increase from values around 0.3 (as in the 100 m test) to stable values in the range of 0.35–0.45 till the failure. However, the coating breakdown of the CrN samples occurs earlier than CrN + MB. This can be easily identified by the sharp increase in the friction coefficients in Figure 7. The resulting average breaking point distances equal 6690 ± 136 m and 8470 ± 223 m for the CrN and CrN + MB, respectively.

The improved behavior of the coating after microblasting can be also confirmed by the analysis of the worn area (Figure 8).

Before the coating failure, the worn area increases almost linearly with the sliding distance for both coatings but with different slopes (Figure 8). In particular, microblasting seems to reduce the wear during the test.

As already reported, hardness is the same for both coatings (Table 2) as well as the wear track morphologies during tests at 9 N (Figure 6) and the wear mechanism. It follows that the worn area should be almost the same, contrary to the
measurements. This can be explained by the different residual compressive stress, induced by microblasting. As reported in [8], in fact, the compressive residual stress is related to an increased stiffness of the substrate. Consequently, a lower deformation is expected when the substrate is stiffer, that is, in the CrN + MB samples.

Hence, the worn area measured during the experiments does not only depend on the effectively removed material, but also depend on the subsidence of the coating into the substrate. The coating is not able to follow the deformation of the substrate, in particular in the CrN samples, causing the earlier failure of the coating.

4. Conclusions

In this paper the influence of microblasting applied to CAE CrN coatings on corrosion and wear resistance was investigated. A preliminary study allowed fixing at 2 bar the proper microblasting pressure able to clean the surface from the droplets, without inducing cracks or delaminations.

Subsequent corrosion tests showed that, notwithstanding the improved cleanliness of the coating surface, samples after microblasting are less resistant. This is because the corrosion phenomenon occurs more easily where the coating thickness is reduced by the removal of the droplets.

Concerning the wear resistance, a better behavior was assessed as a consequence of the higher compressive residual stress in the steel substrate, that is, higher stiffness, due to the microblasting.

Competing Interests

The authors declare that they have no competing interests.

Acknowledgments

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References


