EFFECT OF TEMPERATURE AND OXIDATION IN CUTTING ZONE ON WEAR OF CUTTING TOOLS WITH MULTILAYER COMPOSITE NANO-STRUCTURED COATINGS AT HIGH SPEED TURNING

M. Volosova1, A. Vereschak1, A. Okunkova1, N. Sitnikov2, N. Andreev3, F. Milovich3, and J. Bublikov4

1Moscow State Technological University STANKIN, Moscow, Russia
2National Research Nuclear University MEPhI (Moscow Engineering Physics Institute), Moscow, Russia
3 National University of Science and Technology "MISIS," Moscow, Russia
4 IKTI RAN, Vadkovsky per. 18-1A, Moscow, 127055, Russia

*Corresponding author; e-mail: dr.a.veres@yandex.ru

Abstract
The article discusses the ability of multilayer composite nano-structured coatings based on the Cr-CrN-(Ti,Cr,Al,Si)N system to resist wear factors activated with increasing temperature, such as diffusion and oxidation processes, as well as spinodal decomposition of the coating material. Investigators were carried out in turning C45 steel at the following cutting conditions: f = 0.2 mm/rev, aΔ = 1.0 mm, vC = 300 m/min. SEM and TEM techniques were used to investigate changes in nano-structures of coatings, as well as diffusion and oxidation processes. Of particular interest was the study of a layer of nanometric thickness on the rake surface of the tool, between the adhered material being machined and the surface nanolayers of the coating. The content of oxygen, iron, and metals the coating was investigated. A lot significant difference was revealed in the content of elements in the layer investigated, depending on the location of the area studied around a crater on the rake face of the tool.

Keywords:
Multilayered nanostructured coatings; wear; metal cutting tools; nanolayer; diffusion processes; oxidation

1 INTRODUCTION
As is known, an increase in cutting speed is associated with a marked increase in temperature in the cutting zone. Thus, an increase in heat resistance of wear-resistant coatings is a significant resource for increasing the cutting speed and, consequently, the overall performance of cutting. Cutting tools with the (Ti,Al,Cr,Si)N coatings are widely used due to their high hardness and wear resistance, as well as considerably high thermal resistance of the above coatings [Chang 2009, Ezura 2008, Ichijo 2007, Zhang 2011, Jung 2013, Kuo 2017, Yamamoto 2003, Yamamoto 2005]. Moreover, the introduction of Si into the composition of the above coatings contributes to an additional increase in thermal resistance, crack resistance, and resistance to brittle fracture due to the formation of nanocomposite structures. It was found [Chang 2009] that when the total content of Al and Si in the (Ti,Al,Cr,Si)N coating was 60 at%, a cubic structure of the NaCl-type was formed, and with an increase in the content of Al and Si, the cubic structure transformed into a hexagonal one. Meanwhile, the maximum hardness was shown by a coating with the cubic structure [Ezura 2008, Ichijo 2007, Zhang 2011, Jung 2013, Kuo 2017, Sobol’ 2012, Metel 2011], and the coating stayed resistant to the oxidation until 1000°C [Yamamoto 2003, Yamamoto 2008]. Investigation into the resistance to thermal oxidation of the coating revealed the formation of a thin oxide layer with the high Si content between the main oxide layer and the coating [Yamamoto 2008, Fominski 2012]. Chang et al. [Chang 2009] considered the (Cr0.36Al0.64Si0.02)N and (Cr0.40Ti0.22Al0.36Si0.02)N coatings with nanocrystalline structures. When the coating was heated up to 900°C, the oxide phases of TiO2, Al2O3, and Cr2O3 were formed on its surface. It was found that the protective oxide layer formed by oxides of Al2O3 and Cr2O3 prevented the diffusion of oxygen into the coating structure. Meanwhile, the oxidation resistance was improved with an increase in the content of (Al,Si)N. Zhang et al. [Zhang 2013] studied the nanolayer (Cr,Al,Si)N coating. An important feature of the coating is the very low friction coefficient (0.1-0.2). The authors explain the above figures by the fact that the coating had a very high cohesive energy of interfacial relationships, while the epitaxial growth created a field of alternating stresses that restrained the movement of dislocations and increased the hardness of the coating. Nguyen et al. [Nguyen 2009] investigated the thermal oxidation of the (Ti,Al,Cr,Si)N coating at the
temperature of 1000°C. It was found that under those conditions, the oxide phases of Cr₂O₃, α-Al₂O₃, SiO₂, as well as TiO₂ (rutile) were formed on the surface of the coating. In that case, TiO₂ was registered predominantly in the outer part of the oxide layer, while Cr₂O₃ and α-Al₂O₃ in its inner part. The content of the SiO₂ phase in the oxide layer was very low due to its low mobility.

Thus, it can be argued that the addition of Si in the (Ti,Cr,Al)N coating contributes to the formation of nanograin structure, increase in hardness, heat resistance, and resistance to thermal oxidation. The formation of nanolayer structure of the coating contributes not only to an increase in its hardness, heat resistance, and resistance to thermal oxidation due to the formation of alternating interlayer interfaces that prevent the oxygen diffusion, but such a formation also provides a rational combination of high hardness with high strength and toughness.

The above features of the coatings based on the (Ti,Al,Cr,Si)N system contribute to an assumption about the prospects of using it as a coating for a cutting tool.

2 MATERIALS AND METHODS

The Filtered Cathodic Vacuum Arc Deposition (FCVAD) [Vereschaka 2013, Vereschaka 2016] technology was implemented using VIT-2 installation (IKTI RAN – MSTU STANKIN, Russia). VIT-2 was equipped with a system for filtration of droplets formed at the evaporation of refractory metals such as Al, Ti, Cr; a mechanism for suppression of micro arcs; a dynamic gas mixing system and a generator of high energy ions.

Samples for metallographic studies were prepared on equipment for metallographic sections manufactured by BUEHLER Company (Isomet 1000 precision cutting machine, Simplimet 1000 automatic hydraulic mounting press, and EcoMet 250 + AutoMet 250 automatic grinder and polisher machine).

An analysis of the microstructure and nanostructure of the specimens was studied using JEM 2100 (JEOL, Japan) high-resolution transmission electron microscope (TEM) with an accelerating voltage of 200 kV. The chemical composition of the samples was studied using an energy-dispersive X-ray spectroscopy (EDXS; INCA Energy; Oxford Instruments, England). Sample preparation for the TEM was carried out using a focused ion beam (FEI Strata FIB 201) with an accelerating voltage of 30 kV.

The tests to study the tool life and wear dynamics for the carbide inserts SNUN ISO 1832:2012 (H10 grade – WC + 5% TiC + 6% Co; KZTS, Russia) were conducted at a CU 500 MRD lathe manufactured by Silven Company with the ZMM CU500MRD variable speed drive in turning workpieces made of AISI C45 steel (400 mm long and an initial diameter of 250 mm). No coolants or lubricants were used in the cutting process. The parameters of the carbide inserts in a cutting tool holder were as follows: γ = −7°, α = 7°, λ = 0°, and r = 0.4 mm. In addition, the cutting conditions were as follows: f = 0.25 mm/rev, a₀ = 1.0 mm, v₁ = 300 m/min. The flank wear land VBₘₙ = 0.45 mm was assumed as a wear criterion. Each test was repeated at least five times; when determining the average wear values, the minimum and maximum values were discarded.

3 RESULTS AND DISCUSSION

The study was focused on the carbide cutting tool with the multilayer composite nano-structured Cr-CrN-(Ti,Cr,Al,SI)N coating which showed its good performance properties in previous studies [Vereschaka 2018, Vereschaka 2019].

The coating structure (Fig.1) included the adhesive Cr layer with the thickness of about 20 nm, the transition CrN layer with the thickness of 300 nm, and the wear-resistant (Ti,Cr,Al,SI)N layer with the thickness of 2.4 μm, which had a nanolayer structure [Vereschaka 2018, Vereschaka 2019, Vereschaka 2016].

With a larger magnification (Fig. 2), it can be seen that in turn, each nanolayer consists of several subnanolayers with the thicknesses of 2-7 nm. The formation of these subnanolayers can be associated with the planetary rotation of the tool set during the coating deposition.

Figure 3 presents the results of the investigation into the cutting properties of the tool with the coating under study in turning C45 steel. The tool with this coating showed a higher tool life period both in comparison with the uncoated tool and with the tool with the commercial monolithic (Ti,Al)N coating.

The dynamics of tool wear with the test coating is shown in Fig. 4a.

To investigate the diffusion and oxidation processes in the cutting zone, as well as the influence of high temperatures on the chemical composition of the coating, the study was focused on the area of a bridge between the crater and the main cutting edge (Fig. 4b). This area is usually associated with the formation of an adherent and a stagnant zone that protect the bridge from wear and destruction.
Fig. 3. Relationship between the flank wear and the cutting time in turning a workpiece made of C45 steel.

Fig. 4. The tool rake wear (a) and 3-D scan of the wear area on the carbide insert with the coating (b) after 21 minutes of cutting.

Fig. 5. Cross-section of the coating (a), nanolayer structure of the wear-resistant layer of the coating (b) partial detachment of the adherent of the material being machined from the coating (c).

Area A (Fig. 6) can be selected in the area under study to examine the chemical analysis of the coating destruction layer. The following distribution of elements in the CDL is typical for the area under study (Fig. 6). The low oxygen presence of a layer of the spinodal decomposition of the coating, referred to as the coating destruction layer (CDL). The thickness of the CDL in the area under study is about 700 nm. There is a clear boundary between the destruction of the nanolayer structure of the coating and the transition to the CDL, and this fact can be considered as an indirect proof of the influence of the internanometer interfaces on the inhibition of the development of the CDL. In Fig. 5c, it is possible to notice partial detachment of the adherent of the material being machined from the coating.
content in the composition of the above layer can be explained by the obstructed access of oxygen to the area in connection with the presence of the inhibited layer. It is possible to note the high content of Al with the low content of Ti and the very low content of Cr. This fact can be associated with the process of separation of Al from the solid solution of (Ti,Cr,Al)N and the formation of the wurtzite AlN phase at high temperatures, as well as with the process of the interface-directed spinodal decomposition, during which atoms of Al move to the nearest interface and accumulate in it [Povstugar 2013]. Meanwhile, the coating nanolayers adjacent to the coating destruction layer are characterized by a lower content of Al and higher contents of Ti and Cr. There is insignificant diffusion of Fe from the material being machined in the coating destruction layer in the area under study.

While considering the boundary area of the CDL (dashed line in Fig. 7), it is possible to notice a fairly clear boundary of the nanolayer structure of the coating, as well as rather large (50-200 nm) crystals which formed the CDL. In the very structure of the CDL, it is possible to note traces of the nanolayer structure conserved fragmentally.

![Fig. 6. Cross-cut of the coating (a) and content of chemical elements (b) in area A.](image)

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![200 nm CDL (Ti, Cr, Al, Si)N](image)

**Fig. 7. Boundary of the decomposition of the coating nano-structure in the area under study.**

4 CONCLUSIONS

The studies of boundaries of crater wear on the carbide tool with the Cr-CrN-(Ti, Cr, Al)N coating revealed a layer with a lost nano-structure, referred to as the coating destruction layer (CDL). The studies also found out the thickness of the layer which reached about 700 nm. Since cutting occurs at a sufficiently high cutting speed \(v_c = 300 \text{ m/min}\), oxidation processes in combination with temperature effects have a significant impact on the wear pattern. The result of this impact is the formation of an extensive crater wear on the rake surface.

The coating is formed by simultaneous exposure to high temperatures, diffusion and oxidation processes, as well as wear processes.

The studies found that the area (CDL) is quite clearly limited by the boundaries of the nanolayer, and this fact suggests that the interfaces between nanolayers restrain the development of the coating destruction layer.

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