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THERMAL-CYCLING BEHAVIOR OF CoNiCrAIY BONDS COATED WITH THERMAL BARRIER COATINGS (TBCs) PRODUCED WITH ATMOSPHERIC PLASMA SPRAYING (APS)

OBNAŠANJE CoNiCrAIY PREVLEKE MED TERMIČNIM UTRUJANJEM

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Thermal barrier coatings (TBCs) are commonly applied as a thermal insulation in order to protect against environmental influences in the components of high-temperature gas turbines and jet engines. Gas-turbine components are affected by aggressive conditions of the environment during the service, being subjected to failures such as corrosion, thermal shock and oxidation. Plasma-spray technology is used to produce metallic bonds and ceramic top coats as a cost-effective method to prolong the lifetime of TBCs. In the present research, CoNiCrAIY bond-coat and YSZ top-coat powders included in the TBCs were deposited onto Inconel 718 superalloy substrates using atmospheric plasma spraying (APS). The TBCs were exposed to a furnace cycling test at 1150 °C and one-hour cycles. The presence of porosity and cracks facilitating the diffusion of oxygen in the top-coating structure led to the formation of thermally grown oxides (TGOs) at the interface. In addition, the most effective factors of failures were the formation and growth of mixed oxides at the bond/top-coat interface and the presence of imperfections. According to the test results, an increasing number of thermal cycles resulted in a decrease in the lifetime of the TBCs related to the sintering of the top coating due to the effect of high temperature.

Keywords: thermal barrier coatings (TBCs), thermal cycling, Inconel 718 superalloy, atmospheric plasma spraying (APS), CoNiCrAlY, ytrria-stabilized zirconia (YSZ)

Prevleke s toplotno pregrado (angl. TBCs) se običajno uporabljajo kot toplotna izolacija, ki ščiti komponente visoko temperaturnih plinskih turbin ali reaktivnih motorjev. Na komponente plinskih turbin vplivajo agresivni vplivi okolja, katerim so izpostavljene med obratovanjem, zato pride do okvar, kot so: korozija, termični šok in oksidacija. Tehnologija plazemskega naprševanja je cenovno ugodna tehnologija za izdelavo kovinskih in keramičnih prevlek. V članku avtorji predstavljajo raziskavo v kateri sta bila na podlago iz zlitine Inconel 718 s tehnologijo plazemskega naprševanja v zračni atmosferi nanešena vezivna prevleka CoNiCrAlY in prekrivni sloj iz YSZ prahu. Izdelani prevleki sta bili nato izpostavljeni termičnem utrujanju v peči pri 1150 °C z večimi enournimi cikli. Prisotnost poroznosti in razpok olajša difuzijo kisika v zgornjo plast prevleke kar vodi do nastanka termično induciranega oksida (TGO) na površini prevleke. Poleg tega sta bila najpomembnejša dejavnika za nastanek napak tvorba mešanih oksidov na meji med vezno in vrhno plastjo ter prisotnost drugih nepravilnosti. Rezultati preizkusov so pokazali, da povečanje števila termičnih ciklov zmanjšuje dobo trajanja TBC prevlek, zaradi sintranja zgornje plasti pri povišanih temperaturah.

Ključne besede: prevleke s toplotno pregrado, toplotni (termični) cikel, superzlitina Inconel 718, plazemsko naprševanje, CoNiCrAlY, cirkon stabiliziran z itrijem

1 INTRODUCTION

Atmospheric plasma spraying (APS) is one of the most important methods in the thermal-spray family. The APS technique, which is a type of the plasma spray technique, provide many advantages, such as an economic, quick and easy production; besides, almost all materials can be sprayed with this method. Plasma-sprayed particles quickly cool down from very high temperatures to low temperatures. Plasma-spray coatings consist of microstructured, layered and porous structures.^{1,2}

Thermal barrier coatings (TBCs) are widely employed in gas turbines and its components operating at high-temperature service conditions. The main function of TBCs is to provide a protective thermal shield to protect the main material components, especially turbine

blades and vanes, from the negative effects caused by high temperatures in the environment. There are three main layers in TBC systems, including a nickel-based superalloy substrate, an MCrAlY metallic bond coat and partially yttria-stabilized zirconia (YSZ), acting as the ceramic top coat.^{3,4} The bond coat, which consist of MCrAlY (M = Ni and/or Co), is extensively applied to minimize the thermal mismatch between the superalloy substrate and the ceramic top coat.^{5,6}

TBCs are subject to sudden temperature changes and thermal stress during service conditions. In high-temperature applications, TBCs are adversely affected and tend to suffer because of failures such as oxidation, hot corrosion and thermal cycling/shock so that they lose structural integrity. Residual stresses due to the thermal-expansion mismatch during the cooling and sintering of

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the top coat, and oxidation of the bond coating cause the coating to become damaged during thermal cycling.^{7,8}

In this study, the TBC system consists of a CoNiCrAlY metallic bond and YSZ ceramic top coat deposited with the APS technique on an Inconel 718 superalloy substrate. The effects of oxidation and TGO-growth behavior on the thermal durability were investigated through a furnace thermal cyclic test. The microstructure evolution and thermal durability were evaluated before and after the furnace thermal-cycling tests.

2 EXPERIMENTAL PART

2.1 Material

The Inconel 718 nickel-based superalloy was used as the substrate material. Superalloy disk-shaped samples with a diameter of 25.4 mm and thicknesses of 4 mm were used as substrates. The chemical composition of the nickel-based superalloy substrate is given in **Table 1**.

Table 1: Chemical composition of Inconel 718 Ni-based superalloy⁹

Elements, Chemical Composition, in mass fractions (w/%)							
Ni	Cr	Nb	Ti	Mo	Al	Co	Si
53.53	18.45	5.31	0.95	3.02	0.48	0.27	0.09
Cu	Mn	C	Ta	P	В	S	Fe
0.04	0.06	0.03	0.005	0.007	0.005	0.001	Balance

2.2 Spraying processes

Commercial CoNiCrAlY metallic bond-coat powder (Amdry 9951, Sulzer-Metco) with a 5–37 μm particle size and ZrO $_2$ + 8 %Y $_2$ O $_3$ (YSZ) ceramic top-coat powder (Metco 204NS, Sulzer Metco) with a -45+20 μm particle size were used as the feedstocks. As previously mentioned, the APS technique was used for depositing both the metallic bond and ceramic top coatings. The top layer of YSZ with a thickness of 300 μm was applied on the CoNiCrAlY layer with a thickness of 100 μm on the superalloy substrates.

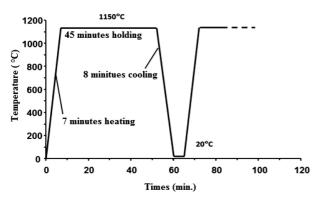


Figure 1: Thermal-cycling diagram during the furnace cycling test

2.3 Furnace thermal-cycling test

The lifetime of the TBCs tested with the furnace thermal-cycling test using rapid temperature furnace 1710 BL was as follows: The samples were heated to 1150 °C in 7 min, held at this temperature for 45 min and then cooled to 20 °C in air within 8 min. The thermal-loading diagram of the samples coated during thermal cycling is given in **Figure 1**. During isothermaloxidation cycles, the temperature changes in the working furnace did not exceed 5 °C. The coating lifetime was determined with the thermal cycling carried out until the cracking point of the outer ceramic layer. This was considered to be cracked when 25 % of the sample surface was blistering. The surface failure was visually determined based on the percentage of the separation of the outer ceramic layer from the sample surface. At the end of the thermal-cycling tests, the samples were investigated from both the top surface and cross-sectional area via a scanning-electron-microscopy (SEM) analysis.

3 RESULTS AND DISCUSSION

3.1 As-sprayed TBC

A SEM microstructure image of an as-deposited TBC is shown in **Figure 2**. The APS-applied CoNiCrAlY metallic bond and the YSZ top coat exhibit a porous structure and some cracks and open discontinuities due to the deposition process. The metallic-bond and top-coat structures show typically characteristic properties of the plasma-spray technique.

In **Figure 3**, the top surface of the TBC is shown. There are some micro-cracks due to heating and rapid cooling of the sprayed particles during the deposition process as well as melted and unmelted particles.

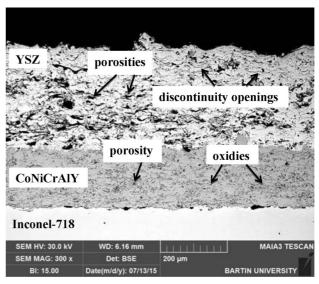


Figure 2: As-deposited TBC cross-sectional SEM microstructure

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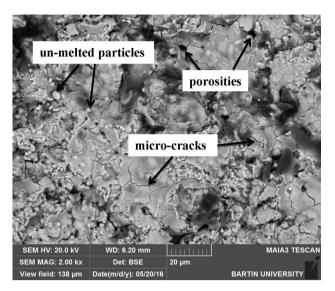


Figure 3: Top-coat SEM image of the as-sprayed TBC

3.2 Thermal-cycling test of the coated specimens

At high temperatures, oxygen can easily diffuse from the top coat to the bond coat due to the ionic conductivity of YSZ. Also, the coating includes many imperfections such as cracks and porosity, which accelerate the penetration of oxygen to the metallic bond coat. As a result, the bond coat oxidizes easily at high temperatures in spite of the ceramic coat. According to the Ellingham diagram, Y and Al are the elements that enter the first reaction with oxygen in the CoNiCrAlY bond coat due to low-standard free energies. With the effect of high temperature, the particles are still oxidized in flight and during the deposition on the substrate material.¹⁰

Using the APS method, there are many internal oxide formations in the coating because of the excessively high temperatures due to plasma during the deposition of the coating. Since the concentration of the Y element is much lower than those of the other elements, Y can be entirely depleted. Thus, after the deposition of a TBC, Al is oxidized at the interface and Al_2O_3 is formed. 11–13

In the early stages of the cyclic oxidation, the TGO layer mainly occurred as the Al₂O₃ phase. This oxidation causes thermal stresses at the interface between the bond and the top coat due to both the phase transformations of Al_2O_3 (γ , δ , θ and α) and a lower thermal-expansion coefficient compared to YSZ. Al₂O₃ (especially in the α -Al₂O₃ form) is thermodynamically a very stable phase compared to the other phases such as Cr₂O₃, (Co, Ni)O or (Co,Ni)(Al,Cr)₂O₄.¹³ In addition, it provides a better resistance against the oxygen expansion into the bond coat. Depending on the time, the Al concentration decreases to a critical level and Cr₂O₃ and (Co, Ni)O phases form at the interface. Afterwards, these oxides react with Al₂O₃ and spinel phases occur. The thermal-expansion coefficient of the spinel phases is relatively lower than that of Al₂O₃, which means it leads to a higher stress. 14,15

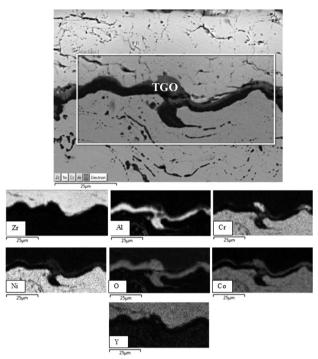


Figure 4: Elemental mapping microstructure belonging to the YSZ TBC after a 40-h thermal-cycling test

In addition, the spinel phases grow faster than Al_2O_3 and show larger volume changes.¹⁶

In **Figure 4**, an elemental mapping microstructure belonging to the YSZ TBC after a 40-h thermal-cycling test is shown. The coating structure almost preserved its uniformity. At the interface, the Al₂O₃ phase is dominant and a mixture of oxide formations like Cr₂O₃, (Co, Ni)O and spinels can be observed. In the bond coat, dark stringers represent the Al₂O₃ phase. In addition, the top coat was sintered due to the effect of high temperature

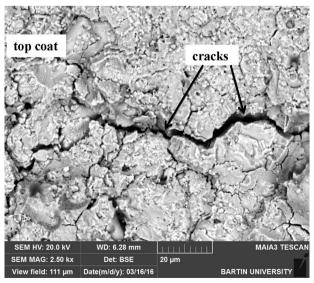


Figure 5: Top-surface SEM image of the TBC after the thermal-cycling test

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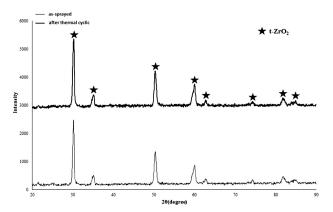


Figure 6: XRD analysis of the TBC before and after the thermal-cycling test

depending on time. This can be explained with decreasing discontinuity openings.

In **Figure 5**, a top-surface SEM image of the TBC after the thermal-cycling test is given. After 40 thermal cycles, the top-surface crack area of the TBC covers approximately 25 %. In **Figure 5**, a failed surface of the TBC is shown. According to Figure 5, there are large crack formations and sintered areas.

Figure 6 shows that there is only one phase in the as-sprayed TBC. After the thermal-cycling test, there is not any phase transformation so the structure is similar to the as-sprayed TBC. The semi-stable tetragonal ZrO₂ phase is durable up to approximately 1200 °C, which means that the crack propagation does not progress into the bond coat.⁸ This can also be understood from a cross-sectional SEM image of the failed TBC.

J. A. Haynes et al.¹⁷ applied furnace thermal-cycling tests to YSZ TBCs including a NiCrAlY bond coat produced with the APS method. The tests were carried out for one hour in a furnace at 1150 °C with periods of 15-min heating and 30-min cooling. The test result shows that after 100 thermal cycles, the APS TBCs failed. However, in the present study, the heating and cooling times of the thermal-cycling tests are shorter. Thus, the CoNiCrAlY TBC is exposed to damage earlier than in the above study. R. Chen et al. used the APS technique to produce a CoNiCrAlY bond coated with the YSZ TBC on an Inconel 625 substrate. Their thermal-cycling test at 1050 °C showed a parabolic TGO growth. The growth rate of the TGO structure increased proportionally as the thermal-cycling period increased. This result supports the present study.

A. El-Turki et al.¹⁸ studied the thermal-cycling behavior of a VPS NiCoCrAlY bond coated with APS YSZ at 1050 °C, with 25-min heating, 300-h holding and 30-min cooling. The TBCs were damaged after 21 cycles. The damage of the TBCs started at the same time as the damage to TGO interface areas. The use of the VPS technique allowed a low porosity and oxidation during the deposition of the bond coat, and their total holding time was very long compared to the present study.¹⁹ J. Sun et al.²⁰ produced their bond coat using the HVOF technique; afterwards they carried out the deposition of

APS YSZ. Their thermal cycling was performed at 1100 °C for one hour, with 10-min cooling. After 30 thermal cycles, the TBC underwent a significant failure. In contrast, the present-study condition is more destructive, but the thermal-cycling lifetime is longer. As a result, the used bond-coat powder type, production technique and thermal-cycling conditions can cause significant differences in the lifetime of TBCs.²⁰ In the present study, the APS TBC separated from the bond-coat surface of the ceramic top coat after 40 h of the thermal cycling and was spalled and damaged. Furthermore, a TGO structure formed during the thermal cycling, causing failure of the TBC systems. The growth of the oxide layer formed the TGO structure and an increase in the stresses increased the failure formation.

4 CONCLUSIONS

In this study, TBCs including CoNiCrAlY bonds and YSZ were deposited using the APS technique. Furnace thermal-cycling tests were performed to predict the thermal-cycling lifetime of the TBCs and to observe their microstructural changes. The results obtained at the end of the observations are given below.

TBCs were damaged, with their cracks covering 25 % of the top coat after 40 cycles.

The most effective factor of failure was the formation of mixed oxides and spinels at the interface of the bond and the top coat, according to the cross-sectional investigations.

The porosity and cracks of the top coat caused the formation of TGOs at the interface.

The top coat was sintered as a result of high temperature depending on time.

At the interface, the Al_2O_3 phase is dominant, but a mixture of oxide formations like Cr_2O_3 , (Co, Ni)O and spinels is also observed. In the bond coat, dark stringers represent the Al_2O_3 phase. The mixture of oxides formed at the interface was mainly responsible for the failure of the TBCs after the thermal cycling.

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