## Microstructure characterisation of double glass-ceramic coatings on TIMETAL 834

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TIMETAL 834, a near- $\alpha$  titanium alloy, offers increased tensile strength and creep resistance up to 600°C together with improved fatigue strength [1]. This alloy is applied for fabrication of last stages compressor parts in modern gas turbines [2]. One of the major limitations of TIMETAL 834 alloy restricting its use at high temperatures is a poor oxidation resistance at temperature above 600°C. To improve it, a surface treatment is frequently applied. Glass-ceramic based coatings are very promising materials for protection of titanium alloys against oxidation at high temperature [3]. Their processing by a simple and low cost slurry method is an excellent method to manufacture reproducibly protective coatings with complex and homogenous chemical composition, uniform morphology and coating thickness [3,4].

The aim of the present work was to characterise a microstructure and phase composition of the double glass-ceramic coating produced on TIMETAL 834 alloy by slurry technique. The coating was characterised by SEM and TEM (SAED, EDS, STEM-EDS). TEM investigations were performed using JEM-2010 ARP on cross-section thin foils prepared by PIPS. The SAED patterns were interpreted with the JEMS software [5].

The coating consists of two layers with a different microstructure and phase composition: outer SiO<sub>2</sub>-P<sub>2</sub>O<sub>5</sub>-B<sub>2</sub>O<sub>3</sub>-CaO-K<sub>2</sub>O-Na<sub>2</sub>O layer (about 140  $\mu$ m thick) and intermediate SiO<sub>2</sub>-Al<sub>2</sub>O<sub>3</sub>-CaO-Na<sub>2</sub>O one (about 215  $\mu$ m thick) (Figs 1a,b). Several crystalline phases were found in the outer layer: mainly SiO<sub>2</sub> (tetragonal primitive), Na<sub>2</sub>Ca<sub>3</sub>Si<sub>6</sub>O<sub>16</sub> (triclinic primitive) and CaSiO<sub>3</sub> (triclinic primitive). Additionally an amorphous phase, containing mainly Si and O as well as some Ca and K, was identified. The fine particles (50-100 nm in size) of the Ca<sub>5</sub>P<sub>2</sub>Si<sub>1</sub>O<sub>12</sub> (orthorhombic primitive) in SiO<sub>2</sub> grains were observed. Also some amorphous particles containing mainly Si and O (up to 100 nm) in amorphous phase were found. The CaSiO<sub>3</sub> grains in amorphous matrix are shown on Fig. 2a. The intermediate layer is composed of the crystalline phases: CaSiO<sub>3</sub> (monoclinic primitive), Na<sub>2</sub>Ca<sub>3</sub>Si<sub>6</sub>O<sub>16</sub> (sporadically) and amorphous phase (containing mainly O, Si, Al and some Ca). The NaAlSiO<sub>4</sub> grains in amorphous matrix are shown on Fig. 2b. The microstructure of the asreceived alloy consists mainly of the  $\alpha$  phase (hcp), a small amount of the  $\beta$  one (body-centred cubic, bcc) and precipitates of the Zr<sub>5</sub>Si<sub>4</sub> phase (tetragonal primitive) [6].

The applied coating has a good adhesion to the substrate and significantly improves the oxidation resistance of the TIMETAL 834 alloy at 800°C. No weight change of the coated alloy was observed after oxidation during 200 hours at 800°C, in contrary to a significant weight change for the uncoated specimens.

- 1. M.J. Thomas et al., Mater. Characterisation 55 (2005) p388.
- 2. W. Kaysser, Surf. Eng. 17 (2001) p305.
- 3. T. Moskalewicz et al., Surf. Coat. Tech. in press, DOI: 10.1016/j.surfcoat.2009.02.016
- 4. F. Smeacetto et al., Carbon 47 (2009) p1511.
- 5. P. Stadelmann, (2004) JEMS Java Electron Microscopy Software, http://cimewww.epfl.ch.
- 6. T. Moskalewicz et al. J. Microscopy 223 (2006) p195.
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**Figure 1.** Cross-section of an outer (a) and an intermediate (b) layer of double glass-ceramic coating on TIMETAL 834 substrate, SEM electron back scattered images.



**Figure 2.** Microstructure of the outer (a) and intermediate layer (b) of the double glassceramic coating on TIMETAL 834 as well as SAED patterns and their identification, TEM, cross-section thin foil.