1 INTRODUCTION

Ceramic materials have been used for centuries as the materials with a very wide range of mechanical and thermal properties suitable for various applications. Today, their common industrial use is more related to the electronics for mechanical parts and biomedicine applications (hip prosthesis, dental implants). One of the main reasons for this is the improvement of the fracture toughness that allows a ceramic material to perform better when it is subjected to the operating conditions. Sometimes these operating conditions may involve rapid changes in the temperature, or a high-temperature application as well as the conditions that include cavitation erosion. This could be related to the technical ceramics, for example, the operations of bearings, injectors or valves. Cavitation, i.e., the appearance of vapor cavities inside an initially homogeneous liquid medium, occurs in very different situations. Hence, the study of the cavitation and cavitation-erosion mechanisms of technical ceramics is of importance to improve their performance in real applications.1–4

Metallic materials are the most common choice when cavitation-erosion resistance is required. Research results concerning the applications of different classes of materials, including ceramics and composite materials for a similar use, were published in the last decade.1–10 Different types of ceramics based on silicon nitride and zirconia, as well as alumina-based ceramic materials were investigated in the conditions of cavitation erosion.2,3,4,9 In numerous papers1–12 related to the investigations of the erosion rate of these materials, attempts were made to...
investigate the type of cracking as well as the influence of the grain sizes and the contents of different compounds and phases on the resistance of the materials to the erosive wear.

The goal of our investigation was to study the possibilities of using a silicon carbide/cordierite based ceramic material as a potential cavitation-resistant material.

2 MATERIALS

The properties of cordierite/silicon carbide based ceramics that enable a widespread use in high-temperature conditions are a particularly low coefficient of thermal expansion, a high thermal conductivity, an excellent thermal fire resistance and an excellent resistance to a thermal shock. Their target application is in the furnaces for the use at the temperatures of over 1000 °C. An addition of silicon carbide improves the physical and mechanical properties of refractory cordierite. The high thermal conductivity of silicon carbide reduces the thermal stresses within a cordierite ceramic body. A mixture of commercially available spinel (MgAl₂O₄), quartz (SiO₂) and alumina (Al₂O₃) corresponding to a cordierite stoichiometric composition was attrition milled using Al₂O₃ balls and ethyl alcohol as the media for four hours; henceforth, this mixture was labeled as KS. Mixture KS was used for preparing the cordierite/SiC composite ceramics with the mass ratio of 30 : 70 and 50 : 50, respectively. After the milling with the Al₂O₃ balls in DI water in a polyethylene bottle for 24 h, the samples were uniaxially pressed and sintered at 1300 °C and 1250 °C, for 3 h, and the ceramic composite samples were marked as KS 30 and KS 50, respectively. The typical values of the selected properties of the dense constituents used in the refractory materials investigated are listed in the previous papers.13–15

An XRD analysis of the KS 30 sample (Figure 1) shows a presence of three crystalline phases: SiC, cordierite and sapphirine and a presence of a certain amount of a glassy phase. In the case of sample KS 50, the phase analysis shows a presence of four crystalline phases: SiC, cordierite, sapphirine and enstatite. There is also a small amount of a glassy phase, which affects the formation of cordierite and sapphirine. One can conclude that an increase in the added amount of the mixture with a composition corresponding to stoichiometric cordierite improves the quantity and crystallization of cordierite.

The microstructures of samples KS 30 and KS 50 (Figure 2) were examined with scanning electron micro-

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**Figure 1:** XRD of the: a) KS 30 and b) KS 50 samples
**Slika 1:** XRD-posnetka vzorcev: a) KS 30 in b) KS 50

**Figure 2:** SEM of the: a) KS 30 and b) KS 50 samples
**Slika 2:** SEM-posnetka vzorcev: a) KS 30 in b) KS 50
scopy using a VEGA TS 5130 mm (TESCAN) device; the samples were coated with a layer of the Au-Pd mixture. A porous structure was observed, with the particles of different sizes and pronounced necks formed during the densification. The pores that are not spherical indicate the initial stages of sintering. The pores present have different sizes and shapes.

3 CAVITATION-EROSION TESTING

The experimental methodology used for the cavitation-erosion testing is explained in7,10,16,17. With respect to the used equipment, the diameter of the horn was 10 mm and the distance between the horn and a sample was 1 mm. The samples were discs with a diameter of 3 cm and a height of 1 cm.

4 RESULTS AND DISCUSSION

4.1 Mass loss and level of destruction during the testing

The mass losses of the test specimens were determined using the analytical balance with an accuracy of ± 0.1 mg. The measurements were performed after subjecting each test specimen to cavitation for 30 min. The duration of the tests was 150 min. The light microscopy technique was applied to analyze the effect of the erosion and to interpret the results of the cavitation tests (Figure 3).

The mass loss was below 1.3 mg, which is similar to metallic15-25 and ceramic5,6,24,25 materials. From Figure 2 it can be seen that sample KS 30 exhibited a better resistance to erosive wear, as after 150 min the mass loss for KS 30 was 0.12 mg and for KS 50 it was 1.23 mg.

The next experiment consisted of the measurements of the degradation levels before and during the testing. In order to carry out an image analysis determining the level of destruction before and during the testing, the samples were photographed as shown in Figure 4.

The results of the image analysis of samples KS 50 and KS 30 are given in Figure 5.

Based on the obtained results, similar conclusions regarding the level of degradation and the mass loss of the samples can be given. Samples KS 50 exhibited a better erosion resistance as their level of damage was 4.4 % compared to 6.9 % for the KS 30 samples. The differences in the sample behavior can be related to the crack nucleation and propagation during the cavitation testing.

As can be seen from Figure 5, samples KS 50 exhibited a stable increase in the level of degradation and similar rates of degradation for the entire time range. At
the end of the experiment, after 150 min, the level of degradation was below 5%.

For the KS 30 samples, two regimes can be observed on Figure 5. The first regime is characterized by a slow increase in the level of destruction over the period from 0 min to 60 min. The second regime, after 60 min of the testing, is characterized by a more rapid change in the damage level. However, it has to be mentioned that the total level of degradation, after 150 min of the testing, did not exceed 8%, which is an excellent result for the cavitation resistance.

From these results it can be concluded that samples KS 30 have a better resistance to crack nucleation and propagation up to the 60th minute of the experiment. After this critical time, the crack propagation is very rapid. The increased cordierite amount and decreased SiC amount influenced the crack nucleation and propagation, as the higher values of cordierite in samples KS 50 caused lower rates of crack propagation during the entire time (0–150 min) and the total degradation of the samples after 150 min of the testing was lower.

4.2 Erosion-ring diameter and erosion-ring area

The samples exposed to cavitation erosion (Figure 3) were also monitored in order to measure the diameters and the areas of the erosion-ring regions. These measurements were performed using the following steps:

Step 1: Making appropriate micrographs that include the erosion ring areas (Figure 3)
Step 2: Measuring the ring diameters: at this point two types of measurements were taken: diameters \( d_1 \) and \( d_2 \) of the cavitation-ring erosion measured with the Image-Pro Plus program. The effective diameter of each ring is calculated according to the following equation:

\[
\frac{d_1 + d_2}{2}
\]  

Step 3: Calculating the effective area of the erosion ring using the values of the diameters measured in Step 2:

\[
P_{\text{eff}} = \frac{\pi d_1 d_2}{4}
\]

Step 4: Using the Image-Pro Plus program to determine the average erosion-surface area of the ring, \( P_{\text{meas}} \).

4.3 Erosion-ring measurements

The measurements of the average diameter of the erosion ring were made after (30, 45, 60, 90, 120 and 150) min.

Samples KS 50: The results presented in Figure 6 show that the formation of an erosion ring can be observed after 45 min of the testing. The average erosion-ring diameter increased from 5.68 mm after 45 min to 6.03 mm after 150 min of the testing.

Samples KS 30: The erosion-ring formation of this sample was observed after 30 min. A rapid increase in the ring diameter was observed after 90 min, when the ring diameter increased from 3.29 mm after 60 min to 7.04 mm after 90 min of the testing. After 90 min the increase in the ring diameter was very slow so that after 150 min the ring diameter reached only 7.41 mm.

The results for the erosion-ring measurement are in a strong correlation with the results for the level of destruction (Figure 5) supporting the conclusions about the influence of cordierite and SiC content on the level of degradation. The same conclusions are valid if the erosion-ring measurements are taken into account. With respect to the energy for crack nucleation, lower values are expected for the KS 30 samples where the formation of the erosion ring was observed after 30 min, while for the KS 50 samples it was formed after 45 min. This difference could be important for some specific applica-

![Figure 6: Average diameter of the erosion ring during the testing](image)

![Figure 7: Average area of the erosion ring based on diameter measurements and area measurements](image)
4.4 Average erosion-surface area

According to the procedure described above, Figure 7 shows the values obtained for the average erosion-surface areas based on monitoring the erosion-ring diameters.

Samples KS 50: The formation of the erosion ring was detected after 45 minutes and the average erosion-surface area was $P_{\text{meas}} = 20.73 \text{ mm}^2$ and $P_{av} = 25.23 \text{ mm}^2$. Both values for the erosion ring area were slowly increased, reaching the values of $P_{av} = 28.35 \text{ mm}^2$ and $P_{\text{meas}} = 33.42 \text{ mm}^2$ after 150 min of the testing. The results showed the differences caused by implementing the method for the erosion-ring-area determination (during the above steps) and these differences apply to the whole experiment.

Samples KS 30: As for the ring-diameter measurements, after 60 min of the testing the erosion ring was detected, with the erosion-surface area of $P_{\text{meas}} = 10.56 \text{ mm}^2$ and $P_{av} = 8.49 \text{ mm}^2$. After 90 min a rapid change in the diameter and the surface of the erosion ring, but by the end of the testing (150 min) this increase is slowed down.

Based on the obtained results it can be concluded that samples KS 30 are more sensitive to the formation of the erosion ring. Also, after 90 min of the testing there are rapid changes in the diameter and the surface of the erosion ring, but by the end of the testing (150 min) this increase is slowed down.

Samples KS 50 exhibited a better erosion resistance as the ring diameter and the average area of erosion were lower, and their increase was slower over the testing time.

The formation of the erosion ring for the KS 50 samples is visible after 30 min. This can be related to the influence of the crack nucleation caused by the erosion experiment.

5 CONCLUSION

Ceramic composite samples based on the SiC/cordierite ceramic material were synthesized in order to investigate the sample resistance to the erosive wear. Two samples with different mass contents of cordierite and SiC were used in this investigation: KS 50 ($w = 50 \%$ of cordierite) and KS 30 ($w = 30 \%$ of cordierite).

The results for the mass loss as well as for the level of degradation indicate a similar (or even better) erosion resistance compared to the metallic and low-alumina samples.

Both samples exhibited an excellent erosive resistance, but sample KS 50 exhibited the better resistance of the two as its mass loss was lower, as were the level of degradation, the erosion-ring diameter and the erosion area.

These experiments showed that composite ceramic materials based on SiC/cordierite can be used as erosion-resistant materials, used for new applications.

Also, an implementation of non-destructive testing such as an image analysis to determine the erosion-ring diameter and the erosion-ring area improved the reliability of predicting the sample behavior in the conditions of erosive wear.

Acknowledgement

This research was financed by the Ministry of Education, Science and Technological Development of Serbia as part of project III 45012.

6 REFERENCES


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