NASA Technical Memorandum 88870

(NASA-TM-888/0) NDE RELIABILITY AND PROCESS CONTROL FOR STRUCTURAL CERAMICS (NASA) 17 p CSCL 14D N87-12910

Unclas G3/38 44727

NDE Reliability and Process Control for Structural Ceramics

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Prepared for the 32nd International Gas Turbine Conference and Exhibition sponsored by the American Society of Mechanical Engineers Anaheim, California, May 31—June 4, 1987



NDE RELIABILITY AND PROCESS CONTROL FOR STRUCTURAL CERAMICS

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ABSTRACT

The reliability of microfocus x-radiography and scanning laser acoustic microscopy for detecting microvoids in silicon nitride and silicon carbide was statistically evaluated. Materials- and process-related parameters that influenced the statistical findings in research samples are discussed. The use of conventional x-radiography in controlling and optimizing the processing and sintering of an Si_3N_4 -SiO₂-Y₂O₃ composition designated NASA 6Y is described. Radiographic evaluation and guidance helped develop uniform high-density Si_3N_4 modulus-of-rupture bars with improved four-point flexural strength (857, 544, and 462 MPa at room temperature, 1200 °C, and 1370 °C, respectively) and reduced strength scatter.

INTRODUCTION

Advanced nondestructive evaluation (NDE) techniques to reliably detect critical flaws are needed if fracture mechanics principles are to be applied effectively in the design of ceramic heat engine components (1,2). These techniques are also needed to monitor and control the fabrication process in the materials development program to make stronger and more reliable ceramics (2). Reliable ceramics require fabrication process control to reduce the incidence of various flaws and to ensure that any flaws that occur are noncritical. The dominant critical flaw type in monolithic ceramics is a pore (3.4.5), which can cause wide strength variations and unacceptably low strength. Recent studies at the NASA Lewis Research Center (6.7) have established the reliability of microfocus x-radiography and scanning laser acoustic microscopy for detecting seeded internal voids in Si_3N_4 and SiC test specimens. Preliminary radiographic characterization of sintered Si_3N_4 specimens (2) revealed the presence of large density gradients (a high-density case with a low-density core structure), which were believed to be detrimental to strength properties. Therefore a program was undertaken at NASA Lewis (3) to incorporate conventional x-radiographic techniques in an extensive investigation of one Si_3N_4 -SiO₂-Y₂O₃ composition in order to monitor its fabrication process.

This paper describes the materials- and process-related factors that affected the reliability of microfocus x-radiography and scanning laser acoustic microscopy for detecting internal voids in Si_3N_4 and SiC. The capability of the conventional x-radiographic technique in guiding powder processing and sintering parameter changes to improve the absolute density of sintered Si_3N_4 and to eliminate detrimental density variations is emphasized. Improvements in flexural strengths and reductions in strength scatter are also described.

MATERIALS AND PROCEDURES

The preparation of SiC and Si₃N₄ specimens needed for the NDE reliability determinations is shown in Fig. 1. The Si₃N₄ powder had the Si₃N₄-SiO₂-Y₂O₃ composition. The SiC powder contained sintering aids and binder materials of boron and carbonaceous resins. Styrene divinyl benzene microspheres of various sizes (50, 80, 115, 220, and 528 μ m diam) were seeded in green specimens and later burned out to create voids within the green and sintered specimens. The seeded test bars had the same densities, compositions, and shapes as typical modulus-of-rupture (MOR) bars of the same materials. Details on specimen fabrication, void characterization, dimension of specimens and voids, and density determination are given in Ref. 6. Microfocus radiography (Fig. 2) and scanning laser acoustic microscopy (Fig. 3) were used to detect the seeded voids (20 to 477 μ m in diam) in the fabricated ceramic samples. References 6 and 7 describe in detail the radiography and SLAM systems, respectively.

The powder-processing steps of the NASA 6Y SigNa composition, containing 6 wt % each of Y_2O_3 and SiO_2 , are shown in Fig. 4. Powders were milled (grinding time, t_g , 24, 100, or 300 hr), die pressed into bars at 21 MPa, and then isopressed at 414 MPa. A total of 690 bars (23 batches) were sintered 15 at a time at 2140 °C in a tungsten cup. High-purity BN disks separated the bars from one another and from contact with the tungsten cup. Sintering times t_s were 1, 1.25, 1.5, and 2 hr. Nitrogen overpressures P_N of 2.5, 3.5, and 5.0 MPa were employed. All batches were radiographically evaluated at all stages of fabrication as described in Fig. 5. Test bars were radiographed (Fig. 6) in two modes: (1) the (W,L) mode, where x-rays are transmitted through the thickness of the bar, and (2) the (T,L) mode, where x-rays are transmitted through the width of the bar. Radiographic feedback on the density uniformity of the bars was used to adjust the powder-processing and sintering parameters. Four-point flexural strength tests were conducted in air at room and elevated temperatures (1200 and 1370 °C) with inner and outer spans of 9.53 and 19.05 mm, respectively. Strength data and radiographic density data were used to further modify the processing and sintering variables.

NDE RELIABILITY

Reliability assessment of microfocus x-radiography and scanning laser acoustic microscopy is probabilistic because of the combined uncertainties associated with the equipment, the operator, the flaw characteristics, etc. Therefore a statistical approach must be used to determine the detection reliability. This examination of the seeded specimens was based on either detecting or not detecting known existing voids. Since only two outcomes from this examination were possible, the probability of detection (POD) can be described by a binomial distribution. Detection data were grouped into size intervals and further rearranged by the optimized probability method (8) to calculate the POD values at the 0.95 confidence level.

Microfocus X-Radiography

NDE reliability data are presented in Figs. 7 and 8 in the form of plots of POD versus void size, expressed as a percent of total specimen thickness.

For green materials the POD curves in Fig. 7(a) indicate that the detection sensitivity of internal voids in SiC was about the same as in SigN₄ (i.e., 2.5 percent of thickness at a POD of 0.9). However, surface voids were easier to detect than internal voids in both materials (Figs. 7(b) and (c)). This was due to the internal voids being partially filled with powder, which reduced the contrast on the x-ray film. Even though internal voids of this type would simulate an interconnected porosity structure in similar materials, it is difficult to establish reliability statistics for naturally occurring internal voids. Hence only the POD data for surface voids presented herein would apply to naturally occurring internal voids in green dry-pressed, injection-molded, or slip-cast materials. For sintered materials (Fig. 8) the detection sensitivity was about 1.5 percent of thickness for surface and internal voids in SiC. 1.5 percent of thickness for surface voids in SiaNa. and better than 1.5 percent of thickness for internal voids in Si_3N_4 at a POD of 0.9. This high sensitivity to internal voids in sintered Si $3N_A$ was due to void morphologies that enhanced the x-ray attenuation depending on the density, chemical composition, and thickness of the shell surrounding the voids. Thus the image contrast of some voids was improved. A dense shell formation could occur for materials where high-atomic-number additives are used to promote the sintering process. When local chemical or density variations are not present, as is the case in sintered SiC, the detection reliability for internal voids can be inferred from the POD curves for surface voids (Fig. 8(b)).

Scanning Laser Acoustic Microscopy

Figure 9 shows how specimen thickness and surface condition affected the POD (at 0.95 confidence level) of surface-connected voids in sintered Si_3N_4 (9). For as-fired 2-, 3-, and 4-mm-thick specimens (with surface roughness of 8 μ m, peak to valley, the 0.9 POD was not attained. After the same specimens were polished to a surface roughness of 2 μ m, 0.9 POD was achieved for all thicknesses. This acoustical opacity of the as-fired specimens was attributed to the surface roughness. However, near-surface pore distribution might have scattered the sound wave and resulted in poor void detectability. Figure 10 shows how void size, void depth below the laser-scanned surface, and matrix material affected the POD of internal voids in sintered specimens having diamond ground surfaces (7). The boundaries of the bar graphs indicate the smallest void sizes and the maximum depths at which 0.9 POD (0.95 confidence level) was achieved. The detection sensitivity was greater for Si_3N_A than it was for SiC. This can be attributed to the coarser grain structure and greater porosity of SiC. These microstructural differences might have increased the ultrasonic scatter and resulted in more ultrasonic attenuation in SiC.

NDE PROCESS CONTROL

Processing and sintering conditions were modified on the basis of feedback from radiography to obtain the uniform high-density MOR bars associated with improved strength and reduced strength scatter. Figure 11 shows how these modified procedures affected the flexural strength of NASA 6Y Si_3N_4 sintered at 2140 °C. The sensitivity of radiography to the within-bar porosity distribution in the baseline and improved materials is shown in Fig. 12. Efforts were made to eliminate the case-core structure (Figs. 6 and 12) and thereby strengthen the material and reduce its strength scatter.

The case-core structure was examined as a function of the powder fineness and the sintering variables grinding time, sinter cup height, nitrogen overpressure, and degree of BN setter contact. Radiography showed that increasing the powder fineness (by using 100- and 300-hr milling times) resulted in thinner and less distinct cases and thus improved both sinterability and uniformity. Minimizing the BN setter contact resulted in more uniform densification as a consequence of more uniform heating. But increasing nitrogen overpressure from 2.5 to 3.5 to 5.0 MPa had no effect on the case-core structure. When using 100-hr milling time, smaller BN disks, and a nitrogen overpressure of 5.0 MPa. increasing the sintering time from 1 to 2 hr and raising the sinter cup into a more uniform temperature zone were very effective in greatly reducing density gradients in sintered bars. The most uniform material, batch 31 (Fig. 12), was the result of the cumulative positive effects of increasing the powder grinding time from 24 to 300 hr, increasing the sintering time from 1 to 2 hr, minimizing the BN setter contact, adjusting the sinter cup height, and using powder wet-sieving procedures. Wet sieving was used to eliminate agglomerates and to reduce the size of impurity particles in green-pressed bars that vaporize during sintering and leave voids in the sintered bars.

Microstructures for the baseline batch and the optimized batch (31) were compared and related to position in the bar by reference to radiographs (Fig. 12). The top and bottom surfaces (W,L) planes of batch 31 had essentially pore-free zones of equal thickness. However, the (W,L) planes of the baseline batch contained zones of dissimilar thickness and appearance and had fine porosity. The internal porosity of batch 31 (3.24 g/cm³) was considerably less than that of the baseline batch (3.12 g/cm³).

The individual and combined effects of powder fineness and each of the sintering variables on the flexural strength of NASA 6Y at room temperature, 1200 °C, and 1370 °C are explained in detail in reference 3. Only the cumulative positive effects of all the variables on flexural strength are shown in Fig. 11. In processing from batch to batch (baseline to 28 to 29 to 31) the room-temperature strength continually increased, with an overall improvement of 56 percent and more than a threefold reduction in the standard deviation. Strength improved 38 and 21 percent at 1200 and 1370 °C, respectively. All successive improvements in the mechanical properties of sintered Si_3N_4 were guided by x-radiographic characterization. For the most improved material the previously dominant failure-causing voids were replaced by large columnar grains, which are less detrimental to strength properties.

CONCLUSIONS

The reliability of microfocus x-radiography for nondestructively evaluating sintered ceramics was affected by the seeded void morphologies, which enhanced the x-ray attenuation depending on the density, chemical composition, and thickness of the shell surrounding the voids. The reliability of scanning laser acoustic microscopy was affected by the specimen surface roughness and microstructural characteristics. Rough surfaces, large grains, and high porosity reduced the detection capability. The successful use of conventional x-radiography in guiding the fabrication process resulted in denser and more uniform Si_3N_4 over the baseline material. The improved material reached four-point average flexural strength/ standard deviation values of 857/36, 544/33, and 462/59 MPa at room temperature, 1200 °C, and 1370 °C, respectively. These strengths represented improvements of 56, 38, and 21 percent over the baseline properties at the three test temperatures. Further, previously dominant failure-causing voids were replaced by large grains, which are less detrimental to strength properties.

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Figure 1. - Fabrication of silicon nitride and silicon carbide test specimens with seeded internal voids.



Figure 2. - Schematic configuration of microfocus projection radiography, where D is thickness of defect, T is thickness of sample, μ_1 is attenuation coefficient of matrix, and μ_2 is attenuation coefficient of defect.



Figure 3. - Scanning laser acoustic microscopy (SLAM) of ceramic test bars containing seeded internal defects.



Figure 4. - Powder processing of Si_3N_4 - SiO_2 - Y_2O_3 composition NASA 6Y.



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Figure 5. - Characterization, testing, and evaluation of NASA 6Y Si_3N_4 .



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Figure 6. - Schematic configuration of conventional radiography employed for characterization of NASA 6Y Si_3N_4 test bars (L = 3.0 cm, W = 0.56 cm, T = 0.28 cm).

















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Figure 11. - Effect of modified processing/sintering procedures on flexural strength of NASA 6Y Si_3N_4 sintered at 2140 $^{\circ}C$.

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1. Report No.	2. Government Accessio	n No.	3. Recipient's Catalog No	0.
NASA TM-88870				
4. Title and Subtitle			5. Report Date	
NDE Reliability and Proces	s Control for Structural			
Ceramics			6. Performing Organizatio	on Code
			533-05-11	
7. Author(s)	· · · · · · · · · · · · · · · · · · ·		8. Performing Organizatio	on Report No.
George Y. Baaklini			F-3276	
			L-3270	
		1	0. Work Unit No.	
9 Performing Organization Name and Address				
National Apropautics and Space Administration			1. Contract or Grant No.	
Lewis Research Center				
Cleveland, Ohio 44135		1	3. Type of Report and Pe	riod Covered
12 Sponsoring Agency Name and Address	- · · · · · ·		Tochnical Mor	norandum
National Aeronautics and Space Administration			4. Sponsoring Agency Co	ode
Washington, D.C. 20340				
15. Supplementary Notes				
Prepared for the 32nd Inte sponsored by the American May 31 - June 4, 1987.	ernational Gas Tu Society of Mecha	irbine Conference inical Engineer:	ce and Exhibit s, Anaheim, Ca	ion, lifornia,
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17. Key Words (Suggested by Author(s))		18. Distribution Statemer	it un]imi+ad	
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Unclassified	Unclassified			
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