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Impact Sintering Temperatures on Mechanical Properties of Ba1-xSrxTiO3

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Abstract: This study investigates the influence of sintering temperatures on the mechanical properties of Ba1-xSrxTiO3 ceramics. Ba1-xSrxTiO3(x=0.2) ceramics are synthesized via Sol-Gel and sintered at varying temperatures. We use the appropriate testing methods to describe mechanical qualities like hardness, fracture toughness, and elastic modulus. The results show that the mechanical behavior changes a lot when the sintering temperature changes. This shows how important processing conditions are for customizing the mechanical performance of Ba1-xSrxTiO3 ceramics that could be used in advanced electronic and structural materials. XRD pattern exhibited a tetragonal phase, and the crystal size increased with increasing sintering temperature. The BST sample's surface morphology appears uniform and homogeneous, with moderate sintering temperatures. High Sintering Temperature increases mechanical strength due to improved densification and reduced porosity, producing higher hardness because of increased density and larger, well-formed grains and improved fracture toughness as the material becomes denser and grain boundaries are better formed, enhancing crack resistance. Also, (c/a ratio) was found to decrease with an increase in sintering temperatures.

Keywords: Ba1-xSrxTiO3, Mechanical Properties, Sintering Temperatures, hardness, toughness.

1 Introduction

Ba1-xSrxTiO3 ceramics have received considerable interest due to their varied uses in electronic devices, capacitors, and structural materials because of their remarkable dielectric and ferroelectric properties [1][2][3]. А comprehensive comprehension of the impact of manufacturing parameters, such as the sintering temperature, on the mechanical behavior of these ceramics is essential for their functionality in a variety of applications [4][5][6]. The sintering temperature is an important factor in ceramic processing because it affects the growth of grains, the development of microstructures, and the formation of defects [7][6]. These changes have a direct effect on mechanical traits like hardness, fracture toughness, and elastic modulus [8][9].

Ba1-xSrxTiO3 (BST) compounds, which are part of the perovskite family, have a diverse range of uses in different disciplines because of their outstanding dielectric and ferroelectric properties [10]. The composition and processing conditions, particularly the sintering temperature, significantly influence the tuning of these properties [11][12]. Among the primary elements determining these qualities, sintering temperature stands out as a critical parameter controlling the microstructure and, as a result, the material's mechanical behavior [13][14].

This study examines the influence of sintering temperatures

ranging from 1000°C to 1400°C on the mechanical characteristics of BST ceramics. The primary objective is to analyze how different sintering temperatures affect the mechanical properties of BST ceramics. Understanding these relationships is essential for optimizing the material's performance in practical applications. This research aims to provide insights into how variations in sintering temperature affect the mechanical characteristics of BST ceramics, thereby contributing to the broader understanding and enhancement of their functional properties [15]. Understanding how variations in sintering temperature influence the mechanical performance of BST ceramics is essential for optimizing their structural integrity and reliability in practical applications [16][17].

This work aims to understand thoroughly the effects of sintering conditions and mechanical properties (such as elastic modulus, attenuation, and velocity of ultrasonic waves) of Ba(1-x)Sr(x)TiO3ceramics sintered at 1000 to 1400°C for 3h. X-ray diffraction and ultrasonic techniques (at a frequency of 2 MHz) were used to characterize the structure and phase transitions of the prepared BST ceramics.

2. Experimental procedures

 $Ba_{0.8}Sr_{0.2}TiO_3$ ceramics were prepared by the new method called the sol-gel method. The Barium acetate, strontium acetate, and Ti isoproxide (IV) were mixed in the appropriate molar ratios (1:1). First, the Ba and Sr acetate solution was mixed and stirred at 60°C. Second, the Ti isopropoxde with 2-



methanol was mixed and added to the first solution. The mixture solution was calcined at 700°C for 3 h in open air. The produced fine powders were pressed into disc-shaped pellets (D:10 mm and Th: 2 mm) at an isostatic pressure of 12 tons. The pelletized samples were sintered at different sintering temperatures.

Characterization:

"X-ray powder diffraction (XRD) was conducted using Cu Ka (k = 1.54056 A) radiation at 40 kV and 40 mA. The microstructure was visualized using scanning electron microscopy (FESEM, JSM EMP-800, JEOL, Japan) [12]. The ultrasonic wave velocities v(in ms-1) propagated in the samples and attenuation $\alpha(\text{dB/cm})$ were measured at room temperature, using pulse-echo technique MATEC Model MBS8000 DSP (ultrasonic digital signal processing) system with 2 MHz resonating.

3. Results and Discussion

Finding the sweet spot for BaSrTiO₃ sintering temperature involves juggling densification, grain development, and optimizing mechanical properties. When the temperature is too high, it can cause grain growth and compromise mechanical qualities; when the temperature is too low, densification is inadequate. Ultimately, the sintering temperature is crucial in establishing the mechanical characteristics of BaSrTiO₃ as shown in Fig. 1. By manipulating the sintering temperature, it is possible to customize the density, grain size, hardness, fracture toughness, dielectric characteristics, and mechanical strength of the material to fulfill specific application demands. The densification factor exhibited a positive correlation with temperature, increasing until it reached its maximum concentration. A positive densification factor indicates that ceramics shrink during the sintering process.



Fig. 1: The sintering temperature vs density of Ba_{0.8}Sr_{0.2}TiO₃

The density of BaSrTiO₃ (barium strontium titanate) is strongly influenced by sintering temperature. Generally, as the sintering temperature increases, the density of the material also increases. This relationship is due to the processes of densification and grain growth that occur during sintering.

$$DF = \frac{\rho_{m-\rho_g}}{\rho_{t-\rho_g}} \tag{1}$$

"Where ρ_g represented the density of green pellets, ρ_t theoretical density and ρ_m is the measured density"

The shrinking measurement checked the diameters of both pre-sintering samples

$$D_{shrinking} = \frac{d_o - d_s}{d_0} \tag{2}$$

"Where d_0 and d_s are the diameter of the green and sintered pellets, respectively"

4. Microstructural Evolution

4.1 Morphology and Surface Characteristics

Fig. 2 displays the FESEM picture of the BaTiO3 ceramic system at various sintering temperatures. The BaTiO3 ceramic, heated to 1150 °C, exhibits an open structure due to the loose packing of particles. This makes the ceramic less dense. At a temperature of 1200 °C, the particles start to form bonds, but the density of pores remains high. At a temperature of 1250 °C, the particles coalesced, and no empty spaces were detected. Nevertheless, the surface microstructure and the particle boundaries remain ambiguous. At a sintering temperature of 1300 °C, the pores are nearly eliminated, resulting in a strongly bonded grain boundary and the formation of a high-density ceramic.

As the temperature continues to rise until it reaches 1400 °C, the grain size increases, and holes appear around the edge and inside the seed. The morphology and surface features of the material significantly determine the mechanical properties of Ba0.8Sr0.2TiO3 (BST) ceramics. Differences in the temperature during the sintering process control these qualities. Field emission scanning electron microscopy (FESEM) pictures showed that the sintering temperature had a big effect on the size and spread of the grains. Lower sintering temperatures, between 1000°C and 1200°C, resulted in smaller and more evenly distributed grains in BST ceramics (Fig. 2). As the temperature of the sintering rose to 1400 °C, grain growth was seen.



Fig. 2: Images of FESEM for surface morphology Ba0.8Sr0.2TiO2

4.2 Phase Composition

X-ray diffraction (XRD) analysis validated the phase purity and crystallinity of BST ceramics across various sintering temperatures. There were unique peaks that corresponded to the creation of specific phases, such as tetragonal and cubic phases, at different temperatures (Fig. 3) Illustration. Ceramics with Ba1-xSrxTiO3 (x=0.2) X-ray diffraction patterns Fig. 3 depicts the indexing of all the peaks. These indices relate to the reflections of the various polycrystalline orientations, with the 110 indices being the high-intensity main peak. It shows that as the sintering temperatures increased from 1000 to 1400°C, the peak positions at 20° angles 31.83251°, 31.88799°, and 32.06737° were confirmed by a closer examination of the 110 peaks for all samples. Scherrer's equation is used to estimate the crystallite size D (hkl) by referring to the broadening of the x-ray line's half-width. The table below lists the evolution of crystal structure, lattice parameters, and crystallite size obtained from XRD data.Table 1 shows that increasing the sintering temperatures leads to an increase in both lattice parameters and crystallite size. The crystal structure showed tetragonal phases for all tested samples and the tetragonality ratio (c/a) decreased as the sintering temperature increased as reported [12].



Fig. 3: XRD pattern for Ba0.8Sr0.2TiO3 with different sintering temperatures

Table 1: The sintering temperatures vs lattice parameters, crystallite size and Tetragonality

| Sintering Temp. C | a (Å) | c (ϟ) | Phase | The Tetragonality ratio (c/a) | Volume(Å ³) | D (nm) |
|----------------------|---------|--------|------------|-------------------------------------|-------------------------|--------|
| 1000 | 3.9771 | 3.9885 | Tetragonal | 1.0038 | 64.08 | 22.78 |
| 1200 | 3.97815 | 3.9922 | Tetragonal | 1.003531 | 63.89 | 32.11 |
| 1400 | 3.97872 | 3.9941 | Tetragonal | 1.00286 | 63.82 | 36.60 |

5. Mechanical Properties

Indentation tests revealed variations in the hardness and fracture toughness of BST ceramics as a function of sintering temperature. Generally, higher sintering temperatures led to increased hardness due to improved densification and grain growth, whereas fracture toughness showed a more complex relationship influenced by microstructural features (Fig. 5). The ultrasonic wave velocities v were calculated by taking the elapsed time between the initiation and the receipt of the pulse appearing on the screen. It can be calculated using the following equation:

$$\nu = \frac{2x}{\Delta t} \tag{3}$$

"Where x is the sample thickness and Δt is the time interval. The measurements were repeated three times to check the reproducibility of the data. The estimated accuracy of the velocity measurement is about 0.04%. The values of elastic moduli; namely longitudinal modulus (L), shear modulus (G), Young's modulus (E), bulk modulus (B), and Poisson's ratio (σ) were obtained from the longitudinal (VL)2 and shear (VS)2 ultrasonic velocities and calculated from the following equations:"[7]

$$L = \rho v_L^2 \tag{4}$$

$$G = \rho v_S^2 \tag{5}$$

$$E = \frac{G(3L - 4G)}{(L - G)}$$
(6)

Table 2: The sintering temperatures vs the mechanical properties (L, G, and E), particle size

| Sintering Temp. °C | Density (Kg/m ³) | Long. Velocity V _L (m/s) | Elastic Modulus L(GPa) | longitudinal modulus (L) | shear modulus (G) | Particles size(nm) |
|-----------------------|---------------------------------|---|------------------------------|-----------------------------|----------------------|-----------------------|
| 1000 | 5.83 | 634.56 | 4.17 | 7.56 | 2.22 | 78.71 |
| 1200 | 5.89 | 1178.89 | 11.45 | 14.78 | 4.67 | 132.11 |
| 1400 | 5.93 | 1278.57 | 16.34 | 18.98 | 6.98 | 141.60 |



Fig. 4: Elastic Moduli E, G, L vs sintering temperatures Effects of Sintering Temperature An increase in the sintering temperature can cause changes in the material's



microstructure, including changes in grain size and porosity. In general, greater sintering temperatures tend to decrease porosity and increase grain size, which ultimately leads to an increase in the longitudinal modulus. Increased density and uniformity in a structure typically lead to increased stiffness in the direction of the applied force. Longitudinal Modulus (L): Increases with higher sintering temperature due to reduced porosity and increased grain size. Shear Modulus (G): Increases with higher sintering temperature as the material becomes denser and more homogeneous. Young's Modulus (E): Typically increases with higher sintering temperature due to improved structural integrity and stiffness. Bulk Modulus (B): Increases with higher sintering temperature as the material becomes more dense and less compressible.

6. Conclusion

The observed changes in morphology and surface characteristics underscore the intricate relationship between processing parameters (sintering temperature) and the microstructural evolution of BST ceramics. Higher temperatures significantly impact mechanical properties such as hardness, fracture toughness, and flexural strength. Understanding these relationships is crucial for tailoring BST ceramics with optimized mechanical performance for diverse applications in electronics, telecommunications, and beyond. Overall, the results highlight the importance of precise control over sintering parameters to achieve desired microstructural features and mechanical properties in BaixSrxTiO3 ceramics, paving the way for further advancements in materials design and application-specific optimizations. This structured approach integrates both results and discussion, emphasizing the interplay between sintering temperature and mechanical properties of BST ceramics.

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Conflict of Interest

The project has no financing, and the authors have declared no competing financial interests.

Availability of data materials

We won't disclose the data because it helps with the study of choosing the right forms for the right applications.

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