Enhancement of the Optical and Dielectric Properties at Low Frequency of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, (0 ≤ x ≤ 0.06) Structure Ceramics

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Abstract: Low loss Ruddlesden–Popper (RP) series, i.e., (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0.0 ≤ x ≤ 0.06, has been synthesized by a mixed oxide route. In this work, the substitution of Ca$^{2+}$ cation in Sr$_5$Ti$_4$O$_{13}$ sintered ceramics was chosen to enhance the structural, optical, and dielectric properties of the product. It was found that the Ca$^{2+}$ content has significant effects on enhancing the dielectric properties as compared to Mn and glass additions. It was observed that the relative density, band gap energy, and dielectric loss (tangent loss) increase while relative permittivity decreases along with Ca$^{2+}$ content. High relative density (96.7%), low porosity, and high band gap energy (2.241 eV) values were obtained in (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0.0 ≤ x ≤ 0.06 sintered ceramics. These results will play a key role in the application of dielectric resonators.

Keywords: (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$ ceramics; XRD; FT-IR optical; electric properties

1. Introduction

Recently, the Sr based Ti family has come to play a key role in the development of wireless communication technologies [1]. The low loss dielectric ceramics with good dielectric properties and temperature stability are widely used in the applications of dielectric resonator antennas [2–5]. To overcome the miniaturized devices requirements, the ceramic dielectrics must have maximum values of relative permittivity ($\varepsilon_r$) and low dielectric losses [6–9]. Besides the poor sintering temperature, the barium magnesium tanatate material have optimum dielectric properties, which is used further in the applications of dielectric resonator devices [10–12]. On the other side, compound, i.e., CaTiO$_3$-MgTiO$_3$, ceramics posses good dielectric properties, i.e., ($\varepsilon_r = 21.2$, $Q \times f = 56,200$ GHz, $\tau_f = \pm 0$ ppm/$^{°}C$). Lately, the RP series ceramics have been the subject of numerous scientific studies, with results showing their interesting dielectric properties [13–16]. $A_{n+1}B_nO_{3n+1}$ is the general formula of RP-series, and the phase analysis consists of n—blocks of octahedra (BO6) corner-sharing which construct a layered perovskite like structure. In previous research, it has been reported that the compound, i.e., SrLn$_2$Al$_2$O$_7$ and MLnAlO$_4$ (Ln = Nd, Ce, Pr, Ba)
La, & Sm, M = Ca, Sr), ceramics along with n = 1 and n = 2 show good dielectric properties. The RP series posses bulk structure and optimum dielectric properties, i.e., $\varepsilon_r = 16.0$ to 19.0, $Q \times f = 54,600–69,500$ GHz and $\tau_f = -1–32$ ppm/$^\circ$C for n = 1; $\varepsilon_r = 17.9$ to 21.6, $Q \times f = 64,680–71,680$ GHz & $\tau_f = +4–22.1$ ppm/$^\circ$C for n = 2, have been reported [13–17]. Fan Yi et al. reported the modification in phase, microstructure, and dielectric properties of RP-series [14,15]. Actually, this impression has been realized in $M^{2+}/Ti^{4+}$ cation substitution in MLnAlO$_4$ calcined ceramics, which further modified the phase, microstructure, optical, and dielectric properties, i.e., ($\varepsilon_r = 18.2$ to 21.4, $Q \times f = 75,000$ GHz to 96,500 GHz and $\tau_f \sim \pm 0$ ppm/$^\circ$C) [18–20]. On the other hand, the dielectric properties, especially quality factor value, are not too good. In RP-series, at n = 2, the ceramic compound, i.e., SrLn$_2$Al$_2$O$_7$ generally has low dielectric losses and good relative permittivity values as compared to MLnAlO$_4$ ceramics. Moreover, many researchers have reported the optimum dielectric properties among the RP series [13–21]. The RP-series, such as Sr$_2$La(A$_3$ + B$_4$)O$_7$ ($A = Fe, Cr, B = Mn, Ti$), has been investigated already in a literature review, and a new RP compound, (Sr$_2$LaAlTiO$_7$), has been synthesized and analyzed as a new low loss ceramic material [22–26].

In the present work, the good results on structural, optical, and dielectric properties (at low frequency) of Sr$_3$Ti$_4$O$_{13}$ based structure ceramics will be studied. These results will be modified by making some doping elements at A-site cation in the base product.

2. Experimental Procedure

The solid solution of (Sr$_{1-x}$Ca$_x$)$_3$Ti$_4$O$_{13}$, 0 $\leq x \leq$ 0.06 ceramic was processed by using high grade pure carbonates and oxide powders, i.e., SrCO$_3$ (99.95%), CaCO$_3$ (99.9%), and TiO$_2$ (99.5%) as raw materials. The resultant stoichiometric ratio of the raw materials were mixed properly and then milled using horizontal ball milling with zirconia media in distilled water for 24 h and then calcined for 3 h in air at 980 °C. After re-milling, the calcined powder was mixed along with polyvinyl alcohol (PVA) solution at 4 wt.% and then made into green pellets of 2–3 mm thickness and 10–12 mm diameter. Then, the green pellets were kept in high energy furnace at 1200 °C sintering temperature for 3 h in air to dense the pellets. After sintering, the pellets were cooled to 600 °C at the rate of 10 °C/min and then cooled to room temperature inside the furnace further. The bulk density was calculated by using the Archimedes principle method for all the pellets. The phase analysis was identified by using X-ray diffraction (XRD, RIGAKU D/max 2550/PC, Rigaku Co-Tokyo Japan) with CuK$_{α}$ radiation. The surface morphology of the thermally etched and gold coated samples was studied using scanning electron microscopy (SEM, S3400; Hitachi, Tokyo, Japan). The relative permittivity ($\varepsilon_r$) and tangent loss were measured by the parallel plate capacitor method using vector-network Analyzer (E8363B, Agilent Technologies Inc., Santa Clara, CA, USA) [27]. At least four samples have been analyzed to ensure the accuracy of data. The reciprocal of Q-factor is the tangent loss (tan$\delta$) [28].

3. Results and Discussion

3.1. Phase Analysis

Figure 1 shows the XRD patterns of RP series of (Sr$_{n+1}$Ti$_n$O$_{3n+1}$) sintered ceramic for n = 4. The patterns revealed the tetragonal structure of RP series along with space-group (I4/mmm) matched to PDF card number 89-1383. The structure of the phase (at n = 4) was attained by put in a rock-salt type Sr-O layers, the strontium based titanates along with direction [001], resulting consecutive perovskite pieces due to shifting by direction $\frac{1}{2}$ [111], w.r.t the unit cell of RP series. The known RP structure has closely alike lattice parameters i.e., (a = b = 0.385 to 0.389 nm) but c = 2.812 nm for n = 4 [27–32]. The variation of lattice parameters and volume of the synthesized samples with Ca$^{2+}$ contents as shown in Table 1. The shifting of peaks to lowest Bragg’s angles were due to the difference of ionic radii of Sr$^{2+}$ and Ca$^{2+}$ cations as shown in Figure 1b. No secondary phase has been observed and revealed the single phase of Sr$_{n+1}$Ti$_n$O$_{3n+1}$ (n = 4) sintered ceramic. Figure 2 shows the variation of relative density with Ca$^{2+}$ contents of Sr$_{n+1}$Ti$_n$O$_{3n+1}$ (n = 4) sintered
ceramics. It has been noted that the relative density increases with the Ca$^{2+}$ content, which further modified the optical and dielectric properties. The highest values of relative density is (96.7%) of Sr$_{n+1}$Ti$_n$O$_{3n+1}$ ($n = 4$) sintered ceramics was observed at $x = 0.06$ content.

Figure 1. (a) XRD pattern of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0 $\leq x \leq 0.06$ sintered ceramics (b) zoomed view of peak (1 0 4) shifted toward the lowest angle.

Figure 2. The relative density of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0 $\leq x \leq 0.06$ ceramics.
Table 1. Volume and Lattice parameters of Sr$_{n+1}$Ti$_n$O$_{3n+1}$ (n = 4) sintered ceramics.

| Contents | a = b (Å) | c (Å)   | c/a    | Error   | Volume of Unit Cell (Å$^3$) |
|----------|-----------|---------|--------|---------|-----------------------------|
| 0.00     | 3.8512    | 28.1253 | 7.3411 | ±0.8631 | 417.1471                    |
| 0.02     | 3.8518    | 28.3951 | 7.3719 | ±0.8643 | 421.2800                    |
| 0.04     | 3.8523    | 28.5505 | 7.4113 | ±0.8651 | 423.6955                    |
| 0.06     | 3.8615    | 28.8975 | 7.4837 | ±0.8664 | 430.8958                    |

3.2. Surface Morphology

Figure 3 shows the SEM images of the gold coated samples of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0.00 ≤ x ≤ 0.06 sintered ceramics. The variation of relative densities and grain size of all the samples has been investigated. The SEM micrographs of (Sr$_5$Ti$_4$O$_{13}$) green pellets with doping of Mn or glasses at different sintering temperature were studied by many scientific researchers [26]. It has been reported that the base product have small crystallite size and less porosity, which may be affected by the surface strain. However, new grains and porosity were produced by adding some dopant elements in (Sr$_5$Ti$_4$O$_{13}$) sintered ceramic [27–30]. Ca$^{2+}$ concentration has been observed to increase the porosity and grain size of all samples in (Sr$_5$Ti$_4$O$_{13}$) sintered ceramic. These factors will affect the structure, optical, and dielectric properties of the base product. In order to improve these properties, numerous studies have examined the synthesis settings used to create various dopants in the base product [31,32].

Figure 3. SEM images of polished & thermally etched samples of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0 ≤ x ≤ 0.06 sintered ceramics (a) x = 0.00, (b) x = 0.02, (c) x = 0.04 and (d) x = 0.06.
3.3. Fourier Transform Infra-Red (FTIR) Spectroscopy

Figure 4 shows the FTIR spectra of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, $0 \leq x \leq 0.06$ sintered ceramic. FTIR spectrometer plays a key role to characterize the vibrational stretching and un-stretching mode of the base sample synthesized by chemical reaction route [33,34]. The vibrational stretching mode (O-H) was observed with variable wave number ($K = 2\pi/\lambda$) i.e., 900.0 cm$^{-1}$, and 3200.0 cm$^{-1}$. This mode of vibration is produced by the absorption of vapors during synthesis process. Only asymmetric mode at wave number (3700.0 per cm) was recorded in the base product which shown carboxylates family [35]. In this characterization the normal stretching mode was observed at wave number (500.0 per cm).

![Figure 4. FTIR spectra of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, $0 \leq x \leq 0.06$ sintered ceramics.](image)

3.4. UV Spectroscopy

Figure 5 shows the UV-spectra of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, $0.00 \leq x \leq 0.06$ sintered ceramics. Many of the researchers reported that the Sr$_5$Ti$_4$O$_{13}$ base sample was found to be transparent for white light [36]. It is very important to note that the compound, i.e., Sr$_5$Ti$_4$O$_{13}$, is translucent for visible light. The band structure and electronic transition were characterized using photon energy [37]. The electron needs to execute the inner shell transition in order to obtain the optical bandgap energy. This optical bandgap energy strongly depends upon the coefficient of absorption ($\alpha$), which was calculated using Equation (1) [25].

$$\alpha = A \frac{(hv - E_g)^{1/2}}{hv}$$  \hspace{1cm} (1)

where $E_g$ = bandgap energy, $A$ = constant of proportionality, and $hv$ = photon energy.

The coefficient of absorption will be defined how distant light of specific wavelength can be penetrated into material before being absorbed. When light absorbed poorly by material have low coefficient of absorption looks to be thin at specific wavelength. The unit
of absorption coefficient is (cm$^{-1}$). The band gap energy of all the samples was calculated using the Tauc plots method. It was reported that the values of band gap energy increase with Ca$^{2+}$ content.

![Figure 5](image_url)

Figure 5. Band gap energies of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0.0 $\leq$ x $\leq$ 0.06 ceramics i.e., (a) 1.684 eV (b) 2.093 eV (c) 2.216 eV & (d) 2.241 eV.

3.5. Photoluminescence (PL) Spectroscopy

Figure 6 shows the photoluminescence (PL) spectroscopy of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0.00 $\leq$ x $\leq$ 0.06 sintered ceramics. The emission line spectrum will be produced by the recombination of holes and electron charge carriers. Using the equation (E = hc/$\lambda$), where E = optical excitation energy, h = Plank’s constant (~6.63 $\times$ 10$^{-34}$ J$s$) c = speed of light (3 $\times$ 10$^8$ m/s) and $\lambda$ is the emission wavelength, we can find the value of excitation energy of all the samples.

Emission at photoluminescence peak of the samples has been noted at the range of ~400–550 nm. Multiple photonic processes such as PL have certain common uses, and PL is a multiple photonic process that has some typical applications, i.e., (i) determination of band gap energy, (ii) material quality, as well as (iii) molecular structure and crystallinity, reported by many researchers [38,39]. It has been observed that the broader emission spectra were located near to ~2.48 eV (excitation energy) and wavelength (~500 nm) which is larger than bandgap energy of all the samples may be occurs due to the presence of impurities. In the photoluminescence spectrum, the cyan color may occur due to the oxygen vacancy [40].
3.6. Low Frequency Dielectric Properties

The low frequency dielectric properties of all the synthesized samples sintered at 1200 °C for 3 h in air were better due to their high relative densities. The variation of relative permittivity ($\varepsilon_r$) and tangent loss (tan$\delta$) with varying temperature was measured at 100 Hz–1 MHz for (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0.0 ≤ x ≤ 0.06 sintered ceramics using the vector network analyzer, as shown in Figure 7. Strong irregularity in relative permittivity ($\varepsilon_r$) and tangent loss (tan$\delta$) were observed for the contents (at x = 0.0 and 0.02), which shows the transition of ferroelectric to Paraelectric phases. The same behavior was recorded in the values of ‘$\varepsilon_r$’ and ‘tan$\delta$’ for Ba$_{5-x}$Sr$_x$DyTi$_3$V$_7$O$_{30}$ (0 ≤ x ≤ 5) sintered ceramics at temperatures of 430 °C, 350 °C, 325 °C, 85 °C, and 42 °C, respectively [41,42]. The lowest value of $\varepsilon_r$ (~1400) was observed for (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, (composition with x = 0.02) at 100 Hz frequency, and found to decrease with increasing operating frequency, which may be due to the interfacial polarizations. Moreover, the value of $\varepsilon_r$ decreased with increasing Ca$^{2+}$ contents, which is due to the difference of ionic polarizabilities of Ca$^{2+}$ (3.16 Å$^3$) and Sr$^{2+}$ (4.24 Å$^3$) [43–45]. It has been revealed that the value of tangent loss increases with temperature due to the process of conductivity and different types of polarizations at low frequency [21]. The lower value of the tangent loss was reported at 1 MHz operating frequency for the base sample. The variations in both the quantities may be due to the difference in the values of dielectric polarizabilities [46]. Generally, tan$\delta$ decreases when high cation ions are replaced by smaller cation ions [47].

The complex impedance spectroscopy mechanism is generally used to investigate the structural properties and bonding of the various types of materials, comprising the ferroelectric, ionic insulator, and linked ceramics under different experimental conditions [36]. The variation in real impedance $Z'$ and imaginary impedance $Z''$ of (Sr$_{1-x}$Ca$_x$)$_5$Ti$_4$O$_{13}$, 0.0 ≤ x ≤ 0.06 sintered ceramics is shown in Figure 8. Initially, it was revealed that the magnitude of $Z''$ increases with $Z'$ and then decreases due to the release of space charge polarization [37]. It was observed that the magnitude of $Z''$ decreases by increasing the $Z'$ and Ca$^{2+}$ contents.
Figure 7. Cont.
Figure 7. Variation of $\varepsilon_r$ and tan$\delta$ with temperature for the $(\text{Sr}_{1-x}\text{Ca}_x)_{5}\text{Ti}_4\text{O}_{13}$, $0.0 \leq x \leq 0.06$ sintered ceramics i.e., (a) $x = 0.00$, (b) $x = 0.02$, (c) $x = 0.04$ & (d) $x = 0.06$.

Figure 8. Cole-Cole Plots of $(\text{Sr}_{1-x}\text{Ca}_x)_{5}\text{Ti}_4\text{O}_{13}$, $0 \leq x \leq 0.06$ sintered ceramics.

4. Conclusions

The solid solutions of $(\text{Sr}_{1-x}\text{Ca}_x)_{5}\text{Ti}_4\text{O}_{13}$, $0.0 \leq x \leq 0.06$ sintered ceramics was synthesized by conventional solid state method. The structural, microstructural, optical, and dielectric properties of all the samples have been investigated. The XRD patterns revealed the tetragonal phase with space group (I4/mmm). The SEM image revealed that the grain size and porosity increase with increasing Ca$^{2+}$ contents, which was due to the difference of ionic radii. The results of UV spectroscopy declared that the bandgap energy increases from 1.68 eV to 2.24 eV along with increasing Ca$^{2+}$ concentrations. The good values of dielectric properties (i.e., $\varepsilon_r \sim 250$, and tan$\delta$ = near to zero) in the frequency range from 100 Hz to 1 MHz was observed. It has been observed that the magnitude of $Z''$ in-
creases with $Z'$ and Ca$^{2+}$ contents. The overall findings are suitable for the application of dielectric devices.

**Author Contributions:** This work was carried out in collaboration among all authors. S.J.A., A.H.A., V.T. and A.A. (Ali Algahtani) Writing, review, editing and Experimentation analysis. A.Z. and A.A. (Asad Ali) did the final review, corrections, and editing. A.A. (Aiyeshah Alhodaib), I.B. and M.A. Formal Analysis and did the final review, corrections, and editing. M.K. and Abdullah Data curation and Resources. Resources V.T. and A.Z.; Visualization A.A. (Ali Algahtani); and Funding acquisition, V.T. and A.H.A. All authors have read and agreed to the published version of the manuscript.

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