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An International Journal of the Materials Science and Technology Society of Nigeria

# Nigerian Journal of Materials Science and Engineering (NJMSE)

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## Impedance and Modulus Spectroscopy of Nanocrystallite Barium Titanate Ceramic Using Mechanochemical Method.

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#### Abstract

Nanocrystalline BaTiO<sub>3</sub> (BT) powder was synthesized by a combination of the solid-state and mechanochemical method. X-ray diffraction, field emission scanning electron microscopy, and impedance spectroscopy utilised appropriately to characterize the BT sample (ceramic). The X-ray diffraction confirmed a single-phase perovskite compound of cubic symmetry with space group *Pm-3m*. The crystallite size and crystal cell volumes were found to be 25.7 nm and 64.250 Å<sup>3</sup> respectively. The average grain size estimated from FE-SEM was found to be 144.5 nm by using intercept technique. Electrical parameters like impedance, modulus, and electrical conductivity of the ceramic were obtained from AC complex impedance spectroscopy technique in the frequency and temperature range of 40Hz – 1MHz and 30 to 150°C, respectively. Both impedance and modulus plots showed the negative temperature coefficient of resistance (NTCR) character in the sample at 70, 110, 130, and 150°C with are similar to a semiconductor. This can be used for the fabrication of highly sensitive thermistors. The Cole-Cole (Nyquist) plot represents the grain and grain boundary conduction which indicates the ideal non-Debye type dielectric relaxation. The modulus analysis suggested the temperature-dependent relaxation process in the BT ceramic. A positive temperature coefficient of resistance (PTCR) character was observed at 90, 50, and 30°C. The activation energy values are found to be 1.12 eV and 1.07 eV for Z'', and M'' while it's 0.46 eV and 0.12 eV for  $\tau_g$  and  $\tau_{gb}$  respectively. A possible suggestion that the carrier transport is due to hopping conduction. The AC and DC conductivity spectra were found to rise with increasing temperature and frequency obey the Jonscher's power law. The conduction process was observed to be thermally activated and followed by Arrhenius law.

Keyword: Keyword: Barium titanate, Nanocrystalline, Impedance spectroscopy, AC conductivity

## INTRODUCTION

Barium titanate (BT) compounds are a standout amongst other perovskite ferroelectric compounds. Broadly, in all ferroelectrics, the associated physical properties and nature of conductivity are recognized by the study of electrical conductivity. In some previous studies (Jaffe 1971; Haertling 1999; Wei and Yao 2007; Hoshina *et al.* 2008), it was found that the interior defects like A-site and oxygen vacancies typically have excellent effect on ferroelectric fatigue or ionic conductivity of the material. In view of the fact that the solid defects play vital role in the thorough applications, it is pertinent for us to acquire a primal intellect of their conductive mechanism.

One of the tools of interest is Complex impedance spectroscopy (CIS) (Macdonald 1987). It is a nondestructive method accustomed to break the grain boundary and grain-electrode effects, which conventionally remain the places of a trap for oxygen vacancies and other flaws. It is also utilized in fixing space charge polarization and its relaxation mechanism, by properly assigning different practical values of resistance and capacitance to the electro-active regions (grain and grain boundary). It typically comes to moves into the shape of complex planes as successive semicircles as observed by (Bidault *et al.* 1994; Li *et al.* 2006; Khan *et al.* 2013). The semicircles symbolised the

electric phenomena involved within the polycrystalline material allegedly owing to grain, grain boundaries, electrical interface effect, and correlate between the dielectric and electrical characteristics. The complex electric modulus formalism has been utilised within the analysis of the electrical properties because it typically offers the first direct response of the majority of the used sample, abolishing the results owing to the electrode and electrical contacts. To be specific, it's peculiarly fitted to carefully extract data owing to the electrodes and ascertaining conductivity relaxation times (Hannachi *et al.*2010).

Both complex impedance and modulus spectroscopy are gainfully employed during this scholarly work because Z'' vs Z' and M'' vs M' plots highlight develop the phenomenon of most massive resistance and littlest capacitance, respectively (Jacob *et al.* 2015). In this paper, we reported a comprehensive study of the conduction mechanism in nanocrystalline Barium titanate ceramic.

## **EXPERIMENTAL PROCEDURES**

BaTiO<sub>3</sub> nanocrystalline ceramics were synthesised by a combination of solid-state reaction and high energy ball milling technique (HBM). The respective masses of the starting powders BaCO<sub>3</sub> and TiO<sub>2</sub> were weighed according to nominal composition using a digital



analytical balance with an accuracy of  $\pm 0.0001$  g. The overall balance equation is given in Equation 1.

$$BaCO_3 + TiO_2 \rightarrow BaTiO_3 + CO_2$$
(1)

A Stoichiometric amount of the weighed oxides (BaCO<sub>3</sub> and  $TiO_2$ ) were mixed in an agate mortar and then ground using a pestle to obtain a first step homogeneous small particle size. The mixture plus zirconia balls were placed in a plastic jar and ball-milled for 12 h at 120 rpm using Isopropyl alcohol as a mixing agent. The weight ratio of the zirconia balls to the oxide powders was 10:1 for better milling which produced homogeneous powder of fine particles. After the mixing process, the mixture was put in a petri dish and placed in an oven and then dried at 90°C for 12 h. The dried powder was grounded again using pestle and mortar to break the caking and produced fine powder for calcination. The mixed powders of BaCO<sub>3</sub> and TiO<sub>2</sub> were placed in a high-temperature ceramic boat (crucible). The loaded ceramic boat was then placed in Carbolite furnace (Vecstar Ltd, U. K) and calcined at 1050°C in the air. The calcined sample and isopropyl alcohol as wetting media were put in SPEX 8000 mixer/miller machine and grind for 7 h. The milling was stopped for 15 min after every 60 minutes of milling to cool down the system. After the grinding process, the mixture (slurry) was put in a 1330 GX multipurpose oven (Sheldon manufacturing, Inc. U. S. A), and dried at 90°C for 12 h. The dried powder of the sample was compacted at 5 tons into pellet of 15 mm diameter using 5 wt% polyvinyl alcohol (PVC) as a binder. The pellets of BT sample were placed in alumina crucibles and sintered in a programmable Carbolite furnace (Vecstar Ltd, U.K) at temperatures of 1190°C for 2 h.

The phase identification of sintered BT sample was carried out using an X-ray diffractometer with monochromatic CuK $\alpha$  radiation ( $\lambda$  = 1.54178 Å) under 40 kV/30 mA over a 2 $\theta$  range 20° to 80° at a scanning rate of 0.01° for 0.10 sec at room temperature (Phillips XPERT-PRO diffractometer model 7602 EA Almelo). The morphology of the BT sample is studied by Field Emission Scanning Electron Microscope (FE-SEM) using FE-SEM (JEOL-7600F) operated at 15 kV.

For electrical measurement, the BT pellets were coated with silver paint on the upper and bottom surfaces and fired at 550°C for 30 min. Impedance measurements were made over the frequency range of 40Hz - 1MHz in the temperature range of 30-150°C, using Agilent A 4294 Impedance Analyzer. An applied voltage of 100 mV is used. The famous software package EC-Lab version 1040 has been employed to analyse the impedance spectroscopy (IS) data based on equivalent circuits.

#### **RESULT AND DISCUSSION**

#### X-ray Diffraction

BT ceramic X-ray diffraction pattern taken at room temperature is shown in Figure 1(a). A single-phase perovskite structure is seen lacking any visible trace of impurity within the background. The magnified XRD pattern in the range of  $2\theta$  from 44.5 to 46.0° (Figure 1(b)) illustrated that the crystal structure of BT was assigned

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to the cubic phase with Pm-3m space group verified from JCPDS database (card no 96-150-7758), since the (200)/ (002) peaks are unseparated (Buttner and Maslen 1992; Veselinovic *et al.* 2014) as reported by other researchers (Lazarevi *et al.* 2010; Itasaka *et al.*, 2018).

The lattice constant of the sample was calculated using the Equation 2

$$\mathbf{a}^2 = \frac{\lambda}{2} \frac{\left[ (\mathbf{h}^2 + \mathbf{k}^2 + \mathbf{l}^2) \right]}{\sin^2 \theta} \tag{2}$$

and the Debye Scherrer equation for calculating the crystal size is given by (Scherrer 1918).

$$\mathbf{D} = \frac{\mathbf{0.98\lambda}}{\beta\cos\theta} \tag{3}$$

where  $\lambda$  is the wavelength of the x-ray ( $\lambda = 1.54178$  Å),  $\theta$  is the Bragg angle and hkl are the Miller indices of the corresponding planes, D is the crystallite size, 0.98 is the crystallite shape factor and  $\beta$  is the full width at half maxima (FWHM) at Bragg's angle (2 $\theta$ ).

The lattice parameters, crystallite size, and unit cell volume of BT ceramic sintered at 1190°C for 2 h were estimated. The lattice parameters and unit cell volume of BT were found to be a = b = c = 4.00522 Å and 64.250 Å<sup>3</sup> respectively. The crystallite size of BT was calculated using equation 2 is found to be 25.7 nm and is similar to the one obtained by other researchers (Al-Naboulsi *et al.* 2016; Yun *et al.* 2019).

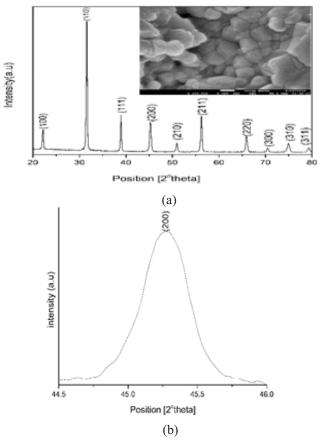


Figure 1: (a) Room Temperature XRD Patterns and inset FESEM Micrograph at x200,000 Magnification for BT Ceramic; (b) Enlarged XRD Patterns from 44.5 to 46.0° of BT Sintered at 1190°C

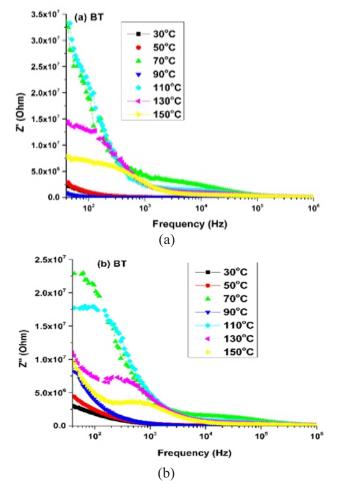


## Surface Morphology

The inset of Figure 1(a) shows the FESEM micrograph of BT ceramics sintered at 1190°C for 2 hrs. It can be sighted that the sintered ceramic sample is dense with the presence of voids and varying microstructures. The appearance of voids in the FESEM image revealed the presence of pores in the sample, an indication of particle agglomeration. The grain size and boundary were observed clearly in non-agglomerated regions while the shape of the grains was indistinguishable. The average grain size of BT ceramic obtained by using the intercept technique (Abrams 1971) was found to be 144.5 nm.

#### Complex Impedance Spectroscopy

Figure 2(a) and 2(b) show the observed variation of real Z' and imaginary Z'' part of complex impedance with a frequency of BT ceramic at a temperature range of 30-150°C. At a temperature of 30, 50, 90°C, and 70, 110, 130, 150°C a monotonous decrease of Z' is seen with increasing frequency which became almost frequency-independent above ~100 Hz and 1000 Hz respectively. It may reasonably be due to the possible release of space charges as a potential consequence of a decrease in the barrier properties of BT ceramics with increased temperatures as



**Figure 2:** Variation of (a) Real (*Z*') and (b) Imaginary (*Z*") with Frequency for BT Ceramic

inferred by (Biswal *et al.* 2014; Mandal *et al.* 2016). An observed increase of Z' at temperature of 30, 50, and 90°C indicated lower conductivity and polarization of the material at temperature. Figure 2(b) shows a decreasing Z' with increasing frequencies up to ~1000 Hz above which it merged seamlessly at a higher frequency. The observed value of Z'' increases at low-frequency at a temperature of 70, 110, 130, and 150°C reached the maxima (Z'<sub>max</sub>) at an elevated temperature of 110, 130, and 150°C. The Z'<sub>max</sub> shifted to higher frequencies with increasing temperature with the presence of relaxation (peaks) at these temperatures. This amply demonstrates the non-Debye nature in the material (Kumar *et al.* 2014).

Conversely, Z" was discovered to decrease monotonously with increasing frequency and decreasing temperatures of 90, 50, and 30°C. This demonstrated that the material exhibited a positive temperature coefficient of resistivity (PTRC) (Kumari, *et al.* 2016). The maximum and minimum values of Z' were 33.3 M $\Omega$ , and 0.93 M $\Omega$ , while values of Z' were 22.8 M $\Omega$ , and 3.00 M $\Omega$  respectively. The maximum and minimum values of Z' were 33.3 M $\Omega$  and 0.93 M $\Omega$ , while values of Z' were 22.8 M $\Omega$  and 3.00 M $\Omega$  respectively.

The likeliest value of the relaxation frequency was d using the relation

$$2\pi f_{max} C = (2\pi f_{max})\tau = 1 \tag{4}$$

e  $\tau = \text{RC}$  is the the likeliest relaxation time lapanda  $et_{\tau} = \tau_o \exp\left(\frac{\Delta E_a}{K_B T}\right)$  st likely relaxing is in line

:e  $\tau_{o}$  is the relaxation time at a pro exponential r, Is a constant associated with the charge carrier ity. By using Equation 4, the activation energy  $(E_{a})$  aluated.

Figure 3 shows the variation of normalized meters of  $Z''/Z''_{max}$  as a function of logarithmic nalized frequency  $f/f_{max}$  and Arrhenius plot of log

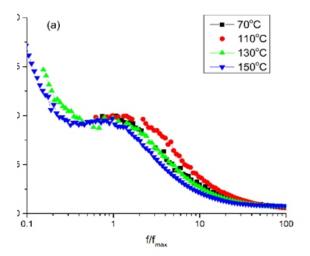


Figure 3: Normalized Parameter of  $Z''/Z''_{max}$  versus  $\log (f/f_{max})$  Plots at various Temperatures.



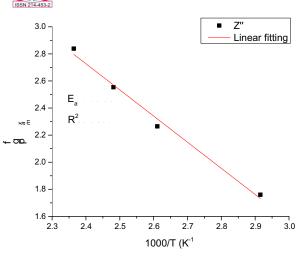


Figure 4: Arrhenius Plot of  $\log (f_{max})$  versus 1000/T for BT Ceramic.

versus 1000/T. In this figure, it can be observed that different peaks nearly coalesce at the same peak irrespective of the temperature. The overlapping of peaks showed relaxation time was temperature independent suggesting the presence of long-range and localized system relaxations. For a localized electron to be mobilised, the support of lattice vibrations was indispensable. For such specific cases, mobile electrons were regarded not to move by vibration, but by hopping motion actuated by vibration from the lattice. Figure 4 presents the Arrhenius plot drawn between  $\log(f_{max})$  versus 1000/T found from Z' plot. The activation energy ( $E_a$ ) determined from the linear fit was found to be 1.12 eV for Z''.

The importance of Nyquist (Cole-Cole) plot is to examine the relaxation and conduction procedures. This was useful in the determination of various criteria relating to the effects of grain and grain boundaries. Figure 5(a) and 5(b) shows the Nyquist (Cole-Cole) plot of imaginary Z" against the real Z at various temperatures. At temperature of 30°C, 50°C, and 90°C, it was observed that the line gradient decreased with a decrease of temperature, and the curve proceeded towards the real Z' axis indicating reduced sample conductivity. This showed evidence of a positive temperature coefficient of resistance (PTCR) effect at the temperature of 30°C, 50°C and 90°C. At temperature of 70, and 110-150°C two depressed semicircles symbolizing both the grain and grain boundaries were discovered.

The high and low-frequency semicircle represented the tangible contribution, respectively, due to grains and the grain boundary effect. (Humera *et al.* 2020). Also, the semicircles comprised of depression, indicating that the distribution of relaxation times remains a non-Debye type. This was due to grain orientation, grain boundary, stress-strain phenomenon, and distribution of atomic defects (Sen *et al.* 2007). To find the parameters concerning grains and grain boundaries, experimental data adhere to an analogous circuit comprising two parallel resistance-capacitance (RC) elements or

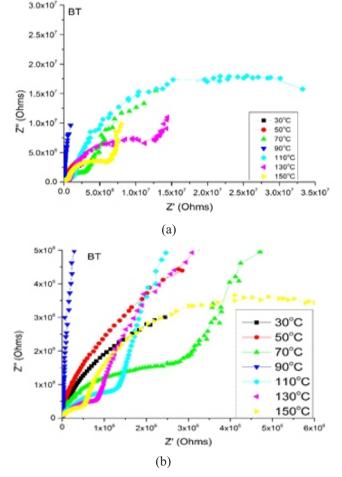


Figure 5: (a) Variation of Z'' with Z' (Nyquist or Cole-Cole) Plot for BT Ceramic.(b) when (*Enlarged*).

replacing the capacitor with a constant phase element (CPE) as a parallel-connected resistance-constant phase element (R-CPE) network (Mondal et al. 2017). The Constant phase element (CPE, denoted by Q) was used to overcome dispersion and non-linearity in the value from ideal behavior or because the semicircles exhibit a non-Debye nature. It is defined as  $C = (R^{1-n} Q)^{1/n}$  where *n* represents the degree of deviation with respect to the value of the pure capacitor. It is zero (0) and unity (1) for an ideal resistor and capacitor respectively (Barick et al. 2013). The circuit fitting parameter was performed using EC-Lab Software V10.40 with the modeled circuits as shown in Figure 6. Table 1 provides the fitting parameters for the equivalent circuit (R-CPE). The relaxation time for the grain  $(\tau_{a})$  and grain boundary  $\tau_{\rm gb}$  was calculated from Equation 4. It was observed that the values of grain resistance  $(R_g)$  decreased with an increasing temperature this indicated an increase of conductivity with rising temperature.

#### **Complex Dielectric Modulus**

Modulus analysis typically represents an alternative method for properly examining the material's electrical properties and improving the various impacts present in the ceramic to specific time constants for relaxation. The real (M') and imaginary (M'') parts of the modulus are calculated using Equations 6 and 7:



 
 Table 1: Parameters Obtained from Temperature Dependent Impedance Spectroscopy Data for the BT Ceramic.

Temp.	$R_g  imes 10^6$		<sup>12</sup> n <sub>1</sub>	$\rm R_{gb} \times 10^{6}$	$Q_2 \times 10^{-12}$	$\mathbf{n}_2$	$\tau_g \times \ 10^{-6}$	$\tau_{gb} \times ~10^{-3}$
(°C)	(Ω)	(F)		(Ω)	(F)		(s)	(s)
70	2.96	5.16	0.92	7.38	2.03	0.79	5.52	4.86
110	1.98	33.1	0.79	3.69	26.8	1.0	5.73	0.99
130	1.01	23.9	0.83	1.32	29.7	0.99	2.75	5.36
150	0.64	63.1	0.77	0.70	37.7	0.97	2.00	2.89

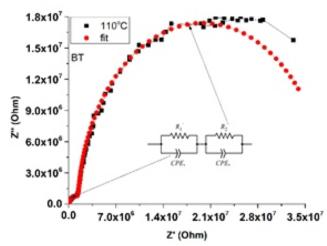
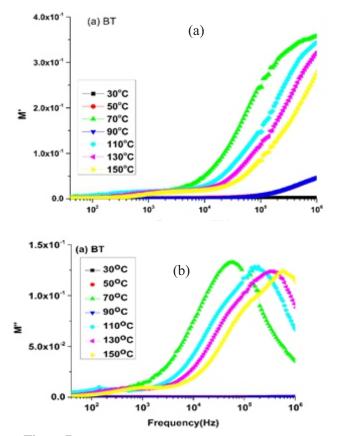


Figure 6. The Model of an Equivalent Circuit used for Fitting of Impedance Spectra



**Figure 7**: Frequency-temperature Dependence of (a) Real Part (*M*') and (b) Imaginary Part (*M*'') for BT Ceramic.

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$$\mathbf{M}' = \frac{\mathbf{\varepsilon}'}{\mathbf{\varepsilon}'^2 + \mathbf{\varepsilon}''^2} \tag{6}$$

$$\mathbf{M}^{\prime\prime} = \frac{\boldsymbol{\varepsilon}^{\prime\prime}}{\boldsymbol{\varepsilon}^{\prime 2} + \boldsymbol{\varepsilon}^{\prime\prime 2}} \tag{7}$$

Figure 7(a) and 7(b) show the real (M) and imaginary (M) frequency dependence parts of electrical modulus at 30-150°C. It was noticed that the M value at lower frequencies is almost zero, affirming the presence of a significant electrode and/or ionic polarization in the temperature ranges examined. The M value increased continuously with increasing frequency and the dispersion changes to a higher frequency towards a high-frequency limit. This merely occurred because of the presence of the conduction procedure owing to the short-range mobility of the carriers in charge.

It can be observed from Figure 7(b) that M' values increased with increasing frequency and temperature. It reached a maximum  $(M'_{max})$  with the appearance of impedance relaxation peaks which appeared at 70°C,  $110^{\circ}$ C,  $130^{\circ}$ C, and  $150^{\circ}$ C at frequencies below  $10^{4}$  Hz. At that point, M' started to decrease with an increasing frequency above the maxima. For rising temperatures, the peaks shifted to a higher-frequency side. This means the relaxation processes are temperaturedependent in these materials. The intensity of this peak decreased slowly because the impedance values decreased with temperature rise. The asymmetric module peaks change to a higher frequency side revealing the association of mobile ion movements (Jacob et al. 2015; Sharma et al. 2015). The broadening of the peaks is because of the spread of relaxation time with the different time constants (Mondal et al. 2017) and so relaxation was of a non-Debye type. The area below and above the relaxation peaks in M' defined the field where the load carrier can travel for both long and short distances (Naidu et al. 2018). The difference in the charging carriers indicates a hopping mechanism makes the electrical conductivity important at a higher temperature. At a temperature of 30, 50, and 90°C the value of M' appeared to become independent of both the frequency and temperature. It presented a possible release of space charge.

Figures 8 shows the modulus scaling behavior by plotting normalized parameters and the Arrhenius relation of BT ceramic. The scaling action of the modulus provided an insight into the dielectric processes within the material (Das *et al.* 2007). In Figure 8 it was observed that the peaks overlapped at 70-130°C, suggesting temperature-independent behavior of the complex processes in the ceramics (Saha and Sinha



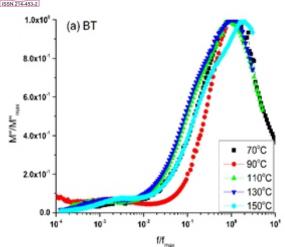
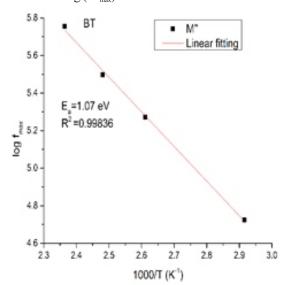


Figure 8: Normalized parameters M"/M"<sub>max</sub> versus  $\log(f/f_{max})$ 



**Figure 9:** Arrhenius plot of  $\log (f_{max})$  versus 1000/T for BT ceramic.

2002). The peak that appeared at 150°C shifted towards a higher frequency side, activating another relaxation process. The dielectric analysis of material can, therefore, be investigated by means of modulus scaling (Das *et al.* 2007). Figure 9 shows the Arrhenius plot of found to be 1.07 eV which was close to the activation the resistance of bulk  $(R_h)$ energy value found from Z" in Figure 4.

Figure 10 shows the Arrhenius plot of  $\ln \tau_{o}$  and  $\ln AC$  Conductivity Analysis  $\tau_{ab}$  with the inverse of absolute temperature (1000/T) The AC conductivity ( $\sigma_{ac}$ ) was computed using Equation 8. for BT ceramic. The activation energies for relaxation were calculated from Equation 8 to be 0.46 and 0.12 eV for grain and grain boundary respectively.

The temperature dependence of the complex modulus spectrum (i.e. M' vs M') spectrum of BT ceramic at 30-150°C is shown in Figure 11. At a temperature of 70°C, 110, 130, and 150°C two semicircular arcs in low and high frequency small and wide semicircular arcs are due to poor three clearly distinguishable areas in the conducting

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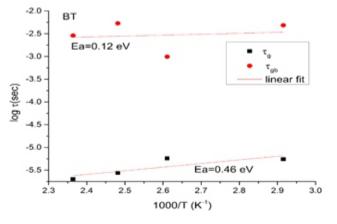


Figure 10: Arrhenius Plot of ln  $\tau_{_g}$  and ln  $\tau_{_{gb}}$  versus 1000/T for BT Ceramic.

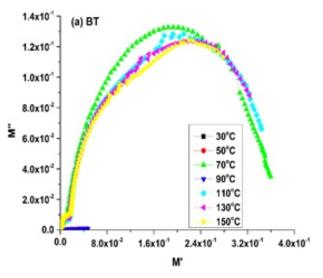


Figure 11: Plot of *M*' versus *M*' for BT Ceramic.

grain boundary effects rather than dominant grain effects and bulk or grain reaction, respectively (Lvovich 2012; Saghrouni et al, 2014). The relaxation in the material was non-Debye form as the semicircular arc centers are below the actual M' axis. It appeared that the intercept of the semicircles on the real axis tend to change with increasing temperature to higher values of M. It showed that capacitance increased with increasing temperature. This  $log(f_{max})$  versus 1000/T found from M'. The activation affirmed the NTCR type behavior of the BT ceramic since energy obtained from the slope of the straight line was the capacitance of bulk  $(C_b)$  was inversely proportional to

$$\sigma_{ac} = \omega \varepsilon' \varepsilon_{o} \tan \delta \tag{8}$$

where  $\omega = 2\pi f$ , f is the frequency,  $\varepsilon$ ' the real dielectric constant,  $\varepsilon_{0}$  permittivity of free space and tan  $\delta$  is the loss tangent.

The plot of log  $\sigma_{ac}$  vs log f at different temperatures is shown in Figure 12. It depicted wide dispersion and frequencies regions were observed. Low and high- the conductivity increased as temperature increased with



regions. However, at high-frequency, the conductivity exhibits frequency dispersion. The AC conductivity increased with a frequency and temperature rise similar to the behavior observed in many of the polymers and semiconductors (Dutta *et al.* 2008). This is due to the change in the mobility of the sample charging carriers as the temperature increased. In the high-frequency field, however, the conductivity showed dispersion with an increasing slope for all the temperatures. The frequency at which the slope changed is known as the frequency of hopping which shifted with an increasing temperature towards the higher values. The power-law equation expressed the total conductivity ( $\sigma$ ) of a given material at a given temperature (Jonscher 1996),

$$\sigma_{ac} = \sigma_{dc} + A\omega" \tag{9}$$

where  $\sigma_{dc}$  is the extrapolated conductivity value of the frequency (at  $\omega = 0$ ). independent part, and A is the AC coefficient,  $n \ (0 \le n \le 1)$  the correlation exponent of the ions. Thermally activated quantities are typically given as A and n, therefore, electrical conduction remained a thermally activated operation. As Jonscher stated, the possible source of frequency dependence on conductivity lies within the relaxation phenomena that arose from mobile charging carriers.

Figure 13 shows the Arrhenius plot of AC and DC

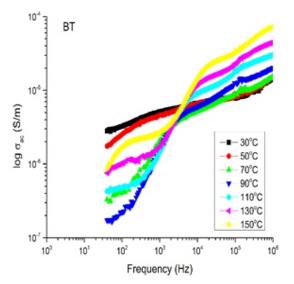


Figure 12: Variation of AC Conductivity with Frequency for BT Ceramic.

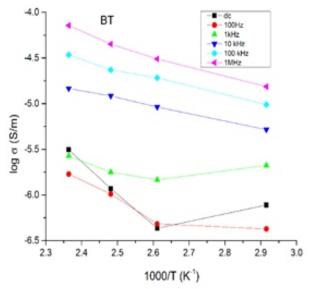


Figure 13: Variation of AC Conductivity with an Inverse Absolute Temperature at Various Frequencies

conductivity versus inverse temperature of BT ceramic. It was observed that the conductivity of the high-temperature component which can be related to the bound charge carriers in the ceramics increased. On top of that, it was observed that the slopes of AC conductivity at a frequency of 100 Hz, 1 kHz, and DC conductivity were not linear but showed little alterations at different temperature regions. This may be attributed to contributions from various areas, like grain and grain boundary (Mahboob *et al.* 2006). The AC and DC conductivity activation energies were carefully determined with the following formula:

$$\sigma_{ac} = \sigma_{o} exp\left(\frac{-E_{ac}}{k_{B}T}\right)$$
(10)

$$\sigma_{dc} = \sigma_{o} exp\left(\frac{-E_{dc}}{k_{B}T}\right)$$
(11)

whence

$$\ln \sigma_{\rm dc \, or \, ac} = \ln \sigma_{\rm o} \tag{12}$$

where  $\sigma_{o}$  is pre-exponential factor, k is Boltzmann constant, T is the absolute temperature,  $E_{ac}$  and  $E_{dc}$  represent activation energy of AC and DC conductivity, respectively.

The apparent activation energy  $E_a$  is obtained from the linear lowest-square fit slope of the conductivity data in Equation 5, and the result is shown in Table 2. At higher temperature ranges, the activation energies DC and AC (100 and 1 kHz) were greater than those at lower

Table 2: The AC and DC conductivity activation energies (eV) for BT ceramic.

Sample		BT				
Conductivity Activation Energy (eV)						
AC						
Temperature	DC	100 kHz	l kHz	10 kHz	100 kHz	1 MHz
(°C)						
70-150	-	-	-	0.48	0.56	0.68
70-110	0.48	0.10	0.30	-	-	-
130-150	2.02	1.28	0.60	-	-	-



temperature ranges. The  $E_a$  values 1 kHz and 1 MHz were in agreement with the earlier report (Humera et al. 2020; Tomar *et al.* 2020). Accordingly, the calculation of activation energy was based on the considerable degree of ionization of the oxygen vacancy (Moretti and Michel, 1987). Activation energy > 1.0 eV related to single ionized vacancies (Ang et al., 2000) and/or electronic mobility in space charge regions (Ortega et al. 2007). In this manner, the conduction cycle within this temperature range could be due to the BT ceramic's hopping of charge carriers and or individually ionized oxygen vacancies (Badapanda et al. 2014). In perovskite ferroelectric, oxygen vacancies are typically regarded as one of the mobile charging carriers and so the ionization of oxygen vacancies produced electrons for conduction.

## CONCLUSION

A combination of solid-state and high energy ball milling technique has been used to synthesise nanocrystalline BaTiO<sub>3</sub> powder. A single-phase perovskite cubic was observed at room temperature from the XRD pattern. FE-SEM image showed that sample was dense and had different microstructures with a certain amount of porosity. Both impedance and modulus analysis at 70, 110, 130 and 150°C supported the typical behavior of a negative temperature coefficient of resistance (NTCR) of the materials which would be effective for the fabrication of a highly sensitive thermistor and suggested the relaxation to be non-Debye type. Conversely, at a temperature of 30, 50, and 90°C the material showed a PTCR effect. The activation energy values suggested the carrier transport was due to hopping conduction. The AC conductivity was found to obey the universal power law, as hinted by Jonscher. The magnitude of the activation energy followed the Arrhenius relationship.

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