

Investigations of impedance and electric modulus properties of $\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$ ceramics

Rajiv Ranjan¹, Nawnit Kumar², Banarji Behera³, R. N. P. Choudhary^{4*}

¹Department of Physics, Jamshedpur Co-operative College, Kolhan University, Jharkhand 831036, India

²Department of Physics and Meteorology, Indian Institute of Technology, Kharagpur, India 721302

³School of Physics, Sambalpur University, Jyoti Vihar, Odisha 768019, India

⁴Department of Physics, ITER, S.O.A. University, Odisha 751030, India

*Corresponding author. Tel: (+91) 674 2350181; Fax: (+91) 6742351880; E-mail: crnpfl@gmail.com

Received: 06 January 2013, Revised: 20 March 2013 and Accepted: 16 April 2013

ABSTRACT

$\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$ (PSZT; $x = 0.00, 0.03$ and 0.06) ceramic samples were prepared by high temperature mixed oxide method. Using complex impedance spectroscopy (CIS) technique, the complex impedance (Z^*) and modulus (M^*) properties of the materials were analyzed within a wide range of temperature and frequency. Impedance analysis indicates the presence of mostly bulk resistive (grain) contributions which is found to decrease with the increase in temperature. It suggests about the negative temperature coefficient of resistance (NTCR) type behaviour of the materials. Complex modulus plots exhibit the presence of grain (bulk) as well as grain boundary contributions in the materials and also support their NTCR type behaviour. Both the complex impedance and complex modulus plots confirm the presence of non-Debye type of relaxation in the materials. At higher temperatures, bulk resistance is found to increase with the increase in Sm^{3+} concentration in PSZT along with the increase in relaxation phenomenon. Copyright © 2014 VBRI press.

Keywords: Ceramics; complex impedance spectroscopy (CIS); impedance; modulus.



Rajiv Ranjan is an Assistant Professor in the Department of Physics, Jamshedpur Co-operative College, Jamshedpur under Kolhan University, Jharkhand, India. He has completed his Ph. D degree in the field of materials physics from Ranchi University with the great support of Prof. R. N. P. Choudhary. For future research work in the field of ferroelectric and multiferroic materials, he has developed research facilities inside his department supported by University Grant Commission, India. He is trying to develop research capabilities in M.Sc. students in the field of material science. He is also supervising Ph. D scholars.



Nawnit Kumar was born in Kahalgaon, Bhagalpur, Bihar, India in 1981. He received M. Sc. (Physics) degree from Tilka Manjhi Bhagalpur University, Bhagalpur, Bihar, India in 2005. He is now doing Ph. D on Multiferroic materials in the Department of Physics and Meteorology, Indian Institute of Technology, Kharagpur, India.

R. N. P. Choudhary is one of the pioneer researchers and contributors in the field of ferroelectric, multiferroic and related advanced materials in India and abroad. He has done Ph. D from University of Edinburgh, U.K. He has been actively engaged in teaching and research for the last 40 years at the institutes of national and international repute such as



National Institute of Technology, Jamshedpur (India), Indian Institute of Technology, Kharagpur (India). He has successfully completed many research projects. He has guided more than four dozens Ph. D students and published more than 600 research papers in national/international journals. He is the member of editorial board of many national/international journals. He is one of the referees of research schemes from DST and CSIR. At present, he is working as a Professor in Department of Physics, Institute of Technical Education and Research (SOA University) Bhubaneswar (India) and many research scholars are doing their research works under his supervision.

Introduction

Now a days, Lead zirconate titanate [PZT; $\text{Pb}(\text{Zr}_{1-y}\text{Ti}_y)\text{O}_3$] [1-2] is an interesting dielectric, piezoelectric and ferroelectric material that has tremendous applications in the form of sensor, actuator and transducer [3-5]. It has perovskite ABO_3 structure (A = mono or divalent, B = trihexavalent ions) in which A-site is occupied by Pb^{2+} and B-site by Zr^{4+} and/or Ti^{4+} [6-8]. It is the solid-solution of ferroelectric PbTiO_3 ($T_c = 763$ K) & antiferroelectric PbZrO_3 ($T_c = 503$ K) presenting tetragonal phase in Ti-rich composition and rhombohedral in Zr-rich composition. Its

electrical and electromechanical properties rise to a great extent at Morphotropic Phase Boundary (MPB) between these two phases [9-11].

Depending upon specific requirements, physical properties and device parameters of PZT based compounds may be considerably modified with the variations in Zr/Ti ratio, synthesis techniques, particle size, calcination & sintering temperatures, etc [1, 12-16]. With suitable substitution(s) at A and/or B site(s), its material properties can be significantly changed. Interestingly, PZT modified with higher valence additives (like La^{3+} , Sm^{3+} or Nb^{5+}) results in the increased value of dielectric constant, electromechanical properties, etc. where electroneutrality is maintained by production of Pb-vacancies. On the other hand, PZT modified with lower valence additives (like K^+ , Fe^{3+} or Mg^{2+}) results in the reduced value of dielectric constant, electromechanical properties, etc [17-18] by the production of O-vacancies.

Since the past few years, electronic and electro-optic device applications of La-modified PZT ceramics [19-22] have encouraged the research works in this field with others additives. The rare earth element samarium (Sm^{3+}) also fulfils the basic requirements for its substitution to PZT [23-24] such as stability of perovskite phase, appropriate ionic size, high Curie-Weiss temperature, etc. But, only few works have been reported on PZT ceramics modified with samarium [25-29]. Interestingly, it is found that PSZT; Zr/Ti = 45/55 has shown a very high dielectric constant and low tangent loss [30]. In this paper, we report about the analysis of complex impedance and modulus properties of pure and Sm^{3+} -modified PZT ceramics having Zr/Ti ratio 45/55 using complex impedance spectroscopy (CIS) technique.

Complex impedance spectroscopy (CIS) [31] is an effective experimental technique used to characterize micro-structural and electrical properties of polycrystalline electro-ceramic materials. It helps to separate real and imaginary components of complex electrical parameters so as to get the true picture of material properties. It enables us to resolve the relaxation contributions, like, bulk effects, grain boundaries and electrode interface effects in the frequency domain of materials.

In complex impedance spectroscopy, frequency dependent behaviour of the materials may be expressed in terms of complex impedance (Z^*), complex electric modulus (M^*), complex dielectric constant (ϵ^*) and tangent loss ($\tan\delta$), which are related to each other through the relations: $Z^* = Z' - jZ''$, $M^* = 1/\epsilon^*(\omega) = M' + jM''$, $\epsilon^* = \epsilon' - j\epsilon''$ and $\tan\delta = \epsilon''/\epsilon'$, where (Z' , M' , ϵ') and (Z'' , M'' , ϵ'') are the real and imaginary components of impedance, modulus and permittivity respectively, $j = \sqrt{-1}$ the imaginary factor, $C_0 =$ vacuum capacitance.

Experimental

The polycrystalline ceramic samples of $\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$ [where $x = 0.00, 0.03$ and 0.06] were prepared by high temperature mixed oxide method using ingredient oxides; PbO , ZrO_2 , TiO_2 and Sm_2O_3 (99.9% purity, M/S Loba Chemie, Inc. Bombay, India) in appropriate stoichiometry. 3% of extra lead oxide was added to the mixture to compensate lead loss at high

temperature calcinations and sintering. The oxide powders were mixed thoroughly, first in, dry air and, then, in alcohol (methanol) medium using agate mortar and pestle. The well-mixed oxides were, first, calcined at 950°C for about 10 hr in alumina crucibles. The processes of grinding and calcination were repeated at temperature intervals of 50°C and the formation of desired compounds has been confirmed (through XRD technique) with the mixture calcined at 1100°C for 10 hr. Using small amount of polyvinyl alcohol (PVA) as binder, the calcined powders were cold pressed into thin cylindrical pellets in a steel die punch with the help of a hydraulic press at a pressure of $4 \times 10^6 \text{ N/m}^2$. These pellets were, subsequently, sintered at an optimized temperature 1200°C for 10 hr in alumina crucibles so as to get nearly 97% of theoretical density.

For the measurement of electrical properties, smooth flat surfaces of each pellet were electroded with high purity air-drying silver paint. The pellets were, then, dried at 150°C for 4 hr to remove moisture. Now, using computer based arrangements of phase sensitive multimeter (PSM N4L; Model 1735), furnace, variac, etc., a number of electrical data were recorded at different temperatures and frequencies through which various electrical parameters were calculated.

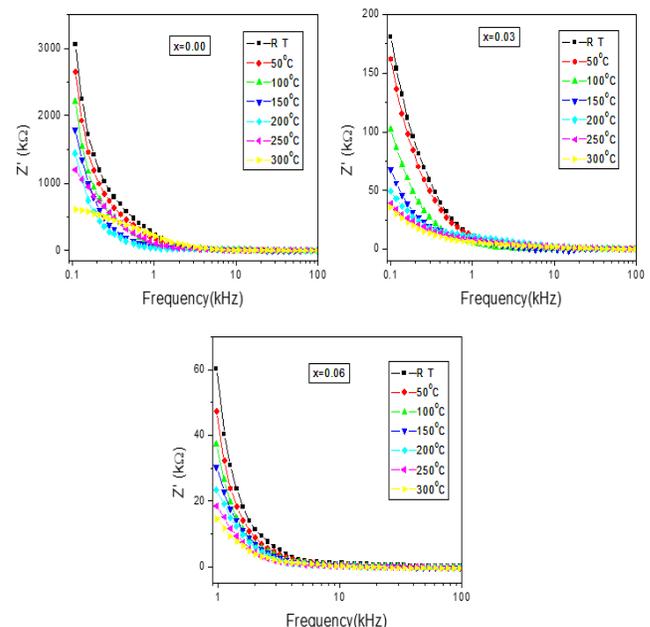


Fig. 1. Variation of Z' with frequency at different temperatures of $\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$; $x = 0.00, 0.03$ and 0.06 .

Results and discussion

Complex impedance analysis

The variation of real part of impedance (Z') of the ceramic samples $\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$ [$x = 0.00, 0.03$ and 0.06] with frequency at different temperatures is shown in **Fig. 1**. Z' is found to decrease with the increase in temperature and achieves a low value at higher temperatures. It exhibits low frequency dispersion mainly due to polarization and then monotonous decrease with the increase in frequency. This shows increase in material's ac-conductivity with the increase in temperature as well as frequency. At higher frequencies, Z' achieves nearly a very

low constant value and becomes almost independent of frequency as well as temperature. At lower frequencies, Z' decreases with the increase in frequency supporting a slow dynamic relaxation process in the materials probably due to space charge that gets released at higher frequencies [32-36]. It is also observed that Z' exhibits a decreasing trend with the increase in Sm^{3+} -concentration in PSZT from $x = 0.00$ to 0.06 upto about 300°C .

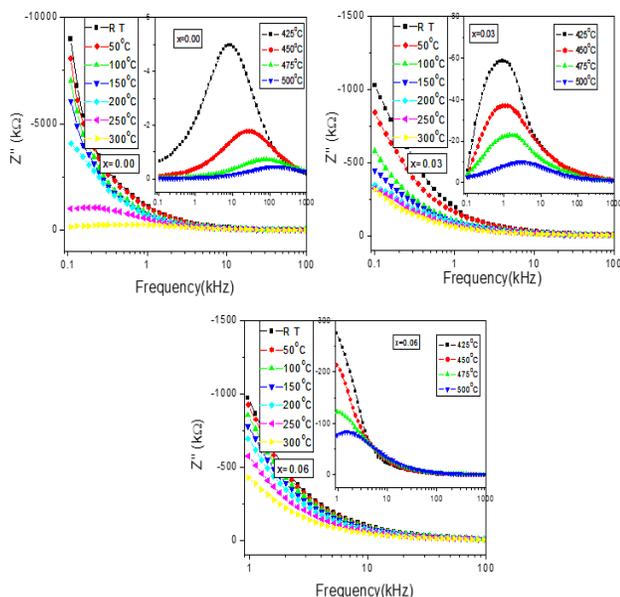


Fig. 2. Variation of Z'' with frequency at different temperatures of $\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$; $x = 0.00, 0.03$ and 0.06 .

Fig. 2 shows the variation of imaginary part of impedance (Z'') of the pure and Sm-modified PZT compounds with frequency at different temperatures. At lower temperatures, Z'' shows behaviour similar to Z' but some peaks are observed at higher temperatures. These peaks are found to shift towards higher frequencies with the increase in temperature exhibiting a decreasing trend of peak height which finally merge in the high frequency region. It suggests about thermally activated relaxation process in the materials and indicates reduction in bulk resistance with the increase in temperature. It may be due to presence of space charge polarization at lower frequencies and its elimination at higher frequencies. The observed asymmetric broadening of peaks indicates about the presence of some electrical processes in the materials with spread of relaxation time. This may be due to the presence of immobile species at low temperature and defects at higher temperatures [37-38]. Though, these peaks have not been observed at lower temperatures which may be due to the weak current dissipation in the materials or beyond the frequency of measurement. With the increase in Sm^{3+} -content from $x = 0.00$ to 0.06 , the peak height gradually increases and found to shift towards the shorter frequencies.

Fig. 3 shows the complex impedance spectrum ($Z' \sim Z''$; Nyquist plot) of PSZT compounds at selected higher temperatures. Appearance of single arc or semicircle at a particular temperature shows that electrical properties in the materials arise mainly due to the contribution of bulk effects. At lower temperatures only single arcs (not shown)

are obtained which are transformed into semicircles at higher temperatures. The formation of full, partial or no semicircles mainly depends on the strength of relaxation and also experimentally available frequency range [39]. The electrical process taking place within the materials may be modelled (as an RC circuit) on the basis of the brick-layer model [31]. In Nyquist plots, intercept of semi-circular arcs on real Z' -axis gives the value of bulk resistance (R_b) which is found to decrease with increase in temperature. It suggests about the negative temperature coefficient of resistance (NTCR) type behaviour of the PSZT compounds. These plots exhibit depressed semicircles having centres lying below the real axis (not shown), this confirms the presence of non-Debye type of relaxation phenomenon in the materials [40-41]. At higher temperatures, R_b is found to increase with the increase in Sm^{3+} -concentration in the materials.

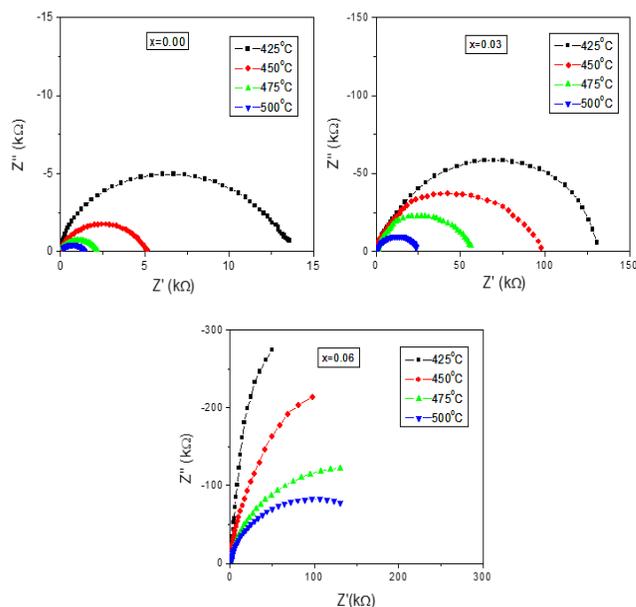


Fig. 3 Complex impedance plots ($Z' \sim Z''$) of $\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$; $x = 0.00, 0.03$ and 0.06 .

Complex modulus analysis

The variation of real part of electric modulus (M') of $\text{Pb}_{1-x}\text{Sm}_x(\text{Zr}_{0.45}\text{Ti}_{0.55})_{1-x/4}\text{O}_3$ [$x = 0.00, 0.03$ and 0.06] with frequency at different temperatures is shown in **Fig. 4**. M' is found to be very low at lower frequencies and exhibits an increasing trend with the increase in frequency. It shows continuous dispersion on increasing frequency which may be due to short range mobility of charge carriers under the action of an induced electric field. But at higher temperatures, M' increases rapidly with the increase in frequency as well as temperature which attributes temperature dependent relaxation process in the materials. Also it is observed that the dispersion region shifts towards higher frequencies suggesting long-range mobility of charge carriers [42]. The observed plateau region or its tendency indicates about frequency invariant dc behaviour of the materials. With Sm^{3+} substitution to $\text{Pb}(\text{Zr}_{0.45}\text{Ti}_{0.55})\text{O}_3$ from $x = 0.00$ to 0.06 , M' is found to high when $x = 0.03$.

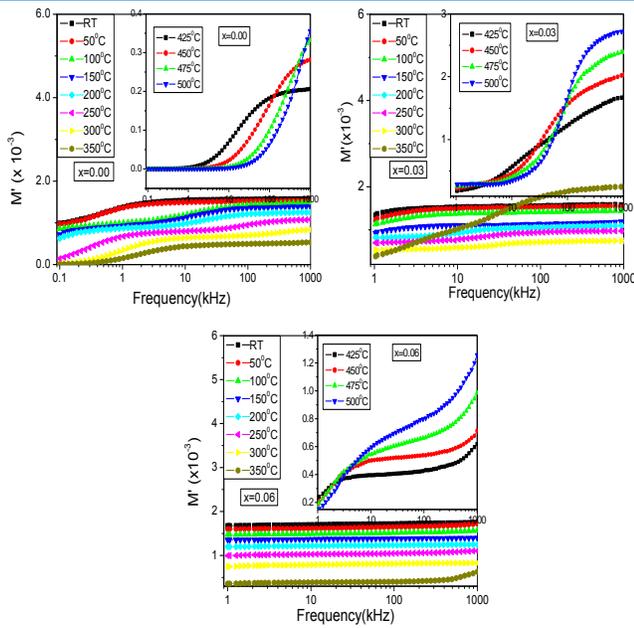


Fig. 4. Variation of M' with frequency at different temperatures of $Pb_{1-x}Sm_x(Zr_{0.45}Ti_{0.55})_{1-x/4}O_3$; $x = 0.00, 0.03$ and 0.06 .

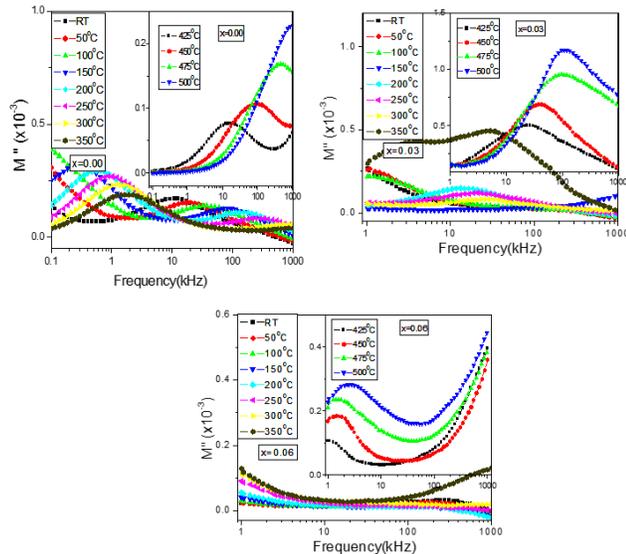


Fig. 5. Variation of M'' with frequency at different temperatures of $Pb_{1-x}Sm_x(Zr_{0.45}Ti_{0.55})_{1-x/4}O_3$; $x = 0.00, 0.03$ and 0.06 .

Fig. 5 shows the variation of imaginary part of electric modulus (M'') of the PSZT compounds with frequency at different temperatures. It exhibits asymmetric modulus peaks or tendency at low as well as at high frequencies. The low frequency peak suggests that ions can move over long distances whereas high frequency peak supports the confinement of ions in their potential well [43]. The observed asymmetry in peak broadening indicates the spread of relaxation time with different time constant which supports the non-Debye type of relaxation in the materials. It is also observed that the relaxation frequencies shift towards higher frequencies with increase in peak height as temperature increases. It indicates thermally activated behaviour of relaxation time. The low value of M'' observed at lower frequencies may occur due to the absence of electrode polarization phenomena. It is observed that M''

corresponding to the peaks is maximum in case of PSZT with $x = 0.03$.

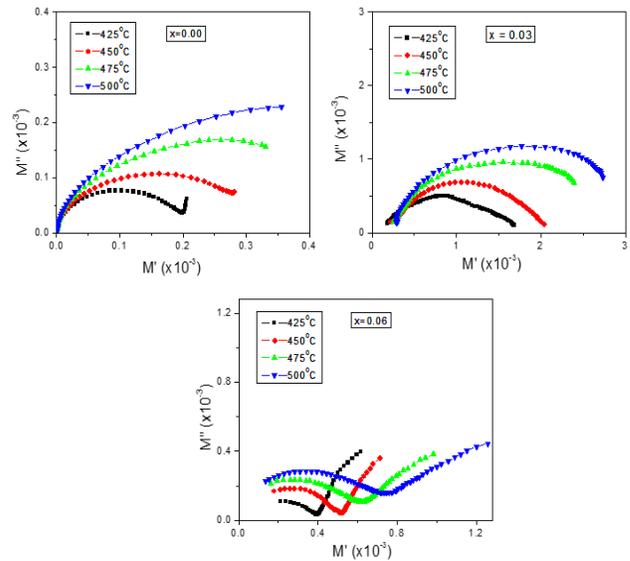


Fig. 6. Complex modulus plots ($M''-M'$) of $Pb_{1-x}Sm_x(Zr_{0.45}Ti_{0.55})_{1-x/4}O_3$; $x = 0.00, 0.03$ and 0.06 .

The temperature dependence of complex modulus spectrum (i.e. M' vs. M'' spectrum) of PSZT is shown in **Fig. 6**. At higher temperatures, semi-circular arcs are observed in the form of two semicircular arcs (or tendency) which indicates the presence of both grain and grain boundary contributions in the materials. Their centres appear to lie below the real M' -axis so it indicates spread of relaxation with different (mean) time constant and hence supports the non-Debye type of relaxation in the materials. Intercepts of these semicircles on real axis appear to shift towards higher values of M' with the increase in temperature. It indicates increase in capacitance with the increase in temperature and supports NTCR type behaviour of the materials since the bulk capacitance (C_b) is inversely proportional to the bulk resistance (R_b). Though, at higher temperatures and lower frequencies, tails are not fully evolved to its true shape which may be due to frequency and temperature limitations.

Conclusion

Polycrystalline samples of $Pb_{1-x}Sm_x(Zr_{0.45}Ti_{0.55})_{1-x/4}O_3$; $x = 0.00, 0.03$ and 0.06 were prepared by mixed oxide method at $1200^\circ C$ sintering temperature. Real and imaginary parts of complex impedance and modulus properties of the materials were investigated by using complex impedance spectroscopy (CIS) technique. At a particular temperature, the observed single arc in the form of single/double semicircles (or tendency) in both the complex impedance and modulus plots confirms about the formation of samples in single phase. Impedance analysis indicates the presence of mostly bulk (grain) resistive contributions in the materials whereas complex modulus plots shows the presence of grains as well as grain boundary contributions in the materials. It is due to the fact that impedance plot highlights the phenomenon with largest resistance whereas electric modulus plot highlights the phenomenon with smallest capacitance. Due to the large difference between

resistive values of grains and grain boundaries, it is not possible to get two semicircles on the same impedance plot [44]. Both impedance and modulus analysis support the typical behaviour of negative temperature coefficient of resistance (NTCR) of the materials. They also confirm the presence of non-Debye type of relaxation phenomenon in the materials. With the increase in Sm^{3+} concentration in $\text{Pb}(\text{Zr}_{0.45}\text{Ti}_{0.55})\text{O}_3$ upto 6%, the bulk resistance exhibits an increasing trend at higher temperatures along with the increase in relaxation phenomenon in the materials.

Acknowledgements

The author, R. Ranjan would like to acknowledge University Grant Commission for MRP-PSJ-001/11-12 and N. Kumar would like to acknowledge CSIR, New Delhi (India) for financial support.

Reference

- Shirane, G.; Suzuki, K. *J. Phys. Soc. Jpn.* **1952**, 7, 333.
DOI: [10.1143/JPSJ.7.333](https://doi.org/10.1143/JPSJ.7.333)
- Lines, M. E.; Glass, A. M. *Principles and Applications of Ferroelectrics and Related Materials*, (Oxford University Press, Oxford, **1977**).
- Sambasivarao, K.; Prasad, T. N. V. K. V.; Subrahmanyam; A. S. V.; Lee, J. H.; Kim, J. J.; Cho, S. H. *Mater. Sci. Eng. B* **2003**, 98, 279.
DOI: [10.1016/S0921-5107\(03\)00064-3](https://doi.org/10.1016/S0921-5107(03)00064-3)
- Neurgaonkar, R. R.; Oliver, J. R.; Nelson J. G. *Mater. Res. Bull.* **1991**, 26, 771.
DOI: [10.1016/0025-5408\(91\)90066-U](https://doi.org/10.1016/0025-5408(91)90066-U)
- Cross, L. E. *Mater. Chem. Phys.* **1996**, 43, 108.
DOI: [10.1016/0254-0584\(95\)01617-4](https://doi.org/10.1016/0254-0584(95)01617-4)
- Hiremath, B. V.; Kingon, A. I.; Biggers, J. V. *J. Am. Ceram. Soc.* **1983**, 66, 790.
DOI: [10.1111/j.1151-2916.1983.tb10564.x](https://doi.org/10.1111/j.1151-2916.1983.tb10564.x)
- Ohno, T.; Takahashi, M.; Tsubouchi, N. *Jpn. Soc. Powder Powder Metall.* **1973**, 20, 154.
- Chandratreya, S. S.; Fulrath, R. M.; Pask, J. A. *J. Am. Ceram. Soc.* **1981**, 64, 422.
DOI: [10.1111/j.1151-2916.1981.tb09883.x](https://doi.org/10.1111/j.1151-2916.1981.tb09883.x)
- Jaffe, B.; Crook, W. R.; Jaffe, H. *Piezoelectric Ceramics*, (Academic Press, New York, **1971**)
- Zhong, W. L.; Wang, Y. G.; Zhang, P. L.; Qu, B. D. *Phys. Rev. B* **1994**, 50, 698.
DOI: [10.1103/PhysRevB.50.698](https://doi.org/10.1103/PhysRevB.50.698)
- Mabud, S. A. *J. Appl. Cryst.* **1980**, 13, 211.
DOI: [10.1107/S0021889880011958](https://doi.org/10.1107/S0021889880011958)
- Smyth, A. P. *Dielectric Behavior and Structure*, (McGraw Hill Publishers, New York, **1955**)
- Haertling, G. H. *J. Am. Ceram. Soc.* **1999**, 82, 797.
DOI: [10.1111/j.1151-2916.1999.tb01840.x](https://doi.org/10.1111/j.1151-2916.1999.tb01840.x)
- Haertling, G. H. *Ferroelectrics* **1987**, 75, 25.
DOI: [10.1080/00150198708008208](https://doi.org/10.1080/00150198708008208)
- Snow, G. S. *J. Am. Ceram. Soc.* **1973**, 56, 91.
DOI: [10.1111/j.1151-2916.1973.tb12365.x](https://doi.org/10.1111/j.1151-2916.1973.tb12365.x)
- Camargo, E. R.; Leite, E. R.; Longo, E. *J. Alloys Compd.*, **2009**, 469, 523.
DOI: [10.1016/j.jallcom.2008.02.008](https://doi.org/10.1016/j.jallcom.2008.02.008)
- Qiu, W.; Hng, H. H. *Mater. Chem. Phys.* **2002**, 75, 151.
PII: [S0254-0584\(02\)00045-7](https://doi.org/10.1016/S0254-0584(02)00045-7)
- Ramam, K.; Lopez, M. *J. Alloys Compd.* **2008**, 466, 398.
DOI: [10.1016/j.jallcom.2007.11.055](https://doi.org/10.1016/j.jallcom.2007.11.055)
- Dutta, S.; Choudhary, R. N. P.; Sinha, P. K. *J. Alloys Compd.* **2007**, 430, 344.
DOI: [10.1016/j.jallcom.2006.05.040](https://doi.org/10.1016/j.jallcom.2006.05.040)
- Shannigrahi, S. R.; Choudhary, R. N. P. *J. Eur. Ceram. Soc.* **2004**, 24, 163.
DOI: [10.1016/S0955-2219\(03\)00316-9](https://doi.org/10.1016/S0955-2219(03)00316-9)
- Rai, R.; Mishra, S.; Singh, N. K. *J. Alloys Compd.* **2009**, 487, 494.
DOI: [10.1016/j.jallcom.2009.07.161](https://doi.org/10.1016/j.jallcom.2009.07.161)
- Chiou, B. S.; Kuo, J. N.; Dai, H. T. *Journal of Electronic Materials*, **1990**, 19(4), 393.
DOI: [10.1007/BF02651302](https://doi.org/10.1007/BF02651302)
- Chon, U.; Bum, K.; Jang, H. M.; Yi, G. C. *Appl. Phys. Lett.* **2001**, 79, 3137.
DOI: [10.1063/1.1415353](https://doi.org/10.1063/1.1415353)
- Pandey, S. K.; Kumar, S.; Chatterjee, S. N.; Kumar, U.; Prakash, C.; Chatterjee, R.; Goel, T. C. *Physica B* **2007**, 388, 404.
DOI: [10.1016/j.physb.2006.06.167](https://doi.org/10.1016/j.physb.2006.06.167)
- Mahato, D. K.; Choudhary, R. K.; Srivastava, S. C. *J. Appl. Sci.* **2006**, 6, 716.
- Prakash, C.; Thakur, O. P. *Material Letters* **2003**, 57, 2310.
DOI: [10.1016/S0167-577X\(02\)01217-X](https://doi.org/10.1016/S0167-577X(02)01217-X)
- Pramila, C.; Goel, T. C.; Pillai, P. K. C. *Mater. Sci. Engg. B* **1994**, 26, 25.
SSDI: [0921-5107\(94\)01069-T](https://doi.org/10.1016/S0921-5107(94)01069-T)
- Pandey, S. K.; Thakur, O. P.; Bhattacharya, D. K.; Prakash, C.; Chatterjee, R. *J. Alloys Compd.* **2009**, 468, 356.
DOI: [10.1016/j.jallcom.2008.01.006](https://doi.org/10.1016/j.jallcom.2008.01.006)
- Tripathi, A. K.; Goel, T. C.; Prakash, C. *Mater. Sci. Engg. B* **2002**, 96, 19.
DOI: [10.1016/S0921-5107\(02\)00338-0](https://doi.org/10.1016/S0921-5107(02)00338-0)
- Ranjan, R.; Kumar, R.; Choudhary, R. N. P. *Research Letters in Physics* **2009**, Article ID 382578, 4 pages.
DOI: [10.1155/2009/382578](https://doi.org/10.1155/2009/382578)
- Macdonald, J. R. *Impedance Spectroscopy*, (John Wiley and Sons, **1987**)
- Suchanicz, J. *Mater. Sci. Eng. B.* **1998**, 55, 114.
DOI: [10.1016/S0921-5107\(98\)00188-3](https://doi.org/10.1016/S0921-5107(98)00188-3)
- Plocharski, J.; Wiczoarek, W. *Solid State Ionics* **1988**, 979, 28.
DOI: [10.1016/0167-2738\(88\)90322-0](https://doi.org/10.1016/0167-2738(88)90322-0)
- Suman, C. K.; Prasad, K.; Choudhary, R. N. P. *J. Mater. Sci.* **2006**, 41, 369.
DOI: [10.1007/s10853-005-2620-5](https://doi.org/10.1007/s10853-005-2620-5)
- Rangaraju, M. R.; Choudhary, R. N. P. *J. Mater. Sci.* **2004**, 39, 1765.
DOI: [10.1023/B:JMSS.0000016182.42768.cf](https://doi.org/10.1023/B:JMSS.0000016182.42768.cf)
- Rawat, M.; Yadav, K. L.; Kumar, A.; Patel, P. K.; Adhlakha, N.; Rani, J. *Adv. Mat. Lett.* **2012**, 3(4), 286.
DOI: [10.5185/amlett.2012.2322](https://doi.org/10.5185/amlett.2012.2322)
- Jonscher, A. K. *Nature.* **1977**, 267, 673.
DOI: [10.1038/267673a0](https://doi.org/10.1038/267673a0)
- Behera, A. K.; Mohanty, N. K.; Behera, B.; Nayak, P. *Adv. Mat. Lett.* **2013**, 4(2), 141.
DOI: [10.5185/amlett.2012.6359](https://doi.org/10.5185/amlett.2012.6359)
- Garhardt, R. *J. Phys. Chem. Solids* **1994**, 55, 1491.
DOI: [10.1016/0022-3697\(94\)90575-4](https://doi.org/10.1016/0022-3697(94)90575-4)
- Sinclair, D. C.; West, A. R. *J. Mater. Sci.* **1994**, 29, 6061.
DOI: [10.1007/BF00354542](https://doi.org/10.1007/BF00354542)
- Nobre, M. A. L.; Lanfredi, S. *J. Phys. Chem. Solids* **2003**, 64, 2457.
DOI: [10.1016/j.jpcs.2003.08.007](https://doi.org/10.1016/j.jpcs.2003.08.007)
- Macedo, P. B.; Moynihan, C. T.; Bose, R. *Phys. Chem. Glasses* **1972**, 13, 171.
DOI: [10.1021/j100599a016](https://doi.org/10.1021/j100599a016)
- Borsa, F.; Torgeson, D. R.; Martin, S. W.; Patel, H. K. *Phys. Rev. B* **1992**, 46, 795.
DOI: [10.1103/PhysRevB.46.795](https://doi.org/10.1103/PhysRevB.46.795)
- Victor, P.; Bhattacharyya, S.; Krupanidhi, S. B. *J. Appl. Phys.* **2003**, 94, 5135.
DOI: [10.1063/1.1606509](https://doi.org/10.1063/1.1606509)

Advanced Materials Letters

Publish your article in this journal

ADVANCED MATERIALS Letters is an international journal published quarterly. The journal is intended to provide top-quality peer-reviewed research papers in the fascinating field of materials science particularly in the area of structure, synthesis and processing, characterization, advanced-state properties, and applications of materials. All articles are indexed on various databases including DOI and are available for download for free. The manuscript management system is completely electronic and has fast and fair peer-review process. The journal includes review articles, research articles, notes, letter to editor and short communications.

