

## Lanthanum doped BaTiO<sub>3</sub> ceramics

Korobova N., Soh Deawha\*

\*SMSE, Seoul National University & CPI, Kazakhstan

\*\*Dept. of Electronics Eng., Myongji University, Korea

E-mail : dwhs0h@mju.ac.kr

### Abstract

Sol-gel processing of BaTiO<sub>3</sub> ceramics doped with La(0.01~1.00 at.%) were prepared from metal barium, titanium n-butoxide and lanthanum iso-propoxide. Characterization of the sol-gel-derived powder using XRD, SEM is also reported. The obtained results showed that insulator to semiconductor transition for highly donor-doped barium titanate was closely related to the incorporation of donor into the grains and to the resultant grain size, which were significantly affected by the sinterability of BaTiO<sub>3</sub> powders and sintering conditions used.

Keywords : Sol-gel processing, BaTiO<sub>3</sub> ceramics, donor-doped, lanthanum titanate, semiconductor

### I. Introduction

It is well known that BaTiO<sub>3</sub> can be converted to an n-type semiconductor when it is doped with a small proportion (0.1~0.3 at.%) of trivalent or pentavalent cations, such as La<sup>3+</sup>, Y<sup>3+</sup>, or Nb<sup>5+</sup>, and that semiconducting barium titanate ceramics exhibit an anomalous increase in resistivity above the Curie point (T<sub>c</sub>~120°C)[1]. Among the many methods of the preparation of fine-grained barium titanate or lanthanum titanate, the alkoxide method is known to give a stoichiometric precursor of the oxide[2,3]. Sol-gel processing of oxide powders has gained much interest because of its simplicity, low processing temperature, and chemical homogeneity[4]. This is especially important for applications of the oxides as electronic devices. We chose a reaction between Ti, Ba and La alkoxides to obtain La<sub>x</sub>Ba<sub>1-x</sub>TiO<sub>3</sub> system, expecting better stoichiometric control by replacing the ion by La and/or Ba ions. The primary objective of this work is to determine the relation between

crystallite size and phase at a given temperature as a function of the composition in the lanthanum-barium titanate system for a well-characterized La<sub>x</sub>Ba<sub>1-x</sub>TiO<sub>3</sub> powder.

### II. Experimental procedure

#### 1. Sample preparation

As a first step a fine particles sol was prepared by dissolving barium metal (Furuuchi Chemical Tokyo, Japan) in absolute butanol alcohol in a mixed solution with 2-methoxyethanol (2-MOE, Aldrich Chemical Co) and stoichiometric amount of titanium (IV) butoxide Ti(n-OBu)<sub>4</sub> (Tri-chemical, Yamanashi, Japan) refluxed at 125°C for 4h under a dry argon atmosphere until the barium metal was completely dissolved. Then the solution was cooled to 80°C. The Ba-Ti complex alkoxide used as a starting material for the (100-x)BaO<sub>x</sub>TiO<sub>2</sub> compositions with La<sub>2</sub>O<sub>3</sub>. Lanthanum isopropoxide was diluted with

isopropyl alcohol and added slowly to barium-titanium complex alkoxide. The mixture was stirred and refluxed at 125°C for 1 h. The handling of chemicals and procedures was conducted in a dry argon chamber. The complex alkoxide was diluted in a closed vessel with anhydrous isopropanol in a molar ratio to the alkoxide of 13. Subsequently, water diluted with anhydrous isopropanol was slowly added drop wise with stirring in order to avoid the local formation of precipitates. The total amount of isopropanol in the final solution was in a molar ratio to the alkoxide of 20. The molar ratio of added water to Ti in the alkoxide was 1.5. No base or acid catalyst was used. The solution increased in viscosity as the hydrolysis-condensation reaction preceded, and set to gel after 5h to 7h depending on the alkoxide composition. Then gel dried at 100°C and heated from 200°C to 600°C at a rate of 10°C/min. compared drying methods, such as air, vacuum, freeze drying to show the surface reactivity of synthesized powders (Table 1). Phase transition was measured by XRD. SEM was used to observe the morphology of the powders and the particle size. A few samples were observed by TEM and STEM to confirm that each primary particle was essentially a crystallite so that we could regard the calculated particle size from the BET measurement as the crystallite size.

### III. Results and Discussions

Table 1 shows that the specific surface area is dependent on drying method for the powders. The X-ray data obtained from this study is found in Table 2. All data are recorded for single-phase samples. It has been found that  $\text{La}_x\text{Ba}_{1-x}\text{TiO}_3$  is a simple cubic perovskite throughout most of its composition range. It was found that "cubic" lattice parameter extrapolated to room temperature the thermal expansion data available for temperatures above 120°C.

**Table 1.** BET Values (Surface Area) for  $\text{La}_x\text{Ba}_{1-x}\text{TiO}_3$  powder

Composition $\text{La}_x\text{Ba}_{1-x}\text{TiO}_3$	Surface area (m <sup>2</sup> /g) of dried gel			Surface area of calcined powder
	Air dry	Vacuum dry	Freeze dry	
$\text{La}_{0.1}\text{Ba}_{0.9}\text{TiO}_3$	136.4	151.9	154.6	59.9
$\text{La}_{0.5}\text{Ba}_{0.5}\text{TiO}_3$	142.1	163.2	174.3	48.1
$\text{La}_{0.9}\text{Ba}_{0.1}\text{TiO}_3$	129.6	131.7	144.2	46.4

At both ends,  $x=0$  and  $x=0.9\sim 1.0$  the system becomes non-cubic. At the end where  $x=0$  there is the well-known compound  $\text{BaTiO}_3$  which is tetragonal at room temperature and cubic above 120°C. Apparently 5 % or less La substitution is sufficient to decrease the transformation point to below room temperature.

**Table 2.** Compositions and their lattice parameters

Nominal composition	Lattice parameter	Wt. % $\text{Ti}^{3+}$ Calc.	Wt. % $\text{Ti}^{3+}$ Found
$\text{BaTiO}_3$		0	0.27
$\text{La}_{0.02}\text{Ba}_{0.98}\text{TiO}_3$		0.41	0.90 ± 0.06
$\text{La}_{0.05}\text{Ba}_{0.95}\text{TiO}_3$	3.997	1.03	1.31 ± 0.03
$\text{La}_{0.1}\text{Ba}_{0.9}\text{TiO}_3$	3.990	2.05	2.06 ± 0.02
	3.989		2.08 ± 0.02
	3.978		2.18 ± 0.02
$\text{La}_{0.25}\text{Ba}_{0.75}\text{TiO}_3$	3.968	5.13	4.78 ± 0.06
$\text{La}_{0.5}\text{Ba}_{0.5}\text{TiO}_3$	3.968	10.2	8.9 ± 0.2
	3.958		8.0 ± 0.1
$\text{La}_{0.75}\text{Ba}_{0.25}\text{TiO}_3$	3.960	15.3	13.9 ± 0.06
	3.957		13.6 ± 0.05
$\text{La}_{0.85}\text{Ba}_{0.15}\text{TiO}_3$	3.957		12.6 ± 0.11
$\text{La}_{0.9}\text{Ba}_{0.1}\text{TiO}_3$	3.958	17.4	
$\text{La}_{0.75}\text{TiO}_3$	3.923		14.6 ± 0.05
$\text{LaTiO}_3$	3.895		

A preparation of  $\text{La}_{0.9}\text{Ba}_{0.1}\text{TiO}_3$  has similar symmetry and almost an identical parameter indicating that the symmetry change occurs between  $x=0.85$  and  $x=0.90$ . The resolution of lines in  $\text{LaTiO}_3$  is not felt sufficient to discriminate between the orthorhombic a and b axes. It might be pointed out that orthorhombic symmetry for  $\text{LaTiO}_3$  has been suggested by Roth[5] on the basis of ionic radii. Figure 1

shows the as-sintered micro-structures of specimens having different La-doping contents.

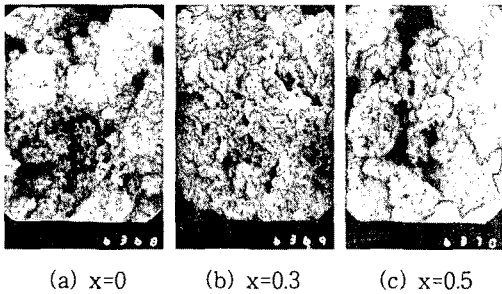


Fig. 1. SEM microstructure of  $Ba_{1-x}La_xTiO_3$  : calcined at  $600^\circ C$  for 2h after freeze drying.

It was found, that powders prepared by air-drying had the strongest agglomerate and freeze-dried powders had the weakest agglomerates. A change of grain size with increasing doping content can be clearly observed. Of particular significance is the observation by Ward and Cotton[6] that  $LaTiO_3$  is the only instance in which perovskite has been made having La defects such that the value  $z$  is 0.25 and 0.2 in the formula  $La_{1-z}Ti_{1-3z}Ti_{3z}O_3$ . The perovskite lattice parameter increased linearly with the lanthanum content and reported a single phase region existing from  $La_{0.75}TiO_3$  to  $LaTiO_3$ . The increase in lattice parameter with  $La^{3+}$  is what would be expected since  $La^{3+}$  is filling vacancies and small  $Ti^{4+}$  ions are being converted to larger  $Ti^{3+}$  ions. The composition  $La_{0.75}TiO_3$  had approximately the same lattice constant as for  $LaTiO_3$  which might suggest the possibility that the products were to a certain extent La deficient. One of the most useful tools for examining the electrical properties of this system is the thermoelectric power measurement. The measurement is not highly sensitive to inevitable variations in sample density and the possibility of small amounts of surface oxidation. The thermoelectric power,  $\alpha$ , might be naively expected to be related to:  $k/e \cdot \ln(x/(x-1))$  resulting in high negative values of  $\alpha$  at low values of  $x$ ,  $\alpha=0$  at  $x=0.5$ , and high positive values of  $\alpha$  at high

values of  $x$ . The experimental results are shown in Fig. 2.

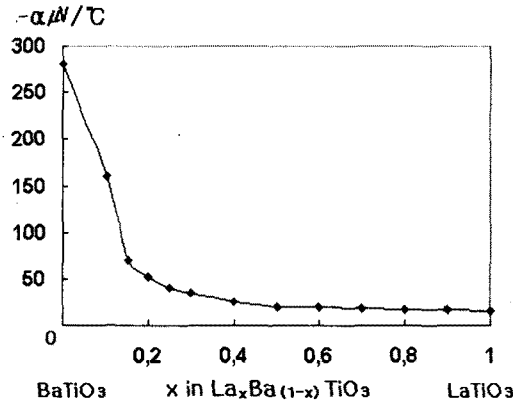


Fig. 2. Thermoelectric power vs. composition.

At small lanthanum contents the thermoelectric power is negative as a result of the formation of  $Ti^{3+}$ , which acts as an electronic conductor. The thermoelectric power also decreases with  $x$  as would be expected. Beyond the vicinity  $x=0.5$  the thermoelectric power does not change sign and does not increase to large positive values with  $x$ . Such experimental results are contrary to what would be expected if the material were to behave as a hole conductor on the basis of the presence of a relatively small amount of  $Ti^{4+}$  in a lattice made up of predominantly  $Ti^{3+}$ . Instead, the thermoelectric power remains as a low negative value suggesting that  $LaTiO_3$  and compositions nearby are metallic in their electrical behavior. This may be interpreted in terms of unpublished data of Mazelsky who studied a number of mixed valence perovskites. Mazelsky found almost without exception that when the transition metal of the perovskite existed primarily in a very stable valence state (e.g.,  $BaTi^{4+}O_3$ ,  $LaFe^{3+}O_3$ ,  $LaCr^{3+}O_3$ ,  $KNb^{5+}O_3$ ) the material was an insulator and when doped became a normal semiconductor. However, when the transition metal existed in an unstable valence state (which are incidentally frequently hard to prepare) (e.g.,  $LaTi^{3+}O_3$ ,  $LaMn^{3+}O_3$ ,  $SrNb^{4+}O_3$ ) the materials

exhibited either metallic behavior or rapidly went intrinsic. This is no doubt due to the ease of the thermally excited reaction of the unstable transition metal ion to the more stable valence state. (e.g.,  $Ti^{3+} \rightarrow Ti^{4+} + e$ ). According to the formula  $La_x^{3+}Ba_{1-x}^{2+}Ti_{1-x}^{4+}O_3^{2-}$  these materials contain reduced titanium in proportion to the amount of lanthanum added. The instability of  $Ti^{3+}$  ion is the reason for prior investigations being limited to very low lanthanum additions and is what necessitates the precautions cited in the preparation procedure.  $La_{0.1}Ba_{0.9}TiO_3$ , fired in the graphite protection tube have values of the thermoelectric power of  $-165 \mu V/^\circ C$  as can be seen on Fig. 2. If the graphite is omitted, the thermoelectric power rises to  $-470 \mu V/^\circ C$  since the number of charge carriers (which may be equated with  $Ti^{3+}$ ) are decreased. Even at room temperature oxidation is a problem. Finely powdered samples were observed to turn from pure black to gray color in a matter of days. Analytical chemical results support the oxidation in these cases. Despite all precautions it will be noted from the tables that practically all samples appear to be somewhat oxidized according to the  $Ti^{3+}$  analysis. Oxidation of the sample as it is ground to a powder immediately prior to analysis, oxidation during storage and handling, and oxidation during the preparation probably all contribute to these results. In general, only  $Ti^{3+}$  was determined. By assuming a formula of the type  $La_{1-y}^{3+}Ti_{1-3y}^{3+}Ti_{3y}^{4+}O_3^{2-}$ , it is possible to calculate the value x on the basis of either La or  $Ti^{3+}$  content.

#### IV. Conclusions

In the present study, the incorporation of La in  $BaTiO_3$  has been studied with higher concentrations. The present method has shown the possibility of preparing submicrometer barium-lanthanum titanate system by simple sol-gel technique. The following conclusions can be drawn from this study:

(1) stoichiometric  $LaTiO_3$  is very probably

orthorhombic; (2)  $LaTiO_3$  with lanthanum defects is very close to simple cubic perovskite with the lattice parameter increasing with the lanthanum content, the presence of additional phases possibly depending on the conditions of preparation, and (3) lanthanum may be lost in the course of  $LaTiO_3$  preparations in some unknown manner; (4) firing time is not a critical determination of particle size: through particles have a slight tendency to grow over longer holding periods.

#### Acknowledgements

This work was supported by the KISTEP grant of M6-0011-00-0043 for int'l. JRP.

#### References

- [1] Gallagher P.K., Schrey F., DiMarcell F.V., Preparation of semiconducting titanates by chemical methods, J. Am. Ceram. Soc., Vol. 46, No. 8, pp. 359, 1963.
- [2] Kuwabara M., Effect of microstructure on the PTCR effect in semiconducting barium titanate ceramics, J. Am. Ceram. Soc., Vol. 64, No. 11, pp. 639, 1981.
- [3] Phule P.P., Risbud S.H., Sol-gel synthesis of barium titanate powders using barium acetate and titanium (IV) isopropoxide. Adv. Ceram. Mater., Vol. 3, No. 2, pp. 183, 1988.
- [4] Korobova N. The metal alkoxides as source materials in the sol-gel technique; Book2 Barium titanate sol-gel materials: ceramics and thin films, Almaty, p. 110, 1998.
- [5] Roth R.S., J. Res. National Bureau of Standards, Vol. 88, p. 75, 1987.
- [6] Ward R. and Cotton F.A., Progress in Inorganic Chemistry. Interscience, New York, 1989.