

## Decizie de indexare a faptei de plagiat la poziția 00385 / 03.07.2017 și pentru admitere la publicare în volum tipărit

care se bazează pe:

**A. Nota de constatare și confirmare a indiciilor de plagiat prin fișa suspiciunii inclusă în decizie.**

Fișa suspiciunii de plagiat / Sheet of plagiarism's suspicion	
Opera suspicionată (OS)	Opera autentică (OA)
Suspicious work	Authentic work
OS	Ioachim Andrei, Toacșan Mariana Irina, Banciu Marin Gabriel, Nedelcu Liviu, Dutu Constantin Augustin, Alexandru V. Horia, Antohe Stefan, Andronescu E, Jinga Sorin, Niță Petru. Synthesis and properties of Ba (Zn 1/3 Ta 2/3) O3 for microwave and millimeter wave applications. <i>Thin Solid Films</i> . 2008 Feb 15;516(7):1558-1562.  <b>Note:</b> a) the source of suspicion: <a href="http://www.integru.org">www.integru.org</a> . b) This work was supported by Ministry of Education and Research under contract CEEX No. 4 / 2005. c) The papers was RETRACTED by the editor.
OA	Ioachim A, Toacșan M.I, Banciu M.G, Nedelcu L, Duțu C.A, Feder M, Plăpcianu C, Lifei F, and Niță P. Effect of the sintering temperature on the Ba (Zn 1/3 Ta 2/3) O3 dielectric properties. <i>Journal of the European Ceramic Society</i> . 2007 Dec 31;27(2):1117-1122. (Available on-line in 12 June 2006).  <b>Note:</b> This work was supported by Ministry of Education and Research under contract nr: 09N/505.2003.
Incidența minimă a suspiciunii / Minimum incidence of suspicion	
p.1559:02s – p.1559: 35s	p.1117:10d – p.1118:23s
p.1559:37s – p.1559:08d	p.1118:26s – p.1118:35s
p.1559:Fig.2	p.1118:Fig.1
p.1559:09d – p.1560:14s	p.1118:37s – p.1118:25d
p.1560:19d – p.1560:31s	p.1119:21s – p.1119:32s
p.1560:04d – p.1560:14s	p.1119:33s – p.1119:45s
p.1560:21d – p.1561:03s	p.1119:34s – p.1119:03d
p.1561:10s – p.1561:21s	p.1120:08d – p.1120:19d
p.1561:22s – p.1561:23d	p.1120:21d – p.1120:44d
p.1561:Fig.7	p.1121:Fig.9
p.1561:Fig.6	p.1121:Fig.7
p.1562:34d – p.1561:38d; p.1562:10s – 1562:03d	p.1121:06d – p.1122:05s
Fișa întocmită pentru includerea suspiciunii în Indexul Operelor Plagiate în România de la Sheet drawn up for including the suspicion in the Index of Plagiarized Works in Romania at <a href="http://www.plagiate.ro">www.plagiate.ro</a>	

**Notă:** Prin „p.72:00” se înțelege paragraful care se termină la finele pag.72. Notăția „p.00:00” semnifică până la ultima pagină a capitolului curent, în întregime de la punctul inițial al preluării.

**Note:** By „p.72:00” one understands the text ending with the end of the page 72. By „p.00:00” one understands the taking over from the initial point till the last page of the current chapter, entirely.

**B. Fișa de argumentare a calificării de plagiat alăturată, fișă care la rândul său este parte a deciziei.**

Echipa Indexului Operelor Plagiate în România

## Fișa de argumentare a calificării

Nr. crt.	Descrierea situației care este încadrată drept plagiat	Se confirmă
1.	Preluarea identică a unor pasaje (piese de creație de tip text) dintr-o operă autentică publicată, fără precizarea întinderii și menționarea provenienței și însușirea acestora într-o lucrare ulterioară celei autentice.	✓
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### Notă:

a) Prin „proveniență” se înțelege informația din care se pot identifica cel puțin numele autorului / autorilor, titlul operei, anul apariției.

b) Plagiul este definit prin textul legii<sup>1</sup>.

„...plagiul – expunerea într-o operă scrisă sau o comunicare orală, inclusiv în format electronic, a unor texte, idei, demonstrații, date, ipoteze, teorii, rezultate ori metode științifice extrase din opere scrise, inclusiv în format electronic, ale altor autori, fără a menționa acest lucru și fără a face trimitere la operele originale...”.

Tehnic, plagiatul are la bază conceptul de **piesă de creație** care<sup>2</sup>:

„...este un element de comunicare prezentat în formă scrisă, ca text, imagine sau combinat, care posedă un subiect, o organizare sau o construcție logică și de argumentare care presupune niște premise, un raționament și o concluzie. Piesa de creație presupune în mod necesar o formă de exprimare specifică unei persoane. Piesa de creație se poate asocia cu întreaga operă autentică sau cu o parte a acesteia...”

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- Opera autentică a fost făcută publică anterior operei suspicioase.
- Cele două opere conțin piese de creație identificabile comune care posedă, fiecare în parte, un subiect și o formă de prezentare bine definită.
- Pentru piesele de creație comune, adică prezente în opera autentică și în opera suspicioasă, nu există o menționare explicită a provenienței. Menționarea provenienței se face printr-o citare care permite identificarea piesei de creație preluate din opera autentică.
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- În opera suspicioasă se identifică un fir sau mai multe fire logice de argumentare și tratare care leagă aceleași premise cu aceleași concluzii ca în opera autentică...”

<sup>1</sup> Legea nr. 206/2004 privind buna conduită în cercetarea științifică, dezvoltarea tehnologică și inovare, publicată în Monitorul Oficial al României, Partea I, nr. 505 din 4 iunie 2004

<sup>2</sup> ISOC, D. Ghid de acțiune împotriva plagiatului: bună-conduită, prevenire, combatere. Cluj-Napoca: Ecou Transilvan, 2012.

<sup>3</sup> ISOC, D. Prevenitor de plagiat. Cluj-Napoca: Ecou Transilvan, 2014.

# Synthesis and properties of $\text{Ba}(\text{Zn}_{1/3}\text{Ta}_{2/3})\text{O}_3$ for microwave and millimeter wave applications



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## Abstract

High dielectric materials have gained an important position in microwave electronics by reducing the size and cost of components for a wide range of applications from mobile telephony to spatial communications.  $\text{Ba}(\text{M}_{1/3}\text{Ta}_{2/3})\text{O}_3$  (BZT) is an  $\text{A}(\text{B}'\text{B}'')\text{O}_3$  type perovskite material, showing ultra high values of the quality factor  $Q$ . Ceramic-based BZT dielectric materials were prepared by solid state reaction. The samples were sintered at temperatures in the range  $1400 \div 1600$  °C for 4 h. Compositional, structural and morphological characterization were performed by using XRD, SEM and EDX analysis. The dielectric properties were measured in the microwave range ( $6 \div 7$  GHz). An additional annealing at 1400 °C for 10 h has improved some dielectric parameters. For samples sintered at temperatures higher than 1500 °C, the permittivity values were obtained in the interval  $30 \div 35$  and almost do not change the value after the annealing. The  $Q \times f$  product substantially increases up to about 135,000 GHz, exhibiting a low temperature coefficient of the resonant frequency ( $\tau_f$ ) in microwaves. The best parameters were obtained for the samples sintered at 1600 °C with additional annealing. The achieved high values of the  $Q \times f$  product recommend these materials for microwave and millimeter wave applications.

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**Keywords:** Tantalates; Thermal treatment; Microstructure; Microwave dielectric properties; Very high quality factor

## 1. Introduction

Complex perovskites  $\text{Ba}(\text{M}_{1/3}\text{B}''_{2/3})\text{O}_3$  are very promising materials for electronic applications owing to their attractive properties, especially the ultra high values of the quality factor  $Q$ .  $\text{Ba}(\text{Zn}_{1/3}\text{Ta}_{2/3})\text{O}_3$  compound (BZT) has potential for applications as satellite broadcasting at frequencies higher than 10 GHz and as very high  $Q$  dielectric resonators (DR) in mobile phone base stations or combiner filter for PCS applications [1]. The BZT ceramics exhibit a dielectric constant of 29, a  $Q \times f$  product of  $100,000 \div 160,000$  GHz and a low temperature coefficient of resonant frequency  $\tau_f$  close to zero in the presence of such additives as strontium–gallium [2].

The long range ordering (LRO) of cations, zinc oxide evaporation, point defects and stabilization of microdomain bound-

aries have been considered to be the factors which influence the  $Q$  values of such perovskites as  $\text{Ba}(\text{M}_{1/3}^{2+}\text{Ta}_{2/3}^{5+})\text{O}_3$  with  $\text{M}=\text{Mg}, \text{Zn}$  [2,3]. The high  $Q$  values were explained from the point of view of the lattice vibrations of its trigonal superstructure. Sagala and Namb [4] calculated the dielectric loss tangent at microwave frequencies from the equation of ion motions, which was a function of B-site ordering. Galasso and co-workers [5] concluded that the B-site ordering increased as the difference in charge and size between  $\text{B}'$  and  $\text{B}''$  atoms increased. The ordering of complex perovskite BZT is important because the 1:2 ordering of the  $\text{B}'$  and  $\text{B}''$  cations along the  $\langle 111 \rangle$  axis is believed to be closely related to the high- $Q$  property of BZT. There is a strong correlation between the cation ordering degree, domain growth, zinc loss and sintering parameters. The synthesis of a BZT material, its structural and morphologic characteristics as well as its microwave dielectric properties correlated to the thermal treatments parameters are reported.

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## 2. Experimental

Ceramics with molar formula  $\text{Ba}(\text{Zn}_{1/3}\text{Ta}_{2/3})\text{O}_3$  were prepared by solid-state reaction. The starting materials were  $\text{BaCO}_3$ ,  $\text{ZnO}$  and  $\text{Ta}_2\text{O}_5$ . Stoichiometric quantities were weighted, ground, homogenized and milled in an agate mill in water for 2 h. The powders were calcined at  $T=925^\circ\text{C}$  for 2 h. Then the powders were milled for 2 h and calcined at  $1000^\circ\text{C}/2\text{ h}$ . The triple calcined powders were mixed with 1% polyvinyl alcohol (PVA) and dried at  $T=80^\circ\text{C}$ , then were pressed into cylindrical samples of 12 mm diameter and 10 mm height. The pellets were slowly dried at  $80^\circ\text{C}$  in order to eliminate the PVA. The third calcination was carried out at  $T=1150^\circ\text{C}/2.5\text{ h}$ , in air.

The density of green ceramics was  $\rho=4400\text{ kg/m}^3$ . The sintering treatment for the BZT samples was performed in air for 4 h at five temperature values:  $1400^\circ\text{C}$ ,  $1450^\circ\text{C}$ ,  $1500^\circ\text{C}$ ,  $1525^\circ\text{C}$  and  $1600^\circ\text{C}$ . In order to increase the dielectric properties, especially the quality factor  $Q$ , the samples were treated supplementary for 10 h at  $1400^\circ\text{C}$ . The pellets were polished in order to remove the superficial zone and to obtain correct values of the microwave dielectric parameters.

The relative bulk density of sintered disks was measured by the Archimedes method. Morphological, structural and compositional analyses were performed on 5 sets of samples by X-ray diffraction (XRD) analysis and electron microscopy (SEM, EDX). The patterns were recorded in a  $2\theta$  range from  $20^\circ$  to  $60^\circ$  on a Seifert Debye Flex 2002 diffractometer into the  $2\theta$  mode. Measurements were performed at room temperature using  $\text{Cu K}\alpha$  radiation, Ni filter and a detector scanned of  $7^\circ/\text{min}$ .

The dielectric parameters, i.e. the dielectric constant  $\epsilon'$ , the quality factor  $Q$  and the loss tangent  $\tan\delta$  were measured in the microwave range by Hakki–Coleman method. A computer-aided measurement system containing an HP 857 C scalar network analyzer and an HP 8350 sweep oscillator was employed.

## 3. Results and discussion

The bulk density of sintered BZT ceramics was measured after grinding and polishing. The temperature dependence of densification after 4 h sintering in air is shown in Fig. 1.

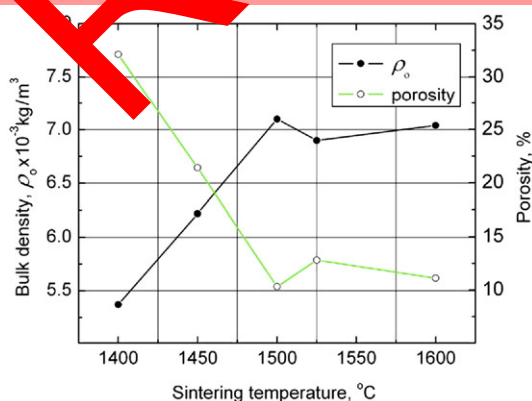


Fig. 1. Bulk density versus sintering temperature for BZT samples.

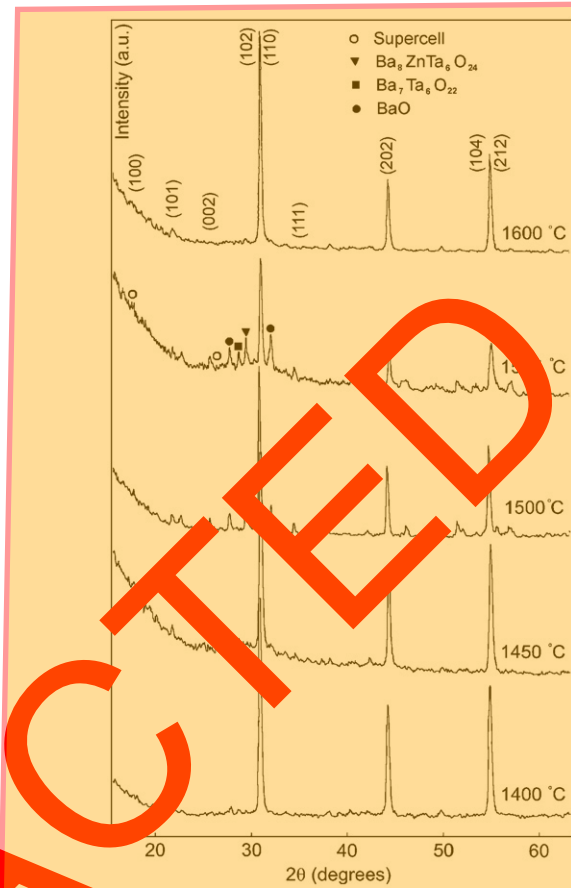


Fig. 2. XRD patterns of  $\text{Ba}(\text{Zn}_{1/3}\text{Ta}_{2/3})\text{O}_3$  system versus sintering temperature.

BZT ceramics sintered between  $1500^\circ\text{C}$  and  $1600^\circ\text{C}$  were well sintered.

The X-ray density of  $\text{Ba}(\text{Zn}_{1/3}\text{Ta}_{2/3})\text{O}_3$  compound was considered as  $\rho_t=7920\text{ kg/m}^3$  [6]. The most dense ceramics exhibit a porosity value of 10%. An abnormal grain growth together with an accentuated  $\text{ZnO}$  evaporation occurred above the sintering temperature  $T_s=1525^\circ\text{C}$ , so the bulk density slightly decreased. Above  $T_s=1525^\circ\text{C}$ , the bulk density continued to increase.

The crystal structure of  $\text{Ba}(\text{Zn}_{1/3}\text{Ta}_{2/3})\text{O}_3$  is the structure of a classic ordered perovskite with two transition metals ( $\text{B}'$  and  $\text{B}''$ ) on the octahedral site. The  $\text{B}'$  and  $\text{B}''$  ions occur on alternating (111) planes in a 1:2 ratio. Each (111) plane contains only one type of small oxygen octahedron, and the planes are ordered in the 1:2 stoichiometric ratio, in the sequence  $\text{Zn-Ta-Ta-Zn-Ta-Ta} \dots$ . The resulting symmetry, due to the (111) type plane stacking and accompanying small oxygen displacement strongly, depends on processing parameters. Consequently, the ordering of Zn and Ta depends of firing temperature and of the sintering time. The BZT samples sintered at low temperatures ( $T_s < 1300^\circ\text{C}$ ) exhibit a pseudocubic perovskite type structure where Zn and Ta cations are in disorder. At higher sintering temperatures ( $T_s > 1300^\circ\text{C}$ ), BZT presents a hexagonal structure, with Zn and Ta ions showing 1:2 order in the B site. The order of Zn and Ta cations expands the original perovskite unit cell along the  $\langle 111 \rangle$  axis and contracts the unit cell in the (111) plane. Consequently,  $c/a$



ratio has a value greater than  $\sqrt{3}/2 \approx 1.2247$  and the unit cell is distorted.

The increase of cation ordering on B site in BZT compounds creates a superlattice with a trigonal supercell. The degree of long range-structural order (LRO) can be evaluated by X-ray methods and it can be expressed as the ratio between the intensity of the super structure reflection peak and that of a basic unit cell reflection peak. The X-ray diffraction patterns for BZT compound, sintered at five sintering temperatures, using Cu K $\alpha$  radiation are given in Fig. 2. The patterns confirm the formation of the hexagonal structure, which is the majority phase. For the BZT compounds sintered at  $T_s = 1525$  °C the patterns reveal the presence of secondary phases Ba $_7$ Ta $_6$ O $_{22}$ , Ba $_8$ ZnTa $_{24}$  and BaO, which disappear at higher sintering temperatures. The superlattice peaks at  $2\theta = 17.6^\circ$  and  $26.4^\circ$  increase in intensity with the increase of  $T_s$  up to 1525 °C. At  $T_s = 1600$  °C they practically disappear. For temperatures up to 1525 °C, the LRO gradually increases with the temperature increase.

The perfectly ordered structure is produced by the following process: (a) the perovskite-type structure is formed, but Zn and Ta cations are in disorder; (b) Zn and Ta cations are partially in order; and (c) Zn and Ta cations are completely in order and the lattice is distorted.

At high sintering temperatures, ZnO evaporates, thus a supplementary distortion can be produced by the ZnO loss. The Zn $^{2+}$  ions on B'-site are partially replaced by Ba $^{2+}$ , which has a much larger ionic radius of 0.161 nm comparatively to 0.074 nm

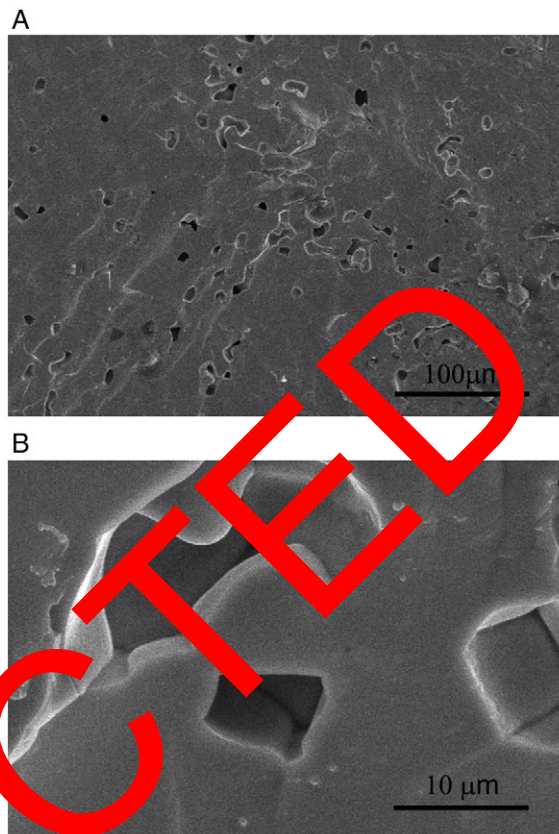


Fig. 4. SEM micrographs of BZT sample sintered in air for 4h at 1600 °C: (A) general view; (B) detail.

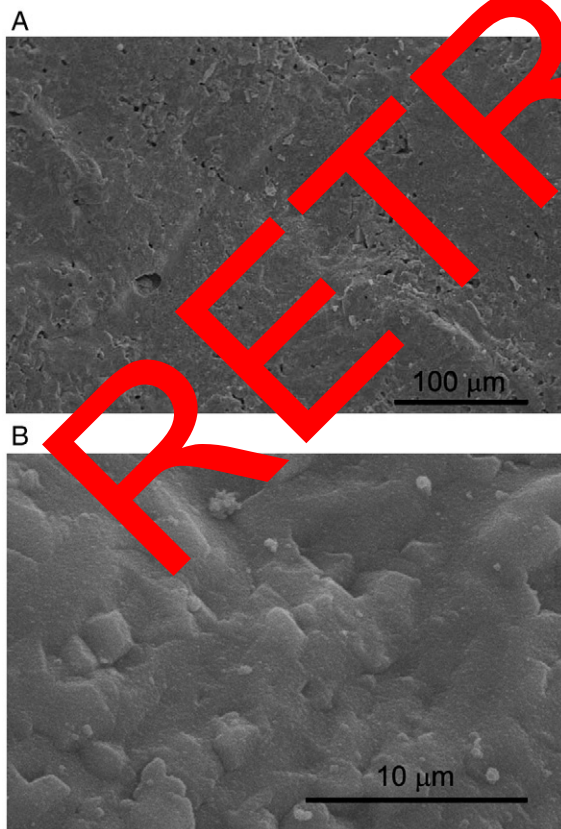


Fig. 3. Micrographs of the BZT sample sintered 1500 °C/4 h: (A) general view; (B) detail.

for Zn cations. This substitution creates a supplementary unit cell distortion, even when the LRO is saturated and leads to an increase of  $Q$ .

The microstructure of BZT ceramics sintered in air at 1500 °C/4h and 1600 °C/4 h was investigated by using SEM. The images are presented in Figs. 3 and 4. The BZT samples sintered at  $T_s > 1500$  °C present grains with concentric ordered shells. BZT samples have a large porous structure due to ZnO evaporation during the sintering process. In Fig. 3, micrographs of the sample sintered in air at 1500 °C/4 h are presented. The SEM image in Fig. 3A reveals an extended inter-aggregates porosity, with pore size in the range (10–30) μm. In addition, few small pores with sizes between 3 and 5 μm are located between grains. Submicron pores appear only on the grain surfaces. The large grains of a polyhedral shape, well faceted and with sizes in the range (50–100) μm, are present in all the samples. Micron grains are located on the grain boundaries or on the grain surfaces; some grains with size in the range 3–6 μm are polyhedral with rounded corners and not well faceted and others are spherical with sizes up to 1.5 μm.

The SEM image of the central zone of BZT sample sintered at 1600 °C for 4 h is shown in Fig. 4. In this case, the porous structure is reduced comparatively with the samples sintered at 1500 °C. The large pores disappear. There are inter-aggregates pores with sizes between 5 and 10 μm and smaller ones with sizes (1–3) μm located between three or four grains.

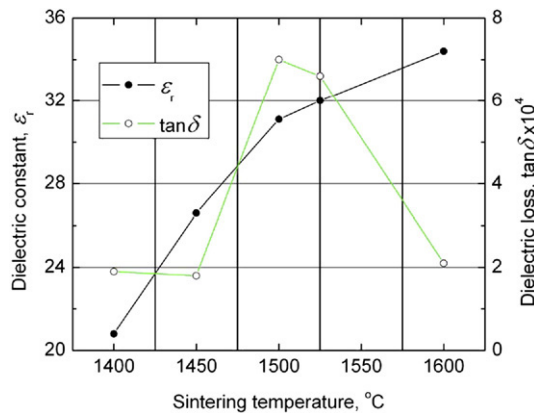


Fig. 5. Dielectric properties versus sintering temperature for BZT samples without annealing.

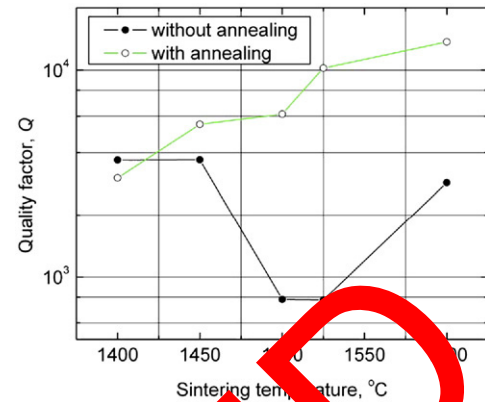


Fig. 7. Quality factor normalized at 10 GHz versus sintering temperature for BZT samples with and without annealing.

The micrograph presents a bimodal distribution of the grains size. Well-faceted polyhedral grains with smooth surfaces and edges and dimensions in the range (30 ÷ 80)  $\mu\text{m}$  are present. The abnormal granular growth together with the ZnO evaporation can explain the non-monotonic variation of the BZT bulk density with  $T_s$ , for  $T_s > 1500$  °C. The relatively small difference between 1500 °C and 1600 °C sintering temperatures has as effect a strong granular growth, as can be noticed in Figs. 3B and 4B.

We used microanalysis (EDX) in an attempt to examine whether a composition trend such as a “paucity” of the Zn element could explain the correlation between the structural long range ordering and dielectric properties. The BZT cylinders sintered at 1600 °C exhibit two different zones: a central axial light-colored zone and a peripheral dark-colored zone. In the outer zone, where the ZnO evaporation is more possible, the Zn atomic concentration indicates a non-stoichiometric composition. On the other hand, the experimental data indicate a variation of  $\text{Ba}^{2+}$  content with higher concentration at the sample boundary. In this case,  $\text{Ba}^{2+}$  can enter on the Zn-octahedron sites, producing a supplementary deformation of the unit cell.

Microwave measurements on dielectric constant and loss tangent were carried out on the BZT samples. The obtained data

revealed a dominant influence of the sintering temperature on the complex dielectric constant. The variation of the dielectric constant and dielectric loss with the sintering temperature can be observed in Fig. 5. This can be considered as an effect of the reduced porosity resulting in a better densification at high sintering temperatures, as can be seen in Fig. 1.

The loss tangent exhibits values around  $2 \times 10^{-4}$  for samples sintered at 1400 °C and 1450 °C. With the sintering temperature increase, the dielectric loss tangent increases up to  $7 \times 10^{-4}$  for  $T_s = 1500$  °C, then decreases down to  $2 \times 10^{-4}$  for  $T_s = 1600$  °C, as can be noticed in Fig. 5. This variation could be related to the Zn and Ta cation ordering process and with the secondary phases, which appear for sintering temperatures between 1450 °C and 1525 °C as shown in Fig. 2. For temperatures higher than 1525 °C, these phases disappear and, consequently, the dielectric loss became lower. The additional thermal treatment at 1400 °C for 10 h for all the samples resulted in the reduction of the BZT dielectric loss as can be seen in Fig. 6. For annealed samples, the dielectric loss continuously decreases with the increase of the sintering temperature. The dielectric constant values were not substantially modified by the annealing treatment, with only one exception of the BZT samples sintered at 1600 °C.

The quality factor  $Q$  values normalized at 10 GHz for the BZT samples are presented in Fig. 7. The  $Q$  is higher for annealed samples than for the samples without thermal treatment; this difference is more significant especially for BZT samples sintered beyond 1525 °C, which indicate the necessity of post sintering thermal treatment. The BZT samples sintered at 1600 °C exhibit a dielectric constant of about 31, a temperature coefficient of resonance frequency  $\tau_f$  between 3 and 7 ppm/°C and a quality factor normalized to 10 GHz up to 13,500.

#### 4. Conclusions

$\text{Ba}(\text{Zn}_{1/3}\text{Ta}_{2/3})\text{O}_3$  ceramics with low loss in microwave domain were obtained by solid state reaction in the temperature range 1400 ÷ 1600 °C. The increase of the sintering temperature leads to a normal granular growth, with a polyhedral grains size development.

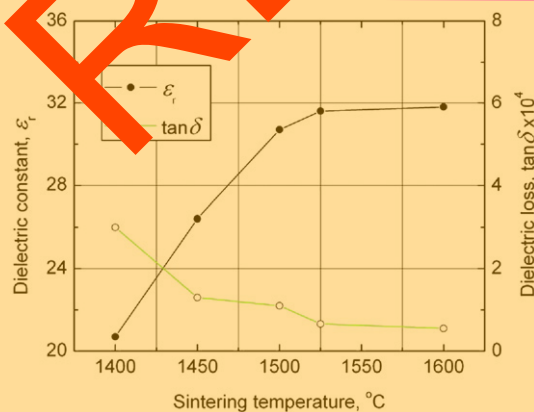


Fig. 6. Dielectric properties versus sintering temperature for annealed BZT samples.

For sintering temperatures higher than 1500 °C, the XRD patterns reveal the BZT multiphase compositions with the presence of trigonal supercell peaks and low Zn content secondary phases. The last two disappear at temperatures higher than 1600 °C.

The investigations on dielectric properties revealed that the increase of the sintering temperature decreases the dielectric loss especially when an annealing of 10 h at 1400 °C is performed.

The quality factor  $Q$  strongly depends on the BZT crystalline structure, that is on the unit cell distortion and cationic order. Dielectric loss decreases with the increase of ordering degree in the structure and with the disappearance of secondary phases. Lowest loss is obtained for a Zn and Ta completely ordered BZT ceramic with a strongly distorted unit cell. Porosity has small effect on dielectric loss of BZT material.

Well-sintered and annealed BZT samples exhibit a dielectric constant around 31, a  $Q \times f$  product up to 135,000 GHz and a temperature coefficient of resonance frequency  $\tau_f$  between 3 and 7 ppm/°C, similar to literature data. The achieved very high

quality factor  $Q$  and small  $\tau_f$  recommend the BZT materials for such microwave and millimeter wave applications as resonators and filters for wireless communication systems.

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