Phase evolution and microwave dielectric properties of ceramics with nominal composition Li$_{2x}$(Zn$_{0.95}$Co$_{0.05}$)$_{2-x}$SiO$_4$ for LTCC applications

Xiangyu Du,$^a$ Hua Su,$^{ab}$ Huaiwu Zhang,$^a$ Xiuting Liu$^a$ and Xiaoli Tang$^a$

Li$_{2x}$(Zn$_{0.95}$Co$_{0.05}$)$_{2-x}$SiO$_4$ (0 $\leq$ x $\leq$ 1) ceramics were prepared through the conventional solid-state route. A fixed amount of Li$_2$O–MgO–ZnO–B$_2$O$_3$–SiO$_2$ (LMZBS) glass was used as a sintering aid to help lower the sintering temperature to around 900 °C. The effects of part lithium-ion substitution on the phase formation, sintering behaviour, microstructures and microwave dielectric properties of the ceramics were systematically investigated. When x = 0.25, the sample achieved a dense microstructure and exhibited excellent microwave dielectric properties of $\varepsilon_r$ = 6.47, $Q$ × $f$ = 131 579 GHz and $\tau_f$ = -27.12 ppm °C$^{-1}$. For its practical application in low-temperature co-fired ceramics (LTCC), large positive $\tau_f$ of CaTiO$_3$ was used to adjust the $\tau_f$ value of the composite ceramic to nearly zero. The composite ceramic of 0.97Li$_2$O·(Zn$_{0.95}$Co$_{0.05}$)$_{2.5}$SiO$_4$·0.025CaTiO$_3$ sintered at 900 °C also presented good microwave dielectric properties of $\varepsilon_r$ = 6.773, $Q$ × $f$ = 69 177 GHz and $\tau_f$ = -2.45 ppm °C$^{-1}$.

1. Introduction

Microwave dielectric materials are widely investigated for their extensive applications as filters, duplexer and antennas in microwave communication industries.$^{1,2}$ Materials with low permittivity ($\varepsilon_r$), high quality factor ($Q$ × $f$) and near-zero temperature coefficient of resonant frequency ($\tau_f$) are required to reduce cross-talk, propagation delay time, noise and power dissipation.$^{3-5}$ Furthermore, these materials should be sintered at temperatures lower than 950 °C or even 900 °C to be used in LTCC multi-layer devices for the miniaturisation and integration of microwave components because of silver (Ag) electrode (around 961 °C), which is typically used as metallic electrode in LTCC materials.$^{5,7}$ Although a number of materials (such as Al$_2$O$_3$ (ref. 4), MgTiO$_3$ (ref. 8) and Mg$_2$SiO$_4$ (ref. 3)) have low permittivity and dielectric loss, these materials cannot be used in LTCC multi-layer devices because of the high sintering temperature.$^8$ Zn$_2$SiO$_4$ is one of these materials and has a low dielectric constant (6.6), a high quality factor value (219 000 GHz) and a temperature coefficient of resonant frequency of -61 ppm °C$^{-1}$. Analogously, the high sintering temperature of 1340 °C limits the application of Zn$_2$SiO$_4$ in LTCC technology.$^{10}$

Recently, Chen reported that the co-substitution (Zn$_{1-x}$Co$_x$)$_2$SiO$_4$ ceramics lowers the sintering temperature of Zn$_2$SiO$_4$ from 1340 °C to 1300 °C and restrains the formation of ZnO phase.$^{11}$ Zhou lowered the sintering temperature of (Zn$_{0.95}$Co$_{0.05}$)$_2$SiO$_4$ ceramics to 900 °C by adding 1.5 wt% LBBS glass. The ceramics sintered at 900 °C showed $\varepsilon_r$ = 6.16, $Q$ × $f$ = 33 000 GHz and $\tau_f$ = -59 ppm °C$^{-1}$. Nonetheless, the ceramics have a large negative $\tau_f$ value, which restricts its potential for practical application. Furthermore, the quality factor is still too low to keep up with the high-speed development of LTCC technology. Although the addition of low softening point glass is the most inexpensive way to lower the sintering temperature of dielectric ceramics, previous studies concluded that a large amount of glass addition leads to either high microwave dielectric loss or crack formation.$^{13}$ Hence, new strategies combined with low amount of sintering aids should be developed to further lower the sintering temperature and dielectric loss of (Zn$_{0.95}$Co$_{0.05}$)$_2$SiO$_4$ ceramics.

The lithium-substitution on Li$_2$CaSiO$_4$,$^{14}$ LiAlSiO$_4$ (ref. 15) and Li$_2$MgSiO$_4$ (ref. 16) ceramics presented good dielectric properties of low dielectric constant (<10) and high $Q$ × $f$ value. However, the phase evolution, sintering behaviour and optimal lithium-substitution content on (Zn$_{0.95}$Co$_{0.05}$)$_2$SiO$_4$ ceramics are not yet reported. Therefore, in this work, the ceramics with nominal composition Li$_{2x}$(Zn$_{0.95}$Co$_{0.05}$)$_{2-x}$SiO$_4$ was synthesised to systematically study the phase formation, sintering behaviour, microstructures and microwave dielectric properties. A fixed amount of LMZBS glass was used as a sintering aid to help lower the sintering temperature at around 900 °C. Moreover, a common modifier with large positive $\tau_f$ of CaTiO$_3$ was used to adjust the $\tau_f$ value of the ceramic, which exhibited the best microwave dielectric property, in the ceramics with nominal composition Li$_{2x}$(Zn$_{0.95}$Co$_{0.05}$)$_{2-x}$SiO$_4$ to nearly zero, thereby
making these ceramics promising candidates for LTCC materials.

2. Experimental

2.1. Preparation of ceramic samples

Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics were prepared through the traditional solid-state ceramic route. The schematic plot of the fabrication process of ceramics is shown in Fig. 1. A previous study reported that the cobalt ions substituted for zinc ions in the lattice sites of ZnO exist as Co$^{2+}$ ions when Co$_2$O$_3$ was used as raw powders.\textsuperscript{17} Hence, high-purity oxides Li$_2$O (99%), ZnO (99%), SiO$_2$ (99%), Co$_2$O$_3$ (99%), CaCO$_3$ (99%) and TiO$_2$ (99%) were used as the raw materials. These raw powders, which were weighed according to respective stoichiometric ratio of Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ (0 ≤ x ≤ 1) and CaTiO$_3$, were ball-milled in nylon jars with zirconia balls in distilled water for 6 h. The resultant slurry was dried, sieved and calcined at 1050 °C for 4 h and 1100 °C for 3 h in air, respectively. A fixed amount of 1.5 wt% LMZBS glass was doped to the pre-sintered Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ (0 ≤ x ≤ 1) powders and then re-milled in distilled water medium for 12 h. With 25 wt% polyvinyl alcohol (PVA) solution as a binder, the powders were dried at 120 °C, well-ground, granulated and pressed into disks under a pressure of 9 MPa and then sintered at 850 °C to 950 °C for 3 h.

After determining the optimal x value to obtain the best microwave dielectric properties of Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics sintered at 900 °C, different amounts of pre-sintered CaTiO$_3$ powders were added to Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ (x was determined) powders with 1.5 wt% LMZBS glass to adjust the $\tau_f$ value of Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ ceramic to nearly zero. The subsequent process was similar to the above-mentioned procedure.

2.2. Preparation of glass

The Li$_2$O–MgO–ZnO–B$_2$O$_3$–SiO$_2$ (LMZBS) glass was synthesized using a quenching method. High-purity grade raw materials (>99%) were mixed, ball-milled and melted at 1350 °C for 1 h using an alumina crucible at the molar ratio of Li$_2$O : MgO : ZnO : B$_2$O$_3$ : SiO$_2$ = 20 : 20 : 20 : 20 : 20. The solution was quickly removed from the furnace and was quenched with cold distilled water to obtain the glass.

2.3. Sample characterization

The bulk densities of these sintered specimens were measured using the Archimedes method. Relative densities were obtained by using the ratio of the bulk and theoretical densities. The phase compositions of these sintered samples were determined by X-ray diffraction (XRD:DX-2700) using Cu K$_\alpha$ radiation. The microstructure of the ceramics was examined by scanning electron microscopy (SEM: Hitachi S-3400N). The microwave dielectric properties of the sintered specimens were investigated using an Agilent N5230A network analyser (300 MHz to 20 GHz) in a resonant cavity. The quality factor and relative permittivity were measured by the resonant-cavity method and the Hakki–Coleman method, respectively.\textsuperscript{18} The temperature coefficient of resonant frequency ($\tau_f$) was measured from the invar cavity using the following formula in the temperature range of 20–80 °C:\textsuperscript{19}

$$\tau_f = \frac{f_f - f_0}{(T - T_0)} \times 10^6 \quad (1)$$

where $f_f$ and $f_0$ are the resonant frequencies at 80 °C and 20 °C, $T$ and $T_0$ are 80 °C and 20 °C, respectively.

3. Results and discussion

Fig. 2a shows the XRD diffraction patterns of Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics doped with 1.5 wt% LMZBS and sintered at 900 °C for 3 h in air. Fig. 2b shows the schematic plot of phase evolution of Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ with various amounts of lithium-ion substitution. Only a rhombohedral structure (Zn,Co)$_2$SiO$_4$ phase (●, PDF #46-1316) appeared in the XRD pattern of the sample with no lithium doped. However, when 0.125 ≤ x ≤ 0.625, the Li$_{1.6}$Zn$_{1.2}$SiO$_4$ phase (●, PDF #24-0676) co-existed with (Zn,Co)$_2$SiO$_4$ phase in the sintered samples. The peak intensities of Li$_{1.6}$Zn$_{1.2}$SiO$_4$ phase gradually increased with the increasing x value. When 0.125 ≤ x ≤ 0.375, the crystalline phase of (Zn,Co)$_2$SiO$_4$ was the main phase, and the phase of Li$_{1.6}$Zn$_{1.2}$SiO$_4$ was the minor phase. However, both of the crystalline phases became the main phase when x was increased to 0.5. When the x value exceeded 0.625, the phase of (Zn,Co)$_2$SiO$_4$ disappeared (Fig. 2a(g)), and another two crystalline phases of tetragonal structure Li$_2$Zn$_{1.2}$SiO$_4$ (*, PDF #15-0056) and orthorhombic structure Li$_2$CoSiO$_4$ (\(\bigvee\), PDF #24-0611) appeared in the sintered sample when x = 1 (Fig. 2a(h)). Fig. 2 shows that various amounts of lithium-ion substitution induce different phase-formations in Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics.

Fig. 3 illustrates the SEM images of samples doped with 1.5 wt% LMZBS glass and sintered at 900 °C with various x values. Generally, the microstructures present considerable densification that might be due to the sufficient liquid phase, which resulted from the glass addition. In detail, with the increasing x value to 0.25, the pores decreased, and the grain was enlarged (see Fig. 2a–c). Combined with the XRD patterns shown in Fig. 2a, this finding explained that a moderate amount of Li substitution might reduce the sintering temperature of the ceramics with nominal composition Li$_2$Zn$_{0.95}$Co$_{0.05}$SiO$_4$.
via synthesising Li$_{1.6}$Zn$_{1.2}$SiO$_4$. The sintering temperature of (Zn$_{0.95}$Co$_{0.05}$)$_2$SiO$_4$ was 1300°C, which is almost similar to that of Zn$_2$SiO$_4$. It was reported that lithium-ions partly substituted for metal cations significantly reduced the sintering temperature compared with that of initial silicate materials. Hence, the sintering temperature of Li$_{1.6}$Zn$_{1.2}$SiO$_4$ was lower than that of (Zn$_{0.95}$Co$_{0.05}$)$_2$SiO$_4$. However, with the successive addition of lithium, the pores increased (Fig. 3d–f), and a large amount of liquid phase (see Fig. 3f) and evident closed pores in the grains (see Fig. 3e) were observed. These phenomena are attributed to the fast growth of the grains; thus, some pores were trapped in the bodies of the abnormal growth grains. In addition, the liquid phase implied that lithium-ion substitution reduces the sintering temperature and thereby resulted in excessive liquid phase. When $x = 0.75$, a dense microstructure reappeared; however, the abnormal growth grain with closed pores and the liquid phase were still observed in Fig. 3g. When $x$ was increased to 1, the shape of grains became rectangular, and the intergranular pores increased (Fig. 3h). This outcome is attributed to the formation of tetragonal structure of Li$_2$ZnSiO$_4$. The optimal content of Li$^+$ substitution was set at 0.25 to obtain the most compact and uniform microstructure.

Fig. 4 shows the relative density as a function of $x$ value for Li$_2$(Zn$_{0.95}$Co$_{0.05}$)$_{2-x}$SiO$_4$ ($0 \leq x \leq 1$) ceramics doped with 1.5 wt% LMZBS glass and sintered at 850–950°C for 3 h. The relative density was calculated using the formula $\rho_s = \frac{\rho}{\rho_\text{x}}$, where $\rho$ and $\rho_\text{x}$ are the bulk and theoretical density values, respectively. The theoretical density $\rho_\text{x}$ was calculated as follows:

$$\rho_\text{x} = \left( \frac{w_1}{\rho_\text{x1}} + \frac{w_2}{\rho_\text{x2}} + \frac{w_3}{\rho_\text{x3}} \right)$$

where $w_1$, $w_2$, and $w_3$ are the weight percentages of crystalline phases and LMZBS glass in Li$_2$(Zn$_{0.95}$Co$_{0.05}$)$_{2-x}$SiO$_4$ ($0 \leq x \leq 1$).
The variations in the relative density of these samples sintered at 900 °C were significantly lower than at 850 °C and 950 °C. This phenomenon is attributed to two reasons. One reason is the tight relationship between the sintering temperature and relative density. The other reason is that the softening temperature of LMZBS is around 900 °C, which is higher than the sintering temperature of these samples. Furthermore, a lower sintering temperature shifts the obtainable maximum relative density to higher lithium doped content. This outcome suggested that the substitution of lithium ions markedly lowers the sintering temperature of the ceramics with nominal composition Li$_2$ZnSiO$_4$.

Fig. 5 shows the relative permittivity ($\varepsilon_r$) values of the Li$_{1-x}$Zn$_{(0.95-x)0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics sintered at 850–950 °C for 3 h. The variation in relative permittivity of specimens sintered at 850–950 °C is generally in agreement with the trend of the relative density. Densification plays a crucial role in determining the relative permittivity. The $\varepsilon_r$ of Li$_{1-x}$Zn$_{(0.95-x)0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics sintered at 900 °C increased with the increasing x value, reached a maximum value (6.47) at x = 0.25 and then decreased to a minimum value (5.84) at x = 1. The relative permittivity of specimens sintered at 850 °C (6.09 and 5.83, respectively). Combined with the XRD patterns shown in Fig. 2a, this finding is due to the synthesis of Li$_x$ZnSiO$_4$, which exists as the major phase. The $\varepsilon_r$ of Li$_x$ZnSiO$_4$ is 5.8, which is lower than that of (Zn$_{0.95}$Co$_{0.05}$)$_2$SiO$_4$ (6.16).

Fig. 6 shows the variations in $Q \times f$ values of Li$_{1-x}$Zn$_{(0.95-x)0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics with different x contents sintered at 850–950 °C for 3 h. The variations in $Q \times f$ values are generally consistent with the variations in the relative density of Li$_{1-x}$Zn$_{(0.95-x)0.05}$SiO$_4$ (0 ≤ x ≤ 1) ceramics. The $Q \times f$ values were all low when the samples were sintered at 850 °C (less than 49 000 GHz), which is mainly attributed to the low relative densities. When the sintering temperature was increased to 900 °C, the $Q \times f$ values...
increased with the increasing $x$ value and achieved the maximum of 131 579 GHz at $x = 0.25$. This outcome might be due to the high densification (96.46%) of the sintered sample when $x = 0.25$. According to the SEM images shown in Fig. 3c, the large grain resulted in less grain boundary, which indicates less lattice mismatch and lower dielectric loss. The $Q \times f$ values of the samples sintered at 850–950 °C remarkably decreased to the minimal values of around 6800 GHz with the variation in $x$ from 0.75 to 1. Microwave dielectric loss can be divided into intrinsic and extrinsic losses. Intrinsic losses are mainly dependent on the crystal structure and are usually caused by the lattice vibration modes. According to the XRD analysis shown in Fig. 2a, tetragonal structure $\text{Li}_2\text{ZnSiO}_4$ (PDF #15-0056) was the major phase of the sintered sample when $x = 1$. In a previous study, $\text{Li}_2\text{ZnSiO}_4$ ceramics doped with 25 wt% ZB glass sintered at 950 °C exhibited a $Q \times f$ value of 10 800 GHz, which is approximated to the $Q \times f$ value in the present work. Hence, in addition to the influence of extrinsic losses caused by lower densification, the high intrinsic losses of $\text{Li}_2\text{ZnSiO}_4$ thereby induced the dramatic decrease of $Q \times f$ values.

Fig. 7 shows the $\tau_f$ values of the $\text{Li}_{1-x}\text{Zn}_x\text{SiO}_4$ ($0 \leq x \leq 1$) ceramics with different $x$ values doped with 1.5 wt% LMZBS and sintered at 900 °C for 3 h. The $\tau_f$ values initially increased and then decreased with the increasing $x$ content. The temperature coefficient of composite ceramics was obtained from the following logarithmic rule:

$$\tau_f = \nu_1\tau_{f1} + \nu_2\tau_{f2}$$

where $\tau_{f1}$ and $\tau_{f2}$ are the $\tau_f$ values of the ($\text{Zn}_{0.95}\text{Co}_{0.05})_2\text{SiO}_4$ and $\text{Li}_{0.5}\text{Zn}_{1.75}\text{SiO}_4$, respectively. The $\tau_f$ value of the sample, whose main crystalline phase was the phase of $\text{Li}_{1.6}\text{Zn}_{1.2}\text{SiO}_4$ when $x = 0.75$, was $-9.04$ ppm °C$^{-1}$; while the $\tau_f$ value of ($\text{Zn}_{0.95}\text{Co}_{0.05})_2\text{SiO}_4$ was $-51.54$ ppm °C$^{-1}$. Hence, the $\tau_f$ value of $\text{Li}_{1.6}\text{Zn}_{1.2}\text{SiO}_4$ was higher than that of ($\text{Zn}_{0.95}\text{Co}_{0.05})_2\text{SiO}_4$. Combined with the XRD patterns shown in Fig. 2a and function (3), the volume percentage of $\text{Li}_{1.6}\text{Zn}_{1.2}\text{SiO}_4$ gradually increased by increasing $x$ to 0.75, which might be the reason for the increase of the $\tau_f$ values of composite ceramics when $0 \leq x \leq 0.75$. Analogously, when the $\tau_f$ values of $\text{Li}_2\text{ZnSiO}_4$ was $-96.6$ ppm °C$^{-1}$, the $\tau_f$ values of $\text{Li}_{1-x}(	ext{Zn}_{0.95}\text{Co}_{0.05})_2\text{SiO}_4$ ($0.75 \leq x \leq 1$) ceramics decreased with the excessive $x$ content.

Considering the above-mentioned analysis, the optimal $x$ value was set at 0.25 because of the best quality factor of $\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$. One of the most important dielectric properties when considering the practical applications for LTCC is the near-zero temperature coefficient of the resonant frequency ($\tau_f$). Large positive $\tau_f$ of $\text{CaTiO}_3$ (+859 ppm °C$^{-1}$) was used as a modifier and was added into $\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$ powders doped with 1.5 wt% LMZBS to further adjust the $\tau_f$ value ($-27.12$ ppm °C$^{-1}$) to around 0 ppm °C$^{-1}$. The five weight percentages of $\text{CaTiO}_3$ were calculated by using the logarithmic rule, where the $\tau_f$ value of $\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$ was set at $-15$, $-21$, $-27$, $-33$ and $-39$ ppm °C$^{-1}$.

Fig. 8 presents the X-ray diffraction patterns of ($1 - x)\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$ wt% $\text{CaTiO}_3$ ceramics doped with 1.5 wt% LMZBS and sintered at 900 °C for 3 h. All the samples exhibited the mixture of ($\text{Zn}_{0.95}\text{Co}_{0.05})_2\text{SiO}_4$, $\text{CaTiO}_3$, ($\text{Zn}_{0.95}\text{Co}_{0.05})_2\text{SiO}_4$ and $\text{CaTiO}_3$. The CaTiO$_3$ phase co-exists with the $\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$ ceramic. Furthermore, no other new phase was produced during sintering.

Fig. 9 shows the theoretical and measured $\tau_f$ values of ($1 - x)\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$ wt% $\text{CaTiO}_3$ ceramics doped with 1.5 wt% LMZBS and sintered at 900 °C for 3 h. The theoretical $\tau_f$ values of ($1 - x)\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$ wt% $\text{CaTiO}_3$ composite ceramics were calculated using the function (3). As seen in Fig. 9, the theoretical $\tau_f$ value of the sample increased with the increasing $\text{CaTiO}_3$ content, which resulted from the high $\tau_f$ value of $\text{CaTiO}_3$. The change in the measured $\tau_f$ values showed a similar tendency with the theoretical values, which indicated that the addition of $\text{CaTiO}_3$ shifts the $\tau_f$ values of $\text{Li}_{0.5}(	ext{Zn}_{0.95}\text{Co}_{0.05})_1.75\text{SiO}_4$ ceramics to positive values. When the
weight percentage of CaTiO$_3$ increased from 1.79 wt% to 4.53 wt%, the measured $\tau_f$ values increased from $-14.6$ ppm °C$^{-1}$ to 10.74 ppm °C$^{-1}$. A near-zero $\tau_f$ value ($-2.45$ ppm °C$^{-1}$) was obtained when $x = 2.49$ wt% and the ceramic was sintered at 900 °C for 3 h.

Table 1 shows the theoretical and measured microwave dielectric properties of (1 – $x$)Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$-x wt% CaTiO$_3$ ceramics doped with 1.5 wt% LMZBS and sintered at 900 °C for 3 h. The theoretical permittivities of the composite ceramics were calculated using the function below:

$$\log \varepsilon = v_1 \log \varepsilon_1 + v_2 \log \varepsilon_2$$

(4)

where $v_1$ and $v_2$ are the volume fractions and $\varepsilon_1$ and $\varepsilon_2$ are the relative permittivities of the Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$ and CaTiO$_3$ ceramics, respectively. Table 1 shows that the variation of the measured $\varepsilon_r$ of (1 – $x$)Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$-x wt% CaTiO$_3$ composite ceramics agrees well with the theoretical value. Given the relatively high $\varepsilon_r$ of CaTiO$_3$ (12 000 GHz),$^{39}$ although CaTiO$_3$ adjusted the $\tau_f$ values of the Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$ ceramics to nearly zero, the $Q \times f$ values sharply decreased when CaTiO$_3$ was added.

4. Conclusion

The phase evolution and microwave dielectric properties of ceramics with nominal composition Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$ were investigated. Lithium ions were added to partly substitute for the zinc ions, and a fixed amount of 1.5 wt% LMZBS glass was used as a sintering aid to help lower the sintering temperature of Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$ (0 ≤ $x$ ≤ 1) ceramics of around 900 °C. The XRD patterns indicated that various amounts of lithium-ion substitution induce different production patterns of other phases in the Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$ ceramic. A compact and uniform microstructure with few pores, high relative density and $Q \times f$ value was obtained when $x$ was set at 0.25. The sample exhibited excellent microwave dielectric properties of $\varepsilon_r = 6.47$, $Q \times f = 131 579$ GHz and $\tau_f = -27.12$ ppm °C$^{-1}$. In considering the practical application for LTCC, a positive $\tau_f$ of CaTiO$_3$ was used to adjust the $\tau_f$ value of the composite ceramics to nearly zero. The 1.5 wt% LMZBS-doped 0.975 Li$_{0.5}$(Zn$_{0.95}$Co$_{0.05}$)$_{1.75}$SiO$_4$-0.025CaTiO$_3$ (weight ratio) composite ceramics also presented good dielectric properties of $\varepsilon_r = 6.773$, $Q \times f = 69 177$ GHz and $\tau_f = -2.45$ ppm °C$^{-1}$.

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