Optical Properties of SiO₂ – TiO₂ – La₂O₃ – Na₂O – Y₂O₃ Glasses and A Novel Process of Preparing the Parent Glass-Ceramics

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Received: 18 December 2020 / Accepted: 3 February 2021 / Published online: 26 February 2021 © Springer Nature B.V. 2021

Abstract
Quaternary glasses with the composition 50SiO₂ – 25TiO₂ – 5La₂O₃ – (20-x) Na₂O – xY₂O₃ where x: (0 ≤ x ≥ 10) were synthesized using the melt-quench technique. XRD examined the nature of prepared glasses. UV-spectroscopic of investigated glass system studied at room temperature. Both optical bandgap and refractive index of the present glass have been increased. The polarizability and basicity were determined. Thermal parameter values increased as Y₂O₃ increased. Under controlling heat, the glass-ceramic were prepared and confirmed using XRD. Glass-ceramics are examined using SEM to evaluate a microstructure. Ultrasonic velocities and elastic-moduli of glass-ceramic samples are increased because of the increase in internal energy. The role of Y₂O₃ modifier in the glass system is clearly demonstrated. Y₂O₃ also works as an excellent nucleating agent that can induce crystallizations, supporting in the creation of the sub-phase of glass-ceramics.

Keywords Yttrium · UV · Glass-ceramics · XRD · SEM

1 Introduction
Glasses containing transition metals oxides attract the attention of several researchers for excellent infrared transmission compared with the conventional glasses. It makes an ideal candidate for various applications such as infrared transmission components, ultra-fast optical switches, and photonic devices [1–3]. Silicate glasses containing transition metal ions exhibit unique, versatile structural properties in physical and spectral (optical & FTIR) studies, and light activated bioactive glass [4–11]. In recent years, the studies on transition metal ions containing glasses have been increased due to their admirable improvements in semiconducting properties and optoelectronic electronic devices. The abundance of multiple valence states of transition metal ions which arises from unfilled d- orbitals made them a potential candidate for extending their applicability in electrical memory switching, photo-conducting, solid-state batteries, and electronic devices [12]. Transition metal ions containing glass are considered semiconducting substances. Nowadays The glasses are considered potential applicants for electronic, mechanical, and optical [12].

TiO₂ containing glasses show remarkable properties like low phonon frequency, high dielectric, non-linear optical, electric, and magnetic. Under these characteristics glass doped TiO₂ has extend applications in the field of optics, photonics, optoelectronics, and telecommunication devices. In glasses, usually TiO₂, are intermediate, crystallizing agents and it may be observed in the Ti⁴⁺ and involved in the structural units of TiO₄, TiO₆ and TiO₅ [13–18].

The incorporation of Y₂O₃ into sodium-silicate glasses causes the replacement of weak Si-O-Na bonds with strong Si-O-Y bonds lead to enhances the thermal, optical & chemical stability of host glasses. The introduction of Y₂O₃ into the glass network enhanced the optical and physical properties of the glass [19]. Glasses containing rare-earth ions have several optical and photonic applications available [20–28]. It is highly possible for UV optics and solid-state batteries applications because of the good ionic conductivity of these glasses [20–28]. Considering the importance of sodium titanium
Silicate glasses in scientific and technological, characteristics like ionic conductivity in power generation glasses modified with various oxides are strongly required [28, 29]. In contrast, the incorporation of transitional or rare earth oxides into sodium titanium silicate glass structures are enhanced optical, electrical, thermal, mechanical and radiation protection characteristics [30–33]. Glasses containing rare-earth ions have attracted a great deal of interest because of their benefits [34–39]. Intermediate oxides such as Y₂O₃ can act either as a glass modifier or former, depending on their concentration in the glass matrix. Y₂O₃ enhances the host glass matrix’s physical structure and mechanical strength [40–46]. The existence of Y₂O₃ enhances the capacity to form glass and reduces devitrification. The existence of TiO₂ and Y₂O₃ impacts UV-spectroscopic in glass systems. These glasses possess lower photon energy and higher refractive index than other glasses. Scientifically and technologically, the recent innovation of titanium silicate glasses containing Y₂O₃ and La₂O₃ is very significant (Table 1).

In the 1950s, Stookey and Kingery [47] discovered the first glass-ceramic that could be manufactured industrially by adding the nucleating agent TiO₂ to control the devitrification of glass. Even at the end of the nineteenth century, Mc. Millan used crystallization [48]. After Stookey’s discovery of the controlled nucleation and crystallization of glass-ceramics, several studies on sintered glass-ceramics were published [49, 50]. Glass-ceramics are polycrystalline materials formed by heat treatment of glasses of appropriate compositions. Crystallization is an important topic in glass science as well as in glass technology. The crystallization behavior and the final properties of glass ceramics parts are mainly affected by the configuration of the parent glass, the nucleating system, and the crystallization conditions. In comparison to those of the parent glass, the mechanical properties of glass-ceramics are better. Glass-ceramics can also, exert other unusual characteristics that are valuable for purposes, as shown by the incredibly small thermal expansion coefficient, which is therefore acceptable for thermal shock-resistant implementations. The goal of our article is preparing of lanthanum titanium silicate glass structures are enhanced optical, electrical, thermal, mechanical and radiation protection characteristics and TiO₂ with high purity. All chemicals used for the glass preparation obtained from Sigma-Aldrich. The starting materials were mixed by grinding the mixture repeatedly to obtain a fine powder. Firstly, the starting materials have been heated to 450 °C for 4 h to eliminate H₂O, and CO₂. The temperature has been raised to 1200 °C for 30 min. The glasses were annealed at 450 °C for 2 h to relieve the internal stresses and allowed to cool gradually to room temperature. The weight losses were found to be less than 1%.

The amorphous state of the glasses and glass-ceramics were checked using X-ray diffraction. A Philips X-ray diffractometer PW1710 with Ni-filtered Cu-Kα radiation (λ = 1.542 Å) powered at 40 kV and 30 mA was used. UV-spectroscopic of investigated glass system is studied by spectrophotometer (type JASCO V-670). Archimedes principle describes the density as the following relation: \( \rho = \rho_0 \left( \frac{C}{C_0} \right) \), the weight of samples in air and liquid is C and C1 respectively, the density of glass-ceramic sample is \( \rho \), and toluene is \( \rho_0(0.865)\) g/cm³. DTA-50 (type Shimadzu) used for the thermal investigation. By heating the specimen at specified temperatures on two steps, first, at 500 °C, the glass-ceramics are produced, the sconed is 1 h at Tc °C, the crystal growth. A pulse-echo method is used to study mechanical measurements for glass-ceramics by (KARL DEUTSCH Echograph model-1085). The model (JEM-100 CX 11 JAPAN), a scanning microscope (SEM), has been used to examine the morphology of investigated glass-ceramic.

### Table 1 Chemical composition of prepared glasses (mol. %)

| Sample Name | SiO₂ | TiO₂ | Na₂O | La₂O₃ | Y₂O₃ |
|-------------|------|------|------|-------|------|
| G1          | 50   | 25   | 20   | 5     | 0    |
| G2          | 50   | 25   | 18   | 5     | 2    |
| G3          | 50   | 25   | 16   | 5     | 4    |
| G4          | 50   | 25   | 14   | 5     | 6    |
| G5          | 50   | 25   | 12   | 5     | 8    |
| G6          | 50   | 25   | 10   | 5     | 10   |

#### 3 Results and Discussions

#### 3.1 XRD Observations

Figure 1. shows the X-ray results of the studied glasses. These diffractograms show no discrete lines, no sharp peaks, and indicate that glass samples have a high degree of glassy state. The slight shift in the hump at \((15–30)\) 2θ° values with respect to Y₂O₃ concentration can be related to the decrease in the bond length and to the higher coordination number with oxygens.

#### 3.2 UV-Visible Absorption Spectra

UV-Vis-NIR absorption spectroscopy is a completely beneficial approach to characterize the optical of different substances which include thin films, filters, pigments, glass, and...
glass ceramics. Optical absorption spectra of various Y₂O₃ co-doped glasses in wavelength range 200–2700 nm were investigated. Figure 2 exemplifies the absorbance (A) and transmittance (T) of glass samples. Figure 3 exemplifies the reflectance (R) of these glasses. There have indications of increasing the absorption coefficient as Fig. 4. Therefore, Y₂O₃ is accounted as BO development [51]. The absorption coefficient \( \alpha \) was calculated by:

\[
\alpha(\lambda) = \frac{1}{x} \ln \left( \frac{1-R}{T} \right),
\]

Where \( x \) is sample thickness.

3.2.1 Optical Band Gap \( E_{opt} \)

Absorption spectra of glasses in the ultraviolet and visible regions have been used to calculate optical band gap.

Optical band gap is determined by \( (\alpha \cdot h\nu)^{1/2} = B(h\nu - E_{opt}) \). where \( E_{opt} \) is optical band gap, B is an energy independent constant and \( h\nu \) is photon energy. By plotting the \( (\alpha \cdot h\nu)^{1/2} \) versus \( h\nu \) as Fig. 5. The intercept of \( \sqrt{\alpha h\nu} \) versus \( h\nu \) at \( \sqrt{\alpha h\nu} = 0 \) denoted the value of \( E_{opt} \). It was noted with increasing Y₂O₃ content the energy gap increases as in Table 2. This increase can be explained as oxygen bridges (BO) are generated that bind energized electrons more strongly than non-bridging oxygen (NBO). Also, this increasing of \( E_{opt} \) may be due to change in composition of glass matrix and increase the interconnection.

3.2.2 Refractive Index (N)

According to the theory of reflectivity of light, the refractive index \( (n) \) values obtained as a function of the reflectance \( (R) \) and the extinction coefficient \( (k) \) as: \( n = (1-R)^{2+k^2/(1+R)} \)
$2 + k^2$, where $k$ values have been determined using the relation $(k = \alpha \sqrt{4\pi})$. According to the Lorentz–Lorenz equation, the density of the material affects the refractive index in a direct proportion. Thus, the increase in the values of the refractive index is ascribed to the increase in density of the glass. The refractive index of studied glasses shown in Fig. 6. The refractive index of the studied glasses increases with increasing of the wavelength and with increasing of $Y_2O_3$ content.

### 3.2.3 Dispersion Parameters

Molar polarization, and polarizability of glasses have been projected as $R_m = \langle n^2 - 1 \rangle n^2 + 2 \rangle Vm$, $\alpha_m = (34\pi N)R_m$, and $\alpha = \frac{1}{N} \left( \frac{2}{n^2+2} \right) \Sigma_{\text{cat}}$ [52–56] where $Vm$ is molar volume, $n$ is refractive index, $N$ Avogadro number, and $No$ number of oxygen atom. The optical basicity of the samples prepared was linked to polarization; $\Lambda = 1.67 \left( \frac{1}{c^2} \right)$. Figs. 7, 8 and 9 present the Molar Polarizability $\alpha_m$, polarizabilities and optical basicity respectively of prepared samples, it was observed the same trend of refractive index with increasing concentrations of $Y_2O_3$ [53–56].

The molar refractivity depends on $E_{opt}$, $R_m = Vm \left( 1 - \sqrt{E_{opt} / 20} \right)$ and molar polarizability ($\alpha_m$). Reflection loss $R_L = \frac{R_m}{V_m}$. These values of $(R_m)$ ($\alpha_m$) and $(R_L)$ decrease with yttrium because of the decrease in the molar volume are presented in Table 2. The criterion for metallization is predicted as $M = 1 - \frac{R_m}{V_m}$, the metallization value rises with $Y^+3$. The electronegativity ($\chi$) is predicted as $\chi = 0.2688E_{opt}$, where $E_{opt}$ bandgap. Thus, with $Y^+3$ increasing, the electronegativity ($\chi$) values increase. The electron polarizability is predicted as $\alpha = -0.9 \chi + 3.5$ and optical basicity $\Lambda = -0.5 \chi + 1.7$. $\alpha$ and $\Lambda$ have the inverse trend of ($\chi$) thus, with $Y^+3$ increase.

### Table 2 Various physical parameters of the studied glasses

| Samples | G 1 | G 2 | G 3 | G 4 | G 5 | G 6 |
|---------|-----|-----|-----|-----|-----|-----|
| Number of oxygen atom | 1.85 | 1.89 | 1.93 | 1.97 | 2.01 | 2.05 |
| Molar Refractivity $R_m$ (cm$^3$/mol) | 14 | 13.16 | 13.32 | 12.97 | 12.5 | 12.05 |
| Molar Polarizability $\alpha_m$ (A$^3$) | 5.55 | 5.22 | 5.2812 | 5.14 | 4.96 | 4.78 |
| Metallization criterion (M) | 0.36 | 0.362 | 0.365 | 0.367 | 0.37 | 0.374 |
| Reflection loss ($R_L$) | 0.639 | 0.637 | 0.635 | 0.633 | 0.63 | 0.626 |
| Electronegativity ($\chi$) | 0.699 | 0.71 | 0.72 | 0.725 | 0.74 | 0.753 |
| Electron Polarizability ($\alpha^o$) | 2.87 | 2.863 | 2.85 | 2.847 | 2.84 | 2.82 |
| Optical basicity ($) | 1.35 | 1.346 | 1.34 | 1.34 | 1.332 | 1.32 |
| Indirect Optical band gap (eV) | 2.601 | 2.635 | 2.669 | 2.699 | 2.74 | 2.8 |
| Ionicity ($I_o$) | 0.58 | 0.578 | 0.576 | 0.574 | 0.571 | 0.569 |
| Covalency ($I_c$) | 0.42 | 0.422 | 0.424 | 0.426 | 0.429 | 0.431 |

![Fig. 5](image-url) Plot of $(\alpha h\nu)^{1/2}$ and $(\alpha h\nu)^2$ against photon energy ($h\nu$) to calculate the optical band gap from the intercept of the curves.

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α° and Λ decrease. This explanation is linked to the value of the optical basicity of Y₂O₃ (0.99) and Na₂O s (1.15) \[55–58\]. Table 2 shows the variation of these values.

### 3.3 Thermal Analysis

DTA-thermograms are the best way to show thermal properties. The first thermal attribute is the transition temperature of samples Tg, while the following thermal property consists of Tc andTp crystallization and peak temperatures. DTA-thermograms of glasses are presented in Fig. 10. It is observed that these parameters have been increased as the Y₂O₃ increases. This is related to increase in the connection of the glass structure. According to these parameters, the glass-ceramics have been prepared. The estimated thermal stability values as: \(\Delta T = (Tc - Tg)\), \(Hg = \frac{\Delta T}{Tg}\), and \(S = (Tp - Tc) \frac{\Delta T}{Tg}\) are listed in Table 3.

It indicated that, the increasing with increasing of Y₂O₃ content is due to the transformation of Si–O–Na into Si–O–Y, and Na–O bond strength is (20 KCal/mol) is much lower than Y–O (50KCal/mol) \[59\]. This behavior is linked to the modification of the coordination number with increasing Y₂O₃, the growth in average constant force and cross-link density. As Y₂O₃ increases at expense of Na₂O, the molar volume decreases and the density increases, making the glass structure more compact.

### 3.4 XRD and SEM Analysis for Consistent Ceramic-Glass

Glass-ceramics have been investigated using XRD for further analysis, as seen in the Fig. 11. The XRD pattern of G1 and G6 are similar, as noticeable from the Fig. 11, excluding that diffraction peaks are stronger. High transparency, uniform color, and good chemical and mechanical properties were...
exhibited due to an increasing the content of $Y_2O_3$. The network of sodium silicate units could be broken by $Y_2O_3$ and act as a glass stabilizer, thereby increasing chemical structure and mechanical hardness. Because the transformation of $Si-O-Na$ into $Si-O-Y$, and $Na-O$ bond strength is (20KCal/mol) is much lower than $Y-O$ (50KCal/mol).

The strongest phases are Lanthanum Titanium Silicate, $La_2(Ti_2SiO_9)$, card No. 01–082-1490, Lanthanum Titanium Silicate Oxide, $La_4Ti(Si_2O_7)2$ (Ti$_3$O$_{16}$) card No. 01–079-2299 and less strongly phase Lanthanum Titanium Silicate Oxide $La_4Ti_5(Si_2O_7)2O_8$ card No. 01–083-023. New phases

### Table 3  Thermal parameters of the studied glasses (°C)

| Sample name | $T_g$ | $T_c$ | $T_p$ | $\Delta T$ | $H_g$ | $(T_p-T_c)$ | $S$ |
|-------------|-------|-------|-------|-----------|-------|-------------|----|
| G 1         | 604   | 690   | 768   | 86        | 0.14  | 78          | 11.11 |
| G 2         | 606   | 701   | 780   | 95        | 0.157 | 79          | 12.38 |
| G 3         | 608   | 706   | 785   | 98        | 0.16  | 79          | 12.73 |
| G 4         | 631   | 731   | 815   | 100       | 0.16  | 84          | 13.31 |
| G 5         | 653   | 780   | 866   | 127       | 0.19  | 86          | 16.73 |
| G 6         | 693   | 806   | 901   | 113       | 0.16  | 95          | 15.5 |

![Fig. 10 The DTA curves of the prepared glasses](image-url)
have emerged through the increase in yttrium oxide Lanthanum Yttrium Titanium Oxide La0.5Y0.5TiO3 card No. 01-073-0070 and Yttrium Titanium Silicon Oxide (Trimounsite-(Y)) Y2Ti2SiO9 card No. 00-046-1375. The microcracks had been easily obtained during the recrystallization process based on differences in the coefficients of thermal expansion between the phases of yttrium and the impeccable standards phase, helping to make the glass-ceramic much harder.

Glass-ceramics had been examined using SEM, as seen in Figs. 12a and b to evaluate a microstructure. The size of the crystal is listed in Tables 4 and 5. SEM photograph of selected glass-ceramic samples. It is indicated that uniform distributions in the glass composition. Figure 12 show the micrographically (G1). It is indicated a nearly unchanging on the surface. Lanthanum Titanium Silicate was crystallized in occasional crystalline texture that includes comparatively large interstitial pores reflecting the remaining glassy matrix. Figure 13 illustrates the micrographically (G6). It is indicated that uniform distributions in the glass composition [60]. With the increase of Y2O3 content, the possibility of crystallization is increased, and a glass sub-phase has been created to increase the internal energy.

3.5 Mechanical Properties of Glass-Ceramics

Figures 14 and Tables 6 represented sound velocities of glass-ceramics. This found that Y2O3 increases those velocities. Due to increase in density, bonding strength, and cross-link density. The increase in sound velocities because the increase in the internal energy.

To determine elastic-moduli as, 
\[ L = \rho v_l^2, \quad G = \rho v_s^2, \]
\[ Y = (1 + \sigma)2G, \quad and, \quad K = L - \left(\frac{Y}{2}\right)G. \]
Elastic-moduli of glass-ceramic have been estimated and represented in Fig. 15 and Table 6. It indicated that, these moduli increasing with the Y2O3 increase. Tis due to transformation of Si–O–Na into Si–O–Y, and Na–O bond strength is (20 KCal/mol) is much lower than B–O (50 KCal/mol).

Poisson’s ratio is projected as \( \sigma = \frac{1}{2} \left(\frac{1}{\frac{4\pi}{1+\sigma}}\right) \). Microhardness is projected as \( H = (1-2\sigma) \frac{\rho v_l^2}{\rho v_s^2 - \rho v_l^2} \). Thermal Expansion \( (\alpha_P) \), \( \alpha_P = 23.2 \times 10^{-6} \) and acoustic impedance \( (Z) \). All these parameters are shown in Table 6 and they increased as the yttrium content increase because of the role of a Y2O3 modifier in the glass system is clearly demonstrated.

Table 4 Peak list and grain-size of the glass-ceramics (G1)

| Pos. [2θ°] | Height [cts] | FWHM °2Th. | d-spacing [Å] | Rel. Int. [%] | Tip width [2θ°] | G size nm | Phase and its Crystalline size nm |
|------------|-------------|------------|---------------|--------------|----------------|-----------|---------------------------------|
| 22.6391    | 709.82      | 0.3542     | 3.92773       | 29.53        | 0.3600         | 80.6      | La2(Ti2SiO9),                   |
| 32.0778    | 2403.5      | 0.5314     | 2.79032       | 100.00       | 0.5400         | 54.78     | 70.49888                        |
| 39.5811    | 895.01      | 0.3542     | 2.27695       | 37.24        | 0.3600         | 83.95     | 73.6                             |
| 46.3840    | 532.26      | 0.4133     | 1.95762       | 22.14        | 0.4200         | 75.35     | 60.23398                        |
| 52.1153    | 243.96      | 0.4133     | 1.75502       | 10.15        | 0.4200         | 77.12     |                                 |
| 57.4043    | 695.72      | 0.4133     | 1.60526       | 28.95        | 0.4200         | 77.12     |                                 |
| 66.8322    | 198.36      | 0.4133     | 1.39988       | 8.25         | 0.4200         | 81.1      |                                 |
| 71.6383    | 96.53       | 0.4723     | 1.31733       | 4.02         | 0.4800         | 73.1      |                                 |
| 75.0820    | 38.81       | 0.9446     | 1.26522       | 1.61         | 0.9600         | 37.4      |                                 |
| 76.9051    | 157.72      | 0.5904     | 1.23972       | 6.56         | 0.6000         | 60.5      | La4Ti5(Si2O7)2O8                 |
| 81.2975    | 87.59       | 0.4723     | 1.18347       | 3.64         | 0.4800         | 78.1      | 69.94197                        |
| 85.1379    | 91.56       | 0.7200     | 1.13869       | 3.81         | 0.6000         | 52.76     |                                 |
Table 5  Peak list and grain-size of the glass-ceramics (G 6)

| Pos. [2θ°] | Height [cts] | FWHM [2θ°] | d-spacing [Å] | Rel. Int. [%] | Tip width [2θ°] | G size [nm] | Phase and its Crystalline size [nm] |
|------------|--------------|-------------|---------------|--------------|----------------|------------|-----------------------------------|
| 19.5792    | 76.95        | 0.2952      | 4.53410       | 3.93         | 0.3000         | 96.18      | La₀.₅Y₀.₅TiO₃, (65.91078)          |
| 22.6109    | 481.73       | 0.3542      | 3.93256       | 24.58        | 0.3600         | 80.55      |                                   |
| 25.4780    | 103.27       | 0.3542      | 3.49615       | 5.27         | 0.3600         | 80.98      |                                   |
| 28.7768    | 221.71       | 0.2952      | 3.10243       | 11.31        | 0.3000         | 97.84      | Y₂Ti₂SiO₇, (76.76907)              |
| 30.2113    | 161.59       | 0.2952      | 2.95832       | 8.24         | 0.3000         | 98.2       |                                   |
| 31.1414    | 223.12       | 0.2952      | 2.87206       | 11.38        | 0.3000         | 98.4       |                                   |
| 31.9839    | 1960.04      | 0.5314      | 2.79830       | 100.00       | 0.5400         | 54.77      |                                   |
| 38.6616    | 83.20        | 0.2952      | 2.32896       | 4.25         | 0.3000         | 100        |                                   |
| 39.5320    | 752.34       | 0.3542      | 2.27967       | 38.38        | 0.3600         | 83.9       |                                   |
| 43.9363    | 98.11        | 0.2952      | 2.06083       | 5.01         | 0.3000         | 102.2      |                                   |
| 45.1712    | 139.29       | 0.4723      | 2.00732       | 7.11         | 0.4800         | 64.2       |                                   |
| 46.3327    | 440.51       | 0.3542      | 1.95967       | 22.47        | 0.3600         | 85.92      |                                   |
| 47.3145    | 103.10       | 0.2952      | 1.92127       | 5.26         | 0.3000         | 103.5      |                                   |
| 49.2890    | 44.07        | 0.3542      | 1.84883       | 2.25         | 0.3600         | 86.9       |                                   |
| 52.0561    | 197.19       | 0.3542      | 1.75688       | 10.06        | 0.3600         | 87.9       |                                   |
| 56.6288    | 236.43       | 0.3542      | 1.62539       | 12.06        | 0.3600         | 89.7       |                                   |
| 57.3691    | 576.42       | 0.3542      | 1.60616       | 29.41        | 0.3600         | 90         |                                   |
| 58.5723    | 103.19       | 0.3542      | 1.57600       | 5.26         | 0.3600         | 90.56      |                                   |
| 66.7314    | 175.71       | 0.5314      | 1.40175       | 8.96         | 0.5400         | 63         |                                   |
| 67.6088    | 119.93       | 0.3542      | 1.38568       | 6.12         | 0.3600         | 95         |                                   |
| 71.5406    | 96.67        | 0.3542      | 1.31888       | 4.93         | 0.3600         | 97.4       |                                   |
| 75.0633    | 35.16        | 0.7085      | 1.26549       | 1.79         | 0.7200         | 49.8       |                                   |
| 76.8601    | 138.16       | 0.5314      | 1.24033       | 7.05         | 0.5400         | 67.21      |                                   |
| 81.2243    | 78.94        | 0.5904      | 1.18435       | 4.03         | 0.6000         | 62.4       |                                   |
| 85.1694    | 72.57        | 0.7200      | 1.13835       | 3.70         | 0.6000         | 52.78      |                                   |

Fig. 12  SEM backscattered electron images of the produced glass-ceramics (G 1)
Conclusions

Quaternary glasses with the composition of 50SiO$_2$ – 25TiO$_2$ – 5La$_2$O$_3$ – (20-x) Na$_2$O – xY$_2$O$_3$ have been manufactured using conventional melt-quenching methods. The optical, thermal, crystallization, and mechanical variables have been examined for these glasses. XRD measurements established the amorphous nature of glasses. Optical absorption was quantified to understand the optical characteristics of the prepared glasses. With increasing Y$_2$O$_3$ content the energy gap increases. This grow can be explained as oxygen bridges (BO) are generated that bind energized electrons more strongly than non-bridging oxygen (NBO). Refractive index of investigated glasses are increases as density increase. Molar polarization, polarizability, and optical basicity of these glasses having the

| Sample No. | $v_L$ (m s$^{-1}$) | $v_T$ (m s$^{-1}$) | $\sigma$ (Gpa) | $\alpha_P$ (K m$^{-2}$ s$^{-1}$ K$^{-1}$) |
|------------|-------------------|-------------------|----------------|-----------------------------------|
| G 1        | 5425              | 2885              | 31.703         | 0.303                             |
| G 2        | 5455              | 2900              | 35.4           | 0.303                             |
| G 3        | 5500              | 2920              | 37.18          | 0.304                             |
| G 4        | 5550              | 2935              | 40             | 0.306                             |
| G 5        | 5610              | 2955              | 157.75221      | 0.308                             |
| G 6        | 5655              | 2970              | 46.9           | 0.310                             |

Fig. 13 B SEM backscattered electron images of the produced glass ceramics (G 6)

Fig. 14 Dependence of the longitudinal and shear ultrasonic velocities $v_L$ and $v_T$ of the investigated glass-ceramics against of Y$_2$O$_3$ by mol. %

Table 6 values of sound velocities ($v_L$ and $v_T$), elastic moduli, Poisson’s ratio, micro hardness, acoustic impedance ($Z$) and thermal expansion coefficient ($\alpha_P$), of the studied glass-ceramics
same trend of refractive index with concentration of Y$_2$O$_3$ increased. Metallization of these glasses was enhanced because of increment of TiO$_2$. Thermal stability of these glasses was increased as the Y$_2$O$_3$ increases. These increases related to increase in the connection of the glass structure. Under controlling heat, the glass-ceramic was prepared and investigated by XRD, SEM, and mechanical properties. XRD results showed all the expected phases due to the crystallization process. SEM photograph of selected glass-ceramic samples indicated that uniform distributions in the glass composition. Ultrasonic velocities and elastic-modules of glass-ceramic samples are increased because of increase in internal energy.

Acknowledgments The authors extend their appreciation to the Deanship of Scientific Research at King Khalid University for funding this work through research groups program under grant number R.G.P. 2/93/41.

Author Contributions Kh. S. Shaaban: performing XRD, UV measurements and analysis, Writing-review, writing manuscript, Methodology, Software, and writing – discussion.
A.F. Abd El-Rehim: Writing-review, writing – discussion and editing manuscript.
E. A. Abdel Wahab: Writing-review, writing – discussion and editing manuscript.
M. M. Abou Halaka: Writing-review, writing – discussion and editing manuscript.

Funding Statement There are currently no Funding Sources in the list.

Availability of Data and Material My manuscript and associated personal data will be shared with Research Square for the delivery of the author dashboard.

Declarations

The manuscript has not been published elsewhere and that it has not been submitted simultaneously for publication elsewhere.

Consent to Participate The authors consent to participate.

Consent for Publication The authors consent for publication.

Conflict of Interest The authors declare that they have no conflict of interest.

Declaration of Competing Interest The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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