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Influence of molybdenum oxide on structural, optical and physical properties of oxychloride glasses for nonlinear optical devices

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Abstract
The unconventional Heavy Metal Oxide Glasses (HMOG) are characterized by a low phonon energy, large infrared range transmission, high refractive index and nonlinear optical properties. Ternary glasses have been synthesized and studied in the Sb\textsubscript{2}O\textsubscript{3}–MoO\textsubscript{3}–ZnCl\textsubscript{2} system. Further, the glass formation compositional limits are reported and some glass samples with the formula: (90-x)Sb\textsubscript{2}O\textsubscript{3} -xMoO\textsubscript{3}–10 ZnCl\textsubscript{2} (10\%\leq x<50\%, mole \%) were elaborated. Thermal properties have been measured and indicating that the glass transition temperature decreases with increasing proportions of molybdenum oxide. The evolution of density, microhardness and elastic modulus has been studied as functions of parameter x and Raman spectra measurements have been shown the partial conversion of MoO\textsubscript{6} octahedral units into MoO\textsubscript{4} tetrahedral.

Keywords: Oxychloride glasses; Density; Raman, Elastic properties; Optical properties

1. Introduction
Over the past few years, several researches have been carried out on antimony glasses. Binary and ternary Sb₂O₅-based glasses have been studied, including various oxide [1-6], oxysulphide [7, 8] and oxychloride systems [9, 10]. These glasses are used in nonlinear optical devices particularly in broadband optical amplifiers functioning about 1.5μm [11].

Molybdenum oxide is considered as a glass former that requires the presence of other elements to form a glass. Molybdate glasses are obtained by introducing another network-forming oxide such as Sb₂O₅, P₂O₅, TeO₂ etc. The molybdate glasses are used for high-density memories, light modulation, large area display devices like smart windows and other electrochromic systems [12,31,32]. The different oxidation states of molybdenum are: Mo⁶⁺, Mo⁵⁺, Mo⁴⁺ and Mo³⁺. In the molybdate glasses, different works suggest the existence of molybdenum in the Mo⁶⁺ form which is the most stable [13]. Rada et al [14] suggest that a part of the Mo⁶⁺ ions is converted into Mo⁵⁺ and Mo⁴⁺ ions. This process causes the depolymerisation of host glass network. The molybdenum oxide is formed in the glass by different structures, like structural units of octahedron and tetrahedron units such as MoO₆ and MoO₄ respectively. Krapchanska et al [15] showed a partial conversion of MoO₆ to MoO₄ units in the MoO₃-TiO₂-Bi₂O₃ system. Subcik et al [16] also showed that the incorporation of MoO₃ units in the ZnO-B₂O₃-P₂O₅ glasses revealed the increase of the MoO₆/ MoO₄ ratio.

Sb₂O₃ contributes in the glass network with SbO₃ structural units and it appears tetrahedra with the oxygens to be found at three corners and the lone pair of electrons of antimony Sb³⁺ at the fourth corner localized in the third equatorial direction of the Sb atom. There may be a possibility of existence of antimony ions in the Sb⁵⁺ state and participate in the glass network forming with Sb⁵⁺O₄ structural units may enhance the nonlinear optical properties [11]. In Sb₂O₃ based glasses the addition of transition metal
oxide MoO$_3$ improves functionality of glasses for their desired optical applications. Particularly, MoO$_3$ shows interesting optoelectronic features and can control phase separation in glasses [43].

The aim of this paper is focuses on the elaboration of a new antimony oxychloride glass from the Sb$_2$O$_3$-MoO$_3$- ZnCl$_2$ system. We are interested by molybdenum rates incorporated into the glass, its effect on the different physical properties that have been measured, and especially in the structural study by Raman measurements. Physical properties such as: density, energy dispersive spectroscopy (EDS), differential scanning calorimetry (DSC), optical transmission, microhardness and elastic moduli have also been considered in terms of molybdenum influence.

2. Experimental

In this present study, the starting materials Sb$_2$O$_3$≥99% (Acros), ZnCl$_2$ ≥99.8% (Merck) and MoO$_3$≥ 99% (Acros) are used for the synthesis of the glass samples. Melt-quenching is the process of synthesis. The powders are mixed thoroughly in a mortar with the mean batch weight around 5 g. The mixture is inserted into a silica glass tube, about 10 mm in diameter and then subjected to a flame until a clear liquid obtaining. The bath of fusion is cast in a mould under preheated brass plate around 200 °C. The samples obtained (thickness of a few millimetres) are placed in the furnace at a temperature lower than the temperature of vitreous transition with a slow heating rate. After annealing, the samples were polished for physical measurements.

The glass composition was analyzed by the (EDS) energy dispersive spectroscopy (JSM 6400 Jeol and Oxford link ISIS). With the help of crystalline materials references, experimental error is estimated to be 1% for Sb, Mo and Zn, 2%
for Cl and 5% of oxygen. Density was determined using a helium pycnometer (Micromeritics, AccuPyc 1330) with ±0.001 g/cm³ accuracy.

Thermal properties of the samples were measured by the differential scanning calorimetry (DSC 2010 from TA Instruments) at the heating rate of 10 K/min with the sensitivity better than 0.1 °C. The estimated accuracy is ±2 K for the glass transition temperature Tg, crystallization temperature Tx and ± 1 K for the temperature of the crystallization of peak. Tx-Tg provides the stability against devitrification. The DSC signals showed no crystallization peaks indicating that these glasses are having a very low tendency to crystallization and exhibiting extremely high thermal stability. Thermal stability is the important property of glasses and gives the measure of the degree of disorder of glassy state.

The analysis of the vitreous samples was carried out by the Raman spectroscopy in the range of 1100 cm⁻¹ to 100 cm⁻¹ by Sentra- Bruker. Varian Cary 5 spectrometer was used to record the optical transmission in the UV–visible spectrum between 400 and 800 nm and Bomem Michelson 100 IR spectrometer was employed for the infrared spectrum in between 500 and 4000 cm⁻¹.

The Matuzawa MXT 70 digital microhardness was used with 100 g as load to measure the microhardness values. Ultrasonic measurements were taken by the pulse – echo method using a Panametrics model 5800 pulser/receiver with a quartz transducer. Both X-cut transducer and Y-cut transducer (with 10 MHz frequency) were employed for both longitudinal and shear modes. A Hewlett-Packard model 54502A oscilloscope recorded the pulse transient. Young's modulus E, bulk modulus K, and shear modulus G were measured by the following equations [33]:
\[ E = \frac{\rho V_T^2 (3V_L^2 - 4V_T^2)}{V_L^2 - V_T^2} \quad (1) \]

\[ K = \frac{\rho (3V_L^2 - 4V_T^2)}{3} \quad (2) \]

\[ G = \rho V_T^2 \quad (3) \]

Where \( \rho \) = Density of the glass sample

\( V_T \) = Transversal ultrasonic velocity

\( V_L \) = Longitudinal ultrasonic velocity

The uncertainty is estimated to be about ± 2% for elastic modulus.

3. Results and discussion

3.1. Physical properties

Chemical composition, some experimental properties of SMZ glasses are provided in Table 1. Fig. 1 shows the Oxy-halogenated glass area for ternary system. The glass matrix \( \text{Sb}_2\text{O}_3 \)-MoO\(_3\)-ZnCl\(_2\) was characterized by its extensive glass area and exceptional thermal stability. We have chosen to vary the molybdenum oxide, which is an excellent effect in heating homogenization [17, 18]. It is used for its electrochromic properties [19] and especially for improving the conductivity by the ionic and electronic input [20, 21]. A series of ternary system bulk samples were prepared for the physical measurements and were represented by the lines in Fig. 1 with the formula: \((90-x)\) \( \text{Sb}_2\text{O}_3\)-xMoO\(_3\)-10ZnCl\(_2\) \((10\leq x \leq 50)\) called «SMZ». The molar concentration of MoO\(_3\) is given by the number that follow the letters SMZ.

Glass compositions of the prepared samples were studied by the EDS spectra. Fig. 2 shows the chemical composition spectra of the sample SMZ20. In Table 1 it is clear from the chemical composition of the glasses that the differences between the
nominal and analysed cations compositions remain weak. These results indicate that the losses induced by evaporation during fusion are very limited. Silicon traces lower than 1% are observed in all the samples, due to crucible contamination. The measurement of concentration refers to the three analysed elements. Discrepancies are larger for anions; this can be explained by the limited accuracy of the EDS technique for light elements, particularly for the oxygen. Uncertainty related to electronic scan microscopy results is about 1%.

The density and molar volume of the glass samples decreases linearly with the substitution of the antimony oxide and molybdenum oxide. The density of a solid glass is a function of its composition; in fact, it is directly depends on the atomic weight of each component as well as the volume occupied by these components [22]. This reduction is due to the lower atomic weight of the molybdenum than the antimony.

3.2. Thermal properties

DSC curves for different glasses are shown in Fig. 4. The characteristic temperatures of the glass samples have been measured for the glass series (90-x)Sb₂O₃-xMoO₃-10ZnCl₂. The thermal stability Tx-Tg, exceeds 100 °C for all the glass samples (Table 2). The numerical values obtained from the DSC curves provides the information on stability against devitrification (20 K/min). As shown in Table 2, from the results of DSC, when the concentration of MoO₃ increases from 10 to 50 %, the glass transition temperature decreases linearly (Fig. 5), indicates the decreasing bonding strength of the material. Goumeidane et al. [10] was mentioned that an excess of anionic oxygen which causes the increase of the nonbridging oxygen in the vitreous network. This conclusion could explain the decreased values of transition temperature Tg in the glass system.
3.3. Optical properties

In the infrared spectrum, the glasses can transmit up to 1800 cm\(^{-1}\) (Fig. 6). The spectrum shows a large absorption band around 3300 cm\(^{-1}\), originates from OH groups. A second extrinsic absorption band is observed between 1700 and 1900 cm\(^{-1}\), which limits the effects of the optical transmission of these glasses. It corresponds to the first harmonic of the fundamental of Si-O vibration band at 1984 cm\(^{-1}\) and arises probably from silicate anions from glass crucibles. The multiphonon absorption cannot be identified accurately under these conditions. Generally, due to low phonon energy of the oxyfluoride or oxychloride glasses, when rare earth ions are doped into them, the multiphonon relaxation rates are reduced from emitting levels to lower states, which causes the quantum efficiencies to increase for various photonic applications[44,45]. UV-visible depends on the energy band gap between the valence and conduction electronic layers of material. It has been determined by the Urbach method. The UV–visible spectrum for 50Sb\(_2\)O\(_3\)- 40MoO\(_3\)-10 ZnCl\(_2\) glass sample (2.5 mm in thickness) has been shown in Fig. 7 and the UV–visible transmission edge was around 516 nm. Fig. (7) shows the Urbach plot of 50Sb\(_2\)O\(_3\)- 40MoO\(_3\)-10 ZnCl\(_2\) glass sample. The value of optical band gap obtained from the extrapolation of the curve is 2.39 eV.

3.4. Raman spectra

The Raman spectra (Fig. 8) of the ternary system’s vitreous samples are recorded at ambient temperature in the spectral range 1500 – 50 cm\(^{-1}\). The broad peaks and Boson peak confirm the glassy state below 100 cm\(^{-1}\), which is ubiquitous in the Raman spectra of disordered solids, especially of glasses. The presence of the Boson peak in the Raman spectra affirms the vitreous state of material [6] which has not an equivalent in the crystalline materials. The Raman spectra of SMZ 10 glasses reveal the
presence of the band at 416 cm\(^{-1}\), related to symmetric stretching vibrations of the
pyramids SbO\(_3\) and ZnO\(_4\) units. This band intensity decreases with the increase of
MoO\(_3\) from 10% to 30%. We recorded a small shift in the SMZ 20 and SMZ 30 samples,
associated with the same vibration mode, they are toward 424 cm\(^{-1}\) and 413
cm\(^{-1}\) respectively. Moreover, the shoulder at 183–205 cm\(^{-1}\) is assigned to the vibration
of ZnO\(_6\) units.

In previous papers [34–42], it has been established that MoOn polyhedral like
MoO\(_4\) tetrahedral units and MoO\(_6\) octahedra units gave strong Raman bands. In the
region 700–1000 cm\(^{-1}\), Raman bands corresponding to symmetric stretching modes \((v_1)\)
and asymmetric stretching modes \((v_3)\) are observed. Yet, Raman bands attributed to
symmetric bending modes \((v_2)\) and asymmetric bending modes \((v_4)\) are noticed in the
wavenumber range of 300–520 cm\(^{-1}\) [26]. In figure 8, the spectra’s peak position of
glasses containing molybdate oxide indicates that the strong band at 918-936 cm\(^{-1}\) could
be attributed to the deformed vibrational mode of MoO\(_6\) [24]. Beyond of 20% of MoO\(_3\),
a partial conversion of MoO\(_6\) units to MoO\(_4\) units. This transformation is assisted by
Zn\(^{2+}\) or Zn\(^{3+}\) [23]. We noticed that the intensity of MoO\(_6\) increases, whereas the band of
SbO\(_3\), ZnO\(_4\) decreases may be the presence of Mo\(^{5+}\)O\(_3\)\(^{-}\) complexes in the structural
network of the glass series and generates the depolymerisation of zinc and/or
antimonite chains in the vitreous samples [24]. Additionally, the \(v_1\) and \(v_2\) vibrational
bands of monomeric MoO\(_4\)\(^{-}\) tetrahedral units have also been located at about 862-869
cm\(^{-1}\) and 327-346 cm\(^{-1}\) respectively.

For the ZnO-B\(_2\)O\(_3\)-MoO\(_3\) system, Alexandrov et al [25] deconvolution of the Raman
spectrum in the regions 600–1100 cm\(^{-1}\) into 6 peaks with positions at: 830, 880, 920 and
950 cm\(^{-1}\), showing the presence of Mo\(^{6+}\) ions in MoO\(_4\)\(^{-}\) with more coordination, one
peak at 780 cm\(^{-1}\) with isolated MoO\(_4\) units, and the last band corresponds to the
stretching vibration of BO$_4$ at 1000-1200 cm$^{-1}$. In our case, the spectra are fitted with a sum of Gaussian curves by means of the least square method as shown in Fig. 9, the deconvolution is performed in the same interval. We found three peaks at 780 cm$^{-1}$ assigned to isolated MoO$_4$ units, 860 cm$^{-1}$ and 920 cm$^{-1}$ indicate that Mo$^{6+}$ ions are present in (MoO$_4$)$_2^-$ tetrahedral units for the glass sample SMZ20. The analysis of the spectra of SMZ30, SMZ40 and SMZ50 glasses shows the appearance of a new peak at 820 cm$^{-1}$ in addition to the previous ones (780, 860 and 920 cm$^{-1}$). Probably, this shows a structural change at 20% of molybdenum with the formation of Mo$^{6+}$ ions in different coordination’s, namely low symmetry, highly distorted, or isolated in MoO$_4^{2-}$[26].

Finally, we notice that the Zinc chloride is involved in the vitreous network by ZnCl$_4^{2-}$ tetrahedra and as shown by the Raman analysis, zinc chloride portion becomes an oxide. The peaks at 233 and 275 cm$^{-1}$ are assigned to the symmetric Zn–Cl stretching vibrations in type A and B respectively [27]. The peaks are not resolved in this work because of the small amount of zinc chloride in the series of glasses and the intrinsic peak broadening of the Raman spectrum corresponding to the state of an amorphous solid (glassy state).

3.5. Microhardness

Microhardness ($H_V$) was measured for all the samples, as shown in Fig. 10 the state of the microhardness curve is adjusted by the expansion of the MoO$_3$ content. The evaluation is monotonic for the MoO$_3$/Sb$_2$O$_3$ substitution, it drops from 229 to 172 N/mm$^2$ between 10 and 50 mol %. This evolution was explained by the variation of the bond strength and also the dissociation energy. The microhardness of the samples increases with the molybdenum content in the glasses, corresponding to a consolidation of the glass by an increasing of the Mo-O strength bonds. Furthermore, the
molybdenum enters in the glass network as a glass former and contributes to the formation of rigid structural units as well as strengthening the rigidity of the glass structure.

3.6. Elastic properties

Elastic moduli E, G, K have been calculated for the SMZ glass series. A general observation of these curves shows (Fig.11) that elastic moduli increase with the substitution of Sb$_2$O$_3$ by MoO$_3$. There are several models for the interpretation of elastic moduli of oxide glasses. In literature, few studies referred to elastic moduli of oxychloride based antimonite Iezid et al. [9]. The series of glasses comprises 90% of the oxides and 10% of ZnCl$_2$. This chloride has a low binding energy compared to oxides and we assume that the ZnCl$_2$ effect on the elastic modulus is negligible. So, elastic moduli of oxide glasses apply in a good approximation in our case. The model of Mackashima and Mackanzi [28, 29] explains the Young's modulus by increasing the dissociation energy and the packing density factor following the increase in the content of MoO$_3$.

The limited variation of shear modulus can be explained by the model of Bridge B et al [30]. We assume that the cross-link densities are run by an almost constant recovery among bridging oxygen and nonbridging, which limits the variation of G.

The evaluation of K can be explained by the Bulk compression model [30] in the sense that the insertion of MoO$_3$ increases the stretching bond strength since the dissociation energy of M-O is greater than Sb-O.

4. Conclusion
A ternary glass system is synthesised and the broad vitreous area is observed in the Sb$_2$O$_3$ -MoO$_3$- ZnCl$_2$ system. A glass samples in which Sb$_2$O$_3$ is substituted by MoO$_3$ have been characterized. The differential scanning calorimetry analysis shows that all the compositions have a large thermal stability (T_x-T_g > 100 °C). The IR transmission is spread out from the visible to infrared due to the low phonon energy of the antimonite. The general trend is that MoO$_3$ incorporation leads to the decrease of density from 5.038 to 4.722 g/cm$^3$, glass transition temperature T_g from 279 to 259 °C whereas the microhardness (172 to 229 N/mm$^2$) and the rigidity (elastic moduli) increases. The Raman spectroscopy provides the partial conversion of molybdenum ions from MoO$_6$ octahedral to MoO$_4$ tetrahedral positions from 20% of MoO$_3$. The quality of these high stability glasses makes them a potential application for optoelectronic devices.

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References


Fig. 1. Glass formation in the Sb2O3-MoO3-ZnCl2 system.

Fig. 2. EDS spectra of glass sample SMZ20 oxychloride glass.

Fig. 3. DSC curves for the SMZ glasses (heating rate 20 K/min).

Fig. 4. Evolution of glass transition temperature versus molybdenum oxide concentration in the (90-x)Sb2O3-xMoO3-10ZnCl2 system.

Fig. 5. Infrared transmission of a 50Sb2O3-40MoO3-10 ZnCl2 glass sample, 2.5 mm in thickness.

Fig. 6. UV-Visible transmission of a 50Sb2O3-40MoO3-10 ZnCl2 glass sample, 2.5 mm in thickness.

Fig. 7. Urbach plot of 50Sb2O3-40MoO3-10 ZnCl2 glass sample.

Fig. 8. Raman spectra of SMZ glasses.
Fig. 9. Peak deconvolution of Raman spectra of (90-x) Sb₂O₃-xMoO₃-10 ZnCl₂ (x=20,30,40 and 50) glasses.

Fig. 10. Microhardness versus composition parameter x for SMZ glasses.

Fig. 11. Evolution of elastic moduli versus composition parameter x for SMZ glasses. E is Young’s modulus, K is bulk modulus and G shear modulus.

Table 1. Composition and some experimental properties of SMZ glasses.

<table>
<thead>
<tr>
<th>Glass</th>
<th>Cation %</th>
<th>Anion %</th>
<th>Cation%</th>
<th>Anion%</th>
<th>Hv N/m²</th>
<th>ρ g/cm³</th>
<th>V_m cm³/mol</th>
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<tbody>
<tr>
<td></td>
<td>Sb</td>
<td>Zn</td>
<td>Mo</td>
<td>O</td>
<td>C</td>
<td>Sb</td>
<td>Zn</td>
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<tr>
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<td>9</td>
<td>55</td>
<td>5</td>
<td>.1</td>
<td>9</td>
<td>55</td>
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<tr>
<td>172</td>
<td>5.0</td>
<td>11.03</td>
<td></td>
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<tr>
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<td>5</td>
<td>11.</td>
<td>93</td>
<td>6</td>
<td>.1</td>
<td>9</td>
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<td>194</td>
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<td></td>
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<tr>
<td>SMZ 30</td>
<td>75</td>
<td>6.</td>
<td>18.</td>
<td>93</td>
<td>6.</td>
<td>72.</td>
<td>6.</td>
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<tr>
<td>SMZ 40</td>
<td>66.</td>
<td>6.</td>
<td>26.</td>
<td>93</td>
<td>6.</td>
<td>58.</td>
<td>6.</td>
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<td>SMZ 50</td>
<td>57.</td>
<td>7.</td>
<td>35.</td>
<td>93</td>
<td>6.</td>
<td>55.</td>
<td>8.</td>
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</table>

Table 2. Characteristic temperatures (°C) and thermal stability range of SMZ glasses: Tg for glass transition, Tx for onset of crystallization and Tp for exothermic maximum.

<table>
<thead>
<tr>
<th>Glass</th>
<th>Sb₂O₃</th>
<th>MoO₃</th>
<th>ZnCl₂</th>
<th>Tg</th>
<th>Tx</th>
<th>Tp</th>
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<td>401</td>
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<tr>
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<td>10</td>
<td>274</td>
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<tr>
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<td>10</td>
<td>270</td>
<td>405</td>
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<tr>
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<td>10</td>
<td>259</td>
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</table>
Figure 1

Figure 2
Figure 3

Figure 4
Figure 5

Figure 6
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Figure 9
Figure 10

Figure 11