UNIVERSITI TEKNOLOGI MARA

ELASTIC AND STRUCTURAL PROPERTIES OF V₂O₅-PbO-ZnO AND V₂O₅-TeO₂-Li₂O GLASS SYSTEMS

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Thesis submitted in fulfillment of the requirements for the degree of **Master of Science**

Faculty of Applied Sciences

September 2015

AUTHOR'S DECLARATION

I declare that the work in this thesis/dissertation was carried out in accordance with the regulations of Universiti Teknologi MARA. It is original and is the result of my own work, unless otherwise indicated or acknowledged as referenced work. This thesis has not been submitted to any other academic institution or non-academic institution for any other degree or qualification.

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Thesis/Dissertation	:	Elastic and Structural Properties of V2O5-PbO
Title		ZnO and V ₂ O ₅ -TeO ₂ -Li ₂ O Glass Systems
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ABSTRACT

Ternary $(55-x)V_2O_5-45PbO_{x}ZnO (x = 0.15 mol\%)$ and $35V_2O_5-(65-x)TeO_2-$ (x)Li₂O (x = 10-50 mol%) glass series were prepared by the melt-quenching method. Elastic and structural properties of the glasses were investigated by measuring sound velocity using the pulse-echo-overlap technique and Fourier Transform Infrared (FTIR) spectroscopy, respectively. Properties such as density (ρ), molar volume (V_a), and glass transition temperature (T_g) were also reported. For $(55-x)V_2O_5$ -45PbO-(x)ZnO glass series, results from the study showed that both longitudinal and shear velocities decreased at $x = 5 \mod \%$ with the increase of ZnO concentration. The independent longitudinal and shear moduli, C_L and μ and Young's modulus (Y) also showed decreasing trend at $x = 5 \mod \%$ as the fraction of ZnO increases. FTIR analysis showed an increase in non-bridging oxygen (NBO) as indicated by the increase in intensity of VO₄ assigned peaks at x = 5 mol% while the increase in intensity of VO₅ assigned peaks at x > 5 mol% indicates increasing bridging oxygen (BO). The glass transition temperature (T_g) showed a decrease at x = 0.5 mol% which implies increase in non-bridging oxygen (NBO) while an increase of T_g for x = 10-15mol% indicates the decrease in NBO. Analysis of the experimental results using the bulk compression and ring deformation models showed that the calculated value of K_{bc} was higher than that of K_e but K_{bc}/K_e ratio drops from 2.16 (x = 5 mol%) to 1.94 (x = 10 mol%) indicating that bond bending or ring deformation was slightly reduced with addition of ZnO. For $35V_2O_5$ -(65-x)TeO₂-(x)Li₂O glass series, both longitudinal (v_L) and shear velocity (v_S) showed an increase of 6.57% and 16.52%, respectively, at x = 20 mol%. The initial increase in C_L and μ at x = 20 mol% was suggested to be attributed to the increase in number of bridging oxygen (BO) which caused stiffness and rigidity of the glass network to increase while the decrease in C_L and μ between x = 20 and x = 30 mol% was suggested to be due to the increase in number of nonbridging oxygen (NBO). FTIR analysis showed an increase in intensity of TeO_4 and VO₅ assigned peaks at x = 10-20 mol% which indicates increasing BO while at the same time the decrease in intensity of TeO3 and VO4 assigned peaks indicates decreasing NBO. The increase in intensity of TeO₃ and VO₄ assigned peaks between x = 30-50 mol% which was accompanied by the decrease in intensity of TeO₄ and VO₅ assigned peaks indicate increases in NBO and decrease in BO, respectively. Therefore in the present study, formation of more BO compared to NBO can be the reason for the increase in stiffness above and vice versa. Our suggestion above is further supported by the variation of glass transition temperature (T_g) results. The increase of glass transition temperature (T_g) at x = 10-20 mol% implies decrease in non-bridging oxygen (NBO) while the decrease of T_g for x = 30-50 mol% indicates the increase in NBO. Quantitative analysis based on bulk compression and ring deformation models showed that the K_{bc}/K_e ratio drops from 2.60 (x = 10 mol%) to 1.74 ($x = 20 \mod \%$) which indicates that bond bending or ring deformation was reduced with the increase of Li₂O. Meanwhile, the slight increase in value of K_{bc}/K_e ratio from 1.74 (x = 20 mol%) to 2.06 (x = 30 mol%) before dropping to 1.93 (x = 40mol%) and later increasing slightly to 1.96 (x = 50 mol%) indicates variation in ring deformation.

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