PAPERS (1)

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Poster Presentation

Vol 1, 2021 -143667

SOLID-STATE NMR INVESTIGATION OF Na₅RESi₄O₁₂-based (RE = Sc, Y) GLASS-CERAMICS

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Keywords: Solid-State NMR, Solid electrolytes, Glass-ceramics.

The design of Na-ion all-solid-state batteries requires the optimization of materials to be used as solid electrolytes. Fast ion-conducting glass-ceramics are suitable for this purpose since they present the features of both glass phase and fast ion-conducting crystallites[1]. Glass-ceramics based on superionic crystalline structure Na₅RESi₄O₁₂ (N5, RE = Sc. Y) can be produced by controlled crystallization of sodium silicate glasses with specific compositions [2] and can reach ionic conductivities up to 0.1 S/cm [3]. Their compositions and properties can be isostructurally modified by Si -> P substitution. Precursor glasses of compositions Na_{3.9}RE_{0.6}P_{0.3}Si_{2.7}O₉ (NREPS) and Na_{4.2}RE_{0.6}Si₃O₉ (NRES), (RE = Sc, Y), were synthesized by the melting-quenching technique, and glass-ceramics were prepared according to the crystallization temperatures found by DSC. X-ray diffraction pattern of the glass-ceramics showed that the N5 crystal structure was found for NScS and NScPS glass-ceramics annealed at 905 and 925 °C, respectively. Multinuclear (29Si, 23Na, 31P and 45Sc) MAS NMR was used to monitor the conversion of the glassy precursor phase into the multi-phase glass ceramics at different stages of thermal annealing, confirming the successful substitution of (SiO₄)⁴⁻ by (PO₄)³⁻ ions in the N5 structure. Impedance spectroscopy and static ²³Na experiments were used to investigate ion dynamics of glass-ceramics containing the N5 structure.

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Acknowledgements: Authors acknowledge financial support from CeRTEV (Center for Research, Technology, and Education on Vitreous Materials, FAPESP 2013/00793-6). I.D.A. Silva acknowledges the financial support from FAPESP grant 2017/17800-0.