## **Poster Contributions**



## PXRD and PDF analysis of multifunctional lanthanide nitrilotrismethylphosphonate-based proton conductors

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Metal phosphonates are multifunctional solids which possess tunable properties, such as H-bond networks, while exhibiting high chemical and thermal stability<sup>1</sup>.

Depending on the protonation of the ligand, different crystalline phases can be obtained. Here, we report three different families of proton conductors based on lanthanide nitrilotrismethylphosphonates. Compounds having cationic layers compensate by chloride or sulfate anions were isolated:  $[Ln(H_4NMP)(H_2O)_2]Cl\cdot 2H_2O^2$  and  $Ln(H_5NMP)]\cdot SO_4 \cdot 4H_2O$  [H<sub>6</sub>NMP = nitrilotris(methylphosphonic acid)]. The crystal structure of Gd-(H<sub>3</sub>NMP)]·SO<sub>4</sub>·4H<sub>2</sub>O was solved ab initio from synchrotron powder diffraction data ( $\lambda$ =0.4124 Å, beamline BL04-MSPD ALBA) and refined by the Rietveld method. Chloride containing phases show two irreversible solid state transformations take place: (1) a crystalline-to-crystalline phase transition, {Ln-H<sub>4</sub>NMP  $\rightarrow$  [Ln<sub>2</sub>(H<sub>3</sub>NMP)<sub>2</sub>(H<sub>2</sub>O)<sub>4</sub>]·4.5H<sub>2</sub>O for Ln= La, Pr}, and (2) crystalline-to-amorphous phase transition, {LnH<sub>4</sub>NMP  $\rightarrow$  [Ln(H<sub>3</sub>NMP)]·1.5H<sub>2</sub>O for Ln= Gd - Ho}, both implies the loss of HCl and structural rearrangements of the frameworks. Variations in average and local structure have been monitored by high resolution powder diffraction and PDF analysis, upon exposure the samples at high relative humidity and temperature (95% RH and 80 °C), in order to understand their behavior as proton conductors.

Acknowledgements. The work at UMA was funded by MAT2016-77648-R, MAT2013-41836-R (MINECO, Spain) and FQM-1656 (Junta de Andalucía) research grants and CELLS-ALBA Synchrotron is thanked for the provision of synchrotron X-ray beamtime at BL04-MSPD.

<sup>&</sup>lt;sup>1</sup> P. Ramaswamy, N. E. Wong and G. K. H. Shimizu, *Chem. Soc. Rev.* **2014**, *43*, 5913-5932.

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