

Cite this: *Dalton Trans.*, 2012, **41**, 4045

www.rsc.org/dalton

PAPER

Multifunctional lanthanum tetrakisphosphonates: Flexible, ultramicroporous and proton-conducting hybrid frameworks†

Rosario M. P. Colodrero,^a Pascual Olivera-Pastor,^a Enrique R. Losilla,^a Miguel A. G. Aranda,^a Laura Leon-Reina,^b Maria Papadaki,^c Alistair C. McKinlay,^d Russell E. Morris,^d Konstantinos D. Demadis*^c and Aurelio Cabeza*^a

Received 20th October 2011, Accepted 13th December 2011

DOI: 10.1039/c2dt11992g

A new flexible ultramicroporous solid, $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**), has been crystallized at room temperature using the tetrakisphosphonic acid H_8DTMP , hexamethylenediamine-*N,N,N',N'*-tetrakis(methylenephosphonic acid). Its crystal structure, solved by synchrotron powder X-ray diffraction, is characterised by a 3D pillared open-framework containing 1D channels filled with water. Upon dehydration, a new related crystalline phase, $\text{La}(\text{H}_5\text{DTMP})$ (**2**) is formed. Partial rehydration of **2** led to $\text{La}(\text{H}_5\text{DTMP})\cdot 2\text{H}_2\text{O}$ (**3**). These new phases contain highly corrugated layers showing different degrees of conformational flexibility of the long organic chain. The combination of the structural study and the gas adsorption characterization (N_2 and CO_2) suggests an ultramicroporous flexible framework. NO isotherms are indicative of a strong irreversible adsorption of NO within the pores. Impedance data indicates that **1** is a proton-conductor with a conductivity of $8 \times 10^{-3} \text{ S cm}^{-1}$ at 297 K and 98% of relative humidity, and an activation energy of 0.25 eV.

Introduction

Lanthanide Metal–Organic Frameworks (Ln-MOFs) are receiving growing interest owing to their potential applications in fields such as photoluminescence, magnetism, catalysis and gas adsorption/sensing.¹ The rare-earth ions possess rich coordination chemistry, functionality and, additionally, their properties may be modulated across the lanthanide series, which is key for designing advanced multifunctional solid state compounds. Compared to d-block transition metal-containing MOFs,² Ln-MOFs have been relatively much less studied. Porous frameworks are rather scarce, because they usually present less

oriented metal-linker bonds and the bulky lanthanides usually exhibit high coordination numbers, which result in dense solids.³

For solid state applications, such as solid sensors, light-emitting materials or proton conductors, two- or three-dimensional network compounds are required. Due to the high affinity of the rare-earth ions for oxygen-containing species, carboxylate-based ligands have been often used as the linkers of preference, yielding robust framework architectures. For example, Harbuzaru *et al.*^{1d} have prepared 3D microporous Ln-MOFs, by appropriate selection of the carboxylate ligand under hydrothermal conditions. The products combine a set of well-defined properties in the same crystal, including high gas adsorption, anisotropic photoluminescence and magnetic properties.

Multifunctional phosphonate ligands are alternative organic linkers, leading to a variety of thermally and chemically stable Ln-MOFs architectures.⁴ However, exercising control of the dimensionality and crystal structure of phosphonate-based Ln-MOFs still remains a difficult task. So far, only a few examples of three-dimensional networks have been reported. In these cases, the frameworks are constructed by chains of edge-sharing lanthanide polyhedral connected by phosphonate groups.⁵

Recent progress in design of new polyphosphonic ligands has opened the way to the tailor-made synthesis of hybrid inorganic–organic materials.⁶ Diamino tetrakisphosphonic acids are versatile organic linkers because they may contain diverse functionalities in between the peripheral phosphonate groups and hence new compounds with chemical and structural peculiarities may be prepared. In fact, a number of diamino tetrakisphosphonates with different alkaline or transition metals showing chain, layered or three-dimensional open framework structures have already been

^aDepartamento de Química Inorgánica, Universidad de Málaga, Campus Teatinos s/n, Málaga, 29071, Spain. E-mail: aurelio@uma.es; Tel: +34 952131870 + 34 952131877

^bServicios Centrales de Apoyo a la Investigación, Universidad de Málaga, Campus Teatinos s/n, Málaga, 29071, Spain

^cCrystal Engineering, Growth and Design Laboratory, Department of Chemistry, University of Crete, Voutes Campus, Crete, GR-71003, Greece. E-mail: demadis@chemistry.uoc.gr; Fax: +30 2810 545001; Tel: +30 2810 545051

^dSchool of Chemistry, University of St Andrews, Purdie Building, St Andrews, UK, KY16 9ST

† Electronic supplementary information (ESI) available: Fig. S1–S3: XRPD Rietveld plots for compounds **1**–**3**; Fig. S4: Comparison of the raw powder patterns for $\text{Ln}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (Ln = La, Sm); Fig. S5: framework view for **1** showing the 1D channels; Fig. S6: lattice water molecule chain within the channel of **1**. Tables S1–S3: Bond lengths and H-bonding distances for compounds **1**–**3**. CCDC reference numbers 810315–810317. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c2dt11992g

reported.⁷ Some of these show interesting luminescent properties,^{1,5,8} excellent selective cation sorption and exchange capacity,⁹ and scale/corrosion inhibition.¹⁰ Certain metal diamino tetraphosphonates, for example Ga^{III}-H₅TDTMP (H₈TDTMP = tetramethylenediamine-*N,N,N',N'*-*tetrakis*(methylenephosphonic acid)), have drawn the interest of the medical community because of their ability to suppress metastatic tumours^{11a} and others, such as Yb-EDTMP, were used for bone palliation.^{11b}

Extending previous work on CaH₆DTMP, an adsorbate-responsive material with 2D topology exhibiting a “breathing” phenomenon,¹² herein we report the synthesis and characterization of the flexible multifunctional La(H₅DTMP)·7H₂O [H₈DTMP = hexamethylenediamine-*N,N,N',N'*-*tetrakis*(methylenephosphonic acid)], and its gas adsorption as well as its proton conductivity properties.

Experimental

Materials

All starting compounds were from commercial sources and used as received, without further treatment. Deionized (DI) water was used for all syntheses and procedures. La₂O₃ (99.5 %), was from Aldrich, USA and SmCl₂·6H₂O (99+ %) was from Alfa-Aesar, USA. K₆(H₂DTMP) (Dequest 2054, 33.5 % stock solution) was from ThermPhos-Dequest, Belgium.

Preparation of lanthanide tetraphosphonate products

La(H₅DTMP)·7H₂O (1). 1 mL of a 33.5% stock solution of K₆(H₂DTMP) (0.58 mmol) was added to 20 mL of DI water. The pH was then adjusted to ~ 2 by using consecutively 5 M and 1 M HCl solutions. Separately, a quantity of La₂O₃ (0.127 g, 0.39 mmol) was suspended in 30 mL of DI water and then dissolved by consecutive addition of 5 M and 1 M HCl solutions, adjusting the pH to about 0.9. The two solutions were then mixed and the final pH was adjusted to 1.2 with a 1 M NaOH stock solution. The resulting clear colourless solution was left undisturbed for 2 months, after which time a microcrystalline white solid appeared. Alternatively, solid H₈DTMP (Dequest 2051) can be used instead of the K₆(H₂DTMP) stock solution. Anal. Calcd (%) for LaP₄N₂O₁₉C₁₀H₃₉: 15.92%C, 5.21%H, 3.71%N. Found: 15.07%C, 4.92%H, 3.62%N. IR data/cm⁻¹: 3429 (br), 3013 (br), 2965 (br), 2779 (w), 2628 (sh), 2366 (w), 2346 (w), 1632 (m), 1482 (m), 1459 (sh), 1438 (m), 1380 (m), 1342 (w), 1297 (m), 1224 (s), 1190 (s), 1162 (s), 1101 (s), 914 (s), 846 (w), 825 (w), 802 (w), 775 (m).

Sm(H₅DTMP)·7H₂O. The same procedure was used as for La(H₅DTMP)·7H₂O (1), except that the metal source was SmCl₂·6H₂O and the final pH was 0.76.

By following the same procedure as above for Ce(NO₃)₃·6H₂O, no Ce-containing product was isolated, except halite (NaCl), as confirmed by its powder diffraction pattern.

Crystal structure determination

Laboratory X-ray powder diffraction (XRPD) patterns were collected on a PANanalytical X'Pert Pro diffractometer in a

Bragg–Brentano reflection configuration by using a Ge(111) primary monochromator (Cu Kα₁) and the X'Celerator detector. XRPD patterns were auto-indexed using the DICVOL06 program¹³ and the space groups derived from the observed systematic extinctions were consistent with *P* 2₁/*m* or *P* 2₁.

For the structural determination of **1**, powder X-ray diffraction data were collected at the high resolution beamline ID31 of the ESRF, European Synchrotron Radiation Facility, (Grenoble, France). A wavelength of 0.2998 Å was selected with a double-crystal Si (111) monochromator and calibrated with Si NIST (*a* = 5.43094 Å). The Debye–Scherrer configuration was used with the sample loaded in a rotating borosilicate glass capillary of diameter of 1.0 mm. The overall measuring time was ≈100 min in order to have satisfactory statistics over the angular range 1.0–15° (in 2θ). The data from the multi-analyser Si(111) stage were normalised and summed into 0.003° step size with local software. Its crystal structure was solved following an *ab initio* methodology. Initially the centrosymmetric space group *P* 2₁/*m* was selected for the extraction of the integrated intensities using the Le Bail method¹⁴ implemented in the GSAS package.¹⁵ The structure determination was carried out by direct methods using the programs Sir2006¹⁶ by default setting. The best solution gave the atomic positions for the lanthanum atom, two phosphorus atoms and some oxygen atoms of both phosphonate groups. The missing atoms were localized by difference of Fourier maps. However, not all water molecules could be found in *P* 2₁/*m*. The crystal structure could be satisfactorily completed in the non-centrosymmetric space group *P* 2₁ which is in agreement with the hydrogen-phosphonate stoichiometry of **1**. Furthermore, the fit in space group *P* 2₁ was better than in *P* 2₁/*m*. The structure was optimised by the Rietveld method¹⁷ using the program GSAS¹⁵ and the graphic interface EXPGUI.¹⁸ The following soft constraints were imposed in order to preserve chemically reasonable geometries for the phosphonate, amine groups and alkyl chains: PO₃C tetrahedron/P–O (1.53(1) Å), P–C (1.80(1) Å), O...O (2.55(2) Å), O...C (2.73(2) Å), N(CH₂)₃ amine group/N–C (1.50(1) Å), C...C (2.45(2) Å) and/alkyl chain/C–C (1.50(1) Å), C_{chain}...C_{chain} (2.50(2) Å). Hydrogen atoms were not included in the refinements. The final weight factor for the soft constraints was 20. Only three isotropic atomic displacement parameters were refined, one for the La atom, a second for the P atoms and a third one for the rest of the atoms. The crystal structures for the anhydrous, La(H₅DTMP) (**2**), and rehydrated phases, La(H₅DTMP)·2H₂O (**3**), were obtained by Rietveld refinement using as starting model the crystal structure of **1**. The Rietveld plots for **1–3** are given in the Electronic Supplementary Information (ESI†) and the relevant crystallographic data are presented in Table 1. Crystal structures have been deposited at CCDC and the reference codes are also given in Table 1.

Thermal analysis

Differential thermal analysis (DTA) and thermogravimetric analysis (TGA) data were recorded on an SDT-Q600 analyzer from TA instruments. Initially, the temperature was varied from RT to 1273 K at a heating rate of 10 K min⁻¹. Measurements were carried out on a sample in an open platinum crucible under a flow of air. Secondly, a set of measurements were carried out

Table 1 Crystal data and selected structure refinement parameters for compounds 1–3

Compound reference	1	2	3
Compound abbreviation	La(H ₅ DTMP)·7H ₂ O	La(H ₅ DTMP)	La(H ₅ DTMP)·2H ₂ O
Chemical formula	C ₁₀ H ₃₉ LaN ₂ O ₁₉ P ₄	C ₁₀ H ₂₅ LaN ₂ O ₁₂ P ₄	C ₁₀ H ₂₉ LaN ₂ O ₁₄ P ₄
Formula Mass	753.91	627.91	663.96
Crystal system	Monoclinic	Monoclinic	Monoclinic
<i>a</i> /Å	13.3444(5)	12.2967(15)	12.5035(6)
<i>b</i> /Å	18.7820(4)	17.7483(21)	18.1874(10)
<i>c</i> /Å	5.848903(23)	5.9887(8)	5.82000(35)
α (°)	90.0	90.0	90.0
β (°)	97.711(4)	104.626(6)	101.923(4)
γ (°)	90.0	90.0	90.0
Unit cell volume/Å ³	1452.71(10)	1264.7(4)	1295.95(11)
<i>V</i> _{non-H-atom} /Å ³	20.2	21.8	20.9
Temperature/K	293	473	293
Space group	<i>P</i> 2 ₁	<i>P</i> 2 ₁	<i>P</i> 2 ₁
No. of formula units per unit cell, <i>Z</i>	2	2	2
No. of independent reflections	1041	1072	1098
Data/Restraints/Parameters	4734/68/139	4547/65/105	5134/74/127
<i>R</i> _{WP}	0.0741	0.0812	0.0946
<i>R</i> _P	0.0532	0.0604	0.0604
<i>R</i> _F	0.0506	0.0504	0.0630
CCDC number	810316	810315	810317

under N₂ flow, saturated with water at 298 K, to characterise the release and uptake of water by **1**.

On the other hand, a thermogravimetric study for **1** was carried out for the sample loaded in an Anton Paar HTK 1200 N chamber under static air. Data were collected at 303, 423, 453 and 473 K with a heating rate of 5 K min⁻¹ and a delay time of 10 min to ensure thermal stabilization. The data acquisition range was 4–80° (2 θ) with a step size of 0.008° and an equivalent counting time of 100 s step⁻¹.

Textural characterization

The nitrogen sorption–desorption isotherms were measured in a Micromeritic ASAP 2020 apparatus. The samples were degassed under high vacuum and the isotherms, obtained at 77 K, were analyzed by the Brunauer–Emmett–Teller (BET) method. Carbon dioxide sorption isotherms were measured in a Micromeritic ASAP 2020 apparatus. The samples were degassed at 463 K for 2 h and the isotherms obtained at 273 K were analyzed by the Dubinin–Radushkevich (DR) method.¹⁹ The adsorption study of NO (at 298 K) was carried out following the experimental procedure described elsewhere.²⁰ The samples were outgassed at 463 K under vacuum for 2 h.

Conductivity characterization

Electrical characterization was carried out on a cylindrical pellet (~10 mm of diameter and ~1 mm of thickness) obtained by pressing ~0.02 g of sample at 1000 MPa, for 2 min. The pellet was pressed between porous C electrodes (Sigracet, GDL 10 BB, no Pt). Impedance spectroscopy data were collected using a HP4284A impedance analyzer over the frequency range from 20 Hz to 1 MHz with an applied voltage of 0.2 V. Electrical measurements were taken at 289, 291, 293, 295 and 297 K under a 5% H₂–Ar flow and at 98% relative humidity over the

temperature range. All measurements were electronically controlled by the winDETA package of programs.²¹

Results and discussion

La(H₅DTMP)·7H₂O (**1**) was obtained by room temperature crystallization from acidic solutions containing La³⁺ and H₅DTMP. The latter exists in a zwitterionic form (each phosphonate group is monodeprotonated, the two N atoms are protonated) in pH regions < 3, with a total charge of “–2”. However, in the presence of metal ions (trivalent, in particular) further phosphonate deprotonation occurs. In this case, one of the four phosphonate groups is fully (doubly) deprotonated, hence the ligand acquires a “–3” total charge. Thus, the charges (“+3” from the lanthanum and “–3” from the ligand) are balanced, and the final product is a neutral framework. The kinetics of the crystallization process is very slow (2 months), however this time can be reduced by using more concentrated solutions of the reagents without extensive compromise in the crystallinity of the final product. A similar synthesis for the Sm derivative was also carried out by following the same synthetic procedure. The samarium derivative was isolated and its powder pattern showed lower crystallinity. However, the habit of the Sm-hybrid pattern is similar to that of **1**, which points out the isostructural nature of this compound, see Fig. S4 in the ESI.†

Crystal structure

Compound **1** crystallizes in a monoclinic unit cell with 36 non-H atoms in the asymmetric part of the unit cell. As shown in Fig. 1, the asymmetric unit contains one crystallographically unique La³⁺ ion, one [H₅DTMP]³⁻ anion, and 7 lattice waters. The ligand can be described as a “zwitter ion”, as the N atoms remain protonated. The La³⁺ centre is coordinated exclusively by six phosphonate oxygen atoms (Figure 1) from six different

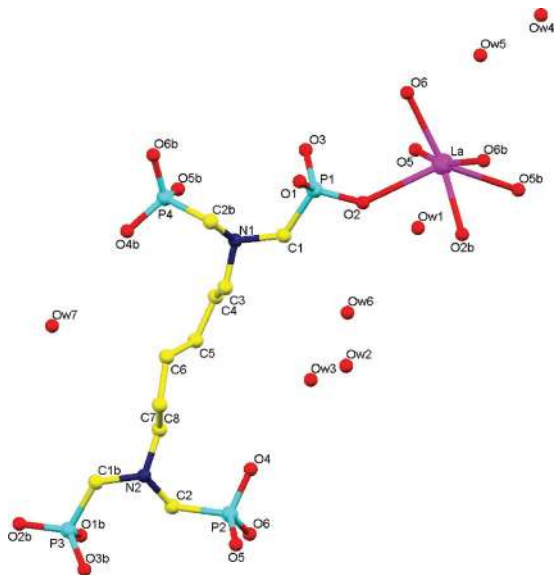


Fig. 1 Structural unit of $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**) with atom labelling.

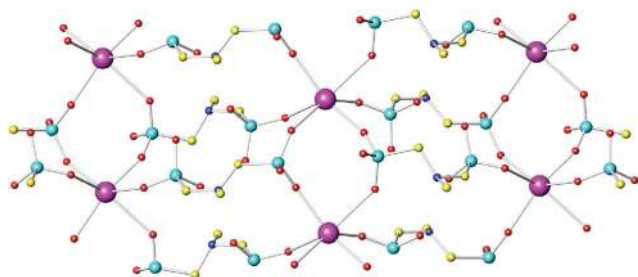


Fig. 2 Ball and stick view of a layer (*bc* plane) for $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**) showing the 8-membered rings (vertical) and the 16-membered rings (horizontal). Colour codes, La: purple; P: shy-blue; O: red; C: small yellow spheres; N: small blue spheres.

$[\text{H}_5\text{DTMP}]^{3-}$ anions. The La–O distances are in the range 2.28(3)–2.80(3) (Table S1†).

There are four crystallographically independent phosphorous atoms, with pairs (P1, P4) and (P2, P3) located in the opposite extremes of the ligand molecule. P1 and P3 are bonded to the La^{3+} centres through only one oxygen atom, whereas the remaining two phosphonate oxygens are non-coordinating. Higher connectivity is displayed by P2 and P4 groups, which bridge two LaO_6 polyhedra, through two oxygen atoms. This type of linkage gives rise to infinite chains along the *c*-axis, forming 8-member rings (see Fig. 2). The third phosphonate oxygen atoms of P2 and P4 are uncoordinated and point out of the plane. The chains of LaO_6 polyhedra, interconnected by bridging phosphonate tetrahedra, build a corrugated layer in the plane *bc*, containing inside 16-membered rings (see Fig. 2). Neighbouring layers are connected by the organic linker, resulting in an open 3D pillared framework, in which the 1-D channels are filled by seven lattice water molecules (see Fig. 3a and S5†). Based on the above structural data the precise molecular formula of $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**) can be expressed as $\text{La}[(\text{O}_3\text{PCH}_2)(\text{HO}_3\text{PCH}_2)\text{N}(\text{H})(\text{CH}_2)_6\text{N}(\text{H})(\text{CH}_2\text{PO}_3\text{H}_2)]\cdot 7\text{H}_2\text{O}$.

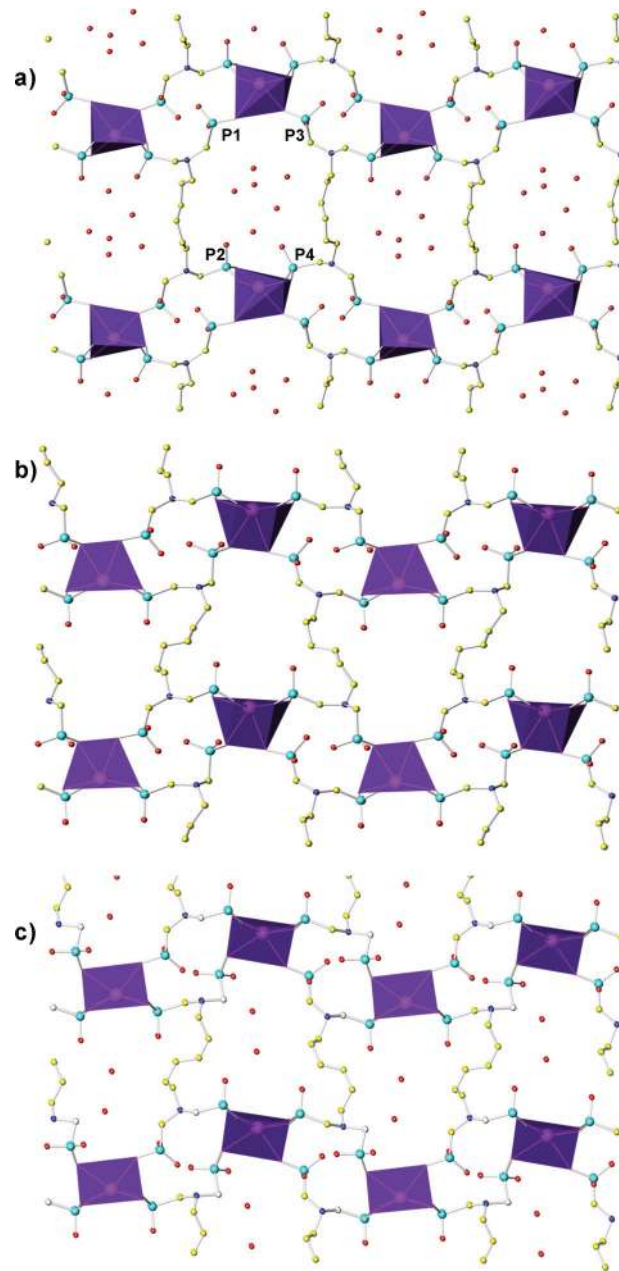


Fig. 3 Crystal structures of the hybrids showing the structural changes during the dehydration/rehydration processes: (a) $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**); (b) $\text{La}(\text{H}_5\text{DTMP})$ (**2**); (c) $\text{La}(\text{H}_5\text{DTMP})\cdot 2\text{H}_2\text{O}$ (**3**). Selected atoms are labelled.

Five lattice waters are situated close to the centre of the channels interacting to each other by H-bonds (see Fig. S6 and Table S2†). The two remaining lattice waters, Ow1 and Ow5, are closer to the lanthanum atoms and interacting strongly by H-bond with the oxygen and of two phosphonate groups (see Table S2†).

The way the organic linker is coordinated to La^{3+} is markedly different from the coordination mode adopted in $\text{Ca}(\text{H}_6\text{DTMP})$ and $\text{Gd}(\text{III})$ -diaminotetraphosphonates,^{5b} despite the fact that ions La^{3+} and Ca^{2+} are quite similar in size and possess the same metal : linker ratio in the corresponding derivatives. Thus, for the

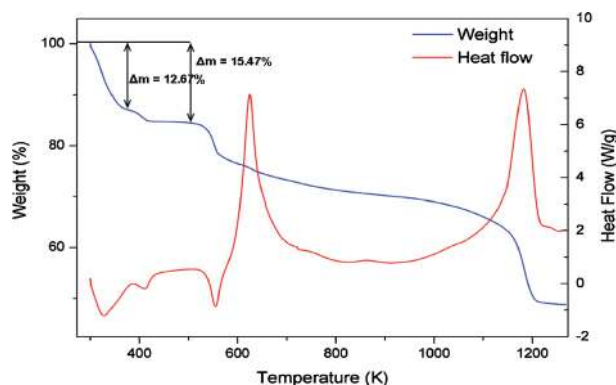


Fig. 4 TGA-DTA curves of **1**.

layered $\text{Ca}(\text{H}_6\text{DTMP})$ derivatives,¹² each amino-bis(methylene-phosphonate) moiety coordinates to a Ca^{2+} centre. Each H_6DTMP ligand bridges two Ca^{2+} centres giving rise to infinite chains, connected to each other through Ca–O bonds stemming from the remaining phosphonate O's. However, in compound **1** only phosphonate groups bridge lanthanum polyhedra in infinite chains with 8-membered rings separating the metal centres (see Fig. 2). These structural features (8-atom and 16-atom rings) are also present in $\text{Zn}(\text{H}_6\text{DTMP})\cdot\text{H}_2\text{O}$ but the resulting framework is different.^{7c} Other metal tetramethylene-diamine-*N,N,N',N'*-tetra-kismethylenephosphonates display pillared frameworks but with different connectivities.²²

Thermal behaviour

Coupled TGA-DTA curves for **1** are depicted in Fig. 4. The TGA curve shows that complete dehydration proceeds through two stages, leading to a weight loss of 15.5% (calcd 16.7%) at 423 K. The first step occurs from RT up to 373 K and corresponds to the release of five lattice waters (weight loss exp. 12.7%, calcd 11.3%). The second step, from 373 to 413 K, corresponds to the loss of two additional water molecules.

The water release pattern measured by TGA is in agreement with the structural study. The first mass loss corresponds to the loss of the five lattice waters located in the centre of the channels. The second mass loss is due to the two water molecules strongly H-bonded to the phosphonate groups. There is a plateau from ~ 423 to ~ 503 K where no mass loss is observed. For temperatures higher than 523 K, the material decomposes in two steps (523–673 K and 673–1223 K). The total observed mass loss of $\sim 51.1\%$ at 1273 K corresponds well to the calculated 50.17% (based on the formation of $\text{La}(\text{PO}_3)_3$, which was identified by its powder pattern, PDF #00-033-0717).

The dehydration process has been followed by thermodiffraction (see Fig. 5). In agreement with the TGA study, full dehydration takes place at 423 K. Heating at this temperature results in a crystalline dehydrated phase, $\text{La}(\text{H}_5\text{DTMP})$ (**2**). The process is only partially reversed after cooling at room temperature and exposing **2** to the air for 1 h. Under these conditions, only two water molecules are incorporated within the framework. This partial rehydration leads to a third crystalline phase, $\text{La}(\text{H}_5\text{DTMP})\cdot 2\text{H}_2\text{O}$ (**3**) that seems to remain stable at room temperature. **3** was kept at room temperature for 7 days in a closed

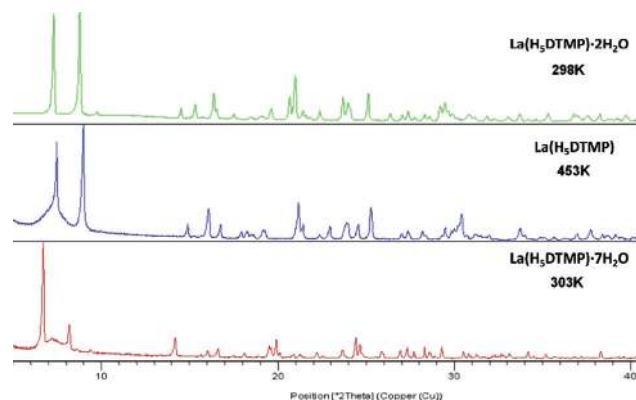


Fig. 5 Selected thermodiffraction data for $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**).

vessel under a humid atmosphere, produced by saturated NaCl, and with the powder pattern of the solid remaining unchanged. Both phases, **2** and **3**, were autoindexed using the DICVOL06 program in monoclinic unit cells related to the initial cell.

The fundamental topology and dimensionality of the network, characteristic of phase **1** remain basically unchanged in phases **2** and **3** (Fig. 3b and 3c, respectively). However, the loss of lattice water and further partial rehydration entails changes in the unit cell parameters, with *a* and *b* being shrunk upon removing water from the framework (Table 1). The structural changes may be followed by comparing four selected representative distances (see Fig. 3 and Table 2). The removal of water results in several structural changes: (1) Partial shrinkage of the structure occurs, as inferred from lower interlamellar La...La distance. (2) Higher deformation of the inorganic layers, deduced from higher sheet corrugation distances (calculated as the height of the triangle defined by three adjacent lanthanums within a sheet). (3) Changes in the relative positions of the phosphonate groups with respect to the metal centres (P1...P3 and P2...P4 distances). It is noteworthy to highlight that the “long” P1...P3 distance shortens, but, in contrast, the “short” P2...P4 distance lengthens on dehydration. These two distances are quite similar in the anhydrous **2**. All these changes are possible due to the partially flexible nature of the C_6H_{12} chains. The free solvent accessible volume in **2**, derived from PLATON²³ routine analysis (filling of the cavities with probe spheres of radius 1.2 Å and with hydrogen atoms situated geometrically), was found to be 18.7%. The lattice water molecules in **3** are localized in triangular cavities, interacting with oxygen atoms of the phosphonate groups through H-bonds (Fig. 3c, Table S3†).

In order to study the affinity of LaH_5DTMP for water molecules a thermal analysis with two cycles of heating and cooling

Table 2 Key structural parameters for $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**), $\text{La}(\text{H}_5\text{DTMP})$ (**2**) and $\text{La}(\text{H}_5\text{DTMP})\cdot 2\text{H}_2\text{O}$ (**3**)

	1	2	3
$(\text{La}\cdots\text{La})_{\text{inter}}$ (Å)	15.27	12.30	12.50
h^a (Å)	4.18	5.06	4.73
P1...P3 (Å)	6.18(5)	4.58(3)	5.64(3)
P2...P4 (Å)	4.36(3)	4.61(4)	4.90(3)

^a Height for a La...La...La intralayer triangle

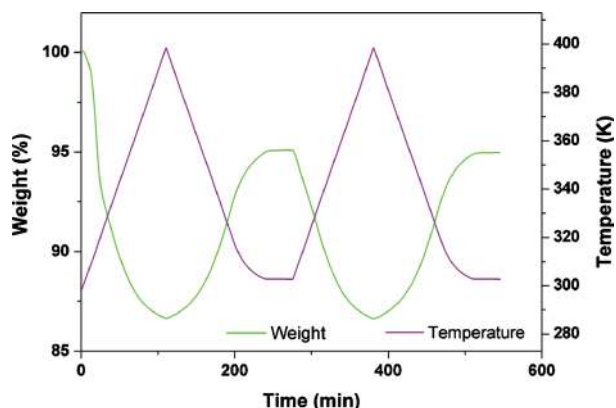


Fig. 6 TGA evolution (green curve) of **1** under a water saturated N_2 flow.

(at 1 K min^{-1} up to 403 K) was carried out under a N_2 flow saturated with water. As shown in Fig. 6, during the heating process the initial weight loss observed is 13.40% (calcd 13.1%) corresponding to removal of 5.5 lattice waters. After cooling, and in the presence of a high relative humidity, a weight gain of 8.43% is observed. This percentage corresponds to the adsorption of 3.5 lattice waters, an amount that is higher than that observed under normal conditions. Therefore, at the end of this experiment, a stoichiometry close to $\text{La}(\text{H}_5\text{DTMP})\cdot 5\text{H}_2\text{O}$ is expected. Moreover, the process is completely reproducible after a second cycle. Therefore, all the structural changes occurring during the dehydration/rehydration processes demonstrate that the framework is flexible as it may adopt different configurations as a function of the presence/absence of guest molecules and the environmental conditions.

A further study to confirm the flexibility of the framework upon water exposure was performed at higher temperatures. In this second study, **1** was heated under water saturated N_2 flow up to 473 K , which led to the formation of **2**. With cooling under water saturated N_2 showed an increase of weight of $10.97\text{ wt}\%$ which is in agreement with the formation of $\text{La}(\text{H}_5\text{DTMP})\cdot 5\text{H}_2\text{O}$.

Microporosity characterisation

1 was outgassed under high vacuum at room temperature for 15 h or at 473 K for 2 h . Both treatments resulted in the anhydrous material, **2**, as determined by the mass losses. On the one hand, the N_2 isotherm gave no porosity with specific surface lower than $3\text{ m}^2\text{ g}^{-1}$. On the other hand, the CO_2 isotherm (not shown), collected at 273 K and under pressure up to 1 bar , gave a micropore surface area of $84\text{ m}^2\text{ g}^{-1}$, deduced from the Dubinin–Radushkevich¹⁹ equation, and an adsorption capacity of 0.35 mmol g^{-1} .

We highlight that the isotherm is not saturated at 1 bar and higher pressures would be necessary to access the full pore volume of the structure. This is in agreement with the shape of the 1D channels in **2**. Higher pressures would allow a gate-opening mechanism that may allow a higher gas uptake. A high-pressure synchrotron powder diffraction study is planned in order to characterise this flexible framework as a function of the loaded species (CO_2 , N_2 , etc.).

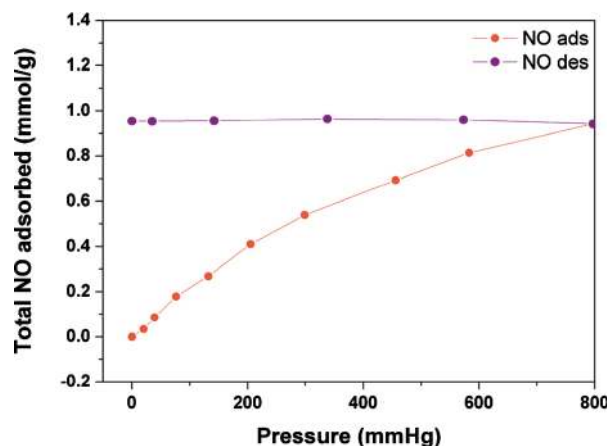


Fig. 7 NO adsorption–desorption measurement for $\text{La}(\text{H}_5\text{DTMP})$ (**2**) at 298 K .

Some preliminary studies on NO adsorption have been also carried out for **2**. Fig. 7 shows the isotherms for NO adsorption–desorption at 298 K . The NO adsorption capacity at $\sim 1\text{ bar}$ was 0.95 mmol g^{-1} (or $2.85\text{ wt}\%$), similar to that found in zeolites²⁴ and $\text{Sn}(\text{IV})$ metal diphosphonates.²⁵ The hysteresis of the isotherm is significant, indicating a strong irreversible adsorption of NO inside the pores. This irreversible adsorption is likely due to the bonding of the NO molecules to the lanthanum cations.

Proton-conducting characterisation

A preliminary ionic conductivity study of **1** has been carried out as the structural study showed 1D channels filled with water molecules (see Fig. 3 and S6†) that may allow proton-conductivity to take place. The impedance study shows that indeed **1** exhibits proton conductivity. Fig. 8 shows the real (Z') and imaginary (Z'') parts of the complex-plane impedance data for **1**. Cole–Cole circular arc fitting gave a conductivity (σ) of $8\text{ }10^{-3}\text{ S cm}^{-1}$ at

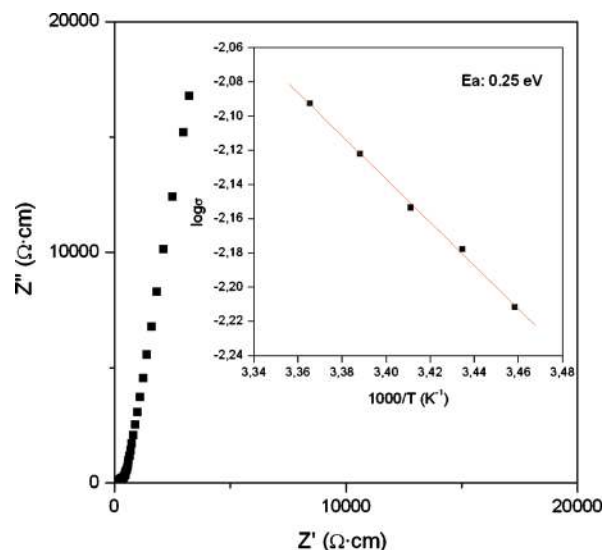


Fig. 8 Complex impedance plane plot for $\text{La}(\text{H}_5\text{DTMP})\cdot 7\text{H}_2\text{O}$ (**1**) at 297 K and 98% RH. The inset shows the Arrhenius plot over the $289\text{--}297\text{ K}$ temperature range in the same conditions.

297 K and 98% of relative humidity (RH). The temperature dependent study is also given in Fig. 8. Arrhenius plot shows a linear behaviour with an activation energy of 0.25 eV. This small number is within the range typically attributed to a Grotthuss transfer mechanism *via* water molecules, 0.1–0.4 eV.²⁶ A full proton-conductivity study is out of the scope of this work and it will be reported elsewhere.

The RT obtained value of the proton conductivity is higher than that recently reported for a metal phosphonate MOF material, $\sigma = 3.5 \times 10^{-5} \text{ S cm}^{-1}$ at 298 K and 98 % RH.²⁷ The activation energy for the proton transfer process derived from bulk conductivity was 0.17 eV. Values of the same magnitude for the proton conductivity, $\sigma = 1.1 \times 10^{-3} \text{ S cm}^{-1}$, at room temperature have been very recently reported for a carboxylate-MOF.²⁸ In this case, the activation energy of the proton conductivity process was nearly the same 0.23 eV.

Conclusions

Two open-framework microporous materials Ln(H₅DTMP) (Ln = La and Sm) are reported. The focus in this paper was on the La derivative, which was fully characterized. The as-synthesised compound contains seven lattice waters which may be removed at moderate temperatures while the crystalline metal–organic framework is maintained. The anhydrous material shows high affinity for water but the rehydrate stoichiometries depend upon the experimental conditions. The crystal structures indicate that the lanthanum coordination environment is made-up exclusively by phosphonate oxygens, *i.e.* none of the water molecules are coordinated to the La³⁺ centres. This structural feature may have implications in the luminescent properties to be studied. The incorporation of a ligand possessing an aliphatic $-(\text{CH}_2)_6-$ chain provides substantial framework flexibility as the structures are adapted to the guest molecule (water) content. The anhydrous material is ultramicroporous and the specific surface for CO₂ adsorption (at 273 K and approximately 1 bar of CO₂) is 84 m² g⁻¹. Finally, **1** is a proton-conductor with a conductivity of $8 \times 10^{-3} \text{ S cm}^{-1}$ at RT and an activation energy of 0.25 eV.

Acknowledgements

The work at UMA was funded by MAT2010-15175 research grant (MICINN, Spain). The work at the UoC was supported by a grant from the Research Committee of the University of Crete, ELKE, (KA 2573). ESRF is thanked for the provision of synchrotron X-ray beamtime at ID31 beamline.

Notes and references

- (a) J. Rocha, L. D. Carlos, F. A. Almeida Paz and D. Ananias, *Chem. Soc. Rev.*, 2011, **40**, 926; (b) S. Ma, D. Yuan, X.-S. Wang and H.-C. Zhou, *Inorg. Chem.*, 2009, **48**, 2072; (c) B. V. Harbuzaru, A. Corma, F. Rey, P. Atienzar, J. L. Jordá, H. García, D. Ananias, L. D. Carlos and J. Rocha, *Angew. Chem., Int. Ed.*, 2008, **47**, 1080; (d) B. V. Harbuzaru, A. Corma, F. Rey, J. L. Jordá, D. Ananias, L. D. Carlos and J. Rocha, *Angew. Chem., Int. Ed.*, 2009, **48**, 6476; (e) J.-C. Ma, Y.-Y. Liu, J. Yang, Y.-Y. Liu and J.-F. Ma, *CrystEngComm*, 2011, **13**, 3498; (f) P. Silva, F. Vieira, A. C. Gomes, D. Ananias, J. A. Fernandes, S. M. Bruno, R. Soares, A. A. Valente, J. Rocha and F. A. Almeida Paz, *J. Am. Chem. Soc.*, 2011, **133**, 15120.
- (a) L. D. Carlos, R. A. S. Ferreira, V. d. Z. Bermudez and S. J. L. Ribeiro, *Adv. Mater.*, 2008, **20**, A1–26; (b) D. T. de Lill and C. L. Cahill, *Prog. Inorg. Chem.*, 2007, **55**, 143.
- (a) B. Cai, P. Yang, J.-W. Dai and J.-Z. Wu, *CrystEngComm*, 2011, **13**, 985; (b) F. Costantino, P. L. Gentili and N. Audebrand, *Inorg. Chem. Commun.*, 2009, **12**, 406.
- (a) J.-G. Mao, *Coord. Chem. Rev.*, 2007, **251**, 1493; (b) N. Zhang, Z. Sun, Y. Zhu, J. Zhang, L. Liu, C. Huang, X. Lu, W. Wang and F. Tong, *New J. Chem.*, 2010, **34**, 2429; (c) G. Pawlicki and S. Lis, *Opt. Mater.*, 2011, **33**, 1544.
- (a) R.-C. Wang, Y. Zhang, H. Hu, R. R. Frausto and A. Clearfield, *Chem. Mater.*, 1992, **4**, 864; (b) S.-M. Ying, X.-R. Zeng, X.-N. Fang, X.-F. Li and D.-S. Liu, *Inorg. Chim. Acta*, 2006, **359**, 1589; (c) S. Bauera, T. Bein and N. Stock, *J. Solid State Chem.*, 2006, **179**, 145; (d) J.-M. Rueff, N. Barrier, S. Boudin, V. Dorcet, V. Caignaert, P. Boullay, G. B. Hix and P.-A. Jaffrès, *Dalton Trans.*, 2009, 10614.
- M. Plabst, L. B. McCusker and T. Bein, *J. Am. Chem. Soc.*, 2009, **131**, 18112.
- (a) F. Costantino, A. Ienco, P. L. Gentili and F. Presciutti, *Cryst. Growth Des.*, 2010, **10**, 4831; (b) N. Stock, M. Rauscher and T. Bein, *J. Solid State Chem.*, 2004, **177**, 642; (c) K. D. Demadis, C. Mantzaridis, R. G. Raptis and G. Mezei, *Inorg. Chem.*, 2005, **44**, 4469; (d) K. D. Demadis, E. Barouda, N. Stavgiannoudaki and H. Zhao, *Cryst. Growth Des.*, 2009, **9**, 1250; (e) K. D. Demadis, E. Barouda, H. Zhao and R. G. Raptis, *Polyhedron*, 2009, **28**, 3361; (f) K. D. Demadis, S. D. Katarachia, H. Zhao, R. G. Raptis and P. Baran, *Cryst. Growth Des.*, 2006, **6**, 836.
- A. Mondry and R. Janicki, *Dalton Trans.*, 2006, 4702.
- J. Wu, H. Hou, H. Han and Y. Fan, *Inorg. Chem.*, 2007, **46**, 7960.
- (a) K. D. Demadis, E. Barouda, R. G. Raptis and H. Zhao, *Inorg. Chem.*, 2009, **48**, 819; (b) K. D. Demadis, C. Mantzaridis and P. Lykoudis, *Ind. Eng. Chem. Res.*, 2006, **45**, 7795; (c) E. Akyol, M. Öner, E. Barouda and K. D. Demadis, *Cryst. Growth Des.*, 2009, **9**, 5145.
- (a) M. Su, Y. Qiu and W. Jia, *Adv. Ther.*, 2005, **22**, 297; (b) B. Mathew, S. Chakraborty, T. Das, H. D. Sarma, S. Banerjee, G. Samuel, M. Venkatesh and M. R. A. Pillai, *Appl. Radiat. Isot.*, 2004, **60**, 635.
- R. M. P. Colodrero, A. Cabeza, P. Olivera-Pastor, A. Infantes-Molina, E. Barouda, K. D. Demadis and M. A. G. Aranda, *Chem.–Eur. J.*, 2009, **15**, 6612.
- A. Boultif and D. Louer, *J. Appl. Crystallogr.*, 2004, **37**, 724.
- A. Le Bail, H. Duroy and J. L. Fourquet, *Mater. Res. Bull.*, 1988, **23**, 447.
- A. C. Larson and R. B. Von Dreele, “General Structure Analysis System (GSAS)”, Los Alamos National Laboratory Report LAUR 86-748 (2004)
- M. C. Burla, R. Caliandro, M. Camalli, B. Carrozzini, G. L. Casciarano, C. De Caro, C. Giacovazzo, G. Polidori, D. Siliqi and R. Spagna, *J. Appl. Crystallogr.*, 2007, **40**, 609.
- H. M. Rietveld, *J. Appl. Crystallogr.*, 1969, **2**, 65–71.
- B. H. Toby, *J. Appl. Crystallogr.*, 2001, **34**, 210.
- M. Dubinin and L. V. Radushkevich, *Proc. Acad. Sci. USSR*, 1947, **55**, 331.
- B. Xiao, P. S. Wheatley, X. Zhao, A. J. Fletcher, S. Fox, A. G. Rossi, I. L. Regli, K. M. Thomas and R. E. Morris, *J. Am. Chem. Soc.*, 2007, **129**, 1203.
- winDETA*, Novocontrol GmbH, Hundsangen, Germany, 1995
- (a) N. Stock, A. Stoll and T. Bein, *Microporous Mesoporous Mater.*, 2004, **69**, 65; (b) N. Stock, M. Rauscher and T. Bein, *J. Solid State Chem.*, 2004, **177**, 642; (c) N. Stock and T. Bein, *Angew. Chem., Int. Ed.*, 2004, **43**, 749.
- A. L. Spek, *Acta. Crystallogr., Sect. D*, 2009, **65**, 148.
- P. S. Wheatley, A. R. Butler, M. S. Crane, A. G. Rossi, I. L. Megson and R. E. Morris, *J. Am. Chem. Soc.*, 2006, **128**, 502.
- M. M. Gomez-Alcantara, A. Cabeza, P. Olivera-Pastor, F. Fernandez-Moreno, I. Sobrados, J. Sanz, R. E. Morris, A. Clearfield and M. A. G. Aranda, *Dalton Trans.*, 2007, 2394.
- P. Colomban, Proton Conductors: Solids, Membranes and Gels Materials and Devices, *Chemistry of Solid State Materials*, Vol. 2; Cambridge University Press, Cambridge, U.K., 1992.
- J. M. Taylor, R. K. Mah, I. L. Moudrakovski, C. I. Ratcliffe, R. Vaidhyanathan and G. K. H. Shimizu, *J. Am. Chem. Soc.*, 2010, **132**, 14055.
- E. Pardo, C. Train, G. Gontard, K. Boubekour, O. Fabelo, H. Liu, B. Dkhil, F. Lloret, K. Nakagawa, H. Tokoro, S. Ohkoshi and M. Verdager, *J. Am. Chem. Soc.*, 2011, **133**, 15328.

