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Sequestration of orthophosphate by Ca₂Al-NO₃ Layered Double Hydroxide – Insight into reactivity and mechanism.

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Abstract

This study reports the mechanism of reactivity of phosphate anions (HPO₄²⁻) with Ca₂Al-NO₃ LDH. Sequestration of PO₄ by Ca₂Al-NO₃ was quantified by fitting the adsorption isotherm with a Langmuir model. Adsorption capacity (65.2 mgP.g⁻¹ of LDH) is greater than the theoretical anion exchange capacity indicating further mechanism of sequestration. Partial dissolution of CaAl-NO₃ results in precipitation of multiple Ca-PO₄ phases. Combined characterizations by XRD, ²⁷Al and ³¹P NMR, FTIR/Raman and SEM highlight the competitive precipitation of hydroxyapatite and brushite within the LDH structure. Multiple environment of PO₄ anions offer promising perspective for CaAl-NO₃/PO₄ as slow release fertilizer.

Keywords: Layered Double Hydroxides; $\text{Ca}_2\text{Al-NO}_3$ hydrocalumite, orthophosphate; adsorption, hydroxyapatite, brushite.

1. Introduction

Inorganic orthophosphate (PO_4) is an essential resource for agriculture and animal feed additives. Indeed, 82% and 7% of their production are used respectively as fertilizers and food production (Schroeder et al., 2009). Unfortunately, PO_4 is a non-renewable resource and the impoverishment of world stocks threaten the food supply to the world population (EU Report, 2013). With a mean production of 160 Mt per year over 16 000 Mt estimated reserves, in 100 years PO_4 scarcity will be a severe problem. Added to these human and economic dangers, a major environmental concern arises from the dissemination of PO_4 in the environment and its accumulation in surface waters through land farming and waters discarded from wastewater treatment plants (WWTP) causing harmful eutrophication of inland and coastal waters (Oliveira et al., 2013; Farley et al, 2012). Consequently, the development of sustainable PO_4 management processes is of high societal concern.

Recovery of PO_4 from WWTP and processing as recycled PO_4 slow release fertilizers (SRF), are under intensive investigations to both limit phosphate wastes and improve circular economic model of this critical raw material (Cieslik et al., 2017; Desmidt et al., 2015). Most sustainable water-cleaning processes in WWTPs target the precipitation of phosphate as struvite ($\text{NH}_4\text{MgPO}_4 \cdot 6\text{H}_2\text{O}$) (Ye et al., 2014) or hydroxyapatite ($\text{Ca}_5(\text{PO}_4)_3(\text{OH})$, HAP) (Chen et al., 2009), for subsequent reuse as fertilizers. However struvite production is limited to large scale WWTPs and has so far failed to reach a sustainable economic model (Talboys et al., 2016).

Moreover, the low solubility of struvite and hydroxyapatite in water (Ricardo et al., 2012) limit their use as efficient SRF. Selective adsorption of PO₄ from wastewaters by inorganic adsorbents that can be recycle once loaded with PO₄ as SRF appears as an alternative green technology.

Layered double hydroxides (LDH) are natural and synthetic inorganic solids with a layered structure and general chemical formula $[M^{2+}_{1-x}M^{3+}_x(OH)_2]^{x+}[A^{n-}_{x/n}.mH_2O]^{x-}$ (M^{2+} , M^{3+} and A^{n-} being divalent and trivalent metals and anionic species). They display unique anion exchange properties largely investigated for the removal of oxoanions (NO₃⁻, PO₄³⁻, CrO₄²⁻, AsO₄³⁻,...) (Lazaridis et al., 2003) as well as they rank amongst the most selective and efficient adsorbents for PO₄ anions (Xu et al., 2010; Zhou et al., 2011; Bernardo et al., 2017; Everaert et al., 2018). Among the LDH, hydrocalumite-like LDH, a Ca₂Al-LDH display a very high chemical reactivity with PO₄ leading to great quantity of PO₄ removed from water, i.e. as high as 400 mg PO₄ per gram of LDH exceeding the theoretical anion exchange capacities (Zhou et al., 2011). This behavior makes Ca₂Al-LDH a very promising PO₄ sorbents and potential SRF. Few studies have investigated PO₄ release from Ca₂Al-PO₄ phases prepared by PO₄ loading on Ca₂Al LDH and their use as slow-release fertilizers. Interestingly, PO₄-loaded hydrocalumite has demonstrated synergistic performances for plant growth and development of bacteria (*Bradyrhizobium elkanii*) responsible for N₂ fixation (Jia et al., 2016).

The mechanism of adsorption of phosphate by Ca-based LDH adsorbents is still under examination (Radha et al., 2005). Some authors claimed that PO₄ adsorption follows a topotactic anion exchange reaction with intercalation of HPO₄²⁻ anions in between the LDH layers (Radha et al., 2005). However, most convincing results clearly show that Ca₂Al LDH undergo a progressive dissolution followed by the immediate precipitation of various Ca-PO₄ phases (Lu et al., 2016; Zhou et al., 2012). The dissolution/precipitation phenomenon is driven by the

difference between the solubility's of CaAl-LDH and most of the common calcium phosphate phases (Table 1). Hydroxyapatite ($\text{Ca}_5(\text{PO})_4(\text{OH})$) and brushite ($\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$) (often referred to DCPD for dicalcium phosphate dihydrate) are the most commonly reported calcium phosphate phases that can form from a reaction between $\text{Ca}_2\text{Al-LDH}$ and PO_4 (Khitous et al., 2016; Vieille et al., 2003). Brushite was obtained when PO_4 adsorption was performed at acidic pH (pH 5) while in basic medium (pH 11) HAP predominantly precipitates (Tsuji et al., 2014). Jeong-Woo Son et al. (Jeong-Woo Son et al., 2015) confirmed these results for phosphate removal by hydrocalumite at basic pH but they observed the precipitation of other phases (gibbsite, boehmite, calcite). The total conversion of hydrocalumite into one of these phases should result in more than 388 mg of adsorbed PO_4 (or 127 mg P) per g of hydrocalumite, values much higher than reported in literature (Lu et al., 2016). The structure and composition of the precipitated Ca- PO_4 phase as well as the rate of $\text{Ca}_2\text{Al-LDH}$ conversion are strongly dependent on various parameters such as $\text{Ca}^{2+}/\text{PO}_4^{3-}$ molar ratio, pH, PO_4 concentration. Moreover, the mechanisms of PO_4 adsorption and phase conversion are mostly ignored.

Conversion into hydroxyapatite is a limiting factor for a 100% PO_4 recovery and may hinder applications of these materials as SRF because extreme pH conditions would be needed to desorb PO_4 from HAP (Vieille et al., 2003).

As $\text{Ca}_2\text{Al-LDH}$ have a great ability for PO_4 sequestration and potential release, there is a need for further investigations in order to better understand the mechanism involved in this reaction and the role of chemical composition and morphology to control PO_4 adsorption and release. Therefore, our study focused on $\text{Ca}_2\text{Al-NO}_3$ LDH phase whose reactivity toward PO_4 adsorption has not been yet investigated. Combination of XRD, solid state NMR spectroscopy,

and electron microscopy characterization was implemented to identify the CaPO₄ phases formed and to assess the mechanism of reactivity of phosphate anions with Ca₂Al-NO₃ LDH materials.

2. Materials and methods

2.1. Materials

All the reagents were analytical grade and used without further purification. Ca(NO₃)₂·4H₂O, Al(NO₃)₃·9H₂O, NaOH, K₂HPO₄, (L)-ascorbic acid, hexaammonium heptamolybdate (NH₄)₆Mo₇O₂₄·4H₂O, sulfuric acid (H₂SO₄), potassium antimony(III) oxide tartrate (K(SbO)C₄H₄O₆·0.5H₂O) were purchased from Sigma-Aldrich. Adsorption studies were performed using K₂HPO₄. Deionized water was used exclusively in all procedures.

2.2. Synthesis of Ca₂Al-NO₃

Ca₂Al-NO₃ was synthesized by the co-precipitation method in a reactor using peristaltic pumps for simultaneous addition of Ca²⁺/Al³⁺ metal salts and NaOH in order to keep the pH constant following the procedure published by Vieille et al. (Vieille et al., 2003). The Ca²⁺/Al³⁺ molar ratio was fixed to 2 and the reagent concentrations equal to 1M and 2M respectively for ([Ca²⁺] + [Al³⁺]) and NaOH. The synthesis was performed at a pH = 11.5 and under N₂ flow to prevent CO₃²⁻ contamination. The reaction was vigorously stirred and completed in 4-5 h. The resulting slurry was aged at room temperature for 24 h. The mixture was centrifuged and the solid washed thoroughly with deionized water and dried at 40 °C for 24h.

2.3. Adsorption isotherms and modeling

The adsorption isotherm experiments were carried out in 2-ml Eppendorf vials. 15 mg of Ca₂Al-NO₃ LDH was suspended in 1.5 ml of K₂HPO₄ aqueous solution with concentrations ranging from 10 to 1000 mg P.L⁻¹ (0, 50, 100, 200, 300, 400, 500, 600, 700, 800, 900, 1000 mgP.L⁻¹) (P-refers to phosphorous). Suspensions were shaken for 2h00 at 20°C under air atmosphere in order to achieve the adsorption equilibrium (Khitous et al., 2016). The supernatants were separated using centrifugation (14000 rpm, 21475 G-force, 20 min) and pH values were measured to be in the range of 8.1±0.4 throughout the adsorption experiment. Concentrations of phosphate in the solution was determined using a Nicolet Evolution 500 UV-Vis spectrophotometer at $\lambda = 880$ nm, following the Molybdenum blue method (Broberg et al., 1988).

The adsorption capacity of the sorbent was measured by the difference between the initial and remaining concentrations of phosphate. The equilibrium adsorption concentrations, q_e (mg.g⁻¹), were calculated according the following equation $q_e = \frac{V(c_0 - c_e)}{W}$, where c_0 is the initial concentration of (mg P. L⁻¹); c_e is the equilibrium concentration (mg P.L⁻¹); V is the volume of the solution used (L); and W is the weight of the adsorbents (g). Sorption isotherms were obtained by plotting experimental q_e versus c_e . The adsorption isotherms were analyzed using the linearized equations of Langmuir and Freundlich, sorption models, expressed as:

$$\text{Langmuir: } \frac{c_e}{q_e} = \frac{1}{q_m K} + \frac{c_e}{q_m} \quad (1)$$

$$\text{Freundlich: } q_e = K_F c_e^{1/n} \quad (2)$$

where c_e ($\text{mg}\cdot\text{L}^{-1}$) is the concentration of phosphate at equilibrium and q_e ($\text{mg}\cdot\text{g}^{-1}$) is the sorption capacity of the solid at equilibrium. K (L/mg) is Langmuir constant, q_m ($\text{mg}\cdot\text{g}^{-1}$) is the saturated sorption capacity of sorbent. K_F and n are Freundlich parameters. K_{DR} (mol^2/J^2) is the activity coefficient. R is the universal gas constant ($8.314 \text{ J}/\text{mol K}$), and T (K) is temperature.

2.4. Phosphate Exchange with LDH

The phosphate exchange experiments were carried out in 30 ml adsorption vials with 90 mg of LDH and 9 ml of a K_2HPO_4 aqueous solution at 20°C under air atmosphere. The K_2HPO_4 concentrations equal to 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, and 0.9 g $\text{HPO}_4^{2-}\cdot\text{L}^{-1}$. The mixtures were stirred for 5h. Solids were separated by centrifugation (14000 rpm, 21475 G-force, 20 min) and washed, repeatedly. Precipitates were dried at 40°C for 24h. pH values of supernatants were measured to be in the range of 8.2 ± 0.3 throughout the exchange experiment. The concentration range was selected in order to vary the initial $\text{HPO}_4^{2-}/\text{Ca}_2\text{Al-NO}_3$ mass ratio from 32.6 to 97.9 $\text{mgP}\cdot\text{g}^{-1}$ of CaAl-NO_3 . Considering that such concentration correspond to mass ratio from the early stage of adsorption to saturation we then prepared a series of PO_4 loaded samples representative of all the steps of reaction.

2.5. Characterization techniques

Chemical analyses of the LDH were conducted using the ICP-OES after dissolving the samples in 1 M HNO_3 solution. X-ray diffraction (XRD) measurements were performed on a Philips X'Pert Pro diffractometer using $\text{CuK}\alpha$ radiation ($\lambda = 1.5405 \text{ \AA}$). PXRD patterns were recorded over the $5\text{--}70^\circ$ 2θ range at an accelerating voltage of 40 kV, a current of 30 mA, a scan speed of 2°min^{-1} . Fourier-transform infrared (FTIR) spectra were recorded with a Nicolet-5700

spectrophotometer in the range of 400–4000 cm^{-1} using the KBr pellet technique. The Raman spectra were recorded from 150 to 1500 cm^{-1} at room temperature using a confocal micro-Raman spectrometer (T64000 Jobin Yvon) with an excitation wavelength of 514.5 nm (argon-ion laser-line). Spectra were acquired at a spectral resolution of 1 cm^{-1} with charge coupled device (CCD) multichannel detector cooled by liquid nitrogen to 140 K coupled with Olympus confocal microscope. SEM images were recorded using a JSM-7500F FESEM operating at 3 KV. Samples to be imaged were mounted on conductive carbon adhesive tabs. ICP-AES analyses were realized with a HORIBA-Jobin-Yvon ULTIMA C spectrometer.

^{27}Al MAS and ^{27}Al 3QMAS NMR spectra were recorded on a Varian INOVA 600 MHz NMR spectrometer (14.1 T) equipped with a 3.2 mm triple resonance MAS probe using 15 kHz spinning speed and a 1s relaxation delay. ^{27}Al MAS NMR spectra were recorded using a short, selective $\pi/18$ pulse to ensure quantitative spectra. Quantitative ^{31}P MAS NMR spectra were recorded on a JEOL 500 MHz NMR spectrometer (11.7 T) using a 3.2 mm double resonance probe, a $\pi/4$ pulse, a spinning speed of 13 kHz, and a relaxation delay (typically 480-540 s) optimized for each sample. 85 % phosphoric acid (0.0 ppm) and 1 M AlCl_3 have been used as reference for ^{27}Al and ^{31}P , respectively. Deconvolutions of the ^{27}Al and ^{31}P MAS NMR spectra has been performed using QuadFit (Kemp et al., 2009) and MestReNova.

3. Results and Discussion

3.1. Structure and chemical analysis of $\text{Ca}_2\text{Al-NO}_3$

The chemical composition ($\text{Ca}_{1.91}\text{Al}(\text{OH})_{5.82}(\text{NO}_3) \cdot 1.63\text{H}_2\text{O}$) of the as-prepared $\text{Ca}_2\text{Al-NO}_3$ LDH phase, determined from ICP-AES and TGA data, is in good agreement with the chemical formula expected for a pure synthetic NO_3 -hydrocalumite. The high pH value (11.5)

used for the synthesis ensured a total coprecipitation of $\text{Ca}^{2+}/\text{Al}^{3+}$ metal ions. Indeed, the experimental $\text{Ca}^{2+}/\text{Al}^{3+}$ ratio (1.91) is close to the ideal composition ($\text{Ca}^{2+}/\text{Al}^{3+} = 2$) expected from initial molar ratio used. The powder X-ray diffractogram (PXRD) (Fig. 1A) and the FTIR spectrum (Fig. 1B) of $\text{Ca}_2\text{Al-NO}_3$ LDH confirmed the formation of the desired material. Indeed PXRD exhibits the structural features including sharp diffraction lines of a well-crystallized $\text{Ca}_2\text{Al-NO}_3$ hydrocalumite phase (Renaudin et al., 1999). The cell parameters were refined in the trigonal space group $P\bar{3}c1$ (Fig. 1). The main structural difference compared to the hydrotalcite structure arises from the distortion of the octahedral layer due to the hepta-coordination of Ca^{2+} ions (6+1) bonded to six OH groups and an intercalated H_2O molecule (Fig. SI-1). This corrugated layered structure is associated to typical Ca,Al–OH, Ca,Al–O and (Ca,Al)–O–(Ca,Al) lattice vibrations located at 782, 592, 527 and 420 cm^{-1} , respectively (Zhang et al., 2011). Intercalation of NO_3^- anions was confirmed by the strong infrared absorption at 1380 cm^{-1} and 854 cm^{-1} relative respectively to the antisymmetric stretching ν_3 and out-of-plane bending ν_2 vibrations of the D_{3h} NO_3^- anions. The broad vibrations between 3400 cm^{-1} and 3650 cm^{-1} are characteristics of all LDH and arises from the symmetric and asymmetric stretching ($\nu(\text{OH})$) of the hydroxide layers and intercalated water molecules (Fig. 1B), 3640 cm^{-1} and 34493 cm^{-1} , respectively for free (Ca₂Al)-O-H groups and H-bonded interlayered water molecules. The bending mode of interlayer water molecules appears between 1622-1637 cm^{-1} . $\text{Ca}_2\text{Al-NO}_3$ was precipitated as large hexagonal platelets with sizes between 1-3 μm and thickness ranging between 25-110 nm leading to a high aspect ratio (Fig. 7A) and a low BET specific surface area of 11 $\text{m}^2\cdot\text{g}^{-1}$. ^{27}Al MAS NMR experiment was employed to assess the purity of the parent $\text{Ca}_2\text{Al-NO}_3$ LDH (Fig. SI-2). The spectrum was simulated to determine the purity of the $\text{Ca}_2\text{Al-LDH}$. The isotropic chemical shift $\delta_{\text{iso}}(^{27}\text{Al}) = 11.0(4)$ ppm is in agreement

with the value reported by Qiu et al. (Qiu et al., 2015) and Jensen et al. (Jensen et al., 2016) for Ca₂Al-LDH intercalated respectively with NO₃⁻ or para-aminosalicylate, respectively. From 90-95 % of the intensity originates from Ca₂Al-LDH, when the contribution from the ²⁷Al satellite transitions (ca 5 %) to the center band is included (Table SI-1 and Fig. SI-2a). We note that a Lorentzian line broadening gave the best fit. Moreover, the spinning sidebands in the ²⁷Al MAS NMR spectrum had a clear indication of dynamics on the time scale of the 1st order quadrupole interaction (MHz) (Fig. SI-2) as earlier reported (Di Bitetto et al., 2017). It slightly differs from the value $\delta_{\text{iso}}(^{27}\text{Al}) = 9.5(4)$ ppm reported for Ca₂Al-Cl by Andersen et al. (Andersen et al., 2002). This small variation is due to a change of anion, as also seen for MgAl- and ZnAl-LDH (Pushparaj et al, 2015; Sideris et al., 2012; Sideris et al., 2008).

3.2. Adsorption isotherm of PO₄ by Ca₂Al-NO₃

In order to monitor the reactivity of Ca₂Al-NO₃ material with phosphate and to quantify the amount of sequestered PO₄, Ca₂Al-NO₃ was suspended at 20°C and for 2h00 with solutions containing increasing amounts of HPO₄²⁻ anions (section 2.2) and the PO₄ content adsorbed per gram of sorbent versus PO₄ equilibrium concentration (q_e vs c_e) was plotted (Fig. 2A). The isotherm plot is typical of an H-type adsorption isotherm curve according Giles's classification (Giles et al., 1974). This adsorption behavior clearly indicates a high affinity of PO₄ anions toward the Ca₂Al-LDH matrix. Both Langmuir and Freundlich models have been used to fit phosphate adsorption by LDH (Hatami et al., 2018; Wan et al., 2017). The Langmuir model assumes monolayer adsorption on a homogeneous surface with no interactions between the adsorbed molecules; while the Freundlich model is often used to describe chemisorption on heterogeneous surface (Chen et al., 2015). Both models were used to fit our experimental data.

The Langmuir model give the best linear fit (with $R^2 > 0.9980$) adsorption constants were determined from the slope and intercept of the plots (Fig.2B and Table SI-2). The experimental PO₄ adsorption capacity of Ca₂Al-NO₃ LDH is equal to 65.20 mgP.g⁻¹, a value higher than the one expected for a pure anion exchange reaction of HPO₄²⁻ (52.75 mgP.g⁻¹), 1.23 time higher, and much higher if we consider PO₄³⁻ exchange (35.17 mgP.g⁻¹, 1.85 time higher). Such a high PO₄ adsorption capacity was already reported in the literature (Jia et al., 2016, Lu et al., 2016) for Ca₂Al-Cl LDH (59.55 mg.g⁻¹). Nature of the intercalated anion (NO₃⁻ vs Cl⁻) does not affect the total PO₄ sequestration but seems to play a role on the mechanism of interaction (Langmuir vs Freundlich)..

It must be mentioned that in our case PO₄ adsorption experiments were performed at kinetic equilibrium and that no Ca²⁺ nor Al³⁺ ions were detected in the supernatants. Indeed Ca²⁺ release percentage compared to total Ca²⁺ amount was negligible, less than 2.82% at 0 mgP/L, and decreased to zero till 0.6 mgP/L (Fig. SI-3). Moreover, the amount of phosphate uptake by the Ca₂Al-NO₃ LDH are much lower than the values expected for a total conversion of Ca₂Al-NO₃ into hydroxyapatite (126.6 mg P.g⁻¹) or brushite (211.0 mgP.g⁻¹). Dissolution/precipitation is probably not total and the mechanism of transformation of CaAl LDH appears much more complex than described in the literature. Consequently, we performed XRD and ²⁷Al SSNMR on samples loaded with increasing amounts of PO₄ in order to follow structural change.

3.3. *Dynamic and Reactivity of PO₄ anions with Ca₂Al-NO₃ LDH*

3.3.1. *Ex-situ and in-situ XRD experiments.*

Ex-situ PXRD patterns (Fig. 3-A) were recorded for Ca₂Al-NO₃ LDH contacted with HPO₄²⁻ solutions under batch reaction conditions. HPO₄²⁻ anions react with the LDH structure

from a very low PO₄ loading (32.6 mg P.g⁻¹ LDH, Ca-0.3P). At 43.5 mg P.g⁻¹ LDH (Ca0.4P), (002) and (004) the most intense reflections of Ca₂Al-NO₃ pristine material have vanished meanwhile some new diffraction reflections appear that accounts for the formation of new phases. These new phases were identified from comparison of PXRD diffractograms of calcium phosphate phases known to appear during PO₄ reaction with hydrocalumite, Ca₅(PO₄)₃(OH) (hydroxyapatite; 09-0432 JCPDS file), CaHPO₄ (monetite; monoclinic, 09-0080 JCPDS file), Ca(HPO₄)₂·H₂O (Brushite; triclinic 09-0077 JCPDS file), octacalcium phosphate (Ca₈H₂(PO₄)₆·5H₂O (OCP) and Al(OH)₃ (Gibbsite; 33-0018 JCPDS file). Table SI-3 display the Bragg distances for the reflections of all Ca-xP samples and reference phases (Brushite, hydroxyapatite, gibbsite). A comparison of the PXRD patterns evidence the multiple reactivity of phosphate anions with Ca₂Al-NO₃ giving rise to a competition between exchange reactions into the Ca₂Al structure and precipitation of Ca/PO₄ phases.

First, the presence of the characteristic (002)/(110) diffraction doublet from Gibbsite (4.814 Å and 4.390 Å), appear at the lowest PO₄ loading (Ca-0.3P), which confirm dissolution of Ca₂Al-NO₃. However, from the point of view of the PXRD data, the amount of Gibbsite does not increase noticeably when increasing phosphate loading, suggesting that dissolution of the pristine layered structure is only partial. This dissolution is confirmed by the occurrence at higher PO₄ loading of some reflections from the mineral hydroxyapatite including the (002), (102), and the (121)/(211)/(112)/(300)/(202) reflections at 2θ = 25.90(3), 28.19(3)° and 31–35° 2θ respectively. Reflections assigned to the HAP phase are broad with a low intensity, due to particles with small sizes and low crystallinity. For Ca₂Al-Cl hydrocalumite, PO₄ adsorption leads to a collapse of the structure and the formation of hydroxyapatite following a dissolution/precipitation mechanism (Jeong-Woo et al. 2016). Formation of brushite is also suggested by the sharp

reflection at $29.42(3)^\circ$, whose intensity increases with PO₄ adsorption. This reflection is unambiguously assigned to the (-141) reflection from brushite.

Interestingly, the two intense reflections at 7.564 Å and 3.782 Å with a spacing ratio of 2.0 are observed for all samples. They cannot account for any known calcium phosphate phases but can be explained by a shift of the Ca₂Al-NO₃ (002) and (004) reflection to high 2θ values. Simultaneously intense carbonate vibration bands are present for all phases. This cannot account for the precipitation of calcium carbonate phase such as calcite reported by Jeong-Woo et al. (Jeong-Woo et al., 2016) as corresponding diffraction lines are not observed in the XRD patterns. This CaAl phase with a mean interlayer spacing or basal spacing (d_{bs}) of 7.564 Å must arise from the topotactic transformation of the pristine Ca₂Al-NO₃ LDH ($d_{bs} = 8.513$ Å) into a Ca₂Al-CO₃ LDH that display a shortest basal spacing (Renaudin et al., 1999; François et al., 1998). Contamination by CO₂ and conversion into CO₃²⁻ is favored by the basicity of the CaAl matrix. This reactivity can be explained by the high stability of the monocarboaluminate phase against both dissolution and phosphate reaction. Obviously phosphate anions cannot be exchanged and stabilized in the Ca₂Al galleries.

In conclusion, the XRD patterns of all samples clearly show a mixture of one or more following phase Ca₂Al-NO₃, Ca₂Al-CO₃, Al(OH)₃, Ca₅(PO₄)₃(OH) and Ca(HPO₄).2H₂O. These results indicate a different reactivity toward HPO₄²⁻ for Ca₂Al-NO₃ compared to Ca₂Al-Cl. Indeed, Zhou et al. (Zhou et al., 2012) reported that under PO₄ adsorption and for a similar range of pH Ca₂Al-Cl undergoes a complete transformation into a mixture of gibbsite, hydroxyapatite and tricalciumphosphate (Ca₃(PO₄)₂). When adsorption is performed at lower pH values using H₂PO₄⁻ aqueous solution again transformation of Ca₂Al-Cl is total. However, the resulting calcium phosphate formed deviate between the reported studies. For example, brushite is the

main calcium phosphate phase and gibbsite is converted into AlPO_4 for Jia et al. (Jia et al., 2016)) and Lu et al. (Lu et al., 2015) while Quian et al. (Quian et al., 2012) reported the conversion of hydrocalumite into HAP, CaCO_3 and $\text{Al}(\text{OH})_3$ at $\text{pH} = 5.0$

In-situ PXRD experiments were conducted on the $\text{Ca}_2\text{Al-NO}_3$ sample, previously wetted as a paste (solid:water mass ratio = 2:1), loaded in a home-made diffraction cell (Fig. SI-4). The solid was in contact with a 2M HPO_4^{2-} solution (Fig. 3-B) in order to force the exchange by HPO_4^{2-} and minimize the dissolution reaction. Under such conditions of in-situ XRD exchange reaction, the high solid/liquid ratio must limit the dissolution of the NO_3 -hydrocalumite and prevent the precipitation of Ca- PO_4 phase. The designed cell insures a fast PO_4 diffusion to the LDH adsorption sites. A topotactic phase transformation of the pristine material occurs in less than 14 min as shown by the in-situ XRD pattern (Fig. 3-B). A group of three new diffraction lines (10.64(5) Å, 5.28(2) Å, and 3.50(1) Å), with integer spacing ratio ($L1/L2 = 2.02$; $L1/L3 : 3.04$), clearly accounts for the formation of a layered phase with a basal spacing mean value of 10.57(5) Å. This 1.957 Å increase of the interlayer distance was also observed when $\text{Ca}_2\text{Al-NO}_3$ was contacted with pure water. This structural expansion is reversible under hydration/dehydration process. Thus, this well-defined structural expansion arises from the hydration of the interlayer and the intercalation of a single layer of water molecules but has only previously been observed for $\text{Ca}_2\text{Al-NO}_3$ phase. The fast hydration reaction is probably a key step in the dissolution and phosphate reaction of the NO_3 -hydrocalumite as H_2O molecules can diffuse easily in the interlayer domains. While NO_3 -containing LDH are known to be the most reactive LDH phases towards anion exchange reactions and particularly with orthophosphate anions, condensation of HPO_4^{2-} with the LDH layer occurs prior to exchange preventing the formations of CaAl- PO_4 phase.

3.3.2. FTIR and Raman analysis

The strong structural changes undergone by the $\text{Ca}_2\text{Al-NO}_3$ phase upon reaction with phosphate, evidenced by XRD analysis, are clearly confirmed by analysis of the Raman (Fig. 4) and FTIR (Fig. SI-5) spectra. First of all, PO_4 the vibration bands (Table 2) related to PO_4 groups increase during PO_4 adsorption as shown on both FTIR and Raman spectra. Loading of HPO_4^{2-} leads to the removal of NO_3^- anions which is linked to the decrease of the $\nu_3(\text{NO}_3)$ stretching band intensity (1055 cm^{-1}). The formation of $\text{Ca}_2\text{Al-CO}_3$ LDH, as suggested by the XRD data (section 3.3.1), is confirmed by the appearance of a broad and intense $\nu_3(\text{CO}_3)$ multiplet band ($1350\text{-}1550\text{ cm}^{-1}$) (Fig. SI-5). The large feature of the CO_3 vibration bands, unusual for LDH material, must account for the presence of other carbonated phase such as carbonated substituted HAP (Gibson et al., 2002). At low frequencies, the symmetric (ν_2) (433 and 450 cm^{-1}) and asymmetric (ν_4) (582 , 593 , 609 cm^{-1}) bending modes of the PO_4^{3-} anions (Fig. SI-5) overlap with the lattice vibrations $\nu_{\text{Al(OH)}_6}$ of gibbsite (534 cm^{-1}) and hydrocalumite-like phases (580 , 528 , 425 cm^{-1}). The splitting of the ν_4 PO_4 vibration band into 3 bands accounts for the multiplicity of environment in the HAP, brushite and substituted- CO_3 HAP structure as reported by the literature (Swamiappan et al., 2016). Resolution of several MO-H vibration feature increases from Ca-03P to Ca-09P, due to a clear observation of the ν_{as} and ν_{s} vibrations of Al(OH)_3 Gibbsite (3470 , 3527 and 3619 cm^{-1}) superimposed on the HAP bands (3527 and 3619 cm^{-1} , $3100 - 3400\text{ cm}^{-1}$).

3.3.3 ^{27}Al NMR and ^{31}P MAS NMR characterizations

^{27}Al and ^{31}P SSNMR were used to investigate the structural changes of $\text{Ca}_2\text{Al-NO}_3$ during reaction with HPO_4^{2-} and to quantify the relative concentration of the different phosphate phases formed. The ^{27}Al MAS NMR spectra (Fig. 5) all show one resonance in the octahedral range in agreement with the coordination of Al in the LDH structure. In addition, a small amount of tetrahedral Al is seen at $\delta_{\text{iso}}(^{27}\text{Al}) \approx 73$ ppm, constituting less than 5% in all the ^{27}Al NMR spectra except for the $\text{Ca}_2\text{Al-NO}_3$ with 0 and 0.9P loading. Upon phosphate exposure of 0.3P an upfield shoulder to the LDH resonance assigned to AOH appears (Jensen et al., 2016); Pushparaj et al., 2015). Different models for simulation of the 0.3P ^{27}Al MAS NMR spectrum has been applied but an unambiguous model could not be identified. However, it was estimated that 15-30 % of the Al belonged to the LDH. Furthermore, a slight shift in $\delta_{\text{iso}}(^{27}\text{Al})$ in the LDH resonance from 11.0(4) to 11.3(4) ppm was observed indicating a change in the intercalated anion, e.g., carbonate as seen by PXRD (Fig. 3-A). At higher phosphate loadings the AOH phase is clearly the dominating phase and consequently the center of the signal is located below 10 ppm. Moreover, the total width of the spinning side band manifold (not shown) changes from ≈ 400 kHz to ≈ 600 kHz implying a 50 % increase of the quadrupole interaction of the predominant ^{27}Al resonance. Thus, at high P loading the predominant Al species is AOH, which has a stronger quadrupole interaction (2.5 MHz) than $\text{Ca}_2\text{Al-LDH}$ (1.6 MHz) (Jensen et al., 2016), c.f., Table SI-2. However, $\text{Ca}_2\text{Al-LDH}$ might still be present with relative Al concentration up to 10-15 % but not distinguished due to the overlap between the LDH and AOH resonances. Hence, the ^{27}Al MAS NMR confirms dissolution of the LDH and precipitation of aluminum hydroxide (AOH) upon phosphate exposure.

The ^{31}P MAS NMR is a very sensitive probe of the phosphorus speciation and it is therefore possible to distinguish between e.g., different mineral phases.

The ^{31}P MAS NMR spectra for the $\text{Ca}_2\text{Al-NO}_3$ all show one resonance at 3 ppm with one set of weak spinning sideband which is indicative of orthophosphate in agreement with earlier studies (Bernardo et al., 2017). The resonance can be adequately fitted with four resonances as seen in the deconvolutions in Fig. 6 and the obtained parameters from deconvolutions are summarized in Table 3. The dominating resonance at $\delta_{\text{iso}}(^{31}\text{P}) = 3.0(3)$ ppm constitutes 71-79 % of the phosphates species and is based on the isotropic shift assigned to hydroxyapatite ($\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$) (Aue et al., 1984; Rothwell et al., 1980). This is in agreement with the PXRD, where formation of $\text{Ca}_5(\text{PO}_4)_3(\text{OH})$ (HAP) is clearly observed. The small resonance at $\delta_{\text{iso}}(^{31}\text{P}) = 1.4-1.7$ ppm is assigned to brushite constituting 6-9% of the sample (Aue et al., 1984; Rothwell et al., 1980). The two resonances at $\delta_{\text{iso}}(^{31}\text{P}) \approx 4.3-4.5$ and $5.6-6.0$ ppm constitute a total of 16-21 % of the samples. The latter resonance ($\delta_{\text{iso}}(^{31}\text{P}) = 5.6-6.0$ ppm) agrees well with a dehydrated oxyhydroxyapatite, $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_{2-2x}\text{O}_x$, which may precipitate during phosphate adsorption experiments. Thus, that ^{31}P NMR confirms that the CaAl-NO_3 LDH degrades during the adsorption experiments and mainly hydroxyapatite is formed, which are in agreement with earlier studies for Cl-hydrocalumite (Gibson et al., 2002; Swamiappan et al., 2016; Renaudin et al., 1999).

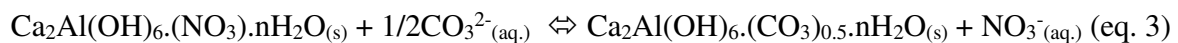
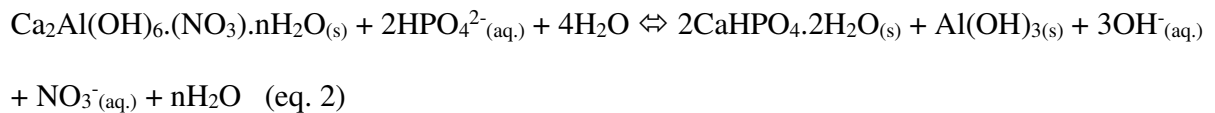
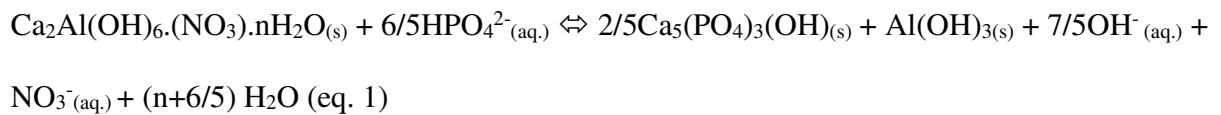
3.3.4 Morphological evolution under PO_4 adsorption

Adsorption of HPO_4^{2-} causes strong changes of the $\text{Ca}_2\text{Al-NO}_3$ morphology as shown by SEM analysis (Fig. 7). Indeed, the large hexagonal platelets of the pristine material (Fig.7-A),

with sizes in the range 1-2 μm , undergo a partial dissolution followed by a nucleation/crystallization process occurring at the surface of CaAl-NO_3 particles. Fig.7-B,C and D display these small HAP nanocrystals (size < 100nm) with needle shape homogeneously dispersed all over the solid. Moreover, at 0.6P loading (Fig 7-C) well defined crystals are clearly observed whose parallelepiped forms look like that of the brushite crystals (Onac et al., 2009). It is clear that some of the large hexagonal platelets are still present at high PO_4 loading. These observations confirms the hypothesis of formation of $\text{Ca}_2\text{Al-CO}_3$, $\text{Ca}_5(\text{PO}_4)_3(\text{OH},\text{CO}_3)$ and $\text{Ca}(\text{HPO}_4)\cdot 2\text{H}_2\text{O}$ (Fig. 7D).

3.4 Mechanism of HPO_4^{2-} reactivity with $\text{Ca}_2\text{Al-NO}_3$ LDH

According results from combined PXRD, ^{27}Al and ^{31}P SSNMR, Raman, FTIR and SEM analysis, HPO_4^{2-} anions react spontaneously with $\text{Ca}_2\text{Al-NO}_3$ LDH in solution. While PO_4 anion exchange was reported into Zn_2Al and Mg_2Al LDH to give respectively $\text{Zn}_2\text{Al-HPO}_4^{2-}$ (Stefel et al., 2018) and $\text{Mg}_2\text{Al-HPO}_4^{2-}$ (Everaert et al., 2016), $\text{Ca}_2\text{Al-HPO}_4$ LDH was not formed, indicating that if HPO_4^{2-} anions intercalates in between the hydrocalumite layers they react with the layer to destroy the structure. Reaction proceeds according the competing formation of HAP (eq. 1), brushite (eq.2) and $\text{Ca}_2\text{Al-CO}_3$ (eq.3):



This result is supported by the adsorption data (section 3.2) that indicates an adsorption capacity greater than expected for a pure anion exchange reaction. However the Langmuir adsorption behavior does not evidence this multiple reactivity because both brushite and HAP formations are strongly thermodynamically favorable.

Unlike most of the data published on hydrocalumite/phosphate reactivity that evidence the conversion of $\text{Ca}_2\text{Al-Cl}$ into hydroxyapatite, our results show that i) brushite is also formed and ii) part of Ca_2Al LDH remains due to a stabilization by CO_3^{2-} intercalation.

A phosphate concentration of only 400 P mg.L^{-1} is sufficient for the $\text{Ca}_2\text{Al-NO}_3$ structure to collapse, i.e. a $\text{HPO}_4^{2-}/\text{Ca}^{2+}$ molar ratio of 0.189 while for a total conversion of hydrocalumite into brushite or hydroxyapatite $\text{HPO}_4^{2-}/\text{Ca}^{2+}$ molar ratio of 1.0 and 0.6 are needed. At maximum adsorption, 48.0% of $\text{Ca}_2\text{Al-NO}_3$ were converted into calcium phosphate phases, HAP ($\text{Ca}_5(\text{PO}_4)_3(\text{OH})$) and brushite ($\text{Ca}(\text{HPO}_4)\cdot 2\text{H}_2\text{O}$). Obviously, when $\text{Ca}_2\text{Al-NO}_3$ is contacted with a solution of HPO_4^{2-} salt, HPO_4^{2-} diffuse easily inside the layered structure up to the Ca^{2+} reactive sites thanks to a fast hydration of the structure (in-situ PXRD) that increases the interlayer distance and opened the structure for PO_4 diffusion. The various phosphorus environments shown by ^{31}P NMR indicate that HPO_4^{2-} reaction probably occurs at the Ca_2Al layer surface through an initial complexation of Ca^{2+} ions assisted by both the expulsion of the labile coordinated water and the hydrolysis of Ca-OH bonds. Clearly the corrugated Ca_2Al layers do not stabilize enough the Ca^{2+} ions inside the hepta-coordinated sites. Ca^{2+} ions can easily diffuse out of the layers. Chemical extraction of Ca^{2+} ions and formation of hydrated $[\text{Ca-PO}_4]$ complexes are the preliminary steps of condensation of Ca-PO_4 phases (Fig. 8). Costless energy surface nucleation is then favored. Competition occurs between precipitations of hydroxyapatite and brushite minerals. Even if HAP ($\text{PO}_4^{3-}/\text{Ca}^{2+} = 3/5$) must be preferentially favored when

surface $\text{PO}_4^{3-}/\text{Ca}^{2+}$ ratio increases Brushite ($\text{PO}_4^{3-}/\text{Ca}^{2+} = 1/1$) was also forms as a secondary phase even at such high pH ($\frac{pK_2+pK_3}{2} = 9.94$). The presence of carbonate in the medium may quenched the formation of calcium phosphate phase, resulting in the stabilization of the CO_3 -hydrocalumite like structure (PXRD, FTIR and Raman data). Interestingly, as shown by scanning electron microscopy, brushite and HAP crystallites with a wide distribution of particle size from nanometer to micrometer are dispersed all over the remaining CO_3 -hydrocalumite phase. Such composite material with embedded Ca- PO_4 phases could give promising result for slow-released phosphate fertilizers. The recycling and reuse of phosphorus could therefore be achieved.

4. Conclusion

This study highlights the specific reactivity of $\text{Ca}_2\text{Al-NO}_3$ hydrocalumite-like LDH toward phosphate sequestration. Up to $65.20 \text{ mgP}\cdot\text{g}^{-1}$ was loaded into the resulting composite, values greater than the anion exchange capacity of $\text{Ca}_2\text{Al-NO}_3$. A mixture of brushite, HAP and $\text{Ca}_2\text{Al-CO}_3$ LDH phases was obtained allowing a distribution of PO_4 structures and morphologies. As HAP (Giroto et al., 2015; Xiong et al., 2018) and Brushite (Hermassi et al., 2017) have been investigated as potential slow release fertilisers, HAP and brushite nanoparticle embedded inside a Ca_2Al matrix may offer promising PO_4 slow release properties.

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Tables and Figures

Figure legends

Fig. 1: (A) XRD pattern and (B) FTIR spectrum of $\text{Ca}_2\text{Al-NO}_3$.

Fig. 2: A) Experimental and calculated adsorption isotherms of $\text{Ca}_2\text{Al-NO}_3/\text{HPO}_4^{2-}$ system. B) Insert linear form of the Langmuir fit.

Fig.3: A) ex-situ XRD of $\text{Ca}_2\text{Al-NO}_3/\text{HPO}_4^{2-}$ series for various HPO_4^{2-} loading at 0; 0.3; 0.4; 0.5; 0.6; 0.7; 0.8; 0.9 g/L (∇ $\text{Ca}_2\text{Al-NO}_3$; \blacktriangle $\text{Ca}_2\text{Al-CO}_3$; \diamond Al(OH)_3 Gibbsite; \blacklozenge $\text{Ca}_5(\text{PO}_4)_3(\text{OH})$ Hydroxyapatite; \star $\text{Ca}(\text{HPO}_4)\cdot 2\text{H}_2\text{O}$ Brushite) and B) in-situ XRD of $\text{Ca}_2\text{Al-NO}_3$ in contact with a 2M HPO_4^{2-} solution.

Fig. 4: Raman spectra of the $\text{Ca}_2\text{Al-NO}_3/\text{HPO}_4^{2-}$ series with increasing HPO_4^{2-} loading (from 0.3 to 0.9 g/L) at A) medium and B) low energy range.

Figure 5: The ^{27}Al MAS NMR spectra of the LDH exposed to the different phosphate concentrations. The dotted line shows the position of resonance for $\text{Ca}_2\text{Al-NO}_3$ LDH.

Fig. 6: Deconvolutions of the ^{31}P MAS NMR spectra of the $\text{Ca}_2\text{Al-NO}_3$ LDH with different phosphate loadings a-g): $\text{Ca}_2\text{Al-NO}_3$ with 0.3-0.9P, h): hydroxyapatite and j) brushite, two resonances from impurities are observed at 0.1 and -1.4 ppm. The top line is the experimental spectrum (Exp), the second is the simulated (Sim) and the bottom line is the residual (Dif).

Fig. 7: SEM images of A) pristine $\text{Ca}_2\text{Al-NO}_3$ material and loaded with various amounts of PO_4 , B) $\text{Ca}0.3\text{P}$, C) $\text{Ca}0.6\text{P}$, D) $\text{Ca}0.9\text{P}$.

Fig. 8: Initial step of the conversion of $\text{Ca}_2\text{Al-NO}_3$ LDH into calcium phosphate minerals.

Fig. 2

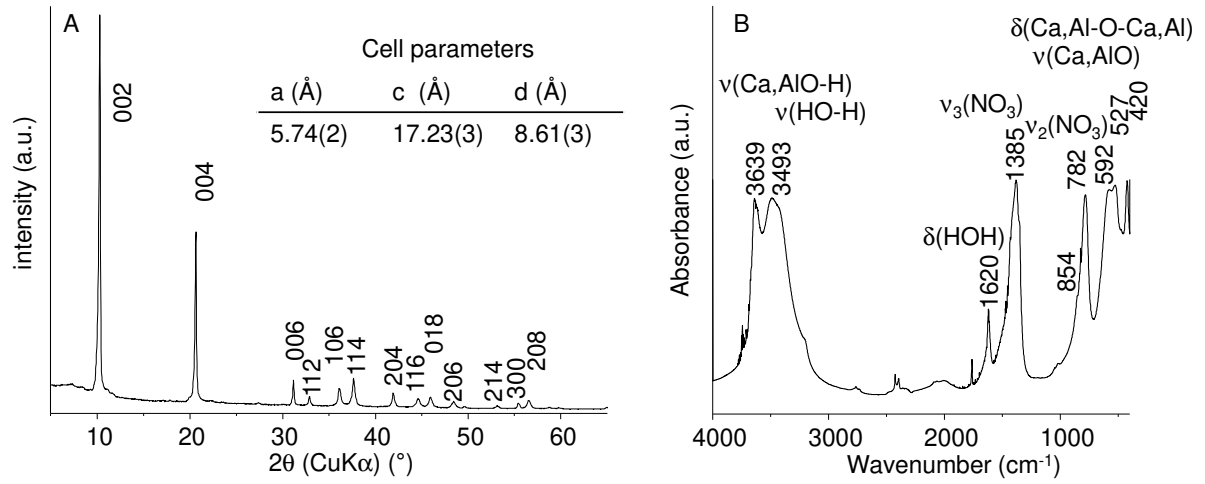


Fig. 2

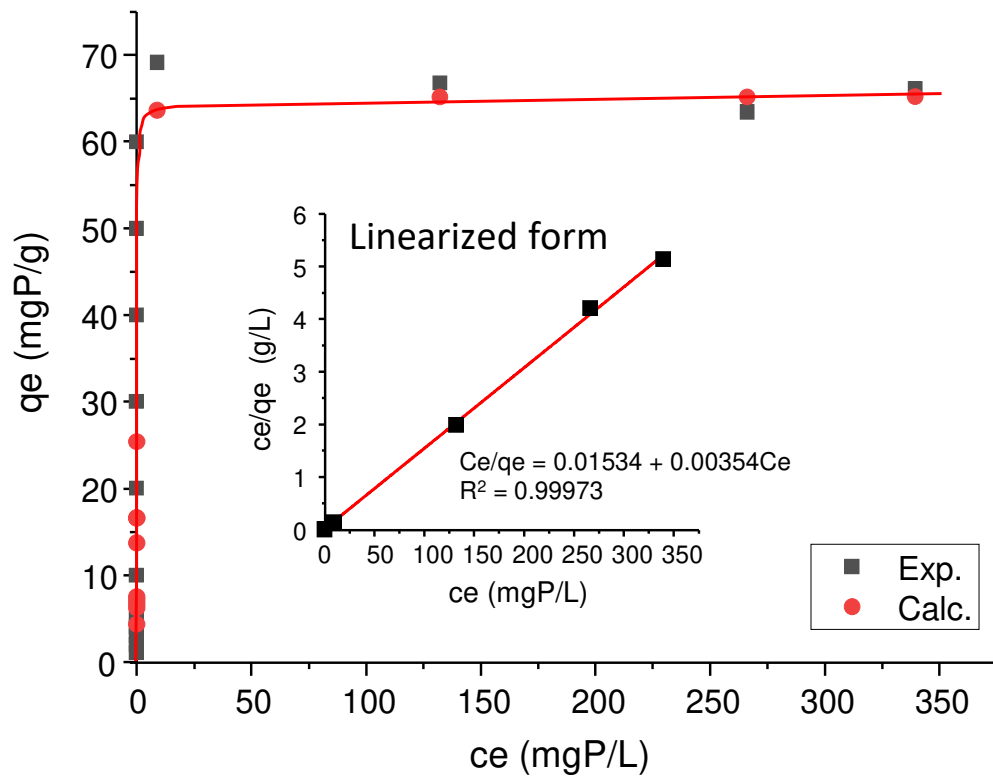


Fig. 3

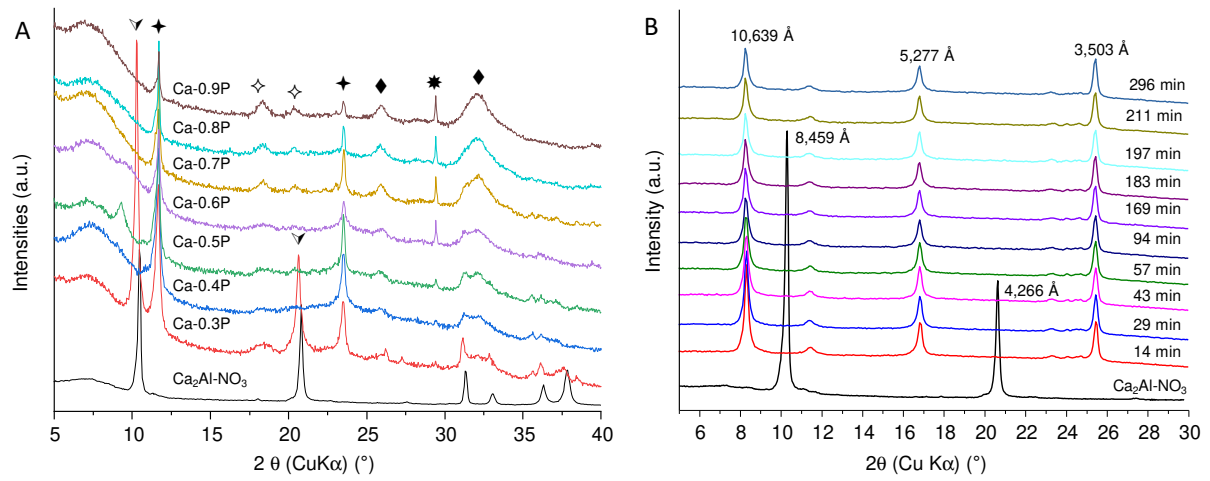


Fig. 4

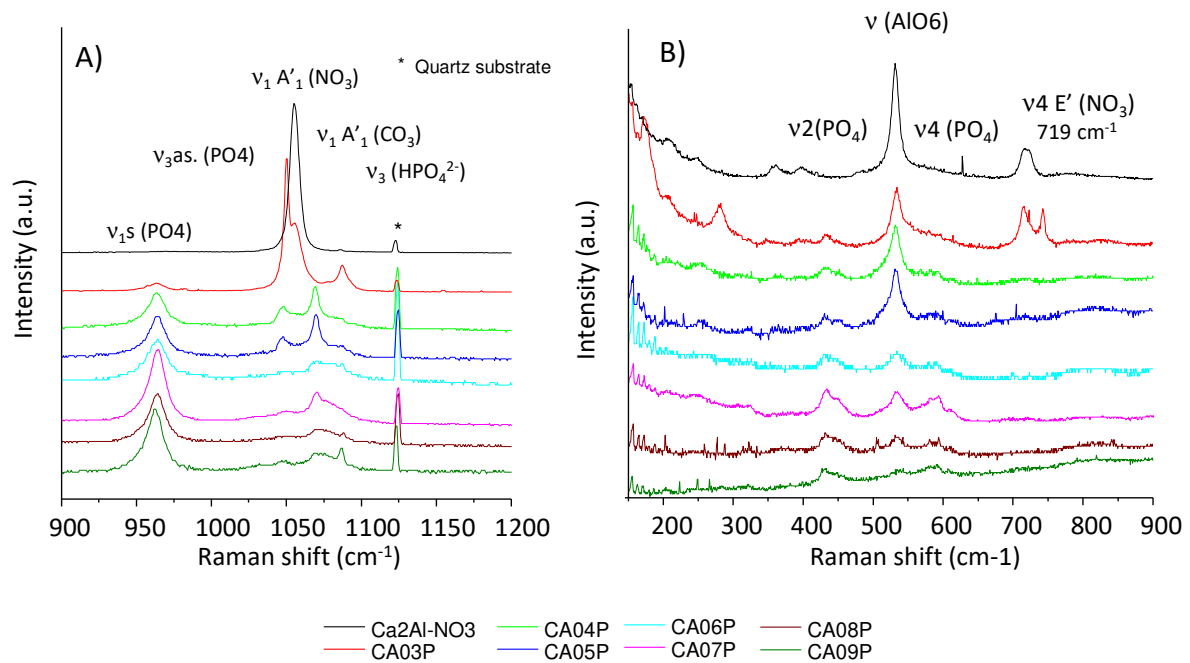


Fig. 5

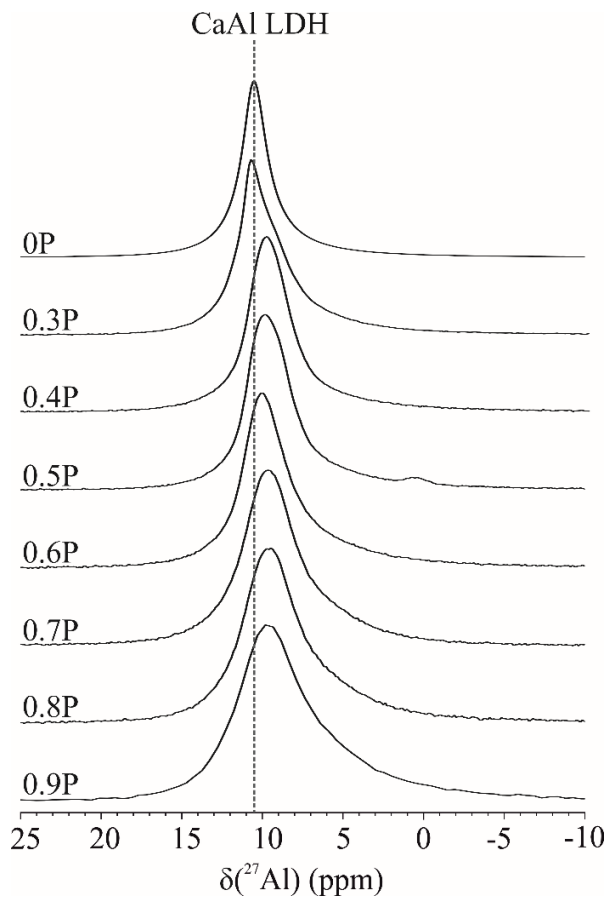


Fig. 6

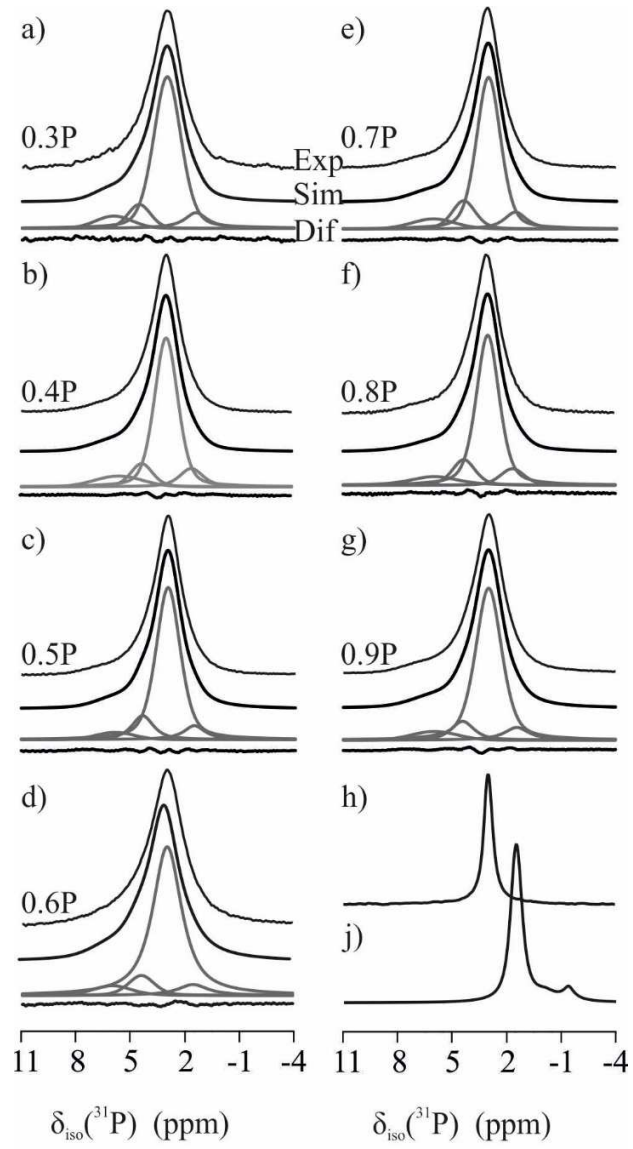


Fig. 7

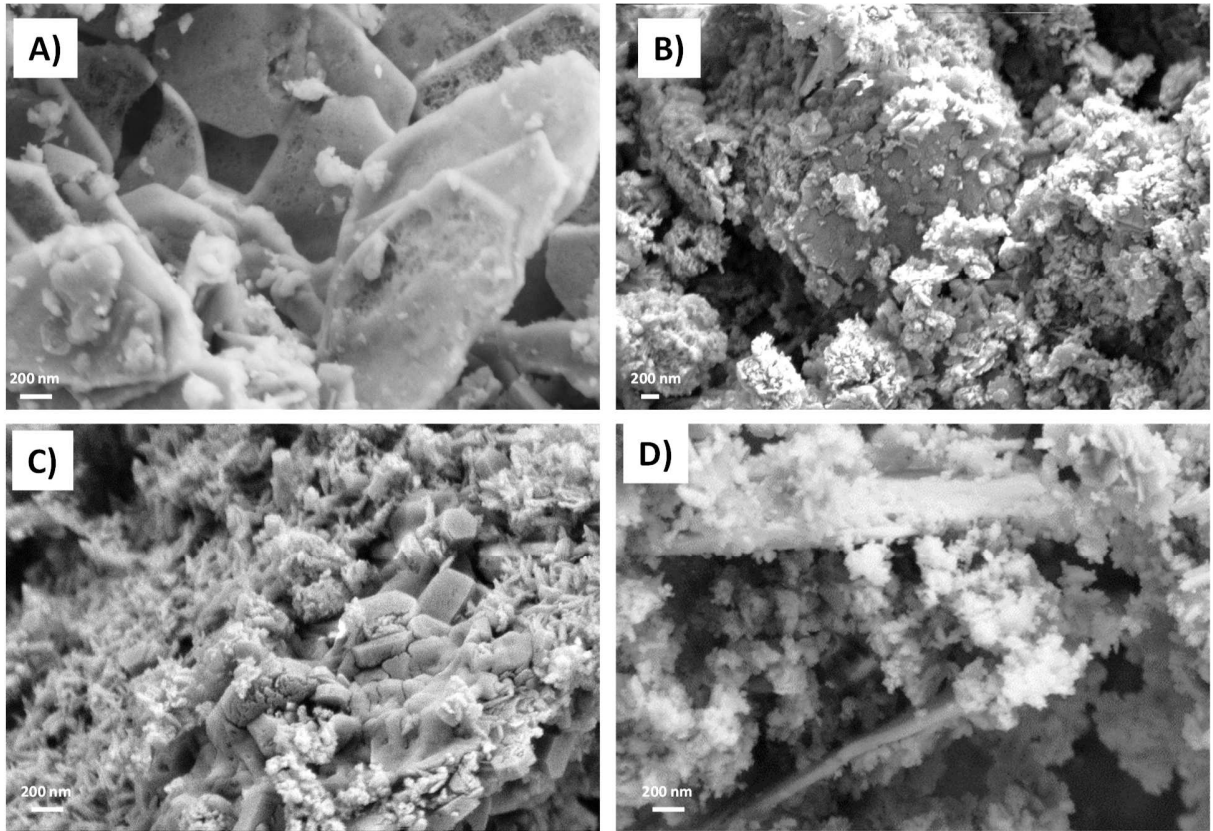
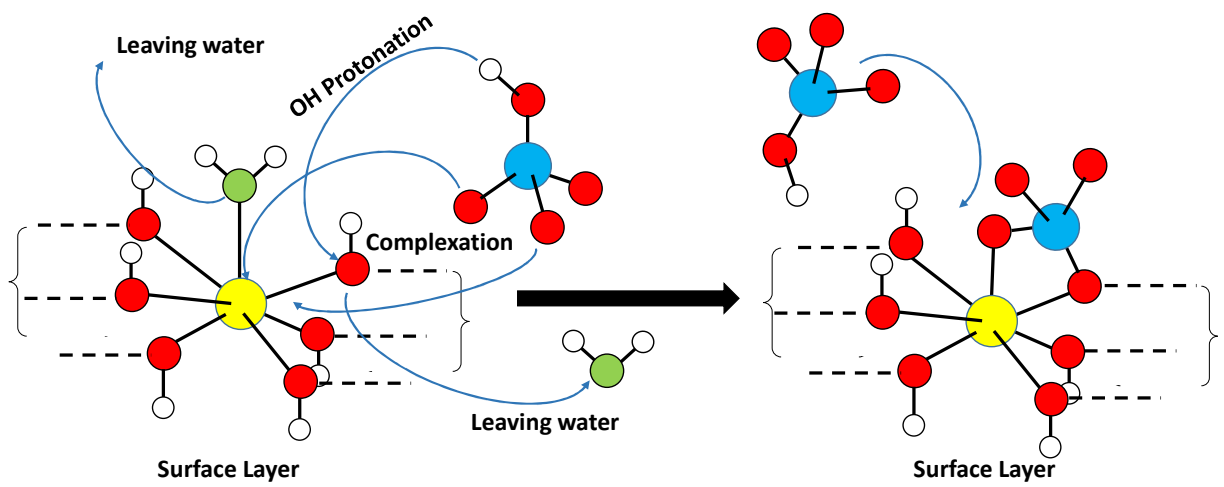


Fig. 8



Tables and Figures

Table legends

Table 1: Solubility products of some calcium phosphate compounds.

Table 2: Assignment of PO₄ infrared and Raman vibration bands.

Table 3: Parameters obtained from the deconvolutions of the ³¹P MAS NMR spectra for Ca₂Al-NO₃.

Table 1

Calcium phosphate phases	pK _{sp}	Ref.
Ca ₅ (PO) ₄ (OH)	-58.4	(Bell et al., 1978)
Ca ₃ (PO ₄) ₂	-25.5	(Fowler et al., 1986)
Ca ₈ H ₂ (PO ₄) ₆ .5H ₂ O	-96.6	(Tung et al., 1988)
CaHPO ₄ .2H ₂ O	-6.59	(Gregory et al., 1970)
Ca ₂ Al(OH) ₆ (OH).2H ₂ O	-11.4	(Gacsi et al., 2016)

Table 2

Vibrations (cm ⁻¹)	Infrared	Raman
ν ₃ (PO ₄) _{as}	1025, 1063, 1105, 1138	1070; 1081
ν ₁ (PO ₄) _s		962; 1050

Table 3:

Sample	Assignment	$\delta_{\text{iso}}(^{31}\text{P})(\text{ppm})$	I(%)	
	Brushite	1.4(5)	8(5)	
	Hydroxyapatite	3.0(3)	75(6)	
		4.5(6)	10(5)	
Ca ₂ Al-NO ₃ 0.3P	Oxyhydroxyapatite phase	6.0(7)	8(5)	
	Brushite	1.6(5)	9(5)	
	Hydroxyapatite	3.0(3)	71(6)	
		4.3(6)	10(5)	
	Ca ₂ Al-NO ₃ 0.4P	Oxyhydroxyapatite phase	5.6(7)	10(5)
		Brushite	1.5(5)	7(6)
Hydroxyapatite		3.0(3)	77(8)	
		4.3(6)	11(6)	
	Ca ₂ Al-NO ₃ 0.5P	Oxyhydroxyapatite phase	5.9(7)	5(6)
		Brushite	1.5(5)	6(5)
Hydroxyapatite		3.0(3)	79(6)	
		4.4(6)	9(5)	
	Ca ₂ Al-NO ₃ 0.6P	Oxyhydroxyapatite phase	6.0(7)	6(5)
		Brushite	1.5(5)	8(5)
Hydroxyapatite		3.0(3)	71(6)	
		4.4(6)	13(5)	
	Ca ₂ Al-NO ₃ 0.7P	Oxyhydroxyapatite phase	6.0(7)	8(5)
		Brushite	1.7(5)	8(5)
Hydroxyapatite		3.0(3)	71(6)	
		4.4(6)	12(5)	
	Ca ₂ Al-NO ₃ 0.8P	Oxyhydroxyapatite phase	6.0(7)	9(5)
		Brushite	1.4(5)	7(5)
Hydroxyapatite		3.0(3)	77(6)	
		4.4(6)	9(5)	
	Ca ₂ Al-NO ₃ 0.9P	Oxyhydroxyapatite phase	6.0(7)	7(6)

The dominating resonance at $\delta_{\text{iso}}(^{31}\text{P}) = 3.0(3)$ ppm constitutes 71-79 % of the phosphates species and is based on the isotropic shift assigned to hydroxyapatite

Graphical abstract

