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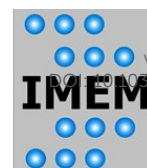


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Department of Chemical Engineering

## Sustainability Spotlight

The catalytic conversion of carbon dioxide (CO<sub>2</sub>) into chemical added-value products (CO<sub>2</sub>-revolarization) is interesting because of its potential to close the carbon cycle while generating green circular economies. However, truly sustainable CO<sub>2</sub>-revolarization processes are normally restricted by the use of metallic-noble-based catalysts and/or intensive reaction conditions. Accordingly, we explore the use of ultraporous permanently polarized hydroxyapatite (upp-HAp) as a catalyst for green CO<sub>2</sub>-revalorization. upp-HAp is abundant and biocompatible (main inorganic component of biological hard tissues). Besides, upp-HAp is capable of performing successful reactions under mild conditions (achievable through renewable sources) and in batch/continuous configurations, representing a viable alternative to conventional catalysts. This work is aligned with the SDG7 (affordable and clean energy) and SDG13 (climate action) UN sustainable development goals.

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RSC Sustainability Accepted Manuscript

## ARTICLE

# Establishing Ultraporous Permanently Polarized Hydroxyapatite as Green and Highly Efficient Catalyst for Carbon Dioxide Conversion in Continuous-Flow under Mild Conditions

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Marc Arnau,<sup>a</sup> Jordi Sans,<sup>\*a</sup> Pau Turon<sup>\*b</sup> and Carlos Alemán<sup>\*ac</sup>

We present the use of ultraporous permanently polarized hydroxyapatite (upp-HAp) catalyst for continuous and highly efficient production of formic acid (predominant) and acetic acid using wet CO<sub>2</sub> (i.e. CO<sub>2</sub> bubbled in liquid water) as reagent. In all cases, reactions were conducted at temperatures ranging from 95 to 150 °C, using a CO<sub>2</sub> constant flow of 100 mL/s, and without applying any external electrical field and/or UV radiation. Herein, we study how to transfer such catalytic system from batch to continuous reactions, focusing on the water supply (proton source): 1) wet CO<sub>2</sub> or; 2) liquid water in small amounts is introduced in the reactor. In general, the reduction of CO<sub>2</sub> to formic acid predominates over the C–C bond formation reaction. On the other hand, when liquid water is added, two interesting outcomes are observed: 1) the yield of products is higher than in the first scenario (> 2 mmol/g<sub>c</sub>-min) while the initial liquid water remains largely available due to the mild reaction temperature (95 °C); and 2) a high yield of ethanol (> 0.5 mmol/g<sub>c</sub>-min) is observed at 120 °C, as a result of the increased efficiency of the C–C bond formation. Analyses of kinetic studies through temporal and temperature dependence show that CO<sub>2</sub> fixation is the rate limiting step, discarding the competing effect of proton adsorption on the binding sites and confirming the crucial role of water. The activation energy for the CO<sub>2</sub> fixation reaction has been determined to be 66 ± 1 kJ/mol, which is within the range of conventional electro-assisted catalysts. Finally, mechanistic insights on the CO<sub>2</sub> activation and role of the binding sites of upp-HAp are provided through isotopic-labeled (<sup>13</sup>CO<sub>2</sub>) and near-ambient pressure X-ray photoelectron spectroscopy (NAP-XPS) studies.

## Introduction

The concentration of carbon dioxide (CO<sub>2</sub>), the most significant greenhouse gas in the atmosphere, has significantly increased in recent years mainly due to the combustion of fossil fuels.<sup>1-4</sup> Such environmental impact has accelerated research activities to capture, avoid or minimize the CO<sub>2</sub> release to the atmosphere. Two research approaches have been prioritized looking for net zero emissions: 1) CO<sub>2</sub> capture and long term geological storage or reutilization in several innovative ways;<sup>5-10</sup> and 2) catalytic conversion of CO<sub>2</sub> into added-value compounds of chemical and industrial interest using efficient catalysts and low energy consuming processes (i.e. CO<sub>2</sub> reutilization and valorization).<sup>11-19</sup>

Although many noble metal-based catalysts have been investigated for CO<sub>2</sub> conversion into chemicals of industrial

interest (e.g. Rh, Ru, Pt, Au and Pd),<sup>15,20-31</sup> simple systems based on non-precious metal catalysts are preferred due to their lower cost and higher sustainability (e.g. Fe, Cu, Zn, Co among others).<sup>32-40</sup> Similarly, although efficient catalytic processes have been developed to transform CO<sub>2</sub> into valuable chemicals using harsh reaction conditions (i.e. high temperature and/or pressure), processes involving mild reaction conditions are the most desired to avoid contradictions related to energy mass balance inefficiencies, as for example that the energy consumption and CO<sub>2</sub> emissions associated with the elimination of this greenhouse gas is higher than the CO<sub>2</sub> transformed.<sup>41-45</sup> A huge amount of studies have been devoted to the catalytic conversion of CO<sub>2</sub> into organic products (e.g. alcohols and acids with one or more carbon atoms), the main results being compiled in recent reviews.<sup>28,46-54</sup>

In a recent study we proposed a polarized bioceramic, a sustainable metal-free catalyst, for the conversion of CO<sub>2</sub> into ethanol using mild reaction conditions.<sup>55</sup> Such catalyst consisted of hydroxyapatite (HAp), a naturally occurring mineral with the formula Ca<sub>5</sub>(PO<sub>4</sub>)<sub>3</sub>(OH), that is non-toxic and biocompatible and, therefore, frequently used for biomedical applications.<sup>56-59</sup> HAp was permanently polarized by applying a thermal stimulated polarization (TSP) process at high temperature. The resulting material (named pp-HAp) became catalytically active due to extraordinary charge accumulation

<sup>a</sup> Departament d'Enginyeria Química, EEBE and Barcelona Research Center in Multiscale Science and Engineering, Universitat Politècnica de Catalunya, C/ Eduard Maristany, 10-14, Ed. I2, 08019, Barcelona, Spain.

<sup>b</sup> B. Braun Surgical, S. A. U. Carretera de Terrassa 121, 08191, Rubí (Barcelona), Spain.

<sup>c</sup> Institute for Bioengineering of Catalonia (IBEC), The Barcelona Institute of Science and Technology, Baldri Reixac, 10-12, 08028, Barcelona, Spain.

† Footnotes relating to the title and/or authors should appear here.

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on its surface. Thus, pp-HAp was found to exhibit not only vacancies and charge accumulation at the HAp crystal lattice but also charge accumulation at the boundaries among the microscopic HAp crystal grains.<sup>60,61</sup> Compact pp-HAp discs, which were prepared by applying a pressure of 620 MPa to HAp powder before applying the TSP, were found to convert CO<sub>2</sub> into a mixture of ethanol (predominant) and some other minority compounds (*i.e.* acetic acid, formic acid, methanol and acetone) using a starting mixture of CO<sub>2</sub> and CH<sub>4</sub> (3 bar each) and 1 mL of liquid water.<sup>55</sup> However, the yield of the reaction (expressed in  $\mu\text{mol}$  per gram of catalyst;  $\mu\text{mol/g}_c$ ), which was performed in a batch reactor at 140 °C, was very low after 72 h:  $18.6 \pm 1.9$ ,  $2.7 \pm 0.6$ ,  $1.9 \pm 0.6$ ,  $1.5 \pm 0.3$  and  $0.9 \pm 0.2$   $\mu\text{mol/g}_c$  for ethanol, acetic acid, formic acid, methanol, acetone, respectively.

More recently, the performance of pp-HAp catalyst was significantly improved by modifying the synthesis process to provide ultra-porosity and printability.<sup>62</sup> Thus, printable inks were prepared by mixing HAp powder with Pluronic® F-127 hydrogel. The latter was removed after 3D printing the scaffold (but before applying the TSP process) by calcining the samples at 1000 °C. The resulting catalyst exhibited spherical HAp nanoparticles with an average diameter of  $159 \pm 43$  nm. More specifically, such printed catalyst consisted of a 3D network forming branches made of coalesced nanoparticles separated by nanometric pores (average size:  $167 \pm 51$  nm). The performance of the ultraporous pp-HAp catalyst (hereafter named upp-HAp) for the production of ethanol was studied by loading a CO<sub>2</sub> : CH<sub>4</sub> mixture (3 bar each) and 1 mL of liquid water into the batch reactor and applying 140 °C for 48 h. The ethanol yield increased up to  $55.0 \pm 4.9$   $\mu\text{mol/g}_c$ , which represented an outstanding increment of the catalytic performance (by a factor of 2.9).

Despite that, most of the studies reported focus on the enhanced electric properties of the material, rather than elucidating the catalytic mechanisms involved. In this work, we go one step further by transferring the use of upp-HAp catalyst from the batch reactor to a continuous process where the kinetics of the conversion of CO<sub>2</sub> into ethanol have been studied. Because of the extremely high yields obtained, we have focused on the catalytic activity of upp-HAp to settle down, for the first time, the fundamentals of upp-HAp as a highly efficient and sustainable catalyst. For this purpose, first we have examined the formation of products in the different phases (*i.e.* adsorbed on the catalyst surface, liquid water and gas phase, as this is a heterogeneous catalytic process) as a function of the time using a batch process. Next, the influence of water, which is rapidly consumed in the continuous process, and how is supplied have been examined (*i.e.* wet CO<sub>2</sub> or liquid water). Third, the catalytic performance of upp-HAp has been studied by varying the temperature and reaction time to determine the kinetics of the reaction. Finally, isotope-labeled <sup>13</sup>CO<sub>2</sub> reactions and near ambient pressure X-ray photoelectron spectroscopy (NAP-XPS) studies have been performed to support the results presented. Accordingly, the catalytic mechanism of CO<sub>2</sub> activation has been experimentally

determined further establishing upp-HAp as a green catalyst for CO<sub>2</sub> fixation.

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## Experimental Methods

### Synthesis of ultraporous pp-HAp catalyst

**Hydroxyapatite (HAp).** 15 mL of 0.5 M (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub> in de-ionized water were added at a constant rate of 2 mL/min to 25 mL of a 0.5 M of Ca(NO<sub>3</sub>)<sub>2</sub> solution in ethanol with pH previously adjusted to 11 using NH<sub>4</sub>OH solution. The mixture was left aging for 1 h under gentle agitation (150 rpm) at room temperature. Hydrothermal treatment at 150 °C was applied using an autoclave Digestec DAB–2 for 24 h. The autoclave was allowed to cool down before opening. The precipitates were separated by centrifugation and washed with water and a 60/40 v/v mixture of ethanol/water (twice). After lyophilizing it for 3 days, a white powder was obtained. The HAp powder was extensively grinded to reduce aggregates and homogenize the grain size, this step being crucial for the nanostructure of the HAp scaffold (see below).

**Synthesis of Pluronic® F-127 hydrogel.** 25 g of distilled water was mixed with 25 g of Pluronic® F-127 polymer using a FlackTek SpeedMixer at 3500 rpm for 5 minutes. After that, 50 g of Pluronic® polymer were added and vigorously stirred using the same conditions. The resultant hydrogel was stored at 4 °C.

**Ultraporous pp-Hap (upp-HAp).** HAp powder was extensively grinded to reduce aggregates and to homogenize the grain size, such step being crucial for achieving the nanostructure of the HAp scaffold. Then, 60% wt. of previously prepared Pluronic® F-127 hydrogel was slowly added to the grinded HAp powder in a cold room at 4 °C. In order to achieve a homogeneous mixture, such additive process was periodically interrupted for stirring at 2500 rpm for a 2 min minutes using a Fisherbrand™ Digital Vortex Mixer. The obtained white paste was left aging at 4 °C for 24 h to ensure the homogeneous distribution of the hydrogel. The resulting ink was modeled using a cold spatula (< 4 °C) to obtain 3D HAp scaffolds with the desired shape. The utilization of the cold spatula allowed to reduce the friction between the metal and the paste without compromising the homogeneity of the shaped samples. Finally, the mixture was calcined at 1000 °C using a muffle Carbolite ELF11/6B/301 for 2 h. In this step, the Pluronic® F-127 hydrogel was completely removed from the resulting HAp modeled structure.

Calcined HAp porous structures were thermally and electrically activated to achieve the catalytic properties by placing the samples between two stainless steel plates (AISI 304), which acted as electrodes. The HAp sample was left in control with negative electrode, while the positive electrode was separated 4 cm from the calcined sample. A constant DC voltage of 500 V was applied for 1 h with a GAMMA power supply, while the temperature was kept at 1000 °C. Samples were allowed to cool down maintaining the applied electric potential for 30 minutes, and finally, all the system was powered off and left to cool down overnight.



### Characterization of the upp-HAp catalyst

Raman analyses were performed by means of an inVia Qontor confocal Raman microscope (Renishaw) equipped with a Renishaw Centrus 2957T2 detector and a 532 nm laser. In order to obtain representative data, 32 single point spectra were averaged.

X-ray diffraction (XRD) studies were conducted using a Bruker D8 Advance model with Bragg-Brentano  $2\theta$  configuration and Cu  $K_{\alpha}$  radiation ( $\lambda = 0.1542$  nm). Measurements were performed in a  $2\theta$  range of  $20^{\circ} - 60^{\circ}$  in steps of  $0.02^{\circ}$ , and a scan speed of 2 s using a one-dimensional Lynx Eye detector.

Scanning electron microscopy (SEM) images were obtained using a Zeiss Neon40 microscope equipped with a SEM GEMINI column.

High resolution transmission electron microscopy (HRTEM) was performed in a JEOL 2010F microscope equipped with a field emission electron source and operated at an accelerating voltage of 200 kV. The point-to-point resolution was 0.19 nm, and the resolution between lines was 0.14 nm. Samples were dispersed in an aqueous (Milli-Q) suspension in an ultrasonic bath, and a drop of the suspension was placed over a grid with holey-carbon film. Images were not filtered or treated by means of digital processing and they correspond to raw data.

Near-Ambient Pressure X-ray Photoelectron Spectroscopy (NAP-XPS) experiments were performed with a SPECS Surface Nano Analysis GmbH (Berlin Germany) system. The spectrometer was equipped with an Al (XR50) anode operating at 400 W and with a Phoibos 225 (SPECS) detector (hemispherical energy analyzer, HAS). The pass energy of the hemispherical analyzer was set to 20 eV and the energy step of high resolution spectra was set at 0.1 eV. The temperature of the analysis was controlled using an Electron Beam Heater (EBH-150, SPECS Surface Nano Analysis GmbH) which has been monitored during the whole experiment through a thermocouple sensor. The C 1s peak was used as internal reference with a binding energy of 284.5 eV. The obtained experimental spectra were fitted using a Gaussian-Lorentzian curve.

### Batch reactions

The reactor consisted in an inert reaction chamber (120 mL) coated with a perfluorinated polymer where both the catalyst and water were placed. All surfaces were coated with a thin film of perfluorinated polymer in order to avoid any contact among the reactants, catalyst and the stainless steel reactor surfaces, in order to discard other catalytic effects. The reactor was equipped with an inlet valve for the entrance of gases and an outlet valve to recover the gaseous reaction products.

Both upp-HAp catalyst and de-ionized liquid water (20 mL) were introduced into the reactor. After exhaustive purge with  $\text{CO}_2$ , the chamber pressure was increased up to 6 bar of  $\text{CO}_2$  (measured at room temperature). The reaction was conducted for 1, 4, 8 and 18 h at  $120^{\circ}\text{C}$ . All processes were performed in triplicate. Isotope-labeled experiment was carried out using  $^{13}\text{C}\text{O}_2$  purchased from Cambridge Isotope Laboratories INC (purity:  $^{13}\text{C}$  99%). 1 bar of  $^{13}\text{C}\text{O}_2$  and 5 bar of  $^{12}\text{C}\text{O}_2$  were

introduced in the reactor. The reaction temperature was set at  $120^{\circ}\text{C}$  for 72 h.

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### Continuous flow reactions

The continuous  $\text{CO}_2$  conversion into valuable chemicals was carried out using a commercial borosilicate glass continuous-flow reactor novaLight CUBE 100 (Peschl Ultraviolet GmbH, Germany). All reactions were conducted using a constant  $\text{CO}_2$  flow of 100 mL/s. Despite the reactor was designed to allow UV irradiation, all reactions were conducted without UV light since previous studies using a batch reactor showed that such energy supply is not required when pp-HAp or upp-HAp catalysts are used.<sup>55,62</sup> Reactions were performed introducing 1 mL of liquid water into the reactor. When performed without liquid water is explicitly indicated for each case below. All the studied processes were performed in triplicate.

### Analysis of the reaction products

The reaction products dissolved in the liquid water (batch reactions), the reaction products condensed by the cold trap (continuous reaction), and the reaction products adsorbed on the catalyst were analyzed by  $^1\text{H}$  NMR spectroscopy. In order to desorb the reaction products from the catalyst, samples were dissolved in a solution containing 100 mM of HCl and 50 mM of NaCl with the final addition of deuterated water. All  $^1\text{H}$  NMR spectra were acquired with a Bruker Avance III-400 spectrometer operating at 400.1 MHz. The chemical shift was calibrated using tetramethylsilane as internal standard. 256 scans were recorded in all cases to obtain an optimum signal-to-noise-ratio for proper area integration. In order to compare the different products obtained from the studied reactions, the areas associated to the proton contribution were normalized and regularly calibrated through external references to ensure a high accuracy of the measurements.

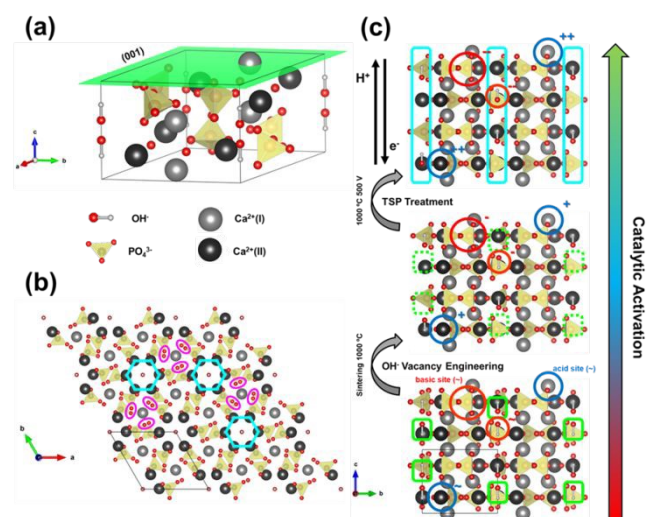
After the batch reactions, the gas was collected by means of a MSHA portable pump and stored into a Tedlar<sup>®</sup> gas sampling bag (LB-2 Septa) and analyzed on a Trace1300 gas chromatograph equipped with a valve for gas injection fitted with a thermal conductivity detector (ThermoFisher Scientific). Prior to the measurements, the samples were transferred from the Tedlar<sup>®</sup> bags to multi-sorbent bed tubes (Carbotrap, Carboxen X and Carboxen 569) to ensure proper storing and manipulation.

## Results and Discussion

### Design and characterization of the catalyst

The upp-HAp catalyst was prepared as is described in the Methods section. The new properties of such catalytic active ceramic rely on the enhanced charge accumulation and boosted electron transport properties achieved through intensive lattice refinement. As depicted in Scheme 1, while vacancy engineering (i.e. generation of  $\text{OH}^{\cdot}$  vacancies) is considered to increase the quantity of available charges (Figure S1 and S2), and thus, the activity of the naturally occurring acid binding sites ( $\text{Ca}^{2+}$  as Lewis and P-OH as Brønsted acidic sites) and basic Lewis binding sites ( $\text{O}^{2-}$  atoms

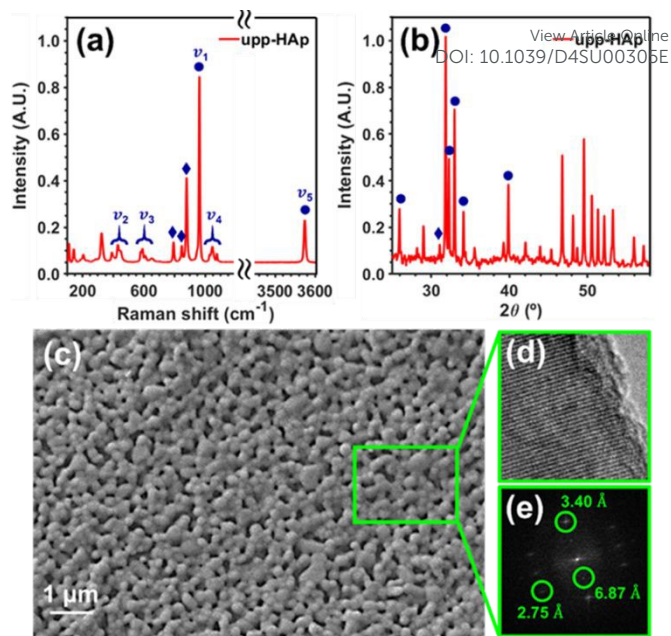




**Scheme 1.** Structure of HAP as a catalyst. (a) HAP unit cell  $P6_3/m$ . (b) (001) crystallographic plane observed from the c-axis direction. The  $\text{OH}^-$  channels and the Ca(I) 6-fold coordination are highlighted. (c) Structural changes and lattice refinement strategies adopted to achieve the catalytic activation of HAP.

of  $\text{PO}_4^{3-}$  and  $\text{OH}^-$ ) of HAP;<sup>63</sup> the TSP treatment imposes a specific global orientation to the hydroxyl groups oriented in columns along the c-axis that boosts the electron transport (Figure S3), and consequently, favors the catalytic conversion of carbon dioxide towards new products. Note that although upp-HAp could be strictly referred as a thermal catalyst (i.e. neither external electric currents nor UV irradiation are needed for the reaction) the nature of the catalyst strongly relies on its refined electrical properties. Hence, upp-HAp finds more similarities with the classical electroreduction mechanisms and formulations reported in the literature. In this sense, such dual approach postulates upp-HAp as a sustainable green candidate to replace the current metal-based thermo catalysts, further competing directly with the efficiencies obtained by means of conventional electrocatalysis. Such features have been widely studied in the literature in terms of the customization of the enhanced electrical properties (material characterization approach), reporting exhaustive structural analyses to ensure its proper catalytic activation.<sup>55,60-62,64-69</sup>

Accordingly, the Raman spectrum of upp-HAp (Figure 1a) displays the characteristic active modes of HAp with  $\nu_1 = 962 \text{ cm}^{-1}$  (P–O symmetric stretching),  $\nu_2 = 400\text{--}480 \text{ cm}^{-1}$  (doubly degenerated O–P–O bending),  $\nu_3 = 570\text{--}625 \text{ cm}^{-1}$  (P–O triply degenerated asymmetric stretching),  $\nu_4 = 1020\text{--}1095 \text{ cm}^{-1}$  (triply degenerated O–P–O bending mode) and  $\nu_5 = 3574 \text{ cm}^{-1}$  (O–H stretching).<sup>63</sup> Besides, the peaks at 878, 844 and  $794 \text{ cm}^{-1}$  are attributed to the formation of brushite ( $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$ ) due to the polarization process, thus proving the correct catalytic activation of HAp.<sup>66</sup> Consistently, the XRD diffractogram of upp-HAp (Figure 1b) shows the typical HAp reflections at  $2\theta = 25.9^\circ$  (002),  $31.8^\circ$  (211),  $32.3^\circ$  (112) and  $33.0^\circ$  (300), while the presence of brushite is confirmed by the peak at  $31.1^\circ$  (121).<sup>66</sup> Also, comparison with the Raman spectrum and XRD diffractogram of calcined HAp samples (i.e. just before applying the TSP process), which are shown in Figure S4,



**Figure 1.** (a) Raman spectrum, (b) XRD diffractogram, (c) SEM and (d) HRTEM micrographs, and (e) fast Fourier Transform of the HRTEM micrograph for upp-HAp. In (a) the characteristic HAp ( $\nu_1$ – $\nu_5$ ) and brushite (diamonds) Raman vibrational modes are displayed. In (b), the characteristic HAp (spots) and brushite (diamonds) reflections are indicated.

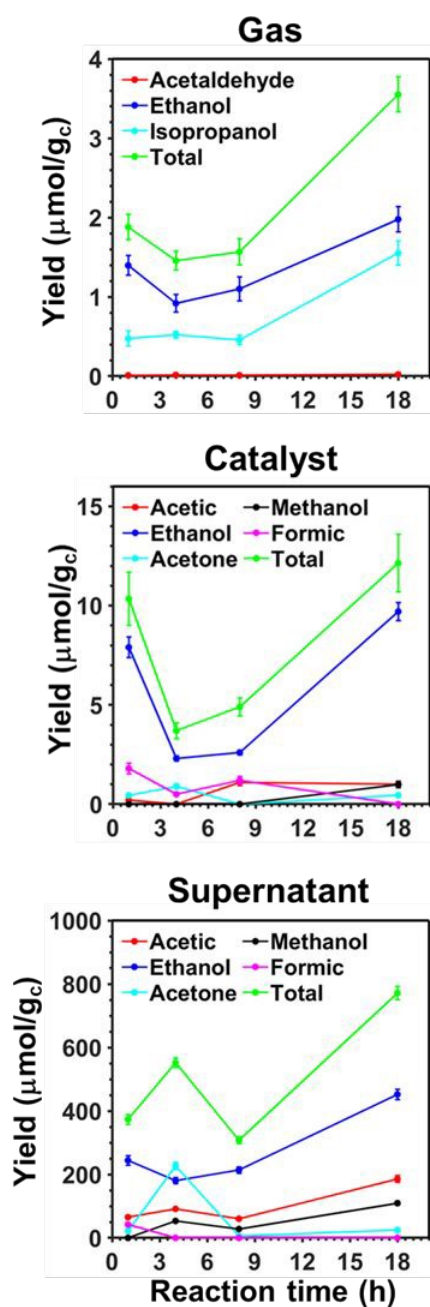
confirms the success of the treatment used to induce permanent polarization.

SEM and HRTEM micrographs of upp-HAp are shown in Figures 1c–e. As it can be seen, SEM image reflects the ultraporous structure of the catalyst with average pore size of  $156 \pm 32 \text{ nm}$ . The porosity, which was estimated considering five independent SEM images, was  $17\% \pm 2\%$ . The crystallinity of the samples was confirmed by the lattice fringes observed in HRTEM images (Figure 1d). Fast Fourier Transform of the upp-HAp fringes (Figure 1e) allowed not only the distinction between the (002) and (112) of HAp crystallographic planes at  $3.40 \text{ \AA}$  and  $2.75 \text{ \AA}$ , respectively, but also the identification of the characteristic polarized superstructure (001) at  $6.87 \text{ \AA}$  d-spacing.

#### From batch to continuous reactions

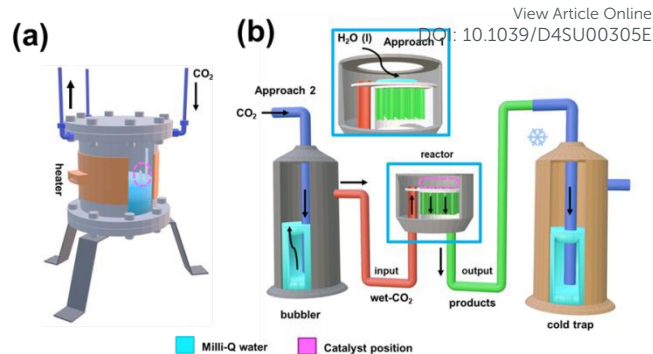
In previous studies, the efficiency of pp-HAp and upp-HAp catalyst to convert  $\text{CO}_2$  alone<sup>67</sup> or mixed with  $\text{CH}_4$ <sup>55,62</sup> into ethanol (predominant) and other valuable chemicals was examined quantifying by  $^1\text{H}$  NMR the amount of reaction products dissolved in the liquid water (hereafter named “supernatant”) and adsorbed on the catalyst. Thus, we assumed that the content of products in the gas phase was negligible in comparison to those in the supernatant and on the catalyst. In this work, as a first step and before transferring the process to a continuous-flow reactor, we experimentally examined the time evolution of the gas composition compared to the supernatant and the catalyst of the batch reactions. The results presented in Figure 2 are expected to provide better understanding about the distribution of products among the different phases in the continuous-flow process described in the next section.





**Figure 2.** Variation of the different product yields in the gas, adsorbed on the catalyst and dissolved in the supernatant vs the reaction time obtained from batch reactions catalyzed by upp-HAp. Reactions were conducted at 120 °C using 6 bar of CO<sub>2</sub> and 20 mL of liquid water. The total yield, which corresponds to the sum of the yields of all the individual products, is represented as well. The yields are expressed in μmol of product per gram of catalyst (μmol/g<sub>c</sub>).

The variations of the product yields through time confirm that the resulting species are partially adsorbed on the solid catalyst and mainly dissolved in the supernatant. Thus, the yields of the different products are one and two orders of magnitude smaller in the gas than in the catalyst and the supernatant, respectively. Furthermore, the three analyzed phases present reaction products containing two or three carbon atoms (C2 and C3, respectively), while products with one carbon atom (C1) are only detected in the catalyst and

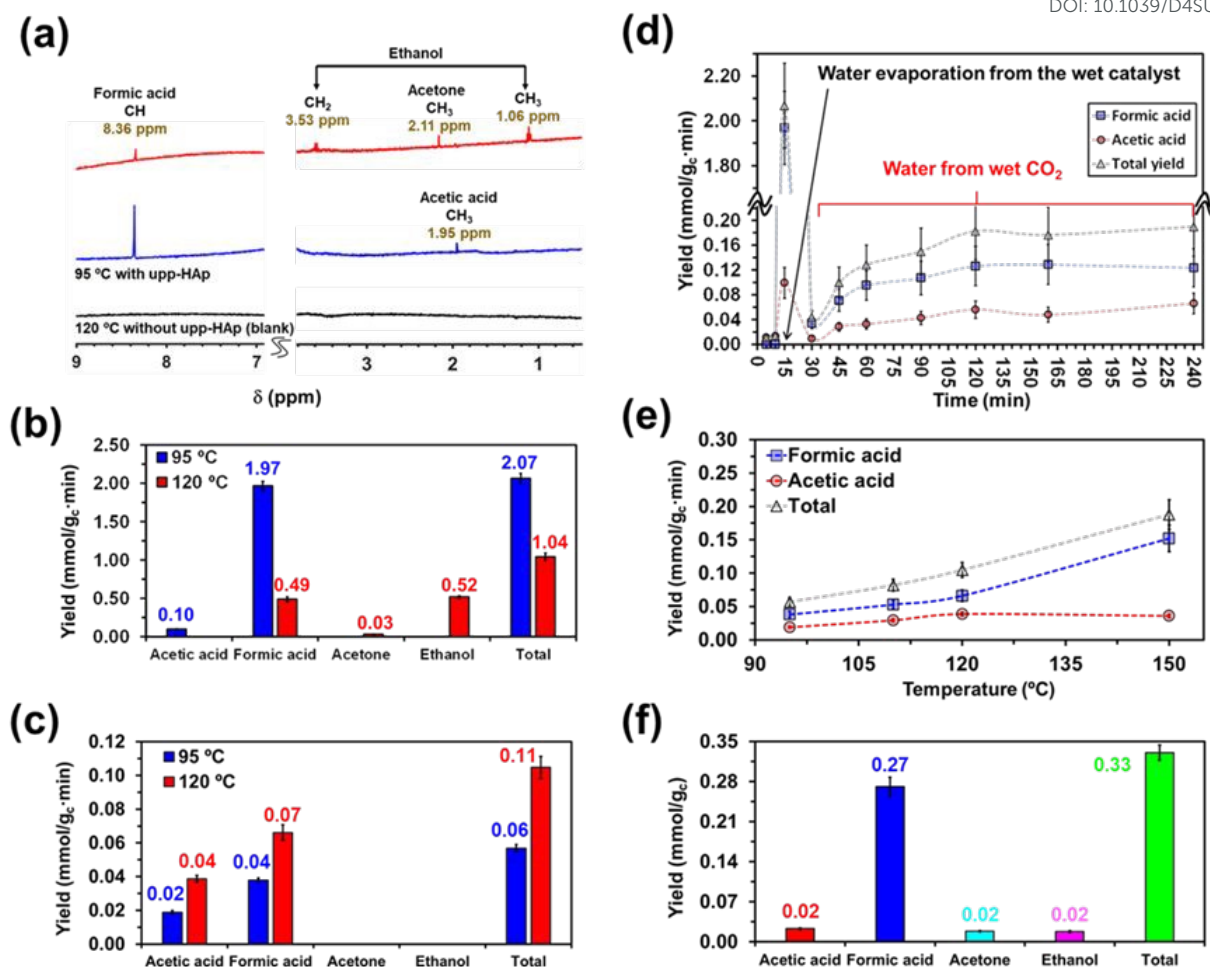


**Scheme 2.** (a) Scheme of batch reactor used, (b) Scheme of continuous reactor used. The continuous reactor was filled with four upp-HAp catalysts (3.60 × 5.00 × 0.44 mm<sup>3</sup> each) and 1 mL of de-ionized water which was homogeneously distributed (approach 1). The outlet valve of the reactor was connected to a cold trap to condense the reaction products obtained in the gas phase, while the inlet valve was connected to a CO<sub>2</sub> cylinder to achieve a constant flow of 100 mL/s, which was controlled using a back-pressure regulator. This high flow was selected to mimic the conditions at pre-pilot scale. To have the required moisture, as water is the proton source, CO<sub>2</sub> was bubbled in liquid water to produce wet-CO<sub>2</sub>, the bubbler being located between the CO<sub>2</sub> cylinder and the inlet valve (approach 2).

supernatant (*i.e.* methanol and formic acid). This feature suggests that C1 products evolve towards C2 and C3 before migrating to the gas phase. The thermodynamics of water splitting and CO<sub>2</sub> reduction into formic acid, which is an exhaustively studied reaction,<sup>68-76</sup> have been discussed in a recent review.<sup>77</sup> Besides, all C2 products seem to be inter-related in the sense that acetic acid and acetaldehyde probably come from the complete or partial oxidation of ethanol, respectively. Similarly, acetone is catalytically hydrogenated to isopropanol at low pressure and temperature when the mass transfer between the liquid and the gas is high enough.<sup>78</sup> Further analysis of the results is discussed in the ESI. On the other hand, the water content available in the reactor also merits special attention when moving from batch to continuous reactions. That is because water not only provides a media to collect the products desorbed from the catalyst (*i.e.* supernatant) but also acts as the proton source. Accordingly, in Scheme 2 and Figure S5 are compared the batch and continuous reactor systems used. Thus, for the continuous reactions two different scenarios were considered: 1) Liquid water is directly introduced inside the reactor (1 mL); and, 2) CO<sub>2</sub> is bubbled in liquid water to produce wet-CO<sub>2</sub> (see Scheme 2b). Both reaction approaches were carried out at 95 °C and 120 °C analyzing the products obtained in the cold trap (Scheme 2b) after 15 minutes. Representative <sup>1</sup>H NMR spectra are presented in Figure 3a (complete spectra are shown in Figure S6), while the yields of the reaction are provided in Figures 3b and 3c for scenarios 1) and 2), respectively. Moreover, a blank (*i.e.* reaction without upp-HAp catalyst at 120 °C using the same experimental conditions that the upp-HAp catalyzed reactions) was also conducted (Figure 3a) to discard other carbon sources.

As it can be observed, the collected spectra showed some important differences, which suggested that the evaporation rate of water plays a crucial role in the reaction mechanisms. The reaction at 95 °C led to the formation of formic acid identified by the singlet at 8.36 ppm, as the predominant





**Figure 3.** Continuous reaction studies. (a) Representative  $^1\text{H}$  NMR spectra used to identify the reaction products for the continuous processes catalyzed by upp-HAp and performed at 95 and 120 °C using a  $\text{CO}_2$  flow of 100 mL/s and 1 mL of liquid water. The spectrum of a blank (*i.e.* without catalyst) at 120 °C is also displayed. Each spectrum corresponds to the liquid collected in the cold trap after 15 min of reaction. (b) Yield of the reaction products for the continuous process described in (a). The yields were calculated from the  $^1\text{H}$  NMR spectra of the liquids collected in the cold trap after 15 min of reaction. The total reaction yield (*i.e.* sum of the yields of the individual products) is also displayed. (c) Yield of the reaction products for the continuous process catalyzed by upp-HAp at 95 °C and 120 °C but without including liquid water in the reactor. The yields were derived from the  $^1\text{H}$  NMR spectra of the liquids collected in the cold trap after 15 min of reaction. (d) Evolution of the non-accumulated yields of the reaction products identified by  $^1\text{H}$  NMR against the reaction time for the continuous processes conducted including 1 mL of liquid water in the reactor and catalyzed by upp-HAp at 95 °C. (e) Variation of the temperature for a processes conducted up to 30 min. In all cases, the reactions were catalyzed by upp-HAp and using wet  $\text{CO}_2$  but without including liquid water in the reactor. (f) Reaction products adsorbed on the upp-HAp catalyst after several continuous-flow catalytic processes at 120 °C without including liquid water in the reactor. The amount of product was calculated from the  $^1\text{H}$  NMR spectrum collected after dissolve the catalyst using an aqueous solution containing 100 mM of HCl and 50 mM of NaCl. The total yield, which is the sum of the yields of all the individual products, is also represented. The yields are expressed mmol of product per gram of catalyst and min ( $\text{mmol/g}_c\cdot\text{min}$ ).

product, and acetic acid denoted by the  $\text{CH}_3$  singlet at 1.95 ppm as residual product. Thus, the reaction exhibited not only a high yield of formic acid (expressed as mmol of product per gram of catalyst and minutes),  $1.97 \pm 0.06 \text{ mmol/g}_c\cdot\text{min}$ , but also a very high selectivity ( $\sim 95\%$ , comparable to the photocatalysts)<sup>79</sup> towards such C1 product being the ratio of formic acid / acetic acid yields around 20 (Figure 3b). Compared to the literature, the results reported here represent a compelling step forward in the field of heterogeneous catalysis being extremely competitive not only when considering the total yield obtained ( $124 \text{ mmol/g}_c\cdot\text{h}$  for upp-HAp compared with the standard range of 2–20  $\text{mmol/g}_c\cdot\text{h}$ )<sup>79</sup> but also because of the greener and more sustainable reaction conditions achieved (*i.e.* mild conditions, use of water instead of  $\text{H}_2$ , avoids the use of electrical supply or UV irradiation, use of a green and abundant catalyst). Table 1 compares all the features obtained with respect the conventional heterogeneous catalysts used (more detailed

information can be also found in Table S1). Surprisingly, the total yield of products at 95 °C ( $2.07 \pm 0.07 \text{ mmol/g}_c\cdot\text{min}$ ) decreased by half at 120 °C ( $1.04 \pm 0.05 \text{ mmol/g}_c\cdot\text{min}$ ) and the reaction became much less selective. Thus, in addition of formic acid ( $0.49 \pm 0.03 \text{ mmol/g}_c\cdot\text{min}$ ), both ethanol (C2) and acetone (C3) were clearly detected at 120 °C. The yield of ethanol ( $0.52 \pm 0.02 \text{ mmol/g}_c\cdot\text{min}$ ), which was identified by the quartet from  $\text{CH}_2$  at 3.53 ppm and the triplet from  $\text{CH}_3$  at 1.06 ppm (Figure 3a), was comparable to that of formic acid. Conversely, acetone, with the  $\text{CH}_3$  singlet at 2.11 ppm (Figure 3a), remained as a residual product ( $0.03 \pm 0.01 \text{ mmol/g}_c\cdot\text{min}$ ). These observations indicate that such conditions (*i.e.* temperature high enough to promote the rapid evaporation of liquid water) favor the catalytic formation of C–C bonds, although it is detrimental to the efficiency of the overall  $\text{CO}_2$  conversion process (Figure 3b).





**Table 1.** Comparison between the most relevant features of this work and the standard conditions reported for heterogeneous catalysis.<sup>79</sup>

	Catalyst	Proton Source	T (°C)	Maximum Selectivity (%)	Productivity (mmol/g <sub>c</sub> -h)
<i>Heterogeneous Catalysis</i> <sup>79</sup>	Pd, Au, Cu-ZnO, M-ZrO <sub>x</sub> ,...	H <sub>2</sub>	220-300	> 70 (methanol)	2-29 (total products)
<i>This Work</i>	upp-HAp	H <sub>2</sub> O	95	~ 95 (formic acid)	124 (total products)

In the batch reactions, which were conducted using 20 mL of liquid water, ethanol was the predominant product in both the supernatant and the catalyst, while the content of formic acid was residual (Figure 2). Instead, ethanol and formic acid were formed with similar yields when only 1 mL of liquid water was introduced in the continuous reactor. This feature suggests that, in the batch process, the excess of evaporated water favors the formation of ethanol (*i.e.* generation of a C–C bond and hydrogenation) over the reduction of CO<sub>2</sub> to formic acid (C1), while the continuous flow of CO<sub>2</sub> and the decrease of available water molecules promote the conversion of CO<sub>2</sub> to formic acid instead of producing ethanol.

Such hypothesis can be further corroborated by analyzing the yields of the products obtained using wet-CO<sub>2</sub> (approach 2). As seen in Figure 3c, both formic and acetic acid are obtained, whereas ethanol and acetone were not detected. Comparison of the yields, which are shown in Figure 3c, with Figure 3b reveals a reduction in the total yield of two and one order of magnitude at 95 and 120 °C, respectively. Furthermore, in absence of liquid water, the yields of the two reaction products increased with temperature, as expected. Overall, those results demonstrate that liquid water is responsible for both the variety of products and the large yields reflected in Figure 3b.

All these features can be clearly observed when merging both approaches (*i.e.* wet-CO<sub>2</sub> and addition of 1 mL of liquid water in the reactor) in a single long term (4 hours) continuous reaction depicted in Figure 3d and exhaustively described in the ESI. Considering a CO<sub>2</sub> flow of 100 mL/s (0.24 mol/min), the yields reported at 15 min in Figure 3d in the lab scale laboratory indicate that conversion of CO<sub>2</sub> into products is of around 1 %. The yields at 95 °C (Figure 3d) stabilized at 7.4 ± 0.4 and 4.0 ± 0.4 mmol/g<sub>c</sub>-h for formic acid and acetic acid, respectively. Such values were significantly higher than those reported in the literature for the photocatalytic and the electrocatalytic reduction and hydrogenation of CO<sub>2</sub>.<sup>80,81</sup>

To better understand the reaction sequence, the yields of the individual products and the total yield were analyzed as a function of the temperature after 30 min of reaction. Results for reactions performed at 95, 110, 120 and 150 °C, which are plotted in Figure 3e, showed that all yields increased with the temperature, even though two regimes were observed. At temperatures ≤ 120 °C, the yields of formic acid and acetic acid, as well as the total yield, increased linearly with the

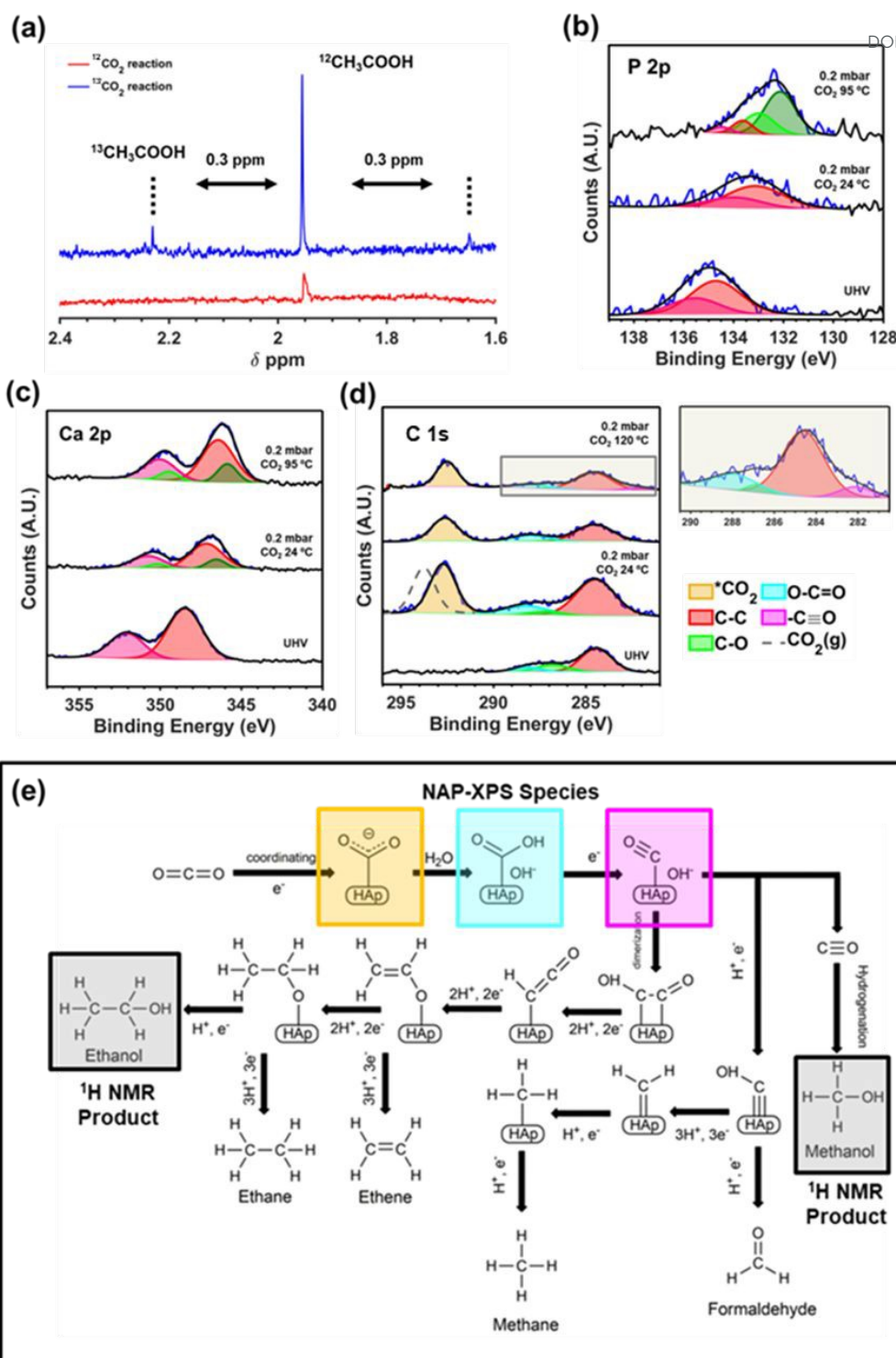
temperature. However, at 150 °C, the yield of acetic acid remained unchanged while that of formic acid increased significantly, suggesting an enhancement in the dominance of the CO<sub>2</sub> reduction to formic with respect to the formation of C–C bonds. After such assays, the products adsorbed in the catalyst were analyzed by <sup>1</sup>H NMR. Accordingly, results presented in Figure 3f have been further discussed in the ESI. Another important observation refers to the catalytic stability of upp-HAp for both batch and continuous processes, which was evidenced by the fact that the yields were maintained in all repetitions carried out (*i.e.* the same catalyst samples were always used). Figure S7, which compares the Raman spectra of the catalyst before and after reaction, confirms such stability. As it can be observed, the main vibration ( $\nu_1$  peak at 962 cm<sup>-1</sup>), which was assigned to the main symmetric stretching P–O mode, does not show any significant difference. It is worth mentioning that any loose of electrical and, consequently, catalytic activity would be detected by a widening and/or splitting of the  $\nu_1$  band in the Raman spectrum. In this sense, this study is in agreement with the results reported in a recent work from where the complete stability of the upp-HAp catalysts is studied through different characterization techniques and under a wide range of temperatures.<sup>82</sup>

#### Mechanistic insights on CO<sub>2</sub> activation: NAP-XPS studies and isotope-labeling reaction

Due to the novelty of the catalyst and the relevance of converting the thermodynamically stable CO<sub>2</sub> molecule in added value molecules, significant effort has been put on experimentally exploring the initial steps of CO<sub>2</sub> adsorption and activation on upp-HAp surface. Firstly, we performed isotope-labeled experiments by loading <sup>13</sup>CO<sub>2</sub> and 5 bar of <sup>12</sup>CO<sub>2</sub> in a batch reactor (20 mL liquid water and 120 °C for 72 hours) to isotopically confirm the CO<sub>2</sub> fixation reaction. Figure 4a compares the <sup>1</sup>H NMR spectra collected from the supernatant of a standard (*i.e.* 6 bar of <sup>12</sup>CO<sub>2</sub>) and the isotope-labeled reaction, focusing on the region of acetic acid (1.6–2.4 ppm). As it can be seen, the distinctive doublet of <sup>13</sup>CH<sub>3</sub>COOH can be observed shifted ~0.3 ppm from the <sup>12</sup>CH<sub>3</sub>COOH peak at 1.95 ppm.<sup>83</sup>

Additionally, near-ambient pressure X-ray photoelectron spectroscopy (NAP-XPS) studies using 0.2 mbar of CO<sub>2</sub> and at different reaction temperatures have been further performed to better understand the role of the binding sites of upp-HAp. Accordingly, Figures 4b and 4c show the high resolution spectra of the P 2p and Ca 2p regions, respectively. As soon as the CO<sub>2</sub> is introduced in the analysis chamber, the characteristic peaks of P 2p at 134.7 eV (P 2p<sub>3/2</sub>) and 135.5 eV (P 2p<sub>1/2</sub>) experience a clear shift of 1.6 eV attributed to the absorption of CO<sub>2</sub> on the PO<sub>4</sub><sup>3-</sup> basic sites. Moreover, at 95 °C new species at 132.1 eV (P 2p<sub>3/2</sub>) and 132.9 eV (P 2p<sub>1/2</sub>) shifted an additional 1 eV, suggesting the subsequent fixation steps of the adsorbed CO<sub>2</sub> molecules. On the other hand, Ca 2p (Figure 4c) is also affected by the CO<sub>2</sub> atmosphere, shifting from the original binding energy in ultra-high vacuum (UHV) conditions of 348.5 eV (Ca 2p<sub>3/2</sub>) and 352.1 eV (Ca 2p<sub>1/2</sub>) 1.3 eV and 2 eV.





**Figure 4.** (a) isotope-labeling experiments. High resolution NAP-XPS studies using  $^{12}\text{CO}_2$  of the P 2p (b), Ca 2p (c) and C 1s (d). (e) Fixation mechanism derived from  $^1\text{H}$  NMR and NAP XPS results.

As expected due to its acidic nature, the Ca species remain stable (compared to P) when temperature is increased. Thus, the shift observed is attributed to the 6-fold coordination with the oxygen atoms of the lattice. Moreover, the fact that two different species appear is in agreement with the fact that exists two distinctive crystallographic sites for calcium cations. More specifically, Ca(I) is strictly coordinated with the oxygens

from the  $\text{PO}_4^{3-}$  tetrahedra, while Ca(II) is also coordinated with a hydroxyl group.<sup>84</sup>

The C 1s region has been also examined to further confirm the results obtained. Several interesting insights can be derived from the acquired spectra presented in Figure 4d. Firstly, the shift of 1.1 eV observed when comparing the  $\text{CO}_2$  gas measured far from the surface of upp-HAp (dashed grey line;



characteristic peak at 293.7 eV for C 1s which is in agreement with the literature)<sup>85</sup> confirms the adsorption and activation of the CO<sub>2</sub> molecule (\*CO<sub>2</sub>) on the surface of upp-HAp, showing an outstanding atomic \*CO<sub>2</sub>/P ratio of ~3. Such results suggest that apart from the PO<sub>4</sub><sup>3-</sup> sites, the CO<sub>2</sub> adsorption might also occur on the OH<sup>-</sup> binding sites. This preliminary experimental evidence of the active role of several binding sites might support the strong affinity towards the catalytic production of C2 and C3 products obtained. This conclusion merits special attention due to the possibility of developing upp-HAp dual site catalysts.

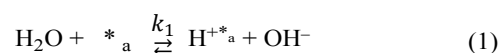
As suggested in Figure 4d, the initial steps of CO<sub>2</sub> activation can be intuited at 95 °C. Accordingly, an increment of the peak at 288.1 eV, attributed to the O–C=O species, can be observed showing a ratio with the adventitious carbon (284.5 eV) of 0.31 (24 °C) and 0.41 (95 °C). Note that during all the experiments samples were kept inside the analysis chamber avoiding the presence and absorption of other carbon-based products on the upp-HAp surface. On the contrary desorption of C–O (contamination during sample preparation) due to temperature can be also observed. To obtain more insights on the fixation mechanism, the temperature of the experiment was finally risen to 120 °C. As it can be observed in the inset of Figure 4d, a small peak appears at 282.1 eV which has been attributed to a partially dissociated C≡O species anchored on the catalyst (resulting in carbidic species). Such result supports the unique structural and electronic properties derived from the vacancy engineering and TSP treatments (i.e. superior crystallinity and high charge delocalization) favoring the fast evolution of the generated carbon intermediates. Overall, combined <sup>1</sup>H NMR and NAP-XPS results allow proposing the first catalytic CO<sub>2</sub> fixation mechanism on upp-HAp (see Figure 4e) which not only supports experimentally the theoretical pathways proposed for HAp<sup>67</sup> and/or other classical electro-mechanisms<sup>28</sup> but also proves the feasibility of performing extremely efficient CO<sub>2</sub> fixation reactions under strictly mild reaction conditions using only upp-HAp as a sustainable and green catalyst. It is worth highlighting the fact that the mechanism proposed involves the C–C coupling by means of a dimerization process, supported by the literature.<sup>67</sup> Such mechanistic pathway requires a high abundance of HCOO<sup>-</sup> species simultaneously adsorbed on the surface of the catalyst. According to Figure 4e the presence of water is crucial to do so, which further corroborates the experimental results discussed above.

### Kinetics of the CO<sub>2</sub> fixation reaction

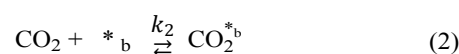
Herein, we present a kinetic model for the CO<sub>2</sub> fixation reaction over upp-HAp, which has been developed combining the fixation mechanisms described in the literature<sup>66</sup> with the experimental results obtained in this work. The main challenge resides in discerning the crucial reaction steps governing the kinetics of the CO<sub>2</sub> fixation, which is essential to discern among the complex known network of intermediate pathways.<sup>67</sup> Figure 3d shows that both the total yield and the yields for the individual products present an initial linear behavior (first-

order reaction) followed by a stabilization regime (zero-order reaction). This fact clearly indicates a competing phenomenon between the adsorption of reactants and desorption of products. For this reason, the Langmuir-Hinshelwood-Hougen-Watson (LHHW) and the Eley-Rideal (ER) approaches, which are frequently used to model heterogeneous catalysis,<sup>86,87</sup> have been considered to derive the expression of the reaction rate, according to a classical electroreduction mechanism, as follows:

Water splitting:



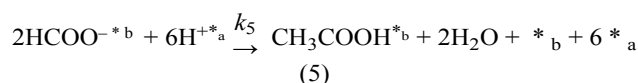
CO<sub>2</sub> adsorption:



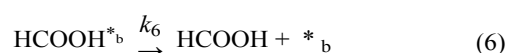
CO<sub>2</sub> fixation:



Chain reaction:



Product desorption:



where \* (written as plain text) corresponds to a free binding site while when written as a superscript it is used to denote an adsorbed specie (i.e. HCOOH<sup>\*<sub>b</sub></sup>). It should be emphasized that the already reported unique electric and charge storage properties of upp-HAp are, in general, consistent with such classical reduction mechanism.<sup>51,52</sup>

In Eqs 1-7, we propose a four step chain mechanism conformed by:<sup>53,54,67</sup> 1) Initial water splitting reaction in the acid sites (\*<sub>a</sub>) corresponding to the Ca<sup>2+</sup> sites of upp-HAp (Eq 1), occurring simultaneously with CO<sub>2</sub> adsorption in the basic sites (\*<sub>b</sub>) mainly attributed to the OH<sup>-</sup> and PO<sub>4</sub><sup>3-</sup> sites (Eq 2);<sup>87,88</sup> 2) CO<sub>2</sub> fixation into HCOO<sup>-</sup> due to the addition of a neighboring proton (Eq 3), which is going to be initially considered as the rate determining step (RDS); 3) Chain reaction towards adsorbed formic acid (Eq 4) or less likely, to adsorbed acetic acid (Eq 5) due to the C–C coupling of two HCOO<sup>-</sup> adsorbed species through a dimerization process;<sup>53</sup> and 4) Products desorption (Eqs 6-7). Foremost and derived from point (1), the fraction of vacant sites (θ) defined according to the LHHW model has to be described separately for both acid and basic binding sites:

$$\theta_{\text{a}} = \frac{1}{1 + k_1 \cdot [\text{H}_2\text{O}]} \quad (8)$$



**Table 2.** Statistical parameters obtained from the rate of the expressions considering different assumptions. The principal differences compared to the Eqs 4-9 have been highlighted in different colors. (Red: different RDS; and Blue: water splitting).

Approach	Assumption	SSE ( $\times 10^{-7}$ )	MSC	Variance error ( $\times 10^{-7}$ )
A1	$\text{CO}_2^{\text{b}} + \text{H}^{\text{a}} \xrightarrow{k_3} \text{HCOO}^{\text{b}} + \text{*}_a$ (RDS)	6.38	2.953	1.06
	$\text{HCOO}^{\text{b}} + \text{H}^{\text{a}} \xrightarrow{k_4} \text{HCOOH}^{\text{b}} + \text{*}_a$			
	$2\text{HCOO}^{\text{b}} + 6\text{H}^{\text{a}} \xrightarrow{k_5} \text{CH}_3\text{COOH}^{\text{b}} + 2\text{H}_2\text{O} + \text{*}_b + 6\text{*}_a$			
A2	$\text{CO}_2^{\text{b}} + \text{H}^{\text{a}} \xrightarrow{k_3} \text{HCOO}^{\text{b}} + \text{*}_a$ (RDS)	2.30	4.196	0.37
	$\text{HCOO}^{\text{b}} + \text{H}^{\text{a}} \xrightarrow{k_4} \text{HCOOH}^{\text{b}} + \text{*}_a$			
A3	$2\text{HCOO}^{\text{b}} + 6\text{H}^{\text{a}} \xrightarrow{k_5} \text{CH}_3\text{COOH}^{\text{b}} + 2\text{H}_2\text{O} + \text{*}_b + 6\text{*}_a$ (RDS) (modeling the kinetics of total $\text{CO}_2$ fixation)	3.56	3.759	0.56
A3*	$2\text{HCOO}^{\text{b}} + 6\text{H}^{\text{a}} \xrightarrow{k_5} \text{CH}_3\text{COOH}^{\text{b}} + 2\text{H}_2\text{O} + \text{*}_b + 6\text{*}_a$ (RDS) (modeling the kinetics of $\text{CH}_3\text{COOH}$ alone)	0.20	6.380	0.03
A4	$\text{H}_2\text{O} + \text{*}_a \xrightleftharpoons[k_{-1}]{k_1} \text{H}^{\text{a}} + \text{OH}^-$ (RDS) – Experimentally discarded	-	-	-
A5	$\text{H}_2\text{O} + \text{*}_a \xrightleftharpoons[k_{-1}]{k_1} \text{H}_3\text{O}^+ + \text{*}_a$	2.27	4.194	0.37
	$\text{CO}_2^{\text{b}} + \text{H}_3\text{O}^+ \xrightarrow{k_3} \text{HCOO}^{\text{b}} + \text{H}_2\text{O}$ (RDS)			

$$\theta_b = \frac{1}{1 + k_2 \cdot [\text{CO}_2] + k_6 \cdot [\text{HCOOH}] + \sum_j k_j \cdot [\text{products}]_j} \approx \frac{1}{1 + k_2 \cdot [\text{CO}_2] + k_6 \cdot [\text{HCOOH}]} \quad (9)$$

According to the results displayed in Figure 3f, the term  $\sum_j k_j \cdot [\text{products}]_j$  in Eq 9, which accounts for the contribution of the rest of products adsorbed in the catalyst (including acetic acid), has been assumed to be negligible in front of the adsorbed formic acid content.

Due to the complexity of the system, interesting questions arise related to the contribution of the different chain reactions to the overall kinetics (which might regulate the RDS) and to the competing nature of the water splitting reaction (as a result of the huge proton demand needed to generate the reported products). In order to shed some light, we modeled the kinetics of the  $\text{CO}_2$  fixation reaction under different hypotheses and compared the resulting statistical parameters. For this purpose, the minimum value for the sum of squared errors (SSE) was chosen. Also, the variance of the error and the Model Selection Criterion (MSC, Eq S1),<sup>89</sup> commonly used in kinetic analyses, were calculated to further confirm the results. The minimum value for variance and higher value for MSC were chosen. The comparison of the different assumptions (hereafter denoted A#) are summarized in Table 2.

As a starting point, assumption A1 was modeled considering that the RDS was Eq 6 and assuming chain reaction towards both formic and acetic acids.<sup>53</sup> Results showed the worst data correlation (Table 2) and was discarded. In scenarios A2 and A3, the chain reaction towards formic acid alone and the dimerization process (Eq 5), respectively, were considered as

RDS.<sup>53</sup> Although both assumptions provided much better statistical data than A1, A2 showed lower SSE and higher MSC values, indicating that the kinetic of the  $\text{CO}_2$  fixation is primarily governed by the first fixation step (Eq 3) and its evolution to formic acid alone. Such conclusion is supported by recently reported density functional theory (DFT) calculations used to study for the conversion of  $\text{CO}_2$  into formic acid.<sup>90</sup> Moreover, discarding the A1 approach is also in agreement with other  $\text{CO}_2$  fixation mechanisms proposed for both electrocatalysts and thermocatalysts.<sup>53,54,90,91</sup>

On the other hand, the synthesis of acetic acid remains extremely limited to Eq 5 (C–C coupling), which acts as the RDS for the latter product in agreement with the experimental results. Accordingly, the expression of the rate from A3 was used to fit the experimental data obtained for acetic acid only (instead of the total  $\text{CO}_2$  fixation, A3\*), showing excellent statistical results. Despite of the fact that the aforementioned conclusion explains the small presence of C2 and C3 adsorbed products (Figure 3f), the non-negligible amount of acetic acid detected might be contradictory. While remarkable differences in the desorption constants of C2 products (*i.e.* acetic acid vs ethanol) have been discarded, the possibility of re-adsorption of formic acid and its subsequent reaction with adsorbed  $\text{HCOO}^-$  is more plausible. In order to demonstrate such hypothesis, a batch reaction consisting on an inert  $\text{N}_2$  atmosphere, with 1 mL of formic acid, 20 mL of water, at 95 °C and for 2 h was carried out. The recorded  $^1\text{H}$  NMR showed the unequivocal existence of acetic acid both adsorbed in the catalyst and dissolved in the supernatant (Figure S8). Hence, although A3\* provides good statistical results, it has to be considered as a very initial approach to the specific kinetics of C2 and C3 products, as it has been demonstrated that the re-



adsorption and subsequent reaction of C1 products play a considerable role in the final model, increasing remarkably the complexity of the system. This mechanism is currently under study through *in situ* and *in operando* conditions to support such hypothesis with experimental evidences.

Finally, the nature of the water splitting reaction has been explored. It was experimentally discarded as the RDS since a first-order reaction regime was found by changing the water content from 1 mL to 80 mL in the batch reactor (Figure S9), which is in discrepancy with the yields reported in Figure 3d. Despite of that fact, the extensive proton demand suggests that  $k_1 \gg k_j$ . Thus, A5 assumes an almost instantaneous desorption of protons, which results in a non-competing behavior, resembling the ER approach. Surprisingly, statistical data showed slightly better results than A2, implying that the generation of products linearly depends on the initial water concentration without expecting any stagnation. This result is in total agreement with the experimental measurements presented in Figure 3d. The high proton reactivity and mobility were previously explained from a structural perspective since upp-HAp showed enhanced proton conductivity along different crystalline domains.<sup>60</sup> Therefore, protons might not be chemisorbed in the acid sites or completely desorbed in the reaction atmosphere but electrostatically attracted to the surface of the catalyst due to the reported charge surface accumulation. Consistently, the null catalytic CO<sub>2</sub> fixation activity found for non-polarized HAp should be related with the lack of protons supplied to the adsorbed CO<sub>2</sub>, clearly limiting the RDS.

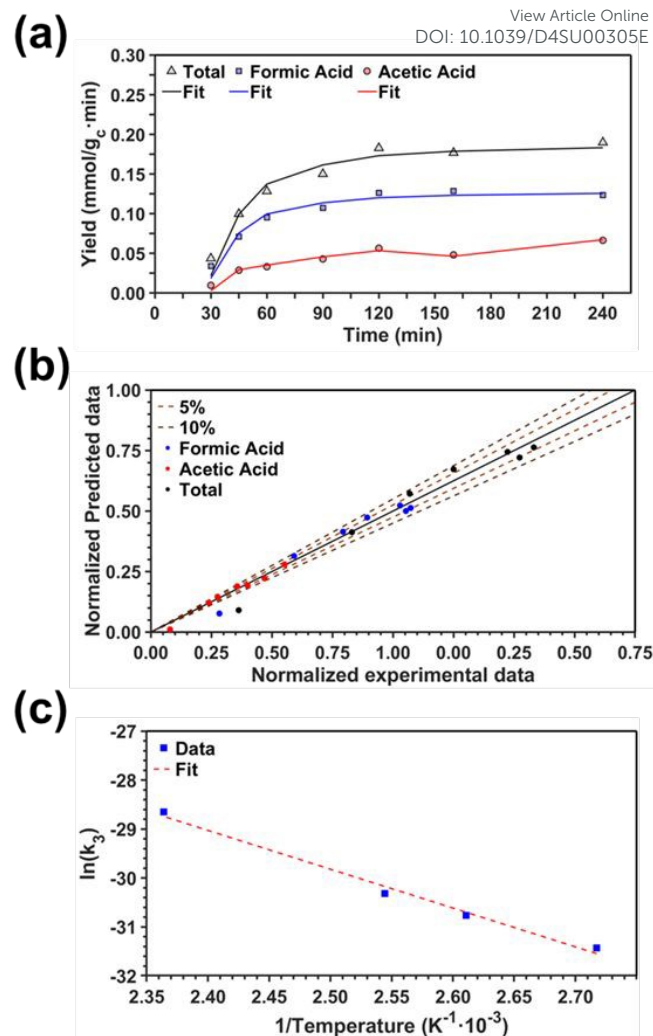
Due to the resemblance among some results (Table 2) and the mathematical restrictions, the final expression for the rate of total CO<sub>2</sub> fixation reaction was expressed as follows considering A2:

$$r = k_3(T) \cdot [\text{CO}_2^{*b}] \cdot [\text{H}_3\text{O}^+] \cdot \theta_a \cdot \theta_b \cong \\ \cong k_3(T) \cdot k_1 \cdot 2 \cdot k_6 \cdot [\text{HCOOH}] \cdot [\text{H}_2\text{O}] \cdot \theta_b \cdot \theta_a \quad (10)$$

where  $k_3(T)$  is the specific constant of the reaction rate ( $\text{mmol}_{\text{product}} \cdot \text{g}_{\text{cat}}^{-1} \cdot \text{s}^{-1}$ ),  $k_1$  is the water splitting reaction constant ( $\text{m}^3 \cdot \text{mmol}^{-1}$ ),  $k_6$  is the formic acid desorption constant ( $\text{mmol} \cdot \text{m}^{-3}$ ) and  $[\text{HCOOH}]$  and  $[\text{H}_2\text{O}]$  are the concentrations of the formic acid and water, respectively ( $\text{mmol} \cdot \text{m}^{-3}$ ).

The activation energy  $E_a$  was obtained using the temperature studies reported in Figure 3e and the Arrhenius relation (Eq S2). Analysis of the kinetic models supports the conclusions derived from the behavior observed in Figure 3e. Thus, the linear increase of formic acid with temperature, while acetic acid remains stable, is in agreement with assumptions A2 and A3.

Figure 5a displays the fittings obtained from both total carbon fixation and acetic acid models in conjunction with the experimental results. For the sake of completeness and due to their intrinsic similarities, the same model (Eq 10) was used to fit the synthesis of formic acid alone. In order to show the good correlation between the experimental data and the points estimated with the kinetic model, a parity plot is



represented in Figure 5b. As it can be observed, the predicted model shows an excellent correlation with the experimental

**Figure 5.** Comparison between the experimental data and the fitting model: (a) fitting of the different experimental yields obtained from the kinetic models proposed; (b) parity plot; and (c) linearization of the Arrhenius expression.

**Table 3.** Kinetic parameters with their confidence intervals of the CO<sub>2</sub> fixation reaction over p-HAp.

Parameter	
$k_1$	$15.17 \pm 1.5 \text{ m}^3/\text{mmol}$
$k_6$	$0.37 \pm 0.06 \text{ mmol}/\text{m}^3$
$k_3(T=95 \text{ }^\circ\text{C})$	$1.43 \pm 0.23 \cdot 10^{-2} \text{ mmol}_{\text{product}}/(\text{g}_{\text{c}} \cdot \text{s})$
$k_0$	$2.20 \pm 0.07 \cdot 10^7 \text{ mmol}_{\text{product}}/\text{g}_{\text{c}} \cdot \text{s})$
$E_a$	$66 \pm 1 \text{ kJ}/\text{mol}$

results without showing any systematic deviations. Thus, almost all of the points are found inside the  $\pm 10\%$  interval range, with more than 50% of the points located in the  $\pm 5\%$  interval range. Finally, Figure 5c proves the good correlation ( $R^2 = 0.989$ ) between the experimental data and the Arrhenius relation.

The resulting kinetic parameters are reported in Table 3.  $k_3(T=95 \text{ }^\circ\text{C})$  presents the smallest value (supporting the RDS A2 assumption), while  $k_1$  (related with the water splitting



reaction) is two orders of magnitude higher than the RDS. The value for the activation energy ( $E_a = 66 \pm 1$  kJ/mol) falls within the range of values reported in the literature (40 – 100 kJ/mol).<sup>92-95</sup> This can be considered as an outstanding result, as upp-HAp (a biocompatible and cheap ceramic) shows the same catalytic potential than conventional metal-based catalysts (e.g. Cu- and Ni-based)<sup>93,94</sup> with the great advantage that no electric potential (which has been demonstrated to clearly affect the  $E_a$ ) is applied.

## Conclusions

We have shown the use of metal-free upp-HAp as catalyst for the continuous production of formic acid and acetic acid using wet  $\text{CO}_2$  as reagent. Firstly, the distribution of reaction products in the gas, liquid and solid phases, as a function of the reaction time, were investigated for the  $\text{CO}_2$  fixation heterogeneous catalytic process using mild reaction conditions (95-120 °C) and both  $\text{CO}_2$  gas (6 bar) and liquid water (20 mL) as reagents. Results evidenced the key role of upp-HAp as catalytic substrate able to adsorb  $\text{CO}_2$  extremely high ratio at regime temperature and the water in the formation of chemical valuable products that were found to be C1, C2 and C3 reflecting the competition of two processes: the reduction of  $\text{CO}_2$  and the formation of C–C bonds.

A noticeable increment of the reaction yield was observed when the  $\text{CO}_2$  conversion process was transferred from batch to continuous-flow mode, using identical catalysts. Certainly, the yield obtained ( $> 2$  mmol/g<sub>c</sub>·min) supposes a great step forward for the use of competitive green heterogeneous catalyst under strictly mild conditions. Two factors were found to be crucial for the continuous-flow process: 1) the addition of liquid water at the beginning of the process, which can provide an excess of water molecules compared to the process using wet  $\text{CO}_2$  as a source of hydrogen, and therefore, enormously increasing the yield of the reaction; and 2) the reaction temperature, depending on whether it is performed above or below 100 °C, since it drastically alters the liquid-gas balance of the water molecules coming from the liquid supplied at the beginning of the process and/or from the wet  $\text{CO}_2$ , affecting the distribution of the reaction products and the selectivity.

The thermodynamic and kinetic studies performed for the continuous-flow process without considering liquid water as reactant have provided information on the relevant reaction steps of the mechanism, primarily pointing out the first  $\text{CO}_2$  fixation step as the RDS. Although the C–C bond formation is extremely challenging, the kinetic model highlights the fast and independent generation of formic acid (C1 product), boosting the total  $\text{CO}_2$  fixation yield. Besides, the competitive adsorption of species in the binding sites has exhaustively been examined, discarding the contribution of proton adsorption to the overall kinetics, which represents a great advantage in terms of yield optimization and industrial scalability. Finally, the activation energy for the total  $\text{CO}_2$  fixation, which has been estimated to be  $66 \pm 1$  kJ/mol, is within the range of conventional electro-assisted catalysts. In

this sense, NAP-XPS experiments further confirm the pathway mechanism for  $\text{CO}_2$  activation occurring in the basic sites of upp-HAp ( $\text{PO}_4^{3-}$  and  $\text{OH}^-$ ). Therefore, such preliminary results suggest a synergy between the different binding sites (i.e. dual site catalysis) which would promote the generation of C2 and C3 products. Indeed, the outstanding catalytic activity of upp-HAp for  $\text{CO}_2$  fixation and to generate organic low carbon molecules has been demonstrated, representing a feasible, highly efficient, sustainable, green, cheap, and energetically low-demanding alternative to conventional electro-assisted catalysts.

## Conflicts of interest

Authors declare that the preparation and application of permanently polarized hydroxyapatite as catalyst was patented by the Universitat Politècnica de Catalunya and B. Braun Surgical S.A. (EP16382381, EP16382524, P27990EP00, PCT/EP2017/069437, P58656 EP, P59205 EP, P59091 EP, P59008 EP, P59528 EP, 2021P00074 EP, P 59985 EP).

## Data availability

The data supporting this article have been included as part of the Supplementary Information.

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## Data Availability Statement

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The data supporting this article have been included as part of the Supplementary Information.

