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# Engineering L-asparaginase for spontaneous formation of calcium phosphate bioinspired microreactors†

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Active bioinspired materials are appealing biotechnological targets, and their study is gaining momentum. These materials, which comprise of an inorganic matrix and one or more biomolecules, are extremely variable and therefore may result difficult to characterize in their intimate structure. In this work we have prepared a hydroxyapatite–L-asparaginase composite, with the perspective of using it in acute leukemia treatment. We demonstrate that the use of electron microscopy and powder X-ray diffraction, combined with the atomic-resolution information coming from solid-state NMR, allows us to understand the topology of the material and how the different components interplay to obtain an active composite.

## Introduction

Biological drugs (biologics<sup>1</sup>) are the fastest-growing category of approved therapeutics, with most biologics being represented by proteins.<sup>2</sup> L-Asparaginases, which catalyze the hydrolysis of L-asparagine (Asn) to L-aspartate (Asp), are one of the earliest examples of biologics. The depletion of Asn from circulating blood induces cell death in acute lymphoblastic leukemia (ALL) cells, which lack the Asn-synthetase enzyme.<sup>3,4</sup> The most commonly used biologics in therapy are type-II bacterial

L-asparaginases (ANSII), with *E. coli* ANSII being in clinical use since 1967.<sup>5</sup> ANSII is indispensable in ALL treatment protocols and is used for remission induction and intensification in all pediatric and most adult regimens.<sup>6</sup> However, treatment with ANSII can lead to severe side effects that comprise hepato- and pancreas toxicity, coagulation disorders and, most relevant, allergic reactions. ANSII can be administered both intravenously (IV) or intramuscularly (IM), the latter route being better tolerated and less often resulting in increased hypersensitivity reactions.<sup>7</sup> The relevance of antibodies directed against ANSII in children with ALL has been highlighted in the Berlin–Frankfurt–Münster (BFM) studies, and recently reviewed:<sup>8</sup> adverse reactions to the enzyme mediated by either IgG or IgE antibodies, or both, have been described.<sup>9</sup> The appearance of antibodies increases ANSII clearance and neutralizes the catalytic activity of the enzyme.<sup>10</sup> Therefore, it is of capital importance to reduce the interaction of the enzyme with the immune system of the patients. Several strategies have been devised to reduce immunogenicity, the most common being PEGylation.<sup>11</sup> However, PEG may cause severe side effects in patients with progressive chronic kidney disease<sup>12</sup> and can induce allergic reactions even when administered IM.<sup>7</sup>

A radically different strategy would be to avoid contact with the immune system of the patient altogether. Extracorporeal processing of blood has been proposed almost forty years ago, with limited follow-up in clinical practice, mostly because of the limited compliance of the treatment.<sup>13</sup> Nowadays, better-designed biomedical devices that can be used to physically confine active components within a membrane are available.

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This possibility mitigates the need of obtaining an enzyme that is able to evade immune recognition. On the other hand, the incorporation of an enzyme within a device requires that the formulation has a long lifetime, both on the shelf and *in operando*, that it is easily handled (weighed, transferred, *etc.*) and that the active site is accessible. For these reasons, the possibility of extracorporeal processing has triggered a strong interest in the protein immobilization community.<sup>14,15</sup>

Here, we describe the design of a biomaterial that carries an ANSII chimera attached to a hydroxyapatite (HA) support, because HA-based composites have already been used as new biomedical devices,<sup>16–18</sup> drugs and vaccine carriers,<sup>19–24</sup> and because immobilization of enzymes often results in stabilization.<sup>25</sup>

The characterization of bioinspired materials is a complex accomplishment because it needs to rely upon the characterization of both the inorganic matrix and the biomolecular component.<sup>26–31</sup> In this context, NMR plays a very important role, as it is the only methodology that can access atomic-level information on both components.<sup>32–41</sup>

## Results and discussion

Native, therapeutically active ANSII is a homotetrameric assembly of about 140 kDa with D<sub>2</sub> symmetry (a dimer of two intimate dimers)<sup>42</sup> with four catalytic sites per tetramer. We have previously provided the NMR characterization of ANSII in its free, PEGylated,<sup>43</sup> sugar-conjugated forms<sup>44</sup> and bound to gold nanoparticles.<sup>45</sup> We fused ANSII with a C-terminal tail encompassing the HA-promoting peptide (HABP) W6p (RWRLEGTTDDKEEPESQRRIGRFG; Fig. S1, ESI<sup>†</sup>).<sup>46</sup> Gel-electrophoresis and dynamic light scattering show that ANSII-HABP is smaller (~80 kDa) than the native enzyme, and likely corresponds to a single intimate dimer (Fig. S2–S4, ESI<sup>†</sup>). However, it still retains about one-fourth of the catalytic activity per monomer (Fig. S5, ESI<sup>†</sup>).

The HA/ANSII-HABP composite was prepared by self-promoted biomimetic mineralization of HA driven by HABP (see ESI<sup>†</sup>). After repeated washings to remove the unbound enzyme, the activity of HA/ANSII-HABP was assayed using NMR (Fig. S6 and S7, ESI<sup>†</sup>), under the same conditions used for metabolomic profiling of serum (Fig. S7, ESI<sup>†</sup>).<sup>47</sup> HA/ANSII-HABP was still active after more than three months of storage at 4 °C. The efficacy of HA/ANSII-HABP was then tested *in vitro* on human ALL cells (697 cell line), and compared to either the wild-type (WT) enzyme or the commercial product Oncaspar (*i.e.* PEG-ANSII). All ANSII preparations were added at 0.74 U mL<sup>-1</sup> concentration, according to previous reports.<sup>48,49</sup> HA/ANSII-HABP was kept separated from leukemic cells *via* a 3.5 kDa cut-off dialysis membrane. HA/ANSII-HABP reduced the leukemic cell viability to the same extent (after 24 hours of incubation) and even more (at 48 hours of incubation) compared to either the free enzyme or the commercial product (Fig. 1).

HA/ANSII-HABP was further characterized using powder X-ray diffraction (XRD), scanning electron microscopy and solid-state NMR. The XRD profile (Fig. 2), shows the typical diffraction pattern of HA,<sup>50</sup> while the diffraction at low  $2\theta$  suggests the simultaneous presence of some octacalcium phosphate (OCP).<sup>51</sup>

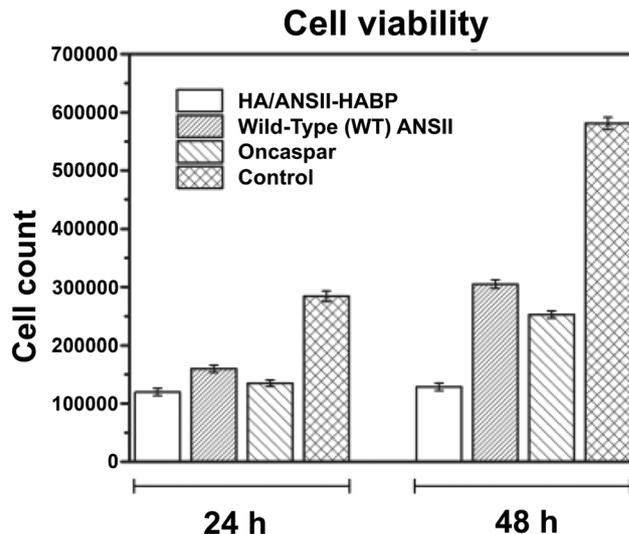


Fig. 1 Effects of different ANSII preparations on the proliferation of ALL cells. ALL 697 cells were treated with different ANSII preparations (HA/ANSII-HABP, WT ANSII and Oncaspar) at a concentration of 0.74 U mL<sup>-1</sup> for 24 and 48 hours and the number of live cells, determined *via* the Trypan Blue exclusion test, was measured. HA/ANSII-HABP was confined in a device with walls consisting of a 3.5 kDa cut-off dialysis membrane. Data are mean values  $\pm$  standard error of the mean of three independent experiments. 48 hours: HA/ANSII-HABP vs. WT ANSII,  $p = 0.00116$ ; HA/ANSII-HABP vs. Oncaspar,  $p = 0.00449$ , as tested by one way ANOVA.

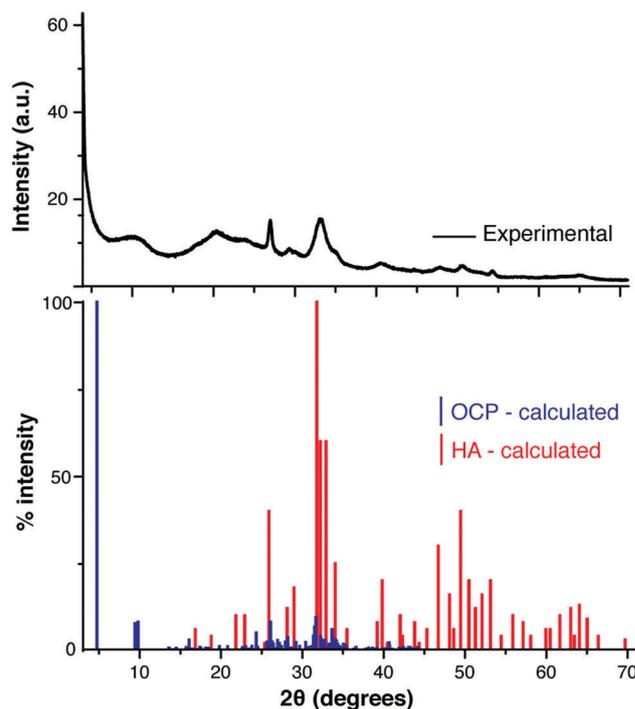
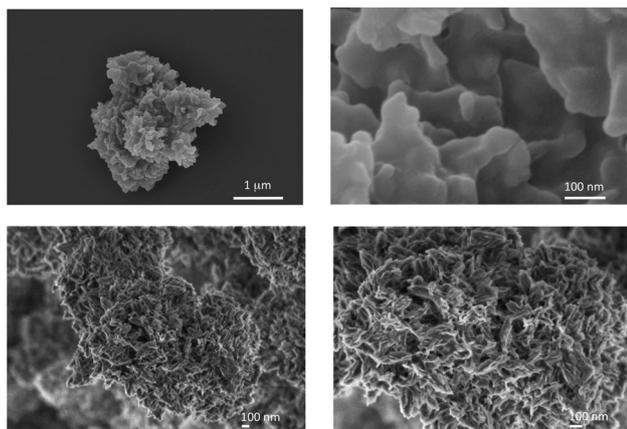


Fig. 2 Top: XRD profile of the HA/ANSII-HABP composite (black line). Bottom: Calculated XRD diffractogram of hydroxyapatite (red) and octacalcium phosphate (blue) shown as stick plots.

The diffraction peaks are broad, as expected for crystallites of small size. By elemental analysis, the protein content was found to be on the order of  $40 \pm 1\%$  in the dried material.





**Fig. 3** SEM images of the HA/ANSII-HABP composite, taken with (upper panels) and without (lower panels) conductive staining, revealing a highly porous structure, which is most suitable for maintaining the accessibility of the enzyme. The high protein loading at the surface reduces the achievable resolution.

SEM micrographs show that the material is composed of fused structures, homogeneous in shape and size (Fig. 3), which more closely resemble HA nanosheets,<sup>52</sup> rather than OCP spherical particles.

Combustion elemental analysis provided the elemental composition and a quantitative determination of the protein content, found to be of the order of 40% in the dried material. After reacting 1 mL of the mixture, and rinsing it several times, about 28 mg of wet material are obtained. From the content of the reaction mixture (see ESI†), the theoretical yield of the dry material can be calculated: assuming that only HA will form, with a 40% w/w protein content in the material, 4 mg of the composite should precipitate. If, on the contrary, only OCP is formed, about 2 mg of the composite should precipitate. Thus, in the wet material, we can estimate that 6.5–13% will consist of HA/ANSII-HABP.

HA/ANSII-HABP was further characterized in terms of their elemental composition using energy dispersive X-ray spectroscopy (EDS). Inductively coupled plasma (ICP) emission was additionally used for elemental analysis of the total Ca and P content in the sample (Table 1). The ICP measurement shows a Ca/P ratio of 1.78, higher than the theoretical HA value of 1.67 (Tables 1 and 2). Such a phosphate depletion has been also observed<sup>53</sup> for HA prepared in the presence of a mineral binding peptide, and is possibly due to a disordered phase in addition to the ordered HA phase.<sup>53</sup>

In order to fully understand the activity of the material, it is necessary to link the bulk observations to the molecular scale, answering the following questions:

- are the ordered and the disordered layers in contact with each other?
- is the protein in contact with the ordered phase, with the disordered one or with both?
- is the protein exposed to the solvent, thus accessible to the substrate?

We here demonstrate how solid-state NMR is the methodology of choice to answer these questions.

**Table 1** Secondary electron emission analysis of HA/ANSII-HABP

Element	Average wt%	Std%	Average atom count	Std
C	23	3	1.9	0.2
N	10	3	0.7	0.2
Ca	26	4	0.7	0.1
P	14	2	0.44	0.06

**Table 2** Summary of the elemental composition of HA/ANSII-HABP, compared with expected values

Ratio	Measured – EDS	Measured – ICP	Expected for HA	Expected for OCP
Ca/P	1.6 ± 0.3	1.78	1.67	1.33
Ratio	Measured	Expected for current protein chimeric construct		
C/N	2.8 ± 0.9	3.57		
Protein%	47 ± 7			

A Bloch decay (BD), a homonuclear decoupled <sup>1</sup>H spectrum of HA/ANSII-HABP and a BD spectrum of HA are shown in Fig. 4. The presence of the HA phase is confirmed by the BD spectrum (red), where HA/ANSII-HABP has the same OH<sup>-</sup> resonance at 0.2 ppm as synthetic HA (green). The BD spectrum shows a broad intense water resonance at 5.3 ppm (consistent with the high water content in HA/ANSII-HABP); however, as will be shown, these water molecules are not necessarily residing at the enzyme–mineral interface. The peak at 1.3 ppm is associated with the aliphatic side-chain protons of the protein chimera but may include some contribution from HA as can be seen by comparing with the mineral spectrum. In the homonuclear decoupled wPMLG spectrum (violet) of HA/ANSII-HABP, the water line is extensively attenuated, exposing three clear bands at 1.3 ppm, 4.5 ppm and 7.2 ppm from ANSII-HABP protons.

Although the wPMLG experiment is intended to enhance resolution by <sup>1</sup>H–<sup>1</sup>H decoupling, in this biomaterial sample we find that it improves the capability to observe the protein proton lines *via* a simple solvent suppression mechanism. Exposing the protons to a strong effective field tilted at the magic angle causes exclusive broadening of the water signal beyond detection due to coincidental matching of the water motional time scale with the field's decoupling cycle period (15 μs). The motions experienced by ANSII-HABP are quite different and therefore are not affected similarly by the homonuclear decoupling. The widths of the ANSII-HABP proton lines are 3–6 ppm and they result from immobile protein molecules: these lines are collectively associated with ANSII-HABP protons belonging to side-chains, H $\alpha$  and amide protons. The peak at 1.3 ppm may mask a smaller contribution from HA (see the spectrum of synthetic HA).

The deconvolution<sup>54</sup> of <sup>1</sup>H–<sup>31</sup>P cross-polarization (CP)<sup>53</sup> spectra of HA/ANSII-HABP obtained under MAS (Fig. 5a and Fig. S8, ESI†) reveals two phosphate species, as synthetic HA,<sup>53</sup> that can be attributed to two different layers. The peak at 2.9 ppm (FWHM 1.2 ppm) is typical of the phosphate resonance in HA crystals (denoted hereafter as P<sub>HA</sub>) while the peak at



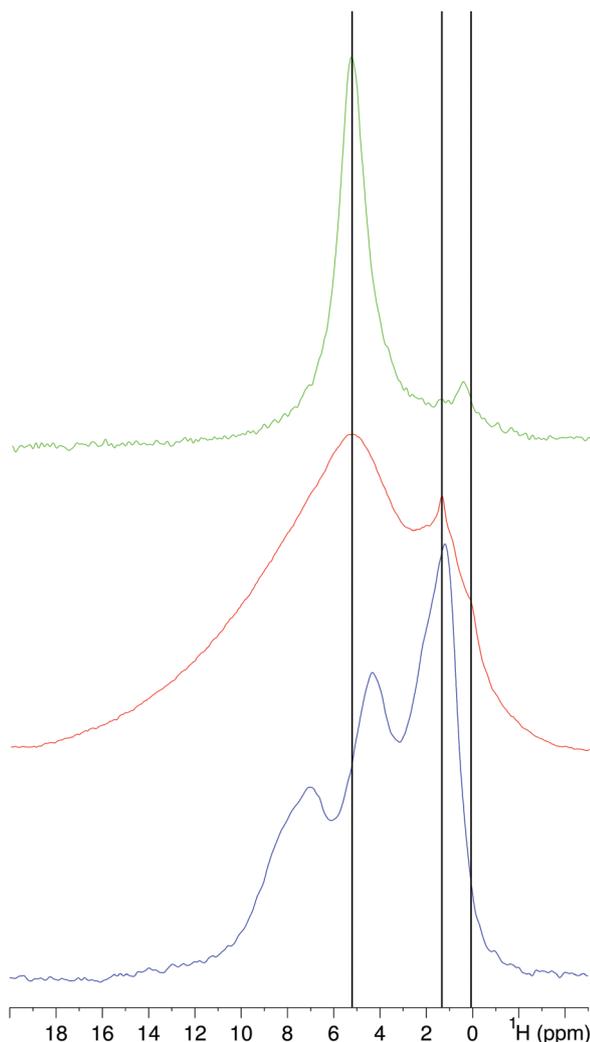


Fig. 4  $^1\text{H}$  NMR spectra of ANSII-HABP acquired using wPMLG for homonuclear-decoupling (violet) and a simple  $90^\circ$ -acquisition (*i.e.* a Bloch decay) (red). A  $^1\text{H}$  Bloch decay spectrum (green) of synthetic hydroxyapatite prepared without calcination is also shown for comparison.

2.6 ppm (FWHM 3 ppm) is attributed to hydrogen phosphate ions ( $\text{HPO}_4^{2-}$ ) (denoted hereafter as  $\text{P}_{\text{HP}}$ ) which are part of the disordered calcium phosphate phase that resides at the HA/ANSII-HABP interface<sup>55</sup> (this peak cannot be attributed to beta-glycerophosphate because it exhibits a higher chemical shift<sup>56</sup>). Mono-acid phosphates were shown to adopt a large range of chemical shifts, depending on the surroundings ions. In HA/ANSII-HABP this is reflected in the significant width, 7 ppm (FWHM), of the  $\text{P}_{\text{HP}}$  peak.<sup>55</sup>

To characterize the proximity of interfacial and bulk protons and phosphates,  $^{31}\text{P}$  CP build-up curves were analyzed.<sup>53</sup>  $^{31}\text{P}$  CP build-up measurements were carried out by varying the contact time from 100  $\mu\text{s}$  to 10 ms, and the intensities of the  $\text{P}_{\text{HA}}$  (black) and  $\text{P}_{\text{S}}$  (red) peaks were plotted against the contact time (see Fig. 5b). Recently, similar magnetization build-up curves were used to isolate magnetization transfers between interfacial protons and phosphates and bulk protons and phosphates in apatite samples prepared with a bone

protein binding peptide which displays similar properties of having a crystalline phase of HA and a disordered phase containing calcium phosphate ions and the biomolecule.<sup>53</sup> Here, we follow a similar strategy.

These intensity buildup curves show the rate of magnetization transfer from adjacent protons to the two phosphate species. The relative line intensity in Table 3 is derived from the fit to the experimental phosphate line, in Fig. 5 as described below. The integral (integrated intensity) of the experimental phosphate peak for each contact time point was calculated and the values were normalized to the highest integral to obtain the buildup normalized to 1. Each spectrum was then deconvolved into the two phosphate contributions and the calculated area under each deconvolved peak was taken as a fraction of one. Therefore, the intensity per phosphate peak and per contact time point was overall normalized to the fractional intensity of that phosphate species within each contact time spectrum and to the intensity of the highest signal in the CP buildup curve. The curve of  $\text{P}_{\text{HA}}$  was fitted using eqn (1) and that of  $\text{P}_{\text{HP}}$  was fitted using the extended eqn (2) which accounts for relaxation processes which further influence transfer.

$$I(t) = I_0 \times \left[ 1 - e\left(-\frac{t}{\tau_{\text{CP}}}\right) \right] \quad (1)$$

$$I(t) = \frac{I_0}{\left(1 - \frac{\tau_{\text{CP}}}{T_{1\rho}^{\text{H}}}\right)} \times \left[ e\left(-\frac{t}{T_{1\rho}^{\text{H}}}\right) - e\left(-\frac{t}{\tau_{\text{CP}}}\right) \right] \quad (2)$$

In these equations,  $I_0$  is the overall intensity of phosphate magnetization,  $\tau_{\text{cp}}$  is the typical CP time and ( $T_{1\rho}^{\text{H}}$ ) is the longitudinal rotating frame relaxation time of the protons.

The kinetic parameters, summarized in Table 1, were deduced from best-fitted curves calculated by minimizing the deviations of signal intensity  $I_0$ , the magnetization transfer time constant ( $\tau_{\text{cp}}$ ) and the  $^1\text{H}$  rotating frame relaxation time ( $T_{1\rho}^{\text{H}}$ ) in calculations and in experiments.

The relatively low normalized  $\text{P}_{\text{HA}}$  intensity, 0.27, is indicative of the low content of crystalline HA in ANSII-HABP. This line typically has an intensity  $>0.6$  in other co-precipitation procedures of proteins with HA.<sup>53</sup> The extracted magnetization transfer time of  $\text{P}_{\text{HA}}$ , 3.1 ms (Table 3), closely resembles the value for synthetic HA (see Fig. 6),<sup>53</sup> and is somewhat intermediate between hydroxyl-to-phosphate transfer time in apatite crystals grown with an osteonectin binding peptide and interfacial transfer time from protons outside the crystals to crystalline phosphates in that same sample.<sup>53</sup>

This finding suggests that either water or protein molecules are located close to the faces of the HA crystal and transfer magnetization to ordered (crystalline) phosphate ions in exposed crystal surfaces. The transfer rate of the  $\text{P}_{\text{HP}}$  line, 0.5 ms, is similar to the magnetization transfer observed for interfacial phosphates in the aforementioned apatite grown with an osteonectin binding peptide.



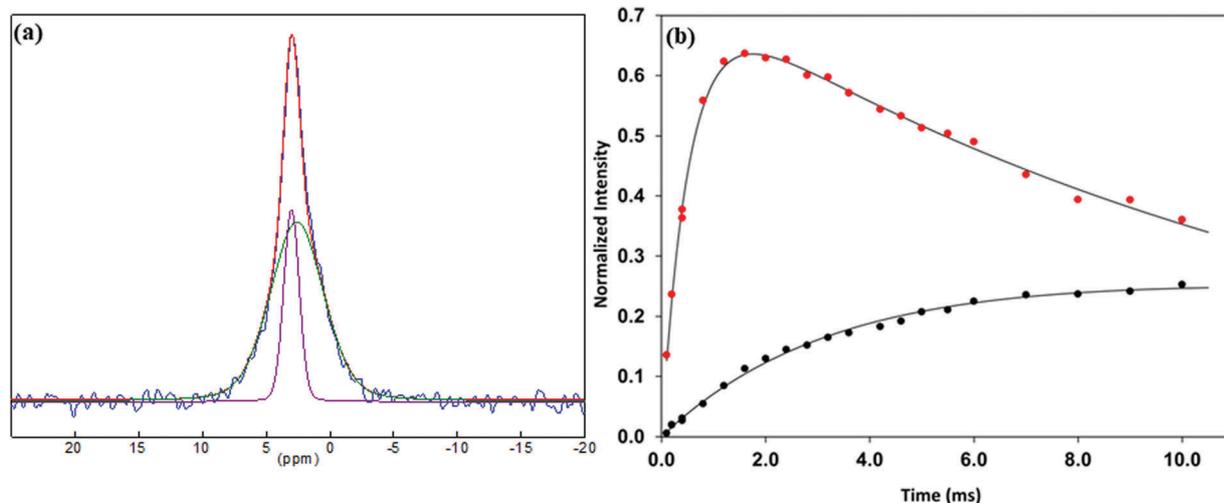


Fig. 5 (a) An example of the  $^{31}\text{P}$  CP spectrum of HA/ANSII-HABP obtained using 3.6 ms contact time deconvoluted using the DMFIT program. The experimental spectrum (blue) is shown with the sum of simulated peaks (red) and individual lines at 2.6 ppm (green) and 2.9 ppm (purple). (b)  $^{31}\text{P}$  intensity buildup with increasing CP contact time of  $\text{P}_{\text{HA}}$  phosphates (black) and  $\text{P}_{\text{HP}}$  phosphates (red).

Table 3 Kinetic parameters of  $^{31}\text{P}$  magnetization buildup of phosphates in HA/ANSII-HABP

Peak position (ppm)	Assignment	Width (ppm)	$\tau_{\text{cp}}$ (ms)	$T_{1\rho}$ (ms)	Normalized intensity
2.9	$\text{P}_{\text{HA}}$	1.4	3.1	—	0.27
2.6	$\text{P}_{\text{HP}}$	5.0	0.5	13.1	0.73

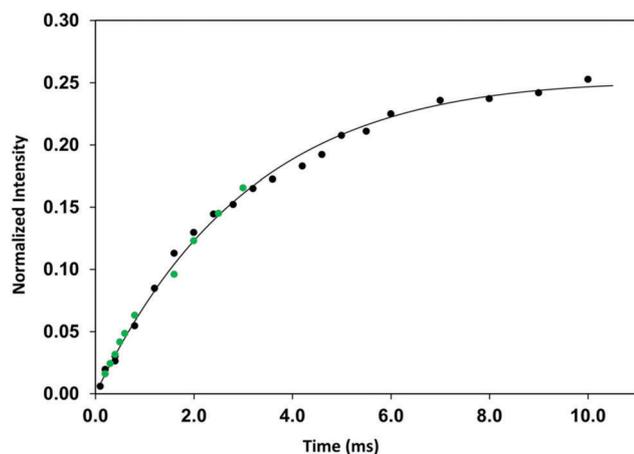


Fig. 6  $^{31}\text{P}$  CP buildup curves of  $\text{P}_{\text{HA}}$  (black) and of the HA phosphate line in hydroxyapatite prepared via a titration method at 310 K (green<sup>53</sup>). It shows that the rate of magnetization buildup of  $\text{P}_{\text{HA}}$  in ANSII-HABP is similar to that of HA, suggesting that this line represents a  $^{31}\text{P}$  nucleus in a crystalline environment.

To further probe the local environment sensed by phosphate groups and identify their distribution within the sample, we recorded two-dimensional  $^1\text{H}$ - $^{31}\text{P}$  HETCOR MAS NMR spectra (Fig. 7).

During the  $^1\text{H}$  magnetization evolution under the chemical shift in  $t_1$ , the PMLG decoupling sequence was used to suppress  $^1\text{H}$ - $^1\text{H}$  dipolar couplings. Fig. 7 shows an intense cross peak

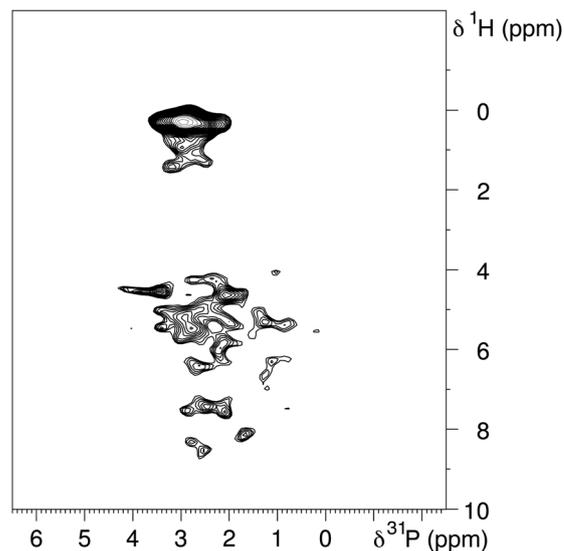


Fig. 7  $^1\text{H}$ - $^{31}\text{P}$  heteronuclear correlation (HETCOR) spectra of HA/ANSII-HABP using contact times of 0.5 ms.

between HA hydroxyl protons at 0.2 ppm and phosphates at 2.9 ppm (*i.e.*  $\text{P}_{\text{HA}}$ ). It also shows a group of narrower cross peaks correlating  $^1\text{H}$  at 1.3 ppm, 4.0–6.5 and 7.0–8.5 ppm along F1 with  $\text{P}_{\text{HP}}$  phosphates. These resonances are attributed to the protein protons that transfer magnetization to proximal  $\text{P}_{\text{HP}}$  phosphates in the mineral. The homonuclear decoupling employed in  $t_1$  effectively removes water proton excitation, and therefore no correlations are observed between water and phosphate species. This is quite different from other preparations of apatite minerals,<sup>53,57</sup> whereby a strong correlation of water protons with phosphates in a disordered calcium phosphate layer on HA crystallites was observed. The absence of the water resonance in the 2D HETCOR spectra is further demonstrated in the  $^1\text{H}$  projections (taken along the phosphate maximal intensity) of



these measurements as compared to similar measurements in synthetic HA where the relative intensity is much different (see Fig. S9, ESI†). Similar  $^1\text{H}$ - $^{31}\text{P}$  2D HETCOR measurements without PMLG decoupling did not show any water cross peaks with phosphates, so it is apparent that water molecules are distantly located from phosphates.

In summary, these spectra show that the protein resonances correlate with  $P_{\text{HP}}$  in the disordered layer, which in turn is connected with the  $P_{\text{HA}}$  phase. The latter crystalline phase is mostly excluded from water, suggesting also that this bulk mineral phase is overall distant from the biomolecule. From these data, we can conclude that ANSII-HABP is interacting with a layer of interfacial phosphates ( $P_{\text{HP}}$ ), probably of the mono-acid type, and has no direct interactions with HA crystallites. Finally, no correlations are here observed between water and phosphate species: this means that the water molecules are either located far from the phosphates or are too dynamic to be detected in dipolar-based magnetization transfer experiments.

The  $^1\text{H}$ - $^{13}\text{C}$  CP<sup>58,59</sup> MAS spectrum with ramped  $^1\text{H}$  amplitude (ramp-CP<sup>60</sup>) has low intensity and resolution (Fig. 8, middle trace) that are both increased when ramp-CP is hybridized with the nuclear Overhauser effect (NOE, Fig. 8, top trace):<sup>61</sup> this sequence results in the appearance of a larger number of signals. Most of these additional peaks appear in the aliphatic region (0–75 ppm) and some peaks from aromatic side chains also appear (110–140 ppm). Moreover, some new peaks can be observed also in the carbonyl region (150–180 ppm). The observed peaks around 153 ppm belong to  $C_\zeta$  of the arginine side-chains, while those around 178 ppm relate to  $C_\gamma$  or  $C_\delta$  of

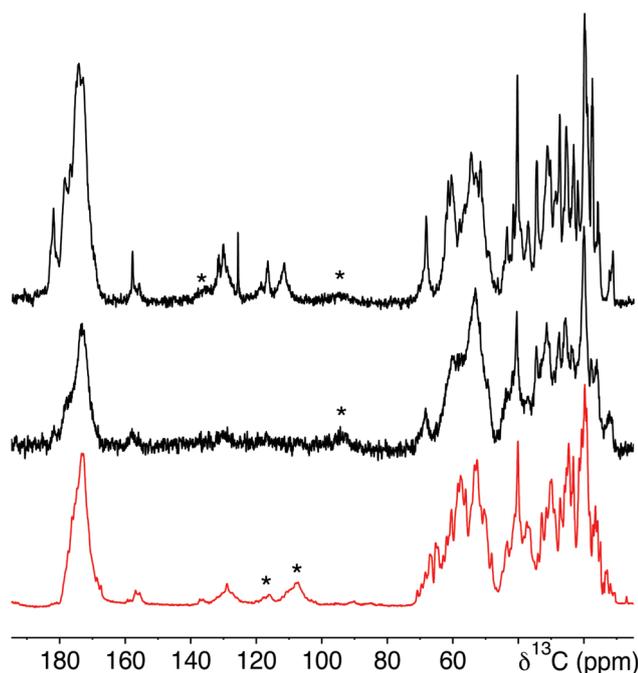


Fig. 8  $^{13}\text{C}$  NMR spectra of HA/ANSII-HABP (black) and PEGylated wtANSII<sup>43</sup> (red) obtained under MAS using CP only (middle and bottom spectra) or CP-NOE (topmost). The spinning sidebands are marked as '\*'. NOE mixing was set to 1.3 s and the CP contact time was 1.2 ms.

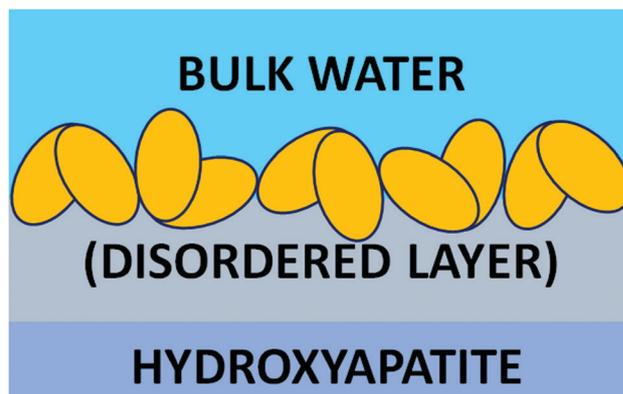


Fig. 9 Scheme of the arrangement of the components of HA/ANSII-HABP.

the aspartate and asparagine or glutamate and glutamine side-chains, respectively (Fig. S10, ESI†). These observations suggest that the additional peaks, observed *via* NOE, originate from those chemical groups or amino acid residues that undergo significant motion on the milliseconds (or faster) timescale, and are therefore not detectable by the dipolar-coupling based CP. One needs to consider the hypothesis of a substantial water-protein magnetization transfer (either through intermolecular NOE, which is prevailing at high spinning rates and low temperature, or through chemical exchange followed by spin diffusion, which is most probably dominant under the present experimental conditions<sup>62</sup>) at the time of the NOE mixing, plus the contribution due to direct  $^{13}\text{C}$  excitations which is intrinsic for the pulse sequence applied.<sup>61</sup> Either of the two mechanisms does not detract from the idea, but rather corroborates it, that a large share of the protein is sizably exposed to water.

The topology of HA/ANSII-HABP (Fig. 9) that can be derived from our integrated analysis is summarized below.

(a) The microscopic characterization demonstrates that the material is composed of small lamellar structures that closely resemble hydroxyapatite, and PXRD confirms this conclusion.

(b) The secondary electron analysis shows that there is a phosphate depletion, consistent with previous reports on HA/peptide composites.

(c) Solid-state NMR investigation of the inorganic part shows that the ordered HA phase is not in contact with the protein and the disordered layer is in intimate contact with the protein. This further shows that the two inorganic phases are in contact with each other, and water is closer to the protein and largely excluded from the mineral phases.

(d) Protein solid-state NMR indicates that the enzyme is not completely immobile in the material, but undergoes extensive motions (Fig. S10, ESI†), suggesting that it is mainly sitting at the surface of the particles, rather than being entrapped within the material (Fig. 9). Despite the preserved mobility, the observed chimeric ANSII-HABP is stably linked to the mineral particles, as discussed above.

(e) The tests against the cells and those *in vitro* confirm that the protein is stable in the construct for several months and demonstrate that the material is active.



Nowadays ANSII is still indispensable<sup>6</sup> in the treatment of ALL patients and new ANSII preparations are needed to overcome the limitations of the current ANSII therapy, especially immunogenicity. The results obtained in an *in vitro* assay on human leukemic cells with HA/ANSII-HABP, and the protocols for its characterization, provide new perspectives for the development of microreactors to be used as possible innovative biomedical devices able (i) to increase the stability of ANSII, (ii) to extend Asn depletion in the serum and at the same time (iii) virtually to reduce immunogenicity. Collectively, our results show that enzymes processing small substrates can be efficiently and profitably bound to the surface of a HA matrix by engineering the wild-type construct with HA promoting peptides.

## Author contributions

AL and MF designed the chimeric construct; AL and SG expressed and purified the protein and prepared the composite; IM, LC, ER, MF, GG, AR and CL performed and interpreted the NMR experiments; IM, AC and GG recorded and interpreted the EM data; SG, SP and AA performed and interpreted the in-cell analysis; ER, MF, AR, GG and CL designed the research; and all authors wrote the manuscript.

## Abbreviations

SSNMR	Solid-state nuclear magnetic resonance
HA	Hydroxyapatite
ANSII	Type-II bacterial asparaginase
HABP	Hydroxyapatite binding peptide W6p RWRLEGTDDKEEPESQRRIGRFG
HA/ANSII-HABP	HA-bound ANSII-HABP chimera composite
OCF	Octacalcium phosphate
CP	Cross-polarization
HETCOR	Heteronuclear correlation

## Conflicts of interest

There are no conflicts to declare.

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