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1 Collagen Suprafibrillar Confinement Drives the Activity of Acidic 2 Calcium-Binding Polymers on Apatite Mineralization

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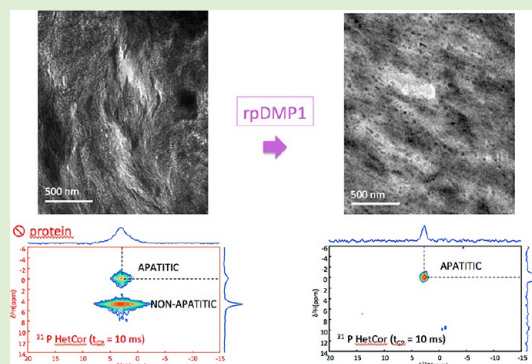


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Supporting Information

ABSTRACT: Bone collagenous extracellular matrix provides a confined environment into which apatite crystals form. This biomineralization process is related to a cascade of events partly controlled by noncollagenous proteins. Although overlooked in bone models, concentration and physical environment influence their activities. Here, we show that collagen suprafibrillar confinement in bone comprising intra- and interfibrillar spaces drives the activity of biomimetic acidic calcium-binding polymers on apatite mineralization. The difference in mineralization between an entrapping dentin matrix protein-1 (DMP1) recombinant peptide (rpDMP1) and the synthetic polyaspartate validates the specificity of the 57-KD fragment of DMP1 in the regulation of mineralization, but strikingly without phosphorylation. We show that all the identified functions of rpDMP1 are dedicated to preclude pathological mineralization. Interestingly, transient apatite phases are only found using a high nonphysiological concentration of both collagen and additives ensures specific chemical interaction of bone components in mineralization.



centration of additives. The possibility to combine biomimetic al interactions and offers perspectives for understanding the role

21 ■ INTRODUCTION

22 Bone is a complex composite material, which closely associates
23 cells and an extracellular matrix (ECM). The bone ECM
24 combines various components, that is, collagen, noncollage-
25 nous proteins (NCPs), carbonated hydroxyapatite (HA)
26 nanoparticles, and water. Although bone is one of the most
27 studied living materials, the exact role of its components,
28 noticeably the organic one, is still debated. Both intra-¹⁻³ (~40
29 nm) and interfibrillar⁴ (~1.5 nm) spaces in collagen are
30 proposed to be nucleation sites for bone mineral. The
31 interfibrillar confinement generated by the collagen matrix
32 increases as the ECM becomes mineralized, thus forming the
33 mature bone (Figure 1A) with the characteristic twisted
34 plywood pattern⁵ (Figure 1B). This suprafibrillar three-
35 dimensional (3D) architecture affects the physical properties
36 of the ECM such as mechanical response^{6,7} and microenviron-
37 ment (e.g., size and spatial distribution of apatite, local
38 hydrated environment of phosphate ions).⁸ In addition, *in vitro*
39 investigations on NCPs activity have indeed yielded contra-
40 dictory findings when the protein is studied either in solution
41 or in gel.^{9,10} Upon working with gels, loose collagen matrices
42 imply large interfibrillar spaces due to the lack of collagen
43 packing,⁸ meaning that bone is only reproduced at the fibril
44 level.^{11,12} Consequently, aside *in vivo*-based experiments,
45 biomimetic models in terms of confinement are needed to
46 ensure that investigations of NCPs activity are conducted

under native environment. As demonstration, only models mimicking both collagen density and geometry described for mature bone (250 mg/mL)⁸ tissues led to the typical apatite/collagen co-alignment (Figure 1C), but noticeably with higher concentrations of calcium ions (Figure 1D, left) than that described in extracellular fluid (ECF).¹³ Indeed, simulated body fluid (SBF) failed to produce intrafibrillar mineral; instead, apatite in the form of spherulite was found (Figure 1D, right). This result suggested that Ca-rich proteins from bone ECM concentrate ions in the gap regions.

To go further on the effect of bone ECM confinement on NCPs activity, we pursue the study by entrapping dentin matrix protein-1 (DMP1), an acidic calcium-binding polymer¹⁴ within our models. DMP1 is a phosphorylated ECM protein produced by osteocytes and odontoblasts that is commonly accepted as mediator in bone mineralization processes.^{15–17} Indeed, DMP1 knockout mice display defective mineralization of dentin^{18–21} and bone^{22,23} resembling phenotypes observed in human genetics. Interestingly, the

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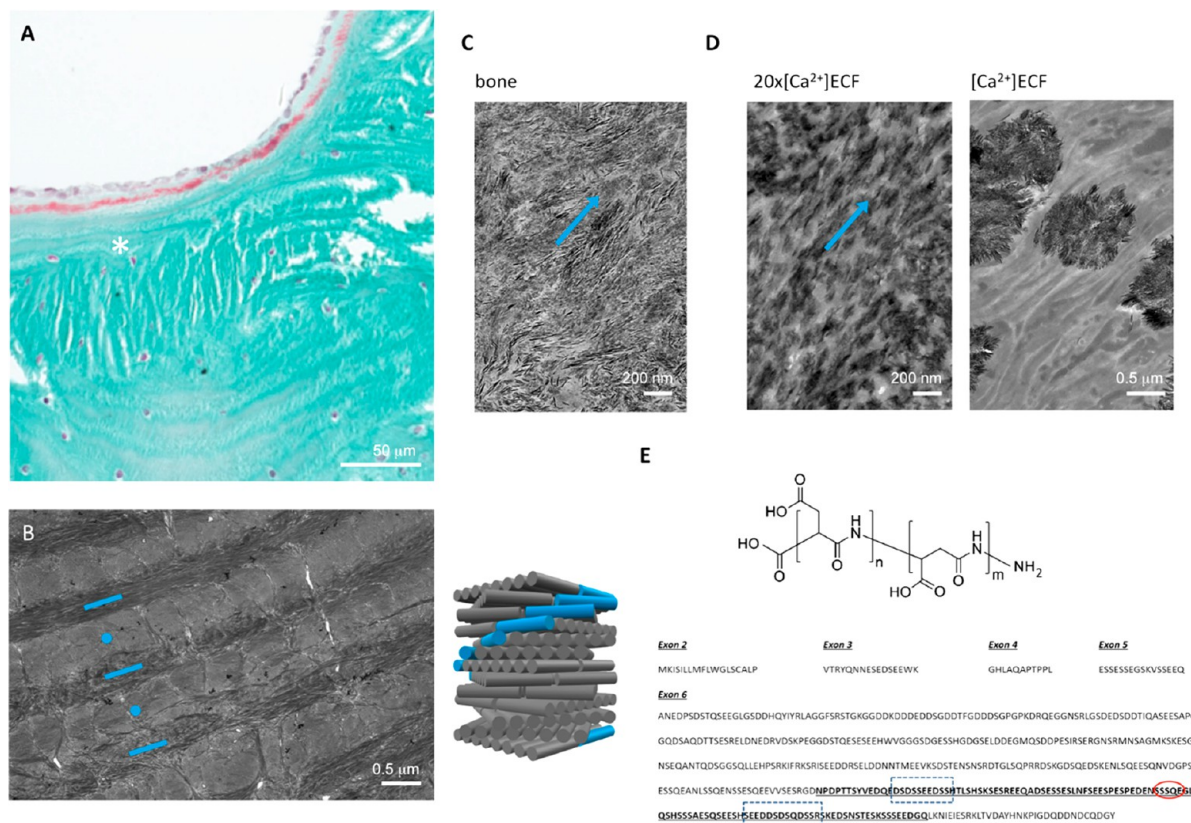


Figure 1. Comparison of collagen/apatite relationship in bone and in synthetic models reproducing the collagen confinement (density and order) found in bone. (A) Thin section of bone stained with Masson's trichrome. From top to bottom: Osteoblasts, osteoid tissue (light blue and red), and mature bone (*). Transmission electron microscopy ultrathin sections of (B) mature decalcified human bone that is characterized by a twisted plywood organization of collagen fibrils (blue dot and bars represent molecules that are perpendicular and parallel to the observation plan, respectively) schematically represented, (C) unstained sheep bone where the alignment of apatite platelets along the fibril is observed (the blue arrow shows the orientation), (D) Coll/CHA/SBF (left) and Coll/SBF/SBF (right) where $[Ca^{2+}] = 20[Ca^{2+}]ECF$ and $[Ca^{2+}] = [Ca^{2+}]ECF$, respectively. The collagen/apatite co-alignment similar to that in bone is observed in Coll/CHA/SBF (left) while spherulitic apatite crystals are obtained in Coll/SBF/SBF (right). In (E), the acidic calcium-polymer used in the biomimetic model, that is, the polyaspartate formula and the human DMP1 sequence (in bold: target peptide; blue boxed texts: collagen binding sites; red encircled text: motif found as unchanged during 220 My). The influence of organic polymers on apatite formation is studied at fixed physiological concentration of calcium, that is, $[Ca^{2+}] = [Ca^{2+}]ECF$ with two different concentrations for rpDMP1 as follows: 2.5 $\mu\text{g/mL}$ (1rpDMP1) and 25 $\mu\text{g/mL}$ (10rpDMP1).

66 role of DMP1 has been investigated *in vitro* through different
67 models (e.g., in solution,^{24,25} adsorbed onto glass plates,^{14,26} in
68 gelatin-gel systems,^{15,27} on a transmission electron microscopy
69 (TEM) grid coated with recombinant spider silks,²⁸ with
70 preassembled collagen fibrils,^{29,30} and with early calcifying
71 bone-like matrix),³¹ reaching the conclusion that the protein
72 could be involved at different levels including in apatite
73 nucleation control^{27,32} and growth inhibition,³³ in mediating
74 the size of collagen fibrils,²⁹ and in stabilizing amorphous
75 calcium phosphate (ACP) phase.^{33,34} To clarify such func-
76 tional diversity, a DMP1 recombinant peptide (rpDMP1)
77 containing those two collagen binding sites²⁹ and the peptide
78 ⁴²⁷SSSQE⁴³¹ was synthesized and used here (unphosphory-
79 lated, 121 amino acids, 13 kDa, pI = 4.14) (Figure 1E, down).
80 Indeed, four DMP1 peptides are identified in dentin and bone
81 extracts: (1) the full-length protein, (2) a N-ter fragment of 37
82 kDa, (3) a C-ter fragment of 57 kDa,³⁵ and (4) a proteoglycan
83 derived from the N-ter fragment and known as DMP1-PG
84 found in rat and mouse.³⁶ Importantly, five highly conserved
85 motifs during mammalian evolution were identified including
86 ⁴²⁷SSSQE⁴³¹, suggesting the importance of this acidic peptide
87 for the protein structure and/or function.³⁷ Moreover, it was
88 reported that the 57 kDa fragment recapitulates the function of

full-length DMP1 in regulation of mineralization and osteocyte
89 maturation.³⁸ Finally, phosphorylation appears to be dedicated
90 to the organization of mineral when there is no collagen
91 fibrillar packing³⁹ since both phosphorylated and non-
92 phosphorylated rDMP1 are proposed as apatite nucleators.⁴⁰ 93

In addition to the confinement criterion, the concentration
94 of protein closer to physiological conditions^{41,42} is of
95 importance to access to their native properties. Noticeably,
96 differences in range of concentrations (factor 10 to 100) can
97 lead to opposite activities^{14,15,24,26,29} blurring the conclusions. 98

To validate the specificity of the 57-KD fragment of DMP1,
99 experiments were also carried out with a synthetic calcium-
100 binding polymer, that is, the biomimetic poly-L-aspartic acid
101 (polyAsp, 1.2 kDa, pI = 2.98) (Figure 1E, up).^{43,44} Nowadays,
102 polyAsp is commonly used in biomineralization models for
103 mimicking acidic proteins in biological calcified tissues (nacre,
104 bone, dentin). This polymer is described to allow the
105 intrafibrillar infiltration of both DMP1⁴⁵ and apatite ion
106 precursors⁴⁶ and consequently the collagen/mineral co-align-
107 ment.⁴⁷ 108

Here, we show that collagen suprafibrillar confinement
109 drives the activity of acidic calcium-binding polymers on apatite
110 mineralization. The difference in mineralization between 111

rpDMP1 and polyAsp illustrates the specificity of NCPs amino acids sequence. Under confinement (i.e., intra- and inter-brillar spaces), we show that all the identified functions of rpDMP1 are dedicated for proper calcification to occur, namely the collagen/apatite co-alignment.^{48,49} Indeed, we show that while collagen nucleates apatite, rpDMP1 (i) concentrates the apatite ion precursors locally, interfering as a supporting agent for collagen (ii) to induce the formation of a first highly crystalline apatite crystal. In addition, we discard that rpDMP1 stabilizes the possible transient precursors of bioapatite^{50,51} since they only form at high nonphysiological concentrations of protein. Finally, (iii) we confirm that it inhibits the apatite growth, but (iv) also show that it might inhibit the homogeneous nucleation irrespective of confinement. Overall, the work demonstrates that aside from cellular and biochemical processes, physicochemical parameters take part in the control of bone biomineralization.

MATERIALS AND METHODS

Synthesis and Purification of the Recombinant DMP1 Peptide. Cloning. The coding sequence of our targeted human DMP1 peptide (i.e., amino acids 367–481) was isolated from a human primary osteoblast cDNA (Promocell). It was amplified in a thermal cycler (Mastercycler pro, Eppendorf) by PCR using the oligonucleotides 5' ATGC CATATG AACCCCGACCCCA 3' and 5' GCAT CTCGAGTCA GTG GTG GTG GTG GTG GTG GTG CAACTGGCCATCTTC 3' to create NdeI and XhoI restriction sites at the start and stop codon, respectively. Cycling conditions were initial denaturation at 94 °C for 2 min, followed by 35 cycles, each cycle consisting of 30 s of denaturation at 94 °C, 45 s of annealing at 60 °C and 45 s of elongation at 72 °C. The final elongation lasted for 2 min at 72 °C. PCR products were analyzed by 1% agarose gel electrophoresis and observed in an analyzer Gel Doc (BIORAD, France) after ethidium bromide staining. The amplified fragment was purified using the QIAquick PCR purification kit (Qiagen SA) and digested with NdeI and XhoI restriction enzymes. The resulting products were separated by a migration in a 1.5% agarose gel with ethidium bromide and cloned into the pET22b vector (Novagen/VWR International S.A.S) which had been beforehand digested similarly. Competent Novablue cells were transformed with the ligation mix and selected by overnight growth on LB agarose plates containing 70 µg/mL ampicillin. Positive clones were checked for the presence of the 0.8 kb fragment by NdeI and XhoI restriction analysis.

Expression. BL21 (DE3) cells transformed with pET-DMP1 were grown on a rotating table (220 rpm) overnight at 37 °C in LB medium with 50 µg/mL ampicillin. When the optical density at 600 nm of the bacterial broth reached 0.6–0.8, the induction was realized with an addition of 1 mM of isopropyl β-D-1-thiogalactopyranoside. After 3 h, the cells were centrifuged at 5000g for 10 min, and the pellets were frozen at –20 °C until purification.

Purification. Pellets were resuspended in 5 mL of buffer A (PBS 1×, 50 mM imidazole, complete inhibitor mix (Roche), lysozyme, DNase) and lysed by sonication three times. The bacterial extract was loaded on a 5 mL bed-volume HisTrap column using an Akta Purifier 10 (GE-Healthcare). Buffer A was flowed through the column at 1 mL/min until A₂₈₀ of the flow through reached a stable value. A 60 mL linear gradient of 0–100% Buffer B (PBS 1×, 500 mM imidazole, complete inhibitor mix (Roche), lysozyme, DNase) in buffer A was applied, and 1 mL fractions were collected. Aliquots were analyzed by 10% SDS-PAGE and the gels stained with Coomassie blue to determine the quality of purified rpDMP1 (12 kDa, pI = 4.14) (Figure S6).

Sample Preparation. Collagen Extraction. A solution of type I collagen at ~3 mg/mL in 0.5 M acetic acid was prepared as previously described.⁵² The collagen was extracted from rat tail tendons. After a washing step with PBS, tendons were solubilized in 0.5 M acetic acid, and the solution was clarified by centrifugation (21,000 rpm, 2 h, 11 °C). The supernatant was selectively precipitated with 0.3 and 0.6 M

of NaCl by two centrifugations (21,000 rpm, 3 h, 11 °C then 4400 rpm, 45 min, 11 °C), in order to remove proteins other than type I collagen and collagen, respectively. The pellets were solubilized in 0.5 M acetic acid and dialyzed against 0.1 M acetic acid in order to remove salts from the solution. A final centrifugation was performed (21,000 rpm, 4 h, 11 °C), and the concentration was adjusted to a final stock concentration of ~3 mg/mL. The final concentration of type I collagen solution was estimated by hydroxyproline titration.⁵³

Synthesis of Collagen/Apatite Matrices. The matrices were prepared according to a procedure that combines injection and reverse dialysis processes (patent WO2011151587A2). The mineralization conditions for collagen matrices (concentrations of components) are summarized in Table 1. Collagen samples were disk-shaped

Table 1. Mineralization Conditions for Collagen Matrices^a

sample designation	end collagen concentration (mg/mL)	acidic polymer (µg/mL)	HA ion precursors in dialysis solution (acetic acid)	additional SBF bath
Coll	250	–	–	–
Coll/SBF	250	–	SBF	–
Coll/SBF/SBF	250	–	SBF	+
Coll/CHA/SBF	250	–	CHA	+
Coll/SBF(rpDMP1)	250	2.5	SBF	–
Coll/SBF(rpDMP1)/SBF	250	2.5	SBF	+
Coll/10rpDMP1	250	25	–	–
Coll/SBF(10rpDMP1)	250	25	SBF	–
Coll/SBF(10rpDMP1)/SBF	250	25	SBF	+
Coll/SBF(polyAsp)	250	5.75	SBF	–
Coll/SBF(polyAsp)/SBF	250	5.75	SBF	+
CollOsteoid/SBF/SBF	40	–	SBF	+
CollOsteoid/SBF(10rpDMP1)/SBF	40	25	SBF	+

^aIonic composition of SBF and CHA solutions are detailed in Table S1.

with a thickness of ~1 mm and a diameter of ~10 mm. A soluble acidic collagen solution (1 mg/mL, 0.5 M acetic acid) supplemented by a 1× SBF solution mimicking the ionic compounds found in human plasma was prepared by diluting the stock solution (3 mg/mL, 0.5 M acetic acid) with a 1.5× SBF acidic solution (0.5 M acetic acid). SBF was prepared as previously described.⁵⁴ The concentrations of the salts precursors are summarized in Table S1. Two concentrations of rpDMP1 were added to this solution: 2.5 µg/mL (low) and 25 µg/mL (high) to form the matrices referred as Coll/SBF-rpDMP1 and Coll/SBF-10rpDMP1, respectively. The rpDMP1 control matrix with the highest concentration of rpDMP1 and without mineral (Coll/10rpDMP1) was obtained by diluting the stock solution with acetic acid (0.5 M). Two matrices used as control were prepared: (i) without any organo-mineral additives (Coll) and (ii) with apatite ion precursors but without addition of any acidic polymer (Coll/SBF). The matrices supplemented by polyaspartate (Coll/SBF-polyAsp) were obtained in a similar way, by adding this acidic polymer (5.75 µg/mL, Lanxess-Bayer, BaypureDS100, 1200 g/mol, pI = 2.98) to the acidic collagen solution. To mimic the osteoid tissue⁵⁵ CollOsteoid/SBF/SBF, 40 mg/mL collagen matrices were also formed. In this case, 25 µg/mL rpDMP1 was added to the acidic collagen solution forming the matrix referred as CollOsteoid/SBF-10rpDMP1/SBF. All these solutions were continually injected in a closed dialysis chamber for 1 week. The bottom of the chamber contained a dialysis membrane with a molecular weight cut off of 1 kDa. The reverse dialysis process⁵⁶ was set against polyethylene glycol (PEG, 35 kDa, Fluka) to

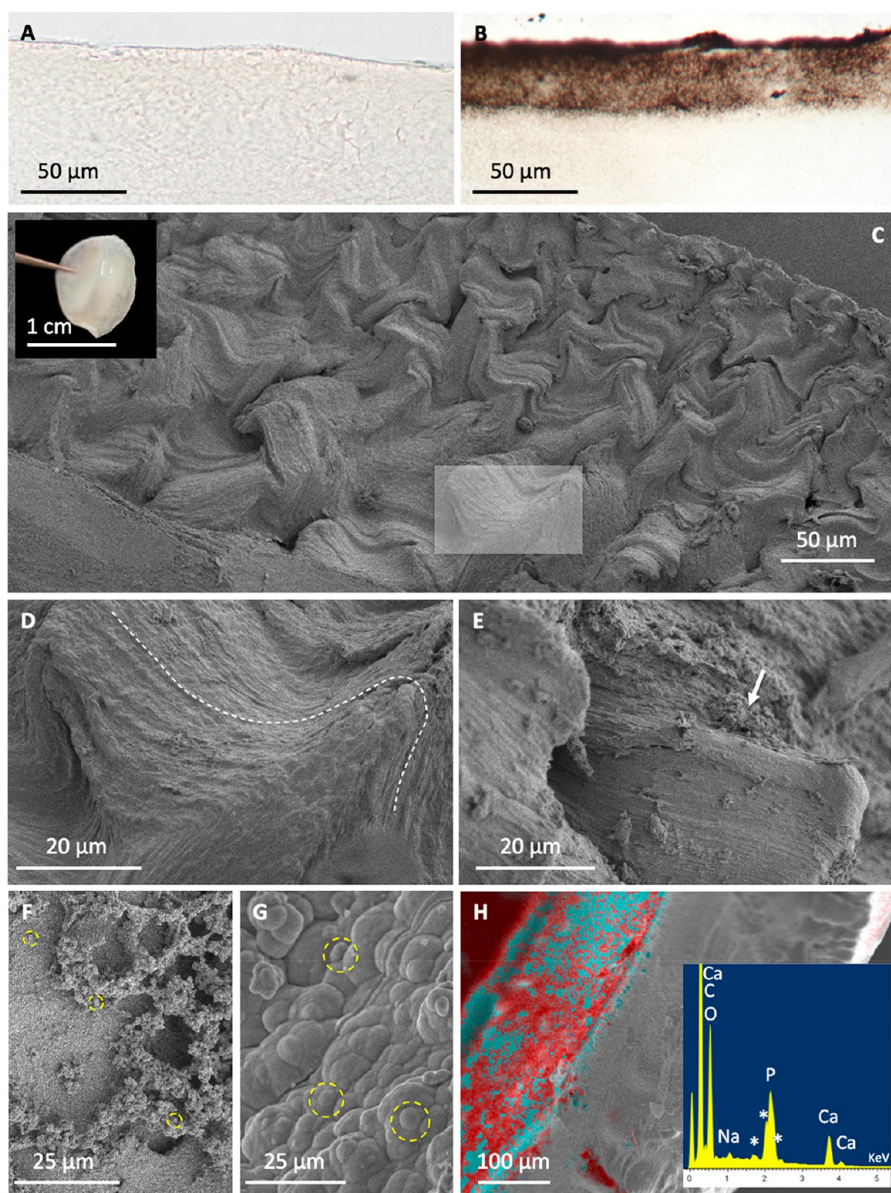


Figure 2. Structural characterizations of the hybrid collagen/apatite matrices in the presence of rpDMP1 containing the highly conserved motif. Histological sections of Coll/SBF(1rpDMP1) (A) and Coll/SBF(1rpDMP1)/SBF (B) stained with von Kossa. SEM observations of the dense collagen matrix (250 mg/mL) at low rpDMP1 content (C–F). Examination of the fractured interior of the disk-shaped matrix (C, inset) before SBF bath (Coll/SBF(1rpDMP1)) (C, D; rectangle in C indicates the enlarged section that is shown in D). Examination after SBF bath (Coll/SBF(1rpDMP1)/SBF): interior (E) and surface (F). In (E), the roughness appears to increase after SBF (arrow). (G) Surface of the collagen matrix precipitated without rpDMP1 (Coll/SBF/SBF). The size and the amount of spherulitic crystals (yellow dashed circles) at the surface of the matrices decrease with rpDMP1. (H) Analysis of Coll/SBF(1rpDMP1)/SBF by EDX coupled with SEM; calcium ions are in red, and phosphorus ions in blue (SEM gold coating*).

control the final concentration of collagen. The PEG was dissolved in 0.5 M acetic acid up to ~300 mg/mL for all Coll matrices or ~50 mg/mL for all CollOsteoid matrices. To form the mineralized matrices, the ionic precursors of SBF or CHA were dissolved in the PEG/acetic acid solution. After injection of the total amount of collagen, dialysis was continued for 1 week in order to obtain a homogeneous concentration in the samples and a good maturation of the peptides. The pH was then increased to a range of 9–10 by ammonia gas diffusion for 3 days to induce collagen fibrillogenesis, stabilize the liquid crystalline organization into dense fibrillar matrices, and precipitate the mineral phase. These matrices were then removed from the dialysis chamber and washed several times in sterile double-distilled water until complete neutralization. The final concentration of type I collagen in the different collagen matrices was assessed by hydroxyproline titration and found to be ~250 mg/mL

for all Coll matrices and ~40 mg/mL for all CollOsteoid matrices. The different matrices were split into halves, and the mineralization degree of one-half was increased by a bath in 45 mL of 1.5× SBF solution at pH = 7.4 under mild rotary stirring (220 rpm) at 37 °C for 1 week.⁸

Mineral Characterization. Histology. Bone samples were fixed in 4% paraformaldehyde and embedded in paraffin for sectioning. Ten μ m-thick serial sections perpendicular to the cell layer were dewaxed, rehydrated, and stained specifically by von Kossa, thus identifying divalent ions. The slides were rinsed, dehydrated, and mounted for observation with an optical microscope (Nikon E600 POL) or an epifluorescence microscope (AXIO 100 Zeiss).

Scanning Electron Microscopy and Energy-Dispersive X-ray Spectroscopy. Each sample was fixed in 3.6% glutaraldehyde in a cacodylate/saccharose buffer solution (0.05 M/0.6 M -pH 7.4). The

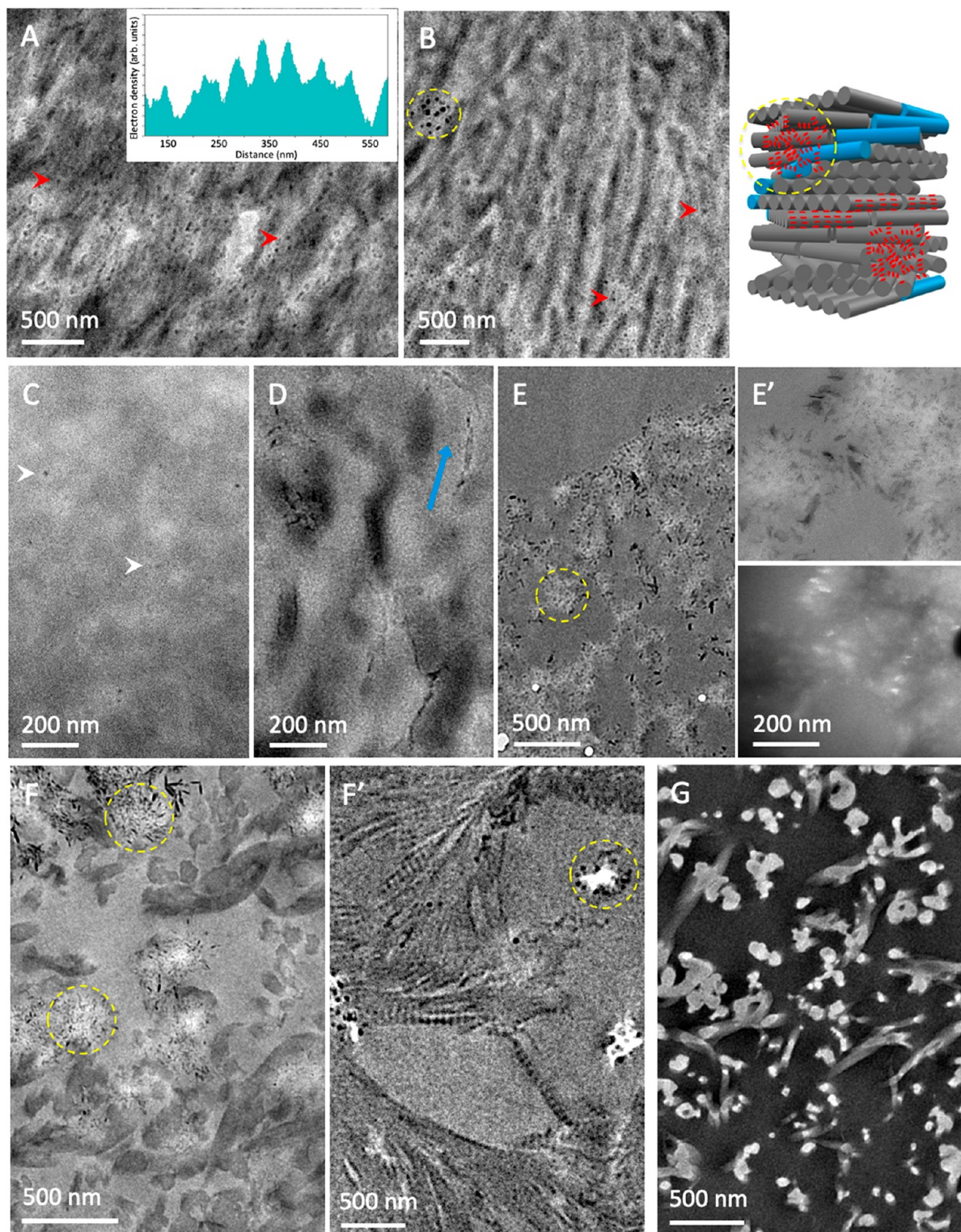


Figure 3. Investigations by TEM of calcium phosphate precipitation in rpDMP1 containing a mature bone-like matrix. Observations of unstained ultrathin sections of (A) Coll/SBF(1rpDMP1) and (B) Coll/SBF(10rpDMP1) before the SBF bath. Both matrices present nanometric electron-dense precipitates (red arrows) along and/or within the collagen fibrils with periodicity highlighted by the intensity profile. In addition, aggregates of the “dot-like” precipitates are seen in Coll/SBF(10rpDMP1); the scheme illustrates the corresponding distribution of dot-like precipitates (red) related to the organization of collagen fibrils. They are found along the fibrils or as aggregates (yellow dashed circle). (C) A control matrix prepared with the high concentration of rpDMP1 (25 $\mu\text{g}/\text{mL}$) and without ionic apatite precursors (Coll/10rpDMP1) shows that collagen fibrils are hardly observed in contrast to the precipitated proteins (yellow arrows). After immersion in SBF, (D) axial alignment of platelets are observed inside the matrix. (E) Observations within the most superficial layer of the matrix show the presence of spherulitic crystals (E') according to bright-/dark-field images. Observations at the surface of Coll/SBF(10rpDMP1)/SBF (F) without and (F') with staining where the dashed yellow circles highlight the presence of spherulite. Observations at the surface of (G) Coll without staining.

248 samples were dehydrated through successive ethanol bath (50%, 70%,
249 80%, 90%, and 100%), and a supercritical CO₂ drying process was
250 performed on a BAL-TEC 030. Then samples were sputter-coated
251 with a gold layer of 10 nm and observed in a Hitachi S-3400N
252 operating at 3 kV.

253 Energy-dispersive X-ray (EDX) microanalysis was used for the
254 mapping of mineral deposits inside the matrices. The EDX instrument
255 X-Max (Oxford Instruments) was coupled to a scanning electron
256 microscope Hitachi S-3400N operating at 12 kV, and the Oxford
257 Microanalysis Group XAN.70 software was used for this analysis.

258 **Transmission electron microscopy.** This protocol is similar to the
259 protocol for the scanning electron microscopy (SEM). Then, samples
260 were rinsed, dehydrated, and embedded in Epon 812. For a few
261 samples (always mentioned in the text), an additional postfixation was
262 carried in 2% osmium tetroxide in a cacodylate/saccharose buffer
263 solution (0.4 M/0.6 M-pH 7.4) during 1 h at 4 °C, otherwise neither
264 osmium nor uranyl acetate (staining) were added to avoid artifacts on
265 ultrathin sections. Sections (~80 nm) were observed with a Tecnai
266 spirit G2 operating at 120 kV.

267 **Wide-Angle X-ray Diffraction (Transmission Mode).** Matrices
268 were inserted in X-ray cylindrical borosilicate capillary tubes. The
269 tubes were wax-sealed to keep the samples hydrated and placed
270 directly in the vacuum chamber beam. X-ray diffraction experiments
271 were performed with a S-MAX 3000 RIGAKU using a mono-
272 chromatic CuK α radiation. The diameter of the cylindrical beam
273 dimension of the specimen was 400 μ m. The data were collected in
274 the 3–60° range (2 θ). The sample-to-detection distance was 0.059 m
275 with a diameter of capillary tubes at 1 mm or 0.058 m with a diameter
276 of 2 mm. The two-dimensional (2D) wide-angle X-ray diffraction
277 (WAXD) patterns were collected with imaging plates then scanned.
278 The data were analyzed using Image (LPS, U-psud) software.

279 **ζ -Potential Measurements.** The matrices were washed before
280 characterization precluding the involvement of free ions in the
281 resulting global charge. Samples were crushed in liquid nitrogen, and
282 the resulting powders were dispersed in PBS solution. Measurements
283 were carried out using a Malvern Zetasizer Nano ZS90.

284 **Thermogravimetric Analysis.** Samples were dried under a laminar
285 hood overnight to minimize the mass loss of loosely bounded water.
286 The analysis was performed on a thermo-microbalance instrument
287 (NETZSCH STA 409PC). The measurement was performed from
288 room temperature to 1000 °C in an oxidizing atmosphere (air) with a
289 heating rate of 5 °C/min.

290 **Nuclear Magnetic Resonance.** Solid-state nuclear magnetic
291 resonance (ssNMR) experiments were realized on hydrated samples.
292 Magic angle spinning (MAS) spectra were acquired at a frequency of
293 8 kHz, with samples packed into 4 mm zirconia rotors. ¹H–³¹P cross-
294 polarization (CP) experiments were performed on an Avance 300
295 Bruker spectrometer operating at frequencies of 300.13 MHz (¹H)
296 and 121.50 MHz (³¹P). The contact times (CT) were set at 10 and 1
297 ms, and the recycle delay (RD) at 2 s. A 2D ¹H–³¹P heteronuclear
298 correlation (HETCOR) was performed with the following param-
299 eters: RD = 2 s, CT = 10 and 1 ms, 1280 transients for each 128 t1
300 increments. ¹H and ³¹P chemical shifts were referenced (δ = 0 ppm)
301 to adamantane and to 85%w aqueous H₃PO₄, respectively.

302 ■ RESULTS AND DISCUSSION

303 **Structural Characterizations of rpDMP1-Loaded Min-**
304 **eralized Collagen Matrices.** Experimental conditions and
305 matrices composition (see [Materials and Methods](#)) are
306 summarized in [Table 1](#). It includes collagen matrices without
307 additives used as control, namely Coll, Coll/SBF ([Figure S1](#)),
308 Coll/CHA/SBF ([Figure 1D](#), left), and Coll/SBF/SBF ([Figure](#)
309 [1D](#), right).⁸ This means that after coprecipitation of collagen
310 with either CHA (~20 times more concentrated in apatite ion
311 precursors see [Table S1](#)) or SBF, these two last matrices were
312 then immersed in SBF to mimic further steps of mineral
313 growth. With additives (rpDMP1, polyAsp), fibril precipitation
314 was only performed in SBF ([Ca²⁺] = [Ca²⁺]ECF)⁵⁷ to set

physiological-like conditions. Two concentrations of rpDMP1 315
were used here, for which we will refer to low (2.5 μ g/mL, 316
1rpDMP1) and high (25 μ g/mL, 10rpDMP1). For a better 317
reading, data for 10rpDMP1 are displayed in the main text 318
only when differences in mineral formation are observed 319
between the two concentrations. Additional data for 320
10rpDMP1 are shown in [Figure S2](#). 321

We first investigated the spreading of mineral within 322
collagen matrices before and after immersion in SBF for 323
both rpDMP1 concentrations. For this purpose, we used von 324
Kossa staining on histological sections as it allows observations 325
at large scale. Interestingly, staining is hardly observed without 326
SBF extra-bath (Coll/SBF(1rpDMP1), [Figure 2A](#)), whereas a 327
strong staining is observed for both matrices at the superficial 328
layer (~50 μ m thick) after immersion (Coll/SBF(1rpDMP1)/ 329
SBF, [Figure 2B](#)). Because von Kossa staining may interfere 330
with any divalent ions,⁵⁸ further characterizations were 331
performed to confirm the presence of mineral. 332

To identify the distribution of the minerals at a lower scale, 333
investigations were performed at the surface and inside 334
fractured disk-shaped matrices ([Figure 2C–G](#)) using SEM 335
coupled with EDX microanalysis ([Figure 2H](#)). Observations 336
inside the matrix reveal that the entrapment of rpDMP1 does 337
not disturb the self-assembly of collagen fibrils since twisted 338
plywood structures form over large distances ([Figure 2C,D](#) 339
versus [Figure S1](#), Coll/SBF). After immersion in SBF (Coll/ 340
SBF(1rpDMP1)/SBF), the surface roughness is more 341
pronounced by the deposition of particles on the closely 342
packed oriented fibrils ([Figure 2D](#) versus [2E](#), arrow). On the 343
surface of the matrices, spherulites usually consisting of 344
aggregates crystals⁵⁹ are observed, and, interestingly, they 345
appear to be smaller in the presence of the protein (~0.5 μ m 346
versus ~4 μ m in Coll/SBF/SBF, yellow dashed circles in 347
[Figure 2F,G](#)). In addition, they appear less abundant since they 348
do not cover the whole surface. EDX analysis shows that the 349
spherulites are mainly composed of calcium and phosphorus 350
atoms with an average Ca/P ratio of about 1.66–1.71, 351
suggesting the formation of apatite in the presence of 352
rpDMP1 ([Figure 2H](#)).⁶⁰ 353

Confinement Effect on Apatite/rpDMP1 Distribution. 354
TEM investigations were performed on thin sections of Coll/ 355
SBF(1rpDMP1) and Coll/SBF(10rpDMP1) to access higher 356
magnifications. TEM sections were not stained to avoid the 357
presence of staining deposits that are difficult to distinguish 358
from CaP crystals.⁸ Observations evidence the presence of 359
nanometric electron-dense (“dot-like”) precipitates (~30 nm) 360
with low and high rpDMP1 concentrations (red arrows in 361
[Figure 3A,B](#), respectively) within the dense collagen network at 362
this scale. The striated pattern (67 nm) is observed locally 363
([Figure 3A](#), inset), indicating that some minerals localize inside 364
the gap regions.^{8,61} Interestingly, additional aggregates of 365
nanoprecipitates are observed for high rpDMP1 content 366
([Figure 3B](#), yellow dashed circle). 367

To help identify the nature of the “dot-like” precipitates, a 368
control matrix was prepared without apatite ion precursors 369
(Coll/10rpDMP1) ([Figure 3C](#)). Likewise, “dot-like” particles 370
are observed (white arrows), confirming that the nano- 371
precipitates are also composed of the protein, in agreement 372
with previous observations.¹⁴ Although it is difficult to 373
conclude due to the low contrast of organic components in 374
TEM, the precipitates appear less abundant and not distributed 375
along the fibrils ([Figure 3C](#) versus [Figure 3A,B](#)). Nevertheless, 376
spherulites are observed without rpDMP1 in SBF (Coll/SBF/ 377

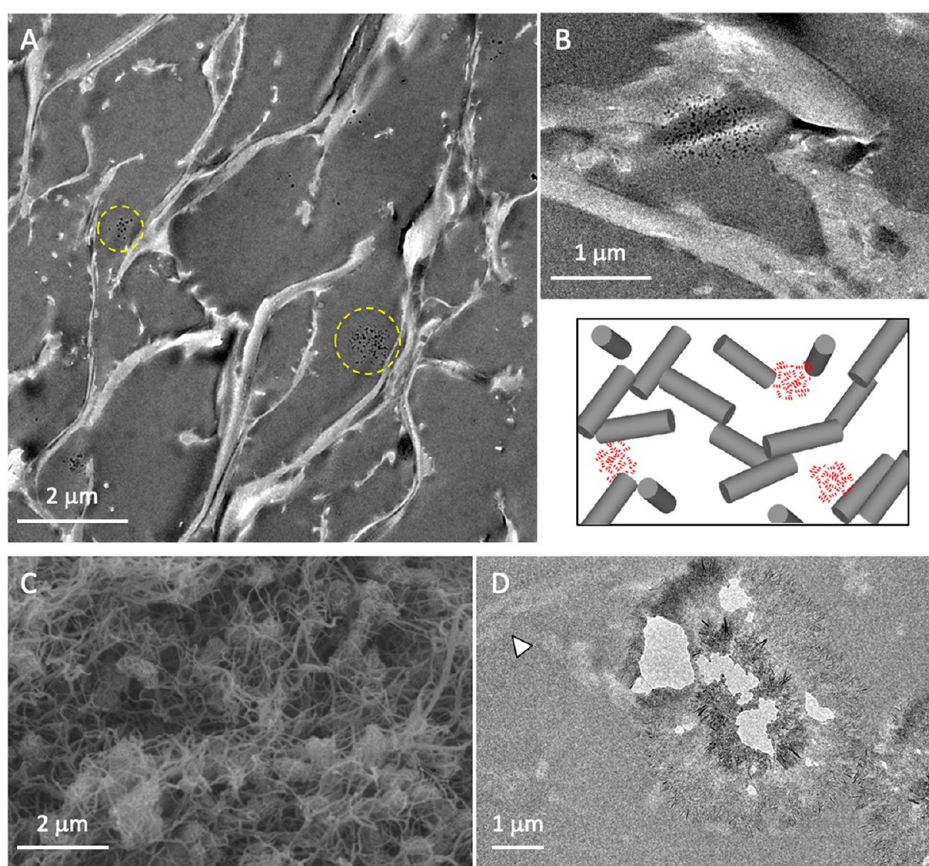


Figure 4. Investigations of calcium phosphate precipitation in rpDMP1 containing osteoid-like matrix. TEM observations of unstained ultrathin sections of CollOsteoid/SBF(10rpDMP1)/SBF (A) at low magnification where “dot-like” precipitates aggregate locally (yellow dashed circle). They are clearly seen (B) at higher magnification. Below, the corresponding relationship between mineral and collagen is represented in the scheme. (C) SEM and (D) TEM micrographs of CollOsteoid/SBF/SBF. The absence of staining makes the observation of cross-striated fibrils usually seen in a longitudinal cut (white arrow) difficult.

SBF in Figure 1D and Coll/SBF⁸), indicating that rpDMP1 may interfere with some ions to localize specifically inside the gap zone.

After a SBF bath, few platelets with axial alignment (blue arrow in Figure 3D) are observed for both rpDMP1 concentrations together with spherulitic particles at the surface and within the most superficial layer of the matrix (dashed yellow circles in Figure 3E,F). Noticeably, they appear smaller in size (~300 nm versus ~2 μm) than those found without rpDMP1 (Figure 1D, right), but remain crystalline according to the contrast observed in the dark-field TEM image (Figure 3E'). Further observations at the surface of Coll/SBF(10rpDMP1)/SBF allow the visualization of both the resin and individual fibril due to a lower density of collagen locally. A comparison between mineralized collagen fibrils without or with staining (Figure 3F,F', respectively) confirms the precipitation of fibrils with the cross-striated pattern. Observations of an unmineralized collagen matrix without staining (Coll, Figure 3H), where the resin is darker than collagen, confirm that the contrast in mineralized samples comes from the spreading of mineral (ions or precipitates) over the collagen fibrils. Note that we cannot conclude on the involvement of DMP1 in mediating the size of fibrils,²⁹ since the average diameter of collagen fibrils appears unmodified (~100 nm) regardless the rpDMP1 concentration.

Going further, thermogravimetric analysis (TGA) investigations were performed to better characterize the effect of

rpDMP1 concentration on apatite formation (Figure S3). Considering the standard deviation recorded for the matrices (± 5 °C), the difference in mineralization degree should be considered as a trend here. After SBF bath, the mineral content increases for low rpDMP1 concentrations (from ~3.5% to ~18%), whereas it does not change significantly at a high content of rpDMP1 (from ~12.5% to ~15%). Since both matrices exhibit spherulites on the surface after SBF bath (Figure 2F and Figure S2C), it indicates that the main mineral content is localized inside the matrix in the form of nanoprecipitates.

In addition, the ζ -potential was investigated to help understand the interactions between collagen, rpDMP1, and the mineral. Before the SBF bath, the global surface charge tends to be neutral in the presence of rpDMP1 whatever the concentration of the protein ($-0.79 (\pm 0.66)$ mV and $0.22 (\pm 0.78)$ mV) for low and high rpDMP1 concentrations, respectively). Indeed, the Coll/SBF matrix exhibits a negative ζ -potential ($-4.3 (\pm 2.5)$ mV).⁸ This result agrees with a partial coating of collagen by the selected DMP1 domain, which includes the two collagen binding sites²⁹ and the highly conserved acidic peptide.³⁷ Indeed, it is reported that DMP1 binds to the N-telopeptide region in collagen.^{29,45} After immersion in SBF, Coll/SBF(1rpDMP1)/SBF exhibits a more negative ζ -potential ($-9.77 (\pm 0.48)$ mV), while it is slightly unchanged for Coll/SBF(10rpDMP1)/SBF ($-1.79 (\pm 0.33)$ mV). The ζ -potential of HA being negative between pH 5 and

8 (−5 to −37 mV respectively)⁶² is in agreement with the TGA analysis.

To go deeper in the understanding of how the suprafibrillar confinement affects the spreading of rpDMP1 over collagen, the peptide was entrapped in a scaffold mimicking the fibrillar arrangement of osteoid tissue (Figure 4A,B). As mentioned above, this means that (i) the fibrils network is not as dense as the previous mature bone model with larger interfibrillar spaces which are above the micron size (macroporous gel) and (ii) the collagen fibrils are not organized (no plywood geometry). Note that spherulitic crystals form in the control matrix without peptide (Coll/Osteoid/SBF/SBF) as shown by SEM (Figure 4C) and TEM (Figure 4D), which is in agreement with previous observations on a matrix with a collagen gradient concentration.⁸

Interestingly, “dot-like” precipitates are also observed, but they do not localize inside the fibrils (Figure 4A). Although most of the investigations have focused on electrostatic interactions to explain the intrafibrillar infiltration of both mineral and proteins in collagen, this difference in apatite crystals distribution in osteoid- and mature bone-like matrices evidence that the suprafibrillar confinement provided by the collagen assembly in bone tissue (i.e., cholesteric geometry) plays a key role on this phenomenon. Thus, it may explain the need of other proteins to infiltrate collagen in models lacking biomimetic interfibrillar spaces (<1.7 nm).⁴⁵ The fact that collagen here is continuously in contact with ions (even before mineralization) as occurring in bone strengthens the need of a balance between osmotic equilibrium and electroneutrality for intrafibrillar mineralization.⁶³ In addition, the use of biomimetic collagen interfibrillar spaces (in both osteoid- and mature-like matrices) contradicts that rpDMP1 favors the templating of crystal growth.¹⁴ Conversely, according to (i) TEM and SEM investigations where it is observed that the size of spherulites consist of either “dot-like” aggregate or “mature” apatite platelets with (Figures 2F, 3B, and 4A,B) or without rpDMP1 (Figures 2G and 4C,D) respectively, (ii) the mineral amount found by TGA, and (iii) the resulting surface charge probed by ζ -potential, we confirm its role as a growth inhibitor.³³ This effect may be related to the confinement which promotes the protein folding by destabilizing the unfolded state^{64,65} and thus further specific interactions between rpDMP1 and apatite nuclei; the structural characteristics of growing apatite being driven by the involvement of ionic substitutions (specifically from carbonate ions).⁶⁶

Toward Evidence of the Specific Activity of rpDMP1 under Confinement. When polyAsp is supplemented to collagen, it is difficult to observe a difference in mineral spreading over the collagen matrix before immersion in SBF (Coll/SBF(polyAsp), Figure 5A) as compared to rpDMP1. Here also, a lower concentration of polyAsp (5.75 $\mu\text{g/mL}$) was used than that in the literature (usually between 10 and 100 $\mu\text{g/mL}$) (i) to reach at least 2 magnitude order lower amounts of negative charge and (ii) to reproduce a more “realistic” collagen to NCPs ratio since such a high polyAsp concentration can be used for one single fibril.³ Note that our samples were not lyophilized before characterization, as it is commonly performed with this polymer in the literature.⁶⁷ This is of importance since the resulting phase and degree of mineralization are here related to a hydrated biomimetic process. The difference between polyAsp and rpDMP1 in apatite mineralization becomes clearer after the SBF bath (Coll/SBF(polyAsp)/SBF, Figure 5B,C). TEM observations

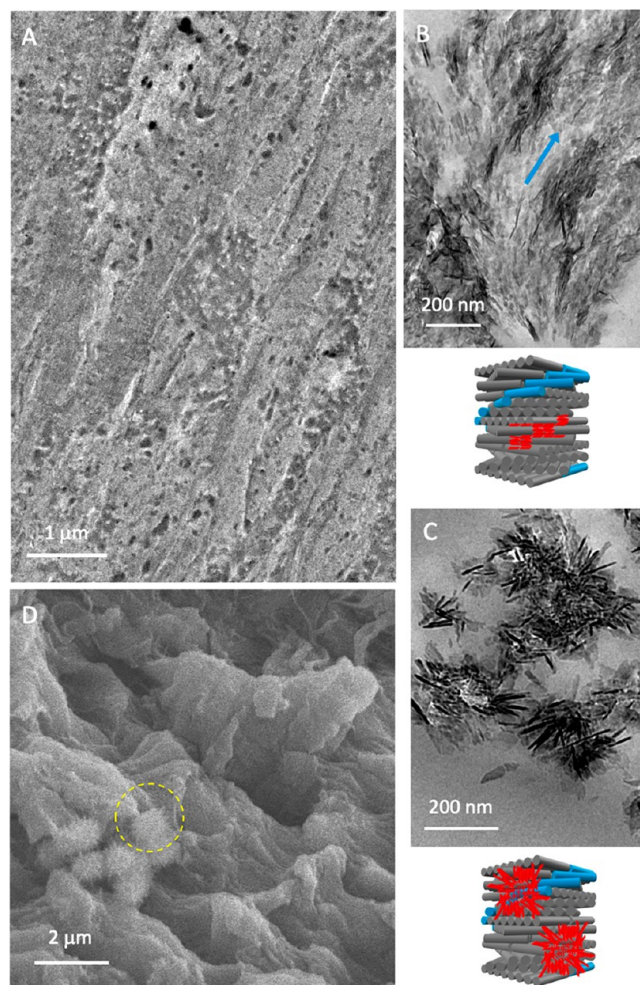


Figure 5. Calcium phosphate precipitation investigated by EM in the presence of polyAsp containing mature bone-like matrix. Observations by TEM of unstained ultrathin sections of matrices before and after SBF immersion: (A) Coll/SBF(polyAsp) and (B,C) Coll/SBF(polyAsp)/SBF, respectively. The corresponding schematic representation of the apatite crystal (red)/collagen fibril (gray) relationship in a 3D perspective is shown. The lack of staining in (A) indicates that there is an homogeneous spreading of SBF ions over the collagen. After immersion in SBF (B, C), HA platelets are observed in the matrix; they either (B) co-align with the long axis of the fibril (blue arrow) or (C) form spherulites as schematically presented, respectively, below the (B) and (C) TEM micrographs. (D) SEM micrograph of Coll/SBF(polyAsp)/SBF showing that spherulites (yellow dashed circle) are also observed inside the matrix.

reveal large domains where the mineral platelets co-align with the collagen fibrils (Figure 5B), as seen in mature bone (Figure 1C) strengthening the affinity of polyAsp for gap regions in line with previous conclusions in the literature.^{44,45,68} However, in contrast to rpDMP1 results, nonbiomimetic spherulitic crystals are also observed in the matrix (Figure 5C,D), strengthening that only small molecules such as osteocalcin can penetrate alone the intrafibrillar space.⁶⁹

These observations evidence the specificity of biological rpDMP1 versus synthetic polyAsp amino-acids sequences in mediating apatite mineralization and also suggest that, in addition to its role as growth inhibitor, it may prevent homogeneous nucleation.³³ Finally, because the rpDMP1 sequence is not phosphorylated, this result tends to strengthen

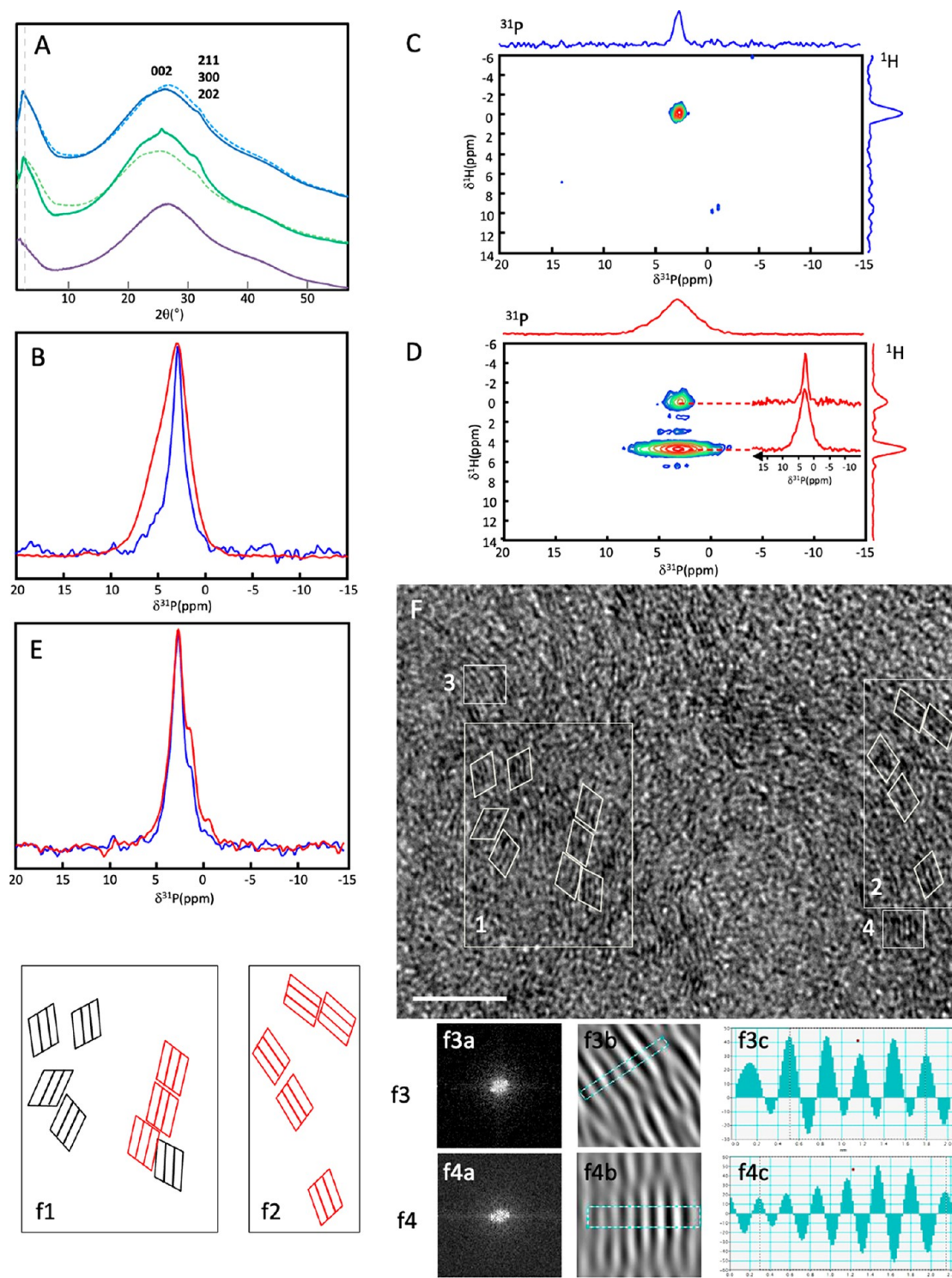


Figure 6. Characterization of the mineral phase. (A) 1D radial average of the WAXD patterns of matrices at 2.5 $\mu\text{g/mL}$, without (Coll/SBF(1rpDMP1), dashed green) and with (Coll/SBF(1rpDMP1)/SBF, plain green) immersion in SBF; at 25 $\mu\text{g/mL}$, without (Coll/SBF(10rpDMP1), dashed blue) and with (Coll/SBF(10rpDMP1)/SBF, full blue) immersion in SBF, and a collagen matrix as control (purple). The vertical gray dashed line points out the shift of the reflection which corresponds to the lateral distance between collagen molecules. (B) 1D ^{31}P CP MAS spectra (CT = 10 ms) of the hybrid matrices Coll/SBF(1rpDMP1) (blue) and Coll/SBF(1rpDMP1)/SBF (red). (C) 2D ^1H - ^{31}P HETCOR spectrum (CT = 10 ms) of Coll/SBF(1rpDMP1) with extracted ^{31}P slices at $\delta(^1\text{H}) = 0$ and 4.9 ppm. (D) 2D ^1H - ^{31}P HETCOR spectrum (CT = 10 ms) of Coll/SBF(1rpDMP1)/SBF. (E) 1D ^{31}P CP MAS spectra (CT = 10 ms) of the hybrid matrices Coll/SBF(10rpDMP1) (blue) and Coll/SBF(10rpDMP1)/SBF (red). (F) TEM micrograph of Coll/SBF(1rpDMP1)/SBF (scale bar = 5 nm) showing crystalline HA nanodomains. (f1) and (f2) are schematic representations of the selected areas 1 and 2 showing the orientation of the HA (210) planes; some crystalline nanodomains appear to align along the same direction (red). (f3) and (f4) FFT (f3a and f4a) and inverse FFT (f3b and f4b) performed on the selected areas 3 and 4, confirming the presence of crystalline nanodomains. The d spacings are measured using the inverse FFT profile along one direction (f3c and f4c).

that the organic phosphate concern may be attributed to polyadenosine diphosphate ribose.⁷⁰

Structural Features of rpDMP1-Mediated Mineral under Confinement. WAXD studies were performed to better characterize the mineral phase and collagen network. The 1D radial average of the WAXD patterns is shown in Figure 6A. Without the SBF extra bath, the signal from collagen fibrils is strong, revealing a low degree of mineralization (dashed green and blue lines). The characteristic HA diffraction peaks, that is, (002) and the merged (211), (300), (202) reflections, are observed more clearly after the SBF bath (full green and blue lines), especially for Coll/ SBF(1rpDMP1)/SBF, confirming an increase in mineral content. In addition, the lateral distance between collagen molecules decreases from 1.5 to 1.1 nm in the matrices loaded with rpDMP1 (with or without SBF bath) in comparison to the pure collagen matrix (full purple line) as observed by the peak shift toward high angles (vertical gray dashed line). This distance is shorter than that reported for a mineralized dense collagen scaffold without protein,⁷ but agrees with densely packed microfibrils⁷¹ and fibrillar collagen in a dry state.⁷² Since the sample is studied in its native hydrated form, this decrease in the intermolecular distance in addition to the contrast observed in fibrils by TEM confirm that rpDMP1 localizes inside the gap zones. The peptide may colocalize with the mineral as shown for polyelectrolytes.⁶³ Besides confinement, SBF ions are needed for the protein to localize in intrafibrillar spaces, especially calcium ions, considering the model of a periodic assembly of DMP1 into a β -sheet template with high calcium-binding capacity.¹⁴ Then the Ca-binding protein can interact with a significant fraction of the π , that is, possibly covalently bond to the collagen.⁷³ At a critical concentration of Ca^{2+} , the apatite nucleation which strongly depends on the degree of confinement⁷⁴ that occurs inside collagen. However, since apatite forms without organic additives under collagen confinement,⁸ it shows that rpDMP1 increases the local ions concentration but does not initiate the nucleation here.

To investigate the presence of calcium-phosphate minerals described as transient phases in bone formation^{51,75,76} and better understand the local phosphate environment in collagen-containing rpDMP1, ^{31}P ssNMR experiments were performed (Figure 6B–E). The ^{31}P CP MAS spectrum of Coll/SBF(1rpDMP1) displays a single ^{31}P resonance centered at 2.8 ppm, typical of apatite (Figure 6B, blue). Interestingly, the 2D ^1H – ^{31}P HETCOR spectrum of Coll/SBF(10rpDMP1) does not show the correlation resonance characteristic from HPO_4^{2-} (Figure 6C), excluding the presence of a hydrated disordered surface layer.^{66,77} In addition, the line width of the ^{31}P resonance (LW = 1 ppm) is thinner than matrices prepared without rpDMP1 (Coll/SBF, LW = 2 ppm) (Figure S4). Both the thinner resonance and the absence/low amount of HPO_4^{2-} surface species demonstrate that the mineral “dot-like” precipitates of the matrices are highly crystalline and that rpDMP1 drives the higher crystallinity of apatite. This result is in agreement with previous works proposing that DMP1 impacts on the initial stages of apatite formation, providing a structural local order.¹⁴ Interestingly, DMP1 is found more abundant in the boundary between the dentin and enamel where apatite is well crystallized.³⁰

Finally, it is interesting to note that such “dot-like crystals” were found at the very early stages of normal *in vivo* calcification.¹ After the SBF bath, the ^{31}P LW in CP MAS

spectrum increases (Figure 6B, red), reaching the value found for bone apatite (4.1 ppm).⁷⁸ The fact that the 2D ^1H – ^{31}P HETCOR spectrum of Coll/SBF(1rpDMP1)/SBF (CT = 10 ms) (Figure 6D) displays the two typical resonances of bone minerals, that is, $\delta(^{31}\text{P}) = 2.8$ and 3.2 ppm, shows that the apatite spherulites observed at the surface of the matrices by microscopy (Figure 2F) dominate the spectroscopic signature here.

At a higher rpDMP1 content, 1D ^{31}P CP-MAS spectra (Figure 6E) display two shoulders (centered at 1.5 and –1 ppm) in addition to the apatite resonance at 2.8 ppm before and after SBF. These resonances correspond to the HPO_4^{2-} ions from the OCP phase. Noticeably, the presence of a relatively low content of rpDMP1 leads to the physiological single apatitic phase, whereas a high “nonphysiological” concentration of protein stabilizes an additional CaP phase like OCP. Such high concentrations are classically used in the literature, questioning the role of DMP1 in stabilizing the ACP phase.^{33,34}

HRTEM observations performed on ultrathin sections of Coll/SBF(1rpDMP1)/SBF (Figure 6F and Figure S5) confirm the highly crystalline nature of the “dot-like” precipitates. Spaced thin layers that are mostly concentric, that is, onion-like morphology that appears to be composed of crystalline nanodomains (~ 1 nm) (some are depicted in areas 1 and 2), are observed. Schematic representations of two selected areas (1 and 2) show the orientation of possible crystal planes (f1 and f2). Some crystalline nanodomains appear to align along the same direction (in red). Their morphology is difficult to identify; they are presented here as parallelogram. In (f3) and (f4), fast-fourier transform (FFT) (f3a and f4a) and inverse FFT (f3b and f4b) performed on the (3 and 4) selected areas confirm the presence of crystalline nanodomains. The crystal planes were indexed by comparing the measured d spacings using the FFT profile along one direction with calculated values of HA (f3c and f4c). Two lattice spacings were measured as 0.31 and 0.34 nm, corresponding to the known (210) and (002) crystal planes of apatite (Figure 6F and Figure S5, respectively). Among the 20 analyzed planes, 80% correspond to the (210) and the remaining 20% to the (002).

The possible consequence of a crystalline apatite nano-domain versus an ACP phase can prevent the formation of pathological calcifications. Indeed, ACP as the first solid phase of calcium phosphate formed in bone would delay apatite formation, allowing the precipitation of other transient phases. This hypothesis is in agreement with the fact that the mineralization process of bone is described to occur rapidly as soon as collagen molecules self-assemble into collagen fibrils in the extracellular space.⁷⁹ In fact, according to our knowledge, there is no report on bone pathologies related to the presence of one of the transient apatite phases, although it is extensively described in synthetic models lacking interfibrillar confinement *in vitro* as discussed above.

CONCLUSION

The bone ECM is a dynamic (in terms of structure, remodelling, and ECF) and complex environment where the confinement effect occurs that is not only critical for the cells behavior⁸⁰ but also for the proteins and even the solvent (water).⁶⁴ Indeed, confinement provided by suprafibrillar organization of collagen strongly impacts the activity of mineralizing polymers. The results suggest the occurrence of

sequential cooperative effects during the early stages of bone apatite precipitation through the formation of Ca-DMP1 complex, which then binds to collagen allowing the concentration of apatite ion precursors in gap regions. Formation of highly apatite nuclei and inhibition of both homogeneous nucleation and crystal growth are consistent with a common role of DMP1 controlling the physiological (versus pathological) bone formation. Strikingly, these effects are reached with the nonphosphorylated 57-KD amino acid sequence under confinement. This versatile bone-like model will be useful to provide insights into the role of other bone components (e.g., citrate, proteoglycans, or even different sequences related to NCPs kinetic of maturation (i.e., post-translation modification)) during the successive events that orchestrate mineralization.

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.biomac.1c00206>.

Composition of the solutions used to mineralize collagen matrices, SEM and TEM images, EDX spectra, and TGA curves (PDF)

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Notes

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