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Recent advances in the synthesis, functionalization and biomedical applications of hydroxyapatite: a review

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Hydroxyapatite (HA) is a member of the Ca apatite family. It resembles natural bone in both structure and chemical composition. HA, owing to its bioactive and biocompatible properties, has been commonly used as an implant material in bone tissue regeneration (osteogenesis), and as a drug carrier in drug and gene delivery systems. With the advances in research on the use of HA, an increasing number of researchers are exploring new synthesis processes, characterization and functionalization techniques for HA and its potential role in various fields such as magnetic resonance, controlled delivery of therapeutic drugs, cell separation, bio imaging and treatment of hyperthermia. Therefore, in this review, we highlight the composition of HA, the advances in its synthesis processes, characterization and functionalization techniques, and its importance in the biomedical field in general, and in emerging areas such as implants, drug delivery, composites, coatings, and ceramic materials in particular. The idea behind writing this review was to collect and summarize the most recent studies involving HA, so that researchers can easily find HA-related information compiled in a single document. In addition, we have also discussed the future prospects of HA. We believe that readers will not only conveniently obtain the desired information from this review, but will also get to the core of the information more easily.

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1. Introduction

Knowledge regarding the structure and composition of natural bone from the macro to the nanoscale has been acquired in a stepwise manner.¹ Ancient civilizations lacked sufficient knowledge about the nature of bone and used bone as a fine tool. Concrete information about natural bone became clearer



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with the emergence of microscopic techniques.² With the advances in microscopy, we can now study bone at the micro-scale level. Research is currently being conducted to study bone at the nanoscale level, and more specifically, at the basic unit or building block level (which is called collagen fibril). One of the difficulties in studying bone at the nanoscale level is the extremely small size of the bone crystals, which can be as smaller as a few nanometers in width and thickness.³ Therefore, bone crystals are the smallest crystals formed biologically. Bone has a hierarchically ordered structure, which is similar to that of

other biological structures such as the intestine, muscles, etc.^{3,4} The lowest stage of this hierarchy is the uniform, and highly ordered arrangement of HA nanocrystals in collagen fibrils (Fig. 1). Depending on the nature and shape of the bone; nanocrystals are arranged in a highly ordered manner in the collagen fibril, either in a lamellar or in a layer-by-layer form. The HA nanocrystals are arranged in such a manner that the *c*-axes of the HA nanocrystals are oriented along the fibril axes (Fig. 1).⁴

Chemically, HA is a naturally occurring mineral form of calcium (Ca) apatite (apatite: group of phosphate minerals). Hence, calcium phosphates and its derivatives are of great interest to various fields of science such as biology, medicine, chemistry, and geology. The attempts to determine their chemical structure and composition started in the middle of the 18th century. However, it was Hausen in the 19th century who proposed the existence of different calcium phosphate phases (earlier all calcium phosphates were called apatites). Among the various calcium phosphates, HA is extremely important owing to its structural and chemical similarities to bone phosphates. In addition, HA is the most stable derivative of calcium phosphate salts within the pH range of 4–14 at room temperature.⁷ It is owing to these properties that HA has found numerous applications in the fields of protein chromatography, fertilizers, pharmaceuticals, water treatment, and biomedicine.^{7,8}



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textiles, synthesis of nanoparticles and their role in drug delivery, hydrogels packs for skin care, polymer composite.



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of scaffolds for bone tissue engineering. Although few attempts have been made by harvesting cells from donor/patient to vascularize the scaffolds before their implantation in the patient but area still need intensive research to develop scaffolds with proper vascularization. The worldwide research has achieved success in tissue engineering but matrix induced autologous chondrocyte implantation has received little success in repair of cartilage. The bioactivity of scaffolds totally depends on surface topology, microstructure, chemistry and mechanical properties of scaffolds. Thus controlling the cells behavior and remodelling is a critical step in development of next generation scaffolds for tissue engineering. The functionalized scaffolds loaded with certain amount of growth factor and therapeutics to stimulate cells proliferation and treatment of infection after surgical operations also strongly needed to achieve success in the area of tissue engineering and regeneration. The increasing interest to incorporate a drug delivery function in the scaffolds is emerged as a new field of scaffolds engineering. The drug loaded scaffolds are able to provide sufficient amount of drug at a specific place in comparison to systematic drug delivery systems. The scaffolds loaded with growth factors such as cytokines, hormones, morphogens and proteins need to be developed to enhance cells adhesion, proliferation and differentiation in tissue engineering. The mechanical behavior of scaffolds has important implication on tissue regeneration; stiff scaffolds mimic pre calcified bone and elastic scaffolds mimic the elasticity of muscles, which cause mesenchymal stem cell differentiation down to a myogenic pathway. Thus scaffolds with optimized mechanical strength need to be fabricated for enhanced bone tissue engineering. The critical review of the literature reported during last one decade has clearly indicated that there is lot of scope to improve the properties and functionality of the scaffolds for bone tissue engineering using various biomaterials and their fabrication techniques.



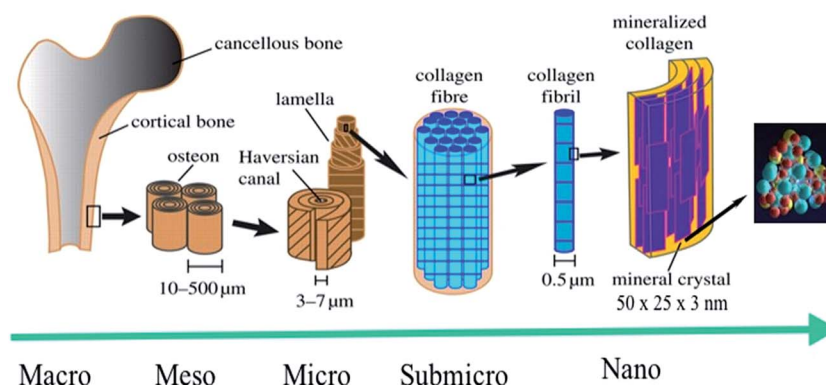


Fig. 1 Hierarchical structure of bone at various size scales. The microstructure of cortical bone consists of osteons with Haversian canals and lamellae, whereas at the nanoscale, the structural units are collagen fibers composed of bundles of mineralized collagen fibrils (this figure was modified from the work of Zhang *et al.* and Rho *et al.*).^{5,6}

HA has found broad applications in the biomedical field⁷ owing to its bioactive and biocompatible nature. It has been frequently used in bone tissue regeneration. Several researchers have reported enhanced osteogenesis using HA.⁷ In addition to enhancing osteogenesis and acting as an implanting material, HA has been commonly used as a coating material as well. HA is usually coated on the surface of the implanting materials to enhance their bioactivity. Furthermore, HA is also a suitable material for carrying drugs (specially proteins) to the targeted sites, or for a slow and sustained release of drugs, which will ultimately help in stimulating the growth of osteoblastic cells.^{9,10} Owing to its wide range of applications in the biomedical field, and more specifically in bone tissue regeneration (osteogenesis), HA has been under intense study and several researchers have published their findings on the use of HA in various reputed journals and conference proceedings (Fig. 2). Therefore, the aim of this review is to compile all the information related to HA, discuss and summarize it in a structured manner, and provide future prospects, so that readers can easily access the desired information.



Fig. 2 Articles indexed in Scopus from 2006 to 2016 with HA.

1.1 Chemical composition and synthesis of HA

As mentioned earlier, HA is a member of the apatite family (consists of Ca and phosphates) with the general formula $\text{Ca}_5(\text{PO}_4)_3\text{OH}$, and unit cell formula $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$. In the unit cell of HA, Ca and phosphates are arranged in such a manner that four Ca atoms are surrounded by nine O atoms of the phosphate moieties at the M1 position, and the other six Ca atoms are surrounded by the remaining six O atoms of the phosphate moieties at the M2 position. M1 and M2 are the crystallographic positions for all Ca atoms (Fig. 3).^{4,11,12} Regardless of its origin, HA contains traces of impurities such as phosphite ions (PO_3^{3-}), chloride ions (Cl^-), fluoride ions (F^-), and hydroxyl ions (OH^-). PO_3^{3-} and Cl^- have been reported to weaken the HA structure, whereas F^- and OH^- are known to enhance the apatite strength.

Owing to the importance of HA in bone tissue regeneration and as a drug delivery material, various techniques have been reported for the preparation of HA nanoparticles and nanorods. Particle size and morphology are two important factors for suitable biomedical applications of HA.¹³ High emphasis has been placed on both HA synthesis and particle size, but very few reports are available on the morphology control of HA. The techniques commonly used for the preparation of HA particles include chemical precipitation (CP),^{14,15} hydrothermal (HT),^{16,17} electrospinning (ES),¹⁸ electrospraying (ESp),¹⁹ solid state (SS),^{14,15,20,21} microwave irradiation (MI),^{22–24} self-propagating combustion (SPC),²⁵ emulsion and microemulsion (EMe),²⁶ surfactant-assisted precipitation,^{24,27–29} chemical vapor,²⁹ and flux cooling (FC).³⁰ All these methods used for the preparation of HA particles result in different morphologies (Table 1) and chemical compositions.³¹ The number of annual publications using these methods is shown in Fig. 4. As observed from the figure, the most commonly used methods for the synthesis of HA are CP, HT, and sol-gel (SG). However, the preparation of HA nanoparticles with the right stoichiometry, high aspect ratio, and high crystallinity remains difficult to obtain. Conventional wet mechano-chemical methods are able to control the stoichiometry of the final product. In these methods, precursor solutions of $\text{Ca}_3(\text{PO}_4)_2$ and $\text{Ca}(\text{OH})_2$, are co-





Fig. 3 Crystal structure of HA showing the *c*-axis perpendicular to 3 *a*-axes lying at 120° of each other (left), and projection of the HA structure on the 001 plane (right) (this figure was reproduced with permission from John Wiley and Sons).⁴

precipitated and then calcined at 1000–1200 °C to refine the crystal structure. If the Ca/P stoichiometry is not adjusted to 1.67 during the precipitation step, then other phases such as β -tricalcium phosphate (TCP) and CaO will be formed at lower and higher values, respectively. Furthermore, it is difficult to control the simultaneous nucleation, crystal growth, coarsening, and/or agglomeration, which occur during the precipitation of the HA particles.³² To synthesize HA particles with a tailored size, various additives such as allylamine hydrochloride,³³ cetyltrimethyl ammonium bromide,³⁴ and poly(acrylic acid) have been used.³⁵

Kehoe *et al.* synthesized HA powders using a fractional factorial, resolution IV *via* a two-level experimental design to assess the critical process parameters (reagent addition rate, reaction temperature, stirring speed, ripening time, initial Ca concentration, and the presence of an inert atmosphere) and their effect (main and interaction) on the final HA powder characteristics, such as phase composition, purity, crystallinity, crystallite size, lattice parameters, particle size, and particle size distribution. This investigation of selected parameters showed an influence on one or more of the responses investigated, as either a main or an interaction effect. However, the ripening time and stirring speed significantly affected the majority of the five responses. In addition, the reaction temperature has

a significant effect on the final phase composition, lattice parameters, and particle size as well.³⁶ Low-temperature HT methods showed regulated ageing and a controlled size.³⁷ Despite being a relatively simple technique for the synthesis of HA, SS methods have barely been reported (Fig. 4). The main limitations of this technique are the use of high temperature and longer reactions times, which allow the formation of different calcium phosphate phases (α - and β -TCP ($\text{Ca}_3(\text{PO}_4)_2$)). The presence of these phases in the final product adversely affects the mechanical properties of HA, and thus, make its performance unpredictable.³⁸ The SG technique is another common method to synthesize HA. This technique has advantages over the above-mentioned methods as it allows the use of low processing temperatures (lower than 400 °C), results in a homogeneous molecular mixing, and generates nanosized particles. K. Agrawal *et al.* fabricated HA using the solution gel technique. They used phosphoric pentoxide (P_2O_5) and calcium nitrate tetrahydrate ($\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$) as precursors. After the mixing and gelation steps, they calcined the sample at 400 and 750 °C for 8 h to remove residues and enhance the crystallinity of the HA. They then characterized their final compound (HA) *via* various spectroscopic techniques.³⁹ In general, to prepare uniform small-sized HA nanoparticles/nanorods, fast precipitation methods are preferred, which have the advantage of

Table 1 Synthetic techniques used for the synthesis of HA and characteristics of the resulting material

Method	Processing time (</>24 h)	Temp (°C)	Size (μm)	Degree of crystallinity	Phase purity	Morphology	Size distribution	Cost	Ref.
CP	>	RT	>0.1	Low	Variable	Diverse	Variable	Low	37 and 41–44
ESp	>	—	75 × 40 nm	—	—	—	—	Low	19
ES	>	—	10 × 10–30	—	—	Fibers	Variable	Variable	18
MI	—	—	100 × 25 nm	High	High	Diverse	Narrow	Variable	22–24
FC	<	500	18.0 × 2.1	High	—	Hexagonal cylinders	—	Variable	30
HT	<	150–400	>0.05	High	High	Needle-like	Wide	High	16 and 17
SS	>	1050–1250	>2.0	High	Low	Diverse	Wide	Low	14, 15, 20 and 21
SG	>	37–85	>0.001	Variable	Variable	Diverse	Narrow	Variable	39
SPC	<	170–500	>0.45	Variable	High	Diverse	Wide	Low	25
EMe	>	RT	>1.0 (emulsion), >0.005 (micro)	Low	Variable	Needle-like	Narrow	High	26





Fig. 4 Articles indexed in Scopus from 2007 to 2016 with the synthesis of HA using different techniques.

avoiding nucleation and crystal growth (which are the basis for attaining monodisperse particles). In addition to fast precipitation methods, aging techniques can also be used for the preparation of HA nanoparticles/nanorods.⁴⁰ However, the synthesis of nanosized HA particles with the proper stoichiometry, high aspect ratio and good crystallinity are still difficult to obtain.

1.2 Characterization techniques

Various characterization techniques have been used to characterize HA in the form of nanoparticles, rods, discs, and powder. These characterization techniques are broadly classified as

spectroscopic and direct visualization techniques. The details about the advantages and limitations of these techniques are given in Table 2. Fourier transform infrared (FTIR) and Raman spectroscopies, and X-ray techniques have been regularly used for the characterization of HA. Spectroscopic techniques such as FTIR and Raman are used to confirm the chemical composition of the HA. Both techniques use characteristics specific to the material to determine the composition. For example, HA shows various characteristic peaks in the FTIR spectrum at 3572 cm^{-1} (OH stretching), 1087, 1046, 1032 cm^{-1} (P–O stretching) and 631, 602, and 574 cm^{-1} (O–P–O bending) (Fig. 5I(a)). On the other hand, Raman spectra show characteristic HA peaks in the 1045–963 cm^{-1} region. The peak at 1045–1043 cm^{-1} is attributed to the asymmetric stretching of P–O, whereas the peak at 963 cm^{-1} confirms the presence of tetrahedral PO_4^{3-} (Fig. 5I(b)). The presence of this peak at this position further confirms that HA is ordered and highly crystalline.⁴⁵ X-ray photoelectron spectroscopy (XPS) is a surface-sensitive quantitative spectroscopic technique that provides elemental composition at parts per thousand range, and information about the empirical formula, chemical state, and electronic state of the elements within the material. In the XPS spectra, HA shows photoelectron signals (Fig. 5I(c)) at 347.9 eV (Ca 2p), 536.1 eV (O 1s), and 133.2 eV (P 2p). These signals are commonly used to confirm the presence of HA.⁴⁰ The X-ray diffraction (XRD) profile of HA usually shows characteristic diffraction peaks of nHA at 26.1, 28.45, 30.1, 32.90, 35.97, 40.19, 41.82, 53.56, 55.75, 57.40, 69.12, 74.45, and 77.56°, which correspond to the 002, 102, 210, 112, 300, 212, 130, 213, 321, 004, and 104 planes of the hexagonal HA unit cell (Fig. 5I(e)).⁴⁶ Energy dispersive spectroscopy (EDS) is another very important technique used in combination

Table 2 Classification, advantages, and limitations of various techniques used to characterize HA nanoparticles and inorganic powder materials (these are normally analyzed by Raman spectroscopy rather than by FTIR)

Technique	Advantages	Limitations
Spectroscopic techniques		
FTIR	Measures the intensity over a narrow range of wavelengths at a time; no external calibration is required; provides accurate results, and identifies even small concentrations of contaminants	Inorganic materials are not easily analyzed by FTIR spectroscopy
Raman spectra	Highly specific (provides a chemical fingerprint of the material); inorganic materials are easier to analyze by Raman spectroscopy than by FTIR	The Raman effect is very weak. The detection requires a sensitive and highly optimized instrument; fluorescence of the impurities or of the sample itself can hide the Raman spectrum; sample heating by the intense laser radiation can destroy the sample or cover the Raman spectrum
XPS	Provides unique information about the chemical composition of a material	Slow, poor spatial resolution, requires high vacuum
XRD	Powerful and rapid (<20 min) technique, provides an unambiguous mineral determination; data interpretation is relatively straightforward	Homogeneous and single phase materials are best for identification of an unknown; peak overlay may occur, and it is worse for high angle reflections
EDS	Chemical microanalysis technique used in conjunction with SEM; provides unique peaks characteristic of the atomic structure of the atoms; quick and versatile technique	Comparatively lower precision
Direct visualization		
SEM	Direct visualization, high resolution	NP aggregation during the sample preparation
TEM	Direct visualization, high resolution	NP aggregation during the sample preparation, electron beam damage, preference for electron-dense atomic species
AFM	High size resolution, 3D profile	Slow speed, limited scanning area





Fig. 5 Techniques used to characterize HA: I (a–e) spectroscopic (reproduced with permission from Springer and INTECH)^{10,45,48} and II ((a–e) ((b) is the unpublished figure owned by authors)) direct visualization (reproduced with permission from Hindawi).^{47,49}



with scanning electron microscopy (SEM). In EDS technique, peaks appear due to the unique atomic structure of the atoms. For HA, the unique peaks of Ca, P, and O appear at their characteristic positions, as shown in Fig. 5I(d). Direct visualization using SEM, transmission electron microscopy (TEM), and atomic force microscopy (AFM) provides information about the morphology, shape, size, and dispersion of the HA nanoparticles (Fig. 5II(a–e)). Occasionally, the information obtained from SEM is not very conclusive, and thus, supplementary techniques such as TEM and AFM are required to determine the dispersion, shape, and size of the HA nanoparticles.⁴⁷

2. Surface modification of HA nanoparticles

To control HA nanoparticle dispersion, delay the dissociation from the composite materials, and make them application-specific, researchers have studied several options. One of these options is surface modification. The aggregation of nanomaterials is a common issue encountered by researchers. Aggregation depends on the type of nanomaterials, surface energy, reagents or method used for nanoparticle synthesis and hydrophilic–hydrophobic interface between the solutions (polymeric or non-polymeric nanoparticles). To overcome this difficulty in HA nanoparticles, various emulsifying agents (like detergent or surfactant) and bridging agents (chemicals or coating the nanoparticles surface with some molecule) have been used. For example, Kim *et al.* used hydrostearic acid⁵⁰ as a bridging agent between hydrophilic–hydrophobic HA and poly(L-lactic acid) (PLA). Bridging agents (mediators) are mainly amphiphilic compounds that contain both hydrophobic and hydrophilic moieties. Therefore, to achieve better HA dispersions in solution, the surface modification of HA *via* these bridging agents is promising not only for the fabrication of HA nanoparticles with uniform dispersions, but also for the preparation of composite scaffolds. In addition, another positive aspect of the surface modification is that it delays the dissociation of HA nanoparticles from the composite materials. Since HA nanoparticles are mostly used in biomedical applications, these bridging agents must fulfill certain requirements. Hence, they should not: (i) compromise the biocompatibility, (ii) be cytotoxic, and (iii) alter the physiological or biological

properties of the nanoparticles or fillers. Different materials and methods have been used for the surface modification of HA (Table 3). For example, Tanaka *et al.* modified HA using decanoic and hexanoic acids, and these compounds bonded strongly to P–OH *via* hydrogen bonding. These acids can be removed from the sample (restoring the original P–OH surface) by outgassing the sample at 500 °C. The use of carboxylic acids has two advantages: (i) using long chain acids (chain with more carbon atoms than acetic acid) would convert HA from hydrophilic to hydrophobic, and (ii) since animals contain different types of carboxylic acid moieties in the form of proteins, metabolic inhibitors, vitamins, and fats, HA interacts strongly with the carboxylic acids of these species.⁵¹ Tanaka *et al.* further improved their experiments and modified the surface of HA using pyrophosphoric acid (PP). PP reacted with the P–OH surface of HA, resulting in the formation of additional P–OH moieties along with phosphoric acids.⁵² Followed by Tanaka *et al.*, Kim *et al.* used oleic acid (belongs to the fatty acids group, and it is amphiphilic and non-cytotoxic to human osteoblastic cells) to modify the surface of HA.^{53,54} Shimabayashi used the amphiphilic properties of the surfactant sodium dodecyl sulphate (SDS) to modify the surface of HA.⁵⁶ They further immobilized bovine serum albumin and polyvinyl pyrrolidone (PVP) on the SDS-modified HA (HA-s); in the first case, SDS interacts electrostatically with the HA, while in the latter case, HA interacts hydrophobically with PVP. A similar but reverse mechanistic design was also reported by Haider *et al.* using carbon nanotubes and gelatin systems.⁵⁵

Inorganic materials such as silanes are also suitable surface modifiers, as they stimulate adhesion, have a bonding ability, and can act as bridging agents for both inorganic and organic materials. Silanes anchored on HA will remain non-cytotoxic, compared to free silanes, which are considered cytotoxic.^{57,58} It is for this reason that silanes have been used in the fabrication of dental⁵⁹ and bone implanting materials.⁶⁰ Researchers have also used grafting (using polymers to improve dispersion and growth factors to enhance cell growth) to modify the surface of HA. For example, Hong *et al.* grafted L-lactide (LA) to HA using ring opening polymerization.⁶¹ The same authors also grafted poly(L-lactide) (PLLA) to HA *via* the carbonyl moieties of PLLA. They reported that the HA particles could be easily dispersed in PLLA using this method.⁶² Grafting techniques

Table 3 Different techniques used to modify the surface of HA

Surface modifying agent	Technique	Ref.	Surface modifying agent	Technique	Ref.
Hydrostearic acid	Direct reaction	50	PLLA	Grafting	62
Decanoic and hexanoic acids	Direct reaction	51	BMP-2	Grafting	49
Oleic acid	Direct reaction	54	Insulin	Grafting	10
SDS	Direct reaction	54	Serum albumin and PVP	Grafting	56
PP	Direct reaction	52 and 64	Vinylphosphonic acid	Grafting	65
Silanes	Coating	58	Ethylene glycol methacrylate phosphate	Polymerization	66
Glycidoxy propyltrimethoxy silane and [3-(2-aminoethylamino)propyl]trimethoxysilane	Coating	67	LA	Polymerization	61
Polyacrylic acid	Coating	68	γ-Benzyl-L-glutamate <i>N</i> -carboxyanhydride	Polymerization	69
γ-Methacroyloxytrimethylsilane	Coating	70	L-Phenylalanine <i>N</i> -carboxyanhydride	Polymerization	71





Fig. 6 Schematic representation of the grafting of (a) BMP-2, and (b) insulin on the surface of nHA (reproduced with permission from Springer and Hindawi).^{10,63}

have been used recently by Haider *et al.* as well to immobilize growth factors such as bone morphogenetic protein (BMP-2) and insulin on HA (Fig. 6(a) and (b)). They have not only observed good dispersion HA, but also an enhanced cell proliferation on the HA-based scaffolds. The figure was reproduced with permission from Springer.^{10,49}

3. Ion-doping of HA nanoparticles

Doping of HA with foreign ions is another method that has been focused by researchers in recent years. Even though doping of foreign ions into HA has been documented in literature, however, not many studies were focused on cell culture or these were only focused on one type of doping ion. This lack of research on the doping of foreign ions into HA for use in cellular culture made it difficult for researchers to evaluate the effect of doping ions on the HA interaction with cell. Recently various ions such as strontium (Sr^{2+}), magnesium (Mg^{2+}), iron (Fe^{3+}), manganese (Mn^{2+}), zinc (Zn^{2+}), carbonate (CO_3^{2-}), silicate (SiO_4^{4-}), *etc.*, have been doped into the structure of HA. The presence of these

dopants in the structure of HA are significant for the bio-structure and biochemistry of the HA to be similar to natural bone.^{72,73} The doping of these ions into HA did not disturb the structure configuration, as HA is known for accommodating different ions: however, in some cases the crystallinity, morphology and solubility of HA were changed. Studies have shown that these changes did not affect cell behaviour, though in some cases cellular biocompatibility properties were changed. This issue was well highlighted by Zhao *et al.* who have shown that the selection of doping ions has a strong impact on cell behaviour *e.g.* a minute amount of Mg^{2+} ions in the HA (~ 1.5 wt%) could lead to a significant cytotoxicity in MG63 cells, but not in rMSCs. They further elaborated that HA internalization (endocytosis) is affected by the stabilization of negative surface charge. This effect was more pronounced in MG63 as compared to rMSCs. Research with Mg^{2+} doped HA further suggested that these nanoparticles could be used to kill cancer cells. This use of such particles will not only eliminate the need of conjugating anticancer drugs onto the nanoparticles but will also solve the issue associated with drug



toxicity.⁷⁴ Thus a proper selection of the foreign doping ions is imperative, if HA is intended for use in cell culture. In another study Subhadip *et al.* studied the effect of Sr^{2+} and Mg^{2+} dopants on the structural stability and biological properties of HA. They observed that the presence of Mg^{2+} and Sr^{2+} not only enhanced the stability of the particles but also improved the osteoblast response activities and density of the viable cell on a negatively charged surface for all cultural periods. Thus, they have concluded that metal ion dopants affect mineral metabolism in the bone remodelling process and enhance the osteoclast apoptosis as well as preosteoblastic cell proliferation. The presence of these dopants in the HA could lead to the designing of bone graft materials that could be of potential use in tissue regeneration for faster healing.⁷⁵ Li *et al.* doped HA with manganese (Mn^{2+}) and iron (Fe^{3+}) ions using wet chemical method and ions exchange mechanism. The samples showed no deviation from the structure, morphology and crystal size, and were single phased and B-type carbonated HA (carbonate groups partially replaced phosphate groups in the lattice structure). Both dopants showed no toxic effect on osteoblast cells. Fe^{3+} doped HA illustrated enhanced osteoblast cell adhesion and increased negative surface charge as compared to Mn^{2+} HA and pristine HA.⁷⁶ The evidences in the above mentioned studies is clearly showing that metal cations dopants can extract more physicomechanical and biological benefits, when incorporated into at issue-engineering construct. Most of the cations dopants are non-harmful even at considerable high concentration. Having said this and as described above some cations can cause toxicity if their concentration is increased beyond a certain level. Furthermore, appropriate considerations must be given to the type of a chemical compound that intended for use in metal doping of the biomaterials. This precaution is imperative because some chemical entities might cause toxicity in the host tissue, although the cations used their selves are not toxic. Besides, very little information is on hand regarding the bioavailability of the metal dopants when released from biomaterials *in vivo* during biodegradation. In addition, the mechanism of action of dopants at the cellular level has not well reported in the literature. Despite all these limitations and drawbacks, the existing research has provided exciting evidences, which suggests that further work with metal doped bioceramics is not only going to be well-timed but also worthy of fast healing the tissue implant.⁷⁷

4. Applications of HA nanoparticles

The interaction between inorganic materials and protein molecules is of pivotal interest for researchers working in the field of biomaterials, biomineralization, biosensors, biochemistry, and industry.⁷⁷ Scientists working in the field of drug delivery and tissue regeneration, more specifically in bone tissue regeneration, have acquired sufficient knowledge to determine the interaction between HA and various proteins. HA crystals have two binding sites, *i.e.* "C" sites (Ca^{2+}) and "P" sites (PO_4^-) owing to its chemical composition and specific orientation. Within a crystal, the "C" sites are arranged in

a rectangular manner, whereas "C" sites are arranged along the *c*-axis surfaces in the hexagonal configuration (Fig. 7).^{79–81} By suspending HA in aqueous media, the "C" sites are exposed and they acquire positive charge, which is mainly attributed to the release of OH^- ions from the crystal surface. The "C" sites, after the release of the OH^- ions, are left with positively charged Ca^{2+} ions. These positively charged Ca^{2+} ions interact with the negatively charged proteins. Chen *et al.* reported that $-\text{COO}^-$ ions with their negatively charged oxygen interact with the "C" sites of HA.⁸² Shaw *et al.* also confirmed *via* solid state NMR that terminal $-\text{COO}^-$ amelogenin ions interact with the HA surface.⁸³ On the other hand, "P" sites are negatively charged ions ($\text{P}(\text{PO}_4^-)$), which interact with the positively charged proteins. In addition to their reactivity with proteins, HA is also used under physiological conditions. Therefore, HA has also been used in chromatography for separating various proteins.^{84,85} Since HA, due to its biocompatible properties, has a higher potential for applications in the biomedical field, scientists working in the orthopedic field have been using HA on the bone implanting material. Besides its use as an implanting material, HA has been used in gene delivery, cancer therapy and as a bio imaging tool. Gene therapy is a long recognised promise for the treatment of a wide range of incurable diseases through replacement of the missing or defective genes, catalyzing the destruction of cancer cells, causing the cancer cells to revert back to normal tissue, promoting the growth of new tissue or stimulating regeneration of the damaged tissue.^{86,87} However, this promise still remains unfulfilled, primarily due to the lack of a safe and competent vehicle for the delivery of gene. HA is considered a potential candidate for this purpose as HA possess characteristics of an ideal gene vehicle. The potential of the HA as gene delivery vehicle was evaluated by Tan *et al.*, who compared the non-viral gene delivery of neutrally charged HA and positively charged zirconia (ZrO_2). In their work they found that a mixture of highly monodispersed aqueous suspension of HA nanoparticles, coated with protamine sulfate (PS), complexed efficiently with plasmid DNA and significantly



Fig. 7 Hexagonal crystal structure of HA with "C" and "P" sites. The figure was redrawn by the authors from the work of Uddin *et al.* and Okada and T. Matsumoto (reproduced with permission from INTECH).^{90,91}



enhanced trans gene expression *in vitro* as compared to ZrO_2 .⁸⁶ Similarly, Liu *et al.* proposed that HA-based nanoparticles could be used for bioimaging of living cells as well. For example, the authors prepared well dispersed fluorescein isothiocyanate-labeled HA nanoparticles using SiO_2 as template for imaging tissues or intracellular structures.⁸⁸ Thus with fluorescence imaging method using nontoxic fluorescent hydroxyapatite (fHA) nanomaterials, visualization of the complete organ in complicated biological processes is possible.⁸⁹ A detailed description of some of the most important applications is given below.

4.1 HA as a drug delivery carrier

Most biological Ca apatites are known for their irregular morphologies, which result from the presence of residue materials in the apatites. However, the surface of HA is rough, which favours the use of HA in biomedical applications. The rough surface together with the "P" and "C" sites of HA facilitates protein binding during the process of mineralization.⁹² The importance of the rough surface and the "P" and "C" sites has been confirmed by the interaction of amino acids with HA.⁹² During the last decade, research has been focused on the use of HA as a drug delivery carrier, and this is evident from the increasing number of research articles each year (Fig. 8) (this data was obtained from Scopus). Several researchers are investigating the use of HA nanorods/nanoparticles as a drug delivery carrier for various proteins (growth factors) and drugs.^{10,49,93} For example, Haider *et al.* modified the surface of HA with a spacer using an amino acid (glutamic acid). They immobilized proteins growth factors (protein drug) on the free terminal groups (carboxylic ($-\text{COOH}$) or amine ($-\text{NH}_2$) moieties) of the spacer. The drug-loaded/modified HA nanorods/nanoparticles were then blended with a polymeric solution to carry the drug-loaded HA to the target site.⁵¹ Yewle *et al.* oriented proteins on the surface of HA using bifunctional hydrazine bisphosphonate (HBP) with hydrazone bonds (Fig. 9(a)). They first modified the surface of HA with HBP, followed by the immobilization

of the protein with aldehyde functionality. The aldehyde functionality was obtained *via* oxidation of the N-terminal of serine and threonine. From the analysis of the results, they concluded that the oriented proteins showed better activity as compared to the absorbed proteins.⁹⁴ Furthermore, Li *et al.* fabricated Gd-doped luminescent HA and mesoporous SrHA nanorods and reported that their HA-modified aptamers could be used as a therapeutic agent against cancerous cells (Fig. 9(c)).⁹⁵ Son *et al.* used HA coated with Ti discs (Fig. 9(b)) for the delivery of bioactive drug molecules (dexamethasone (DEX)) using biodegradable poly(lactide-co-glycolide) (PLGA) nanoparticles. In their study, they coated the surface of Ti with HA and electrostatically immobilized DEX-loaded PLGA nanoparticles on the coated surface.⁹⁶ The sample with the lowest surface roughness S_3 (as shown in Fig. 9(b)) showed not only a sustained release of DEX over a period of 30 days, but also developed more extracellular matrix on it. Recently, Ibrahim *et al.* used a very simple methodology and produced mesoporous HA nanopowder with a pore volume of $1.4 \text{ cm}^3 \text{ g}^{-1}$ and a surface area and of $284.1 \text{ m}^2 \text{ g}^{-1}$ from raw eggshells at room temperature using a simple two-step procedure. The as-prepared HA nanopowder showed high loading of ibuprofen drug ($1.38 \text{ g g}^{-1} \text{ HA}$), which was used as model drug. Good dissolution and controlled release of the drug *via* solute-saturated supercritical CO_2 was observed (Fig. 9(d)).⁹⁷ Besides these studies, various research groups around the world are working on the sustainable use of HA as a drug delivery carrier on the surface of implanting materials (in one way or the other) to improve the bioactivity of the implanting material.^{98–101}

4.2 HA as a coating material

To restore a damaged bone, it is imperative to design a bone implanting material that can be used not only for curing the bone defect, but also for reconstructing the bone. There are several difficulties in fabricating suitable bone implanting materials. For example, the first difficulty that commonly appears is the synchronization between the implanting material and bone during the bone remodelling process (resorption and reparative process). The implanting material should not have any adverse effects on the immune system. To date, the methods used for managing bone defects are allografting, autografting, and synthetic (fabricated using HA or naturally occurring biocompatible substances) bone implantation techniques. However, all these techniques have their limitations. For example, the implanting materials sometimes resorb before the osteogenesis, rendering the material ineffective.^{103,104} In other cases, infections due to the implants result in incidence of disease and even death, which are very costly to the patient in particular and society in general.¹⁰⁵ Owing to these reasons, it is necessary to devise a way to design and fabricate suitable bone tissue regenerating implants. For the fabrication of an implanting material, four vital aspects of an implant should be considered: (i) the implanting material should have a signal carrier, (ii) the implanting material should present a morphogenetic signal, (iii) responsive cells for the signals should be present in the implanting sites, and (iv) the existence of a well



Fig. 8 Articles indexed in Scopus from 2007 to 2016 with the application of HA in drug delivery systems.





Fig. 9 (a) Schematic for the oriented proteins on the surface of HA using bifunctional HBP and immobilization of proteins (reproduced with permission from ACS),⁹⁴ (b) three-dimensional (3D) interactive display images obtained by optical profiling of the Ti disc surfaces. R_a : roughness average, R_t : roughness height. (S1) Resorbable blast media (RBM) on a Ti disc, (S2) RBM on a HA-Ti disc, and (S3) DEX-loaded PLGA particles immobilized on a HA-Ti disc (reproduced with permission from John Wiley Sons),⁹⁶ (c) schematic representation of HA-based nanoprobe targeting bioimaging and drug delivery (reproduced with permission from Springer),¹⁰² and (d) illustration of the methodology, morphology, and drug release profiles of the spongy-like mesoporous HA prepared using raw waste eggshells.⁹⁷

vascularised host bed.¹⁰⁶ Recently, a technique has been used where an implant is coated with a biocompatible and bioactive material to enhance the compatibility and bioactivity of the implanting material. For bone tissue regeneration and implanting, the most common coating agent used is HA. The importance of HA as a coating material is evident from the number of articles that have been published in the last decade using HA as coating material (Fig. 10). HA has been used as a coating material because it chemically resembles natural bone. Furthermore, coating of the implanting material surfaces (usually metallic prosthesis) with HA enhances the osteointegration with bone.¹⁰⁷ Sintered HA can adhere tightly to the bone, thus hindering degradation. J. He *et al.* used a thin coating of HA on a Ti implanting surface *via* a plasma spraying method.¹⁰⁹ The advantage of this method is that it avoided a sub-optimal fatigue of the sintered HA. Other well-known methods that have been used to coat HA on the surface of implanting materials (metallic) sputter coating, include dip coating, pulsed laser deposition, hot pressing and hot isostatic pressing, thermal spraying, electrostatic spraying, electrophoretic deposition,

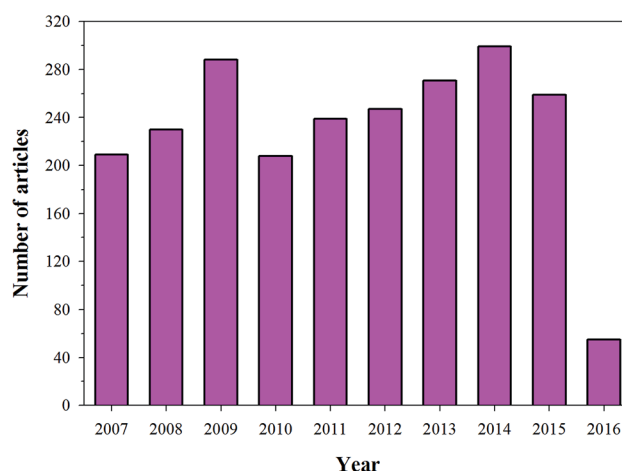


Fig. 10 Articles indexed in Scopus from 2007 to 2016 using HA as a coating material.

solution gel, pulse laser deposition, and an electrostatic layer-by-layer sputtering method.^{105,109} The advantages and disadvantages of these coating techniques are presented in Table 4. Researchers working in the field of bone tissue regeneration have concluded that the shear strength of a Ti alloy coated with HA is similar to that of cortical bone.¹¹⁰ Rigo *et al.* coated a silicon nitride surface with HA to enhance its bioactivity and biocompatibility. Silicon nitride has good mechanical strength, but it has limited biomedical applications due to its inertness.¹¹¹ LeGeros concluded that osteoblastic cells could easily interact with the HA-coated surface.¹¹² These interactions will lead to the formation of osteoids and thus to chemical and biological interactions between the implanting material and bone in a safe manner. The authors also concluded that besides the osteoconductive properties, HA (with a 3D morphology) also shows osteoinductive properties when it binds and interacts with bone morphogenetic proteins (BMPs).¹¹² All the techniques used for coating HA on the surface of metal alloy prosthesis and other orthopedic implanting materials helped in stimulating new bone growth under optimized conditions.¹⁰⁵ However, limitations associated with the necessity and effectiveness of

HA coatings for various anatomic sites, the strength of HA coatings to withstand physiological loads without fragmentation, and the issues related to third-body wear by HA particles restrict its applicability.¹⁰⁵

4.3 HA-based composite materials

As mentioned earlier, polymers are of great use in the field of tissue regeneration. The polymers used in bone regeneration can be either biodegradable or non-degradable and can have a natural or synthetic origin. In the latter half of the 19th century, Urist *et al.* concluded that polymers induced bone formation in animals.^{113–115} Similarly, Kulkarni *et al.* concluded that the morphology of polymer-based implants and the implanting sites were pivotal in stimulating bone tissue regeneration. They reported that tubular absorbable polymer implants could enhance bone tissue regeneration in long bones.¹¹⁶ Various synthetic polymers have been tested directly or in combination with HA to mimic natural bone matrix. Among them, PLGA,¹¹⁷ PLA,^{118,119} and poly(caprolactone)^{120,121} are the most frequently used polyesters. In most polymeric composites, HA has been used as an enforcer to provide

Table 4 HA coating techniques and their characteristics (reproduced with permission from INTECH)¹⁰⁸

Techniques	Advantage	Disadvantage	Thickness
Dip coating	Inexpensive Coatings applied quickly	High sintering temperatures Thermal expansion results in an amorphous coating	0.05–2 mm
Sputter coating	Coats 3D complex porous substrates Uniform coating thickness on flat substrates	Fragile due to the thickness Only coats visible areas Expensive and time-consuming Unable to coat complex 3D porous substrates	0.02–2 µm
Pulsed laser deposition	Uniform coating thickness on flat substrates	Risk of amorphous coatings Only coats visible areas Expensive and time-consuming Unable to coat complex 3D substrates	1–10 µm
Electrostatic spray deposition	Uniform coating thickness on flat Substrates Relatively cheap	Only coats visible areas	1–10 µm
Electrophoretic deposition	Uniform coating thickness High deposition rates Coats complex 3D porous substrates	Fragile Cracks during coating High sintering temperatures	0.1–200 µm
Electrostatic spraying, plasma spraying	High deposition rates	Only coats visible areas Coating decomposition due to high temperature Rapid cooling may result in amorphous coatings	30–200 µm
SG	Coats 3D complex porous substrates Low processing temperatures Relatively cheap and forms very thin coatings	Processing in controlled atmosphere	<1 µm
Hot pressing, hot isostatic pressing and sintering	Dense coatings	Unable to coat complex 3D porous substrates Differences in coating elasticity Expensive Interaction with or changes due to the encapsulation of the material	0.1–10 mm
Layer-by-layer coatings	Controlled loading efficiency and release kinetics of the encapsulated biomolecules, various	Require the use of a few hundred layers to avoid a burst release of biomolecule Requires an acidic solution for an effective loading Low mechanical stability	0.1–0.61 µm



mechanical strength and bioactivity to the composite material.^{10,49,93} The importance of HA in bone tissue regeneration has been reported extensively. Even though research on the application of HA composites is advancing slowly, it has been steadily increasing. This is evident from the number of articles published on this topic over the last decade (Fig. 11). Various researchers have fabricated HA/polymer scaffolds for use as a drug delivery carrier, improved bone tissue regeneration (grafting proteins and drugs on the surface of HA) and treatment of various bone defects such as osteoporosis. Haider *et al.* immobilized insulin and BMP-2 on the surface of HA. The modified HA was incorporated in PLGA to prepare HA/PLGA composites. They concluded that after the protein was immobilized on HA and its composting with PLGA, the composite enhanced osteoblastic cell growth and accelerated osteogenesis.⁴⁹ In another study, the authors also immobilized pamidronic acid on the surface of HA to treat osteoporosis. Due to excessive osteoclastic activity, bones are prone to break; hence, they concluded from their experiments that pamidronic acid reduced the osteoclastic cell activity. They also showed that the presence of HA enhanced the osteoblastic cell activity.¹²² Ignjatović *et al.* fabricated a HA/PLLA composite, whereas Wang *et al.* synthesized a HA/polyethylene composite. They concluded that these composites have sufficient mechanical strength to be used in bone regeneration.^{123,124} Bakos fabricated a HA/collagen composite by blending to mimic natural bone. The composite showed bend strength of 5.37 kPa, and they concluded that this material could find applications in cases where better cohesion and shape stability are required.^{123,125–127} Dalby *et al.* prepared a HA/poly(methylmethacrylate) (PMMA) composite and revealed that this material acts as a substrate for human osteoblast-like cells.^{123,125–127} In another study, a group of researchers tried a variety of combinations of chitosan and HA to improve the biocompatibility and mechanical strength of chitosan-based materials. For example, a composite containing high molecular weight chitosan and HA particles was prepared

by freezing and lyophilisation. The composite showed better mechanical properties than medium molecular weight chitosan ones.¹²⁸ Furthermore, various collagen compositions with HA have been reported as well, which have shown better biocompatibility and enzymatic degradability.^{129–132}

4.4 HA-based ceramics in bone tissue regeneration

Natural bone is a ceramic composite comprised of calcium phosphate, collagen, and water. Apart from these major constituents, polysaccharides, lipids, and proteins are also found in natural bone in small amounts. Calcium phosphate is present in bones in the form of HA crystals, which provides rigidity to the bone. The HA components are similar to those of bone with some impurities of Na⁺, Mg²⁺, and F[−] ions. HA crystals in bone are in the nanometer range for length and width. The presence of HA in an implant not only enhances the osteoinductive behaviour of the implant, but it also improves the osteoconductive properties between the implant and the bone cells (osteoblastic cells).¹²⁵ Owing to the similarity between HA and inorganic cement of natural bone, HA has been frequently used either in pristine, composite, or ceramic form for bone tissue regeneration. This is evident from the number of articles published on this topic between 2007 and 2016 (Fig. 12). To achieve the desired goal (safe and satisfactory bone regeneration), the selected materials for an implant should have similar properties to those of natural bone. The knowledge of the natural environment of bone osteoblastic cells is imperative, if infection-free bone regeneration is desired. Several HA bioceramics have been prepared to mimic natural bone. However, as mentioned above, it is necessary that the implants provide an environment close to the natural one. For example, besides other factors, the implant must have a mechanical strength close to that of the bone. Hence, the difference in mechanical strength between the implant and bone should not exceed a factor of 55 (between moduli) with respect to cortical bone, or it will have an adverse effect on the living cells of the bone (due to the bone shielding by the bioceramic implanting

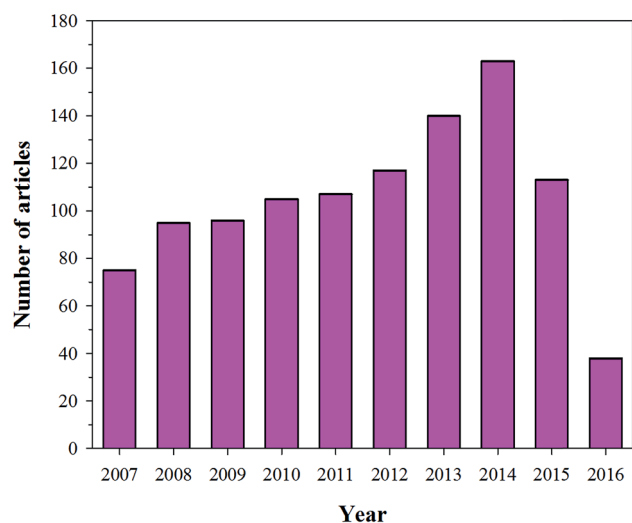


Fig. 11 Articles indexed in Scopus from 2007 to 2016 with HA-based composite materials.



Fig. 12 Articles indexed in Scopus from 2007 to 2016 with the application of HA in tissue engineering.



Table 5 Synthesis methods and various characteristics of HA bio-ceramics (reproduced with permission from MDPI)¹⁴⁵

Process	HA form	Sintering	Compressive strength	Pore size	Porosity
HA + ammonium methacrylate in polytetrafluoroethylene mould, freeze dried and then sintered	Powder	1300 °C	16 MPa 65 MPa 145 MPa	Open Unidirectional 50–150 µm	>60% 56% 47%
Polymer foams coated, compressed after infiltration, and then calcined	Powder	700–1300 °C	—	100–200 µm	—
Directly injectable calcium orthophosphate cement, self-hardens, mannitol as porogen	Nanocrystalline	—	2.2–4.2 MPa (flexural)	0–50% macroporous	65–82%
Slurries of HA prepared: gel-casting + polymer sponge technique, and then sintered	Powder	600 °C (1 h) 1350 °C (2 h)	0.5–5 MPa	200–400 µm	70–77%
Slurry of HA with polyoxyethylenelaurylether (cross-linked) and then sintered	Powder	1200 °C (3 h)	2.25 MPa 4.92 MPa 11.2 MPa	500 µm 200 µm Interconnected	~77%
Electrophoretic deposition of HA, and then by sintered	Powder	1200 °C (2 h)	860 MPa	0.5 µm 130 µm	~20%

material).¹³³ In addition, excessive hardness of the bioceramic implanting materials will lead to wearing of the bone. HA-based bioceramics can be fabricated by various methods according to the requirements of the experiment or implanting site (Table 5). For the preparation of an implanting material with high tensile strength, sintered (dense) (Fig. 13(a)) ceramics (bending strength 38–250 MPa, compressive strength 120–900 MPa, and tensile strength 300 MPa), are used instead of porous ceramics (bending strength 2–11 MPa, compressive strength 2–100 MPa, and tensile strength ~3 MPa (ref. 125 and 134)). Porous ceramics (Fig. 13(a)) with interconnected pores fabricated by burning the organic materials^{135,136} are mainly used in drug delivery, bone in growth, bone fixation, *etc.*^{135,137–139} Research is growing on the preparation of bio-ceramics to mimic natural bone cement. For that purpose, various reinforcing agents such as metals, nanoparticles, and fibers have been tested. However, adding these materials may sometimes affect the biocompatibility of HA. To overcome this issue, Ahn *et al.* fabricated HA-based Zr bioceramic materials with a fine structure and enhanced mechanical properties, and suggested that it could be used both in dental and orthopedic implanting materials.¹⁴⁰ Yamashita *et al.* used an electrospinning technique to control the morphology of HA-based ceramic materials. Using this technique, the HA crystals on the negatively charged surface of the HA ceramic were grown in simulated body fluid.¹⁴¹ Several

research groups have been studying various systems to fabricate an ideal scaffold, and thus, they have been testing different materials such as HA/collagen,^{126,142} HA/PLGA and HA/PLA,^{123,143} HA/PMMA, among others.^{127,144} However, despite the recent advances, there is a necessity for more research to prepare an infection-free implanting material with sufficient mechanical strength.

5. Future prospects

With the advances in materials science and nanotechnology, new materials have been regularly fabricated for several purposes. HA is used as an orthopedic implant, coating on metallic materials, and drug delivery carrier. Substantial advances have been made regarding our knowledge and understanding of cellular reactions to HA and related ceramics. However, HA still has limited applications in load-bearing systems. Hence, researchers are now focusing on the development of materials that could bear loads (sufficient mechanical strength). They are using various strategies to combine HA and related calcium phosphate materials with natural polymers to fabricate composite materials with mechanical strength, elasticity, and toughness similar to that of bone. Moreover, scientists are also facing a challenge to develop strategies to avoid possible infections. Research has been advancing in this direction using nanotechnology. HA coating is also playing a vital part in the enhancement of bone formation, as these provide an osteoconductive and osteoinductive approach. The biological properties may be amplified further by grafting growth factors and other useful molecules to HA to fabricate a real osteoinductive platform. However, limitations related to the necessity and effectiveness of HA coatings in different anatomic sites, robustness of HA coatings to withstand physiological loads, and third-body wear by HA particles, restrict its application. Further research to answer these questions will improve the mechanical and biologic aspects of HA-based implants and optimize their safety and efficiency.



Fig. 13 Example of sintered HA used as an artificial bone substitute (reproduced with permission from MDPI)¹⁴⁶ and graded porous HA materials using tape-casting and lamination (reproduced with permission from MDPI).¹⁴⁷



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