Translation of bone wax and its substitutes: History, clinical status and future directions

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Abstract  Bone wax, primarily composed of beeswax and softening agent, is a century-old material used to control bleeding of disrupted bone surfaces by acting as a mechanical barrier to seal the wound. The current bone wax products are commonly packed in easy-to-open foil in the form of sterile sticks or plates, with excellent malleability and smooth consistency, enabling cost-effective and easy handling approach for bleeding control. It has also been reported that the inert nature of bone wax causes complications including foreign body reaction, infection promotion and bone healing inhibition. With the advances in biomaterials and the market boost of bone haemostatic materials, the arena of bone wax substitute research has expanded to a wide spectrum of material formulations and forms. However, the development of substitutes of bone wax for translation is a pivotal yet challenging topic because currently a potential candidate is recommended to be just as simple to use, effective and inexpensive to produce as traditional bone wax but also be absorbable and osteogenic. This review provides an overview of bone wax including its history, clinical applications and associated
Introduction

Bone contains abundant channels for blood and bone marrow. When it is surgically incised or traumatically fractured, osseous haemorrhage can be a difficult problem to control, especially in the highly vascular bones of the spine and sternum. Medical sterile bone wax is an essential material for haemostasis of bone during orthopaedic surgeries, thoracic surgeries, neurological surgeries and so on. Along with its widespread use, complications such as foreign body reaction, bone healing inhibition and infection promotion associated with bone wax are observed. With the growing knowledge in biomaterials and the boost of market of bone haemostatic materials, bone wax substitute research is thriving. An overview of bone and its substitutes together with evolution of their design criteria is carried out in this work, providing information for the innovation and translation of bone haemostatic agents in the near future.

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The applications of bone wax

Both cancellous and cortical bones contain vascular tissues, and when the bone is incised or fractured, damage to its vasculature can cause osseous haemorrhage that is...
sometimes too severe to be controlled by natural haemostasis. In this case, haemorrhage should be effectively controlled to avoid further pathologic consequences such as tissue necrosis and eventually mortalities due to blood loss [11]. At the present time, some options can be used in clinical settings for bone haemostasis: (a) the use of classical absorbable haemostatic agents such as collagen and oxidised cellulose, (b) the application of electrocautery and (c) use of bone wax. However, the use of oxidised cellulose is limited because of its inappropriate knitted fabric form and lack of adherence within the bone, causing problems in sealing irregular surfaces and pores of defective bone; collagen, on the other hand, in various forms, alone or in combination with fibrin and suspended in various delivery vehicles, has been proposed as a bone haemostatic agent but problems with storage stability, cohesiveness and biocompatibility have prevented practical fruition [12,13]. The use of electrocautery, which thermally sears oozing blood vessels and closes them, is time-consuming and can easily induce severe thermal damage to tissues, which may further delay osteogenesis and allow soft tissue ingrowth that interferes with normal bone union [13,14]. Instead, bone wax is highlighted by its ease of operation, satisfactory cohesion to bone, malleability and cost-effectiveness. It is mainly applied for bone haemostasis during orthopaedic surgeries [15], thoracic surgeries [16] and neurological surgeries [3], but can be occasionally extended to dental and jaw surgeries [17–19]. In addition to directly being applied to the wound site, bone wax can be used to modify surgical tools for blood control purposes. For example, in percutaneous endoscopic cervical discectomy via the anterior transcorporeal approach for cervical intervertebral disc herniation, bone wax was smeared onto the endoscopic burr to control bleeding without obvious interference with bone healing [20]. Similarly, it can also be used to prevent the leakage of blood through the lumen of a cannulated screw after arthroscopic repair of the anterior cruciate ligament [21].

Concerns of bone wax

Clinical practice has uncovered numerous complications associated with bone wax since its development. Although bone wax has potential and promise for haemostasis application, the related complications may outweigh its benefits. As shown in Fig. 2, a series of complications in surgeries were reported, such as failed bone healing, foreign body reaction, granuloma growth, thrombosis, infection and nerve compression [22–46].

A rat calvarial bone model was used to define the local reactions of bone to bone wax [47]. It was confirmed bone regrowth was markedly impaired by the presence of bone wax. Inert bone wax most typically encompassed the bone margins in a collar-like fashion, and active bone production was observed only beyond this collar and then connected mostly to the external periosteum. Moderate-to-severe inflammation, foreign body reaction and fibrous reaction were observed in the lesion. The observations are in consistent with histologic findings associated with the implantation of bone wax in a rat tibia model, which typically includes foreign body reactions and a lack of bone formation [48]. Sorrenti et al. [49] investigated the response of human tibia to bone wax. At the early stage, a nonspecific

![Figure 1](image)

**Figure 1** Operating process of bone wax. (A) Bone wax is a sterile and flat slice wrapped in a tiny bag; (B), (C) and (D) softened bone wax is malleable and easy to be shaped.
Inflammatory response was noted (6 months), followed by an increase in fibrous tissue with foreign-body giant cells (9 months). After 13 months, mature fibrous tissue with no inflammatory response was observed. As bone wax dramatically interferes with bone healing, it should be used sparingly and restricted in the sites where fusion is highly desired.

The effect of bone wax on the ability of cancellous bone to clear bacteria was examined using a Staphylococcus aureus and rabbit model, indicating that as a foreign body, bone wax can significantly diminish the ability of bone to clear bacteria [50]. Similarly, in a rat model of chronic S. aureus osteomyelitis, the infection-promoting potential of sterile bone wax was also observed [51]. In a case series of 19 patients presenting with osteomyelitis of the sternum after cardiac surgery, bone wax applied to the oozing sternum halves was postulated to be the possible cause of this problem [52]. Because of the risk of induction of infection, bone wax must never be used in contaminated fields. Besides, cleansing of bone wax in iodine is recommended after manual manipulation during clinical practice.

In addition to these postoperative complications, the efficiency of control of bone bleeding using bone wax is also in dispute. For example, in a prospective randomised study on 400 thoracic surgical patients undergoing isolated coronary bypass surgery, bone wax application after median sternotomy showed no benefits in blood loss control [9]. In contrast, in a report of total knee arthroplasty, the application of bone wax was reported to be safe and effective for reducing total blood loss and maintaining higher haemoglobin levels. No consensus is achieved yet, and surgeons may readily use bone wax at their own experience and discretion.

So far, the status of bone wax has been overviewed, and a table summarising the advantages and disadvantages of bone wax is thereby presented as a guideline for the development of a new generation of bone haemostatic materials (Table 1).

**Substitutes of bone wax**

It has been recognised that the major factor causing the aforementioned postoperative complications such as bone union prevention, infection promotion and foreign body reaction is the intrinsic inertness and poor biocompatibility of bone wax. The attempt of developing absorbable bone wax can be dated back to 1950, when Geary and Frantz [8] reported experimental haemostatic bone wax by combining Carbowax, PEG and oxidised cellulose together. Although

| Advantages and disadvantages of bone wax. | | |
|---|---|---|
| **Advantages** | **Disadvantages** |
| Low cost | Inertness |
| Easy handling | Bone union prevention |
| Malleability | Foreign body reaction induction |
| Inertness | Granuloma growth induction |
| Sealing capacity | Infection promotion |
| Bone adherence | Lack of inherent haemostatic quality |
| Long clinical history | Undesired immigration |
| | Thrombosis induction |
this formulation failed in dental surgery studies [15], this work guided the direction of development of bone wax substitutes by clarifying that absorbability is the top priority in the design of substitutes.

From 1980 to 2000, numerous bone wax substitute prototypes have been reported in the literature, such as fatty acid salts [53], fibrin/collagen paste [54,55], gelatin paste [56], glycolic or lactic acid/glycerol oligomers [57,58], partially deacetylated chitin hydrochloride [59], PEG/microfibrillar collagen paste [60], polydioxanone/natural oils [61] and polyorthoester [54]. Unfortunately, none of these formulations are in widespread use or launched in market, which suggests that it has been difficult to combine the beneficial characteristics of traditional bone wax with the advantages of an absorbable material.

In the 2000s, an ideal bone wax substitute was suggested to be just as simple to use, effective and inexpensive to produce as traditional bone wax but would also be fully absorbable, noninflammatory and biocompatible. According to this criterion, water soluble wax composed solely of alkyene oxide block copolymers (Pluronics) was developed in 2001, which has material, application and haemostatic characteristics that are similar to those of bone wax, but its absorbable property avoids the negative biological effects [62]. Inspired by this pioneering work, a commercially available water-soluble alkyene oxide copolymer—based bone wax substitute (Ostene, “absorbable bone wax”; Ceremed, Inc., Los Angeles, CA, USA) was launched in 2006 [63]. Similar to bone wax, Ostene can be softened by manual manipulation before use and sticks well to bleeding bone as a tamponade. The material dissolves in the implanted site within 24–48 h, allowing the early phases of bone healing to occur. Because of its solubility, Ostene provides the potential to address adverse reactions associated with inert bone wax [64,65] (Fig. 3). This formula was later revised to develop new products in the literature, such as a putty-like mixture of alkyene oxide copolymers and carboxymethylcellulose sodium salt (Absorbable haemostatic bone putty; Abyrx, Inc., Irvington, NY, USA) [66] or a glue-like miscible blend of PEG–polypropylene glycol–PEG (PEG–PPG–PEG) copolymer and pregelatinized starch [67].

Back in 1992, haemostatic agents including microfibrillar collagen flour, absorbable gelatin sponge and oxidised regenerated cellulose powder were applied as alternatives to bone wax in iliac bone procurement, showing no bone regeneration inhibition as planned [68]. Such extended applications of existing haemostatic agents, including hydrated gelatin powder [69], gelatin paste [56], gelatin-thrombin matrix sealant [70], fibrin solution [71], patient-derived fibrin sealant [72], autologous platelet-poor plasma gel [73], fibrin dressing [74] and gel-like mixture of agar and coagulation factors [75], for control of bone bleeding were popular in research. However, whether these agents can act as alternatives to bone wax in bone haemostasis is still questionable because of limited animal studies and lack of clinical trials.

In the past few years, the reciprocal influence between the conceptual strategy of developing absorbable substitutes and the better understanding of haemostasis and bone regeneration has led to the evolution of bone wax substitutes from a sole haemostatic agent to hybrid agent with both haemostatic and bone regeneration capabilities. On the one hand, some fast absorbable haemostatic agents are not free of risk of complications, possibly causing allergic reaction and retarding bone regeneration to some extent, making it a must-addressed issue in designing bone wax substitute [76]. On the other hand, in many surgical scenarios, scaffold-induced bone healing is highly desirable [77,78]. One typical strategy in the present day is the adoption of bioceramic cement–based paste/putty in bone haemostasis, whose phase and as-formed matrix can help to both stop bleeding and enhance osteogenesis [79–81]. To further improve blood clotting efficiency and handling properties, supplements such as alginate [82], cellulose [83] and chitosan [84] can be blended. For example, currently Zhang et al. reported a self-curing bone wax substitute by mixing tricalcium silicate (C₃S) cement and 585 bioactive glass/chitosan/carboxymethyl cellulose with KH₂PO₄ setting solution, which enables haemostasis, injection and bone cell proliferation [80]. Calcium Apatite bone tamponade (CAAP; Skeletal Kinetics, LLC., Cupertino, CA, USA) is an FDA-approved product composed of calcium phosphate, sodium silicate solution and a mixing system (mixing bowl, pestle and spatula). Its operation procedure is kind of similar to that of cement: (1) open powder vial, pour powder into the mixing bowl, and gently tap the vial to ensure maximum transfer of powder; (2) slowly pour the

![Figure 3](image-url)  
Figure 3  The rabbit tibia model was inoculated with *Staphylococcus aureus*, introduced into the intramedullary canal through a defect created at the anteromedial facet of the proximal tibia. After 4 weeks, the cross section of the rabbit tibia shows normal bone development in the cortical window of the Ostene and control samples. The bone wax cross section shows signs of osteomyelitis with no sign of bone healing [64].
liquid vial into the mixing bowl; (3) use the pestle to vigorously mix in circular motion the powder and liquid for approximately 1 min, and make sure to reincorporate the material collected on the pestle into the mixing process to achieve a proper mix and (4) when mixed together, it forms a cement-like paste that can be applied directly to sites of bleeding bone; the resulting hardening scaffold from the paste is composed of hydroxyapatite similar to the mineral phase of native bone tissue, enabling bony ingrowth and bone regeneration [79,85]. The drawback of this formula is the necessity to manually mix the powder and setting solution before use, increasing operation steps and contamination risk. As a solution to this issue, a ready-to-use paste of calcium phosphate cement, PEG and pregelatinised starch was reported [86] (Fig. 4). After exposure to a humid environment, the PEG phase dissolved and was exchanged by penetrating water that interacted with the calcium phosphate precursor to form highly porous, nanocrystalline hydroxyapatite via a dissolution/precipitation reaction. Simultaneously, pregelatinised starch could gel and supply the mixture with liquid-sealing features. The novel formulation was found to be cohesive and malleable, and after hardening under aqueous conditions, it had a mechanical performance (~2.5 MPa compressive strength) that is comparable to that of cancellous bone. The concerns of this formula are the sensitivity of calcium phosphate cement to moisture during storage and the lack of clinical evaluations. Nevertheless, the ready-to-use design demonstrates high translational potential as it would simplify the surgery and make the material closer to the handling characteristics of bone wax.

Owing to the potential risk of infection, antibiotics have been incorporated into bone wax substitutes [87,88]. Such an attempt paves the possibility of using bone wax substitutes to deliver therapeutic agents to enhance blood clotting or bone regeneration. For example, water-soluble bone wax substitutes were suggested to serve the additional purpose of acting as an absorbable matrix for short-term drug delivery (e.g., bone morphogenic proteins, antibiotics and cytokines) to the damaged bone, providing additional benefits by promoting osteogenesis, improving fusion rates, reducing inflammation and preventing postsurgical infections [62]. The challenges mainly rely on the shelf life of therapeutic agents and their release and cost control. There is increasing amount of evidence over the past decade demonstrating that the delivery of selected bioactive ions can trigger specific biological responses such as microbial inhibition, blood clotting stimulation, angiogenesis and osteogenesis [89–91]. In reported bone wax substitutes, bioceramics such as bioactive glass, tricalcium silicate and calcium phosphate have been added to promote bone regeneration, which are also potential candidates for bioactive ion delivery. The design of bone wax substitutes as bioactive ion carriers requires the control of different ion release profiles after the material is exposed to blood and analysis of their consequent biological responses in situ in blood clotting and bone regeneration.

Outlook and summary
Since the introduction of bone wax over 125 years, this century-old haemostatic agent is still being used for controlling bone bleeding and sealing. Clinical complications originated from the nature of bone wax spur the continuing research of substitutes, and a potential candidate is recommended to be just as simple to use, effective and

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**Figure 4** Traditional calcium phosphate cement (CPC) requires the addition of a liquid curing agent and manual mixing to finally form HA, whereas ready-to-use bone wax substitute can be much simpler and easier to use [86]. After exposure to blood, the PEG phase dissolved and was exchanged by penetrating blood, which stimulates HA matrix formation in situ. HA, hydroxyapatite; PEG, polyethylene glycol.
in vivo standardised evaluation of the adhesive and sealing capability of agency. This controversy definitely calls for the stand-achieved between the relevant surgeons and regulatory judgement. Contradictory evaluation results are sometimes such failure reports are largely based on the subjective awareness of end-users. In addition, market players in the US are more active in research and development of or introducing new products with improved efficacy to the market. Europe is the second major market for bone wax. The market in these regions is driven by the increase in the number of surgical procedures, innovations in bone wax products, rise of bone diseases and accidental fracture cases. The bone wax market in the Asia-Pacific region is also growing rapidly at a growth rate of 3% during the forecast period, driven primarily by the developing countries such as India and China because of the soaring market needs.

With the growing knowledge and technology advances in biomaterials and the boost of market, the arena of bone wax substitute research has expanded to a wide spectrum of material formulations and forms to meet the evolving design criteria. The innovation and translation of bone wax substitutes is expected to thrive in the near future.

As the outcomes of research move towards translation and commercialisation, it will also be important to eluci-date thorough standardised evaluation systems of bone wax and its substitutes, which will lead to improved material design and generation. From an operational point of view, it is highly suggested that bone wax and its substitutes can easily detach from gloves when pressed into cavity but can exhibit strong adhesion to the bleeding bone surfaces or the capability to seal the bleeding defects. In FDA enforcement reports of bone wax, the failure of bone wax is largely attributed to its loss of adhesion to the host site after storage [93]. However, according to the FDA information, such failure reports are largely based on the subjective judgement. Contradictory evaluation results are sometimes achieved between the relevant surgeons and regulatory agency. This controversy definitely calls for the standardised evaluation of the adhesive and sealing capability of bone wax. So far, the haemostatic performance has been generally studied through in vivo animal experiments by drilling holes in the bone and plugging with bone wax. Instead of complicated animal testing, an in vitro model has been proposed by Suwanprateeb et al [94] for sealing capability testing. In brief, acrylic glass tubes with a length of 2.00 mm and an inner diameter of 3.00 mm were filled with water up to a height of 1.91 m, which corresponds to systolic blood pressure (18.68 kPa, 140 mmHg), and sealed with cone-shaped samples; the constructs were stored at room temperature and monitored until their failure. However, there are drawbacks for this model design: (1) in body, the blood actually diffuses from the damaged cancellous bone, different from the flowing of water in a tube, and (2) water is far different from blood, unable to reflect blood clotting and other haemodynamic behaviours. Besides, the relevant testing of shelf life and lone-term adhesiveness, preservation of samples is commonly lacking in the research and development of bone wax substitutes. Taken together, reliable and standardised testing methods for bone haemostatic agents with haemostatic and adhesive capabilities are needed in the future.

Conflict of interest

The authors have no conflicts of interest to disclose in relation to this article.

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