The role of CKIP-1 in osteoporosis development and treatment

Osteoporosis is a systemic skeletal disorder characterized by reduced bone mass and deterioration of bone microarchitecture, which results in increased bone fragility and fracture risk. Casein kinase 2-interacting protein-1 (CKIP-1) is a protein that plays an important role in regulation of bone formation. The effect of CKIP-1 on bone formation is mainly mediated through negative regulation of the bone morphogenetic protein pathway. In addition, CKIP-1 has an important role in the progression of osteoporosis. This review provides a summary of the recent studies on the role of CKIP-1 in osteoporosis development and treatment.

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Article focus
- An overview of the role of CKIP-1 in osteogenesis and osteoclastogenesis and its relationship with osteoporosis.
- Reveal the possible treatment of osteoporosis through manipulating the biological function of CKIP-1.

Key messages
- The structure and function of casein kinase 2-interacting protein-1 (CKIP-1).
- CKIP-1 negatively regulates bone formation through bone morphogenetic protein pathway.
- Progress in research on the role of CKIP-1 in osteoporosis treatment.

Strengths and limitations
- This review summarizes the mechanism of action of CKIP-1 in the pathogenesis of OP and predicts CKIP-1 siRNA therapy may be a novel treatment option for OP.
- Whether CKIP-1 can interact with other signaling molecules to regulate the BMP pathway or regulate bone metabolism via other pathways remains unconfirmed.
- The osteoblast-targeted treatment using CKIP-1 siRNA has only been tested in animals and similar drugs have not yet been developed for human application.

Introduction
Osteoporosis (OP) is a systemic multifactorial skeletal disorder characterized by reduced bone mass, deterioration of bone microarchitecture and increased bone fragility, resulting in a tendency of fracture and leading to possible lifelong disability or death. Therefore, with an expanding ageing population, OP has become an important public health problem and the understanding of the physiological and biochemical regulations of osteoblast-osteoclast balance, along with the mechanisms of bone destruction and loss, and the development of possible novel therapeutic strategies has become of real importance.

Structure and function of casein kinase 2-interacting protein-1 (CKIP-1)
CKIP-1 is a transcription factor that was first identified in the gene expression profile in the liver of fetuses. This gene is commonly known as CKIP-1 or Pleckstrin Homology domain-containing family O member 1 (PLEKHO1). The full-length CKIP-1 cDNA encodes a protein of 409 amino acids, of approximately 46 kD; the N-terminus of CKIP-1 protein contains a pleckstrin homology (PH) domain, and the C-terminus contains a hypothetical leucine zipper (LZ) motif (352 amino acids (aa) to 380 aa), along with five proline-rich motifs throughout the protein. CKIP-1 acts as a scaffold adaptor to mediate the interactions with multiple signalling and cellular proteins, such as CK2a, CPa, PI3K/Akt, c-Jun/JunD, ATM/p53, IFP35/Nmi, Smurf1. CKIP-1 therefore...
CKIP-1 induces osteoporosis through negative regulation of bone formation

In order to maintain their normal functions bone tissues undergo constant metabolic turnover. However, with ageing, the loss of bone components gradually leads to a negative balance in bone mass. There is increased activity of osteoclasts along with loss of osteogenic function which contributes to an overall reduction in bone formation. This imbalance of ‘bone loss > bone formation’ forms the cellular basis for the development of OP. 

Lu et al demonstrated that bone density and bone mass increased with the age of CKIP-1-knockout (KO) mice and the activity of osteoblasts in these mice was significantly higher than those of wild-type mice. Zhou et al have shown that in a rat mandibular distraction osteogenesis model, the small interfering RNA of CKIP (siRNA) significantly promoted bone formation both in vitro and in vivo. This indicates that the physiological level of CKIP-1 is an important negative regulator of bone formation, suggesting that high levels of CKIP-1 can lead to OP.

**CKIP-1 and primary osteoporosis.** Primary OP is the most common type of OP and is mainly caused by ageing, decreased physiological functions of organs and reduced secretion of sex hormones. There are two types of primary OP, postmenopausal (Type I) and senile (Type II). Type I is a high-turnover OP (bone resorption > bone formation). Using an ovariectomized rat model, Zhang et al evaluated the efficacy of CKIP-1 siRNA delivered by a bone-targeted delivery system (DSS₆-liposome) and examined bone morphometric parameters, bone mass, and bone structures. They found that therapeutic CKIP-1 siRNA intervention could significantly promote bone formation without affecting bone resorption, indicating that CKIP-1 plays a significant role in reversing bone loss during Type 1 OP.

Type II OP, is mainly found among individuals aged ≥ 70 years with a male to female ratio of 1:2. Ling et al studied the efficacy of CKIP-1 siRNA for treating Type II OP in old male Sprague-Dawley rats, (also assessed by examining bone morphometric parameters, bone mass and bone structures), and concluded that there was inhibition of bone formation by osteoblasts.
hand, Liu et al\textsuperscript{21} reported that, with aging, expression of CKIP-1 was increased in patients with bone fractures and that there was an association between reduced bone morphogenetic protein (BMP)-Smad signalling and bone formation. Hence using genetic approaches, the loss of CKIP-1 in osteoblasts, could promote BMP-Smad signalling and thereby alleviate the reduction in bone formation in Type II OP.

**CKIP-1 and secondary osteoporosis.** Secondary OP induced by diseases or drugs, can be classified as: (1) congenital e.g. osteogenesis imperfecta or homocystinuria (2) endocrinological (3) nutritional deficiency; induced by vitamin D and K deficiency, long-term calcium insufficiency, or long-term insufficiency of protein or elements including magnesium, manganese, strontium, and zinc (4) blood disease-induced e.g. monoclonal gammopathy of uncertain significance, multiple myeloma, systemic mastocytosis, beta thalassemia major (5) drug-induced, e.g. long-term use of adrenal cortex hormones (6) renal, caused by chronic kidney disease (7) due to respiratory diseases and (8) weightlessness or disuse-induced, caused by long-term immobilization or from space flight.\textsuperscript{1}

Long-term usage of glucocorticoids can inhibit osteoblast differentiation and mineralization, promote osteoblasts and osteocytes undergoing apoptosis and reduce trabecular bone mass. In addition, it can inhibit bone matrix protein synthesis, suppress the action of vitamin D, reduce calcium absorption, increase calcium secretion and stimulate parathyroid hormone (PTH) secretion, leading to enhanced bone resorption and loss of bone mass, in turn contributing to OP.\textsuperscript{22} Glucocorticoid-induced OP (GIOP) is the most common drug-induced OP and one of the most serious side effects of glucocorticoid use. Liu et al\textsuperscript{6} showed that high expression of CKIP-1 during extracorporeal glucocorticoid treatment could inhibit osteoblast differentiation and mineral deposition in osteoblasts by disrupting the Smad-dependent BMP signaling pathway.

**CKIP-1 and mechanical stimulation on osteogenesis.** Weightlessness-induced OP is a type of disuse-induced OP as gravity has an important role in normal growth and function of the human skeletal system. In a microgravity environment, reduced mechanical stimulation in the skeletal system enhances osteoclast function and attenuates osteoblast function.\textsuperscript{23} Zhang et al\textsuperscript{24} reported that during a tail suspension test, which simulated the effect of a microgravity environment, the femoral bone in CKIP-1 KO mice was not changed significantly, suggesting that loss of CKIP-1 could alleviate microgravity-induced OP by influencing the process of bone formation.

Distraction osteogenesis (DO) is also becoming an effective therapy for bone defects, by stimulating endogenous bone regeneration. Zhou et al have shown that CKIP-1 silences inhibited cell apoptosis and improves calcification through the induced expression of Wnt-3a, \(\beta\)-catenin and osteocalcin (OCN) in rat mandibular distraction osteogenesis.\textsuperscript{18} Also in fracture repair in animals, Wnt signalling may be involved in the bone formation of chitosan/si-CKIP-1 in DO.\textsuperscript{25}

In summary, CKIP-1 can decrease osteogenic differentiation and cause primary OP through negative regulation of bone formation and can reduce bone formation by negatively regulating the BMP pathway in GIOP and Wnt-3a/\(\beta\)-catenin pathway in DO.

**CKIP-1 regulates bone formation through bone morphogenetic protein pathway**

Osteogenesis is comprised of two basic processes, bone formation and bone resorption,\textsuperscript{26} both of which involve various complex factors. These factors include numerous cytokines and signalling pathways, such as nuclear receptor-interacting protein 1 (NIRIP1), interleukin-1 receptor-associated kinase 3 (IRAK3), bone morphogenetic protein 7 (BMP7), SMAD family member 1 (SMAD1), MAPK3/C-X-C chemokine receptor type 4 (CXCR4), OPG/RANKL/RANKL pathway, Wnt/\(\beta\)-catenin pathway, cathepsin K pathway, and BMP pathway.\textsuperscript{27,28} The BMP pathway is the most important signalling during these processes, as it can induce osteogenic differentiation, enhance bone formation without activating resorption and can promote bone repair without affecting non-skeletal tissues.\textsuperscript{29}

BMP’s are members of the transforming growth factor-\(\beta\) (TGF-\(\beta\)) superfamily and are significant extracellular molecules which promote bone formation and induce osteoblast differentiation.\textsuperscript{30} There are two types of transmembrane serine/threonine kinase receptors for BMPs, namely, the type I BMP receptor (BMPR-I) and the type II BMP receptor (BMPR-II).\textsuperscript{31} As one of the most widely studied and most potent osteogenic activity-inducing members of the BMP family, BMP-2, plays an important role in osteogenesis by activating Smad signal transduction and regulating osteogenic gene transcription.\textsuperscript{32} The process of BMP-2 signal transduction begins with its binding to BMPR-II. Subsequently, BMPR-II phosphorylates BMPR-I and the activated BMPR-I activates downstream Smads.\textsuperscript{33} Both in vitro and in vivo studies have confirmed that the BMP-Smad signalling could regulate various aspects of the osteoblast lifecycle, including mesenchymal stem cell (MSC) differentiation into osteoblasts, osteoprogenitor proliferation, osteoblast mineralization and osteoblast-osteoclast coupling.\textsuperscript{34} Defects in the BMP-Smad signalling pathway often result in reduced bone formation and mass, with the development of OP. In addition, the activation of BMP-Smad signalling can lead to a bone sclerosis phenotype.\textsuperscript{34}

After the BMP-2 has mediated the activation of Smads (Smad1, Smad5 and Smad8), it initiates the Runt-related gene 2 (RUNX2) expression through the distal P1 and proximal P2 promoters of the proteins. RUNX2 plays a
vital role in bone formation and remodelling and it has been shown that RUNX2-KO mice display non-mineralized bone formation. This complete blockage of intramembranous and endochondral ossifications in mice has demonstrated the importance of RUNX2 in bone development. Smad ubiquitination regulatory factor 1 (Smurf1) is a member of the Nedd4 family, Homologous to the E6-associated protein Carboxyl Terminus (HECT) E3 ligases. It can promote the ubiquitination and degradation of substrates such as Smad1/5, mitogen-activated protein kinase kinase kinase 2 (MEKK2), and RUNX2; moreover, it has an important function during the osteoblast differentiation. Smurf1 is comprised of one HECT domain and two WW domains. A highly conserved cysteine residue at the C-terminus of the HECT domain can form a thioester bond with ubiquitin. However, once this highly conserved cysteine residue is mutated to alanine or glycine, the ubiquitination and protein-specific degradation activities of Smurf1 are completely lost. The WW domain is another important feature of Smurf1, located between 236 aa and 311 aa, and is approximately 30 amino acids long. It consists of two highly conserved tryptophan residues and one highly conserved proline residue and is capable of binding to small proline-rich (PPxY motif) peptides. Zhang et al have shown that mutation of the PY motif in the Smad protein (proline- and tyrosine-rich domain) abrogated the interaction between Smad and Smurf1, resulting in the inhibition of Smad degradation.

Lu et al reported that CKIP-1 could increase the affinity between the WW domain and its substrates by binding to the linking region between the two WW domains of Smurf1. This enhanced the ubiquitin ligase activity of Smurf1, promoting the degradation of Smad1/5 and negatively regulating the BMP signalling pathway. Moreover, CKIP-1 could bind to the proteasome subunit Rpt6 as a linker and be coupled with the Smurf1 proteasome to enhance degradation of ubiquitinated protein substrates and inhibit osteoblast differentiation. Therefore, CKIP-1 plays dual roles in the interaction between Smurf1 and its substrates and the recruitment of substrates to the 26S proteasome.

Taken together, CKIP-1 enhances ubiquitin ligase activity by binding to Smurf1, which in turn accelerates the degradation of bone formation-related substrates and results in the inhibition of new bone formation. Therefore, CKIP-1 plays an important negative regulatory role in the process of bone formation (Figure 2).

**Progress in research on the role of CKIP-1 in osteoporosis treatment**

Osteoporosis is easily misdiagnosed and can be difficult to completely cure. Ageing and oestrogen deficiency are the key pathophysiological mechanisms. Although a number of antiresorptive drugs are thought to be effective for OP prevention, they do not provide comprehensive protection, which may be due to postmenopausal state and age, both of which reduce the ability of recruitment of stem cells and osteogenesis. Currently, the antiresorptive drugs, such as selective oestrogen receptor modulators (SERMs), bisphosphonate and denosumab, inhibit bone resorption. Wang et al conducted a systematic review of ten different therapies used for postmenopausal OP and showed that PTH and zoledronic acid (ZOL) have the highest probability of treatment efficacy in preventing clinical vertebral fractures. However, these drugs have
several drawbacks. SERMs (such as raloxifene) when used long term, can lead to breast and endometrial cancer.47 Bisphosphonates have relatively high gastrointestinal toxicity which can induce, abdominal pain, gastritis and oesophagitis, and can also lead to jaw necrosis, femoral fractures and atrial fibrillation.48 To overcome these shortcomings, new therapeutic drugs for OP are being developed with optimized efficacy, reduced side effects, and, hopefully, enhanced patient compliance.49

Regulatory imbalance between the activity of osteoblasts and osteoclasts is thought to be the main cause of OP. There have been several studies which have explored the mechanisms of regulation of bone metabolism by osteoclasts.51-52 However, osteoclast-targeted treatments for OP are still a problem, as they are only able to reduce or decelerate the rate of bone resorption and are unable to effectively promote new osteogenesis.51-52 It is known that CKIP-1 acts as a negative regulator of bone formation and its gene knockdown could significantly increase bone density and bone mass.

The siRNA delivery is a promising approach for the treatment of various diseases. Zheng et al53 reported that CKIP-1 siRNA could reduce the expression of CKIP-1 mRNA and promote osteoblast gene expression and osteoblast mineralization. Guo et al54 also found that the cross-species CKIP-1 siRNA promoted osteoblast differentiation in human, rhesus monkey, rat and mouse osteoblast-like cells in vitro, with stimulated bone formation but without elevating bone resorption in healthy rodents and osteoporotic mice in vivo.54 Zhang et al55 reported that a chitosan/si-CKIP-1-biofunctionalized titanium implant significantly improved the in vitro osteogenic differentiation of MSCs and led to dramatically enhanced osseointegration in the in vivo rat model. Moreover, it has been shown in animals that biweekly intravenous injections of 7.5 mg/kg CKIP-1 siRNA could maintain long-term low expression levels of CKIP-1 mRNA.19,54 Zheng et al developed a liposome-based bone tissue-specific delivery system to enable osteoblast-targeted delivery of CKIP-1 siRNA, which was highly effective in inhibiting CKIP-1 expression in osteoblasts, promoting osteogenic gene expression, increasing bone mass and improving bone microarchitecture.53 Considering the highly efficient and specific silencing effect of siRNA, CKIP-1 siRNA therapy may be a novel treatment option for OP. However, this strategy has drawbacks as the introduction of external siRNA into mammalian cells can potentially induce an immune response. Also CKIP-1 plays a role in many other areas, such as in the regulation of the cytoskeleton, cell growth and apoptosis. Therefore, its clinical application will need further investigation.

Future prospects
This review summarizes the mechanism of action of CKIP-1 in the pathogenesis of OP. CKIP-1 reduces bone formation mainly through binding to Smurf1 and the subsequent specific negative regulation of the BMP pathway. However, whether CKIP-1 can interact with other signaling molecules to regulate the BMP pathway or regulate bone metabolism via other pathways remains unconfirmed. In addition, the osteoblast-targeted treatment using CKIP-1 siRNA has only been tested in animals and similar drugs have not yet been developed for human application. Furthermore, it is largely unknown whether CKIP-1 siRNA will have human specificity or will induce side effects. Therefore, further studies are required to fully understand the mechanism of CKIP-1 regulation in OP pathogenesis and its potential as a targeted treatment for OP in humans.

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Conflict of Interest Statement
None declared

Author Contribution
X. Peng: Designing the review.
X. Wu: Reviewing the literature, Writing the manuscript.
J. Zhang: Revising the manuscript.
G. Zhang: Revising the manuscript.
G. Li: Designing the study.
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