

# Estrogen-Related Receptor $\alpha$

Subjects: Biochemistry & Molecular Biology

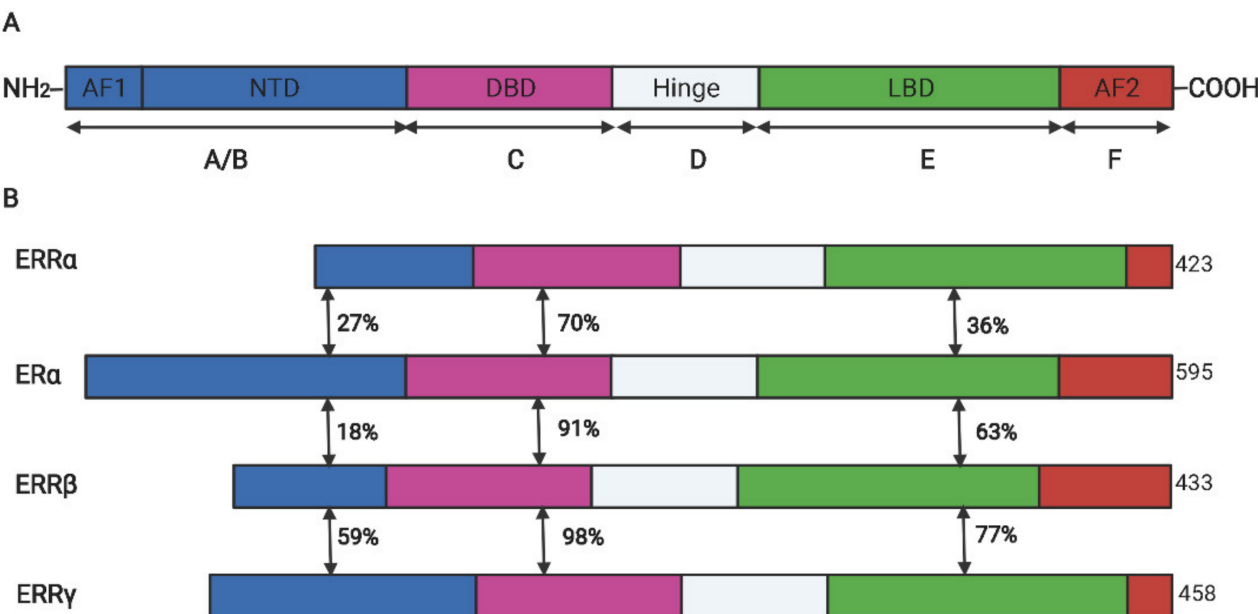
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Bone homeostasis is maintained with the balance between bone formation and bone resorption, which is involved in the functional performance of osteoblast and osteoclast. Disruption of this equilibrium usually causes bone disorders including osteoporosis, osteoarthritis, and osteosclerosis. In addition, aberrant activity of bone also contributes to the bone metastasis that frequently occurs in the late stage of aggressive cancers. Orphan nuclear receptor estrogen-related receptor (ERR $\alpha$ ) has been demonstrated to control the bone cell fate and the progression of tumor cells in bone through crosstalk with various molecules and signaling pathways.

Keywords: ERR $\alpha$  ; bone ; osteoclast ; osteoblast

## 1. Structure and Function of Estrogen-Related Receptor $\alpha$ (ERR $\alpha$ )

Estrogen-related receptor  $\alpha$  (ERR $\alpha$ ) belong to a subfamily of nuclear receptors and consist of three members: ERR $\alpha$ , ERR $\beta$ , and ERR $\gamma$ . Owing to lack of endogenous ligands *in vivo*, they are also called orphan nuclear receptors [1]. ERR $\alpha$  was firstly identified from a cDNA library by using the DNA-binding domain (DBD) sequence of estrogen receptor  $\alpha$  (ER $\alpha$ ) as a probe [1]. Similar to other steroid hormone receptors, ERR $\alpha$  has an N-terminal located ligand-independent transcription activation domain, a centrally located DBD encompassing two zinc fingers that mediate the interaction between the specific DNA motif and the receptor, and a C-terminal located domain that is responsible for ligand dependent transaction (also called ligand binding domain (LBD)). Between the DBD and LBD, a poorly conserved hinge domain mainly mediates the homodimer formation [2][3], the schematic diagram of structure domains between human ERRs and ER $\alpha$  is illustrated in **Figure 1**. Despite the high structural conservation among ERR $\alpha$  and ER $\alpha$ , the ligand of ER $\alpha$ , such as 17 $\beta$ -estradiol, cannot activate its transcriptional activity. Subsequent research revealed that ERR $\alpha$  recognized and binded on the estrogen-related response element (ERRE) but not the estrogen r element (ERE) on the promoter region of target genes to activate the transcription expression [4]. However, many target genes of ER $\alpha$  (like *pS2* and *LTF*) were simultaneously regulated by the two receptors in breast cancer cells [5][6]. In addition, the transcriptional activity of ERR $\alpha$  is largely dependent on its co-regulators such as peroxisome proliferator-activated receptor  $\gamma$  coactivator-1 (PGC-1)  $\alpha$  and PGC-1 $\beta$ , which is also distinguished from ER $\alpha$  that prefers to cooperate with SRC/p160 family [7].



**Figure 1.** The schematic diagram for the protein domains of ER $\alpha$  and ERR isoforms. (A). The conserved structure of nuclear receptors. (B). The domain similarity between ERRs and ER $\alpha$ . DBD, DNA binding domain; LBD, ligand binding

domain; AF, activation functional domain; NTD, N-terminal domain; The double-headed arrows indicated the sequence similarity between two receptors.

ERR $\alpha$  has also been clarified to exhibit various regulatory functions in physiological and pathological processes, including energy metabolism [8], organ development, immune response [9][10], and tumorigenesis [11][12]. ERR $\alpha$  is strongly expressed in high energy consumption tissues (such as heart, liver, muscle, and adipose tissues), and it can regulate lipid metabolism and thermogenesis in both white adipose tissues (WAT) and brown adipose tissues (BAT) [13]. Mice with ERR $\alpha$  knockout (KO) rapidly exhibited hypothermia under low-temperature conditions, and the expression and activity of uncoupling protein 1 (UCP1) induced by adrenalin in BAT was largely dependent on the presence and activity of ERR $\alpha$  [14][15]. In addition, ERR $\alpha$  was proved to modulate glycolytic metabolism via regulating the expression of essential enzymes including phosphofructokinase, hexokinase 2, and enolase 1 [16]. ERR $\alpha$  cooperating with PGC-1 $\alpha$  controlled the mitochondria metabolism and oxidative response to sensor the cellular ROS levels [17]. Beyond the function in metabolism, ERR $\alpha$  also acted in the innate immune response via regulating mitochondria ROS production and autophagy [9][10]. Furthermore, the high abundance of ERR $\alpha$  was frequently detected in many types of solid tumors (including colon, endometrium, bladder, ovary, breast, and prostate cancers), and its upregulation was associated with more aggressive phenotypes, drug resistance, and poor prognosis [11][18]. Mechanism exploration found that ERR $\alpha$  facilitated the proliferation and metastasis of tumor cells via upregulating oncogenes transcription and inducing metabolic reprogramming [12]. However, the role of ERR $\alpha$  in different cancers appears to be inconsistent. For example, ERR $\alpha$  almost functioned as an oncogenic factor in many types of carcinomas, but could serve as a favorable biomarker for the tamoxifen sensitivity in the patients with triple-negative breast cancer (TNBC) [19]. An alternative probability is that the functional performance of ERR $\alpha$  is tightly dependent on the specific cellular environment and post-translational modifications (PTMs) [18][20]. In addition, many specific inhibitors or natural products that function as agonists or antagonists of ERR $\alpha$  were identified, and these compounds have been implicated in the treatment of cancers or metabolism-disorder diseases [3][21]. Therefore, it is essential to explore the defined role of ERR $\alpha$  in a certain tumor model and achieve the precise therapy using the specific pharmacotherapy targeting on ERR $\alpha$ .

## 2. The Regulatory Role of ERR $\alpha$ in Osteoblasts

Epidemiological investigation showed that single nucleotide polymorphism (SNP) within *ESRRA* was closely associated with increased bone mineral density (BMD) of premenopausal women [22], and in situ hybridization assay found that *ESRRA* was consistently expressed in ossification zones during the mouse embryonic development stages, and it could directly bind to the promoter and activate the *OPN* transcription. These results indicated that ERR $\alpha$  may be involved in the process of the osteoblast progenitors differentiation and bone metabolism [23]. Indeed, ERR $\alpha$  was initially indicated to serve as a positive modulator for osteoblast differentiation and bone formation [24]; in vitro differentiation of osteoblast progenitors originating from rat calvaria or bone marrow was also accompanied with the continuous expression of ERR $\alpha$ , knockdown of ERR $\alpha$  by shRNA significantly impaired the proliferation and differentiation of osteoblast progenitors, while overexpression of this receptor increased the differentiation markers [25][26]. Moreover, estrogen could enhance the *ESRRA* transcription in a dose-dependent manner in the early differentiation stage of osteoblast progenitor cells [27][28], inhibition of ERR $\alpha$  in a culture medium induced a decreased colony number of mineralized cells [25], and the progenitor cells with ERR $\alpha$  deficiency were inclined to differentiate into adipocytes as evidenced through the increasing lipid droplet accumulation and expression of adipocyte markers (such as PPAR $\gamma$  or aP2). These studies suggested that ERR $\alpha$  served as a switch factor to favor the precursor cells differentiating into osteoblasts but not adipocytes [28][29]. Furthermore, other models from mice and humans also supported the promoted effect of ERR $\alpha$  on the osteoblast differentiation and functional exhibition, Rajalin et al., isolated the mouse mesenchymal stem cell from bone marrow and conducted ex vivo differentiation culture; they found that knockdown of ERR $\alpha$  induced the decreased cell proliferation, osteoblastic differentiation, and mineralization with reduced expression of bone sialoprotein (BSP) [30]. Cai et al., demonstrated that the mRNA and protein levels of ERR $\alpha$  were significantly increased during the late stage of osteogenic differentiation of human periodontal ligament cells (hPDLSCs), and the receptor maintained the osteogenic differentiation via regulating the ALP activity and osteogenesis-related genes expression [31], and another group established that ERR $\alpha$  interacted cooperatively with PGC-1 $\alpha$  in the promoter region of *osteocalcin* and enhanced its transcription expression to facilitate the osteogenesis [32]. In the human MSCs model, Huang et al., demonstrated that ERR $\alpha$  could directly bind to ERRE on the *Gls* promoter and transactivate its transcription, then further regulated glutamine anaplerosis and mitochondrial function, and this effect was required for osteogenic differentiation and impaired by aging [33]. Overall, these data indicated that ERR $\alpha$  functioned as a driving factor for osteoblast differentiation and was resistant to bone loss.

However, an increasing number of studies proposed that ERR $\alpha$  preferred to repress osteoblast differentiation and caused bone loss in vitro and in vivo. Firstly, the association between *ESRRA* SNP and increased BMD was not replicable in

another set of independent clinical data [34]. Two different groups consistently demonstrated that *ERRα*-KO mice exhibited increased BMD and enhanced osteoblast activity when compared with control, osteoblast progenitors derived from mice with *ERRα* deficiency had elevated expression of differential markers and were more liable to differentiate to matured phenotype but not adipocyte lineage [35][36]. One of groups further showed that the animals with *ERRα*-KO had resistance to the bone loss induced by aging or estrogen withdrawal mimicked with ovariectomy, indicating that targeting the receptor in menopause could prevent excessive bone loss and reduce the rate of fracture [36]. Moreover, Gallet et al., generated a conditional *ESRRA*-KO mouse in which *ERRα* was normally presented during the early differentiation stage of osteoblasts but lost its function in maturation; they found that *ERRα* could effectively respond to estrogen deficiency and inhibit the osteoblast differentiate and osteogenesis in the later maturation stage but not the early stage [37]. This phenomenon was consistent with the regulatory role of *ERRα* investigated in the mineralization of osteoblasts [35][38]. *OPN* had been verified to be the inhibitor of mineralization and one of the target genes of *ERRα*, *ERRα* deficiency was accompanied with a dramatic reduction of *OPN* and impairment of mineralization [39]. Consistently, *OPN*<sup>-/-</sup> mice were also resistant to the bone loss induced by estrogen withdrawal, this phenotype was similar to *ERRα*-KO animals [40]. Mechanism investigation revealed that *ERRα* might serve as a dual functional factor to modulate the activation of signaling pathways and osteogenesis-related genes expression (such as *Runx2* and *OPN*), and this effect was mainly dependent on the presence of its cofactor PGC-1β [41][42]. Studies using different models, such as different cell lines or mouse strains, may obtain inconsistent and contrary results. Therefore, more comprehensive research deserves to be conducted to uncover the defined role of *ERRα* in osteoblast differentiation and bone homeostasis.

### **3. The Role of *ERRα* in Osteoclasts**

Bone remodeling is not only associated with osteoblasts-mediated bone formation, but involves bone resorption regulated by the osteoclasts. Bonnelye et al., conducted the in vitro culture experiments and revealed that *ERRα* exhibited little effect on osteoclast differentiation but promoted adhesion and migration of osteoclasts via regulating the expression of *OPN* and integrin β3 chain and the alteration of cytoskeleton [43]. This conclusion was partially consistent with the role of *ERRα* in breast cancer [11]. However, in vivo studies using *ERRα*-KO animals challenged this conclusion. They found a specific phenotype with increased bone formation and reduced bone absorption in an *ESRRA*<sup>-/-</sup> mouse model owing to the decreased number and activity of osteoclasts [35][44]; mechanism investigation revealed that *ERRα* cooperated with PGC-1β and controlled the mitochondria biosynthesis and oxidative activity, which indirectly enhanced the osteoclast differentiation [44]. They further demonstrated that *ERRα* served as the key factor in modulating the RANKL- and rosiglitazone-mediated bone loss and functional performance of osteoclasts [44]. Consistently, PGC-1β was parallel with *ERRα* expression during the osteoclasts transition and the knockout of PGC-1β in mice exhibited osteopetrosis with abnormal morphology and the impaired bone-resorption activity of osteoclasts [45]. Subsequently, the significant role of *ERRα* in cholesterol metabolism and bone homeostasis was confirmed by Wei et al. [46] who identified that cholesterol acted as an endogenous ligand for *ERRα* to activate its transcriptional activity; this effect induced the inhibition of proinflammatory cytokines production by macrophage and facilitated the osteoclast differentiation [46]. Concomitantly, pharmacological inhibition on cholesterol synthesis and bone resorption by statin and bisphosphonate were also modulated by *ERRα* [46]. Considering the origin of osteoclast and macrophage from embryogenesis, *ERRα* may act as rheostat factor in the differentiation of macrophage and osteoclast. Congruously, MYC promoted the oxidative metabolism and drove osteoclastogenesis, the process of which was largely dependent on the *ERRα* action [47]. These fundamental studies suggested that the inhibition of *ERRα* by molecules or natural products may prevent or improve the bone loss induced by aging or hormone withdrawal. Indeed, Zheng et al., identified that carnolic acid, isolated from *Rosmarinus officinalis* L., could decrease the transcription activity of *ERRα* via reducing the interaction between *ERRα* and PGC-1β and triggering proteasomal degradation of *ERRα*, then suppress RANKL-mediated osteoclastogenesis and ovariectomy-induced bone loss [48]. Andrographolide, another natural compound targeted on *ERRα* from virtual docking screening, had been proven to suppress the formation of *ERRα*/PGC-1β complex and further reduce glutaminase metabolism during osteoclast differentiation. Inhibition on this regulatory axis attenuated osteoclasts-mediated bone resorption and bone loss in vivo [21]. Taken together, these data illustrated that *ERRα* functioned as a promoting effector to facilitate osteoclast differentiation and bone absorption involving various mechanism networks, targeting on the receptor would be a promising strategy for degenerative bone disorders.

### **4. The Role of *ERRα* in Cartilage and Osteoarthritis**

Osteoarthritis (OA) is characteristically associated with chronic inflammatory pathological feature and closely associated with articular cartilage dysfunction (including abnormal proliferation and erosion of cartilage) [29][49][50]. During the development of osteoarthritis, the elevated iNOS, IFNs, and TNFs levels in the microenvironment were detected, and LPS or TNF-γ induced the aberrant activation of NF-κB signaling in the pro-inflammatory (M1) macrophages, which could

generate pro-inflammatory cytokines (such as TNF- $\gamma$ , IL-6 and IL-12) to form a positive feedback loop and result in the degradation of cartilage [51].  $ERR\alpha$  has been reported to be involved in the innate immune response and functional performance of macrophage via modulating autophagy or TLR4/TRAFF6/NF- $\kappa$ B axis [10][52]. Consistently,  $ERR\alpha$  also performed a dual role in cartilage formation and OA progression. The effect of  $ERR\alpha$  on cartilage formation was initially involved in the regulation of SOX-9, Bonnely et al., reported that the receptor upregulated Sox-9 transcription and induced the differentiation and proliferation of chondrocytes [53]. The chondrocytes isolated from the OA patients treated with XCT790 (the specific reverse agonist of  $ERR\alpha$ ) showed the downregulation of SOX-9 expression in a dose dependent manner. These results were proven by Kim et al., in a zebrafish embryo model, where  $ERR\alpha$  knockdown led to the malformation of pharyngeal arch cartilage [54]. In addition,  $ERR\alpha$  also contributed to the cartilage degradation which is associated with IL-1 $\beta$  and MMP-13. IL-1 $\beta$  stimulation could increase the  $ERR\alpha$  expression via the PGE2/cAMP/PKA signaling pathway,  $ERR\alpha$  reversely activated the MMP-13 expression in OA progression [55]. This process may involve the macrophage action mediated by  $ERR\alpha$ /NF- $\kappa$ B signaling, but this possibility needs to be further investigated. The comprehensive effect of  $ERR\alpha$  on cartilage and OA had been reviewed by Bonnelye and Tang [29][56], but more systematic research is indispensable to reveal the crosstalk between inflammatory signaling pathways and  $ERR\alpha$  action in cartilage.

## **5. The Role of $ERR\alpha$ in Bone Metastasis**

According to the different mechanisms involved, bone metastasis is mainly classified into two types: osteoblastic (bone-forming) metastasis and osteolytic (bone-resorbing) metastasis, which are closely associated with abnormal activity of osteoblasts or osteoclasts, respectively. Considering the essential role of  $ERR\alpha$  in cancer metastasis and osteocyte differentiation, the role of  $ERR\alpha$  in bone metastasis was also investigated. Similar to the complicated action of  $ERR\alpha$  in bone metabolism, its role in bone metastasis was also controversial. Although the studies regarding the regulation of  $ERR\alpha$  in bone metastasis were mainly produced from the same research group, the conclusion appears to be evolutionary and inconsistent. Initially,  $ERR\alpha$  was shown to enhance the growth in situ but repress the proliferation of breast cancer cells in bone via upregulating the osteoclastogenesis inhibitor OPG and angiogenic growth factor VEGF in a mouse xenograft model of metastatic human breast cancer. A primary breast tumor with overexpression of  $ERR\alpha$  exhibited highly vascularized and increased growth ability, while the receptor inhibited the differentiation and activity of osteoclasts, and further limited the tumorigenesis in bone [57]. However, the study conducted in castration-resistant prostate cancer (CRPC), which had higher aggression and frequently occurring bone metastasis, obtained a distinctive conclusion: they found that the overexpression of  $ERR\alpha$  significantly increased the progression of CRPC and metastasis in bone, and mechanism investigation revealed that the receptor enhanced the bone remodeling via increasing the expression of metastatic factors (such as VEGF-A, WNT5A, TGF $\beta$ 1, and periostin) and generating a favoring-growth tumor environment [58]. In addition, Vargas et al., also found that  $ERR\alpha$  promoted orthotopic breast cancer cell dissemination to bone but not lung metastasis in a mouse breast cancer model through increasing the expression of RANK, which acted as the receptor of RANKL and guided the cancer cells' migration into the bone microenvironment. In line with this, RANKL stimulation induced a higher activity of mTOR signaling in 4T1 cells with  $ERR\alpha$  overexpression, and pharmaceutical inhibition of  $ERR\alpha$  by C29 complex inhibited the tumorigenesis and bone metastatic events [59]. Interestingly, Bouchet et al., subsequently revealed that  $ERR\alpha$  inhibited the growth of breast cancer cells after tumor cell anchorage in the bone via activating the immune response in bone microenvironment. This phenomenon could be achieved via  $ERR\alpha$ -mediated induction of chemokines (CCL17 and CCL20) and reduction of TGF $\beta$ 3, which recruited the CD8 positive T cells to trigger immune cytotoxic action and metastasis repression [60]. The reason that  $ERR\alpha$  appears to function inconsistently or reversely in different cancer types or stages of bone metastasis is a critical issue to be explored. One alternative probability is that prostate cancer predominantly prefers osteoblastic lesions, and the metastasis from breast cancer mostly exhibits osteolytic lesions [61], which may explain the diverse role of  $ERR\alpha$  between prostate cancer and breast cancer. With regard to the different function of  $ERR\alpha$  between dissemination of tumor cells to bone and the growth of metastases, the distinguished microenvironment and signaling stimulation may account for these results [62][63]. Therefore, an appropriate metastatic model and comprehensive analysis of regulatory network will illustrate the important effect of  $ERR\alpha$  on the initiation and progression of bone metastasis.

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## **References**

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