

Dual Function of ERR α in Breast Cancer and Bone Metastasis Formation: Implication of VEGF and Osteoprotegerin

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Abstract

Bone metastasis is a complication occurring in up to 70% of advanced breast cancer patients. The estrogen receptor-related receptor alpha (ERR α) has been implicated in breast cancer and bone development, prompting us to examine whether ERR α may function in promoting the osteolytic growth of breast cancer cells in bone. In a mouse xenograft model of metastatic human breast cancer, overexpression of wild-type ERR α reduced metastasis, whereas overexpression of a dominant negative mutant promoted metastasis. Osteoclasts were directly affected and ERR α upregulated the osteoclastogenesis inhibitor, osteoprotegerin (OPG), providing a direct mechanistic basis for understanding how ERR α reduced breast cancer cell growth in bone. In contrast, ERR α overexpression increased breast cancer cell growth in the mammary gland. ERR α -overexpressing primary tumors were highly vascularized, consistent with an observed upregulation of angiogenic growth factor, the VEGF. In support of these findings, we documented that elevated expression of ERR α mRNA in breast carcinomas was associated with high expression of OPG and VEGF and with disease progression. In conclusion, our results show that ERR α plays a dual role in breast cancer progression in promoting the local growth of tumor cells, but decreasing metastatic growth of osteolytic lesions in bone. *Cancer Res*; 71(17); 5728–38.   2011 AACR.

Introduction

Bone metastasis is a frequent complication of cancer, occurring in up to 70% of patients with advanced breast cancer. Bone metastasis is not a direct cause of death but is associated with significant morbidity (1). For cancer cells to grow in bone, malignant cells recruit and activate osteoclasts (OC; bone resorbing cells) to resorb the bone matrix. Indeed, osteolytic breast cancer metastasis are characterized by an increase in OC number and activity at the bone metastatic site, where excessive bone destruction provides a permissive microenvironment for breast cancer cells to proliferate and expand (2, 3). Unfortunately, current treatments for bone metastasis that rely on antiresorptive agents are only palliative, raising the need for a better understanding of the

molecular mechanisms involved in this pathology so as to design potential alternative therapies (3, 4).

Nuclear steroid receptors are transcription factors that comprise both ligand-dependent molecules such as estrogen receptors (ER) and a large number of so-called orphan receptors, for which no ligand has yet been determined (5). Three orphan receptors, estrogen receptor-related receptor alpha (ERR α), ERR β , and ERR γ , share structural similarities with ER α and ER β (5), but they do not bind estrogen (6, 7). Sequence alignment of ERR α and the ERs reveals a high similarity (68%) in the 66 amino acids of the DNA binding domain, but only a moderate similarity (36%) in the ligand-binding domain, which may explain the fact that ERR α recognizes the same DNA binding elements as ERs but does not bind estrogen (8). Although ERR α activity is decreased by the synthetic molecule XCT790, no natural ligand has yet been found (9–11).

ERR α is known to regulate fatty acid oxidation and the adaptive bioenergetic response (12, 13). It is widely expressed in normal tissues, but several RNA expression studies show its presence in a range of cancerous cells including breast, prostate, endometrial, colorectal, and ovarian tumor tissues (14–20). ERR α was markedly increased in neoplastic versus normal tissues, and ERR α -positive tumors were associated with more invasive disease and higher risk of recurrences (14, 15). On the contrary, ER α and ER β were significantly lower in neoplastic versus normal tissue and were associated with better prognosis (14, 17, 18). ERR α is also highly expressed in skeletal tissues (21, 22) and has been reported to regulate

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Note: Supplementary data for this article are available at Cancer Research Online (<http://cancerres.aacrjournals.org/>).

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doi: 10.1158/0008-5472.CAN-11-1431

  2011 American Association for Cancer Research.

osteoblast, OC differentiation, and bone formation *in vitro* (21, 22, 23, 24) and *in vivo* (25–27). Consistent with these observations, osteopontin (OPN) has been reported to be a direct target gene of ERR α in osteoblastic cell lines (28–30). The role of ERR α in bone metastasis formation is currently unknown.

In the light of these findings, we asked here whether ERR α is involved in breast cancer bone metastasis formation and progression, and whether modulating its activity abrogates bone destruction.

Materials and Methods

Ethics statement

BALB/c and NMRI mice were purchased from Charles River laboratories. All procedures involving animals, including housing and care, the method by which they were killed, and experimental protocols, were conducted in accordance with a code of practice established by the local ethical committee (CREEA: comite Regionale d'Ethique pour l'Experimentation Animale). Studies involving human primary breast tumors were carried out according to the principles embodied in the Declaration of Helsinki. Patients were included anonymously in this study. All human experiments were approved by the Experimental Review Board from the Laennec School of Medicine.

Breast cancer tissue specimens

The autopsy files of the Department of Pathology (Pr. J. Boniver, Centre Hospitalier Universitaire of Liège, Belgium) were searched for diagnosis of disseminated breast cancer with histologically proven bone metastasis during the period 1991 to 1998. Slides were retrieved, and clinical history was obtained. Two breast cancer patients who died with disseminated disease, including bone metastasis, were selected for immunohistochemistry. Soft tissue metastasis (TM) was fixed with formalin, dehydrated, and paraffin embedded.

Breast cancer cohort of patients

In the cohort, patients ($n = 251$) were selected according to the following criteria: primary breast tumor without inflammatory features and no previous treatment (31). Breast cancer tissue biopsies were obtained by surgery, selected by the pathologist, and immediately stored in liquid nitrogen until processing. The biopsies were pulverized using a MikroDis-membrator (B.Braun Biotech International), and total RNA was extracted using TRI Reagent (Sigma). RNA quality was verified using an Agilent Bioanalyser 2100 (Agilent Technologies). Real-time reverse transcriptase PCR (RT-PCR) was carried out.

Cell lines and transfection

MDA-BO2-FRT (BO2) cells and stably transfected clonal derivatives were cultured in complete Dulbecco's modified Eagle's medium (Invitrogen), 10% FBS (Perbio), and 1% penicillin/streptomycin (Invitrogen) at 37°C in a 5% CO₂ incubator. Characteristics of MDA-MB-231/BO2-FRT (BO2) breast cancer cells were previously described (32). To avoid potential effects of different insertion sites, a pcDNA5/FRT vector (Invitrogen) was used to obtain the stable BO2-ERR α WT, BO2-ERR α Δ AF2, and BO2 (CT) cell lines. Human ERR α cDNA [wild type (WT) and

Δ AF2-AD] was obtained from mRNA extracted from BO2-FRT cells, by using RT-PCR with specific primers [(NM_004451.3): ERR α upstream (177bp): GGG AAG CTT AGC GCC ATG TCC AGC CAG; ERR α downstream (WT; 177-1461 bp): GGG GGA TCC CCA CCC CTT GCC TCA GTC C; ERR α downstream (Δ AF2-AD): GGG GGA TCC TCA TGT CTG GCG GAG GAG (177-1350 bp; helix11-12 deletion (32 amino acids)]. Amplimers were sequenced for verification. The pcDNA5/FRT/ERR α -WT and pcDNA5/FRT/ERR α - Δ AF2-AD constructs were cotransfected with the plasmid POG44 (Invitrogen) conferring the specific integration into the FRT site present in the BO2 cells. For clonal selection, cells were cultured for 4 weeks in the presence of hygromycin (20 mg/mL; Invitrogen). Conditioned medium from all clones and from BO2 treated with the inverse agonist XCT-790 at 5.10⁻⁷ mol/L (Sigma) was obtained after 48 hours in α -MEM supplemented with 0.5% of serum, then filter sterilized and proteins quantified to use equal concentration of proteins for each condition (25 μ g).

Animal studies

Tumor fat pad experiments were carried out using BO2-ERR α WT-1, BO2-ERR α Δ AF2 (pool of AF2-1, -2, and -3 clones), and BO2 (CT1/2) cell lines (10⁶ cells in 50 μ L of PBS) injected into the fat pad of the fourth mammary gland of female 4-week-old NMRI nude mice (Charles River). Tumor progression was followed by bioluminescence (NightOwl, Berthold), then tumor size and weight were determined after sacrifice at 66 days.

Bone metastasis experiments using the same pool of clones were carried out in 4-week-old BALB/c nude mice as previously described (33). Cells were suspended at a density of 5 \times 10⁵ in 100 μ L of PBS and inoculated intravenously into animals. Radiographs (LifeRay HM Plus, Ferrania) of animals were taken at 35 days after inoculation using X-ray (MX-20; Faxitron X-ray Corporation). Animals were sacrificed; hind limbs were collected for histology and histomorphometrics analyses. Tibiae were scanned using microcomputed tomography (Skyscan1076, Skyscan) with an 8.8 voxel size, and three-dimensional (3D) reconstructions were carried out with a dedicated visualization software (Amira 5.2, Visage Imaging Inc.). The area of osteolytic lesions was measured using the computerized image analysis system MorphoExpert (Exploranova). The extent of bone destruction for each animal was expressed in square millimeter.

Bone histomorphometry and histology

Hind limbs from animals were fixed and embedded in paraffin. Five millimeter sections were stained with Goldner's Trichrome and processed for histomorphometric analyses to calculate the bone volume/tissue volume (BV/TV) ratio and the tumor burden/tissue volume (TB/TV) ratio. The *in situ* detection of OC was carried out on sections of bone tissue with metastasis using the tartaric-resistant acid phosphatase (TRAP) activity Kit assay (Sigma). The resorption surface (Oc.S/BS) was calculated as the ratio of TRAP-positive trabecular bone surface (Oc.S) to the total bone surface (BS) using the computerized image analysis system MorphoExpert (Exploranova).

Osteoclastogenesis assay

Bone marrow cells from 6-week-old OF1 male mice were cultured for 7 days in differentiation medium: α -MEM medium containing 10% fetal calf serum (Invitrogen), 20 ng/mL of macrophage colony-stimulating factors (M-CSF; R&D Systems), and 200 ng/mL of soluble recombinant receptor activator of nuclear factor κ B ligand (RANKL; ref. 34). Cells were continuously (day 1–7) exposed to conditioned medium extracted (25 μ g proteins for each conditions) from BO2 clones. After 7 days, mature multinucleated OC were stained for TRAP activity (Sigma-Aldrich) and counted as OC when containing 3 or more nuclei.

Immunofluorescence

BO2 cultures were fixed in culture wells with 3.7% paraformaldehyde (Sigma) in PBS for 10 minutes and permeabilized with 0.2% Triton X-100 in PBS. Immunodetection was carried out using a goat polyclonal antibody against human ERR α (Santa Cruz, Tebu) and the secondary antibody (fluorescein isothiocyanate-conjugated donkey anti-goat; Rockland, Tebu-bio). The distribution of F-actin was visualized using phalloidin (Molecular Probes; ref. 14). Cells were observed using an LMS510 laser scanning confocal microscope (Zeiss) with a 63 \times (numerical aperture 1.4) Plan Neo Fluor objective.

Immunoblotting

Cell proteins were extracted, separated in 4% to 12% SDS-PAGE (Invitrogen), then transferred to nitrocellulose membranes (Millipore) using a semidry system. For ERR α and α -tubulin detection, the same goat polyclonal antibody ERR α (Santa Cruz) and a mouse polyclonal antibody against human α -tubulin (Sigma-Aldrich) were used. Membrane was incubated with secondary antibody horseradish peroxidase (HRP)-conjugated donkey anti-goat (Santa Cruz) and anti-mouse (Amersham), respectively. An ECL kit (PerkinElmer) was used for detection.

Immunocytochemistry

Five micrometer sections were subjected to immunohistochemistry using the same goat polyclonal antibody ERR α (Santa Cruz) and a rabbit polyclonal antibody against human osteoprotegerin (OPG; Abbiotec). Sections were incubated with secondary antibody HRP-conjugated donkey anti-goat and anti-rabbit, respectively (Amersham; dilution 1/300) for 1 hour. After washing, the sections were revealed by 3,3'-diaminobenzidine (Dako).

Real-time RT-PCR

Total RNA was extracted with Trizol reagent (Sigma) from cancer cells and OCs. Real-time RT-PCR was carried out on a

Table 1. Clinical and biological characteristics and ERR α mRNA expression in breast cancer patients

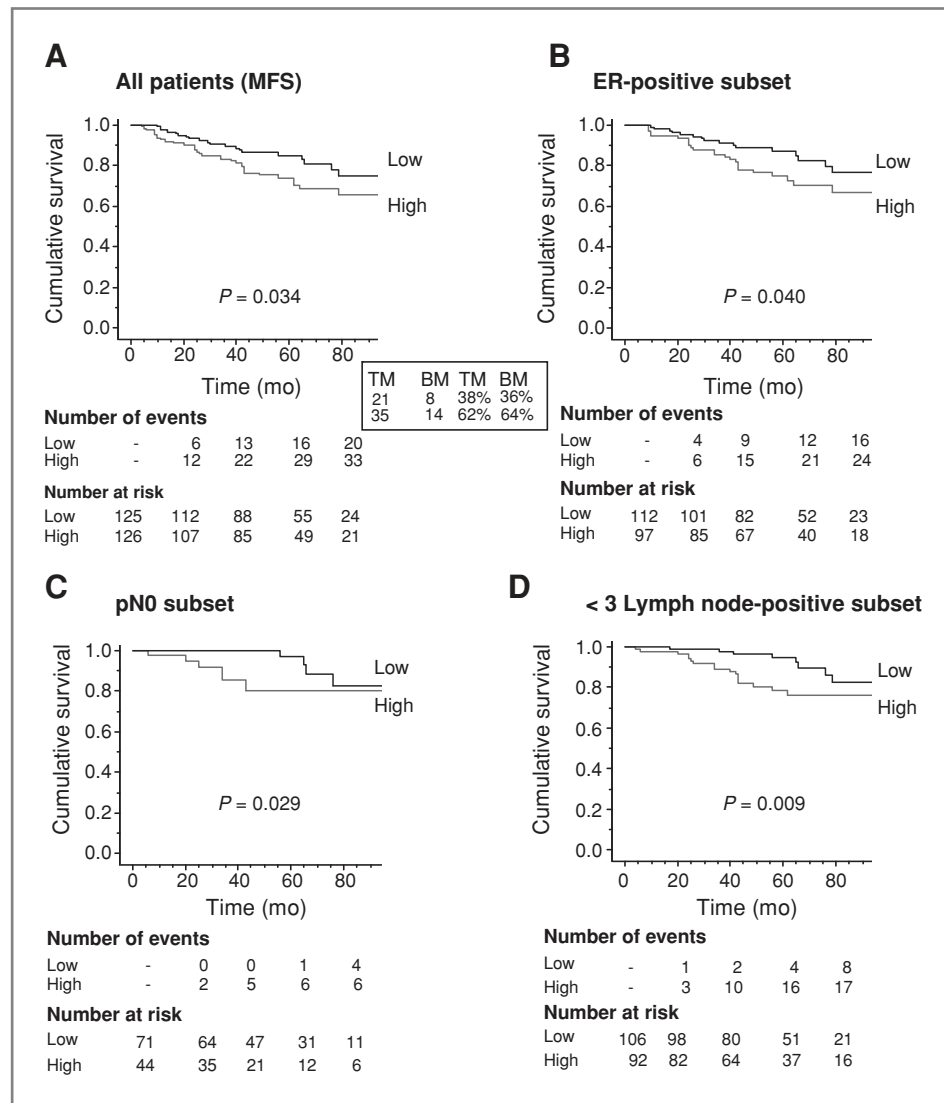
Characteristics		All patients N = 251			pN0 patients N = 115			pN+ patients N = 136		
		n	Median	P	n	Median	P	n	Median	P
Menopausal status	Pre	105	2.68	0.425	48	2.59	0.020	79	3.10	0.229
	Post	146	2.59		67	2.04		57	2.72	
Surgical tumor size	<20 mm	101	2.53	0.256	58	2.15	0.662	43	2.89	0.967
	\geq 20 mm	142	2.69		55	2.46		87	2.95	
	ND	8								
Histologic type	Ductal	205	2.68	0.026	91	2.32	0.208	114	3.03	0.058
	Lobular	37	2.09		16	1.71		21	2.21	
	Others	9								
Histologic grade ^a	1	27	2.32	0.341	14	2.22	0.718	13	2.37	0.040
	2	107	2.72		53	2.41		54	3.18	
	3	58	2.84		22	2.16		36	3.12	
	ND	13								
Node status	Negative	115	2.31	<0.001			0.439	83	2.72	0.439
	1–3	83	2.72					53	3.12	
	>3	53	3.12							
RE status	Negative	42	3.17	<0.001	16	3.47	0.003	26	3.17	0.106
	Positive	209	2.53		99	2.12		110	2.78	
VEGF status ^b	Low	92	2.35	0.002	38	1.99	0.018	54	2.62	0.010
	High	93	3.08		42	2.57		51	3.23	
	ND	66								

NOTE: P values correspond to Mann–Whitney test or Kruskal–Wallis test (histologic grade and node status).

^aHistologic grade defined only in ductal carcinomas.

^bLow: <50% quartile, high: \geq 50% quartile.

Figure 1. *ERR* α is a bad prognostic marker. Kaplan–Meier curves show correlation between high expression of *ERR* α , categorized with median value, and MFS in patients in (A) the whole population ($N = 251$), (B) the ER-positive subset ($N = 209$), (C and D) the pN0 subset patients ($N = 115$) and the <3 lymph node-positive subset ($N = 198$). Low: $\leq 50\%$ quartile; high: $\geq 50\%$.



Roche Lightcycler Module (Roche) with specific primers (see Supplementary Table S2). Real-time RT-PCR was carried out by using SYBR Green (Qiagen) on the LightCycler system (Roche) according to the manufacturer's instructions. Amplifiers were all normalized to corresponding L32 values. Data analysis was carried out using the comparative C_t method.

Real-time RT-PCR on breast cancer tissue biopsy mRNA was carried out using primers specific for human *L32* (101 bp): 5'-CAAGGAGCTGGAAGTGTGCTGC-3', 5'-CAGCTCTTCCACGATGGCT-3'; TATA-box binding protein (*TBP*; 138 bp) 5'-TGGTGTGCACAGGAGCAAG-3', 5'-TTCACATCACAGTCCCCAC-3'; *ERR* α (101 bp): 5'-ACCGAGAGATTGTGGT-CACCA-3', 5'-CATCCACACGCTCTGCAGTACT-3' and *OPG* (Supplementary Table S2) and SYBR green (Invitrogen) in 96-well plates on a Mastercycler EP system (Realplex2, Eppendorf) according to the manufacturer's instructions with an initial step for 10 minutes at 95°C followed by 40 cycles of 20 seconds at 95°C, 15 seconds at T_m (*L32*: 62°C, *TBP*: 67°C,

ERR α : 59°C), and 10 seconds at 72°C. *ERR* α and *OPG* expression were normalized with the average of the genes expression encoding the ribosomal protein *L32* and the *TBP*.

Cell invasion assay

Invasion assays were carried out using Bio-Coat migration chambers (Becton Dickinson) with 8 μ m filters coated with Matrigel as described previously (35). BO2 cells (5×10^4) were plated in the upper chambers and the chemoattractant (10% FBS) in the lower chambers. After 24 hours at 37°C in 5% CO₂ incubator, cells that had migrated through the filters were fixed and stained. Cells were counted (200 \times magnification). All experiments were run in triplicate, and invasion was expressed in cells/square millimeter.

OPG ELISA

Conditioned medium obtained from BO2-CT(1/2), BO2-ERR α -WT-1, and BO2-FRT-ERR α -AAF2 (pool of AF2-1, -2

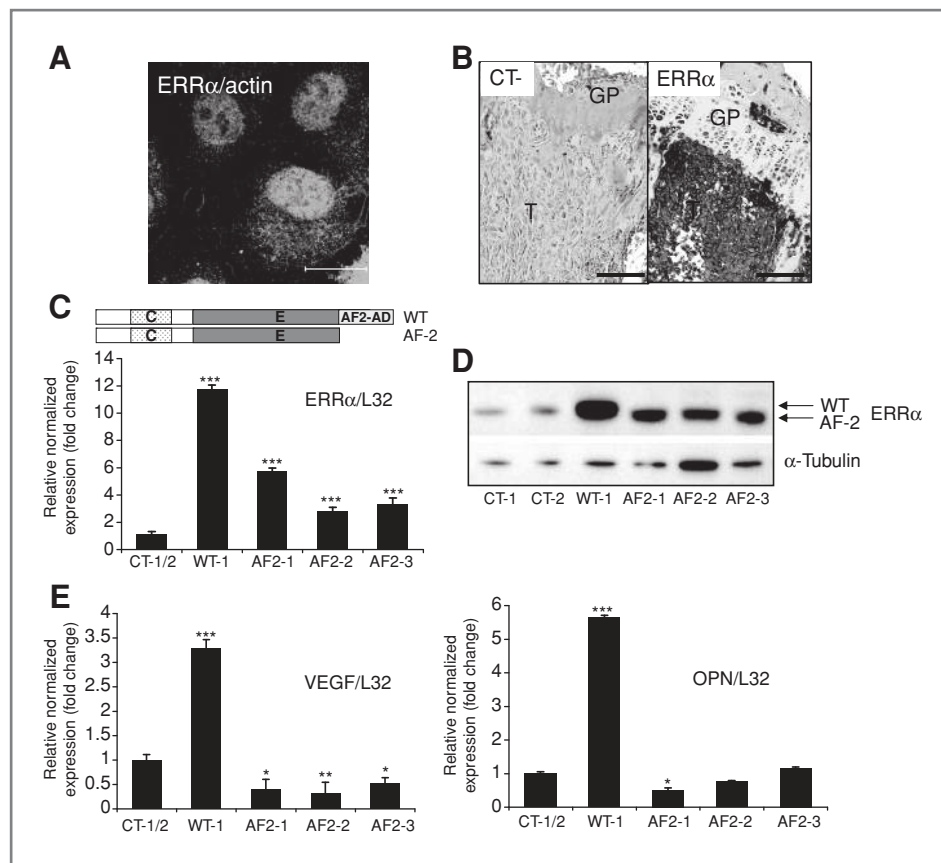


Figure 2. Modulation of *ERRα* in BO2 breast cancer cell line. **A**, *ERRα* protein expression (nucleus and cytoplasm) in BO2 cells by immunofluorescence and confocal microscopy and **(B)** *in vivo* by immunohistochemistry in bone metastasis present 30 days after intravenous injection of BO2 cells. **C**, isolation after stable transfection of three independent BO2-*ERRαΔAF2* clones (*ERRα* dominant-negative form), one clone BO2-*ERRαWT* and two controls (CT-1 and CT-2) BO2-CT (empty vector). *ERRα* expression was assessed by real-time PCR on triplicate samples and normalized against that of the ribosomal protein gene *L32* (ANOVA, $P < 0.0001$) and **(D)** by Western blotting. **(E)** *VEGF* and *OPN* expression was increased in BO2-*ERRαWT* and decreased or not regulated in BO2-*ERRαΔAF2* (ANOVA, $P < 0.0001$ for *VEGF* and *OPN* in WT-1 or $\Delta AF2$ versus CT). **(A)** bar = 20 μm and **(B)** bar = 200 μm . T, tumor; GP, growth plate.

and -3 clones) were diluted following the manufacturer's instructions, and OPG concentration was evaluated using the ELISA Kit (RayBiotech).

Statistical analysis

Data were analyzed statistically by one-way ANOVA followed by post hoc *t*-tests to assess the differences between groups for *in vitro* and *in vivo* studies. Concerning the cohort, the median follow-up at the time of analysis was 54 months. The criterion for statistical analyses was the metastasis free survival (MFS), that is, the delay between the time of primary surgery and the first event: nodal or distant metastasis or death. Analysis of the distribution of *ERRα* expression in relation to the usual prognostic parameters was carried out using the Mann-Whitney or Kruskal-Wallis test. Survival probabilities were estimated using Kaplan-Meier estimators and were compared using the log-rank test. Univariate analysis was carried out using the Cox proportional hazard model. Results of $P < 0.05$ were considered significant.

Results and Discussion

ERRα mRNA and protein expression in human primary breast tumors and bone metastasis

We analyzed *ERRα* mRNA expression by real-time RT-PCR in a cohort of 251 breast tumor biopsies (Supplementary

Table S1; ref. 31). As reported previously by others (14, 15, 17, 18), a statistically significant association was detected in all patients analyzed between *ERRα* expression and histologic type, node status, and ERs (radioligand method; $P = 0.026$, $P < 0.001$, $P < 0.001$; Table 1). The Kaplan-Meier curve was constructed after segmentation into 2 groups on the basis of the median value for *ERRα* expression (Fig. 1A-D). It was observed that high levels of *ERRα* mRNA expression were related to a decrease in MFS ($N = 251$, $P = 0.034$; Fig. 1A). Sixty-two percent of patients (35/56) with high *ERRα* expression levels exhibited liver, lung, bone, and soft tissue metastasis compared with 38% of patients (21/56) having low *ERRα* levels (Fig. 1A, see frame). This paralleled the frequencies seen in patients ($n = 22$) who had developed "only" bone metastasis (BM), that is, 64% (high *ERRα*) and 36% (low *ERRα*; Fig. 1A) suggesting that *ERRα* is an overall bad prognostic factor that is not a determinant of metastasis location of breast cancer cells. Moreover, high *ERRα* expression correlated with a higher risk of recurrence at an early stage of the disease in the ER-positive group ($N = 209$), the pN0 subset, and in the pN < 3 lymph-node-positive subset ($P = 0.04$; $P = 0.029$, and $P = 0.009$; log-rank test), when compared with low *ERRα* (Fig. 1B-D) suggesting that *ERRα* may be a very useful early prognostic marker in breast cancer. Finally, as previously described (15), *ERRα* protein was present *in situ* and in invasive breast carcinoma cells (Supplementary Fig. S1B and C, respectively)

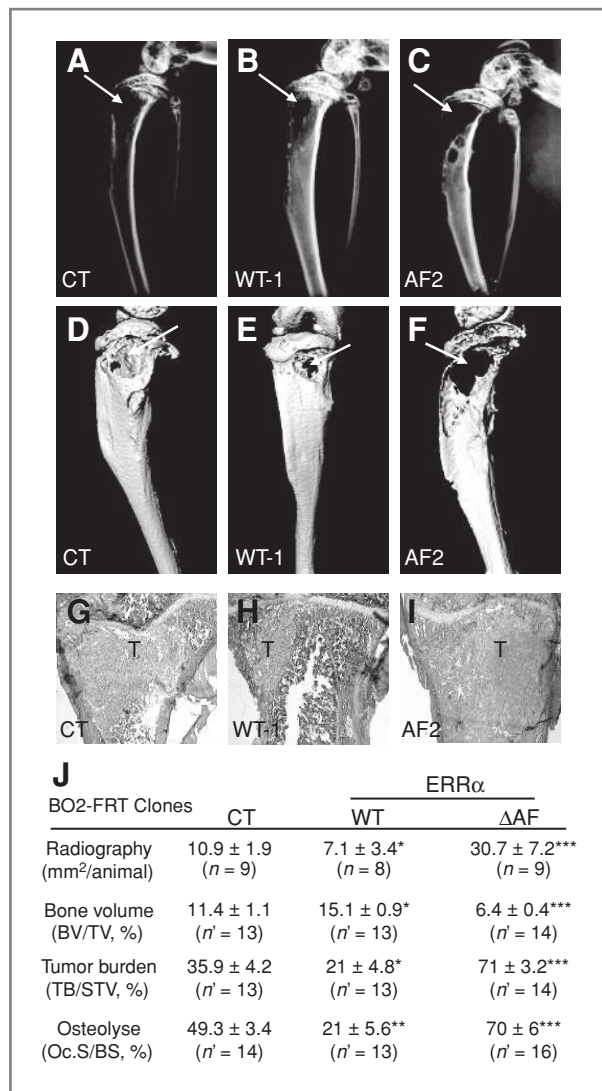


Figure 3. Overexpression of ERR α inhibits development of bone metastasis. A–C, BO2-ERR α WT-1, BO2-ERR α Δ AF2 (pool), or BO2-CT (pool) cells were inoculated into BALB/c nude mice; 35 days postinoculation, radiography revealed smaller osteolytic lesions in mice injected with BO2-ERR α WT-1 cells and much larger lesions in mice injected with BO2-ERR α Δ AF2 cells compared with mice injected with CT cells (see white arrows). D–F, 3D micro-CT reconstructions of tibiae and (G–I) histology after Goldner's Trichrome staining confirmed the radiography results. J, quantification of bone destruction. T, tumor.

but not in normal breast epithelial cells (Supplementary Fig. S1A). ERR α was also clearly present in breast cancer cells that metastasized to bone (Supplementary Fig. S1D see T). As previously reported by us (21), ERR α was also detected in osteocytes embedded in the bone matrix.

ERR α expression in breast cancer cells reduces their ability to induce osteolytic lesions *in vivo*

To assess whether ERR α is involved in bone metastasis formation, we used MDA-BO2-FRT (BO2) cells, a subpopulation of the human MDA-231 breast cancer cell line, that was

selected for the high efficiency with which it metastasizes to bone (32). ERR α protein was seen in the nucleus and cytoplasm of BO2 cells *in vitro* (Fig. 2A) and *in situ* in bone metastasis from legs of animals, 30 days after intravenous tumor cell inoculation (Fig. 2B).

To establish a functional role for ERR α in bone metastasis development, we next transfected BO2 cells with a full-length (WT) ERR α or a truncated version of ERR α lacking the coactivator binding domain AF2, ERR α Δ AF2, which acts as a dominant-negative form (22, 23, 36; Fig. 2C). Constructs of human ERR α WT and ERR α Δ AF2 were stably transfected into the genomic FRT site present in the BO2 cells. Three independent BO2-ERR α Δ AF2 (1, 2, 3), one BO2-ERR α WT, and two BO2-CT (empty vector) clones were obtained, named AF2-1, AF2-2, AF2-3, WT-1, CT-1, and CT-2, respectively. As judged by real-time PCR, total ERR α mRNA expression was increased when compared with CT-1/2 clones (Fig. 2C). Western blotting detected a band of approximately 50 kD for ERR α protein in CT1-2 and WT-1 that was increased in WT-1 and AF2-1, AF2-2, and AF2-3 cells. The presence of a band with a slightly lower molecular weight in AF2-1, AF2-2, and AF2-3 cells corresponded well with the expected size for truncation of the AF2 domain (42 amino acids; Fig. 2D). mRNA expression levels of the ERR α target genes *VEGF* and *OPN* were statistically significantly increased in WT-1 cells compared with CT-1/2 cells (Fig. 2E). By contrast, *VEGF* and *OPN* mRNA levels remained reduced or unchanged in AF2 clones (Fig. 2E), confirming the increased activity and the dominant-negative functions of the WT and truncated ERR α Δ AF2 constructs, respectively.

To assess the involvement of ERR α in bone metastasis formation, CT (pool of CT-1 and -2 clones), WT-1, and AF2 (pool of AF2-1, -2 and -3 clones) cells were inoculated intravenously into female BALB/c nude mice. Thirty-five days after tumor cell injection, radiographic analysis revealed that animals bearing WT-1 tumors had osteolytic lesions that were 40% smaller than those of mice bearing CT tumors (Fig. 3A,B, and J). By contrast, there was a 3-fold increase in the extent of osteolytic lesions in animals bearing AF2 tumors, when compared with control (Fig. 3A, C, and J). The inhibitory effect of ERR α on cancer-induced bone destruction was confirmed using 3D micro-CT reconstruction (Fig. 3D–F), histology (Fig. 3G–I), and histomorphometric analyses of tibiae (BV/TV; skeletal tumor burden, TB/STV; Fig. 3J). Taken together, our results indicated that overexpression of ERR α in breast cancer cells reduced the formation of osteolytic lesions.

Regulation of OC formation by ERR α -expressing BO2 cells

Given these data, we next asked whether modulation of ERR α in breast cancer cells could alter OCs, the bone resorbing cells. TRAP staining of tibial sections of metastatic legs from animals bearing WT-1 and AF2 tumors showed a 43% decrease and a 143% increase of TRAP-positive OC surface (Oc.S/BS) at the bone/tumor cell interface, respectively, when compared with CT tumors (Fig. 4A and 3J; Supplementary Fig. S2). Consistent with these *in vivo* data, the treatment of primary mouse bone marrow cell

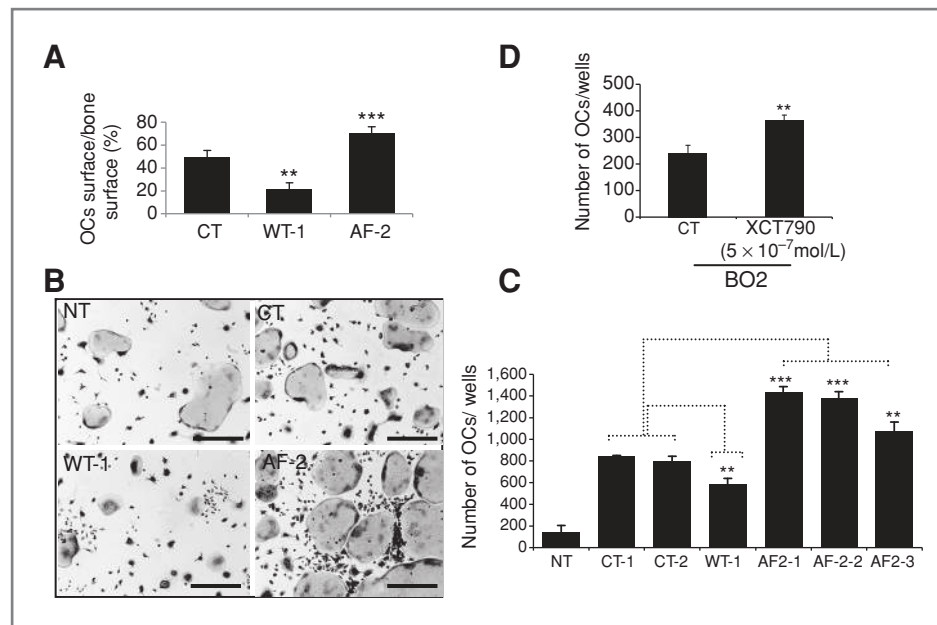


Figure 4. *ERRα* expression in BO2 cells regulates OCs formation. A, TRAP staining of OCs in sections of tibiae taken from mice injected with BO2-*ERRα*WT-1, BO2-*ERRα*ΔAF2 (pool), or BO2-CT (pool) cells shows decreased and increased surface of OCs in BO2-*ERRα*WT-1 and BO2-*ERRα*ΔAF2, respectively compared with CT (ANOVA, $P < 0.0001$). B and C, primary mouse bone marrow cells were cultured in the presence of RANKL and M-CSF and treated or not with medium conditioned by BO2-*ERRα*WT-1, BO2-*ERRα*ΔAF2, or BO2-CT cells. Fewer OCs formed in cultures treated with BO2-*ERRα*WT-1 conditioned medium, whereas more formed in cultures treated with BO2-*ERRα*ΔAF2 conditioned medium, compared with CT (refs. 1, 2) conditioned medium (ANOVA, $P < 0.0001$). D, conditioned medium obtained from parental BO2 cells treated with the *ERRα* inverse agonist XCT-790 increased OCs formation, mimicking the results obtained with BO2-*ERRα*ΔAF2 conditioned medium (ANOVA, $P < 0.001$). Bar, 100 μ m.

cultures with RANKL and M-CSF together with the conditioned medium of WT-1 cells inhibited the formation of TRAP-positive multinucleated OCs compared with that observed with the conditioned medium of CT cells (Fig. 4B and C). By contrast, the conditioned medium from AF2 cells stimulated OC formation (Fig. 4B and C). In addition, the conditioned medium from parental BO2 cells treated with the inverse agonist XCT-790, which blocks *ERRα* activity, increased OC formation compared with control (dimethyl sulfoxide; Fig. 4D), confirming our osteoclastogenesis data obtained with the conditioned medium of AF2 cells.

***ERRα* regulates *OPG* expression in breast cancer cells**

We showed that BO2 breast cancer cells overexpressing wild-type *ERRα* markedly inhibited osteolysis *in vivo* (Fig. 3J) and reduced OC formation *in vitro* (Fig. 4). We quantified several markers involved in osteoblasts and OC differentiation, and we found that the *OPG*, a soluble decoy receptor for RANKL that inhibits osteoclastogenesis, was regulated by *ERRα* (Fig. 5A; ref. 37). By immunohistochemistry, we show that *OPG* expression was higher in skeletal WT-1 tumors compared with that observed in AF2 and CT tumors (Fig. 5B). In addition, as judged by ELISA, WT-1 cells secreted higher amounts of *OPG* compared with CT-1/2 and AF2 cells (pool of AF2-1, -2 and -3 clones; Fig. 5C). *OPG* mRNA expression was also quantified by real-time RT-PCR in the cohort of 251 patients. *OPG* levels were statistically significantly higher

in *ERRα*-positive tumors compared with *ERRα*-negative tumors (Fig. 5D; $P = 0.013$). Moreover, there was a positive correlation between high mRNA expression levels of both *ERRα* and *OPG* (*ERRα*⁺/*OPG*⁺) and a decrease in relapse-free survival ($P = 0.028$, log-rank test; Fig. 5E). All together, the significant correlation between high *ERRα* and *OPG* in patients and the regulation of *OPG* by *ERRα* in BO2 cells provide a mechanistic basis for the reduction of osteoclastogenesis *in vitro* and *in vivo*. Interestingly, *OPG* in our preclinical data suggest that, alone it had no prognostic value in breast carcinomas (Fig. 5F) whereas in association with high *ERRα* mRNA levels, a correlation with a poor clinical outcome in patients was found (Fig. 5E). *OPG* is not only an osteoclastogenesis inhibitor, but also a survival factor for human breast cancer cells (38, 39). It also promotes angiogenesis (40), and its overexpression in human MCF-7 breast cancer cells enhances tumor growth following orthotopic inoculation in animals (41). *ERRα* has been implicated in tumor progression, and the positive association between high *ERRα*/*OPG* mRNA levels and increased risk of recurrences in patients (Fig. 5E) suggested that *ERRα* could play a role on primary tumor expansion.

ERRα* stimulates tumor growth and angiogenesis *in vivo

To address this hypothesis, orthotopic tumors were induced with CT (pool of CT-1 and -2 clones), WT-1, or AF2 (pool of AF2-1, -2 and -3 clones) cells upon inoculation within the mammary fat pad of NMRI nude female mice.

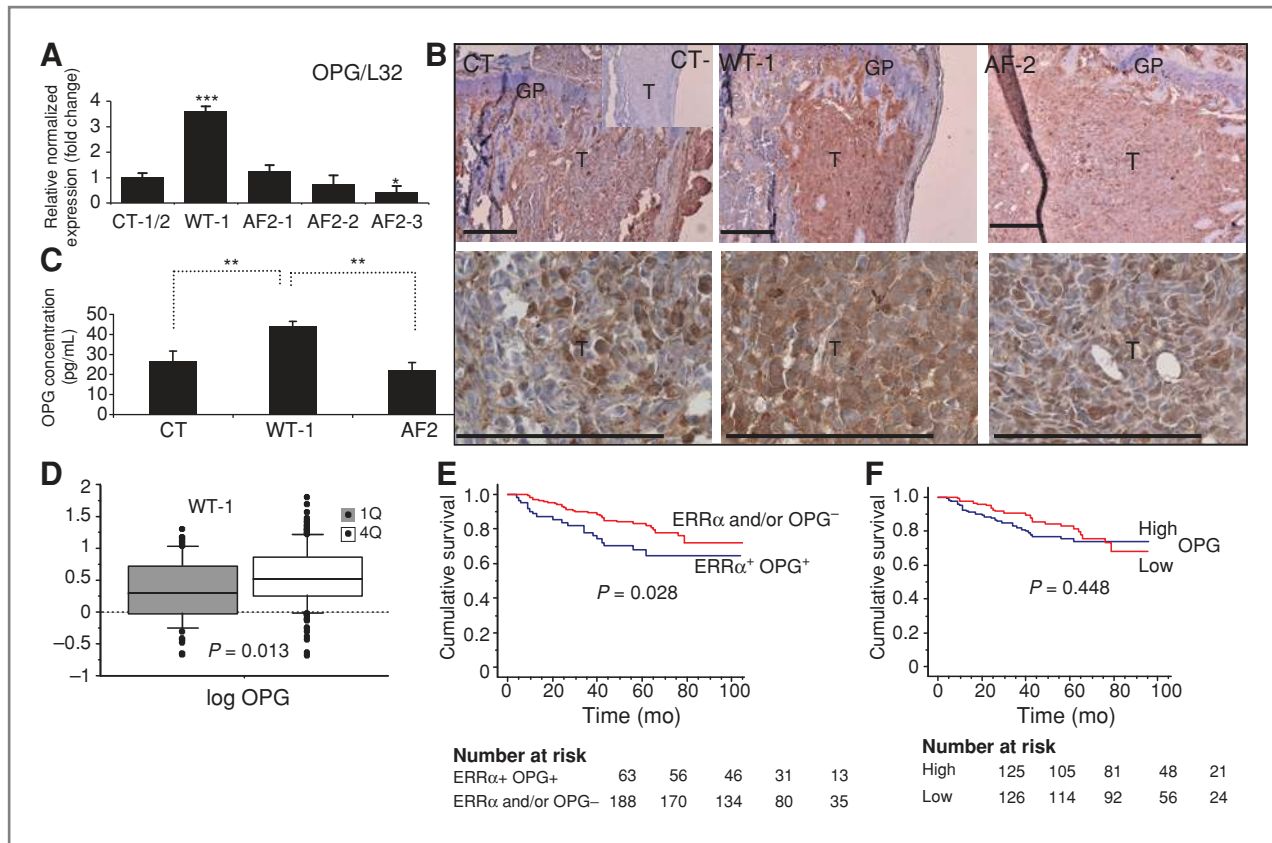


Figure 5. Correlation of ERR α and OPG in BO2 cells and breast cancer patients. **A**, real-time PCR carried out on RNA extracted from BO2 clones showed increased expression of OPG by ERR α (ANOVA, $P < 0.0001$). **B**, staining for OPG is higher in bone metastasis induced by BO2-ERR α WT cells compared with BO2-CT and BO2-ERR α ΔAF2 cells; tissues collected 35 days postcell inoculation. **C**, ELISA quantification confirmed the increased secretion of OPG by BO2-ERR α WT compared with BO2-CT (pool) and BO2-ERR α ΔAF2 (pool) cells (ANOVA, $P = 0.0064$; $P < 0.01$ CT versus WT-1 and WT-1 versus AF-2). **D** and **E**, a significant correlation was also found between levels of ERR α mRNA and median values of OPG mRNA in the cohort (ERR α 1st quartile and median OPG = 2.03; ERR α 2nd–4th quartile and median OPG = 3.45). Kaplan-Meier curves show that ERR α ⁺/OPG⁺ expression was associated with a decrease in MFS. **F**, OPG alone was not associated with MFS.

Bioluminescence analysis from day 5 to day 66 revealed a dramatically greater tumor progression in WT-1 tumor-bearing animals compared with that observed with CT and AF2 tumor-bearing animals a modest increase in AF2 tumor burden was also observed at day 62 and 66 (Fig. 6A and B). Tumor weight/size at day 66 (Fig. 6C and D) correlated well with bioluminescence quantification (Fig. 6B and C). Interestingly, WT-1 tumors were highly vascularized compared with CT and AF2 tumors (Fig. 6E), an observation correlating with higher *VEGF* mRNA levels observed in WT-1 versus AF2 or CT tumors (Fig. 6E). Moreover, if these results are in agreement with previous data describing *VEGF* as a target gene for ERR α in breast cancer (42), we show for the first time a positive association between high levels of ERR α and *VEGF* in breast tumors from patients ($P = 0.002$; Table 1). Interestingly, OPG expression that can be stimulated by VEGF in endothelial cells, is also known to be a positive regulator of microvessel formation *in vivo* (43) and therefore can participate to the neovascularization observed in WT-1 tumors. We also observed that

ERR α promoted BO2 breast cancer cell invasion *in vitro* (Supplementary Fig. S3A) but has a slightly effect on proliferation (data not shown). Consistent with this we found matrix metalloproteinases *MMP1* and *MMP13* regulated by ERR α (Supplementary Fig. S3B). These results were in agreement with previous findings showing that the silencing of ERR α dramatically reduced the *in vitro* migratory capacity of breast cancer cell lines (44). Taken together, these results strongly suggested that ERR α promoted tumor growth, mainly through the stimulation of angiogenesis and invasion. Based on our results on high ERR α /OPG/VEGF in our preclinical study, we propose that OPG worked in concert with VEGF to stimulate tumor angiogenesis which, in turn, promoted the growth of BO2-ERR α WT cells. Conversely, in bone metastasis although the angiogenic factor *VEGF* was overproduced in BO2-ERR α WT cells, tumor-derived VEGF had probably a low impact on progression of osteolytic lesions. Indeed, recent studies have shown that hypoxia was nonessential for bone metastasis while promoting angiogenesis in lung metastasis and primary tumor

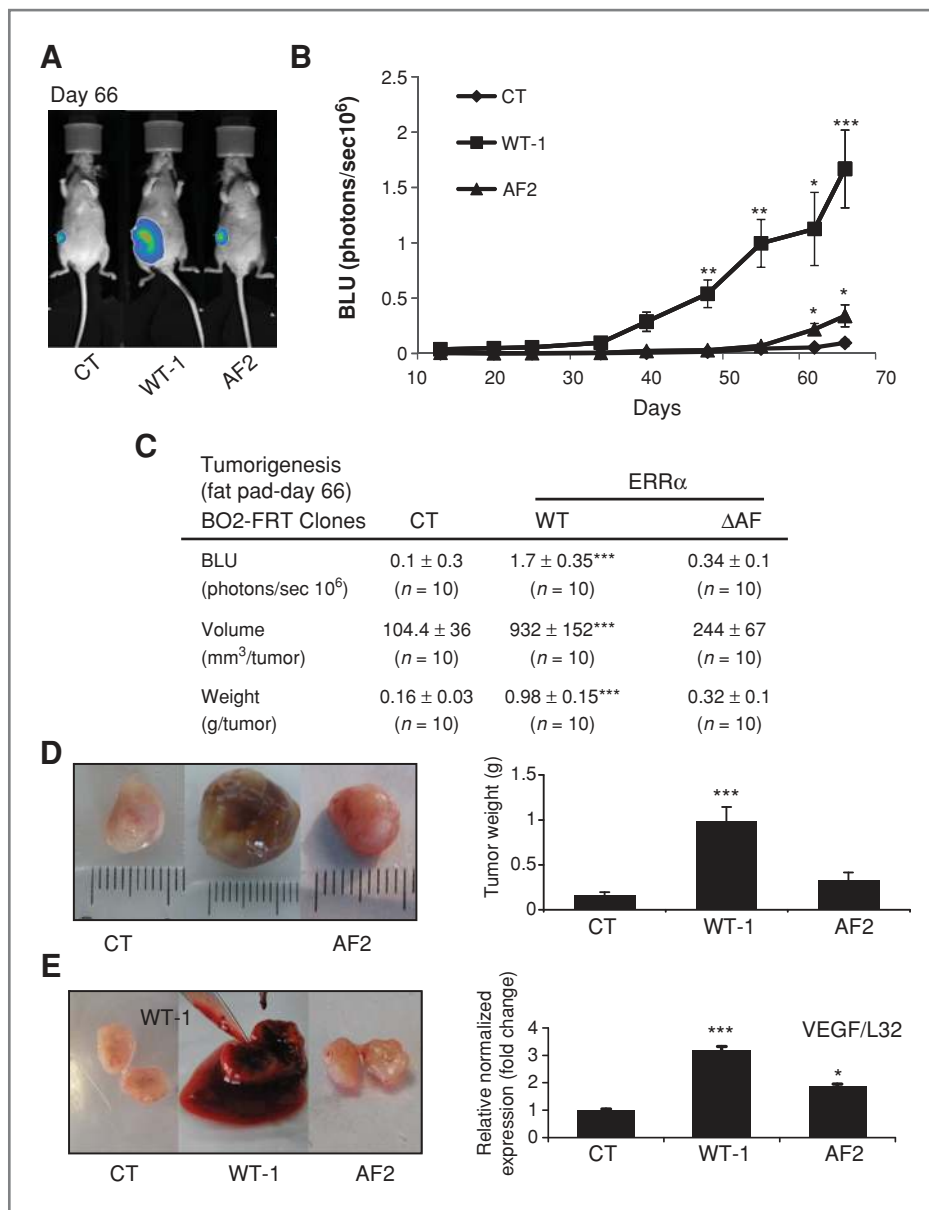


Figure 6. Stimulation of tumor progression and angiogenesis by ERR α *in vivo*. A–C, BO2-ERR α WT, BO2-ERR α Δ AF2 (pool), or BO2-CT (pool) cells were inoculated into the fat pad of NMRI nude mice. Tumor progression was followed by bioluminescence from day 5 to 66. Greater tumor expansion was observed in mice with BO2-ERR α WT-1 compared with BO2-ERR α Δ AF2 (pool) or BO2-CT (pool) cells. C and D, weight and volume of tumors dissected at endpoint (E) and VEGF expression normalized against L32 (ANOVA, $P < 0.001$) within tumors (pool of $n = 3$ for each condition) correlated with greater tumor vascularization.

growth (45). Therefore, modulating angiogenesis through VEGF and the proangiogenic role of OPG may have no impact on angiogenesis in bone, as bone is already extremely vascularized (46), but have dramatic impact on vascularization and progression of primary breast tumors or metastasis to nonbone sites. These data provide novel insights into how ERR α can be a bad prognostic factor in the primary tumor (angiogenesis via VEGF and OPG) but a favorable biomarker in the very special case of bone metastasis (inhibition of OC formation through OPG).

In conclusion, our results show for the first time that ERR α plays a dual role, promoting the progression and invasion of primary tumors, but decreasing osteolytic lesions in bone. In addition, our data show that OPG is modulated by ERR α that

probably contributes to the overall negative clinical outcome which is associated with the expression of ERR α in human breast carcinomas.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

Acknowledgments

We are very grateful to C. Lionnet and C. Chamot from PLATIM (IFR 128 Lyon Biosciences) for their help with imaging experiments and to Blandine Deux, Vincent Gonin, and Pascale Heneaux for their technical help. The authors also thank the CeCIL platform (Faculté de Médecine Laennec, Lyon, France) for technical assistance.

Grant Support

This work was supported by the CNRS (EB), Inserm, the University of Lyon, "Ligue Regionale contre le Cancer" (Isère; E. Bonnelye). A. Fradet is supported by the Ligue Nationale contre le Cancer. B. Depalle by region Rhône Alpes, and A. Bellahcène from the National Fund for Scientific Research, Belgium.

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Received April 29, 2011; revised June 29, 2011; accepted June 29, 2011; published OnlineFirst July 6, 2011.

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Cancer Res 2011;71:5728-5738. Published OnlineFirst July 6, 2011.

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