Loss of ERα induces amoeboid-like migration of breast cancer cells by downregulating vinculin

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Oestrogen receptor alpha (ERα) is a well-known target of endocrine therapy for ERα-positive breast cancer. ERα-negative cells, which are enriched during endocrine therapy, are associated with metastatic relapse. Here we determine that loss of ERα in the invasive front and in lymph node metastasis in human breast cancer is significantly correlated with lymphatic metastasis. Using in vivo and in vitro experiments, we demonstrate that ERα inhibits breast cancer metastasis. Furthermore, we find that ERα is a novel regulator of vinculin expression in breast cancer. Notably, ERα suppresses the amoeboid-like movement of breast cancer cells by upregulating vinculin in 3D matrix, which in turn promotes cell-cell and cell-matrix adhesion and inhibits the formation of amoeboid-like protrusions. A positive association between ERα and vinculin expression is found in human breast cancer tissues. The results show that ERα inhibits breast cancer metastasis and suggest that ERα suppresses cell amoeboid-like movement by upregulating vinculin.
The pathogenesis of breast cancer is associated with oestrogen receptor alpha (ERα), which is activated by sex hormones and contributes to the aberrant proliferation of breast cancer cells. The classical mechanism of ERα action involves regulating the transcription of oestrogen-responsive genes by binding to the oestrogen-responsive element (ERE) within the promoters of the target genes.

Endocrine therapy with selective oestrogen receptor modulators, such as tamoxifen, has been widely used to antagonize ERα in breast cancer tissues. However, tamoxifen appears to decrease the risk of ERα-positive contralateral breast tumours and to increase the risk of ERα-negative contralateral tumours. Therefore, loss of expression of ERα during adjuvant endocrine treatment for ERα-positive breast cancer allows for resistance to common adjuvant endocrine therapies and is associated with ERα-negative metastatic relapse. Nevertheless, how ERα loss is associated with metastasis remains to be elucidated, particularly in a three-dimensional (3D) environment, which can better mimic human breast cancer metastasis in vivo.

The migration of invasive cells in vivo consists of the mesenchymal mode, in which invasive cells are elongated and require pericellular matrix proteolysis, and the amoeboid mode, in which carcinoma cells with low adhesion and round morphology require subcellular localization of myosin II behind the cell nucleus to drive actomyosin contractility, independently of matrix metalloprotease. Moreover, mesenchymal cells usually burst after entering the bloodstream, whereas highly metastatic cancer cells efficiently penetrate the blood vessels through amoeboid-like migration with high actomyosin contractility, which provides cells with mechanical strength to resist shear forces in the circulation. In addition, cells undergoing amoeboid migration have higher velocity than those undergoing mesenchymal migration. Despite this observation, the molecular mechanisms that affect the amoeboid motility in a 3D environment require more in-depth study.

Vinculin (VCL), a membrane cytoskeletal protein found in focal adhesion plaques, is involved in the linkage of the ECM to the actin cytoskeleton. Vinculin-null cells transfected with vinculin cDNA show markedly decreased motility and tumorigenicity. In addition, downregulation of vinculin was found in several highly metastatic cancer cells. Loss of vinculin induced protection from apoptosis in an anchorage-independent manner and enhanced cell motility. However, few studies have described the involvement of vinculin in the regulation of breast cancer amoeboid movement.

In this study, we show that loss of ERα promotes tumour metastasis through in vitro experiments, in vivo tumour xenograft assays and the analysis of clinical breast cancer samples. Furthermore, we find that ERα is a novel regulator of vinculin expression in breast cancer and that loss of ERα induces amoeboid-like migration of breast cancer cells by regulating vinculin in a 3D matrix.

Results

Loss of ERα is correlated with breast cancer metastasis. Metastasis occurs when tumour cells detach from their primary location and move to the lymph nodes and then to distant organs. Therefore, the lymph node usually serves as a bridge allowing the metastatic dissemination of tumours. We examined ERα expression in human primary breast cancer tissues and the corresponding lymph node metastasis from 124 ERα-positive breast cancer patients. This analysis demonstrated that, in contrast to the abundant expression of ERα in the primary tumour, 54.8% of samples lost the expression of ERα in the corresponding lymphatic metastasis. In addition, loss of ERα expression in lymphatic metastasis was also positively associated with the clinical stage and the number of lymph node metastases and the loss of progesterone receptor expression. However, there was no significant association with age, tumour size or HER2 expression.

We further examined the expression of ERα at the invasive front where the infiltration of CD68-positive tumour-associated macrophages is found and where at least 50% of the cell surface of tumour cells contacts the matrix and the non-invasive front of the primary tumour tissues. The results showed that ERα was minimally expressed in breast cancer cells at the invasive front, whereas an increased intensity of ERα staining was observed at the non-invasive front. Collectively, these findings suggest that decreased ERα expression might promote breast cancer metastasis.

ERα inhibits breast cancer metastasis in vivo and in vitro. To investigate whether ERα might inhibit the metastasis of breast cancer cells, we first chose ERα-positive MCF-7 and Cas9-ERα MCF-7 cells to examine the lung metastasis in a tail vein injection model, which mimics the process of loss of ERα during metastasis in patients. The expression of ERα in the two cell lines was detected by western blot analysis. The cell proliferation assay confirmed that the Cas9-ERα MCF-7 cells grew more slowly in vitro. Biofluorescence was examined at different time points to monitor the location and growth of tumour xenografts in the lungs. We found that the normalized photon flux of Cas9-ERα MCF-7 cells, which represents the pulmonary metastasis focuses, was significantly higher than that of control cells, as evidenced by H&E staining. Therefore, the Cas9-ERα MCF-7 cells showed a more profound increase in metastatic potential to the lungs.

Furthermore, control MDA-MB-231 cells and ERα-expressing MDA-MB-231 cells were also used to examine the metastasis in an orthotopic injection model. The expression levels of ERα in these two cell lines were quantified in vitro. The normalized photon flux of control cells was lower than that of ERα-expressing cells. The cell proliferation assay also verified that the control cells grew more slowly in vitro. To observe the tumour metastasis, we covered the primary tumour site to avoid strong signal interference from the primary tumour. We found that the tumours derived from control cells were more metastatic than the tumours from ERα-expressing cells.

Transwell assays were performed with MDA-MB-231 and ERα-expressing MCF-7 cells with gain or loss of ERα, and the results showed that ERα inhibited breast cancer cell invasion in vitro. We also conducted transwell assays with MCF-7 cells that were treated with different concentrations of fulvestrant, an ERα downregulator. The results showed that, although 0.29 and 10 nm fulvestrant significantly downregulated ERα expression and inhibited cell proliferation, it promoted cell invasion. In addition, we found that the invasive capacity of tamoxifen-resistant MCF-7 (MR) cells with lower ERα expression was significantly stronger than that in parental MCF-7 cells.
These observations indicated that loss of ERα expression promotes the invasive and metastatic ability of breast cancer cells.

**Loss of ERα induces the amoeboid migration of MCF-7 cells.** The migration of invasive cells in the 3D environment consists of the mesenchymal mode and the amoeboid mode. Next, we tracked and analysed the migration of different MCF-7 cell populations of control, Cas9-ERα and GM6001 (matrix metalloproteinase inhibitor)-treated Cas9-ERα in the 3D matrix by using phase holographic imaging assays. The result showed that loss of ERα in MCF-7 cells promoted the invasive ability of tumour cells in the 3D matrix and that GM6001 did not impair the migration speed of Cas9-ERα cells (Fig. 3a,b). In addition, GM6001 did not affect the migration speed of MCF-7 cells either (Supplementary Fig. 2a,b).

We also found that the control and Cas9-ERα MCF-7 cells did not express active MMP-2 or MMP-9 enzymes by zymography assay (Supplementary Fig. 2c). Using phase holographic imaging assays, we observed the following five phenotypes in the different populations of MCF-7 cells: 'stable adhesion', 'unstable lamellipodia', 'bleb', 'stable bleb' and 'unstable pseudopod' (Supplementary Movies 1–5). The 'bleb', 'unstable pseudopod' and 'stable bleb' phenotypes are known as amoeboid-like protrusions, which are associated with enhanced amoeboid movement. The incidence of a 'stable adhesion' phenotype was significantly reduced, and the amoeboid-like protrusions were more predominant in Cas9-ERα- or GM6001-treated Cas9-ERα cells compared with the control (Fig. 3c). Amoeboid migration requires high levels of phosphorylation of myosin light chain (p-MLC) and subcellular localization of p-MLC behind the cell nucleus to drive actomyosin contractility. Confocal assays showed an...
Figure 2 | ERα inhibits breast cancer metastasis in vivo and in vitro. MCF-7-luc2 (control) or CRISPR/Cas9-mediated ESR1-deleted MCF-7-luc2 cells (Cas9-ERα) were injected via tail veins into nude mice (n = 5). (a) Bioluminescence imaging of the control or the Cas9-ERα group at different time points was used to evaluate tumour progression in the lung. The lung metastases were determined using H&E staining. (b) Luciferase counts of the metastasis sites of mice on week 4. MDA-MB-231-luc2 (control) or ERα-overexpressing MDA-MB-231-luc2 (ERα) cells were injected into nude mice to generate xenograft models (n = 5). (c) Bioluminescence imaging at different time points was used to evaluate tumour progression. (d) Luciferase counts of the primary tumours of mice at different time points. (e) Representative images of mice on week 4 are shown after shielding the primary tumour. The lymphatic metastases were determined with H&E staining. (f) The lifetimes of mice injected with control or ERα-overexpressing cells. (g) A transwell assay (top) was performed to determine the effect of ERα on cell invasion by gain or loss of ERα in MDA-MB-231 or MCF-7 cells (n = 3). The expression of ERα (bottom) in different groups of MDA-MB-231 or MCF-7 cells was detected by western blotting. (h) A transwell assay (top) was performed to determine the invasive capability of parental MCF-7 and MR cells (n = 3). The expression of ERα (bottom) was detected by western blotting. (b,d,g,i) Graphs show mean ± s.e.m. **P<0.01, ***P<0.001. (b,d,g,i) Unpaired t-test; (f) log-rank test; (g,h) analysis of variance (ANOVA) with Dunnett t-test.
increased expression of the active form (phospho-Ser19) of myosin light chain (p-MLC) in Cas9-ERα MCF-7 cells, and more Cas9-ERα MCF-7 cells showed p-MLC rear distribution (Fig. 3d–f). In addition, the roundness index of Cas9-ERα cells was higher than that of control cells (Fig. 3g). Together, these results indicated that loss of ERα induces the amoeboid-like migration of MCF-7 cells.

**ERα is a transcriptional promoter of vinculin.** To elucidate the molecular mechanism of ERα action on the metastasis of breast cancer, RNA-sequencing was performed using MDA-MB-231-ERα cells and MDA-MB-231-vector cells. The main metastasis-associated genes that showed significantly altered expression levels are shown in Supplementary Table 3. The vinculin transcript was most significantly altered, and this result
ERα upregulates vinculin expression in breast cancer cells. MCF-7 cells cultured in oestrogen-depleted medium were treated with 0 or 5 nm oestrogen. We observed that the mRNA levels of vinculin were higher in MCF-7 cells treated with 5 nm oestrogen than in control cells (Supplementary Fig. 4a). In addition, the mRNA and protein levels of vinculin were higher in ERα-positive breast cancer cell lines than in ERα-negative cell lines (Fig. 5a; Supplementary Fig. 4b). To further investigate the role of ERα in regulating vinculin transcription and expression, real-time PCR and western blotting were performed to assess the effect of gain or loss of ERα expression on the level of vinculin mRNA and protein. The results showed that silencing endogenous ERα by using short interfering RNAs (siRNAs) led to a downregulation of vinculin in MCF-7 and ZR-75-1 cells (Fig. 5c). Knockdown of ERα in MDA-MB-231 and MCF-7 cells (Fig. 5d). Collectively, these results demonstrated that ERα directly facilitates vinculin transcription by specifically binding to the ERE region of the vinculin promoter.

Vinculin downstream of ERα is important for metastasis. To investigate the role of vinculin, downstream of ERα, in breast cancer metastasis, we performed tail vein injections of control or Cas9-vinculin MCF-7-luc2 cells in athymic mice and examined the expression levels of vinculin by western blotting (Supplementary Fig. 5a,b). We found that, compared with control cells, the Cas9-vinculin cells showed a more profound metastatic potential to the lungs (Fig. 6a,b). In addition, extravasation is a crucial step during tumour metastasis. To test whether vinculin involved in this process, the lung extravasation assay was performed. Similar numbers of control or vinculin-depleted MCF-7 cells lodged in the lung capillaries 0.5 h after injection (Fig. 6c). However, after 24 h the number of vinculin-depleted MCF-7 cells that remained in the lung parenchyma was more than the number of control cells (Fig. 6c,d).

We further studied an orthotopic mouse model of breast cancer, using MDA-MB-231 cells stably expressing ERα with or without vinculin knockdown (Supplementary Fig. 5c). Vinculin expression levels were detected by western blotting (Supplementary Fig. 5d,e). The results showed that tumours formed by vinculin knockdown cells were more metastatic compared with those of the control cells, and, after 4 weeks, lymph node metastasis was observed in the mice injected with vinculin knockdown cells (Supplementary Fig. 5f,g). The mice inoculated with vinculin knockdown cells had shorter lifetimes, probably because of the distant metastasis (Supplementary Fig. 5h). In addition, a transwell assay also showed that vinculin knockdown in MDA-MB-231 cells stably expressing ERα rescued the invasive capacity of the cells (Fig. 6e).

Furthermore, it has been reported that MDA-MB-231 cells can migrate through either the mesenchymal or amoeboid mode, and the amoeboid morphology might be predominant after protease inhibition. Using zymography on conditioned two-dimensional (2D) or 3D medium, we observed that MDA-MB-231 cells expressed active MMP-2 or MMP-9 enzymes (Supplementary Fig. 5i). We also observed that GM6001 indeed decreased the invasive capacity of MDA-MB-231 cells in transwell assays (Supplementary Fig. 5j,k). Moreover, we observed that GM6001 induced round cell morphology and that depleted vinculin expression reversed the decreased invasive capacity of the GM6001-treated MDA-MB-231 cells to some degree (Supplementary Fig. 5j,k). Next, we performed orthotopic injection of control, GM6001-treated control or GM6001-treated Cas9-vinculin MDA-MB-231 cells in athymic mice. Tumours from GM6001-treated cells were less metastatic than those from control cells, and tumours from GM6001-treated Cas9-vinculin cells and control cells had similar levels of metastasis (Fig. 6f). In addition, GM6001-treated cells in primary tumours showed round cell morphology with a higher average roundness score than control cells in vivo (Fig. 6h). In addition, the mice inoculated with control or GM6001-treated Cas9-vinculin cells had shorter lifetimes than those inoculated with GM6001-treated control cells (Supplementary Fig. 5l). These results showed that GM6001 can induce the rounded-amoeboid morphology of cells and loss of vinculin in GM6001-treated breast cancer cells is associated with increased metastatic potential.

Loss of vinculin promotes amoeboid features of cancer cells. The amoeboid features contained membrane blebbing, cell rounding, high actomyosin contractility and increased invasion. To investigate whether loss of vinculin could promote the amoeboid features of cells, we tracked and analysed the migration speed of different MCF-7 cell populations of control and Cas9-vinculin in the 3D matrix by using phase holographic imaging assays (Fig. 7a). We found that loss of vinculin in MCF-7 cells promoted the instantaneous speed of tumour cells in the 3D matrix (Fig. 7b). In addition, after depletion of vinculin, the activity of MLC increased, thus indicating that loss of vinculin promoted actomyosin contractility in breast cancer cells
We also found that the percentage of amoeboid-like protrusions, including ‘bleb’, ‘stable bleb’ and ‘unstable pseudopod’, in Cas9-vinculin cells significantly increased (Fig. 7e; Supplementary Movies 6–10). Confocal assays further showed that the control MCF-7 cells formed actin stress fibres, which were concomitant with p-MLC being evenly distributed in the cytoplasm, whereas in vinculin-depleted MCF-7 cells, few stress fibres were observed, and p-MLC was mostly polarized to the rear part of the cell (Fig. 7f). Moreover, Cas9-vinculin cells exhibited cell rounding and dynamic blebbing (Fig. 7f,g) and, compared with control cells, more Cas9-vinculin cells displayed p-MLC rear distribution (Fig. 7h).

**ERα correlates with vinculin in breast cancer tissues.** To investigate in a clinical setting whether ERα suppresses breast cancer metastasis by facilitating vinculin expression, we measured the expression of vinculin in the same 124 human primary breast

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**Figure 4 | ERα is a transcriptional promoter of vinculin.** (a) The schematic diagrams of human vinculin promoter containing four putative ERα-binding sites and the amplification regions of ChIP primers are presented in the top panel. The bottom panel shows the amounts of DNA fragments that were normalized to the total input genomic DNA from MCF-7 cells precipitated by either anti-ERα monoclonal antibody or control IgG (n = 3). (b) Schematic diagrams of vinculin promoter truncation. FL indicates the full length of the vinculin promoter, and the corresponding truncation is represented by F1, F2 and F3. (c) Luciferase activity was measured in MCF-7 cells co-transfected with truncated vinculin promoter and ERα siRNA. pGL3-basic plasmid was used as a negative control (n = 3). (d) Luciferase activity was measured in MDA-MB-231 cells co-transfected with the truncation of the vinculin promoter and the ERα vector (n = 3). (e) Schematic diagrams of a 23-bp ERE in the vinculin promoter and respective mutation. Mut1: deletion of GGCC sequence; Mut2: replacing GGCC with TTAA; Mut3: totally deleting ERE. (f) Luciferase activity was measured in MDA-MB-231 cells transfected with mutations of the vinculin promoter and ERα vector (n = 3). (g) Luciferase activity was measured in MCF-7 cells transfected with mutations of the vinculin promoter (n = 3).

(a,c,d,f,g) The data are shown as the mean ± s.e.m. **P < 0.001 (a,c,d,f,g) ANOVA with Dunnett t-test.
**Figure 5 | ERα upregulates the expression of vinculin in breast cancer cells.** (a) Real-time PCR detecting the transcription levels of vinculin in breast cancer cell lines. The results were normalized to GAPDH (n = 3). (b) Western blotting was conducted to detect the protein levels of vinculin and ERα in four breast cancer cell lines. (c) MCF-7 and ZR-75-1 cells were transfected with ERα siRNAs or scrambled RNA and subjected to quantitative reverse transcriptase PCR (qRT-PCR) assay (n = 3). (d) MDA-MB-231 and SK-BR-3 cells that were transfected with ERα vector or vector subjected to qRT-PCR assay (n = 3). (e) Western blotting to detect the protein level of vinculin and ERα protein by the gain or loss of ERα in breast cancer cell lines. (f) Confocal assay for ERα and vinculin expression in MCF-7 (control or si-ERα) cells. Nuclear staining with 4,6-diamidino-2-phenylindole (DAPI) is also shown. Scale bar, 10 μm. (a, c, d) The data are shown as the mean ± s.e.m. **P<0.01, ***P<0.001 (a, c) ANOVA with Dunnett t-test; (d) unpaired t-test.
cancer tissues and human breast cancer lymph node metastases that were used in the ERα analysis. The results showed that vinculin abundance was also significantly higher in human primary breast cancer tissues than in lymphatic metastases (Fig. 8a). A positive association was found between vinculin and ERα in both primary tumour \((P < 0.001, R^2 = 0.528)\) and lymphatic metastasis \((P < 0.001, R^2 = 0.366; \text{Supplementary Tables 4 and 5})\). The expression levels of vinculin are shown in Supplementary Fig. 6a.

We further compared the expression of these two molecules at the invasive and non-invasive fronts in primary breast cancer tissue. Similarly to the trend of ERα expression level, diminished
vinculin expression was also detected at the invasive front, whereas increased staining of vinculin was observed at the non-invasive front (Fig. 8b–e). These findings indicated that ERß might suppress human breast cancer metastasis by facilitating vinculin expression.

**Discussion**

Among all breast cancers, ERß-positive (ER+) tumours constitute the largest proportion, ~70% (ref. 30). Although sporadic publications have shown that the expression of ERß in MDA-MB-231 cells inhibits proliferation in vitro33, a plethora of laboratory and epidemiological data have demonstrated that ERß, by binding to oestrogen, is the major driving factor for growth in ERß+ breast cancers3,32. In the present study, we also found that expression of ERß indeed promoted proliferation of MDA-MB-231 cells in vitro and in vivo. ERß induces cell proliferation by increasing the expression of MYC and cyclin D1 (refs 33,34). Therefore, ERß has been used as a key target for endocrine therapy of ERß-positive breast cancer to block the proliferation of cancer cells35.

However, an increasing number of the clinical epidemiological investigations show that ERß-positive primary breast cancer patients have an increased frequency of ESR1 mutations in metastatic ERß+ breast cancer tissues and even ERß-negative metastatic relapse after receiving endocrine therapy1,7. Although a decline in ERß levels has been detected in invasive breast cancers5, the relationship between ERß and tumour metastasis is still far from clear. In the present study, we observed that, compared with those in the primary tumour, breast cancer cells in lymph node metastases expressed lower levels of ERß. More importantly, loss of ERß in lymphatic metastases was also positively associated with clinical stages and lymph node metastases. Even in the primary tumour, the expression of ERß at the invasive front was lower than that at the non-invasive front. Therefore, these findings indicated that ERß expression is inversely correlated with breast cancer metastasis. We further demonstrated that ERß is indeed capable of inhibiting breast cancer metastasis by using athymic mouse models and transwell assays. This finding is very important because it suggested that breast cancer cells with loss of expression of ERß should be not only be resistant to common adjuvant endocrine therapy but also have stronger invasion and metastasis capabilities. Consistently with this possibility, we found that an appropriate concentration of fulvestrant promoted cell invasion in vitro by downregulating ERß and that the invasive capacity of tamoxifen-resistant MCF-7 cells with lower ERß expression was stronger than that of parental MCF-7 cells. Collectively, we believe that repeated biopsies are necessary to reassess the receptor status in metastatic disease to guide endocrine therapy with greater precision.

Several studies have reported that ERß suppresses cellular motility and invasion51 by inhibiting the epithelial–mesenchymal transition in 2D conditions36,38–39. However, with regard to cell shape and movement, the 3D environment resembles the *in vivo* condition40. Some tumour cells can utilize amoeboid-like migration as an alternative to mesenchymal migration during migration in 3D cultures and *in vivo*41,42. To date, few studies have focused on the relationship between the amoeboid-like migration of breast cancer cells in a 3D environment and ERß expression. We demonstrated that ERß-positive MCF-7 cells expressed fewer active MMP-2 and MMP-9. Loss of ERß also promoted a more round shape and increased cell motility in the 3D matrix in vitro, and these effects were not impaired by the MMP inhibitor GM6001. The formation of amoeboid-like protrusions, such as blebs, often corresponds with the amoeboid phenotype and enhanced invasion and metastasis and blebbing movement has been widely promoted as a cancer cell migration strategy43–45. We observed the significantly increased formation of amoeboid-like protrusions in ERß-depleted MCF-7 cells in the present study. During the initiation of blebs, local dissociation of the membrane from the cortex or a local rupture of the actin cortex causes the bleb to form in the direction of the desired flow46,47. Furthermore, it has been reported that high contractility of the cell rear triggers bleb initiation33,50. Indeed, we observed that loss of ERß in MCF-7 cells induced the polarization of p-MLC at the cell rear, thereby resulting in high contractility of the cell rear, the formation of amoeboid-like protrusions and enhanced amoeboid-like migration. Even in the MDA-MB-231 cells that use mixed migration modes in a 3D matrix, ERß suppressed amoeboid-like migration after MMP inhibition, although overexpression of ERß did not alter the mesenchymal-like morphology of MDA-MB-231 cells in a 2D substrate (Supplementary Fig. 6b,c). Together, our results show that ERß inhibits the amoeboid-like migration of breast cancer cells in a 3D matrix.

With respect to the mechanisms by which ERß inhibits cancer metastasis, we focused on the regulation of ERß to vinculin expression based on transcriptome sequence analysis. Our data show that ERß is a novel regulator of vinculin expression in breast cancer and that vinculin is involved in ERß-mediated inhibition of breast cancer cell metastasis. As above, vinculin is a key regulator of focal adhesions and loss of vinculin in cells promotes focal adhesion turnover51. It was reported that focal adhesions could promote the formation of persistent stress fibres, which prevent the transition to amoeboid migration by competing with the cell cortex for recruitment of the actomyosin contractile machinery52. In the present study, our results showed that the depletion of vinculin in MCF-7 cells induced decreased cell–matrix adhesion (Supplementary Fig. 6d). More importantly, we found that in a 3D matrix vinculin depletion leads to decreased stress fibres and p-MLC mostly polarizes to the rear part of the cell, thus resulting in high contractility of the cell rear and induced bleb initiation.

Metastasis occurs when tumour cells detach from the epithelial sheets and invade surrounding tissue. It was reported that vinculin was important for cadherin-mediated cell–cell junctions53 and might suppress metastasis formation *in vivo* by

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**Figure 6 | Vinculin downstream of ERß is important for metastasis.** MCF-7-luc2 cells and CRISPR/Cas9-mediated VCL-deleted MCF-7-luc2 cells were injected into nude mice via the tail vein to generate xenografts (*n* = 5). (a) Bioluminescence imaging of the control or the Cas9-vinculin group. Tumour formation in the lung was determined using H&E staining. (b) Luciferase counts in the lungs of mice on week 4. (c) Representative confocal images of mouse lungs 0.5 and 24 h after tail vein co-injection of control MCF-7 cells (red) and Cas9-vinculin MCF-7 cells (green). Scale bar, 100 μm. (d) Quantification of cells retained in the lung after tail vein injection (*n* = 4 mice). (e) A transwell assay was performed in ERß-overexpressing MDA-MB-231 cells infected with lentivirus containing vinculin short hairpin RNA (shRNA; sh-VCL) or scrambled RNA (shramble; *n* = 3). Control MDA-MB-231-luc2 cells, GM6001-treated control cells or GM6001-treated and CRISPR/Cas9-mediated VCL-deleted MDA-MB-231 cells were injected into nude mice to generate xenografts (*n* = 5). (f) Representative bioluminescence images of different groups on week 4 are shown. The cell morphology of the primary tumour was determined by H&E staining. Scale bar, 50 μm. The particular section of H&E image was enlarged to highlight the cell morphology. (g) Luciferase counts of metastasis sites on week 4. (h) Roundness index of corresponding MDA-MB-231 cells from the primary tumour in *f* (*n* = 200 cells). The data are shown as the mean ± s.e.m. ns *P* > 0.05, **P < 0.01, ***P < 0.001. (b, d, e, g, h) Unpaired t-test; (b, d, e, g, h) ANOVA with Tukey’s post hoc test.
promoting cadherin-mediated retention of tumour cells in primary tumours. Meanwhile, we found that depletion of vinculin in MCF-7 cells induced loss of cell–cell contact (Supplementary Fig. 6e,f). However, we also found that loss of vinculin in MCF-7 cells induced cell rounding, increased actomyosin contractility, fast shape changes and increased invasion. These observations indicated that the effect of vinculin on amoeboid features of cells might contribute to the process of local invasion or even the metastasis formation besides the effect on cell–cell adhesion. Moreover, it was reported that high levels of actomyosin contractility in cancer cells could promote and efficient lung colonization or seeding and rapid extravasation was associated with the amoeboid features. Our in vivo extravasation assay showed that the number of vinculin-depleted cells that extravasated into the lung parenchyma after 24 h was more than the number of control cells. These observations indicated a potential involvement of vinculin in efficient retention in the lungs via its regulation of amoeboid features. More importantly, we do not discard the possibility that reduced cell–cell adhesion and anoikis resistance that resulted from loss of vinculin may be involved in metastasis; however, we believe that we have identified amoeboid features regulated by vinculin, which are associated with metastasis.

In summary, our results demonstrate that ERα suppresses breast cancer metastasis by regulating vinculin. Our findings provide novel insight into the association of ERα loss during endocrine therapy with enhanced invasive and metastatic ability of breast cancer cells and should aid in a more comprehensive understanding of the role of vinculin in breast cancer metastasis.
understanding of the effects of endocrine therapy in clinical treatment.

Methods

Antibodies and inhibitors. Antibodies and dilutions used were as follows: ERα (ab32063; immunohistochemistry, 1:150; immunoblotting, 1:750; immunofluorescence, 1:150) from Abcam; vinculin (ab18058; immunohistochemistry, 1:100; immunoblotting, 1:400; immunofluorescence, 1:150) from Abcam; CD68 (916104; immunohistochemistry, 1:100) from BioLegend; GAPDH (CW0101; immunoblotting, 1:1,000) from CWBIOTECH; p-MLC (3671; immunoblotting, 1:750; immunofluorescence, 1:50) from Cell Signaling Technology; MLC (10906-1-AP; immunoblotting, 1:500) from Proteintech; F-actin (40734ES75; immunofluorescence, 1:100) from YEASEN; fluorescein isothiocyanate (FITC)-conjugated anti-rabbit and Cy3-conjugated anti-mouse antibodies from ZHUANGZHIBIO.

The inhibitors are as follows: in vivo: GM6001 (MedChem Express, Monmouth Junction, NJ, USA) was subcutaneously injected every 3 days at 100 mg per kg body weight. In vitro: GM6001 (27 nm).

Cell lines and culture. Human breast cancer cell lines MCF-7, ZR-75-1, MDA-MB-231 and SK-BR-3 were obtained from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). The breast cancer cell...
line MDA-MB-231-1uc2, which expresses luciferase, was a kind gift from Dr Xia Habin of Shannxi Normal University. The parental MCF-7 cells and tamoxifen-resistant MCF-7 cells were gifted from Dr Zhang of the Fourth Military Medical University. All cell lines were authenticated by the analysis of short tandem repeat (STR) profiles and 100% matched the standard cell lines in the DSMZ data bank. All cells were tested negative for cross-contamination of other human cells and mycoplasma contamination.

The plasmid GV392 (Lenti-Case9-sgRNA-puromyacin) was purchased from GeneChem (Shanghai, China). The parental MCF-7 cells and NATURE COMMUNICATIONS | DOI: 10.1038/ncomms14483 | www.nature.com/naturecommunications

Plasmid construction and RNA interference. The pC2DNA3.1(–) plasmid was retained by our laboratory. The synthesized nucleotides encoding wild-type, truncated or mutant vinculin promoters were digested with Kpn1 and Xho1 and cloned into a pG3L-basic vector. Three ERα siRNA molecules (Gene Pharma, Shanghai, China) were used to knockdown ERα expression in breast cancer cells. The vinculin short hairpin RNAs were designed and synthesized by GeneCHEM (Shanghai, China). No negative control for non-sequence-specific effects. All sequences are listed in Supplementary Tables 6 and 7. siRNAs and plasmids were transfected into cells using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA) following the manufacturer’s instructions. The plasmid GV392 (Lenti-Case9-sgRNA-puromyacin) was purchased from GeneCHEM.

Animal studies. Female athymic mice, 4-week of age, were selected. The animal study was performed with the approval of the Institutional Animal Care and Use Committee of the FMMU. For mammary fat pad injection, 3 × 10^5 viable breast cancer cells were injected into the mammary fat pads of athymic mice (n = 5). For tail intravenous injection, 1 × 10^6 viable breast cancer cells were injected into athymic mice from tail vein (n = 5). Then, the mice were anaesthetized and injected with luciferin and imaged using an IVIS-100 System at the UAB Small Animal Imaging Core Facility. Light emission from animal tissue was measured using a software provided by the vendor (Xenogen). In parallel, an equivalent number of tumour cells were inoculated into the same sites of other athymic mice to record their survival time. The operator who performed injection of tumour cells was blinded with the group allocation. At least two independent experiments were performed.

Cell proliferation assay. The proliferation ability of the cells was measured by Cell Counting Kit-8 (CCK-8) solution (Tea biotech, Shanghai, China). Briefly, breast cancer cells were seeded on 96-well plates (Corning, USA) at a concentration of 2 × 10^4 cells per well and incubated at 37°C overnight. The Cell Counting Kit-8 reagents were then added to a subset of wells when cells grew for 24, 48, 72 or 96 h. After the cells were incubated for 2 h at 37°C, we quantified the absorbance at 450 nm using a microplate reader (Bio-Rad). Each group was made in quintuplicate.

Zymography. The assessment of MMP-2 and MMP-9 activity was performed using a Gelatin Zymography Kit (Xin Sails Biotechnology, Shanghai, China). Briefly, conditioned serum-free medium of breast cancer cells grown on plastic for 36 h was loaded into the polyacrylamide gels containing 10% gelatin and then electrophoresed at 20 mA per gel for 90 min. Following electrophoresis, gels were rinsed twice with 10 ml 1 × Buffer A for 30 min at room temperature and incubated with 10 ml 1 × Buffer B for 3 h at 37°C. Gels were then stained with 0.5% Coomassie Blue for 2 h and destained five times for 20 min with destaining solution. The pore-molecular weight of MMP-2, MMP-9 and pro-MMP-9 is 66–72, 92 and 130 kDa.

Phase holographic imaging assay. Overall, 3 × 10^5 MCF-7 cells of different groups were embedded in Matrigel (diluted with serum-free medium) at 37°C overnight. Half an hour before the experiment, the transwell insert containing Matrigel (diluted with 10% FBS) was seeded on 3D matrix as a chemottractant, which was in the right side of the observation point. Then, HoloMonitorTM M4 (PerkinElmer) was used to track and record related parameters of the movement of cells in 3D matrix in 4 h. Motility and protrusion data were obtained using × 40 objective. The cells were imaged every 2 min and the movies were played back at 15 frames per second. The main criteria used to group cells into the different phenotypes were the size of protrusions (<2 μm or not) and the lifetime of protrusions (<2 min or not).

Calculation of cell roundness. It was assessed by dividing the shortest diameter of each cell by the longest one (ratio/b/ratio-a) to produce a score between 0 and 1, with perfectly round cells having a score of 1 (ref. 9).

ChIP assay. ChIP experiments were performed using the EZ ChIP Chromatin ImmunoPrecipitation Kit (Millipore, Billerica, USA). Four primer sets were designed to flank-related putative ERα-binding sites in the promoter region of vinculin. Details of the primer sequence are listed in Supplementary Table 9. Briefly, cells were fixed with 1% paraformaldehyde and sonicated seven times for 10 s each using a sonicator with a microtip in a 1.5 ml tube. Anti-ERα antibody or control human IgG was applied to pull down the chromatin associated with ERα. The chromatin–antibody complexes were collected with Protein G-Agarose. After washing and elution of the complexes from the beads, the DNA–protein crosslinks were reversed at 65°C overnight. The amounts of the specific DNA fragment were then quantified by real-time PCR and normalized against the genomic DNA preparation from the same cells. Each group was made in triplicate.

Luciferase reporter assay. Briefly, breast cancer MCF-7 or MDA-MB-231 cells were seeded in 24-well plates at 50% confluence and transfected with either ERα siRNAs or ERα plasmid using Lipofectamine 2000 and then co-transfected with pRL-TK and vinculin promoter (pGL3-basic-vinculin). Thirty-six hours later, the cells were lysed in a passive lysis buffer (Promega, San Luis Obispo) and the luciferase activity was measured. Each group was made in triplicate.

Quantitative real-time PCR. Total RNA was isolated from cultured cells with RNAiso Plus (Takara, Dalian, China), and cDNA was synthesized with the PrimeScript RT Reagent Kit (Takara). Then, 2 μl of cDNA was used for real-time PCR reactions in a Prism 7500 real-time thermocycler (Applied Biosystems, Foster City, CA, USA) with SYBR Green Ex Taq (Takara) according to the manufacturer’s instructions. The primer sequences are provided in Supplementary Table 10. Each group was made in triplicate.

Western blot analysis. In brief, proteins were transferred to polyvinylidene difluoride membranes after SDS–PAGE using a Bio-Rad Semi-Dry electrophoretic cell. Western blot analyses were performed using specific antibodies followed by horseradish peroxidase–conjugated IgG antibody. Enhanced chemiluminescence (Pierce) was used for immunoreactive protein visualization. Uncropped scans of the blots are shown in Supplementary Figs 7 and 8.

Immunofluorescence analysis. For the observation of the subcellular localization of p-MLC and F-actin, 4 × 10^5 breast cancer cells were embedded in Matrigel at 37°C overnight. For the observation of the expression of ERα and vinculin, 4 × 10^5 breast cancer cells were seeded on glass plates at 37°C overnight. The cells were then washed twice with cold PBS and fixed with 4% paraformaldehyde for 20 min, followed by permeabilization with 0.2% Triton X-100 diluted with PBS for 30 min at room temperature. The cells were incubated overnight at
Lung extravasation assay. Overall, 5 × 10⁵ Cas9-vinculin MCF-7 cells (enriched green fluorescent protein, eGFP) and 5 × 10⁵ control MCF-7 cells (monomeric red fluorescent protein, mRFP) were mixed and injected into the tail vein of nude mice (female, 6 weeks old). Mice were killed after 0.5 or 24 h and the lungs were fixed (4% formaldehyde for 24 h) and examined for fluorescently labelled cells under a confocal microscope (FluoView FV1000, Olympus, Tokyo, Japan).

Cell adhesion assay. In all, 1 × 10⁴ cells were seeded on Matrigel-coated 96-well plate and cultured for 2 h. After washing, adhesion cells were counted using optical microscope (Olympus). The adhesion cells distributed in a randomly selected view were counted under a microscope (× 20) and averaged.

Binding of cells to recombinant E-cadherin. A 96-well plate was coated with 100 μl human recombinant E-cadherin Fc chimera (648-EC-100, R&D Systems)/well at 1.5 μg/ml in PBS at 37°C for 1 h. The plate was washed with PBS 3 times and blocked by adding 100 μl per well 1% BSA in PBS at 37°C for 30 min. Then 3 × 10⁵ cells per well are added to Recombinant Human E-Cadherin Fc Chimera coated plates and the plate was kept at 37°C for 90 min. After washing, adhesion cells were counted by optical microscope (Olympus, Tokyo, Japan). The adhesion cells distributed in a randomly selected view were counted under a microscope (× 40). Each group was made in octuplicate.

Statistical analysis. Statistical analysis was performed using the SPSS statistical software (SPSS16.0, Chicago, CA, USA). A value of P < 0.05 was considered statistically significant. A random number table was used to randomize the mice into control and treatment groups. The numbers of mice (in vivo) were determined on the basis of our pre-tests and previous experience with similar experiments. Sample size was chosen to ensure adequate and statistically significant results. Investigators who determined the expression levels of ER were blinded with respect to the treatment allocation. The in vitro experiments were repeated at least three times. The statistical tests were two-sided.

Data availability. The authors declare that the main data supporting the findings of this study are available within the article and its Supplementary Information Files. Extra data are available from the corresponding authors upon request.

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