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Prolyl-4-hydroxylase α subunit 2 promotes breast cancer progression and metastasis by regulating collagen deposition

Gaofeng Xiong1, Lei Deng1, Jieqing Zhu1, Piotr G Rychahou1,2 and Ren Xu1,3*

Abstract

Background: Increased collagen deposition provides physical and biochemical signals to support tumor growth and invasion during breast cancer development. Therefore, inhibition of collagen synthesis and deposition has been considered a strategy to suppress breast cancer progression. Collagen prolyl-4-hydroxylase α subunit 2 (P4HA2), an enzyme hydroxylating proline residues in -X-Pro-Gly- sequences, is a potential therapeutic target for the disorders associated with increased collagen deposition. However, expression and function of P4HA2 in breast cancer progression are not well investigated.

Methods: Gene co-expression analysis was performed in the published microarray datasets to identify potential regulators of collagen I, III, and IV in human breast cancer tissue. Expression of P4HA2 was silenced by shRNAs, and its activity was inhibited by 1, 4-DPCA, a prolyl-4-hydroxylase inhibitor. Three-dimensional culture assay was used to analyze roles of P4HA2 in regulating malignant phenotypes of breast cancer cells. Reduced deposition of collagen I and IV was detected by Western blotting and immunofluorescence. Control and P4HA2-silenced breast cancer cells were injected into fat pad and tail vein of SCID mice to examine effect of P4HA2 on tumor growth and lung metastasis.

Results: Using gene co-expression analysis, we showed that P4HA2 was associated with expression of Col1A1, Col3A1, and Col4A1 during breast cancer development and progression. P4HA2 mRNA levels were significantly upregulated in breast cancer compared to normal mammary tissue. Increased mRNA levels of P4HA2 correlated with poor clinical outcome in breast cancer patients, which is independent of estrogen receptor status. Silencing P4HA2 expression or treatment with the P4HA inhibitor significantly inhibited cell proliferation and suppressed aggressive phenotypes of breast cancer cells in 3D culture, accompanied by reduced deposition of collagen I and IV. We also found that knockdown of P4HA2 inhibited mammary tumor growth and metastasis to lungs in xenograft models.

Conclusion: These results suggest the critical role of P4HA2 in breast cancer progression and identify P4HA2 as a potential therapeutic target and biomarker for breast cancer progression.

Keywords: Tumor microenvironment, Breast cancer, Collagen deposition, Cancer progression, Cell proliferation
Background
Extracellular matrix (ECM) is an important component of tumor microenvironment and plays critical roles in cancer development [1-3]. Collagens are the major structural ECM proteins and form fibers or networks in tumor tissue [4-6]. Cell-collagen interaction controls a variety of cellular activities including proliferation, migration, and invasion through integrin and discoidin domain receptor [7-9]. Enhanced expression and deposition of collagens are associated with tumor development and progression [10-12]. Recent studies demonstrate that increased collagen deposition and crosslinking enhance the stiffness and density of mammary tissue [5,10,13], which is an important risk factor for breast cancer development. Type I collagen has been identified as a prognosis marker and is associated with cancer recurrence in human breast cancer patients [14]. Collagen VI knockout mice have reduced primary tumor formation and growth [12], while enhancing collagen deposition and inhibiting collagen degradation significantly enhances tumor initiation and tumor growth [5,10]. In addition, cancer cell invasion usually occurs at tumor-stromal interfaces with oriented collagen fibers, and aligned collagen fibers can facilitate cell migration and metastasis [5,10,11,15]. These results indicate that increased collagen expression and deposition promotes breast cancer development and progression by enhancing tumor growth and invasion. Therefore, inhibiting collagen synthesis or deposition is a promising strategy to suppress breast cancer progression.

Collagen biosynthesis is a multistep process that involves several post-transcription modification enzymes, and one of the most important members of these enzymes is collagen prolyl-4-hydroxylase [16]. It catalyzes the formation of 4-hydroxyproline by hydroxylating proline residues in -X-Pro-Gly- sequences [17-20]. Collagen prolyl-4-hydroxylase resides within the lumen of the endoplasmic reticulum (ER) [18] and its expression is used as a marker for collagen synthesis [21,22]. Because 4-hydroxyproline residues formed in this reaction are essential for triple helix formation and stabilization of collagen [22-24], inhibiting the prolyl-4-hydroxylases activity efficiently blocks collagen synthesis and deposition. All known vertebrate collagen prolyl-4-hydroxylases are α2β2 tetromers consisting of two α subunits and two β subunits. Each α subunit contains the peptide substrate binding domain and the two catalytic sites of the enzyme, and the β subunits have been identified as protein disulfide isomerases [17,19,25]. Three types of collagen prolyl-4-hydroxylases α isoforms (P4HA1, P4HA2 and P4HA3) have been identified in human tissue. P4HA1 is expressed in most cell types; P4HA2 is mainly expressed in chondrocytes, osteoblasts, and capillary endothelial cells; P4HA3 expression is detected in adult and fetal tissues at very low levels compared to P4HA1 and P4HA2 [18,26]. Increased P4HA2 expression has been detected in many solid tumors, including oral cavity squamous cell carcinoma, papillary thyroid cancer, and breast cancer [27-30], however, the function of P4HA2 in cancer progression largely remains to be determined.

Here, we showed that expression of P4HA2 and collagen genes (Col1A1, Col3A1, and Col4A1) is significantly correlated during breast cancer development and progression, and that increased mRNA levels of P4HA2 are associated with poor prognosis in breast cancer patients. Silencing P4HA2 or treatment with the P4HA inhibitor attenuates cell proliferation and suppresses aggressive 3D phenotypes, tumor growth, and cancer metastasis, which are accompanied by reduced collagen deposition. These results suggest that P4HA2 promotes breast cancer progression by enhancing collagen deposition and it may serve as a potential therapeutic target for breast cancer.

Methods
Antibodies and reagents
The Click-iT® EdU Alexa Fluor® 488 Imaging Kit and Alexa Fluor® 594 phalloidin were from Invitrogen. Matrigel (lrECM) and Type I collagen were from BD Bioscience. ShP4HA2 plasmids were purchased from Sigma. 1, 4-DPCA was purchased from Cayman Chemical. Masson's trichrome stain kit was purchased from Polysciences, Inc. The following antibodies were obtained as indicated: integrin α6 (Millipore); collagen I (Abcam); collagen IV (Abcam); P4HA2 (Santa Cruz); tubulin (Millipore).

Cell culture and virus preparation
HMT-3522 T4-2 cells (a kind gift from Dr. Mina J. Bissell) were maintained on tissue culture plastic as previously described [31]. MDA-MB-231 cells were propagated in DMEM/F12 (Sigma) with 10% fetal bovine serum (Invitrogen). MDA-MB-157 cells and ZR-75-1 cells were propagated in DMEM (Sigma) with 10% fetal bovine serum. ZR-75-1 cells: ER-positive and PR positive; T4-2 cells, MDA-MB-231 cells and MDA-MB-157 cells: ER-negative and PR negative.

3D laminin-rich extracellular matrix (3D lrECM) on-top cultures were prepared by trypsinization of cells from tissue culture plastic, seeding of single cells on top of a thin gel of Engelbreth-Holm-Swarm (EHS) tumor extract (Matrigel: BD Biosciences, 354230), and addition of medium containing 5% EHS. T4-2 cells were seeded at a density of 2.1 × 10^5 cells per cm^2; MDA-MB-157 cells, ZR-75-1 cells, and MDA-MB-231 cells were seeded at 1.4 × 10^4 cells per cm^2. T4-2 cells were maintained in their propagation medium with media change every 2 days. MDA-MB-157 cells, ZR-75-1 cells and MDA-
MB-231 cells were maintained in H14 medium with 1% fetal bovine serum. The cell colonies cultured in 3D were imaged and used for immunofluorescence staining at Day 4 after seeding.

HEK293 FT cells were transfected with scrambled RNA sh-control vector or sh-P4HA2-1 (CCGGGCCG AATTCCTCACCTCTATTTCGGAATAGGTTG AAAGAATCGGCTTGG), sh-P4HA2-2 (CCGGGCGAT CTCGGAAAGATACATCTCGAGATGTACTCTTT CAGAGACTTGTCTTTTG) plus packaging lentivector using lipofectamine (Invitrogen). Cancer cells were infected with lentivirus and selected by puromycin 48 h after infection.

**Immunofluorescence and Masson’s trichrome staining**

Cells in IrECM gel were smeared on slides, dried briefly, and fixed with 4% paraformaldehyde and permeabilized with 0.5% Triton X-100. Immunostaining was performed as previous described [32]. Stained samples were imaged with a Nikon upright epifluorescence microscope or a confocal system comprised of an Olympus IX81 microscope.

Xenograft tumor sections were de-paraffined and hydrated from xylene, 100% ethanol, 95% ethanol, 85% ethanol and 70% ethanol to distilled water. For Masson’s trichrome staining, slides were re-fixed with Bouin’s solution at 60°C for 60 minutes. Slides were washed in running tap water for 5 minutes and stained in Weigert’s working hematoxylin for 10 minutes. Then they were washed in running tap water for 5 minutes and stained in Biebrich scarlet-acid fuchsin solution for 5 minutes. Slides were rinsed in distilled water and differentiated in phosphomolybdic-phosphotungstic acid solution for 10 minutes, transferred to aniline blue solution and stained for 5 minutes. Slides were rinsed in distilled water and images were taken with a Nikon microscope. The percentage of collagen was quantified by calculating the ratio of blue staining (collagen) area in the total area of the tumor section using Imagescope analysis software [33].

**Western blot analysis**

Cells grown on plastic were lysed in situ in 2% SDS in PBS buffer containing phosphatase and protease inhibitor cocktails (Calbiochem). Protein concentration was measured using DC™ protein assay (Bio-Rad). Control and shP4HA2 cells were trypsinized and counted; equal amounts of conditional medium (normalized to cell number) were precipitated by pre-cooled acetone. Equal amounts of protein lysates and cell conditional medium were subjected to SDS gel electrophoresis, immunoblotted, and detected with an ECL system (Pierce). Western blotting results were quantified using AlphaInnotech analysis software.

**Transwell invasion assay**

The Transwells (Corning) were coated with 60 mL 1 mg/mL Matrigel and incubated for 30 minutes at 37°C. Sh-control or sh-P4HA2 silencing MDA-MB-231 cells (1 × 10^5 cells in 200 μL medium) were plated on the top of the Transwell filter and incubated in 37°C 5% CO2 for 24 h. The invaded cells on the bottom face of the filter were fixed by methanol and stained with 8% crystal violet. Images were taken with a Nikon microscope and the number of invaded cells was counted.

**Xenograft experiment**

Female SCID mice (6 weeks old) were randomly grouped and injected with 2 × 10^6 sh-control or shP4HA2-1 MDA-MB-231/Luc cells via tail vein. To detect lung metastasis, bioluminescent images were taken day 30 after cancer cells injection with IVIS Spectrum. Mice were sacrificed week 5 after cancer cells injection.

**Lung metastasis experiment**

Female SCID mice (6 weeks old) were randomly grouped and injected with 1 × 10^6 (in 200 μL PBS) sh-control or sh-P4HA2-1 MDA-MB-231/Luc cells via tail vein. At the experimental endpoint, tumors were harvested and fixed with 4% PFA for paraffin-embedded section. All procedures were performed within the guidelines of the Division of Laboratory Animal Resources at the University of Kentucky.

**Kaplan Meier survival analysis and other statistical analyses**

Kaplan-Meier survival analysis was performed in a large combined breast cancer dataset [34]. Breast cancer patients were grouped by estrogen receptor (ER)-positive (n = 1452) and ER-negative (n = 473), and tumor samples were equally grouped into low and high P4HA2 expression based on the mRNA levels. Significant differences in overall survival time were assessed with the Cox proportional hazard (log-rank) test.

Analysis of P4HA2 mRNA levels in normal and malignant tissues was performed in the TCGA breast cancer dataset that was downloaded from Oncomine. The association between mRNA levels of P4HA2 and collagen genes was evaluated by the Spearman correlation analysis. All experiments were repeated at least twice. Results are reported as mean ± S.E.M; the significance of difference was assessed by independent Student’s t-test. P < 0.05 represents statistical significance and P < 0.01 represents sufficiently statistical significance. All reported P values were 2-tailed. Statistical analysis was conducted.
with SigmaPlot (Systat Software, Inc.) and SAS (version 9.2; SAS Institute Inc.).

Results and discussion
P4HA2 is associated with collagen expression and poor prognosis in human breast cancer
To determine which genes contribute to collagen deposition during cancer development and progression, we performed gene co-expression analysis using the published microarray datasets generated from human breast cancer tissues [35]. P4HA2 was identified as one of the genes associated with ECM protein expression [35]. By analyzing gene expression in the TCGA breast cancer dataset downloaded from Oncomine, we found expression of Col1A1, Col3A1, and Col4A1 was significantly correlated with P4HA2 levels in normal and malignant breast tissues (p < 0.001, Figure 1A, B and C). Similar results were also obtained in another published microarray dataset [36] (Additional file 1: Figure S1). To assess whether P4HA2 expression is associated with breast cancer development and progression, we analyzed mRNA levels of P4HA2 in the TCGA human breast cancer

Figure 1 P4HA2 is associated with collagen expression and poor prognosis in human breast cancer. (A-C) Scatterplot of correlated mRNA levels between P4HA2 and (A) Col1A1, (B) Col3A1, and (C) Col4A1 in normal and malignant breast tissues (n = 593). The mRNA levels of P4HA2 were acquired from the TCGA microarray dataset generated from human breast cancer. (D) P4HA2 mRNA levels in IBC (invasive ductal breast carcinoma) (n = 76), IDBC (invasive ductal breast carcinoma) (n = 398) and ILBC (invasive lobular breast carcinoma) (n = 36) is higher than normal breast tissue (n = 61). (E) P4HA2 mRNA expression is higher in ERBB2 (epidermal growth factor receptor 2) negative tumors than in ERBB2-positive cancer tissues. (F) P4HA2 mRNA levels are associated with stages of breast cancer. (G) Kaplan-Meier survival analysis showed the association between P4HA2 expression and clinical outcomes. Breast cancer patients were grouped by ER status (ER-positive, n = 1452; ER-negative, n = 473). Tumor samples were classified into low and high P4HA2 expression based on the mRNA levels. Tumor had high P4HA2 expression levels in a shorter overall survival period. The association of P4HA2 expression and clinical outcome is ER status independent (*p < 0.05; **p < 0.01; ***p < 0.001).
microarray dataset. \textit{P4HA2} expression was significantly upregulated in invasive breast carcinoma (p < 0.001), invasive ductal breast carcinoma (p < 0.001) and invasive lobular breast carcinoma (p < 0.001) compared to normal breast tissue (Figure 1D). We also analyzed the association of \textit{P4HA2} with a number of molecular markers of breast cancer, such as ERBB2 (epidermal growth factor receptor 2), ER, and PR. \textit{P4HA2} expression was significantly upregulated in ERBB2-positive breast cancers compared to ERBB2-negative breast cancers (p < 0.01) (Figure 1E), but \textit{P4HA2} levels had no significant difference between ER- or PR-positive and negative cancer (data not shown). These data confirm the previous finding that \textit{P4HA2} is associated with ERBB2 in human breast cancer cells [29]. We also found that high stage breast cancer had significantly increased \textit{P4HA2} expression (Figure 1F). These results indicate that breast cancer development and progression is accompanied by activation \textit{P4HA2}, which may contribute to collagen synthesis.

A number of genes encoding collagen proteins have been identified as prognostic markers for human breast cancer [37,38]. Since expression of \textit{P4HA2} and collagen genes is correlated in human breast cancer tissues, we asked whether \textit{P4HA2} expression is associated with clinical outcome in human breast cancer patients. Breast cancer patients were divided into two groups based on \textit{P4HA2} mRNA levels (low and high). Kaplan-Meier log rank analysis showed that patients whose tumors had high \textit{P4HA2} expression levels had a significantly shorter overall survival period (Figure 1G). Moreover, the association of \textit{P4HA2} with clinical outcome is ER status independent (see Figure 1G).

\textbf{Inhibition of \textit{P4HA2} suppresses the malignant phenotypes of breast cancer cells in 3D culture}

Increased expression of \textit{P4HA2} has been detected in many cancers [27-30], but roles of \textit{P4HA2} in cancer progression remain to be determined. To examine the function of \textit{P4HA2} in breast cancer progression, we silenced \textit{P4HA2} expression in a panel of breast cancer cell lines (HMT-3522 T4-2, MDA-MB-231, ZR-75-1, and MDA-MB-157) with two different shRNAs (shP4HA2-1 and shP4HA2-2). The \textit{P4HA2} knockdown efficiency in T4-2 cells was examined by Western blotting (Figure 2A). Similar knockdown efficiency was also obtained in MDA-MB-231 and ZR-75-1 cells (Additional file 1: Figure S2). 3D culture models have been widely used to examine the malignant mammary tissue morphogenesis [31], and the specific 3D phenotypes of breast cancer cells are associated with tumor development and cancer invasiveness. The breast cancer cell lines were classified into four groups based on their phenotypes in 3D culture: round, mass, grape-like, and stellate [39]. Malignant T4-2 cells usually form mass-like structures without apical-basal polarity in Matrigel. Disruption of polarized acinar structure is an early cellular event of tumor development, and 3D culture of T4-2 cells has been used to monitor this process. Knockdown of \textit{P4HA2} reprogrammed T4-2 cells to form polarized spheroid structures with reduced colony size (Figure 2B, C, D). Integrin α6 subunit has been used as a basal marker to detect basal polarity in mammary epithelial cells [40,41]. Immunofluorescence staining of α6 integrin showed that knockdown of \textit{P4HA2} reprogrammed the T4-2 cells to form polarized acinar-like structures (basal staining of α6 integrin), while the control cells formed the unpolarized (lateral staining of α6 integrin) and mass-like morphology (Figure 2B, C). Since disruption of polarized acinar structure is an early event during breast cancer development, these results suggest that \textit{P4HA2} contributes to the early stage of breast cancer progression. Knockdown of \textit{P4HA2} in T4-2 and ZR-75-1 cells also significantly reduced colony size in 3D culture (Figure 2B, D). To determine whether reduced colony size is due to growth inhibition, cell proliferation was examined by an Edu (5-ethyl-2′-deoxyuridine) labeling assay as described previously [42,43]. We found that Edu positive cells were significantly reduced in \textit{P4HA2}-silenced ZR-75-1 and T4-2 cells compared to control cells (Figure 2E).

MDA-MB-231 and MDA-MB-157 cells form stellate structures in 3D culture, which reflects decreased cell-cell interactions and enhanced cell invasiveness [39]. Silencing \textit{P4HA2} in MDA-MB-231 cells and MDA-MB-157 cells significantly reduced invasive branches compared to the control cells (Figure 3A, B). \textit{P4HA2}-silenced cells also had significantly reduced cell invasion in the Transwell assay (Figure 3D, E). Surprisingly, knockdown of \textit{P4HA2} had little effect on cell proliferation in MDA-MB-231 in 3D culture system (Figure 3C). Thus, \textit{P4HA2} activity in regulating cell proliferation may be cancer stage- and/or subtype-dependent. Nevertheless, reduced invasive branches in \textit{P4HA2}-silenced cells indicate that \textit{P4HA2} contributes to malignant tissue morphogenesis and cancer cell invasion in 3D culture.

1,4 dihydrophenononthrolin-4-one-3-carboxylic acid (1,4-DPCA) has been identified as a high efficiency inhibitor of prolyl-4-hydroxylase [44-46]. To determine whether \textit{P4HA2} is a potential therapeutic target for breast cancer, we treated breast cancer cells with 1,4-DPCA in 3D culture. 1,4-DPCA treatment significantly reduced the colony sizes of T4-2 and ZR-75-1 cells (Figure 4A, B). Immunofluorescence staining results showed that 1,4-DPCA-treated T4-2 cells form polarized spheroids in 3D culture (Figure 4C). Treatment with 1,4-DPCA significantly reduced invasive branches in MDA-MB-231 and MDA-MB-157 cells (Figure 4A, D). Additionally, proliferation of T4-2, ZR-75-1, MDA-MB-157 and MDA-MB-231 cells was
all inhibited by 1,4-DPCA (Figure 4E). 1,4-DPCA is not a P4HA2-specific inhibitor, and it also inhibits activity of P4HA1 and P4HA3. A recent study shows that P4HA1 and P4HA3 also contribute to breast cancer progression [33]. Therefore, this small molecule may suppress the malignant phenotypes of breast cancer cells in 3D culture by inhibiting all three P4HA isoforms. Further investigation of 1,4-DPCA activity with an in vivo mammary tumor model may lead to discovery of a new drug to inhibit breast cancer development and/or progression.

Daniele M. Gilkes et al. reported that knockdown of P4HA2 or treatment MDA-MB-231 cells with hydroxylase inhibitor DHB inhibits tumor growth in vivo, but...
little inhibitory effect on cell proliferation was detected in 2D culture assay [33]. 3D culture has been considered a better model for testing drugs and investigating cancer biology compared to 2D culture, and different drug responses between these two culture systems have recently been reported [47–49]. For example, MDA-MB-231 cells in 3D culture are more sensitive to MEK inhibition compared to cells in 2D culture [47]. Using the 3D culture model, we showed that reducing P4HA2 expression or inhibiting its activity significantly inhibited cell proliferation and suppressed the malignant phenotypes in multiple breast cancer cell lines.

Figure 3 Knockdown of P4HA2 suppresses breast cancer cells invasiveness in 3D culture. (A) Phase images of control and P4HA2-silenced MDA-MB-157 and MDA-MB-231 cells in 3D culture for 4 days. The right column shows immunofluorescence images of F-actin (red) and DAPI (blue) staining. P4HA2-silenced MDA-MB-157 and MDA-MB-231 cells formed smaller and less invasive cell clusters than control cells (scale bar, 50 μm). (B) Quantification of the invasive branches of control and P4HA2-silenced MDA-MB-157 and MDA-MB-231 cells in 3D culture by counting the branches in at least 50 colonies. Knockdown of P4HA2 reduced the invasive branches. (C) EdU-staining was used to analyze the proliferation of control and P4HA2 knockdown MDA-MB-231 cells in 3D culture. Knockdown of P4HA2 had little effect on the proliferation of MDA-MB-231 cells (**p < 0.01). (D) Transwell cell invasion assay analysis of control and P4HA2-silenced MDA-MB-231 cells (scale bar, 200 μm). (E) Quantification of invasion analysis of control and P4HA2-silenced MDA-MB-231 cells. Knockdown P4HA2 significantly inhibited cell invasion in MDA-MB-231 cells compared with control group.
Reducing P4HA2 expression or inhibiting its activity impairs deposition of collagen I and IV

Since increased collagen deposition promotes tumor progression by modulating tumor growth and invasion, we asked whether P4HA2 regulates 3D malignant phenotypes of breast cancer cells through enhancing collagen expression and deposition. The conditioned medium was collected from the control and P4HA2-silenced cells (sh-P4HA2-1). Western blot analysis showed that knockdown of P4HA2 in T4-2 cells reduced the protein levels of collagen I and IV in the conditioned medium (Figure 5A). Deposition of collagen I and IV was also reduced in the P4HA2-silenced and 1,4-DPCA-treated T4-2 cells compared to the control T4-2 cell in 3D culture (Figure 5B, C). These results indicate P4HA2 is crucial for secretion and deposition of collagen I and IV in T4-2 cells. Cancer cells produce a significant amount of ECM proteins and remodeling enzymes [13,50-53]. Dr. Massague’s group recently demonstrated that tenacin-c produced by MDA-MB-231 cells enhances dissemination and survival of tumor cells during the early steps of metastasis by generating a metastatic niche [52]. In addition, breast cancer cells express lysyl oxidases (LOXs), which promote cell invasion by increasing tissue tension and ECM rigidity [13,53]. These results indicate that the ECM microenvironment remodeled by cancer cells is...
critical for cancer progression. Thus, both inhibiting P4HA2 activity and reducing its expression are potential strategies to suppress collagen-dependent cancer progression.

**P4HA2 regulates tumor growth and metastasis in vivo**

Our data showed that silencing P4HA2 in breast cancer cells suppressed their malignant phenotypes and inhibited cell proliferation in 3D culture. We also found that P4HA2 expression is positively associated with breast cancer progression, thus we hypothesized that knockdown of P4HA2 reduces tumor growth and metastasis in vivo. To test this hypothesis, we performed xenograft tumor experiments using control and P4HA2-silenced (sh-P4HA2-1) MDA-MB-231 cells. Knockdown of P4HA2 (sh-P4HA2-1) significantly inhibited primary tumor growth in SCID mice (Figure 6A, B and C). HE staining showed that control group tumors had aggressive invasion at primary tumor margins, while tumors in the P4HA2-silenced (sh-P4HA2-1) group had no such invasion (Figure 6D). Collagen deposition in the tumors was assessed by Masson’s Trichrome staining. A significant amount of collagen fibers was detected in the tumors and at the tumor invasion margins in the control group compared to the P4HA2-silenced (sh-P4HA2-1) tumors (Figure 6E). It has been shown that increased collagen deposition and/or orientation/alignment of collagen fibers around tumors enhances cancer progression [5,10,12,14]. Cancer cell invasion usually occurs at the tumor-stromal interface with oriented collagen fibers, and aligned collagen fibers can facilitate cell migration and metastasis [5,10,15,54]. These results suggest that P4HA2 promotes tumor growth and invasion through enhancing collagen deposition.

To determine whether P4HA2 promotes breast cancer lung metastasis in vivo, the control and P4HA2-silenced (sh-P4HA2-1) MDA-MB-231 cells were injected into the tail veins of SCID mice. Lung colonization of the cancer cells was monitored by IVIS imaging. We showed that the mice injected with control cells developed lung metastasis within 6 weeks, while no metastasis was detected in the P4HA2-silenced group (Figure 6F). HE staining...
further confirmed that knockdown of P4HA2 blocked the lung colonization of MDA-MB-231 cells in SCID mice (Figure 6G).

**Conclusion**

In the present study, we show that P4HA2 is associated with expression of collagen I, III, and IV during breast cancer progression. Increased mRNA levels of P4HA2 correlate with poor prognosis in human breast cancer patients. Silencing P4HA2 or inhibiting its activity suppresses breast cancer progression by reducing tumor growth and metastasis, and this process is accompanied by reduced collagen deposition. During preparation of this manuscript, Daniele M. Gilkes *et al.* reported that hypoxia-inducible factor 1 activates the transcription of P4HA1 and P4HA2 during breast cancer development, and this activation enhances collagen fiber alignment and breast cancer progression [33,55]. Taken together, these findings indicate that P4HA2 is a promising therapeutic target to inhibit ECM-dependent breast cancer progression.

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**Figure 6** Knockdown of P4HA2 suppresses tumor growth and metastasis in vivo. (A) Control or P4HA2-silenced (sh-P4HA2-1) MDA-MB-231/Luciferase cells were injected into the mammary fat pad in SCID mice. IVIS images show representative mice from each group (n = 6). (B) Tumor growth curve shows that knockdown of P4HA2 (sh-P4HA2-1) inhibited tumor growth in SCID mice (n = 6). (C) Tumor volume formed by P4HA2 knockdown (sh-P4HA2-1) cells was significantly reduced compared with control MDA-MB-231/Luc cells. Tumor volume was obtained by quantifying IVIS images. (D) Tumor sections were stained with hematoxylin and eosin. Arrows point to invasion area at primary tumor margins (scale bar, 200 μm). (E) Masson’s Trichrome staining of tumor sections (blue, collagen fibers; black, nuclei; red, cytoplasm). A significant amount of collagen fibers (arrows) were detected in the control tumors, but not in the P4HA2-silenced (sh-P4HA2-1) tumors (scale bar, 200 μm). The right bar graph is the quantification of Masson’s Trichrome staining results. (F) Mice received a tail vein injection of control or P4HA2 knockdown (sh-P4HA2-1) MDA-MB-231/Luc cells (n = 5). IVIS image showed lung metastasis of control and P4HA2-silenced (sh-P4HA2-1) MDA-MB-231 cells in SCID mice. Lung metastasis can be detected in control but not in the knockdown P4HA2 group (*p < 0.05). (G) Lung colonization was analyzed by hematoxylin and eosin staining. Arrows point to the tumor formed in control group lung tissue (Scale bar, 500 μm).
Additional file

Additional file 1: Figure S1. Scatterplot of correlated mRNA levels between P4HA2 and (A) Col1A1, (B) Col3A1 and (C) Col4A1. Plots indicate the correlation between P4HA2 and Col1A1, Col3A1, Col4A1 expression in malignant breast tissues (n=118). The mRNA levels of P4HA2 were acquired from the Chin K’s breast cancer dataset [38]. Figure S2. Five shP4HA2s were tested in MDA-MB-231 cells by Western blot. ShP4HA2-1 and shP4HA2-2 showed best knockdown efficiency. Figure S3. Western blotting experiments shown P4HA2 knock out efficiency in shP4HA2-1 and shP4HA2-2 infected ZR-75-1 cells and MDA-MB-157 cells.

Abbreviations
P4HA2: Prolyl 4-hydroxylase subunit 2; ECM: Extracellular matrix; ER: Estrogen receptor; PR: Progesterone receptor; RRBB2: Ependymal growth factor receptor 2, 1,4-DPCA, 1,4-dihydrophenonthrolin-4-one-3-Carboxylic acid; EdU: 5-ethylthyl-2'-deoxyuridine.

Competing interests
The authors declare that they have no conflict of interest.

Authors’ contributions
GX participated in the functional assays, in vivo experiments with shP4HA2 breast cancer cells and drafted the manuscript. LD participated in the functional assays. JZ conducted Western blotting and helped draft the manuscript. RX conceived of the study, supervised its design and coordination, conducted the bioinformatics analysis and wrote the manuscript. All authors read and approved the final manuscript.

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