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Combinatorial targeting of microRNA-26b and microRNA-101 exerts a synergistic inhibition on cyclooxygenase-2 in brain metastatic triple-negative breast cancer cells

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Abstract

Purpose Extravasation of triple-negative (TN) metastatic breast cancer (BC) cells through the brain endothelium (BE) is a critical step in brain metastasis (BM). During extravasation, metastatic cells induce alteration in the inter-endothelial junctions and transmigrate through the endothelial barrier. Transmigration of metastatic cells is mediated by the upregulation of cyclooxygenase-2 (COX-2) that induces matrix metalloproteinase-1 (MMP-1) capable of degrading inter-endothelial junctional proteins. Despite their important role in BM, the molecular mechanisms upregulating COX-2 and MMP-1 in TNBC cells remain poorly understood. In this study, we unraveled a synergistic effect of a pair of micro-RNAs (miR-26b-5p and miR-101-3p) on COX-2 expression and the brain transmigration ability of BC cells.

Methods Using a gain-and-loss of function approach, we modulated levels of miR-26b-5p and miR-101-3p in two TNBC cell lines (the parental MDA-MB-231 and its brain metastatic variant MDA-MB-231-BrM2), and examined the resultant effect on COX-2/MMP-1 expression and the transmigration of cancer cells through the BE.

Results We observed that the dual inhibition of miR-26b-5p and miR-101-3p in BC cells results in higher increase of COX-2/MMP-1 expression and a higher trans-endothelial migration compared to either micro-RNA alone. The dual restoration of both micro-RNAs exerted a synergistic inhibition on COX-2/MMP-1 by targeting COX-2 and potentiated the suppression of trans-endothelial migration compared to single micro-RNA.

Conclusion These findings provide new insights on a synergism between miR-26-5p and miR-101-3p in regulating COX-2 in metastatic TNBC cells and shed light on miR-26-5p and miR-101-3p as prognostic and therapeutic targets that can be exploited to predict or prevent BM.

Keywords miR-26b · miR-101 · COX-2 · Breast cancer · Brain metastasis · Blood–brain barrier

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Introduction

Brain metastasis is an incurable end-stage of breast cancer and remains invariably a major cause of morbidity and mortality in spite of the major advances that have been made by systemic therapies [1]. The metastatic potential of BC cells as well as their response to therapy varies with the molecular subtype of the disease; each molecular subtype displays distinct patterns of metastatic spread with notable differences in survival after relapse [2]. Brain metastasis develops more frequently in triple-negative (25%-27%) and HER2 + subtypes (11%-20%), while the incidence of BM in hormone receptor-positive is much lower (8%-15%) [2, 3]. BM in patients with the TN subtype is associated with the worse median survival (3–4 months) due to several factors including the absence of approved effective targeted therapy [4]. Despite its clinical importance, the molecular mechanism of brain metastasis remains poorly understood. Understanding the biology of BM and the molecular makeup of tumor cells, particularly TNBC, more prone to disseminate to the brain, will open avenues for both the prediction of patients at high risk to develop BM and the discovery of new drug targets.

Brain metastasis is a complex process during which cancer cells detach from the primary tumors, survive in the bloodstream, cross the blood–brain barrier (BBB) and finally colonize the brain [5]. One of the critical steps in breast cancer brain metastasis (BCBM) is the passage of BC cells through the BBB, a process known as extravasation. The BBB is a unique multicellular structure located at the level of brain microvasculature at the interface between the blood and the brain. It is composed of brain endothelial cells (BEC), pericytes, astrocytes, and a basement membrane. The brain endothelial cells are considered the key element restricting the barrier permeability [6]. Unlike their homologue in the periphery, the BEC are tightly attached to each other by junctional complexes, comprising tight and adherens junctions that restrict the paracellular permeability and make the BBB one of the tightest barriers in the body. The tight junctions (TJ) completely seal the inter-endothelial cleft, while the adherens junctions (AJ) maintain contact between neighboring endothelial cells. Structurally, the TJ complexes are composed of transmembrane proteins (including occludin, claudins (1-, 3-, and -5) and junctional adhesion molecule (JAM)) connected to the actin cytoskeleton via cytoplasmic plaque proteins (including Zonula Occludens (ZO-1, -2, and -3)). Adherens junctions are located below the tight junctions and are composed of transmembrane proteins (including cadherins) linked to the cytoskeleton by cytoplasmic proteins (including catenins) [6, 7]. During extravasation, BC cells adhere to the brain endothelium, induce changes in

the inter-endothelial tight junctional proteins (including occludin, claudin-5, and ZO-1) [8, 9] and adhesion junctional proteins (including the VE-cadherin/catenin complex) [10], and then transmigrate between two neighboring cells to reach the brain. Disruption of TJs and AJs by tumor cells increases the BBB permeability and allows the paracellular passage of cancer cells through the endothelial barrier [5]. However, the molecular mechanism by which BC cells acquire their migrative properties allowing them to induce alteration in the inter-endothelial junctions and cross the BBB remains poorly understood.

The cyclooxygenase-2 (COX-2)/matrix metalloproteinase-1 (MMP-1) pathway was shown to play a critical role in the extravasation of TNBC cells through the BBB. TNBC patients with high COX-2 and MMP-1 were found to have the greatest risk of brain metastasis [11]. Mechanistic studies revealed that brain metastatic cells express high levels of COX-2 and MMP-1 compared to primary tumor cells [12] and that overexpression of COX-2 promotes brain metastasis by inducing MMP-1 expression [11] which in turn degrades the inter-endothelial junctions, enhances the BBB permeability, and allows the transmigration of metastatic cells through the brain endothelium [11, 13]. Despite their important role in brain metastasis, the molecular mechanisms leading to upregulation of COX-2/MMP-1 in TN brain metastatic cells remain poorly understood. A better understanding of these molecular mechanisms will open avenues for better prognostic and therapeutic strategies.

In recent years, micro-RNAs have emerged as important regulators of gene expression. Micro-RNAs are a class of non-coding RNA that negatively regulate gene expression at mRNA and protein level [14]. Dysregulation of micro-RNAs is involved in tumorigenesis, metastasis, and drug resistance [15, 16]. In case of BCBM metastasis, brain metastatic tumors were shown to express distinct micro-RNAs signature compared to primary tumors, suggesting a potential role of micro-RNAs dysregulation in the development of brain metastases. In a previous study, we showed that miR-101-3p is downregulated in TN brain metastatic breast cancer cells which upregulates COX-2 and increases the transmigration of metastatic cells through the brain endothelium. Restoring miR-101-3p in brain metastatic cells reduces tumor cells migration by directly targeting and reducing COX-2 expression, however, without total suppression of cancer cells transmigration (11).

Mechanistic studies on micro-RNAs action revealed that the 3'UTR of genes are frequently targeted by multiple micro-RNAs and that simultaneous modulation of miRNA pairs can exert synergistic effects on the target genes leading to a potentiation of their biological effects compared to single micro-RNAs [17–22]. Interestingly, a pair of micro-RNAs (miR-101-3p and miR-26b-5p), shown to target COX-2 (PTGS2) gene in multiple types

of cancers [23–28], are both downregulated in brain metastatic tumors compared to primary tumors [29]. Based on these data, we hypothesized that a combinatorial targeting of miR-101-3p and miR-26b-5p might exert a synergistic repression on COX-2 gene in TN metastatic cancer cells and suppress their transmigration through the brain endothelium to a greater extent than single micro-RNA. To test this hypothesis, we compared the anti-transmigrative effect of miR-101 and miR-26b alone and in combination in *in vitro* models of brain endothelial barrier.

Materials and methods

Animals

Adult female mice ([C57BL/6 J], 8–10 weeks of age) were purchased from Jackson Laboratory (USA) and maintained at the accredited animal facility in the University of Sharjah (UOS). All experiments were approved by the UOS animal care and use committee and conducted according to the UOS directives for animal care.

Cell lines

The non-metastatic (ER⁺, PR[±], HER2⁻) MCF-7 and the triple-negative MDA-MB-231 human breast carcinoma cell lines were obtained from American Type Culture Collection (Rockville, MD). The TN brain metastatic MDA-MB-231-BrM2 cell variant and the parental MDA-MB-231-TGL cells with low brain metastatic propensity were obtained from Dr Joan Massagué (Memorial Sloan-Kettering Cancer Center, New York, USA). The MDA-MB-231-BrM2 cells have an increased propensity to form brain metastases and were isolated from the brain of nude mice following multiple injections of the parental MDA-MB-231-TGL into the internal carotid artery [12]. Cells were cultured in Dulbecco's Modified Eagle Medium (DMEM) supplemented with 10% FBS, 1X L-Glutamine, and antibiotics (1000 U Penicillin/1000 U Streptomycin). The human cerebral microvascular endothelial cell line (hCMEC/D3) was obtained from Cedarlane (TebuBio, France) and cultured in EndoGROTM-MV Complete Medium (cat# SCME004, EMD Millipore, USA) supplemented with 1 ng/mL FGF-2 (cat# GF003, EMD Millipore) and antibiotics. hCMEC/D3 were cultured on flasks pre-coated with thin collagen I (cat# 08–115, EMD Millipore) and fibronectin (cat# F1141, Sigma-Aldrich) coating. All cells were cultured in a humidified incubator with 5% CO₂ at 37 °C and regularly tested for mycoplasma contamination and phenotypic changes.

Establishment of *in vitro* brain endothelial barriers from hCMEC/D3 cell line

To establish the endothelial barriers from hCMEC/D3 cells, (5×10^4) cells were seeded on the upper side of a collagen and fibronectin-coated polyester Transwell membrane (Costar, pore size 8 μm ; growth area 1.12 cm^2) in endothelial basal medium (EndoGROTM-MV Complete Medium). Under these experimental conditions, brain endothelial cells form a confluent monolayer within 72 h with the highest transendothelial electrical resistance (TEER) values between days 3 and 7 [30, 31].

Establishment of *in vitro* brain endothelial barriers from primary cells

Primary brain endothelial cells were isolated from adult female mice ([C57BL/6 J], 8–10 weeks of age) as previously described [32, 33]. The purity of isolated brain endothelial cells was examined by measuring the gene expression of three cell markers: CD31 or pecam-1 (marker of endothelial cells), glial fibrillary acidic protein or gfap (marker of glial cells), and α -actin or acta2 (marker of pericytes). The following mouse primers from Macrogen (South Korea) were used: Pecam1 Mouse qPCR Primer Pair (NM_008816), F: CCAAAGCCAGTAGCATCATGGTC, R: GGATGGTGAAGTTGGCTACAGG; Gfap Mouse qPCR Primer Pair (NM_010277), F: CACCTACAGGAAATTGCTGGAGG, R: CCACGATGTTCTCTTGAGGTG; Acta2 Mouse qPCR Primer Pair (NM_007392), F: TGCTGACAGAGG CACCACTGAA, R: CAGTTGTACGTCCAGAGGCATAG. To establish the endothelial barriers from primary cells, (5×10^4) of isolated BEC cells were seeded on the upper side of a collagen and fibronectin-coated polyester transwell membrane (Costar, pore size 8 μm ; growth area 1.12 cm^2) in endothelial basal medium (EGM-2MV Endothelial Med BulletKit, cat # CC-3202, Lonza, Switzerland) containing 0.1% human recombinant epidermal growth factor (hEGF), 0.04% hydrocortisone, 0.1% human recombinant insulin-like growth factor, 0.1% ascorbic acid, 0.1% gentamicin, 0.1% amphotericin-BN, and 5% fetal bovine serum). Under these experimental conditions, brain endothelial cells form a confluent monolayer within 12 days.

Transendothelial electric resistance (TEER) measurement

TEER of endothelial barriers was measured with an Endohm 12 chamber and an Endohmeter EVOMX (World Precision Instruments). Media were refreshed and inserts were kept at room temperature for 20 min prior to TEER measurement to exclude interference of temperature and media. Background of the coated transwell membrane and medium

was subtracted from each reading. The TEER values were expressed as $\Omega \cdot \text{cm}^2$ (surface area of the Transwell inserts).

Cell transfection

To introduce miR-26b-5p and miR-101-3p, cells were transfected with 2.5 nM of miR-26b-5p mimic (Syn-hsa-miR-26b-5p miScript miRNA Mimic, Cat# 219,600, MSY0000083: 5'UUCAAGUAAUUCAGGAUAGGU, Qiagen, Germany), miR-101-3p mimic (Syn-hsa-miR-101-3p miScript miRNA Mimic, Cat# 219,600, MSY0000099: 5'UACAGUACUGUGAUAACUGAA, Qiagen), or its (scrambled) control (AllStars Negative Control, cat# SI03650318, Qiagen) using HiPerFect transfection agent (Qiagen) in culture media for 48 h. For inhibition studies, 15 nM of miR-26b-5p inhibitor (Anti-hsa-miR-26b-5p miScript miRNA Inhibitor, cat# 219,300, MIN0000083), miR-101-3p inhibitor (Anti-hsa-miR-101-3p miScript miRNA Inhibitor, cat# 219,300, MIN0000099), or its negative control (miScript Inhibitor Negative Control, cat# 1,027,271, Qiagen) were transfected using HiPerFect transfection agent in culture media for 48 h. Down- and upregulation of miR-26b-5p and miR-101-3p were assessed by qPCR.

Cell viability assay

MDA-MB-231 cells (5×10^3 cell/well) were seeded in 96-well plate. After 24 h, cells were treated with 20 μM of celecoxib (ab141988, abcam). Six hours later, cells were transfected with miR-26b-5p and miR-101-3p mimics or inhibitors, and cell viability was measured by MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) assay after 48 h; 100 μL of MTT (5 mg/mL) was added and cells were incubated for 4 h. Supernatant was then removed and 50 μL of DMSO (dimethyl sulfoxide) was added to each well. The absorbance was measured at 540 nm wavelength using Multiskan Go microplate spectrophotometer (Thermo Fisher, Waltham, MA, USA).

Quantitative real-time PCR

Total RNAs were extracted from cells using the miRNeasy Micro Kit (cat# 217,084, Qiagen, Germany) following the manufacturer's instructions. micro-RNAs were reversed transcribed into cDNA using the miScript II RT Kit (cat# 218,161) from Qiagen. mRNAs were reversed transcribed using the RT² First Strand Kit (Cat#, 330,401, Qiagen). The miScript SYBR Green PCR Kit (Cat# 218,075) was used to detect miRNA expression, and the RT² SYBR Green ROX qPCR Mastermix (cat# 330,522) was used for mRNA expression according to the manufacturer's protocol. PCR amplifications were performed on the "Applied Biosystems® StepOne™ Real-Time PCR System." The specific primers

from Qiagen were as follows: Hs_miR-26b_1 miScript Primer Assay (Cat# MS00003234), Hs_miR-101_3 miScript Primer Assay (Cat# MS8300072), RT² qPCR Primer Assay for Human PTGS2 (cat# PPH01136F), and RT² qPCR Primer Assay for Human MMP1 (cat# PPH00120B-200). Hs_RNU6-2_11 miScript Primer Assay (cat# MS00033740) and RT² qPCR Primer Assay for Human GAPDH (Cat# PPH72843A) were used as internal control for miRNA and mRNA, respectively. The specificity of each reaction was assessed by melting curve analysis. The relative gene expression was determined using the $2^{-\Delta\text{Ct}}$ and $2^{-\Delta\Delta\text{Ct}}$ methods [34].

Trans-endothelial migration assay

Trans-endothelial migration assay was performed as previously described [35]. hCMEC/D3 and primary brain endothelial cells were plated on the upper side of an 8 μm pore size insert, pre-coated with collagen and fibronectin. On the day of the experiment, TEER was monitored to assess the monolayer tightness. TEER values of control endothelial monolayers hCMEC/D3 were around 70 $\Omega \text{ cm}^2$ between days 5 and 7 of culture. Chambers with a TEER lower than 65 $\Omega \text{ cm}^2$ were discarded. TEER values of control primary brain endothelial cells were around 500 $\Omega \text{ cm}^2$ on day 12 of culture. For transmigration assay, 5×10^4 transfected or control cells were pre-labeled with 5 μM of CellTracker™ Green CMFDA fluorescent dye (cat# C2925, Thermo Fisher Scientific, USA) and then added gently to the apical (upper) side of the transwell. Four hours later, non-adherent cells were washed with PBS. After 24 h, TEER was measured again and non-migrated cells remaining in the upper chamber were carefully removed with a cotton swab. Cancer cells that have transmigrated through the brain endothelium to the basal compartment were fixed with 4% PFA for 20 min for imaging and counting under Olympus BX43 fluorescence microscope, or lysed with RIPA 1X buffer for fluorescence quantification [28, 36]. The fluorescence signal in the lower compartment was quantified in Varioskan Flash microplate reader (Thermo Fisher scientific) at wavelength excitation/emission: 492/517 nm.

Cell sorting and western blot

Protein expression of COX-2 and MMP-1 in cancer cells as well as the inter-endothelial junctions (Claudin-5, VE-cadherin, ZO-1, and β -catenin) in brain endothelial cells was assessed by western blot following standard procedures and as previously described [28]. To measure the expression of inter-endothelial junctions in BECs, MDA-MB-231-TGL and MDA-MB-231-Brm2 cells, which are stably transfected with a lentivirus expressing a triple fusion reporter (TGL) encoding herpes simplex virus thymidine kinase

1, green fluorescent protein (GFP) and firefly luciferase [12], were co-cultured with hCMEC/D3 cells. After 24 h of co-culture, cells were collected with accutase (Corning, USA) and sorted with the BD FACSAria™ III cell sorter (BD Biosciences, USA) using the 488 nm laser to separate GFP-expressing cancer cells from unlabeled hCMEC/D3. The following primary antibodies from Abcam were used: Anti-COX-2 (ab15191) 1/1000, Anti-MMP-1 (ab38929) 1/5000, Anti-Claudin-5 (ab15106) 1/500, Anti-VE Cadherin (ab33168) 1/1000, Anti-ZO-1 (ab216880) 1/1000, and Anti- β -catenin (ab32572) 1/3500. Anti-GAPDH (ab9485) 1/7500 was used as loading and transfer control.

Immunofluorescence

Cells were fixed with 4% paraformaldehyde for 10 min and permeabilized with 0.15% Triton™ X-100. After blocking with 2% BSA for 1 h, cells were incubated overnight at 4 °C with anti-COX-2 primary antibody (ab15191) 1/500. Cells were then washed and incubated for 1 h with secondary antibody donkey F(ab')₂ anti-rabbit IgG H&L (Phycocerythrin) (ab7007) 1/200. Samples were then mounted in ProLong® Gold Antifade Reagent with DAPI (cat# 8961, Cell Signaling, USA) and examined with Olympus BX43 fluorescence microscope.

Enzyme-linked immunosorbant assay (ELISA)

Culture supernatants of transfected BC cells were collected and frozen at – 20 °C. For quantification of Prostaglandin E₂ (PGE₂) and MMP-1 levels, the Prostaglandin E₂ Parameter Assay Kit (cat# KGE004B, R&D Systems) and the Human Total MMP-1 DuoSet (cat# DY901B, R&D Systems) were used, respectively, according to manufacturer's instructions. The detection limits for PGE₂ were 39 pg/mL and for MMP1 were 62.5 pg/mL. Signal saturation was observed at concentrations > 2500 pg/mL for PGE₂ and > 4000 pg/mL for MMP-1. The optical density was measured using a Multiskan™ GO Microplate Spectrophotometer (Thermo Scientific, USA) at a wavelength of 450 nm. PGE₂ and MMP-1 concentrations were then calculated by generating a four parameter logistic (4-PL) curve-fit.

Dual luciferase reporter assay

For miR-26b target validation, the wild-type (WT) putative binding site of miR-26b-5p in the 3'UTR of COX-2 predicted by TargetScan (version 7.2) (position 257–264 of COX-2 3' UTR), and the mutant (Mut) 3'UTR of COX-2 with the seed region deleted were cloned into pmirGLO Dual-Luciferase miRNA Target Expression Vector (pmirGLO-empty, Promega, USA) downstream of the firefly luciferase gene (XbaI sites) to obtain Luc Reporter Constructs.

The following primers were used for cloning: 26_PTGS2_WT_F: 5'CTAGACCAGAGAGAAATGAGTTTTTGAC GTC TTTTACTTGAATTTCAACTTATATTATAAGAACGA AAGTAA3'; 26_PTGS2_WT_R 5'CTAGTT ACTTTCGTT CTTATAATATAAGTTGAAATTCAAGTAAAAAGACGT CAAAACCTCATTCTCTCTGGT3'(WT). The mutant primers were: 26_PTGS2_MUT_F: 5'CTAGACCAGAGAGAA ATGAGTTTTGACGCTTTT TTTTCAACTTATATTATA AGAACGAAAGTAA3'; 26_PTGS2_MUT_R 5'CTAGTT ACTTTCGTTCTTATAA TATAAGTTGAAAAAAGA CGTCAAAACCTCATTCTCTCTGGT3' (MUT). For miR-101-3p target validation, the wild-type (WT) putative binding site of miR-101-3p in the 3'UTR of COX-2 predicted by TargetScan 7.2 (position 1738–1744 of PTGS2 3' UTR) and the mutant (MUT) 3'UTR of COX-2 with the seed region deleted were cloned into pmirGLO Dual-Luciferase miRNA Target Expression Vector as previously described [28]. The following primers were used:

101_PTGS2_WT_F 5'CTAGCCACATCTCATTGTGAC TGACATTTAATGGTACTGTATATTACTTAATTTATT GAAGATTATTATTTA3'; 101_PTGS2_WT_R 5'CTAGTA AATAATAATCTTCAATAAATTAAGTAATATACAGTA CCATTAAATGTCAGTGACAATGAGATGTGG3';

101_PTGS2_MUT_F 5'CTAGCCACATCTCATTGT CACTGACATTTAATGGATTACTTAATTTATTGAAGA TTATTATTTA3'; 101_PTGS2_MUT_R 5'CTAGTAAAT AATAATCTTCAATAAATTAAGTAATCCATTAAATGT CAGTGACAATGAGATGTGG3'.

MDA-MB-231 cells (ATCC) were plated at a density of 10⁴ cells/well in a 96-well plate and co-transfected with pmirGLO-Mut (50 ng), pmirGLO-WT (50 ng), miR-26 mimic (10 nM), and/or miR-101 mimic (10 nM) or negative scrambled control depending on treatments, and following Lipofectamine 3000 reagent protocol. 24 h after transfection, cell lysates were harvested and used to sequentially measure the Firefly and Renilla luciferase activities using the Dual-Luciferase® Reporter Assay System analysis (cat# E2940, Promega, USA) according to the manufacturer's instructions. Firefly activities were normalized with Renilla luciferase.

Statistical analysis

Data are expressed as mean \pm SD (standard deviation) from three or four independent experiments with (2–3) replicates each. Unpaired Student's *t* test was used for statistical comparison between two groups, and one-way ANOVA followed by Bonferroni post hoc test was used for comparison among multiple groups. Differences were considered to be statistically significant at probability levels of $p < 0.05$ (*), $p < 0.01$ (**), and $p < 0.001$ (***). Calculations and figures were generated using the statistical software GraphPad Prism (version 8.2.0).

To identify whether the combination had a synergistic effect, the Combination Index (CI) was calculated according to the Chou-Talalay method [37–40]. Cells were treated with a combination of miR-26b and miR-101 mimics using the method of constant ratio drug combination proposed by Chou and Talalay. The two mimics were used at the concentration of 1.25, 2.5, and 5 nM each. Then using gene expression data from real-time PCR, the CI values for each mimic pair were calculated using the CompuSyn algorithm. $CI < 1$ denotes a synergism; $CI = 1$ denotes an addition, and $CI > 1$ denotes an antagonism.

Results

Breast cancer cells with high brain metastatic propensity express lower levels of miR-26b-5p and miR-101-3p compared to non-metastatic cells

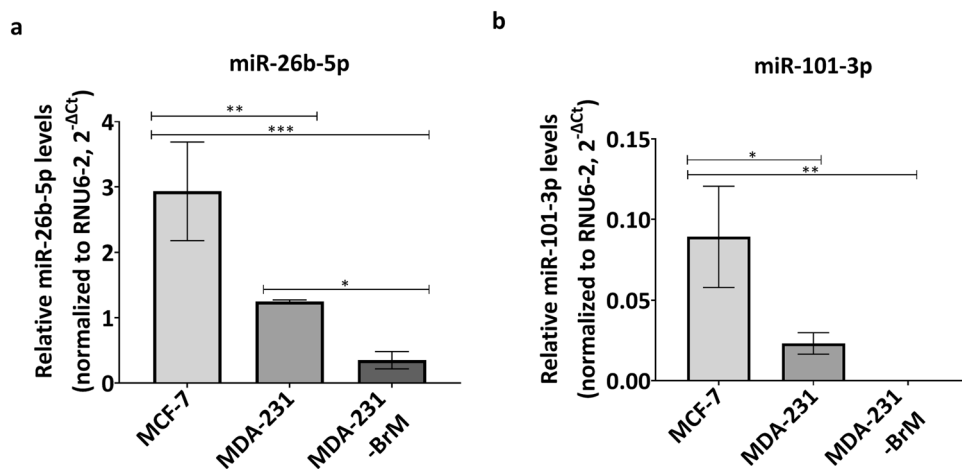
Previous studies showed that two micro-RNAs (miR-26b-5p and miR-101-3p), known to target the 3'UTR of COX-2, are significantly downregulated in brain metastatic tumors compared to primary tumors in patients samples [29]. We measured the expression levels of miR-26b-5p and miR-101-3p by real-time PCR in three breast cancer cell lines with different brain metastatic propensities: MCF-7 (ER⁺, PR[±], HER2⁻, non-metastatic), MDA-MB-231-TGL (MDA-231 for brevity, TN, metastatic), and MDA-MB-231-BrM2 (MDA-231-BrM for brevity, TN, brain metastatic). Our results showed that levels of both micro-RNAs, miR-26b-5p and miR-101-3p, were lower in TN MDA-231-BrM cells compared to parental TN MDA-231 (miR-26: fold change = 0.28, $p = 0.0475$; miR-101: fold change = 0.01) and non-metastatic (ER⁺, PR[±], HER2⁻) MCF-7 (miR-26: fold change = 0.13, $p = 0.0004$; miR-101: fold change = 0.002, $p = 0.0011$) (Fig. 1). These results are in accordance with previous results obtained from patients' samples showing that miR-101-3p and miR-26b-5p

are significantly downregulated in brain metastatic tumors compared to primary tumors [29], which suggests that loss of these micro-RNAs could exert a potential role in brain metastasis.

MiR-26b-5p and miR-101-3p cooperatively target the 3'-UTR of COX-2 mRNA in breast cancer cells

The 3'UTR of COX-2 (PTGS2) mRNA was shown to be targeted by miR-26b-5p or miR-101-3p in multiple types of cancer [23–28]. The putative target sequences of miR-26b-5p and miR-101-3p in the 3'UTR of COX-2 mRNA predicted by TargetScan 7.2 are shown in Fig. 2a and b. We conducted luciferase reporter assays in MDA-MB-231 cells to confirm the previous observations. miR-26b-5p or miR-101-3p mimics were co-transfected with luciferase constructs containing the putative (wild-type WT) or mutated (Mut) binding site of miR-26b-5p (pmirGlo-PTGS2 (seed-miR-26)-3'UTR) and miR-101-3p (pmirGlo-PTGS2 (seed-miR-101)-3'UTR) in the 3'UTR of COX-2. For the luciferase assay, we used MDA-MB-231 cells (from ATCC) with no endogenous luciferase activity; the two other cell lines, MDA-MB-231-TGL and MDA-MB-231-BrM2, could not be used as they are stably transduced with a lentivirus expressing a triple fusion reporter (TGL) encoding firefly luciferase. The luciferase assays showed that mimics of both micro-RNAs significantly reduced the luciferase activity in cells transfected with the pmirGlo-PTGS2-3'UTR-WT construct compared to control cells (miR-26b-5p: 43.14% reduction, $p < 0.0001$; miR-101-3p: 34.07% reduction, $p < 0.0027$), whereas the two mimics had no inhibitory effects on luciferase activity in cells transfected with construct containing mutated seed sequences (Fig. 2a and b). These data confirmed results obtained from previous studies showing that miR-26b-5p and miR-101-3p directly target COX-2 mRNA which suggests a potential role of these micro-RNAs in regulating COX-2 expression in metastatic breast cancer cells.

Fig. 1 miR-26b-5p and miR-101-3p levels are reduced in brain metastatic breast cancer cells compared to non-metastatic cells. Expression levels of miR-26b-5p (a) and miR-101-3p (b) in breast cancer cells with different brain metastatic propensities measured by real-time PCR. The small nuclear RNA (RNU6-2) was used as an internal standard. Data represent mean \pm SD from three independent experiments. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$



Interestingly, the combination of miR-26-5p and miR-101-3p mimics results in greater reduction of the luciferase activity in cells co-transfected with both constructs containing the putative seed sequences of miR-26b-5p and miR-101-3p and this by around 77.64% ($p < 0.0001$) compared to control treatment (Fig. 2c). In comparison to single transfections, the combination (miR-26-5p and miR-101-3p) reduced the luciferase signal by around 35.25% ($p < 0.0001$) compared to miR-26b-5p and by around 45.30% ($p < 0.0001$) compared to miR-101-3p (Fig. 2c). These results suggested that the combination (miR-26-5p and miR-101-3p) tend to synergistically regulate the COX-2 expression in BC cells.

The dual loss of miR-26 and miR-101 in TN breast cancer cells produces a higher increase of COX-2/MMP-1 expression compared to either micro-RNA alone

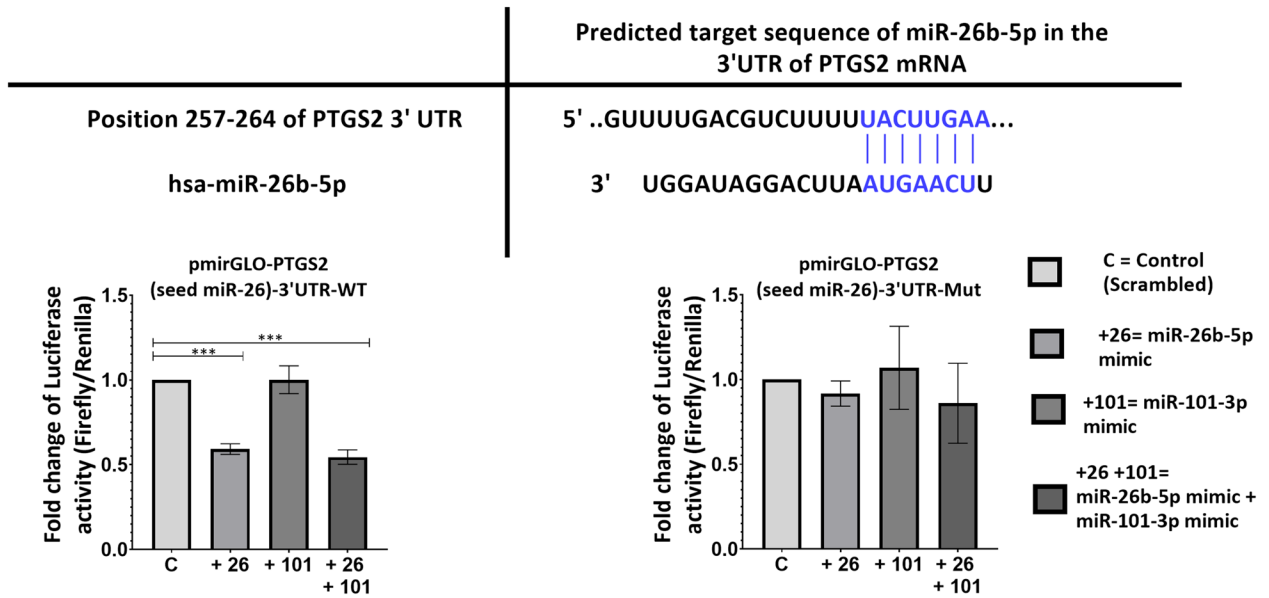
We next investigated the effect of a dual inhibition of miR-26b-5p and miR-101-3p on COX-2 expression in TNBC cells. MDA-MB-231-TGL (MDA-231) cells were transfected with miR-26b-5p inhibitor (15 nM), miR-101-3p inhibitor (15 nM), or both inhibitors (15 nM each) simultaneously. Results are represented in Fig. 3 and show that treatment with miR-26b-5p inhibitor alone significantly reduced miR-26b-5p expression ($p < 0.001$) without affecting miR-101-3p expression, while the co-transfection with both inhibitors significantly reduced expression of miR-26b-5p and miR-101-3p (Fig. 3a). Same observations were made with miR-101-3p inhibitor alone or combined with miR-26b-5p inhibitor ($p < 0.001$) (Fig. 3b). The dual inhibition of miR-26b-5p or miR-101-3p in MDA-231 cells expressing low levels of COX-2 produced a significant increase of COX-2 gene expression (fold change = 4.10, $p = 0.0003$) that is higher than each micro-RNA alone (Fig. 3c). This higher increase was also observed at the level of protein expression, measured by western blot and immunofluorescent staining where the dual inhibition of both micro-RNAs increased COX-2 expression [fold change = 5.14-fold ($p = 0.0027$) and 5.60 ($p = 0.0041$)] compared to miR-26b-5p or miR-101-3p alone, respectively (Fig. 3d and e). As COX-2 is known to mediate cancer cells transmigration by inducing MMP-1 expression, we assessed expression of MMP-1. Our results showed that the dual inhibition of both micro-RNAs increased MMP-1 protein expression by 3.25-fold and 3.30-fold compared to miR-26b-5p or miR-101-3p alone, respectively (Fig. 3f). Rescue experiments were performed by treating MDA-231 cells with celecoxib (20 μ M), a selective COX-2 inhibitor. Six hours later, cancer cells were transfected with miR-26b-5p inhibitor alone, miR-101-3p inhibitor alone, or with both inhibitors simultaneously, and expression of COX-2 and MMP-1 was assessed 48 h later. Results showed that treatment with celecoxib rescued the effect of micro-RNAs

inhibition on COX-2 and MMP-1 expression (Supplementary Figures S1). In addition, we assessed the effect of micro-RNAs inhibition and celecoxib treatment on cell viability as previous studies reported that celecoxib reduces viability of BC cells at high doses (40 μ M) [41]. Our results represented in (Supplementary Figures S2) showed that treatment of cancer cells with celecoxib at the dose of 20 μ M and miR-26b-5p and miR-101-3p inhibitors (15 nM) does not significantly affect cell viability. Taken together, these results showed that miR-26 and miR-101 cooperatively target COX-2 in breast cancer cells, and the dual inhibition of both micro-RNAs produces a higher increase of COX-2/MMP-1 expression compared to either micro-RNA alone.

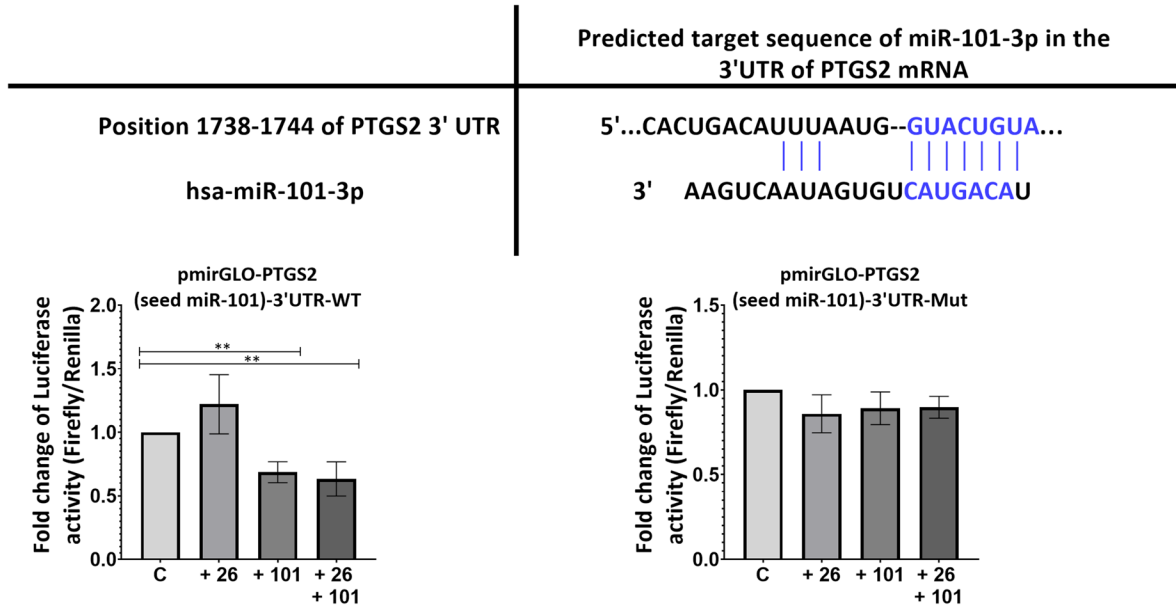
The dual inhibition of miR-26 and miR-101 in TN breast cancer cells results in greater increase of the transmigration of BC cells through the brain endothelium compared to either micro-RNA alone.

We next evaluated the effect of miR-26b-5p and miR-101-3p dual inhibition on the transmigration of MDA-231 cells through the brain endothelium. Two models of *in vitro* brain endothelial barriers were established using primary brain endothelial cells or the immortalized human hCMEC/D3 cell line known to retain the morphological characteristics of primary BEC and to form a tight monolayer [30, 31]. Results are represented in Fig. 4 and show that mono-transfection of BC cells with 15 nM of miR-26b-5p or miR-101-3p inhibitors alone do not cause a significant increase in the transmigration of BC cells through the primary brain endothelium compared to negative control, while the dual inhibition of both micro-RNAs resulted in significantly higher increase of trans-endothelial migration by 2.71-fold ($p = 0.003$) compared to negative control and by 2.12-fold ($p = 0.0074$) and 2.06-fold ($p = 0.0129$) compared to miR-26b-5p and miR-101-3p, respectively (Fig. 4a). Using the *in vitro* model of brain endothelial barriers established with hCMEC/D3 cells, results showed that trans-endothelial migration of BC cells was increased by 2.51-fold ($p = 0.0049$) and 2.93-fold ($p = 0.0092$) when MDA-231 cells were mono-transfected with miR-26b-5p or miR-101-3p alone, respectively, while co-transfection with both micro-RNAs further increased transmigration of cancer cells by 6.95 times ($p < 0.001$) compared to control and by 2.77 times ($p = 0.0006$) and 2.38 times ($p = 0.0019$) compared to either miR-26b-5p or miR-101-3p, respectively (Fig. 4b and c). Rescue experiments showed that treatment with celecoxib rescued the effect of micro-RNAs inhibition on the trans-endothelial migration (Supplementary Figures S3). Taken together, these results show that the dual inhibition of miR-26b-5p and miR-101-3p produces a higher increase in COX-2 expression which results in greater transmigration of BC cells through the

a



b



c

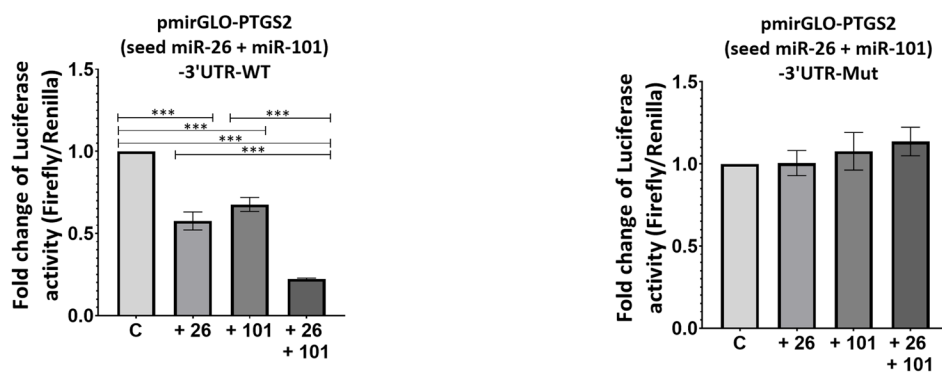


Fig. 2 miR-26b-5p and miR-101-3p cooperatively target the 3'-UTR of COX-2 mRNA. **a** Binding site of miR-26b-5p with the 3'UTR of COX-2 (PTGS2) as predicted by TargetScan (version 7.2). Complementary sequence is represented in blue. Luciferase reporter assay in MDA-MB-231 cells co-transfected with miR-26b-5p mimic (or control) and constructs carrying PTGS2 3'UTR luciferase reporter [pmirGLO-PTGS2 (Seed miR-26b)-3'UTR-WT] or mutant of miR-26b-5p binding site [pmirGLO-PTGS2 (seed miR-26b)-3'UTR-Mut]. **b** Binding site of miR-101-3p with the 3'UTR of COX-2 (PTGS2) as predicted by TargetScan 7.2. Complementary sequence is represented in blue. Luciferase reporter assay in MDA-MB-231 cells co-transfected with miR-101-3p mimic (or control) and constructs carrying PTGS2 3'UTR luciferase reporter [pmirGLO-PTGS2 (Seed miR-101)-3'UTR-WT] or mutant of miR-101-3p binding site [pmirGLO-PTGS2 (seed miR-101)-3'UTR-Mut]. **c** Luciferase reporter assay in MDA-MB-231 cells co-transfected with miR-26b-5p and miR-101-3p mimics (or negative control) and constructs carrying PTGS2 3'UTR luciferase reporter [pmirGLO-PTGS2 (Seed miR-26b)-3'UTR-WT and pmirGLO-PTGS2 (Seed miR-101)-3'UTR-WT] or deletion mutant of miR-26b-5p and miR-101-3p binding sites [pmirGLO-PTGS2 (seed miR-26b)-3'UTR-MUT and pmirGLO-PTGS2 (seed miR-101)-3'UTR-Mut]. Data represent mean \pm SD from at least three independent experiments. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$

brain endothelium compared to either micro-RNA alone. We next investigated the effect of miR-26b-5p and miR-101-3p inhibition on the integrity of brain endothelium. Our results showed that co-transfection of MDA-231 cells with both micro-RNA inhibitors results in greater reduction of the TEER compared to each micro-RNA alone (Fig. 5a and b). In primary BECs, co-culture with MDA-231 cells reduced TEER by around 25.42% ($p = 0.0014$) compared to monoculture, while co-culture with MDA-231 cells transfected with either miR-26-5p or miR-101-3p inhibitor alone reduced the TEER by 51.96% ($p = 0.0009$) and 49.48% ($p = 0.002$) compared to control, respectively. Co-culture with MDA-231 cells transfected with both inhibitors further reduced the TEER by 82.58% ($p < 0.001$) compared to control and by 30.62% ($p = 0.0086$) and 33.10% ($p = 0.0106$) compared to cells mono-transfected with either miR-26-5p or miR-101-3p inhibitors, respectively (Fig. 5a). Similar trend was observed using the in vitro model of BECs established with hCMEC/D3 cell line (Fig. 5b). To confirm the effect of both micro-RNAs on the barrier integrity, we evaluated the protein expression of four junctional proteins (claudin-5, VE-cadherin, β -catenin, and ZO-1). Our results showed that these proteins were further reduced when BC cells were transfected with both micro-RNA compared to single inhibition (Fig. 5c-f). Taken together, our data showed that the dual inhibition of miR-26b-5p and miR-101-3p results in greater induction of COX-2/MMP-1 and enhances the transmigration of BC cells through the brain endothelium by exacerbating the disruption of inter-endothelial junctions and further reducing the barrier integrity compared to each micro-RNA alone.

Combination of miR-26b-5p and miR-101-3p exerts a synergistic repressive effect on COX-2 expression in TN brain metastatic BC cells

To determine whether the combination of miR-26b-5p and miR-101-3p exerts a synergistic effect in terms of repressing COX-2 expression, MDA-231-BrM2 cells were first transfected with 1.25 nM, 2.5 nM, or 5 nM of miR-26b-5p mimic alone, miR-101-3p mimic alone, or a combination of miR-26b-5p mimic and miR-101-3p mimic at the same concentrations. Forty-eight hours later, mRNA levels of COX-2 were examined by real-time PCR, and the Combination Index (CI) was calculated according to the Chou-Talalay method [37–40]. Results showed that the different combinations of micro-RNAs at lower concentrations exert a greater repression on COX-2 expression compared to single micro-RNA at higher concentrations, with the greater reduction observed at the combination of 2.5 nM (Supplementary figures S4). Regarding the combination index, our results showed that the CI using 1.25 or 2.5 nM of each mimic are 0.59 and 0.81, respectively, indicating a synergism. However, CI using 5 nM of each mimic is 1.63 indicating an antagonism, which is probably due to an undesirable off-target effect caused by the usage of higher doses of mimics. Indeed, several previous studies showed that transfection of miRNA mimics at high concentrations causes non-specific alterations in gene expression [41, 42]. Based on these first results, the dose 2.5 nM that showed the highest reduction in COX-2 expression was used for the rest of the study to confirm the synergistic effect of miR-26b and miR-101 on COX-2 expression. Indeed, using the concentration of 2.5 nM, transfection with miR-26b-5p mimic alone significantly increased miR-26b-5p expression ($p = 0.0003$) without affecting miR-101-3p expression, while the co-transfection with both mimics significantly increased expression of both miR-26b-5p and miR-101-3p ($p < 0.001$) (Fig. 6a). Same observations were made with miR-101-3p mimic alone or combined with miR-26b-5p mimic (Fig. 6b). Compared to control cells, COX-2 (PTGS2) gene expression was reduced by 3.63-fold in cells transfected with miR-26b-5p alone (fold change = 0.27, $p < 0.001$), by 3.44 times in cells transfected with miR-101-3p mimic (fold change = 0.29, $p < 0.001$), and by 27-fold in cells transfected with both miR-26b-5p and miR-101-3p mimics (fold change = 0.035, $p < 0.001$). The dual inhibition of both micro-RNAs reduced COX-2 gene expression by 7.68-fold (fold change = 0.13, $p = 0.0005$) compared to single mimics (Fig. 6c). Similarly, COX-2 protein levels were more reduced in cells transfected with both mimics (fold change = 0.03, $p = 0.051$) than in cells transfected with either miR-26b-5p (fold change = 0.25, $p = 0.0128$) or miR-101-3p (fold change = 0.25, $p = 0.136$) alone compared to control (Fig. 6d and e). We also measured levels of PGE2 released in culture media. Levels of

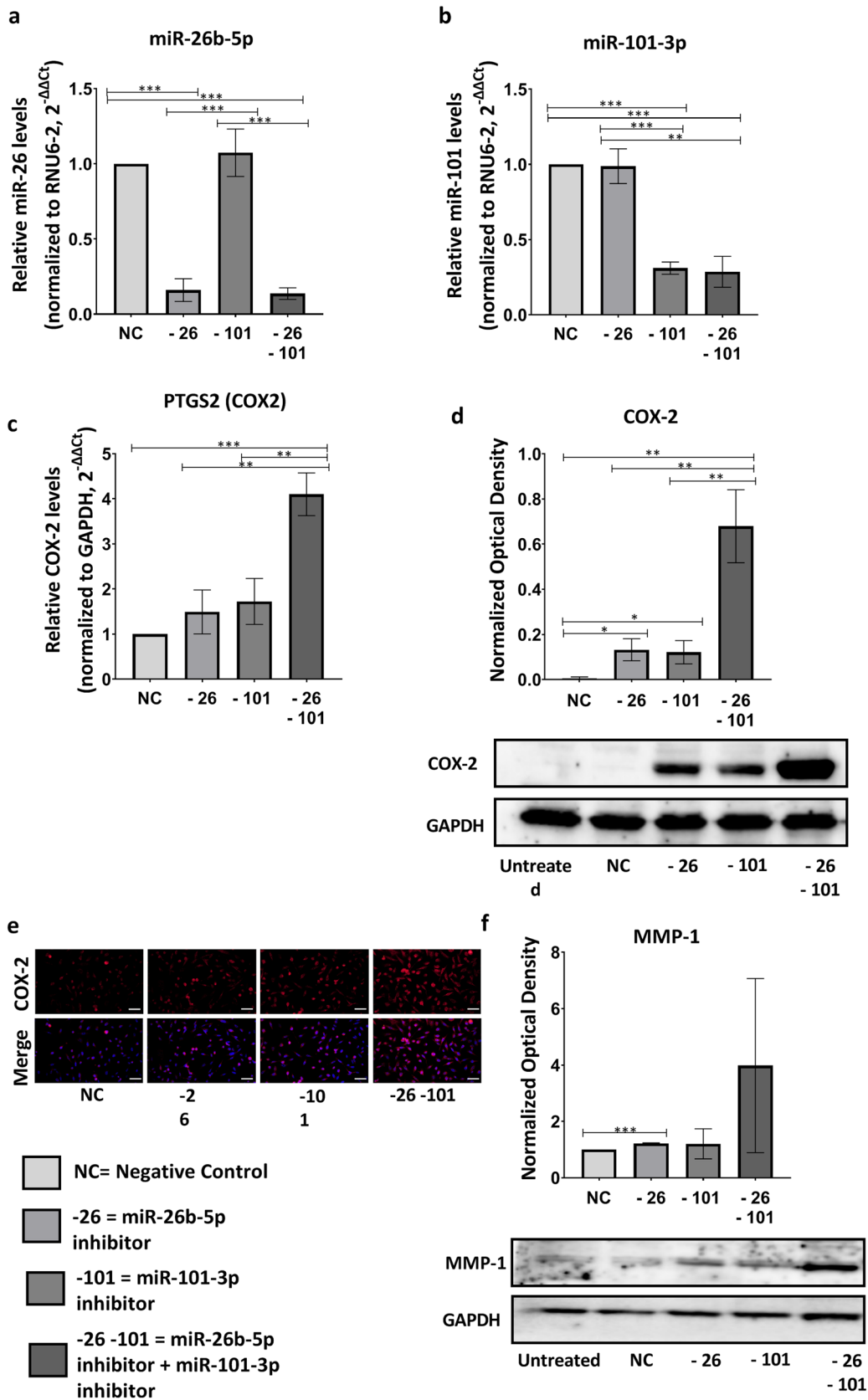


Fig. 3 The dual inhibition of miR-26b-5p and miR-101-3p produces higher increase in COX-2/MMP-1 expression compared to single micro-RNA. MDA-MB-231-TGL cells were transfected with miR-26b-5p/miR-101-3p inhibitors or negative control. **a, b** Relative levels of miR-26b-5p (**a**) and miR-101-3p (**b**) measured by real-time PCR. The small nuclear RNA (RNU6-2) was used as an internal standard. **c** Relative levels of COX-2 (PTGS2) mRNA levels measured by real-time PCR. GAPDH was used as an internal standard. **d** Western blot analysis of COX-2 protein. Optical densities were analyzed with Image Lab 6.0.1 software (Bio-Rad) and normalized to GAPDH. **e** Immunofluorescence staining of COX-2. **f** Western blot analysis of MMP-1 protein. Data represent mean \pm SD from three independent experiments. Scale bar = 50 μ m. * p < 0.05, ** p < 0.01, *** p < 0.001

PGE2 were further reduced in cells transfected with both mimics (fold change = 0.03, p < 0.001) compared to cells transfected with either miR-26b-5p (fold change = 0.25, p = 0.0004) or miR-101-3p (fold change = 0.29, p = 0.014) alone compared to control (Fig. 6f). The combination also exerted an enhanced repressive effect on MMP-1 levels. Relative to control cells, MMP-1 protein level was further reduced in cells co-transfected with both inhibitors (fold change = 0.25, p = 0.066) compared to cells transfected with miR-26b-5p (fold change = 0.76, p = 0.0013) or miR-101-3p (fold change = 0.52) alone (Fig. 6g). Similarly, relative to control cells, levels of MMP-1 released in the media were further reduced in cells co-transfected with both mimics (fold change = 0.05, p < 0.001) than with single miR-26b-5p (fold change = 0.56, p = 0.0092) or miR-101-3p (fold change = 0.50, p = 0.0348) (Fig. 6h). Taken together, these results demonstrate that COX-2/MMP-1 repression was

greater when cells are transfected with both micro-RNAs (miR-26b-5p and miR-101-3p) than single micro-RNA.

The combinatorial restoration of miR-26 and miR-101 inhibits the transmigration of TNBC cells through the brain endothelium and preserves the endothelial barrier integrity

We next investigated whether the synergistic effect of combining miR-26b-5p and miR-101-3p on COX-2 repression resulted in greater suppression of BC cells transmigration through the brain endothelium. Relative to control cells, transmigration of cancer cells through primary brain endothelial cells was 37.24% lower when MDA-MB-BrM2 cells are transfected with miR-26b-5p alone (p < 0.001), 33.55% lower when MDA-MB-BrM2 cells are transfected with miR-101-3p alone (p = 0.02), and 76.75% lower when cells are transfected with the combination of miR-26b-5p and miR-101-3p mimics (p < 0.001) (Fig. 7a). Similarly, transmigration of cancer cells through hCMEC/D3 cells was 62.27% lower when MDA-MB-BrM2 cells are transfected with miR-26b-5p alone (p < 0.001), 36.99% lower when MDA-MB-BrM2 cells are transfected with miR-101-3p alone (p = 0.0113), and 91.70% lower when cells are transfected with the combination of miR-26b-5p and miR-101-3p mimics (p < 0.001) (Fig. 7b and c). These results indicate that combining miR-26b-5p and miR-101-3p result in greater suppression of the transmigration of metastatic BC cells through the brain endothelium than did either miR-26b-5p or miR-101-3p alone. We then evaluated effect

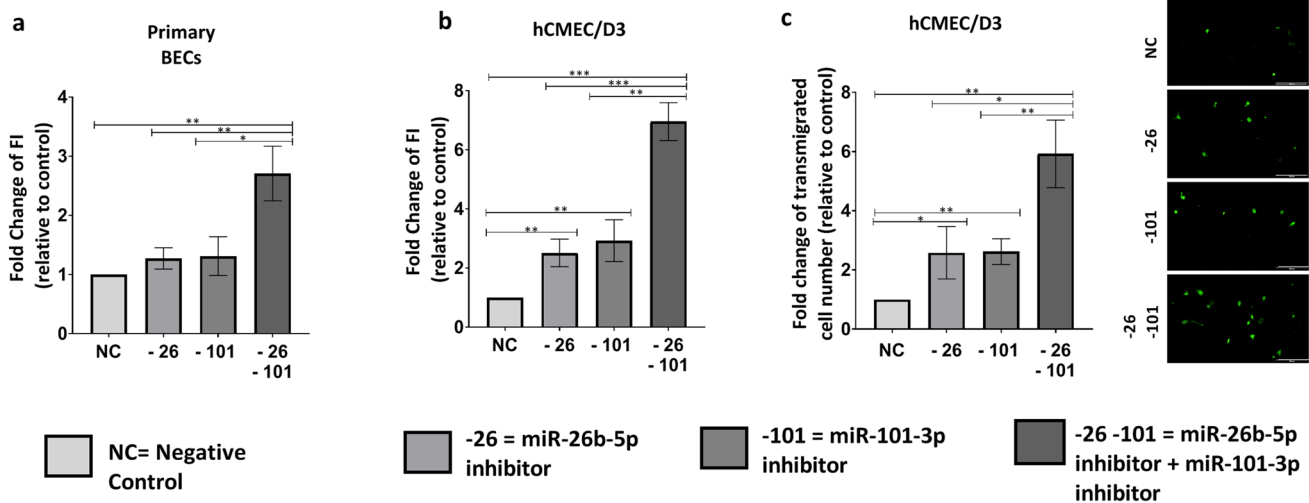


Fig. 4 The dual inhibition of miR-26b-5p and miR-101-3p results in higher transmigration of TNBC cells through the brain endothelium compared to single micro-RNA. **a, b** Fluorescently labeled MDA-MB-231-TGL cells transfected with miR-26b-5p/miR-101-3p inhibitors (or negative control) were seeded on top of transwell inserts coated with **(a)** primary BECs or **(b)** hCMEC/D3 cells. The fluorescent

intensity in the lower chambers was measured after 24 h. **c** The number of invaded cells was counted in three different fields per insert. Representative images of transmigrated cells (small green cells) are shown. Data represent mean \pm SD from three independent experiments. * p < 0.05, ** p < 0.01, *** p < 0.001

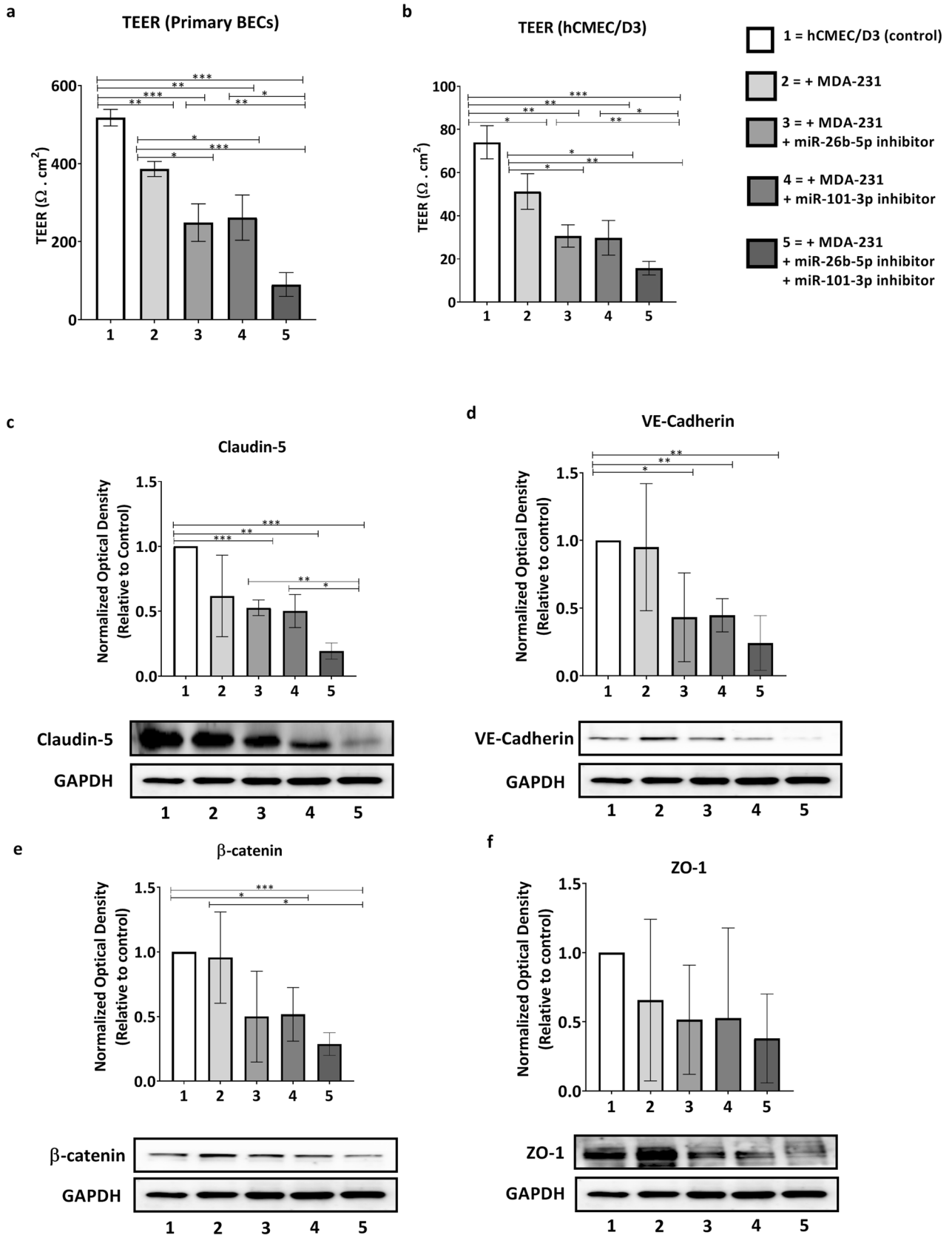


Fig. 5 The dual inhibition of miR-26b-5p and miR-101-3p in TNBC cells exacerbates disruption of the brain endothelium integrity. **a,b** Transendothelial electrical resistance (TEER) of primary brain endothelial cells (**a**) or hCMEC/D3 (**b**) co-cultured with MDA-MB-231-TGL cells pre-transfected with miR-26b-5p/miR-101-3p inhibitors or negative control after 24 h of co-culture. **c–f** Western blot analysis of the inter-endothelial junctional proteins (claudin-5, VE-cadherin, β -catenin, and ZO-1) of hCMEC/D3 cells co-cultured with transfected MDA-MB-231-TGL cells. Optical densities were analyzed with Image Lab 6.0.1 software (Bio-Rad) and normalized to GAPDH. Data are mean \pm SD from three independent experiments. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$

of combining miR-26b-5p and miR-101-3p on the barrier integrity and its inter-endothelial junctions. As shown in Fig. 8, combination of both micro-RNAs restores normal TEER values and normal protein levels of inter-endothelial junctions. These results show that the dual restoration of miR-26b-5p and miR-101-3p potentiated the suppression of COX-2 and trans-endothelial migration of metastatic cells, which is in accordance with previous studies showing that the biological effect of micro-RNAs can be potentiated by the synergistic effect of combining two micro-RNAs that target a same gene.

Discussion

In the present study, we provide first-time evidence of a synergistic effect exerted by a pair of micro-RNAs (miR-26b-5p and miR-101-3p) on COX-2 expression in brain metastatic triple-negative breast cancer cells. The dual inhibition of miR-26b-5p and miR-101-3p results in higher increase in COX-2/MMP-1 and grants metastatic cells a higher trans-migrative capacity through the brain endothelium. The combinatorial restoration of both miR-26b-5p and miR-101-3p exerted a synergistic repressive effect on COX-2/MMP-1 by targeting COX-2 and potentiated the suppression of trans-endothelial migration compared to either micro-RNA alone. These findings provide new insights on the regulation of COX-2 by micro-RNAs and shed light on the pair (miR-26b-5p and miR-101-3p) as novel prognostic and therapeutic tool that can serve to predict TNBC patients at high risk of brain metastasis and to develop novel anti-metastatic therapeutic strategies with enhanced efficacy.

The role of COX-2 in breast cancer pathogenesis has been widely studied. Cyclooxygenase enzymes catalyze the conversion of arachidonic acid to prostaglandins. The isoform COX-1 is constitutively expressed in normal tissues, whereas COX-2, undetectable in normal physiological conditions, is inducible in response to stimuli such as mitogens, cytokines, growth factors, or hormones [43]. COX-2 is overexpressed in several malignant tumors and was shown to promote carcinogenesis and cancer cell resistance to chemo- and radiotherapy [44, 45]. COX-2 overexpression converts arachidonic acid

(AA) into prostaglandin E₂ (PGE₂) which promotes cancer progression through inducing migration [46], stem-like cell (SLC) formation [47], and angiogenesis [48, 49]. In case of breast cancer metastasis, a number of studies showed that COX-2 expression is correlated with the metastatic spread of cancer cells and has a strong potential as prognostic indicator of disease severity and progression [50–54]. In addition, triple-negative breast cancer patients with high levels of COX-2 were found to have the greatest risk of developing brain metastasis [11]. Mechanistic studies showed that upregulation of COX-2 in TNBC metastatic cells induces MMP-1 which degrades the inter-endothelial junctions and promotes passage of cancer cells through the BBB [11–13].

Due to its important role in cancer pathogenesis, several preclinical and clinical studies tested the use of COX-2 inhibitors alone and in combination with other agents for the treatment and prevention of breast cancer, and in the adjuvant, neo-adjuvant, and metastatic treatment settings ([43, 44, 49]. Preclinical studies demonstrated that celecoxib suppress the proliferation and growth of breast cancer, while clinical studies suggested that celecoxib administration is related to reduced incidence of breast cancer in women without disease, reduced recurrence risk and mortality in women with breast cancer, and a better prognosis. However, despite promising preclinical data, trials on COX-2 inhibitors in patients with established breast tumor were unsuccessful in improving overall survival [49, 54, 55]. Similarly, clinical trials with MMP inhibitors failed to prove a survival benefit in patients with advanced disease. Batimastat, a multi-MMPs inhibitor, showed promising preclinical anti-tumor effect, however significant toxicity was observed. The clinical trials on another MMPs inhibitor, Marimastat, failed to prove a survival benefit with severe side effects. A more selective MMPs inhibitor was also tested for the treatment of metastatic lung, breast, and prostate carcinomas. However, the trials failed to demonstrate a positive effect on survival [56]. Among the possible reasons explaining the failure of COX-2 and MMP-1 inhibitors in clinical trials is that these trials were performed in patients with advanced metastatic disease, however, COX-2 and MMP-1 inhibitors are expected to be effective in early pre-metastatic stages and may not exert beneficial therapeutic effect if used in advanced stages [49, 56]. In addition, the role of COX-2 in promoting brain metastasis was documented in the TN subtype specifically [11, 12]. However, several pieces of evidence indicate that the molecular mechanism driving brain metastasis depends on the molecular subtype of BC cells. For instance, Palmiery et al., showed that HER2 overexpression promotes extravasation of HER2+ BC cells through the blood–brain barrier [57]; however TNBC do not express HER2 and therefore differences in the molecular mechanisms are evident. Therefore, COX-2 and MMP-1 inhibitors need to be tested for their metastasis preventive effect in

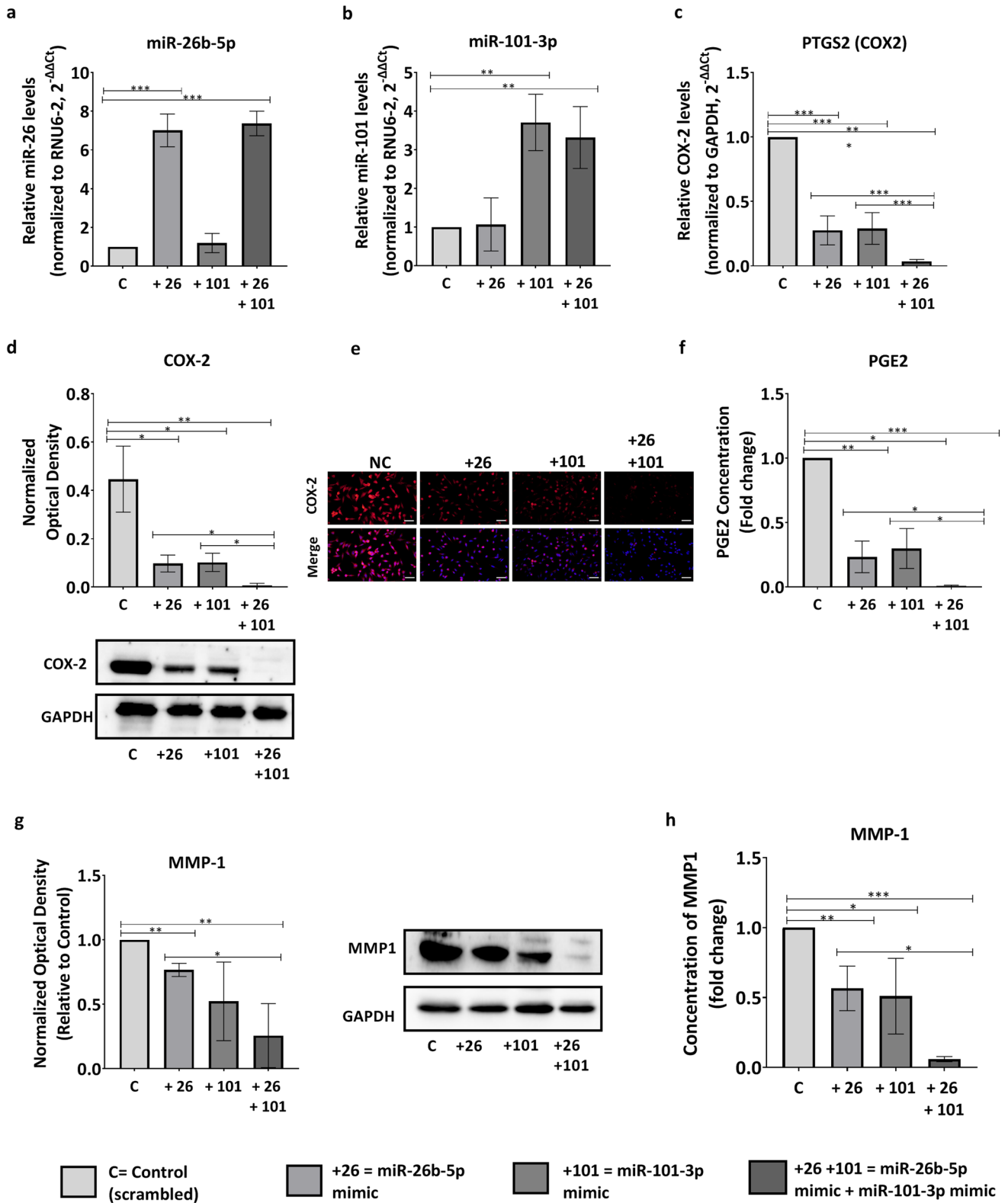


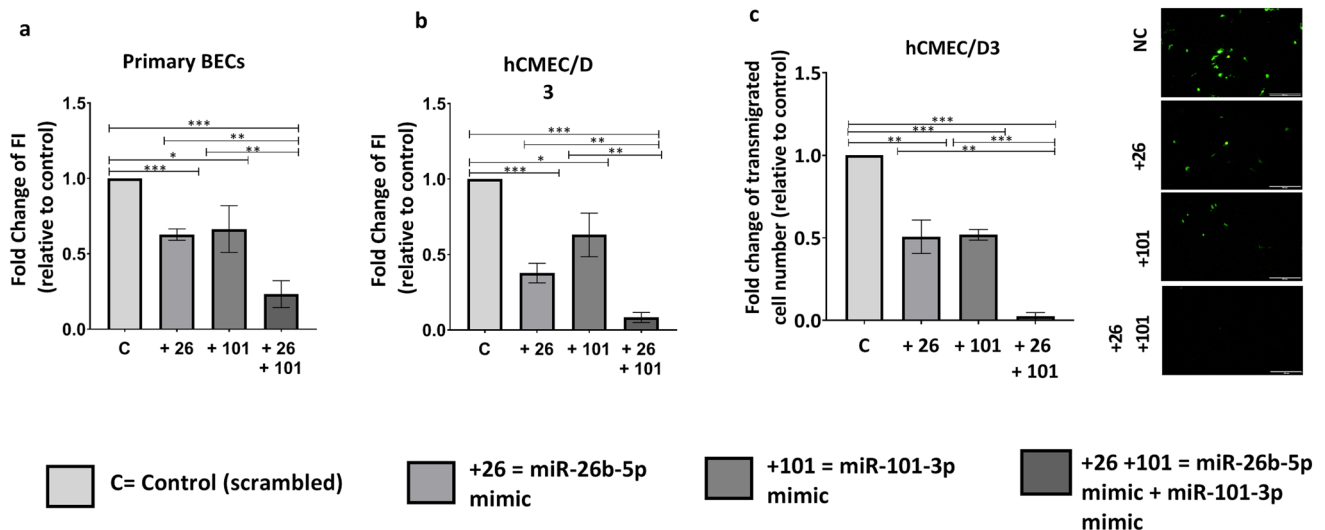
Fig. 6 Combination of miR-26 and miR-101 exerts a synergistic repressive effect on COX-2 expression in brain metastatic TNBC cells. MDA-MB-231-BrM2 cells were transfected with miR-26b-5p/miR-101-3p mimics or control. **a, b** Relative levels of miR-26b-5p (**a**) and miR-101-3p (**b**) measured by real-time PCR. The small nuclear RNA (RNU6-2) was used as an internal standard. **c** Relative levels of COX-2 (PTGS2) mRNA levels measured by real-time PCR. GAPDH was used as an internal standard. **d** Western blot analysis of COX-2 proteins. Optical densities were analyzed with Image Lab 6.0.1 software (Bio-Rad) and normalized to GAPDH. **e** Immunofluorescence staining of COX-2. **f** PGE2 levels released in the culture media quantified by ELISA. **g** Western blot analysis of MMP-1 proteins. **h** MMP-1 levels released in the culture media were quantified by ELISA. Data represent mean \pm SD from three independent experiments. Scale bar = 50 μ m. * p < 0.05, ** p < 0.01, *** p < 0.001

early stages before metastases are established and in the TN subtype specifically.

However, for this strategy to become within reach, the molecular mechanisms upregulating COX-2 and MMP-1 in triple-negative breast cancer cells and rendering them more prone to disseminate to the brain need to be characterized in order to identify patients at high risk of developing brain metastasis and for better therapeutic interventions.

In this study, we identified a synergistic effect of two micro-RNAs (miR-26b-5p and miR-101-3p) in upregulating COX-2 expression in metastatic TNBC cells. miR-26b-5p and miR-101-3p are downregulated in brain metastatic cells compared to parental cells. These results are in accordance with previous studies showing that miR-26b-5p and miR-101-3p are downregulated in brain metastatic tumors

compared to primary tumors in patients [29]. The roles of miR-101 and miR-26 were studied in several types of cancer [24, 58–61]. In triple-negative breast cancer cells, miR-101 was shown to exert tumor suppressor effects [62] and to increase sensitivity to paclitaxel [63]. miR-101-3p was also reported to have anti-metastatic effect in breast cancer [62–64], glioblastoma [65], lung cancer [66], hepatocellular carcinoma [67], and gallbladder carcinoma [68]. Similarly, miR-26b was shown to exert anti-tumor and anti-metastatic effects in several types of cancer including breast [26], lung [27], and human tongue carcinoma [24]. miR-26 was also found to be downregulated in advanced inflammatory breast cancer [69]. Interestingly, several studies showed that COX-2 is a direct target of miR-26b-5p and miR-101-3p. Specifically, miR-101-3p was shown to target COX-2 mRNA in endometrial carcinoma [23], colon cancer [25], and breast cancer [28], while miR-26b-5p was found to target COX-2 in breast cancer [26], lung cancer [27], and human tongue carcinoma [24]. However, role of these two micro-RNAs in regulating COX-2 expression in metastatic breast cancer cells and their dual role in brain metastasis remain to be elucidated. In a previous study, our group demonstrated that loss of miR-101-3p in metastatic breast cancer cells induces COX-2 and promotes transmigration of TNBC cells through the brain endothelium [28]. Mechanistic studies on the action of micro-RNAs reported that micro-RNAs tend to collaboratively or synergistically control gene expression [19, 21]. It was suggested that such synergistic gene regulation may maximize miRNAs' efficiency and



lower chambers was measured after 24 h. **c** The number of invaded cells was counted in three different fields per insert. Representative images of transigrated cells (small green cells) are shown. Data represent mean \pm SD from three independent experiments. * p < 0.05, *** p < 0.001

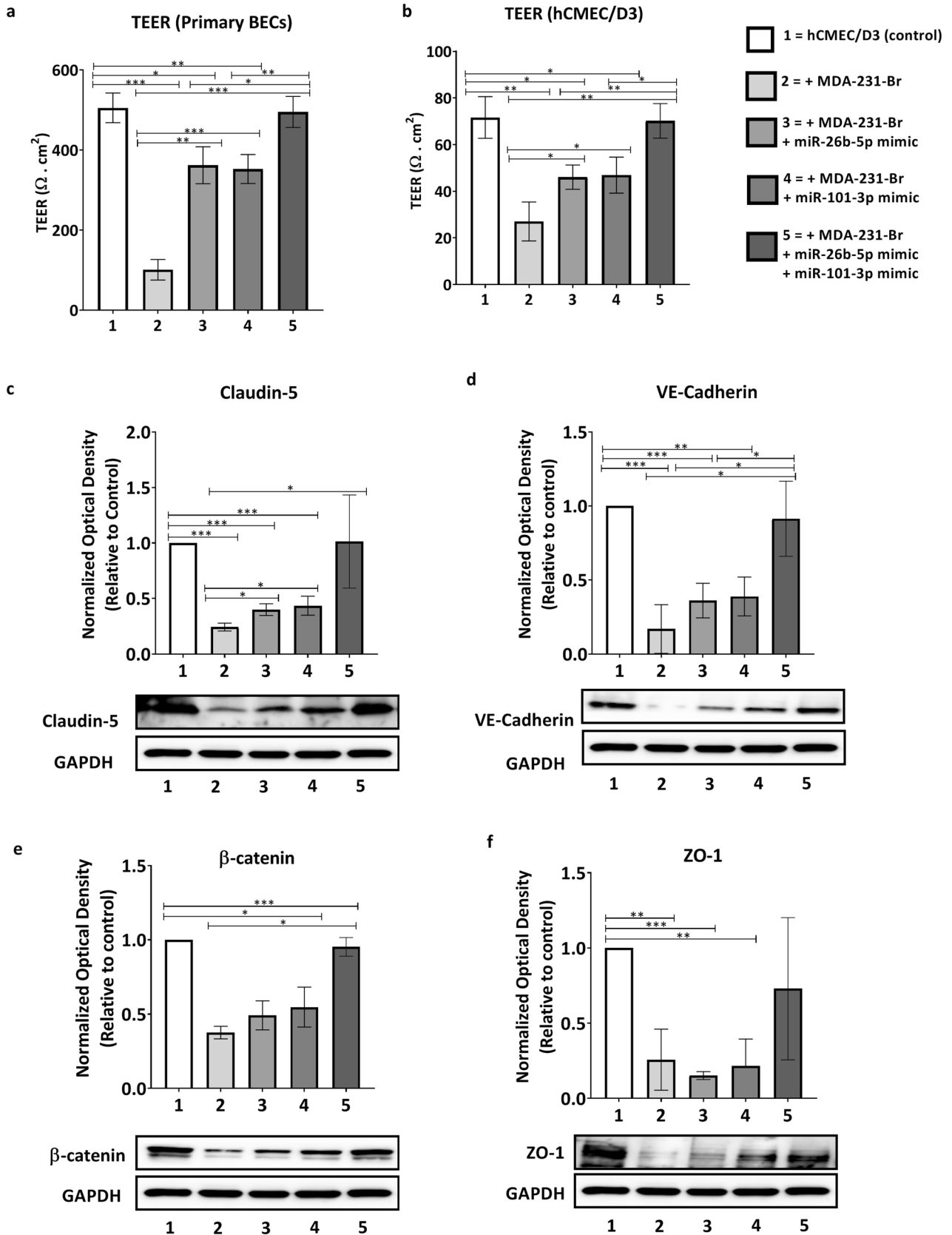


Fig. 8 Combination of miR-26 and miR-101 preserves the endothelial barrier integrity. **a,b** Transendothelial electrical resistance (TEER) of primary brain endothelial cells (**a**) or hCMEC/D3 (**b**) co-cultured with MDA-MB-231-BrM2 cells pre-transfected with miR-26b-5p/miR-101-3p mimics or control after 24 h of co-culture. **c–f** Western blot analysis of the inter-endothelial junctional proteins (claudin-5, VE-cadherin, β -catenin, and ZO-1) of hCMEC/D3 cells co-cultured with transfected MDA-MB-231-BrM2 cells. Optical densities were analyzed with Image Lab 6.0.1 software (Bio-Rad) and normalized to GAPDH. Data are mean \pm SD from three independent experiments. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$

that the collaborative miRNA–miRNA interaction may be an important regulation mode of endogenous miRNA. As miR-26b-5p and miR-101-3p are downregulated in brain metastases and are both validated to target the 3'UTR of COX-2, we examined their potential collaboration in regulating COX-2 expression in metastatic BC cells and on tumor cells transmigration capacity. We also tested whether a combinatorial restoration of miR-26b-5p and miR-101-3p could exert a synergistic repressive effect on COX-2 expression and suppress transmigration of TNBC cells through the brain endothelium, key step in brain metastasis. Our findings showed that the dual inhibition of both micro-RNAs (miR-26b-5p and miR-101-3p) results in higher increase of COX-2/MMP-1 and a higher transmigration of tumor cells through the brain endothelium suggesting a potential collaboration between the two micro-RNAs in the regulation of COX-2 expression. An examination of the inter-endothelial junctions revealed that the dual loss of both micro-RNAs exacerbated the disruption of the endothelial barrier compared to single inhibition. In contrast, restoration of both miR-26b-5p and miR-101-3p collaboratively targets COX-2 and exerts a synergistic repressive effect on COX-2/MMP-1. In addition, restoration of miR-26b-5p and miR-101-3p suppressed the transmigration of brain metastatic TNBC cells through the brain endothelium and preserved the barrier integrity. Importantly, the trans-endothelial suppression was potentiated when both micro-RNAs were restored simultaneously compared to single micro-RNA. These findings confirm previous reports suggesting that the simultaneous modulation of miRNA pairs exerts synergistic effects on the target genes which leads to the potentiation of their biological effects compared to single micro-RNAs.

Conclusion

The present study adds to our knowledge of the molecular mechanisms upregulating COX-2 expression in metastatic TNBC cells and granting them a higher brain propensity. Our findings imply that the pair (miR-26b-5p and miR-101-3p) could be used as a potential prognostic tool to

identify TNBC patients at high risk of developing brain metastasis. In addition, we shed light on (miR-26b-5p and miR-101-3p) as potential therapeutic targets that can be exploited by subsequent in vivo and translational studies to develop better therapeutic strategies that prevent breast cancer brain metastasis to date incurable.

Supplementary Information The online version contains supplementary material available at <https://doi.org/10.1007/s10549-021-06255-y>.

Author contributions RH designed the study, performed research, analyzed, and interpreted the data and wrote the manuscript; AT, MM, and GK contributed to the luciferase reporter assay; AT, AM, GK, and RH provided support with analysis of data and interpretation of results. All authors read and approved the final manuscript.

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Data availability The authors confirm that the data supporting the findings of this study are available within the article and/or its supplementary material.

Declarations

Conflict of interest The authors declare that there are no conflicts of interest/competing interests.

Ethical approval All experiments were approved by the University of Sharjah animal care and use committee and conducted according to the UOS directives for animal care.

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