A novel small-molecule antagonizes PRMT5-mediated KLF4 methylation for targeted therapy

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ABSTRACT

Background: Triple negative breast cancers (TNBCs) have a poor prognosis and are not amenable to endocrine- or HER2-targeted therapies. The malignant and invasive feature of TNBCs is correlated with its high cancer stem cell population. Recent results from us and others have unveiled an oncogenic role for the PRMT5-KLF4 axis in regulating tumor progression by orchestrating the stemness in mammary tumor cell as well as genome stability. Methylation of KLF4 by PRMT5 leads to KLF4 stabilization, resulting in promoting mitogenesis.

Methods: We have developed a small molecule inhibitor, WX2–43, that specifically intercepts the interaction between PRMT5 and KLF4, thereby enhancing KLF4 degradation.

Findings: Results from our characterization demonstrate that WX2–43 binds to the region between amino acids L400-M500 on PRMT5. Degradation of KLF4 down-regulates KLF4-mediated genes transcription. We have characterized the potent effect for WX2–43 in inhibiting PRMT5-KLF4 binding that, in turns, suppresses tumor progression and induces tumor cell death in both TNBC cultured-cell and animal models.

Interpretation: WX2–43-mediated inhibition of KLF4 methylation by PRMT5 could be a potential strategy for anti-TNBC treatment.

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1. Introduction

Kruppel-like factor 4 (KLF4) acts as a pivotal transcription factor and regulates various biological processes such as cell growth, proliferation, apoptosis, metabolism and stem cell self-renewal, and maintenance of pluripotency [1]. KLF4 with other three transcriptional factors, OCT3/4, SOX2, and c-MYC could switch somatic cells into stem cells [2]. Surprisingly, results from recent genetic and pathological studies have uncovered an ambivalent nature for KLF4 in tumorigenesis as either a tissue-specific tumor suppressor or an oncogene with mystery underlying mechanism [3]. KLF4 has been reported to have tumor-suppressive properties in lung cancer, gastric cancer, colorectal cancers, urothelial cancer, cervical cancer, bladder, prostate and stomach, while it acts as an oncogenic factor in breast, oral, osteosarcoma, melanoma and skin squamous cell carcinomas [4–7]. KLF4 has a pivotal function in cancer stem cells maintenance and differentiation in various types of cancer including breast, colon, and osteosarcoma [3,8–11]. Interestingly in colon cancer, KLF4 has been recognized as a well-characterized tumor suppressor, it still functions as an oncogene for cancer stem cell [8,10,12].
Breast cancer self-renewal of breast cancer stem cells and cell motility could abolish doxorubicin-induced cancer stem cells and reverses the study reported that using simvastatin to downregulate KLF4 expression maintenance leads it to be an attractive target for cancer therapy. Recent cancer stem cell [19,20]. It is upregulated in various human malignancies, including lymphomas, lung cancer, breast cancer, and colorectal cancer [23,24]. A phase I study of GSK3326595 regarding its safety and activity is continuing in solid tumors and non-Hodgkin lymphoma (NCT02783300). An ongoing phase I dose escalation study of GSK3326595 aims to identify the safety and activity of the PRMT5 inhibitor GSK3326595 in solid tumors and non-Hodgkin lymphoma (NCT02783300). This study was initiated 2 years' ago and is expected to be completed in 2019. However, the current PRMT5 inhibitors designed against its enzymatic activity lacks specificity for the individual downstream substrate.

We have recently demonstrated that KLF4 is a rapidly turned over protein with its half-life governed by VHL-VBC ubiquitin-protein ligase. We have further revealed that KLF4 could be methylated by PRMT5, whose methylation on amino acids R374, R376, and R377 counteracts VHL/VBC-mediated KLF4 ubiquitylation [4]. We further observed that adding up methyl group to KLF4 alters KLF4 protein conformation that, in turn, prevents KLF4 ubiquitylation VHL/VBC E3 ligase through affecting the recognition of KLF4 by VHL/VBC, thereby stabilizing KLF4. In response to DNA damage signal, PRMT5-mediated KLF4 methylation causes failure of DNA damage-induced KLF4 accumulation that, in turn, reduces p21-mediated DNA damage response. In tumorigenesis, downregulation of KLF4 methylation leads to increased Bax levels and reduced p21 expression levels that further suppress several critical oncogenes, including cyclin D2, cyclin B1, Myc, EGF, and IGF1, resulting in inhibition of tumor initiation and tumor progression. Results from TCGA and proteomic studies have indicated that PRMT5 upregulation closely associated with KLF4 accumulation in mammary cancer [4]. Thus, inhibition of KLF4 function through dissociating interaction between PRMT5 and KLF4 could be an ideal approach to suppress breast cancer. In the present study, we report a novel in silico identified and in vitro validated small-molecule that blocks PRMT5-KLF4 binding and inhibits breast cancer progression. Our development of a small-molecule inhibitor that specifically intercepts the interaction between PRMT5 and KLF4 provides a promising candidate drug that suppresses breast cancer growth. Our findings could potentially lead to the development of more specific PRMT5 inhibitor for anti-TNBC treatment.

2. Experimental procedures

2.1. Construction of the 3D structural model of PRMT5 and its substrates

The 3D structure of human KLF4 was constructed by using reported KLF4 crystal structures (PDB ID: 2WBS, 2WBU, and 4M9E). The 3D crystal structure of human p53 (PDB ID: 3Q01) was retrieved from Protein Data Bank (www.rcsb.org). The sequence (residues from 112 to 199) of small nuclear ribonucleoprotein-associated protein B (Drosophila melanogaster) was downloaded from UniprotKB (https://www.uniprot.org/) and constructed using I-TASSER server (https://zhanglab.ccmb.med.umich.edu/I-TASSER/) [30]. The 3D structures modeling of KLF4-PRMT5 binding was conducted on the basis of our previous report [4], and the 3D structural models of PRMT5-p53 and PRMT5-SMB were constructed by using Z-DOCK server (http://zdock.umassmed.edu/) [31].

2.2. Development of KLF4 methylation inhibitor

We adopted the MOLCAD module implemented in SYBYL-X 1.3 to explore the potential binding pockets based on the complex of KLF4-PRMT5 and KLF4-P53.
PRMT5. Sequentially, we performed virtual docking screenings against our 3D chemical database using our reported protocol in order to identify small molecules that can interfere with PRMT5/KLF4-binding [32]. Briefly, the docking parameters used were as follows: (a) the “number of starting conformations per ligand” was set to 3, and the “number of max conformations per fragment” was set to 20; (b) the “maximum number of rotatable bonds per molecule” was set to 100; (c) flags were turned on at “pre-dock minimization”, “post-dock minimization”, “molecule fragmentation”, and “soft grid treatment”; (d) “activate spin alignment method with density of search” was set to 9.0; and (e) the “number of spins per alignment” was set to 12.

2.3. Cell lines and cell culture

HEK293T, MCF10A, MCF12A, 184-B5, MDA-MB-468, MDA-MB-453, HCC1937, HCC38, MCF-7, and MDA-MB-231 were obtained from the American Type Culture Collection (Manassas, VA). The KLF4+/+ MEF and KLF4−/− MEF were gifts from Dr. Engda Hagos (Colgate University). HEK293T, MDA-MB-468, MDA-MB-453, HCC1937, HCC38, MCF-7, MDA-MB-231, KLF4+/+ MEF and KLF4−/− MEF were maintained in DMEM supplemented with 5 or 10% FBS, 1× antibiotic/antimycotic solution (100 units/ml streptomycin and 100 units/ml penicillin) (all from Invitrogen). MCF10A, MCF12A, 184-B5 were maintained in monolayer...
2.5. Lentiviral infection

transfected using Lipofectamine 2000 (Invitrogen). The plasmids pLenti6/FLAG-KLF4WT, pLenti6/FLAG-KLF43K, GST fused PRMT5 were co-transfected with packaging plasmids pVSV-G, pRRE, and pRSV-REV into HEK293T cells by using Lipofectamine 2000. The generated lentiviral particles were collected, mixed with polybrene, and then added to target cells. The stable cell lines were established by culturing the cells in the medium containing antibiotic blasticidin (10 μg/ml).

2.6. RNA interference

The synthesized siRNA sequences are as following: Luciferase, 5′-CGT ACCGGGAATCCTGCA-3′; SipRMT5–1# 5′-GCCAGTGTAGATGCCTATAT-3′; SipRMT5–2# 5′-CCGTATTCGGACCTGACCC-3′; p53 (Clone DO-1, Santa Cruz Biotechnology) and HRP-conjugated goat anti-mouse or antirabbit secondary antibody (Promega). The SYM10 antibody against Symmetric Di-Methyl Arginine motif (SDMA) was purchased from Millipore. Signals were detected with ECL reagents (Millipore). For immunoprecipitation assay, cell lysate was incubated with KL4F (Santa Cruz Biotechnology), PRMT5 (Millipore, 07–405), SmD3 (Abgent), p53 (Clone DO-1, Santa Cruz Biotechnology) antibody overnight at 4 °C on a rotator, followed by the addition of protein A/G plus agarose (Santa Cruz Biotechnology) to the reaction for 2 h at 4 °C. After five washes with lysis buffer supplemented with protease inhibitor mixture, complexes were released by boiling for 5 min in 2× SDS-PAGE loading buffer. Western blotting was used to detect SDMA modification.

2.7. Western blotting and immunoprecipitation assay

Cells were harvested and lysed in radioimmune precipitation assay (RIPA) lysis buffer (Cell Signaling Technology) containing protease inhibitor mixture (Sigma) or 1× SDS loading buffer. The protein concentration was determined using Bio-Rad protein assay reagent. Western blotting was performed using anti-KLF4 (Santa Cruz Biotechnology), PRMT5 (Millipore, 07–405), p21 (BD Bioscience, 556,430), FLAG (Sigma), β-actin (Sigma), SmD3 (Abgent), SmB (clone 12F5, Sigma), p53 (Clone DO-1, Santa Cruz Biotechnology) and HRP-conjugated goat anti-mouse or antirabbit secondary antibody (Promega). The SYM10 antibody against Symmetric Di-Methyl Arginine motif (SDMA) was purchased from Millipore. Signals were detected with ECL reagents (Millipore). For immunoprecipitation assay, cell lysate was incubated with KL4F (Santa Cruz Biotechnology), PRMT5 (Millipore, 07–405), SmD3 (Abgent), p53 (Clone DO-1, Santa Cruz Biotechnology) antibody overnight at 4 °C on a rotator, followed by the addition of protein A/G plus agarose (Santa Cruz Biotechnology) to the reaction for 2 h at 4 °C. After five washes with lysis buffer supplemented with protease inhibitor mixture, complexes were released by boiling for 5 min in 2× SDS-PAGE loading buffer. Western blotting was used to detect SDMA modification.

2.8. Recombinant GST-KLF4 and GST -PRMT5 pull-downs experiments

GST-fused KL4F and PRMT5 were expressed in the E. coli strain BL21 and purified using glutathione-Sepharose 4B resin following manufacturer’s protocol. 2 μg of purified recombinant protein was incubated at 4 °C for 1 h with 500 μg of cell lysates from MDA-MB-231 cells or 2 μg His tagged KL4F purified from E. coli strain BL21. Glutathione resin was added and interacting protein was pulled down. After five washes with 1× TBS buffer, the resin was eluted with 1× SDS loading buffer.

2.9. Biotin-WX2-43 pull-downs assay

2 μg/ml biotin-WX2-43 was incubated at 4 °C for 1 h with 20 μg 1 μg/ml GST-PRMT5 fragments or GST-PRMT5 (GST-PRMT1–9) purified from E. coli. Avidin-agarose resin (Sigma) was then added and incubated at 4 °C for 1 h. Biotin-WX2-43 interacting protein was pulled down by centrifuge. After five washes with 1× TBS buffer, the resin was eluted with 1× SDS loading buffer.

2.10. WX2-43 binding ELISA assay

Purified human GST-PRMT5 proteins were coated at a 96-well microplate at 2 μg/ml (100 μl/well) for overnight in 4 °C and washed three times with washing buffer (PBS with 0.2% Tween-20). Wells were blocked with 1% BSA in PBS, incubated for 1 h, and then rinsed with Wash Buffer. Biotinylated-WX2-43 (100 μl) with a linear range concentration of 0.1 nM–10 μM diluted in 1% BSA / PBS were added into the plates and further incubated for 2 h at room temperature. Plates were washed three times with washing buffer (PBS with 0.2% Tween-20). After rinsing, 100 μl of Streptavidin–HRP at a concentration of 50 ng/ml in 1% BSA in PBS was added and incubated for 20 min at room temperature. Then plates were washed three times with washing buffer (PBS with 0.2% Tween-20). 100 μl of Substrate TMB (3,3′,5,5′-Tetramethylbenzidine, Sigma) Solution was added and incubated for 20 min at room temperature. 50 μl of the Stop Solution was added to each well. The absorbance of light with the wavelength of 450 nm was measured with 800TS absorbance reader (Biotek). For freedom WX2-43 competitive ELISA, 5 μM biotin-WX2-43 in 1% BSA / PBS was mixed with freedom WX2-43 at 1:1 (5 μM biotin-WX2-43 : 0 μM WX2-43), 1:2 (5 μM biotin-WX2-43: 10 μM WX2-43), 1:5 (5 μM biotin-WX2-43: 25 μM WX2-43).
25 μM WX2-43), 1:10 (5 μM biotin-WX2-43: 50 μM WX2-43), 1:20 (5 μM biotin-WX2-43: 100 μM WX2-43) (mol: mol) and then incubated with GST-PRMT5 coated 96-well plates for 2 h.

2.11. Animal experiments assay

All animal experiments were performed according to the protocols approved by the Institutional Animal Ethics Committee of Northwestern University. The investigation conforms to the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health (2011, 8th Ed.). In the orthotypic model, 5 × 10^6 of MDA-MB-231 cells mixed with matrigel (1:1 volume) were injected into the mammary fat pads of 8-week-old female Crl: Nu-Foxn1(nu) nude mice. 1 week later when the tumor was palpable, WX2-43 and vehicle (10% N,N-Dimethylacetamide in saline) were intraperitoneally administrated twice per week at the dose of 40 mg/kg, 80 mg/kg. Tumor volume was measured twice a week. After 4 weeks continued administration, mice were sacrificed and tumor tissues were collected for western blot.

2.12. Flow cytometry analysis

The flow cytometry analysis was conducted as the previous report [4], the breast cancer cells at the logarithmic growth phase were digested with 0.25% trypsin and washed with PBS for three times, followed by being re-suspended in 100 μl PBS, and then stained with anti-CD44-PE and anti-CD24-FITC or stained with their isotype controls at room temperature for 40 min. The samples were then washed by PBS three times and finally re-suspended in 200 μl PBS. Flow cytometry analysis was performed on a BD Accuri™ C6 Flow Cytometer (BD Bioscience). The expression ratios of CD44 to CD24 (CD44/CD24) in different subtypes of breast cancer cell lines were calculated from the percentage of CD44 and CD24 positive subpopulations in the flow cytometry analysis.

2.13. Aldefluor assay

The ALDEFLUOR kit (StemCell Technologies, Durham, NC, USA) was used to analyze the population with high ALDH enzymatic activity according to the manufacturer's instructions. Briefly, the cells were incubated in the ALDEFLUOR assay buffer containing ALDH substrate (BAAA, 1 mmol/l per 1 × 10^6 cells), and incubated for 40 min at 37°C. In each experiment, an aliquot of cells was incubated under identical conditions, using 50 mmol/l of diethylamino benzaldehyde (DEAB), as a specific ALDH inhibitor, as a negative control.

2.14. Chemistry and compounds

PRMT5 inhibitor epz015666 was purchased from the Selleckchem Company. The compounds to be tested as possible blockers of PRMT5-KLF4 complexation, including WX2-43, were synthesized in-house. All compounds were prepared in DMSO stock solution at a concentration of 10 mM and stored at −20°C before use. The compounds were then diluted in Iscove’s modified Dulbecco’s medium (GBICO) to make a working solution, as needed. The final DMSO concentration was <0.1% in the working solution.

2.15. Statistical analysis

The significance of the differences in the assays was analyzed by Student’s t-test or one or two-way ANOVA, followed by Tukey’s multiple comparisons test. A value of P < .05 was considered significant.

3. Results

3.1. Abnormal accumulations of PRMT5 and KLF4 are observed in triple negative breast cancer

We have recently reported that deregulated KLF4 methylation by PRMT5 results in genome instability and mammary tumorigenesis. To decipher the abnormal accumulation of PRMT5 and KLF4 in breast cancer, we have measured the protein expression levels of KLF4 and PRMT5 in various molecular subtypes of breast cancer cells as well as human breast tumor tissue. As shown in Fig. 1, while KLF4 and PRMT5 expressions are relatively low in mammary epithelial cell lines MCF10A and moderate in HER2 positive breast cancer cell lines SKBR3 and BT474, significant accumulation of KLF4 is observed in triple negative and ER positive type of breast cancer cell lines MDA-MB-231, MDA-MB-468, MCF7 and T47D, especially in triple negative breast cancer (TNBC) cells MDA-MB-231 (Fig. 1a). We further analyzed the expression levels of both PRMT5 and KLF4 in various types of breast patient specimens using a tissue array with BC081120c, which contains 10 adjacent normal breast tissue (normal), 62 ER/PR positive breast cancer (ER+/PR+), 19 HER2 positive breast cancer (HER2) and 23 TNBC tissues. As shown in Fig. 1b–c, in general, both PRMT5 and KLF4 are overexpressed in various breast cancer tissues but severe PRMT5 and KLF4 protein accumulation were obviously detected in the TNBC. In addition, the Kaplan-Meier survival analyses for PRMT5 and KLF4 indicates that patients with overexpression of PRMT5 correlate well with poor prognosis in comparison with patients with low expression of PRMT5 or KLF4 in TNBC patients (Fig. 1d & e). Thus, the abnormal accumulation of PRMT5 and KLF4 protein could be oncogenic causation for mammary tumor initiation and progression. Targeting PRMT5-KLF4 axis could be a novel strategy anti-TNBC therapy.

3.2. Methylation of KLF4 contributes to cell survival in response to DNA damage and cancer stem cell maintenance

The current principal clinical therapy for TNBCs is relayed on chemotherapy following by radiation therapy [33]. High subpopulation of cancer stem cell existing in TNBC tissue has been a major issue affecting the therapeutic efficiency [27]. To ask if accumulation of KLF4 due to the PRMT5-mediated methylation regulates cancer cell survival and enhances cancer stem cell expansion, we have measured the impact of KLF4 methylation, by functional interference of KLF4 methylation through expression of a KLF4-methylation deficient mutant, KLF4-3 K, in various types of breast patient specimens (Fig. 2a, b). In the Kaplan-Meier survival analyses for PRMT5 and KLF4 indicates that patients with overexpression of PRMT5 correlate well with poor prognosis in comparison with patients with low expression of PRMT5 or KLF4 in TNBC patients (Fig. 1d & e). Thus, the abnormal accumulation of PRMT5 and KLF4 protein could be oncogenic causation for mammary tumor initiation and progression. Targeting PRMT5-KLF4 axis could be a novel strategy anti-TNBC therapy.
Fig. 4. Characterization of blockade effect of WX2–43 on the PRMT5-KLF4 interaction and PRMT5-mediated KLF4 methylation. (a) WX2–43 efficiently intercepts the interaction between KLF4 and PRMT5, and blocks KLF4 methylation. (b) Blockade of KLF4 methylation further leads to downregulation of p21 and upregulation of Bax. (c) Specificity of WX2–43, LLY-283 and EPZ015666. MDA-MB-231 cells were treated with WX2–43, LLY-283 and EPZ015666 at indicated dose for 24 h and then expressions of SDMA, p53, KLF4, SMB were detected. All three compounds could highly efficiency abrogated the whole cell SDMA modification at 1 μM level. WX2–43 shows less efficiency at 0.1 μM level to inhibit total SDMA modification. However, WX2–43 shows more capability in inhibiting SDMA modification on KLF4. (d) Comparison of the substrate specificity for WX2–43 and that of the non-specific PRMT5 inhibitor EPZ015666.
the radiation-induced upregulation of KLF4. We further observed that elevated expression of KLF4 leads to increased cancer cell survival against chemo-drug Doxorubicin and Cisplatin but suppression of KLF4 methylation results in significant attenuation of cancer cell survival (Fig. 2b). Meanwhile, knockdown PRMT5 expression by siRNA also decreases KLF4 expression and sensitizes MDA-MB-231 cells to chemo-drug Doxorubicin and Cisplatin (Fig. 2c & d). In addition, we have further examined the role of KLF4 methylation in the maintenance of cancer stem cell using 3D mammosphere formation assay. As shown in Fig. 2e–i, in both triple negative and ER positive breast cancer cells, elevated expression of KLF4 largely enhances cancer stem cell population (CD44+/CD24− or ALDH positive), but prevention of KLF4 methylation leads to significant suppression of breast cancer stem cells. Taken together, the above results clearly show that the methylation of KLF4 by PRMT5 is critical to regulating cell survival and cancer stem cell maintenance.

3.3. Development of an inhibitor that specifically blocks the interaction between PRMT5 and KLF4

Based on the critical effect of KLF4 methylation in promoting breast tumor progression, cancer cell survival after chemotherapy and cancer stem cell expansion, we decided to develop small molecule inhibitor that could efficiently block PRMT5-KLF4 complexation as a new anti-breast cancer therapeutic approach. As shown in Fig. 3a (left), we first looked into the detailed information of the binding sites in PRMT5. We found that the key residues involved in the binding site of EPZ015666 included Phe327, Glu444, Phe577, Phe580. Moreover, we observed that EPZ015666 forms strong hydrogen bond with Glu444 (2.6 Å), Phe577 (3.3 Å), and Phe580 (2.8 Å), while interacting with Phe327 (3.5 Å) via a strong n–π interaction. The key PRMT5 residues involved in the binding site of SAM included Phe327, Tyr334, Glu392, Asp419, Met420, Glu435, Glu444, and Cys449. Furthermore, we observed that SAM formed a strong hydrogen bond with Asp419 (3.3 Å), and Phe580 (2.8 Å), while interacting with Phe327 (3.5 Å) via a strong n–π interaction. The key residues involved in the binding site of SAM included Phe327, Tyr334, Glu392, Asp419, Met420, Glu435, Glu444, and Cys449. Furthermore, we observed that SAM formed a strong hydrogen bond with Asp419 (3.3 Å), and Phe580 (2.8 Å), while interacting with Phe327 (3.5 Å) via a strong n–π interaction. The key residues involved in the binding site of SAM included Phe327, Tyr334, Glu392, Asp419, Met420, Glu435, Glu444, and Cys449. Furthermore, we observed that SAM formed a strong hydrogen bond with Asp419 (3.3 Å), and Phe580 (2.8 Å), while interacting with Phe327 (3.5 Å) via a strong n–π interaction.

3.4. Characterization of blockade effect of WX2-43 on the PRMT5-KLF4 interaction and PRMT5-mediated KLF4 methylation

To further validate whether WX2-43 could block the binding between PRMT5 with its substrates including KLF4, P53 and SMB, we first detected the capacity of WX2-43 in dissociation of interaction between PRMT5 with KLF4, P53 and SMB by using functional cell lysate-based GST-fused PRMT5 pull-down assay. Interestingly, all three compounds including WX2-43, LLY-283 and EPZ-0156666 could have capacity to dissociate PRMT5-p53 and PRMT5-SMB binding, while only the compounds WX2-43 has an obvious capacity to block PRMT5-KLF4 binding in MDA-MB-231 cell lysates in a dose-dependent manner (Fig. 3e). Meanwhile, we further used purified His-tagged KLF4 protein instead of MDA-MB-231 lysates identified only the WX2-43, other than LLY-283 and EPZ-0156666 could prevent PRMT5-KLF4 binding (Fig 3f). To further detect the capacity of WX2-43 in the dissociation of interaction between PRMT5 with KLF4, we further use functional cell lysate-based GST-KLF4 pull-down assay following by measurement of the KLF4 methylation status as well as binding of PRMT5 to KLF4 by western blotting. As shown in Fig. 4a, we observed that GST-KLF4 could efficiently pull down PRMT5 proteins from MDA-MB-231 cell lysates and the GST-KLF4 protein also could be methylated by endogenous PRMT5 present in cell lysates. We further observed that pre-addition of the WX2-43 to the lysates significantly decreased the efficacy for KLF4 to pull-down PRMT5 and suppresses KLF4 methylation in a dose-dependent manner. We next measured the effect of WX2-43 on the KLF4 downstream target p21 and Bax. As shown in Fig. 4b, we observed that pretreatment of MDA-MB-231 cells with WX2-43 for 24 h, significantly decreased KLF4 protein levels and its transcriptional activation in upregulating p21 expression and downregulating Bax gene expression. These data are in support of the interference of WX2-43 with the methylation of KLF4 by PRMT5, thus promoting KLF4 ubiquitylation and degradation.

To compare our newly developed PRMT5-KLF4 specific inhibitor with recently reported non-specific PRMT5 inhibitor LLY-283 and EPZ0156666, we have tested all three compounds side by side (Fig. 4c and d). By staining of SYM10 antibody against Symmetric DI-Methyl Arginine Motif (SDMA), the whole cell symmetric di-methylation modification could be measured. All three compounds could highly efficiently abrogated the whole cell SDMA modification at 1 μM level, but WX2-43 shows less potent at 0.1 μM dose (Fig. 4c). However, when
immunoprecipitated KLF4 and detected the SDMA modification, WX2-43 shows more capacity to inhibit KLF4 modified by PRMT5, which indicated WX2-43 is more KLF4 specific PRMT5 inhibitor. WX2-43 is discovered here, on the other hand, based on specific blockade of interaction between PRMT5 and KLF4, and is therefore anticipated to suppress KLF4-mediated downstream effects exclusively. We further compared WX2-43 with EPZ015666 inhibits the catalytic activity of PRMT5 in general and thereby inhibits the methylation of all PRMT5 substrates.
indiscriminately. As shown in Fig. 4d, while WX2-43 specifically inhibits KLF4 methylation, EPZ015666 reduces the methylation of all three substrates. Based on these results, we have identified a specific PRMT5-KLF4 binding inhibitor. Thus, WX2-43 could be defined as a specific PRMT5-KLF4 inhibitor. Taken together, the above experiments validated that WX2-43, identified with guidance from in-silico studies, is an effective compound that specifically blocks PRMT5-KLF4 binding, and affects KLF4 stability and suppresses KLF4 cellular activity.

3.5. WX2-43 is a potent PRMT5 specific inhibitor

Given the presence of nine members in the protein arginine methyltransferases family, we next ask if our newly developed PRMT5 inhibitor also affect other members. Previous works have sketched the non-redundant feature for the nine members (PRMT1-9) (Fig. S3) [37]. To date, we expressed all nine GST fused PRMTs and then do the Biotin-labelled WX2-43 pull down. Interestingly, only the PRMT5 shows affinity to WX2-43 (Fig. 5a). To further study how WX2-43 binds to PRMT5 that in turn interrupts PRMT5-KLF4 binding, we have generated a series of GST-PRMT5 fragment deletions and then detected the exact molecular motif that binds to WX2-43 by using pulldown assay [38]. To date, we subjected biotinylated-WX2-43 conjugated with Streptavidin-agarose to a set of GST-PRMT5 fragments including 1-400Aa, 1-500Aa, 500-637Aa and the full-length, and then pull-down biotinylated-WX2-43 coupled Streptavidin-agarose following by measuring retaining of GST-PRMT5 fragment fusion protein abundance by Coomassie blue staining and western blotting (Fig. 5a). As shown in Fig. 5b, we only observed the GST-PRMT5 1–500 and the full length PRMT5 could be pulled down by biotin-WX2-43, indicating that the amino acids stretch between L400-M500 is involved in binding WX2-43. This result is consistent with the core amino acids predicted in our molecular docking model, including Glu392, Met420, Asp419, Glu435, Glu444 localized in this area. We further mutated the PRMT5 Glu392, Asp419 and Glu 435 to alanines (EDE/AAA) and then did the biotin-WX2-43 pulldown. Mutated these three amino acids abrogated WX2-43 binds to PRMT5, which indicated there three amino acids facilitate WX2-43 binding (Fig. 5c). We further measured the binding coefficient of biotin-WX2-43 with GST-PRMT5 using ELISA and found the binding $K_d$ is $11.0 \pm 1.03$ nM (Fig. 6a). In addition, the binding of biotin-WX2-43 to PRMT5 was observed to be dose-dependent by free WX2-43 in the competition ELISA assay, suggesting the binding of biotin-WX2-43 to PRMT5 is specific (Fig. 6b).

3.6. WX2-43 is an inhibitor that specifically blocks KLF4-mediated transcription and cell survival

We next examined whether WX2-43 inhibits KLF4 function via specifically affecting PRMT5-KLF4 axis. As shown in Fig. 6c–e, we observed

![Fig. 7. WX2-43 is a potent inhibitor that suppresses breast tumor progression. (a, b) MDA-MB-231 cells were implanted into the mammary fat pad of nude mice. WX2-43 were treated twice a week at the indicated dose. Tumor volume was measured once a week and mice were sacrificed 32 days after injection. Tumor volume growth curve and tumor weight were measured. The asterisk indicates significant difference ($p < .01$) between vehicle, 40 mg/kg and 80 mg/kg treatment group. Data are mean ± SEM; one-way ANOVA was used for the statistical analysis. (c) Measurement of expression of PRMT5, KLF4 and P21 in xenograft tumor. After tumor harvest at 32 days, the tumors were lysed and detected by Western blotting using indicated antibodies.](image-url)
that WX2-43 significantly decreases KLF4 abundance and its targeted gene p21 expression as well as cell proliferation, and this effect is very similar to that of PRMT5 knockdown, indicating the WX2-43 has no additive effect on KLF4-p21 expression and cell proliferation other than via PRMT5. To further determine whether WX2-43 inhibits cell viability especially via KLF4, we have tested cell viability in response to cisplatin by using the KLF4 knockout MEFs (KLF4+/−/− MEFs) cells. We observed that, while wild-type KLF4 MEFs has obvious dose response to WX2-43 inhibition, lack of KLF4 in MEFs attenuates the dose responsive manner, suggesting the effect of WX2-43 is through KLF4 (Fig. 6e). We previously observed that the KLF4 expression levels in mammary gland epithelial cells are much lower in comparison with TNBC cells. We wondered if the response to WX2-43 is different between MCF10A and MDA-MB-231. We tested the IC50 of WX2-43 in mammary epithelial cell lines including 184-B5, MCF10A, MCF12A and breast cancer cell lines MDA-MB-231, MDA-MB-468, MCF7, HCC-1957, HCC38 and MDA-MB-453, and observed that triple negative breast cancer cell lines such as MDA-MB-231, HCC1937 cells are much more sensitive to WX2-43 than normal epithelial cancer cell lines MCF10A, MCF12A and 184-B5. The observed WX2-43 killing selectivity between mammary gland epithelial and TNBC cells suggest that WX2-43 could be a selective inhibitor for anti-TNBC therapy (Fig. 6f).

3.7. WX2-43 is a potent inhibitor that suppresses breast tumor progression

The above characterization has shown that WX2-43 is a potent inhibitor that specifically disrupts PRMT5-KLF4 interaction and selectively targets TNBC cells for growth inhibition. To determine the physiological and clinical relevance of WX2-43 inhibitor in breast cancer progression, we have examined the impact of WX2-43 in TNBC using human breast cancer xenograft model [39]. MDA-MB-231 cells were subcutaneously injected into athymic nude mice, WX2-43 where intraperitoneal (i.p.) administrated and tumor growth was monitored. The administration of 40, 80 mg/kg WX2-43 to mice had no obvious toxic effect and mice gained weight normally. 80 mg/kg WX2-43 reduced approximately by 60%, and 40 mg/kg WX2-43 reduced by about 40% both the size and weight of breast cancer xenograft tumor in comparison with vehicle (Fig. 7a-b). We further measured the expression levels of KLF4, p21 and PRMT5 in the isolated grafted tumors using Western blotting, and confirmed the effect of WX2-43 on inhibiting KLF4 function upon WX2-43 administration (Fig. 7c). The result from the in vivo analysis shows that WX2-43 is a potent inhibitor that suppresses TNBC progression. Altogether, our results indicate that PRMT5-KLF4 plays a critical role in promoting breast tumor initiation and progression through regulating cell genome stability and stem cell expansion. Blockade of the PRMT5-KLF4 oncogenic axis by our newly developed PRMT5-KLF4 inhibitor could be a novel strategy to treat TNBC.

4. Discussion

KLF4 is a pivotal cellular fate determinant, whose deregulation has been linked to a variety of human disease through orchestrating various cellular processes [1]. Previous studies from TCGA and pathological analyses unveiled the correlation of abnormal regulation of KLF4 with poor prognosis of breast cancer, although the underlying mechanism has been reminded until recently. We have demonstrated that the management of KLF4 protein levels is accurate and controlled by a sophisticated interplay between ubiquitylation and protein methylation [4,40]. We have shown the KLF4 protein turnover is regulated by VHL/VBC-mediated ubiquitylation following by degradation. Intriguingly, we have observed that the KLF4 protein degradation is counteracted by protein methylation, wherein the addition of methyl group to the KLF4 my arginine methyltransferase PRMT5 stabilizes KLF4 stability through antagonizing its ubiquitylation due to interruption the binding between KLF4 with its ubiquitin E3 ligase VHL because of alteration of its intracellular molecular confirmation [4,40]. Results from our pathological analyses have drawn attention to the unexpected accumulation of both PRMT5 and KLF4 in breast cancer tissue and their significant correlation [4]. Elevation of KLF4 or PRMT5 leads to malignant mammary epithelial cell transformation and tumor progression in preclinical models [4]. Evidence from multiple layers clearly implies that the suppression of KLF4 function by blockade of its methylation catalyzed by PRMT5 could be a valuable approach to inhibit breast tumor progression, especially for TNBC. In the present study, the development of a specific KLF4 methylation inhibitor provides a promising strategy to suppress breast tumor progression by targeting PRMT5-KLF4 interaction. Our characterization has further confirmed the specificity, selectivity as well as efficacy in killing TNBC cells in both cultured-cell and animal models.

PRMT5 is the major symmetric arginine methyltransferase in mammals [19]. Previous pathological studies have demonstrated the impact of PRMT5 in promoting various oncogenic processes [25,41]. Targeting PRMT5 recently became a hot spot for anti-cancer drug development. Various approaches such as the high-throughput screen, pharmacophore- and docking-based virtual screening have been utilized to search for PRMT5 small-molecule inhibitor. Such efforts have been further encouraged by the recently reported development of PRMT5 inhibitors by several companies [42]. Approximately 50 candidate PRMTs inhibitors have been documented, including EPZ015666 (IC50 = 22 nM), Compound 9 (IC50 = 35 μM), compound 17 (IC50 = 0.33 μM), Compounds CMP5 and HLCL-61 [21,22,28,29]. Among these, EPZ015666 shows outstanding selectivity for PRMT5 with promising potency (IC50 = 0.022 μM) but limited to other PRMTs. PRMT5 is reported to modulate transcription through the methylation of histone H2 arginine 3 (H2R3), histone H3 arginine 8 (H3R8) and histone H4 arginine 3 (H4R3). The major effects of these inhibitors are more rely on histone methylation and transcriptional regulation. WX2-43, our newly developed PRMT5-KLF4 specific inhibitor is unique and precise because of its specificity on the blockade of PRMT5-facilitated KLF4 methylation, which determines KLF4 protein stability and KLF4-mediated oncogenesis. WX2-43, in preclinical models, has shown its attractive potency in killing TNBC cells with IC50 at 11 nM, and EC50 at 2 μM. Importantly, WX2-43 is not a PRMT5 pan-methyltransferase inhibitor but only specifically targets PRMT5-mediated KLF4 methylation. Therefore, WX2-43 is more specific than other inhibitors such as EPZ015666 to target the PRMT5-KLF4 axis for suppressing breast tumor progression. The sister inhibitor of EPZ015666, GSK3326595, is ongoing phase 1 trial in solid tumors and non-Hodgkin lymphoma, which implies the emerging clinical value of PRMT5 inhibitor. Likely, WX2-43 has its advantage and potential in targeting specificity in comparison with other PRMT5 pan-methyltransferase inhibitors in future’s anti-breast cancer treatment.

We demonstrated the role of PRMT5-KLF4 axis in regulating DNA damage response and breast tumorigenesis [4,43]. In response to genotoxic stress, precise control of KLF4 abundance determines the cellular fate either undergoing survival or committing to death through regulating the ratio between p21 and Bax [4]. Under the circumstance of oncogenesis, our chip assay revealed the elevated KLF4 levels leads to upregulation of a series of oncoproteins such as EGF, IGF1, Myc and CDKs [4]. These observations imply that manipulation of KLF4 by blockade of PRMT5-KLF4 axis could potentially synergize KLF4 with other critical targeted drugs such as CDK4/6 inhibitor and PARP inhibitors in potent killing breast cancer cells. We observed, in high KLF4 expression set breast cancer cell lines, the elevated KLF4 levels leads to reduced drug response [4]. Our work inspires a novel clinical intervention strategy in combination of PRMT5-KLF4 inhibitor with PARP inhibitor, talaraparib/olaparib, or CDK4 inhibitor for anti-TNBC therapy.

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Author contributions

Y.W. and XX are the PIs who managed the whole project. Y.W., Z.Z., Z.F. and XX designed the experiments and wrote the manuscript. Z.Z. performed the experiments. D.H. provided some plasmons and participated in some experiments. Z.F. and P.Y. performed the development of PRMT5-KLF4 inhibitor. M.G. participated in structure simulation of PRMT5-KLF4. M.C. and W.J.G. provided consultation on some drug related experiments. I.B. supervised M.G. in molecular modeling of PRMT5-KLF4.

Declaration of competing interests

Dr. Xiangqun Xie issued a patent US8772541.

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