Comprehensive analysis of PPPCs family reveals the clinical significance of PPP1CA and PPP4C in breast cancer

Wenjun Xie¹b,#, Ying Sun¹c,#, Yu Zeng³e,#, Linfei Hu³, Jingtai Zhi³, Hang Ling³, Xiangqian Zheng³, Xianhui Ruan³, and Ming Gao⁴f

¹Department of Thyroid and Neck Tumour, Tianjin Medical University Cancer Institute and Hospital, National Clinical Research Center for Cancer, Key Laboratory of Cancer Prevention and Therapy, Tianjin’s Clinical Research Center for Cancer, Tianjin, China; ²Provincial Clinical College, Fujian Medical University, Fuzhou, China; ³Chongqing University Cancer Hospital, Chongqing Key Laboratory of Translational Research for Cancer Metastasis and Individualized Treatment, Chongqing, China; ⁴Department of Otolaryngology-Head and Neck Surgery, Tianjin First Center Hospital, Nankai District of Tianjin, Institute of Otolaryngology of Tianjin, Key Laboratory of Auditory Speech and Balance Medicine, Key Clinical Discipline of Tianjin (Otolaryngology), Otolaryngology Clinical Quality Control Centre, Tianjin, China; ⁵Department of Pathology, Fujian Provincial Hospital, Fuzhou, China; ⁶Department of Thyroid and Breast Surgery, Tianjin Union Medical Center, Tianjin, China

ABSTRACT
The phosphoprotein phosphatase catalytic subunit (PPPCs) family has been shown to play an important role in the development and progression of various malignancies, but its expression patterns and biological functions in breast cancer (BC) remain unclear. Therefore, we aimed to investigate the clinical significance and biological functions of the PPPCs family to understand its possible significance in the diagnosis, prognosis and treatment of breast cancer. We comprehensively investigated the expression levels, diagnostic accuracy, prognostic outcomes, biological functions and effects on immune cell infiltration of the PPPCs family in breast cancer using online databases. Except for PPP1CB, PPP1CC, PPP5C and PPEF1, the mRNA expression levels of the PPPCs family in breast cancer tissues were significantly different from those in paracancerous tissues. The differentially expressed genes (DEGs) were associated with the clinicopathological parameters and prognosis of breast cancer. The DEGs were mainly associated with the WNT signaling pathway, antigen presentation and DNA repair. In addition, the DEGs significantly affected the infiltration of immune cells in breast cancer tissues. Among the PPPCs family, PPP1CA and PPP4C played a prominent role in the progression of breast cancer, and inhibition of PPP1CA and PPP4C expression by siRNA can significantly inhibit breast cancer cells proliferation and migration. In conclusion, the PPPCs family, especially PPP1CA and PPP4C, could be used as new biomarkers to improve diagnostic accuracy, predict prognosis and novel targets for the treatment of breast cancer.

Introduction
Breast cancer has become the most prevalent tumor worldwide, with 2.3 million new cases expected in 2020, accounting for 11.7% of cancer cases and becoming the fifth leading cause of death from cancer [1]. The prognosis of breast cancer has been significantly improved by targeted therapy, radiotherapy and immunotherapy [2].

However, some breast cancer patients still suffer a poor prognosis; approximately 20% of metastatic breast cancer patients survive for only 5 years [3]. Given the high incidence of breast cancer and the poor prognosis of some patients, the search for new biomarkers for breast cancer diagnosis, prognosis and treatment remains the focus of breast cancer research. With the continuous advancement and application of bioinformatics analysis, protein families associated with breast cancer have been discovered, such as the APRO protein family [4] and the AQP family [5], and the discovery of these indicators has implications for breast cancer diagnosis, prognosis and treatment policy.

Protein phosphatase can be classified into four gene families: serine/threonine phospho-
protein phosphatase (PPP); Mg\(^{2+}\)-dependent protein phosphatase (PPM/PP2C); phospho-
yrosine phosphatase (PTP) and asp-based pro-
tein phosphatase [6]. The PPP family consists of
conserved catalytic subunits. At present, thir-
teen different isoforms of the PPP catalytic sub-
unit (PPPCs) family have been identified: PPP1CA,
PPP1CB, PPP1CC, PPP2CA, PPP2CB, PPP3CA,
PPP3CB, PPP3CC, PPP4C, PPP5C, PPP6C, PPEF1
and PPEF2. The PPCPs family can regulate a var-
iety of signaling pathways, and dysregulation of these genes leads to
aberrant processes, including uncontrolled pro-
liferation, differentiation and metastasis [7].
Numerous subsequent studies have found that
the PPCPs family is aberrantly expressed in
various tumors [8]. However, the role of the
PPCPs family in breast cancer has not been
systematically explored.

In this study, we hypothesized that the PPCPs
family plays an important role in the pathogenesis
and prognosis of breast cancer. Therefore, we
comprehensively analyzed the role of the PPCPs
family in breast cancer through online databases.
We subsequently inhibited the expression of
PPP1CA and PPP4C in breast cancer cells by
siRNA and observed changes in their proliferation
and migration abilities, and verified through these
results that these two genes played an outstanding
role in the development of breast cancer. These
studies will help to find new biomarkers relevant
for the diagnosis, prognosis and treatment of
breast cancer.

Materials and methods

TCGA database

The fragments per kilobase of per million (FPKM)
of breast cancer transcriptome (including 1109
tumor samples and 113 normal samples) were
downloaded from The Cancer Genome Atlas
(TCGA, https://tcgadata.nci.nih). The relevant
data were then processed by Rstudio software
v3.6.3 (https://www.rstudio.com/). The receiver
operating characteristic (ROC) curves of genes
were described by the pROC package [9], and
their area under the receiver operating charac-
teristic curve (AUROC) scores were ranked from
high to low. DeLong’s test was used to determine
the diagnostic efficacy of each gene, and P < 0.05
was considered significant.

Oncomine database

Oncomine [10] (https://www.Oncomine.org) is an
online cancer gene expression profile database
containing 715 datasets and 86,733 samples. We
used this database to analyze the transcript levels
of the PPCPs family in breast cancer.

cBioPortal database

cBioPortal [11] (http://www.cbioportal.org/) is an
open resource for the interactive exploration of
multiple cancer genomic datasets. We used the
database to explore the PPCPs family alteration
frequency, and the correlation with the prognosis
of breast cancer patients was explored [12].

GEPIA database

The Gene Expression Profiling Interactive Analy-
sis (GEPIA) V2.0 database [13] (http://gepia.
cancerpku.cn/) provides customizable functions such as tumor/
normal differential expression analysis, similar gene
detection, and gene correlation detection. We used
this database to detect gene correlations in the PPCPs
family and the top 70 genes in breast cancer tissues
that were similar to genes in the PPCPs family.

Kaplan–Meier Plotter database

The Kaplan–Meier (K-M) Plotter database [14]
(https://kmplot.com/analysis/), as a meta-analysis-
based biomarker assessment tool, was able to assess
54 K gene expression levels on the prognosis of 21
tumors, including breast cancer. Using the K-M
Plotter, we analyzed the prognostic value of the
PPCPs family gene mRNA expression levels in breast
cancer.

Breast cancer gene-expression Miner
(bc-GenExMiner) v4.5 database

Bc-GenExMiner v4.15 [15] (http://bcgenex.ico.unin
cancer.fr) is a statistical mining tool that contains
breast cancer transcriptomic data and prognostic
data. We analyzed the relationship between the PPPCs family and breast cancer clinicopathological parameters and prognosis using this database. The subtypes of parameters include age, nodal status, estrogen receptor (ER), progesterone receptor (PR), human epidermal growth factor receptor 2 (HER-2) and triple-negative status.

**Metascape database**

Metascape [16] (http://metascape.org/) is a web-based portal for the analysis and annotation of gene lists. Metascape combines feature-rich, interactive group analysis, gene annotation, and member search by combining more than 40 independent knowledge bases in one integrated portal. In this study, we used it to visualize the results of the enrichment analysis.

**TIMER database**

TIMER [17] (https://cistrome.shinyapps.io/timer/) is a database designed to analyze immune cell infiltration in a wide range of cancers. The database uses statistical methods validated by pathological examination to estimate tumor immune infiltration by neutrophils, macrophages, dendritic cells, B cells and CD4/CD8 T cells. We used this database to explore the relationship between the PPPCs family and the extent of infiltration of specific immune cell subpopulations.

**Immunohistochemistry and evaluation of the immunostaining intensity**

Paraffin-embedded breast cancer tissues were obtained from the Department of Pathology of Tianjin Cancer Hospital and Fujian Provincial Hospital. The study included 120 samples from 60 patients diagnosed with breast cancer. The sections were incubated with anti-PPP1CA (ab150782, 1:200, Abcam, UK) or anti-PPP4C (ab195371, 1:200, Abcam, UK) at 4°C overnight. The intensity of immunohistochemistry (IHC) staining was scored as 1 (weak); 2 (medium); and 3 (strong). The degree of staining was scored (from 0 to 4) according to the percentage of immunoreactive tumor cells (<5%, 5–25%, 26–50%, 51–75% and >75%). By multiplying the staining degree score with the staining intensity score, a score between 0 and 12 was calculated for each example [18]. All sections were scored by 2 independent pathologists.

**Cell culture**

The human breast cancer cell lines MCF-7 and MDA-MB-468 were purchased from the Cell Bank of the Chinese Academy of Sciences (China). MCF-7 and MDA-MB-468 cell lines were cultured in DMEM high-glucose medium (Gibco, USA) containing 10% fetal bovine serum.

**siRNA transient interference assay**

siRNA was purchased from GenePharma (GenePharma, China) (Supplementary Table 1). MCF-7 or MDA-MB-468 cells (1 × 10^6) were cultured in six-well culture plates (NEST, China). A total of 5 µL of siRNA and 5 µL of RNAiMAX Reagent (Thermo Fisher, USA) were diluted in 125 µL of Opti-MEM I Reduced Serum Medium (Gibbon, USA) and incubated for 5 min separately. The solution was mixed and then analyzed for transient interference according to the manufacturer’s protocol.

**CCK-8 cell proliferation assay**

Cell proliferation was determined using the Cell Counting Kit-8 (CCK-8) assay. MCF-7 or MDA-MB-468 cells (2 × 10^5/well) were plated in 96-well plates (Nest, China). The absorbance of each well at 450 nm was measured using a multimode enzyme marker (SYNERGY H1, BioTek, USA).

**Scratch assay**

MCF-7 or MDA-MB-468 cells (8 × 10^5 cells/well) were plated in 6-well plates and cultured until cell adherence. A sterile 20-µL pipette tip was used for uniform scratching and the aspiration of free cell debris. Wound healing was evaluated using a microscope (Nikon, Japan) after 0 h and 48 h.

**RNA extraction and RT-qPCR analysis**

RNA from 40 breast tissue specimens was extracted by TRIzol (Thermo Fisher, USA) and reverse transcribed to cDNA using the
PrimeScript RT kit (Takara, Japan). RT-qPCR was performed using the SYBR Premix Ex Taq kit (Takara, Japan) with primers (Supplementary Table 2) according to the manufacturer’s protocol.

**Western blot**

After cell lysis, protein concentrations were determined by using the BCA protein quantification kit. Protein lysates were electrophoresed on SDS-polyacrylamide gels and then transferred to PVDF membranes. The membranes were then closed with 5% w/v skim milk powder in 1x TBST for 2 h, followed by incubation with primary antibodies, PPP1CA (1:1000, Abcam, UK), PPP4C (1:1000, Abcam, UK) and GAPDH (1:4000, Abcam, UK), at 4°C overnight. Then, the membranes were incubated for 2 h at room temperature with the secondary anti-sheep horseradish peroxidase (HRP)-conjugated antibody (1:5000, Abcam, UK). After the addition of HRP substrate, the membrane fluorescence was checked using an image acquisition and analysis system (Tanon-5220, Tanon Science & Technology, China).

**Statistical analysis**

SPSS 22.0 software (SPSS Inc., Armonk, NY, USA) was used for statistical analysis. In vitro experimental data are presented as the mean ± standard deviation (SD) of 3 independent replicates. Student’s t-test was performed to compare differences. P < 0.05 was considered a significant difference in all circumstances.

**Results**

To investigate whether the PPCPs family changes promoted breast cancer development, we performed bioinformatic analyses and in vitro experiments. For the bioinformatic analyses, we used online public databases to compare the differences in the expression of the PPCPs family members between tumors and non-tumors and the impact of the PPCPs family on breast cancer diagnosis, clinicopathological parameters, prognosis, biological functions and signaling pathways in breast cancer. We also validated the changes we identified by establishing siRNA knockout models for PPP1CA and PPP4C to investigate the biological impact in breast cancer cell culture.

**Transcription levels of the PPCPs family genes in breast cancer**

In the Oncomine database, when compared with normal breast tissues, PPP1CA, PPP2CA, PPP4C and PPEF1 were significantly elevated in breast cancer tissues, while PPP1CB, PPP2CB, PPP3CA, PPP3CB, PPP3CC and PPP6C were significantly decreased in breast cancer tissues. PPP1CC, PPP5C and PPEF2 were not aberrantly expressed in breast cancer tissues (Figure 1, Supplementary Table 3). We verified the above results in the TCGA database and found that PPP1CA, PPP2CA, PPP4C and PPEF1 were significantly expressed at higher levels in breast cancer tissues than in normal tissues. PPP2CB, PPP3CA, PPP3CB, PPP3CC and PPP6C were significantly expressed at lower levels in breast cancer tissues (Figure 1). PPP1CA, PPP2CA, PPP2CB, PPP3CA, PPP3CB, PPP3CC, PPP4C, PPP6C and PPEF1 were selected as the differentially expressed genes (DEGs) for the next analysis.

**The PPCPs family genes alterations and coexpression in breast cancer**

We examined the PPCPs family alterations in the cBioPortal database. Three datasets, breast cancer (METABRIC, Nature 2012 & Nat Commun 2016), breast invasive carcinoma (TCGA, PanCancer Atlas) and breast invasive carcinoma (TCGA, Firehose Legacy, 2015), showed that the PPCPs family variation rates were 16.86%, 24.26% and 26.48%, respectively (Figure 2). Among the gene alterations, PPP1CA and PPP4C gene alterations occurred most frequently (Figure 2). The alterations of PPP1CA, PPP3CA, PPP3CB and PPP6C were significantly associated with the prognosis of breast cancer (Figure S1).

We analyzed the correlation between the PPCPs family members in the GEPIA database using Pearson regression. We identified numerous genes in the PPCPs family that were significantly and positively correlated (Figure 2). The above results suggested that the functions of multiple members of the PPCPs family in breast cancer were significantly related.
Figure 1. Transcription of the PPPCs family in breast cancer. (a) Comparison of the PPPCs family between normal breast tissues and breast cancer tissues using the Oncomine database. Red and blue colors represent relatively high and low expression of mRNA in the corresponding group, respectively. P-value < 0.05, |fold change| > 1.5, and gene ranking in the top 5% were set as the thresholds. (b) Comparison of the PPPCs family between normal breast tissues and breast cancer tissues using the TCGA database. Two paired samples Student’s t-test or the Wilcoxon signed rank test was used to compare the mRNA difference of two paired groups. *P < 0.05, **P < 0.01, ***P < 0.001 and ns: not significant.
Diagnostic accuracy of the DEGs for breast cancer

In analyzing the TCGA database, ROC curve analysis revealed that the area under the curve for the DEGs was significant. Among the highly expressed genes, PPP4C had the highest diagnostic efficacy. PPP1CA and PPEF1 had comparable diagnostic efficacy, and PPP2CA had the lowest diagnostic efficacy. Among the genes with low expression, PPP2CB, PPP3CB and PPP3CC had the highest diagnostic efficacy, while PPP3CA had higher diagnostic efficacy than PPP6C. We found that the PPPCs family had high diagnostic accuracy for breast cancer, with PPP1CA, PPP4C

Figure 2. Gene alterations and coexpression analysis of the PPPCs family in breast cancer. (a-b) Genetic and transcriptional alterations of the PPPCs family in the METABRIC dataset, TCGA (PanCancer Atlas dataset) and TCGA (Firehose Legacy, 2015 dataset) analyzed using the cbioPortal database. (c) Genetic coexpression of the PPPCs family in breast cancer using the GEPIA database. Darker colors indicated stronger associations. Linear dependence was measured using Pearson’s correlation coefficient. P-value <0.05 was set as the thresholds.
and PPEF1 having the highest diagnostic accuracy (Figure 3, Supplementary Table 4–5).

**Correlations between the DEGs and the clinical and pathological parameters of breast cancer**

For clinicopathological parameter parameters, PPP2CA and PPEF1 had higher mRNA expression levels, and PPP3CA had lower mRNA levels in patients over 51 years of age. The lymph node metastasis rate was positively correlated with PPP1CA and PPP4C expression levels and negatively correlated with PPP2CB expression levels. ER-positive status was positively correlated with PPP2CA, PPP2CB, PPP3CA, PPP3CB, PPP4C, PPP6C and PPEF1. PR-positive status was positively correlated with PPP2CA, PPP2CB, PPP3CA, PPP3CB and PPP6C. HER-2-positive status was positively correlated with PPP1CA and PPEF1 and negatively correlated with PPP2CA, PPP2CB, PPP3CB, and PPP3CC (Supplementary Table 6). Based on these results, we found that the DEGs were strongly associated

![Figure 3](image-url)
with factors related to breast cancer prognosis, especially the lymph node metastasis rate.

**Correlations between the DEGs and the prognosis of breast cancer**

Since the PPPCs family for breast cancer had an impact on the type of clinicopathology, we subsequently analyzed whether it affected the prognosis of breast cancer patients. In the bc-GenExMiner database, poor overall survival (OS) was associated with high expression of PPP1CA, PPP4C and PPEF1 and low expression of PPP2CB, PPP3CA, PPP3CB and PPP3CC (Figure 4, Supplementary Table 7). In the K-M plotter database, we found that the results were consistent with the above results, except that PPP2CA and PPP3CA were not associated with poor OS (Figure S2, Supplementary Table 8). From the above results, we found that the overexpressed PPP1CA and PPP4C had the highest hazard ratio (HR) with poor OS, while among the genes with low expression, PPP2CB had the highest HR. In addition, we analyzed the DEGs with DMSF in these databases (Supplementary Tables 7–8). The above results

**Figure 4.** Prognostic analysis of the PPPCs family for breast cancer. The correlations between the PPPCs mRNA expression levels and overall survival in breast cancer of the Affymetrix and METABRIC datasets were calculated using the bc-GenExMiner database. The results are displayed in Kaplan-Meier survival plots. Hazard ratios and 95% confidence levels were calculated automatically by the web tool. P-values <0.05 were set as the thresholds.
showed that the DEGs were significantly associated with breast cancer prognosis.

**The mRNA expression levels and protein levels of PPP1CA and PPP4C were significantly higher in breast cancer tissues**

We examined the expression of PPP1CA and PPP4C, in clinical samples. The mRNA expression level of PPP1CA in 40 breast cancer patients and the protein expression level of PPP1CA in 60 breast cancer patients were significantly higher in breast cancer tissues than in paracancerous tissues (Figure 5). Similarly, the mRNA expression level of PPP4C in 40 breast cancer patients and the protein expression level of PPP4C in 60 breast cancer patients were significantly higher in breast cancer tissues than in paracancerous tissues (Figure 5).

**Significantly reduced proliferation and migration of breast cancer cells after the inhibition of PPP1CA or PPP4C expression**

After intervention with siRNA in breast cancer cells, the mRNA expression levels and protein expression levels of PPP1CA or PPP4C in MCF-7 and MD-MB-468 cell lines were significantly decreased (Figure 6). The relative cell viability of the MCF-7-siPPP1CA or MDA-MB-468-siPPP1CA group was significantly lower than that of the MCF-7-siPPP1CA-Control group (Figure 6) or MDA-MB-468-siPPP1CA-Control group (Figure 6). The wound closure area of the MCF-7-siPPP1CA or MDA-MB-468-siPPP1CA group was significantly lower than that of the MCF-7-siPPP1CA-Control group (Figure 6) or MDA-MB-468-siPPP1CA-Control group (Figure 6). The relative cell viability of the MCF-7-siPPP4C or MDA-MB-468-siPPP4C group was significantly lower than that of the MCF-7-siPPP4C-Control group (Figure 6) or MDA-MB-468-siPPP4C-Control group (Figure 6). The wound closure area of the MCF-7-siPPP4C or MDA-MB-468-siPPP4C group was significantly lower than that of the MCF-7-siPPP4C-Control group (Figure 6) or the MDA-MB-468-siPPP4C-Control group (Figure 6).

**Potential biological functions and related signaling pathways of the DEGs in breast cancer**

We performed functional analysis of the DEGs and their analogues (Supplementary Table 9).

![Figure 5. PPP1CA and PPP4C expression in human breast cancer tissues. The mRNA levels (a) and protein levels (b) of PPP1CA in breast cancer tissues and paracancerous tissues. The mRNA levels (c) and protein levels (d) of PPP4C in breast cancer tissues and paracancerous tissues. The data are presented as the means (minimum-maximum). All data are representative of three independent experiments. *P < 0.05, **P < 0.01, ***P < 0.001.](image-url)
According to GO term analysis, these genes were mainly related to ubiquitin-dependent protein processes, DNA repair and circadian rhythms. KEGG pathway analysis showed that these genes were enriched in membrane trafficking, oocyte meiosis, signaling by WNT in cancer and class I MHC-mediated antigen processing and presentation (Figure 7). Based on these results, these genes were shown to play important roles in biological functions and signaling pathways closely related to breast cancer development. In addition, Cytoscape was used to determine the relationship of enriched terms and to build a network diagram (Figure 7).
Correlations between the DEGs and immune cell infiltration of breast cancer

Immune cell infiltration was closely associated with the development of breast cancer. Given that the PPPCs family was associated with immune system diseases [19] and immune system tumors [20], we analyzed whether these factors affected the immune cell infiltration of breast cancer tissues. Among the highly expressed genes in

Figure 7. Top 20 GO and KEGG enriched terms of the DEGs. (a) Heatmap of GO and KEGG analyses of the DEGs and their 70 most analogous genes, with Orange representing enrichment terms colored by -log10(P-value). (b) Interaction network of the top enrichment terms colored by cluster ID, with different colors representing different enrichment pathways of these genes. The thresholds for Min Overlap, P-value and Min Enrichment in the Metascape database were set to 3, 0.05 and 3, respectively. The item with the greatest statistical significance within the cluster was selected as the item representing that cluster.
breast cancer tissues, PPP1CA was negatively correlated with CD8+ T cell and macrophage infiltration, while PPP2CA was positively correlated with infiltration. PPP4C expression was negatively associated with CD8+ T cell, macrophage, neutrophil and dendritic cell infiltration. PPEF1 expression was positively correlated with CD8+ T cell, macrophage and dendritic cell infiltration. Among the genes with low expression in breast cancer tissues, PPP3CB and PPP6C were positively associated with CD8+ T cell, macrophage and neutrophil infiltration. PPP2CB was positively correlated with CD4+/CD8+ T cell, macrophage and neutrophil infiltration; PPP3CA was positively correlated with CD8+ T cell, macrophage, neutrophil and dendritic cell infiltration, and PPP3CC was positively correlated with B cell, CD4+/8+ T cell, macrophage, neutrophil and dendritic cell infiltration (Figure 8). With these results, we suggest for the first time that the DEGs of the PPPCs family, except for PPP2CA and PPEF1, lead to reduced infiltration of immune cells in breast cancer tissues, which can affect the development and immunotherapy of breast cancer.

**Discussion**

Previous studies have demonstrated that the PPPCs family plays essential roles in tumor cell proliferation [21], metastasis [22] and resistance to chemotherapy [23]. However, the significance of the PPPCs family in the development of breast cancer is still largely unknown. Therefore, we integrated several publicly available data into one comprehensive analysis to explore the diagnostic, prognostic and therapeutic value of the PPPCs family in breast cancer for the first time.

We first examined the expression levels of the PPPCs family members in breast cancer and their correlation with clinicopathological parameters of breast cancer. In the Oncomine and TCGA databases, there were nine genes among the PPPCs family that were differentially expressed in breast cancer tissues compared with normal breast tissues (elevated expression of PPP1CA, PPP2CA, PPP4C and PPEF1 and reduced expression of PPP2CB, PPP3CA, PPP3CB, PPP3CC and PPP6C). Abnormal expression of the PPPCs family was associated with older patients and a higher rate of lymph node metastasis of breast cancer, both of which were independent risk factors for worse prognosis in breast cancer [24], suggesting that the PPPCs family may be closely associated with the development and prognosis of breast cancer. To verify the credibility of these results, we analyzed the prognostic impact of the DEGs in breast cancer tissues and found that abnormal expression of these genes was strongly associated with breast cancer prognosis.

As the emergence and progression of breast cancer presents a multistep and multifactorial process involving progressive accumulation of genetic, epigenetic and microenvironmental alterations [25,26], we further developed a comprehensive analysis of the alterations of the PPPCs family in breast cancer.

Analysis of data obtained from the cBioPortal database showed that alterations of the PPPCs family were found in breast cancer. Alterations in mRNA expression, amplification and profound deletions were also commonly observed. Alterations in multiple genes in the PPPCs family were associated with breast cancer prognosis. These data confirmed that cumulative genetic alterations of the PPPCs family were involved in the development and progression of breast cancer.

In the above data, we found that among the PPPCs family, PPP1CA and PPP4C played the most significant roles in the development of breast cancer. We therefore further analyzed the functions of these two genes. PPP1CA is one of the three isoforms of PPP1C. PPP1CA was associated with aggressive metastasis and poor prognosis in a variety of tumors [27]. PPP1CA promoted tumor cell proliferation and metastasis by activating the MAPK signaling pathway [28]. This result could partially explain why the bc-GenExMiner database indicated a high rate of lymph node metastasis in patients with high PPP1CA expression. In addition, PPP1CA can bind to cyclin D1 to phosphorylate RB [29] or can dephosphorylate breast cancer susceptibility protein-1 (BRCA1) [30], all of which induce cell cycle deregulation and promote tumor cell proliferation. Therefore, inhibition of PPP1CA may play an important role in inhibiting breast cancer proliferation and metastasis. Especially for triple-negative breast
Figure 8. Relationship between the expression levels of the DEGs and immune cell infiltration. The correlations between the mRNA expression levels of the DEGs and immune cell infiltration were calculated by the TIMER database. |Partial correlation|>0.1 and P-value <0.05 were set as the thresholds.
cancer (TNBC) patients with high mutation of BRAC1 [31], a promising oncogenic effect may be achieved by inhibiting PPP1CA. PPP4C is a ubiquitous serine/threonine phosphatase that affects DNA repair function through nonhomologous end joining (NHEJ) and homologous recombination (HR) to affect DNA repair functions [32]. High PPP4C expression can improve tumor resistance to platinum by improving nuclear factor kappa B subunit 1 (NF-κB) expression [33], and inhibition of PPP4C leads to sustained phosphorylation and ubiquitination of γ-microtubule proteins, resulting in blocked microtubule nucleation, which enhances the anticancer effect of paclitaxel [34]. In addition, PPP4C promotes the expression of matrix metallopeptidase (MMP)-2 and MMP-9 through the PI3K/AKT signaling pathway, thus promoting the invasion and metastasis of cancer cells [35].

To verify that PPP1CA and PPP4C indeed promote the development of breast cancer, we conducted in vitro experiments on these genes. We examined the mRNA and protein expression levels of PPP1CA and PPP4C in clinical samples and showed that both genes were expressed at significantly higher levels in cancer tissues than in noncancerous tissues. Inhibition of these genes in the TNBC cell line MDA-MB-231 and the non-TNBC cell line MCF-7 can inhibit the proliferation and migration of these cells. We therefore concluded that both PPP1CA and PPP4C have a significant effect on the proliferation and metastasis of breast cancer, but whether these genes lead to a worse prognosis in breast cancer needs to be supported by more clinical data.

To investigate the mechanisms by which the PPCs family influenced the development of breast cancer, we investigated the biological functions of the DEGs in breast cancer tissues. We found that the DEGs were mostly located in the cell membrane and nucleus and involved in DNA repair [36] and several signaling pathways related to extracellular signal transduction, such as the WNT signaling pathway [37]. These biological functions and signaling pathways were closely related to the development of breast cancer. In addition, we found that the biological functions of the DEGs were related to circadian rhythm, which affected the development of breast cancer mainly by influencing ER signaling networks and DNA repair [38]. Interestingly, we found that ER transport and DNA repair functions were enriched in the DEGs. This finding suggested that the DEGs also influence the development of breast cancer through circadian rhythms. Future studies on these functions could unravel the mechanisms by which aberrant expression of the PPCs family promotes breast carcinogenesis.

According to our results, several members of the PPCs family were closely associated with breast cancer prognosis, and inhibition of the expression of these genes may offer the possibility of improving the survival of breast cancer patients. We demonstrated in vitro that inhibition of PPP1CA and PPP4C expression significantly inhibited breast cancer proliferation and migration. Thus, chemotherapy or surgery combined with specific the PPCs-targeted drugs may improve the therapeutic outcome of breast cancer. In addition, we found that aberrant expression of the PPCs family members can inhibit the infiltration of immune cells in breast cancer; therefore, using specific the PPCs-targeted drugs to promote immune cell infiltration in breast cancer may enhance the effectiveness of immune checkpoint inhibitors in treating breast cancer [39].

Our study has some limitations. Since the data used for the analysis were obtained from multiple online bioinformatics sources, although we used multiple databases for iterative validation, there may still be bias due to confounding factors. Although we validated the expression levels of PPP1CA and PPP4C in breast cancer tissues and their effects on the biological behavior of breast cancer cells, further investigation of their molecular mechanisms is needed. Furthermore, we did not validate the role of other PPCs family members in breast cancer or the PPCs family in different molecular subtypes of breast cancer, which should be fully explored in subsequent studies.

Conclusions

In our study, the diagnostic value, prognostic value and biological functions of the PPCs family were comprehensively assessed. The DEGs of the PPC
family in breast cancer were associated with clinicopathological parameters and prognosis, which may be related to effects on the WNT signaling pathway, antigen expression, circadian rhythm and immune cell infiltration in breast cancer tissues. In addition, the detection of the DEGs can play a positive role in improving the accuracy of breast cancer diagnosis. Based on the above findings, the PPPCs family, especially PPP1CA and PPP4C, could be the most promising diagnostic and prognostic biomarkers and therapeutic targets for breast cancer.

**Highlights**

- The PPPCs family was significantly differentially expressed in BC tissues.
- The PPPCs family correlates with the lymph node metastasis rate of BC.
- The PPPCs family correlated with prognosis and immune cell infiltration in BC.
- PPP1CA and PPP4C may be prognostic biomarkers and therapeutic targets of BC.

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**Disclosure statement**

No potential conflict of interest was reported by the author(s).

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**Data availability statement**

All the data pertaining to this work is available within the text and no additional data is required.

**Ethical statement**

The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

**CRedit authorship contribution statement**

Conceptualization, Wenjun Xie, Xianhui Ruan and Ming Gao; Data curation, Jingtai Zhi; Formal analysis, Wenjun Xie, Yu Zeng, Linfei Hu, Hang Ling and Ming Gao; Funding acquisition, Ying Sun, Xiangqian Zheng and Ming Gao; Investigation, Wenjun Xie; Methodology, Wenjun Xie; Project administration, Ming Gao; Resources, Xiangqian Zheng and Ming Gao; Software, Wenjun Xie, Yu Zeng and Jingtai Zhi; Supervision, Xiangqian Zheng and Ming Gao; Validation, Wenjun Xie, Ying Sun and Yu Zeng; Visualization, Linfei Hu, Jingtai Zhi, Hang Ling and Xiangqian Zheng; Writing – original draft, Wenjun Xie, Ying Sun and Yu Zeng; Writing – review & editing, Xianhui Ruan.

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