Protein tyrosine phosphatase SHP-1 sensitizes EGFR/HER-2 positive breast cancer cells to trastuzumab through modulating phosphorylation of EGFR and HER-2

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Background: Trastuzumab resistance in HER-2 positive breast cancer cells is closely related to overexpression of both epidermal growth factor receptor (EGFR) and human epidermal receptor (HER-2). SHP-1 has been demonstrated to downregulate tyrosine kinase activity including EGFR via its phosphatase function, but its effect on HER-2 activity is still unknown. Here, we examined the hypothesis that SHP-1 enhances the anticancer efficacy of trastuzumab in EGFR/HER-2 positive breast cancer cells through combining dual inhibition of EGFR and HER-2.

Methods: Trastuzumab-resistant breast cancer SKBr-3 cells were generated by long-term in vitro culture of SKBr-3 cells in the presence of trastuzumab. The SHP-1 was ectopically expressed by stable transfection. The activity and expression of EGFR, HER-2, and downstream signaling pathways were tested by Western blot. Cell viability was examined by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay, and apoptosis was examined by flow cytometry. The binding between SHP-1 and EGFR/HER-2 was evaluated by immunoprecipitation assay and bimolecular fluorescence complementation. The effects of SHP-1 on tumorigenicity and trastuzumab sensitivity were confirmed via in vivo xenograft model.

Results: Trastuzumab-resistant SKBr-3 cells showed aberrant co-expression of EGFR and HER-2. Introduction of wild-type SHP-1 inhibited cell proliferation, clone formation, and promoted the apoptosis induced by trastuzumab. Meanwhile, SHP-1 overexpression reduced phosphorylation levels of EGFR and HER-2 both in parental and trastuzumab-resistant SKBr-3 cells. In vivo study showed an increased antitumor effect of trastuzumab in SHP-1 overexpressed xenografts. At last, we discovered that SHP-1 can make complexes with both EGFR and HER-2, and both phospho-EGFR and phosphor-HER-2 levels in wild-type SHP-1 immunoprecipitates were less than those in phosphatase-inactive SHP-1 (C453S) immunoprecipitates, indicating that EGFR and HER-2 are potential substrates of SHP-1.

Conclusion: Taken together, we have demonstrated that the SHP-1 is a negative regulatory factor of the tyrosine kinase activity of HER-2 and EGFR through inhibiting phosphorylation. Dual targeting of EGFR and HER-2, by combining trastuzumab with SHP-1 overexpression, may improve response in HER-2 overexpressing breast cancer cells that also express high levels of EGFR.

Keywords: breast cancer, trastuzumab, EGFR, HER-2, SHP-1, drug resistance

Background
Breast cancer is a leading cause of cancer-related death in women; it is estimated that there will be 39,620 breast cancer related deaths among women in the United States1 and 88,886 in Europe2 in 2013. HER-2 is a transmembrane tyrosine kinase
receptor that belongs to the human epidermal growth factor receptor (EGFR/HER) family, which activates various intracellular downstream oncogenic signaling pathways. HER-2 amplification occurs in approximately 20%-30% of patients with breast cancer, and it correlates with more aggressive biological features of tumor such as poor clinical outcome, high rates of metastasis, and resistance to conventional treatment modalities. Trastuzumab (Herceptin®), a humanized recombinant monoclonal antibody that binds to the extracellular domain of HER-2, has been recommended as first-line treatment for HER-2 positive patients and shown to improve outcome. However, not all patients with HER-2 positive respond to trastuzumab treatment; nearly 15% of patients will relapse with distant disease within 12 months of treatment with trastuzumab, and only about 26% of HER-2-overexpressing metastatic breast cancer patients showed response to trastuzumab monotherapy as first-line treatment. The innate and acquired trastuzumab resistance has increasingly occurred and remains the biggest clinical challenge. Epidermal growth factor receptor (EGFR) is overexpressed in 35%-65% of HER-2-positive breast cancers. There is evidence that the EGFR overexpression influences response to HER-2 inhibition, impairs response to trastuzumab in HER-2 positive breast cancers, and correlates with poor survival. Therefore, combining trastuzumab with EGFR-targeted strategy may improve response to trastuzumab treatment. It is well known that phosphorylation of proteins at specific tyrosine residues has an important role in regulating the activity of EGFR/HER family proteins and downstream pathways. Biologically, the de-phosphorylation of tyrosine residues in proteins is precisely regulated by protein tyrosine phosphatases (PTPs), and defects in PTPs can lead to the intracellular accumulation of tyrosine-phosphorylated proteins, which can cause abnormal cell proliferation and differentiation. HERs are major substrates of PTPs in breast cancer cells, and ectopic expression of some PTPs in human breast cancer cells decreased cell proliferation and tumorigenicity, which suggests their role as a tumor suppressor. In consideration of the dual inhibitory effect of PTPs on both EGFR and HERs, it is reasonable to speculate that some members of PTPs might be potential therapy targets and markers. SHP-1, a non-receptor PTP containing an SH2 domain, is a key regulator that controls the intracellular phosphotyrosine level. Extensive studies on SHP-1 revealed that the expression of SHP-1 protein was decreased or absent in many kinds of cancer cell lines and tissues. Abolished expression in leukemia and lymphoma has been related to both malignant transformation and malignant tumor behavior. Ectopic expression of SHP-1 in the breast cancer cell line MDAMB-231 impaired cell proliferation two- to threefold. In addition, SHP-1 has also been shown to be involved in regulating several signals including EGFR and other tyrosine kinase receptors. The findings mentioned earlier suggests a potential key role of SHP-1 in resistance to trastuzumab, but the role of SHP-1 in trastuzumab resistance remains unclear. The aim of this study is to elucidate the role of SHP-1 protein expression in relation to trastuzumab treatment and resistance in HER-2 positive breast cancer.

Materials and methods

Cell culture

The human breast cancer SKBr-3 and human embryonic kidney 293 (HEK293) cell lines were obtained from American Type Culture Collection (Manassas, VA, USA). SKBr-3 cells were cultured in RPMI 1640 media (Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 10% fetal bovine serum and HEK293 cells were cultured in DMEM high glucose medium (Thermo Fisher Scientific) containing 10% fetal bovine serum in a humidified 5% CO2 atmosphere at 37°C.

Establishment of trastuzumab-resistant cells

Trastuzumab/Herceptin (Hoffman-La Roche Ltd., Basel, Switzerland) was dissolved in sterile water. Trastuzumab-resistant cells were generated through continuous culture of SKBr-3 cells in the presence of 5 μg/mL trastuzumab for 6 months, as reported previously. Subsequently, the established trastuzumab-resistant SKBr-3 cells were cultured in medium containing 5 μg/mL trastuzumab.

Plasmids and virus infection

SHP-1 cDNA was amplified by polymerase chain reaction (PCR) from total RNA extracted from SHP-1 overexpressed MCF-7 cells and ligated into pCDNA3.1. The catalytically inactive (C453S) SHP-1 was generated by introducing a point mutation using standard PCR techniques. Lentiviral particles were produced by transfecting 239T cells, the viral supernatants were collected after 72 hours following transfection, and the aliquots were stored at -80°C. The titers of the concentrated virus particles were measured before using. For bimolecular fluorescence complementation (BiFC), the pBiFC-VC155 plasmid (#22011, deposited by Chang-Deng Hu) and
pBiFC-VN173 plasmid (#22010, deposited by Chang-Deng Hu) were obtained from Addgene (Cambridge, MA, USA). The SHP-1 gene was amplified from MCF-7 cells, and the PCR product was digested with KpnI and EcoRI, and then ligated into pBiFC-VN173. EGFR, HER-2, HER-3, and HER-4 genes were amplified from total RNA extracted from SKBr-3 cells and ligated into pBiFC-VC155.

**Cell proliferation analysis**

MTT substrate (Sigma-Aldrich, St Louis, MO, USA) was used to assay cell proliferation according to the manufacturer’s instructions. Briefly, a total of 5×10^3 cells were seeded into 96-well dishes and allowed to adhere overnight; trastuzumab of indicated concentration was then added into the medium. The medium was replaced by 100 μL fresh serum-free medium containing 0.5 g/L MTT 24 hours later, followed by 4-hour incubation. Then the MTT medium was replaced by 50 μL of DMSO for further 10 minutes incubation at 37°C. The A490 value of each sample was measured.

**Colony formation assay**

For colony formation assay, transfected cells were placed in a fresh six-well plate with a density of 500 cells per well and maintained in DMEM medium containing 10% fetal bovine serum and trastuzumab of indicated concentration at 37°C for 10 days. Colonies were fixed with methanol and stained with 0.1% crystal violet and counted.

**Quantitative reverse transcription PCR analysis**

Total RNA was extracted from each cell line using TRIzol reagent (Thermo Fisher Scientific) according to the manufacturer’s protocol. Reverse transcription was performed using M-MLV reverse transcriptase (Thermo Fisher Scientific) and quantitative PCR was performed with SYBR Green Master Mix (Hoffman-La Roche Ltd.). Relative expression values were calculated (ΔΔCT method) using GAPDH as a normalizer. The PCR primers sequences were:

- **SHP-1**, forward, 5'-GGAGAAGTTCTGCACTC TGAC-3'; reverse, 5'-GCCGGTACTTGAGGTGGATG-3';
- **EGFR**, forward, 5'-CCACTCATGCTCTACAACCC-3'; reverse, 5'-TCGCACCTTCTACCTGCGG-3';
- **HER-2**, forward, 5'-TGCAGGGAAACCTGGAACTC-3';
- **HER-3**, reverse, 5'-GCGGGTACTTGAGGTGGATG-3';
- **HER-4**, forward, 5'-ACAGGGGTGTATTGTCAGC-3'.

**Flow cytometry**

Cells were plated into six-well plates at a density of 3×10^5 cells per well, and 24 hours later cells were treated with trastuzumab of indicated concentrations. After 24 hours exposure to trastuzumab, cells were stained with Annexin V-FITC and propidium iodide, and then flow cytometry was performed to detect apoptosis of the transfected cells.

**Immunoblotting**

Cells were treated as indicated earlier and were placed on ice and washed with PBS. Washed cells were lysed in RIPA lysis buffer (50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1% sodium deoxycholate, 1% Triton X-100, and 0.1% SDS) plus protease inhibitor cocktail (Hoffman-La Roche Ltd.). Equal amounts of protein sample were subjected to sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS–PAGE), transferred to polyvinylidene difluoride (PVDF) membrane (EMD Millipore, Billerica, MA, USA). Membranes were blocked with Tris-buffered saline containing Tween-20 and 5% (w/v) fat-free milk powder, and then incubated with appropriate primary antibodies at 4°C overnight followed by horseradish peroxidase-conjugated secondary antibodies. All antibodies were purchased from Cell Signaling Technologies: phospho-HER2 (Tyr1248) rabbit polyclonal antibody (#2247), HER2 rabbit polyclonal antibody (#2242), phospho-EGFR (Tyr1173) rabbit monoclonal antibody (#4407), EGFR rabbit monoclonal antibody (#4405), phospho-Erk1/2 (Thr202/Tyr204) rabbit monoclonal antibody (#4370), Erk1/2 rabbit monoclonal antibody (#4695), phospho-Stat3 (Tyr705) rabbit monoclonal antibody (#9145), Stat3 rabbit monoclonal antibody (#4904), phospho-Akt (Ser473) rabbit monoclonal antibody (#4060), and Akt (pan) rabbit monoclonal antibody (#4685).

Quantification and comparison of the intensity of bands of Western blot was done by using ImageJ software. Briefly, the images of Western blot results were converted to grayscale, the background were subtracted, the blot areas were selected, and the corresponding area and mean gray value (GV) were recorded. The GV intensity = area × mean GV. The GV of each protein was normalized to that of GAPDH, and the GV of each phosphorylated protein was normalized to the GV of its own corresponding total protein.

**Co-immunoprecipitation**

The co-immunoprecipitation experiment was carried out with the Pierce® Co-Immunoprecipitation Kit according to the manufacturer’s protocol using the following antibodies: EGF Receptor (D38B1) XP® Rabbit mAb (#4267, Cell Signaling Technology, 1:500), HER2 (D8F12) XP® Rabbit mAb (#4290, Cell Signaling Technology, 1:500), HER3 (1B2E) Rabbit mAb (#4754, Cell Signaling Technology,

In vivo tumor growth and mouse survival assay
BALB/c nu/nu mice were randomly grouped to monitor tumor growth (n=10) and mouse survival (n=20). Trastuzumab-resistant SKBr-3 cells stably transfected vectors expressing SHP-1 or blank vectors and were subcutaneously injected (2×10⁶) into the right mammary fat pad of each nude mice. After 1 week, mice were intravenously injected with 10 mg/kg trastuzumab twice a week. Tumor size was measured with the following formula: volume = (L×W²)/2, where L and W are the longest and shortest diameters, respectively. Mice were sacrificed 45 days post-first trastuzumab injection, and tumors were separated for weighing. For survival assay, the survival of mice in each group was recorded, and the ratios of surviving mice were plotted. All experimental procedures involving animals were in accordance with the Guide for the Care and Use of Laboratory Animals and were performed according to the institutional ethical guidelines for animal experiment. The study protocol was also approved by the Committee on the Use of Live Animals in Teaching and Research, Southern Medical University, Guangzhou, Guangdong, People’s Republic of China.

Results
EGFR and HER-2 are aberrantly expressed and phosphorylated in trastuzumab-resistant cells.

The HER-2 overexpressed human breast cancer SKBr-3 cells were cultured in medium containing 5 μg/mL trastuzumab continuously for 6 months, which resulting in the acquisition of trastuzumab resistance. Compared with the parental SKBr-3 cells, MTT assay showed that the trastuzumab-resistant cells displayed a significantly higher viability or proliferative capacity in the presence of every different concentration of trastuzumab (Figure 1A). Besides, the resistant SKBr-3 cells displayed dramatically increased colony formation ability under the treatment of trastuzumab (Figure 1B). We next assessed EGFR and HER-2 expression in the parental cells and trastuzumab-resistant cells, and found that both the mRNA and protein levels of EGFR and HER-2 were increased in the trastuzumab-resistant cells (Figure 1C and D). Results also showed that the phosphorylation levels of EGFR and HER-2 as well as the key kinases of their downstream pathways which including STAT3, Akt, and ERK were significantly enhanced (Figure 1D and E).

SHP-1 overexpression promotes trastuzumab sensitivity in breast cancer cells

To further investigate the role of SHP-1 in trastuzumab resistance of breast cancers, stable SHP-1 overexpressing SKBr-3 cells were established through retrovirus infection. In order to assess the role of the phosphatase activity of SHP-1 in the regulation of trastuzumab resistance, vectors expressing a catalytic domain mutant (C453S) of SHP-1 (SHP-1 mutant-type, SHP-1 MT) were established subsequently. We first treated cells with different concentrations of trastuzumab, and evaluated cell viability by using MTT assay. Results showed that, compared with cells expressing control scramble shRNA, both parent cells and resistant cells with wild-type SHP-1 overexpression displayed enhanced sensitivity to trastuzumab (Figure 2A); as expected, cells transfected with mutant-type SHP-1 showed similar trastuzumab sensitivity with controls. We also examined the apoptosis of cells after treatment with 5 μg/mL trastuzumab for 24 hours. Wild-type SHP-1 overexpression caused a significant increase in apoptosis of both parental cells and trastuzumab-resistant cells (Figure 2B). Notably, trastuzumab-resistant cells with overexpression of SHP-1 displayed a parallel trastuzumab sensitivity compared with SHP-1 overexpression parental cells. In addition, overexpression of wild-type SHP-1 also significantly suppressed in vitro colony formation by both the parental and trastuzumab-resistant cells (Figure 2C). At last, we investigated the effect of SHP-1 overexpression on the expression levels and activity of EGFR and HER-2 proteins through Western blot analysis. We found that the protein levels of EGFR and HER-2, as well as their phosphorylation levels, were significantly decreased in cells stably expressing wild-type SHP-1 compared with controls (Figure 2D and E).

SHP-1 binds to EGFR and HER-2 proteins

To verify the interactions between SHP-1 and EGFR, as well as HER-2, we conducted immunoprecipitation studies in order to ascertain whether SHP-1 and EGFR could be coprecipitated — an indicator of their co-association or heterodimerization. Anti-EGFR and anti-HER-2 antibodies were used to detect EGFR and HER-2 proteins through Western blot analysis. We found that the protein levels of EGFR and HER-2, as well as their phosphorylation levels, were significantly decreased in cells stably expressing wild-type SHP-1 compared with controls (Figure 3A and B). We also detected the interaction between SHP-1 and phosphorylated EGFR (pEGFR) or pHER-2 in anti-SHP-1 immunoprecipitates. As expected, the corresponding bands were observed (Figure 3C and D). Notably, both pEGFR and pHER-2 protein levels...
in the wild type SHP-1 immunoprecipitates were less than those of C453S SHP-1. Besides, we also examined whether SHP-1 could interact with HER-3 and HER-4. We did not detect any specific band by using anti-HER-3 (Figure 3E) and anti-HER-4 antibodies (Figure 3F). The interaction between SHP-1 and both EGFR and HER-2 was further investigated by using BiFC. Cells were cotransfected with pBiFC-VC155-EGFR vectors and pBiFC-VN173-SHP1 vectors, and green
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Figure 2 Overexpression of SHP-1 partly restores trastuzumab sensitivity in vivo.

Notes: (A) MTT assays of modified SKBr-3 cells after treatment with increasing concentrations of trastuzumab. (B) Parental and resistant cells were treated with 5 μg/mL trastuzumab for 24 hours. Apoptosis rate in response was evaluated by using flow cytometry assay. (C) Plate colony formation assays of modified parental and trastuzumab-resistant cells were performed in the presence of 5 μg/mL trastuzumab. (D) Parental and resistant SKBr-3 cells were treated with 5 μg/mL trastuzumab, and then the total or phosphorylated EGFR and HER-2 protein levels were determined by Western blot. (E) Quantification and comparing the gray value of Western blot results between parental group and trastuzumab-resistant group. The gray value of each protein was normalized to that of GAPDH, and the GV of each phosphorylated protein was normalized to the GV of its own corresponding total protein before comparing, the gray value of proteins in NC group were set as 1. Error bars represent the mean ± SD of triplicate experiments. Statistical significance was calculated using the Student’s t-test or ANOVA tests. ***P<0.001.

Abbreviations: MTT, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; EGFR, epidermal growth factor receptor; HER-2, human epidermal receptor; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; GV, gray value; NC, nonsense control; SD, standard deviation; ANOVA, analysis of variance; OD, optical density; WT, wild-type; MT, mutant-type.
Figure 3 SHP-1 binds to EGFR and HER-2 protein.

Notes: (A, B) SKBR3 cells were transfected with control (NC), SHP-1 WT, or SHP-1 MT vector for 48 hours. EGFR and HER-2 proteins were pulled down by immunoprecipitation and the protein level was detected by immunoblotting. The whole gel is shown, and no specific band of HER-3 or HER-4 was detected.

Abbreviations: EGFR, epidermal growth factor receptor; HER-2, human epidermal receptor; NC, nonsense control; WT, wild-type; IP, immunoprecipitation.
fluorescence was detected (Figure 3G), which indicated protein interactions between EGFR and SHP-1. Fluorescent was also detected in cells cotransfected with pBiFC-VC155-HER2 vectors and pBiFC-VN173-SHP1 vectors (Figure 3E). However, we did not detect fluorescence in cells cotransfected with pBiFC-VN173-SHP1 and either pBiFC-VC155-HER3 or pBiFC-VC155-HER4 (Figure 3G).

**SHP-1 promotes trastuzumab sensitivity in vivo**

To further investigate the effect of SHP-1 on the trastuzumab resistance in vivo, xenograft models were generated using trastuzumab-resistant SKBr-3 cells modified to overexpress SHP-1 or control vectors. Compared with mice bearing tumors derived from SKBr-3 cells expressing control vectors, mice inoculated with SHP-1-modified SKBr-3 cells displayed significantly suppressed tumor development and growth (Figure 4A–C). Besides, we also evaluated the effect of SHP-1 overexpression on the survival of mice with tumor burden. A Kaplan–Meier survival analysis showed a better overall survival in mice bearing tumors developed from cells overexpressing SHP-1, compared with those injected with the control cells after treatment with trastuzumab (P=0.0036, Figure 4D). These data suggest that the overexpression of SHP-1 may sensitize trastuzumab-resistant breast cancers to trastuzumab in vivo.

**Discussion**

Over the past decade, the humanized monoclonal erbB2/HER-2 antibody trastuzumab (Herceptin) has been successfully used for clinical treatment of patients with HER-2-positive breast cancers. Nevertheless, primary or acquired resistance to this antitumor antibody has become the major obstacle to its clinical efficacy. Here, we provided insight into the role of SHP-1 in the treatment of trastuzumab-resistant breast cancer. We showed that increased expression of SHP-1, a phosphatase, induces dephosphorylation of both EGFR and HER-2 and contributes to trastuzumab sensitivity of HER-2-positive, trastuzumab-resistant breast cancer cells.

Human EGFR 1 is a key member of the HER signaling pathway that may be associated with trastuzumab resistance.

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Figure 4 SHP-1 promotes trastuzumab sensitivity in vivo.

Notes: Nude mice were inoculated in the mammary fat pad with trastuzumab-resistant SKBr-3 cells overexpressing SHP-1 or control scramble shRNA to allow tumor development. Mice were intravenously injected with 10 mg/kg trastuzumab twice a week. (A) One cohort of mice consisting five individuals in each treatment group were sacrificed after 45 days after first trastuzumab injection. (B) Tumor volume was recorded every 3 days, and (C) tumors were weighted and compared at last. (D) For survival assay, the survival of the other cohort containing ten mice in each group was recorded. Kaplan–Meier survival curves of the trastuzumab-treated mice are shown. ***P<0.001.

Abbreviations: shRNA, small hairpin RNA; NC, nonsense control.
In earlier research, Ritter et al report an association between active involvement of EGFR initiated signal pathways and trastuzumab resistance; they determined that EGFR activation is causally connected to HER-2 phosphorylation, and the trastuzumab resistance cells exhibits higher levels of p-EGFR and EGFR/HER-2 heterodimers. They draw the conclusion that EGFR activation is a potential significant mechanism for sustaining the proliferation of trastuzumab-resistant breast cancer cells which continue to proliferate even in the presence of trastuzumab. Several previous cell-line model studies have also reported that overexpression and activation of EGFR induces trastuzumab resistance in HER-2-positive cell lines. Most recently, a randomized trial (NCCTG N9831) showed that high expression of EGFR appears to be associated with decreased benefit from adjuvant concurrent trastuzumab. These findings are supported by evidence showing that such trastuzumab-resistant breast cancer cells showed sensitivity in the presence of EGFR tyrosine kinase inhibitor. Ritter et al also reported that phosphorylation of HER-2 was inhibited by EGFR-specific concentrations of the EGFR tyrosine kinase inhibitors. The mentioned earlier findings emphasize the strategy of inhibiting EGFR activation in the treatment of trastuzumab-resistant cases.

The Src-homology PTP SHP-1 is a non-receptor phosphatase that contains two SH2 domains at its N-terminus, a single catalytic domain, and a C-terminal tail that contains tyrosine residues. SHP-1 has been suggested to contribute to the termination of mitogenic signals of growth factors by dephosphorylating critical phosphorylated molecules. SHP-1 has also been shown to be associated with dephosphorylation of cell membrane receptors such as platelet-derived growth factor receptor and the interleukin-3 receptor and/or other downstream substrates such as JAK/STAT3.

Many observations suggest that SHP-1 is associated with proliferation of hematopoietic malignancies and epithelium-derived carcinomas. In breast cancer, Wu et al have reported that introduction of SHP-1 into breast cancer cells with low endogenous expression levels resulted in decreased cell proliferation. Several studies have demonstrated that enhanced SHP-1 expression inhibits STAT3 phosphorylation in breast cancer cells.

In this study, we validated the effect of SHP-1 on the activation of EGFR and HER-2 in breast cancer cells for the first time. We observed an obvious inhibition effect of SHP-1 on EGFR and HER-2 phosphorylation, and found that introduction of SHP-1 to trastuzumab-resistant SKBr-3 cells dramatically promoted trastuzumab sensitivity. Also HER-3 and HER-4, the other members of ErbB family, may also mediate the acquirement of resistance to trastuzumab in breast cancer; we did not observe a combination between SHP-1 and neither HER-3 nor HER-4, which suggested SHP-1 mainly affected ErbB1 and ErbB2 among proteins of this family.

Besides the direct effect of SHP-1 expression level on cell proliferation and apoptosis, SHP-1 seems to be a target of some antitumor drugs. For example, Liu et al reported that SC-2001, a novel obatoclax derivative, inhibited STAT3 phosphorylation through enhancing SHP-1 expression and induces apoptosis in human breast cancer cells. Regorafenib, an inhibitor of multiple protein kinases, dramatically down-regulated p-STAT3 expression and resulted in significant suppression of growth in wild-type SHP-1-expressing colorectal cancer cells through enhancing the phosphatase activity of SHP-1. Other drugs, such as dovitinib, sorafenib derivatives, γ-tocotrienol, and Capillarisin all showed a SHP-1-mediated antitumor activity. Our findings may provide potential association between the antitumor mechanism of these drugs and SHP-1 mediated EGFR inhibition. In addition, it is reasonable to speculate that drugs displaying effective upregulation and/or activation effect on SHP-1 might have potential antitumor effect in trastuzumab-resistant breast cancer cells.

**Conclusion**

To our knowledge, this is the first study to reveal the role of SHP-1 in EGFR/HER-2 positive trastuzumab-resistant breast cancer cells. We discovered that the antitumor activity of trastuzumab was significantly triggered by introduction of SHP-1, accompanied with dephosphorylation of both EGFR and HER-2. SHP-1 may be a useful target for treatment of EGFR/HER-2 positive trastuzumab-resistant breast cancer.

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**Authors’ contributions**

RC Luo conceived of and carried out experiments, analyzed and interpreted data, and drafted the manuscript; YF Wu and R Li conceived of experiments, analyzed, and interpreted data, and wrote the manuscript; JY Zhang,
G Wang, B Liu, XF Huang, and T Zhang analyzed and interpreted data. All authors read, revised, and approved the final manuscript.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**


