



# A case report of low-dose apatinib in the treatment of advanced triple-negative breast cancer

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**Background:** The current study shows that the incidence rate of triple-negative breast cancer accounts for 10–17% of invasive ductal carcinoma of the breast. There is no specific treatment target, the age of onset is relatively small, and the recurrence rate is relatively fast. The prognosis of breast cancer in different subtypes is the most unsatisfactory, with a 5-year survival rate of less than 15%. We report a typical case of metastatic advanced triple-negative breast cancer who responded well to apatinib mesylate after chemotherapy failure and achieved significant progression-free survival, which is relatively rare in triple-negative breast cancer with limited treatment means.

**Case Description:** A 55-year-old female was surgically diagnosed as triple-negative breast cancer on April 17, 2015. After surgery, she had lung metastasis after standard adjuvant chemotherapy and radiotherapy. After receiving the NX regimen (vinorelbine, capecitabine) for 8 cycles, she progressed. Because the patient refused later, she was adjusted to apatinib mesylate, and serious adverse reactions occurred during the treatment process. By adjusting the drug dose, and low-dose apatinib treatment, the lung lesions were close to complete response (CR), reaching a progression-free survival period of 45 months.

**Conclusions:** Low-dose apatinib may be a promising anti-tumor drug for triple-negative breast cancer patients, which needs more samples to verify. This case may provide a reference for the treatment selection of triple-negative metastatic breast cancer in the future.

**Keywords:** Metastatic breast cancer; angiogenesis inhibitors; apatinib mesylate; case report

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## Introduction

Triple negative breast cancer (TNBC) is a special type of breast cancer, which lacks the expression of estrogen receptor (ER), progesterone receptor and human epidermal growth factor receptor 2 (HER2) genes. The incidence rate of TNBC accounts for 10–17% in breast invasive ductal carcinoma. The treatment of TNBC is particularly challenging due to the lack of recognized molecular biological targets. Unfortunately, TNBC is highly proliferative and invasive, the disease progresses rapidly,

the risk of recurrence is high, and the prognosis is poor (1). Chemotherapy is the main treatment of TNBC, and new drugs and strategies are urgently needed. At present, many studies have shown that angiogenesis inhibitors play an important role in advanced TNBC. There are also some small molecule tyrosine kinase (TK) inhibitors with anti-angiogenic effects, such as apatinib, which have also shown some efficacy in clinical studies.

Angiogenesis is one of the signs of cancer, which is crucial for tumor growth, development and metastasis.

Anti-angiogenesis is an important anti-cancer strategy (2). Vascular endothelial growth factor (VEGF) signal plays an important role in angiogenesis by activating VEGF receptor (VEGFR) (2). The VEGFR family involves three molecular subtypes (VEGFR-1, VEGFR-2 and VEGFR-3), which are type II transmembrane proteins characterized by TK activity (3). Among them, VEGFR-2 is mainly related to the pathological excessive formation of blood vessels in a variety of solid tumors (4).

Apatinib is a new generation of small molecule TK inhibitors, which can highly selectively inhibit the phosphorylation of VEGFR-2, mainly through the anti-angiogenesis pathway, and inhibit tumor proliferation and metastasis. Recently, apatinib has shown satisfactory efficacy in many cancers, such as gastric cancer (5), breast cancer (6), and nasopharyngeal cancer (7). At the same time, it has acceptable toxicity. It also shows that the bioavailability in the human body is relatively high, and it is convenient to take orally in clinical practice, with a certain degree of safety, which is easily tolerated by patients (8). Here, we report a typical case of TNBC with lung metastasis. After chemotherapy failure, oral apatinib reached clinical complete response (CR), and reached 45 months of progression-free survival. This case provides a reference for the treatment of triple-negative metastatic breast cancer in the future. We present this case in accordance with

the CARE reporting checklist (available at <https://tbc.amegroups.org/article/view/10.21037/tbcr-23-24/rc>).

## Case presentation

The patient was diagnosed with TNBC at the age of 55 in April 2015. After right breast-conserving surgery, the pathological stage was T1N0M0. After the operation, patients received AC regimen (pirarubicin hydrochloride 100 mg, 50 mg/m<sup>2</sup>, dl and cyclophosphamide 1,000 mg, 60 mg/m<sup>2</sup>, 21 days as a cycle) for 4 cycles, followed by T regimen [paclitaxel 300 mg (175 mg/m<sup>2</sup>, 14 days as a cycle) for 4 cycles]. After that, she received adjuvant radiotherapy, completed 6MV-X Dt 5,000 cGy/25 f, and locally added to Dt 6,000 cGy/30 f. Cellular immunotherapy was performed from October 2015 to January 2016. On March 22, 2017, when right lower lung metastasis was found, NX regimen (vinorelbine capsule 100 mg, 56 mg/m<sup>2</sup>, dl, 8 and capecitabine 3,500 mg, 1,989 mg/m<sup>2</sup>, days 1–14, 21 days as a cycle) was selected as the first-line chemotherapy for 8 cycles, with the best effect reaching disease stabilization (SD). The main adverse reactions were anorexia, liver function damage, and weight loss. On October 19, 2017, there was progress in the disease, which showed that the lung metastasis was enlarged. The patient refused chemotherapy. On October 21, 2017, the patient began to take 500 mg of apatinib mesylate orally for treatment. Grade 1 hand foot syndrome, grade 2 proteinuria, grade 3 diarrhea, and grade 3 hypertension occurred. After 5 months, 500 mg of apatinib mesylate was taken for 5 days and 2 days off, but still had grade 2 diarrhea and grade 1 proteinuria, the blood pressure was well controlled. After 2 months, it was reduced to 500 mg of apatinib mesylate for 4 days and 3 days off. In September 2019, the pulmonary lesions were close to CR (4 mm). According to RECIST 1.1 (9), the treatment evaluation result is a partial response (PR; *Figure 1*). On September 9, 2019, the patient reduced the dosage of the drug to 500 mg of apatinib mesylate for 3 days and 4 days off. During this period, the adverse reactions of the patients were mild (grade 1 hypertension, grade 1 proteinuria, grade 1 diarrhea, grade 1 hand-foot syndrome, grade 1 hypothyroidism), with good tolerance (*Table 1*). By June 2021, a reexamination of positron emission tomography (PET) computed tomography (CT) showed that lymph nodes in the neck, mediastinum, and right hilar lung were metastatic, and left adrenal gland mass was considered to be metastatic. Metastatic tumor of the left frontal lobe. The progression-free survival was 45 months (*Figure 2*).

### Highlight box

#### Key findings

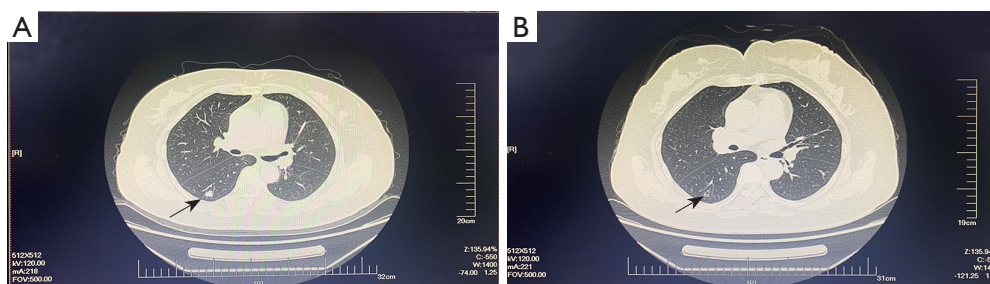
- Low-dose apatinib has achieved good results in a patient with metastatic triple-negative breast cancer who failed chemotherapy, and the treatment is well tolerated.

#### What is known and what is new?

- Recent studies have shown that anti-angiogenic drugs play an important role in advanced triple-negative breast cancer. Some small molecule tyrosine kinase inhibitors with anti-angiogenic effects, such as apatinib, have also shown partial efficacy in clinical studies. However, in clinical practice, the efficacy of using it alone is limited.
- The low dose of apatinib in this case achieved a progression free survival period of 45 months, with good long-term oral tolerance, and is relatively rare in triple negative cancers with limited treatment options.

#### What is the implication, and what should change now?

- The good management of this case takes into account the long-term anti-tumor efficacy, safety, and tolerance, providing a reference for future treatment choices for triple negative metastatic cancer.



**Figure 1** Comparison of lung metastases in breast cancer patients before and after oral apatinib mesylate treatment. (A) Before oral apatinib mesylate treatment for breast cancer patients (October 2017). (B) After oral apatinib mesylate treatment in breast cancer patients (September 2019). Black arrow indicates pulmonary metastasis.

**Table 1** Treatment process of apatinib

Course of treatment	Treatment process			
	2017.10–2018.03	2018.03–2018.05	2018.05–2019.09	2019.09–2021.06
Dose (apatinib)	500 mg q.d.	500 mg for 5 days and 2 days off	500 mg for 4 days and 3 days off	500 mg for 3 days and 4 days off
Efficacy evaluation	PR	PR	PR	PR
Adverse reactions	Grade 1 hand-foot syndrome, grade 2 proteinuria, grade 3 diarrhea and grade 3 hypertension	Grade 2 diarrhea and grade 1 proteinuria, the blood pressure was well controlled	Grade 1 hypertension, grade 1 proteinuria, grade 1 diarrhea, grade 1 hand foot syndrome, grade 1 hypothyroidism	

PR, partial response.



**Figure 2** Time line for treatment of breast cancer patients. NX, vinorelbine, capecitabine; T, albumin paclitaxel; SD, disease stabilization. PR, partial response.

This study was conducted in accordance with the Helsinki Declaration (revised in 2013). This study has been approved by the Ethics Committee of General Hospital of Ningxia Medical University (Ethics Committee registration number 2023-28) and obtained informed consent from the patient.

### Discussion

TNBC accounts for 10–15% of all breast cancer cases. TNBC is a particularly aggressive and heterogeneous breast cancer. Compared with other breast cancer subtypes,

because there is no specific therapeutic target, the prognosis is usually poor. At present, chemotherapy is still the standard treatment method for TNBC. In recent years, some progress has been made in the application of new drugs for specific subtypes of programmed death ligand 1 (PD-L1) + tumor or embryonic BRCA mutation. However, only a small part of these patients have responded to immune checkpoints or Poly (ADP-ribose) polymerase (PARP) inhibitors, and even patients who have responded often have drug resistance and relapse, so the treatment of TNBC remains a clinical challenge, especially, patients who fail in first-line or second-line treatment, they urgently need new drugs and treatment strategies.

Angiogenesis is one of the signs of cancer, which is crucial for tumor growth, development and metastasis. Anti-angiogenesis is an important anti-cancer strategy. VEGF signal plays an important role in angiogenesis by activating VEGFR. The VEGFR family involves three molecular subtypes (VEGFR-1, VEGFR-2 and VEGFR-3), which are type II transmembrane proteins characterized by TK activity. Among them, VEGFR-2 is mainly related to the pathological excessive formation of blood vessels in various

solid tumors.

Apatinib is a drug independently researched and developed in China. It is a new generation of small molecule TK inhibitors. It has potential anti-angiogenesis and anti-tumor functions. It can selectively bind and inhibit VEGFR-2, inhibition of VEGF-stimulated endothelial cell migration and proliferation, and reduce tumor microvessel density, thereby inhibiting tumor proliferation and metastasis. In addition, apatinib can inhibit a variety of TKs related to tumor genesis and progression, thereby inhibiting tumor angiogenesis, in order to achieve the purpose of anti-tumor (10,11).

Apatinib is the first small molecule anti-angiogenesis targeted drug that has been proved to be safe and effective for advanced gastric cancer that has failed standard chemotherapy in the world. In October 2014, it was approved by the State Food and Drug Administration to be marketed for the third-line or above treatment of advanced gastric adenocarcinoma or gastric-esophageal junction adenocarcinoma. At present, although apatinib has not been approved for the treatment of breast cancer, clinical trials for breast cancer have been carried out. In recent years, a series of studies have shown that apatinib has anti-tumor activity in several other solid tumors, including non-small cell lung cancer (12) and breast cancer (6), and has good tolerance in previous studies (13,14). In a multicenter phase II study, the efficacy and safety of apatinib in patients with metastatic TNBC after multi-line therapy were also evaluated (6). The long-term benefit of single-agent apatinib is rarer. The patients reported in this article chose single drug apatinib after failure of first-line chemotherapy, and achieved clinical CR after 2 years of oral apatinib treatment. The overall progression free survival time reached 45 months, which is a very rare clinical success case.

The adverse reactions of long-term oral of apatinib are also the focus of attention. The standard dose of 750 or 500 mg is once a day, and the common adverse reactions such as proteinuria, hypertension and hand-foot syndrome occur frequently. In clinical practice, patients often feel relieved after taking medicine, but have to stop treatment due to uncontrollable hypertension and proteinuria, which affects the overall effect of treatment. A phase IIb clinical study found that the reduction of drug concentration can reduce the occurrence and degree of adverse reactions, and the efficacy is not affected (6). The patient took 500 mg q.d. orally at the time of initial diagnosis, and the main adverse reactions were: elevated blood pressure, hand-foot syndrome, hypothyroidism, proteinuria, diarrhea. Especially

diarrhea and blood pressure rise to grade 3. After 5 months, the condition was evaluated to be stable. Because of adverse reactions, after 2 months, the drug dose was adjusted to 500 mg for 4 days and 3 days off. The clinical CR was reached in the reexamination in September 2019, and the dosage was adjusted to 500 mg for 3 days and 4 days off. The adverse reactions were all grade 1. The overall progression-free survival time reached 45 months, with mild adverse reactions. It can be seen that the reduction of the drug concentration of apatinib can reduce the occurrence and degree of adverse reactions, and the efficacy is not affected. It is consistent with the clinical research results. At the same time, low-dose apatinib not only alleviates the adverse reactions and economic burden of patients, but also achieves better efficacy, and obtains a longer disease progression-free survival period, which provides a new strategy for the treatment of advanced breast cancer.

The patient was adjusted to albumin paclitaxel monotherapy after the disease progressed again, and the gene detection was improved. The results are shown in *Figure 3*. KEYNOTE-355 research results show that for advanced triple-negative breast cancer with high PD-L1 expression, pembrolizumab combined with chemotherapy can significantly improve progression-free survival compared with chemotherapy alone. The detection result of this patient shows that PD-L1 is highly expressed. If the efficacy of albumin paclitaxel was poor, chemotherapy combined with PD-1 treatment could be considered.

Apatinib mesylate monotherapy for multidrug-resistant advanced breast cancer has a significant short-term effect, and has good drug resistance. The adverse reactions are less and can be controlled, thus improving the quality of life of patients. It has become an effective treatment for multidrug resistant breast cancer. This case proves that apatinib mesylate monotherapy has an ideal effect in the treatment of multidrug-resistant advanced breast cancer. Low-dose medication can reduce the adverse reactions of drugs and the economic burden of patients, achieve the same treatment effect, and have better tolerance. It is worthy of clinical promotion and application.

## Conclusions

Here, we report a case of advanced metastatic TNBC. After the first-line chemotherapy progress, she was adjusted to apatinib mesylate, and serious adverse reactions occurred during the treatment. By adjusting the drug dosage and low-dose apatinib treatment, the efficacy is close to clinical

Solid tumor		
Detection content	Detection significance	Detection result
Tumor mutation load (TMB)	Patients with high mutation load have a better prognosis after receiving immunotherapy FDA approved pembrolizumab for the treatment of unresectable or metastatic solid tumors with TMB-H and disease progression after previous treatment	13.68 Muts/Mb, bTMB is high
Immunohistochemical analysis (PD-L1)	FDA approves the use of pembrolizumab for all approved adult indications On November 13, 2020, the FDA approved the combination of pembrolizumab and chemotherapy for the treatment of tumor expression Patients with PD-L1 CPS $\geq 10$ locally recurrent, unresectable or metastatic TNBC. For PD-L1 positive TNBC patients, atezolizumab + albumin bound paclitaxel can also be recommended	PD-L1 TPS 1% PD-L1 CPS 20
MSS detection (MSI)	FDA approves the use of pembrolizumab for solid tumor patients with MSI-H (or dMMR) NCCN guidelines recommend that pamumab be used in breast cancer patients with MSI-H (or dMMR)	MSS

**Figure 3** Patient gene test report. TMB, tumor mutation burden; FDA, Food and Drug Administration; TMB-H, high tumor mutation burden; bTMB, blood-tumor mutation burden; PD-L1, programmed death ligand 1; CPS, combined positive score; TNBC, triple negative breast cancer; TPS, tumor cell proportion score; MSI, microsatellite instability; MSI-H, high-frequency MSI; dMMR, d-mismatch repair; NCCN, National Comprehensive Cancer Network; MSS, microsatellite stability.

CR, achieving a progression-free survival of 45 months.

Through the treatment of this patient, we found that low-dose apatinib may be a promising anti-tumor drug for treating triple-negative cancer patients, requiring more samples for validation. This case can provide a reference for the treatment selection of triple negative metastasis of cancer in the future.

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### Footnote

**Reporting Checklist:** The authors have completed the CARE reporting checklist. Available at <https://tbc.amegroups.org/article/view/10.21037/tbcr-23-24/rc>

**Peer Review File:** Available at <https://tbc.amegroups.org/article/view/10.21037/tbcr-23-24/prf>

**Conflicts of Interest:** All authors have completed the ICMJE

unified disclosure form (available at <https://tbc.amegroups.org/article/view/10.21037/tbcr-23-24/coif>). The authors have no conflicts of interest to declare.

**Ethical Statement:** The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. This study was conducted in accordance with the Helsinki Declaration (revised in 2013). This study has been approved by the Ethics Committee of General Hospital of Ningxia Medical University (Ethics Committee registration number 2023-28) and obtained informed consent from the patient.

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