

A Remarkable and Durable Response to Sintilimab and Anlotinib in the First-Line Treatment of an Anaplastic Thyroid Carcinoma without Targetable Genomic Alterations: A Case Report

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Abstract: Anaplastic thyroid carcinoma (ATC) is a rare and highly aggressive fatal tumor. Most ATC patients using traditional surgery or radio-chemotherapy have poor prognosis and experience recurrence in a very short time. There is no optimal therapy for ATC, and the median survival time is about 5 months. We report a 67-year-old ATC patient, who experienced rapid local recurrence after radical thyroidectomy. The resected tumor tissue was sent for immunohistochemistry analysis and targeted next-generation sequencing. The results indicated high PD-L1 expression, a tumor mutation burden of 0.48 muts/Mb, microsatellite stable, and somatic mutations of *TERT* promoter, *EIF1AX*, *NRAS* and *TP53*. However, none of the mutations indicated corresponding target therapy. An immediate operation was unsuitable because of rapid recurrence after surgery. The patient was also not in a condition to tolerate chemotherapy. Based on the high expression of PD-L1, an optimum strategy was used, combining immunotherapeutic agent, sintilimab, with an anti-angiogenesis drug, anlotinib. The patient obtained remarkable tumor shrinkage and an 18.3-month-sustained remission period. This is an effective case of using immunotherapy and anti-angiogenesis agent in the first-line treatment of ATC. It demonstrates a feasible and novel therapeutic option for future treatment of ATC patients.

Keywords: anaplastic thyroid carcinoma, PD-1 inhibitor, anti-angiogenesis drug, *TERT* promoter mutations, *TP53* mutations, *NRAS* mutation, next-generation sequencing

Introduction

Anaplastic thyroid carcinoma (ATC) is a rare and highly fatal malignancy with characteristic rapid progression, early metastasis, and poor prognosis, accounting for about 1–2% of all thyroid cancers.¹ ATC tumor comprises heterogeneous mixtures of spindle-shaped, epithelioid, and pleomorphic giant cells. Histopathological features of ATC include tumor necrosis, obvious nuclear pleomorphism, and high frequency of atypical mitosis. Major metastases usually occur in the lungs, followed by bones, intrathoracic and neck lymph nodes.^{2,3} ATC is most prevalent in the 60–70-year age group, and its frequency is higher in females than males (2:1.5).¹

Traditional clinical management of ATC involves surgery, radiotherapy and/or chemotherapy.⁴ However, these approaches often do not control the disease because many patients are diagnosed with inoperable or distant metastases.⁴ The

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disease-specific mortality rate is nearly 100%, and the median survival time, after diagnosis, is approximately 5 months; only 20% of patients survive more than one year.¹ Therefore, it is urgent to establish novel diagnostic and standard strategies for ATC.

Multiple driver mutations of ATC have been identified with the development of next-generation sequencing, such as oncogenic mutations of MAPK pathway (*BRAF*, *RAS*), mutations in *TP53* and the *TERT* promoter, and fusions in *RET* and *NTRK*.⁵ Dabrafenib, a selective inhibitor of *BRAF* mutant, and trametinib, a selective MEK inhibitor, were approved by the US Food and Drug Administration (FDA) in 2018, for the treatment of ATC patients with *BRAF* mutation. Other studies have shown that PD-L1 was exclusively expressed on ATC tumor cells, and T cells infiltrated the microenvironment.⁶ As such, several novel monoclonal antibodies, such as nivolumab, pembrolizumab and sintilimab, are under clinical trials and are proposed to be highly promising treatment modalities.⁵ It has been reported that immunotherapy combined with anti-angiogenic drugs have synergistic anti-tumor activity, because the drugs could promote T cell infiltration.⁷ Multi-target tyrosine kinase inhibitor (MTKI) anlotinib, a novel anti-angiogenesis agent, is a popular drug for its efficacy against multiple cancers and low side effects. Previous reports showed that anlotinib could effectively inhibit ATC cell lines.⁸ We reported an ATC patient who had recurrence after radical surgery, having *TP53* and *NRAS* mutation, and high expression of PD-L1, but not *BRAF* mutations. PD-1 inhibitor, sintilimab, and anti-angiogenesis drug, anlotinib, were used, and major remission (about 18 months) was obtained. This report was the first case that investigated and proved the efficacy of immunotherapy combined with anti-angiogenesis agent for ATC, but it still needs a larger investigation.

Case Presentation

A 67-year-old female presented with the chief complaint of rapidly growing neck mass and hoarseness. Contrast-enhanced computed tomography (CT) revealed a neoplastic lesion of 5 cm in the left lobe of thyroid gland and multiple small lymph nodes in the neck but no distant metastases. A surgery was performed on March 20, 2019. The tumor was found to be closely related to muscles and esophagus. The pathology of intraoperative freezing indicated high-grade malignant tumor. She underwent total thyroidectomy and central neck lymph node dissection. The surgical pathology showed multifocal, bilateral

anaplastic thyroid carcinoma (ATC). The tumors broke through the thyroid capsule and involved the surrounding muscles, and the left central lymph nodes were involved (pT4N1M0, IVB). The immunohistochemistry (IHC) was Ki-67 (80% positive), EGFR, P53, AE1/AE3, P16 positive, and TG, P40, EBR, TTF-1, P63, LCA negative. Eleven days later, she underwent a debridement because of esophageal leakage. Unfortunately, two weeks after the second surgery, the patient had a feeling of suffocation. A mass with rapidly progressive enlargement was observed in the operated area with a range of 6.7×6.5×2.8 cm on magnetic resonance imaging (MRI), and was thought to be tumor recurrence. Multiple metastatic lymph nodes were found in the bilateral neck, the biggest one about 1.6×1.0 cm (Figure 1A). The patient was referred to the Cancer Hospital, Chinese Academy of Medical Sciences & Peking Union Medical College. The patient had a poor Eastern Cooperative Oncology Group (ECOG) performance status of 3 with difficulty in breath, unable to lie flat, swollen face and hoarse voice. So, she could not tolerate chemotherapy, radiotherapy or an immediate operation.

Tumor tissue from the thyroidectomy was sent to Genetron Health (Beijing) for molecular testing and immunohistochemical analysis (IHC) of programmed death-ligand 1 (PD-L1). DNA from a paraffin-embedded sample of ATC and the patient's peripheral blood (as control) were analyzed with the Onco PanScan™ (Genetronhealth, Beijing, China), which interrogates 825 cancer-related genes by next-generation sequencing (NGS). The results revealed a tumor mutation burden of 0.48 muts/Mb, microsatellite stable (MSS), and somatic mutations of *TERT* promoter, *EIF1AX*, *NRAS* and *TP53* (Table 1). PD-L1 IHC 22C3 pharmDx assay (Agilent Technologies, Carpinteria, CA, USA) was used to assess the expression of PD-L1 in formalin-fixed thyroid tumor samples, evaluated by both the tumor proportion score (TPS) and the combined positive score (CPS). At least 100 tumor cells must be present in the sample for evaluation. The tumor PD-L1 TPS was 60% and CPS was 70.

The patient was unsuitable for an immediate operation as subsequent treatment given that recurrence occurred shortly after surgery. Also, the patient was not in a condition to tolerate chemo- or radio-therapy. Sintilimab, an immune checkpoint inhibitor, was suggested as an effective treatment option because of the identified association between high PD-L1 expression (TPS ≥ 50%) and improved responses to immune

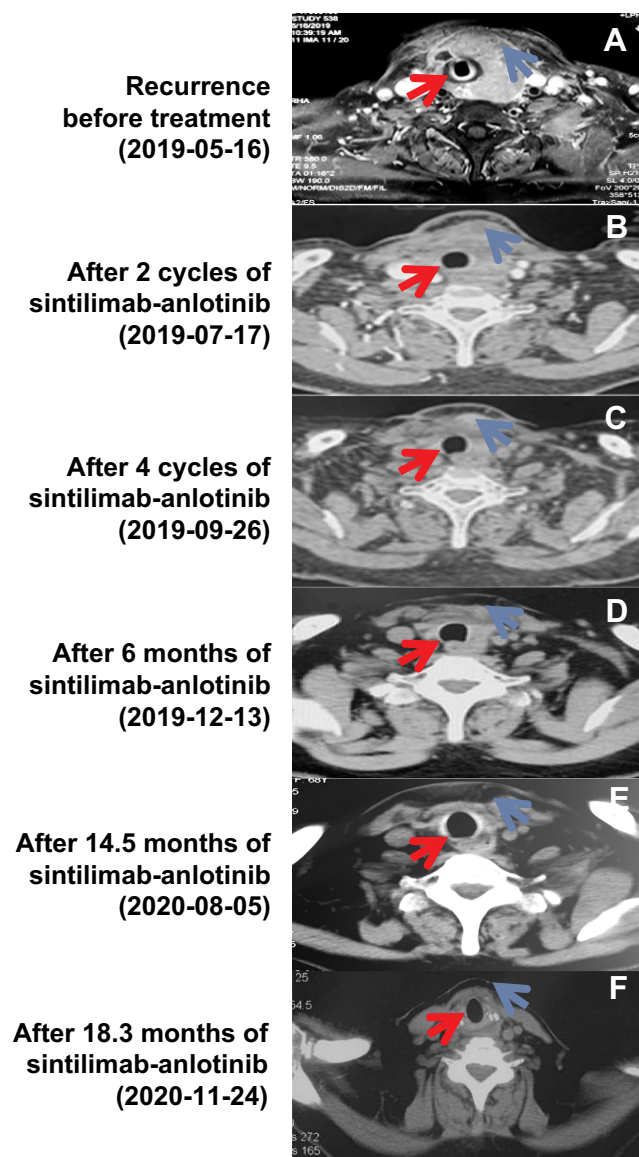


Figure 1 Radiological monitoring of the patient before and after treatment with sintilimab plus anlotinib. (A) Magnetic resonance imaging (MRI) was performed on the recurrence before treatment (Baseline), on May 16, 2019. (B and C). Computerized tomography (CT) on July 17, 2019 (B) and September 26, 2019 (C) showed the tumor shrank after 2 and 4 cycles of sintilimab-anlotinib, respectively. (D) CT on December 13, 2019 identified a little residue of the recurrence mass. (E) CT after 14.5 months of the treatment showed a sustained and prominent reduction of the tumor. (F) The tumor shrinkage was continued over 18.3 months as shown by the CT on November 24, 2020. The red arrow indicates tracheal compression, whereas the blue arrow indicates the tumor.

checkpoint inhibitors. Furthermore, it was proved that anti-angiogenesis drug could improve the immunotherapeutic response; thus, an anti-angiogenesis drug, anlotinib, was added. Starting from June 6, 2019, the immunotherapy agent sintilimab was given 200 mg every 3 weeks, combined with anti-angiogenesis drug, oral anlotinib 12 mg, once daily, 2-week on/1-week off. After 2 cycles (6 weeks) of treatment, CT scanning showed significant

Table 1 NGS* Gene Mutation Profiling in Primary Thyroid Lesion

Gene	Mutation Type	Genovariation	Amino Acid Variation	Allelic Frequency
<i>EIF1AX</i>	Missense mutation	c.25G>C	p. G9R	7.2%
<i>NRAS</i>	Missense mutation	c.182A>G	p. Q61R	11.0%
<i>TP53</i>	Missense mutation	c.584T>C	p. I195T	19.7%
<i>TERT</i>	Upstream promoter mutation	C228T	/	22.58%

Abbreviation: *NGS, next-generation sequencing.

reduction of the mass in the neck, with a maximum cross-sectional area (MCSA) shrinking from 6.7×6.5 cm to 3.7×3.6 cm. Metastatic lymph nodes in the left neck were narrowed, with a short diameter shrinking from 1.0 cm to 0.7 cm in the largest one, and no swelling on the right side (Figure 1B). Partial response was achieved according to RECIST v1.1. In addition, the mass continued to shrink with treatment (Figure 1C). Six months later, CT scan showed little remains with the MCSA of about 1.9×1.4 cm (Figure 1D). A follow-up CT scan at 14.5 months and 18.3 months after treatment demonstrated prominent and sustained reduction of the tumor (Figure 1E and F). The esophagus was damaged during surgery (radiation may cause esophagus injury), and the patient was unwilling to undergo consolidation radiotherapy. Grade 1 rash was the only adverse event observed during treatment. Patient-reported outcome surveys, at approximately 18.3 months of treatment, indicated an excellent quality of life, including continued normal physical exercises such as swimming and jogging.

Discussion

ATC is one of the most fatal cancers. The median survival time after diagnosis is only about 5 months.⁹ Currently, surgery is one of the common treatments for ATC. However, majority of patients with ATC at diagnosed are distant metastasis or locally advanced, which are not amenable to completing resection.^{1,10} Besides, anaplastic carcinoma surgical intervention is affected by more severe complication, including recurrent laryngeal nerve (RLN) palsy,¹¹ hypocalcemia¹² and lymph nodal involvement.¹³ Majority of ATC patients have recurrence soon after surgery. Thus, the benefit of surgical treatment to patient survival is limited. The patient reported here relapsed shortly after surgery, having a poor condition with ECOG score of 3, all of which showed no indication to

commence chemotherapy or radiotherapy. Precise treatment guided by gene diagnosis to determine the efficacy of targeted therapies, such as BRAF inhibitor for *BRAF V600E* mutant positive patients¹⁴ and TRK inhibitor for *NTRK* fusion positive patients¹⁵ and inhibitors for *RET* or *ALK* rearrangement patients,^{16,17} may be an appropriate strategy. However, NGS results of this patient did not show any mutations or rearrangements that had corresponding targeted therapeutic agents; therefore, multi-target anti-angiogenic drugs were considered. Recently, Multi-target tyrosine kinase inhibitor (MTKI) lenvatinib was approved by FDA for the management of advanced differential thyroid cancer after radioactive iodine treatment, and also proposed for ATC patients who do not have other effective therapeutic options. However, the effective rate of lenvatinib for ATC patients was only 17–24%.¹⁸ Anlotinib is a novel MTKI and antiangiogenic drug that can inhibit platelet-derived growth factor receptors, Ret, Aurora-B, epidermal growth factor receptors (EGFR), and fibroblast growth factor receptor.⁸ Anlotinib showed anti-cancer activity on ATC cell lines. It can inhibit tumor cell proliferation by causing ATC cells to have abnormal spindle assembly, G2/M phase arrest, TP53 activation, thereby suppressing cell migration and growth of xenograft thyroid tumors by disturbing the formation of F-action.⁸ According to current clinical reports, anlotinib is safe and effective in the treatment of advanced refractory thyroid cancer.¹⁹ For 58 patients with locally advanced or metastatic medullary thyroid cancer who received anlotinib, 56.9% had a partial response, and the progression-free survival rate at 48 weeks was 85.5%.²⁰ Therefore, anlotinib potentially offers an effective treatment option for ATC patients.

The patient had high PD-L1 expression (TPS 60%, CPS 70). Meta-analysis in 2018 showed that PD-L1 overexpression in patients with non-medullary thyroid cancer is significantly associated with poor survival, disease recurrence and concurrent thyroiditis.²¹ This suggests that PD-L1 may be a prognostic biomarker, and anti-PD-1/PD-L1 immunotherapy may have promise in the PD-L1 positive locally advanced, unresectable, or metastatic thyroid cancer patients. PD-L1 IHC positivity has been implicated as a predictive biomarker in PD-1/PD-L1 checkpoint blockade immunotherapy in a variety of malignancies. It was reported that *BRAF V600E* mutation and PD-L1 expression (3+) were identified in the tumor tissue of a 62-year-old male ATC patient, with local thyroid recurrence, distant metastases after initial surgery and no recovery

after first- and second-line chemotherapies.²² Subsequent treatment with vemurafenib and nivolumab led to substantial regression of tumor nodules. A distinct decrease in metastatic lesions and complete remission was observed 20 months after initiation of nivolumab, demonstrating the effectiveness of immunotherapy for this patient. And an initially unresectable, BRAF-mutated ATC patient with a PD-L1 TPS of 95% was treated with dabrafenib, trametinib, and pembrolizumab. A meaningful response was achieved and the tumor was completely resected. MD Anderson cancer center retrospectively reviewed 12 ATC patients treated with pembrolizumab in combination with kinase inhibitor at the time of disease progression; 42% had a partial response and 33% had stable disease; pembrolizumab was considered an effective salvage treatment.²³ Besides, previous studies showed that melanoma patients with *NRAS* mutations in tumor had better response to immunotherapy and better outcomes than patients whose tumors had other genetic subtypes.²⁴ Here, the ATC patient had mutation of *NRAS*, which may be one of the reasons why the patient was suitable for immunotherapy. Currently, several clinical trials on PD-1/PD-L1 inhibitors for ATC, as single or combination therapy strategies, are ongoing, including NCT02688608, NCT03211117, NCT03246958, and NCT03181100. Sintilimab is a highly selective, humanized, monoclonal antibody that blocks the interaction between PD-1 and its ligands. The agent was approved for the treatment of relapsed or refractory classical Hodgkin lymphoma in China, and it has been tested on multiple advanced solid tumors.²⁵ Therefore, PD-1/PD-L1 immunotherapy using sintilimab was suitable for this patient.

It was recently found that many cancer patients could not benefit from immunotherapy alone, due to vascular abnormalities that facilitate immune evasion in most solid tumors. Judicious use of drugs targeting proangiogenic factors (VEGF, bFGF,^{16,17} PDGF) and their receptors (VEGFR, FGFR, PDGFR) can improve therapeutic response. This is likely due to the normalization of tumor vasculature, therefore increasing the infiltration of immune cells and improving the immunosuppressed microenvironment.⁷ Many studies have evaluated the combination of immune checkpoint inhibitors with the anti-angiogenic drug, bevacizumab, across various tumor types, and most demonstrated marked improvement in the objective response rate, progression-free survival and overall survival.²⁶ Similarly, the antitumor activity of lenvatinib plus anti-PD-1 antibodies was greater than that of either single agent.²⁷ Several clinical trials with

these combined treatments are being tested for thyroid cancer, but no conclusions have been reached. MD Anderson cancer center is studying the efficiency of lenvatinib and pembrolizumab for the treatment of stage IVC metastatic ATC, in an ongoing clinical trial NCT04171622. The National Cancer Institute (NCI) is evaluating the effect of lenvatinib and pembrolizumab combined treatment on differentiated thyroid cancer (DTC), in the NCT02973997 trial. NCI is also testing the effectiveness of the combination of MTKI agent cabozantinib, nivolumab and ipilimumab for advanced DTC, in the NCT03914300 trial. In this study, the tumor shrunk remarkably on treatment with the antiangiogenic drug anlotinib combined with the immunotherapeutic agent sintilimab, and the clinical benefit for the ATC patient has continued over 18.3 months. This case provides evidence of the effectiveness of this combination in ATC patients.

Conclusion

This case reported an ATC patient, who had recurrence within a short period after radical surgery, had high expression of PD-L1, and *TP53* and *NRAS* mutation, but was TMB-low, MSS and did not have genomic alterations with level 1 to 4 of evidence according to the OncoKB knowledge database. PD-1 inhibitor, sintilimab, along with anti-angiogenesis drug, anlotinib, were administered to the patient. Major and sustained remission was obtained. This report was the first case that investigated and proved the efficacy of immunotherapy combined with anti-angiogenesis agent in the first-line treatment of ATC, and the regimen is worthy of further study in ATC.

Ethics Approval

The study was approved by the Institutional Review Board of the Cancer Institute/Hospital, Chinese Academy of Medical Sciences & Peking Union Medical College.

Consent to Publication

Written informed consent for the publication of this case details was obtained from the patient.

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Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

The authors declare no competing interests.

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