Long-Term Favorable Outcome With Nivolumab in a Case of Advanced Non-Small Cell Lung Cancer: A Case Report

Vijay Ketan Reddy, Dhan B. Shrestha, Suman Gaire, Wasey Ali Yadullahi Mir, Mohammed Kassem

1. Department of Internal Medicine, Mount Sinai Hospital, Chicago, USA
2. Department of Emergency Medicine, Palpa Hospital, Palpa, NPL
3. Department of Hematology and Oncology, Mount Sinai Hospital, Chicago, USA

Corresponding author: Dhan B. Shrestha, medhan75@gmail.com

Abstract

Non-small cell lung cancer (NSCLC) constitutes around 85% of lung cancer cases. Advanced non-small cell lung cancer has a poor prognosis. Immunotherapy plays a pivotal role in managing advanced non-small cell lung cancer not positive for driver mutations. Nivolumab is a monoclonal antibody against programmed death-ligand 1 (PD-L1). It is approved as a second-line treatment for patients with advanced non-small cell lung cancer who progress on or after chemotherapy. We present a case of a 71-year-old female with advanced non-small cell lung cancer without any driver mutations diagnosed four years ago. Her disease progressed while on conventional chemotherapy, and she was started on nivolumab three and a half years ago. Her lung nodules resolved, she did not show signs of progression, and her performance status improved while on nivolumab. This case report highlights the current role of nivolumab in the management of NSCLC. Patients whose condition worsens while on conventional chemotherapy can respond very well to modern targeted immunotherapy.

Introduction

Lung cancer is the leading cause of cancer deaths worldwide and in the United States [1,2]. Non-small cell lung cancer (NSCLC) constitutes about 85% of lung cancer cases [3]. The prognosis of lung cancer is poor with a five-year survival of 10%-20% after the diagnosis. Despite the progress in treatment and survival in NSCLC, the two-year survival in distant metastatic NSCLC is still around 20% only [1]. There are targeted therapies for various driver mutations, such as epidermal growth factor receptor (EGFR), anaplastic lymphoma kinase (ALK), and receptor tyrosine kinase (ROS1), in NSCLC, which have improved the prognosis of patients with driver mutations [4,5]. For patients without driver mutations, immunotherapy has played a pivotal role in the management of NSCLC. Several immune checkpoint inhibitors targeting programmed death-1/programmed death-ligand 1 (PD1/PD-L1) have been approved for NSCLC [4,5]. These immune checkpoint inhibitors have increased the survival of patients suffering from NSCLC [6].

Nivolumab is a monoclonal antibody against PD-L1. It was approved for use in advanced NSCLC for patients who have progressed on or after platinum-based chemotherapy [5]. Compared with other PD1 inhibitors, nivolumab has demonstrated similar efficacy with a lesser incidence of severe adverse effects than pembrolizumab or atezolizumab [7].

We present a case of a 71-year-old female suffering from advanced NSCLC without any driver mutation who was managed with nivolumab.

Case Presentation

A 71-year-old female with no significant past medical history had presented four and a half years ago with a weight loss of about 50 pounds over six months prior to presentation. She complained of occasional shortness of breath but denied any cough. Physical examination revealed enlarged right axillary and supraclavicular lymph nodes.

Mammography was performed, revealing multiple enlarged right axillary lymph nodes, but no suspicious lesions were seen in either breast. An ultrasound-guided core biopsy of the right axillary lymph node was performed. The histopathology report showed poorly differentiated metastatic adenocarcinoma. The biopsy was cytokeratin 7 (CK7) and thyroid transcription factor-1 (TTF-1) positive and negative for CK20 and mammaglobin, respectively. Epidermal growth factor receptor (EGFR), ROS1, and anaplastic lymphoma kinase (ALK) were negative, and PD1 and PD-L1 expressions were less than 1%.

A computed tomogram (CT) of the chest with contrast enhancement was done. There were multiple irregular...
nodular opacities in bilateral lungs, with the largest measuring 10 mm, and extensive bilateral mediastinal lymphadenopathy and supraclavicular lymphadenopathy, with the largest measuring 1.8 cm (Figures 1-3). Magnetic resonance imaging (MRI) brain and bone scans were negative for any metastases. However, a positron emission tomography (PET) scan revealed hypermetabolic nodules in the lung and multiple suspicious nodes in the neck, chest, and abdomen, suggesting advanced stage IV lung cancer.

FIGURE 1: CT chest lung window showing right upper lobe nodule on 06/12/2017
She was initially treated with carboplatin, pemetrexed, and paclitaxel for a total of four cycles. She responded well to the treatment with a decrease in the size of pulmonary tumor nodules (from 11.12 to 7.97 mm) noted on the follow-up CT scan (Figures 4-5). She went on to complete eight cycles of maintenance pemetrexed, and on follow-up, a CT scan revealed the right upper lobe lung nodule to have increased in size.
from the previous 1 x 0.7 cm to 1.5 x 0.8 cm (Figure 6). At this time, she was started on nivolumab (3 mg/kg/dose). Six months following commencement of nivolumab, complete resolution of lung lesions was seen on CT chest (Figures 7-9). The lung lesions remain unremarkable since then (Figure 10). Her performance status was stable at ECOG-PS-0 over the four years of treatment, with weight gain from improvement in her appetite. She had resolution of all her respiratory symptoms and B symptoms.

FIGURE 4: CT chest lung window showing right upper lobe lung nodule during chemotherapy on 11/27/2017
FIGURE 5: CT chest soft tissue window showing resolving enlarged right axillary lymph nodes (marked with a black circle) during chemotherapy on 11/27/2017.

FIGURE 6: CT chest lung window showing the increased size of the lung nodule to 14.61 x 8.46 mm on 03/12/2018.
FIGURE 7: CT chest lung window showing resolving lung nodule on 10/26/2018

FIGURE 8: CT chest soft tissue window showing resolved right subclavian lymphadenopathy on 10/26/2018
After being on treatment for almost four years, the patient developed a pruritic, hyperpigmented rash on the upper and lower extremities and trunk. The rash was associated with dermographism and minimal erythema, suggesting grade II-III maculopapular rash, a well-known side effect often seen in patients being
treated with checkpoint inhibitors. At this time, she received a treatment break from nivolumab while she was treated with steroids (1 mg/kg/day of prednisone) and hydroxyzine.

On the resolution of her rash, she was restarted on nivolumab. The patient presented to ER and was subsequently admitted for significant posterior epistaxis and melena along with a drop in her hemoglobin. A CT head scan was performed as part of the workup for severe posterior epistaxis, which showed a 10 mm enhancing lesion in the posterior left frontal lobe, suspicious for metastatic disease. MRI brain confirmed a 0.9 cm enhancing lesion at the posterior left frontal lobe (Figure 11). For the incidental metastatic lesion in her brain, she received stereotactic radiosurgery, given her solitary lesion. The timeline of the events is shown in Figure 12.

FIGURE 11: MRI brain showing incidental brain metastatic lesion on 07/08/2021
Discussion

The treatment of advanced stage NSCLC is palliative in intent to prolong survival and maintain quality of life. With the advent of immunotherapy, immune checkpoint inhibitors have been incorporated into the first-line management of advanced NSCLC. Furthermore, immune checkpoint inhibitors such as nivolumab can be used in patients who progress on or during standard chemotherapy or those who did not receive checkpoint inhibitors as first-line therapy [8].

Our patient at diagnosis had stage IV adenocarcinoma of the lung. She received palliative chemotherapy, followed by pemetrexed maintenance therapy. However, she was started on nivolumab monotherapy due to subsequent disease progression.

Nivolumab is a monoclonal antibody against PD-L1. Tumor cells in NSCLC express various neoantigens that can evade T cells and are sensitive to multiple immune checkpoint inhibitors [9]. PD-L1 is expressed in lung cancer, ovarian cancer, colon cancer, and melanoma [10]. PD-L1-PD1 interaction prevents the activated CD8 T cells from lysing their target cells and promotes CD8 T cell apoptosis [11]. Thus, antibodies that block PD1 or PD-L1, such as nivolumab, potentiate the antitumor immune response.

The Checkmate 227 trial, which compared nivolumab/ipilimumab, nivolumab alone, and chemotherapy in patients suffering from NSCLC, reported higher overall survival, progression-free survival, and objective response rate in nivolumab/ipilimumab as compared with chemotherapy. The benefits were present regardless of PD-L1 level [12]. The combination, therefore, can be used as first-line therapy. However, nivolumab alone is recommended only as second-line therapy. When used as second-line therapy for advanced NSCLC, nivolumab was found to have better efficacy in terms of progression-free survival (hazard ratios [HR]: 0.70, P = 0.05) and overall survival (HR: 0.70, P < 0.00001) as compared to docetaxel in a meta-analysis that pooled six studies [13]. In a real-world pooled analysis of 2585 patients, the median overall survival in previously treated patients with NSCLC on nivolumab was found to be 11.3 months (95% CI: 10.5-12.2) [14].

Our patient showed resolution of lung lesions on CT and no disease progression on nivolumab for three and a half years. However, the MRI brain scan showed a new solitary metastasis in the brain, which was treated with stereotactic radiosurgery, as nivolumab, similar to other monoclonal antibodies, are expected to have no blood-brain barrier penetration [15,16].

Conclusions

Immunotherapy plays a vital role in the management of patients with NSCLC, especially those lacking driver mutations. Nivolumab is an effective second-line immunotherapeutic agent for advanced NSCLC regardless of PD-L1 expression. For the management of NSCLC, nivolumab is used if the patient shows progression on or after chemotherapy. The case report shows the current role of nivolumab in the management of NSCLC. It also highlights the promise of immunotherapy in maintaining the quality of life and prognosis of driver mutation-negative NSCLC.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the
submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

Acknowledgements

We want to acknowledge our patient without whom this report would not have been possible. Additionally, we would like to thank all the treating healthcare personnel involved in patient care.

References

1. Sung H, Ferlay J, Siegel RL, Laversanne M, Soerjomataram I, Jemal A, Bray F: Global cancer statistics 2020: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. CA Cancer J Clin. 2021, 71:209-49. 10.3322/caac.21660
2. Siegel RL, Miller KD, Fuchs HE, Jemal A: Cancer statistics, 2021. CA Cancer J Clin. 2021, 71:7-33. 10.3322/caac.21654
3. Molina JR, Yang P, Cassivi SD, Schild SE, Adjei AA: Non-small cell lung cancer: epidemiology, risk factors, treatment, and survivorship. Mayo Clinic Proc. 2008, 83:584-94. 10.4065/83.5.584
4. Suh J, Blumenthal GM, Jiang X, He K, Keegan P, Pazdur R: FDA approval summary: pembrolizumab for the treatment of patients with metastatic non-small cell lung cancer whose tumors express programmed death-ligand 1. Oncologist. 2016, 21:645-50. 10.1634/theoncologist.2015-0498
5. Kazandjian D, Suzman DL, Blumenthal G, et al.: FDA approval summary: nivolumab for the treatment of metastatic non-small cell lung cancer with progression on or after platinum-based chemotherapy. Oncologist. 2016, 21:654-42. 10.1634/theoncologist.2015-0507
6. Nishijima TF, Shachar SS, Nyrop KA, Muss HB: Safety and tolerability of PD-1/PD-L1 inhibitors compared with chemotherapy in patients with advanced cancer: a meta-analysis. Oncologist. 2017, 22:470-9. 10.1634/theoncologist.2016-0419
7. Passiglia F, Galvano A, Rizzo S, Incorvaia L, Listi A, Bazan V, Russo A: Looking for the best immune-checkpoint inhibitor in pre-treated NSCLC patients: an indirect comparison between nivolumab, pembrolizumab and atezolizumab. Int J Cancer. 2018, 142:1277-84. 10.1002/ijc.31136
8. Ettinger DS, Wood DE, Chair V, et al.: NCCN guidelines panel disclosures NCCN guidelines version 5.2021 non-small cell lung cancer. 2021.
9. McGranahan N, Furness AJ, Rosenthal R, et al.: Clonal neoantigens elicit T cell immunoreactivity and sensitivity to immune checkpoint blockade. Science. 2016, 351:1463-9. 10.1126/science.aaf1490
10. Dong H, Strome SE, Salomao DR, et al.: Tumor-associated B7-H1 promotes T-cell apoptosis: a potential mechanism of immune evasion. Nat Med. 2002, 8:793-800. 10.1038/rmm730
11. Drake CG, Jaffee E, Pardoll DM: Mechanisms of immune evasion by tumors. Adv Immunol. 2006, 90:51-81. 10.1016/S0065-2776(06)90002-9
12. Hellmann MD, Paz-Ares L, Bernabe Caro R, et al.: Nivolumab plus ipilimumab in advanced non-small-cell lung cancer. N Engl J Med. 2019, 381:2020-31. 10.1056/NEJMoa1910251
13. Xu Z, Yi F, Yu D, Xu J, Wei Y, Zhang W: Nivolumab provides improved effectiveness and safety compared with docetaxel as a second-line treatment for advanced non-small cell lung cancer: a systematic review and meta-analysis. Cancer Med. 2019, 8:829-42. 10.1002/cam4.1966
14. Debieuve D, Juergens RA, Asselain B, et al.: Two-year survival with nivolumab in previously treated advanced non-small-cell lung cancer: a real-world pooled analysis of patients from France, Germany, and Canada. Lung Cancer. 2021, 157:40-7. 10.1016/j.lungcan.2021.04.022
15. Tahrizi M, Bornstein GG, Suria H: Biodistribution mechanisms of therapeutic monoclonal antibodies in health and disease. AAPS J. 2010, 12:33-45. 10.1208/s12248-009-9157-5
16. van Bussel MT, Beijnen JH, Brandsma D: Intracranial antitumor responses of nivolumab and ipilimumab: a pharmacodynamic and pharmacokinetic perspective, a scoping systematic review. BMC Cancer. 2019, 19:519. 10.1186/s12885-019-7541-y

2021 Reddy et al. Cureus 13(10): e18526. DOI 10.7759/cureus.18526 10 of 10