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**Relapsed/refractory classical Hodgkin lymphoma effectively treated with low-dose decitabine plus tislelizumab: a case report**

Ding XS *et al*. decitabine plus tislelizumab in Hodgkin lymphoma

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**Abstract**

BACKGROUND

Academic studies have proved that anti-programmed death-1 (PD-1) monoclonal antibodies demonstrated remarkable activity in relapsed/refractory classical Hodgkin lymphoma (cHL). However, most patients ultimately experienced failure or resistance. It is urgent and necessary to develop a novel strategy for relapsed/refractory cHL. The aim of this case report is to evaluate the combination approach of low-dose decitabine plus a PD-1 inhibitor in relapsed/refractory cHL patients with prior PD-1 inhibitor exposure.

CASE SUMMARY

The patient was a 27-year-old man who complained of enlarged right-sided cervical lymph nodes and progressive pain aggravation of the right shoulder over the past 3 mo before admission. Histological analysis of lymph node biopsy was suggestive of cHL. The patient experienced failure of eight lines of therapy, including multiple cycles of chemotherapy, PD-1 blockade, and anti-CD47 antibody therapy. Contrast-enhanced CT showed that the tumors of the chest and abdomen significantly shrunk or disappeared after three cycles of treatment with decitabine plus tislelizumab. The patient had been followed for 11.5 mo until March 2, 2021, and no progressive enlargement of the tumor was observed.

CONCLUSION

The strategy of combining low-dose decitabine with tislelizumab could reverse the resistance to PD-1 inhibitors in patients with heavily pretreated relapsed/refractory cHL. The therapeutic effect of this strategy needs to be further assessed.

**Key Words:** Immunotherapy; hypomethylating agent; classical Hodgkin lymphoma; survival; case report

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**Core Tip:** We report a 27-year-old man who complained of enlarged right-sided cervical lymph nodes and progressive pain aggravation of the right shoulder over the past 3 mo before admission. Histological analysis of lymph node biopsy was suggestive of classical Hodgkin’s lymphoma. The patient experienced failure of eight lines of therapy. computed tomography showed that the tumors of the chest and abdomen significantly shrunk or disappeared after three cycles of treatment with low-dose decitabine plus tislelizumab. The patient had been followed for 11.5 mo until March 2, 2021, and no progressive enlargement of the tumor was observed.

**INTRODUCTION**

Classical Hodgkin’s lymphoma (cHL) is a largely curable malignancy of the lymphatic system. Patients with newly diagnosed cHL are often treated with empirical combination chemotherapy regimens, such as ABVD (doxorubicin, bleomycin, vinblastine, and dacarbazine). Approximately 20% to 30% of patients will experience progression after treatment or fail to respond to induction therapy[1,2]. For these patients, < 20% were exposed to autologous stem-cell transplantation stem-cell transplantation (ASCT) and even fewer to brentuximab vedotin (BV) in China. In recent years, immune checkpoint inhibitors (ICIs) have provided very impressive results with prolonged remission or disease stabilization in many patients. Unfortunately, the majority of these patients will experience resistance or failure, and subsequent therapy is challenging[3,4]. Recently, clinical data found that resistance to ICIs may be reversed by hypomethylating agents, such as decitabine, in heavily pretreated cHL patients[5,6]. The aim of this case report is to evaluate the combination approach of low-dose decitabine plus a programmed death-1 (PD-1) inhibitor in relapsed/refractory cHL patients with prior PD-1 inhibitor exposure.

**CASE PRESENTATION**

***Chief complaints***

The patient was a 27-year-old man who complained of enlarged right-sided cervical lymph nodes and progressive pain aggravation of the right shoulder in 2014.

***History of present illness***

The patient’s right-sided cervical lymph nodes appeared significantly enlarged approximately 3 mo before admittance, and lymph node biopsy was suggestive of cHL. The patient experienced failure of eight lines of therapy. The treatment procedure is provided in Table 1.

***History of past illness***

The patient was diagnosed with subclinical hypothyroidism in September 2018, and levothyroxine (25 µg/d) was prescribed at the Chinese People's Liberation Army General Hospital. The patient’s family history was negative.

***Personal and family history***

Other symptoms were not present and the patient did not drink alcohol, smoke, or have a history of surgery.

***Physical examination***

On admission, the patient’s blood pressure was 120/74 mmHg, pulse rate was 80 beats/min, and temperature was 36.1 °C. His body weight was 77.5 kg, and his height was 172 cm (BMI 26.2 kg/m2, body surface area). His performance status was 2 according to the criteria of the Eastern Cooperative Oncology Group, and his pain was 6 out of 10 according to the numeric rating scale (NRS). No superficial lymphadenopathy was noted. The left buttock was fuller than the right buttock, and no soft tissue tumor was palpated in the left buttock. Left hip tenderness was noted.

***Laboratory examinations***

On admission, routine blood tests revealed significant leukocytosis, moderate anemia, decreased albumin, and slightly elevated blood urea nitrogen. The patient’s blood biochemical tests were indicative of impaired liver function with a slight elevation of alkaline phosphatase and glutamyl transferase levels. The coagulation function test showed a marked elevation in plasma fibrinogen and activated partial thromboplastin time levels. The results are presented in Table 2.

***Imaging examinations***

A computed tomography (CT) scan of the chest revealed that multiple lung nodules significantly decreased or disappeared after treatment with three cycles of combination therapy of decitabine and tislelizumab (Figure 1). Similarly, a CT scan of the abdomen and pelvis showed that abdominal lymph nodes decreased in size (Figure 2).

**FINAL DIAGNOSIS**

The patient was finally diagnosed with cHL by biopsy of the right cervical lymph node. The subtype was nodular sclerosis, Ann Arbor stage IVB, involving the left iliac muscle, left piriformis muscle, sacrum, bilateral ilium, bone marrow, left supraclavicular lymph nodes, mediastinal lymph nodes, right hilar lymph nodes, retroperitoneal lymph nodes, and bilateral lymph nodes near iliac blood vessels. Immunohistochemistry revealed the following: CD15 (+), CD163 (-), CD3 (-), CD30 (+), EBER (-), EBV (-), EMA (-), Ki67 (+), LCA (-), PAX-5 (weakly +), CD20 (+), CD57 (-), ALK (-), and CD68 (-).

**TREATMENT**

The patient was treated with decitabine (10 mg/d, days 1 to 5) plus tislelizumab (200 mg, day 6) every 3 wk. A total of 12 courses of combination therapy were delivered.

**OUTCOME AND FOLLOW-UP**

To date, the patient has completed 12 cycles of decitabine in combination with the PD-1 inhibitor tislelizumab, and the date of the last cycle was November 17, 2020. Abdominal pain was significantly alleviated, and the NRS score decreased to 1 to date. Partial remission was achieved after treatment until March 2, 2021, and progression-free survival was 11.5 mo. The treatment was safe and well tolerated.

**DISCUSSION**

It is difficult to treat relapsed/refractory cHL patients with a heavy pretreatment history due to the lack of standard treatment options. PD-1 inhibitor is considered as a promising option. Programmed death-ligand 1 (PD-L1)/PD-L2 protein is often upregulated in Hodgkin lymphoma cells due to frequent copy-number gains of CD274 (PD-L1) and PDC1LG2 (PD-L2) on chromosome 9p24.1 and Epstein-Barr virus (EBV) infection, which binds the PD-1 receptor on T cells, inducing T-cell exhaustion through the inhibition of T-cell activation and proliferation[4]. In patients with relapsed/refractory cHL who experienced failure with both ASCT and BV, PD-1 blockade has an objective response rate of 65% to 87%[7,8]. However, most patients will ultimately experience progression with a median progression-free survival (PFS) of approximately 12 mo. Thus, an unmet medical need for effective therapies in patients who experienced failure with PD-1 blockade therapy remains.

Recently, a close association between epigenetic aberrations and immune escape has been explored in cHL. Ghoneim *et al*[9] reported that *de novo* DNA methylation programming is causal in reinforcing the development of T-cell exhaustion and establishes a stable cell-intrinsic barrier to PD-1 blockade, causing a decrease in the efficacy of anti-PD-1 antibodies, whereas methylation inhibition could enhance the antitumor activity of PD-1 blockade-mediated T-cell rejuvenation. Falchi *et al*[5] published their experience on a few patients with relapsed/refractory cHL, suggesting that treatment with a PD-1 inhibitor might result in higher complete remission rates as observed in 5/5 patients who were previously exposed to 5-azacitidine. A similar clinical conclusion was obtained by a single-center, two-arm, open-label phase II trial, which included 86 patients with relapsed/refractory cHL after failure of a median of four lines of therapy. In total, 25/86 had been previously exposed to PD-1 inhibitors, and nivolumab was the most commonly used drug, accounting for 72% of cases[6]. The PD-1 inhibitor used in this trial was camrelizumab, and the hypomethylating agent was decitabine, which has been approved to treat myelodysplastic syndromes and acute myeloid leukemia in the United States. The 25 patients with prior PD-1 inhibitor exposure were treated with a combination of 10 mg/d decitabine on days 1-5 and 200 mg camrelizumab on day 8 every 3 wk, and the objective response rate (ORR) and complete remission rate were 52% and 28%, respectively. The ORR was higher in patients with acquired resistance to PD-1 blockade compared to patients with primary resistance (62% *vs* 42%). At one year, the PFS and duration of response rates were 59% and 81%, respectively. Combination therapy could potentially reverse PD-1 resistance due to low-dose decitabine changing the epigenetic status of both tumors and immunocytes, increasing the infiltration of both CD8+ and CD4+ T cells, boosting T-cell function, enhancing tumor immunogenicity, and synergizing with anti-PD-1 antibodies to restore immunosurveillance[10-12]. Thus, hypomethylating agents might have a suppressive effect on tumoral immune escape in relapsed/refractory cHL. Another explanation for the reversal of PD-1 resistance may be attributed to the difference in the structure of different anti-PD-1 monoclonal antibodies. Anti-PD-1 Abs, including pembrolizumab, nivolumab, and other anti-PD-1s, all harbor wild-type Fc regions in the antibody structure. Binding to FcγR on macrophages compromises the antitumor activity of PD-1 monoclonal antibodies with the wild-type Fc region through activation of antibody-dependent macrophage-mediated killing of T effector cells[13,14]. Tislelizumab is a humanized IgG4 anti–PD-1 antibody specifically engineered to minimize binding to FcγR on macrophages. Preclinical data showed that in macrophage- and T cell-enriched conditions, tislelizumab did not induce antibody-dependent cellular phagocytosis; thus, its antitumor activity was not compromised[15].

Here, we report the case of a relapsed/refractory cHL patient who experienced failure with eight lines of therapy and was treated with a combination of low-dose decitabine plus tislelizumab and demonstrate that this combination approach is effective and safe. Further studies are needed to assess the therapeutic effect of this combination therapy in a larger cohort of patients with relapsed/refractory cHL.

**CONCLUSION**

The strategy of combining low-dose decitabine with tislelizumab could reverse the resistance to PD-1 inhibitors in patients with heavily pretreated relapsed/refractory cHL.

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**Footnotes**

**Informed consent statement:** Written informed consent was obtained from the patient for publication of the clinical data and accompanying images.

**Conflict-of-interest statement:** The author reports no conflicts of interest in this work.

**CARE Checklist (2016) statement:** The authors have read the CARE Checklist (2016) and the manuscript was prepared and revised according to the CARE Checklist (2016).

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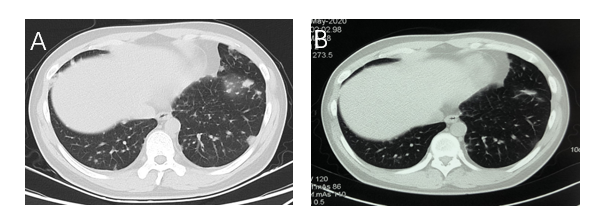
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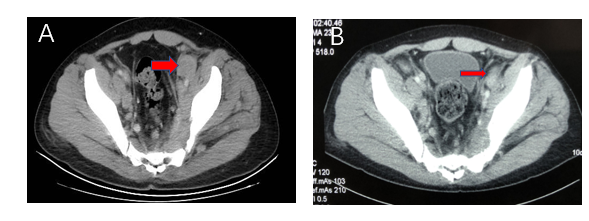
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**Figure Legends**



**Figure 1 Computed tomography scan of the chest before and after combination therapy.** a: Imaging features before treatment; B: Imaging changes after three cycles of treatment.



**Figure 2 Computed tomography scan of the abdomen and pelvis before and after combination therapy.** a: Imaging features before treatment; B: Imaging changes after three cycles of treatment.

**Table 1 The entire treatment process after the diagnosis**

|  |  |  |  |
| --- | --- | --- | --- |
| **Time of therapy** | **Treatment** | **Best efficacy** | **PFS (mo)** |
| November 2014 | ABVD × 6 cycles | CR | 16 |
| March 2016 | AVD × 1 cycle | unknown | - |
| September 2016 | AVD × 1 cycle | unknown | - |
| October 2016 | GVD × 4 cycles | PD | 4.8 |
| October 2017 | GVD × 6 cycles | SD | 8.8 |
| July 2018 | ESHAP × 1 cycle | PD | 0.9 |
| October 2018 | AK105 × 9 cycles | PR | 3.5 |
| July 2019 | IBI188 × 16 wk | SD | 3.6 |
| November 2019 | DICE × 2 cycles | PD | 1.3 |
| February 2020 | F0002-ADC × 1 cycle | PD | 0.6 |
| March 2020 | decitabine plus tislelizumab | PR | 11.5 |

PFS: progression-free survival; CR: complete remission; PR: partial remission; PD: progression disease; SD: stable disease; ABVD: doxorubicin, bleomycin, vinblastine, and dacarbazine; AVD: doxorubicin, vinblastine, and dacarbazine; GVD: gemcitabine, vinorelbine, liposomal doxorubicin; ESHAP: etoposide, methylprednisolone, high-dose cytarabine, and cisplatin; DICE: dexamethasone, ifosfamide, cisplatin, and etoposide.

**Table 2 Laboratory blood test results at hospital admission**

|  |  |  |
| --- | --- | --- |
|  | **Result** | **Normal range** |
| Hematology | | |
| White blood cell count | 20.81 × 109/L | 4.0-10.0 |
| Red blood cell count | 3.17 × 1012/L | 3.5-5.5 |
| Hemoglobin | 87 g/L | 120.0-160.0 |
| Hematocrit | 28.8% | 37.0-49.0 |
| MCV | 91 fl | 82.0-92.0 |
| MCH | 27.3 pg | 27.0-31.0 |
| MCHC | 300 g/L | 320.0-360.0 |
| Platelets | 384 × 109/L | 100.0-350.0 |
| Lymphocytes | 0.96 × 109/L | 1.0-5.0 |
| Monocytes | 0.24 × 109/L | 0.2-0.8 |
| Neutrophils | 19.59 × 109/L | 2.0-8.0 |
| Eosinophils | 0 × 109/L | 0.1-0.5 |
| Basophils | 0.02 × 109/L | 0.0-0.1 |
| ESR |  |  |
| Coagulation | | |
| APTT | 59.3 s | 24.0-39.0 |
| Thrombin time | 16.6 s | 14.0-21.0 |
| Prothrombin time | 12.6 s | 11.0-14.0 |
| INR | 1.11 | 0.8-1.5 |
| Prothrombin activity | 81.9% | 70.0-130.0 |
| Fibrinogen | 942.1 mg/dl | 200.0-400.0 |
| Biochemistry |  |  |
| C-reactive protein | 106.3 mg/L | < 8.0 |
| Procalcitonin | 0.47 ng/mL | < 0.5 |
| Glucose | 7.24 mmol/L | 3.6-6.1 |
| Creatinine | 57 µmol/L | 50.0-130.0 |
| MDRD GFR | > 60 ml/min | > 60 |
| Uric acid | 318 µmol/L | 90.0-420.0 |
| Lactate dehydrogenase | 260 IU/L | 110.0-240.0 |
| ALB | 40.1 g/L | 35.0-55.0 |
| TP | 69.4 g/L | 60.0-80.0 |
| Alkaline phosphatase | 178 IU/L | 40.0-160.0 |
| Serum iron | 9.9 µmol/L | 10.6-28.3 |
| Serum ferritin | 7585 µg/L | 30.0-400.0 |
| UIBC | 20 µmol/L | 19.7-66.2 |
| TIBC | 29.9 µmol/L | 40.8-76.6 |
| Total cholesterol | 4.66 mmol/L | 2.84-5.68 |
| Triglycerides | 0.97 mmol/L | 0.56-1.7 |
| Hormones | | |
| TSH | 3.47 mIU/L | 0.27-4.2 |
| FT3 | 3.59 pmol/L | 3.1-6.8 |
| FT4 | 17.34 pmol/L | 12.0-22.0 |
| Electrolytes | | |
| Sodium | 137 mmol/L | 135.0-145.0 |
| Potassium | 4.47 mmol/L | 3.5-5.3 |
| Calcium-total | 2.35 mmol/L | 2.12-2.75 |
| Phosphates inorganic | 0.83 mmol/L | 0.69-1.6 |
| Liver enzymes | | |
| ALT | 25 IU/L | 0.0-40.0 |
| AST | 23 IU/L | 0.0-45.0 |
| GGT | 151 IU/L | 10.0-50.0 |
| DBIL | 3.0 µmol/L | 0.0-6.0 |
| Total bilirubin | 9.4 µmol/L | 1.7-20.0 |
| Virology tests | | |
| Anti-HCV | Negative |  |
| HbsAg | Negative |  |
| Anti-HIV | Negative |  |

MCV: Mean corpuscular volume; MCH: Mean corpuscular hemoglobin; MCHC: Mean corpuscular hemoglobin concentration; ESR: Erythrocyte sedimentation rate; APTT: Activated partial thromboplastin time; INR: International normalized ratio; MDRD GFR: Modification of diet in renal disease glomerular filtration rate; TP: total protein; UIBC: Unsaturated iron binding capacity; TIBC: Total iron binding capacity; TSH: Thyroid stimulating hormone; ALT: Alanine aminotransferase; AST: Aspartate aminotransferase; GGT: Gamma-glutamyl transferase; DBIL: direct bilirubin; HbsAg: Surface antigen of hepatitis B virus; HCV: hepatitis B virus; HIV: human immunodeficiency virus.



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