Immunoexpression of PD-L1 and PD-1 and Its Clinicopathological Correlation in Urothelial Carcinomas

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J Lab Physicians 2022;14:197–201.

Abstract

Introduction Urothelial carcinomas are the most common types of bladder tumors that have recently shown a changing trend in treatment protocols with the introduction and approval of immune checkpoint inhibitors. The most important immune checkpoint lies with the PD-1–PD-L1 axis. Although multiple drugs have been approved, there is uncertainty about patient selection criteria and diagnostic assays. Recent studies related to the laboratory-developed tests have opened up the horizon of PD-1 and PD-L1 immunohistochemistry even at resource-constrained laboratories. We propose to study these immunohistochemistry markers in our laboratory using newer clones.

Materials and Methods We selected 116 consecutive cases of transurethral bladder tumor resection from our laboratory archive and applied PD-1 and PD-L1 immunohistochemistry. The study was approved by the institution’s ethics committee.

Results We found high expression of PD-1 and PD-L1 in urothelial carcinoma even with different cut-offs of PD-L1. Muscle invasion, lamina invasion, and grade of carcinoma had a statistically significant effect on the expression; however, age and sex did not affect the expression.

Conclusion Based on our current study, we can conclude that the clones used in our study show high expression in urothelial carcinoma and can aid in patient selection and treatment protocol, irrespective of age and sex.

Keywords
- immune checkpoint inhibitors
- PD-1
- PD-L1
- urothelial carcinoma

Introduction

Programmed cell death protein 1 (PD-1 or CD 279) and its ligand programmed cell death ligand 1 (PD-L1 or CD 274 or B7-H1) are cell-surface transmembrane molecules that play an important role in downregulating the immune system. Cancer cells express PD-L1 to avoid immunorecognition and destruction by imitating the signaling process of healthy cells. Currently, multiple immune checkpoint inhibitors including anti-PD-1 and anti-PD-L1 have been approved in
metastatic and advanced urothelial carcinoma expanding the scope of treatment of urothelial carcinoma. Identifying patients who may or may not respond to PD1/PDL1 inhibitors is important as the majority of patients in different clinical trials did not have an overall response. Immunohistochemistry is the most commonly used assay in different clinical trials for patient selection. Although five immune checkpoint inhibitors have been approved in urothelial carcinoma, only two companion diagnostic immunohistochemistry assay has been approved by the US FDA. The Canadian Association of Pathologists-Association Canadienne Des Pathologistes (CAP-ACP) has recently published a guideline for Fit-For-Purpose PD-L1 biomarker testing for patient selection. This has opened up the space for validating new clones for PD-1 and PD-L1 immunohistochemistry assays. We tried to examine the expression of PD-L1 and PD-1 in our laboratory using new clones that have not yet been validated for patient selection for treatment in urothelial carcinoma.

Methodology

This study was a retrospective study and 116 consecutive cases of urothelial carcinoma received as TURBT (transurethral resection of urinary bladder tumors) chips were taken from the archived tissue blocks after the removal of duplicate cases. The cases included in the study did not receive any chemotherapy before surgery. Histopathological and immunohistochemical analyses were performed by two histopathology consultants independently. The examiners were blinded to histopathological diagnosis while interpreting immunohistochemistry slides. Clinical characteristics such as age, gender, and, smoking history were obtained from hospital records.

PD-L1 and PD-1 immunostaining were performed manually using prediluted antibodies for PD-1 (clone NAT105, isotype mouse IgG, Biocare Medical, USA) and PD-L1 (clone CAL10, isotype Rabbit IgG, Biocare Medical, USA). Four-micron-thick paraffin sections were brought to water. Heat-induced epitope retrieval method under steam pressure was used for antigen retrieval with 1% EDTA (pH 8.0) as the retrieval buffer. This was followed by peroxidase blocking using Biocare’s Peroxidase blocking system for 5 minutes and washing the sections twice. The slides were flooded with a primary antibody and incubated in dark at room temperature for 1 hour. After washing, slides were treated with horseradish peroxidase (HRP) for 10 minutes, followed by detection using the DAB detection system as chromogen and counterstained with hematoxylin. Known positive and negative controls were used with each batch of IHC staining.

Immunohistochemical staining was then assessed for PD-L1 and PD-1. Membranous positivity in tumor cells and any cytoplasmic/membranous staining in mononuclear cells was considered as positive for PD-L1 (Fig. 1A, B). PD-L1 immunohistochemistry was scored as the percentage of positive cells. The cases were analyzed using a cut-off of both 1% and 5% for both lymphocytes and tumor cells separately. PD-1 was evaluated in mononuclear cells and any cytoplasmic/membranous positivity was taken as positive (Fig. 1C). The staining percentage was calculated on the slides examined for the immunohistochemistry by calculating the percentage of lymphocytes or tumor cells showing staining.

The complete data were divided based on clinical and pathological features including age (0–40, 40–60, > 60 yrs), smoking history (present, absent), gender (male, female), grade (high, low), lamina invasion (invasive, non-invasive), and muscle invasion (present, absent) (Table 1).

R version 3.5.3 was used for statistical analysis. Fisher’s exact test was used for comparison between different groups.

Results and Interpretation

Our study consisted of 116 patients. The clinicopathological characteristics of the study population are shown in Table 1. Deep muscle biopsy could not be evaluated in 10 patients due to either the absence or marked crushing of deep muscle tissue. Smoking history was known in 90 cases only, which were used for analysis.

PD-1 expression was noted in 32.7% (38 of 116) of cases. The expression was seen in 43% (33 of 77) of high-grade cases and 49% (21 of 43) of muscle-invasive cases (Table 1).

PD-L1 expression in tumor cells was seen in 62% (72 of 116) and 50.8% (59 of 116) of cases, respectively, when the cut-offs of 1% and 5% were taken as positivity, respectively. PD-L1 expression in tumor-infiltrating lymphocytes was seen in 79.3% (92 of 116) and 68.1% (79 of 116) of cases, respectively, when the cut-offs of 1% and 5% were taken for positivity (Table 1).

Immunoreactivity of PD-1 and PD-L1 showed no statistical difference in different age groups, gender, or smokers. PD-1 showed a higher expression in high-grade tumors.
(p = 0.001) and tumor involving deep muscle (p = 0.006), while it showed a non-significant difference in the expression in lamina invasive and non invasive tumors (p = 0.07). PD-L1 immunoexpression in tumor cells shows a higher expression in tumors with higher grade, lamina invasion, and deep muscle invasion; however, PD-L1 immunoexpression in TILs showed higher expression in higher grade and lamina invasive tumor but no difference in tumors involving deep muscle (for p-values, refer to Table 2).

### Discussion and Conclusion

Although five immunotherapeutic agents (PD-1/PD-L1 inhibitors) have been approved by the US-FDA for use in bladder tumors, the appropriate assay for patient selection has remained controversial to date. Immunohistochemistry for PD-L1 and PD-1 is the most commonly used assay; however, the guidelines for positivity and clones to be used remain vague. A recent guideline by CAP-ACP allows the use of laboratory-developed tests (LDTs) if properly validated clinically and in the laboratory to be used for the specific purpose for which it has been validated. In our study, we have tried to evaluate the expression of PD-1 and PD-L1 in urothelial carcinoma using clones NAT105 and CAL10, respectively, and compare it with previous studies. We also plan to compare the immunoexpression of these antibody clones with already established assays in further studies. Although these clones are evaluated in other tumors such as lung, breast, and renal cell carcinomas, the literature regarding its evaluation in bladder carcinoma is limited.  

PD-1 immunohistochemistry in the present study showed positivity in 32.7% of cases with a high expression in high-grade and muscle-invasive carcinomas. A high expression in high-grade and muscle-invasive carcinomas has also been shown in the previous study by Kawahara et al. Although previous studies have shown a slightly higher expression, this may be due to different clones used, a higher number of invasive and high-grade carcinomas, tumor heterogeneity, and the different cut-offs for positivity. We also experienced marked difficulty in establishing the staining protocol and interpretation of PD-1 immunohistochemistry. We observed that the staining intensity is mild to moderate with

### Table 1 Clinicopathological variables used in this study along with the positive number of cases in different categories

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Number of cases</th>
<th>PD-1 positive</th>
<th>PD-L1 positive TILs (1% cut-off)</th>
<th>PD-L1 positive TILs (tm 1% cut-off)</th>
<th>PD-L1 positive TILs (5% cut-off)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0–40</td>
<td>13</td>
<td>4</td>
<td>10</td>
<td>8</td>
<td>9</td>
</tr>
<tr>
<td>&gt; 40–60</td>
<td>53</td>
<td>16</td>
<td>39</td>
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<td>37</td>
</tr>
<tr>
<td>&gt; 60</td>
<td>50</td>
<td>18</td>
<td>43</td>
<td>32</td>
<td>33</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Male</td>
<td>106</td>
<td>35</td>
<td>83</td>
<td>64</td>
<td>71</td>
</tr>
<tr>
<td>Female</td>
<td>10</td>
<td>3</td>
<td>9</td>
<td>8</td>
<td>8</td>
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<tr>
<td>Muscle invasion</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>43</td>
<td>21</td>
<td>37</td>
<td>33</td>
<td>30</td>
</tr>
<tr>
<td>Absent</td>
<td>63</td>
<td>14</td>
<td>46</td>
<td>32</td>
<td>41</td>
</tr>
<tr>
<td>Lamina invasion</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>95</td>
<td>35</td>
<td>81</td>
<td>64</td>
<td>70</td>
</tr>
<tr>
<td>Absent</td>
<td>21</td>
<td>3</td>
<td>11</td>
<td>8</td>
<td>9</td>
</tr>
<tr>
<td>Grade</td>
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<tr>
<td>High</td>
<td>77</td>
<td>33</td>
<td>68</td>
<td>59</td>
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<tr>
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<td>39</td>
<td>5</td>
<td>24</td>
<td>13</td>
<td>20</td>
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<tr>
<td>Smoking</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>44</td>
<td>14</td>
<td>37</td>
<td>23</td>
<td>30</td>
</tr>
<tr>
<td>Absent</td>
<td>46</td>
<td>18</td>
<td>36</td>
<td>30</td>
<td>31</td>
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</tbody>
</table>

Abbreviations: TIL, tumor infiltrating lymphocytes; tm, tumor.

### Table 2 p-Values of Fisher’s Exact Test for clinicopathological variables

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>PD-1</th>
<th>PD-L1 TILs (1% cut-off)</th>
<th>PD-L1 tm (1% cut-off)</th>
<th>PD-L1 TILs (5% cut-off)</th>
<th>PD-L1 tm (5% cut-off)</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>0.885</td>
<td>0.292</td>
<td>0.962</td>
<td>0.959</td>
<td>0.627</td>
<td>116</td>
</tr>
<tr>
<td>Muscle Invasion</td>
<td>0.006</td>
<td>0.150</td>
<td>0.008</td>
<td>0.677</td>
<td>0.005</td>
<td>106</td>
</tr>
<tr>
<td>Gender</td>
<td>1</td>
<td>0.685</td>
<td>0.314</td>
<td>0.499</td>
<td>0.743</td>
<td>116</td>
</tr>
<tr>
<td>Grade</td>
<td>0.001</td>
<td>0.001</td>
<td>&lt; 0.001</td>
<td>0.011</td>
<td>&lt; 0.001</td>
<td>116</td>
</tr>
<tr>
<td>Lamina invasion</td>
<td>0.070</td>
<td>0.002</td>
<td>0.002</td>
<td>0.009</td>
<td>0.030</td>
<td>116</td>
</tr>
<tr>
<td>Smoking</td>
<td>0.514</td>
<td>0.593</td>
<td>0.284</td>
<td>1</td>
<td>1</td>
<td>90</td>
</tr>
</tbody>
</table>
marked tumor heterogeneity in the expression, which has been described in previous studies. 

PD-L1 expression in tumor cells was seen in 62% of cases when a cut-off of 1% was taken and 51% of cases when a cut-off of 5% was taken for positivity. Similarly, PD-L1 expression in tumor-infiltrating lymphocytes was seen in 79% of cases when a cut-off of 1% was taken as positivity and 68% of cases when a cut-off of 5% was taken for positivity. The clone used in this study showed a relatively higher expression as compared with that reported in previous studies; however, we cannot evaluate the optimal cut-off values and propose to do the same in further studies using established assays and follow-ups. PD-L1 expression at a cut-off of both 1% and 5% in TILs was higher in high-grade and lamina invasive tumors \( (p < 0.05) \) and there was no statistical difference in the expression in muscle-invasive or noninvasive tumors.

PD-L1 expression in tumor cells showed a high expression in high-grade, lamina invasive, and also muscle-invasive tumors \( (p < 0.05) \). The correlation of PD-L1 expression with grade and muscle invasion has also been shown previously by Kawahara et al and other authors. Differences in age, gender, or smoking history did not show any significant difference in the expression of either PD-1 or PD-L1. A previous study by Holland et al correlating clinicopathological features with PD-1 and PD-L1 expression has also shown no impact of age and sex on the expression.

Higher PD-L1 expression is related to a better overall response to treatment and survival as in multiple studies done previously. Recently, the US FDA has approved the use of Atezolizumab and Pembrolizumab for first-line therapy-ineeligible patients only in PD-L1-positive tumors. This has opened up the scope of compulsory PD-L1 testing in patients with urothelial carcinoma ineligible for platinum-based therapies; however, the availability of companion diagnostic approved is scarce and laboratory dependent. There is a marked demand for the development and validation of laboratory-dependent tests so that these may be used before treatment. This will promote the use of immunotherapeutic agents on a larger scale even in resource-poor settings.

Our current study is limited by the absence of correlation with a validated diagnostic assay. Moreover, we have not ascertained a fixed cut-off criterion for PD-L1 but this is important for further studies so that the data can be correlated. We also tried to follow up the patients; however, only 8 out of the total patients can be followed up and because these patients belonged to different categories, a valid statistical study cannot be done. This is the first study of immunoeexpression of both PD-1 and PD-L1 in the literature in bladder tumors in the Indian subcontinent to the best of our knowledge and can be used as a baseline study.

Based on the present study we conclude that PD-L1 clone CAL10 showed excellent expression and appears to be comparable to other clones; however, this needs to be further confirmed using validated assays. The high prevalence of PD-1 and PD-L1 positivity irrespective of age and sex also points to the possible use of immunotherapeutic agents in all sex and age groups.

Authors’ Contributions
Utupal Kumar contributed to the concept, data collection, literature search, immunohistochemistry, primary manuscript, and editing of the final manuscript. Michael Leonard Anthony contributed to the concept, data collection, and editing of the final manuscript. Rishabh Sahai contributed to the concept, data collection, literature search, immunohistochemistry, and editing of the final manuscript. Ankur Mittal contributed to the concept, data collection, and editing of the final manuscript. Prashant Durgapal contributed to the concept, primary reporting, primary manuscript, and editing of the final manuscript. Sanjeev Kishore contributed to the concept, primary reporting, scholarly inputs, and editing of the final manuscript.

Institutional Ethics Committee Approval
The study was approved by the Institution’s Ethics Committee (approval no: AIIMS/IEC/20/185).

Conflict of Interest
None declared.

Acknowledgments
I thank Mr. Chandan and Mr. Tarakeshwar for extending their technical support.

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