The DPY30-H3K4me3 Axis-Mediated PD-L1 Expression in Melanoma

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Background: DPY30 is a common subunit of the human SET1/MLL complex and is an essential protein required for the activity of SET1/MLL methyltransferase. DPY30 regulates the histone H3K4 modification, and dysfunction of DPY30 might contribute to the regulation of cancer immune evasion. However, the functions and regulation of DPY30 in the expression of programmed cell death ligand 1 (PD-L1) is still not completely explored.

Methods: Various online databases were used for data processing and visualization, including UALCAN, Oncomine, cBioPortal, SangerBox, TISIDB, TIMER, and GEPIA databases. The expression of DPY30 and PD-L1 in melanoma tissues were evaluated by IHC. Chromatin Immunoprecipitation (ChIP), RT-PCR and flow cytometry were used to elucidate the underlying molecular mechanism of PD-L1 expression regulation and its function.

Results: The mRNA level of DPY30 in melanoma was higher than in normal tissues. The expression of DPY30 was positively associated with TMB, neoantigens and PD-L1 expression. Furthermore, DPY30 expression showed significant positive correlations with immune suppressor cells and ICP genes involved in T-cell exhaustion. IHC showed that the positive rates of DPY30 and PD-L1 in melanoma tissues were 62% and 58%, respectively. Correlation analysis revealed that DPY30 over-expression was positively associated with PD-L1 expression. Silencing of DPY30 by specific siRNA significantly inhibited PD-L1 expression. ChIP analysis revealed that H3K4me3 levels were enriched in the proximal PD-L1 promoter region in tumor cells. Inhibition of DPY30 still suppressed the PD-L1 level in IFN-γ treated MMAC-SF cells. Furthermore, the apoptosis of PD1+ T-cells in co-culture with MMAC-SF cells by knockdown of DPY30 were markedly reduced.

Conclusion: This study shows the roles of DPY30 in regulating the cancer immune evasion in melanoma. Targeting the DPY30-H3K4me3 axis might be an alternative approach to enhance the efficacy of checkpoint immunotherapy.

Keywords: DPY-30, H3K4me3, PD-L1, melanoma, immunotherapy

Introduction

Melanoma stands out as one highly immunogenic cancer that approaches durable response to checkpoint immunotherapy, leading to the first approval of immune checkpoint inhibitors in the treatment of melanoma.1,2 Immune checkpoint inhibitors targeting the cytotoxic T lymphocyte antigen 4 (CTLA-4), programmed cell death protein 1 (PD-1) and programmed cell death ligand 1 (PD-L1) led to a considerable improvement of survival in patients with metastatic melanoma.3 Unfortunately, about 60% of patients with melanoma showed primary resistance to PD-1 blockade, and 20–30% of initial responders would develop secondary drug resistance.4 It has been suggested that the expression of PD-L1 in tumor cells is considered a predictive biomarker for response to immunotherapy,5 but the mechanism underlying melanoma resistance to PD-1/PD-L1 inhibitors is still elusive.

PD-L1 is expressed on the surface of several cell types, including T-cells, B-cells, antigen-presenting cells (APCs), monocytes, and epithelial cells, and is upregulated in response to IFN-γ.6 IFN-γ is a key cytokine produced by activated...
T cells, as well as natural killer (NK) and NK T cells, in the tumor microenvironment. Notably, tumor cells can adaptively increase the expression of PD-L1 under the control of IFN-γ. IFN-γ binds with receptor and subsequently activates JAK/STAT signaling pathway, which leads to the downstream expression and activation of IRF-1, further inducing PD-L1 expression on tumor cells. Furthermore, high PD-L1 expression detected in cancer cells and associated with poor prognosis. The binding of PD-L1 to PD-1 in T cells induced T cell apoptosis and reducing immunocompetence, which thus helps cancer cells escape immune surveillance and killing. Moreover, it has been reported that MYC and Sox2 overexpression, and the loss of PTEN regulate constitutive PD-L1 expression in tumor cells. PD-L1 has become a key protein in immunotherapy, and its regulatory mechanisms are being intensively studied.

Histone H3K4 methylation is one of the most prominent epigenetic modifications, and plays critical roles in regulating target gene transcription. Especially histone H3K4 trimethylation (H3K4me3) selectively localizes at gene promoters and transcriptional start sites and directly regulates gene transcription. As the most notable histone H3K4 methylation enzymes in mammals, the SET1/MLL complexes comprise Set1a or −1b or Mll1, −2, −3, or −4 as the catalytic subunit, and Wdr5, Rbbp5, Ash2l, and Dpy30 as integral core subunits necessary for the full methylation activity. DPY30 directly facilitates genome-wide H3K4 methylation and involves in regulating fundamental cellular processes. It has been reported that DPY30 significantly promotes cell proliferation, migration, invasion, and epithelial–mesenchymal transition of epithelial ovarian cancer. Similarly, DPY30 plays a critical role in the proliferation and appropriate differentiation of hematopoietic progenitor cells. Moreover, DPY30 regulates cell senescence through ID protein expression. However, the roles and mechanisms of DPY30 in the development of melanoma are still unclearly known.

In this study, we analyzed the expression of DPY30 in melanoma and investigated the correlation between DPY30 and progression, and cancer immune evasion in melanoma on the basis of various databases. Finally, we evaluated DPY30 and PD-L1 levels in melanoma tissues by IHC analysis, and tested the hypothesis that PD-L1 expression is regulated by an epigenetic mechanism in melanoma.

**Materials and Methods**

**Cell Culture and Reagents**

Human melanoma MMAC-SF (RRID: CVCL_1420) and WM-115 (RRID: CVCL_0040) cells were propagated in DMEM medium and MEM medium at humid atmosphere (5% CO₂, 37°C), respectively. The culture medium was supplemented with 10% FBS, penicillin (100 u/mL) and streptomycin (100 mg/mL). FITC Annexin V Apoptosis Detection Kit I (cat# 556547) was supplied by BD Biosciences (San Jose, CA, USA). Chromatin Immunoprecipitation Kit (17–10086) was obtained from Merck (Millipore, CA, USA). H3K4Me3 (9751S) was purchased from Cell Signaling Technology (Danvers, MA, USA). IFN-γ (300–02) was supplied by Peprotech (Jiangsu China). PE anti-mouse PD-L1 (124307) and PE anti-mouse PD-1 (135205) were purchased from Biolegend (San Diego, CA, USA). Anti PD-L1 antibody (ab205921) was obtained from Abcam (Cambridge, MA, USA). Anti DPY30 antibody (MA5-32900) was acquired Invitrogen (Carlsbad, CA, USA).

**Gene Analysis in Public Datasets**

The Oncomine database (http://www.oncomine.org) and UALCAN (http://ualcan.path.uab.edu) database were used to search and analyze the expression levels of DPY30 reported by other studies. The TIMER (http://TIMER.cistrome.org) online tool was used to observe the pan-cancer analysis of DPY30 mRNA expression. DNA methylation of DPY30 in skin cutaneous melanoma (SKCM) was estimated and visualized using MEXPRESS (https://mexpress.be) and UALCAN. cBioPortal (https://www.cbioportal.org/) was used to search and download mutation information. SangerBox (https://www.SangerBox.com/) was used to analyze immune checkpoint (ICP), tumor mutational burden (TMB), and neoantigens. TIMER (cistrome.shinyapps.io/timer) was used to investigate the correlations between DPY30 and tumor infiltrating immune cells. TISIDB (http://cis.hku.hk/TISIDB/index.php) and GEPIA (http://gepia.cancer-pku.cn) were used to explore the correlations between DPY30 expression and immune or molecular subtypes of different cancer types.
Tissue Microarray and Evaluation of Immunostaining

The commercial melanoma tissue arrays were constructed by Avilabio Co. Ltd. in Shaanxi, China. Briefly, the tissue microarrays including 50 melanoma patients were prepared from archival formalin-fixed, paraffin embedded tissue blocks. Immunohistochemical (IHC) staining was performed as previously described. Protein expression levels were determined using this formula: overall score = positive percentage score × intensity score. The staining intensity was scored as negative (0), weak (1), moderate (2) or strong (3). The percentage of positive cells was scored as 0 (0–10%), 1 (10–25%), 2 (25–50%) or 3 (50–75%) or 4 (>75%). DPY30 in nucleus was defined as negative (combined score from 0 to 6) or positive (combined score from 8 to 12). PD-L1 expression was considered positive when ≥5% membranous staining of tumor cells were positive, as described in previous studies.

Real-Time PCR (RT-PCR) Analysis

Total RNA was isolated using the Trizol reagent (15596–026; Invitrogen, USA). Reverse transcription (RT) was performed using the TransScript First-strand cDNA Synthesis SuperMix (TransGen), according to the manufacturer’s protocol. The primer sequences were as follows: DPY30 forward primer: 5′-AACGCAGGTTGCAGAAAATCCT−3′, Reverse primer, 5′-TCTGATCCAGGTAGCCACGAG−3′; PD-L1 forward primer: 5′-GCTGTTGAAGGACGACGTCT-3′, Reverse primer, 5′-TGAGGATGTGCCAGAGGTA-3′; GAPDH forward primer: 5′-AATCCCATCACCACCTTCC-3′, Reverse primer, 5′-CATCAGCAGTCCATCACCACCTTCC-3′.

Chromatin Immunoprecipitation (ChIP)

Interaction of H3K4me3 in the PD-L1 promoters was examined using Chromatin Immunoprecipitation Kits. Briefly, MMAC-SF and WM-115 cells were used for each ChIP. DNA-protein complex was crosslinked with 1% formaldehyde, lysed with IP lysis buffer, digested with micrococcal nuclease, ultrasonicated, and incubated with anti-H3K4me3. After incubation with protein G beads and washing with wash buffers, chromatin fragments were harvested using the elution buffer. Finally, after decrosslinking and protein digestion, the resulting DNA fragments were purified and fold enrichment was examined using qPCR.

Knockdown by siRNA

Small interfering RNAs for gene DPY30 were purchased from Qiagen (validated FlexiTube siRNA). Transfection of MMAC-SF and WM-115 cells were carried out using Lipofectamine® RNAiMAX vehicle according to the manufacturer’s instruction following optimization. As a negative control, a non-targeting sequence siRNA was employed. Knockdown at mRNA level was confirmed by RT-PCR. RT-PCR was carried out on an ABI PRISM 7900HT sequence detection system (Applied Biosystems).

Preparation of Human T Cells and Co-Culture

The human T cells were isolated from the three healthy volunteers using a human T cell isolation kit (Miltenyi Biotec) according to the manufacturer’s protocol. T cells were cultured in Human ImmunoCult-XFT Cell Expansion medium (Stem Cell) with penicillin (100 u/mL) and streptomycin (100 mg/mL) at 37°C with 5% CO₂. T cells were prestimulated with IFN-γ (Peprotech) for 48 hours before co-culture with MMAC-SF cells. After 24 hours of co-culture, PD1⁺ T cells apoptosis was determined by flow cytometry.

Flow Cytometry

Melanoma cells and human T cells were collected and incubated with the appropriate antibodies in room temperature for 30 min. After washing, the appropriate FITC or DyLight649-conjugated antibodies were used to combine with the first antibodies and were analyzed using FlowJo software (v10, BD, USA). For IFN-γ treatment, cells were first transfected with siRNA mimics as described above, they were treated with 1 ng/mL or 5 ng/mL IFN-γ. Then, cells were collected for flow cytometry. Co-culture with MMAC-SF cells and T cells were harvested, and washed twice then resuspended in 1 mL binding buffer as per BD Biosciences manufacturer’s protocol. 5 μL Annexin V-FITC and
5 μL PI were added to the cell suspension for staining in the dark for 15 min. Cells were then analyzed by flow cytometry.

**Western Blotting**

Total protein was extracted using RIPA lysis buffer supplemented with 1% cocktail and 1% PMSF. The concentration of proteins was measured by the BCA protein assay kit. The quantified proteins were resolved by 10% SDS-PAGE gels electrophoresis and then transferred to a PVDF membrane. Blots were blocked with 5% non-fat dry milk for 1 h at 37°C, and then primary antibodies were probed for 2 h against DPY30 (1:500), PD-L1 (1:2000) and β-actin (1:1000) at a room temperature. Corresponding secondary antibodies were labeled with horseradish peroxidase (1:1000) and incubated for 1.5 h at 37°C. Finally, the blots were placed on film using the enhanced chemiluminescent system (ECL, Bio-Ras Laboratories, Hercules, CA, USA).

**Statistical Analysis**

Statistical analyses were performed using Student’s t-test comparing 2 groups. Statistical analysis was performed with SPSS 23.0 statistical software (Chicago, IL, USA). Here, the difference was indicated statistically significant when $P < 0.01$ and $P < 0.05$.

**Results**

**DPY30 Is Upregulated in Melanoma**

To explore the roles of DPY30 in melanoma progression, we firstly analyzed its expression in Oncomine database. The results showed that the mRNA level of DPY30 in melanoma was higher than in normal tissues (Figure 1A). From the UALCAN database we also identified that the mRNA level of DPY30 was higher in metastatic melanoma compared with primary melanoma (Figure 1B). Meanwhile, we found that DPY30 had a high expression level in almost all cancer cell lines (Figure 1C). In addition, we analyzed the genetic alterations of DPY30 on the outcomes of SKCM patients via the cBioPortal database. Based on TCGA, we found that DPY30 was altered in 5% of SKCM patients (Figure 1D), which included shallow deletion, diploid, gain, and amplification (Figure 1E and F). However, the genetic alterations of DPY30 represented no significant impact on the overall survival rates of SKCM patients (Figure 1G). Then, we analyzed the methylation status of DPY30 on the progression of SKCM patients through cBioPortal and UALCAN database. The results revealed that expression of DPY30 mRNA was significantly negatively correlated with its methylation status (Figure 2A). From the UALCAN database we also identified that the degree of DPY30 gene methylation was inversely related to the tumor’s stage (Figures 2B and C). By using the MEXPRESS to explore, we found that among 22 predicted CpG sites of DPY30, 8 CpG sites, including cg16143319 ($r = -0.139$, $P < 0.01$), cg17781879 ($r = -0.102$, $P < 0.05$), cg17678946 ($r = -0.092$, $P < 0.05$), cg02727605 ($r = -0.126$, $P < 0.01$), cg04302752 ($r = -0.185$, $P < 0.001$), cg07656391 ($r = -0.117$, $P < 0.05$), cg06995575 ($r = -0.205$, $P < 0.001$), and cg24220907 ($r = -0.127$, $P < 0.01$), exhibited negative correlation with DPY30 expression (Figure 2D).

**DPY30 Expression Is Related to ICP, TMB and Neoantigen**

ICP, TMB, and neoantigens in tumor microenvironment were related to antitumor immunity and could predict the therapeutic efficacy of tumor immunotherapy.\(^{25-27}\) Then, we analyzed the correlations between DPY30 expression and ICP, TMB, and neoantigens. We utilized SangerBox and TISIDB to explore the relationships between DPY30 and immune checkpoint genes involved in T-cell exhaustion. Our results indicated that DPY30 expression was positively associated with CD274 ($r = 0.489$, $P = 0.0326$), CTLA4 ($r = 0.098$, $P = 0.0342$), CD160 ($r = 0.147$, $P = 0.001$), HAVCR2 ($r = 0.1$, $P = 0.029$), IDO2 ($r = 0.14$, $P = 0.002$), and PDCD1LG2 ($r = 0.2$, $P < 0.0001$) (Figure 3A). By using the SangerBox to explore, we identified that DPY30 was positively associated with TMB levels ($r = 0.20$, $P < 0.0001$) (Figure 3B). By analyzing the relationships between DPY30 expression and neoantigens, we found that DPY30 expression was positively related to neoantigens ($r = 0.20$, $P = 0.049$) (Figure 3C). The above results demonstrated that DPY30 might be involved in cancer immune evasion in melanoma.
Figure 1 DPY30 expression in melanoma. (A) DPY30 gene copy number in Haqq melanoma vs normal tissues. (B) Comparison of DPY30 expression in primary and metastatic melanoma. (C) DPY30 expression in different cancers and normal tissues in TCGA data. (D) DPY30 gene expression and mutation analysis in SKCM. (E) The distribution of DPY30 genomic alterations in SKCM. (F) The graph of the correlation between DPY30 expression and copy number alterations in SKCM. (G) Kaplan–Meier curve of OS in SKCM patients with altered (red) and unaltered (blue) DPY30 gene. *P < 0.05, **P < 0.01, ***P < 0.001.
Thus, we further performed correlation analyses to investigate the association between DPY30 and immunosuppressive TME by using TIMER and GEPIA database. The results indicated that DPY30 expression was positively correlated with immunosuppressive cells including myeloid-derived suppressor cells (MDSCs) (CD33: $r = 0.315$, $P = 0.0014$; ITGAM: $r = 0.1$, $P = 0.029$; FUT4: $r = 0.36$, $P < 0.0001$), tumour-associated macrophages (TAMs) (CD163: $r = 0.15$, $P < 0.001$; CCL2: $r = 0.12$, $P = 0.011$; IL10: $r = 0.21$, $P < 0.001$), and regulatory T cells (Tregs) (STAT5B: $r = 0.19$, $P < 0.0001$; CCR8: $r = 0.11$, $P = 0.018$; ITGAE: $r = 0.35$, $P < 0.0001$) (Figure 4A).

We further assessed the correlation of DPY30 expression with chemokines using TISIDB database. The result revealed that DPY30 was negatively associated with chemokines, such as CCL14 ($r = -0.115$, $P = 0.012$), CX3CL1 ($r = -0.178$, $P = 0.0001$), and CCL18 ($r = -0.134$, $P < 0.001$).
Taken together, our findings revealed that DPY30 was closely associated with the immunosuppressive TME in SKCM.

**DPY30 Over-Expression Is Positively Associated with PD-L1 Expression**

PD-L1 played essential roles in reprogramming the immunosuppressive TME, so we firstly analyzed the association between DPY30 and CD274 in melanoma by using TIMER and GEPIA. We found that the expression of DPY30 and CD274 was positively correlated (Figure 5A, Cor = 0.274, p = 1.41e-9; Figure 5B, R = 0.22, p = 1.3e-6). We further evaluated the levels of DPY30 and PD-L1 in melanoma tissues in human melanoma tissue arrays. The results showed that 31 of 50 of the primary lesions exhibited positive staining for DPY30, and DPY30 protein was localized in the nucleus of melanoma cells. Then, we detected PD-L1 in the same type of tissue arrays. The results demonstrated that the positive rates of PD-L1 was 58% (29/50), and PD-L1 protein was detected on the cell membrane and in the cytoplasm of melanoma cells (Figure 5C). Pearson correlation analysis demonstrated that the expression of DPY30 was closely associated the level of PD-L1, and DPY30 was positively correlated with PD-L1 (r = 0.336, p = 0.017) (Table 1).
The DPY30-H3K4me3 Axis-Mediated PD-L1 Expression

To study the regulation between DPY30 and PD-L1, we used the siRNA targeting DPY30 in MMAC-SF cells. Silencing of DPY30 by specific siRNA significantly inhibited PD-L1 expression in MMAC-SF cells, and this was supported by Western blot and RT-PCR (Figure 6A and B). To determine whether this decrease in PD-L1 expression was due to DPY30-mediated chromatin modifications, ChIP analysis revealed that H3K4me3 levels were enriched in the proximal PD-L1 promoter region in MMAC-SF cells (Figure 6C). In addition, TISIDB was used to analyze the function of DPY30 and found that DPY30, as part of the MLL1/MLL complex, is involved in the methylation of histone H3K4 especially trimethylation (Supplementary Table 1). These results suggested that DPY30 may mediate PD-L1 expression through H3K4me3.

Figure 4 Correlations of DPY30 with immune infiltration level in SKCM. (A) Correlations of DPY30 expression with immunosuppressive cells and markers of (MDSCs, TAMs, and Tregs). (B) Correlation between DPY30 expression and chemokines.
DPY30 Regulates the Expression of PD-L1

IFN-γ induces PD-L1 expression and disruption of this pathway might sensitize melanoma patients to immune-based therapies. To further study the regulation between DPY30 and PD-L1, we detected the PD-L1 level by IFN-γ treatment.

![Figure 5](https://doi.org/10.2147/JIR.S377678)

**Figure 5** The expression of DPY30 correlates with PD-L1 in human melanoma tissues. (A and B) Spearman correlation analysis of DPY30 expression with CD274 expression. (C) Representative IHC staining of DPY30 and PD-L1 in the same melanoma tissue. Red arrows point to the location of DPY30 protein and PD-L1 protein.

**Table 1** Correlation Between the Expression of DPY30 and PD-L1

|        | PD-L1     |
|--------|-----------|
| **DPY30** | Correlation Coefficient | 0.336* |
|        | Sg. (2-tailed)          | 0.017  |
|        | N                      | 50     |

**Note:** *Correlation is significant at the 0.05 level (2-tailed).
Figure 6: Function of DPY30 in regulation of PD-L1 expression. (A and B) MMAC-SF cells were transiently transfected with DPY30-specific SiRNA, and the protein levels of DPY30 and PD-L1 were analyzed by Western blotting, and the mRNA levels of DPY30 and PD-L1 were detected by RT-PCR. (C) The H3K4me3 levels in the PD-L1 promoter region in MMAC-SF and WM-115 cell lines were analyzed by ChIP. (D and E) MMAC-SF cells and MMAC-SF Si-DPY30 cells were treated with IFN-γ at the prescribed dose for 24 h or 48h, and analyzed for PD-L1 mRNA level by RT-PCR. (F and G) MMAC-SF cells and MMAC-SF Si-DPY30 cells were treated with IFN-γ at the prescribed dose for 24 h or 48h, stained with PD-L1 specific MAb, and analyzed by flow cytometry. **P<0.01.
and down-regulation of DPY30 in MMAC-SF cells. The results of RT-PCR showed that the level of PD-L1 in MMAC-SF si-DPY30 cells remained unchanged in the 1ng IFN-γ treatment group and slightly increased in the 5ng IFN-γ treatment group compared with the non-IFN-γ treatment group at 24 hours of IFN-γ treatment. The overall PD-L1 level at 48 h after IFN-γ treatment was higher than at 24h, but the trend kept the same as that at 24h (Figures 6D and E). Moreover, the results of flow cytometry indicated that knockdown of DPY30 gene could significantly suppress PD-L1 levels after IFN-γ (1 or 5 ng) treated MMAC-SF cells for 24 hours. After 48 hours of IFN-γ treatment, Knockdown of DPY30 still suppressed PD-L1 levels (Figures 6F and G). To further validate the effect of down-regulation of DPY30 in MMAC-SF cells, and we detected the apoptosis of PD1+ T cells by flow cytometry in a co-culture model. We detected PD-1 level in human T cells from 3 normal samples by flow cytometry (Figure 7A), and the apoptosis of PD1+ T-cells in co-culture with MMAC-SF cells by knockdown of DPY30 were markedly reduced (Figure 7B).

**Discussion**

As the major H3K4 methylation enzymes in mammals, the SET1/MLL complexes are considered potential pharmacologic targets in epigenetic therapeutics. DPY30, a core subunit of SET1/MLL complexes, is important for facilitating genome-wide H3K4 methylation. Accumulating evidence have revealed significant upregulation of DPY30 mRNA expression were observed in hepatocellular carcinoma, cervical squamous cell carcinoma, ovarian cancer, and cholangiocarcinoma. Our study showed that the mRNA level of DPY30 in melanoma was higher than in normal tissues. In addition, DPY30 could induce epithelial to mesenchymal transition by activating Wnt/β-catenin
signaling in the progression of cervical squamous cell carcinoma. Similarly, our research also proved that the mRNA level of DPY30 was higher in metastatic melanoma compared with primary melanoma. These findings suggested that DPY30 over-expressed in melanoma, which might play essential roles in the progression of melanoma.

DNA methylation is an important biological process involving in cancer development, and changes in DNA methylation in cancer have been regarded as diagnostic or prognostic biomarkers. A previous study indicated that 36 DNA methylation markers from 304 patients with hepatocellular carcinoma could predict poor survival in patients with hepatocellular carcinoma. Furthermore, comprehensive genome-scale analysis of aberrant DNA methylation enabled classification of colorectal carcinoma patients based on their prognosis. In this study, we analyzed the methylation status of DPY30 gene through the UALCAN database. The analysis presented that the degree of DPY30 gene methylation was negatively correlated with tumor stage, and the expression of DPY30 mRNA was significantly negatively correlated with its methylation status. Furthermore, among 22 predicted CpG sites of DPY30, 8 CpG sites exhibited negative correlation with DPY30 expression. These findings suggested that the methylation status of DPY30 might be a novel direction for prognosis evaluation of melanoma.

Immune checkpoint play essential roles in regulating the immunosuppressive TME. Immune checkpoint inhibitors, including anti-PD-1 and anti-CTLA4 antibodies, have revolutionized the management of melanoma and significantly improved the clinical outcomes for patients with melanoma. However, immune checkpoint inhibitors are only effective in a part of patients with melanoma, and two-thirds of patients are resistant and will require further treatment. Our study indicated that DPY30 expression had significant positive associations with ICP genes involved in T-cell exhaustion, including CD274, CTLA4, CD160, HAVCR2, IDO2, and PDCD1LG2. Further studies indicated that DPY30 was positively correlated with immunosuppressive cells in SKCM including MDSCs, TAMs, and Tregs, suggesting that DPY30 was closely associated with the immunosuppressive TME in SKCM. IHC results showed that the positive rates of DPY30 and PD-L1 in melanoma tissues were 62% and 58%, respectively. Correlation analysis revealed that DPY30 was positively associated with PD-L1 expression. Furthermore, we found that silencing of DPY30 by specific siRNA significantly inhibited PD-L1 expression. Therefore, we believe that DPY30 might be involved in cancer immune evasion in melanoma.

Histone H3K4 methylation is one of the most significant epigenetic modifications, and plays critical roles in regulating target gene transcription. The most notable H3K4 methylation enzymes in mammals are the SET1/MLL complexes, which have a profound effect on numerous biological processes including cancer progression. SET1/MLL family complexes comprise Set1a, Set1b, MLL1, MLL2, MLL3, and MLL4 as the catalytic subunit, and Wdr5, Rbpb5, Ash2l, and DPY30 as the common core subunits. The roles of the MLL/SET1 complexes in transcriptional regulation have been extensively studied. Among these subunits, MLL1 has been shown to bind to the CD274 promoter to catalyze H3K4me3 to activate PD-L1 transcription in tumor cells. Inhibition of MLL1 activity or silencing of MLL1 expression reduced H3K4me3 level in the CD274 promoter region and downregulated PD-L1 expression in tumor cells. Previous study showed that MLL3 was located in the enhancer of PD-L1 and regulated the H3K4me1 level in the enhancer regions. Depletion of MLL3 reduced the binding of H3K4me1 in the PD-L1 enhancer regions and Pol II Ser-5p in PD-L1 promoter regions. Furthermore, it has been reported that WDR5 was positively correlated with PD-L1 expression, and WDR5 inhibitor inhibited PD-L1-based immune evasion by blocking the WDR5-MLL complex mediating H3K4me3 of target genes. DPY30 is a core subunit of SET1/MLL complexes, and it has been reported that DPY30 promoted the expression of endogenous MYC by regulating chromatin accessibility. Previous study also showed that DPY30 regulated angiogenesis and tumor growth in an H3K4me3-dependent manner, and PDE4B was a key downstream effector of DPY30-H3K4me3 axis. Targeting DPY30-PDE4B axis was an effective approach for treatment of glioblastoma. DPY30 promoted the expression of Hif-1α and its targeted glycolytic genes by regulating H3K4me3 modification. In this study, we found that H3K4me3 levels were enriched in the proximal PD-L1 promoter region in melanoma cells. Inhibition of DPY30 suppressed the PD-L1 level in IFN-γ treated melanoma cells. Our study demonstrated that DPY30 might regulate PD-L1 expression through H3K4me3, which might be a promising therapeutic target in melanoma.

**Conclusion**

In summary, our study firstly demonstrated that the elevated DPY30 levels in melanoma tissues, and the expression of DPY30 was closely related to the level of PD-L1. DPY30 was associated with cancer immune evasion by regulating PD-L1 expression through epigenetic mechanism in melanoma.
Abbreviations
APCs, antigen-presenting cells; ChIP, Chromatin Immunoprecipitation; CTLA-4, cytotoxic T lymphocyte antigen 4; H3K4me3, histone H3K4 trimethylation; IHC, Immunohistochemical; ICP, immune checkpoint; MDSCs, myeloid-derived suppressor cells; NK, natural killer; PD-1, programmed cell death protein 1; PD-L1, programmed cell death ligand 1; TAMs, tumour-associated macrophages; TMB, tumor mutational burden; Tregs, regulatory T cells.

Data Sharing Statement
The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found in the article.

Ethics Approval and Informed Consent
The study’s protocol was approved by the ethics committee of Cancer Hospital, Chinese Academy of Medical Sciences, Peking Union Medical College, and informed consent was obtained from the study participants. Our experiments followed the guidelines outlined in the Declaration of Helsinki.

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Disclosure
The authors declare no conflicts of interest for this work.

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