Original Article

Magnetic resonance imaging monitoring therapeutic response to dendritic cell vaccine in murine orthotopic pancreatic cancer models

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Abstract: Pancreatic ductal adenocarcinoma (PDAC) carries the worst prognosis and caused one of the highest cancer-related mortalities. Dendritic cell (DC) vaccination is a promising cancer immunotherapy; however, the clinical outcomes are often poor. The administration route of DC vaccine can significantly alter the anti-tumor immune response. Here we report on the cytotoxic T lymphocyte (CTL) responses induced by DC vaccination administered via intraperitoneal (IP) for murine PDAC, and the longitudinal assessment of tumor growth and therapeutic responses using magnetic resonance imaging (MRI). In this study, we established murine orthotopic Panc02 models of PDAC and delivered apoptotic Panc02 cell-pulsed DCs via IP injection. The migration of Panc02-pulsed DCs into spleens significantly increased from 6 h to 12 h after initiation of treatment (P = 0.002), and Panc02-pulsed DCs injected via IP induced a significantly higher level of CTL responses against Panc02 cells compared to unpulsed DCs. Tumor size and tumor apparent diffusion coefficient (ADC) were measured on MR images. Tumor sizes were significantly smaller in the treated mice than in the untreated mice (P < 0.05). The reduction of tumor ADC was less in the treated mice than in the untreated mice (P < 0.05), and the changes in tumor ADC showed significant negative correlation with the changes in tumor volume (r = -0.882, 95% confidence interval, -0.967 to -0.701, P < 0.0001). These results demonstrated the efficacy of DC vaccination administered via IP injection in murine PDAC, and the feasibility of ADC measurement as an imaging biomarker for assessment of therapeutic responses in immunotherapy.

Keywords: Pancreatic ductal carcinoma, dendritic cell, vaccination, intraperitoneal injection, magnetic resonance imaging

Introduction

Pancreatic ductal adenocarcinoma (PDAC) accounts for over 95% of all pancreatic malignancies. The 5-year survival rate is approximately 7.1% and the median overall survival (OS) is 4-6 months for all patients with PDAC [1, 2]. Surgical resection, the only potentially curative treatment, is applicable in only 5% to 25% patients; even after surgery, the 5-year survival rate is less than 20% [3]. Local ablation and systemic chemotherapy have offered little benefit [4]. Thus, new approaches are urgently needed to treat this deadly disease effectively.

Recently, many immunotherapies are based on the vaccination of cancer patients with autologous DCs loaded with tumor antigens (DC vaccines) [5-7]. DCs are the main type of antigen-presenting cells (APCs), and the aim of DC vaccination is to induce tumor-specific effector T cells that can specifically reduce the tumor mass and induce immunological memory to control tumor recurrence [8-10]. DC vaccination has clinically relevant mechanisms of action
with a great potential for the systemic treatment of cancers in a clinical setting. However, clinical trials have demonstrated its poor therapeutic efficacy [11-13]. The efficacy of DC vaccination is strongly influenced by DC vaccine migration to the draining lymph nodes (LNs). It has been reported that the administration route of DC vaccine can significantly alter the anti-tumor immune response and immune memory [13, 14]. Administration routes including subcutaneous, intradermal, intratumoral, or intranodal injection have been applied in both preclinical research and clinical trials [15, 16]. However, only a small number of DCs can migrate to LNs and lymphoid organs by subcutaneous or intradermal injection [17, 18]. A direct delivery of DCs by intratumoral injection or by intranodal injection could impair the efficacy of DC vaccine [18, 19]. Given that a large number of LNs in the abdomen composes the most important secondary lymphoid organ and the fact that spleen is the largest lymphatic organ in the abdomen, we have very recently tested the migration of DCs to LNs via intraperitoneal (IP) injection, and the results indicated that IP injection could increase the amount of viable DC vaccines actively absorbed by LNs and improve outcomes.

Magnetic resonance imaging (MRI), which can provide anatomic and functional information about tumors, has a well-established role in the evaluation of patients with PDAC [20]. Diffusion weighted MRI (DW-MRI), which reflects the diffusion of water molecules within biological tissues, provides microstructural information at the cellular level [21, 22]. The apparent diffusion coefficient (ADC), which can be quantified from DW-MRI, decreases when water molecule movements are hindered by any obstacle; in particular, it has been proved to reduce in tissues with high cell density [23]. Studies on a variety of tumor types reported that tumor ADC could act as an imaging biomarker for early detection of cancer treatment response [24-27]. To the best of our knowledge, no studies have assessed DC vaccine therapeutic responses in PDAC using DWI.

**Materials and methods**

Our study was approved by the Institutional Animal Care and Use Committee (IACUC) of Northwestern University.

**Cell lines and culture**

The mouse Panc02 cell line is derived from amethylcholanthrene-induced pancreatic ductal adenocarcinoma in C57BL/6 mice and was purchased from the American Type Culture Collection (ATCC; Rockville, MD). Panc02 cells were cultured in RPMI 1640 medium (Life Technologies, Carlsbad, CA) supplemented with glutamine (2 mmol/l, Life Technologies), pyruvate (1 mmol/l, Sigma-Aldrich, St. Louis, Mo), penicillin and streptomycin (100 IU/ml, Sigma-Aldrich), and 10% fetal bovine serum (FBS, Sigma-Aldrich). The cells were maintained in a humidified atmosphere of 5% CO₂ at 37°C. Before implantation, cell viability was assessed using trypan blue (Sigma-Aldrich) staining (cell viability of > 90% was confirmed prior to tumor implantation).

**UV-B irradiation of Panc02 cells**

Panc02 cells were diluted in phosphate-buffered saline (PBS) at a final concentration of 2 × 10⁶/ml, and were placed in a 6-well plate at 0.5 ml per well. Panc02 cells were irradiated using UV-B light with 0.75 J/cm². After irradiation, cells were maintained in culture medium. 12 h later, Panc02 cell apoptosis was confirmed by flow cytometric analysis using Annexin V/PI staining (Figure 1). The inactivated Panc02 cells were collected for co-incubation with DCs.

**Generation and antigen loading of bone marrow-derived DCs**

The mouse bone marrow-derived DCs (BMDCs) were prepared as described in a previous publication with some modifications [28]. Briefly, female C57BL/6 mice (5-6 weeks of age, weighted between 16 and 19 g; Charles River, Wilmington, MA) were sacrificed by CO₂ narcosis and cervical dislocation, and the tibias and femurs were harvested. After immersion in 70% ethanol for 5 mins, the bones were carefully dissected, and each half was flushed with FBS-free RMPI 1640 (Gibco, Waltham, MA) in a cell
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culture hood. The bone marrow cells were collected, and erythrocytes were lysed with ammonium chloride buffer (BD Biosciences, San Jose, CA). A total of $2 \times 10^6$ cells was cultured in 10 mL RPMI supplemented with 10% FBS (Gibco, Waltham, MA), 100 units/mL penicillin, 100 µg/mL streptomycin, 0.25 µg/mL amphotericin b (antibiotic-antimycotic 100 ×, Gibco, Waltham, MA), 10 ng/mL rm-GM-CSF, and 1 ng/mL rm-IL-4 (both from Shenandoah Biotechnology, Warwick, PA). Equal volume of the same culture media was added after 48 h. At day 7, lipopolysaccharide (LPS, 250 ng/ml) and interferon γ (IFNγ, 100 ng/ml) were added for 24 h. Loosely adherent cells were then harvested, and the expression of DC markers was analyzed by fluorescence-activated cell sorting (FACS) as described below. BMDCs were co-incubated with the inactivated Panc02 cells at a DC/Panc02 cell ratio of 10:1 for 24 hours. At day 9, loosely adherent cells (Panc02-pulsed DCs) were harvested, washed and resuspended in PBS, and analyzed by FACS.

FACS

Pan02-pulsed DCs were collected after 8 d of in vitro culture and washed with cold PBS. BMDC were stained by incubation for 40 mins at 4°C with 2 µg/3 × 10^5 cells anti-mouse PerCP-Cy5.5 CD11c monoclonal antibody (mAb), PerCP-Cy5.5 CD11b mAb, PE CD80 mAb, APC CD86 mAb, PE H2Db mAb, FITC H2Kb mAb (all from BD Bioscience, San Jose, CA), PE MHC class II mAb (Southern Biotech, Birmingham, AL), and appropriate isotype controls. The expression of DC markers was quantified by fluorescence-activated cell sorting (FACS) using flow cytometry (BD LSRFortessa™ cell analyzer, San Jose, CA) and analyzed using FlowJo (Ashland, OR).

Migration of Panc02-pulsed DCs to spleens via IP injection

Migration of Panc02-pulsed DCs to spleens after IP injection was visualized, as spleen is the largest lymphoid organ in the abdominal cavity. $1 \times 10^7$ Panc02-pulsed DCs in 100 µL Diluent C were mixed with 0.4 µL PKH26 dye in equal volume of Diluent C for 5 mins (Sigma-Aldrich, St. Louis, MO). After adding 200 µL FBS and washing 3 times with PBS, the DCs were labeled with PKH26. Twelve C57BL/6 mice were IP injected with $1 \times 10^7$ PKH26-labeled Panc02-pulsed DCs in 100 µL PBS. 6 mice were randomly euthanized at each time point (6 h and 12 h after IP injection) to harvest the spleens for fluorescence staining. The collected spleen tissues were embedded in OCT (Fisher HealthCare, Houston, TX) infused modes that were placed on dry ice and frozen at -80°C after 2 mins. The frozen samples were cut into 5 µm thick slices with the microtome and placed on slides. Then the slides were mounted by cover glasses with ProLong Glod Antifade Reagent with DAPI (Cell Signaling Technology, Danvers, MA) and visualized by fluorescent microscope (Axioimager Z1, Carl Zeiss, Ontario, CA). Three representative fluorescent microscopy images were acquired from each sample with the same filter settings at the same magnifications. Total cells counts and PKH26 positive cells counts from the fluorescence microscopy images were quantified using ImageJ software (NIH, Bethesda, MD). The ratio of PKH26 positive cells to total number of cells was calculated for each sample.

Orthotopic mouse model of pancreatic cancer

Female C57BL/6 mice were used for establishing orthotopic pancreatic cancer models. The orthotopic mouse model of pancreatic cancer was prepared using previously published protocols with modifications [29]. $1 \times 10^5$ viable Panc02 cells were gently mixed with ice-cold...
Matrigel (Sigma-Aldrich) at a ratio of 3:1 to produce a homogeneous suspension. Anesthesia was induced and maintained in each mouse by inhalation of 2% isoflurane in oxygen at a rate of 1 L/min (Isoflurane Vaporizer, Vaporizer Sales and Services, Rockmart, GA). After local shaving and disinfection, the abdominal cavity was opened by a 1.5-cm longitudinal incision at the left upper quadrant. The tail of the pancreas was identified by mobilization of the spleen. 5 μL of Panc02 cell-Matrigel suspension was then slowly injected into the parenchyma of pancreatic tail. To prevent further leakage, the needle was kept in the injection site for 30 s prior to removal. The spleen and pancreas were placed back into the abdominal cavity, and the abdominal cavity was closed by a running two-layer silk suture. Postoperative status and wound healing were monitored every day for one week. After one week, a visible nodule at the location of pancreatic tail was detected by magnetic resonance imaging (MRI) in all eighteen mice (protocols described below), which indicated that the orthotopic pancreatic cancer models were established successfully.

**Therapeutic strategy**

Twelve mice with orthotopic pancreatic cancer were randomly divided into two groups: treatment group (n = 6) and control group (n = 6). Therapeutic strategy was started one week after tumor implantation. The mice in the treatment group underwent IP injection of $1 \times 10^7$ Panc02-pulsed DCs in 100 μL PBS for therapeutic vaccination, and the mice in the control group underwent IP injection of 100 μL saline solution. IP injection was administered once a week for 5 weeks.

**CTL assay**

Twelve orthotopic pancreatic cancer mice and six age-matched normal C57BL/6 mice were used for CTL assay. One week after tumor implantation, six of the mice with orthotopic pancreatic cancer were IP injected with $1 \times 10^7$ Panc02-pulsed DCs in 100 μL PBS, and the other six were IP injected with $1 \times 10^7$ unpulsed mature DCs in 100 μL PBS. IP injection was administered once a week for 3 weeks. The six normal C57BL/6 mice that did not undergo any operation were used as another control group. After sacrifice, spleens were harvested and homogenized to release splenocytes in RPMI 1640 medium. The single-cell suspension was prepared, and CD8+ T cells were isolated using magnetic cell sorting by negative selection (CD8a+ T cell isolation Kit, Miltenyi Biotec) according to the manufacturer’s instructions. Panc02-pulsed DC-induced, unpulsed DC-induced, and normal murine CD8+ T cells were prepared. Flow cytometry confirmed > 95% purity. Lactate dehydrogenase (LDH) assay was performed to determine the cytotoxicity of the purified CD8+ T cells using Cytotoxicity Detection KitPLUS (Sigma-Aldrich). For LDH assay, Panc02 cells (target cells) were collected and plated at 3000 cells/well in a flat-bottom 96-well cell culture plate (BD Biosciences, San Diego, CA, USA). Then, CD8+ T cells (effector cells) were added to the 96-well plates at different effector-to-target (ET) ratios (5:1, 10:1, 20:1 and 40:1), and 2% Triton X-100 lysis solution was added to the wells that were prepared for maximal LDH release measurement. The plate was incubated at 37°C for 4 h. After incubation, the plates were centrifuged for 10 min at 330 × g, and 100 μL/well cell-free supernatant was transferred into the corresponding wells of an optically clear, flat-bottom 96-well plate. Then, 100 μL mixed detection kit reagent was added to each well, and cells were incubated at room temperature in the dark for 30 min. After incubation, the absorbance was measured using a multiwell plate reader at 490 nm. The percentage of cytotoxicity was calculated according to the following equation: cytotoxicity (%) = \( \left( \frac{ET - E}{Max - T} \right) \times 100 \), where \( ET \) is the LDH release by effector-target reaction, \( E \) is the spontaneous LDH release of effector cells, \( T \) is the spontaneous LDH release of target cells, and \( Max \) is the maximal LDH release from complete target cell lysis.

**MRI examination**

MRI examinations were performed by using a 7.0 T small-animal MRI scanner with a commercial mouse coil (ClinScan, Bruker Biospin). One week after tumor implantation, all the mice underwent MRI examination to confirm the successful establishment of orthotopic pancreatic cancer models. The mice in the treatment group and the control group underwent MRI examination at one week, three weeks, and five weeks after initiation of treatment for monitoring tumor growth and evaluating the response to therapies in vivo. Mice were anesthetized by
inhalation of a mixture of 2% isoflurane and oxygen at 1 L/min, and body temperature was monitored continuously and was controlled with a water bed (SA Instruments, Stony Brook, NY). The MRI sequences and parameters were as follows: (a) axial T1-weighted imaging (T1WI): repetition time (TR)/echo time (TE) = 630/20 ms; field of view (FOV) = 27 mm × 30 mm; matrix size = 134 × 192; slice thickness (ST) = 0.8 mm; gap = 0 mm; (b) axial T2-weighted imaging (T2WI): TR/TE = 1581/40 ms; FOV = 21 mm × 30 mm; matrix size = 154 × 192; ST = 0.8 mm; gap = 0 mm; (c) coronal T2WI: TR/TE = 2500/40 ms; FOV = 23 mm × 32 mm; matrix size = 142 × 192; ST = 0.8 mm; gap = 0 mm; (d) sagittal T2-weighted imaging: TR/TE = 2500/40 ms; FOV = 24 mm × 30 mm; matrix size = 154 × 192; ST = 0.5 mm; gap = 0 mm; (e) axial diffusion-weighted imaging (DWI): TR/TE = 3500/40 ms; FOV = 28 mm × 28 mm; matrix size = 82 × 82; ST = 1 mm; gap = 0 mm; b-value = 0 and 800 s/mm². DWI was performed in 3 orthogonal directions of the diffusion gradients.

MR image analysis

MR images were analyzed by a radiologist with more than 5 years of experience. In vivo tumor size measurement was performed by using an open source software ITK-SNAP (version 3.6.0, University of Pennsylvania) by measuring both the longest diameter and tumor volume. The longest diameter of each orthotopic tumor was measured on either axial, coronal or sagittal T2-weighted images. For tumor volume measurement, free-hand regions of interest (ROIs) were traced along the tumor margin on each slice of axial T2-weighted images containing a orthotopic tumor, and then the three-dimensional volume was calculated [30]. The change in the longest diameter (Δdiameter) after five weeks of treatment was calculated according to the equation: Δdiameter = diameter_{5w} - diameter_{1w}, where diameter_{1w} and diameter_{5w} are the longest diameters that measured at one week and five weeks after initiation of treatment, respectively. The change in tumor volume (Δvolume) after five weeks of treatment was calculated according to the equation: Δvolume = volume_{5w} - volume_{1w}, where volume_{1w} and volume_{5w} are the tumor volumes that measured at one week and five weeks after initiation of treatment, respectively.

Diffusion-weighted images were postprocessed to generate ADC maps in Matlab R2016b (Mathworks, Natick, MA). Using axial T2-weighted images as reference, a free-hand ROI was drawn inside the tumor on the slice of ADC maps with the largest tumor area, avoiding regions with necrosis and artifacts, and the mean ADC for each orthotopic tumor was recorded. The change in tumor ADC (ΔADC) after five weeks of treatment was calculated according to the equation: ΔADC = ADC_{5w} - ADC_{1w}, where ADC_{1w} and ADC_{5w} are, respectively, the tumor ADC that measured at one week and five weeks after initiation of treatment.

Statistical analysis

Statistical analysis was performed using SPSS (version 22.0; SPSS, Chicago, IL). The results are presented as mean ± standard deviation (SD). Mann-Whitney U test was applied to compare the difference between the ratio of PKH26 positive DCs to the total cells in the spleens harvested 6 h and 12 h after IP injection. Repeated measures ANOVA analysis was applied to compare the differences in the longest diameter and tumor volume between the treatment and the control group and the changes over time. Student’s t test was used to compare the differences in ΔADC and Δvolume between the treatment group and the control group. The differences in the cytotoxicity between normal, MDI, and PDI CD8+ T cells were determined using One-Way ANOVA analysis followed by LSD test for multiple comparisons. Pearson correlation coefficient (r) and 95% confidence interval (CI) was calculated to assess the relationship between Δdiameter and ΔADC, and the relationship between Δvolume and ΔADC. For the absolute value of r, 0-0.19 is regarded as very weak, 0.2-0.39 as weak, 0.40-0.59 as moderate, 0.6-0.79 as strong, and 0.8-1.0 as very strong correlation. P < 0.05 was considered to indicate a significant difference in all statistical tests.

Results

Purity and maturity of bone marrow-derived DCs and Panc02-pulsed DCs

The purities of DCs in BMDCs were analyzed on FACS by assessing the population of cells with positive CD11b and CD11c expression. As shown in Figure 2, the population of DCs in
Figure 2. The immunophenotypes of bone marrow-derived cells in vitro culture for 8 days and Panc02-pulsed DCs. A. The immunophenotypes of bone marrow-derived cells in vitro culture for 8 days. B. The immunophenotypes of Panc02-pulsed DCs. The results showed that the purity of CD11b+ DCs and CD11c+ DCs were more than 75% for both bone marrow-derived cells 8 days after in vitro culture and Panc02-pulsed DCs. The high expression of the maturation surface markers CD80, CD86, H2Db, H2Kb and MHC class II indicated that the DCs were in a mature state.
BMDCs that being cultured in vitro for 8 days (Figure 2A) and in BMDCs that being pulsed by Panc02 cells (Figure 2B) both achieved higher than 75%. The maturation of DCs was evaluated by measuring the expression level of maturation surface markers CD80, CD86, H2Db, H2Kb, and MHC class II, indicating that the DCs after in vitro culture were in a mature state that ready to present antigen to T cells (Figure 2).

Migration of fluorescence-labeled Panc02-pulsed DCs to spleens via IP injection

In order to track the migration of Panc02-pulsed DCs to spleen, the levels of PKH26-labeled DCs in the spleen were detected by fluorescence microscopy. For spleens harvested 6 h and 12 h after IP injection, the mean ratios of PKH26 positive DCs to the total number of cells were $(1.24 \pm 0.12) \times 10^{-2}$ and $(2.52 \pm 0.26) \times 10^{-2}$, respectively (Figure 3A and 3B). The ratio of positive DCs in spleens harvested 12 h after IP injection was significantly higher than that harvested 6 h after IP injection ($P = 0.002$, Figure 3C), indicating the accumulative migration of Panc02-pulsed DCs in spleens.

CTL responses induced by IP injection of Panc02-pulsed DCs

To study the anti-tumor toxicity of CTLs that stimulated by Panc02-pulsed DCs, the CD8+ T cells are purified after IP injection of Panc02-pulsed DCs, and its cytotoxicity was evaluated by measuring the LDH that released by co-cultured Panc02 cells. Normal murine CD8+ T cells were used as controls. IP injection of Panc02-pulsed DCs induced a significantly higher level of CTL response against Panc02 cells compared to IP injection of unpulsed DCs or normal murine CD8+ T cells ($P < 0.0001$).
Therefore, the Panc02-pulsed DCs are capable to trigger strong anti-tumor immunity of CD8+ T cells.

**In vivo tumor size measurement on MR images**

Representative axial T2-weighted images of the pancreatic tumors in the treatment group and control group at different therapeutic time points were shown in Figure 5A-F. At one week, three weeks, and five weeks after initiation of treatment, the mean longest diameters were 2.29 ± 0.50 mm, 5.04 ± 1.06 mm, and 8.83 ± 0.97 mm for the treatment group and 3.04 ± 0.87 mm, 6.10 ± 0.48 mm, and 10.93 ± 1.62 mm for the control group respectively (Figure 5G). The mean Δdiameters were 6.21 ± 1.42 mm for the treatment group and 7.89 ± 2.05 mm for the control group. There were no significant differences in the longest diameters between the treatment and the control group one week ($P = 0.097$) and three weeks after initiation of treatment ($P = 0.500$). The longest diameters of the pancreatic tumors in the control group were significantly longer than those in the treatment group five weeks after initiation of treatment ($P = 0.021$). (H) Changes of tumor volumes in the treatment and the control group at different therapeutic time points (error bars indicate SD, *$P < 0.05$). There were no significant differences in tumor volumes between the treatment and the control group one week after initiation of treatment ($P = 0.076$). Tumor volumes in the control group were significantly larger than those in the treatment group three weeks ($P = 0.004$) and five weeks after initiation of treatment ($P = 0.001$).
group and the control group ($P = 0.129$). On the other hand, the tumor volumes have experienced a more dynamic change. The mean tumor volumes were $4.64 \pm 1.50$ mm$^3$, $25.75 \pm 10.64$ mm$^3$, and $110.23 \pm 24.30$ mm$^3$ for the treatment group and $10.08 \pm 6.58$ mm$^3$, $54.42 \pm 16.02$ mm$^3$, and $209.89 \pm 48.69$ mm$^3$ for the control group ($P = 0.076$). Tumor volume in the control group was significantly higher than that in the treatment group at three weeks ($P = 0.004$, Figure 6G). There was no significant correlation between $\Delta$volume and $\Delta$diameter ($r = -0.440$, 95% CI, -0.848 to 0.509, $P = 0.152$), but a very strong negative correlation between $\Delta$ADC and $\Delta$volume ($r = -0.882$, 95% CI, -0.967 to -0.701, $P < 0.0001$, Figure 6I).

ADC measurement

Representative axial ADC maps of pancreatic tumors in the treatment group and control group at different therapeutic time points were shown in Figure 6A-F. The mean $\Delta$ADCs were $(-0.14 \pm 0.13) \times 10^{-4}$ mm$^2$/s for the treatment group and $(-0.54 \pm 0.22) \times 10^{-4}$ mm$^2$/s for the control group. $\Delta$ADC in the treatment group was significantly higher than that in the control group ($P = 0.004$, Figure 6G). There was no significant correlation between $\Delta$ADC and $\Delta$diameter ($r = -0.440$, 95% CI, -0.848 to 0.509, $P = 0.152$, Figure 6H), but a very strong negative correlation between $\Delta$ADC and $\Delta$volume ($r = -0.882$, 95% CI, -0.967 to -0.701, $P < 0.0001$, Figure 6I).
Discussion

This study provided an effective DC vaccination in a murine PDAC model and successfully monitored the tumor response using ADC measurement. Firstly, our results demonstrated that Panc02-pulsed DC vaccination administered via IP injection induces a tumor-specific CTL response and inhibits tumor growth in a murine PDAC model. Following five weeks of DC vaccination, there was a significant difference in the reduction of tumor ADC between the treatment group and the control group; furthermore, a strong correlation was observed between the change in tumor ADC and the change in tumor volume.

In this study, the Panc02 murine orthotopic pancreatic tumor model was used, which is the animal model mimics both histologic characteristics and biological behaviors of human PDAC, and is ideal for testing the efficacy of DC vaccination in the treatment of PDAC [28, 31, 32]. In this study, apoptotic Panc02 cells generated by UV-B irradiation was used for DC vaccine preparation due to its advantage in antigen loading effectiveness and the capacity of loaded DC in T-cells activation [33]. Our results showed that there was a significant increase in the level of CTL response induced by Panc02-pulsed DCs compared with that induced by unpulsed DCs. As for the administration route of DC vaccine, previous studies have shown that DCs failed to migrate into secondary lymphoid tissues after intratumoral injection, resulting in an inferior capacity to induce adaptive immunity [19, 34]. Lesterhuis et al [18] found that the more laborious and variable intranodal route did not offer an advantage over intradermal route. Although intradermal and subcutaneous injection are the most common routes for vaccination in human clinical trials, low migration rate is the major obstacle that leads to low efficacy [35]. IP injection of DC vaccine in this study is easy to apply, and could deliver a relatively large volume of DC vaccine can be injected at multiple time points. The migration of Panc02-pulsed DCs to the spleen and LNs after IP injection was tested, and a large amount of labeled Panc02-pulsed DCs were observed in those lymphoid organs 12 h after IP injection. Our results indicated that IP injection of DC vaccine could be a potential regimen for pancreatic cancer.

In most mouse PDAC therapeutic studies, tumor size was conventionally assessed by standard dissection techniques with the manual caliper measurements, which could only provide the endpoint data. Compared with the conventional assessment method, MRI could provide noninvasive longitudinal in vivo measurements of tumor growth with higher accuracy and repeatability. Besides, the evaluation of cancer therapeutics is conventionally based on measurements of the longest tumor diameters in clinical settings, which are adopted by the revised Response Evaluation Criteria in Solid Tumors guideline (RECIST, version 1.1) [36]. Volume-based size measurements are more effective than one-dimensional or two-dimensional measurements in clinical settings, and the change in tumor volume could be used to predict the overall survival and recurrence-free survival of patients in clinical trials [37]. According to our results, the tumor volume was more sensitive in detecting the changes of tumor size compared to the longest diameter. The results of this study indicated that the measurement of tumor volume could better reflect therapeutic response of DC vaccination in PDAC than that of the longest diameter. Zhang et al [38] showed that quantitative ADC measurement reflected pancreatic tumor microstructure in a rat model. In this study, the value of ADC measurement in monitoring therapeutic response of DC vaccination in PDAC was investigated. The results showed that the reduction of tumor ADC in the treatment group was less than that in the control group. Moreover, the change in tumor ADC was correlated negatively with the change in tumor volume after five weeks of treatment; however, there was no significant correlation between the change in tumor ADC and the change in the longest diameter. These results indicated that ADC measurement could provide a valuable imaging biomarker of therapeutic response to DC vaccination in pancreatic cancer.

The current study had several limitations. First, the sample size of mice pancreatic tumor model was small, and we will enlarge the sample size in the future. Second, MRI examinations were performed at only three time points after initiation of treatment; further investigations will include more time points as well as the over survival rate. Third, we also found that programmed cell death receptor ligand 1 (PD-
L1) was highly expressed in the murine pancreatic tumor model, which was considered to be involved in the mechanism of downregulating the antitumor response [39]. However, we did not explore it in depth in this study. DC vaccination combining with immune checkpoint inhibitors, such as PD-1/PD-L1-inhibitors, is the future direction of our work.

In conclusion, this study demonstrated the efficacy of DC vaccination administered via IP injection for immunotherapy in murine PDAC model, and the feasibility of ADC measurement as an imaging biomarker for assessment of therapeutic responses in immunotherapy.

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Disclosure of conflict of interest

None.

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