A unique MUC1-2-VNTR DNA vaccine suppresses tumor growth and prolongs survival in a murine multiple myeloma model

YUHUI WENG1,3, LIANG SHAO2, HONGMEI OUYANG1,5, YUEBO LIU1, JIN YAO1, HONG YANG1, YUNJIAO LUO1,4, HAO WANG1, ZHENYU ZHAO1, HONG MOU1, ZEPING ZHOU1 and YOU ZHANG1

1Department of Hematology, Second Affiliated Hospital of Kunming Medical University, Kunming 650101; 2Department of Medicine, The University of Hong Kong, Queen Mary Hospital, Hong Kong; 3Department of Hematology, Fuqing Hospital, Fuqing 350300; 4Children's Hospital of Kunming, Kunming 650000; 5First People's Hospital of Yunnan, Kunming 650000, P.R. China

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Abstract. Multiple myeloma (MM) is a clonal B-cell malignancy characterized by the aberrant proliferation of malignant plasma cells in the bone marrow. MM is still an incurable malignancy. In this regard, novel treatments are urgently required. MUC1 (mucin 1), a type I transmembrane protein, is overexpressed and aberrantly glycosylated in many carcinomas particularly in MM resulting in an antigenically distinct molecule and may be a potential target for specific immunotherapy. In this study, we first designed a unique DNA vaccine, termed MUC1-2-VNTR (various number tandem repeats) to investigate whether the vaccine could specifically suppress tumor growth in a murine multiple myeloma model. Our results showed that the constructed DNA vaccine pcDNA3.1-VNTR elicited both humoral and cellular tumor-specific immune responses in the MM mouse model leading to delay in tumor growth and prolonged survival of the mice. Consequently, our study indicates that this DNA vaccine shows promise to be used as a novel strategy for the treatment of MM.

Introduction

Multiple myeloma (MM) is characterized by clonal proliferation of malignant plasma cells. The median survival of MM patients is approximately 3 years (1). Despite recent improvements in the treatment of MM, present therapies still cannot cure the majority of patients with MM and they eventually relapse, mainly due to MRD (minimal residual disease). Many efforts have been made to find new treatments for this disease. Tumor vaccine has been one of the developments recently. Mucin 1 (MUC1), a kind of tumor-associated antigen (TAA) and a member of mucin family, is a highly glycosylated type I transmembrane protein normally presented on the luminal surface of secretory glands (2) as well as normal hematopoietic cell lines (3). The extracellular domain of MUC1 primarily consists of 25-125 variable number of tandem repeats (VNTR) of 20 amino acids: VTSAPDTRPAPGSTAPPAHG (4). However, some tumors such as breast, prostate, lung, colon, ovary, pancreas, MM, AML (acute myeloid leukemia), expressed an aberrant form of MUC1 (5-16). The polarized expression of MUC1 is lost and the normally glycosylated protein is overexpressed in hypoglycosylated forms (17), resulting in exposure of cryptic epitopes that have the potential to induce humoral and cellular immunity (18,19). It has also been reported that MUC1 specific immune response could be successfully induced both in mice and human (10-13). Therefore, MUC1 should be an attractive and applicable target strategy to treat MM.

Here, we firstly constructed recombinant vectors encoding MUC1-2-VNTR to investigate the effect of this vaccination on the immune system and its antitumor effects in MM mouse model.

Materials and methods

Construction of eukaryotic expression vector of MUC1. MUC1-2-VNTR coding gene was used as a research gene, a KOZAK sequence was engineered before the gene HindIII and XbaI restriction sites. The full length of the gene was 140 bp, synthesized by Takara, which subcloned into plasmid PUC18, and transformed into E. coli JM109 competent cells. pcDNA3.1/Myc-hisB vector was purchased from Invitrogen, containing multiple cloning site (MCS), BGH reverse priming site, myc epitope, C-terminal polyhistidine tag, SV40 early promoter and origin, polyadenylation signal and ampicillin resistance gene. PUC18/MUC1-2-VNTR plasmid from E. coli JM109 was carried out following the manufacturer's instructions. Then the PUC18/MUC1-2-VNTR plasmid and the pcDNA3.1/Myc-hisB vector were separately digested with XbaI and HindIII (Takara Biotechnology Co., Ltd., Dalian, China).
China), and ligated by T4 DNA ligase (Takara) to construct a plasmid termed pcDNA3.1-2-VNTR/myc-hisB. After that, the ligation mixture was transformed into E. coli DH5α competent cells, culturing on an LB Petri dish with 100 µg/ml ampicillin at 37°C overnight. Ampicillin-resistant transformants were selected to bacterial colony. After the expected bands were observed by electrophoresis gel, the plamid isolated from positive colony was sequenced by Takara.

**COS-7 cell transfection.** COS-7 cells were routinely cultured in DMEM (Gibco) with 10% fetal bovine serum (FBS) (HyClone Laboratories, Logan, UT). The day before transfection, 5x10⁵ cells/ml was plated in each well of a 6-well plate without antibiotics until 90-95% confluent at the time of transfection. The mixture of 5 µg plasmid GFP, 2.5 mol/l CaCl₂, 2X HBS-HEPES, 5 µg plasmid pcDNA3.1-2-VNTR, or empty vector was diluted in 250 µl of serum-free medium (SFM) and incubated for 20 min at room temperature with a mixture of 10 µl Lipofectamine 2000 and 250 µl SFM and the resultant complex mixture was added to a monolayer of pre-confluent cells seeded in a 6-well plate. The cells were incubated at 37°C and in 5% CO₂ for 6 h, at which time the transfected medium was replaced with fresh and complete one. The cells transfected with vector pcDNA3.1 (+) were generated as a negative control.

**COS-7 cell transfection assay by western blot analysis.** Forty hours after cell transfection, the supernatant of medium for detection was collected and 1x10⁷ cells were prepared in 0.5 ml lysis buffer (10 mmol/l Tris-Cl, 5 mmol/l EDTA, 50 mmol/l NaCl, 50 mmol/l Na₂PO₄, 50 mmol/l NaF, 0.1 mmol/l Na₂VO₃, 1 mmol/l PMSF, 5 mg/ml aprotinin, 0.1% Triton X-100, pH 7.6). The cells were incubated on ice for 15 min and then centrifuged at 14,000 rpm at 4°C for 20 min. The supernatant was then collected followed by denaturing at 100°C for 5 min. For western blot analysis, the samples were loaded onto a 12% SDS-polyacrylamide gel and transferred to a nitrocellulose membrane. The membrane was incubated with a mouse anti-human MUC1 antibody, followed by a horseradish peroxidase-conjugated rat anti-mouse IgG antibody. Then the membrane was reacted with enhanced chemiluminescence detection reagent for 5 min and exposed to X-ray film at room temperature.

**The construction of multiple myeloma cell line 653-MUC1.** P3X63Ag.653 cells (mouse myeloma cell line) from CellBank of Chinese Academy of Sciences were grown in RPMI-1640 medium (HyClone) containing 10% fetal calf serum (FBS, Hyclone), 2 mM L-glutamine, 100 U/ml penicillin and 100 µg/ml streptomycin. The cells were incubated at 37°C in a humidified atmosphere of 5% CO₂ for 6 h, at which time the transfected medium was replaced with fresh and complete one. The cells transfected with vector pcDNA3.1 were used as the control. The mixture of 5 µg plasmid GFP, 2.5 mol/l CaCl₂, 2X HBS-HEPES, 5 µg plasmid pcDNA3.1-2-VNTR, or empty vector was diluted in 250 µl of serum-free medium (SFM) and incubated for 20 min at room temperature with a mixture of 10 µl Lipofectamine 2000 and 250 µl SFM and the resultant complex mixture was added to a monolayer of pre-confluent cells seeded in a 6-well plate. The cells were incubated at 37°C and in 5% CO₂ for 6 h, at which time the transfected medium was replaced with fresh and complete one. The cells transfected with vector pcDNA3.1 (+) were generated as a negative control.

**RNA isolation and RT-PCR.** Total RNA was prepared using TRIzol reagent (Invitrogen) as described by the manufacturer. Total RNA was reverse transcribed and amplified using the AccessQuick™ RT-PCR System (Promega) following the manufacturer's instructions. A pair of PCR primers, 5'-GGGG AGGATTTGGGAAGAC-3' (forward) and 5'-CCGAGATAGGG GTTGAGTTGT-T-3' (reverse), was designed according to the sequence of pcDNA3.1-2-VNTR, which amplified a 442 bp β-actin and 653 cells transfected without pcDNA3.1-2-VNTR was used as controls. The primers, β-actin, 5'-ATGCCATCCTGGTCGTCCCTGGACCTGGC-3' (forward), 5'-AGCATTGGCGGT GCACGATGGAGG-3' (reverse), amplified a 607 bp. The amplification protocol consisted of a 5-min denaturation at 95°C followed by 30 cycles of denaturation at 95°C for 50 sec, annealing at 54°C for 50 sec and 72°C for 50 sec and a final extension at 72°C for 5 min.

**Mice and immunization.** The BALB/c mice, female, 7-8 weeks of age, from the Shanghai Laboratory Animals Center, Science Academy of China, were bred in the Animal Lab center (Kunming Medical University, Yunnan). The facility was approved by the Association for Assessment and Accreditation of Laboratory Animal Care International, and all procedures were carried out in accordance with the Guidelines and Regulations for Use and Care of Animals. The mice were divided randomly into 2 groups (n=10). Plasmid DNA (100 µl/100 µg) was injected into the spleen of the mice on Day 0. A booster immunization was given on the fifth week by injecting into each hind quadriceps muscle. The mice treated with the empty plasmid pcDNA3.1 were used as the control. Mice were bled by retroorbital venous orbita at 1, 2, 4, 6 and 8 weeks after the spleen immunization. Sera were collected after centrifugation at 13,000 rpm at 4°C for 20 min, then cryopreserved at -80°C.

**MUC1 antibodies detected by ELISA.** Polyvinyl chloride microtiter plates (Costar, Inc., Corning, NY) coated with 500 ng/well MUC1-VNTR was incubated at 4°C, overnight. The plates were blocked (0.5% porcine gelatin, 4% BSA in PBS), followed by incubation with diluted mouse sera in serum samples (1:200). Anti-MUC1 antibodies were detected with horseradish peroxidase (HRP)-labeled anti-mouse IgG (H+L) (Pierce Biotechnology, Inc., Rockford, IL) and developed with 3-ethylbenz-thiazoline-6-sulfonic acid (ABTS) solution. Absorbance was read at 405 nm. Values (1/titer) were scored as positive for the presence of MUC1 antibodies if the OD readings were at least three-times over PBS control wells.

**Proliferative assay by CCK-8 method.** Four weeks after the second immunization, mice were sacrificed, and single-cell suspension of splenocytes was prepared. They were plated at 500,000/well in 96-well U-bottom plates (#3799: Costar Co., Cambridge, MA) and incubated with MUC1-VNTR (10 µg/ml) at 37°C in 5% CO₂ for 3 days. Sixteen hours before termination of the culturing, 20 µl of CellTiter 96® AQueous One Solution Reagent (Promega) was added into each well. Stimulation index (SI) was calculated to detect spleen lymphocyte proliferation activity. SI, cpm of the experimental pair/cpm of negative control. All experiments were performed in
triple samples. ConA and RPMI-1640 were used as positive and negative control, respectively.

Cytotoxicity assay by LDH releasing method. Cytotoxicity of CTL against the target cells was detected by a non-radioactive lactate dehydrogenase (LDH)-releasing assay (CytoTox 96, Promega, USA). Single-cell suspension of splenocytes were collected as effector cells, 653 cells stably transfected with MUC1-VNTR as target cells. The effector cells were co-cultured with target cells at different effector cell/target cell ratios (10:1, 40:1 and 80:1). Then, 1x10^5 target cells per well were mixed with a varying number of effector cells in 96-well round-bottom plates with total volume reaching 200 µl and incubated for 4 h at 37˚C. At the end of the assay, supernatants (50 µl/well) were harvested and transferred to a fresh plate to test LDH releasing rate according to the manufacturer. The killing activity of CTL cells against tumor cells was calculated with the following formula: killing activity (%) = (experimental - effector spontaneous - target spontaneous)/(target maximum - target spontaneous) x 100%. Spontaneous and maximal releases were determined in the presence of either medium or 1% Triton X-100, respectively.

Animal studies. The BALB/c mice, female, 7-8 weeks old, were purchased from the Shanghai Laboratory Animals Center, Science Academia of China. The mice were randomly divided into 2 groups (n=10, respectively). The mice were injected with plasmid DNA (100 µl/100 µg) as mentioned above. The empty plasmid pcDNA3.1 was used as control. Seven days after the second immunization, each mouse was inoculated with 1x10⁶ 653-MUC1 cells at the interterm of the left anterior leg armpit. The tumor development in individual mouse was monitored every 2-3 days and the tumor volume (in mm³) was calculated by the following formula: 0.5 x length (mm) x width (mm)². The survival time after the tumor challenge was recorded for 60 days until the death of the mouse.

Statistical analysis. Data are presented as mean ± SEM. Statistical differences between two groups were evaluated by the unpaired Student's t-test. The survival time was calculated by the Kaplan-Meier method. The survival rates were compared by the log-rank test (SPSS 13.0). P-values <0.05 were considered significant.

Results

Identification of the MUC1-2-VNTR and eukaryotic expressing vector pcDNA3.1/myc-hisB. The MUC1-2-VNTR and eukaryotic expressing vector pcDNA3.1/myc-hisB were cut by the double restriction enzyme Xbal and HindIII. Then the enzyme products were electrophoresed using agar gels. The objective fragment 140 bp (Fig. 1) and fragment 5417 bp were obtained (Fig. 2).

Identification of the recombinant plasmids. Four transformants from LB Petri dish with 100 µg/ml ampicillin using T7 promoter/BGHrev primer for PCR were selected, then the PCR products were electrophoresed using the agar gels. The objective fragment 140 bp (Fig. 1) and fragment 5417 bp were obtained (Fig. 2).

Identification of the recombinant plasmids. Four transformants from LB Petri dish with 100 µg/ml ampicillin using T7 promoter/BGHrev primer for PCR were selected, then the PCR products were electrophoresed using the agar gels. The objective fragment 140 bp (Fig. 1) and fragment 5417 bp were obtained (Fig. 2).

The sequencing includes whole reading frame of pcDNA3.1/myc-his B and the 2-VNTR inserted DNA sequence (underlined). Together, the construction of recombinant plasmid pcDNA3.1-2-VNTR was successful. 5’-CCTACTTTGG CAGTACATCTACGTTATTAGTCATCGCTATTACCATGG GATGCGGTGTGGCAGTACATCAATGGGCGTGGAGC
Figure 4. DNA sequencing of recombinant plasmid pcDNA3.1-2 VNTR/myc-his B.

Figure 5. Co-transferred GFP observed by fluorescence microscopy. (A) Untransfected COS-7 cells. (B) COS-7 cells transfected with vacant plasmid pcDNA3.1/myc-his B+GFP. (C) COS-7 cells transfected with recombinant plasmid pcDNA3.1-2 VNTR/myc-his B+GFP.

GGTGTGACTCACGAGATTTCCAAGTCTCCCCCATTTG
ACGTCAATGGGAGTTTGTTTTGGCCAAAATCAACGG
GACCTTTCAAAATGTCGCAACTCGCCCCATTTGACG
CAGGAGCCGAGGACGCCGTACGTAGGGACTTATATA
TAAGCAGGCTCTCTTGACTAAGAAGAAGCCACTG
CTTACTGCTTTATCGAAATTAATACGACTCAGTTAGTT

GGAGACCCAAGCTGGCTAGTTAAGCTTGCACCACAT
GGTCACCTCGGCCCCGGACACCCAGGGGCCCCGG
GCTCCACGCCCCCCCCACGCTGTCACCTCG
GCCCCGACCCAGGGCCGCCCCGGGCTCCACCACGC
CCCCCCACCCACCGTCTAGAGGGCCCGCGTTCG
AACAAAAACTCATC-3'.
Cotransfected COS-7 cells with pcDNA3.1+GFP. COS-7 cells were co-cultured with vacant plasmid pcDNA3.1/myc-hisB or recombinant plasmid pcDNA3.1/myc-hisB+GFP for 2 days, followed by the detection of fluorescence microscopy. The results showed that recombinant pcDNA3.1/myc-his B and pcDNA3.1-2VNTR-hisB were successfully transfected into COS-7 cells (Fig. 5).

Detection of expression of pcDNA-2-VNTR protein by western blot analysis. The result of western blot analysis showed MUC1 protein was expressed in the cells transfected with pcDNA plasmid. After probing with antibodies against VNTR, the molecular weight of objective band was 4 kDa (Fig. 6).

Monoclonal cell line verification by RT-PCR. P3X63Ag.653 transfected by pcDNA3.1-2-VNTR showed much stronger expression of VNTR. Band 2 was about 121 bp as expected, whereas, we detected no expression of VNTR in the control group (P3X63Ag.653 transfected by vector pcDNA3.1(+)) (Fig. 7).

Elevated level of specific antibodies in the sera of vaccinated mice. Mice were bled by retro-orbital venous in week 1, 2, 4, 6 and 8 after the spleen immunization. Specific anti-MUC1 antibody can be detected in the first week after the spleen vaccination, and the antibody level reached a peak in the eighth week. Compared with the empty vector group, significantly higher titres of anti-MUC1 antibodies were detected in the sera of pcDNA3.1-VNTR-vaccinated mice (P<0.01) (Fig. 8). The result indicated that pcDNA3.1-VNTR can efficiently induce a specific humoral immune response via induction of specific antibodies.

Induction of specific antibody response. Mice were immunized with pcDNA3.1-VNTR or pcDNA3.1 and sera were harvested in week 1, 2, 4, 6 and 8 after the spleen immunization. The MUC1-specific IgG antibodies in sera of different groups were determined at a 1:200 dilution by ELISA. As shown in Fig. 8, relatively higher titres of anti-MUC1 antibodies were detected in the serum of pcDNA3.1-VNTR vaccinated mice compared to that in the sera of the empty vector vaccinated groups in week 1, 2, 4, 6 and 8 (P<0.01).

Effects of pcDNA3.1-MUC1 on the proliferation of lymphocytes. To test the effect of pcDNA3.1-MUC1 vaccine on the proliferation of lymphocytes, lymphocytes were isolated from the spleens of immunized mice four weeks after the second immunization. We chose to optimize and qualify the colorimetric cell proliferation assay using MTS (aqueous soluble form of the tetrazolium salt based assay) with the Cell Titer96® Aqueous One assay kit from Promega Corp. The results showed that proliferation activity of lymphocytes from mice...
vaccinated with recombinant plasmid group was significantly higher than that of the empty vector treated mice (14.94±2.74 vs. 7.61±1.59, P<0.01) (Fig. 9).

Splenocytes from immunized mice were isolated four weeks after the second immunization. CTLs were from mice immunized with pcDNA3.1-VNTR or pcDNA3.1. 653-MUC1 cells were used as targets. The CTLs induced by the pcDNA3.1-VNTR could specifically kill the 653-MUC1 cells.


To determine whether immunization with pcDNA3.1-VNTR could induce strong CTL response, we detected the cytotoxicity response by non-radioactive LDH-releasing assay. Effector cells were co-cultured with target cells in different ratios 80:1, 40:1 and 10:1. There was significant difference in the ratios of both 80:1 and 40:1, with the killing rate of 26.23±5.02 and 16.88±1.98, respectively, compared with 14.36±4.13 and 10.67±2.11 in the empty vector control group. However, there was no significant difference in the ratio of 10:1 (E:T) (10.30±1.65 vs. 8.98±2.16, P>0.05) between the two groups (Fig. 10). These results confirmed that pcDNA3.1-VNTR are capable of inducing MUC1-specific CTL effect in vitro.

Effect of pcDNA3.1-VNTR vaccine on multiple myeloma in vivo.

To determine whether pcDNA3.1-VNTR vaccine is effective in suppressing multiple myeloma in vivo, cohorts of mice were inoculated with pcDNA3.1-VNTR twice before challenging with 1x10⁶ pcDNA3.1-VNTR or pcDNA3.1, respectively. As shown in Fig. 11, tumor of the mice immunized with pcDNA3.1-VNTR grew slower compared with mice treated with pcDNA3.1. Furthermore, immunized mice were tumor-challenged and monitored for tumor growth for 60 days. As shown in the Fig. 12, the mice immunized with pcDNA3.1-VNTR survived longer compared with mice treated with pcDNA3.1. Moreover, immunized mice were tumor-challenged and monitored for tumor growth for 60 days. As shown in the Fig. 12, the mice immunized with pcDNA3.1-VNTR survived longer than mice in control group (P<0.01 by log-rank test). Mean survival time of pcDNA3.1-VNTR immunized mice was 41 days, higher than the 28 days of pcDNA3.1 immunized mice, demonstrating that the pcDNA3.1-VNTR vaccine could efficiently induce suppressive effect on tumor growth.

Discussion

MM is still considered as an incurable malignancy. Many efforts have been made to find a new way to treat the disease. In general, the growth of malignant cells in MM is relatively lower than that of other hematologic malignant cells such as leukemic cells, which may result in the low sensitivity to chemotherapy but, at the same time, may lead to be more suitable for the use of immunotherapy especially after CR (complete remission) to eliminate the MRD (minimal residual disease).

MUC1, a tumor-associated antigen (TAA), is composed of a large ectodomain that is heavily glycosylated through GalNAc O-linkages, a single-pass transmembrane region, and an intracellular cytoplasmic tail (CT) (20-22). The extracellular domain of MUC1 primarily consists of 25-125 various number tandem repeats (VNTR) of 20 amino acids and varies among individuals or directly derived tissues (4), which is...
the most immunogenic part of MUC1 and many T and B cell antigen recognition sites generated to date almost exclusively recognize epitopes in that immunogenic region (23-25). Thus, MUC1-VNTR sequences may be expected to induce host immune specific CTL responses and as a possible target treatment approach for MM.

In the mid-80s MUC1 was considered to be a tumor marker when it was detected in the serum of most breast cancer patients (26). The presence of MUC1 was demonstrated in subgroups of human lymphomas, including plasma cell myeloma in the early 1980s (27,28).

Non-MHC-restricted cytotoxic T cell activity has been described on multiple myeloma cells by Takahashi et al (15). In a published study, 44% of myeloma patients exhibited elevated frequencies of MUC1-specific CD8 T cells in peripheral blood as well as bone marrow (29).

Recently, MUC1 immunization studies mainly focus on peptides vaccine (30-34), virus vaccine (35-37), dendritic cells (DC) and DC/tumor cell fusions vaccine (38-42), RNA vaccine (38) and DNA vaccines (43-45). MUC1-DNA immunization has been actively investigated since the demonstration of tumor protective immune response after intramuscular injection of naked MUC1-DNA (46).

In our previous studies, we have confirmed that MUC1 was highly expressed in MM patients by immunohistochemistry, ELISA and RT-PCR (47-49). On the other hand, the low tumor burden in the CR patients with MM and the low growth rate of MM cells may benefit the most from the DNA vaccine for treatment of MM.

Based on the evidence, we supposed that the approach of using MUC1 as a target to further eradicate MRD of MM may be reasonable. The eukaryotic expression vector pcDNA3.1/myc-hisB used in this study is typical and efficient vector which can be expressed in the majority of mammals. DNA vaccines have been widely investigated in a variety of disease models and clinical trials including autoimmune disease, infectious disease as well as carcinoma. However, we realized that DNA vaccines may generate weak immune response without any booster or modification. So, in order to enhance the immunogenicity of MUC1, we designed a unique 2-fold VNTR sequence (2-VNTR) as a target antigen to construct the immunogenicity of MUC1, we designed a unique 2-fold VNTR sequence (2-VNTR) as a target antigen to construct the immunogenicity of MUC1.

To investigate the humoral and cellular immune responses induced by MUC1-VNTR vaccine in multiple myeloma, we used pcDNA3.1-VNTR immunized female BALB/C mice to detect both of them. Detection of specific antibodies by ELISA confirmed that humoral immune responses were elicited by pcDNA3.1-VNTR and cytotoxicity assay showed that splenocytes from the pcDNA3.1-VNTR immunized mice could specifically kill the 653-MUC1 tumor cells.

In the suppression experiment of the model, we found that the median survival of vaccinated mice was significantly longer than that of the control group. In addition, the tumor growth of the mice was markedly suppressed. These results showed that MUC1-2-VNTR vaccine may effectively inhibit or kill MM tumor cells in vivo and potentially to eliminate the MRD of MM, at the end, to cure the disease. The mechanism involved in this strong effect may be mainly due to the unique design of the 2-fold VNTR sequence which greatly enhance the specific CTL effect and lymphocyte proliferation as well as boosting specific anti-MUC-1 antibody produced.

However, the vaccine may not be strong enough to abrogate all the MM tumor cells, as all the mice that received pcDNA3.1-2-VNTR vaccine still died within 60 days. Future work might be focused on further enhancing the effectiveness of the vaccine by, for example, designing three-fold VNTR sequences, or to transfect siRNA (small interfering RNA) into DC (dendritic cell) pulsed with MUC1-2-VNTR to silence the expression of SOCS1 (suppressor of cytokine signaling).

Our study, for the first time, uses pcDNA3.1-2-VNTR vaccine to suppress the growth of MM cells in mouse model demonstrating that our hypothesis above may be correct. It may provide a potential therapeutic approach to treat MM patients in the future.

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