**Immune responses in mice vaccinated with a DNA vaccine expressing a new elastase from *Trichinella spiralis***

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**Abstract:** The elastase, which belongs to the serine protease family, hydrolyses various proteins and may be involved in the parasite invasion. In this study, complete sequence of elastase-1 (TsE) the nematode *Trichinella spiralis* (Owen, 1835) was cloned into the plasmid pcDNA3.1 as TsE DNA vaccine. After intramuscular vaccination, serum anti-*Trichinella* antibodies (IgG and subclass IgG1/ IgG2a, and IgA), total and specific intestinal mucosal slgA in mice vaccinated with pcDNA3.1/TsE were measured by ELISA. The results showed that vaccination with pcDNA3.1/TsE induced a systemic humoral immune response (high levels of serum IgG and subclass IgG1/IgG2a and IgA) and local intestinal mucosal immune responses (high levels of TsE-specific slgA). Vaccination of mice with TsE DNA vaccine also triggered a systemic and local concomitant Th1/Th2 response, as demonstrated by significant elevation of Th1 (IFN-γ and IL-2) / Th2 (IL-4 and IL-10) cytokine levels after the spleen, mesenteric lymph node and Peyer’s patch cells from vaccinated mice were stimulated with recombinant TsE (rTsE). The vaccination of mice with pcDNA3.1/TsE displayed a 17% reduction of intestinal adult worms and a 39% reduction of muscle larvae. Our results indicated that TsE DNA vaccine elicited a systemic concomitant Th1/Th2 response and an enteral local slgA response, and produced a partial protection against infection with *T. spiralis*. The TsE may be regarded as a potential candidate vaccine target against *Trichinella* infection. The oral polyclonal vaccines should be developed to improve the protective efficacy of anti-*Trichinella* vaccines.

**Keywords:** *Trichinella*, trichinellosis, vaccination, protective immunity

**Species of *Trichinella* Railliet, 1895 are intestinal and tissue-dwelling nematodes which can infect over 150 species of mammals, birds and reptiles over the world (Feidas et al. 2014). It has known that pigs and other domestic animals (horses, dogs, cats etc.) are the main natural hosts of *Trichinella spiralis* (Owen, 1835). Human with infection *T. spiralis* occurs when raw or uncooked meat containing encapsulated infective larvae is consumed. *Trichinella spiralis* is a severe public health threat and a tremendous hazard on animal food safety (Cui and Wang 2011, Cui et al. 2013a, Jiang et al. 2016, Rostami et al. 2017).

In the mainland China, the infected domestic pork is still the primary infectious source of human trichinellosis (Cui et al. 2011). It is difficult to prevent and control this zoonotic parasitosis because of its wide animal reservoirs, complicated life cycle and different state-specific antigens (Poziol 2007, Liu et al. 2015c). Until now, there are no effective preventive vaccines available for the food animal production (Zhang et al. 2018a). Hence, it is necessary to exploit anti-*Trichinella* vaccines to eliminate *Trichinella* infection in domestic pigs and block the transmission from pigs to humans (Li et al. 2015, 2018).

In recent years, some significant immune protection has been reported by using DNA vaccines to prevent parasitic infections (Xu et al. 2017, Qi et al. 2018a, Wang et al. 2018). Furthermore, the DNA vaccine is easy to be constructed, prepared and administered. In our previous studies, a new elastase gene of *T. spiralis* (TsE, GenBank: XP_003377838.1), which belongs to the serine protease family, was obtained from the *T. spiralis* draft genome (Mitreva et al. 2011). It was cloned, expressed and purified in our laboratory. Bioinformatic analysis showed that complete TsE cDNA sequence was 1,350 bp encoding 449 amino acid with a weight of 47.3 kDa.

The TsE protein expression level at *T. spiralis* muscle larvae (ML) and intestinal infective larvae (ILI) stages was higher than those of other worm stages (Hu et al. 2020). The recombinant TsE (rTsE) facilitated the penetration of larvae into the *in vitro* intestinal epithelium cell (IEC) monolayer, whereas anti-rTsE antibodies and siRNA inhibited larval penetration of IEC (Zhang et al. 2020). The results suggested that the novel TsE might be a promising target molecule of vaccines against *T. spiralis* infection. In the present study, the complete TsE cDNA sequence was cloned into the eukaryotic expression vector pcDNA3.1 and the pcDNA3.1/TsE was used to vaccinate BALB/c mice by intramuscular injection. The systematic and intestinal mucosal immune responses, and immune protection induced by vaccination using TsE DNA vaccine were ascertained.
MATERIALS AND METHODS

Worms and hosts

Trichinella spiralis isolate (ISS534) used in this study was obtained from a domestic pig in central China and passed in our laboratory (Wang et al. 2012). Female BALB/c mice (four weeks old) were purchased from the Zhengzhou University Experimental Animal Center (Zhengzhou, China), and fed in individual ventilated cages (IVC, Suzhou, China) (Cui et al. 2019). All animal experiment at procedures were authorised by the Life Science Ethics Committee, Zhengzhou University (No. SCXK 2017–0001).

Collection of various life-cycle stages

Muscle larvae (ML) were collected from infected mouse skeletal muscles at 42 days post-infection (dpi) by artificial digestion as previously reported (Jiang et al. 2012). The intestinal infective L1 larvae (ILL) were recovered from the intestine of infected mouse at six hours post-infection (hpi) (Liu et al. 2015c). Adult worms (AW) were isolated from the intestine at 3 dpi (Yang et al. 2015a). Female adults were cultured in RPMI-1640 at 37°C in 5% CO₂ for 72 h, and then newborn larvae (NBL) were harvested as described (Zhang et al. 2013, Wu et al. 2016).

Preparation of rTsE and anti-rTsE serum

Complete TsE gene was cloned and recombinant plasmid pQE-80L-TsE was transformed into Escherichia coli BL21 (Novagen) and induced by 1 mM IPTG at 37°C for six hours (Liu et al. 2017). Recombinant TsE protein (rTsE) was purified with the aid of Ni-NTA-Sefinose resin (Sangon Biotech, Shanghai, China) in our laboratory (Hu et al. 2020). Fifteen mice were subcutaneously injected by 20 μg rTsE mixed 1 : 1(v/v) with complete Freund’s adjuvant. Two immunisation boosts were given as the same rTsE dosage with incomplete Freund’s adjuvant at a two-weeks interval (Long et al. 2014, Sun et al. 2018b). At two weeks following the final immunisation, tail blood was taken and anti-rTsE sera were isolated (Cui et al. 2013b).

Plasmid construction and sequence analysis of recombinant expression plasmids

The total RNA from T. spiralis ML was extracted using Trizol (Invitrogen, USA), and reversely transcribed to the first strand cDNA (Hu et al. 2000). Complete TsE cDNA sequence (GenBank: XP_003377838.1) was obtained by PCR amplification with the following primers carrying Bam HI and Xhol(bold) (5’–CGGATCCGCCACCATGATCCTTTTCAAGTGC-3’, 5’–CGGCTCGA GTTAGCGAAAAAGGTG AAT GA-3’).

The DNA encoding complete TsE was cloned into pcDNA3.1 as the TsE DNA vaccine (pcDNA3.1/TsE). In order to permit efficient initiation of translation, an artificial Kozak sequence (GCCACC) was incorporated into the sense primers (Liu et al. 2015a). The clones with expected size inserts were selected by restriction digestion and subjected to DNA sequencing. DNA and predicted amino acid sequences were analysed by Lasergene 7.1 software (DNASTAR Inc., Madison, Wisconsin, USA). The plasmid DNA was isolated using alkaline cleavage method (Tang et al. 2012).

RT-PCR and immunofluorescence test (IFT) for detecting TsE mRNA and protein expression in vitro

Baby hamster kidney cell line 21 (BHK-21) were cultivated in culture plates (Nunc) with DMEM medium containing 100 U/ml penicillin, 100 μg/ml streptomycin and 10% fetal bovine serum at 37°C in 5% CO₂. When the cells were grown to 90% confluency, the cells were transfected by pcDNA3.1/TsE with a Lipofectamine 2000 (Invitrogen, USA) at 37°C for 48 h. Total RNA was isolated from transfected cell and TsE mRNA expression was measured by RT-PCR with TsE-specific primers as mentioned above (Liu et al. 2015a). TsE protein expression in transfected BHK-21 cells was examined by IFT (Liu et al. 2018). In brief, BHK-21 were culture up to cell monolayer and fixed using 4% formaldehyde for 20 min. After washes with PBS, the cells were permeabilised using 0.1% Triton-X-100 for 15 min, blocked using 5% normal goat serum at 37°C for one hour, and incubated with anti-TsE serum (1 : 10) at 4°C overnight. After washes with PBS, the cells were incubated with FITC-anti-mouse IgG conjugate (1 : 100; Santa Cruz, USA) at 37°C for one hour (Song et al. 2018a). The cells were re-dyed using 4’, 6-diamidino-2-phenylinodole (DAPI) for seven min. The cells were examined on fluorescence microscopy (Olympus, Tokyo, Japan) (Long et al. 2015, Xu et al. 2018).

Vaccination regime and sample collection

One hundred and eighty mice were divided into three groups (60 animals per group). Pre-immune serum samples (normal serum) were obtained via tail bleeding before the first vaccination (Liu et al. 2015a). Three days before vaccination, all mice were first injected in left and right quadriceps with 25μl of 5% bupivacaine hydrochloride. All mice were vaccinated through bilateral intramuscular injection into the quadriceps three times with different time interval. Each mouse of vaccine group was injected with 100 μg pcDNA3.1/TsE in 100 μl sterile PBS at each vaccination. As negative and blank control, mice were injected with 100 μg empty plasmid pcDNA3.1 or 100 μl PBS alone. At weeks 0, 4, 6 and 8 post-vaccination, ten mice of each group were sacrificed. Serum, spleen, mesenteric lymph nodes (MLNs), Peyer’s patches (PPs), 1 cm long duodenum segments and enteral flushing fluid were recovered to assess systematic and intestinal local immune response (Sun et al. 2019b).

RT-PCR and IFT for detecting TsE mRNA and protein expression in vivo

TsE mRNA expression in spleen and MLNs of vaccinated mice was determined using RT-PCR at two weeks after the final vaccination. Total RNAs were extracted from spleens and MLNs with Trizol regents (Invitrogen, Carlsbad, USA). Mouse β-actin was also amplified as an internal control (Liu et al. 2014). PCR products were analysed on 1% agarose gels (Wang et al. 2013). To investigate the TsE protein expression, 3 μm tissue sections of spleen and MLNs of vaccinated mice were prepared. The sections were fixed with acetone and blocked with 5% normal goat serum at 37°C for one hour. Following washes with PBS, the sections were probed with anti-rTsE serum (1 : 10 in PBS) at 4°C overnight. After washes with PBS again, they were stained using FITC-conjugated anti-mouse IgG (1 : 100), and observed in fluorescence microscope (Olympus, Japan) (Song et al. 2018b, Sun et al. 2018a).
Determination of anti-TsE antibodies

Anti-rTsE antibodies (total IgG, IgG1, IgG2a and IgA) in serum samples of vaccinated mice were measured using indirect ELISA (Cui et al. 2015, Sun et al. 2015). In brief, the ELISA plate was coated with 1 μg/ml rTsE for total IgG and 2 μg/ml rTsE for IgG1, IgG2a and IgA overnight at 4 °C. Blockage was performed with 5% nonfat milk at 37 °C for one hour. After washes with PBST, the plates were incubated at 37 °C for 2 h with various dilute serum samples (1:100 for detecting IgG, 1:50 for detecting IgG1, IgG2a and IgA), followed by incubation with HRP-conjugated anti-mouse IgG (1:10,000), IgG1, IgG2a and IgA (1:5000; Sigma-Aldrich Co.) for one hour at 37 °C (Sun et al. 2019a). Colouration was performed by using OPD (Sigma-Aldrich Co.) plus 0.15% H2O2, and reaction was finished by the addition of 2 M H2SO4. The absorbance at 492 nm was measured by a microplate reader (Tecan, Schweiz, Switzerland) (Li et al. 2015, Liu et al. 2015d).

Recognition of natural TsE at various stages of Trichinella spiralis by IFT

The recognition of natural TsE at diverse life cycle stages of T. spiralis was investigated using IFT with 3 μm thick cross-sections of worms as reported (Pompa-Mera et al. 2014, Qi et al. 2018b). The blockage was performed using 5% normal goat serum at 37 °C for one hour. After washing with PBS, the sections were incubated with various immune serum (1:10 dilution) at 37 °C for one hour, stained using FITC-labelled anti-mouse IgG (1:100; Santa Cruz, USA), and observed under fluorescence microscope (Olympus, Japan) (Xu et al. 2020).

Assay of enteral mucosal secretory IgA (sIgA) response

To evaluate enteral local mucosal sIgA response on vaccination with pcDNA3.1-TsE, total and TsE-specific secretory sIgA in enteral fluid were determined (Liu et al. 2015b, Bermúdez-Cruz et al. 2016). In brief, a 20 cm long pieces of the small intestine was incised, and content was flushed by 1 ml of cold PBS containing 1% protease inhibitor (Sangon Biotech, China). The eluting fluid was recovered, centrifuged at 10000 g for 5 min, and supernatants were collected (Li et al. 2018). Total sIgA was measured by a sandwich ELISA (Abcam, Cambridge, UK), and TsE-specific sIgA was determined by a conventional ELISA with 2 μg/ml of rTsE as reported (Sun et al. 2015). Colouration with OPD and determination of absorbance at 492 nm were performed as described before (Liu et al. 2013, Ren et al. 2018).

Investigation of enteral goblet cells/mucins

One centimetre of the duodenum of vaccinated mouse was cut off at two weeks after the last vaccination. The duodenum was fixed using parafomaldehyde, embedded in paraffin, and cut into 3-μm thick sections (Long et al. 2015). To identify the generation of enteral mucus, sections were stained with alcian blue-periodic acid Schiff reagent at 37 °C for 15 min, and stained using hematoxylin. The goblet cells/mucins were examined under light microscopy, and goblet cells per intact intestinal villus were numbered.

Determination of cytokine response

To analyse cellular immune responses to pcDNA3.1-TsE vaccination, four cytokines (IL-2, IL-4, IL-10 and IFN-γ) from spleen, MLN and PP from vaccinated mice were measured at weeks 0, 4, 6 and 8 after vaccination. Spleen, MLNs and PPs were homogenised using complete RPMI-1640 medium (Gibco, Auckland, New Zealand), the pellets were obtained after being centrifuged at 300 g for 5 min, and the cells were harvested (Dea-Ayuela et al. 2006). These cells (2 × 106 cells/ml) were stimulated using 10 μg/ml of rTsE at 37 °C for 72 h, supernatant was collected and the cytokine concentration (pg/ml) was measured using sandwich ELISA (Cui et al. 2013b).

Challenge infection and evaluation of immune protection

To determine the immune protective efficacy generated from pcDNA3.1/TsE vaccination, all vaccinated mice were challenged orally with 300 T. spiralis ML two weeks after the third vaccination. Ten mice from each group were euthanised at 7 dpi, adult worms were collected from the intestines and counted (Zhang et al. 2016). The remaining ten mice from each group were sacrificed at 42 dpi, each carcass was weighed, digested and the ML were collected and numbered as reported (Wang et al. 2015, Yang et al. 2019). The immune protective efficacy was ascertained as parasite burden reduction of intestinal AWs and larvae per gram (LPG) of muscles from of pcDNA3.1-TsE vaccinated mice relative to that of PBS group (Xu et al. 2017).

Statistical analysis

Statistical analysis was performed with SPSS for Windows, version 20.0. The data were shown as mean ± standard deviation (SD). Difference among various groups was analysed by one-way ANOVA. P < 0.05 was regarded as a statistical significance.

RESULTS

Identification of recombinant plasmid pcDNA3.1-TsE

Recombinant plasmid pcDNA3.1-TsE was digested using BamHI/Xhol. Electrophoresis of digestion products revealed that recombinant plasmid with an insert of about 1,350 bp was successfully constructed. Sequence analysis demonstrated that the amplified TsE gene fragment consisting of 1,350 bp was correctly cloned into the pcDNA3.1 (Fig. 1), and had 99.81% identity to the TsE sequence in GenBank (XP_003377838.1).

The in vitro TsE mRNA and protein expression

The in vitro TsE mRNA expression in BHK-21 cells was determined using RT-PCR. The results showed that amplified TsE fragments were detected in the cells transfected by pcDNA3.1/TsE, but not in the empty pcDNA3.1-transfected cells. TsE protein expression in the BHK-21 cells was examined by IFT with anti-TsE serum. Positive green fluorescence in pcDNA3.1/TsE transfected cells was observed by using anti-TsE serum and infection serum, but not in pcDNA3.1 transfected control cells (Fig. 2).
The in vivo TsE mRNA and protein expression

Total RNAs were extracted from vaccinated mouse spleen and MLNs at two weeks after the final vaccination. TsE mRNA expression in the tissues was ascertained by RT-PCR. The results showed that TsE mRNA was expressed in spleen and MLNs from mice immunised with pcDNA3.1/TsE, but not from mice injected with only empty pcDNA3.1 or PBS alone (Fig. 3). The IFT results revealed that fluorescent staining was observed in spleen and MLN sections of TsE DNA vaccine group, but not in empty pcDNA3.1 or PBS group. Furthermore, when the immunised mouse spleen and MLN sections were incubated with pre-immune serum, no fluorescence staining was seen. The results demonstrated that TsE mRNA and protein were expressed in spleen and MLNs of pcDNA3.1/TsE immunised mice.

Systemic TsE-specific antibody response elicited by vaccination with pcDNA3.1/TsE

As shown in Fig. 4, specific anti-TsE antibodies (total IgG, IgG1 and IgG2a) of pcDNA3.1/TsE vaccinated
mice at 4, 6 and 8 weeks after vaccination were evidently higher than those of empty pcDNA3.1 and PBS groups \((P < 0.0001)\), suggesting that a mixed Th1/Th2 immune response was triggered by pcDNA3.1-TsE vaccination. Moreover, TsE-specific IgA response was also elicited in the vaccinated mice.

Recognition of natural TsE at various stages of *Trichinella spiralis* by IFT

The IFT results indicated that natural TsE on cross-sections of IIL, adult worms and muscle larvae was recognised by anti-TsE serum from pcDNA3.1/TsE vaccinated mice. Immunostaining was principally located at the cuticle, stichosome and around female embryos of this parasite (Fig. 5). However, no worm tissues were identified by using serum from mice injected with empty pcDNA3.1 or PBS alone.

Enteral local mucosal sIgA response

To investigate local enteral mucosal sIgA response to pcDNA3.1/TsE vaccination, total and TsE-specific sIgA were assayed by ELISA. The results showed that total sIgA level in pcDNA3.1-TsE vaccinated mice was remarkably higher than those of only pcDNA3.1 or PBS group (Fig. 6A) \((F_{4w} = 186.037, F_{6w} = 478.646, F_{8w} = 804.413, P < 0.0001)\). TsE-specific sIgA level in pcDNA3.1-TsE vaccinated mice was also significantly higher than those of empty pcDNA3.1 or PBS group (Fig. 6B) \((F_{4w} = 126.359, F_{6w} = 321.75, F_{8w} = 360.426, P < 0.0001)\). No TsE-specific sIgA response was observed in mice inoculated with empty pcDNA3.1 or PBS alone.
Enteral goblet cells/mucins

The duodenum was obtained at two weeks after the last vaccination. Duodenal tissue section was dyed with AB-PAS, a plentiful of acidic mucin was dyed bluish violet, and identified in enteral mucosa from pcDNA3.1-TsE vaccinated mice. A number of acidic mucin was also observed in empty pcDNA3.1 group. However, a little of mucin was detected in PBS group (Fig. 7). The amount of enteral goblet cells/mucins in pcDNA3.1-TsE vaccinated mice was obviously higher than that of empty pcDNA3.1 or PBS group (Figs. 7, 8) ($F = 65.572$, $P < 0.0001$).

Evaluation of cytokine response

The results of cytokine assay revealed that the level of Th1 (IFN-γ, IL-2)/Th2 (IL-4, IL-10) cytokines at 4, 6 and 8 weeks after pcDNA3.1/TsE vaccination was significantly higher than those of only pcDNA3.1 or PBS injection ($P < 0.0001$) (Fig. 9). The results indicated that pcDNA3.1-TsE vaccination elicited the concurrent Th1/Th2 responses in the light of TsE-specific IgG subclass and cytokine response. It was also suggested that intramuscular inoculation of pcDNA3.1-TsE induced both systemic (spleen) and intestinal local (MLNs and PPs) cellular immune response.

Immune protective efficacy of pcDNA3.1/TsE vaccination

Compared to PBS blank control mice, the mice vaccinated with pcDNA3.1/TsE exhibited a 17% intestinal AW reduction at 7 dpi (Fig. 10a) and a 39% ML reduction at 42 dpi (Fig. 10b) after challenge with 300 *Trichinella spiralis* ML ($F_{AW} = 50.302$, $P < 0.01$; $F_{ML} = 8.925$, $P < 0.0001$).
DISCUSSION

Serine protease is a major protease family and plays an important role in protection against helminthoses. The protease can degrade different host proteins and enteral mucin, participates in the larval intrusion, growth and molting (Nagano et al. 2003, Yang et al. 2015b). Elastases belong to a trypsin-like serine protease which has the ability to hydrolyse various tissue components, such as fibronectin, laminin, collagen and elastin. The elastases are involved in the parasite invasion and digestion, and might a crucial role act in worm invasion (El-Faham et al. 2017). In a previous study, the TsE was cloned and expressed in Escherichia coli, and the rTsE might be a novel target molecule for anti-

T. spiralis vaccines (Hu et al. 2020). Nevertheless, the proteins expressed in prokaryotic plasmids might differ from the natural active proteins on the basis of their structure and function (Liu et al. 2014).

The recombinant protein fold and post-translational modification (e.g., glycosylation) could be obtained in DNA vaccine with eukaryotic expression plasmid, and the surface-exposed recombinant antigenic epitopes would be also exhibited correctly like the native protein structures (Wang et al. 2016). The DNA vaccine is more attractive due to its ability to induce a broad and lasting immune response (Liu et al. 2015a). An effective DNA vaccine usually requires to be administered more than one boost vaccination. Moreover, the DNA vaccine is commonly used through intramuscular injection (Wang et al. 2006, Xu et al. 2017).

In the present work, the complete TsE sequence was cloned into pcDNA3.1, and TsE mRNA and protein expres-
sion was observed in pcDNA3.1/TsE transfected- BHK-21 cells in vitro, suggesting that the TsE DNA vaccine was successfully prepared. The results of RT-PCR and IFT demonstrated that the TsE mRNA and protein were also expressed in spleen and MLNs of pcDNA3.1/TsE immunised mice. Cellular uptake of DNA is relatively inefficient following intramuscular injection, and therefore expression of the encoded antigen is low. Strong promoters have been shown to increase expression levels and the immediate/early promoter from cytomegalovirus (CMV) is the most effective and frequently used. Muscle-damaging agents, such as bupivacaine, have increased gene expression (Xu et al. 2017). In the present study, the pcDNA3.1 vaccine vector which contains the strong CMV promoter was used for the construction of TsE DNA vaccine and 5% bupivacaine hydrochloride was injected into the quadriceps of mice three days before intramuscular vaccination. The results showed that the TsE protein expression in muscular and other cells of inoculated mice was detected. However, their transport mechanism within cells is still unknown.

Antibody assay demonstrated that intramuscular vaccination with TsE DNA triggered not only the systemic specific antibody response (IgG and subclass IgG1/IgG2a, and IgA) and also enteral local mucosal sIgA responses. Anti-rTsE sera from vaccinated mice identified the natural TsE at the epicuticle, stichosome of IIL larvae and around female embryos of this nematode. Anti-Trichinella antibodies (mainly IgG) binding to the epicuticle of enteral invasive stage larvae (6 h IIL) of Trichinella spiralis might be involved in the partial blockade of larval intrusion of intestinal epithelia, because the antigen-antibody complex formation at the larval anterior end could physically separate the larval direct contact with intestinal epithelia and interrupt the larval penetration into the intestinal mucosa (McVay et al. 1998, Wang et al. 2013).

Additionally, the sIgA plays a vital role for mucosal barrier defense and prevents from parasite invasion into the enteral epithelia (Wang et al. 2006, Martinez-Gomez et al. 2009, Li et al. 2018). The T. spiralis-specific sIgA participated and accelerated the parasite expulsion from the intestines. Passive immunisation of mice with anti-Trichinella antibody IgA exhibited an obvious immune protection against challenge infection (Inaba et al. 2003). The generation of mucosal sIgA is Th2-dependent; in particular, IL-10 is a main cytokine that enhances the IgA response (Pompa-Mera et al. 2014). In the present study, the correlation of high intestinal sIgA levels and high IL-10 levels demonstrate that cytokine IL-10 might enhance the intestinal mucosal sIgA response.

Our results also showed a plentiful of enteral mucosal goblet cells/mucins induced in pcDNA3.1-TsE vaccinated mice. Previous studies showed that the enteral stage of T. spiralis was trapped in intestinal mucus of immunised mice (Castillo Alvarez et al. 2013). The main function of enteral mucus is to play a physical barrier to protect intestinal epithelium. It can also cover the worm bodies and limits the worm activity (Hasnain et al. 2012, Sharpe et al. 2018, Sun et al. 2019b). The results also showed that vaccination with TsE DNA elicited the concomitant Th1/Th2 immunity, as shown by a clearly ascended level of Th1 cytokines (IFN-γ and IL-2) and Th2 cytokines (IL-4 and IL-10) following spleen, MLN and PP cells from vaccinated mice being stimulated by rTsE. The concomitant Th1/Th2 response is also important for immune protection against T. spiralis infection (Pompa-Mera et al. 2014).

After being challenged with 300 T. spiralis ML, the mice vaccinated with pcDNA3.1/TsE showed a 17% intestinal AW reduction at 7 dpi and a 39% ML reduction at 42. The worm burden reduction observed in this work is similar to those of intramuscular vaccination with DNA vaccines of T. spiralis enolase, Ts45 protein and a T. spiralis multicystatin-like domain protein (TsMCD-1) (Tang et al. 2012, Zhang et al. 2018b), but it was lower than the protective efficacy of DNA vaccines of T. spiralis nudix hydrolase (54% larval reduction) (Liu et al. 2015a) and newborn larval serine protease-like protein (78%) (Xu et al. 2017). The significant difference between adult and larval burden observed in this study is likely because the TsE DNA vaccine might have an obvious inhibitory effect on the female fecundity and newborn larvae. Previous stud-

Fig. 10. Protective efficacy of pcDNA3.1/TsE vaccination after challenge with 300 T. spiralis infective larvae of Trichinella spiralis in a murine model. A – Intestinal adult worm burden; B – Muscle larvae burden (larvae per gram, LPG). The parasite burden is shown as mean ± SD from pcDNA3.1-TsE-vaccinated mice, empty pcDNA3.1 or PBS group (n=10). * P < 0.01 compared with empty pcDNA3.1 or PBS group.

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ies indicated that intestinal slgA inhibited the female adult reproductive capacity of T. spiralis (Liu et al. 2015a). Additionally, anti-Trichinella antibodies also participated in killing of newborn larvae through an ADCC mode (Song et al. 2018a, Zhang et al. 2020). Hence, although the adult worm reduction in the intestine of mice vaccinated with pcDNA3.1/TsE is low, larval burden reduction in muscle of vaccinated mice is more evident.

Trichinella spiralis is a multicellular parasitic nematode, it has the diverse life-cycle stages (muscle larvae, adult worms and newborn larvae), and transient intestinal and migratory stages (intestinal infective L1-L4 larvae, migratory larvae and pre-encapsulated larvae) (Despommier 1983). During natural T. spiralis infection, each life-cycle stage has its stage-specific antigens (Parkhouse and Ortega-Pierres 1984, Ren et al. 2019). Vaccination of animals with an individual T. spiralis protein molecule only induced a partial protection following challenge infection. The DNA vaccines containing multiple antigenic molecules (or immune costimulatory molecules, such as macrophage migration inhibitory factor, murine CR2 binding domain of C3d and Th2 cytokines) have significantly improved the immune protection (Pompa-Mera et al. 2014).

To prevent and control T. spiralis infection, the vaccination with preventive anti-Trichinella vaccines should interrupt the IIL larval penetration into the intestinal mucosa, to block the IIL larval development to adulthood, to expel the adults from intestine, to impede the NBL generation and kill NBL (Ortega-Pierres et al. 2015). Additionally, Trichinella infection is acquired mainly from the oral ingestion of infected meat. Therefore, the oral polyvalent anti-Trichinella vaccines should be developed. These vaccines should include multiple epitopes from different T. spiralis life-cycle stages (Gu et al. 2020, Qi et al. 2018a)

Vaccination of mice with attenuated Salmonella expressing a T. spiralis gp43 antigen-derived 30-mer peptide fused to the molecular adjuvant C3d-P28 produced a 93% reduction of enteral AW burden after challenge (Pompa-Mera et al. 2014). When the mixed plasmid of pVAX1-Ts43 and pVAX1-Ts45 were used to vaccinate the mice, their larval reduction (52 and 34%) was increased to 76% (Wang et al. 2018).

The multiple epitope vaccines (MEP) have been studied in several parasite infections. In lymphatic filariasis, thioredoxin-transglutaminase MEP produced a significantly higher protection of 63% than the whole protein cocktail vaccine (56%) in a model of jird (Immanuel et al. 2017). This multiple antigenic epitope vaccine has strong immunogenicity and could enhance the protective immune response in vaccinated animals. Screening and identification of these antigenic epitopes are a crucial for the design of a multiple epitope vaccine. The immunogenic epitopes of the proteins might be screened and identified by epitope prediction algorithms with bioinformatics or by corresponding antibodies. These epitopes should be verified for their capacity to trigger strong and long-lasting humoral and cellular immune responses against the infection (Gu et al. 2020).

In conclusion, the results of the present work indicated that intramuscular vaccination of mice with pcDNA3.1/TsE induced a systemic concomitant Th1/Th2 immune response and a local intestinal mucosal slgA response. The immune responses elicited by vaccination with TsE DNA vaccine produced a partial protection against T. spiralis challenge infection. The oral polyvalent vaccine including multiple epitopes from various life-cycle stages of T. spiralis should be developed to improve the protective efficacy of anti-Trichinella vaccines.

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