

Eukaryotic Plasmids with *Toxoplasma gondii* Dense Granule Antigen (GRA 5) and Microneme 3 (MIC3) Genes as a Cocktail DNA Vaccine and Evaluation of Immune Responses in BALB/C Mice

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Abstract

Toxoplasma gondii is an obligate intracellular protozoan that causes toxoplasmosis in human and animal. This parasite is worldwide spread and about one third of people are seropositive. Toxoplasmosis in immuno compromised patients causes serious symptoms. *Toxoplasma gondii* has a lot of various immunogenic antigens. Excreted/secreted antigens could stimulate of the cell mediated immune response and hence it appears to be a good candidate for vaccine in toxoplasmosis. In this study Microneme3 (MIC3) and GRA5 of *Toxoplasma gondii* are used as DNA vaccine.

The results indicated that survival rate of mice that immunized by recombinant plasmid have significant differences with control groups. IgG and IgG_{2a} assay approved significant different between case and control groups (P<0.05). Cytokine assay indicated high level of IFN-γ and low level of IL4 for immunized groups. These results indicated DNA vaccine encoded MIC3 and GRA5 genes of *Toxoplasma gondii* capable to induced partially protection against toxoplasmosis.

Keywords: *Toxoplasma gondii*; DNA vaccine; MIC3; GRA5

Introduction

In toxoplasmosis excretory - secretory antigens and surface antigens have important role in stimulating protective immunity. For this reason antigen excretion - secretion are suggested as candidates for immunization studies [1]. Microneme antigens bind and invade host cells before rhoptry antigens and have a major role in the pathogenicity [2]. One of the most important antigens of Microneme is Microneme protein 3 (MIC3), which is an important protein intakes during the invasion of the host cell, secreted by Toxoplasma. These antigens can be expressed in the tachyzoites, bradyzoites and sporozoites forms. MIC3 is synthesized as a 40 kDa protein and polymerized as double-stranded form with 90 kDa molecular weight protein [3]. GRA antigens are candidates for vaccine and GRA5 have been found in the tachyzoites, bradyzoites and sporozoites forms too [4-11]. GRA5 is secreted as a soluble antigen into the parasitophorous vacuole and may be related to a critical function in parasite-host interactions [12].

The aim of this study was to determine whether DNA vaccination with dense granule protein GRA5 and microneme MIC3 could prime the immune system of BALB/c mice as well as to examine the immunogenicity and protective efficacy of this DNA vaccine against lethal challenge infection with the highly virulent RH strain of *T. gondii*.

Materials and Methods

Parasite

The tachyzoites of *Toxoplasma gondii* RH strain that injected intraperitoneally by serial passage in BALB /c mice then collected and used for mice challenge. For antigen preparation the tachyzoites of *Toxoplasma gondii* RH strain washed with PBS and were stored in -20°C freezer.

Recombinant plasmid construction

The DNA sequence of the Microneme 3 gene obtained from data

bank website NCBI with accession No. AJ132530. All genes encoding the MIC3 about 2247bp and we selected 1052 bp from the entire gene that located within 724 to 1775 and primers were designed with Gene Runner Software as follows [13].

Forward: 5'- CA CAAGCTTATGGCGCTCACCTTCATGGGGG - 3'

Reverse: 5'- ACAGATATCTCACGTCACGGTGTGGGCATGGT - 3'

GRA5: The sequence of GRA5 gene of *T. gondii* RH strain (complete code: 363 bp) was obtained from Gen Bank, with the accession No. EU918733. The size of PCR product was 363 bp, primers were designed with Gene Runner Software as follows [14].

Forward primer 25nt, 5'- AAG CTT ATG GCG TCT GTA AAA CGC G - 3'

Hind III

Reverse primer 27nt, 5'- GAA TTC TTA CTC TTC CTC GGC AAC TTC - 3'

EcoR I

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The recombinant plasmids were transformed into *E. coli*, strain TG1, and following mass replication of the bacterium were extracted from the bacteria using a plasmid extraction kit (Qiagen, Germany).

Grouping of mice

Mice were grouped according to the type of material that the name and number of each group is shown in the following table (Table 1).

Immunization

The mice used in this study were 6 to 8 week-old female BALB/c mice that purchased from Iran's Razi Serum and Vaccine Production Research Institute. Immunization in 3 times within 3 weeks interval (days zero, 21 and 42) and injection of 100 µl per injection intramuscularly in the quadriceps of mice. For injection we used insulin syringe with needle gauge 30 in volume of 100 µl for each injection (containing 100 µg of the plasmid) (Table 1).

Four weeks after the last immunization and control groups, mice inoculated with 10⁴ of live tachyzoites of strain RH were challenged intraperitoneally. The survival rate of different groups of mice was recorded daily.

Evaluation of IgG and subclasses

Evaluation of humoral immunity in mice by measuring the serum levels of total IgG, IgG1 and IgG2a against *Toxoplasma gondii* in two blood samples using ELISA experiment was performed.

To determine the subclasses of antibodies Monoclonal Antibody Isotyping Reagents kit from Sigma Co. was used. The method that used was ELISA and performed according to the instructions Co. For ELISA *Toxoplasma gondii* lysate antigen (TLA) was used with 10 µg/ml concentration.

Antigen preparation

For *Toxoplasma gondii* lysate antigen (TLA) preparation, the tachyzoites were obtained from peritoneal infected mice. Phenyl-methane-sulfonyl-fluoride (PMSF) as anti- protease with concentration 1 mM was added to tachyzoites and then Freeze & Thaw and sonication were done.

Cellular immunity

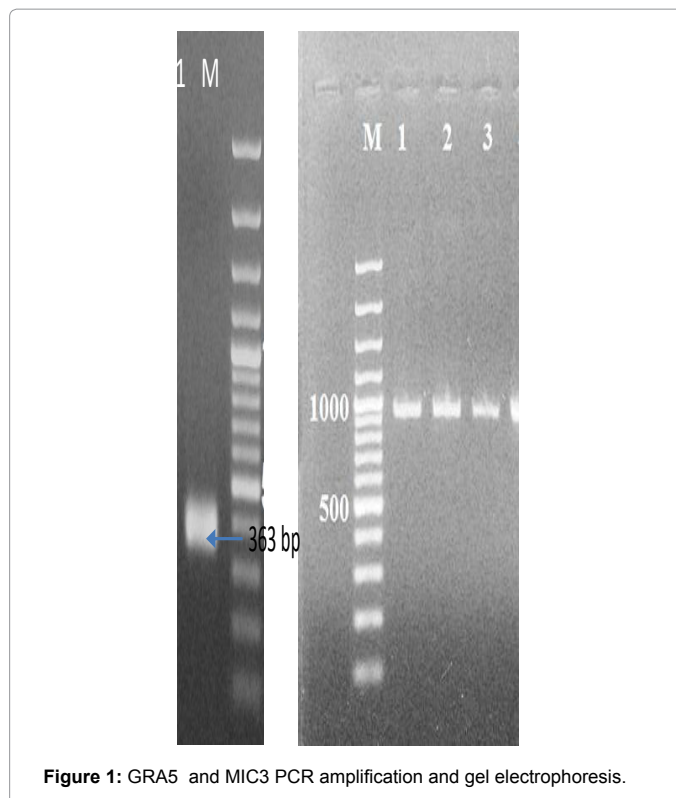
Mouse spleen lymphocyte cultured for measuring the MTT and cytokines assay.

The presence of cytokines IL-4 and IFN-γ in cell culture of mouse spleen lymphocytes were evaluated in five groups of BALB/c mice (immunized and control groups). For this purpose, spleen cells were cultured according to the procedure below.

500 ml lymphocyte suspension from each mouse in two wells of 24-

Amount of injected plasmid µg/100 µl	No of mice for immunological assay	No of mice for survival assay	Groups
50+50	5	5	pcMIC3 + pcDNA3
50+50	5	5	pcGRA5+ pcDNA3
50+50	5	5	pcMIC3+pcGRA5
100	5	5	pcDNA3
100 µl	5	5	PBS

Table 1: Grouping of mice and amount of injected plasmid according to different inserted plasmids



well culture plate was cultured and stimulated with 50 µg/ml of TLA. Plates were incubated for 72 hours in 37°C incubator with 5% CO₂. After this period, supernatants were collected and kept in -70°C freezer.

Cytokine assay

The presence of cytokines IL-4 and IFN-γ in the supernatant of cells from mouse spleen lymphocytes was measured with kit (UcyTech Netherlands) according to manufacturer's recommendations.

Statistical evaluation of the results

To statistical analysis, the results of measurements of total IgG, IgG2a, IgG1, IFN-γ and IL4 and tested to investigate the survival of mice in different groups were entered in SPSS software information page.

As well as the survival of the rats tested in SPSS software information page compiled and compared with each other. Parametric test Kruskalwallis, Kaplan Meiere and Mann-whitney. To draw graphs and Excel software were used.

Result

Results of PCR amplification using plasmid DNA extracted from parasites using primers shown in Figure 1. This figure shows that 363 bp DNA fragment amplified by PCR was about the same size of GRA5 gene of *Toxoplasma gondii*. The primers designed to amplify is specific for GRA5 gene.

The results obtained by use of PCR on plasmid gene-specific primers MIC3 2 shows the DNA fragment amplified by PCR was about 1052 bp and MIC3 *Toxoplasma* genes are of similar size and any genes other than gene amplification has not MIC3, so MIC3 gene -specific primers designed for amplification of *Toxoplasma* MIC3 gene specific primers PCR products in 1% agarose gel ; line 1:PCR Product (1052 bp gene fragment MIC3), M: marker 100 bp.

MIC3 sequencing analysis of *Toxoplasma gondii* cloned in pTZ57R / T using the website www.ncbi.nlm.nih.gov / blast revealed that 1052 bp fragment was cloned in the plasmid is MIC3 gene of *Toxoplasma gondii* (Figure 1).

Lanes M 100bp DNA ladder. Lane 1, PCR product of GRA5 (363 bp) and MIC3 (1052 bp).

Survival assay

The results of survival assay in BALB/c mice after challenged with 10^4 of tachyzoites of *Toxoplasma gondii* RH strain are shown in Fig. 2. The mean days of survival is 4, 4.6, 7.8, 7,8 and 8.6 for PBS, pc GNA3, GRA4, MIC3 and GRA4+MIC3 respectively (Figure 2).

Results of humoral immunity

Results of total IgG measurement: The lowest mean OD value for total IgG in the both of samplings related to the pc-DNA3 and PBS groups, and the highest values were obtained in pcGRA5 group. Furthermore, the cut-off value determined according to measurement of total IgG in serum samples of seronegative mice was (Mean + $3 \times$ SD) 0.234 (Table 2).

Results of the measurement of IgG1 isotype: With regard to the results provided in (Table 3), in the first and second blood samplings, the lowest mean OD values related to PBS group. The highest values related to pcGRA5+ pcMIC3 and pcMIC3 groups, respectively. However, the differences were not statistically significant ($p > 0.05$). Moreover, the cut-off value determined according the IgG1 measurement in serum sample of seronegative mice was $0.165 + 3 \times 0.006 = 0.183$ (Table 3).

Results of the measurement of IgG2a isotype : According to Table 3, the lowest mean OD values in the first and second blood samplings

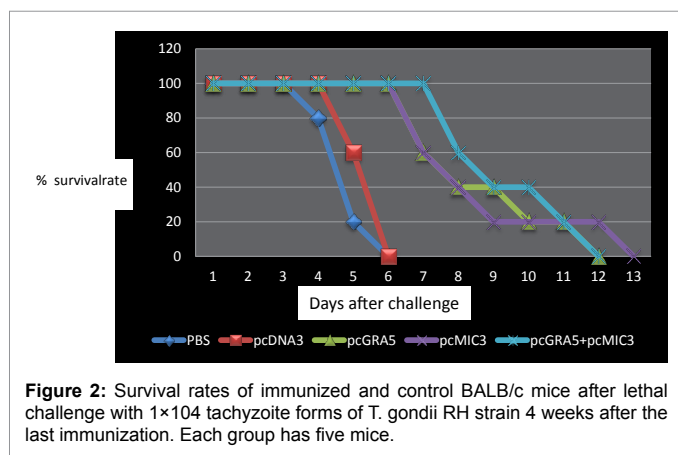


Figure 2: Survival rates of immunized and control BALB/c mice after lethal challenge with 1×10^4 tachyzoite forms of *T. gondii* RH strain 4 weeks after the last immunization. Each group has five mice.

Number of group	Immunization regimen	Total IgG on day 42	
		SD \pm Mean	sig(P<0.05) with groups
1	pcGRA5	0.024 \pm 0.27	* (4,5)
2	pcMIC3+pcGRA5	0.018 \pm 0.29	* (4,5)
3	pcMIC3	0.025 \pm 0.255	* (4,5)
4	PBS	0.027 \pm 0.153	*(1,2,3)
5	pcDNA3	0.016 \pm 0.167	*(1,2,3)

Significantly different from the groups considered in the parentheses according to ANOVA and Mann-whitneytests ($p < 0.05$).

Table 2: Comparison of mean OD values for total IgG determined using ELISA test in serum samples of mice for two blood samplings.

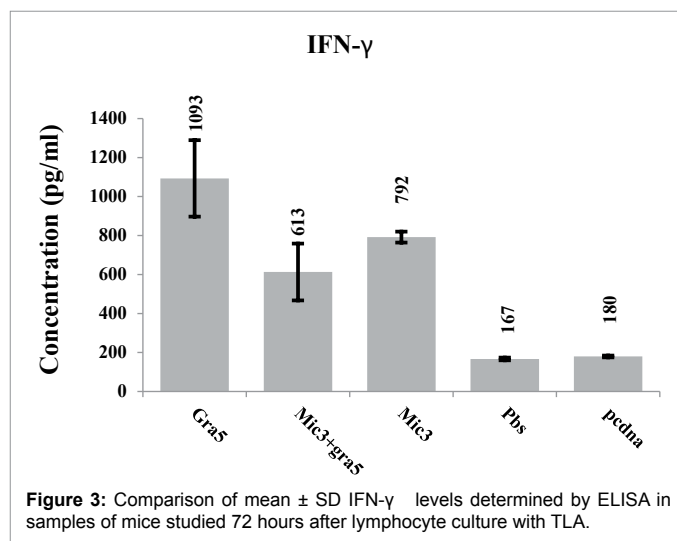


Figure 3: Comparison of mean \pm SD IFN- γ levels determined by ELISA in samples of mice studied 72 hours after lymphocyte culture with TLA.

Number of group	Immunization regimen	IgG1 on day 42	
		SD \pm Mean	sig(P<0.05) with groups
1	pcGRA5	0.009 \pm 0.192	* (2,3)
2	pcMIC3+pcGRA5	0.187 \pm 0.311	* (1,4,5)
3	pcMIC3	0.104 \pm 0.282	* (1,4,5)
4	PBS	0.006 \pm 0.165	*(2,3)
5	pcDNA3	0.024 \pm 0.178	*(2,3)

Significantly different from the groups considered in the parentheses according to ANOVA and Mann-whitneytests ($p < 0.05$).

Table 3: Comparison of mean OD values of IgG1 determined by ELISA in the mice in two blood samplings.

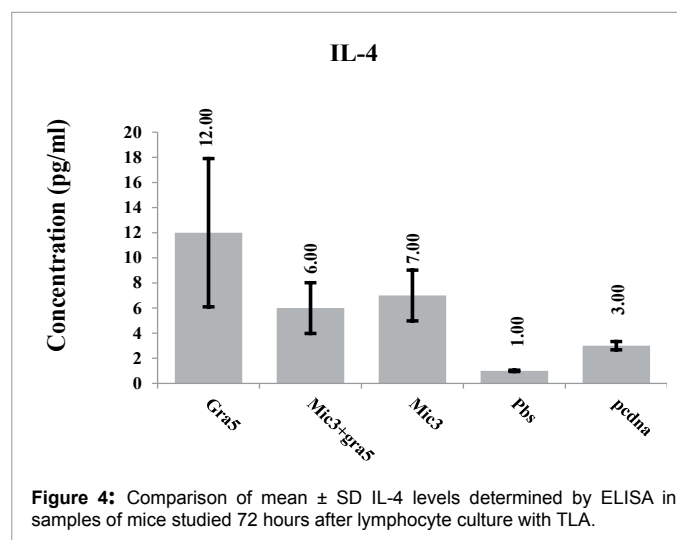


Figure 4: Comparison of mean \pm SD IL-4 levels determined by ELISA in samples of mice studied 72 hours after lymphocyte culture with TLA.

were obtained for the pcDNA3 and PBS groups, respectively. This is while the highest values related to pcMIC3 and pcGRA5+ pcMIC3 groups respectively. Furthermore, the cut-off value obtained according to IgG2a measurement in serum sample of seronegative mice was $0.149 + 3 \times 0.005 = 0.164$ (Table 4).

Results of cellular immunity

Results of IFN- γ and IL-4 measurement: As shown in (Figure 3,4) the mice immunized with pcGRA5 elicited stronger IFN- γ and

Number of group	Immunization regimen	IgG2a on day 42	
		SD ± Mean	sig(P<0.05) with groups
1	pcGRA5	0.050 ± 0.225	*(4,5)
2	pcMIC3 +pcGRA5	0.051 ± 0.247	*(4,5)
3	pcMIC3	0.028 ± 0.257	*(4,5)
4	PBS	0.005 ± 0.149	*(1,2,3)
5	pcDNA3	0.004 ± 0.144	*(1,2,3)

Significantly different from the groups considered in the parentheses according to ANOVA and Mann-whitneytests ($p < 0.05$).

Table 4: Comparison of mean OD values for IgG2a determined using ELISA test in serum samples of mice for two blood samplings.

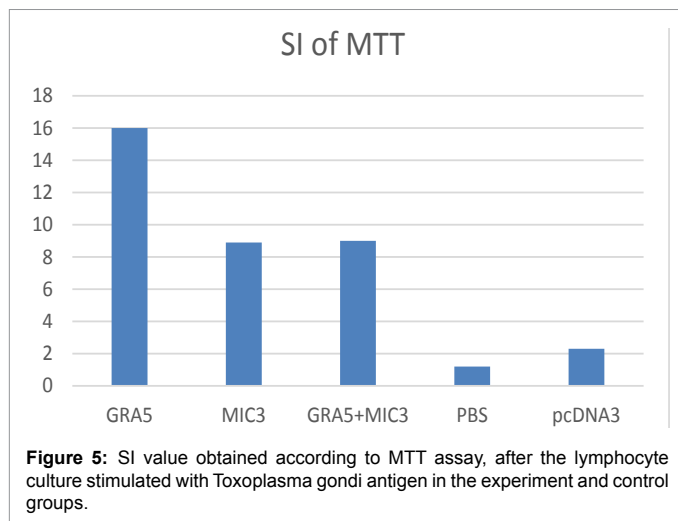


Figure 5: SI value obtained according to MTT assay, after the lymphocyte culture stimulated with *Toxoplasma gondii* antigen in the experiment and control groups.

IL-4 responses than other groups. Significant high level of IFN- γ was observed in spleen cell cultures in mice immunized with pcGRA5 compared with control groups ($P < 0.05$).

Results of MTT assay: The stimulation index (SI) calculated according MTT assay for all groups after the stimulation of lymphocytes with antigen is shown in fig. The results showed the most SI is for GRA5 group that were obtained 16 (Figure 5).

Discussion

Toxoplasmosis is one of the most common infections of humans during its importance in immuno compromised patients is congenital. Since the mother is the time period of pregnancy complications is different. Thus, clearly indicate the need for the development of a more effective vaccine [15]. Research has shown that DNA vaccines could protect humans and animals against intracellular parasites. In recent years, extensive research in finding the right candidates that can induce a protective immune response that has been done. The previous study showed that excretory secretory antigens of *T. gondii* have a main role in stimulating the protective immune system [16]. MIC3 and GRA5 have been found in all stages of the parasite. These results indicated DNA vaccine encoded GRA5 and MIC3 genes of *Toxoplasma gondii* capable to induced partial protection against Toxoplasmosis.

MIC3 is an efficient and strong antigen of *T. gondii* that have been secreted in all stages of the parasite and for this reason is suitable candidate for vaccine [2]. The results show that recombinant plasmids contain GRA5 and MIC3 genes separately and in combination with each other capable to stimulate the immune system and increase the survival rate. Recombinant plasmid inserted with MIC3 gene of *T.*

gondii capable to stimulate cellular immunity with production of high level of IFN- γ and low level of IL-4, MIC3.

The protein of MIC3 has a main role in recognize and attachment to host cells. This protein secreted from microneme in progress of host cell by parasites. MIC3 is very immunogenic antigen in rapid diagnosis latex agglutination test [17]. Ismael et al. found that vaccination of CBA/J mice with inserted plasmid with MIC3 gene could produce high level of IgG against MIC3 and the immune response increased with injection of cloned plasmid with PGM-CSF gene [18].

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Conflict of interest

The authors have no conflicts of interest to declare.

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