Evaluation of the oxidative stress in liver of mice vaccinated with gamma radiation-attenuated Toxoplasma gondii

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ABSTRACT

Background: Researches on attenuated Toxoplasma gondii vaccine have shown its protective effect against challenged infection in addition to its therapeutic rather than preventative activity against some autoimmune and malignant diseases. Rise in oxidative stress is one of the most common signs of toxoplasmosis; hence this study aims to determine local oxidative stress on hepatocytes as well as associated immune variations in mice vaccinated with gamma radiation-attenuated Toxoplasma gondii.

Materials and Methods: Thirty female Swiss albino mice are divided into: normal control; mice infected with T. gondii cysts (infected control) and mice vaccinated with the same infective dose attenuated with 0.4 KgY of gamma radiation (vaccinated group).

Results: Hepatic superoxide dismutase (SOD), Malondialdehyde (MDA) and inducible nitric oxidase (INOs) levels were significantly increased with a significant decrease in reduced glutathione (GSH) in liver tissues of infected control mice. Besides, serum levels of interlekin-10 (IL-10) and tumor necrosis factor-α (TNF-α) were significantly elevated. Moreover, histopathological examinations revealed distinct alterations with decreased nuclear expression of DNA in hepatocytes. Conversely, vaccinated group exhibited an improvement in oxidative stress and elevation in nuclear expression of DNA in hepatocytes collimated with an amelioration of the histopathological alterations. Furthermore, reduced serum IL-10 and TNF-α level were detected.

Conclusion: It is concluded that attenuation of Toxoplasma gondii with gamma radiation exerted protective effect in liver tissues against oxidative stress and DNA damages of hepatocytes with an improvement of host’s immune response.

Keywords: Toxoplasma gondii, Gamma radiation, Liver, Oxidative stress, DNA damage.

INTRODUCTION

Toxoplasma gondii is a zoonotic protozoan parasite that can cause morbidity and mortality in humans as well as in domestic animals. Infection occurs as a result of drinking or eating vegetables washed with water from faucets (¹, ²).

In acute infection, the parasite metabolizes glucose for energy production and its growth, so oxidative stress and free radicals production occur (³). Oxidizing radicals leading to pathological processes can directly damage those free radicals. Immune system is stimulated resulting in defense mechanisms to limit free radical production. These mechanisms employ antioxidant enzymes as SOD and GSH which are inducible by oxidant stress (⁴).

SOD is a first line of defense antioxidant, as it prevents formation of new free radical and converts the existing into less harmful
molecules. It catalyzes the dismutation of superoxide radical into hydrogen peroxide ($\text{H}_2\text{O}_2$) and molecular oxygen participates in deletion and neutralization of toxic reactive oxygen species (ROS). Glutathione is the most abundant non-protein thiol source in the cell. It acts as a substrate for several enzymes, including glutathione peroxidase. It plays an important role in counteracting oxidative damage and maintaining cell health (5-7).

Lipid peroxidation is a degenerative process that alters the fluidity of biological membranes and then causes cell degradation. It causes production and propagation of free radical reactions that involve membrane polyunsaturated fatty acids and damages the enzyme system as well as nuclear DNA (8).

Inducible nitric oxide is an inducible immune inflammatory factor expressed in response to interferon gamma (IFN-γ) which is an important cytokine that eliminate intracellular Toxoplasma gondii. This enzyme can be found in activated macrophages and hepatocytes that generate NO during infection and thereby contributes to control replication or killing of pathogens. It was identified as an anti-Toxoplasma host factor in mice and its genetic deletion or inhibition from the beginning of infection increased the number of T. gondii cysts in the brain (9-11).

Host's immune reaction against toxoplasmosis depends on cellular immunity through secretion of cytokines such as interleukin-12 (IL-12), IFN-γ, IL-10 and TNF-α which are key regulator of immune response cascade (12). However, T. gondii can manipulate this host immunity through deregulation of signaling pathways and controlling host gene transcription by the parasite effector proteins secreted from rhoptries (13). It was reported that TNF-α is a proinflammatory cytokine involved in mediating the pathology and makes up the acute phase reaction of toxoplasmosis. While IL-10 counters the harmful inflammatory effect as a result of type-1 immune response (14, 15).

Vaccination of animals with gamma radiation-attenuated Toxoplasma not only protects against challenged infection but it circumvents the danger of un-irradiated vaccine that remains life-threatening (16). Non replicating Toxoplasma strain is capable of preventing development of some autoimmune diseases as lupus nephritis and arthritis in experimental animals (17, 18). Furthermore, its antitumor activity against ovarian carcinoma, leukemia, sarcoma, Lewis lung and melanoma had been demonstrated (19-21).

Attenuation of T. gondii with radiation induces marked morphological changes such as decrease in rhoptries, disappearance of golgi complex and mitochondria in addition to decreased number of electron dense granules. Severe damage to various intracellular organelles including the nucleus occurs so the parasites lose their reproductive ability while still maintaining immunogenicity (22).

Based on previous studies that reported the immunoprotective effect of gamma radiation-attenuated Toxoplasma vaccine (23) and our recent model demonstrated that using this vaccine produced immune protection against ovarian infiltration in Ehrlich ascites carcinoma-bearing mice (24), this study is undertaken to spot the light on the effect of vaccination on hepatic oxidative stress and the host's immune response.

**MATERIALS AND METHODS**

Thirty Swiss albino female mice 8-10 weeks-old with 22-25g weight were used in the current study. They were obtained from the breeding unit of the Egyptian Organization for Biological Products and Vaccines, housed in cages (5 animals/cage) and maintained on a commercial standard pellet diet and tap water ad libitum. They were also maintained under standard environmental conditions, continually monitored for survival and clinical condition till the end of experiment. Low virulence T. gondii Me49 cysts, which were harvested from the brain of mice that had been intraperitonealy inoculated according to Atmaca et al. (25) were used to induce acute infection. Ten mice were orally infected with ten T. gondii cysts (20) as infected group. The second group was vaccinated with the same dose of T. gondii cysts.
that were exposed to attenuating dose (0.4 KGY) of gamma radiation (26) as vaccinated group and the third group was used as normal control. The study was conducted in the National Center for Radiation Research and Technology (NCRRT) Cairo, Egypt and the animal care was taken in accordance with the recommendations of the Scientific Research Ethical Committee.

Three weeks post infection, the experiment was terminated by anaesthetizing the animals with ether. Liver tissue samples were collected half of them were fixed in 10% phosphate buffered formalin (pH 7.4), embedded in paraffin wax and sections of 5-micrometer thickness were cut. Slides were stained with Haematoxylin and eosin for histological examination (27) and with Feulgen’s stain for histochemical demonstration of DNA (28). The other half of liver tissues of each group were weighted and placed on PBS buffer (1:1) and subsequent homogenization was done then centrifuged at 4000g for 20 minutes. The supernatant was collected for biochemical study. Blood samples were collected, centrifuged and sera were separated for determination of TNF-α and IL-10.

Chemicals and reagent in this study were obtained from Sigma-Aldrich Chemical Co., St. Louis, MO, USA. All ELISA assays for SOD, GSH, MDA and iNOS levels were performed according to the manufacturer’s instructions of commercial kit (Bio-diagnostic, Dokki, Giza, Egypt) and serum cytokines according to MyBioSource kit (Glory Science Co., Ltd, USA).

**Determination of SOD activity**

The assay for super oxide dismutase activity involved inhibition of nitroblue tetrazolium (NBT) reduction, with xanthine oxidase. One unit SOD was defined as the amount of protein that inhibits the rate of NBT reduction by 50%. It was determined at 560 nm and expressed as U/g (30).

**Determination of GSH**

Reduced glutathione was determined as previously described by Beutler, (31). The method based on the reduction of 5,5'-dithiobis-2-nitrobenzoic acid (DTNB) with glutathione to produce a yellow compound. The reduced chromogen was directly proportional to GSH concentration and its absorbance was determined at 405 nm and expressed as mmol/g tissue.

**Determination of lipid peroxidation**

Levels of thiobarbituric acid reacting substance (TBARS) as a marker of lipid peroxidation were determined according to the method described by Ohkawa et al. (29). The formation of pink color, as a result of the reaction was measured at 535 nm and expressed as nmol/g liver tissue.

**Inducible Nitric oxidase measurement**

iNOS in the crude liver tissue homogenate was determined using a commercial kit (Bio diagnostic, Egypt).

**Determination of serum TNF-α and IL-10**

Serum levels of TNF-α and IL-10 were assayed by Enzyme linked-Immunosorbent Assay (ELISA) using the MyBioSource kit. Measurements were done according to methods of Sutterwala et al. (32).

**Statistical analysis**

All data were expressed as mean ±SE (standard error). Data were assessed by using a one-way ANOVA using SPSS 15.0 program and p<0.05 was considered statistically significant.

**RESULTS**

**Histopathological findings**

Figure 1 represents H&E stained liver sections where normal control group exhibited average central veins surrounded by hepatocytes arranged in single-cell cords with intervening blood sinusoids, and normal portal tracts with average bile ducts and portal veins (figure 1, a, a1). Liver sections of infected control group showed portal tract with markedly dilated portal veins and inflammatory cellular infiltration around the portal area. Dilated central vein with detached lining and peri-venular inflammatory infiltrate can be also
detected (figure 1, b, b1). Vaccinated group revealed average central veins and mild intra-lobular inflammatory infiltrate. Also, dilated congested portal veins with average hepatocytes in peri-portal area were clearly seen (figure 1, c, c1).

**Histochemical expression of DNA Feulgen’s stain**

Expression of hepatic DNA content was shown in figure 2, table 1. Normal group revealed marked positivity for Feulgen (14.5±1.2), average DNA distribution and intact nuclear membrane in most of hepatocytes. *Toxoplasma*-infected group revealed mild positivity for Feulgen (2.9±0.6) with loss of nuclear membrane in most of hepatocyte nuclei in peri-venular area and high significant decrease (*P*<0.001) compared to normal control. Moderate positivity for Feulgen (9±0.8) with few reduction of DNA in hepatocyte nuclei and significant decrease was detected in vaccinated group compared to normal (*P*<0.01) and high significant increase compared to infected group (*P*<0.001).

Biochemical analysis of oxidative stress in liver tissues of infected control group showed significant increase in SOD levels compared to normal control while vaccinated mice group revealed significant decrease compared to infected control (figure 3a). GSH levels were significantly decreased in infected control mice group compared to normal control however; their levels were significantly increased in vaccinated group (figure 3b). A significant increase in MDA was detected in infected control group compared to the normal group on contrary to vaccinated group showed a significant decreased compared to infected control one (figure 3c) (*P*<0.001).

Regarding INOs, the mean±SE was 94.8±3.8 in infected group while it was 43.6±0.9 in normal control with high significant increase (*P*<0.001). In vaccinated group it was 64.2±1.4 showing high significant decrease versus infected control group (*P*<0.001) and significant increase (*P*<0.01) versus normal one (table 1).

To gain insight into the immunological mode of action of irradiated *T. gondii*, we next examined the influence of vaccination on the serum inflammatory cytokines (table 2). TNF-α and IL-10 levels were 42.3±0.8 and 67.8±4.9 respectively in infected control group while they were 9.8±0.5 and 140.2±3 respectively in normal control with high significant increase as *P*<0.001. On contrary, they were 18.4±0.7 and 409.2±5.3 respectively in vaccinated group showing high significant decrease (*P*<0.001) versus infected control one.

![Figure 1. Liver section of normal group showing average portal tract with average portal vein (PV), average bile duct (black arrow), average central vein (red arrow) and average hepatocytes (blue arrow) (a, a1). Infected group showed expanded portal tract with portal inflammatory infiltrate (black arrow) and markedly dilated portal veins (PV). Dilated central vein (CV) with detached lining (blue arrow) and peri-venular inflammatory infiltrate (red arrow) can be detected (b, b1). Vaccinated group showed portal tract with dilated congested portal vein (PV), average central veins (black arrow), and mild intra-lobular inflammatory infiltrate (blue arrow), markedly dilated portal vein (PV), and average hepatocytes in peri-portal area (red arrow) (c, c1) (a-c X200, a1-c1 X400).](https://ijrr.com/article-19.2.339)

![Table 1. Expression of INOs and DNA in hepatocytes of experimental groups.](https://ijrr.com/table-19.2.339)

**Table 1.** Expression of INOs and DNA in hepatocytes of experimental groups.

<table>
<thead>
<tr>
<th>Groups</th>
<th>INOs</th>
<th>DNA</th>
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<tbody>
<tr>
<td>Normal control</td>
<td>43.6±0.9</td>
<td>14.5±1.2</td>
</tr>
<tr>
<td>Infected control</td>
<td>94.8±3.8</td>
<td>2.9±0.6</td>
</tr>
<tr>
<td>Vaccinated group</td>
<td>64.2±1.4</td>
<td>9.8±0.8</td>
</tr>
</tbody>
</table>

Data are expressed as mean ±SE.

* Significant compared to normal group *P* < 0.01.

** High significant compared to normal group *P* < 0.001.

+++ High significant compared to infected group *P*<0.001.
Figure 2. Normal control group showing marked positivity for Feulgen (+++) with average DNA distribution in most of rounded hepatocyte nuclei with intact nuclear membrane in peri-venular area (red arrows) [a, a1]. Infected group showed mild positivity for Feulgen (+) with marked reduction of DNA and loss of nuclear membrane in most of hepatocyte nuclei in peri-venular area (red arrows) [b, b1]. Vaccinated group showed moderate positivity for Feulgen (++) with reduction of DNA in few of hepatocyte nuclei with loss of nuclear membrane in peri-portal area (red arrows) [c, c1] (Feulgen stain a-c X 400, a1-c1X1000).

Figure 3. Levels of oxidative stress markers in different studied groups. (a) level of SOD, (b) level of GSH and (c) level of MAD. Each value represents the mean ± S.E. ** Significant difference versus normal control group. ++ Significant difference versus infected control group at P < 0.001.

Table 2. Mean±SE of IL-10 and TNF-α in the sera of experimental mice. Data are expressed as mean ± SE.
* Significant compared to normal group.
+ Significant compared to infected group.
P < 0.001
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**DISCUSSION**

Most of previous studies have focused on the attenuation of *Toxoplasma gondii* by radiations that help maintaining their integrity, antigenicity while increasing immunogenicity so eliciting an immune-protective effect. For instance, it has been previously reported that vaccination with gamma radiation-attenuated *Toxoplasma* tachyzoites resulted in an amelioration of histopathological changes in liver, lung, kidney, spleen and testis of experimentally infected mice (22). Recently, using gamma radiation-attenuated *T. gondii* cysts vaccine has demonstrated immune protection against ovarian infiltration in mice-bearing Ehrlich ascites carcinoma (24). As one of the most common signs of toxoplasmosis is a rise of hepatic oxidative stress, hence the aim of the present study is to determine the impact of gamma irradiated *Toxoplasma* cyst vaccine on hepatic oxidative stress, INOs and immune response in mice.

The data of this study demonstrated a significant decrease level of GSH in infected group compared to normal. This is in agreement with previous reports that revealed a decrease of GSH and increase of GPX activity in different tissues of *Toxoplasma*-infected hosts (33-35). A significant increase of SOD in the infected group was noted, which is in accordance to the study of Turkoglu et al. (36) who reported an increase of SOD in liver tissues of the infected mice thirty days post infection. In the literature, SOD level was reported to be increased several weeks post-infection with other parasites as *Fasciola hepatica*, *Trichinella spiralis*, *Schistosoma mansoni* and *Trypanosoma cruzi* several weeks post-infection (37-39).

Lipid peroxidation, determined by MDA concentration, was significantly increased in infected group. In accordance of these results is the previous works which reported higher level of MDA in liver of infected rats (33, 36). Significant increase of INOS expression in hepatocytes was detected in our study. This is compatible with the previous study of Dincel and Atmaca (40) who reported a remarkable increase of it during *T. gondii* encephalitis in mice 74 days after inoculation. It was reported to be strongly stimulated by IFN-γ produced by parasite-infected monocytes (41). In agreement to our results, another study reported a significant increase of hepatic lipid peroxidation and NO and decrease in GSH activity in liver homogenates of control infected group (42).

In *Toxoplasma*-vaccinated group, a high significant increase in GSH with decrease of SOD, MDA and INOs levels were detected compared to infected group. Similar to mice vaccinated with gamma radiation-attenuated *Toxoplasma*, rats vaccinated with gamma radiation-attenuated *Toxocara canis* larvae revealed a significant decrease in SOD and MDA with increase of GSH in kidney tissues compared to infected control group (43).

Regarding histopathological changes and histochemical expression of nuclear DNA, dilated central and portal veins with inflammatory cellular infiltrate were clearly observed in infected group. Furthermore, mild positivity for Feulgen’s stain with marked reduction of DNA and loss of nuclear membrane in most of hepatocyte nuclei in peri-veinular area was also detected. It has been shown that free radicals from peroxidase in Kupffer cells play a predominant role in hepatic pathogenesis and the decrease in the levels of liver SOD may lead to excess superoxide radicals which may interact with hydrogen peroxides and organic peroxides with generation of highly reactive entities that can attack DNA (44).

In the present study, vaccinated group revealed an amelioration of these histopathological changes and increased expression of nuclear DNA in hepatocytes compared to infected group. This was in consistence with previous study by Amin and Hafez (2015) who reported minimal histopathological alterations with normal architecture of the hepatic lobules and minimal degeneration of nuclear DNA in mice vaccinated with either gamma or UV radiation-attenuated *Toxoplasma*. This was explained by the direct effect of radiations that weaken the parasites and induce changes leading to decreased or abolished reproduction by damaging of the DNA molecules. However irradiated parasite

maintains viability, a respiratory response and preserved protein and nucleic acid synthesis so, it present antigens to the host’s immune system and elicit cellular immunity and cytokine responses in a highly similar way to natural infection [22, 45].

In current study, an increasing level of TNF-α was recorded in infected group compared to normal one. This proapoptotic molecule and others as nitric oxide (NO) are produced by macrophages as a result of invasion with Toxoplasma parasite to maintain the viability of the cell during its residence (46). Also, an elevation of IL-10 level was detected which was in line with Mammary et al. (47) who reported that ROP16, a molecule secreted by rhoptries, stimulates IL-10 expression and induces activation of macrophages thus promoting Th-2 response. Likewise, previous studies reported a significant increase of TNF-α and IL-10 in T. gondii-infected patients compared with healthy control (48, 49).

On the other hand, vaccinated group showed a significant decrease of their levels compared to infected group and significant increase compared to normal control one which may be due to the immunogenic effect of living attenuated parasite that can be recognized as protein antigen. Similar results were recorded by Moawad et al. (10) who reported a significant decrease in serum levels of IL-10 and TNF-α in mice vaccinated with either UV or gamma-radiation attenuated Toxoplasma tachyzoites compared to infected control.

**CONCLUSION**

From the results obtained, it is concluded that gamma radiation-attenuated non replicating vaccine to exclude the delirious effects of imposing infection by using live one.

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**Conflicts of interest:** Declared none.

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