



Zirconium oxide nanoparticles: Protocidal effect against hydatid cyst protoscoleces with interleukins evaluation in mice

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Abstract

The current study investigated the effect of zirconium oxide nanoparticles NPs on the levels of interleukins in mice inoculated with protoscoleces of *Echinococcus granulosus*, after exposure to ZrO₂ NPs at the doses, 5, 10, 50, and 100 μ g/ml for 30 minutes, and different periods of infection, on the other hand, control groups inoculated with unexposed protoscoleces to the nanoparticles. Following the infection, blood serum was obtained from animals to determine interleukin levels using sandwich ELISA technique three, four, and five months following the infection. The outcome of the study demonstrated a significant decrease in IL-5, IL-12 levels, and a significant increase in IFN- γ levels in treated mice compared with control groups. The lowest value of IL-5 was 11.164 ng/ml at the dose of 100 μ g/ml compared to the control group 47.446 ng/ml, four months after infestation, while the lowest value for IL-12 was 12.824 ng/ml, at 100 μ g/ml, compared to 112.485 ng/mL in the control group, five months to infection. IFN- γ was significantly increased in the treated mice, 207.39 ng/ml, at 100 μ g/ml, compared to 110.07 ng/mL in the control group, five months after injury. Based on the research outcome, ZrO₂ nanoparticles could be exploited as a consequential titer for assessing the immune situation and, consequently as a dynamic curative of hydatidosis.

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Introduction

Cystic echinococcosis is a zoonotic disease induced by *Echinococcus granulosus* larval stage infection. The disease causes health problems worldwide and causes economic losses estimated at 3\$ billion annually (1,2). Hydatid cysts develop in different host organs, such as the liver 70%, lungs 30%, and even in the heart, brain, bones, spleen and kidneys, but at a lower rate, leading to death (3-5). Iraq is one of the countries affected by this endemic disease, transmitted to humans by eating food contaminated with *E. granulosus* eggs or by contact with infected dogs (6,7). The diagnosis of this disease is still challenging. Many cases show symptoms after years due to the absence of clear pathological signs. Symptoms implicate abdomen discomfort, nausea,

obstruction of the bile ducts, in the chest, breathing difficulties, and neurological signs due to seizures as a result of brain injury (8,9). Some of the most common methods for detecting hydatid cysts in humans are X-ray, ultrasound, computed tomography, and magnetic resonance imaging (10-12). These methods are characterized by an inability to differentiate cysts from abscesses. Therefore, early diagnosis must be confirmed through other high-quality tests, such as serological techniques, such as immunoelectrophoresis, double diffusion test, and direct immunofluorescence assay, as new sensitive techniques (13,14). The accidental rupture of cysts or rupture during surgery is a serious medical problem due to the ability of the protoscoleces to develop into new cysts. Thus, many chemicals were used as protoscolicidal pre cyst removal (15,16). Infection with

Echinococcus granulosus prompts a humoral and cellular response in the host, increasing in serum antibodies as an essential titer for diagnosing the disease and Th1 and Th2 cytokines (17). The interaction between the host and the parasite is significantly impacted by host immunity. The parasite secretes substances that impact immunological cells, releasing antibodies, activating T cells, and inducing an inflammatory immune response. Antibodies are essential to combat the parasites, protect the host, and prevent disease. *E. granulosus* induce a biphasic response, represented by Th2 response implicates cytokines, as well as Th1 cytokines. While Th17 cells are potential pro-inflammatory cells contribute the secretion of IL-22 and IL-21 (18,19). Roles of cytokines in host immunity differ with different species of parasites, their size, location within the host, and their metabolic products. Heavy infection with parasites stimulates Th2 cytokines (20). Most studies on mice and humans confirmed the dominance of Th1 cytokine response, characterized by the IFN- γ release produced by dendritic cells with IL-12. It was found that both have an effective role in eliminating parasites during early infection. The parasite can influence the host's immune response by secretory excretory secretions, resulting in a Th2 response and the parasite's survival, associated with the cytokines IL-4, IL-5, and IL-10, depending on the parasite's receptor ability, which leads to chronic infection (21-23). Cytokines represent one of the pharmacological agents capable of improving the immune response to ABZ. In an experimental study on mice, the mixture of ABZ with IFN- γ and IL-12 had prophylactic effects of 100% and 97%, respectively against echinococcosis, indicating the importance of cytokine therapy in preventing disease (24). Patients with cystic echinococcosis who did not react to treatment exhibited Th2 type response, while patients who responded to treatment revealed a Th1 type 1 response. Researchers explored interleukin -10 as a quantitative signal of the parasite's escape from the host's immune response (25). Nanoparticles have immunomodulatory properties that help enhance weak conjugated antigens' antigenic properties. They act as adjuvants, antigenic properties differ according to their size and surface charge, and the size of nanoparticles determines whether the loaded nanoparticles induce Th1 cells with interferon- γ and Th2 cells with IL-4, IL-10 when recognized by the immune system. Several studies confirm that nanostructured vaccine formulations provide certain advantages against pathogens in the induced immune response, as they generally work to stabilize and increase the supply of antigens or act as modulators of the immune system (26-30). Other research applied to mice treated with a group of antihelminthic drugs Oxfendazol, Praziquantel, and Albendazole after infection with hydatid cyst showed an increase in the level of IFN- γ in the hepatic and spleen inflammatory cells in the treated groups, these results confirmed that effective treatment reduces IL-10 in human infections leading to in situ immune activation by increasing

L-2 IFN- γ . It was confirmed that elimination of the parasite will restore the host cell response (31,32).

The current study targeted investigating levels of interleukins, as a considerable indicator of immune response, in mice inoculated with protoscoleces of hydatid cysts exposed to ZrO₂ nanoparticles as an alternate protoscolicidal therapy during surgical operations.

Materials and methods

Ethical approve

The study was accomplished in accordance with the declaration of Helsinki, and the protocol was approved by the Ethics committee of College of Veterinary Medicine at University of Mosul in ethical approval code UM.VET.2022.083 in 19/10/2022.

Animals

Seventy-five male albino Scottish mice, 3-4 weeks old, were used and reared in the animal house of The College of Veterinary Medicine, University of Mosul. Mice were inoculated with 2000 protoscoleces viable and exposed to different concentrations of ZrO₂, 5, 10, 50, and 100 μ g/ml for 30 minutes. Control groups were administered with 2000 vital unexposed protoscoleces. After 3, 4, and 5 months of infection, mice were anesthetized, blood was obtained from the ophthalmic Venous Plexus, according to Waynforth (33), and serum was obtained according to authenticated methods kept in the freezer at -20 centigrade till use.

Cytokine assays

Sandwich ELISA technique, ELISA kit from Bioassay Technology Laboratory (BT Lab), China, used to detect interleukins in the blood serum of infected animals by using a 96-hole outlet. Serum samples, all reagents, and standard solutions were prepared at room temperature. The standard protein for each interleukin was prepared, five numbered Eppendorf tubes were used, and 120 μ l of the standard diluent was added to each tube. The number of strips required for measurement was determined. The strips were inserted in the places designated for use. 50 μ l of standard solutions and 40 μ l of samples were added to the holes. 10 μ l of diluted serum was added to the samples in the pit, then 50 μ l of streptavidin HRP to the samples and the standard. The plate was covered with sealer, incubated posteriorly for 60 minutes at 37°C, and washed five times with at least 0.35 μ l of the washing solution for a minute each. 50 μ l of substrate solution A and 50 μ l of substrate solution B were added to each well. After that the plate was incubated for ten minutes at 37 °C. Subsequently, 50 μ l of the stop solution was added to each hole, so the blue color turned yellowish immediately. The optical density was determined immediately using a microplate reader set at a wavelength of 450 nanometers within 10 minutes of adding the stop solution.

Statistical analysis

Data were analyzed by complete random design and analysis of variance. Degrees of differences were estimated by Duncan’s multiple range test. Statistical analysis Software version nine was used (34).

Results

The current study showed significant differences ($P \leq 0.01$) in the level of cytokines IL-5, IL-12, and IFN- γ ($P \leq 0.05$) in mice administrated with protoscoleces treated with zirconium nanoparticles compared to the control group (Table 1). Table 2 demonstrates substantial variations in the level of cytokines ($P \leq 0.01$), represented by a decrease in the level of IL-5 in the treated collection, 11.164 ng/ml at 100 $\mu\text{g/ml}$, four months postinfection, compared to untreated ones, 75.171 ng/ml, five months next infection. Table 3 detected a significant decrease in IL-12 level, 12.824 ng/ml at 100 $\mu\text{g/ml}$ throughout infection, compared with the

control collection of 112.485 ng/ml after the fifth month of infection. Table 4 detected substantial variations represented by an increase in the level of IFN- γ in the treated collection, 207.39 ng/ml, five months after infection, compared to the control group, 110 ng/ml, three months post infection.

Discussion

The outcome of the current research showed a significant rise in the interferon-gamma level and a decrease in the level of interleukins IL-5 and IL-12. Cytokine therapy is one of the effective ways to prevent the formation of hydatid cysts and to treat cases in which surgeries cannot be performed. TH1 cytokines are essential in initiating protective immune responses against pathogens comprising bacteria, viruses, and parasites. IFN- γ IL-12 stimulates increased macrophages and NK cells as antigen-presenting cells to destroy pathogens (35).

Table 1: Level of cytokines IL-5, IL-12, and IFN- γ in experimental mice

Parameters	Degree of freedom	Duration	Handling	Duration \times handling	SE
		two	four	eight	Sixty
IL-5	Summation of square	1370.290**	9066.392**	4397.281**	3325.80991
	medium square	685.1452**	2266.5981**	549.66020**	
	F amount	12.36	40.89	9.92	
IL-12	Summation of square	22.48057	60063.527**	5006.26708**	1684.03633
	medium square	11.24029	15015.88185**	625.78338**	
	F amount	0.4	535.00	22.30	
IFN- γ	Summation of square	1580.56506	5331.61120	92223.2389*	300620.479
	medium square	790.2825	1332.902	11527.904*	
	F amount	0.16	0.27	2.30	

** Considerable significance $P \leq 0.01$. *Considerable significance $P \leq 0.05$.

Table 2: Level of IL-5 in experimental mice

Duration	Three months	Four months	Five months	Medium
Control	45.507cb	47.446b	75.171a	56.041a
First group (5 $\mu\text{g/ml}$)	40.057bcd	42.383bc	37.833bcd	40.273b
Second group (10 $\mu\text{g/ml}$)	42.039bc	35.369cd	33.977f	37.128b
Third group (50 $\mu\text{g/ml}$)	35.325cd	29.611ed	38.276bcd	34.404b
Fourth group (100 $\mu\text{g/ml}$)	18.950fg	11.164g	22.619ef	17.733c
Medium	36.3756b	33.1946b	41.5752a	

Similar letters indicate no significant difference and different letters indicate significant differences.

Table 3: Level of IL-12 in experimental mice

Duration	Three months	Four months	Five months	Medium
Control	73.545c	96.242b	112.485a	94.090a
First group (5 $\mu\text{g/ml}$)	49.071d	36.880ef	43.359de	43.103b
Second group (10 $\mu\text{g/ml}$)	35.021f	32.024gf	24.450h	30.498c
Third group (50 $\mu\text{g/ml}$)	26.634gh	19.842hi	19.446hi	21.974d
Fourth group (100 $\mu\text{g/ml}$)	15.790i	13.208i	12.824i	13.940d
Medium	78.9415a	66.8475b	63.2238c	

Similar letters indicate no significant difference and different letters indicate significant differences.

Table 4: Level of IFN- γ in experimental mice

Duration	Three months	Four months	Five months	Medium
Control	110.07ab	148.25ab	175.09ab	144.47a
First group (5 μ g/ml)	149.88ab	69.04b	101.18ab	106.7a
Second group (10 μ g/ml)	119.64ab	133.80ab	116.70ab	123.38a
Third group (50 μ g/ml)	122.77ab	111.75ab	202.40a	145.64a
Fourth group (100 μ g/ml)	126.09ab	160.30ab	207.39a	164.593a
Medium	125.69c	124.628b	160.552a	

Similar letters indicate no significant difference and different letters indicate significant differences.

Recent research suggests that Th1 and Th2 reactions act together during infection with CE. Th2 response is connected with disease vulnerability (Active cyst) while preventive immunity is associated with Th1 response (Inactive cyst). The biological importance of the cellular response appears in some infectious diseases, where the responses of T-cells of the first type TH1 and the second type TH2 are associated with resistance and sensitivity, and immunological studies that have been applied to humans have indicated mixed immune responses are associated with infection, or susceptibility to disease in the case of active cysts, whereas TH1 responses are associated with protective immunity or resistance to disease in the case of inactive cysts, patients with cystic echinococcosis who did not respond well to chemotherapy usually showed a TH2 response, while patients who did not show TH1 response (36-38).

Antigen B (Ag B) affected the immune system response in rabbits immunized with AgB and full Freund's adjuvant. Three intramuscular booster shots were carried out again after two weeks following the immunization using incomplete Freund's adjuvant, and five rabbits were treated as the control group. IgG, IFN- γ , and IL-10 levels were determined using ELISA. IgG levels in the immunized rabbits were significantly higher 45.9 ng/ml than in the control group 16.6 ng/ml. The differences in Th1 (IFN- γ) and Th2 (IL-10) cytokine levels between the vaccinated and control groups were statistically significant, and the mean of IFN- γ values in the immunized rabbits 28.6 pg/ml were less than those in the control groups 83.6 pg/ml. The findings showed that AgB immunization-induced Th2 immune response in immunized rabbits, as evidenced by a large increase in total IgG and IL-10 and a significant decrease in IFN- γ concentration (39). Siracusano *et al.* (40) showed that IFN- γ improved the ability of macrophages to kill protozoa while IL-4, IL-10 reduced the activity. IFN- γ is the key to phagocytic cell function through NO production and inhibition of helminth growth and function. In addition to other pathogens, it thus plays a role in establishing protective immunity by TH1 during *Echinococcus granulosus* infestation, and the destructive role of nitric oxide associated with the production of TH1 response acting to reduce the vitality of the parasite. Elevated levels of IFN- γ and NO are found both in vitro and in vivo during human infection, while levels are not detectable upon reinfection.

Through studies in humans and mice, most researches addressed the dominance of the Th1 response, characterized by IFN- γ release. After being produced by DCS dendritic cells with IL-12, it was found that both are effective in eliminating the parasite at an early stage. TH2 and parasite survival associated explicitly with IL-5, IL-4, IL-10 and transitional growth factor (TGF) are generally associated with the receptive ability of the parasite and lead to chronic infection (40-42). It was also revealed that when the cyst dies naturally, is killed by chemotherapy, or is surgically removed, Th2 cell responses rapidly decline and Th1 cell response is dominant. This can be explained by the fact that Th1 lymphocytes contribute significantly to the inactive stages of the disease, with Th2 lymphocytes being more critical in the transitional and active stages (5,43). A recent study concurs with research concerned with pre-treatment IL-5 levels to monitor the change of cytokines in mice infected with various *Echinococcus* components, ELISA technique registered a shift in blood levels of IL-2, Interferon-gamma, TNF-alpha, IL-4, IL-5 and IL10 during 220-day infection compared to the group of animals that were not infected. Cytokine levels increased significantly at 260 days, IL-2 level reached a peak 80 days after infection, posteriorly quickly collapsed after 140 days post-injury, compared to uninfected series, a high amount of TNF-alpha was determined after 40 days; peaked 100 days after infection and dropped off swiftly after 140 days. After eighty days post-infection, IFN-gamma levels peaked and steadily started declining. Before 80 days, the levels of Interleukin -4, Interleukin -5, and Interleukin -10 stayed lower and significantly increased following a hundred days. At 100 days post-injury, the levels of IL-4 and IL-10 peaked, while IL-5 peaked at 140 days post-injury. The evidence pointed to vital host defense mechanism against metacestodes as the Th2 AMI (antibody-mediated immunity) is significant in the later stages of infection and has an essential role in the initial phases of infection (3).

Conclusion

According to the research, CE had each Th1 and Th2 cytokine profiles, with Th2 predominating throughout the active stage of the illness, and significantly decreasing in laboratory mice that responded to remediation with the

nanoparticles. The results demonstrated a considerable protoscolicidal activity of Zirconium oxide nanoparticles against hydatid cysts in mice by modulating the levels of interleukins through increasing INF- γ and decreasing IL-12 and IL-5, respectively. These nanoparticles can be used as a favorable alternative for hydatid cyst remedies to overcome the disadvantages of drug side effects.

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Conflict of interest

The authors profess that they have no conflict of interest.

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أكسيد الزركونيوم النانوي وتأثيره كقاتل لرؤيسات الكيس العدري في الفئران مع تقييم الانترليوكينات فيها

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و لؤي جميل رشان^٢

^١ قسم علوم الحياة، كلية التربية للعلوم الصرفة، جامعة الموصل، الموصل، العراق، ^٢ مركز البحوث بجامعة ظفار، صلالة، سلطنة عمان

الخلاصة

تحررت الدراسة الحالية تأثير أكسيد الزركونيوم النانوية في مستوى الانترليوكينات في الفئران المجرعة بالرؤيسات الأولية للمشوكة الحبيبية، المعرضة لأوكسيد الزركونيوم النانوية عند الجرعات ٥، ١٠، ٥٠، و ١٠٠ ميكروغرام/ مليلتر، لمدة ثلاثين دقيقة، و لفترات زمنية مختلفة من الاصابة، من جهة أخرى جرعت مجاميع السيطرة بالرؤيسات الأولية غير المعرضة للمادة النانوية. بعد ثلاثة، أربعة، وخمسة أشهر من الاصابة، تم الحصول على مصل الدم من الحيوانات لتحديد مستوى الانترليوكينات باستخدام تقنية الاليزا متعدد الطبقات. أظهرت النتائج نقصانا معنويا في مستوى انترليوكين -٥ وانترليوكين-١٢، وزيادة معنوية في مستوى انترفيرون-كاما، في الفئران المعاملة مقارنة بمجاميع السيطرة. بلغ أدنى مستوى ل-انترليوكين-٥، ١١,١٦٤ نانوغرام/مل عند الجرعة ١٠٠ مايكروغرام/مل مقارنة بمجموعة السيطرة ٤٧,٤٤٦ نانوغرام/مل بعد أربعة أشهر بعد الاصابة، بينما بلغ أدنى مستوى لانترليوكين-١٢، ١٢,٨٢٤ نانوغرام/مل، عند الجرعة ١٠٠ ميكروغرام/مل، بالمقارنة مع ١٢,٤٨٥ نانوغرام/مل في مجموعة السيطرة، بعد خمسة أشهر من الاصابة. ازداد مستوى إنترفيرون- كما معنويا في الفئران المعالجة، حيث بلغ ٢٠٧,٣٩ نانوجرام/مل، عند الجرعة ١٠٠ مايكروغرام/مل مقارنة بـ ١١٠,٠٧ نانوغرام/مل في مجموعة السيطرة، بعد خمسة أشهر من الاصابة. استنادا الى نتائج البحث، فان أكسيد الزركونيوم النانوي يمكن أن يستعمل بوصفه معيارا فعالا في تقدير الحالة المناعية، وبالتالي علاجا فعالا لداء الأكياس العدري.