Al-Qadisiyah Journal of Pure Science

Al-Qadisiyah Journal of Pure Science

ISSN(Printed): 1997-2490

ISSN(Online): 2411-3514

DOI: /10.29350/jops.

<http://qu.edu.iq/journalsc/index.php/JOPS>

Study of some histopathological changes occurring in white laboratory mice infected with Cutaneous Leishmaniasis in Al – diwaniyah province, Iraq.

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1. Introduction

Leishmaniasis one of the important diseases for humans, which is caused by the parasite *Leishmania* Which belongs to the order Kinetoplastida family Trypanosomatid. *Leishmania* parasite two forms during its life cycle Promastigote form It is an infectious stage and is seen in females of the sand fly *Phlebotomus* (Carrier host) in the ancient world .And *Lutzomyia* In the new world (18,19). As well as in industrial agricultural communities. Amastigote form It is found in humans and storing hosts and lives inside macrophages in the skin, mucous membranes, lymph nodes, bone marrow and spleen.(10,4). Leishmaniasis is characterized by a variety of clinical features of the disease from Cutaneous Leishmaniasis (CL) Mucocutaneous Leishmaniasis (MCL) Visceral Leishmaniasis (VL) (6,12) . Cutaneous leishmaniasis appears in two types, the first Zoonotic Cutaneous Leishmaniasis (ZCL) It is caused by a parasite L. major And it appears in rural areas and causes a wet sore . and Anthroponotic Cutaneous Leishmaniasis (ACL) When the parasite is transmitted from one person to another and causes it L. tropica This type causes a dry sore (7,9)

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Leishmaniasis represents a major public health problem in the Eastern Mediterranean Region (EMR) of the World Health Organization (WHO) (13) , leishmaniasis causes 70,000 deaths per year. Clinical features depend on the species of *Leishmania* involved and the immune response of the host .Infection of mice with *L. major* is a well-established model for the investigation of factors that control disease development. Most inbred strains of mice, termed resistant or healer strains, develop a local inflammation at the site of *L. major* inoculation (17).

Material and Methods

Samples Collection

Leishmaniasis samples were collected from the edge of ulcers before treatment for patients with cutaneous leishmaniasis, and arrivals to Diwaniyah Teaching Hospital.

Fig (1): Represents some of the skin lesions caused by infection with leishmaniasis dermal parasites.

Diagnosis of Samples

Clinical diagnosis : Clinical diagnosis was made by a dermatologist.

Laboratory diagnosis: The direct smear method prepared from the edge of a pigmented ulcer using Giemsa was used and examined by a high-strength microscope using an oil immersion.(8)

Results

1- Gross Changes

The current study showed the emergence of skin infections in mice experimentally infected with *Leishmania* parasite, which were observed visually in the injection areas a month after the infection. These lesions were accompanied by changes in the behavior of the affected animals, such as lack of movement and lack of desire for food in addition to lethargy.

2- Histological Changes

The first group (control group) showed that normal liver tissue consisted of hepatocytes arranged radially in hexagonal shapes, with the bile duct appearing normal, Fig (2).

Fig (2): A cross section of mouse liver tissue from the control group shows the normal tissue of the liver. With the appearance of the bile duct normal (yellow arrow) (. (H & E, 10X)

The results of the study show that the histological sections of the liver of laboratory mice infected with *L. major*, hepatocytes are irregular in shape as a result of severe fatty degeneration of hepatocytes, which creates vacuoles (expansion of hepatic sinuses) leading to the loss of the radial arrangement of the hepatocytes, with the appearance of heavy infiltration Inflammatory cells, especially macrophages, with Hyperplasia and congestion of the bile duct.fig(3,4)

Fig (3): A cross section of the liver tissue of the affected experiment group observes the disappearance of the radial arrangement between the hepatocytes (green arrow) and the infiltration of inflammatory cells, especially phagocytes (large macrophage) (yellow arrow) with severe fatty degeneration, the nucleus takes a peripheral position (red and blue arrow)). Hepatic sinus enlargement (orange arrow) and biliary enlargement and congestion (black arrow) (H&E, 10x).

Fig (4) a cross-section in the liver tissue of the affected experiment group. The radial arrangement between the hepatocytes (green arrow) and the infiltration of inflammatory cells, especially phagocytic cells (large macrophage) (yellow arrow) with severe steatosis, the nucleus takes a peripheral position (red and blue arrow) . Hepatic sinus enlargement (orange arrow) and bile duct enlargement and congestion (black arrow) (H & E, 40X).

The study of histological sections of skin lesions taken from the ear, foot and tail revealed epidermal ulcerative ulcers accompanied by severe infiltration in the dermis layer mediated by neutrophil cells and polymorphic lymphocytes, with the appearance of severe hemorrhage in the area of the dermis layer, with necrosis in the Epidermal cells for all skin lesions in the ear, foot and tail, Fig (5,6,7).

Fig (5) A cross-section in the ear of the affected experimental group, in which skin ulcers are observed with severe inflammatory infiltration of the dermis layer by neutrophils and polymorphic lymphocytes (red and blue arrow). With severe hemorrhage in the dermis (yellow arrow). Necrosis of the epidermal cells in the ear (black arrow). (H&E,40X) .

Fig (6) A cross section of the foot of the affected experimental group, in which skin ulcers are observed with severe inflammatory infiltration of the dermis layer by neutrophils and polymorphic lymphocytes (red and blue arrow). With severe hemorrhage in the dermis (yellow arrow). Necrosis of the epidermal cells in the ear (black arrow) (H&E, 400X) .

Fig (7) A cross section in the tail of the affected experimental group, in which skin ulcers are observed with severe inflammatory infiltration of the dermis layer by neutrophils and polymorphic lymphocytes (red and blue arrow). With severe hemorrhage in the dermis (yellow arrow). Necrosis of the epidermal cells in the ear (black arrow). (H&E,40X).

Discussion

The results of histopathology of the livers of infected mice showed that the cutaneous leishmaniasis parasite is widespread in the liver and organs close to the surface of the skin, although it appears in the dermal region. In the study of (5) that the parasites *L. amazonensis* spread in the liver of laboratory mice, although the parasites do not reproduce in these organs.

the study of (16) indicated that the pathological effects of the liver are the most prominent with infiltration of inflammatory cells and the appearance of blood congestion and granuloma, and (3) noted the observation of hyperplasia of Koffer cells and infiltration of lymphocytes with enlarged liver cells, and (1) indicated that there are significant tissue changes in Collections of affected mice including hepatocyte enlargement, inflammatory cell accumulation, and lymphocyte infiltration.

Histological sections of the skin lesions visible on the ear, foot and tail regions showed the presence of the parasite in the dermis region, and this is consistent with what was mentioned by (11) that Leishmania parasites are not found inside the epidermis. Heavy infiltration of inflammatory cells such as lymphocytes and macrophage cells was also observed, and this is consistent with what was reported by (15) Abscesses as a result of the breakdown of inflammatory cells, especially neutrophils, due to their short life span compared to other cells, It is possible to explain why neutrophil cell debris produces antibiotic-like substances called defensins that release strong oxidizing substances that kill pathogens (2).

The results of the present study were in agreement with (20) of severe infiltration in the dermis layer mediated by neutrophil and polymorph lymphocytes, with the emergence of severe hemorrhage in the area of the dermis, with Necrosis in the epidermal cells penetrated by inflammatory cells.

And the increase in glutathione levels through NAC treatment improves the BALB/c mice response to infection as analysed using several critical parameters: histopathological outcome of the footpad lesion and cytokine profile of popliteal lymph node cells (14).

Conflict of Interest: The authors declare that they have no conflict of interest.

Acknowledgements:

We are very grateful for the by the University of Al-Qadisiyah. We also deeply thank the locals and persons who helped us during the continuous surveys that were conducted during the study period.

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