Clinical and Laboratory Studies in Correlation to Oxidative Stress Indices in Demodicosis Affected Dogs By

Nafie, T.S.*; Mahmoud M.M.** and Dina A. Abdelkhalek*

*Department of Veterinary medicine (internal Medicine), Faculty of Veterinary Medicine Suez Canal University, Ismailia.

tnafie@hotmail.com

**Department of Veterinary medicine (Infectious diseases), Faculty of Veterinary Medicine Suez Canal University, Ismailia. Mmostafa76@yahoo.com

Corresponding author: dina_amin@vet.suez.edu.eg

Abstract:

Skin diseases are the most common health problem among dogs. Demodicosis is the most important one. The study aimed to detect the clinical effect of demodicosis on dogs which come to veterinary clinic and the correlation between some blood parameters and some oxidative stress indices. Clinical and laboratory examination of the diseased dogs was carried out. The clinical study was conducted on two groups of dogs. The first one includes seven dogs (7.77%) out of 90 diseased dogs of different ages. The second group 14 dogs were clinically healthy used as a control group. Animals were subjected to physical examination followed by special dermatological examination. Skin scrapings were examined under microscope for mite's detection. Blood samples were taken for hematological examination and biochemical analysis. Hematological parameters of demodicosis affected dogs showed a highly significant decrease in hemoglobin values. Regarding differential leukocytes, a significant increase in granulocytes absolute number was recorded. The biochemical findings of blood plasma obtained from dogs affected by demodicosis showed a significant increase in plasma SOD when compared with control ones and a highly significant increase in plasma MDA values. Correlation analysis revealed positive correlation between plasma zinc and copper levels and antioxidant enzymes as SOD, GPx and negative correlation between plasma SOD and plasma MDA levels. The study revealed the importance of oxidative stress

indicators variations in diagnosis and pathophysiology of demodicosis.

Keywords: Dogs, Demodicosis, hematological changes, oxidative stress markers, correlation analysis

Introduction:

Demodicosis is a very common parasitic skin disease in dogs (*Mueller*, 2004), in which a large number of Demodex mites are found in the skin (Gortel, 2006). Mueller et al. (2012) and Beigh et al. (2013) described clinical signs of demodicosis that they are erythema, scaling, partial or complete alopecia, papules. follicular casts, pustules, and in cases. furunculosis. severe crusting, exudation and ulceration with focal draining tracts. Generally, the lesions begin on the face and limbs, but they may become generalized. Moreover, Sivajothi et al (2015) observed similar results with lichenification and cellulitis. The authors added that the clinical distribution of lesions was observed on face, around the eyes and ears, chin region, fore limbs, neck and lateral abdomen. diagnosis of mange The is typically based on clinical signs and is confirmed by the presence of mites in skin scrapings (Carter, 2001).

Deb et al (2000) and Jyotsna et al. (2005) reported that dogs affected with generalized demodicosis revealed significant reduction in total erythrocyte count, hemoglobin and packed cell volume resulting to anemia. Affected dogs also showed leukocytosis accompanied by neutrophilia, eosinophilia and lymphopenia with low absolute lymphocyte count.

Nair æ Nauriyal (2007)recorded lower level of hemoglobin concentration and total erythrocytic count as well as normal level of total leukocvtic count. neutrophil. lymphocyte and monocyte cells with presence of eosinophilia Dadhich (2008) observed in a study on dogs with demodicosis slight decrease in total a erythrocytic count, hemoglobin and packed cell volume and an increase in total leukocytic count and eosinophilia. Similar results were obtained by Sakina et al (2012) who reported that the mean values of Hb. PCV and TEC were significantly lower. however MCV and MCH were significantly higher in dogs suffered from demodectic mange as compared to control group indicated macrocytic anemia in affected groups. The author reported that anemia might be due to the stress arising from the disease. The authors added that there was a significant increase in leukocytic count, neutrophil and eosinophil values.

Portugal et al. (2007) reported that in skin diseases, the bodies antioxidant such as SOD, CAT, GSH, GSH-peroxidase (GPx) act synergistically to cause sequential degradation of peroxides and free radicals to combat oxidative damage. Gurer et al. (1998) reported that the measurement of antioxidant enzymes like SOD and CAT are appropriate indirect ways to assess the status of antioxidant defense and the estimation of Malondialdehyde (MDA)а byproduct of lipid peroxidationcontinues to be a reliable method to assess the degree of oxidative damage to cell membranes. Moreover, Fang et al. (2002) mentioned that estimation of antioxidant enzymes activities and levels of endogenous antioxidants in blood are indirect reliable methods but for assessment of free radicals' activity and oxidative stress. as well. Additionally, Tomsič et al (2016) reported that total antioxidant capacity (TAC) and antioxidant enzymes, GPx and are commonly used SOD. markers of antioxidant status and oxidative stress.

Materials and methods:

the clinical study was conducted on two groups of dogs of different breeds and ages. the first group was fourteen apparently healthy dogs used as control group. the second group was the diseased included seven dogs out of ninety diseased dogs were affected with demodicosis. The study was carried out in the clinic of both faculties of veterinary medicine Suez Canal university, Ismailia, and Cairo university, Egypt.

definitive diagnosis was carried by case history, physical examination, inspection of skin, skin coat, skin scraping. as well as body temperature, pulse, respiratory rate, examination of superficial lymph nodes DIAGNOSIS was carried according to Birchard æ Sherding (2005).

Multiple skin scrapings were taken from the periphery of the lesions or alopecic areas until oozing of blood. After samples collections several drops of 10% sodium hydroxide (NaOH) solution were added then gently heated and examined under light microscope for the identification of mites (*Houston, 2000*).

Anti-coagulated blood samples drained then were sent immediately for hematological analysis for complete blood pictures and separation of plasma. Plasma freeze at -80°c for biochemical analysis. The hematological procedures were carried out using fully automatic blood cell counter.

Zinc, copper, SOD, Catalase, GPX and MDA were estimated using colorimetric test kits (Bio-Diagnostic Company Egypt) according to manufacturer's instructions (Hayakawa, 1961, Ventura and King 1951, Nishikimi et al. 1972, Aebi 1984, Paglia and Valentine 1967 and Ohkawa et al 1979) respectively.

Statistically, the obtained data was analyzed using statistical program of social science (SPSS) for windows, Version 24. Values of measured the parameters were expressed as mean value ± slandered error (S.E) and the difference between means of the two groups was determined by using one tail ttest and the significance was considered at P values < 0.05 or 0.01. Correlations between variables were evaluated using Pearson Correlation Coefficient. The interpretation of Pearson Coefficient correlation according to (Mukaka, 2012).

Results and discussion:

The clinical investigation revealed that demodicosis was diagnosed in seven (7.77%) out of ninety dogs suffered from dermatological problems. These results coincided with Auila (2000) who reported that the prevalence of canine demodicosis was 6.04% out of 281 examined dogs during the study. Similar factors were reported by Deepa et al. (2005) who stated that demodicosis is influenced by numerous factors such as alimentation, presence of stress factors, fitness and other diseases or pathogens. *Shrestha et al (2015)* found that prevalence rate of demodicosis was higher in stray dogs who suffer from poor management, poor body condition and improper nutrition.

Dogs affected by demodicosis showed pale mucus membrane, elevated slightly pulse and respiratory rates. Demodicosis was manifested by diffuse alopecia. excessive scaling. characteristic erythema, offensive odor and severe itching. In most cases the lesions accompanied by secondary pyoderma. Similar signs were reported by Gortel (2006). Mueller et al. (2012), and Beigh et al. (2013) reported similar as signs erythema, scaling. complete alopecia, papules, follicular casts, pustules and in some cases furunculosis, crusting. exudation and ulceration with focal draining Microscopically tracts. examination of skin scrapings revealed presence of *Demodex* canis in samples taken from diseased dogs. These results were similar to those of Carter (2001) and Mueller et al. (2012) who reported that the diagnosis of demodicosis is made by deep skin scraping. Mueller and Bettenay (2010) stated that finding more than one mite is strongly suggestive of clinical demodicosis. Moreover.

Katariya et al (2018) screened all dogs in their study for the presence of mites. The authors recorded the presence of mites in eight dogs.

Hematological parameters of demodicosis affected dogs showed a highly signific ant decrease in hemoglobin values. Similar results were recorded by Deb et al (2000), Nair and Nauriyal (2007), Dadhich, (2008) and Sudhakara et al (2014). Janus et al. (2014) interpreted the decrease in Hb level and PCV values to the deteriorated condition of affected dogs owing to reduced food intake, systemic illness, toxemia and septicemia caused by mites as well as secondary bacterial infection. Similar explanation reported by Pathak and Bhatia (1986). Regarding differential leukocytes, a significant increase in granulocytes absolute number was recorded this results agree with the results obtained by Dadhich (2008), Sakina et al (2012), Reddy et al (2015) and Katariya et al (2018). A highly significant increase in monocyte which coincided with the results obtained by Sudhakara et al (2014). Also, Rebar (1998) who mentioned that eosinophilia and monocytosis are the best indicators of dermatitis in case of demodicosis.

The biochemical findings of blood plasma obtained from

dogs affected by demodicosis showed a significant increase in plasma SOD when compared with control ones. These findings agree with the results obtained by Dimri et al. (2008) Salem et al. (2020) and Kubesy et al (2020). They reported that the increase in plasma SOD activity could be attributed to upregulation in its synthesis to free radicals. counteract Moreover.

Additionally, Singh et al. (2014) and Abdulaziz et al. (2019) mentioned that the reason of the observed elevation in plasma SOD levels in mange affected dogs may be attributed to its role as the first-line of defense antioxidant as it helps the body superoxide to remove the radicals by converting them to hvdrogen peroxide (H2O2). However. some authors suggested that the increase in the antioxidant enzymes is not necessarily desirable. as the antioxidant enzymes are not alwavs decreased in some disease conditions as reported by Russo and Bracarense (2016). The study revealed a highly significant increase in plasma MDA values. These findings were supported by Gurer et al. (**1998**) who reported that estimation of plasma malondialdehyde considered to be a reliable method to assess the degree of peroxidative damage of cell membrane, as it is the

most abundant aldehyde formed as a by-product during this process. Similarly, Cini et al. (1994) and Gurer et al. (1998) reported that estimation

plasma Malondialdehv de (MDA), a byproduct of lipid peroxidation, continues to be a reliable method to assess the degree of oxidative damage to cell membranes. The obtained findings were similar to that obtained by Beigh et al. (2013), Kubesy et al (2020) and Salem et al (2020) who observed increased plasma MDA levels in their study on demodicosis of dogs. The authors added that increased plasma MDA levels increased reflects membrane peroxidation. providing lipid evidence of enhanced free radicle generation in generalized demodicosis. Beigh et al. (2016) attributed the increase of plasma MDA to the oxidative stress interceded by pro-inflammatory during cytokine liberation external parasitic infestation.

of

Demodicosis affected dogs moderate 3) revealed (table correlation positive between plasma zinc level and plasma Cu level and a moderate positive correlation between plasma Cu level and plasma SOD activity. Beigh et al (2013) recorded a strong positive correlation between SOD activity and Cu in both healthy and diseased dogs. The author added that a positive correlation between Zn and SOD

in both healthy and diseased These results were dogs. explained by AL-Qudah et al. (2011) who reported that trace elements are required for the activity of a number of enzymes, including antioxidant enzymes. Ewans and Halliwell (2001) stated that copper along with zinc are essential components of the body's antioxidant defense. Additionally. and Ighodaro Akinlove (2018) mentioned that SOD is a metalloenzyme and hence, requires a metal cofactor for its activity as iron, Zn and Cu. The correlation analysis in the present study revealed a high correlation between negative plasma MDA activity and plasma SOD activity in dogs affected with demodicosis. Similar results were obtained by Beigh et al (2013) who recorded that MDA levels were strongly negatively correlated with SOD activity in diseased dogs. Rice-Evans and Burdon (1994) mentioned that antioxidants function to delay or prevent ROS-induced cellular damage, and work by reducing local concentrations. oxygen impairing chain initiation reactions. However, Cini et al. (1994) reported that malondialde hyde (MDA) is a byproduct of lipid peroxidation that resulting in oxidative damage to cell membranes. The previous authors as well as the recorded results of table 3

confirmed the negative correlation between antioxidant enzymes such as (SOD, Catalase and GPx) and MDA as a byproduct of lipid peroxidation.

Table (1): Blood Parameters of Demodicosis Affected DogsCompared with Clinically Healthy

Parameter	Control (n=14) (Me an ±SE)	Demodicosis (n=7) (Mean ±SE)		
Hb (g/dl)	14.86 ±0.53	11.73**±0.91		
PCV (%)	39.00±1.34	34.73±2.27		
RBCs (10^6/µL)	5.98±0.25	5.28±0.23		
MCV (fL)	67.28±2.25	64.20±2.02		
MCHC (%)	38.15±2.31	33.50±1.03		
WBCs (10^3/µL)	9.72±0.86	13.61±1.94		
Granulocyte /cubic mm	1.10±0.01	2.6*±0.07		
Lymphocyte /cubic mm	7.01±0.05	9.06±0.11		
Monocyte /cubic mm	0.65±0.01	13.60**±1.85		
*Significance p (0.05)	** Highly significance p (0.01)			

Table (2): Biochemical Findings of Demodicosis Affected DogsCompared with Clinically Healthy Dogs.

Parame te r	Control (n=14) (Mean ±SE)	Demodicosis (n=7) (Mean ±SE)	
Zn (µ/ml)	0.822±0.040	0.786±0.052	
Cu (µmole/ml)	0.087±0.012	0.079±0.015	
Catalase (U/ml)	0.289±0.020	0.364±0.039	
SOD (U/ml)	2.280±0.170	2.770*±0.124	
GPx (u/ml)	0.212±0.014	0.225±0.022	
MDA (nmole/L)	0.212±0.014	0.225**±0.022	

^{*}*Significance* p (0.05 ***Highly* significance p (0.01)

	Zn	Cu	CAT	SOD	GPX	MDA
Zn	1	0.670 ^c	-0.112	0.382	-0.012	0.120
Cu	0.670°	1	-0.406	0.568°	0.040	0.408
CAT	-0.112	-0.406	1	0.277	-0.772 ^b	-0.002
SOD	0.382	0.568°	0.277	1	-0.266	-0.809 ^b
GPx	-0.012	-0.040	-0.772 ^b	-0.266	1	-0.009
MDA	0.120	-0.408	-0.002	-0.809 ^b	-0.009	1

 Table (3) Correlation Between Trace Elements and Antioxidants of Dogs Affected with Demodicosis.

1. Very high positive/negative correlation (0.9 to 1.0)/(-1.0to-0.9)

2. *High positive/negative correlation* (0.7to0.9)/(-0.9to-0.7)

3. Moderate positive/negative correlation (0.5 to 0.7)/(-0.7 to -0.5)

References:

Abdulaziz AR, Almuzaini AA, Hassan AA (2019). Evaluation of the antioxidative activity and trace elements concentrations in *Demodex canis* infected dogs. Peer Res Nest 1(2):1–6.

Aebi H (1984). Catalasein vitro. In: Colowick SP, Kaplan NO (eds) Methods in enzymology, vol 105. Academic Press, New York, pp 121–126.

AL-Qudah, K.M., Gharaibeh, AL-Shyvab, M.M. A.A., (2011). Trace minerals status antioxidant and enzymes activities in calves with dermatophytosis. Biol. Trace Elem. Res. 136, 40–47.

Aujla, R. S., Singla, L. D., Juval, P. D., & Gupta, P. P. (2000). Prevalence and pathology of mange-mite infestations in dogs. Journal of Veterinary Parasitology, 14(1), 45-49.

Beigh, S. A., Soodan, J. S., Singh, R., & Khan, A. M. (2013). Trace minerals status and antioxidative enzyme activity in dogs with generalized demodecosis. Veterinary parasitology, 198(1-2), 180-186.

Beigh S.A., Soodan J.S. and Bhat A.M. (2016). Sarcoptic mange in dogs: its effect on liver, oxidative stress, trace. Vet Parasitol 227:30–34.

Birchard, S. J., & Sherding, R. G. (2005). Saunders Manual of Small Animal Practice-E-Book. Elsevier Health Sciences.

Carter G.R. (2001). External parasitic diseases of dogs and

cats. In: A concise guide to infectious diseases of dogs and cats (Ed: Carter GR), International Veterinary Information Service, Ithaca, NY.

Cini, M., Fariello, R.Y., Bianchetti, A. and Moretti, A. (1994): Studies on lipidperoxidation in the rat brain. Neurochem. Res. 19, 283–288.

Dadhich,H.(2008).Pathological,haemato-biochemicaland immunologicalstudiesofcutaneousectoparasitosesindogs.measurement, 40, 58-65.

Deb, A. R., Jha, M. K., & Prasad, K. D. (2000). Clinical and haematological changes in dogs infected with *Demodex* canis. Journal of Research, Birsa Agricultural University, 12(2), 281-283.

Deepa, L.; Kalita, D.N. and Neog, R. (2005). Epizootiology of demodectic mange in dogs. Indian Vet. J., 82:501-502.

Dimri, U., Ranjan, R., Kumar, N., Sharma, M.C., Swarup, D., Sharma, B., Kataria, M. (2008). Changes in oxidative stress indices, zinc and copper concentrations in blood in canine demodicosis. Vet. Parasitol. 154 (1–2), 98–102.

Ewans, P. and Halliwell, B. (2001). Micronutrients: oxidant/antioxidant status.Br. J. Nutr. 85, 57–74. Fang, Y.-Z., Yang, S. and Wu, G. (2002). Free radicals, antioxidants and nutrition. Nutrition 18, 872–879.

Gortel, K. (2006). Update on canine demodicosis. Veterinary Clinics of North America: Small Animal Practice.36, 229–241.

Gurer, H., Ozgunes, H., Neal, R., Spitz, D.R. and Ercal, N. (1998). Antioxidant effects of Nacetyl cysteine and succimer in red blood cells from lead exposed rats. Toxicology 128, 181–189.

Hayakawa, R. (1961). Determination of zinc. *Jap J Toxic Environ Health*, 8, 14-18.

Houston, D. M. (2000). Clinical examination of dogs and cats. In: Radostits, O.M.; Mayhew, I.G.; and Houston, D.M.: Veterinary Clinical Examination and Diagnosis. W.B. Saunders, China, pp., 125-138.

Ighodaro, O. M., & Akinlove, O. A. (2018). First line defense antioxidants-superoxide dismutase (SOD), catalase (CAT) and glutathione peroxidase (GPX): Their fundamental role in the entire antioxidant defense grid. Alexandria Journal of Medicine, 54(4), 287-293.

Janus, A.; Tresamol, P. V.; Mercev, K. A.; Biju P. H. and Shameem H. (2014). Study on clinical and hematological parameters in canine demodicosis. Ind. J. of Canine Pract. Vol. 6.

Jyotsna, P., Maiti, S. K., Sanyal, P. K., & Tiwari, S. P. (2005). Haematobiochemical and mineral profiles in generalized canine demodicosis. Intas Polivet, 6(2), 331-334.

Katariya, A., Arora, N., Rajora, V. S., Ilyas, W., Mrigesh, M., Das, A. K., & Prasad, A. (2018). Hematobiochemical alterations in ectoparasite infestation in canines.

Kubesy, A. A. M., Yehia, S. G., Salem, S. I., & Rabah, M. (2020). Altered blood oxidative stress markers in association with antioxidant supplemented therapy for mange, tick, and flea allergic dermatitis of dogs. Comparative Clinical Pathology, 1-7.

Mueller RS (2004). Treatment Protocols for Demodicosis: An Evidence-Based Review. Veterinary Dermatology 2004; 15(2) 75-89.

Mueller, R.S. and Bettenay, S.V. (2010). Skin scrapings and skin biopsies. In: Ettinger SJ, Feldman EC, eds. Textbook of Veterinary Internal Medicine. Philadelphia, W.B. Saunders; 368–371.

Mueller, R.S.; Bensignor, E.; Ferrer, L.; Holm, B.; Lemarie, S.; Paradis, M. and Shipstone, M.A. (2012). Treatment of demodicosis in dogs: Clinical Practice Guidelines.Vet. Dermatol., 23(2): 86-96.

Mukaka M. M. (2012). Statistics corner: A guide to appropriate use of correlation coefficient in medical research. Malawi medical journal: the journal of Medical Association of Malawi, 24(3), 69-71.

Nair, S. S., & Naurival, D. S. (2007). Diagnostic significance of haematological changes associated with various canine dermatoses. Intas polivet, 8(1), 68-72.

Nishikimi M. Roa NA and Yagi K (1972). The occurrence of superoxide anion in the reaction of reduced phenazinemethosulfate and molecular oxygen. Biochem Biophys Res Commun 46:849– 854.

Ohkawa H, N Ohishi and K Yagi (1979). Assay of lipid peroxides in animal tissues by thiobarbituric acid reaction, Annals of Biochemistry. (95):351-358.

Paglia, D.E. and Valentine, W. N. (1967). Studies on the quantitative and qualitative characterization of erythrocyte glutathione peroxidase. J lab Clin Med.70:158-169. Pathak, K. M. L. and Bhatia, B. B. (1986). Haematobiochemical and pathological changes in a dog with generalized demodecosis. Ind. J. Vet. Med., 6(1): 26-28.

M., Barak, V., Portugal, Ginsburg, I. and Kohen, R. (2007). Interplay among oxiantioxidants. and dants. cytokines skin disorders: in present statusand future considerations. Biomed. Pharmacother. 61, 412–422.

Rebar, A. H.

(**1998**). Hemogram interpretation for dogs and cats. Ralston Purina International.

Reddv. B. S., Kumari, K. N., & Sivajothi, S. (2015). Haematobiochemical findings and thyroxin levels in canine demodicosis. Comparative Clinical Pathology, 24(2), 287-290.

Rice-Evans CA and Burdon RH (1994). *Free Radical Damage and its Control.* Amsterdam: Elsevier Science B.V., 1994, pp. 113–114.

Russo C and Bracarense A (2016). Oxidative stress in dogs. Semina Ciências Agrárias Londrina 37(3):1431–1440.

Sakina, A., Mandial, R. K., & Mudasir, Q. (2012). Haematobiochemical changes in canine demodecosis. Vet Scan, 7, 75-78. Salem, N. Y., Abdel-Saeed, H., Farag, H. S., & Ghandour, R. A. (2020). Canine demodicosis: Hematological and biochemical alterations. Veterinary World, 13(1), 68.

Shrestha, D., Thapa, B., Rawal, G., Dhakal, S., & Sharma, B. (2015). Prevalence of demodectic mange in canines of Kathmandu Valley having skin disorder and its associated risk factors. International Journal of Applied Sciences and Biotechnology, 3(3), 459-463.

Singh Z., Karthigesu I.P., Singh P. and Kaur R. (2014). Use of malondialdehyde as a biomarker for assessing oxidative stress in different disease pathologies: a review. Iran J Publ Health 43(3):7–16.

Sivajothi, S., Reddy, B. S., & Rayulu, V. C. (2015). Demodicosis caused by *Demodex canis* and Demodex cornei in dogs. Journal of parasitic diseases, 39(4), 673-676.

Sudhakara, B. R.; Nalini, K. K.: Sivajothi, S. and Venkatasivakumar, R. (2014). Dermatitis due to mixed Demodex and Sarcoptes mites in dogs. Case reports in Vet. Med. Vol. http://dx.doi.org/10.1155/2014/7 68242.

Tomsič, K., Seliškar, A., Lukanc, B., & Svete, A. N. (2016). Plasma total antioxidant capacity and activities of blood glutathione peroxidase and superoxide dismutase determined in healthy dogs by using commercially available kits. Acta Veterinaria, 66(4), 534-548.

Ventura S. and King, E.J. (1951). Biochem. J. 48 IXI= Eden A. Green HH. (1940). Biochem. J. 34: 1202.

دراسات اكلينيكية ومعملية مع الارتباط بمؤشرات الإجهاد التأكسدي في داء الجرب في الكلاب ثروت سالم نافع، محمد مصطفى محمود، دينا امين عبد الخالق محمد

الملخص العربى

الأمر اض الجلدية هي أكثر المشاكل الصحية شبوعًا بين الكلاب. داء الجرب من اهمها. هدفت الدر اسة إلى الكشف عن التأثير الاكلينيكي للجرب على الكلاب التي تأتي إلى العبادة اليبطرية. والعلاقة بين بعض مقاييس الدم وبعض مؤشرات الإجهاد التأكسدي. تم إجراء الفحص الإكلينيكي والمعملي للكلاب المريضة. تم أخذ عينات مختلفة من الحيو إنات المريضة لتحديد مشكلة المرض بدقة. تحديد بعض مؤشر ات الإجهاد التأكسدي للكشف عن تأثير الجرب على مستوياته في بلاز ما الدم. أجريت الدر اسة الاكلينيكية على مجمو عتين من الكلاب. الأول يضم ٧ كلاب (٧,٧٧٪) من بين ٩٠ كلباً مريضة من مختلف الأعمار . المجموعة الثانية ٢٤ كلبًا كانت صحية سريريًا استخدمت كمجموعة ضابطة. تم إخضاع الحيوانات للفحص الجسدي تلاه فحص جلدي خاص. تم فحص قشور الجلد تحت المجهر للكَشف عن الديمو ديكس. تم أخذ عينات الدم لفحص الدم والتحليل الكيميائي الحيوي للزنك والنحاس وعلامات الإجهاد التأكسدي بما في ذلك الكاتلاز ، GPx ، SOD و MDA. أظهرت صورة الدم للكلاب المصابة داء الجرب انخفاضًا معنويًا للغاية في قيم الهيموجلوبين. فيما يتعلق بالكريات البيض التفاضلية تم تسجيل زيادة كبيرة في العدد المطَّلق للخلايا المحببة. أظهرت النتائج البيوكيميائية لبلازما الدم المأخوذة من الكلاب المصابة بداءالجرب زيادة معنوية في SOD في البلازما عند مقارنتها مع مجموعة التحكم وزيادة معنوية عالية في قيم MDA في البلازما. كشف تحليل الارتباط عن وجود علاقة أرتباط موجبة بين مستويَّات الزنك في البَّلازما ومستويات النحاس وأنز يمات مضادات الأكسدة مثل SOD و GPx و العلاقة السلبية بين مستويات SOD في البلازما ومستويات MDA في البلازما. كشفت الدراسة عن أهمية تباين مؤشرات الإجهاد التأكسدي في التشخيص.