Assessment of the level of interleukin-10 in patients with COVID-19

Qusay M. Saleh¹, Nihad A. Jafar², Sabah M. Salih³

¹University of Tikrit / College of Veterinary Medicine / Microbiology Department ²University of Tikrit / Collage of Veterinary Medicine ³University of Kirkuk / Collage of Pharmacy Prof.dr..al.aubaidi@gmail.com

Abstract

The current study aimed to assess the level of Interleukin-10 among Covid-19 patients by using the ELISA technique. For this purpose, (100) serum samples were collected from patients infected with Covid-19 disease in the city of Kirkuk for the period from February 20, 2021 to July 8, 2021, with ages ranging from 22 to 93 years, fifty males and fifty females. The patients were divided according to the severity of the disease into three groups, the first group had mild symptoms (19%), the second group had moderate severity (41%), and the third group was severe (40%). The results of the study showed a significant increase in the levels of interleukin-10 (73 ± 8.2 pg/ml) among Covid-19 patients compared to the control group (P.Valeu ≤ 0.05). Interleukin-10 also recorded the highest level in severe cases (121.1 ± 3.11 pg/ml) compared to moderate and mild cases (P.Valeu ≤ 0.05). Also, the level of interleukin-10 increased significantly among Covid-19 patients with diabetes, and the highest increase was recorded in severe cases, where it was (142.9 ± 7.27 pg/ml, P.Valeu ≤ 0.05).

Keywords

COVID-19, Corona virus, interleukin-10

I. INTRODUCTION

Coronavirus disease 2019 (COVID-19) is caused by a new coronavirus first identified in Wuhan, China, in December 2019 ⁽¹⁾. Corona virus belongs to the genus Beta Corona, under the family Coronaviridae, which is a group of RNA viruses that cause diseases in mammals and birds ^(2,3). It was first discovered in 1965 by Terrell and Benoy and the first virus to be discovered was the bronchitis virus of an adult with a cold ⁽⁴⁾. Corona viruses are mainly associated with respiratory diseases, in humans it primarily causes respiratory infections that can range from mild to fatal. Mild illnesses in humans include some cases of the common cold (also caused by other viruses, mostly rhinoviruses), while more deadly types can cause SARS, MERS, and Novel Coronavirus ^(2,5). After the emergence of the coronavirus that

causes severe acute respiratory syndrome (SARS-CoV) and the coronavirus that causes Middle East respiratory syndrome (MERS-CoV), the emerging coronavirus (SARS-CoV-2) is the seventh member of the coronavirus family to be a new member of the The genus Beta Corona virus, which results in the disease Covid-19 that infects humans ^(6,7). The common symptoms of patients were fever, shortness of breath, cough, phlegm, headache, muscle pain, chest pain, diarrhea, and pharyngitis ⁽⁸⁾.

The name Corona virus is derived from the Latin word Corona, which means crown ⁽⁹⁾. The new virus was named severe acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) by the International Committee on Classification of Viruses (ICTV) as the name for the new virus on February 11, 2020. This name was chosen because the virus is genetically related to the coronavirus responsible for the SARS outbreak 2003. The disease it causes was named Coronavirus Disease 2019 (Covid-19) by the World Health Organization (WHO) in the International Classification of Diseases (ICD) as the name for this new disease on February 11, 2020, following the guidelines previously established with The World Organization for Animal Health (OIE) and the Food and Agriculture Organization of the United Nations (FAO) ⁽¹⁰⁾.

The innate immune response is the first step in the defense mechanism against the emerging corona virus, as host cell receptors recognize the virus's RNA and initiate a range of immune and inflammatory responses including the production of cytokines, which in turn recruit immune cells such as macrophages, neutrophils, T cells and B cells to The site of infection to increase the fight against the virus. Anti-inflammatory cytokines such as interleukin-10 are produced during the course of infection to regulate inflammation and maintain immune homeostasis ⁽¹¹⁾. Interleukin-10 is a pleotrophic, anti-inflammatory cytokine, originally identified as being produced by helper T cells (Th2) and is now known to be produced by many myeloid and lymphoid-derived immune cells involved in both Innate and adaptive immunity ⁽¹²⁾, This includes macrophages such as cardiac macrophages, microglia, natural killer cells, mast cells, epithelial cells, monocytes, neutrophils, and eosinophils, as well as T cells and B cells ^(13,14). The primary function of interleukin-10 during infection is to inhibit the host's immune response to avoid excessive immune activation during infection with a pathogen (viral, bacterial, and fungal), thereby mitigating tissue damage and reducing autoimmune damage, by inhibiting the production of pro-inflammatory cytokines. It also limits the excessive activation and proliferation of T cells through its interaction with a specific receptor that is highly expressed on the membrane of monocytes, macrophages, and

T cells ^(12,15). One of the main causes of death in COVID patients is the overproduction of pro-inflammatory cytokines resulting from excessive activation of immune cells, referred to as a 'cytokine storm'. Therefore, an early elevation of interleukin-10, which acts as an anti-inflammatory, is a hallmark of excessive inflammation during severe infection, as several studies indicate that elevation in the level of interleukin-10 is evidence of poor outcome in patients infected with the virus based on its role as an anti-inflammatory cytokine and an inhibitor of cytokines. Pro-inflammatory ⁽¹²⁾. This was observed in patients infected with the virus, as they showed a severe increase in the level of interleukin 10 ⁽¹⁶⁾.

I. MATERIALS AND METHODS

110 serum samples were collected from the city of Kirkuk for the period from February 20, 2021, to July 8, 2021, their ages ranged between 22 to 93 years, 100 samples of patients infected with COVID-19 were 50 males and 50 females. And 10 samples were from healthy subjects, representing the control group, were 5 males and 5 females. The samples recorded in this study were divided into three groups, mild symptoms (19), moderate symptoms (41), severe symptoms (40), the serum sample used to assess the level of interleukin-10 in Covid-19 patients by using the ELISA technic, by use the kit manufactured by the chines Shanghai Biological Company, to assess the level of interleukin-10, The same steps were used according to the leaflet attached to the test kit. The results were significant at the probability value (P. value ≤ 0.05).

I. RESULT AND DISCUSSION

The results of the statistical analysis showed a significant increase in the mean of IL-10 in COVID-19 patients when compared with the control group, where it was (61.4 pg/ml, 7.03pg/ml, respectively, P. value ≤ 0.05) as shown in Table (1).

Standards	patients	control group	P. value
IL-10	73 ± 8.2 pg/ml	$14.78 \pm 1.4 \text{ pg/ml}$	0.0004

Table (1): Mean level of IL-10 in covid-19 patients and control group

Cytokines play an important role in eliminating pneumonia-causing pathogens that can be synthesized by many other immune or non-immune cells. However, if their level becomes too high, tissue damage and host death can occur. Therefore, the presence and quantity of cytokines can give a clear indication of the understanding of the pathogen and immune response of the host ⁽¹⁷⁾. This increase indicates the effect of the pneumonia in the patient group on the level of cytokines, this inflammatory response can become detrimental if it is excessive or not sufficiently balanced by anti-inflammatory agents ⁽¹⁸⁾. The results of the current study agreed with what researchers found in London, where a study in London found that patients with Covid-19 disease show elevated levels of interleukin-10 ⁽¹⁹⁾, and this matches what was reported by an Egyptian clinical study in individuals with Covid-19 disease about High levels of anti-inflammatory cytokines, including interleukin-10 ⁽²⁰⁾.

The results of the statistical analysis also showed significant increase in the meano of IL-10 in COVID-19 patients according to the severity of the disease, where the average in the following cases (severe symptoms, moderate symptoms, mild symptoms) were as follows (121.1 pg/ml, 50.3 pg/ml, 20.5 pg/ml, respectively, p value ≤ 0.05) as shown in Table (2).

Table (2): Mean level of IL-10 in covi	patients according	g to severity of disease
--	--------------------	--------------------------

Standards	mild symptoms	moderate symptoms	severe symptoms	P. value
IL-10	$20.5\pm0.98~pg/ml$	50.3 ± 2.94 pg/ml	121.1 ± 3.11 pg/ml	0.0007

The results of the current study are consistent with those in Wuhan, China, the first descriptive primary study evaluating the immunological characteristics of patients with laboratory confirmed COVID infection, where the study found that serum concentrations of IL-10 in most Severe cases were significantly higher than those in moderate and mild cases, and the cytokine storm corresponded to disease severity ⁽²¹⁾, and researchers in Italy also found that in severe cases of COVID patients had high levels of IL-10 ⁽²²⁾, the present findings are also consistent with a study in India that found that COVID patients were associated with moderate increases in a wide range of cytokines including IL-10 that were significantly associated with disease severity ⁽²³⁾.

The results of the statistical analysis also showed a significant increase in the mean of interleukin-10 in Covid-19 patients with diabetes, (severe symptoms, moderate symptoms, mild symptoms) were as follows (142.9 pg/ml, 61.59 pg/ml, 22.89 pg/ml, respectively, p value ≤ 0.05) compared with patients without diabetes (110.7 pg/ml, 46.65 pg/ml, 19.38 pg/ml, respectively, p value ≤ 0.05) as shown in Table (3).

Standards	severity of disease	with diabetes	Without diabetes	P. value
IL-10	mild	$22.89 \pm 1.63 \text{ pg/ml}$	$19.38\pm0.82\ pg/ml$	0.0004
	moderate	$61.59\pm5.09~pg/ml$	$46.65\pm1.94~pg/ml$	0.0004
	severe	$142.9 \pm 7.27 \text{ pg/ml}$	$110.7\pm4.93~\text{pg/ml}$	0.0004

Table (3): Mean level of IL-10 in covid patients with and without diabetes

Studies have shown that people with diabetes have a dysregulated innate and adaptive immune response and have chronic low-grade inflammation making them more susceptible to a cytokine storm ⁽²⁴⁾, A study conducted in Italy also agreed with the results of the current study, which revealed that the average interleukin-10 increased significantly in Covid-19 patients with diabetes when compared to patients without diabetes, where high blood sugar was associated with a high level of interleukin-10, which indicates the presence of inflammatory signs, which increases the production of interleukin 10 ⁽²⁵⁾, so, when infected with the Corona virus, this pre-existing inflammation in diabetics can be exacerbated, predisposing to hyperinflammatory syndrome or cytokine storm, which is the most common factor. significance in disease progression ⁽²⁶⁾, so it has been suggested that COVID patients with diabetes are more susceptible to an 'inflammatory storm', which is associated with rapid deterioration and higher mortality risk ⁽²⁷⁾.

Reference

- World Health Organization, 2020. Clinical management of severe acute respiratory infection (SARI) when COVID-19 disease is suspected: interim guidance, 13 March 2020 (No. WHO/2019-nCoV/clinical/2020.4). World Health Organization.
- Zhu, N., Zhang, D., Wang, W., Li, X., Yang, B., Song, J., Zhao, X., Huang, B., Shi, W., Lu, R. and Niu, P., 2020. A novel coronavirus from patients with pneumonia in China, 2019. New England journal of medicine.
- Lavi, E. and Weiss, S.R., 1989. Coronaviruses. In Clinical and Molecular Aspects of Neurotropic Virus Infection (pp. 101-139). Springer, Boston, MA.
- 4. Kahn, J.S. and McIntosh, K., 2005. History and recent advances in coronavirus discovery. The Pediatric infectious disease journal, 24(11), pp.S223-S227.
- 5. Brian, D.A. and Baric, R.S., 2005. Coronavirus genome structure and replication. Coronavirus replication and reverse genetics, pp.1-30.

- Yang, P. and Wang, X., 2020. COVID-19: a new challenge for human beings. Cellular & molecular immunology, 17(5), pp.555-557.
- Gao, Y., Yan, L., Huang, Y., Liu, F., Zhao, Y., Cao, L., Wang, T., Sun, Q., Ming, Z., Zhang, L. and Ge, J., 2020. Structure of the RNA-dependent RNA polymerase from COVID-19 virus. Science, 368(6492), pp.779-782.
- Wu, Z. and McGoogan, J.M., 2020. Characteristics of and important lessons from the coronavirus disease 2019 (COVID-19) outbreak in China: summary of a report of 72 314 cases from the Chinese Center for Disease Control and Prevention. Jama, 323(13), pp.1239-1242.
- 9. Singhal, T., 2020. A review of coronavirus disease-2019 (COVID-19). The indian journal of pediatrics, 87(4), pp.281-286.
- 10.World Health Organization, 2020. Naming the coronavirus disease (COVID-19) and the virus that causes it. Brazilian Journal of Implantology and Health Sciences, 2(3).
- 11.Dhar, S.K., Vishnupriyan, K., Damodar, S., Gujar, S. and Das, M., 2021. IL-6 and IL-10 as predictors of disease severity in COVID-19 patients: results from meta-analysis and regression. Heliyon, 7(2), p.e06155.
- 12.Islam, H., Chamberlain, T.C., Mui, A.L. and Little, J.P., 2021. Elevated interleukin-10 levels in COVID-19: potentiation of pro-inflammatory responses or impaired antiinflammatory action?. Frontiers in Immunology, 12, p.2485.
- 13.Silva, D.P.L.J., Carriche, G.M., Saraiva, M., Castro, A.G. and Roque, S., 2016. Balancing the immune response in the brain: IL-10 and its regulation.
- 14.Hulsmans, M., Sager, H.B., Roh, J.D., Valero-Muñoz, M., Houstis, N.E., Iwamoto, Y., Sun, Y., Wilson, R.M., Wojtkiewicz, G., Tricot, B. and Osborne, M.T., 2018. Cardiac macrophages promote diastolic dysfunction. Journal of Experimental Medicine, 215(2), pp.423-440.
- 15.Ni, G., Chen, S., Yuan, J., Cavezza, S.F., Wei, M.Q., Li, H., Pan, X., Liu, X. and Wang, T., 2019. Comparative proteomic study reveals the enhanced immune response with the blockade of interleukin 10 with anti-IL-10 and anti-IL-10 receptor antibodies in human U937 cells. Plos one, 14(3), p.e0213813.
- 16.Sun, X., Wang, T., Cai, D., Hu, Z., Liao, H., Zhi, L., Wei, H., Zhang, Z., Qiu, Y., Wang, J. and Wang, A., 2020. Cytokine storm intervention in the early stages of COVID-19 pneumonia. Cytokine & growth factor reviews, 53, pp.38-42.

- 17.Gómez-Laguna, J., Salguero, F.J., Pallarés, F.J., De Marco, M.F., Barranco, I., Cerón, J.J., Martínez-Subiela, S., Van Reeth, K. and Carrasco, L., 2010. Acute phase response in porcine reproductive and respiratory syndrome virus infection. Comparative immunology, microbiology and infectious diseases, 33(6), pp.e51-e58.
- 18.Khattab, A.A., El-Mekkawy, M.S., Shehata, A.M. and Whdan, N.A., 2018. Clinical study of serum interleukin-6 in children with community-acquired pneumonia. Egyptian Pediatric Association Gazette, 66(2), pp.43-48.
- 19.Chan, J.F., Chan, K.H., Kao, R.Y., To, K.K., Zheng, B.J., Li, C.P., Li, P.T., Dai, J., Mok, F.K., Chen, H. and Hayden, F.G., 2013. Broad-spectrum antivirals for the emerging Middle East respiratory syndrome coronavirus. Journal of Infection, 67(6), pp.606-616.
- 20.Ragab, D. et al. (2020) 'The COVID-19 Cytokine Storm; What We Know So Far', Frontiers in Immunology, 11(June), pp. 1–4. doi: 10.3389/fimmu.2020.01446.
- 21.Chen, G. et al. (2020) 'Clinical and immunological features of severe and moderate coronavirus disease 2019', Journal of Clinical Investigation, 130(5), pp. 2620–2629. doi: 10.1172/JCI137244.
- 22.Cristina, S., Concetta, R., Francesco, R. and Annalisa, C., 2020. SARS-Cov-2 infection: Response of human immune system and possible implications for the rapid test and treatment. International immunopharmacology, 84, p.106519.
- 23.Dhar, S.K., Vishnupriyan, K., Damodar, S., Gujar, S. and Das, M., 2021. IL-6 and IL-10 as predictors of disease severity in COVID-19 patients: results from meta-analysis and regression. Heliyon, 7(2), p.e06155.
- 24. Apicella, M., Campopiano, M.C., Mantuano, M., Mazoni, L., Coppelli, A. and Del Prato, S., 2020. COVID-19 in people with diabetes: understanding the reasons for worse outcomes. The lancet Diabetes & endocrinology.
- 25.Sardu, C., D'Onofrio, N., Balestrieri, M.L., Barbieri, M., Rizzo, M.R., Messina, V., Maggi, P., Coppola, N., Paolisso, G. and Marfella, R., 2020. Outcomes in patients with hyperglycemia affected by COVID-19: can we do more on glycemic control?. Diabetes care, 43(7), pp.1408-1415.
- 26.Erener, S. (2020) 'Diabetes, infection risk and COVID-19', Molecular Metabolism. Elsevier GmbH, 39(June), p. 101044. doi: 10.1016/j.molmet.2020.101044.
- 27. Acharya, D., Lee, K., Lee, D.S., Lee, Y.S. and Moon, S.S., 2020, September. Mortality rate and predictors of mortality in hospitalized COVID-19 patients with diabetes. In Healthcare (Vol. 8, No. 3, p. 338). Multidisciplinary Digital Publishing Institute.