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DERLEME
REVIEW

The effect of tocilizumab on severe COVID-19 infection: Review of current evidence

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ABSTRACT

The effect of tocilizumab on severe COVID-19 infection: Review of current evidence

The COVID-19 outbreak that spread in December 2019 has caused the death of millions of people in a short time. Many studies published recently have shown that many cytokines (interleukin (IL) IL-1, IL-2, IL-6, TNF and IFN-) are significantly increased in COVID-19 patients with pneumonia, and especially IL-6 in combination with other cytokines has shown to be the main cause of the cytokine storm. Since IL-6 level is associated with clinical worsening in COVID-19 patients, anti-IL-6 therapy is seen as a promising treatment. Tocilizumab, a widely used IL-6 antagonist, was approved by the FDA in 2017 for Cytokine Storm Syndrome (CSS). Its addition to the treatment in COVID-19 patients with increased blood IL-6 levels and oxygen saturation <92% has been recommended due to bilateral widespread pulmonary infiltration. We summarized the studies on tocilizumab and COVID-19 disease published in the literature. In the light of current information, data on the mechanism of the action of tocilizumab, which is still widely used in COVID-19 patients, appropriate patient selection, effectiveness in treatment and side effects are presented.

Key words: COVID-19; cytokine storm syndrome; interleukin-6; interleukin-6 antagonist; Tocilizumab

ÖZ

Ağır COVID-19 enfeksiyonunda tosilizumab'ın etkinliği: Güncel çalışmaların derlemesi

Aralık 2019'da başlayan COVID-19 salgını, kısa sürede milyonlarca insanın ölümüne yol açmıştır. Yakın zamanda yayınlanan çalışmalar, pnömoni ile seyreden COVID-19 hastalarında birçok sitokin (IL-1, IL-2, IL-6, TNF ve IFN- γ) önemli ölçüde arttığını ve özellikle Interlökin 6 (IL-6)'nın diğer sitokinlerle birlikte sitokin fırtınasının ana nedeni olduğunu göstermiştir. IL-6 düzeyi, COVID-19 hastalarında klinik kötüleşme ile ilişki bulunduğu için, anti-IL 6 tedavisi umut vaat eden tedavi olarak görülmektedir. Yaygın kullanılan IL-6 antagonisti olan Tosilizumab'ın 2017 yılında FDA tarafından Sitokin Fırtınası Sendromu için kullanımı onaylanmıştır. Kanda IL-6 düzeyi yükselmiş, akciğerde bilateral yaygın infiltrasyonu nedeniyle oksijen saturasyonu <%92 olan COVID-19 hastalarında tedaviye eklenmesi önerilmektedir. Bu çalışmada, 20 Ekim 2020'ye kadar literatürde yayınlanan Tosilizumab ve COVID-19 ile ilgili araştırmalar, meta-analizler ve randomize çalışmalar tarandı. Güncel bilgiler ışığında halen yaygın kullanımda olan Tosilizumab'ın COVID-19 hastalarında etki mekanizması, uygun hasta seçimi, tedavideki etkinliği ve yan etkileri hakkında veriler sunuldu.

Anahtar kelimeler: COVID-19; sitokin fırtınası sendromu; interlökin-6; interlökin-6 antagonisti; tosilizumab

INTRODUCTION

The COVID-19 (Coronavirus Disease-2019) outbreak started with the reporting of pneumonia of unknown cause in Wuhan City, China, in December 2019. It has spread all over the world in a very short time, causing the death of more than 1 million people and approximately 37 million people to become infected as of October 2020, according to WHO data (1). Due to high mortality and morbidity rates, researchers are in search of the best treatment for COVID-19 infection.

Interleukin 6 (IL-6) levels have been found to be higher in patients with severe COVID-19 infection (septic shock, multiorgan failure, high oxygen demand) compared to patients with mild and moderate COVID-19 disease (2,3). Since IL-6 level has been found to be associated with clinical worsening in COVID-19 patients (4), anti-IL-6 therapy is considered as a promising treatment (5).

Tocilizumab, sarilumab and siltuximab are IL-6 antagonists with different pharmacological properties. Sarilumab has been approved only for rheumatoid arthritis (RA) and siltuximab for Castleman disease. However, tocilizumab has also been approved for the treatment of Juvenile Idiopathic Arthritis (JIA) and Giant Cell Arthritis (GCA) in addition to these diseases (6). Moreover, it was approved by the FDA in 2017 for the treatment of Cytokine Storm Syndrome (CSS) (7).

In this review, it was aimed to discuss the mechanism of the action of tocilizumab, which is still in widespread use in COVID-19 patients, appropriate patient selection, effectiveness in treatment and its side effects in the light of current evidence.

Literature Search Method

An electronic literature screening was conducted using Web of Science, PubMed and Google Scholar databases to find retrospective, observational and randomized controlled studies and meta-analyses published on tocilizumab and COVID-19. As Medical Subject Headings (MeSH) search items of "Coronavirus", "Tocilizumab", "COVID-19 and Tocilizumab", "Severe Acute Respiratory Syndrome", "SARS-CoV-2" were created. Reference lists of the studies were also checked to find greater number of research studies related to our search terms. Articles in languages other than English were excluded.

The Pathogenesis and Importance of Cytokine Storm in COVID-19 Disease

Cytokines, which are small proteins secreted by our immune system, have various functions such as communication between cells, regulation of innate and adaptive immunity, hematopoiesis, and repair of damaged tissues (8). Recent studies published have shown that many cytokines (IL-1, IL-2, IL-6, TNF, and IFN-) are significantly increased in COVID-19 patients with pneumonia, and IL-6 is the main cause of the cytokine storm together with other cytokines (9-10). As a trigger of inflammation, IL-6 is highly expressed in COVID-19 patients and induces differentiation of B lymphocytes, antibody production from B lymphocytes, as well as proliferation and differentiation of T (11). B and T lymphocytes, which proliferate and differentiate in excess, enter the pulmonary circulation by causing cytokine storm, leading to lung alveolar damage and respiratory failure (12). It is thought that the definition of cytokine storm was first made in 1993, in graft versus host disease (13). Later, it was shown that cytokine storm due to the



H5N1 virus caused multiple organ failure (14). In CSS, refractory hypotension, respiratory failure, coagulopathy, kidney and liver failure occur due to highly elevated immune activation of lymphocytes, macrophages or myeloid cells. It is thought that the increase in IL-6 and IFN gamma cytokines initiates the cytokine storm by increasing the differentiation of helper T (T helper) lymphocytes (15). Since IL-6 is known as an important mediator of CSS, it is thought that targeted drugs that inhibit IL-6 can block cytokine storm (16).

Possible Mechanism of the Action of Tocilizumab in COVID-19

Mechanism of the action of tocilizumab in the treatment of COVID-19 is still unclear. Previous studies have shown that IL-6 is found in small amounts in fibroblasts, endothelial cells and tumor cells in addition to immune system cells such as B lymphocytes, T lymphocytes, macrophages, and monocytes (17). Normally, very low levels of IL-6 are synthesized rapidly due to reasons such as infection or injury, and its

level in the blood increases. The higher the peak concentration of IL-6 in COVID-19 patients, the more severe the CSS. IL-6 binds to its receptor IL-6R to form a complex, and this complex then binds to signal-converting glycoprotein 130 (gp-130) to initiate signal transduction and triggers gene expression (18). The IL-6 signaling pathway is mediated by two systems. The first pathway is mediated by the JAK/STAT tyrosine kinase system, while the other is mediated by the Ras/mitogen-activated protein kinase (MAPK)/NF-KB-IL-6 pathway. Studies conducted to block signal pathways have been effective in various preclinical, chronic, and autoimmune disease models (19). Tocilizumab, a humanized anti-IL-6R monoclonal antibody, is thought to be effective for the treatment of cytokine storm in COVID-19 disease by inhibiting signaling pathways (Table 1).

Patient Selection and Drug Administration for Tocilizumab in the Treatment of COVID-19 Disease

Early recognition of CSS and starting treatment in COVID-19 patients is important to decrease mortality

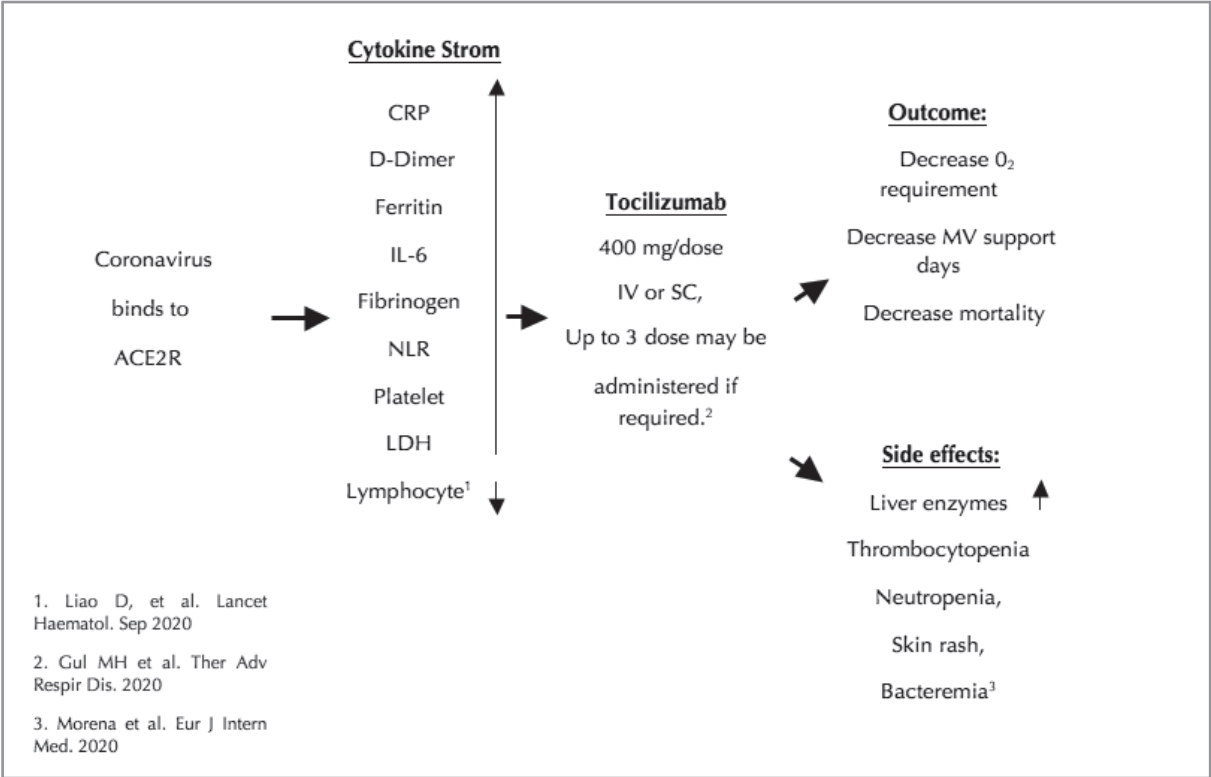


Table 1. The potential mechanism of cytokine storm and Tocilizumab as medical treatment of CSS on COVID-19.

IL: interleukin; ACE2R: ACE 2 Receptors; CRP: C Reactive Protein; NLR: Neutrophil/Lymphocyte Ratio; LDH: Lactat Dehydrogenase Enzyme; O₂: Oxygen; MV: Mechanic Ventilation CSS: Cytokine Storm Sendrome.

and morbidity. CSS should be considered if patients experience clinical worsening and have increased blood IL-6, ferritin, CRP, fibrinogen levels, neutrophil counts, and decreased lymphocyte and platelet counts (20,21). Neutrophil/Lymphocyte (NLR) ratio has also been shown to be important in the diagnosis of CSS (21).

Tocilizumab, which has a monoclonal antibody structure that blocks IL-6 receptors, has been shown to be effective in CSS (22,23). Therefore, tocilizumab has received FDA approval for CSS in addition to rheumatological indication (7). It is recommended to add tocilizumab to the treatment in patients with increased blood IL-6 levels and bilateral diffuse pulmonary infiltration (24).

Tocilizumab can be used alone or in combination with glucocorticoids in adults with severe or life-threatening CSS and in pediatric patients aged >2 years (7). The recommended doses of tocilizumab are 400 mg in patients weighing 50-60 kg; 600 mg for those weighing 60-85 kg, and 800 mg in patients weighing >85 kg (25). Doses exceeding 800 mg per infusion are not recommended (7,24). It is administered intravenously or subcutaneously. If there is no clinical improvement in signs and symptoms after the first dose, up to 3 additional doses may be administered at intervals of at least 8 hours. Infusion time should be at least one hour (7).

Pharmaceutical Effectiveness of Tocilizumab in COVID-19 Disease

Since March 2020, when the pandemic spread all over the world, many studies have been published investigating the effectiveness of tocilizumab in COVID-19 patients (Table 2). In a study conducted by Xu et al. in China, tocilizumab was given to respiratory failure patients with 17 severe and 4 critically ill COVID-19 disease in addition to routine treatment. All patients were discharged after tocilizumab treatment and no drug-related side effects were observed (26). In a study conducted by Sarhan et al. in Egypt, tocilizumab was given to patients with severe COVID-19 disease. Improvement was observed in all laboratory parameters (IL 6, CRP, LDH, ferritin) of the patients and reduction of the length of stay in hospital and ICU, need for MV and mortality rate after double dose Tocilizumab treatment (27). Another prospective study included 85 patients with respiratory failure not requiring mechanical ventilation (MV), better survival

rates were obtained in the tocilizumab group compared to the control group ($p=0.004$). While discharge rate was 92% in the group receiving tocilizumab, it was found to be 42.1% in the control group (28). Similarly, a multi-center retrospective study involving 544 patients, it was found that the need for invasive MV was significantly reduced in the tocilizumab group compared to the standard treatment group ($p=0.020$) (29). Three doses of tocilizumab were administered to 100 patients with respiratory failure, and clinical improvement was observed at a rate of 58% on the 72nd hour, and 77% of on the 10th day of tocilizumab treatment (30). Moreno-Garcia et al. enrolled 171 patients and investigated the requirement for intensive care stay and intubation. They found lower intensive care unit stay and intubation in the tocilizumab group ($p=0.005$ and $p=0.001$, respectively) (31). In a multi-center study including 1221 patients, the primary endpoints were 14-day and 30-day mortality rates. The mortality rates were found to be similar at the end of 14 days, whereas lower 30-day mortality rates were observed in the tocilizumab group ($p=0.52$, and $p<0.001$, respectively) (32). Somers et al. included 154 intubated COVID-19 patients, tocilizumab was found to reduce the risk of death (hazard of death) by 45% [hazard ratio= 0.55 (95% CI 0.33, 0.90)] (33). 1229 COVID-19 patients were divided into two groups according to their CRP levels. In the group with CRP>150 mg/L, tocilizumab significantly reduced the risk of death and number of admissions to intensive care units (aHR 0.34, $p=0.005$; aHR 0.38, $p=0.011$). However, in the group with CRP≤150 mg/L, the mortality rate was similar (34). In the study conducted by Price et al. on 239 patients, tocilizumab was given to all patients considered to have CSS and stratified as those moderately and severely in need of oxygen therapy (>3 Lt O₂). 14-day survival rates of the patients were 91% and 83% in moderately and severely ill patients, respectively, and 86% in all. Interestingly, unlike other studies, survival was found to be better in Afro-Americans and Hispanics than in the Caucasian race (35). Lastly, other group showed better survival in intubated COVID-19 patients who received tocilizumab (36). The effectiveness of anakinra, sarilumab, siltuximab and tocilizumab drugs used in cytokine storm were evaluated in a meta-analysis study. Seventy-one studies and a total of twenty two thousand patients were included. In prospective studies, tocilizumab was associated with improved

Table 2. Characteristics of the studies about tocilizumab effectiveness on COVID-19 treatment

Study	Study type	Population (N)	Treatment	Comparison	Main Findings
Xu et al. (26)	Multi-center retrospective	N: 21 Severe: 17 Critic: 4	TCZ: Up to 800 mg. In case of fever within 12 h, single additional dose	No comparison	All patients discharged on average 15.1 day after TCZ. 75.0% of patients had lower oxygen intake 5 days after TCZ infusion.
Sarhan et al. (27)	Single-center prospective	N: 25	N: 25 TCZ twice doses (up to 800 mg)	No comparison	All patients showed significantly lower median IL-6, LDH, CRP, ferritin, TLC at $p < 0.001$ and D-Dimer at $p = 0.22$ than their baseline. The number of patients who required MV decreased from 11 to 8. Only 5 patients died after TCZ treatment.
Capra et al. (28)	Single-center, retrospective	N: 85	N: 62, TCZ: 400-800 mg	N: 23 Not receiving TCZ	TCZ group had greater survival rate compared to control ($p = 0.004$).
Guaraldi et al. (29)	Multi-center retrospective	N: 544	N: 179, TCZ twice doses (up to 800 mg)	N: 365 Not receiving TCZ	Tocilizumab reduced risk of IMV or death ($p = 0.020$).
Toniati et al. (30)	Single-center prospective	N: 100	TCZ: Up to 800 mg two infusions 12 h apart. A third infusion, 24 h apart from if required.	No comparison	At day 10, 77% improved or stabilized and 23% worsened (of whom 20% died).
Moreno-Garcia et al. (31)	Single-center retrospective	N: 171 Non-ICU patients with COVID-19	N: 77, TCZ: 400 mg ≤ 75 kg and 600 mg > 75 kg. Up to 3 doses if partial response	N: 94 Not receiving TCZ	Tocilizumab group had less ICU admissions ($p = 0.005$) and less invasive ventilation ($p = 0.001$).
Somers et al. (33)	Single-center retrospective	N: 154 severe COVID-19 under mechanical ventilation	N: 78, TCZ (up to 800 mg)	N: 76 Not receiving TCZ	TCZ associated with adjusted lower hazard of death (HR: 0.55)
Perrone et al. (32)	Multi-center, open-label trial, phase 2 study	N: 1221 Hospitalized patients with COVID-19	TCZ up to 800 mg, (Second dose, if required)	No comparison	Mortality was 18.4% and 22.4% at 14 and 30 days in phase 2.
Martinez-Sans et al. (34)	Multicenter retrospective	N: 1229 Hospitalized patients with COVID-19	N: 260, TCZ: median dose 600 mg	N: 969 Not receiving TCZ	TCZ group had higher risk of death (HR 1.53, $p = 0.001$); ICU/death (HR 1.77, $p < 0.001$).
Price et al. (35)	Single-center prospective nonrandomized observational	N: 153 Hospitalized patients with COVID-19	TCZ: Up to 800 mg; second dose if required	No comparison	Moderate and severe patients survival were %91 and %83 respectively at 14 days.
Rossi et al. (36)	Single-center retrospective	N: 246 Patients with severe COVID-19	N: 106, TCZ: 400 mg, single dose	N: 140 Not receiving TCZ	Tocilizumab group had fewer mortality and IMV (H= 0.49 $p = 0.005$)

Table 2. Characteristics of the studies about tosilizumab effectiveness on COVID-19 treatment (continue)

Study	Study type	Population (N)	Treatment	Comparison	Main Findings
Khan et al. (37)	Meta-analysis study	71 studies, (6 random trials) Total 22,058 patients were included	58 studies were about only TCZ treatment. 13 studies were about Anakinra (6), Sarilumab (4), Siltuximab (1), Anakinra and TCZ (1) Sarilumab and TCZ (1).	No comparison	TCZ was associated with a lower relative risk of mortality in prospective studies, but effects were inconclusive for other outcomes. Current evidence for the efficacy of anakinra, siltuximab or sarilumab in COVID-19 is insufficient.
Gupta et al. (38)	Multi-center cohort study	N: 3924 Patients with severe COVID-19	N: 433 (11%) patients received TCZ in the first 2 days of ICU admission.	N: 3491 did not receive TCZ within 2 days of ICU admission	The 1544 patients who died included 125 of the 433 patients (28.9%) treated with TCZ and 1419 of the 3491 patients (40.6%) not treated with TCZ (unadjusted HR, 0.64; 95% CI, 0.54-0.77).
Salama et al. (39)	Randomized, placebo-controlled trial	N: 389 Hospitalized Patients with COVID-19 randomized.	N: 249 patients in the TCZ group	N: 128 patients in the placebo group	Percentage of patients who had received MV or who had died by day 28 was 12.0% in the TCZ group and 19.3% in the placebo group (HR for MV or death, 0.56; p= 0.04 by log-rank test).
Rajendram et al. (40)	Propensity Matched Analysis Study	N: 444 Patients with severe COVID-19	102 patients (23%) received TCZ. 82 patients in each arm were matched between two group.	342 patients (77%) did not receive TCZ	TCZ use was associated with a significant decrease in ICU mortality at 28 days in critically ill coronavirus disease 2019 patients with severe hypoxemic respiratory failure
Kow et al. (41)	Meta-analysis study of randomized controlled trials	Six RCTs were included	Two trials with an overall low risk of bias and four trials had some concerns regarding the overall risk of bias.	No comparison	Despite no clear mortality benefits in hospitalized patients with COVID-19, tocilizumab appears to reduce the likelihood of progression to mechanical ventilation.
Campochiaro et al. (42)	Single-center, prospective	N: 65	N: 32, TCZ: Single dose 400 mg (Second dose, if required)	N: 33 Not receiving TCZ	Discharge from hospital and clinical improvement similar (p= 0.32; p= 0.61, respectively).
Colaneri et al. (43)	Single-center retrospective	N: 112 Hospitalized patients with COVID-19	N: 21, TCZ: up to 800 mg per dose, repeated after 12 h if no side effects	N: 91 Not receiving TCZ	TCZ group had similar ICU admission (p= 0.22) and 7-day mortality (p= 0.84) compared with standard group.
Ip et al. (44)	Multi-center, retrospective, observational	N: 547 ICU patients with COVID-19	N: 134, TCZ: 400 mg (96%), followed by 800 mg (1%), 8 mg/kg (1%), 4 mg/kg (1%), and missing dosing (1%).	N: 413 Not receiving TCZ	TCZ group associated with greater survival (HR= 0.76). 30 day mortality with and without TCZ of 46% versus 56%.
Stone et al. (45)	Randomized, double-blind, placebo-controlled trial	N: 243 Patients with severe COVID-19	N: 161, TCZ up to 800 mg	N: 82 Placebo group	TCZ group had similar worsening of disease (%18.0, %14.9, p= 0.73) and median time to discontinuation of supplemental O ₂ therapy compared with placebo group (5.0 days, 4.9 days, p= 0.69).

IMV: Invasive mechanic ventilation, TCZ: Tocilizumab, HR: Hazard ratio, ICU: Intensive care unit, RCTs: Randomized controlled trials.

survival (HR 0.83, 95% CI 0.72-0.96), but conclusive benefit was not demonstrated for other drugs. However, in the conclusion of this meta-analysis study, it was emphasized that the fact that exact benefits of anakinra, siltuximab and sarilumab could not be demonstrated may be due to the insufficient evidence available (37). A multi-center cohort study conducted by Gupta et al. on 3924 critically ill patients with COVID-19 showed that early administration of tocilizumab was associated with prolonged survival. Patients treated with tocilizumab had a lower risk of death compared to those not treated with tocilizumab (HR, 0.71; 95% CI, 0.56-0.92). The estimated thirty day mortality in the tocilizumab and non-tocilizumab group were found 27.5% and 37.1% respectively (38). In a randomised study conducted by Salama et al. on the modified intention-to-treat population included 249 patients in the tocilizumab group and 128 patients in the placebo group. The cumulative percentage of patients who had received MV or died by day 28 was lower in the tocilizumab group than the placebo group (12.0%; 19.3%; $p=0.04$). However, death from any cause by day 28 was similar between the two groups in this study (10.4%; 8.6%, respectively) (39). In a propensity match analysis study, 82 patients in each arm were matched with patient groups who received tocilizumab and did not receive tocilizumab. ICU mortality was lower in the tocilizumab group (23.2% vs 37.8%) with more ICU, hospital, and vasoactive free days at day 28 compared with those who did not receive tocilizumab (40). In a meta-analysis study including six randomized controlled trials, the overall effect of tocilizumab on the risk of mortality was summarized among patients with COVID-19. Despite no benefits in the mortality of hospitalized patients with COVID-19, tocilizumab appears to reduce the likelihood of progression to MV in this study ($H=0.62$; $n=749$) (41).

In contrast, tocilizumab did not show the expected benefit in some recent studies (42-45). In a prospective study conducted by Campochiaro et al. on 65 COVID-19 patients, the patients were divided into two groups as those receiving standard therapy or tocilizumab. Although survival in the first 28 days was better in the group receiving tocilizumab, discharge rates and clinical improvement rates were similar ($p=0.15$, $p=0.32$, $p=0.61$, respectively) (42). In the same line, Colaneri et al. did not find any difference in tocilizumab and standart therapy group

in terms of need for intensive care and 7-day survival ($p=0.84$, $p=0.22$, respectively) (43). Ip et al. retrospectively investigated 547 COVID-19 patients hospitalized in intensive care units. They found survival similar in the tocilizumab and the control group (44). A total of 243 patients who did not need MV were included in the randomized controlled double-blind study. Tocilizumab was given to one group in addition to the standard treatment, and the other group received only placebo. After 14 days, COVID-19 disease worsened in comparable percentages of the patients in both groups (18% and 14.9%, respectively; $p=0.73$). In addition, the median time required to get rid of the need for oxygen support was similar in both groups (5.0 days, 4.9 days, respectively; $p=0.69$) (45). However, Leaf et al. criticized this study. They signified that this study was severely underpowered because in the placebo group, the percentage of patients with a primary outcome event (i.e., intubation or death) was 12.5%, far lower than the anticipated 30%. So, Leaf et al. emphasized that these results should not be extrapolated to other populations of patients with COVID-19, particularly the critically ill (46).

Tocilizumab has been found to be effective in many studies, but not in other studies which may be related to variables such as the methodology of the studies, the disease severity, heterogeneity in critical methodological issues, late initiation of tocilizumab treatment, comorbidities and age of the patients included in the study. Timing of tocilizumab treatment is very important. It should be evaluated on mild-moderate disease in high-risk patients before any signs of pneumonia. It is recommended that tocilizumab treatment should be initiated within the first two days of CSS since irreversible organ injury can occur after cytokine storm is initiated (38). Findings from other studies emphasized early use of tocilizumab in those with severe disease (29,35). During the early infection period (viral load), tocilizumab has no benefit beacuse CSS is not initiated. In addition, adverse effect and secondary infection risk may increase in the early period.

Drug Safety and Side Effects of Tocilizumab

It has been shown that tocilizumab can be used safely in many autoimmune diseases such as Takayasu arteritis and giant cell arteritis (47-48), but since these studies have been conducted in the restricted patient groups, study data related to a large population are

lacking. Possible side effects of tocilizumab due to its long-term use in autoimmune patients include occurrence of opportunistic infection due to its anti-inflammatory activity (pneumonia, invasive aspergillosis, urinary tract infection, diverticulitis, cystitis), gastrointestinal perforation (especially in concurrent non-steroidal anti-inflammatory users), elevated liver enzymes, infusion-related hypertension, headache, and allergic skin reactions (26,49). However, since short-term use is planned in COVID-19 patients, it is not clear whether the side effects that occur in long-term use also develop in its short-term use.

As expected, serum IL-6 level may increase after tocilizumab administration in COVID-19 patients due to the blockage of receptor IL-6R (50). Although it is recommended to use tocilizumab carefully in those with absolute neutrophil count below $<2 \times 10^9/L$, its use in those with neutrophil counts below $0.5 \times 10^9/L$ is not recommended. In addition, its use is not recommended in patients with platelet levels $<50 \times 10^3/\mu L$. Treatment is recommended to be discontinued in patients whose liver AST/ALT level reaches 5 times the normal (26).

In a study by Campochiaro et al. regarding side effects of tocilizumab in COVID-19 patients, the incidence of adverse events was found to be similar to the group receiving standard therapy (25%, 27%, respectively, $p=0.99$) (42). In the study conducted by Salama et al., serious adverse events occurred similar between tocilizumab and placebo group (38 of 250 patients (15.2%) in the tocilizumab group and 25 of 127 patients (19.7%) in the placebo group) (39). Similarly, retrospective studies revealed no serious side effects related with tocilizumab treatment (43-44).

On the other hand, development of new infection was found to be significantly higher in the group that received tocilizumab compared to the control group (13%, and 4%, respectively; $p<0.001$) (29). In another study including 100 patients, two patients died from septic shock and one from gastrointestinal perforation (30). In a propensity-match analysis study, rates of secondary infections were higher in the tocilizumab cohort (26.5% vs 15.8%) before matching. However, after matching, there was no difference in rates of secondary infection (25.6% vs 25.6%) (40).

Moreover, tocilizumab may increase liver enzymes level, thrombocytopenia, neutropenia, skin rash and

bacteremia (35,51). However, neutropenia was also found in 6-8% of COVID-19 patients who did not receive tocilizumab (52-53). Due to these different data, studies with larger participation are required to evaluate side effects of tocilizumab as neutropenia and bacteremia. In addition, since high transaminase levels may be associated with the progression of COVID-19 disease and CSS in patients receiving tocilizumab, randomized studies are needed so as to be able to claim the presence of tocilizumab-induced hepatotoxicity in these patients.

The Use of Tocilizumab in Special Populations

It has been stated that tocilizumab may have a negative effect on the immune development of the fetus in pregnant women. Caution is recommended in patients over 65 years of age, as it may increase the risk of severe infections. Caution should be exercised in terms of neutropenia and thrombocytopenia. It can be used safely without dose adjustment in mild and moderate renal failure (creatinine clearance >30 mL/min) (26). There are insufficient data for its use in liver failure, but if liver enzymes triple the upper limit of normal due to tocilizumab treatment, it is recommended to wait for these levels to drop down to normal levels (54).

CONCLUSION

Despite significant advances made worldwide in the fight against the COVID-19 pandemic; it is still unclear why some people experience cytokine storm and what the treatment strategy is in these patients. One of the key cytokines in CSS associated with COVID-19 infection is IL-6. Tocilizumab can play an active role in the treatment of the CSS caused by COVID-19 by blocking inflammation via binding IL-6R. However, there is still not enough evidence regarding the clinical efficacy and safety of tocilizumab in COVID-19 patients. Further randomised controlled studies with big sample size are urgently warranted to elucidate the role of tocilizumab in COVID-19 patients.

REFERENCES

1. World Health Organization (WHO). Coronavirus Disease (COVID-19) Dashboard. Accessed date: 15 Nov 2020. Available from: <https://covid19.who.int>
2. Zhou F, Yu T, Du R, Fan G, Liu Y, Liu Z, et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: a retrospective cohort study. *Lancet* 2020; 395: 1054-62.



3. Ruan Q, Yang K, Wang W, Jiang L, Song J. Clinical predictors of mortality due to COVID-19 based on an analysis of data of 150 patients from Wuhan, China. *Intensive Care Med* 2020; 46: 846-8.
4. Coomes EA, Haghighyan H. Interleukin-6 in Covid-19: A systematic review and meta-analysis. *Rev Med Virol* 2020; 30(6): 1-9.
5. Cortegiani A, Ippolito M, Ingoglia G, Einav S. Chloroquine for COVID-19: rationale, facts, hopes. *Crit Care* 2020; 8; 24(1): 210.
6. Tufan A, Avanoğlu Güler A, Matucci-Cerinic M. COVID-19, immune system response, hyperinflammation and repurposing antirheumatic drugs. *Turk J Med Sci* 2020; 21; 50(SI-1): 620-32.
7. The U.S. Food and Drug Administration. Information for ACTEMRA®. Accessed date: 15 Nov 2020. Available from: https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/125276s114lbl.pdf.
8. Tisoncik JR, Korth MJ, Simmons CP, Farrar J, Martin TR, Katze MG, et al. Into the eye of the cytokine storm. *Microbiol Mol Biol Rev* 2012; 76(1): 16-32.
9. Huang C, Wang Y, Li X, Ren L, Zhao J, Hu Y, et al. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. *Lancet* 2020; 395(10223): 497-506.
10. Liu J, Li S, Liu J, Liang B, Wang X, Wang H, et al. Longitudinal characteristics of lymphocyte responses and cytokine profiles in the peripheral blood of SARS-CoV-2 infected patients. *EBioMed* 2020; 55: 102763.
11. Zhai H, Liu Y, Wang Y, Li X, Li T. The pathogen distribution and its influence on inflammatory factors in old patients with heart failure complicated with pulmonary infection. *Tianjin Med J* 2018; 46(9): 952-5.
12. Yonggang Z, Binjing F, Xiaohu Z, Dongsheng W, Changcheng Z, Yingjie Q, et al. Pathogenic T-cells and inflammatory monocytes incite inflammatory storms in severe COVID-19 patients. *Nat Sci Rev* 2020; 7(6): 998-1002.
13. Ferrara JL, Abhyankar S, Gilliland DG. Cytokine storm of graft-versus-host disease: a critical effector role for interleukin-1. *Transplant Proc* 1993; 25(1 Pt 2): 1216-17.
14. De Jong MD, Simmons CP, Thanh TT, Hien VM, Smith GJ, Chau TN, et al. Fatal outcome of human influenza A (H5N1) is associated with high viral load and hypercytokinemia. *Nat Med* 2006; 12(10): 1203-7.
15. Teachey DT, Lacey SF, Shaw PA, Melenhorst JJ, Maude SL, Frey N, et al. Identification of predictive biomarkers for cytokine release syndrome after chimeric antigen receptor T-cell therapy for acute lymphoblastic leukemia. *Cancer Discov* 2016; 6(6): 664-79.
16. Tanaka T, Narazaki M, Kishimoto T. Immunotherapeutic implications of IL-6 blockade for cytokine storm. *Immunotherapy* 2016; 8(8): 959-70.
17. Jones SA, Jenkins BJ. Recent insights into targeting the IL-6 cytokine family in inflammatory diseases and cancer. *Nat Rev Immunol* 2018; 18(12): 773-89.
18. Braun GS, Nagayama Y, Maruta Y, Heymann F, van Roeyen CR, Klinkhammer BM, et al. IL-6 trans-signaling drives murine crescentic GN. *Am Soc Nephrol* 2016; 27(1): 132-42.
19. Rose-John S. The soluble Interleukin 6 receptor: advanced therapeutic options in inflammation. *Clin Pharmacol Ther* 2017; 102(4): 591-8.
20. Zhu Z, Cai T, Fan L, Lou K, Hua X, Huang Z, et al. Clinical value of immune-inflammatory parameters to assess the severity of coronavirus disease 2019. *Int J Infect Dis* 2020; 95: 332-9.
21. Henry BM, de Oliveira MHS, Benoit S, Plebani M, Lippi G. Hematologic, biochemical and immune biomarker abnormalities associated with severe illness and mortality in coronavirus disease 2019 (COVID-19): a meta-analysis. *Clin Chem Lab Med* 2020; 25; 58(7): 1021-8.
22. Fitzgerald JC, Weiss SL, Maude SL, Barrett DM, Lacey SF, Melenhorst JJ, et al. Cytokine release syndrome after chimeric antigen receptor T cell therapy for acute lymphoblastic leukemia. *Crit Care Med* 2017; 45: 124-31.
23. Maude SL, Laetsch TW, Buechner J, Rives S, Boyer M, Bittencourt H, et al. Tisagenlecleucel in children and young adults with B-cell lymphoblastic leukemia. *N Engl J Med* 2018; 378(5): 439-48.
24. Wei P-F. Diagnosis and treatment protocol for novel coronavirus pneumonia (Trial Version 7). *Chin Med J* 2020; 133: 1087-95.
25. Gul MH, Htun ZM, Shaikat N, Imran M, Khan A. Potential specific therapies in COVID-19. *Ther Adv Respir Dis* 2020; 14: 1753466620926853.
26. Xu X, Han M, Li T, Sun W, Wang D, Fu B, et al. Effective treatment of severe COVID-19 patients with tocilizumab. *Proc Natl Acad Sci USA* 2020; 19; 117(20): 10970-5.
27. Sarhan RM, Madney YM, Abou Warda AE, Boshra MS. Therapeutic efficacy, mechanical ventilation, length of hospital stay, and mortality rate in severe COVID-19 patients treated with tocilizumab. *Int J Clin Pract* 2021; 6: 14079.
28. Capra R, De Rossi N, Mattioli F, Romanelli G, Scarpazza C, Sormani MP, et al. Impact of low dose tocilizumab on mortality rate in patients with COVID-19 related pneumonia. *Eur J Intern Med* 2020; 176: 31-35.
29. Guaraldi G, Meschiari M, Cozzi-Lepri A, Milic J, Tonelli R, Menozzi M, et al. Tocilizumab in patients with severe COVID-19: a retrospective cohort study. *Lancet Rheumatol* 2020; 12; 8: 474-84.
30. Toniati P, Piva S, Cattalini M, Garrafa E, Regola F, Castelli F, et al. Tocilizumab for the treatment of severe COVID-19 pneumonia with hyperinflammatory syndrome and acute respiratory failure: a single-center study of 100 patients in Brescia, Italy. *Autoimmun Rev* 2020; 19: 102568.

31. Moreno Garcia E, Rico Caballero V, Albiach L, Agüero D, Ambrosioni J, et al. Tocilizumab is associated with reduction of the risk of ICU admission and mortality in patients with SARS-CoV-2 infection. *medRxiv* 2020; 20113738.
32. Perrone F, Piccirillo MC, Ascierto PA, Salvarani C, Parrella R, Marata AM, et al. Tocilizumab for patients with COVID-19 pneumonia. The single-arm TOCIVID-19 prospective trial. *J Transl Med* 2020; 18(1): 405.
33. Somers EC, Eschenauer GA, Troost JP, Golob JL, Gandhi TN, et al. Tocilizumab for treatment of mechanically ventilated patients with COVID-19. *Clin Infect Dis* 2020; ciaa954.
34. Martínez-Sanz J, Muriel A, Ron R, Herrera S, Pérez-Molina JA, et al. Effects of tocilizumab on mortality in hospitalized patients with COVID-19: a multicentre cohort study. *Clin Microbiol Infect* 2020; 27(2): 238-43.
35. Price CC, Altice FL, Shyr Y, Koff A, Pischel L, Goshua G, et al. Tocilizumab treatment for cytokine release syndrome in hospitalized patients with coronavirus disease 2019: survival and clinical outcomes. *Chest* 2020; 158(4): 1397-408.
36. Rossi B, Nguyen LS, Zimmermann P, Boucenna F, Dubret L, Baucher L, et al. Effect of tocilizumab in hospitalized patients with severe COVID-19 pneumonia: a case-control cohort study. *Pharmaceuticals (Basel)* 2020; 17; 13(10): 317.
37. Khan FA, Stewart I, Fabbri L, Moss S, Robinson K, Smyth AR, et al. Systematic review and meta-analysis of anakinra, sarilumab, siltuximab and tocilizumab for COVID-19. *Thorax medRxiv* 2020; 20076612.
38. Gupta S, Wang W, Hayek SS, Chan L, Mathews KS, Melamed ML, et al. Association between early treatment with tocilizumab and mortality among critically ill patients with COVID-19. *JAMA Intern Med* 2021; 181(1): 41-51.
39. Salama C, Han J, Yau L, Reiss WG, Kramer B, Neidhart JD et al. Tocilizumab in patients hospitalized with Covid-19 pneumonia. *N Engl J Med* 2021; 384(1): 20-30.
40. Rajendram P, Sacha GL, Mehkri O, Wang X, Han X, Vachharajani V, Duggal A. Tocilizumab in coronavirus disease 2019-related critical illness: a propensity matched analysis. *Crit Care Explor* 2021; 3(1): 0327.
41. Kow CS, Hasan SS. The effect of tocilizumab on mortality in hospitalized patients with COVID-19: a meta-analysis of randomized controlled trials. *Eur J Clin Pharmacol* 2021; 2: 1-6.
42. Campochiaro C, Della-Torre E, Cavalli G, De Luca G, Ripa M, Boffini N, et al. Efficacy and safety of tocilizumab in severe COVID-19 patients: a single-center retrospective cohort study. *Eur J Intern Med* 2020; 76: 43-9.
43. Colaneri M, Bogliolo L, Valsecchi P, Sacchi P, Zuccaro V, Brandolino F, et al. Tocilizumab for treatment of severe COVID-19 patients: preliminary results from SMAtteo Covid19 REgistry (SMACORE). *Microorganisms* 2020; 8: 695.
44. Ip A, Berry DA, Hansen E, Goy AH, Pecora AL, Sinclair BA, et al. Hydroxychloroquine and tocilizumab therapy in COVID-19 patients-an observational study. *PLoS One* 2020; 13; 15(8): 0237693.
45. Stone JH, Frigault MJ, Serling-Boyd NJ, Fernandes AD, Harvey L, Foulkes AS, et al. Efficacy of tocilizumab in patients hospitalized with Covid-19. *N Engl J Med* 2020; 383: 2333-44.
46. Stone JH, Horick NK, Healy BC. Tocilizumab in Covid-19 Reply. *N Engl J Med* 2021; 384(1): 87.
47. Nakaoka Y, Isobe M, Takei S, Tanaka Y, Ishii T, Yokota S, et al. Efficacy and safety of tocilizumab in patients with refractory Takayasu arteritis: results from a randomised, double-blind, placebo-controlled, phase 3 trial in Japan (the TAKT study). *Ann Rheum Dis* 2018; 77(3): 348-54.
48. Samson M, Devilliers H, Ly KH, Maurier F, Bienvenu B, Terrier B, et al. Tocilizumab as an add-on therapy to glucocorticoids during the first 3 months of treatment of giant cell arteritis: a prospective study. *Eur J Intern Med* 2018; 57: 96-104.
49. Helleberg M, Steensen M, Arendrup MC. Invasive aspergillosis in patients with severe COVID-19 pneumonia. *Clin Microbiol Infect* 2021; 27(1): 147-8.
50. Nishimoto N, Terao K, Mima T, Nakahara H, Takagi N, Kakehi T. Mechanisms and pathologic significances in increase in serum interleukin-6 (IL-6) and soluble IL-6 receptor after administration of an anti-IL-6 receptor antibody, tocilizumab, in patients with rheumatoid arthritis and Castleman disease. *Blood* 2008; 15; 112(10): 3959-64.
51. Morena V, Milazzo L, Oreni L, Bestetti G, Fossali T, Bassoli C, et al. Off-label use of tocilizumab for the treatment of SARS-CoV-2 pneumonia in Milan, Italy. *Eur J Intern Med* 2020; 76: 36-42.
52. Feng Y, Ling Y, Bai T, Xie Y, Huang J, Li J, et al. COVID-19 with different severities: a multicenter study of clinical features. *Am J Respir Crit Care Med* 2020; 201(11): 1380-8.
53. Goyal P, Choi JJ, Pinheiro LC, Schenck EJ, Chen R, Jabri A, et al. Clinical characteristics of Covid-19 in New York City. *N Engl J Med* 2020; 11; 382(24): 2372-4.
54. Barlow A, Landolf KM, Barlow B, Yeung SYA, Heavner JJ, Claassen CW, et al. Review of emerging pharmacotherapy for the treatment of coronavirus disease 2019. *Pharmacother* 2020; 40(5): 416-37.