

Letter to the Editor

Can we use Mesenchymal Stem Cell transplantation for COVID-19 patients in puerperium period?

Şahin et al. Mesenchymal Stem Cell transplantation for COVID-19

Ayça Sultan Şahin¹, Ebru Kaya¹, Gürsel Turgut², Ali Kocataş³

¹Department of Anesthesiology and Reanimation, Kanuni Sultan Süleyman Training and Research Hospital, İstanbul, Turkey

²Department of Plastic and Reconstructive Surgery, Genkord-Umbilical Cord Blood Banking, İstanbul, Turkey

³Department of General Surgery, Kanuni Sultan Süleyman Training and Research Hospital, İstanbul, Turkey

Address for Correspondence: Ayça Sultan Şahin,

Phone: +90 505 398 04 19 e-mail: aycasultan@gmail.com

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To the Editor,

The COVID-19 pandemic has affected the whole world, and pregnant women and women in the postpartum period have not been spared from its effects. Pregnant women during this pandemic pose an unknown risk to their babies. We followed up pregnant and post-pregnancy COVID-19 patients in this period in the intensive care unit and used Mesenchymal Stem Cell (MSCs) transplantation in suitable patients. We aimed to investigate patients with COVID-19 pneumonia and who were hospitalized in the intensive care unit (ICU) after C-section in the puerperium period and either given or not given MSCs treatment.

Case 1

A 33 year-old multiparous woman with a history of pregnancy at 33 gestation weeks, admitted with fever ($>38^{\circ}\text{C}$) and cough, with a presumptive diagnosis of COVID-19. Chest CT showed patchy ground-glass opacity (Figure-1). On the third day of hospitalization, she had severe dyspnea and tachypnea and delivery was advised, so she underwent C-section with spinal anesthesia. The patient was transferred to ICU after labor with respiratory distress and X-rays showed patchy ground-glass opacities. She was intubated. D-dimer, C-reactive-protein, and ferritin levels continued to increase up to the tenth day after the C-section. The patient's clinical situation was evaluated as cytokine storm. After tocilizumab treatment, MSC treatment was given twice with two days intervals. At the end of the third week the patient was extubated. X-rays showed dramatic healing of the patchy ground-glass opacity. When she could breath air unassisted, she was discharged from ICU and subsequently to home.

Case 2

A 34 year-old woman was admitted to ICU after cesarean section with a diagnosis of COVID-19. She had tachypnea and shortness of breath in the postoperative period. High flow nasal cannula oxygenation, intermittent non-invasive ventilation and intermittent prone positioning was applied at ICU follow up. The patient did not respond to seven days of this treatment so

she was intubated. Again, D-dimer, C-reactive-protein, and ferritin levels continued to increase. On the eleventh and twelfth days, she received immune plasma therapy. After 21 days, despite all these interventions, she unfortunately died.

MSCs are tolerated by the recipient immune system while also being immunomodulatory. MSCs are known to function via several mechanisms relevant to acute lung injury. When administered intravenously they sequester in the lung. The effects of MSC are known to include: anti-inflammatory mechanisms; inhibition of lung fibrosis; lung tissue regeneration; and an anti-apoptotic effect in injured cells (1). After MSC injection, cytokine-secreting immune cells, including NK cells, were markedly reduced and pro-inflammatory cytokine TNF- α was decreased (2). MSC transplantation may be an appropriate choice of therapy in patients with severe COVID-19 to prevent severe morbidity and mortality in critically ill patients in the future.

References

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Figure 1. Thorax CT image before ICU admission