THE EFFICACY OF EARLY ADDITIONAL HEMOPERFUSION THERAPY FOR SEVERE AND CRITICAL COVID-19 PATIENTS IN A TERTIARY HOSPITAL : A PROSPECTIVE COHORT STUDY

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Objectives

 To investigate the effects of early HA-330 hemoperfusion in combination with standard therapy in severe COVID-19 patients.

Methods

- Single center, prospective cohort study on severe COVID-19 patients who admitted to ICU

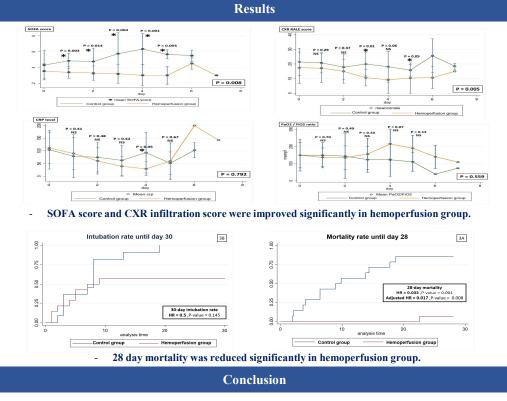
- Compared patients in **"Hemoperfusion group"** (defined as patients who were treated with hemoperfusion therapy at least 3 sessions in combination with standard therapy : N = 15) with **"Control group"** (defined as patients who received standard treatment alone or received less than 3 sessions of hemoperfusion therapy : N = 14)

- Hemoperfusion method: HA-330 disposable hemoperfusion 1 session/day (Jafron Biomedical)

: Duration of each session 3-4 h for 3 consecutive day

- Primary outcomes : Daily C-reactive protein , PaO_2 and ratio of PaO_2/FiO_2 , severity scoring

- of lung infiltration on the chest x-ray (CXR RALE score) and organ failure (SOFA)
- Secondary outcomes : Ventilator free day, Hospital mortality and 28-day mortality



In severe COVID-19 patients, the addition of at least 3 sessions of hemoperfusion therapy to standard therapy seemed to improve severity of organ failure, CXR severity score, ventilator-free day and reduced the mortality rate.

