## CLINICAL CHARACTERISTICS AND OUTCOMES OF PATIENTS WITH LIVER FAILURE TREATED WITH MOLECULAR ADSORBENT RECIRCULATING SYSTEM (MARS) - A SINGLE CENTER EXPERIENCE -

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## INTRODUCTION

- Molecular Adsorbent Recirculating System (MARS<sup>™</sup>) is an albumin-based dialysis system has been utilized in patients with severe liver failure with AKI and Hepatic Encephalopathy at our center for the last 3 years.
- The aim of this retrospective study was to identify change in total bilirubin, ammonia, and PSE stage in patients with acute liver failure (ALF) and acute-on-chronic liver failure (ACLF) who received MARS.

## METHODS

- This is a single center retrospective study performed at Methodist Dallas Medical Center.
- Data included 42 patients with alcoholic hepatitis (AH), ACLF or ALF who underwent at least one session of MARS from January 2019 to March 2022.
- MARS was performed 12 hrs/day for 5 days MARS was interrupted in the case of hemodynamic instability, clinical deterioration, or in cases of substantial clinical improvement.
- Measured outcomes included change in total bilirubin, ammonia, MELD-Na scores and PSE stage pre- and post-MARS. We assessed their overall mortality after receiving MARS.
- Wilcoxon signed-rank test was used to compare continuous variables. McNemar's test was used for categorical variables.



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# IMAGES

#### Variable

Age [years, median (Q1-Q3)]

Etiology of Liver Failure (%)

Sessions of MARS therapy [median (Q1-Q3]

ICU Length of Stay [days, median (Q1-Q3)]

Length of Stay [days, median (Q1-Q3)]

28-day transplant free survival (n, %)

30-day readmission (n, %)

Table 1: Patient demographics



Figure 2: Frequency of patients in portosystemic encephalopathy stages 0-4 at baseline and after MARS therapy.

Total Patients (N=42)
<b>47</b> (38-57)
81.8% AH 57.1% AoCLF 38.1% ALF
4 (3-5)
<b>7.5</b> (4-14)
<b>16.5</b> (11-32)
<b>15</b> (38.4%)
<b>18</b> (42.9%)



Figure 1: Line graph showing outcomes after treatment with MARS therapy.

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### RESULTS

- We studied data on 42 patients.
- Median age was 47 years old.
- Etiology of liver failure comprised of 57.1% ACLF and 38.1% ALF; 81.8% had AH.
- Mean sessions of MARS was 4.
- 28-day transplant free survival was 38.4% (15/39) 3 patients underwent liver transplant. Overall survival was 42.8% (18/42).
- There was a statistically significant difference in total bilirubin (median 20.35 vs 17) and ammonia levels (median 67 vs 31) after MARS.
- 30 patients had available pre- and post-MARS ammonia levels – this population was not significantly different from the total population. The proportion of patients with stage 3 or 4 PSE significantly decreased after MARS therapy (28/42 vs 15/42).
- There was no statistically significant difference in MELD-Na pre- and post-MARS (mean 38 vs 37).

## CONCLUSIONS

- In this descriptive and exploratory analysis, MARS therapy led to a significant decrease in total bilirubin and ammonia levels, and in the proportion of patients with PSE stage 3 or 4.
- There was not a significant change in MELD-Na.
- Overall mortality was 57.1% (The predicted mortality of this group based on this MELD score is 70%<sup>4</sup>). A more meaningful analysis could be achieved with historical controls based on age and MELD, which is in process.

### REFERENCES

- 1. Reuben et al. Ann Intern Med. 2016, 164 (11) 724-732
- 2. Kamath, PS *Clin Liver Dis.* **2017**, 9 (4), 84-88
- Hernaez et al. *Gut.* **2017**, *6*6 *(*3) 541-553
  UNOS. https://optn.transplant.hrsa.gov/
- 5. Saliba, F. *Crit Care.* **2006**, 10 (1)
- 6. Garcia et al. Ann Intensive Care. 2018, 109 (2018)