ABSTRACT

Background and Objective: SARS Cov 2 virus are spreading havoc throughout the world and is responsible for the present pandemic. The clinical presentation of this pandemic has been varied from simple flu like symptoms to severe pneumonia. It is also found to affect multiple organ systems in our body. This study was conducted to find out the variations in the biochemical parameters among SARS Covid patients admitted in our hospital.

Materials and Methods: This study was a retrospective cross sectional survey of medical records of 258 SARS Cov 2 patients admitted in Chettinad hospital and research institute from July 2020 to September 2020. Among them 183 were males, 75 were females. The details of their liver profile, renal profile, cardiac profile, ferritin levels and LDH levels were retrieved and studied.
**Results:** In this study we found a significant difference in the levels of ferritin, LDH, serum albumin, troponin I, CK-MB and BNP levels between patients admitted in ICU and ward. The patients were divided into three groups based on their age as 20-50, 50-80 and more than 80 years. We found significant difference in the levels of total protein, serum albumin, ALT, BUN, troponin I and CK-MB values. Regression analysis showed ferritin, LDH and serum albumin to be independent markers in predicting disease severity (p value <0.05).

**Conclusion:** This study has shown that ferritin, LDH and serum albumin are promising biochemical markers in the diagnosis and prognosis of SARS Covid 19 infection.

**Keywords:** Cardiac profile; ferritin; liver profile; renal profile; sars covid; lactate dehydrogenase.

1. **INTRODUCTION**

SARS COVID viruses are a family of viruses that can cause respiratory and intestinal diseases in animals and humans [1]. These viruses tend to attack the cells of the upper respiratory tract. While most of the Covid-19 cases are found to be mild, some patients also present with respiratory failure, septic shock and/or multi organ failure and death. SARS Cov 2 infection has been declared as pandemic by the World Health Organization (WHO) having led to deaths and hospitalization of millions of people worldwide. Laboratory diagnostic tests play an important role in the diagnosis, assessment of prognosis, severity and therapeutic monitoring of different diseases among which the role of biochemical blood tests in assessing the severity and prognosis of the present pandemic is valuable [2]. According to the centre of disease control and prevention, elderly patients with comorbidities such as hypertension, diabetes and cardiac problems had a higher chance of contracting SARS Covid infection [3]. Some studies have shown that SARS Covid infected patients with cardiovascular morbidity had a greater risk of progressing to severe stages [4]. The pathophysiology behind multi organ involvement in SARS Cov 2 may be due to the immune response specific to each organ leading to the development of multi organ failure [5]. Recent studies have shown the involvement of liver and that Angiotensin Converting Enzyme - 2(ACE-2) is the cell entry receptor for SARS Covid-19 virus and this enzyme is expressed in a higher concentration in the type II alveolar cells of lungs and also in the bile duct cells [6,7]. Early detection of patients with chances of deterioration should be done to prevent mortality from SARS Covid-19 cases. If a link between the biochemical parameters and severity of SARS Covid-19 cases are found out it can be useful in the better management of SARS Cov 2 patients in the near future.

1.1 **Aim**

Aim of this study was to analyze the variations in the liver profile, cardiac profile, renal profile, ferritin and LDH among SARS Cov 2 patient.

2. **MATERIALS AND METHODS**

2.1 **Study Design**

This is a single centre retrospective cross sectional study done over a period of three months from July to September 2020.

2.2 **Inclusion Criteria**

Details of 500 patients of the age group 20-80 years of both sex who attended Chettinad Hospital and Research Institute from July 2020-September 2020 diagnosed as SARS Cov 2 were included in the study. Among them 242 subjects were excluded from the study due to incomplete data.

2.3 **Exclusion Criteria**

SARS Covid-19 patients with previous H/O liver diseases, renal diseases, cardiac problems, hematological disorders, HIV infection, pregnant women, patients less than 18 years of age, patients with cancer, were excluded from this study.

The medical records of all patients were collected and examined. The details of biochemical investigations, clinical, drug history and demographic details of the patients were retrieved. The patients in our hospital were grouped based on their admission in ICU or ward. Details regarding patients age, gender, previous comorbidities, previous drug history, present clinical profile were collected. Data regarding biochemical investigations like liver profile, lactate dehydrogenase, serum creatinine, blood urea, troponin I, CK-MB and serum ferritin were...
3. RESULTS

The study group included 258 cases of SARS Covid patients admitted to Chettinad Hospital and research institute. The group included 187 males and 77 females. In this study we found a significant difference in the levels of ferritin (454.12 +/- 396.95) (323 +/- 362.4) (p value 0.006), LDH (402.6 +/- 199.3) (328.16 +/- 250.62) (p value 0.0084), serum albumin (3.24 +/- 0.5) (3.46 +/- 0.5) (p value 0.009), troponin I (155.84 +/- 528.2) (44.916 +/- 140.9) (p value 0.03), CK-MB (5.8 +/- 10.4) (3.3 +/- 5.9) (p value 0.02) and BNP levels (357.4 +/- 996) (146.5 +/- 490) (p value 0.036) among patients admitted in ICU and ward.

The patients were divided into three groups based on their age as 20-50, 50-80 and more than 80 years. We found out a significant difference in the levels of total protein (7.02 +/- 0.64) (6.72 +/- 0.59) (6.92 +/- 0.5) (p value 0.0008), serum albumin (3.57 +/- 0.5) (3.2 +/- 0.52) (3.25 +/- 0.24) (p value 0.0001), ALT (58.04 +/- 63.86) (43 +/- 30.51) (24.5 +/- 6.14) (p value 0.02), BUN (12.79 +/- 10.49) (21.4 +/- 20.18) (35.5 +/- 28.34) (p value 0.0001), troponin I (55.36 +/- 271.36) (386.94 +/- 2267.6) (2103.9 +/- 3941.9) (p value 0.05) and CK-MB (2.46 +/- 4.63) (5.61 +/- 9.24) (17.7 +/- 29.42) (p value 0.0002) values. Regression analysis showed ferritin, LDH and serum albumin to be independent markers in predicting disease severity.

4. DISCUSSION AND CONCLUSION

This study was done among 258 SARS Covid patients admitted in Chettinad hospitals. Among them 183 were males and 75 were females. This number shows that males are more affected than their female counterparts. A study done by Dr. Kyle Sue shows the effect of hormones on the immunologic responses of males leading to a less robust immune response among males [8]. X chromosomes carry the largest number of immune related genes causing superior immune response in females [9]. Animal models have shown that the ACE 2 receptor which has been proposed to be the entry point of SARS Covid to be more expressed among males [10]. The variables did not show much statistical difference between males and females, except for albumin, BUN creatinine and troponin I. Multiple organ involvement has been found to be increasing the morbidity of SARS Covid 19 patients. Covid 19 pandemic has given way for understanding various biochemical alterations in the liver and cardiac profile of patients suffering from this viral disease. Various studies are being done to understand the correlation between these parameters and the severity of the disease. Ferritin has been considered as a marker for finding out the severity of SARS Covid 19 patients. Ferritin is the storage form of iron in tissues. It has two subunits H & L. The levels of these two subunits vary from tissue to tissue. Among these subunits, H subunit is found to have an inflammatory and immunomodulatory function. Ferritin has been found to increase the levels of inflammatory markers in the blood like interleukins. The increase in the levels of serum ferritin during inflammation may be due to its active secretion from the hepatocytes or by macrophages through a non-classical pathway [11,12,13]. In our study we found out that the levels of ferritin among SARS Covid patients were higher than the normal reference interval and when compared between the patients admitted in ICU (454.12 +/- 396.95) and ward (323 +/- 362.4), the difference between the levels were statistically significant. (p value = 0.006) (Table 3). Free iron released from ferritin is a pro inflammatory mediator, responsible for alteration in the coagulation profile leading to vascular thrombosis in various organs especially in lungs. Hence iron chelation therapy may be helpful in decreasing the iron induced coagulopathy. This may form the therapeutic basis for iron chelating agents in the treatment of severe SARS Covid infection as evidenced by the increased levels of ferritin in the biochemical profile [14,15,16]. A recent study done among 39 SARS Covid 19 patients showed that serum ferritin levels correlated with the severity of the disease [17]. This study shows a
Albumin levels were found to be mildly elevated than the normal values. Total protein levels and albumin levels were found to have a significant difference between different age groups (Table 3). Albumin levels were found to have a significant difference between patients admitted in ICU (328.16+/−250.62) (p value=0.008) (Table 3). Ferritin and LDH were independently associated with the disease as shown by the regression analysis. (Fig. 2). SARS Covid 19 has been found to affect multiple organs in our body especially liver. Studies have shown Liver profile derangements were found among 50% of patients infected with SARS Covid 19 [19]. Studies have found out that SARS Covid 19 virus infect the Angiotensin converting enzyme (ACE) receptor on cholangiocytes leading to its dysfunction and inturn causing widespread systemic inflammatory response [20]. We have analyzed the liver profile of all the SARS Covid 19 patients which included Total protein, albumin, Aspartate transaminase (AST), Alanine amino transferase (ALT), Alkaline phosphatase (ALP) and GGT (Gamma glutamyl transferase). In this study, the various liver parameters were found to be mildly elevated than the normal values. Total protein levels and albumin levels were found to have significant difference between different age groups (Table:3). Albumin levels were found to have a significant difference between patients admitted in ICU (3.24+/−0.5) than those admitted in ward (3.46+/−0.5) (p value=0.009). Albumin is a negative acute phase protein. Cytokines released during inflammation caused by SARS Covid 19 affect the liver which is the main organ producing all the proteins in our body and result in pre translational modification of gene expression of all acute phase proteins released from liver thereby decreasing the levels of negative acute phase proteins and increasing the levels of positive acute phase proteins like ferritin and CRP [21]. Liver is the sole source of albumin. Albumin synthesis in liver depends on the amino acids reaching the intestine from the various food intakes, but during infections, the amount of food intake by the patients will be less which inturn leads to a decrease in the synthesis of albumin [22]. Albumin levels were found to be independently associated with serum ferritin levels and as the ferritin levels increased, the albumin levels were found to decrease.  

This again emphasis the use of albumin as an independent marker for the prediction of disease severity. ALT levels were found to have a significant difference among patients of different age groups 20-50 years 58.04 ± 63.86, 50-80 years 43 ± 30.51 and >80 years -24.5 ± 6.149 (p value=0.02). There was no significant difference in the levels of other liver enzymes among the different age groups and also between those admitted in ICU and ward as shown by some recent studies [23, 24]. Cardiac parameters like Troponin I, BNP and CK-MB were found to have significant difference between the patients admitted in ICU and ward pointing to the risk of developing cardiac abnormalities among SARS Covid patients. A study done by Lippi G, Pebani M has shown that 8-12% of the SARS Covid 19 patients had severe cardiac abnormalities [25]. Another study done by Huang C, Wang Y, Li X, et al has shown that myocardial injury among SARS Covid patients presented as marginal increase in the troponin I levels [26].

In this study we assessed the variations in the levels of renal parameters like Blood urea nitrogen and creatinine levels. It was found out that the mean levels of creatinine was 1.84 ± 5.04 and the mean value of BUN was 18.38 ± 17.88 respectively. ANOVA test done among the different age groups showed a significant difference in the levels of BUN among the different age groups (p value =0.0007), but creatinine values did not show much difference between the different age groups. There was no significant difference in the values of BUN and creatinine levels among patients admitted in ward and ICU. Although the exact mechanism of the involvement of kidney in SARS Covid remain unclear, a number of causes like hyperferritinemina, hypercoagulability, hypercomplementemia, damage to the kidney by direct injury to the tubular cells has been proposed [27]. ACE2 receptors, the source of entry of SARS Covid virus has been found in kidney cells. [28, 29] Studies also report the isolation of SARS Covid virus in the blood and urine of patients affected with the disease [30]. This study has thrown light on the importance of biochemical parameters in the treatment of SARS Covid patients. Ferritin, serum albumin and LDH levels seem to be promising markers in assessing the severity of SARS Covid patients. Proper monitoring of these parameters can help treating physicians in finding out the progress of the disease condition.
Table 1. Shows the descriptive statistics of variables among Covid 19 patients

| Variable       | N   | N Miss | Mean ± STD   |
|----------------|-----|--------|--------------|
| Total Protein  | 258 | 0      | 6.84 ± 0.63  |
| Albumin        | 258 | 0      | 3.34 ± 0.54  |
| Total_bilirubin| 258 | 0      | 0.6 ± 0.8    |
| Direct_bilirubin| 258 | 0      | 0.18 ± 0.17  |
| AST            | 258 | 0      | 48.97 ± 50.76|
| ALT            | 258 | 0      | 48.37 ± 46.37|
| ALP            | 258 | 0      | 76.71 ± 31.54|
| GGT            | 258 | 0      | 63.44 ± 85.48|
| FERRITIN       | 258 | 0      | 394.25 ± 386.55|
| LDH            | 258 | 0      | 367.98 ± 227.26|
| Creatinine     | 258 | 0      | 1.84 ± 5.04  |
| BUN            | 258 | 0      | 18.38 ± 17.88|
| Trop_I         | 257 | 1      | 289.8 ± 184.54|
| CK_Myopathy    | 254 | 4      | 4.65 ± 8.7   |
| BNP            | 258 | 0      | 259.29 ± 807.34|

*AST-Aspartate transaminase, ALT-Alanine amino transferase, ALP-Alkaline phosphatase, GGT-Gammaglutamyltransferase, BUN-Blood urea nitrogen, CK-MB-creatine kinase, BNP-Brain natriuretic peptide (p value<0.05 is considered significant)

Table 2. ANOVA analysis to compare the variables between the three groups divided based on age 20-50, 51-80 and >80 years

| Variables       | Total N = 258 | Age_Group (20-50) N = 97 | Age_Group (51-80) N = 157 | Age_Group (> 80) N = 4 | Pval   |
|-----------------|---------------|--------------------------|---------------------------|-------------------------|--------|
| Protein (g/dl)  | 97            | 7.02 ± 0.64              | 6.72 ± 0.59               | 6.9 ± 0.5               | 0.0008 |
| Albumin (g/dl)  | 97            | 3.57 ± 0.5               | 3.21 ± 0.52               | 3.25 ± 0.24             | <.0001 |
| Total_bilirubin (mg/dl) | 97 | 0.59 ± 0.82 | 0.61 ± 0.8 | 0.48 ± 0.23 | 0.9457 |
| Direct_bilirubin (mg/dl) | 97 | 0.17 ± 0.19 | 0.19 ± 0.15 | 0.12 ± 0.09 | 0.5614 |
| AST(IU/L)       | 97            | 44.66 ± 33.78            | 51.68 ± 59.3              | 47.5 ± 20.14            | 0.5652 |
| ALT(IU/L)       | 97            | 58.04 ± 63.86            | 43 ± 30.51                | 24.5 ± 6.14             | 0.0243 |
| ALP(IU/L)       | 97            | 73.3 ± 19.63             | 78.92 ± 37.2              | 72.75 ± 16.17           | 0.3755 |
| GGT(mg/dl)      | 97            | 65.02 ± 63.57            | 63.51 ± 97.49             | 22.25 ± 10.14           | 0.6199 |
| FERRITIN        | 96            | 368.72 ± 426.33          | 411.03 ± 362.25           | 348.5 ± 364.6           | 0.682  |
| LDH (U/L)       | 97            | 345.04 ± 280.04          | 379.14 ± 187.87           | 486.5 ± 193.32          | 0.2941 |
| Creatinine      | 97            | 1.08 ± 1.52              | 2.3 ± 6.3                 | 2.23 ± 2.08             | 0.1701 |
| BUN(mg/dl)      | 97            | 12.79 ± 10.49            | 21.4 ± 20.18              | 35.5 ± 28.34            | 0.0001 |
| Trop_I (pg/ml)  | 96            | 55.36 ± 271.36           | 386.94 ± 2267.66          | 2103.9 ± 3941.93        | 0.0538 |
| CK_Myopathy     | 93            | 2.46 ± 4.63              | 5.61 ± 9.24               | 17.7 ± 29.42            | 0.0002 |
| BNP(pg/ml)      | 97            | 215.25 ± 790.27          | 282.43 ± 827.43           | 419.25 ± 756.46         | 0.7517 |

*AST-Aspartate transaminase, ALT-Alanine amino transferase, ALP-Alkaline phosphatase, GGT-Gammaglutamyltransferase, BUN-Blood urea nitrogen, CK-MB-creatine kinase, BNP-Brain natriuretic peptide (p value<0.05 is significant)
Table 3. Independent sample t test shows the comparison between variables among patients admitted in ICU and ward

| Variable                  | Mean+/−SD Cases admitted in ICU | Mean+/−SD Cases admitted in ward | P-value |
|---------------------------|---------------------------------|----------------------------------|---------|
| Total protein(g/dl)       | 6.8+/−0.67                      | 6.8+/−0.6                        | 0.19    |
| Albumin(g/dl)             | 3.24+/−0.5                      | 3.46+/−0.5                       | 0.009*  |
| Total bilirubin(mg/dl)    | 0.54+/−0.3                      | 0.67+/−1.13                     | 0.17    |
| Direct bilirubin(mg/dl)   | 0.19+/−0.13                     | 0.17+/−0.19                     | 0.41    |
| AST(IU/L)                 | 52.6+/−44.6                     | 44.7+/−56.8                     | 0.22    |
| ALT(IU/L)                 | 49.9+/−54.35                    | 46.6+/−35.18                    | 0.57    |
| ALP(mg/dl)                | 77.26+/−31.9                    | 76.08+/−31.2                    | 0.76    |
| GGT(mg/dl)                | 64.8+/−54.8                     | 61.8+/−111                      | 0.78    |
| Ferritin(ng/ml)           | 454.12+/−396.95                 | 323+/−362.4                     | 0.006*  |
| LDH(u/l)                  | 402.6+/−199.3                   | 328.16+/−250.62                 | 0.008*  |
| Creatinine(mg/dl)         | 1.63+/−2.7                      | 2.08+/−6.8                      | 0.48    |
| BUN(mg/dl)                | 20.09+/−19.7                    | 16.4+/−15.3                     | 0.09    |
| Troponin I(pg/ml)         | 155.84+/−528.2                  | 44.916+/−140.9                  | 0.03*   |
| CK-MB(mg/ml)              | 5.8+/−10.4                      | 3.3+/−5.9                       | 0.02*   |
| Ferritin(ng/ml)           | 363.28+/−405.06                 | 47.088                          | 0.89    |
| LDH(U/L)                  | 49.54+/−38.4                    | 2.946                           | 0.131   |
| AST(IU/L)                 | 51.20+/−56.38                   | 6.554                           | 0.17    |
| ALT(IU/L)                 | 69.75+/−97.44                   | 7.473                           | 0.17    |
| ALP(U/L)                  | 363.28+/−405.06                 | 47.088                          | 0.89    |
| GGT(mg/dl)                | 396.19+/−385.63                 | 29.577                          | 0.88    |
| LDH(U/L)                  | 354.71+/−250.018                | 19.175                          | 0.97    |
| CREAT(mg/dl)              | 1.224+/−1.539                   | 0.179                           | 0.05*   |
| BUN (mg/dl)               | 2.123+/−6.0938                  | 0.467                           | 0.018   |
| Troponin I(pg/ml)         | 576.651+/−3232.1630             | 375.7313                        | 0.04*   |
| CK-MB(mg/ml)              | 4.908+/−10.6705                 | 1.2404                          | 0.94    |
| BNP(pg/ml)                | 207.31+/−603.943                | 70.207                          | 0.28    |

Table 4. Independent sample T test showing the difference in variables among male and female SARS Covid patients

| SEX | Variable          | Mean | Std. Deviation | Std. Error Mean | P value |
|-----|-------------------|------|----------------|-----------------|---------|
| ALB (g/dl) | F | 3.342 | 0.4626 | 0.0538 | 0.019* |
| M | 3.355 | 0.5679 | 0.0436 |
| Total bilirubin(mg/dl) | F | 0.547 | 0.9004 | 0.1047 | 0.711 |
| M | 0.638 | 0.7847 | 0.0602 |
| Direct bilirubin(mg/dl) | F | 0.146 | 0.1398 | 0.0162 | 0.271 |
| M | 0.202 | 0.1804 | 0.0138 |
| AST(IU/L) | F | 48.88 | 45.959 | 5.343 | 0.835 |
| M | 49.11 | 54.363 | 4.169 |
| ALT(IU/L) | F | 45.36 | 61.777 | 7.181 | 0.698 |
| M | 49.78 | 38.406 | 2.946 |
| ALP(U/L) | F | 74.54 | 23.334 | 2.713 | 0.131 |
| M | 78.78 | 34.794 | 2.669 |
| GGT(mg/dl) | F | 51.20 | 56.383 | 6.554 | 0.17 |
| M | 69.75 | 97.434 | 7.473 |
| FERRITIN(ng/ml) | F | 363.28 | 405.064 | 47.088 | 0.89 |
| M | 396.19 | 385.631 | 29.577 |
| LDH(U/L) | F | 389.81 | 177.561 | 20.641 | 0.88 |
| M | 354.71 | 250.018 | 19.175 |
| CREAT(mg/dl) | F | 1.224 | 1.5393 | 0.179 | 0.05* |
| M | 2.123 | 6.0938 | 0.4674 |
| BUN (mg/dl) | F | 15.21 | 13.828 | 1.607 | 0.018* |
| M | 19.35 | 19.133 | 1.467 |
| Troponin I(pg/ml) | F | 576.651 | 3232.1630 | 375.7313 | 0.04* |
| M | 183.101 | 784.7633 | 60.1886 |
| CK-MB(mg/ml) | F | 4.908 | 10.6705 | 1.2404 | 0.94 |
| M | 4.739 | 7.9537 | 0.6100 |
| BNP(pg/ml) | F | 207.31 | 603.943 | 70.207 | 0.28 |
| M | 262.52 | 843.434 | 64.688 |

*AST-Aspartate transaminase, ALT-Alanine amino transferase, ALP-Alkaline phosphatase, GGT-Gammaglutamyltransferase, BUN-Blood urea nitrogen, CK-MB-creatinekinase, BNP-Brain natriuretic peptide (p value<0.05 is significant)
Fig. 1. Shows the regression analysis of ferritin and LDH

Fig. 2. Shows the regression analysis of ferritin and albumin

CONSENT
It is not applicable.

ETHICAL APPROVAL
Institute ethical committee approval was obtained before commencement of the study.(proposal No:076/IHEC/July2020).

COMPETING INTERESTS
Authors have declared that no competing interests exist.

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