# RIFLE classification as a predictor of short term prognosis in critically ill cirrhotic patient

#### **Thesis**

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## **Summary**

End-stage liver disease is frequently complicated by renal function disturbances. Cirrhotic patients with renal failure admitted to intensive care units (ICUs) have high mortality rates. So, early detection of Acute Kidney Injury (AKI) in these patients is very important to prevent complication and eventually decreases mortality rate.

Most of the commonly used clinical scoring systems evaluate renal function according to the serum creatinine level which is unable to detect AKI in the early stages where the serum creatinine may be low while GFR is markedly reduced since there may not have been sufficient time for the creatinine to accumulate.

The RIFLE (risk of renal failure, injury to the kidney, failure of kidney function, loss of kidney function, and end-stage renal failure) classification was first proposed by the Acute Dialysis Quality Initiative (ADQI) group at the second ADQI conference in Vicenza, Italy, in May 2002, in an attempt to standardize the study of ARF. The RIFLE criteria classify ARF into three groups (risk, injury, and failure) according to changes in SCr and/or urine output and thus can detect small and early injury that might occur to the kidney.

This study was performed on 100 cirrhotic patients admitted to Intensive Care Unit in the hospital of Theodore Bilharz Research Institute. The objective of this study is to identify the association between hospital mortality and RIFLE criteria in critically ill cirrhotic patient. Other scoring systems, including Child score, SOFA score, MELD score and APACHE II score are also used for comparison with the RIFLE classification. The studied population was classified according to the primary outcome of this study (Hospital



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## **List of Abbreviations**

AFP : Alpha Feto-Protein
AKI : Acute Kidney Injury

**AKIN** : Acute Kidney Injury Network

**AIDS** : Acquired Immunodeficiency Syndrome

**Alb** : Albumin

**ALF** : Acute Liver Failure

**ALKM**: Anti-Liver Kidney and Microsomes

ALP : Alkaline Phosphatase
ALT : Alanine Transaminase
ANA : Antinuclear Antibody,

**ANCA**: Antineutophil Cytoplasmic Antibody

**ANOVA**: Analysis of Variance

**ANZICS:** Australia New Zealand Intensive Care Society **APACHE:** Acute Physiology And Chronic Health Evaluation

APD : Adult Patient DatabaseAPS : Acute Physiology ScoreARF : Acute Renal Failure

**ASMA** : Antismooth Muscle Antibody

**AST** : Aspartate Transaminase

**AUROC**: Area Under Receiver Operation Curve

**BIL**: Bilirubin

**CBC**: Complete Blood Count

**COPD**: Chronic Obstructive Pulmonary Disease

CRF : Chronic Renal FailureCT : Computed Tomography

**CTP** : Child-Turcotte-Pugh Classification

**DIC**: Disseminated Intravascular Coagulopathy

**DM** : Diabetes Mellitus

**ECMO**: Extra-Corporeal Membrane Oxygenation

**ESR** : Erythrocyte Sedimentation Rate

ESRD : End Stage Renal Disease
 F<sub>i</sub>O<sub>2</sub> : Fraction of Inspired Oxygen
 FSH : Follicle Stimulating Hormone
 GABA : Gamma-Amino-Butyric Acid

## **List of Abbreviations** (Cont.)

GCS : Glasgow Coma Scale

**GFR**: Glomerular Filtration Rate

**GGT** : Gamma Glutamyle Transpeptidase

Hb : HemoglobinHBV : Hepatitis B Virus

**HCC**: Hepatocellular Carcinoma

HCO3 : BicarbonateHct : HematocriteHCV : Hepatitis C Virus

**HDU**: High Dependency Unit

HH : Hereditary Hemochromatosis.HIV : Human Immune Deficiency Virus

**HLA**: Human Leukocyte Antigen

**HOA** : Hypertrophic Osteo-Arthropathy

**HRS**: Hepatorenal Syndrome

**HTN**: Hypertension

**ICU**: Intensive Care Unit

INR : International Normalized RatioLDLT : Living Donor Liver Transplantation

LFTs : Liver Function Tests LH : Luteinizing Hormone

**LVP** : Large Volume Paracentesis

**MAP**: Mean Arterial Pressure

MDRD : Modification of Diet in Renal DiseaseMELD : Model for End -Stage Liver Disease

MPM : Mortality Probability ModelsNASH : Non Alcoholic Steato-Hepatitis

NO : Nitric Oxide

**NSAIDs**: Non Steroidal Anti-Inflammatory Drugs

**NYHA**: New York Heart Association

**OR** : Odds Ratio

**P-ANCA**: Perinuclear Anti-Neutrophil Cytoplasmic Antibody

PaO<sub>2</sub>: Arterial Oxygen Tension PAO<sub>2</sub>: Alveolar Oxygen Tension

## **List of Abbreviations** (Cont.)

PBC : Primary Biliary Cirrhosis
 PC : Prothrombin Concentration
 PCO<sub>2</sub> : Carbon Dioxide Tension
 PCR : Polymerase Chain Reaction

**PGE2**: Prostaglandin E2

**PLT**: Platlets

**PSC**: Primary Sclerosing Cholangitis

PT : Prothrombin Time
P value : Significance Level
PVD : Portal Vein Diameter

**RCTs**: Randomized Controlled Trials

**RIFLE**: Risk-Injury-Failure-Loss of kidney function-End stage

renal disease

**ROC** : Receiver Operation Curve

**RR** : Relative Risk

RRT : Renal Replacement Therapy
 SAAG : Serum-Ascites Albumin Gradient
 SAPS : Simplified Acute Physiology Score
 SBP : Spontaneous Bacterial Peritonitis

SCr : Serum CreatinineSO2 : Oxygen Saturation

**SOFA**: Sequential Organ Failure Assesment

**S.D.** : Standard Deviation

SGOT : Serum Glutamic Oxaloacetic TransaminaseSGPT : Serum Glutamic Pyruvic Transaminase

TIPS: Transjugular Intrahepatic Portosystemic Shunt

TPO: Thrombopoietin UK: United Kingdom

**UNOS**: United Network for Organ Sharing

**UO** : Urine Output

**USA**: United States of America

**WBC's**: White Blood Cells

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## RIFLE Classification as a predictor of short term prognosis in critically ill cirrhotic patient

#### **Introduction:**

Liver cirrhosis represents the final common pathway of virtually all chronic liver disease and is characterized by an accumulation of extracellular matrix rich in fibrillar collagen(1).

A feature of liver cirrhosis is the existence of disturbances in systemic circulation characterized by marked arterial vasodilation that occurs principally in the splanchnic circulation and generates a reduction in total peripheral vascular resistance and arterial pressure and a secondary increase in cardiac output. These abnormalities are central to the development of several major complications of cirrhosis, such as hepatorenal syndrome, ascites, spontaneous bacterial peritonitis, dilutional hyponatremia, and hepatopulmonary syndrome. Renal failure is the most clinical relevant of these conditions as its appearance generally indicates very poor prognosis (2, 3, 4, 5).

Elevated serum creatinine (SCr) levels of > 1.3 or 1.5mg/dl have been identified as a predictor of poor prognosis in patients with advanced liver cirrhosis (6).

The common used scoring systems for predicting the outcome in critically ill cirrhotic patients, such as Child-Pugh score (7), Sequential Organ Failure Assessment (SOFA) (8), Model for End-stage Liver Disease (MELD) (9), and Acute Physiology, Age, Chronic Health Evaluation II (APACHE II)(10) evaluate renal function according to the serum creatinine.

The RIFLE (risk of renal failure, injury to the kidney, failure of kidney function, loss of kidney function, and end-stage renal failure) classification was first proposed by the Acute Dialysis Quality Initiative (ADQI) group at the second ADQI conference in Vicenza, Italy, in May 2002, in an attempt to standardize the study of ARF. The RIFLE criteria classify ARF into three groups (risk, injury, and failure) according to changes in SCr and urine output (UO) (11). To date, the RIFLE classification has been applied in critical ill patients receiving renal replacement therapy, cardiac surgery patients, heterogeneous patients from intensive care units (ICUs), heterogeneous population of hospitalized

patients, and unique populations such as patients requiring extracorporeal membrane oxygenation for post-cardiotomy

## Aim of the work:

cardiogenic shock(11).

The objective of this study is to identify the association between hospital mortality and RIFLE criteria in critically ill cirrhotic patient.

Other scoring systems, including Child score, SOFA score, MELD score and APACHE II score are also used for comparison with the RIFLE classification.

#### **Patients and Methods:**

This study will be performed on 100 cirrhotic patients admitted to Intensive Care Unit in the hospital of Theodore Bilharz Research Institute.

Exclusion criteria are:

- \* Pediatric patients  $\leq$  18 years of age.
- \* Uremic patients undergoing renal replacement therapy.
- \* Patients who had undergone liver transplantation.

The following data will be collected for each patient on the 1<sup>st</sup> day of admission:

- → Demographics.
- → Reason for ICU admission.
- → Acute diagnosis.
- $\rightarrow$  RIFLE classification.
- → Child score.
- $\rightarrow$  SOFA score.
- $\rightarrow$  MELD score.
- → APACHE II score.

And finally, the duration of hospitalization and the outcome of each patient will be recorded.