DISSEMINATED INTRAVASCULAR COAGULATION (DIC)
BEFORE AND AFTER OFSOPHAGEAL SCLEROTHERAPY

Thesis Submitted for Partial Fulfilment of the M.Sc. Degree in

Internal Medicine

رسالت

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LIST OF ABBREVIATIONS

ALT : Alanine transaminase

AP : Alkaline phosphatase

AST : Aspartate transaminase

AT-III : Antithrombin III

Bilharziasis or Bilharzial

Bl. urea : Blood urea

D.b : Direct serum bilirubin

DIC : Disseminated intravascular coagulation

EACA : Epsilon Amino-caproic acid

EVL : Endoscopic variceal ligation

FDPs : Firbin(ogen) degradation products

Hb : Haemoglobin

HB s Ag : Hepatitis B surface antigen

HB s Ab : Hepatitis B surface antibody

HB e Ag : Hepatitis B "e" antigen

I.V. : Intravenous

0₂ : Oxygen

PAI : Plasminogen activator inhibitor

PLT : Platelet

PT : Prothrombin time

PTT : Partial thromboplastin time

r.p.m. : Rotation per minute

S. alb. : Serum albumin

S. creatin. : Serum creatinine

S.D. : Standard deviation

T.b. : Total bilirubin

T.E. : 0.75% tetradecyl sulfate +

47% ethanol and saline

T.E.S : 1% tetradecyl sulfate +

32% ethanol and saline

TLC : Total leucocytic count

TP : Total serum proteins

TT : Thrombin time

TTD : 0.5% tetradecyl sulfate +

50 units/ml topical thrombin in 50%

dextrose

VWF : Von Willibrand factor.

> : More than

Less than

INTRODUCTION AND AIM OF THE WORK

INTRODUCTION

Haemorrhagic manifestations are a common complication and a major cause of morbidity and mortality in patients with chronic liver diseases, mainly when complicated by portal hypertension and oesophageal varices.

(Classen et al., 1986)

Recently, it has been reported that patients with chronic liver diseases, especially cirrhosis, demonstrate impairment in their fibrinogen survival as well as their fibrinolytic system. This makes these patients more prone to develop DIC on exposure to precipitating factors including sepsis, shock, hypotension, endothelium injury as well as others. (Van-Dewater et al., 1986; Carr, 1989).

Endoscopic sclerotherapy represents a therapeutic advance in the management of oesophageal varices for both the acute bleeding and the prevention of recurrent bleeding.

(Hootegem et al., 1984)

Endoscopic sclerotherapy carries a complication rate of 5-15%. These include local complications at the site of injection as well as systemic complications due to dissemination of injected material mainly with intravariceal injection. (Agres et al., 1983; Bellary and Isaacs, 1990)

Although coagulation abnormalities including DIC have been suspected following sclerotherapy, still this

has not been definitely proved, and various results have been reported by different workers.

(Hedberg et al., 1982)

AIM OF THE WORK

The aim of the present study is to examine patients with oesophageal varices subjected to endoscopic sclerotherapy for the possible development of post-sclerotherapy disseminated intravascular coagulopathy.

REVIEW OF LITERATURE