Faculty of Medicine Ain Shams University

COMPARATIVE STUDY ON THE COAGULATIVE AND BLEEDING PROFILE IN PATIENTS WITH BLEEDING OESOPHAGEAL VARICES TREATED BY INJECTION SCLEROTHERAPY VERSUS BAND LIGATION

Thesis
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Internal Medicine

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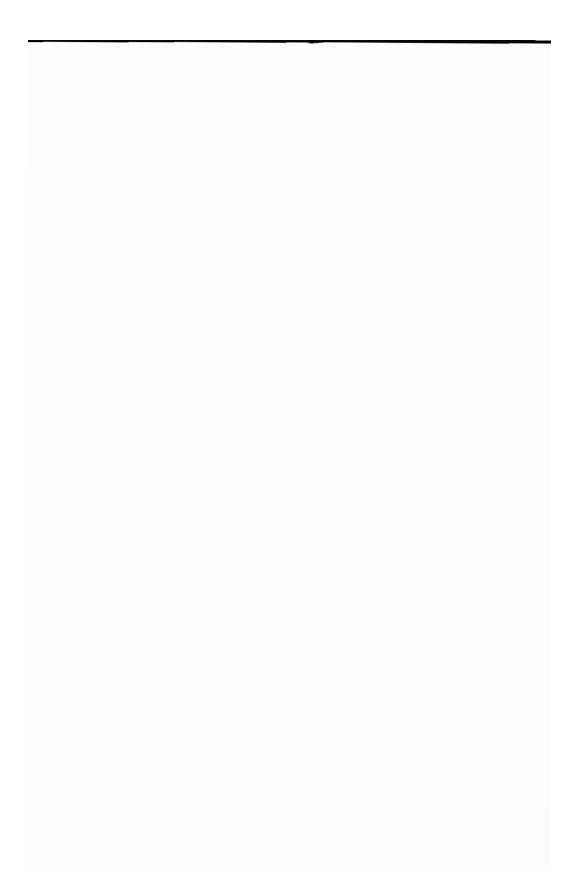
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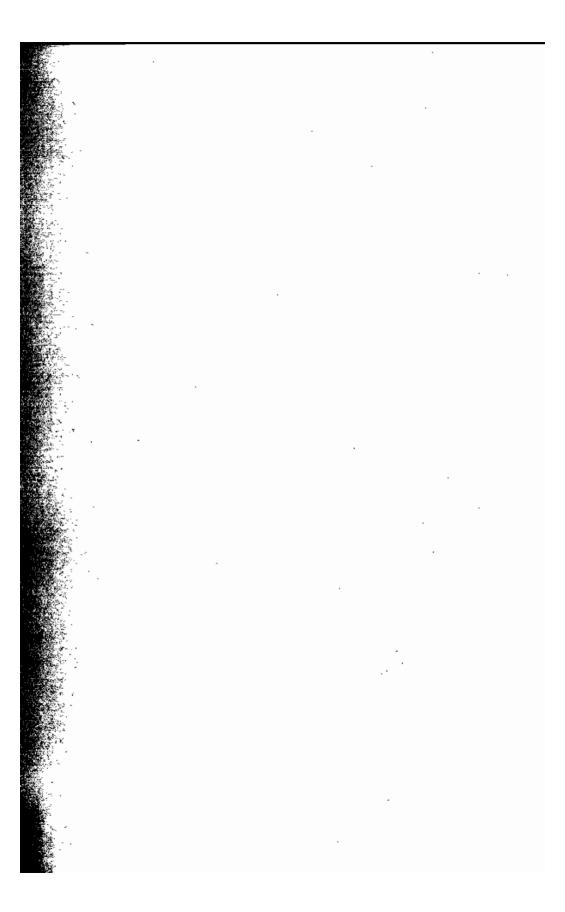
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INTRODUCTION AND AIM OF THE WORK



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Introduction

The liver is the principle site for synthesis of all coagulation proteins with the exception of Von Willebrand Factor and Fibrinolytic proteins. The liver also synthesizes protease inhibitors which modulate the coagulation cascade, such as antithrombin III, protein C and heparin co-factor II. The liver also clears activated clotting factors from the blood.

Impaired coagulation is the major haematological manifestation of liver cirrhosis. The decreased activity of any or all of these factors slows the coagulation process and, in conjunction with thrombocytopenia and various potential bleeding sites, poses a lethal threat to decompensated patients with cirrhosis (Sherlock and Dooley, 1997).

Intravariceal sclerotherapy results in transient activation of the clotting system with mild consumptive coagulopathy probably precipitated by inducing platelet agglutination and by indirect activation of contact clotting factors secondary to endothelial injury (Tang et al., 1986).

Endoscopic injection sclerotherapy did not lead to disseminated intravascular coagulation although transient activation of coagulative and fibrinolytic systems did occur.

With a significant decrease in platelet count, and almost complete reversion was noted about 18 hours after endoscopic injection sclerotherapy (Bernardi et al., 1984).

Platelet aggregation after intravariceal injection of 5% ethanolamine oleate was significantly suppressed by a humoral response (Yamaga et al., 1989).

Aim of the Work

To compare the systemic coagulopathy and bleeding profile following sclerotherapy of oesophageal varices with 5% ethanolamine oleate versus that following band ligation.