

شبكة المعلومات الجامعية







شبكة المعلومات الجامعية التوثيق الالكتروني والميكروفيلم



شبكة المعلومات الجامعية

جامعة عين شمس

التوثيق الالكتروني والميكروفيلم

قسم

نقسم بالله العظيم أن المادة التي تم توثيقها وتسجيلها على هذه الأفلام قد أعدت دون أية تغيرات



يجب أن

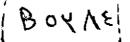
تحفظ هذه الأفلام بعيدا عن الغبار في درجة حرارة من ١٥-٥٠ مئوية ورطوبة نسبية من ٢٠-٠٠% To be Kept away from Dust in Dry Cool place of 15-25- c and relative humidity 20-40%



بعض الوثائـــق الإصليــة تالفــة



بالرسالة صفحات لم ترد بالإصل



EVALUATION OF SOME DIAGNOSITC AND THERAPEUTIC ASPECTS OF PORTAL HYPERTENSION IN CHILDREN

Thesis
Submitted in Partial Fulfillment of the M.D. Degree in Pediatrics

Ву

MOHIEE EL-DEEN ABD EL-AZIZ AWAD

(M.B.,B.CH.)

(M.Sc. Pediatrics)

Assistant Lecturer of Pediatrics
Tanta Faculty of Medicine

SUPERVISORS

Prof. Dr.

MOHAMED A. HAMAM

Prof. and Head of Pediatric Dep.

Faculty of Medicine Tanta University

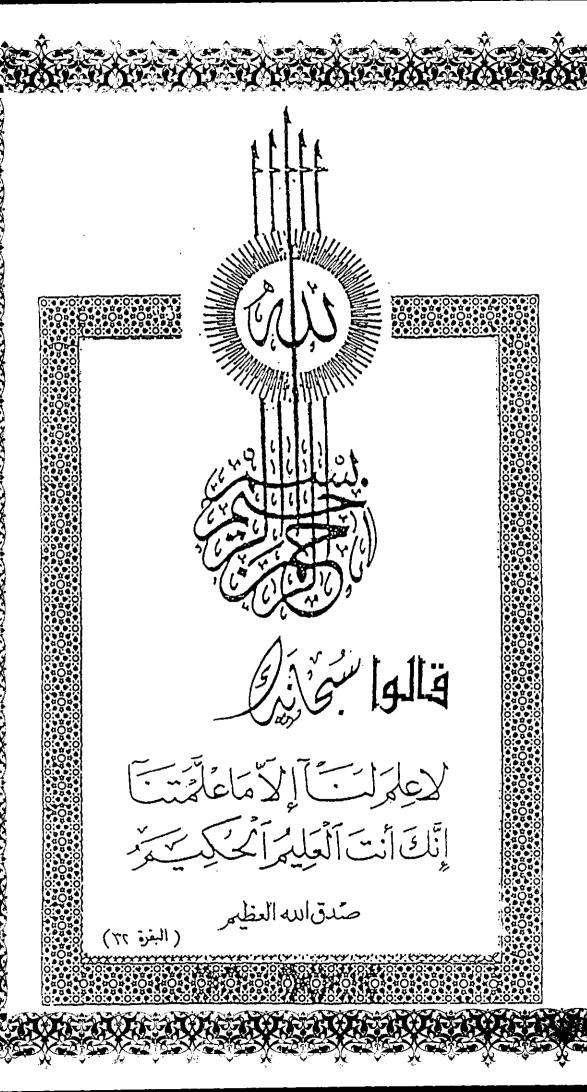
PROF. DR.
MOHAMED M. KANTOOSH

Prof. of Pediatrics Faculty of Medicine Tanta University. PROF. DR.
SHERIF LOTFY BAYOUMI

Prof. and Head of Pathology Dep.
Faculty of Medicine
Tanta University.

FACULTY OF MEDICINE TANTA UNIVERSITY

1998



ABBREVIATIONS

PH= Portal Hypertension

PVS=Portal Venous System

PVF= Portal Venous Flow

PVP= Portal Venous Pressure

PV= Portal Vein

SV= Splenic Vein

SMV= Superior Mesenteric Vein

SMA= Superior Mesenteric Artery

IVC=Inferior Vena Cava

WHVP=Wedged Hepatic Venous Pressure

HVPG= Hepatic Venous Pressure Gradient

US= Ultrasonography, Ultrasound, Ultrasonic

PVT= Portal Vein Thrombosis

PHG= Portal Hypertensive Gastropathy

BFV=Blood Flow Velocity

GIT=Gastrointestinal Tract

IV= Intravenous

MCL= Mid Clavicular Line

ML = Mid Line

LS=Longitudinal Section

TS=Transverse Section

Bilh.= Bilharzial

ALT=Alanine Aminotransferase

AST=Aspartate Aminotransferase

PT=Prothrombin Time

R.B.Cs= Red Blood Corpuscles

W.B.Cs=White Blood Cells

Fig.= Figure

Tab.=Table

ACKNOWLEDGEMENT

First and foremost, thanks are due to ALLAH.

I would like to express my deepest sincere thanks and gratitude to Prof. Dr. **MOHAMED A. HAMAM,** Prof. and Head of Pediatric Dep., Faculty of Medicine, Tanta University, for proposing the subject of this thesis, his unlimited patience, meticulous discussion, sound explanation and sincere help during the performance and writing of this thesis.

I wish also to express my deepest sincere thanks and gratitude to Prof. Dr. MOHAMED M. KANTOOSH, Professor of Pediatrics, Faculty of medicine, Tanta University for his great care, continuos supervision, scientific criticism and kind guidance during this study.

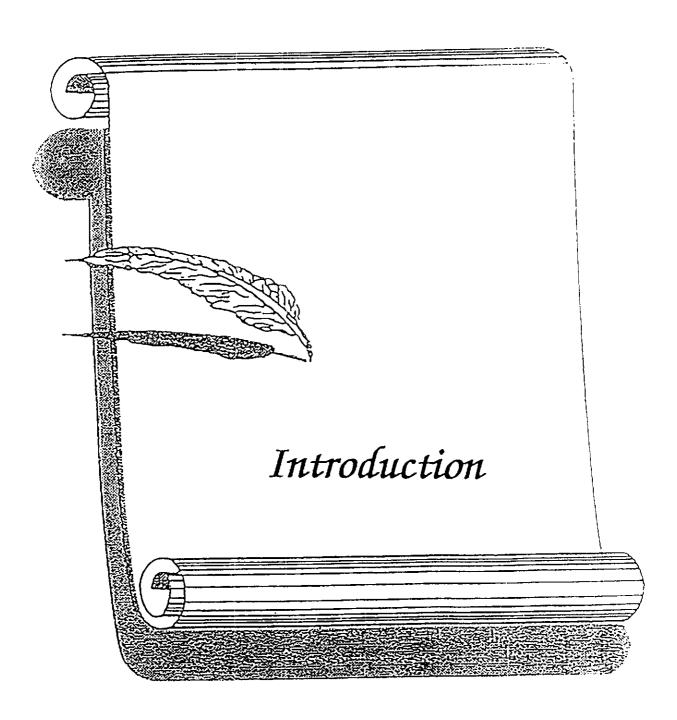
It is my pleasure indeed to express my deep gratitude and thanks to Prof. Dr. SHERIF LOTFY BAYOUMI, Prof. and Head of Pathology Dep., Faculty of Medicine, Tanta University for his valuable efforts and assistance during performance of the laboratory procedures and during writing of this thesis.

I wish also to express my deep gratitude and appreciation to Dr. Saleh Amin, Lecturer of Pediatrics, Faculty of Medicine, Tanta University, for his kind support and sincere help during the work and during writing of this thesis.

Also, my sincere thanks to all members of our department and every one who helped and advised me during this work.

CONTENTS

INTRODUCTION	1
REVIEW OF LITERATURE	••••
-Anatomy and circulation of the portal venous system	3
-Factors regulating portal venous flow and pressure	
-Portal Hypertension *Definition and etiology	16
*Collateral circulation	
*Clinical picture	
*Methods of portal pressure measurements.	
-Ultrasonography of portal venous system	
-Esophageal changes in portal hypertension	
-Stomach and gastric changes in portal hypertension	
-Liver biopsy in portal hypertension	
-Long-term control of portal hypertension	
AIM OF THE WORK	115
SUBJECTS AND METHODS	116
RESULTS	132
DISCUSSION	194
SUMMARY AND CONCLUSIONS	215
REFERENCES	222
ARABIC SUMMARY	



Introduction

INTRODUCTION

Portal hypertension in children is a serious problem because it is an important cause of gastrointestinal bleeding especially from esophagogastric varices (1).

The documentation of portal hypertension and the resulting collaterals by invasive and time-consuming procedures such as direct venoportography or arteriography can result in significant morbidity among patients with abnormal coagulation tests and other medical problems (2).

The hazards of such invasive investigations have stimulated some investigators (3,4,5) to study the portal venous hemodynamics by measuring the portal and splenic vein diameters as well as blood velocity and flow in these vessels utilizing recent non invasive real-time and Doppler ultrasonography.

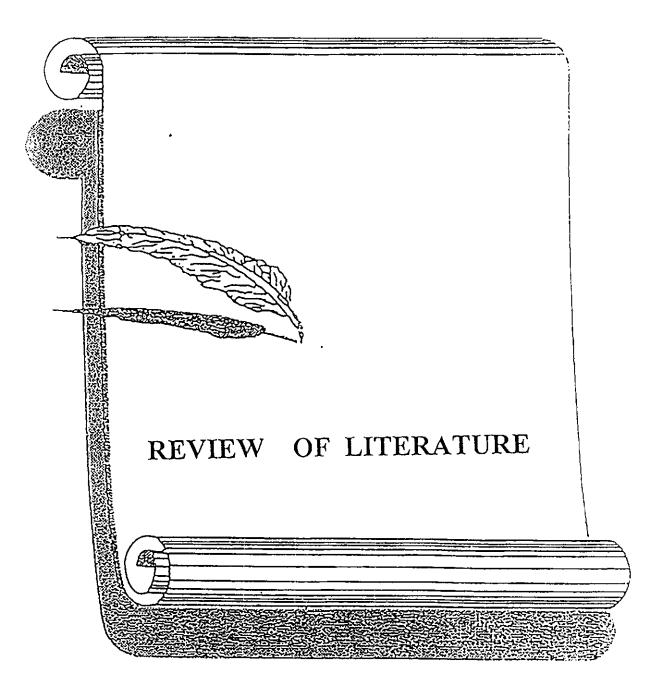
Upper gastrointestinal tract endoscopy is the best method of visualizing esophageal and gastric vasrices(6). The use of endoscopy for diagnosis and sclerotherapy of bleeding varices combined with regular endoscopic follow-up has provided a unique opportunity to study the progression of changes occurring in the portal hypertensive gastric mucosa(7).

Gastric mucosal lesions are common in portal hypertension and are an important source of bleeding(8). The nature of portal hypertensive gastric mucosa varied among different studies in adults(9,10), but there are few studies reported on gastric mucosal lesions in infants and children with portal hypertension.

Introduction

Reduction of portal hypertension could be achieved by continuous medical treatment. Different agents that effectively lower portal pressure are available. Of these, propranolol a non-selective β -blocker provoked considerable interest among several investigators (11,12). However, the effect of propranolol therapy on portal hypertension and the resulting gastropathy in children was reported by a few investigators (13).

The present work was conducted to show some light on these important diagnostic and therapeutic aspects of portal hypertension in children.



Review of literature

THE ANATOMY OF THE PORTAL VENOUS SYSTEM

The portal venous system (PVS) includes all veins draining the abdominal part of the alimentary tract, spleen, pancreas, and gall bladder. The portal vein (PV) is formed by union of the superior mesenteric vein (SMV) and splenic vein (SV) just posterior to the neck of the pancreas. At the liver hilum, the PV divides into two major branches one to each lobe (right and left branches) (14). These major branches undergo several intrahepatic segmental divisions which supply the liver in a segmental manner and terminate in small branches which pierce the limiting plate of the portal tracts and enter the adjacent sinusoids through short channels (15). After passing to the sinusoids, the hepatic blood is collected to the inferior vena cava (IVC) by the hepatic venous system (14).

The tributaries of the extrahepatic portion of the PV (Fig.1) include:

- 1) splenic vein (SV)
- 2) superior mesenteric vein (SMV)
- 3) left gastric vein (coronary vein) 4) right gastric vein
- 5)paraumbilical veins
- 6)cystic vein

Fig. (2) shows a diagram of the histological structure of the normal human liver demonstrating the territories of the classic hepatic lobules and the portal vein territories which are shared by branches of the hepatic artery and tributaries of the hepatic duct (16).