

بسم الله الرحمن الرحيم





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شبكة المعلومات الجامعية التوثيق الالكتروني والميكروفيلم



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Risk Factors and Predictors of Occlusion of Stent of Trans Jugular Intrahepatic Porto-systemic Shunt (TIPSS) in Egyptian Patients with Primary Budd Chiari Syndrome

Thesis

Submitted for Partial Fulfillment of Master Degree in **Tropical Medicine**

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2021



سورة البقرة الآية: ٣٢

Acknowledgments

First and foremost, I feel always indebted to **Allah** the Most Beneficent and Merciful.

I wish to express my deepest thanks, gratitude and appreciation to **Prof. Dr. Fatma Ahmed Ali El-Din**, Professor of Tropical Medicine, Faculty of Medicine, Ain Shams University, for her meticulous supervision, kind guidance, valuable instructions and generous help.

Special thanks are due to **Prof. Dr. Mohamed El-Ghareeb Abo Elmaaty**, Professor of Intervention
Radiology, Faculty of Medicine, Ain Shams
University, for his sincere efforts, fruitful
encouragement.

I am deeply thankful to **Dr. Heba Rashad Atteya**, Lecturer of Tropical Medicine, Faculty of
Medicine, Ain Shams University, for her great help,
outstanding support, active participation and
guidance.

I would like to express my hearty thanks to all my family for their support till this work was completed.

Ibrahim Abd El-Hakim Ibrahim Hindy Dawoud

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Tist of Abbreviations

Abb.	Full term
AID	A 11
ALB	
	Alanine Aminotransferase
	Angiogenic myeloid metaplasia
	Antiphospholipid syndrome
	Aspartate Aminotransferase
	Budd Chiari syndrome
BD	
	Blood urea nitrogen
	Contrast induced nephropathy
	Chronic myclogenous leukemia
CRP	C-Reactive Protein
CT	Computed tomography
DVT	Deep venous thrombosis
EASL	European Association for the Study of the
	Liver
ESR	Erythrocyte Sedimentation Rate
	Essential thrombocytosis
	Factor V leiden mutation
HBG	Hemoglobin
	Hepatocellular carcinoma
	Hepatic encephalopathy
	Hepatic vena cava budd chiari syndrome
	Hepatic venous outflow tract obstruction
	International normalized ratio
	Inferior vena cava
	Model for end stage liver disease
MF	
	Mean pulmonary artery pressure
	Myeloproliferatine diseases
	Myloproliferative neoplasms
	Magnetic resonance
	Methylenetetrahydrofolate Reductase
1,1111111111111111111111111111111111111	Deficiency
	20110101109

Tist of Abbreviations cont...

Abb.	Full term
PGM	.Progthrombin G mutation
PLT	.Platelets
PNH	.Paroxysmal noctural hemoglobinuria
PRV	.Polycythemia rubra vera
PV	.Portal vein
PVT	.Portal vein thrombosis
TIPSS	.Transjugular intrahepatic portosystemic
	shunt
US	
WBC	.White blood cells

ABSTRACT

Budd Chiari syndrome is a rare disorder which is characterized by hepatic venous outflow tract obstruction. The classical triad of presentation are ascites, hepatomegaly and abdominal pain.

Early diagnosis and proper treatment improve the overall survival and decreases morbidity and mortality, anticoagulation is main line of treatment, angioplasty, stenting and TIPSS provides way of decompression of the congested liver.

TIPSS is used as the most common line of treatment in treating BCS patients with good outcome in case of good patient selection.

TIPSS occlusion remains a major problem after intervention which require regular follow up and shunt revision for proper identification and restoring patency.

The prevalent thrombophilia etiology in the studied group was FVLM (69.7%) then MTHFRD (67.1%).

The overall rate of stent occlusion in 6 months among the patients was 50 % with highest rate of occlusion at 7 days post TIPSS and 3 months.

After analyzing the results, we found that there is no significant relation between age, sex and TIPSS stent occlusion. Different etiologies of BCS may be a risk factor for stent occlusion. Patients with AT III deficiency and PRV carried a higher rate of occlusion. Also cirrhotic and coarse liver by ultrasound may predict stent occlusion.

Keywords: Occlusion, Trans Jugular Intrahepatic Porto-systemic Shunt, Egyptian Patients, Primary Budd Chiari Syndrome

Introduction

Budd-Chiari syndrome (BCS) is a rare disorder caused by obstruction of the hepatic venous outflow tract at any level between the small hepatic veins and the right atrium, hence also known as hepatic venous outflow tract obstruction (HVOTO). This obstruction leads to venous stasis resulting in congestive hepatopathy (*Mukund & Gamanagatti*, 2011).

Congestive hepatopathy results in increased sinusoidal pressure with hepatic sinusoidal thrombosis as evidenced by fibrin deposition within sinusoids. This sinusoidal thrombosis in turn causes reduced hepatic perfusion leading to ischemia and necrosis of hepatocytes especially in perivenular zones leading to fibrosis and portal hypertension (*Simonetta et al.*, 2015).

Budd-Chiari syndrome (BCS) may result due to primary venous problem or may be secondary to compression/invasion of venous outflow by a space occupying lesion within or outside the liver (*Van Wettere et al.*, 2018).

Primary BCS is often related to multiple thrombophilic conditions, including primary myeloproliferative disorders, paroxysmal nocturnal haemoglobinuria, inherited thrombophilia (i.e., deficiencies of natural anticoagulants and mutations of factor V Leiden and Factor II), antiphospholipid antibodies, and circumstances at risk, such as pregnancy (*Valla*, *2018*).