# A study of the effect of intraperitoneal chemohyperthermia versus intraperitoneal chemotherapy in patients with malignant ascites

#### THESIS

Submitted in partial fulfillment for the requirement of M.D degree in Tropical Medicine

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#### Acknowledgement

greatly honored to work under the supervision of Prof. Dr. Laila **Ahmed Mohammed** Professor of Tropical Medicine. Faculty of Medicine, Cairo University. She has given me much of her time, her guidance, advice and scientific attitude which help to make this work come to light. No words can be sufficient express my deep gratitude and indebtedness to her.

It was a great pleasure to record my sincere thanks and gratefullness **to Prof. Dr. Hosny Salama** Professor of Tropical Medicine, Faculty of Medicine, Cairo University. His distinguished care, supervision and noble attitude could not be expressed in words.

I would like to express my thanks and deepest gratitude to **Prof. Dr. Farouk Ahmed Haggag** Professor of Oncology, Faculty of Medicine, Cairo University, for his continuous guidance, help and support throughout this work.

My deepest gratitude goes to **Prof. Dr. Serag Zakaria** Head of Tropical Medicine Department, Faculty of Medicine, Cairo University, for his continuous support and faithful guidance

I wish to express my great appreciation and sincere thanks to **Prof. Dr. Hanan Abd El Halim Marzouk** Assistant Professor of Tropical Medicine, Faculty of Medicine. Cairo University. for her valuable suggestions and continuous support which make the accomplishment of this work possible.

I would like to express my sincere thanks to **Prof. Dr. Hany khatab** Professor of Pathology, Faculty of Medicine. Cairo University, for his great help support and guidance.

My thanks should extend to all my eminent professors and staff members in Tropical Medicine Department with special thanks to **Prof. Dr. Sohair Zakaria**, whom I am very proud to gain knowledge at her hands and whose kindness and support cannot be denied.

Special thanks to my friends and colleagues in Tropical Medicine Department for their generous encouragement and support.

I wish I could express my profound gratitude and sincere thanks to my family for their encouragement, goodness and kindness. I dedicate this work to them hoping that they would be proud of me.

My deepest thanks goes to the patients who are suffering, hoping that this work will give them hope for a better future with a good health.

#### **Abstract**

Malignant ascites is one of the most serious complications of many malignant tumours. Many treatment modalities have been tried to eradicate cancer cells in ascitic fluid. In this study we compared effect intraperitoneal the of chemohyperthermia intraperitoneal chemotherapy in patients with malignant ascites. .We compared their effects on ascitic fluid cytology, quality of life and intraperitoneal progression. We concluded that disease chemohyperthermia can affect the viability of malignant cells in with minimal with malignant ascites adverse effects. patients Moreover intraperitoneal chemohyperthermia exerts better effects than intraperitoneal chemotherapy alone. We have also shown that quality of life in our patients has improved as patients experienced some relief of ascites related symptoms with regression of ascites.

Key Words: - Hyperthermia

- Intraperitoneal

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### List of abbreviations

•	ADM	Adriamycin.
•	AMSA	American Medical Student Association.
•	Ara-C	Cytarabine.
•	b-FGF	basic-fibroblastic growth factor.
•	BLM	-Bleomycin.
•	CA-125	Cancer antigen-125.
•	CCS	Completeness of cytoreduction score.
•	CEA	-Carcinoembryonic antigen.
•	CT	-Computed tomography.
•	DACH	-Diamino cyclohexane.
•	DNA	-Deoxynucleic acid.
•	DPAM	-Disseminated peritoneal adenomucinosi.
•	DTIC direct intra ar	terial chemotherapy.
•	EPIC	Early postoperative intraperitoneal chemotherapy.
•		Food and Drug Administration.
•		Federation of Gynecology and Obstetrics.
•	5FU	5-florouracil.
•		Gastrointestinal tract.
•	HIICHyperthe	rmic intraoperative intraperitoneal chemotherapy.
•	HIPECIntraoper	rative hyperthermic intraperitoneal chemotherapy.
•	HMB45	Mouse monoclonal antibody.
•		Investigational Device Exemption
•	IHCP	- Intraperitoneal hyperthermic chemoperfusion.
•	IL-1	
•	IL-2	
•	IP	Intraperitoneal
•		Intraperitoneal chemohyperthermia.
•	ISM	Industrial, scientific, and medical.
•	IV	
•	L-PAM	L-phenylalanine mustard.
•		Lactate dehydrogenase.
•	LS	
•		Long-term intraperitoneal chemotherapy.
•	MMC	Mitomycin C.

• Mo-----Month.

•	• NCI	National Cancer Institute.
•	NIH	National Institute of Health.
•	NK	Natural killer cells.
•	PC	Peritoneal cancer.
•	• PCIP	Peritoneal Cancer Index.
•	• PMCAP	eritoneal mucinous carcinoma.
•	PVSPe	eritoneovenous shunting.
•	• RNA	Ribonucleic acid.
•	SPCI	Simplified Peritoneal Carcinomatosis.
•	TNF- α	Tumour necrosis factor alpha.
•	TNFT	Sumour necrosis factor.
•	UCHL-1U	Jbiquitin carboxyl-terminal esterase L1.
•	• VEGFV	ascular endothelial growth factor.

• VPF-----Vascular permeability factor/

# Introduction

#### INTRODUCTION

Malignancy is the cause of ascites in 10% of patients who present with it (*Runyon.*, 1994).

Malignant ascites most commonly results from peritoneal "studding" and occlusion of the diaphragmatic lymphatics leading to impaired reabsorbtion of abdominal fluid (Zervos & Rosemurgy, 2001). Malignant ascites may also result from inferior vena cava or hepatic vein obstruction secondary to primary or metastatic tumor infiltration of the liver; but this is less common (Zervos & Rosemurgy, 2001). Ovarian cancer is by far the most common cause of malignant ascites, accounting for almost 50% of all cases (Ringenberg et al., 1989). Occult malignancy accounts for an additional 20% of the cases and pancreatic, gastric, colon, lung, breast, and lymphoma comprise the remaining cases (Ringenberg et al., 1989).

When malignant ascites is suspected, every attempt should be done to document the nature and source of the ascites utilizing radiographic studies, cytology, fluid studies, and tumor markers (Zervos & Rosemurgy, 2001). Computed tomography scan should be obtained early because it is helpful not only in assessing the origin and extent of the malignant process but also the degree of peritoneal involvement (Zervos & Rosemurgy, 2001). Cytology may direct the workup, as do tumor markers in the case of occult or unknown primary processes, the most useful of which is CA-125 in women with suspected ovarian cancer (Zervos & Rosemurgy, 2001). Fibronectin levels and serum protein to ascites ratio are biochemical markers