

سورة البقرة الأية: ٣٢

## Mini Dose Versus Large Dose Spinal Anesthesia For Laparoscopic Cholecystectomy

### **Ehesis**

Submitted for the Partial Fulfillment of MD in Anesthesiology

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#### **List of Abbreviations**

ASA : American society of anesthesiologist

**ACTH** : Adreno corticotrophic hormone

**ADH** : Antidiuretic hormone

**AVP** : Arginine vasopressor

BMI : Body mass index

**COPD** : Chronic obstructive pulmonary disease

CI : Cardiac index

CO : Cardiac output

CO<sub>2</sub> : Carbon dioxide

**CSE** : Combined spinal epidural

**CSF** : Cerebro-spinal fluid

**CVP** : Central venous pressure.

**DVT** : Deep venous thrombosis

**ECG** : Electro-cardiogram

**EtCO<sub>2</sub>** : End tidal carbon dioxide

FRC : Functional residual capacity

GA : General anesthesia

GB : Gall bladder

IAP : Intra-abdominal pressure

**ICP** : Intra-cranial pressure

**IOP** : Intra-ocular pressure

**IPPV** : Intermittent positive pulmonary ventilation

IV : Intravenous

IVC : Inferior vena cava

LDSA : Large-dose spinal anesthesia

LDS-group : Large dose spinal anesthesia group

LC : Laparoscopic cholecytectomy

LMA : Laryngeal mask airway

MAC : Minimum alveolar concentration

MAP : Mean arterial pressure

MB : Motor block

MBP : Mean Blood Pressure

MBS : Modified Bromage scale

MDSA : Mini-dose spinal anesthesia

MDS-group: Mini dose spinal anesthesia group

**MV** : Minute volume

NIBP : Non-invasive blood pressure

NMDA : N-methyl-D- aspartate

NSAIDs : Nonsteroidal Anti-inflammatory Drugs

PCA : Patient controlled analgesia

PCWP : Pulmonary capillary wedge pressure

PDPH : Postdural puncture headache

PO : Postoperative

PONV : Post-operative nausea and vomiting

PPP : Positive pressure pneumoperitoneum

**PVR** : Pulmonary vascular resistance

PaCO<sub>2</sub> : Partial pressure of carbon dioxide in arterial

blood

PeCO<sub>2</sub> : Partial pressure of carbon dioxide in expired air

RAP : Right atrial pressure

SA : Spinal anesthesia

SB : Sensory block

SBP : Systolic blood pressure

**SpO**<sub>2</sub> : Arterial oxygen saturation

**SVR** : Systemic venous resistance

TIVA : Total intravenous anesthesia

TV : Tidal volume

VC : Vital capacity

V/Q : Ventilation/Perfusion

VR : Venous return

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### Introduction

Laparoscopic cholecystectomy was first introduced by Phillipe Mouret in 1987 and is now generally performed by many surgeons (*Vecchio et al., 2000*). Laparoscopic cholecystectomy has become firmly established as the method of choice for the treatment of asymptomatic gall stones because of obvious advantages of minimally invasive character of the procedure associated with less pain, reduced hospital stay, and earlier return to daily activities (*Savas et al., 2004*).

However, considerable difficulties in anesthetic management could be encountered as wide hemodynamic fluctuation may develop due to pneumoperitoneum and position changes during the operation. Also, serious complications such as tissue damage or air embolism could occur (*Pursani et al.*, 1998).

Laparoscopic cholecystectomy is classically performed under general anesthesia to prevent aspiration and respiratory problems due to the pneumoperitoneum (Savas et al., 2004). Laparoscopic cholecystectomy done under spinal anesthesia may have several advantages over laparoscopic surgery done under general anesthesia (Sinha et al., 2008).

#### Introduction

Since the patient is awake, there is early detection of complications. Also it provides excellent post-operative analgesia with a lower incidence of post-operative nausea and vomiting. However, extensive sensory block (T4) is required to abolish the discomfort of manipulation of the upper gastrointestinal structures (*Sood & Kumara*, 2003).

### **Aim of the Work**

This study aimed at comparing the effect of mini dose spinal anesthesia (MDSA) versus large dose spinal anesthesia (LDSA) on intraoperative hemodynamics and blood gases changes in patients undergoing laproscopic cholecystectomy. It also compared the motor effect, the duration of anesthesia and the hospital stay in both procedures.

## Neuroanatomy of Extrahepatic Biliary System

The innervation of the gall bladder (GB) in humans involved three routes: the anterior and posterior hepatic plexus and the phrenic nerves (Yi et al., 2005).

### Via the anterior hepatic plexus

The hepatic division of the vagus arises from the anterior vagal trunk, runs through the hepatogastric ligament near the edge of the liver (caudal liver), and joins the anterior hepatic plexus in the hepatoduodenal ligament. The plexus, containing parasympathetic and sympathetic fibers arises from the celiac plexus and around the common hepatic artery, then sends some branches to the GB via the deep and superficial branches of the cystic artery, which are distributed in the peritoneal aspect of the GB and the site of attachment of this organ in the bed between the GB and hepatic artery (figure 1) (Yi et al., 2005).

On the other hand, the anterior hepatic plexus sends some branches directly to the GB along the cystic duct, i.e., forward to the neck, body, fundus of the GB. However, the branches of anterior hepatic plexus is concentrated mainly in the cystic duct and the neck of the GB compared with the

#### Review of Literature

body and fundus of the GB. However, it was not observed that the branches arising from the hepatic division of the vagus were sent directly to the GB (Yi et al., 2005).

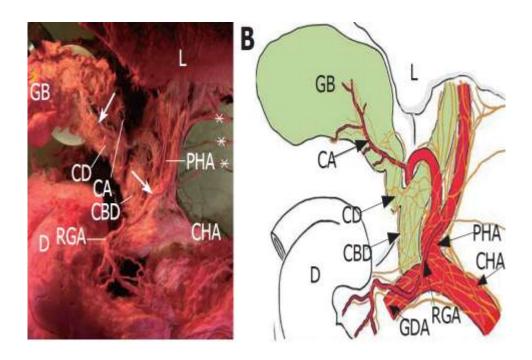


Figure (1): Innervation of the gallbladder (GB) from the ventral aspect (A) in a cadaver and a schematic representation of it (B) (Yi et al., 2005).

The branches innervating the GB originate from the anterior hepatic plexus, and run along the cystic duct (CD) and the cystic artery (CA). The hepatic divisions of the vagus join in the anterior hepatic plexus in the proper hepatic artery (PHA). Arrows indicate nerve branches. CBD: common bile duct; CHA: common hepatic artery; D: duodenum; GDA: gastroduodenal artery; L: liver; RGA: right gastric artery.

### Via the posterior hepatic plexus

The posterior hepatic plexus or the dorsal hepatic plexus arises from the celiac plexus on its right side and runs along the dorsal side of celiac plexus to the portal vein. The posterior hepatic plexus is composed of 4-5 nerve fascicles, divided into two groups of nerve bundles, and the thickest branches, about 80% of the nerve fibers, extend along the upper part of the common bile duct and portal vein, join the liver and the GB, or descend, sending branches to the proximal side of the descending part of the duodenum and the lower common bile duct. Abundant communicating rami behind the common bile duct and portal vein between the ascending and descending plexuses are observed, showing the existence of direct bidirectional neural connections between the duodenal papilla and the biliary tract containing the GB (*Yi et al.*, 2005).

As in the case of the anterior hepatic plexus, the branches sent to the GB are distributed mainly in the cystic duct and the neck of the GB. Moreover, abundant communicating rami are seen between the anterior and posterior hepatic plexuses around the cystic duct (*Yi et al.*, 2005).