

**THE EFFECT OF ADDING MAGNESIUM SULPHATE,  
DEXMEDETOMIDINE AND KETAMINE TO EPIDURAL  
ANESTHESIA FOR LOWER URINARY TRACT  
SURGERIES**

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# Contents

Acknowledgement .....	I
Contents.....	II
List of Abbreviations .....	IV
List of Figures.....	VII
List of Tables.....	VIII
Abstract.....	IX
Introduction.....	1
Chapter (1): <i>Anatomy of epidural space</i> .....	3
Chapter (2): <i>Physiological effects of neuraxial block</i> .....	14
Chapter (3): <i>Pain physiology</i> .....	17
Chapter (4): <i>Epidural anesthesia</i> .....	26
Chapter (5): <i>Pharmacology</i> .....	51

Patients and Methods.....	68
Results.....	73
Discussion.....	81
References .....	86
English Summary .....	98

## **List Of Abbreviations**

<b>ACTH .....</b>	<b>Adrenocorticotropic Hormone</b>
<b>ADH .....</b>	<b>Antidiuretic Hormone</b>
<b>ASA .....</b>	<b>American Society of Anesthesiologists</b>
<b>C2.....</b>	<b>Second Cervical vertebra</b>
<b>C7.....</b>	<b>Seventh Cervical vertebra</b>
<b>CSF .....</b>	<b>Cerebro Spinal Fluid</b>
<b>CNS .....</b>	<b>Central Nervous System</b>
<b>CV .....</b>	<b>Closing Volume</b>
<b>ECG .....</b>	<b>Electrocardiogram</b>
<b>FRC.....</b>	<b>Functional Residual Capacity</b>
<b>GABA.....</b>	<b>Gamma-aminobutyric Acid</b>
<b>ICP.....</b>	<b>Increase intracranial pressure</b>
<b>ICU .....</b>	<b>Intensive Care Unit</b>
<b>IM.....</b>	<b>Intramuscular</b>
<b>INR .....</b>	<b>International Normalized Ratio</b>
<b>IV.....</b>	<b>Intravenous</b>
<b>LMWH.....</b>	<b>Low-Molecular-Weight Heparin</b>
<b>Mg<sup>2+</sup> .....</b>	<b>Magnesium ion</b>

<b>MgSO<sub>4</sub></b> .....	<b>Magnesium Sulphate</b>
<b>MPQ</b> .....	<b>McGill Pain Questionnaire</b>
<b>NMDA</b> .....	<b>N-Methyl-D-aspartate</b>
<b>NSAIDS</b> .....	<b>Non Steroidal Anti-inflammatory Drugs</b>
<b>PONV</b> .....	<b>Postoprative nausea &amp; vomiting</b>
<b>PCA</b> .....	<b>Patient Controlled Analgesia</b>
<b>PT</b> .....	<b>Prothrombin Time</b>
<b>PTT</b> .....	<b>Partial Thromboplastin Time</b>
<b>VAS</b> .....	<b>Visual Analogue Scale</b>

## List Of Figures

<b>Fig</b>		<b>Page</b>
1	Anatomy of the sacral hiatus and dorsum of the sacrum	5
2	Normal curvature of the vertebral column	6
3	A lumbar vertebra in (a) lateral and (b) antero-superior views	7
4	Sagittal section of the lumbar region	8
5	Ascending pain pathways	19
6	Pain measurement scales	21
7	Surface landmarks for identifying spinal levels	27
8	Sitting position for neuroaxial blockade	28
9	The effect of flexion on adjacent vertebrae	29
10	Lateral decubitus position for neuroaxial blockade	29
11	Lumbar epidural anesthesia; midline approach	31
12	Thoracic epidural anesthesia; paramedian approach	32
13	Technique of paramedian approach	33
14	Sensory dermatomes	35
15	Epidural needles	41
16	Epidural needles with catheter assortment	42
17	Epidural catheters	43
18	Onset of block (mean $\pm$ SD)	76
19	Time to two-segment regression (mean $\pm$ SD)	76
20	Total dose of bupivacaine (mean $\pm$ SD)	77
21	Onset of postoperative pain (mean $\pm$ SD)	77
22	Recovery of motor power (mean $\pm$ SD)	78

## **List Of Tables**

<b>Table</b>		<b>Page</b>
1	General comparison between the four types of Neuraxial blocks	16
2	Guidelines for epidural placement and removal during anticoagulant therapy	38
3	Contraindications to neuroaxial blockade	40
4	Recommended location of catheter insertion for surgical procedures	48
5	Complications of neuroaxial blockade	49
6	Incidence of serious complications from spinal and epidural anesthesia	50
7	Demographic data	73
8	Study Parameters	75
9	MAP measurements throughout the study period	79
10	Heart rate measurements throughout the study period	80
11	Side effects throughout the study period	82

## **Abstract**

Epidural anesthesia is a safe and inexpensive technique with the advantage of providing surgical anesthesia and prolonged postoperative pain relief. It is also an effective treatment of operative pain as it blunts autonomic, somatic and endocrine responses. This study showed the effect of adding magnesium sulphate, dexmedetomidine and ketamine to epidural bupivacaine in anesthesia for lower urinary tract surgeries, with respect to onset of action, potency, duration of the block and the total dose of bupivacaine consumption during surgery as well hemodynamic changes.

**Methods:** 80 patients, ASA I/II were enrolled into the study. All patients had epidural anesthesia and randomly allocated into one of four groups. **Group C:** (20 patients) The patient received 10 ml of bupivacaine 0.5%, **Group M:** (20 patients): The patient received 10 ml of bupivacaine 0.5% plus 1.0 ml of magnesium sulphate 10% (100 mg) , **Group D:** (20 patients): The patient received 10 ml of bupivacaine 0.5% plus dexmedetomidine in a dose of 1 µg/kg diluted in 1 ml normal saline and **Group K:** (20 patients) The patient received 10 ml of bupivacaine 0.5% plus ketamine in a dose of 0.3 mg/kg diluted in 1 ml normal saline. In all groups assessment was done after 10 minutes aiming to achieve T10 sensory level and G3 motor block. Otherwise incremental doses of 5 ml bupivacaine 0.5% were given without additives.

**Results:** This study demonstrated that a more rapid onset of action of the epidural block was achieved in the magnesium group, while in the ketamine and dexmedetomidine groups, a more prolonged duration of action was recorded. In all groups, total dose consumption of the local anesthetic was lower than that of the control group. Hemodynamic variables were stable and there were no significant differences between groups.

**Conclusion:** Magnesium, ketamine and dexmedetomidine are effective as useful adjuvants to local anesthetic for epidural anesthesia. Magnesium sulphate is associated with a shorter onset of action of the epidural block. While both ketamine and dexmedetomidine have a prolonged duration and an increased potency of the block. All the studied drugs effectively decreased the total dose requirements of local anesthetic drugs. No significant side effects were observed throughout the study period.

**Keyword:** ADH, GABA, DEXMEDETOMIDINE, KETAMINE

## **INTRODUCTION**

Epidural anesthesia is a safe and inexpensive technique with the advantage of providing surgical anesthesia and prolonged postoperative pain relief. It is also an effective treatment of operative pain as it blunts autonomic, somatic and endocrine responses. It has become a common practice to use polypharmacy approach for treatment of intra and postoperative pain, because no drug has yet been identified that specifically inhibits nociception without associated side effects. <sup>(1)</sup> Research continues concerning different techniques and drugs that could provide better surgical anesthesia and postoperative pain relief.

Magnesium is the fourth most plentiful cation in our body. It has antinociceptive effects in animal and human models of pain. <sup>(2)</sup> It has been mentioned in a systematic review that it may be worthwhile to further study the role of supplemental magnesium in providing perioperative analgesia, because this is a relatively harmless molecule, is not expensive and also because the biological basis for its potential antinociceptive effect is promising. <sup>(3)</sup> These effects are primarily based on physiological calcium antagonism, that is voltage-dependent regulation of calcium influx into the cell, and noncompetitive antagonism of N-methyl-D-aspartate (NMDA) receptors. <sup>(4)</sup>

The addition of magnesium to epidural bupivacaine and fentanyl in women undergoing elective caesarean section with combined spinal-epidural anesthesia improved intraoperative conditions and the quality of postoperative analgesia. <sup>(4)</sup> In patients undergoing orthopedic surgery, supplementation of spinal anesthesia with combined intrathecal and epidural magnesium sulfate significantly reduced patients' post-operative analgesic requirements. <sup>(5)</sup>

Dexmedetomidine is a new addition to the class of alpha-2 agonist which has got numerous beneficial effects when used through epidural route. <sup>(6)</sup> It acts on both pre and post synaptic sympathetic nerve terminal and central nervous system thereby decreasing the sympathetic outflow and nor-epinephrine release causing sedative, anti-anxiety, analgesic, sympatholytic and hemodynamic effects. <sup>(7)</sup> Dexmedetomidine does cause a manageable hypotension and bradycardia but the striking feature of this drug is the lack of opioid-related side effects like respiratory depression, pruritis, nausea, and vomiting. <sup>(8)</sup>

Ketamine, a non-competitive NMDA receptor antagonist, inhibits central sensitization, thus potentiating the analgesic effect of epidural bupivacaine. <sup>(9)</sup>

### **Aim of work**

The main goal of our study is to compare the efficacy of magnesium sulphate, dexmedetomidine or ketamine when added to 0.5% bupivacaine in epidural anesthesia for lower urinary tract surgery.

The comparison will include onset time, duration, quality and dermatomal spread of epidural anesthesia

# **ANATOMICAL CONSIDERATIONS IN**

## **NEURAXIAL BLOCKADE**

### **The vertebral column:**

The spine is one of the most important parts of our body. The main functions of the spine are: Protecting the spinal cord, nerve roots and several of the body's internal organs, providing structural support and balance to maintain an upright posture and enabling flexible motion.<sup>(10)</sup>

### **Regions of the spine:**

The spine is divided into 4 main regions; cervical, thoracic, lumbar and sacral. Each region has specific characteristics and functions

#### ***- Cervical spine:***

This region consists of 7 vertebrae. These vertebrae protect the brain stem and the spinal cord, support the skull and allow for a wide range of head movement.<sup>(11)</sup>

#### ***- Thoracic spine:***

Beneath the last cervical vertebra are the 12 vertebrae of the thoracic spine. The first thoracic vertebra (T<sub>1</sub>) is the smallest and the last thoracic vertebra (T<sub>12</sub>) is the largest. The thoracic vertebrae are larger than the cervical bones and have longer spinous processes. In addition, rib attachments add to the thoracic spine's strength. These structures make the thoracic spine more stable than the cervical or lumbar regions. In addition, the rib cage and ligament systems limit the thoracic spine's range of motion and protect many vital organs.<sup>(12)</sup>

**- Lumbar spine:**

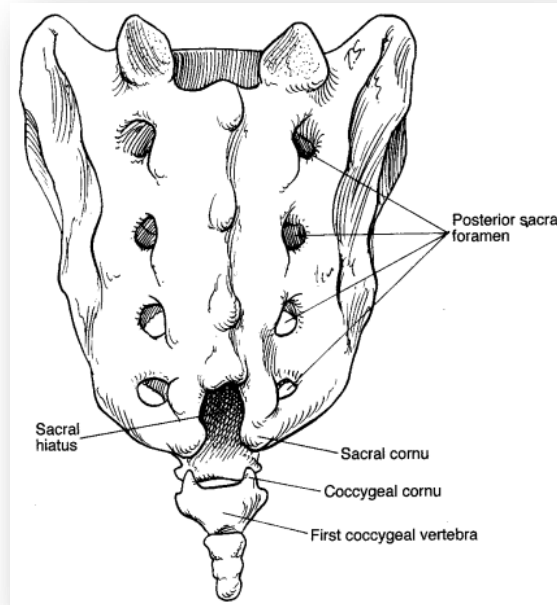
The lumbar spine has 5 vertebrae. The size and shape of each lumbar vertebra is designed to carry most of the body's weight. Each structural element of a lumbar vertebra is bigger, wider and broader than similar components in the cervical and thoracic regions. The lumbar spine has more range of motion than the thoracic spine, but less than the cervical spine.<sup>(13)</sup>

The lumbar facet joints allow for significant flexion and extension movement as they are oriented somewhat parasagittally but limit rotation. Lumbar vertebrae also contain small mammillary and accessory processes on their bodies. These bony protuberances are sites of attachment of deep back muscles .<sup>(14)</sup>

The nearly perpendicular orientation of the spinous process in the lumbar area and the downward angular orientation in the thoracic area define the angle required for placement and advancement of a needle intended to access the vertebral canal. <sup>(15)</sup>

**- Sacral spine:**

The sacrum is located behind the pelvis. 5 bones fused into a triangular shape. The sacrum fits between the two hip bones connecting the spine to the pelvis. The last lumbar vertebra (L5) articulates with the sacrum. Immediately below the sacrum are 5 additional bones, fused together to form the coccyx (Fig. 1). <sup>(10)</sup>



**Fig. (1):** Anatomy of the sacral hiatus and dorsum of the sacrum.<sup>(10)</sup>

### **Spinal curves:**

The normal spine has S like curve when looking at it from the sagittal plane. This allows for an even distribution of weight. The cervical and the lumbar spines curve slightly inward, the thoracic and the sacral spines curve outward. Even though the lower portion of the spine holds most of the body's weight, each segment relies upon the strength of the others to function properly.<sup>(16)</sup>

The thoracic convexity (kyphosis) and the lumbar concavity (lordosis) are of major importance to the distribution of local anesthetic (LA) solution in the subarachnoid space (Fig. 2).<sup>(15)</sup>