**Original Article** 

# Article Postoperative Complications in Lichtenstein Repair Under Spinal

Postoperative Complications in Lichtenstein Repair

## Anesthesia

**Muhammad Fahad** 

## **ABSTRACT**

Objective: To describe postoperative complications in Lichtenstein repair under spinal anesthesia.

**Study Design:** Descriptive, prospective, cohort study

**Place and Duration of Study:** This study was conducted at the Abbasi Shaheed Hospital, Karachi for six months duration from 07.02,2023 till 06.08.2023.

**Methods:** It is a descriptive, prospective<sup>1</sup>, cohort study with both qualitative and quantitative aspects.my research is focus on postoperative complications occurs after Lichtenstein repair. Sample size was derived from formula<sup>2</sup> and through internet based calculator.net which is 194.The no of participants selected through consecutive method of sampling and follow ups are done postoperatively in wards and on out-patient department bases. The postoperative complications were taken as variables and analysed through SPSS. The patients are between 21 and 85 years old. We categorized the individuals into two distinct groups F1 and F2. In the F1 group, we used the European hernia society classification (EHS), and in the F2 group, we used ultrasound and the EHS classification to determine accurate defect size. Lichtenstein repair is the procedure of choice. Postoperative complications were evaluated.

**Results:** According to our results, we found that out of the 97/194 patients in the F1 group, 51 have lateral inguinal hernia and 41 have medial inguinal hernia. In the F2 group, 43 patients have lateral inguinal hernia, and 49 patients have medial inguinal hernia. Postoperative complications. In our study, many patients who developed complications had Lichtenstein repair done by postgraduate surgeons under supervision and surgical registrars. Experienced surgeons like Professor and Associate Professor have performed Lichtenstein repairs on patients with negligible or fewer complications.

**Conclusion:** Accurate estimation of defect size, adequate spinal anesthesia with experienced anesthetists, proper fixation and overlapping of mesh, proper suture material, and instruments with proper lightning and sterilization techniques—surgical experience noted >300 lichtenstein repairs can reduce postoperative complications in Lichtenstein repair.

Key Words: Postoperative Complications, Lichtenstein Repair, Spinal Anesthesia

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## INTRODUCTION

Across the world 20 million patients did Lichtenstein repair yearly. It is one of the best and frequent surgical procedure offered in the world. The rate of occurrence of inguinal hernia is 27-43% for male and 3-6% for female<sup>3,10</sup>. Lichtenstein repair is a type of open inguinal hernia repair with tension free mesh repair technique.

The tension free concept got its description with Irving Lichtenstein (1920-2000) belong to Los Angeles in the

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Accepted: January, 2024 Printed: May, 2024 second edition of his famous hernia article. He formulated a tension free repair by implanting prosthetic material to fix the gap between the muscular and ligament tissues.

His repair evenly used R/Marlex as mesh in a classic anterior inguinal approach. He made the intervention to an state of art procedure. The efforts of Lichtenstein ends with flying colours that it is still one of the best evidence based mesh repair technique in the world. Prolene Hernia System is an adaptation of the Lichtenstein technique in which implantation of a double sided prolene mesh is done before and after the muscle by an open incision. In 1987 Lichtenstein bring out his data registry configuration. It had his own observations with over 6000 demonstrations and he also incorporated his classification system.

Despite all advances, postoperative complications in Lichtenstein repair is still the matter of debate. A clinical study showed the rate of developing intraoperative complications is 6% and immediate postoperative complications was seen in 12% Patients Postoperative surgical complications are classified

usually according to Clavien Dindo Classification (CDC)<sup>7,8</sup>. This classification is constituted to categorize the severity of a surgical outcome. It is based on the line of management needed to correct the complications<sup>9</sup>. The scale divided into several grades

**Grade 1:** Any deviation from the normal postoperative course without the need for pharmacological treatment or surgical, endoscopic and radiological interventions.

**Grade 2:** Requiring pharmacological treatment with drugs other than such allowed for Grade I complications. Blood transfusions and total parenteral nutrition are also included.

**Grade 3:** Requiring surgical, endoscopic or radiological intervention.

**Grade 3 a:** Intervention not under general anesthesia.

**Grade 3 b:** Intervention under general anesthesia.

**Grade 4:** Life-threatening (complication including those affecting the brain) requiring intensive care management.

Grade 4 a: Single organ dysfunction (including dialysis)

Grade 4 b: Multi organ dysfunction.

**Grade 5:** Death of patient

In Lichtenstein repair usually grade 1,2 and 3 complications are reported in our study. Grade 4 and 5 complications are not usually encountered.

The postoperative complications in Lichtenstein repair is usually divided into early and late postoperative complications. The early post operative complications are postoperative pain, urinary retention, postoperative bleeding, hematoma, seroma and wound infection late postoperative complications are recurrence ,chronic postoperative pain, postoperative neuralgias and testicular atrophy.

#### **METHODS**

In surgical ward of our hospital, I have conducted a prospective cohort study which is both qualitative and quantitative. The objective of research study was to see the impact of accurate defect size estimation and overlapping of mesh on postoperative complications in Lichtenstein repair. I have selected 194 patients of direct and indirect inguinal hernia through consecutive sampling and divided into two groups F1 and F2 .In first group F1, I have estimated accurate defect size through European hernia society (EHS) and in group F2 ultrasound and EHS classification is used for determination of defect size in inguinal hernia .Postoperative complications in both group have been evaluated upto three months in both groups.

## RESULTS

According to our results, we found that out of the 97/194 patients in the F1 group, 51 have lateral inguinal hernia and 41 have medial inguinal hernia. In the F2 group, 43 patients have lateral inguinal hernia, and 49 patients have medial inguinal hernia.

Table No.1: Postoperative complications.

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Postoperative	F1	F2	Analysis	Treatment		
complications						
1)hematoma	47	O7	P:0. 001, OR:13.67,	Conservative, guided drainage,		
			95%CI:5.753,32.51, Kappa:0.443	and antibiotics		
2)wound infection	63	07	P:0.001, OR:26.11,	Conservative, antibiotics		
			95%CI:10.85,62.82,kappa:0.598	anti-inflammatory		
3)Postop pain	64	07	P:0.001, OR:23.68,	Conservative		
			95%CI:10.22,54.86	analgesics		
			kappa:0.598	anti-inflammatory		
4)postop neuralgia	60	07	P:0.001, OR:43.83,	Analgesic		
			95%CI:17.74,108.24,kappa:0.701	gabapentin		
5)recurrence	01	00	-	Observation		
				surgical management		
6)Testicular atrophy	01	00	-	Observation		

Postoperative Complications In:

## 1) Haematoma:

group * haematoma Cross tabulation Count					
Count					
		haematoma			
		yes	no	Total	
group	f1=no	50	47	97	
	f2=yes	7	90	97	
Total		57	137	194	

**F1 Group:** 50 patients developed haematoma in the early postoperative period resolved conservatively.

**F2 group:** 07 patients developed haematoma resolved conservatively.

Statistical analysis: P-value: 0.001

Odd's Ratio: 13.67

95% Confidence interval: 5.75,32.51,kappa:0.443

#### 2) Wound infection:

Group * wound infection Cross tabulation Count					
		Wound Infection			
		yes	no	Total	
group	f1=no	65	32	97	
	f2=yes	7	90	97	
Total		72	122	194	

**F1 group:** 65 patients developed wound infection in F1 group which are treated conservatively by daily dressing, antibiotics and anti-inflammatory medication. The most common organisms found are staphylococcus aureus, Ecoli, and klebsiella.

**F2 group:** O7 patients developed wound infection treated. Mostly are superficial surgical site infection treated conservatively with daily dressing and antibiotics.

Statistical analysis: P value: 0.001 Odds ratio: 26.11

95% confidence interval: 10.85,62.82, kappa:0.598

#### 3) Postoperative Pain:

group * postoperativepain Crosstabulation						
Count						
		postoperativepain				
		pain	no pain	Total		
group	f1=no	66	31	97		
	f2=yes	8	89	97		
Total		74	120	194		

**F1 group:** 66 patients have mild to moderate pain in first 10 days with paper based VAS (visual analog score) for pain is 4-8. 11,13

No patient developed pain till 03 months.

All the patients treated conservatively.

**F2 group:** 08 patients had mild to moderate pain with paper based VAS (visual analog score) was 4-8. 11,4

## **Statistical Analysis:**

P value:0.001

Odds Ratio:23.68

95% Confidence interval:10.22,54.86,kappa:0.5998.

## 4) Postoperative Neuralgia:

**F1 group:** 64 patient had complains of postoperative neuralgia (pricking, numbness, burning at surgical site and in groins, spinal headache. Treated conservatively with anti inflammatory, analgesics and gabapentin.

**F2 group:** 08 patients developed postoperative neuralgias for 10 days upto 01 month treated conservatively with anti-inflammatory ,spinal headache with caffeine, cola drinks and gabapentin.

#### **Statistical Analysis:**

P-value:0.001.

Odds ratio:21.57.

95% Confidence interval: 9.34,49.80. kappa: 0.577.

**5)Testicular atrophy:** Only 01 patient reported in F1 group

**6) Recurrence:** Only 01 patient reported in F1 group.

In our study many patients who developed complications have Lichtenstein repair done by postgraduate surgeons under supervision and by surgical registrars. The lichtenstein repair on patients carried out by experienced surgeon have developed negligible or less complications. The level of experience noted should be >300 lichtenstein repair. Combination of ultrasound and EHS classification is found to be sensitive in reducing post-operative complications by accurate estimation of defect size and with the help of this junior as well as senior surgeons can plan to overcome the incident alomas and intra and postoperative complications<sup>14</sup>.

## **DISCUSSION**

Inguinal hernia has been the disease ever since the mankind existed 1.04 decades before Lichtenstein developed a state of art tension free mesh repair known as Lichtenstein Repair 1. In the year 2004, European hernia society annual meeting held in Capri, Italy in which the standard inguinal hernia classification system was orchesterated. In this simple and comprehensive classification of hernia 15 in which direct, indirect and femoral hernia were marked and defect size will be evaluated by taking index fingerbreadth as a criteria of measurement. Ultrasound helps in diagnosing occult inguinal hernia 16. Significance of clinically hard inguinal hernia with help of ultrasound is confirmed by European hernia society.

A European study led by Mathews proposed that in patients with normal or doubtful clinical examination, the preoperative ultrasound can be considered diagnostic for evaluation. It is now affirmed that ultrasound has high accuracy in the diagnosis of inguinal hernia including differentiating the type of hernia in doubtful cases. Post-operative complications in tension free mesh repair is still the problem in Lichtenstein repair. In one study 12 % cases of different post-operative complications were identified in Lichtenstein repair.

The most common post-operative complication world wide in open inguinal hernia repair is recurrence but due to major advances and expertise it is declining. Other post-operative complications such as wound infection, urinary retention, haematoma <sup>17</sup>, seroma, postoperative pain, postoperative neuralgia, testicular atrophy is also present. In my study, I found 54(27%) patient developed haematoma <sup>17</sup>, 70(36%) patient developed wound infection <sup>5</sup>, postoperative pain <sup>5</sup> is present in 70(36%) of cases mostly pain of short duration, no case of chronic postoperative pain <sup>19</sup> was reported in the duration of study all cases were managed conservatively,

In this study, 67(34%) cases of postoperative neuralgia were reported included patients having mild pricking, burning sensation of short duration in the area of nerve distribution all treated conservatively. Common nerves

encountered in Lichtenstein repair are three Ilioinguinal nerve, genital branch of genitofemoral nerve and iliohypogastric nerves. Cases of spinal headache due to inappropriate spinal anesthesia were also reported and increased the length of stay. Approximately 38(20%) cases of spinal headache were reported in our study. All cases were treated conservatively. Cases of urinary retention were 6(3%) also found in the duration of study.01 patient of testicular atrophy and 01 patient of recurrence<sup>20</sup> were reported in duration of our study. In this study I found Lichtenstein repair done by expert surgical hands causing less complications. The level of experience should be >300 lichenstein repairs. Many risk factors were associated with development of postoperative complications in this study including age<sup>21</sup>, intercurrent illneses, lack of expertise of surgeons and anesthetist, inappropriate suture material, fixation of mesh over the edge which should be 0.5 cm away from edge, improper closure technique by junior surgeons. The author concluded that use of spinal anaesthesia in elderly patient is not supported by existing evidence.

## **CONCLUSION**

Accurate estimation of defect size, adequate spinal anesthesia with experienced anesthetist, proper fixation and overlapping of mesh, Availability of proper suture material, and instruments with proper lightning and sterlisation techniques, surgical expertise can reduce postoperative complications in Lichtenstein repair.

#### **Author's Contribution:**

Concept & Design of Study: Muhammad Fahad Drafting: Muhammad Fahad Data Analysis: Muhammad Fahad Revisiting Critically: Muhammad Fahad Final Approval of version: Muhammad Fahad

**Conflict of Interest:** The study has no conflict of interest to declare by any author.

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