



EFFICACY OF SINGLE DOSE VERSUS MULTIPLE DOSE ANTIBIOTIC IN ELECTIVE INGUINAL HERNIA SURGERIES

Dr. Girish B. K¹, Dr. Harish. C², Dr. Hemand N. M³, Dr. Shibin Sankar^{4*}

^{1,3} Junior resident 3rd year, Rajarajeshwari Medical College and Hospital, Bengaluru-560074, Karnataka, India.

² Senior Resident General Surgery, Rajarajeshwari Medical College and Hospital, Bengaluru-560074, Karnataka, India. Email - dr.girishkrishnamoorthi@gmail.com

^{4*} Junior resident 3rd year, Rajarajeshwari Medical College and Hospital, Bengaluru-560074, Karnataka, India.

***Corresponding Author:** Dr. Shibin Sankar

*Junior resident 3rd year, Rajarajeshwari Medical College and Hospital, Bengaluru-560074, Karnataka, India.

INTRODUCTION

A hernia is an abnormal protrusion of the whole or part of a viscus through normal or abnormal opening in the walls of its containing cavity.¹ Inguinal hernias are among the most common problems seen in surgical clinics. Inguinal hernias comprise 75% of all abdominal hernias. The lifetime risk of developing an inguinal hernia is 27% in males. The lifetime risk of developing an inguinal hernia is 3% in females². They can be congenital or acquired, complete or partial, external or internal, reducible or irreducible, direct or indirect, and primary or recurrent³. Patients typically present with a bulge in the groin that is associated with pain in two-thirds of cases.⁴ Painful hernias are most frequently described as a dull aching, heavy, dragging, or burning sensation. Maneuvers that increase intra-abdominal pressure, such as straining, lifting, or coughing, may exacerbate pain or hernia size by causing intra-abdominal contents to be pushed through the fascial defect⁵. Some patients may complain of worsening symptoms at the end of the day or after increased activity. Minor symptomatic cases may be temporarily improved by lying down or reducing the hernia manually. Severe or unbearable pain, that is, sudden onset, suggests possible complicated hernia and should be treated as an emergency.

Inguinal hernias are primarily diagnosed by history and physical examination with secondary imaging rarely needed⁶. Traditionally almost all inguinal hernias are referred for surgical treatment following diagnosis. Progression of a hernia by time is natural and most surgeons prefer repairing all inguinal hernias as soon as possible. Inguinal hernia is a benign disease and its repair results in only rare and minor complications in elective setting. Nevertheless complications developed after emergency repairs may be more dramatic and frequent, even mortality may be recorded^{7,8}.

Table 1: A classification of current repair techniques for inguinal hernias-⁹

A. Tension-free prosthetic repairs	
1.	Anterior repairs
a.	Lichtenstein repair and its modifications
b.	Plug repairs
c.	Patch and plug repairs
d.	Double-layer devices
2.	Posterior (preperitoneal) repairs
a.	Open techniques via inguinal incision
b.	Stoppa repair
c.	Laparoscopic/endoscopic repairs
i.	Transabdominal preperitoneal
ii.	Total extraperitoneal
B. Tissue-Suture repairs	
1.	Bassini-Shouldice technique and its modifications
2.	Marcy repair

The open repair with mesh (Lichtenstein tension-free) technique is the current gold standard of care for most patients with an inguinal hernia.¹⁰ The use of prosthetic mesh is recommended because of its association with a 50%-75% lower risk of hernia recurrence, lower risk of chronic pain post-operatively, and an earlier return to work compared with a sutured repair.¹¹

Other surgical repairs like non-mesh open repairs, the Shouldice technique is recommended due to its lower risk of recurrence compared with other pure tissue repairs (e.g., McVay or Bassini techniques).¹¹ The recurrence rate with the Shouldice techniques is higher than that with the mesh techniques [odds ratio (OR) 3.80, 95% confidence interval (CI) 1.99–7.26] but lower than other pure tissue repairs (OR 0.62, 95% CI 0.45–0.85).¹²

Hernia repair is typically classified as a "clean" procedure, meaning the surgical site has minimal contamination risk¹³.

Despite this classification, surgical site infections (SSIs) remain a potential complication, leading to increased morbidity, healthcare costs, patient discomfort and Mental stress.

Antibiotic prophylaxis, the administration of antibiotics before surgery to prevent infection, plays a crucial role in minimizing SSIs after elective inguinal hernia repair¹⁴.

Different techniques of antibiotic prophylaxis, the dose and nature of drug use, timings of the usage were studied by various cross sectional studies and based on which standard protocol for antibiotic prophylaxis for each procedure were made¹⁵.

The approach of providing antibiotic prophylaxis as a single dose, given just before the surgical incision is made, has become increasingly popular and widely adopted.

This single-dose prophylactic antibiotic regimen is favored due to its simplicity, as it involves only a one-time administration of the antibiotic. The convenience offered by the single-dose regimen, in contrast to multi-dose or prolonged antibiotic

administration, contributes to its growing popularity and acceptance¹⁶. In public healthcare facilities grappling with overcrowded conditions and suboptimal hygiene standards, a widespread practice involves administering antibiotics for an extended duration of 7 to 10 days, even for clean and clean-contaminated surgical procedures.

This practice stems from concerns over the potential development of surgical site infections. However, such prolonged multi-dose antibiotic regimens not only impose a substantial financial burden on the hospital but also contribute to the alarming rise of antimicrobial resistance.

Our study aims to evaluate the effectiveness of a single-dose antibiotic prophylaxis regimen, administered 30 minutes prior to hernia surgery, utilizing a standard, carefully selected antibiotic agent. This approach will be compared to the conventional practice of administering the same antibiotic in multiple doses until the patient is discharged from the hospital.

The primary objective is to determine whether a single-dose prophylactic regimen can achieve comparable efficacy to the multi-dose approach, thereby offering the potential to reduce costs while concurrently mitigating the risk of fostering antimicrobial resistance.

AIMS & OBJECTIVES:

This study aims to compare the efficacy of single dose versus multiple dose antibiotics in elective Inguinal hernia surgery. More precisely,

1. To assess the efficacy of single dose versus multiple dose antibiotics in preventing surgical site infection in elective Inguinal hernia surgery.
2. To assess safety and cost efficacy of using single dose versus multiple dose antibiotics prophylaxis.

MATERIALS & METHODS:

A. Source of Data :This study was conducted on the patients admitted in the department of general surgery for inguinal hernia at Rajarajeswari Medical College and Hospital

B. Method of Data Collection

Study design: Randomized case-control prospective study Study period: August 2022 to February 2024

Study Centre : Rajarajeswari Medical College and Hospital

SampleSize :60 (30 for single dose and 30 for multiple dose)

Study group/ Group A :Patients undergoing Elective Inguinal hernia repair will be given a single dose of 1.5gm Cefaperazone + sulbactam intravenously half an hour before operation.

Control group /Group B :Patients undergoing Elective Inguinal hernia repair will be given multiple doses of 1.5gm Cefaperazone + sulbactam intravenously for 5 days post operatively.

Inclusion Criteria:

1. Patients with the age group between 18-60 years of either sex.
 2. Those willing to give informed consent (annexure 1) and posted for following surgeries are included in the study.
 3. Reducible Inguinal hernia of all types undergoing elective inguinal hernia repair.
- Exclusion Criteria:

1. Age group less than 18 years and more than 60 years.

2. Patients posted for emergency inguinal surgeries.
3. History of hypersensitivity to cephalosporin group of antibiotics.
4. Patient on steroid medications or those who have immunodeficiency.
5. Patients not willing to give informed consent.

Methodology:

After obtaining approval and clearance from the institutional ethics committee, the patients fulfilling the inclusion criteria will be enrolled for the study after obtaining informed consent. (Annexure – 1) Case record form with follow up chart will be maintained (Annexure – 2). This study will be conducted as a randomized case control prospective study in the Department of Surgery in Rajarajeswari medical college and Hospital, Bangalore. Totally 60 patients fulfilled the inclusion criteria for elective inguinal hernia surgeries were admitted in our hospital. The patients will be randomized into study group A and B. Patients in group A will be given a single dose of 1.5gm Cefaperazone + sulbactam intravenously half an hour before operation and patients in group B will be given multiple doses of antibiotic intravenously for 5 days post operatively. All the surgeries will be carried out in the same theatre environment, and same preoperative safety protocol, and post-operative care will be followed for all patients. Temperature and vitals are monitored periodically, and the charts are maintained. Based on the Southampton scoring system on post-operative day 3rd, 5th, 7th, 14 th and 30th days the wounds will be inspected and the infection grades will be noted. Patients will be followed up with the drugs to be administered and ensured the antibiotics are given at appropriate time as per the protocol.

Assessment tools:

- -Development of infection will be measured based on Southampton grade.
- -Outcomes will be measured in terms of rate of surgical site infection in single dose and multiple dose antibiotic groups.

STATISTICAL ANALYSIS:

The collected data were entered in the Microsoft Excel 2016 and analysed with IBM SPSS Statistics for Windows, Version 29.0.(Armonk, NY: IBM Corp).To describe about the data descriptive statistics

frequency analysis, percentage analysis were used for categorical variables and the mean & S.D were used for continuous variables. To find the significant difference between the bivariate samples in Independent groups the Independent sample t-test was used. To find the significance in qualitative categorical data Chi-Square test was used similarly if the expected cell frequency is less than 5 in 2×2 tables then the Fisher's Exact was used. In all the above statistical tools the probability value .05 is considered as significant level.

RESULTS:

Table: Age distribution

Age distribution		
	Frequency	Percent
Upto 30 yrs	10	16.7
31 - 40 yrs	13	21.7
41 - 50 yrs	17	28.3
51 - 60 yrs	20	33.3
Total	60	100.0

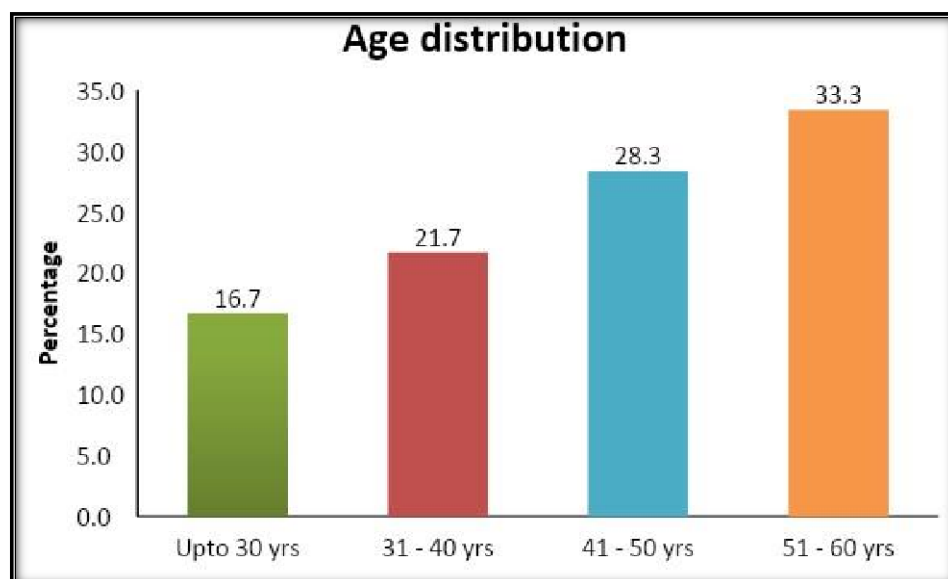


Figure 24: Age distribution between the two groups

The above table shows Age distribution were <30 years is 16.7%, 31-40 years is 21.7%, 41-50 years is 28.3%, 51-60 years is 33.3%.

Table: Gender distribution

Gender distribution		
	Frequency	Percent
Female	14	23.3
Male	46	76.7
Total	60	100.0

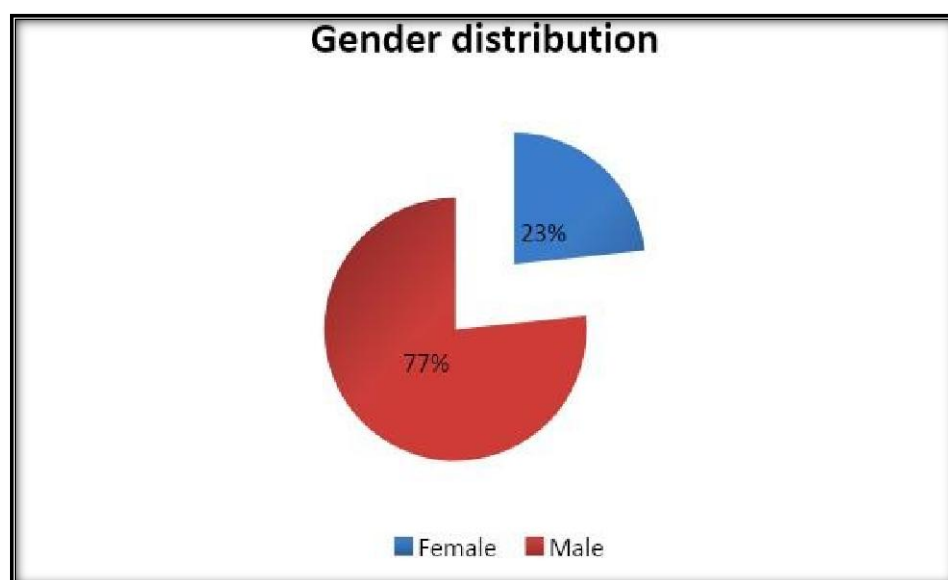


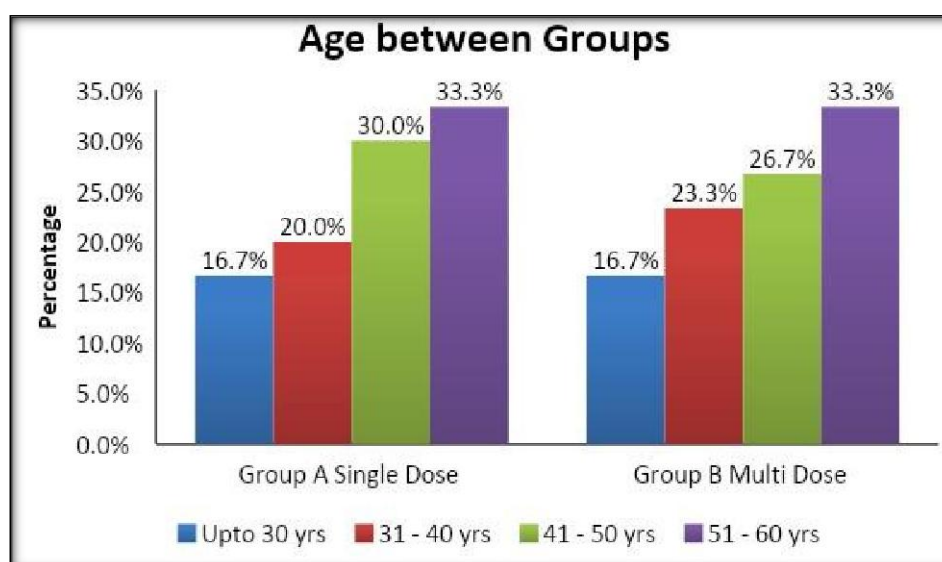
Figure 25: Gender distribution between the two groups

The above table shows Gender distribution were Female is 23.3%, Male is 76.7%.

Table: Comparison of Age between Groups by Pearson's Chi-Square test

			Groups		Total	χ^2 - value	p-value
			Group A Single Dose	Group B Multi Dose			
Age	Upto 30 yrs	Count	5	5	10	0.136	0.987 #
		%	16.7%	16.7%	16.7%		
	31 - 40 yrs	Count	6	7	13		
		%	20.0%	23.3%	21.7%		
	41 - 50 yrs	Count	9	8	17		
		%	30.0%	26.7%	28.3%		
Total		Count	30	30	60		
		%	100.0%	100.0%	100.0%		

No Statistical Significance at $p > 0.05$ level

**Figure: Comparison of Age between Groups by Pearson's Chi- Square test**

The above table shows comparison of Age between Groups by Pearson's Chi-Square test were $\chi^2 = 0.136$, $p = 0.987 > 0.05$ which shows no statistical significance association between Age and Groups. The mean \pm standard deviation of the age in GROUP A SINGLE DOSE were 43.5 ± 11.6 years and in GROUP B MULTI DOSE were 42 ± 11.4 years

Table: Comparison of Gender between Groups by Pearson's Chi- Square test

			Groups		Total	χ^2 - value	p-value
			Group A Single Dose	Group B Multi Dose			
Gender	Female	Count	7	7	14	0.000	1.000 #
		%	23.3%	23.3%	23.3%		
	Male	Count	23	23	46		
		%	76.7%	76.7%	76.7%		
Total		Count	30	30	60		
		%	100.0	100.0	100.0		

No Statistical Significance at $p > 0.05$ level

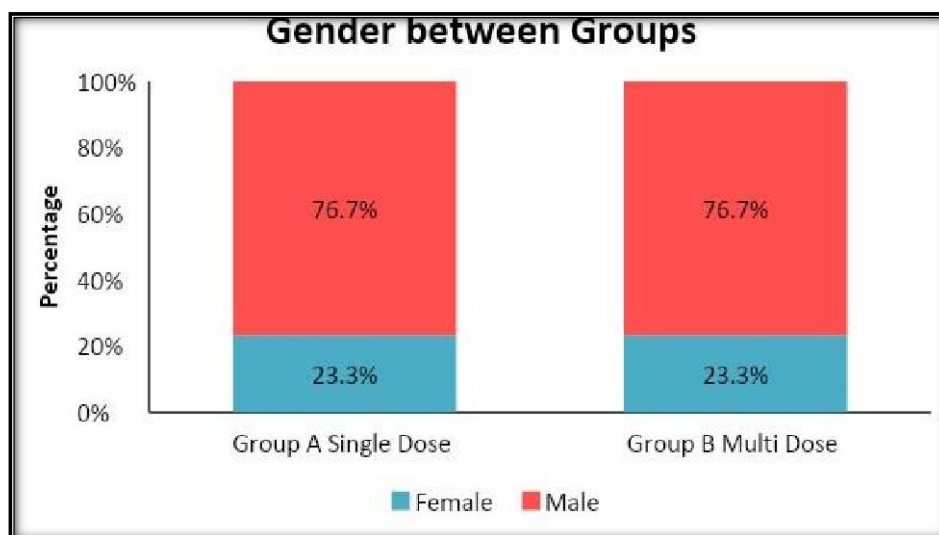


Figure: Comparison of Gender between Groups by Pearson's Chi- Square test

The above table shows comparison of Gender between Groups by Pearson's Chi-

Square test were $\chi^2=0.000$, $p=1.000>0.05$ which shows no statistical significance association between Gender and Groups.

Table : Comparison of Type of Hernia between Groups by Pearson's Chi-Square test

			Groups		Total	χ^2 - value	p-value
			Group A Single Dose	Group B Multi Dose			
Type of Hernia	Bilateral	Count	0	5	5	5.455	0.052 #
		%	0.0%	16.7%	8.3%		
	Unilateral	Count	30	25	55		
		%	100.0 %	83.3%	91.7%		
Total		Count	30	30	60		
		%	100.0 %	100.0 %	100.0 %		
		# No Statistical Significance at p > 0.05 level					

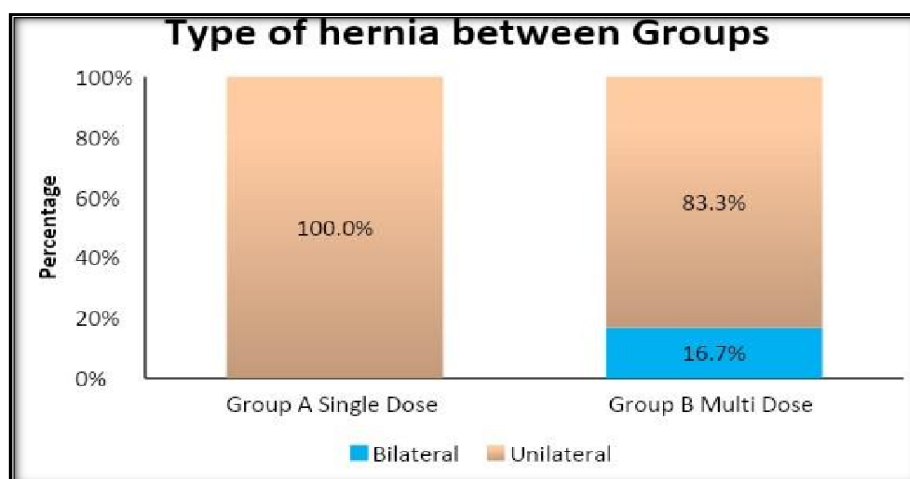


Figure: Comparison of Type of Hernia between Groups by Pearson's Chi- Square test

The above table shows comparison of Type of Hernia between Groups by Pearson's Chi-Square test were $\chi^2=5.455$, $p=0.052>0.05$ which shows no statistical significance association between Type of Hernia and Groups.

Table: Comparison of Side of Hernia between Groups by Pearson's Chi- Square test

			Groups		Total	χ^2 - value	p-value
			Group A Single Dose	Group B Multi Dose			
Side of Hernia	Left	Count	15	15	30	0.000	1.000 #
		%	50.0%	50.0%	50.0%		
	Right	Count	15	15	30		
		%	50.0%	50.0%	50.0%		
Total		Count	30	30	60		
		%	100.0	100.0	100.0		
		%	%	%			
# No Statistical Significance at p > 0.05 level							

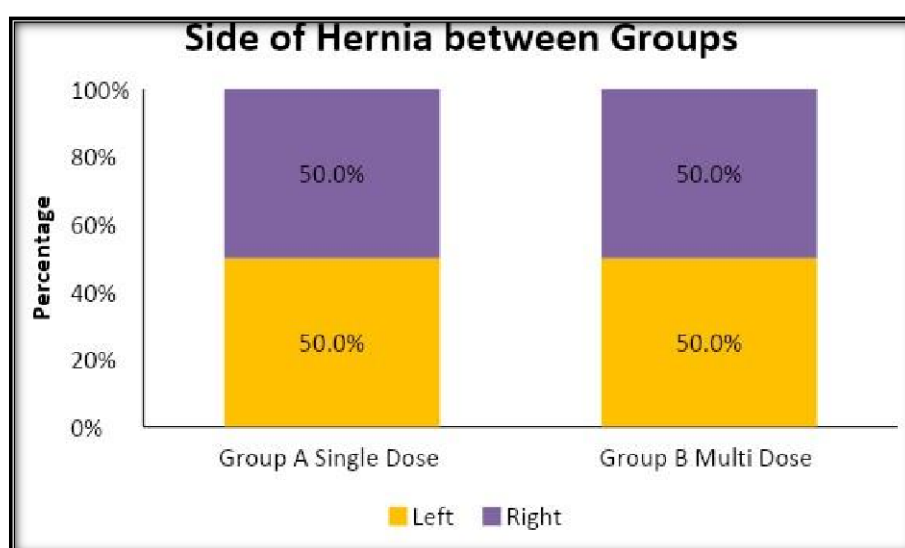


Figure: Comparison of Side of Hernia between Groups by Pearson's Chi- Square test

The above table shows comparison of Side of Hernia between Groups by Pearson's Chi-Square test were $\chi^2=0.000$, $p=1.000>0.05$ which shows no statistical significance association between Side of Hernia and Groups.

Table: Comparison of SSI between Groups by Pearson's Chi-Square test.

			Groups		Total	χ^2 - value	p-value
			Group A Single Dose	Group B Multi Dose			
	Day 3	Count	2	1	3		
		%	66.7%	100.0%	75.0%		
	Day 5	Count	1	0	1		
		%	33.3%	0.0%	25.0%		
	Day 7	Count	0	0	0		
		%	0.0%	0.0%	0.0%		
		Count					
		%					

SSI		Count	0	0	0	0.444	1.000 #
	Day 14	%	0.0%	0.0%	0.0%		
	Day 30	Count	0	0	0		
		%	0.0%	0.0%	0.0%		
Total		Count	3	1	4		
		%	100.0%	100.0%	100.0%		
# No Statistical Significance at p > 0.05 level							

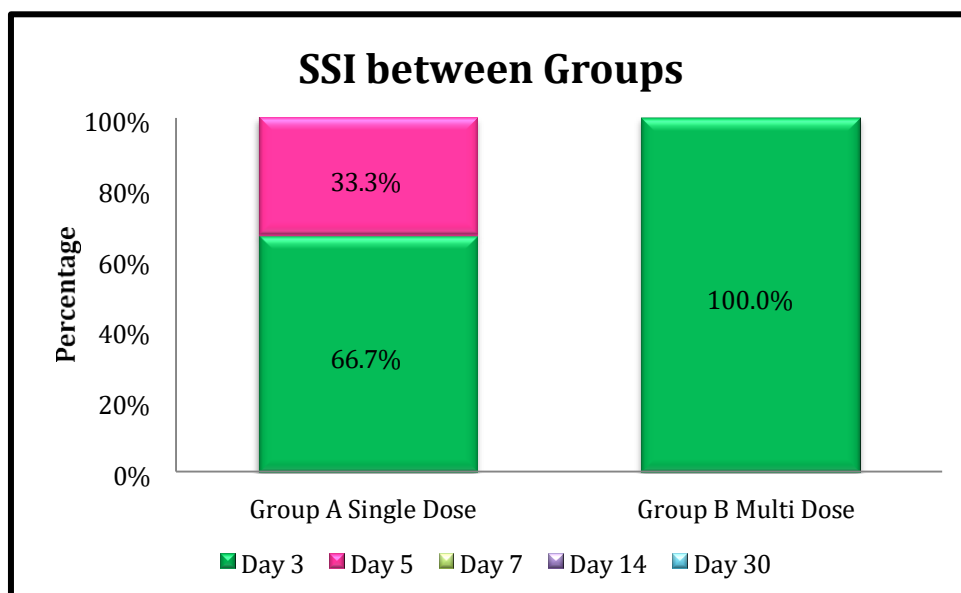


Figure: Comparison of SSI between Groups by Pearson's Chi-Square test

The above table shows comparison of SSI between Groups by Pearson's Chi-Square test were

$\chi^2 = 0.313$, $p = 1.000 > 0.05$ which shows no statistical significance association between SSI and Groups.

Table: Comparison of Deep SSI between Groups by Fisher's exact test

			Groups		Total	χ^2 - value	p-value	
			Group A Single Dose	Group B Multi Dose				
Deep SSI	No	Count	29	30	59	1.017	1.000 #	
		%	96.7%	100.0 %	98.3%			
	Yes	Count	1	0	1			
		%	3.3%	0.0%	1.7%			
	Total		Count	30	30			60
			%	100.0 %	100.0 %			100.0 %
# No Statistical Significance at p > 0.05 level								

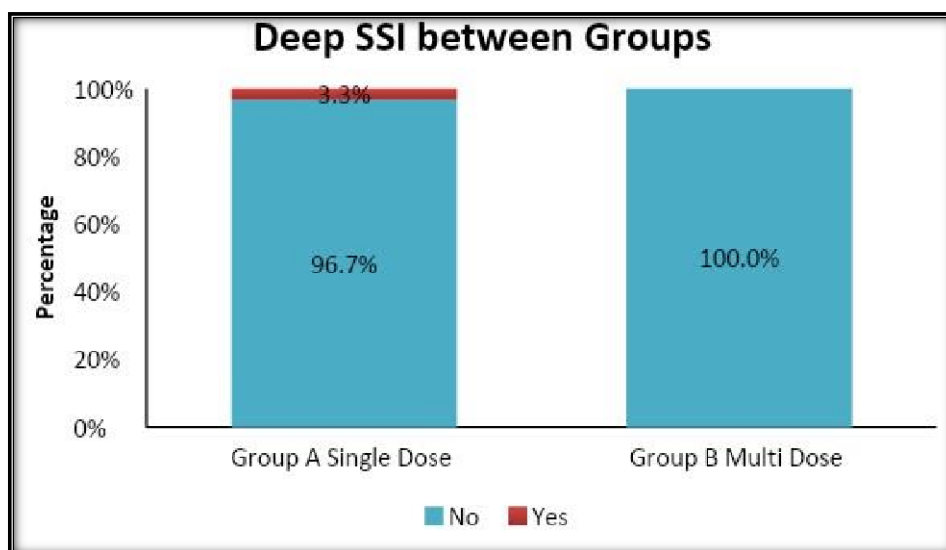


Figure: Comparison of Deep SSI between Groups by Fisher's exact test

The above table shows comparison of Deep SSI between Groups by Fisher's exact test were $\chi^2 = 0.017$, $p = 1.000 > 0.05$ which shows no statistical significance association between Deep SSI and Groups.

Table: Comparison of Superficial SSI between Groups by Fisher's exact test

			Groups		Total	χ^2 - value	p-value
			Group A Single Dose	Group B Multi Dose			
Superficial SSI	No	Count	29	29	58	0.000	1.000 #
		%	96.7%	96.7%	96.7%		
	Yes	Count	1	1	2		
		%	3.3%	3.3%	3.3%		
Total		Count	30	30	60		
		%	100.0%	100.0%	100.0%		

No Statistical Significance at p > 0.05 level

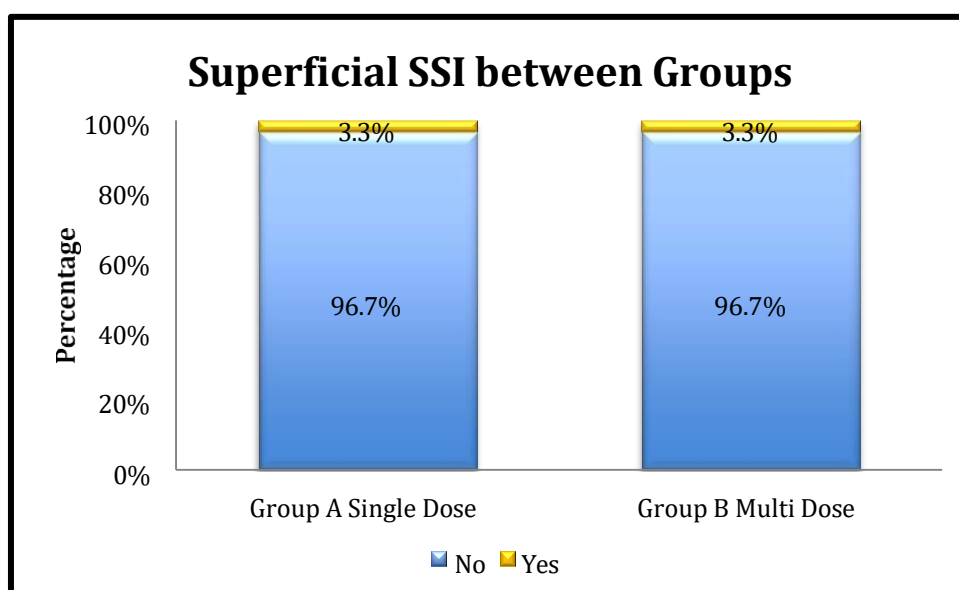


Figure: Comparison of Superficial SSI between Groups by Fisher's exact test

The above table shows comparison of Superficial SSI between Groups by Fisher's exact test were

$2=0.741$, $p=0.671>0.05$ which shows no statistical significance association between Superficial SSI and Groups.

Table: Comparison of Conversion form SD to MD between Groups by Fisher's exact test

			Groups		Total	χ^2 - value	p-value
			Group A Single Dose	Group B Multi Dose			
Conversion form SD to MD	No	Count	29	30	59	1.017	1.000 #
		%	96.7%	100.0%	98.3%		
	Yes	Count	1	0	1		
		%	3.3%	0.0%	1.7%		
Tot 1		Count	30	30	60		
		%	100.0%	100.0%	100.0%		

No Statistical Significance at $p > 0.05$ level

No Statistical Significance at $p > 0.05$ level

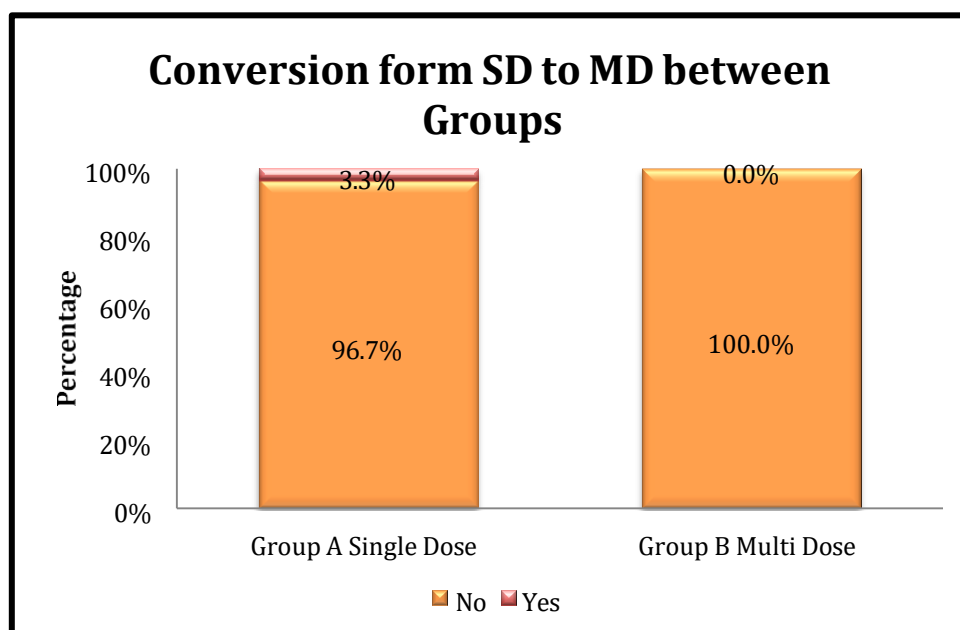


Figure: Comparison of Conversion form SD to MD between Groups by Fisher's exact test

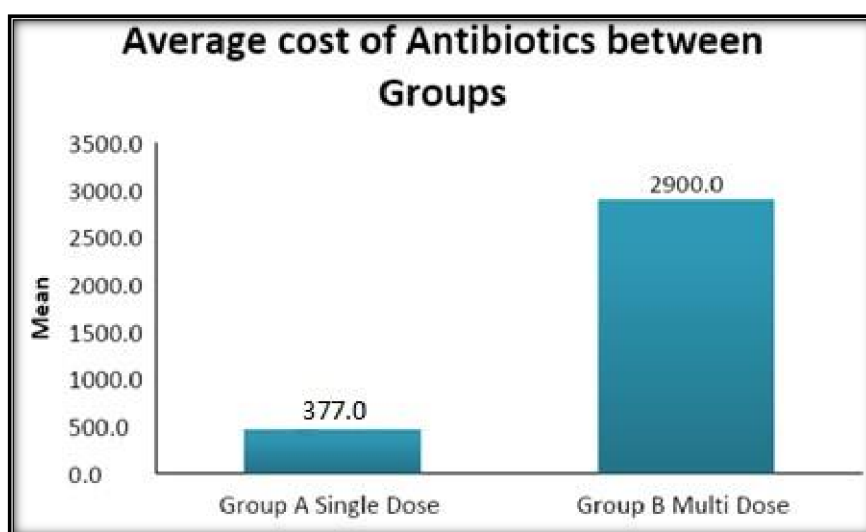
The above table shows comparison of Conversion form SD to MD between Groups by Fisher's exact test were $2=2.069$, $p=0.492>0.05$ which shows no statistical significance association between Conversion form SD to MD and Groups.

□

Table: Comparison of Average cost of Antibiotics between Groups by Independent sample t-test

Variable	Groups	N	Mean	SD	t-value	p-value
Average cost of Antibiotics	Group A Single Dose	30	377.0	662.2	20.149	0.0005**
	Group B Multi Dose	30	2900.0	0.0		

** Highly Statistical Significance at $p < 0.01$ level

**Figure 34: Comparison of Average cost of Antibiotics between Groups by Independent sample t-test**

The above table shows comparison of Average cost of Antibiotics between Groups by Independent sample t-test were $t\text{-value}=20.149$, $p\text{-value}=0.0005 < 0.01$ which shows highly statistical significance difference at $p < 0.01$ level.

Table: Group Statistics of Average cost of Antibiotics between Groups in Conversion from SD to MD

Group Statistics					
	Groups	Conversion from SD to MD	N	Mean	SD
Average cost of Antibiotics	Group A Single Dose	Yes	1	2900.0	0.0
Average cost of Antibiotics	Group A Single Dose	Yes	1	2900.0	0.0
		No	29	290.0	0.0
	Group B Single Dose	Yes	0	0.0	0.0
		No	30	2900.0	0.0

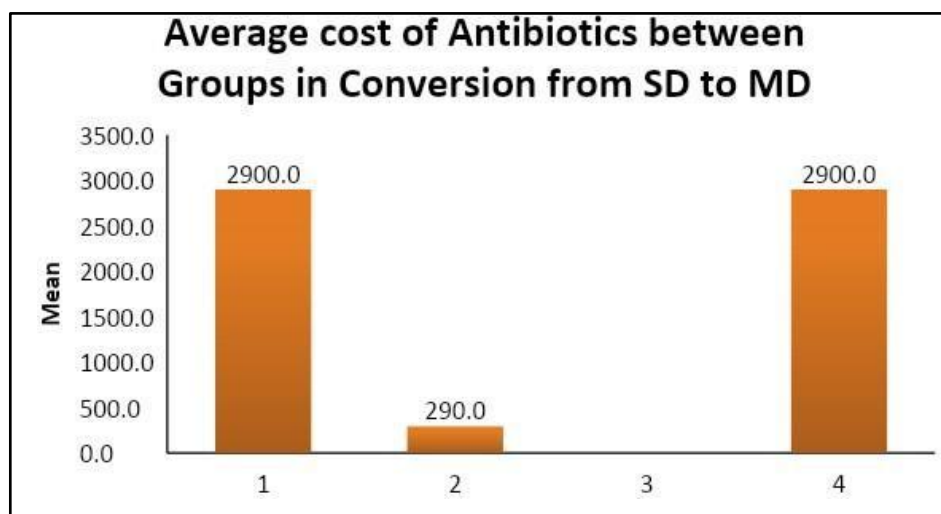


Figure: Group Statistics of Average cost of Antibiotics between Groups in Conversion from SD to MD

The above table shows the mean \pm standard deviation of the Antibiotics between Groups in Conversion from SD to MD.

Summary

- The Age distribution were <30 years is 16.7%, 31-40 years is 21.7%, 41-50 years is 28.3%, 51-60 years is 33.3%.
- The Gender distribution were Female is 23.3%, Male is 76.7%.
- The Age between Groups by Pearson's Chi-Square test were $\chi^2=0.136$, $p=0.987>0.05$ which shows no statistical significance association between Age and Groups. The mean \pm standard deviation of the age in Group A Single Dose were 43.5 ± 11.6 years and in Group B Multi Dose were 42 ± 11.4 years
- The Gender between Groups by Pearson's Chi-Square test were $\chi^2=0.000$, $p=1.000>0.05$ which shows no statistical significance association between Gender and Groups.
- The comparison of Type of Hernia between Groups by Pearson's Chi-Square test were

$\chi^2=5.455$, $p=0.052>0.05$ which shows no statistical significance association between Type of Hernia and Groups.

- The Side of Hernia between Groups by Pearson's Chi-Square test were $\chi^2=0.000$, $p=1.000>0.05$ which shows no statistical significance association between Side of Hernia and Groups.
- The comparison of SSI between Groups by Pearson's Chi-Square test were $\chi^2=0.313$, $p=1.000>0.05$ which shows no statistical significance association between SSI and Groups.
- The Deep SSI between Groups by Fisher's exact test were $\chi^2=1.017$, $p=1.000>0.05$ which shows no statistical significance association between Deep SSI and Groups.
- The Superficial SSI between Groups by Fisher's exact test were $\chi^2=0.741$, $p=0.671>0.05$ which shows no statistical significance association between Superficial SSI and Groups.
- The Conversion form SD to MD between Groups by Fisher's exact test were $\chi^2=2.069$, $p=0.492>0.05$ which shows no statistical significance association between Conversion form SD to MD and Groups.
- The Average cost of Antibiotics between Groups by Independent sample t-test were t-value=20.149, p-value=0.0005<0.01 which shows highly statistical significance difference at $p < 0.01$ level.
- The mean \pm standard deviation of the Antibiotics between Groups in Conversion from SD to MD.

DISCUSSION:

Surgeries performed on elective basis are generally clean ones.

The factors playing a major role in averting surgical site infection other than risk factors associated with patients, are the atmosphere and sterility of operating room, the sterility of instruments, surgeons due efforts to maintain asepsis during the surgery.

The operating surgeon must not have the freedom of prescribing antibiotics due to faulty techniques as it can never be a substitute for a clean aseptic environment. In clean elective surgeries, the source of infection in case of wound sepsis is often from an exogenous source like the nostril or oral cavity of surgeons or skin of patient.

In our study patient risk factors like diabetes mellitus, hypertension, immunocompromised state, hypersensitivity to any drugs and any other comorbidities that may hamper the results have been strictly excluded.

The literature is of the opinion that the rate of infection in clean surgeries is as low as 1.5%. Also the studies involving hernia show even lower fraction of infection. This study was performed to evaluate the worth and efficacy of single dose prophylactic antibiotics compared to conventional antibiotics.

Study results of 60 patients studied with no loss to follow up revealed the following findings.

Out of the 30 patients belonging to group A i.e, who were given a single dose of prophylactic antibiotic, 2 developed signs of SSI within 1st week. And in group B, i.e, those who were given conventional multi dose antibiotics, out of 30 patients, only 1 developed SSI within 1st week. The overall p value was 1.000 which was not significant statistically.

As per CDC guidelines, study group i.e, group A, 1 had deep and 1 had superficial infections which was not significant as compared to the control group i.e, group B with a p value of 0.671

In our study the average cost of antibiotic between both the groups was 377 Rs in group A and 2900Rs in group B which was Highly Statistical Significance with p value

0.0005.

SSI grading as per Southampton grade was also insignificant with p value of 0.337. However, out of the 60 cases group 1 were converted to conventional antibiotics. The use of prophylactic antibiotic in all surgical cases has been advocated ever since, the concept of use of antibiotic preoperatively to curtain and prevent wound infection was postulated by Bernard and Cole in 1964.

A study conducted by **Jayalal JA et al**³², in which the patients in study group undergoing surgeries were given 1gm of injection cefotaxime after test dose 60 min prior to the surgery whereas in the control group, the patients were given 3 days of injection ciprofloxacin 200 mg intravenously twice a day, injection metronidazole 500mg intravenously thrice a day. The infection rate was similar in both groups with no significant differences.

Also a network meta-analysis by **T. Boonchan et al**⁸³ about antibiotic prophylaxis for prevention of surgical-site infection after groin hernia surgery which opined that beta lactam antibiotics were most effective SSI prophylaxis for groin hernia repair.

Naz et al⁸⁴ in a comparative study between a single-dose cefradine as the prophylactic antibiotics versus conventional dose of antibiotics in major gynaecological procedures have stated prophylactic antibiotic use is adequate provided standard principles of operative surgery are adhered.

A randomized clinical study conducted by **N Vinoth et al** on the role of antibiotic prophylaxis in open inguinal hernioplasty revealed that out of the sixty patients under 77 study 5 developed SSI which was 8.3 percent of which 3 were in the case group and 2 in control. They developed only superficial SSI with ODD'S ratio of 0.6429 which was statistically insignificant.

An operating surgeon should weigh the potential risk and also the benefits of giving an antibiotic after a particular procedure especially after a clean and uncontaminated surgery where the chances of SSI is very low.

Any improvement in quality of medical treatment can be attained by proper use of antibiotic which will be effective in preventing and controlling infection. The drug regimens should be optimised depending on the surgical procedure as it becomes burden on the economy.

However our study which was done to assess the effectiveness of a single dose of prophylactic antibiotic versus multiple doses antibiotics has shown no significant difference in the wound infection rate in both the studied groups.

CONCLUSION:

By the end of this study we come to a conclusion that use of single dose of antibiotic is as effective as multi dose antibiotic for a clean surgery of inguinal hernia repair in terms of surgical site infection. Additionally use of multi dose antibiotic will increase overall health care cost.

As this study involved only a smaller group of patients from a single institution the effect of operating room, duration of surgery and the surgeon leading to bias could not be assessed.

SUMMARY:

Randomized controlled study among 60 patients undergoing elective inguinal hernia surgeries in hospitals attached to RajaRajeswari Medical College, Bengaluru. 60 patients were randomized into two groups by randomization.

Patients in group A will be given single dose of 1.5gm cefaperazone + sulbactam intravenously half an hour before operation.

Patients in group B will be given multiple doses of antibiotic intravenously for 5 days post operatively.

1. Majority of the patients were in the age group of 50-60 years in both groups. There was no significant difference between the control and study group based on age.
2. General profile were comparable between two groups. In our study most of the patients were male . Again there was no significant difference between both the groups in sex wise distribution
3. Incidence of SSI was 3 in total among 60 patients in which 2 SSI were among (1 superficial SSI and 1 deep SSI) single dose group as compared to 1 superficial SSI in multiple dose group and the incidence of SSI among the two groups was statistically insignificant.
4. Among 3 patients who had SSI in single dose group ,1 patients had grade 3 , 1 had grade 4 SSI Southampton grades . whereas 1 patient with SSI in multiple dose group had grade 3 SSI.
5. In Patients receiving single dose antibiotic 29 patients were continued in the study group where as 1 patients who had grade 4 SSI was converted to full dose antibiotic.
6. The average cost of antibiotic in single dose group was significantly lesser when compared to multiple dose group.

REFERENCE

1. Malangoni MA, Rosen MJ. Hernias. Sabiston textbook of Surgery. 19th edn. Saunders Publication 2012: p. 1114-5.
2. Brunicaudi F, Andersen D, Billiar T, et al. Schwartz's principles of surgery. 10th edn. McGraw-Hill Education/Medical 2014: p. 1495-591.
3. Friedman DW, Boyd CD, Norton P, Greco RS, Boyarsky AH, Mackenzie JW, Deak SB. Increases in type III collagen gene expression and protein synthesis in patients with inguinal hernias. *Ann Surg* 1993;218(6): 754-760.
4. Öberg S, Andresen K, Rosenberg J. Etiology of Inguinal Hernias: A Comprehensive Review. *Front Surg*. 2017;4:52. Published 2017 Sep 22. doi:10.3389/fsurg.2017.00052
5. LeBlanc KE, LeBlanc LL, LeBlanc KA. Inguinal hernias: diagnosis and management. *Am Fam Physician*. 2013;87(12):844-848.
6. Kraft BM, Kolb H, Kuckuk B, et al. Diagnosis and classification of inguinal hernias. *Surg Endosc*. 2003;17(12):2021-2024. doi:10.1007/s00464-002-9283-y
7. Kulah B, Kulacoglu IH, Oruc MT, Duzgun AP, Moran M, Ozmen MM, et al. Presentation and outcome of incarcerated external hernias in adults. *Am J Surg*. 2001;181:101–104. [PubMed] [Google Scholar]
8. Akinci M, Ergül Z, Kulah B, Yilmaz KB, Kulacoglu H. Risk factors related with unfavorable

- outcomes in groin hernia repairs. *Hernia*. 2010;14:489–493. [PubMed] [Google Scholar]
9. Kulacoglu H. Current options in inguinal hernia repair in adult patients. *Hippokratia*. 2011;15(3):223-231.
 10. Shukla A, Mathur RK, Sheikh Z, et al. Nbutyl-2-cyanoacrylate glue versus suture for mesh fixation in open inguinal hernioplasty. *J. Evolution Med. Dent. Sci*. 2019;8(48):3575-3578.
 11. Hakeem A, Shanmugam V. Inguinodynia following Lichtenstein tension-free hernia repair: a review. *World J Gastroenterol*. 2011;17(14):1791-1796. doi:10.3748/wjg.v17.i14.1791.
 12. Amato B, Moja L, Panico S, et al. Shouldice technique versus other open techniques for inguinal hernia repair. *Cochrane Database Syst Rev*. 2012;2012(4):CD001543. Published 2012 Apr 18. doi:10.1002/14651858.CD001543.pub4
 13. Scott NW, McCormack K, Graham P, Go PM, Ross SJ, Grant AM. Open mesh versus non-mesh for repair of femoral and inguinal hernia. *Cochrane Database Syst Rev*. 2002;(4):CD002197. doi:10.1002/14651858.CD002197.
 14. Gaynes RP, Culver DH, Horan TC, Edwards JR, Richards C, Tolson JS. Surgical site infection (SSI) rates in the United States, 1992-1998: the National Nosocomial Infections Surveillance System basic SSI risk index. *Clin Infect Dis*. 2001 Sep 1;33 Suppl 2:S69-7
 15. Ghafur A, Jayalaletal JA. The Chennai declaration. *Indian J Cancer* 2012;49:71
 16. Erdas E, Medas F, Pisano G, Nicolosi A, Calò PG. Antibiotic prophylaxis for open mesh repair of groin hernia: systematic review and meta-analysis. *Hernia*. 2016;20(6):765-776. doi:10.1007/s10029-016-1536-0
 17. Ranjan A, Singh R, Naik PC. A comparative study of single-dose preoperative antibiotic prophylaxis versus routine long-term postoperative prophylaxis in elective general surgical cases. *Int J Med Sci Public Health* 2016;5:1083-1087.
 18. Basavarajappa M, Mathad AS, Deepak G. A Comparative Study of Effect of Single-Dose Prophylactic Antibiotic versus Conventional Antibiotic Therapy in Selected Clean Surgical Cases in A Tertiary Care Centre.
 19. Basant RK, Kumar R, Pandey VK, Saxena A, Singh V, Madeshiyaa S. A comparative study of single dose preoperative antibiotic prophylaxis versus fiveday conventional postoperative antibiotic therapy in patient undergoing elective surgical procedure. *International Surgery Journal*. 2019 Jan 28;6(2):409-15.
 20. Sumukha SR. Efficacy of single dose versus multiple dose antibiotic prophylaxis in anterior abdominal wall hernia repair at a tertiary hospital. *International Journal of Surgery*. 2020;4(3):230-2
 21. Resident NC. Outcome of Evidence-based Allocation of Single-dose Antibiotic extended to Three- dose Antibiotic Prophylaxis in Surgical Site Infection. *International Journal of Recent Surgical and Medical Sciences*. 2017 Dec;3(02):079-84.
 22. Shah YD, Thekdi PI, Raut S, Patel KG. Single shot versus multiple shot antibiotic therapy in patients undergoing laparoscopic surgery: our experience. *Int J Res Med Sci* 2013;1:252-6
 23. Shaikh SA, Iqbal M. Comparison of Single Dose with Multiple Dose Antibiotic Prophylaxis with Cefuroxime in Open Cholecystectomy. *Journal of Islamabad Medical & Dental College*. 2012;1(1):2-5.
 24. Madhu BS, HB SK, Reddy AV, Kalabhairav S. Effect of single dose pre-operative antibiotic prophylaxis versus conventional antibiotic therapy in patients undergoing lichtenstein tension free mesh repair. *International Surgery Journal*. 2017 Jan 25;4(2):738-42.
 25. Khan GJ, KHAN RA, Rasheed A. Comparison between single and multiple days treatment with parenteral cefuroxime in inguinal hernia mesh repair surgery. *Biomedica*. 2013 Jul;29(3):169-74.
 26. Surahio AR, Khan AA, Farooq MU, Fatima I. Single versus 3-dose antibiotic prophylaxis in clean and clean contaminated operations. *Journal of Ayub Medical College Abbottabad*. 2010 Dec1;22(4):91-4.
 27. Garg S, Garg S, Shoaib M. Efficacy of single dose versus multiple dose antibiotic prophylaxis in clean and clean-contaminated herniorrhaphy surgery. *J. Evid. Based Med. Healthc*. 2018; 5(39), 2786-2789.

28. Sutariya PK, Thekdi PI. Single dose versus multiple dose prophylactic antibiotic in laparoscopic cholecystectomy: a comparative study. *International Surgery Journal*. 2016 Dec 8;3(2):633-6.
29. Jeong WK, Park JW, Lim SB, Choi HS, Jeong SY. Cefotetan versus conventional triple antibiotic prophylaxis in elective colorectal cancer surgery. *J. Korean Med. Sci.* 2010; 25(2): 429–34.
30. Borade SV, Syed O. Single dose antibiotic prophylaxis for prevention of surgical site infection in elective surgery. *International Surgery Journal*. 2017 Dec 26;5(1):27-33.
31. Shaheen S, Akhtar S. Comparison of single dose versus multiple doses of antibiotic prophylaxis in elective caesarian. *J Postgrad Med Inst* 2014; 28(1):83-6.
32. Jayalal JA, Selwyn, Thambithurai D. Effect of Single-Dose Antibiotic Prophylaxis versus Conventional Antibiotic Therapy in Surgery: A Randomized Controlled Trial in a Public Teaching Hospital. *Int J Sci Stud* 2015;3(8):109-113.
33. Dr. Soundara Rajan, Aswathy Harikumar, Sreelekshmi. To assess the efficacy of single dose prophylactic antibiotics in preventing SSI after clean elective surgery. *J Case Rep Sci Images* 2021;3(1):01-03
34. K. Saravanan, A. Muthukumar. Prophylaxis pre-operative single dose antibiotic versus conventional antibiotic therapy in lichtenstein tension free mesh repair. *International Journal of Contemporary Medicine Surgery and Radiology*. 2019;4(2):B171-B173.
35. Thapa SB, Kher YR, Tambay YG. Single dose Intraoperative Antibiotics versus Postoperative Antibiotics for Patient Undergoing Laparoscopic Cholecystectomy for Symptomatic Cholelithiasis: a Randomized Clinical Trial. *Journal of Lumbini Medical College*. 2017 Jul 5;5(1):13-7.
36. Diwaker A, Datey A, Verma D, Bandi A. A comparative study of Single dose versus multiple doses of antibiotic prophylaxis in open inguinal hernioplasty. *IOSR J Dental Med Sci*. 2018;17(5):28-32.
37. Chaudhary R, Sharma S, Chaudhary S, Thakur S, Shukla A, Sharma A. A prospective study comparing single with multiple antibiotic prophylaxis dose in elective cholecystectomy. *Ann Int Med Den Res*. 2015;1(1):29-33.
38. Subburathinam DA, Karthick MP, Princy J. Effectiveness of single over triple dose antibiotic prophylaxis in open inguinal hernioplasty. *International Journal of Surgery*. 2021;5(1):297-300.
39. Javid PJ, Brooks DC. Hernias. In: Zinner MJ, Ashley SW, eds. *Maingot's Abdominal operations*, 11th Ed. New York: Mc Graw Hill, 2007: 103-140.
40. Bell RL, Seymour NE. Abdominal wall, omentum, mesentery, retroperitoneum. Brunicaudi FC, ed. *Schwartz's principles of surgery*, 8th Ed. New York: Mc Graw Hill, 2005:1317- 1328.
41. Haeger K. *The illustrated history of Surgery*. London: Harold Starke, 1988.
42. Sinnatamby Cs. Anterior abdominal wall. *Last's anatomy regional and applied*, 10th ed. Churchill Livingstone, 2000; 215- 226.
43. Russell RCG, Williams NS, Bulstrode CJK. Hernias, umbilicus and abdominal wall, *Bailey & Love's – short practice of surgery*, 24th ed. London, Arnold, 2004; 1272- 1293.
44. Chaurasia BD. Anterior abdominal wall, *Human Anatomy Regional & Applied*, 3rd Ed, CBS Publishers, 2001; 163-180.
45. Singh IB. *Textbook of Anatomy with Colour Atlas*. 5th ed. Delhi. jaypee Brothers; 2007. Chapter 39 p550
46. Lytle WJ. Inguinal anatomy. *J Anat*. 1979 May;128(Pt 3):581-94
47. Chaurasia BD. Anterior abdominal wall, *Human Anatomy Regional & Applied*, 3rd Ed, CBS Publishers, 2001; 163-180
48. Sinnatamby CS. Anterior abdominal wall. *Last's anatomy regional and applied*, 10th Ed. Churchill Livingstone, 2000; 215-226.
49. Decker GAG. The groin & scrotum. *Lee Mc Gregor's Synopsis of Surgical Anatomy*. 12th ed, Varghese Publishing house, 1999; 118-136.
50. Russell RCG, Williams NS, Bulstrode CJK. Hernias, umbilicus and abdominal wall. *Bailey & Love's - Short practice of surgery*, 25th Ed. London: Arnold, 2008; 968-

51. Read RC, Anatomy of Abdominal Herniation: The Parietoperitoneal spaces. In: Nyhus LM, Baker RJ, Fischer JE, Mastery of Surgery, 3rd ed. Little, Brown and Company, 1997; 1795-1806.
52. Javid PJ, Brooks DC. Hernias. In: Zinner MJ, Ashley SW, eds. Maingot's Abdominal operations, 11th Ed. New York: Mc Graw Hill, 2007: 103-140.
53. Casten DF. Functional anatomy of the groin area as related to the classification and treatment of groin hernias. Am J Surg. 1967 Dec; 114(6):894-9.
54. Halverson K. Inguinal and Femoral Hernioplasty: A 22-Year Study of the Authors' Methods. Arch Surg. 1970 Aug 1; 101(2):127.
55. Ponka JL. Surgical management of large bilateral indirect sliding inguinal hernias Am J Surg. 1966 Jul; 112(1):52-7.
56. The Aachen classification of inguinal hernia [Internet]. Available from: https://www.researchgate.net/publication/298655224_The_Aachen_classification_ofinguinal_hernia
57. Bendavid R. Classification of inguinal hernias: The T.S.D. Classification – A Nomenclature for Groin Hernias. Surg Laparosc Endosc Percutan Tech. 1994 Dec; 4(6):467.
58. Chowbey Pradeep K., et al. Complications in groin hernia surgery and the way out. Journal of minimal access surgery. 2006; 3:2:174- 177
59. S Das. A manual on clinical surgery. Calcutta 13th edition 2018; 594-610.
60. S Das. A manual on clinical surgery. 4th ed. Calcutta; 1996. 428-444 p.
61. A C. Demonstration of physical signs in clinical surgery. 17th ed. 1992. 260-272.
62. NL B. An introduction to the symptoms and signs of surgical diseases. 2nd ed. ELBS; 1991. 310-321 p.
63. J A. Hernias. Maingot's abdominal operations. 10th ed. McGraw Hill; 2001. 479-580 p.
64. Devlin HB. Inguinal hernia in adults. In: Carter D, Russell RCG, Pitt HA. Atlas of General Surgery. 3rd ed. London: Arnold 1996; 38-49.
65. Farquharson M, Moran B eds. Surgery of the groin and external genitalia. In: Farquharson's Textbook of Operative General Surgery. 9th Ed. London: Edward Arnold, 2005: 459-484.
66. Farquharson M, Moran B eds. Surgery of the groin and external genitalia. In: Farquharson's Textbook of Operative General Surgery. 9th Ed. London: Edward Arnold, 2005: 459-484.
67. Shenoy KR, ed. Hernia. In: Manipal manual of surgery, 2nd Ed. New Delhi: CBS, 2005: 551-573.
68. Palanivelu C, ed. Transabdominal preperitoneal hernioplasty. In: Operative manual of laparoscopic hernia surgery, 1st Ed. Coimbatore: GEM, 2004: 83-98.B
69. Palanivelu C, ed. Totally extraperitoneal hernioplasty. In: Operative manual of laparoscopic hernia surgery, 1st Ed. Coimbatore: GEM, 2004: 99-118.
70. Ayliffe. G.A.J., Fraiese A.P., et al, Control of hospital infection, 4th edn, Arnold, Newyork.
71. Ad hoc committee on Trauma, Division of Medical Sciences, National Academy of sciences, National Research Council 'Post operative wound infections. The influence of ultraviolet irradiation of the operating room and of various other factors'. Ann surg 1964; 160 (Supp 13); 1-32.
72. Cruse. P.J.E, Foord R, „The epidemiology of wound infection. A ten year prospective study of 62, 939 wound“. Surg Clin North Am 1980; 60; 27-40.
73. Edwards.L.D., „The Epidemiology of 2056 Remote site infections and 1966 Surgical wound infections occurring in 1865 patients.“ Ann.surg. 1976, 184 (10): 758-66.
74. Hunt K.T., Reid.V.Muller, „Inflammation, Infection & antibiotics Chapter 8 in Medical Management of the surgical patient. Edtd., by Michael Lubig et al, 3rd edn., J.Blippincott co., Philadelphia.
75. Olson M.M., James. T.Lee 'Continuous, 10-year wound Infection Surveillance; results, advantages and unanswered questions' Arch Surg. 1990: 794-803.
76. Nichols Ronald lee "Post Operative wound infections" The New England Journal of Medicine, 1982; 307: 1701-02.
77. Mangram AJ, Nichols RL. Recognition, preparation, surveillance and management of surgical

- site infection. Hospital infection control practices advisory Committee. Infect Control Hosp Epidemiol 1999; 20:250.
78. Peter J. Morris, Ronald A. Mact. Wound infection, Oxford Text Book of Surgery. Vol. 1 p. 27-43. 1994.
 79. Mackie, McCartney. Practical Medical Microbiology : 13th edition, Wound, Skin and deep sepsis p. 619.
 80. Ahrenholz DH, Simmons RL. Mixed and synergistic infection, in Howard RJ, Simmons RL (eds) : Surgical Infectious diseases, 2nd ed, Norwalk, CT, Appelton and Lange, 1991 ; Chapt. 2, P. 187.
 81. Sawyer Robert .G., and Timothy L. pruet, "Wound Infections". Surgical Clinics of North America, 1994, 74; 519 –36
 82. Page C.P., et al, 'Antimicrobial Prophylaxis for surgical wounds; Guidelines for clinical care' Arch surg- 1993 Jan; 128; 79-88
 83. Boonchan T, Wilasrusmee C, McEvoy M, Attia J, Thakkestian A. Network metaanalysis of antibiotic prophylaxis for prevention of surgical-site infection after groin hernia surgery. Br J Surg. 2017;104(2):e106-e117. doi:10.1002/bjs.10441.
 84. Garg, S., Garg, S., & Shoaib, M. (2018). Efficacy of single dose versus multiple dose antibiotic prophylaxis in clean and clean-contaminated herniorrhaphy surgery. *Journal of Evidence Based Medicine and Healthcare*, 5(39), 2786–2789. <https://doi.org/10.18410/jebmh/2018/570>