

A Quasi Experimental Study to Assess the Effectiveness of Abdominal Binder on Ambulatory Pain among Post-operative Patients at Selected Hospital, Coimbatore

Amirthaveni, K.¹, Kanchana. K², Nirmala T³

¹M. Sc Nursing Student, College of Nursing, Sri Ramakrishna Institute of Paramedical Sciences, Coimbatore, Tamil Nadu, India.

²Professor, College of Nursing, Sri Ramakrishna Institute of Paramedical Sciences, Coimbatore, Tamil Nadu, India.

³Principal, College of Nursing, Sri Ramakrishna Institute of Paramedical Sciences, Coimbatore, Tamil Nadu, India.

ABSTRACT

Abdominal binders are effective in reducing ambulatory pain and improving early ambulation. The study aimed to identify the effect of abdominal binder on ambulatory pain among post-operative patients with abdominal surgery. Quasi-experimental post-test-only control group design was adopted in this study. 30 patients in experimental group and 30 patients in control group were selected based on inclusive and exclusive criteria by using purposive sampling technique. The demographic variables and clinical variables were collected from the postoperative patients. Abdominal binder was applied to the experimental group and the control group received routine postoperative care. Post-test was done for both groups to assess the ambulatory pain by using a numerical pain rating scale twice a day for first three postoperative days. It was identified on the first day that the mean levels of ambulatory pain for the experimental and control group were 8.35 and 9.07 with a mean difference of 0.72. Standard deviations were 0.787 and 0.435 respectively. The calculated value of 0.286 was less than the table value at 0.05 level of significance. On the second day mean levels of ambulatory pain for the experimental and control group were 5.93 and 7.28 with a mean difference of 1.35. Standard deviations were 0.81 and 0.692 respectively. The calculated value of 6.035 was greater than the table value at a 0.001 level of significance. On the third day mean levels of ambulatory pain for the experimental and control group were 3.28 and 5.18 with a mean difference of 1.9. Standard deviations were 0.663 and 0.748 respectively. The calculated value of 3.653 was greater than the table value at a 0.001 level of significance. Hence, the researcher concludes that the abdominal binder is a non-elastic, cost effective method on reducing the ambulatory pain among postoperative patients.

KEYWORDS: Abdominal binder, post-operative patients, early ambulation, postoperative care, abdominal surgery

INTRODUCTION

Pain is an unpleasant sensory and emotional experience which may be associated with actual or potential tissue damage. Surgical trauma induces hyperalgesia which could lead to chronic pain in the postoperative period when left unattended (Sundeeep, 2018). The effective relief of pain is of the utmost importance to treating patients undergoing surgery. Pain relief has significant physiological benefits hence, monitoring of pain relief is increasingly becoming an important postoperative quality measure. The goal for postoperative pain management is to reduce or eliminate pain and discomfort with a minimum of side effects (Veerabhadram Garimella, 2013). Post-Surgical Pain is common and expected after surgery. Effective post-surgical pain management is associated with patient satisfaction, earlier mobilization, shortened hospital stays, and reduced costs. The goal of pain management following a surgical procedure is to prevent and control pain (Saramma, 2010).

NEED FOR THE STUDY

Globally, a staggering 310 million major surgeries are performed each year. It is estimated that 1–4% of these patients will die, up to 15% will have serious postoperative morbidity, and 5–15% will be readmitted within 30 days. (Dobson GP, 2020). After abdomen surgery, participants should support the incision area with a pillow or their hands during mobilization. Although, it is not possible to provide constant support all the time; hence, using an abdominal binder is a practical and common application that helps mobility and recovery. It has been reported that using abdominal binder might decrease the pain following major abdominal surgery by limiting motion and supporting abdominal wall during recovery period, compression at surgical site, increases blood flow and reduces inflammation, accordingly improves rapid tissue repair. Some studies specified that additional benefits of abdominal binder including the prevention of herniation, wound seroma and hematoma (Boonploeng et al., 2021). These facts stimulated the researcher to investigate the effectiveness of abdominal binder on ambulatory pain among postoperative patients.

STATEMENT OF THE PROBLEM

A Quasi Experimental Study to Assess the Effectiveness of Abdominal Binder on Ambulatory Pain among Post-operative Patients at Selected Hospital, Coimbatore.

OBJECTIVES OF THE STUDY

- To assess the level of ambulatory pain among postoperative patients
- To evaluate the effectiveness of abdominal binder on ambulatory pain among postoperative patients.
- To find out the association between the level of ambulatory pain and selected variables among postoperative patients.

OPERATIONAL DEFINITION

Effectiveness

Effectiveness refers to the changes in the level of ambulatory pain after the application of abdominal binder among postoperative patients. It is measured by Numerical Pain Rating Scale (NPRS) twice a day for first three postoperative days.

Abdominal Binder

Abdominal binder refers to a non-elastic belt that encircles abdomen. It is used to support the incisional

site by compressing and stabilizing the abdominal muscles for patient who underwent abdominal surgeries.

Ambulatory Pain

It refers to the pain experienced by the patient during ambulation after an abdominal surgery, as measured by numerical pain rating scale.

Postoperative patients

It refers to patients who underwent open abdominal surgery with vertical incision.

Hypothesis

H₁ -There is a significant difference in the level of ambulatory pain after the application of abdominal binder in experimental group and control group.

H₂ - There is a significant association between the levels of ambulatory pain and selected variables among postoperative patients.

METHODOLOGY

RESEARCH APPROACH: Quantitative Approach

RESEARCH DESIGN: Quasi- experimental post test only control group design.

RESEARCH SETTING: The study was conducted in, surgical, post-operative ,gynecology and special wards at Sri Ramakrishna Hospital, Coimbatore

TARGET POPULATION

Patients who underwent abdominal surgery in Coimbatore

ACCESSIBLE POPULATION

Patients who underwent abdominal surgery at Sri Ramakrishna Hospital, Coimbatore

SAMPLING TECHNIQUE

Non probability Purposive sampling (n=60)

SAMPLING SIZE:

60 samples who underwent abdominal surgery.

Criteria for Sample Selection

The purposive sampling method is employed to pick 60 patients. Patients were chosen for the study based on inclusion and exclusion criteria.

Inclusion criteria

- Patients with vertical incision in the abdomen.
- Patients with laparotomy.
- Patients with hysterectomy
- Patients who are alert and cooperative.
- Patients who can be ambulated after getting surgeon's opinion.

Exclusion criteria

- Patients with Cesarean section.
- Patients who are critically ill
- Patients undergoing laparoscopic surgery

- Patients with orthopedic, neuromuscular and circulatory disorders.
- Epidural analgesics

DATA COLLECTION INSTRUMENT

Non- randomized quasi experimental post test only control group design was adopted. By using the purposive sampling technique 60 study participants were selected based on inclusion and exclusion criteria were assigned alternately so as to have 30 in experimental and 30 in control group. The intervention for the study was application of abdominal binder. Abdominal binder was applied for the patients on the first postoperative day onwards and continued in the experimental group and control group received routine nursing care. For the first three postoperative days, ambulatory pain assessed using the Numerical Pain Rating Scale (NPRS) twice daily for both the experimental and controls groups.

**DATA ANALYSIS AND INTERPRETATION
SECTION I**

Demographic details of post-operative patients with abdominal surgery

(n =60)

S.No	in years	Number of patients			
		Experimental group (n=30)		Control group (n=30)	
		Frequency	Percentage (%)	Frequency	Percentage (%)
Age					
1	18 - 30	4	13.3	3	10.0
2	31 - 40	5	16.7	6	20.0
3	41 - 50	11	36.7	6	20.0
4	51 - 60	7	23.3	7	23.3
5	Above 60	3	10.0	8	26.7
Gender					
1	Male	8	26.7	8	26.7
2	Female	22	73.3	22	73.3
Religious Status					
1	Hindu	25	83.3	21	70
2	Muslim	4	13.3	5	16.7
3	Christian	1	3.3	4	13.3

Dietary pattern					
1	Vegetarian	8	26.7	6	20
2	Non-Vegetarian	22	73.3	24	80

Educational status					
1	Illiterate	4	13.3	6	20.0
2	Primary	9	30.0	2	6.7
3	Secondary	11	36.7	11	36.7
4	Graduate and above	6	20.0	11	36.7
Occupation					
1	Government	3	10.0	1	3.3
2	Private	8	26.7	6	20.0
3	Business	3	10.0	2	6.7
4	Daily Wages	1	3.3	1	3.3
5	Unemployed	15	50.0	20	66.7
Marital Status					
1	Single	3	10	2	6.7
2	Married	21	70	21	70.0
3	Widow	6	20	4	13.3
4	Separated	-	-	3	10
Type of Family					
1	Joint	14	47	18	60
2	Nuclear	16	53	12	40
Family Monthly Income					
1	Less than 10000	1	3.3	-	-
2	10,001 to 20,000	3	10	2	6.7
3	20,001 to 30,000	5	16.7	4	13.3
4	30,000 above	21	70	24	80

SECTION II

CLINICAL VARIABLES	No. of Patients			
	Number of patients		Number of patients	
	Experimental group (n=30)	Experimental group (n=30)	Control group (n=30)	Control group (n=30)

Diagnosis	Abdominal blunt injury				
	Abdominal Pelvic Mass	1	3.3	1	3.3
	Abnormal Uterine Bleeding	1	3.3		
	Appendicitis	5	16.6	4	13.3
	Hernia	2	6.6	1	3.3
	Fibroid uterus	8	26.6	10	33.3
	Cancer in uterus	3	10	4	13.3
	Pancreatitis	3	10	5	16.6
	Cancer in abdomen	2	6.6	1	3.3
	Jejunal perforation	4	13.3	2	6.6
	Recurrent pyogenic cholangitis			1	3.3
	Sigmoid diverticulitis	1	3.3		
	Surgical Procedure	Emergency laparotomy	3	10	3
Elective laparotomy		9	30	10	33.33
Abdominal Hysterectomy		10	33.33	10	33.33
Hernia repair		8	26.67	7	23.34
Body Temperature (°F)	Hypothermia (<95)	-	-	-	-
	Normothermia(96-98.9)	23	76.7	21	70
	Hyperthermia(99-100)	7	23.3	9	30
Pulse Rate (beats per minute)	70-80	16	53.33	12	40
	80-90	12	40	16	53.33
	90-100	2	6.67	2	6.67
Respiratory Rate (breaths per minute)	18- 22	25	83.3	21	70
	22 - 30	5	16.7	9	30
Blood Pressure (mm.hg) Systolic Blood Pressure	100-110	13	43.3	7	23.33
	111-120	3	10	10	33.33
	121-130	10	33.3	11	36.67
	131-140	2	6.6	1	3.33
	> 141	2	6.6	1	3.33
Blood Pressure Diastolic Blood Pressure	60-70	11	36.6	10	33.33
	71-80	13	43	17	56.67
	81-90	6	20	3	10

Body Mass Index	Underweight (18.5)	5	16.7	4	13.3
	Normal (18.5-24.5)	15	50	13	43.3
	Over weight (25.0-29.9)	3	10	6	20
	Obesity (Above30)	7	23.3	7	23.3
Comorbidity	Yes	15	50	17	56.67
	No	15	50	13	43.33
Previous Surgery	Yes	22	73.3	22	73.3
	No	8	26.7	8	26.7
Duration of Surgery	Less than 3Hrs	16	53.3	20	66.7
	More than 3 Hrs	14	46.7	10	33
Anesthesia	General	17	56.7	11	36.7
	Spinal	8	26.7	6	20
	Both	5	16.7	13	43.3
Recovery from Anesthesia	≤ 3 Hours	12	40	18	60
	>3 Hours	18	60	12	40
Abdominal Girth	≤85cm	17	56.7	15	50
	>85cm	13	43.3	15	50
Length of Surgical Incision	Less than 15 cm	16	53.3	17	56.7
	More than 15 cm	14	46.7	13	43.3
Surgical Dressing	Grip Band Plaster	21	70	21	70
	Gauze Pad	9	30	9	30
Presence of Drain	No drain tube	8	26.7	10	33.3
	Romovac drain	7	23.3	8	26.7
	Romovac and pvc drain	15	50.0	12	40
Post Operative Diet	Empty Stomach	13	43.3	10	33.3
	Liquid Diet	15	50.0	8	26.7
	Bland Diet	2	6.7	12	40

Assessment on level of ambulatory pain among post operative patients with abdominal surgery n=60

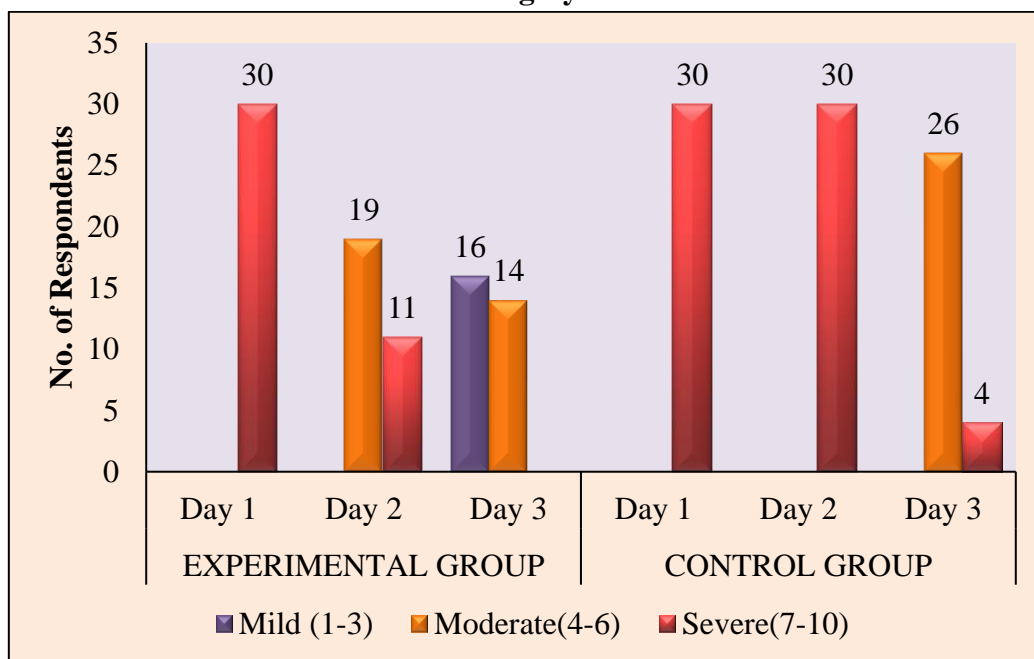
S. No	LEVEL OF AMBULATORY PAIN	Number of Patients					
		Experimental group (n=30)			Control group (n=30)		
		POD 1	POD 2	POD 3	POD 1	POD2	POD 3
		Frequency	Frequency	Frequency	Frequency	Frequency	Frequency

1	Mild (1-3)	-	-	16	-	-	-
2	Moderate (4-6)	-	19	14	-	-	26
3	Severe (7-10)	30	11	-	30	30	4

The above table 4.3.1 depicts the assessment on level of ambulatory pain among Post-Operative Patients with Abdominal surgery. The results showed that in the experimental and control group 30 (100%) patients had severe pain on the first postoperative day. On the second day, 19 (63.33%) patients had moderate pain and 11 (36.67%) patients had severe pain in the experimental group. On the other hand, 30 (100%) patients had severe pain in the control group.

On the third day, 16 (53.3%) patients had mild pain, and 14 (46.67%) patients had moderate pain in the experimental group. On the other hand, 26 patients had moderate pain, and 4 (13.33%) patients had severe pain in the control group.

Chart- Assessment on level of ambulatory pain among post operative patients with abdominal surgery



Effect of abdominal binder on ambulatory pain among post operative patients with abdominal surgery in the experimental and control group
n=60

POD*	Group	Mean	SD	Mean difference	'z' Value	Table value
I	Experimental group	8.35	0.787	0.72	0.286	1.96

	Control group	9.07	0.435			
II	Experimental group	5.93	0.81	1.35	6.037	3.29***
	Control group	7.28	0.692			
III	Experimental group	3.28	0.663	1.9	3.653	3.29***
	Control group	5.18	0.748			

*-POD-Post Operative Day

***significant at 0.001 level

RESULTS AND DISCUSSION

The analysis showed that the mean level of pain among experimental and control group was 8.35 and 9.07 respectively with mean difference of 0.72. The calculated ‘z’ value 0.286 was less than table value of 1.96 at 0.05 level of significance. Hence, the hypothesis H_1 is rejected there is no significant difference in the level of ambulatory pain among post operative patients with abdominal surgery in the experimental and control groups on the first post operative day.

The analysis showed that the mean level of pain among experimental and control group was 5.93 and 7.28 respectively with mean difference of 1.35. The calculated ‘value 6.037 was greater than table value of 2.58 at 0.001% level of significance. Hence, hypothesis H_1 is accepted, there is a highly significant difference in the level of ambulatory pain among post operative patients with abdominal surgery in the experimental and control groups on second post operative day.

CONCLUSION

Postoperative ambulatory pain is common for abdominal surgeries. The planned post-operative nursing care will reduce the ambulatory pain and improve the early ambulation. The abdominal binders help to reduce the post-operative ambulatory pain and preserve the hemodynamic stability. The study shows that application of abdominal binder on incisional site results in reduction of ambulatory pain during post-operative period. Hence it is concluded that the abdominal binder is a non - elastic, cost effective method in reducing the ambulatory pain among post operative patients.

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