**Research Article** 

# Effectiveness of Guided Imagery Technique on pain among terminally Ill Patients at selected Hospitals of Jaipur

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**Techniques** 

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## Abstract

Terminal illness is an incurable disease that cannot be adequately treated. Pain at terminal illness is significant such as cancer, late HIV disease, and degenerative diseases. The aim of study to evaluate the effectiveness of guided imagery technique regarding pain, among terminally ill patients. A quantitative experimental research approach with Quasi-experimental, non-equivalent control group design was used with 210 terminally ill patients (105 in experimental groups and 105 in control group) to evaluate the effectiveness of guided imagery technique regarding pain. Consecutive sampling technique was used. A socio-demographic data, pain assessment scale, was developed by the researcher to evaluate pain, among terminally ill patients. Researcher used Guided imagery technique as intervention for study. The guided imagery technique was found effective to reduce pain. In pain experimental group pre-test mean  $\pm$  S.D was 1.94  $\pm$ 1.36, mean difference was -3.7, t= 13.34, p=<.01, Significant, whereas in control group pre-test mean  $\pm$  S.D was 5.28  $\pm$  2.85, in follow up mean  $\pm$  S.D was 4.40  $\pm$ 2.22, mean difference was -.8, t=2.4, p=<.05, Significant. Hence, the result revealed that guided imagery technique was more effective to reduce the pain in experimental group as compared to control group.

**Keywords:** Effectiveness; Guided Imagery Technique; Pain; Terminally ill patients

# Introduction

Terminal illness is an incurable disease that cannot be adequately treated and is reasonably expected to result in the death of the patient within a short period of time. The term terminally illness is more commonly used for progressive diseases such as cancer or advanced heart disease than for trauma. A patient has such an illness may be referred to as a terminal patient, terminally ill or simply terminal. There is no standardized life expectancy for a patient to be considered terminal, although it is generally months or less [1].

Patients are nearing the end of their lives frequently feel severe pain. Unresolved pain has been highlighted as indicative of low quality end-of-life care. The data on that judgment is founded, however, is limited. Despite the fact that half of terminally ill patients were in moderate to severe pain, just 30% desired extra pain therapy from their primary-care physician. The number of patients in agony is still far too large. However, the figure is not as huge as it appears. Furthermore, most people are ready to put up with discomfort. Furthermore, pain is a consistent feeling across most terminal illnesses [2].

In the 1970s, Dr. David Bressler and Dr. Martin Rossman began establish support for guided imagery as an effective approach for the treatment of chronic pain, cancer, and other serious illnesses. Their work led them to co-found the Academy for Guided Imagery in 1989 [3].

#### Need of study

Terminal illness is a medical term popularized in the 20th century to describe an active and malignant disease that cannot be cured or adequately treated and that is reasonably expected to result in the death of the patient. Life expectancy for terminal patients is a rough estimate given by the physician based on previous data and does not always reflect true longevity [4].

Pain at terminal illness is significant such as cancer, late HIV disease, degenerative diseases; most people equate pain at the end of life

with terminal illness. Surveys of adult cancer patients with advanced disease—often performed in a hospice or palliative care setting indicate that the prevalence of pain ranges from 50% to 90%.<sup>16</sup> It is observed that 40-50% of those with pain from cancer report it to be severe while 25-30% describe it to be very severe [5].

There are many pharmacological and non-pharmacological treatments available for treatment of terminal illness. The pharmacological treatment have many adverse effects and develops tolerance which results in pain, among terminal ill patients but non-pharmacological treatment such as guided imagery technique is cheap, easy and free from any side effects. So, health personnel should educate terminally ill respondents regarding the benefits of non-pharmacological treatment.

#### The objective of the study

1. To assess the level of Pain, among terminally ill patients admitted in selected hospitals at Jaipur.

2. To evaluate the effectiveness of guided imagery technique on Pain, among terminally ill patients admitted in selected hospitals at Jaipur.

3. To find out the association of Pain, with selected demographic variables of terminally ill patients admitted in selected hospitals at Jaipur.

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#### Aim of the study

To evaluate the effectiveness of guided imagery technique regarding pain, among terminally ill patients.

#### Hypothesis

> H<sub>0</sub>1- There is no statically significant relationship between the obtained score of pain, among terminally ill patients admitted at selected hospitals in Jaipur.

 $H_0^2$ - There is no statically significant relationship between effect of guided imagery technique and score obtained on level of pain, among terminally ill patients admitted at selected hospitals in Jaipur.

ightarrow H<sub>0</sub>3- There is no statically significant relationship between pre-test score obtained on level of pain, with socio demographic variables among terminally ill patients admitted at selected hospital in Jaipur.

# **Review of Literature**

The review of literature in this study is divided in following sections:

Section-A: Prevalence of terminal illness

Section-B: Pain in terminal illness

Section-C: Guided Imagery Technique on terminally illness

#### Material and methods

A quantitative experimental research approach with Quasiexperimental, non-equivalent control group design was used which help the researcher to evaluate the effectiveness of guided imagery technique regarding pain. 216 terminally ill patients were selected with Consecutive sampling technique. The researcher consider three type of variable under study -Dependent Variable, Independent Variable, Attributed variable .the study was carried out at 3 selected hospitals of Jaipur. A socio-demographic data, pain assessment scale, was developed by the researcher to evaluate pain, among terminally ill patients. Socio-Demographic Performa consist of 14 items which include information of respondents about Age, Gender, Religion, Marital status, Habitat, Education qualification, Occupation, Family type, Monthly family income, Source of major income in family, Area of hospitalization, Duration of taking treatment, Types of disease and Level of dependency. Researcher was use standardized Numerical universal pain assessment scale. The scale was created by Mc Gill University's Dr. Ronald Melzack and Dr. Warren Torgerson [6].

Researcher used Guided imagery technique as intervention for study. Tool and Guided Imagery Technique was validated with 14 different subject expert Numerical pain assessment scale is a standard tool and reliability already established and tested by the valedictory authority. Physiopedia a nonprofit organization and worldwide physiotherapist community established reliability of numerical pain assessment scale with test – retest method r=.95. 97.

A formal permission was obtained from the concern authority. The data were collected from Dec. 2020 to July 2021. The researcher collected the data from 210 terminally ill patients. A total of 210 (105 in experimental group and 105 in control group) respondents were selected for the study. Researcher obtained informed written consent from each respondent. Ethical approval was obtained from the institutional ethical committee to conduct the study. Direct face to face interview was conducted and confidentiality of study was assured. The data was

analyzed in the term of objectives of the study using descriptive and inferential statistic. A master sheet was prepared by the researcher as response is given by respondent. The effectiveness of Guided Imagery Technique was analyzed by using t-test for experimental and control group. The association between pre-test score of numerical pain assessment, with selected socio demographic variables was analyzed by chi square test for experimental and control group. Experimental and control group data were presented in table, graph and diagram.

#### Result

Section-1: Distribution of the socio-demographic variables.

**Section-2:** Pre-test assessment of pain among terminally ill patient.

Section-3: Effectiveness of guided imagery technique on pain.

**Section-4:** Association between pre-test score of pain, with selected socio-demographic variables.

**SECTION-1:** Distribution of the socio-demographic variables. (Table 1)

Table 1: Distribution of the socio-demographic variable
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SI. No.	Socio- Demographic		MENTAL OUP	CONTROL FROUP		
	Variable	Frequency (f)	Percentage (%)	Frequency (f)	Percentage (%)	
1	Age in Years					
	21-30 Years	25	23.8	26	24.8	
	31-40 Years	23	21.9	22	21.0	
	41-50 Years	23	21.9	19	18.1	
	51-60 Years	25	23.8	26	24.8	
	Above 60 Years	09	8.60	12	11.4	
2	Gender					
	Male	55	52.4	58	55.2	
	Female	50	47.6	47	44.8	
	Trans-Gender	0	0	0	0	
3	Religion					
	Hindu	42	40.0	54	51.4	
	Muslim	35	33.3	22	21.0	
	Christian	03	2.9	05	4.8	
	Any Other	25	23.8	24	22.9	
4	Marital Status					
	Married	94	89.5	96	91.4	
	Unmarried	04	3.8	05	4.8	
	Separated	06	5.7	03	2.9	
	Divorced	01	1.0	01	1.0	
5	Habitat					
	Urban	39	37.1	36	34.3	
	Semi-Urban	24	22.9	25	23.8	
	Rural	42	40	44	41.9	
6	Educational Qualification					
	Non-Formal Education	02	1.9	03	2.9	
	Primary Education	06	5.7	15	14.3	
	Secondary Education	20	19.0	24	22.9	
	Senior Secondary	32	30.5	36	34.3	
	Graduate And Above	45	42.9	27	25.7	

7	Occupation				
	Unemployed	06	5.7	09	8.6
	Unskilled Labor	20	1.9	06	5.7
	Govt. Employee	20	19.0	14	13.3
	Private Employee	24	22.9	28	26.7
	Self Employed	53	50.5	48	45.7
8	Family Type				
	Nuclear	11	10.5	12	11.4
	Joint	59	56.2	50	47.6
	Extended	35	33.3	43	41.0
9	Monthly Income				
	Less Than 5000 Inr	06	5.7	02	1.9
	5001-10000 Inr	06	5.7	15	14.3
	10001-15000 Inr	37	35.2	18	17.1
	15001-20000Inr	29	27.6	29	27.6
	More Than 20000Inr	27	25.7	41	39.0
10	Source of Major Income				
	Patient Self	65	61.9	68	64.8
	Parents	13	12.4	07	6.7
	Sibling	11	10.5	07	6.7
	Child	09	8.6	13	12.4
	Any Other	07	6.7	10	9.5
11	Area of Hospitalization				
	Palliative Unit	97	92.4	96	91.4
	Art Centre	08	7.6	09	8.6
12	Duration of Taking Treatment				
	Less Than 1 Yr	62	59.0	56	53.3
	1 To 2 Yrs	32	30.5	28	26.7
	More Than 2 Years	11	10.5	21	20.0
13	Level of Dependency				
	Independent	22	21.0	12	11.4
	Partially Dependent	76	72.4	71	67.6
	Completely Dependent	07	6.7	22	21.0

Table 1 showed the distribution of respondent according to age. In experimental group 23.8% belonged to 21-30yrs and 51-60 yrs., 23.% belonged to 31-40 yrs. and 41-50yrs and only 8.6% respondent belonged to above 60yrs of age whereas in control group most of respondent 24.8% belonged to age group of 21-30yrs and 51-60 yrs. 21% belonged to 31-40yrs, 18.1% belonged to 41-50 yrs. and only 11.4% respondent belonged to above 60 years age. In experimental group majority of respondent were males 52.4%, females 47.6%, whereas in control group majority of respondent were also males 55.2%, females 44.8% and transgender 0.0% were in both experimental and control group. majority of sample in experimental group were Hindus 40%, Muslim 33.3%, any other 23.8%, Christian 3% whereas in control group majority of respondent were Hindu 51.4%, any other 22.9%, Muslim 21%, Christian 4.8%.

In experimental group majority of respondents were married 89.5%, separated 5.7%, unmarried 3.8% and divorced 1% whereas in control group majority of respondents were married 91.4%, unmarried 4.8%, separated 2.9% and divorced 1%. In experimental group most of respondents belonged to rural area 40.0%, urban area 37.1%,

semi-urban 24.0% area whereas in control group most of respondent belonged to rural area 41.9%, urban area 34.3% and semi-urban area 23.8%. In experimental group most of respondent were Graduate and above 42.9%, followed by senior secondary 30.5%, secondary education 19%, primary education 5.7%, and 1.9% respondent had non-formal education whereas in control group most of respondent were senior secondary 34.3% followed by graduate and above 25.7%, secondary education 22.9%, primary education 14.3% and only 2.9% respondent had non-formal education.

In experimental group most of respondent were self-employed 50.5%, private employee 22.9%, Govt. employee 19%, unemployed 5.7%, unskilled labor 1.9% were whereas in control group most of respondents were self employed 45.7%, private employee 26.7%, Govt. employee 13.3%, unemployed 8.6% and unskilled labor 5.7%. In experimental group majority of respondents belonged to joint family 56.2%, extended family 33.3%, and nuclear family 10.5% whereas in control group majority of respondents belonged to joint family 47.6%, extended family 41%, and nuclear family 11.4%. In experimental group majority of respondents i.e.; 35.2% had 10001-15000 Inr monthly income, 27.6% had 15001-20000 Inr monthly income, 25.7% had more than 20000 Inr monthly income and only 5.7% respondent had 5001-10000 Inr and 8.8% respondents had less than 5000 Inr monthly income whereas in control group majority of respondent i.e.; 39% had more than 20000 Inr monthly income, 27.6% had 15001-20000, 17.1% had 10001-15000 Inr monthly income, 14.3% had 5001-10000 inr monthly income and 1.9% respondent had less than 5000 INR monthly income.

In experimental group majority of respondent 61.9% were patient self, 12.4% were parents, 10.5% were siblings, 8.6% were child and 6.7% respondent were any other whereas in control group most of respondent 64.8% were patient self, 12.4% were child, 9.5% were any other and 6.7% respondent were parents and siblings. In experimental group majority of respondent were from Palliative unit 92.4%, ART Centre 7.6% whereas in control group majority of respondent were from Palliative unit 91.4%, ART Centre 8.6%. in experimental group most of respondent 59% were taking treatment less than 1 Yr., 30.5% were taking treatment from 1 to 2 Yrs. and 10.5% respondent were taking treatment more than 2 Yrs. whereas in control group most of respondent 53.3% were taking treatment less than 1Yr, 26.7% were taking treatment from 1 to 2 Yrs., and only 20% respondent were taking treatment more than 2 Years. in experiment group most of respondents i.e.; 31.4% had renal failure followed by 23.8% had liver cirrhosis,7.6% had cancer and road accident, 6.7% had dementia and HIV, 5.7% had stroke, 4.8% had Parkinson's disease, 3.8% had paralysis, and 1.9% had brain tumor whereas in control group most of respondents i.e.; 30.5% had renal failure, 21.9% had liver cirrhosis, 11.4% had stroke, 7.6% had HIV, 6.7% had road accident, 5.7% had paralysis, 4.8% had dementia and Parkinson's disease, 3.8% had cancer and 2.9% had brain tumor. In experimental group majority of respondent were partially dependent 72.4%, independent 21%, completely dependent 6.7% whereas in control group majority of respondent were partially dependent 67.6%, completely dependent 21%, independent 11.4%.

# **SECTION-2:** Pre-test assessment of pain among terminally ill patient. (Table 2 and Figure 1)

Table-2 and figure-1 described the level of pain before application of Guided imagery technique in experimental group most of respondents i.e.: 41% had severe pain, 40% had moderate pain, 16.2% had mild pain and only 2.9% had no pain whereas in control group most of respondents i.e.: 38.1% had severe pain, 35.2% had moderate pain, 23.8% had mild pain and only 2.9% had no pain. It showed that

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	rable 2: Pre-test assessment of pain among terminally in patient.							
LEVEL OF PAIN		EXPERIME	NTAL GROUP	CONTR	OL GROUP			
	PAIN SCORE	Frequency (f)	Percentage (%)	Frequency (f)	Percentage (%)			
NO PAIN	0	03	2.9	03	2.9			
MILD PAIN	1-2	17	16.2	25	23.8			
MODERATE PAIN	3-6	42	40.0	37	35.2			
SEVER PAIN	7-10	43	41.0	40	38.1			
TOTAL		105	100.0	105	100.0			

 Table 2: Pre-test assessment of pain among terminally ill patient.

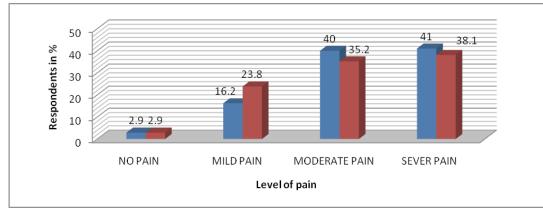


Figure 1: Pre-Test Level of Pain among Terminally III Patients.

LEVEL OF	PAIN			Experime	ntal Group					Contro	ol group		
PAIN	SCORE	Pre	-test	Pos	t-test	Follo	w up	Pre	-test	Pos	t-test	Folle	ow up
		Frequency (f)	Percentage (%)										
NO PAIN	0	03	2.9	07	6.7	18	17.2	03	2.9	05	4.8	05	4.8
MILD PAIN	1-2	17	16.2	33	31.4	54	51.4	25	23.8	28	26.7	35	33.3
MODERATE PAIN	3-6	42	40	62	59	33	31.4	37	35.2	57	54.3	61	58.1
SEVERE PAIN	7-10	43	41	03	2.9	0	0	40	38.1	15	14.3	04	3.8
TOTAL		105	100.0	105	100.0	105	105	105	100.0	105	100.0	105	100

there was a significant relationship between pain score with terminally ill patients at selected hospitals, Jaipur. Hence, null hypothesis  $H_01$  was rejected.

**SECTION-3:** Effectiveness of guided imagery technique on pain. (Table 3 and Figure 2)

Table-3 and figure-2 described the level of pain among terminally patients. In experimental group pre-test 41% respondent had ill severe pain, followed by 40% respondent had moderate pain, 16.2% respondent had mild pain and 2.9% had no pain whereas in post- test 59% respondent had moderate pain, 31.4% respondent had mild pain, 6.7% respondent had no pain and 2.9% respondent had severe pain, in follow-up 51.4% respondent had mild pain, 31.4% respondent had moderate pain and 17.2% respondent had no pain,. In control group pre-test 38.1% respondent had severe pain, 35.2% respondent had moderate pain, 23.8% respondent had mild pain and 2.9% had no pain whereas in post-test 54.3% respondent had moderate pain, 26.7% respondent had mild pain 14.3% respondent had severe pain and 4.8% had no pain, and in follow-up 58.1% respondent had moderate pain, 33.3% respondent had mild pain, 4.8% respondent had no pain, and 3.8% had severe pain. It showed significant relationship between effect of guided imagery technique and score obtained on level of pain. Hence, null hypothesis H<sub>0</sub>2 was rejected.

# Table-4: Effectiveness of Level of Pain among Terminally IllPatients. (Table 4 and Figure 3)

Table-4 and figure-3 revealed that the mean value of pre-test, post-test, follow-up test level of pain among terminally ill patients in experimental group and control group. Pre-test mean score in experimental group was 5.68, whereas in control group mean score was 5.25, t= -1.16, p=0.24.Post-test mean score in experimental group was 2.98, whereas in control group mean score was 4.93, t=6.8, p=<.01, Significant. Follow-up in experimental group mean score was 1.94, whereas in control group mean score was 4.4, t=9.7, p=<.01, Significant.

# **SECTION-4:** Association between pre-test score of pain, with selected socio-demographic variables. (Table 5)

Table-5 revealed that the calculated chi square value of Age  $\chi 2=$  25.89, marital status  $\chi 2=$ 21.4, area of hospitalization  $\chi 2=$ 7.6, duration of taking treatment  $\chi 2=$ 12.84, level of dependency  $\chi 2=$ 15.08 was found significant at level of 0.05 and 0.01. So, there was significant association between experimental group level of pain. Hence, null hypothesis H<sub>0</sub>3 was rejected for above socio-demographic variable. (Table 6)

Table-6 described that the calculated chi square value for, marital status=26.14, educational qualification  $\chi$ 2=27.91 family type  $\chi$ 2=21.63,

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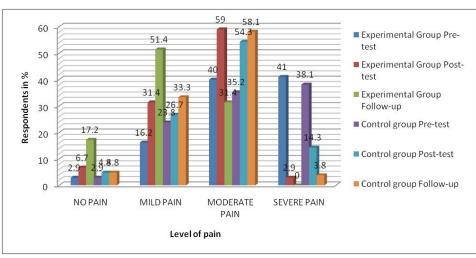


Figure 2: Level of Pain among Terminally III Patients.

Test	Group	Mean	Standard Deviation	Mean Difference	Standard Error	t -Value	P Value
Pre-Test	Experimental Group	5.68	2.53	0.43	0.37	1.16	0.24
	Control Group	5.25	2.85				
Post-Test	Experimental Group	2.98	1.68	1.95	0.28	6.8	<.01
	Control Group	4.93	2.39				
Follow-Up	Experimental Group	1.94	1.36	2.46	0.25	9.7	<.01
	Control Group	4.4	2.22				

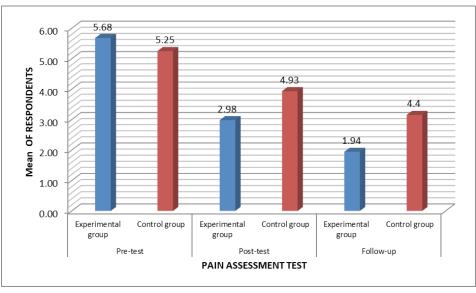


Figure 3: Effectiveness of Level of Pain among Terminally III Patients.

area of hospitalization  $\chi 2=14.41$ , duration of taking  $\chi 2=14.02$  level of dependency  $\chi 2=15.09$ . So, there was significant association between control group levels of pain. Hence, the null hypothesis H03 was rejected for above socio-demographic variable.

## Discussion

The aim of study to evaluate the effectiveness of guided imagery technique regarding pain, among terminally ill patients. In pre-test in experimental group most of respondents i.e.: 41% had severe pain,40% had moderate pain, 16.2% had mild pain and only 2.9% had no pain

whereas in control group most of respondents i.e.: 38.1% had severe pain, 35.2% had moderate pain, 23.8% had mild pain and only 2.9% had no pain. It showed there was a significant relationship between pain score with terminally ill patients at selected hospitals, Jaipur, hence null hypothesis H01 was rejected. A similar study was conducted on "Prevalence of Pain in terminally Ill Cancer Patients: A Prospective Nonrandomized Observational Study" at Shri Mahant Indiresh Hospital, Dehradun, Uttarakhand, India. 126 patients incorporated in the study. The study revealed that majority of respondents had moderate pain 75.40% followed by severe pain 62.70%, chronic pain

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SL NO	DEMOGRAFIC CHARACTERSTICS	CHI-SQUIRE	d.f	CALCULATED "P" VALUE	INFERANCE
1	AGE	25.89	12	.01	Significant
2	GENDER	1.3	6	.97	Not significant
3	RELIGION	13.60	9	.13	Not significant
4	MARITAL STATUS	21.4	9	.01	Significant
5	HABITAT	11.84	6	.06	Not significant
6	EDUCATION QUALIFICATION	18.71	12	.09	Not significant
7	OCCUPATION	20.2	12	.07	Not significant
8	FAMILY TYPE	10.82	6	.09	Not Significant
9	MONTHLY FAMILY INCOME	15.25	12	.22	Not Significant
10	SOURCE OF MAJOR INCOME	19.2	12	.08	Not Significant
11	AREA OF HOSPITALIZATION	7.6	3	.05	Significant
12	DURATION OF TAKING TREATMENT	12.84	6	.04	Significant
13	LEVEL OF DEPENDENCY	15.08	6	.01	Significant

Table 5: Distribution of Experimental Group Respondents by Pre-Test Level of Pain and Socio-Demographic Variable.

Table 6: Distribution of Control Group Respondents by Pre-Test Level of Pain and Socio-Demographic Variable.

SL NO	DEMOGRAFIC CHARACTERSTICS	CHI-SQUIRE	d.f	CALCULATED "P" VALUE	INFERANCE
1	AGE	7.6	12	.80	Non- Significant
2	GENDER	2.89	6	.82	Non- Significant
3	RELIGION	16	9	.06	Non- Significant
4	MARITAL STATUS	26.14	9	< .01	Significant
5	HABITAT	10.55	6	.11	Non- Significant
6	EDUCATION QUALIFICATION	27.91	12	< .01	Significant
7	OCCUPATION	17.2	12	.14	Non- Significant
8	FAMILY TYPE	21.63	6	< .01	Significant
9	MONTHLY FAMILY INCOME	14.5	12	.26	Non- Significant
10	SOURCE OF MAJOR INCOME	10.36	12	.58	Non- Significant
11	AREA OF HOSPITALIZATION	14.41	3	< .01	Significant
12	DURATION OF TAKING TREATMENT	14.02	6	.02	Significant
13	LEVEL OF DEPENDENCY	15.09	6	.01	Significant

26.98%, neuropathic pain 13.49%. The average duration of pain was 171.16  $\pm$  716.50 days. Totally 46.03% and 42.01% patients had at least one and more than equal to 2 neuropathic pain symptoms, respectively [7].

In experimental group pre-test 41% had severe pain, 40% had moderate pain, 16.2% had mild pain and 2.9% had no pain whereas in posttest only 2.9% had severe pain, 59% had moderate pain, 31.4% had mild pain and 6.7% had no pain. In control group pre-test 38.1% had severe pain, 35.2% had moderate pain, 23.8% had mild pain and only 2.9% had no pain whereas in post-test 14.3% had severe pain, 54.3% had moderate pain, 26.7% had mild pain, and only 4.8% had no pain. It showed significant relationship between effect of guided imagery technique and score obtained on level of pain, hence, null hypothesis H02 was rejected. In experimental group pre-test, follow-up mean difference was -3.7 and t= 13.34, p=<.01, Significant, whereas in control group pre-test- follow-up mean difference was -.85 and t=2.4, p=<.05, Significant. Pre-test mean score in experimental group was 5.68, whereas in control group mean score was 5.25, t= 1.16 p=0.24 non-significant .Post-test mean score in experimental group was 2.98, whereas in control group mean score was 4.93, t=6.8 p=<.01, Significant. Follow-up in experimental group mean score was 1.94, whereas in control group mean score was 4.4, t=9.7, p=<.01, Significant. Hence, the result revealed that guided imagery technique was more effective to reduce the pain in experimental group as compared to control group.

A similar study was conducted on "Effect of Guided Imagery Technique on Pain Intensity in Post Sectio Caesarea Patients" at Permata Bunda Hospital. The study showed before guided imagery techniques data were obtained by 71.4% respondents on the medium pain scale and after intervention 71.4% were on the light pain scale. In the control group, the first measurement obtained pain intensity in 85.8% respondents were on the moderate pain scale and the 2nd measurement of pain intensity were 71.4% increased on the severe pain scale. There was a difference in the intensity of pain in the decreased intervention group reaching 2 scales and the control group experienced a mean 1 increase in scale. After t test dependent test obtained p = 0.001significant.. Researcher concluded thar guided imagery techniques had a significant effect on the pain levels of post-operative patients of sectio Caesarea [8].

In the present study experimental group was significant association between level of pain score and selected socio-demographic variable such as age  $\chi 2= 25.89$ , marital status  $\chi 2=21.4$ , area of hospitalization  $\chi 2= 7.6$ , duration of taking treatment  $\chi 2= 12.84$  and level of dependency  $\chi 2= 15.08$ . In control group was significant association between level of pain score and selected socio-demographic variables such as marital status  $\chi 2=26.14$ , educational qualification  $\chi 2= 27.91$ , family type  $\chi 2=$ 21.63, area of hospitalization  $\chi 2= 14.41$ , duration of taking treatment  $\chi 2= 14.02$  and level of dependency  $\chi 2= 15.09$ . Hence, null hypothesis H03 rejected.

A study was carried out on "Pain level associated to sociodemographic and clinical variables in people living with human immunodeficiency virus and acquired immunodeficiency Syndrome" at Brazil. The data revealed significant relationship between pain and age p=0.02, the age groups between 21 and 30 years p<0.046, 41 and 50 years p<0.023 and 51 and 60 years p<0.030 were protection factors for

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the presence of moderate pain [9].

## Conclusion

The study concluded that the guided imagery technique was found effective to reduce pain. In pain experimental group pre-test mean  $\pm$  S.D was 5.68  $\pm$  2.53, in follow up mean  $\pm$  S.D was 1.94  $\pm$ 1.36, mean difference was -3.7, t= 13.34, p=<.01, Significant, whereas in control group pre-test mean  $\pm$  S.D was 5.28  $\pm$  2.85, in follow up mean  $\pm$  S.D was 4.40  $\pm$ 2.22, mean difference was -.8, t=2.4, p=<.05, Significant. Hence, the result revealed that guided imagery technique was more effective to reduce the pain in experimental group as compared to control group.

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# **Conflicts of interest**

There are no conflicts of interest

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