

Impact of Nursing Educational Program on Health Promotion for Patients after Intracranial Surgery

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Abstract

Intracranial surgery means surgery performed inside skull to treat problems in brain and surrounding structures. **This study aims** to: determine physical, social and emotional problems of patients after intracranial surgery, identify needs of patients after intracranial surgery and develop and implement nursing educational program based on patient's needs. **Hypotheses:** Studied participants patients post application of nursing educational program will exhibit more knowledge, health improvement and fewer complications. **Subjects and methods:** Sixty adult patients after intracranial surgery (burr hole, craniotomy and craniectomy). Their age 18-65 years old, fully conscious, both sexes. Patients divided randomly into two groups; control group (30 patients received routine care) and study group (30 patients received nursing educational program). Two tools were used for data collection; patient assessment sheet and patient evaluation sheet. **Research design:** Quasi- experimental research design. **Results:** Good improvement in total knowledge scores of study group patients after application of program. Majority of patients in study group showed health improvement while less than half of patients in control group showed health improvement. **Conclusion:** Improving patients' knowledge in study group had a favorable effect in improving their health than in control group. **Recommendation:** Establishment of educational center in neurosurgery department to educate patients about their conditions.

Key words: *Intracranial Surgery & Nursing Educational Program.*

Introduction

The neurosurgical cranial procedures are relatively frequent in the daily practice and a wide range of neurosurgical techniques has been developed (burr holes, craniotomy and craniectomy) to treat patients with intracranial disorders. The entire treatment plan drains the patient physically, mentally and emotionally. It is generally the weakest and the most vulnerable phase of a patient's life (Guilbert, 2014).

Surgical burr hole is a hole made in the skull by a special drill to gain direct access of the brain (Wilson et al., 2013). A craniotomy is a surgery during which a piece of the skull (bone flap) is removed in order to allow direct access to the brain. After the surgery is performed, the bone flap is returned to its previous location. Craniectomy procedure includes removal of part of the skull bone (Heisler, 2014).

Complications after intracranial surgery may include intracranial bleeding, brain abscess, further neurological impairment, behavioral changes, infection, and seizures. Hemorrhage at the operative site can occur within hours after surgery. Bleeding may occur in the subdural or subarachnoid space or within the ventricles (Yarbro et al., 2011).

The patient needs comfort and reassurance at all times after intracranial surgery. Despite normal feelings of anxiety and fear, the caregivers should act

strong and provide support with proper guidance and accurate information from the medical team. Postoperative care after intracranial surgery might not end soon in some cases; it could last for months or even for years. The caregiver should be mentally prepared for this fact and not spend all their energy right at the beginning. Also, clarify with the doctor the nature of symptoms; those which are a natural part of the recuperation process and the ones which would signal an emergency (Suzanne et al., 2010).

Because of nurses act as a vital members of the health care team and considered as a vital component in the overall patient outcome based on the expert neurological assessment, they are involved to care for intracranial surgery patient at various levels of intervention and health education. The nurse must have a thorough understanding of patients' needs to provide optimal nursing intervention and education and thus improve patients' health (McGlinsey & Kirk, 2014).

Patient education is defined as any set of planned educational activities designed to improve patients' health behaviors and health status. Its main purpose is to maintain or to improve patient health or, in some cases, to slow deterioration. An informed and educated patient can actively participate in his or her own treatment, improve outcomes, help identify

errors before they occur, and reduce his or her length of stay. Medical component of health education involves medical information and preventative measures concerning health and well-being. Research has demonstrated that effective health education begins with identification of various important needs for the patients (Haddad et al., 2013). Health promotion is the science and art of helping people change their lifestyle to move toward a state of optimal health. Optimal health is defined as a balance of physical, emotional, social, spiritual and intellectual health. Lifestyle change can be facilitated through a combination of efforts to enhance awareness, change behavior and create environments that support good health practices (World Health Organization, 2012).

According to hospital records at Assiut University Hospital during the period of one year (1/1/2013-1/1/2014), it has been found that 998 patients admitted to the neurosurgery department and perform intracranial surgery (Assiut University Hospital Record, 2013-2014). Not present international incidence related to intracranial surgery but there is incidence for the diseases that require intracranial surgery.

Significance of the study

Patients after intracranial surgery have physical, social, and emotional problems that affect greatly their life. Those patients are needed for special nursing care and health teaching to improve their condition. So, this study will be the first study in this geographical location which will help those patients improve their condition

Aim of the study

This study aimed to:

1. Determine the physical, social and emotional problems of patients after intracranial surgery.
2. Identify the needs of patients after intracranial surgery.
3. Develop and implement the nursing educational program based on patient's needs.

Research hypotheses

To fulfill the aim of this study, the following research hypotheses were formulated:

1. The knowledge of studied patients after application of the nursing educational program will be higher than their knowledge before the application of it.
2. Studied participants patients post application of nursing educational program will exhibit higher health improvement than control group.
3. Studied participants patients post application of nursing educational program will exhibit fewer

symptoms or complications; physical, social or emotional problems than control group.

Subjects and methods

Research design:

Quasi experimental research design was utilized to fulfill the aims of this study.

Technical design

Setting

This study was conducted in neurosurgery department and neurosurgery out patient clinic at Assiut University Hospital.

Subjects

A random sample was obtained; sixty adult patients after intracranial surgery (burr hole, craniotomy and craniectomy) were included in this study. They had the following criteria; the age ranged from 18-65 years old, fully conscious, both males and females. The patients divided randomly into two groups: the control group (30 patients who received routine care) and the study group (30 patients who received nursing educational program). The number of males include: (18 in control and 17 in study groups) while the number of females include: (12 in control and 13 in study groups). Data were collected during the period from (December 2012 to June 2014); this period included both the time of assessment and follow up. Patients were assessed immediately after intracranial surgery (burr hole, craniotomy and craniectomy) and daily during hospitalization. Those patients were followed up before discharge in the neurosurgery department, 3 months and 6 months after intracranial surgery in the neurosurgery out patient clinic. Their ages ranged from 18 to 65 years with a mean value of (43.27±6.38) for control and (41.25±5.83) for study groups.

Exclusion criteria

- Disoriented patient.
- Uncooperative patient.
- Patients on mechanical ventilation.

Tools

Tool I: Postoperative intracranial surgery patient assessment sheet: This tool consisted of four parts:

Part 1: Socio-demographic characteristics of patients.

Part 2: Patients' nursing needs: This included structured items to identify patients' nursing needs; it is used to assess:

1. Medical history.
2. Neurological physical examination (level of consciousness by Glasgow coma scale (GCS) (Teasdale and Jennett, 1974), cranial nerves, motor and sensory function, sphincter control and vital signs).

3. Psychological problems (social and emotional problems) such as anxiety, fear, depression or social isolation, sensitivity to noise or people in crowded places, apathy or crying spells.
4. Laboratory investigation and diagnostic procedures. It includes certain investigation such as hemoglobin level, red and white blood cells count, platelets count, blood sugar level, kidney function (urea and creatinine), prothrombin time and concentration and/or serum electrolytes (sodium, potassium). Certain procedures such as computed tomography, magnetic resonance imaging and cerebrospinal fluid analysis.

Part 3: Systemic and/or neurosurgical postoperative complications that may develop for patients after intracranial surgery. Systemic complications (cardiovascular, gastrointestinal, metabolic and wound infection). Neurosurgical complications (behavioral changes, cerebrospinal fluid leak, postoperative hematoma, seizures, residual neurological problems, visual disturbances brain abscess and insomnia)

Part 4: Patients' knowledge regarding intracranial surgery: It included 39 questions concerning types of intracranial surgery, treatment, postoperative complications, postoperative care, follow up and health promotion after intracranial surgery.

Nursing educational program after intracranial surgery: Developed by researcher based on patient's assessment needs after reviewing current national and international literature to maintain health promotion for patients and reduce or prevent postoperative complications. It includes:

- Brief anatomy of the brain.
- Definition and indication for intracranial surgery.
- Various diagnostic procedures and how to be prepared for it.
- Benefits of surgical management and types of intracranial surgery.
- Postoperative complications related to anesthesia and surgery.
- Physical, social and emotional problems that the patient may complain after intracranial surgery.

- Information about how to promote health through:
- Medical therapy after intracranial surgery.
- How to deal with seizures.
- Care of wound site.
- Routine follow up and when it is necessary to seek medical help and immediately go to the hospital
- Life style modification:
 - Nutrition.
 - Weight control.
 - Rest.
 - Physical activity and exercises.
 - Smoking cessation.
 - Stress reduction.
 - Effective communication.
 - Control of diabetes mellitus and hypertension.

Tool II: Patient evaluation sheet: This tool consisted of three parts:

Part 1: Patients' knowledge regarding intracranial surgery, it included 39 questions concerning types of intracranial surgery, treatment, postoperative complications, postoperative care, follow up and health promotion after intracranial surgery.

Part 2: Rand short form (SF) 36 items questionnaire. It contains 36 questions and measures health status. It divided into eight scales plus one health comparison question (health change).

1. Physical functioning.
 2. Role limitations due to physical problems.
 3. Role limitations due to emotional problems.
 4. Vitality (Energy / fatigue).
 5. Mental health (Emotional well being).
 6. Social functioning.
 7. Pain.
 8. General health.
- Health comparison question (Health change).

Scoring system for Rand short form 36 questionnaire (Ware and Sherbourne, 1992): Each scale contains from 2-10 items. All questions are scored on a scale from 0 to 100, with 100 representing the highest level.

How to score Rand short form-36 questionnaire

Step 1: Scoring questions

Scoring system		
Question number	Original response	Recorded value
1, 2, 20, 22, 34, 36	1	100
	2	75
	3	50
	4	25
	5	0

Scoring system		
Question number	Original response	Recorded value
3, 4, 5, 6, 7, 8, 9, 10, 11, 12	1	0
	2	50
	3	100
Scoring system		
Question number	Original response	Recorded value
13, 14, 15, 16, 17, 18, 19,	1	0
	2	100
Scoring system		
Question number	Original response	Recorded value
21	1	100
	2	80
	3	60
	4	40
	5	20
Scoring system		
Question number	Original response	Recorded value
23, 26, 27, 30	1	100
	2	80
	3	60
	4	40
	5	20
	6	0
Scoring system		
Question number	Original response	Recorded value
24, 25, 28, 29, 31	1	0
	2	20
	3	40
	4	60
	5	80
	6	100
Scoring system		
Question number	Original response	Recorded value
32, 33, 35	1	0
	2	25
	3	50
	4	75
	5	100

Step 2: Average items to form 8 scales

Scale	Number of items	Average the following items (Question numbers)
Physical functioning; Limitations in physical activity because of health problems	10	3, 4, 5, 6, 7, 8, 9, 10, 11, 12
Role limitations – physical; Limitations in usual role activities because of physical health problem	4	13, 14, 15, 16
Role limitations – emotional; Limitations in usual role activities because of emotional problems.	3	17, 18, 19
Vitality; Energy and fatigue	4	23, 27, 29, 31

Scale	Number of items	Average the following items (Question numbers)
Mental health; Psychological distress and well-being.	5	24, 25, 26, 28, 30
Social functioning; Limitations in social activities because of physical or emotional problems	2	20, 32
Pain; Presence of pain and limitations due to pain	2	21, 22
General health perception	5	1, 33, 34, 35, 36
Health comparison question; Health change	1	2

Step 3: Figuring scores

Rand recommends the following straightforward approach to scoring the Rand 36-items health survey. All questions are scored on a scale from 0 to 100, with 100 representing the highest level.

Aggregate scores are compiled as a percentage of the total points possible, using the Rand scoring table (step 1).

The scores from those questions that address each specific area of functional health status (step 2) are then averaged together, for a final score within each of the 8 scales measured (e.g. pain, physical functioning etc.)

For example, to measure the patient's energy/fatigue level, add the scores from questions 23, 27, 29, and 31. If a patient circled 4 on 23, 3 on 27, 3 on 29 and 1 on 31, use (Step 1) to score them.

An answer of 4 to question 23 is scored as 40, 3 to question 27 is scored as 60, 3 to question 29 is scored as 40 and 1 to question 31 is scored as 0. The score for this block is $40+60+40+0=140$. Now we divide by the 4 answered questions to get a total of 46.7. Since a score of 100 represents high energy with no fatigue, the lower score of 46.7% suggests the patient is experiencing a loss of energy and is experiencing some fatigue. All 8 scales are scored in the same way.

Part 3: Systemic and/or neurosurgical postoperative complications that may develop for patients after intracranial surgery. Systemic complications (cardiovascular, gastrointestinal, metabolic and wound infection). Neurosurgical complications (behavioral changes, cerebrospinal fluid leak, postoperative hematoma, seizures, residual neurological problems, visual disturbances brain abscess and insomnia)

II. Operational design

Procedure:

Phase I: Preparatory phase: An official permission to proceed with the proposed study was obtained from the head of the neurosurgery department. The researcher designed and tested health promotion program after reviewing extensive literature.

Content validity and reliability

Content validity was done by five expertise from the medical staff and medical –surgical nursing staff. Two expertise from medical–surgical nursing staff

and three expertise from neurosurgery staff who reviewed the tools and the nursing educational program for clarity, relevance, comprehensiveness, understanding, applicability and easiness for administration. The content is valid and reliable. Reliability was assessed by correlation coefficient with >0.8 considered strong.

Pilot study:

It was conducted on 10% of sample (6 patients) in a selected setting for testing clarity and applicability and feasibility of the study tools. The purpose of the pilot study was to detect any particular problem in the statements clarity, feasibility, and applicability of the tool. The data obtained from the pilot study were analyzed, no change was done in the assessment sheet, so the 10% of subjects selected for the pilot study were included in the main study.

Phase II: implementation phase: At initial interview the researcher introduce herself to initiate communication, explain the nature and purpose of the study.

The study group visited daily during hospitalization to fill out the patient assessment sheet (tool I) to assess patients` knowledge and needs and implement nursing educational program.

Regarding control group, they received the routine postoperative hospital care. They visited daily during hospitalization to fill out (tool I).

The nursing educational program was introduced to patients through individualized sessions. 10 educational sessions were conducted for each patient. The duration of each session was 30 – 45 minutes, including 10 minutes for discussion and feedback. Each session usually started by a summary of what has been taught during the previous sessions and the objectives of the new topics. Feedback and reinforcement of teaching was performed according to the patients` needs to ensure their understanding. Each session ended by a summary of its contents and feedback of the patient through discussion and questions.

Phase III: Evaluation phase: After application of the nursing educational program, the patients` knowledge and condition have been evaluated by the researcher through filling the patient evaluation sheet (tool II). Follow up has been done for patients in

study and control groups for a period of 6 months on three phases (before discharge, 3 months and 6 months after intracranial surgery).

Ethical considerations

Informed consent was obtained from patients who are willing to participate in the study after explanation of the nature and purposes of the study. Patients' privacy and confidentiality were certainly assured.

Statistical design

Data were collected and analyzed by computer program SPSS (version 17). Data expressed as mean, standard deviation, number and percentage. T-test is used to determine significant for numeric variable. Chi-square test is used to determine significant for

non-parametric variable.

A probability level of <0.05 was adopted as a level of significance for testing the research hypotheses.

Limitations of the study

1. The patient's anxiety may interfere with the ability to learn information provided.
2. During preparation of a teaching booklet, the level of literacy in the patient population was considered. A lack of reading skills limited the ability of patients to access and use critical information.
3. Participants suffered from transportation and financial problems. So, some patients interview after 6 months by telephone.

Results

Table (1): Distribution of socio-demographic characteristics of patients (control and study groups).

Socio-demographic characteristics	Control (n =30)			Study (n =30)		
	No.	%	X + SD	No.	%	X + SD
Age						
18< 35 yrs	1	3.3		3	10	
35<50yrs	19	63.3	43.27±6.38	17	56.7	41.25±5.83
50-65yrs	10	33.3		10	33.3	
Sex						
Male	18	60	-	17	56.7	-
Female	12	40		13	43.3	
Marital status						
Single	11	36.7		4	13.3	
Married	19	63.3	-	26	86.7	-
Level of education						
High education	2	6.7		2	6.7	
Secondary school education	6	20	-	8	26.7	-
Read and write	5	16.7		6	20	
Illiterate	17	56.7		14	46.7	
Occupation						
Office work	3	10		5	16.7	
Machinery work	6	20		2	6.7	
Manual work	14	46.7	-	11	36.7	-
Housewife	6	20		9	30.0	
Not working	1	3.3		3	10	

Table (2): Distribution of patients' medical history (control and study groups).

Patients' medical history	Control (n =30)		Study (n =30)	
	No.	%	No.	%
Surgical management				
Burr holes	8	26.7	12	40.0
Craniotomy	18	60.0	13	43.33
Craniectomy	4	13.33	5	16.7

Table (3): Distribution of patients (study and control groups) as regard level of consciousness (GCS).

GCS	Control (n =30)			Study (n =30)		
	No.	%	X ± SD	No.	%	X ± SD
Mild	30	100	15.00±15.00	30	100	15.00±15.00
Moderate	-	-		-	-	
Severe	-	-		-	-	

Table (4): Distribution of patients (study and control groups) as regard cranial nerves.

Cranial nerves	Control (n =30)				Study (n =30)				P-value
	Normal		Abnormal		Normal		Abnormal		
	No.	%	No.	%	No.	%	No.	%	
I- Olfactory	30	100	-	-	30	100	-	-	P=0.694 n.s
II- Optic	27	90	3	10	28	93.3	2	6.7	
III- Oculomotor	30	100	-	-	30	100	-	-	
IV- Trochlear	30	100	-	-	30	100	-	-	
V- Trigeminal	30	100	-	-	30	100	-	-	
VI- Abducens	30	100	-	-	30	100	-	-	
VII- Facial	30	100	-	-	30	100	-	-	
VIII- Acoustic	29	96.7	1	3.3	30	100	-	-	
IX- Glosopharyngeal	30	100	-	-	30	100	-	-	
X- Vagus	30	100	-	-	30	100	-	-	
XI- Accessory	30	100	-	-	30	100	-	-	
XII- Hypoglossal	30	100	-	-	30	100	-	-	

N.S: Non significant ($P>0.05$).*: Significant ($P<0.05$).** : Moderate significant ($P<0.001$).***: Highly significant ($P<0.0001$).**Table (5): Distribution of patients (study and control groups) as regard sensory function.**

Sensory function	Control (n =30)				Study (n =30)				P-value
	Normal		Abnormal		Normal		Abnormal		
	No.	%	No.	%	No.	%	No.	%	
Pain and temperature sensation	30	100	-	-	30	100	-	-	P= 0.762n.s
Position sense (Proprioception)	29	96.7	1	3.3	30	100	-	-	
Light touch	28	93.3	2	6.7	29	96.7	1	3.3	

Table (6): Distribution of patients (study and control groups) as regard sphincter control.

Sphincter control	Control (n =30)				Study (n =30)				P-value
	Present		Absent		Present		Absent		
	No.	%	No.	%	No.	%	No.	%	
Bladder control	29	96.7	1	3.3	30	100	-	-	P= 0.794n.s
Bowel control	30	100	-	-	30	100	-	-	

Table (7): Distribution of patients (study and control groups) as regard motor function.

Motor function	Control (n =30)		Study (n =30)		P-value
	No.	%	No.	%	
1. Muscle size					
Normal	30	100	30	100	-
Muscle atrophy	-	-	-	-	
2. Muscle tone					
Normal	30	100	30	100	-
Spastic	-	-	-	-	
Rigid	-	-	-	-	
Flaccid	-	-	-	-	
3. Muscle power					
Grade 5: Normal power	28	93.3	30	100	P=0.859n.s
Grade 4: Active movement against gravity and resistance	2	6.7	-	-	
Grade 3: Active movement against gravity	-	-	-	-	
Grade 2: Active movement with gravity eliminated	-	-	-	-	
Grade 1: Trace of contraction	-	-	-	-	
Grade 0: Absent, no muscle contraction	-	-	-	-	
5. Involuntary movements					
Yes	2	6.7	1	3.3	P=0.793n.s
No	28	93.3	29	96.7	

Table (8): Distribution of patients (study and control groups) as regard vital signs.

Vital signs	Control (n =30) X±SD	Study (n =30) X±SD	P-value
Body temperature			
1 st day	36.86±0.13	36.83±0.19	P =0.531 n.s
2 nd day	36.94±0.08	36.93± 0.15	P = 0.425 n.s
3 rd day	37.85±0.39	36.92±0.15	P<0.01*
Pulse			
1 st day	76.53±12.00	78.63±6.50	P = 0.429 n.s
2 nd day	78.83± 11.91	80.50±5.32	P =0.372 n.s
3 rd day	77.00± 11.84	79.80± 4.55	P <0.01*
Respiration:			
1 st day	18.16±1.17	18.63±1.86	P =0.441 n.s
2 nd day	18.83±1.46	18.56±1.77	P =0.588 n.s
3 rd day	18.93±1.38	18.20±1.39	P <0.04*
Blood pressure			
	Systolic	Systolic	
1 st day	116.50±8.00	121.17±9.79	P =0.522 n.s
2 nd day	117.17±5.97	116.40± 20.25	P =0.621 n.s
3 rd day	116.67±5.77	115.57±19.66	P <0.04*
	Diastolic	Diastolic	
1 st day	80.16±8.55	81.50±6.58	P =0.502 n.s
2 nd day	80.03±4.77	81.50± 5.27	P =0.264 n.s
3 rd day	78.73±4.86	77.63±3.87	P <0.02*

Table (9): Distribution of patients (study and control groups) as regard psychological assessment (social and emotional problems).

Psychological assessment (More than one)	Control (n =30)		Study (n =30)		P-value
	No.	%	No.	%	
1. Anxiety	28	93.3	27	90.0	P =0.246 n.s
2. Fear	28	93.3	27	90.0	P =0.335 n.s
3. Depression (social isolation)	2	6.7	-	-	P =0.246 n.s
4. Sensitivity to noise	5	16.7	2	6.7	P =0.212 n.s

Table (10): Comparison between patients (study and control groups) as regard laboratory investigation.

Laboratory investigation	Control (n =30) X±SD	Study (n =30) X±SD	P-value
Hemoglobin	13.64± 1.26	13.66±1.05	P =0.962 n.s
White blood cells	7.08±2.75	6.92±2.11	P =0.791 n.s
Red blood cells	5.04±0.506	4.85±0.52	P =0.151 n.s
Platelets	239.33±53.32	248.53±55.70	P =0.516 n.s
Blood sugar	5.94±2.25	6.31±1.21	P =0.432 n.s
Urea	5.27±1.54	5.43±1.25	P =0.667 n.s
Creatinine	81.08±17.75	85.34±16.93	P =0.346 n.s
Prothrombin time	16.20±2.48	12.06±0.46	P =0.317 n.s
Prothrombin concentration	94.07±18.33	97.40±5.22	P =0.343 n.s
Sodium	144.52±4.11	139.97±3.95	P =0.117 n.s
Potassium	4.23±0.32	4.04±0.43	P =0.489 n.s

Table (11): Comparison between patients (study and control groups) as regard diagnostic procedure.

Diagnostic procedure (More than one)	Control (n =30)		Study (n =30)		P-value
	No.	%	No.	%	
Computed tomography	3	10	5	16.7	P=0.35 n.s
Magnetic resonance imaging	2	6.7	1	3.3	P=0.50 n.s
Cerebrospinal fluid analysis	1	3.3	-	-	P=0.50 n.s
Electroencephalogram	-	-	-	-	-
X-rays	-	-	-	-	-
Others	-	-	-	-	-

Table (12): Distribution of patients (study and control groups) as regard systemic postoperative complications at assessment.

Systemic postoperative complications (More than one)	Control (n=30)		Study (n =30)		P-value
	No.	%	No.	%	
1. Cardiovascular complications					P=0.273 n.s
- Hypotension	-	-	-	-	
- Hypertension	5	16.7	2	6.7	
- Bradycardia	-	-	-	-	
- Tachycardia	5	16.7	2	6.7	
- Myocardial infarction	-	-	-	-	
- Arrhythmias	-	-	-	-	
- Heart failure	-	-	-	-	
2. Gastrointestinal complications					
- Nausea	2	6.7	4	13.3	
- Vomiting	2	6.7	4	13.3	
- Gastric irritation	-	-	-	-	
- Gastric stress ulceration	-	-	-	-	
- Hemorrhage	-	-	-	-	

Table (13): Distribution of patients (study and control groups) as regard neurosurgical postoperative complications at assessment.

Neurosurgical postoperative complications (More than one)	Control (n =30)		Study (n =30)		P-value
	No.	%	No.	%	
1. Cerebral edema	-	-	-	-	P=0.338 n.s
2. Brain or nerve damage	-	-	-	-	
3. Stroke	-	-	-	-	
4. Behavioral changes	2	6.7	-	-	
5. Cerebrospinal fluid leak	1	3.3	1	3.3	
6. Postoperative subdural hematoma	2	6.7	-	-	
7. Pneumocephalus	-	-	-	-	
8. Seizures	3	10.0	2	6.7	
9. Others	-	-	-	-	

Table (14): Comparison between systemic postoperative complications of patients (study and control groups) at follow up.

Systemic postoperative complications (More than one)	Before discharge				After 3 months				After 6 months			
	Control (n =30)		Study (n =30)		Control (n =30)		Study (n =30)		Control (n =30)		Study (n =30)	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
1. Cardiovascular												
- Hypotension	-	-	-	-	-	-	-	-	-	-	-	-
- Hypertension	-	-	-	-	3	10.0	1	3.3	1	3.3	-	-
- Bradycardia	-	-	-	-	-	-	-	-	-	-	-	-
- Tachycardia	3	10.0	1	3.3	-	-	-	-	-	-	-	-
- Myocardial infarction	-	-	-	-	-	-	-	-	1	3.3	-	-
- Arrhythmias	-	-	-	-	-	-	-	-	-	-	-	-
- Heart failure	-	-	-	-	-	-	-	-	-	-	-	-
2. Gastrointestinal												
- Nausea	3	10.0	2	6.7	6	20	2	6.7	7	23.3	2	6.7
- Vomiting	3	10.0	-	-	-	-	-	-	-	-	-	-
- Gastric irritation	-	-	-	-	12	40	3	10	9	30	1	3.3
- Constipation	2	6.7	1	3.3	5	16.7	1	3.3	6	20.0	-	-
3. Metabolic												
- Hyperglycemia	-	-	-	-	-	-	-	-	1	3.3	-	-
- Hypoglycemia	-	-	-	-	-	-	-	-	-	-	-	-
- Hyponatremia	-	-	-	-	-	-	-	-	-	-	-	-
4. Wound complications												
- Wound infection	1	3.3	-	-	2	6.7	-	-	-	-	-	-
- Impaired wound healing	1	3.3	-	-	-	-	-	-	-	-	-	-
p- value	P=0.252n.s				P<0.03*				P<0.04*			

Table (15): Distribution of patients (study and control groups) as regard neurosurgical postoperative complications at follow up.

Neurosurgical postoperative complications (More than one)	Before discharge				After 3 months				After 6 months			
	Control (n =30)		Study (n =30)		Control (n =30)		Study (n =30)		Control (n =30)		Study (n =30)	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
1. Behavioral changes	4	14.3	-	-	3	10.0	-	-	3	10.0	-	-
2. Postoperative subdural hematoma	-	-	-	-	-	-	-	-	1	3.3	-	-
3. Seizures	3	10.0	1	3.3	5	16.7	2	6.7	8	26.7	-	-
4. Residual neurological problems	4	14.3	4	14.3	6	20	4	14.3	7	23.3	3	10.0
5. Visual disturbances	-	-	-	-	2	6.7	2	6.7	6	20	2	6.7
6. Brain abscess	-	-	-	-	-	-	-	-	1	3.3	-	-
7. Insomnia	4	14.3	2	6.7	3	10	1	3.3	4	14.3	1	3.3
p- value	P=0.485n.s				P=0.237n.s				P<0.01*			

Table (16): Comparison between assessment and follow up total score of patients` knowledge (study and control groups).

Patients` knowledge	Control (n =30)			Study (n =30)			P-value
	No.	%	X±SD	No.	%	X±SD	
At assessment							
Satisfactory	-	-	13.73±5.97	-	-	12.26±4.50	P =0.273 n.s
Unsatisfactory	30	100		30	100		
Before discharge							
Satisfactory	-	-	14.46±4.84	30	100	47.23±3.16	P <0.0001***
Unsatisfactory	30	100		-	-		
After 3 months							
Satisfactory	-	-	17.80±5.12	28	93.3	45.73±4.57	P <0.0001***
Unsatisfactory	30	100		2	6.7		
After 6 months							
Satisfactory	-	-	19.49±5.63	27	90.0	43.36±4.83	P <0.0001***
Unsatisfactory	30	100		3	10.0		

N.B. total score of knowledge is 50.

Table (17): Comparison between short form-36 questionnaire (8 scales plus health comparison question) of studied patients (study and control groups) after 6 months.

Scales	Control (n =30) X±SD	Study (n =30) X±SD	P-value
1. Physical functioning	72.83±8.51	89.83±10.54	P <0.02*
2. Role limitation due to physical problems	55.00±3.8	89.67±11.4	P <0.001**
3. Role limitation due to emotional problems	31.85±6.41	97.0±3.00	P <0.0001***
4. Vitality; energy /fatigue	32.70 ±9.72	53.16±11.48	P <0.04*
5. Emotional well being	48.28±15.58	66.26±9.71	P <0.03*
6. Social functioning	57.41±11.5	85.0±14.8	P <0.04*
7. Pain	60.50±3.02	88.45±10.25	P <0.0001***
8. General health perception	39.50±12.86	73.67 ±18.70	P <0.001**

Table (18): Comparison between health change (Health promotion) of studied patients (study and control groups) after 6 months.

Health change (Health promotion)	Control(n=30)		Study (n =30)		P-value
	No.	%	No.	%	
Much better now than 6 months ago (100)	6	20.0	15	50.0	P <0. 0001***
Somewhat better now than 6 months ago (75)	7	23.3	12	40.0	
About the same as 6 months ago (50)	7	23.3	2	6.67	
Somewhat worse than 6 months ago (25)	6	20.0	1	3.33	
Much worse than 6 months ago (0)	4	13.3	0	0.0	

Table (1): This table illustrates that, more than half of patients were males in both control (60.0%) and study (56.7%) groups. More than half of patients in control (63.3 %) and study (56.7%) groups were having an age ranged from (35<50) years with a mean value of (43.27±6.38) and (41.25±5.83). Regarding the patients` marital status; the majority of patients were married which include (63.3%) in control and (86.7%) in study groups. Regarding education; (20.0%) in control and (26.7%) in study groups were secondary school education and (56.7%) in control and (46.7%) in study groups were illiterate. Regarding occupation; (46.7%) of patients in control and (36.7%) in study groups were manual work.

Table (2): Illustrates that, more than half of patients in control (60.0%) and less than half of patients in study (43.33%) groups performed craniotomy.

Table (3): GCS of all patients in both study and control groups presented with a mean value of (15.00±15.00).

Table (4): There was non significant difference between study and control groups as regard cranial nerves assessment.

Table (5): There was non significant difference between study and control groups as regard sensory function assessment.

Table (6): There was non significant difference between study and control groups as regard sphincter control assessment.

Table (7): There was non significant difference between study and control groups as regard motor function assessment.

Table (8): Shows the difference between patients` vital signs recording chart for both study and control group. There were significant differences during the 3rd day postoperative in vital signs including temperature, pulse, respiration and blood pressure [systolic/diastolic] (P<0.01, P<0.01, P<0.04, [P<0.04, P <0.02]) respectively. A mean value for control group (37.85±0.39, 77.00± 11.84, 77.00± 11.84, [116.67±5.77, 78.73±4.86]) respectively. A mean value for study group (36.92±0.15, 79.80± 4.55, 18.20±1.39, [115.57±19.66, 77.63±3.87]) respectively.

Table (9): shows that, the majority of patients in both control (93.3%) and study (90.0%) groups were having anxiety and fear.

Table (10): There was non significant difference between study and control groups as regard laboratory investigation.

Table (11): There was non significant difference between study and control groups as regard diagnostic procedure.

Table (12): There was non significant difference (P=0.273) between study and control groups as regard systemic complications at assessment.

Table (13): There was non significant difference (P=0.338) between study and control groups as regard neurosurgical complications at assessment.

Table (14): There was non significant difference (P=0.252) between study and control groups as regard systemic complications before discharge. After 3 and 6 months of follow up there was significant difference regarding systemic complications (P <0.03, P <0.04) respectively.

Table (15): There was non significant difference between study and control groups as regard neurosurgical complications before discharge and after 3 months of follow up (P=0.485, P=0.237). There was significant difference (P <0.01) after 6 months of follow up regarding neurosurgical complications.

Table (16): This table shows that, all patients (study and control groups) had unsatisfactory level of knowledge at the time of assessment and had non significant difference (P =0.273). At the time of follow up, there was high significant difference (P <0.0001) between patients` knowledge (study and control groups) in relation to total knowledge score.

Table (17): This table illustrates that, there was high significant difference (P <0.0001) between study and control groups as regard role limitation due to emotional problems and pain. There was moderate significant difference (P <0.001) between study and control groups as regard role limitation due to physical health problems and general health perception. There was significant difference between study and control groups as regard physical

functioning, vitality, emotional well being and social functioning ($P < 0.02$, $P < 0.04$, $P < 0.03$, $P < 0.04$) respectively.

Table (18): Illustrates that, there was high significant difference ($P < 0.0001$) between study and control groups as regard health change. The majority of patients (90 %) in study group were having health improvement while less than half of patients in control group were having health improvement.

Discussion

The present study aimed to:

1. Determine the physical, social and emotional problems of patients after intracranial surgery.
2. Identify the needs of patients after intracranial surgery.
3. Develop and implement the nursing educational program based on patient's needs.

One of the most critical surgeries performed on the human body is intracranial surgery. The entire treatment plan affects the patient physically, mentally and emotionally. To accomplish quality care and best possible outcome after intracranial surgery; nurses should be knowledgeable of the type of surgery planned, its course and possible complications (Guilabert, 2014).

The results of the present study showed that more than half of patients in both control and study groups were males their age ranged from 35<50 years old and the majority of them were married and employ.

In this regard this result agree with the study of (Bin-Madhi, 2012) entitled as "Brain tumors excision guided by neuronavigation: Practical application and results " which revealed that intracranial surgery is more common in males than females with a mean age of 47 years old.

The results in the present study revealed that patients' GCS=15 at the time of assessment after intracranial surgery indicate good prognosis for both study and control groups. Most of patients after intracranial surgery are clinically improved because pressure of the tumor, haematoma, abscess or cyst is relived.

In the same line this result agree with the study of (Abbass et al., 2007), entitled as (Glasgow coma scale on admission is correlated with postoperative Glasgow outcome scale in chronic subdural hematoma) which revealed that their is positive correlation between admission GCS and GCS in chronic subdural hematoma.

Through researcher's assessment, found that blood sample may be drawn after intracranial surgery to determine the level of red blood cells, white blood cells, platelets, hemoglobin, concentration of sodium and potassium because certain changes may be occurred after intracranial surgery. Also, certain

procedures may be performed to evaluate patients' conditions and this is performed according to their clinical presentation.

The results in the present study showed that there were non significant differences at the time of assessment between patients and their neurological physical assessment after intracranial surgery such as cranial nerves, motor function, sensory function, sphincter control. There were non significant differences between patients and their laboratory investigation and diagnostic procedures after intracranial surgery.

In the same line (Brem, 2014) stated that frequent neurological checks will be performed by the nursing and medical staff to test the brain function and to make sure the body systems are functioning properly after surgery. Patients will be asked to follow a variety of basic commands, such as moving arms and legs, to assess brain function. Pupils will be checked with a flash light, and patients will be asked questions to assess their orientation (such as name, date, and place). The strength of arms and legs will also be tested.

These results supported by (Godoy, 2013) who reported that laboratory investigations and skull computed tomography should be obtained to establish a diagnosis of the complications and any postoperative neurological deteriorations detected during neurological examination and to plan new intervention, if necessary. (Edlinger et al., 2012) stated that to manage a postoperative neurosurgical patient the care provider requires knowledge of how the central nervous system reacts to stress and anesthesia as well as the potential complications associated with each specific procedure.

As regard vital signs in the present study we found that there was significant difference of body temperature during the third postoperative day. Regarding pulse there was moderate significant difference during the second postoperative day and high significant difference during the third postoperative day. Regarding respiration there was significant difference during the third postoperative day. Regarding blood pressure there was moderate significant difference in systolic and high significant difference in diastolic blood pressure during the third postoperative day.

So, the results of the present study show no signs of increased intracranial pressure postoperative. The abnormal changes that occur in vital signs may be attributed to emotional stress and fear in control group than in study group during the second and third postoperative days and this is also the neurosurgeon opinion in the neurosurgery department.

The presence of non significant differences in vital signs especially pulse and blood pressure during the first postoperative day can be attributed to anxiety and fear in both groups and these can produce hypertension and tachycardia as in the study sample. However, the study of **(Smith and Timby, 2013)** demonstrated that the blood pressure, pulse, respiration, and temperature are closely monitored on all patients with a potential or actual neurological disorder. The temperature often needs to be monitored every hour because central nervous system disorders can affect the temperature-regulating center of the hypothalamus. A sudden increase or decrease in any of the vital signs indicates a change in the neurological status, and the physician is notified immediately.

The results of the present study showed that there were non significant differences related to systemic and neurosurgical complications at the time of assessment; before application of the nursing educational program. This may be attributed to that the majority of studied patients suffer from anxiety and fear in addition to the immediate postoperative effect of anesthesia and this is also the opinion of the neurosurgeon and anesthesiologist in the neurosurgery department.

The results of the present study showed significant difference as regard systemic complications after application of the nursing educational program at the time of follow up (after 3 and 6 months). As regard neurosurgical complications, there was significant difference after application of the nursing educational program at the time of follow up (after 6 months).

The study finding was supported by **(Buttaro, 2008)** who demonstrated that unmanaged stress is linked to hypertension, heart diseases and gastrointestinal problems. Also, unmanaged stress is linked to some emotional health disorders. An elevated level of stress increases the frequency of abnormal behaviours.

(Silver et al., 2011) said that patients who have a successful outcome after intracranial operation may still experience residual neuropsychological symptoms that may be not noticed by the examiner but devastating to the patient. Over the years, many patients have expressed the opinion that some discussion of such possible sequelae before and after surgery would have been helpful to them. Neurosurgeons should routinely discuss possible neurological sequelae with patients before and after brain surgery. Providing patients with a simple brochure describing possible temporary or long-term neuropsychological consequences is suggested. Such information must be shared in a manner that does not frighten or upset the patient but provides true informed consent.

In the present study, all patients in both study and control groups before application of nursing educational program for patients after intracranial surgery had unsatisfactory level of knowledge regarding intracranial surgery.

This could be explained by the fact that, patients didn't receive enough information from health care team. Some patients were lacking interest to know any information while others were interested to know but they didn't find the person who had enough time to provide them with enough information. In the same line this result agree with **(Desoky, 2014)** study who said that "Impact of a designed nursing teaching protocol on quality of life of patients with chronic lower limb ischemia at Assiut University Hospital" which revealed that all patients in both study and control groups had unsatisfactory level of knowledge before application of a designed nursing teaching protocol.

After application of the nursing educational program for patients after intracranial surgery (before discharge, after 3 months and after 6 months), patients' knowledge score levels regarding intracranial surgery were highly significantly improved for study group patients. This improvement emphasis the fact that, most patients have a strong desire to learn more knowledge about their conditions and show the effect of the program.

This result is in the same line with the study of **(Ali, 2004)** entitled as "Impact of a designed nursing intervention protocol on performing self-care activities among rheumatoid arthritic women" which revealed that all studied sample had unsatisfactory level of knowledge about their disease at initial assessment, however, after application of a designed nursing intervention protocol, founded that, there was a significant increase in knowledge of study group patients regarding disease.

The results of the present study showed positive effect for the nursing educational program on patients' recovery and health for study group patients. Health promotion in this study was evaluated by assessing postoperative complications at the time of follow up and by using SF-36 questionnaire. The SF-36 self-evaluated health transition items (five response categories ranging from "much better" to "much worse") has been shown to be useful in estimating average changes in health status. It is individualized questionnaire; measured according to patient's response and patient's response differ from one patient to another.

These results were supported by **(Piper and Stewart, 2009)** who revealed that effective health educational program will result in changes that demonstrate increased knowledge about specific medical and health-related issues for a prolonged period of time.

Effective health education will yield both short-term and long-term changes in behavior that reduce risky behavior and/or improve quality-of-life. These changes in behavior can be recorded through evaluator observations and learner feedback, or through more formal means such as questionnaires.

Before discharge there was improvement in the scales concerned with role limitations due to emotional problems, social functioning and general health. After 3 months there was improvement in the scale concerned with role limitations due to emotional problems, vitality, mental health, social functioning, and general health. After 6 months there was improvement in all 8 scales plus health comparison question (health change).

Regarding health comparison question, this question was asked to the patients at the last time of follow up (after 6 months) to show the effect of the nursing educational program. The majority of patients (90 %) in study group were having health improvement while less than half of patients (43%) in control group were having health improvement.

In this regard this result agree with the study of **(Bin-Madhi, 2012)** entitled as "Brain tumors excision guided by neuronavigation: Practical application and results " which revealed 85% of postoperative patients showed an improvement of their neurological status.

However, **(Krug, 2008)** who conducted study entitled as "Functional outcome and self-perceived overall health status following surgery to remove primary brain tumor" which revealed that postoperative complications may delay improvement, though at three months, these complications and their effect on function should have been overcome. Functional recovery following surgery for brain tumor was not significant overall. When examined individually, a majority of subjects demonstrated improvement of a clinically relevant nature.

Conclusions

Based on the result of the present study, it can be concluded that:

- Patients after intracranial surgery; burr holes, craniotomy and craniectomy are at high risk for systemic and/or neurosurgical postoperative complications which had a bad effect on patients` health.
- Before application of the nursing educational program all patients in both study and control groups had unsatisfactory level of knowledge.
- The 1st hypothesis was supported as patients` knowledge score levels were highly significantly improved for study group patients after application of nursing educational program. Control group

patients who did not receive the nursing educational program had unsatisfactory level of knowledge at the time of follow up.

- The 2nd hypothesis was supported as a highly significant difference between study and control groups in relation to health improvement. The majority of patients in study group were having health improvement while less than half of patients in control group were having health improvement after application of nursing educational program.
- The 3rd hypothesis was supported as the study group patients suffered from less symptoms or complications (systemic or neurosurgical); physical, social or emotional problems than control group patients during follow up periods.

Recommendations

Based on results of the present study the following can be recommended:

For patients

Patients are to be provided with sufficient information to remind them with specific instructions regarding treatment, how to cope with postoperative periods; wound care, activity, rest, diet, stress and maintain effective communication to avoid certain postoperative complications that may develop and when they must notify physician.

For nurses

Nurses should be aware by instructions that given to patients before discharge and inform patients about them to improve their health.

For administration

Establishment of health care educational center in the neurosurgery department to educate patients about necessary instructions regarding their conditions using booklet and illustrated pamphlets for each patient especially those who cannot read and write.

For research

Similar studies should be replicated on longitudinal bases till one year as a minimum time period for follow up.

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