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# EFFECT OF APPLYING SPONTANEOUS PUSHING TECHNIQUE DURING SECOND STAGE OF LABOR ON WOMEN'S EARLY POSTPARTUM FATIGUE

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#### Abstract:

The aim of this study was to evaluate the effect of applying of spontaneous pushing technique during second Stage of labor in early postpartum fatigue. Methods: A quasiexperimental research design was used to carry out this study at the Labor and delivery unit of Mansoura University Hospital on 100 primigravida women at ≤37 gestational weeks, who were selected by purposive sampling technique. They were free from any medical and obstetric problems, can read and write. They were assigned either to the control or intervention group (n= 50 per each group). The intervention group: had received spontaneous pushing during second stage of labor, while the control group had received the routine pushing technique. Three tools for data collection were used; an interviewing questionnaire schedule, Visual analogue scale for fatigue and women's satisfaction questionnaire. Results: The study showed significant different between the two groups regarding the duration of second stage ,newborn health status & Apper score (7.8  $\pm 1$  versus 4  $\pm 1.4$  respectively, 0<0.001). Women expressed greater satisfaction with spontaneous pushing (98% versus 44% respectively, p<0.001). Conclusion spontaneous pushing technique is an effecting method in reducing women's fatigue. Recommendation Educating women about the spontaneous pushing technique in the first stage of labor and providing support for spontaneous pushing in the second stage for decrees postpartum fatigue.

Key word: second stage of labor, pushing technique, postpartum Fatigue

#### **Introduction:**

Labor and childbirth represent important life events fraught with strain and anxiety. Laboring women naturally seek the assistance of professionals who are experienced and qualified to secure an optimal birth outcome (15,10). She needed to explanation and give health education about labor and identified proper pushing techniques.

The second stage of labor is the stage of expelled of fetus from mother's body. The second stage subdivided into two phase, the latent phase full dilatation of the cervical os is recorded ,but the presenting parte may not yet have reached the pelvic outlet.

Women in this situation may not experience strong expulsive urge until the head has descended sufficiently to exert pressure on rectum and perineal tissues<sup>(1,2)</sup>. active pushing(bearing down) during latent phase versus the women's natural feeling to push and not achieve much. active pushing in this stage result increased exhaustion<sup>(3)</sup>.

Active phase during this phase the fetal head is lower in the pelvic floor and is stretching the perineum. the fetal head has rotated and started to descend (4,5,6).

Expulsion of the fetus out of the maternal body relies on pushing forces. Power forces are of two sources: the upper uterine contraction is contracted and relaxed the power from uterine contractions less due to the short ended upper uterine muscles. during this time the fetal head moves downward to the pelvic floor, about half of the fetal body has passed from the upper uterus (1,7). So the power from the existent short upper uterine contractions alone cannot push the fetus through the pelvic cavity (8).

Maternal bearing down (pushing) is the second force efforts, whose occur with contractions of the diaphragm and abdominal muscles. This is help the fetus to move downward to compress the pelvic floor, perineum and rectum, after that the sacrum and obdurate nerves as well as stretch receptors causes to an urge to push <sup>(7,8)</sup>.

There are two basic pushing methods commonly employed during the second stage of labour. directive pushing women are encouraged to take a deep breath, hold it and strongly push for as long as possible without releasing air or making any noise (10.9). This cause increases intrathoracic pressure, decrease in the oxygen supply to the fetus results in a marked fetal hypoxic, unnecessary instrumental delivery due to maternal exhaustion, burst blood capillaries in maternal eyes, and damage to the muscles of the vagina and perineum(10,11).

A woman who uses a spontaneous pushing approach is self-directed in her bearing-down techniques. She may push with an open glottis and vocalisation or use an intermittent exhalation technique. She will typically push three to five times for 5–7 seconds followed by a breath and release of air<sup>(12.13)</sup>.

The spontaneous pushing method has been found to have advantages over directed pushing, such as greater perineal integrity, fewer fetal complications including acidosis and heart rate alterations fewer maternal complications such as a change in blood pressure, less fatigue, and higher maternal satisfaction. Nonetheless, the spontaneous pushing method is not widely used in local maternity units. (9,10,12)

Postpartum fatigue has a major impact on women's lives, affecting their relationships with others and their ability to fulfil new roles (10,19,17,20).

Nursing care for women during second-stage labor is complex. It involves frequent and intelligent assessments of mother and fetus, promoting fetal descent, and supporting each woman's ability to cope with labor and pushing (13,12).

Good 'pushing techniques' nonmedical practice areas that can bring about significant improvements in maternal and fetal well-being<sup>(15)</sup>. relationship between postpartum fatigue and the pushing technique used during the second stage remains unclear. This study was evaluated effect of applying spontaneous pushing technique during the second stage of labor on women's postpartum fatigue, to assess their effect on women's perceived level of maternal fatigue immediately after and about 2 hours, 24 hours post birth.

# Research hypothesis

Spontaneous pushing technique used during second stage of labor less early post-partum fatigue

## **Subjects and Method**

**Study Design:** Aquasi-experimental design was utilized.

**Study Setting:** The study was conducted at the Labor and delivery unite of Mansoura University Hospital in Mansoura City.

**Subjects:** purposive sample, was selected according to the following formulae (*Senn*, 2007):  $n=2(Z_{\alpha/2}+Z_{\beta})^2\sigma^2/\Delta^2$ 

and include the criteria, gestation age at ≤37 gestational weeks they were free from any medical and obstetric problems, can read and write. The eligible 100 laboring women were equally divided into two groups: Intervention group (n=50): received a Spontaneous Pushing Technique during Second Stage of Labor while the control group (n=50): had received

the routine pushing (directive pushing).

**Tools of Data Collection:** three tools were used for data collection:

# Tool I: A Structured Interviewing Questionnaire Schedule:

This tool was developed and used by the researcher. It consists of 2 parts. **Part I:** Entails the participant's sociodemographic data, (e.g., age, education, occupation, marital status, residence, family income and telephone number).

Part II: Represents the current labour and delivery data for both mother and newborn. Maternal data includes duration of labour stages, use of Oxytocin, duration of pushing, episiotomy, tear and its degree, and degree of perineal pain (i.e. mild, moderate, or severe). Newborn data includes.

Tool II: Apgar Score. This is a simple method to assess the condition of the newborn infant. It is performed in the first minute and after five minutes of fetal expulsion (Appar et al, 1958). It is based on assessment of five physical signs, namely: heart rate, respiratory effort, reflex irritability, muscle tone, and color. The total score ranges from 0 to 10. A score of zero means none of these signs is present and 10 means a completely healthy infant. If the infant score ranges from 7 to 10, this indicates good infant condition. A score from 4 to 6 indicates moderate infant condition (moderate asphyxia), and from 0 to 3

indicates very bad infant condition (sever asphyxia).

# Tool III: Visual analogue scale for fatigue (VAS-Fatigue):

The original VAS-Fatigue scale consists of 18-items (13 items for fatigue and 5 items for energy). It is a valid tool which was previously tested for its validity and reliability (Lee et al., 1991). VAS-Fatigue was evaluated three times; immediately; 2-hours and 24-hours after labor; by asking each respondent to mark her level of fatigue/energy on 100 mm scale, and then the researcher measure the marking point and length (100 mm) using a standard metal ruler with mm marking to demonstrate the scores for fatigue and energy at each time point. It took less than 2 minutes to complete.

# Tool IV: Women's Satisfaction **Ouestionnaire**:

This tool was developed by **Yurachai.** (2006) and used by the researcher to assess the level of women's satisfaction regarding the pushing technique used during labor. It consist of 6 items. It was demonstrated by selecting one of two options; either satisfied score 1 or not satisfied score0.

#### **Ethical Considerations:**

- Ethical approval was obtained from the Research Ethics Committee of the Faculty of Nursing, Mansoura University.
- Official permissions were obtained from the head of the Obstetrics and Gynecology department and the

- Director of Mansoura University Hospital.
- The aim of the study was explained to the studied groups and informed consents were obtained.
- Ethical issues were considered in dealing with the obtained information.
- Women had the right to withdraw from the study at any time, and their data were confidential.
- Tool of data collection didn't touch religious, culture or ethical issues among mothers and mother's dignity was considered.

## **Pilot Study:**

After preparing the tools, a pilot study was conducted on 10 Parturient women (5 per each group). The results of the pilot indicated that the statements of the questionnaire were clear and relevant, and few words and items were modified. No modification with done.

#### Field Work:

■ The data were collected over a period of five months. During the study period; from June 2013 to October 2013, the researcher visited the study setting 3 days/week from 9 am to 3 pm. The researcher introduced herself and briefly explained the purpose and method of this study to the eligible women. The researcher then divided the sample into two groups beginning with the control group.

### **Control Group:**

This group was subjected to the conventional pushing (directed pushing) during the second stage of labor. The Obstetricians on duty determined the progression of labor by vaginal examination, when they detected that the cervix fully dilated, laboring women were encouraged by the researcher to push at the beginning of each uterine contraction, whether they had an urge to push or not. The detected researcher the uterine contractions by placing the palm of the hand over the woman's abdomen at the funds. As a contraction started, the laboring women were asked to take a deep breath and hold it while both hands hold on the bedside rails and push strongly for as long as possible (closed glottis) and were instructed to repeat the same procedure with every contraction until birth, and when the crown was visible at 2-3 cm, the laboring woman was sent to the delivery room using the lithotomy position to complete the birthing process.

# **Study Group:**

Parturient women in this group were subjected to spontaneous pushing technique. During the first stage, they were instructed by the researcher to relax during uterine contractions by inhaling deeply-slowly and exhaling deeply-slowly until the contraction had ceased (breathing exercise), while in second stage they were instructed to push spontaneously, by pushing only during contraction when they felt the urge to do so and rest in between, without any specific instructions about

the timing and duration of pushing. As in the control group subjects, when the crown was visible at 2–3 cm, the laboring woman was sent to the delivery room to complete the birthing process.

#### For both group

#### **Maternal assessment:**

- Levels of fatigue and energy were assessed using VAS-Fatigue immediately, at 2 and 24 hours postpartum. As well as, women's satisfaction regarding the technique that was assessed on the discharge time were the primary outcomes.
- Duration of the second stage, perineum status; either episiotomy incision or tear, and the level of perineal pain were the secondary outcomes.

#### **Neonatal assessment:**

- Apgar score was assessed at the 1<sup>st</sup> and 5<sup>th</sup> minute postpartum.
- Admission to the neonatal intensive care unit (NICU).

#### **Statistical analysis:**

After data were collected, it coded, organized, categorized, and then transferred into especially designed formats. The statistical analysis of data was done by using SPSS program (statistical package for social science) version 17.0. The data was tabulated and presented. The description of the data was done in form of mean and standard deviation for quantitative data (duration of pregnancy), frequency and proportion for qualitative data. Analysis of the data was performed to

test statistical significant difference between variables for both groups (intervention and control). For disease duration (mean and standard deviation) independent t- Test was used to compare between the two groups. For qualitative data (frequency and proportion), Chi- square test was used. Statistical significant difference was considered at P<0.05, and highly significant difference at P<0.001.

#### **Results:**

Table (1) Distribution of the general characteristics of both intervention and control groups according to their demographic characteristics. Data reveals no statistically significant differences observed among the studied groups related to age, education level, occupation and residence (P >0.05).

Table (2) Comparison of the current labor and delivery data between the both intervention and control groups. Data reveals that significantly longer in the directed pushing group compared to the spontaneous pushing women in the directed pushing group needed analgesics while none of the women in the spontaneous pushing needed analgesics. This difference significant (p=0.006).

Table (3) Comparison of the maternal and newborn condition between the both intervention and control groups: reveals that the duration of pushing during second stage was significantly longer in the directed pushing group than in the spontaneous pushing group (50.3 ±5.8)

versus  $29.5 \pm 6.6$ , p<0.001) reveals that the duration of pushing during second stage was significantly longer in the directed pushing group than in the spontaneous pushing group (50.3  $\pm$ 5.8 versus  $29.5 \pm 6.6$ , p<0.001). Women in the directed pushing group 3 (8.1%) had mild tear, 25 (43.2%) had moderate tear and 24 (48.6%) had severe tear (p<0.001).

Table (4) Comparison of the women's satisfaction between the both intervention and control groups: find statistical significant in the intervention group that is in control group.

Figure (1) The changes of VASfatigue total score of the intervention and the control groups immediately, two hours and 24 hours postpartum It is clear from this table that the VASfatigue total score of the spontaneous pushing group was lower (104.2±5.7,  $57.1\pm7.8$ ,  $28.7\pm5.9$ ) and that of the group directed pushing was  $(112.2\pm4.0,$  $71.7\pm7.6$  $40.8\pm6.3$ ) immediately, two hours and 24 hours postpartum respectively.

**Figure (2)** the changes of VAS-energy total score of the spontaneous and the directed pushing groups immediately, two hours and 24 hours postpartum. It is clear from this table that the VAS-energy total score of the spontaneous pushing group was  $(44.0\pm2.1,\ 34.6\pm4.4,\ 16.6\ \pm3.2)$  and that of the directed pushing group was  $(41.1\pm3.3,\ 29.7\pm5.3,\ 12.0\ \pm4.6)$  immediately, two hours and 24 hours postpartum respectively

**Table 1.** Distribution of the general characteristics of both intervention and control groups according to their demographic characteristics

	Inter	vention	Caret		Chi a	
	g	roup	Cont	rol group	Chi squ	are test
	N	%	n	%	$X^2$	P
Age (years) (mean $\pm$ SD)	22.6 ±	3.4	22.8	±3.5	0.233*	0.816
Residence						
Rural	31	62	29	58	0.167	0.683
Urban	19	38	21	42		
Marital status						
Married	48	80	47	84	0.344	0.842
Divorced	1	0	1	0		
Widow	1	0	2	0		
Education level						
Illiterate	2	4	3	6	0.597	0.742
Primary/preparatory	6	12	8	16		
Secondary/university	42	84	39	78		
Occupation						
Housewife	25	50	21	42	0.644	0.422
Working	25	50	29	58		
Income						
Enough	15	30	10	20	1.333	0.248
Not enough	35	70	40	80		

<sup>\*</sup> Student's t test

**Table 2.** Comparison of the current labor and delivery data between the both intervention and control groups

	Inter- group	vention	Contr		Student	's t test
	Mear	n±SD	Mean	±SD	t	P
Duration of stages of labor						
1 <sup>st</sup> stage (minutes)	15.1	$\pm 0.8$	15.0	$\pm 0.8$	0.369	0.713
2 <sup>nd</sup> stage (minutes)	104.8	$3 \pm 3.4$	115.2	$2 \pm 2.9$	16.614	< 0.001
3 <sup>rd</sup> stage (minutes)	20.3	$\pm 1.4$	20.0	$\pm 1.5$	0.893	0.374
Use of analgesics (n, %)	0	0	7	14	7.527*	0.006
Use of oxytocin (n, %)	30	60	45	90	12.00	< 0.001
Spontaneous membrane rupture	50	100	50	100	0*	1
Amniotic fluid characteristics						
Clear	50	100	45	90	5.236*	0.022
Meconium	0	0	5	10		

<sup>\*</sup> Chi square test

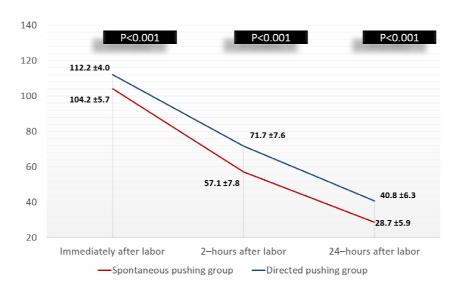
**Table 3.** Comparison of the maternal and newborn condition between the both intervention and control groups

intervention a		·	~		~1.	
Variables	Intervent	ion group	Contro	ol group	Chi squ	are test
	N	%	n	%	$X^2$	P
Duration of pushing						
<60 seconds	50	100	0	0	88.104	< 0.001
60 - 90 seconds	0	0	36	72		
>90 seconds	0	0	14	28		
Degree of tear**						_
No	38	95	0	0	72.193	< 0.001
Mild	2	5	3	8.1		
Moderate	0	0	16	43.2		
Severe	0	0	18	48.6		
Episiotomy	10	20	13	26	0.508	0.476
Perineal pain						
Mild	46	92	0	0	86.207	< 0.001
Moderate	4	8	25	50		
Severe	0	0	25	50		
Apgar score						
1 minute	$7.8 \pm 1$		$4 \pm 1.4$		15.628*	< 0.001
postpartum						
5 minutes postpartum	$9.5 \pm 0.5$		9.4		0.750*	0.455
			±0.8			
Admission to the NICU	2	4	9	18	5.005	0.025

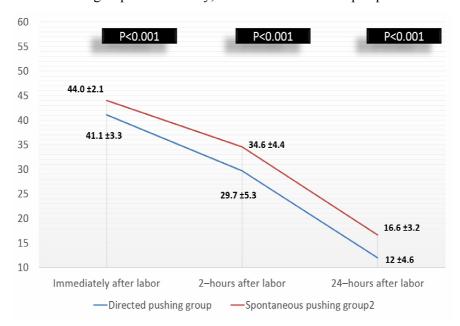
<sup>\*</sup> Student's t test, \*\* n=40 in the intervention group, n=37 in the control group

**Table 4.** Comparison of the women's satisfaction between the both intervention and control groups

_			Control	group	Chi square test		
	n	%	n	%	$X^2$	P	
xperence has shown that I can							
ave appropriate and adequate							
ontrol over my care	48	96	39	78	7.16	0.007	
he pirsons responsible for my							
re were caring and							
ompassionate	45	90	37	74	4.34	0.037	
roblems that have arisen up to							
ow have been dealt with							
fectively	42	84	31	62	6.14	0.013	
ly needs have been addressed							
ith appropriate consideration for							
y time	44	88	36	72	4.00	0.046	
he overall organization of my							
are has been appropriate	47	94	39	78	5.32	0.021	
ould choose the same type of							
re for my next labor	49	98	22	44	35.4	< 0.001	



**Figure 1.** The changes of VAS-fatigue total score of the intervention and the control groups immediately, one hours and 2 hours postpartum



**Figure 2.** The changes of VAS-energy total score of the spontaneous and the directed pushing groups immediately, two hours and 24 hours postpartum.

Table (2): Represents comparison between control group and study (SCG) group regarding alopecia questions that all women (100%) in control group and the majority of the women (96.7%)studygroup(SCG) were suffered from hair loss, and all the women (100%) in both groups suffering hair loss more than a month. There is no significant difference between the study group (SCG) & the control group regarding alopecia questions (P<0.05). As regard to the degree of hair loss according to

WHO scaling system for hair loss there is highly significance difference between the two groups regarding degree of hair loss (p=.000) as in the control group there were (46.7%) suffering total alopecia in comparison to study group (SCG) there were (3.3%) suffering total alopecia. This means that the study group (scalp cooling group) suffered less hair loss after receiving scalp cooling than the control group who receive the routine hospital care.

**Table (2):** comparison between control group and study group regarding alopecia questions

questions						
		rol group		ly group		
Alopecia questions	(N	V=30)	(N	0 = 30	X <sup>2</sup>	P - value
	No	%	No	%		
Suffering from hair loss:-						
Yes	30	100.0%	29	96.7%	1.017	.313
No	0	.0%	1	3.3%		
Time of hair loss:-						
More than a month	30	100.0%	30	100.0	-	-
				%		
Degree of hair loss:-						
0-Not significant hair loss	0	0%	1	3.3%		
1-Minor hair loss not	0	0%	19	63.3%	37.06	.000**
requiring a wig	4	16.7%	6	20.0%	7	
2-Moderate hair loss not						
requiring a wig		40.0%				
3-sever hair loss, requiring	12	43.3%	3	10.0%		
a wig	14		1	3.3%		
4- Total alopecia						
Other scalp problems						
before chemotherapy:						
No	30	100.0	30	100.0	-	-
		%		%		

**Table (3):** Represents the prevalence of scalp cooling side effect, where 100% of study group(scalp cooling group) suffered from feeling cold and

70% needed blanket for warmth during doing this procedure. And 100% are suffering from headache and feel boring

**Table (3):**Prevalence of side effects of scalp cooling among scalp cooling group.

Scalp cooling side effects		dy group No =30)
	No	%
Feeling cold:-		
Yes	30	100.0%
No	0	0%
Blanket for warmth:-		
Yes	21	70.0%
No	9	30.0%
Headache :-		
Yes	30	100.0%
No	0	0%
Feeling boring:-		
Yes	30	100.0%
No	0	0

**Table (4)**: Represents comparison between control group and study group(SCG) in grades of hair loss before the first cycle and after the six cycle of chemotherapy there were (63.3%&0%) have grade1 from the study group (SCG)andcontrol group respectively. While there were

(43.3%&3.3%) have grade 4(total alopecia) from the control group and study group(SCG) respectively after the six cycle of chemotherapy. There is highly significance difference between the two groups regarding grades of hair loss. P<.05

**Table (4):** Comparison between control group &study group (scalp cooling group) in grades of hair loss according to WHO before the first cycle and after the six cycle of chemotherapy.

Grades of		Contro	l grou	р		Study g	roup (SC	G)	X 2		P - value	)
hair loss	Befo	fore cycle		After cycle 6		fore cle 1	After cycle 6		Before cycle 1	After cycle 6	Befor e cycle	After cycle 6
	No	%	No	%	No	%	No	%			1	
0	30	100%	0	0%	30	100	1	3.3%	1.017	35.777	.313	.000**
1	0	0%	0	0%	0		19	63.3%				
2			5	16.7%		0%	6	20.0%				
3			12	40.0%			3	10.0%				
4			13	43.3%			1	3.3%				

Table (5):Represents the relation between the type of chemotherapy and grades of hair loss after the sixth cycle of chemotherapy in both control group and study group (SCG) that there were from the women with Taxol-carpolinate in study group have gradeI hair loss and also 40% have grade II hair loss while 60% from the women who treated with taxolcarpolinate in the control group have grade III hair loss. In relation to the women treated with FAC Adriamycin-(Floururacil

Cyclophosphamide {Endoxan}) there were 60% from the women in study group(SCG)have grade I hair loss while in the control group there were 70.6% have grade IV hair loss. As regard to the women who treated with Floxthere were 80% from the women in the study group have grade I hair loss while 62.5% from the women in the control group have grade III. There were significance differences between the two groups in grades of hair loss in relation to type of chemotherapy

**Table (5)** The relation between grades of hair loss and type of chemotherapy after the six cycle of chemotherapy among study group (scalp cooling group) and control group.

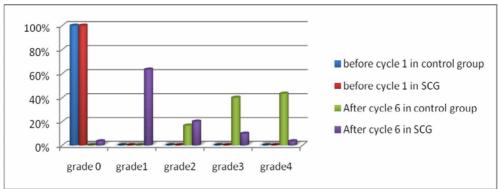
		Туре	of chemotl	herapy in SCG	group			Type of	chemoth	erapy in contro	ol group									
	Ta	axol –	FAC(F	Floururacil-	F	oulox	Ta	axol –	FAC(I	Floururacil-	F	oulox								
Grades	сат	oolinate	Adr	iamycin-			carı	oolinate	Adı	riamycin-				X 2	P -	· value				
of hair			Cyclop	hosphamide						Cyclophosphamide										
loss		l	{En	idoxan})					{Endoxan})		{Endoxan})		{Endoxan})							
	No	%	No	%	No	%	No	%	No	%	No	%	SCG	Control	SCG	Control				
Grade 0	0	0%	0	0%	1	10.0%	-	-	-	-	-	-								
Grade 1	2	40.0%	9	60.0%	8	80.0%	-	-	-	-	-	-								
Grade 2	2	40.0%	3	20.0%	1	10.0%	1	20.0%	1	5.9%	3	37.5%	7.061	15.962	.530	.003*				
Grade 3	1	20.0%	2	13.3%	0	0%	3	60.0%	4	23.5%	5	62.5%								
Grade 4	0	0%	1	6.7%	0	0%	1	20.0%	12	70.6%	0	0%								

**Table** (6):Represents the relation between age & grades of hair loss after the 6th cycle among study group(scalp cooling group)(SCG) that all women whose age ranged from (18 – 39) years old have gradeI hair loss and 46.2% from the women whose age ranged from(40 – 49) have gradeI

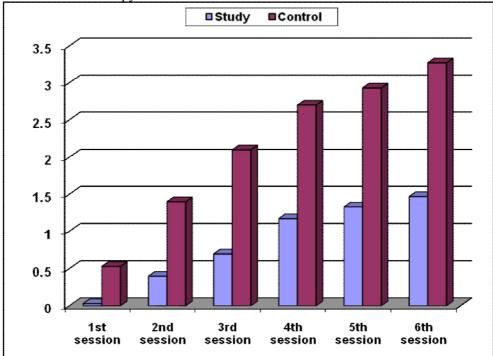
hair loss.Otherwise 40% from women aged from(50 – 60) have gradeII hair loss. This means that there was a significance difference between younger age women (most of them have gradeI) & the older age women (which they have gradeII,III,IV) and the grades of hair loss. (P<0.05)

**Table (6)** Relation between age & grades of hair loss after 6<sup>th</sup> cycle among study group (scalp cooling group).

Condon								
Grade of	18	8 < 39	4	0 < 49	5	0 < 60	X <sup>2</sup>	P - value
hair loss	No	%	No %		No	%		
Grade 0	0	0%	1	7.7%	0	0%		
Grade 1	12	100%	6	46.2%	1	20.0%	17.174	.028*
Grade 2	0	0%	4	30.8%	2	40.0%	17.174	.020
Grade 3	0	0%	0% 2 15.4% 1		20.0%			
Grade 4	0	0%	0	0%	1	20%		



**Fig (1)** Comparison between control group &study group (SCG) in grades of hair loss according to WHO before the first cycle and after the six cycle of chemotherapy.



**Fig (2)** Comparison between the control and study group(SCG) in the six cycles of chemotherapy in relation to hair loss

# Discussion:

The current study was aimed to evaluate effect of applying spontaneous pushing technique during second stage of labor in early postpartum fatigue.

The present study found no significant differences between the two groups in their demographics as

regards the (age of the participants, residence, marital status, educational level, occupation status and income).

Several causes of maternal fatigue in the early postpartum ,include physical factors such as length of labour, type of delivery and blood loss associated with delivery, maternal hormonal shifts, episiotomy healing, pain , mode of delivery, and pushing technique used during labour also affect maternal fatigue .

present study found The significantly higher fatigue in the directed pushing at immediately, two, 24-hours postpartum. These can be related to the study findings reveals that there the 2<sup>nd</sup> stage of labor was significantly longer and increase of duration pushing time in the directed pushing group compared to the spontaneous pushing group. The spontaneous pushing causes decrease pain sensation and fatigueon early postpartum period. Accordingly, the study hypothesis is accepted. These findings are consistent with those of Christine and Susan . (2010) and Gillesby et al. (2010) reported that directed pushing longer and strong push every uterine contraction and immediate full cervical dilatation during the second stage of labour led to higher fatigue scores at 24 hours after birth(15,3). Yildirim and Beji (2008); Su-Chen Kuo, et al. (2011) and Yurachai . (2006) who reported that women use spontaneous pushing low post partum fatigue(10,12). The present study was agreed with the Methodius et al, (2012) there reported that second stage of labor duration less for the spontaneous pushing, compared

with the directive pushing. Similarly, Lai et al. (2009) the researcher found that the duration of the expulsion phase were significantly longer with directed pushing.

On another hand the present study found spontaneous pushing technique improved circulation (increased blood supply of uterus) and decrease sensation of pain and decrease of needed analgesics. This finding supported by Albers and Borders. (2007) and Su-Chen Kuo, et al. (2011)who reported directive pushing during the second stage of labor in mothers receiving analgesics. Lai et al. (2009) there reported that women in the directive pushing had a higher pain.

The present study found spontaneous pushing improved placenta perfusion, fetal oxygenation cause the Apgar score at 1 minute was significantly higher in the babies of the women of spontaneous pushing group. However, at 5 minutes, Apgar score did not differ significantly between the two groups. The study supported by Lai et al., (2009) who reported that spontaneous group had significantly higher one minute Apgar scores (x2 =8.696, p < .01). There was no between difference groups for 5minute Appar scores (p = 0.001) or cord pH levels (p = 0.004). Samer, N(2005): Yildirim and Beji. (2008) with spontaneous pushing, with higher 1- and 5-minute Apgar scores, and higher umbilical cord pH and Po<sub>2</sub> levels (22,12). Osborne. & Hanson. (2012) Directive versus Supportive Approaches Used by Midwives When Providing Care during the Second

Stage of Labor. The researcher findings the group use spontaneous pushing fetal and neonatal outcomes include improved fetal oxygenation (11).

The present study findings women who are uses bear down spontaneously report higher levels of satisfaction The study supported by. **Ibrahim,S.** (2012): Yurachai . (2006): Yildirim and Beji (2008). Reported that spontaneous pushing technique increased satisfaction levels for women giving birth(10, 23, 18).

#### **Conclusion:**

The findings of this study the spontaneous pushing reduce of fatigue on women's early postpartum.

### **Recommendations:**

On the basis the most important findings of the study the recommendation following are suggested:

- Educating women about the spontaneous pushing technique in the first stage of labor and providing support for spontaneous pushing in the second stage for decrees postpartum fatigue
- Dissemination of the present study finding to all hospital and MCH health services at Mansoura city.

# **Further study**

 Rising awareness for caregivers working in the delivery room spontaneous pushing during the second stage of labor.

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