Maternal and fetal outcomes in women undergoing caesarean section under general and spinal anesthesia

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ABSTRACT

Background: Both spinal and general anaesthesia used for caesarean section have certain advantages and disadvantages and there is no method which is completely ideal. The most important factors for choice of anaesthesia are; pregnant systemic problems and wishes, the urgency of the operation, and the surgeon and the anesthetists experience.

Aim of the Work: These studies aimed at comparing maternal and fetal outcomes in women undergoing elective caesarean section and have spinal anaesthesia with those having general anaesthesia.

Subjects and Methods: This study was carried out at Ain Shams University Maternity Hospital during the period from December 2017 to August 2018 after approval of the hospital health ethical committee. It included 186 patients who had C.S and were subdivided into 2 groups according to a randomization scale. On the day of the operation, each randomly received a closed opaque envelope for the selection of the procedure (spinal versus general).

Results: We noted that the mean haemoglobin and haematocrit values at the 24th hour were higher in the spinal anaesthesia group. The estimated blood loss volume was significantly higher in the general anaesthesia group. The median apgar score at the first and the fifth minutes were significantly higher, and the time that elapsed until the first requirement for analgesia was significantly longer in the spinal anaesthesia group.

Conclusion and Recommendations: General anaesthesia could be thought the quickest anaesthesia method in an emergency since it avoids the possibility of a failed regional block. Meanwhile, it is associated with higher possibility of blood loss and low Apgar score. Thus, using spinal anaesthesia for elective caesarean section is recommended provided that adequate maternal hydration is established and sparing general anaesthesia for emergency caesarean sections or whenever spinal anaesthesia is contraindicated (e.g. coagulopathy, sever thrombocytopenia, anticoagulation or sever degree of malformation of spine).

Keywords: Accidental awareness during general anesthesia; combined spinal and epidural; General anesthesia.

INTRODUCTION

Delivery by caesarean section is by far one of the most commonly performed operations all over the world. Approximately 18.5 million caesarean sections are performed yearly worldwide⁽¹⁾.

About 40% of the countries have CS rates <10%, about 10% have CS rates between 10 and 15%, and approximately 50% have CS rates>15%. Countries with CS rates <10% account for only 25% (4.5 millions) of the global CS but for 60% (77 millions) of the total number births worldwide. On the other hand, 73% (13.5millions) of the total number of CS are performed in the countries with CS rates >15% where 37.5% (48.4 millions) of the total number of births occur⁽¹⁾.

In Egypt, more than one-half of deliveries in the five-year period before the 2014 were by caesarean section. Caesarean deliveries were more common in the urban areas compared to in rural areas (60% and 48% respectively)⁽²⁾. When medically justified, caesarean section can effectively prevent maternal and perinatal mortality and morbidity. However, there is no evidence showing the benefits of caesarean delivery for women or infants who do not require the procedure. As with any surgery, caesarean sections are associated with short and long term risk which can extend many years beyond the current delivery and affect the health of the woman, her child and future pregnancies. These risks are higher in woman with limited access to comprehensive obstetric care ⁽³⁾.

This operation requires effective anaesthesia which can be regional (epidural or spinal) or a general anaesthesia. The type of anaesthesia used and the care with which it is administered is an important determine of the outcome of caesarean section ⁽⁴⁾.

Both spinal and general anaesthesia used for caesarean section have certain advantages and disadvantages and there is no method which is completely ideal. The most important factors for choice of anaesthesia are; pregnant systemic problems and wishes, the urgency of the operation, and the surgeon and the anesthetists experience⁽⁵⁾.

General anaesthesia refers to the loss of the ability to perceive pain associated with loss of consciousness produced by intravenous or inhalational anaesthetic agents. For caesarean section, this involves the use of thiopentone for induction, tracheal intubation facilitated by suxamethonium, positive-pressure ventilation of the lung with a nitrous oxide/ oxygen mixture plus a volatile agent, and a muscle relaxant⁽⁶⁾.

Spinal anaesthesia refers to the use of local anaesthetic solutions to produce circumscribed area of loss of sensation. The spinal anaesthesia used for caesarean section involves the infiltration of a local anaesthetic agent, usually bupivacaine, into the surroundings of the spinal cord through the lower back of the woman (the drug is injected directly into the subarachnoid space)⁽⁶⁾.

Over the last 30 years, the use of spinal anaesthesia is rapidly increasing ⁽⁷⁾.

Spinal anaesthesia is relatively easy to perform, gives excellent anaesthesia a low potential of toxicity, allows mother to be awake and interact immediately after the birth of the baby. Compared to general anaesthesia it offers less maternal morbidity, comparable less blood loss ⁽⁸⁾.

It also enables early recovery of gastrointestinal functions, prolonged interval to first analgesic requirement, less analgesic consumption and early ambulation⁽⁹⁾.

However, spinal anaesthesia is not free from side effects and has its own complication like maternal hypotension, hypothermia, post-operative headache, accidental total spinal anaesthesia and patients at risk of heavy peripartum haemorrhage may not tolerate the haemodynamic effects of regional anaesthesia⁽¹⁰⁾.

General anaesthesia is a more quickly administered procedure and is often preferred in cases where speed is important ⁽¹¹⁾.

It also used in certain situation like contraindication to regional anaesthesia, failed regional anaesthesia and maternal request ⁽¹²⁾.

The risks include aspiration of stomach contents, awareness to surgical procedure (due to inadequate anaesthesia), failed intubations, and respiratory problems for both mother and baby ⁽¹³⁾.

In general, general anaesthesia is preferred for emergency caesarean section since it provides rapid onset of action and more stabilization of the patient's circulation and vital signs, on the other hand regional anaesthesia is preferred for elective operations because of its lower risk of drugs complications to the mother and the fetus⁽¹⁴⁾.

PATIENTS AND METHODS

Type of study:

This study was carried out at Ain Shams University Maternity Hospital during the period from December 2017 to August 2018 after approval of the hospital health ethical committee. It included 186 patients who had C.S and were subdivided into 2 groups according to a randomization scale. On the day of the operation each randomly received a closed opaque envelope for the selection of the procedure (spinal versus general). The study was approved by the Ethics Board of Ain Shams University and an informed written consent was taken from each participant in the study.

Inclusion Criteria: Full term. Singleton pregnancy. Maternal age from 20-35. Elective caesarean section.

Exclusion criteria: Multiple pregnancies. Grand multiparity (more than 4 deliveries). Macrosomia (more than or equal 4500 grams). Polyhydramnios (defined as amniotic fluid index more than 25 centimeter). Placental abnormalities such as placental abruption. Preterm rupture of membrane. Preterm delivery (before the 37thweak of pregnancy). Post term delivery (pregnancy exceeding 40th gestational weak). Preeclampsia. Gestational diabetes. Unsuitability for regional anesthesia

Elimination criteria: Refusal to participate after counseling. Any intraoperative complication. Presence of intestinal or omental adhesions. Insertion of intra-peritoneal drain. Excessive small bowel manipulation. The study was prospective, randomized, controlled trial.

Randomization: Computer generated randomization of the two groups (spinal anesthesia and general anesthesia) was done. Before the operation each participant was received a closed opaque envelope for the selection of the procedure. (Spinal versus general).

Methodology:

Patients were admitted from the outpatient clinic at Ain Shams University Maternity Hospital and were subjected to the following: Counseling about the two types of anesthesia and explanation of the procedure. An informed consent to this participation. Full history taking: Full name, age, gravidity and parity. Past medical history of hypertension, DM and other endocrine diseases, also surgical history of abdominal operations. Examination: Vital signs (pulse, temperature, blood pressure and respiratory rate). Measurement of weight (Kg), height (m) to calculate the BMI. Full laboratory investigations especially (CBC, liver and kidney function and RBS). Ultrasound (for assessment of fetal wellbeing). All women fasted at least 6h prior to the surgery.

Steps:

On arrival to the operation room, standard monitoring was applied with noninvasive blood pressure measurement, electro cardiography and pulse oximetry.

General anesthesia:

Following *Diemunsch and Noll*⁽¹⁵⁾ parturients in this group received standard rapid sequence induction with pre-oxygenation for 3 minutes followed by 4-5 mg/kg succinylcholine, cricoid pressure was applied throughout induction once necessary. After correct placement of the tracheal tube was confirmed, anesthesia was maintained with up to 1.5% isoflurane and oxygen, neuromuscular blockade was maintained with 0.4 mg/kg atracurium.

Spinal anesthesia:

Following *Armstrong*⁽¹⁶⁾, parturients in this group were rehydrated with 500ml lactated ringer solution intravenously within 15 min in the sitting position. Low back was prepared and draped in a sterile fashion with betadine solution 10%.

Spinal anesthesia was performed at L2-3 or L3-4 Inter vertebral space using a fine spinal needle (size 22G 3.5 inch). Injection of local anesthetics into the subarachnoid space, Bupivacaine (Marcaine) (1.5-3.5ml) was used.

Operative data: C.S was done by the senior resident according to standard technique demonstrated by *Louis et al.* ⁽¹⁷⁾. The skin was opened with the modified pfannenstiel incision. The anterior abdominal wall was opened in layers. The peritoneum was opened by elevating it with two clamps placed about 2 cm apart. The peritoneum is incised sharply superiorly to the upper pole of the incision and down warded to just above the peritoneal reflection over the bladder. The lower flap of visceral peritoneum was opened in the bladder is gently separated by blunt dissection from the underlying myometrium. The uterus was opened in the lower segment (the lower uterine segment is incised transversely). The baby was delivered: a hand is slipped into the uterine

cavity. The head is elevated with the fingers and palm through the incision then the shoulder and baby was delivered. After the shoulders are delivered, an intravenous infusion containing (20 units) of oxytocin per liter of crystalloid is infused at 10 ml/min until the uterus contracts satisfactorily. The placenta was then delivered by spontaneous delivery, with some cord traction. The uterine incision was then closed with two layers using blunt needle and continuous absorbable suture (Vicryl No.0) intra abdominally. The visceral and parietal peritoneum were closed using continuous absorbable suture (Vicryl No. 0). The recti muscles were approximated with two figure-of-The rectus sheath and eight sutures of 0 Vicryl. subcutaneous tissue was sutured using continuous absorbable suture (Vicryl 1) and the skin was closed by subcuticular suture (Prolene 2.0). The evaluation of the newborn was performed by the pediatrician who was present in the operating room. The apgar scores in the 1st and 5th minutes after the birth were recorded.

After the operation: After the operation all patients were transferred to post-operative room for 6 hours where they were under close observation for vital data, vaginal bleeding and urine output and then transferred to the word until discharge. Both groups had the same hospital fluid regimen which is 500cc of 5% glucose every 6 hours, 500cc of ringer every 12 hours and 500cc of saline 9% every 24 hours. All participant received the same intra operative prophylactic antibiotic Amoxicillinetrihydrate +Flucloxacilline monohydrate 1:1 (Flumox) vial 1gm before skin incision that had been repeated every 8hrs for the first 24hrs and from the same formula one capsule 500mg tds for one weak was recommended. For postoperative analgesia, intramuscular doses of 75mg diclofenac sodium (Voltaren, Novartis Pharma, Egypt), a nonsteroidal anti-inflammatory medication, were offered. The first was given once needed after waning of the effect of anesthesia and the second 12 hours later. The time needed for first analgesic request was recorded. Auscultation for intestinal sound was started 2 hours after operation and was performed at one hour interval till normal bowel sounds were detected. Patients were observed if they experienced nausea or not(the patient reported that she had sensation of the need to vomit) at 6 hours intervals after the operation. The presence of vomiting or not was observed and recorded at 6 hours intervals after the operation. The presence of shivering or not was observed and recorded. Patients were asked if they experienced headache attacks. CBC was done 24 hours postoperative. No oral or rectal bowel stimulants were given after surgery. Urinary catheter was removed 6 hours postoperatively and patients were encouraged to ambulate. Eligible criteria for hospital discharge included, stable vital signs with no febrile morbidity for at least 24 hours, ability to ambulate and urinate without assistance, passage of a bowel motion, ability to tolerate solid food without emesis and absence of unresolved other postoperative complications.

Statistical Methods:

Data were analyzed using MedCalc[©] version 18.2.1 (MedCalc[©] Software bvba, Ostend, Belgium).

Numerical variables were presented as mean and SD and between-group differences were compared using the unpaired t test.

Categorical variables were presented as number and percentage and differences were compared using Fisher's exact test. Ordinal data were compared using the chi-squared test for trend.

Time to event analysis was done using the Kaplan-Meier method P-values <0.05 were considered statistically significant.

RESULTS

The current study was conducted on 186 pregnant women at Ain Shams University Maternity Hospital from December 2017 to August 2018 to compare the maternal and fetal outcomes after general versus spinal anaesthesia in caesarean section.

Demographic data of the patients participated in the study as shown in the next table (1).

 Table (1): Demographic characteristics of patients

 in both groups.

Variable	Spinal anesthesia (n=93)		General anesthesia (n=93)		Difference	95%CI	P-value*	
	Mean	SD	Mean	SD				
Age (years)	28.5	4.7	27.8	5.3	-0.7	-2.1 to 0.7	0.341	
BMI (kg/m ²)	26.8	1.8	26.7	1.6	-0.1	-0.6 to 0.4	0.651	

Data are mean and standard deviation (SD).

95% CI = 95% confidence interval.

*Unpaired t test.

There were no significant differences between women of both groups regarding age and body mass index.

 Table (2): Obstetric history of patients in both groups.

Parity	Spina	l anesthesia (n=93)	Gener	al anesthesia (n=93)	χ2(df=1)	P-value*
-	Ν	%	n	%		
PG	16	17.2%	26	28.0%		
P1	22	23.7%	19	20.4%	1.942	0.175
P2	25	26.9%	22	23.7%	1.042	0.175
P3	30	32.3%	26	28.0%		

Data are number (n) and percentage (%).

 $\chi 2$ = chi-squared s statistic, df = degree of freedom.

*Chi-squared test for trend.

Table (2) illustrates that there is no statistically significant difference between study groups regarding the parity.

Table (3): Indication for CS in both study groups.

Indication for CS	an (Spinal esthesia n=93)	(ar	General nesthesia (n=93)	P-value*
	Ν	%	Ν	%	
Previous CS	44	47.3%	41	44.1%	
Obstructed labor	13	14.0%	9	9.7%	
Failed progress	9	9.7%	21	22.6%	
Breech presentation	14	15.1%	9	9.7%	0.289
Transverse lie	1	1.1%	3	3.2%	
Infertility	7	7.5%	6	6.5%	
Cardiac disease	3	3.2%	2	2.2%	
Previous cervical repair	2	2.2%	1	1.1%	
Epilepsy	0	.0%	1	1.1%	

Data are number (n) and percentage (%).

*Fisher's exact test.

Table (3) illustrates that there is no statistically significant difference between study groups as regards to the indication of caesarean section.

Table (4): Maternal outcomes in both study
groups.

Variable	Spinal anesthesia (n=93)		General anesthesia (n=93)		Differenc	95% C	P-
	Mea n	SD	Mea n	SD	e	1	value*
Postoperative hemoglobin (g/dl)	10.4	1.1	9.4	1.5	-0.9	-1.4 to - 0.6	<0.000 1
Absolute Hemoglobin drop (g/dl)	0.91	0.66	1.29	0.95	0.38	0.1 to 0.6	0.002
Postoperative hematocrit (%)	32.2	3.2	30.1	4.0	-2.0	-3.1 to - 1.0	< 0.001
Absolute Hematocrit drop (%)	2.9	1.9	3.4	2.8	0.5	-0.2 to 1.2	0.133
EBL (ml)	411.6	100. 3	501.2	119.7	89.5	6.3 to 172.8	0.035
TFA request (min)	2.4	1.8	1.7	1.2	0.7	0.3 to 1.1	0.002
Time to recover intestinal sounds (h)	6.0	1.3	7.1	1.5	1.1	0.7 to 1.5	<0.000 1

Data are mean and standard deviation (SD); 95% CI = 95% confidence interval.

*Unpaired t test.

Table (4) displays that there is statistically significant differences with p- value (< 0.0001) between study groups as regards to the

postoperative hemoglobin with high mean among spinal anesthesia group.



Fig (1): Postoperative hemoglobin in both study groups. Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations.

Table (4) reveals that there is statistically significant difference with p-value (0.002) between study groups as regards the absolute hemoglobin drop (g/dl) with high mean among general anesthesia group.



Fig (2): Drop in hemoglobin in both study groups. Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations

Table (4) demonstrates that there is statistically significant difference with p-value (0.001) between study groups as regards to the postoperative haematocrit with high mean among spinal anesthesia group.



Fig (3): Postoperative hematocrit in both study groups. Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations

Table (4) shows that there is no statistically significant difference with p-value (0.133) between study group as regards to the absolute haematocrit drop values (%).



Fig (4): Drop in hematocrit in both study groups. Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations.

Table (4) illustrates that there is a statistically significant difference with p-value(0.035) between study groups as regards to the estimated blood loss (ml) with high mean among general anesthesia group.



Fig (5): EBL in both study groups. Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations.

Table (4) manifests that there is statistically significant difference with p-value (0.002) between study groups as regards to the first analgesic request (hr) with high mean among spinal anesthesia group.



Fig (6): TFA request in both study groups. Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations.

Table (4) establishes that there is statistically significant difference with p-value (0.0001) between study groups as regards to the first intestinal sound (hr) with high mean among general anesthesia group.



Fig (7): Time to recover intestinal sounds in both study groups. Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations.

Table (5): Fetal outcomes in both study groups.

Variable	Spinal anesthesia (n=93)		General anesthesia (n=93)		Difference	95% CI	P- value*
	Mean	SD	Mean	SD			
Apgar 1	6.8	1.3	6.0	1.6	-0.8	-1.2 to -0.4	< 0.001
Apgar 5	8.6	0.6	8.3	1.2	-0.4	-0.6 to -0.1	0.010

Data are mean and standard deviation (SD).

95% CI = 95% confidence interval.

*Unpaired t test.

Table (5) expresses that there is statistically significant difference between study group as regards to apgar score at 1 minute (p<0.001) and apgar score at 5 minute(p=0.010) with high mean among spinal anesthesia group.



Fig (8): Apgar score at 1 minute and 5 minutes in both study groups.

Horizontal line (black) represents the mean. Error bars (green) represent the standard error (SE). Markers represent individual observations.

Adverse	Spinal a (n	nnesthesia =93)	General a (n=	anesthesia =93)	χ2(df=1)	P-value*	
outcome	N	%	n	%			
Low Apgar 1	31	33%	53	57%	10.507	0.001	
Low Apgar 5	1	1.1%	2	2.2%	-	1.000#	
Nausea	8	8.6%	13	14.0%	1.342	0.247	
Vomiting	2	2.2%	1	1.1%	-	1.000#	
Headache	23	24.7%	14	15.1%	2.733	0.098	
Shivering	44	47.3%	29	31.2%	5.073	0.024	
Pain requiring analgesic	91	97.8%	93	100%	-	0.497#	

 Table (6): Incidence of maternal and fetal adverse outcomes in both study groups

Data are number (n) and percentage (%).

 $\chi 2$ = chi-squared statistic, df = degree of freedom.

*Pearson Chi-squared test unless otherwise indicated.

#Fisher's exact test.

Table (6) illustrates that there is statistically significant difference with p-value (0.001) as regards to incidence of low apgar score at 1 minute with high mean among general anesthesia group.

Table (6) indicates that there is statistically significant difference between both group with p- value (0.024) as regards to the incidence of shivering with high mean among spinal anesthesia group, whereas there is no statistical difference regarding the incidence of nausea (p=0.247), vomiting (p=1.000), headache (p=0.098) and pain requiring analgesic (p=0.497).



Fig (9): Incidence of maternal and fetal adverse outcomes in both study groups.

Table	(7):	Risk	analysis	for	the	Incidence	of
adverse	e outc	omes	in both stu	udy g	group	os.	

Outcome	RR	95% CI	z	P-value	NNT / NNH	95% CI
Low Apgar 1	0.58	0.42 to 0.82	3.1 16	0.002	4.2	2.7 (Benefit) to 10.2 (Benefit)
Low Apgar 5	0.50	0.05 to 5.42	0.5 70	0.569	93.0	39.3 (Harm) to 21.3 (Benefit)
Nausea	0.62	0.27 to 1.41	1.1 43	0.253	18.6	27.1 (Harm) to 6.9 (Benefit)
Vomiting	2.00	0.18 to 21.68	0.5 70	0.569	93.0*	21.3 (Harm) to 39.3 (Benefit)
Headache	1.64	0.90 to 2.99	1.6 24	0.104	10.3*	4.7 (Harm) to 58.4 (Benefit)
Shivering	1.52	1.05 to 2.20	2.2 06	0.027	6.2*	43.7 (Harm) to 3.3 (Harm)
Pain requiring analgesic	0.98	0.95 to 1.01	1.4 14	0.157	46.5*	19.6 (Benefit) to 125.4 (Harm)

RR = relative risk, 95% CI = 95% confidence interval, Z = z-statistic, NNT = number needed to treat (or to benefit), NNH = number needed to harm.*NNH.

Table (7): illustrates that spinal anesthesia is associated with significantly lower risk for low (<7) Apgar score at 1 minute (RR=0.58, 95%CI =0.42-0.82, P=0.002), but it is associated with significantly higher risk for shivering (RR=1.52, 95% CI=1.05-2.20, P=0.027).

DISCUSSION

Delivery by caesarean section is by far one of the most commonly performed operations all over the world. Approximately 18.5 million caesarean sections are performed yearly worldwide⁽¹⁾.

This operation requires effective anaesthesia which can be regional (epidural or spinal) or a general anaesthesia. The type of anaesthesia used and the care with which it is administered is an important determinant of the outcome of caesarean section ⁽⁴⁾.

Both spinal and general anaesthesia used for caesarean section have certain advantages and disadvantages and there is no method which is completely ideal. The most important factors for choice of anaesthesia are: systemic problems, the urgency of the operation, the surgeon and the anaesthetists experience and wishes⁽⁵⁾.

This prospective randomized controlled study compared between general and spinal anaesthesia regarding the maternal and fetal outcomes after caesarean section. In this study there was no demographic difference between women in both groups regarding the mean age, BMI, parity and indication of caesarean section (p>0.05).

Also there was no significant difference between both groups as regard to preoperative blood pressure, heart rate and temperature (p>0.05).

This study showed non significant difference between both groups as regard to preoperative haemoglobin levels as p=0.586; the mean preoperative haemoglobin levels was 11.3 ± 1.2 g/dl in spinal anaesthesia group vs. 11.2 ± 1.3 g/dl in general anaesthesia group.

As regard to the postoperative haemoglobin there was a significant difference between both groups (p<0.0001), there was less reduction in haemoglobin level in spinal group than general group (the mean postoperative haemoglobin level was 10.4 ± 1.1 g/dl vs. 9.4 ± 1.5 g/dl respectively). There was also a significant difference between both groups as regard to the absolute drop in haemoglobin (g/dl) as p 0.002, there was less drop in spinal group compared to the general group (the mean absolute drop in haemoglobin level was 0.91±0.66g/dl vs. 1.29±0.95 g/dl respectively).

In this study, there was no significant difference between both groups as regard preoperative haematocrit values (p=0.352), the mean preoperative haematocrit values were $35.1\pm3.5\%$ in spinal group vs. $34.6\pm3.8\%$ in the general group.

regard the postoperative to As haematocrit values, there were significant differences between both groups (p<0.001); there was less drop in haematocrit values in spinal anaesthesia group than general anaesthesia group (the mean haematocrit values was 32±3.2% vs. 30.1±4 respectively). There was no significant difference between both groups as regard to the absolute drop in haematocrit values (%) as p=0.133.

The results of the present study goes with those of the previous study conducted by *Ezzatalsadat et al.*⁽¹⁸⁾ which showed that mean loss of haemoglobin in spinal group was

significantly lower than in general group (p=0.017). Mean loss of haematocrit in spinal group was significantly lower than in general group (p=0.035).

Another study conducted by *Marzouni et al.*⁽¹⁹⁾showed that the amount of decrease in haemoglobin and haematocrit level after caesarean section in parturients who were undergoing general anaesthesia, significantly higher than those who were undergoing spinal anaesthesia. According to the result of this study, the amount of decrease in haemoglobin and haematocrit in the general anaesthesia group was $0.8\pm0.03g/dl$ and 4.4 ± 2.2 % and the amount of decrease in the spinal anaesthesia group 0.67±0.1 g/dl and4±0.6% (p=0.002 and CI=95%).

Regarding to the amount of estimated blood loss, a statistical significance between the two groups was found (p=0.035) in which spinal anaesthesia group had less estimated blood loss than general anaesthesia group. The amount of estimated blood loss in spinal group declared a mean value of 411.6 ± 238.3 ml vs. 501 ± 329.7 ml in general anaesthesia group.

This conclusion agrees with the result of *Jeong et al.*⁽²⁰⁾ who showed that women underwent caesarean section under spinal anaesthesia had lost blood less than those under general anaesthesia (1.160 ± 710 ml vs. 1.230 ± 650 ml respectively).

This study showed that the parturient who received spinal anaesthesia had a significant longer time interval to first analgesic request. The mean time interval for the first analgesic request was longer in spinal group than general group (2.4 ± 1.8 hours vs 1.7 ± 1.2 respectively). There was a significant difference (p=0.002).

The results of the present study confirm those found in previous randomized controlled trial conducted by *Lada and Adriana*⁽²¹⁾ who stated that the time till first request for postoperative analgesia was longer with spinal anaesthesia than general anaesthesia as the mean time till first analgesic request with spinal anaesthesia group (n=35) was 159 minutes while with general anaesthesia group was 119 minutes.

This study showed that there was statistically significant effect of spinal anaesthesia versus general anaesthesia in term of shorter mean time interval to normal intestinal sound (6 ± 1.3 vs. 7.1 ± 1.5 hours). This agrees with the result of *Saygi et al.*⁽²²⁾study which included a total of 100 patients divided into two groups of 50 each, named the general anaesthesia and spinal anaesthesia groups, according to the route of administration of anaesthesia. The study revealed that the starting time for bowel sounds (22.08 ± 7.48 vs. 18.75 ± 9.2 ; p= 0.049) were significantly longer in the general anaesthesia group.

In this study, there was no significant difference between both groups as regard to the incidence of postoperative nausea (p=0.247). However, it was slightly more frequent in general group than spinal group (14% vs. 8.6% respectively).

As regard to the incidence of postoperative vomiting, this study showed no significant difference between both groups (p=1.000).

In this study, there was no significant difference between both groups as regard to the incidence of postoperative headache (p=0.098). However, it was slightly more frequent in the spinal group than general group (24.7% vs 15.1% respectively).

As regard to the incidence of postoperative shivering, it was more frequent in spinal group than general group (47.3% vs. 31.2% respectively) with significant difference (p=0.024).

A prospective observational study done by *Luggya et al.*⁽²³⁾ showed that 22 out of 270 patients undergoing caesarean section under spinal anaesthesia developed postoperative shivering giving prevalence of 8.15% with intraoperative hypotension and hypothermia as main associated factors. This study noted that postoperative shivering can be effectively controlled by 25 mg intravenous pethedine.

In this study there was statistically significant difference between the study groups as regards the mean of Apgar score at 1 minute (6.8 ± 1.3 in the spinal group vs. 6 ± 1.2 in the general group(P<0.001) and

significant difference between them as regards the mean of Apgar score at 5 minute $(8.6\pm0.6 \text{ in spinal} \text{group vs. } 8.3\pm1.2 \text{ in general group. } P=0.01).$

This conclusion agrees with that of *Mekonnen and Deska* ⁽²⁴⁾ who stated that the first and fifth Apgar score were better in neonates delivered under spinal anaesthesia as compared to general anaesthesia.

The present work also agrees that done by *Hogan et al.* ⁽²⁵⁾ which showed that there significantly increased risks of low apgar score if the delivery was performed under general anaesthesia rather than spinal anaesthesia.

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