



Acute postoperative pain in cesarean section and tubal ligation

Dolor agudo postoperatorio en cesárea y ligadura tubárica

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ABSTRACT

Background: Postoperative pain is a common complication in patients undergoing cesarean section and tubal ligation, affecting recovery and satisfaction. This study evaluates the intensity of postoperative pain at 2, 24, and 48 hours in patients who underwent cesarean section, cesarean with tubal ligation, or tubal ligation alone, under spinal anesthesia. **Methods:** A prospective cohort observational study was conducted in 73 patients at the Instituto Materno Infantil de Bogotá. Patients with pre-existing chronic pain or critical conditions were excluded. Postoperative pain was measured using the numeric pain scale at 2, 24, and 48 hours. Fisher's exact test was used for categorical comparisons and the Wilcoxon test with continuity correction for ordinal variables, applying Bonferroni correction for multiple comparisons. **Results:** At 2 hours postoperatively, 89% of patients reported no pain, while 20.5% experienced severe pain at 24 hours, and 16.5% reported severe pain at 48 hours. No statistically significant differences were found between pain levels at 24 and 48 hours ($p = 0.4094$). Furthermore, no significant differences were observed in pain levels between the three types of procedures (cesarean section, cesarean with tubal ligation, and tubal ligation alone) at any of the measured time points (2 hours, $p = 0.1037$; 24 hours, $p = 0.9685$; 48 hours, $p = 0.88$). **Conclusion:** Postoperative pain increased between 2 and 24 hours, remaining elevated at 48 hours, with no significant differences between procedures. The need to improve postoperative pain management regardless of the type of surgery is highlighted.

Keywords: Acute pain, pain, postoperative, cesarean section, sterilization, tubal, chronic pain.

RESUMEN

Antecedentes: El dolor postoperatorio agudo es una complicación común en pacientes sometidas a cesárea y ligadura tubárica, afectando la recuperación y la satisfacción. Este estudio evalúa la intensidad del dolor posoperatorio a las 2, 24 y 48 h en pacientes sometidas a cesárea, cesárea con ligadura tubárica, o ligadura tubárica sola, bajo anestesia subaracnoidea. **Métodos:** Se realizó un estudio observacional de cohorte prospectivo en 73 pacientes del Instituto Materno Infantil de Bogotá. Se excluyeron pacientes con dolor crónico preexistente o en estado crítico. El dolor postoperatorio se midió utilizando la escala numérica del dolor a las 2, 24 y 48 h. Para el análisis estadístico se empleó la prueba exacta de Fisher para comparaciones categóricas y la prueba de Wilcoxon con corrección de continuidad para variables ordinales, aplicando corrección de Bonferroni en comparaciones múltiples. **Resultados:** A las 2 h postoperatorias, el 89% de las pacientes no reportaron dolor, mientras que el 20,5% experimentó dolor severo a las 24 h, y el 16,5% reportó dolor severo a las 48 h. No se encontraron diferencias estadísticamente significativas entre los niveles de dolor a las 24 y 48 h ($p = 0,4094$). Además, no se observaron diferencias significativas en los niveles de dolor entre los tres tipos de procedimientos (cesárea, cesárea con ligadura tubárica, y ligadura tubárica sola) en ninguno de los momentos medidos (2 h, $p = 0,1037$; 24 h, $p = 0,9685$; 48 h, $p = 0,88$). **Conclusión:** El dolor postoperatorio aumenta entre las 2 y 24 h, manteniéndose elevado a las 48 h, sin diferencias significativas entre los procedimientos. Se destaca la necesidad de mejorar el manejo del dolor postoperatorio independientemente del tipo de cirugía.

Palabras clave: Dolor agudo, dolor postoperatorio, cesárea, esterilización tubaria, dolor crónico.

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Introduction

Postoperative pain is a major concern in the management of patients undergoing surgical procedures such as cesarean section and tubal ligation. According to reports from the World Health Organization (WHO), annually, 18.5 million cesarean sections are performed globally, of which approximately 6 million are considered unnecessary. It is believed that the cesarean section rate should not exceed 15% anywhere in the world[1].

In Colombia, in 2016, the proportion of births by cesarean section was 46.4% at the national level, with a slight decrease to 44.6% by 2020. In public health institutions (IPS), the proportion of cesarean sections increased from 26.2% in 1998 to 42.9% in 2014, while in private institutions it increased from 45.0% to 57.7% in 2013. The prevalence ratio of cesarean sections in private institutions compared to public ones was 1.57 (95% CI: 1.56-1.57)[2]. In Brazil, between 2014 and 2017, it was observed that the cesarean section rate was 80.0% in patients without prenatal care, 45.2% in those with inadequate prenatal care, 43.0% for those with adequate care, and 50.5% in the group with "adequate plus" care[3], showing similar proportions of cesarean sections reported in Latin America. Cesarean section rates above 30% in Latin America are concerning due to their association with higher perioperative morbidity and mortality[4].

High-efficacy contraceptive measures, such as tubal ligation, can significantly contribute to improving post-cesarean morbidity and mortality rates. This procedure is increasingly common among women, especially in the immediate postpartum period, particularly in those with higher parity[5]. Tubal ligation not only offers a permanent contraceptive method but also reduces the risk of complications in future pregnancies[6].

Although these surgical procedures are considered to have lower pain scores, postoperative pain for tubal ligations has shown average scores of 4.74 and for cesarean section 6.14 on the numerical pain scale[7]. This pain is associated with decreased patient satisfaction, delayed ambulation, the development of chronic pain, and increased morbidity and mortality[8].

The primary objective of this study is to evaluate the intensity of acute postoperative pain in patients undergoing cesarean section, with or without tubal ligation, under spinal anesthesia. Secondary objectives include characterizing sociodemographic, clinical, and surgical variables, as well as pain management. Additionally, the study aims to establish relationships between pain intensity (mild, moderate, or severe) and the type of procedure performed, evaluating these parameters at three key postoperative moments, up to 48 hours after the procedure.

Methods

This prospective cohort observational study was approved by the Ethics Committee of the Universidad Nacional de Colombia (act No. 011-191-17) and the Ethics and Research Committee of the Hospital Materno Infantil - Subred Centro Oriente (act No. 231 of November 27, 2017). It was conducted at the Maternal and Child Institute in Bogotá, where data were collected between November 2017 and 2018. Pregnant patients over 18 years old who underwent cesarean section with tubal

ligation, cesarean section alone, or tubal ligation alone, all under spinal anesthesia, were included. Patients in critical condition with mechanical ventilation, postoperative neurological complications, pre-existing chronic pain, or those undergoing simultaneous surgeries were excluded.

Using a statistical power of 80%, an expected correlation of 0.5, a two-tailed hypothesis, and a significance level of 0.05, a minimum sample size of 56 participants was calculated[9]. Adjusting for a 20% non-response rate, a total of 73 participants were required. A form with three categories of information was used for prospective data collection: sociodemographic, clinical, and related to the surgical procedure and anesthesia. Follow-up was conducted from admission to the operating room until 48 hours after the surgical procedure. Postoperative pain intensity was measured upon admission to the post-anesthesia care unit, at 24 and 48 hours, using the numerical pain scale[10]. The collected physical data were stored in a file under the custody of the principal investigator. A Microsoft Excel® database was built for data processing and analysis, which was performed using the R programming language (R Foundation®).

The data were analyzed using descriptive statistics. Numerical variables were presented as means and standard deviations, while nominal and qualitative variables were expressed as absolute numbers and percentages. A bivariate analysis was performed to evaluate differences in pain levels between the different procedures (cesarean section, cesarean section with tubal ligation, and tubal ligation), using Fisher's exact test[11], for categorical comparisons and the Wilcoxon test with continuity correction for ordinal variables[12]. In addition, Bonferroni correction was applied to adjust the significance values in multiple comparisons with a p-value of 0.017 for repeated measures comparison[13]. Bivariate results were considered statistically significant when the p-value was less than 0.05.

Results

In the analysis of sociodemographic and clinical data from the 73 patients, the average age was $26.5 \pm$ standard deviation (SD) of 6.2, while the average body mass index (BMI) was $26.9 \pm$ SD 3.9 kg/m². Most procedures were tubal ligation (54.8%), followed by cesarean section (24.7%) and cesarean section with tubal ligation (20.5%). 82.2% of patients had between 1 and 3 previous pregnancies, and 90.4% had had between 1 and 3 vaginal deliveries. 52.1% of patients had no history of cesarean section, while 39.7% had had between 1 and 2 previous cesarean sections. Labor before the cesarean section occurred in only 21.2% of patients (Table 1).

For pre-cesarean analgesia, only 15.1% of patients received epidural anesthesia and 9.1% received intravenous anesthesia. The indication for cesarean section was elective in 72.3% of cases, while 27.7% were emergency cesarean sections. The most commonly used anesthetic technique was spinal anesthesia (98.6%), and in terms of neuroaxial opioids, an average of 19.2 ± 22.3 mcg of fentanyl and 54.5 ± 51.4 mcg of morphine were administered. The average surgical time was 32.2 ± 18 minutes, and intraoperative complications (hypotension, nausea, and vomiting) occurred in 6.8% of cases.

For intraoperative analgesia, the most frequently administered intravenous drug was diclofenac, which was used in

Table1. Sociodemographic and Clinical Characterization

Characteristic	n = 73 (%)
Age (years) ± SD	26.5 ± 6.2
Weight (kg) ± SD	67.1 ± 10.7
Height (meters) ± SD	1.58 ± 0.06
BMI (kg/m ²) ± SD	26.9 ± 3.9
Procedure	
- Cesarean section	18 (24.7)
- Cesarean section and Tubal Ligation	15 (20.5)
- Tubal Ligation	40 (54.8)
Previous pregnancies	
- 1-3	60 (82.2)
- ≥ 4	13 (17.8)
Previous vaginal deliveries	
- 1-3	66 (90.4)
- 4	7 (9.6)
Previous cesarean sections	
- 0	38 (52.1)
- 1-2	29 (39.7)
- ≥ 3	6 (8.2)
Labor prior to cesarean section	7 (21.2)
Pre-cesarean analgesia	
- Epidural	5 (15.1)
- Intravenous	3 (9.1)
Indication for cesarean section	
- Elective	24 (72.3)
- Emergency	9 (27.7)
Anesthetic technique	
- Spinal	72 (98.6)
- Epidural	1 (1.4)
Neuroaxial opioid	
- Fentanyl (mcg) ± SD	19.2 ± 22.3
- Morphine (mcg) ± SD	54.5 ± 51.4
Surgical time (minutes) ± SD	32.2 ± 18
Complications (Hypotension, nausea, and vomiting)	5 (6.8)
Intraoperative analgesia	
- Diclofenac	50 (68.5)
- Dipyrrone	11 (15.1)
Immediate postoperative analgesia	
- Diclofenac	1 (1.4)
- Dipyrrone	32 (43.8)
Post-anesthesia care unit pain (2 hours) (NRS)	
Mean: 0.37 SD 1.33 Median 0 (IQR 0-0)	
- No pain (0)	65 (89.0)
- Mild (1-3)	5 (6.9)
- Moderate (4-6)	2 (2.7)
- Severe (7-10)	1 (1.4)
Pain at 24 hours (NRS)	
Mean: 4.03 SD 2.73 Median 4 (IQR 3-5)	
- No pain (0)	10 (13.7)
- Mild (1-3)	25 (34.3)
- Moderate (4-6)	23 (31.5)
- Severe (7-10)	15 (20.5)
Pain at 48 hours (NRS)	
Mean: 3.85 SD 2.32 Median 4 (IQR 2-5)	
- No pain (0)	9 (12.3)
- Mild (1-3)	25 (34.3)
- Moderate (4-6)	27 (36.9)
- Severe (7-10)	12 (16.5)

SD: Standard Deviation; BMI: Body Mass Index; NRS: Numeric Rating Scale for pain; IQR: Interquartile Range.

68.5% of patients, while 15.1% received dipyrrone. In immediate postoperative analgesia, only 1.4% received diclofenac, and 43.8% received dipyrrone. In terms of postoperative pain, 89% of patients reported no pain at 2 hours. At 24 hours, 13.7% of patients had no pain, and 20.5% had severe pain. At 48 hours, a similar distribution was observed, with only 12.3% having no pain, while 16.5% had severe pain.

In the bivariate analysis of pain levels in the post-anesthesia care unit (PACU) at 2 hours postoperatively, no statistically significant differences were found between the three types of procedures (cesarean section, cesarean section with ligation, and tubal ligation alone) at any of the observation times: 2, 24, and 48 hours (Table 2).

When comparing pain measurements at 2, 24, and 48 hours postoperatively, the Wilcoxon test with continuity correction was used, and applying the Bonferroni correction ($p < 0.017$), statistically significant differences were observed between pain levels at 2 hours and 24 hours ($p < 0.01$), as well as between 2 hours and 48 hours ($p < 0.01$). No significant differences were found in pain levels between 24 and 48 hours ($p = 0.4094$) (Figure 1).

Discussion

This study showed that acute postoperative pain in the first 48 hours after the procedure does not vary significantly among patients undergoing cesarean section, cesarean section with tubal ligation, and tubal ligation alone. Two hours postoperatively, 89% of patients did not report pain, whereas at 24 hours, this percentage decreased to 13.7%, and 20.5% of patients reported severe pain. At 48 hours, only 12.3% of patients did not report pain, while 16.5% still experienced severe

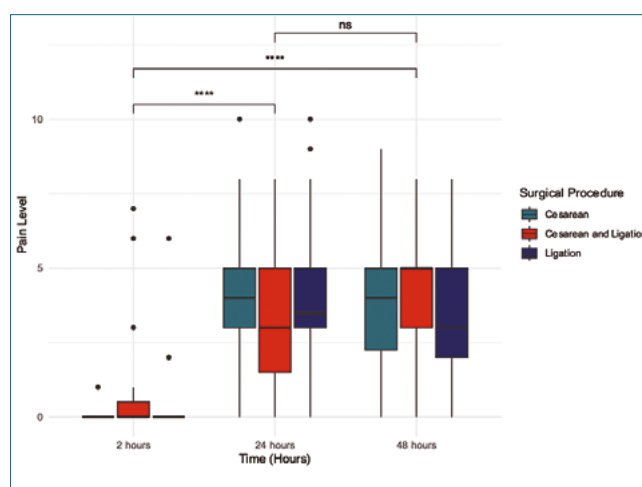


Figure 1. Distribution of pain at 2, 24, and 48 hours by procedure. Comparisons of pain levels at 2, 24, and 48 hours using the Wilcoxon test with continuity correction. Bonferroni correction ($p < 0.0167$) was applied to adjust the significance values with 3 comparisons. **** The results showed statistically significant differences between pain levels at 2 hours and 24 hours ($p < 0.01$) and between 2 hours and 48 hours ($p < 0.01$). Ns: No significant difference was found between pain levels at 24 and 48 hours ($p = 0.4094$).

Table 2. Bivariate analysis of pain and procedure

Type of Procedure (Mean ± SD)	Pain in PACU				Value p
2 hours postoperatively					
	No Pain	Mild	Moderado	Severo	0.1037*
Cesarean section (0.1± 0.32)	16 (88.9)	2 (11.1)	0 (0)	0 (0)	
Cesarean section and Tubal Ligation (1.1 ± 2.32)	11 (73.3)	2 (13.3)	1 (6.7)	1 (6.7)	
Ligadura de trompas (0.2± 0.99)	38 (95)	1 (2.5)	1 (2.5)	0 (0)	
24 hours postoperatively					
Cesarean section (4.2 ± 2.67)	2 (11.1)	5 (27.7)	7 (38.9)	4 (22.2)	0.9685*
Cesarean section and Tubal Ligation (3.5 ± 2.77)	3 (20)	5 (33.3)	4 (26.7)	3 (20)	
Tubal Ligation (4.1 ± 2.78)	5 (12.5)	15 (37.5)	12 (30)	8 (20)	
48 hours postoperatively					
Cesarean section (3.9±2.46)	2 (11.1)	6 (33.3)	7 (38,9)	3 (16,7)	0.88*
Cesarean section and Tubal Ligation (4.2± 1.98)	1 (6.7)	4 (26.7)	8 (53,3)	2 (13,3)	
Tubal Ligation (3.6 ± 2.39)	6 (15)	15 (37.5)	12 (30)	7 (17,5)	

PACU: Post-anesthesia care unit; *: Fisher's exact test; SD: Standard deviation.

pain. These findings are consistent with previous studies indicating that postoperative pain in cesarean sections is a critical factor affecting recovery and patient satisfaction[14].

The bivariate analysis of pain levels between the three procedures showed that, at 2 hours, there were no statistically significant differences between the groups ($p = 0.1037$), reinforcing the idea that tubal ligation is not a procedure exempt from relevant postoperative pain. At 24 hours, no significant differences were found in pain levels among the three procedures ($p = 0.9685$), with mean pain scores of 4.2 for cesarean section, 3.5 for cesarean section with ligation, and 4.1 for tubal ligation. These data highlight the need to improve pain management strategies in all patients undergoing tubal ligation[15]. This suggests that postoperative pain does not depend on the procedure but on the time elapsed after surgery, which is consistent with existing literature on the evolution of postoperative pain in patients undergoing cesarean section and tubal ligation[16],[17].

This study presents some limitations that should be considered when interpreting the results. The relatively small sample size ($n = 73$) could affect the generalization of the results. However, the sample size calculation was adequate to maintain the accepted statistical power, aiming to reduce alpha and beta errors, thereby increasing the validity of the findings within this specific context. Another important limitation was the lack of long-term follow-up to assess the incidence of chronic postoperative pain, which is known to affect a significant percentage of women undergoing cesarean section. Future studies should

explore different follow-up periods for patients undergoing tubal ligation to establish its association with potential persistent postoperative pain, with a larger sample size that could detect a statistical difference.

Conclusion

Acute postoperative pain did not show significant differences between cesarean section, cesarean section with ligation, and tubal ligation alone at 2 hours. However, there was an increase in pain intensity at 24 and 48 hours, highlighting the importance of postoperative pain management regardless of the surgical procedure.

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