Evaluation of pain during diagnostic and surgical minihysteroscopy under local anesthesia

Keywords

hysteroscopy, ketoprofen, visual analog scale, paracervical anesthesia, lignocaine

Abstract

Introduction

Hysteroscopy is currently considered as gold standard for the diagnosis and treatment of lesions in the uterine cavity and cervical canal. Currently, smaller diameter hysteroscopes are used, which enable the procedure to be performed without general anesthesia. Despite the use of smaller operating tools, some patients report pain.

Material and methods

The study included 142 patients who were divided into two groups: diagnostic hysteroscopy (86) and surgical hysteroscopy (56). Before the start of the procedure, ketoprofen intravenously and lignocaine paracervically were administered to the patient. The level of pain was measured using the visual analog scale (VAS).

Results

The average intensity of pain during the procedure in both groups was rated at 3 points (3.03 ± 2.25 points). There was no statistically significant difference between the two groups (VAS score 2.85 ± 2.15 vs 3.3 ± 2.38). However it was noted that as the duration of the procedure increased (regardless of the type), the level of pain experienced increased.

Conclusions

Minihysteroscopy under local, paracervical anesthesia using lignocaine, with prior administration (approximately 30 minutes before surgery) of 100 mg ketoprofen, seems to be an optimal approach for perioperative pain management. The results of the study suggest that hysteroscopes with a smaller diameter and paracervical block can be successfully used in outpatient medical practice.

Introduction

Hysteroscopy is a common endoscopic procedure performed under general and local anesthesia. It is currently considered as gold standard for the diagnosis and treatment e.g. abnormal uterine bleeding, endometrial polyps, or submucous myomas [1-6]. As hysteroscopy allows direct visualization as well as treatment of lesions, it is an ideal alternative to the procedures of uterine cavity curettage performed under general anesthesia [7]. Initially, hysteroscopy was carried out using an endoscope with a diameter of 10 mm or more, a speculum, tenaculum forceps, and dilators, which enabled the dilation of the cervical canal [8]. However, due to accompanying pain, this procedure required general anesthesia [8,9]. Nowadays, resectoscopes with a reduced diameter are widely used for hysteroscopy. The advent of modern minihysteroscopes, which do not necessitate the dilatation of the cervical canal, has made it possible to perform hysteroscopy with only local anesthesia rather than short-term general anesthesia [10,11]. Hysteroscopy under local anesthesia offers significant advantages over the procedure performed under general anesthesia e.g. a reduction in the risk associated with general anesthesia, decreased costs, and shorter hospital stay [12,13]. The use of hysteroscopes with smaller diameters does not guarantee a painless procedure [14] also surgery under local anesthesia is not always successful due to increased pain and risk of vasovagal reaction [8,15]. Data from the literature indicate that 1.3–5.2% of procedures cannot be performed due to pain intolerance [16]. The severity of pain may be determined not only by technical aspects but also by the experience of the surgeon, duration of the procedure, abnormalities of the genital tract, and psychological profile of the patient [1].

Studies on major gynecological procedures often use the Visual Analogue Scale (VAS) [17]. VAS is a proven and sensitive tool for the assessment of pain. It avoids vague descriptions of pain that can be difficult for a patient to understand and also allows a comparison of

measurements over time [19]. However, only a few studies have examined the level of pain experienced during hysteroscopy under local anesthesia, and thus, it is important to investigate this aspect. With increasing availability of ultrasound examination, the uterine cavity abnormalities are being increasingly diagnosed, and hence there is a need to determine the best treatment option for patients [18].

Aim

The aim of this study was to assess the level of pain experienced by patients during minihysteroscopy under local anesthesia, with the use of VAS.

Materials and methods

The study included patients who were undergoing diagnostic and surgical minihysteroscopy in the Gynecological-Obstetrical Clinical Hospital of Poznan University of Medical Sciences, during 2020–2021.

Procedure of qualifying patients for surgery

The eligibility criteria for minihysteroscopy procedures under local anesthesia (with lignocaine) included abnormal uterine bleeding, endometrial polyps or submucous myomas, and endometrial hyperplasia. Furthermore, the patients included in the study were in the first phase of the menstrual cycle or in the postmenopausal period and qualified for surgery after gynecological and ultrasound examination. All women reported good health without cardiovascular diseases and negative history of lignocaine and ketoprofen allergy. Those diagnosed with heavy genital bleeding, vaginal and/or cervical inflammation, and inability to exclude pregnancy during qualification for the procedure were excluded.

Data collecting

A medical history was collected during the qualifying examination, and the patients were asked about their age, body weight and height, history of operations, allergies, number and type of deliveries, number of miscarriages, previous cervical and endometrial procedures, and general health status.

Pain was assessed using VAS. During qualification, before the procedure, the principles of VAS pain assessment were explained to each examined woman. After the procedure, the patients were asked to indicate their pain severity based on VAS. VAS was used to assess the pain experienced during the procedure on a scale of 0–10. The pain level was classified as follows: 0 points—no pain, 1–3 points—mild pain, 4–7 points—moderate pain, and 8–10 points—severe pain [19].

Surgery procedures

After gynecological examination and transvaginal ultrasound, signed consent for the hysteroscopic procedure was obtained from each patient. Thirty minutes before the start of the procedure, 100 mg ketoprofen was intravenously administered to the patient. During the procedure, vital functions, such as heart rate, blood pressure, O₂ saturation, and respiratory rate, were monitored. Ten minutes before the minihysteroscope was inserted into the cervical canal, 10 ml of 0.1% lignocaine solution was paracervically administered in two injections (the first 10 ml at 4 o'clock and the second 10 ml at 8 o'clock) at a needle insertion depth of about 2 mm. Hysteroscopy was performed without inserting tenaculum forceps and speculum in the vagina.

The operators who had experience in hysteroscopy performed the procedure. The patients were first placed on a "treatment table" in a position used for gynecological examination. The uterine cavity was dilated with 0.9% NaCl solution administered with a

continuous flow at a pressure of 110 mmHg. Hysteroscopy was performed using the GUBBINI Mini Hystero-Resectoscope system with a working diameter of 16 Fr. Being a certified center and working on the Gubini system, we adapted the procedure of pericervical anesthesia to Polish conditions, using lignocaine (a preparation with a less cardiotoxic effect than

bupivacaine) and ketoprofen.

Based on the reported symptoms and abnormalities found on ultrasound examination and the type of procedure performed, the patients were divided into two groups: diagnostic hysteroscopy (HD) and surgical hysteroscopy (HO). The HD group consisted of women (n=86) aged 46.29 ± 9.55 years (median (Me): 45 years), who reported abnormally heavy uterine bleeding. These patients had an endometrial biopsy before the planned laparoscopic supracervical uterine removal surgery. During the procedure, 3 pieces of endometrium with a volume up to 0,5 ml were collected. The HO group consisted of women (n=56) aged 45.38 ± 13.04 years (Me: 43 years), with ultrasound-confirmed diagnosis of endometrial polyps or submucous myomas. The diameter of the lesions within the uterine cavity ranged from 0.5 cm to 1.5 cm. During the procedure, the patients were conscious and could observe the procedure in real time on a monitor.

Statistical analysis

Data analysis was carried out using Statistica (TIBCO Software Inc. 2017, data analysis software system, version 13) and Microsoft Excel (version 2019, Microsoft Office). The distribution of variables was assessed using the Shapiro–Wilk test. Homogeneity of variance was assessed using Levene's and Brown–Forsythe tests. Student's *t*-test, Mann–Whitney *U* test, and Kruskal–Wallis analysis of variance were used for comparisons between groups. The level of significance was set at p<0.05 for all calculations.

Results

Characteristics of the study group

A total of 142 women were included in the study. The age of the subjects ranged from 22 to 80 years, and their average age was 44 years (45.93 \pm 11.02). The body mass index (BMI) values of the subjects ranged from 17.72 to 41.98 kg/m², and their mean BMI was 24.97 kg/m² (26.14 \pm 5.28). No statistically significant differences were noted between the type of hysteroscopic procedure performed and the patient age, BMI, body mass score, or presence of comorbidities (Table I).

Among the studied patients, 90.7% in the HD group and 67.86% in the HO group declared obstetric history of pregnancy. Women in the HD group were characterized by a higher number of past pregnancies (p<0.001). A total of 25 subjects (17.61%) declared a history of miscarriage, including 17 (19.77%) in the HD group and 8 (14.29%) in the HO group. Sixty-one (70.93%) patients in the HD group and 32 (57.14%) in the HO group declared natural childbirth. No correlation was observed between the type of minihysteroscopy procedure performed and the number of natural deliveries. Among the studied subjects, 29 (20.42%) declared delivery by cesarean section. The patients in the HD group reported a significantly higher number of deliveries by cesarean section (24 vs 5; p=0.006). The obstetric history indicated that women in the HO group had significantly fewer offspring (p=0.001) than those in the HD group (Table II).

Hysteroscopy

The procedures lasted from 13 to 37 minutes. The mean duration of hysteroscopy was 25 minutes (23.54 \pm 5.38). There were no significant differences in the procedure time based

on the type of hysteroscopy. No association was found between the procedure time and patient BMI or age. However, it was noted that in the HD group, a shorter procedure time was associated with a higher number of natural deliveries (τ =-0.15; *p*=0.035). In contrast, in the HO group, a higher number of miscarriages was associated with a shorter procedure time (τ =-0.27; *p*=0.003). Among the studied women, 26 (18.31%) had previously undergone minor gynecological procedures (uterine cavity curettage), including 15 (17.44%) in the HD group and 11 (19.64%) in the HO group. No significant relationship was observed between the type of minihysteroscopy and the history and/or number of minor gynecological procedures (Table III).

VAS

The average intensity of pain during the procedure was rated at 3 points (3.03 ± 2.25 points) (Figure 1). No differences in pain scores were found with respect to the type of minihysteroscopy procedure performed (Table III). During the assessment of pain after surgery, 21 (14.8%) patients reported no pain, while mild pain (1–3 points) was reported by 65 (45.8%), moderate pain (4–7 points) by 49 (34.5%), and severe pain (8–10 points) by 7 (4.9%). No statistically significant relationship between the VAS score and BMI or age of the subjects was noted. In the HO group, a higher number of deliveries by cesarean section was associated with a lower pain score (τ =–0.2; p=0.03). It was also noted that as the duration of the procedure increased (regardless of the type), the level of pain experienced increased (r=0.17; p=0.0495).

Discussion

Hysteroscopy has been the gold standard for evaluating the pathology of the cervical canal and uterine cavity in both premenopausal and postmenopausal patients. Minihysteroscopy, performed using miniature hysteroscopic equipment, shortens the recovery time and effectively integrates clinical practice with a "see-and-treat" mode, while avoiding the risk of anesthesia and inconvenience of the operating room [20]. Many efforts have been made in the past 20 years to promote the wide use of the minihysteroscopy procedure. However, minihysteroscopy is still considered painful, with some women reporting discomfort at various stages of the procedure [21]. In addition, unpleasant sensations may be compounded by high levels of anxiety about the procedure [21]. It has been shown that pain is the most common reason for the discontinuation of the surgical procedure [22]. For this reason, pharmacological (administration of nonsteroidal anti-inflammatory drugs (NSAIDs), cyclooxygenase-2 inhibitors, antispasmodics, local anesthetics, prostaglandins, and opioids) and nonpharmacological (percutaneous electrical nerve stimulation, uterine relaxation, adequate fluid pressure during uterine entry, heating and type of medium used for the procedure, hypnosis, music, talking to the patient) approaches of intraoperative and postoperative pain relief have been used in women undergoing minihysteroscopy, with results varying in different groups [22]. Abis et al. [23] reviewed the different types of pain relief methods available. Their study indicated that routine pre- or intraoperative analgesia was used by 34% of surgeons (67% of them used NSAIDs, 12% paracetamol, 7% opioids, 13.5% other). Among the operators who routinely used intraoperative analgesia, 46.1% used paracervical block and 15.4% used intracervical block, 3.8% used local anesthesia (spray or gel on the cervical surface), and 19.2% used another method, while 15.4% encouraged patients to listen to music during the surgery. Regarding misoprostol, 75% of hysteroscopists reported that they do not use it routinely [23]. In our study, we compared the level of pain reported by patients undergoing diagnostic and surgical minihysteroscopy procedures under local, paracervical anesthesia using lignocaine with prior administration (approximately 30 minutes before surgery) of 100 mg ketoprofen. The mean pain reported by female patients in this study based on VAS was 3, which implies that the pain perceived was mild or moderate pain. The rest of the patients in the study reported 4 or more points on the scale, which may support the hypothesis that hysteroscopy may be moderately painful in some patients. Similar results were obtained in the study by Pegoraro et al. [24], in which 41.7% of patients who underwent hysteroscopy for infertility diagnosis rated pain at 4 or more points [24] (Table IV). Other studies of mixed populations (with different indications for hysteroscopy) demonstrated that a wide range of women (21-88%) reported pain at 4 points on VAS, with a mean pain score of 1.8–6.2 points. The pain values may depend, among others, on reproductive status, operator experience, dilating medium, ethnicity, patient stress, and diameter of surgical instruments [14,18,25-27] (Table IV). According to De Silva [28], saline should be administered at the lowest pressure during minihysteroscopy to dilate the uterus for a satisfactory view and pain reduction [28]. Pain induced by the passage of minihysteroscope through the cervical canal is also associated with cervical dilation. Women who had natural deliveries have a more dilated cervix, and thus experience less pain during the procedure and the procedure time is also shorter [27],[29]. In our study, we observed that in the HD group, a shorter procedure time was associated with a higher number of natural deliveries in the history, whereas in the HO group, a higher number of miscarriages was associated with a shorter procedure time. In turn, a shorter procedure time was associated with a lower VAS score. Samy et al. [30] demonstrated that the use of lidocaine and tramadol resulted in effective pain reduction in postmenopausal patients during and after diagnostic minihysteroscopy. However, after lidocaine infusion into the uterine cavity, the timing of the minihysteroscope passage through the cervical canal was associated with less pain as assessed by VAS [30] (Table IV). In a similar study, Hassan obtained similar VAS results in the groups of patients in which tramadol and placebo were used [31]. In the study of Al-Sunaidi, the VAS values were found at the level of 2.1 and 3.2 with the use of lorazepam *p. o.*, intra- and paracervical anesthesia with bupivacaine [32] (Table IV). According to Ying Law, listening to music may influence pain assessment by VAS [33]. In a study on the Chinese population, the author examined whether listening to music during hysteroscopy reduced pain. The results indicated that music reduced pain by 1.3 points, and therefore routine use of music during hysteroscopy can improve patient care [33]. These results are not confirmed by Mak earlier work, in which he indicates that listening to music may be a factor distracting the operator [34] (Table IV).

Other studies have revealed that vasovagal reactions occur during hysteroscopy. Coimbra [35] observed vasovagal reactions such as bradycardia, hypotension, syncope, sweating, nausea, or vomiting during or after the procedure [35]. In our study, we did not record such a reaction during the our procedure. The occurrence of vasovagal reaction may be associated with the use of perioperative anesthesia with lidocaine, which has been confirmed by Zupi [36] and Elasy et al. [37].

Besides pain, another frequently studied factor is the anxiety level of patients, which was not assessed in this study. Anxiety before surgery is assessed using the Illness Perception Questionnaire [29,38,39]. Anxiety, in addition to pain, may hinder the procedure and thus translate into a negative patient experience [29]. The factors associated with pain during hysteroscopy have not been studied in detail so far [40]. There is also consistent evidence clearly indicating the safest and most effective pain relief option for women undergoing hysteroscopy [41]. Therefore, efforts should be made to help identify those who are at a higher risk of experiencing pain. The overarching goal is to provide adequate pain management during minihysteroscopy procedures, in order to reduce discomfort and satisfy both the patient and the operator.

The main advantages of this study at the Hysteroscopy Center under Local Anesthesia

are	
	- conducting the study on a large number of patients,
	 repeatability of treatment conditions,
	- performing treatments by a small group of qualified operators based on the same
	technique and equipment
	- using the same anesthesia for each patient.
As	a team, we are convinced that further research will lead to the popularization of this
me	thod and the introduction of standards in hysteroscopic procedures.

Conclusions

Diagnostic and surgical minihysteroscopy under local, paracervical anesthesia using lignocaine, with prior administration (approximately 30 minutes before surgery) of 100 mg ketoprofen, seems to be an acceptable approach for perioperative pain management. The results of the study suggest that hysteroscopes with a smaller diameter and paracervical block can be successfully used in outpatient medical practice.

Data availability

Data supporting the findings of this study are available from the corresponding author upon request.

Conflicts of interest

The authors declare no conflicts of interest with respect to the authorship and/or publication of this article.

Ethics statement

The methods used for patient enrollment and collection and storage of research material were approved by the Bioethics Committee at the Poznan University of Medical Sciences (Resolution No. 828/20 of 14.01.2021).

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Table I. Clinical characteristics

	Type of hysteroscopy		
Age	Diagnostic minihysteroscopy	Surgical minihysteroscopy	
Ν	86	56	
M ± SD	46.29 ± 9.55	45.38 ± 13.04	
Min–Max	27–76	22–80	
Me [Q1–Q3]	45 [41–50]	43 [37–50]	
U	2119.5		
BMI			
M ± SD	26.14 ± 5.22	26.14 ± 5.42	
Min–Max	17.72-40.23	18.96-41.98	
Me [Q1–Q3]	25.105 [22.31–28.84]	24.61 [21.62–29.38]	
U	2365		
Body weight assessment	Diagnostic	Surgical	
Underweight	1 (1.16%)	0 (0%)	
Normal	42 (48.84%)	28 (50%)	
Overweight	27 (31.4%)	15 (26.79%)	
Obesity	16 (18.6%)	13 (23.21%)	
	p=	p=0.74	
Comorbidities			
Yes	59 (68.6%)	35 (62.5%)	
No	27 (31.4%)	21 (37.5%)	
	p=0.45		

Table II. Obstetric history vs type of hysteroscopy

	Type of hysteroscopy		
Past pregnancies	Diagnostic minihysteroscopy	Surgical minihysteroscopy	
Yes	78 (90.7%)	38 (67.86%)	
No	8 (9.3%)	18 (32.14%)	
	p<0.001	!	
Natural childbirth			
Yes	61 (70.93%)	32 (57.14%)	
No	25 (29.07%)	24 (42.86%)	
	p=0.09		
Past miscarriage		~	
Yes	17 (19.77%)	8 (14.29%)	
No	69 (80.23%)	48 (85.71%)	
	p=0.4		
Past cesarean section			
Yes	24 (27.91%)	5 (8.93%)	
No	62 (72.09%)	51 (91.07%)	
	p=0.006	5	
Number of cesarean sections			
0	62 (72.09%)	51 (91.07%)	
1	17 (19.77%)	3 (5.36%)	
2	7 (8.14%)	2 (3.57%)	
	p=0.02		
Parity			

Childless	9 (10.47%)	20 (35.71%)
Primipara	27 (31.4%)	12 (21.43%)
Multipara	50 (58.14%)	24 (42.86%)
	p=0.00	01
Occurrence of preterm delivery		
Yes	3 (3.49%)	3 (5.36%)
No	83 (96.51%)	53 (94.64%)
	p=0.5	9

Table III. Previous procedures, time and VAS vs type of hysteroscopy

	Type of hysteroscopy		
Undergoing a minor gynecological procedure	Diagnostic minihysteroscopy	Surgical minihysteroscopy	
Yes	15 (17.44%)	11 (19.64%)	
No	71 (82.56%)	45 (80.36%)	
	p=0.74	l	
Number of minor gynecological procedures performed			
0	71 (82.56%)	45 (80.36%)	
1	12 (13.95%)	9 (16.07%)	
2	2 (2.33%)	1 (1.79%)	
3	1 (1.16%)	0 (0%)	
4	0 (0%)	1 (1.79%)	
	p=0.67	,	
Duration of the procedure			
M ± SD	22.8 ± 4.61	24.66 ± 6.27	
Min–Max	13–35	15–37	
Me [Q1–Q3]	22 [20–25]	25 [20–30]	
U	2006		
Pain assessment—VAS			
M ± SD	2.85 ± 2.15	3.3 ± 2.38	
Min–Max	08	0–10	
Me [Q1–Q3]	3 [1-4]	3 [1–5]	
U	2149		
	*p<0.05; **p<0.01; ***p<0.001		

Table IV. VAS results in other studies

Author	Type of hysteroscopy	Number of patients	Type of anesthesia/ pain reducting factor	VAS
Angelis	Diagnostic	102	-	2.3 ± 2.1
Pluchino	Diagnostic	42	-	2.07 ± 0.76
	Diagnostic	52	lidocaine <i>i. v.</i>	4.0 ± 2.4
Samy	2.08.0000	52	tramadol <i>p. o.</i>	4,7 ± 2,1
		52	placebo	5.8 ± 2.3
Law	Diagnostic / surgical	53	music	4.54 ± 2.89
		50	-	5.88 ± 2.9
Rolim	Diagnostic	252		5.6 ± 3.2
		39	music	5.7 ± 2.6
Mak	Diagnostic	42	-	5.2 ± 2.7
		70	tramadol p. o.	4.37 ± 1.77
Hassan	Diagnostic	70	celecoxib <i>p. o.</i>	4.63 ±1.63
		70	placebo	5.92 ± 2.26
Kokanali	Diagnostic	148	-	5.3 ± 2.8
		42	lorazepan 10 mg p.o +	3.2 ± 0.3
			bupivacainum 2 ml 0,5 % paracervical	
Al- Sunaidi	Diagnostic	42	lorazepan 10 mg p.o. + bupivacainum 2 ml 0,5 % paracervical and intracervical	2.1 ± 0.2



Figure 1 Pain level during minihysteroscopy based on VAS scores



Diagnostic and surgical minihysteroscopy under local, paracervical anesthesia using lignocaine, with prior administration (approximately 30 minutes before surgery) of 100 mg ketoprofen, seems to be an acceptable approach for perioperative pain management.

Table I. Clinical characteristics

	Type of hysteroscopy		
Age	Diagnostic minihysteroscopy	Surgical minihysteroscopy	
Ν	86	56	
$M\pm SD$	46.29 ± 9.55	45.38 ± 13.04	
Min–Max	27–76	22–80	
Me [Q1–Q3]	45 [41–50]	43 [37–50]	
U	2119.5		
BMI			
$M\pm SD$	26.14 ± 5.22	26.14 ± 5.42	
Min–Max	17.72-40.23	18.96-41.98	
Me [Q1–Q3]	25.105 [22.31–28.84]	24.61 [21.62–29.38]	
U	2365		
Body weight assessment	Diagnostic	Surgical	
Underweight	1 (1.16%)	0 (0%)	
Normal	42 (48.84%)	28 (50%)	
Overweight	27 (31.4%)	15 (26.79%)	
Obesity	16 (18.6%)	13 (23.21%)	
Significance	<i>p</i> =0.74		
Comorbidities			
Yes	59 (68.6%)	35 (62.5%)	
No	27 (31.4%)	21 (37.5%)	
Significance	p=0.45		

Table II. Obstetric history vs type of hysteroscopy

	Type of hysteroscopy		
Past pregnancies	Diagnostic minihysteroscopy	Surgical minihysteroscopy	
Yes	78 (90.7%)	38 (67.86%)	
No	8 (9.3%)	18 (32.14%)	
Significance	p<0.0	01	
Natural childbirth			
Yes	61 (70.93%)	32 (57.14%)	
No	25 (29.07%)	24 (42.86%)	
Significance	<i>p</i> =0.	09	
Past miscarriage			
Yes	17 (19.77%)	8 (14.29%)	
No	69 (80.23%)	48 (85.71%)	
Significance	<i>p=0</i>	.4	
Past cesarean section			
Yes	24 (27.91%)	5 (8.93%)	
No	62 (72.09%)	51 (91.07%)	
Significance	<i>p=0.0</i>	06	
Number of cesarean sections			
0	62 (72.09%)	51 (91.07%)	
1	17 (19.77%)	3 (5.36%)	
2	7 (8.14%)	2 (3.57%)	
Significance	p=0.02		
Parity			
Childless	9 (10.47%)	20 (35.71%)	

Primipara	27 (31.4%)	12 (21.43%)
Multipara	50 (58.14%)	24 (42.86%)
Significance	<i>p</i> =0.001	
Occurrence of preterm delivery		
Yes	3 (3.49%)	3 (5.36%)
No	83 (96.51%)	53 (94.64%)
Significance	<i>p=0.59</i>	

Table III. Previous procedures, time and VAS vs type of hysteroscopy

	Type of hysteroscopy	
Undergoing a minor gynecological procedure	Diagnostic minihysteroscopy	Surgical minihysteroscopy
Yes	15 (17.44%)	11 (19.64%)
No	71 (82.56%)	45 (80.36%)
Significance	p=0.74	4
Number of minor gynecological procedures performed		
0	71 (82.56%)	45 (80.36%)
1	12 (13.95%)	9 (16.07%)
2	2 (2.33%)	1 (1.79%)
3	1 (1.16%)	0 (0%)
4	0 (0%)	1 (1.79%)
Significance	<i>p</i> =0.67	
Duration of the procedure		
M ± SD	22.8 ± 4.61	24.66 ± 6.27
Min–Max	13–35	15–37
Me [Q1–Q3]	22 [20–25]	25 [20–30]
U	2006	·
Pain assessment—VAS		
$M \pm SD$	2.85 ± 2.15	3.3 ± 2.38
Min–Max	0-8	0-10
Me [Q1–Q3]	3 [1–4]	3 [1–5]
U	2149	
	*p<0.05; **p<0.01; ***p<0.001	

Table IV. VAS results in other studies

Author	Type of hysteroscopy	Number of patients	Type of anesthesia/ pain reducting factor	VAS
Angelis	Diagnostic	102	-	2.3 ± 2.1
Pluchino	Diagnostic	42	-	2.07 ± 0.76
	Diagnostic	52	lidocaine <i>i</i> . <i>v</i> .	4.0 ± 2.4
Samy		52	tramadol p. o.	4.7 ± 2.1
		52	placebo	5.8 ± 2.3
Law	Diagnostic / surgical	53	music	4.54 ± 2.89
		50		5.88 ± 2.9
Rolim	Diagnostic	252		5.6 ± 3.2
		39	music	5.7 ± 2.6
Mak	Diagnostic	42	-	5.2 ± 2.7
		70	tramadol p. o.	4.37 ± 1.77
Hassan	Diagnostic	70	celecoxib <i>p. o.</i>	4.63 ±1.63
		70	placebo	5.92 ± 2.26
Kokanali	Diagnostic	148	-	5.3 ± 2.8
		42	lorazepan 10 mg p.o +	3.2 ± 0.3
			bupivacainum 2 ml 0,5 % paracervical	
Al- Sunaidi	Diagnostic	42	lorazepan 10 mg p.o. +	2.1 ± 0.2
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Pain level during minihysteroscopy based on VAS scores