RESEARCH ARTICLE

Postoperative urinary retention - prevalence and risk factors: prospective, cohort study

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ABSTRACT

Introduction. Ensuring perioperative urination maintenance can often be challenging, as postoperative urinary retention is frequently overlooked in favor of more clearly defined goals such as successful surgery, comprehensive postoperative pain control, reducing the risk of postoperative cardiorespiratory complications and shortening the patient's overall hospital stay. However, the inability to initiate urination and empty the bladder in the early postoperative period may negatively affect each of the listed success criteria.

Material and methods. A single-center, prospective, observational, cohort study was conducted, enrolling elderly patients without severe comorbidities. A total of 127 complete datasets were analyzed. Anthropometric parameters, type of surgery, duration of anesthesia and surgery; and several parameters previously reported as risk factors for postoperative urinary retention were recorded. The main objective wasto identify the prevalence of postoperative urinary retention in a surgical group in the Republic of Moldova. The secondary objective was to test the predictive value of a series of parameters (modifiable and non-modifiable) related to the patient or surgical treatment received as risk factors for urinary retention in the first 24 hours postoperatively. Statistical software used: Social Science Statistics (https://www.socscistatistics.com).

Results. The studied surgical population was homogeneous in terms of body mass, height, duration of surgery and anesthesia; heterogeneous by gender (62.2% male) and type of anesthesia (64% general anesthesia). Depending on the definition criteria, the prevalence of postoperative urinary retention varied between 5.5% and 7.9%. The preoperative unmodifiable risk factors for postoperative urinary retention: positive history for hypertension OR = 9.0 (X^2 (1, N = 127) = 5.6, p = 0.017), diabetes mellitus OR = 5.1 (X^2 (1, N = 127) = 5.36, p = 0.021) and stroke OR = 4.83 (X^2 (1, N = 127) = 2.098, p = 0.148).

Conclusions. The prevalence of postoperative urinary retention in a single-center surgical population from the Republic of Moldova varies between 5.5% and 7.9%, depending on the criteria for postoperative urinary retention applied. This variation highlights the need for a consensus on diagnostic criteria for postoperative urinary retention is needed. Patients with hypertension and diabetes mellitus were more likely to develop postoperative urinary retention. Patients with pre-existing neurological disorders such as positive history for stroke and diabetic polyneuropathy were more susceptible for postoperative urinary retention.

Keywords: postoperative urinary retention, risk factors, postoperative complications.

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Key messages

What is not yet known on the issue addressed in the submitted manuscript

Currently no national studies would report the prevalence and risk factors for postoperative urinary retention.

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The research hypothesis
To identify the prevalence of postoperative urinary retention in a surgical population in the Republic of Moldova, with identification of risk factors for this complication.

The novelty added by manuscript to the already published scientific literature
Hypertension is an independent risk factor for postoperative urinary retention. Diabetes mellitus in combination with hypertension predisposes to the development of postoperative urinary retention. Patients with pre-existing neurological disorders such as a history of stroke and diabetic polyneuropathy are more susceptible to postoperative urinary retention.

Introduction

Regardless of the type of surgery, the post-operative period is a critical one, during which many complications can occur, such as acute post-operative pain, respiratory and/or hemodynamic instability, nausea and post-operative vomiting. Postoperative urinary retention (POUR) is one of the complications that can develop in the first 24 hours after surgery and often remains unrecognized and underestimated. Essentially, POUR is the inability to urinate in the presence of a full bladder in the early postoperative period. The prevalence of the complication varies widely, from 5 to 70% which indicates on the one hand a lack of consensus on clear criteria for POUR, and on the other hand confirming the multitude of risk factors related to this complication [1, 2]. Over the years, several risk factors associated with POUR have been studied and reported: unmodifiable (male [2-4], female [4-7], age [1, 2, 4, 8-10], comorbidities (diabetes mellitus (DM) [1, 2, 4], pre-existing neurological disorders [2, 4, 10]) and modifiable, related to the type of surgery (anorectal, colorectal, urogenital surgery [1, 2, 4, 6, 8]) and duration of surgical treatment (consisting of surgery under the protection of a certain type of anesthesia) [2-4, 10], intraoperative fluid volume [1, 2, 4, 6, 10], medication used (opioids [4, 8, 10], anticholinergics and antipsychotics [10]) and high American Society of Anesthesiologists score (ASA) [9], among others .

If left undiagnosed and untreated in a timely manner, POUR can lead to increased morbidity due to urinary tract infections, detrusor muscle dysfunction, arrhythmias and delirium, with prolonged hospital stay [2]. These complications associated with POUR have increased attention on early diagnosis of the phenomenon with the development of prevention strategies. Recently, preventive ultrasound diagnosis of POUR has gained significant importance [11].

The perioperative period may affect micturition with precipitation of POUR. As a spinal reflex controlled by the brainstem, the urination process is a complex one, consisting of 2 phases - the storage phase (mediated by sympathetic innervation) and the emptying phase (provided by parasympathetic fibers). The bladder itself is a container with a flexible muscular wall, which can hold an increasing volume of urine without large variations in pressure until a certain threshold is reached. On average, normal bladder capacity is 400-600 ml [10]. The first impulse to urinate occurs when the bladder volume is about 150 ml, and the sensation of fullness occurs at 300 ml volume, which, once exceeded, transmits the information through the pelvic splanchnic system, e activating emptying process by parasympathetic fibers [4]. For bladder emptying to occur, inhibition of the motor cortex must be absent and the contraction of the detrusor muscle. Motor cortex disinhibition is triggered by pudendal afferents as soon as urine enters the posterior urethra. As a result, relaxation of the pelvic floor muscle, descent of the levator ani muscle and voiding of urine occurs [1, 4, 10].

Surgical and intra-anesthetic stressor factors, acute postoperative pain, medications used, and patient comorbidities can complexly interfere with the physiological pathways of micturition, resulting in POUR [3, 5, 8, 10]. For instance, opioids may alleviate the sensation of bladder fullness through parasympathetic inhibition and increased sphincter tone due to increased sympathetic activity [4, 5, 10]. Beyond smooth muscle relaxation with reduced bladder contractility, multimodal general anesthesia also predisposes to POUR by causing autonomic bladder tone dysregulation [3-5]. Neuraxial techniques interfere with both the afferent and efferent pathways of micturition, with the prevalence of POUR being directly proportional to the duration of action of the local anesthetic molecule introduced intrathecally [2-5, 10]. Neuraxial opioids are associated with an increased prevalence of POUR compared with their intravenous administration [4].

The study aimed to estimate the prevalence of POUR in a medical institution with a surgical profile from Moldova and to identify a series of risk factors.

Material and methods

Study design and parameters
The prospective, observational, monocentric, cohort study, designed to elucidate the prevalence and characteristics of POUR in a medical institution in Moldova, was conducted from June 1, 2022, to December 31, 2022, at *Valeriu Ghereg* Anesthesiology and Resuscitation Department No.1, and the Institute of Emergency Medicine. All included patients signed informed consent to participate in the study.

Participants

A number of surgical patients were evaluated weekly according to eligibility criteria. The inclusion criteria were age > 18 years, signed informed consent; undergoing general, neuroaxial, locoregional or combined type of anesthesia; surgery lasting > 30 minutes, ASA score I - III. Patients were excluded if they expressed their wish to leave the study, positive history of renal failure, benign prostatic hypertrophy with obstruction, required bladder catheterization from the beginning of the surgery or lasting surgery > 3 hours or were scheduled for ambulatory surgery.

The Consort flow chart with the pattern of refusals, enrolment, and follow-up in the first 24 hours postoperatively is shown in Figure 1.

After obtaining informed consent to participate, the study questionnaire, developed based on the literature review, was completed with the patient's personal and clinical data regarding risk factors for POUR. The questionnaire included 3 compartments: general patient data (comorbidities, chronic medication, surgical profile), data related to peri-anesthesic management and a stipulation of study's inclusion and exclusion criteria. The time points of recording the parameters are represented in the study diagram (Figure 2).
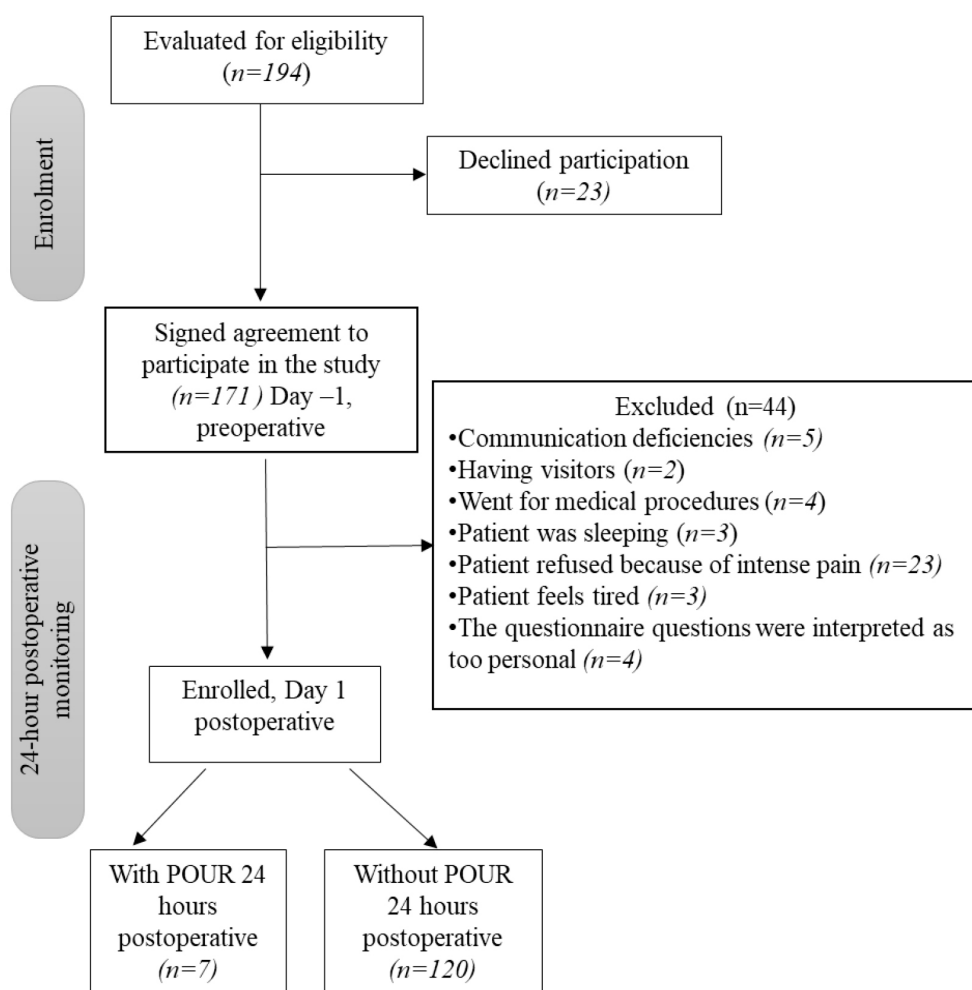

Fig. 1 CONSORT flowchart: refusal scheme, enrolment, postoperative follow-up
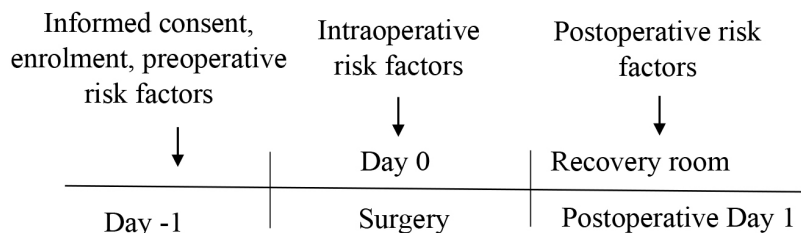**Note**: *POUR – postoperative urinary retention.*

Fig. 2 Study design diagram.

Recorded parameters and statistical analysis

The following general parameters were recorded: age, sex, height, body mass, type of surgery, duration of anesthesia, duration of surgery; and parameters tested as risk factors: preoperative, intraoperative and postoperative factors, directly related to patient or the medical act itself and the medication administered - their nominal detail are presented in the results. The values of the parameters were numerated via de-identification in an Excel table and then imported into the statistical analysis software Social Science Statistics (https://www.socscistatistics.com). Data are presented as absolute and relative values or mean and standard deviation. The Fisher exact test, the hi-squared independence test (X^2) including Yates correction, odds ratio (OR) were calculated, and a $p<0.05$ was considered statistically significant.

All respondents agreed to participate in the research and signed the informed consent for participation in the study, as well as the informed consents for hospitalization, surgery and anesthesia. The present study observes individuals and measures variables of interest but does not attempt to influence the treatment, lacking randomization of exposure. Also, the observational data are reported in grouped format (counts and percentages), ensuring that participants cannot be identified from the results. The nature of the data reported in the manuscript is not of a sensitive nature, and the study is generally considered a low-risk project.

Results

The general characterization of the 127 patients enrolled in the study is presented in Table 1. The mean age of the patients was 47.0±19 years (ranging from 18 to 83 years). The mean age of patients who did not develop POUR versus those who developed POUR was 45.8±18.6 and 60.0±10.9 (p = 0.951) years, respectively. The surgical population studied was homogeneous in terms of anthropometric criteria and heterogeneous in terms of gender distribution, with a predominance of males.

Table 1. General characterization of patients enrolled in the study.

Parameter	Enrolled (n = 127)	Non-POUR (n = 120)	POUR* (n = 7)	No difference
Age, years	47.0±19 (18 – 83)	45.8±18.6 (18 – 83)	60.0±10,9 (42 – 70)	
18 – 24	19 (15.0 %)	19(15.8 %)	0 (0 %)	
25 – 44	41 (32.3 %)	40 (33.4 %)	1 (14.3%)	p* = 0.951
45 – 64	35 (27.5 %)	33 (27.5 %)	2 (28.6%)	
>65	32 (25.2 %)	28 (23.3 %)	4 (57.1 %)	
Men/ Women	79 (62.2 %)/ 48/ (37.8 %), 1.6/1	76 (63.3 %)/ 44 (36.7 %), 1.7/1	3 (42.9 %)/ 4 (57.1%), 0.75/1	X^2 = 1.046 p* = 0.306
Body mass, kg	77.9 (50 – 108)	77.91 (50 – 108)	76.8 (60 – 101)	t = 0.176 p* = 0.860
BMI<25	47 (37 %)	45 (37.5 %)	2 (28.6 %)	X^2 = 0.226 p* = 0.942
BMI>25	80 (63 %)	75 (62.5 %)	5 (71.4 %)	
Height, cm	173.8 (157 – 192.0)	174.0 (157 – 192)	170.3 (160 – 187)	t = 0.331 p* = 0.740

Note: data are presented as absolute and relative values, mean and standard deviation, POUR - acute postoperative urinary retention (POUR* - where POUR was considered the inability to urinate in the presence of bladder globus accompanied by specific clinical signs with the need for bladder catheterization [1]); BMI – body mass index, *p < 0.05 statistically significant, t – T-student test.

The surgical procedures included: abdominal surgery - appendectomy (34/127 [26.8%]), cholecystectomy (6/127 [4.7%]), inguinal/ventral hernioplasty (8/127 [6.3%]), laparotomies for acute abdomen (15/127 [15.8%])); pelvic surgery (3/127 [2.35%]); traumatology surgery - upper/lower limb (25/127 [40.9%], of which 11 cases of septic trauma); thoracotomies (3,127 [2.35%]; neurosurgery (3/127 [2.35%]); oromaxillofacial surgery (3/127 [2.35%]). The average duration of anesthesia was 122.1±31.4 minutes and the average duration of surgery was 111.3±30.3 minutes. The group comprised 3 types of anesthesia: general anesthesia 64% (81/127), spinal anesthesia 26% (33/127), peripheral nerve block 10% (13/127).

In our study, we identified a POUR prevalence of 5.5%. Patients with ASA IV-V anesthetic risk, who have severe preoperative comorbidities, were not included in the present study. As a result, this limitation will not allow us to generalize the obtained POUR prevalence. In addition, the exclusion of patients with ASA IV-V anesthetic risk also eliminates the risk of overestimating POUR prevalence.

The literature reports a wide range of POUR prevalence, from 5% to 70% [2]. The large discrepancy between the reported prevalence may indicate the complexity and multitude of risk factors contributing to the development of POUR on the one hand, and on the other hand, the heterogeneity of criteria used to diagnose POUR. Thus, in the present study, POUR was defined as the postoperative clinical situation characterized by inability to urinate requiring bladder catheterization. However, Cataldo P. [12] and Paulsen E. [13] establish the diagnosis of POUR even in the absence of micturition for ≥ 8 hours postoperatively. In addition to patients who required the insertion of Foley catheter, we recorded 3 patients who reported their first voiding at ≥ 8 hours after surgery (8, 13 and 9.5 hours), which would contribute to an increased prevalence of POUR in the present study from 5.5% (7/127) to 7.9% (10/127). Dobbs S. defines POUR when patient is unable to urinate for ≥12 hours postoperatively [14]. Applying this criterion in our study, POUR would have a prevalence of 6.3% (8/127) (Table 2).

Table 2. Prevalence of postoperative urinary retention depending on the definition criteria.

Postoperative urinary retention definition criteria	International studies	Moldova (present study)
Bladder catheterization [1]	5 -70%	5,5% (7/127)
Absence of urination ≥ 8 hours postoperatively [1, 12, 13]	52%	7,9% (10/127)
Absence of urination ≥ 12 hours postoperatively [1, 14]	36%	6,3% (8/127)

The first mobilization of patients with POUR was at 13.1±5.0 hours after surgery compared to 9.7±4.4 hours (Student's t = 1.97, p = 0.05) for non-POUR patients.

The secondary outcome parameters of the study in-volved testing a number of previously reported risk factors to determine whether their presence perioperatively would be associated with POUR. From the preoperative patient-re-lated parameters, none were confirmed as a risk factor for POUR: DM, higher anesthetic risk (ASA III versus ASA I-II), ≥ 3 deliveries in women, female gender, chronic beta-block-er medication, Class II NYHA HF, hypothyroidism, age ≥ 60 years or age ≥ 55 years, previous abdominal surgery, myo-cardial infarction in patient's history, cerebrovascular acci-dent (CVA) in medical history.

Among the patient-related preoperative risk factors that are unmodifiable, essential hypertension (HTN) was detect-ed as an independent risk factor for POUR (Table 3).

Table 3. Preoperative risk factors for acute postoperative urinary retention

Risk factors	Fisher exact	X^2	p^*	Odds ratio	X^2 Yates correction	p^*
Diabetes mellitus (DM)	0.114	3.712	0.054	4.93	1.526	0.217
≥ 3 childbirths	0.305	1.731	0.188	0.23	0.623	0.429
Anesthetic risk ASA III	0.680	0.380	0.538	1.62	0.040	0.842
Age ≥ 60 years old	0.205	2.258	0.132	3.11	1.175	0.278
Age ≥ 55 years old	0.111	3.189	0.074	4.17	1.926	0.165
BMI > 25	1.000	0.226	0.634	1.50	0.005	0.942
Female gender	0.425	1.180	0.277	2.30	0.147	0.493
HTN	0.041	5.655	0.017	9.00	3.939	0.047
Beta-blocker medication	0.425	1.180	0.277	2.30	0.470	0.493
NYHA II HF	0.426	0.591	0.441	1.77	0.118	0.730
Hypothyroidism	0.151	2.710	0.996	3.96	1.010	0.315
Association HTN+DM	0.026	10.786	0.001	23.0	5.354	0.020
Previous abdominal surgery	0.248	1.852	0.173	3.06	0.947	0.330
Previous myocardial infarction	0.539	0.132	0.716	1.50	0.077	0.781
Previous CVA	0.250	2.098	0.148	4.83	0.201	0.654

Note: DM - diabetes mellitus, BMI - body mass index, ASA - American Society of Anesthesiology, CVA – cerebrovascular accident/ stroke, HF II NYHA - NYHA heart failure grade II, HTN – arterial hypertension; *p < 0.05 statistically significant.

At the same time, DM, a risk factor often reported in the literature, was parametrically at the limit of statistical sig-nificance. This risk factor was further analyzed by applying the extended POUR definition (using the criterion of „lack of micturition in the first 8 hours postoperatively" [12, 13]), proving to be a statistically significant risk factor (OR = 5.1; X^2 (1, N = 127) = 5.36, p = 0.021). Returning to the study group where only the need for bladder catheterization was considered as POUR, we analyzed the association between HTN and DM, in our study they were detected as a combina-tion of risk factors for POUR. Similarly, stroke was tested as a risk factor on the group that included both patients with POUR (7/127) and those who had no voiding in the first 8 hours postoperatively (3/127) with confirmation of risk factor status (OR = 4.83 (X^2 (1, N = 127) = 7.405, p = 0.006).

From the intraoperative and intra-anesthetic parame-ters tested for the quality of risk factors for POUR none was confirmed (Table 4).

Table 4. Intraoperative and intra-anesthetic risk factors for acute postoperative urinary retention

Risk factors	Fisher exact	X^2	p^*	Odds Ratio	X^2 Yates correction	p^*
Fasting for fluids ≤ 6 hours	0.706	0.291	0.589	1.52	0.023	0.880
Surgery duration > 90 min.	1.000	0.016	0.899	1.11	0.087	0.767
Spinal anesthesia	1.000	0.026	0.872	1.15	0.080	0.777
Fentanyl ≥ 0,5 mg total	1.000	0.022	0.881	0.87	0.100	0.752
Atropine administration	0.103	3.564	0.059	4.48	2.211	0.137
Fentanyl ≥ 0,5 mg + 0,1 mg p/o	1.000	0.156	0.693	1.78	0.134	0.714
Anesthesia ≥ 95 min.	1.000	0.045	0.832	0.83	0.056	0.813
Abdomen + lower limb	1.000	0.073	0.787	1.35	0.055	0.815
Without NSAIDs	0.129	2.801	0.094	5.88	1.297	0.255
Opioid vs. non-opioid analgesia	0.678	0.070	0.792	0.80	0.036	0.850
Fluids ≥ 1500 ml i/ op	0.707	0.291	0.589	1.520	0.023	0.881
Bleeding volume ≤ 400 ml	0.478	0.296	0.586	0.55	0.0216	0.883
Analgesia (promedol + NSAIDs)	0.330	1.471	0.225	5.17	0.071	0.789
Fentanyl vs. Morphine p/o	1.0	0.037	0.847	1.27	0.173	0.677

Note: NSAIDs - non-steroidal anti-inflammatory drugs, min - minutes, i/op - intraoperative, p/o - postoperative; *p < 0.05 statistically significant.

Discussions

The present study aimed to test the risk factor associated with POUR for a number of perioperative parameters.

The surgical population studied inadvertently included predominantly male subjects (79/127 (62.2%)). However, according to our results, the prevalence of POUR was equivalent in both women (4/7) and men (3/7). Thus, sex remains a controversial risk factor. Some studies report that male sex is more frequently associated with POUR [2, 3], particularly due to specific pathologies such as benign prostatic hypertrophy [6]. Other studies report that the female sex is more susceptible to POUR [8]. Regarding patients' chronic medication, in our study patients with benign prostatic hypertrophy were excluded, as they were chronically medicated with alpha-receptor blockers (e.g., tamsulosin), which could have been an important factor of bias.

Advancing age increases the risk of POUR, with reports that patients aged ≥ 50 years develop POUR 2.4 to 2.8 times more frequently, which is attributed to progressive neuronal degeneration with bladder dysfunction [2-4, 6]. In our study, however older age (≥ 60 years/ ≥ 50 years) did not confirm its status as a risk factor for POUR. The explanation could lie in the fact that we excluded from the outset the anesthetic population with ASA score > III and orthopedic surgery.

Among the patient-related preoperative risk factors that are unmodifiable, HTN has been detected as an independent risk factor for POUR, consistent with previous studies.

Thus, in our study, patients with pre-existing neurological disorders such as stroke and diabetic neuropathy more frequently developed POUR. Pathophysiologically, the association of POUR and DM can be explained by impaired bladder sensation, decreased capacity, and contractility [15]. Similarly, Keita H. [3] confirmed that pre-existing neurological disorders (cerebral palsy, multiple sclerosis, alcoholic or diabetic polyneuropathy, poliomyelitis) as a risk factor for POUR. The same study reports a direct correlation between bladder volume ≥ 270 ml after surgery and POUR. In our study, there was no possibility to estimate bladder volume using ultrasound, but among the whole group of patients, only one patient stopped the intake of clear liquids 2 hours preoperatively, 60 patients did not ingest clear liquids 6 hours preoperatively, 67 respondents abstained more than 6 hours from preoperative fluid intake. In addition, an infusion volume ≥ 1500 ml intra-anesthetic or hemorrhage ≤ 400 ml were not identified as risk factors for POUR. No patient with peripheral nerve block developed POUR.

Although opioids are known to cause urinary retention, this concept was not confirmed in our study. At the same time, opioid-associated POUR phenomenon has been described at fentanyl doses above 1 mg [16]. In our study, regarding the total dose of fentanyl used perianesthetically no statistically significant differences were identified between fentanyl ≥ 0.5 mg + 0.1 mg postoperatively (10/127) versus fentanyl < 0.5 mg + 0.1 mg postoperatively (17/127). In addition, all patients had intra-anesthetic summary dose of fentanyl ≤ 0.8 mg.

Although non-steroidal anti-inflammatory drugs (NSAIDs) have often been questioned as contributing factors not only to the induction of acute kidney injury and to the precipitation of POUR, in our study no significant differences in the prevalence of POUR were identified between groups of patients who received NSAIDs intraoperatively and/or postoperatively versus those who were not exposed to these drugs. However, some studies describe the protective role of NSAIDs against the development of POUR, with NSAIDs being used in multimodal analgesia regimens to avoid opioid administration [4].

Patients' postoperative analgesia was also studied. Thus, 24.4% patients received nonopioid analgesia with NSAIDs (31/127) with 2 cases of POUR (2/31), 30.7% patients beneficiated of promedol (39/127) with 2 cases of POUR recorded (2/39), 27.6% fentanyl (35/127) with another 2 cases of POUR (2/35) and 17.3% morphine (22/127) with only 1 case of POUR (1/22). Cases of POUR were observed both in the opioid analgesia group, regardless of the morphine administered, and in the non-opioid analgesia group.

When analyzing risk factors that may be related to surgical treatment per se, it has been reported that the prevalence of POUR varies depending on the type of surgery and the likelihood of autonomic nerve injury (as a result of total mesorectal excision), pelvic nerve injury or reflex increase in internal sphincter tone (caused by pain in patients undergoing anorectal surgery) [17]. In the present study, no statistical differences were detected between the groups of patients who underwent abdominal and lower limb versus thoracic and upper limb surgery.

Previous studies support that increased volumes of intraoperatively administered fluids would be associated with higher prevalence for POUR [2-4]. In the present trial, we did not determine statistically significant differences between study groups in which ≥ 1500 mL infusible fluids versus < 1000 mL was administered intraoperatively. It is noteworthy that none of the patients infused with 500 ml-developed POUR.

Strengths and potential biases of the study

One of the strengths is the prospective methodology of the study. Also, to standardize data collection, patients were assessed postoperatively by the same investigator.

The given study has some limitations that need to be taken into consideration. One factor of bias would reside in the small sample and single-center study. Also, in the institution where the study took place, orthopedic surgery patients undergoing spinal anesthesia are inserted with urinary catheters from the beginning of surgery. Therefore, this large group of patients was excluded from the study. Also, patients who developed complications, with admission to Intensive Care Unit, were excluded from the study.

Conclusions

The prevalence of POUR in a single-center surgical population from the Republic of Moldova varies between 5.5% and 7.9%, depending on which criteria for POUR we apply.

In this regard, a consensus on diagnostic criteria for POUR is needed.

Patients with HTN and DM developed more frequently POUR. Patients with pre-existing neurological disorders such as positive history for stroke and diabetic polyneuropathy are more susceptible to POUR. It was also determined that the first mobilization of patients with POUR was later compared to patients who did not develop this complication.

Competing interests

None declared.

Authors' contribution

NB developed the study concept, literature review and survey questionnaire, analyzed the collected data, wrote, translated into English and approved the final version of the manuscript. CL recruited the eligible patients, completed the study questionnaire, monitored the patients in the first 24 hours postoperatively, numerated the data in Excel, and contributed to the literature review. Both authors approved the final version of the manuscript.

Informed consent for publication

Obtained.

Ethics approval

Not needed for this study.

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