

# Effects of Low Intraperitoneal Pressure on Quality of Postoperative Recovery after Laparoscopic Surgery for Genital Prolapse in Elderly Patients Aged 75 Years or Older

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2	Effects of low intraperitoneal pressure on quality of postoperative recovery after
3	laparoscopic surgery for genital prolapse in elderly patients age 75 years and older.
4	
5	Short title: laparoscopy and elderly patients age 75 years and older
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36	corresponding author (SM), on reasonable request.
37	

#### 38 Abstract

#### 39 STUDY OBJECTIVE

- 40 Previous clinical trials for laparoscopic surgery have included few elderly patients age ≥75
- 41 years. We aimed to evaluate the quality of postoperative recovery after laparoscopic surgery
- 42 using low intraperitoneal pressure (IPP) (6 mmHg) and warmed, humidified carbon dioxide
- 43 (CO<sub>2</sub>) gas for genital prolapse in elderly patients age  $\geq$ 75 years.

#### 44 DESIGN

45 Prospective consecutive case series.

#### 46 SETTING

47 University hospital

#### 48 **PATIENTS**

- 49 Consecutive patients (n=30) age ≥75 years planning to undergo laparoscopic surgery for
- 50 genital prolapse by the same surgeon were recruited from October 2016 to December 2019.

#### 51 **INTERVENTIONS**

- 52 Laparoscopic promontofixation for the treatment of genital prolapse was performed using low
- 53 IPP and warmed, humidified CO<sub>2</sub> gas<sup>-</sup> When a promontory could not be easily identified,
- 54 laparoscopic pectopexy was alternatively performed.

55

#### 56 **MEASUREMENTS**

- 57 The primary outcome was the QoR-40 (Quality of Recovery–40) score at 24 hours
- 58 postoperatively. Secondary outcomes were postoperative pain using a 100-mm visual
- analogue scale (VAS) and length of hospital stay after surgery (LHSS).

#### 60 MAIN RESULTS

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For the global QoR-40 score and for the four dimensions of the QoR-40, "emotional state,"

- 62 "physical comfort," "psychological support," and "pain," no differences were observed
- 63 between the baseline score and at 24 hours. The score for the "physical independence"
- 64 dimension was significantly lower at 24 hours than the baseline score (p<.001). No patient
- had VAS pain scores >30 of 100 at 12 hours or later. LHSS was <48 hours in 22 patients
- 66 (73.3%) and <72 hours in 8 patients (26.7%). Multivariable analysis showed that the odds of
- a LHSS >48 hours were more than 8 times higher in patients who were discharged from the
- operating room (OR) in the afternoon compared with those with a morning discharge.

### 69 CONCLUSION

- The use of a low IPP is feasible, safe and has clinical benefits for elderly patients age  $\geq$ 75
- 71 years who undergo laparoscopic surgery for genital prolapse.
- 72
- 73
- 74 KEYWORDS: length of hospital stay after surgery; pneumoperitoneum; Quality of
- 75 Recovery-40
- 76

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#### 77 Introduction

The chance of a woman experiencing a prolapse increases with age [1]. The lifetime 78 79 risk of undergoing a single operation for prolapse or incontinence by age 80 is 11.1% [1]. A 80 recent study demonstrated that the population undergoing surgery is aging at a faster rate than the general population [2]. By 2030, it is estimated that one-fifth of patients  $\geq$ 75 years 81 will undergo surgery each year in England [2]. Therefore, the number of very elderly patients 82 (age  $\geq$ 75 years) who require surgery for genital prolapse is also expected to rise [1, 2]. The 83 elderly are often excluded from clinical trials [2, 3], particularly surgical trials [2]. Therefore, 84 the best available evidence frequently does not apply to the elderly patient [2, 3]. 85 Our previous randomized 2x2 factorial trial controlled trial (RCT) showed that low 86

intraperitoneal pressure (IPP) (8 mmHg) and warmed, humidified carbon dioxide (CO<sub>2</sub>) gas independently decreased postoperative pain in patients <75 years old who underwent laparoscopic surgery for genital prolapse [4]. In the present prospective study, we attempted to evaluate benefits and harms of low IPP and warmed, humidified CO<sub>2</sub> gas in very elderly patients (age  $\geq$ 75 years) who underwent laparoscopic surgery for genital prolapse.

92

#### 93 Materials and Methods

#### 94 *Methods*

95 The study was designed as a prospective, single-center, observational trial. Consecutive patients age  $\geq$ 75 years who underwent laparoscopic surgery for genital 96 prolapse by the same surgeon (RB) were recruited at CHU Clermont-Ferrand from October 97 2016 through December 2019. Informed written consent was obtained from each patient 98 prior to surgery. The study protocol was approved on 29<sup>th</sup> September 2016 by the IRB 99 100 (00008526) of the Committee for Protection of Persons in Biomedical Research (CPP SE VI) 101 of the Auvergne (France) (2016 / CE 57) and registered on ClinicalTrials.gov 102 (NCT03095716).

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Anesthetic management was performed by staff anesthesiologists and 103 anesthesiology residents or nurse anesthetists under their supervision. Standard ASA 104 105 anesthetic monitors were placed in the operating room. All patients received the 106 same general technique, as we previously reported [4]. After induction of anesthesia, all 107 patients received forced-air warming blanket, which was maintained until the end of the 108 operation. Intraoperative core temperatures were recorded at 15-minute intervals by a probe 109 inserted into the distal esophagus throughout surgery. Paracetamol (1g) and ketoprofene 110 (50mg) were injected 30-minutes before the end of the operation to prevent postoperative 111 pain.

All operations were performed by the same surgeon (RB) with the assistance of a 112 gynecological surgical resident. Insufflation of CO<sub>2</sub> gas was performed using a Storz 113 electronic endoflator (KARL STORZ SE & Co. KG, Tuttlingen, Germany). All patients 114 received warm (37 °C), humidified (98% relative humidity) CO<sub>2</sub> insufflation gas delivered 115 by the Fisher & Paykel MR860 Laparoscopic Humidification System (HumiGard, Fisher & 116 117 Paykel Healthcare, Auckland, New Zealand) [4]. When the IPP reached 15 mmHg, four trocars were inserted. Then, the IPP was immediately decreased to 6 mmHg, at which it was 118 maintained for the duration of surgery. We used 6 mmHg instead of 8 mmHg for low IPP in 119 120 the present study, according to the European Association for Endoscopic Surgery clinical 121 practice guideline on pneumoperitoneum for laparoscopic surgery, which recommends using 122 the lowest IPP possible that allows adequate exposure of the operative field rather than 123 using a routine pressure [5].

For all patients, 5 mL ropivacaine hydrochloride solution (2 mg/mL) were infiltrated around the trocar wounds before incisions [1]. In addition, 20 mL ropivacaine solution (2 mg/mL) were infused under the right hemidiaphragm just after insertion of the four trocars [4]. We have been systematically performing infusion of ropivacaine solution under the right hemidiaphragm for over 10 years [6]. During this time, we have had no or few complaints about shoulder pain after laparoscopic surgery [1]. Laparoscopic promontofixation with or without (for patients who have already undergone hysterectomy) sub-total hysterectomy

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using the same surgical technique as we previously described [7]. When a promontory as 131 well as surrounding blood vessels and/or nerves could not be very easily identified, 132 laparoscopic pectopexy (the iliopectineal ligament on both sides is used for the mesh 133 134 fixation) [8] was alternatively performed to avoid potential surgical complications such as blood vessels and/or nerves injuries. A RCT showed that both surgical 135 techniques are equally effective in treatment for genital prolapse [8]. We did not attempt to 136 increase IPP prior to making the decision to proceed to pectopexy, because we know from 137 138 our experiences that increased IPP does not enable us to identify a promontory and surrounding vessels and nerves more easily [4]. 139

All patients were treated according to our standardised postoperative pain 140 management as we previously reported [4]. After arrival of patients in the postanesthesia 141 care unit (PACU), they were asked to rate intensity of their pain at rest using a 100-mm 142 visual analogue scale (VAS) every 20 minutes. When the VAS pain score was >30 of 100 143 144 (clinically relevant moderate to severe pain), patients received an IV bolus of 2 to 3 mg morphine, and then 1 to 2 mg IV every 10 minutes to achieve a VAS pain score  $\leq$ 30 of 100 145 146 (no or mild pain). After arrival of patients in the ward, they were asked to rate intensity of their pain at rest using a VAS, every 4 hours for the first 24 hours postoperatively, and then 3 147 times daily until discharge. The patients received intravenous paracetamol (1g) and 148 ketoprofene (50 mg) every 6 hours for the first 24 hours postoperatively. 149

The quality of postoperative functional recovery was evaluated using the Quality of Recovery–40 questionnaire (QoR-40) [4, 9]. The QoR-40 the quality of postoperative recovery through five dimensions (emotional state, physical comfort, psychological support, physical independence, and pain) [9]. This tool has been confirmed as the most suitable one for perioperative clinical studies [10].

155 The day before surgery (between 7:00 and 8:00 pm), patients were asked to 156 complete QoR-40 questionnaire as a measure of baseline score. The QoR-40 was then administered three times, 24 hours and 48 hours postoperatively. Surgical complications

were categorized according to the Clavien-Dindo classification [11].

The quality of the surgical conditions (the operative technical difficulty, working space, visibility, and pain experienced by the surgeon during surgery) was rated by the operating surgeon at the end of surgery using a 100-mm VAS [4].

162

#### 163 Statistical analysis

Statistical analysis was performed using the STATA program version 13.1
(StataCorp, College Station, TX, USA). The power calculation of the present trial was based
on our previous RCT [4].

VAS pain scores were categorized as either  $\leq$ 30 of 100 or >30 of 100, providing a 167 binary variable, as previously reported [4, 12]. Comparisons of the global QoR-40 scores and 168 the dimensions of the QoR-40 questionnaire between the day before surgery (baseline) and 169 at 24 hours were made using Wilcoxon's matched pairs test. To identify factors that influence 170 the length of hospital stay after surgery (LHSS), the first set of analyses compared the 171 characteristics of patients in the two length of stay groups (<48 hours versus <72 hours). 172 173 Continuous variables were compared between groups using either the unpaired t-test, if found to be normally distributed, or the Mann-Whitney test or one-way analysis of variance 174 (ANOVA) otherwise. Categorical variables were compared between groups using Fisher's 175 exact test. The second stage examined the joint association between patient factors and 176 177 LHSS in a multivariable analysis. As there were only two different lengths of stay (<48 hours versus <72 hours), the analysis was performed using logistic regression. Due to the relatively 178 small sample size, only variables showing some association with LHSS from the initial 179 analyses (p<.1) were included in this stage of the analysis. A backwards selection procedure 180 181 was used to retain only the statistically significant variables in the final model. This involved omitting non-significant variables, one at a time, until only significant variables remain. 182

183 Statistical significance was accepted at the .05 level.

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#### 185 Results

186	A total of 32 consecutive patients age $\geq$ 75 years underwent surgical treatment for
187	genital prolapse by the same surgeon (RB) during the study period. Among them, 2 patients
188	declined participation in completing the QoR-40 questionnaire. The present analysis
189	therefore included 30 patients. Patient and surgical characteristics are presented in Table 1.
190	A total of 27 patients underwent laparoscopic promontofixation and the remaining 3 patients
191	underwent pectopexy. A total of 6 patients had previously undergone total hysterectomy. The
192	surgery could be performed with 6 mmHg and the IPP was maintained for the duration of
193	surgery in all the patients.
194	
195	Quality of Recovery–40 score

Results are shown in Table 2. For the global QoR-40 score and for the four
dimensions of the QoR-40, "emotional state," "physical comfort," "psychological support," and
"pain," no differences were observed between the basal score (the day before surgery,
between 7:00 and 8:00 pm) and at 24 hours (Table 2). The score for the "physical
independence" dimension was significantly lower at 24 hours than the basal score (Table 2).

For further evaluation of the effects of a low IPP in elderly patients, we compared the 201 present results with those of our previous RCT in elderly patients age  $\geq 65$  years [4] 202 (Supplementary Table 1). In our previous RCT [4], we included only patients who underwent 203 laparoscopic sub-total hysterectomy with promontofixation. Thus, we performed a subgroup 204 analysis of patients underwent the same surgical procedure (n=21) for comparison 205 (Supplementary Table 1). The operating time was significantly longer in our previous RCT 206 207 compared with those of the present study (Supplementary Table 1). The results of the QoR40 scores (Supplementary Table 2) in these patients (n=21) were nearly identical to 208 those in the full cohort of patients (Table 2). In the low IPP group (8 mmHg, n=17) of our 209

184

previous RCT study [4], the scores for the two dimensions of the QoR-40, "physical comfort,"
and "physical independence," were significantly lower at 24 hours than the basal score
(Supplementary Table 2). In the standard IPP group (12 mmHg, n=20) of our previous RCT
[4], the scores for the global QoR-40, and for the three dimensions of the QoR-40, "physical
comfort," "physical independence," and "pain," were significantly lower at 24 hours than the
basal score (Supplementary Table 2).

To evaluate the effects of additional surgical procedures (Burch colposuspension [Burch] or tension-free vaginal tape procedure [TVT]), we compared the QoR40 scores between patients with (n=18) versus without (n=12) additional surgical procedures (Supplementary Table 3). The operating time was significantly longer in patients with additional surgical procedures (Supplementary Table 3). There was no significant difference in the QoR40 scores between patients with versus without additional surgical procedures (Supplementary Table 4)

223

#### 224 VAS pain score

No patients had a VAS pain score >30 the day before surgery. No patients had postoperative shoulder pain. In the PACU, 7 (23%) patients had VAS pain scores >30. Upon arrival in the ward, 4 patients (13%) had VAS pain scores >30. No patient had VAS pain scores >30 at 12 hours or later (Supplementary Table 5).

229

#### 230 Intraoperative core temperature

Results are shown in Table 1. We observed that 15 of 30 (50.0%) patients were already hypothermic just before starting the surgical procedure. Of these, 10 patients remained hypothermic at the end of surgery. A total of 15 patients (50.0%) were hypothermic at the end of surgery.

235

#### 236 Intraoperative and postoperative complications

237 One patient who had previously undergone laparoscopic total hysterectomy following 238 radiation therapy and chemotherapy due to uterine adenocarcinoma experienced an 239 intraoperative complication (bladder perforation). This was not related to low IPP, but rather 240 due to postsurgical and/or radiation-related adhesion. The patient was discharged on 241 postoperative day 2 with an indwelling urinary catheter after an uneventful postoperative 242 recovery. No patients experienced postoperative complications. No patients required re-243 admission.

244

#### 245 *Quality of surgical conditions*

Scores rated by the surgeon for operative technical difficulty and working space for dissection and suturing, surgical field visibility, pain experienced by the surgeon, are shown in Supplementary Table 6.

249

### 250 Factors influencing length of hospital stay after surgery

LHSS was <48 hours in 22 patients (73.3%) and <72 hours in 8 patients (26.7%). The 251 results of analyses comparing the characteristics of patients with a LHSS <48 hours versus 252 253 those >48 hours are summarized in Table 3. Only operating room (OR) exit time significantly differed between patients with a LHSS <48 hours versus those >48 hours (Table 3). The 254 majority (n=7, 88%) of patients with a LHSS >48 hours were discharged from the OR in the 255 afternoon, whilst only one third of those (n=7, 32%) with a LHSS <48 hours were discharged 256 in the afternoon (Table 3). In addition to this significant factor, some evidence suggested that 257 operating time and QoR-40 scores at both baseline and 24 hours varied between the two 258 discharge time groups (Table 3). However, the results for all these variables did not quite 259 reach statistical significance. 260

Next, we examined the joint association between select factors and LHSS. A
 backwards selection procedure was used to retain only the significant variables in the final

analysis (Table 4). The results of the multivariable analysis showed that both OR exit time 263 and QoR-40 score at baseline were associated with LHSS (Table 4). After adjusting for these 264 265 two variables, no additional effects of either operating time or QoR-40 score at 24 hours were observed, both of which showed some association with LHSS in the initial analyses. OR exit 266 time was again associated with an increased chance of a LHSS >48 hours. The odds of a 267 LHSS >48 hours were more than 8 times higher in those with an afternoon discharge from 268 269 the OR compared with those with a morning discharge (Table 4). Higher baseline QoR-40 270 scores were associated with a lower chance of a LHSS >48 hours (Table 4). A one-unit increase in this score was associated with a 20% reduction in the odds of a LHSS >48 hours 271 (Table 4). 272

273

#### 274 Discussion

275 The present study showed the feasibility, safety, and clinical benefits of use of a low IPP in very elderly patients who undergo laparoscopic surgery for genital prolapse, in 276 agreement with the results of our previous RCT [4]. Laparoscopic surgery confers several 277 benefits, including decreased postoperative pain, shorter LHSS, and a more rapid return to 278 279 regular daily activities [13, 14]. However, previous clinical trials for laparoscopic surgery have 280 included few very elderly patients age  $\geq$ 75 years [15,16], who may benefit the most from 281 laparoscopic surgery [13, 14]. The present clinical trial could provide useful information for 282 very elderly population who require surgical treatment for genital prolapse.

Several limitations of the present study should be considered, and thus, the present findings may not be generalized. First, the present study included only a small number of patients and the same surgeon performed all procedures. Second, there were no agematched controls. Because our previous RCT [4] has already showed that a low IPP could result in better clinical outcomes, it was not ethically possible to include age-matched elderly patients who undergo laparoscopic surgery under a standard IPP (12mmHg). However, 289 compared with those of our previous RCT in elderly patients age  $\geq 65$  years, the present findings suggested that the use of a low IPP could result in better quality of postoperative 290 291 recovery in elderly patients. Third, there may be biases associated with the heterogeneity of 292 the present study population. However, we found no significant differences in the QoR 40 293 scores between patients with versus without additional surgical procedures. In a subgroup analysis of patients who underwent laparoscopic sub-total hysterectomy with 294 295 promontofixation, results of the QoR40 scores were nearly identical to those in the full cohort 296 of patients. Furthermore, the objective of the present study was not to identify factors that 297 influence LHSS; thus, we did not calculate the required number of patients for this. A large, prospective, multicenter study is required to confirm the external validity of the present 298 results. 299

300 Aging is associated with impaired thermoregulatory function [17-19]; thus, elderly 301 people are at greater risk of perioperative hypothermia [17-19]. The present study showed no 302 significant effect of warmed CO<sub>2</sub> gas on the prevention of intraoperative hypothermia, also consistent with our previous RCT [4]. Studies have suggested that extending systemic 303 304 warming into the preoperative period could be more beneficial than limiting it to surgery alone [20-22]. Further studies are required to investigate whether the preoperative plus 305 306 intraoperative systemic warming could be more effective for prevention of hypothermia in elderly patients. 307

308 LHSS after surgery can be influenced by many different factors, including medical 309 and social factors [23-32]. The presence of comorbidity has been a well-known risk factor 310 [28-30]. In the present study, no patients had ASA III-IV. Because genital prolapse is not a life-threatening disease, only selected elderly patients who are in relatively good condition 311 312 might be referred to our hospital for surgical treatment. In the present study as well as in our previous RCT [4], all patients were informed of expected discharge on postoperative day 3. 313 314 Social reasons requiring a longer hospital stay are more common in elderly patients [23, 24, 26, 27, 31]. Furthermore, studies have shown that older age can influence LHSS after 315

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surgery [26, 31, 32]. Nevertheless, in the present study, 77.3% (22 of 30) patients were 316 317 discharged within 48 hours after surgery, whereas among elderly patients age ≥65 years of 318 our previous RCT [4], only 25.0% (5 of 20) patients in the 12-mmHg group and 23.5% (4 of 319 17) in the 8 mmHg group were discharged within 48 hours (Supplementary Table 1). 320 Shortening LHSS after surgery benefits patients who prefer to be at home [33]. Furthermore, in two 10-year age groups (68–77 years and 78–87 years), better functional outcomes were 321 322 associated with short admissions [34]. Finally, shortening LHSS benefits the healthcare 323 system in terms of cost and resource utilization [35, 36]. Hospital health care resource 324 utilization and costs are disproportionately incurred within the first 3 days of hospitalization 325 [35, 36]. It is therefore clinically important for both elderly patients and the healthcare system 326 to identify factors influencing LHSS. In the present study population, the odds of a LHSS >48 327 hours were more than 8 times higher in patients with an afternoon discharge from the OR compared with those with a morning discharge. Furthermore, we observed approximately a 328 329 40-minute decrease both operation time and CO<sub>2</sub> pneumoperitoneum in the present t study compared to those in our previous RCT [4]. Previous studies suggested operation time [26, 330 32, 37, 38] and postoperative pain [28, 38, 39] as factors that influence LHSS. Thus, at least 331 three modifiable factors exist at an institutional and/or individual surgeon level-332 postoperative pain, operative time, and OR exit time-that may potentially influence LHSS in 333 334 very elderly patients after laparoscopic surgical treatment for genital prolapse. To reduce postoperative pain, a lower IPP [4, 28, 40] could be used. To reduce operative time, 335 improving surgeon efficiency is required: at an institutional level by improving coordination of 336 337 the surgical team and limiting unnecessary delays [41-45]. At the individual surgeon level, this could be achieved by improving technical skill and operative efficiency [46, 47]. 338 Achievement of a morning discharge from the OR is accomplished by a shorter operation 339 time and by scheduling the procedure as the first case in the morning [45]. Because the 340 341 number of very elderly patients who require surgical treatment is expected to rise [2], 342 improving the operating schedule and OR efficiency are worthy of careful consideration for both patients and healthcare systems. 343

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## Table 1. Baseline Characteristics

Variable	6 mmHg warmed, humidified CO2 gas (n=30)
Vallablo	
Age, mean (SD), y	80.4 (3.5)
Hight, mean (SD), cm	156.7 (19.4)
Body weight, Mean (SD), kg	59.3 (12.7)
Body mass index, mean (SD)	23.4 (3.7)
ASA status, No. (%) ASA 1	4 (13.3)
ASA 2	26 (86.7)
ACCI	3.6 (0.50)
Basal QoR 40, median [IQR]	191 [185, 197]
Additional surgical procedure, No. (%)	

None Burch TVT	12 (40.0) 15 (50.0) 3 (10.0)
Operating time, mean (SD), min Total operating time Pneumoperitoneum time	124.3 (21.7) 112.3 (23.2)
Volume CO2 used, median [IQR], L	160 [80, 240]
Trendelenburg position, mean (SD), degree	22.6 (3.6)
Intraoperative sufentanyl usage, mean (SD), μg	41.0 (6.8)
Core temperature,	
median [IQR], °C Start temperature	35.9 [35.2, 36.6]
Minimum temperature	35.7 [35.2, 36.2]
Maximum temperature	36.0 [35.4, 36.6]
Final temperature	35.9 [35.4, 36.4]
PACU LOS, mean (SD), min	118.6 (24.6)

Morphine requirement in the PACU	
Patients requiring morphine, No. (%)	7 (23.3)
Morphine usage, median [IQR], mg	0 (1)
Patients requiring morphine in the ward, No. (%)	0 (0)
Hospital LOS after surgery, No. (%)	
<48h >48h <72h	22 (73.3) 8 (26.7)

## Abbreviations

IPP: intraperitoneal pressure, IQR: interquartile range, ASA: American Society of Anesthesiologists

ACCI: age-adjusted Charlson comorbidity index, QoR 40: Quality of Recovery-40

LOS: length of stay , PACU: postanesthesia care unit

Burch: Burch colposuspension, TVT: tension-free vaginal tape procedure

## Table 2: QoR-40 Scores

Outcome	Baseline	24h	P-value
	(n=30)	(n=30)	
Global score	191 [185, 197]	190 [181, 194]	.07
C1: Emotional State	42 [38, 43]	41.5 [38, 44]	.76
C2: Physical Comfort	57.5 [54, 59]	58 [53, 58]	.33
C3: Psychological Support	35 [35, 35]	35 [35, 35]	.34
C4: Physical independence	25 [24, 25]	23.5 [22, 25]	<.001

C5: Pain	34 [31, 35]	33 [32, 34]	0.24
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Results are shown as median [inter-quartile range]

Abbreviations

QoR-40: Quality of Recovery-40

Variable	Category	<48 hours (n=22)	>48h <72h (n=8)	P-value
Age	-	80.4 ± 3.8	80.4 ± 2.6	.98
ASA score	1 2	3 (14%) 19 (86%)	1 (12%) 7 (88%)	1.00
ACCI		$3.5 \pm 0.5$	$3.7 \pm 0.5$	.32
ЗМІ	-	23.6 ± 3.6	$24.0 \pm 4.4$	.80
CO2 duration (mins)	-	108 ± 19	120 ± 25	.18
Operating time (mins)	-	119 ± 20	134 ± 22	.09
Volume CO2 (L)	-	165 [115, 200]	154 [121, 198]	.98
Additional procedure	No Yes	10 (45%) 12 (55%)	1 (12%) 7 (88%)	.20
Surgical difficulty	-	35 ± 16	33 ± 16	.71
OR exit time	AM PM	15 (68%) 7 (32%)	1 (12%) 7 (88%)	.01
Weekend discharge	No Yes	14 (64) 8 (36%)	5 (62%) 3 (38%)	1.00
LOS in PACU	-	118 ± 24	120 ± 27	.83
Pain in PACU	≤ 30 > 30	15 (68%) 7 (32%)	5 (62%) 3 (38%)	.77

## Table 3: Comparisons of patients with a LHSS <48 hours versus those >48 hours

Temperature start	-	36.0 [35.6, 36.3]	36.1 [35.9, 36.2]	.83
Temperature end		36.0 [35.5, 36.2]	35.8 [35.5, 36.2]	.96
QoR40 score baseline	-	193 [189, 197]	182 [177, 190]	.06
QoR40 score 24 hrs		191 [182, 196]	184 [179, 190]	.08

Results are shown as mean ± standard deviation, median [inter-quartile range], or number (percentage)

Abbreviations

LHSS; Length of hospital stay after surgery, ASA: American Society of Anesthesiologists, ACCI: age-adjusted Charlson comorbidity index,

BMI: body mass index, OR: operating room, QoR-40: Quality of Recovery-40, LOS: length of stay, PACU: postanesthesia care unit

## Table 4: Multivariable analysis of factors associated with LHSS

Variable	Category	Odds Ratio (95% CI) (*)	P-value
OR exit time	AM PM	1 8.6 (1.7, 4461)	.03
QoR40 score baseline	_	0.80 (0.65, 0.99)	.04

(\*) Odds ratios represent the odds of a LHSS >48 hours.

## Abbreviations

LHSS; Length of hospital stay after surgery, OR: Operating room, QoR-40: Quality of Recovery-40