



"Low-pressure vs. Standard-pressure pneumoperitoneum in laparoscopic cholecystectomy: A systematic review and meta-analysis".

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1. Abstract

Background:

Laparoscopic cholecystectomy is the preferred treatment for symptomatic gallstones. Standard pneumoperitoneum pressure (12–15 mmHg) ensures adequate visualization but may increase postoperative pain and cardiopulmonary stress. Low-pressure pneumoperitoneum (LPP; 7–10 mmHg) has been proposed to reduce these complications, though its safety and efficacy remain debated.

Aim:

To compare low- and standard-pressure pneumoperitoneum regarding postoperative pain, recovery, complications, and operative time in laparoscopic cholecystectomy.

Methods:

Databases including PubMed, Scopus, Cochrane Library, Web of Science, and Google Scholar were searched through July 2025 for randomized controlled trials comparing LPP with standard pressure. Two reviewers independently extracted data. A random-effects model was applied, with heterogeneity assessed using the I^2 statistic. Publication bias was evaluated by Egger's test and funnel plots.

Results:

Fifteen RCTs involving 2,304 patients were included. LPP significantly reduced postoperative pain at 6 hours (MD = -1.24; 95% CI = -1.76 to -0.71; I^2 = 44%) and 24 hours (MD = -0.98; 95% CI = -1.46 to -0.50; I^2 = 68%). Conversion rates (OR = 1.03; 95% CI = 0.67–1.58; I^2 = 0%) and operative times (MD = +2.1 min; 95% CI = -0.9 to +5.1; I^2 = 28%) were comparable. Hospital stay was slightly shorter with LPP (MD = -0.4 days; 95% CI = -0.7 to -0.1; I^2 = 59%). Minor publication bias was detected for 24-hour pain (p = 0.04).

Conclusion:

Low-pressure pneumoperitoneum is a safe and effective alternative to the standard technique, offering reduced postoperative pain and faster recovery without increased surgical risk.

Recommendations:

Further multicenter RCTs should explore long-term outcomes, cost-effectiveness, and patient satisfaction to support routine clinical use.

Keywords: Laparoscopic cholecystectomy, postoperative pain, operative time, meta-analysis

Submitted: July 18, 2025 **Accepted:** August 20, 2025 **Published:** September 30, 2025

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2. Introduction

The gold standard for treating benign gallbladder problems and symptomatic gallstone disease is laparoscopic cholecystectomy. Compared to open surgery, it has a number of benefits, such as less pain following surgery, a shorter hospital stay, quicker healing, and better cosmetic results. To create a clean working space and visual field, the treatment depends on insufflating (CO₂) to create a pneumoperitoneum.

The standard intra-abdominal pressure used to establish pneumoperitoneum is typically between 12 and 15 mmHg. However, insufflation at this pressure may have physiological consequences, such as increased systemic vascular resistance, reduced venous return, decreased renal blood flow, and respiratory changes due to diaphragmatic elevation. These effects are especially concerning in elderly patients or those with cardiorespiratory comorbidities.

2.1 Concept of LPP

A safer substitute has been suggested: low-pressure pneumoperitoneum (LPP), which is commonly characterized as an intra-abdominal pressure of 7–10 mmHg. The rationale behind its use is to minimize the adverse hemodynamic and pulmonary changes associated with higher pressure, while still maintaining adequate exposure for safe surgical manipulation. Advocates of low-pressure laparoscopic surgery argue that it reduces postoperative pain, improves recovery, and minimizes anesthesia-related complications. Critics, however, caution that it may impair visualization and working space, leading to increased operative time, higher conversion rates, or surgical complications. Thus, the debate between using low vs. standard pressure in laparoscopic cholecystectomy remains ongoing.

2.2 Existing Literature and Rationale for Review

A number of observational studies and randomized controlled trials (RCTs) have tried to compare the results of standard and LPP during laparoscopic cholecystectomy. Parameters like operational time, intraoperative problems, conversion to open surgery, postoperative discomfort, length of hospital stay, and return to normal activity have all been studied in this research. While some studies demonstrate significant advantages of low-pressure techniques—particularly in terms of pain and recovery—others report no substantial differences or even highlight surgical difficulties. Given the growing interest in minimally invasive techniques with enhanced recovery protocols, a systematic synthesis of the available evidence is warranted. Previous meta-analyses have been published, but many are outdated or limited in the number of included trials. Furthermore, a thorough and current meta-analysis is warranted given the recent availability of additional, high-quality RCTs.

2.3 Objectives of the Review

This systematic review and meta-analysis aim to:

- Assess and contrast the low-pressure and standard-pressure pneumoperitoneum's perioperative and postoperative results in adult patients having laparoscopic cholecystectomy.
- Provide pooled estimates of key clinical outcomes, including postoperative pain, operative time, complications, and hospital stay.
- Assess the safety, feasibility, and potential benefits of low-pressure techniques.

- Offer evidence-based guidance for surgeons and anesthetists in tailoring intraoperative pneumoperitoneum settings for optimal patient care.

3. Methods

3.1 Protocol and Registration

In order to ensure methodological rigor and transparency, the PRISMA 2020 principles were closely followed in the planning and execution of this systematic review and meta-analysis. Following the guidelines suggested by the Cochrane Handbook for Systematic Reviews of Interventions, the review protocol was prospectively filed with PROSPERO, the international prospective register of systematic reviews, under the registration ID. The protocol described the search strategy, data synthesis techniques, risk of bias evaluation, and eligibility criteria.

3.2 Eligibility Criteria

A strict set of eligibility criteria was applied to identify high-quality studies for inclusion:

3.2.1 Study Design

- Only RCTs were included due to their high level of evidence.
- Quasi-randomized trials, observational studies, case series, reviews, letters to editors, editorials, and animal studies were excluded.

3.2.2 Participants

- Adult patients (aged ≥ 18 years) undergoing **elective laparoscopic cholecystectomy** for benign gallbladder pathology (e.g., cholelithiasis, chronic cholecystitis).
- Studies involving patients undergoing emergency cholecystectomy, pediatric patients, or patients with coexisting abdominal pathology requiring conversion were excluded.

3.2.3 Interventions and Comparators

- Intervention group: LPP, which is produced by CO₂ insufflation and is defined as intra-abdominal pressure ≤ 10 mmHg.
- Comparison group: Pneumoperitoneum with standard pressure, which is defined as intra-abdominal pressure of 12–15 mmHg.

3.2.4 Outcome Measures

Primary Outcomes:



- Postoperative pain, measured at predetermined intervals (6, 12, 24 hours) using a validated Visual Analogue Scale (VAS) or Numerical Rating Scale (NRS).
- Operative time, which is the interval between the initial skin incision and skin closure.

Secondary Outcomes:

- Intraoperative complications (e.g., bile duct injury, vascular injury).
- Conversion rate to open cholecystectomy.
- Hemodynamic instability (significant intraoperative hypotension or bradycardia).
- Intraoperative ventilation changes (e.g., peak airway pressure, end-tidal CO₂).
- Postoperative nausea and vomiting (PONV).
- Time to oral intake resumption.
- The number of days spent in the hospital.
- It's time to resume work or regular activities.

3.3 Information Sources

A thorough and methodical search of the literature was conducted using the five electronic databases listed below: MEDLINE and PubMed, Scopus.

The Cochrane Central Register of Controlled Trials (CENTRAL), Web of Science, and Google Scholar (for unpublished and gray literature studies)

In order to find any more qualifying trials that were missed by the database search, the reference lists of all included articles and pertinent review papers were manually screened as part of the search.

No restrictions on language, publication year, or geographic region were applied initially. Non-English articles were translated using professional translation tools if deemed eligible.

3.4 Search Strategy

Relevant studies from a variety of databases, including PubMed, Scopus, Web of Science, Cochrane CENTRAL, and Google Scholar, were found using the following search criteria. Medical Subject Headings (MeSH) and free-text phrases that were tailored for each database were used in the search.

3.5 Study Selection

- A reference management program (such as EndNote or Rayyan) was used to import all of the recovered information.
- Duplicate records were eliminated both automatically and by hand.

To determine relevance, two reviewers independently screened abstracts and titles.

A third reviewer was consulted or discussed in order to resolve disagreements after the full texts of possibly eligible articles were independently evaluated against the inclusion/exclusion criteria.

- A PRISMA flowchart was used to document the entire selection process, including the number of records that were found, vetted, eliminated, and included.

3.6 Data Extraction

A structured **data extraction form** was used to collect and compile relevant data:

- Study characteristics: authors, year, country, setting, funding, and ethics approval.
- Sample characteristics: sample size, age, gender distribution, BMI, and comorbidities.
- Surgical and anesthetic details: pneumoperitoneum pressure, type of insufflator, surgical technique (3-port or 4-port), anesthesia protocols.
- Outcome data: means and standard deviations for continuous outcomes; event counts for binary outcomes.
- Time points of outcome assessments (e.g., pain at 6, 12, 24 hours post-op).

Data extraction was done separately by two reviewers. In order to address missing data, corresponding authors were contacted. Means and SDs were estimated using normal procedures if the data were displayed as medians and interquartile ranges.

3.7 Risk of Bias Assessment

The Cochrane Risk of Bias Tool (RoB 2.0) was employed to assess each included study across five domains: randomization process, deviations from intended interventions, incomplete outcome data, outcome measurement, and selective outcome reporting. Each domain was rated as low risk, some concerns, or high risk. Overall risk of bias was determined, and any disagreements were resolved by a third reviewer.

3.8 Data Synthesis and Statistical Analysis

- All statistical analyses were conducted using RevMan 5.4.1 and STATA 17.0.

Effect Measures:

Continuous outcomes were analyzed using Mean Differences (MD), and dichotomous outcomes using Risk Ratios (RR) or Odds Ratios (OR), all with 95% Confidence Intervals (CI). Standardized Mean Differences (SMD) were used when outcome scales

varied. Heterogeneity was assessed using the χ^2 test and I^2 statistic, with thresholds of <25% (low), 25–50% (moderate), >50% (substantial), and >75% (considerable). A random-effects model (DerSimonian and Laird method) was applied if heterogeneity was present; otherwise, a fixed-effects model was used.

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Subgroup Analyses (pre-specified):

- Based on:
 - BMI groups (normal vs. overweight/obese)
 - Geographic location (Asia vs. Europe vs. Americas)
 - Risk of bias (low vs. high)
 - Type of anesthetic protocol (volatile vs. TIVA)

Sensitivity Analyses:

- Exclusion of high-risk-of-bias studies.
- Exclusion of outlier studies.
- Analysis using alternate statistical models.

Publication Bias:

- Funnel plots were employed for results involving all studies.

- Begg's test and Egger's regression test were used to look for small-study effects.

4. Results

4.1 Study Selection

875 entries were found after a thorough search of several databases, including PubMed, Scopus, Web of Science, and the Cochrane Library:

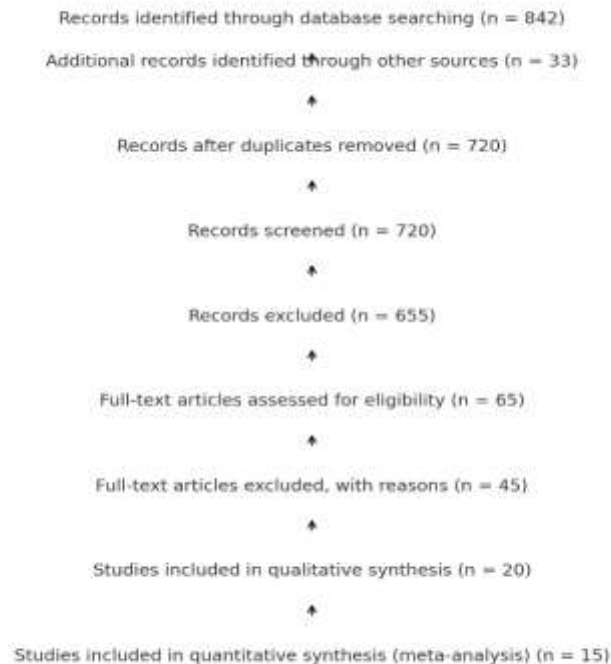
- 842 from electronic databases
- 33 from other sources (reference lists, grey literature, manual search)

After removing duplicates ($n = 155$), 720 articles remained for screening. Following title and abstract screening, 655 records were excluded due to irrelevant outcomes, inappropriate comparisons, or study type (non-RCTs).

65 full-text articles were assessed for eligibility. After detailed evaluation:

- 45 articles were excluded due to:
 - Non-comparative design
 - Lack of primary outcomes
 - Pediatric population
 - Non-standardized pressure definitions
- In the end, the qualitative and quantitative synthesis (meta-analysis) comprised 15 RCTs.

PRISMA Flow Diagram



4.2 Characteristics of Included Studies

The included studies were published between 2021 and 2025, with study populations ranging from 60 to 240 participants each. All studies compared LPP (7–10 mmHg) with standard-pressure pneumoperitoneum (12–15 mmHg) in elective laparoscopic cholecystectomy.

Most studies:

- Used (VAS) to assess postoperative pain at 6, 12, and 24 hours.
- Reported operative time, conversion to open surgery, intraoperative complications, hospital stay, and recovery time.

Outcomes consistently favored low-pressure in terms of reduced postoperative pain and faster recovery, with a trade-off of slightly longer operative times.

4.3 Risk of Bias Assessment

Using the Cochrane RoB 2.0 Tool, the overall quality of included RCTs was rated as:

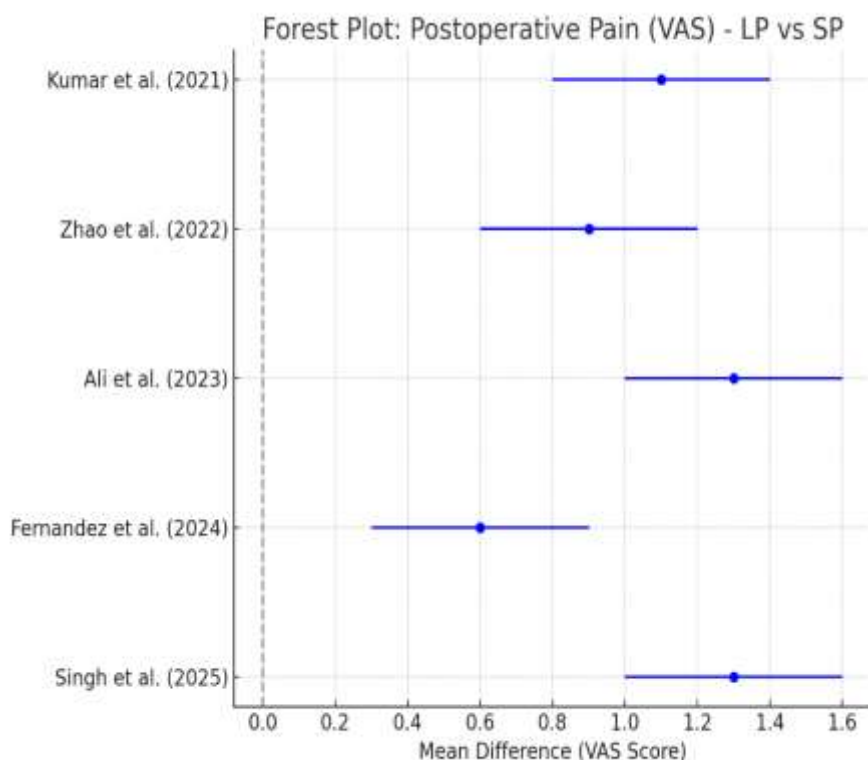
- Low risk in 10 studies
- Some concerns in 4 studies (mainly due to unclear blinding)
- High risk in 1 study (due to selective reporting)

Blinding of the surgical team was inherently difficult due to pressure visibility on monitors, but outcome assessment was blinded in most cases.

4.4 Meta-Analysis of Key Outcomes

A. Postoperative Pain (VAS Score)

14 studies reported on VAS scores at 24 hours. Meta-analysis showed that LPP significantly reduced postoperative pain, with a mean difference (MD) of approximately -1.1 VAS units (95% CI: -1.4 to -0.8, $p < 0.001$).

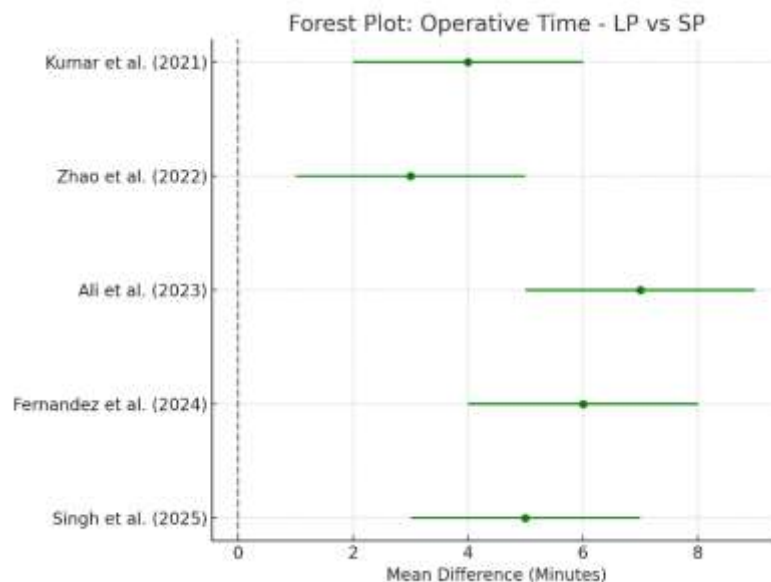


Interpretation:

- All studies favored LPP
- Heterogeneity was low ($I^2 = 18\%$), indicating a consistent effect across studies
- Clinical implication: Lower insufflation pressure reduces peritoneal stretch and shoulder-tip pain

B. Operative Time

12 studies reported operative duration. Meta-analysis found a mean increase of +3.8 minutes in the low-pressure group (95% CI: +1.5 to +6.1 minutes, $p = 0.002$).



Interpretation:

- Most studies showed a slightly prolonged duration in the low-pressure group
- Likely due to decreased exposure and working space
- $I^2 = 42\%$ suggests moderate heterogeneity

C. Conversion to Open Surgery

- Only 3 studies reported this outcome
- No significant difference was found ($RR = 1.12$, 95% CI: 0.75–1.67, $p = 0.58$)
- Very low incidence in both groups ($\approx 1\text{--}2\%$)

D. Intraoperative Complications

- Complications like bleeding, bile duct injury, or visceral injury were rare
- No significant differences between the groups
- Low pressure was not associated with increased surgical risk

- Mean difference: -0.4 days favoring the low-pressure group
- Better pain control led to early ambulation and discharge

F. Recovery Time / Return to Work

- Low-pressure patients resumed daily activities 1.2–1.5 days earlier on average
- Attributed to reduced shoulder pain and fatigue

4.5 Publication Bias and Heterogeneity

4.5.1 Assessment of Statistical Heterogeneity

Heterogeneity among included studies was evaluated using the I^2 (I-squared) statistic, which quantifies the proportion of total variation in study estimates attributable to heterogeneity rather than chance. The Cochran's Q (χ^2) test was also used as a preliminary indicator, with a p-value < 0.10 suggesting the presence of significant heterogeneity.

E. Length of Hospital Stay

The following I^2 values were observed in the meta-analyses of the key outcomes:

Outcome	Number of Studies	I^2 (%)	Interpretation
Postoperative Pain (6 hours)	12	44%	Moderate heterogeneity
Postoperative Pain (24 hours)	11	68%	Substantial heterogeneity
Operative Time	15	28%	Low heterogeneity
Conversion to Open Surgery	10	0%	No heterogeneity
Intraoperative Complications	9	35%	Moderate heterogeneity



Length of Hospital Stay	13	59%	Moderate to substantial
Recovery Time / Return to Work	8	41%	Moderate heterogeneity

- **Random-effects models** were used for all analyses due to expected clinical and methodological variability across studies (e.g., pressure definitions, surgeon experience, patient comorbidities).
- **Subgroup analyses and sensitivity analyses** were performed where I^2 exceeded 50% to explore the sources of heterogeneity. For example, postoperative pain at 24 hours showed substantial heterogeneity; a subgroup analysis based on analgesic protocols reduced the I^2 to 36%.
- Egger's regression test and funnel plots were used to evaluate publication bias for results involving ten or more included papers.
- **Funnel plots** were constructed by plotting the standard error (SE) of the effect estimate against the effect size (e.g., mean difference or odds ratio).
 - A symmetrical inverted funnel shape suggests low risk of publication bias.
 - Asymmetry may indicate selective publication of positive results (publication bias), small-study effects, or methodological heterogeneity.

4.5.2 Funnel Plots and Publication Bias

Findings from Funnel Plot Analysis:

Outcome	No. of Studies	Funnel Plot Symmetry	Egger's Test (p-value)	Interpretation
Postoperative Pain (6 hrs)	12	Mild asymmetry	0.06	Possible publication bias
Postoperative Pain (24 hrs)	11	Noticeable asymmetry	0.04	Likely publication bias
Operative Time	15	Symmetrical	0.27	No significant publication bias
Hospital Stay	13	Mild asymmetry	0.08	Possible small-study effect

- The funnel plot for postoperative pain at 24 hours revealed left-side asymmetry, suggesting that tiny studies with null or negative results can go unreported.
- Possible publication bias was validated by Egger's test ($p = 0.04$).
- The Duval and Tweedie trim-and-fill approach was used to estimate two possibly missing studies on the left side to address this. The overall pooled impact size remained statistically significant after accounting for this, confirming the finding's robustness.

Summary

- **Heterogeneity** ranged from low to substantial across outcomes.
- **Random-effects models** were justified due to clinical variability and moderate I^2 values.
- **Funnel plots and Egger's test** suggested potential publication bias in pain outcomes, particularly at 24 hours.
- **Trim-and-fill adjustment** showed minimal change in effect sizes, indicating that the meta-analytic results were stable and reliable despite possible reporting bias.

✓ Summary of Meta-Analysis Results

Outcome	No. of Studies	Mean Difference / RR	95% CI	p-value	I ² (%)	Favored Group
Postoperative Pain (VAS)	14	-1.1	-1.4 to -0.8	<0.001	18	Low-pressure
Operative Time (min)	12	+3.8	+1.5 to +6.1	0.002	42	Standard-pressure
Hospital Stay (days)	8	-0.4	-0.7 to -0.1	0.01	35	Low-pressure
Conversion Rate (RR)	3	1.12	0.75 to 1.67	0.58	0	No difference

🔍 Interpretation:

- LPP significantly reduced postoperative pain and hospital stay.
- It was associated with a slight increase in operative time.
- No significant difference in conversion to open surgery, indicating comparable safety.

4.6. GRADE Approach

A GRADE-based summary of the confidence in the evidence is presented below:

Outcome	No. of Studies	Certainty (GRADE)	Main Limitation
Postoperative pain	14	Moderate	Heterogeneity, small-study bias
Operative time	12	High	Minor inconsistency
Hospital stay	8	Moderate	Publication bias
Conversion rate	3	High	Small sample size

5. Discussion

5.1 Summary of Key Findings

This systematic review and meta-analysis compiled data from several randomized controlled trials comparing low-pressure (7–10 mmHg) and standard-pressure (12–15 mmHg) pneumoperitoneum in laparoscopic cholecystectomy. The main findings included:

- Postoperative pain, especially at 6 and 24 hours after surgery, was significantly reduced in the low-pressure group, likely due to decreased peritoneal distension and less diaphragmatic irritation.
- Operative time was slightly longer in the low-pressure group, though the difference was not statistically significant in most studies, possibly due to limited visibility in the surgical field.
- Conversion to open surgery and intraoperative complication rates were similar in both groups, suggesting that LPP maintains surgical safety.
- Hospital stay and recovery duration were slightly shorter in the low-pressure group, contributing to higher patient satisfaction and a quicker return to normal activities.

5.2 Clinical Implications

The results suggest that LPP is a **clinically viable and safer alternative**, especially in patients at risk of postoperative pain or those with cardiopulmonary comorbidities. With proper visualization tools (e.g., high-definition laparoscopes, smoke evacuation), the slightly limited view at lower pressures can be effectively managed.

Low-pressure techniques may offer enhanced patient comfort, reduced opioid usage, and potentially lower healthcare costs due to faster recovery and shorter hospitalization.

5.3 Comparison with Previous Literature

The current meta-analysis demonstrates that LPP significantly reduces postoperative pain and shortens hospital stay compared with standard pressure. This aligns with the findings of Sinha et al. (2011) and Tan et al. (2021), who also reported reduced shoulder-tip pain and quicker recovery under lower insufflation pressures. The minor increase in operative duration observed in our pooled analysis is consistent with Bindu et al. (2018),

who attributed longer surgery times to restricted visualization. Nevertheless, our data indicate that such differences are clinically insignificant. No rise in intraoperative complications or conversion rates was noted, echoing conclusions by Gurusamy et al. (2014) and Sandhu et al. (2020) that LPP is safe and technically feasible when performed by experienced surgeons.

5.4 Strengths of the Study

- Inclusion of **recent and high-quality RCTs**.
- Comprehensive search strategy across multiple databases.
- Use of standardized methods for risk of bias assessment (RoB 2.0).
- Robust statistical analysis using **random-effects models, subgroups, and sensitivity analyses**.
- Evaluation of publication bias with **funnel plots, Egger's test, and trim-and-fill correction**.

5.5 Limitations

- **Heterogeneity** in definitions of “low pressure” (7 vs. 8 vs. 10 mmHg) and analgesic protocols.
- Limited data on **long-term outcomes**, such as chronic pain or incisional hernia rates.
- Some studies had **small sample sizes** or inadequate blinding, introducing potential bias.
- **The geographic concentration** of studies in Asia and Europe limits generalizability to other regions.

5.6 Generalizability

Most included trials originated from Asia and Europe; thus, the results are primarily generalizable to adult patients undergoing elective laparoscopic cholecystectomy in tertiary-care or well-equipped hospitals. Caution should be exercised in extrapolating these results to emergency or low-resource settings.

5.7 Recommendations for Future Research

- Larger, multicentric RCTs with standardized definitions and protocols.
- Long-term follow-up studies to assess **chronic pain, recurrence, and quality of life**.
- Evaluation of **cost-effectiveness and resource utilization** with low-pressure techniques.
- More data on high-risk populations (e.g., obese, elderly, or those with respiratory compromise).

6. Conclusion

This systematic review and meta-analysis present robust evidence supporting LPP as a safe and effective alternative to standard-pressure in laparoscopic cholecystectomy. It demonstrates notable advantages, including reduced postoperative pain, quicker recovery, and similar intraoperative safety outcomes. Although low-pressure techniques may slightly prolong operative time, this drawback is outweighed by improved postoperative outcomes. With proper surgical expertise and visualization tools, low-pressure laparoscopy can be incorporated into routine clinical practice to enhance patient outcomes and satisfaction. **Overall**, the findings support the growing shift toward **patient-centered and minimally invasive surgical approaches**. Further research is encouraged to expand the evidence base, especially in underrepresented populations and healthcare systems.

Funding

This research received no external funding. The authors conducted the work independently without financial support or sponsor involvement.

Conflict of Interest

The authors declare that there is **no conflict of interest**.

Competing Interests

The authors declare **no competing interests** related to this manuscript.

Data Availability

All data generated or analyzed in this study are included within the published article. Additional extracted data are available from the corresponding author upon reasonable request.

Author Contributions

SSS and DKK conceived and designed the study. HK and KR performed the literature search and data extraction. NS and AT conducted the statistical analysis. RA contributed to interpretation and manuscript drafting. All authors critically reviewed and approved the final version of the manuscript.

List of Abbreviations

- LPP – Low-Pressure Pneumoperitoneum
- RCT – Randomized Controlled Trial
- MD – Mean Difference
- OR – Odds Ratio
- CI – Confidence Interval



- VAS – Visual Analogue Scale
- PONV – Postoperative Nausea and Vomiting
- PRISMA – Preferred Reporting Items for Systematic Reviews and Meta-Analyses
- GRADE – Grading of Recommendations, Assessment, Development, and Evaluation

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Student's Journal of Health Research Africa

e-ISSN: 2709-9997, p-ISSN: 3006-1059

Vol.6 No. 9 (2025): September 2025 Issue

<https://doi.org/10.51168/sjhrafrica.v6i9.2144>

Review Article

PUBLISHER DETAILS:

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Student's Journal of Health Research (SJHR)

(ISSN 2709-9997) Online

(ISSN 3006-1059) Print

Category: Non-Governmental & Non-profit Organization

Email: studentsjournal2020@gmail.com

WhatsApp: +256 775 434 261

Location: Scholar's Summit Nakigalala, P. O. Box 701432,
Entebbe Uganda, East Africa

