Non-invasive pulse wave analysis for monitoring the cardiovascular effects of CO₂ pneumoperitoneum during laparoscopic cholecystectomy- a prospective case-series study

Péter Sárkány¹, Szabolcs Lengyel², Réka Nemes¹, Lívia Orosz¹, Dénes Páll², Csilla Molnár¹ and Béla Fülesdi¹*

Abstract

Background: Peritoneal insufflation results in hemodynamic changes during laparoscopic cholecystectomy. The aim of the present work is to test whether non-invasive applanation tonometry is suitable for reflecting these hemodynamic alterations.

Methods: 41 patients undergoing laparoscopic cholecystectomies were monitored using the SphygmoCor pulse wave analysing system. Peripheral blood pressures (PBP), central aortic blood pressures (CBP), augmentation index (ALX@HR75) and subendocardial viability ratio (SVR) were measured at rest (Phase 1), after anesthetic induction (Phase 2), after peritoneal inflation (Phase 3) and after peritoneal deflation (Phase 4).

Results: Induction of anesthesia resulted in a statistically significant reduction in both the peripheral blood pressure and central aortic pressures, accompanied by a decrease in augmentation pressure and augmentation index. Peripheral blood pressures did not change along with the peritoneal cavity insufflation, except for a moderate increase in systolic blood pressure. In contrast to this, an increase could be observed in central aortic pressure (106.77 ± 18.78 vs. 118.05 ± 19.85 mmHg, P < 0.01) which was accompanied by increased augmentation pressure (18.97 ± 10.80 vs. 31.55 ± 12.01; P < 0.001) and augmentation index (7.31 ± 5.59 vs. 12.61 ± 7.56, P < 0.001), indicating a rise in peripheral arterial stiffness.

Conclusions: The Sphigmocor pulse wave analysis system can be reliably used for detecting and monitoring cardiovascular changes occurring during laparoscopic cholecystectomy.

Keywords: Laparoscopic cholecystectomy, Hemodynamic changes, Applanation tonometry

Background

It has been proven by previous studies that inducing a positive pressure within the intraperitoneal cavity during laparoscopic surgical interventions results in numerous cardiovascular, neuroendocrine and renal changes [1]. These changes include an increase in systemic and pulmonary vascular resistance and a consequent decrease in cardiac output, which may be attributed to direct mechanical factors due to intraperitoneal pressure rise as well as to humoral changes evoked by the procedure [2].

During preoperative anesthesiological consultation it is a frequent question whether laparoscopic cholecystectomy can be performed safely in patients with known cardiovascular risk factors. It is worth mentioning that in a study of low risk patients undergoing laparoscopic cholecystectomy, 2 out of 16 patients had acute ST changes on their ECG [3]. In view of this, preoperative cardiovascular risk stratification as well as proper intraoperative monitoring may be of high importance in patients at risk.

So far, cardiovascular consequences occurring during laparoscopic procedures have been assessed either during animal experiments [4] or by using invasive intraoperative
hemodynamic monitoring of humans [5-9]. However, invasive hemodynamic monitoring techniques may also have side effects and thus are not indicated in all patients undergoing otherwise relatively short and low risk surgical procedures.

Along these lines, we tested the hypothesis that cardiovascular changes caused by CO₂-pneumoperitoneum may be accurately assessed intraoperatively by non-invasive applanation tonometry. Our results were compared with data obtained from the literature.

Patients and methods

Patients

A total of 41 consecutive patients undergoing elective laparoscopic cholecystectomy for symptomatic cholelithiasis without cholangiography or cholecystotomy were enrolled in this prospective case-series. The patients were all in good health, classified as ASA I and II. This study was conducted with approval from the University of Debrecen Medical Ethics Committee (Registration number: DEOEC RKET/IKET 2312-2010, responsible person: József Szentmiklósí; Department of Pharmacology, University of Debrecen, 98.Nagyerdei krt. Debrecen, Hungary, Phone: +3652411600). A written informed consent to participate was obtained from all patients included in the study. Patients with diabetes mellitus, untreated hypertension, atrial fibrillation, morbid obesity (body mass index [BMI] > 35), infection, psychiatric or neurologic conditions impairing patients’ ability to cooperate were excluded from the study.

Anesthesia and CO₂ pneumoperitoneum

General anesthesia was administered to all patients according to the same protocol. During the patient’s stay in the preparation area, 15 ml/kg/BW of Ringer’s acetate solution was infused for a period of 2 hours. As premedication, oral midazolam (0.15 mg/kgBW) was administered 30 minutes before the induction of anaesthesia. Following preoxygenation (2 minutes) by face mask, anesthesia was induced with intravenous propofol (2 mg/kg BW) and fentanyl (3-5 μg/kgBW). Rocuronium 0.6 mg/kgBW was used to facilitate tracheal intubation and maintain muscle relaxation. After intubation, the lungs were ventilated with a mixture of air/oxygen (50/50%). For maintenance of anesthesia sevoflurane (2 vol%) and intermittent doses of fentanyl were applied. Sevoflurane was titrated in order to keep the bispectral index (BIS) values between 40 and 50. Ventilation was mechanically controlled at a frequency and tidal volume sufficient for maintaining normocapnia. PEEP was not administered (ZEEP). End expiratory CO₂ was used to ensure normoventilation (end-tidal carbon dioxide level was kept between 35 and 38 mmHg). Intraoperative crystalloid infusion was administered at 7 mL/kg per hour. During anesthesia body temperature was maintained between 36.0 and 36.5°C by heating blankets.

The surgical technique was similar for all patients. CO₂ intraperitoneal pressure was maintained automatically at a recommended 12-14 mmHg by a CO₂ insufflator at an insufflation rate of 1 to 1.5 L/min with the patients placed in the 20° reverse Trendelenburg position (rT). On confirming the appropriate placement of the video laparoscope, each patient’s position was changed to a left lateral tilt (10°-15°). Once the surgery was completed, the abdomen was deflated and each patient was returned to the horizontal position.

Routine intraoperative patient monitoring included continuous five-lead electrocardiography, pulse oximetry, non-invasive blood pressure measurements, peak airway pressures, capnography, as well as BIS monitoring for assessment of depth of anesthesia. Neuromuscular monitoring was performed to control the neuromuscular block throughout the course of anesthesia using TOF Watch SX acceleromyograph.

Monitoring cardiovascular function

The SphygmoCor pulse wave analysing system was used for monitoring cardiovascular function, which is a non-invasive method based on applanation tonometry [10]. During the present study we measured systemic and central aortic pressure, augmentation pressure, augmentation index, ejection duration and subendocardial viability ratio.

- **Measurement of central aortic pressure and aortic pressure waveform**: A conventional cuff pressure measurement was used for calibration. After applanation tonometry SphygmoCor derived a complete waveform for the whole cardiac cycle for the aortic pulse. A combination of the two methods makes it possible to analyse the coupling between the ejecting heart and the pressure load.
- **Measurement of augmentation pressure and augmentation index**: Augmentation pressure is based on the principle that there is a reflected pressure from the periphery that appears in the aortic pressure waveform. The amount of augmentation reflects the stiffness of the peripheral arterial tree; it increases along with higher stiffness. In order to make the value of augmentation index independent from the individual changes of pulse rate, the device calculates a corrected augmentation index (ALX@HR75).
- **Subendocardial viability ratio**: This parameter is calculated by the device by dividing the area under the systolic and diastolic part of the curve. A ratio under 100% reflects underperfusion of the subendocardium.
Hemodynamic measurements were repeated in different phases of the procedure: Before induction of anesthesia (resting phase, Time 1); 5 minutes after induction of anesthesia (Time 2); 5 minutes after inflation of the peritoneal space (Time 3) followed by repeated measurements every 10 minutes and 5 minutes after deflation of the peritoneal cavity (Time 4).

Statistical analysis
Means and standard deviation were calculated for all values. Repeated measure analysis of variance was used for all values to check the time main effect of the laparoscopic procedure, i.e. whether laparoscopic cholecystectomy overall had any significant hemodynamic effect. Pairwise comparisons of all parameters were performed in order to check the effect of inflation and deflation of the peritoneal space by taking the values obtained after induction of anesthesia as reference value. A p < 0.05 was considered as a statistically significant difference.

Results
A total of 41 patients entered the study. There were 33 females and 8 males with an average age of 52.3 ± 15.4 years.

The effect of anesthetic induction on hemodynamic parameters
As shown in Table 1, induction of anesthesia resulted in a statistically significant reduction in both peripheral blood pressure and central aortic pressures, accompanied by a decrease in augmentation pressure and augmentation index.

| Parameter                        | Before induction | After induction | p-value |
|----------------------------------|------------------|----------------|---------|
| **Peripheral blood pressure**    |                  |                |         |
| Systolic                         | 132.47 ± 18.87   | 116.80 ± 18.61 | P < 0.001|
| Diastolic                         | 78.60 ± 10.40   | 72.71 ± 13.54  | 0.01    |
| Pulse pressure                    | 52.12 ± 16.12   | 43.65 ± 11.96  | P < 0.01|
| **Central (aortic) blood pressure** |                |                |         |
| Systolic                         | 120.80 ± 19.10  | 106.77 ± 18.78 | P < 0.001|
| Diastolic                         | 80.00 ± 10.38   | 73.88 ± 13.79  | P < 0.01|
| Pulse pressure                    | 41.05 ± 14.70   | 33.11 ± 11.08  | P < 0.001|
| Ejection duration                 | 41.0 ± 4.93     | 38.40 ± 6.71   | 0.01    |
| **Pressure augmentation**        |                  |                |         |
| Augmentation pressure            | 1052 ± 8.52     | 731 ± 5.59     | P < 0.01|
| Augmentation index                | 23.62 ± 10.58   | 18.97 ± 10.80  | P < 0.01|
| Subendocardial viability ratio (%)| 121.85 ± 22.7   | 142.5 ± 38.2   | P < 0.01|

Means and standard deviations are shown.

The effect of peritoneal insufflation on hemodynamic parameters
Table 2 summarizes the parameters that were obtained before and after peritoneal cavity insufflation. Peripheral blood pressures did not change markedly along with peritoneal cavity insufflation, except for a moderate increase in systolic blood pressure. In contrast to this, a marked increase could be observed in central aortic pressure values which was accompanied by increased augmentation pressure and augmentation index, indicating a rise in peripheral arterial stiffness. Despite changes in the central aortic blood pressure, subendocardial viability ratio remained relatively stable during and after peritoneal cavity insufflation.

Comparison of hemodynamic parameters at preinduction and peritoneal insufflation phase
Peripheral and central blood pressures returned to the pre-induction values after peritoneal insufflation (comparison of Phase 1 and 3). The only parameter that showed a gradual increase was augmentation index (Table 3).

After deflation of the abdominal cavity both peripheral and aortic pressure values returned to the levels observed after induction of anesthesia. Although augmentation pressures were still higher than before inflation, augmentation index (the main indicator of peripheral arterial stiffness) also returned to the pre-insufflation value. Figure 1 depicts and summarizes the changes of all parameters during the entire course of the study.

Discussion
In this cohort study we used a new non-invasive technique for assessing the cardiovascular changes that occur during laparoscopic cholecystectomy, the Sphigmocor pulse wave

Table 1 The effect of anesthetic induction on peripheral and central (aortic) blood pressures, ejection duration and pressure augmentation

| Parameter                        | Before induction | After induction | p-value |
|----------------------------------|------------------|----------------|---------|
| **Peripheral blood pressure**    |                  |                |         |
| Systolic                         | 132.47 ± 18.87   | 116.80 ± 18.61 | P < 0.001|
| Diastolic                         | 78.60 ± 10.40   | 72.71 ± 13.54  | 0.01    |
| Pulse pressure                    | 52.12 ± 16.12   | 43.65 ± 11.96  | P < 0.01|
| **Central (aortic) blood pressure** |                |                |         |
| Systolic                         | 120.80 ± 19.10  | 106.77 ± 18.78 | P < 0.001|
| Diastolic                         | 80.00 ± 10.38   | 73.88 ± 13.79  | P < 0.01|
| Pulse pressure                    | 41.05 ± 14.70   | 33.11 ± 11.08  | P < 0.001|
| Ejection duration                 | 41.0 ± 4.93     | 38.40 ± 6.71   | 0.01    |
| **Pressure augmentation**        |                  |                |         |
| Augmentation pressure            | 1052 ± 8.52     | 731 ± 5.59     | P < 0.01|
| Augmentation index                | 23.62 ± 10.58   | 18.97 ± 10.80  | P < 0.01|
| Subendocardial viability ratio (%)| 121.85 ± 22.7   | 142.5 ± 38.2   | P < 0.01|

Means and standard deviations are shown.
The method has already been tested in different clinical conditions such as arterial hypertension, diabetes mellitus, systolic heart failure and preeclampsia [11-13]. In anesthesiological practice, this is the first report on the use of the technique.

Similar to previous reports we were able to detect a significant decline of the systemic blood pressure after anesthetic induction which was accompanied by a decrease in augmentation index, reflecting the stiffness of the peripheral vessels [3,5,9,14,15]. This initial reduction in blood pressure and peripheral resistance may be due to the direct myocardial depressant and vasodilatory effects of the anesthetics together with the loss of sympathetic tone [2]. During the next phase of the procedure, after inflating the abdomen and tilting the patient to a reverse Trendelenburg position, the most important observations were increases in central aortic pressures accompanied by an increase of augmentation index. This is in line with previous observations reporting on an increase in mean arterial pressure and peripheral resistance [1,2,5,9,14] after peritoneal insufflation. To transform these results to our observation, we have to mention that systolic central aortic pressure increased by 10.6%, whereas augmentation index referring to peripheral resistance increased by 66% on average after peritoneal insufflation. It has to be noted that hemodynamic parameters in this phase returned to the baseline, preinduction values (Table 3). It seems that the effect of inducing pneumoperitoneum counteracts the

Table 2 The effect of peritoneal cavity insufflation on peripheral and central (aortic) blood pressures, ejection duration and pressure augmentation

|                         | Before insufflation | After insufflation | p-value |
|-------------------------|---------------------|--------------------|---------|
| **Peripheral blood pressure** |                     |                    |         |
| Systolic                | 116.80 ± 18.61      | 125.17 ± 20.21     | 0.02    |
| Diastolic               | 72.71 ± 13.54       | 78.92 ± 16.94      | 0.07    |
| Pulse pressure          | 43.65 ± 11.96       | 44.64 ± 13.23      | 0.24    |
| **Central (aortic) blood pressure** |                     |                    |         |
| Systolic                | 106.77 ± 18.78      | 118.05 ± 19.85     | P < 0.01|
| Diastolic               | 73.88 ± 13.79       | 81.82 ± 12.28      | P < 0.01|
| Pulse pressure          | 33.11 ± 11.08       | 36.41 ± 12.60      | 0.04    |
| Ejection duration       | 38.40 ± 6.71        | 39.15 ± 5.51       | 0.34    |
| **Pressure augmentation** |                     |                    |         |
| Augmentation pressure   | 7.31 ± 5.59         | 12.61 ± 7.56       | P < 0.001|
| Augmentation index      | 18.97 ± 10.80       | 31.55 ± 12.01      | P < 0.001|
| Subendocardial viability ratio (%) | 142.5 ± 38.2 | 137.11 ± 26.3 | P = 0.48 |

Means and standard deviations are shown.

Table 3 Comparison of peripheral and central (aortic) blood pressures, ejection duration and pressure augmentation values at phases before induction and after peritoneal insufflation

|                         | Before induction | After insufflation | p-value |
|-------------------------|------------------|--------------------|---------|
| **Peripheral blood pressure** |                   |                    |         |
| Systolic                | 132.47 ± 18.87   | 125.17 ± 20.21     | 0.10    |
| Diastolic               | 78.60 ± 10.40    | 78.92 ± 16.94      | 0.91    |
| Pulse pressure          | 52.12 ± 16.12    | 44.64 ± 13.23      | 0.02    |
| **Central (aortic) blood pressure** |                 |                    |         |
| Systolic                | 120.80 ± 19.10   | 118.05 ± 19.85     | 0.53    |
| Diastolic               | 80.00 ± 10.38    | 81.82 ± 12.28      | 0.47    |
| Pulse pressure          | 41.05 ± 14.70    | 36.41 ± 12.60      | 0.13    |
| Ejection duration       | 41.0 ± 4.93      | 39.15 ± 5.51       | 0.11    |
| **Pressure augmentation** |                  |                    |         |
| Augmentation pressure   | 10.52 ± 8.52     | 12.61 ± 7.56       | 0.25    |
| Augmentation index      | 23.62 ± 10.58    | 31.55 ± 12.01      | <0.01   |
| Subendocardial viability ratio (%) | 121.85 ± 22.7 | 137.11 ± 26.3 | <0.01   |

Means and standard deviations are shown.
Figure 1 Changes of hemodynamic parameters during the study: Means and standard deviations are shown. \( \times \times \) indicate \( p < 0.01 \), \( \times \times \times \) indicate \( p < 0.001 \) compared to preinduction value; \( \ddagger \ddagger \) \( p < 0.01 \), \( \ddagger \ddagger \ddagger \) indicate \( p < 0.001 \) statistical difference compared to preinsufflation value.
hemodynamic depressant effects of the anesthetics suggesting that changes in ASA I-II patients are clinically most probably not relevant. In a recent study, Cinnella and co-workers also demonstrated that hemodynamic stability after administering pneumoperitoneum is maintained even if moderate (5 cm H2O) is applied [16].

In a previous review Wahba an co-workers [2] summarized the hemodynamic effects and suggested that direct mechanical, neurohumoral processes play a role, slightly modified by the effect of the resorbed CO2 during pneumoperitoneum. Mechanical effects of pneumoperitoneum may decrease renal flow, activating the renin-angiotensine-aldosteron system, may result in the compression of the abdominal veins and the aorta. It has also been proven that inducing pneumoperitoneum leads to an increased production of vasopressin, adrenalin, noradrenalin, renin and cortisol, which is in correlation with the changes of mean arterial pressure and systemic resistance [15]. According to previous reports, this is the phase of laparoscopic cholecystectomy where patients of different ASA severity (ASA I-II vs. ASA III-IV) may react differently to pneumoperitoneum. In more severe patients (ASA III-IV) a pressure rise in the abdomen resulted in a more pronounced increase in mean arterial pressure and decreased oxygen delivery [17]. Consequently, left ventricular stroke work index increases, which causes higher oxygen demand of the myocardium [7]. In our series we included ASA I and II patients and no subendocardial viability ratio reflecting the potential underperfusion of the subendocardium could be detected after and during the course of abdominal inflation.

The principal basis of the pulse wave analysis system is that the peripheral arterial pressure waveform may be used for the reconstruction of central (aortic) pressure. The method behind this is applanation tonometry, which ensures the sensitive detection of the radial artery pulse waveform. It is generally accepted that the characteristics of the peripheral pulse reflect the changes in arterial diameters, wall elasticity, wall thickness and the condition of the peripheral vascular beds. The main attribute of SphygmoCor is its ability to derive the central aortic pressure waveform non-invasively from the pressure pulse recorded at a peripheral site, usually at the upper arm (radial artery) [10].

Limitations

The intraoperative use of the device is limited by the position of the radial artery, i.e. in some surgical scenarios it may disturb the surgical team, making monitoring impossible. Another limitation to be mentioned is operator-dependency: for reliable monitoring it is necessary to have previous experience with the technique. Finally, the main limitation of this study is the lack of a control group, i.e. other hemodynamic measurements were not used. However, this trial is a pilot application of applanation tonometry in this field. As this is a non-invasive method, in this first, pilot step of our investigations we intended to compare hemodynamic changes with those that used invasive monitoring techniques reported in the literature.

Conclusion

In conclusion: in this study we have shown that the SphygmoCor pulse wave analysis system can be reliably used for detecting and monitoring cardiovascular changes occurring during laparoscopic cholecystectomy. Further studies are needed to prove whether the method may be helpful in delineating critical situations in patients with limited cardiovascular reserve (ASA III-IV patients) by defining cut-off values of safety and to help in guiding abdominal insufflation and tilting during the procedure as suggested in previous reports [18].

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

PS: Intraoperative measurements, drafting of the manuscript. SL: Intraoperative measurements, statistical analysis. LO: Intraoperative measurements, drafting of the manuscript. DP: Study protocol consultation. CM: Study protocol, drafting of the manuscript. BF: Study protocol, drafting of the manuscript. All authors read and approved the final manuscript.

Author details

1Department of Anesthesiology and Intensive Care, University of Debrecen, Medical and Health Science Centre, Nagyerdei krt. 98, H-4032 Debrecen, Hungary. 21st Department of Medicine, University of Debrecen, Medical and Health Science Centre, Nagyerdei krt. 98, H-4032 Debrecen, Hungary.

Received: 1 April 2014 Accepted: 30 September 2014

Published: 31 October 2014

References

1. Struthers AD, Cuschieri A: Cardiovascular consequences of laparoscopic surgery. Lancet 1998, 352:568–570.
2. Wahba RW, Béïque F, Kleinman SL: Cardiopulmonary function and laparoscopic cholecystectomy. Can J Anaesth 1995, 42:51–63.
3. O’Leary E, Hubbard K, Tomney W, Cunningham AJ: Laparoscopic cholecystectomy: haemodynamic and neuroendocrine responses after pneumoperitoneum and changes in position. Br J Anaesth 1996, 76:640–644.
4. Blachereine F, Machado SB, Fonseca EB, Ottuque D, Auler JO Jr, Richard F: Pulse pressure variation as a tool to detect hypovolaemia during pneumoperitoneum. Acta Anaesthesiol Scand 2007, 51:1269–1272.
5. Turkistani AA: Cardiodynamic monitoring during laparoscopic cholecystectomy. Middle East J Anaesthesiol 2005, 18:435–439.
6. Galizia G, Prizio G, Lieto E, Castellano P, Pelosi L, Imperatore V, Ferrara A, Pignatelli C: Hemodynamic and pulmonary changes during open, carbon dioxide pneumoperitoneum and abdominal wall-lifting cholecystectomy. A prospective, randomized study. Surg Endosc 2001, 15:477–483.
7. Treig B, Berger DH, Dupuis JF: Hemodynamic effects of CO2 abdominal insufflation (CAI) during laparoscopy in high-risk patients. Anesth Analg 1994, 78:5109.
8. Joshi GP, Hein HA, Macarenhas WL, Ramsay MA, Bayer O, Klotz P: Continuous transesophageal echo-Doppler assessment of hemodynamic function during laparoscopic cholecystectomy. J Clin Anesth 2005, 17:17–121.
9. Joris JL, Nointot DP, Legrand MJ, Jacquet NI, Lamy ML: Hemodynamic changes during laparoscopic cholecystectomy. Anesth Analg 1993, 76:1067–1071.
10. O’Rourke MF, Pauca AL: Augmentation of the aortic and central arterial pressure waveform. Blood Press Monit 2004, 9:179–185.
11. Weber T, Auer J, O’Rourke MF, Kvas E, Lassnig E, Lamm G, Stark N, Rammer M, Eber B: Increased arterial wave reflections predict severe cardiovascular events in patients undergoing percutaneous coronary interventions. Eur Heart J 2005, 26:2657–2663.

12. Agnoletti D, Lieber A, Zhang Y, Protogerou AD, Borghi C, Blacher J, Safar ME: Central hemodynamic modifications in diabetes mellitus. Atherosclerosis 2013, 230:315–321.

13. Carty DM, Neisius U, Rooney LG, Dominiczak AF, Delles C: Pulse wave analysis for the prediction of preeclampsia. J Hum Hypertens 2014, 28:98–104.

14. Hömme R: Anesthesia for laparoscopic interventions (German). Anaesthesist 2011, 60:175–187.

15. Joris JL, Chiche JD, Canivet JL, Jacquet NJ, Legros JJ, Lamy ML: Hemodynamic changes induced by laparoscopy and their endocrine correlates: effects of clonidine. J Am Coll Cardiol 1998, 32:1389–1396.

16. Cinnella G, Grasso S, Spadaro S, Rauseo M, Mirabella R, Salatto P, De Capraris A, Napoli L, Greco P, Dambrosio M: Effects of Recruitment Maneuver and Positive End-expiratory Pressure on Respiratory Mechanics and Transpulmonary Pressure during Laparoscopic Surgery. Anesthesiology 2013, 118:114–122.

17. Safran D, Spambati S, Orlando R 3rd: Laparoscopy in high-risk cardiac patients. Surg Gynecol Obstet 1993, 176:548–554.

18. Dhoste K, Lacoste L, Karayan J, Lebeau MS, Thomas D, Fusciardi J: Haemodynamic and ventilatory changes during laparoscopic cholecystectomy in elderly ASA III patients. Can J Anaesth 1996, 43:783–788.

doi:10.1186/1471-2253-14-98
Cite this article as: Sárkány et al.: Non-invasive pulse wave analysis for monitoring the cardiovascular effects of CO2 pneumoperitoneum during laparoscopic cholecystectomy- a prospective case-series study. BMC Anesthesiology 2014 14:98.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at www.biomedcentral.com/submit