The association of acute hypercarbia and plasma potassium concentration during laparoscopic surgery: a retrospective observational study

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Abstract

Background: It is uncertain whether increases in PaCO\textsubscript{2} during surgery lead to an increase in plasma potassium concentration and, if so, by how much. Hyperkalaemia may result in cardiac arrhythmias, muscle weakness or paralysis. The key objectives were to determine whether increases in PaCO\textsubscript{2} during laparoscopic surgery induce increases in plasma potassium concentrations and, if so, to determine the magnitude of such changes.

Methods: A retrospective observational study of adult patients undergoing laparoscopic abdominal surgery was performed. The independent association between increases in PaCO\textsubscript{2} and changes in plasma potassium concentration was assessed by performing arterial blood gases within 15 min of induction of anaesthesia and within 15 min of completion of surgery.

Results: 289 patients were studied (mean age of 63.2 years; 176 [60.9%] male, and mean body mass index of 29.3 kg/m\textsuperscript{2}). At the completion of the surgery, PaCO\textsubscript{2} had increased by 5.18 mmHg (95% CI 4.27 mmHg to 6.09 mmHg) compared to baseline values (P < 0.001) with an associated increase in potassium concentration of 0.25 mmol/L (95% CI 0.20 mmol/L to 0.31 mmol/L, P < 0.001). On multiple regression analysis, PaCO\textsubscript{2} changes significantly predicted immediate changes in plasma potassium concentration and could account for 33.1% of the variance ($r^2 = 0.331$, f(3,259) = $38.915$, P < 0.001). For each 10 mmHg increment of PaCO\textsubscript{2} the plasma potassium concentration increased by 0.18 mmol/L.

Conclusion: In patients receiving laparoscopic abdominal surgery, there is an increase in PaCO\textsubscript{2} at the end of surgery, which is independently associated with an increase in plasma potassium concentration. However, this effect is small and is mostly influenced by intravenous fluid therapy (Plasma-Lyte 148 solution) and the presence of diabetes.

Trial registration Retrospectively registered in the Australian New Zealand Clinical Trials Registry (Trial Number: ACTRN12619000716167).

Keywords: Anaesthesia, Potassium, Hypercarbia, Laparoscopic, Surgery

Background

Increased partial pressure of arterial carbon dioxide (PaCO\textsubscript{2}) (hypercarbia) is common in the setting of mechanical ventilation and surgery [1, 2]. The effect of hypercarbia and respiratory acidosis on plasma
potassium concentrations ([K⁺]ᵢ) is unclear with some studies showing an increase in concentration and others showing no effect [3–10]. Further, a body of longstanding evidence indicates that the hyperkalaemia-acidaemia relationship is more complex than the relatively simplistic, but commonly accepted notion that hyperkalaemia develops due to an increase in extracellular acidity and subsequent exchange of extracellular hydrogen ions for intracellular potassium ions [8]. Challenging this theory, early studies have shown that the directional flux of potassium during acute acid–base disorders is not uniform among various tissues [11–13]. During acute respiratory acidosis, potassium was shown to move extracellularly in skeletal muscle [12, 14, 15] and liver tissue [16, 17], but intracellularly in cardiac tissue [11, 15, 18, 19]. Whilst mild hyperkalaemia intraoperatively is usually asymptomatic, high plasma levels of potassium may result in cardiac arrhythmias, muscle weakness or paralysis. Understanding this relationship has specific implications for the prevention of severe hyperkalaemia and the immediate monitoring and management of hyperkalaemia in the intra- and postoperative periods, as timely recognition and treatment of complications that arise from hyperkalaemia is imperative.

These conflicting findings, together with a lack of information from large scale studies concerning the quantitative and temporal relationships between changes in PaCO₂ and [K⁺]ᵢ, have clinical relevance during anaesthesia. Accordingly, a retrospective study was performed to describe the association between changes in PaCO₂ and acute changes in [K⁺]ᵢ in patients undergoing laparoscopic surgery. Laparoscopic surgery is a well-described model of carbon dioxide absorption and subsequent hypercarbia during anaesthesia [1, 2, 20–22]. The following hypotheses were made. First, in alignment with the current literature [1, 2], patients undergoing laparoscopic surgery would develop acute respiratory acidosis. Second, the development of any respiratory acidosis would cause acute respiratory acidemia. Finally, there would be an acute rise in [K⁺]ᵢ in response to the respiratory acidemia. To test these hypotheses, patients undergoing extraperitoneal and intraperitoneal laparoscopic surgery were retrospectively studied in a university hospital.

**Methods**

The Austin Health Human Research Ethics Committee approved this study (LNR/19/Austin/41) and provided a waiver for participant consent. Between February 2015 and February 2019, data was collected from the medical records of all patients undergoing extraperitoneal and intraperitoneal laparoscopic abdominal surgery at university hospital. Patients were included if they underwent surgery of greater than two hours duration, received an arterial line as part of anaesthesia care and were hospitalised for at least one postoperative night. To accurately compare changes in PaCO₂ and [K⁺]ᵢ during surgery to baseline values, data was collected from patients who had an arterial blood gas sampled within 15 min of induction of anaesthesia and a subsequent arterial blood gas sampled within 15 min of completion of surgery. The study was registered the study with the Australian New Zealand Clinical Trials Registry (Trial number ACTRN12619000716167; registered 13th May 2019; available at [http://www.anzctr.org.au/Trial/Registration/TrialReview.aspx?id=377303&showOriginal=true&isReview=true](http://www.anzctr.org.au/Trial/Registration/TrialReview.aspx?id=377303&showOriginal=true&isReview=true)).

Conduct of anaesthesia, was at the discretion of the treating anaesthetist and in accordance with existing protocols for patients undergoing major laparoscopic surgery at our institution. All patients received balanced crystalloid fluid therapy with the buffered fluid Plasma-Lyte 148 solution (Baxter Healthcare, Toongabbie, Australia). The physiochemical and electrolyte composition of Plasma-Lyte 148 is similar to human plasma, with a pH of 7.4 and a potassium concentration of 5 mmol/L.

During surgery, the handling and transportation of blood gas aspirates were performed using a standard hospital sampling protocol. A 3–5 mL sample of arterial blood was slowly aspirated into a 5 mL discard syringe minimising any force or strain during the aspiration process. Then, 1.0 mL of arterial blood was aspirated from the arterial line over a 2 s period and placed into a safePICO™ blood gas aspirator (Radiometer Medical Aps, Brønshøj, Denmark) containing 80 IU electrolyte balanced heparin. This blood sampling technique is designed to minimise any in-vitro haemolysis, which can falsely increase values of plasma constituents, especially potassium. The safePICO aspirator has a clear 1.0 mL label designed for accurate blood sampling, which is necessary to produce reliable results.

The blood sample syringe was transported horizontally and at standard operating room temperature (22 °C) to an automated blood gas analyser (ABL800, Radiometer, Denmark). [K⁺]ᵢ, PaCO₂, and pH were measured directly by the analyser and automatically entered into the patient’s electronic medical record. The same blood gas analyser was used for all measurements with regular calibration performed to ensure consistency in sample analysis. As a result, high analytical performance in the measurement of pH, PaCO₂, [K⁺]ᵢ and oximetry parameters were obtained. The system uses an automatic mixing system to obtain a homogenous sample for correct results thereby avoiding vigorous manual mixing, which can also lead to a haemolysed blood gas sample. The analyser prewarmed all samples to
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37 °C prior to measurement, and the alpha-stat model was applied for all blood gases analyses.

The primary aim of the study was to describe the relationship between \( P_{aCO2} \) and \([K^+]_p\). To investigate the independent association between acute hypercarbia and hyperkalaemia, we collected the following preoperative patient variables: age (years), gender, body mass index (kg/m²), current smoker status, ASA physical status classification, history of obstructive pulmonary lung disease, presence of other comorbidities, and surgery urgency. Intraoperative variables collected included the type of laparoscopic surgery, duration of surgery and amount of intraoperative fluid administered. We also collected the ARISCAT score, adjusted for the laparoscopic surgery, which predicts the risk of pulmonary complications after surgery [23, 24].

Inferential statistics were performed with IBM SPSS Statistics for Windows, version 23 (IBM Corp., Armonk, NY, USA). Missing value assessments were performed with R software 3.5.2 (R Development Core Team, Vienna, Austria, 2018) using “mice” and “missMDA” packages. Figure 1 was created using MS Word 365 (16.0.13127.20402) and Fig. 2 was created using SigmaPlot for Windows version 12.0 (Systat Software Inc., 2011, U.S.A).

For all continuous variables, data exploration for outliers and missing values was performed. If there were

**Fig. 1** Study flow diagram

**Fig. 2** pH, partial pressure of arterial carbon dioxide (\( P_{aCO2} \)) and plasma potassium concentrations \([K^+]_p\) changes during laparoscopic surgery. * indicates \( P < 0.05 \) with paired t-test.
extreme outliers, winsorization was applied. Only two values from each of baseline $[K^+]_p$ and pH at the end of surgery were extreme outliers and were replaced with the most adjacent values. If the missing value rate was below 5%, we planned a complete case analysis. Normality assumptions were examined with a visual check of the quantile–quantile (Q–Q) plot. Variables of apparent violated normality assumption were treated with non-parametric statistical analysis methods.

Baseline characteristics were presented using descriptive statistics. Values at baseline and at the end of surgery were compared using the paired t-test. To investigate the relationship between the magnitude of PaCO$_2$ changes, $[K^+]_p$ changes, and patient characteristics, correlation analysis was performed. For the relationship between the magnitude of PaCO$_2$ changes and $[K^+]_p$ changes, visual checking of the scatter plot and curve-fit analysis using regression were performed with various types of equations. Based on these results, linear regression was performed to clarify the relationship between the magnitude of PaCO$_2$ changes and $[K^+]_p$ changes, and other patient characteristics.

To establish the final model, the stepwise selection method was used, and assumptions of linear regression were tested with a histogram and a Q–Q plot for multicollinearity, Durbin–Watson statistics for autocorrelation, and visual checking of the scatter plot for each independent variable for homoscedasticity. Regression diagnostics were performed with normality test of residuals. Finally, the characteristics of patients who presented with hypercarbia were also compared to baseline values ($P < 0.001$, Cohen’s $d=1.26$). pH also decreased reciprocally to the increased PaCO$_2$ ($P < 0.001$, Cohen's $d=0.99$), with a similarly large effect size observed. $[K^+]_p$ increased by 0.25 (95% CI: 0.20 to 0.31) mmol/L at the end of surgery compared to baseline values ($P < 0.001$, Cohen's $d=0.03$). The effect size of this change was small. Before and after changes in these variables are presented graphically in Fig. 2. Bicarbonate and standard base excess values remained within normal laboratory limits at all time points (Table 2).

Correlation analysis between PaCO$_2$ changes, $[K^+]_p$ changes and other baseline patient characteristics are presented in Table 3. Increases in PaCO$_2$ correlated weakly with increasing BMI ($r=0.12, P=0.049$), and higher $[K^+]_p$ were associated with type II diabetes mellitus (rho = 0.13, $P=0.023$). According to curve-fit analysis results, a linear model ($R^2=0.217$, $P<0.001$) was suitable because quadratic and squared models resulted in the same proportion of the variance for the $[K^+]_p$ change associated with PaCO$_2$ changes ($R^2=0.217$, $P<0.001$ for the quadratic and squared models).

Multiple regression analysis evaluating whether PaCO$_2$ changes during surgery significantly predicted immediate changes in $[K^+]_p$ at completion of surgery demonstrated that a final fitted regression model could predict 33.1% of the observed variance in PaCO$_2$ ($R^2=0.331,$
Table 1 Preoperative and intraoperative variables

| Parameters                                                                 | Patients (n = 289) | Missing rate (%) |
|---------------------------------------------------------------------------|--------------------|------------------|
| Age (years)                                                               | 63.2 ± 11.5        | 0.00             |
| Male gender                                                               | 176 (60.9%)        | 0.00             |
| Body mass index (kg/m²)                                                   | 29.3 ± 6.8         | 0.35             |
| Corrected ARISCAT score                                                   | 29.5 ± 11.8        | 4.84             |
| Current smoker                                                            | 51 (17.6%)         | 0.00             |
| American Society of Anesthesiologists status                             |                    |                  |
| Class 1                                                                   | 21 (7.3%)          |                  |
| Class 2                                                                   | 124 (42.9%)        | 2.08             |
| Class 3                                                                   | 129 (44.6%)        |                  |
| Class 4                                                                   | 9 (3.1%)           |                  |
| No of patients with ≥ 1 chronic disease                                   | 214 (74.0%)        | 0.00             |
| Types of chronic disease*                                                 |                    |                  |
| Obstructive lung disease                                                  | 84 (29.1%)         |                  |
| Type II diabetes mellitus                                                 | 57 (19.7%)         |                  |
| Essential hypertension                                                    | 149 (51.6%)        |                  |
| Renal disease (creatinine > 120 mol/L)                                    | 25 (8.7%)          | 0.00             |
| Hepatic disease (bilirubin > 20 μmol/L)                                   | 25 (8.7%)          |                  |
| Coronary artery disease                                                   | 46 (15.9%)         |                  |
| Emergency surgery                                                         | 6 (2.1%)           | 0.00             |
| Types of laparoscopic procedures                                          |                    |                  |
| Anterior resection                                                       | 61 (21.1%)         |                  |
| Left or right hemicolectomy                                               | 60 (20.8%)         |                  |
| Small bowel resection                                                    | 2 (0.7%)           |                  |
| Intestinal bypass                                                        | 2 (0.7%)           | 0.00             |
| Adrenalectomy                                                             | 4 (1.4%)           |                  |
| Colostomy or adhesiolysis                                                | 6 (2.1%)           |                  |
| Cholecystectomy                                                           | 10 (3.5%)          |                  |
| Diaphragmatic hernia repair                                               | 1 (0.3%)           |                  |
| Fundoplication                                                           | 5 (1.7%)           |                  |
| Sleeve gastrectomy gastric bypass                                         | 17 (5.9%)          |                  |
| Splenectomy                                                               | 2 (0.7%)           |                  |
| Pancreatectomy                                                            | 1 (0.3%)           |                  |
| Hernia repair                                                             | 7 (2.4%)           |                  |
| Liver resection                                                           | 20 (6.9%)          |                  |
| Nephrectomy                                                              | 45 (15.6%)         |                  |
| Radical prostatectomy                                                     | 35 (12.1%)         |                  |
| Pyeloplasty                                                               | 1 (0.3%)           |                  |
| Rectopexy                                                                | 1 (0.3%)           |                  |
| Removal of pelvic or peritoneal mass                                      | 3 (1.0%)           |                  |
| Exploratory procedures                                                   | 6 (2.1%)           |                  |
| Duration of surgical procedure (min)                                      | 184.0 (137.3 to 240.0) | 0.35 |
| Intraoperative fluid administration (l)                                   | 2.0 (1.0 to 2.1)   | 1.73             |

Values are expressed as mean ± standard deviation or median (interquartile range) for continuous variables, number (percentile) for categorical variables. Missing data rate of each variable is presented.

* Some patients had > 1 chronic disease

F(3,259) = 38.915, P < 0.001). Regression analysis showed that PaCO₂ changes, intraoperative Plasma-Lyte 148 administration volume, and a history of type II diabetes mellitus significantly predicted changes in [K⁺]ₚ at completion of surgery (β = 0.018 (95% CI: 0.011 to 0.025), P < 0.001; β = 0.061 (95% CI: 0.031 to 0.091), P < 0.001; β = 0.154 (95% CI: 0.023 to 0.284), P = 0.021 respectively). There was no association between [K⁺]ₚ and duration of surgery (P = 0.102, Table 4).

Thus, for each 10 mmHg increment of PaCO₂ from baseline during surgery, [K⁺]ₚ increased by 0.18 mmol/L. Furthermore, [K⁺]ₚ increased by 0.061 mmol/L for each litre of Plasma-Lyte 148 administered, and patients with type II diabetes mellitus experienced a 0.154 mmol/L greater rise in [K⁺]ₚ compared to those without.

Discussion

In a single centre retrospective study describing the association of acute changes in PaCO₂ with acute changes in [K⁺]ₚ in patients undergoing laparoscopic surgery, CO₂ pneumoperitoneum induced small but significant increases in PaCO₂ and small but significant decreases in pH during surgery. Moreover, [K⁺]ₚ increased slightly but significantly at the end of surgery compared to baseline. After adjustment for multiple potential confounders, for every 10 mmHg increment in PaCO₂ from baseline, [K⁺]ₚ increased by almost 0.2 mmol/L. Finally, changes in [K⁺]ₚ were additionally positively affected by the amount of Plasma-Lyte 148 administered and the presence of type II diabetes mellitus. Given the magnitude of the changes observed, our findings imply that any acute hypercarbic or hyperkalaemic state induced by laparoscopic surgery is probably of limited clinical significance.

Our findings imply that during surgery clinicians should not ascribe large changes in [K⁺]ₚ to hypercarbia. Other aetiological factors should be considered as a more likely cause of hyperkalaemia. Such differentials should include renal failure, non-renal and endocrine causes, medications and surgical factors. The main hormonal system regulating renal potassium excretion is the renin–angiotensin–aldosterone system. Medications used in the perioperative setting that inhibit this system include angiotensin converting enzyme inhibitors, angiotensin II receptor blockers, nonsteroidal anti-inflammatory drugs, and adrenergic beta-antagonists. Other common anaesthesia medications known to cause hyperkalaemia during surgery include suxamethonium, beta-blockers, digoxin, mannitol and some intravenous penicillins (high potassium content). Finally, surgical causes that can result in the release of potassium from injured cells include ischaemia–reperfusion injury, rhabdomyolysis from surgical associated muscle damage, and high-volume blood
In many patients, during surgery the cause of hyperkalaemia is multifactorial. Our findings however are at variance with the case series of 24 patients undergoing cardiac surgery who were exposed to a 15 min period of apnoeic or low tidal volume ventilation [9]. Over this time period, PaCO₂ increased from 43.6 mmHg at baseline to 83.9 mmHg with [K⁺]ᵣ increasing marginally from 4.16 mmol/L at baseline to 4.28 mmol/L at 15 min. This study was limited by a small sample size and exposure to significant hypercarbia for only a 15 min duration. Similarly, Nataliani et al. [10] compared the acute changes in [K⁺]ᵣ in acutely hypercapnic patients undergoing rigid bronchoscopy. The sampling time to evaluate the effects of PaCO₂ on hyperkalaemia was also of short duration (20 min), and acute respiratory acidosis did not affect [K⁺]ᵣ. These findings suggest that there appears to be both a quantitative and temporal relationship between changes in [K⁺]ᵣ and hypercarbia under anaesthesia, and that acute hypercarbia of short duration may have less of an effect on [K⁺]ᵣ compared to prolonged exposure. As there was no significant association between [K⁺]ᵣ and duration of surgery in this study, and only patients who had a surgical duration of two hours were included, it is likely that the temporal onset of [K⁺]ᵣ change due to a change in PaCO₂ is less than two hours. Our study has several strengths. Data on a relatively large sample size was provided and a detailed description of the acute effects of hypercarbia and its associations with [K⁺]ᵣ during laparoscopic surgery were

Table 2 Comparisons of pH, partial pressure of arterial carbon dioxide (PaCO₂) and plasma potassium concentrations ([K⁺]ᵣ between baseline and end of surgery

| Variable | Baseline | End of surgery | Paired differences | P value | Cohen’s d |
|----------|----------|----------------|-------------------|---------|-----------|
| pH       | 7.40 ± 0.053 | 7.34 ± 0.069 | −0.061 (−0.068 to −0.054) | <0.001* | 0.99 |
| PaCO₂ (mmHg) | 41.44 ± 5.84 | 46.61 ± 8.83 | 5.18 (4.27 to 6.09) | <0.001* | 1.26 |
| [K⁺]ᵣ (mmol/L) | 3.91 ± 0.416 | 4.16 ± 0.591 | 0.251 (0.197 to 0.305) | <0.001* | 0.03 |
| Bicarbonate (mmol/L) | 24.96 ± 3.58 | 23.90 ± 3.61 | 1.06 (0.54 to 1.58) | <0.001* | 0.19 |
| Standard base excess (mmol/L) | 0.87 ± 2.97 | -0.93 ± 2.74 | 1.80 (1.44 to 2.16) | <0.001* | 0.24 |

Paired differences were presented with 95% confidence intervals. The effect size is presented with Cohen’s d.

* Indicates P < 0.05 with paired t-test

Table 3 Correlation analysis between partial pressure of arterial carbon dioxide (PaCO₂) changes, plasma potassium concentrations ([K⁺]ᵣ changes and other parameters

| Variables | PaCO₂ changes | [K⁺]ᵣ changes |
|-----------|---------------|---------------|
|           | Coefficient   | P value       | Coefficient | P value       |
| Age       | −0.03         | 0.642         | −0.01       | 0.807         |
| ASA class | −0.06         | 0.315         | 0.04        | 0.546         |
| BMI       | 0.12*         | 0.049         | 0.03        | 0.632         |
| Current smoking status | 0.06 | 0.347 | 0.03 | 0.571 |
| Lung disease | 0.06 | 0.348 | 0.09 | 0.132 |
| Coronary disease | −0.04 | 0.553 | −0.01 | 0.829 |
| Type II diabetes mellitus | −0.04 | 0.534 | 0.13* | 0.023 |
| Essential hypertension | −0.05 | 0.394 | 0.02 | 0.679 |
| Renal disease | 0.03 | 0.576 | −0.07 | 0.215 |
| Hepatic disease | −0.03 | 0.568 | 0.09 | 0.128 |
| Elective / Emergency operation | 0.08 | 0.161 | −0.02 | 0.735 |
| Duration of surgery | 0.02 | 0.783 | 0.10 | 0.102 |
| Corrected ARISCAT score | 0.06 | 0.356 | 0.06 | 0.337 |
| Intraoperative fluid amount | 0.01 | 0.928 | 0.10 | 0.113 |

Coefficients are presented as Pearson correlation for continuous variables or Spearman’s rho for categorical variables.

* Indicates P < 0.05
analysed. \( \text{PaCO}_2 \), \([K^+]_p\), bicarbonate and standard base excess values were collected directly from arterial blood samples and were not amenable to ascertainment bias or derivation. The timing of blood gases in relation to the start and completion of surgery was manually cross-checked with the patients’ medical records by two independent investigators, allowing for accurate confirmation of these data points at the start and at completion of surgery. The large dataset examines the impact of laparoscopic surgery on \( \text{PaCO}_2 \) and \([K^+]_p\) by directly measuring \( \text{PaCO}_2 \) and \([K^+]_p\) at baseline and at the completion of surgery, allowing a greater understanding of the temporal relationship of changes in \([K^+]_p\) and \( \text{PaCO}_2 \) to be observed.

Our study has several limitations. It was a single centre study, limiting the external validity of our findings. However, given laparoscopic surgery with \( \text{CO}_2 \) insufflation is utilised worldwide, our findings are likely relevant and externally generalisable. We cannot extrapolate our results to shorter surgical times, to paediatric patients or to patients at high risk of developing hyperkalaemia (renal failure patients, rhabdomyolysis, tumour lysis syndrome etc.). However, the biological principles and physiochemical changes described are likely to apply to all hypercarbic patients undergoing general anaesthesia. As only two blood gases were sampled, the exact onset and rate of change of the findings observed cannot be evaluated. This study was not designed to determine whether there was an association between changes in \( \text{PaCO}_2 \) or \([K^+]_p\) and the occurrence of detrimental clinical outcomes. Similarly, we did not evaluate the association between changes in \( \text{PaCO}_2 \) or \([K^+]_p\) with specific medications that are known to be associated with hyperkalaemia such as preoperative use of certain antihypertensive medications e.g. angiotensin-converting enzyme inhibitors, angiotensin II receptor blockers, and/or potassium-sparing diuretics. This data was unable to be collected and we acknowledge this as a limitation of the study. However, this study was designed to evaluate the change in \( \text{PaCO}_2 \) from baseline rather than the absolute value of \([K^+]_p\) at baseline, which is more likely to be effected by these medications. As the anaesthesia records in our institution are hand-written, and not part of the hospital’s electronic medical record system, corresponding end-tidal \( \text{CO}_2 \) and acid–base responses to an increased \( \text{PaCO}_2 \).

For all cases the initial \( \text{CO}_2 \) insufflation occurred at a rate of 1 to 5 L/min to achieve an intraabdominal pressure of 12 to 15 mmHg. We used a constant \( \text{CO}_2 \) flow of approximately 200 to 400 mL/min to maintain pneumoperitoneum perioperatively. Given this standard practice for all patients, we cannot make any inferences about the relationship of intraabdominal pressure and changes in \( \text{PaCO}_2 \). Finally, due to the heterogeneity of clinical precipitants for potassium disorders, and the observational nature of this study, our findings are merely associations and do not infer causation.

| Table 4 Coefficients estimated using linear regression for predicting plasma potassium concentrations \([K^+]_p\) changes |
| Variables | Coefficient | P value |
| --- | --- | --- |
| Included in the final regression model | | |
| \( \text{PaCO}_2 \) changes | 0.018 (0.011–0.025) | <0.001 |
| Intraoperative Plasma-Lyte 148 administration volume | 0.061 (0.031–0.091) | <0.001 |
| Type II diabetes mellitus | 0.154 (0.023–0.284) | 0.021 |
| Excluded in the final regression model | | |
| Age | 0.084 | 0.464 |
| American Society of Anesthesiologists status | 0.029 | 0.683 |
| Body mass index | 0.046 | 0.703 |
| Smoking history | −0.008 | 0.889 |
| Obstructive lung disease | 0.058 | 0.348 |
| Coronary artery disease | −0.041 | 0.464 |
| Hypertension | 0.014 | 0.847 |
| Renal disease (creatinine > 120 \( \text{umol/L} \)) | −0.102 | 0.058 |
| Hepatic disease (bilirubin > 50 \( \text{umol/L} \)) | 0.07 | 0.199 |
| Emergency surgery | −0.036 | 0.497 |
| Duration of surgical procedure | 0.156 | 0.235 |
| Corrected ARISCAT score | 0.082 | 0.475 |

Coefficient values are expressed with 95% confidential intervals when corresponding variable is included in the final regression model.
Conclusion
During laparoscopic surgery, for every 10 mmHg increase in PaCO2 from baseline, [K+]p increased by almost 0.2 mmol/L. [K+]p was further elevated with the administration of the crystalloid solution Plasma-Lyte 148 and the presence of type II diabetics mellitus. Clinicians should not ascribe larger changes in [K+]p to hypercarbia in this setting and should consider other aetiological factors in their differential diagnosis.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s12893-020-01034-w

Additional file 1: Table S1. Database_Final_CO2_Study_OctR4_2020.

Abbreviations
PaCO2: Partial pressure of arterial carbon dioxide; CO2: Carbon dioxide; CI: Confidence interval; [K+]p: Plasma potassium concentrations; QQ: Quantile–quantile.

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Authors’ contributions
CRediT (Contributor Roles Taxonomy). LW, DK, RB: Conceptualization, Methodology, Writing–Reviewing and Editing. DKL, DL: Formal analysis, Writing–Reviewing and Editing. CG, AH, LF, DB, ST, DJ: Data curation, Writing–Original Draft Preparation. All authors read and approved the final manuscript.

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Availability of data and materials
The deidentified datasets analysed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
The study was approved by the Austin Health Human Research Ethics Committee (LNR/19/Austin/41) and no additional administrative permissions were required to access the raw data. Given the retrospective design, a waiver of consent for publication is Not applicable.

Competing interests
The authors declare that they have no competing interests.

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