Impact of Comorbidities on Beneficial Effect of Lactated Ringers Versus Saline in Sepsis Patients

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Research

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Abstract

**Background:** Lactated Ringers reduced mortality more than saline in sepsis patients but increased mortality more than saline in traumatic brain injury patients.

**Method:** This prospective cohort study included sepsis patients and identified heart, lung, liver, kidney, and endocrine comorbidities by detailed history taking and routine admission survey such as HbA1C, and liver functions. We evaluate resuscitation response with central venous pressure, central venous oxygen saturation, and serum lactate level simultaneously. Propensity-score matching and Cox regression were used to estimate 60-day mortality. The competing risk model compared the lengths of hospital and ICU stays with the subdistribution hazard ratio (SHR). Mixed-effect linear models were used to fit clinical variables and electrolyte trends.

**Results:** Overall, 874 patients were included in the analysis; 636 patients were in the saline group, and 302 patients were in the lactated Ringers group. The lactated Ringers group had a lower mortality rate (adjusted hazard ratio, 0.59; 95% CI 0.43-0.81) and shorter lengths of hospital (SHR, 1.39; 95% C.I. 1.15-1.67) and ICU stays (SHR, 1.41; 95% CI 1.17, 1.71) than the saline group; the differences were greater in patients with chronic pulmonary disease and small and nonsignificant in those with chronic kidney disease, moderate to severe liver disease and cerebral vascular disease. The resuscitation efficacy was the same between fluid types, but serum lactate levels were significantly higher in the lactated Ringers group than in the saline group (0.12 mg/dL/hour; 95% C.I.: 0.03, 0.21), especially in chronic liver disease patients receiving lactated Ringers. The serum potassium level increased within the first few hours and recovered more slowly in patients with chronic kidney disease regardless of fluid type. Compared to the saline group, the lactated Ringers group achieved target glucose level earlier in both diabetes and non-diabetes patients.

**Conclusion:** Patients receiving lactated Ringers had lower mortality and shorter lengths of hospital and ICU stays than those receiving saline, especially patients with chronic pulmonary disease, but there were no differences in those with chronic kidney disease, chronic liver disease and cerebral vascular disease. Comorbidities are important for clinicians to consider before choosing a fluid type.

Introduction

The fourth edition of the Surviving Sepsis Campaign suggests using either balanced crystalloids or saline for fluid resuscitation in sepsis patients\(^1\). Recently, a large, pragmatic randomized trial found that critically ill adult patients receiving balanced crystalloids had lower rates of the composite outcome of death and renal adverse events than those receiving saline.\(^2\) Our latest network meta-analysis also found a lower risk of mortality in sepsis patients treated with balanced crystalloids than in those treated with saline\(^3\). In contrast, balanced crystalloids increased mortality in traumatic brain injury patients\(^3\). Therefore, both the fluid types and patient condition need to be considered when choosing an optimal fluid treatment.
Saline and balanced crystalloids, such as lactated Ringers, have different fluid osmolalities (saline: isotonic, 308 mOsm/kg; lactated Ringers: hypotonic, 273 mOsm/kg) and contain different electrolyte composites and lactate metabolites, resulting in varying risks for ineffective increases in central venous pressure, hyperkalemia, acidosis or glucose instability, especially in sepsis patients with comorbidities\(^4\). Saline solution contains no potassium or other chemical compounds, while lactated Ringers solution contains 4 mmol/L potassium and 28 mmol/L sodium lactate. The excessive potassium in lactated Ringers may specifically lead to hyperkalemia in chronic kidney disease patients\(^5\). Seventy percent of serum lactate undergoes gluconeogenesis, which can potentially lead to an increase in blood glucose levels and instability, and these effects might be more significant in diabetes patients\(^6\),\(^7\). Thirty percent of serum lactate undergoes oxidation in the liver, producing \(\text{HCO}_3^-\), which helps limit acidosis during sepsis. However, lactate metabolism is impaired in chronic liver disease patients, and thus, the acidosis prevention effect of lactated Ringers in chronic liver disease patients is questionable\(^8\),\(^9\).

This study aimed to investigate differences in mortality and lengths of stay in the intensive care unit (ICU) and hospital in sepsis patients treated with lactated Ringers or saline. Subgroup analyses of sepsis patients with different comorbidities, including chronic pulmonary disease, chronic kidney disease, chronic liver disease and diabetes, were also performed. The differences in trends of central venous pressure, central venous oxygen saturation, and the serum lactate level between fluid types were also investigated to compare their resuscitation efficacy. During and after the resuscitation periods, we compared the trends of serum potassium in patients with and without chronic kidney disease, the trends of blood glucose levels in diabetes and nondiabetes patients, and the trends of serum lactate in patients with and without chronic liver disease.

**Methods**

**Study population and care protocols**

Since 2011, every critically ill patient admitted to the ICU in the Veteran General Hospital Taichung has been screened for sepsis according to the sepsis criteria of the 2001 international sepsis conference\(^10\). For sepsis patients who met the criteria, we implemented bundled care and included them in this study. The sepsis bundles include (1) guided fluid resuscitation considering lactate clearance, mixed venous oxygen saturation, and urine volume; (2) early empiric antibiotic prescription rules, (3) glucose control protocols, (4) protective ventilator settings, (5) c-reactive protein, procalcitonin, B-type natriuretic peptide, HbA1C, and albumin levels before fluid resuscitation, and (6) lactate and mixed venous oxygenation levels, which are checked every 6 hours for every sepsis patient in the first 24 hours. Among the bundles, the glucose target was 140 ~ 180 mg/dl. An insulin sliding scale was applied for glucose levels higher than 180 mg/dl. The frequency of the one-touch glucose examination was every 4 hours if the blood glucose level was lower than 180 mg/dl or every 2 hours if the blood glucose level was higher than 180 mg/dl.
In this prospective cohort study, we retrieved relevant patient data from electronic medical records; the protocol was approved by the institutional review board of the Veteran General Hospital Taichung (IRB no. CF16017A). We collected basic patient characteristics, biochemical blood test results, fluid volumes and relevant medications prescribed for every patient from 2011–2017. According to their resuscitation fluid profiles in the first 24 hours, patients were divided into two groups: the saline group, who received mainly saline for resuscitation and less than 500 ml of lactated Ringers, and the lactated Ringers group, who received more than 500 ml of lactated Ringers in the first 24 hours. In the sensitivity analysis, we further separated patients into three groups: the saline only, saline predominant (more saline in the total fluid amount) and lactated Ringers predominant groups (more lactated Ringers in the total fluid amount). Definitions of comorbidities, including chronic kidney disease, diabetes, mild or moderate to severe liver disease, congestive heart failure, and cerebral vascular disease, followed those of the Charlson Comorbidity Index \(^{11}\) (Supplementary Table 1). We also recorded blood transfusion volumes, including red blood cells and fresh frozen plasma, and hemodialysis events for every patient.

**Statistical methods for mortality and lengths of stay in the ICU and hospital**

The differences in patient characteristics between two fluid type groups were compared by using Pearson's chi-squared test for categorical variables and Student's t-test for continuous variables. As our cohort was not randomly assigned to the two fluid type groups, we used propensity-score matching by fluid type to control for potential confounding factors and selection bias \(^{12}\), thereby optimizing comparability between the saline group and lactated Ringers group \(^{13}\). We entered the APACHE score, age and HbA1c level into the logistic regression analysis to compute propensity scores for each participant. The propensity score represented the probability of a patient with sepsis being assigned to the lactated Ringers group. Based on the propensity score, patients who received lactated Ringers were matched with two patients who received saline, and then we created propensity score-matched sets for the Cox proportional hazard model to estimate the hazard ratio of 60-day mortality. In addition, we also performed a logistic regression analysis to compare 60-day mortality between the saline group and lactated Ringers group \(^{14}\). Finally, we used the competing risk model to estimate the subdistribution hazard ratios (SHRs) for the lengths of ICU and hospital stays \(^{15, 16}\). We considered death a competing risk, and discharge from the ICU or hospital was the event of interest. We hypothesized that resuscitation fluid with fewer complications would lead to a shorter length of ICU stay and earlier discharge. We also compared the lengths of ICU stay and hospital stay between patients with and without comorbidities.

**Statistical method for repeated measured data**

Mixed-effect linear models were used to analyze changes in the following clinical variables during and after the resuscitation period: central venous pressure, central venous oxygen saturation, serum bicarbonate, serum lactate, creatinine, urine output, serum potassium and blood glucose. Interaction terms between fluid types and changes in those clinical variables were included to investigate whether patients receiving these two fluids showed different trends. Regarding the confounding effect of kidney
function on serum potassium levels, we performed separate regression analyses in patients with and without chronic kidney disease. In addition, we also performed analyses to investigate the differences in blood glucose trends in patients with and without diabetes to avoid confounding effects between blood glucose and diabetes status. All statistical tests were performed with Stata version 14.0 (StataCorp, Texas, USA).

**Statistical method for glycemic variability**

We used the mean amplitude of glycemic excursions (MAGE)\(^{17}\) and coefficient of variation (CoV)\(^{18}\) to assess glycemic variability. The MAGE represents the mean blood glucose value exceeding the standard deviation from the 24-hour mean blood glucose level, and CoV represents the ratio of the standard deviation to the mean glucose level. A high blood glucose index and low blood glucose index represent the risks of hyperglycemia and hypoglycemia, respectively\(^{19}\). The glucose index was derived from a logarithmic transformation of the blood glucose scale that assigned maximum risk to blood glucose of 20 and 600 ml/dl and zero risk to 112.5 mg/dl\(^{20}\). The above glycemic parameters were calculated using EasyGV software\(^{21}\).

**Results**

**Baseline patient characteristics**

A total of 874 sepsis patients were admitted to the Taichung Veteran General Hospital ICUICU from 2011 to 2017. The saline group and lactated Ringers group included 636 and 302 patients, with male percentages of 76.1% and 73.8%, mean ages of 71.9 and 70.7, and mean Acute Physiology and Chronic Health Evaluation (APACHE) scores of 26.0 ± 6.9 and 29.0 ± 6.4, respectively (Table 1). The total fluid resuscitation volume on day 1 was not significantly different between the saline and lactated Ringers groups (4591 ml vs. 4959 ml, p = 0.157). The percentages of chronic kidney disease (saline group, 4.1%; lactated Ringers group 10.9%, p = 0.057), chronic liver disease (saline group, 4.6%; lactated Ringers group 5.0%, p = 0.869), type 2 diabetes mellitus (saline group, 49.7%; lactated Ringers group 44.7%, p = 0.162), and other comorbidities were not significantly different between the two groups. The results of laboratory tests, including HbA1c, procalcitonin, c-reactive protein, B-type natriuretic peptide and albumin, before fluid resuscitation were not significantly different between the two groups (Table 1).
Table 1
Patient’s characteristics according to types of fluid resuscitation

|                           | Saline group | Lactated Ringer's group | P value |
|---------------------------|--------------|-------------------------|---------|
| No. of case               | 636          | 302                     |         |
| Gender, n (%)             |              |                         |         |
| Male                      | 484 (76.1)   | 223 (73.8)              | 0.466   |
| Female                    | 152 (23.8)   | 79 (26.2)               |         |
| Age, mean (SD)            | 71.9 (15.8)  | 70.7 (15.3)             | 0.277   |
| Body mass index, mean (SD)| 23.2 (4.4)   | 23.0 (4.6)              | 0.613   |
| APACHE score, mean (SD)   | 26.0 (6.9)   | 29.0 (6.4)              | < 0.001 |
| Total fluid resuscitation in day 1, mean (SD) | 4591 (3778) | 4959 (3602) | 0.157   |
| NS fluid amount in day 1, mean (SD) | 4587 (3776) | 1787 (2362) | < 0.001 |
| LR fluid amount in day 1, mean (SD) | 3.9 (44.2) | 3172 (2442) | < 0.001 |
| Comorbidity               |              |                         |         |
| Coronary artery disease, n (%) | 68 (10.7)    | 26 (8.6)                | 0.353   |
| Congestive heart failure, n (%) | 156 (24.5)   | 68 (22.5)               | 0.513   |
| Cerebral vascular disease, n (%) | 214 (33.6)   | 83 (31.0)               | 0.413   |
| Chronic pulmonary disease, n (%) | 318 (50)     | 141 (46.7)              | 0.364   |
| Chronic kidney disease, n (%) | 45 (7.1)     | 33 (10.9)               | 0.057   |
| Chronic liver disease, n (%) | 29 (4.6)     | 15 (5.0)                | 0.869   |
| Diabetes, n (%)           | 316 (49.7)   | 135 (44.7)              | 0.162   |
| Laboratory exam before resuscitation |          |                         |         |
| HbA1C (%), mean (SD)      | 6.3 (1.4)    | 6.5 (1.5)               | 0.186   |
| Procalcitonin (ng/mL), mean (SD) | 18.7 (31.0)  | 18.8 (31.3)            | 0.979   |
| c-reactive protein (mg/dL), mean (SD) | 15.7 (11.7)  | 14.5 (11.3)            | 0.162   |
| B-type natriuretic peptide (pg/mL), mean (SD) | 8911 (10659) | 8385 (10859) | 0.562   |
| Albumin (g/dL), mean (SD) | 2.6 (0.7)    | 2.7 (0.6)               | 0.013   |

Mortality and lengths of ICU and hospital stays
The propensity score distributions between the saline group and lactated Ringers group were similar after matching (supplementary table 2). In the propensity-score matched Cox regression analysis, the lactated Ringers group had a lower risk of mortality than the saline group (adjusted hazard ratio, 0.59; 95% CI 0.43–0.81). In the logistic regression analysis for 60-day mortality, the lactated Ringers group also had lower odds of mortality than the saline group (adjusted odds ratio, 0.68; 95% CI 0.51–0.92). In the competing risk regression analysis for length of ICU stay, patients resuscitated with lactated Ringers were discharged from the ICU (SHR, 1.41; 95% CI 1.17–1.71) and hospital (SHR, 1.39; 95% CI 1.15–1.67) earlier than those resuscitated with saline. We divided patients into three subgroups according to the saline fluid amount they received. The results showed that both the lactated Ringers predominant group (SHR, 1.49; 95% CI 1.18–1.88) and saline predominant group (SHR, 1.32; 95% CI 1.02–1.71) had significantly shorter ICU stays than the saline group (Table 2). Regarding the subgroup analyses stratified by comorbidities (Table 3), the use of lactated Ringers, compared to the use of saline, significantly reduced the length of stay in the ICU in patients with chronic pulmonary disease (SHR, 1.80; 95% CI 1.37, 2.36), without chronic kidney disease (SHR, 1.48; 95% CI 1.21, 1.80), without moderate to severe liver disease (SHR, 1.44; 95% CI 1.19, 1.75), and without cerebral vascular disease (SHR, 1.54; 95% CI 1.22, 1.95). No significant differences were found between the two types of fluids in patients without chronic pulmonary disease (SHR, 1.17; 95% CI 0.89, 1.53), with chronic kidney disease (SHR, 0.77; 95% CI 0.35, 1.70), with moderate to severe liver disease (SHR, 1.01; 95% CI 0.32, 3.17), and with cerebral vascular disease (SHR, 1.27; 95% CI 0.92, 1.76).
Table 2
Mortality analysis and competing risk analysis for length of stay in intensive care unit and hospital

| Outcome                        | Comparison                                      | Adjusted hazard ratio | P value   |
|--------------------------------|-------------------------------------------------|-----------------------|-----------|
| 60 days mortality              | Lactated Ringer’s group vs saline group          | 0.59 (0.43–0.81)      | < 0.001   |

B. Logistic regression analysis for 60-days mortality

| Outcome                        | Comparison                                      | Adjusted odd ratio    | P value   |
|--------------------------------|-------------------------------------------------|-----------------------|-----------|
| 60 days mortality              | Lactated Ringer’s group vs saline group          | 0.68 (0.51, 0.92)     | 0.013     |
|                                | APACHE                                          | 1.09 (1.07, 1.12)     | < 0.001   |

C. Logistic regression analysis for 60-days mortality with three saline exposure level

| Outcome                        | Comparison                                      | Adjusted odd ratio    | P value   |
|--------------------------------|-------------------------------------------------|-----------------------|-----------|
| 60 days mortality              | Saline predominant vs saline only group          | 0.74 (0.50, 1.11)     | 0.142     |
|                                | Lactated Ringer’s predominant vs saline only group | 0.64 (0.44, 0.93)     | 0.019     |
|                                | APACHE                                          | 1.09 (1.07, 1.12)     | < 0.001   |

D. Competing risk analysis for ICU and hospital stay

| Outcome                        | Comparison                                      | SHR (95% C.I.)        | P value   |
|--------------------------------|-------------------------------------------------|-----------------------|-----------|
| Length of ICU day              | Lactated Ringer’s group vs saline group          | 1.41 (1.17, 1.71)     | < 0.001   |
|                                | APACHE                                          | 0.95 (0.93, 0.96)     | < 0.001   |
| Length of hospital day         | Lactated Ringer’s group vs saline group          | 1.39 (1.15, 1.67)     | < 0.001   |
|                                | APACHE                                          | 0.95 (0.93, 0.96)     | < 0.001   |

E. Competing risk analysis for ICU and hospital stay with three saline exposure level

*Subdistribution hazard ratio
## A. Propensity score matching cox regression analysis for 60-days mortality

|                          |                          | Hazard Ratio | p-value |
|--------------------------|--------------------------|--------------|---------|
| **Length of ICU day**    | Lactated Ringer’s predominant vs saline only group | 1.49 (1.18, 1.88) | 0.001   |
|                          | Saline predominant vs saline only group            | 1.32 (1.02, 1.71) | 0.034   |
|                          | APACHE                                                 | 0.95 (0.93, 0.96) | < 0.001 |
| **Length of hospital day** | Lactated Ringer's predominant vs saline only group | 1.44 (1.14, 1.81) | 0.002   |
|                          | Saline predominant vs saline only group            | 1.32 (1.01, 1.71) | 0.037   |
|                          | APACHE                                                 | 0.95 (0.93, 0.96) | < 0.001 |

*Subdistribution hazard ratio*
Table 3
Competing risk analysis for length of stay in intensive care unit and hospital in different comorbidities

| Comorbidities                  | No. of cases | Length of ICU day | Length of hospital day |
|-------------------------------|--------------|-------------------|------------------------|
|                               |              | SHR* (95% C.I.)   | P value                | SHR* (95% C.I.)   | P value                |
| Chronic pulmonary disease (CPD) |              |                   |                        |                        |                        |
| No CPD                        | 467          | 1.17 (0.89, 1.53) | 0.258                  | 1.13 (0.87, 1.48)   | 0.356                  |
| With CPD                      | 447          | 1.80 (1.37, 2.36) | < 0.001                | 1.78 (1.36, 2.34)   | < 0.001                |
| Chronic kidney disease (CKD)  |              |                   |                        |                        |                        |
| No CKD                        | 838          | 1.48 (1.21, 1.80) | < 0.001                | 1.44 (1.18, 1.75)   | < 0.001                |
| With CKD                      | 76           | 0.77 (0.35, 1.70) | 0.525                  | 0.82 (0.36, 1.85)   | 0.635                  |
| Acute kidney injury or CKD    |              |                   |                        |                        |                        |
| No AKI or CKD                 | 373          | 1.35 (1.04, 1.76) | 0.026                  | 1.34 (1.02, 1.75)   | 0.032                  |
| With AKI or CKD               | 528          | 1.28 (0.96, 1.70) | 0.094                  | 1.25 (0.94, 1.66)   | 0.125                  |
| Mild liver disease (LD)       |              |                   |                        |                        |                        |
| No liver disease              | 676          | 1.38 (1.11, 1.72) | 0.004                  | 1.36 (1.09, 1.69)   | 0.006                  |
| With mild liver disease       | 238          | 1.49 (1.02, 2.18) | 0.039                  | 1.42 (0.97, 2.06)   | 0.068                  |
| Moderate to severe LD         |              |                   |                        |                        |                        |
| No liver disease              | 873          | 1.44 (1.19, 1.75) | < 0.001                | 1.41 (1.16, 1.71)   | < 0.001                |
| With moderate to severe LD    | 41           | 1.01 (0.32, 3.17) | 0.984                  | 1.01 (0.33, 3.40)   | 0.924                  |
| Cerebral vascular disease (CVD)|              |                   |                        |                        |                        |
| No CVD                        | 615          | 1.54 (1.22, 1.95) | < 0.001                | 1.52 (1.20, 1.93)   | 0.001                  |

*SHR, Subdistribution hazard ratio; lactated Ringer’s group compared to saline group (reference group).
| Comorbidities                        | No. of cases | Length of ICU day | Length of hospital day |
|-------------------------------------|--------------|-------------------|------------------------|
|                                     |              | SHR* (95% C.I.)   | P value                | SHR* (95% C.I.)   | P value                |
| With CVD                            | 299          | 1.27 (0.92, 1.76) | 0.148                  | 1.24 (0.90, 1.71) | 0.183                  |
| Congestive heart failure (CHF)      |              |                   |                        |                        |
| No CHF                              | 696          | 1.39 (1.11, 1.72) | 0.003                  | 1.36 (1.09, 1.68)  | 0.006                  |
| With CHF                            | 218          | 1.54 (1.05, 2.27) | 0.028                  | 1.55 (1.05, 2.28)  | 0.026                  |
| Diabetes mellitus                   |              |                   |                        |                        |
| No diabetes mellitus                | 472          | 1.41 (1.09, 1.81) | 0.008                  | 1.43 (1.10, 1.84)  | 0.007                  |
| With diabetes mellitus              | 442          | 1.43 (1.07, 1.91) | 0.015                  | 1.34 (1.01, 1.77)  | 0.041                  |
| Rheumatology disease (RD)           |              |                   |                        |                        |
| No RD                               | 838          | 1.32 (1.08, 1.61) | 0.006                  | 1.30 (1.07, 1.59)  | 0.009                  |
| With RD                             | 76           | 2.71 (1.34, 5.51) | 0.006                  | 2.35 (1.26, 4.38)  | 0.007                  |
| Malignancy                          |              |                   |                        |                        |
| No Malignancy                       | 505          | 1.47 (1.16, 1.87) | 0.002                  | 1.46 (1.14, 1.85)  | 0.002                  |
| With Malignancy                     | 409          | 1.33 (0.98, 1.80) | 0.071                  | 1.27 (0.94, 1.71)  | 0.122                  |

*SHR, Subdistribution hazard ratio; lactated Ringer’s group compared to saline group (reference group).

### Relationships between fluid types and clinical variables associated with resuscitation

Central venous pressure trends consistently increased in the first 24 hours (0.19 cmH₂O/hour; 95% C.I.: 0.15, 0.55) in both the saline group and lactated Ringers group, and the trends were not significantly different between the two groups (0.04 cmH₂O/hour; 95% C.I.: -0.02, 0.11) (Fig. 1A). Central venous oxygen saturation trends were also not significantly different between the two groups (0.52%/hr; 95% C.I.: -1.2, 1.19) (Fig. 1B). Serum bicarbonate progressively decreased in the saline group (-0.58 mmol/L/hour; 95% C.I.: -0.73, -0.42) but increased significantly in the lactated Ringers group (0.85 mmol/L/hour; 95% C.I.: 0.64, 1.06) (Fig. 1C). The serum lactate levels decreased in the first few hours in both groups; it
continued to decrease for 72 hours in the saline group (-0.35 mg/dL/hour; 95% C.I.: -0.41, -0.30) but remained abnormally high in the lactated Ringers group from 12 to 72 hours (0.12 mg/dL/hour; 95% C.I.: 0.03, 0.21) (Fig. 1D). For the first 7 days, serum creatinine and urine output trends were not significantly different between the saline group and lactated Ringers group (Fig. 1E, Fig. 1F).

Relationships between fluid types and serum potassium and blood glucose

Serum potassium levels were higher than normal on admission to the ICU and gradually decreased in the first 72 hours in the saline subgroup without chronic kidney disease (-0.02 mmol/L/hour; 95% CI: -0.02, -0.01) but remained in the normal upper limit in the saline subgroup with chronic kidney disease (-0.01 mmol/L/hour; 95% CI: -0.02, 0.01). The serum potassium trends between the saline group and lactated Ringers group were not significantly different in patients with or without chronic kidney disease (Fig. 2A). Blood glucose in the first 24 hours consistently remained higher than normal in the saline group in both diabetes (0.15 mg/dl/hour; 95% CI: -0.28, 0.31) and nondiabetes patients (0.28; 95% CI: 0.06, 0.51); in the lactated Ringers group, the blood glucose level approached the target level in both patients with diabetes (-1.25 mg/dl/hour; 95% CI: -1.78, -0.71) and patients without diabetes (-0.83 mg/dl/hour; 95% CI: -1.21, -0.44) (Fig. 2B). In patients receiving lactated Ringers, the serum lactate level was significantly higher in patients with moderate to severe liver disease than in patients without liver disease (p = 0.045) (Fig. 2C).

Relationships between fluid types and glycemic variability (MAGE, CoV)

Compared to the saline group, the lactated Ringers group had a significantly lower glucose CoV (-1.78%, 95% CI: -3.34, -0.23) and a lower MAGE (-0.51%, 95% CI: -0.89, -0.13). The high blood glucose index was significantly lower in the lactated Ringers group than in the saline group (-3.17, 95% CI: -4.40, -1.95), but the low blood glucose index was not significantly different (0.03, 95% CI: -0.66, 0.71) (Table 4).

Table 4

| Diabetic mellitus | APACHE | Lactated Ringer’s group vs Saline only |
|-------------------|--------|--------------------------------------|
| CoV (%)           | 1.64 (0.21, 3.08)* | 0.14 (0.03, 0.24)** | -1.78 (-3.34, -0.23)* |
| MAGE              | 0.41 (0.06, 0.75)* | 0.02 (-0.01, 0.05)  | -0.51 (-0.89, -0.13)** |
| HBGi              | 5.47 (4.35, 6.60)** | 0.05 (-0.04, 0.13)  | -3.17 (-4.40, -1.95)** |
| LBGI              | -0.22 (-0.85, 0.41) | 0.09 (0.04, 0.13)** | 0.03 (-0.66, 0.71)    |

*<0.05, **<0.01, ***<0.001.

Abbreviations: CoV, coefficient of variation; MAGE, mean amplitude of glycemic excursion; HBGi, High Blood Glucose Index; LBGI, Low Blood Glucose Index.
The proportions of patients requiring hemodialysis in the first 7 days were 27.4% in the saline group and 28.5% in the lactated Ringers group. The red blood cell transfusion volumes in the first 7 days were 443.4 ml (95% CI: 405.79, 480.99) in the saline group and 427.98 ml (95% CI: 370.75, 485.21) in the lactated Ringers group. The fresh frozen plasma transfusion volume in the first 7 days was significantly higher in the saline group (0.55 units, 95% CI: 0.45, 0.64) than in the lactated Ringers group (0.36 units, 95% CI: 0.22, 0.50) (Supplementary Table 3).

Discussion

This prospective cohort study found that using lactated Ringers solution for resuscitation in sepsis patients decreased mortality and shortened the lengths of ICU and hospital stays compared with using saline, especially in patients with chronic pulmonary disease, without chronic kidney disease, without moderate to severe liver disease and without cerebral vascular disease. The trends for central venous pressure and oxygen saturation during the resuscitation period were similar between the saline group and lactated Ringers group, but the serum lactate level after resuscitation was significantly higher in the lactated Ringers group, especially in the chronic liver disease subgroup. The serum potassium level increased in the first few hours but recovered more slowly in patients with chronic kidney disease regardless of fluid type. Blood glucose reached the target level faster and the glycemic variability was lower in the lactated Ringers group, which may suggest that lactated Ringers helps stabilize blood glucose levels during and after resuscitation.

Our analysis further confirmed that sepsis patients treated with saline had increased mortality compared to those treated with lactated Ringers\textsuperscript{2}. Evidence has also suggested that saline prolongs the ICU stay and hospital stay due to the increased risk of hyperchloremia acidosis, endothelial glycocalyx damage-related interstitial edema, renal vessel constriction-related kidney injury, gastrointestinal edema, ileus, bleeding events and the need for blood product transfusion\textsuperscript{22}. However, the reduced risk of acidosis and shortened length of ICU stay associated with the use of lactated Ringers were observed in only sepsis patients without chronic kidney disease and without chronic liver disease. This implies that intact kidney and liver functions play important roles in the efficacy of lactated Ringers in inducing acidosis prevention effects. On the other hand, the use of lactated Ringers shortened the ICU length of stay in sepsis patients with chronic pulmonary disease but not in those without; this phenomenon seems to suggest that the acidosis prevention effect of lactated Ringers is especially important for patients with chronic pulmonary disease who often and easily develop respiratory acidosis.

Osmolality is isotonic in saline (308 mOsm/kg) but hypotonic in lactated Ringers (273 mOsm/kg). The difference in osmolality did not seem to affect resuscitation efficacy, as the trends of central venous pressure and central venous oxygen saturation were not significantly different between the two fluid groups. However, the lactate levels during and after resuscitation were higher in the lactated Ringers group. This indicates that lactated Ringers may increase the serum lactate level during the resuscitation period. Thus, we cannot use the decreasing trend of serum lactate as a surrogate for perfusion.
restoration and resuscitation efficacy, especially in patients who receive lactated Ringers as the main resuscitation fluid\textsuperscript{23}.

Saline solution contains no potassium, but lactated Ringers solution contains 4 mmol/L of potassium. The risk of hyperkalemia in patients who receive lactated Ringers, especially those with chronic kidney disease, is a concern. Our analysis found that the serum potassium level increased in the first few hours, and the time to return to normal was not different between the saline and lactated Ringers groups. However, the rate of recovery was slower in chronic kidney disease patients. Another study found that the hyperkalemia risk was lower in chronic kidney disease patients among patients who received lactated Ringers\textsuperscript{24}. This implies that the acid-base effects of saline are more important for serum potassium homeostasis than those induced by the small amount of potassium in lactated Ringers fluid.

Seventy percent of lactate undergoes gluconeogenesis, resulting in a transient increase in blood glucose. Thirty percent of lactate clearance occurs via oxidation by the consumption of hydrogen ions\textsuperscript{25}, leaving \( \text{OH}^- \) to bind to \( \text{CO}_2 \) to form \( \text{HCO}_3^- \). Thus, in patients with acidosis, fluid resuscitation with lactated Ringers help limit and reverse acidosis. This effect is reversed in saline, as a high chloride load exacerbates acidosis\textsuperscript{26}. Lactated Ringers solution delays glycemic recovery in diabetic ketoacidosis, but acidosis recovery is faster with the administration of lactated Ringers solution\textsuperscript{7}. Our study found that in the lactated Ringers group, the initially high glucose level decreased to the target glucose level faster than that in the saline group. This finding is significant for both diabetes and nondiabetes patients. Lactated Ringers also decreased glycemic variability and decreased the risk for high blood glucose. This could be explained by the fact that the balanced electrolyte distribution in lactated Ringers may help stabilize blood glucose levels. Finally, in patients with moderate to severe liver disease, lactate metabolism and thus the acidosis-prevention effect were impaired. This may explain why the lactate level remained high during and after resuscitation in liver disease patients and that the lengths of ICU and hospital stays were not significantly different between the lactated Ringers and saline groups.

The strength of this study was that we collected detailed comorbidity data and implemented standard care bundles, including regularly evaluating resuscitation targets to adjust fluid volume administration, collecting baseline laboratory data, and implementing standard glucose control and glucose target protocols. This helped to control all possible confounding factors. Our study also had some limitations. First, the choice of lactated Ringers or saline was dependent on physician preference. However, physicians have favored lactated Ringers in recent years and favored saline in previous years, but our care bundles remained the same throughout the whole research period. Second, resuscitation fluid volume and types before ICU admission were not available. The APACHE score in the lactated Ringers group was higher than that in the saline group. We consider this confounding factor in the adjusted regression model.

Conclusions
Lactated Ringers use was associated with lower mortality and shorter lengths of ICU and hospital stays, and these beneficial effects were observed in patients with chronic pulmonary disease, without chronic kidney disease, without chronic liver disease and without cerebral vascular disease. Lactated Ringers use was associated with higher serum lactate levels, especially in liver disease patients, and hyperkalemia was more likely to occur in chronic kidney disease patients regardless of fluid type. The glucose levels during resuscitation were highest in diabetes patients receiving saline and lowest in nondiabetes patients receiving lactated Ringers. Comorbidities are important for clinicians to consider when choosing a fluid type.

**Abbreviations**

SHR
subdistribution hazard ratio

ICU
intensive care unit

APACHE
Acute Physiology and Chronic Health Evaluation

HbA1c
Hemoglobin A1c

CoV
coefficient of variation

MAGE
mean amplitude of glycemic excursion

**Declarations**

**Ethical Approval and Consent to participate**

Not applicable

**Consent for publication**

Not applicable

**Availability of supporting data**

All data generated or analysed during this study are included in this published article and its supplementary files.

**Competing interests**

The authors declare that they have no competing interests.
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Authors’ contributions

CH, TT and MC performed analyzed data and did the tables and figures. CH and MC confirmed comorbidities for every sepsis patients and audit the sepsis protocol implementation. KY and SM collected clinical data from electric medical records. CH and YK were major contributor in writing the manuscript. All authors read and approved the final manuscript.

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References

1. Rhodes A, Evans LE, Alhazzani W, et al. Surviving Sepsis Campaign: International Guidelines for Management of Sepsis and Septic Shock: 2016. Critical Care Medicine. 2017;45(3).
2. Semler MW, Self WH, Wanderer JP, et al. Balanced Crystalloids versus Saline in Critically Ill Adults. N Engl J Med Mar. 2018;1(9):829–39. doi:10.1056/NEJMoa1711584. 378 ).
3. Tseng CH, Chen TT, Wu MY, Chan MC, Shih MC, Tu YK. Resuscitation fluid types in sepsis, surgical, and trauma patients: a systematic review and sequential network meta-analyses. Crit Care Dec. 2020;14(1):693. doi:10.1186/s13054-020-03419-y. 24 ).
4. Myburgh JA, Mythen MG. Resuscitation fluids. N Engl J Med Dec. 2013;19(25):2462–3. doi:10.1056/NEJMmc1313345. 369 ).
5. Self WH, Semler MW, Wanderer JP, et al. Balanced Crystalloids versus Saline in Noncritically Ill Adults. N Engl J Med Mar. 2018;1(9):819–28. doi:10.1056/NEJMoa1711586. 378 ).
6. Cohen RD, Simpson R. Lactate metabolism. Anesthesiology Dec. 1975;43(6):661–73. doi:10.1097/00000542-197512000-00013.
7. Van Zyl DG, Rheeder P, Delport E. Fluid management in diabetic-acidosis—Ringers lactate versus normal saline: a randomized controlled trial. Qjm Apr. 2012;105(4):337–43. doi:10.1093/qjmed/hcr226.
8. Sterling SA, Puskarich MA, Jones AE. The effect of liver disease on lactate normalization in severe sepsis and septic shock: a cohort study. Clin Exp Emerg Med. 2015;2(4):197–202. doi:10.15441/ceem.15.025.
9. Jeppesen JB, Mortensen C, Bendtsen F, Møller S. Lactate metabolism in chronic liver disease. Scand J Clin Lab Invest. 2013;73(4):293–9. doi:10.3109/00365513.2013.773591.
10. Levy MM, Fink MP, Marshall JC, et al. 2001 SCCM/ESICM/ACCP/ATS/SIS International Sepsis Definitions Conference. Crit Care Med. Apr 2003;31(4):1250-6. doi:10.1097/01.Ccm.0000050454.01978.3b.

11. Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. J Chronic Dis. 1987;40(5):373–83. doi:10.1016/0021-9681(87)90171-8.

12. Wyss R, Ellis AR, Brookhart MA, et al. The role of prediction modeling in propensity score estimation: an evaluation of logistic regression, bCART, and the covariate-balancing propensity score. Am J Epidemiol Sep. 2014;15(6):645–55. doi:10.1093/aje/kwu181. 180).

13. Rosenbaum PR, Rubin DB. Reducing Bias in Observational Studies Using Subclassification on the Propensity Score. J Am Stat Assoc. 1984;79(387):516–24. doi:10.2307/2288398.

14. Schoenfeld D. Survival methods, including those using competing risk analysis, are not appropriate for intensive care unit outcome studies. Crit Care (London England). 2006;10(1):103–3. doi:10.1186/cc3949.

15. Resche-Rigon M, Azoulay E, Chevret S. Evaluating mortality in intensive care units: contribution of competing risks analyses. Crit Care (London England). 2006;10(1):R5–5. doi:10.1186/cc3921.

16. Fine JP, Gray RJ. A Proportional Hazards Model for the Subdistribution of a Competing Risk. J Am Stat Assoc. 1999/06/01 1999;94(446):496–509. doi:10.1080/01621459.1999.10474144.

17. Service FJ, Molnar GD, Rosevear JW, Ackerman E, Gatewood LC, Taylor WF. Mean amplitude of glycemic excursions, a measure of diabetic instability. Diabetes Sep. 1970;19(9):644–55. doi:10.2337/diab.19.9.644.

18. Lanspa MJ, Dickerson J, Morris AH, Orme JF, Holmen J, Hirshberg EL. Coefficient of glucose variation is independently associated with mortality in critically ill patients receiving intravenous insulin. Crit Care Apr. 2014;30(2):R86. doi:10.1186/cc13851. 18).

19. Kovatchev BP, Cox DJ, Gonder-Frederick LA, Clarke W. Symmetrization of the blood glucose measurement scale and its applications. Diabetes Care Nov. 1997;20(11):1655–8. doi:10.2337/diabcare.20.11.1655.

20. Kovatchev BP, Cox DJ, Kumar A, Gonder-Frederick L, Clarke WL. Algorithmic evaluation of metabolic control and risk of severe hypoglycemia in type 1 and type 2 diabetes using self-monitoring blood glucose data. Diabetes Technol Ther. 2003;5(5):817–28. doi:10.1089/152091503322527021.

21. Hill NR, Oliver NS, Choudhary P, Levy JC, Hindmarsh P, Matthews DR. Normal reference range for mean tissue glucose and glycemic variability derived from continuous glucose monitoring for subjects without diabetes in different ethnic groups. Diabetes Technol Ther Sep. 2011;13(9):921–8. doi:10.1089/dia.2010.0247.

22. Lobo DN, Awad S. Should chloride-rich crystalloids remain the mainstay of fluid resuscitation to prevent 'pre-renal' acute kidney injury?: con. Kidney Int Dec. 2014;86(6):1096–105. doi:10.1038/ki.2014.105.
23. Jones AE, Shapiro NI, Trzeciak S, Arnold RC, Claremont HA, Kline JA. Lactate clearance vs central venous oxygen saturation as goals of early sepsis therapy: a randomized clinical trial. Jama Feb. 2010;24(8):739–46. doi:10.1001/jama.2010.158. 303 ) .

24. Toporek AH, Semler MW, Self WH, et al. Balanced Crystalloids versus Saline in Critically Ill Adults with Hyperkalemia or Acute Kidney Injury: Secondary Analysis of a Clinical Trial. Am J Respir Crit Care Med Jan 27 2021;doi:10.1164/rccm.202011-4122LE.

25. Hartmann AF, Senn MJ. Studies in the metabolism of sodium r-lactate. III. Response of human subjects with liver damage, disturbed water and mineral balance, and renal insufficiency to the intravenous injection of sodium r-lactate. J Clin Invest Mar. 1932;11(2):345–55. doi:10.1172/jci100416.

26. Yunos NM, Kim IB, Bellomo R, et al. The biochemical effects of restricting chloride-rich fluids in intensive care. Crit Care Med Nov 2011;39(11):2419–24. doi:10.1097/CCM.0b013e31822571e5.

**Figures**
Figure 1

Trends for central venous pressure, central venous oxygen saturation, serum bicarbonate, lactate, creatinine changes and urine output among fluid types and diabetes status. Abbreviations: LR, lactated Ringer's.
Figure 2

Trends for (A) serum potassium among fluid types and chronic kidney status, (B) blood sugar changes among fluid types and diabetes status, (C) serum lactate in patient with or without moderate to severe liver disease

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