Hyperchloremia in Intensive Care Unit Mortality: An Underestimated Fact

Zahid Siddique Shad ¹, Muhammad Shehaib Saffar Qureshi ¹, Aayesha Qadeer ¹, Azmat Abdullah ¹, Kamran Munawar ¹, Muhammad Tariq Khan ¹, Luqman Saeed ¹, Syed Waqar Hussain ²

1. Internal Medicine, Shifa International Hospital, Islamabad, PAK 2. Internal Medicine, Khan Research Laboratories Hospital, Islamabad, PAK

Corresponding author: Zahid Siddique Shad, zahids80@gmail.com

Abstract

Objective

The goal of the study was to determine the percentage of hyperchloremia in patients who died in medical intensive care unit (ICU) and thus emphasizing the need of avoiding chloride-rich solutions due to their deleterious effects.

Study design

We conducted a retrospective study of data from 206 patients who expired in medical ICU in one year from January 2017 to December 2017 in the department of critical care medicine at Shifa International Hospital, Islamabad.

Material and methods

The study included 206 patients: 93 (43.1%) men and 123 (56.9%) women, over the age of 18 years who expired in medical ICU in one year from January 2017 to December 2017. Patients included for the study were all those who expired with any diagnosis but those who remained admitted in ICU for at least 72 hours and received intravenous fluids. The serum chloride levels of the patients at the time of admission and at 72 hours of stay in ICU were collected. The patients who were having serum chloride levels of 107 milliequivalent per deciliter (meq/dl) or more were labeled as having hyperchloremia. The data were analyzed using IBM SPSS Statistics for Windows, Version 23.0 (IBM Corp., Armonk, NY, USA). The mean and standard deviations were calculated for continuous variables while frequency and percentages were calculated for qualitative variables.

Results

Among 206 patients who expired in our ICU, 109 (50.5%) patients had hyperchloremia at 72 hours of admission in ICU while 107 (49.5%) patients did not had hyperchloremia. Hyperchloremia was more frequent in patients with sepsis or septic shock.

Conclusion

Higher percentage (50.5%) of hyperchloremia at 72 hours of admission among patients (who expired in our medical ICU) indicates excessive use of chloride-rich intravenous fluids. This finding may have significant impact on mortality along with other contributing factors that lead to death of the patients. Keeping in view the findings of the study, chloride-rich solutions should be used carefully to counter the effects of hyperchloremia in patients requiring large volume fluid resuscitation in ICU. Fluids with lower content of chloride such as lactated ringer may be equally good in large volume fluid resuscitation with advantage of avoiding hyperchloremia.

Introduction

Chloride is one of the most important and the most abundant anion in the extracellular fluid contributing approximately one-third extracellular fluid tonicity [1]. Chloride plays several physiologically important roles in the body which include acid-base balance, immune modulation, osmosis and muscular activity [2]. Acid base homeostasis is the main function, chloride plays in the body. Chloride levels are inversely related to bicarbonate which is thought to be the major buffer in human body. So, increasing or decreasing levels of chloride have direct impact on acid base status, resulting in non-anion gap metabolic acidosis and metabolic alkalosis, respectively.

How to cite this article

Shad Z, Qureshi M, Qadeer A, et al. (May 28, 2019) Hyperchloremia in Intensive Care Unit Mortality: An Underestimated Fact. Cureus 11(5): e4770. DOI 10.7759/cureus.4770
In gastrointestinal tract, chloride has two unique functions: first, it is secreted as a part of hydrochloric acid in the stomach and helps in protein digestion, keeping a check on microorganism and absorption of some of the important nutrients/minerals [2]; second, it maintains the gastrointestinal osmotic gradient and fluid secretion. Other effects of chloride are on unloading of oxygen from hemoglobin, immune modulation and cardiovascular stability.

Serum chloride levels are maintained in normal range mainly by the kidneys and gastrointestinal system. Dyschloremia in intensive care unit (ICU) is a common observation in patients who are receiving intravenous fluids or diuretics. Hyperchloremia is somewhat more observed as compared to hypochloremia in ICU patient. Despite its physiologically important functions, chloride has caught little or no attention by the researchers until recently [3] when association of chloride-rich solutions to hyperchloremia metabolic acidosis has been established [4, 5]. It has been found that hyperchloremia increases short-term mortality after non-cardiac surgery [6, 7].

Normal saline (0.9% saline) is the chloride-rich solution most commonly used in clinical practice especially in critically ill patients and preoperatively. Contrary to the name, 0.9% saline in reality is not normal or neutral solution [8]. It has got chloride in supra physiological amount when we compare it with plasma (154 vs. 100 meq/L, respectively) [9, 10]. The liberal use of such a chloride-rich solution results in hyperchloremic metabolic acidosis in critically ill patients [11, 12] with several detrimental effects [13, 14] especially in patients with severe sepsis and septic shock [15]. Hyperchloremia has been associated with acute kidney injury in ICU patients and likewise severe acidosis when present can result in decreased efficacy of inotropes as well as cardiovascular dysfunction. All these deleterious effects of hyperchloremia increase in mortality. Septic patients in critical condition are commonly exposed to 0.9% saline during the salvage phase of septic shock and are at risk of hyperchloremic metabolic acidosis and other detrimental effects of hyperchloremia in the post-resuscitation phase. Observational studies evaluating the association of hyperchloremia with hospital mortality have shown conflicting results and included only a small number of septic patients in the ICU [16-18].

In our ICU, we commonly use normal saline to resuscitate patients. The purpose of the study was to find in what percentage of expired patients hyperchloremia is present so that we may change our practice of using chloride-rich solution and prefer solution with acceptable levels of chloride (lactated ringer or plasmalyte) to avoid this iatrogenic complication which may benefit our patients in future.

Materials And Methods

We conducted a retrospective review of patients' medical records at the department of critical care medicine at Shifa International Hospital Islamabad. The ethical review board of Shifa International Hospital Islamabad provided ethical approval for the study. We reviewed mortality data in ICU from January 2017 to December 2017. The inclusion criteria entitled all patients eligible who expired in ICU due to any disease/diagnosis in the specified duration of study but an essential stay in ICU for at least 72 hours before expiration and receiving intravenous fluids was a must. Two hundred and six patients met the inclusion criteria for the research. Of these 93 (43.1%) were male and 123 (56.9%) were female. The minimum age was 18 years and maximum age was 100 years. The mean age of the patients was 60.62 years with standard deviation of 14.784. Mean chloride level of the patients at the time of admission in ICU was 107.5093 meq/dl. We divided the patients, on the basis of diagnosis, into two groups: 1) Sepsis and septic shock group, and 2) diagnosis other than sepsis. One hundred and nineteen patients were in sepsis and septic shock group and 97 patients were in group with diagnosis other than sepsis. All these patients received intravenous fluids. Hyperchloremia was present in 109 (50.5%) patients and 107 (49.5%) patients were not having hyperchloremia at 72 hours of ICU admission. Hyperchloremia was present among 97 patients out of 119 patients in sepsis/septic shock group and 12 patients out of 85 in patients with diagnosis other than sepsis. Among men 50 (55.76%) out of 93 developed hyperchloremia and among women 59 (47.96%) out of 125 developed hyperchloremia at 72 hours of ICU admission. The detailed results are shown in Tables 1-6.
|                         | N   | Minimum | Maximum | Mean   | Std. Deviation |
|-------------------------|-----|---------|---------|--------|----------------|
| Age                     | 216 | 18      | 100     | 60.62  | 14.784         |
| Chloride level at admission | 216 | 84.00   | 115.00  | 99.7963| 3.99362        |
| Chloride levels at 72 hours | 216 | 92.00   | 119.00  | 107.5093| 5.86514        |
| Valid N (listwise)      | 216 |         |         |        |                |

**TABLE 1:** Descriptive statistics for age, chloride level at admission and chloride levels at 72 hours.

|                         | N   | Minimum | Maximum | Mean   | Std. Deviation |
|-------------------------|-----|---------|---------|--------|----------------|
| Age                     | 216 | 18      | 100     | 60.62  | 14.784         |
| Chloride level at admission | 216 | 84.00   | 115.00  | 99.7963| 3.99362        |
| Chloride levels at 72 hours | 216 | 92.00   | 119.00  | 107.5093| 5.86514        |
| Valid N (listwise)      | 216 |         |         |        |                |

**TABLE 2:** Chloride levels at admission and at 72 hours.

|                           | Frequency | Percent | Valid Percent | Cumulative Percent |
|---------------------------|-----------|---------|---------------|--------------------|
| Hyperchloremia present    | 109       | 50.5    | 50.5          | 50.5               |
| Hyperchloremia not present| 107       | 49.5    | 49.5          | 100.0              |
| Total                     | 216       | 100.0   | 100.0         |                    |

**TABLE 3:** Frequency of hyperchloremia among patients at 72 hours of admission.

|                           | Frequency | Percent | Valid Percent | Cumulative Percent |
|---------------------------|-----------|---------|---------------|--------------------|
| Male                      | 93        | 43.1    | 43.1          | 43.1               |
| Female                    | 123       | 56.9    | 56.9          | 100.0              |
| Total                     | 216       | 100.0   | 100.0         |                    |

**TABLE 4:** Frequency of gender of the patients.
Diagnosis

|                  | Sepsis/septic shock | Other than sepsis | Total |
|------------------|---------------------|-------------------|-------|
| Hyperchloremia   | 97                  | 12                | 109   |
| Hyperchloremia not present | 22          | 85                | 107   |
| Total            | 119                 | 97                | 216   |

TABLE 5: Hyperchloremia and diagnosis--cross tabulation.

| Gender       | Total |
|--------------|-------|
| Male         | 93    |
| Female       | 123   |
| Total        | 216   |

TABLE 6: Hyperchloremia and gender of the patient–cross tabulation.

Discussion

Chloride is one of the most important and the most abundant anion in the extracellular fluid contributing approximately one-third extracellular fluid tonicity [1]. Chloride plays several physiologically important roles in the body which include acid-base balance, immune modulation, osmosis and muscular activity [2]. Acid base homeostasis is the main function chloride plays in the body. Large volume fluid resuscitation is common in ICU and selection of fluid type is very important in this regard. If chloride-rich solutions are used for large volume resuscitation in septic patients, it may result in hyperchloremia and its untoward effects.

A number of studies have pinpointed the hazardous effects of hyperchloremia in critically ill patients. In a large retrospective study by Neyra et al., deleterious effects of hyperchloremia in ICU patients were studied. The study included 1940 patients. Of these 615 (31.7%) patients had hyperchloremia on ICU admission and 1325 (68.3%) patients were having no hyperchloremia at the time of admission in ICU. Patients were observed from the time of admission in ICU until discharge or death. During observation period, 431 (23.9%) patients died that included 147 (23.9%) patient in subgroup of hyperchloremia and 284 (21.4%) patients in subgroup without hyperchloremia. This significant mortality difference in both groups suggests hyperchloremia increases mortality in ICU patients. Moreover, patients in hyperchloremia subgroup required more vasopressors support, higher number of blood transfusions, prolonged stay in ICU due to more days on mechanical ventilation and these patients were having more frequent oliguria. These observations along with other studies signify the importance of maintaining physiological range of serum chloride in ICU patients [19].

A large, cluster-randomized, multiple-crossover trial conducted by Semler et al., compared normal saline and balanced crystalloids (lactated ringer or plasma lyte A) for fluid resuscitation in ICU. In this research, 15802 patients in five ICUs were randomized to receive either normal saline (0.9% sodium chloride solution) or balanced crystalloids fluids (lactated Ringer’s solution or Plasma-Lyte A). The primary outcome of the study was to find a major adverse kidney event within 30 days, new renal replacement therapy (RRT) or persistent renal dysfunction defined as elevation of serum creatinine level to ≥200% of baseline or death from any cause. All information were analyzed at the time of discharge/death or 30 days whichever came first. The results were in favor of subgroup with balanced crystalloid administration versus normal saline. Among the 7942 patients in balanced-crystalloid group, 14.3% (1139 patients) had a major kidney-related event while in the normal saline group the percentage of the same was 15.4% (1211 out of 7860 patients). Considering other parameters, 30 days mortality in hospital was 10.3% in the balanced saline group and 11.1% in the saline group (p-value 0.06). The incident of new renal replacement therapy was 2.5% in the balanced saline group and 2.9% in the normal saline group (p-value 0.08) and the incidence of persistent renal dysfunction was 6.4% and 6.6% (p-value 0.60), respectively. The researchers concluded that balanced crystalloid solution has an edge over normal saline in terms of causing less organ dysfunction [20].
In our study, we reviewed the mortality data of those patients who remained in ICU for at least 72 hours before expiration and essentially received intravenous fluids during the stay. The fluid was normal saline exclusively. The purpose was to find how many patients developed hyperchloremia at 72 hours of stay in ICU while receiving intravenous fluids. The study results were astonishing: a significantly higher percentage of hyperchloremia (50.5%) was noted among patients at 72 hours of stay in ICU. It signifies that we are overusing chloride-rich solution: almost exclusively 0.9% saline which has supra physiological content of chloride that can lead to dose dependent hyperchloremic metabolic acidosis and other hazards. Reviewing the available literature on effects of hyperchloremia in critically ill patients, the situation seems alarming and emphasizes that we have to be careful in selection of fluid while we need to use large volume fluid resuscitation in our patients. One more interesting aspect of the study result was lower frequency of hyperchloremia in patients who were having diagnosis other than sepsis that included all other patients except sepsis or septic shock. The reason was overall less fluid administration when compared to the group with diagnosis of sepsis.

Conclusions

Keeping in view the findings of the study, chloride-rich solutions should be used carefully to counter the effects of hyperchloremia in patients requiring large volume fluid resuscitation in ICU. Balanced crystalloids (lactated ringer or plasmalyte) are equally good choices for large volume resuscitations in critically ill patients with lower risk of hyperchloremia and its effects. The study has some weaknesses as regard to relation of hyperchloremia and total amount of fluids administered. Yet the finding of higher frequency of hyperchloremia in patients who died in ICU clearly advocates using intravenous fluids with low content of chloride to avoid the evitable iatrogenic adverse effect of fluid resuscitation when appropriate.

Additional Information

Disclosures

Human subjects: Consent was obtained by all participants in this study. Institutional Review Board & Ethics Committee Shifa International Hospital, Islamabad, Pakistan issued approval IRB#994-269-2018. Institutional Review Board and Ethics Committee, Shifa International Hospital, Islamabad, approved your study entitled as ‘Hyperchloremia in ICU Mortality: An Underestimated Fact’. Animal subjects: All authors have confirmed that this study did not involve animal subjects or tissue. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

Acknowledgements

I am highly obliged for the continuous support of Dr. Ayesha Qadeer and Dr. Shehar Bano throughout the research work.

References

1. Hall JE, Guyton AC: Guyton and Hall Textbook of Medical Physiology. Saunders/Elsevier, Philadelphia; 2011.
2. Berend K, van Hulsteijn LH, Gans RO: Chloride: the queen of electrolytes?. Eur J Intern Med. 2012, 23:203-211. 10.1016/j.ejim.2011.11.015
3. Yunon NM, Bellomo R, Story D, Kellum J: Bench-to-bedside review: chloride in critical illness. Critical Care. 2010, 14:226. 10.1186/cc9502
4. Scheinragher S, Rehn M, Sehmisch G, Finsterer U: Rapid saline infusion produces hyperchloremic acidosis in patients undergoing gynecologic surgery. Anesthesiology. 1999, 90:1265-1270.
5. Mann C, Held U, Herzog S, Baenziger O: Impact of normal saline infusion on postoperative metabolic acidosis. Paediatr Anaesth. 2009, 19:1070-1077. 10.1111/j.1460-9592.2009.03126.x
6. Silva Junior JM, Neves EF, Santana TC, Ferreira UP, Marti YN, Silva JM: The importance of intraoperative hyperchloremia. (Article in English, Portuguese). Rev Bras Anestesiol. 2009, 59:304-315. 10.1590/S0034-79942009005000005
7. McCluskey SA, Karkoulti K, Wijeyunadera D, et al.: Hyperchloremia after noncardiac surgery is independently associated with increased morbidity and mortality: a propensity-matched cohort study. Anesth Analg. 2015, 117:412-421. 10.1213/ANE.0b013e318295d81e
8. Story DA, Thistlethwaite P, Bellomo R: The effect of PVC packaging on the acidity of 0.9% saline . Anaesth Intensive Care. 2000, 28:287-292. 10.1111/j.1460-9592.2000.tb08006.x
9. Guidet B, Soni N, Deilla Rocca G, Koeck S, Vallet B, Annane D, James M: A balanced view of balanced solutions. Critical Care. 2010, 14:325. 10.1186/cc9230
10. Veech RL: The toxic impact of parenteral solutions on the metabolism of cells: a hypothesis for physiological parenteral therapy. Am J Clin Nutr. 1986, 44:519-551. 10.1093/ajcn/44.4.519
11. Moviat M, van den Hoogaard M, Intven F, van der Voort P, van der Hoeven H, Pickkers P: Stewart analysis of apparently normal acid-base state in the critically ill. J Crit Care. 2013, 28:1048-1054. 10.1016/j.jcrc.2013.06.005
12. O’Dell E, Tibby SM, Durward A, et al.: Hyperchloremia is the dominant cause of metabolic acidosis in the postresuscitation phase of pediatric meningococcal sepsis. Crit Care Med. 2007, 35:2390-2394. 10.1097/01.CCM.0000284588.17760.99

13. Lobo DN, Awad S: Should chloride-rich crystalloids remain the mainstay of fluid resuscitation to prevent ‘pre-renal’ acute kidney injury?: con. Kidney Int. 2014, 86:1096-1105. 10.1038/ki.2014.105

14. Krajewski ML, Raghunathan K, Paluszakwicz SM, Schermer CR, Shaw AD: Meta-analysis of high- versus low-chloride content in perioperative and critical care fluid resuscitation. Br J Surg. 2015, 102:24-36. 10.1002/bjs.9651

15. Noritomi DT, Soriano FG, Kellum JA, Cappi S, Biselli P, Libório A, Park M: Metabolic acidosis in patients with severe sepsis and septic shock: a longitudinal quantitative study. Crit Care Med. 2009, 37:2733-2739. 10.1097/CCM.0b013e3181a59163

16. Bonitatti MM, Cardoso PR, Castilho RK, Vieira SR: Is hyperchloremia associated with mortality in critically ill patients? A prospective cohort study. J Crit Care. 2011, 26:175-179. 10.1016/j.jcrc.2010.04.013

17. Tani M, Morimatsu H, Takatsu F, Morita K: The incidence and prognostic value of hypochloremia in critically ill patients. Scientific World J. 2012, 2012:7. 10.1100/2012/474185

18. Shaw AD, Raghunathan K, Peyerl FW, Munson SH, Paluszakwicz SM, Schermer CR: Association between intravenous chloride load during resuscitation and in-hospital mortality among patients with SIRS. Intensive Care Med. 2014, 40:1897-1905. 10.1007/s00134-014-3505-3

19. Neyra JA, Canepa-Escarco F, Li X, Manullo J, Adams-Huet B, Yee J, Yessayan L: Association of hyperchloremia with hospital mortality in critically ill septic patients. Critical Care Med. 2015, 43:1938-1944. 10.1097/CCM.0000000000001161

20. Semler MW, Self WH, Wanderer JP, et al.: Balanced crystalloids versus saline in critically ill adults. N Engl J Med. 2018, 378:829-839. 10.1056/NEJMoA1711584