Ion-selective electrode and anion gap range: What should the anion gap be?

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Background: Using flame photometry technique in the 1970s, the normal value of anion gap (AG) was determined to be $12 \pm 4$ meq/L. However, with introduction of the autoanalyzers using an ion-selective electrode (ISE), the anion gap value has fallen to lower levels.

Methods: A retrospective study of US veterans from a single medical center was performed to determine the value of the anion gap in subjects with normal renal function and normal serum albumin and in patients with lactic acidosis and end-stage renal disease on dialysis.

Results: In 409 patients with an estimated glomerular filtration rate $\geq 60$ mL/min/1.73 m$^2$ body surface area and serum albumin $\geq 4$ g/dL, the mean AG was $7.2 \pm 2$ (range 3–11) meq/L. In 299 patients with lactic acidosis (lactate level $\geq 4$ meq/L) and 68 patients with end-stage renal disease on dialysis, the mean AG was 12.5 meq/L and 12.4 meq/L, respectively. A value $< 2$ meq/L should be considered a low anion gap and a possible clue to drug intoxication and paraproteinemic disorders.

Conclusion: With the advent of ISE for measurement of analytes, the value of the anion gap has fallen. Physicians need to be aware of the normal AG value in their respective institutions, and laboratories need to have an established value for AG based on the type of instrument they are using.

Keywords: acidosis, electrolytes, ESRD

Introduction

The anion gap (AG) represents the difference between the measured anion and cation concentrations in the serum, and is used by clinicians to diagnose acid base problems, especially in unsuspected cases of acidosis, alkalosis, drug poisoning, and disease states, particularly in emergency rooms and intensive care units,\textsuperscript{1–5} and as a quality control measure by laboratories. Although the concept of AG arose from the gamblegram described in 1939, it did not gain widespread recognition by physicians until the 1970s after the introduction of autoanalyzers and rapid availability of measuring multiple analytes. Originally, it was the difference between serum sodium and potassium on the one hand and chloride and bicarbonate concentrations on the other, but later on, it was simplified to the difference between sodium and the sum of chloride and bicarbonate concentrations, mainly because of the narrowness of the range of normal potassium concentration.

Prior to the introduction of the ion-selective electrode in the 1970s, flame photometry was the technique most often used to measure serum electrolytes. In a study of 1047 hospitalized patients, Wittle et al\textsuperscript{6} reported a mean AG value of 12 meq/L with a 90% range of 8–17 meq/L. This and other studies established the normal value.
for AG as 12 ± 4 meq/L. Compared with flame photometry, the ion-selective electrode (ISE) tends to give a higher concentration of serum chloride, so the AG value has fallen over time. We and others have shown that the actual value of the AG is much narrower than the currently published value3-11,13,14 (Table 1). For this reason, we examined the AG value in a random sample of patients at a single United States Veterans Administration hospital.

Materials and methods
This study was a retrospective review of serum electrolytes and anion gap in 409 outpatients aged older than 18 years with an estimated glomerular filtration rate (GFR) of > 60 mL/1.73 m² body surface area and serum albumin ≥4 g/dL. We also reviewed serum electrolytes in 68 dialysis-dependent outpatients and 299 inpatients with lactic acidosis, defined as serum lactate level ≥4 meq/L. The value of 4 meq/L was chosen to avoid including patients who had hyperlactatemia but not lactic acidosis. Serum electrolyte review was done on one random sample. Chemical analysis of serum was done using a Unicell DXC 800 autoanalyzer (Beckman Coulter, Inc., Fullerton, CA, USA) measuring sodium (Na), potassium (K), and chloride (Cl) by indirect ion-selective potentiometry.15 Serum albumin was measured by bromocresol purple methodology and serum creatinine by the selective potentiometry.

Results
In the 409 patients with normal kidney function and serum albumin, the following was found: mean age ± 1 standard deviation (SD) 52.8 ± 15 years, male/female ratio 93.4%/6.6%, Na 139.4 ± 2.3 meq/L, K 4.3 ± 0.3 meq/L, Cl 103.6 ± 3.7 meq/L, CO₂ content 28.5 ± 2.3 meq/L, glucose 100 ± 30.8 mg/dL, calcium 9.5 ± 0.3 mg/dL, albumin 4.2 ± 0.17 g/dL, blood urea nitrogen 13 ± 13 mg/dL, and creatinine 0.96 ± 0.1 mg/dL.

The mean anion gap in this group was 7.2 ± 2 (range 3.2–11.2) meq/L. In patients with lactic acidosis (serum lactate ≥4 meq/L), mean age was 68 ± 11.7 years, with a male to female ratio of 96%/4%, serum Na 137.9 ± 5.8 meq/L, K 4.5 ± 1 meq/L, Cl 105 ± 7.3 meq/L, CO₂ 19.9 ± 6.0 meq/L, glucose 161 ± 89 mg/dL, calcium 8.3 ± 5.3 mg/dL, albumin 2.4 ± 0 g/dL, blood urea nitrogen 36 ± 26.7 mg/dL, creatinine 2.2 ± 1.6 mg/dL, and lactate 7.0 ± 3.8 meq/L. In this group, uncorrected for serum albumin, the mean anion gap was 12.5 ± 8.0 (range 2–32) meq/L. Uncorrected AG was >15 in 94 (31%), 10–14 in 124 (41.4%), and <10 in 81 (27%) patients. In patients with end-stage renal disease, the mean age was 65.4 ± 9.6 years, with a male/female ratio of 98.6%/1.4%, diabetes 77.9%, hypertension 100%, congestive heart failure 21%, hepatitis C infection 11.7%, Na 138.1 ± 2.7 meq/L, K 4.9 ± 0.7 meq/L, Cl 100.6 ± 3.5 meq/L, CO₂ 25.1 ± 2.9 meq/L, glucose 144 ± 65.3 mg/dL, albumin 3.4 ± 0.5 g/dL, blood urea nitrogen 56.5 ± 20.5 mg/dL, and creatinine 9.4 ± 3.4 mg/dL.

In this group, mean AG was 12.4 ± 3.2 (range 6–26) meq/L (Tables 2 and 3). Uncorrected for serum albumin, AG was >15 in 21 (31%), 10–14 in 34 (50%), and <10 meq/L in 13 (19%) patients. An AG <2, although rare, would be considered to be a low anion gap and a possible clue to disease entities like immunoglobulin G myeloma, and bromide or lithium intoxication.3,12

Discussion
Serum AG is a useful parameter for physicians to decipher acid base disorders and gain clues to the presence of unsuspected cases of alkalosis, acidosis, mixed acid base disorders, drug intoxication, and sometimes a clue to the presence of serious diseases. Other methods, like Stewart’s method of measuring strong ion gap and strong ion difference,16-18 acid base nomograms, the slide rule,19 and base excess, are more cumbersome, complicated, and not readily accepted or used by physicians. Some physicians, particularly those in critical care, feel more comfortable with

Table 1 Studies describing anion gap values measured by an ion-selective electrode

| Author            | Year of publication | Patients (n)          | Anion gap (mean ± SD or range), meq/L | Instrument used       |
|-------------------|---------------------|-----------------------|---------------------------------------|-----------------------|
| Winter et al15     | 1990                | 29 healthy individuals | 5.9 ± 1.2                             | ASTRA                 |
|                    |                     | 120 blood donors      | 6.0 ± 1.4                             | Beckman               |
| Sadjadi3          | 1995                | 222                   | 6.6 ± 2.0                             | Beckman Synchron CX5  |
| Lolekha et al10    | 2001                | 124                   | 5–12                                  | Beckman Synchron CX7  |
| Hassan et al12     | 2004                | 286                   | 7.0 ± 2.2                             | Beckman Synchron CX5  |

Abbreviation: SD, standard deviation.
the Stuart formula and find it more helpful than the AG for detection of unmeasured anions.

However, the critical question is what is the normal mean and range for the AG. If the traditional value of 12 ± 4 meq/L is used, clues to the presence of occult acid base problems will be missed, particularly in emergency rooms and intensive care units where time is of the essence for appropriate and efficient provision of medical care. Some research suggests that lower AG values have more sensitivity and specificity in the diagnosis of acid base problems, while others do not. This study, along with our previous study incorporating 631 patients, shows the actual value of the AG range for each test.

Table 2 Patient demographics and laboratory data, shown as the mean ± SD

| Parameter                  | Normal kidney function | Lactic acidosis | End-stage renal disease |
|----------------------------|------------------------|----------------|-------------------------|
| n                          | 409                    | 299            | 68                      |
| Age, years                 | 52.8 ± 15              | 68 ± 11.7      | 65.4 ± 9.6              |
| Male/female, %             | 93.4/6.6               | 96/4.0         | 98.6/1.4                |
| Sodium, meq/L              | 139.4 ± 2.3            | 137.9 ± 5.8    | 138.1 ± 2.7             |
| Potassium, meq/L           | 4.3 ± 0.3              | 4.5 ± 1.0      | 4.9 ± 0.7               |
| Chloride, meq/L            | 103.6 ± 3.7            | 105.7 ± 7.3    | 100.6 ± 3.5             |
| CO₂, meq/L                 | 28.5 ± 2.3             | 19.9 ± 6       | 25.1 ± 2.9              |
| Glucose, mg/dL             | 100 ± 30.8             | 161 ± 89       | 144 ± 65.3              |
| Albumin g/dL               | 4.3 ± 0.2              | 2.4 ± 0        | 3.4 ± 0.5               |
| BUN mg/dL                  | 13 ± 13                | 36 ± 26.7      | 56.5 ± 20.5             |
| Creatinine, mg/dL          | 0.96 ± 0.1             | 2.2 ± 1.6      | 9.4 ± 3.5               |

Abbreviations: SD, standard deviation; BUN, blood urea nitrogen.

Table 3 Mean anion gap in patients with normal renal function, ESRD, and lactic acidosis

| Group                        | AG, meq/L; mean ± SD | AG range, meq/L Uncorrected for serum albumin |
|------------------------------|----------------------|-----------------------------------------------|
| eGFR >60 mL per minute       | 7.2 ± 2.0            | 3–11                                          |
| ESRD                         | 12.4 ± 3.2           | 6–26                                          |
| Lactic acidosis              | 12.5 ± 8.0           | 2–32                                          |

Abbreviations: ESRD, end-stage renal disease; eGFR, estimated glomerular filtration rate; SD, standard deviation; AG, anion gap.

have a normal AG range of their own and use the value recommended by the manufacturer of the instrument, which could vary from 10 to 20 meq/L. For this reason, we believe the value of the AG needs to be lowered to 6–7 meq/L and that every laboratory should have an established value for AG. Clinicians should know which instrument is used in their laboratory and what is the normal AG value for their institution. In a review of 11,597 adults who participated in the National Health and Nutrition Examination Survey, conducted between 1999 and 2004 and using ISE, the mean value for AG was 12 meq/L. However, it is not mentioned in this report if the value of chloride was corrected for ISE overestimation.22

Why is the anion gap narrower? This relates to the fact that ISE gives a higher reading for chloride concentration compared with flame photometry and colorimetric techniques. In the study reported by Winter et al,7 the average serum chloride of 423 hospital employees measured using the ASTRA autoanalyzer (Beckman Coulter) was 107 meq/L, while the reference serum chloride done using the Technicon instrument (Technicon Corporation, Tarrytown, NY, USA) in a previous study was 101 meq/L. Lolekha et al point out that the lower AG value obtained with ISE analyzers compared with continuous flow analyzers is partly due to a greater loss of carbon dioxide with the latter types of instrument.8 The AG value is also dependent on the type of instrument used to measure its components. In the study reported by Roberts et al, the AG value was 5–10 with the Synchrom CX3 analyzer (Beckman Coulter), 9–14 for the Hitachi 717 analyzer (Boehringer Mannheim, Mannheim, Germany), and 8–13 meq/L for the Vitros 950 analyzer (Johnson and Johnson, New Brunswick, NJ, USA).14

Another issue is the effect of serum albumin and other proteins on the anion gap. As calculated by Siggard-Anderson et al, albumin accounts for 73%, globulins for 22%, and phosphate for 5% of the buffer capacity of the blood.23 This has traditionally required an upward adjustment of 2.04 to 3 meq/L24–28 in the anion gap for every g/dL decrease in serum albumin. In a study of 5328 patients, 70% of whom had normal serum albumin (3.5–4.7 g/dL), 21.7% had hypoalbuminemia and 8% had hyperalbuminemia; the relationship between serum albumin and AG was linear, with a slope of 2.3 meq/L change in AG per g/dL change in serum albumin concentration.29

Some authors have questioned the value of correcting AG for serum albumin while others have found it to be useful. In their study of intensive care patients, Dinh et al did not find much improvement in detecting hyperlactatemia using
albumin corrected or uncorrected AG.30 pH also affects the charge on albumin but the effect is small, in the range of 0.5 meq/L per 0.1 unit change in pH.31 As pointed out by Rastegar, in spite of its pitfalls, AG when corrected for serum albumin, is more sensitive than the Stuart method or base excess for delineation of acid base disorders.16

As expected, in our patients with lactic acidosis and end-stage renal disease, AG was higher, with a mean value of 12.5 meq/L. Because our study was retrospective and serum lactate levels and arterial blood gases were not routinely measured in all critically ill patients with lactic acidosis, we did not attempt to calculate the sensitivity and specificity of our findings. Some studies have reported sensitivity and specificity data based on retrospective observations. In our view, this is not appropriate and should be avoided. Certainly, no system is perfect, but we believe the time has come to change the AG value to a lower one to maintain its utility.32

Our study has some limitations. It was done in a single Veterans Administration medical center where the overwhelming majority of patients are elderly males, so our results may not be applicable to all patient groups, but the study also has some strengths. Combined with our previous research,8 it examines the value of the anion gap in almost 1000 patients using the current generation of autoanalyzers and compares the AG value in patients with normal kidney function, lactic acidosis, and end-stage renal disease.

Disclosure
The authors report no conflicts of interest in this work.

References
1. Kraut J, Medias N. Serum anion gap: its uses and limitations in clinical medicine. Clin J Am Soc Nephrol. 2007;2:162–174.
2. Van Hoeven KH, Joseph RE, Gaughan WJ, et al. The anion gap and routine serum protein measurements in monoclonal gammapathies. Clin J Am Soc Nephrol. 2011;6:2814–2821.
3. Vasuyattakul S, Lertpattanasuwan N, Vareesangthi K, et al. A negative anion gap as clue to diagnosis of bromide intoxication. Nephron. 1995;69:311–313.
4. Gabow PA. Disorders associated with an altered anion gap. Kidney Int. 1983;27:472–485.
5. Fenves AZ, Kirkpatrick HM, Patel VV, Sweetman L, Emmett M. Increased anion gap metabolic acidosis as a result of 5-oxoproline (pyroglutamic acid): a role for acetaminophen. Clin J Am Soc Nephrol. 2006;1:441–447.
6. Wittle DL, Rodgers JL, Barrett DA. The anion gap: its use in quality control. Clin Chem. 1976;22:643–648.
7. Winter SD, Pearson JR, Gabow PA, Schultz AL, Lepoff RB. The fall of serum anion gap. Arch Intern Med. 1990;150:311–313.
8. Sadjadi SA. A new range for anion gap. Ann Intern Med. 1995;123:807.
9. Lolekha PH, Vanavan SAN, Lolekha S. Update on value of anion gap in clinical diagnosis and laboratory evaluation. Clin Chim Acta. 2001;307:133–136.
10. Hassan H, Joh JH, Bacon BR, Bastani B. Evaluation of serum anion gap in patients with liver cirrhosis of diverse etiologies. Mt Sinai J Med. 2004;71:281–284.
11. Jurado RL, del Rio C, Nassar G, Navarettes J, Pimental IL. Low anion gap. South Med J. 1998;91:624–629.
12. Kelleher SP, Raciti A, Arbib LA. Reduced or absent serum AG as a marker of severe lithium carbonate intoxication. Arch Intern Med. 1986;146:1939–1940.
13. Lee S, Kang KP, Kang SK. Clinical usefulness of the serum anion gap. Electrolyte Blood Press. 2006;4:44–46.
14. Roberts WL, Johnson RD. Serum anion gap. Has the reference value really fallen. Arch Pathol Lab Med. 1997;121:568–572.
15. Fortgens P, Pillay TS. Pseudohyponatremia revisited, a modern day pitfall. Arch Pathol Lab Med. 2011;135:516–519.
16. Rastegar A. Clinical utility of Stewart’s method in diagnosis and management of acid base disorders. Clin J Am Soc Nephrol. 2009;4:1267–1274.
17. Lloyd P. Strong ion calculator – a practical bedside application of modern quantitative acid-base physiology. Crit Care Resusc. 2004;6:285–294.
18. Kaplan LJ, Frangos S. Clinical review: acid-base abnormalities in the intensive care unit. Crit Care. 2005;9:198–203.
19. Di Iorio C, Rufolo L, Mellilo EM, Granata A, Mellilo G. The slide rule: a new method for the assessment of acid base equilibrium disorders. Minerva Anestesiol. 2007;73:339–342.
20. Singh RN, Singh NC, Hutchinson J, Moses GC. Lower anion gap increases sensitivity in predicting elevated lactate levels. Clin Intensive Care. 1994;5:221–224.
21. Adams BD, Bonzani TA, Hunter CJ. The anion gap does not accurately screen for lactic acidosis in emergency department patients. Emerg Med. 2006;23:179–182.
22. Abramowitz M, Hostetter T, Melamed ML. The serum anion gap is altered in early kidney disease and associates with mortality. Kidney Int. 2012;82:701–709.
23. Siggard-Anderson O, Roth M, Strickland DAP. The buffer value of plasma, erythrocyte fluid and whole blood, In: Workshop On pH and Blood Gases. Washington, DC: National Bureau of Standards; 1977.
24. Figge J, Jabor A, Kazda A, Fencl V. Anion gap and hypoalbuminemia. Crit Care Med. 1998;26:1807–1809.
25. Nicholson JP, Wolmarans MR, Park GF. The role of albumin in critical illness. Br J Anaesth. 2000;85:599–610.
26. Salem MM, Mujais SK. Gaps in the anion gap. Arch Intern Med. 1992;152:1625–1629.
27. Carvounis C, Feinfeld DA. A simple estimate of the effect of serum albumin level on the anion gap. Arch Intern Med. 2000;20:369–372.
28. Hatherill M, Waggie Z, Purves L, Reynolds L, Argent A. Correction of anion gap for albumin in order to detect occult tissue anions in shock. Arch Dis Child. 2002;87:526–529.
29. Feldman M, Soni N, Dickson B. Influence of hypoalbuminemia or hyperalbuminemia on the serum anion gap. J Lab Clin Med. 2005;146:317–320.
30. Dinh CH, Ng R, Grandinetti A, Joffe A, Chow DC. Correcting the anion gap for hypoalbuminemia does not improve detection of hyperlactatemia. Emerg Med J. 2006;23:627–629.
31. Paulson WD. Effect of acute pH change on serum anion gap. Am J Nephrol. 1996;7:357–363.
32. Lolekha PH, Vanavan N, Teerakanjana N, Chaichana Jaremkl. Reference ranges of electrolytes and anion gap on the Beckman E4A, Beckman Synchron CX5, Nova CRT, and Nova Stat Profile Ultra. Clin Chem Acta. 2001;307:87–93.
