

Hypotonic and isotonic intravenous maintenance fluids in hospitalised paediatric patients: a randomised controlled trial

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ABSTRACT

Objective To compare the changes in serum sodium and acid/base status in patients receiving hypotonic and isotonic solutions.

Design A randomised, controlled and double-blind clinical trial.

Setting Department of Paediatrics in a tertiary general hospital (Hospital Universitario Austral) in Buenos Aires, Argentina.

Patients Children between 29 days and 15 years of age who were hospitalised in the paediatric intensive care unit and general hospital between 12 January 2010 and 30 November 2016, and who required exclusively parenteral maintenance solutions for at least 24 hours.

Interventions A hypotonic solution with 77 mEq/L sodium chloride (0.45% in 5% dextrose) and isotonic solution with 150 mEq/L (0.9% in 5% dextrose) were infused for 48 hours and were labelled.

Main outcome measure The main outcome was to evaluate the incidence of hyponatraemia between patients treated with parenteral hydration with hypotonic or isotonic fluids. The secondary outcome was to estimate the incidence of metabolic acidosis induced by each of the solutions.

Results The 299 patients in the present study were randomised to groups that received the hypotonic solution (n=154) or isotonic solution (n=145). The mean serum sodium concentration measurements at 12 hours were 136.3±3.9 mEq/L and 140.1±2.3 mEq/L in the hypotonic and isotonic groups, respectively, with a hyponatraemia incidence of 8.27% (n=12) and 18.8% (n=29) (p<0.001). At 24 hours, 12.4% (n=18) of the isotonic group had developed hyponatraemia compared with 46.1% (n=71) of the hypotonic group (p<0.001). The mean serum sodium concentration measurements were 134.4±5.6 and 139.3±3.1, respectively. No patient developed hypernatraemia (serum sodium concentrations >150 mEq/L) or other adverse outcomes. The relative risk in the hypotonic group was 3.7 (95% CI 2.3 to 5.9), almost four times the risk of developing hyponatraemia than those who received isotonic fluids. There were also no significant differences between the groups with regard to the development of metabolic acidosis. Hypotonic solution, age <12 months and postoperative abdominal surgery were risk factors associated with hyponatraemia.

What is known about the subject?

- ▶ There have been numerous publications that describe the development of hyponatraemia in children who receive hypotonic fluids.
- ▶ Previous studies have observed deaths and neurological disorders associated with hyponatraemia in hospitalised children.

What this study adds?

- ▶ This is the first large study in Latin America evaluating the incidence of metabolic acidosis using hypotonic versus isotonic solutions.
- ▶ It also adds understanding about the risks of adverse effects using these solutions.

Conclusions The incidence of iatrogenic hyponatraemia was greater with the administration of hypotonic fluids compared with that of isotonic fluids. There were no significant differences in the incidence of metabolic acidosis between the groups.

INTRODUCTION

In the paediatric population, the administration of parenteral therapy to maintain homeostasis of sodium and water has always been of interest. Hyponatraemia, defined as a serum sodium level of <135 mEq/L, is a frequent disorder in hospitalised children.¹ In this age group, decreases in serum chloride and sodium concentrations of 5–6 mEq/L have been observed within the first 48 hours of hospitalisation when hypotonic solutions were administered or when other sources of free water existed.^{1,2}

Maintenance solutions must guarantee the supply of water and electrolytes in patients unable to take oral ingestion; however, there is not yet a universal consensus regarding the composition of these solutions in the



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paediatric population despite growing evidence against the use of intravenous hypotonic solutions as maintenance fluids in this population.²⁻⁵

A number of studies have assessed the incidence of hyponatraemia in children,⁶⁻¹⁰ reporting that the administration of hypotonic solutions in children increases the risk of hyponatraemia in contrast to that for isotonic solutions.^{2 7 9 11 12}

The present study could be an important evidence in Latin America, where hypotonic solutions are still frequently used for hydration of paediatric patients despite studies that support the use of isotonic solutions. The differences in the risk of hyponatraemia in both solutions have never been compared in Latin America.

This study analysed the incidence of hyponatraemia between patients treated with parenteral hydration with hypotonic or isotonic fluids. Additionally, as recent studies¹³⁻¹⁵ suggest that the use of isotonic fluids with the same proportions of sodium and chloride concentrations can induce metabolic acidosis, we measured the blood pH of all enrolled patients to estimate the incidence of metabolic acidosis using each of the solutions.

MATERIALS AND METHODS

Design

This is a randomised, controlled and double-blind clinical trial. The trial was registered in March 2009 on RENIS (Registro Nacional Argentino de Investigaciones en Salud; <https://sisa.msal.gov.ar>).

Patient and public involvement

A member of the Institutional Evaluation Committee who belongs to the community as a representative of the patients participated in the design of the study. The following issues were considered: definition of the research questions, data confidence, data access, outcome and public dissemination of the results.

Patients

This study included children between 29 days and 15 years of age who had normal serum sodium concentrations on admission, were hospitalised in the Department of Pediatrics of the Hospital Universitario Austral (Pilar, Buenos Aires, Argentina) between 12 January 2010 and 30 November 2016, and required exclusively parenteral maintenance solutions for at least 24 hours. The exclusion criteria were patients with acute and chronic heart failure; immediate postoperative cardiac surgery; acute or chronic renal failure; nephritic or nephrotic syndromes; cirrhosis and/or immediate liver transplant status; history of syndrome of inappropriate secretion of antidiuretic hormone, bronchopulmonary dysplasia or pulmonary hypertension (diagnosed by echocardiography); prescription of diuretics, converting enzyme inhibitors and calcium channel blockers; and the presence of hyponatraemia prior to the administration of the solutions.

The exclusion criteria also included the following: children who did not undergo two determinations of serum sodium concentrations, those who had received an infusion of at least one dose of diuretics, the appearance of heart or kidney failure during the course of the study, interruption of fluid therapy by the attending physician, urinary osmolarity greater than 100 mOsm/kg and a weight change >2% in the first 24 hours.

The presence of the following variables was recorded: sodium (mEq/L), mild hyponatraemia (sodium (Na) ≤ 135 mEq/L), moderate hyponatraemia (Na ≤ 130 mEq/L), hypernatraemia (Na > 150 mEq/L), blood pH, metabolic acidosis (pH < 7.30), age (months), sex, death and baseline diagnosis (table 1).

Each of the patients included in this study had three sodium measurements on admission and at 12 and 24 hours. Additionally, we analysed blood pH, electrolytes, urea and glucose on admission and at 12 and 24 hours in all patients.

Intervention

Hypotonic solution with 77 mEq/L sodium chloride (0.45% in 5% dextrose) and isotonic solution with 150 mEq/L (0.9% in 5% dextrose) were infused for 48 hours and were labelled. In both groups, the volume of liquids to be infused was calculated according to the Holliday-Segar formula.³ The solutions were administered by peripheral, central, implantable or semi-implantable venous catheters.

The hydration solutions were prepared and masked in the pharmacy by two pharmacy technicians coordinating the research. The solutions were made under laminar flow, complying with the standards of quality and biosafety present in the clinical practice guidelines and procedures of the Austral Hospital. They were stored in individual closed bags until their delivery with the respective scrambling code number.

Per-protocol analysis was performed for both groups. The randomisation was performed by a sequence of random numbers generated on a computer (Microsoft Excel 2010) in blocks of four. The randomisation code was only known to the pharmacy coordinator and was hidden from all medical and nursing staff responsible for patient care.

The researchers who carried out the recruitment, processing and subsequent analysis of the data did not participate in the care and clinical monitoring of the patients.

Statistical analysis

A sample size was calculated based on the difference in proportions for categorical data. In order to detect a difference of at least 10% between the incidences of hyponatraemia in the groups, with a power of 90% and a level of significance of 5%, a minimum of 50 patients per treatment group was required.

Continuous data were expressed as mean \pm SD or as median and IQR. According to the verification of the normality assumptions of the population, the samples were compared using Student's t-test or Wilcoxon test.

Table 1 Baseline characteristics of both groups showing a balanced sample without significant differences between the groups

| Baseline characteristics | Group with hypotonic solution n=151 | Group with isotonic solution n=143 | P value |
|--|--|---------------------------------------|---------|
| Age in months* | 21, 3–109 | 18, 2–110 | 0.13† |
| Age in months* | 21, 3–109 | 18, 2–110 | 0.13† |
| <12 | 56 | 67 | |
| 12–24 | 53 | 40 | |
| >24 | 45 | 38 | |
| Sex | 73 | 59 | 0.22‡ |
| Male | | | |
| Time at study enrolment* | 6±3.2 | 5±2.6 | 0.43† |
| Serum sodium concentrations on admission (mEq/L) | 142 (141–145) | 143 (139–146) | 0.75† |
| Potassium values (mEq/L) | 4 (3.4–4.4) | 3.9 (3.5–4.8) | 0.71† |
| Glucose values in blood (mg/dL) | 121 (109–171) | 134 (110–168) | 0.69† |
| Blood pH on admission | 7.41 (7.35–7.45) | 7.402 (7.39–7.51) | 0.67† |
| Total volume infused by weight (mL/kg/day) | 69 (45.2–90.3) | 105 (35.8–112.8) | 0.62† |

| Diagnosis | Total N=294 | Group with hypotonic solution n=151 | Group with isotonic solution n=143 | P value |
|--------------------------------------|----------------|---|--|---------|
| Asthma | 13 | 6 | 7 | 0.11‡ |
| Acute respiratory infectious disease | 109 | 53 | 56 | 0.72‡ |
| Postbone marrow transplant | 23 | 14 | 11 | 0.66‡ |
| Postsurgical | | | | |
| Neurosurgical | 21 | 10 | 11 | 0.78‡ |
| Thoracic | 25 | 11 | 14 | 0.26‡ |
| Abdominal | 34 | 19 | 12 | 0.77‡ |
| Oncological chemotherapy plan | 37 | 23 | 13 | 0.54‡ |
| Seizure disorder | 32 | 15 | 19 | 0.86‡ |

*Median, IQR (25–75).

†Wilcoxon test.

‡Pearson's χ^2 test.

Categorical data were expressed as proportions and compared using the χ^2 or Fisher's tests. A p value <0.05 was considered statistically significant. Multivariable logistic regression was conducted to predict the incidence of hyponatraemia in each of the groups by studying the influence of the other variables.

Stata V.8.0 was used for statistical analyses.

RESULTS

According to the computerised records of the Austral University Hospital, the Department of Pediatrics had a total of 12 200 admissions for hospitalisation between 1 December 2010 and 30 October 2016, including 2590 to the paediatric intensive and intermediate care units.

A total of 318 patients (figure 1) were eligible and randomised for the study, of whom 19 were excluded according to the pre-established criteria. Five did not undergo a second measurement for serum sodium concentrations, three had received at least one dose of diuretics, four had associated comorbidity, and the

parents of the remaining seven refused to sign the informed consent.

Five patients were lost to follow-up, resulting in 151 and 143 patients in the hypotonic and isotonic solution groups, respectively. The median serum sodium concentration measurements at 12 hours were 136.3 mEq/L (IQR: 133–139) and 140.1 mEq/L (IQR: 138–142) in the hypotonic and isotonic groups, respectively. The incidence of mild hyponatraemia was 18.8% (n=29) in the hypotonic group and 8.2% (n=12) in the isotonic group, with a statistically significant difference (table 2).

At 24 hours, the median serum sodium concentrations were 134.4 mEq/L (IQR: 128–139) and 139.3 mEq/L (IQR: 136–142) in the hypotonic and isotonic groups, respectively (table 2). The hypotonic group developed mild hyponatraemia in 46.1% (n=71) compared with 12.4% (n=18) of the isotonic group, and this difference was statistically significant.

There were only five patients with moderate hyponatraemia, and all of them belong to the hypotonic group.

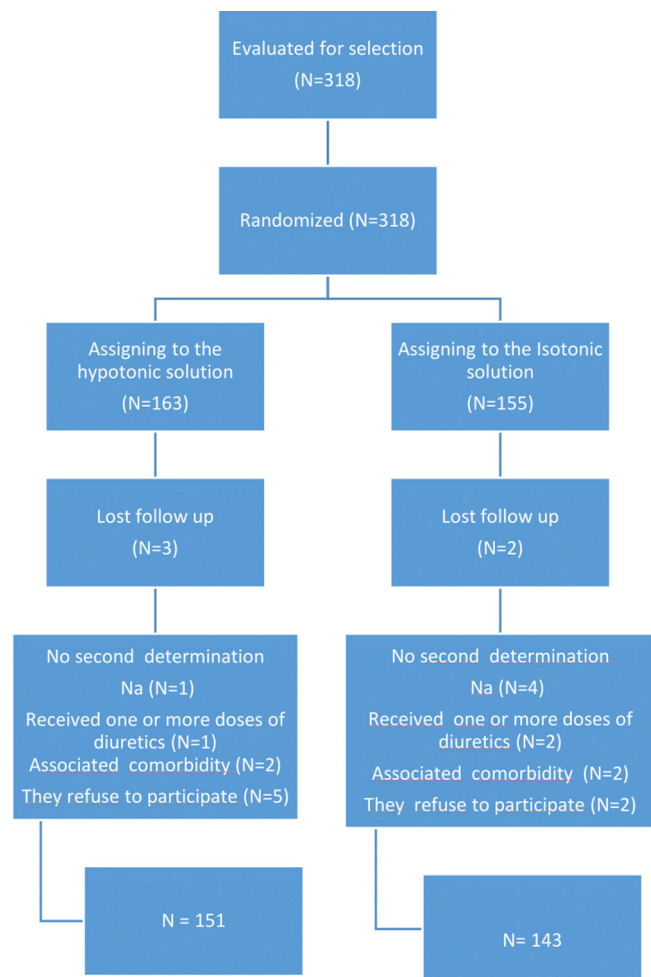


Figure 1 Flow chart of subject assignment and selection. Na, sodium.

We did not find cases of hypernatraemia in any of the groups (serum sodium concentrations >150 mEq/L) or other adverse outcomes. At 12 and 24 hours we did not

observe a statistically significant difference in the blood pH (table 2).

Comparison of the group that received hydration with the isotonic solutions in relation to the hypotonic solution group revealed a risk difference of -0.33 . The relative risk (RR) of developing hyponatraemia in the hypotonic group was 3.7 (95% CI 2.33 to 5.91); therefore, patients who received hypotonic fluids had almost four times the risk of developing hyponatraemia than those who received isotonic fluids.

There were also no significant differences between groups and 12 and 24 hours with regard to the development of metabolic acidosis (table 2). The proportions of patients with blood pH <7.30 were 17.9% (n=26) and 26.2% (n=38) in the isotonic group at 12 and 24 hours and 11.6% (n=18) and 20.7% (n=32) in the hypotonic group.

Although the restriction imposed by the inclusion and exclusion criteria could control for some confounding variables and effect modifiers, we improved on this aspect by performing multivariable logistic regression analysis.

Based on the particular behaviour of each of the variables in univariate analysis (table 3), the variables with statistical significance were selected for multivariable analysis. We observed that hypotonic solution (OR: 1.98, 95% CI 1.69 to 4.91, p=0.04), age <12 months (OR: 1.07, 95% CI 1.05 to 1.08, p=0.03) and postoperative abdominal surgery (OR: 2.13, 95% CI 1.02 to 3.03, p=0.05) were associated with hyponatraemia. The performance of the multivariable analysis was tested by a receiver operating characteristic curve (figure 2).

DISCUSSION

In this study, the use of isotonic solutions reduces the development of iatrogenic hyponatraemia in

Table 2 Results at 12 and 24 hours

| | Hypotonic group | Isotonic group | P value |
|--|------------------|------------------|---------|
| Patients with mild hyponatraemia (Na <135) at 12 hours | 29/154 (18.8%) | 12/145 (8.2%) | <0.01* |
| Patients with mild hyponatraemia (Na <135) at 24 hours | 66/154 (42.8%) | 18/145 (12.4%) | <0.01* |
| Patients with moderate hyponatraemia (Na <130) at 12 hours | 0/154 | 0/145 | |
| Patients with moderate hyponatraemia (Na <130) at 24 hours | 5/154 (3.2%) | 0/145 | |
| Patients with metabolic acidosis (pH <7.30) at 12 hours | 18/154 (11.6%) | 26/145 (17.9%) | 0.32* |
| Patients with metabolic acidosis (pH <7.30) at 24 hours | 32/154 (20.7%) | 38/145 (26.2%) | 0.71* |
| Sodium serum (mEq/L) at 12 hours† | 136.3 (133–139) | 140.1 (138–142) | 0.02‡ |
| Sodium serum (mEq/L) at 24 hours† | 134.4 (128–139) | 139.3 (136–142) | 0.03‡ |
| pH in blood at 12 hours† | 7.36 (7.31–7.42) | 7.34 (7.27–7.40) | 0.18‡ |
| pH in blood at 24 hours† | 7.39 (7.35–7.40) | 7.36 (7.29–7.39) | 0.31‡ |
| Serum bicarbonate (mmol/L) at 12 hours† | 23 (21–31) | 24 (19–28) | 0.23‡ |
| Serum bicarbonate (mmol/L) at 24 hours† | 28 (24–33) | 25 (22–31) | 0.19‡ |

Categorical variables are expressed as percentages.

* χ^2 test.

†Median, IQR (25–75).

‡Wilcoxon test.

Na, sodium.

Table 3 Univariate analysis

| Univariate analysis | Hyponatraemia | | |
|--|---------------|-------------|---------|
| | Variables | OR, 95% CI | P value |
| Age <12 months | 3.24 | 2.7 to 5.3 | <0.01 |
| Age 12–24 months | 2.13 | 0.8 to 9.3 | 0.75 |
| Age >24 months | 1.24 | 0.4 to 5.9 | 0.46 |
| Male | 1.75 | 1.0 to 2.6 | 0.02 |
| After thorax surgery | 0.52 | 0.3 to 1.7 | 0.07 |
| After neurosurgery | 2.01 | 0.7 to 4.7 | 0.06 |
| After abdominal surgery | 2.03 | 1.9 to 4.5 | 0.01 |
| Convulsive syndrome | 1.47 | 0.8 to 4.7 | 0.07 |
| Acute infectious respiratory disease | 2.34 | 1.1 to 5.2 | 0.05 |
| Asthma | 0.56 | 0.3 to 12.1 | 0.78 |
| Treatment with chemotherapy (oncological and transplanted patients of bone marrow) | 0.65 | 0.2 to 1.8 | 0.06 |

hospitalised paediatric patients receiving exclusive parenteral hydration during 24 hours compared with children who received hypotonic fluids. The RR for developing hyponatraemia was 3.17, and this is similar to other studies, including a meta-analysis⁷ involving almost 1000 children where hypotonic fluids were associated with an RR of 2.37 for developing mild hyponatraemia and 6.2 for developing moderate hyponatraemia.

There has been much discussion about the routine use of hypotonic fluids and the risk of hyponatraemia in hospitalised children.^{16–18} Although numerous clinical studies have shown differences between children treated with both solutions, they do not always reach statistical significance, perhaps due to the sample size.^{17–19} However, McNab *et al* have recently produced

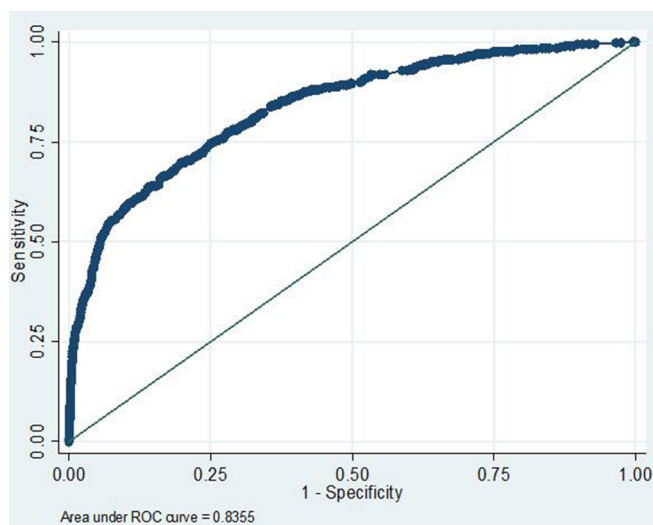


Figure 2 Area under the receiver operating characteristic (ROC) curve, 0.83 (95% CI 0.819 to 0.848).

categorical evidence showing prevention of hyponatraemia with the use of isotonic fluids.²⁰

The multivariable analysis revealed that age <12 months and postsurgical abdominal surgery were significantly associated with the development of hyponatraemia.

Young children present difficulties in maintaining a hypertonic renal medulla, which could explain the association of age <12 months with hyponatraemia. Among abdominal postoperative patients, excess interstitial fluid immediately postintervention as well as the presence of colostomies (with possible sodium loss) in some of these children could justify the association. However, both situations had an increased incidence of hyponatraemia, regardless of the infused solution.

Moritz and Ayus^{19,21} reported an incidence of death in patients with iatrogenic hyponatraemia of 8.4% (95% CI 1.85 to 15.01) and that 1 in 12 patients who acquired in-hospital hyponatraemia could die, thus showing the potential for severe consequences of the administration of hypotonic solutions. In our study, no patients died, probably due to the fact that they were mainly clinically stable patients, including those admitted to the intensive care unit who had a low risk of expected mortality, with a paediatric mortality index classification of 2 on admission. We did not observe neurological symptoms in either group based on the serum sodium concentration values, as might occur in cases of abrupt fluctuations of serum sodium or hyponatraemia with values lower than 130–125 mEq/L.

Several studies have suggested that the main factor associated with iatrogenic hyponatraemia is the continuous intravenous infusion of hypotonic solutions for more than 6 hours.^{18, 19, 21} Holliday *et al* reported that maintaining isotonic fluids for 6–12 hours in postsurgical patients restored blood volume and reduced ADH (Antidiuretic hormone) secretion; however, the authors recommend changing to hypotonic solutions after 12 hours due to the potential risk of hypernatraemia.¹⁸ The possibility of inducing hypernatraemia with isotonic fluids is very low in contrast to the high risk of generating hyponatraemia by maintaining hypotonic fluids. Numerous studies have reported a protective effect against hyponatraemia with the infusion of isotonic solutions from the time of admission and for more than 12 hours.^{17, 19, 21–23} Our study focused only on patients who received parenteral support exclusively. We did not perform evaluations after 24 hours of treatment when children are frequently fed via the enteral route, and the normal compensatory mechanisms are expected to normalise the serum sodium concentrations.

The American Academy of Pediatrics Clinical Practice Guideline: Maintenance Intravenous Fluids in Children, recently published, recommends that patients 28 days–18 years of age requiring maintenance intravenous fluids should receive isotonic solutions with appropriate potassium chloride and dextrose because

they significantly decrease the risk of developing hyponatraemia. In this paper the authors comment on the importance of analysing whether the use of 0.9% saline fluids for maintenance can increase the incidence of metabolic acidosis.²⁴

One of the contributions of our study was its analysis of the risk of developing hyperchloraemic metabolic acidosis after the infusion of isotonic solutions with the same concentrations of chloride and sodium. This adverse collateral effect has been reported in different series^{13 18 22} and clinical trials.^{14 15 23} In our study, we did not observe a statistically significant difference in the incidence of metabolic acidosis.

Finally, in our study we excluded children with renal or cardiac comorbidities; however, the potential for the development of hyponatraemia following the administration of hypotonic solutions should also be considered in these patients with difficulties in the excretion of free water. This could be a consideration for future studies.

CONCLUSIONS

The incidence of iatrogenic hyponatraemia was greater with the administration of hypotonic fluids compared with isotonic fluids. There were no significant differences in the incidence of metabolic acidosis between the groups.

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Competing interests None declared.

Patient consent for publication Obtained.

Ethics approval The study was approved by the Institutional Evaluation Committee (IEC) of the Austral University Hospital (HUA). The parents and/or guardians of the children authorised their participation in the study after having read and understood the informed consent prepared under the rules of the IEC.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available in a public, open access repository.

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