The Effects of Salt and Water Loading on Kidney Function in Healthy Undergraduates

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To cite this article:
Ehmann Peter J., Brush Christopher J., Bozzi Brittany, Dowden Robert A., Ogilvie Anna, Wisniewski Paul J., Bernard Laura P., Merrill Gary F. The Effects of Salt and Water Loading on Kidney Function in Healthy Undergraduates. Advances in Applied Physiology. Vol. 4, No. 2, 2019, pp. 11-18. doi: 10.11648/j.aap.20190402.11

Received: October 8, 2019; Accepted: November 6, 2019; Published: November 17, 2019

Abstract: Rutgers University juniors and seniors enrolled in Systems Physiology Laboratory (01:146:356) Spring, 2019. One-hundred forty-five students volunteered to test the effects of salt and water loading on kidney function. Students were investigated beginning at 8:40 a.m., 1:40 p.m., and 6:40 p.m. They were approximately evenly distributed between male and female genders and were of multiple cultural/ethnic backgrounds. Upon entering the laboratory baseline data were collected. Approximately half of the volunteers then consumed 200 mL of a 2.0% NaCl solution (n=77). The others directly consumed 1.0 L of tap water (n=68). Upon entering the laboratory baseline data were collected. The others directly consumed 1.0 L of tap water. Kidney function was monitored each 30 minutes for the next 90 minutes. Under baseline conditions, urine osmolality was elevated in all students (≥ 280 mOsm/kg) but was significantly (P<0.05) greater in the 8:40 a.m. group than in either of the p.m. groups. Urine specific gravity and sodium excretion were also elevated in the 8:40 a.m. group compared to the others. Urine flow rate was least in the 8:40 group and greatest in the 1:40 p.m. group. We conclude that in these students renal function was influenced by a diurnal pattern. We also conclude that without compelling incentives (e.g. financial, grade-influencing) it was virtually impossible to get students to comply with pre-experimental instructions (e.g. no salt-laden meals after 6:00 p.m. before the day of experimentation).

Keywords: Dehydration, Salt Loading, Urine Osmolality, Water Balance, Water Loading, Sodium Excretion, Urine Flow

1. Introduction

Physiological kidney function is one of the sine qua non of good health [1, 2]. In medicine clinicians have been using the estimated glomerular filtration rate (eGFR) for the past 15-20 years as an indicator of renal function [3-5]. There are many other estimates of kidney function as well, including volume of urine excreted and frequency of urination. The latter two, coupled with measures of urine osmolality, urine specific gravity, and urinary sodium excretion, are excellent and standard measures of body water content and therefore of the degree of hydration/dehydration [6-8]. We use these measures routinely in a laboratory experiment designed to teach kidney function to juniors and seniors planning careers in medicine and related health sciences professions (e.g. physician assistants, physical/occupational therapy, etc.).

This is not only a useful experiment to help students understand kidney function, but it also helps demonstrate important physiological principles related to body water regulation. The physiology and truth about body water are long-standing discussions fueled in more recent decades by the bottled water industry and the ubiquitous commercialization of exercise gyms and gym memberships. Both are big businesses funded (and founded) by the misinformed, uninformed and under-educated in all things physiology and body function. In this article we hope to correct some of the misinformation that continues to circulate about kidney function, body water, and physiological health.
2. Methods

Experimental subjects were students enrolled in Systems Physiology Laboratory (01:146:357) at Rutgers University, New Brunswick/Piscataway, New Jersey, in the Spring Semester, 2019. Upon arriving in the lab subjects (n=145) were divided into two, salt- and water-consuming groups. As they arrived in the laboratory all students were sent to the restrooms to empty their bladders and to obtain urine samples. After this students in the salt group (n=77) quickly consumed 200 mL of a 2.0% salt solution. Students in the water group (n=68) drank 1.0 L of tap water as quickly as possible. Thereafter, timers were set and each student subsequently collected additional urine samples at 30, 60 and 90 minutes post-consumption of their respective drinks. Measurements of urine flow rate, specific gravity, NaCl concentration and osmolality were made/calculated on each sample. Students handled only their own urine, and this and other experiments in this experimental physiology laboratory course have been reviewed and approved by Rutgers University.

Specimen containers were graduated and full at 100-120 ml. If the volume of urine voided exceeded 100-120 ml, students dumped the first volume in the toilet and continued collecting the remaining volume. To estimate their rates of urine flow (mL/min), students noted the volume markings on specimen cups. They then divided the volume (mL) by the time elapsed (min) since the last urine collection.

Specific gravity---Specific gravities of each urine sample were estimated using refractometry (refractometers, ExTech Instruments, model RF20, China). With the prism cover slip open and the prism cleaned, students calibrated their refractometers by carefully dispensing 2-3 drops of distilled, deionized water (d_{H2O}) onto the prism. They moved the cover slip to the closed position and waited several seconds for the fluid to be evenly distributed across the prism. Holding the refractometer to an eye and parallel to the floor, they used the adjustable screw to set the refractive index scale to 1.000. The prism was once again cleaned, and students carefully dispensed 2-3 drops of urine onto the prism. They moved the cover slip to the closed position and waited several seconds for the urine sample to be evenly distributed across the prism. They then observed and recorded the specific gravity for each sample.

Sodium chloride concentration (mg/mL)---Using micropipettes students transferred approximately 500 µL of urine into clean test tubes and then added 100 µL potassium chromate. To this they added 100 µL aliquots of a 2.9% silver nitrate solution (AgNO₃), incrementally, while gently shaking the sample as each aliquot was added, until the yellow-colored urine turned a brown hue. They recorded the total volume of silver nitrate needed to make the colorimetric change. Each 100 µL of 2.9% silver nitrate added represented approximately 0.15 mg NaCl in 500 µL (i.e., 100 µL of AgNO₃ = 0.3 mg NaCl/mL). They calculated the NaCl concentration (mg/mL) in the urine as: [NaCl] = [(µL AgNO₃) X (0.3mg NaCl/ml urine)] / [500 µL].

Osmolality---Osmolality---Under baseline conditions and at each 30-minute timed interval graduate student teaching assistants (TAs) received 20 µl urine samples from experimental subjects and analyzed osmolalities (mOsm/L, model 2020 osmometer, Advanced Instruments, Norwood, MA). Osmometers were previously calibrated using standard sodium chloride solutions of 50, 280, 750 and 1200 mOsm/L.

Between each 30-minute collection/calculation period, TAs led 10-minute discussions on the physiology of kidney function (power point mini-lectures). Students were asked to not consume excess salt and/or water during the 24 hours prior to initiation of the experiment.

Statistics

As data were collected they were recorded on a white board so all participants could see and make records of them. No records were kept that would allow any participant to be identified (data were confidential). The grouped data were later used by students to complete written laboratory reports.

Baseline data (3 levels; 8:40 a.m., 1:40 p.m., 6:40 p.m.) were analyzed with a one-way ANOVA. Pairwise comparisons were conducted using Tukey’s HSD test for renal variables with significant ANOVA results. Effects over time (4 levels; baseline, 30min, 60min, 90min) for each experimental group (2 levels; salt, water) were analyze with a repeated-measures ANOVA. Pairwise comparisons were conducted using Tukey’s HSD test for renal variables with a significant Time x Group interaction. All data are reported as mean plus or minus one standard deviation. Statistically significant results were identified at P<0.05.

3. Results

3.1. Baseline Data

Results of the one-way ANOVA and descriptive statistics are presented in Table 1. Pairwise comparisons for each variable with significant ANOVA results are presented in Table 2. Flow rate was significantly lower at 8:40 a.m. compared to the p.m. times. Urine specific gravity was only significantly different between 8:40 a.m. and 6:40 p.m. times, with the former being greater. No differences were observed over time in urine NaCl concentration at baseline. NaCl excretion rate was significantly different between 8:40 a.m. and 1:40 p.m. times, with the former being lower. Urine osmolality was significantly greater at 8:40 a.m. than both the 1:40 p.m. and 6:40 p.m. time points.

3.2. Group Differences over Time

Figures 1-5 display group differences in each variable over time (bounded by 95% confidence interval). Importantly, no differences were observed between groups for any variable at baseline (indicating proper random group assignment).

Urine flow rate---Under baseline conditions and on the average, urine flow rates in both groups were about 1.1 ml/min. These rose significantly in the water group to more than 3.6 and 6.5 ml/min at 30 and 90 minutes, respectively. At 90 minutes urine flow in the salt group declined
significantly to about 0.8 ml/min (P<0.05, Figure 1).

Figure 1. Influence of tap water vs a 2.0% solution of NaCl on the rate of urine flow.

Urine specific gravity---On the average and relative to baseline conditions, the specific gravity of urine in the water group declined significantly (P<0.05) at 30, 60, and 90 minutes post-consumption. Conversely, there was a trend (at 60 and 90 minutes post-consumption) for specific gravity to increase in the group that consumed 2.0% NaCl which did reach statistical significance relative to baseline (P<0.05, Figure 2).

Figure 2. Influence of tap water vs a 2.0% solution of NaCl on urine specific gravity.
Urine sodium chloride concentrations---Urinary sodium chloride concentrations increased significantly in the salt group and decreased significantly in the tap water group (Figure 3).

For example, under baseline conditions, salt concentrations in the urine were about 2.5 mg/dl in both groups. In the salt group this rose to 2.7, 3.5 and 3.9 mg/dl, on the average at 30, 60, and 90 minutes, respectively, post-consumption. Conversely, corresponding figures in the water group at 30, 60, and 90 minutes were, 1.7, 0.7 and 0.6 mg/dl.

Urinary sodium chloride excretion---Relative to baseline conditions, the rates of excretion of sodium chloride in both groups of subjects were elevated throughout the experiment (i.e. 30-90 minutes following consumption of tap water and a 2.0% solution of NaCl). At 60 minutes the rates were significantly lower (P<0.05) in the salt group than in the water group (Figure 4).

These rates were, however, still elevated above baseline control values at 90 minutes in both groups.
Urban osmolality---On the average urine osmolality was close to 560 mOsm/kg under baseline conditions in both groups of students. It rose steadily but modestly to about 800 mOsm/kg in the salt group at 90 minutes, and declined significantly to about <100 mOsm/kg at the same time in the water group. In the salt group urine osmolality was only statistically significant at 90 minutes (relative to baseline, Figure 5), whereas in the water group it was significantly different than baseline at 60 and 90 minutes.

4. Discussion

The kidneys are the chief regulators of the internal environment of the body (internal milieu as opposed to the external milieu). They help control the concentration of ions, water, and pH in the several body water compartments (intracellular space, interstitial space, intravascular space). In addition, kidneys eliminate waste products of metabolism. The nearly one million nephrons in each kidney contain two main structures: the glomeruli (glomerulus, singular) and the renal tubules. As blood passes through the kidneys it is filtered by the glomeruli at a rate of approximately 120 ml/min, (in healthy adult humans; known as the glomerular filtration rate, GFR).

Chemically, the glomerular filtrate is similar to blood plasma except that large molecules having molecular weights of more than 70 kD are excluded (e.g., plasma proteins). As this glomerular filtrate (also known as an ultrafiltrate) passes along the proximal and distal tubules most of the water is reabsorbed, and many other inorganic and some organic chemicals are actively or passively reabsorbed back into the bloodstream. Toxic byproducts of metabolism, as well as substances in excess, are retained in the ultrafiltrate or are secreted into the ultrafiltrate and finally excreted in the urine (urine flow rate is approximately 0.3 – 3.0 ml/min; diurnal variation). Thus the final composition of urine is quite different from that of the glomerular ultrafiltrate and reflects the integrity of kidney function and changes in blood composition.

An analysis of urine can yield valuable information about the health of the kidney and of the body in general. Various diseases are characterized by altered metabolism, which causes unwanted byproducts of metabolism to appear in the urine. For example, phenyl pyruvic acid appears in the urine in phenylketonuria (PKU), a disease resulting in mental retardation. In diabetes mellitus, deficient production of insulin by the pancreas results in the appearance of glucose in the urine (glucosuria). Thus, the volume of urine produced, its rate of flow, urine osmolality and specific gravity, and related qualities give information on the state of hydration or dehydration of the body.

Urine flow rate---Under baseline conditions and on the average, urine flow rates in both groups were about 1.0 ml/min. These rose significantly in the water group to more than 6 ml/min and declined markedly in the salt group to about 0.5 ml/min. This 12-fold difference in urine flow is best explained by the physiology of osmoreceptors in the anterior hypothalamus, the production and release of arginine vasopressin (AVP or antidiuretic hormone, ADH) from the posterior pituitary, and AVPs actions on distal components of the renal tubular system (9-13). In the water group the aim of these interactive physiological mechanisms was to eliminate excess water. In the salt group, oppositely-directed actions of the same organs/tissues were directed at conserving water by

![Figure 5. Differences in urine osmolality in subjects after consuming 1.0 liters of tap water and 200 mL of a 2.0% NaCl solution.](image-url)
reducing its excretion in the urine.

Arginine vasopressin, so named because its vasopressor (vascular) effects were identified before its renal effects were, enhances the production and subcellular movement of proteins known as ‘aquaporins’ [13, 14-16]. These protein water channels are inserted into the apical membranes of tubular epithelial cells where they enhance the ability of tubules to reabsorb water, thus conserving it. Although we did not measure AVP, aquaporins, or related compounds in this experiment, our speculation is reasonable because it is based on irrefutable, supporting evidence [17-21].

Urine osmolality—Significantly reduced urine flow rates at 8:40 a.m. were not expected. Based on diurnal patterns in humans, urine flow rates are lowest in the late p.m. to early a.m. hours, and highest in the late a.m. to early p.m. hours [22, 23]. We suspect the reduced flow was caused by hyperosmolality as baseline data suggest (e.g. see Figure 5, baseline values in both water and salt groups). University students, as a rule, lead unregulated dietary lifestyles. This includes eating at any time of the day, e.g. ordering pizza with extra sauce, pepperoni, and cheese after midnight (personal communications). We believe it is their generally-poor dietary habits that best explain high urine osmolality at 8:40 a.m. in university students.

Hypothalamic osmoreceptors would have been stimulated by elevated plasma osmolality (pizza, sushi, salt) leading to increased release of AVP from the neurohypophysis. The subsequent increased insertion of aquaporins into apical membranes of renal tubular cells would have promoted tubular water conservation leading to increased urine osmolality. Support for this reasoning can be confirmed by carefully reviewing the data in Figures 1-4.

Water and salt balance---The distribution and regulation of body water is tightly coupled to fluid osmolality. Likewise, the regulation of osmolality is physiologically determined by the regulation of body water [24-27]. For example, intracellular fluid volume is determined, in part, by the osmolality of extracellular fluid, including plasma osmolality. As extracellular fluid sodium concentration increases post-consumption of a salt-loaded meal, so too will extracellular osmolality increase. Among other physiological events, this will lead to withdrawal of water from the intracellular compartment (by osmosis), and to stimulation of hypothalamic osmoreceptors. Thirst mechanisms will be simultaneously activated driving the person to consume water.

During a 24-hour period following salt loading, salt concentration in the intravascular compartment will begin to decline (with renal water conservation and corresponding thirst-driven water consumption). Decreasing salt concentrations in the extracellular compartment will be accompanied by a decrease in the osmolality of the hypothalamic, interstitial spaces and to a reduction in the release of neurohypophysyal AVP. With reduced AVP, epithelial cells in the distal renal tubules will recycle apical aquaporins and water reabsorption will decrease.

Physiological homeostasis for the distribution of body water and salt in the student will be re-achieved until the next craving pizza/sushi in the late p.m./early a.m. hours.

5. Conclusions

Students consuming a 2% solution of NaCl had lower rates of urine flow and higher urine osmolalities than those consuming tap water. Principles of renal water conservation and renal excretion of excess salt were confirmed in this routine experiment on body water and salt homeostasis. Disappointingly, most students had hyperosmotic urine even during the baseline period. This suggests the unwillingness of students to comply with experimental instructions, e.g. please do not consume excess salt and/or water during the 24 hours prior to this classic renal experiment.

Appendix

Table 1. One-way ANOVA results and descriptive statistics (M ±SD) analyzing effect of time of day on renal variables.

| Flow Rate * | n | Mean | SD | df₁ | df₂ | F    | p   |
|-------------|---|------|----|-----|-----|------|-----|
| 8:40am      | 51| 0.62 | 0.61| 2   | 142 | 11.07| <.001|
| 1:40pm      | 39| 1.67 | 1.53|     |     |      |     |
| 6:40pm      | 55| 1.24 | 1.00| 2   | 131 | 4.05 | .02  |
| Specific Gravity * | 8:40am | 51 | 1.033 | 0.013 | 2 | 142 | 1.59 | .21  |
| 1:40pm      | 28 | 1.025 | 0.016 |     |     |      |     |
| 6:40pm      | 55 | 1.025 | 0.016 | 2   | 142 | 4.45 | .01  |
| [NaCl]      | 8:40am | 51 | 2.78 | 1.31 | 2   | 142 | 4.45 | .01  |
| 1:40pm      | 39 | 2.36 | 1.23 |     |     |      |     |
| 6:40pm      | 55 | 2.38 | 1.41 | 2   | 142 | 14.76| <.001|
| NaCl ex. Rate * | 8:40am | 51 | 1.44 | 1.12 | 2   | 88  | 14.76| <.001|
| 1:40pm      | 39 | 3.17 | 2.89 |     |     |      |     |
| 6:40pm      | 55 | 2.66 | 3.84 | 2   | 88  |      |     |
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Mean  SD  df1  df2  F  p  df
8:40am 38  738  279
1:40pm 22  481  230
6:40pm 31  412  260

Note: Sample size is much lower for Osmolality because graduate student TA did have enough time to collect all students’ samples. Tukey HSD pairwise comparisons for significant results are presented in Table 2.

Table 2. Tukey HSD pairwise comparisons for variables with significant ANOVA results.

|       | difference | 95% CI lower | 95% CI upper | p adj |
|-------|------------|--------------|--------------|-------|
| Flow Rate | 2 – 1 * | 1.048 | 0.511 | 1.586 | <.001 |
|        | 3 – 1 * | 0.618 | 0.127 | 1.109 | <.01 |
|        | 3 – 2 | -0.431 | -0.960 | 0.098 | .13 |
| Specific Gravity | 2 – 1 | -0.007 | -0.016 | 0.001 | .09 |
|        | 3 – 1 * | -0.008 | -0.144 | -0.001 | .03 |
|        | 3 – 2 | -0.001 | -0.008 | 0.008 | .99 |
| NaCl exr. rate | 2 – 1 * | 1.728 | 0.279 | 3.178 | .01 |
|        | 3 – 1 | 1.223 | -0.102 | 2.547 | .08 |
|        | 3 – 2 | -0.506 | -1.933 | 0.920 | .68 |
| Osmolality | 2 – 1 * | -257 | -424 | -90 | <.01 |
|        | 3 – 1 * | -326 | -477 | -175 | <.001 |
|        | 3 – 2 | -69 | -242 | 105 | .61 |

Note: 1 = 8:40am, 2 = 1:40pm, 3 = 6:40pm. Comparisons are significant if zero does not fall within the 95% CI. Adjusted p-values are shown.

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