Proceedings

Optimal Hydration Biomarkers: Consideration of Daily Activities

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Introduction

Water intake and hydration status have recently gained attention as one of the many, and potentially manipulable, factors associated with disease development and wellbeing [1–7]. Researchers have explored a variety of physiological dysfunctions implicated with chronic low water intake and hypohydration, but particular interest resides in the study of obesity and related disease states due to overwhelming prevalence. However, the current lack of a hydration assessment gold standard greatly impedes our attempts to link water intake and negative health outcomes as well as to make public dietary guidelines.

Current hydration biomarkers include blood, saliva, and urine sampling subjected to a variety of quantification methods (osmolality, specific gravity, volume, etc.). Recent work from our laboratory [8] contributed to the ongoing quest to identify optimal selection of hydration biomarkers, but through consideration of the context under which body water loss occurs. We present two scenarios when selecting optimal hydration biomarkers: i) active versus passive body water loss, and ii) single versus serial measurements.

Methodology

We examined 23 healthy, recreationally active males (age = 22 ± 3 years; mass = 77.3 ± 12.8 kg; height = 179.9 ± 8.8 cm; body fat = 10.6 ± 4.5%). Participants completed one familiarization trial, recorded their complete dietary intake for 3 consecutive days, and underwent

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two experimental trials. Individual water intake was prescribed for the day prior to experimental trials as an average intake from their 3-day dietary records. Furthermore, participants replicated all food consumption the day prior to experimental trials.

Two experimental trials were conducted in a hot environment (about 36° C, approximately 50% relative humidity) to examine: i) passive dehydration (PAS), where participants simply sat in the hot environment for 5 h, and ii) active dehydration (ACT), where participants cycled in the hot environment at a moderate intensity (about 68% of age-predicted maximal heart rate) for 5 h. Participants did not consume foods or water during these trials. Upon each percent body mass loss, participants provided blood, saliva, and urine samples. Researchers measured blood serum osmolality (S_{osm}), salivary osmolality (V_{osm}), urine osmolality (U_{osm}), volume (U_{vol}), and specific gravity (U_{sg}).

Passive versus Active Dehydration

We uncovered that the context under which body water losses occur does influence the ability of hydration biomarkers to assess dehydration. Largely, we found during PAS that progressive and large changes were observed more so in urinary compared to serum and salivary variables. Conversely, we found during ACT that serum and salivary markers responded more precisely to body water losses than did urinary variables (fig. 1). These findings are likely explained through autonomic regulation and hemoconcentration during heat and exercise exposure [9], which concentrates the extracellular compartment, primary saliva [10], and diverts cardiac output away from the kidneys [11–13]. In support of these findings, we were able to compare mean differences between PAS and ACT trials at 1% body mass loss, and found that even mild dehydration elicited statistically and physiologically meaningful differences between the two methods of body water loss (table 1). In essence, these findings indicate that exercise exaggerated changes in S_{osm} and V_{osm}, and that urinary markers best represented water losses during PAS.

Single versus Serial Measurement

Statistical tools for measuring diagnostic accuracy and reference change values (RCV) permitted evaluation of hydration biomarker utility with availability of single versus serial measurements, respectively.

Diagnostic Accuracy

In order to evaluate hydration biomarker utility with a single measurement, we operationally defined a dehydrated state as 2% body mass loss, as this degree of body water loss typically provokes physiological and psychological dysfunction [5, 14–17]. Diagnostic accuracy statistics included: i) receiver operating characteristic (ROC) to define criterion values for each diagnostic test, ii) area under the ROC curve (AUC) to summarize diagnostic accuracy, and iii) sensitivity, specificity, and positive likelihood ratio (+LR) to assist in clinical decision making. The best combination of sensitivity (the probability of a true positive) and specificity (the probability of a true negative) allows the practitioner to use the ROC-derived criterion value with confidence to diagnose dehydration with the respective hydration biomarker. Great practicality exists in the +LR, as high ratios indicate that true positives are much more likely than false positives.

Diagnostic accuracy was conducted for ACT only, as PAS resulted in an insufficient sample size greater than 2% body mass loss. We found that V_{osm} and S_{osm} most accurately assessed

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Fig. 1. Hydration biomarker change (Δ means ± SE) with body mass loss via PAS (left) and ACT (right): **a** serum osmolality, **b** salivary osmolality, **c** urine osmolality, **d** urine volume, and **e** urine specific gravity. Significant differences (p < 0.05) indicated with * compared to 0% mass change, † compared to –1% mass change, ‡ compared to –2% mass change, and § compared to –3% mass change.
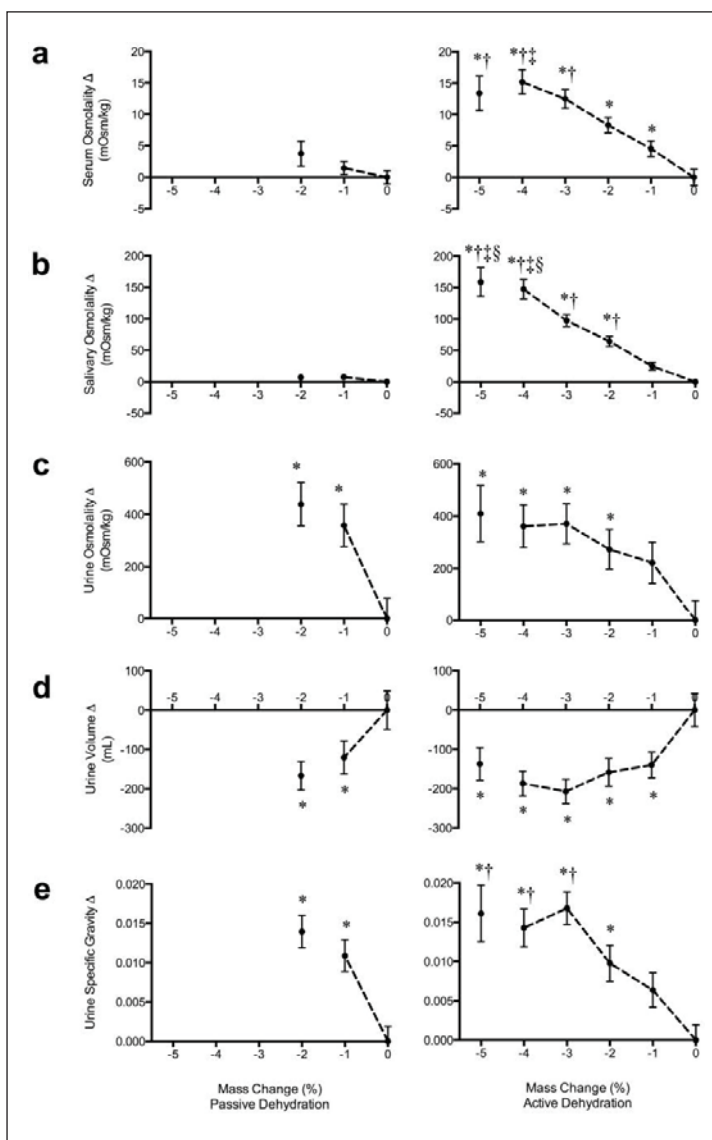
Table 1. Hydration indices at 0% (pre-trial) and –1% body mass change during PAS and ACT

Hydration biomarker	0% mass change		–1% mass change		n	p
	PAS	ACT	PAS	ACT		
Serum osmolality, mOsm·kg⁻¹	295 ± 4	296 ± 4	296 ± 4	301 ± 4	22	<0.01ᵃ
Salivary osmolality, mOsm·kg⁻¹	68 ± 14	64 ± 9	74 ± 14	90 ± 24	21	<0.01ᵃ
Urine osmolality, mOsm·kg⁻¹	466 ± 267	439 ± 252	895 ± 207	661 ± 192	11	0.01ᵃ
Urine volume, ml	266.8 ± 165.4	231.0 ± 130.1	106.4 ± 79.4	91.2 ± 46.4	11	0.59
Urine specific gravity	1.011 ± 0.006	1.012 ± 0.006	1.023 ± 0.006	1.018 ± 0.006	11	0.06

[a]Indicates differences between PAS and ACT at –1% body mass change (p < 0.05).
PAS = Passive dehydration; ACT = active dehydration.
Variation in sample size (n) at –1% body mass change attributable to number of paired samples (sample provided by subject for both PAS and ACT)

Table 2. Receiver operating characteristic curve analysis of hydration biomarkers during active dehydration when only a single measurement is available

Hydration biomarker	AUC	Estimate SE	Criterion value	Sensitivity, %	Specificity, %
Serum osmolality, mOsm·kg⁻¹	0.91[a]	0.03	303	83	83
Salivary osmolality, mOsm·kg⁻¹	0.94[a]	0.03	108	86	91
Urine osmolality, mOsm·kg⁻¹	0.80[a,b]	0.05	631	80	65
Urine volume, ml	0.87[a]	0.04	90	79	79
Urine specific gravity	0.89[a]	0.04	1.020	81	81

[a]Denotes that AUC was significantly different from change.
[b]Denotes that AUC was significantly different from salivary osmolality ($p < 0.05$); a trend was observed between serum and urine osmolality ($p = 0.06$).
AUC = Area under curve; SE = standard error.

Table 3. Positive likelihood ratios (+LR) and corresponding values for hydration biomarkers during active dehydration when only a single measurement is available

Hydration biomarker	+LR	Measured value
Serum osmolality	11.73	306 mOsm·kg⁻¹
	4.88	303 mOsm·kg⁻¹
	1.82	298 mOsm·kg⁻¹
Salivary osmolality	10.02	116 mOsm·kg⁻¹
	9.65	108 mOsm·kg⁻¹
	1.92	73 mOsm·kg⁻¹
Urine osmolality	8.38	1,146 mOsm·kg⁻¹
	2.29	631 mOsm·kg⁻¹
	1.97	582 mOsm·kg⁻¹
Urine volume	10.53	31 ml
	3.70	90 ml
	1.99	143.2 ml
Urine specific gravity	10.50	1.024
	4.22	1.020
	1.81	1.014

dehydration (table 2), which supports our observation that these variables responded more drastically to body water loss than urinary variables during ACT. The +LR for S_{osm} (table 3) tells the practitioner that, e.g., a value of 303 mOsm·kg⁻¹ occurs 4.88 times more often in a dehydrated than sufficiently hydrated individual who has been exercising.

Reference Change Values

RCV statistics evaluate the utility of a test with serial measurement. This requires preliminary measures of analytical and biological variation, which includes coefficients of variation for the analytical device (CV_A), and interindividual (CV_I) and intraindividual (CV_G) variation. The derived RCV (table 4) in this investigation indicates a meaningful change (expressed as a percentage or unit) in the hydration biomarker indicative of a dehydrated state. Subsequent analyses included the index of individuality (II) which indicates usefulness for population-based reference intervals, where a value greater than 1.4 signifies high usefulness and less than 0.6 suggests low usefulness. The index of heterogeneity (IH) assists in validating the RCV, which, in our investigation, was an IH greater than 2.

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Table 4. Analytical and biological coefficients of variation and indices of variability for hydration biomarkers when serial measurements are available

Hydration biomarker	CV_A[a]	CV_I[b]	CV_G[c]	II[d]	IH[e]	RCV, %[f]	RCV, unit[f]	Decision level
Serum osmolality, mOsm·kg⁻¹	0.3	0.9	0.9	0.96	0.96	2.12	6	302 ± 4
Salivary osmolality, mOsm·kg⁻¹	2.5	10.1	18.4	0.56	7.34	24.21	16	84 ± 16
Urine osmolality, mOsm·kg⁻¹	0.4	23.9	48.2	0.49	16.87	55.67	255	708 ± 257
Urine volume, ml	negligible	41.1	15.0	2.74	29.08	−95.96	−242	9 ± 150
Urine specific gravity	negligible	0.3	0.3	0.79	0.20	0.65	0.007	1.018 ± 0.006
Body mass, % loss	negligible	0.5	10.9	0.05	0.36	−1.20	−1.20	−1.2 ± 0.2

[a]CV_A = Coefficient of variation for the analytical device. When possible, CV_A was measured with standard solutions specific to the measurement range; otherwise, actual samples were used. All values are expressed as a percentage.

[b]CV_I = Intraindividual coefficient of variation.

[c]CV_G = Interindividual coefficient of variation. CV_I and CV_G measured with all pre-trial samples from passive dehydration and active dehydration trials and expressed as percentages.

[d]II = Index of individuality.

[e]IH = Index of heterogeneity.

[f]RCV = Reference change value expressed as a percentage, or specific to the unit of measurement.

Due to the nature of RCV statistics, the values reported are irrespective of the method through which body water loss occurred (active vs. passive). We found that only three hydration biomarkers demonstrated valid RCV (IH > 2): S_{osm}, U_{sg}, and body mass loss. For example, a change in serial U_{sg} measurement of 0.007 would indicate a dehydrated state according to our findings (table 4). Furthermore, only U_{vol} had a high enough II considered useful for population-based reference intervals, while it did not result in a valid RCV. This implies good utility for interpretation of hydration state among healthy populations based on typical variation, but not for serial measurement with acute dehydration, respectively.

Conclusion

This investigation elucidated the need to consider previous or concurrent activities, and the frequency of sample measurement when selecting hydration biomarkers. Largely, we found that S_{osm} and V_{osm} during ACT in the heat, and urinary markers during PAS in the heat, were the most suitable biomarkers given the context or means of dehydration. When only single measurements are available during ACT, S_{osm} and V_{osm} established the greatest efficacy. Regardless of the method of body water loss, S_{osm}, U_{sg}, and body mass loss resulted as the most valuable serial measurement dehydration indices. In essence, the context surrounding body water loss should dictate selection of hydration biomarkers to appropriately identify hydration status before research examining the influence of chronic water intake on health outcomes progresses.

Disclosure Statement

The authors declared no conflict of interest.

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