Beverage consumption, appetite, and energy intake: what did you expect?1–3

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ABSTRACT

Background: Beverage consumption is implicated in the overweight/obesity epidemic through the weaker energy compensation response it elicits compared with solid food forms. However, plausible mechanisms are not documented.

Objective: This study assessed the cognitive and sensory contributions of differential postigestive responses to energy- and macronutrient-matched liquid (in beverage form) and solid food forms and identifies physiologic processes that may account for them.

Design: Fifty-two healthy adults [mean ± SD age: 24.7 ± 5.5 y; BMI (in kg/m²): 26.3 ± 6.3] completed this randomized, 4-arm crossover study. Participants consumed oral liquid and solid preloads that they perceived, through cognitive manipulation, to be liquid or solid in their stomach (ie, oral liquid/perceived gastric liquid, oral liquid/perceived gastric solid, oral solid/perceived gastric liquid, or oral solid/perceived gastric solid). However, all preloads were designed to present a liquid gastric challenge. Appetite, gastric-emptying and orocecal transit times, and selected endocrine responses were monitored for the following 4 h; total energy intake was also recorded.

Results: Oral-liquid and perceived gastric-liquid preloads elicited greater postprandial hunger and lower fullness sensations, more rapid gastric-emptying and orocecal transit times, attenuated insulin and glucagon-like peptide 1 release, and lower ghrelin suppression than did responses after oral-solid and perceived gastric-solid treatments (all P < 0.05). Faster gastric-emptying times were significantly associated with greater energy intake after consumption of perceived gastric-liquid preloads (P < 0.05). Energy intake was greater on days when perceived gastric-liquid preloads were consumed than when perceived gastric solids were consumed (2311 ± 95 compared with 1897 ± 72 kcal, P = 0.007).

Conclusions: These data document sensory and cognitive effects of food form on ingestive behavior and identify physical and endocrine variables that may account for the low satiety value of beverages. They are consistent with findings that clear, energy-yielding beverages evoke limited compensatory dietary responses (ie, failure to adjust intake at subsequent eating occasions for energy supplied by the beverages) in comparison to solid food forms (4, 5). Whereas extensive available data have clearly focused concern on sweetened beverages, evidence that beverages containing different energy sources elicit weak dietary compensation (6–9) suggests that the food form (ie, liquid in beverage form compared with semisolid or solid physical state), and not the macronutrient or energy source (ie, carbohydrate, fat, or protein), is likely responsible for the association between energy-yielding beverage consumption and positive energy balance (10).

Contrary, a small body of evidence fails to support the link between energy-yielding beverages and weak appetitive or dietary responses (11, 12). Discrepant findings between studies could be attributable to variations in study design, most notably failure to isolate effects of food form on postprandial responses (11, 13, 14). Comparisons of dissimilar foods and beverages are confounded by expectations and properties such as palatability and nutrient sources. There is a need for direct comparison between responses for beverages and solids with the use of appropriately designed test loads (ie, equally palatable, isocaloric, macronutrient-matched). In addition, mechanistic explanations of why food forms elicit differential regulatory responses are lacking. Beverages require less oral processing, have more rapid gastric-emptying and orocecal transit times (15, 16), and evoke lower expected satiation values (17) (ie, the degree to which an individual expects a particular food to be satiating) (18). Short-term feeding studies show that cognitive manipulations (eg, time, energy content, food labeling, portion size) significantly influence appetitive ratings and subsequent energy intake (19–22). Indeed, the perceived energy content of a food may better predict self-reported appetitive

INTRODUCTION

“We cannot entertain a doubt that every change in our sensations and ideas must be accompanied by some corresponding change in the organic matter of the body.” —Sir Humphry Davy

The rise in obesity and overweight closely parallels the increase in energy-yielding beverage consumption over the past 3 decades (1–3). Children and adults now consume ~400 kcal/d of energy-yielding beverages, accounting for ~20–25% of daily energy intake (3). The contribution of energy-yielding beverages to the promotion of positive energy balance and weight gain remains controversial. However, most studies reveal that beverages, particularly clear varieties, hold weak satiety properties and evoke limited compensatory dietary responses (ie, failure to adjust intake at subsequent eating occasions for energy supplied by the beverages) in comparison to solid food forms (4, 5). Whereas extensive available data have clearly focused concern on sweetened beverages, evidence that beverages containing different energy sources elicit weak dietary compensation (6–9) suggests that the food form (ie, liquid in beverage form compared with semisolid or solid physical state), and not the macronutrient or energy source (ie, carbohydrate, fat, or protein), is likely responsible for the association between energy-yielding beverage consumption and positive energy balance (10).

Conversely, a small body of evidence fails to support the link between energy-yielding beverages and weak appetitive or dietary responses (11, 12). Discrepant findings between studies could be attributable to variations in study design, most notably failure to isolate effects of food form on postprandial responses (11, 13, 14). Comparisons of dissimilar foods and beverages are confounded by expectations and properties such as palatability and nutrient sources. There is a need for direct comparison between responses for beverages and solids with the use of appropriately designed test loads (ie, equally palatable, isocaloric, macronutrient-matched). In addition, mechanistic explanations of why food forms elicit differential regulatory responses are lacking. Beverages require less oral processing, have more rapid gastric-emptying and orocecal transit times (15, 16), and evoke lower expected satiation values (17) (ie, the degree to which an individual expects a particular food to be satiating) (18). Short-term feeding studies show that cognitive manipulations (eg, time, energy content, food labeling, portion size) significantly influence appetitive ratings and subsequent energy intake (19–22). Indeed, the perceived energy content of a food may better predict self-reported appetitive

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sensations than the true energy content (23, 24). Whether expected appetitive effects differ between beverage and solid food forms or between populations potentially at risk for positive energy balance associated with beverage consumption is poorly characterized.

In comparison with lean individuals, the obese reportedly have higher beverage intake and experience greater weight loss (25) or gain (26) with reduced or increased beverage consumption, respectively. Obese individuals may have less precise regulatory systems (27) and more rapid gastrointestinal motility, leading to increased energy intake due to a rapid loss of satiety (28). These responses may specifically hold after beverage compared with solid food consumption as numerous studies have failed to document differences in energy intake after solid preload consumption (6). In addition, habitual exercisers have more accurate dietary compensation after beverage preloads than do nonexercisers (29), suggesting that exercise not only increases energy expenditure but also improves appetite control sensitivity (30).

The primary aim of the present trial was to contrast appetitive, dietary, gastric emptying, orocecal transit time, and selected endocrine (insulin, GLP-1, CCK, and ghrelin) responses to pre-ingestive (cognitive and orosensory) properties of energy-matched liquid (in beverage form) and solid food forms to identify plausible mechanisms for beverage-specific effects on body weight. The primary hypotheses tested were that consumption of an oral liquid and expectation it would remain a liquid in the gastrointestinal tract would lead to weaker appetitive effects (ie, greater hunger and lower fullness sensations), more rapid gastric-emptying and orocecal transit times, lower satiety hormone release (ie, GLP-1, CCK), lesser orexigenic hormone suppression (ie, ghrelin), and reduced energy intake compensation compared with when a solid food was consumed and expected to remain solid in the gastrointestinal tract. Furthermore, it was predicted that beliefs about food form would have especially weak effects in obese and unfit participants compared with their lean and fit counterparts.

SUBJECTS AND METHODS

Participant eligibility

Eligibility criteria included the following: age of 18–50 y, BMI (in kg/m2) of 18–23 (lean) or 30–35 (obese), body fat percentage in the lower (lean) or upper (obese) tertile for age and sex, cardiorespiratory fitness (estimated maximum aerobic power) in the upper (fit) or lower (unfit) tertile for age and sex (31, 32), a dietary restraint score <11 (33), consistent diet and activity patterns, not pregnant or lactating, glucose tolerant, not taking medications known to influence appetite or metabolism, and a self-reported breakfast and lunch consumer. All participants signed an informed consent form approved by the Purdue University Institutional Review Board and received monetary compensation.

Experimental design and procedures

The study followed a 4-arm, randomized crossover design with a 1-wk washout period between sessions. Analyses were based on a mixed-model, repeated-measures design with cognitive information and sensory properties related to food form as within-subject factors and body fat percentage (lean compared with obese) and cardiorespiratory fitness (fit compared with unfit) as between-subject factors. Equal numbers were recruited in each participant group: lean/fit, lean/unfit, obese/fit, or obese/unfit. Participant height was measured without shoes or socks with a Holtain stadiometer (Holtain Ltd). Fasting-state body weight was measured to the nearest 0.1 kg after the participant had voided. Fasting-state whole-body density was determined by using a whole-body plethysmography system (BodPod; Life Instrument Inc). Whole-body percentage body fat was estimated from body density by using the 2-compartment Siri equation (34).

To assess cardiorespiratory fitness, participants completed the YMCA Submaximal Cycle Ergometer Test (35) to estimate maximal oxygen uptake (VO2max) (36).

To standardize testing conditions, participants consumed their customary breakfast and reported to the laboratory at their habitual lunchtime after refraining from eating for >3 h on each test day. Appetitive sensations were rated, and testing continued if hunger was rated greater than “strong” and fasting glucose concentrations were <6.1 mmol/L. An indwelling catheter was inserted, and after a 15-min rest, breath and blood samples and subjective appetite ratings were obtained. Participants were shown the session’s preload, informed of its postingestive properties through a demonstration, and allowed 10 min to consume the preload. Breath, blood, and subjective sensory and appetite ratings were collected immediately after preload consumption (time = 0) and at 15, 30, 45, 60, 90, 120, 180, and 240 min. During this time, participants were semisupine and isolated from all food-related cues. At the end of the session, participants consumed an ad libitum, weighed challenge meal of macaroni and cheese (380 kcal/100 g) and 350 mL water. They were instructed to eat as much as it took to reach a comfortable level of fullness (3 on a 9-point scale: 1 = extremely full, 9 = not full at all), and intake was recorded. Participants also completed diet records and appetite ratings for the remainder of the testing day.

Study preloads

The preloads corresponded to ~10% of individual daily energy requirements (37) (0% fat, 88.4% carbohydrate, 11.6% protein). Participants were placed into 1 of 3 groups according to their estimated energy needs: 175, 225, or 275 kcal. One test session required consumption of a clear, cherry-flavored unthickened beverage (viscosity of ~10 mPa·s). The treatment demonstration involved pouring the preload into a clear liquid that participants were told was gastric acid but was actually tap water. This session was referred to as the “liquid to liquid” (L-L) session because participants consumed a liquid in the form of a beverage and believed it would remain liquid in their stomach.

Another session, referred to as “liquid to solid” (L-S), involved the researcher pouring a 1% alginic solution that resembled the cherry-flavored beverage into a 5% calcium chloride solution (“gastric acid”). This resulted in an instantaneous formation of a solid mass that participants were informed would occur in their stomach. However, the actual preload consumed was identical to the previously described session and only differed in the expected gastric food form (ie, liquid or solid).
A third session involved consumption of 1” × 1” × 1” cherry-flavored gelatin cubes. Texture analysis of the dense gelatin cubes measured an average peak bloom strength of 313.8 ± 12 g (TA.XTplus; Stable Microsystems Ltd). The demonstration involved placement of a cube into warm water (“gastric acid”), resulting in liquefaction in <10 s, and was referred to as the “solid to liquid” (S-L) session. Whereas the cubes were solid in the oral cavity and masticated at a fixed rate, timed to a metronome, they were isocaloric to the beverage preload and assumed to liquefy in the stomach in seconds based on simulated gastric models.

The fourth session, “solid to solid” (S-S), involved the same gelatin cubes, but participants were informed that the consistency would remain solid in the stomach. To demonstrate this, a cube was placed into cold water (“gastric acid”) where it remained solid, thereby leading to the expectation that it would remain a gastric solid when, in fact, it would rapidly liquefy and resemble all other preloads.

To retain the rheologic properties while keeping the macronutrient composition equal between food forms, participants consumed 20–25 capsules (on the basis of energy needs) filled with unflavored gelatin or maltodextrin with 150–200 mL water. Capsules consumed on oral-liquid testing days contained gelatin present in the oral-solid food forms. Capsules consumed on oral-solid testing days contained maltodextrin that was dissolved in the oral-liquid preloads. Participants were not aware of capsule contents.

**Sensory and appetitive ratings**

Viscosity, hunger, fullness, and desire to eat (38) were measured on 100-mm visual analog scales with end anchors of “not at all” to “extremely.”

**Gastric-emptying and orocecal transit times**

Immediately after preload consumption, participants consumed 10 g liquid lactulose (39) and 1.5 g liquid acetaminophen (40) to permit estimation of orocecal transit times and gastric emptying, respectively. Serum acetaminophen was quantified via enzymatic colorimetry by using the Cobas Integra 400 Analyzer (Roche Diagnostics). Orocecal transit times were measured through hydrogen analysis of end-alveolar air samples (39, 41) (QuinTron SC MicroLyzer; QuinTron Instrument Co).

**Biochemical analyses**

Blood was collected into ice-cooled, evacuated EDTA-coated tubes, and the protease inhibitors dipeptidyl peptidase-IV (Millipore), 4-(2-aminoethyl) benzenesulfonyl fluoride hydrochloride (Roche Diagnostics), and aprotinin (Phoenix Pharmaceuticals) were added immediately per the manufacturers’ instructions to prevent GLP-1, ghrelin, and CCK degradation, respectively. Samples were centrifuged at 4°C, separated into aliquots, and frozen at −80°C until analyzed. In addition, after centrifugation plasma ghrelin was acidified with HCl (42). Commercial ELISAs were used to determine active plasma GLP-17–36 (EGLP-35K; Millipore), active n-octanoyl ghrelin (EZGRA-88K; Millipore), and CCK26–33 (FEK-069–04; Phoenix Pharmaceuticals). The limits of detection were 1.97, 7.40, and 5.21 pmol/L with intraassay CVs of 7%, 1.7%, and 10%, respectively. All samples for each participant were analyzed in duplicate on the same assay plate. Serum insulin concentrations were measured via electrochemiluminescence immunoassays by using an Elecsys 2010 analyzer (Roche Diagnostics), and serum acetaminophen and glucose concentrations were measured by enzymatic colorimetry via the Cobas Integra 400 Analyzer (Roche Diagnostics). The limits of detection were 1.38 pmol/L, 1.32 μmol/L, and 0.12 mmol/L with intraassay CVs of 1.9%, 3.1%, and 0.4%, respectively.

**Dietary intake**

Energy intake was recorded on test and nontest days and analyzed with the use of the University of Minnesota Nutrition Data System for Research 2009.

**Statistical analysis**

Statistical analyses were performed with the use of SPSS software, version 17.0 (SPSS Inc). Significance was defined as P < 0.05, 2-tailed. All data were expressed as means ± SEMs unless stated otherwise. Treatment effects were tested by repeated-measures ANOVA with a Bonferroni correction for multiple comparisons. When significant effects were noted, AUC was determined by using the trapezoidal rule. Associations between appetite and energy intake with study outcomes were assessed via Pearson’s correlation coefficients.

**RESULTS**

**Participant characteristics**

Three hundred eighty-one participants completed the initial screening questionnaire. The 81 individuals meeting initial eligibility criteria completed additional screening procedures, and 57 participants met all inclusion criteria. Five individuals discontinued participation for the following reasons: scheduling conflicts (2 individuals), initiated an exercise regimen (1 individual), and unable to set catheters (2 individuals). Contrary to the hypotheses, subgroup analyses were unremarkable, so all participant data were pooled.

Fifty-two adults (23 men, 29 women; mean age 65.5 ± 6.1 y) as the least viscous (all different from each other, P < 0.05). Orosensory (actual food form in the oropharynx) and cognitive (expected food form in the gastrointestinal tract) contributions were observed. To further examine the
Gastric-emptying and orocecal transit times resulted from consumption of an oral-solid (mean of S-L and S-S; 121.6 min, $P < 0.001$) or perceived gastric-solid preload (mean of S-L and S-S; 97.8 min, $P = 0.005$). There was a trend for a cognitive contribution ($P = 0.08$), and faster gastric-emptying times were associated with greater energy intake after consumption of perceived gastric-liquid preloads (L-L: $r = -0.489$, $P < 0.001$; S-L: $r = -0.313$, $P = 0.03$).

Consumption of the S-S preload resulted in delayed orocecal transit times (137 ± 6.5 min) compared with all other treatments (L-L: 83.5 ± 8.3 min, $P < 0.001$; L-S: 104 ± 6.9 min, $P < 0.001$; S-L: 112 ± 7.6 min, $P = 0.047$) (Figure 2B). Oro sensory and cognitive effects were noted. Greater orocecal transit times resulted from consumption of an oral-solid (mean of S-L and S-S; 125 ± 5.3 min) or a perceived gastric-solid preload (mean of L-S and S-S; 121 ± 5.7 min) and compared with an oral-liquid (mean of L-L and L-S; 93.9 ± 6.1 min, $P < 0.001$) or perceived gastric-liquid preload (mean of L-L and S-L; 97.8 ± 5.9 min, $P = 0.004$). Prolonged orocecal transit times after consumption of oral-solid preloads were associated with lower hunger ratings (S-L: $r = -0.331$, $P = 0.02$; S-S: $r = -0.286$, $P = 0.04$).

### Glucose and endocrine responses

The magnitude of change in glucose $C_{\text{max}}$ ($C_{\text{max}}$ minus baseline) was greater after the L-L preload (3.02 ± 0.14 pmol/L) compared with all other treatments ($L-L: r = -0.489, P < 0.001$; $L-S: r = -0.313, P = 0.03$). Consumption of the S-S preload resulted in decreased plasma glucose levels (mean ± SEM $C_{\text{max}}$: 95.6 ± 5.3 min) compared with all other treatments (L-L: 97.8 ± 5.6 min, $P = 0.005$). There was a trend for a cognitive contribution ($P = 0.08$), and faster gastric-emptying times were associated with greater energy intake after consumption of perceived gastric-liquid preloads (L-L: $r = -0.489$, $P < 0.001$; S-L: $r = -0.313$, $P = 0.03$).

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compared with both S-L (2.55 ± 0.13 pmol/L, P = 0.003) and S-S (2.63 ± 0.12 pmol/L, P = 0.012) preloads. Main effects of treatment and treatment-by-time interactions were observed for insulin [F(3,153) = 33.1, P < 0.001; F(27,1377) = 5.63, P < 0.001], GLP-1 [F(3,153) = 6.72, P < 0.001; F(27,1377) = 4.02, P < 0.001], and ghrelin [F(3,150) = 5.80, P = 0.001; F(27,1350) = 1.88, P = 0.004] (Figure 3, A–C). Insulin and GLP-1 AUC_t were lower after L-L and L-S preloads than after S-L and S-S preloads (all P < 0.05). Opposite responses were observed for ghrelin AUC_t, and nadir concentrations were lower after S-S (45.9 ± 3.4 pmol/L, P = 0.019) compared with L-L (55.0 ± 3.3 pmol/L, P = 0.001) and L-S (57.9 ± 4.4 pmol/L, P = 0.001) preloads. Higher insulin (P < 0.001) and GLP-1 (P = 0.011) and lower ghrelin AUC_t (P < 0.001) were noted with oral solids (mean of S-L and S-S) compared with oral liquids (mean of L-L and L-S), which is consistent with an orosensory effect. We did not observe significant treatment effects for cholecystokinin.

Energy intake

Challenge meal intake was greater after L-L (720 ± 40 kcal) compared with L-S (583 ± 35 kcal, P = 0.004) and S-S (562 ± 38 kcal, P < 0.001) preloads as well as after S-L (643 ± 44 kcal) compared with S-S (P = 0.008) preloads. The same pattern was observed for the remainder of the testing-day energy intake, which resulted in greater total testing-day intake after L-L (2370 ± 101 kcal) and S-L (2252 ± 113 kcal) preloads than after both L-S (1940 ± 77 kcal) and S-S (1853 ± 82 kcal) preloads (all P < 0.01). Consequently, energy intake was higher by ~21.8% on days when perceived gastric-liquid preloads were consumed (mean of L-L and S-L; 2311 ± 95 kcal) compared with perceived gastric solids (mean of L-S and S-S; 1897 ± 72 kcal, P = 0.007). Water intake at the challenge meal and throughout the study visit day was not significantly different between study treatments.

DISCUSSION

The high energy intake from beverages is potentially problematic; evidence from animal behavioral studies (43, 44) and
short-term mechanistic human trials (17, 6, 45), suggest that fluids stimulate weak appetitive and compensatory dietary responses compared with energy-matched semisolids or solid items. The primary aim of this study was to isolate and characterize cognitive and orosensory influences stemming from consumption of liquid (in the form of a beverage) and solid food forms. These forms of input are known to evoke neurally mediated physiologic responses to food exposure (ie, cephalic phase responses), with implications for energy balance (46). Stimulation was accomplished by exposures to beverage and solid food forms orally and by providing information about the impending physical state of the test foods in the participant’s gastrointestinal tract. In addition to the fact that this cognitive manipulation led to multiple objectively measured differential responses, the effectiveness of the intervention was supported by numerous confirming spontaneous subjective comments from study participants (Table 1). The findings indicate that the mere expectation that food will be in one form or another in the gastrointestinal tract produces behavioral and physiologic responses likely to contribute to lower satiety effects (Figure 1) and weaker dietary compensation after beverage ingestion.

Initially, viscosity ratings of the 2 identical oral-liquid samples and the 2 oral-solid samples were higher if participants expected the preload to transform into or remain a solid in their gut. Previous work suggests that there is a direct relation between viscosity and postprandial hunger suppression and 24-h energy intake (4, 47, 48), purportedly through increased gastric viscosity and prolonged gastric emptying. However, gastric processing may rapidly reduce viscosity and result in similar gastric-emptying times (49). Thus, perceived oral and gastric meal viscosity may be an important mediator of these effects (16). The present findings also document a strong orosensory effect because oral-liquid stimulation led to more rapid gastric-emptying and orocecal transit times (Figure 2), a smaller increase of GLP-1 and insulin, as well as a smaller reduction in ghrelin compared with oral-solid stimulation (Figure 3). Slower gastric-emptying (48) and orocecal transit (50) times are associated with enhanced satiety, whereas insulin (51) and GLP-1 (52) are purported satiety hormones, and ghrelin is reportedly an orexigenic hormone (53). Thus, all noted responses would favor the observed weaker satiety effect for the oral-liquid stimulus that also led to a greater energy intake. This could reflect differential cephalic phase activation. A weaker insulin response to a beverage compared with a food has been documented previously, and preand postabsorptive responses were correlated (54). Hence, the rheologic properties of beverages provide a second mechanism by which beverages may hold weaker satiety properties and facilitate greater energy intake.

Although differential responses were noted on the basis of food form, these were further modified by cognitive manipulation. Energy intake was greater after L-L preload ingestion than after L-S preload ingestion, and after S-L than after S-S preload ingestion. This is supported by the cognitive influence on gastric-emptying and orocecal transit times, which were shorter with expectations that the gastrointestinal challenge would be liquid. A sensory contribution was also present because the differences were greatest for the L-L condition and weakest for the S-S preload.

Overall, it is notable that, for the appetitive and gastrointestinal transit time responses, both oral compared with stomach (cognitive) and beverage compared with solid (sensory) differences were observed. In contrast, the gastric-emptying and hormonal responses were more closely aligned with the sensory difference with no cognitive effect. Whether this shows differences between sensory and cognitive influences on these processes is worthy of further study.

This study sought to determine the cognitive and sensory contributions of differential responses to beverage and solid food forms in groups defined according to adiposity and fitness that may have varying sensitivities to cognitive or sensory food cues. However, similar to other studies (6, 20, 47), these data showed no distinct response patterns between such groups. This suggests processes other than cognition and orosensory stimulation mediate reported response differences in these specific groups.

The measurement precision afforded by conducting this trial in a laboratory setting is perhaps the trial’s primary weakness because its external validity remains to be established. However, the findings provide initial mechanistic support for the observed differential appetitive and dietary responses to beverage and solid foods (4, 17) and their likely influence on body weight.

The authors’ responsibilities were as follows—BAC: conceived and designed the study, conducted participant recruitment and testing, performed sample and data analyses, interpreted data, and generated the initial manuscript; RVC: performed hormone analyses and interpretation; and RDM: conceived and designed the study and performed data analyses and interpretation. All authors discussed the results and contributed to the final manuscript. None of the authors had a personal or financial conflict of interest.

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