Postprandial hypotension in older adults: Can it be prevented by drinking water before the meal?

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Background & aims: An important consequence of ageing is a tendency for postprandial blood pressure to decline, which can lead to fainting. As a possible countermeasure, we investigated in healthy older adults the impact of drinking water before a breakfast meal on postprandial cardiovascular and autonomic functions.

Methods: After a stable cardiovascular baseline recording for at least 20 min, twelve older adult (67 ± 1 y) test subjects ingested, in a crossover study design, either 100 mL or 500 mL of tap water over 4 min, which was followed by the consumption of the test breakfast meal (1708 kJ) over a period of 15 min. Then, cardiovascular recordings were resumed for 90 min after the meal. Eleven young (25 ± 1 y) and healthy subjects served as a control group. Measurements included beat-to-beat blood pressure, heart rate, impedance cardiography and autonomic variables.

Results: In older adults, systolic and diastolic blood pressure started to decline around 30 min after the meal, with the lowest values around 60 min; these effects were not observed in the young control group. Postprandial systolic blood pressure decreased between 30 and 90 min to a greater extent in response to 100 mL than to 500 mL (6.4 vs. 3.3 mmHg, P < 0.05). Drinking 500 mL of water tended to increase stroke volume, cardiac output and vagal markers to a greater extent than 100 mL.

Conclusions: Our data suggest that drinking a large volume (500 mL) of water before a meal may attenuate postprandial hypotension in older adults.

1. Introduction

An important consequence of ageing is the tendency for blood pressure to fall after eating a meal. Clinically, postprandial hypotension is defined as a reduction in blood pressure of 20 mmHg within 2 h after having a meal [1] and about two-thirds of older adults admitted to geriatric departments were found to have postprandial hypotension [1]. The most common symptoms of postprandial hypotension are dizziness, nausea, weakness and light-headedness [1] but more severe consequences such as syncope have also been reported. Meal composition appears to play a role in postprandial hypotension because carbohydrates, particularly glucose, have the greatest blood pressure lowering effect [2]. Even older adult individuals with no history or symptoms of postprandial hypotension tend to show some degree of blood pressure reduction after ingesting a carbohydrate meal [3].

Ingesting a meal causes pooling of blood in the abdominal vasculature that is accompanied by a drop in systemic peripheral resistance, activation of the sympathetic nervous system and increases in heart rate and cardiac output (CO) [4,5], but the rise in CO is not sufficient to counteract the dropping blood pressure, which could lead to presyncopal symptoms or even syncope. Drinking approximately 500 mL of water acutely increased blood pressure in autonomic failure patients as well as in older subjects [6] and improved orthostatic tolerance in healthy humans [7]. Moreover, in patients suffering from primary autonomic failure, ingestion of 480 mL of tap water in combination with a meal markedly improved the postprandial drop in blood pressure [8]. However, to
date, the precise mechanisms of postprandial hypotension are not well characterized, though inadequate autonomic nervous system responses to meal ingestion have been implicated [1].

In the study reported here, we investigated the effects of tap water in combination with a breakfast meal on cardiovascular parameters and cardiac vagal tone in older adults. We chose a light meal because older adults often consume small meals of about 400 kcal [9]. Furthermore, a breakfast meal is the least likely meal to be skipped by older adults [9]. Therefore, we evaluated heat-to-beat hemodynamic parameters and autonomic responses in a group of older individuals drinking either 100 ml or 500 ml before a light breakfast meal. The responses were compared to a control group comprised of young and healthy adults. It was hypothesized that ingestion of a higher amount of water would prevent the postprandial drop in blood pressure in older and generally healthy individuals.

2. Methods

2.1. Participants

Eleven young (four females; Age: 25 ± 1 years; Height: 171 ± 3 cm; Weight: 75 ± 3 kg; Body Mass Index: 25.8 ± 0.8 kg m⁻²) and twelve older (five females; Age: 67 ± 1 years; Height: 167 ± 2 cm; Weight: 75 ± 2 kg; Body Mass Index: 27.0 ± 0.8 kg m⁻²) non-smoking individuals were recruited. The participants in the young control group were recruited from local students and their friends, whilst older adults were recruited from relatives and friends of our local staff. Seven of the older subjects were normotensive, whilst five were taking medication for hypertension (β1 selective beta blockers (n = 2), angiotensin II receptor antagonists (n = 2) and a calcium-antagonist (n = 1)). The participants were requested to avoid alcohol and caffeine for 24h and were studied in the morning starting at 8am after an overnight (12h) fast. The treated hypertensive subjects refrained from taking their medication on the morning of the study. Written informed consent was obtained from each participant and the study protocol received local ethics committee approval in accordance with the Declaration of Helsinki.

2.2. Study design

All measurements were performed between 08.00 and 09.00 h in a temperature-controlled (22 ± 1 °C) quiet laboratory with the study participants seated in a comfortable armchair. Every older adult attended two separate experimental sessions, with each session (100 ml or 500 ml tap water before a meal) separated by at least two days according to a crossover study design; the young group served as a control for the condition with 100 ml before the meal. On arrival at the laboratory, subjects were asked to empty their bladders if necessary and to sit in a comfortable armchair. Subjects clothing consisted of a t-shirt, trousers and shoes. The equipment for cardiovascular monitoring was then attached.

Following a period for cardiovascular stability (20–30 min), a baseline recording was then made for 20 min. Then, the older adults ingested over four minutes either 100 ml or 500 ml of tap water at room temperature (the control group drank 100 ml water) which was followed by the consumption of the test meal over a period of 15 min and cardiovascular recordings were resumed for 90 min after the meal. The meal comprised a light breakfast of 100 g of toast-bread, over which was spread 12 g of butter and 30 g of jam. The total energy content was 1708 kJ, with carbohydrate, fat and protein respectively contributing to 60%, 31% and 9% of the energy intake. Such small meals are often consumed by older adults [9].

2.3. Cardiovascular recordings

Cardiovascular recordings were performed using a Task Force Monitor (TFM) (CNSystems, Medizintechnik, Graz, Austria) with data sampled at a rate of 1.000 Hz [10]. Continuous blood pressure was monitored using the Penaz principle from either the index or middle finger of the right hand and was calibrated to oscilometric brachial blood pressure measurements on the contralateral arm. Impedance cardiography measurements were performed using an improved estimate of thoracic volume [11], which allows calculation of stroke volume (SV). Electrode strips were placed at the neck and thoracic regions, the latter specifically at the midclavicular at the xiphoid process level (CNSystems standard electrode kits).

2.4. Autonomic measurements

High frequency (HF: 0.17–0.40 Hz) power components of RR-intervals (HF_RRI) were evaluated and given in absolute values (ms²). We used changes in the HF range of heart rate variability to assess parasympathetic activity because HF_RRI is primarily mediated by parasympathetic nerve modulation [12,13]. Powers of HF_RRI were analyzed after natural logarithmic transformation. Baroreflex sensitivity (BRS) was determined from spontaneous fluctuations in blood pressure and cardiac interval using the sequence technique [14].

2.5. Data and statistical analysis

Values of cardiac interval, systolic blood pressure (SBP), mean blood pressure and diastolic blood pressure (DBP)); SV and autonomic variables were averaged every 10 min during the baseline period and every 15 min during the 90 min post-drink period. CO was calculated as the product of SV and heart rate (HR), where HR was calculated from the appropriate cardiac interval. Total peripheral resistance (TPR) was calculated as mean blood pressure (MBP) divided by CO, where MBP was calculated as the result of DBP + 1/3 (SBP-DBP). Baseline values and responses to the intervention were statistically not different between the treated hypertensive and the normotensive group and were, therefore, combined for the final analysis.

Blood pressure variables started to decline significantly below baseline values around 30 min postprandial in older adults, therefore we chose the period from 30 to 90 min postprandial as appropriate for a statistical discrimination between the drink volumes used (Figs. 1–3, right panel).

Statistical analyses were performed using GraphPad Prism (Version 5, San Diego, CA, USA). All values are reported as means ± SE. Repeated-measures ANOVA with Dunnett’s post hoc testing were used to test for changes over time from baseline values. One-way ANOVA with Newman–Keuls post hoc testing was used to compare baseline values between the three groups. A paired t-test was used to compare mean changes of the 30–90 min post-drink period with baseline values and between the conditions. All reported P values are two-sided. For all tests, significance was set at P < 0.05.

3. Results

Baseline resting values were similar for hemodynamic and autonomic measurement parameters in older adults between the drinks. Baseline DBP and TPR were higher and SV, BRS and HF_RRI lower in the respective older adult group compared to the younger controls (Table 1). None of the test subjects showed unpleasant effects after ingesting the test meal regardless of the corresponding drink type. We could not find statistical differences related to blood
pressure variables — or to other observed variables (data not shown) — between the hypertension-treated and the untreated healthy older adults (Table 2).

3.1 Hemodynamic changes

Figure 1 shows changes for SBP, DBP and HR in response to ingestion of 100 mL of tap water before a meal in young and older adult humans, and also in the same older adults ingesting 500 mL tap water. In older adults, ingestion of 100 mL or 500 mL water before a breakfast meal led to decreased blood pressure variables below baseline levels around 30 min postprandial with the lowest values around 60 min (SBP 100 mL: -8.9 mmHg; SBP 500 mL: -5.2 mmHg; DBP 100 mL: -8.7 mmHg; DBP 500 mL: -5.2 mmHg) but not in younger controls. A comparison of mean responses between the two drink conditions between 30 and 90 min postprandial revealed a more pronounced drop for SBP when ingesting the lower volume (SBP: -6.4 vs -3.3 mmHg, \( P < 0.05 \)), whilst the drop in DBP was not significantly different between the drinks (DBP: -6.7 ± vs -4.2 mmHg, \( P = 0.32 \)). A comparison of mean responses in all drink groups over the 90 min study period found SBP as significantly higher in the young controls than in older adults (\( P < 0.05 \)), whilst DBP was found different comparing young controls with older adults ingesting 100 mL (\( P < 0.05 \)) but not 500 mL. In older adults, drinking 100 mL of water before the meal started to increase HR soon after ingestion with the
highest values around 60 min (+4.6 beats min⁻¹). The 500 mL condition showed a latency in the rise of HR but rose to similar levels as with 100 mL.

Figure 2 shows changes for SV, CO and TPR in response to ingestion of 100 mL of tap water before a meal in young and older adult humans, and also in the same older adults ingesting 500 mL tap water. Water ingestion (500 mL) before a breakfast meal increased SV soon over baseline values with a maximum around 30 min (+10.8 mL), whilst drinking 100 mL water before a breakfast meal increased SV more gradually with the highest value around 60 min (+7.0 mL). SV increased to a larger extent in older adults ingesting 500 mL compared to the young control group (P < 0.05), whilst no difference between the older adults drink conditions could be observed.

3.2. Autonomic changes

Figure 3 shows changes for BRS and HF_RRI in response to ingestion of 100 mL of tap water before a meal in young and older adult humans, and also in the same older adults ingesting 500 mL tap water. Water ingestion (500 mL) before a breakfast meal tended to increase BRS slightly over baseline levels, but drinking 100 mL before the breakfast meal significantly decreased BRS in older adults as well as in the younger controls (−0.83 and −2.6 ms mmHg⁻¹, respectively). The time course for HF_RRI showed an early peak value (0.66 ln ms²) shortly after ingestion.
500 mL water and then quickly returned towards baseline values, whilst ingestion of 100 mL (in older adults less pronounced than in the younger controls) resulted in a slow progressive decrease. A comparison between mean responses over the 90 min post-drink period observed a tendency for HF_RRI to increase after ingestion of 500 mL compared with 100 mL water in older adults ($+0.22 \pm 0.10$ ln ms$^2$, $P = 0.07$).

4. Discussion

To our knowledge, this is the first study where the hemodynamic and autonomic responses of water ingestion before a meal were investigated in older adults under controlled conditions in a crossover study design. We assessed blood pressure variables by continuous beat-to-beat cardiovascular measurements and observed (i) a postprandial reduction in SBP and DBP in older adults drinking 100 mL before the meal, which was not seen in a younger control group and (ii) an attenuation of postprandial hypotension, significant for SBP and a trend for DBP, when drinking a much larger drink volume of water. Thus, our results suggest that the amount of water drunk before a meal significantly influences postprandial blood pressure in older adults.

The postprandial blood pressure decrease in our older subjects is considered a normal response in older adults [3,15] and, although this blood pressure drop may have serious clinical consequences, effective countermeasures are still elusive. The physiological responses to water ingestion in humans have considerable impact on hemodynamics. In patients suffering from severe autonomic failure, drinking water increased blood pressure substantially [6,16] accompanied by a rise of plasma norepinephrine [6]. However, the pressure response due to ingestion of similar amounts of water is not seen in young and healthy subjects [6,17,18] despite an augmented muscle sympathetic nerve activity and a rise in plasma norepinephrine levels [18]. In our study, a standard light meal (1708 kJ), which was preceded by ingestion of 100 mL water, produced significant decreases in beat-to-beat measured SBP and DBP variables starting around 30 min postprandial in older adults but was not seen in a younger control group. Ingesting 500 mL of water before the meal significantly reduced the postprandial drop in SBP whilst DBP showed a trend towards an attenuation of the postprandial pressure drop. Therefore, it is conceivable that the amount of water ingested before a meal influences positively the postprandial SBP in older adults. In agreement with our results, a previous study observed no significant changes in SBP compared to baseline levels but a small drop in DBP after a breakfast meal plus water ingestion (350 mL) at a comparable composition (63% carbohydrates, 27% fat and 10% proteins) and energy content (1301 kJ) [19]. Our observation of a water-induced attenuation of the postprandial decrease in blood pressure is further supported by a study where gastric distension raised volume dependent blood pressure variables during intraduodenal glucose infusion [20]. Taken together, in response to the higher water drink volume, we observed an attenuation of the decrease in SBP by 3.1 mmHg compared to the lower drink volume. Whether this blunting is of clinical relevance remains to be determined in patients presenting symptoms of postprandial hypotension.

In the present study, we observed that postprandial SV and CO increased in response to water ingestion before a meal both in young controls and in older adults, with the latter’s response being greater, particularly with the large water volume. TPR decreased in response to the higher water drink volume, we observed an attenuation of the decrease in SBP by 3.1 mmHg compared to the lower drink volume. Whether this blunting is of clinical relevance remains to be determined in patients presenting symptoms of postprandial hypotension.
Table 1
Baseline data recorded 20 min before ingesting either 100 mL or 500 mL of tap water which was followed by consumption of a standard light meal (1708 kJ) in older adults and young controls drinking 100 mL.

<table>
<thead>
<tr>
<th></th>
<th>Controls 100 mL</th>
<th>Controls 500 mL</th>
<th>Older adults 100 mL</th>
<th>Older adults 500 mL</th>
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<tbody>
<tr>
<td>Systolic blood pressure, mmHg</td>
<td>112 ± 2</td>
<td>120 ± 3</td>
<td>119 ± 4</td>
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<tr>
<td>Diastolic blood pressure, mmHg</td>
<td>72 ± 2</td>
<td>83 ± 2*</td>
<td>81 ± 3*</td>
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<tr>
<td>Heart rate, beats min⁻¹</td>
<td>61 ± 3</td>
<td>62 ± 2</td>
<td>62 ± 2</td>
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<tr>
<td>Stroke volume, ml</td>
<td>78 ± 3</td>
<td>60 ± 2*</td>
<td>61 ± 3</td>
<td>61 ± 3</td>
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<tr>
<td>Cardiac output, L min⁻¹</td>
<td>4.7 ± 0.1</td>
<td>3.7 ± 0.2</td>
<td>3.7 ± 0.2</td>
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<tr>
<td>Total peripheral resistance, mmHg L⁻¹ min⁻¹</td>
<td>18.4 ± 0.8</td>
<td>26.5 ± 1.4</td>
<td>25.6 ± 1.7</td>
<td>25.6 ± 1.7</td>
</tr>
<tr>
<td>Baroreflex sensitivity, ms mmHg⁻¹</td>
<td>262.6 ± 16</td>
<td>103.3 ± 12</td>
<td>9.0 ± 0.6</td>
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<tr>
<td>High frequency power of RRI, ms²</td>
<td>10.57 ± 233</td>
<td>188 ± 62</td>
<td>159 ± 44</td>
<td>159 ± 44</td>
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</tbody>
</table>
| RRI, RR-interval (s); t, p < 0.05 compared to controls; t, p < 0.005 compared to young controls using one-way ANOVA with Newman–Keuls multiple comparison post-hoc testing. Data are presented as means ± SE.

Table 2
Baseline and mean cardiovascular data averaged from 30 to 90 min postprandial relative to baseline values and presented as a delta (Δ) (i.e. average from 30 to 90 min postprandial, minus the average over the 20 min baseline period).

<table>
<thead>
<tr>
<th></th>
<th>Hypertensives baseline</th>
<th>Normotensives baseline</th>
<th>Hypertensives Δ</th>
<th>Normotensives Δ</th>
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</thead>
<tbody>
<tr>
<td>SBP, mmHg (100 mL)</td>
<td>124 ± 6</td>
<td>118 ± 3</td>
<td>-7.2 ± 1.2</td>
<td>-5.8 ± 1.7</td>
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<tr>
<td>DBP, mmHg (100 mL)</td>
<td>85 ± 5</td>
<td>81 ± 1</td>
<td>-7.9 ± 1.4</td>
<td>-5.8 ± 1.5</td>
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<tr>
<td>SBP, mmHg (500 mL)</td>
<td>124 ± 8</td>
<td>115 ± 5</td>
<td>-4.4 ± 2.5</td>
<td>-2.5 ± 2.6</td>
</tr>
<tr>
<td>DBP, mmHg (500 mL)</td>
<td>85 ± 6</td>
<td>77 ± 3</td>
<td>-6.1 ± 3.3</td>
<td>-3.6 ± 2.4</td>
</tr>
</tbody>
</table>
| SBP: Systolic blood pressure; DBP: Diastolic blood pressure; Hypertensives refer to treated hypertensive subjects; Normotensives refer to normotensive subjects. 100 mL and 500 mL refer to the amount of water before the breakfast meal. An unpaired t-test was used to compare postprandial cardiovascular changes between normotensive (n = 7) and hypertensive (n = 5) older adults. No significant changes have been found. Data are presented as means ± SE and statistical significance was set at P < 0.05.

5. Conclusions
The present study evaluated beat-to-beat hemodynamic parameters and autonomic responses in a group of older individuals ingesting either 100 mL or 500 mL before a meal and in a young control group ingesting 100 mL. Our data suggest that drinking a larger volume of water before a meal attenuates the postprandial decrease in blood pressure in older adults.

Conflict of interest
The study authors declare no conflict of interests.

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Contributions of the authors
JPM and AGD designed research; BG and GV conducted research and recruited subjects; EKG analyzed the data and wrote the first draft of the manuscript; All authors contributed towards, read and approved the final manuscript.

References


