Urinary Sodium and Potassium Levels and Blood Pressure in Population with High Sodium Intake

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Received: 21 September 2020; Accepted: 6 November 2020; Published: 10 November 2020

Abstract: The purpose of this study was to examine the association of urinary sodium-to-creatinine ratio and potassium-to-creatinine ratio with blood pressure in a cross-sectional study comprising Korean adults who participated in the Healthy Twin Study. The participants consisted of 2653 men and women in the Healthy Twin Study aged ≥ 19 years. Participants’ urinary excretion of sodium, potassium, and creatinine was measured from overnight half-day urine samples. Food intake was assessed using a validated food frequency questionnaire. We examined systolic and diastolic blood pressures according to sodium- or potassium-to-creatinine ratios using the generalized linear model. We determined food groups explaining high urinary sodium- or potassium-to-creatinine ratio using the reduced rank regression and calculated sodium- or potassium-contributing food score. We observed that systolic blood pressure was higher among men and women in the highest quintile of urinary sodium-to-creatinine ratio or sodium-to-potassium ratio than it was in the lowest quintile. Geometric means (95% CIs) of the lowest and the highest quintiles of systolic blood pressure (mmHg) were 113.4 (111.8–115.0) and 115.6 (114.1–117.2; \( P \) for trend = 0.02), respectively, for sodium-to-creatinine ratio. The association between urinary sodium-to-creatinine and systolic blood pressure was more pronounced among individuals whose body mass index (BMI) was less than 25 kg/m² (\( P \) for interaction = 0.03). We found that vegetables, kimchi and seaweed intake contributed to high sodium intake and a sodium-contributing food score were associated with increased blood pressure. In our study, we identified the food groups contributing to high sodium intake and found that high urinary sodium levels were associated with increasing blood pressure among Korean adults.

Keywords: sodium; potassium; systolic blood pressure; diastolic blood pressure; Healthy Twin Study

1. Introduction

Hypertension is a major risk factor for cardiovascular disease [1], some cancer types [2–4], and diabetes [5]. Global Burden of Disease reported that diet may contribute to cardiovascular disease death by 52.9% [6]. Dietary factors are strongly associated with hypertension [7,8], and a reduction in
sodium intake has been regarded as a promising strategy to prevent hypertension [8,9]. The Korea National Health and Nutrition Examination Survey (KNHANES) in 2018 reported 28.3% of the Korean adults aged over 30 years had hypertension [10]. The prevalence of hypertension was similar to that in American adults aged over 18 years (29.0% in the NHANES 2015–2016) [11]. Given the high sodium intake in Korea, with the mean intake being 3255.0 mg/d in 2018, which is more than 1.5 times the recommended amount of World Health Organization [10], reduction in sodium intake has been a major public health message to decrease the risk of chronic disease in Korea.

Dietary assessment of sodium intake tends to underestimate the actual sodium intake [12] because sodium is added in cooking or at the table while eating and discretionary sodium intake is higher than nondiscretionary sodium intake in Koreans [13]. Therefore, measuring 24-h urinary sodium excretion was regarded as a desirable method for estimating sodium intake [14] and for disease association studies. As it is challenging to collect 24-h urine samples and extract urinary sodium in a large population, spot urine or half-day urine specimen are often collected.

The associations between urinary sodium levels and hypertension or blood pressure have been studied in several Western [15–18] and Korean cross-sectional studies [19–23]. These Korean studies including 148 to 19,476 participants found an increase in both systolic and diastolic blood pressure in individuals with high sodium excretion. A few randomized controlled trials including Dietary Approaches to Stop Hypertension (DASH) [9] and Trials of Hypertension Prevention Collaborative Research Group (TOHP) [24] found that sodium intake reduction lowered blood pressure levels. Also, several randomized controlled trials suggest the evidence that potassium supplementation decreased blood pressure [25].

In this study of more than 2000 adult participants, we examined the association between the sodium-to-creatinine ratio and potassium-to-creatinine ratio in urine and blood pressure in a cross-sectional study of Korean adults participating in the Healthy Twin Study, a large cohort study of twin families. We also determined food groups contributing to the higher sodium-to-creatinine ratio in urine, calculated the reduced rank regression (RRR) pattern score, and examined their associations with systolic and diastolic blood pressures.

2. Materials and Methods

2.1. Study Population

The Healthy Twin Study comprised of same-sex adult twin pairs aged 30 years or older and their first-degree family members; these participants were the ones who volunteered in response to a nationwide media advertisement and mailing campaign open to the general population. From August 2005 to January 2012, a total of 3320 study participants (1333 men and 1987 women) have been enrolled. Study design and methodology have been described elsewhere [26].

When we examined the association of blood pressure with urinary concentrations of sodium and potassium; we excluded participants if they were not adults (under age of 19 years; \( n = 9 \)), did not have information on blood pressure \( (n = 8) \), or did not have measurements of urinary sodium/potassium or creatinine \( (n = 275) \). We also excluded those who had been diagnosed with hypertension \( (n = 324) \) or cancer \( (n = 51) \) to avoid the potential effect of low sodium intake among participants with previous illnesses. Consequently, a total of 1034 men and 1619 women were included in the analysis of urinary sodium and potassium.

In the analysis of food groups and blood pressure, we excluded study participants who did not answer questions pertaining to rice items or left questions for more than 50 items unanswered on the food frequency questionnaire (FFQ) \( (n = 164) \), as well as those whose total energy intake was beyond three standard deviations from the loge-transformed mean energy intake \( (n = 27) \). Consequently, a total of 961 men and 1501 women were included in the analysis. The study was approved by the ethics committees at the Samsung Medical Center (IRB No. 2005-08-113), Busan Paik
Informed consent was obtained from all study participants.

2.2. Ascertainment of Blood Pressure and Urinary Sodium, Potassium and Creatinine Excretion

Blood pressure measurements were performed twice with a standard manual sphygmomanometer with participants in a sitting position after 5 min rest and the average of two readings was used. Participants voided all urine at 7 pm and collected urine in plastic containers until the next day morning. They recorded the start and the end time of urine collection. All participants collected urine for more than eight hours. We also considered creatinine concentrations to adjust for different volume. When we adjusted for hours of urine collection, the results did not differ. Urinary sodium and potassium after >8 h of urine collection were measured using indirect ion-selective electrodes, and urinary creatinine concentration was measured using Jaffè reaction on Siemens Advia 1800 analyzers (ADVIA1800, Siemens, Malvern, PA, USA). We collected overnight urine because participant burden is lower compared to 24-h urine collection and sodium from overnight urine has been considered as an alternative approach to replace 24-h urine [27]. In our previous validation study, we reported high correlation coefficient ($r = 0.837$) between sodium excretion directly from 24-h urine and estimated sodium excretion from overnight half-day urine using Kawasaki formula in 44 participants [28]. We did not use Kawasaki formulas in the main analysis because the estimated 24-h sodium excretion may be overestimated [29]. However, on performing a sensitivity analysis using Kawasaki formulas, the results were similar.

2.3. Assessment of Dietary and Non-Dietary Factors

Food intake of each participant was assessed using a validated semi-quantitative food-frequency questionnaire [30]. Participants were asked how frequently they consumed each food item during the previous year, and the frequencies were classified into the following nine categories: never, 1 time per month, 2–3 times per month, 1–2 times per week, 3–4 times per week, 5–6 times per week, 1 time per day, 2 times per day, and 3 times per day. Responses on frequencies of a specified serving size for each food item were converted to average daily intake. We consolidated 106 food items into 32 food groups based on their macronutrient composition and food preparation. Foods that did not match with a group or individual foods (e.g., nut, eggs, and coffee) remained as individual categories.

Participants provided information on smoking status, alcohol consumption, education level, marital status, and medical history and menopausal status in women through a questionnaire. Additionally, a trained and experienced interviewer conducted face-to-face interview to clarify incomplete or equivocal responses. Each participant visited one of the centers to undergo physical examination, biochemical assessment, and anthropometric measurements. We calculated the body mass index (BMI) by dividing the weight (kg) by the height squared ($m^2$). Pack-years smoked were calculated by multiplying the number of years smoked by the average number of cigarettes smoked per day. A woman was considered as postmenopausal if she did not experience menstruation for more than 12 months. If menopausal status was not reported ($n = 33$), we defined a woman as postmenopausal if the age was ≥50 years (average menopausal age in Korean women) when menopausal status was included as a covariate. The estimated glomerular filtration rate (eGFR) was determined based on the equation developed by Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) research group [31]. Circulating creatinine levels for eGFR calculation were measured using Jaffè reaction on the Siemens Advia 1800 analyzers.

2.4. Statistical Analysis

We calculated geometric means and 95% confidence intervals (CIs) of systolic and diastolic blood pressures according to quintiles of urinary sodium-to-creatinine ratio, potassium-to-creatinine ratio, and sodium-to-potassium ratio using the generalized linear model (GLM). We categorized the exposure by quintile for men and woman, separately. We added a random effect to the model to take family
into account. We loge-transformed systolic and diastolic blood pressures to improve normality and exponentiated them. To test for trend across quintiles, participants were assigned the median value of their quintile level. This variable was entered as a continuous term, the coefficient for which was evaluated by the Wald test. Models were adjusted for age (years, continuous), age square (years², continuous), BMI (kg/m², continuous), total alcohol consumption (men: none, <15, 15–30, ≥30 g/day; women: none, <3.75, 3.75–7.5, ≥7.5 g/day), marital status (unmarried, married and living together, and divorced or widowed or separated), pack-years of smoking (men: never smoker, <10, 10–20, ≥20 pack-years; women: never smoker, <3, 3–6, ≥6 pack-years), education level (less than high school graduate, high school graduate, and college or above), and menopausal status (premenopause, postmenopause) for women. We additionally adjusted for total energy intake in food group analysis, urinary potassium (mmol/L, continuous) in sodium analysis, and urinary sodium (mmol/L, continuous) in potassium analysis. We examined whether the associations varied by BMI (<25, ≥25 kg/m²), smoking status (never smoker, ever smoker), alcohol consumption (non-drinker, current drinker), eGFR levels (<90, ≥90 mL/min/1.73 m²), urinary potassium (<37, ≥37 mmol/L, median), education level (high school graduate or college above, less than high school graduate), and marital status (married, non-married). The statistical significance of interaction was tested by using a Wald test of the beta coefficient of the cross-product term. We applied the RRR method to determine food groups explaining urinary sodium- or potassium-to-creatinine ratio [32]. RRR extracts factors that explain maximum response variation. The coefficient vectors are eigenvectors of a covariance matrix and the eigenvalue explains the variation in the corresponding linear function of predictors. The factors are sorted by decreasing eigenvalues and the first factor explains the most variation in response. We used the intake of 32 food groups as a predictor and logarithmically transformed urinary sodium excretion as responses. We derived linear combinations of 32 food groups that explained as much variation of sodium-to-creatinine ratio and potassium-to-creatinine ratio as possible. SAS PROC PLS command was used to derive the factor loading value for each food group. Food groups with an absolute factor loading of ≥0.20 were selected as groups contributing to urinary sodium excretion. The intake of each selected food group was standardized by deducing the mean and dividing the standard error (standardized Z score), and the food score was calculated by multiplying the standardized Z score by factor loading of each food group. Weighted Z-standardized food score for each food group was summed and considered as a sodium-contributing food score. We examined the association between a sodium-contributing food score and systolic and diastolic blood pressures using the GLM. All analyses were conducted using SAS version 9.4 (SAS Institute, Cary, NC, USA). Significance was set at a two-sided P value < 0.05.

3. Results

The mean overnight half-day urinary sodium excretion was 134.7 mmol/L in men and 122.0 mmol/L in women. The mean overnight half-day urinary potassium excretion was 42.3 mmol/L in men and 41.7 mmol/L in women. Baseline characteristics of participants according to the quintile of urinary sodium-to-creatinine ratio, potassium-to-creatinine ratio, and sodium-to-potassium ratio are presented in Table 1. Compared to the participants with low urinary sodium-to-creatinine ratio, those with a high ratio were more likely to be older, never smokers, divorced, widowed, or separated and less likely to have attended college or higher education. Compared to the individuals with low urinary potassium-to-creatinine levels, those with high urinary potassium were more likely to be older, never smoker, and less likely to drink alcohol and have attended college or higher education. The baseline characteristics by sex was presented in the Table S1.
Table 1. Baseline characteristics according to the sodium-to-creatinine ratio, potassium-to-creatinine ratio and sodium-to-potassium ratio in men and women combined 1.

| Sodium-to-creatinine ratio | Quintile 1 | Quintile 2 | Quintile 3 | Quintile 4 | Quintile 5 |
|----------------------------|------------|------------|------------|------------|------------|
| No. of participants       | 530        | 531        | 531        | 531        | 530        |
| Sodium-to-creatinine ratio (range) | 1.60–10.40 | 10.40–14.20 | 14.20–18.32 | 18.34–24.51 | 24.51–102.48 |
| Age (years)               | 37.89 (11.16) | 40.78 (11.66) | 41.56 (11.72) | 44.41 (12.47) | 46.57 (13.28) |
| Sodium (mmol/L)           | 101.74 (44.62) | 121.55 (47.71) | 130.05 (53.62) | 138.31 (52.85) | 143.02 (54.41) |
| Potassium (mmol/L)        | 52.01 (27.62) | 45.01 (23.12) | 40.38 (20.80) | 38.23 (19.71) | 34.21 (17.56) |
| BMI (kg/m²)               | 23.69 (3.26) | 23.30 (3.20) | 23.20 (3.16) | 23.61 (3.21) | 23.45 (3.21) |
| Smoking status 2          | 301 (56.79) | 337 (63.47) | 330 (62.15) | 353 (66.48) | 398 (75.09) |
| Alcohol consumption 2     | 97 (18.30) | 125 (23.54) | 136 (25.61) | 161 (30.32) | 169 (31.89) |
| Education level           | 56 (10.57) | 81 (15.25) | 92 (17.33) | 123 (24.86) | 167 (31.51) |
| Marital status 2          | 170 (32.08) | 120 (22.60) | 120 (22.60) | 80 (15.07) | 63 (11.89) |
| Sodium-to-creatinine ratio (range) | 1.2–3.46 | 3.46–4.46 | 4.46–5.6 | 5.6–7.25 | 7.26–30.17 |
| Age (years)               | 37.05 (10.96) | 40.80 (11.75) | 42.24 (12.00) | 44.29 (12.33) | 46.86 (12.89) |
| Sodium (mmol/L)           | 127.82 (52.89) | 128.87 (53.20) | 123.81 (51.10) | 126.10 (51.53) | 128.09 (55.24) |
| Potassium (mmol/L)        | 32.69 (16.93) | 38.21 (18.21) | 41.83 (22.83) | 45.55 (22.81) | 51.58 (27.37) |
| BMI (kg/m²)               | 23.87 (3.24) | 23.64 (3.13) | 23.29 (3.18) | 22.22 (3.39) | 22.22 (3.07) |
| Smoking status 2          | 241 (45.39) | 304 (57.36) | 345 (64.97) | 397 (74.62) | 432 (81.66) |
| Alcohol consumption 2     | 77 (14.50) | 104 (19.62) | 145 (27.31) | 162 (30.45) | 200 (37.81) |
| Education level           | 59 (11.11) | 80 (15.09) | 98 (18.46) | 126 (23.68) | 165 (31.19) |
| Marital status 2          | 170 (32.08) | 120 (22.60) | 120 (22.60) | 80 (15.07) | 63 (11.89) |
| Sodium-to-potassium ratio (range) | 0.32–5.83 | 0.93–8.95 | 1.23–10.65 | 1.57–11.93 | 1.57–13.42 |
| Age (years)               | 41.22 (11.75) | 42.04 (11.98) | 43.50 (13.18) | 42.67 (12.67) | 41.79 (12.49) |
| Sodium (mmol/L)           | 100.54 (45.82) | 118.62 (47.57) | 129.47 (51.04) | 138.31 (50.59) | 146.73 (55.79) |
| Potassium (mmol/L)        | 63.59 (29.72) | 47.05 (19.03) | 40.27 (15.89) | 34.12 (12.58) | 24.82 (10.18) |
| BMI (kg/m²)               | 23.38 (3.36) | 23.20 (3.31) | 23.28 (3.05) | 23.57 (3.21) | 23.83 (3.28) |
| Smoking status 2          | 375 (70.62) | 370 (68.91) | 350 (66.16) | 341 (64.10) | 283 (53.30) |
| Sodium-to-potassium ratio (range) | 0.32–5.83 | 0.93–8.95 | 1.23–10.65 | 283 (53.30) | 161 (30.32) |
Table 1. Cont.

| Quintile 1 | Quintile 2 | Quintile 3 | Quintile 4 | Quintile 5 |
|------------|------------|------------|------------|------------|
| Alcohol consumption | | | | | |
| Never | 150 (28.25) | 133 (25.09) | 143 (27.03) | 144 (27.07) | 118 (22.22) |
| Past | 49 (9.23) | 49 (9.25) | 59 (11.15) | 34 (6.39) | 42 (7.91) |
| Current | 331 (62.34) | 348 (65.66) | 327 (61.81) | 354 (66.54) | 371 (69.87) |
| Education level | | | | | |
| Less than high school graduate | 88 (16.57) | 101 (19.06) | 112 (21.17) | 120 (22.56) | 107 (20.15) |
| High school graduate | 164 (30.89) | 186 (35.09) | 176 (33.27) | 184 (34.59) | 191 (35.97) |
| College or above | 277 (52.17) | 240 (45.28) | 241 (45.56) | 225 (42.29) | 232 (43.69) |
| Marital status | | | | | |
| Never married | 119 (22.41) | 114 (21.51) | 96 (18.15) | 108 (20.30) | 116 (21.85) |
| Married | 382 (71.94) | 391 (73.77) | 389 (73.53) | 380 (71.43) | 376 (70.81) |
| Divorced or widowed or separated | 29 (5.46) | 25 (4.72) | 44 (8.32) | 40 (7.52) | 36 (6.78) |

Abbreviation: BMI, body mass index. 1 Values are means (SD) or number (percentage). 2 A few participants had missing values.

Table 2 shows the relationships of urinary sodium-to-creatinine ratio, potassium-to-creatinine ratio and sodium-to-potassium ratio with systolic and diastolic blood pressures. We observed that urinary sodium-to-creatinine ratio was positively associated with systolic blood pressure, while potassium-to-creatinine ratio was inversely associated with systolic blood pressure in men and women combined. Geometric means (95% CIs) of the lowest and the highest quintiles of systolic blood pressure (mmHg) were 113.4 (111.8–115.0) and 115.6 (114.1–117.2; P for trend = 0.02), respectively, for sodium-to-creatinine ratio, 115.1 (113.5–116.8) and 113.1 (111.5–114.7; P for trend = 0.04), respectively, for potassium-to-creatinine ratio, and 112.4 (110.9–113.9) and 115.6 (114.0–117.1; P for trend < 0.001), respectively, for sodium-to-potassium ratio. When we separated men and women, we observed increasing systolic blood pressure with increasing half-day urinary sodium-to-potassium ratio with systolic blood pressure in both men and women (P for trend < 0.05) (Table S2).

Table 2. Multivariate-adjusted geometric means (95% CIs) 1 of systolic and diastolic blood pressures (mmHg) according to urinary sodium to creatinine, potassium to creatinine, or sodium to potassium ratios

| Quadrant 1 | Quadrant 2 | Quadrant 3 | Quadrant 4 | Quadrant 5 |
|------------|------------|------------|------------|------------|
| Sodium-to-creatinine ratio | | | | | |
| Range | 1.6–10.4 | 10.4–14.2 | 14.2–18.3 | 18.3–24.5 | 24.5–102.5 |
| SBP (111.8–115.0) | (112.4–115.3) | (112.5–115.5) | (112.3–115.4) | (114.1–117.2) | 0.02 |
| DBP 72.3 (71.3–73.4) | 72.4 (71.4–73.5) | 72.5 (71.4–73.5) | 72.5 (71.4–73.5) | 73.2 (72.1–74.3) | 0.19 |
| Potassium-to-creatinine ratio | | | | | |
| Range | 1.2–3.5 | 3.5–4.5 | 4.5–5.6 | 5.6–7.3 | 7.3–30.2 |
| SBP (113.5–116.8) | (113.0–116.2) | (112.9–115.8) | (112.9–116.0) | (111.5–114.7) | 0.04 |
| DBP 73.1 (71.9–74.3) | 72.9 (71.8–74.0) | 72.5 (71.4–73.5) | 72.9 (71.8–73.9) | 71.9 (70.8–73.0) | 0.08 |
| Sodium-to-potassium ratio | | | | | |
| Range | 0.3–2.2 | 2.2–2.9 | 2.9–3.6 | 3.6–4.7 | 4.7–13.4 |
| SBP (110.9–113.9) | (112.1–115.1) | (113.1–116.1) | (113.2–116.3) | (114.0–117.1) | <0.001 |
| DBP 71.4 (70.4–72.4) | 73.1 (72.1–74.1) | 72.9 (71.9–74.0) | 72.6 (71.5–73.6) | 73.0 (71.8–74.1) | 0.06 |

Abbreviation: CI, confidence interval; SBP, systolic blood pressure; DBP, diastolic blood pressure; BMI, body mass index. 1 Adjusted for age, age², BMI (kg/m²; continuous), pack years of smoking (never, <10, 10 to <20, ≥20 for men and never, <3, 3 to <6, ≥6 for women), alcohol consumption (none, <15, 15 to <30, ≥30 g/day for men and none, <3.75, 3.75 to <7.5, ≥7.5 g/day for women), marriage status (never, married, divorced or widowed or separated), education level (less than high school graduate, high school graduate, college or above), and menopause status (pre, post) for women. 2 We further adjusted for urinary potassium (mmol/L; continuous) in the model. 3 We further adjusted for urinary sodium (mmol/L; continuous) in the model.
We examined whether the associations of urinary sodium-to-creatinine ratio and potassium-to-creatinine ratio with blood pressure differed by BMI (<25, ≥25 kg/m²), smoking status (never smoker, ever smoker), alcohol consumption (none, current) (Table 3), eGFR (<90, ≥90 mL/min/1.73 m²), or urinary potassium (<37, ≥37 mmol/L, median) (Table S3).

Table 3. Multivariate-adjusted geometric means (95% CIs) of systolic and diastolic blood pressures (mmHg) according to urinary sodium-to-creatinine and potassium-to-creatinine ratios by BMI, smoking status, and alcohol consumption

|                      | n     | Quintile 1 (95% CI) | Quintile 2 (95% CI) | Quintile 3 (95% CI) | Quintile 4 (95% CI) | Quintile 5 (95% CI) | P Trend | P Interaction |
|----------------------|-------|---------------------|---------------------|---------------------|---------------------|---------------------|---------|--------------|
| Sodium-to-creatinine ratio |       |                     |                     |                     |                     |                     |         |              |
| SBP                  |       |                     |                     |                     |                     |                     |         |              |
| BMI (kg/m²)          |       |                     |                     |                     |                     |                     |         |              |
| <25                  | 1886  | 110.8 (108.9–112.7) | 111.1 (109.4–112.8) | 111.7 (110.0–113.4) | 112.1 (110.3–113.9) | 113.3 (111.5–115.1) | 0.01    | 0.03         |
| ≥25                  | 767   | 119.8 (116.5–123.1) | 120.3 (117.5–123.2) | 119.5 (116.7–122.4) | 118.9 (115.9–121.9) | 120.4 (117.5–123.4) | 0.91    |              |
| Smoking status       |       |                     |                     |                     |                     |                     |         |              |
| Never smoker         | 1719  | 113.1 (111.3–115.0) | 113.4 (111.7–115.0) | 114.1 (112.4–115.9) | 113.5 (111.8–115.2) | 115.3 (113.5–117.0) | 0.04    | 0.34         |
| Ever smoker          | 925   | 114.0 (110.7–117.4) | 114.2 (111.2–117.4) | 113.6 (110.5–116.7) | 114.2 (111.1–117.4) | 114.6 (111.3–118.1) | 0.72    |              |
| Alcohol consumption  |       |                     |                     |                     |                     |                     |         |              |
| Non-drinker          | 920   | 112.1 (109.0–115.2) | 113.5 (111.1–116.0) | 114.4 (111.8–117.1) | 114.0 (111.4–116.7) | 114.0 (111.4–116.6) | 0.39    | 0.52         |
| Current drinker      | 1732  | 113.8 (111.8–115.7) | 113.5 (111.7–115.4) | 113.5 (111.6–115.4) | 113.3 (111.5–115.2) | 116.2 (114.2–118.1) | 0.03    |              |
| DBP                  |       |                     |                     |                     |                     |                     |         |              |
| BMI (kg/m²)          |       |                     |                     |                     |                     |                     |         |              |
| <25                  | 1886  | 70.4 (69.3–71.9)    | 71.0 (69.8–72.2)    | 70.8 (69.6–72.1)    | 71.1 (69.8–72.3)    | 71.4 (70.2–72.6)    | 0.30    | 0.65         |
| ≥25                  | 767   | 75.9 (73.9–77.9)    | 75.6 (73.6–77.7)    | 76.0 (73.9–78.1)    | 76.3 (74.1–78.5)    | 76.9 (74.8–79.1)    | 0.32    |              |
| Smoking status       |       |                     |                     |                     |                     |                     |         |              |
| Never smoker         | 1719  | 72.4 (71.7–73.7)    | 72.3 (71.0–73.5)    | 72.5 (71.2–73.8)    | 72.2 (70.9–73.5)    | 73.0 (71.7–74.3)    | 0.38    | 0.75         |
| Ever smoker          | 925   | 72.0 (70.7–74.2)    | 72.3 (70.2–74.4)    | 71.8 (69.7–73.9)    | 72.3 (70.2–74.5)    | 72.2 (69.8–74.7)    | 0.87    |              |
| Alcohol consumption  |       |                     |                     |                     |                     |                     |         |              |
| Non-drinker          | 920   | 72.4 (70.3–74.5)    | 72.8 (71.7–74.5)    | 73.7 (71.8–75.6)    | 73.2 (71.3–75.1)    | 72.5 (70.5–74.5)    | 0.85    | 0.83         |
| Current drinker      | 1732  | 72.1 (70.9–73.4)    | 71.8 (70.6–73.1)    | 71.6 (70.3–73.0)    | 71.7 (70.4–73.1)    | 75.3 (72.0–74.6)    | 0.10    |              |
| Potassium-to-creatinine ratio |     |                     |                     |                     |                     |                     |         |              |
| SBP                  |       |                     |                     |                     |                     |                     |         |              |
| BMI (kg/m²)          |       |                     |                     |                     |                     |                     |         |              |
| <25                  | 1886  | 112.9 (110.9–114.9) | 112.1 (110.2–113.9) | 112.5 (110.9–114.1) | 112.4 (110.6–114.3) | 110.3 (108.5–112.2) | 0.03    | 0.94         |
| ≥25                  | 767   | 120.6 (117.4–124.0) | 120.0 (116.9–123.1) | 118.6 (115.8–121.6) | 116.4 (114.4–122.1) | 120.4 (117.5–123.3) | 0.89    |              |
| Smoking status       |       |                     |                     |                     |                     |                     |         |              |
| Never smoker         | 1719  | 114.3 (112.3–116.3) | 113.9 (112.1–115.7) | 114.1 (112.4–115.7) | 114.3 (112.6–116.1) | 113.1 (111.3–114.8) | 0.29    | 0.02         |
| Ever smoker          | 925   | 116.1 (112.6–119.7) | 115.0 (111.8–118.3) | 114.2 (111.0–117.4) | 114.4 (111.1–117.7) | 112.1 (108.9–115.3) | 0.03    |              |
| Alcohol consumption  |       |                     |                     |                     |                     |                     |         |              |
| Non-drinker          | 920   | 113.5 (110.3–116.7) | 113.2 (110.6–116.0) | 115.1 (112.5–117.8) | 114.2 (111.6–116.8) | 112.6 (110.0–115.3) | 0.35    | 0.06         |
| Current drinker      | 1732  | 115.3 (113.3–117.3) | 114.8 (112.8–116.8) | 113.9 (112.1–115.8) | 114.1 (112.2–116.1) | 112.9 (111.0–114.9) | 0.03    |              |
Table 3. Cont.

| Quintile 1          | Quintile 2          | Quintile 3          | Quintile 4          | Quintile 5          | P Trend | P Interaction |
|---------------------|---------------------|---------------------|---------------------|---------------------|---------|---------------|
| **BMI (kg/m²)**     |                     |                     |                     |                     |         |               |
| <25                 | 71.6 (70.2–73.0)    | 71.2 (70.0–72.5)    | 71.3 (70.2–72.5)    | 71.4 (70.2–72.7)    | 69.8 (68.5–71.0) | 0.01 | 0.32 |
| ≥25                 | 76.6 (74.4–78.8)    | 76.1 (74.0–78.3)    | 74.8 (72.7–77.1)    | 74.3 (74.3–78.4)    | 69.8 (74.9–79.2) | 0.61 |       |
| **Smoking status**  |                     |                     |                     |                     |         |               |
| Never smoker        | 73.2 (71.8–74.6)    | 72.4 (71.1–73.7)    | 72.2 (70.9–73.5)    | 72.6 (71.4–73.9)    | 72.1 (70.8–73.3) | 0.36 | 0.15 |
| Ever smoker         | 75.0 (70.7–75.4)    | 73.1 (70.9–75.3)    | 72.4 (70.7–75.3)    | 72.9 (68.0–72.3)    | 70.1     | 0.03 |       |
| **Alcohol consumption** |                   |                     |                     |                     |         |               |
| Non-drinker         | 73.1 (70.9–75.4)    | 72.6 (70.7–74.5)    | 73.4 (71.5–75.4)    | 73.1 (71.4–74.8)    | 72.4 (70.5–74.3) | 0.47 | 0.12 |
| Current drinker     | 72.8 (71.4–74.2)    | 72.9 (71.5–74.2)    | 71.9 (70.6–73.2)    | 72.4 (71.1–73.7)    | 71.3 (70.0–72.6) | 0.05 |       |

Abbreviation: CI, confidence interval; SBP, systolic blood pressure; DBP, diastolic blood pressure; BMI, body mass index. 1 Adjusted for age, age², BMI (kg/m²; continuous), pack years of smoking (never, <10, 10 to <20, ≥20 for men and never, <3, 3 to <6, ≥6 for women), alcohol consumption (none, <15, 15 to <30, ≥30 g/day for men and none, <3.75, 3.75 to <7.5, ≥7.5 g/day for women), marriage status (never, married, divorced or widowed or separated), education level (less than high school graduate, high school graduate, college or above), and menopause status (pre, post) for women. 2 We further adjusted for urinary sodium (mmol/L; continuous) in the model. 3 We further adjusted for urinary potassium (mmol/L; continuous) in the model.

When we fitted RRR to determine food groups contributing to a high urinary sodium-to-creatinine ratio, we found the intake of vegetables, soy paste, kimchi (Korean-style fermented vegetables), and seaweed to be positively and the intake of poultry and pizza to be inversely associated with high urinary sodium-to-creatinine ratio (Table 4). A higher sodium-contributing food score was associated with increasing systolic and diastolic blood pressure in men and women combined (Table 5). The associations of higher sodium-contributing food score with systolic or diastolic blood pressure did not vary by BMI (<25, ≥25 kg/m²) (Table S4). Table 6 shows the associations of selected food groups with blood pressures. There were significant associations of systolic and diastolic blood pressures with intakes of vegetables, kimchi, and seaweed (P for trend < 0.03). We found positive association for intakes of vegetables, potato and sweet potato, and fruit, but negative association for intakes of ramen, soft drink, poultry, red meat and beef soup, processed meat, noodle, and pizza, with urinary potassium-to-creatinine ratio (Table S5). Potassium-contributing food score was not associated with blood pressure (Table S6).

Table 4. Factor loading values of food groups contributing to urinary sodium-to-creatinine ratio through reduced rank regression analysis.

| Food Groups       | Factor Loadings 1 |
|-------------------|-------------------|
| **Positive associations** |                     |
| Vegetables        | +0.461            |
| Soy paste         | +0.304            |
| Kimchi            | +0.284            |
| Seaweed           | +0.233            |
| **Inverse associations** |                   |
| Poultry           | −0.295            |
| Pizza             | −0.273            |

1 The food groups with the absolute value of factor loading greater than 0.20 were selected.
### Table 5. Multivariate-adjusted geometric means (95% CIs) of systolic and diastolic blood pressures (mmHg) according to sodium-contributing food score.

|       | Quintile 1          | Quintile 2          | Quintile 3          | Quintile 4          | Quintile 5          | P Trend |
|-------|---------------------|---------------------|---------------------|---------------------|---------------------|---------|
| All   | 113.2 (111.6–114.8) | 113.1 (111.5–114.7) | 113.6 (112.1–115.2) | 115.2 (113.5–116.9) | 114.5 (112.9–116.2) | 0.03    |
|        | 71.7 (70.6–72.8)    | 71.8 (70.7–73.0)    | 72.3 (71.3–73.3)    | 73.3 (72.2–74.5)    | 72.9 (71.7–74.1)    | 0.02    |
| Men   | 119.6 (116.8–122.6) | 118.0 (115.1–120.9) | 119.4 (116.7–122.1) | 120.0 (117.0–123.2) | 121.4 (118.5–124.4) | 0.07    |
|        | 74.5 (72.6–75.6)    | 73.7 (71.9–75.6)    | 74.5 (72.8–76.2)    | 75.8 (73.9–77.7)    | 76.3 (74.3–78.4)    | 0.02    |
| Women | 108.4 (106.0–110.9) | 109.3 (107.0–111.7) | 109.9 (107.6–112.2) | 110.8 (108.1–113.5) | 110.7 (107.4–112.1) | 0.21    |
|        | 68.9 (67.3–70.5)    | 69.8 (68.1–71.4)    | 70.3 (68.7–71.9)    | 70.8 (69–72.6)      | 70.1 (68.5–71.8)    | 0.15    |

Abbreviation: CI, confidence interval; SBP, systolic blood pressure; DBP, diastolic blood pressure. 1 Adjusted for age, age², BMI (kg/m²; continuous), pack years of smoking (never, <10, 10 to <20, ≥20 for men and never, <3, 3 to <6, ≥6 for women), alcohol consumption (none, <15, 15 to <30, ≥30 g/day for men and none, <3.75, 3.75 to <7.5, ≥7.5 g/day for women), marriage status (never, married, divorced or widowed or separated), education level (less than high school graduate, high school graduate, college or above), menopause status (pre, post) for women, and urinary potassium (mmol/L; continuous).

### Table 6. Multivariate-adjusted geometric means (95% CIs) of systolic and diastolic blood pressure according to each food group with a high sodium contributing food score.

|       | Quartile 1          | Quartile 2          | Quartile 3          | Quartile 4          | P Trend |
|-------|---------------------|---------------------|---------------------|---------------------|---------|
| SBP   | 113.7 (112.2–115.2) | 113.6 (112.1–115.1) | 113.3 (111.8–114.7) | 115.3 (113.8–117.0) | 0.02    |
| Vegetables | 113.4 (111.8–114.9) | 113.2 (111.8–114.7) | 114.8 (113.4–116.3) | 114.3 (112.8–115.9) | 0.23    |
| Soy paste | 113.0 (111.4–114.1) | 112.7 (113.2–116.1) | 114.6 (113.4–116.7) | 115.0 (112.6–116.7) | 0.007   |
| Kimchi | 113.1 (111.7–114.5) | 114.2 (112.2–115.2) | 113.7 (112.3–116.2) | 115.2 (113.6–116.7) | 0.02    |
| Seaweed | 114.0 (112.4–114.6) | 114.7 (112.2–115.2) | 113.7 (112.6–115.2) | 113.9 (112.3–115.4) | 0.61    |
| Poultry | 114.4 (113.1–115.5) | 115.2 (113.1–117.4) | 113.1 (111.6–114.8) | 113.4 (111.9–115.0) | 0.11    |
| Pizza  | 113.1 (113.1–115.8) | 113.3 (113.1–117.4) | 113.3 (113.1–117.4) | 113.4 (111.6–114.8) | 0.11    |

Abbreviation: CI, confidence interval; SBP, systolic blood pressure; DBP, diastolic blood pressure. 1 Adjusted for age, age², BMI (kg/m²; continuous), pack years of smoking (never, <10, 10 to <20, ≥20 for men and never, <3, 3 to <6, ≥6 for women), alcohol consumption (none, <15, 15 to <30, ≥30 g/day for men and none, <3.75, 3.75 to <7.5, ≥7.5 g/day for women), marriage status (never, married, divorced or widowed or separated), education level (less than high school graduate, high school graduate, college or above), menopause status (pre, post) for women, total energy intake (kcal/d; continuous), and urinary potassium (mmol/L; continuous).
4. Discussion

We observed that urinary sodium-to-creatinine and sodium-to-potassium ratios were positively associated with systolic blood pressure and potassium-to-creatinine ratio was inversely associated with systolic blood pressure in 2653 men and women aged 19 years or older. We found more pronounced positive associations for sodium-to-creatinine ratio in the participants with BMI < 25 kg/m², never smokers, or current alcohol drinkers. We identified food groups contributing to urinary sodium-to-creatinine ratio. Intake of vegetables, kimchi, and seaweed were identified to explain the variability of urinary sodium-to-creatinine ratio and were positively associated with increased blood pressure.

Collecting spot urine samples reduces the low compliance problem in large population study. Therefore, large epidemiologic studies often used 12-h urine replaced for 24-h urine and several studies suggested that 12-h urine sodium excretion may reflect dietary sodium intake [33]. For example, estimated correlation coefficient between mean 24-h and mean overnight sodium excretion was 0.72 in 142 male participants [27,34].

Our findings of positive association between urinary sodium-to-creatinine ratio and blood pressure are consistent with previous studies, which have reported a positive association between urinary sodium concentration and blood pressure in general populations. In a cross-sectional, community-based study, the 24-h urine sodium-to-potassium ratio was linearly associated with nighttime systolic blood pressure (beta-coefficient = 0.142 in linear regression) and diastolic blood pressure (beta-coefficient = 0.144) in 217 participants aged ≥ 55 years [23]. In the Korea National Health and Nutrition Examination Survey (KNHANES) 2009–2010, systolic blood pressure levels were 122.9 ± 17.4 mmHg in the top quartile but 114.3 ± 13.3 mmHg in the bottom quartile of urinary sodium-to-creatinine ratio assessed from spot urine analysis in men [35]. In the Korea Genome and Epidemiology Study (KoGES) cohort, KoGES Ansan and Ansung study, the calculated 24-h urinary sodium levels had positively linear association with systolic and diastolic blood pressure levels (P for trend < 0.001) [22]. The Norfolk cohort of the European prospective investigation into cancer (EPIC–Norfolk) study showed that high urinary sodium-to-creatinine ratio measured from casual urine samples was positively associated with the systolic and diastolic blood pressure [36]. Differences in systolic and diastolic blood pressures were 7.2 mmHg and 3.0 mmHg between the top and bottom quintiles of urinary sodium-to-creatinine ratio, respectively. The Melbourne Collaborative Cohort Study collected 24-h urine samples from 587 participants and found that an increment of urinary sodium of 100mmol/day was associated with 2.3 mmHg increase in systolic blood pressure [37]. In a meta-analysis of randomized trials, a reduction of 100 mmol/day in salt intake led to a decrease in systolic/diastolic blood pressure by 7.11/3.88 mmHg in hypertensive individuals and 3.57/1.66 mmHg in normotensive individuals [38]. The possible mechanism by which high urinary sodium increases blood pressure may be related to renin–angiotensin system and renal function. Excessive sodium retention leads to decrease in kidney capacity in maintaining osmotic pressure of the plasma, interstitial fluid volumes, acid–base balance, and electrical activity of cells, thus resulting in increased blood pressure [39].

Several epidemiologic and intervention studies reported an inverse association between potassium intake or urinary potassium excretion and blood pressure. The INTERSALT study observed that systolic and diastolic blood pressure levels increased by 3.4 mmHg and 1.9 mmHg, respectively, according to 50 nmol per day of 24-h urinary potassium excretion among 10,079 participants [15]. A recent meta-analysis of randomized-controlled trials, where participants in intervention arm were provided with potassium supplements for 4 to 24 weeks, found reduction in systolic blood pressure by 4.7 mmHg and diastolic blood pressure by 3.5 mmHg [25]. Activation of sodium-chloride cotransporter has been suggested as a potential mechanism through which low potassium increases blood pressure. Low potassium stimulates the activation of sodium-chloride cotransporter in the kidney, which promotes the sodium retention and increases blood pressure [40]. Also, sodium-hydrogen exchanger type 3 in kidney may be enhanced by potassium depletion, resulting in increase in sodium retention [41].
The reasons for the significant interaction by BMI were not clear. The US National Health and Nutrition Examination Survey (NHANES) Epidemiologic Follow-up Study reported increase in cardiovascular disease mortality only among overweight individuals [42]. However, the subgroup analysis of DASH diet did not find any difference in low sodium effect on SBP by obesity [43]. The association between sodium intake and blood pressure warrants further investigation.

RRR, a new method of dietary pattern analysis, identifies linear combinations of dietary intake variables that explain the variance in a set of intermediated response variables [32]. We determined food groups contributing to urinary sodium-to-creatinine ratio using RRR and the associations of these food groups with systolic or diastolic blood pressures. Intakes of vegetables, soy paste, kimchi, and seaweed were associated with increased systolic and diastolic blood pressures. Kimchi appeared to be top food contributing to high sodium intake in Korean populations [44]. One of the most common ways to cook vegetables is to blanch them with soy sauce or salt. Roasted laver is commonly seasoned with salt and sesame oil in Korea, and dried kelp has high salts content. Significant association with systolic or diastolic blood pressure observed for RRR-derived sodium score may suggest a possible association between sodium intake and hypertension.

There are some limitations in our study. We did not take into account dietary intake of sodium or correlation between urinary sodium intake and dietary sodium intake. Measurement error in estimating food intake from FFQs could attenuate to some extent the associations for food groups contributing to urinary sodium levels. Our cross-sectional study design may not infer the temporal relationship between urinary sodium and blood pressure. Further prospective studies are warranted to replicate the association we observed. The strengths of our study include a large number of participants and measurements of overnight urine for over 2500 participants.

5. Conclusions

Our study included a large number of participants in Korean population whose sodium intake is high and found that sodium intake was associated with increased levels of blood pressure. Although the association between sodium intake and blood pressure has been supported by several epidemiologic studies, urinary sodium, a better indicator for sodium intake than dietary intake, has not been well studied especially in Asian populations. Our study supports that even in population with high sodium intake, blood pressure increases with sodium intake.

Our study suggests that sodium intake and food groups with high sodium levels lead to high urinary sodium concentration and are associated with increasing blood pressure among Korean healthy adults.

Supplementary Materials: The following are available online at http://www.mdpi.com/2072-6643/12/11/3442/s1, Table S1: Baseline characteristics according to the quintiles of sodium-to-creatinine ratio, potassium-to-creatinine ratio or sodium-to-potassium ratio in men and women, Table S2: Multivariate-adjusted geometric means (95% CIs) of systolic and diastolic blood pressures (mmHg) according to urinary sodium-to-creatinine, potassium-to-creatinine, or sodium-to-potassium ratios in men and women, Table S3: Multivariate-adjusted geometric means (95% CIs) of systolic and diastolic blood pressures (mmHg) according to urinary sodium-to-creatinine and potassium-to-creatinine ratios by eGFR, urinary potassium level, education level, and married status, Table S4: Multivariate-adjusted geometric means (95% CIs) of systolic and diastolic blood pressures (mmHg) according to sodium contributing food score by BMI category, Table S5: Factor loading values of food groups contributing to urinary potassium-to-creatinine ratio through reduced rank regression analysis, Table S6: Multivariate-adjusted geometric means (95% CIs) of systolic and diastolic blood pressures (mmHg) according to potassium-contributing food score.

Author Contributions: Conceptualization, J.S. and J.E.L.; formal analysis, D.Y.S., J.Y. and K.K.; investigation, J.S. and J.E.L.; writing—original draft preparation, D.Y.S., J.Y., J.S. and J.E.L.; writing—review and editing, D.Y.S., J.Y., K.K., J.S. and J.E.L.; funding acquisition, J.S. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Acknowledgments: The authors thank all participants who volunteered for Healthy Twin Study and staff for supporting the cohort study.
Conflicts of Interest: The authors declare no conflict of interest.

References

1. Kannel, W.B. Blood pressure as a cardiovascular risk factor: Prevention and treatment. *JAMA* 1996, 275, 1571–1576. [CrossRef] [PubMed]
2. Grossman, E.; Messerli, F.H.; Boyko, V.; Goldbourt, U. Is there an association between hypertension and cancer mortality? *Am. J. Med.* 2002, 112, 479–486. [CrossRef]
3. Liang, Z.; Xie, B.; Li, J.; Wang, X.; Wang, S.; Meng, S.; Ji, A.; Zhu, Y.; Xu, X.; Zheng, X.; et al. Hypertension and risk of prostate cancer: A systematic review and meta-analysis. *Sci. Rep.* 2016, 6, 31358. [CrossRef]
4. Han, H.; Guo, W.; Shi, W.; Yu, Y.; Zhang, Y.; Ye, X.; He, J. Hypertension and breast cancer risk: A systematic review and meta-analysis. *Sci. Rep.* 2017, 7, 44877. [CrossRef] [PubMed]
5. Fuller, J.H. Epidemiology of hypertension associated with diabetes mellitus. *Hypertension* 1985, 7, II3–II7. [CrossRef]
6. Afshin, A.; Sur, P.J.; Fay, K.A.; Cornaby, L.; Ferrara, G.; Salama, J.S.; Mullany, E.C.; Abate, K.H.; Abbafati, C.; Abebe, Z.; et al. Health effects of dietary risks in 195 countries, 1990–2017: A systematic analysis for the global burden of disease study 2017. *Lancet* 2019, 393, 1958–1972. [CrossRef]
7. Appel, L.J.; Moore, T.J.; Obarzanek, E.; Vollmer, W.M.; Svetkey, L.P.; Sacks, F.M.; Bray, G.A.; Vogt, T.M.; Cutler, J.A.; Windhauser, M.M.; et al. A clinical trial of the effects of dietary patterns on blood pressure. *Dash Collab. Res. Group N. Engl. J. Med.* 1997, 336, 1117–1124. [CrossRef]
8. Appel, L.J. Lifestyle modification as a means to prevent and treat high blood pressure. *J. Am. Soc. Nephrol.* 2003, 14, S99–S102. [CrossRef]
9. Sacks, F.M.; Svetkey, L.P.; Vollmer, W.M.; Appel, L.J.; Bray, G.A.; Harsha, D.; Obarzanek, E.; Conlin, P.R.; Miller, E.R., 3rd; Simons-Morton, D.G.; et al. Effects on blood pressure of reduced dietary sodium and the dietary approaches to stop hypertension (Dash) diet. *Dash-Sodium Collab. Res. Group N. Engl. J. Med.* 2001, 344, 3–10. [CrossRef]
10. Korea Health Statistics 2018: Korea National Health and Nutrition Examination Survey (Knhanes vii-3); Korea Centers for Disease Control and Prevention, 2019. Available online: https://knhanes.cdc.go.kr/knhanes/sub04/sub04_03.do?classType (accessed on 9 November 2020).
11. Fryar, C.D.; Ostchega, Y.; Hales, C.M.; Zhang, G.; Kruszon-Moran, D. Hypertension prevalence and control among adults: United states, 2015–2016. *NCHS Data Brief.* 2017, 289, 1–8.
12. Loria, C.M.; Obarzanek, E.; Ernst, N.D. Choose and prepare foods with less salt: Dietary advice for all americans. *J. Nutr.* 2001, 131, 536S–551S. [CrossRef] [PubMed]
13. Kim, Y.S.; Paik, H.Y. Measurement of na intake in korean adult females. *Korean J. Nutr.* 1987, 20, 341–349.
14. Ji, C.; Sykes, L.; Paul, C.; Dary, O.; Legetic, B.; Campbell, N.R.; Cappuccio, F.P. Systematic review of studies comparing 24-hour and spot urine collections for estimating population salt intake. *Rev. Panam. de Salud Pública* 2012, 32, 307–315. [CrossRef] [PubMed]
15. Intersalt Cooperative Research Group. Intersalt: An international study of electrolyte excretion and blood pressure. Results for 24 hour urinary sodium and potassium excretion. Intersalt cooperative research group. *BMJ* 1988, 297, 319–328. [CrossRef] [PubMed]
16. Jackson, S.L.; Cogswell, M.E.; Zhao, L.; Terry, A.L.; Wang, C.Y.; Wright, J.; Coleman King, S.M.; Bowman, B.; Chen, T.C.; Merritt, R.; et al. Association between urinary sodium and potassium excretion and blood pressure among adults in the united states: National health and nutrition examination survey, 2014. *Circulation* 2018, 137, 237–246. [CrossRef] [PubMed]
17. Welsh, C.E.; Welsh, P.; Jhund, P.; Delles, C.; Celis-Morales, C.; Lewsey, J.D.; Gray, S.; Lyall, D.; Iliodromiti, S.; Gill, J.M.R.; et al. Urinary sodium excretion, blood pressure, and risk of future cardiovascular disease and mortality in subjects without prior cardiovascular disease. *Hypertension* 2019, 73, 1202–1209. [CrossRef]
18. Dolson, G.M.; Ellis, K.J.; Bernardo, M.V.; Prakash, R.; Adrogué, H.J. Acute decreases in serum potassium augment blood pressure. *Am. J. Kidney Dis.* 1995, 26, 321–326. [CrossRef]
19. Oh, J.; Lee, J.; Koo, H.S.; Kim, S.; Chin, H.J. Estimated 24-hour urine sodium excretion is correlated with blood pressure in korean population: 2009–2011 korean national health and nutritional examination survey. *J. Korean Med Sci.* 2014, 29 (Suppl. 2), S109–S116. [CrossRef]
20. Koo, H.S.; Kim, Y.C.; Ahn, S.Y.; Oh, S.W.; Kim, S.; Chin, H.J. Analysis of correlation between 24-hour urinary sodium and the degree of blood pressure control in patients with chronic kidney disease and non-chronic kidney disease. *J. Korean Med Sci.* 2014, 29 (Suppl. 2), S117–S122. [CrossRef]

21. Shin, J.; Xu, E.; Lim, Y.H.; Choi, B.Y.; Kim, B.K.; Lee, Y.G.; Kim, M.K.; Mori, M.; Yamori, Y. Relationship between nocturnal blood pressure and 24-h urinary sodium excretion in a rural population in Korea. *Clin. Hypertens.* 2014, 20, 9. [CrossRef]

22. Park, Y.M.; Kwock, C.K.; Kim, K.; Kim, J.; Yang, Y.J. Interaction between single nucleotide polymorphism and urinary sodium, potassium, and sodium-potassium ratio on the risk of hypertension in Korean adults. *Nutrients* 2017, 9, 235. [CrossRef] [PubMed]

23. Kim, M.K.; Kwon, M.; Rhee, M.Y.; Kim, K.I.; Nah, D.Y.; Kim, S.W.; Gu, N.; Sung, K.C.; Hong, K.S.; Cho, E.J.; et al. Dose-response association of 24-hour urine sodium and sodium to potassium ratio with nighttime blood pressure at older ages. *Eur. J. Prev. Cardiol.* 2019, 26, 952–960.

24. Kumanyika, S.K.; Cook, N.R.; Cutler, J.A.; Belden, L.; Brewer, A.; Cohen, J.D.; Hebert, P.R.; Lasser, V.I.; Raines, J.; Raczyński, J.; et al. Sodium reduction for hypertension prevention in overweight adults: Further results from the trials of hypertension prevention phase II. *J. Hum. Hypertens.* 2005, 19, 33–45. [CrossRef] [PubMed]

25. Binia, A.; Jaeger, J.; Hu, Y.; Singh, A.; Zimmermann, D. Daily potassium intake and sodium-to-potassium ratio in the reduction of blood pressure: A meta-analysis of randomized controlled trials. *J. Hypertens.* 2015, 33, 1509–1520. [CrossRef] [PubMed]

26. Sung, J.; Cho, S.I.; Lee, K.; Ha, M.; Choi, E.Y.; Choi, J.S.; Kim, H.; Kim, J.; Hong, K.S.; Kim, Y.; et al. Healthy twin: A twin-family study of Korea—Protocols and current status. *Twin Res. Hum. Genet.* 2006, 9, 844–848. [CrossRef]

27. Liu, K.; Dyer, A.R.; Cooper, R.S.; Stamler, R.; Stamler, J. Can overnight urine replace 24-hour urine collection to assess salt intake? *Hypertension* 1979, 1, 529–536. [CrossRef]

28. Kho, M.; Lee, J.E.; Song, Y.-M.; Lee, K.; Kim, K.; Yang, S.; Joung, H.; Sung, J. Genetic and environmental influences on sodium intake determined by using half-day urine samples: The healthy twin study. *The Am. J. Clin. Nutr.* 2013, 98, 1410–1416. [CrossRef]

29. Charlton, K.; Ware, L.J.; Chidumwa, G.; Cockeran, M.; Schutte, A.E.; Naidoo, N.; Kowal, P. Prediction of 24-hour sodium excretion from spot urine samples in South African adults: A comparison of four equations. *J. Hum. Hypertens.* 2020, 34, 24–33. [CrossRef]

30. Ahn, Y.; Kwon, E.; Shim, J.E.; Park, M.K.; Joo, Y.; Kimm, K.; Park, C.; Kim, D.H. Validation and reproducibility of food frequency questionnaire for Korean genome epidemiologic study. *Eur. J. Clin. Nutr.* 2007, 61, 1435–1441. [CrossRef]

31. Levey, A.S.; Stevens, L.A.; Schmid, C.H.; Zhang, Y.L.; Castro, A.F., 3rd; Feldman, H.L.; Kusek, J.W.; Eggers, P.; Van Lente, F.; Greene, T.; et al. A new equation to estimate glomerular filtration rate. *Ann. Intern. Med.* 2009, 150, 604–612. [CrossRef] [PubMed]

32. Hoffmann, K.; Schulze, M.B.; Schienkiewitz, A.; Nothlings, U.; Boeing, H. Application of a new statistical method to derive dietary patterns in nutritional epidemiology. *Am. J. Epidemiol.* 2004, 159, 935–944. [CrossRef] [PubMed]

33. Watson, R.L.; Langford, H.G. Usefulness of overnight urines in population groups. *Pilot studies of sodium, potassium, and calcium excretion. Am. J. Clin. Nutr.* 1970, 23, 290–304. [CrossRef] [PubMed]

34. Hu, Y.; Zong, G.; Liu, G.; Wang, M.; Rosner, B.; Pan, A.; Willett, W.C.; Manson, J.E.; Hu, F.B.; Sun, Q. Smoking Cessation, Weight Change, Type 2 Diabetes, and Mortality. *New Engl. J. Med.* 2018, 379, 623–632. [CrossRef] [PubMed]

35. Lee, S.-G.; Lee, W.; Kwon, O.H.; Kim, J.-H. Association of urinary sodium/creatinine ratio and urinary sodium/specific gravity unit ratio with blood pressure and hypertension: KNHANES 2009–2010. *Clin. Chim. Acta* 2013, 424, 168–173. [CrossRef]

36. Khaw, K.T.; Bingham, S.; Welch, A.; Luben, R.; O’Reilly, S.; Wareham, N.; Day, N. Blood pressure and urinary sodium in men and women: The Norfolk Cohort of the European Prospective Investigation into Cancer (EPIC-Norfolk). *Am. J. Clin. Nutr.* 2004, 80, 1397–1403. [CrossRef] [PubMed]

37. E Huggins, C.; O’Reilly, S.; Brinkman, M.; Hodge, A.; Giles, G.G.; English, D.R.; Nowson, C.A. Relationship of urinary sodium and sodium-to-potassium ratio to blood pressure in older adults in Australia. *Med J. Aust.* 2011, 195, 128–132. [CrossRef]
38. He, F.J.; A MacGregor, G. Effect of modest salt reduction on blood pressure: A meta-analysis of randomized trials. *Implications for public health. J. Hum. Hypertens.* **2002**, *16*, 761–770. [CrossRef]

39. World Health Organization. *The Global Burden of Disease: 2004 Update*; WHO: Geneva, Switzerland, 2008.

40. Vallon, V.; Schroth, J.; Lang, F.; Kuhl, D.; Uchida, S. Expression and phosphorylation of the Na+-Cl− cotransporter NCC in vivo is regulated by dietary salt, potassium, and SGK1. *Am. J. Physiol. Physiol.* **2009**, *297*, F704–F712. [CrossRef]

41. Soleimani, M.; A Bergman, J.; A Hosford, M.; McKinney, T.D. Potassium depletion increases luminal Na+/H+ exchange and basolateral Na+:CO3−:HCO3− cotransport in rat renal cortex. *J. Clin. Investig.* **1990**, *86*, 1076–1083. [CrossRef]

42. Whelton, P.K. Dietary Sodium Intake and Subsequent Risk of Cardiovascular Disease in Overweight Adults. *JAMA* **1999**, *282*, 2027–2034.

43. Vollmer, W.M.; Sacks, F.M.; Ard, J.; Appel, L.J.; Bray, G.A.; Simons-Morton, D.G.; Conlin, P.R.; Svetkey, L.P.; Erlinger, T.P.; Moore, T.J.; et al. Effects of Diet and Sodium Intake on Blood Pressure: Subgroup Analysis of the DASH-Sodium Trial. *Ann. Intern. Med.* **2001**, *135*, 1019–1028. [CrossRef] [PubMed]

44. Song, D.Y.; Park, J.E.; Shim, J.E.; Song, S. Trends in the major dish groups and food groups contributing to sodium intake in the Korea National Health and Nutrition Examination Survey 1998–2010. *Korean J. Nutr.* **2013**, *46*, 72–85. [CrossRef]

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