Dietary Salt Intake and Mortality in Patients With Type 2 Diabetes

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OBJECTIVE—Many guidelines recommend that patients with type 2 diabetes should aim to reduce their intake of salt. However, the precise relationship between dietary salt intake and mortality in patients with type 2 diabetes has not been previously explored.

RESEARCH DESIGN AND METHODS—Six hundred and thirty-eight patients attending a single diabetes clinic were followed in a prospective cohort study. Baseline sodium excretion was estimated from 24-h urinary collections ($24hU_{Na}$). The predictors of all-cause and cardiovascular mortality were determined by Cox regression and competing risk modeling, respectively.

RESULTS—The mean baseline $24hU_{Na}$ was 184 ± 73 mmol/24 h, which remained consistent throughout the follow-up (intraindividual coefficient of variation [CV] $23 \pm 11\%$). Over a median of 9.9 years, there were 175 deaths, 75 (43%) of which were secondary to cardiovascular events. All-cause mortality was inversely associated with $24hU_{Na}$, after adjusting for other baseline risk factors ($P < 0.001$). For every 100 mmol rise in $24hU_{Na}$, all-cause mortality was 28% lower (95% CI 6–45%, $P = 0.02$). After adjusting for the competing risk of noncardiovascular death and other predictors, $24hU_{Na}$ was also significantly associated with cardiovascular mortality (sub-hazard ratio 0.65 [95% CI 0.44–0.95]; $P = 0.03$).

CONCLUSIONS—In patients with type 2 diabetes, lower 24-h urinary sodium excretion was paradoxically associated with increased all-cause and cardiovascular mortality. Interventional studies are necessary to determine if dietary salt has a causative role in determining adverse outcomes in patients with type 2 diabetes and the appropriateness of guidelines advocating salt restriction in this setting.

I n patients with type 2 diabetes, hypertension is associated with a range of adverse outcomes including cardiovascular disease (CVD) and premature mortality. Consequently, clinical guidelines recommend that patients with type 2 diabetes undertake measures to maintain a blood pressure at or below target levels. Among the interventions advocated to assist in achieving these targets, most guidelines recommend a reduced intake of salt, as dietary sodium intake is positively correlated with blood pressure levels in the general population (1).

Moreover, in patients with type 2 diabetes, salt restriction confers a modest reduction in blood pressure (2), and salt supplementation reduces the antihypertensive efficacy of blood pressure–lowering agents in the short term (3). However, the precise relationship between salt intake and mortality in patients with type 2 diabetes has not been previously explored. It is widely assumed that any blood pressure lowering associated with reduced dietary salt intake may be translated into protection from end-organ damage in the context of diabetes.

However, there is also evidence that reduced sodium intake is associated with activation of metabolic and neurohormonal pathways, including the sympathetic nervous system (4) and the renin-angiotensin-aldosterone system (RAAS) (4), as well as increases in total and LDL cholesterol (4) and reduced peripheral insulin sensitivity (5). In the context of type 2 diabetes, each of these factors may offset or even outweigh gains achieved from blood pressure lowering. Hence, in this study we explored the association between dietary salt intake, the best estimate of which is 24-h urine collection as ~90% of dietary sodium intake is renally excreted (6), and all-cause and cardiovascular mortality in patients with type 2 diabetes.

RESEARCH DESIGN AND METHODS

Patient recruitment
This study was initiated in July 2000 as a prospective survey of patients with type 2 diabetes who were in long-term follow-up in a single diabetes clinic at Austin Health, Melbourne, Australia. Austin Health is a major university teaching hospital and a tertiary referral center, serving a population of approximately 700,000. Long-term follow-up was defined by the patients having at least three previous estimations of urinary albumin excretion rate (AER) performed on 24-h urine collections with at least one AER having been performed within the year 2000. Patients with type 1 diabetes or diabetes secondary to medication or pancreatitis were specifically excluded from the current study. Using these criteria, 665 patients with type 2 diabetes were eligible to participate in the survey. Informed consent was obtained from participating patients as approved by the Austin Health Human Research Ethics Committee.

Measurement of baseline characteristics
Baseline data collection was performed during routine clinical visits between January and December 2001. Baseline clinical and biochemical characteristics of all participants were ascertained including a full clinical history, medication use, anthropometric data, and smoking habits. The presence of preexisting CVD was

defined on the basis of a clinical history of myocardial infarction, unstable angina requiring hospitalization, coronary revascularization (including coronary artery bypass grafting, angioplasty, or coronary stenting), heart failure, stroke, carotid artery surgery, peripheral revascularization (including bypass grafting, angioplasty, or stenting), and amputation for critical limb ischemia. Fasting blood samples were taken for glucose, HbA_{1c}, lipid profiles, and creatinine. The glomerular filtration rate was estimated (eGFR) using the chronic kidney disease-epidemiology collaboration (CKD-EPI) formula.

All regularly attending patients at the Austin Health diabetes clinics perform a 24-h urine collection before each visit for the determination of urinary sodium, potassium, and albumin excretion using standard methods in the Department of Laboratory Medicine at Austin Health. Baseline levels for each participant were determined as the mean excretion rates across all urine samples performed during 2001. Data on $24hU_{Na}$ were available in 638 participants (96%) who formed the study cohort (Supplementary Figure 1). The clinical characteristics of these patients were not different from those in whom $24hU_{Na}$ was not measured.

Patients were given general dietary advice as part of their routine care at an initial assessment by a dietitian. However detailed assessment of dietary salt intake was not performed. During follow-up, all patients continued to have standard medical care including antihypertensive, lipid-lowering, and antidiabetic medications according to recommended guidelines.

Mortality outcomes

The primary outcome was death from any cause until the 31 March 2010. Timing and cause of death were identified from a search of hospital patients' records and the Australian Institute of Health and Welfare Death Registry. To evaluate the independent predictors of all-cause mortality in patients with type 2 diabetes, we used Cox proportional hazards models. The final model variables were determined by sequential penalized likelihood (Akaike information criterion) (7). The functional form (in particular, nonlinearity) of continuous variables in the final model was explored by both fractional polynomials and restricted cubic splines (7). Categorical variables were parameterized as simple indicator variables; the potential for multicollinearity was assessed using the variance inflation factor and condition number. Overall Cox model fit was assessed by approximation of cumulative Cox-Snell residuals to (-log) Kaplan-Meier estimates, residual plots, and testing of the proportional hazards assumption and goodness-of-fit test (8). The cumulative hazard of all-cause mortality was graphically displayed using Nelson-Aalen estimates stratified by percentiles of covariate(s) of interest (9).

Table 1—Baseline characteristics of patients with type 2 diabetes, stratified according to tertiles (T) of 24-h urinary sodium excretion

Baseline parameter	T1 <150 mmol/24 h	T2 150–208 mmol/24 h	T3 >208 mmol/24 h
Age (years)	67 ± 12*	64 ± 11	61 ± 12*
Sex (% male)	42*	56	70*
Diabetes duration (years)	14 ± 9*	12 ± 8	11 ± 8
Obese (BMI >30 kg/m^2) (%)	41	45	55*
Macrovascular disease (%)	49	43	44
Coronary heart disease (%)	34	32	37
Atrial fibrillation (%)	20	20	12*
Congestive cardiac failure (%)	17	11	15
C-reactive protein (geometric mean; IU/mL)	2.4	2.1	2.6
Systolic blood pressure (mmHg)	141 ± 17	140 ± 17	140 ± 16
Diastolic blood pressure (mmHg)	77 ± 10*	80 ± 9	78 ± 10*
Antihypertensive therapy (%)	78	78	76
ACE inhibitor (%)	45*	56	57
Angiotensin receptor blocker (%)	13	13	10
Diuretic (%)	38	32	42
β-Blocker (%)	23*	15	20
Calcium channel blocker (%)	34	28	34
α-Blocker (%)	5	5	5
HbA_{1c} (%)	7.8 ± 1.7	7.8 ± 1.3	7.7 ± 1.5
Fasting plasma glucose (mmol/L)	9.6 ± 4.0	9.3 ± 3.2	9.3 ± 3.7
Metformin (%)	50	57	58
Sulfonylurea (%)	37*	51	40*
Insulin (%)	49%*	36%	39%
LDL cholesterol (mmol/L)	2.6 ± 0.8	2.7 ± 0.8	2.6 ± 0.8
Triglycerides (mmol/L)	2.0 ± 1.8	2.0 ± 1.4	2.2 ± 1.6
HDL cholesterol (mmol/L)	1.2 ± 0.4	1.2 ± 0.3	1.1 ± 0.3
Statin (%)	53	52	49
eGFR (mL/min/1.73 m^2)	67 ± 26*	73 ± 25	78 ± 24*
Albuminuria (micro and macro, %)	42	42	50
Hemoglobin (g/dL)	13.2 ± 1.5	13.4 ± 1.5	13.7 ± 1.5*

Data are mean ± SD unless otherwise indicated. *$P < 0.01$ vs. middle tertile (T2).

To identify the independent predictors of the cumulative incidence of cardiovascular mortality, we used the Fine and Gray model, which extends the Cox proportional hazards model to competing risks data (10) by considering the subdistribution hazard while adjusting for the competing risk of noncardiovascular deaths. The strength of the association between each variable and the outcome was assessed using the sub-hazard ratio, which is the ratio of hazards associated with the cumulative incidence function in the presence and absence of a prognostic factor. The Fine and Gray model was implemented in Stata statistical software (V11, 2009; College Station, TX) using the stcrreg module.

RESULTS

Cohort characteristics

The mean age of participants was 64 years, 56% were men, the median duration of diabetes was 11 years, and 47%

were obese (BMI >30 kg/m^2). Complications of diabetes were common with 45% of participants having previously experienced a cardiovascular event, including 35% who had a myocardial infarction. One-third had background retinopathy on ophthalmological review (33%), 30% had an eGFR <60 mL/min/1.73 m^2, and 45% had elevated urinary albumin excretion (>20 μg/min), including 13% with macroalbuminuria (>200 μg/min). Eighty-five percent of patients had hypertension (defined by the use of antihypertensives and/or blood pressure >140/90). Achievement of therapeutic targets was low, reflecting the tertiary referral nature of the clinic, with two-thirds of patients having an HbA$_{1c}$ >7.0%, 55% with an LDL cholesterol level >2.5 mM, and 44% with a systolic blood pressure >140 mmHg.

24-h urinary sodium excretion

Baseline urinary sodium excretion was estimated in 638 patients from a median

of two collections performed during the year 2001 (range 1–5). The mean urinary sodium excretion was 184 mmol/24 h, similar to previous global population surveys (11). One-third of participants had a sodium excretion <150 mmol/24 h (lowest tertile), and one-third had an excretion >208 mmol/24 h (highest tertile) with the middle tertile falling between 150 and 208 mmol/24 h. Patients with urinary sodium excretion in the highest tertile were younger, with a shorter duration of diabetes, more likely to be obese, but with better renal function, and hemoglobin, when compared with patients in tertile 2 (Table 1). Sex was also a strong predictor of sodium excretion, consistent with the higher sodium intake described in men in global population surveys (11). On multivariate analysis, 24hU$_{Na}$ (as a continuous variable) was positively correlated with eGFR, BMI, male sex, and the use of ACE inhibitors and was negatively correlated with age (full regression model [Supplementary Table 1]). The presence
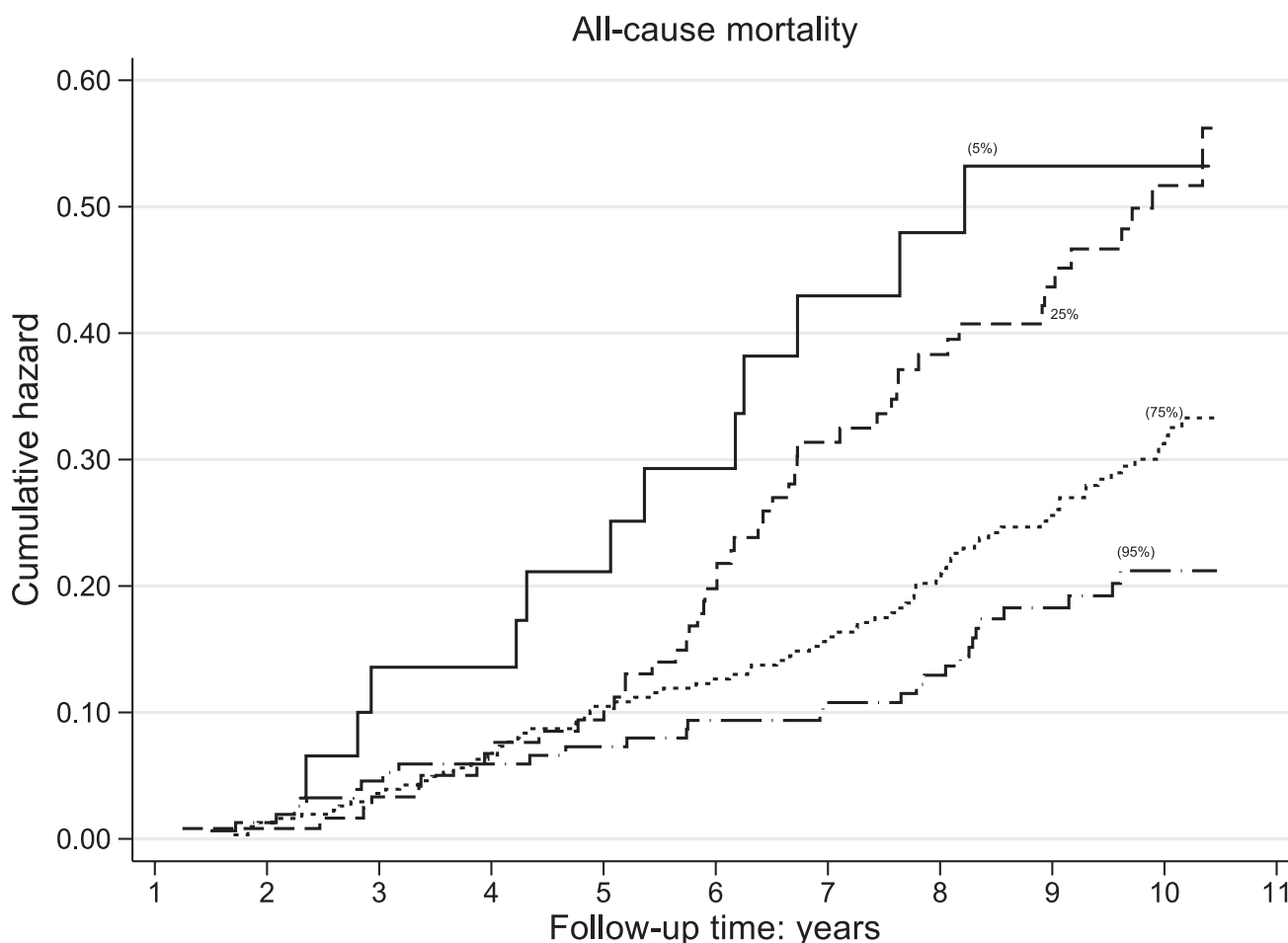
Figure 1—*Cumulative hazard (Nelson-Aalen) of all-cause mortality, stratified by percentiles (5th, 25th, 75th, and 95th) of 24-h urinary sodium excretion. All-cause mortality was inversely associated with 24-h urinary sodium excretion.*

or absence of heart failure, atrial fibrillation, and the use of diuretics or β-blockers was not associated with $24hU_{Na}$ after adjusting for age and sex.

Variability in sodium excretion

Baseline $24hU_{Na}$ excretion was strongly correlated with the yearly mean $24hU_{Na}$ during follow-up. Over two-thirds of individuals with $24hU_{Na}$ in the highest tertile at baseline remained in the highest tertile in subsequent years. Similarly, over two-thirds of individuals with $24hU_{Na}$ in the lowest tertile remained in the lowest tertile in subsequent years (data not shown). Less than 5% of individuals changed from the lowest to the highest tertile or the highest to the lowest tertile on an annual basis. Overall, the intraindividual CV of mean annual $24hU_{Na}$ was $23 \pm 11\%$ over 8 years.

Mortality and 24-h urinary sodium excretion

Vital status was determined in 620 patients with the remaining 18 (3%) lost to follow-up. The median follow-up for survival analysis was 9.9 years (5,475 patient-years) during which 175 deaths were recorded (28%). CVD was listed as a major contributing cause in 75 deaths (43% of all deaths) with the other major causes of death including cancer ($n = 36$) and infection ($n = 19$). In 17 patients, although death was verified, no cause of death could be ascertained.

All-cause mortality was inversely associated with $24hU_{Na}$ such that individuals with the lowest $24hU_{Na}$ had the highest cumulative hazard for mortality (Fig. 1). After adjusting for parameters associated with $24hU_{Na}$, as well as other factors independently associated with all-cause mortality (including age, sex, duration of diabetes, atrial fibrillation, the presence and severity of CKD (eGFR and log AER) for every 100 mmol rise in 24hUNa, all-cause mortality was 28% lower (95% CI 6–45%, $P = 0.02$) (Table 2). Overall, $24hU_{Na}$ explained 7% of the variability ($R^2 = 0.07$, 95% CI 0.02–0.14) in all-cause mortality in this cohort. No

significant interactions between categorical variables or between categorical and continuous variables were demonstrated in the model implying the relative hazard associated with sodium intake was uniform regardless of the presence and severity of hypertension, CVD, CKD, and other risk factors. Moreover, its effect was stronger than for other reversible risk factors including LDL cholesterol and HbA_{1c}, and was equivalent to that of systolic blood pressure (Table 2). Interestingly, blood pressure levels were inversely correlated with all-cause mortality with no convincing evidence of nonlinear effects including J-curve associations.

Cardiovascular mortality was also significantly associated with $24hU_{Na}$ (Fig. 2) after adjusting for the competing risk of noncardiovascular death and other predictors (Table 2). No association was observed between $24hU_{Na}$ and noncardiac deaths (data not shown). When modeled as a continuous variable, the association between $24hU_{Na}$ and all-cause and cardiovascular mortality was linear. No convincing evidence of nonlinear effects, including J-shaped associations, was demonstrated as judged by residual-by-time analysis, fractional polynomials, and (cubic) regression splines (7). Again, no significant interactions between categorical variables or between categorical and continuous variables were demonstrated, implying the relationship between $24hU_{Na}$ and mortality was independent of BMI, age, sex, and the type of antihypertensive therapy or achieved level of blood pressure control. Furthermore, the association between $24hU_{Na}$ and mortality was independent of 24-h potassium excretion (data not shown).

CONCLUSIONS—Advice to reduce dietary salt intake is a key element of many lifestyle intervention programs for type 2 diabetes. However, the association between dietary salt intake and mortality outcomes has not been previously studied specifically in the context of type 2 diabetes. In this article, we show that $24hU_{Na}$ was independently associated with all-cause and cardiovascular mortality in patients with type 2 diabetes such that the highest mortality risks were observed in individuals with the lowest sodium intake and vice versa.

Although such data may seem contrary to established dogma, in fact previous observational studies, both in the general population and in hypertensive patients, have failed to give consistent results regarding the association of salt

Table 2—Independent associations with all-cause mortality and cumulative incidence of cardiovascular mortality in individuals with type 2 diabetes

	All-cause mortality		
Baseline parameter	Hazard ratio	P	95% CI
24-h urinary sodium excretion			
(per 100 mmol/day)	0.72	0.017	0.55–0.94
Age (per decade)	1.05	<0.001	1.03–1.07
Male sex (yes/no)	1.51	0.013	1.09–2.09
Pre-existing CVD (yes/no)	1.85	0.001	1.30–2.64
eGFR (per 10 mL/min/1.73 m^2)	0.988	0.002	0.980–0.996
Atrial fibrillation (yes/no)	1.97	<0.001	1.39–2.81
Log_{10} AER	1.71	<0.001	1.38–2.12
Systolic blood pressure (mmHg)	0.986	0.015	0.974–0.997
Diabetes duration (decades)	1.02	0.010	1.01–1.04

	Cardiovascular mortality		
Baseline parameter	Sub-hazard ratio	P	95% CI
24-h urinary sodium excretion			
(per 100 mmol/day)	0.65	0.026	0.44–0.95
Male sex (yes/no)	1.93	0.011	1.17–3.20
Pre-existing CVD (yes/no)	1.88	0.014	1.14–3.11
eGFR (per 10 mL/min/1.73 m^2)	0.985	0.001	0.98–0.99
Atrial fibrillation (yes/no)	2.78	<0.001	1.71–4.53
Log_{10} AER	1.76	<0.001	1.28–2.42
Systolic blood pressure (mmHg)	0.97	<0.001	0.96–0.99
Diabetes duration (decades)	1.05	<0.001	1.02–1.08

All-cause mortality: independent associations with all-cause mortality in individuals with type 2 diabetes in a multivariate Cox model. The model explained 52% of the variation in all-cause mortality (95% CI 0.42–0.64) and was well specified (Harrell's C: 0.79; PH test: $P = 0.136$; goodness-of-fit test: $P \geq 0.37$). PH, proportional hazard. Cardiovascular mortality: independent associations with the cumulative incidence of cardiovascular mortality in individuals with type 2 diabetes in the Fine and Gray (proportional hazards) model after accounting for the competing risk of noncardiovascular death.
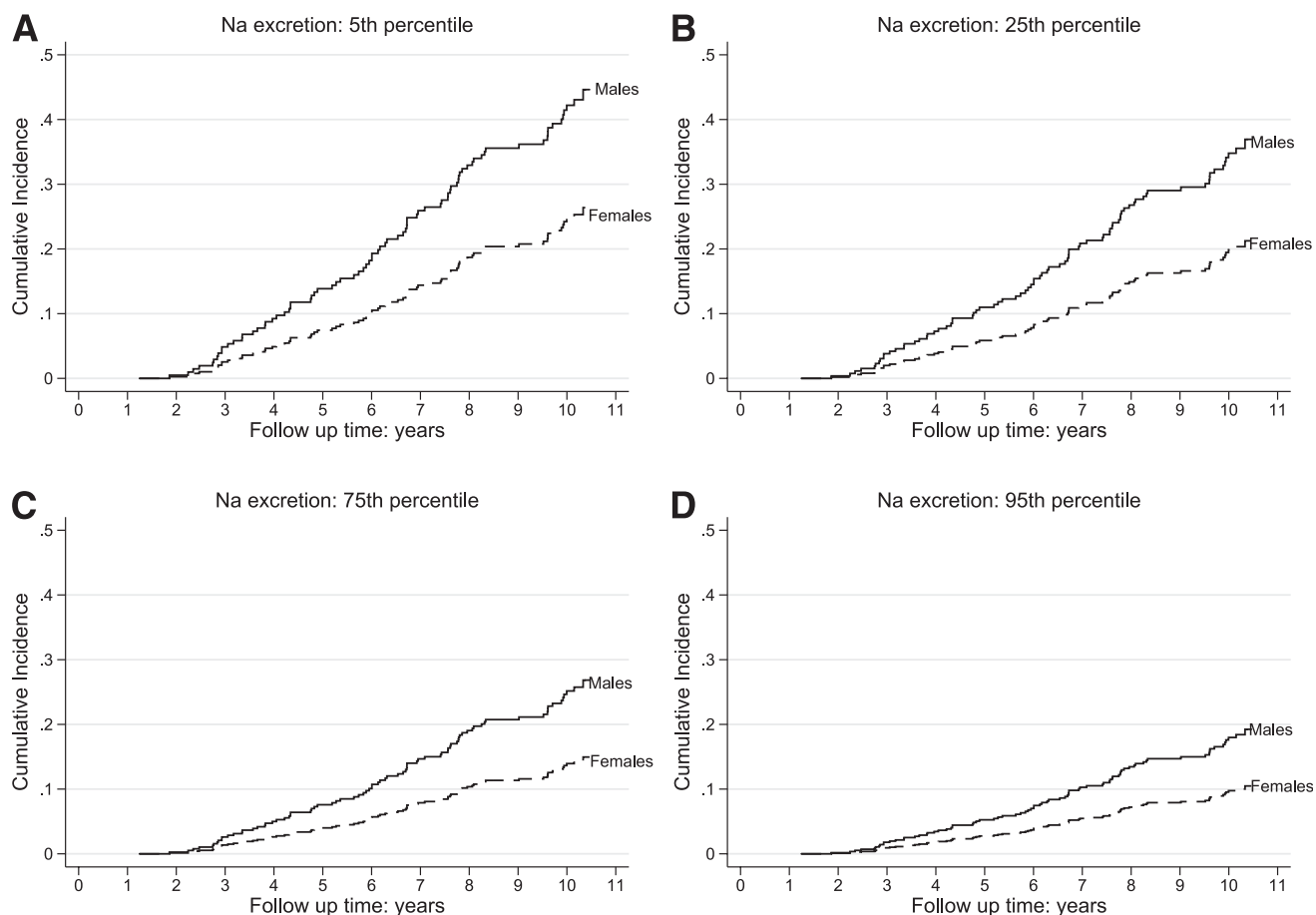
Figure 2—*The cumulative incidence (Fine and Gray) of cardiovascular mortality over the 5th, 25th, 75th, and 95th percentile (A–D, respectively) of 24-h urinary sodium excretion in men and women (solid line and dotted line, respectively), adjusted for other covariate predictors (Table 2) and accounting for noncardiovascular mortality as the competing risk. The other predictors are set at: eGFR = 76.6 mL/min/1.73 m^2 (median); atrial fibrillation = yes; preexisting cardiovascular disease = yes; Log$_{10}$ AER = 1.2 (median); systolic blood pressure = 140 mmHg (mean); diabetes duration = 10.4 years (median).*

and mortality. In some studies, higher salt intake has been associated with an increased risk of cardiovascular events (12–15), while in others—including the National Health and Nutrition Examination Surveys (NHANES) I, II, and III— lower salt intake has been associated with an increased risk of cardiovascular events and or mortality (16,17). Other studies in the general community have failed to demonstrate any association of dietary salt intake with cardiovascular events (18–20). Although hypertension and CVD are common in patients with type 2 diabetes, the issue of whether dietary salt intake influences mortality and morbidity has not been studied specifically in the context of type 2 diabetes, and any inference from outcomes in other populations should be viewed with caution.

The major strength of the current study is the use of multiple 24-h urine collections to estimate dietary sodium intake. With few exceptions (13,16), the majority of previous studies that have examined the association between salt and mortality, including NHANES (17), have relied on dietary recall, which can underestimate dietary salt intake by up to half (21). All patients were trained in urine collection and were experienced in performing 24-h urine collections at the start of the study. Moreover, in this population, the annual intraindividual CV of 24hU$_{Na}$ was low (23% over an 8-year period). In addition, the urinary sodium excretion observed was very similar to previous global population surveys (11), confirming that the patients were not sodium-restricted, which may otherwise confound analysis by indication.

The chief limitations of our data are that they are context-specific. The findings of our study specifically detail outcomes in a high-risk subpopulation of patients with longstanding type 2 diabetes attending a tertiary referral center. Nonetheless, these are precisely the patients in whom more aggressive lifestyle interventions are often applied. Our patients were mostly obese, hypertensive, with a history of poor metabolic control, and multiple comorbidities, including a significant burden of renal impairment and preexisting CVD at baseline. In all patients, cardiac risk factors were treated aggressively. It is possible that in this clinical setting, nontraditional risk factors have a stronger relationship with outcomes or that paradoxical associations are observed (22). For example, most of our patients were hypertensive despite treatment, and it has been previously shown in treated hypertensive men that low 24hU$_{Na}$ is associated with an increased risk of cardiovascular morbidity and mortality (16) and that tight blood pressure control in diabetic patients is associated with increased all-cause mortality (23).

However, in our study, the relationship between urinary sodium excretion and mortality was independent of the presence and severity of hypertension.

In our statistical analysis, we specifically assessed the nonlinear effects of predictive variables modeled for potential interactions and, in the case of cardiovascular mortality, modeled within the paradigm of a formal competing risks (Fine and Gray) model, which estimated the cumulative incidence of cardiovascular deaths while taking into consideration the competing risk of other causes of death, which may otherwise confound results (24). This strategy is especially important in patients with diabetes, as diabetes is also associated with increased noncardiovascular mortality (25), which may potentially confound cause-specific analysis. Although such strategies have a number of advantages, it is nevertheless possible that associations demonstrated in this study may be because of confounding by unmeasured factors or those that are difficult to quantify. Variability in dietary sodium intake may be associated with a range of differences in diet composition, processing, and preparation that may themselves impact on adverse outcomes in diabetic individuals. For example, sodium intake is dominated by sodium added in manufactured foods in Western diets, (~75% of intake) (11). In addition, we cannot exclude the possibility that salt appetite and cardiovascular risk are linked to a common unidentified etiological factor.

Any pathophysiological mechanisms that may underlie our observed association between mortality and salt intake in patients with type 2 diabetes remain to be established. Certainly, RAAS activity (4), insulin resistance (5), catecholamine levels (4), and lipids (4) may be influenced by sodium intake, and each of these potential mediators may be of particular relevance in the setting of type 2 diabetes and/or established atherosclerosis. For example, dietary sodium restriction leads to increased levels of angiotensin II and aldosterone, chiefly via an increase in plasma renin activity. Given the primacy of RAAS in the development and progression of diabetes complications, as adjudged by the efficacy of RAAS blockade, it is perhaps not surprising that activation of RAAS by reducing sodium intake may also be associated with adverse outcomes. The same may also be said for increased sympathetic activity, insulin resistance, and dyslipidemia associated with sodium restriction.

In summary, we show that 24hU$_{Na}$, the best marker of dietary sodium intake, was negatively associated with all-cause mortality specifically in the setting of type 2 diabetes, after adjusting for baseline risk factors. This may reflect the special status of dietary salt and the pathways it regulates in diabetic pathophysiology. Such data call into question universal recommendations that all adults should endeavor to reduce their salt intake (12). Ultimately, the explanation for our findings needs to be tested experimentally in clinical trials performed specifically in diabetic patients.

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