

# Association between lower serum sodium levels and clinical outcomes in insomnia patients

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## Research article

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# Abstract

**Background** The association of lower serum sodium levels with clinical outcomes in insomnia patients remains unclear. We explored whether lower serum sodium is associated with poor clinical outcomes in patients with insomnia.

**Methods** We retrospectively enrolled patients with a diagnosis of insomnia from January 2011 to December 2012. We divided participants into three groups according to initial serum sodium level: tertile 1 ( $< 138$  mmol/L), tertile 2 (138.0–140.9 mmol/L), and tertile 3 ( $\geq 141.0$  mmol/L). To calculate the relative risk of death, hazard ratios (HRs) and 95% confidence intervals (CIs) were obtained using Cox proportional hazard models.

**Result** A total of 412 patients with insomnia were included, of whom 13.6% ( $n = 56$ ) had hyponatremia. Patients with lower serum sodium concentrations were older and had lower hemoglobin, calcium, phosphorus, and albumin levels. At the median follow-up of 49.4 months, 44 patients had died and 62 experienced acute kidney injury (AKI). Kaplan-Meier analysis showed significantly higher mortality in patients in the lowest tertile for serum sodium. Lowest tertile of serum sodium, AKI, and chronic respiratory disease were associated with all-cause mortality. In addition, lowest tertile of serum sodium was also significantly associated with AKI.

**Conclusions** The lowest tertile of serum sodium was associated with a higher mortality and AKI rate in insomnia patients. Our results suggest serum sodium level could be used one of the prognostic factor in insomniacs and physicians should be careful to take care of them when they present lower sodium level.

## Introduction

Insomnia is a disorder characterized by at least one nocturnal sleep symptom, as well as one daytime or waking symptom attributable to poor sleep [1]. The prevalence of insomnia disorder in the general population is approximately 10–20%, with about 50% of cases having a chronic course [2]. Insomnia is not only associated with poor quality of life, but also with a risk of cognitive dysfunction [3], hypertension (HT) [4], metabolic diseases [5], and coronary artery disease (CAD) [6]. Although the pathophysiology of insomnia is complex, neurohormonal factors, sociocultural factors, and medical illness are all associated with the condition. In addition, various medical illnesses are associated with insomnia and it is important to identify precipitating factors.

Serum sodium level is important for neuronal function and osmoregulation between cells and the extracellular fluid [7]. Sodium level is the main contributor to plasma osmolality and these disorders are typically characterized by hyponatremia and hypernatremia, respectively. To maintain optimal sodium concentrations, osmoreceptors in the hypothalamus and the kidneys tightly control water homeostasis [8]. Recent studies have shown that mild hyponatremia is associated with attention deficit, gait disturbance, and falls in patients admitted to the emergency room [9, 10]. Additionally, even mild hyponatremia is believed to be associated with risk of fracture [11] and mortality in adults living in the

community [12–14]. However, the association between lower serum sodium levels and clinical outcomes in patients with insomnia remains unclear. We hypothesized that lower serum sodium level would be associated with important clinical outcomes such as overall mortality and acute kidney injury (AKI) in insomnia patients. We also investigated other factors affecting clinical outcomes in insomniacs.

## Materials And Methods

### Study population

This study retrospectively enrolled 774 insomnia adults (age  $\geq$  18 years old) patients admitted to Gyeongsang National University Hospital between January 2011 and December 2012. Patient were determined to have insomnia of the following conditions were all met, 1) ICD code G470 is listed on the discharge form as a diagnosis, and 2) patients prescribed pharmacological treatments for insomnia (benzodiazepines, benzodiazepine receptor agonists, melatonin). Data on demographic and clinical characteristics, laboratory findings, and comorbidities were obtained from the medical records at the time of admission. Patients with no available data on serum sodium level, and those who underwent renal replacement treatment, were being treated for cancer or had a history of cancer, or were lost to follow-up within 3 months were excluded.

### Definitions and clinical outcome measurements

Hyponatremia and hypernatremia were defined as a serum sodium level below 135 and above 145 mmol/L, respectively. Serum sodium level was corrected based on the serum glucose level in patients with hyperglycemia; the corrected sodium level was calculated as measured sodium + [(serum glucose – 100)  $\times$  0.016] [15]. The estimated glomerular filtration rate (eGFR) was calculated using the Modification of Diet in Renal Disease (MDRD) study formula [ $1.86 \times (\text{plasma creatinine}) - 1.154 \times (\text{age}) - 0.203$ ]  $\times$  (0.74 if female)  $\times$  (1.210 if black). Creatinine was measured using Jaffe one and serum sodium level was measured using an indirect ion-specific electrode.

Chronic kidney disease (CKD) was defined as an eGFR  $<$  60 mL/min/1.73 m<sup>2</sup>. AKI was defined as an increase in serum creatinine level  $\geq$  0.3 mg/dL within 48 hours, or an increase in serum creatinine to  $\geq$  1.5 times the baseline value, either documented or presumed to have occurred within the previous 7 days [16]. Cardiovascular disease (CVD) was defined as having a history of CAD such as angina and myocardial infarction or other CVDs include stroke, heart failure, hypertensive heart disease, rheumatic heart disease, cardiomyopathy, abnormal heart rhythms, congenital heart disease, valvular heart disease, peripheral artery disease, thromboembolic disease. Chronic respiratory disease (CRD) was defined as having a history of asthma, chronic obstructive pulmonary disease (COPD), pulmonary fibrosis, tuberculosis. Alcoholic liver disease (ALD) was defined as having liver manifestations of alcohol overconsumption such as fatty liver, alcoholic hepatitis, and chronic hepatitis with liver cirrhosis. We extracted the mortality data from medical records and Statistics Korea [17]. To evaluate differences in demographic, laboratory and clinical outcomes data among insomnia patients, we divided them into tertile groups according to serum sodium level (tertile 1:  $<$  138.0 mg/dL, tertile 2 [reference]: 138.0–

140.9 mg/dL, tertile 3:  $\geq 141.0$  mg/dL). We also compared patients divided into hyponatremia and non-hyponatremia groups. The primary outcome was all-cause mortality and the secondary outcome was AKI incidence according to tertiles of serum sodium. The study protocol was approved by the Institutional Review Board of Gyeongsang National University Hospital (IRB No. 2018-11-013-001).

## Statistical Analysis

Data are presented as the mean  $\pm$  standard deviation or frequency (count and percentage). Differences among the three tertile groups were determined using the chi-square test for categorical variables and analysis of variance (*t*-test) for continuous variables. To assess the association between serum sodium level and clinical factors, univariate and multivariate linear regression analyses were performed. To explore the association between serum sodium level and all-cause mortality, Kaplan-Meier curves were plotted for the three serum sodium groups. We also fit a restricted cubic spline function. Survival differences were compared using the log rank test. To calculate the relative risk of death, hazard ratios (HRs) and 95% confidence intervals (CIs) were derived based on Cox proportional hazards models. Factors showing a significant association ( $P < 0.10$ ) after univariate analysis, or were of clinical concern, were included in Cox proportional hazards models. Variables were selected using a backward conditional method. Statistical analyses were performed using SPSS for Windows (ver. 21.0; SPSS Inc., Chicago, IL, USA) and R software (ver. 3.2.3; R Development Core Team, Vienna, Austria). Statistical significance was defined as  $P < 0.05$ .

## Results

### Baseline characteristics according to serum sodium level

A total of 412 patients were included in the final analysis: 362 patients were excluded for various reasons. Figure 1 showed the distribution of serum sodium in the cohort. The proportion of patients with hypernatremia was much lower than that with hyponatremia. Most of the insomnia patients' serum sodium levels were within the normal range. The mean age was 61.5 years and 56.1% ( $n = 231$ ) of patients were male. The mean follow-up duration was 49.4 months. The mean serum sodium level was 138.9 mmol/L. Patients in the lowest serum sodium group (tertile 1) significantly had lower hemoglobin, calcium, phosphorus, total protein, albumin, and uric acid levels than the other two groups. Body mass index (BMI), systolic blood pressure (SBP), heart rate (HR) were not significantly different according to tertiles of serum sodium. Comorbid conditions and the number of patients taking thiazide medications did not differ among the tertiles (Table 1). The percentage of patients with hyponatremia, defined as  $<$  serum sodium 135 mmol/L, was 13.6% ( $n = 56$ ). Baseline characteristics of the insomnia patients according to hyponatremia status are shown in Supplementary Table.

Table 1  
Baseline characteristics of insomnia patients by tertiles of serum sodium levels

Variables	Total (N = 412)	< 138.0 mg/dL (N = 147)	138.0– 140.9 mg/dL (N = 136)	≥ 141.0 mg/dL (N = 129)	<i>P</i>
Age (yr)	61.5 ± 14.8	63.4 ± 14.2	60.4 ± 16.1	60.4 ± 13.8	0.150
Men (%)	231 (56.1)	93 (63.3)	77 (56.6)	61 (47.3)	0.028
Body mass index (kg/m <sup>2</sup> )	23.5 ± 2.2	23.3 ± 2.0	23.6 ± 2.3	23.6 ± 2.2	0.431
Systolic blood pressure (mmHg)	124.2 ± 12.1	123.8 ± 11.9	124.7 ± 12.1	124.1 ± 12.5	0.826
Diastolic blood pressure (mmHg)	79.8 ± 7.8	79.9 ± 7.3	80.0 ± 8.0	79.5 ± 8.1	0.906
Serum sodium, (mmol/L)	138.9 ± 3.7	135.0 ± 2.9	139.6 ± 0.9	142.5 ± 1.6	< 0.001
Serum potassium (mmol/L)	4.1 ± 0.5	4.2 ± 0.5	4.2 ± 0.5	4.1 ± 0.5	0.570
Hemoglobin (g/dL)	12.6 ± 2.0	12.2 ± 1.9	12.8 ± 2.0	12.7 ± 2.0	0.012
Calcium (mg/dL)	8.9 ± 0.7	8.7 ± 0.6	9.0 ± 0.6	9.0 ± 0.6	< 0.001
Phosphorus (mg/dL)	3.5 ± 0.8	3.4 ± 0.9	3.6 ± 0.7	3.7 ± 0.7	0.025
Glucose (mg/dL)	129.9 ± 50.2	137.7 ± 59.6	130.9 ± 46.4	120.1 ± 40.1	0.015
Total Protein (g/dL)	6.5 ± 0.8	6.3 ± 0.8	6.6 ± 0.8	6.5 ± 0.7	0.043
Albumin (g/dL)	3.8 ± 0.7	3.5 ± 0.7	4.0 ± 0.7	4.0 ± 0.6	< 0.001
Cholesterol (mg/dL)	165.9 ± 45.2	159.6 ± 52.6	170.9 ± 41.7	171.5 ± 37.4	0.007
Uric acid (mg/dL)	4.6 ± 1.7	4.0 ± 1.9	4.8 ± 1.5	5.0 ± 1.6	< 0.001
eGFR (mL/min/1.73 m <sup>2</sup> )	88.3 ± 24.8	89.0 ± 28.1	88.4 ± 22.4	87.4 ± 22.4	0.787
Follow up duration (month)	49.4 ± 29.0	42.5 ± 29.2	53.8 ± 27.7	52.9 ± 28.9	0.001
Comorbid diseases					
eGFR; estimated glomerular filtration rate					

Variables	Total (N = 412)	< 138.0 mg/dL (N = 147)	138.0– 140.9 mg/dL (N = 136)	≥ 141.0 mg/dL (N = 129)	<i>P</i>
Diabetes mellitus (%)	107 (26.0)	46 (31.3)	34 (25.0)	27 (20.9)	0.140
Hypertension (%)	162 (39.3)	60 (40.8)	56 (41.2)	46 (39.3)	0.586
Chronic kidney disease (%)	35 (8.5)	13 (8.8)	12 (8.8)	10 (7.8)	0.935
Cardiovascular disease (%)	114 (27.7)	45 (30.6)	38 (27.9)	31 (24.0)	0.474
Chronic respiratory disease (%)	62 (26.8)	30 (30.3)	16 (25.8)	16 (22.9)	0.548
Alcoholic liver disease (%)	30 (13.0)	15 (15.2)	9 (14.5)	6 (8.6)	0.418
Use of thiazide (%)	32 (7.8)	12 (8.2)	11 (8.1)	9 (7.0)	0.920
eGFR; estimated glomerular filtration rate					

## Clinical parameters affecting the serum sodium level

We measured parameters affecting the serum sodium level in the insomnia patients. Male sex, hemoglobin, calcium, uric acid, albumin, and cholesterol were positively correlated with serum sodium level, whereas use of thiazides was negatively correlated with serum sodium level based on univariate analysis. However, age, BMI, SBP, DBP, HR and co-morbid diseases were not significantly associated with serum sodium level. In a backward multivariate linear model, sex, calcium, albumin, and eGFR were significantly associated with serum sodium level (Table 2).

Table 2  
Baseline parameters associated with serum sodium level in insomnia patients

	$\beta$	95% CI	<i>P</i>	Adjusted $\beta$	95% CI	<i>P</i>
Age (yr)	-0.02	(-0.05, 0.00)	0.072			
Sex (ref. male)	0.92	(0.20, 1.64)	0.013	1.53	(0.30, 2.76)	0.015
Systolic blood pressure (mmHg)	-0.01	(-0.03, 0.03)	0.932			
Diastolic blood pressure (mmHg)	-0.02	(-0.07, 0.03)	0.420			
Hemoglobin (mg/dL)	0.23	(0.05, 0.41)	0.011			
Calcium (mg/dL)	1.16	(0.54, 1.79)	< 0.001	-1.30	(-2.52, -0.08)	0.037
Phosphorus (mg/dL)	0.45	(-0.08, 0.98)	0.095			
Glucose (g/dL)	-0.01	(-0.01, 0.00)	0.086			
Uric acid (mg/dL)	0.35	(0.12, 0.58)	0.003			
Albumin (g/dL)	1.65	(1.12, 2.18)	< 0.001	2.27	(1.05, 3.49)	< 0.001
Cholesterol (mg/dL)	0.02	(0.01, 0.02)	< 0.001			
eGFR (mL/min/1.73 m <sup>2</sup> )	0.01	(-0.01, 0.02)	0.843	0.05	(0.02, 0.08)	0.002
Diabetes mellitus	-0.37	(-1.19, 0.45)	0.381			
Hypertension	-0.14	(-0.88, 0.60)	0.716			
Chronic kidney disease	-0.21	(-1.44, 1.02)	0.743			

$\beta$ ; regression coefficient with serum sodium level. eGFR; estimated glomerular filtration rate

Adjusted for age, sex, hemoglobin, calcium, phosphorus, cholesterol, albumin, eGFR, uric acid, hypertension, diabetes, cardiovascular disease, chronic respiratory disease, alcoholic liver disease, use of thiazide.

	$\beta$	95% CI	<i>P</i>	Adjusted $\beta$	95% CI	<i>P</i>
Cardiovascular disease	-0.67	(-1.48, 0.13)	0.101			
Chronic respiratory disease	-1.00	(-2.18, 0.19)	0.098			
Alcoholic liver disease	-0.93	(-2.53, 0.66)	0.250			
Use of thiazide	-0.03	(-1.33, 1.40)	< 0.001			
$\beta$ ; regression coefficient with serum sodium level. eGFR; estimated glomerular filtration rate						
Adjusted for age, sex, hemoglobin, calcium, phosphorus, cholesterol, albumin, eGFR, uric acid, hypertension, diabetes, cardiovascular disease, chronic respiratory disease, alcoholic liver disease, use of thiazide.						

## Prediction of all-cause mortality based on the serum sodium level

We evaluated factors associated with all-cause mortality. During the median follow-up of 49.4 months, 44 (10.7%) patients died. We also examined how the risk of death varies with the overall serum sodium level. Figure 2 illustrates the nonlinear mortality risk according to the serum sodium level after adjusting for clinical covariates such as age, sex, hemoglobin, albumin, eGFR, and co-morbid diseases. There was a U-shaped association between serum sodium level and adjusted log-hazards ratio (HR). The HR was lowest at a serum concentration of 140–143 mg/dL; outside of this range, the HR increased in both directions (Fig. 2). The association of serum sodium tertile with all-cause mortality was evaluated using Kaplan-Meier analysis (Fig. 3). The results showed a significant difference in all-cause mortality among tertile groups. The lowest tertile of serum sodium (< 138.0 mg/dL) had a significantly higher mortality rate compared than the other two tertile groups. To explore the effect of serum sodium level on all-cause mortality, we performed Cox regression analyses. In multivariate analysis, being in the lowest serum sodium group (tertile 1; HR, 2.36 [95% CI: 0.96–5.79]) was an independent predictor of all-cause mortality in the insomnia patients, even after adjusting for covariates (Table 3). In addition, AKI (HR, 4.72 [95% CI: 2.39–9.34]), and CRD (HR, 2.92 [95% CI: 1.40–6.12]) were all significantly associated with all-cause mortality (Table 3).

Table 3  
Hazard ratios for all-cause mortality risk factors in insomnia patients.

	All-cause mortality	
	HR (95% CI)	P
Tertiles of serum sodium (ref. serum sodium 138.0–140.9 mg/dL)		0.014
Serum sodium < 138.0 mg/dL	2.36 (0.96–5.79)	
Serum sodium ≥ 141.0 mg/dL	0.50 (0.13–2.01)	
Acute kidney injury (ref. No)	4.72 (2.39–9.34)	< 0.001
Chronic respiratory disease (ref. No)	2.92 (1.40–6.12)	0.004
HR; hazard ratio, CI; confidence interval. Adjusted for age, Hemoglobin, albumin, eGFR, tertiles of serum sodium, diabetes, cardiovascular disease, chronic respiratory disease, alcoholic liver disease, acute kidney injury		

## Prediction of acute kidney injury

Acute kidney injury occurred in 15.0% (n = 62) of patients. Table 4 summarizes the results of multivariate logistic regression analyses. The lowest serum sodium group (tertile1; HR 2.62 [95% CI: 1.00–6.85]), eGFR (HR, 0.97 [95% CI: 0.95–0.98], DM (HR, 2.42 [95% CI: 1.01–5.40]), and ALD (HR, 3.62 [95% CI: 1.31–10.02]) were significantly associated with AKI.

Table 4  
Odds ratio for acute kidney injury risk factors in insomnia patients.

	AKI	
	OR (95% CI)	P
Tertiles of serum sodium (ref. serum sodium 138.0–140.9 mg/dL)		0.017
Serum sodium < 138.0 mg/dL	2.62 (1.00–6.85)	
Serum sodium ≥ 141.0 mg/dL	0.69 (0.20–2.38)	
Estimated glomerular filtration rate	0.97 (0.95–0.98)	< 0.001
Diabetes mellitus (ref. No)	2.42 (1.01–5.40)	0.032
Alcoholic liver disease (ref. No)	3.62 (1.31–10.02)	0.013
OR; odds ratio, CI; confidence interval. Adjusted for age, hemoglobin, albumin, eGFR, tertiles of serum sodium, diabetes, cardiovascular disease, chronic respiratory disease, alcoholic liver disease		

## Discussion

Our study showed a U-shaped relationship between overall serum sodium level and mortality and the lowest tertile of serum sodium was significantly associated with increased all-cause mortality and AKI even after adjusting for covariates. To the best of our knowledge, our results are the first to demonstrate an independent association between serum sodium level and all-cause mortality in insomnia patients.

The association between lower serum sodium level and insomnia has not been studied previously. We hypothesized that insomnia may be associated with a lower serum sodium level for the following reasons: first, the comorbidities of patients with insomnia may themselves be associated with lower serum sodium levels. In our study, 61.4% (n = 258) of patients had comorbidities, the most common of which was HT, followed by CVD, CRD, and DM. These comorbidities are known to be associated with decreased serum sodium levels and commonly cause hyponatremia. In other words, lower serum sodium levels may not be due to insomnia itself, but rather to comorbidities. Second, activated sympathetic nerve activity due to insomnia [18, 19], leading to increased renin release [20, 21] and tubular fluid reabsorption [22], may be associated with low serum sodium.

Previous studies indicated that hyponatremia is an independent predictor of increased mortality in the general population [19], as well as in patients with a variety of diseases such as acute ST-elevation myocardial infarction [23], heart failure [24], and liver disease [25]. It has not yet been determined whether hyponatremia is simply an indicator of disease severity, or itself affects the disease. Chawla et al. suggested that serum sodium is seldom the cause of death but rather a marker of the severity of underlying disease [26]. Another study suggested that hyponatremia is an independent predictor of mortality even after adjusting for age, gender, and several comorbidities in the general outpatient population [27].

In our study of patients with insomnia, the lowest serum sodium tertile had the highest risk of all-cause mortality. The exact mechanism underlying increased mortality in these patients remains unclear. However, it is possible that activation of the autonomic nervous system in insomnia patients could be associated with both lower serum sodium levels and increased mortality risk. Hyperarousal is also considered a key pathophysiological mechanism in insomnia [1], increasing the whole-body metabolic rate during sleep, high-frequency electroencephalographic activity during non-rapid eye movement sleep, and cortisol and adrenocorticotrophic hormone levels during the early sleep period, and decreasing parasympathetic tone and heart rate variability [28, 29]. These hyperarousal states may be associated with increased cardiovascular activity, and insomnia is known to be associated with both CVD risk and mortality [30]. In our study, we could only identify HR as a factor related to the sympathetic nerve activity, but there was no significant association between HR and serum sodium, mortality. Due to our study enrolled hospitalized patients, it is difficult to identify sympathetic nerve activity with HR alone.

Another hypothesis is that hyponatremia may be associated with various medical conditions including bone fractures, falls [9, 10], cardiovascular events [31], and cognitive dysfunction [3, 32], eventually leading to a high mortality rate [27]. Our study showed that the lowest tertile of serum sodium had a higher proportion of comorbidities, although not statistically significantly. Also, certain demographic,

hematologic, and biochemical parameters, such as older age and lower serum hemoglobin, calcium, phosphorus, protein, albumin, and uric acid levels, were commonly seen in our insomnia patients in the lowest tertile of serum sodium. These variables also had a direct or indirect impact on mortality.

Interestingly, all-cause mortality was significantly associated with CRD in insomnia patients. Previous studies showed that poor sleep quality was common among patients with COPD [33, 34] and disturbed sleep was associated with mortality and adverse COPD outcomes [35]. Severe hypoxemia was observed during sleep in COPD patients [36], which not only causes insomnia, but also might be associated with poor clinical outcomes. Consistent with previous studies, our study shows that inpatients with insomnia are associated with high mortality in the presence of CRD as an underlying disease. And in light of this, when complaining of insomnia in patients with CRD, the exacerbation of respiratory disease should be considered and factors to be corrected should be sought.

Previous studies have shown that hyponatremia is associated with the development of AKI in hospitalized patients [37]. Other reports have suggested that hyponatremia is a significant prognostic factor for renal replacement therapy in CKD patients treated with diuretics, eventually leading to AKI [38]. Furthermore, one report showed that serum sodium itself would not have a significant effect on kidney function [39]. However, no study has explored the relationship between AKI incidence and lower serum levels in insomnia patients. In our study, lowest tertile of serum sodium was significantly related with AKI in insomnia patients, and this is also explained by the factors involve in the death of hyponatremia mentioned above.

There were several limitations to our study. First, since it used a single-center retrospective design and relied on data from medical records, we could not tightly control certain factors that may affect the serum sodium level such as volume status, drugs (excluding thiazide), and hormone levels, and our results may thus not be generalizable. Second, we obtained serum sodium levels at baseline only; we could not obtain them at follow-up. Therefore, we could not monitor changes in the serum sodium level. Third, we enrolled insomnia patients based only on the ICD code and did not use other tools such as polysomnography or sleep habit questionnaires. However, we believe that these limitations were ameliorated by the large number of patients enrolled and the use of robust statistical methods. Relatively similar laboratory tests were applied and patients were followed-up at the same facility, since this was a single-center study. For the same reason, our results should not be generalized and further studies remain to be elucidated.

In conclusion, our study showed a relationship between lower serum sodium levels and mortality in insomnia patients. The lowest tertile of serum sodium level was associated with mortality and AKI in these patients. Further studies are required to verify the mechanism by which insomnia, low serum sodium level, and poor clinical outcomes are associated. In conclusion, physicians should consider serum sodium as a prognostic factor in patients with insomnia.

## Abbreviations

ALD: Alcoholic liver disease; AKI: acute kidney injury; BMI: Body mass index; CAD; Coronary heart disease; CKD: Chronic kidney disease; CVD: Cardiovascular disease; COPD: Chronic obstructive pulmonary disease; CRD: Chronic respiratory disease; DBP: Diastolic blood pressure; HT: Hypertension; HR: Heart rate; SBP: Systolic blood pressure

## **Declarations**

### **Ethics approval and consent to participate**

We respected all patients' rights to privacy and protected their identity. The study protocol was approved by the Institutional Review Board of Gyeongsang National University Hospital (IRB No. 2018-11-013-001), who waived the need for informed consent. All individual information was de-identified.

### **Consent for publication**

Not applicable.

### **Availability of data and materials**

The datasets used and/or analyzed during this study are available from the corresponding author on reasonable request.

### **Competing interests**

The authors declare that they have no competing interests.

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### **Authors' contributions**

EB and TWL designed the study, drafted the manuscript and performed the statistical analyses. HNJ, HSC, and SC collected and interpreted the data. EB and DJP prepared, reviewed, and revised the manuscript. DJP further supervised the work. All authors contributed to and approved the final manuscript.

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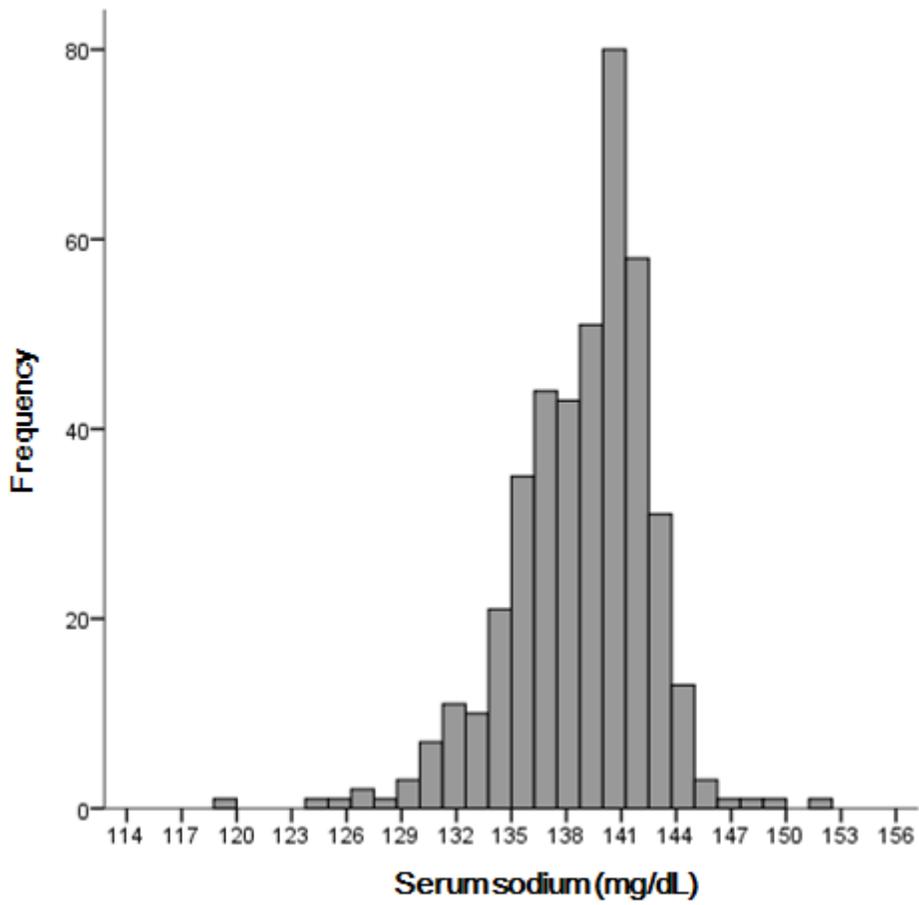
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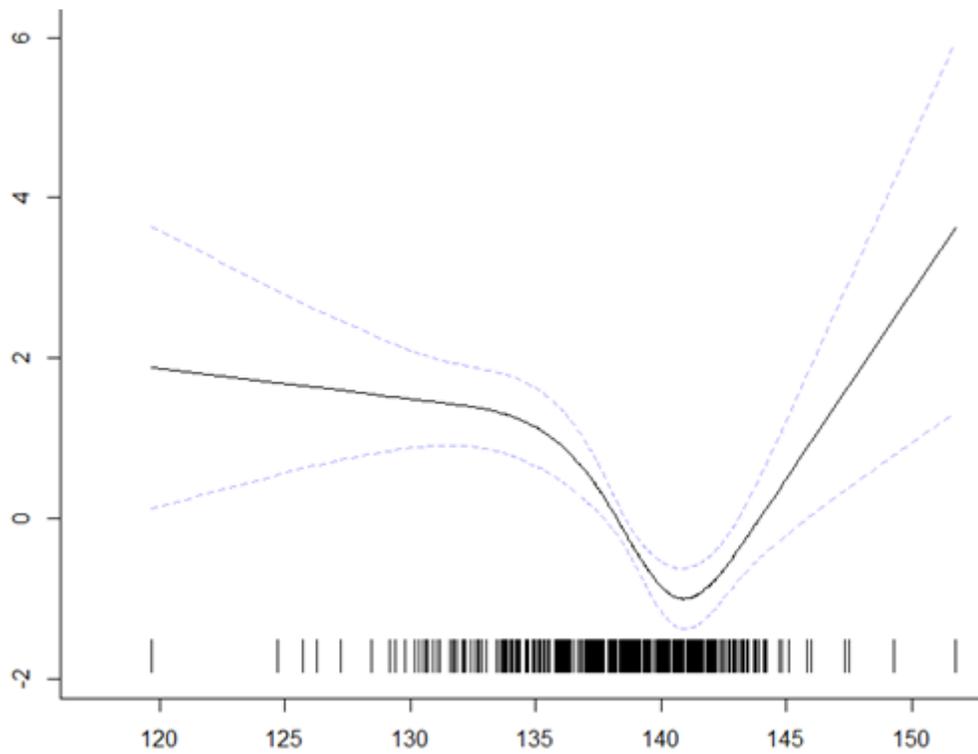
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## Figures



**Figure 1**

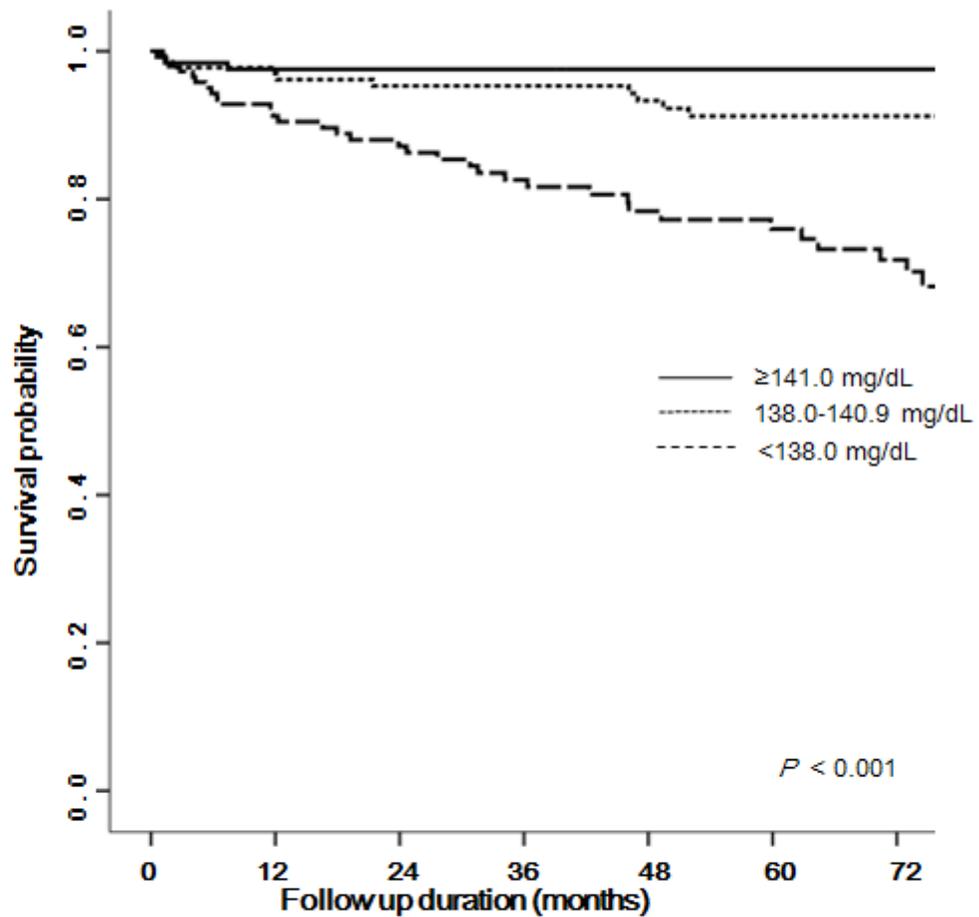
Distribution of patients relative to the serum sodium ranges.



**Figure 2**

Association between serum sodium level and hazard ratios for all-cause mortality. The log hazard ratios for all-cause mortality (solid line) and 95% confidence index (dashed lines) are presented.

Figure 3.



Number At Risk (n)

Serum sodium $\geq 141.0$ mg/dL	129	105	96	94	89	77	51
Serum sodium 138.0-140.9 mg/dL	137	121	107	98	91	78	60
Serum sodium <138.0 mg/dL	151	115	102	85	69	58	45

Figure 3

Kaplan-Meier analysis of survival probabilities for tertile of serum sodium level.

## Supplementary Files

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