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24-hour Urinary Mineral Excretion and Its Developmental Trajectory and Longitudinal Changes in Relation to Renal Deterioration in Patients with CKD

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Objectives : The aim of this study was to investigate the trajectories and longitudinal changes of 24-hour urine mineral excretion (24-hour UME) and their association with chronic kidney disease (CKD) progression in patients with CKD stages 1-4.

Methods : The study collected clinical data from patients admitted to the PLA General Hospital between January 2014 and July 2022, who had at least two 24-hour UME measurements over a year-long interval. Latent class linear mixed models (LCLMM) were used to model and visualize the 24-hour UME trajectories over three years. Multinomial regression and joint models were employed to assess the associations of 24-hour UME trajectories and longitudinal changes with rapid kidney function decline and CKD progression, respectively.

Results : The study included 1,212 participants with a mean follow-up of 30 (IQR, 19-48) months. Of these, 317 (26.2%) exhibited CKD progression. The optimal linear trajectories of 7,896 UME values for each mineral were categorized into different groups. Specifically, 3 groups were identified for sodium, chloride, potassium and magnesium; 5 for calcium; and 4 for phosphorus. Multinomial logistic regression demonstrated that patients with steady lower levels of urinary calcium (OR: 10.000, 95% CI: 4.739-21.277, P<0.001) or a decreasing trajectory (OR: 9.901, 95% CI: 3.788-25.641, P<0.001) had a significantly higher risk of rapid kidney function decline than the reference group. In joint models, baseline and longitudinal 24-hour urinary potassium and calcium independently predicted CKD progression.

Conclusions : This study discovered unique patterns of 24-hour urine mineral excretion in non-dialysis-dependent CKD patients. Baseline and longitudinal 24-hour urine excretion of potassium and calcium were determined to be significant factors in CKD progression, while sodium did not show a significant association.

Figure 1.jpg

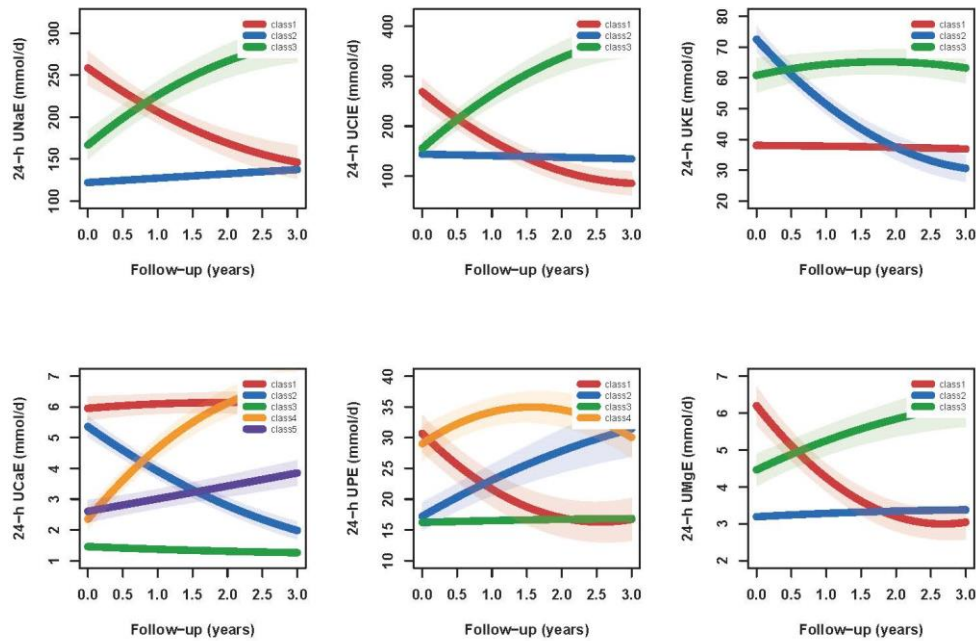


Figure 1.jpg

Table 1. The association between urine mineral excretion and CKD progression (eGFR decline \geq 50%, dialysis or kidney transplantation).

		Model 0		Model 1		Model 2	
		HR (95% CI)	P	HR (95% CI)	P	HR (95% CI)	P
UNa (mmol/d)	Baseline UNa (per 1SD)	1.178 (1.057, 1.314)	0.003	1.052 (0.937, 1.181)	0.392	1.048 (0.930, 1.182)	0.441
	Longitudinal UNa (per 1SD)	0.834 (0.662, 1.050)	0.122	0.945 (0.748, 1.192)	0.631	0.929 (0.728, 1.184)	0.550
UCI (mmol/d)	Baseline UCI (per 1SD)	1.059 (0.944, 1.189)	0.328	0.964 (0.852, 1.091)	0.557	0.956 (0.841, 1.087)	0.488
	Longitudinal UCI (per 1SD)	0.474 (0.371, 0.606)	<0.001	0.608 (0.470, 0.788)	<0.001	0.582 (0.445, 0.763)	<0.001
UK (mmol/d)	Baseline UK (per 1SD)	1.051 (0.952, 1.161)	0.325	0.925 (0.833, 1.026)	0.140	0.882 (0.789, 0.986)	0.027
	Longitudinal UK (per 1SD)	0.421 (0.322, 0.551)	<0.001	0.606 (0.460, 0.799)	<0.001	0.566 (0.426, 0.752)	<0.001
UCa (mmol/d)	Baseline UCa (per 1SD)	0.553 (0.467, 0.654)	<0.001	0.838 (0.706, 0.994)	0.043	0.809 (0.672, 0.975)	0.026
	Longitudinal UCa (per 1SD)	0.390 (0.298, 0.509)	<0.001	0.489 (0.374, 0.639)	<0.001	0.482 (0.368, 0.631)	<0.001
UP (mmol/d)	Baseline UP (per 1SD)	0.996 (0.877, 1.131)	0.950	0.997 (0.866, 1.147)	0.963	0.963 (0.833, 1.115)	0.617
	Longitudinal UP (per 1SD)	0.449 (0.367, 0.549)	<0.001	0.565 (0.458, 0.696)	<0.001	0.583 (0.470, 0.722)	<0.001
UMg (mmol/d)	Baseline UMg (per 1SD)	0.949 (0.845, 1.066)	0.379	0.916 (0.814, 1.032)	0.150	0.899 (0.795, 1.017)	0.089
	Longitudinal UMg (per 1SD)	0.636 (0.497, 0.814)	<0.001	0.729 (0.561, 0.947)	0.018	0.720 (0.551, 0.939)	0.015

Notes: HRs and 95% CIs for the effect of a SD increase in urine mineral excretion on CKD progression, adjusted for confounders. The unit for baseline and longitudinal mineral is a SD increase in urine mineral excretion. Estimates of Model 0 were without adjustment. Estimates of model 1 were adjusted from multivariable joint model for age, sex, eGFR, and 24-hour proteinuria (Ln-transformed) at baseline. Besides the variables in model 1, model 2 was also adjusted for serum mineral (Na, K, Cl, Ca, P, Mg), hemoglobin, and albumin, 24-hour urine creatinine, hypertension, diabetes mellitus, and diuretics usage at baseline.