Clinical utility of urinary levels of catecholamines and their fraction ratios related to heart rate and thyroid function

Naruhiko Sunada, Yoshihisa Hanayama, Koichiro Yamamoto, Yasuhiro Nakano, Takahiro Nada, Hiroyuki Honda, Kou Hasegawa, Hideharu Hagiya and Fumio Otsuka

Department of General Medicine, Okayama University Graduate School of Medicine, Dentistry and Pharmaceutical Sciences, Okayama 700-8558, Japan

Abstract. Urinary catecholamines (CAs) have been examined for the screening of pheochromocytomas. The decision to perform screening is based on symptoms suggesting secondary hypertension or hyperactivities of the sympathetic nervous system. To elucidate the usefulness of urinary fractions and ratios of CAs, 79 patients in whom 24-h excretions of urinary CAs including adrenaline (AD), noradrenaline (NA) and dopamine (DA) had been examined from 2015 until 2020 were retrospectively analyzed. There were no significant differences in urinary CA levels between two age groups, gender groups and two BMI groups. Patients with histories of preexisting hypertension and diabetes showed significantly higher levels of urinary NA excretion, and the urinary ratio of NA/DA was also increased in the patients with a history of hypertension. Heart rate (HR) was significantly correlated with the urinary ratio of NA/DA. Serum free thyroxine (FT4) concentration and ratio of FT4/thyrotropin (TSH) were correlated with the level of urinary AD. The levels of TSH and FT4/TSH showed negative and positive correlations, respectively, with the urinary NA/DA ratio. Thus, increases of HR are related to the enhanced conversion of DA to NA and increased thyroid hormones are involved in the increase in urinary AD and the conversion of DA to NA. History of lifestyle-related diseases and changes of HR and thyroid functions need to be considered for the evaluation of urinary CAs and their ratios.

Key words: Adrenaline, Catecholamine, Hypertension, Noradrenaline, Thyroid function

LEVELS of catecholamines (CAs) in plasma and urine are often examined to evaluate the possibility of pheochromocytomas and/or paragangliomas (PPGLs) in hypertensive patients [1]. The patients with PPGLs have various symptoms: so-called “5-H” symptoms that arise from tumor-producing CAs including hypertension (HT), headaches, hyperhidrosis, hypermetabolism and hyperglycemia [2-4]. Therefore, plasma and/or urinary levels of CAs are often examined in patients with such symptoms. Other symptoms in patients with PPGLs include anxiety, nausea, vomiting, and weakness [5].

CAs include dopamine (DA), noradrenaline (NA), and adrenaline (AD), which are synthesized from tyrosine. Tyrosine hydroxylase (TH) converts tyrosine to 3,4-dihydroxyphenylalanine (DOPA), and this is followed by conversion by DOPA decarboxylase of DOPA to DA, conversion by dopamine β-hydroxylase of DA to NA, and conversion by phenylethanolamine N-methyltransferase (PNMT) of NA to AD [6-8]. In this cascade process, the action of TH is considered to be a rate-limiting step [9]. In clinical situations, plasma CA concentrations and urinary levels of CA excretion are often utilized to determine the systemic secretory activities of CAs that are produced by the adrenal medulla and various sympathetic ganglions. The biochemical diagnosis of PPGLs consists of the demonstration of hypersecretion of CAs or their O-methylated metabolites, the latter of which show a high sensitivity for detecting PPGLs [10].
However, the clinical usefulness and significance of measurements of urinary CAs other than for examining the possibilities of PPGLs have not been established. In the present study, we retrospectively investigated the clinical significance of urinary excretions of CAs and the relative ratios of CA fractions.

**Patients and Methods**

**Study design**

We retrospectively screened the medical records of 79 patients whose levels of urinary CAs were examined during the period from January 2015 to December in 2020 at the Department of General Medicine, Okayama University Hospital. Of those patients, 5 patients with active PPGLs, 1 patient with adrenal hyperplasia, 1 patient with adrenal lymphoma and 6 patients who had been prescribed catecholaminergic agonists were excluded. As a result, 66 patients (27 males, 39 females) were included in the present analysis. Examination of urinary CAs was decided by each physician when hyper- or hypo-secretion of CAs was clinically suspected on the basis of each patient’s symptoms, laboratory data, and past medical history. All of the urinary collections for CAs excretions were performed in hospitalized conditions for one to two days in our hospital. Data for other biochemical markers were also obtained within a few days of measurement of urinary CA excretions. These examinations were performed for the patients under adequate insurance coverage. Information regarding the present study was provided on our hospital wall and on the website of our hospital, and patients who wished to opt out were offered that opportunity. This study was approved by the Ethical Committee of Okayama University Hospital (KEN 2106-016) and adhered to the Declaration of Helsinki.

**Analysis of clinical parameters**

Information on the patients’ chief complaints was obtained from hospital medical records. Data on age, gender, body mass index (BMI), self-rated depression scale (SDS), and frequency scale for symptoms of gastroesophageal reflux disease (FSSG) were also obtained. Diagnosis of hypertension was based on the definition of hypertension: having a systolic blood pressure (SBP) ≥140 mmHg, having a diastolic blood pressure (DBP) ≥90 mmHg, or taking BP lowering agents [11]. Values for the following blood biochemical parameters were obtained from records, when available: white blood cells (WBC), hemoglobin, and platelets for blood cell counts; total bilirubin (T.Bil), albumin, aspartate aminotransferase (AST), alanine aminotransferase (ALT), alkaline phosphatase (ALP), lactate dehydrogenase (LDH), creatinine (Cr), urea nitrogen (UN), estimated glomerular filtration rate (eGFR), sodium, potassium, and corrected calcium (Ca) for liver and renal functions; blood glucose, hemoglobin A1c (HbA1c), low-density lipoprotein cholesterol (LDL-C) and high-density lipoprotein cholesterol (HDL-C) as metabolic markers; and thyrotrpin (TSH), free thyroxine (FT4), free triiodothyronine (FT3), and ratio of TSH/FT4 as thyroid function markers. eGFR was calculated by using the following formulas: 194 × Age−0.287 × Cr−1.094 for males and 194 × Age−0.287 × Cr−1.094 × 0.739 for females. Urinary CA levels were determined by high-performance liquid chromatography (HPLC; BML, Inc., Saitama, Japan) in 24-h collected urine samples with 6-M hydrochloric acid to prevent degradation of collected urinary CAs [12]. The normal ranges of urinary CAs are as follows: AD: 3.0–41.0 μg/day; NA: 31.0–160.0 μg/day; and DA: 280.0–1,100.0 μg/day. All other levels were determined using an auto-analyzer system at the Central Laboratory of Okayama University Hospital.

**Statistical analysis**

EZR, version 1.40 (Saitama Medical Center, Jichi Medical University, Saitama, Japan), which is a graphical user interface for R (The R Foundation for Statistical Computing, Vienna, Austria), was used in all statistical analyses [13]. It is modified from R commander, which is designed to add frequently used functions in biostatistics. The Mann-Whitney U test or Spearman’s rank correlation coefficient was used to statistically analyze continuous measurements. All tests were performed as two-sided, and p < 0.05 was regarded as statistically significant.

**Results**

The mean age of the patients was 64 years (interquartile range: IQR, 45–69 years), the male/female ratio was 39 (59%)/27 (41%), and mean BMI was 24.1 kg/m² (IQR, 21.6–26.0 kg/m²). As shown in Fig. 1, it was found that there were no significant differences in urinary levels of AD, NA and DA or the ratios of AD/NA and NA/DA between the two age groups (<65 vs. ≥65 years), two gender groups (M vs. F) and two BMI groups (<25 vs. ≥25 kg/m²). In patients with urinary NA higher than the normal upper limit (>160 μg/day), 52%, 61% and 45% of the patients were older patients (≥65 years), female patients and patients with high BMI (≥25 kg/m²), respectively, whereas 46%, 57% and 34% (slightly lower percentages) of the patients with normal urinary NA (≤160 μg/day) were older patients (≥65 years), female patients and patients with high BMI (≥25 kg/m²), respectively.
The mean values of SBP, DBP and heart rate (HR) were 129 mmHg (IQR, 119–144 mmHg), 78 mmHg (75–85 mmHg) and 74 bpm (64–78 bpm), respectively. The major symptoms for which it was decided to perform CA measurements were headaches in 13 cases (19.7%), palpitation in 8 cases (12.1%), hyperhidrosis in 6 cases (9.1%), and body weight loss in 6 cases (9.1%). The medical histories of the patients included HT in 53 patients (80.3%), glucose intolerance (HbA1c ≥6.5) in 8 patients (12.1%), and chronic kidney diseases (CKD) in 12 patients (18.2%).

Next, the impacts of clinical backgrounds including histories of HT, CKD and diabetes mellitus were assessed in relation to the levels of urinary levels of AD, NA and DA or the ratios of AD/NA and NA/DA. As shown in Fig. 2, among the medical histories of HT, CKD and diabetes mellitus, patients with preexisting HT and diabetes (hemoglobin A1c ≥6.5) showed significantly higher levels of urinary NA excretion. In patients with urinary NA higher than the normal upper limit (>160 μg/day), 90%, 20% and 20% of the patients had preexisting HT, CKD and diabetes, respectively, whereas 71%, 17% and 7% (slightly lower percentages) of the patients with normal urinary NA (≤160 μg/day) had HT, CKD and diabetes, respectively. Of interest, urinary ratios of AD/NA were significantly lower in the patients with medical histories of HT and diabetes than in the patients without these disorders. On the other hand, urinary ratio of NA/DA in the patients with a history of HT was significantly higher than that in the patients without a history of HT, suggesting that the conversion step from DA to NA could be clinically linked to the etiology of HT.

Fig. 1 Urinary levels of catecholamines and their fraction ratios based on the characteristics of patients in the present study. Urinary levels of all catecholamines (CAs; A) including adrenaline (AD), noradrenaline (NA) and dopamine (DA) and the urinary ratios (B) were compared in two age groups (<65 vs. ≥65 years), two gender groups (male vs. female), and two body mass index (BMI) groups (<25 vs. ≥25 kg/m²). In each panel, the upper horizontal line of the box is the 75th percentile, the lower horizontal line of the box is the median, the upper horizontal bar outside the box is the maximum value within 1.5 times the interquartile range, and the lower horizontal bar outside the box is the minimum value within 1.5 times the interquartile range.
The relationships of various clinical data with urinary levels of CAs and ratios of CAs are shown in Table 1. Considering the process of the production of CAs, we also focused on the enzymatic conversion steps from DA to NA and the following step caused by PNMT from NA to AD. To characterize the converting activities in the adrenal medulla and sympathetic ganglia, urinary ratios of AD/NA and NA/DA were calculated and statistically analyzed. It was found that urinary ratios of NA/DA were significantly correlated to HR, serum LDL-C levels and thyroid functions. In addition, blood glucose levels were positively correlated with urinary levels of NA but negatively correlated with urinary ratios of AD/NA, whereas HbA1c levels were not significantly correlated with either urinary level of CAs or the ratios.

The key clinical symptoms related to the actions of CAs were compared to the urinary levels of CAs. As shown in Fig. 3A, individual levels of SBP were not significantly correlated with either urinary level of AD or NA. On the other hand, HR was significantly (r = 0.285; *p < 0.05) correlated with the ratio of urinary levels of NA/DA (Fig. 3B).

Furthermore, to try to determine the factors involved in the regulation of HR, results of thyroid function tests in relation to changes of HR were analyzed. As shown in Fig. 4, it was notable that serum FT4 concentration (r = 0.443; **p < 0.01; Fig. 4A) and ratio of serum FT4/TSH (r = 0.291; *p < 0.05; Fig. 4C) were significantly correlated with the level of urinary AD excretion. The level of serum TSH (r = -0.344; **p < 0.01; Fig. 4B) and ratio of serum FT4/TSH (r = 0.373; **p < 0.01; Fig. 4C) showed significant negative and positive correlations, respectively, with the ratio of urinary levels of NA and DA.
Table 1  Relationships of clinical data with urinary levels of catecholamines and their ratios

| CA, Ratios | U-AD | | | U-NA | | | U-AD/NA | | | U-NA/DA | | |
|------------|------|------|------|------|------|------|------|------|------|------|------|------|
|            | \( r \) | \( p \) values | \( r \) | \( p \) values | \( r \) | \( p \) values | \( r \) | \( p \) values | \( N \) |
| **Patients' profile** | | | | | | | | | | | | |
| Age | 0.0559 | 0.656 | 0.0949 | 0.449 | \(-0.0236\) | 0.851 | 0.201 | 0.105 | 66 |
| BMI | 0.000773 | 0.995 | 0.26 | 0.0349** | \(-0.18\) | 0.148 | \(-0.0507\) | 0.686 | 66 |
| SDS | 0.34 | 0.104 | \(-0.0262\) | 0.903 | 0.252 | 0.234 | 0.018 | 0.934 | 24 |
| FSSG | \(-0.322\) | 0.134 | \(-0.147\) | 0.504 | \(-0.0367\) | 0.868 | 0.065 | 0.768 | 23 |
| SBP | \(-0.189\) | 0.128 | \(-0.0832\) | 0.506 | \(-0.112\) | 0.372 | 0.0503 | 0.688 | 66 |
| HR | 0.0447 | 0.722 | 0.231 | 0.0618 | \(-0.152\) | 0.224 | 0.285 | 0.0205* | 66 |
| **Laboratory data** | | | | | | | | | | | | |
| WBC | \(-0.00355\) | 0.978 | 0.207 | 0.101 | \(-0.0759\) | 0.551 | \(-0.0134\) | 0.916 | 64 |
| Hemoglobin | 0.0732 | 0.565 | 0.161 | 0.204 | \(-0.101\) | 0.427 | \(-0.0501\) | 0.694 | 64 |
| Platelets | 0.101 | 0.428 | 0.248 | 0.0479* | \(-0.00365\) | 0.977 | 0.0689 | 0.589 | 64 |
| T. Bil | \(-0.0713\) | 0.591 | \(-0.215\) | 0.102 | 0.0414 | 0.756 | \(-0.0318\) | 0.811 | 59 |
| Albumin | 0.0747 | 0.558 | \(-0.116\) | 0.36 | 0.0734 | 0.564 | \(-0.0922\) | 0.469 | 64 |
| AST | \(-0.128\) | 0.318 | 0.219 | 0.0847 | \(-0.239\) | 0.0594 | 0.0291 | 0.821 | 63 |
| ALT | \(-0.104\) | 0.408 | 0.225 | 0.0717 | \(-0.274\) | 0.0274* | \(-0.0663\) | 0.6 | 65 |
| ALP | \(-0.163\) | 0.201 | 0.0597 | 0.642 | \(-0.096\) | 0.454 | 0.197 | 0.121 | 63 |
| LDH | 0.0141 | 0.913 | 0.219 | 0.0852 | \(-0.116\) | 0.366 | 0.0335 | 0.795 | 63 |
| Creatinine | 0.00554 | 0.965 | \(-0.00674\) | 0.958 | 0.0681 | 0.59 | 0.173 | 0.168 | 65 |
| Urea nitrogen | \(-0.0399\) | 0.752 | 0.145 | 0.25 | \(-0.121\) | 0.337 | 0.103 | 0.414 | 65 |
| eGFR | \(-0.0165\) | 0.896 | \(-0.0925\) | 0.463 | \(-0.0217\) | 0.864 | \(-0.195\) | 0.12 | 65 |
| Sodium | \(-0.0721\) | 0.568 | \(-0.00298\) | 0.981 | \(-0.0974\) | 0.44 | \(-0.0699\) | 0.58 | 65 |
| Potassium | 0.069 | 0.585 | 0.14 | 0.266 | \(-0.0558\) | 0.659 | \(-0.0229\) | 0.856 | 65 |
| Corrected Ca | 0.2 | 0.112 | 0.163 | 0.197 | 0.0544 | 0.67 | \(-0.000289\) | 0.998 | 64 |
| **Endocrine and metabolic data** | | | | | | | | | | | | |
| LDL-C | 0.057 | 0.679 | 0.0905 | 0.511 | \(-0.0224\) | 0.871 | \(-0.278\) | 0.0402* | 55 |
| HDL-C | 0.0448 | 0.752 | 0.0802 | 0.572 | \(-0.0696\) | 0.624 | 0.149 | 0.291 | 52 |
| Glucose | 0.052 | 0.706 | 0.0551 | 0.0000131** | \(-0.355\) | 0.000787** | 0.229 | 0.093 | 55 |
| HbA1c | \(-0.0437\) | 0.74 | 0.22 | 0.0906 | \(-0.223\) | 0.0867 | 0.0469 | 0.722 | 60 |
| TSH | \(-0.211\) | 0.125 | \(-0.0983\) | 0.479 | \(-0.147\) | 0.289 | \(-0.344\) | 0.0108* | 54 |
| FT3 | 0.0307 | 0.901 | \(-0.193\) | 0.427 | 0.0894 | 0.716 | \(-0.105\) | 0.67 | 19 |
| FT4 | 0.443 | 0.000896** | 0.162 | 0.248 | 0.222 | 0.109 | 0.205 | 0.141 | 53 |
| FT4/TSH | 0.291 | 0.0347* | 0.109 | 0.437 | 0.202 | 0.146 | 0.373 | 0.00592** | 53 |
| BNP | 0.216 | 0.31 | 0.0805 | 0.709 | 0.112 | 0.602 | \(-0.207\) | 0.332 | 24 |

** \( p < 0.01 \) and * \( p < 0.05 \) between the indicated factors (data underlined). ALP, alkaline phosphatase; ALT, alanine aminotransferase; AST, aspartate aminotransferase; BMI, body mass index; BNP, brain natriuretic peptide; Ca, calcium; eGFR, estimated glomerular filtration rate; FSSG, frequency scale for symptoms of gastroesophageal reflux disease; FT3, free triiodothyronine; FT4, free thyroxine; HbA1c, hemoglobin A1c; HDL-C, high-density lipoprotein cholesterol; HR, heart rate; LDH, lactate dehydrogenase; LDL-C, low-density lipoprotein cholesterol; SBP, systolic blood pressure; SDS, self-rating depression scale; T. Bil, total bilirubin; TSH, thyrotropin and WBC, white blood cells.

**Discussion**

In the present study, urinary NA levels were found to be increased in patients with histories of HT and diabetes, in which the conversion step from DA to NA is likely to be involved in the medical history of HT. Of note, it was revealed that increases of HR are related to enhancement of the conversion of urinary DA to NA and that increased thyroid hormones are potentially involved in the increase in urinary AD and the conversion of DA to NA.

As a background of the patients, patients with preexisting HT showed significantly higher levels of urinary NA excretion and urinary ratios of NA/DA. In this
regard, Esler et al. showed the presence of a high level of activation of renal sympathetic outflow in patients with untreated essential HT [14], suggesting that renal sympathetic activation is substantially related to the pathogenesis of HT. Regarding the clinical significance of the sympathetic nervous system as assessed by urinary CAs, Missouris et al. showed an interrelationship between urinary NA levels and blood pressure in patients with essential HT [15]. In the present study, it was found that changes of HR were correlated with the urinary ratio of NA/DA. Given the finding that urinary NA excretion and resting HR are increased in relation to the severity of HT [15], these findings imply that sympathetic overactivity may play an important role in the etiology and progression of essential HT.

As for the interrelationships between urinary levels of CAs and glucose tolerance, Chamapaneri et al. showed that men, but not women, with diabetes had significantly lower urinary levels of CAs than those in men without diabetes among the various neuroendocrine parameters they examined [16]. The present study also showed that patients with preexisting diabetes (hemoglobin A1c ≥6.5) had significantly higher levels of urinary NA excretion. Also, the levels of blood glucose were correlated to the levels of urinary NA and the ratios of AD/NA. It is thus thought that the pathophysiology of diabetes is, at least in part, associated with neuroendocrine dysregulation, though there might be some differences depending on the situations and gender.

In view of the relationship between urinary CAs and

![Fig. 3](https://example.com/fig3.png)

Interrelationships between the levels of blood pressure, heart rate and urinary catecholamines. The patients were further examined for interrelationships between urinary levels of catecholamines (CAs), including adrenaline (AD) and noradrenaline (NA), and levels of systolic blood pressure (SBP; A). The interrelationships between urinary levels of CAs or ratios of urinary AD/NA and NA/DA and heart rate (HR; B) were also analyzed. * p < 0.05, statistically significant between the indicated factors.

![A](https://example.com/fig3a.png)

![B](https://example.com/fig3b.png)
metabolic activity, Hollstein et al. demonstrated a functional link between CAs and energy expenditure [17]. They showed that NA rather than AD was independently associated with both energy expenditure and sleeping metabolic rate. Thus, sympathetic nervous system activity, mediated via NA, is one of the determinants of human energy expenditure in a basal non-stress condition [17]. In the present study, it was revealed that increases of HR are related to the enhancement of urinary NA and that increased thyroid hormones are also involved in enhanced conversion from DA to NA. Taken together, the results indicate that the systemic level of NA might be a useful marker to know the basal metabolic status related to susceptibility to metabolic syndrome.

The mutual interaction between the levels of urinary CAs and thyroid hormone levels may be involved in some critical disorders. For instance, thyroid hormones enable sensitization of the heart for CAs by stimulating β1 receptor expression in cardiomyocytes [18, 19]. An excess of thyroid hormones might potentiate the effects of CAs in cardiovascular tissues, leading to increased sensitivity to stress events, which might be involved in the etiology of Takotsubo cardiomyopathy [19]. The results of our study may also indicate the presence of a functional sympatho-adrenergic system in various patients who visit a general medicine department. Unfortunately, since the sample number for FT3 measurement was much smaller than that for FT4 measurement in this retrospective study, a prospective study for assessment of the interrelationships between levels of FT3 and CAs is needed.

In the present study, there was no significant difference on urinary DA levels between the groups based on

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**Fig. 4** Interrelationships between thyroid functions and urinary levels of catecholamines. The patients were further examined for interrelationships between urinary levels of catecholamines (CAs), including adrenaline (AD) and noradrenaline (NA), or ratios of urinary AD/NA and NA/DA and levels of serum free thyroxine (FT4; A) and thyrotropin (TSH; B) and ratios of FT4/TSH (C). **p < 0.01 and * p < 0.05, statistically significant between the indicated factors.
various clinical background. However, the sympatho-renal system including local DA contents could also be related to the pathophysiology of arterial hypertension and cardiovascular damages [20]. On the other hand, sympathetic neural factors including NA and AD are potentially involved in the pathophysiology of life-threatening cardiac disorders, especially in patients with heart failure, obesity, sleep apnea, or CKD [21, 22], which are closely related to marked activation of sympathetic stimuli [23].

For instance, both NA and AD levels were elevated in plasma and urine in obstructive sleep apnea patients [24, 25]. Animal studies also suggested that intermittent hypoxia facilitates catecholamine secretion from the adrenal medulla [25]. Persistent sympathetic activation and hypertension associated with sleep apnea are functionally linked to sympathetic activation and action of elevated circulating levels of CA on the vasculature. Hence, it is important to take care of latent life-threatening conditions accelerated by sympathetic neural activities that are detected by a urinary NA excess.

Collectively, the results of the present study revealed that urinary NA excretion and/or the conversion of DA to NA are increased in patients with histories of HT and diabetes. Increases of HR, but not blood pressure, are linked to enhancement of urinary NA excretion, and increased thyroid hormones are also involved in the increases in urinary AD levels and the conversion of DA to NA. On the other hand, when a decision is made to measure urinary CAs, the patient’s history of lifestyle-related diseases and the patient’s HR and thyroid functions should also be taken into account for evaluating activity of the sympathetic nervous system in the clinical setting of general practice.

In the present study, the conversion steps from DA to NA and from NA to AD were analyzed, for the first time, as urinary ratios of AD/NA and NA/DA in addition to the levels of urinary CAs. However, there were several limitations in the present cross-sectional study. Urinary CA levels might be affected by various foods, stress conditions, and ingredients such as caffeine, nicotine, and amine-rich foods [26], although the sampling conditions were carefully performed in each patient in our study. Urinary CA levels can be affected by medications including antihypertensive drugs and tricyclic antidepressant drugs, and it has also been reported that urinary NA excretion can be increased in hypertensive patients taking single-drug therapy with either a long-acting dihydropyridine calcium antagonist or a β-blocker [15].

Serum levels of FT3 in a larger number of patients should also be analyzed to determine the direct interaction between thyroid hormones and CAs, though there is some difficulty for measuring FT3 under the coverage of Japanese medical insurance. In addition, more specific and sensitive methods such as LC-MS/MS for quantitative determination of urinary CAs would be ideal to exclude the effects of structurally related drugs and metabolites [27]. Finally, this study was performed as a retrospective single-center study in which sample sizes were relatively small and blood samplings were not always performed in the same periods. Further studies including a multi-center and prospective study in age- and gender-matched cohorts would be ideal to clarify the effectiveness of measurement of urinary CAs.

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Author Contributions

F.O. and Y.H. conceived and designed the study; N.S., Y.H., K.Y. and Y.N. performed data collection; T.N., H.H. and K.H. analyzed the data; and N.S. and F.O. wrote and H.H. revised the paper. All authors have read and agreed to publish the final version of the manuscript.

Disclosure

The authors have nothing to disclose.

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