Angiopoietin-2, Angiopoietin-1 and subclinical cardiovascular disease in Chronic Kidney Disease

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Angiopoietins (Angpt) and vascular endothelial growth factor (VEGF) have been associated with cardiovascular disease. The study enrolled 270 pre-dialysis stage 3–5 CKD patients to assess the link between circulating Angpt2, Angpt1 and VEGF-A and subclinical measures of cardiovascular structure and function. Serum markers of angiogenesis were measured using commercial enzyme-linked immunosorbent assays. Cardiac structure and function were examined by echocardiography. Brachial-ankle pulse wave velocity (baPWV) was measured by the ankle-brachial index. The adjusted mean of left ventricular mass index (LVMI) was 2.05 in patients of Angpt2 quartile 4 and 1.99 in those of Angpt2 quartile 1 (P = 0.04). Angpt2 was significantly associated with LV hypertrophy (LVH) (Angpt2 quartile 4 compared with Angpt2 quartile 1: adjusted OR: 2.68, 95% CI: 1.15–6.20). Angpt1 was negatively correlated with left atrial diameter (adjusted mean of LAD: 3.59 in Angpt1 quartile 4, 3.92 in Angpt1 quartile 1, P = 0.03). A positive and significant correlation was found between Angpt2 level and baPWV in spearman’s correlation, but not in adjusted model. In conclusion, high Angpt2 and low Angpt1 levels were positively associated with abnormal cardiac structure in stages 3–5 CKD patients, which is compatible with the viewpoint that angiopoietins participates in cardiovascular burdens.

Angiopoietins, as one of the endothelial growth factors, modulates vascular development and remodeling during angiogenesis and inflammation process1. There are two major types of angiopoietins: angiopoietin-1 (Angpt1) and angiopoietin-2 (Angpt2), and both of them bind to the same endothelial receptor, Tie-2. Angpt1 binds to Tie2 receptor and then activates downstream signaling, thereby stabilizing endothelial and vascular structure. However, Angpt2 possesses opposite physiological properties and expressions of Angpt1. Angpt2 interrupts Ang-1-Tie-2 signaling, and then contributes to structural and functional changes of vessels through the effect of vascular endothelial grower factor (VEGF)2.

Accumulating evidence has demonstrated that high circulating Angpt2 and VEGF levels were shown in cardiovascular diseases, including congestive heart failure3 and coronary artery disease4. Elevated Angpt2 and VEGF levels were also significantly associated with traditional risk factors for cardiovascular diseases (CVD), such as blood pressure and metabolic syndrome5,6. Furthermore, a significant relationship between Ang-2 and cardiovascular mortality had been mentioned in general population7. Our previous report also found that high Angpt2 level was significantly associated with major adverse cardiac events (MACEs) in chronic kidney disease (CKD) not on dialysis. Angpt2 was as an independent predictor of cardiovascular burdens8.

CKD patients have higher risk of developing CVD and all-cause mortality9,10. Apart from the traditional risk factors, endothelial dysfunction has been shown to associates of cardiovascular morbidity and mortality11. In the study by David et al., Angpt2 was notably elevated in CKD patients either on dialysis or not12. Circulating Angpt2, not Angpt1 level was positively correlated with coronary artery disease and peripheral artery disease scores in dialysis and transplant patients13. The mechanisms mediating the increased cardiovascular burdens are not well-known. Shroff et al. indicated a significant association of Angpt2 with intima media thickness in children on dialysis14. Angpt2 was also correlated with ventricular dysfunction and the clinical stages of heart failure in

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Table 1. The clinical characteristics of study subjects stratified by circulating angiopoietin-2 quartile. Data are expressed as number (percentage) for categorical variables and mean ± SD or median (25th, 75th percentile) for continuous variables, as appropriate. Abbreviations: CKD, chronic kidney disease; ACEI, angiotensin converting enzyme inhibitors; ARB, angiotensin II receptor blockers; eGFR, estimated glomerular filtration rate; hsCRP, high-sensitivity C-reactive protein. Bonferroni post hoc analysis was utilized for analysis between two groups. *P < 0.05 compared with quartile 1; †P < 0.05 compared with quartile 2; ‡P < 0.05 compared with quartile 3. *Angiopoietin-2 quartile cut at 1538.2, 1990.7, and 2753.2 pg/ml. ‡P-value calculated from one-way analysis of variance (ANOVA) test.
patients with Angpt2 quartile 3 than those with Angpt2 quartile 1. The proportion of usage of calcium channel blocker and β-blocker was the highest in CKD patients with Angpt2 quartile 4. Serum albumin and hemoglobin levels were lower in CKD patients with Angpt2 quartile 4 than those with Angpt2 quartile 1. Urine protein was higher in CKD patients with Angpt2 quartile 4 than other groups. Patients with Angpt2 quartile 4 had the highest incidence of CKD stage 5 and the lowest incidence of CKD stage 3 among 4 groups.

**Correlation between circulating Angpt2, Angpt1 and VEGF-A levels and cardiac structure and function.** Table 2 shows the distribution of echocardiographic parameters in study patients stratified by Angpt2 quartiles. Of all patients, the median of left ventricular mass index (LVMI) was 133 g/m² and 25.9% of those had concentric left ventricular hypertrophy (LVH) and 40.7% of those had eccentric LVH. The medians of aortic root diameter and left atrium diameter were 3.2 and 3.7 cm, respectively. 73.9% of patients had peak early transmitral filling wave velocity (E)/peak late transmitral filling wave velocity (A) less than 1. There was a significant difference of LVMI and the proportion of LVH were higher in Angpt2 quartile 4 compared with Angpt2 quartile 1. The rate of eccentric LVH was higher in Angpt1 quartile 4 than in those with Angpt2 quartile 1. Urine protein was higher in CKD patients with Angpt2 quartile 4 than other groups. Patients with Angpt2 quartile 4 had the highest incidence of CKD stage 5 and the lowest incidence of CKD stage 3 among 4 groups.

**Table 2. The echocardiographic parameters of study patients stratified by circulating angiopoietin-2 quartile.** Data are expressed as number (percentage) for categorical variables and mean ± SD or median (25th, 75th percentile) for continuous variables, as appropriate. Abbreviations: LVMI, left ventricular mass index; LVH, left ventricular hypertrophy; SV, stroke volume; LVEF, left ventricular ejection fraction; LVFS, Left ventricular fraction shortening; E/A, peak early transmitral filling wave velocity/peak late transmitral filling wave velocity. Bonferroni post hoc analysis was utilized for analysis between two groups. *P < 0.05 compared with quartile 1; †P < 0.05 compared with quartile 2; ‡P < 0.05 compared with quartile 3. Angiopoietin-2 quartile cut at 1538.2, 1990.7, and 2753.2 pg/ml. $P$-value calculated from one-way analysis of variance (ANOVA) test.

![Table 2](image_url)
still significantly associated with LVH in CKD patients with hsCRP ≤ 1 mg/L (P-trend = 0.02) and those with eGFR ≥ 20 ml/min/1.73 m² (P-trend = 0.007) after adjustment of age and sex, but not in those with hsCRP > 1 mg/L and those with eGFR < 20 ml/min/1.73 m².
Correlation between circulating Angpt2, Angpt1 and VEGF-A levels and brachial-ankle pulse wave velocity (baPWV).

A positive and significant correlation was found between baPWV and circulating Angpt2 level (spearman $r = 0.20$, $P = 0.02$). Besides, baPWV was significantly correlated with Angpt1 ($spearman r = -0.27$, $P = 0.01$) and Angpt2/Angpt1 ($spearman r = 0.28$, $P = 0.007$), but not with VEGF-A. However, there was no significant association of baPWV with Angpt2, Angpt1 or Angpt2/Angpt1 in adjusted analysis (Fig. 2 and Table 3).

Table 4. Relation of circulating Angiopoietin-1 level with left atrial diameter (LAD) in CKD stages 3–5 subjects. Multivariate adjusted model: adjusted for age, sex, diabetes mellitus, heart disease, cigarette smoking, estimated glomerular filtration rate, serum hemoglobin and cholesterol levels. $^{*}P$-value $< 0.01$ compared with quartile 1, $^{**}P$-value $< 0.05$ compared with quartile 1. Angiopoietin-1 quartile cut at 5948.9, 11378.6 and 23148.7 pg/ml.

Table 5. Relation of circulating Angiopoietin-2 level with left ventricle mass index (LVMI) in CKD stages 3–5 subjects. Multivariate adjusted model: adjusted for age, sex, diabetes mellitus, heart disease, cigarette smoking, estimated glomerular filtration rate, serum hemoglobin and cholesterol levels. $^{*}P$-value $< 0.01$ compared with quartile 1, $^{**}P$-value $< 0.05$ compared with quartile 1. Angiopoietin-2 quartile cut at 1538.2, 1990.7, and 2753.2 pg/ml.

Table 6. Relation of circulating Angiopoietin-2 level with left ventricle hypertrophy (LVH) in CKD stages 3–5 subjects. Multivariate adjusted model: adjusted for age, sex, diabetes mellitus, heart disease, cigarette smoking, estimated glomerular filtration rate, serum hemoglobin and cholesterol levels. $^{*}P$-value $< 0.01$ compared with quartile 1, $^{**}P$-value $< 0.05$ compared with quartile 1. Angiopoietin-2 quartile cut at 1538.2, 1990.7, and 2753.2 pg/ml.

Discussion

This study assessed the relationship between serum markers of angiogenesis including Angpt2, Angpt1, and VEGF-A and measures of cardiovascular structure and function as subclinical features of CVD in CKD patients not on dialysis. The results show that high circulating Angpt2 level was significantly associated with the increase of LVMI and LVH and the negative correlation between circulating Angpt1 level and LAD was also shown in patients with stages 3–5 CKD after adjustment for traditional cardiovascular risk factors. There was no significant association of baPWV with Angpt2, Angpt1 or Angpt2/Angpt1 in adjusted analysis (Fig. 2 and Table 3).

Circulating Angpt2 level has been a potential predictor of the accelerated development of cardiovascular burden. Overexpression of Ang-2 leaded to endothelial apoptosis, and promoted of expression of adhesion molecules, such as intercellular adhesion molecule 1 (ICAM-1) and vascular cell adhesion molecule 1 (VCAM-1), thereby playing a crucial role in the progression of cardiac hypertrophy and interstitial fibrosis in animal model. Lee et al. found these patients had higher Angpt2 level than control group and further observed a serial changes in circulating Angpt2 level from the acute to the chronic stages in myocardial infarction. Eleuteri et al. investigated the correlation between Angpt2 and the index of cardiac function impairment and showed that Angpt2 level was significantly correlated with the severity of congestive heart failure. The study of Health in Pomerania (SHIP) that included 3204 normal subjects with a median age of 55 years to evaluate the relationship between Ang-2 and...
cardiac function revealed that high circulating Angpt2 level was associated with low LVFS value as a measure of LV systolic function, but not with LVM as a measure of LV structure\(^ {19}\). However, the information between Angpt2 and associated markers of angiogenesis and actual cardiac structure and function status is limited in CKD population.

This present study shows that high circulating Angpt2 level, not Angpt1 or VEGF-A was significantly associated with high LVM level and increased risk for LVH, but not with either LV systolic or diastolic function in CKD patients, whereas these findings were inconsistent with those of SHIP in general population. Comparing with SHIP, our participants were older, had lower baseline eGFR, higher proportion of diabetes and anti-hypertensive drugs usage. A large dissimilarity of the participants may partially explain the inconsistency of results between the two studies. Structural and functional impairment of heart is frequently mentioned in CKD patients and these patients are easier to have high burdens of cardiovascular morbidity and mortality. There was no big difference of LVM across Angpt2 quartiles 1–3 in our CKD patients. Patients with Angpt2 quartile 4 had the lowest eGFR and highest serum hsCRP level than others. Angpt2, impaired renal function, and inflammation might have a synergic effect on cardiac hypertrophy. The significant association between Angpt2 and LVH was only consistent in patients with eGFR \(\geq 20\) ml/min/1.73 m\(^2\). Thus, further studies are needed to explore the mechanisms of the interaction between Angpt2, renal function and cardiac structure.

Chronic inflammation has been considered as a crucial factor in atherosclerosis development. Masiha\ et al.\ reported that inflammatory markers such as high-sensitivity c-reactive protein (hsCRP), ICAM-1, and VCAM-1 were elevated in concentric LVH, and hsCRP was correlated with LV diastolic function, while inflammation might play a pathogenetic role in abnormal LV geometry and function\(^ {20}\). On the other hand, Angpt2 is associated with pro-inflammatory responses with P-selectin, E-selectin and VCAM-1\(^ {12}\). Angpt2 triggers the infiltration of neutrophils and monocytes and Angpt2-induced signaling leads to induction of adhesion molecules and vascular leakage\(^ {21}\). It is possible that the influence of the interaction between inflammation and Angpt2 on cardiac function and structure exists. Our subgroup analysis found that circulating Angpt2 level was significantly associated with LVH in CKD patients with hsCRP \(\leq 1\) mg/L not in those with hsCRP \(> 1\) mg/L. There might be a complicated interaction between Angpt2 and inflammation in cardiac remodeling.

The Angpt-Tie system has been considered as a major regulator of vascular and endothelial cell maintenance. Vascular and endothelial dysfunction would contribute to cardiovascular morbidity and mortality. The study conducted by Chang\ et al.\ supported that Angpt2 was positively associated with baPWV, a clinical marker of arterial stiffness, in CKD patients not on dialysis. However, the significant linear correlation between Angpt2 and baPWV was not consistent in our CKD patients after adjusting well-known variables. The present results show that baPWV seemed to increase from Angpt2 quartile 1 to 3, but decreased in Angpt2 quartile 4. We supposed that poor renal function and inflammation might interfere in the association of Angpt2 with baPWV. Additionally, David\ et al.\ has speculated a better relationship between Angpt2 and coronary artery disease than those between Angpt2 and peripheral artery disease in patients with CKD stage 4 and dialysis\(^ {23}\). The dissimilarity is probable that the alterations in the Ang/Tie axis might be different between coronary arterial endothelium and peripheral arterial endothelium\(^ {24}\). Further study is needed to investigate the mechanism of association between Angpt2 and the different vascular beds for these inconsistent results.

Among angiopoietin families, Angpt2 had been more strongly related to increased risks for cardiovascular burdens than Angpt1 in previous clinical reports\(^ {3,4,13}\). The information of Angpt1 in subclinical cardiovascular disease is limited. We demonstrated the novel finding of a negative association between Angpt1 and LAD, as a predictor of adverse cardiovascular outcome in CKD patients\(^ {35}\). The balance between Angpt1 and Angpt2 regulates vascular development and remodeling\(^ {1}\). Angpt1 could restrain inflammatory process and abnormal angiogenesis\(^ {1}\). Decreased Angpt1 might have a potential influence on the change of cardiac structure. Conversely, there
was no significant correlation between VEGF-A and cardiovascular structure in our CKD cohort. Nevertheless, VEGF-A still has played a role in modulation of Angpt2-mediated signaling in pathophysiology.

Some limitations must be considered in this study. Firstly, serum markers of angiogenesis and echocardiography parameters were measured once at enrollment. The effect of the time-varying serum markers of angiogenesis levels on cardiac structure and function might be underestimated. Although we found the significant association of Angpt2 with cardiac hypertrophy and correlation between Angpt1 and LAD, the p-values for trend seemed to be low. The relative small sample size probably diminished the statistic power. Nevertheless, Angpt2 and Angpt1 seemed to be a potential marker of cardiac remodeling.

In conclusion, our study demonstrates that high circulating Angpt2 is associated with increased risks for LVH and Angpt1 was negatively correlated with LAD in stages 3–5 CKD patients. These findings promote the notion that Angpt2 and Angpt1 were potential biomarkers of cardiac remodeling in CKD patients. Future studies will be necessary to investigate the pathogenic role of Angpt2 and Angpt1 in cardiac structure and function.

Methods
Study Samples. This observational study enrolled 270 CKD stages 3–5 patients, who had been enrolled in our integrated CKD program for more than 3 months. These patients were scheduled for a study interview after they have given and written informed consent between 2006 and 2011 at Kaohsiung Medical University Hospital in Southern Taiwan. CKD was staged based on Kidney Disease Outcomes Quality Initiative (K/DOQI) guidelines. The eGFR was calculated according to Levey et al. This study protocol was approved by the Institutional Review Board of the Kaohsiung Medical University Hospital (KMUH-IRB-990198). The methods were carried out in accordance with the relevant guidelines, including any relevant details.

Clinical measurements. Information on socio-demographic characteristics and clinical data including age, gender, cigarette smoking, alcohol, medication, and co-morbidities were obtained by medical records and interviews with patients at enrollment. Diabetes was defined by self-report, use of anti-diabetic drugs, or blood glucose values according to the American Diabetes Association criteria. Hypertension was defined as patients with a history of hypertension, taking antihypertensive drugs, or the measurement of blood pressure ≥140/90 mmHg. Heart disease was defined as a history of myocardial infarction, ischemic heart disease or congestive heart failure. Information regarding patients’ medications, such as angiotensin converting enzyme inhibitors (ACEI), angiotensin II receptor blockers (ARB), β-blockers, and calcium channel blockers before and after enrollment was collected from medical records. Blood pressure was measured in the seated position after 5 min rest, using one single calibrated device. Blood pressure was recorded as the mean of three consecutive measurements with 5 min intervals.

Laboratory measurements. Fasting blood samples were collected at enrollment for the biochemistry study, including hemoglobin, glycated hemoglobin, albumin, phosphate, calcium, uric acid, cholesterol, triglyceride, hsCRP, and parathyroid hormone. The markers of angiogenesis including Angpt1, Angpt2, and VEGF-A levels were measured using commercial enzyme-linked immunosorbent assays (R&D Systems Inc). The sensitivity of Angpt1, Angpt2, and VEGF-A assay was 10.3, 0.12, and 5 pg/ml. The interassay and intraassay coefficients of variation of Angpt1, Angpt2, and VEGF-A assay was 10.3, 1.20, and 5 pg/ml. All assays were assessed in duplicate by investigators blinded to the characteristics and clinical outcome of patients.

Evaluation of cardiac structure and function. The echocardiographic examination was performed at the same time as blood and urine sample collection by experienced cardiologists using VIVID 7 (General Electric Medical Systems, Horten, Norway), while patients breathed quietly in the left decubitus position. The cardiologists were blind to the clinical characteristics and laboratory data of patients. Two-dimensional and M-mode images were recorded from the standardized views. The echocardiographic measurements including LAD, AoR, left ventricular posterior wall thickness in diastole and systole (LVPWTd and LVPWTSs), left ventricular internal diameter in diastole and systole (LVIDd and LVIDs), interventricular septal wall thickness in diastole and systole (IVSd and IVDs), E, and A were obtained. E/A ratio less than one was defined as diastolic dysfunction. Left ventricular systolic function was assessed according to LVFS and LVEF. In addition, left ventricular mass (LVM) was calculated using Devereux-modified method, while LVMi was calculated by dividing left ventricular mass by body surface area. The definition of LVH was based on the 2007 European Society of Hypertension/European Society of Cardiology guidelines. Left ventricular relative wall thickness (LVRWT) was calculated as the ratio of 2 × LVPWTd/LVIDd. Concentric LVH was defined as LVMI ≥125 g/m² in men and ≥110 g/m² in women, with LVRWT ≥0.45; eccentric LVH was defined as LVMI ≥125 g/m² in men and ≥110 g/m² in women, with LVRWT < 0.45. Stroke volume was calculated using measurements of ventricle volumes and subtracting the volume of the blood in the ventricle at the end of a beat (end-systolic volume) from the volume of blood just prior to the beat (end-diastolic volume). Based on the unmeasured E/A ratio in atrial fibrillation, this study excluded CKD patients with atrial fibrillation.

Assessment of brachial-ankle pulse wave velocity. BaPWV, as a fine parameter of arterial stiffness, was measured by the ankle-brachial index (ABI)-form device (Colin VP1000, Komaki, Japan) at enrollment. This device automatically and simultaneously measured blood pressures in both arms and ankles using an oscillometric method. Pulse waves obtained from the brachial and tibial arteries were recorded simultaneously, and the transmission time (ΔTba), which was defined as the time interval between the initial increase in brachial and ankle waveforms, was determined for baPWV measurement. The transmission distance from the brachium to ankle was calculated according to body height. The path length from the suprasternal notch to the brachium (Lb)

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was obtained using the following equation: \( L_b = 0.2195 \times \text{height of the patient (in cm)} - 2.0734 \). The path length from the suprasternal notch to the ankle (\( L_a \)) was obtained using the following equation: \( L_a = (0.8129 \times \text{height of the patient [in cm]} + 12.328) \). Finally, the following equation was used to automatically obtain baPWV: \( \text{baPWV} = (L_a - L_b)/\Delta T_{ba} \). The higher baPWV value was used to represent arterial stiffness for each patient.

**Statistical Analysis.** Statistical results of baseline characteristics of all patients were stratified by quartiles of Angpt2, with cutoff values of 1538.2, 1990.7, and 2753.2 mg/ml respectively based on the non-normal distribution of Angpt2. Continuous variables were expressed as mean ± SD or median (25th, 75th percentile) as appropriate. Skewed distribution of continuous variables, such as Angpt2, VEGF and LVMI was log-transformed to approximate normal distribution. Categorical variables were expressed as percentages. The differences in continuous variables between groups were evaluated by one-way analysis of variance (ANOVA) or the Kruskal-Wallis H test as appropriate. Bonferroni post hoc analysis was also used for evaluation between two groups after ANOVA or the Kruskal-Wallis H test. The differences in categorical variables between groups were tested by Chi-square test. Furthermore, the Spearman correlation and adjusted partial correlation were performed to analyze the correlation between serum markers of angiogenesis, baPWV and echocardiography parameters. Univariate and multivariate linear regression and logistic regression were utilized to investigate the association between serum markers of angiogenesis and echocardiography parameters. Adjusted variables included age, sex, smoking history, diabetes mellitus, heart disease, eGFR, and serum hemoglobin and cholesterol levels. Statistical analyses were conducted using SPSS 18.0 for Windows (SPSS Inc., Chicago, Illinois). Statistical significance was set at a two-sided p-value of less than 0.05.

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Author Contributions
Dr. Yi-Chun Tsai was in charge for data collection and preparing the manuscript and Figure. Drs Mei-Chuan Kuo, Yi-Wen Chiu, Hung-Tien Kuo, Chee-Siong Lee, Professor Shang-Jyh Hwang, and Professor Hung-Chun Chen contributed to the concept of this study especially in the field of medical utilizations and statistics. Miss Su-Chu Lee helped Dr. Tsai in data collection and the experimental works. Dr. Mei-Chuan Kuo created the idea of this study, helping Dr. Tsai in manuscript preparation, and takes responsibility for the integrity of the data and accuracy of the data analysis. All authors reviewed the manuscript.

Additional Information
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