Circulating ANGPTL2 Levels Increase in Humans and Mice Exhibiting Cardiac Dysfunction

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Background: Recently, it was reported that angiopoietin-like protein 2 (ANGPTL2) secreted from a pathologically stressed heart accelerates cardiac dysfunction in an autocrine/paracrine manner, and that suppression of ANGPTL2 production in the heart restored cardiac function and myocardial energy metabolism, thereby blocking heart failure (HF) development. Interestingly, circulating ANGPTL2 concentrations reportedly increase in HF patients, suggesting a possible endocrine effect on cardiac dysfunction. However, it remains unclear why circulating ANGPTL2 increases in those subjects and whether circulating ANGPTL2 alters cardiac function in an endocrine manner.

Methods and Results: It was found that circulating ANGPTL2 levels are positively correlated with left atrial diameter and pulmonary capillary wedge pressure, and are inversely proportional to the percent of ejection fraction in patients with dilated cardiomyopathy. Furthermore, in mice, circulating ANGPTL2 concentrations increased as HF developed following transverse aorta constriction (TAC), and were inversely correlated with the percent of fractional shortening. Interestingly, although circulating ANGPTL2 concentrations significantly increased in transgenic mice overexpressing keratinocyte-derived ANGPTL2, no pathological cardiac remodeling was seen. Furthermore, it was observed that there was no difference in HF development between transgenic mice and controls following TAC surgery.

Conclusions: Circulating ANGPTL2 levels increase in subjects experiencing cardiac dysfunction. However, circulating ANGPTL2 does not promote cardiac dysfunction in an endocrine manner, and increased levels of circulating ANGPTL2 seen during HF are a secondary effect of increased ANGPTL2 secretion from stressed hearts in HF pathologies.

Key Words: ANGPTL2; DCM; Heart failure; Mouse TAC model

Heart failure (HF) is a leading cause of mortality in the developed world, with a 5-year mortality rate of approximately 50%, a rate equal to that of many cancers. Recently, the number of HF patients has rapidly increased to an estimated 26 million people worldwide, an outcome now recognized as a HF pandemic. HF prevalence increases among older people due to cumulative effects of cardiovascular risk factors over a lifetime, and HF represents a major public health problem in developed countries. Notably, Japan has the highest proportion of older people in the world: 26.3% of the population were aged 65 years or older in 2015 (www.worldbank.org). Accordingly, a HF pandemic in Japan is predicted by 2030. Therefore, clarification of molecular mechanisms underlying HF development is necessary to develop effective therapeutic strategies to combat this outcome.

Angiopoietin-like protein 2 (ANGPTL2), a member of the ANGPTL family, is a secreted glycoprotein that contributes to tissue homeostasis by enhancing tissue remodeling in an autocrine/paracrine manner, thereby promoting chronic inflammation and subsequent irreversible pathological tissue remodeling, resulting in various lifestyle- or aging-associated diseases, such as obesity, metabolic syndrome, type 2 diabetes, atherosclerotic vascular disease, and some cancers. Several studies report that ANGPTL2 is expressed...
ANGPTL2 in circulation itself did not promote cardiac dysfunction in these mice, suggesting that increased levels of circulating ANGPTL2 seen under HF conditions are a secondary effect of increased ANGPTL2 secretion from a stressed heart in HF pathologies.

**Methods**

**Human Studies**

Sixty-three consecutive patients with dilated cardiomyopathy (DCM) (42 men and 21 women; mean age ± SEM, 54.8 ± 12.5 years; Table 1) were enrolled in the study. Twenty-six were classified as New York Heart Association (NYHA) class I, 31 as class II, and 6 as class III. Individuals with an episode of acute HF within the previous 3 months or with renal dysfunction (estimated glomerular filtration rate [eGFR] of <30 mL/min/1.73 m²) were excluded from the study. All subjects underwent coronary angiography to exclude coronary artery disease, as well as an endomyocardial biopsy to exclude myocarditis or specific muscle disease. DCM was defined as the presence of both a left ventricular (LV) ejection fraction (EF) of <50% (as revealed by contrast left ventriculography) and a dilated LV cavity in the absence of coronary artery stenosis of >50%, valvular heart disease, arterial hypertension, and secondary cardiac muscle disease attributable to any known systemic condition. No patient had a history of acute viral myocarditis or familial DCM. There was also no evidence that immune triggers influenced DCM.

<p>| Table 1. Clinical Characteristics of Patients With Dilated Cardiomyopathy |</p>
<table>
<thead>
<tr>
<th>Variable</th>
<th>n (%) or median (IQR); range</th>
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<tr>
<td>n</td>
<td>63</td>
</tr>
<tr>
<td>Age</td>
<td>58 (44–64); 30–75</td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>42/21</td>
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<tr>
<td>NYHA (I/II/III)</td>
<td>26/31/6</td>
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<tr>
<td>BNP (pg/mL)</td>
<td>89.75 (52.25–281.25); 3.9–1,515.9</td>
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<tr>
<td>Ultrasonic echocardiographic analysis</td>
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<tr>
<td>EF (%)</td>
<td>30 (24–40); 13–55</td>
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<tr>
<td>LVDd (mm)</td>
<td>62 (56–66); 46–88</td>
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<tr>
<td>LVDs (mm)</td>
<td>51 (45–56); 35–83</td>
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<tr>
<td>LAD (mm)</td>
<td>40 (35–46); 23–62</td>
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<tr>
<td>E/A</td>
<td>2.58 (1.74–3.39); 0.70–5.36</td>
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<tr>
<td>LVMI (g/m²)</td>
<td>156 (121–199); 94–409</td>
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<tr>
<td>Cardiac catheterization analysis</td>
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<tr>
<td>HR (beats/min)</td>
<td>77 (69–90); 52–117</td>
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<tr>
<td>CO (L/min)</td>
<td>4.45 (3.70–5.325); 1.90–7.70</td>
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<tr>
<td>CI (L/min/m²)</td>
<td>2.6 (2.2–3.1); 1.2–4.3</td>
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<td>sBP (mmHg)</td>
<td>120.5 (105.75–137); 79–182</td>
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<tr>
<td>dBP (mmHg)</td>
<td>74.5 (64.75–87); 50–107</td>
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<tr>
<td>PCWP (mmHg)</td>
<td>11 (7–14); –2 to 28</td>
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<tr>
<td>sPA (mmHg)</td>
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<td>dPA (mmHg)</td>
<td>20 (7–14); –1 to 30</td>
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<tr>
<td>RA (mmHg)</td>
<td>6 (3–8); –2 to 13</td>
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Values are presented as mean ± SEM or number (n) and percentage (%) for categorical variables. BNP, brain natriuretic peptide; CI, cardiac index; CO, cardiac output; dBP, diastolic blood pressure; DCM, dilated cardiomyopathy; dPA, diastolic pulmonary artery pressure; E/A, ratio of peak early mitral inflow velocity to peak late diastolic mitral inflow velocity during atrial contraction; EF, ejection fraction; HR, heart rate; LAD, left atrial dimension; LVDd, left ventricular end-diastolic dimension; LVDs, left ventricular end-systolic dimension; LVMI, left ventricular mass index; NYHA, New York Heart Association; PCWP, pulmonary capillary wedge pressure; RA, mean atrial pressure; sBP, systolic blood pressure; sPA, systolic pulmonary artery pressure.
development in any patient. Patients were in a stable condition before referral to Nagoya University Hospital for cardiac catheterization. Written informed consent was obtained from each patient before cardiac catheterization, and the study was approved by the Human Ethics Committee of the Nagoya University School of Medicine, Japan (protocol approval No. 359-7; March 16, 2015).

**Cardiac Catheterization Analysis**

All patients underwent diagnostic right and left heart catheterization, as described previously. In brief, pulmonary arterial wedge pressure (PCWP) and cardiac output (CO) were measured using a Swan-Ganz catheter inserted through the right internal jugular vein. Cardiac index (CI) was calculated as follows: CI = CO per body surface area (L/min/m²). Central aortic systolic and diastolic blood pressure were measured using a 6F fluid-filled pigtail catheter inserted through the right radial or femoral artery. Serum ANGPTL2 levels were determined using human ANGPTL2 ELISA kits (IBL, Fujioka, Japan).

**Ultrasonic Echocardiographic (UCG) Analysis of Human Patients**

Two-dimensional UCG analysis was performed using a ViVid 7 system (ViVid 7; GE Healthcare, Milwaukee, WI, USA), as described previously. Analysis included measurement of LV end-diastolic dimension (LVDd), LV endsystolic dimension (LVDs), and left arterial dimension (LAD). Percentage of EF (%EF) was calculated from LVDd and LVDs. LV mass index (LVMi) was calculated from two-dimensional measurements based on a formula approved by the American Society of Echocardiology.

**Animal Studies**

All experimental procedures were approved by the Kumamoto University Ethics Review Committee for Animal Experimentation. Mice were fed a normal diet (ND; CE-2; CLEA, Tokyo, Japan), bred in a mouse house with automatically controlled lighting (12 h on, 12 h off), and maintained at a stable temperature of 23°C. Genetically engineered mice used in the study were: Angptl2 conditional knockout mice (Angptl2floxed/floxed) on a C57BL/6N background, transgenic (Tg) mice overexpressing Cre driven by the murine aMHC promoter (aMHC-Cre) on a C57BL/6N background, and Tg mice overexpressing Angptl2 in keratinocytes (K14-Angptl2) on an FVB/N background. The db/db, KK-Ay, and KK mice were purchased from CLEA Japan.

**Establishment of Hypertrophy Mouse Models**

Male mice approximately 10 weeks old (body weight of 23–25 g) were subjected to pressure overload using transverse aorta constriction (TAC) surgery, as described previously. Echocardiography analysis was performed after TAC surgery, as described previously.

**Histological Analysis**

Histological analysis was performed, as described previously. In brief, mouse heart tissue sections were stained with hematoxylin and eosin (H&E; Wako, Osaka, Japan) to evaluate morphology or Masson’s Trichrome (Muto Pure Chemicals, Tokyo, Japan) to evaluate cardiac fibrosis. The fibrotic area was calculated, using Image J software (National Institutes of Health), as the summation of blue-stained areas divided by total ventricular area. Slides were mounted and examined using a BIORÉVO BZ-9000 microscope (Keyence, Osaka, Japan).

**Real-Time Quantitative Reverse-Transcription-Polymerase Chain Reaction (RT-PCR) Analysis**

Real-time quantitative RT-PCR analysis was performed, as described previously. Relative transcript abundance was normalized to that of 18S rRNA levels in mouse and human samples. Primer sets used for RT-PCR are listed in Table S1.

**Western Blot Analysis**

Western blot analysis was performed, as described previously. In brief, total protein (20 µg) or serum (0.1 µl) proteins were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to poly vinylidene fluoride (PVDF) membranes. Membranes were incubated with anti-AKT (#9272S), anti-p-AKT (s473) (#9271), anti-p-AKT (T308) (#40656; Cell Signaling Technology, Danvers, MA, USA), or anti-Serca2a (ab3625; Abcam, Cambridge, MA) antibodies diluted 1:1,000 at 4°C overnight. After washing with phosphate-buffered saline with 0.1% Tween 20 (PBST), membranes were incubated with 1,2,000 diluted horseradish peroxidase (HRP)-conjugated sheep anti-rabbit IgG (GE Healthcare Life Science, Piscataway, NJ, USA) antibodies at room temperature for 60 min. For Angptl2 immunoblotting, membranes were incubated with biotinylated goat anti-Angptl2 antibody (BAF1444; R&D Systems, Minneapolis, MN, USA) diluted 1:3,000 at 4°C overnight. After washing with PBST, membranes were incubated with HRP-conjugated streptavidin (Thermo Fisher Scientific Inc., Waltham, MA, USA) at room temperature for 60 min. Internal controls were mouse anti-Hsc70 (sc-7298; Santa Cruz Biotechnology, Santa Cruz, CA, USA) and diluted HRP-conjugated sheep anti-mouse IgG (GE Healthcare Life Science) antibodies (both diluted 1:2,000), which served as primary and secondary antibodies, respectively. Hsc70 was used for normalization.

**Recombinant Adeno-Associated Virus (AAV) Treatment**

Recombinant AAV treatment was performed, as described previously. In brief, production and purification of recombinant AAV6 vectors were conducted at Takara Bio Inc. In brief, for shRNA synthesis, single-stranded DNA oligos harboring mouse Angptl2-targeting siRNA and complementary strands were designed as follows: top 5'-CTAGAGCCAGAAGAGGGAGATCTATATA GTGCTCCCTGTGTGTATAGTACTCGCTTTCTGTGCTTTTTTTA-3' and bottom 5'-CTAGTAAAAAGGCCAGAAAAGCGAGGTACTATATA CACTATATAGTACTCGCTTTCTGTGCTTTTTTTA-3'. The db/db, KK-Ay, and KK mice were purchased from CLEA Japan.
sonic echocardiographically derived parameters, and hemodynamic measurement). For multiple linear regression analyses, a generalized linear model was used. Statistical analyses were performed using JMP Pro 12 software (SAS

Figure 1. Relationship of circulating angiopoietin-like protein 2 (ANGPTL2) levels to heart failure (HF)-associated clinical parameters. Scatter plots show correlation between circulating ANGPTL2 levels and brain natriuretic peptide (BNP) (A), echocardiographically derived parameters (left atrial diameter (LAD), left ventricular end-diastolic dimension (LVEDd), left ventricular end-systolic dimension (LVESd), percent of ejection fraction (EF), and left ventricular mass index (LVMI) (B); right heart catheterization analysis-derived parameters (pulmonary capillary wedge pressure (PCWP), CO (cardiac output), and CI (cardiac index) (C, Upper rows), and left heart catheterization analysis-derived parameters (systolic blood pressure (sBP), diastolic blood pressure (dBP); and C, Bottom rows). Also shown is Pearson’s correlation coefficient for rho.

Statistical Analysis
In human studies, Pearson’s test and correlation coefficients were used to evaluate relationships between serum ANGPTL2 levels and clinical variables (e.g., BNP, ultrasonic echocardiographically derived parameters, and hemodynamic measurement). For multiple linear regression analyses, a generalized linear model was used. Statistical analyses were performed using JMP Pro 12 software (SAS
In animal studies, no statistical methods were used to determine sample size, but sample sizes were similar to those used in previous reports. No exclusion/inclusion criteria were applied to mice used in this study. Group allocation and outcome assessment were performed in a blinded manner. In vitro experiments were repeated at least three times. All values were reported as mean±SEM. Data were assessed with 2-group comparisons of variables using an unpaired 2-tailed t-test, with multiple tests.

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<th>Parameters</th>
<th>β (95% CI)</th>
<th>P value</th>
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<tr>
<td>LAD</td>
<td>5.830 (0.921, 10.740)</td>
<td>0.0207</td>
</tr>
<tr>
<td>EF</td>
<td>−5.685 (−12.225, 0.855)</td>
<td>0.0872</td>
</tr>
<tr>
<td>PCWP</td>
<td>3.912 (0.183, 7.640)</td>
<td>0.0401</td>
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Table 2. Generalized Linear Regression Analysis (Adjusted for Age and Sex)

Abbreviations as in Table 1.

Figure 2. Circulating ANGPTL2 levels increase in mice following transverse aorta constriction (TAC) surgery. (A) Echocardiographically left ventricular end-diastolic internal diameter (LVID;d; Left), diastolic left ventricular posterior wall thickness (LVPW;d; Middle), and percent fractional shortening (FS%; Right) before surgery (0W) and 2, 4, and 6 weeks after surgery in sham and TAC-operated wild-type (WT) mice (n=8–10 per group). (B) Representative images of a heart 6 weeks after sham surgery and 2, 4, and 6 weeks after post-TAC surgery (Left). Calculation of heart weight/body weight (HW/BW) ratio (Right) at 2, 4, and 6 weeks after sham or TAC surgery (n=8–10 per group). Scale bar: 5 mm. (C) Relative expression of mRNAs associated with HF and cardiac fibrosis. Sham-operated control values were set to 1 (n=8–10 per group). (D) Enzyme-linked immunosorbent assay (ELISA) for circulating ANGPTL2 protein before surgery and at 2, 4, and 6 weeks after surgery in sham and TAC-operated WT mice (n=8–10 per group). (E) Representative western blots and quantitation of serum ANGPTL2 protein from WT mice 6 weeks after TAC or sham surgery (n=13 per group). Albumin served as the loading control. Sham-operated control values were set to 1. (F) Scatter plots showing the correlation between serum ANGPTL2 protein and %FS from WT mice at 2, 4, and 6 weeks after TAC (n=8–10 per group). Data are presented as mean±SEM. Statistical significance was determined by using a Student’s t-test. *P<0.05, **P<0.01, †P<0.001 between groups. Abbreviations as in Figure 1.
P<0.05 was considered statistically significant. Comparisons by 1- or 2-way ANOVA with Tukey’s multiple comparisons test between each group. A value of P<0.05 was considered statistically significant.

Results

Circulating ANGPTL2 Levels Correlate With LAD and PCWP in Patients With DCM
To determine whether circulating ANGPTL2 levels were correlated with cardiac function, we assessed brain natriuretic peptide (BNP) levels and undertook UCG analysis and hemodynamic measurement in DCM patients (Figure 1A–C). The study population consisted of 63 patients with DCM, and their characteristics are shown in Table 1. Interestingly, LAD and PCWP were positively correlated with circulating ANGPTL2 levels (r=0.268, P=0.034 and r=0.324, P=0.012, respectively), while %EF was inversely correlated with those levels (r=−0.257, P=0.046) in DCM patients. The statistically significant positive correlation between circulating ANGPTL2 concentrations and LAD and PCWP remained unchanged after adjustment for age and sex (P=0.0207, P=0.0401, respectively; Table 2). Taken together, we found that circulating ANGPTL2 levels are positively correlated with cardiac dysfunction in patients with DCM.

Circulating ANGPTL2 Levels Increase in Mice Following TAC Surgery
To assess whether circulating ANGPTL2 levels change in a HF mouse model, we measured those levels in the blood of mice showing cardiac dysfunction induced by TAC surgery-dependent pressure overload. After TAC surgery, we evaluated pathological cardiac remodeling by analyzing UCG (LV dimension in end-diastolic phase (LVDd)) and thickness of the LV posterior wall (LVPW), percent of fractional shortening of LV (% of FS), heart weight/body weight (HW/BW) ratio, and expression of markers of HF (ANP, BNP, and Myh7) and cardiac fibrosis (CTGF, CollI, and Col3a1) (Figure 2A–C). Enzyme-linked immunosorbent assay (ELISA) of mouse sera revealed gradually increasing circulating ANGPTL2 concentrations after TAC surgery (Figure 2D). Six weeks after TAC, western blotting also revealed that circulating ANGPTL2 concentrations were significantly increased compared to sham-operated controls (Figure 2E). Circulating ANGPTL2 levels were also inversely correlated with %FS in the TAC model (r^2=0.3716, P<0.001, Figure 2F). These results indicate overall that in mice, circulating ANGPTL2 levels are positively correlated with cardiac dysfunction.

Increased Circulating ANGPTL2 Following TAC Surgery Is Derived From Cardiomyocytes
We next asked which tissues contribute to increased circulating ANGPTL2 levels after TAC surgery in mice. Expression of Angptl2 mRNA in the heart was significantly increased compared to that of sham-operated controls throughout a 2-, 4- and 6-week period of TAC-induced HF development (Figure 3A). We then asked whether lowering ANGPTL2 expression in the heart would decrease circulating ANGPTL2 concentrations after TAC surgery by employing systemic delivery of recombinant adenoviral-associated virus serotype 6 (AAV6) vectors, which preferentially transduce cardiac muscle.20 We then compared circulating ANGPTL2 protein levels after TAC surgery between mice delivered Angptl2 shRNA cassettes to the heart by intravenous injection of a recombinant AAV6 vector harboring Angptl2 shRNA (3×10^10 vg/mouse) and controls similarly injected with scramble control shRNA (3×10^10 vg/mouse). ANGPTL2 protein levels in the hearts of mice receiving scramble control shRNA after TAC surgery were higher than those seen in sham-operated mice, and those increases were blocked in mice receiving Angptl2 shRNA (data not shown), as previously reported.20 Relevant to circulating ANGPTL2, levels seen in scramble control shRNA in TAC-operated mice increased remarkably compared with those in sham-operated mice, whereas increases were significantly attenuated in TAC-operated mice injected with Angptl2 shRNA (Figure 3B). Moreover, we compared circulating ANGPTL2 levels after TAC between cardiomyocyte-specific Angptl2 conditional knockout mice (Angptl2^floxpfllox;αMHC-Cre) and (Angptl2^floxpfloxflox) controls. ANGPTL2 protein levels in the hearts of control mice increased compared with levels in sham-operated mice, and those increases were attenuated in...
ANGPTL2 in Circulation and Heart Failure

ANGPTL2 Protein Levels Increase in Mouse Visceral White Adipose Tissue (vWAT) and Brown Adipose Tissue (BAT) Following TAC Surgery

Increases in circulating ANGPTL2 levels are not completely blocked after cardiomyocyte-specific ANGPTL2 knockdown (Figure 3B), suggesting that other tissues produce ANGPTL2 in these mice. To test this possibility, we compared ANGPTL2 protein levels in various tissues of TAC- and sham-operated mice. Four weeks after surgery, ANGPTL2 protein levels significantly increased in vWAT and BAT, as well as in the hearts of TAC-relative to sham-operated controls (Figure 4A,B), suggesting that mice adipose tissue is an additional source of circulating ANGPTL2 levels following TAC-induced cardiac dysfunction.

Increased Circulating ANGPTL2 Protein Does Not Promote Pathological Hypertrophy or Cardiac Dysfunction

We next asked whether increased circulating ANGPTL2 has an endocrine effect on cardiac dysfunction. As reported previously,16,26 we confirmed that circulating ANGPTL2 protein levels increased approximately 3-fold in transgenic mice overexpressing ANGPTL2 in the skin under the control of the K14 promoter (K14-Angptl2 Tg mice) compared to those seen in wild-type controls (Figure 5A). In contrast, cardiac ANGPTL2 protein levels were comparable in both groups (Figure 5B). Interestingly, the increased ANGPTL2 levels in circulation were comparable to those in wild-type mice with HF pathologies.

We recently reported that ANGPTL2 secreted from cardiomyocytes perturbs cardiac function by inactivating the AKT-SERCA2a Ca2+ handling cascade in transgenic mice overexpressing ANGPTL2 in cardiomyocytes under control of the αMHC promoter (αMHC-Angptl2 Tg mice).20 Therefore, we questioned whether circulating ANGPTL2 has an endocrine effect on that activity. Western blotting analysis revealed that the AKT-SERCA2a Ca2+ handling cascade was not inactivated in K14-Angptl2 Tg mice (Figure 5C). Furthermore, analysis of UCG and HW/BW, as well as histology and RT-PCR, revealed no differences in cardiac phenotypes or marker expression between K14-Angptl2 Tg and wild-type controls with or without TAC (Figure 5D–H), suggesting that increased ANGPTL2 levels in circulation itself does not have effect on cardiac remodeling in HF pathologies.

Given that adipose tissues also contribute to increased circulating ANGPTL2 levels (Figure 4), we questioned whether adipose tissue-derived ANGPTL2 protein promoted cardiac dysfunction. We previously reported that diet-induced obese mice showed increased ANGPTL2 expression in adipose tissue and increased levels of
Figure 5. Circulating ANGPTL2 protein has no effect on cardiac function or development of cardiac hypertrophy in K14-ANGPTL2 Tg mice. (A) Representative western blots and quantification of serum ANGPTL2 protein levels in K14-Angptl2 Tg or WT littermate mice (n=5–6 per group). Albumin serves as a control. WT values were set to 1. (B) Representative western blots and quantification of ANGPTL2 protein and ANGPTL2 mRNA levels in hearts of K14-Angptl2 Tg and WT mice. Hsc70 serves as a control. WT values were set to 1 (n=6 per group). (C) Representative immunoblot and quantitation of p-AKT, AKT and SERCA2A protein levels in hearts of K14-Angptl2 Tg and WT mice (n=6 per group). Hsc70 serves as a control. Values from WT were set to 1. (D) Shown are representative M-mode echocardiography recordings from K14-Angptl2 Tg and WT littermate mice 4 weeks after sham surgery (Upper row) or TAC (Bottom row) surgery. (E) Echocardiographic quantitation of LVPW;d (Left), LVID;d (Middle), and fractional shortening (FS; Right). (F) Heart weight/body weight (HW/BW) ratio (Left) and lung weight/body weight (LW/BW) ratio (Right) of K14-Angptl2 Tg or WT littermate mice. (G) Sections of Masson's Trichrome-stained heart tissue from K14-Angptl2 Tg or WT littermate mice 4 weeks after sham surgery (Upper row) or TAC (Bottom row) surgery. Scale bar: 100 μm. (H) Relative expression of mRNAs associated with HF and cardiac fibrosis (n=6 per group). WT values were set to 1. Data are presented as mean±SEM. Statistical significance was determined by using a Student’s t-test. *P<0.05 between genotypes. Abbreviations as in Figures 1, 2.
circulating ANGPTL2 protein. Here, using 2 models of genetically obese diabetic mice (db/db and KK-Ay), we observed elevated ANGPTL2 protein levels in circulation compared with controls (Figure 6A,E), whereas ANGPTL2 protein levels in heart tissues of both models were comparable to controls (Figure 6B,F). UCG analysis revealed no cardiac hypertrophy, ventricular dilatation, or cardiac dysfunction in db/db or KK-Ay mice compared with non-obese controls (Figure 6C,G). Moreover, in both models, ANP, BNP, and Myh7 expression in the heart was unchanged relative to the controls (Figure 6D,H). We conclude that increased ANGPTL2 protein derived from adipose tissue does not promote cardiac dysfunction, which also suggests no endocrine effects of ANGPTL2 in circulation on cardiac function.

Discussion

This study demonstrates that circulating ANGPTL2 levels positively correlate with cardiac dysfunction in DCM patients and in a TAC-induced mouse HF model. Our study also demonstrates that, in mice, increased ANGPTL2 lev-
els are primarily due to secretion by cardiomyocytes in conditions of TAC-induced cardiac dysfunction, although adipose tissues also contribute to these effects. Our findings indicate that increased ANGPTL2 in circulation likely does not cause or accelerate pathological hypertrophy or cardiac dysfunction in mice, but is rather a result of increased secretion from stressed heart in HF pathologies.

Recently, Huang et al reported that circulating ANGPTL2 levels in HF patients were higher than those in control individuals, and that cardiac function was inversely correlated with levels of ANGPTL2 circulating in the periphery. These authors noted that circulating ANGPTL2 levels paralleled levels of NT-pro BNP and of LAD as measured by UCG, and that those parameters were inversely correlated with %EF, as measured by UCG. In this study, we also observed a positive correlation of circulating ANGPTL2 levels with LAD and an inverse correlation of those levels with %EF in DCM patients. Taken together, circulating ANGPTL2 concentrations increase in subjects with cardiac dysfunction, and may correlate with cardiac dysfunction not only in DCM patients but in patients showing other types of HF, although larger-scale patient studies are required for confirmation. Moreover, additional investigations are needed to determine whether circulating ANGPTL2 concentrations decrease as cardiac dysfunction improves.

A recent study by our group demonstrates that pathological stimuli, such as hypertension, activate ANGPTL2 production in cardiomyocytes, and that heart-derived ANGPTL2 directly contributes to increased circulating ANGPTL2 in HF conditions. Interestingly, the present study revealed that cardiomyocytes are a major source of circulating ANGPTL2 protein in mice after TAC surgery. However, suppression of cardiomyocyte-derived ANGPTL2 in mice did not lower circulating ANGPTL2 levels to baseline levels seen before TAC surgery (Figure 3C). Relevant to this, it is noteworthy that ANGPTL2 protein production significantly increased in mouse vWAT and BAT, as well as heart tissues after TAC surgery relative to sham-operated controls, suggesting that after TAC, adipose tissue contributes to increased levels of circulating ANGPTL2. More interestingly, ANGPTL2 protein production was significantly decreased in mice lung and liver after TAC surgery relative to sham-operated controls (Figure 4A); however, mechanisms underlying this change remain unclear. We conclude that increased levels of circulating ANGPTL2 after TAC are likely secondary phenomena caused by altered ANGPTL2 secretion from various organs as a result of HF, and that increased ANGPTL2 levels are not an adequate biomarker for HF.

ANGPTL2 is an adipocyte-derived inflammatory mediator that promotes chronic adipose tissue inflammation and systemic insulin resistance. Thus, it is noteworthy that ANGPTL2 expression in adipose tissue increases when HF occurs and is promoted by TAC. This outcome might be exacerbated by induction of adipose tissue inflammation and subsequent resistance in HF conditions, an idea supported by observations that circulating ANGPTL2 levels increase in subjects with adipose tissue inflammation and/or insulin resistance. Overall, given these findings, it would be useful to determine whether ANGPTL2 mediates adipose tissue inflammation and subsequent systemic insulin resistance in patients with HF. In contrast, insulin resistance and/or diabetes predisposes patients to HF, creating a vicious cycle of HF and anomalous glucose metabolism that likely accelerates progression of both conditions and is potentially exacerbated by ANGPTL2. This possibility should be addressed in future studies.

Here, we found that circulating ANGPTL2 levels in sham-operated mice injected with AAV6-shANGPTL2 were comparable to that of shRNA controls, as anticipated (Figure 3B). We also found that circulating ANGPTL2 levels in sham-operated Angptl2 Flox/Flox; αMHC-Cre mice were comparable to those seen in sham-operated Angptl2 Flox/Flox; αMHC-Cre mice (Figure 3C), suggesting that cardiomyocyte-secreted ANGPTL2 has a lower contribution to circulating ANGPTL2 concentrations in mice without HF.

Our recent study demonstrated that ANGPTL2 secreted from cardiomyocytes perturbs LV contractile ability by inactivating AKT-SERCA2a signaling and decreasing myocardial energy metabolism in αMHC-Angptl2 Tg mice. However, in that paper, it has not been clarified whether ANGPTL2 affects cardiomyocyte in an endocrine or an autocrine/paracrine manner, because ANGPTL2 protein is a secreted glycoprotein. In this study, we clarified that ANGPTL2 affects cardiomyocytes in an autocrine/paracrine manner, but does not show an endocrine effect on cardiomyocytes. This is consistent with our findings of the in vitro cardiomyocyte experiment where effective ANGPTL2 protein concentration levels were more than 5 μg/mL and greatly exceeded circulating ANGPTL2 protein concentration levels (~1.2 ng/mL) in mice with HF. Thus, increased levels of circulating ANGPTL2 seen in mice with HF are a secondary effect of increased ANGPTL2 secretion from stressed hearts in HF pathologies.

In the present study, we show that based on analysis of K14-Angptl2 Tg mice and genetically obese diabetic mice, circulating ANGPTL2 alone is not sufficient enough to induce a pathological cardiac response. An alternative way to address this question would be to chronically infuse mice with the ANGPTL2 protein to assess a potential endocrine effect on heart function; unfortunately, we were unable to prepare sufficiently high concentrations of recombinant ANGPTL2 protein required to conduct such an analysis. Nonetheless, further investigations of this type may be required to assess potential endocrine effects of ANGPTL2 on cardiac function.

In summary, circulating ANGPTL2 levels increase in HF patients and mouse models of HF. Those increases are primarily due to enhanced ANGPTL2 secretion from cardiomyocytes, although adipose tissue-derived ANGPTL2 also contributes to those increases. However, we observed no endocrine effect of circulating ANGPTL2 on cardiac function, which suggests that increased circulating ANGPTL2 levels in HF patients reflect, rather than promote, cardiac dysfunction.

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**Disclosures**

None.
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References


Supplementary Files

Supplementary File 1

Table S1. Primer sequences used in quantitative RT-PCR

Please find select supplementary file(s): http://dx.doi.org/10.1253/circj.CJ-17-0327