Research paper

Lipidomics reveals accumulation of the oxidized cholesterol in erythrocytes of heart failure patients

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A R T I C L E   I N F O

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A B S T R A C T

Lipids play an important role in the pathogenesis of cardiovascular disease. Changes in lipids of erythrocytes are indicative of the outcome of pathophysiological processes. In the present study, we assessed whether the lipid profiles of erythrocytes from heart failure (HF) patients are informative of their disease risk. The lipidomes of erythrocytes from 10 control subjects and 29 patients at different HF stages were analyzed using liquid chromatography time-of-flight mass spectrometry. The lipid composition of erythrocytes obtained from HF patients was significantly different from that of normal controls. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins decreased in HF erythrocytes as compared with those of control subjects; however, the levels of lysoPCs, lysoPEs, and ceramides increased in HF erythrocytes. Notably, the oxidized cholesterol 7-ketocholesterol (7KCh) accumulated to higher level in HF erythrocytes than in plasma from the same patients. We further validated our findings with a cohort of 115 subjects of control subjects (n=28) and patients (n=87). Mechanistically, 7KCh promoted reactive oxygen species (ROS) formation in cardiomyocytes; and induced their death, probably through an ATF4-dependent pathway. Our findings suggest that erythrocytic 7KCh can be a risk factor for HF, and is probably implicated in its pathophysiology.

1. Introduction

Cardiovascular disease is a major health problem and the leading cause of death globally. Cardiac function deterioration hampers the ability of the heart to support blood circulation, resulting in heart failure (HF). The pathogenic mechanism leading to this end stage is complicated. Myocardial infarction, hypertension, cardiomyopathy, valvular heart disease, and inflammation-induced oxidative stress are known risk factors for disease progression [1,2].

Several biochemical pathways, including the pentose phosphate pathway, anaplerotic metabolism, ketone body metabolism, lipotoxic intermediate metabolism, and glycolysis, are affected in patients with HF [3–5]. Changes in metabolites have been identified in plasma and are associated with clinical outcomes in patients with HF [6,7]. These findings suggest that metabolic remodeling in patients may occur during HF progression, and the metabolite profile can thus be used as a biomarker panel for a variety of assessment purposes.

Lipid metabolism alterations have been increasingly demonstrated to underlie the pathogenesis of cardiovascular disease [8]. Currently, research on lipids has focused on the analysis of plasma lipids such as cholesterol, triacylglyceride, and phospholipids [8,9]. Reports seldom indicate specific fatty acids and total cholesterol in erythrocytes as a predictor of cardiovascular events [10–12]. Additionally, erythrocytes are involved in reverse cholesterol transport (RCT), particularly in the low high-density lipoprotein (HDL) state [13]. Given the relatively long life (approximately 120 days) of erythrocytes, any change in the lipid profile of erythrocyte membrane may reflect pathophysiologic changes associated with disease progression.

Few studies have reported on the comprehensive assessment of the metabolome and lipidome of RBCs [14–16], especially in the scenario...
of HF. The aim of this study was to identify lipid profiles of HF erythrocytes using high-throughput liquid chromatography time-of-flight mass spectrometry (LC-TOF-MS). Our findings suggested that the erythrocyte lipid profiles of patients with HF were significantly different from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects. The levels of phosphatidylcholines (PCs), phosphatidylethanolamines (PEs), and sphingomyelins (SMs) decreased from those of normal subjects.

2. Materials and methods

Chemicals were purchased from Sigma-Aldrich (St. Louis, MO), unless otherwise stated. Antibodies to ATF-4 (11815; Cell Signaling Technology, Danvers, MA, USA), CHOP (sc-7351; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), LC3A/B (12741; Cell Signaling Technology, Danvers, MA, USA), Cleaved Caspase-3 (9664; Cell Signaling Technology, Danvers, MA, USA), Cleaved Caspase-3 (9662; Cell Signaling Technology, Danvers, MA, USA), Actin (A5441, A2103; Sigma-Aldrich, St. Louis, MO, USA), and 7-ketocholesterol (Clone #35 A, Japan Institute for the Control of Aging, Shizuoka, Japan) were purchased from respective vendors.

2.1. Blood sample collection and erythrocyte preparation

Patients with HF were classified as stages A, B, and C according to the American College of Cardiology and the American Heart Association Heart Failure (ACC/AHA HF) classification system [17]: patients in stage A were at high risk and asymptomatic but did not have structural heart disease; patients in stage B had structural heart disease but were asymptomatic; and patients in stage C had been hospitalized due to acute or decompensated chronic HF. Patients aged 39–85 years were enrolled in this study. Patients with hypothyroidism, decompenated liver cirrhosis, systemic lupus erythematosus, or serum creatinine of > 3 mg/dL were excluded. Informed consent was obtained from all patients. The study was designed and performed in accordance with the principles of the Declaration of Helsinki and with approval from the Ethics Review Board of Chang Gung Memorial Hospital.

Blood samples were collected in EDTA-containing tubes before the patients were discharged from the hospital. RBCs were prepared for LC-MS analysis. The plasma level of B-type natriuretic peptide (BNP) was measured in triplicate with the principles of the Declaration of Helsinki and with approval from the Ethics Review Board of Chang Gung Memorial Hospital.

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2.2. Lipidomics analysis by LC-MS

For human erythrocyte lipids, a modified method was employed [19]. In brief, 100 μL of erythrocyte lysates was transferred to a glass tube, and 100 μL of water was added. The sample was vortexed and then placed on ice for 15 min. After 1.1 mL of isopropanol was added, the sample was vortexed 4 times for 30 s. Approximately 700 μL of chloroform was added, and the sample was vortexed 4 times for 30 s again. It was subsequently centrifuged at 700 × g for 30 min at 4 °C. The supernatant was transferred to a new tube, dried under nitrogen gas, and stored at −80 °C. Prior to analysis, the sample was dissolved in 500 μL of isopropanol/acetonitrile/water (2:1:1).

For lipid separation, an ACQUITY CSH C18 (2.1 mm × 100 mm, 1.7 μm) column was used. The column temperature was set at 55 °C and the flow rate at 400 μL/min. Mobile phase buffer A was acetonitrile/water (60/40) with 10 mM ammonium formate and 0.1% formic acid, whereas buffer B was isopropanol/acetonitrile (90/10) with 10 mM ammonium formate and 0.1% formic acid. The initial LC gradient conditions were 40% buffer B, increasing to 70% B within 12.1 min, and then to 99% B for 6 min before re-equilibration for 2 min at 40% B. The lyophilized sample was diluted with 500 μL of isopropanol/acetonitrile/water (2:1:1). Each sample was analyzed in triplicate. MS was performed on a Waters QTOFMS (SYNAPT G1 HDMS, Waters MS Technologies, Manchester, UK) operating in the positive or negative ion mode. The desolvation gas was set at 800 L/h at a temperature of 400 °C; the cone gas was set at 25 L/h; and the source temperature was set at 100 °C. The capillary voltage and cone voltage were set to 3000 and 35 V, respectively. Leucine encephalin was used as the lock mass standard (an [M+H]+ ion at 556.2771 Da in electrospray ionization (ESI) positive mode; an [M+H]+ ion at 554.2615 Da in ESI negative mode).

MS data were processed using MassLynx V4.1 and Progenesis QI software (Waters Corp., Milford, Massachusetts, USA). The intensity of each mass ion was normalized with respect to the total ion count to generate a data matrix that included the retention time, m/z value, and normalized peak area. The multivariate data matrix was analyzed using SIMCA-P software (version 13.0, Umetrics AB, Umea, Sweden).

Significant metabolites were subjected to a database search using an in-house database, the Human Metabolome Database (HMDB) (http://www.hmdb.ca/), or the METLIN Metabolomics Database (metlinscripps.edu/index.php). For the identification of specific metabolites, MS/MS spectra were collected and confirmed by comparison with the spectra of chemical standards or by searching the HMDB and METLIN databases.

2.3. Quantification of erythrocyte and plasma 7KCh by LC-MS-MS

Erythrocyte 7KCh was extracted using a modified method [19] as described above. A sample was dissolved in isopropanol: acetonitrile-water (2:1:1 (v/v/v)) containing the deuterated forms of 7KCh-D7 and cholesterol-D6, which were employed as internal standards for quantification. For analysis of total-form sterols and free-form sterols in plasma, plasma samples (10 μL) were preincubated with or without cholesterol ester hydrolase at 37 °C for 15 min in 10 mM phosphate buffer containing taurocholic acid [20]. The samples were extracted with 500 μL of methanol containing cholesterol-D6 and 7-KCh-D7, and then centrifuged at 12,000 × g at 4 °C for 30 min. The resulting supernatant was collected in a new microtube and dried with nitrogen. The sample was dissolved in 100 μL of 100% methanol and was centrifuged at 12,000 × g at 4 °C for 30 min for 7KCh, cholesterol, lanosterol, lanosterol, and 7-dehydrocholesterol quantification.

All the samples were analyzed using ultra-high-performance LC coupled with Xevo TQ-S MS (Waters Corp.) as previously described with modifications [21]. MS with atmospheric pressure chemical ionization (APCI) was performed in positive-ion multiple-reaction-monitoring mode. The optimized parameters were as follows: corona discharge current at 1 μA; probe temperature at 600 °C; source temperature at 150 °C; and gas flow at 800 L/h. Chromatographic separation was achieved on a pentfluorophenyl (PPF) column (100 mm × 2.1 mm; with particle size of 1.8 μm; Waters Corp.) at 25 °C with eluent A (acetonitrile: water 25:75 (v/v) with 0.1% formic acid) and eluent B (methanol); the flow rate was set at 0.3 mL/min. The gradient profile was as follows: isocratic 70% B, 1 min; linear gradient 70–75.7% B, 8 min; linear gradient 75.7–100% B, 1 min; and 100% B, 2 min. The column was then re-equilibrated for 8 min for the next analysis.

2.4. Cell culture and viability assay

HL-1 atrial myocytes were cultured in fibronectin–gelatin-coated
flasks containing Claycomb medium, supplemented with 10% fetal bovine serum, 100 U/mL of penicillin, 100 µg/mL of streptomycin, 2 mM L-glutamine, and 0.1 mM norepinephrine in a humidified atmosphere containing 5% CO₂ at 37 °C, as previously described [22]. For 7KCh treatment, 2 × 10⁴ cells were incubated with 10, 20, and 50 µM 2µM MitoSOX red or 5 µM MitoSox green or DCFH-DA for 30 min at 37 °C, washed twice with PBS, and trypsinized

ROS formation was analyzed quantitatively by cytometric analysis. HL-1 cells were fixed and permeabilized with 4% paraformaldehyde, 0.1% Triton-X100 in PBS for 2 h. After PBS wash, cells were treated with PBS containing 0.1% Triton X-100 and 5% BSA for 1 h at room temperature. Cells were stained with primary antibody of 7KCh (1:100; Clone #35 A, Japan Institute for the Control of Aging, Shizuoka, Japan) overnight at 4 °C. Secondary antibody, conjugated to Alexa-488 (Thermo Fisher Scientific Inc., Waltham, MA, USA), was added at a 1:200 dilution and incubated for 1–2 h. Hoechst 33342 (Thermo Fisher Scientific Inc., Waltham, MA, USA) was used to stain cellular nuclei.

2.7. Western blotting assay

SDS-PAGE and western blotting were performed as previously described [25]. The cells were rinsed with cold PBS, scraped and collected by centrifugation. Cells were immediately lysed in lysis buffer (20 mM Tris, HCl (pH 8), 1% Triton X-100, 137 mM NaCl, 1.5 mM MgCl₂, 10% glycerol, 1 mM EDTA, pH 8) and centrifuged at 16,000 g to precipitate the ghosts. The erythrocyte membrane was incubated with 7KCh or cholesterol at 37 °C for 4 h, after which the membrane-bound 7KCh or cholesterol in the ghost pellet was diluted to the indicated concentrations for a Western blot assay. Cell viability was determined by a neutral red assay, as previously described [23].

2.8. Statistical analyses

To maximize the number of differentially abundant metabolites among groups, the orthogonal projection to latent structures discriminant analysis (OPLS-DA) model was applied using SIMCA-P software (version 13.0, Umetrics AB, Umea, Sweden). The variable importance in the projection (VIP) value of each variable in the model was calculated to indicate its contribution. S-plots were constructed from the OPLS-DA data. Metabolites were chosen based on their contribution to the variation and correlation within the data set of the control and the stage C HF groups. The VIP values of variables greater than 1.0 are considered significantly different.

Results are expressed as the mean ± SD for continuous variables and as the number (percentage) for categorical variables. Data were compared by two-sample t-tests. A p value of < 0.05 was considered significant.

3. Results

3.1. Baseline characteristics

Initially, a total of 115 subjects were enrolled in this study, comprising 28 normal controls and 87 patients at stages A (n = 29), B (n = 29), and C (n = 29). The baseline characteristics and laboratory data are shown in Table 1. Notable differences in these parameters were observed between patients at stages A, B, and C. Compared with the
normal controls, patients at stage C had remarkably higher glucose, creatinine, and BNP levels, but they had lower total cholesterol, low-density lipoprotein (LDL) cholesterol, HDL cholesterol, sodium, hemoglobin and albumin levels, and lower estimated glomerular filtration rate. The percentage of male subjects was higher in the patient groups.

For the untargeted lipid profile study, sex- and age-matched samples from 10 controls and 29 patients with HF at stage A (n = 10), stage B (n = 9), and stage C (n = 10) were chosen (Supplementary Table S1).

3.2. Changes in lipidomes of erythrocytes in patients with heart failure

To investigate whether the erythrocyte lipid composition differs between patients with HF and normal controls, 39 RBC samples were subjected to untargeted metabolomic analysis using LC-TOF-MS. The typical spectra of erythrocyte extracts were obtained in positive (Fig. 1A) and negative (Fig. 1B) ion modes. After data processing with Progenesis QI, 457 and 438 molecular features obtained in ESI positive and negative modes were extracted, respectively. The data were subjected to SIMCA-P analysis. The derived OPLS-DA score plot showed remarkable separation between the control, stage A, stage B, and stage C groups for data sets obtained in both ESI positive and negative modes (Fig. 1C, E). To identify lipophilic metabolites discriminating between the samples from the normal controls and those from patients in group C, we reanalyzed these data sets. The OPLS-DA plots (Fig. 1D, F) and S-plots (Fig. 1G, I) for the data sets obtained in both ESI positive and negative modes are shown herein. From the S-plots, features with a p (corr) value of > 0.5 or < −0.5 were selected. We observed that 30 and 23 metabolites acquired in both ESI positive and negative modes had VIP scores of > 1.0 and revealed significant differences (p < 0.01) between the patients in stage C and the healthy controls (Fig. 1H, J). Clearly, sterols, phospholipids, and ceramides are important discriminators of the normal controls and patients in stage C.

Fig. 1. Liquid chromatography time-of-flight mass spectrometry-based lipidomics analysis of heart failure (HF) erythrocytes. Erythrocytes from normal control subjects and patients with stages A, B, and C were isolated for time-of-flight mass spectrometry analysis in electrospray ionization (ESI) positive and negative modes. Basal peak chromatograms of patients with HF at different stages and normal control subjects were obtained in ESI positive (A) and negative (B) modes, respectively. The molecular features were identified in samples (n = 39) by using Progenesis QI software and further data processing and statistical analysis were performed through SIMCA-P. Orthogonal partial least squares discriminant analysis (OPLS-DA) was performed for all samples, and the score plots for data sets obtained in ESI positive (C) and negative modes (E) are shown. The data for normal control and patients with stage C were reanalyzed using OPLS-DA, and the score plots for data sets obtained in ESI positive (D) and negative modes (F) are shown. Metabolites with significant differences in abundance in ESI positive and negative modes between normal control and patients with stage C are presented in S-plots (G and I), respectively. The Venn diagrams of the features obtained in ESI positive (VIP > 1.0 and p < 0.01) and ESI negative (VIP > 1.0 and p < 0.01) modes are shown in panels H and J, respectively. The normal control (n = 10), stage A (n = 10), stage B (n = 9), and stage C (n = 10) groups are marked in green, yellow, orange, and red, respectively. The ellipse shown in the model represents the Hotelling T2 with 95% confidence.
as a biomarker for the early identification of subjects at risk of HF. To ensure whether 7KCh accumulates in HL-1 cells, we performed intracellular level of 7KCh in HL-1 cell (Fig. 4E). The 7KCh accumulated from erythrocytes to cause cellular damage.

We observed significant increases in the levels of 7KCh 3.4. Intracellular 7KCh accumulation enhances ROS production and reduces the viability of cardiomyocytes

To examine whether 7KCh affects the viability of HL-1 cells, we performed an immunofluorescence assay with anti-7KCh antibody to detect the intracellular level of 7KCh in HL-1 cell (Fig. 4E). The 7KCh accumulated in cytosol of HL-1 cell upon 24 h treatment with 20 μM 7KCh.

To determine whether 7KCh causes ROS generation in HL-1 cells, we treated cells with 7KCh for 24 h, stained with H2DCFDA, and analyzed the results using flow cytometer. As expected, the DCF fluorescence increased in cells treated with 10 μM and 20 μM 7KCh (Fig. 4F). H2O2- treated HL-1 cells served as positive control. It was consistent with the results for cells stained with MitoSOX Red, a mitochondrion-specific probe for ROS. The 7KCh-treated HL-1 cells showed higher ROS production than positive control (i.e. HL-1 cells treated with 5 mM H2O2).

3.5. 7KCh induces cell death through activation of transcription factor 4 (ATF4) pathway

ATF4 is induced by various stressors, such as endoplasmic reticulum (ER), amino acid deprivation, and oxidative stress. 

Table 2

| No. | Metabolites                | Fold change Log2 (A/N) | p (corr) | p-value ANOVA | p-value (N vs. C) | VIP (N vs. C) | Trend of HF |
|-----|----------------------------|------------------------|----------|---------------|-------------------|--------------|-------------|
| 1   | LysoPC C16:0               | 0.86                   | 0.97     | 0.67          | 0.64              | 1.19E-08     | 1.67E-07    | 2.52        | Increased   |
| 2   | LysoPC C18:1               | 1.23                   | 1.44     | 1.44          | 0.70              | 2.62E-10     | 6.26E-08    | 1.02        | Increased   |
| 3   | LysoPC C16:0               | 3.43                   | 3.77     | 3.18          | 0.51              | 9.75E-09     | 2.39E-05    | 1.03        | Increased   |
| 4   | LysoPC C16:0               | 1.00                   | 1.20     | 0.86          | 0.67              | 7.66E-10     | 2.70E-08    | 1.59        | Increased   |
| 5   | 7-Dehydrocholesterol       | 3.28                   | 3.57     | 2.76          | 0.65              | 9.22E-15     | 3.71E-07    | 1.62        | Increased   |
| 6   | 7-Ketocholesterol           | 5.23                   | 5.58     | 4.60          | 0.67              | 7.48E-18     | 3.59E-07    | 6.87        | Increased   |
| 7   | Cer C22:0                  | 0.81                   | 0.74     | 1.01          | 0.57              | 4.69E-06     | 1.66E-06    | 1.57        | Increased   |
| 8   | Cer C24:0                  | 1.06                   | 0.97     | 1.46          | 0.65              | 2.50E-08     | 2.49E-08    | 2.69        | Increased   |
| 9   | Cer C24:1                  | 0.50                   | -0.49    | 1.61          | 0.53              | 1.27E-09     | 5.44E-05    | 1.33        | Increased   |
| 10  | LysoPE C20:4               | -2.31                  | -1.82    | -1.21         | -0.53             | 2.95E-11     | 4.75E-05    | 1.64        | Decreased   |
| 11  | PE C34:0                   | -0.94                  | -0.86    | -0.97         | -0.61             | 7.37E-12     | 3.47E-07    | 1.32        | Decreased   |
| 12  | PE C34:2                   | -1.57                  | -1.10    | -1.34         | -0.63             | 8.53E-17     | 7.58E-08    | 12.87       | Decreased   |
| 13  | PC C36:2                   | -1.43                  | -1.72    | -1.03         | -0.55             | 1.74E-11     | 1.61E-06    | 2.89        | Decreased   |
| 14  | PC C36:2                   | -1.09                  | -0.98    | -1.21         | -0.64             | 2.07E-18     | 1.76E-07    | 6.72        | Decreased   |
| 15  | PC C36:3                   | -2.25                  | -2.62    | -0.96         | -0.61             | 1.59E-24     | 2.03E-07    | 2.29        | Decreased   |
| 16  | PC C36:4                   | < -6.00                | < -6.00  | -2.85         | -0.62             | 1.69E-16     | 7.48E-07    | 1.14        | Decreased   |
| 17  | PE C34:1                   | -3.54                  | -5.06    | -3.23         | -0.51             | 5.45E-09     | 1.66E-05    | 1.24        | Decreased   |
| 18  | PE C34:2                   | -2.76                  | -2.88    | -2.20         | -0.77             | 2.07E-21     | 2.29E-09    | 3.98        | Decreased   |
| 19  | PE C36:1                   | -1.72                  | -1.46    | -1.44         | -0.52             | 1.72E-07     | 0.00027     | 1.42        | Decreased   |
| 20  | PE C36:2                   | -2.13                  | -2.92    | -1.30         | -0.60             | 3.49E-14     | 2.87E-05    | 2.01        | Decreased   |
| 21  | PE C36:3                   | -3.82                  | -4.19    | -2.64         | -0.80             | 4.90E-30     | 1.22E-10    | 3.78        | Decreased   |
| 22  | PE C36:4                   | -2.80                  | -3.32    | -1.33         | -0.60             | 1.46E-19     | 4.69E-07    | 3.37        | Decreased   |
| 23  | SM C24:2                   | -0.75                  | -0.69    | -0.59         | -0.65             | 7.55E-18     | 1.75E-08    | 2.25        | Decreased   |
| 24  | SM C24:1                   | -1.21                  | -0.98    | -1.56         | -0.64             | 3.37E-08     | 1.84E-07    | 2.00        | Decreased   |

PC, phosphatidylcholine; PE, phosphatidylethanolamine; Cer, ceramide; SM, sphingomyelin; VIP, The variable importance in the projection.

(Supplementary Tables S2 and S3). The levels of lysophospholipids (such as lysoPC and lysoPE), ceramides, and oxysterols (such as 7KCh) were higher in the HF erythrocytes than in the control erythrocytes. By contrast, the levels of phospholipids (such as PCs, PEs, and sphingomyelins) decreased in the HF erythrocytes compared with those of the normal cells (Table 2).

3.3. Erythrocyte and plasma 7KCh levels in patient with heart failure

The distribution of 7KCh between plasma and erythrocytes was reported for healthy volunteers [26]. Of the samples from the normal controls and patients in the current study, the 7KCh level was substantially lower than the levels of lanosterol, lathosterol, and 7-dehydrocholesterol (intermediates in cholesterol biosynthetic pathway) in plasma. By contrast, the 7KCh level was significantly higher than the levels of these metabolites in erythrocytes (Figs. 2 and 3).
oxidative stress accentuate each other, which have been shown to activate apoptotic signaling in vitro and in vivo models [28]. To test the possibility that 7KCh acts through activation of ATF4, we examined the expression level of ATF4 in 7KCh-treated HL-1 cells. The ATF4 level increased in a time-dependent manner (Fig. 5B, C). Moreover, the levels of CHOP and active caspase 3 increased significantly at 36 h after treatment (Fig. 5B, D, E), whereas that of LC3A/B increased slightly during the same period (Fig. 5B, G). These findings suggest that induction of ATF4 and CHOP contribute to death of cardiomyocytes.

4. Discussion

The lipidomes of erythrocytes from patients with HF were significantly different from those of normal controls. Moreover, 7KCh, lysoPCs, lysoPEs, and ceramides significantly increased in abundance in HF erythrocytes, whereas the levels of PCs, PEs, and SMs significantly decreased. Of these lipids, 7KCh was best at discriminating between patients and controls, and may thus serve as a biomarker for early identification of individuals at risk of HF. Additionally, 7KCh may be implicated in death of cardiomyocytes and HF pathogenesis.

Chronic inflammation is associated with HF progression. A number of proinflammatory cytokines, such as tumor necrosis factor α, interleukin (IL)-1, and IL-6, were implicated in this process [29]. In general, chronic inflammation leads to increased oxidative stress and damage and probably accounts for some of the observed changes in HF erythrocytes. Oxidative stress induces phospholipase activity, which leads to a decline in phospholipid levels and an increase in lysophospholipid levels. Moreover, 7KCh, an oxidation product of cholesterol, accumulates as a consequence of oxidative stress. Previous studies have revealed that oxidative damage products, such as oxidized LDL and oxysterols, are found in patients with cardiovascular disease [30,31]. 7KCh is considered an important metabolite for monitoring cardiovascular disease outcomes and mortality [32] as well as for predicting the incidence of cardiovascular disease events in general population [33]. Accumulation of 7KCh in HF erythrocytes suggests that 7KCh is a risk factor for HF, with a potential for clinical applications.

7KCh is mainly derived from oxidation of biomembranes and lipoproteins. Dietary intake of food oxysterols are not probably an important source of 7KCh, as dietary 7KCh can be rapidly metabolized to bile acid in liver and excreted [34–36]. Erythrocyte 7KCh may be derived from oxidation of membrane of erythrocytes. Moreover, elevation in erythrocyte 7KCh levels may be related to the changes in its catabolism in patients with HF. CYP7A1, the rate-limiting enzyme in bile acid biosynthesis, is the only hepatic enzyme known to be involved in 7KCh catabolism [37]. CYP7A1 deficiency causes premature atherosclerosis in humans [38,39]. The extrahepatic metabolism of 7KCh occurs by esterification to fatty acids through cytosolic sterol O-acyltransferase (SOAT) and subsequent selective efflux to HDL [37]. HDL...
transfers esterified 7KCh back to liver for further catabolism. Relatively low SOAT1 and SOAT2 expression in heart tissue leads to poor esterification and accumulation of free-form 7KCh. A decrease in reverse cholesterol transport (RCT) via HDL ensues [37]. The situation is aggravated by lower plasma HDL content in HF patients. It has been recently shown that erythrocytes play an important role in RCT. Animal study has shown that erythrocytes acquired tritiated cholesterol from subcutaneously injected foam [13]. Also, erythrocytes can exchange cholesterol with lipoproteins [40]. It is likely that erythrocytes may take up 7KCh from peripheral tissues and/or from other lipoproteins. Erythrocytes can then transport 7KCh directly to liver for catabolism, or transfer it to HDL via the exchange mechanism. The latter process may be less effective in light of the decrease in plasma HDL in HF patients. An increase in 7KCh abundance in erythrocytes may reflect not only increased oxidative stress but also an inefficient removal through RCT to HDL in HF patients.

7KCh causes cellular damage and induces oxidative stress in several types of cells, such as endothelial and cardiac cells [41,42]. Exposure to 7KCh promotes inflammation [43], ER stress [44,45], and lysosomal dysfunction [46]. 7KCh triggers autophagy through inhibition of Atg4B activity [44], and induces lysosomal dysfunction in vascular smooth muscle cells through an increase in oxidative stress [42]. Our findings show that 7KCh induces ROS formation and causes cardiomyocyte death, which is probably mediated by the ATF4/CHOP pathway. Previous studies have shown that ER stress leads to activation of PERK, phosphorylation of eukaryotic translation initiation factor 2, and subsequent translation of mRNAs, including ATF4. ATF4 itself can activate the transcription of CHOP, which is essential to ER stress-induced cell death [47–53].

Our findings may have interesting implications about the transport functions of erythrocytes. Erythrocytes may act as a transporter of substances other than oxygen. The free-form cholesterol of the erythrocyte plasma membrane can be bidirectionally exchanged with that of plasma lipoprotein and cellular plasma membrane. Erythrocytes were postulated to play a role in RCT, particularly in the low HDL state [13]. Our unpublished findings have shown an inverse correlation between erythrocyte 7KCh and plasma HDL in HF patients. Erythrocytes, in addition to HDL, may perform RCT function. Another intriguing possibility is that erythrocytes may transport 7KCh to cardiac tissue and inflict damage to cardiac cells. The proposed scheme is summarized in Fig. 6.

The current study has several limitations. One of the limitations pertains to the small cohort size of patients at different stages of HF. Although we demonstrated that 7KCh induces cardiomyocyte death, the
evidence supporting the causal relationship between an increase in erythrocyte 7KCh and HF development is far from complete. An animal model is needed in further study to validate our findings.

In conclusion, blood 7KCh is concentrated in erythrocytes in patients with HF. 7KCh-laden erythrocyte ghost induces cardiomyocyte death, suggesting its involvement in the pathogenesis of HF. Moreover, our findings support the importance of erythrocyte 7KCh as a risk factor for HF.
Fig. 6. A proposed model of the effect of erythrocyte 7KCh on cardiomycocytes. Erythrocytes may transport 7KCh to cardiac tissue, and inflicts damage to cardiac cells.

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Author Disclosure Statement

The authors declare no competing financial interests.

Appendix A. Supporting information

Supplementary data associated with this article can be found in the online version at http://dx.doi.org/10.1016/j.redox.2017.10.020.

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