Expression of connexin 43 protein in cardiomyocytes of heart failure mouse model

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Heart failure (HF) is the end stage of various cardiovascular diseases, with high morbidity and mortality, and is associated with a poor prognosis. One of the primary causes of HF is aortic valve disease, manifested by progressive aortic valve stenosis (AVS), resulting in increased left ventricular load, ventricular hypertrophy, ultimately ventricular dysfunction, and HF. Early assessment of the degree of cardiomyopathy and timely intervention is expected to improve patients’ cardiac function and delay or even avoid the occurrence of HF. The Wnt signaling pathway is mainly involved in regulating myocardial insufficiency after valve stenosis. Connexin 43 protein (Cx43) is an essential target of Wnt signaling pathway that forms gap junction (GJ) structures and is widely distributed in various organs and tissues, especially in the heart. The distribution and transformation of Cx43 among cardiac cells are crucial for the development of HF. To specifically label Cx43 in vivo, we established a new Cx43-BFP-GFP mouse model with two loxp sites on both sides of the tag BFP-polyA box, which can be removed by Cre recombination. This double-reporter line endowed us with a powerful genetic tool for determining the area, spatial distribution, and functional status of Cx43. It also indicated changes in electrical conduction between cells in a steady or diseased state.

KEYWORDS
heart failure, aortic valve stenosis, Cx43, Cx43-BFP-GFP, mouse model

Introduction

Heart failure (HF) is a chronic disease associated with poor outcomes and high mortality rates (1). It is the main cause of non-elective hospitalization in developed countries for patients over 65 years of age (2). HF is caused by myocardial damage for various reasons, of which heart valve disease is one of the major causes (3). Aortic valve stenosis (AVS) is an independent factor in left ventricular failure (4). Roughly 10% of patients with AVS suffer from heart failure with reduced ejection fraction (HFrEF) (5).
Materials and methods

Human myocardium samples

Myocardial tissues were collected from three patients who underwent heart transplantation at Zhongshan Hospital Affiliated with Fudan University. Patients with diabetes, hypertension, rheumatic valvular disease, and other cardiomyopathy were excluded from this study. Written informed consent was obtained from each patient before surgery with the approval of the Ethics Committee of Zhongshan Hospital, Affiliated with Fudan University (Ethical approval number: B2022-031R). Tissues were collected during surgery, and patient’s medical records were reviewed to assess clinical data and were performed in accordance with the Declaration of Helsinki.

Experimental animals

The animal study was reviewed and approved by the Ethics Committees of Zhongshan Hospital, Affiliated with Fudan University for the Care and Use of Animals for Research Purposes. These mouse lines were generated by Shanghai Biomodel Organism Science and Technology Development, Shanghai, China. Mice were maintained in a C57BL/6J genetic background and bred with a normal diet.

For generation of the Cx43-BFP-GFP mouse line, the CRISPR/Cas9 technology was used to insert loxp-tagBFP-WPRE-polyA-loxp-eGFP-WPRE-polyA cassette at the start codon site of Gja1 gene through homologous recombination. The brief process was as follows: Cas9 mRNA and gRNA were obtained by in vitro transcription. Homologous recombination vector was constructed by in-fusion cloning method, which contains a 3.5 kb 5′ homology arm, loxp-tagBFP-WPRE-polyA-loxp-eGFP-WPRE-polyA, and 3.2 kb 3′ homology arm. Cas9 mRNA, gRNA, and donor vector were microinjected into the fertilized eggs of C57BL/6J mice to obtain F0 generation mice. The positive F0 generation mice identified by PCR amplification and sequencing were mated with C57BL/6J mice to obtain positive F1 generation mice for the experiment.

Genomic polymerase chain reaction

Genotyping of mice was achieved by PCR amplification. Genomic DNA was extracted from the toes of mice. These toes were digested by lysis buffer with proteinase K at 55°C oven overnight, followed by centrifuging at 15,000 rpm for 8 min to obtain the supernatant, added with isopropanol to precipitate the DNA, and then washed in 70% ethanol twice. The PCR primers used to identify wild-type mice were 5′-AGTTGGGCTGGCTTCATTCC-3′ and 5′-TGCGGTGTTCTTCAATCCCAT-3′ (wild-type allele). For the Cx43-BFP-GFP/+ line, we used the genotyping primers 5′-GATTGGGCTGGTGATTCTGTTTTG-3′ and 5′-CGCCCCGTCTTGGATGT-3′ to detect.
Tissue whole-mount fluorescence microscopy

Isolated organs and tissues were washed several times in PBS to clear blood, and the excess tissues were removed under the bright field of the microscope. Then placed on transparent agar in an anatomical direction to obtain the bright-field whole-mount images and fluorescent images by the Zeiss stereoscope (AxioZoom V16). We applied the automatic z-stack image function by the Zeiss stereoscope (AxioZoom V16) to obtain the magnification of a specific area.

Tissue collection and Immunofluorescence staining and imaging

The isolated organs were fixed in 4% paraformaldehyde (PFA) for 1 h and washed 3 times in PBS, then placed them in 30% sucrose solution to dehydrate at 4°C overnight. The samples were embedded and frozen in OCT compound (Sakura, 4,583) for freezing and then stored at –80°C. For immunofluorescence staining, the frozen sections were cut into 10 µm thick and placed in a fume hood for 30–60 min. Subsequently, the slices were blocked with 5% normal donkey serum and 0.1% Triton X-100 PBS for 30 min at room temperature after 3 times of PBS washing. Then sections were incubated with primary antibodies overnight at 4°C. After that, sections were incubated with corresponding secondary antibodies for 40 min at room temperature in the dark after 3 times of PBS washing. After three washes, the stained sections were fixed with 50% glycerol and stored for further analysis. The immunostained images were obtained by Zeiss confocal microscope system (LSM710) and Olympus laser scanning confocal microscope (Fluoview 1200). The primary antibodies used for immunofluorescence staining were as follows: Cx43 (Sigma-Aldrich, C6219, 1:100); α-SMA (Sigma, F3777, 1:100); Cdh5 (R&D, AF1002, 1:100); PDGFRα (R&D, AF1062, 1:100); Troponin I (Abcam, ab56357, 1:100); CD45 (eBioscience, 17-0451-82, 1:100); DAPI (Vector Laboratories, 1:1,000). Used Alexa-conjugated secondary antibody (Invitrogen) at a concentration of 1:1,000.

Hematoxylin-esosin staining

Tissue slices were soaked in PBS for 10 min and then placed in hematoxylin for 5 min, and placed in hydrochloric acid ethanol for 1 min after rinsing with running water. Next, put the sections in ammonia water for 5 min after dyeing, then in 95% ethanol for 1 min, eosin dye for 8 s, and rinsed in deionized water. Subsequently, placed the stained slices in 95% ethanol for 1 min, 100% ethanol for 4 min, and xylene for 10 min. Mounted the slices with mounting medium.

Sirius red staining

Tissue slices were soaked in PBS for 15 min and then fixed in 4% PFA for 15 min, then washed 3 times with PBS. Next, the cryo sections were incubated overnight at room temperature in Bouins’ solution, then washed 3 times in PBS. It was then sequentially incubated in 0.1% Fast Green for 5 min and 0.1% Sirius red for 3 min. The slides were sequentially dehydrated in ethanol for 5 min and in xylene for 5 min after being washed 3 times in PBS. Finally, mounted the slices with mounting medium.

Transverse aortic coarctation

The mice were placed in a container containing isoflurane (2% isoflurane dissolved in 100% O₂, 0.8–1.2 L/min) for
anesthesia and then fixed on the operating table. Then the mice were connected to an animal ventilator at 120–140 times/min and 0.2 mL tidal volume after endotracheal intubation. Thoracotomy was performed on the second rib under a microscope after disinfecting the surgical site with 75% alcohol. Passed the 6.0 thread through the aortic arch between the innominate artery and the left carotid artery, placed a 26 G pad over the intended ligation site of the aortic arch, then wrapped the ligature around the aorta and ligate with a 26 G pad, and carefully withdraw the pad. Closed the chest after confirming that there was no hemothorax or pneumothorax. Removal of the tracheal tube when mice breathe spontaneously.

**Animal echocardiography**

Depilated the mice before the procedure. The mice were placed in a container containing isoflurane (2% isoflurane dissolved in 100% O\textsubscript{2}, 0.8–1.2 L/min) for anesthesia, and then fixed on the operation board of a cardiac ultrasound machine to expose the chest. Then the mice limbs were connected to sensing electrodes and added conductive liquid to record ECG signals. Data collection was performed by a professional echocardiographer. Echocardiography was conducted with the VisualSonics Vero 2100 system. The measurement parameters included ejection fraction (EF), fractional shortening (FS), left ventricular internal diameter at end-diastole (LVIDd), and left ventricular internal diameter at end-systole (LVIDs).

**Quantification and statistical analysis**

The obtained images were processed and analyzed using PhotoLine software. Statistical analyses were performed using Student's t-test with GraphPad Prism 8 software. All data were representative of at least three independent experiments and presented as mean ± SD. Student’s tests were applied for comparison between two groups. *p-value of < 0.05, **p-value of < 0.01, ***p-value of < 0.001, and ****p-value of < 0.0001 were considered as statistically significant difference.

**FIGURE 2**

Generation of Cx43-BFP-GFP/+ reporter line. (A) Schematic diagram showing the targeting strategy of Cx43-BFP-GFP allele by homologous recombination using CRISPR/Cas9; (B) polymerase chain reaction (PCR) of genomic DNA, P1 + P2 means primer1 + primer2; P3 + P4 means primer3 + primer4; Heterozygous amplified a 564 bp band.
Results

Myocardial characteristics of aortic valve disease

Immunohistochemical results showed that cardiomyocytes in patients with severe AVS were abnormally hypertrophic and disordered, with uneven cytoplasmic staining and decreased intercellular density compared with normal myocardial tissues (Figure 1A). In addition, severe myocardial collagen deposition in AVS patients. The fibrosis area in AVS patients was approximately 2.5 times higher than that of the control group (Figures 1B,C). Progressive myocardial fibrosis in aortic stenosis drives the progression of HF (21). Therefore, we stained myocardial tissues for Cx43 and the cardiomyocyte-specific marker protein Troponin I and found that Cx43 expression was significantly reduced and unevenly distributed in AVS myocardium compared with normal myocardium (Figure 1D).

FIGURE 3
Generation and Characterization of Cx43-BFP-GFP/+ reporter line. (A,B) Whole-mount bright field and epifluorescence images of visceral organs from 6-weeks Cx43-BFP-GFP/+ mice; (C) immunostaining for BFP and Cx43 on heart, lung, brain, stomach, kidney, liver, spleen, and pancreas sections. The high magnification images of organs in areas labeled with the square dotted frame, respectively; (D) quantification of the labeling efficiency by the percentage of BFP⁺ Cx43⁺ in BFP⁺ cells. Data are the mean ± SD; n = 5 mice per group; (E) quantification of the labeling specificity by the percentage of BFP⁺ Cx43⁺ in Cx43⁺ cells. Data are the mean ± SD; n = 5 images per group. Yellow scale bar = 1 mm, white scale bar = 100 µm.
The fluorescence intensity of both Cx43 and Troponin I decreased by more than 20% compared to the control ones (Figure 1E). Information of patients and donors is provided in Supplementary Table 1.

**Generation of Cx43-BFP-GFP reporter line**

In order to further explore the distribution and expression of Cx43 in valvular stenotic cardiomyopathy, we constructed a tool mouse Cx43-BFP-GFP to specifically label Cx43. To generate Cx43-BFP-GFP knock-in mouse line, we inserted a loxp-tagBFP-WPRE-polyA-loxp-eGFP-WPRE-polyA cassette into the 3′-UTR of the Gja1 gene through homologous recombination. In the targeting construct, the tagBFP and polyA stop cassette (tagBFP-polyA) is flanked by two loxp sites, and the enhanced GFP is followed by a second loxp as illustrated in Figure 2A.

To optimize the genotyping protocol for Cx43-BFP-GFP knock-in, we re-designed four PCR primers [P1 and P2 to amplify the wild-type allele (666 bp) and P3 and P4 to amplify the inserted site (564 bp)], which were located outside or inside of the loxp-tagBFP-WPRE-polyA-loxp-eGFP-WPRE-polyA cassette sequence (Figure 2B). Genotyping...
by PCR using the forward and reverse primers showed that the loxp-tagBFP-WPRE-polyA-loxp-eGFP-WPRE-polyA cassette was successfully inserted into the Cx43 locus. No defects were observed in the generated heterozygous Cx43-BFP-GFP reporter mice, compared with wild-type littermates.

Characterization of Cx43-BFP-GFP reporter line at 6 weeks

We examined the Cx43-BFP-GFP reporter expression by fluorescence imaging of a whole mount of organs and immunostaining of frozen organ sections from Cx43-BFP-GFP mice at 6 weeks. Although tag BFP reporter expression was not obvious in the whole-mount views (Figures 3A,B), we found that tag BFP was widely expressed in all organs. To further characterize the Cx43-BFP-GFP mouse line, BFP and Cx43 were widely expressed in tissues including the heart, liver, lung, stomach, pancreas, brain, kidney, brain, and spleen (Figure 3C). Microscopic quantification showed that 94 ± 0.71% and 94.2 ± 0.58% of BFP+ Cx43+ express BFP in the heart and brain, respectively (Figure 3D). It is shown that 92.8 ± 0.97% and 81 ± 2.12% of BFP+ Cx43+ express Cx43 in the heart and brain, respectively (Figure 3E). These data demonstrate that the Cx43-BFP-GFP mouse line has high labeling specificity and efficiency.

Characterization of Cx43-BFP-GFP reporter line at the embryonic stage

To demonstrate the expression of our tool in embryos, we proceeded to analyze the Cx43-BFP-GFP mice at the embryonic stage. At embryonic day (E) 16.5, we detected that BFP signals were robustly expressed in the dorsal neural tube, heart, kidney, lung, and brain but not obvious in other organs from the whole-mount views (Figures 4A,B). Similarly, we immunostained the genetic markers BFP and Cx43 on the tissue sections showing that plenty of BFP+ and Cx43+ cells were identified in the heart, brain, kidney, and lung at E16.5 (Figure 4C). Microscopic quantification showed that 96.4 ± 0.24% and 94.6 ± 0.51% of BFP+ Cx43+ express BFP in the heart and brain, respectively (Figure 4D). It is shown that 94.4 ± 0.51% and 95.2 ± 0.37% of BFP+ Cx43+ express Cx43 in the heart and brain, respectively (Figure 4E). These data indicate that the Cx43-BFP-GFP mice line exhibits high labeling efficiency in tissues at the early embryonic stage.

Expression of Connexin 43 in different cells with Cx43-BFP-GFP mice at 6 weeks

Cx43 is widely distributed in various cells of organs throughout the body, especially cardiomyocytes (22). To precisely demonstrate the cell type which expresses BFP reporter in the heart, co-staining assays of heart sections with BFP and either cardiomyocyte marker Troponin I or endothelial cell marker Cdh5 antibodies at 6-weeks Cx43-BFP-GFP mice were produced showing that BFP labeled most cardiomyocytes and endothelial cells. However, we did not find BFP expression in coronary vascular SMCs and immune cells, as illustrated in Figure 5A by immunostaining for the smooth muscle cell marker α-SMA and the immune cell marker CD45 on heart sections, consistent with previous reports (23).

In addition, to further assess the expression of Cx43 in cells of various tissues of Cx43-BFP-GFP mice at P6W, we performed...
co-immunostaining for BFP and other cellular markers, including α-SMA, Cdh5, CD45, and the fibroblast marker PDGFRα on other tissue sections showing in Figures 5B–D. The results showed that Cx43 was widely expressed in endothelial cells and fibroblasts in the liver but less expressed in immune cells and smooth muscle cells (Figure 5B). BFP targeted most endothelial cells, immune cells, and fibroblasts on kidney sections. Moreover, our observations confirmed that fewer SMCs were marked by BFP in the adult kidneys, as illustrated in Figure 5C. Similarly, we
stained the spleen sections with antibodies against BFP, Cdh5, PDGFRα, or CD45 demonstrating that BFP was detected in many endothelial cells, fibroblasts, and immune cells of the spleen cord. Interestingly, no BFP-labeled SMCs were found in splenic sinuses shown in Figure 5D. We also stained the Cx43-BFP-GFP mouse brain, lung, stomach, and thymus with BFP, Cdh5, PDGFRα, and CD45 antibodies to observe the co-localization of BFP and Cx43 in these tissue cells (supplementary Figure 1), suggesting that BFP reporter expression was confined to specific cells in different organs.

Connexin 43 protein expression in the hearts of heart failure mice

The murine transverse aortic constriction (TAC) is a commonly used experimental model to mimic AVS and HF (24). So, we evaluated the cardiac function of the mice by echocardiography 4 weeks after TAC in Cx43-BFP-GFP mice. Figure 6A displays representative M-mode echocardiograms showing evidence of HF in TAC group. Similarly, we evaluated the EF, FS, and LVID in mice, as shown in Figures 6B–E. The results showed that EF and FS were significantly reduced, LVIDd, and LVIDs increased significantly in TAC mice. TAC mice had markedly enlarged hearts and massive collagen fibril deposition in the myocardium (Figures 6F–H). The fibrosis area in the HF mice was approximately 100 times higher than that of the control ones (Figure 6I). Similarly, we stained myocardial tissues with the fluorescent marker BFP and found that the expression of Cx43 was widely and uniformly distributed in normal myocardial tissue; however, the fluorescence expression of Cx43 is decreased and distributed unevenly in the myocardial tissues of HF mice (Figures 6J,K). The co-staining of BFP and other cell markers could further clarify the main cellular transformation of Cx43 in the decompensation of intercellular signaling in cardiomyopathy caused by valvular disease, which is beneficial to the subsequent determination of the mechanism of HF caused by other factors such as valvular disease.

Conclusion

In our study, we found that AVS disease, which is an independent factor for HF, had marked fibrosis and uneven distribution of GJs in the myocardium. To better explore this pathological process, we established the Cx43-BFP-GFP mouse model. In this reporter line, Cx43+ cells can be labeled with BFP, with high labeling efficiency in important tissues and organs, especially cardiac cells. The efficacy of this reporter line has been validated in the TAC HF model with decreased expression and uneven distribution of Cx43 in cardiomyocytes of mice with HF. This dual reporter gene line provides a unique genetic tool for understanding the pathological process of myocardial remodeling and provides a new treatment direction for delaying HF.

Discussion

Various factors that cause myocardial injury may lead to cardiac systolic dysfunctions, resulting in HF. Among them, the pathological injury of AVS is one of the leading factors (25–27). Previous reports have found that moderate aortic stenosis was related to poor prognosis in patients with HFref (28). The myocardium of valvular stenosis has obvious fibrosis. This process involves the participation of a variety of cells, among which the coupling and signal transduction mechanisms between cardiomyocytes, endothelial cells, fibroblasts, and other cells are not fully understood yet. Previous studies pointed out that the Wnt signaling pathway had an important role in the pathogenesis of HF. The Wnt signaling pathway is mainly transduced in cells through two different pathways, the canonical Wnt/β-catenin pathway and the non-canonical Wnt/Ca2+ pathway (29). Persistent activation of endothelial Wnt/β-catenin signaling induces HF (30). Besides, cardiac fibroblasts are the main cell types in cardiac fibrosis (31); interestingly, Wnt/β-catenin signaling also had vital functions in regulating fibroblast activation and ECM gene expressions (32). Thus, studying the Wnt/β-catenin pathway is the key to unveiling the pathogenesis in HF.

Cx43 is considered to be a target of Wnt/β-catenin transcription, which interacts with the β-catenin (33, 34). One of the critical mechanisms of cardiac remodeling is the change in intracellular electrical coupling caused by aberrant expression of Cx43 protein (35, 36). Moreover, its ubiquitous expression at the poles of ventricular and atrial myocytes and in certain regions of the conduction system (22, 37) has significantly correlated with cardiac arrhythmia. Thus, the observation of Cx43 in vivo is beneficial to a better understanding of the pathogenetic mechanism and is essential for drug screening and other clinical practices in HF.

Several Cx43 Cre-loxp and Cx43 Cre-ER (T)/+ (38, 39) mouse tools were generated previously by transgene and applied to study diseases such as arrhythmia (40). Cx43kiECFP (41) and Cx43kiLacZ (42) mouse tools have been used in specific subsets of cells. The advantage of the Cx43-BFP-GFP line developed in this study is that Cx43+ cells can be labeled with BFP in a natural state, and the GFP could be expressed by crossing the specific Cre mice if available. Therefore, the dual reporter system is expected to explore further the interaction and transformation of cardiac fibroblasts, endothelial cells, cardiomyocytes, and other cells in HF or other heart-related diseases.

Our research has introduced a double-reporter Cx43-BFP-GFP line. This mouse line cannot only display the existence of Cx43 but also be used as a conditional knockout line with the second reporter of eGFP. Meanwhile, to date, the experiments from Cx43-BFP-GFP: Cre mice remain lacking. Crossing the
reporter line with Cre mice to verify the advantage of the dual reporter system in HF still needs further research.

Data availability statement

The original contributions presented in this study are included in the article/Supplementary material, further inquiries can be directed to the corresponding author/s.

Ethics statement

The studies involving human participants were reviewed and approved by the Ethics Committee of Zhongshan Hospital, Affiliated With Fudan University. The patients/participants provided their written informed consent to participate in this study. The animal study was reviewed and approved by the Ethics Committees of Zhongshan Hospital, Affiliated With Fudan University.

Author contributions

SL and YL performed material preparation, data collection, and analyses. QZ and YZ edited and commented on the manuscript. KL and TL bred the mice, performed the experiments, and analyzed the data. JG supervised the study and analyzed the data. YY contributed to the study conception and design. All authors reviewed the manuscript and agreed to be accountable for the content of the work.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fcvm.2022.1028558/full#supplementary-material

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