Effects of acute ischemia and hypoxia in young and adult calsequestrin (CSQ2) knock-out and wild-type mice

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Abstract

Calsequestrin (CSQ2) is the main Ca2+-binding protein in the sarcoplasmic reticulum of the mammalian heart. In order to understand the function of calsequestrin better, we compared two age groups (young: 4–5 months of age versus adult: 18 months of age) of CSQ2 knock-out mice (CSQ2(−/−)) and littermate wild-type mice (CSQ2(+/+)). Using echocardiography, in adult mice, the basal left ventricular ejection fraction and the spontaneous beating rate were lower in CSQ2(−/−) compared to CSQ2(+/+). The increase in ejection fraction by β-adrenergic stimulation (intraperitoneal injection of isoproterenol) was lower in adult CSQ2(−/−) versus adult CSQ2(+/+). After hypoxia in vitro (isolated atrial preparations) by gassing the organ bath buffer with 95% N2, force of contraction in electrically driven left atria increased to lower values in young CSQ2(−/−) than in young CSQ2(+/+). In addition, after global ischemia and reperfusion (buffer-perfused hearts according to Langendorff; 20-min ischemia and 15-min reperfusion), the rate of tension development was higher in young CSQ2(−/−) compared to young CSQ2(+/+). Finally, we evaluated signs of inflammation (immune cells, autoantibodies, and fibrosis). However, whereas no immunological alterations were found between all investigated groups, pronounced fibrosis was found in the ventricles of adult CSQ2(−/−) compared to all other groups. We suggest that in young mice, CSQ2 is important for cardiac performance especially in isolated cardiac preparations under conditions of impaired oxygen supply, but with differences between atrium and ventricle. Lack of CSQ2 leads age dependently to fibrosis and depressed cardiac performance in echocardiographic studies.

Keywords Calsequestrin · Hypertrophy · Aging · Ischemia · Hypoxia
**Introduction**

Ca\(^{2+}\) is crucial for excitation–contraction coupling in the mammalian heart notably the human heart. In the sarcoplasmic reticulum (SR) of the cardiomyocytes, calsequestrin is important for the storage of Ca\(^{2+}\) and the release of Ca\(^{2+}\) from the SR (more details are shown in Fig. 1; for review: [1, 2]). Calsequestrin 2 (CSQ2) is the cardiac isoform of calsequestrin coded by a different gene than the isoform CSQ1 that is found in smooth muscles but also in skeletal muscles [1, 3–5]. There are undoubtedly at least a dozen Ca\(^{2+}\)-binding proteins in the heart notably calreticulin, which might compensate for CSQ2 in its function as a Ca\(^{2+}\)-binding protein. However, CSQ2 that binds Ca\(^{2+}\) with high capacity is quantitatively the main Ca\(^{2+}\)-binding protein in the heart [6]. Moreover, CSQ2 is hardly expressed in non-cardiomyocytes (e.g., in the esophagus or skeletal muscle), which one might interpret as indicative of a special cardiac role of CSQ2. In vitro data with recombinant proteins in artificial membranes have convincingly shown that CSQ2 can reduce the opening of the SR Ca\(^{2+}\) release channel (cardiac ryanodine receptor = RyR2) for Ca\(^{2+}\) [7, 8]. That should reduce cardiac force generation via the mechanism delineated in Fig. 1. In addition, CSQ2 might not only be relevant for physiological force generation in the heart but CSQ2 might also be important for cardiac arrhythmias: altered levels or inappropriate timing of Ca\(^{2+}\) release from the SR during diastole could induce the sarcolemmal Na\(^+\)/Ca\(^{2+}\) exchanger to extrude Ca\(^{2+}\) from the cytosol of the cardiomyocyte into the interstitium. Being electrogenic, an activated Na\(^+\)/Ca\(^{2+}\) exchanger should lead to cardiac depolarization. Thus, cardiac depolarization might occur in a delayed fashion and this delayed after depolarization can bring about deadly arrhythmias.

In line with these theoretical mechanisms, several mutations in the CSQ2 gene have become known in some patients with catecholaminergic polymorphic ventricular tachycardia (CPVT) and might, at worst, lead to the death of the patients (even children) due to ventricular fibrillation [2, 9–11]. In susceptible patients, CPVT may occur for the first time after stimulation of β-adrenoceptors. This β-adrenoceptor stimulation may result from physical exercise or emotional stress [2]. An overview of incidences, manifestations, and treatment strategies for CPVT can be found elsewhere [12, 13].

Because of the putative important role of CSQ2 in cardiac force generation and in the heart beat, we ourselves and others developed and characterized homozygous CSQ2 knock-out mice (CSQ2(−/−)), using different genetic targeting strategies [14, 15]. The data of both groups are in broad agreement [14, 15]. For example, Glukhov and co-workers found interstitial fibrosis in the sinoatrial node of CSQ2(−/−) and noted altered cardiac pacemaker activity in the sinus node. They concluded that these functional and structural alterations could contribute to sinoatrial node dysfunction as well as to atrial fibrillation in patients with loss of function mutations of CSQ2(−/−) [16]. As Ca\(^{2+}\) is crucial for cardiac contractility, manipulation of cardiac CSQ2 levels might be thought to lead to altered responses to stressors that alter force of contraction by changing the Ca\(^{2+}\) homeostasis in the heart. Such stressors are cardiac hypoxia or cardiac ischemia, which we therefore studied here following established procedures in our laboratory.

Hence, we thought it worthwhile to study how fast cardiac preparations from CSQ2(−/−) mice respond with reduced contractility and enhanced arrhythmias to acute ischemia and acute hypoxia but also whether reduced force of contraction and/or arrhythmogenesis are fully reversible. To put it otherwise, we were interested how CSQ might contribute to reperfusion injury and we
hypothesized that under these conditions, altered cardiac function in CSQ2(−/−) might become apparent, which might be relevant for myocardial infarction, an important clinical disease. To test these hypotheses, we used isolated cardiac preparations. Moreover, it also is known that immunological phenomena can contribute to cardiac dysfunction and even cardiac failure. This can occur in acute cardiac failure (e.g., in sepsis) or chronic heart failure. Chronic immunological activation of the heart can worsen congestive heart failure. From a mechanistically point of view, altered Ca²⁺ homeostasis can change the activity of Ca²⁺-dependent kinases and these kinases can alter the activity of transcription factors that enhance the transcription of proinflammatory genes leading to cardiac inflammation. Therefore, we hypothesized that the loss of CSQ2 might lead to immunological alterations in the heart of young and/or adult mice. Moreover, aging per se is accompanied by inflammation in many tissues, also in the heart [17]. Cardiac aging leads to functional impairment, to which fibrosis, death of cardiac cells, and subsequent exposure of antigens on cardiac cells to the immune system contribute [17–19]. For instance, in mice, CD4⁺ T cells can mediate cardiac inflammation and functional impairment upon aging [20]. In addition, cardiac aging might involve cardiac-resident macrophages [21, 22]. These hypotheses were addressed by several in vitro techniques, like flow cytometry, immunofluorescence, and assessment of fibrosis by histology. Finally, we wanted to test the hypothesis that ablation of CSQ2 might alter echocardiographic functions like left ventricular ejection fraction or left ventricular myocardial strain parameters in young as well as in adult mice.

### Materials and methods

#### Mouse model

Calsequestrin (CSQ2)-deficient mice (CSQ2(−/−)), with ablation of the gene for calsequestrin 2 in the heart have been described before [14]. CSQ2(−/−) survive birth, are fertile, and show altered contractile parameters in isolated atrium or ventricle and react on stimulation with β-adrenoceptor agonists in vivo with cardiac arrhythmias [14]. Moreover, in initial data, we noted an impaired basal (that means in absence of β-adrenergic stimulation) contractility of older CSQ2(−/−) mice compared to age-matched controls [14]. These mice were used in the present study and compared with wild-type littermate mice CSQ2(+/+).

We have not selected for gender. In other words, male and female mice were randomly selected. That is, in all subsequent studies, about the same number of male and female mice was included in the CSQ2(+/+) group as well as in the CSQ2(−/−) group. Table 1 gives additional information on the mice used in this study. Total number of mice were as follows: CSQ2(+/+) $n = 38$ young and $n = 29$ adult and CSQ2(−/−), $n = 43$ young and $n = 28$ adult. The numbers of mice used in the specific experiments are given in the corresponding results section. The investigation conforms to the Guide for the Care and Use of Laboratory Animals published by the National Research Council (2011) [23]. Animals were handled and maintained according to approved protocols of the animal welfare committee of the University of Halle-Wittenberg, Halle, Germany (approval reference number 42502-2-1506 MLU). The manuscript was prepared as per the ARRIVE guidelines [24].

#### Table 1 Age and gravimetric measurements of the mice studied

| Genotype | CSQ2(+/-) | CSQ2(−/−) |
|----------|----------|----------|
|          | Young    | Adult    | Young    | Adult    |
| Age (d)  | 139.67 ± 30.26 | 538.75 ± 35.64$^\text{g}$ | 118.54 ± 12.35 | 458.18 ± 35.11$^\text{g}$ |
| Numbers (n) | 9       | 12       | 13       | 11       |
| Body weight (g) | 22.56 ± 1.14 | 25.87 ± 1.61 | 21.32 ± 1.37 | 26.05 ± 0.79$^\text{g}$ |
| Heart weight (mg) | 157.30 ± 12.19 | 218.69 ± 13.06$^\text{g}$ | 174.65 ± 13.25 | 240.46 ± 15.58$^\text{g}$ |
| Rel. weight heart (mg/g) | 6.99 ± 0.44 | 8.75 ± 0.61$^\text{g}$ | 8.27 ± 0.50 | 8.90 ± 0.50 |
| Rel. weight lung (mg/g) | 6.98 ± 0.48 | 7.51 ± 0.93 | 8.81 ± 0.72 | 7.31 ± 0.29 |
| Rel. weight liver (mg/g) | 53.10 ± 1.52 | 51.38 ± 4.02 | 54.85 ± 2.64 | 53.96 ± 1.54 |
| Rel. weight spleen (mg/g) | 3.61 ± 0.38 | 4.39 ± 0.56 | 3.64 ± 0.28 | 3.93 ± 0.22 |
| Rel. weight kidney (mg/g) | 15.05 ± 0.59 | 15.43 ± 0.67 | 15.21 ± 0.42 | 14.35 ± 0.56 |

Genotype indicates CSQ2(+/-) and CSQ2(−/−) mice. Numbers of animals are indicated in the second row

$^\text{g}p < 0.05$ vs. young age group. Relative (Rel.) weight means weight of organ divided by body weight
Histology

Tissue samples of young and adult mouse hearts were fixed in buffered 4% formaldehyde and routinely embedded in paraffin. Four-µm-thick paraffin tissue sections were deparaffinized with xylene and graded in ethanol. For pathohistological analysis, tissue sections were routinely stained with hematoxylin/eosin, elastica van Gieson staining, and Masson–Goldner-three-color staining. For visualizing and image processing, microscopic images were captured using an AxioCam digital microscope camera and the AxioVision image processing software (Carl Zeiss Microscopy, Oberkochen, Germany). The images were acquired at 96 DPI and submitted with the final revision of the manuscript at 300 DPI. Images shown are representative of at least three independent experiments, which gave similar results.

Immunofluorescence to detect cardiac auto-reactivity

Heart-specific autoantibodies were detected by incubating histological heart slices prepared from immunoglobulin-deficient animals (AID/µS −/−) with the plasma of C57BL/6j (+/+ ) and C57BL/6j (−/−) animals. Next, plasma reactivity against cardiac antigens was assessed using anti-mouse IgM-Alexa 555 and anti-mouse IgG-Alexa 488 antibodies (both from Thermo Fisher Scientific, Waltham, MA, USA). As control, some heart slides were incubated with secondary antibodies only. In the absence of plasma, immunofluorescence histology analysis was performed with a Zeiss Axioskop 2 plus (Carl Zeiss Microscopy, Oberkochen, Germany) and the quantification was done with the open source software ImageJ (National Institute of Health, USA).

Contraction experiments

Contraction experiments in isolated atrial preparations from young mice were performed as reported [23]. In brief, electrically driven (1 Hz) left atrial preparations were mounted in an organ bath. The bathing solution of the organ baths contained (in mM) NaCl, 119.8; KCl, 5.4; CaCl2 1.8; MgCl2, 1.05; NaH2PO4, 0.42; NaHCO3, 22.6; Na2EDTA, 0.05; ascorbic acid, 0.28; and glucose, 5.05. Contractile parameters were continuously measured at constant muscle length. Spontaneously beating right atrial preparations from mice were used to study any chronotropic effects. After an equilibration time of 30 min, buffer was bubbled with a mixture of 95% N2 and 5% CO2, in order to induce hypoxia. Under these conditions, the atrium was maintained at 37 °C and pHi at 7.4. The organ bath contained double glass walls allowing pre-warmed water to circulate and this allowed us to maintain 37 °C in the organ bath before, during, and after hypoxia. This protocol was successfully used and published previously by our group [25–27].

Contraction experiments in isolated whole-perfused hearts from young mice were also performed using established protocols in our laboratory [26, 27] with modifications of the original description [28]. In brief, spontaneously beating hearts were retrogradely perfused under constant flow (2 ml/min) and the force of contraction was measured mechanically at the apex of the heart by means of a force transducer. This signal was digitized as described above for atrial preparation. After 20 min of equilibration, hypoxia was induced for 20 min by stop of the perfusion pump followed by reperfusion by starting the perfusion pump again for 15 min. Contractile parameters were continuously electronically stored. Before stop of perfusion (basal parameters), at the end of stop of perfusion, and after reperfusion (hypoxia parameters) force parameters were measured and subjected to statistical analysis. The hearts remained at room temperature during this global ischemia and were not warmed. Thereafter, the pump was started again to simulate cardiac reperfusion.

Flow cytometry

A cell suspension of whole-heart samples was obtained after collagenase type II digestion (1.000 IU/mL. 37 °C. 30 min) and then ground in Hanks’ balanced salt solution containing 1% (wt/vol) BSA (BSS/BSA) using a 40-µm cell strainer, according to previous descriptions [29]. Lymph node samples were ground and prepared in BSS/BSA [5% (vol/vol) FCS]. Cells were stained using the following fluorescently labeled antibodies (in different combinations): anti-CD4 (clone RM4-5), anti-CD8 (clone 53–6.7), anti-CD44 (clone IM7), anti-CD45 (clone 30-F11), anti-CD45/B220 (clone RA3-6B2), and anti-TCRβ (clone H57-597). All antibodies were purchased from BioLegend (BioLegend GmbH, Koblenz, Germany). Measurements were made using an LSR-Fortessa cell analyzer (Becton Dickinson GmbH, Heidelberg, Germany) and the data were analyzed using FlowJo.
Echocardiography

Transthoracic echocardiographic measurements were performed in spontaneously breathing young and adult mice anaesthetized with 1.5% isoflurane using a Vevo 2100 system equipped with a MS 550D transducer (Visual Sonics, Toronto, Canada). Analysis of the myocardial wall motion was performed using the VisualSonics VevoStrain™ Software (Visual Sonics, Toronto, Canada) following the instructions of the manufacturer. After induction of anesthesia the mice were positioned on a 37-°C heating pad. Two-dimensional images and M-mode tracings from the parasternal long-axis view were recorded. Cardiac dimensions were measured and the ejection fraction of the hearts was calculated. Isoproterenol was injected, where indicated, at 1 mM in 100 µl, as reported before [30].

Data analysis

Data shown are means ± SEM. Statistical significance was estimated using T tests or for multiple comparisons by analysis of variance followed by Bonferroni's two-sided t test. A p-value < 0.05 was considered significant.

Drugs and materials

Isoproterenol was obtained from Sigma-Aldrich (Taufkirchen, Germany). All other chemicals were of analytical grade. Deionized water was used throughout the experiments. Stock solutions were freshly prepared daily.

Results

Gravimetric results

In young mice, body weights were similar in CSQ2(−/−) or littermate wild-type mice CSQ2(+/+) (Table 1). However, the relative heart weights (heart weight divided by body weight) were lower in young CSQ2(−/−) than in littermate young CSQ2(+/+) mice (Table 1). Aging increased heart weight in CSQ2(−/−) and CSQ2(+/+) (Table 1). Absolute body weights were larger in adult KO mice than in adult CSQ2(+/+) mice (Table 1). In contrast, relative heart weights were increased in adult CSQ2(+/+) but not adult CSQ2(−/−) (Table 1). As a control, we determined also relative spleen, kidneys, liver, and lung weights, which were not significantly different between genotypes or ages (Table 1).

Basal echocardiography

Next, echocardiographic measurements in narcotized animals were performed in order to study cardiac function in vivo. It is noteworthy that the ejection fractions under basal conditions were not different between genotypes in young mice, whereas in adult CSQ2(−/−) mice, the basal ejection fraction as well as the beating rates were lower than in adult CSQ2(+/+) mice, arguing for an age-dependent reduction in left cardiac ejection fraction due to CSQ2 ablation (Table 2) which confirms an earlier study [12] on a different group of animals from our lab. Here, we extend those data by a rather complete assessment of noninvasive echocardiographic data, notably flow measurements of vessels, functional assessment of the mitral valve, tissue Doppler of the left ventricle, and septal and left ventricular systolic and diastolic diameters.

Here, it is noteworthy that under basal conditions the velocity time integral (VTI) through the aorta descendens was lower in adult CSQ2(−/−) compared to young CSQ2(−/−), suggesting an involvement of aging. In adult CSQ2(−/−) the VTI through the pulmonary artery was lower than in young CSQ2(−/−) but notably also than in adult CSQ2(+/+) suggesting both, an involvement of aging and gene ablation. Moreover, left ventricular end-systolic diameter was larger in adult CSQ2(−/−) than in adult CSQ2(+/+) indicative of a role of CSQ2 knock-out. Furthermore, the wall motion was assessed by regional strain parameters using under these basal echocardiographic conditions (Table 3). Of note, effects of genotype and gene ablation behaved in some parameters differently. For instance, radial lateral wall strain increased with aging in CSQ2(−/−) as well as in CSQ2(+/+) (Table 3). Mid longitudinal posterior strain decreased with aging in CSQ2(+/+) but increased with aging in CSQ2(−/−). Mid longitudinal posterior strain rates were higher in adult CSQ2(−/−) than in young CSQ2(−/−). However, mid longitudinal posterior strains and mid longitudinal strain rates were lower in CSQ2(−/−) than in CSQ2(+/+) (Table 3).

Echocardiography under isoproterenol

Next we assessed the response to an intraperitoneal injection of isoproterenol, which stimulates β-adrenoceptors, in a dose that we have used before in echocardiographic studies of the heart of other transgenic mouse lines [30, 31]. Isoproterenol led to higher peak velocity in the aorta descendens of adult versus young CSQ2(−/−). Interestingly, while injected isoproterenol increased ejection fraction, in adult and young mice both transgenic and CSQ2(+/+), and the percentile increase in injection fraction was smaller in adult (but not young) CSQ2(−/−) compared to adult CSQ2(+/+) (Table 2). After injection
of isoproterenol, the left ventricular end-systolic diameters were diminished in all genotypes; but less so in adult CSQ2(−−) compared to adult CSQ2(+/+), in line with the findings on ejection fraction and suggesting a gene knock-specific effect of aging (Table 2). In addition, supposedly as a consequence of impaired stimulation of left ventricular function by injection of isoproterenol, likewise the pulmonary artery velocity time integral exhibited lower values in CSQ2(−−) compared to CSQ2(++) after injection of isoproterenol (Table 2).
Fibrosis

We tested the hypothesis that upon aging, fibrosis might be detectable histologically in adult CSQ2(−/−). Employing Masson–Goldner three-color staining, we found a heavy fibrosis in the ventricles (n = 3) from adult CSQ2(−/−) but no signs of fibrosis in the ventricles (n = 3) from adult CSQ2(+/+) (Fig. 2). We display here (Fig. 2) only one of three adult CSQ2(+/+) but failed to detect fibrosis in the other samples from adult CSQ2(+/+). We failed to detect fibrosis histologically in samples from young mice of CSQ2(−/−) and CSQ2(+/+) (n = 3 each, data not shown).

Immune responses

In addition, we sought to assess whether immune-mediated responses could contribute to the cardiac decline observed in CSQ2(−/−). Therefore, we performed flow cytometry analyses to assess the distribution of all major lymphocyte subsets in collagenase-digested hearts and mediastinal lymph nodes (heart -draining) obtained from CSQ2(+/+) and CSQ2(−/−) mice. As shown in Fig. 3 and Table 4, we did not observe any differences in the composition of cardiac T- and B-lymphocytes subsets across different genotypes, indicating that in situ inflammation might not contribute to CSQ2-mediated cardiac decline. Nevertheless, CSQ2(−/−) presented an enlargement of the heart-draining lymph nodes, especially associated with an expansion of the B-cell compartment (Fig. 3, Table 4). These alterations in the B-cell compartment led us considering a possible role for autoantibody-mediated cardiac damage in CSQ2(−/−). To assess the putative role of autoantibodies in this model, we incubated plasma obtained from CSQ2(+/+) and CSQ2-deficient mice with heart slices obtained from immunoglobulin-deficient mice. The putative presence of heart-reactive antibodies in the sera of CSQ2(+/+) versus CSQ2(−/−) was then detected using an anti-mouse IgG and IgM conjugated with fluorescent dyes, followed by inspection in the fluorescence microscope. As shown in Fig. 3 and summarized in Table 4, the plasma obtained from CSQ2(+/+) and CSQ2(−/−) showed the same cardiac autoantibody pattern. These data suggest that autoantibodies are not likely to play a role in the CSQ2-mediated cardiac functional decline.

Hypoxia

As a further stressor for cardiac function in young CSQ2(−/−), we induced hypoxia in paced isolated left and spontaneously beating right atrial preparations by replacing oxygen with nitrogen in the organ bath. We have used this method successfully in other transgenic mice to mimic hypoxia in vivo [26, 27]. This hypoxia in the organ bath was able to reduce force of contraction in the left atrial preparations within 30 min nearly completely, consistent with our earlier publications [31]. Thereafter, normoxia was re-instated but developed force of contraction and its first derivative did not return to the values observed before hypoxia, in both CSQ2(+/+) and CSQ2(−/−) (data not shown). Of note, at 10 min of reperfusion, the decrease in developed force of contraction (expressed in percentage of pre-hypoxic values) and its first derivative with regard to time were more pronounced (that is after previous hysteresis).
hypoxia) in CSQ2(−/−) than in age-matched CSQ2(+/+): 28.1 ± 5.92% of pre-hypoxic levels versus 10.1 ± 1.59% of pre-hypoxic values or 34.8 ± 10.2 mN/sec and 17.2 ± 2.89 mN/sec, respectively (p < 0.05; n = 8–9), suggesting that CSQ2 is important for the mouse atrium to sustain better short-term hypoxia. Time to peak tension was unchanged in hypoxia in both genotypes. However, time of relaxation was prolonged in both, CSQ2(+/+) and CSQ2(−/−), under hypoxia and under re-oxygenation, it was normalized only in CSQ2(+/+) but in CSQ2(−/−), and time of relaxation fell below the pre-hypoxic values: 33.9 ± 1.75 ms versus 30.0 ± 1.68 ms (p < 0.05; n = 8–9).

Heart rates in spontaneously beating right atrial preparations were increased at the beginning of hypoxia and all preparations of CSQ2(−/−) and CSQ2(+/+) went into arrhythmias at the end of the maximum time of hypoxia, but returned to sinus rhythm and pre-hypoxia rates. Heart rates in CSQ2(+/+) and CSQ2(−/−) remained below pre-hypoxic heart rate (data not shown). Thus, no difference with respect to heart rate were notable in isolated right atria between the genotypes studied.

Ischemia

Next, we studied cardiac ischemia in an in vitro model as a further stressor. Here, in order to assess left ventricular function, protocols were used that have been published before [26, 27]. We studied global ischemia in the isolated spontaneously beating retrogradely perfused mouse heart (Langendorff heart, [28]): after stabilization of the contractile function in the isolated perfused hearts for 30 min, global ischemia was induced by stopping perfusion of the heart and this rapidly led to a cessation of contraction. The time to the stop of any heart beat amounted to 65 ± 7.3 s and 145 ± 15 s in young CSQ2(−/−) and CSQ2(+/+), respectively (n = 5, p < 0.05), suggesting a protective role of CSQ2. Reperfusion led to the reappearance of spontaneous beating of the hearts of both genotypes. After 15 min of reperfusion, developed tension was lower in CSQ2(−/−) than in CSQ2(+/+) (15.6 ± 3.23 mN and 25.8 ± 2.3) but also its first derivate versus time was lower in CSQ2(−/−) than in CSQ2(+/+) (dF/dt: 458 ± 81.7 mN/s and 776 ± 62.1 mN/s; n = 5 each, p < 0.05), suggesting that
CSQ2 is possibly detrimental for maintenance of left ventricular force after global ischemia.

From these data one could argue that CSQ2 is relevant in cardiac hypoxia and aging in a mouse model. Data in patients are awaited with interest.

**Discussion**

Novel findings in this report include a systematic comparison of echocardiographic parameters of CSQ2(−/−) with age-matched controls. This was done with mice at younger and older age. Moreover, we present new data on ventricular fibrosis as a possible mechanism that might explain some of the echo findings in older CSQ2(−/−). Another new finding in this report is that CSQ2(−/−) are less able to withstand the stress of hypoxia and ischemia under the present experimental conditions.

We had deleted constitutionally the coding region of exon [14, 32] and, using classical methods, we finally achieved one heterozygous founder, the offspring of which was used also in the present studies. In the initial description of this mouse, we reported the absence of CSQ2 on protein level in Western blots of atria and ventricles from CSQ2(−/−), implying successful deletion of CSQ2 [14]. The observation that homozygous CSQ2(−/−) were viable and fertile, most likely means that CSQ2 is not absolutely required for Ca2+ storage and handling in the mammalian heart. This suggests that the cardiac function of CSQ2 must be more subtle than originally assumed from biochemical studies: the function of CSQ2 as a Ca2+ buffer is apparently overtaken, at least in part, by other proteins, which act as Ca2+ stores in the SR. However, it needs to be elucidated how other proteins can compensate, exempli gratia, for the modulatory role of CSQ2 on RYR2 function. Structurally, in accordance with patient data and animal data [15, 33], the hearts at half a year of age from CSQ2(−/−) were normal based on gross inspection, echocardiography and HE staining. But the increased left atrial weights of CSQ2(−/−) at half a year of age which we reported previously [14] suggested to us a beginning heart disease and motivated us to study older (adult) CSQ2(−/−) in the present work. We hypothesized...
that older CSQ2(−/−) might develop cardiac hypertrophy and signs of heart failure which might be explained by an increased diastolic Ca2+ due to impaired SR Ca2+ release. An increase in diastolic Ca2+ can lead not only to arrhythmias but also activates Ca2+ kinases and thus induces gene transcription programs leading to cardiac hypertrophy and heart failure (overview in [34]). In agreement with other groups, we found bradycardia in CSQ2(−/−) in ECG [15, 35]. The bradycardia might indicate a role of CSQ2 in the cardiac clock, the pacemaker of the heart, which is known to be tightly linked to SR Ca2+ release [34]. In agreement with other groups, we found bradycardia in CSQ2(−/−) in ECG [15, 35]. The bradycardia might indicate a role of CSQ2 in the cardiac clock, the pacemaker of the heart, which is known to be tightly linked to SR Ca2+ release [34].

In order to confirm this, we measured beating rates in living adult animals. The literature suggests that the beating rate of adult animals is about 6–10 beats per minute [36]. In our study, we measured about 5–7 beats per minute in young CSQ2(−/−) and about 3–5 beats per minute in older CSQ2(−/−). This is consistent with the literature and the expected age-related decrease in beating rate.

Using fluorescence-activated cell sorting (FACS) with FlowJo Single-Cell Analysis Software, we analyzed leucocytes in the samples. Mean intensity of immunofluorescence signals using anti-mouse IgG, IgM, and DAPI conjugated with fluorescent dyes was measured using a Zeiss Axioskop 2 Plus and ImageJ software. Genotype indicates CSQ2(+/+) and CSQ2(−/−) mice. Adult means mice about 18 months old and young means about six months old. See Table 1 for exact ages. Numbers of animals are indicated in the last column.

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reported reduction of the expression of CSQ2 in acute hypoxia in neonatal rat cardiomyocytes [39] or an increase in isolated rat hearts after ischemia and reperfusion [40].

We would like to point out that we observed in the present report opposite effects of CSQ2 deletion on response to hypoxia in atrium versus ventricle, at least in the mouse. The fact that we present evidence for a detrimental effect of CSQ2 in the left ventricle and beneficial effects of CSQ2 in atrium might be related to the observation that the Ca$^{2+}$ handling is different in atrium versus ventricle in the mammalian heart. Anatomically, it is well established that the SR is larger and much more developed in the mammalian ventricle than in the mammalian atrium [34]. Furthermore, excitation–contraction coupling shows functional regional differences: in the mouse atrium, an increase in the beating rate will reduce force of contraction, whereas in the isolated mouse ventricular preparation, an increase in the beating rate will increase force (negative and positive “treppe” or staircase phenomena) [14, 34]. In addition, another difference between atrium and ventricle lies in the observation that the sarcoplasmic Ca$^{2+}$ flow is thought to be more important in atrium than ventricle of the mouse and rat to initiate excitation–contraction coupling [34]. How exactly CSQ2 would alter Ca$^{2+}$ handling in hypoxic atrial cells versus ventricular cells would be important to study in the future. Moreover, it needs to be confirmed that these regional differences occur also in carriers of loss of function mutations of CSQ2. We would predict a similar pattern in these patients.

It is currently unclear, why we noted fibrosis in CSQ2(−/−) ventricles. However, these findings are consistent with reports of others that fibrosis occurs in the pacemaker region in three- and twelve-month adult CSQ2(−/−) hearts [16]. They suggested that the increased basal cytosolic Ca$^{2+}$ levels in CSQ2(−/−) might activate Ca$^{2+}$-dependent enzymes that promote gene transcription of pro-fibrotic proteins. The same arguments could be made for the ventricular fibrosis that we report here. But this is just speculative as strong evidences are still missing. Alternatively, they speculated that the higher incidence of arrhythmias in CSQ2(−/−) led to arrhythmia-induced hypertrophy [16]. The fibrosis noted in this study may contribute to the impaired ejection fraction in the 18-month-old CSQ2(−/−), which we report here.

Meanwhile, it is accepted that cardiovascular diseases are associated with cardiovascular inflammation. This includes ischemic heart diseases like myocardial infarction where pro-inflammatory macrophages release pro-inflammatory cytokines and other immune cells, like T cells, for example, CD8 + and CD4 + T cells are involved [41]. But also hypoxia and reperfusion lead to cytokine release [41]. Moreover, cardiac hypertrophy as well as aging is associated with inflammation processes that are involved in the progress of non-ischemic heart failure [41]. In wild-type C57BL/6 J mice (the same genetic background as the CSQ2(−/−) mice), it was noted that anti-inflammatory macrophages decrease with age, whereas pro-inflammatory macrophages increase with age in the heart [42]. Others found that in cardiac aging of C57BL/6 J mice, the number of macrophages declined from 2–3 months to 12–15 months, whereas the number of granulocytes in the heart increased [20]. This was accompanied by a decline in fractional shortening and increased end-diastolic anterior wall thickness [20]. In cardiac lymph nodes, increased inflammation was seen upon aging because the number of CD4 + T cells rose [20]. Therefore, we expected that also in CSQ2(−/−) hearts, at least in the adult group, signs of inflammation should be visible because of the hypertrophy, fibrosis, and impaired cardiac function. But contrary to our expectations, we could not found indications to cardiac inflammation. In addition, one could ask whether calsequestrin was also present in blood cells like B cells and might be affected by our general knock-out of CSQ2 and might explain hypertrophy in the aging CSQ2(−/−) hearts. However, to the best of our knowledge, in blood cells only CSQ1 is present, which is coded by a different gene and that should not be altered in our CSQ2-deficient mice [14, 43, 44].

Limitations: a direct link between CSQ2 and fibrosis is missing and remains to be established. A direct mechanism starting in fibrocytes appears unlikely given the observation by others that cardiac fibrocytes do not express CSQ2 [4]. Therefore, indirect mechanisms like cardiac hypertrophy or pro-fibrotic mediators produced by the CSQ2(−/−) cardiomyocytes have to be considered. However, in future work it would be informative to measure intracellular Ca$^{2+}$ transients in cardiomyocytes from younger and older CSQ2(−/−) (also in hypoxia) and the effects of isoproterenol on these transients. Moreover, the biochemical interaction of RYR2 with triadin and junctin in younger and older CSQ2(−/−) should be studied. This could be combined with the study of ryanodine on intracellular Ca$^{2+}$ transients in cardiomyocytes of these mice, which was regrettably beyond the scope of the present paper.

In summary, the present data argue: CSQ2 is relevant in cardiac hypoxia and aging in the mammalian heart. Data in patients with functional ablation of CSQ2 are awaited with interest. The present work might motivate careful study of patients with functional loss of CSQ2 with the questions in mind, whether these patients are more likely to develop impaired ejection fraction in echocardiography than control patients and this information might be clinically helpful for risk stratification.

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Author contributions JN designed the research. KB, CS, and GR performed research; UG, UH, and SF analyzed the data. UG and JN wrote the paper. All authors have read and have approved submission of the manuscript.

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Data availability Data are available on reasonable request.

Declarations

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval The investigation conforms to the Guide for the Care and Use of Laboratory Animals published by the National Research Council (2011). Animals were handled and maintained according to approved protocols of the animal welfare committee of the University of Halle-Wittenberg, Halle, Germany (approval reference number 42502-2-1506 MLU). The manuscript was prepared as per the ARRIVE guidelines.

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