KCNJ11 downregulation stimulates cardiac cell apoptosis in myocarditis

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One of the inflammatory heart diseases, viz. acute myocarditis, occurs due to cardiac cell death. However, the molecular mechanism underlying cell death remains largely unexplored. In this study, we report that the down-regulation of KCNJ11, a central subunit of the ATP-sensitive potassium (KATP) channel plays a key role in the reduction of blood glucose, and is involved in apoptosis of cardiac cells. Using proteomics analysis of experimental autoimmune myocarditis (EAM), we show that the KCNJ11 level marked by decreased, whereas Camk2a expression increased significantly in EAM tissues by 16 and 20 days, compared to control tissues. Using 1H-MAS NMR we also show that glucose levels were slightly elevated in EAM tissues. In vitro assays using H9c2 cardiac cells revealed that both lipopolysaccharide (LPS) and high glucose treatment decreased cell viability, in which toxicity was attenuated by treatment with KATP pharmacological openers, but not by the KATP blockers (Gli and 5-HD). Apoptosis induced by LPS or high glucose treatment was suppressed by Ca2+ chelator (BAPTA-AM) treatment. We found that KCNJ11 levels had decreased in cardiac cells by LPS or high glucose treatment, and siRNA-mediated knockdown of KCNJ11 expression further stimulated the LPS- or high glucose-induced apoptosis. Together, our results demonstrate first that KCNJ11 is down-regulated under inflammation and high glucose conditions and its inactivation facilitates cardiac cell apoptosis. We assume that down-regulation of KCNJ11 has an effect on the development of myocarditis.

Keywords: Apoptosis, blood glucose lipopolysaccharide, cardiac cells, myocarditis.

Inflammatory heart disease, including myocarditis, is caused by infection of virus and protozoa, autoimmune responses and drugs1–3. Acute myocarditis is a heart disease that cannot be predicted because several clinical courses are asymptomatic or a variety of symptoms1,2. Acute myocarditis is induced by massive cardiac cell death, in which the accumulation of fibres results in fibrosis and heart attack1,4. However, the signalling pathway of cardiac cell death remains largely elusive.

One of the well-known myocarditis models is experimental autoimmune myocarditis (EAM), in which myosin injection triggers autoimmunity in heart tissues. The rat EAM shows elevated levels of toll-like receptor 4 (TLR4), caspase-3 cleaved form and tumour necrosis factor-α (TNF-α), proof that rat EAM mimics viral myocarditis1,5,6. The TNF-α related apoptosis-inducing ligand (TRAIL), Fas (CD95/APO1) and TNF-α cell death receptors can control cell apoptosis and also trigger intrinsic apoptosis pathway7–9. Previous studies showed that mitochondrial Ca2+ stimulates an increase in reactive oxygen species (ROS) and induces mitochondrial permeability and apoptosis10,11. High levels of glucose are also known to trigger mitochondrial dysfunction, caspase activation and apoptosis12,13.

The ATP-sensitive potassium channels (KATP) regulate insulin secretion in pancreatic beta cells14,15. The octameric complex containing the Kir6.2 (KCNJ11) tetramer at the centre is the KATP channel, surrounded by four SUR1 subunits, which responds to glucose concentration via its opening/closing for regulating blood glucose levels16–18. Under high glucose conditions, the KATP channel is closed, resulting in cell membrane depolarization, Ca2+ uptake and apoptosis16,19,20. Studies also showed that inactivation of KCNJ11, the central subunit of the KATP channel, is implicated in cardiac cell death and heart diseases21,22.

To explore the mechanistic basis of cardiac cell death leading to acute myocarditis, we analysed the proteomics profiles using rat EAM model. We show that the KCNJ11 level is downregulated during acute myocarditis and its inactivation facilitates cardiac cell apoptosis triggered by inflammation or high glucose-induced stresses.

Material and methods

Experimental autoimmune myocarditis induced by myosin in rats

The EAM was induced by injecting 1 mg/100 μl of porcine heart myosin and the Freund’s adjuvant to footpad of...
seven-week-old Lewis rats ($n=15$) twice on the zeroth and sixth day, in addition to injecting 500 ng of *Bordetella pertussis* toxin intraperitoneally on the first day. The cardiac tissues were obtained at 9, 16 and 20 days after the first injection. The vehicle without *B. pertussis* toxin or pig heart myosin was injected into the control group ($n=5$). All animal procedures were performed with permission from the ‘Institutional Animal Care and Use Committee at the Korea Basic Science Institute’ (KBSI-AEC1602).

**Reagents treatment and H9c2 cardiac cell culture**

Under condition 5% CO$_2$ humidifier at 37°C and incubator, the cells were cultured in culture flasks. The cells were obtained from rat embryonic heart tissue (ATCC, Manassas, VA, USA). Penicillin–streptomycin (P/S) 1% and foetal bovine serum (FBS) 10% were added to Dulbecco’s modified Eagle’s medium (DMEM, Gibco, Basel, Switzerland) 23. Cytotoxicity was activated under glucose or LPS conditions. The reagents (Sigma-Aldrich, St. Louis, USA) were used in the treatment of cardiac cells at 37°C for 0, 6, 18 and 24 h. Diaxoxide (DZ), glyburide (Gli), pinacidil (Pin) and 5-hydroxy decanoic acid (5-HD) were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA) 23.

**1H HR-MAS NMR analysis in EAM heart tissue**

$^1$H HR-MAS NMR analysis in EAM heart tissue was done according to previous experiments 23. A 4 mm diameter zirconia rotor with inserts (total volume of 40 μl) was used, where 5–10 mg of heart tissue samples was placed 23. Distilled water containing 0.05% (wt) of 3-(trimethylsilyl)-propionic-$d_4$ acid (TSP) to provide $^2$H field locking was used to fill the rest of the rotor volume 23. The $^1$H HR-MAS NMR spectra were recorded (Unity Inova 500 spectrometer; Agilent Technologies Inc., USA) under the operating conditions at 500.13 for $^1$H, a spin rate of 2300 Hz, and temperature of 283 K and 4 mm gHX nano-probe head was attached for acquisition 23. The Carr–Purcell–Meiboom–Gill (CPMG) spin–echo pulse sequence was used to collect the $^1$H NMR spectra with total spin–spin relaxation time of 300 ms and spin–echo delay ($\tau$) of 400 μs (refs 23, 24). For each sample, 128 transients into 32 K data points were collected through a spectrum width of 8004.8 Hz, an acquisition time of 2 sec with water pre-saturation during the relaxation delay. A line-broadening function of 0.3 Hz was used to multiply the spectra and was then phase-adjusted, Fourier-transformed (FT) and manually baseline-corrected, and referenced to TSP (0.00 ppm) 23. Signal assignments for representative tissue samples were achieved using heteronuclear single quantum correlation (HSQC), total correlation spectroscopy (TOCSY) and two-dimensional $^1$H–$^1$H correlation spectroscopy (COSY), the 500 MHz library from Chenomx NMR suite version 6.0 and compared with the literature 23,25,26.

**Protein level analysis in heart tissues and cardiac cells**

The total proteins of cardiac cells and EAM heart tissues were extracted in the RIPA buffer. Next, 10–15% SDS-PAGE gel was used to separate the 20–30 μg of total proteins and then transferred to a polyvinylidene difluoride (PVDF) membrane. The primary antibodies for GAPDH (CST #2118), cytochrome c (CST #4272), caspase-3 (CST #14220), β-actin antibody (CST #3700) and Kir6.2 (KCNJ11; PA5-26218) were purchased from Thermo Scientific (Waltham, MA, USA), Cell signalling (CST, Danvers, MA, USA) and Santa Cruz Biotechnology (Santa Cruz, Calif. USA).

**mRNA analysis in heart tissues and cardiac cells**

Total RNA was extracted from EAM hearts tissues or the H9c2 cardiac cells using 0.2 ml of TRIzol reagent (Thermo Scientific) and converted to cDNA using the First Strand cDNA Synthesis Kit (Thermo Scientific). The qRT-PCR reaction was duplicated in 10 μl of cDNA diluted 10 times in water using SYBR-Green PCR Master Mix (Thermo Scientific). Using Step One Plus Real-Time PCR System (Thermo Scientific), the qRT-PCR working conditions were as follows: 3 min at 95°C followed by 40 cycles at 95°C for 15 sec (denaturation), 60°C for 25 sec (annealing temperature), and 72°C for 25 sec (extension). The 18S rRNA or GAPDH level was used to normalize the quantification of PCR products of the target genes. The primers used were – Camk2a sense primers: 5’-CGAGGgcctggactttca-3’, and Camk2a antisense primers: 5’TGACGGGCTTgctttca-3’. and Camk2a antisense primers: 5’T-GACCGGGCttgctttca-3’. and Camk2a antisense primers: 5’T-GACCGGGCttgctttca-3’. and Camk2a antisense primers: 5’T-GACCGGGCttgctttca-3’.

**siRNA transfection into H9c2 cells**

The cardiac cells were seeded in six-well plates without antibiotics. A transient transfection with lipofectamine 2000 (Invitrogen, Waltham, MA, USA) was applied when the cells were 60–75% confluent and the experiment was in accordance with the manufacturer’s instructions 23. In this experiment, siRNA (10 pmol) and lipofectamine 2000 (2.5 μl) were individually diluted in 125 μl serum-free Opti-MEM (GIBCO/Invitrogen) and incubated at
25°C for 5 min, and then combined. The siRNA–Lipofectamine mixture was incubated at 25°C for 20 min and added to the cell culture in a 5% CO₂ humidifier at 37°C for 16 h. The next day, PBS was used for cell wash and the cells were cultured in fresh medium with 1% antibiotics. At 24, 48 and 72 h after transfection, the cells were harvested. The siRNA knockdown constructs used in the study were obtained from Cosmogenetech (Seoul, Korea) and their sequences are listed here: Kcnj11-rat-335 sense primers: 5′-CCGCCUUGGUCCAAAGAAATT-3′ and antisense primer: 5′-UUUCUUGGACACGGAAGC-GGTT-3′; Kcnj11-rat-689 sense primer: 5′-GCAGAAUAUCGUAGGCUATT-3′ and antisense primer: 5′-UAGCCCUAGCAUUCUGCTT-3′; Kcnj11-rat-1384 sense primer: 5′-GCAUCUCCCGGAAUCCUUTT-3′ and antisense primer: 5′-AAGGAUCCCGAGAGAGUG-CTT-3′. The random sequence control mixtures were also provided by the manufacturer and treated as instructed.

Cell viability assay in H9c2 cells

The MTT proliferation assay kit (ATCC) was used for cell viability analysis, according to the manufacturer’s instructions. The 10% FBS/DMEM medium was used to culture cardiac cells in 96-well plates for 16 h. At the end of the treatment, 10 μl of MTT reagent (0.5 mg/ml MTT) was treated to cells with or without the exposure of siRNA constructs, LPS or glucose, which was then incubated in a 5% CO₂ humidifier at 37°C for additional 4 h.

Statistics

At least three independent experiments were performed to obtain the mean ± standard deviation (SD) of the data. The statistical significance of the differences was confirmed using an unpaired Student’s t-test. The SPSS Statistical Software version 24 (IBM, Armonk, NY, USA) was used for statistical analysis and one-way analysis of variance (ANOVA) was used to determine the P value. A P-value of less than 0.05 was considered statistically significant (**).

Results

Apoptosis induction in EAM heart tissues

To characterize cardiac cell apoptosis involved in myocarditis pathogenesis, we used the rat EAM model in which myocarditis was induced by myosin injection. Following injection of myosin or adjuvant (control) into rats, the EAM heart tissues were analysed on days 9, 16 and 20. As expected, myosin-induced EAM lesions (discoloured surface) were clearly seen at day 20 post-injection (Figure 1a). To assess apoptosis induction, we performed immunoblot analyses of cleavage caspase-3 and cytochrome c, which are elevated through activation of the mitochondrial pathway of apoptosis. Compared to control tissues, EAM tissues at day 20 showed markedly increased levels of both cytochrome c and cleaved caspase-3, confirming that active apoptosis is accompanied by myocarditis development (Figure 1b and c).

Elevated levels of glucose and KCNJ11 down-regulation in EAM tissues

Previous studies demonstrated that the inactivation of KATP channel, a key regulator of calcium and glucose, is associated with the development of heart disease. In the present study, 1H HR–MAS NMR revealed that glucose levels were increased in EAM tissues, compared to control (Figure 2a). We thus measured the level of KCNJ11, a subunit of the KATP channel which controls calcium transport in response to glucose levels. An immunoblot assay showed that the level of KCNJ11 protein gradually decreased in EAM (Figure 2b and c). In addition, we observed that the mRNA level of Camk2a (Ca²⁺ associated calmodulin) gene was significantly increased in EAM tissues compared to control (Figure 2d). Given that elevated calcium in mitochondria activates mitochondrial apoptosis signalling, the results suggest that myocarditis pathogenesis might be associated with the elevation of glucose and calcium, and subsequent mitochondrial dysfunction and apoptosis.

KATP openers attenuate cardiac cell death induced by LPS or high glucose

To confirm whether abnormal KATP function is associated with cardiac cell death, we tested the effects of KATP opener (diazoxide (DZ) and pinacidil (Pin)), KATP-blocker glyburide (Gli) and 5-hydroxy decanoic acid (5-HD) or Ca²⁺ chelate (BAPTA-AM) on the cell viability of H9c2 cells exposed to LPS or high glucose levels. MTT assay showed that compared to cells treated with LPS or glucose alone, cells co-treated with DZ (100 μM) or Pin (50 μM) exhibited 19–26% increase in viability (Figure 3a and b). Conversely, cells co-treated with Gli (100 μM) or 5-HD (100 μM) exhibited 43% reduction in viability compared to LPS- or glucose-treated control cells. Cells treated with BAPTA-AM showed 17–24% increase in viability compared to control (Figure 3a and b). Figure 3c and d shows the effects of KATP openers, blockers or Ca²⁺ chelate BAPTA-AM on cell death. Consistent with MTT assay results, cells treated with DZ, Pin or BAPTA-AM were protected against LPS or glucose alone compared to control cells.
Figure 1. Changes in heart morphology and in apoptosis gene in the experimental autoimmune myocarditis (EAM) model during EAM induction. a, EAM rat heart on days 9, 16 and 20. b, Protein levels of cytochrome-c and cleaved caspase-3 proteins EAM heart. c, Quantitative confirmation of cytochrome-c and cleaved caspase-3 protein levels confirmed in EAM heart at 20 days. Data represent mean ±SD (n = 3). *P < 0.05, **P < 0.01 in two-tailed Student’s t-test.

Figure 2. Changes in glucose levels by 1H HR-MAS NMR and KCNJ11 and Camk2a levels in EAM model. a, Changes in glucose level by 1H HR-MAS NMR in EAM heart tissue. b, Western blot showing KCNJ11 protein levels at 9, 16 and 20 days in the EAM model. c, Quantification of KCNJ11 protein levels confirmed on days 9, 16 and 20 in the EAM heart. d, EAM cardiac tissue confirming the level of Camk2a mRNA at 9, 16 and 20 days. Data represent mean ±SD (n = 3). *P < 0.05, **P < 0.01 in two-tailed Student’s t-test.

KCNJ11 reduction facilitates cardiac cell apoptosis induced by LPS or high glucose levels

We tested whether reduced expression of KCNJ11 was involved in cardiac cell apoptosis. Western blot assay showed that LPS or glucose treatment downregulated KCNJ11 expression in cardiac cells, which was accompanied by an increase in cleaved caspase-3 levels (Figure 4a). A drastic decrease in KCNJ11 was observed in cells treated with LPS, compared to those treated with glucose (Figure 4b). We also observed that the mRNA level of Camk2a was increased in cells treated with glucose or LPS for 24 h (Figure 4c). Moreover, siRNA-mediated knockdown of KCNJ11 expression further promoted glucose- or LPS-mediated apoptosis (Figure 4d and e).

Discussion

Pro-inflammatory cytokines such as TNF-α play an important role in cardiac cell apoptosis triggered by pathogen invasion and inflammation. It has been also reported that LPS-exposed heart injury is related to mitochondrial dysfunction and cell death in the EAM model. However, the molecular mechanism underlying cardiac cell death during myocarditis development remains unknown. In this study using myosin-induced rat EAM model, we have identified that KCNJ11 downregulation is implicated in induction of heart cell apoptosis under inflammatory conditions.

It is known that lactate levels are elevated in EAM tissues and increased lactate levels are associated with heart attack. Lactate moves to the liver and produces glucose through gluconeogenesis, and this glucose is again used as an energy source for muscles. It is known that high levels of glucose cause mitochondrial dysfunction. In response to high glucose levels, the
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Figure 3. Cell viability assay by treatment with diazoxide (DZ), glyburide (Gli), pinacidil (Pin), 5-hydroxy decanoic acid (5-HD) and Ca$^{2+}$ chelator (BAPTA-AM) on H9c2 cells under lipopolysaccharide (LPS) or high glucose conditions. a, b, MTT analysis confirmed that the cells treated with glucose or LPS showed change in cell viability. However, glucose or LPS toxicities were significantly by 50 μM Pin or 100 μM DZ or 0.5 μM BAPTA-AM. Glucose or LPS toxicity was found to increase with 100 μM Gli or 100 μM 5-HD. c, d, Cells were treated without or with glucose or LPS, and 100 μM DZ, 50 μM Pin, 100 μM Gli, 100 μM 5HD or 0.5 μM BAPTA-AM to confirm cell morphology. Data represent mean ±SD (n = 3).

Figure 4. The exposure of LPS or glucose induces cleaved caspase-3 gene expression and the knock-down of KCNJ11 induces cytotoxicity of LPS or glucose in H9c2 cells. a, In Western blot, the levels of KCNJ11 and cleaved caspase-3 protein were identified in H9c2 cells after 10 μg LPS or 50 mM glucose exposure. b, In protein quantification, cleaved caspase-3 levels increased in 50 mM glucose or 10 μg LPS for 24 h, while KCNJ11 levels decreased in 24 h. c, Checking the Camk2a mRNA level in cells using qRT-PCR while being exposed to glucose or LPS for 24 h. d, In immunoblot, 50 mM glucose or 10 μg LPS treatment in the cells induced the expression of cleaved caspase-3 and siRNA of KCNJ11 increased the glucose or LPS effects. The loading control was β-actin. e, Quantification of caspase-3 and KCNJ11 protein levels in H9c2 cells with siRNA. In the quantification of cleaved caspase-3 and KCNJ11 proteins in H9c2 cells, 50 mM glucose or 10 μg LPS treatment substantially significantly increased the level of cleaved caspase-3, decreased the level of KCNJ11 and the glucose or LPS effects were almost completely increased by the siRNA of KCNJ11. Data are presented as mean ±SD (n = 3). *P < 0.05, **P < 0.01 in two-tailed Student’s t-test.
Ca^{2+}-calmodulin-dependent proteolytic enzyme II (CaMKII) is a serine/threonine protein kinase that phosphorylates multiple excitation–contraction binding proteins^{36–41}. CaMKII is important in the pathogenesis of hypertrophy and heart dysfunction^{42}. In the present study, we found that EAM tissues have elevated levels of glucose, reduced expression of KCNJ11 and increased expression of CaMKII. Using cultured cardiac cells, we also verified that LPS- or glucose-induced reduction of cardiac cell viability was attenuated when the cells were co-treated with KATP openers (DZ and Pin). Consistently, cardiac cell viability was attenuated when the cells were treated with LPS or high glucose. The immunoblot assay of cleaved caspase-3 also revealed that the treatment using LPS or high glucose was further elevated by combined treatment with KATP openers. This suggests that the treatment using LPS or high glucose induces mitochondrial dysfunction and apoptosis in heart cells. We conclude that when KCNJ11 was inhibited, apoptosis occurs in heart cells, which affects the development of myocarditis.

**Conflict of interest:** The authors declare no competing financial interests.

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