A porcine model of heart failure with preserved ejection fraction

Charles, Christopher J.; Lee, Philip; Li, Renee R.; Yeung, Teresa; Mazlan, Stephane M. Ibrahim; Tay, Zhi Wei; Abdurrachim, Desiree; Teo, Xing Qi; Wang, Wei-Hsin; de Kleijn, Dominique P.

Published in:
ESC Heart Failure

DOI:
10.1002/ehf2.12536

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher's PDF, also known as Version of record

Publication date:
2019

Link to publication in University of Groningen/UMCG research database

Citation for published version (APA):

Copyright
Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

Take-down policy
If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): http://www.rug.nl/research/portal. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.
A porcine model of heart failure with preserved ejection fraction: magnetic resonance imaging and metabolic energetics

Christopher J. Charles1,2,3*, Philip Lee4, Renee R. Li1,2, Teresa Yeung1,2, Stephane M. Ibrahim Mazlan1,2, Zhi Wei Tay4, Desiree Abdurrachim4, Xing Qi Teo4, Wei-Hsin Wang4, Dominique P.V. de Kleijn5,6, Patrick J. Cozzone4, Carolyn S.P. Lam1,7,8,9,10 and A. Mark Richards1,3

1Cardiovascular Research Institute, National University Heart Centre, Singapore; 2Department of Surgery, Yong Loo Lin School of Medicine, National University of Singapore, Singapore; 3Christchurch Heart Institute, University of Otago, Christchurch, New Zealand; 4Singapore Bioimaging Consortium, Agency for Science, Technology and Research, Singapore; 5Department of Vascular Surgery, University Medical Centre, Utrecht, Utrecht, The Netherlands; 6Netherlands Heart Institute, Utrecht, The Netherlands; 7National Heart Centre Singapore, Singapore; 8Faculty of Medicine, Duke-National University Singapore, Singapore; 9University Medical Centre Groningen, Gronigen, The Netherlands; 10The George Institute for Global Health, Sydney, Australia

Abstract

Aims A significant proportion of heart failure (HF) patients have HF preserved ejection fraction (HFpEF). The lack of effective treatments for HFpEF remains a critical unmet need. A key obstacle to therapeutic innovation in HFpEF is the paucity of pre-clinical models. Although several large animal models have been reported, few demonstrate progression to decompensated HF. We have established a model of HFpEF by enhancing a porcine model of progressive left ventricular (LV) pressure overload and characterized HF in this model including advanced cardiometabolic imaging using cardiac magnetic resonance imaging and hyperpolarized carbon-13 magnetic resonance spectroscopy.

Methods and results Pigs underwent progressive LV pressure overload by means of an inflatable aortic cuff. Pigs developed LV hypertrophy (50% increase in wall thickness, \( P < 0.001 \), and two-fold increase in mass compared to sham control, \( P < 0.001 \)) with no evidence of LV dilatation but a significant increase in left atrial volume (\( P = 0.013 \)). Cardiac magnetic resonance imaging demonstrated T1 modified Look-Locker inversion recovery values increased in 16/17 segments compared to sham pigs (\( P < 0.05-P < 0.001 \)) indicating global ventricular fibrosis. Mean LV end-diastolic (\( P = 0.047 \)) and pulmonary capillary wedge pressures (\( P = 0.008 \)) were elevated compared with sham control. One-third of the pigs demonstrated clinical signs of frank decompensated HF, and mean plasma BNP concentrations were raised compared with sham control (\( P = 0.008 \)). Cardiometabolic imaging with hyperpolarized carbon-13 magnetic resonance spectroscopy agreed with known metabolic changes in the failing heart with a switch from fatty acid towards glucose substrate utilization.

Conclusions Progressive aortic constriction in growing pigs induces significant LV hypertrophy with cardiac fibrosis associated with left atrial dilation, raised filling pressures, and an ability to transition to overt HF with raised BNP without reduction in LVEF. This model replicates many aspects of clinical HFpEF with a predominant background of hypertension and can be used to advance understanding of underlying pathology and for necessary pre-clinical testing of novel candidate therapies.

Keywords Left ventricular pressure overload; Hypertrophy; Cardiac fibrosis; CMRI hyperpolarizer; BNP; Animal model

Introduction

Heart failure (HF) is the final common pathway of a myriad of metabolic and cardiovascular diseases and imposes a significant healthcare burden worldwide. Although there have been significant improvements in clinical management and outcomes, morbidity and mortality remain high, and there is an indisputable need for improved treatment options. The pathophysiology of HF is complex. Whilst many patients present clinically as HF with reduced ejection fraction (HFrEF) (≤40%
EF), a significant proportion of HF patients have preserved EF (HfPfEF) demonstrating a reduced ability of the heart to relax but with EF above 50%.[9-10] Our recently reported prospective multi-ethnic study shows that HfPfEF accounts for ~30% of all HF cases in parallel New Zealand and Singapore cohorts. HfPfEF is characterized by older age, a greater prevalence of females, and predominant background hypertension (78%) compared with HFrEF.[9-10] Recent reports of HfPfEF have demonstrated mortality or HF hospitalization.।

Clinical trials in patients with HfPfEF have failed to demonstrate improvements in mortality. A recent meta-analysis examining data from 25 trials (>18,000 patients) showed no beneficial effect from angiotensin-converting enzyme inhibitors, angiotensin receptor blocker, or mineralocorticoid antagonists on mortality or HF hospitalization.[11] The lack of effective treatments for HfPfEF remains a critical unmet need. A significant obstacle to therapeutic innovation in HfPfEF is the absence of pre-clinical models including large animal models that, unlike rodent models, permit detailed instrumentation and extensive imaging and sampling protocols.[9-13] Although several large animal models have been reported,[9-13] none fulfill all the features present in human disease, and few demonstrate progression to HF. We hypothesized that enhancing a porcine model of progressive left ventricular pressure overload (LVPO)”[9-13] with greater characterization would demonstrate incipient HF with some animal progressing to definitive HF confirming that this model is suitable for ongoing pre-clinical studies of HfPfEF. Accordingly, we established and enhanced this model of LVPO and characterized HF in this model with clinical observations, filling pressures and BNP measurements and also advanced cardiometabolic imaging using cardiac magnetic resonance imaging (CMRI) and hyperpolarized carbon-13 magnetic resonance (hp13CMRI) spectroscopy.

Materials and methods

The study protocol was approved by the Institutional Animal Care and Use Committees of National University of Singapore and Biological Research Centre, Agency for Science, Technology and Research. We built on a previously reported progressive LVPO model”[9-13] with key enhancements to progress pigs to evidence-based HfPfEF. Thirty-nine female Yorkshire × Landrace pigs (18–23 kg) were housed individually receiving standard care. For echocardiography and CMRI, pigs were pre-medicated with intramuscular ketamine (Ceva Animal Health, Glenorie NSW, Australia) (10 mg/kg), atropine (Atrosite, Troy Laboratories, Glendenning NSW, Australia) (0.40 mg/kg), and midazolam (Dormicum, Hoffmann-La Roche, Basel, Switzerland) (0.6 mg/kg), intubated, ventilated, and maintained with inhalation isoflurane (Attane, Piramal Critical Care, Bethlehem, PA). Following Day 0 imaging, pigs were switched to total intravenous (i.v.) anaesthesia [midazolam 2.5–4 mg/kg/h, alfentanil (Rapifen, GlaxoSmithKline, Torrile, Italy) 250–400 μg/kg/h, and pancuronium (Rotexmedica, Trattau, Germany) 0.25–0.3 mg/kg/h] for surgical implantation of aortic cuff. Other medications included oral amiodarone (Cordarone, Sanofi-Aventis, Paris, France) (200 mg daily), clopidogrel (Ceruvin Ranbaxy, Kedah, Malaysia) (75–300 mg), and aspirin (Cardiprin, Reckitt Benckiser Healthcare, Hull, UK) (100 mg daily). Pigs also received augmentin (SmithKline Beecham, Worthing, UK) (15 mg/kg i.v. then 312 mg orally for 5 days) and fentanyl (Durogesic®, Janssen Pharmaceutical, Beere, Belgium) transdermal patches (5 μg/h for 4 days). All pigs underwent left lateral thoracotomy and pericardiotomy to isolate the aorta. In 33 pigs, an inflatable cuff (Access Technologies, Skokie, IL, www.norfalkaccess.com) was placed around the aortic root and connected to an injectable port tunnelled subcutaneously adjacent to the scapula. Sham pigs (n = 6) had thoracotomy without cuff placement. Pigs recovered for 1 week prior to commencement of cuff inflation.

Pigs underwent weekly echocardiography prior to progressive cuff inflation over 4 weeks titrated to achieve incremental trans-constriction pressure gradients (PGs) of 20, 40, 60, and 80 mmHg (assessed echocardiographically). From week 4, the PG was maintained at 80 mmHg until completion of final magnetic resonance imaging (MRI) and HfPfEF measurements after which pigs were terminated. Cardiac left ventricle (LV) and lungs were excised, weighed, and samples taken for subsequent histology. Sham pigs underwent identical procedures. Trans-thoracic and trans-oesophageal echo were used to acquire standard cardiac structure and function measures (Philips E PIQ 7 machine). Trans-oesophageal echo, with tissue Doppler imaging, of the aortic root displayed the cuff-induced aortic constriction and defined PGs enabling titration of cuff inflation up or down as necessary.

CMRI was performed prior to thoracotomy and repeated either Day 35 or Day 42 (sham and 2nd half cuff pigs) using Siemens 3T MRI scanner (MAGNETOM Skyra, Siemens Healthineers AG, Erlangen, Germany). Standard CMRI protocol included cine imaging [global and regional LV systolic function including LV volumes, mass, and LV ejection fraction (LVEF)], and T1-mapping from Modified Look-Locker Imaging (MOLLI). MRI analysis was performed using SEGMENT v2.2 R6338 (http://segment.heiberg.se). A subset of pigs underwent tandem CMRI and CMRI hyperpolarizer on Day 42.

HfPfEF spectroscopy was performed in the same scanner as CMRI, using 13C TX quadrature volume coil/13C RX 16-channel flex array coil (Rapid Biomedical, Rimpar, Germany). [1-13C]pyruvate (7.18 mmol/L; Cambridge Isotope Laboratories) doped with AH111501 (15 mM) was inserted into polarizer (SPINLab, GE Healthcare, Chicago, IL) for microwave irradiation. The polarized sample was subsequently
dissolved yielding a solution of 250 mmol/L [1-13C]pyruvate (pH = 7.2–7.7) for immediate injection into animals at a dose of 0.2 mmol/kg body weight. 13C-MRS started just before i.v. bolus injection of hyperpolarized [1-13C]pyruvate. The build-up of metabolites was followed for ~2 min after injection, with spectra sampled every two heartbeats. Quantification of metabolites was performed using AMARES in the jMRUI software package (http://www.jmrui.eu). 13C-MR spectra were summed over 40 s upon metabolite appearance with peaks fitted for [1-13C]lactate, [1-13C]pyruvate hydrate, [1-13C]alanine, and [13C]bicarbonate all normalized to the [1-13C]pyruvate peak.

Ethlylenediaminetetraacetic acid blood samples drawn at baseline and pre-termination were centrifuged, and plasma was stored at −80°C prior to assay for porcine BNP at Christchurch Heart Institute using a well-validated assay.3 A subset of pigs (six sham and five cuff) underwent invasive pressure monitoring immediately after final echocardiography. Sheaths were placed in a jugular vein and carotid artery to allow passing under fluoroscopic guidance a Swan Ganz catheter into the pulmonary artery for measurement of pulmonary capillary wedge pressure (PCWP) and a fluid-filled catheter into the LV for measurement of end-diastolic pressure (LVEDP). LV tissue harvested post-mortem was fixed in buffered 10% formaldehyde prior to paraffin embedding. Picrosirius red staining was performed by standard methods to allow assessment of myocardial fibrosis using automated image analysis (IMAGEJ).

**Statistics**

Results are expressed as mean ± standard error of mean. Primary analysis was independent t-tests of time-matched data between the sham and cuff pigs. P < 0.05 was the threshold for statistical significance.

**Results**

Studies in sham pigs proceeded as planned with data collection complete. Among the cuff-implanted pigs, three experienced cuff failure (Days 21–28) and were excluded from analyses. A further five pigs were excluded due to premature death (<Day 28) with post-mortems revealing two fatal aortic occlusion deaths (last recorded PG >80 mmHg), one aortic rupture, one with large aortic and LV thrombi, and one pig died of bacterial pneumonia. A further five pigs successfully completed the protocol but exhibited no significant increase in LV mass despite equivalent PGs. Since LV hypertrophy (LVH) was the primary driver for development of HFpEF, these pigs were also excluded from further analysis. Twenty

---

**Figure 1** Serial body weight, pressure gradients across aortic arch, left ventricular mass/body weight (LVmass/BW) ratios, and echocardiography derived serial left ventricular dimensions including interventricular septum thickness at diastole, left ventricular posterior wall thickness at diastole and left ventricular internal diameter at diastole in 20 pigs with progressive inflation of aortic cuff to induce left ventricular hypertrophy (●) and six sham control (○) pigs. Values shown are mean ± standard error of the mean. Significant differences at time-matched points between the sham and aortic cuff to induce left ventricular hypertrophy pigs are indicated as follows: *P < 0.05, **P < 0.01 and ***P < 0.001.
pigs undergoing aortic cuffing completed the full protocol and developed LVH (cuff-LVH group). Data from this group are compared to animals undergoing sham procedures.

Both sham and cuff-LVH pigs gained weight throughout the protocol (Figure 2)[OAurl][?]–>][OAurl][?]–>]. The PG remained minimal (<6 mmHg) in the sham group but rose stepwise (in ~20 mmHg increments) over 4 weeks and thereafter remained stable (Figure 2)[OAurl][?]–>][OAurl][?]–>].

From 2 weeks, LV mass indexed to body weight (LVmass/BW) increased in the cuff-LVH group to be significantly (“two-fold”) higher than sham controls by the end of the study (Figure 4)[OAurl][?]–>][OAurl][?]–>]. Measures of LV mass by echocardiography, CMRI, and direct weighing at post-mortem correlated well. LV wall dimensions were stable in sham pigs but increased in the cuff-LVH pigs. Thickness of both interventricular septums and LV posterior walls in dias-tole increased in the cuffed pigs to be significantly higher than sham by Day 21 and eventually ~50% thicker (Figure 4)[OAurl][?]–>][OAurl][?]–>]. The LV internal diameter showed some increase in the cuff group, but the dimension was only greater than sham at Day 35 (Figure 4)[OAurl][?]–>][OAurl][?]–>].

LVEF remained stable with values for both sham and LVH-cuff group remaining in the 60–65% range (Figure 4)[OAurl][?]–>][OAurl][?]–>]. There was no significant change in either group in LV end-systolic volumes (LVESVs) or LV end-diastolic volumes (LVEDVs) (Figure 4)[OAurl][?]–>][OAurl][?]–>]. LV structure and function data measured by CMRI at baseline and the end of study corroborated changes measured by echocardiography with no significant changes comparing cuff-LVH to sham pigs for LVEDV, LSV, stroke volume, LVEF, cardiac output, or peak ejection rate (Figure 4)[OAurl][?]–>][OAurl][?]–>]. In contrast, left atrial end-diastolic volume (P = 0.013 compared to baseline) and peak filling rate (P < 0.01 compared with sham) increased in the cuff-LVH (Figure 4)[OAurl][?]–>][OAurl][?]–>].

Native T1 values derived from MOLLI did not differ significantly between sham and cuff-LVH pigs at Day 0 for 16/17 segments of the LV (with the mid-inferior segment the exception; Figure 4)[OAurl][?]–>][OAurl][?]–>]. By the end of study T1 values were significantly higher in the cuff-LVH pigs compared with sham for 16/17 segments with the apical-lateral segment the sole exception. Picrosirius red staining of sections from LV free wall (Figure 4)[OAurl][?]–>][OAurl][?]–>][OAurl][?]–>]. showed an increase in both interstitial and perivascular collagen deposition in the tissue from the cuff-LVH pig with mean intensities measured under a bright field showing a trend for difference and under polarized light being significantly different (P = 0.044) between LVH and sham groups.

Invasive intra-cardiac pressures (sub-study) showed normal range pressures in sham pigs, but both PCWP (P = 0.008) and LVEDP (P= 0.047) were raised in cuff-LVH pigs (Figure 4)[OAurl][?]–>][OAurl][?]–>]. Plasma BNP levels remained in the normal range in sham control pigs but on average increased by end of study in the cuff-LVH pigs being significantly higher than sham (P = 0.008) and baseline (Day 0) values (P = 0.02) (Figure 4)[OAurl][?]–>][OAurl][?]–>]. Clinical signs of HF were observed in at least one-third of the pigs including tachypnea, laboured breathing, cough, lung congestion on auscultation, cyanosis, and lethargy. One pig demonstrated major falls in arterial oxygen saturation (SPO₂ < 40%) on induction of anaesthesia for Day 35 imaging with extreme cyanosis and severe pulmonary oedema and pleural effusions. This pig underwent drainage of accumulated pleural fluid and received additional oxygen therapy and treatment with 2 mg/kg i.v. frusemide (Furosemide Fresenius Kabi, Bad Homburg, Germany) before stabilizing and proceeding to MRI and echo, followed by partial removal of glycerol from the aortic cuff to stabilize PG. It recovered to survive to end of the protocol (Day 42).

Representative cardiac ³¹C-MR spectra after injection of hyperpolarized [1-13C]pyruvate from a normal (sham) pig
Figure 3  Cardiac magnetic resonance imaging derived left ventricular end diastolic volume, end systolic volume, stroke volume, ejection fraction, peak filling and ejection rates, cardiac output and left atrial end-diastolic (ED) volume in 20 aortic cuff to induce left ventricular hypertrophy (●) and six sham control (◯) pigs. Values shown are mean ± standard error of the mean. Left atrial end-diastolic volume was significantly increased at 42 days compared with baseline (*P < 0.05) and left ventricular peak filling rate increased in aortic cuff to induce left ventricular hypertrophy pigs compared with sham (**P < 0.01).
demonstrates that [1-13C]pyruvate is metabolized into [1-13C]lactate, [1-13C]alanine, [13C]bicarbonate via lactate dehydrogenase, alanine aminotransferase (ALT), and pyruvate dehydrogenase (PDH), respectively with metabolic products visible within 2 min (Figure 4). Compared to sham pigs, cuff-LVH pigs showed increases in [1-13C]alanine ($P < 0.01$) and [13C]bicarbonate ($P < 0.05$) with no increase in [1-13C] lactate representing ~2.6-fold and ~6-fold increase in ALT and PDH activity, respectively (Figure 4).

**Figure 4** Top panel – cardiac MRI T1 MOLLI values across 17 segments of the LV at baseline and end of study in 20 cuff-LVH (●) and 6 sham control (◯) pigs. Bottom panels – examples on picrosirius red staining of sections of the LV free wall and group mean intensities from sham and cuff-LVH pigs measured under bright field and polarizing light. Values shown are mean + SEM. Significant differences at time-matched points between the sham and cuff-LVH pigs are indicated as follows: *$P < 0.05$, †$P < 0.01$ and ‡$P < 0.001$. 

Discussion

This porcine HFpEF model demonstrates significant LVH (doubling of LV mass) without LV dilation similar to that previously reported. Unique features of our model that document HF include comprehensive serial imaging (echocardiography and MRI) clearly demonstrating global ventricular fibrosis by MRI T1 MOLLI, a significant increases in plasma BNP, and significantly elevated PCWP and LVEDP. Hyperpolarized $^{13}$C-MR studies showed increased
manifested in clinical symptoms, raised LV HF with frequent progression to frank decompensated HFpEF.

ALT and PDH activity in HFpEF pigs. Pigs developed incipient HF with frequent progression to overt HF con- straining that this provides a clinically relevant model suitable for ongoing preclinical studies.

Figure 6 Metabolic signals from hyperpolarizing 13C magnetic resonance imaging (A) 13C spectra from a normal (sham) pig and (B) metabolite/pyruvate signal from 11 aortic cuff to induce left ventricular hypertrophy (*) and six sham control (o) pigs. Values shown are mean ± standard error of the mean. The units are in arbitrary units and normalized to the [1-13C] pyruvate peak amplitude of each animal. Significant differences at time-matched points between the sham and aortic cuff to induce left ventricular hypertrophy pigs are indicated as follows: *P < 0.05 and **P < 0.01.

ALT and PDH activity in HFpEF pigs. Pigs developed incipient HF with frequent progression to frank decompensated HFpEF manifested in clinical symptoms, raised LV filling pressures and, in some pigs, frank pulmonary oedema. Taken together, full characterization of this enhanced LVPO model demonstrates development of incipient HF with some animals progressing to overt HF confirming that this provides a clinically relevant model suitable for ongoing preclinical studies of HFpEF.

Our model is based the porcine LVPO model first reported by Spinale et al.10 [OAurl] –10 impaired LV contraction and systolic dysfunction;11 [OAurl]–11 arrhythmias, [OAurl]–12 and impaired coronary blood flow13 [OAurl]–13 and is a powerful indicator of adverse outcomes in heart disease.14,15

Spinale/Zile had previously demonstrated LVH in the LVPO model is associated with increased myocardial stiffness (three-fold) and total collagen (1.5-fold).10 [OAurl]–10 [OAurl]–6 MRI T1 MOLLI results here clearly demonstrated global increases fibrosis across 16/17 segments of the LV. Only the apical lateral segment failed to show a definitive rise in T1 value. Nevertheless, picrosirius red staining at this site demonstrated significant collagen deposition (Figure 6) [OAurl]–>6 [OAurl]–>6, but rather than representative, it likely underestimated overall ventricular fibrosis in this model.

Invasive pressures measured in a subset of the LVH-cuff animals indicated elevations of PCWP and LVEDP to levels equivalent to those observed in mild–moderate clinical HF. The BNPs are long established biomarkers for the
diagnosis and prognosis of HF."")[OAurl]([?–]?>)[OAurl][?–]?>16 We used our well-validated assay for porcine/ovine BNP")[OAurl]([?–]?>)[OAurl][?–]?> to find elevated mean BNP levels with three samples from three pigs in the mild range (6–10 pmol/L or 21–35 pg/mL) and five in the moderate–severe HF range (>10 pmol/L or >35 pg/mL, range of 39–119 pg/mL). Using the identical assay, sheep with post-myocardial infarction LV dysfunction have BNP levels of 10–12 pmol/L (~35–42 pg/mL).")[OAurl]([?–]?>)[OAurl][?–]?>17 and sheep with LVPO leading to pre-morbid (euthanasia) severe HF/death have BNP of 15.3 ± 3.6 pmol/L (~53 pg/mL).")[OAurl]([?–]?>)[OAurl][?–]?>18 Thus, levels of BNP measured in the pigs in this study were indicative of moderate to severe HF.

A highly novel aspect of this study is the HP$^{13}$CMR imaging. Advanced cardiometabolic imaging allows detection of novel structural and metabolic cardiac changes. HP$^{13}$CMR spectroscopy allows real-time study of myocardial energetics and substrate utilization to aid in understanding the pathophysiology of HFpEF and may identify novel therapeutic strategies. HP$^{13}$CMR results showed significant increases in hyperpolarized [1$^{13}$C]alanine and [1$^{13}$C]bicarbonate levels in the cardiac tissue of HFpEF pigs. Because these [1$^{13}$C]labeled metabolite signals of [1$^{13}$C]lactate, [1$^{13}$C]alanine, and [1$^{13}$C]bicarbonate are obtained by conversion from the hyperpolarized injected [1$^{13}$C]pyruvate via lactate dehydrogenase, ALT, and PDH, respectively, we deduced that ALT and PDH activity increased in the HFP EF pigs. The failing heart switches its energy source from fatty acids towards glucose,")[OAurl]([?–]?>)[OAurl][?–]?>18 directed towards glycolysis, whilst the change in glucose oxidation associated with PDH flux is related to the stage and pathogenesis of HF.")[OAurl]([?–]?>)[OAurl]([?–]?>)[OAurl][?–]?>19 In rats with aortic constriction, glucose oxidation is initially increased, but eventually decreases as cardiac systolic dysfunction develops.")[OAurl]([?–]?>)[OAurl][?–]?>20 This may explain the increase in PDH flux in our HFP EF pig model. Others have demonstrated that in 8 weeks, from pressure-overload LVH non-diabetic rats, more active PDH complex was extracted with healthy hearts.")[OAurl]([?–]?>)[OAurl][?–]?>21 Increased cardiac glucose oxidation is also observed in spontaneously hypertensive rats")[OAurl]([?–]?>)[OAurl][?–]?>22 and in dogs with pacing-induced HF.")[OAurl]([?–]?>)[OAurl][?–]?>23 Concomitantly, increased ALT activity may reflect increased supply of amino acids and nucleic acids for hypertrophic growth.")[OAurl]([?–]?>)[OAurl][?–]?>24 Taken together, the HP$^{13}$CMR results indicate our porcine HFpEF model displays metabolic changes consistent with HF.

The unmet need for models of HFpEF allowing both elucidation of the biology of this syndrome and providing a test bed for candidate novel therapies has triggered interest in developing large animal models of HFpEF. Efforts in this direction have had varying degrees of success. In addition to the progressive LVPO model,")[OAurl]([?–]?>)[OAurl][?–]?>6 and our previously described sheep model of acute aortic constriction, which induced rapid deterioration to HF/death.")[OAurl][?–]?>25 others have also employed aortic constriction in pigs. Acute onset aortic constriction beyond the carotid bifurcation resulted in rapid induction of LVH with reduced LVEF and acute HF by Day 7 (including death in 4/22 pigs), which then over 7 weeks evolved into a compensated phase of concentric LVH with restored cardiac function.")[OAurl](?)[OAurl][?–]?>25 Thus, there is need to induce LVPO in a progressive nature as implemented in this study to induce progressive LVH without systolic dysfunction. Other authors banded ascending aorta of minipigs (25–30 kg) but showed no significant LVH or change in left atrial volume or LVEDP over 20 weeks and did not report of plasma BNP. Hence this effort produced no objective evidence of HF.")[OAurl](?)[OAurl][?–]?>26 The Mayo group showed that renal wrapping of old dogs induced chronic hypertension with secondary LVH and fibrosis with impaired LV relaxation but no increase in LV diastolic stiffness claiming to mimic HFP EF.")[OAurl][?–]?>27 However, the requirement for old animals (aged 8–13) makes this model relatively impractical and expensive.

The traditional rodent deoxycorticosterone acetate (DOCA)–salt model has been upscaled to pigs, with variable results.")[OAurl][?–]?>28 DOCA-salt pigs demonstrated significant increases in LV mass and wall thickness and also left atrial volumes at rest compared with control pigs.")[OAurl][?–]?>28 During dobutamine stress, changes in LVEF and LVEVS demonstrated normal contractile reserve but there were differences in cardiac index and LVEDV compared to normal pigs indicating inadequate increase in myocardial perfusion reserve during dobutamine stress perhaps indicating early-stage HFpEF but there was no change in T1 mapping of fibrosis.")[OAurl][?–]?>28 When DOCA-induced hypertension is combined with high salt, fat, cholesterol, and sugar diet, pigs develop concentric LVH and left atrial dilation with no change in LVEF or HF symptoms at rest.")[OAurl][?–]?>29 While there was some evidence of pressure volume relationship being perturbed, LVEDP was not elevated unless pigs were challenged with pacing plus dobutamine. A recently reported study took an even more complex approach with multiple comorbidities modelled by combining streptozocin-induced diabetes, renal artery embolization to induce kidney dysfunction and hypertension, and a high fat/salt diet in six pigs followed for 6 months.")[OAurl][?–]?>30 They documented evidence of systemic inflammation, coronary endothelial dysfunction, and a rise in reactive oxygen species with disruption of nitric oxide along with myocardocyte stiffness and cardiac fibrosis. This model claimed to be a clinically relevant model LV diastolic dysfunction and a precursor for HFP EF, but no pigs demonstrated overt HF. Thus, although
several large animal models have been proposed, none of these fulfill all the features present in human disease. That being said, human HFpEF is a complex syndrome with multiple contributing aetiologies with varying occurrence and severity of comorbidities. Animal models invariably model some but not all aspects of pathology, thus, researchers need to select combinations of pathological features according to the specific aims of their study. Many of the models noted above replicate pre-clinical disease with no objective evidence of HF; thus, they represent early stages of the disease process and not HFpEF per se.

In conclusion, we have described a steadily progressive, but sufficiently stable and controllable (via titration of cuff/constriction) model of LVPO with novel data evidencing fibrosis based on MRI T1 MOLLI mapping, elevated filling pressures, metabolic changes, and progression to frank HF with prolongation of the model. This is a clinically relevant model of HFpEF on a predominant background of hypertension with clear utility in advancing knowledge of underlying pathophysiology of HFpEF and testing novel candidate therapies.

Acknowledgements

We would like to thank staff of Comparative Medicine, National University Singapore, particularly Denise Mathew, for expert assistance with animal care and procedures. We also thank Professor Mike Zile and Frank Spinale, University South Carolina for consultation and advice on set up of the pig model. We also thank Janise Lalic for help with her assistance with animal setup at the Singapore Bioimaging Consortium as well as Ong Sing Yee and Nurul Farhana Salleh for help with HP C13 preparation.

Conflict of interest

None declared.

Funding

Grant support was provided by the ATTRaCT SPF grant from Biomedical Research Council of Singapore.

References


4. Roh J, Houstis N, Rosenzweig A. Why don’t we have proven treatments for HFpEF. *Circ Res* 2017; 120: 1243–1245.


Porcine model of heart failure with preserved ejection fraction


