



Effect of Empagliflozin on Left Ventricular Volumes in Patients With Type 2 Diabetes, or Prediabetes, and Heart Failure With Reduced Ejection Fraction (SUGAR-DM-HF)

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BACKGROUND: Sodium-glucose cotransporter 2 inhibitors reduce the risk of heart failure hospitalization and cardiovascular death in patients with heart failure and reduced ejection fraction (HFrEF). However, their effects on cardiac structure and function in HFrEF are uncertain.

METHODS: We designed a multicenter, randomized, double-blind, placebo-controlled trial (the SUGAR-DM-HF trial [Studies of Empagliflozin and Its Cardiovascular, Renal and Metabolic Effects in Patients With Diabetes Mellitus, or Prediabetes, and Heart Failure]) to investigate the cardiac effects of empagliflozin in patients in New York Heart Association functional class II to IV with a left ventricular (LV) ejection fraction $\leq 40\%$ and type 2 diabetes or prediabetes. Patients were randomly assigned 1:1 to empagliflozin 10 mg once daily or placebo, stratified by age (<65 and ≥ 65 years) and glycemic status (diabetes or prediabetes). The coprimary outcomes were change from baseline to 36 weeks in LV end-systolic volume indexed to body surface area and LV global longitudinal strain both measured using cardiovascular magnetic resonance. Secondary efficacy outcomes included other cardiovascular magnetic resonance measures (LV end-diastolic volume index, LV ejection fraction), diuretic intensification, symptoms (Kansas City Cardiomyopathy Questionnaire Total Symptom Score, 6-minute walk distance, B-lines on lung ultrasound, and biomarkers (including N-terminal pro-B-type natriuretic peptide).

RESULTS: From April 2018 to August 2019, 105 patients were randomly assigned: mean age 68.7 (SD, 11.1) years, 77 (73.3%) male, 82 (78.1%) diabetes and 23 (21.9%) prediabetes, mean LV ejection fraction 32.5% (9.8%), and 81 (77.1%) New York Heart Association II and 24 (22.9%) New York Heart Association III. Patients received standard treatment for HFrEF. In comparison with placebo, empagliflozin reduced LV end-systolic volume index by 6.0 (95% CI, -10.8 to -1.2) mL/m² ($P=0.015$). There was no difference in LV global longitudinal strain. Empagliflozin reduced LV end-diastolic volume index by 8.2 (95% CI, -13.7 to -2.6) mL/m² ($P=0.0042$) and reduced N-terminal pro-B-type natriuretic peptide by 28% (2%–47%), $P=0.038$. There were no between-group differences in other cardiovascular magnetic resonance measures, diuretic intensification, Kansas City Cardiomyopathy Questionnaire Total Symptom Score, 6-minute walk distance, or B-lines.

CONCLUSIONS: The sodium-glucose cotransporter 2 inhibitor empagliflozin reduced LV volumes in patients with HFrEF and type 2 diabetes or prediabetes. Favorable reverse LV remodeling may be a mechanism by which sodium-glucose cotransporter 2 inhibitors reduce heart failure hospitalization and mortality in HFrEF.

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Key Words: clinical trial ■ diabetes mellitus ■ empagliflozin ■ heart failure ■ magnetic resonance imaging ■ myocardium ■ sodium-glucose transporter 2 inhibitors ■ ventricular remodeling

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Clinical Perspective

What Is New?

- Although sodium-glucose cotransporter 2 inhibitors improve clinical outcomes in patients with heart failure with reduced ejection fraction, their effects on left ventricular remodeling are uncertain.
- To our knowledge, this is the first adequately powered cardiovascular magnetic resonance trial using a sodium-glucose cotransporter 2 inhibitor in patients with heart failure with reduced ejection fraction.
- In a total of 105 patients with heart failure with reduced ejection fraction and type 2 diabetes, or prediabetes, randomly assigned to empagliflozin 10 mg once daily or matching placebo and treated for 36 weeks, empagliflozin led to significant reductions in left ventricular end-systolic and end-diastolic indexed volumes of 6.0 and 8.2 mL/m², respectively, and reductions in N-terminal pro-B-type natriuretic peptide by 28% (2%–47%), *P*=0.038.

What Are the Clinical Implications?

- Clinicians should be aware that the sodium-glucose cotransporter 2 inhibitor, empagliflozin, causes favorable cardiac remodeling in patients with heart failure with reduced ejection fraction.
- This may represent a mechanism by which sodium-glucose cotransporter 2 inhibitors reduce heart failure hospitalizations and cardiovascular death.

Adverse left ventricular remodeling, reflected by increased volumes and reduced contractility, is at the core of the pathophysiology of heart failure with reduced ejection fraction (HFrEF).^{1,2} Cardiac remodeling in HFrEF progresses over time, and larger left ventricular volumes are associated with worse non-fatal and fatal outcomes.^{3–10} Drug and device therapies that slow the progression of, or even reverse, remodeling are associated with better clinical outcomes in HFrEF. Sodium-glucose cotransporter 2 (SGLT2) inhibitors recently have been shown to reduce the risk of heart failure hospitalization and cardiovascular death in patients with HFrEF.^{11,12} The mechanism or mechanisms underlying these benefits are uncertain, including whether SGLT2 inhibitors have a favorable effect on cardiac remodeling. However, the reduction in NT-proBNP (N-terminal pro-B-type natriuretic peptide) observed with SGLT2 inhibitors is consistent with such an action.^{11,12} Empagliflozin has been shown to reduce left ventricular mass (indexed to body surface area) in people with type 2 diabetes and coronary artery disease, but the participants in this trial (the EMPA-HEART CardioLink-6 trial [Effects of Empagliflozin on Cardiac

Structure in Patients with Type 2 Diabetes]) had normal left ventricular volumes and ejection fraction, and the decrease in left ventricular mass likely reflected reduction in weight and blood pressure, effects not associated with improved outcomes or favorable left ventricular remodeling in HFrEF.¹³ Only 1 prior trial has examined the effect of a SGLT2 inhibitor on left ventricular remodeling specifically in patients with HFrEF. The participants had relatively modest left ventricular enlargement and the trial was underpowered, with only 28 patients per treatment group.¹⁴ Consequently, we designed a larger, multicenter randomized, double-blind, placebo-controlled trial powered to investigate the effects of SGLT2 inhibition on left ventricular volumes in patients with HFrEF and type 2 diabetes or prediabetes.

METHODS

The data that support the findings of this study may be available from the corresponding author on reasonable request.

Fifteen hospitals in Scotland took part in the trial (details in the [Data Supplement](#)), each of which had ethical committee approval, and all participants gave written informed consent.

Patients

Patients ≥18 years of age with HFrEF and type 2 diabetes (documented history of diabetes or previously undiagnosed diabetes with glycohemoglobin ≥48 mmol/mol [≥6.5%]), or prediabetes (glycohemoglobin 39–47 mmol/mol [5.7%–6.4%]) were potentially eligible. Participants had to be in New York Heart Association (NYHA) functional class II to IV and have a left ventricular ejection fraction (LVEF) ≤40%. The dose of angiotensin-converting enzyme inhibitor, angiotensin receptor blocker, or sacubitril/valsartan and β-blocker had to be unchanged for at least 4 weeks. Patients with type 2 diabetes were required to have a glycohemoglobin ≤97 mmol/mol (≤11%) and, if they were treated with glucose-lowering therapy, the treatment dose had to be unchanged for at least 6 weeks. The key exclusion criteria were type 1 diabetes, estimated glomerular filtration rate <30 mL/min/1.73m², and history of diabetic ketoacidosis (a complete list of inclusion and exclusion criteria are provided in the [Data Supplement](#)).

Randomization and Stratification

Patients were randomly assigned 1:1 to either empagliflozin 10 mg once daily or matching placebo by using an interactive web response system. The randomization sequence was performed in blocks of 4 and stratified by (1) age (<65 years, ≥65 years) and (2) diabetes or prediabetes status.

Schedule of Study Visits

Patients attended for 6 visits: screening, baseline, weeks 2, 12, 36, and 40 ([Table I in the Data Supplement](#)).

Outcomes

Copriary Outcomes

The copriary outcomes were (1) left ventricular end-systolic volume indexed (LVESVi) to body surface area and (2) left

ventricular global longitudinal strain (LV GLS). Between treatment group differences in the change from baseline to 36 weeks were calculated for both outcomes.

Secondary Outcomes

The secondary end points assessed at 36 weeks included additional cardiovascular magnetic resonance (CMR) measurements (body surface area–indexed left ventricular end-diastolic volume [LVEDVi], LVEF, body surface area–indexed left ventricular [LV] mass index, LV global function index, body surface area–indexed left atrial volume, myocardial blood flow, and extracellular volume fraction), diuretic intensification, assessment of symptoms using the Kansas City Cardiomyopathy Questionnaire Total Symptom Score (KCCQ-TSS), 6-minute walk distance, number of B-lines on lung ultrasound (a marker of lung congestion), and a range of biomarkers (as described in Biomarker Assessments and in the [Data Supplement](#)).^{15–17}

CMR Acquisition and Analysis

ECG-gated CMR was performed at 1 center (Queen Elizabeth University Hospital) at baseline prerandomization and week 36 using a 3.0 Tesla scanner (MAGNETOM Prisma, Siemens Healthcare). A week 40 scan after withdrawal of treatment was also planned but could not be performed in many patients because of the coronavirus disease 2019 (COVID-19) pandemic. The imaging protocol ([Table II in the Data Supplement](#)) included cine imaging, native T1 mapping (modified Look-Locker inversion-recovery), and delayed enhancement sequences. All scan acquisitions were spatially coregistered. CMR analysis methods are detailed in the [Data Supplement](#).

Biomarker Assessments

Glycohemoglobin, estimated glomerular filtration rate, uric acid, high-sensitivity cardiac troponin I, and galectin-3 (i1000SR, Abbott Laboratories, Abbott Diagnostics) were measured, along with NT-proBNP and growth differentiation factor-15 (e411, Roche Diagnostics) at baseline, 12 weeks, and 36 weeks. All of these assays were performed by using the manufacturers' calibrators and quality controls.

Statistical Analysis

Statistical analyses were conducted at the study data center (Robertson Center for Biostatistics, University of Glasgow) according to a prespecified Statistical Analysis Plan. All analyses were performed according to the intention-to-treat principle, including all randomly assigned participants with postrandomization data available for the outcome of interest at any given time point, irrespective of their subsequent participation in the study and their adherence to randomized treatment.

For all outcomes, analysis of covariance, with adjustment for randomized group, baseline value, and the stratification variables (age and diabetes/prediabetes status) were used to determine the between-group differences in the outcomes at 36 weeks. For the coprimary end points of LVESVi and LV GLS, the α was split so that a *P* value of 0.025 was considered statistically significant. No interim analyses were performed. For all secondary outcomes, a *P* value of 0.05 was considered statistically significant. All analyses were conducted using SAS version 9.4 (SAS Inc).

RESULTS

Recruitment took place between April 2018 and August 2019; follow-up visits were completed in May 2020. Of 166 patients screened, 105 were randomly assigned (52 to empagliflozin and 53 to placebo). Of these, 82 patients (78.1%) had diabetes and 23 (21.9%) had prediabetes.

Baseline Characteristics

The mean (SD) age of participants was 68.7 (11.1) years, 77 patients (73.3%) were male, and 81 (77.1%) patients were in NYHA functional class II and 24 (22.9%) in NYHA class III. The median (interquartile range) duration of heart failure was 2.1 (1.0–4.8) years, and 52 patients (49.5%) had a history of heart failure hospitalization. Most patients had coronary artery disease (74 patients, 70.5%) and received standard heart failure therapy (Table 1). The mean (SD) CMR LVEF was 32.5% (9.8%), and median (interquartile range) NT-proBNP was 466 (177–1120) pg/mL.

Diabetes treatments and other medical history are shown in Table 1.

Completeness of Follow-Up and Adherence

Of the 52 patients randomly assigned to empagliflozin, 42 remained on randomized therapy and had complete coprimary end point data at baseline and week 36 ([Figure I in the Data Supplement](#)). Of the 53 patients randomly assigned to placebo, 50 remained on randomized therapy and had complete coprimary end point data at baseline and week 36. There were 2 deaths in the empagliflozin group, one attributable to newly diagnosed pancreatic cancer and one attributable to cardiogenic shock. There were no deaths in the placebo group.

Coprimary Outcomes

LVESVi decreased by 7.9 mL/m² between baseline and 36 weeks in the empagliflozin group in comparison with a reduction of 1.5 mL/m² in the placebo group: adjusted between-group difference –6.0 (95% CI, –10.8 to –1.2) mL/m²; *P*=0.015 (Table 2 and Figure 1). There was no difference in LV GLS between the empagliflozin and placebo groups: adjusted between-group difference 0.35% (95% CI, –0.25% to 0.95%); *P*=0.25. In view of potential imbalances in sex, NYHA class, and history of heart failure hospitalization at baseline, we did an additional post hoc adjustment for these variables that did not change our findings: adjusted between-group difference in LVESVi –5.9 (95% CI –11.0 to –0.9) mL/m²; *P*=0.023 and adjusted between-group difference in LV GLS 0.45% (95% CI, –0.17% to 1.08%); *P*=0.15.

Table 1. Baseline Characteristics of Randomized Patients

Characteristic	All	Empagliflozin	Placebo
	(n=105)	(n=52)	(n=53)
Age, mean (SD), y	68.7 (11.1)	68.2 (11.7)	69.2 (10.6)
Male, n (%)	77 (73.3)	34 (65.4)	43 (81.1)
Body mass index, mean (SD), kg/m ²	30.7 (5.5)	30.9 (5.9)	30.4 (5.1)
Systolic blood pressure, mean (SD), mm Hg	128.0 (20.0)	125.8 (18.2)	130.3 (21.6)
Heart rate, mean (SD), beats/min	66.4 (11.5)	68.0 (12.9)	64.8 (9.8)
Glycemic status			
Type 2 diabetes, * n (%)	82 (78.1)	40 (76.9)	42 (79.2)
Prediabetes, n (%)	23 (21.9)	12 (23.1)	11 (20.8)
Glycohemoglobin, mean (SD), %	7.2 (1.5)	7.5 (1.6)	7.0 (1.4)
Heart failure status			
Cardiovascular magnetic resonance left ventricular ejection fraction, mean (SD), %	32.5 (9.8)	32.1 (10.3)	32.9 (9.3)
Duration of heart failure, median (IQR), y	2.1 (1.0–4.8)	2.3 (1.0–5.3)	2.0 (1.0–4.1)
Prior heart failure hospitalization, n (%)	52 (49.5)	21 (40.4)	31 (58.5)
N-terminal pro-B-type natriuretic peptide, median (IQR), pg/mL	466 (177–1120)	475 (198–1117)	466 (156–1148)
New York Heart Association Class II, n (%)	81 (77.1)	37 (71.2)	44 (83.0)
New York Heart Association Class III, n (%)	24 (22.9)	15 (28.8)	9 (17.0)
New York Heart Association Class IV, n (%)	0 (0.0)	0 (0.0)	0 (0.0)
Other medical history			
Coronary artery disease, n (%)	74 (70.5)	36 (69.2)	38 (71.7)
Prior myocardial infarction, † n (%)	62 (83.8)	27 (75.0)	35 (92.1)
Percutaneous coronary intervention, † n (%)	42 (56.8)	17 (47.2)	25 (65.8)
Coronary artery bypass graft, † n (%)	18 (24.3)	10 (27.8)	8 (21.1)
Hypertension, n (%)	74 (70.5)	39 (75.0)	35 (66.0)
Stroke, n (%)	11 (10.5)	3 (5.8)	8 (15.1)
Creatinine, mean (SD), μmol/L	100.2 (31.8)	96.4 (33.4)	104.0 (30.1)
Estimated glomerular filtration rate—Chronic Kidney Disease Epidemiology Collaboration, mean (SD), mL/min/1.73m ²	67.3 (22.0)	69.5 (22.3)	65.1 (21.8)
Hemoglobin, mean (SD), g/L	131.8 (13.5)	131.0 (12.3)	132.7 (14.7)
Heart failure therapy			
Angiotensin-converting enzyme inhibitor/angiotensin receptor blocker/angiotensin-receptor neprilysin inhibitor, n (%)	100 (95.2)	49 (94.2)	51 (96.2)
Angiotensin-converting enzyme inhibitor, n (%)	49 (46.7)	25 (48.1)	24 (45.3)
Angiotensin receptor blocker, n (%)	15 (14.3)	3 (5.8)	12 (22.6)
Angiotensin-receptor neprilysin inhibitor, n (%)	36 (34.3)	21 (40.4)	15 (28.3)
β-Blocker, n (%)	96 (91.4)	46 (88.5)	50 (94.3)
Mineralocorticoid receptor antagonist, n (%)	63 (60.0)	32 (61.5)	31 (58.5)
Loop diuretic, n (%)	60 (57.1)	31 (59.6)	29 (54.7)
Digoxin, n (%)	5 (4.8)	4 (7.7)	1 (1.9)
Ivabradine, n (%)	11 (10.5)	8 (15.4)	3 (5.7)
Type 2 diabetes			
Duration of diabetes, median (IQR), y	9.0 (3.2–14.8)	8.2 (3.6–13.6)	10.0 (3.2–15.6)
Diabetes therapy, n (%)	61 (74.4)	31 (77.5)	30 (71.4)
Biguanide (metformin), n (%)	46 (75.4)	25 (80.6)	21 (70.0)
Sulfonylurea, n (%)	25 (41.0)	8 (25.8)	17 (56.7)
Insulin, n (%)	9 (14.8)	6 (19.4)	3 (10.0)
Dipeptidyl dipeptidase-4 inhibitors, n (%)	22 (36.1)	13 (41.9)	9 (30.0)
Glucagon-like peptide-1 receptor agonist, n (%)	5 (8.2)	3 (9.7)	2 (6.7)

*One patient had newly diagnosed diabetes at screening.

†Values are n (%) for prior myocardial infarction, percutaneous coronary intervention, and coronary artery bypass graft are for subset of those who have coronary artery disease.

Table 2. Change in CMR Parameters with Empagliflozin 10 mg/d or Placebo From Baseline to Week 36

Variable*	Empagliflozin				Placebo				Between-group difference (95% CI)†	P value
	n	Baseline	Week 36	Change	n	Baseline	Week 36	Change		
Coprimary CMR outcomes										
LV end-systolic volume index, ‡ mL/m ²	42	80.8 (37.2)	72.9 (37.0)	−7.9 (11.8)	50	76.6 (29.3)	75.2 (29.2)	−1.5 (11.3)	−6.0 (−10.8 to −1.2)	0.015
LV global longitudinal strain, § %	42	−7.04 (2.11)	−7.09 (2.11)	−0.05 (1.57)	50	−7.79 (2.54)	−7.97 (2.31)	−0.18 (1.49)	0.35 (−0.25 to 0.95)	0.25
Secondary CMR outcomes										
LV end-diastolic volume index, ‡ mL/m ²	42	114.7 (37.0)	105.7 (37.6)	−9.0 (12.4)	50	111.4 (29.2)	110.9 (28.3)	−0.4 (14.1)	−8.2 (−13.7 to −2.6)	0.004
LV ejection fraction, %	42	31.7 (9.9)	33.5 (10.3)	1.8 (5.7)	50	33.0 (9.5)	34.2 (9.7)	1.2 (3.8)	0.3 (−1.7 to 2.3)	0.75
LV mass index, ‡ g/m ²	42	61.2 (16.1)	58.6 (16.2)	−2.7 (6.1)	50	65.4 (19.6)	64.1 (18.3)	−1.3 (7.3)	−1.9 (−4.7 to 0.8)	0.17
LV global function index, %	42	23.4 (7.7)	24.4 (7.9)	1.0 (3.8)	50	23.6 (7.4)	24.8 (7.5)	1.1 (3.3)	−0.2 (−1.7 to 1.2)	0.76
Left atrial volume index, ‡ mL/m ²	42	40.5 (13.3)	38.6 (13.5)	−1.9 (8.5)	50	43.7 (12.5)	43.0 (11.9)	−0.7 (12.1)	−2.4 (−6.5 to 1.8)	0.26
Myocardial blood flow, mL/g/min	32	0.80 (0.18)	0.81 (0.24)	0.01 (0.22)	37	0.85 (0.24)	0.93 (0.30)	0.08 (0.27)	−0.08 (−0.20 to 0.04)	0.17
Extracellular volume fraction, %	32	31.8 (4.5)	31.0 (4.7)	−0.8 (3.5)	36	31.6 (4.8)	31.0 (5.1)	−0.7 (3.5)	0.004 (−1.7 to 1.7)	1.00
Exploratory CMR outcomes										
LV end-systolic volume, mL	42	157.5 (68.1)	142.3 (70.9)	−15.1 (24.0)	50	152.9 (58.4)	150.1 (57.7)	−2.8 (23.7)	−11.9 (−21.9 to −1.9)	0.021
LV end-diastolic volume, mL	42	224.8 (72.2)	207.5 (75.3)	−17.3 (24.8)	50	222.7 (60.1)	222.1 (59.3)	−0.6 (29.2)	−16.4 (−27.8 to −5.0)	0.005
LV mass, g	42	121.2 (36.5)	116.1 (37.1)	−5.1 (12.7)	50	131.9 (44.9)	129.5 (42.9)	−2.5 (14.8)	−3.8 (−9.6 to 1.9)	0.19
Left atrial volume, mL	42	79.0 (24.3)	75.5 (26.3)	−3.5 (17.2)	50	87.9 (27.5)	86.3 (25.6)	−1.5 (24.1)	−5.1 (−13.4 to 3.2)	0.22

CMR indicates cardiovascular magnetic resonance; and LV, left ventricular.

*Mean (SD).

†Treatment effect calculated using an analysis of covariance model adjusted for the treatment group, age at baseline, diabetes status, and baseline value of the outcome.

‡Indexed to baseline body surface area.

§Feature tracking.

||Myocardial blood flow corrected for rate-pressure product.

Secondary Outcomes

Cardiovascular Magnetic Resonance

LVEDVi decreased by 9.0 mL/m² between baseline and 36 weeks in the empagliflozin group in comparison with a reduction of 0.4 mL/m² in the placebo group: adjusted between-group difference −8.2 (−13.7 to −2.6) mL/m²; $P=0.004$. There were no significant changes in the other CMR variables (LVEF, LV mass index, LV global function index, indexed left atrial volume, myocardial blood flow, or extracellular volume fraction) after 36 weeks of treatment with empagliflozin, in comparison with placebo (Table 2 and Figure 2).

Diuretic Intensification, KCCQ-TSS, 6-Minute Walk Distance, and B Lines

There were no between-group differences in the intensification of diuretic therapy, change in KCCQ-TSS,

6-minute walk distance, or total number of B lines, between baseline and 36 weeks (Table 3).

Biomarkers

In comparison with placebo, empagliflozin reduced serum uric acid ($P<0.0001$) and NT-proBNP ($P=0.038$), and increased galectin-3 ($P=0.013$) and hematocrit ($P<0.0001$). There were no between-group differences in other biomarkers (Table 3).

Safety and Exploratory Outcomes

There were no between-group differences in safety outcomes (Table III in the Data Supplement). Exploratory outcomes (blood pressure, heart rate, body weight, and blood ketones) are presented in Table IV in the Data Supplement.

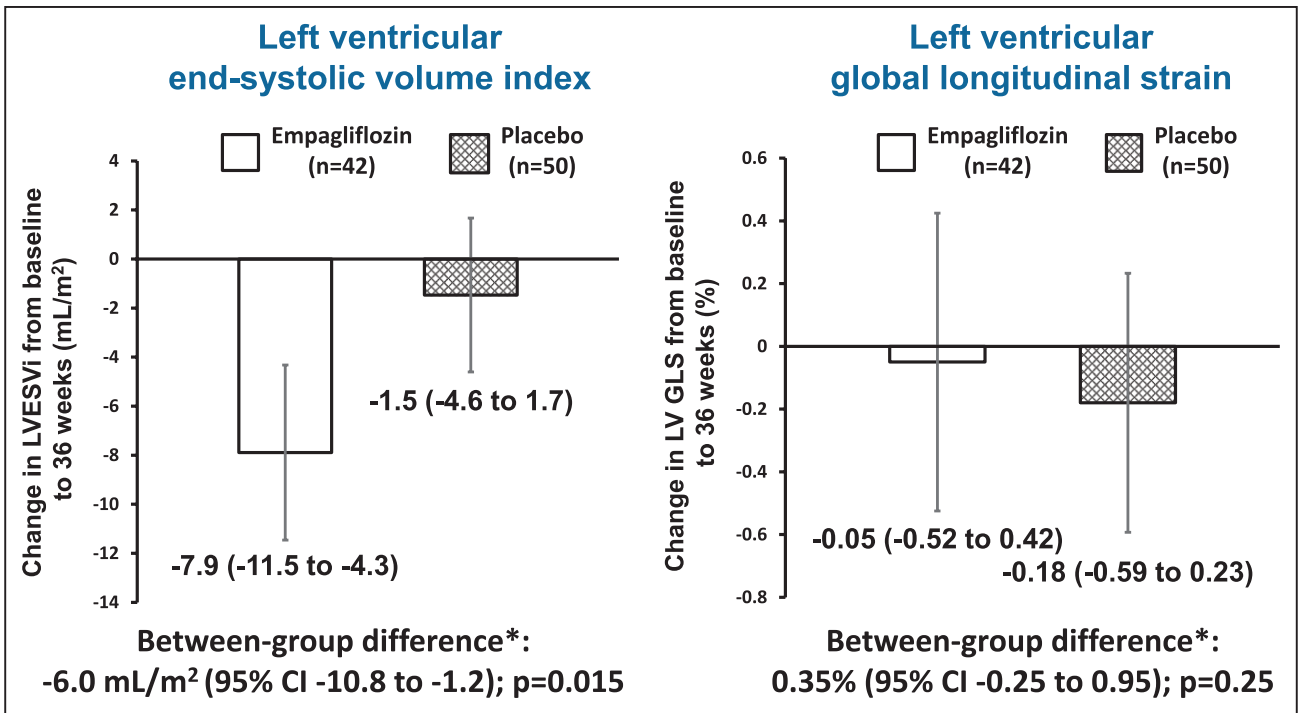


Figure 1. Change in coprimary cardiovascular magnetic resonance outcomes from baseline to week 36. Mean (95% CI). *Treatment effect calculated using an analysis of covariance model adjusted for treatment group, age at baseline, diabetes status, and baseline value. LVESVi indicates left ventricular end-systolic volume index; and LV GLS, left ventricular global longitudinal strain.

DISCUSSION

We found that treatment of patients with HFrEF for 36 weeks with the SGLT2 inhibitor empagliflozin led to a

significant reduction in LV volumes (LVESVi and LVEDVi) in comparison with placebo, but no improvement in LV GLS. Empagliflozin also reduced NT-proBNP. These favorable changes with empagliflozin were observed

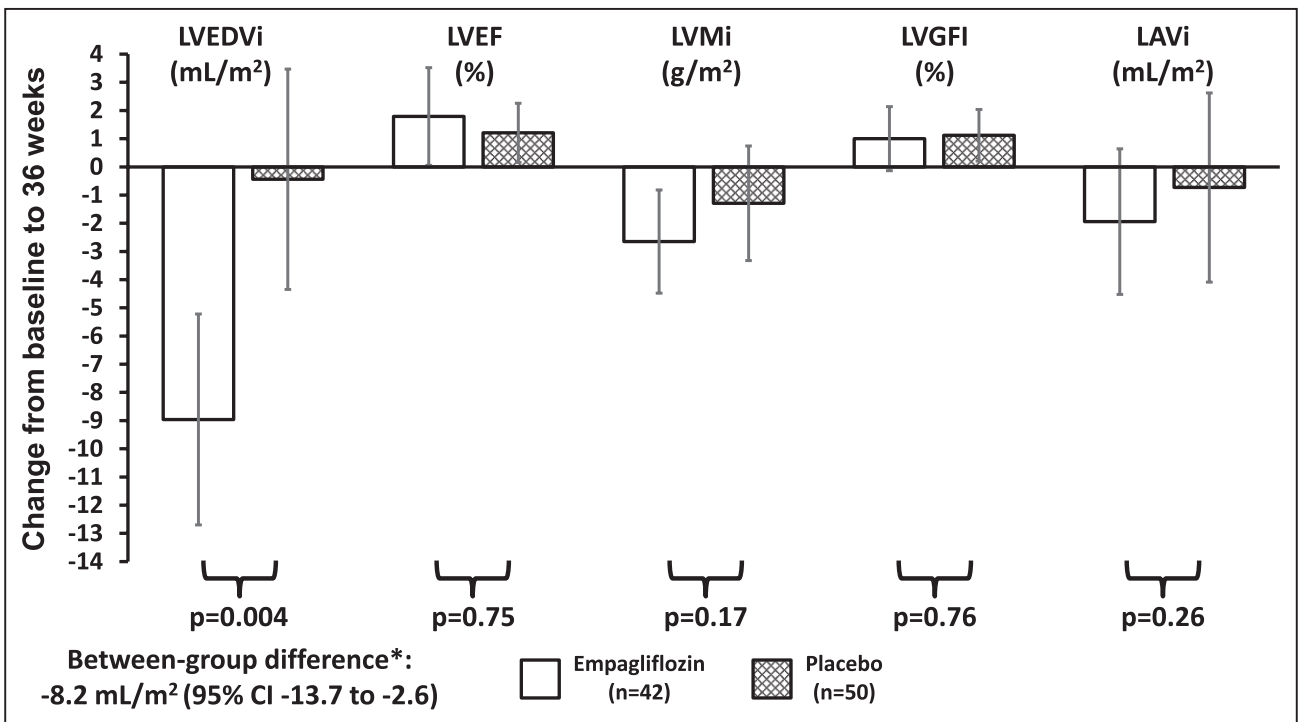


Figure 2. Change in secondary cardiovascular magnetic resonance outcomes from baseline to week 36. *Treatment effect calculated using an analysis of covariance model adjusted for treatment group, age at baseline, diabetes status, and baseline value. LAVi indicates left atrial volume index; LVEDVi, left ventricular end-diastolic volume index; LVEF, left ventricular ejection fraction; LVGFI, left ventricular global function index; and LVMI, left ventricular mass index.

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Table 3. Change in Secondary Outcomes with Empagliflozin 10 mg/d or Placebo from Baseline to Week 36

Secondary outcome variables	Empagliflozin				Placebo				Between-group difference (95% CI)*	P value
	n	Baseline	Week 36	Change	n	Baseline	Week 36	Change		
Diuretic intensification†	28	–	3 (10.7)	3 (10.7)	29	–	4 (13.8)	4 (13.8)	–	1.00
Kansas City Cardiomyopathy Questionnaire Total Symptom Score‡	44	73.4 (22.0)	74.1 (22.9)	0.7 (17.5)	51	74.7 (19.5)	78.9 (21.6)	4.2 (13.8)	–4.0 (–10.2 to 2.1)	0.19
6-minute walk distance,‡ meters	35	360.2 (105.8)	385.7 (86.9)	25.4 (60.5)	42	353.5 (93.8)	387.1 (109.9)	33.6 (50.7)	–9.9 (–34.4 to 14.7)	0.43
Total B lines‡§	44	5.86 (5.00)	7.14 (5.81)	1.27 (4.06)	49	6.55 (6.32)	7.71 (5.13)	1.16 (4.42)	–0.14 (–1.70 to 1.43)	0.86
Glycohemoglobin,‡ %	44	7.37 (1.55)	7.14 (1.29)	–0.23 (1.08)	50	6.99 (1.39)	7.29 (1.66)	0.30 (1.21)	–0.36 (–0.80 to 0.09)	0.11
Estimated glomerular filtration rate–Chronic Kidney Disease Epidemiology Collaboration,‡ mL/min/1.73 m ²	44	68.3 (22.4)	67.7 (23.0)	–0.54 (9.0)	50	64.1 (22.0)	62.4 (23.9)	–1.7 (8.8)	1.4 (–2.2 to 5.0)	0.44
Uric acid,‡ μmol/L	44	391.6 (132.5)	323.9 (113.9)	–67.7 (81.9)	50	405.8 (106.3)	402.8 (109.0)	–3.08 (56.0)	–66.2 (–91.9 to –40.5)	<0.0001
N-terminal pro-B-type natriuretic peptide,‡ pg/mL	44	1236 (2342)	970 (1918)	–266 (955)	50	1148 (1905)	1373 (3317)	224 (2247)	–28% (–47% to –2%)	0.038
High-sensitivity cardiac troponin I,‡ pg/mL	44	11.4 (19.9)	10.6 (14.8)	–0.8 (9.6)	50	11.0 (20.6)	10.2 (12.0)	–0.7 (12.4)	–1% (–19% to 21%)	0.94
Growth differentiation factor-15,‡ pg/mL	44	2819 (2002)	2908 (2116)	89 (1295)	50	2949 (2046)	2899 (2017)	–50 (1173)	1% (–10% to 15%)	0.82
Galectin-3,‡ ng/mL	44	17.8 (6.2)	19.1 (6.3)	1.3 (3.6)	50	16.6 (5.9)	16.4 (6.1)	–0.2 (4.3)	1.94 (0.41 to 3.47)	0.013
Hematocrit,‡ L/L	44	0.39 (0.04)	0.43 (0.04)	0.03 (0.03)	50	0.40 (0.04)	0.40 (0.04)	0.01 (0.03)	0.027 (0.015 to 0.038)	<0.0001

*Treatment effect calculated using an analysis of covariance model adjusted for the treatment group, age at baseline, diabetes status, and baseline value of the outcome. Between-group differences are reported as mean change (95% CI) for glycohemoglobin, estimated glomerular filtration rate Chronic Kidney Disease Epidemiology Collaboration, uric acid, galectin-3, and hematocrit. Between-group differences are reported as % change (95% CI) for N-terminal pro-B-type natriuretic peptide, high-sensitivity cardiac troponin I, and growth differentiation factor-15.

†Values are n (%). Total N is of those that provided a diuretic dose. Diuretic intensification was defined as either an increase in diuretic dose, or initiation of diuretic therapy. Bumetanide 1 mg was considered equivalent to furosemide 80 mg orally.

‡Mean (SD).

§Eight-zone lung ultrasound. One subject is withdrawn from lung ultrasound analyses because of having known pulmonary fibrosis.

||The follow-up and baseline values of the outcome that are used in the analysis model have been log-transformed and the treatment effect estimate is reported as the relative difference (% difference) with 95% CI and P value.

despite excellent conventional heart failure therapy, including sacubitril/valsartan in more than one-third of patients, and may explain, at least in part, the improvement in clinical outcomes observed with SGLT2 inhibitors in HFrEF. We did not observe an improvement in symptoms (using the KCCQ-TSS) or functional capacity (6-minute walk distance), although we had limited power to show a change in either of these measures. B-line number on lung ultrasound did not differ between treatment groups, and there was no difference in diuretic use, suggesting no major difference in surrogates of congestion between treatment groups.

Deleterious LV remodeling, including increases in end-diastolic and end-systolic volume and reduction in LVEF, is pathognomonic of HFrEF, and the extent of adverse remodeling correlates with risk of hospitalization and death.^{1,2,18} LV volumes and contractility worsen progressively over time, and slowing of this progression,

or even reversal of adverse remodeling, is believed to be a key mechanism through which several pharmacological therapies improve clinical outcomes in HFrEF, as does cardiac resynchronization therapy. The magnitude of change in LV volumes observed at 36 weeks with empagliflozin compares favorably with the effects of other beneficial therapies in HFrEF and was incremental to those effects as our patients were comprehensively treated with renin-angiotensin system blockers and β -blockers. The more recently introduced therapies ivabradine and sacubitril/valsartan also reduced LV volumes when added to renin-angiotensin system blockers and β -blockers. In comparison with placebo, ivabradine reduced LVESVi by 5.8 mL/m² and LVEDVi by 5.5 mL/m² over 8 months in a substudy of SHIFT (Systolic Heart Failure Treatment With the I_f Inhibitor Ivabradine Trial).⁵ There have been 2 randomized trials of sacubitril/valsartan and remodeling. In the EVALUATE-HF trial (Effect

of Sacubitril-Valsartan versus Enalapril on Aortic Stiffness in Patients With Heart Failure and Reduced Ejection Fraction), sacubitril/valsartan, in comparison with enalapril, reduced LVESVi by 1.6 mL/m² and LVEDVi by 2.0 mL/m² over 12 weeks.⁶ In the PRIME trial (Pharmacological Reduction of Functional, Ischemic Mitral Regurgitation), patients with an ejection fraction between 25% and <50% and mitral regurgitation were randomly assigned to sacubitril/valsartan or valsartan.¹⁹ The addition of neprilysin inhibition did not result in a reduction in LVESVi, but a reduction in LVEDVi (7 mL/m²) was observed. In SHIFT, EVALUATE-HF, and PRIME, LV remodeling was evaluated by using transthoracic echocardiography, but in the present study, we used the gold standard method of assessing cardiac structure and function with CMR.

We know of only 1 other trial examining the effect of a SGLT2 inhibitor on LV remodeling in patients with HFrEF. That trial, REFORM (Research Into the Effect of SGLT2 Inhibition on Left Ventricular Remodeling in Patients With Heart Failure and Diabetes Mellitus), did not show any improvement in LV remodeling with dapagliflozin.¹⁴ The likely explanations for the difference between the trials are that patients in REFORM had a higher LVEF (45.5% versus 32.5%), better NYHA class distribution (45% versus 0% NYHA class I), less LV enlargement (LVESVi 52 versus 77 mL/m²), and, especially, the much smaller number of randomly assigned (56 versus 105 patients), resulted in limited power to detect a difference between treatments.

A key remaining question is how does SGLT2 inhibition lead to favorable reverse remodeling? Empagliflozin may have reduced LV afterload, although the decreases in systolic and diastolic blood pressure were small and there was no significant reduction in LV mass, which might have been anticipated if there was an important and sustained reduction in afterload. Alternatively, SGLT2 inhibitors may cause a reduction in preload by inducing a diuresis; the consequent reduction in LV stretch may lead to a reduction in LV volumes. In the present study and in others, SGLT2 inhibitors have been shown to reduce levels of NT-proBNP, which is a surrogate marker for the degree of LV wall stress.²⁰ The extent to which SGLT2 inhibitors exert a diuretic effect is debated, however, and we found no evidence of change in markers of pulmonary decongestion on lung ultrasound, no difference in conventional diuretic therapy, and no significant reduction in indexed left atrial volume. Alternatively, several additional mechanistic benefits of SGLT2 inhibitors on the myocardium have been proposed and these might also explain reduced LV volumes. A switch in myocardial energetics has been proposed but not proven in patients with HFrEF. SGLT2 inhibitors have also been proposed to reduce cardiac oxidative stress and inflammation through promotion of the actions of sirtuin-1 and upregulation

of hypoxia-inducible factors signaling, but such hypotheses require confirmation.²¹ SGLT2 inhibitors do not appear to exert their actions by altering myocardial blood flow or extracellular volume, because neither was changed in the present trial.

We did not observe improvements in LV GLS with SGLT2 inhibition. Our patients had LV GLS values considerably below normal and consistent with those described in other studies in patients with HFrEF.^{6,22} As a relatively new measure of myocardial systolic function, the only other trial of LV remodeling in HFrEF to investigate LV GLS was the EVALUATE-HF trial.⁶ In comparison with enalapril, sacubitril/valsartan had no effect on LV GLS in that trial. Strain has not been reported in the older trials of LV remodeling, so it is unclear whether LV GLS is expected to change in parallel with LV volumes. We did not observe an increase in LVEF with empagliflozin treatment. In general, increases in LVEF have paralleled reductions in LV volumes in prior trials with β -blockers and ivabradine.³⁻⁵ It is unclear why we did not see an increase in LVEF, although this was not observed in the trials with sacubitril/valsartan mentioned earlier.^{6,19}

Limitations

We did not include patients with atrial fibrillation or patients with cardiac devices to avoid image degradation. We did not recruit any patients with NYHA functional class IV, although such patients were eligible. Treatment was given for only 36 weeks, and favorable remodeling in response to pharmacological therapy may continue over a longer period.¹ Finally, in any modest-sized study such as this, imbalances in baseline characteristics can occur and potentially influence interpretation of the effects of randomized trials. To address this, we prespecified that we would adjust for the baseline value of the variable of interest, and factors used to stratify the randomization (age and diabetes status), as recommended in most guidelines for the analysis of clinical trials.

Conclusion

In summary, treatment with the SGLT2 inhibitor empagliflozin led to favorable reverse LV remodeling in patients with HFrEF and type 2 diabetes or prediabetes. This finding may, at least in part, explain the beneficial effect of SGLT2 inhibitors on clinical outcomes in HFrEF.

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Supplemental Materials

Data Supplement Methods
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REFERENCES

- Kramer DG, Trikalinos TA, Kent DM, Antonopoulos GV, Konstam MA, Udelson JE. Quantitative evaluation of drug or device effects on ventricular remodeling as predictors of therapeutic effects on mortality in patients with heart failure and reduced ejection fraction: a meta-analytic approach. *J Am Coll Cardiol*. 2010;56:392–406. doi: 10.1016/j.jacc.2010.05.011
- Cohn JN, Ferrari R, Sharpe N. Cardiac remodeling—concepts and clinical implications: a consensus paper from an international forum on cardiac remodeling. Behalf of an International Forum on Cardiac Remodeling. *J Am Coll Cardiol*. 2000;35:569–582. doi: 10.1016/s0735-1097(99)00630-0
- Bellenger NG, Rajappan K, Rahman SL, Lahiri A, Raval U, Webster J, Murray GD, Coats AJ, Cleland JG, Pennell DJ; CHRISTMAS Study Steering Committee and Investigators. Effects of carvedilol on left ventricular remodelling in chronic stable heart failure: a cardiovascular magnetic resonance study. *Heart*. 2004;90:760–764. doi: 10.1136/hrt.2003.015552
- Groenning BA, Nilsson JC, Sondergaard L, Fritz-Hansen T, Larsson HB, Hildebrandt PR. Antiremodeling effects on the left ventricle during beta-blockade with metoprolol in the treatment of chronic heart failure. *J Am Coll Cardiol*. 2000;36:2072–2080. doi: 10.1016/s0735-1097(00)01006-8
- Tardif JC, O'Meara E, Komajda M, Böhm M, Borer JS, Ford I, Tavazzi L, Swedberg K; SHIFT Investigators. Effects of selective heart rate reduction with ivabradine on left ventricular remodelling and function: results from the SHIFT echocardiography substudy. *Eur Heart J*. 2011;32:2507–2515. doi: 10.1093/eurheartj/ehr311
- Desai AS, Solomon SD, Shah AM, Claggett BL, Fang JC, Izzo J, McCague K, Abbas CA, Rocha R, Mitchell GF, EVALUATE-HF Investigators. Effect of sacubitril-valsartan vs enalapril on aortic stiffness in patients with heart failure and reduced ejection fraction: a randomized clinical trial. *JAMA*. 2019;322:1–10.
- Wong M, Staszewsky L, Latini R, Barlera S, Volpi A, Chiang YT, Benza RL, Gottlieb SO, Kleemann TD, Rosconi F, et al; Val-HeFT Heart Failure Trial Investigators. Valsartan benefits left ventricular structure and function in heart failure: Val-HeFT echocardiographic study. *J Am Coll Cardiol*. 2002;40:970–975. doi: 10.1016/s0735-1097(02)02063-6
- Greenberg B, Quinones MA, Koilpillai C, Limacher M, Shindler D, Benedict C, Shelton B. Effects of long-term enalapril therapy on cardiac structure and function in patients with left ventricular dysfunction. Results of the SOLVD echocardiography substudy. *Circulation*. 1995;91:2573–2581. doi: 10.1161/01.cir.91.10.2573
- Colucci WS, Koliass TJ, Adams KF, Armstrong WF, Ghali JK, Gottlieb SS, Greenberg B, Klibaner MI, Kukin ML, Sugg JE; REVERT Study Group. Metoprolol reverses left ventricular remodeling in patients with asymptomatic systolic dysfunction: the REversal of VEntricular

- Remodeling with Toprol-XL (REVERT) trial. *Circulation*. 2007;116:49–56. doi: 10.1161/CIRCULATIONAHA.106.666016
10. Cleland JG, Daubert JC, Erdmann E, Freemantle N, Gras D, Kappenberger L, Tavazzi L; Cardiac Resynchronization-Heart Failure (CARE-HF) Study Investigators. The effect of cardiac resynchronization on morbidity and mortality in heart failure. *N Engl J Med*. 2005;352:1539–1549. doi: 10.1056/NEJMoa050496
 11. McMurray JJV, Solomon SD, Inzucchi SE, Køber L, Kosiborod MN, Martinez FA, Ponikowski P, Sabatine MS, Anand IS, Bøhlhåvek J, et al; DAPA-HF Trial Committees and Investigators. Dapagliflozin in patients with heart failure and reduced ejection fraction. *N Engl J Med*. 2019;381:1995–2008. doi: 10.1056/NEJMoa1911303
 12. Packer M, Anker SD, Butler J, Filippatos G, Pocock SJ, Carson P, Januzzi J, Verma S, Tsutsui H, Brueckmann M, et al; EMPEROR-Reduced Trial Investigators. Cardiovascular and Renal outcomes with empagliflozin in heart failure. *N Engl J Med*. 2020;383:1413–1424. doi: 10.1056/NEJMoa2022190
 13. Verma S, Mazer CD, Yan AT, Mason T, Garg V, Teoh H, Zuo F, Quan A, Farkouh ME, Fitchett DH, et al. Effect of empagliflozin on left ventricular mass in patients with type 2 diabetes mellitus and coronary artery disease: the EMPA-HEART CardiLink-6 randomized clinical trial. *Circulation*. 2019;140:1693–1702. doi: 10.1161/CIRCULATIONAHA.119.042375
 14. Singh JSS, Mordi IR, Vickneson K, Fathi A, Donnan PT, Mohan M, Choy AMJ, Gandy S, George J, Khan F, et al. Dapagliflozin versus placebo on left ventricular remodeling in patients with diabetes and heart failure: the REFORM trial. *Diabetes Care*. 2020;43:1356–1359. doi: 10.2337/dc19-2187
 15. Green CP, Porter CB, Bresnahan DR, Spertus JA. Development and evaluation of the Kansas City Cardiomyopathy Questionnaire: a new health status measure for heart failure. *J Am Coll Cardiol*. 2000;35:1245–1255. doi: 10.1016/s0735-1097(00)00531-3
 16. Demers C, McKelvie RS, Negassa A, Yusuf S; RESOLVD Pilot Study Investigators. Reliability, validity, and responsiveness of the six-minute walk test in patients with heart failure. *Am Heart J*. 2001;142:698–703. doi: 10.1067/mhj.2001.118468
 17. Platz E, Campbell RT, Claggett B, Lewis EF, Groarke JD, Docherty KF, Lee MMY, Merz AA, Silverman M, Swamy V, et al. Lung ultrasound in acute heart failure: prevalence of pulmonary congestion and short- and long-term outcomes. *JACC Heart Fail*. 2019;7:849–858. doi: 10.1016/j.jchf.2019.07.008
 18. Konstam MA, Kramer DG, Patel AR, Maron MS, Udelson JE. Left ventricular remodeling in heart failure: current concepts in clinical significance and assessment. *JACC Cardiovasc Imaging*. 2011;4:98–108. doi: 10.1016/j.jcmg.2010.10.008
 19. Kang DH, Park SJ, Shin SH, Hong GR, Lee S, Kim MS, Yun SC, Song JM, Park SW, Kim JJ. Angiotensin receptor neprilysin inhibitor for functional mitral regurgitation. *Circulation*. 2019;139:1354–1365. doi: 10.1161/CIRCULATIONAHA.118.037077
 20. Maeda K, Tsutamoto T, Wada A, Hisanaga T, Kinoshita M. Plasma brain natriuretic peptide as a biochemical marker of high left ventricular end-diastolic pressure in patients with symptomatic left ventricular dysfunction. *Am Heart J*. 1998;135(5 pt 1):825–832. doi: 10.1016/s0002-8703(98)70041-9
 21. Packer M. Cardioprotective effects of sirtuin-1 and its downstream effectors: potential role in mediating the heart failure benefits of SGLT2 (sodium-glucose cotransporter 2) inhibitors. *Circ Heart Fail*. 2020;13:e007197. doi: 10.1161/CIRCHEARTFAILURE.120.007197
 22. Jung IH, Park JH, Lee JA, Kim GS, Lee HY, Byun YS, Kim BO. Left Ventricular Global Longitudinal Strain as a Predictor for Left Ventricular Reverse Remodeling in Dilated Cardiomyopathy. *J Cardiovasc Imaging*. 2020;28:137–149. doi: 10.4250/jcvi.2019.0111
 23. Petersen SE, Aung N, Sanghvi MM, Zemrak F, Fung K, Paiva JM, Francis JM, Khanji MY, Lukaschuk E, Lee AM, et al. Reference ranges for cardiac structure and function using cardiovascular magnetic resonance (CMR) in Caucasians from the UK Biobank population cohort. *J Cardiovasc Magn Reson*. 2017;19:18. doi: 10.1186/s12968-017-0327-9
 24. Le TT, Tan RS, De Deyn M, Goh EP, Han Y, Leong BR, Cook SA, Chin CW. Cardiovascular magnetic resonance reference ranges for the heart and aorta in Chinese at 3T. *J Cardiovasc Magn Reson*. 2016;18:21. doi: 10.1186/s12968-016-0236-3
 25. Dreisbach JG, Mathur S, Houbois CP, Oechslin E, Ross H, Hanneman K, Wintersperger BJ. Cardiovascular magnetic resonance based diagnosis of left ventricular non-compaction cardiomyopathy: impact of cine bSSFP strain analysis. *J Cardiovasc Magn Reson*. 2020;22:9. doi: 10.1186/s12968-020-0599-3
 26. Liu H, Wang J, Pan Y, Ge Y, Guo Z, Zhao S. Early and quantitative assessment of myocardial deformation in essential hypertension patients by using cardiovascular magnetic resonance feature tracking. *Sci Rep*. 2020;10:3582. doi: 10.1038/s41598-020-60537-x
 27. Newton N, Opdahl A, Choi EY, Almeida AL, Kawel N, Wu CO, Burke GL, Liu S, Liu K, Bluemke DA, et al. Left ventricular global function index by magnetic resonance imaging—a novel marker for assessment of cardiac performance for the prediction of cardiovascular events: the multi-ethnic study of atherosclerosis. *Hypertension*. 2013;61:770–778. doi: 10.1161/HYPERTENSIONAHA.111.198028
 28. Eitel I, Pöss J, Jobs A, Eitel C, de Waha S, Barkhausen J, Desch S, Thiele H. Left ventricular global function index assessed by cardiovascular magnetic resonance for the prediction of cardiovascular events in ST-elevation myocardial infarction. *J Cardiovasc Magn Reson*. 2015;17:62. doi: 10.1186/s12968-015-0161-x
 29. McComb C, Berry C. Prognostic importance of a new measure of global systolic heart function in healthy adults. *Hypertension*. 2013;61:762–764. doi: 10.1161/HYPERTENSIONAHA.112.199992
 30. Sourbron S. paresmedima/PMI-0.4-SUGAR: Outcomes. October 13, 2020. Accessed October 14, 2020. <http://doi.org/10.5281/zenodo.4086055>.
 31. Haaf P, Garg P, Messroghli DR, Broadbent DA, Greenwood JP, Plein S. Cardiac T1 mapping and extracellular volume (ECV) in clinical practice: a comprehensive review. *J Cardiovasc Magn Reson*. 2016;18:89. doi: 10.1186/s12968-016-0308-4
 32. Romano S, Judd RM, Kim RJ, Kim HW, Klem I, Heitner JF, Shah DJ, Jue J, White BE, Indorkar R, et al. Feature-tracking global longitudinal strain predicts death in a multicenter population of patients with ischemic and nonischemic dilated cardiomyopathy incremental to ejection fraction and late gadolinium enhancement. *JACC Cardiovasc Imaging*. 2018;11:1419–1429. doi: 10.1016/j.jcmg.2017.10.024