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Rationale and design of the United Kingdom Heart Failure with Preserved Ejection Fraction Registry

UK HFpEF Collaborative Group

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ABSTRACT

Objective Heart failure with preserved ejection fraction (HFpEF) is a common heterogeneous syndrome that remains imprecisely defined and consequently has limited treatment options and poor outcomes.

Methods The UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF) is a prospective data-enabled cohort and platform study. The study will develop a large, highly characterised cohort of patients with HFpEF. A biobank will be established. Deep clinical phenotyping, imaging, multiomics and centrally held national electronic health record data will be integrated at scale, in order to reclassify HFpEF into distinct subgroups, improve understanding of disease mechanisms and identify new biological pathways and molecular targets. Together, these will form the basis for developing diagnostics and targeted therapeutics specific to subgroups. It will be a platform for more effective and efficient trials, focusing on subgroups in whom targeted interventions are expected to be effective, with consent in place to facilitate rapid recruitment, and linkage for follow-up. Patients with a diagnosis of HFpEF made by a heart failure specialist, who have had natriuretic peptide levels measured and a left ventricular ejection fraction >40% are eligible. Patients with an ejection fraction between 40% and 49% will be limited to no more than 25% of the cohort.

Conclusions UK HFpEF will develop a rich, multimodal data resource to enable the identification of disease endotypes and develop more effective diagnostic strategies, precise risk stratification and targeted therapeutics.

Trial registration number NCT05441839.

INTRODUCTION

Heart failure (HF) is a major public health problem, with large and growing individual, societal and economic impacts.^{1,2} In Western-type and developed countries, lifetime risk for HF is high, estimated to be between 1 in 5 and 1 in 3, and prevalence is expected to increase by around 50% over the next 20 years.^{3,4} HF is the leading cause of hospitalisation for people aged over 65, with a subsequent 1-year mortality rate of more than 30%.^{5,6} Quality of life is markedly impaired compared with other chronic diseases.⁷ Importantly, there are substantial geographic, socio-economic and ethnic disparities in HF incidence and outcomes.^{8,9} The economic burden of HF on health-care systems is considerable; in 2012, the estimated global cost of HF was \$108 billion per annum, and in 2019 estimated costs were more than \$24 000 per patient per year in the USA.^{10,11}

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ Heart failure with preserved ejection fraction (HFpEF) is a common heterogenous systemic syndrome with limited treatment options.

WHAT THIS STUDY ADDS

⇒ UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF) is a prospective data-enabled cohort and platform study.
⇒ Rich, multimodal data resource to enable identification of endotypes and develop more effective diagnostic strategies, precise risk stratification and targeted therapeutics.
⇒ Platform for more effective and efficient trials, targeting interventions, consent in place for rapid recruitment, linkage for outcomes.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

⇒ UK HFpEF is a unique resource that will become a key platform for collaborative UK clinical and translational HFpEF research.

Approximately half of patients with HF have a left ventricular ejection fraction (LVEF) that is not markedly abnormal. The Candesartan in HF Assessment of Reduction in Mortality and Morbidity (CHARM) trial programme gave rise to the term ‘preserved’ EF, referring to patients with an EF >40%. Rather than being based on biology, it was a pragmatic approach to distinguish this group from the better studied group of patients with HF and a lower EF, for whom evidence-based therapies already existed and for whom placebo was thus not an appropriate comparator, and because EF was available in all patients.^{12,13} More recently, HF guidelines use the term HFpEF (heart failure with preserved ejection fraction) to designate a group with an LVEF >50%, and HF with mildly reduced EF for LVEF 41–49%, while also recognising that EF is a continuum, that its measurement is associated with error and that imaging guidelines from the same societies use different thresholds.^{1,2}

Instead of being a single diagnosis, it is clear that HFpEF represents a heterogenous systemic syndrome. A wide range of cardiovascular and systemic disease mechanisms are described.¹⁴ Patients typically have a range of long-term cardiometabolic and other conditions. Outcomes are also variable; around half of



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deaths in patients with HFpEF are non-cardiovascular, and fewer than 20% of hospital admissions are due to HF.^{15 16}

Understanding of HFpEF remains limited. For example, it is unclear why some people with cardiometabolic conditions develop HFpEF but most do not; there is a lack of pathophysiological features that reliably distinguish HFpEF from its risk factors and normal ageing; the marked heterogeneity in cardiovascular and ‘extra-cardiovascular’ phenotypes is unexplained; and predictors of outcomes are poorly defined.

As a consequence, identification and management of patients with HFpEF remains challenging. A variety of diagnostic guidelines and algorithms exist, each comprising differing variables with varying measurement thresholds.^{17 18} No single classification or score is able to determine the presence or absence of HFpEF. Treatment options are limited. Almost all phase III trials have been neutral, in part because study design has considered HFpEF to be a single disease entity. Indeed, it is noteworthy that subgroup analyses suggest that some agents might be effective in subgroups of patients with particular characteristics.^{19–22} Sodium-glucose co-transporter-2 inhibitors are undoubtedly moving the field forward, improving quality of life and reducing the risk of HF hospitalisation, as well as prompting discussion around organisation of care.^{15 16} No therapies, however, reduce mortality. The heterogeneity of patients with the same diagnostic label, lack of unified diagnostic criteria and limited treatment options have resulted in patients receiving inconsistent care.

It is with these factors in mind that the UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF) has been conceived and designed. The overarching goal is that developing a large, deeply characterised cohort will enable novel understanding of HFpEF and identification of distinct disease endotypes, which will form the basis for more effective diagnostic strategies, precise risk stratification and targeted therapies, that will lead to improved quality of life and outcomes.

DESIGN AND METHODS

Overall study design and aims

UK HFpEF is a national prospective data-enabled cohort and platform study. It is a unique resource that will become a key platform for collaborative UK clinical and translational HFpEF

research. The overarching objective is to provide genotyping and deep phenotyping, linked to outcomes, in a large cohort of patients. This will enable machine learning techniques to be applied in order to reclassify HFpEF into more distinct diagnoses. It will be a platform for the development of diagnostics specific to the different HFpEF subgroups, and for more effective trials that will target subgroups in whom new, repurposed or previously discarded treatments are expected to be effective. Moreover, it will provide cohorts of patients readily available for recruitment to such trials, with linkage in place for follow-up. It will enable scaled investigation aimed at understanding the causes of HFpEF, improving risk stratification and facilitating preventative intervention, and will leverage commercial funding and participation, facilitated by simplified, single-point, UK-wide access. Overall study design and aims are summarised in [figure 1](#) and [box 1](#).

Oversight and governance

The study sponsor is Manchester University NHS Foundation Trust. The Executive Steering Committee is responsible for oversight of study conduct. The study was designed by the Executive Steering Committee, working in conjunction with a dedicated patient advisory group and a working group that includes representatives from sites expressing an interest in participating and other expertise such as data science, data governance and statistics. Membership of the study management committees is given in the online supplemental appendix.

Study set-up and initial recruitment are funded by the UK National Institute for Health and Care Research (NIHR) (NIHR301848). The funder had no role in study design other than through their external peer review processes. The study is supported by the NIHR–British Heart Foundation (BHF) Cardiovascular Partnership and the British Society for Heart Failure, is adopted onto the NIHR Clinical Research Network Portfolio (Central Portfolio Management System reference 52749) and registered at ClinicalTrials.gov (NCT05441839). Study website: <https://www.ukhfpef.org/>

Patients

Eligibility criteria are summarised in [box 2](#). Enrolment began on 7 October 2022 after approval. The study is being conducted in

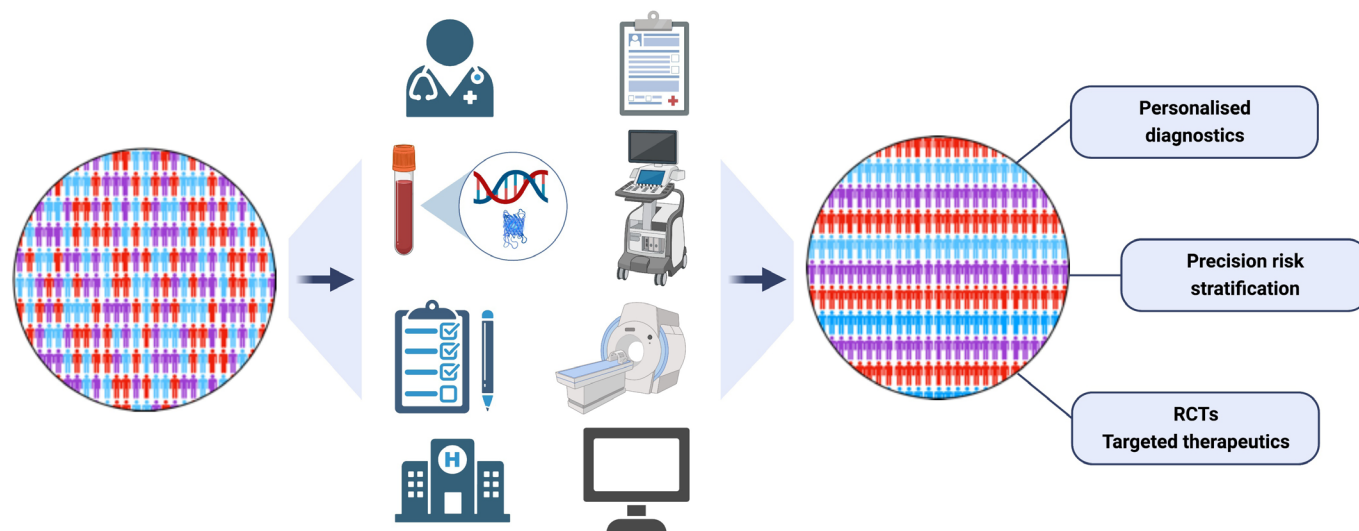


Figure 1 UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF) overview. UK HFpEF will develop a rich, multimodal data resource to enable the identification of disease endotypes and develop more effective diagnostic strategies, precise risk stratification and targeted therapeutics. Figure created with BioRender.com. RCTs, Hurandomised controlled trials.

Box 1 Aims of UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF)

To develop a large, deeply characterised cohort that will be a platform for collaborative clinical and translational HFpEF research, in order to:

- ⇒ Reclassify HFpEF into distinct diagnoses, where possible, based on disease mechanisms, clinical factors and outcome.
- ⇒ Evaluate whether patients in the distinct groups respond differentially to treatments, with the aim of predicting individual patient treatment response.
- ⇒ Create a platform for clinical trials that:
 - Matches mechanism of action of therapies (new, repurposed or previously discarded) with HFpEF subgroup/anticipated treatment response.
 - Provides groups of patients readily available for recruitment to trials.
 - Has data linkage in place for clinical outcomes.
- ⇒ Create a platform for identifying phenotypic and genetic factors that could be used as the basis for:
 - Improving understanding of the causes of HFpEF.
 - Developing diagnostics.
 - Improving risk stratification.
- ⇒ Facilitate industry engagement by providing a single point of access for industry.

HFpEF, heart failure with preserved ejection fraction.

accordance with the principles of Good Clinical Practice and the World Medical Association Declaration of Helsinki. All participants provided written informed consent.

Study procedures

The study protocol, participant information sheet and consent form are available at <https://www.ukhfpef.org/>

Recruitment

It is expected that the majority of patients will be recruited via HF services, including outpatient clinics and inpatient wards. Patients can also be identified by primary care physicians with HF expertise, or express an interest in participating directly via patient-centred recruitment platforms such as CardioTrials (<https://cardiotrials.org/>), in which case they will be invited to the most appropriate secondary care centre. Reasons for non-recruitment will be recorded in a screening log.

Baseline evaluations

Study assessments and procedures are illustrated in [figure 2](#). The study uses data collected as part of clinical care supplemented with study-specific data.

Personal data

Participant personal details are collected in order to retrieve medical, health and social care data from local, regional and national data systems and organisations, and to allow participants to be contacted regarding stage 2 studies (see below).

Medical, health and social care information

Data collected include demographics, medical history, HF history, medications, laboratory investigations, ECG, echocardiography and other investigations that participants may undergo as part of their clinical care, for example, cardiac catheterisation,

Box 2 Eligibility criteria for UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF)**Inclusion criteria**

- ⇒ Written informed consent.
- ⇒ Diagnosis of HFpEF by an HF specialist (eg, a cardiologist with HF expertise, a primary care physician with HF expertise, a secondary/tertiary care physician with HF expertise, an HF nurse specialist, a specialist HF pharmacist).*
- ⇒ Natriuretic peptide levels measured.

Exclusion criteria

- ⇒ LVEF ever <40%.† (For clarity, patients with a previous LVEF below 40%, which has since improved to above 40%, are excluded.)‡
- ⇒ Known infiltrative cardiomyopathy (eg, amyloid, sarcoid, lymphoma, endomyocardial fibrosis).
- ⇒ Known active myocarditis, constrictive pericarditis or cardiac tamponade.
- ⇒ Known genetic hypertrophic cardiomyopathy or obstructive hypertrophic cardiomyopathy.
- ⇒ Known arrhythmogenic right ventricular cardiomyopathy.
- ⇒ Known severe primary valvular heart disease.
- ⇒ Known idiopathic, heritable or drug-induced pulmonary arterial hypertension.
- ⇒ Heart transplantation or ventricular assist device.
- ⇒ Complex congenital heart disease.

*Original text (diagnosis of HFpEF by a cardiologist with HF expertise, or a primary care physician with HF expertise, or a heart failure nurse) clarified in a protocol amendment.

†Note regarding LVEF: recruitment will be centrally monitored; the proportion of participants with LVEF 40–49% will be limited to no more than 25% of the cohort.

‡Original text (LVEF<40% (at screening or any previous measurement)) clarified in a protocol amendment.

HF, heart failure; LVEF, left ventricular ejection fraction.

cardiopulmonary exercise testing, exercise echocardiogram, heart rhythm monitoring, nuclear scintigraphy. Date of each item is also captured.

To standardise the clinical evaluation that patients with HFpEF receive across the UK, and data collection, a core set of laboratory investigations are advised (online supplemental appendix). Similarly, there is a standardised echocardiography protocol, in line with the British Society of Echocardiography minimum data set (online supplemental appendix).

Pseudonymised ECGs and echocardiogram digital imaging and communications in medicine (DICOM) images are uploaded to the study database, where they are available for central analysis, including using automated artificial intelligence (AI) algorithms.

Physical status

Data collected include HF symptoms and signs, New York Heart Association (NYHA) class, blood pressure, pulse rate, height, weight and Rockwood Clinical Frailty Scale.²³

Patient-reported outcome measure

To characterise patient health-related quality of life, the Minnesota Living with Heart Failure Questionnaire is conducted.²⁴

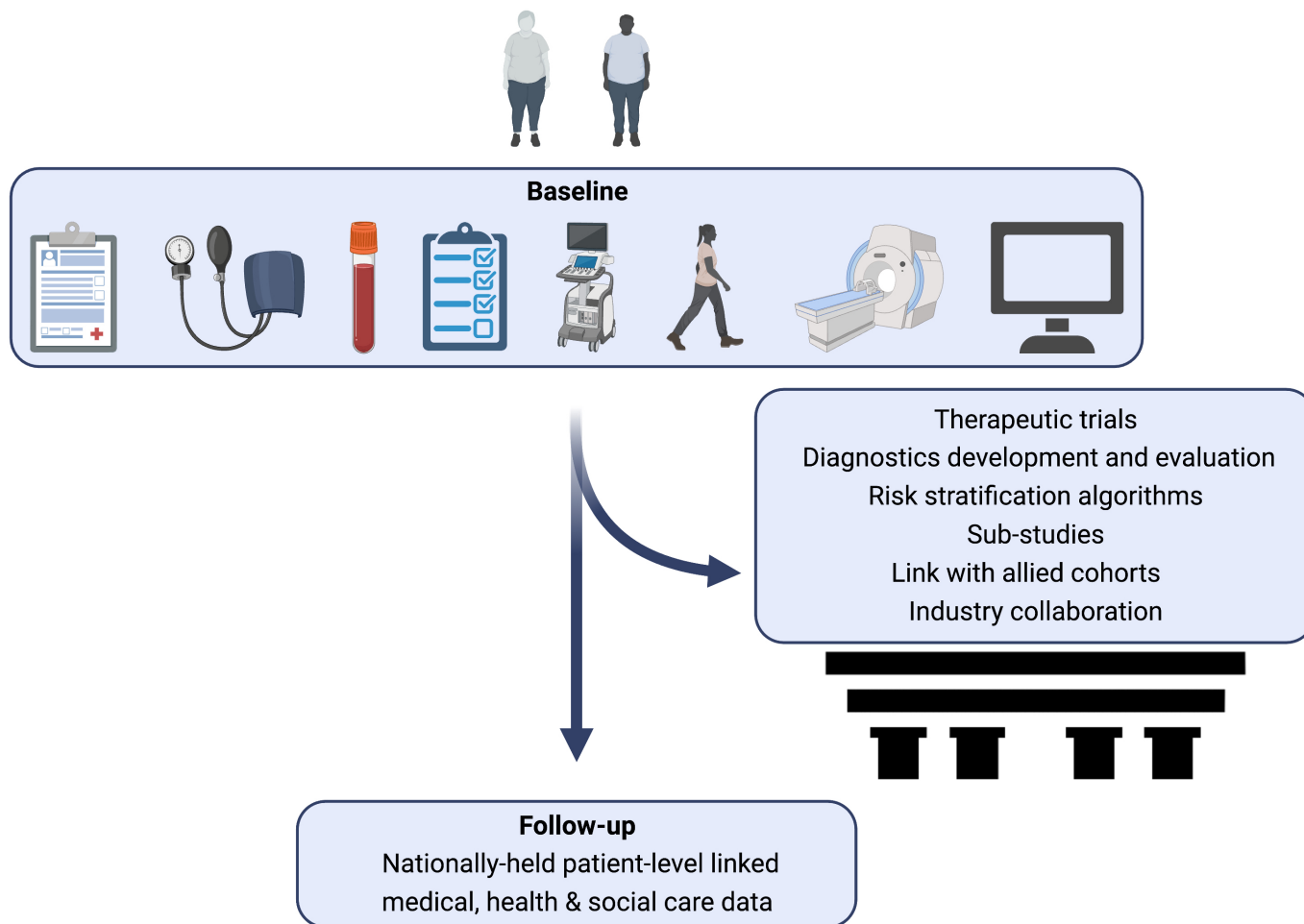


Figure 2 UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF) study design. Baseline assessments use data collected as part of clinical care supplemented with study-specific data. Recruitment to other heart failure with preserved ejection fraction (HFpEF)-related studies, such as clinical trials and evaluation of diagnostics, will be supported. Follow-up will be incorporated from national healthcare data services. See text for further details. Figure created with BioRender.com.

Blood sampling

Up to 50 mL of blood is collected and aliquoted (plasma (10 aliquots), serum (10 aliquots), buffy coat (4 aliquots)), before being transferred for central storage at NIHR National Biosample Centre.

Substudies

Substudies will focus on specific aspects of HFpEF in addition to the core data set, involving investigators and sites with a particular interest. This approach ensures that the registry population is as representative of HFpEF as possible, while also providing a platform for more specific evaluations. Substudies will benefit from the data present in the wider registry, and the wider registry will benefit from data collected as part of the substudies. Substudies may include, for example, invasive assessments. Example substudies included from the outset are:

Exercise capacity

Where possible, in terms of site logistics and participant characteristics, 6-min walk testing is performed.²⁵

Cardiovascular MRI

Where participants are undergoing cardiovascular MRI (CMR) as part of their clinical care, a standardised protocol is advised

(online supplemental appendix). This includes approximately 5 min of additional research imaging above that considered part of the clinical scan. The pseudonymised DICOM images are uploaded to the study database, where they are available for central analysis.

Blood sample analyses

Multiple types of analyses will be performed on the donated samples including genomic, transcriptomic, proteomic, lipidomic, metabolomic and biochemical analyses, dependent on future funding. Participants provide consent for sequencing up to the level of the whole genome. Data generated from the samples will be linked with the other data.

Data flow and management

Figure 3 provides an overview of the flow and management of data. Data collected at sites are entered via a secure Research Electronic Data Capture (REDCap, version 12.4.11) web application.²⁶ Consent forms and pseudonymised ECGs are uploaded via REDCap, and pseudonymised echocardiogram and CMR DICOM files are uploaded via REDCap using a secure file transfer system (ownCloud V.10.11.0). In addition, patient-level data will be incorporated from national healthcare data services, such as NHS England, Digital Health and Care Northern Ireland

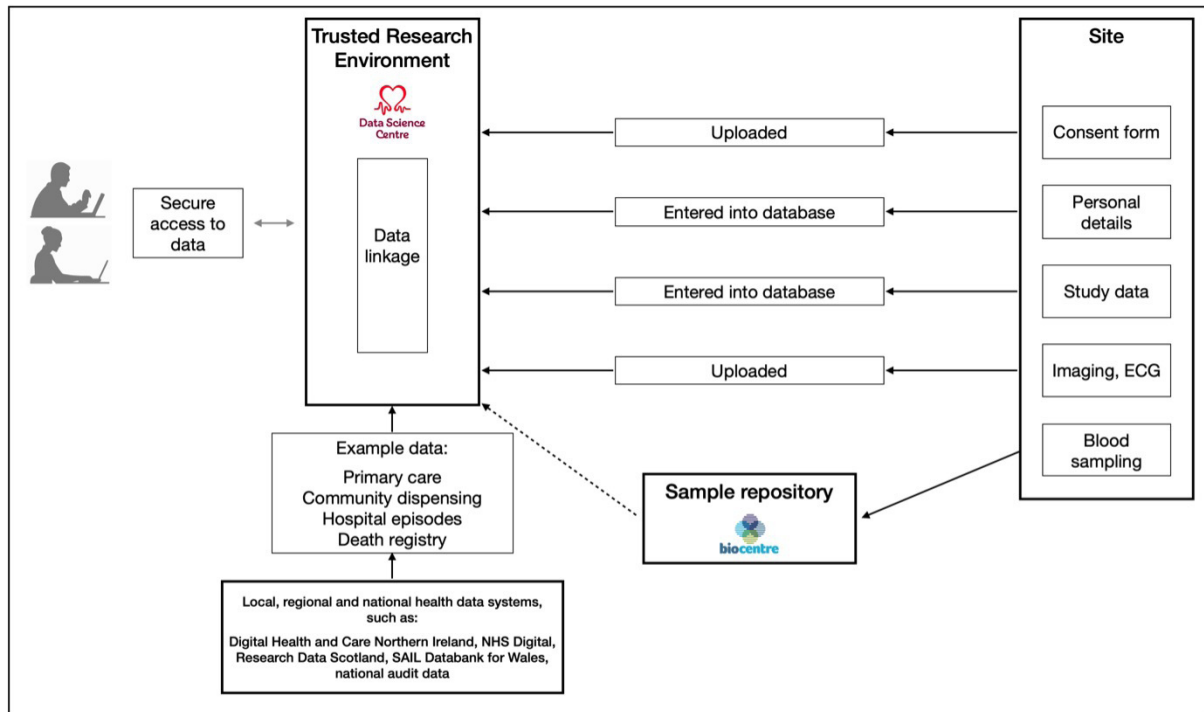


Figure 3 UK Heart Failure with Preserved Ejection Fraction Registry (UK HFpEF) data flow and management. Data collected at sites are entered and uploaded via a secure web application. Patient-level data will be incorporated from national healthcare data services. Data are stored in an accredited Trusted Research Environment in collaboration with the British Heart Foundation Data Science Centre. See text for further details. NHS, National Health Service; SAIL, Secure Anonymised Information Linkage.

(Northern Ireland), Research Data Scotland (Scotland) and Secure Anonymised Information Linkage Databank for Wales (Wales). Such data include primary care, community dispensing, hospital episode and death registry data.

Data are stored in an accredited Trusted Research Environment in collaboration with the BHF Data Science Centre, part of Health Data Research UK, the UK's Institute for Health data science. The Trusted Research Environment enables highly secure and privacy-preserving data storage, access, sharing, analysis and linkage in a well-governed environment. Access, determined by the Executive Steering Committee, is controlled, secure and role based on a named person basis only, with individual user authentication. The Trusted Research Environment provides an accredited platform to enable data sharing framework agreements with national data organisations and services, allowing the data described above to be incorporated and linked at participant level. It also provides an analysis environment that supports statistical software. Activity is audited and outputs are reviewed by the Executive Steering Committee before they are released. REDCap and ownCloud are hosted within the Trusted Research Environment.

Participant confidentiality

Participants are given a unique participant identification number. Deidentified, pseudonymised study data are held in the 'UK HFpEF Research Data' database in the Trusted Research Environment. Personal data and uploaded consent forms are held in a separate 'UK HFpEF Consent & Personal Information' database in the Trusted Research Environment. The link between the pseudoidentifiers and participant personal details is kept securely in the Consent & Personal Information database. Only a small number of appropriately trained, named study team members

determined by the Executive Steering Committee can access the identifiable information.

Stage 2 studies

A key aim of the study is to be a platform to support recruitment to other HFpEF-related studies, such as clinical trials of novel and repurposed therapies and evaluation of diagnostics. The genotyping and deep phenotyping mean that patients who meet recruitment criteria for other studies can be readily identified and contacted, enabling efficient recruitment and more effective research, for example, allowing mechanism of action of a new therapy to be matched with anticipated individual treatment response. The data linkage may be used to support collection of clinical outcomes for these other studies. Importantly, the process provides patients who would like to take part in other studies the opportunity to do so.

UK HFpEF participants are asked to provide consent to being contacted regarding up to four stage 2 studies in any 12-month period. It is generally expected that data generated from stage 2 studies will be deposited in the UK HFpEF database.

Industry collaboration

In line with the NIHR–BHF Cardiovascular Partnership strategy, an important objective is to develop appropriate industry collaborations. Private sector partnership provides opportunity for resource and expertise to support HFpEF research, and to access novel technologies at an early stage to facilitate innovative research and more rapid translation. The Executive Steering Committee will retain control of commercial relationships, which are anticipated to be on a project-by-project basis. Multiple industry partners are expected rather than exclusivity. As a prerequisite, data generated must be deposited back into the

study database. All participants are made aware of the potential for industry collaboration in the participant information sheet and are specifically asked to provide consent to their data and samples being shared with industry.

Additional methods are found in the online supplemental appendix.

DISCUSSION

UK HFpEF realises the full potential of the UK healthcare system and clinical research infrastructure for cardiovascular research. The study combines research-specific data, with clinical data available via individual patient records at sites, nationally held patient-level healthcare data, contemporary imaging and biobanking, at scale, from all four nations. It is supported by the NIHR-BHF Cardiovascular Partnership, which brings together NIHR and BHF research infrastructure, and the NIHR Clinical Research Network. It is a vanguard for the BHF Data Science Centre, specifically its 'Enabling Cohorts' thematic area, and samples are stored centrally in the NIHR National Biosample Centre. The study aims to link with similar projects internationally, such as the National Heart, Lung and Blood Institute (NHLBI) HeartShare programme,²⁷ and other large studies investigating allied pathophysiology, such as the UK Pulmonary Arterial Hypertension Cohort Study.

Precision medicine requires detailed characterisation, at scale, and digital technologies to make sense of the data optimally. UK HFpEF will develop a rich, multimodal data resource that integrates deep clinical phenotyping, imaging, multiomics and electronic health records in a large cohort. Machine learning algorithms will be applied to reclassify HFpEF into subgroups, improve understanding of disease mechanisms underlying the development and progression of HFpEF and identify novel biological pathways, new molecular targets and validation of existing targets. Together, these will form the basis for developing diagnostics and targeted therapeutics specific to subgroups. Moreover, the study will be a platform for more effective and efficient trials, focusing on groups of patients in whom interventions are expected to be effective, with consent in place to facilitate rapid recruitment, and linkage in place for follow-up. While existing HF registries aim to address important knowledge gaps, UK HFpEF will focus on HFpEF and combine extensive phenotyping with genetic data at scale to enable systems biology approaches. Similar approaches have yielded important translational insights into other high burden diseases such as type 2 diabetes; however, a lack of available data sets has stymied the application of these approaches to HFpEF.

Previous attempts to reclassify HFpEF with machine learning techniques suggest that distinct subgroups may exist, but have been limited by sparse characterisation and often small retrospective cohorts. The resulting clusters have, therefore, been superficial and not advanced the field. The analysis of genetic susceptibility to HFpEF is likely to be important for deconvoluting causal factors and therapeutic targets: to date, however, genetic investigations have been restricted by sample size and limited phenotyping.

Recruitment criteria are pragmatic. There are no widely accepted evidence-based HFpEF diagnostic criteria, and the field is rapidly evolving. Requiring a diagnosis of HFpEF by a healthcare professional with specialist HF expertise provides specificity, and means the cohort reflects current practice. It was felt important to document natriuretic peptide levels in all patients, but given the lack of consensus regarding appropriate thresholds, and because levels are confounded by factors common to this

patient group, no threshold is specified. Instead, the cohort will be used to develop new criteria, while also evaluating existing criteria and scoring systems.

An LVEF threshold of 40% was chosen, after extensive Steering Committee and working group discussion, for a number of reasons: (1) the guideline-designated LVEF cut-offs are recognised within the guidelines as being 'arbitrary';¹; (2) LVEF measurement is associated with substantial variation²⁸; (3) of the eight phase III HFpEF trials currently listed on ClinicalTrials.gov as 'recruiting' or 'active not recruiting', seven have an entry LVEF criterion of less than 50%, and thus an LVEF threshold of 40% permits investigation of the patients that are being included in HFpEF trials²⁹; and (4) fundamentally, the aim of the study is to move the field forward, so that patient evaluation and intervention become based on the underlying biology.

Including patient-level linked electronic medical, health and social care record data from national data services provides several advantages. It will add retrospective information to enrich medical history data, and prospective information to enable longitudinal follow-up of health-related outcomes. Less manual data input is required, and as more data become available via national services, manual data input should reduce further. Community-prescribing data are accurate and contemporary, and hospital episode and death data, which would otherwise be challenging to collect at scale, are comprehensive.

Having raw DICOM image data available centrally will facilitate development and application of current and future AI image analysis algorithms, data from which will be incorporated into the wider machine learning analyses. The study provides the opportunity for synergistic training and development of clinically and non-clinically trained individuals.

In summary, UK HFpEF will develop a rich, multimodal data resource to enable novel understanding of HFpEF, which will form the basis for more effective diagnostic strategies, precise risk stratification and targeted therapies.

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Contributors All authors inputted to study design and are involved with study delivery. The manuscript was drafted by CAM and all authors provided critical input and review. CAM is responsible for the overall content as guarantor.

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Competing interests CAM has served on advisory boards for AstraZeneca, Boehringer Ingelheim and Lilly Alliance, Novartis, and PureTech Health; serves as an advisor for HAYA Therapeutics; and has received speaker fees from Boehringer Ingelheim and Novo Nordisk, conference attendance support from AstraZeneca and research support from Amicus Therapeutics, AstraZeneca, Guerbet Laboratories, Roche and Univar Solutions. RTL has received research grants from Pfizer and has provided consultancy for FITFILE and HealthLumen. SEP provides consultancy to Circle Cardiovascular Imaging, Calgary, Alberta, Canada.

Patient and public involvement Patients and/or the public were involved in the design, or conduct, or reporting, or dissemination plans of this research. Refer to the Methods section for further details.

Patient consent for publication Not applicable.

Ethics approval This study involves human participants and was approved by the NHS Health Research Authority and the London-Fulham Research Ethics Committee (IRAS project ID: 314091) (REC reference: 22/PR/0543). Participants gave informed consent to participate in the study before taking part.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data sharing not applicable as no data sets generated and/or analysed for this study. No data were analysed or available for this registry protocol manuscript.

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**UK Heart Failure with Preserved Ejection Fraction Registry:
rationale and design of UK HFpEF**

Supplemental appendix

The UK HFpEF Collaborative Group

Executive Steering Committee

Name	Affiliation	Expertise
Christopher A Miller	Manchester University NHS Foundation Trust University of Manchester	Chair
Abdallah Al-Mohammad	Sheffield Teaching Hospitals NHS Trust	National Institute for Health and Care Excellence (NICE) guidelines
Janine Beezer	South Tyneside and Sunderland NHS Trust	Specialist Clinical Pharmacist Heart Failure
Emma Columbine	Manchester University NHS Foundation Trust	Sponsor representative
David Cotterell	Manchester University NHS Foundation Trust	Project manager
Simon Fisher	National Institute for Health and Care Research (NIHR) – British Heart Foundation (BHF) Cardiovascular Partnership	NIHR-BHF Cardiovascular Partnership Manager
Nick Hartshorne-Evans	Pumping Marvellous Foundation	Patient representative
Laurence Humphreys-Davies	British Society for Heart Failure	Patient representative
Rebecca Hyland	Wiltshire Health and Care	Heart Failure Nurse Specialist
R Thomas Lumbers	UCL Institute of Health Informatics NIHR University College London Hospitals Biomedical Research Centre Barts Health NHS Trust UCLH NHS Foundation Trust	Genomics; electronic health record; phenotyping
Gerry P McCann	University of Leicester NIHR Leicester Biomedical Research Centre, Glenfield Hospital, Leicester, UK	Cardiovascular magnetic resonance imaging
Maria F Paton Shaun Robinson	University of Leeds Imperial College Healthcare NHS Trust	Echocardiography
Mark C Petrie	University of Glasgow	Trials/therapeutics
Cathie Sudlow	BHF Data Science Centre, Health Data Research UK	Health data (linkages, curation pipelines, trusted research environments); epidemiological design

Patient Advisory Group

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University of Manchester

Niels Peek (Health informatics and data science)

Supplemental methods

Study duration

Participants will remain in the study for 10 years from when they provide consent. It is anticipated that at the end of the study, anonymised study data will be transferred to a managed-access research or scientific archive.

Authorship

A policy for authorship that follows the principles of the International Committee of Medical Journal Editors, written by the Executive Steering Committee and agreed by the Working Group, is in place (available at <https://www.ukhfpef.org/>).

Data Sharing

Participants are asked to provide consent for pseudonymised participant-level study data to be shared for research purposes. Requests for access to data are managed by the Executive Steering Committee. Release of data is subject to scientific review by the Executive Steering Committee and an appropriate Data Transfer Agreement. A Collaboration and Support Policy, written by the Executive Steering Committee and agreed by the Working Group, describes the framework for collaborations (available at <https://www.ukhfpef.org/>).

Supplemental Table 1. Core clinical laboratory investigations

BNP or NT-Pro BNP
hs-Troponin T or I
Haemoglobin
MCV
Haematocrit
White cell count
Sodium
Potassium
Urea
Creatinine
eGFR
Albumin
Alkaline phosphatase
Bilirubin
Alanine aminotransferase
CRP
Iron
Ferritin
Transferrin saturation
Thyroid stimulating hormone
Free T4
HBA1c

Additional clinical laboratory investigations, to record if available

IgG
IgA
IgM
Serum protein electrophoresis
Urine Bence Jones protein
Urine albumin : creatinine ratio
Urine dipstick proteinuria

BNP = Brain natriuretic peptide; CRP C-reactive protein; eGFR = estimated glomerular filtration rate; hs = high sensitivity; Ig = Immunoglobulin; MCV = mean corpuscular volume; NTproBNP = N-terminal pro B-type natriuretic peptide.

Supplemental Table 2. Echocardiography protocol

Standard echo acquisition in line with the British Society of Echocardiography Minimum Dataset.¹
Key views and corresponding measurements are as follows:

Key views	Measurements
Parasternal long axis 2D	LV end-diastolic dimension (cm) LV end-systolic dimension (cm) Maximum wall thickness (mm)
Parasternal long axis RV inflow CWD	TR V_{\max} (m/s)
Apical 4 chamber 2D	LV ejection fraction (%)
Apical 2 chamber 2D	LV ejection fraction (%)
Apical 4 chamber 2D GLS*	Peak GLS (%)
Apical 2 chamber 2D GLS*	Peak GLS (%)
Apical long axis 2D GLS*	Peak GLS (%)
Apical 4 chamber 2D optimised for LA volume	LA volume (cm ³)
Apical 2 chamber 2D optimised for LA volume	LA volume (cm ³)
Apical 4 chamber mitral valve PWD	E V_{\max} (cm/s) A V_{\max} (cm/s) DT (ms)
Apical 4 chamber mitral valve TDI	Lateral e' (cm/s) Septal e' (cm/s)
Apical 5 chamber aortic valve CWD	AV V_{\max} (m/s)
Apical 4 chamber modified for RV/RA 2D	Basal RV diameter Visual assessment of RV function
Apical 4 chamber modified for RV/RA CWD	TR V_{\max} (m/s)
Apical 4 chamber lateral tricuspid valve annulus MM	TAPSE (cm)
Apical 4 chamber right ventricle TDI	RV S' (cm/s)
Subcostal 2D +/- MM	IVC diameter (mm) IVC diameter during inspiration (mm)
Multiple views	Mitral, aortic and tricuspid valve function Pericardial effusion (present/absent)

* As permitted by image quality and local feasibility.

2D = Two-dimensional; A V_{\max} = Peak velocity in late diastole; AV V_{\max} = Aortic valve peak velocity; CWD = Continuous wave Doppler; DT = Flow deceleration time from peak E wave to end of E wave signal; E V_{\max} = Peak velocity in early diastole; GLS = Global longitudinal strain; IVC = Inferior vena cava; LA = Left atrium; LV = Left ventricle; MM = M-mode; PWD = Pulsed wave Doppler; RV = Right ventricle; TAPSE = Tricuspid Annular Plane Systolic Excursion; TDI = Tissue Doppler imaging; TR V_{\max} = Tricuspid regurgitation peak velocity;

1. Robinson et al. A practical guideline for performing a comprehensive transthoracic echocardiogram in adults: the British Society of Echocardiography minimum dataset *Echo Red Pract.* 2020; 4: G59-G93.

Supplemental Table 3. Cardiovascular magnetic resonance protocol

Core

- Localisers
- CH4 cine.
- CH2 cine.
- CH3 cine.
- LVOT cine
- Aortic valve cine
- Gadolinium based contrast agent in line with local policy.
- LV short axis cine stack.
- TI Scout
- LGE segmented inversion recovery and PSIR. CH4, CH2, CH3 and short axis stack

Supplemental

- T1 mapping basal and mid short axis, before and after Gadolinium
- CH4 fat-water sequence.
- T2 mapping. Mid short axis.
- Aortic candy stick cine
- Cine perpendicular to the ascending and descending aorta at pulmonary bifurcation level, with measurement of blood pressure.
- Phase encoded velocity mapping perpendicular to the main pulmonary artery.
- 3D Dixon fat-water sequence, centred over the renal arteries.
- Perfusion imaging if being performed clinically

Notes

- 1.5T or 3T
- The protocol is split into core and supplementary sequences. It is expected that core sequences would be performed as part of a standard clinical CMR.
- As part of site set-up, the central study team will liaise with the site regarding the details of the CMR protocol appropriate for the site, and provide site-specific CMR guidance. The protocol is a guide.

CH = chamber; LGE = late gadolinium enhancement; LV = left ventricle; LVOT = left ventricular outflow tract; PSIR = Phase-sensitive inversion recovery; TI = inversion time.