

Breaking the cycle of heart failure with preserved ejection fraction and atrial fibrillation

Tica, Otilia; Khamboo, Waseem; Kotecha, Dipak

DOI:

[10.15420/cfr.2022.03](https://doi.org/10.15420/cfr.2022.03)

License:

Creative Commons: Attribution-NonCommercial (CC BY-NC)

Document Version

Publisher's PDF, also known as Version of record

Citation for published version (Harvard):

Tica, O, Khamboo, W & Kotecha, D 2022, 'Breaking the cycle of heart failure with preserved ejection fraction and atrial fibrillation', *Cardiac Failure Review*, vol. 8, e32. <https://doi.org/10.15420/cfr.2022.03>

[Link to publication on Research at Birmingham portal](#)

General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.




When citing, please reference the published version.

Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact UBIRA@lists.bham.ac.uk providing details and we will remove access to the work immediately and investigate.

Breaking the Cycle of Heart Failure With Preserved Ejection Fraction and Atrial Fibrillation

Otilia Țica ^{1,2} Waseem Khamboo ¹ and Dipak Kotecha ^{1,3}

1. Institute of Cardiovascular Sciences, University of Birmingham, Medical School, Birmingham, UK;

2. Cardiology Department, Emergency County Clinical Hospital of Oradea, Oradea, Romania;

3. University Hospitals Birmingham NHS Foundation Trust, Birmingham, UK

Abstract

Heart failure with preserved ejection fraction (HFpEF) and AF are two common cardiovascular conditions that are inextricably linked to each other's development and progression, often in multimorbid patients. Current management is often directed to specific components of each disease without considering their joint impact on diagnosis, treatment and prognosis. The result for patients is suboptimal on all three levels, restricting clinicians from preventing major adverse events, including death, which occurs in 20% of patients at 2 years and in 45% at 4 years. New trial evidence and reanalysis of prior trials are providing a glimmer of hope that adverse outcomes can be reduced in those with concurrent HFpEF and AF. This will require a restructuring of care to integrate heart failure and AF teams, alongside those that manage comorbidities. Parallel commencement and non-sequential uptitration of therapeutics across different domains will be vital to ensure that all patients benefit at a personal level, based on their own needs and priorities.

Keywords

Heart failure, heart failure with preserved ejection fraction, AF, comorbidities, management, treatment

Disclosure: DK reports grants from the National Institute for Health Research (NIHR CDF-2015-08-074 RATE-AF, NIHR130280 DaRe2THINK, NIHR132974 D2T-NeuroVascular), the British Heart Foundation (PG/17/55/33087, AA/18/2/34218, FS/CDRF/21/21032), the European Union/European Federation of Pharmaceutical Industries and Associations Innovative Medicines Initiative (BigData@Heart 116074), the European Society of Cardiology supported by educational grants from Boehringer Ingelheim/BMS-Pfizer Alliance, Bayer, Daiichi Sanyo, Boston Scientific, the NIHR/University of Oxford Biomedical Research Centre and the British Heart Foundation/University of Birmingham Accelerator Award (STEEER-AF NCT04396418), Amomed Pharma and IRCCS San Raffaele/Menarini (Beta-blockers in Heart Failure Collaborative Group NCT0083244), as well as advisory board personal fees from Bayer, Amomed, Protherics Medicines Development and Myokardia (all outside the submitted work). DK is an editorial board member for *Cardiac Failure Review*; this did not influence peer review. OT reports funding from EU/EEPIA Innovative Medicines Initiative (BigData@Heart 116074) and Amomed Pharma outside the submitted work. WK has no conflicts of interest to declare.

Received: 11 January 2022 **Accepted:** 9 April 2022 **Citation:** *Cardiac Failure Review* 2022;8:e32. **DOI:** <https://doi.org/10.15420/cfr.2022.03>

Correspondence: Dipak Kotecha, Institute of Cardiovascular Sciences, University of Birmingham, Medical School, Vincent Drive, Birmingham B15 2TT, UK.

E: d.kotecha@bham.ac.uk

Open Access: This work is open access under the CC-BY-NC 4.0 License which allows users to copy, redistribute and make derivative works for non-commercial purposes, provided the original work is cited correctly.

Heart failure (HF) with preserved ejection fraction (HFpEF) and AF are two of the most common cardiovascular conditions encountered in daily practice, and are leading causes of hospitalisation and adverse patient outcomes.^{1,2} Both conditions have increasing prevalence and pose a growing burden on global healthcare systems. They share common risk factors, such as hypertension and obesity, that are themselves increasing in prevalence. Patients with HFpEF or AF are often multimorbid, with advanced age, ischaemic heart disease, diabetes and other non-cardiovascular conditions. HFpEF and AF frequently coexist, and each predisposes to the other. Patients with an already high risk of adverse events (including death) have an even worse prognosis when HFpEF and AF combine.³

In this review we propose various targets to break the cycle between HFpEF and AF, learning lessons from epidemiology, pathophysiology and associated comorbidities to improve diagnosis, treatment and patient well-being. In the rapidly developing fields of HFpEF and AF, we outline where joined-up thinking can help both elements independently, as well

as create a new treatment paradigm for patients with both HFpEF and AF.

Epidemiology of HFpEF and AF

Studies of incident HF in community-based cohorts suggest that HFpEF accounts for between 37% and 53% of HF cases overall, with a higher observed incidence in older participants.⁴⁻⁶ These figures are likely an underestimate of the burden of HFpEF due to the challenges in diagnosing HFpEF, particularly in primary care settings. Hospitalisation due to HF is responsible for a significant proportion of the cost burden of cardiovascular diseases. Although HF with reduced ejection fraction (HFrEF) costs more than HFpEF on an individual basis, the increase in acute admissions is being driven more by HFpEF.^{7,8}

With regard to AF, the projected prevalence is rising rapidly, fuelled by better identification, an ageing population and improved survival from acute coronary syndromes and HF, which, in later life, can increase the risk of AF.⁹ With AF associated with high rates of stroke and thromboembolism, as well as evidence of a clear link with cognitive

decline and vascular dementia, the increase in AF across all communities will have a profound public health impact.¹⁰

The reported prevalence of AF in HFpEF (and vice versa) varies widely, likely due to differences in HFpEF definition, HF severity and, in particular, study selection criteria. For example, research involving AF interventions tends to have lower rates of HFpEF compared with HF interventions reporting AF.^{11–13} In the long-term HF registry of the European Society of Cardiology, the prevalence of AF was 39% among HFpEF patients.¹⁴ Again, this is likely an underestimate because it does not consider the temporal relationship between these conditions.¹⁵ In Framingham participants with new-onset HF (1980–2012), the overall rate of AF in HFpEF (considering previous, concurrent and future AF) was 62%; this was significantly higher than the 55% of patients with AF in the context of HFrEF.¹⁶ Based on these figures, the prevailing notion that AF is common in HFpEF is wrong; in fact, over time, having concomitant HFpEF and AF is actually the norm. This has major implications on the ability of cardiologists and general physicians to improve patient outcomes, as discussed below.

Mechanisms Underlying Both Heart Failure With Preserved Ejection Fraction and AF

There are multilevel links between HF and AF, contributing both simple and complex mechanisms that lead to concurrence in individual patients. The inter-relationship between HFpEF and AF is outlined in *Figure 1* and includes cellular, biohumoral, structural and haemodynamic changes from HFpEF and AF that cause the progression of each condition and increase the likelihood of the other developing. Much has been made of this cyclical relationship in the literature, but probably more important to shared pathophysiology and reciprocal causation is the connection to a set of similar comorbidities. Early ageing, hypertension, coronary artery disease, diabetes, obesity and a range of other comorbidities are all antecedents of both HFpEF and AF, many with inflammation as an underlying trigger. The sequence that links inflammation to HFpEF, AF and both diseases combined includes endothelial dysfunction and oxidative stress, culminating in end-organ manifestations, such as diastolic dysfunction.¹⁷ In addition, multimorbidity is increasing in patients with HF, as demonstrated by a longitudinal study in which 87% had three or more comorbidities in 2012–14, compared to 68% a decade previously.¹⁸ The interactions of these comorbidities will place additional burden on the mechanisms portending to HFpEF and AF.

With regard to more specific cardiac interactions, left atrial structural and functional remodelling is a clear mechanism through which HFpEF leads to AF. Left atrial enlargement and pressure change is commonly associated with a proarrhythmic substrate due to atrial fibrosis, which promotes further electrical remodelling, decreases the atrial effective refractory period and enhances the risk and burden of AF.¹⁹ Subsequent upregulation of the adrenergic and renin–angiotensin–aldosterone systems can accentuate atrial fibrosis, and changes to atrial and ventricular natriuretic peptide release and other neurohormonal activation and haemodynamic changes can trigger the development of ventricular myocardial fibrosis. This and the resultant structural changes (often made worse by valve dysfunction) further worsen HFpEF status and set up a continuum of deteriorating cardiac output. Added to this, persistent tachycardia from uncontrolled AF can contribute to both an atrial and ventricular cardiomyopathy.²⁰

Prognostic Implications

The poor outlook for patients with either HFpEF or persistent forms of AF is further worsened when these conditions combine, augmented by the

impact of interacting comorbidities and varying due to the heterogeneity of HFpEF.^{16,21,22} Adverse event rates are generally increased, most notably death. Incident AF can double mortality risk in patients with HFpEF, independent of underlying risk factors.¹⁵ Extrapolating from the published sources described in *Figure 2*, average absolute mortality rates are approximately 20% at 2 years in patients with combined HFpEF and AF, increasing to around 45% at 4 years. In a meta-analysis of 45,100 patients, the increase in mortality was higher when HFrEF was combined with AF (relative risk 1.24 versus HFpEF-AF; 95% CI [1.12–1.36]; $p < 0.001$).³⁷ However, there was no significant difference between HFrEF-AF and HFpEF-AF in the rates of hospitalisation due to HF, or incident stroke. Patients with HF and AF have poor quality of life, substantially worse across most domains than other long-standing illnesses, with a negative trajectory over time and more patients deteriorating than improving.^{38,39}

Diagnostic Challenges for Heart Failure With Preserved Ejection Fraction and AF

Different trials, observational studies and registries have used varying definitions of HFpEF, including various cut-off points for left ventricular ejection fraction (LVEF). However, the diagnosis of HFpEF requires more than just 'normal' LVEF. Current guidelines from the European Society of Cardiology define HFpEF as patients with a clinical syndrome of HF (with characteristic symptoms and signs), a consistent rise in natriuretic peptides and some objective evidence of diastolic impairment.⁴⁰ Each of these aspects poses particular difficulties in the context of concomitant AF. Symptoms such as dyspnoea and lethargy are common to both HFpEF and AF, and natriuretic peptides are elevated in patients with AF regardless of HF status, especially in those with persistent forms of AF. In a recent healthcare-embedded clinical trial of patients with permanent AF and dyspnoea (New York Heart Association [NYHA] class II or above), the median N-terminal pro B-type natriuretic peptide (NT-proBNP) concentration was 1057 pg/ml (interquartile range 744–1522 pg/ml), a magnitude higher than the usual cut-off point used to exclude HF.⁴¹

Documenting diastolic dysfunction using cardiac imaging is also challenging when AF is present, and there is limited information about what measurement and what value should be used in these patients.⁴² The current guideline-suggested practice of averaging multiple sequential beats in AF to obtain a reasonable mean is not based on scientific principle. Variation between beats for the measurement of E/e' filling pressures is over 40% in AF, meaning that reproducibility is so low that we should question the value of such measurements.⁴³ In contrast, the index beat approach selects appropriate cardiac cycles for measurement, thereby addressing beat-to-beat variation in AF. In a blinded study, this physiology-based approach was more reliable (coefficient of variation reduced to 25% for E/e') and more efficient in terms of echocardiographer time.⁴³

The diagnosis of AF benefits from many new forms of rhythm monitoring, including consumer electronic devices such as smartwatches. However, the validation of these devices remains unclear, and the AF they describe may not be associated with the same degree of impact on stroke and thromboembolism.^{44,45} Incident cardiac and cerebral event rates may also be dependent on the 'severity' of HFpEF and the degree of underlying systolic impairment in AF, even if above an LVEF of 50%.⁴⁶ The value of additional physiological assessment remains unclear (e.g. exercise echocardiography, right heart catheterisation or detailed assessment of atrial function), although these tests can be valuable in particular cases where the HFpEF diagnosis is uncertain. Atrial-specific biomarkers and novel molecular imaging technologies may provide more tangible benefit

in the future.⁴⁷

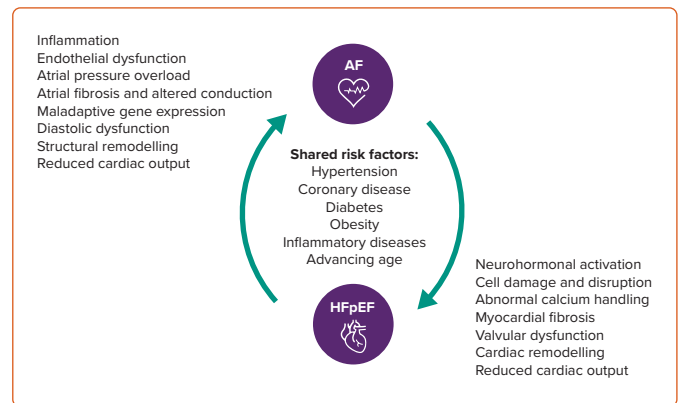
Distinguishing the relative impact of HFpEF and AF on patient symptoms when both conditions are present is also very challenging. This can impair the ability of clinicians to use focused treatments to improve patient quality of life, especially in the presence of comorbidities.³⁸ In some cases, the response to therapy should be evaluated; for example, how well diuretics relieve the dyspnoea and congestion of HFpEF, and perhaps subsequently decrease sympathetic drive and heart rate.¹ Conversely (and where feasible), it may be useful to perform electrical or pharmacological cardioversion to assess the impact of AF rhythm on current HF symptoms, albeit that sinus rhythm may be short lasting. These approaches can help identify patients with HFpEF and AF who may benefit from additional intervention, including advanced HF therapy and AF ablation. Assessment of health-related quality of life can be performed using various tools, including generic assessments such as the EQ-5D and 36-Item Short Form Health Survey (SF-36) or disease-specific questionnaires such as Atrial Fibrillation Effect on Quality-of-Life.⁴¹ For HF, lower scores with the Kansas City Cardiomyopathy Questionnaire have been associated with higher all-cause death and HF hospitalisation in both HFrEF and HFpEF.⁴⁸

Treatment Paradigm in Patients with Heart Failure With Preserved Ejection Fraction and AF

Patients with both HFpEF and AF require a different approach to management that encompasses the key elements of management for each condition, but also respects the interconnected nature of both and their combined effect on therapeutic efficacy. A patient-centred, shared management approach is essential,⁴⁹ focused on the key outcomes of importance to that individual patient, rather than esoteric outcomes taken from clinical research studies. As patients with concomitant HFpEF and AF are often older, more comorbid and already dealing with polypharmacy, it may be more relevant to focus on aspects that improve quality of life. In contrast, some patients will give a clear steer about their desire for prognostic improvement. Whichever approach is prioritised, feedback about progress can inform future clinical decisions, and tools such as quality of life questionnaires can be helpful in evaluating effectiveness and residual impairment.³⁸

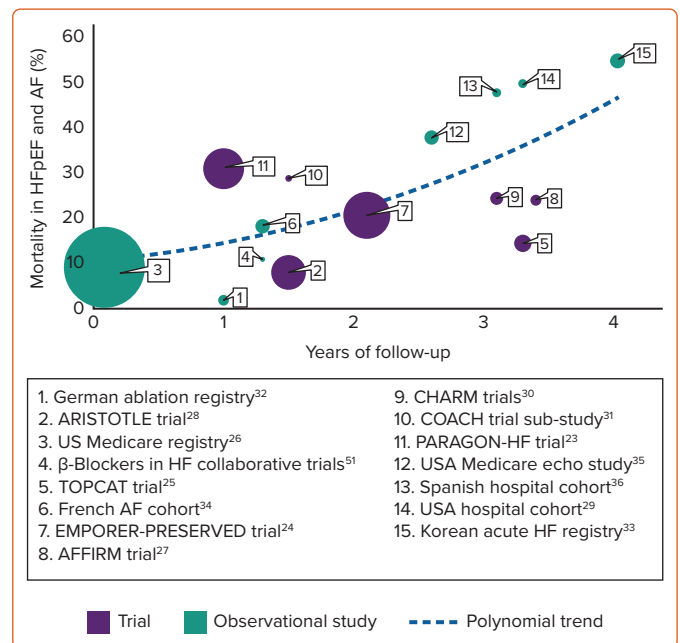
Key steps in the management of patients with HF and AF are presented in *Figure 3* (the CAN-TREAT algorithm).^{3,50} Most treatment steps are similar regardless of the LVEF of the individual patient, reflecting the need to ensure haemodynamic stability first and foremost, more widespread use of anticoagulation to prevent thromboembolism and achieving euvoalaemia. More specific approaches for rate control, heart failure therapy and rhythm control then diverge for patients with HFpEF compared with HFrEF. The intermediate group, also known as HF with mildly reduced ejection fraction, should be treated as though they have HFrEF due to the consistent evidence that they benefit from HFrEF treatments.^{51,52} An often-neglected component of the care of patients with HF and AF is to carefully and systematically address comorbidities; not just hypertension and myocardial ischaemia, but also non-cardiovascular diseases. This requires an integrated approach to achieve the best outcomes, not only between HF and AF clinical teams, but also the spectrum of healthcare professionals. Finally, to achieve the best outcomes, the conventional 'sequential' management approach should be discontinued. Relevant components of the treatment algorithm can be started in parallel; for example, starting new therapies without waiting to fully uptitrate prior drugs. Such an approach has already been advocated for HFrEF.⁵³

Figure 1: Mechanisms and Inter-relationship of Heart Failure With Preserved Ejection Fraction and AF



HFpEF = heart failure with preserved ejection fraction.

Figure 2: All-cause Mortality in Patients with Heart Failure With Preserved Ejection Fraction and AF

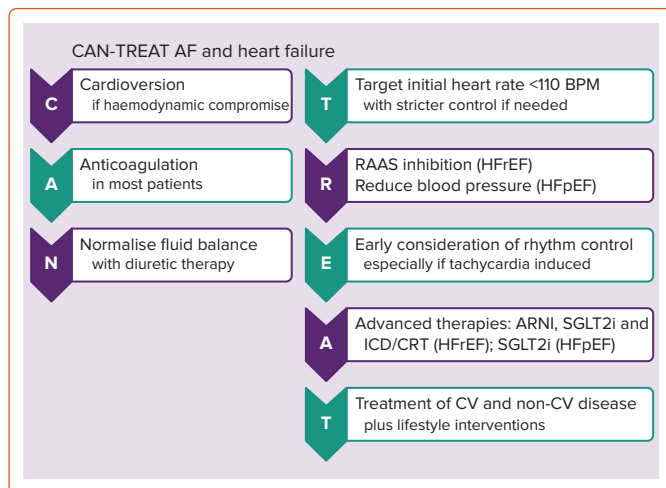


Mortality rates extracted from trials and observational studies assessing patients with concurrent HFpEF and AF. HFpEF = heart failure with preserved ejection fraction. Source: Solomon et al. 2019,²³ Packer et al. 2021,²⁴ Pitt et al. 2014,²⁵ Eapen et al. 2014,²⁶ Badheka et al. 2011,²⁷ McMurray et al. 2013,²⁸ Parkash et al. 2005,²⁹ Olsson et al. 2006,³⁰ Linssen et al. 2011,³¹ Eitel et al. 2019,³² Son et al. 2020,³³ Banerjee et al. 2012,³⁴ Pai et al. 2007,³⁵ Grigorian et al. 2006,³⁶ Cleland et al. 2018.⁵¹

Prevention of Thromboembolism

Anticoagulation is one of the only therapeutic approaches in AF with clear and proven ability to improve prognosis.⁵⁴ Although there are no specific trials in HFpEF, post hoc analysis of the four major trials of direct oral anticoagulants (DOACs) showed similar efficacy in those with HF. Compared with warfarin, DOACs reduced the risk of stroke and systemic embolisms in HF patients by 14%, with a 24% lower risk of major bleeding.⁵⁵ Therefore, except in the case of patients with severe mitral stenosis, mechanical valve prosthesis or end-stage renal dysfunction, DOACs are the first-line approach for the prevention of thromboembolism in HFpEF and AF. The place of percutaneous left atrial appendage closure is unclear due to the lack of trials against DOAC therapy. Current guidelines indicate the use of percutaneous left atrial appendage closure only in cases with

Figure 3: Update to the CAN-TREAT Management Paradigm in Patients with AF and Heart Failure



ARNI = angiotensin receptor–neprilysin inhibitor; CRT = cardiac resynchronisation therapy; CV = cardiovascular; HFpEF = heart failure with preserved ejection fraction; HFrEF = heart failure with reduced ejection fraction; RAAS = renin–angiotensin–aldosterone system; SGLT2i = sodium–glucose cotransporter 2 inhibitor. Source: adapted from Kotecha and Piccini 2015.³ Used with permission from Oxford University Press under a Creative Commons CC BY NC 4.0 licence.

an absolute contraindication for DOAC therapy (e.g. intracranial bleed without a reversible cause).⁵⁶ Where available, thoracoscopic left atrial appendage clipping is also an option in this patient group.⁵⁷

Therapies Targeting the Heart Failure With Preserved Ejection Fraction Component

Attention to fluid balance and diuretic dose is crucial, and euvoalaemia should be achieved as a priority to avoid driving tachycardia and neurohormonal activation. The use of treatments with proven benefit in HFrEF have shown disappointing results in clinical trials for patients with HFpEF. However, few of these studies have had sufficient numbers of patients with concomitant AF to be certain about either benefit or harm (Supplementary Table 1). This applies to angiotensin-converting enzyme inhibitors, angiotensin receptor blockers, β -blockers and mineralocorticoid receptor antagonists.^{51,58,60–63} Specifically, the IMPRESS-AF trial randomised 250 patients with AF and HFpEF (LVEF $\geq 55\%$) to spironolactone or placebo and disappointingly identified no benefit with regard to peak oxygen consumption on cardiopulmonary exercise testing at 2 years or secondary endpoints such as 6-min walk distance, the E/e' ratio or quality of life.⁶⁴ All participants had controlled blood pressure, so there remains potential for these therapies in the context of hypertension, as discussed later.^{59,60}

With regard to newer HF treatments, sacubitril/valsartan did not improve the composite of cardiovascular death and HF hospitalisation in patients with LVEF $\geq 45\%$ (rate ratio 0.87; 95% CI [0.75–1.01]; $p=0.06$), and the effect in those with AF was also not significant (rate ratio 0.83; 95% CI [0.69–1.00]).²³ Interaction analysis suggested those with a history of AF experienced a 13% reduction in primary event rate with sacubitril/valsartan compared with valsartan, but this ranged from a 37% greater benefit to a 21% weaker effect.²³ Results from the EMPEROR-PRESERVED trial provide the first suggestion of hope for the treatment of HFpEF.⁶⁵ Empagliflozin reduced the risk of cardiovascular death or HF hospitalisation (HR 0.79; 95% CI [0.69–0.90]; $p<0.001$) and the effects were maintained in those with AF at baseline (HR 0.78; 95% [0.66–0.93]).²⁴ Interpretation of that trial should consider that the LVEF criterion was $\geq 40\%$. There appeared to be

a 'dose' relationship with LVEF (more effect at lower LVEF), although the effect of empagliflozin remained significant in the subgroup of patients with LVEF 50–60% (HR 0.80; 95% CI [0.64–0.99]). These results suggest that sodium–glucose cotransporter 2 (SGLT2) inhibitors should become standard of care in both HFrEF and HFpEF, with an important role to play in patients with concurrent AF.

Therapies Targeting the AF Component

Heart rate control is often required in the management of patients with HFpEF and AF; however, care should be taken to avoid inducing bradycardia because chronotropic incompetence is common in elderly patients.⁶⁶ The optimal heart rate range is not known (and likely varies for individual patients), but trial evidence suggests that strict control is not beneficial and may even increase the need for hospitalisation.⁶⁷ The close linear association between heart rate and mortality seen in those with HF and sinus rhythm was not demonstrated in patients with concomitant AF.⁶⁸ The choice of rate control agent is between β -blockers (most commonly used), digoxin or calcium channel blockers, such as diltiazem or verapamil. Amiodarone should not be used as a rate control agent due to its non-cardiac side effects. Comparing β -blockers versus low-dose digoxin, the RATE-AF trial randomised 160 patients with permanent AF and symptoms of HF (NYHA class II or above; 81% with LVEF $\geq 50\%$).⁴¹ The trial found no difference in the physical component of quality of life, but there were significant benefits from digoxin on AF symptoms (two-class improvement in 53% of patients versus in 9% of patients with β -blockers; $p<0.001$), NYHA class (mean 2.4, decreasing to 1.5 at 12 months, versus 2.0 with β -blockers; $p<0.001$), and NT-proBNP (960 pg/ml at 12 months versus 1250 pg/ml with β -blockers; $p=0.005$).⁴¹ There were also substantially fewer adverse events with low-dose digoxin (25% of patients with at least one event versus 64% with β -blockers; $p<0.001$). Calcium channel blockers are another option for rate control in those with normal LVEF, but their benefit on exercise capacity and NT-proBNP compared with β -blockers has only been established in a small cross-over trial of patients without HF.⁶⁹ Ablation of the atrioventricular node is an option when other attempts to control heart rate fail, but leads to pacemaker dependency.

For those with ongoing symptoms despite good heart rate control, rhythm control should be considered, balancing the risk of rhythm control approaches (anti-arrhythmic drugs, such as amiodarone and dronedarone, and catheter ablation) against the potential for long-term benefit. This balance is often challenging in patients with established HFpEF and AF because, due to multimorbidity, there may be a lower chance for maintenance of sinus rhythm. Clinical trials are lacking in this patient group, and observational data are not helpful due to the considerable selection and performance biases.⁷⁰ Consideration of early rhythm control, for example within the first year of AF, merits separate attention. In the EAST trial, which randomised patients to early or conventional rhythm control, the subgroup of 798 patients with HF had a similar reduction as non-HF patients for the composite of cardiovascular death, stroke and hospitalisation for HF or acute coronary syndrome (HR 0.74 with an early approach; 95% CI [0.56–0.97]; $p=0.03$).⁷¹ There were insufficient patients to be certain about benefit in those specifically with HFpEF ($p=0.24$) but, pending further trial evidence, an early rhythm approach should be considered, particularly where HFpEF may be a consequence of AF-related irregularity or elevated heart rate (e.g. in tachycardia-related cardiomyopathy). Data specifically on HFpEF are not available, although studies in HFrEF would suggest clinical benefit from AF ablation.

Lifestyle Interventions and Comorbidity Management

Lifestyle changes should be suggested in all patients with HFpEF and AF where relevant to that individual patient; this will enable the patient to take an active role in their management alongside medical therapy. Certain interventions have also demonstrated improvements in patient well-being, although there remains uncertainty about their impact on clinical endpoints.⁷² For example, in a meta-analysis of six small randomised trials (total of 276 patients with HFpEF), exercise training improved peak oxygen uptake (weighted mean difference 2.72 ml/kg/min; 95% CI [1.79–3.65 ml/kg/min]) and quality of life (weighted mean difference –3.97; 95% CI [–7.21, –0.72]).⁷³ In a factorial trial of 100 obese patients with HFpEF, the addition of a low-calorie diet to an exercise regime resulted in further improvements to peak VO₂ (1.2, 1.3 and 2.5 ml/kg/min for exercise, diet and both together, respectively).⁷⁴ The effects of these approaches in patients with combined HFpEF and AF is not yet known.

Adequate control of blood pressure is often asserted as an essential component of HFpEF management, although there remains little trial evidence for an impact on clinical endpoints once HFpEF is established. The aforementioned trials of renin–angiotensin–aldosterone system antagonists (*Supplementary Table 1*) all demonstrate a significant reduction in blood pressure, so these are suitable as first-line agents until the place of SGLT2 inhibitors becomes clearer. With regard to AF, an individual patient-level meta-analysis comprising 22 trials showed that blood pressure-lowering treatment reduces the risk of major cardiovascular events to a similar extent in individuals with and without AF (13,266 and 175,304 participants, respectively).⁷⁵ Each 5-mmHg reduction

in blood pressure lowered the risk of stroke, ischaemic heart disease or HF by 9% during a 4.5-year follow-up. The target for blood pressure control in either HFpEF or AF remains the subject of debate.⁷⁶

Other important comorbidities whose management should be prioritised in the context of HFpEF with AF are underlying ischaemic heart disease, obesity, iron deficiency and glycaemic control in patients with diabetes. Depression is common in HF patients, and although therapies can improve quality of life, the impact on clinical outcomes remains uncertain.⁷⁷ Non-cardiovascular comorbidities, such as pulmonary disease, have a worse impact on mortality in HFpEF than HFrEF, whereas gout and cancer have a similar effect in both HF phenotypes.⁷⁸ For AF, targeted therapy of underlying conditions in a randomised trial led to significant improvements in the maintenance of sinus rhythm.⁷⁹ In HFpEF and AF, mortality and morbidity are frequently from non-cardiovascular causes, and so the management of patients is incomplete without the systematic, individualised assessment and treatment of comorbidities.⁸⁰

Conclusion

HFpEF and AF are increasingly prevalent and, when combined, lead to a substantial increase in mortality and poor patient quality of life. Diagnosis is challenging and conventional therapy is often unable to improve clinical endpoints. A paradigm shift is needed in clinical management that considers the joint effects of both conditions in order to adequately treat patients. This approach should be personalised, use non-sequential treatment prescription targeted to both HFpEF and AF components and integrate attention on underlying comorbidities to prevent progression. □

- Kotecha D, Lam CS, Van Veldhuisen DJ, et al. Heart failure with preserved ejection fraction and atrial fibrillation: vicious twins. *J Am Coll Cardiol* 2016;68:2217–28. <https://doi.org/10.1016/j.jacc.2016.08.048>; PMID: 27855811.
- Vermond RA, Geelhoed B, Verweij N, et al. Incidence of atrial fibrillation and relationship with cardiovascular events, heart failure, and mortality: a community-based study from the Netherlands. *J Am Coll Cardiol* 2015;66:1000–7. <https://doi.org/10.1016/j.jacc.2015.06.1314>; PMID: 26314526.
- Kotecha D, Piccini JP. Atrial fibrillation in heart failure: what should we do? *Eur Heart J* 2015;36:3250–7. <https://doi.org/10.1093/eurheartj/ehv513>; PMID: 26419625.
- Rozen G, Hosseini SM, Kaadan MI, et al. Emergency department visits for atrial fibrillation in the United States: trends in admission rates and economic burden from 2007 to 2014. *J Am Heart Assoc* 2018;7:e009024. <https://doi.org/10.1161/JAHA.118.009024>; PMID: 30030215.
- JE H, Enserro D, Brouwers FP, et al. Predicting heart failure with preserved and reduced ejection fraction: the international collaboration on heart failure subtypes. *Circ Heart Fail* 2016;9:e003116. <https://doi.org/10.1161/CIRCHEARTFAILURE.115.003116>; PMID: 27266854.
- Pfeffer MA, Shah AM, Borlaug BA. Heart failure with preserved ejection fraction in perspective. *Circ Res* 2019;124:1598–617. <https://doi.org/10.1161/CIRCRESAHA.119.313572>; PMID: 31120821.
- Urbich M, Globe G, Pantiri K, et al. A systematic review of medical costs associated with heart failure in the USA (2014–2020). *Pharmacoeconomics* 2020;38:1219–36. <https://doi.org/10.1007/s40273-020-00952-0>; PMID: 32812149.
- Chang PP, Wruck LM, Shahar E, et al. Trends in hospitalizations and survival of acute decompensated heart failure in four US communities (2005–2014): ARIC study community surveillance. *Circulation* 2018;138:12–24. <https://doi.org/10.1161/CIRCULATIONAHA.117.027551>; PMID: 29519849.
- Lane DA, Skjøth F, Lip GYH, et al. Temporal trends in incidence, prevalence, and mortality of atrial fibrillation in primary care. *J Am Heart Assoc* 2017;6:e005155. <https://doi.org/10.1161/JAHA.116.005155>; PMID: 28455344.
- Conen D, Rodondi N, Muller A, et al. Relationships of overt and silent brain lesions with cognitive function in patients with atrial fibrillation. *J Am Coll Cardiol* 2019;73:989–99. <https://doi.org/10.1016/j.jacc.2018.12.039>; PMID: 30846109.
- Groenewegen A, Rutten FH, Mosterd A, Hoes AW. Epidemiology of heart failure. *Eur J Heart Fail* 2020;22:1342–56. <https://doi.org/10.1002/ehfj.1858>; PMID: 32483830.
- Maggioni AP, Dahlström U, Filippatos G, et al. EURObservational research programme: the heart failure pilot survey (ESC-HF pilot). *Eur J Heart Fail* 2010;12:1076–84. <https://doi.org/10.1093/eurjhf/hq154>; PMID: 20805094.
- Groeneweld HF, Tijssen JG, Crijns HJ, et al. Rate control efficacy in permanent atrial fibrillation: successful and failed strict rate control against a background of lenient rate control: data from RACE II (Rate Control Efficacy in permanent atrial fibrillation). *J Am Coll Cardiol* 2013;61:741–8. <https://doi.org/10.1016/j.jacc.2012.11.038>; PMID: 23410544.
- Zafir B, Lund LH, Laroche C, et al. Prognostic implications of atrial fibrillation in heart failure with reduced, mid-range, and preserved ejection fraction: a report from 14 964 patients in the European Society of Cardiology Heart Failure Long-Term Registry. *Eur Heart J* 2018;39:4277–84. <https://doi.org/10.1093/eurheartj/ehy626>; PMID: 30325423.
- Zakeri R, Chamberlain AM, Roger VL, Redfield MM. Temporal relationship and prognostic significance of atrial fibrillation in heart failure patients with preserved ejection fraction: a community-based study. *Circulation* 2013;128:1085–93. <https://doi.org/10.1161/CIRCULATIONAHA.113.001475>; PMID: 23908348.
- Santhanakrishnan R, Wang N, Larson MG, et al. Atrial fibrillation begets heart failure and vice versa: temporal associations and differences in preserved versus reduced ejection fraction. *Circulation* 2016;133:484–92. <https://doi.org/10.1161/CIRCULATIONAHA.115.018614>; PMID: 26746177.
- Franssen C, Chen S, Unger A, et al. Myocardial microvascular inflammatory endothelial activation in heart failure with preserved ejection fraction. *JACC Heart Fail* 2016;4:312–24. <https://doi.org/10.1016/j.jchf.2015.10.007>; PMID: 26682792.
- Conrad N, Judge A, Tran J, et al. Temporal trends and patterns in heart failure incidence: a population-based study of 4 million individuals. *Lancet* 2018;391:572–80. [https://doi.org/10.1016/S0140-6736\(17\)32520-5](https://doi.org/10.1016/S0140-6736(17)32520-5); PMID: 29174292.
- Carlisle MA, Fudim M, DeVore AD, Piccini JP. Heart failure and atrial fibrillation, like fire and fury. *JACC Heart Fail* 2019;7:447–56. <https://doi.org/10.1016/j.jchf.2019.03.005>; PMID: 31146871.
- Packer M, Lam CSP, Lund LH, Redfield MM. Interdependence of atrial fibrillation and heart failure with a preserved ejection fraction reflects a common underlying atrial and ventricular myopathy. *Circulation* 2020;141:4–6. <https://doi.org/10.1161/CIRCULATIONAHA.119.042996>; PMID: 31887078.
- Sartipy U, Dahlström U, Fu M, Lund LH. Atrial fibrillation in heart failure with preserved, mid-range, and reduced ejection fraction. *JACC Heart Fail* 2017;5:565–74. <https://doi.org/10.1016/j.jchf.2017.05.001>; PMID: 28711451.
- O'Neal WT, Sandesara P, Hammadah M, et al. Gender differences in the risk of adverse outcomes in patients with atrial fibrillation and heart failure with preserved ejection fraction. *Am J Cardiol* 2017;119:1785–90. <https://doi.org/10.1016/j.amjcard.2017.02.045>; PMID: 28395886.
- Solomon SD, McMurray JJV, Anand IS, et al. Angiotensin–neprilysin inhibition in heart failure with preserved ejection fraction. *N Engl J Med* 2019;381:1609–20. <https://doi.org/10.1056/NEJMoa1908655>; PMID: 31475794.
- Packer M, Butler J, Zannad F, et al. Effect of empagliflozin on worsening heart failure events in patients with heart failure and preserved ejection fraction: EMPEROR-preserved trial. *Circulation* 2021;144:1284–94. <https://doi.org/10.1161/CIRCULATIONAHA.121.056824>; PMID: 34459213.
- Pitt B, Pfeffer MA, Assmann SF, et al. Spironolactone for heart failure with preserved ejection fraction. *N Engl J Med* 2014;370:1383–92. <https://doi.org/10.1056/NEJMoa1313731>; PMID: 24716680.
- Eapen ZJ, Greiner MA, Fonarow GC, et al. Associations between atrial fibrillation and early outcomes of patients with heart failure and reduced or preserved ejection fraction. *Am Heart J* 2014;167:369–75.e362. <https://doi.org/10.1016/j.ahj.2013.12.001>; PMID: 24576522.
- Badheka AO, Rathod A, Kizilbash MA, et al. Comparison of mortality and morbidity in patients with atrial fibrillation and heart failure with preserved versus decreased left ventricular ejection fraction. *Am J Cardiol* 2011;108:1283–8. <https://doi.org/10.1016/j.amjcard.2011.06.045>; PMID: 21855829.
- McMurray JJ, Ezekowitz JA, Lewis BS, et al. Left ventricular systolic dysfunction, heart failure, and the risk of stroke and systemic embolism in patients with atrial fibrillation: insights from the Aristotle trial. *Circ Heart Fail* 2013;6:451–60. <https://doi.org/10.1161/CIRCHEARTFAILURE.112.000143>; PMID: 23575255.
- Parkash R, Maisel WH, Toca FM, Stevenson WG. Atrial fibrillation in heart failure: high mortality risk even if ventricular function is preserved. *Am Heart J* 2005;150:701–6. <https://doi.org/10.1016/j.ahj.2004.12.014>; PMID: 16209969.
- Olsson LG, Swedberg K, Ducharme A, et al. Atrial fibrillation and risk of clinical events in chronic heart failure with and without left ventricular systolic dysfunction: results from the Candesartan in Heart failure-Assessment of Reduction in Mortality and Morbidity (CHARM) Program. *J Am Coll Cardiol* 2006;47:1997–2004. <https://doi.org/10.1016/j.jacc.2006.01.060>; PMID: 16697316.
- Linssen GC, Rienstra M, Jaarsma T, et al. Clinical and prognostic effects of atrial fibrillation in heart failure patients with reduced and preserved left ventricular ejection fraction. *Eur J Heart Fail* 2011;13:1111–20. <https://doi.org/10.1093/eurjhf/hfr066>; PMID: 21642293.
- Eitel C, Ince H, Brachmann J, et al. Atrial fibrillation ablation strategies and outcome in patients with heart failure: insights from the German ablation registry. *Clin Res Cardiol* 2019;108:815–23. <https://doi.org/10.1007/s00392-019-01411-3>; PMID: 30788620.

33. Son MK, Park JJ, Lim NK, et al. Impact of atrial fibrillation in patients with heart failure and reduced, mid-range or preserved ejection fraction. *Heart* 2020;106:1160–8. <https://doi.org/10.1136/heartjnl-2019-316219>; PMID: 32341140.
34. Banerjee A, Taillandier S, Olesen JB, et al. Ejection fraction and outcomes in patients with atrial fibrillation and heart failure: the Loire Valley Atrial Fibrillation Project. *Eur J Heart Fail* 2012;14:295–301. <https://doi.org/10.1093/eurjhf/hfs005>; PMID: 22294759.
35. Pai RG, Varadarajan P. Prognostic significance of atrial fibrillation is a function of left ventricular ejection fraction. *Clin Cardiol* 2007;30:349–54. <https://doi.org/10.1002/clc.20107>; PMID: 17674374.
36. Grigorian Shamagian L, Roman AV, Seara JG, et al. Atrial fibrillation in patients hospitalized for congestive heart failure: the same prognostic influence independently of left ventricular systolic function? *Int J Cardiol* 2006;110:366–72. <https://doi.org/10.1016/j.ijcard.2005.08.022>; PMID: 16297467.
37. Kotecha D, Chudasama R, Lane DA, et al. Atrial fibrillation and heart failure due to reduced versus preserved ejection fraction: a systematic review and meta-analysis of death and adverse outcomes. *Int J Cardiol* 2016;203:660–6. <https://doi.org/10.1016/j.ijcard.2015.10.220>; PMID: 26580351.
38. Jones J, Stanbury M, Haynes S, et al. Importance and assessment of quality of life in symptomatic permanent atrial fibrillation: patient focus groups from the RATE-AF trial. *Cardiology* 2020;145:666–75. <https://doi.org/10.1159/000511048>; PMID: 32862174.
39. Sepelhrvand N, Savu A, Spertus JA, et al. Change of health-related quality of life over time and its association with patient outcomes in patients with heart failure. *J Am Heart Assoc* 2020;9:e017278. <https://doi.org/10.1161/JAHA.120.017278>; PMID: 32812460.
40. McDonagh TA, Metra M, Adamo M, et al. 2021 ESC guidelines for the diagnosis and treatment of acute and chronic heart failure. *Eur Heart J* 2021;42:3599–726. <https://doi.org/10.1093/eurheartj/ehab368>; PMID: 34447992.
41. Kotecha D, Bunting KV, Gill SK, et al. Effect of digoxin vs bisoprolol for heart rate control in atrial fibrillation on patient-reported quality of life: the RATE-AF randomized clinical trial. *JAMA* 2020;324:2497–508. <https://doi.org/10.1001/jama.2020.23138>; PMID: 33351042.
42. Bunting KV, O'Connor K, Steeds RP, Kotecha D. Cardiac imaging to assess left ventricular systolic function in atrial fibrillation. *Am J Cardiol* 2021;139:40–9. <https://doi.org/10.1016/j.amjcard.2020.10.012>; PMID: 33065079.
43. Bunting KV, Gill SK, Stith A, et al. Improving the diagnosis of heart failure in patients with atrial fibrillation. *Heart* 2021;107:902–8. <https://doi.org/10.1136/heartjnl-2020-318557>; PMID: 33692093.
44. Gill SK, Bunting KV, Sartini C, et al. Smartphone detection of atrial fibrillation using photoplethysmography: a systematic review and meta-analysis. *Heart* 2022. <https://doi.org/10.1136/heartjnl-2021-320417>; PMID: 35277454; epub ahead of press.
45. Kotecha D, Breithardt G, Camm AJ, et al. Integrating new approaches to atrial fibrillation management: the 6th AFNET/EHRA consensus conference. *Europace* 2018;20:395–407. <https://doi.org/10.1093/eurheartj/ehx318>; PMID: 29300976.
46. Siller-Matula JM, Pecan L, Patti G, et al. Heart failure subtypes and thromboembolic risk in patients with atrial fibrillation: the PREFER in AF–HF substudy. *Int J Cardiol* 2018;265:141–7. <https://doi.org/10.1016/j.ijcard.2018.04.093>; PMID: 29706429.
47. Coats AJS, Heymans S, Farmakis D, et al. Atrial disease and heart failure: the common soil hypothesis proposed by the Heart Failure Association of the European Society of Cardiology. *Eur Heart J* 2022;43:863–7. <https://doi.org/10.1093/eurheartj/ehab834>; PMID: 34875053.
48. Johansson I, Joseph P, Balasubramanian K, et al. Health-related quality of life and mortality in heart failure: the global congestive heart failure study of 23 000 patients from 40 countries. *Circulation* 2021;143:2129–42. <https://doi.org/10.1161/CIRCULATIONAHA.120.050850>; PMID: 33906372.
49. Kotecha D, Chua WWL, Fabritz L, et al. European Society of Cardiology smartphone and tablet applications for patients with atrial fibrillation and their health care providers. *Europace* 2018;20:225–33. <https://doi.org/10.1093/eurheartj/ehx299>; PMID: 29040548.
50. Kotecha D, Lainscak M. Co-morbidity (HFpEF and HFpEF): atrial fibrillation. In: Camm A, Lüscher T, Maurer G, et al., eds. *The ESC Textbook of Cardiovascular Medicine*. 3rd ed. Oxford, UK: Oxford University Press; 2018; 1798–801. <https://doi.org/10.1093/med/9780198784906.003.0413>.
51. Cleland JGF, Bunting KV, Flather MD, et al. Beta-blockers for heart failure with reduced, mid-range, and preserved ejection fraction: an individual patient-level analysis of double-blind randomized trials. *Eur Heart J* 2018;39:26–35. <https://doi.org/10.1093/eurheartj/ehx564>; PMID: 29040525.
52. Lund LH, Claggett B, Liu J, et al. Heart failure with mid-range ejection fraction in CHARM: characteristics, outcomes and effect of candesartan across the entire ejection fraction spectrum. *Eur J Heart Fail* 2018;20:1230–9. <https://doi.org/10.1002/ehfj.1149>; PMID: 29431256.
53. McMurray JJV, Packer M. How should we sequence the treatments for heart failure and a reduced ejection fraction? A redefinition of evidence-based medicine. *Circulation* 2021;143:875–7. <https://doi.org/10.1161/CIRCULATIONAHA.120.052926>; PMID: 33378214.
54. Kotecha D, Pollack CV, Jr, De Caterina R, et al. Direct oral anticoagulants halve thromboembolic events after cardioversion of AF compared with warfarin. *J Am Coll Cardiol* 2018;72:1984–6. <https://doi.org/10.1016/j.jacc.2018.07.083>; PMID: 30309478.
55. Xiong Q, Lau YC, Senoo K, et al. Non-vitamin K antagonist oral anticoagulants (NOACs) in patients with concomitant atrial fibrillation and heart failure: a systematic review and meta-analysis of randomized trials. *Eur J Heart Fail* 2015;17:1192–200. <https://doi.org/10.1002/ehfj.343>; PMID: 26335355.
56. Hindricks G, Potpara T, Dagres N, et al. 2020 ESC guidelines for the diagnosis and management of atrial fibrillation developed in collaboration with the European Association for Cardio-Thoracic Surgery (EACTS). *Eur Heart J* 2021;42:373–498. <https://doi.org/10.1093/eurheartj/ehaa612>; PMID: 32860505.
57. van Laar C, Verberkmoes NJ, van Es HW, et al. Thoracoscopic left atrial appendage clipping: a multicenter cohort analysis. *JACC Clin Electrophysiol* 2018;4:893–901. <https://doi.org/10.1016/j.jacep.2018.03.009>; PMID: 30025689.
58. Cleland JG, Tendera M, Adamus J, et al. The perindopril in elderly people with chronic heart failure (PEP-CHF) study. *Eur Heart J* 2006;27:2338–45. <https://doi.org/10.1093/eurheartj/ehl250>; PMID: 16963472.
59. Massie BM, Carson PE, McMurray JJ, et al. Irbesartan in patients with heart failure and preserved ejection fraction. *N Engl J Med* 2008;359:2456–67. <https://doi.org/10.1056/NEJMoa0805450>; PMID: 19001508.
60. Yusuf S, Pfeffer MA, Swedberg K, et al. Effects of candesartan in patients with chronic heart failure and preserved left-ventricular ejection fraction: the CHARM-preserved trial. *Lancet* 2003;362:777–81. [https://doi.org/10.1016/S0140-6736\(03\)14285-7](https://doi.org/10.1016/S0140-6736(03)14285-7); PMID: 13678871.
61. Yamamoto K, Origasa H, Hori M, J-DHF Investigators. Effects of carvedilol on heart failure with preserved ejection fraction: the Japanese Diastolic Heart Failure Study (J-DHF). *Eur J Heart Fail* 2013;15:110–8. <https://doi.org/10.1093/eurjhf/hfs141>; PMID: 22983988.
62. Cikes M, Claggett B, Shah AM, et al. Atrial fibrillation in heart failure with preserved ejection fraction: the TOPCAT trial. *JACC Heart Fail* 2018;6:689–97. <https://doi.org/10.1016/j.jchf.2018.05.005>; PMID: 30007557.
63. Kotecha D, Holmes J, Krum H, et al. Efficacy of beta blockers in patients with heart failure plus atrial fibrillation: an individual-patient data meta-analysis. *Lancet* 2014;384:2235–43. [https://doi.org/10.1016/S0140-6736\(14\)61373-8](https://doi.org/10.1016/S0140-6736(14)61373-8); PMID: 25193873.
64. Shantsila E, Shahid F, Sun Y, et al. Spironolactone in atrial fibrillation with preserved cardiac fraction: the IMPRESS-AF trial. *J Am Heart Assoc* 2020;9:e016239. <https://doi.org/10.1161/JAHA.119.016239>; PMID: 32909497.
65. Anker SD, Butler J, Filippatos G, et al. Empagliflozin in heart failure with a preserved ejection fraction. *N Engl J Med* 2021;385:1451–61. <https://doi.org/10.1056/NEJMoa2107038>; PMID: 34449189.
66. Kotecha D, Calvert M, Deeks JJ, et al. A review of rate control in atrial fibrillation, and the rationale and protocol for the RATE-AF trial. *BMJ Open* 2017;7:e015099. <https://doi.org/10.1136/bmjopen-2016-015099>; PMID: 28729311.
67. Van Gelder IC, Groeneweld HF, Crijns HJGM, et al. Lenient versus strict rate control in patients with atrial fibrillation. *N Engl J Med* 2010;362:1363–73. <https://doi.org/10.1056/NEJMoa1001337>; PMID: 20231232.
68. Kotecha D, Flather MD, Altman DG, et al. Heart rate and rhythm and the benefit of beta-blockers in patients with heart failure. *J Am Coll Cardiol* 2017;69:2885–96. <https://doi.org/10.1016/j.jacc.2017.04.001>; PMID: 28467883.
69. Ullmoen SR, Enger S, Pripp AH, et al. Calcium channel blockers improve exercise capacity and reduce N-terminal pro-B-type natriuretic peptide levels compared with beta-blockers in patients with permanent atrial fibrillation. *Eur Heart J* 2014;35:517–24. <https://doi.org/10.1093/eurheartj/ehx429>; PMID: 24135831.
70. Mulder BA, Rienstra M, Van Gelder IC, Blaauw Y. Update on management of atrial fibrillation in heart failure: a focus on ablation. *Heart* 2022;108:422–8. <https://doi.org/10.1136/heartjnl-2020-318081>; PMID: 34088767.
71. Rillig A, Magnussen C, Ozga AK, et al. Early rhythm control therapy in patients with atrial fibrillation and heart failure. *Circulation* 2021;144:845–58. <https://doi.org/10.1161/CIRCULATIONAHA.121.056323>; PMID: 34328366.
72. Aggarwal M, Bozkurt B, Panjath G, et al. Lifestyle modifications for preventing and treating heart failure. *J Am Coll Cardiol* 2018;72:2391–405. <https://doi.org/10.1016/j.jacc.2018.08.2160>; PMID: 30384895.
73. Pandey A, Parashar A, Kumbhani D, et al. Exercise training in patients with heart failure and preserved ejection fraction: meta-analysis of randomized control trials. *Circ Heart Fail* 2015;8:33–40. <https://doi.org/10.1161/CIRCHEARTFAILURE.114.001615>; PMID: 25399909.
74. Kitzman DW, Brubaker P, Morgan T, et al. Effect of caloric restriction or aerobic exercise training on peak oxygen consumption and quality of life in obese older patients with heart failure with preserved ejection fraction: a randomized clinical trial. *JAMA* 2016;315:36–46. <https://doi.org/10.1001/jama.2015.17346>; PMID: 26746456.
75. Pinho-Gomes AC, Azevedo L, Copland E, et al. Blood pressure-lowering treatment for the prevention of cardiovascular events in patients with atrial fibrillation: an individual participant data meta-analysis. *PLOS Med* 2021;18:e1003599. <https://doi.org/10.1371/journal.pmed.1003599>; PMID: 34061831.
76. Myhre PL, Selvaraj S, Solomon SD. Management of hypertension in heart failure with preserved ejection fraction: is there a blood pressure goal? *Curr Opin Cardiol* 2021;36:413–9. <https://doi.org/10.1097/HCO.0000000000000852>; PMID: 33709982.
77. Jayanthan K, Kotecha D, Thanki D, et al. Effects of cognitive behavioural therapy for depression in heart failure patients: a systematic review and meta-analysis. *Heart Fail Rev* 2017;22:731–41. <https://doi.org/10.1007/s10741-017-9640-5>; PMID: 28733911.
78. Ergatoules C, Schaufelberger M, Andersson B, et al. Non-cardiac comorbidities and mortality in patients with heart failure with reduced vs. preserved ejection fraction: a study using the Swedish Heart Failure Registry. *Clin Res Cardiol* 2019;108:1025–33. <https://doi.org/10.1007/s00392-019-01430-0>; PMID: 30788622.
79. Rienstra M, Hobbelt AH, Alings M, et al. Targeted therapy of underlying conditions improves sinus rhythm maintenance in patients with persistent atrial fibrillation: results of the RACE 3 trial. *Eur Heart J* 2018;39:2987–96. <https://doi.org/10.1093/eurheartj/ehx739>; PMID: 29401239.
80. Tica O, Tica O, Bunting KV, deBono J, Gkoutos GV, Popescu MI, Kotecha D. Post-mortem examination of high mortality in patients with heart failure and atrial fibrillation. *BMC Med*. 2022;20:331 DOI: 10.1186/s12916-022-02533-8. PMID: 36195871.