

NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Single Technology Appraisal

Empagliflozin for treating chronic heart failure with reduced ejection fraction

Final scope

Draft remit/appraisal objective

To appraise the clinical and cost effectiveness of empagliflozin within its marketing authorisation for treating symptomatic chronic heart failure with reduced ejection fraction.

Background

Heart failure is a complex clinical syndrome of signs and symptoms, generally defined as the inability of the heart to supply sufficient blood flow to meet the body's needs. It is caused by structural or functional abnormalities of the heart, commonly resulting from coronary artery disease. Other conditions that can increase the risk of heart failure include; ischemic heart disease, atrial fibrillation, valve disease, hypertension, diabetes, chronic obstructive pulmonary disease, and asthma. Heart failure may be associated with left ventricular systolic dysfunction (that is, reduced left ventricular ejection fraction, where the left pumping chamber's ability to pump is impaired) but may also be associated with reduced ejection fraction, defined as an ejection fraction below 40% in [NICE guideline 106 for chronic heart failure in adults](#).

Symptoms of heart failure commonly include breathlessness, fatigue and ankle swelling. Quality of life is affected by the physical limitations imposed by the symptoms.

More than 550,000 people in England have heart failure¹. There were 94,185 hospitalisations in England for heart failure in 2019/20². 66% of people with heart failure have a reduced left ventricular ejection fraction³. Both the prevalence and incidence of heart failure increase with age. 30 to 40% of people diagnosed with heart failure die within the first year, with a 5-year mortality rate of about 50%⁴.

[NICE guideline 106 for chronic heart failure in adults](#) recommends offering an angiotensin-converting enzyme (ACE) inhibitor and a beta-blocker for people with heart failure with reduced ejection fraction. If ACE inhibitors are contraindicated or not tolerated, an angiotensin receptor blocker (ARB) should be considered. A mineralocorticoid receptor antagonist (MRA) in addition to an ACE inhibitor (or ARB) and beta-blocker should be offered if symptoms continue.

If neither ACE inhibitors or ARBs are tolerated, specialist advice should be sought and treatment with hydralazine in combination with nitrate can be

considered. If symptoms worsen or become severe despite first-line treatment, specialist advice should be sought and treatment with digoxin can be considered.

NICE [technology appraisal guidance 388](#) recommends sacubitril valsartan as an option for treating symptomatic chronic heart failure with reduced ejection fraction, only in people:

- with New York Heart Association (NYHA) class II to IV symptoms and
- with a left ventricular ejection fraction of 35% or less and
- who are already taking a stable dose of ACE inhibitors or ARBs

NICE [technology appraisal guidance 267](#) recommends ivabradine in combination with standard therapy for people:

- with New York Heart Association (NYHA) class II to IV stable chronic heart failure with systolic dysfunction and
- who are in sinus rhythm with a heart rate of 75 beats per minute or more and
- who are given ivabradine in combination with standard therapy including beta-blocker therapy, angiotensin-converting enzyme (ACE) inhibitors and aldosterone antagonists, or when beta-blocker therapy is contraindicated or not tolerated and
- with a left ventricular ejection fraction of 35% or less

NICE [technology appraisal guidance 679](#) recommends dapagliflozin as an option for treating symptomatic chronic heart failure with reduced ejection fraction in adults, only if it is used as an add-on to optimised standard care with:

- angiotensin-converting enzyme (ACE) inhibitors or angiotensin-2 receptor blockers (ARBs), with beta blockers, and, if tolerated, mineralocorticoid receptor antagonists (MRAs), or
- sacubitril valsartan, with beta blockers, and, if tolerated, MRAs.

The technology

Empagliflozin (Jardiance, Boehringer Ingelheim) is a sodium-glucose co-transporter 2 (SGLT-2) inhibitor. The mechanism of action of empagliflozin in heart failure with reduced ejection fraction is not yet fully understood. It is administered orally.

Empagliflozin does not currently have a marketing authorisation in the UK for chronic heart failure with reduced ejection fraction. It is being studied in

randomised controlled trials compared with placebo, in adults with an diagnosis of chronic heart (NYHA functional class II-IV) failure with left reduced ejection fraction of 40% or less.

Intervention(s)	Empagliflozin in combination with standard care (including diuretics, treatment with an ACE inhibitor, ARBs, mineralocorticoid receptor antagonist, beta blockers, cardiac devices and sacubitril valsartan)
Population(s)	Adults for the treatment of symptomatic chronic heart failure with reduced ejection fraction
Comparators	<ul style="list-style-type: none"> • Individually optimised standard care without empagliflozin. Standard care is defined as: <ul style="list-style-type: none"> ○ ACE inhibitors in combination with beta-blockers, and/or mineralocorticoid receptor antagonists ○ ARBs in combination with beta-blockers, and/or mineralocorticoid receptor antagonists ○ Sacubitril valsartan in combination with beta-blockers, and/or mineralocorticoid receptor antagonists • Dapagliflozin as an add on to standard care
Outcomes	<p>The outcome measures to be considered include:</p> <ul style="list-style-type: none"> • symptoms of heart failure • hospitalisation for heart failure • all-cause hospitalisation • mortality • cardiovascular mortality • kidney function • adverse effects of treatment • health-related quality of life

<p>Economic analysis</p>	<p>The reference case stipulates that the cost effectiveness of treatments should be expressed in terms of incremental cost per quality-adjusted life year.</p> <p>If the technology is likely to provide similar or greater health benefits at similar or lower cost than technologies recommended in published NICE technology appraisal guidance for the same indication, a cost-comparison may be carried out.</p> <p>The reference case stipulates that the time horizon for estimating clinical and cost effectiveness should be sufficiently long to reflect any differences in costs or outcomes between the technologies being compared.</p> <p>Costs will be considered from an NHS and Personal Social Services perspective.</p> <p>The availability of any commercial arrangements for the intervention, comparator and subsequent treatment technologies will be taken into account.</p> <p>The cost of background therapies, such as diuretics for people with oedema, should also be included in cost effectiveness analyses.</p>
<p>Other considerations</p>	<p>Guidance will only be issued in accordance with the marketing authorisation. Where the wording of the therapeutic indication does not include specific treatment combinations, guidance will be issued only in the context of the evidence that has underpinned the marketing authorisation granted by the regulator.</p>
<p>Related NICE recommendations and NICE Pathways</p>	<p>Related Technology Appraisals:</p> <p>Dapagliflozin for treating chronic heart failure with reduced ejection fraction (2021) NICE technology appraisal 679</p> <p>Sacubitril valsartan for treating symptomatic chronic heart failure with reduced ejection fraction (2016) NICE technology appraisal 388</p> <p>Ivabradine for treating chronic heart failure (2012) NICE technology appraisal guidance 267</p> <p>Related Guidelines:</p> <p>Chronic heart failure in adults: diagnosis and management (2018) NICE guideline NG106</p> <p>Related Quality Standards:</p> <p>Chronic heart failure in adults (2011) NICE quality standard 9</p>

	<p>Related NICE Pathways:</p> <p>Chronic heart failure (2019) NICE pathway</p>
Related National Policy	<p>The NHS Long Term Plan, 2019. NHS Long Term Plan NHS England (2018/2019) NHS manual for prescribed specialist services (2018/2019)</p> <p>Department of Health and Social Care, NHS Outcomes Framework 2016-2017: Domains 1,2. https://www.gov.uk/government/publications/nhs-outcomes-framework-2016-to-2017</p>

References

1. British Heart Foundation. Heart statistics, BHF statistics fact sheet England. Accessed March 2021. Available at: <https://www.bhf.org.uk/what-we-do/our-research/heart-statistics>
2. NHS Digital (2020) Hospital admitted patient care activity, 2019-20: Primary diagnosis 3 character. Accessed March 2021. Available at: <https://digital.nhs.uk/data-and-information/publications/statistical/hospital-admitted-patient-care-activity/2019-20>
3. National Institute for Cardiovascular Outcomes Research (2019) National heart failure audit 2017/18. Accessed March 2021. Available at: <https://www.hqip.org.uk/wp-content/uploads/2019/09/Ref-129-Cardiac-Heart-Failure-Summary-Report-2019-FINAL.pdf>
4. Taylor CJ, Ordonez-Mena JM, Roalfe AK et al. (2019) Trends in survival after a diagnosis of heart failure in the United Kingdom 2000-2017: population based cohort study. BMJ 364:l223.