Practice Guidelines

NICE Updates Guidelines on Management of Chronic Heart Failure

MARA LAMBERT

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Heart failure is associated with poor prognosis and quality of life, as well as high health care costs. In 2010, the National Institute for Health and Clinical Excellence (NICE) updated its previous guideline on the management of chronic heart failure in adults. This update focuses on the role of signs and symptoms, serum natriuretic peptide levels, and echocardiography in the diagnosis of heart failure. It also discusses pharmacologic treatment of heart failure and management using cardiac resynchronization therapy and implantable cardioverter-defibrillators (ICDs), as well as disease monitoring. These recommendations apply to nonpregnant adults with symptoms of chronic heart failure, but not with acute heart failure or acute exacerbations of chronic heart failure.

Diagnosis

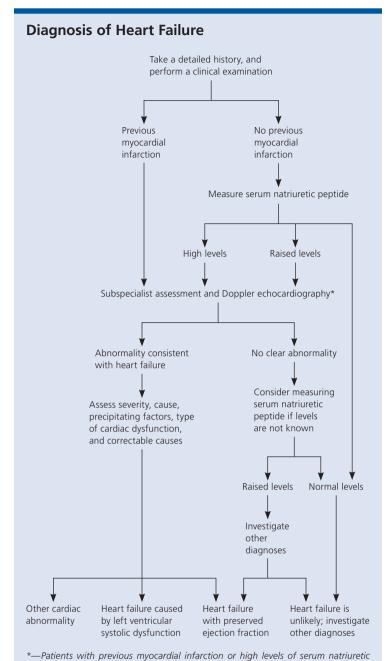
Diagnosis of heart failure begins with a history and physical examination (*Figure 1*). To rule out other diagnoses, electrocardiography and consideration of chest radiography, peak flow measurement or spirometry, and blood tests are appropriate. Clinical signs and symptoms have limited use in diagnosing heart failure. Measurement of serum natriuretic peptide levels (brain natriuretic peptide [BNP] and N-terminal pro-BNP) has a high sensitivity but only moderate specificity for diagnosing heart failure.

In patients without previous myocardial infarction (MI), it is appropriate to measure serum natriuretic peptide levels and perform subsequent echocardiography. Subspecialist evaluation is needed if levels are elevated. In patients with a history of MI, the physician should proceed directly to echocardiography with subspecialist evaluation. If the echocardiogram is normal, measurement of serum natriuretic peptide levels should be considered. Patients with previous MI or with high levels of serum natriuretic peptide should undergo echocardiography and subspecialist evaluation within two weeks of presentation. Those with no previous MI and raised (but not high) levels of serum natriuretic peptide should have echocardiography and subspecialist evaluation within six weeks of initial presentation. If an abnormality is discovered, the physician can begin assessing the severity and possible cause of cardiac dysfunction.

Treatment

Management of chronic heart failure depends on whether the patient has heart failure with preserved ejection fraction or caused by left ventricular systolic dysfunction (*Figure 2*). In patients with preserved ventricular function, physicians should focus on managing comorbid conditions (e.g., hypertension, ischemic heart disease, diabetes mellitus) because there is insufficient evidence to recommend specific therapies in this population. Diuretics may be prescribed to manage fluid retention.

Patients with left ventricular systolic dysfunction should receive an angiotensin-converting enzyme (ACE) inhibitor and a beta blocker, regardless of symptom severity. High-quality evidence has shown that this treatment reduces morbidity and increases survival. Outcomes are the same regardless of whether ACE inhibitor or beta-blocker therapy is started first. If patients cannot tolerate ACE inhibitors, angiotensin receptor blockers (ARBs) may be considered. Second-line therapy options include ARBs, aldosterone antagonists, and combination therapy with hydralazine and a nitrate. Hydralazine in combination with a nitrate is appropriate in patients who cannot tolerate ACE inhibitors or ARBs. When choosing a second-line therapy, physicians should consider the



peptide should have subspecialist assessment and echocardiography within two weeks of presentation. Those without a previous myocardial infarction and raised levels of serum natriuretic peptide should have subspecialist assessment and echocardiography within six weeks of presentation.

Figure 1. Algorithm for the diagnosis of heart failure. Normal serum natriuretic peptide levels are defined as brain natriuretic peptide (BNP) levels < 100 pg per mL (100 ng per L) or N-terminal pro-BNP levels < 400 pg per mL (400 ng per L). Raised serum natriuretic peptide levels are defined as BNP levels between 100 and 400 pg per mL (100 and 400 ng per L) or N-terminal pro-BNP levels between 400 and 2,000 pg per mL (400 and 2,000 ng per L). High serum natriuretic peptide levels are defined as BNP levels > 400 pg per mL or N-terminal pro-BNP levels > 2,000 pg per mL.

Adapted with permission from Mant J, Al-Mohammad A, Swain S, Laramée P. Management of chronic heart failure in adults: synopsis of the National Institute for Health and Clinical Excellence Guideline. Ann Intern Med. 2011;155(4):254.

severity of heart failure, the patient's ethnicity, and comorbid conditions. Monitoring renal function is important in patients taking any of these medications. Potassium levels should be monitored closely in those taking ARBs or aldosterone antagonists.

Patients with left ventricular systolic dysfunction and persistent symptoms should be referred to a subspecialist for possible additional drug therapy (digoxin), cardiac resynchronization therapy, or an ICD. Criteria for cardiac resynchronization therapy include a left ventricular ejection fraction less than 35 percent and a QRS duration on electrocardiography of 150 milliseconds or higher, or between 120 and 149 milliseconds in those with mechanical dyssynchrony on echocardiography. Criteria for an ICD include sustained ventricular tachycardia or nonsustained ventricular tachycardia that is inducible on electrophysiologic testing if the left ventricular ejection fraction is less than 35 percent, or a QRS duration of 120 milliseconds or higher on electrocardiography if the left ventricular ejection fraction is less than 30 percent.

All patients with heart failure should be offered supervised group exercise—based rehabilitation that includes psychological and educational components, as long as they are stable and do not have a condition or device that precludes such a program. Moderate-quality evidence has shown that exercise rehabilitation reduces hospital admissions for heart failure and increases long-term quality of life almost exclusively in patients with left ventricular systolic dysfunction.

Monitoring Patients with Heart Failure

Because the clinical course of heart failure is unpredictable and fluctuating, physicians should monitor patients to make sure they are receiving optimal therapy and adjust treatment as necessary. Serial monitoring of serum natriuretic peptide levels has been shown to be cost-effective when used by subspecialists. Moderate-quality evidence has shown that therapy guided by serum natriuretic peptide levels can lead to a medium-term reduction in hospitalization for heart failure, although there was no reduction in mortality in persons older than 75 years, in quality of life, or in hospitalization for any

Treatment of Heart Failure Heart failure Heart failure Heart failure caused with preserved by left ventricular ejection fraction systolic dysfunction* Offer ACE inhibitors and beta Manage comorbid conditions, such as high blood pressure, blockers licensed for heart ischemic heart disease, and failure as first-line treatment diabetes mellitus, in accordance (consider an ARB if the patient with NICE guidance cannot tolerate ACE inhibitors) If symptoms persist despite optimal first-line treatment, seek subspecialist advice Offer rehabilitation and education, and diuretics for congestion and fluid retention; consider an ICD where appropriate† For second-line treatment, consider adding one of the following: • An aldosterone antagonist licensed for heart failure (especially in patients with moderate to severe heart failure; or myocardial infarction in the past month) • An ARB licensed for heart failure§ (especially in mild to moderate heart failure||) • Hydralazine in combination with a nitrate (especially in persons of African or Caribbean origin¶ with moderate to severe heart failure‡; also may be considered if the patient cannot tolerate ACE inhibitors and ARBs) If symptoms persist, consider the following: CRT (pacing with or without a defibrillator)** Digoxin

- *—For more information on drug treatment, see Appendix J to the NICE clinical guideline on heart failure and the NICE clinical guideline on chronic kidney disease.
- †—Consider using an ICD according to the recommendations described in the NICE technology appraisal on ICDs.
- ‡—New York Heart Association class III to IV.
- §—Not all ARBs are licensed for use in heart failure in combination with ACE inhibitors.
- ||—New York Heart Association class II to III.
- \P —This does not include persons of mixed race. For more information, see the full NICE chronic heart failure guideline.
- **—Consider using CRT according to the recommendations described in the NICE technology appraisal on CRT.

Figure 2. Algorithm for the treatment of heart failure. The term "licensed for heart failure" refers to drugs that have been approved for use for the given indication by regulatory agencies in the United Kingdom. (ACE = angiotensin-converting enzyme; ARB = angiotensin receptor blocker; CRT = cardiac resynchronization therapy; ICD = implantable cardioverter-defibrillator; NICE = National Institute for Health and Clinical Excellence.)

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cause. Telemonitoring can reduce mortality and hospitalization for any cause, but does not improve quality of life or decrease hospitalization for heart failure. However, the guideline does not make a recommendation on the use of telemonitoring because of heterogeneity of results from different trials.

New Evidence

Since publication of the 2010 update of the NICE clinical guideline on heart failure, some new evidence has emerged, although it is unclear whether these results would have affected guideline recommendations. The Eplerenone in Mild Patients Hospitalization and Survival Study in Heart Failure investigated patients with chronic heart failure caused by left ventricular systolic dysfunction with mild symptoms (i.e., New York Heart Association class II heart failure). Results showed significant reductions in hospitalization and mortality when eplerenone (Inspra) therapy was started in patients hospitalized during the preceding six months or in those with persistent moderate elevation in serum natriuretic peptide levels. The Systolic Heart Failure Treatment with the If Inhibitor Ivabradine Trial showed that ivabradine, a blocker of the I_f channel in the sinoatrial node, significantly reduced unplanned hospitalization and mortality in patients with heart failure caused by left ventricular systolic dysfunction who had a heart rate above 70 beats per minute. A large trial of telemonitoring in patients with a recent hospitalization for heart failure found no evidence that it reduced hospital readmission or mortality. The Resynchronisation/Defibrillation for Ambulatory Heart Failure Trial found that adding cardiac resynchronization therapy in patients with an ICD who had mild to moderate heart failure reduced all-cause mortality and hospitalization, but increased adverse events.

Answers to This Issue's CME Quiz

Q1. A	Q6. B, C, I
Q2. B	Q7. B
Q3. C, D	Q8. A
Q4. A	Q9. A, D
Q5. D	Q10. A, B