

Decision Making in Advanced Heart Failure

A Scientific Statement From the American Heart Association

Endorsed by Heart Failure Society of America, American Association of Heart Failure Nurses, and Society for Medical Decision Making

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Shared decision making for advanced heart failure has become both more challenging and more crucial as duration of disease and treatment options have increased. High-quality decisions are chosen from medically reasonable options and are aligned with values, goals, and preferences of an informed patient. The top 10 things to know about decision making in advanced heart failure care are listed in Table 1.

Why Shared Decision Making?

Providers have an ethical and legal mandate to involve patients in medical decisions. Shared decision making recognizes that there are complex trade-offs in the choice of medical care.¹ Shared decision making also addresses the ethical need to fully inform patients about the risks and benefits of treatments.² In the setting of multiple reasonable options for medical care, shared decision making involves clinicians working with patients to ensure that patients' values, goals, and preferences guide informed decisions that are right for each individual patient.

Grounded in the ethical principle of autonomy,³ judicial decisions (eg, *Cruzan v Missouri Department of Health*⁴) and legislative actions (eg, the Patient Self-Determination Act⁵) have repeatedly affirmed the rights of patients or duly appointed surrogates to choose their medical therapy from among reasonable options.⁶ The formal process of informed consent before procedural interventions is an embodiment of this concept in that it underscores the clinician's obligation to ensure that the patient has the opportunity to be informed.³ An informed patient is one who is aware of the diagnosis and prognosis, the nature of the proposed intervention, the risks and benefits of that intervention, and all reasonable alternatives and their associated risks and benefits.⁷ A major purpose of a high-functioning healthcare system is to provide the resources with which an activated, informed patient can engage in productive discussions with a proactive, prepared healthcare team.⁸

Shared decision making moves beyond informed consent. It asks that clinicians and patients share information with each other and work toward patient-centered decisions about

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Table 1. Top Ten Things to Know

1. Shared decision making is the process through which clinicians and patients share information with each other and work toward decisions about treatment chosen from medically reasonable options that are aligned with the patients' values, goals, and preferences.
2. For patients with advanced heart failure, shared decision making has become both more challenging and more crucial as duration of disease and treatment options have increased.
3. Difficult discussions now will simplify difficult decisions in the future.
4. Ideally, shared decision making is an iterative process that evolves over time as a patient's disease and quality of life change.
5. Attention to the clinical trajectory is required to calibrate expectations and guide timely decisions, but prognostic uncertainty is inevitable and should be included in discussions with patients and caregivers.
6. An annual heart failure review with patients should include discussion of current and potential therapies for both anticipated and unanticipated events.
7. Discussions should include outcomes beyond survival, including major adverse events, symptom burden, functional limitations, loss of independence, quality of life, and obligations for caregivers.
8. As the end of life is anticipated, clinicians should take responsibility for initiating the development of a comprehensive plan for end-of-life care consistent with patient values, preferences, and goals.
9. Assessing and integrating emotional readiness of the patient and family is vital to effective communication.
10. Changes in organizational and reimbursement structures are essential to promote high-quality decision making and delivery of patient-centered health care.

treatment.⁹ Shared decision making incorporates the perspective of the patient, who is responsible for articulating goals, values, and preferences as they relate to his or her health care. Shared decision making incorporates the perspective of the clinician, who is responsible for narrowing the diagnostic and treatment options to those that are medically reasonable. Shared decision making is most easily applied to preference-sensitive decisions, in which both clinicians and patients agree that equipoise exists, and decision support helps patients think through, forecast, and deliberate their options. However, in situations in which clinicians hold the view that scientific evidence for benefit strongly outweighs harm, behavioral support (eg, smoking cessation counseling) designed to describe, justify, and recommend specific behavior may also be appropriate and complementary to decision support.¹⁰ Finally, certain therapeutic options may be considered unreasonable and therefore independent of patient demands, although situations of medical futility are relatively rare.⁶ Although not all patients will be able to clearly articulate decisions that are congruent with their stated goals, shared decision making aims to ensure that patients' values, goals, and preferences are explored and incorporated into the medical decision-making process.

Patient-centered medicine has been suggested as the next phase in health care.¹¹ Shared decision making puts into practice the principle of "patient-centered care," which the Institute of Medicine has identified as 1 of the 6 pillars of quality,¹² with patient-centered care defined as "providing care that is respectful of and responsive to individual patient preferences, needs, and values and ensuring that patient values guide all clinical decisions."¹² The Patient Protection and Affordable Care Act devotes 4 pages to patient-centered care, specifically calling for the

Table 2. European Society of Cardiology Criteria for Advanced Chronic Heart Failure

1. Moderate to severe symptoms of dyspnea and/or fatigue at rest or with minimal exertion (NYHA functional class III or IV)
2. Episodes of fluid retention and/or reduced cardiac output
3. Objective evidence of severe cardiac dysfunction demonstrated by at least 1 of the following:
 - Left ventricular ejection fraction <30%
 - Pseudonormal or restrictive mitral inflow pattern by Doppler
 - High left and/or right ventricular filling pressures, or
 - Elevated B-type natriuretic peptide
4. Severe impairment of functional capacity as demonstrated by either inability to exercise, 6-min walk distance <300 m, or peak oxygen uptake <12 to 14 mL · g⁻¹ · min⁻¹
5. History of at least 1 hospitalization in the past 6 mo
6. Characteristics should be present despite optimal medical therapy

NYHA indicates New York Heart Association.

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development of decision aids, shared decision-making programs, and metrics for the quality of decision making.¹³

It will be assumed throughout this document that discussions and decision making with patients also include, when appropriate, the family and other individuals involved, such as caregivers and companions. The approach to decision making outlined in this Scientific Statement takes the perspective of the individual patient rather than that of society in general. Although individual medical decisions taken collectively have implications for distributive justice and resource allocation, it is not the responsibility of clinicians, patients, or families to directly factor these global considerations into individual decisions.¹⁴ Rather, discussions regarding alternative treatment options, including no treatment, should be focused on meeting a specific individual's values, goals, and preferences within the context of societal rules and regulations.

Why Advanced Heart Failure?

Heart failure affects 2.4% of the adult population and over 11% of the expanding population >80 years old.¹⁵ Estimated total heart failure costs in the United States are projected to reach 44.6 billion by 2015.¹⁵ Existing therapies slow, but infrequently reverse, disease progression. As a result, the prevalence of symptomatic heart failure has increased, including a prolongation of the advanced phase of the disease.¹⁶ The American Heart Association characterizes the far end of the heart failure continuum as stage D, or "refractory end-stage heart failure,"¹⁷ further defined by others,^{18,19} including the European Society of Cardiology (Table 2).²⁰ These overlapping definitions describe a group of patients for whom symptoms limit daily life despite usual recommended therapies and for whom lasting remission into less symptomatic disease is unlikely. The increasing prevalence, high symptom burden, and possible disease-exchanging therapies (ie, transplantation and mechanical circulatory support) for patients living with advanced heart failure mandate a systematic and thoughtful approach to decision making.

This Scientific Statement reviews the clinical context for decision making in advanced heart failure and provides

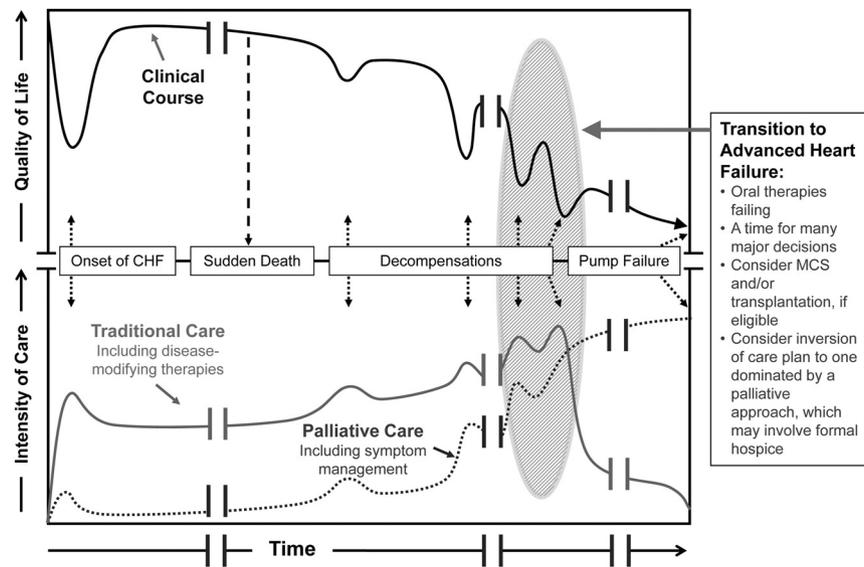


Figure 1. A depiction of the clinical course of heart failure with associated types and intensities of available therapies. Black line: Patients tend to follow a progressive, albeit nonlinear, decline in health-related quality of life as the disease progresses; this course can be interrupted by sudden cardiac death caused by arrhythmia or can end in a more gradual death caused by progressive pump failure. Gray line: At disease onset, multiple oral therapies are prescribed for cardiac dysfunction and/or treatment of comorbidities. As disease severity increases, the intensity of care may increase in parallel, with intensification of diuretics, addition of an implantable cardioverter-defibrillator/cardiac resynchronization therapy for those eligible, and increasing interaction with the medical system through ambulatory visits and hospitalizations, until the time when standard therapies begin to fail (transition to advanced heart failure). Dotted line: Palliative therapies to control symptoms, address quality of life, and enhance communication are relevant throughout the course of heart failure, not just in advanced disease; palliative therapies work hand in hand with traditional therapies designed to prolong survival. The critical transition into advanced heart failure from the medical perspective is often followed by a transition in goals of care from the patient and family perspective, wherein palliative therapies may become the dominant treatment paradigm (for the majority of patients in whom transplantation and mechanical circulatory support are not an option). Clinicians must recognize the transition to advanced heart failure so that therapeutic options can be considered in a timely fashion and patients are able to proactively match medical decisions to clinical realities. CHF indicates chronic heart failure; MCS, mechanical circulatory support. Modified from Lanken et al;²¹ reprinted with permission of the American Thoracic Society. Copyright © 2012, American Thoracic Society.

guidance on communication techniques to support these decisions. Its goal is primarily to help healthcare providers of all types integrate these concepts into their routine practice to promote the delivery of effective, safe, efficient, timely, equitable, and patient-centered care.¹² We recognize that major barriers to the implementation of these concepts are time, training, and resources. We also recognize the limited and inequitable access to experts with formal training in heart failure and palliative care, which leaves many of these responsibilities to be borne by healthcare providers in a general medical setting. If the goals of this document are to be realized, however, the healthcare system will need to make a fundamental commitment to shared decision making, with realignment of incentives to support the tailoring of advanced care to individual patients. Without changes in the structure of the healthcare team and associated reimbursement, these recommendations will remain an unfunded mandate that are unlikely to be fully realized in most practice settings.

Expectations for the Future

- Attention to the clinical trajectory is required to calibrate expectations and guide timely decisions.
- Predictive models can target high-risk populations but leave wide uncertainties around estimates of survival for an individual.
- Difficult discussions now will simplify difficult decisions in the future.

- Uncertainty is inevitable and should be included in discussions with patients and family.

Estimating Prognosis in Heart Failure

Assessment of prognosis is the foundation for selection among therapies for life-threatening disease, but this is particularly challenging for heart failure. The clinical course varies dramatically across the spectrum of disease severity and is relatively unpredictable for individual patients (Figure 1).^{19,21} This contrasts with the more linear decline of patients with advanced cancer, which has traditionally been the model for approaches to end-stage disease. Even late in heart failure, patients often enjoy “good days” and brief interludes of apparent stability, which can lull them and their care providers into postponing vital decisions. Prognosis is further clouded by the unique contrast between unexpected sudden death (ie, lethal arrhythmia) and lingering death with congestive symptoms (ie, progressive pump failure). Frequent reappraisal of the clinical trajectory helps calibrate expectations, guide communication, and inform rational decisions.

More than 100 variables have been associated with mortality and rehospitalization in heart failure.^{22–27} Examples of prognostic factors include demographics (age, sex, race, insurance status), functional status (New York Heart Association functional class and health-related quality-of-life scores), exercise capacity (peak oxygen consumption, 6-minute walk), cardiac structure and function (cardiac cham-

Table 3. Selected Prognostic Models in Heart Failure

	Key Covariates	Outcome
Ambulatory		
Heart Failure Survival Score ²³	Peak $\dot{V}O_2$, LVEF, serum sodium, mean BP, HR, ischemic etiology, QRS duration/morphology	All-cause mortality
Seattle Heart Failure Model ²² (depts.washington.edu/shfm) ^{22a}	NYHA function class, ischemic etiology, diuretic dose, LVEF, SBP, sodium, hemoglobin, percent lymphocytes, uric acid, and cholesterol	All-cause mortality, urgent transplantation, or LVAD implantation
Hospitalized		
EVEREST Risk Model ²²	Age, diabetes, h/o stroke, h/o arrhythmia, β -blocker use, BUN, sodium, BNP, KCCQ scores	The combined end point of mortality or persistently poor quality of life (KCCQ <45) over the 6 mo after discharge
EFFECT ²⁹	Age, SBP, respiratory rate, sodium, hemoglobin, BUN, h/o CVA, h/o dementia, h/o COPD, h/o cirrhosis, h/o cancer	30-d and 1-y mortality
ADHERE ²⁸	BUN, SBP, serum creatinine	In-hospital mortality
ESCAPE Discharge Score ³¹	BNP, cardiopulmonary resuscitation or mechanical ventilation during hospitalization, BUN, sodium, age >70 y, daily loop diuretic dose, lack of β -blocker, 6-min walk distance	6-mo mortality

$\dot{V}O_2$ indicates oxygen consumption; LVEF, left ventricular ejection fraction; BP, blood pressure; HR, heart rate; NYHA, New York Heart Association; SBP, systolic BP; LVAD, left ventricular assist device; EVEREST, Efficacy of Vasopressin Antagonism in Heart Failure Outcome Study with Tolvaptan; h/o, medical history of; BUN, blood urea nitrogen; BNP, B-type natriuretic peptide; KCCQ, Kansas City Cardiomyopathy Questionnaire; EFFECT, Enhanced Feedback for Effective Cardiac Treatment; CVA, cerebrovascular accident; COPD, chronic obstructive pulmonary disease; ADHERE, Registry for Acute Decompensated Heart Failure Patients; and ESCAPE, Evaluation Study of Congestive Heart Failure and Pulmonary Artery Catheterization Effectiveness.

ber size, ejection fraction), assessments of filling pressures, biomarkers (natriuretic peptides, inflammatory markers), renal and liver dysfunction, comorbidities (diabetes, lung disease), clinical events (defibrillator shocks and recent hospitalizations), psychosocial factors (depression, social isolation), and behavioral factors (eg, adherence to the medical regimen).

A variety of multivariable models have been published in an effort to provide more refined predictions of prognosis in patients with heart failure (Table 3). The most commonly used multivariable instruments for estimating prognosis in symptomatic outpatients are the Heart Failure Survival Score²³ and the Seattle Heart Failure Model.²² In patients hospitalized for heart failure, a variety of inpatient models have been developed to predict both in-hospital²⁸ and post-discharge outcomes.^{26,27,29–31} These inpatient models have highlighted the strength of natriuretic peptides, renal function, and low blood pressure as predictors of survival in patients in this setting.²⁴ Recently, the first model to predict both mortality and quality-of-life outcomes after discharge has been published.³² Although all of these models require complex mathematical formulas to generate risks, the increasing use of health information technology in the delivery of care offers the potential to automatically generate risk profiles from the electronic medical record.

The application of commonly used ambulatory heart failure models to the advanced heart failure population can result in miscalibrated estimates of life expectancy, with significant underestimation of risk in certain populations.^{33,34} Therefore, before recommending general use of risk models, adequate discrimination (ie, the ability of a model to accurately distinguish between a patient who will experience the event versus one who will not)^{35,36} and calibration (ie, the ability of the model to accurately predict the observed probability of an event across levels of risk)³⁷ will need to be validated for broader populations than those from clinical trials.^{33,34}

Prognosis for Both Quantity and Quality of Life

Most prognostic models in heart failure focus on mortality, which is easily determined and highly relevant; however, other clinical outcomes also rank high in importance to individual patients (Figure 2). Multiple studies have documented patients’ willingness to sacrifice survival in exchange for symptom relief, a trade-off that varies between patients and within the same patient over time and is correlated loosely with disease severity^{39,40} but strongly with do-not-resuscitate status.⁴¹ A full discussion of prognosis therefore includes not only the risks of death but also the potential burdens of worsening symptoms, limited functional capacity, loss of independence, reduced social functioning, decreased quality of life, and increased caregiver commitment.⁴² Unfortunately, much less is known about the risks of these latter outcomes. The only existing model that estimates the risk of unfavorable future quality of life shows important differences from risk models for death, particularly the relative impor-

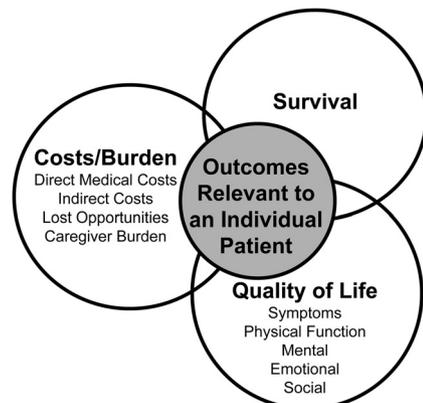


Figure 2. Prognosis is not only about expectations for survival. There are multiple domains that are of varying importance to individual patients. Adapted from Spilker.³⁸

tance of current measures of quality of life.³² More astute anticipation of an unfavorable quality of life until death, in addition to anticipation of death, would better identify patients for whom detailed discussions of prognosis and options are appropriate. In choosing among options, this information gap regarding nonmortality patient-centered outcomes is exacerbated by the lack of rigor in collecting health status information in major trials, although this is improving. Even less is known about the relative impact of the disease and therapies on caregiver burden and quality of life for family members.^{43–46}

Uncertainty for the Individual

Even under these idealized circumstances, most models designed to predict mortality have only modest accuracy.⁴⁷ Further complicating practical use, prediction models represent the average survival for a population of patients with characteristics similar to those of the individual patient. A 70% chance of 2-year survival does not directly translate to an individual who will instead be 100% alive or dead at any point in time. For patients with advanced disease, interest often focuses instead on the expected length of time remaining; patients ask the question, “How long do I have?” This point prediction of survival time⁴⁸ is even more difficult to estimate.^{49,50} Even if a model fits well for a cohort and the estimated survival curve provides a good fit to the data, it is not clear where along the curve an individual patient will lie. As an example of the difficulty in estimating survival duration, one can consider the median survival estimate (50% survival at time x) as an estimation of the time in which half of the patients will live longer and half will live for a shorter time. Parkes⁵¹ defines an “error” in survival as an estimate more than twice as long as the actual survival or less than half the actual survival. That is, if a patient survived for 12 months, a predicted survival of >2 years or <6 months would be considered an error by this definition. This error depends on the variability in survival times for patients, more specifically on the standard deviation of the logarithm of survival time. Using several statistical models of survival, the probability of greater than 2-fold error remains near 50% under realistic assumptions.⁴⁹

Ultimately, the stochastic nature of heart failure conveys a high level of prognostic uncertainty for most patients. Future events have a certain degree of unpredictability, such that improved understanding of risk tends to add incrementally less prognostic information to existing models. Even a perfect model that includes all possible measurements describes only what has already happened. The trajectory can often be steepened by new conditions or life events, such as myocardial infarction, a serious fall, or the death of a spouse. It is vital to acknowledge uncertainty in discussions about future care.

Need for Accurate Estimates of Risk

Despite limitations of prognostic models, they are generally more accurate than clinical intuition, which is prone to bias. A review of survival predictions among terminally ill cancer patients⁵² found that physicians consistently overestimated survival, which has been seen in other studies.^{51,53} For

patients discharged from the hospital with advanced-stage heart failure, both physicians’ and nurses’ survival estimates had modest ability to discriminate those who subsequently died from those who lived (with nurses outperforming physicians), but absolute estimates were significantly miscalibrated, again overestimating survival.⁵⁴ In patients with chronic heart failure, the patient-predicted survival also tended to overestimate survival versus model-based predictions, particularly for younger patients.⁵⁵ Clinicians need to learn how to leverage objective risk models, while recognizing their limitations and adapting them on the basis of their unique clinical and psychosocial features and serial assessments not generally incorporated into such models.

Anticipation, Timing, and Review

- An annual heart failure review with patients should include discussion of current and potential therapies for both anticipated and unanticipated events.
- On the day of hospital admission, it is far better to review rather than introduce advanced care decisions, which requires that patient preferences have been discussed previously and documented in the ambulatory setting.
- Clinical milestones such as implantable cardioverter-defibrillator (ICD) shocks or recurrent hospitalization should trigger interim review and discussion of treatment options and preferences.

Timing of Discussions

Finding appropriate time to discuss preferences, prognosis, and medical options is a formidable challenge. Such discussions require a major commitment of time, focus, and emotional energy, which is not in synchrony with the frenetic pace and frequent interruptions of clinical practice. Current organizational and reimbursement structures provide strong disincentive to such intense encounters.

As a result, formal discussions about prognosis and decision making are often deferred until more emergent and less favorable occasions, when thoughtful decision making may be impaired. For instance, at the time of presentation for hospital admission with decompensated heart failure, patients are frequently uncomfortable and often require urgent, intensive evaluation and management. Clinicians responsible for delivering care in this setting are typically unfamiliar with the patient and overall disease trajectory. Hasty questions such as, “Do you want us to do everything?” and “Would you want to be kept alive as a vegetable?” can yield inaccurate and conflicting answers. It has been shown that patients deciding resuscitation preferences during an acute hospitalization frequently reverse their decisions over the next few months.⁵⁶

Therefore, optimal shared decision making requires that patient preferences have been discussed previously and documented in the ambulatory setting. The day of hospital admission is a time to review and possibly update, rather than introduce, advanced care decisions. On the other hand, once the clinical course has become apparent during hospitalization, clinicians can take advantage of the substantial time they have with the patient and family to further address complex medical decisions before discharge. When the expected survival or quality of life is very limited, hospitalization may

Table 4. Triggers for Formally Assessing Prognosis and Having Conversations About Goals of Care and Voluntary Advance Care Planning

Routine
“Annual Heart Failure Review” with a scheduled clinic visit
Event-driven “milestones” that should prompt reassessment
Increased symptom burden and/or decreased quality of life
Significant decrease in functional capacity
Loss of ADLs
Falls
Transition in living situation (independent to assisted or LTC)
Worsening heart failure prompting hospitalization, particularly if recurrent ⁵⁷
Serial increases of maintenance diuretic dose
Symptomatic hypotension, azotemia, or refractory fluid retention necessitating neurohormonal medication underdosing or withdrawal ⁵⁸
Circulatory-renal limitations to ACEI/ARB
Decrease or discontinuation of β -blockers because of hypotension
First or recurrent ICD shock for VT/VF ⁵⁹
Initiation of intravenous inotropic support
Consideration of renal replacement therapy
Other important comorbidities: new cancer, etc
Major “life events”: death of a spouse
ADL indicates activities of daily living; LTC, long-term care; ACEI, angiotensin-converting enzyme inhibitor; ARB, angiotensin II receptor blocker; ICD, implantable cardioverter-defibrillator; VT, ventricular tachycardia; and VF, ventricular fibrillation.

also afford better access to multidisciplinary teams, palliative care, and other resources than can be marshaled in the outpatient setting. All of these considerations underscore the importance of a proactive, anticipatory, and iterative approach to soliciting patients’ preferences. This should occur both routinely and at the occurrence of milestones that herald a worsening prognosis (Table 4).

Annual Heart Failure Review

The “Annual Heart Failure Review” is a concept based on the annual wellness visits that have long been a successful part of primary care. This reflects the principle and practice of “anticipatory guidance,” the psychological preparation of a person to help relieve the fear and anxiety of an event expected to be stressful. When triggered by a scheduled anniversary in the same way as well baby visits or periodic mammography, an automatic annual review can open a broad dialogue with patients and families without the unvoiced concern that it signifies bad news. In heart failure, this may coincide, for example, with an annual influenza vaccination or at 1-year increments roughly originating from the date of diagnosis. It may be convenient to have this review occur in temporal proximity to an annual general medical evaluation, particularly with regard to screening studies, for which the indications might change in the setting of progressive heart disease.

In the annual review visit (Table 5), a variety of tasks could be accomplished. Patients could summarize their recent symptom burden and quality of life. Goals for the coming year and preferences for outcomes including survival, func-

Table 5. Selected Components That May Be Included in an Annual Heart Failure Review

Characterization of clinical status
Functional ability, symptom burden, mental status, quality of life, and disease trajectory
Perceptions from caregiver
Solicitation of patient values, goals, and general care preferences
Estimation of prognosis
Consider incorporating objective modeling data
Orient to wide range of uncertainty
Review of therapies
Indicated heart failure therapies in appropriate patients (BB, ACEI/ARB, AA, CRT, ICD)
Treatment of comorbidities (AF, HTN, DM, CKD, etc)
Appropriate preventive care, within the context of symptomatic heart failure
Planning for future events/advance care planning
Resuscitation preferences
Desire for advanced therapies, major surgery, hospice
Standardized documentation of the annual review in the medical record
BB indicates β -blocker; ACEI, angiotensin-converting enzyme inhibitor; ARB, angiotensin II receptor blocker; AA, aldosterone antagonist; CRT, cardiac resynchronization therapy; ICD, implantable cardioverter-defibrillator; AF, atrial fibrillation; HTN, hypertension; DM, diabetes mellitus; and CKD, chronic kidney disease.

tional capacity, and quality of life could be solicited. A range of prognosis would be estimated and broadly conveyed. Current treatment with drugs and devices could be reviewed relative to indicated treatment based on the patient’s heart failure type, stage, and trajectory. Similarly, evaluation and management of new relevant comorbidities could be reviewed, such as sleep apnea, anemia, and depression. “Voluntary advance care planning,” including formal designation of a healthcare proxy and do-not-resuscitate status, which has been proposed as part of the initial Medicare visit and subsequent “wellness visits,”^{60(p73406)} would be essential and would take place naturally within the context of an annual review.

This scheduled review would require considerable face-to-face time between the patient, family, and physician. The results of this discussion should be documented specifically in a designated area of the chart available to all who might be involved in the patient’s current and future care.

Responding to Milestones

Although heart failure is a chronic disease, its clinical course often includes sudden changes. There are several “milestones” in the clinical course of heart failure that can represent an “inflection point” in the overall trajectory (Figure 1^{19,21}), such as first ICD shock, rehospitalization, development of cardiorenal syndrome, withdrawal of angiotensin-converting enzyme inhibitors, or intubation. Once the acute condition has been addressed, such events should trigger a focused version of the “Heart Failure Review,” which would include reassessment of prognosis, treatment options, and patient preferences.

Table 6. Framework of Major Medical Decisions in Advanced Heart Failure Faced by Patients and Their Clinicians

Types of Options	Specific Examples of Interventions	Generally Considered Only for HF With Reduced LVEF	Examples of Uncommon Outcomes That Could Be Anticipated With “What If” Discussions in High-Risk Patients
Major interventions that may improve cardiac function	CABG		Worsened cardiac function/inability to come off bypass or IABP: Place MCS?
	Valve surgery		Ventilator dependence: Extubate? When?
	Pericardial stripping		Stroke: Feeding tube? Institutional care?
	Percutaneous valve intervention		Coronary occlusion: Revert to CABG?
	PCI		Unable to place coronary sinus lead: Convert to thoracotomy?
Therapies that only reduce the risk of sudden cardiac death	CRT	X	
	ICD	X	Terminal or permanently disabling disease: Device deactivation?
Adjunctive therapies instituted during acute decompensation with potential chronic dependence	Temporary support devices (IABP, percutaneous VAD, ECMO)	X	Unable to wean: Convert to permanent MCS or withdraw?
	IV inotropes	X	Unable to wean: Transition to home inotropes or discontinue?
Advanced surgical therapies to exchange disease	Renal replacement therapy (dialysis or ultrafiltration)		Failure of acute injury to resolve: Initiate indefinite hemodialysis or discontinue?
	Transplantation	X	Early graft failure or other serious postoperative complications: MCS or withdraw support?
	Permanent MCS/LVAD	X	Later graft failure: Retransplantation? Stroke, infection, or recurrent bleeding: Turn off device?
Noncardiac procedures for comorbidities	Joint replacement		Worsening heart failure causing hemodynamic and/or respiratory collapse: Continue ventilatory support and/or initiate circulatory support?
	Hernia repair		
	Resection of pulmonary nodule Asymptomatic aortic aneurysm repair Screening colonoscopy	Not generally to be done, because risks are thought to outweigh potential benefit	

HF indicates heart failure; LVEF, left ventricular ejection fraction; CABG, coronary artery bypass grafting surgery; IABP, intra-aortic balloon pump; MCS, mechanical circulatory support; PCI, percutaneous coronary intervention; CRT, cardiac resynchronization therapy; ICD, implantable cardioverter-defibrillator; VAD, ventricular assist device; ECMO, extracorporeal membranous oxygenation; and IV, intravenous.

Framework of Options

- Physicians are responsible for defining the range of options that are medically appropriate.
- Presentation of major interventions should always include specific description of alternative approaches, including continuation or withdrawal of ongoing treatments and focus on symptomatic care.
- Discussions should include a range of anticipated outcomes, including not only survival but also major adverse events, independence, functional capacity, and quality of life for both patient and caregiver, even if to acknowledge lack of this information for some interventions.
- Therapies that may lead to dependence should be weighed carefully before initiation even when anticipated to be temporary (eg, intravenous inotropes, renal replacement therapy, and intubation).
- Benefits and risks of noncardiac procedures should be reviewed in the context of competing risks for death and functional limitation attributable to heart failure (eg, hip replacement, repair of asymptomatic aortic aneurysm, or screening tests).
- Decisions for major cardiac and noncardiac interventions should include consideration of “what if” situations of unanticipated adversity.

- Referral to a palliative care team should be considered for assistance with difficult decision making, symptom management in advanced disease, and caregiver support even as patients continue to receive disease-modifying therapies.

In the face of the increasing complexity of diagnostic and treatment options for heart failure, a framework for classifying various medical decision-making scenarios should help clinicians better anticipate those decisions most likely to occur as the disease progresses to an advanced stage (Table 6). Too frequently, the default assumption is that patients would “want everything done.” Rather, it is the clinicians who are responsible for defining the “everything” set of interventions that are medically reasonable. From these, patients and families can choose those most consistent with their values, preferences, and goals. It is increasingly explicit in quality metrics that those groups or institutions offering specific advanced therapies should include palliative care and access to ongoing care regardless of the therapies chosen.

Continuation of Stage C Medical Therapies

While anticipating and addressing new options that accompany the transition to advanced stage D heart failure, medical therapy usually includes all stage C therapies.^{17,61,62} The

initial approach to stage D heart failure is optimization of these treatments. The need to decrease or discontinue neurohormonal antagonists is a milestone, as described in Table 4.

Major Interventions That Might Improve Cardiac Function and Clinical Outcomes

High-Risk Cardiac Surgery

Patients may be considered for cardiac surgery for coronary, valvular, and pericardial disease. These surgeries are particularly high risk as a consequence of the patient's advanced heart failure. These procedures may be pursued either with the hope of improving the heart failure condition or in response to a superimposed diagnosis such as angina or acquired aortic stenosis. It is usually anticipated that there will be substantial residual cardiac dysfunction even if the surgery is successful. Although the intent may be to improve cardiac function, the benefit of cardiac surgery in most patients with advanced chronic heart failure is not established. Even if the cardiac function improves, the surgery and related events may lead to prolonged morbidity and possibly death.⁶³ The potential for protracted postoperative rehabilitation and loss of independence must be considered and included thoughtfully in the shared decision, because surgery inherently increases short-term risk for the prospect of longer-term benefit. Unfortunately, there is limited information about the frequency of these outcomes beyond general estimates of prolonged hospitalization in the Society of Thoracic Surgery risk scores.⁶⁴

Percutaneous Interventions

Less invasive percutaneous approaches for the treatment of coronary and valvular disease may be appealing in advanced heart failure because of the increased surgical risk among these patients. However, potential benefits depend on a variety of factors and are relatively unknown for the advanced heart failure population; meanwhile, risks of contrast-induced nephropathy⁶⁵ and 30-day mortality⁶⁶ are markedly increased in this population. Percutaneous approaches to valvular disease are less well developed than for coronary disease, but the technology is improving rapidly. Catheter approaches to both aortic⁶⁷ and mitral⁶⁸ disease have now been shown to be reasonable alternatives to surgery in certain populations. The benefits of valve repair or replacement are less well established in patients with significant heart failure, especially when treating functional (secondary) mitral regurgitation for patients with a dilated left ventricle.⁶⁹ The acute risk of stroke must be weighed against potential longer-term benefits. Decisions regarding percutaneous interventions should also include consideration of whether emergency "bail out" surgery would be appropriate and feasible.

Pacing Device Therapy

Cardiac resynchronization therapy (CRT) represents a clinical challenge in advanced heart failure. Patients with New York Heart Association functional class IV heart failure have represented a small fraction of patients included in randomized trials of CRT.⁷⁰ Although there are some reports suggesting that CRT can improve outcomes for patients taking intravenous inotropes, these findings have not been consistent, and the reports have methodological limitations.^{71,72}

Before left ventricular lead implantation or ICD device upgrade, the care team should plan for contingencies. For instance, if the lead cannot be placed transvenously, will there be consideration of an open thoracotomy for placement, with the attendant morbidity of a chest tube and possible recurrent pleural effusion in the setting of chronically elevated right atrial pressures? The risk-benefit ratios of the full complement of possible procedural variations, in the context of patient preferences, should be considered before any procedure so that the patient can provide truly informed consent and a response to potential adverse outcomes is planned a priori. Factors likely to modify the risk-benefit ratio of device implantation, such as noncardiovascular morbidity and acute decompensation, should also be recognized and incorporated into these discussions.^{73,74}

Although CRT pacers and ICDs can be packaged together, their purposes are quite different. CRT, like neurohormonal antagonist therapy, is designed to reverse remodeling and improve cardiac performance. Although CRT has been shown to improve survival, it also can have significant effects on symptom reduction and quality of life in select patients. In contrast, ICDs treat life-threatening arrhythmia without improving symptoms. Patients may not understand the distinction until after the device has discharged or the issue of deactivation is raised. Recommendation for a combined CRT-defibrillator (CRT-D) device should prompt separate discussions on the indications for defibrillator capacity versus cardiac resynchronization, as well as differences in need for monitoring, chances for inappropriate shocks and worsening heart failure, risks for infection and lead malfunction, and options for deactivation.

ICDs to Reduce the Risk of Sudden Cardiac Death

ICDs are fundamentally different than many life-saving therapies for patients with chronic heart failure with reduced ejection fraction. Neurohormonal antagonist medications and CRT improve cardiac function, thereby reducing mortality, reducing hospitalization, and improving quality of life. In contrast, ICDs improve survival by aborting lethal arrhythmias but do not improve cardiac function or heart failure symptoms. Additionally, ICDs can create an additional burden for patients, particularly from inappropriate discharges and prevention of a rapid death. Because ICDs involve this trade-off between reduced risk of sudden cardiac death and an increased risk of hospitalization,⁷⁵ potential decrease in quality of life,⁷⁶ and higher likelihood of prolonged death from progressive pump failure, meticulous discussion of absolute risks with and without ICDs are particularly important for informed consent and shared decision making.

Temporary Therapies With Potential Dependence

Some therapies are intended for short-term use to stabilize patients, thereby allowing for recovery from reversible insults or transition to more definitive therapy (ie, cardiac transplantation or permanent mechanical circulatory support). Although initially intended as a temporary intervention, such stabilizing therapies can create indefinite dependence if the patient does not improve as hoped or develops an adverse event (eg, stroke, progressive renal failure) that compromises

options for anticipated definitive therapies. Such scenarios are difficult for patients and clinicians and therefore must be anticipated.

Temporary Mechanical Circulatory Support

Short-term circulatory support with intra-aortic balloon pumps or other devices may be initiated when acute or acute-on-chronic hemodynamic instability requires urgent intervention to avoid permanent end-organ dysfunction or death. It may be instituted with the hope of supporting a reversible underlying condition, such as fulminate myocarditis or right-sided heart failure after acute myocardial infarction. It may also be initiated in patients who might be potential candidates for transplantation or permanent circulatory support, in whom (1) there has not been an opportunity to appropriately evaluate their candidacy for more definitive high-dependence therapies, (2) reversibility of end-organ dysfunction is uncertain, or (3) contraindications to more definitive therapies may resolve in the near future. If end-organ dysfunction or contraindications do not resolve, a decision will need to be made about discontinuation. To whatever degree possible, these issues should be addressed before the initiation of short-term support.

Positive Inotropic Agents

Intravenous inotropic agents are commonly initiated in the acute setting for hemodynamic stabilization and to improve end-organ perfusion. Use is most often anticipated to be temporary, with the hope of either clinical improvement or eligibility for more definitive therapies as above. Regardless of intent, initiation of inotropic support for exacerbation of chronic heart failure should be considered a significant clinical milestone (Table 4). When patients fail to wean from intravenous inotropic support, decisions arise concerning its continued chronic use. Therefore, the goals of temporary inotrope use should be established clearly before initiation, and unexpected dependence on this therapy should prompt direct discussions about overall goals of care and the limited range of options available at this juncture.

The decision to arrange for chronic continuous infusions after hospital discharge should be guided by the need for symptom relief and patient preferences. Agreement on the goals of the therapy and various “what if” scenarios should be reached before initiation. Nonrandomized data suggest that the number of hospital days may decrease after initiation of chronic inotrope infusion,⁷⁷ with an increased risk of sudden cardiac death.^{78,79} However, the majority of patients require hospital readmission after initiation of chronic intravenous inotropic therapy, even if begun with the hope of helping patients to stay at home until death. The use of intermittent infusions to control symptoms is currently not recommended by American Heart Association/American College of Cardiology guidelines (class III recommendation).¹⁷

A strategy to continue intravenous inotropic therapy for symptom relief and return home should not trigger implantation of an ICD, unless the patient is awaiting definitive therapy such as transplantation. The majority of patients on home inotropic infusions die by 6 months, and almost all are dead by 1 year, most often of terminal hemodynamic decompensation.

Renal Replacement Therapy

The prevalence of advanced kidney disease increases dramatically with worsening heart failure,⁸⁰ and measures of renal dysfunction are strong predictors of adverse outcomes in patients with heart failure (Table 3). Conversely, approximately 33% of individuals who commence hemodialysis have a recorded diagnosis of heart failure, and their mortality rates are significantly higher than patients who initiate dialysis without a heart failure diagnosis (adjusted relative risk of 1.22 in the US Renal Data System).⁸¹ Dialysis in the setting of advanced heart failure, especially in older patients with other comorbidities or frailty, has been shown to add to patient burden and in high-risk patients may not extend life.^{82,83} Therefore, the decision to initiate renal replacement therapies (eg, hemodialysis, ultrafiltration) in patients with advanced heart failure should only be made after a clear discussion with the patient about the risks and benefits of dialysis on the patient’s quality of life and prognosis.⁸⁴

Transplantation and Mechanical Circulatory Support: Exchange of Disease

Cardiac transplantation and mechanical circulatory support offer the potential to fundamentally change the clinical course of heart failure by exchanging it for surgical therapy and the need to adjust to living with a different set of benefits, risks, and burdens. In the case of transplantation, patients must adapt to the risks of organ rejection and immunosuppression and its side effects. For permanently implanted mechanical circulatory support, patients are dependent on a device with major complications of infection and stroke, as well as the potential for continued symptoms and required therapies for right-sided heart dysfunction. Thus, for eligible patients, whether to pursue these therapies represents one of the most difficult decisions that patients and clinicians can make. However, these therapies are limited to a highly selected group of patients. The use of cardiac transplantation is constrained by a limited supply of donor hearts, a situation that will not likely change in the foreseeable future. The use of mechanical circulatory support may increase as the technology improves but is likely to remain inappropriate for the majority of patients with heart failure because of the predominance of heart failure with normal ejection fraction, multiple comorbidities, or very advanced age.^{15,16} Detailed clinical practice guidelines are available that address the use of these advanced therapies.^{85–87}

Noncardiac Procedures in the Patient With Advanced Heart Failure

The risks and benefits of interventions for noncardiac conditions may be altered significantly in patients with advanced heart failure. When the likelihood of meaningful recovery without the procedure is small, the increase in procedural risk associated with heart failure may be considered acceptable. Examples include both emergent (eg, laparotomy for perforated viscus) and urgent (eg, hip arthroplasty for fracture) surgical procedures. Other procedures, such as knee replacement for degenerative joint disease, must be considered carefully in the context of patient preferences, because complications of the procedure may or may not outweigh the

potential benefit. Procedures should be discouraged when they do not offer a tangible improvement in quality of life (eg, repair of asymptomatic abdominal aortic aneurysm). Placement of permanent peritoneal and pleural catheters for the control of volume status is not indicated unless incorporated into a comprehensive palliative plan of care. Similarly, routine preventive care screening tests (eg, mammography, prostate-specific antigen) are typically not appropriate in the context of a significant competing risk of mortality caused by advanced heart failure, yet such tests are frequently ordered at the end of life.⁸⁸

Anticipating Decisions for Unanticipated Events

The process of clarifying preferences for cardiopulmonary resuscitation, intubation, feeding tubes, implantable defibrillator deactivation, intensive care unit transfer, and other near end-of-life interventions before the occurrence of a near-terminal event or acute-on-chronic decompensation is an important aspect of shared decision making. There is literature on the type and scope of these discussions,^{89,90} with suggestions to make them an annual event in routine medical care.^{60(p73406)} Such anticipatory discussions should include advance care planning guided by the concept that it is proper to “plan for the worst while hoping for the best.” It is also crucial for the care team to make clear that clinician abandonment will not occur; that is, the clinicians will work with the patient and family in downstream decision making and management.

Palliative Care

Palliative care is interdisciplinary care aimed at improving quality of life for patients by preventing and relieving suffering and supporting families.⁹¹ As such, it can be offered simultaneously with all other appropriate medical therapies. Palliative care is not synonymous with end-of-life care or hospice but can encompass them as the disease advances. Palliative care allows for continued disease-modifying therapies while ensuring symptom relief and interventions that address psychosocial, physical, and spiritual needs. This is done in 2 ways: by treating symptoms and by ensuring that patients' treatment plans match their values and goals.⁹²⁻⁹⁴ The process of shared decision making is a central tenet of palliative care: that the patient and clinician reach an understanding about preferences for life-prolonging therapy, symptom relief, pain control, and end-of-life care. Unlike hospice care (“Use of Hospice Services”), the application of palliative care is based on patient need rather than patient's prognosis or life expectancy.

Although data on palliative care in patients with heart failure are limited, several guidelines and reviews recommend integration of palliative care for all patients with advanced heart failure.^{19,89,90} This can and should be done by all clinicians involved in the care of these patients. However, referral to a palliative care team should be considered for assistance with difficult decision making and refractory symptom management in advanced disease, even as patients continue to receive disease-modifying therapies. Palliative care teams can consist of physicians, nurses, social workers, chaplains, and other professionals who work to ensure that

patient and caregiver needs are assessed and met. Because of the complex and changing nature of heart failure and the complexity of conversations as they change according to the patient's underlying heart disease, it is important to integrate palliative care into the care of patients with heart failure before they enter stage D. Even as patients are being considered for transplantation, mechanical circulatory support, or trials of novel therapeutics and pharmacological agents, palliative care can be increasingly integrated to ensure that patients' symptoms are appropriately controlled and that patients understand the nature of these interventions, as well as the full complement of alternative therapies.^{95,96} The synergistic relationship between palliative care services and the heart failure team for patients with mechanical circulatory support has been reviewed recently.⁹⁵

End-of-Life Care Planning

- Clinicians should take responsibility for initiating the development of a comprehensive plan for end-of-life care consistent with patient values, preferences, and goals.
- Deactivation of an ICD is desirable to avoid unnecessary pain and distress for patients and families at the end of life.
- Active discontinuation of mechanical circulatory support is often appropriate and necessary at the end of life.

Planning for Anticipated Death

Although the prognostic uncertainty inherent in heart failure makes it difficult to accurately anticipate the end of life, some patients enter a terminal phase of the disease that may be relatively apparent to the patients and/or their clinicians. In such situations, when the goals of care often transition from a focus on survival to quality of life and ensuring a good death, clinicians should take responsibility for initiating the process of putting into place a comprehensive plan of care consistent with patient values, preferences, and goals.

Passive Withdrawal of Therapies: Deactivation of ICDs

The option and ease of ICD deactivation should be discussed before implantation and again for major changes in clinical status (Table 4) or transitions in goals of care.⁹⁷ At present, this is done only rarely, thus leaving many patients vulnerable to inappropriate device discharge and unnecessary suffering. A recent survey found that only 1 in 4 next of kin reported that a physician had discussed device deactivation with their deceased family member before death.⁹⁸ In a nationwide survey of 734 physicians, including 292 cardiologists, 60% had fewer than 3 experiences discussing deactivation of ICDs with patients and/or families.⁹⁹ Concordant with those findings, a national survey of hospices found that <10% of hospices have a policy regarding deactivation of ICDs, and >50% of hospices had at least 1 patient who had been shocked within the past year.¹⁰⁰ For a device near its end-of-battery life, the generator should not be changed without careful review of whether or not active defibrillation is consistent with overall goals of care and anticipated duration of good-quality survival.

Active Withdrawal of Therapies When Patients Are Dependent on Them

Although the legal construct of patient autonomy does not recognize different degrees of dependence on therapies to be withdrawn, clinicians, patients, and families may view scenarios in which withdrawal leads to direct and rapid patient demise as unique and emotionally difficult. Examples include withdrawal of renal replacement therapy, feeding tubes, or pacemaker support for patients dependent on cardiac pacing. An increasingly common scenario is the withdrawal of mechanical circulatory support devices, either temporary or durable, in patients who are not expected to recover to return to a quality of life they consider acceptable.⁹⁵ The average life span of a patient after implantable left ventricular assist device (LVAD) placement has been increasing over time,¹⁰¹ but morbidity and mortality remain high. Estimated actuarial survival in the HeartMate II destination therapy trial was 58% at 2 years.¹⁰² With improvements in medical technology and associated outcomes, patients maintained with mechanical circulatory support may not only be susceptible to death attributable to cardiovascular causes but to other life-limiting disease as well. The discussion about discontinuing device therapy should be part of the “what if” informed consent process before implantation. Subsequently, although there are no specific recommendations to direct when these therapies should be discontinued, it appears that declining quality of life, signs of other organ system failure, or an irreversible catastrophic adverse event such as a major stroke or hemorrhage should trigger serious discussions about device deactivation. In a recent small study of characteristics of patients for whom mechanical circulatory support was electively discontinued, the most common triggers for discontinuation included sepsis, stroke, cancer, renal failure, and impending pump failure.²⁰³ Despite the perception that LVADs may impose special ethical dilemmas,⁹¹ the Patient Self-Determination Act still broadly applies, giving the patient, or their surrogate decision maker, full autonomy to withdraw support.

Device deactivation can be performed in the hospital or at home, attended by a device-trained individual (ventricular assist device coordinator) and others as requested (hospice nurse, chaplain, etc). Before device deactivation, a discussion with the patient (if able to participate) and family about the patient’s current condition and prognosis, changes in device benefit profile, how the device would be stopped, how symptoms would be managed, readiness to proceed, and anticipated outcome (ie, rapid death) is valuable. Unlike ICDs, which can be deactivated without immediate effect, LVAD discontinuation can result in rapid decompensation and expedite death, particularly with valveless continuous-flow devices. The average time to death after device deactivation is approximately 20 minutes, which indicates that a thoughtful discussion and careful plan should be in place well before the device is discontinued.⁷¹ This clinical scenario has been likened to withdrawal of endotracheal intubation and ventilatory support, although patients with LVAD support are more likely to be awake and alert at the time of decision to discontinue support. If patients are on multiple forms of support (eg, mechanical ventilatory support and LVAD sup-

port, with an ICD also in place), a coordinated plan to discontinue all of these therapies simultaneously is needed.

Use of Hospice Services

For patients approaching the end of life, hospice may be a viable option to provide symptom care and supportive services for patients and their families, while also ensuring that patients are able to die in their preferred environment. To be enrolled in the Medicare hospice benefit, 2 physicians or a physician and a nurse practitioner (one of whom is often the hospice medical director) must certify that the patient has ≤ 6 months to live if the disease follows its usual course, and the patient must be willing to forego usual medical services aimed at curing the underlying terminal diagnosis.¹⁰⁴ Most private insurers have a hospice benefit similar to that provided under Medicare. Although hospice is provided in a variety of environments,¹⁰⁵ it is most commonly provided for patients at home with the goal of keeping them in their home until death. Hospice can offer a number of benefits to enrollees and their families, including interdisciplinary team management, home visits, respite care, and provision of medications and durable medical equipment. Hospice also includes a nurse who can always be contacted to advise on urgent symptom needs and provide reassurance that interventions are appropriate.

Customized care plans may provide comfort and relief for some patients unwilling to accept formal hospice support. In many cases, patients feel they are “not ready for hospice,” and these patients should be referred to palliative care to ensure expert control of their symptoms, as well as support for the family. Likewise, continued education about the benefits of hospice and the fact that families are often more satisfied with hospice care than care provided in the hospital may also help elucidate its benefits.¹⁰⁶ One study of Medicare beneficiaries with heart failure who received hospice demonstrated a longer survival (by 81 days) than for those heart failure patients who did not receive hospice.¹⁰⁷

Hospice services have been shown to improve patient and family satisfaction with care. Families of those dying with hospice services were more likely to rate their dying experience as “favorable or excellent” than those who died in an institution or at home with only home health services.¹⁰⁶ Fewer than half of all patients with heart failure receive hospice. This is, however, a marked increase from $<20\%$ of heart failure patients being enrolled in hospice a decade ago.¹⁰⁸ Appropriate timing of referral to hospice is important, because the family’s perception of being referred “too late” is associated with greater dissatisfaction and unmet needs.¹⁰⁹

Communication and the Decision-Making Process

- Trust is the basis for the collaborative shared decision-making process.
- Early solicitation of values, goals, and preferences is necessary to guide the range of possible therapy options and decisions.
- Shared decision making is an iterative process that evolves over time as a patient’s disease and quality of life change.

- Assessment and integration of the emotional readiness of the patient and family are vital to effective communication.
- Ask-Tell-Ask provides a useful framework for communicating about prognosis and goals.
- Successful conflict resolution involves early recognition of conflict unfolding, with a shift in focus from winning an argument to trying to understand the reasons for conflict.
- Decision aids are tools that can enhance shared decision making by presenting numeric data in more understandable ways and assisting patients in clarifying their values.

Most patients and families want accurate and honest conversations with their clinicians.^{110,111} One study found that 93% of surrogate decision makers felt that avoiding discussions about prognosis was unacceptable.¹¹⁰ Advanced heart failure, with its high degree of prognostic uncertainty and complex trade-offs in the choice of medical care, demands a thoughtful approach to communication and decision making. Ideally, these interactions are not 1-time events but occur as an evolving series of discussions over time, particularly as a patient's condition changes. Such interactions may be difficult and time consuming, and they often require planning to create a supportive environment for effective communication. These discussions require careful attention to both mental and emotional needs. Clinicians must determine how much quantitative information patients want about prognosis, comparative risks, and benefits for both length and quality of life with available therapies. At the same time, clinicians must attend to the emotional nature of conversations with patients to build trust, clarify core values, and allow for sharper focus on the information. Here, we provide an overview of the tasks and skills, along with sample phrases, that can create high-quality shared decision making (Table 7).

Communication Is Desired, Beneficial, and Dynamic

Open, clear, and accurate communication with patients with heart failure is important for several reasons. First, the majority of patients with serious illness want information about their illness and to be included in the decision making process.^{116–118} Second, when clinicians have conversations with patients about their prognosis and desires, patients are more likely to receive care that is aligned with their goals and preferences.^{119–121} These conversations also improve the patient-clinician relationship.¹²² Finally, when conversations occur, families of deceased patients have better outcomes in terms of the manner in which they cope with loss of their loved one, as well as their own psychological outcome.^{110,121,123} One randomized trial of an advanced care planning intervention demonstrated that the intervention increased the likelihood that the patient's preferences were known and followed (86% versus 30%, $P < 0.001$) and decreased family members' stress, anxiety, and depression.¹²¹ Although many of the data about benefits of communication in patients with advanced illness are from the field of oncology, the evidence base demonstrating similar results in cardiology is increasing.¹²⁴

The biological reality of heart failure makes communication particularly difficult for 2 reasons. First, heart failure is

characterized by unpredictable periods of acute illness, followed by improvement in symptoms and function (Figure 1).^{19,21,125,126} Attending to this uncertainty involves both acknowledging the cognitive aspect of the conversation (eg, explaining to patients and families the unpredictable nature of illness and recognizing the inability of modern medicine to accurately predict life expectancy), while simultaneously addressing the complex emotions associated with the “roller coaster” of heart failure (eg, fear, anxiety, and uncertainty). Second, the chronic nature and unpredictability of heart failure require that communication be viewed as an evolving series of dynamic conversations that take into account the overall goals of the patient and family, the current state of health, and the shifting balance between benefits and burdens of any treatment or test that is either currently being used or that is being considered. Patients' preferences may change over time as their illness progresses and their experience with the disease changes, which further underscores the importance of an ongoing dialogue with patients and their families.^{39,56,127} For example, in 1 study of 936 patients with advanced heart failure, 19% had changed their preferences for resuscitation within 2 months.⁵⁶

To communicate effectively, clinicians must both determine and then readdress over time patients' understanding of their heart failure and their goals and treatment preferences (Table 4 for timing) and then determine how to have these conversations within the scope of clinical care. For example, when a patient is being seen for their first office visit after a hospitalization for heart failure, it may be useful to readdress goals of care from a global perspective, asking how the patient's thoughts about his or her heart disease have changed since the last hospitalization. In the case where emergent decision making is needed, the clinician might acknowledge how the conversation was last addressed and then bring up the specific emergent decision(s) at hand.

A Roadmap to Guide Conversations

Physician training in the conduct of these discussions is limited and needs to be fortified.^{128,129} Furthermore, the work of shared decision making belongs not only to physicians but to other members of the healthcare team as well, specific to their roles and responsibilities. Although additional training and mentored experience are clearly desirable, this section provides an introductory roadmap of how to effectively communicate with patients with advanced heart failure to facilitate the shared decision-making process. It is meant to represent an idealized version of communication, with the realization that this must be balanced with other competing responsibilities and clinicians' limited time. The goal is to offer effective strategies to improve conversations and decision making by demonstrating how complex conversations can be broken down into discrete elements, making them easier to accomplish. Not all conversations will include all of these elements every time. The goal is to offer a simple outline (along with some helpful phrases and tools) that may make conversations simpler for the busy clinician. More comprehensive explanations about how to communicate with patients with advanced illness are available.^{94,112,130}

Table 7. Core Tasks, Skills, and Sample Phrases to Improve Clinician-Patient Communication in Advanced Heart Failure

Steps in the Roadmap	Elements of the Step	Sample Phrases
Establish the setting and participants	Determine who should be present and ensure that all appropriate clinicians are present as well	"In preparation for our meeting tomorrow, I'm going to have the cardiothoracic surgeon there to be a part of our conversation. In terms of your family or support network, who is it important that we make sure is there?"
Determine what patients know and want to know	ASK what patients/families know	"Tell me about your heart disease; how have you been doing lately?" "What is your understanding of what is occurring now and why we are considering the treatment that we have been discussing?"
	ASK what patients/families want to know	"Sometime patients want to know all the details, whereas other times they just want to know a general outline. What kind of person are you?" "How much information would you like to know about what is happening with your heart disease?"
	TELL the patient/family the information in a sympathetic and thoughtful manner while also clearing up any misconceptions or unanswered questions ASK the patient or family to repeat back the information that has been delivered	"I think you have a pretty good understanding of what is happening with your heart, but there are a few points I'd like to review and clarify" "Now that I've clarified a few things about your illness, I want to make sure you understand what I've said. Tell me in your own words what we've been talking about"
Establish goals and preferences	Use open-ended questions to gain understanding of the patient's values to determine what is most important to them	"Help me to understand what is important to you. Some patients say they want to live as long as possible, regardless of quality of life. Sometimes patients tell me they are worried that they will be in a great deal of pain or have other uncontrolled symptoms. What is important to you at this point in terms of your health care?" "What are you hoping for?" "What is important to you now?" "What is your biggest concern right now?" "When you think about the future, what are the things you want to avoid?"
		In cases in which the patient is not involved in the conversation, a useful phrase might be, "What would your loved one say right now if he or she were hearing what we are discussing?"
Work with patient and family to tailor treatments and decisions to goals	Tailor explanation of benefits/burdens of a particular therapy based on goals established	"I think I understand what is important to you now, and it helps me better explain to you the decisions and treatments at hand now. I'd like to take a moment to review the benefits and burdens of each of the treatments based on what you've said is important to you at this point"
	Be willing to make a recommendation based on the patient's goals	"Would it be helpful if I made a recommendation based on what you've said the overall focus of care should be now?" "Based on what you have told me, if you get sicker and need to go back on a breathing machine again to stay alive, that is very unlikely to provide the kind of life you want to lead. Therefore, I think you should not go back on those machines"
	Acknowledge that there is uncertainty in the course of heart failure	"One of the most difficult things about heart disease is that we can never know for sure exactly what will happen in the next (hours, days, weeks, etc). We must make our best guess and decide what to do based on that information. If things change, we can always readdress this discussion at any time in the future"

Adapted from others.¹¹²⁻¹¹⁵

Where Are We on the Road? Ask-Tell-Ask to Determine What Patients Know and Want to Know

Before one can embark on conversations with patients and their families, it is important to establish the right context for the conversation. This includes asking whether the patient

wants to have the conversation by themselves or would like other individuals present, remembering that patients often define family in a myriad of ways. Creating the right setting also involves ensuring that the right clinicians are present, or at least have been consulted, before the conversation begins.

The individual leading the meeting ideally will have spoken to all the clinicians involved in the care of the patient so all points of view are represented and everyone is “on the same page” in terms of the illness and timely decisions.

Begin by asking the patient and family what they know and want to know. In this system, often called Ask-Tell-Ask,^{113,131} the clinician begins by asking patients and their families both what they know about their disease or the treatment being considered and how much information they want. Nearly 80% of patients want information about their illness, and this number rises as patients’ disease progresses. A Cochrane review of decision-making trials demonstrated that as patients learn more about the risks and benefits of therapies, the proportion preferring to take an active role in decision making increases to 85%.¹³² However, the only way to assess patients’ wishes is by asking and providing the patient this locus of control, which generates trust that is essential for collaborative decision making. An explicit way to ask is, “Would you want to know everything about your illness or the treatments we are considering, even if it wasn’t good news?” When patients and families express they do not want certain information, this should be explored further, with the explanation that information may be helpful for improving their understanding and to make sure that the decisions are consistent with the patient’s wishes. Denial and other defense mechanisms should not be ignored but instead carefully addressed and managed (“Emotional Roadblocks” below).

Once basic expectations for information exchange have been established, clinicians convey information to the patient and family in a clear and thoughtful manner, while also clearing up any misconceptions or unanswered questions they might have. This is the “Tell” in Ask-Tell-Ask. It is important to initially focus on the larger picture of the patient’s health, because the ability to cognitively hear information, particularly in stressful situations, is limited.¹³³ Giving all of the medical details may easily overwhelm the patient and may also lead him or her to focus on details that ultimately are not critical. The information should be delivered in simple language with frequent pauses to assess patient and family understanding.

The last “Ask” of the “Ask-Tell-Ask” process involves asking the patient or family to repeat back the information that has been delivered, to assess their understanding. This allows the clinician to determine the level of understanding the patient and/or family have and clarify any elements that may remain unclear in their minds.

The Ask-Tell-Ask technique is meant to be iterative and can be applied to many different levels of the process of communication. For increasingly complicated treatments and situations (eg, destination-therapy mechanical circulatory support), Ask-Tell-Ask is likely to be an extensive recurrent process that will occur over multiple encounters with the patient and family. Particularly complicated decisions may be augmented with decision aids (below).

Where Does the Patient Want the Road to Go? Establishing Values, Goals, and Preferences

One of the core elements of good communication is that it assesses patients’ values, goals, and desired outcomes, thus

allowing treatments and care to be as closely tailored to those desires as possible. This is especially true in light of the fact that physicians are often wrong about patients’ desires for care.⁵⁶ Optimal communication with patients with advanced heart failure does not begin with questions about treatments. Asking a patient during a routine office visit, “Do you want us to try and restart your heart?” is unlikely to be an effective starting point. This task is especially difficult because it involves weighing desired outcomes that may be contradictory (eg, avoiding severe disability while maximizing survival). This step then not only outlines what the patient hopes for but also considers complex trade-offs and situations the patient might consider a “fate worse than death.”

Open-ended questions to gain insight into the patient’s life and values are a useful method for initiating this portion of the conversation. Examples include, “What is important to you now?” or “What are you hoping for?” Another technique involves asking patients to discuss what is important in life outside of the hospital; this helps the clinician to understand what patients are doing in day-to-day life, how much patients value those tasks, and how patients view those tasks in the future. It also may be useful to inquire about a patient’s worries or concerns, using questions such as, “What is your biggest concern right now?” or “When you think about the future, what are the things you want to avoid?” In cases in which the patient is not involved in the conversation, either because of illness severity or because the patient chooses not to take part in the conversation, a useful phrase might be, “What would your loved one say right now if he or she were hearing what we are discussing?”

When advanced heart failure patients discuss their goals, they often discuss quality of life; that is, they typically are not only concerned about how long they will live but also how well they will live. This is especially true given that poor quality of life (limited ability to perform daily activities, significant symptom burden, emotional distress, and social isolation) is often reported by patients with advanced heart failure.^{42,134,135}

After clarifying the patient’s goals, it is often useful to summarize what has been expressed. In addition to ensuring that the clinician has heard and understood these hopes correctly, doing this also demonstrates care for the patient and that the clinician is attending to their needs. This may start with phrases such as, “Let me see if I understand what you are saying.”

Ensuring the Road Is Aligned With the Desired Destination: Tailoring Treatments to Goals

After goals have been clarified, the conversation can then move to discussing the role of specific treatments within the context of the desired outcomes. This involves working with the patient and family to (1) summarize the range of medically reasonable treatments for this particular patient at this particular time and then (2) explain the risks and benefits of each treatment option within the personalized rubric of goals and desires set forth by the patient and the family. Working within this context, the clinician helps the patient understand which treatments are most appropriate or inappropriate, based on their likelihood of getting the patient to the desired

outcome. This is, in fact, the core of shared decision making: The clinician does not dictate treatments, nor does the burden of the decision rest solely with the patient and family. Instead, the 2 parties work together to determine which options or treatments make the most sense given the patient/family's desired outcomes in the context of the current clinical scenario. In some cases, patients or families may be able to come to a decision on their own once the treatments and probable outcomes have been presented. However, even in these cases, families report they want to know what the physician would recommend.¹³⁶ In other cases, the patient and the family may signal they want more guidance (eg, "What would you do if it were your mother?") In these cases, it is appropriate to offer a recommendation based on the patient's stated goals (eg, "Given what you have told me about what is important to her, I think the treatment that makes the most sense to get her to the desired goal is ..."). Uncertainty about outcomes of specific treatments and outcomes should be communicated honestly and openly with patients and their families.

Tips for Navigating Barriers

These complex conversations with patients and families face serious challenges and barriers. This section outlines some of those challenges to inform busy clinicians and increase their awareness of them. Whenever possible, solutions and approaches with which to navigate these potential roadblocks are offered. These elements may be encountered (or solutions utilized) at any point in the decision-making process and thus operate in conjunction with the roadmap described above.

Acknowledging Emotional Roadblocks

Difficult decision making can stimulate complex emotions. Engaging patients in selecting treatments aligned with their informed goals and values requires that clinicians not only present the options clearly but that they also be attentive to patients' emotional needs. Patients are ill, caregivers are exhausted, and there may be a tremendous amount of fear, anxiety, stress, and perceived loss of control.¹³⁷ Neuropsychological studies have shown that when people are emotionally reactive, cognitive information is not processed accurately.^{130,133} One small study of family members of patients in the intensive care unit showed a relationship between communication styles and rates of anxiety in caregivers.¹³⁸ Attending to patients' emotions may improve their ability to process cognitive data and make better decisions. In addition, responding empathetically has been shown to strengthen the patient-clinician relationship, increase patient satisfaction, and make patients more likely to disclose future worries.¹³⁹

The first element of developing an emotional language for conversations is to recognize that patients are having an emotional reaction to the news that is being delivered and then learning to address the emotional content of the conversation. In one study of communication with family members of patients dying in an intensive care unit, clinicians missed opportunities to respond to the emotional content of the conversation in 29% of the conversations.¹⁴⁰ Although at times, patients' emotional reactions may be clear (especially when they use words such as "scared" or "angry"), at other

Table 8. Using the N-U-R-S-E Mnemonic to Help Express Verbal Empathy When Communicating With Patients With Advanced Heart Disease

Technique	Sample Language
Name the emotion	You seem worried about what will happen if we don't implant the LVAD. Can you tell me more about that?
Understand the emotion	I see why you might be fearful of proceeding with the transplant. Can you help me understand what you're afraid of?
Respect the emotion	You have shown a lot of strength up to this point. Tell me more about what keeps you going
Support the patient	Whether or not you choose to have the procedure, I want you to know that I will continue to be your cardiologist and will take care of you no matter what happens
Explore the emotion	You mentioned earlier that you're concerned about what this worsening of your shortness of breath might mean. Can you tell me more about your concerns?

LVAD indicates left ventricular assist device.

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times the exact emotional content of the discussion may be more veiled, such as when a patient or family says, "I don't know if I can handle this anymore." If patients or families keep raising the same issue repeatedly during a meeting, it may indicate that they are having a reaction to an issue that is so strong that it interferes with their ability to process information. For example, when patients ask a question such as, "How did this happen?" it may often be a clue that there is an emotional component to the information and that what is needed is a query about the patient's emotions and not a cognitive response explaining the cause of a disease or a complication. Addressing the reaction of patients and their families to serious illness is a metric that has been proposed to measure the quality of palliative care programs.⁹³

Once the clinician has recognized that there is an emotional component to the patient's or family's reaction, the next essential skill is to respond to it. Although there are data demonstrating that as little as 40 seconds of empathetic comments in conversations can improve patient and family outcomes related to communication, clinicians often need assistance in finding the right words to express their empathy for patients in the course of these complex conversations.^{141,142} One useful mnemonic device that can help clinicians respond empathetically in conversations is the mnemonic N-U-R-S-E.¹⁴¹ As explained in Table 8, NURSE stands for *N*aming the emotion expressed in the conversation, demonstrating *U*nderstanding of the emotion, *R*especting the emotion displayed by the patient or family, *S*upporting the patient/family, and *E*xploring the emotion in the context of the discussion. This assists clinicians in demonstrating verbal empathy and ensures that the complex emotional components of the conversation are addressed. More comprehensive reviews regarding the importance of acknowledging patient and family emotions have been published.^{143,144}

Depression and Anxiety

Depression and anxiety are common in patients with heart failure, with prevalence rates ranging from 13% to 77% and from 50% to 70%, respectively.^{145–147} Furthermore, prevalence rates for depression are almost 4-fold higher in patients with New York Heart Association functional class IV versus class I heart failure and also vary by self-reported symptom severity and health status.¹⁴⁸ These indicators of mental health status may affect the decision-making process. Depression is associated with impaired cognition and so can interfere with processing of information, memory, and executive function, which can affect decision making, especially in older adults.^{149–151} Anxiety, as noted above, can alter processing of information because of emotional reactivity. Thus, in patients with advanced heart failure, these barriers to decision making need to be identified and addressed to enhance discussions about therapeutic options. Screening for depression and anxiety, followed by pharmacological and nonpharmacological interventions (including psychological and/or psychiatric consultation), may be appropriate.

Limitations of Cognition, Literacy, and Numeracy

Current evidence suggests that in general, patients have a poor understanding of their medical interventions and that their preferences are not driving decisions.⁷ A survey of 3010 adults revealed that people were relatively unaware of the risks of 9 common medical conditions. For example, of patients taking a cholesterol medication, 38% did not know that the treatment was lifelong, and 83% could not correctly identify the most common side effect (muscle pain).¹⁵² In the case of ICDs, more than half of patients overrated the benefits of ICD therapy by 500%, thinking that >50 of 100 lives would be saved by the ICD therapy over the next 5 years (the actual estimate is closer to 5–10 per 100).¹⁵³ In the case of elective percutaneous coronary intervention, patients significantly overestimated the benefits; one study demonstrated that 88% of patients believed that percutaneous coronary intervention would reduce the chance of a recurrent myocardial infarction and 82% believed that it would reduce mortality,¹⁵⁴ despite clinical trial data that elective percutaneous coronary intervention has no effect on recurrent myocardial infarction or mortality.¹⁵⁵ Another study demonstrated that fewer than half of the patients could recall at least 1 complication of percutaneous coronary intervention despite the fact that most of them expressed a strong interest in participating in decision making. Together, these data suggest a need for improvement in the decision-making process.¹⁵⁶

Cognitive impairments compound difficulties with communication, comprehension, and decision making. Mild cognitive decline is seen in 25% to 50% of adults with heart failure.¹⁵⁷ A recent comparison of heart failure patients, healthy participants, and medical patients demonstrated that heart failure patients had poorer memory, psychomotor speed, and executive function than the other participant groups.¹⁵⁸ Almost one quarter of the heart failure patients had deficits in ≥ 3 domains of neuropsychological functioning. Those patients most likely to experience cognitive decline were those with the worst heart failure severity.

Limitations in health literacy and numeracy further interfere with understanding and integration of the information discussed as it relates to decision making. Health literacy is the degree to which people have the capacity to obtain, process, and understand the basic health information and services needed to make appropriate health decisions.¹⁵⁹ Nearly 10% of the population is functioning at below basic literacy levels.¹⁶⁰ Those most likely to be in the below basic level had less than a high school education, spoke no English before starting school, or were Hispanic, black, ≥ 65 years of age, or had multiple disabilities. Only 12% of the US population had the skills needed to manage their own health care proficiently.¹⁶¹

Not surprisingly, low health literacy is associated with poor self-care^{162–164} and increased mortality in older adults with chronic illnesses such as heart failure.^{165–167} To offset these poor outcomes, a recent scientific statement from the Heart Failure Society of America specifies that clinicians must recognize the consequences of low health literacy, screen patients at risk, document literacy levels and learning preferences, and integrate into practice effective strategies to enhance patients' understanding.¹⁶⁸ Furthermore, communicating numeric risk in a graphic form has also been shown to improve comprehension among patients with difficulties in literacy and numeracy¹³² ("Patient Decision Aids" at <http://decisionaid.ohri.ca/>¹⁶⁹ for examples).

Family Dynamics

Family dynamics can be a barrier to negotiating goals of care. A recent survey of elders (mean age 83 years) and their adult children (mean age 53 years) revealed that although most family units had discussed end-of-life preferences, important barriers to such discussions included fear of death, trust in others to make decisions, family dynamics, and uncertainty about preferences.¹⁷⁰ Those factors that facilitated the discussion were acceptance of the reality of death, prior experience with death, religion or spirituality, and a desire to help the family. In addition, there are significant psychological burdens associated with surrogate decision making.¹⁷⁰ Interestingly, previous discussion with their loved one about goals appeared to mitigate these burdens, providing an additional reason to raise these issues with both the patient and family early in the course of the illness.

Culture and Religion

Cultural and religious differences in patient preferences are known to exist.¹⁷¹ Awareness of cultural and religious differences can facilitate understanding of patient choices when discussing treatment options, especially when patients decline evidence-based therapies that healthcare professionals perceive as offering more benefit than risk. Although clinicians are not expected to be experts in cultural or religious issues relating to decision making, it is important that they be aware of the influence of these elements on decision making. Conversely, clinicians should also be warned against making assumptions about patient preferences based on perceived cultural or religious expectations. Clinicians should speak candidly with patients using the strategies recommended. Referral to palliative care, chaplaincy, or social work services

may help reveal existing religious and cultural differences as they relate to the decision-making process.

Language Differences

According to the US Census Bureau, the population speaking a language other than English at home has increased steadily for the past 3 decades.¹⁷² The number of languages and dialects spoken across the country presents major challenges to clinicians seeking to have a meaningful conversation about therapeutic options and end-of-life issues. Even among those patients who have learned English, when English is the second language, the subtleties and nuances of a discussion may be missed. Interpreters are often needed, which can be challenging in terms of finding someone linguistically capable, available, and sufficiently sensitive to communicate both the content and the tone.¹⁷³ Often, clinicians rely on family members to interpret for them, but family members have their own needs and emotions surrounding these conversations, which makes their use as interpreters potentially problematic.¹⁷⁴

Time

One of the largest barriers to these conversations is the time-intensive nature of these conversations, coupled with the time constraints faced by both patients and clinicians in a busy medical setting. Currently, clinicians can bill solely on the basis of time when >50% of the encounter (inpatient or outpatient) is spent on counseling the patient/families or coordinating care. Recent efforts aimed at making it easier for providers to be reimbursed for conversations about goal setting were ultimately abandoned for a variety of reasons.¹⁷⁵ Because of the complexity of these conversations, however, realignment of incentives to encourage providers to have these conversations and improve their skills for doing so is necessary and vital.

Resolving Conflict

In some cases, an intervention desired by a patient may appear discordant with the patient's stated goals and/or medical realities, and clinicians must explain why it is not warranted. This is particularly difficult in our national culture of entitlement and denial of morbidity and mortality. Clinicians must work with patients and their families to explain why a particular treatment is inconsistent with the overall goals of care, using patients' preferences as a rubric for why the treatment is not appropriate. These discussions can be emotionally charged and adversarial and may require considerable time. Given both the complexities of these conversations and their considerable length, formal involvement of palliative care teams has been shown to improve patient and caregiver satisfaction, effectively control symptoms, and decrease costs for patients with advanced disease.^{176–179}

There are 3 key elements that characterize discussions about not providing a therapy. First, the emphasis of the conversation should be on what treatments will still be provided that will help accomplish the patient's goals. This ensures the patient does not feel abandoned. Second, one should attend to the emotion behind the request. For example, "I wish I could tell you that doing [specific treatment] will accomplish the goals that you have outlined, but I'm sorry to have to say that it will not." In general, the "I wish" statement

can be beneficial in terms of acknowledging the emotional impact on patients of no longer having options for various therapies (ie, "I wish things were different"). Third, sometimes the patient or family has either misheard or misinterpreted the data that was presented, so it may be helpful to clarify what further information is needed to reconcile any inconsistencies (eg, "Tell me more about how you think CPR would help you"). By asking patients to clarify their reasoning, the clinician can more effectively address misunderstandings and inconsistencies. In this manner, the clinician attends to the emotions to better understand what is behind the request and then uses the NURSE strategy to acknowledge the emotion, rather than creating conflict with patients and their families about the treatment itself.

Decision Support to Assist With Particularly Difficult Conversations

In many cases, the decision at hand may be particularly complex or may require assistive methods to help patients and caregivers understand the potential outcomes and risks. In these cases, a decision support intervention, such as a decision aid or a decision coach, can help enhance conversations between patients and clinicians.

"Decision aids" are tools that help patients and caregivers become involved in decision making by providing information about the options and outcomes and by assisting patients in clarifying their personal values.^{10,180} Decision aids come in various forms, including booklets, pamphlets, videos, and Web-based systems,¹⁶⁹ and are designed to complement, not replace, a clinical encounter. They can be conceptualized broadly as either aids to assist the patient during or independent of the face-to-face encounter.¹⁰ A key difference between decision aids and a simple information pamphlet is that decision aids do not simply provide data about the anticipated risks and benefits but also provide guidance to help patients clarify their personal values and make a decision.¹⁸¹ Decision aids can also help patients clarify their values through a simple pros and cons list or an "imagined future" exercise.¹⁸² Decision aids attempt to present probabilities of the risks and benefits in ways that patients can understand. In fact, recent innovations have included the calculation and presentation of patient-specific outcomes generated from multivariable models (such as those listed in Table 3, among others), in routine clinical care. In the setting of informed consent for angioplasty, such a tool was demonstrated to improve patients' understanding of the risks of treatment, decrease anxiety, and improve satisfaction.¹⁸³ As new models are created that estimate a broader range of outcomes that are important to patients, this concept can be further developed, tested, and applied in advanced heart failure. A Cochrane review of 55 randomized trials of patient decision aids demonstrated that decision aids improved patient knowledge, reduced decisional conflict, increased patients' participation in decision making, and reduced the number of people remaining undecided with no associated adverse health outcomes.¹⁸⁴ However, only 1 trial in this Cochrane review was related to ischemic heart disease, and none were related to heart failure. The fact that substantial evidence suggests that decision aids help patients make better decisions combined with the fact

that patients with heart failure face a multitude of complicated decisions indicates that this is an area in need of significant development and research. The work on developing decision aids should begin with the higher-stakes decisions, including use of ICDs, inotropes, LVADs, and transplantation.

An alternative model to decision aids is the “decision coach,” a trained professional, often a nurse, who assists patients in making medical decisions by helping them prepare for a consultation and by empowering them to ask questions of their provider.¹⁸⁵ Early research suggests that coaching interventions may have modest effects on knowledge and participation in decision making.¹⁸⁶ The Ottawa Decision Support Guide is available for download and can be used for decision coaching (<http://decisionaid.ohri.ca/decguide.html>).¹⁶⁹ Although the use of coaches has not been studied in patients with advanced heart failure, nurses and other providers working with heart failure patients could be trained in decision coaching techniques.

Directions for the Future

- Changes in organizational and reimbursement structure will be necessary to reward and integrate decision making into the delivery of patient-centered health care.
- Mechanisms for standardizing, integrating, and distributing the work of shared decision making among the healthcare team should be developed and evaluated.
- Research to better define comparative functional and quality-of-life outcomes for the major therapies is vital to truly inform and align decision making with patient goals.
- Caregiver burdens and expectations should be assessed for all therapies and included as components of decision making.
- Content and form of decision aids should reflect contemporary outcomes data and interactive technology.
- Skill sets for communication and shared decision making should become part of standard curriculum and training for providers.
- Future research is needed in a variety of areas related to shared decision making, including effective communication training, decision support interventions, group visits, health-related quality-of-life measures, and caregiver burden, needs, and outcomes.

Need for Structural and Reimbursement Changes to Emphasize Shared Decision Making

Many busy clinicians may dismiss the above recommendations as impractical given the considerable time needed to complete the detailed communication processes outlined above. The diverse tasks of physicians, nurse practitioners, and physician assistants involved in primary care, general cardiology, and advanced heart failure management limit the capacity to conduct thorough prognostication, communication, and shared decision making for the various patient-centered outcomes, diverse patient and family preferences, and array of treatment options available to these complex patients. Yet, the unique role of clinicians demands that they

assume the primary responsibility for advancing shared decision making and promoting patient-centered care.

As such, the routine conduct of these activities must be efficiently integrated into routine care. The more clinicians perform shared decision making, the better they will be at making it a natural part of their routine practice of care. Patients and families could be prepared for important discussions before the clinic visit through a variety of possible mechanisms, facilitating better use of time for the busy clinician. Furthermore, palliative care services are an instrumental resource in helping with the more complex decisions and major milestones that arise in the course of advanced heart failure care. Multidisciplinary team-based care is necessary. Guidelines for the most dramatic therapeutic options, for instance, transplantation and mechanical circulatory support, already recognize the need for coordinated interdisciplinary care.^{85,87} Only through diverse inclusion of healthcare providers—nurses, nurse practitioners, physician assistants, primary care physicians, medical ethicists, chaplains, social workers, and others—can this shift adequately take place. A cultural change, particularly within cardiology but also more broadly, is necessary.

However, increased efficiency and dedication on the part of healthcare providers can only partially address the need for quality decision making. Ultimately, the US healthcare system primarily reimburses clinicians for doing things, not for deciding which things should and should not be done; both activities are time consuming and involve the expertise of a skilled clinician, yet only one is valued financially. For a healthcare system that has been criticized for overutilization, placing a greater emphasis on the shared decision-making process is likely to serve as a corrective force to achieve greater value from the system.¹⁸⁷ A start would be to reimburse clinicians more equitably for conducting a comprehensive annual heart failure review, which would include voluntary advanced care planning.^{60(p73406)} Unfortunately, attempts to specifically reimburse clinicians for these types of activities have met resistance for a variety of reasons.¹⁷⁵ Until these policy differences can be reconciled, actions like those by the Institute of Medicine to make shared decision making 1 of the pillars of quality care equate to an unfunded mandate in clinician time and energy.

Training Programs to Improve Communication With Patients With Heart Failure

Skills required to support patients in decision making are not adequately taught in training programs.¹²⁹ Multiple studies have shown variation and deficiencies in the ability of clinicians to communicate with patients and address end-of-life issues.^{188–190} Given the important yet difficult task of communication in clinical practice, improving communication should be a core element of the “performance-based” training and certification processes adopted by the Accreditation Council for Graduate Medical Education. The recent establishment of the secondary subspecialty of advanced heart failure and transplant cardiology¹⁹¹ by the American Board of Medical Specialties is an opportunity to formally add communication techniques and shared decision making to the training and certification process for physicians dedi-

cating their career to the care of these patients. Similar training is especially important for general internists and advance practice nurses who will be caring for a large number of patients with advanced heart failure. Requirements for institutional certification, such as The Joint Commission's Advanced Certification in Heart Failure program,¹⁹² offer yet another opportunity to emphasize formal processes that enhance shared decision making.

Several interventions have successfully improved communication skills for clinicians, particularly with regard to end-of-life care. Studies on the Education for Palliative and End-of-Life Care Project, a standardized multicomponent curriculum with an emphasis on improving communication, demonstrated an improved physician knowledge base and confidence in caring for patients at the end of life.^{193,194} Likewise, work with oncologists found that a program that combined training, role play, and individualized feedback improved participants' communication patterns, outcomes, and transitions across healthcare settings.^{195,196} Some nursing-centered interventions have been shown to be successful in changing nursing practice and improving outcomes for patients. The End-of-Life Nursing Education Consortium (ELNEC) program uses a train-the-trainer model, whereby nurses who complete the program are considered trainers who then go back to their home institutions and teach other nurses the core content aimed at improving care for patients with advanced diseases. Across the country, more than 4500 nurses have been enrolled in ELNEC,¹⁹⁷ and data from ELNEC have shown that the program improved nursing attitudes and knowledge about end-of-life care.¹⁹⁸

The successful programs to improve communication skills in end-of-life care provide a template for comprehensive "communication" training of heart failure clinicians and need to be expanded to the general physician and midlevel provider community. Additionally, mentoring trainees in patient and family meetings can help to ensure that clinicians have obtained the necessary skills to effectively communicate with heart failure patients and their families.

Future Research Directions to Improve Communication With Patients With Heart Failure

Methods for educating clinicians regarding communication and shared decision making remain early in their development. Useful decision aids for commonly encountered medical decisions in heart failure are also largely underdeveloped or unavailable. One company has developed decision aids for patients considering ICD and CRT therapy (<http://www.healthwise.org>),^{198a} and another has developed a general decision aid for patients with heart failure (<http://www.informedmedicaldecisions.org>),^{198b} but these have not been studied formally in real-world settings. Several recently funded National Institutes of Health grants are designed to develop and evaluate decision aids among patients with heart failure, with the recognition that the results of this work are several years away.

At a more basic level, our understanding of how patients with advanced heart failure make choices is limited. There is also no consensus in the literature on the best way to measure whether a medical decision was a "good" one.¹⁹⁹ Decisional

quality, defined as "the extent to which the implemented decision reflects the considered preferences of a well-informed patient,"²⁰⁰ is emerging as another possible measure to assess the quality of decision making, but validated measures to quantify decisional quality are in the early developmental stages.²⁰⁰ An ideal metric to measure decision quality would include domains in knowledge and values and a way to measure value-treatment concordance. Despite lingering questions, decision quality measurement is gaining popularity, and some have proposed that measures of decision quality be included as part of the larger pay-for-performance agenda.¹⁸⁷ The Patient Protection and Affordability Care Act calls for "the development of quality measures that allow for assessment of the experience, quality, and use of information provided to and used by patients, caregivers, and authorized representatives."¹³

The role of palliative care in patients with advanced heart failure has been far less developed than in cancer,²⁰¹ and further work to document the synergistic effect of adding palliative care to the clinical care of patients with advanced heart failure is needed. Our understanding of health-related quality of life—"the functional effect of an illness and its consequent therapy on a patient, as perceived by the patient"³⁸—for patients with advanced heart disease is limited as well.²⁰² Although health-related quality-of-life measurements have been developed for patients with symptomatic heart failure,^{203,204} questions remain about their sensitivity in very advanced stages of disease. There has been some interest in developing self-report instruments to assess quality of life at the end of life, but they have not been thoroughly tested and validated.²⁰⁵ Primarily, there has been relatively slow uptake of health-related quality-of-life instruments in the evaluation of therapies and in the routine care of patients. Our understanding of the burden and quality of life of caregivers of heart failure patients is even more limited, as is knowledge about how best to intervene to maximize caregiver quality of life.

Conclusions

The importance of shared decision making in advanced heart failure cannot be overstated given the complex myriad of treatment options that confront patients, families, and caregivers. We have offered a roadmap for when and how to have conversations with patients to support shared decision making. This process must occur in the context of uncertainties in prognosis, multiple and often competing outcomes, and barriers to communication. Although the promotion of shared decision making may seem daunting to busy practicing clinicians, we have attempted to provide guiding principles and simple tools that can help set future expectations, anticipate major decisions, and promote productive conversations. Our statement is a "call to action," not only to clinicians within our community directly responsible for facilitating shared decision making but also to those on a national level who would reform and restructure the healthcare medical system to truly support patient-centered care.

Disclosures

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*Modest.

†Significant.

Reviewer Disclosures

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KEY WORDS: AHA Scientific Statements ■ communication ■ decision making ■ heart failure ■ heart-assist device ■ palliative care ■ prognosis ■ transplantation

Decision Making in Advanced Heart Failure: A Scientific Statement From the American Heart Association

Larry A. Allen, Lynne W. Stevenson, Kathleen L. Grady, Nathan E. Goldstein, Daniel D. Matlock, Robert M. Arnold, Nancy R. Cook, G. Michael Felker, Gary S. Francis, Paul J. Hauptman, Edward P. Havranek, Harlan M. Krumholz, Donna Mancini, Barbara Riegel and John A. Spertus

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Correction

In the article by Allen et al, “Decision Making in Advanced Heart Failure: A Scientific Statement From the American Heart Association,” which published ahead of print on March 5, 2012, and appeared in the April 17, 2012, issue of the journal (*Circulation*. 2012;125:1928–1952), several corrections were needed:

1. On page 1928, in the endorsements, the Society for Medical Decision Making was added. The endorsement read, “Endorsed by Heart Failure Society of America and American Association of Heart Failure Nurses.” It has been changed to read, “Endorsed by Heart Failure Society of America, American Association of Heart Failure Nurses, and Society for Medical Decision Making.”
2. On page 1949, in the References, the journal name in reference 41 read, “*Eur Heart J*.” It has been changed to read, “*Eur J Heart Fail*.” The corrected reference now reads,
 41. Dev S, Clare R, Felker GM, Fiuzat M, Stevenson LW, O’Connor CM. Link between decisions regarding resuscitation and preferences for quality over length of life with heart failure. *Eur J Heart Fail*. 2012;14:45–53.

These corrections have been made to the print version and to the current online version of the article, which is available at <http://circ.ahajournals.org/content/125/15/1928>.

Executive Summary: Decision Making in Advanced Heart Failure

A Scientific Statement From the American Heart Association

Endorsed by Heart Failure Society of America, American Association of Heart Failure Nurses, and Society for Medical Decision Making

Larry A. Allen, MD, MHS, Co-Chair; Lynne W. Stevenson, MD, Co-Chair; Kathleen L. Grady, PhD, APN, FAHA, Co-Chair; Nathan E. Goldstein, MD; Daniel D. Matlock, MD, MPH; Robert M. Arnold, MD; Nancy R. Cook, ScD; G. Michael Felker, MD, MHS; Gary S. Francis, MD, FAHA; Paul J. Hauptman, MD; Edward P. Havranek, MD; Harlan M. Krumholz, MD, SM, FAHA; Donna Mancini, MD; Barbara Riegel, DNSc, RN, FAHA; John A. Spertus, MD, MPH, FAHA; on behalf of the American Heart Association Council on Quality of Care and Outcomes Research, Council on Cardiovascular Nursing, Council on Clinical Cardiology, Council on Cardiovascular Radiology and Intervention, and Council on Cardiovascular Surgery and Anesthesia

Decision making for advanced heart failure has become more crucial and more challenging as the prevalence of symptomatic disease and the complexity of therapeutic options have increased. Once heart failure advances despite recommended medical therapies, a small, highly select group of patients are faced with a decision to exchange the syndrome of heart failure for an advanced surgical therapy (ie, heart transplantation and/or mechanical circulatory support) with an entirely different set of benefits, risks, and burdens. The vast majority of patients with advanced heart failure, however, are faced with day-to-day decisions in the context of an extended period in which increasing symptoms adversely affect their daily lives, as well as the lives of their caregivers. Shared decision making asks that clinicians and patients share information with each other and work toward decisions about treatment that are aligned with patients' values, goals, and preferences and are feasible given existing medical evidence (Table).

The ethical principle of autonomy recognizes the rights of patients to choose their therapies from among reasonable available options. The formal process of informed consent includes not only the procedural details but also comprehension of the benefits and risks of the offered therapy and available alternatives, including continuation and withdrawal of ongoing treatments. Shared decision making extends beyond informed consent, requiring that clinicians and patients consider information together and work toward consensus. Shared decision making thus follows the principle of "patient-centered care," 1 of the 6

pillars of healthcare quality identified by the Institute of Medicine. The implication is that beneficial therapies and those recommended by guidelines should be offered but need to be discussed as an option among many strategies, with attention given to how they are aligned with patient preferences.

Anticipation is a central tenet of shared decision making, particularly in the setting of a progressive condition with high rates of morbidity and early mortality. Setting expectations for the future for individual patients with advanced heart failure requires attention to the clinical course, including milestones such as recurrent hospitalizations and decreased effectiveness of chronic medical regimens. Marked clinical fluctuations characterize heart failure more than other chronic diseases, with inevitable uncertainty in prognosis. Although predictive models can target high-risk populations, the probability of greater than a 2-fold error rate in either direction for a survival estimate applied to an individual patient (such as predicting survival of <6 months or >2 years for a patient who dies in 12 months) remains near 50% under realistic assumptions. However, this uncertainty should not deter discussions with patients and families but rather should be acknowledged as an inherent part of decisions regarding future care.

Difficult discussions conducted earlier will simplify difficult decisions in the future. We advocate for an annual heart failure review between patients and their primary clinicians, which would encompass discussion of current and potential therapies for both anticipated and unanticipated events (including cardiac

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Table 1. Top Ten Things to Know

1. Shared decision making is the process through which clinicians and patients share information with each other and work toward decisions about treatment chosen from medically reasonable options that are aligned with the patients' values, goals, and preferences.
2. For patients with advanced heart failure, shared decision making has become both more challenging and more crucial as duration of disease and treatment options have increased.
3. Difficult discussions now will simplify difficult decisions in the future.
4. Ideally, shared decision making is an iterative process that evolves over time as a patient's disease and quality of life change.
5. Attention to the clinical trajectory is required to calibrate expectations and guide timely decisions, but prognostic uncertainty is inevitable and should be included in discussions with patients and caregivers.
6. An annual heart failure review with patients should include discussion of current and potential therapies for both anticipated and unanticipated events.
7. Discussions should include outcomes beyond survival, including major adverse events, symptom burden, functional limitations, loss of independence, quality of life, and obligations for caregivers.
8. As the end of life is anticipated, clinicians should take responsibility for initiating the development of a comprehensive plan for end-of-life care consistent with patient values, preferences, and goals.
9. Assessing and integrating emotional readiness of the patient and family is vital to effective communication.
10. Changes in organizational and reimbursement structures are essential to promote high-quality decision making and delivery of patient-centered health care.

and respiratory arrest) within the context of patient values, goals, and preferences. In addition to the annual review, a more focused discussion should be triggered by milestones for review, such as hospitalizations, defibrillator shocks, progressive hypotension or renal dysfunction, significantly increased symptom burden, and limitations in functional capacity and independence.

A framework for types of decisions in advanced heart failure includes (1) cardiac interventions such as valve surgery, percutaneous catheter intervention, or cardiac resynchronization therapy that may improve cardiac performance; (2) implantable defibrillators that reduce the risk of sudden death but do not improve cardiac performance; (3) temporary options for acute decompensation such as intravenous inotropic therapy or hemodialysis that may lead to unanticipated dependence; (4) the surgical therapies of transplantation or mechanical cardiac assist devices that exchange diseases and set patients on different life trajectories; and (5) noncardiac procedures, for which benefits and risks should be reviewed in the context of competing risks from heart failure.

Discussions should include a range of anticipated outcomes, including not only survival but also major adverse events, quality of life, symptom burden, functional limitations, loss of independence, and obligations for caregivers. Major interventions should be presented with a balanced description of alternative approaches, including a focus on symptomatic therapy. Referral to a palliative care team should be considered for assistance with difficult decision making and symptom management, even as patients continue to receive disease-modifying therapies. The use of palliative care services should not be considered equivalent to the withdrawal of disease-modifying therapies.

Planning end-of-life care for anticipated death is a "time of action"; clinicians should take responsibility for initiating the development of a plan of care consistent with the patient's expressed values and goals. Part of the final plan of care may involve consulting palliative care professionals and recommending hospice services, because they can provide effective symptom management for the patient and support for the family, most often in the home.

Communication is essential to the decision-making process. Research has shown that most patients and families want accurate and honest conversations with clinicians; however, these complex conversations face multiple potential barriers. Difficult decisions about life and family stimulate powerful and complex emotions that may prevent processing of important information. Attention to informational preferences, learning styles, health literacy, anxiety and depression, cognitive limitations, cultural and religious differences, and language barriers is critical to facilitate informed discussions. Family and caregiver dynamics, particularly with regard to surrogate decision making, can further complicate the goal of matching therapy to patient values and preferences. Conflict can arise when an intervention desired by the patient or family is not aligned with medical realities or the patient's stated goals.

To navigate these barriers, clinicians must be highly skilled in communication. A roadmap for effective communication can facilitate these discussions. Ideally, shared decision making is an iterative process that evolves over time as a patient's disease and quality of life change. Early elicitation of values, goals, and preferences is necessary to guide future discussions of possible therapeutic options and decision making. Specific skills and tools that clinicians can integrate into their practice include the Ask-Tell-Ask format for communicating difficult information, the N-U-R-S-E mnemonic (Naming the emotion expressed in the conversation, demonstrating Understanding of the emotion, Respecting the emotion displayed by the patient or family, Supporting the patient/family, and Exploring the emotion in the context of the discussion) for dealing with complex emotions, and decision aids for enhancing the communication of difficult quantitative information and integration of patient values, goals, and preferences. Referral to palliative care, chaplaincy, and psychological or social work services may also help to identify and overcome these barriers.

Future efforts to improve the process of shared decision making should include an enhanced research focus on the full range of anticipated outcomes for possible interventions, skills training in clinician communication, and development of decision support tools and decision quality measures. Although clinicians must assume the primary responsibility for advancing shared decision making and patient-centered care, these efforts will be restricted in scope until the healthcare system shifts its emphasis from reimbursing clinicians for providing specific therapies to reimbursing clinicians for helping patients decide which strategies should and should not be pursued; society must realign incentives to support shared decision making and a culture of patient-centered health care.

References

References are available in the full text of this guideline: <http://circ.ahajournals.org/lookup/doi/10.1161/CIR.0b013e31824f2173>.