SPECIAL ARTICLE

Achieving High-Value Cardiac Imaging: Challenges and Opportunities

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Cardiac imaging is under intense scrutiny as a contributor to health care costs, with multiple initiatives under way to reduce and eliminate inappropriate testing. Appropriate use criteria are valuable guides to selecting imaging studies but until recently have focused on the test rather than the patient. Patient-centered means are needed to define the true value of imaging for patients in specific clinical situations. This article provides a definition of high-value cardiac imaging. A paradigm to judge the efficacy of echocardiography in the absence of randomized controlled trials is presented. Candidate clinical scenarios are proposed in which echocardiography constitutes high-value imaging, as well as stratagems to increase the likelihood that high-value cardiac imaging takes place in those circumstances. (J Am Soc Echocardiogr 2014;27:1-7.)

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Cardiac imaging has come under intense scrutiny as a contributor to rising health care costs in the United States. Attention has been focused on the number of cardiac imaging studies performed, including echocardiography. Volume is easy to measure; a far more difficult, and more important, task is to ascertain the value of imaging for specific patients or groups of patients. The critical issue is not how many studies are being done but that they are done in circumstances in which the results will enhance the patient care—and not done when the results will not make a difference—so that studies lead to better outcomes.

Increased demand for testing is due to both patient-related and physician-related factors.^{1,2} Among the drivers are physician training that encourages a culture of completeness regardless of cost or of effects on others; misaligned financial incentives; effective marketing of new technologies to physicians in the absence of comparative effectiveness data with which physicians can assess the value of that technology; and fear of malpractice suits, encouraging the practice of defensive medicine. On the patient side, Americans are enamored of high technology and may perceive that more tests are by definition equal to better care. Direct-to-consumer marketing influences patients' preferences for testing, and a health care system in which patients are insulated from the true fiscal costs of testing also drives demand.

Recent data indicate that the rate at which cardiac imaging is performed not only is no longer increasing but has begun to drop. While the US General Accounting Office reported in 2008 that Medicare spending on imaging services under the Part B physician fee schedule more than doubled from 2000 through 2006, a subsequent Medicare Payment Advisory Commission report to Congress noted that annual rate of growth in the number of echocardiograms provided per Medicare beneficiary was only 2.6% between 2005 and 2009 and decreased by 0.8% per year between 2009

Copyright 2014 by the American Society of Echocardiography. http://dx.doi.org/10.1016/j.echo.2013.08.027 and 2010.³ On the cost side, payments to cardiologists for noninvasive diagnostic imaging decreased by a total of 33% between 2006 and 2010, reversing the increases seen during the preceding 6 years.⁴ Multiple explanations have been cited for this phenomenon, which is sometimes referred to as "bending the cost curve." They include the promulgation of appropriate use criteria for cardiac imaging by professional societies such as the American College of Cardiology Foundation (ACCF) and the American Society of Echocardiography, among others. These documents evaluate the relative benefits and risks of an imaging study to determine whether it is reasonable to consider performing the study for a specific indication.⁵ The terminology used to describe the three appropriateness categories has evolved for greater clarity since their original publication. Studies for specific indications were initially divided into appropriate, uncertain, and inappropriate categories. The terminology has been revised to "appropriate care," "may be appropriate care," and "rarely appropriate care," recognizing that a study that is rarely appropriate may be precisely correct for a specific patient.⁶ Stated another way, the goal for rarely appropriate studies is not zero. Education programs such as the American Board of Internal Medicine's Choosing Wisely campaign have been directed at patients and providers. Commercial insurers have turned to radiology benefits managers in an attempt to reduce test ordering they deem inappropriate, while Medicare has adopted payment reductions to providers.

REDUCING OVERUTILIZATION

The interest in limiting inappropriate cardiac testing stems not just from containing costs. Excess testing carries the potential for downstream ill effects. When a study that may have good specificity is ordered in a population in which a disorder has a low prevalence, the few "abnormal" results are more likely to be false-positives than true-positives. This can cause anxiety on the part of patients and lead to unwarranted further testing, which carries its own inherent risks. Conversely, a false-negative result provides false reassurance and the potential for delayed diagnosis. These concepts are explicitly recognized by the ACCF in its definition of an appropriate imaging

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Abbreviation

ACCF = American College of Cardiology Foundation

study as "one in which the expected incremental information, combined with clinical judgment, exceeds the expected negative consequences by a suffi-

ciently wide margin for a specific indication that the procedure is generally considered acceptable care and a reasonable approach for the indication."⁵

Appropriate use stratagems have been used to examine and vet imaging studies once they have been ordered, to determine whether they are being ordered for appropriate reasons. Methodologies focusing on studies after they have been ordered are suited to reducing overutilization. Research in community as well as academic settings has shown that 9% to 20% of transthoracic and stress echocardiographic studies are ordered for inappropriate indications.⁷⁻¹¹ A much smaller proportion of requested transesophageal studies is rated as inappropriate.¹² The reasons for the disparity have not been studied but might include differences in specialties of the ordering physicians (i.e., cardiologists vs noncardiologists) for transesophageal studies compared with transthoracic or stress echocardiography. The ease with which a transthoracic or stress echocardiographic study can be ordered, contrasted with the fact that transesophageal studies are semi-invasive and are directly performed by cardiologists who must actively assent to their performance, may play a role in differing rates of appropriateness. Applying appropriate use criteria had previously been a manual undertaking, consisting of matching the clinical scenario to a list of criteria on paper and uncovering the appropriateness score. An application for myocardial perfusion imaging is available for both major smart phone platforms, and one for echocardiography has been announced. The American College of Cardiology has designed Imaging in FOCUS, a voluntary, Web-based decision support program designed to reduce inappropriateness in cardiac imaging. FOCUS demonstrated a reduction in inappropriate single-photon emission computed tomographic myocardial perfusion imaging ordering among participants, from 11% of studies before using FOCUS to 5% afterward.¹³ The American Society of Echocardiography has codeveloped a FOCUS module for transthoracic echocardiography. It is reasonable to expect comparable improvements in the degree of study appropriateness when this tool is applied to transthoracic echocardiography, but this hypothesis has yet to be tested beyond a pilot study.¹¹ FOCUS is evolving into a robust, multimodality program that links with commercially available electronic health records and provides integrated decision support at the point of order entry.¹⁴

DEFINING AND IDENTIFYING HIGH-VALUE IMAGING

In the quest for high-value imaging, rooting out cardiac imaging studies that are of questionable appropriateness by looking at the studies is one part of the solution. However, if examining appropriateness begins once a test has been ordered, the process is entered at the midpoint of the dimensions of care framework for evaluating the quality of cardiac imaging described by the ACCF (Figure 1). This framework starts with the patient, recognizing that efforts at enhancing the value of imaging studies must be patient centered rather than test centered. Focusing efforts at the patient level uncovers not only which patients do not need an imaging studies to detect or risk-stratify diseases. Such high-value imaging may lead to management changes that improve outcomes or, alternatively, lead to the



Figure 1 Dimensions of care framework for evaluating quality of cardiovascular imaging. Reproduced with permission from Douglas P, Chen J, Gillam L, Hendel R, Jollis J, Iskandrian AE, et al. Achieving quality in cardiovascular imaging: proceedings from the American College of Cardiology-Duke University Medical Center Think Tank on Quality in Cardiovascular Imaging. J Am Coll Cardiol 2006; 48:2141-2151.

imaging study that most conclusively and efficiently excludes a disease, thereby reducing both patient anxiety and downstream costs. This approach might better be conceptualized as "bending the value curve," because the goal of managing cardiac imaging is not just lower costs but higher value to patients and the health system. The concept of developing an outcomes-based imaging cycle backed by evidence is not new¹⁵ but bears explication, particularly as the American health care system continues to transform.

Value in health care has been defined as health outcomes achieved per dollar spent.¹⁶ Determining what is high-value cardiac imaging requires measurable outcomes that are specific to a given condition. Outcomes, in the numerator, must be achieved efficiently; that is, the total cost of care for the condition must be calculated, and not merely the cost of an individual service. A more expensive test that reduces the overall cost of care may be a good investment of health care dollars. Diagnostic studies do not by themselves cure, or change outcomes. Yet high-value imaging, by being performed in the correct part of the care cycle, conceptually can reduce the overall cost of care if it leads to a better health outcome. Although the most critical outcomes for patients are increased survival, and recovery or improved health, other metrics include time to recovery, avoiding treatment-related side effects, avoidance of complications, sustained health and function, and avoiding care-induced illnesses.

The highest level of evidence for the value of an imaging study would come from a randomized controlled trial that measures specified outcomes. An example of such a study is the Prospective Multicenter Imaging Study for Evaluation of Chest Pain trial, a randomized trial funded by the National Heart, Lung, and Blood Institute of the clinical effectiveness of diagnostic strategies in patients with chest pain, who are randomized to either functional (exercise electrocardiography, stress echocardiography, or stress nuclear imaging) testing versus anatomic testing (coronary computed tomographic angiography).¹⁷ Randomized trials for an accepted technology that is already in clinical use, such as echocardiography, as part of a diagnostic and treatment strategy are unlikely to be conducted because of the large number of conditions for which echocardiography is performed and perhaps also because of the lack of sponsor enthusiasm for investing in what are perceived to be mature technologies.

An alternate, frequently cited paradigm to judge the value of imaging uses a six-tiered, hierarchical model to conceptualize diagnostic imaging as part of a larger system whose goal is to treat patients effectively and efficiently. Level 1 is technical efficacy, comprising variables needed to produce a high-quality image. Level 2 is diagnostic accuracy efficacy, such as the percentage of correct diagnoses, positive and negative predictive value, sensitivity and specificity, as well

Table 1 Hierarchical model for the value of imaging¹⁸

Level 1: technical efficacy

- Variables needed to produce a high-quality image
- Level 2: diagnostic accuracy efficacy
 - Percentage of correct diagnoses (diagnostic accuracy)
 - Positive and negative predictive value
 - Sensitivity and specificity
 - Area under the receiver operating characteristic curve
- Level 3: diagnostic thinking efficacy
 - Percentage of cases in which imaging was helpful in making the diagnosis
 - Difference in the clinician's estimation of posttest vs pretest probability

Level 4: therapeutic efficacy

- Percentage of times imaging affects management or changes the diagnostic or therapeutic plan
- Percentage of times a medical procedure is avoided because of imaging information

Level 5: patient outcome efficacy

- Percentage of patients who improve after the test compared with those without it
- · Morbidities avoided

Change in QALYs deriving from the test; cost per QALY saved

- Level 6: societal efficacy
 - Benefits and costs of the imaging strategy from the societal viewpoint

QALY, Quality-adjusted life-year.

as receiver operating characteristic curves. Level 3, diagnostic thinking efficacy, describes the percentage of cases in which imaging was helpful in making the diagnosis, the difference in the clinician's estimation of posttest versus pretest probability, and the like. Level 4, therapeutic efficacy, consists of the percentage of times imaging affects management or changes the diagnostic or therapeutic plan. Level 5 describes patient outcome efficacy, being the percentage of patients who improve after a test compared with those without it, morbidities avoided, change in quality-adjusted life-years derived from the test, and similar metrics. Finally, level 6, societal efficacy, comprises the benefits and costs of the imaging strategy from a societal viewpoint¹⁸ (Table 1).

The value of cardiac imaging can be assessed if information about the higher levels of efficacy cited in the model is available. For a given condition, an imaging study can lead to changes in diagnostic thinking, such as uncovering the presence and severity of a disorder. Conversely, a test may confirm the absence of a condition, such a decline in left ventricular systolic function during a course of cancer chemotherapy, allowing treatment to continue as planned by the treating clinician. The test results bring about an evidence-based change in management that has been proved in randomized, controlled trials to improve patient outcomes. This approach might constitute an adequate surrogate for randomized, controlled trials of the imaging modality itself.^{19,20}

Aspects of the candidate imaging study itself also enter into the value equation. In all forms of testing, a good candidate predictor should have a favorable risk-benefit ratio, reasonable cost, acceptability, and convenience, all of which are characteristic of echocardiography.²¹ With respect to candidate conditions, a "commonsense checklist" would consist of the following. One should apply the predictor to diseases with major morbidity, for which some effective treatment is available that is not equally effective for all persons. The candidate test should allow more accurate classification of indi-

viduals into categories in which treatment is or is not indicated. The incremental prediction should be beyond what can be achieved with information that is already available. There should be consensus about and standardization of established predictors, and the predictor should be unambiguously defined and measured.²¹

OPPORTUNITIES TO ACHIEVE HIGH-VALUE CARDIAC IMAGING

Examples of cardiac conditions with significant morbidity that fit these criteria abound. In each case, echocardiography reclassifies patients noninvasively, painlessly, and without the use of ionizing radiation, on the basis of standardized criteria such as those for chamber quantification, stress echocardiography and valvular regurgitation that have been promulgated in a series of guidelines published by the American Society of Echocardiography and the ACCF.²²⁻²⁴

An example candidate condition is heart failure, which affects an estimated 5.7 million people in the United States.²⁵ Many class I recommended therapies for heart failure with reduced left ventricular ejection fraction are supported by level of evidence A; that is, they are recommended on the basis of multiple randomized controlled trials or meta-analyses conducted in multiple populations. Therapies including angiotensin-converting enzyme inhibitors or angiotensin receptor blockers, β -blockers, implantable cardioverter-defibrillators for primary and secondary prevention of sudden cardiac death, and cardiac resynchronization have been shown to reduce symptoms, decrease hospitalizations, and increase survival, depending on stage and symptom class. Treatment of systolic heart failure is predicated on identifying the clinical predictor decreased systolic left ventricular function, usually via two-dimensional echocardiography, which is described as "the single most useful diagnostic test in the evaluation of patients with [heart failure]."26

Chronic mitral regurgitation serves as another example. This disorder is characterized by a long latent period in which patients can remain asymptomatic even in the face of developing left ventricular dilation and systolic dysfunction; yet even patients with preserved left ventricular ejection fraction may be at increased risk for death. Mitral valve surgery performed in an asymptomatic individual whose ejection fraction has fallen below 60% is likely to prevent further deterioration in left ventricular function and improve longevity, although the level of evidence is less robust (level B: limited populations evaluated; data derived from a single randomized trial or nonrandomized studies). Once the ejection fraction falls even lower, the risk for surgery increases, and postoperative survival is less. Thus, there exists a golden moment for patients with chronic severe mitral regurgitation, wherein the identification of the severity of regurgitation and tracking left ventricular systolic function lead to surgical therapy, which improves outcomes, and conversely whereby outcomes are appreciably worse if the golden moment is missed.²⁷

Other conditions for which opportunities exist to achieve highvalue cardiac imaging leading to effective, evidence-based treatments are listed in Table 2.^{26,28-31}

ENHANCING THE LIKELIHOOD THAT HIGH-VALUE CARDIAC IMAGING IS PROVIDED

The risk for missed opportunities might increase as health care in the United States reorganizes into episodes of care for a specific patient,

Condition	Echocardiographic parameter	Outcome	Class	Level of evidence	References
Valve disease					
Aortic regurgitation	LVEF < 50%, asymptomatic	Surgery	1	С	27
	LVEDD > 75 mm or LVESD > 55 mm, asymptomatic	Surgery	lla	С	27
Aortic stenosis	LVEF < 50%, asymptomatic	Surgery	1	С	27
Prevention of sudden cardiac death					
Nonischemic CM	LVEF \leq 35%, NYHA class II or III	ICD	1	A	26,29
Prior myocardial infarction	LVEF \leq 35%, NYHA class II or III	ICD	I.	А	26,28,29
Hypertrophic CM	LV wall thickness > 30 mm	ICD	lla	В	29,30
Endocarditis	Valve dysfunction, abscess	Surgery	I	A	27,31

Table 2	Conditions in	which echoo	cardiography c	can lead to	auideline-based	changes in	management
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CM, Cardiomyopathy; *ICD*, implantable cardioverter-defibrillator; *LVEDD*, left ventricular end-diastolic dimension; *LVEF*, left ventricular ejection fraction; *LVESD*, left ventricular end-systolic dimension; *NYHA*, New York Heart Association.

for a specified condition, over a defined period of time. Global payment schemes in which providers assume financial risk can be perceived as carrying with them financial incentives to underuse services, including imaging.³² It is thus incumbent on the cardiology profession to define what constitutes high-value cardiac imaging in each care bundle or episode of care. Even under a fee-for-service system in which there are no incentives to underuse services, underuse of necessary care is common.³³ Nearly 40% of Medicare beneficiaries with newly diagnosed heart failure do not undergo assessments of left ventricular function, a recommended performance measure for these patients.³⁴ Recent studies applying the appropriate use criteria to cardiac testing reveal evidence of missed opportunities to detect and correctly treat heart disease. A retrospective study of appropriate use criteria for coronary revascularization of patients with stable coronary artery disease revealed that only 69% of patients with appropriate indications for revascularization actually underwent revascularization.³⁵ The 2011 appropriate use criteria for echocardiography were applied to 931 consecutive inpatients referred for transthoracic echocardiography in 5 community hospitals in Italy, who were compared with 259 patients who had been discharged without having been referred for echocardiography. The investigators determined that 14.7% of requested studies fell into the inappropriate category. However, among the patients discharged without echocardiographic examinations, 16.2% failed to undergo studies despite appropriate indications, most commonly worsening signs or symptoms of heart failure.¹⁰

The universe of clinical conditions for which echocardiography may be indicated is large, and the task of deciding at what point a study becomes a high-value test is challenging even for physicians trained in cardiology. Determining in the clinic or at the bedside when echocardiography meets "high-value" criteria may be even more difficult for noncardiologists, such as internal medicine specialists or general practice physicians, who order 71% of echocardiographic studies.³⁶ What tools are available to assist clinicians in ordering appropriate imaging tests for a given patient, while refraining from ordering ones of low value? Clinical decision support systems, defined as "any electronic system designed to aid directly in clinical decision making, in which characteristics of individual patients are used to generate patient-specific assessments or recommendations that are then presented to clinicians for consideration," are promising means by which to improve cardiac test ordering.³⁷ Four recent reviews found moderate-strength evidence that decision support systems, integrated into computerized point-of-entry or electronic health

record systems, can improve the appropriate ordering of clinical studies. Decision support systems varied in the effectiveness with which they improved the quality of care, as judged by health care process measures such as performing preventive services, diagnostic test ordering, and prescription of therapies. Data are sparse for effects on patient or economic outcomes. Few of the individual studies reviewed examined imaging.³⁸⁻⁴¹

From the foregoing, a two-sided paradigm for achieving high-value cardiac imaging emerges, as cardiac imaging must be "right sized" in two directions. The issue of inappropriate overutilization has been recognized, and tools to identify and address it are appearing. Those initiatives work toward the important goal of reducing the number of imaging studies for which the ratio of positive outcomes to dollars spent is unfavorable. In a complementary manner, research is needed into scenarios in which inappropriate underutilization takes place. Circumstances in which a favorable ratio of health care value achieved relative to cost indicates that a study is high value must not be missed. A systematic approach for research into the latter would identify controlled trials of cardiac conditions for which therapies that improve patient outcomes require imaging studies to detect candidates for treatment. Then, investigations would delineate methods that start with the patient at the point of care for his or her symptoms or condition and alert care providers (some of whom may not be familiar with the nuances of appropriate cardiac testing) to order precisely the right test. An ideal system would provide realtime feedback to educate as well as inform physicians. Research would later be needed to discover whether beneficial changes take place in practice patterns and, more important, in patient outcomes and societal efficacy.

Progress in this direction is evident from a recent prospective study that evaluated an appropriate use criteria decision support tool for physicians ordering imaging studies for coronary artery disease. The studies included stress echocardiography, as well as myocardial perfusion scintigraphy and coronary computed tomographic angiography. In addition to examining the effects on test appropriateness, the investigators studied the effects of using the tool on intended changes in medical therapy. The tool was used at the point of ordering and took an average of 2 min of physician time to use, and the immediate feedback to the physician provided an educational component. Comparing the first 2 months and the last 2 months of the trial, ordering of rarely appropriate studies decreased from 22% to 6%, the percentage of appropriate studies increased from 49% to 61%, and intended changes in medical therapy increased from 11% to 32%.⁴²

Multimodality, disease-specific appropriate use criteria for imaging are now beginning to appear, with the publication of the 2013 document for cardiovascular imaging in heart failure.⁴³ This appropriate use document differs from prior publications not only because it is the first to encompass multiple imaging modalities. The authors identified five key clinical entry points or scenarios for heart failure-directed imaging, "emphasizing that each indication represents the specific 'point-of-order' for an imaging study." Each scenario reviews the rationale for imaging, delineates the choice of imaging modalities, references the heart failure guidelines, and categorizes the appropriateness of each modality. For the scenario "newly suspected or potential heart failure," appropriate use recommendations drill down to the selection of imaging on the basis of symptoms and signs, for malignancy with cardiotoxic therapy, familial or genetic cardiomyopathy, known adult congenital heart disease, and acute myocardial infarction.

ENVISIONING A SYSTEM FOR PROMOTING HIGH-VALUE CARDIAC TESTING

One can hypothesize the form an ideal system to ensure high-value cardiac imaging might take. The process would use a decision support tool at the point of care or ordering. It would begin with a set of signs and symptoms or a disease state. Logic built into the decision support algorithm could automatically pull patient-specific descriptors that are already present in the database, such as the physical examination, the severity of symptoms, comorbid conditions such as malignancy, and prior relevant imaging or laboratory values such as creatinine, for its impact on choosing a dye study, among others. By referencing appropriate use criteria for multimodality imaging, the system could prompt clinicians to consider ordering an appropriate study for each specific case as indicated. It might be designed to look back to prior, similar testing and, by matching elements in the report (such as mild, moderate, or severe regurgitation) and patient descriptors (such as symptom severity), inform the clinician whether a study meets guideline recommendations to repeat the study or whether to refrain from doing so if these criteria are not met.

A means to harmonize expert opinion from cardiology with guidelines for the same condition by experts in other disciplines might further improve test utilization. Conceptually, at least, clinicians might be most familiar with recommendations in their own specialty's literature. An example is syncope, a condition treated by emergency medicine specialists as well as cardiologists. Syncope is ranked in the echocardiography appropriate use guidelines as appropriate for "clinical symptoms or signs consistent with a cardiac diagnosis known to cause lightheadedness/presyncope/syncope."44 Recommendations similar to these have been made in multidisciplinary guidelines for the investigation of syncope authored by experts in emergency medicine.^{45,46} Using decision rules found in the emergency medicine literature for patients presenting to emergency rooms with syncope may improve the diagnostic and therapeutic efficacy of echocardiography. Applying a syncope diagnostic protocol to patients presenting to a hospital in the United Kingdom increased the percentage of echocardiographic studies performed. Importantly, the authors noted that 75% of echocardiographic studies in the study group were performed for significant clinical findings such as aortic stenosis. Compared with historical controls, the number of examinations needed to make a diagnosis decreased significantly, implying that the percentage of low-yield echocardiographic studies was reduced.47

CONCLUSIONS

Efforts to reduce low-value imaging represent an important start in ensuring the appropriate use of resources. Short-term benefits are relatively easy to quantify, at least economically in terms of dollars saved. We must at the same time recognize and address the thornier problem of missed opportunities for high-value imaging. As the United States moves to value-based purchasing, diagnoses will be bundled into payment groups, and pressure will mount to reduce costs. Physicians must improve the way they order cardiac imaging, moving beyond cost to the concept of value. Inappropriate under-utilization as well as overutilization must be reduced. In the latter case, failure to perform the right test at the right time can lead to an even more unfortunate circumstance for patients, described by hockey great Wayne Gretzky as "you miss 100% of the shots you never take."

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