



Best Practices Guidelines On Imaging Clinical Decision Support Systems

Preface

In light of the evolving technological advances in the area of radiology utilization management, and specifically the emerging importance of clinical decision support¹ (CDS) systems, the Radiology Business Management Association (RBMA) and the American College of Radiology (ACR) determined that the previously published Best Practice Guidelines on Radiology Benefit Management Programs² should be revisited. Although that guideline pointed to the use of evidence-based tools such as CDS, it did not delve into or provide quidance on CDS specifically. Thus, this document on best practices for CDS systems, developed by the ACR and the RBMA, is intended to provide guidance to regulators, payers, vendors, referring physicians, and radiology providers as to implementation, use, and ongoing evaluation of CDS systems. This document delineates optimal business practices for companies that supply CDS systems to ensure that their radiology utilization management products are a continuously improving, cost-effective, efficient, and reliable method of providing clinical and payer-neutral guidance for the appropriate ordering of imaging services. It is essential that these products be based on patients' clinical indications and high-quality, evidence-based clinical imaging guidelines. Such guidelines serve as a benchmark against which CDS systems can be measured. The following best practice recommendations are based on ACR and RBMA member consensus and a review of utilization management practices under development or currently in use by CDS companies. Implementation of these best practices will promote transparency and help create a uniform process that will ease administrative burden on payers, treating clinicians, and radiology providers. This paper is not an endorsement of specific companies that supply CDS systems or their approach to the marketplace.

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¹ For the purposes of these guidelines, the authors consider "clinical decision support," "computerized decision support," "decision support," "computerized imaging order entry," and "computerized physician order entry with clinical decision support (CPOE/CDS)" as analogous terms.

ACR-RBMA Best Practices Guidelines on Radiology Benefits Management Programs (2011); http://www.rbma.org/uploadedFiles/RBMA Web site/Advocacy/Advocacy Payer Relations/ACR-RBMA%20Best%20Practices%20for%20Radiology%20Benefits%20Management%20Programs.pdf

Background

Although imaging unquestionably has significantly improved the quality of health care and increased value by saving money, it is an expensive tool. For numerous reasons inappropriate imaging does occur, which has led to efforts to improve appropriate use through several approaches. One such approach is the use of Radiology Benefit Managers (RBM). RBMs, either independent organizations or payer-owned, are contracted to determine the appropriateness of ordering advanced imaging procedures in the outpatient setting based on the patients' clinical indications (signs, symptoms, or diagnoses) as well as third-party payer internal proprietary guidelines. Determination of appropriateness, ostensibly based on these guidelines, is generally not transparent, may involve added administrative hurdles, and imposes a temporal, administrative, and financial burden on ordering health care providers, radiologists, and insurers. Some payers also implement their own internal RBM program, usually in the form of prior-authorization³. Prior authorization of outpatient services before performance of the imaging study is often required for payment, and may involve selection of the imaging provider by the RBM. This selection process is sometimes referred to as "patient steerage." RBM steerage strategies often ignore patient care integrated networks, provider physician and staff skills, experience, quality, and procedure protocol uniformity.

CDS systems provide a favorable alternative to RBMs and other pre-authorization approaches, as is apparent from a consideration of the characteristics of an ideal system to manage imaging utilization. The functional CDS system incorporates the following:

- Can readily be incorporated into daily work processes.
- Processes requests in real-time with minimal disruption or interference (i.e., moves
 from requests for imaging procedures, to approval and performance if indicated, or to
 communication between the requesting health care provider and the imaging expertise
 inherent in the CDS system if there are questions or concerns).
- It is efficient, user friendly, consistent, and educational, with immediate feedback as to the recommendations.

³ For the purposes of these guidelines, the authors consider "prior-authorization," "pre-certification," and "pre-authorization" as analogous terms.

- It is based on clinical guidelines that are produced using sound methodology, are evidence-based to the greatest extent possible, supplemented as needed by clinical expert opinion, transparent and readily reviewable, and are regularly updated.
- Using sound guidelines and realistic systems, the CDS will allow real-time evaluation of the appropriateness of a requested imaging study and will eliminate the need for any other system of pre-evaluation or pre-certification.
- The CDS system will, through vendors, allow feedback between the creators of the guidelines and the users, and thereby facilitate both improved guidelines and local quality improvement for systems and for individual imaging-ordering providers.
- o Consequently, the CDS system will produce quality measure and outcome data.
- To assess continuous performance improvement longitudinally against benchmark data, the CDS system will supply utilization data to providers that support networks.

These conditions can be met using decision support on the basis of widely accepted evidence-based utilization guidelines, such as the ACR Appropriateness Criteria[®], at point of computer order entry. Proponents note that:

CDS is a cost-effective, efficient, and reliable method for analyzing the clinical indications of a patient and comparing those indications to evidence based data sets, allowing physicians to recommend the most appropriate course of treatment for the patient. This can include a recommendation for no imaging study or to change the requested study to one that is more medically appropriate. The electronic CDS process serves as documentation that the patient is to receive the most appropriate care under the circumstances presented. The benefits of CDS leverage data, drive decision making, improve quality and safety, and help reduce costs by ensuring the right imaging study is recommended.⁴

Several CDS systems are currently commercially available, and use of such systems is being widely considered by health care systems, payers, and regulators.

CDS and RBM companies exhibit similarities and differences in their approach to utilization management. Both claim to utilize national specialty society-approved guidelines, and some supplement these guidelines with their own consensus processes. Unlike the RBM business model, CDS offers real-time, electronically embedded guidance during the imaging ordering

⁴ Computerized Decision Support (CDS) Systems for Advanced Imaging Services, Imaging e-Ordering Coalition.

process, optimally eliminating the need for the referring or treating clinician to take time away from patient care to obtain a pre-authorization and relay that to the imaging provider. RBMs, on the other hand, argue that CDS does not place the same level of accountability on referring physicians for their ordering habits. Several health systems and one state, however, have or are in the process of deploying public data that report and compare physician ordering behavior. CDS vendors and some providers suggest this transparency offers accountability and more effectively influences practice patterns.

The RBM process provides no data about patient care quality or outcomes. RBM efforts are highly focused on controlled spending for imaging. This is appropriate but inadequate with regard to many other important concerns identified by government, health care reform advocates, and providers. No data are available to individual physicians and their groups that would support education allowing identification of the rare "problem provider," or benchmark group performance. No data are available to support integrated care network process improvement or to support accountable care organizations (ACO). There is no data stratification related to patient severity of illness. There is no effort or contribution toward promoting best medical practice standards, that often requires a broader understanding of patient care and evaluation that goes beyond the ordering of an individual imaging test. A properly designed CDS can support all of these vital components.

Clinical Patient Care Guidelines:

The CDS should present the available choice of imaging procedures or range of imaging procedures determined as appropriate based on the medical evidence, to the extent possible, supplemented by clinical expertise. Determining the range of possibly appropriate imaging procedures is based on indications, such as patient symptoms, information from prior exams, the patient/family medical history and risk factors, or clinical circumstances (e.g., emergent case). The degree to which this information is available at the point of ordering an imaging exam varies from patient to patient and procedure to procedure. Additionally, even with substantial information available, the CDS system will not account for every possible clinical scenario. There may be additional factors that impact the final decision for a specific case and there may be more than one procedure that is appropriate for the clinical indications or scenario.

For example, based on indications, history, risk factors, etc., entered into a CDS for a patient with the following:

- Suspected acute pancreatitis
- Severe abdominal pain,
- Elevated amylase lipase,
- Without fever or elevated white blood cell count, but
- With hemoconcentration, oliquria, and tachycardia

The CDS algorithm may find all the following exams as appropriate:

- CT abdomen with contrast,
- · CT abdomen without contrast,
- MRI abdomen without, (including magnetic resonance cholangiopancreatography -MRCP) and with contrast,
- MRI abdomen without contrast with MRCP,
- · CT abdomen without and with contrast or
- US abdomen

At any particular institution or facility, practice variations, available staff, available scanners, patient allergies or other factors may determine the most expedient, efficacious exam choice suitable to answer the clinical question.

The CDS should provide to the ordering clinician any procedure determined appropriate and not restrict selection to a specific exam whether defined by CPT® codes, charge master codes, or other manner that a facility may use to code a procedure. The imaging provider may then determine that no imaging is required or that it is necessary to perform any exam, whether specified within the range of appropriate exams as determined by the CDS or not. In this manner, the CDS should provide the radiologist and treating clinician the flexibility to choose the most appropriate imaging study based on patient indication(s). Furthermore, information from prior exams, the patient medical history and risk factors, family medical history, or clinical circumstances (e.g., emergent case) may necessitate a departure from results presented by a CDS that has been implemented with restrictive adherence to singular or non-inclusive groups of CPT® coding or charge master mapping, which could potentially require a second CDS query.

Allowing a radiologist or other credentialed imaging interpreter to perform such appropriate imaging studies without requiring pre-approval, based on their exigent clinical condition, would avoid repeat studies, delays while attempting to revise an order to permit a more

appropriate study, and additional administrative costs, as well as patient inconvenience and potentially unnecessary or repeated exposure to radiation and contrast.

Education of treating clinicians

CDS systems should provide real-time, 24/7 education and guidance for two purposes: first, to educate referring physicians and other ordering clinicians about the clinical information that needs to be submitted in the imaging prescription, and second, to provide guidance (and education) as to the most appropriate exam, if any, for the specific clinical indications. Both must be accomplished in a way that is evidence-based, non-intrusive, sufficiently user friendly so that the system is accepted by ordering health care providers, payer-neutral and that does not interfere with the physician-patient relationship. Further, referring physicians and their practice personnel should expect uniformity and consistency regarding the type and amount of clinical information needed, regardless of the payer, plan, intermediary, or geography in which the study is performed or ordered. Similarly, in a risk-sharing arrangement such as an ACO, credit should be given to the performing provider when the initial order meets the appropriateness standard.

Use of national specialty guidelines

When a CDS algorithm varies from a specialty-specific guideline (e.g., ACR Practice Guidelines and Technical Standards, ACR Appropriateness Criteria[®]) for any reason, such as to add more specific guidance, this variation must be clearly embedded in the electronic tool and offer transparency to the ordering physician and patient.

Outpatient Study Process

By fully leveraging information available in the patient electronic health record (EHR), CDS systems can provide ordering physicians with additional information to ensure appropriate imaging is ordered. While the CDS provider cannot be expected to have the content to provide clinical information to ordering physicians, CDS and EHR providers are encouraged to develop appropriate interfaces so that both ordering and performing/interpreting physicians have this information readily available through the order entry process. For outpatient procedure authorization and scheduling, provider accreditation and specific payment plan participation for each provider should be incorporated into the order entry database. Many current CDS systems do not include this information. Cumulative patient examination history should be immediately available to the ordering physician to further facilitate the elimination of unnecessary or repeated examinations as well as reduction of

patient exposure to radiation. Common examination requirements such as estimated glomerular filtration rate (eGFR), history of a prior contrast reaction, and other relevant risk factors should also be presented as appropriate. Providers should be able to present specific equipment identification numbers, along with physician, staff, and site accreditation information in digital format when requested.

Compliance with applicable laws and regulations

The utilization management process should be transparent and evidence based. The methodology and data used in constructing the CDS algorithms need to be transparent, sound, and available to the medical community and the public at large. Criteria should be evidence based to the greatest extent possible (through the use of the peer-reviewed literature, specialty guidelines, and sound, widely accepted guideline methodology) and supplemented by expert opinion so that they follow best medical practice. The CDS algorithms and resulting criteria should conform with applicable state and federal laws and regulations and be applied and administered consistently across geographic boundaries and authorizing entities by all third-party payers. The imaging provider should be reimbursed for any medically appropriate studies they perform in accordance with the CDS algorithm, as long as the patient is eligible for coverage under the applicable provider agreement. Most importantly, any study that is found to be medically appropriate in accordance with the CDS algorithm should be deemed to be "pre-authorized" in accordance with applicable federal, state or local laws, rules, or regulations.

Ability to override CDS guidance

CDS should offer a process allowing either the treating clinician or radiologist (with approval of the ordering physician) to override the CDS algorithm guidance within defined guidelines. This is necessary to assure that the physician, who may be aware of a patient's specific medical circumstances that may not have been specifically addressed in the CDS tool, can be the final decision maker on which test is ordered. However, all CDS systems should also have an inherent utilization review and quality improvement (QI) component. That is, their utilization patterns should be shared with the ordering physicians, and analyzed by the health care system locally, to evaluate to what extent ordering of imaging studies conforms to the CDS. This information should first be made available to each physician and the entity where the physician or other clinician practices. Data gathered regionally and nationally, through a CDS system such as that of the ACR Appropriateness Criteria[®], can also be shared locally as an educational and QI tool. Using this approach has several advantages:

efficiency, ease of use, adaptability to individual circumstances that may fall outside the norm or the usual, ready evaluation of the patterns of use of individual ordering health care providers, and use of this information for QI. Further, use of such systems will allow comparison of utilization and of outcomes among institutions and regions to improve quality nationwide. Although CDS providers cannot be expected to anticipate or provide all reports system users may require, all CDS providers should be expected to provide reports regarding utilization patterns of each ordering physician. Additionally, CDS systems should be able to provide raw data to end users so that customized reports can be developed by the institutions according to their specific needs.

Process Consistency

Physicians (treating or interpreting), their patients, and their practice personnel should expect consistent interpretation and application of medically appropriate criteria across geographic boundaries and authorizing entities in accordance with applicable law. Predictability in the determination of medically appropriate studies will promote compliance, help mitigate burdensome administrative costs and promote the delivery of a uniformly high and improving quality of patient care. Conversely, inconsistent standards lead to uneven patient care, frustration on the part of referring and rendering physicians and their staff and patients, and unnecessary administrative cost.

Administrative Process Guidelines:

Continuous ordering improvement process

As noted, CDS systems should allow a referring or rendering physician to override the guidance because such physician(s) may be aware of the need to update or further refine the CDS algorithms/guidelines. Each CDS company should clearly communicate what steps a physician should take to communicate a potential change which he or she recommends. The response to such suggestions by the CDS company should be timely and transparent. These instances of exam order changes or CDS guidance "overrides" need to be part of the CDS system data reports so that appropriate algorithms and process improvements can be made over time.

Procedure Payment Process Consistency

Imaging services that have been conducted after a CDS has been consulted should not have reimbursement denied by a third-party payer based on medical appropriateness grounds, as long as the patient is eligible for coverage under the applicable provider agreement. Providers need confidence that CDS-approved services that have been documented in the EHR or otherwise documented electronically will be reimbursed. Those third-party payers who require a CDS consultation number to be included with the claim must provide a rectification and appeals process for any claim that is denied based on consultation number transmission errors, whether they are human or technical.

Reimbursement criteria should be applied and administered consistently across geographic boundaries and authorizing entities, not only with respect to medically appropriate criteria, but also without regard to a payer's coverage and adjudication policies, i.e., ideally an imaging provider should receive reimbursement for a study that successfully utilized CDS regardless of payer, plan, or other administrative considerations.

Payers should ensure that all services vetted by a CDS can be transferred properly and on time for accurate claims processing

If the CDS vendor and/or their payer clients intend to use the CDS process to deny service or payment, then the payer and/or CDS provider should have sufficient claims adjudication, electronic connectivity systems (i.e., currently a portal), and reconciliation processes in place. Studies found by the CDS algorithm to be medically appropriate should be accurately expedited through the claims processing systems and reimbursed.

<u>Payers should, across all product lines and regardless of any internal utilization</u> <u>management programs, reimburse providers for all imaging studies that have been vetted</u> <u>by the CDS system</u>

Payers should avoid "mixing" utilization management programs for their product lines by avoiding the use of multiple systems, i.e., both CDS and RBM. Such mixing creates added administration costs as well as confusion, since many patients do not know the details of their plans and the variability can be overlooked readily by referring physicians, their staff, or the staff of the imaging facility.

The CDS exam validation period should be from the date of the order by the referring physician through a period of 30–60 days

The validation period should be considered to extend "from the date the tool was consulted" to avoid re-application or payment denial. Furthermore, to promote patient and provider convenience and efficiency, the patients should be afforded the flexibility of scheduling

anytime within the applicable validation period. This is important for some types of advanced imaging services which may not be available in all geographic areas, requiring the patient to arrange for travel for the services.

Transparency Recommendations:

The collection of quality of imaging information, ordering behavior, and cost data from practices should be fair, consistent, and accurate

Payers increasingly are exploring the use of comparative statistics for practices based on quality, ordering behavior and cost. Although this could be a useful resource for patients and employers, providing this information is beyond the scope of CDS. However, if payers and CDS providers use information from physician ordering patterns, the value of such a product will depend largely on the accuracy of the underlying data and the reliability of comparisons. Further, it will require the use of consistent, transparent, high-quality imaging guidelines that are comparable, or identical, for any CDS systems. Accordingly, payers should have processes in place to ensure that the data are collected and shared consistently across practices, geographic boundaries, and authorizing entities using a common, transparent methodology and equivalent data sets and predictive modeling techniques. Further, the data should be subject to review by the practices prior to release and there should be the expectation that there will be frequent and easy updating of such data as the CDS system and the practice changes. A credible data dispute resolution process is essential to ensure fairness to payers, providers, and the public. Resolution of a dispute which changes publicly reported data should be posted within 30 days of the final decision.

CDS should provide regular performance reports over a specific period of time

The CDS system should provide reports to the payer, providers, and each referring physician on the ordering patterns of each referring physician, including his or her adherence to appropriateness guidelines that are the basis of the CDS system. Rules governing which data will be public and which data will be privileged should be clearly stated. Quality assurance data may be subject to legal protection.

Benchmarking

There is presently no credible accreditation process for CDS systems. Further, there are no nationally accepted CDS performance standards or metrics developed at this time. Our expectation is that such criteria will emerge in the future. In the interim, each CDS system

should establish and report performance metrics for system operations, physician, and provider satisfaction. These data should be reported for specific time intervals and as longitudinal trends.

CDS Criteria Revision

The CDS system and its algorithms should be regularly updated for new and revised clinical information (e.g., guidelines), changes in CPT[®] coding, etc. Updates should be clearly communicated, readily apparent, and smoothly incorporated when using the CDS tool.

Technology Issues:

CDS should be part of an integrated health care system solution

Ideally, CDS should integrate seamlessly and on a real-time basis with the computerized clinical and management solutions utilized by the rendering physician, imaging facility (hospital or imaging center), and treating clinician without the need to separately purchase vendor specific HL-7 or other interfaces.

CDS should be capable of mining a patient's medical record for relevant information and include such information in its determination-making process.

A patient's medical record may contain information (e.g., prior studies, personal or family histories) that would be pertinent to the imaging study being evaluated along with the information given by the referring physician. EHR vendors employing CDS systems that are integrated into an overall health care system solution (e.g., EHR, radiology information system, hospital information system) are encouraged to develop interfaces to extract relevant patient information for use at the time of order entry.

CDS can be successfully deployed in a variety of ways

A CDS can be a web-based process that generates an authorization number similar to an RBM program. This approach does not require disparate computer system integration or HL-7 program interfaces. Although less elegant than fully integrated solutions, web-based systems can be deployed relatively quickly and inexpensively. CDS could be deployed by a health plan as an improvement in their current utilization management solution. CDS could also be deployed by providers who then contract with payers to provide this front-end utilization management. This position paper supports the underlying visions that CDS will be fully integrated into the institutional EHR process and delivers a superior decision process.

The notion that all payers must accept an internal provider's CDS process as acceptable is neither mandated nor required today, but would be highly desirable. All of these approaches can add value if properly deployed and supported. All can operate at a lower cost versus RBM methods while achieving similar macro-economic results.

Conclusion

The ACR and the RBMA both are strong proponents of performing the right study, at the right time, in the right way for each individual patient. The ACR has more than a 20year history of developing and updating ACR Appropriateness Criteria®, ACR Guidelines and Standards, the most rigorous accreditation programs in imaging, and numerous quality improvement data registries. The ACR has also collaboratively developed and participated in the Image Gently, Image Wisely, and Choosing Wisely campaigns. ACR activities have contributed significantly to guiding and improving patient imaging care in the United States; similar programs are now being developed in many other parts of the world. The ACR and RBMA firmly believe in and advocate for the adoption of CDS systems as a more effective and less costly process for ensuring appropriate imaging now and in the future. The RBM approach is now recognized to be more costly, less efficient, and inadequate for our future health care policy goals and needs as a nation. This guidance paper describes the most desirable characteristics of CDS systems and provides a framework against which CDS performance, design, and utilization can be measured.