

Harrison's Principles of Internal Medicine, 21e >

Chapter 9: Diagnosis: Reducing Errors and Improving Quality

Gordon Schiff

INTRODUCTION

Diagnosing patients' illnesses is the essence of medicine. Patients present to doctors seeking an answer to the question, "What is wrong with me?" Ideally, no clinician would want to treat a patient without knowing the diagnosis or, worse yet, erroneously treat a misdiagnosed illness. From the earliest moments of medical school, the defining quest toward becoming a knowledgeable and proficient physician is learning how to put a diagnostic label on patients' symptoms and physical findings, and clinicians pride themselves on being "good diagnosticians." Yet the centuries-old paradigm of mastering a long list of diseases, understanding their pathophysiology, and knowing the cardinal ways they manifest themselves in signs and symptoms, while still of fundamental importance, is being challenged by new insights illuminated by the glaring spotlight of diagnostic errors. Basic internal medicine diseases, such as asthma, pulmonary embolism, congestive heart failure, seizures, strokes, ruptured aneurysms, depression, and cancer, are misdiagnosed at shockingly high rates, often with 20–50% of patients either being mislabeled as having these conditions (false-positive diagnoses) or having their diagnosis missed or delayed (false negatives). How and why do physicians so often get it wrong, and what can we do to both diagnose and treat the problem of delayed diagnosis or misdiagnosis?

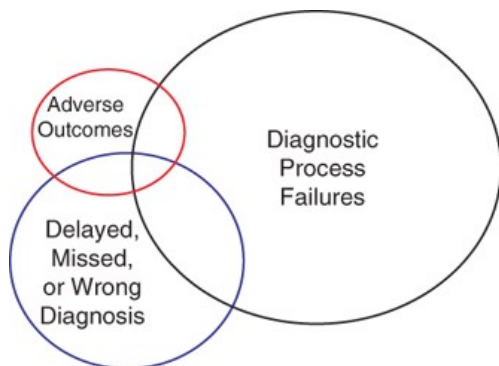
Diagnosis is both an ancient art and a modern science. The current science of diagnosis, however, goes far beyond what typically comes to clinicians' and patients' minds when they conjure up images of state-of-the-art molecular, genetic, or imaging technologies. Improvements in diagnosis are just as likely to come from other areas, many with origins outside of medicine, as they are from advanced diagnostic testing modalities. These diverse sciences that the field of diagnostic safety has, and must, draw from include systems and human factors engineering, reliability science, cognitive psychology, decision sciences, forensic science, clinical epidemiology, health services research, decision analysis, network medicine, learning health systems theory, medical sociology, team dynamics and communication, risk assessment and communication, information and knowledge management, and health information technology, especially artificial intelligence and clinical decision support. A clinician reading this chapter is likely to find this list of overlapping and intersecting domains quite daunting. However, rather than feeling overwhelmed, we urge readers to view them as the basic science supports that will ultimately make their lives easier and diagnosis more accurate and timely. Rather than feeling intimidated, clinicians should feel a sense of relief and assurance in understanding that good diagnosis does not rest entirely on their shoulders. Instead, it is a systems property, where an infrastructure and a team, one that especially includes the patient, can in a coordinated way work together to achieve more reliable and optimal diagnosis.

EMERGENCE OF DIAGNOSIS ERROR AS AN IMPORTANT PATIENT SAFETY ISSUE

Over the past decade, a series of studies culminating in a landmark report from the U.S. National Academy of Medicine (NAM), *Improving Diagnosis in Health Care*, have shone a spotlight on diagnostic errors. Reports from patient surveys, malpractice claims, and safety organizations, such as the ECRI and the National Patient Safety Foundation (now part of Institute for Healthcare Improvement), have found that diagnostic errors are the leading type of medical error. Although errors in diagnosis are defined in various ways, the NAM Committee defined diagnostic error as "the failure to (a) establish an accurate and timely explanation of the patient's health problem(s) or (b) communicate that explanation to the patient." One way to visualize diagnostic errors is through a Venn diagram (**Fig. 9-1**), which illustrates the fact that many things can go wrong in the diagnostic process (e.g., failure to ask an important history question, physical examination sign overlooked, laboratory specimen erroneously switched between two patients, x-ray not followed up), but this usually does not result in a wrong diagnosis or patient harm. Similarly, a patient can be misdiagnosed but unharmed, without any identifiable error in the care received. Our greatest concern is where these three circles intersect, with conservative estimates suggesting that 40,000–80,000 patients die each year in U.S. hospitals alone from diagnostic errors. The NAM report outlined eight recommendations that are the foundation for this chapter (**Table 9-1**).

FIGURE 9-1

What is a diagnosis error? (Adapted from GD Schiff et al: *Diagnosing diagnosis errors: Lessons from a multi-institutional collaborative project*, in *Advances in Patient Safety: from Research to Implementation*. Vol. 2 Concepts and Methodology, Rockville, MD, 2005, pp 255-278, and GD Schiff, L Leape: *Acad Med* 87:135, 2012.)



Source: Joseph Loscalzo, Anthony Fauci, Dennis Kasper, Stephen Hauser, Dan Longo, J. Larry Jameson: *Harrison's Principles of Internal Medicine*, 21e Copyright © McGraw Hill. All rights reserved.

TABLE 9-1

National Academy of Medicine Recommendations for Improving Diagnosis in Health Care

1. Facilitate more effective teamwork in the diagnostic process among health care professionals, patients, and their families.
2. Enhance professional education and training in the diagnostic process in areas such as clinical reasoning; teamwork; communication with patients, families, and other health care professionals; and appropriate use of diagnostic tests.
3. Ensure that health information technologies support patients and health care professionals in the diagnostic process.
4. Develop and deploy approaches to identify, learn from, and reduce diagnostic errors and near misses in clinical practice including providing systematic feedback on diagnostic performance.
5. Establish a work system and culture that supports the diagnostic process and improvements in diagnostic performance.
6. Develop a reporting environment and medical liability system that facilitates improved diagnosis by learning from diagnostic errors and near misses.
7. Design a payment and care delivery environment that supports the diagnostic process.
8. Provide dedicated funding for research on the diagnostic process and diagnostic errors.

NEW WAYS TO THINK ABOUT DIAGNOSIS AND DIAGNOSTIC ERRORS

Medical textbooks have historically given attention to “clinician reasoning” and associated cognitive heuristics and biases. Errors in clinical reasoning can be summarized in three broad groups: (1) hasty judgments, (2) biased judgments, and (3) inaccurate probability estimates. Research from cognitive psychology has identified scores of common mental shortcuts or “heuristics” humans are prone to use in everyday life, many of which are useful for efficient diagnosis but can also lead to biases and errors. **Table 9-2** lists some of the common cognitive biases that can lead diagnosis astray (this topic is discussed further in **Chap. 4**).

TABLE 9-2

Selected Cognitive Biases Contributing to Diagnostic Errors

- 1. Premature closure: accepting a diagnosis before it has been fully verified
- 2. Anchoring: tendency to fixate on a specific symptom or piece of information early in the diagnostic process with subsequent failure to appropriately adjust
- 3. Confirmation bias: tendency to look for confirming evidence to support one’s diagnostic hypothesis, rather than disconfirming evidence to refute it
- 4. Search satisficing: tendency to call off a search, satisfied once a piece of data or presumed explanation is found, and not considering/searching for additional findings or diagnoses
- 5. Availability bias: tendency to give too much weight to diagnoses that come more readily to mind (e.g., recent dramatic case)
- 6. Base-rate neglect: failing to adequately take into account prevalence of a particular disease (e.g., erroneously interpreting a positive test as indicating disease in a low-prevalence population using a test with 5% false-positive rate)
- 7. Knowledge deficit (on part of provider, with accompanying lack of awareness)
- 8. Framing bias: Judgement overly influenced by the way the problem was presented (how it was framed in words, settings, situations)
- 9. Social/demographic/stereotype bias: biases from personal or cultural beliefs about women, minorities, or other patient groups for whom prejudices may distort diagnostic assessment

However, clinicians will also benefit from having a better understanding of diagnosis as a “system” rather than just what takes place in clinicians’ minds. Classic teaching exhorting trainees and practicing physicians to have a broad differential and “high index of suspicion” for various diseases is challenged not only by these unconscious biases but also by limitations of human memory, information shortfalls, constrained encounter time, system process failures, and the myriad nonspecific symptoms that patients bring to clinicians. Many symptoms are self-limited, defy a precise diagnosis or etiology, and do not portend harmful outcomes. Insights from safety and cognitive sciences call for rethinking traditional approaches to diagnosis and suggest new approaches to overcome current limitations (Table 9-3).

TABLE 9-3

New Models for Conceptualizing Diagnosis and Diagnosis Improvement

TRADITIONAL WAYS OF THINKING ABOUT DIAGNOSIS AND DIAGNOSTIC ERROR	NEW PARADIGMS/BETTER WAYS TO THINK ABOUT DIAGNOSIS AND IMPROVING DIAGNOSIS
General	
A good diagnostician gets it right the first time, almost all of the time	Diagnosis is an inexact science with inherent uncertainties Goal is to minimize errors and delays via more reliable systems and follow-up
Lore of masterful/skillful academic expert diagnostician who knows/recalls everything; need to look to them if seeking diagnostic excellence	Less reliance on (fallible) human memory Quality diagnosis is based on well-coordinated distributed network/team of people and reliable processes All patients entitled to receive quality diagnosis, regardless of where and from whom they receive care
Diagnosis is the doctor’s job	Co-production of diagnosis among clinicians (including lab, radiology, specialists, nurses, social workers) and, especially, the patient and family
Patients often viewed as overly anxious, exaggerating, time-consuming, questioning, with sometimes unreasonable demands and expectations	Patients are key allies in diagnosis; hold key information Need to address understandable/legitimate fears, desires for explanations Leveraging patient questions and questioning of diagnosis to stimulate rethinking the

	diagnosis where needed
Diagnosis and treatment as separate stages in patient care (i.e., make a diagnosis, then treat)	Prioritizing diagnostic efforts to target treatable conditions More integrated strategies and timing for testing and treatment depending on urgency for treatment
Clinical practices	
Order lots of tests to avoid missing diagnoses	Judicious ordering: targeted, well-organized data and testing Appreciation of test limitations (false positive or negative, incidental findings, overdiagnosis, test risks) and resulting harms
More referrals to avoid missing rarer/specialized diagnoses; concomitant utilization barriers (copays, prior authorization) to minimize overuse	“Pull systems” to lower barriers and make it easier to pose questions, obtain real-time virtual consults Co-management approaches to enable collaborative watch-and-wait conservative strategies where appropriate
Frequent empirical drug trials when uncertain of diagnosis	Conservative use of drugs to avoid confusing clinical picture or labeling patients with diseases they may not have
Physician attention/efforts to ensure disease screening	Automating, delegating clerical functions; teamwork to free up physician cognitive time
Diagnosis errors and challenges	
Diagnostic error viewed as a personal failing Errors classified as either “system” or “cognitive”	Many errors/delays rooted in processes and system design/failures Errors multifactorial with interwoven, interacting, and inseparable cognitive and system factors
Errors are infrequent; hit-and-miss ways to learn about errors	Errors are common; systematic proactive follow-up is needed to recognize potential for errors Surveilling of high-risk situations and one’s own diagnostic performance and outcomes
Clinicians’ reactions: denial, defensive, others to blame, pointing to others also making similar errors	Culture of actively and nondefensively seeking to uncover, dig deep to learn from, and share errors and lessons
Dreading complex, frustrating diagnostic dilemmas	Welcoming/enjoying intellectual/professional challenges Adequate support (time, help, consultations) for more complex patients
Diagnoses as distinct labels, events	Diagnoses can be indistinct, interacting comorbidities, socially constructed, multifactorial, evolving over time, or have overlapping genotype-phenotype expressions
Documentation/communication	
Viewed as time-consuming, mindless, primarily to document for billing code and/or bulwark against malpractice claims	Documentation as useful tool for reflecting, crafting, sharing assessments, differential diagnosis, reflecting about unanswered questions Opportunities for decision support interacting with computer Notes open for patients to read to help understand and critique diagnosis
Say and write as little as possible about uncertainties, lest it be used against you in malpractice allegation	Share uncertainties to maximize communication and engagement with other caregivers, patients

Don't let patient know about errors so they don't become angry, mistrustful, or sue	Patients have right to honest disclosure; often find out about errors anyway (e.g., cancer evolves); anticipate, engage their concerns
Patients advised to call if not better; no news is good news (test results: "We'll call if anything is abnormal.")	Systematic proactive follow-up to close loop on tests and symptoms, to check how patient is doing, monitor outcomes
Global remedies	
Knowing/memorizing more medical knowledge	Knowing more about the patient (including psychosocial, past history, environmental contexts)
Attention to the "objective" data (physical exam, tests) to reliably make diagnoses	Renewed emphasis on history, history-taking, listening Acknowledgement of ubiquitous subjective cognitive biases; efforts to anticipate, recognize, counteract
Exhortations to have "high index of suspicion" of various diagnoses	Less reliance on memory recall of lectures/reading; more just-in-time info look-up Affordances, alerts to red flags engineered into workflow Delineation of "don't miss" diagnoses with design of context-relevant decision support reminders
Ensuring physician is copied on everything, thorough/voluminous notes, widespread reminders/alerts	Biggest problem is no longer lack of access to information, but rather information overload; strategies to organize, minimize
Continuing medical education (CME) courses to expand medical knowledge	Real-time, context-aware reminders of pitfalls, critical differential diagnoses, and key differentiating features. Ready access to medical references, second opinions
Redundancies, double-checks	Recognition that single, highly reliable systems are often better than multiple halfway solutions. Clear delineation of responsibilities for follow-up tasks
Fear of malpractice suits to motivate physicians to be more careful and practice defensive medicine	Drive out fear, making it safe to learn from and share errors Shared situational awareness of where pitfalls lurk
More accountability, financial incentives, and penalties tied to performance metrics	Clinician engagement in improvement based on trust, collaboration, professionalism, financial neutrality Metric modesty, recognizing many best practices yet to be defined/proven
More rules, requirements; target outliers for better compliance	Standardization with flexibility; learning from deviations
More time with patients	Better time spent with patients: offloading distractions, more efficient history collection/organization, longitudinal continuity, and, where needed, additional time to talk/think/explain during, before, or after visits Easier access for patients to reach or be seen by clinicians when experiencing symptoms
Reflex changes in response to errors	Avoiding "tampering," which entails understanding/diagnosing difference between "special cause" versus "common cause" (random) variation

Source: Modified from GD Schiff: Quality and Safety in Health Care 2013.

UNCERTAINTY IN DIAGNOSIS

Given variations and overlap in ways patients present, illnesses evolve, and tests perform, it is often not possible or practical to “make” a definitive diagnosis, particularly in the primary care setting early in the course of a patient’s illness. Clinicians need to harness these uncertainties to both engineer situational awareness of where things can go wrong and create safety nets to protect patients against harms from delayed diagnosis and misdiagnosis. Terms such as *preliminary diagnosis*, *working diagnosis*, *differential diagnosis*, *deferred diagnosis*, *undiagnosed illness*, *diagnoses with uncertain or multifactorial etiologies*, *intermittent diagnoses*, *multiple/dual diagnoses*, *self-diagnosis*, or at times *contested diagnosis* need to be part of our vocabulary, thinking, and communications with patients to convey that diagnosis is often imprecise. Anxious patients worried about a condition, for example, cancer, COVID-19 infection, or a diagnosis to which a relative or a friend has recently succumbed, come seeking reassurance and may not welcome an uncertain answer. Thus, we have to work with patients, listen to and respect their concerns, and take their symptoms seriously yet modestly acknowledge our limitations. We need to tailor this approach to patients’ differing levels of health literacy, trust in our clinical advice, and experiences with the health system.

DON’T MISS DIAGNOSES AND RED FLAGS

Uncertainty should not be a license for complacency. Particularly for diseases that (1) progress rapidly, (2) require specific treatments that depend on making the correct diagnosis, or (3) have public health or contagion implications, clinicians need to be poised, and systems designed, to consider and, where appropriate, pursue critical “don’t miss” diagnoses. While clinicians are generally aware of more common “don’t miss” diagnoses (e.g., acute myocardial infarction, sepsis), **Table 9-4** illustrates examples of less common diagnoses that warrant similar consideration. Throughout this textbook, readers should orient themselves to recognize such critical diagnoses and think about presentations and syndromes where they may be lurking.

TABLE 9-4

Examples of “Don’t Miss” Diagnoses

INFECTIONS/INFLAMMATION	CARDIAC/ISCHEMIC/BLEEDING	METABOLIC/HEMATOLOGIC/ENVIRONMENTAL
Spinal epidural abscess	Aortic dissection Leaking/ruptured abdominal aortic aneurysm	Diabetes ketoacidosis Hyperosmolar hyperglycemia
Necrotizing fasciitis	Pericardial tamponade	Myxedema/thyrotoxicosis
Meningitis	Wolff-Parkinson-White Prolonged QT	Addison’s disease
Endocarditis	Pulmonary embolism	B ₁₂ deficiency anemia
Peritonsillar abscess	Tension pneumothorax	von Willebrand’s disease
Tuberculosis-active pulmonary, other	Acute mesenteric ischemia Sigmoid volvulus	Hemochromatosis
COVID-19 infection	Esophageal, bowel perforation	Celiac sprue
Guillain-Barré syndrome	Cerebellar hemorrhage	Carbon monoxide poisoning
Ebola infection	Spinal cord compression	Food poisoning
Temporal arteritis	Testicular, ovarian torsion	Malignant hyperthermia
Rhabdomyolysis	Ectopic pregnancy	Alcohol, benzodiazepine, barbiturate withdrawal
Angioedema	Retroperitoneal hemorrhage	Tumor lysis syndrome Hypo-/hypercalcemia

An important related concept is so-called “red flags” or “alarm symptoms.” This construct has its origins in guidelines for back pain but has increasingly been applied to many other problems, such as headache, red eye, swollen joint, or even abdominal pain and chest pain. Examples of widely cited red flags for back pain that should trigger consideration of more serious etiologies include fever, weight loss, history of malignancy or intravenous drug use, or neurologic signs and symptoms. In theory, many presenting syndromes could benefit from identification of such clues to more serious diagnoses. Evidence-based medicine calls for better data on the sensitivity, specificity, yield, and discriminatory ability of various clinical “red flag” clues; yet, few have been rigorously evaluated. Nonetheless, clinicians find them useful as simple ways to reassure themselves and their patients that a common symptom such as back pain or headache is, or is not, likely an indicator of more urgent or serious pathology.

Interwoven with the challenges of not missing critical diagnoses is the problem of overtesting and overdiagnosis—performing unnecessary and even potentially harmful tests whose benefit does not justify the risks or costs or that may lead to diagnoses that would have never caused any symptoms or problems. Thoughtful diagnosticians need to weigh carefully this “other side of the coin” of missed diagnosis to avoid such harms and expenses.

DIAGNOSTIC PITFALLS

One of the important ways of learning in medicine is learning from the missteps of those who have walked the path ahead of us. By learning about commonly missed diagnoses and the ways accurate, timely diagnosis went astray, we can avoid making similar mistakes. Anticipating the potential for

similar types of errors can both create situational awareness of traps to avoid and contribute to learning from our own personal and collective patterns of mistakes. Several studies have examined common or recurring pitfalls in diagnosis. An example of a common disease-specific diagnostic pitfall in breast cancer diagnosis is ordering a mammogram for a woman with a palpable breast lump and, when the mammogram returns as normal, reassuring her that cancer has been “ruled out” by the negative test. Any mass or lesion palpable on physical examination probably needs more careful assessment proceeding all the way to invasive biopsy, if necessary. Diagnostic pitfalls can be classified into a number of generic scenarios ([Table 9-5](#)). We now have large databases that have the potential to track “diagnoses outcomes”—i.e., whether a new diagnosis emerges that suggests an initial diagnosis was incorrect or a diagnosis of a patient’s symptoms was suboptimally delayed. This should, in the future, allow us to more rigorously focus on these cases, to identify contributing factors and recurring patterns, and to help point the way for systemwide improvement strategies.

TABLE 9-5

Generic Types of Diagnostic Pitfalls

PITFALL	EXAMPLES
Disease A mistaken for disease B Diseases often mistaken/misdiagnosed with each other	<ul style="list-style-type: none"> Aortic dissection misdiagnosed as acute myocardial infarction Bipolar disorder misdiagnosed as depression
Misinterpretation of test result(s) False-positive or false-negative results with failure to recognize test limitations	<ul style="list-style-type: none"> Breast lump dismissed after negative mammogram Negative COVID-19 test early or late in course
Failure to recognize atypical presentation, signs, and symptoms	<ul style="list-style-type: none"> Apathetic hyperthyroidism Sepsis in elderly patient who is afebrile or hypothermic
Failure to assess appropriately the urgency of diagnosis Urgency of the clinical situation was not appreciated and/or delays critical diagnoses	<ul style="list-style-type: none"> Compartment syndrome Pericardial tamponade Tension pneumothorax
Perils of intermittent symptoms or misleading evolution Intermittent symptoms dismissed due to normal findings (exam, lab, electrocardiogram) when initially seen	<ul style="list-style-type: none"> “Lucid interval” in traumatic epidural hematoma Paroxysmal arrhythmias Intermittent hydrocephalus (Bruns’ syndrome)
Confusion arising from response/masking by empiric treatment	<ul style="list-style-type: none"> Empiric treatment with steroids, proton pump inhibitors, antibiotics, pain medication erroneously masking serious diagnosis
Chronic disease or comorbidity presumed to account for new symptoms Especially in medically complex patients	<ul style="list-style-type: none"> Septic joint signs misattributed to chronic rheumatoid arthritis Mental status change due to infection or medication misattributed to underlying dementia
Rare diagnosis: failure to consider or know	<ul style="list-style-type: none"> Many; fortunately, by definition, rare, but still warrant consideration especially if urgent or treatable
Drug or environmental factor not considered/overlooked Underlying etiology causing/contributing to symptoms, or disease progression not sought, uncovered	<ul style="list-style-type: none"> Ventricular arrhythmia related to QT-prolonging drug Achilles tendon rupture related to quinolone drugs
Failure to appreciate risk factors for particular disease	<ul style="list-style-type: none"> Family history of breast, colorectal cancer not solicited and/or weighed in diagnostic evaluation or screening
Failure to appreciate limitations of physical exam Now with ↑ telemedicine, missing physical exam entirely	<ul style="list-style-type: none"> Overweighing absence of tenderness, swelling in deep vein thrombosis Missing pill-rolling tremor during telemedicine visit

DIAGNOSIS SAFETY CULTURE

Just as diagnosing bacterial infections relies on a proper culture medium to grow and identify etiologic organisms, good diagnosis also requires a healthy safety culture that will allow it to grow and flourish. While clinicians may be inclined to view “safety culture” as something too subjective to be important in their quest to make a definitive diagnosis, this view is misguided. Multiple studies have demonstrated adverse consequences resulting from organizational cultures that inhibit openness, learning, and sharing and create a climate where staff and patients are afraid to speak up when they observe problems or have questions. Most importantly, patients need to be encouraged to question diagnoses and be heard, particularly when they are not responding to treatment as expected or developing symptoms that are either not consistent with the diagnosis or represent possible red flags for other diagnoses or complications.

Studies examining “high-reliability organizations” outside of medicine and “learning health care organizations” have distilled a series of fundamental properties that are correlated with more reliable and safer outcomes. Just as a thermometer or recording of a pulse can suggest how ill a patient is, we now have instruments that can measure safety culture. These safety measurement tools typically are validated staff surveys that assess (1) communication about errors with staff willingness to report mistakes because they do not feel these mistakes are held against them; (2) openness and encouragement to talk about hospital/office problems; (3) existence of a learning culture that seeks to learn from errors and improve based on lessons learned; (4) leadership commitment to safety, prioritizing safety over production speed and the “bottom line” by providing adequate staffing and resources to operate safely; and (5) accountability and transparency for following up safety events and concerns. Each of these generic culture attributes translates into specific implications for diagnostic safety. These include the following:

- Making it “safe” for clinicians to admit and share diagnostic errors
- Proactive identification, ownership, and accountability regarding error-prone diagnostic workflow processes (particularly around test results, referrals, and patient follow-up)
- Leadership making diagnosis improvement a top priority based on recognition that patients and malpractice insurers report that diagnostic errors are the leading patient safety problem
- Mutual trust and respect for challenges that clinicians often face in making diagnoses and caution in applying the lens of hindsight bias in judging what in retrospect might seem like an “obvious” diagnosis that a clinician initially missed

HEALTH INFORMATION TECHNOLOGY AND THE FUTURE OF DIAGNOSIS

Clinicians now spend more time interacting with computers than they do interacting with patients. This is especially true for diagnosis and will likely be even more so in the future. Interactions with patients, consultants, and other staff are increasingly mediated through the computer. Key activities, such as collecting patients’ history (past and current), interpreting data to make a diagnosis, conveying diagnostic assessments (to others on the team and, increasingly, to the patient via open notes), and tracking diagnostic trajectories as they evolve over time, are now computer based. With the rise of telemedicine, even elements of the physical examination have been rerouted to electronic encounters.

While many complain the computer has “gotten in the way” of good diagnosis, distracting clinicians from quality time listening to patients and miring doctors in reading and writing notes filled with copied/pasted/templated information of questionable currency and accuracy, medicine needs to harness the computer’s capabilities to improve diagnosis (**Table 9-6**). Although these basic diagnosis-supporting capabilities should be the foundation of the design of health information technology and everyday workflow, electronic medical records have historically been largely designed around other needs, such as ordering medications and billing and malpractice documentation. They need to be radically redesigned to better support diagnostic processes, as well as save, rather than squander, clinicians’ time.

TABLE 9-6

Areas Where Health Information Technology Has Potential to Help Improve Diagnosis and Reduce Errors

FUNCTION	EXAMPLES
Facilitate collection/gathering of information	<ul style="list-style-type: none">• Quickly access past history from prior care at same and outside institutions• Electronic collection of history of present illness, review of systems, and social determinant risks in advance of visits
Enhanced information entry, organization, and display	<ul style="list-style-type: none">• Visually enhanced flowsheets showing trends, relationships to treatment• Reorganized notes to facilitate summarization and simplification and prevent items from getting lost
Generating differential diagnosis	<ul style="list-style-type: none">• Automated creation of lists of diagnoses to consider based on patient's symptoms, demographics, risks
Weighing diagnoses likelihoods	<ul style="list-style-type: none">• Tools to assist in calculation of posttest (Bayesian) probabilities
Aids for formulating diagnostic plan, intelligent test ordering	<ul style="list-style-type: none">• Entering a diagnostic consideration (e.g., celiac disease, pheochromocytoma) and computer suggests most appropriate diagnostic test(s) and how to order
Access to diagnostic reference information	<ul style="list-style-type: none">• Info-buttons instantly linking symptom or diagnosis relevant questions to Harrison's, Up-to-Date chapters, references
Ensuring more reliable follow-up	<ul style="list-style-type: none">• Hardwiring "closed loops" to ensure abnormal labs, missed referrals, worrisome symptoms are tracked and followed up
Support screening for early detection	<ul style="list-style-type: none">• Collaborative tools that patients, clinicians, and offices can use to know when due, order and track screening based on individualized demographics, risk factors, prior tests
Collaborative diagnosis; access to specialist	<ul style="list-style-type: none">• Real-time posing/answering of questions• Electronic consults; virtual co-management
Facilitating feedback on diagnoses	<ul style="list-style-type: none">• Feeding back new diagnoses (from downstream providers, patients) that emerge suggesting potential misdiagnosis/errors to clinicians, ERs who saw patient previously

Abbreviation: ERs, emergency rooms.

Source: Modified from G Schiff, DW Bates: N Engl J Med 362:1066, 2010, and R El-Karah et al: BMJ Qual Saf Suppl 2:ii40, 2013.

DIAGNOSIS OF DIAGNOSIS ERRORS AND SAFETY: PRACTICAL CONCLUSIONS

In practice, there are frequent and meaningful opportunities for improving diagnosis in each of the three NAM-defined areas to make it a) more reliable, b) timely, and c) to improve diagnosis-related communication with patients. Clinicians in training, practicing physicians, nurses, and others should develop the habit of regularly asking themselves three questions about individual patients in their care, and another three questions regarding

the systems in which they work. For each patient being assessed, clinicians should ask:

1. What else might this be? (forcing a differential diagnosis to be made)
2. What doesn't fit? (making sure unexplained abnormal findings are not dismissed)
3. What critical diagnoses are important not to miss? (injecting consideration of "don't miss" diagnoses, red flags, and known pitfalls)

and to diagnose safely, each practitioner must recognize that he or she is working within a larger system. Questions to be asking continually, ensuring we are maximizing reliability and timeliness and minimizing potential for errors, include:

1. Do we have reliable "closed loop" systems to provide reliable, ideally automated tracking and following up of patients' symptoms, abnormal laboratory or imaging findings, and critical referrals that we order?
2. What is the culture-of-safety climate in our organization, office, or clinic?
3. How does the electronic (or even paper) medical record as currently implemented help versus impair efficient, timely, accurate, and fail-safe diagnosis, and how can it be improved?

To take these questions to the next stage, an international movement dedicated to studying and improving diagnosis has emerged. These efforts include annual conferences of clinicians, researchers, and patients; the formation of the Society for Improving Diagnosis in Medicine (SIDM); and convening of a broad coalition of organizations, including the American Board of Internal Medicine (ABIM), the American College of Physicians (ACP), and the Society of Hospital Medicine (SHM), committed to increasing awareness and action. Ultimately, collectively tackling the challenges of improving the quality of diagnosis will transform the way clinicians and patients work together to co-produce better diagnoses.

FURTHER READING

Gandhi TK, Singh H: Reducing the risk of diagnostic error in the COVID-19 era. *J Hosp Med* 15:363, 2020. [PubMed: 32490798]

Graber ML et al: The impact of electronic health records on diagnosis. *Diagnosis (Berl)* 4:211, 2017. [PubMed: 29536944]

National Academies of Sciences, Engineering, and Medicine. 2015. *Improving Diagnosis in Health Care*. <https://doi-org.kaplanmc.idm.oclc.org/10.17226/21794>. Adapted and reproduced with permission from the National Academy of Sciences, Courtesy of the National Academies Press, Washington, DC.

Newman-Toker DE et al: Serious misdiagnosis-related harms in malpractice claims: The "big three"—Vascular events, infections, and cancers. *Diagnosis (Berl)* 6:227, 2019. [PubMed: 31535832]

Schiff GD et al: Diagnosing diagnosis errors: Lessons from a multi-institutional collaborative project, in *Advances in Patient Safety: From Research to Implementation. Vol 2: Concepts and Methodology*. Rockville, MD, Agency for Healthcare Research and Quality, 2005.

Schiff GD et al: Ten principles for more conservative, care-full diagnosis. *Ann Intern Med* 169:643, 2018. [PubMed: 30285046]