

Diagnostic Stewardship: Opportunity for a Laboratory–Infectious Diseases Partnership

Robin Patel¹ and Ferric C. Fang²

¹Divisions of Clinical Microbiology and Infectious Diseases, Departments of Laboratory Medicine and Pathology and Medicine, Mayo Clinic, Rochester, Minnesota, and ²Departments of Laboratory Medicine and Microbiology, University of Washington School of Medicine, Seattle.

Recent advances in microbial diagnostics are providing clinicians with information about microbes causing infections and their resistance to antimicrobial agents more rapidly than ever before. Diagnostic stewardship refers to the appropriate use of laboratory testing to guide patient management, including treatment, in order to optimize clinical outcomes and limit the spread of antimicrobial resistance. Fulfilling the promise of diagnostic stewardship requires a seamless partnership between clinical laboratories, pharmacists, and infectious diseases clinicians, so that appropriate tests are ordered and diagnostic information is translated into appropriate management in real time.

Keywords. diagnostics; stewardship; clinical microbiology; infectious diseases; culture-independent.

Rapid precision diagnostics are revolutionizing clinical microbiology and promise to improve patient outcomes and curb the antimicrobial resistance (AMR) crisis by improving the use of antibiotics. For this potential to be fully realized, infectious diseases (ID) clinicians will play an essential role in a collaborative effort referred to as *diagnostic stewardship* (not to be confused with the cost-effective use of laboratory tests which, though part of diagnostic stewardship, is more limited in scope) [1–4]. Diagnostic stewardship requires a serious reconsideration of current practices, as empiricism gives way to diagnostics-guided therapy.

The goal of new diagnostic methods is to improve human health, but technological advances alone cannot achieve this goal. Decisions must be made about which new diagnostics are needed, how they will be used, and whether they are worth paying for. Laboratory tests currently account for only 4% of health-care costs [5] but represent the most rapidly growing segment of the healthcare budget, mainly as a result of new molecular assays [6]. Although conventional clinical microbiology diagnostics are relatively inexpensive, some newer and technologically advanced tests can be costly, elevating the need to address value. It is estimated that approximately one-fifth of available tests are overused, with even more being underused [7]. Overuse of tests adds unnecessary costs, and both overuse and underuse can lead to incorrect diagnoses and inappropriate treatment. Moreover, many microbiology tests have become

outdated, and optimal testing methods are unavailable in many settings. Appropriate use of testing is becoming more challenging as the number of available diagnostic tests increases. ID specialists can help to determine the appropriate tests for specific patients and situations.

ID clinicians currently partner with laboratory scientists to determine which antimicrobial susceptibility results are routinely reported for specific microorganisms and when additional testing should be performed. This facilitates antimicrobial stewardship by encouraging appropriate antibiotic use. With recent diagnostic advances that allow the identification of microorganisms virtually as soon as they are grown on plates or in blood culture bottles, and sometimes even earlier [8], ID expertise plays an essential role in translating this information into appropriate treatment.

Many studies have shown that rapid diagnostics only improve clinical outcomes if they are coupled with stewardship teams that properly interpret results and apply them to treatment decisions [9–20]. This approach may require expanding the hours of laboratory operation and providing real-time ID consultative support. ID physicians and pharmacists may be asked to work alongside their laboratorians on diagnostic management teams [21], or clinical microbial sequencing boards (modeled after tumor boards) (<https://www.genomeweb.com/sequencing/ucsf-lab-readies-launch-metagenomic-ngs-test-infectious-disease>) that assist clinicians with the interpretation of complex test results in a specific clinical field.

ID clinicians can assist laboratories in devising appropriate comments to accompany test results in the electronic medical record (EMR), such as “possible contaminant which may not require antibiotic treatment” when coagulase-negative staphylococci are reported from a single blood culture; such comments can be tailored to the needs and unique epidemiology of

Received 13 December 2017; editorial decision 19 January 2018; accepted 29 January 2018; published online March 14, 2018.

Correspondence: R. Patel, Division of Clinical Microbiology, Department of Laboratory Medicine and Pathology, Mayo Clinic, Rochester, MN 55905 (patel.rob@mayo.edu).

Clinical Infectious Diseases® 2018;67(5):799–801

© The Author(s) 2018. Published by Oxford University Press for the Infectious Diseases Society of America. All rights reserved. For permissions, e-mail: journals.permissions@oup.com. DOI: 10.1093/cid/ciy077

individual institutions [22]. In this way, the EMR can provide a solution to AMR. Integration of laboratories with the EMR, antimicrobial stewardship teams, clinical pharmacists, and clinicians can ensure that treatment decisions are appropriately modified in response to test results in real-time [13]. Likewise, detection of certain drug-resistant organisms can automatically trigger inpatient isolation, preventing the spread of these organisms.

Sophisticated diagnostics can augment antimicrobial stewardship efforts by allowing the replacement of broad-spectrum antimicrobial agents with narrow-spectrum agents that target the microbes specifically responsible for individual patients' infections, facilitating early discontinuation of antimicrobial agents, or abrogating their use in the first place. To support the safety and efficacy of this approach, more studies of the application of precision diagnostics to optimize patient outcomes and reduce AMR (ie, implementation science) will be needed. In the context of AMR, improvements at the patient level promise to provide benefits at the population level.

Test menu selection is another activity in which ID clinicians can partner with their laboratories. An ever-increasing number of powerful but expensive technologies, ranging from point-of-care molecular diagnostics [23, 24] to multiplex panels [25, 26] and next-generation sequencing methods [27], require careful and discerning application. Laboratories can benefit from clinical input to select cost-effective diagnostics that best address patient needs. Similarly, if diagnostics manufacturers market tests directly to ID clinicians, then ID clinicians should work with their clinical microbiologists to ensure that the needs of their patients are optimally served.

As new technologies become available, local laboratory methods and a catalog of send-out tests should continually be reassessed to ensure that patients benefit from the latest diagnostic advances; ID clinicians should help ensure that testing is both available and appropriately ordered (eg, by helping to build smart ordering systems in the EMR). For example, ID clinicians can provide guidance with regard to the appropriate clinical criteria for testing of patients with clinical syndromes, such as acute gastroenteritis [28–30] or suspected *Clostridioides difficile* infections [31, 32].

Finally, by defining important unmet diagnostic needs [33], ID clinicians will play an increasingly important role in defining the future tests that should be developed by industry. For example, the development of syndromic molecular diagnostic panels can benefit from clinical guidance [25, 26]. Molecular diagnostics allow the rapid and sensitive detection of pathogens that were not previously detectable with conventional methods [30], and ID expertise will be required to determine the implications of these diagnoses for specific management.

For example, it is not unusual for multiplex molecular platforms to detect multiple potential pathogens in a single clinical sample [25]. ID physicians can help interpret apparent

coinfections with multiple potential pathogens and determine when a pathogen is likely to be responsible for a patient's symptoms, as well as establish appropriate criteria for the use of multiplex tests and assist in the design of such assays so that appropriate target organisms are included [26]. Newer technologies, such as whole-genome sequencing, shotgun metagenomic methods to diagnose infection, and methods to characterize the host microbiome [34, 35] likewise pose both an opportunity and a challenge, and ID clinicians can help establish interpretive criteria and applications.

Achieving a clinician-laboratorian collaboration will not necessarily be simple. The consolidation of laboratory services [36] has created obstacles for direct clinician-laboratory interactions at a time when such interactions are needed more than ever. In addition, new clinical guidelines and testing algorithms will need to keep pace with the development of novel diagnostic methods, which will require the input of both laboratory scientists and ID clinicians (eg, on guidelines panels).

In a recent commentary in *Clinical Infectious Diseases* [37], Arturo Casadevall pointed out that the role of ID specialists has historically been to provide “intellectual input in the form of consultation.” Nowadays he suggests that ID specialists should “use (their) expertise to command an important position in the information and decision flows in medicine” but worries that empiricism in the use of antimicrobial agents has “fostered a neglect of new diagnostics.” The current diagnostics revolution promises to transform clinical practice to more closely conform with Casadevall's vision of diagnostics-driven therapy for ID.

Developers of new diagnostic technologies will be best served by a team-based approach. ID specialists are ideal partners to develop and implement systems to ensure appropriate diagnostic testing and the seamless translation of laboratory results into personalized treatment. Improved diagnostics may increase the costs of diagnostics, so assessing value will become increasingly important, with regard to both specific tests and approaches to patient care. Diagnostic stewardship means selecting the right test for the right patient at the right time, to optimize clinical care and antimicrobial use [4]. This is a mission for ID specialists and clinical microbiologists to take on together.

Notes

Financial support. This work was supported by the National Institute of Allergy and Infectious Diseases, National Institutes of Health (NIH) (grants UM1 AI104681, R01 AR056647, and R21 AI125870 to R. P.) and the National Institute of Arthritis and Musculoskeletal and Skin Diseases, NIH (grant R01 AR056647 to R. P.).

Potential conflicts of interest. R. P. has participated in research studies supported by CD Diagnostics, BioFire, Curetis, Merck, Hutchison Biofilm Medical Solutions, Accelerate Diagnostics, Allergan, and The Medicines Company; is a consultant to Curetis Specific Technologies, Selux Dx, GenMark Diagnostics, PathoQuest and Genentech (monies are paid to Mayo Clinic); receives travel reimbursement and an editor's stipend from the American Society for Microbiology and honoraria from the National Board of Medical Examiners, Up-to-Date, and the Infectious Diseases Board Review Course; has patents issued on *Bordetella pertussis*/

parapertussis PCR and an anti-biofilm substance and a patent on a device/method for sonication with royalties paid by Samsung to Mayo Clinic; and has served on an Actelion data monitoring board. F. C. F. has participated in research studies supported by BioFire, Cepheid, ELITech, and Luminex (formerly Nanosphere). R. P. and F. C. F. both receive editors' stipends from the Infectious Diseases Society of America. Both authors have submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest. Conflicts that the editors consider relevant to the content of the manuscript have been disclosed.

References

- Goff DA, Jankowski C, Tenover FC. Using rapid diagnostic tests to optimize antimicrobial selection in antimicrobial stewardship programs. *Pharmacotherapy* **2012**; 32:677–87.
- Avdic E, Carroll KC. The role of the microbiology laboratory in antimicrobial stewardship programs. *Infect Dis Clin North Am* **2014**; 28:212–35.
- Morgan DJ, Malani P, Diekema DJ. Diagnostic stewardship—leveraging the laboratory to improve antimicrobial use. *JAMA* **2017**; 318:607–8.
- Messacar K, Parker SK, Todd JK, Dominguez SR. Implementation of rapid molecular infectious disease diagnostics: the role of diagnostic and antimicrobial stewardship. *J Clin Microbiol* **2017**; 55:715–23.
- Riley SB. Trends in laboratory utilization. *Clin Microbiol Newslett* **2017**; 39:69–73.
- Brown S, Dickerson J. The struggle is real: lab leaders discuss utilization challenges during a 2-day summit. *J Appl Lab Med* **2016**; 1:306–9.
- Zhi M, Ding EL, Theisen-Toupal J, Whelan J, Arnaout R. The landscape of inappropriate laboratory testing: a 15-year meta-analysis. *PLoS One* **2013**; 8:e78962.
- Perlin DS, Wiederhold NP. Culture-independent molecular methods for detection of antifungal resistance mechanisms and fungal identification. *J Infect Dis* **2017**; 216:458–65.
- Bauer KA, West JE, Balada-Llasat JM, Pancholi P, Stevenson KB, Goff DA. An antimicrobial stewardship program's impact with rapid polymerase chain reaction methicillin-resistant *Staphylococcus aureus*/S. aureus blood culture test in patients with S. aureus bacteremia. *Clin Infect Dis* **2010**; 51:1074–80.
- Huang AM, Newton D, Kunapuli A, et al. Impact of rapid organism identification via matrix-assisted laser desorption/ionization time-of-flight combined with antimicrobial stewardship team intervention in adult patients with bacteremia and candidemia. *Clin Infect Dis* **2013**; 57:1237–45.
- Bauer KA, Perez KK, Forrest GN, Goff DA. Review of rapid diagnostic tests used by antimicrobial stewardship programs. *Clin Infect Dis* **2014**; 59:S134–45.
- Perez KK, Olsen RJ, Musick WL, et al. Integrating rapid diagnostics and antimicrobial stewardship improves outcomes in patients with antibiotic-resistant Gram-negative bacteremia. *J Infect* **2014**; 69:216–25.
- Banerjee R, Teng CB, Cunningham SA, et al. Randomized trial of rapid multiplex polymerase chain reaction-based blood culture identification and susceptibility testing. *Clin Infect Dis* **2015**; 61:1071–80.
- Lockwood AM, Perez KK, Musick WL, et al. Integrating rapid diagnostics and antimicrobial stewardship in two community hospitals improved process measures and antibiotic adjustment time. *Infect Control Hosp Epidemiol* **2016**; 37:425–32.
- Patel TS, Kaakeh R, Nagel JL, Newton DW, Stevenson JG. Cost analysis of implementing matrix-assisted laser desorption ionization-time of flight mass spectrometry plus real-time antimicrobial stewardship intervention for bloodstream infections. *J Clin Microbiol* **2017**; 55:60–7.
- Minejima E, Wong-Beringer A. Implementation of rapid diagnostics with antimicrobial stewardship. *Expert Rev Anti Infect Ther* **2016**; 14:1065–75.
- Timbrook TT, Morton JB, McConeghy KW, Caffrey AR, Mylonakis E, LaPlante KL. The effect of molecular rapid diagnostic testing on clinical outcomes in bloodstream infections: a systematic review and meta-analysis. *Clin Infect Dis* **2017**; 64:15–23.
- Beganovic M, Costello M, Wiczorkiewicz SM. Effect of matrix-assisted laser desorption ionization-time of flight mass spectrometry (MALDI-TOF MS) alone versus MALDI-TOF MS combined with real-time antimicrobial stewardship interventions on time to optimal antimicrobial therapy in patients with positive blood cultures. *J Clin Microbiol* **2017**; 55:1437–45.
- Rivard KR, Athans V, Lam SW, et al. Impact of antimicrobial stewardship and rapid microarray testing on patients with gram-negative bacteremia. *Eur J Clin Microbiol Infect Dis* **2017**; 36:1879–87.
- Eby JC, Richey MM, Platts-Mills JA, Mathers AJ, Novicoff WM, Cox HL. A healthcare improvement intervention combining nucleic acid microarray testing with direct physician response for management of *Staphylococcus aureus* bacteremia. *Clin Infect Dis* **2018**; 66:64–71.
- Laposata M. The what and why of diagnostic management teams. *CAP Today, January 2017*. Available at: <http://www.captodayonline.com/diagnostic-management-teams/>. Accessed February 13, 2018.
- Banerjee R, Ozenci V, Patel R. Individualized approaches are needed for optimized blood cultures. *Clin Infect Dis* **2016**; 63:1332–9.
- Patel R, Karon BS. Advances afoot in microbiology. *J Clin Microbiol* **2017**; 55:1984–8.
- Simner PJ, Miller S, Carroll KC. Understanding the promises and hurdles of metagenomic next-generation sequencing as a diagnostic tool for infectious diseases. *Clin Infect Dis* **2018**; 66:778–88.
- Hanson KE, Couturier MR. Multiplexed molecular diagnostics for respiratory, gastrointestinal, and central nervous system infections. *Clin Infect Dis* **2016**; 63:1361–7.
- Ramanan P, Bryson A, Binnicker M, Pritt B, Patel R. Syndromic panel-based testing in clinical microbiology. *Clin Microbiol Rev* **2018**; 31:1–28.
- Ellington MJ, Ekelund O, Aarestrup FM, et al. The role of whole genome sequencing in antimicrobial susceptibility testing of bacteria: report from the EUCAST Subcommittee. *Clin Microbiol Infect* **2017**; 23:2–22.
- Riddle MS, DuPont HL, Connor BA. ACG clinical guideline: diagnosis, treatment, and prevention of acute diarrheal infections in adults. *Am J Gastroenterol* **2016**; 111:602–22.
- Shane AL, Mody RK, Crump JA, et al. 2017 Infectious Diseases Society of America clinical practice guidelines for the diagnosis and management of infectious diarrhea. *Clin Infect Dis* **2017**; 65:e45–80.
- Fang FC, Patel R. 2017 Infectious Diseases Society of America infectious diarrhea guidelines: a view from the clinical laboratory. *Clin Infect Dis* **2017**; 65:1974–6.
- Truong CY, Gombar S, Wilson R, et al. Real-time electronic tracking of diarrheal episodes and laxative therapy enables verification of *Clostridium difficile* clinical testing criteria and reduction of *Clostridium difficile* infection rates. *J Clin Microbiol* **2017**; 55:1276–84.
- Fang FC, Polage CR, Wilcox MH. Point-counterpoint: what is the optimal approach for detection of *Clostridium difficile* infection? *J Clin Microbiol* **2017**; 55:670–80.
- Blaschke AJ, Hersh AL, Beekmann SE, Ince D, Polgreen PM, Hanson KE. Unmet diagnostic needs in infectious disease. *Diagn Microbiol Infect Dis* **2015**; 81:57–9.
- Goldberg B, Sichtig H, Geyer C, Ledebner N, Weinstock GM. Making the leap from research laboratory to clinic: challenges and opportunities for next-generation sequencing in infectious disease diagnostics. *MBio* **2015**; 6:e01888–15.
- Besser J, Carleton HA, Gerner-Smidt P, Lindsey RL, Trees E. Next-generation sequencing technologies and their application to the study and control of bacterial infections. *Clin Microbiol Infect* **2017**. doi:10.1016/j.cmi.2017.10.01
- Sautter RL, Thomson RB Jr. Consolidated clinical microbiology laboratories. *J Clin Microbiol* **2015**; 53:1467–72.
- Casadevall A. Crisis in infectious diseases: 2 decades later. *Clin Infect Dis* **2017**; 64:823–8.