

Strategies to Prioritize Clinical Options in Primary Care

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Ann Fam Med 2017;15:10-13. <https://doi.org/10.1370/afm.2027>.

In 2003, McGlynn and colleagues famously identified 439 clinical services recommended by subspecialists and others that, they argue, should be consistently delivered in primary care.¹ Based on lengthy patient interviews and detailed chart audits, they estimated that US adults receive only 55% of these 439 recommended clinical services. They noted that the patients did not receive recommended clinical services, on average, 16 times, and that 1 patient was deficient in recommended services 304 times.

Yarnall et al reviewed US Preventive Services Task Force-recommended preventive services and estimated that a primary care clinician would need to spend 7.4 hours per working day to deliver recommended preventive services to a panel of patients²; doing so would, of course, leave little time for anything else and lead to frustrated clinicians and dissatisfied patients. What is a primary care clinician to do? We are damned if we do not deliver all evidence-based preventive and chronic disease care services with robot-like consistency and sleepless if we do. Proposed remedies include off-loading delivery of preventive or chronic disease care services to office staff, nurse case managers, subspecialty clinics, or even attractive-looking multicolored smartphone apps.

The Case for Prioritization

Perhaps a more reasonable and less-expensive strategy would be to prioritize clinical services related to preven-

tive care and chronic disease care. In this issue of *Annals of Family Medicine*, Maciosek et al present compelling evidence that childhood immunizations and efforts to prevent or stop smoking are dominant population health priorities and rank numerous other services based on potential clinical benefit and cost-effectiveness.³⁻⁵ Once smoking cessation and immunizations are addressed, however, we are left wondering how to prioritize the other 439 evidence-based clinical services based on their potential benefit to an individual patient.

There are several fundamentally sound reasons to prioritize clinical services at the patient level. First, the value of even very strongly evidence-based clinical services varies across patients and with time. For example, the potential benefit of screening for colorectal, lung, cervical, and breast cancer varies up to tenfold based on patient-specific demographic, clinical, behavioral, and genetic factors.⁶ Likewise, the risks and benefits of intensive glucose control in patients with diabetes vary by age, comorbid conditions, cardiovascular risk, distance from personalized glycosylated hemoglobin (HbA_{1c}) goal, and other factors. If an older patient with major comorbidities already on intensive glucose-lowering therapy is not at their personalized HbA_{1c} goal, the risks of further intensifying glucose therapy may well exceed the benefits.^{7,8} The ranks provided by Maciosek et al, which are based on overall population health benefit, must be further personalized to assess relative benefit of these services to an individual patient.

Second, in very large clinical trials, very small clinical benefits may be statistically significant and thus be designated as being evidence based. With respect to cholesterol-lowering medications, the number of study participants who need to be treated with a statin for 5 years to prevent 1 heart attack can vary from 6 patients to more than 240 patients, depending on baseline cholesterol level and baseline coronary heart disease risk.^{9,10}

Conflicts of interest: authors report none.

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In general, the number of patients who need to be treated to prevent 1 heart attack or 1 cancer death varies widely across various evidence-based clinical options.

Third, most clinical trials that assess efficacy of clinical options are limited to consenting patients who meet stringent eligibility requirements. Results observed in these highly selected patients may have low generalizability to most patients. Moreover, because patient eligibility criteria and research protocols are designed to minimize the likelihood of serious adverse events in clinical trials, serious adverse events may be much more common in community care than in published research studies.¹¹

How to Prioritize Evidence-Based Clinical Options

Primary care clinicians have always intuitively prioritized treatment options, but both benefits and risks of treatment are often not estimated accurately. For some clinical services, such as smoking cessation and certain immunizations, intuition is adequate. But beyond smoking and immunizations, intuitive estimation of potential benefit of multiple clinical options is very challenging.

Several alternative methods are available to identify and prioritize evidence-based clinical options with the most potential benefit to a given patient at a given point in time.¹² With respect to cardiovascular risk factor management, risk prediction equations, such as the American College of Cardiology/American Heart Association (ACC/AHA) Cardiovascular Disease risk equations,^{13,14} can be used to estimate the benefit of various clinical actions using the following 3-step approach: (1) use the risk equation to estimate a person's cardiovascular risk using current clinical data, (2) run the risk equation again, replacing 1 suboptimal clinical value (eg, an elevated blood pressure) with a potentially improved clinical value (anticipated improved blood pressure after treatment), and (3) subtract the results to estimate the potential reduction in cardiovascular risk that may be achieved by better blood pressure control. The potential benefits associated with better blood pressure management, cholesterol management, smoking cessation, or other clinical options can then be similarly estimated and then prioritized based on potential clinical benefit.^{15,16}

This approach to prioritization has a number of important limitations. Risk estimates are necessarily derived from groups of people and thus cannot precisely predict future risk for one person. The benefits of stopping smoking are not the same as the benefits of never having smoked. The full benefits of improved cardiovascular risk factor control do not kick in immediately, and benefit estimates assume that the improved risk factor control will be sustained.

Current cardiovascular risk prediction tools are based on relatively small cohort studies that began in the 1950s, when many current drug classes were not available, aspirin use was low, smoking rates were high, and cardiac care was primitive when judged by today's standards.¹⁷ For these and other reasons, the ACC/AHA and most other cardiovascular risk equations are somewhat obsolete and tend to overestimate event and death rates.¹⁸ Despite such limitations, explicit estimation of benefits and risks of treatment options is usually far more accurate than intuitive risk estimation by either clinician or patient.¹⁹ The availability of large databases that include detailed clinical data on millions of patients and novel analytic approaches, such as marginal structural models and machine learning, will likely lead to improved risk prediction and prioritization methods in the near future.²⁰⁻²² It may be difficult, however, to explain these complex statistical approaches to clinicians and patients, who thus may be skeptical of their results.

How to Apply Prioritization in Practice

The potential of electronic health records (EHRs) to improve care has long been recognized but rarely been realized. Prototype EHR-linked, Web-based clinical decision support systems that identify and prioritize clinical options, however, save time, satisfy clinicians, empower patients, have high use rates, and improve care are now up and running in several large health care systems.^{15,16} Web services that include risk prediction equations can receive patient-specific data that are automatically sent from an EHR, perform the multiple computations needed to estimate the relative benefits of alternative treatment options, and display patient-specific prioritized treatment options on the EHR screen within 1 second.

Presenting clinical options to the patient facilitates patient-centered care and shared decision making by informing the patient of clinical options with the most potential benefit and then empowering the patient to select their preferred option(s). Many patients will continue to decline clinical options of high benefit, such as smoking cessation, colorectal cancer screening, or statin treatment. Then we must respect our patient's preferences and remember that patient treatment preferences and readiness to change typically vary with time.²³ Clinical decision support systems update and reprioritize evidence-based treatment options at each subsequent encounter, enabling patients to see progress in some areas and reconsider previous preferences in other areas.

Results of randomized trials show that systems improve blood pressure levels and glucose control in diabetes patients, smoking cessation in dental offices, identification of high blood pressure levels in adoles-

cents, and screening for hepatitis B in high-risk populations, as well as reduce cardiovascular risk in adults at high risk but without a diagnosis of heart disease or diabetes.¹⁶ There is also evidence that such systems are cost-effective and may be cost saving to payers when used on a routine basis in large care delivery systems.²⁴ These clinical decision support systems are used at 70% to 80% of targeted visits, have 94% primary care clinician satisfaction, and are now in use every day at 3 large health care delivery systems that provide care to 1.5 million patients.

Future Challenges

An ongoing major challenge is how to present quantitative risk and benefit information to patients in a comprehensible way. Health literacy and numeracy vary widely across patients, suggesting that presentation of information on potential risks and benefits of clinical actions should be customized to specific groups of patients. Development of effective strategies to clearly communicate risk and benefit information to those with low numeracy is very much a work in progress, and there is plenty of room for new ideas on how to advance this agenda.²⁵

Another ongoing challenge is to develop prioritization methods that can compare benefits across diverse clinical domains. Will a patient who does not like to take a lot of pills benefit more from starting a statin or treating osteoporosis? Prioritizing across diverse clinical domains is challenging because the benefits of lipid and osteoporosis management are very different (reduced risk of a cardiovascular event or death on the one hand and reduced likelihood of fracture and disability on the other). The traditional resolution of this problem is to quantify all benefits in terms of quality-adjusted life expectancy (QALE). Neither clinicians nor patients, however, are usually fluent in the language of QALE, and benefits of even very effective treatments on QALE are often surprisingly small. For example, among patients with type 2 diabetes in the United Kingdom Prospective Diabetes Study study,²⁷ intensive lipid control extends QALE 1.42 years,²⁶ and intensive blood pressure control extends QALE about 1.16 years,²⁶ but intensive glucose control extends QALE by only 0.27 years, and it did not improve QALE at all in the ACCORD Trial.^{8,28}

Recent advances in health care informatics and risk prediction methods enable design of new and more effective types of EHR-linked, Web-based, real-time clinical decision support systems that have high use rates at targeted visits, have high clinician satisfaction rates, and improve patients' clinical outcomes. We anticipate that further progress may occur as risk prediction science improves, better methods of com-

municating results to patients in customized ways are devised, and ways of prioritizing clinical options across a broader set of clinical domains are developed.^{3-5,13,29}

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Key words: disease, prevention & control; health services; economics; prioritization; health impact; cost-effectiveness; cost-savings; immunization; mass screening; behavioral counseling

Submitted November 18, 2016; submitted, revised, November 18, 2016; accepted November 29, 2016.

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