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Health Professions Building, Department
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PUBLISHING OFFICES

Dowden Health Media, Inc,
110 Summit Avenue,
Montvale, NJ 07645
Telephone: (201) 740-6193
Fax: (201) 740-6226



David L. Hahn, MD, MS
Department of Family Practice,
Dean Medical Center,
Madison, Wis

Public reporting needs reform!

Like many of my colleagues, I support President Obama's call to demonstrate value as part of health care reform. One way to do that is through public reporting. The rationale is that public scrutiny of outcomes will motivate the health care "industry" to improve the "product" (outcomes), rather than accelerating valueless economic activity (process) that often benefits providers more than patients.

Fair enough. But does the existing system of quality indicators support the goals of reform identified by the Institute of Medicine (IOM)?¹ That is, does it make the system safer, more effective and efficient, timely, equitable, and patient-centered?

Not necessarily.

The reason is 2-fold. First, the best quality indicators are patient-oriented outcomes (eg, quality of life, morbidity, mortality), but that's not what's being reported. Second, many publicly reported surrogate measures are more harmful than helpful, and in need of serious reform themselves.

My experience

The Wisconsin Collaborative for Healthcare Quality (WCHQ), which I've been involved with for nearly 10 years, is composed of health care organizations, mine included, committed to voluntary reporting of quality metrics. The **TABLE** features a list of the metrics, chosen by the WCHQ, that are reported.

I've rated each metric on 2 criteria:

1. How good is the evidence for the screening tool or intervention? (There is good evidence for colorectal cancer screening, for example, but evidence for low-density lipoprotein [LDL] testing is poor.)
2. How good is the quality indicator itself, including the frequency? (There's good evidence for Pap testing within 3 years, whereas twice-yearly HbA1c testing is opinion-based.)

Some worrisome examples

While the ratings are partly subjective, they're meant to illustrate that not all publicly reported metrics are supported by good evidence.

This is particularly troubling, given the fact that acting on fair or poor evidence may cause more harm than good. Consider these worrisome examples:

LDL control <100 mg/dL. I've known patients who had their first myocardial infarction when their LDL cholesterol was <100 mg/dL. After the event, these patients weren't given a statin because they were already "at goal"; they subsequently had a reinfarction.

LDL should not be used as a quality indicator in secondary prevention for (at least) 2 reasons: First, some LDL-lowering drugs are harmful or have no net benefit (eg, estrogen in women, fibrates).² Second, statin benefit may or may not be

TABLE

How do the “quality indicators” rate?

SCREENING TOOL/INTERVENTION (RECOMMENDED METRIC)	SCREENING TOOL SOR	METRIC SOR
Colorectal cancer screening (various modalities and frequencies)	A	A
Pap smear (within 3 years)	A	A
Tobacco use (documented in the past year)	A	B
DM2: BP control (last BP <130/80)	A	B
BP control in nondiabetics (last BP <140/90)	A	B
DM2: HbA1c testing (at least twice yearly)	B	C
DM2: blood sugar control (HbA1c <7)	B	C
Pneumococcal vaccine (once after age 65)	B	C
Mammography (within 2 years, women ages 40-69)	B	C
Postpartum care (21-56 days after delivery)	C	C
DM2: kidney function monitored (creatinine yearly)	C	C
CVD: LDL testing (yearly)	C	C
CVD: LDL control (LDL<100 mg/dL)	C	C
DM2: LDL testing (yearly)	C	C
DM2: LDL control (LDL<100 mg/dL)	C	C

BP, blood pressure; CVD, cardiovascular disease; DM2, type 2 diabetes; HbA1c, glycosylated hemoglobin; LDL, low-density lipoproteins; SOR, strength of recommendation.

Strength of recommendation (SOR):

A Good-quality patient-oriented evidence

B Inconsistent or limited-quality patient-oriented evidence

C Consensus, usual practice, opinion, disease-oriented evidence, case series

Source: Wisconsin Collaborative for Healthcare Quality (<http://www.wchq.org/>).

FAST TRACK

Many publicly reported surrogate measures are more harmful than helpful.

related to lipid lowering, and the magnitude of benefit is not related to any arbitrary LDL goal.³

There is clear, compelling evidence supporting near-universal statin therapy for patients at high cardiovascular risk regardless of their LDL cholesterol values—but a lack of evidence that titrating lipid therapy to achieve proposed low LDL levels is beneficial or safe.⁴ Receiving the maximum tolerated dose of statin is therefore the appropriate evidence-based surrogate quality indicator, *not* LDL.

Mammography. The Cochrane collaboration has concluded that “for every 2000 women invited for [mammography] screening throughout 10 years, 1 will have her life prolonged,” and 10

healthy women who would not have been diagnosed without the screening will be treated unnecessarily.⁵ The Cochrane review thus concluded that it’s not clear whether mammography screening does more good than harm.⁵

The US Preventive Services Task Force (USPSTF) recently downgraded mammography screening for women over age 50 from an A- to a B-rated recommendation. The fine balance between benefit and harm in this and other USPSTF B-rated preventive measures requires that clinicians educate patients and elicit their preferences. But this doesn’t occur when health plans strive to outdo one another in achieving higher publicly reported screening goals.⁶ Documentation

of valid shared decision-making, not screening rates, is the appropriate quality indicator.

The evidence vs the “business” of medicine

I have no illusions that my recommendations will be adopted easily—or soon. After all, we practice in an environment in which evidence-based practice recommendations can conflict with financial and operational goals perceived as necessary to survive. However, I believe that evidence trumps business in achieving the IOM goals.

It remains to be seen whether we can simultaneously move toward valid evidence-based public reporting and health-care financial reform. But one thing is clear: To insist that evidence-based patient-oriented quality indicators are too difficult to measure, or to ignore or deny

the evidence, puts the lie to claims of patient-centered care and, ultimately, to long-needed health care reform looming on the horizon. ■

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Incretin-related therapies:

Targeting the underlying physiology and cardiometabolic factors of type 2 diabetes

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