

Epocrates MedInsight[®] Treatment Trends in Type 2 Diabetes The Evolving Role of A1c, Metformin & Newer Medications

For more information, contact payerpartnerships@epocrates.com

Treatment Trends in Type 2 Diabetes The Evolving Role of A1c, Metformin & Newer Medications

Abstract

While prescribing trends reflect physician behaviors, understanding the attitudes and perceptions determining medication choices requires additional research.

Expenditures for prescription medications for type 2 diabetes have risen three fold in the past 10 years from 6.7 billion (2001) to 818.9 billion (2011).^{1,2}

This increase may be associated with a number of trends, including fewer patients on monotherapy (82% in 1994 vs. 47% in 2007), a decline in the use of sulfonylureas (64% in 1994 vs. 34% in 2007) and the introduction of newer therapies.² According to National Disease and Therapeutic Index data, DPP-4 inhibitors are the fastest growing oral drug class for type 2 diabetes.¹ Epocrates TapStream, which monitors aggregated drug lookup patterns of Epocrates' physicians, also reflects this trend.

While prescribing trends reflect physician behaviors, such data does not reveal the underlying attitudes and perceptions determining medication choices. To better understand the thinking behind type 2 diabetes prescribing decisions, Epocrates surveyed 50 primary care physicians on topics ranging from A1c targets, the ACCORD (Action to Control Cardiovascular Risk in Diabetes) trial and the importance of blood pressure control.



Epocrates TapStream Data^{*} Diabetes Drug Class LookUps (September - February)

* Epocrates TapStream data reflects number of times clinicians using Epocrates Rx looked up certain classes of type 2 diabetes drugs. Epocrates TapStream data does not represent actual prescribing data.

Findings: Highlights from the Epocrates MedInsight survey appear below.



of physicians agreed that A1c ought to be tightly controlled (<6.5) for most patients. **This perception is inconsistent with 2010/2011 Veterans Administration (VA) and American Diabetes Association (ADA) guidelines.**



physicians agreed that the combination of metformin + glipizide is to be avoided due to high degree of hypoglycemia. **This perception is consistent with marketing claims.**



of physicians agreed that the ACCORD trial showed that intensive glucose control reduces cardiovascular mortality. This perception is inconsistent with ACCORD results showing that intensive control increased mortality and did not significantly reduce major cardiovascular events.



of physicians agreed that metformin is contraindicated for diabetes patients with mild renal impairment due to the potential for lactic acidosis. **This is inconsistent with published evidence from the past ten years.**

Treatment Trends in Type 2 Diabetes The Evolving Role of A1c, Metformin and Newer Medications

Participant Profile

We surveyed a total of 50 primary care physicians using the Epocrates MedInsight platform. Participating physicians treated at least 10 type 2 diabetes patients weekly and wrote 16 or more type 2 diabetes prescriptions per month.

We surveyed doctors from two distinct geographic regions to assess whether there would be geographic variation in physician responses. Physicians surveyed represented various practice settings.



Methods

Physicians were presented with statements concerning type 2 diabetes (questions 1 through 8) and asked to what extent they disagreed or agreed using a 4-part Likert Scale – Strongly Disagree, Disagree, Agree and Strongly Agree. For purposes of analysis, responses were combined in a binary fashion (agree or disagree) and presented as percentages. Question 9 uses a multiple choice format.

Q1 Current evidence suggests that aggressive targets for HbA1c (<6.5%) are desirable for most patients.

74% of total physicians surveyed agreed with this statement, with 26% of respondents disagreeing. The majority response is inconsistent with current evidence and recent guidelines.³⁻⁵ Current ADA and VA guidelines, which are based on evidence from multiple clinical studies, recommend individualizing A1c target (range 7% to 9%) depending on factors including physiologic age, presence/severity of major comorbidities and duration of diabetes.^{3,4} For frail older adults and individuals with a life expectancy of less than five years, the American Geriatrics Society suggests an A1c target of 8 percent.⁶



QZ

Metformin is the first drug of choice for the great majority of diabetes patients.

92% of total physicians surveyed agreed with this statement, with 8% of respondents disagreeing. The majority responded in a manner that is consistent with evidence and guidelines showing that metformin is the preferred first-line oral agent for treatment of type 2 diabetes.⁷



A Metformin + sulfonylurea is the preferred oral combination for patients who do not have adequate glycemic control on monotherapy with either drug.

The majority of respondents agreed with this statement which is consistent with recommendations.⁷ However, 20% of the physicians in the Northeast and 40% of the doctors in the South disagreed.



Q4 The ACCORD trial demonstrated that intensive glycemic control reduced CVD mortality in patients with Type 2 diabetes.

70% of total physicians surveyed agreed with the statement. However, the statement is incorrect; ACCORD did not demonstrate reduced CVD mortality with intensive control.

In ACCORD, 10,250 patients with long-standing type 2 diabetes were randomly assigned to intensive or standard glycemic control.⁸ After a median follow-up of 3.7 years, intensive therapy was stopped due to a higher number of total and cardiovascular deaths in subjects assigned to intensive therapy compared with the standard treatment group.⁸



Q5 Januvia (sitagliptin) + metformin may be preferred to metformin + glipizide to avoid hypoglycemia.

94% of respondents agreed with this statement, which is made in marketing materials. This claim is based on a clinical trial sponsored, and a paper authored, by industry.⁹ In this paper, patients taking metformin + glipizide experienced rates of hypoglycemia higher than



The risk of lactic acidosis with metformin use is substantially elevated for diabetes patients with mild renal impairment (creatinine 1.3-1.8 mg/dL).

68% of total physicians surveyed agreed with this statement; however there appears to be little evidence to support this.^{11,12}

In a 2010 Cochrane Review, 324 (97%) of the 334 prospective studies allowed for the inclusion of patients taking metformin with at least one contraindication, including renal insufficiency. Analysis of these trials and studies (as well as previous Cochrane Reviews) showed no increased risk of lactic acidosis, or increased level of lactate, for metformin compared to other agents.¹¹ In another study of 393 patients with chronic kidney disease (plasma creatinine levels of 1.5 to 2.5 mg/dl), no cases of lactic acidosis occurred over the four year trial duration.¹²



An ARB is preferred to an ACEI for treating hypertension in most patients with Type 2 diabetes.

88% of total physicians surveyed disagreed with this statement, which indicates that the majority of physicians' responses were consistent with evidence. The benefits of ACE inhibitors in patients with diabetes and hypertension are well established, with strong evidence demonstrating their beneficial effects on multiple adverse outcomes, including both macrovascular and microvascular complications.¹³



Q8

For my patients with Type 2 diabetes, good blood pressure control takes priority over getting HbA1c <7%.

64% of respondents disagreed with this statement. However, according to the 2010 VA guidelines, BP control ought to take priority over getting a low A1c in patients with type 2 diabetes.⁴ While glucose control early in the course of diabetes appears to confer some long-term benefit in CVD risk reduction, research clearly indicates that achieving hypertension and dyslipidemia targets confers the greatest benefit.^{4,13}



Q9 In clinical trials, better glycemic control has been associated with reductions in which of the following outcomes?

50% of total physicians surveyed indicated that better glycemic control is associated with both reduced microvascular and macrovascular advantages. This is inconsistent with the evidence which shows that intensive glycemic control is only proven to reduce the risk for diabetes microvascular and neuropathic disease.¹⁴

In contrast, most randomized clinical trials show that intensive therapy does not improve macrovascular outcomes in patients with type 2 diabetes.⁵ VADT (Veterans Affairs Diabetes Trial), ACCORD, and ADVANCE (Action in Diabetes and Vascular Disease: Preterax and Diamicron Modified Release Controlled Evaluation) were all designed to study the impact of intensive vs. conventional therapy on cardiovascular outcomes in patients with long-standing type 2 diabetes (duration 8 to 12 years). None of these trials showed reductions in cardiovascular risk with intensive therapy.⁵ ACCORD, in fact, showed that intensive therapy was associated with significant increases in total and CVD mortality in patients with type 2 diabetes (mean baseline A1c of 7.9%) randomized to intensive (n=18,315) or standard (n= 16,218) treatment, there were no reductions in all-cause mortality or cardiovascular causes of death with intensive treatment and more than a doubling in severe hypoglycemia.¹⁵



Conclusion

According to the Institute of Medicine, it takes "an average of 17 years for new knowledge generated by randomized controlled trails to be incorporated into practice, and even then application is highly uneven."¹⁶ Factors influencing a slow uptake in evidence-based practice include: 1) lack of consistent knowledge and adoption of clinical guidelines, 2) lack of balanced sources of summarized evidence, 3) the inherent bias of patients and prescribers towards use of new therapies, and 4) insulation of consumers and prescribers from health care costs.¹⁷ Numerous reasons have been sought to explain this evidence "adoption gap"- the extended time it takes for research to be incorporated into physician prescribing practices.¹⁸ One key reason is the challenge of transferring evidence-based information to practicing clinicians. This problem arises from information overload and the growing complexity of research findings.¹⁸

Our survey of physician knowledge of type 2 diabetes treatment suggests that a) many physicians are unfamiliar with newer guidelines, b) older evidence (> 15 years) is more likely to be incorporated into practice, c) physician concerns around certain drugs may be unfounded, and 4) physicians have strong recall of claims made in marketing materials.

Physician respondents were unfamiliar with new A1c target goals reflected in recent guidelines. Additionally, over two thirds of respondents misunderstood the results of the ACCORD trial, leading to a perception that cardiovascular mortality could be reduced through intensive glycemic control. Data from the ACCORD trial, in fact, suggests that intensive control of A1c can be unsafe, particularly in patients with a long history of diabetes who are at high risk for cardiovascular disease.⁵

Recommendations such as those for treatment of hypertension in patients with diabetes appear to have better diffusion and adoption. 88% of physicians surveyed were aware, for example, that angiotensin converting enzyme (ACE) inhibitors are as effective as an ARB for treating hypertensive diabetic patients, with benefits that are well established and have been aggressively disseminated by the Joint National Commission.^{18,19}

The majority of physicians saw metformin use as contraindicated in patients with renal impairment due to concerns regarding lactic acidosis. However, the evidence shows that diabetic patients who are treated with metformin and who tolerate it well may continue taking it, even when mild renal impairment develops.^{11,12}

Many attempts to rectify the paradox of high cost/low quality practice have failed due to a failure to address the complex behavioral, cultural, and social contexts of professional practice.²⁰ New strategies for communicating evidence to physicians are required to accelerate adoption of optimal prescribing practices.

References

- 1. National Prescription Audit Plus[™] (NPA). Collegeville, PA: IMS Health; 2011.
- 2. Alexander GC, Seghal NL, Moloney RM, Stafford RS. National trends in treatment of type 2 diabetes mellitus, 1994–2007. *Arch Intern Med.* 2008;168(19):2088-2094.
- American Diabetes Association. Standards of medical care in diabetes—2011. *Diabetes Care*. 2011; 34(suppl 1):S11-S61.
- 4. US Department of Veteran Affairs. VA/DoD clinical practice guideline for the management of diabetes mellitus. http://www.healthquality.va.gov/diabetes/DM2010_FUL-v4e.pdf. Updated August 2010. Accessed April 2, 2012.
- 5. McCulloch DK. Glycemic control and vascular complications in type 2 diabetes mellitus. In: UpToDate, Mulder, JE (Ed), UpToDate, Waltham, MA, 2012.
- 6. Huang ES, Zhang Q, Gandra N, Chin MH, Meltzer DO. The effect of comorbid illness and functional status on the expected benefits of intensive glucose control in older patients with type 2 diabetes: a decision analysis. *Ann Intern Med.* 2008;149(1):11-19.
- Bennett WL, Maruthur NM, Singh S, Segal JB, Wilson LM, Chatterjee R, et al. Comparative effectiveness and safety of medications for type 2 diabetes: an update including new drugs and 2-drug combinations. *Ann Intern Med.* 2011;154(9):602-613.
- Gerstein HC, Miller ME, Byington RP, et al. Effects of intensive glucose lowering in type 2 diabetes. N Engl J Med. 2008;358(24):2545-2559.
- Nauck MA, Meininger G, Sheng D, Terranella L, Stein PP. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor, sitagliptin, compared with the sulfonylurea, glipizide, in patients with type 2 diabetes inadequately controlled on metformin alone: a randomized, double-blind, noninferiority trial. *Diabetes Obes Metab.* 2007;9(2):194–205.
- 10. Micromedex® 2.0 [database online]. Ann Arbor, MI: Thomson Reuters; 2012. http://www. micromedex.com. Accessed March 18, 2012.
- Salpeter SR, Greyber E, Pasternak GA, Salpeter EE. Risk of fatal and nonfatal lactic acidosis with metformin use in type 2 diabetes mellitus. *Cochrane Database Syst Rev.* 2010;(4):CD002967. doi:0.1002/14651858.CD002967.pub4
- Rachmani R, Slavachevski I, Levi Z, Zadok B, Kedar Y, Ravid M. Metformin in patients with type 2 diabetes mellitus: reconsideration of traditional contraindications. *Eur J Intern Med.* 2002;13(7):428-433.
- 13. Bakris GL. Treatment of hypertension in patients with diabetes mellitus. In: UpToDate, Forman JP (Ed), UpToDate, Waltham, MA, 2012.
- 14. Blonde, L. Benefits and risks for intensive glycemic control in patients with diabetes mellitus. *Am J Med Sci.* 2012;343(1):17–20.

- Boussageon R, Bejan-Angoulvant T, Saadatian-Elahi M, et al. Effect of intensive glucose lowering treatment on all cause mortality, cardiovascular death, and microvascular events in type 2 diabetes: meta-analysis of randomised controlled trials. *BMJ.* 2011;343:d4169. doi:1136/bmj.d4169.
- Crossing the Quality Chasm: A New Health System for the 21st Century. Washington, DC: The National Academies Press, 2001. http://www.iom.edu/~/media/Files/Report%20Files/2001/ Crossing-the-Quality-Chasm/Quality%20Chasm%202001%20%20report%20brief.pdf Accessed April 3, 2012.
- 17. Mason A. New medicines in primary care: a review of influences on general practitioner prescribing. *J Clin Pharm Ther.* 2008;33(1):1-10.
- Boissel JP, Amsallem E, Cucherat M, Nony P, Haugh MC. Bridging the gap between therapeutic research results and physician prescribing decisions: knowledge transfer, a prerequisite to knowledge translation. *Eur J Clin Pharmacol.* 2004:60(9):609-616.
- 19. Gansevoort RT, Sluiter WJ, Hemmelder MH, de Zeeuw D, de Jong PE. Antiproteinuric effect of blood-pressure-lowering agents: a meta-analysis of comparative trials. *Nephrol Dial Transplant*. 1995;10(11):1963-1974.
- Forsetlund L, Bjørndal A, Rashidian A, et al. Continuing education meetings and workshops: effects on professional practice and health care outcomes. *Cochrane Database Syst Rev.* 2009;(2):CD003030. doi:0.1002/14651858.CD003030.pub2

About MedInsight

The Epocrates MedInsight Service offers access to one of the largest verified and opted-in panels of physicians and allied healthcare practitioners in the U.S. Using MedInsight's unique online tools, organizations conduct primary research with our specialist panel. Insights gleaned from physician surveys can assist in developing communication programs and interventions designed to improve quality of care and manage prescription drug utilization.

About Health Plan Services

The Epocrates Health Plan Services allows health plans to integrate their formulary information with trusted clinical information in the Epocrates drug references. Physicians and other prescribers can access multiple formularies on their mobile or desktop computer helping to maintain compliance and increasing provider satisfaction. In addition, Health Plan Services offers the ability to deliver clinical messaging, CME content, and interactive learning modules directly to physicians' mobile devices.

About Epocrates

Epocrates is a leading provider of mobile drug reference tools and interactive services to healthcare professionals and the healthcare industry. Epocrates' active user network currently has more than one million healthcare professionals, including more than 50 percent of U.S. physicians. Most commonly used on mobile devices at the point of care, the company's clinical products and services help healthcare professionals make more informed prescribing decisions, enhance patient safety and improve practice productivity.

Thanks to Randall Stafford, MD, PhD, Program Director, Program on Prevention Outcomes and Practices, Stanford Prevention Research Center, Stanford University School of Medicine and RxBalance for assistance with development of this survey and report.

Copyright © 2012, Epocrates Inc. All Rights Reserved. April 2012.