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Many Americans Have Pre-Diabetes and Should Be Considered for Metformin Therapy

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OBJECTIVE — To determine the proportion of the American population who would merit metformin treatment, according to recent American Diabetes Association (ADA) consensus panel recommendations to prevent or delay the development of diabetes.

RESEARCH DESIGN AND METHODS — Risk factors were evaluated in 1,581 Screening for Impaired Glucose Tolerance (SIGT), 2,014 Third National Health and Nutrition Examination Survey (NHANES III), and 1,111 National Health and Nutrition Examination Survey 2005–2006 (NHANES 2005–2006) subjects, who were non-Hispanic white and black, without known diabetes. Criteria for consideration of metformin included the presence of both impaired fasting glucose (IFG) and impaired glucose tolerance (IGT), with ≥1 additional diabetes risk factor: age <60 years, BMI ≥35 kg/m², family history of diabetes, elevated triglycerides, reduced HDL cholesterol, hypertension, or A1C >6.0%.

RESULTS — Isolated IFG, isolated IGT, and IFG and IGT were found in 18.0, 7.2, and 8.2% of SIGT, 22.3, 6.4, and 9.4% of NHANES III, and 21.8, 5.0, and 9.0% of NHANES 2005–2006 subjects, respectively. In SIGT, NHANES III, and NHANES 2005–2006, criteria for metformin consideration were met in 99, 96, and 96% of those with IFG and IGT; 31, 29, and 28% of all those with IFG; and 53, 57, and 62% of all those with IGT (8.1, 9.1, and 8.7% of all subjects), respectively.

CONCLUSIONS — More than 96% of individuals with both IFG and IGT are likely to meet ADA consensus criteria for consideration of metformin. Because ≥28% of all those with IFG met the criteria, providers should perform oral glucose tolerance tests to find concomitant IGT in all patients with IFG. To the extent that our findings are representative of the U.S. population, ~1 in 12 adults has a combination of pre-diabetes and risk factors that may justify consideration of metformin treatment for diabetes prevention.

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Diabetes is a public health epidemic (1) associated with high morbidity, mortality (1), and cost (2). Currently, an estimated 38 million Americans have the disease, nearly 40% of which is undiagnosed, and another 87 million have pre-diabetes: impaired fasting glucose (IFG) and/or impaired glucose tolerance (IGT) (3). Diabetes develops insidiously over several years, during which time glucose metabolism progresses slowly from normal to pre-diabetes and then more rapidly to diabetes. Based on observational and prospective studies, ~25–40% of individuals with pre-diabetes go on to develop diabetes over 3–8 years (4–6), and there is evidence of complications in 50% of patients at the time of diagnosis of diabetes (7).

Because progression from pre-diabetes can be prevented or delayed by lifestyle change and/or medication (4–6), the American Diabetes Association (ADA) has issued a consensus statement recommending early identification and preventive treatment in high-risk individuals (8). The panel statement recommends that individuals with both IFG and IGT and one additional risk factor (age <60 years, BMI ≥35 kg/m², family history of diabetes in first-degree relative, elevated triglycerides, reduced HDL cholesterol, or A1C >6.0%) should be considered for treatment with metformin, in addition to lifestyle modification, which includes weight loss and physical activity.

To determine what proportion of the American population presenting with either IFG or IGT would merit consideration for metformin treatment in accordance with the recent ADA recommendations, we evaluated healthy volunteers without known diabetes who were screened for diabetes/pre-diabetes by the 75-g oral glucose tolerance test (OGTT).

RESEARCH DESIGN AND METHODS — In cross-sectional analyses, we evaluated the likelihood that Americans with previously unrecognized pre-diabetes would meet ADA consensus panel recommendations for consideration of metformin in addition to change in lifestyle. Criteria for consideration of metformin included the presence of both IFG and IGT, with ≥1 additional diabetes risk factor: age <60 years, BMI ≥35 kg/m², family history of diabetes, elevated triglycerides, reduced HDL cholesterol, hypertension, or A1C > 6.0%.

Between 1 December 2005 and 31 March 2008, subjects were recruited to participate in the Screening for Impaired Glucose Tolerance (SIGT) study (9), a cross-sectional study that was approved...
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by the Emory Institutional Review Board. The invitation to participate was extended to employees of the Grady Health System, Emory HealthCare, and Emory University and Morehouse Schools of Medicine as well as to members of the community. Criteria for eligibility were age ≥18 years, non-Hispanic white or black race, no prior diagnosis of diabetes, not pregnant or breast-feeding, not taking glucocorticoids, and being well enough to have worked during the previous week (without requiring actual employment). During recruitment, 4,024 individuals expressed initial interest in the study, among whom 2,111 were scheduled for first visits (selected largely on the basis of need to balance participant sex and race), 1,658 completed first visits, and 1,581 completed the protocol. All study visits were performed in the General Clinical Research Centers at Emory University Hospital and Grady Memorial Hospital. All subjects gave written informed consent before study participation.

We also evaluated subjects who took part in the Third National Health and Nutrition Examination Survey (NHANES III) (10) and the continuous National Health and Nutrition Examination Survey 2005–2006 (NHANES 2005–2006) (11). NHANES is a program of studies conducted by the National Center for Health Statistics of the Centers for Disease Control and Prevention that include both interviews and physical examinations in a nationally representative sample to assess the health and nutritional status of adults aged 18 years, BMI >0 (8). Other risk factors for diabetes that were not specifically defined by the ADA were categorized according to the American Heart Association/National Heart, Lung, and Blood Institute (AHA/NHLBI) diagnostic criteria for metabolic syndrome (12): presence of hypertension by history, systolic blood pressure >130 mmHg or diastolic blood pressure >85 mmHg, triglyceride level ≥150 mg/dl, and HDL cholesterol <40 mg/dl in men and <50 mg/dl in women. Given the high number of subjects in NHANES III and NHANES 2005–2006 whose reporting of the diabetes status of one or more first-degree relatives was either not known or left blank (NHANES III, n = 1,163; NHANES 2005–2006, n = 116), relatives whose diabetes status was missing or not known were assumed to not have diabetes, a method that was also implemented for the analysis of the SIGT study group. In addition, subjects with missing values for the remaining risk factors were excluded from analysis (NHANES III: 1 missing blood pressure measurement or hypertension history, 7 missing A1C values, 27 missing triglyceride values, and 35 missing HDL cholesterol values; NHANES 2005–2006: 32 missing blood pressure measurements or hypertension history, 8 missing BMI measurements, 2 missing A1C values, 4 missing triglyceride values, and 4 missing HDL values), leaving 2,014 subjects in NHANES III and 1,111 subjects in NHANES 2005–2006 to be analyzed for metformin consideration.

Means and frequencies were determined in aggregate and by subgroup analysis of the different glucose tolerance categories. All SIGT analyses were performed using SPSS 15.0 (SPSS, Chicago, IL). All NHANES III and NHANES 2005–2006 analyses were conducted using SUDAAN statistical software (version 10).
RESULTS — Among 1,581 volunteers who completed OGTTs in the SIGT study, average age was 48 years and BMI was 30.3 kg/m², 42% were male, and 58% were black (Table 1). In the selected NHANES III population (n = 2,014), the average age was 55 years and BMI was 27.3 kg/m², 47% were male, and 10% were black, and in NHANES 2005–2006 (n = 1,111), the average age was 46 years and BMI was 28.5 kg/m², 49% were male, and 13% were black (Table 1).

In the SIGT population, 62.1% had normal fasting glucose and NGT, 18.0% had isolated IFG, 7.2% had isolated IGT, 8.2% had both IFG and IGT, and 4.6% had diabetes, similar to the proportions in NHANES III (54.3% had NGT, 22.3% had isolated IFG, 6.4% had isolated IGT, 9.4% had both IFG and IGT, and 7.6% had diabetes) and NHANES 2005–2006 (39.1% had NGT, 21.8% had isolated IFG, 5.0% had isolated IGT, 9.0% had both IFG and IGT, and 5.2% had diabetes). All three populations had a comparable portion with either IFG or IGT (33.4% in SIGT, 38.1% in NHANES III, and 35.8% in NHANES 2005–2006).

When the associated risk factors for diabetes, as specified by the ADA consensus statement (8), were considered, among those with both IFG and IGT, the presence of each risk factor was generally higher among SIGT subjects, compared with subjects in NHANES III and NHANES 2005–2006, with the exception of elevated triglycerides and A1C levels (Table 2). Even with the differences in the prevalence of risk factors, almost all subjects with both IFG and IGT in all three populations had at least one risk factor (99% in SIGT, 96% in NHANES III, and 96% in NHANES 2005–2006), which was similar among those with IFG (isolated or with IGT: 99% in SIGT, 96% in NHANES III, and 94% in NHANES 2005–2006). Among all subjects with IFG (isolated or with IGT), one-quarter to one-third (31% in SIGT, 29% in NHANES III, and 28% in NHANES 2005–2006) met the recommended criteria for metformin treatment, and among all subjects with IGT (isolated or with IFG), one-half to two-thirds (53% in SIGT, 57% in NHANES III, and 62% in NHANES 2005–2006) did so (Fig. 1). Overall, ~1 in 12 individuals in these populations met the criteria for consideration of metformin (8.1% in SIGT, 9.1% in NHANES III, and 8.7% in NHANES 2005–2006).

CONCLUSIONS — In consideration of the enormous public health impact of diabetes and the evidence of benefit from pharmacological treatment for the prevention of diabetes, the ADA issued a consensus statement recommending preventive treatment in individuals at high risk of developing diabetes, defined as those with more severe pre-diabetes.

Table 1—Characteristics of study subjects

<table>
<thead>
<tr>
<th>Category</th>
<th>SIGT</th>
<th>NHANES III</th>
<th>NHANES 2005–2006</th>
</tr>
</thead>
<tbody>
<tr>
<td>n</td>
<td>1,581</td>
<td>2,014 (weighted)</td>
<td>1,111 (weighted)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>48 ± 0.3</td>
<td>55 ± 0.5</td>
<td>46 ± 1.0</td>
</tr>
<tr>
<td>Male sex (%)</td>
<td>42 ± 0.01</td>
<td>47 ± 1.1</td>
<td>49 ± 1.7</td>
</tr>
<tr>
<td>Black (%)</td>
<td>58 ± 0.01</td>
<td>10 ± 0.8</td>
<td>13 ± 2.1</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>30 ± 0.2</td>
<td>27 ± 0.2</td>
<td>28 ± 0.2</td>
</tr>
<tr>
<td>A1C (%)</td>
<td>5.4 ± 0.01</td>
<td>5.4 ± 0.02</td>
<td>5.3 ± 0.02</td>
</tr>
</tbody>
</table>

Data are means ± SEM.

To account for the complex survey design, and all estimates were weighted (RTI International, Research Triangle Park, NC).

Table 2—Prevalence of risk factors for diabetes in study subjects

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>All subjects</td>
<td>1,581</td>
<td>2,014 (weighted)</td>
<td>1,111 (weighted)</td>
</tr>
<tr>
<td>Age &lt;60 years</td>
<td>84</td>
<td>79</td>
<td>76</td>
</tr>
<tr>
<td>BMI ≥ 35 kg/m²</td>
<td>22</td>
<td>27</td>
<td>30</td>
</tr>
<tr>
<td>Family history of diabetes</td>
<td>46</td>
<td>49</td>
<td>54</td>
</tr>
<tr>
<td>Triglycerides ≥ 150 mg/dl</td>
<td>13</td>
<td>18</td>
<td>21</td>
</tr>
<tr>
<td>Reduced HDL cholesterol*</td>
<td>47</td>
<td>54</td>
<td>57</td>
</tr>
<tr>
<td>Hypertension†</td>
<td>49</td>
<td>63</td>
<td>68</td>
</tr>
<tr>
<td>A1C &lt;6.0%</td>
<td>7</td>
<td>10</td>
<td>12</td>
</tr>
<tr>
<td>≥1 risk factor</td>
<td>99</td>
<td>99</td>
<td>99</td>
</tr>
<tr>
<td>Metformin indicated†</td>
<td>8.1</td>
<td>31</td>
<td>52.7</td>
</tr>
</tbody>
</table>

Data are %. Glucose tolerance categories: IFG (isolated or with IGT), IGT (isolated or with IFG), IGT with or without IFG, IFG + IGT, both IFG and IGT. *Reduced HDL cholesterol defined as ≤40 mg/dl in men and ≤50 mg/dl in women. †Hypertension defined by any of the following: history of hypertension, systolic blood pressure ≥130 mmHg, or diastolic blood pressure ≥85 mmHg. ‡Metformin indicated per the ADA consensus statement (8) criteria of the presence of both IFG and IGT and one of the following diabetes risk factors: age <60 years, BMI ≥35 kg/m², family history of diabetes, elevated triglycerides, reduced HDL cholesterol, and A1C >6.0%. Risk factors for diabetes that were not specifically defined by the ADA were categorized according to the AHA/NHLBI diagnostic criteria for metabolic syndrome (12): presence of hypertension by history, systolic blood pressure >130 mmHg or diastolic blood pressure >85 mmHg, triglyceride level ≥150 mg/dl, and HDL cholesterol <40 mg/dl in men and <50 mg/dl in women.
Metformin indicated in many pre-diabetic patients

Figure 1—Prevalence of metformin indication, stratified by glucose tolerance category. Metformin is indicated per the ADA consensus statement criteria of the presence of both IFG and IGT and one of the following diabetes risk factors: age <60 years, BMI ≥35 kg/m², family history of diabetes, elevated triglycerides, reduced HDL cholesterol, and A1C >6.0% (8). Risk factors for diabetes that were not specifically defined by the ADA were categorized according to the AHA/NHLBI diagnostic criteria for metabolic syndrome (12): presence of hypertension by history, systolic blood pressure >130 mmHg or diastolic blood pressure >85 mmHg, triglyceride level ≥150 mg/dl, and HDL cholesterol <40 mg/dl in men and <50 mg/dl in women. Glucose tolerance categories are as follows: IFG 100–109, FPG levels 100–109 mg/dl and 2-h postchallenge plasma glucose <140 mg/dl; IFG 110–125, FPG 110–125 mg/dl and 2-h postchallenge plasma glucose <140 mg/dl; all IFG, isolated IFG (FPG 100–125 mg/dl and 2-h postchallenge plasma glucose <140 mg/dl); IGT, isolated IGT; and IFG 100–125 + IGT, all IFG and IGT.

Diabetes is currently the leading cause of blindness, end-stage renal disease requiring dialysis, and nontraumatic amputations in the U.S. and increases the risk for cardiovascular disease and stroke by two- to fourfold, compared with those without diabetes (1). It is the seventh leading cause of death (1) and in 2007 cost $174 billion in both direct and indirect health care expenditures (2). In addition, the prevalence of diabetes has been on the rise in the adolescent population (13), indicating that the epidemic is likely to continue into the next generation.

Pre-diabetes, the stage preceding the development of diabetes, increases the risk for the development of diabetes, such that 25–39% of patients with IFG or IGT go on to develop diabetes over a period of 5–10 years (14,15). Moreover, pre-diabetes alone has been associated with an increased risk for the development of cardiovascular disease (16,17) and microvascular complications typically seen with diabetes (18). Given these risks, prospective studies have been conducted to identify preventive treatment. In addition to lifestyle modification, pharmacological treatment with acarbose (5), rosiglitazone (6), orlistat (19), or metformin (4) has shown efficacy in preventing or delaying the onset of diabetes in individuals with pre-diabetes. The relative risk reduction for diabetes in the pre-diabetic population was 25% over 3.3 years in patients treated with acarbose (5), 52–62% over 2–4 years with orlistat (19), 62% over 3 years with rosiglitazone (6), and 26–31% over 2.5–2.8 years with metformin (4). However, because many individuals with pre-diabetes are generally healthy, the benefit of preventive treatment must outweigh any associated side effects or additional risks, particularly because none of these medications have U.S. Food and Drug Administration approval for the indication of diabetes prevention. Gastrointestinal side effects are commonly associated with acarbose (5) and orlistat (19), leading to poor patient compliance, whereas an increased risk of bone loss

(both IFG and IGT as well as an additional risk factor) (8). To determine the proportion of individuals who would be targeted by such a recommendation, we examined a relatively healthy population without previously diagnosed diabetes (SIGT) and representative samples of the U.S. population (NHANES III and NHANES 2005–2006) and found that one-quarter to one-third had pre-diabetes. Among those with IFG, nearly one-third of subjects met the criteria for consideration of metformin treatment to prevent diabetes in accordance with the recent ADA consensus statement, more than one-half of all of the subjects with IGT qualified, and almost all of those with both IFG and IGT qualified. Overall, 8–9% met the recommended criteria. Assuming that our data are generalizable to the U.S. population, ~24 million Americans might benefit from pharmacological treatment in addition to lifestyle modification.

The epidemic of diabetes and the insidious onset of its complications have prompted a call for early identification and preventive treatment of the disease. Diabetes care.diabetesjournals.org
(20), worsening or new-onset edema (21), and heart failure (22) are associated with rosiglitazone. Therefore, metformin, which has been used for many years and is both generally well tolerated and relatively safe, has become the leading candidate for preventive treatment. In addition to the recommendations of the ADA, the American College of Endocrinology (ACE) has recently issued their consensus statement on the management of pre-diabetes (23). Similar to the ADA recommendations, the ACE statement recognizes the need for preventive treatment, beginning with lifestyle modification, but also emphasizes the importance of treating relevant comorbid conditions, such as hypertension, hypercholesterolemia, and obesity, and provides a looser set of criteria regarding the initiation of pharmacological treatment. Acarbose and metformin are recommended treatments for individuals who are at high risk of developing diabetes, which include, but are not limited to, those with IFG, IGT, and/or the metabolic syndrome, worsening glycermia, cardiovascular disease, nonalcoholic fatty liver disease, a history of gestational diabetes, or polycystic ovary syndrome. Taking into account the target populations as defined by the ADA and the ACE, >8% of Americans could benefit from pharmacological treatment to prevent or delay development of diabetes.

Use of pharmacological agents for the many Americans who may benefit from preventive treatment would incur substantial costs: at current generic rates for metformin, possibly $4/month × 12 months × 24 million Americans = $1.15 billion per year. However, several studies suggest that diabetes prevention or delay with metformin is likely to be cost-effective and/or cost-saving (24); further evaluation using a variety of cost analysis methods may be required to reach a definitive conclusion regarding the cost of preventive treatment.

To our knowledge, our findings are the first evaluation of the proportion of relatively healthy individuals who might benefit from metformin treatment for the prevention or delay of development of diabetes. However, our study has limitations. Because all SIGT subjects were recruited on a volunteer basis, there may have been a selection bias toward higher family history of diabetes and/or other risk factors for diabetes. Therefore, the SIGT population may represent a group of individuals at higher risk. However, because many SIGT subjects were recruited from university and health care settings, they may also follow healthier lifestyles, which could offset such a bias. Moreover, the proportion with diabetes or prediabetes in SIGT was no higher than that in NHANES III and was comparable to that in the more recent NHANES 2005–2006, both of which represent randomized, stratified samples of the American population.

The morbidity, mortality, and cost of the epidemic of diabetes have prompted a call for primary prevention of diabetes in high-risk individuals by the use of metformin in addition to lifestyle changes. To the extent that our findings are representative of the U.S. population, close to 1 in 12 American adults may meet the recommended guidelines for consideration of metformin treatment for diabetes prevention or delay. Notably, eligibility for metformin use appeared to be almost completely determined by impaired glucose metabolism alone, because 99% of the SIGT population and 96% of the NHANES populations with both IFG and IGT had at least one risk factor. Therefore, once the presence of both IFG and IGT has been established, the presence of additional risk factors could almost be assumed, and initiation of metformin should be considered. Moreover, because nearly one-third of all subjects with IFG met the criteria for metformin treatment, providers should perform OGTTs in all patients with IFG to test for the presence of IGT (or unrecognized diabetes) and thereby determine whether they merit consideration of metformin treatment.

References
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