# Diabetes Risk in a Cuban Primary Care Setting in Persons with No Known Glucose Abnormalities

Adrian A. Naranjo, Ángel Y. Rodríguez MD, Rosa E. Llera MD MS, Ronald Aroche MD

## ABSTRACT

**INTRODUCTION** With 333 million cases worldwide predicted for 2015, type 2 diabetes mellitus presents an important global health challenge. Its rising tide calls for health policies emphasizing prevention at the primary care level, including public education as well as early risk identification and intervention.

**OBJECTIVES** Estimate risk of developing type 2 diabetes in persons with no known glucose abnormalities, registered in a primary care setting in Pinar del Río city, Cuba, using FINDRISK.

**METHODS** A descriptive, cross-sectional study applied FINDRISK to 620 persons aged ≥18 years randomly selected from a universe of 1058 patients with no known glucose abnormalities, registered in family-doctor-and-nurse office No. 23 in the Turcios Lima Teaching Polyclinic health area, Pinar del Río city.

## **INTRODUCTION**

Diabetes mellitus (DM) has become one of the most important global health problems, with accelerated growth in numbers of people affected. Each year, there are seven million new cases and every ten seconds a patient dies from DM. An increase to 333 million cases by 2025 is anticipated, which would involve 6.5% of the world's population.[1]

In Cuba, DM is also a major public health problem, affecting quality of life for diabetics and their families, as well as increasing the burden of direct and indirect health costs to them, the public health system and the national economy. In 2010 there were 22,000 DM hospitalizations. In 2011 DM prevalence was 47.7 per 1000 population, and DM was the eighth cause of death, with a mortality rate of 11.5 per 100,000 population; it was also implicated in the first and third causes of death.[2,3]

Preventing type 1 DM remains a goal for the future, but the possibility of preventing type 2 DM has been demonstrated. It requires action based on sustained lifestyle changes involving diet and physical activity, as well as identification of population groups at greater risk, in order to implement health policies that create environments conducive to achieving these changes.[4-10]

Several tools have been described for predicting DM in individuals with no known glucose metabolism disorder, most based on clinical and anthropometric variables and biochemical measurements.[11–24] However, we have not found any application of these tools or predictive studies in the Cuban medical literature.

The Finnish Diabetes Risk Score (FINDRISK)[25] was the first lifestyle and clinical parameter predictive scale to identify individuals at risk for type 2 DM. It covers eight aspects: age, body mass index (BMI), abdominal circumference, physical activity, diet, antihypertensive drug use, personal history of high blood glucose and family history of DM. It was developed in 1987 by Lindström and Tuomilehto for a population sample in Finland comprised of per**RESULTS** The study population was predominantly aged ≤45 years (53.5%) and 80.2% was overweight or obese. At least moderate risk of diabetes was found in 74.4% of the sample, and 10.5% was at very high risk, meaning an estimated 120 patients in the sample could be expected to develop type 2 diabetes within the next 10 years.

**CONCLUSIONS** Type 2 diabetes prevalence can be expected to increase substantially in this population over the next decade. We recommend design and timely implementation of intensive lifestyle change programs to eliminate or slow development of type 2 diabetes in at-risk individuals. We propose following cohorts identified by FIND-RISK to assess its prognostic value in the Cuban population.

**KEYWORDS** Diabetes mellitus, risk factors, risk prediction, prevention, Cuba

sons who answered a questionnaire and were followed for ten years. The scale was found useful for predicting type 2 DM and it has been the one most widely disseminated and used for this purpose internationally.[25]

In the population served by family doctor-and-nurse office (CMF, the Spanish acronym) No. 23 of the Turcios Lima Teaching Polyclinic health area in Pinar del Río city, the assumption was made of the existence of a group of individuals unknowingly at high risk for developing type 2 DM; identifying them was necessary to map strategies to prevent or slow disease onset. Hence, our decision to conduct this study to answer the question: can we detect such individuals, who have no known glucose abnormalities, using FINDRISK,[25] a scale designed for a different population?

#### **METHODS**

A descriptive, cross-sectional study was done of the population served by CMF No. 23 of the Turcios Lima Teaching Polyclinic health area in Pinar del Río city in 2010, to estimate risk of type 2 DM in individuals with no known glucose abnormalities.

**Population** The study universe consisted of 1058 individuals aged  $\geq$ 18 years who received primary health care at CMF No. 23 and had no personal history of any type of DM or other known carbohydrate metabolism disorder. The final sample consisted of 620 individuals randomly selected using a random numbers table, following sample size determination based on estimates of risk distribution on the FINDRISK scale from a pilot sample of 24 individuals.[25] We estimated that a sample size of 620 would be needed to obtain a confidence interval for the population mean at a significance level of  $\alpha = 0.01$  (at 99%).

**Data collection** Patient data were gathered using a survey modeled on the original FINDRISK[25] score; when necessary, medical records were consulted and updated during followup. Data from both sources were entered into an information collec-

tion form designed for the study, which contained the variables described in Table 1.

**Statistical analysis** A Microsoft Office Access database was created and later exported to SPSS 11.5 statistical software for Windows. Variables were described with absolute and relative frequency distributions.

**Ethical aspects** The project was approved by the medical ethics committee of Turcios Lima Teaching Polyclinic. Individuals' identities were protected and they provided written informed consent prior to inclusion in the study.

## RESULTS

In the study sample, 53.5% (332/620) of participants were aged  $\leq$ 45 years and the number of individuals in each age group decreased with increasing age (Table 2). The study group was predominantly female (59.7%, 370/620).

The majority of the sample, 80.2% (497/620), were overweight or obese, and 61% (226/370) of women and 74.8% (187/250) of men were in the abdominal circumference categories for slight or established visceral obesity (Table 2).

It was found that 33.7% (209/620) exercised and 13.9% (86/620) ate vegetables daily. Some 34.7% (215/620) of those evaluated were taking antihypertensives. Additionally, 17.7% (110/620) had a diabetic relative; 6.6% (41/620) a first-degree relative. Finally, 16.8% (104/620) reported a personal history of hyperglycemia (Table 2).

Table 2: Distribution of FINDRISK<sup>a</sup> variables in CMF No. 23, Turcios Lima Teaching Polyclinic, Pinar del Río, Cuba (n = 620)

Variable	Category	No. (%)		
Age (years)	18–45 46–54 55–64 ≥65	332 (53.5) 209 (33.7) 64 (10.3) 15 (2.4)		
Sex	Male Female	250 (40.3) 370 (59.7)		
BMI	Low or normal weight Overweight Obese	123 (19.8) 317 (51.2) 180 (29.0)		
Abdominal circumference		Female n = 370	Male n = 250	
	No visceral obesity Slight visceral obesity Established visceral obesity	144 (38.9) 113 (30.5) 113 (30.5)	· · · ·	
Daily physical activity <sup>b</sup>	Yes No		209 (33.7) 411 (66.3)	
Daily vegetable consumption <sup>c</sup>	Yes No	86 (13.9) 534 (86.1)		
Antihypertensive use	Yes No	215 (34.7) 405 (65.3)		
History of high blood glucose	Yes No		104 (16.8) 516 (83.2)	
Family history of diabetes	Non first-degree relative First-degree relative		69 (11.1) 41 (6.6)	

CMF: family doctor-and-nurse office

<sup>a</sup> FINDRISK[25] <sup>b</sup> Exercise at least 30 minutes a day, including physical work and calorie-expending activities

° Quantity not considered

Table 3 shows the distribution of FIND-RISK risk levels in the sample, probabilities for each risk level of developing type 2 DM in the next ten years, and the number of resulting new cases of type 2 DM.

Over the next decade, if no risk reduction action is taken, 120 new DM cases can be expected in the sample and 202 in the total CMF No. 23 population, assuming the same underlying risk distribution in both.

## DISCUSSION

The fight against DM, with its high prevalence and both short- and long-term complications, can be won through prevention and early diagnosis.[26–28]

Persons with elevated DM risk should be targeted for education.[2,6] Education of diabetics is essential to optimize metabolic control and prevent appearance and progression of complications.

Like other population screening studies,[29] in this one we chose FINDRISK because of the ease of obtaining the required clinical, anthropometric and hematologic parameters. However, it

#### Table 1: Study variables

Variableª	Description			
	18–45			
Age group (years)	46–54			
	55–64			
	≥65			
Sex	Male/Female			
Body mass index	Low or normal weight	<25.0		
(BMI in kg/m <sup>2</sup> )	Overweight	25.0–29.9		
	Obese	≥30.0		
		No visceral	Slight visceral	Established
Abdominal circumference (cm)		obesity	obesity	visceral obesity
Abdominal circumerence (cm)	Female	<80	80–88	>88
	Male	<94	94–102	>102
Daily physical activity <sup>b</sup>	Yes/No			
Daily vegetable consumption °	Yes/No			
Antihypertensive use	Yes/No			
History of high blood glucose	Yes/No			
	No			
Family history of diabetes	Yes; not first degree			
	Yes; first degree			
	Low	<7		
FINDRISK Score	Slightly elevated	7–11		
	Moderate	12–14		
	High	15–20		

<sup>a</sup> Grouped according to FINDRISK[25]

<sup>b</sup> Physical exercise at least 30 minutes a day including physical work and calorie-expending activities <sup>c</sup>Quantity not considered Table 3: Estimated ten-year increase in diabetics in study population per FINDRISK

Individual risk	No. (%)	Estimated probability of developing diabetes	No. in sample (n = 620)	No. in population* (n = 1058)
Low	94 (15.1)	1/100	1	2
Slightly elevated	65 (10.5)	1/25	3	4
Moderate	295 (47.6)	1/6	49	84
High	101 (16.3)	1/3	34	57
Very high	65 (10.5)	1/2	33	55
Total	620 (100.0)		120	202

\*Estimated total number CMF No. 23 patients likely to develop diabetes, assuming same age distribution and estimated probabilities

would have been ideal to develop a risk scoring adapted to the Cuban population, since methods developed in one country cannot always be applied successfully in another.[25,30]

In multivariate analysis including all parameters of the original FINDRISK[25] cohort (n = 4746), an age >45 years increased DM risk.[25] In our study, the majority of patients were aged  $\leq$ 45 years, and therefore had less age-associated risk.

Another statistically significant predictor of DM in the FIND-RISK[25] validation was male sex (OR 1.58, 95% CI 1.15–2.18). Inclusion or exclusion of sex in predictive models can influence the predictive coefficient of other variables, so some authors have preferred to develop sex-specific risk scores.[24,31]

In some studies assessing DM risk, such as that of Klein,male sex predominated in the sample;[32] female sex predominated in our sample, as in Rahman's validation of the Cambridge Diabetes Risk Score.[33] This is congruent with the greater DM prevalence in Cuba's female population reported in the 2010 statistical yearbook.[34]

Most persons in our sample were classified as overweight by BMI. In Lindström and Tuomilehto's multivariate analysis for FIN-DRISK, overweight was not a statistically significant predictor, [25] but they included it in the final version of the score because it was obvious to them that overweight was an intermediate stage between normal weight and obesity, the latter associated with greater DM risk. Overweight individuals should also be included in lifestyle change efforts, since DM onset is almost unavoidable once they become obese.[25]

The FINDRISK validation report indicates that increased abdominal circumference in both sexes significantly increases DM risk. [25] This is not surprising, since it correlates well with intra-abdominal and retroperitoneal fat. Intra-abdominal fat is metabolically very active and therefore its basal state is one of constant lipolysis, releasing free fatty acids into portal circulation. These free fatty acids travel to the liver and induce insulin resistance, leading to increased hepatic glucose production.[35]

The FINDRISK model also includes physical activity and fruit and vegetable consumption, although they contributed little to its predictive power.[25] They included the two variables to emphasize the importance of physical activity and diet in DM prevention. The fact that the bulk of patients in our sample did not exercise regularly or eat vegetables daily underscores the importance of addressing these factors in prevention programs.

FINDRISK included antihypertensive use[25] because it is an unequivocal marker of clinically evident hypertension and can be determined without the need to measure blood pressure. Antihypertensive use doubled DM risk in the Finnish population,[25] although not in the German population of the KORA study.[30] Our finding of high prevalence (34.7%) of antihypertensive use in the sample is cause for concern.

DM prognostic scales have been used not only to estimate DM risk, but also in diagnosis of established glucose abnormalities.[36] In the Finnish study,[25] half the individuals who reported hyperglycemia history were corroborated as diabetic. Thus, the 104 patients in our study who reported previously elevated blood glucose levels should be given new fasting glucose and oral glucose tolerance tests, to confirm or rule out DM or pre-DM.

A small but important proportion of the sample reported a firstdegree family history of DM, noteworthy given the multifactoral nature of DM etiology involving both genetic and environmental factors.[1,2,27] In the KORA study, family history increased tenyear risk of DM by 1.5 to 2 times, depending on the score used. [30] In a British population using the QDScore,first-degree family history of DM increased risk by 2.3 times in women and 2.7 times in men.[37]

The distribution of risk observed in our sample suggests a substantial increase in type 2 DM prevalence over the next ten years if we do not take effective preventive measures. In the Finnish cohort where the score was first used, overall 10-year increase in DM prevalence was 4.1%.[25]

Prevention is the most reasonable and effective way to avoid the dramatic consequences of DM, and our results confirm the importance of our main recommendation: design and implementation of intensive lifestyle change programs to mitigate the course of the current DM epidemic.

The use of risk scales such as FINDRISK enables us to define the population most at risk of DM, and, as result, to intervene in a timely fashion to eliminate or slow its onset. Efforts to that end include building awareness among primary care physicians, as well as developing public policies for prevention and public education starting at an early age.

The cross-sectional design of the study constitutes a limitation; hence our second recommendation, ten-year followup of the cohort to validate our risk predictions.

# CONCLUSIONS

DM prevalence in our CMF population is expected to increase over the coming decade. Hence the need for timely, intensive lifestyle change programs to eliminate or slow the appearance of type 2 DM in at-risk individuals. We propose following cohorts identified in this study by FINDRISK scoring to determine its predictive value for the Cuban population.

# REFERENCES

- Rydén L, Standl E, Bartnik M, Van der Berghe G, Betteridge J, de Boer MJ, et al. Guidelines on diabetes, pre-diabetes and cardiovascular diseases: executive summary. The Task Force on Diabetes and Cardiovascular Diseases of the European Society of Cardiology (ESC) and of the European Association for the Study of Diabetes (EASD). Eur Heart J. 2007 Jan;28(1):88–136.
- National Health Statistics and Medical Records Division (CU). Anuario Estadístico de Salud 2011. Havana: Ministry of Public Health (CU); 2012. Spanish.
- National Health Statistics and Medical Records Division (CU). Indicadores básicos de salud en Cuba 2011 [Internet]. Havana: Ministry of Public Health (CU); 2012 [cited 2012 Nov 10]. Available from: http://www.sld.cu/sitios/dne/. Spanish.
- Talmud PJ, Hingorani AD, Cooper JA, Marmot MG, Brunner EJ, Kumari M, et al. Utility of genetic and non-genetic risk factors in prediction of type 2 diabetes: Whitehall II prospective cohort study. BMJ. 2010 Jan 1;340:b4838.
- Bell RA, Mayer-Davis EJ, Beyer JW, D'Agostino RB Jr, Lawrence JM, Linder B, et al. Diabetes in non-Hispanic white youth: prevalence, incidence, and clinical characteristics: the SEARCH for Diabetes in Youth Study. Diabetes Care. 2009 Mar;32 Suppl 2:S102–11.
- 6. The Diabetes Prevention Program Research Group. The 10-year cost-effectiveness of lifestyle intervention or metformin for diabetes prevention: an intent-to-treat analysis of the DPP/DPPOS. Diabetes Care. 2012 Apr;35(4):723–30.
- Fonseca VA, Kirkman MS, Darsow T, Ratner RE. The American Diabetes Association diabetes research perspective. Diabetes Care. 2012 Jun;35(6):1380–7.
- Ackermann RT, Finch EA, Brizendine E, Zhou H, Marrero DG. Translating the Diabetes Prevention Program into the community. The DEPLOY Pilot Study. Am J Prev Med. 2008 Oct;35(4):357–63.
- DePalma MT, Rollison J, Camporese M. Psychosocial predictors of diabetes management. Am J Health Behav. 2011Mar–Apr;35(2):209–18.
- Boyle JP, Thompson TJ, Gregg EW, Barker LE, Williamson DF. Projection of the year 2050 burden of diabetes in the US adult population: dynamic modeling of incidence, mortality, and prediabetes prevalence. Popul Health Metr. 2010 Oct 22;8:29.
- Heikes KE, Eddy DM, Arondekar B, Schlessinger L. Diabetes risk calculator: a simple tool for detecting undiagnosed diabetes and pre-diabetes. Diabetes Care. 2008 May;31(5):1040–5.
- Aekplakorn W, Bunnag P, Woodward M, Sritara P, Cheepudomwit S, Yamwong S, et al. A risk score for predicting incident diabetes in the Thai population. Diabetes Care. 2006 Aug;29(8):1872–7.
- Stern MP, Williams K, Haffner SM. Identification of persons at high risk for type 2 diabetes mellitus: do we need the oral glucose tolerance test? Ann Intern Med. 2002 Apr 16;136(8):575–81.
- Glümer C, Carstensen B, Sandbaek A, Lauritzen T, Jørgensen T, Borch-Johnsen K; inter99 study. A Danish diabetes risk score for targeted screening: the Inter99 study. Diabetes Care. 2004 Mar;27(3):727–33.
- Balkau B, Lange C, Fezeu L, Tichet J, de Lauzon-Guillain B, Cezernichow S, et al. Predicting diabetes: clinical, biological, and genetic approaches. Diabetes Care. 2008 Oct;31(10):2056–61.

- Cabrera de León A, Coello SD, Rodríguez Pérez MC, Medina MB, Almeida González D, Díaz BB, et al. A simple clinical score for type 2 diabetes mellitus screening in the Canary Islands. Diabetes Res Clin Pract. 2008 Apr;80(1):128–33.
- Vivek CKS, Prabhakaran RD, Jeemon P, Ramakrishnan L, Shah P, Shah B. Development of a clinical risk score in predicting undiagnosed diabetes in urban Asian Indian adults: a populationbased study. CVD Prevention and Control. 2008 Sep;3(3):141–51.
- Glümer C, Vistisen D, Borch-Johnsen K, Colagiuri S; DETECT-2 Collaboration. Risk scores for type 2 diabetes can be applied in some populations but not all. Diabetes Care. 2006 Feb;29(2):410–4.
- Ramachandran A, Snehalatha C, Vijay V, Wareham NJ, Colagiuri S. Derivation and validation of diabetes risk score for urban Asian Indians. Diabetes Res Clin Pract. 2005 Oct;70(1):63–70.
- Mohan V, Deepa R, Deepa M, Somannavar S, Datta M. A simplified Indian Diabetes Risk Score for screening for undiagnosed diabetic subjects. J Assoc Physicians India. 2005 Sep;53:759–63.
- 21. Al-Lawati JA, Tuomilehto J. Diabetes risk score in Oman: a tool to identify prevalent type 2 diabetes among Arabs of the Middle East. Diabetes Res Clin Pract. 2007 Sep;77(3):438–44.
- Xie J, Hu D, Yu D, Chen CS, He J, Gu D. A quick self-assessment tool to identify individuals at high risk of type 2 diabetes in the Chinese general population. J Epidemiol Community Health. 2010 Mar;64(3):236–42.
- Waugh N, Scotland G, McNamee P, Gillett M, Brennan A, Goyder E, et al. A Screening for type 2 diabetes: literature review and economic modelling. Health Technol Assess. 2007;11:1–125.
- Balkau B, Lange C, Fezeu L, Tichet J, de Lauzon-Guillain B, Czernichow S, et al. Predicting diabetes: clinical, biological, and genetic approaches: data from the Epidemiological Study on the Insulin Resistance Syndrome (DESIR). Diabetes Care. 2008 Oct;31(10):2056–61.
- Lindström J, Tuomilehto J. The diabetes risk score: a practical tool to predict type 2 diabetes risk. Diabetes Care. 2003 Mar;26(3):725–31.
- García R, Suárez R. La educación a personas con diabetes mellitus en la atención primaria de salud. Rev Cubana Endocrinol [Internet] 2007 [cited 2012 Nov 10];18(1). Available from: http:// bvs.sld.cu/revistas/end/vol18\_1\_07/end05107 .htm. Spanish.
- American Diabetes Association. Diagnosis and classification of diabetes mellitus. Diabetes Care. 2012;35 (Suppl 1):S64–71.
- Vermunt PW, Milder IE, Wielaard F, van Oers JA, Westert GP. An active strategy to identify individuals eligible for type 2 diabetes prevention by lifestyle intervention in Dutch primary care: the APHRODITE study. Fam Pract. 2010 Jun;27(3):312–9.
- Saaristo T, Peltonen M, Lindström J, Saarikoski L, Sundvall J, Eriksson JG, et al. Crosssectional evaluation of the Finnish Diabetes Risk Score: a tool to identify undetected type 2 diabetes, abnormal glucose tolerance and metabolic syndrome. Diab Vasc Dis Res. 2005 May;2(2):67–72.
- Rathmann W, Martin S, Haastert B, Icks A, Holle R, Löwel H, et al. Performance of screening questionnaires and risk scores for undiagnosed

diabetes: the KORA Survey 2000. Arch Intern Med. 2005 Feb 28;165(4):436-41.

- Waugh N, Scotland G, McNamee P, Gillett M, Brennan A, Goyder E, et al. A Screening for type 2 diabetes: literature review and economic modelling. Health Technol Assess. 2007 May;11(17):1–125.
- Klein Woothuis EP, de Grauw WJ, van Gerwen WH, van den Hoogen HJ, van de Lisdonk EH, Metsemakers JF, et al. Identifying people at risk for undiagnosed type 2 diabetes using the GP's electronic medical record. Fam Pract. 2007 Jun;24(3):230–6.
- Rahman M, Simmons RK, Harding AH, Wareham NJ, Griffin SJ. A simple risk score identifies individuals at high risk of developing type 2 diabetes: a prospective cohort study. Fam Pract. 2008 Jun;25(3):191–6.
- National Health Statistics and Medical Records Division (CU). Anuario Estadístico de Salud 2010. Havana: Ministry of Public Health (CU); 2011. Spanish.
- Bonné Moreno V, González Löwenberg O, Charques Velasco E, Alonso Martínez MM. Riesgo coronario y prescripción en pacientes con hipercolesterolemia en atención primaria. Aten Primaria. 2000 Mar 15;25(4):209–13. Spanish.
- 36. Franciosi M, De Berardis G, Rossi MC, Sacco M, Belfiglio M, Pellegrini F, et al. Use of the diabetes risk score for opportunistic screening of undiagnosed diabetes and impaired glucose tolerance: the IGLOO (Impaired Glucose Tolerance and Long-Term Outcomes Observational) study. Diabetes Care. 2005 May;28(5):1187–94.
- Hippisley-Cox J, Coupland C, Robson J, Sheikh A, Brindle P. Predicting risk of type 2 diabetes in England and Wales: prospective derivation and validation of QDScore. BMJ. 2009 Mar 17;338:b880.

#### **THE AUTHORS**

Adrian A. Naranjo Domínguez (Corresponding author: adrian90@princesa.pri.sld.cu), fourthyear medical student at the Medical University of Pinar del Río, posted at the Turcios Lima Teaching Polyclinic, Pinar del Río, Cuba.

**Ángel Y. Rodríguez Navarro,** family medicine resident, Turcios Lima Teaching Polyclinic, Pinar del Río, Cuba.

**Rosa E. Llera Almenteros,** physiologist with a master's degree in education. Associate professor, Medical University of Pinar del Río, Cuba.

**Ronald Aroche Aportela**, family physician and cardiologist. Instructor, Medical University of Havana and Medical University of Pinar del Río, Cuba.

Submitted: November 13, 2012 Approved for publication: April 20, 2013 Disclosures: None