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Evaluating the performance of the Indian Diabetes Risk Score in different ethnic groups

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Novelty Statement:

- Indian Diabetes Risk Score (IDRS) is a non-invasive diabetes risk score that uses only four variables and it has been identified as a cost-effective tool for identifying undiagnosed diabetes among Indians
- This study has shown that IDRS Score has good performance not only among Indians but also in other ethnic groups including Hispanic, Non-Hispanic White, Non-Hispanic Black and other American
- IDRS can be used in mass screening programmes by non-medically trained health care workers and as a self-administered tool among public to help in early diagnosis, management and optimal control of diabetes in India and America

Keywords: Diabetes Risk Scores, Undiagnosed Diabetes, Indian Diabetes Risk Score

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Abstract

Aim

To evaluate the performance of Madras Diabetes Research Foundation -Indian Diabetes Risk Score (MDRF-IDRS score) in different ethnic groups including Indians, Hispanic, Non-Hispanic Whites, Non-Hispanic Blacks and other American.

Methods

The MDRF-IDRS score is calculated based on a risk equation that includes age, waist circumference, family history of diabetes and physical activity. The National Health and Nutrition Examination Survey (NHANES) data on American and Chennai Urban Rural Epidemiology Study data on Indians were used in this study. Study participants aged ≥ 20 years with and without type 2 diabetes were included. Performance of the MDRF-IDRS score was assessed using sensitivity, specificity, positive predictive value, negative predictive value and the area under the receiver operating characteristic curve measures within each ethnic group. IDRS scores' performance was then compared with existing noninvasive American diabetes risk scores.

Results

Total number of participants included was 11,035 (2292 Indians and 8743 American). MDRF-IDRS score (cut off≥ 60) performed well in Indians with an AUC, sensitivity and specificity of 0.73, 80.2% and 57.3% respectively. MDRF-IDRS score cut off ≥ 70 had the highest discriminative performance among Hispanic, Non-Hispanic Whites and Non-Hispanic Blacks with sensitivity and specificity of between 70.1-86.9% and 61.2-72.2% respectively. The AUC for American was between 0.77-0.81 with the highest and lowest AUC in Non-Hispanic Black and Non-Hispanic White respectively. With a smaller number of variables, IDRS score showed almost the same performance in predicting diabetes among American compared with the existing non-invasive American diabetes risk score.

Conclusion

The MDRF-IDRS score performs well among Indians and American including Hispanic, Non-Hispanic White, Non-Hispanic Black and other American. It can be used as a screening tool Downloaded by UNIVERSITY COLLEGE LONDON from www.liebertpub.com at 12/27/19. For personal use only.

Diabetes Technology and Therapeutics

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to help in early diagnosis, management and optimal control of diabetes mainly in mass screening programmes in India and America.

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Introduction

Diabetes is a major public health burden in many countries around the world. The number of people with diabetes has risen rapidly over the last few decades and currently, there are around 425 million people with diabetes globally.^{1 2} In addition to those who have been diagnosed with diabetes, there are a large number of people with undiagnosed diabetes.³ The prevalence of undiagnosed diabetes is increasing not only in low-and-middle income countries but also in developed countries and it can be costly and harmful for the individual as well as for communities and health care systems.⁴ Early detection of people with undiagnosed diabetes is important as it enables them to manage their level of diabetes early on and to have a better quality of life by preventing or delaying the serious life threatening complications of diabetes such as heart disease, stroke, blindness, kidney disease and amputation.⁵⁻⁷

The World Health Organization and other health care organizations such as Centres for Disease Control and Prevention and the National Institute for Health and Clinical Excellence (NICE) recommend several different screening tools to identify people with diabetes. These include risk scoring tools and biochemical tests such as urine glucose, random blood glucose, fasting plasma glucose, glycated haemoglobin (HbA1c) and 75-g oral glucose tolerance test (OGTT). However, it is difficult to carry out biochemical tests in mass screening programmes as most of them have a lengthy processing time, require patients to fast overnight, require commitment of nursing/laboratory staff, can be expensive to conduct in some areas of the world and some people may not have access to these tests right away. The second control of the processing time access to these tests right away.

Whereas, non-invasive risk scoring tools can be used as the first step in mass screening programmes and as a self-administered tool for public. For example, The Diabetes UK encourages people to use their online self-administered non-invasive diabetes risk score in identifying the risk of diabetes. Using such tools is useful not only for individuals to make informed lifestyle changes but also for the society and for the healthcare systems as it can prevent or delay serious complications of diabetes among the public.

However, current evidence on the performance of similar scores developed in low-and middle-income countries is rare specially in relation to their applicability in other populations and settings. For example, in 2005, the Madras Diabetes Research Foundation (MDRF) developed the Indian Diabetes Risk score (IDRS) to triage people with undiagnosed diabetes. This was derived from the non-invasive risk factors identified in the Chennai Urban Rural Epidemiology Study (CURES), which include waist circumference, family history of diabetes, age and level of physical activity. Compared with other non-invasive diabetes risk scores developed for Indians, Hors is the most commonly externally validated tool in a large number of studies across India. Has been identified as a cost-effective, simple and easy to use tool mainly in resource restricted settings. According to the MDRF-IDRS model development study by Mohan et al, the sensitivity and specificity of IDRS is 72.5% and 60.1% respectively. Several other studies conducted in different regions of India has shown that the sensitivity and specificity of IDRS in identifying undiagnosed diabetes can vary between 50.8%-97.5% and 17.6%-91.7%.

Despite these existing studies from India, the performance of MDRF-IDRS score has not yet been evaluated in other settings or among other ethnic groups. MDRF-IDRS score is also likely to be applicable among American as it was initially developed using the risk factors recommended by the American Diabetes Association. 11 However, its performance among American has not yet been formally evaluated. There are numerous diabetes risk scores available for American but most of them require some form of clinical measurement such as blood pressure level, cholesterol level, fasting plasma glucose level, etc and hence it is difficult to use them in mass screening programmes, ²⁸ ²⁹ whereas IDRS score is a non-invasive risk score that requires only four variables. In addition, a substantial percentage of people with undiagnosed diabetes in India³⁰ as well as in America³¹ reported to have no contact with a primary care provider. This emphasises the importance of noninvasive diabetes risk scores that can be used in mass screening as well as a selfadministered tool in both settings. Moreover, people with undiagnosed diabetes are likely to be unevenly distributed in different ethnic/racial and socioeconomic groups. 32 For example, the age-standardized percentage of undiagnosed diabetes cases among Hispanics and non-Hispanic Asian are reported to be higher than all other racial/ethnic

groups in America.³² Hence it is important to assess the performance of risk scores in different ethnic groups separately.

Therefore, in this study, we aimed to assess and compare the MDRF-IDRS scores' performance among Indian and American ethnic groups including Hispanic, Non-Hispanic White, Non-Hispanic Black, and other American. Moreover, we aimed to compare MDRF-IDRS score's performance with existing non-invasive American diabetes risk scores.

Methods

MDRF-IDRS score assigns weights for each risk factor (age, abdominal obesity, physical activity level and family history of diabetes) and generates an aggregated weighted score between 0-100 as shown in Table 1. We used this weighted score in the National Health and Nutrition Examination Survey (NHANES) and Chennai Urban Rural Epidemiology Study (CURES) data to assess MDRF-IDRS scores performance in different ethnic groups including Hispanic, Non-Hispanic White, Non-Hispanic Black, other American and Indians.

Table 1: Indian Diabetes Risk Score (MDRF-IDRS Score)

NHANES is a repeated cross-sectional survey which is performed in 2-year cycles among a nationally representative sample of the US civilian, noninstitutionalized US population.³³ We used NHANES data for the years of 2003-2006 for adults aged ≥20. NHANES data on the age, ethnicity, family history of diabetes, waist circumference, and level of physical activity were obtained from relevant data files. Average level of physical activity each day was questioned in NHANES by providing study participants with four options to select. The first is "sits during the day and not walk about very much" this option was matched with the IDRS "no exercise and sedentary work" category. The second and third options of NHANES "stand or walk about a lot during the day but do not have to carry or lift things very often" and "lifts light load or has to climb stairs or hills often" were matched with the IDRS "Exercise (regular) or strenuous work" category. The third NHANES physical activity group "do heavy work or carry heavy loads" were matched with the IDRS "Exercise (regular)+ strenuous work" group. Study participants of NHANES were questioned if "any of close relatives that is blood relatives including father, mother, sisters or brothers had diabetes". The answer to this question was "yes/no" and if they answered yes this was

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matched with IDRS "both parents diabetic" category otherwise it was matched with the IDRS score's "No family history" category.

Within the NHANES cohort, people with type 2 diabetes were identified using their self-reported history of diabetes based on the question, "Have you ever been told by a doctor or health professional that you have diabetes or sugar diabetes?" (Answered yes and borderline were considered as having diabetes) OR if they had glycated haemoglobin $(HbA1c) \ge 6.5$ OR if they were taking anti-diabetes medication according to the medication inventory file. People with type 1 diabetes were excluded from the study population. Type 1 diabetes patients were defined as those who have been diagnosed with diabetes before age 30, currently taking insulin and started taking insulin within one year of the diagnosis.

We also used data from the CURES study, which is an epidemiological study conducted between 2001-2003 among a representative sample of adults (aged≥20 years) of Chennai, India.³⁴ In Phase 1 of this study, the study participants with self-reported type 2 diabetes were identified. Moreover, age and sex matched non-diabetes subjects and all participants with fasting capillary blood glucose of ≥ 110mg/dL underwent an oral glucose tolerance test. Those who confirmed having 2-hour plasma glucose levels of ≥ 200 mg/dL were identified as newly diagnosed diabetes patients. This survey collected data on age, family history of diabetes, waist circumference, and level of physical activity. In this survey physical activity was measured using an estimate for 24-h energy expenditure. 34-36 This estimate was generated using a questionnaire on the physical activity which included leisure time, household chores, work, sleep, sedentary activities and other common daily activities. The 24-hour energy expenditure was then categorised into vigorous, moderate, mild, and sedentary physical activity levels. We mapped "vigorous" energy expenditure level to "Exercise[regular] +strenuous work" category of IDRS score; "moderate" and "mild" levels into "Exercise [regular] or strenuous work" category of IDRS and "sedentary" physical activity level to "No exercise and sedentary work" category of IDRS score.

Descriptive analysis was carried out to identify the characteristics of the study participants with and without diabetes in each ethnic group. Chi squared test was used to identify any significant difference between the groups. The MDRF-IDRS score's highest performance

cut-off point was selected using the Youden Index. The Youden's index combines sensitivity and specificity into a single measure (sensitivity + specificity - 1). The area under the receiver operating characteristic curve (AUC) and sensitivity, specificity, positive predictive value and negative predictive value of the relevant cut-off measures were used to assess the performance of MDRF-IDRS score within different ethnic groups.

We then reviewed existing literature on non-invasive American diabetes risk scores for identifying undiagnosed type two diabetes. We excluded models developed using nonregression techniques (e.g. neural networks, classification trees, and machine learning) due to the lack of transparency in these non-regression model development techniques especially in relation to reproducibility and validation in external datasets.^{37 38} We identified only one non-invasive American diabetes risk score fulfilling this criteria and it is the Bang et al³⁹ score with sensitivity of 79% and specificity of 67% in detecting undiagnosed diabetes among American. This score was developed using logistic regression modelling and included age, sex, family history of diabetes, history of hypertension, body mass index, and physical activity level were included as variables. In this score, study participants were considered having family history of diabetes if their parents or siblings had diabetes. Physical activity was measured based on their answer (Yes/No) to the question "Are you physically active". IDRS scores' performance was then compared with this non-invasive American diabetes risk score. NHANES survey uses a complex, multistage, probability sampling design to produce a nationally representative data set. Therefore, we used NHANES survey weights to take the NHANES complex sampling design into account during the data analysis. Data management and analysis were performed using Stata 15 (Stata Corp., College Station, Texas, USA).

Results

There were 11,035 study participants (2,292 Indians and 8743 American) eligible to include in this study and they were aged ≥20, did not have type I diabetes and they had complete data for all the variables considered in IDRS score. Detailed participant flow diagram is given in Appendices (Figure 2 & Figure 3). The study population characteristics were varied between the ethnic groups (Table 2). Non-Hispanic-White participants had the lowest prevalence of diabetes (9.1%) whereas Indian participants had the highest prevalence of

diabetes (15.5%). The mean age varied between 38.6-47.8 years. The mean waist circumference varied between 83.4-98.9 cm. Physical activity level among Indians were relatively low when compared with the other groups, 80.8% of Indians were in the low physical activity level group whereas in the other ethnic groups 14.7%-25.3% study participants were in the low physical activity group. Indians and Non-Hispanic Black had the lowest (23.7%) and highest (54.2%) prevalence of family history of diabetes respectively.

As shown in Table 3, age, waist circumference, and family history of diabetes were significantly associated with diabetes in all ethnic groups. Study participants with diabetes were older, had higher waist circumference and had higher prevalence of family history compared to those who did not have diabetes and this trend was common in all ethnic groups. There was no significant difference in the gender distribution among American diabetes and non-diabetes study participants. However, the distribution of gender among Indian diabetes and non-diabetes study participants were significantly different (p=0.02), with higher percentage of males in the diabetes group. People with diabetes had lower level of physical activity compared to those who did not have diabetes and this difference was statistically significant in all ethnic groups apart from the Other-American ethnic group.

The MDRF-IDRS score performed well among Indians and American as shown in Figure 1 and Table 4. The AUC for IDRS score among Indians was 0.7345 and the cut off≥ 60 had the highest performance with sensitivity and specificity of 80.2% and 57.3% respectively. MDRF-IDRS score cut off ≥ 70 had the highest discriminative performance among Hispanic, Non-Hispanic White and Non-Hispanic Black ethnic groups with sensitivity and specificity ranging between 70.1-86.9% and 61.2-72.2% respectively. For other-American, IDRS score ≥60 was identified as the highest performing cut off with sensitivity and specificity of 94.8% and 48.9% respectively. The AUC for Hispanic, Non-Hispanic White, Non-Hispanic Black and Other-American ethnicities was 0.7952, 0.7749, 0.8148, and 0.7872 respectively. Moreover, MDRF-IDRS score had a high negative predictive value between 0.94-0.98 for all ethnicities showing that those who get a negative test result for IDRS are highly unlikely to have diabetes.

As given in Table 5, the comparison of IDRS score with existing American non-invasive score for undiagnosed diabetes showed that IDRS score has almost the same performance as Bang et al³⁹ score which uses six risk factors (Age, Gender, Family history of diabetes, self-reported high blood pressure/medication use for hypertension, weight, physical activity level).

Table 2: IDRS variables and diabetes prevalence of the study population by different ethnic group

Table 3: Differences between people with diabetes and people without diabetes by different ethnic group

Figure 1: IDRS AUC for different ethnic groups

Table 4: Discriminative performance of IDRS score for different ethnicities

Table 5: Comparison of IDRS Score's performance with other **non-invasive American diabetes risk scores**

Discussion

This is the first study to assess MDRF-IDRS score's performance among American and to carry out a comparison of its performance among Indian and American. It has shown that this score performs well not only among Indians but also in among American ethnic groups including Hispanic, Non-Hispanic White, and Non-Hispanic Black. The MDRF-IDRS score can be used to identify people with undiagnosed diabetes in mass health screening programmes. This risk score has a high negative predictive value for all ethnicities and therefore it is suitable as a triage tool to rule out those with a negative test result. Those who get positive test results for IDRS can then be prioritised in providing further testing for diabetes to confirm their diabetes status. The current study results are in line with the existing literature on MDRF-IDRS score performance in different regions in India^{13 17-26} and almost all studies showed that 60 is the optimal cut off point for the MDRF-IDRS score (Table 6).

When definitions of physical activity and family history were not identical between the datasets and IDRS score, we tried to use the best available variable within each dataset and categorise them in the best possible way to achieve reasonable consistency across datasets and IDRS score. However, the physical activity categories used in IDRS score seems not specific enough to obtain accurate and comparable measurements across different datasets. This may have resulted in the lower specificity of IDRS score identified in the current study as well as in most of the previously conducted studies. ^{11 13 16-27}

Table 6: Performance of IDRS Score in identifying undiagnosed diabetes according to previously published studies in India

The MDRF IDRS Score was developed using the non-invasive risk factors, recommended by the American Diabetes Association¹¹ and may explain the score's high performance in American ethnicities. Comparison of IDRS Score with existing American non-invasive diabetes scores showed that IDRS has almost the same performance as Bang et al³⁹ score. Moreover, IDRS score uses only four variables whereas Bang et al³⁹ score uses six variables including self-reported hypertension which could be difficult to obtain accurately in resource restricted settings as people in these settings are less likely to be aware of their blood pressure levels. 40 41

United States. In United States in 2017 there were 7.2 million people with undiagnosed diabetes and this represents 23.8% of people with diabetes. People with diabetes in the US incur average medical expenditures that are 2.3 times higher than that in the absence of diabetes and the total estimated cost of diagnosed diabetes in 2017 was \$327 billion. Most of these medical expenditure are due to the diabetes related complications and comorbidities. Delayed diagnosis of diabetes can lead to people having more complications and in return this will cause more burden on the health care system. In India, the economic burden due to diabetes is even higher and it is expected to rise particularly among the economically disadvantaged groups. Hence early identification of diabetes though screening can have a substantial impact on reducing the costs associated with delayed diagnosis in both settings.

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In addition, using non-invasive diabetes risk assessment tool such as MDRF-IDRS Score has the potential to reduce the cost of diabetes diagnosis by ruling out those who are unlikely to have diabetes. For example, as shown in Appendices (Figure 4 (b)) among 1000 Hispanic, 325 get a positive result and 675 get a negative test result for IDRS score. Due to high negative predictive value of IDRS score further testing for diabetes will only be required in 325 patients. Therefore, MDRF-IDRS score can help prioritise in early diagnosis of diabetes which also facilitates early management and optimal control of diabetes and reduced risk of complications and associated economic burden for the healthcare system and society.

In this study we have shown that IDRS performs well in different settings and in different populations regardless of the differences in population characteristics such as age distribution, physical activity levels, and family history. We've used the same Indian dataset that was used to develop IDRS and this is likely to have slightly overestimated the performance of IDRS among Indians as risk scores tend to perform well in the development data set. 38 46 Despite this limitation of internal validation of IDRS score it has performed well in the external validation carried out using the NHANES dataset providing that it has good performance in different settings and populations. This study included a nationally representative sample of study participants from America and it combined two consecutive NHANES 2-year cycles (2003-2006) creating a large sample and therefore, the results of this study are generalisable to the American population. However, in the NHANES dataset, the family history of diabetes question did not distinguish between one parent or both parent having diabetes and therefore, we allocated 20 points to those who said yes for family history of diabetes. Whereas, ideally according to the MDRF-IDRS score 10 points should be allocated to those having a single parent with diabetes and 20 points to those with both parents having diabetes as shown in Table 1. The lack of this information from the dataset might have altered IDRS score's performance among American and future studies could investigate this by using an appropriate dataset.

Conclusion

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The Madras Diabetes Research Foundation-Indian Diabetes Risk Score performs well not only among Indians but also among Hispanic, Non-Hispanic White, Non-Hispanic Black and other American. It can be used as a screening tool to help in early diagnosis, management and optimal control of diabetes mainly in mass screening programmes in India and America.

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Author Contribution

The study was conceived by GN, SS and MN. The study was designed by the Ornate India grant co-applicants (GN, SS, VM, RR). MN and GN provided the statistical input. MN drafted the manuscript and all authors commented on drafts and approved the final version.

Table 1: Indian Diabetes Risk Score (MDRF-IDRS Score)

Particulars	Score
Age	
<35	0
35-49	20
50+	30
Abdominal obesity	
Waist <80cm [female], <90cm [male]	0
Waist ≥80-89cm [female], ≥ 90-99 cm [male]	10
Waist ≥90cm [female], ≥ 100cm [male]	20
Physical activity	
Exercise [regular] + strenuous work	0
Exercise [regular] or strenuous work	20
No exercise and sedentary work	30
Family history	
No family history	0
Either parent	10
Both parents	20
Minimum score	0
Maximum score	100

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Table 2: IDRS variables and diabetes prevalence of the study population by different ethnic group

21

			Ethnicity		
	Indian	Hispanic	Non-Hispanic	Non-Hispanic	Other-
			White	Black	American
	N=229	N*=1786	N*=4479	N*=1849	
	2	(PS=15,358,20	(PS=138,673,83	(PS=21,575,79	N*=629
		3)	2)	4)	(PS=16,772,05
					4)
Diabetes	15.5%	11.1%	9.1%	14.3%	12.3%
(Type II)					
Mean Age	39.6	38.6	47.8	43.4	42.4
(years)					
Mean Waist	83.4	97.1	97.8	98.9	92.9
Circumferen					
ce					
(cm)					
Gender					
Male	46.3%	52.7%	47.9%	44.1%	46.1%
Family					
History of	23.7%	52.5%	43.1%	54.2%	45.3%
Diabetes					
Yes					
Level of					
Physical					
Activity	80.8%	14.7%	23.9%	25.3%	22.7%
Low	17.0%	71.1%	68.1%	67.9%	70.4%
Moderate	2.1%	14.2%	8.0%	6.8%	6.9%
High					

N: CURES dataset total number of individuals sampled using systematic random sampling

N*: NHANES dataset total number of individuals sampled using complex, multistage, probability sampling design

PS: NHANES Survey weight adjusted nationally representative population size

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NOTE: The proportions presented for American ethnicities represent the survey weight adjusted proportion out of the population size for each category

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Table 3: Differences between diabetic and non-diabetic study participants by different ethnic group

		Indian		Hispanic			Non-	Hispanic	nic White Non-Hispan			ic Black Other-Ame		er-Ameri	can
		N=2292			N*=1786	5		N*=447	9		N*=1849			N*=629	
				(PS	=15,358,	203)	(PS	=138,673	,832)	(PS=	=21,575,7	794)	(PS=	16,772,0	154)
	Dial	petic		Diabetic			Dial	oetic		Diabetic			Diabetic		
			P			P			P value			Р			Р
	No	Yes	value	No	Yes	value	No	Yes		No	Yes	value	No	Yes	valu
	n=1925	n=367		PS=13	PS=1,		PS=12	PS=12		PS=18	PS=3,		PS=14	PS=2,	е
	(84%)			,668,	689,4		6,193,	,480,6		,555,1	020,6		,715,4	056,6	
		(16%)		801	02		187	45		82	12		07(88	47	
				(89%)	(11%)		(91%)	(9%)		(86%)	(14%)		%)	(12%)	
Gender															
Male	45%	54%		53%	51%	0.537	48%	50%	0.294	45%	40%	0.162	47%	40%	0.29
			0.001												4
Age															
(years)															
20-35	45%	14%	<0.00	52%	12%	<0.00	27%	5%	<0.001	39%	4%	<0.00	38%	8%	<0.0
35-49	37%	41%	1	33%	32%	1	32%	14%		34%	26%	1	36%	32%	01

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50+	18%	45%		15%	56%		40%	80%		27%	70%		26%	60%	
Waist															
Circumference															
(cm)															
Female	54%	34%	<0.00	21%	6%	<0.00	22%	5%	<0.001	24%	8%	<0.00	34%	14%	<0.0
Male	31%	41%	1	31%	16%	1	27%	9%		23%	14%	1	27%	14%	01
<80	15%	25%		47%	78%		51%	86%		53%	78%		39%	72%	
<90															
80-89															
90-99															
>90															
>100															
Family History															
of Diabetes	22%	34%	<0.00	50%	71%	<0.00	41%	64%	<0.001	50%	79%	<0.00	41%	72%	<0.0
Yes			1			1						1			01
Level of															
Physical															

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															25
Activity	80%	86%		13%	25%	<0.00	23%	31%	<0.001	23%	38%	<0.00	23%	22%	0.45
Low	18%	12%	0.023	71%	67%	1	68%	66%		69%	59%	1	70%	74%	0
Moderate	2%	2%		15%	8%		9%	3%		8%	3%		7%	4%	
High															

N: CURES dataset total number of individuals sampled using systematic random sampling

N*: NHANES dataset total number of individuals sampled using complex, multistage, probability sampling design

PS: NHANES Survey weight adjusted nationally representative population size

NOTE: The proportions presented for American ethnicities represent the survey weight adjusted proportion out of the population size for each category

Table 4: Discriminative performance of IDRS score for different ethnicities

Ethnicity	IDRS cut-off with highest performance	Sensitivity	Specificity	ratio of a positive test (LR+)	ratio of a negative test (LR-)	Youden Index	Probability of having diabetes if test positive (PPV)	Probability of No Diabetes if test negative (NPV)
Indian	60	80.2%	57.3%	1.88	0.34	0.37	0.26	0.94
Hispanic	70	70.1%	72.2%	2.52	0.41 0.42 0		0.24	0.95
Non- Hispanic White	70	86.9%	54.8%	1.92	0.24	0.42	0.16	0.98
Non- Hispanic Black	70	86.6%	61.2%	2.23	0.22	0.48	0.27	0.96
Other- American	60	94.8%	48.9%	1.85	0.11	0.44	0.21	0.98

- IDRS: Indian Diabetes Risk Score
- Youden Index = Sensitivity + Specificity 1

(Youden Index is a single statistic that captures the performance of a diagnostic test)

- PPV Positive Predictive Value
- NPV Negative Predictive Value

Table 5: Comparison of IDRS Score's performance with non-invasive American diabetes risk score

American Ethnic Groups	Area Under the Receiver Ope	rating Curve (AUC) in NHANES
	IDRS Score	Bang et al Score ¹⁴
	AUC (95% CI)	AUC (95% CI)
Hispanic	0.7952 (0.7948 – 0.7955)	0.8060 (0.8056- 0.8062)
Non-Hispanic White	0.7749 (0.7747 – 0.7750)	0.7990 (0.7989 – 0.7991)
Non-Hispanic Black	0.8148 (0.8145 – 0.8150)	0.8221 (0.8218 – 0.8222)
Other American	0.7872 (0.7869 – 0.7875)	0.7647 (0.7644 – 0.7650)

IDRS variables: Age, Waist circumference, Physical activity, Family history of diabetes

Bang et al variables: Age, Gender, Family history of diabetes, self-reported high blood pressure/medication use for hypertension, weight, physical activity level

Table 6: Performance of IDRS Score in identifying undiagnosed diabetes according to previously published studies in India

Paper	Setting	Sample Size (n)	Age	AUC	Cut-	Sensitivity	Specificity	PPV	NPV
			Group		off	(%)	(%)	(%)	(%)
Mohan et al,	Chennai	2350	≥ 20	0.698	≥60	72.5	60.1	17	95.1
2005 ¹¹			years						
Nandeshwar,	Bhopal city, Madhya	250	≥ 25	-	≥60	94.7	44.8	50.8	93.3
2010 ¹⁷	Pradesh		years						
Adhikari et	Mangalore	551	≥ 20	_	≥60	62.2	73.7	-	-
al, 2010 ¹³			years						
Patel et al,	Bhavnagar city	260	≥ 30	0.838	≥60	92.5	62.3	30.8	97.86
2012 ¹⁸			years						
Stanley et al,	Chennai	154	19 - 99	-	≥60	100	17.6	-	-
2012 ¹⁹			years						
Taksande et	Rural community of	478	≥ 45	_	≥60	97.5	87.89	-	-
al, 2012 ²⁰	central India		years						
Sathish et al,	Thiruvananthapuram,	451	15 –	0.80	≥60	85.7	59.4	32.6	94.8
2013 ²¹	Kerala		64						
			years						
Bhadoria et	Jabalpur District	911	Adults:	-	≥60	26.4	91.7	26.1	91.8

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al, 2014 ²²				no a	age						
				rang	ge						
				give	n						
Ahmed et al,	Jaipur	551		≥	20	-	≥60	62.2	73.7	-	-
2015 ²³				year	·s						
Basu et al,	Nationally		567	25	-	-	≥60	50.8	64.0	11.1	93.3
2015 ²⁷	representative		million	65							
	sample of Indians		(simulated	year	s						
			sample								
			using data								
			from 58								
			studies)								
Najeeb et al,	Mullana, Ambala	200		2	30	0.731	≥60	65.0	62.5	-	-
2015 ²⁴				year	rs						
Dudeja et al,	Urban slum, India	155		Mea	an	-	≥60	95.1	28.9	32.5	94.3
2017 ²⁵				age:							
				49.6	8						
				SD:							
				14.8	80						

Kaushal et	Shimla	465	≥ 20	-	≥70	61.3	56.14	23.4	86.8
al, 2017 ²⁶			years						
Oommen et	Tamil Nadu	4896	30-64	-	-	59.1	59.4	8.9	95.6
al, 2017 ¹⁶			years						

AUC: Area Under the Curve, PPV: Positive Predictive Value, NPV: Negative Predictive Value

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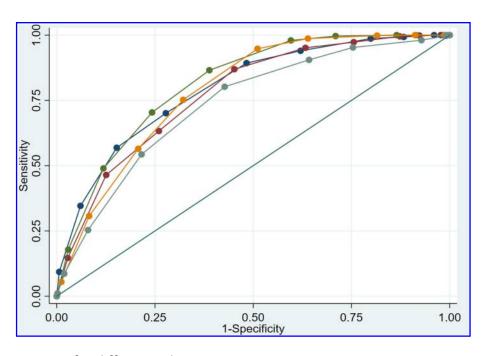


Figure 1: IDRS AUC for different ethnic groups

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Figure 1 Legend



NHANES 2003-2006 20,470 Age <20 = 10,450 Age ≥ 20 10,020 Type 1 diabetes = 23 Adults (Age≥ 20) without Type I diabetes 9,997 Missing data Waist =1049 Physical Activity = 7 Complete data for Adults (Age≥ 20) Family History = 198 without Type I diabetes 8,743 Non-Diabetic Type II diabetic 7,576 1, 167

Figure 2: NHANES Data flow chart

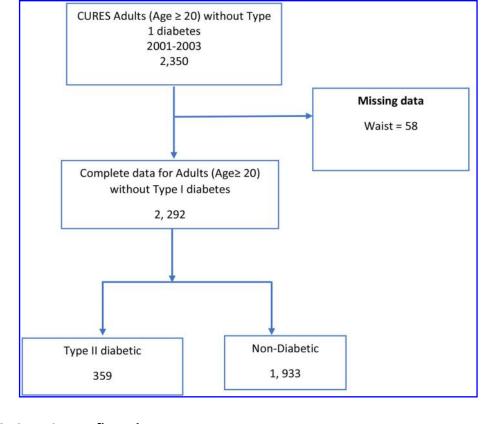


Figure 3: CURES Data flow chart

5(a)

Indian

Table 5: Optimum cut-off for different ethnic groups

			Correctly			Youden
Cutpoint	Sensitivity	Specificity	Classified	LR+	LR-	Index
(>= 0)	100.00%	0.00%	15.66%	1		0.00
(>= 10)	100.00%	0.67%	16.23%	1.0068	0	0.01
(>= 20)	99.72%	0.93%	16.40%	1.0066	0.2991	0.01
(>= 30)	98.05%	7.19%	21.42%	1.0565	0.2712	0.05
(>= 40)	95.26%	24.62%	35.69%	1.2639	0.1923	0.20
(>= 50)	90.53%	35.80%	44.37%	1.4101	0.2646	0.26
(>= 60)	80.22%	57.27%	60.86%	1.8774	0.3453	0.37
(>= 70)	54.32%	78.43%	74.65%	2.5179	0.5825	0.33
(>= 80)	25.35%	92.03%	81.59%	3.1817	0.8111	0.17
(>= 90)	8.64%	98.09%	84.08%	4.5113	0.9315	0.07
(>= 100)	1.11%	99.79%	84.34%	5.3844	0.9909	0.01
(> 100)	0.00%	100.00%	84.34%		1	0.00

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5(b) Hispanic

			Correctly			Youden
Cutpoint	Sensitivity	Specificity	Classified	LR+	LR-	Index
(>= 0)	100.00%	0.00%	11.17%	1		0.00
(>= 10)	100.00%	2.26%	13.18%	1.0232	0	0.02
(>= 20)	100.00%	3.98%	14.71%	1.0415	0	0.04
(>= 30)	99.29%	11.70%	21.48%	1.1245	0.0603	0.11
(>= 40)	98.66%	20.10%	28.88%	1.2349	0.0665	0.19
(>= 50)	94.00%	37.95%	44.22%	1.515	0.1581	0.32
(>= 60)	89.29%	51.70%	55.90%	1.8486	0.2072	0.41
(>= 70)	70.09%	72.24%	72.00%	2.5247	0.414	0.42
(>= 80)	56.86%	84.74%	81.63%	3.7265	0.5091	0.42
(>= 90)	34.62%	93.96%	87.33%	5.7272	0.6959	0.29
(>= 100)	9.32%	99.36%	89.30%	14.5623	0.9126	0.09
(> 100)	0.00%	100.00%	88.83%		1	0.00

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5© Non-Hispanic

White

			Correctly			Youden
Cutpoint	Sensitivity	Specificity	Classified	LR+	LR-	Index
(>= 0)	100.00%	0.00%	9.06%	1		0.00
(>= 10)	100.00%	1.20%	10.15%	1.0121	0	0.01
(>= 20)	100.00%	1.70%	10.60%	1.0173	0	0.02
(>= 30)	99.85%	7.73%	16.07%	1.0821	0.0199	0.08
(>= 40)	99.52%	12.70%	20.56%	1.1399	0.0381	0.12
(>= 50)	97.31%	24.42%	31.02%	1.2875	0.1103	0.22
(>= 60)	95.14%	36.71%	42.00%	1.5033	0.1323	0.32
(>= 70)	86.95%	54.81%	57.72%	1.924	0.2382	0.42
(>= 80)	63.29%	74.00%	73.03%	2.4343	0.4961	0.37
(>= 90)	46.45%	87.42%	83.71%	3.6922	0.6125	0.34
(>= 100)	14.67%	97.20%	89.72%	5.2336	0.8779	0.12
(> 100)	0.00%	100.00%	90.94%		1	0.00

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5(d) Non-Hispanic

Black

			Correctly			Youden
Cutpoint	Sensitivity	Specificity	Classified	LR+	LR-	Index
(>= 0)	100.00%	0.00%	14.25%	1		0.00
(>= 10)	100.00%	0.93%	15.05%	1.0094	0	0.01
(>= 20)	100.00%	1.12%	15.21%	1.0113	0	0.01
(>= 30)	100.00%	8.56%	21.60%	1.0937	0	0.09
(>= 40)	100.00%	13.51%	25.84%	1.1562	0	0.14
(>= 50)	99.65%	29.04%	39.10%	1.4042	0.0121	0.29
(>= 60)	97.97%	40.42%	48.62%	1.6444	0.0502	0.38
(>= 70)	86.55%	61.17%	64.78%	2.2287	0.2199	0.48
(>= 80)	70.39%	75.74%	74.98%	2.9019	0.3909	0.46
(>= 90)	48.95%	88.13%	82.55%	4.1248	0.5792	0.37
(>= 100)	17.83%	97.08%	85.79%	6.1105	0.8464	0.15
(> 100)	0.00%	100.00%	85.75%		1	0.00

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5(e) Other

American

			Correctly			Youden
Cutpoint	Sensitivity	Specificity	Classified	LR+	LR-	Index
(>= 0)	100.00%	0.00%	12.31%	1		0.00
(>= 10)	100.00%	1.20%	13.35%	1.0121	0	0.01
(>= 20)	100.00%	1.34%	13.48%	1.0136	0	0.01
(>= 30)	100.00%	8.80%	20.03%	1.0965	0	0.09
(>= 40)	99.83%	18.49%	28.50%	1.2249	0.0091	0.18
(>= 50)	98.67%	36.08%	43.78%	1.5436	0.0368	0.35
(>= 60)	94.79%	48.91%	54.56%	1.8553	0.1066	0.44
(>= 70)	75.23%	67.82%	68.73%	2.3378	0.3652	0.43
(>= 80)	56.48%	79.27%	76.46%	2.7242	0.5491	0.36
(>= 90)	30.71%	91.76%	84.25%	3.7259	0.7551	0.22
(>= 100)	5.46%	98.82%	87.34%	4.6433	0.9567	0.04
(> 100)	0.00%	100.00%	87.69%		1	0.00

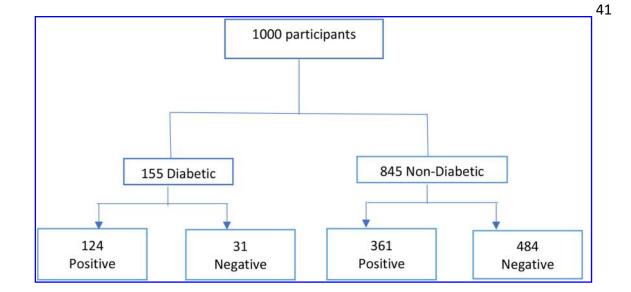


Figure 4: Performance of IDRS Score in Natural Frequencies

Indian

Prevalence = 15.5%; Sensitivity = 80.2%; Specificity = 57.3%

Probability of Diabetes if IDRS test positive = 124/ (124+361) = 0.26

Probability of no diabetes if IDRS test negative = 484/ (484+31) = 0.94

42 1000 participants 889 Non-Diabetic 111 Diabetic 78 33 247 642 Positive Positive Negative Negative

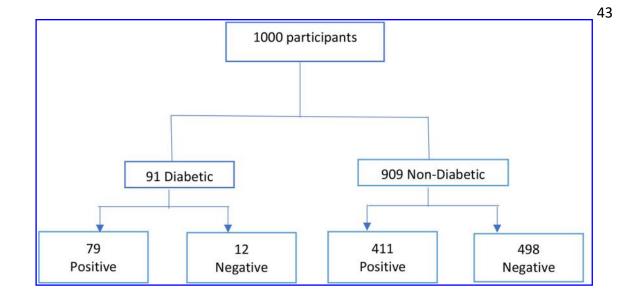
Hispanic

Prevalence = 11.1%; Sensitivity = 70.1%; Specificity = 72.2%

Probability of Diabetes if IDRS test positive = 78/ (78+247) = 0.24

Probability of no diabetes if IDRS test negative = 642/(642+33) = 0.95



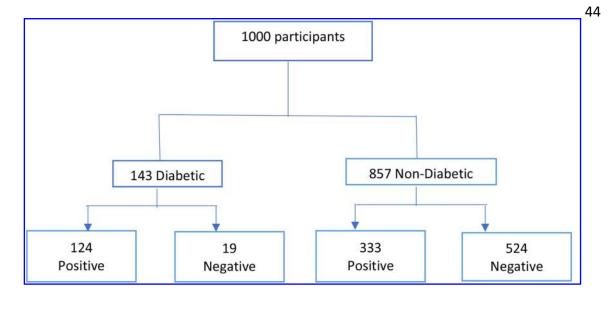


Non-Hispanic White

Prevalence = 15.5%; Sensitivity = 80.2%; Specificity = 57.3%

Probability of Diabetes if IDRS test positive = 79/ (79+411) = 0.16

Probability of no diabetes if IDRS test negative = 498/ (498+12) = 0.98

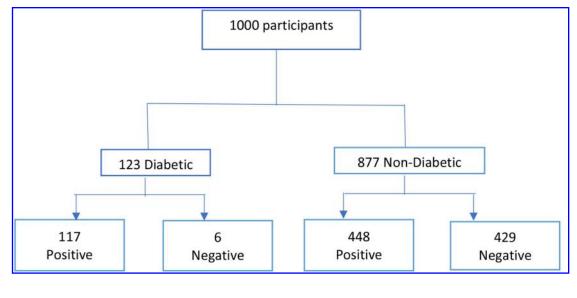


Non-Hispanic Black

Prevalence = 14.3%; Sensitivity = 86.5%; Specificity = 61.2%

Probability of Diabetes if IDRS test positive = 124/ (124+333) = 0.27

Probability of no diabetes if IDRS test negative = 524/ (524+19) = 0.96



Other American

Prevalence = 12.3%; Sensitivity = 94.8%; Specificity = 48.9%

Probability of Diabetes if IDRS test positive = 124/ (124+361) = 0.21

Probability of no diabetes if IDRS test negative = 484/ (484+31) = 0.98