

RESEARCH ARTICLE

Open Access



# Sex-specific prevalence of coronary heart disease among Tehranian adult population across different glycemic status: Tehran lipid and glucose study, 2008–2011

Seyyed Saeed Moazzeni<sup>1</sup>, Hamidreza Ghafelehbashi<sup>1</sup>, Mitra Hashemini<sup>1</sup>, Donna Parizadeh<sup>1</sup>, Arash Ghanbarian<sup>1</sup>, Fereidoun Azizi<sup>2</sup> and Farzad Hadaegh<sup>1\*</sup> 

## Abstract

**Background:** Coronary heart disease (CHD) is one of the leading causes of death. Alarming Iranian populations had a high rank of CHD worldwide. The current study aimed to assess the prevalence of CHD across different glycemic categories.

**Methods:** This study was conducted on 7718 Tehranian participants (Men = 3427) aged  $\geq 30$  years from 2008 to 2011. They were categorized based on glycemic status. The prevalence of CHD was calculated in each group separately. CHD was defined as hospital records adjudicated by an outcome committee. The association of different glycemic categories with CHD was calculated using multivariate logistic regression, compared with normal fasting glucose /normal glucose tolerance (NFG/NGT) group as reference.

**Results:** The age-standardized prevalence of isolated impaired fasting glucose (iIFG), isolated impaired glucose tolerance (iIGT), both impaired fasting glucose and impaired glucose tolerance (IFG/IGT), newly diagnosed diabetes mellitus (NDM), and known diabetes mellitus (KDM) were 14.30% [95% confidence interval (CI): 13.50–15.09], 4.81% [4.32–5.29], 5.19% [4.71–5.67], 5.79% [5.29–6.28] and 7.72% [7.17–8.27], respectively. Among a total of 750 individuals diagnosed as cases of CHD (398 in men), 117 (15.6%), 453 (60.4%), and 317 (42.3%) individuals had a history of myocardial infarction (MI), cardiac procedure, and unstable angina, respectively. The age-standardized prevalence of CHD for the Tehranian population was 7.71% [7.18–8.24] in the total population, 8.62 [7.81–9.44] in men and 7.19 [6.46–7.93] in women. Moreover, among diabetic participants, the age-standardized prevalence of CHD was 13.10 [9.83–16.38] in men and 10.67 [8.90–12.44] in women, significantly higher than corresponding values for NFG/NGT and prediabetic groups.

Across six levels of glycemic status, CHD was associated with IFG/IGT [odds ratio (OR) and 95% CI: 1.38 (1.01–1.89)], NDM [1.83 (1.40–2.41)], and KDM [2.83 (2.26–3.55)] groups, in the age- and sex-adjusted model. Furthermore, in the full-adjusted model, only NDM and KDM status remained to be associated with the presence of CHD by ORs of 1.40 (Continued on next page)

\* Correspondence: [fzhadaegh@endocrine.ac.ir](mailto:fzhadaegh@endocrine.ac.ir)

<sup>1</sup>Prevention of Metabolic Disorders Research Center, Research Institute for Endocrine Sciences, Shahid Beheshti University of Medical Sciences, P.O. Box: 19395-4763, No. 24, Parvaneh Street, Velenjak, Tehran, Iran  
Full list of author information is available at the end of the article



© The Author(s). 2020 **Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>. The Creative Commons Public Domain Dedication waiver (<http://creativecommons.org/publicdomain/zero/1.0/>) applies to the data made available in this article, unless otherwise stated in a credit line to the data.

(Continued from previous page)

(1.06–1.86) for NDM and 1.91 (1.51–2.43) for KDM.

**Conclusion:** The high prevalence of CHD, especially among diabetic populations, necessitates the urgent implementation of behavioral interventions in the Tehranian population, according to evidence-based guidelines for the clinical management of diabetic patients.

**Keywords:** Coronary heart disease, Diabetes mellitus, Prediabetes, Prevalence, Tehran lipid and glucose study

## Background

Coronary heart disease (CHD) is one of the most common causes of death worldwide [1, 2]. Its global fatality rate increased from 7.3 million in 2007 to 8.93 million deaths in 2017 [1]. We previously reported that the overall prevalence of CHD was 21.8% (22.3% among women and 18.8% among men) in 1991–2001 among residents of Tehran, as a metropolitan city [3]. Moreover, in Tehran, the incidence rate of CHD was 10.5 and 6.1 per 1000 person-years among men and women, respectively [4]. Importantly, among Tehranian adults aged  $\geq 30$  years, over 40% of mortality is attributed to cardiovascular disease (CVD) [5].

Type 2 diabetes mellitus (T2DM) is a major leading factor for CHD and its mortality [6, 7]. In addition to the well-known relationship between diabetes and CVD, it has also been shown that prediabetes status could lead to CHD and CVD [8, 9]. Based on national studies in 2011, about 11.4 and 14.6% of Iranian adults had diabetes mellitus (DM) and impaired fasting glucose (IFG), respectively. Furthermore, there was an alarming increase of 35.1% in the prevalence of DM from 2005 to 2011 [10]. A prediabetes tsunami (included both impaired glucose tolerance (IGT) and IFG) was also reported among a Tehranian population, with  $\geq 4\%$  of adult individuals developing prediabetes each year [11].

The current study aims to report the population-based prevalence of CHD among Tehranian adults, aged  $\geq 30$  years, according to their glycemic status in phase IV (2008–2011) of the oldest cohort study in the Middle East and North Africa (MENA) region, the Tehran Lipid and Glucose Study (TLGS) [12].

## Methods

### Study design and population study

The current study was performed within the framework of the TLGS, which is an ongoing cohort study being conducted on a representative sample of Tehranian citizens. The TLGS have the aim of determining the epidemiological aspects of non-communicable diseases (NCDs) and their risk factors. The TLGS also intended to prevent NCDs by developing healthier lifestyles. Further details for the TLGS have been described before [12]. Briefly, after the first baseline examination (1999–2001), participants were followed-up until 2011. For this

study, 8400 individuals aged  $\geq 30$  years were enrolled from phase IV of TLGS (2008–2011). Firstly, we excluded 497 individuals whose glycemic status was not differentiable for us. Secondly, we excluded 177 subjects with missing data on covariates, including body mass index (BMI), total cholesterol (TC), high-density lipoprotein cholesterol (HDL-C), systolic blood pressure (SBP), diastolic blood pressure (DBP), family history of CVD, and smoking status (overlap features between numbers considered). Finally, due to the lack of information on the outcome (CHD) assessment, eight individuals were excluded, and 7718 participants remained eligible for analysis of the current study.

### Clinical and laboratory measurements

Using pretested questionnaires, an interviewer gathered data that included demographic data, smoking status, education level, drug history, past medical history, and family history. Details of blood pressure (BP) and anthropometric parameters measurements in the TLGS setting have been published previously [8]. After over 12 h of fasting, blood samples were drawn between 07:00 AM and 09:00 AM and then analyzed on the same day. Apart from those who had on glucose-lowering medications, a standard oral glucose tolerance test with 75 g glucose was done for all participants. Fasting plasma glucose (FPG) and 2-h post-challenge plasma glucose (2 h-PCPG) were measured by enzymatic colorimetric glucose oxidase method, both inter- and intra-assay coefficient of variations were  $< 2.2\%$ . More details of laboratory measurements have been published elsewhere [8].

### Definition of terms

Participants were categorized into different groups as follows: Normal fasting glucose (NFG)/normal glucose tolerance (NGT),  $FPG < 5.6$  and  $2\text{ h-PCPG} < 7.7$  mmol/L; isolated impaired fasting glucose (iIFG),  $5.6 \leq FPG < 7$  and  $2\text{ h-PCPG} < 7.7$  mmol/L; isolated impaired glucose tolerance (iIGT),  $7.7 \leq 2\text{ h-PG} < 11.1$  and  $FPG < 5.6$  mmol/L; combined IFG and IGT (IFG/IGT),  $5.6 \leq FPG < 7$  and  $7.7 \leq 2\text{ h-PCPG} < 11.1$  mmol/L [9]. Moreover, in the present study, prediabetes status was defined as the presence of IFG or IGT. Finally, newly diagnosed diabetes mellitus (NDM) was defined as  $FPG \geq 7.0$  or  $2\text{ h-PCPG} \geq 11.1$  mmol/L among those participants were not

on glucose-lowering medications and known diabetes mellitus (KDM) as subjects with positive history of taking any glucose lowering medications. Having TC  $\geq$  5.2 mmol/L or using lipid-lowering medications defined as hypercholesterolemia. Low HDL-C was defined as HDL-C  $<$  1.036 mmol/L for men and  $<$  1.295 mmol/L for women, or taking lipid-lowering medications. Based on the seventh report of the Joint National Committee on prevention, detection, evaluation, and treatment of high blood pressure (the JNC 7 report) [13], hypertension was considered as either of having SBP  $\geq$  140 mmHg or DBP  $\geq$  90 mmHg or the usage of any anti-hypertensive medications. Smoking status was categorized into three levels, including current, past, and never smoker. Education levels were classified as  $<$  6 years (reference group), 6–12 years, and  $>$  12 years. By the Modifiable Activity Questionnaire (MAQ), which judged all types of activities [8], physical activity was evaluated. Low physical activity (inactive person) was defined as not achieving a minimum score of 600 MET (metabolic equivalent task)-minutes per week [14]. If there was at least one history of CHD/stroke in a male first-degree relative aged  $<$  55 years or in a female first-degree relative aged  $<$  65 years, the family history of premature CVD is considered positive.

#### Definition of CHD

Details of the collection of outcome data have been reported elsewhere [8]. To summarize, each individual was under continuous surveillance for any medical outcome leading to hospitalization. As a part of the cohort data collection, a trained nurse called all participants annually and recorded any medical events experienced during the last year. A trained physician followed-up any reported event by a home visit for medical data gathering. Collected data were then evaluated by a consulting committee, the outcome committee, included a principal investigator, an internist, an endocrinologist, a cardiologist, an epidemiologist, and the physician that collected the outcome data. Every confirmed event was considered as a NCD outcome based on ICD-10 criteria [8, 15]. In this study, CHD was selected from ICD-10 rubric I20–I25. CHD cases included [15–18]:

- (1) **Myocardial infarction (MI)**, included a) definite MI diagnosed by diagnostic electrocardiogram (ECG) and biomarkers (including CK, CK-MB, CK-MBm, troponin (cTn), and myoglobin), b) probable MI distinguished by positive ECG findings plus cardiac symptoms or signs and biomarkers showing negative or equivocal results.
- (2) **Cardiac procedure**, defined as a) angiography proven CHD with a result of  $\geq$  50% stenosis in at least one major coronary vessel, b) history of angioplasty or bypass surgery.
- (3) **Unstable angina pectoris**, who developed new cardiac symptoms or showed changing symptom patterns and positive ECG findings with normal biomarkers and admitted to coronary care unit (CCU).

#### Statistics

Baseline characteristics are presented as means  $\pm$  standard deviations (SD), median (interquartile range), and number (frequency) as appropriate. ANOVA and Kruskal-Wallis tests were used for comparison of means and medians, respectively. Chi-squared test was applied for comparison of frequencies.

The crude and age-standardized prevalence (95% confidence interval: CI) were calculated for all glycemic status, including NFG/NGT, iIFG, iIGT, IFG/IGT, NDM, and KDM. Regarding differences in the age distributions between the TLGS population from 2008 to 2011 and the Iranian census 2010 (supplementary Table 1), especially in the 30–39-year age-group and those aged  $\geq$  70 years, the age-standardized prevalence was reported, using the Iranian (Tehran province) census 2010.

We also examined the association of different glycemic status with the prevalence of CHD. Using logistic regression analyses, odds ratios (ORs) for this association were calculated in 3 levels of adjustment: 1) without adjustment (crude OR); 2) age and sex adjustment; 3) full adjustment (adjusted for age, sex, BMI, hypercholesterolemia, low HDL-C, hypertension, family history of premature CVD, and smoking status).

Statistical analyses were done using STATA version 14. *P*-values  $<$  0.05 were considered to be statistically significant.

#### Results

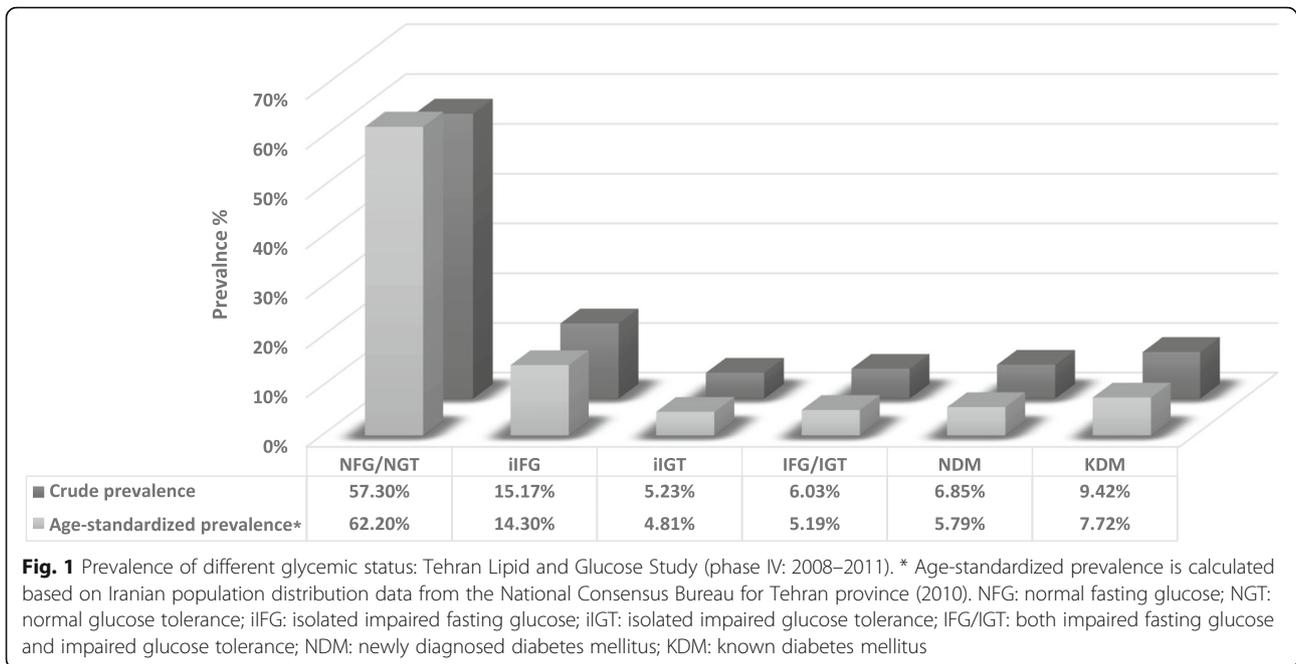
The study sample included 7718 participants (men = 3427) aged  $\geq$  30 years [mean age (SD) 50.1 (13.2) years]. Sex-specific baseline characteristics across glycemic categories are shown in Table 1. Generally, in comparison with the prediabetes and DM groups, participants with NFG/NGT had better cardiometabolic risk profiles, including age, BMI, waist to hip ratio, triglycerides, TC (only among women), SBP, and DBP. Furthermore, compared to the prediabetes and DM groups, participants with NFG/NGT had a better education status, lower prevalence of low physical activity (among women), and lower frequency of lipid-lowering and anti-hypertensive medications usage. Moreover, for most above-mentioned factors, prediabetic participants were ranked between NFG/NGT and DM groups.

Figure 1 shows the crude and age-standardized prevalence of different glycemic status. Among our total population, the age-standardized prevalence of iIFG, iIGT, IFG/IGT, NDM, and KDM were 14.30% (13.50–15.09), 4.81%

**Table 1** Baseline characteristics of participants across glyceemic categories: Tehran Lipid and Glucose Study (phase IV: 2008–2011)

	Men				Women				P-value*
	NFG/NGT	Prediabetes (IFG or IGT)	DM	P-value*	NFG/NGT	Prediabetes (IFG or IGT)	DM	P-value*	
Number of participants	1891	1000	536		2531	1040	720		
Continuous variables, Mean ± SD									
Age (year)	47.2 ± 12.8	53.6 ± 13.7	59.5 ± 12.6	< 0.001	45.2 ± 11.5	52.8 ± 11.9	58.7 ± 10.7	< 0.001	
BMI (kg/m <sup>2</sup> )	26.6 ± 4.0	28.0 ± 4.1	28.1 ± 4.1	< 0.001	28.5 ± 4.7	31.4 ± 18	31.3 ± 5.6	< 0.001	
Waist to hip ratio × 100	95.9 ± 6.3	98.6 ± 5.7	100.6 ± 5.7	< 0.001	90.7 ± 7.6	95.2 ± 7.4	99.2 ± 6.8	< 0.001	
SBP (mmHg)	117.1 ± 15.6	124.0 ± 18.3	130 ± 19.2	< 0.001	111.9 ± 16.2	122.2 ± 19.5	129.8 ± 22.0	< 0.001	
DBP (mmHg)	78.9 ± 10.5	80.8 ± 10.8	82.3 ± 11.0	< 0.001	74.7 ± 10.4	78.4 ± 10.8	79.8 ± 11.6	< 0.001	
Total cholesterol (mmol/L)	4.9 ± 0.9	5.0 ± 1.0	4.8 ± 1.0	< 0.001	5.0 ± 0.9	5.4 ± 1.0	5.3 ± 1.2	< 0.001	
HDL-C (mmol/L)	1.1 ± 0.2	1.1 ± 0.2	1.1 ± 0.2	0.109	1.3 ± 0.3	1.3 ± 0.3	1.2 ± 0.3	< 0.001	
Triglyceride (mmol/L)	1.5 (1.1–2.1)	1.7 (1.2–2.3)	1.8 (1.3–2.5)	< 0.001	1.2 (0.9–1.7)	1.6 (1.2–2.2)	1.9 (1.4–2.8)	< 0.001	
Categorical variables, n (%)									
Smoking status				< 0.001				0.382	
- Never	1055 (55.8%)	578 (57.8%)	303 (56.5%)		2403 (94.9%)	994 (95.6%)	690 (96.0%)		
- Past	345 (18.2%)	227 (22.7%)	140 (26.1%)		45 (1.8%)	16 (1.5%)	15 (2.1%)		
- Current	491 (26.0%)	195 (19.5%)	93 (17.4%)		83 (3.3%)	30 (2.9%)	14 (1.9%)		
Education level				< 0.001				< 0.001	
- < 6 years	283 (15.0%)	256 (25.6%)	198 (36.9%)		578 (22.8%)	448 (43.1%)	450 (62.6%)		
- 6–12 years	1042 (55.1%)	524 (52.4%)	255 (47.6%)		1405 (55.5%)	491 (47.2%)	232 (32.3%)		
- > 12 years	566 (29.9%)	220 (22.0%)	83 (15.5%)		548 (21.7%)	101 (9.7%)	37 (5.1%)		
Low physical activity, yes	723 (39.6%)	401 (43.3%)	203 (41.6%)	0.169	620 (24.6%)	272 (26.6%)	259 (37.6%)	< 0.001	
Family history of premature CVD, yes	78 (4.1%)	43 (4.3%)	28 (5.2%)	0.543	160 (6.3%)	81 (7.8%)	47 (6.5%)	0.276	
Lipid-lowering medication, yes	80 (4.2%)	96 (9.6%)	129 (24.1%)	< 0.001	129 (5.1%)	131 (12.6%)	262 (36.4%)	< 0.001	
Anti-hypertensive medication, yes	126 (6.7%)	121 (12.1%)	148 (27.6%)	< 0.001	194 (7.7%)	215 (20.7%)	317 (44.1%)	< 0.001	

Values are shown as Mean ± SD and number (%), for continuous and categorical variables, respectively; Triglycerides are given as median (interquartile range) due to skewed distribution  
 BMI Body mass index, SBP Systolic blood pressure, DBP Diastolic blood pressure, HDL-C High density lipoprotein cholesterol, CVD Cardiovascular disease, MFG Normal fasting glucose, IGT Impaired fasting glucose, DM Diabetes mellitus  
 \*The comparison p-value between groups was calculated using ANOVA test for normal continues variables, chi-square test for categorical variables and kruskal wallis test for skewed variables

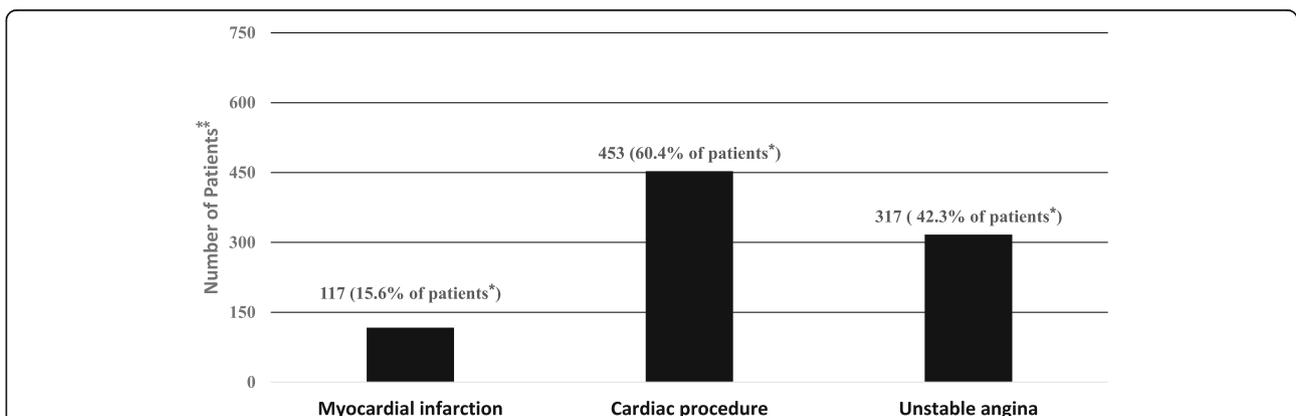


(4.32–5.29), 5.19% (4.71–5.67), 5.79% (5.29–6.28), and 7.72% (7.17–8.27), respectively.

Seven hundred fifty individuals were diagnosed as cases of CHD. The crude and age-standardized prevalence of CHD for the Tehranian population were 9.72% (95% CI: 9.06–10.38) and 7.71% (7.18–8.24), respectively. As is illustrated in Fig. 2, from a total of 750 patients with CHD in this study, 117 (15.6%), 453 (60.4%), and 317 (42.3%) individuals had a history of MI (definite and probable MI),

cardiac procedure, and unstable angina, respectively. It should be noted that the total number of different types of CHD is over 750 (100%), considering that patients might have more than one type of CHD.

The sex-specific prevalence of CHD across glycemc categories is shown in Table 2. Among the total age group, the crude prevalence of CHD was 11.61 (10.54–12.69) in men, which was significantly higher than the corresponding number among women. After age standardization, the



**Fig. 2** Number of patients across different types of coronary heart disease (CHD): Tehran Lipid and Glucose Study (phase IV: 2008–2011). \* Percentage of each type were calculated only among 750 patients with positive coronary heart disease in our data set. \*\* The total number of different type of CHD is over 750 (100%), considering that patients might have more than one type of CHD. CHD events included cases of: (1) Myocardial infarction (MI), included a) definite MI diagnosed by diagnostic electrocardiogram (ECG) and biomarkers (including CK, CK-MB, CK-MBm, troponin (cTn), and myoglobin), b) probable MI distinguished by positive ECG findings plus cardiac symptoms or signs and biomarkers showing negative or equivocal results. (2) Cardiac procedure, defined as a) angiography proven CHD with a result of  $\geq 50\%$  stenosis in at least one major coronary vessel, b) history of angioplasty or bypass surgery. (3) Unstable angina pectoris, who developed new cardiac symptoms or showed changing symptom patterns and positive ECG findings with normal biomarkers and admitted to coronary care unit (CCU)

**Table 2** Prevalence of coronary heart diseases across glycemic categories, by gender: Tehran Lipid and Glucose Study (phase IV: 2008–2011)

	Men			Women		
	Case/Total	Crude prevalence % (95% CI)	Age-standardized prevalence <sup>a</sup> % (95% CI)	Case/Total	Crude prevalence % (95% CI)	Age-standardized prevalence <sup>a</sup> % (95% CI)
<b>NFG/NGT</b>	139/1891	7.35 (6.17–8.53)	7.28 (6.06–8.49)	108/2531	4.27 (3.48–5.05)	5.71 (4.62–6.79)
<b>Prediabetes (IFG or IGT)</b>	126/1000	12.60 (10.54–14.66)	7.95 (6.55–9.36)	88/1040	8.46 (6.77–10.15)	6.62 (5.06–8.19)
<b>DM</b>	133/536	24.81 (21.15–28.47)	13.10 (9.83–16.38)	156/720	21.67 (18.65–24.68)	10.67 (8.90–12.44)
<b>Total</b>	398/3427	11.61 (10.54–12.69)	8.62 (7.81–9.44)	352/4291	8.20 (7.38–9.02)	7.19 (6.46–7.93)

NFG Normal fasting glucose, NGT Normal glucose tolerance, IFG Impaired fasting glucose, IGT Impaired glucose tolerance, DM Diabetes mellitus, CI Confidence interval

<sup>a</sup>Age-standardized prevalence is calculated based on Iranian population distribution data from the National Consensus Bureau for Tehran province (2010)

prevalence decreased to 8.62 (7.81–9.44) in men and 7.19 (6.46–7.93) in women. Among diabetic participants, the age-standardized prevalence of CHD was 13.10 (9.83–16.38) in men and 10.67 (8.90–12.44) in women, significantly higher than corresponding values for prediabetes and NFG/NGT groups.

The prevalence and ORs of CHD across NFG/NGT (as reference group), iIFG, iIGT, combined IFG and IGT, NDM, and KDM groups are presented in Table 3. KDM and NFG/NGT groups had the highest and lowest crude prevalence of CHD, respectively. Moreover, among prediabetic groups, the prevalence of CHD was tended to be more prominent in the combined IFG and IGT. The age-standardized prevalence of CHD was estimated to be 6.39 (5.59–7.19), 6.52 (5.23–7.82), 7.00 (4.73–9.27), 8.04 (5.40–10.67), 8.74 (7.08–10.40), and 14.26 (10.73–17.79) among NFG/NGT, iIFG, iIGT, combined IFG and IGT, NDM, and KDM groups, respectively. After adjustment for age and sex, CHD was more likely to be associated with combined IFG and IGT, NDM, and KDM groups. Furthermore, in the full-adjusted model, NDM and KDM status remained to be significantly associated with the presence

of CHD by ORs of 1.40 (1.06–1.86) for NDM and 1.91 (1.51–2.43) for KDM.

## Discussion

In this population-based study conducted in 2008–2011, 14.30, 4.81, 5.19, 5.79, and 7.72% of Tehranian residents were found to be iIFG, iIGT, IFG/IGT, NDM, and KDM, respectively. The age-standardized prevalence of CHD was about 7.7% among Tehranian residents. Generally, compared to women, the prevalence of CHD was found to be in higher ranges among men. Additionally, the age-standardized prevalence of CHD among diabetic participants was reported to be 13.1% in men and 10.7% in women. After age and sex adjustment, compared to the NFG/NGT group, the presence of IFG/IGT, NDM, and KDM were significantly associated with higher prevalence of CHD. The associations were significant for NDM and KDM, even in the full-adjusted model.

Based on the current study, about 40% of Tehranian adults were in prediabetes or diabetes status in 2008–2011, which was higher than our previous finding in 1990–2001 (about 30%) [19]. In 2005–2011, national

**Table 3** Prevalence and odds ratio of coronary heart diseases across glycemic categories: Tehran Lipid and Glucose Study (phase IV: 2008–2011)

	Case/Total	Crude prevalence % (95% CI)	Age-standardized prevalence <sup>a</sup> % (95% CI)	Crude odds ratio (95% CI)	Age and sex adjusted odds ratio (95% CI)	Full-adjusted odds ratio <sup>b</sup> (95% CI)
<b>NFG/NGT</b>	247/4422	5.59 (4.91–6.26)	6.39 (5.59–7.19)	Reference	Reference	Reference
<b>iIFG</b>	105/1171	8.97 (7.33–10.60)	6.52 (5.23–7.82)	1.67 (1.31–2.11)	1.13 (0.88–1.45)	0.99 (0.76–1.27)
<b>iIGT</b>	45/404	11.14 (8.07–14.21)	7.00 (4.73–9.27)	2.12 (1.52–2.96)	1.17 (0.82–1.67)	0.95 (0.66–1.38)
<b>IFG/IGT</b>	64/465	13.76 (10.63–16.90)	8.04 (5.40–10.67)	2.70 (2.01–3.62)	1.38 (1.01–1.89)	1.07 (0.78–1.47)
<b>NDM</b>	98/529	18.53 (15.21–21.84)	8.74 (7.08–10.40)	3.84 (2.98–4.96)	1.83 (1.40–2.41)	1.40 (1.06–1.86)
<b>KDM</b>	191/727	26.27 (23.07–29.47)	14.26 (10.73–17.79)	6.02 (4.89–7.43)	2.83 (2.26–3.55)	1.91 (1.51–2.43)
<b>Total</b>	750/7718	9.72 (9.06–10.38)	7.71 (7.18–8.24)	–	–	–

NFG Normal fasting glucose, NGT Normal glucose tolerance, iIFG Isolated impaired fasting glucose, iIGT Isolated impaired glucose tolerance, IFG/IGT Both impaired fasting glucose and impaired glucose tolerance, NDM Newly diagnosed diabetes mellitus, KDM Known diabetes mellitus, CI Confidence interval

<sup>a</sup>Age-standardized prevalence is calculated based on Iranian population distribution data from the National Consensus Bureau for Tehran province (2010)

<sup>b</sup>Adjusted for age, sex, BMI, hypercholesterolemia, low high-density lipoprotein cholesterol, hypertension, family history of premature cardiovascular disease, and smoking status

studies found a 35% increase in DM prevalence; the prevalence reached 11.4% in 2011. Importantly, in this period, DM awareness improved, and the nation-wide prevalence of NDM decreased from 45.7 to 24.7% [10]. Additionally, the researchers found that about 14.6% of Iranian adults (15.4% among urban residents) were in IFG status in 2011 [10], which was comparable to our study.

In our previous study, using self-reported history of CHD, Rose angina, and ECG-defined ischemia for CHD definition, a 21.8% prevalence of CHD was reported for Tehranian adults in 1999–2001 [3]. The differences between the prior study and the current study might be attributable to the following factors. Firstly, in our previous report, the silent ischemia and positive history of Rose angina were considered as cases of CHD; however, we did not consider these soft criteria of CHD in the current study. Secondly, for the history of CHD, in contrast to our original report, it was considered positive only when its hospital records were provided and then adjudicated by the outcome committee. Hence, in the current study, using the solid criteria for the definition of CHD led to underestimations of the prevalence of CHD.

It is important to note that due to different diagnostic criteria for CHD and different baseline characteristics of the population study, comparing our results with other population-based studies is somewhat difficult. Abbasi et al. reported that among an Iranian population aged over 20 years, the national prevalence of self-reported CHD was 5.3% (5.6% among urban residents) in 2011 [20]; their values for the prevalence of self-reported CHD were significantly lower than our reports. Compared to developed countries, data from the Quebec Integrated Chronic Disease Surveillance System (QICDSS) showed that the age-standardized prevalence of CHD (CHD death not included) was 7.7% among a Canadian adult population in 2009/2010 [21], which was comparable to ours. The American Heart Association (AHA) reported that the prevalence of total CHD was 6.7% among US adults aged  $\geq 20$  years (7.4% for men and 6.2% for women) [22]. For the United Kingdom (UK), data from the Quality and Outcomes Framework (QOF) indicated that the prevalence of CHD remained constant at about 3% in England and 4% in Scotland, Wales, and Northern Ireland between 2004/2005 and 2014/2015 [23]. Additionally, the prevalence of CHD varied from 2 to 4% in national studies of India [24]. Furthermore, in Saudi Arabia, as a Middle Eastern country, the age-standardized prevalence of CHD was reported to be 5.9 and 4.4% among male and female adults, respectively [25]. Generally, it seems that the estimated prevalence of CHD among the Tehranian population is higher than corresponding figures in US [22], UK [23], India [24] and, Saudi Arabia [25], an issue previously addressed in 2015 by Zhu et al. [26]. As we reported previously, modifiable risk factors (diabetes,

hypertension, smoking, and dyslipidemia) had population attributable fraction of 36.6 and 50.2% for incident CHD among the male and female Tehranian populations, respectively [4]. Other reasons that might justify the high prevalence of CHD among the Tehranian population are related to the impact of air pollution [27, 28] and stress [29], which are common in Tehran.

Focusing on diabetes status, from a national study on Iranian diabetic patients aged  $\geq 18$  years, it was reported that the crude prevalence of CHD was 25.1% for men and 23.2% for women in 2016 [30], which were comparable to our findings (24.81% for men and 21.67% for women). Among diabetic populations of other countries, the age-standardized prevalence of CHD was found to be 4.43 and 4.76% among Chinese men and women with T2DM, respectively [31]. Moreover, the prevalence of CHD among Thai patients with diabetes was 3.54% in 2013 [32]. Additionally, among Swedish diabetic patients aged 45–74 years, the crude prevalence of CHD was reported to be 24.9% for men and 18.0% for women [33]. In addition, a significant racial difference was reported in the prevalence of CHD between White and African diabetic patients in a hospital-based study [34]. It has been suggested that there is also a racial susceptibility for CHD among diabetic patients, which could make Iranian diabetic patients more prone to developing CHD, compared to Asian, African, and European ethnicities. Furthermore, although CVD risk factors among Iranian diabetic populations have been controlled to some degree, during recent years, most diabetic participants still have uncontrolled CVD risk factors [35], which could also lead to a high prevalence of CHD among our diabetic population.

In the current study, as expected, the significantly highest prevalence of CHD was found among participants with KDM. We also found that compared to NFG/NGT group, the presence of NDM status was associated with CHD [adjusted OR = 1.40 (1.06–1.86)]. We have previously reported that during a 7.6 years follow-up, Tehranian adults with NDM exhibited a CHD risk comparable to non-DM with a prior CHD [36]. In a cohort study, conducted on 271,174 participants with T2DM who were in the Swedish National Diabetes Register, it was shown that patients with T2DM who had five risk factors within the target range, appeared to have little or no excess risk of MI, in comparison with the general population [37]. Hence, we suppose that tight control of all CVD risk factors among the Iranian diabetic population should be considered in health policies to halt the increasing burden of CVD events. Regarding prediabetes status, a significant association was found between combined IFG/IGT and CHD in the sex- and age-adjusted model. Among a non-diabetic population, it was also reported that participants with combined IFG and IGT had a higher prevalence of significant CHD and higher

severity of disease in their angiographic results; however, there were no significant differences among subjects with NGT, i-IFG, and i-IGT [38].

Our study has its strengths in adjudicated cases of CHD by an outcome committee and the determination of CHD prevalence across different glycemic status, using the glucose challenge test. Several limitations need to be acknowledged. First, our study shows an optimistic picture of CHD prevalence among our population since the inclusion of subjects in an ongoing study can improve the level of attention paid to controlling their health risks (cohort effect). Therefore, the burden of CHD might be much higher in the context of the community. Second, this investigation was conducted among residents of Tehran as a metropolitan city. Hence, our results might not be generalizable to rural zones.

## Conclusion

The high prevalence of CHD, especially among diabetic populations, necessitates urgent behavioral intervention to be aimed at halting obesity tsunami [39], hypertension [40], and physical inactivity [41] among the Tehranian population, according to evidence-based guidelines for the clinical management of diabetic patients. Last but not least, the impact of environmental and psychosocial factors on CHD in Tehranians should be investigated in future studies.

## Supplementary information

**Supplementary information** accompanies this paper at <https://doi.org/10.1186/s12889-020-09595-4>.

**Additional file 1: Table S1.** Age distribution of the Iranian population for Tehran province (2010) and the sample population of Tehran lipid and glucose study (2008–2011).

## Abbreviations

CHD: Coronary Heart Disease; CVD: Cardiovascular Disease; T2DM: Type 2 Diabetes Mellitus; DM: Diabetes Mellitus; IFG: Impaired Fasting Glucose; IGT: Impaired Glucose Tolerance; MI: Myocardial Infarction; ECG: Electrocardiogram; MENA: Middle East and North Africa; TLGS: Tehran Lipid and Glucose Study; NCD: Non-communicable Disease; BP: Blood Pressure; BMI: Body Mass Index; TC: Total Cholesterol; HDL-C: High Density Lipoprotein Cholesterol; SBP: Systolic Blood Pressure; DBP: Diastolic Blood Pressure; FPG: Fasting Plasma Glucose; 2 h-PCPG: 2-h Post Challenge Plasma Glucose; NFG: Normal Fasting Glucose; NGT: Normal Glucose Tolerance; iIFG: Isolated Impaired Fasting Glucose; iIGT: Isolated Impaired Glucose Tolerance; IFG/IGT: Combined IFG and IGT; NDM: Newly Diagnosed Diabetes; KDM: Known Diabetes Mellitus; MAQ: Modifiable Activity Questionnaire; CCU: Coronary Care Unit; SD: Standard Deviations; CI: Confidence Interval; QICDSS: Quebec Integrated Chronic Disease Surveillance System; UK: United Kingdom; QOF: Quality and Outcomes Framework

## Acknowledgements

The authors wish to acknowledge Ms. Niloofar Shiva and Ms. Reyhane Hizomi Arani for critical editing of English grammar and syntax of the manuscript.

## Authors' contributions

Study conception and design: S.S.M and F.H.; Analysis and interpretation of data: M. H, D. P, and F.H.; Drafting of the manuscript: S.S.M, H. G, and F.H.;

Critical revision: S.S.M, A. G, H. G, F. A, and F.H. All authors read and approved the final manuscript.

## Funding

No funding from any source was obtained for this study.

## Availability of data and materials

The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

## Ethics approval and consent to participate

This study was passed by the Institutional Review Board (IRB) of the Research Institute for Endocrine Sciences (RIES), Shahid Beheshti University of Medical Sciences. Written informed consent was provided for all participants.

## Consent for publication

Not applicable.

## Competing interests

The authors declare that they have no competing interests.

## Author details

<sup>1</sup>Prevention of Metabolic Disorders Research Center, Research Institute for Endocrine Sciences, Shahid Beheshti University of Medical Sciences, P.O. Box: 19395-4763, No. 24, Parvaneh Street, Velenjak, Tehran, Iran. <sup>2</sup>Endocrine Research Center, Research Institute for Endocrine Sciences, Shahid Beheshti University of Medical Sciences, Tehran, Iran.

Received: 11 December 2019 Accepted: 22 September 2020

Published online: 06 October 2020

## References

- Roth GA, Abate D, Abate KH, Abay SM, Abbafati C, Abbasi N, et al. Global, regional, and national age-sex-specific mortality for 282 causes of death in 195 countries and territories, 1980–2017: a systematic analysis for the global burden of disease study 2017. *Lancet*. 2018;392(10159):1736–88.
- Mathers C, Stevens G, Hogan D, Mahanani WR, Ho J. Global and regional causes of death: patterns and trends, 2000–15; 2017.
- Hadaegh F, Harati H, Ghanbarian A, Azizi F. Prevalence of coronary heart disease among Tehran adults: Tehran lipid and glucose study. *East Mediterr Health J*. 2009;15(1):157–66.
- Khalili D, Sheikholeslami FH, Bakhtiyari M, Azizi F, Momenan AA, Hadaegh F. The incidence of coronary heart disease and the population attributable fraction of its risk factors in Tehran: a 10-year population-based cohort study. *PLoS One*. 2014;9(8):e105804.
- Sardarinia M, Akbarpour S, Lotfaliany M, Bagherzadeh-Khiabani F, Bozorgmanesh M, Sheikholeslami F, et al. Risk factors for incidence of cardiovascular diseases and all-cause mortality in a middle eastern population over a decade follow-up: Tehran lipid and glucose study. *PLoS One*. 2016;11(12):e0167623.
- Collaboration ERF. Diabetes mellitus, fasting blood glucose concentration, and risk of vascular disease: a collaborative meta-analysis of 102 prospective studies. *Lancet*. 2010;375(9733):2215–22.
- Naito R, Miyauchi K. Coronary artery disease and type 2 diabetes mellitus. *Int Heart J*. 2017;58(4):475–80.
- Azizi F, Ghanbarian A, Momenan AA, Hadaegh F, Mirmiran P, Hedayati M, et al. Prevention of non-communicable disease in a population in nutrition transition: Tehran lipid and glucose study phase II. *Trials*. 2009;10(1):5.
- Genuth S, Alberti K, Bennett P, Buse J, Defronzo R, Kahn R, et al. Follow-up report on the diagnosis of diabetes mellitus. The expert Committee on the diagnosis and classification of diabetes mellitus. *Diabetes Care*. 2003;26(11):3160–7.
- Esteghamati A, Etemad K, Koohpayehzadeh J, Abbasi M, Meysamie A, Noshad S, et al. Trends in the prevalence of diabetes and impaired fasting glucose in association with obesity in Iran: 2005–2011. *Diabetes Res Clin Pract*. 2014;103(2):319–27.
- Hadaegh F, Derakhshan A, Zafari N, Khalili D, Mirbolouk M, Saadat N, et al. Pre-diabetes tsunami: incidence rates and risk factors of pre-diabetes and its different phenotypes over 9 years of follow-up. *Diabet Med*. 2017;34(1):69–78.

12. Azizi F, Zadeh-Vakili A, Takyar M. Review of rationale, design, and initial findings: Tehran lipid and glucose study. *Int J Endocrinol Metab.* 2018;16(4 Suppl):e84777.
13. Chobanian AV, Bakris GL, Black HR, Cushman WC, Green LA, Izzo JL Jr, et al. The seventh report of the joint national committee on prevention, detection, evaluation, and treatment of high blood pressure: the JNC 7 report. *Jama.* 2003;289(19):2560–71.
14. Committee IR. Guidelines for data processing and analysis of the International Physical Activity Questionnaire (IPAQ)-short and long forms. <http://www.ipaq.ki.se/scoring.pdf>. 2005.
15. Kabootari M, Hasheminiya M, Azizi F, Mirbolouk M, Hadaegh F. Change in glucose intolerance status and risk of incident cardiovascular disease: Tehran lipid and glucose study. *Cardiovasc Diabetol.* 2020;19:1–11.
16. Luepker RV, Apple FS, Christenson RH, Crow RS, Fortmann SP, Goff D, et al. Case definitions for acute coronary heart disease in epidemiology and clinical research studies: a statement from the AHA Council on Epidemiology and Prevention; AHA Statistics Committee; World Heart Federation Council on Epidemiology and Prevention; the European Society of Cardiology Working Group on Epidemiology and Prevention; Centers for Disease Control and Prevention; and the National Heart, Lung, and Blood Institute. *Circulation.* 2003;108(20):2543–9.
17. Moran KE, Ommerborn MJ, Blackshear CT, Sims M, Clark CR. Financial stress and risk of coronary heart disease in the Jackson heart study. *Am J Prev Med.* 2019;56(2):224–31.
18. Keku E, Rosamond W, Taylor HA Jr, Garrison R, Wyatt SB, Richard M, et al. Cardiovascular disease event classification in the Jackson heart study: methods and procedures. *Ethn Dis.* 2005;15(4 Suppl 6):S6–62.
19. Hadaegh F, Bozorgmanesh MR, Ghasemi A, Harati H, Saadat N, Azizi F. High prevalence of undiagnosed diabetes and abnormal glucose tolerance in the Iranian urban population: Tehran lipid and glucose study. *BMC Public Health.* 2008;8(1):176.
20. Abbasi M, Neishaboury M, Koohpayehzadeh J, Etemad K, Meysamie A, Asgari F, et al. National Prevalence of Self-Reported Coronary Heart Disease and Chronic Stable Angina Pectoris: Factor Analysis of the Underlying Cardiometabolic Risk Factors in the SuRFNCD-2011. *Glob Heart.* 2018;13(2):73–82.e71.
21. Blais C, Rochette L. Trends in prevalence, incidence and mortality of diagnosed and silent coronary heart disease in Quebec. *Health Promot Chronic Dis Prev Can.* 2015;35(10):184–93.
22. Benjamin EJ, Muntner P, Alonso A, Bittencourt MS, Callaway CW, Carson AP, et al. Heart disease and stroke Statistics-2019 update: a report from the American Heart Association. *Circulation.* 2019;139(10):e56–e528.
23. Bhatnagar P, Wickramasinghe K, Wilkins E, Townsend N. Trends in the epidemiology of cardiovascular disease in the UK. *Heart.* 2016;102(24):1945–52.
24. Gupta R, Mohan I, Narula J. Trends in coronary heart disease epidemiology in India. *Ann Glob Health.* 2016;82(2):307–15.
25. Al-Nozha M, Arafah M, Al-Mazrou Y, Al-Maatouq M, Khan N, Khalil M, et al. Coronary artery disease in Saudi Arabia. *Saudi Med J.* 2004;25(9):1165–71.
26. Zhu K-F, Wang Y-M, Zhu J-Z, Zhou Q-Y, Wang N-F. National prevalence of coronary heart disease and its relationship with human development index: a systematic review. *Eur J Prev Cardiol.* 2016;23(5):530–43.
27. Khajavi A, Khalili D, Azizi F, Hadaegh F. Impact of temperature and air pollution on cardiovascular disease and death in Iran: a 15-year follow-up of Tehran lipid and glucose study. *Sci Total Environ.* 2019;661:243–50.
28. Luo C, Zhu X, Yao C, Hou L, Zhang J, Cao J, et al. Short-term exposure to particulate air pollution and risk of myocardial infarction: a systematic review and meta-analysis. *Environ Sci Pollut Res Int.* 2015;22(19):14651–62.
29. Steptoe A, Kivimaki M. Stress and cardiovascular disease: an update on current knowledge. *Annu Rev Public Health.* 2013;34:337–54.
30. Esteghamati A, Larijani B, Aghajani MH, Ghaemi F, Kermanchi J, Shahrami A, et al. Diabetes in Iran: prospective analysis from first Nationwide diabetes report of National Program for prevention and control of diabetes (NPPCD-2016). *Sci Rep.* 2017;7(1):13461.
31. Lyu Y, Luo Y, Li C, Guo X, Lu J, Wu H, et al. Regional differences in the prevalence of coronary heart disease and stroke in patients with type 2 diabetes in China. *J Clin Endocrinol Metab.* 2018;103(9):3319–30.
32. Sakboonyarat B, Rangsin R. Prevalence and associated factors of ischemic heart disease (IHD) among patients with diabetes mellitus: a nation-wide, cross-sectional survey. *BMC Cardiovasc Disord.* 2018;18(1):151.
33. Wirehn AB, Ostgren CJ, Carstensen JM. Age and gender differences in the impact of diabetes on the prevalence of ischemic heart disease: a population-based register study. *Diabetes Res Clin Pract.* 2008;79(3):497–502.
34. Kalk WJ, Joffe BI. Differences in coronary heart disease prevalence and risk factors in African and white patients with type 2 diabetes. *Diabetes Res Clin Pract.* 2007;77(1):107–12.
35. Jahangiri-Noudeh Y, Akbarpour S, Lotfaliany M, Zafari N, Khalili D, Tohidi M, et al. Trends in cardiovascular disease risk factors in people with and without diabetes mellitus: a middle eastern cohort study. *PLoS One.* 2014;9(12):e112639.
36. Hadaegh F, Fahimfar N, Khalili D, Sheikholeslami F, Azizi F. New and known type 2 diabetes as coronary heart disease equivalent: results from 7.6 year follow up in a Middle East population. *Cardiovasc Diabetol.* 2010;9(1):84.
37. Rawshani A, Rawshani A, Franzén S, Sattar N, Eliasson B, Svensson AM, et al. Risk factors, mortality, and cardiovascular outcomes in patients with type 2 diabetes. *N Engl J Med.* 2018;379(7):633–44.
38. Yan Q, Gu WQ, Hong J, Zhang YF, Su YX, Gui MH, et al. Coronary angiographic studies of impaired glucose regulation and coronary artery disease in Chinese nondiabetic subjects. *Endocrine.* 2009;36(3):457–63.
39. Barzin M, Valizadeh M, Serahati S, Mahdavi M, Azizi F, Hosseinpah F. Overweight and obesity: findings from 20 years of the Tehran lipid and glucose study. *Int J Endocrinol Metab.* 2018;16(4 Suppl):e84778.
40. Asgari S, Moazzeni SS, Azizi F, Abdi H, Khalili D, Hakemi MS, et al. Sex-specific incidence rates and risk factors for hypertension during 13 years of follow-up: the Tehran lipid and glucose study. *Glob Heart.* 2020;15(1):29.
41. Esteghamati A, Khalilzadeh O, Rashidi A, Kamgar M, Meysamie A, Abbasi M. Physical activity in Iran: results of the third national surveillance of risk factors of non-communicable diseases (SuRFNCD-2007). *J Phys Act Health.* 2011;8(1):27–35.

## Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

**Ready to submit your research? Choose BMC and benefit from:**

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

**At BMC, research is always in progress.**

Learn more [biomedcentral.com/submissions](https://biomedcentral.com/submissions)

