The higher adherence to a healthy lifestyle score is associated with a decreased risk of type 2 diabetes in Iranian adults

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Abstract

Background: The combined roles of lifestyle factors in the risk of type 2 diabetes (T2D) are not fully investigated. In the present study, we aimed to assess the relationship between a healthy lifestyle score (HLS) and the risk of T2D in Tehranian adults.

Methods: A total of 3859 individuals without T2D were recruited from participants of the Tehran Lipid and Glucose Study (2009–2011) who were followed up for a mean period of 6 years. A food frequency questionnaire was used to collect individuals’ dietary intakes at baseline. HLS scores was calculated based on three pre-defined methods with focusing on 4 lifestyle factors including, no current smoking, no obesity, high physical activity, and greater adherence to the healthy diet (determined using the alternate healthy eating index-2010 (AHEI-2010), modified French Programme National Nutrition Santé-Guideline Score(mPNNS-GS), and healthy diet pattern score(HDP)).

Results: Mean ± SD age of participants (44.4% men) was 41.1 ± 12.3 years. After 6-year follow-up of study, 295 (7.6%) new cases of T2D were reported. Based on the age and sex-adjusted model, an inverse association was observed between the higher score of HLS-AHEI-2010 (OR = 0.24; 95%CI: 0.10–0.60), HLS-mPNNS-GS (OR = 0.28; 95%CI: 0.15–0.50), and HLS-HDP (OR = 0.39; 95%CI: 0.24–0.64) and the risk of T2D (P for trend < 0.05). Also, the fully-adjusted model showed that after controlling the effects of various confounders, this inverse association between the higher score of HLS-AHEI-2010 (OR = 0.25; 95%CI: 0.10–0.61, P for trend: 0.001), HLS-mPNNS-GS (OR = 0.29; 95%CI: 0.15–0.55, P for trend: 0.001), and HLS-HDP (OR = 0.36; 95%CI: 0.22–0.61, P for trend < 0.001) and risk of T2D was remained significant.

Conclusion: The results of this study suggested that higher score of HLS, characterized by no smoking, normal body weight, vigorous physical activity, and healthy diet, is related to decreased risk of T2D incidence.

Keywords: Healthy lifestyle score, Dietary pattern, Obesity, Physical activity, Smoking, Type 2 diabetes, Adult

Background

Type 2 diabetes (T2D) is a major health problem with complex metabolic conditions, which is considered the leading preventable risk factor for the development of cardiovascular disease (CVD) and all-cause mortality worldwide [1, 2]. Indeed, T2D is a complex and polygenic disease characterized by chronic hyperglycemia as a consequence of impaired insulin action and secretion, insulin resistance, unfavorable inflammatory condition,
and metabolic susceptibility [3]. In 2017, 8.5% of the global adult population (425 million people) had T2D, and it has been anticipated to be 629 million by 2045 [1]. Recent investigation has reported that the T2D incidence rate in the Iranian population was 36.3 per 1000 person-years, with more than 800,000 new cases per year [4]. Also, it is estimated that in the year 2030, nearly 9.2 million Iranians will be affected by T2D [5]. Lack of proper management of T2D with the increasing duration of this disease threatens people’s lives due to various microvascular complications such as nephropathy, retinopathy, and neuropathy, and macrovascular complications in the form of peripheral vascular disease and coronary artery disease and lead to premature death [1, 6].

The interplay of genetic predisposition and metabolic factors are the main drivers of the global epidemic of T2DM [7]. Along with population aging, modifiable factors, including unhealthy dietary pattern, obesity, physical inactivity, and smoking, are recognized as prominent risk factors that can contribute to the development of T2D [1, 2]. In previous reports, lifestyle risk modification has been suggested as a strong strategy in the prevention of chronic diseases such as T2D [8]; so that it has been suggested that variable levels of each modifiable lifestyle factor, including smoking [9], physical activity [10], normal body weight [11], and adherence to a healthy dietary pattern [12] can be associated with the risk of T2D. Also, some studies have recently investigated the combined role of lifestyle factors, including healthy diet pattern, not smoking, avoiding obesity, and regular exercise, as a single variable, called healthy lifestyle score (HLS), with risk of chronic diseases and mortality [13–16]. Based on the results of previous reports, adherence to HLS is related to the promotion of overall health during aging [13], decreasing the risk of coronary disease [14], and reducing mortality in type 2 diabetic subjects [15, 16]. However, to the best of our knowledge, there is no study on the association of combined lifestyle factors with the risk of T2D incidence in the adult population.

Considering convincing evidence on the high incidence of T2D among Iranian adults [17], the possible beneficial combined role of healthy lifestyle factors as an HLS in the prevention of some chronic diseases or their related mortality, and also due to the lack of data on the association between HLS and the risk of T2D, in the present study, we aimed to assess the relationship between HLS and the risk of T2D incident among Iranian adult population.

**Methods**

**Study population**

The present study was conducted within the framework of the Tehran Lipid and Glucose Study (TLGS), a population-based cohort study that was performed to determine the risk factors for chronic diseases among a representative urban population of Tehran, including 15,005 participants aged ≥ 3 years [18]. The first phase of TLGS (a cross-sectional survey) was initiated in March 1999, and data collection, conducted prospectively at three years intervals, is ongoing; the details of the TLGS have been reported previously [18]. In the fourth survey of the TLGS (2009–2011), of 12,823 participants, 7956 subjects, aged 3–75 years, were randomly selected to be assessed for dietary intakes.

For the present study, after excluding individuals aged < 19 years (n = 1585), 6371 participants, aged ≥ 19 years, with complete data in the fourth phase of the TLGS (baseline examination), were selected. Individuals who under- or over-reported dietary intakes (< 800 kcal/day or > 4200 kcal/day, respectively) or who were on specific diets (n = 470), those with prevalent cancer (n = 12), cardiovascular diseases (n = 49), and pregnant and lactating women (n = 116), were excluded. The individuals with baseline T2D were also excluded (n = 553); some of whom fell into more than one category (n = 142). Next, 5313 healthy individuals (free of T2D) were followed until the sixth examination, for a mean period of 6 years from the baseline phase (fourth survey). Finally, after excluding the participants who left the study (n = 1454), final analyses were performed on the data of 3859 adults (Fig. 1).

**Physical activity assessment**

Data on physical activity was collected using a modifiable activity questionnaire (MAQ), which was previously modified and validated among Iranian adults [19]. Individuals were asked to report and identify the frequency and time spent on activities of light, moderate, hard, and very hard intensity over the past 12 months, based on a list of usual activities of daily life; the ‘participants’ physical activity levels were reported as metabolic equivalent hours per week (MET-h/wk).

**Demographic, anthropometric, and lifestyle assessment**

Details of the TLGS data collection approaches and measurements have been previously reported [18]. The participants were asked to report their information on demographic characteristics, drug use, medical history, and cigarette smoking using a pretest standard questionnaire at baseline of current study. Body weight was measured and recorded to the nearest 100 g using a digital scale while participants wearing minimal clothes and were without shoes. The height of individuals was determined to the nearest 0.5 cm using a stadiometer, without shoes that shoulders were in a normal position. Body mass index (BMI) was considered as a individual’s weight in kilograms divided by the square of their height.
in meters. The systolic blood pressure (SBP) and diastolic blood pressure (DBP) were determined using a standardized mercury sphygmomanometer with an accuracy of two mmHg based on standard protocols which was previously explained [18].

**Biochemical Measurements**

A 10 ml blood sample was taken after 12–14 h of overnight fasting to measure biochemical variables including fasting plasma glucose (FPG), triglycerides (TGs), total cholesterol (TC), and high-density lipoprotein-cholesterol (HDL-C) based on the standard protocol that has been previously described [20]. All blood analyses were performed at the TLGS research laboratory using the Selectra 2 auto-analyzer (Vital Scientific, Spankeren, The Netherlands) to analyze the samples. FPG, TG, TC, and HDL-C were measured using enzymatic colorimetric method and all analyses were done using commercial kits (Pars Azmoon Inc., Tehran, Iran). The Friedewald formula was used to determine low-density lipoprotein-cholesterol (LDL-C) from the serum TC, TG, and HDL-C concentrations.

**Dietary assessment**

Dietary intakes of individuals were collected using a valid and reliable semi-quantitative food frequency at baseline [21]. The trained nutritionist asked participants to designate their consumption frequency for each food
item during the previous year daily, weekly, or monthly; portion sizes of consumed foods, reported in household measures, were then converted to grams. Since the Iranian Food Composition Table (FCT) is incomplete and has limited data on the nutrient content of raw foods and beverages, the United States Department of Agriculture (USDA) FCT was used. The Iranian FCT was used for national foods not listed in the USDA FCT.

**Determination of healthy lifestyle scores**
The score of healthy lifestyle factors for participants was determined using three healthy lifestyle indices, including HLS-­‐ alternate healthy eating index (AHEI)-2010 [15], HLS- modified French Programme National Nutrition Santé-Guideline Score (mPNNS-GS) [13], and HLS- healthy dietary pattern (HDP) [14]. All three indices consist of four factors, including smoking (yes/no), physical activity (active/ inactive), obesity (yes/no), and diet quality, which in each of these scoring procedures, the quality of the diet has been assessed in different ways.

For computing of HLS-AHEI-2010 [15], individuals were divided into two groups based on each lifestyle factor as follows: non-obese (BMI < 25 kg/m²) vs. overweight/obese (BMI ≥ 25 kg/m²); non-current smoking vs. current smoking; physically active (≥ 22.5 MET/h/w) vs. physically low active/sedentary (< 22.5 MET/h/w). A score of 1 was assigned to participants who were normal weight. Also, a score of 1 was considered for subjects who were non-smoked. Furthermore, the physically active participants (≥ 22.5 MET/h/w) received a score of 1. The quality of diet in participants has been assessed based on the AHEI-2010 diet score, which was defined in detail previously [22]. Individuals were divided into quintiles based on the AHEI-2010 score; individuals in the upper two quintiles of the AHEI-2010 score were given a score of 1 (as a higher adherence to a healthy diet). The quality of diet in participants has been assessed based on the AHEI-2010 diet score, which was defined in detail previously [22]. Individuals were divided into quintiles based on the AHEI-2010 score; individuals in the upper two quintiles of the AHEI-2010 score were given a score of 1 (as a higher adherence to a healthy diet). The points for factors were then summed to create the overall healthy lifestyle score, which ranged from 0 (lowest adherence to HLS-AHEI-2010) to 4 (highest adherence to HLS-AHEI-2010).

The HLS-HDP index was also determined according to the strategic goals of the American Heart Association (AHA) using four important lifestyle variables (as binary components), including BMI, physical activity, smoking, diet quality [14]. We assigned a score of 1 to participants who are not obese (BMI < 30 kg/m²), are non-current smoking, and are physically active (≥ 66.5 MET/h/w). The quality of dietary pattern in participants was assessed by focusing on high consumption of fruits, vegetables, fish, whole grains, nuts, and dairy products and low consumption of processed meats, unprocessed red meats, refined grains, trans fats, sugar-sweetened beverages, and sodium. The overall healthy dietary pattern score ranged from 0 to 12. Then, a score of 1 was assigned to individuals if they had a healthy dietary pattern (diet quality score ≥ 6). Finally, we summed the points for each of the factors to create the overall healthy lifestyle score, which ranged from 0 (lowest adherence to HLS-HDP) to 4 (highest adherence to HLS-HDP).

In the present study, lifestyle score (named as HLS- mPNNS-GS) was also determined based on the method reported in the Atallah et al. study [13]. Like the methods mentioned above, HLS-mPNNS-GS were calculated based on four main lifestyle factors: no obesity, vigorous exercise, no smoking, and a healthy diet. We assigned a score of 1 for participants if they are normal weight (BMI = 18.5–24.99 kg/m²) and non-current smoking. Also, participants were divided into quartiles based on physical activity level; individuals in the upper two quartiles of physical activity level were given a score of 1 (as an active people). The diet quality score in participants was calculated based on the French guideline-based dietary index (mPNNS-GS). We have determined the overall healthy diet score based on 10 food groups, including fruit and vegetables, bread and legumes, whole grains, dairy products, red and processed meat, fish and seafood, added fats, fats with the vegetable origin, sugary food, and Sweetened beverage, salt, and MET score. The diet quality score ranged from 0 to 12.5. We assigned a score of 1 to participants if they had a healthy dietary pattern (diet quality score ≥ 6.5). Finally, we summed the points of the factors to create the overall healthy lifestyle score, which ranged from 0 (lowest adherence to HLS- mPNNS-GS) to 4 (highest adherence to HLS- mPNNS-GS).

Since alcoholic drinks intakes such as wine and liquor are unusual among the Iranian people due to religious considerations and were not reported in the TLGS study, we did not include them in computing the three indices mentioned above. As we did not have any food items as low-energy beverages and cream soup in our FFQ, we excluded them from the calculation.

**Definitions**
Diabetes was defined based on the criteria of the American Diabetes Association (ADA) as FPG ≥ 126 mg/dl or 2-h post 75 g glucose load ≥ 200 mg/dl or taking oral hypoglycemic medication [23].

**Statistical analysis**
All analyses of our study were done using the Statistical Package for Social Sciences (Version 20.0; SPSS, Chicago, IL). Kolmogorov-Smirnov test was used to check the variables’ normality. Baseline characteristics of the individuals were presented as the mean ± SD or median (25–75 interquartile) for continuous variables and percentages for categorical variables. We used Chi-square
and independent two-sample t-tests to compare qualitative and quantitative variables, respectively, between individuals with and without T2D. Individuals were also classified according to quartiles of HLS (HLS-AHEI-2010, HLS-mPNNS-GS, and HLS-HDS) cut-off points; Chi-square test and linear regression analyses were used to test the trends of categorical and continuous variables across quartiles of HLS, respectively. Multivariable logistic regression models were used to predict the risk of 6-year incident T2D (as the dependent variable) across quartiles of HLS-AHEI-2010, HLS-mPNNS-GS, and HLS-HDS (as independent variables). Multivariable logistic regression models were used to predict the risk of 6-year incident T2D (as the dependent variable) across quartiles of HLS-AHEI-2010, HLS-mPNNS-GS, and HLS-HDS (as independent variables). The findings of Table 1 showed that the % of employed individuals, smoking, men, and the mean of physical activity, energy inakes, and macronutrients did not differ significantly between the T2D and non-T2D groups.

Participants’ characteristics are also presented across the quartiles of HLS-AHEI-2010, HLS-mPNNS-GS, and HLS-HDP in Tables 2, 3, and 4, respectively. Based on the results of Table 2, participants in the highest quartile of the HLS-AHEI-2010 score significantly to be more female, younger, non-smoked, had higher levels of physical activity, academic education, and HDL-C, and had a lower level of BMI, TGs, LDL-C, SBP, DBP, and lower % incidence of T2D (after 6-years of follow-up) compared to those in the lowest quartile of HLS-AHEI-2010 (P < 0.05). Also, the intakes of vegetables, whole grains, fruits, and nuts and legumes significantly increased across quartile of the HLS-AHEI-2010 (P < 0.05), whereas dietary intakes of red and processed meat, sweetened beverages, trans fatty acid, and sodium were significantly decreased across HLS-AHEI-2010 score quartile (P < 0.05).

According to Table 3, individuals in the highest quartile of the HLS-mPNNS-GS score significantly to be more

### Table 1 Baseline characteristics of participants according to the development of the type 2 diabetes

|                          | Type 2 diabetes (n = 295) | Non-Type 2 diabetes (n = 3564) | P-value |
|--------------------------|--------------------------|-------------------------------|---------|
| **Baseline demographic and biochemical data** |                          |                               |         |
| Age (year)               | 47.4 ± 11.5              | 40.6 ± 12.3                   | <0.0001 |
| Male, %                  | 43.7                     | 44.5                          | 0.423   |
| Body mass index (kg/m²)  | 30.8 ± 5.3               | 27.0 ± 4.4                    | <0.0001 |
| Physical Activity (MET/h/w) | 65.1 (35.7–93.8)     | 67.5 (35.7–99.1)              | 0.660   |
| Academic education (graduated), % | 20.5               | 29.9                          | <0.0001 |
| Employed, %              | 84.1                     | 84.7                          | 0.419   |
| Smoker, %                | 11.9                     | 10.8                          | 0.310   |
| Fasting plasma glucose (mg/dl) | 102.0 ± 10.3          | 92.2 ± 7.9                    | <0.0001 |
| Triglycerides (mg/dl)    | 169 (126–230)            | 113 (80–162)                  | <0.0001 |
| Low density lipoprotein-Cholesterol (mg/dl) | 117.7 ± 32.6           | 111.6 ± 32.5                  | 0.003   |
| High density lipoprotein-Cholesterol (mg/dl) | 44.4 ± 10.5           | 47.9 ± 11.6                   | <0.0001 |
| Systolic blood pressure (mmHg) | 123.4 ± 18.2           | 111.9 ± 15.1                  | 0.001   |
| Diastolic blood pressure (mmHg) | 80.9 ± 12.4           | 74.9 ± 10.7                   | <0.0001 |
| **Dietary intakes**      |                          |                               |         |
| Energy (kcal/day)        | 2359 ± 701               | 2399 ± 709                    | 0.351   |
| Carbohydrate (g/day)     | 348 ± 113                | 354 ± 119                     | 0.394   |
| Protein (g/day)          | 87.3 ± 28.1              | 89.2 ± 48.8                   | 0.508   |
| Fat (g/day)              | 77.5 ± 28.1              | 80.3 ± 48.2                   | 0.315   |

Data are presented as mean ± standard deviation for normally distributed variables and median (25–75 interquartile range) for skewed variables, and percent for categorical variables.
female, younger, high-active, more employed, lower-smoked, had higher level of academic education, and had lower BMI, TGs, FPG, LDL-C, SBP, and DBP levels (at baseline) and lower % incidence of T2D (after 6-years of follow-up) compared to those in the lowest quartile of this score \( (P<0.001) \). Also, based on our results, the intakes of vegetables and fruits, bread and legumes, whole grains, seafood, and vegetable oils significantly increased across quartiles of HLS-HDP score, whereas the intake of red meat significantly decreased across quartiles of HLS-HDP score \( (P<0.05) \), whereas dietary intakes of refined grains, sodium, and sweetened beverage were significantly reduced across quartiles of HLS-HDP score \( (P<0.05) \).

We determined the odds of T2D across the quartiles of HLS-AHEI-2010, HLS-mPNNS-GS, and HLS-HDP and showed main results in Fig. 2 (A-C). Based on the age and sex-adjusted model, the odds of T2D was reduced across quartiles of HLS-AHEI-2010 (OR = 0.24; 95%CI: 0.10–0.60, \( P \) for trend = 0.001), HLS-mPNNS-GS (OR = 0.28; 95%CI: 0.15–0.50, \( P \) for trend = 0.002), and HLS-HDP (OR = 0.39; 95%CI: 0.24–0.64, \( P \) for trend < 0.001). Also, findings of the multivariable-adjusted model indicated that after adjusting age, gender,
Table 3  General characteristics and HLS-mPNNS-GS components of participants according to quartiles of HLS-mPNNS-GS

| Variable                        | Quartiles of HLS-mPNNS-GS | P for trend |
|---------------------------------|---------------------------|-------------|
|                                | Q1 (n = 519)              | Q2 (n = 1044) | Q3 (n = 1757) | Q4 (n = 539) |
| Mean score                      | 1.00±0.00                 | 2.00±0.00    | 3.00±0.00    | 4.00±0.00    | <0.001      |
| Age (year)                      | 44.2±13.2                 | 41.5±12.5    | 41.3±11.6    | 36.8±12.3    | <0.001      |
| Male, %                         | 68.9                      | 60.1         | 51.5         | 47.5         | <0.001      |
| Academic education (graduated),%| 23.1                      | 27.5         | 29.3         | 37.8         | <0.001      |
| Employed, %                     | 76.2                      | 82.4         | 87.9         | 86.2         | <0.001      |
| Fasting plasma glucose (mg/dl)  | 93.8±9.1                  | 93.3±8.9     | 93.2±8.3     | 90.6±7.1     | <0.001      |
| Triglycerides (mg/dl)           | 130.0 (92.0 – 185.0)      | 121.0 (84.0 – 173.0) | 120.0 (87.0 – 171.0) | 90.0 (66.0 – 128.0) | <0.001      |
| Low density lipoprotein-Cholesterol (mg/dl) | 115.8±33.1          | 111.9±33.4    | 113.4±32.0   | 104.1±30.4   | <0.001      |
| High density lipoprotein-Cholesterol (mg/dl) | 47.8±12.4            | 47.5±11.8     | 47.0±11.0    | 49.9±11.2    | 0.064       |
| Systolic blood pressure (mmHg)  | 115.2±17.2                | 112.7±15.5   | 113.7±15.3   | 107.2±13.8   | <0.001      |
| Diastolic blood pressure (mmHg) | 76.4±11.5                | 75.1±11.0    | 76.2±10.9    | 71.8±9.4     | <0.001      |
| Diabetes incidence (%)          | 9.5                       | 9.2          | 7.9          | 2.2          | <0.001      |
| Current smoker (%)              | 19.0                      | 22.2         | 5.1          | 0.0          | <0.001      |
| Physical activity (MET/h/w)     | 26.0 (12.9 – 37.6)        | 38.7 (19.9 – 71.4) | 82.3 (61.5 – 112.6) | 89.3 (67.0 – 128.0) | <0.001      |
| Body mass index (kg/m²)         | 29.3±4.5                  | 27.5±4.5     | 27.9±4.4     | 22.7±1.6     | <0.001      |
| Diet quality score (based on mPNNS-GS) | 6.05±0.69             | 7.01±1.08    | 7.96±0.88    | 8.08±0.80    | <0.001      |

Data are presented as mean ± standard deviation for normally distributed variables and median (25–75 interquartile range) for skewed variables, and percent for categorical variables.

* Percentage of diabetes incidence across quartiles of HLS-mPNNS-GS was determined at the end of follow-up of study.

energy intake, employment status, education, and marital status, the risk of T2D was decreased across the quartiles of HLS-AHEI-2010 (OR=0.25; 95%CI:0.10–0.61, P for trend=0.001), HLS-mPNNS-GS (OR=0.29; 95%CI: 0.15–0.55, P for trend=0.001), and HLS-HDP (OR=0.36; 95%CI: 0.22–0.61, P for trend <0.001).

Discussion

The current study has provided notable results on the association of HLS and the risk of T2D in the framework of longitudinal population-based study after a 6-year follow-up. Our findings reported that a higher score of HLS, assessed by three indices including HLS-AHEI-2010, HLS-mPNNS-GS, HLS-HDP, is related to decreased risk of T2D independent of confounding factors.

Although strong reviews have focused on the association of each lifestyle factor with the risk of T2D incident separately, their findings have shown a significant association of healthy dietary pattern scores such as DASH diet, Mediterranean diet, and healthy eating index [12], vigorous physical activity [10], smoking [9], and elevated BMI with risk of T2D [11], there is no study on the combined role of lifestyle factors score as a single variable with risk of T2D. Recently, some studies have investigated the combined role of HLS in health status or chronic diseases risk, such as the risk of coronary disease [14], healthy aging [13], and mortality [15, 16]. Interestingly, our study results are in agreement with the results of two previous studies that observed an inverse association between the HLS and total mortality in adults with type 2 diabetes patients [15, 16]. Also, in the Atallah et al. study, a one-point increase in the HLS was associated with an 11% higher probability of healthy aging [13]. Furthermore, the Khera et al. study has reported that in
Table 4 General characteristics and HLS-HDP components of participants according to quartiles of HLS-HDP

| Variables | Quartiles of HLS-HDP | P for trend |
|-----------|----------------------|------------|
|           | Q1 (n = 348) | Q2 (n = 1396) | Q3 (n = 1578) | Q4 (n = 537) |
| Mean score | 1.00±0.00 | 2.00±0.00 | 3.00±0.00 | 4.00±0.00 | <0.001 |
| Age (year) | 44.3±11.7 | 40.9±12.4 | 40.7±12.3 | 40.7±12.0 | <0.001 |
| Male, % | 57.3 | 62.8 | 53.0 | 43.4 | <0.001 |
| Academic education (graduated), % | 19.4 | 26.5 | 31.6 | 35.3 | <0.001 |
| Employed, % | 80.9 | 82.8 | 86.1 | 87.3 | 0.006 |
| Fasting plasma glucose (mg/dl) | 95.2±9.2 | 93.0±8.7 | 92.7±8.3 | 92.1±7.8 | <0.001 |
| Triglycerides (mg/dl) | 133.0 (98.0 – 185.0) | 118.50 (82.0 – 169.7) | 113.0 (80.7 – 165.0) | 113.0 (78.0 – 160.0) | 0.002 |
| Low density lipoprotein-Cholesterol (mg/dl) | 115.0±34.1 | 112.5±32.3 | 111.1±32.6 | 111.7±31.5 | 0.099 |
| High density lipoprotein-Cholesterol (mg/dl) | 45.3±10.7 | 47.8±11.3 | 48.0±11.7 | 48.7±11.9 | 0.019 |
| Systolic blood pressure (mmHg) | 116.7±15.2 | 112.7±16.1 | 112.1±15.6 | 112.1±14.5 | <0.001 |
| Diastolic blood pressure (mmHg) | 77.7±10.7 | 75.1±11.2 | 75.0±10.8 | 75.2±10.30 | 0.011 |
| Diabetes incidence (%) | 13.8 | 9.0 | 5.9 | 5.2 | <0.001 |
| Current smoker (%) | 36.0 | 14.8 | 5.6 | 0.0 | <0.001 |
| Physical activity (MET/h/w) | 39.6 (22.9 – 56.1) | 47.3 (23.8 – 71.4) | 78.1 (47.0 – 107.1) | 98.2 (81.1 – 129.1) | <0.001 |
| Body mass index (kg/m²) | 31.8±7.7 | 27.8±4.9 | 26.3±4.0 | 25.8±2.6 | <0.001 |
| Diet quality score (overall healthy dietary pattern) | 4.46±0.81 | 4.82±1.16 | 5.51±1.26 | 6.51±0.72 | <0.001 |

Overall healthy dietary pattern components

| Diet quality score (overall healthy dietary pattern) | Fruit (serving/day) | Vegetables (serving/day) | Dairy products (serving/day) | Whole grains (serving/day) | Fish (serving/day) | Nuts (serving/day) | Red meat (serving/day) | Processed meat (serving/day) | Refined grains (serving/day) | Sweetened beverage (serving/week) | Sodium (mg/day) | Trans fatty acid (g/day) |
|------------------------------------------------------|---------------------|--------------------------|-----------------------------|---------------------------|-------------------|-------------------|----------------------|--------------------------|-----------------------------|-------------------------------|----------------|------------------------|
| Q1 (n = 348) | 1.91 (1.18 – 3.29) | 2.47 (1.71 – 3.52) | 1.42 (0.86 – 2.08) | 2.77 (1.42 – 5.05) | 0.21 (0.11 – 0.42) | 0.13 (0.06 – 0.24) | 0.66 (0.40 – 0.96) | 0.08 (0.02 – 0.15) | 4.02 (2.97 – 6.10) | 1.12 (0.42 – 2.80) | 3655 (3002 – 4748) | 2.35 (1.14 – 4.72) |
| Q2 (n = 1396) | 2.33 (1.38 – 3.98) | 2.53 (1.69 – 3.79) | 1.56 (1.05 – 2.23) | 2.90 (1.56 – 5.17) | 0.21 (0.11 – 0.43) | 0.14 (0.07 – 0.30) | 0.64 (0.38 – 1.02) | 0.08 (0.02 – 0.15) | 4.17 (2.89 – 6.54) | 0.91 (0.35 – 2.10) | 3445 (2719 – 4302) | 2.21 (1.19 – 4.58) |
| Q3 (n = 1578) | 3.11 (1.73 – 5.07) | 3.19 (2.04 – 4.38) | 1.70 (1.11 – 2.52) | 3.84 (2.06 – 6.28) | 0.23 (0.12 – 0.48) | 0.16 (0.07 – 0.33) | 0.63 (0.37 – 1.02) | 0.07 (0.01 – 0.17) | 3.88 (2.80 – 5.98) | 0.77 (0.28 – 1.96) | 3287 (2558 – 4155) | 2.39 (1.25 – 4.71) |
| Q4 (n = 537) | 4.49 (3.17 – 6.44) | 3.66 (2.85 – 4.88) | 2.28 (1.45 – 3.09) | 4.41 (3.03 – 7.30) | 0.29 (0.14 – 0.54) | 0.21 (0.10 – 0.43) | 0.61 (0.37 – 1.03) | 0.07 (0.01 – 0.16) | 3.65 (2.37 – 5.48) | 0.70 (0.21 1.82) | 3336 (2659 – 4118) | 2.35 (1.21 – 4.51) |

Data are presented as mean ± standard deviation for normally distributed variables and median (25–75 interquartile range) for skewed variables, and percent for categorical variables

* Percentage of diabetes incidence across quartiles of HLS-HDP was determined at the end of follow-up of study

individuals with high genetic susceptibility, a healthy lifestyle was associated with a 46% lower risk of coronary disease events than an unfavorable lifestyle [14]. For investigation of the relationship between HLS and T2D risk, we have determined HLS in participants using three indices, including HLS-AHEI-2010, HLS-mPNNS-GS, and HLS-HDP. The rational for using these three indices was as follows: first, the scoring procedure of these indices were different in diet quality assessment or considering the cut point for the classification of participants based on BMI level; e.g. Patel et al. had used the AHEI 2010 to assess the diet quality [15], however, the Atallah et al. had used the PNNS-GS method to determine food quality [13]. Also, in Patel et al. and Atallah et al. studies, subjects were divided into two groups of individuals with normal weight and those with overweight and obesity, but in Khera et al. study, individuals with normal weight and overweight were classified into one group and received the same score [14]. Second, each of these HLS scoring methods has been used and validated in different societies that their population probably was different in general characteristics such as lifestyle and dietary pattern with other population such as people in Middle East and North Africa (MENA) region. Another important point was that in the previous studies, each of these indices has been used to determine the HLS in relation to mortality or various nutrition-related chronic diseases.
risk, but none of previous studies has been focused on risk of the T2D. Therefore, in our study, we assessed the relationship of healthy lifestyle, determined by three HLS indices, with the T2D risk in the Iranian population. Based on the results of all three methods, we were able to show a remarkable association between a higher HLS score and reduced the T2D risk, which indicates that all three methods used have an acceptable and similar ability in the prediction of people’s healthy lifestyle score and its possible relationship with the incidence of T2D.

The results of the current study are further in agreement with the results of previous studies that have summarized evidence on the independent role of individual lifestyle factors in the development of T2D [9–12]; one of the most important components of a lifestyle affecting the incidence of T2D is the dietary pattern of individuals. A systematic review has shown that the higher adherence

![Fig. 2 A-C](image-url)
to healthy dietary patterns such as the Mediterranean diet and the DASH diet, characterized by higher consumption of fruits, vegetables, legumes, nuts, low-fat dairy, whole grain, seafood, dietary fiber, unsaturated fats, and antioxidant nutrients and lower consumption of animal fat, red and processed meat, sweets and desserts, are associated with decreased risk of T2D [12]. Also, the greater adherence to alternative healthy eating index focused on a high intake of vegetables, fruits, nuts, legumes, whole grain, PUFA, and lower intakes of red meat, sweet beverages, trans fatty acids, and sodium, was linked to decreased risk of T2D as well as lower prevalence of general obesity [12].

Several studies have also focused on the independent role of other lifestyle factors, including smoking, physical activity, and obesity, in the pathogenesis of T2D. In a meta-analysis of cohort studies, higher physical activity was related to a substantially lower risk of T2D [10]. It has been suggested that the vigorous physical activity level has a protective role in reducing the risk of T2D via the beneficial effect on insulin sensitivity, improving glucose metabolism via insulin-receptor up-regulation in muscle, and increased insulin and glucose delivery to muscle, increasing total antioxidant capacity and reducing central adiposity through negative energy balance [24, 25]. Also, the Akter et al. study has reported that cigarette smoking is linked with an increased risk of T2D [9]. Smoking leads to inadequate insulin secretion and insulin resistance [26, 27] via various underlying effects such as hormonal imbalance, inflammation, oxidative stress, central adiposity, and endothelial dysfunction [9, 28]. Also, smoking harms the metabolism of nutrients, β-cells dysfunction, and up-regulating inflammatory markers such as C-reactive protein (CRP), which can be related to increased risk of cardiometabolic abnormalities [29, 30]. Furthermore, the elevated body weight and central adiposity is another main underlying risk factor for T2D [11]; because obesity has an important role in various metabolic pathways and in producing several potential risk factors such as increased levels of the chronic inflammatory response (increased inflammatory markers such as interleukin-6 and tumor necrosis factor-alpha), increased risk of impaired hepatic free fatty acids and glucose metabolism, hyperinsulinemia, β-cell dysfunction, insulin resistance, and dyslipidemia which all of them are mostly associated with higher risk of T2D [31, 32].

Given the important individual role of each lifestyle factor in predicting the risk of T2D, we examined the relationship between the combined role of these factors and the risk of T2D in individuals who followed a healthy lifestyle. Our results provide strong evidence that following combined healthy behaviors in the framework of a healthy lifestyle, which are characterized by a healthy dietary pattern rich in fruits and vegetables, legumes and grains, lower consumption of red and processed meat and sweetened beverage, and having regular meals, high physical activity, and abstinence from smoking, can have a remarkable effect size in reducing the risk of T2D incident. Indeed, despite the differences between the previous studies and the current study concerning definitions of HLS (components and scoring), the robust findings reinforced and confirmed the assumption that combined healthy behaviors may positively impact health promotion and decrease risk of chronic diseases such as T2D.

Our study has several important strengths. This study is the first population-based prospective study that assessed the combined role of healthy lifestyle factors in prediction of T2D risk. Also, the current study has a large sample size and relatively long-term follow-up duration. Using valid and reliable food-frequency and physical activity questionnaires for collection the participants’ data on dietary intakes and physical activity levels is another strength of this study. Despite these strengths, some limitations of current study should be mentioned. Similar to epidemiological studies, recall bias and measurement error are inherent limitation of using the FFQ in the present study. Also, although various confounding factors were controlled in multivariable logistic regression model, we cannot ruled out the effects of residual or unmeasured confounders.

Conclusion
In conclusion, our findings have shown that higher adherence to HLS, characterized by a healthy diet, vigorous physical activity, not smoking, and normal BMI, is strongly associated with a decreased risk of T2D in Iranian adults. Further observational studies are suggested to assess the combined protective role of HLS in the development of T2D among other populations.

Abbreviations

| Abbreviation | Description |
|--------------|-------------|
| AHEI-2010 | Alternate healthy eating index-2010, AHA: American Heart Association; BMI: Body mass index; CI: Confidence interval; CVD: Cardiovascular disease; DBP: Diastolic blood pressure; FPG: Fasting plasma glucose; FCT: Food composition table; FFQ: Food frequency questionnaire; HDL-C: High-density lipoprotein cholesterol; HDFP: Healthy diet pattern; HLS: Healthy lifestyle score; LDL-C: Low-density lipoprotein cholesterol; MAQ: Modified Activity Questionnaire; MET-h/wk: Metabolic equivalent hours per week; mPNNS-GS: Modified French Programme National Nutrition Santé-Guideline Score; SBP: Systolic blood pressure; TC: Total cholesterol; TG: Triglyceride; TLGS: Tehran Lipid and Glucose Study; T2D: Type 2 diabetes; USDA: United States Department of Agriculture; WC: Waist circumference. |

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Authors' contributions
HF and GA contributed to the conception, design, and statistical analysis. HF, FT, and EM contributed to data collection and manuscript drafting. PM and FA supervised the study. All authors approved the final version of the manuscript.

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Availability of data and materials
The datasets analysed in the current study are available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate
Informed written consents were obtained from participants. All procedures performed in studies involving human participants adhered to the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. The study protocol was approved by the research council of the Research Institute for Endocrine Sciences, Shahid Beheshti University of Medical Sciences (Ethical number: IR.SBMU.ENDOCRINE.REC.1399.152).

Consent for publication
Not applicable.

Competing of interests
The authors declared there is no conflict of interest.

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References
1. Giovaci D, Fan W, Wong ND. Epidemiology of diabetes Mellitus and cardiovascular disease. Curr Cardiol Rep. 2019;21(4):21.
2. Zheng Y, Ley SH, Hu FB. Global astrology and epidemiology of type 2 diabetes mellitus and its complications. Nat Rev Endocrinol. 2018;14(2):88–98.
3. Ling C. Epigenetic regulation of insulin action and secretion – role in the pathogenesis of type 2 diabetes. J Intern Med. 2020;288(2):153–67.
4. Ramezankhani A, Harati H, Bozorgmanesh M, Tohidi M, Khalili D, Azizi F, et al. Diabetes Mellitus: findings from 20 years of the Tehran Lipid and Glucose Study. Int J Endocrinol Metab. 2018;16(4 Suppl):e84784-e.
5. Mirzaei M, Rahmaninan M, Mirzaei M, Nadjarzadeh A, Dehghanitafti AA. Epidemiology of diabetes mellitus, pre-diabetes, undiagnosed and uncontrolled diabetes in Central Iran: results from Yazd health study. BMC Public Health. 2020;20(1):166.
6. Beckman JA, Creager MA, Libby P. Diabetes and atherosclerosis: epidemiology, pathophysiology, and management. JAMA. 2002;287(19):2570–81.
7. Fan W. Epidemiology in diabetes mellitus and cardiovascular disease. Cardiovasc Endocrinol. 2017;6(1):18–16.
8. Patja K, Joussilhati P, Hu G, Vallet T, Qiao Q, Tuomilehto J. Effects of smoking, obesity and physical activity on the risk of type 2 diabetes in middle-aged Finnish men and women. J Intern Med. 2005;258(4):356–62.
9. Akter S, Goto A, Mizoue T. Smoking and the risk of type 2 diabetes in Japan: a systematic review and meta-analysis. J Epidemiol. 2017;27(12):553–61.
10. Smith AD, Crippa A, Woodcock J, Brage S. Physical activity and incident type 2 diabetes mellitus: a systematic review and dose-response meta-analysis of prospective cohort studies. Diabetologia. 2016;59(12):2527–45.
11. Krishnan S, Rosenberg L, Djoussé L, Cupples LA, Palmer JR. Overall and central obesity and risk of type 2 diabetes in US. Black Women Obesity. 2007;15(7):1860–6.
12. Jannasch F, Kroger J, Schulze MB. Dietary patterns and type 2 diabetes: a systematic literature review and meta-analysis of prospective studies. J Nutr. 2017;147(6):1174–82.
13. Atalafah N, Adjjabade M, Lelong H, Herberg S, Galan P, Assmann KE, et al. How healthy lifestyle factors at midlife relate to healthy aging. Nutrients. 2018;10(7):854.
14. Khera AV, Emdin CA, Drake I, Natarajan P, Bick AG, Cook NR, et al. Genetic risk, adherence to a healthy lifestyle, and coronary disease. N Engl J Med. 2016;375(24):2349–58.
15. Patel YR, Gadrijo TV, Gazzano JM, Djoussé L. Adherence to healthy lifestyle factors and risk of death in men with diabetes mellitus: the physicians’ health study. Clin Nutr. 2018;37(1):139–43.
16. NÖTHLINGS U, FORD ES, KROGER J, BOEING H. Lifestyle factors and mortality among adults with diabetes: findings from the European prospective investigation into cancer and nutrition–potsdam study*. J Diabetes. 2010;2(2):112–7.
17. Sali S, Farhadnejad H, Asghari G, Teymoori F, Mirmiran P, Djazayeri A, et al. Animal based low carbohydrate diet is associated with increased risk of type 2 diabetes in Tehranian adults. Diabetol Metab Syndr. 2020;12(1):87.
18. Azizi F, Ghanbarian A, Mommen 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.
19. Mommen AA, Delshad M, Sarbazi N, REZAEI GN, Ghanbarian A, AZIZI F. Reliability and validity of the Modifiable Activity Questionnaire (MAQ) in an Iranian urban adult population. Arch Iran Med. 2012;15(5):279–82.
20. Farhadnejad H, Asghari G, Mirmiran P, Djazayeri A, et al. Animal based low carbohydrate diet is associated with increased risk of type 2 diabetes in Tehranian adults. Diabetol Metab Syndr. 2020;12(1):87.
21. Azizi F, Ghanbarian A, Mommen 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. Public Health Nutr. 2009;13(5):654–62.
22. Chiueve SE, Fung TT, Rimm EB, Hu FB, McCullough ML, Wang M, et al. Alternative dietary indices both strongly predict risk of chronic disease. J Nutr. 2012;142(6):1009–18.
23. Idiapathic B, Endocrinopathies D. Report of the expert committee on the diagnosis and classification of diabetes mellitus. Diabetes Care. 2003;26:55–20.
24. Weinstein AR, Sesso HD, Lee IM, Cook NR, Manson JE, Buring JE, et al. Relationship of physical activity vs body mass index with type 2 diabetes in women. JAMA. 2004;292(10):1188–94.
25. Hamasaki H. Daily physical activity and type 2 diabetes: a review. World J Diabetes. 2016;7(12):243–51.
26. Attvall S, Fowelin J, Lager I, Von Schenck H, Smith U. Smoking induces insulin resistance—a potential link with the insulin resistance syndrome. J Intern Med. 1993;233(4):327–32.
27. Facchini FS, Hollenbeck CB, Jeppesen J, Chen YD, Reaven GM. Insulin resistance and cigarette smoking. Lancet. 1992;339(8802):1128–30.
28. van der Vaart H, Postma DS, Timens W, Ten Hacken NH. Acute effects of cigarette smoke on inflammation and oxidative stress: a review. Thorax. 2005;60(10):824–30.
29. Stadler M, Tomm L, Storka A, Wolz M, Peric S, Bieglmayer C, et al. Effects of smoking cessation on b-cell function, insulin sensitivity, body weight, and appetite. Eur J Endocrinol. 2014;170(2):219–27.
30. Kolovou GD, Kolovou V, Mavrogeni S. Cigarette smoking/cessation and insulin resistance and type 2 diabetes. Nature. 2004;498(7498):713–21.
31. Stadler M, Tomm L, Storka A, Wolz M, Peric S, Bieglmayer C, et al. Effects of smoking cessation on b-cell function, insulin sensitivity, body weight, and appetite. Eur J Endocrinol. 2014;170(2):219–27.
32. Kolovou GD, Kolovou V, Mavrogeni S. Cigarette smoking/cessation and insulin resistance and type 2 diabetes. Nature. 2004;498(7498):713–21.
32. Bastard J-P, Maachi M, Lagathu C, Kim MJ, Caron M, Vidal H, et al. Recent advances in the relationship between obesity, inflammation, and insulin resistance. Eur Cytokine Netw. 2006;17(1):4–12.

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