

RESEARCH ARTICLE

Joint Association of Screen Time and Physical Activity with Cardiometabolic Risk Factors in a National Sample of Iranian Adolescents: The CASPIANIII Study

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Citation: Heshmat R, Qorbani M, Shahr Babaki AE, Djalalinia S, Ataei-Jafari A, Motlagh ME, et al. (2016) Joint Association of Screen Time and Physical Activity with Cardiometabolic Risk Factors in a National Sample of Iranian Adolescents: The CASPIANIII Study. PLoS ONE 11(5): e0154502. doi:10.1371/journal.pone.0154502

Editor: Fernando Guerrero-Romero, Mexican Social Security Institute, MEXICO

Received: January 5, 2015

Accepted: April 14, 2016

Published: May 11, 2016

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Data Availability Statement: Due to legal restrictions our data are available upon request. Readers may contact Dr. Mostafa Qorbani and Dr. Kelishadi (corresponding authors of present article) to request the data.

Funding: The authors have no support or funding to report.

Competing Interests: The authors have declared that no competing interests exist.

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Abstract

Metabolic syndrome (MetS) and its contributing factors are considered important health problems in the pediatric age group. This study was designed to assess the joint association of ST and PA with cardiometabolic risk factors among Iranian adolescents. A representative sample of 5625 (50.2% boys) school students with a mean age of 14.73 (SD: 2.41) were selected through multistage random cluster sampling method from urban and rural areas of 27 provinces in Iran. ST and PA were assessed by self-administered validated questionnaires. Anthropometric measures (height, weight and waist circumference (WC)) and MetS components (abdominal obesity, elevated blood pressure (BP), low high-density lipoprotein cholesterol (HDL-C), elevated triglycerides (TG) and high fasting blood sugar (FBG)) were measured according to standardized protocols. MetS was defined according to the Adult Treatment Panel III criteria modified for the pediatric age group. Moreover, elevated total cholesterol (TC), elevated low-density lipoprotein cholesterol (LDL-C), and generalized obesity were considered as other cardiometabolic risk factors. Students with high ST levels had significantly higher body mass index z-score (BMI z-score), WC, TG, LDL-C, and BP as well as lower HDL-C level; whereas those with high PA levels had significantly higher HDL-C levels as well as lower BMI z-score, TC, and BP. Adolescents with low PA/ high ST levels

had significantly higher BMI, WC, LDL-C levels, as well as higher SBP and DBP compared to their other counterparts. In Multivariate model, joint effect of low PA/ high ST (compared to the high PA/low ST group) increased the odds of overweight, abdominal obesity and low HDL-C and decreased the odds of elevated TC. The findings of this study showed that joint association of high ST and low PA have direct association with abdominal obesity, overweight and low HDL-C and indirect association with elevated TC.

Introduction

Metabolic syndrome (MetS) is defined as a cluster of risk factors for various non-communicable diseases, namely diabetes mellitus (DM) and cardiovascular diseases (CVD) [1].

Pediatric MetS is no more limited to industrial societies; recent studies have demonstrated that MetS is rapidly increasing in Iranian adolescents [2–9] similar to many other developing and Western countries [10–17]; it has been documented even in those with normal weight [18]. Some studies have explored the notion of persistence of MetS and other cardiometabolic risk factors into adulthood [1,7,19]. Notably, this combination of risk factors, rather than each of them, contributes to additional risks, beyond the sum of risks attributed to each individual risk factors [20].

CVDs and their major risk factors become highly prevalent worldwide and it is of special concern in the Middle East [21]. The evidences of epidemiological studies show that geographical differences and ethnicity are two main factors affecting the prevalence of cardiometabolic risk factors in different populations [22,23]. The most frequent metabolic risk factors in Iranian pediatric population are low levels of HDL-C, hypertriglyceridemia and overweight, respectively [24].

Sedentary lifestyle including low physical activity (PA) and prolonged screen time (ST) are considered as one of the major health problems in the pediatric population of developing and developed countries [25–27]. Previous studies have reported a considerably high prevalence of low PA and prolonged ST in Iranian adolescents [28,29].

Various studies have examined the associations of ST, i.e. leisure time spent in front of television (TV) or computer, and PA with cardiometabolic risk factors [30–33]. It is shown that changes in life style including; reducing routine daily activities of children may contribute to lower PA in this age group [34]. Some controversies exist about the associations of PA and ST with some components of MetS [35]. Some studies found that ST and PA have weak correlations [36–38]; whereas some other studies reported that lower energy expenditure and PA might mostly contribute to higher prevalence of metabolic abnormalities [35, 39]. A study by Mitchell *et al.* argued that sedentary behavior has neglected effect on obesity; and suggested that combined effects PA and ST should be studied [40].

Therefore, the present study was designed to examine the associations of the ST, PA and their joint association with MetS and cardiometabolic risk factors in Iranian adolescents.

Material and Methods

A national representative sample of third survey of the school-based surveillance system entitled “Childhood and Adolescence Surveillance and Prevention of Adult Non-communicable Disease” (CASPIAN) study (2009–2010). The aim and methods of mentioned study is described previously [41].

Participants were 5625 students, aged 10–18 years, who were selected via multistage random cluster sampling method from urban and rural areas of 27 provinces in Iran. Data for some variables was missing. Based on the protocol of the study, the stratification of eligible schools was followed using the information bank of the Ministry of Health and Medical Education. In the next step sampled schools were selected randomly. In later step, in each of selected schools, the sampling of students was randomly. In data gathering phase, all processes of examinations with calibrated instruments and recording of information in validated checklists were designed and conducted under the standard protocol by trained health care professional teams.

The study was approved by the ethical committee of Tehran and Isfahan University of Medical Sciences and other relevant national regulatory organizations (Ministry of Health and Ministry of Education). Participation in the study was voluntarily. Sampling and examinations were begun after complete introducing of project and explanation of the study's protocols for students and their parents. Oral assent was obtained from participants and written informed consent from their parents or legal guardians.

Clinical and Laboratory Measurements

Height and weight were measured, according to standardized protocols, without shoes and with light clothing to the nearest 0.1 unit of measure (cm for height and kg for weight). Body mass index (BMI) was calculated from weight and height [BMI = weight (kg) /height (m²)] [4, 42]. Waist circumference (WC) was measured over skin, midway between the lower border of the rib margin and the iliac crest at the end of normal expiration, to the nearest 0.1 cm. Both of WC and height were measured using a non-elastic tape. Abdominal obesity was defined as waist to height ratio (WHtR) more than 0.5 [43].

Systolic blood pressure (SBP) and diastolic blood pressure (DBP) were measured, using a standardized mercury sphygmomanometer, on the right arm after a 15-minute rest in a sitting position. The first and fifth Korotkoff sounds were respectively recorded as systolic and diastolic blood pressure. The mean of the two measurements was considered as the subject's blood pressure.

For each of participants, a blood sample was drawn between 7:00 and 9:00 AM after 12 to 14 hours overnight fasting. Blood samples were delivered to the laboratory on the same day. Fasting blood glucose (FBG), total cholesterol (TC), triglycerides (TG), high density lipoprotein-cholesterol (HDL-C), and low-density lipoprotein cholesterol (LDL-C) were measured enzymatically by auto-analyzers. HDL-C was determined after dextran sulfate-magnesium chloride precipitation of non-HDL-C [44].

As the highest quality of data was critical to the success of our multi-center data gathering, the different levels of quality assurance and control were exactly considered by Data and Safety Monitoring Board (DSMB) of the project.

Demographic Information

Demographic information was completed for all participant students in the sampled classes of the selected schools through an interview with parents or child. Family based characteristics including: family history of chronic diseases (hypertension, dyslipidemia, diabetes, and obesity), parental level of education (the highest total years of schooling), possessing a family private car and type of home (rented/owned), dietary behaviors, PA, and sedentary lifestyle.

Definition of Terms

Cardiometabolic risk factors. If students had at least three of the following criteria according to Adult Treatment Panel III (ATP III) criteria modified for children and

adolescents, were considered as having MetS. The modified criteria for children and adolescents have been defined as follow: Abdominal obesity as waist to height ratio (WHtR) more than 0.5; Elevated BP: either systolic or diastolic BP \geq 90th percentile for age, sex and height; Low HDL-C: HDL-C \leq 40 mg/dl (except in boys 15–19 years old that the cut off was $<$ 45 mg/dl); High TG: TG \geq 100 mg/dl) was taken as the 90th percentile value for age; High FBG: FBG levels of \geq 100 mg/dl [45]. Five criteria of MetS and TC, LDL-C, and general obesity were included in this study as cardiometabolic risk factors. High TC and LDL-C was defined according to the recent recommendation by the American Heart Association (TC \geq 200 mg/dl, LDL-C $>$ 110mg/dl [46, 47].

The overweight and general obesity definition provided by the Centers of Disease Control and Prevention (CDC) and the percentiles computed in the population studied were used for the classification of the adolescents as overweight (85–94th percentile) and obese ($>$ 95th percentile) [46].

Socioeconomic status (SES). The method and variables, which was used for calculating SES was approved previously in the Progress in the International Reading Literacy Study (PIRLS) [48] Using principle component analysis (PCA) method variables including parental education, parents' job, possessing private car, school type (public/private), and having personal computer in home were summarized in one main component. This main component was categorized into tertiles. The first tertile was defined as a low SES, second tertile as an intermediate and third tertile as a high.

ST and PA. In this study, the ST behavior of the adolescents was assessed through the questionnaire that asked the child to report the average number of hours per day they spent watching TV/VCDs, personal computer (PC), or electronic games (EG) in week days and weekends. Based on that total cumulative spent time for ST was estimated. For the analysis of correlates of ST, according to the international ST recommendations, ST was categorized into two groups; less than 2 hours per day (Low), and 2 hours per day or more (High) [49, 50].

For leisure time PA, the information of past week was collected. Participants reported the weekly frequency of their leisure time PA outside the school. Having PA, considered as at least 30 minutes duration of daily exercises might lead to heavy sweating or large increases in breathing or heart rate. Therefore, we categorized weekly PA habits through available response choices as follow; none, 1–2 days, 3–6 days, and every day. For statistical analysis, PA was categorized into low (0–2 days /week) and high (3–7 days/week) levels [51].

The joint associations of PA and ST were considered based on following possible mixed conditions: Low PA & Low ST, Low PA & High ST, High PA & Low ST, and High PA & High ST.

Statistical Analyses

Quantitative variables are expressed as mean (standard deviation (SD)) and qualitative variables as number (percentage). Differences between means were investigated by T-test or ANOVA test (followed by Tukey's post-hoc tests) and for categorized variables; the Pearson Chi-square test used to compare the percentages. Logistic regression analyses were used to evaluate the joint association of PA and ST categories with odds of cardiometabolic risk factors. In Model I the joint association of PA/ST categories with cardiometabolic risk factors were assessed without adjustment. In Model II the association was adjusted for age and sex and in Model III, additionally family history and SES were adjusted in the model. In Model IV in addition to all variables in the Model III, BMI was adjusted in all abnormalities except obesity. In all models "high PA & low ST" group was considered as reference group because this combination according to the international ST recommendations has lowest risk of cardiometabolic

risk factors compared to other combinations. Results of logistic regression are presented as odds ratio (OR) and 95% confidence interval (CI). In all analysis design of sampling (cluster sampling) were considered. Data was analyzed using "survey data method". All statistical analyses were performed using programs available in the STATA version 10. A *p-value* of less than 0.05 was considered as statistically significant.

Results

The participants of this national study were 5625 students (50.2% boys, 49.8% girls) with a mean age of 14.73 (SD: 2.41). A comparison of baseline characteristics is presented in [Table 1](#). It shows that boys had higher BMI z-score ($p < 0.01$), whereas girls had greater height, weight and WC measurements ($p < 0.001$). Considering the combination of times spent on watching TV and working with the computer, most of the adolescents had low ST levels (54.1% of all participants) which was statistically significant between both sexes (56.8% of boys and 51.3% of girls) ($p < 0.001$); boys and girls spent equal amounts of time in front of TV, while for working with computer, boys spent more time than girls which was statistically significant ($p < 0.001$). Distribution of PA levels among boys and girls pointed out a significant difference among those PA level ($p < 0.001$); the majority of participants of both genders and the overall sample had low PA levels (88.4% of boys, 81.3% of girls and 84.9% of all), while more girls were highly active. Regarding joint association of ST and PA there was a statistically significant difference between boys and girls. In this regard, the combination of low ST and low PA in boys was significantly more prevalent than in girls ($p < 0.001$) ([Table 1](#)).

[Table 2](#) presents the associations between cardiometabolic risk factors with ST, PA and joint association of PA and ST categories. Students with high ST levels had significantly higher BMI z-score, WC, TG, LDL and BP and lower HDL level; those with high PA levels had significantly lower BMI z-score, TC and BP and higher HDL levels. Regarding the joint association of PA and ST, we observed that mean of all cardiometabolic risk factors except WHtR, FBG and TG were significantly different across PA/ST levels. Results of post hoc test show that students with low PA and high ST levels had significantly higher BMI z-score, SBP and DBP, compared to the high PA and low ST levels; these individuals had the lowest serum HDL-C levels among all participants ($p < 0.05$).

[Table 3](#) shows the prevalence of cardiometabolic risk factors according to ST and PA levels categories and their joint association. Individuals with high ST levels had a higher prevalence of having elevated serum TG and LDL levels, being overweight or having lower serum HDL levels; although there was a significant difference between the prevalence of different MetS components in low and high ST levels, prevalence of MetS was similar in the two categories. Considering PA levels, participants with high PA levels had a lower prevalence of low HDL levels (27.2% versus 36.4%) or elevated serum TC levels (5.3% versus 8.7%). When accounting for the joint association of PA and ST, participants with low PA/ high ST levels had the highest prevalence of low serum HDL levels; those with high PA/ low ST levels had the highest prevalence of elevated serum TC levels ($p < 0.01$) ([Table 3](#)).

[Table 4](#) illustrates the comparison of joint effects of PA/ ST levels on cardiometabolic risk factors, against being highly physically active and having low ST levels (high PA/ low ST), using different logistic regression models. Only those with low PA levels had significantly different risks of having certain cardiometabolic risk factors; individuals with low PA/ high ST levels had their odds of being overweight and having abdominal obesity and low HDL-C increased by 73%, 48% and 40–50%, respectively. Their odds of having elevated serum TC levels were decreased 39–42%, based on the regression model. Participants with low PA/ low ST levels had just decreased odds of having elevated serum TC levels (50–52%) ([Table 4](#)).

Table 1. Demographic characteristics and anthropometric measures according to sexes: The CASPIAN III study.

	Boys	Girls	Total	<i>p</i> value
Age (year)	14.69 (2.45)	14.76(2.37)	14.73(2.41)	0.24
Height (cm)	151.80(11.64)	156.54(15.59)	154.16(13.95)	<0.001*
Weight (Kg)	45.93(13.21)	48.42(16.53)	47.17(15.00)	<0.001*
BMI z-score	0.46 (1.0)	-0.46 (1.0)	0(1.0)	0.001*
WC (cm)	67.60(22.18)	69.85(18.99)	68.72(20.69)	<0.001*
WHiR	0.44 (0.14)	0.45(0.12)	0.44(0.12)	0.73
Obesity; n (%)	286 (10.1)	215 (7.7)	501 (8.9)	0.001*
Overweight; n (%)	265 (9.4)	186 (6.6)	451 (8.0)	<0.001*
Abdominal obesity; n (%)	415 (14.7)	471 (16.9)	886 (15.8)	0.02*
Father's education; n (%)				0.48
<6y	1138 (41.4)	1153 (42.6)	2291 (42.0)	
6–9y	662 (24.1)	668 (24.7)	1330 (24.4)	
10–12y	685 (24.9)	626 (23.2)	1311 (24.0)	
>12y	263 (9.6)	257 (9.5)	520 (9.5)	
Mother's education; n (%)				0.50
<6y	1489 (53.6)	1515 (55.6)	3004 (54.6)	
6–9y	580 (20.9)	548 (20.1)	1128 (20.5)	
10–12y	570 (20.5)	534 (19.6)	1104 (20.1)	
>12y	141 (5.1)	128 (4.7)	269 (4.9)	
ST Activity(hours/day)				
Watching TV; n (%)				0.65
<2	991 (35.6)	940 (35.0)	1931 (35.3)	
≥2	1790 (64.4)	1742 (65.0)	3532 (64.7)	
Working with computer; n (%)				<0.001*
<2	2372 (84.3)	2803 (76.5)	4453 (80.4)	
≥2	443 (15.7)	641 (23.5)	1084 (19.6)	
ST; n (%) [¶]				<0.001*
Low	1576 (56.8)	1370 (51.3)	2946 (54.1)	
High	1199 (43.2)	1303 (48.7)	2502 (45.9)	
PA; n (%) [£]				<0.001*
Low	2491 (88.4)	2224 (81.3)	4715 (84.9)	
High	327 (11.6)	510 (18.7)	837 (15.1)	
Joint associations of PA & ST; n (%)				<0.001*
High PA & Low ST	208 (7.5)	262 (9.8)	470 (8.6)	
High PA & High ST	114 (4.1)	235 (8.8)	349 (6.4)	
Low PA & Low ST	1368 (49.3)	1107 (41.4)	2475 (45.5)	
Low PA & High ST	1083 (39.1)	1068 (40.0)	2151 (39.5)	
Family history; n (%)				
HTN	1134 (51.4)	1039 (47.7)	2173 (49.5)	0.02*
Dyslipidemia	909 (42.7)	859 (40.3)	1768 (41.5)	0.13
DM	831 (39.1)	717 (34.2)	1548 (36.7)	0.001*
Obesity	876 (42.8)	747 (37.1)	1623 (40.0)	<0.001*
Type of home; n (%)				0.63
Rented home	531 (19.5)	531 (20.0)	1062 (19.8)	
Owned home	2193 (80.5)	2119 (80.0)	4312 (80.2)	
Private car; n (%)				0.83
Yes	1373 (49.8)	1340 (49.5)	2713 (49.7)	

(Continued)

Table 1. (Continued)

	Boys	Girls	Total	<i>p</i> value
No	1382 (50.2)	1366 (50.5)	2748 (50.3)	
SES; n (%)				0.76
Low	527 (19.6)	525 (20.0)	1052 (19.8)	
Intermediate	1047 (38.9)	1038 (39.5)	2085 (39.2)	
High	1115 (41.5)	1063 (40.5)	2178 (41.0)	

Data are means (SD) unless indicated otherwise.

BMI; Body Mass Index, WC; Waist Circumference, HTN; Hypertension, DM; Diabetes Mellitus, ST; Screen Time, PA; Physical Activity, SES; Socioeconomic status.

[¶] Low ST: <2 hours/day; high ST: ≥2 hours/day

[£] Low PA: 0–2 days/week; high PA: 3–7 days / week

* Statistically significant

doi:10.1371/journal.pone.0154502.t001

Discussion

The current findings on the joint effects of PA and ST levels showed that adolescents with low PA/high ST levels had the higher BMI z-score, WC, LDL-C, SBP, and DBP, as well as the lower

Table 2. Mean (SD) of cardiometabolic values according to ST, PA and joint associations of ST and PA categories: The CASPIAN III study.

	ST [¶]			PA [£]			Joint association of PA and ST				
	Low	High	<i>p</i> -value	Low	High	<i>p</i> -value	High PA/ Low ST	High PA & High ST	Low PA & Low ST	Low PA/ High ST	<i>P</i> -value
BMI z-score	-0.07 (1.0)	0.09 (1.0)	<0.001*	0.01 (1.0)	- 0.07 (1.0)	0.03*	-0.15 (1.0) ^a	0.07 (1.0) ^b	-0.05 (1.0) ^{ab}	0.09 (1.0) ^{bc}	<0.001*
WC (cm)	67.9 (22.8)	69.7 (18.5)	0.002*	68.8 (20.4)	68.3 (22.8)	0.03*	67.8(28.9) ^{ab}	69.4(10.7) ^{ab}	67.9(21.4) ^b	69.7(19.4) ^a	0.023*
WHTR	0.45 (0.15)	0.45 (0.11)	0.96	0.44 (0.13)	0.45 (0.15)	0.67	0.45(0.19)	0.44(0.05)	0.44(0.14)	0.45(0.11)	0.851
FBG (mg/dl)	87.8 (13.4)	87.5 (14.6)	0.45	87.5 (14.1)	88.6 (12.6)	0.06	88.0(12.5)	89.5(12.9)	87.8(13.6)	87.2(14.8)	0.064
TG (mg/dl)	91.4 (38.9)	94.2 (42.8)	0.02*	92.8 (41.1)	91.9 (39.8)	0.59	90.6(38.7)	93.8(41.6)	91.6(39.0)	94.3(43.0)	0.137
HDL-C (mg/dl)	47.1 (14.5)	45.3 (14.3)	<0.001*	45.9 (14.1)	48.5 (14.9)	<0.001*	47.9(14.6) ^a	49.4(15.7) ^a	46.9(14.5) ^a	44.7(13.8) ^b	<0.001*
TC (mg/dl)	148.2 (31.4)	149.1 (32.3)	0.36	151.3 (33.3)	147.9 (31.4)	0.01*	151.7(34.1) ^a	151.4(32.5) ^{ab}	147.5 (30.8) ^b	148.7 (32.2) ^{ab}	0.035*
LDL-C (mg/dl)	82.9 (26.9)	85.7 (27.8)	0.005*	84.2 (27.4)	84.1 (27.3)	0.91	84.0 (27.6) ^{ab}	84.1(27.1) ^{ab}	82.7(26.8) ^b	85.9(27.9) ^a	0.025*
SBP (mmHg)	102.6 (13.8)	103.9 (14.0)	<0.001*	103.3 (13.9)	102.2 (14.1)	0.04*	101.2(14.1) ^a	103.6(13.9) ^{ab}	102.8 (13.7) ^a	104.0 (14.1) ^b	0.001*
DBP (mmHg)	65.3 (10.8)	66.2 (10.6)	0.001*	65.8 (10.6)	64.8 (11.2)	0.01*	64.6(11.5) ^a	65.2(10.9) ^{ab}	65.4 (10.6) ^a	66.4(10.6) ^b	0.001*

Data are mean(SD),BMI; Body Mass Index, WC; Waist Circumference, WHTR; Waist to Height Ratio, FBS; Fasting Blood Sugar, TG; Triglycerides, HDL-C; High Density Lipoprotein-Cholesterol, TC; Total Cholesterol, LDL-C; Low-Density Lipoprotein Cholesterol, SBP; Systolic Blood Pressure; DBP; diastolic Blood Pressure, ST; Screen Time, PA; Physical Activity.

For joint association of PA and ST, within rows, means with different superscript letters are significantly different (by Tukey’s post hoc tests).

[¶] Low ST: <2 hours/day; high ST: ≥2 hours/day

[£] Low PA: 0–2 days/week; high PA: 3–7 days / week

* Statistically significant

doi:10.1371/journal.pone.0154502.t002

Table 3. Prevalence of cardiometabolic risk factors according to ST, PA and joint associations of ST and PA categories: The CASPIAN III study.

	ST			PA			Joint association of ST and PA				
	Low	High	<i>p</i> value	Low	High	<i>p</i> value	High PA/ Low ST	High PA & High ST	Low PA & Low ST	Low PA/ High ST	<i>p</i> value
Abdominal obesity	445 (15.1)	416 (16.7)	0.17	756 (16.1)	123 (14.7)	0.33	66 (14.0)	56 (16.0)	379 (15.3)	360 (16.8)	0.384
Elevated SBP	96(3.5)	94 (4.0)	0.37	168 (3.8)	28 (3.6)	0.9	12 (2.8)	16 (5.0)	84 (3.7)	77 (3.8)	0.464
Elevated DBP	81 (2.9)	76 (3.2)	0.57	128 (2.9)	29 (3.7)	0.26	16 (3.6)	13 (3.9)	65 (2.8)	63 (3.1)	0.582
Elevated BP	140 (5.3)	141 (6.2)	0.16	236 (5.5)	47 (6.2)	0.44	23 (5.4)	24 (7.7)	117 (5.2)	117 (6.0)	0.321
Elevated TG	172 (7.1)	187 (9.2)	0.01*	310 (8.1)	54 (7.8)	0.88	31 (7.9)	22 (7.7)	141 (7.0)	165 (9.4)	0.051
Elevated FBG	356 (15.0)	317 (16.2)	0.27	561 (15.0)	119 (17.4)	0.12	64 (16.5)	53 (19.0)	291 (14.6)	264 (15.8)	0.255
Low HDL-C	694 (33.5)	651 (36.9)	0.03*	1210 (36.4)	159 (27.2)	<0.001*	98 (29.8)	59 (24.4)	596 (34.3)	592 (38.9)	<0.001*
Elevated TC	139 (5.7)	127 (6.0)	0.61	207 (5.3)	60 (8.7)	.001*	37 (9.4)	23 (8.1)	102 (5.0)	104 (5.7)	0.002*
Elevated LDL-C	130 (5.0)	130 (6.9)	0.03*	220 (5.9)	35 (5.0)	0.45	20 (4.9)	20 (5.5)	105 (5.0)	124 (7.1)	0.117
General obesity	257 (8.7)	230 (9.2)	0.57	427 (9.1)	70 (8.4)	0.56	39 (8.3)	31 (8.9)	218 (8.8)	199 (9.3)	0.908
Overweight	217 (7.4)	224 (9.0)	0.04*	390 (8.3)	60 (7.2)	0.30	30 (6.4)	29 (8.3)	187 (7.6)	195 (9.1)	0.135
Number of Mets components			0.04*			0.46					0.301
0	840 (46.2)	630 (40.9)		1259 (43.3)	241 (46.6)		143 (48.3)	92 (43.6)	697 (45.8)	537 (40.4)	
1	695 (38.2)	631 (40.9)		1156 (39.8)	195 (37.7)		111 (37.5)	80 (37.9)	583 (38.3)	551 (41.4)	
2	216 (11.9)	213 (13.8)		367 (12.6)	67 (13.0)		34 (11.5)	33 (15.6)	182 (12.0)	180 (13.5)	
3	58 (3.2)	57 (3.7)		105 (3.6)	11 (2.1)		6 (2.0)	5 (2.4)	52 (3.4)	52 (3.9)	
4	10 (0.6)	11 (0.7)		18 (0.6)	3 (0.6)		2 (0.7)	1 (0.5)	8 (0.5)	10 (0.8)	
Having MetS	68 (3.7)	68 (4.4)	0.34	123 (4.2)	14 (2.7)	0.11	8 (2.7)	6 (2.8)	60 (3.9)	62 (4.7)	0.321

Data are N (%), BP; Blood Pressure, TG; Triglycerides, FBG; Fasting Blood Glucose, HDL-C; High Density Lipoprotein-Cholesterol, TC; Total Cholesterol, LDL-C; Low-Density Lipoprotein Cholesterol, MetS; Metabolic Syndrome, ST; Screen Time, PA; Physical Activity.

Abdominal obesity: WHtR>0.05; Elevated SBP: systolic BP>90th adjusted by age, sex and height; Elevated DBP: diastolic BP>90th adjusted by age, sex and height; Elevated BP: either systolic or diastolic BP>90th adjusted by age, sex and height; Elevated FBS:>100 mg/dl; Elevated TG:>= 100 mg/dl; Low HDL: <50 mg/dl (except in boys 15–19 years old, that cut-off was <45 mg/dl); Elevated TC:>200 mg/dl; Elevated LDL:>110 mg/dl; Overweight: BMI:85th–95th; Obesity: BMI > 95th; Having MetS: having at least three criteria according to modified ATP III criteria; Low ST: <2 hours/day; high ST: ≥2 hours/day; Low PA: 0–2 days/week; high PA: 3–7 days / week.

* Statistically significant

doi:10.1371/journal.pone.0154502.t003

HDL-C levels compared to other combinations of PA/ST. In addition, participants with high PA/low ST had the highest mean serum levels of total cholesterol amongst other counterparts, which could be explained by high levels of HDL-C in this group. To the best of our knowledge,

Table 4. Odds ratios (95% CI) for cardiometabolic risk factors according to joint associations of ST and PA categories: The CASPIAN III study.

		Joint association of ST and PA							
		High PA & Low ST		High PA & High ST		Low PA & Low ST		Low PA & High ST	
		OR	CI	OR	CI	OR	CI	OR	CI
Abdominal obesity									
Model I	1	1.170	0.795–1.722	1.108	0.835–1.469	1.234	0.929–1.639		
Model II	1	1.133	0.769–1.670	1.121	0.844–1.489	1.229	0.923–1.636		
Model III	1	1.204	0.731–1.984	1.283	0.892–1.846	1.485*	1.029–2.144		
Elevated BP									
Model I	1	1.455	0.805–2.629	0.969	0.612–1.534	1.115	0.704–1.766		
Model II	1	1.208	0.664–2.199	0.906	0.569–1.444	0.944	0.592–1.506		
Model III	1	1.202	0.582–2.483	1.063	0.613–1.844	0.991	0.567–1.732		
Model IV	1	1.048	0.499–2.205	1.018	0.580–1.784	0.897	0.507–1.586		
Elevated FBG									
Model I	1	1.191	0.797–1.780	0.871	0.648–1.171	0.949	0.704–1.280		
Model II	1	1.127	0.752–1.689	0.913	0.678–1.231	0.971	0.717–1.314		
Model III	1	1.072	0.678–1.695	0.829	0.595–1.155	0.918	0.656–1.286		
Model IV	1	1.063	0.672–1.682	0.828	0.594–1.153	0.912	0.651–1.278		
Elevated TG									
Model I	1	0.981	0.555–1.732	0.876	0.584–1.313	1.218	0.816–1.817		
Model II	1	0.952	0.538–1.685	0.845	0.563–1.270	1.157	0.772–1.733		
Model III	1	0.663	0.315–1.395	0.867	0.541–1.388	1.091	0.680–1.750		
Model IV	1	0.586	0.274–1.250	0.811	0.500–1.315	0.984	0.605–1.598		
Low HDL-C									
Model I	1	0.760	0.521–1.108	1.228	0.950–1.587	1.502*	1.161–1.944		
Model II	1	0.709	0.486–1.036	1.187	0.917–1.536	1.400*	1.078–1.817		
Model III	1	0.766	0.492–1.193	1.219	0.907–1.640	1.311	0.971–1.771		
Model IV	1	0.748	0.480–1.165	1.218	0.906–1.639	1.297	0.960–1.753		
Elevated TC									
Model I	1	0.842	0.489–1.452	0.502*	0.339–0.744	0.584*	0.394–0.864		
Model II	1	0.896	0.518–1.549	0.499*	0.336–0.741	0.598*	0.401–0.891		
Model III	1	0.956	0.506–1.808	0.500*	0.318–0.788	0.647	0.409–1.023		
Model IV	1	0.908	0.479–1.720	0.487*	0.309–0.768	0.615*	0.388–0.976		
Elevated LDL-C									
Model I	1	1.134	0.486–2.644	1.039	0.567–1.904	1.501	0.826–2.727		
Model II	1	1.243	0.531–2.910	1.067	0.580–1.963	1.645	0.898–3.011		
Model III	1	1.583	0.557–4.501	1.200	0.558–2.584	1.991	0.933–4.248		
Model IV	1	1.542	0.542–4.389	1.192	0.554–2.566	1.958	0.917–4.180		
Overweight									
Model I	1	1.329	0.782–2.259	1.199	0.805–1.786	1.462	0.982–2.177		
Model II	1	1.400	0.822–2.384	1.163	0.779–1.736	1.452	0.972–2.171		
Model III	1	1.577	0.813–3.058	1.224	0.743–2.017	1.773*	1.075–2.923		
General obesity									
Model I	1	1.077	0.658–1.764	1.067	0.748–1.524	1.127	0.787–1.613		
Model II	1	1.160	0.707–1.904	1.073	0.750–1.535	1.181	0.822–1.697		
Model III	1	1.222	0.629–2.378	1.292	0.803–2.079	1.418	0.874–2.302		
Having MetS									
Model I	1	1.054	0.360–3.083	1.477	0.699–3.123	1.760	0.834–3.717		

(Continued)

Table 4. (Continued)

Joint association of ST and PA							
	High PA & Low ST	High PA & High ST		Low PA & Low ST		Low PA & High ST	
		OR	CI	OR	CI	OR	CI
Model II	1	0.898	0.306–2.638	1.410	0.664–2.995	1.557	0.733–3.310
Model III	1	0.991	0.260–3.782	1.599	0.623–4.101	1.571	0.606–4.074
Model IV	1	0.737	0.180–3.023	1.563	0.569–4.292	1.536	0.552–4.274

BP; Blood Pressure, TG; Triglycerides, FBG; Fasting Blood Glucose, HDL-C; High Density Lipoprotein-Cholesterol, TC; Total Cholesterol, LDL-C; Low-Density Lipoprotein Cholesterol, MetS; Metabolic Syndrome, ST; screen time, PA; physical activity.

Abdominal obesity: WHtR>0.05; Elevated BP: either systolic or diastolic BP>90th adjusted by age, sex and height; Elevated FBS:>100 mg/dl; Elevated TG:>= 100 mg/dl; Low HDL: <50 mg/dl (except in boys 15–19 years old, that cut-off was <45 mg/dl); Elevated TC:>200 mg/dl; Elevated LDL:>110 mg/dl; Overweight: BMI:85th-95th; Obesity: BMI > 95th; Having MetS: having at least three criteria according to modified ATP III criteria; Low ST: <2 hours/day; high ST: ≥2 hours/day; Low PA: 0–2 days/week; high PA: 3–7 days / week.

Model I: crude model; Model II is adjusted for age and gender; Model III adjusted additionally for family history and socio-economic status; Model IV is adjusted additionally for BMI in all abnormalities except obesity.

* Statistically significant

doi:10.1371/journal.pone.0154502.t004

this is the first study in Middle Eastern adolescents that reports the combined associations of ST and PA with the odds of having cardio-metabolic risk factors.

Considering the importance and priority of problem, epidemiological aspects of cardiometabolic risk factors become one of the most attractive domain of health research [1,2,39, 46]. Highest prevalence of pediatric metabolic risk factors and MetS of developing countries are reported from the Eastern Europe and the Middle East, and the lowest one from India and Sri Lanka [52].

Our findings are consistent with some previous studies conducted on the joint effect of PA and ST or sedentary behaviors on BMI in Japanese [53], Australian [54], and US adults [55]. Our findings about joint association of PA and ST with cardiometabolic risk factors are contrary to Drenowatz et al. study [36]. Their study reported no significant association between joint effect of PA and ST with cardiometabolic risk factors in 10-year-old children; however, in that study, in line with our results low PA and high ST were related to a higher CVD risk score [36]. Some studies emphasized that higher activity levels were associated with a healthier diet and lower ST indicating an overall healthier lifestyle of this subgroup [33].

The current findings show that adolescents with low PA/ high ST were about 1.5 times more likely to have low levels of HDL-C, with approximately 40% lower risk of elevated TC compared to those with high PA/ low ST. After adjustment for confounding factors, these participants were 1.5 times more likely to have abdominal obesity, and 1.5 to 1.8 times more likely to be overweight than those in the high PA/ low ST group.

Some studies have reported that sufficient PA is associated with reduced cardiometabolic risk in children and adolescents [30, 56]. In a cross-sectional study in 1732 school children from Denmark, Estonia, and Portugal, negative association existed between PA and clustered cardiovascular risk [57]. Pooled data from 14 studies comprising 20871 children and adolescents, aged 4–18 years, also showed that moderate to vigorous PA might be associated with lower cardiometabolic risk factors, regardless of their amount of sedentary behaviors [58].

There is some evidences that the association between PA/ST and MetS differs according to sex and type of sedentary behaviors [59–60]. In the current study, sedentary behaviors included times spent for watching TV/VCDs and playing computer/electronic games, but other

sedentary behaviors such as homework and motorized transport were not considered. A cohort study of Canadian children aged 8–10 years showed that not overall sedentary behaviors, but only ST was independently associated with WC and HDL-C [61]. In the present study, joint association of PA and ST was associated with sex. A cross-sectional study of French adults showed that the relationship of the MetS with ST and PA differs according to sex [60].

The absence of statistically significant associations between the PA/ST and elevated FBG, TG, and BP may be explained by the generally better status of these parameters or small number of adolescents with unfavorable levels of these risk factors.

One plausible mechanism responsible for the adverse association of sedentary behaviors on serum HDL-C is the suppression of the rate-limiting enzyme lipoprotein lipase activity in skeletal muscles. This is an enzyme necessary for HDL-C production and TG uptake [62, 63]. In addition to this physiologic mechanism, it is suggested that students eat unhealthy snacks more frequently while watching TV [63]. These findings suggest that lower BMI and WC, as well as improving HDL-C should be encouraged by highlighting the importance of increasing PA and reducing ST.

As one of the main strengths, present study benefited from a large national representative sample of Iranian adolescents. Moreover, it led exactly based on the World Health Organization- Global School-based student Health Survey (WHO-GSHS) protocol.

We also faced several limitations. First, the cross-sectional design of study which does not demonstrate the causality or specify the direction of causation between the cardiometabolic risk factors and PA/ST. Therefore, it may be that participants with a higher BMI do less PA and engage more extensively in sedentary behaviors like TV viewing. Future studies examining the prospective association of prolonged sitting on the risk of developing cardiometabolic risk factors would be warranted. Second, ST and PA data were based on self-reports that may be subject to recall bias. Finally, the possible contributing factors, as eating snacks or drinking soft drinks during ST, were not integrated into the study. Since some evidences show that association between PA/ST and MetS differs according to sex and type of sedentary behaviors, therefore stratify analysis according to sex and type of sedentary behaviors is suggested for future research.

A next step could be assessing the clustering of multiple life style risk factors associated with high ST levels in Iranian adolescents. This study used a large sample-size which was representative of all parts and strata of Iranian pediatric population. Moreover, as the basic practical evidences for interventional health programs, complementary researches on determinant of differences between boys and girls are recommend.

Conclusion

The findings of this study showed that joint association of high ST and low PA have direct association with abdominal obesity, overweight and low HDL-C and indirect association with elevated TC. The current findings underscore the importance of reducing ST along with increasing PA for reducing the risk of developing cardiometabolic risk factors. Future public interventions are needed focusing on sedentary behaviors and PA from early life.

Acknowledgments

This nationwide survey was conducted in Iran with corporation of the Ministry of Health and Medical education, Ministry of Education and Training, Child Health Promotion Research Center, Isfahan University of Medical Sciences, and Endocrinology and Metabolism Research Institute of Tehran University of Medical Sciences.

Author Contributions

Conceived and designed the experiments: RH RK MEM. Performed the experiments: GA AAJ SD. Analyzed the data: MQ HA. Contributed reagents/materials/analysis tools: AAJ SD. Wrote the paper: AESB MQ. Data acquisition: TA HA FR.

References

1. Zimmet P, Alberti G, Kaufman F, Tajima N, Silink M, Arslanian S, et al. (2007) The metabolic syndrome in children and adolescents—an IDF consensus report. *Pediatr diabete* 8:299–306.
2. Kelishadi R, Ardalan G, Adeli K, Motaghian M, Majdzadeh R, Mahmood-Arabi MS, et al. (2007) Factor analysis of cardiovascular risk clustering in pediatric metabolic syndrome: CASPIAN study. *Ann Nutr Metab* 51:208–15. PMID: [17587791](#)
3. Hajian-Tilaki KO, Sajjadi P, Razavi A (2011) Prevalence of overweight and obesity and associated risk factors in urban primary-school children in Babol, Islamic Republic of Iran. *East Mediterr Health J* 17:109–14. PMID: [21735944](#)
4. Khashayar P, Heshmat R, Qorbani M, Motlagh ME, Aminaee T, Ardalan G, et al. (2013) Metabolic Syndrome and Cardiovascular Risk Factors in a National Sample of Adolescent Population in the Middle East and North Africa: The CASPIAN III Study. *Int J endocrinol* 2013:702095. PMID: [23476647](#). doi: [10.1155/2013/702095](#)
5. Hajian-Tilaki K, Heidari B (2012) Prevalences of overweight and obesity and their association with physical activity pattern among Iranian adolescents aged 12–17 years. *Public Health Nutr* 15:2246–52. doi: [10.1017/S1368980012001048](#) PMID: [22578771](#)
6. Kelishadi R, Ardalan G, Gheiratmand R, Gouya MM, Razaghi EM, Majdzadeh R, et al. (2007) Association of physical activity and dietary behaviours in relation to the body mass index in a national sample of Iranian children and adolescents: CASPIAN Study. *Bull World Health Organ* 85:19–26. PMID: [17242754](#)
7. Kelishadi R, Razaghi EM, Gouya MM, Ardalan G, Gheiratmand R, Delavari A, et al. (2007) Association of physical activity and the metabolic syndrome in children and adolescents: CASPIAN Study. *Horm Res* 67:46–52. PMID: [17035710](#)
8. Mehrkash M, Kelishadi R, Mohammadian S, Mousavinasab F, Qorbani M, Hashemi ME, et al. (2012) Obesity and metabolic syndrome among a representative sample of Iranian adolescents. *Southeast Asian J Trop Med Public Health* 43:756–63. PMID: [23077856](#)
9. Shafiee G, Kelishadi R, Qorbani M, Motlagh ME, Taheri M, Ardalan G, et al. (2013) Association of breakfast intake with cardiometabolic risk factors. *J Pediatr (Rio J)* 89:575–82.
10. Kaler S, Ralph-Campbell K, Pohar S, King M, Laboucan C, Toth E. (2006) High rates of the metabolic syndrome in a First Nations Community in western Canada: prevalence and determinants in adults and children. *Int J Circumpolar Health* 65(5):389. PMID: [17319084](#)
11. Park Y-W, Zhu S, Palaniappan L, Heshka S, Carnethon MR, Heymsfield SB. (2003) The Metabolic Syndrome: Prevalence and Associated Risk Factor Findings in the US Population From the Third National Health and Nutrition Examination Survey, 1988–1994. *Archives of internal medicine* 163(4):427. PMID: [12588201](#)
12. Grundy S. Metabolic syndrome pandemic. (2008) *Arteriosclerosis, thrombosis, and vascular biology* 28(4):629. doi: [10.1161/ATVBAHA.107.151092](#) PMID: [18174459](#)
13. Molnár D. (2004) The prevalence of the metabolic syndrome and type 2 diabetes mellitus in children and adolescents. *International journal of obesity and related metabolic disorders: journal of the International Association for the Study of Obesity* 28:S70.
14. Li Y, Yang X, Zhai F, Kok F, Zhao W, Piao J, et al. (2008) Prevalence of the metabolic syndrome in Chinese adolescents. *The British journal of nutrition* 99(3):565. PMID: [17662161](#)
15. Park J, Hilmers DC, Mendoza JA, Stuff JE, Liu Y, Nicklas TA. (2010) Prevalence of Metabolic Syndrome and Obesity in Adolescents Aged 12 to 19 Years: Comparison between the United States and Korea. *Journal of Korean Medical Science* 25(1):75. doi: [10.3346/jkms.2010.25.1.75](#) PMID: [20052351](#)
16. Kelishadi R. (2007) Childhood overweight, obesity, and the metabolic syndrome in developing countries. *Epidemiologic reviews* 29:62. PMID: [17478440](#)
17. McMahon S, Haynes A, Ratnam N, Grant M, Carne C, Jones T, et al. (2004) Increase in type 2 diabetes in children and adolescents in Western Australia. *The Medical journal of Australia* 180(9):459. PMID: [15115424](#)
18. Qorbani M, Kelishadi R, Farrokhi-Khajeh-Pasha Y, Motlagh ME, Aminaee T, Ardalan G, et al. (2013). Association of anthropometric measures with cardiovascular risk factors and metabolic syndrome in

- normal-weight children and adolescents: The CASPIAN III study. *Obesity facts* 6: 483–492. doi: [10.1159/000356011](https://doi.org/10.1159/000356011) PMID: [24157679](https://pubmed.ncbi.nlm.nih.gov/24157679/)
19. Camhi SM, Katzmarzyk PT (2010) Tracking of cardiometabolic risk factor clustering from childhood to adulthood. *Int J Pediatr Obes* 5:12.
 20. Golden SH, Folsom AR, Coresh J, Sharrett AR, Szklo M, Brancati F. (2002) Risk factor groupings related to insulin resistance and their synergistic effects on subclinical atherosclerosis: the atherosclerosis risk in communities study. *Diabetes* 51:3069–76. PMID: [12351449](https://pubmed.ncbi.nlm.nih.gov/12351449/)
 21. Gehani AA, Al-Hinai AT, Zubaid M, Almahmeed W, Hasani MM, Yusufali AH, et al. (2014) Association of risk factors with acute myocardial infarction in Middle Eastern countries: the INTERHEART Middle East study. *Eur J Prev Cardiol* 21(4):400–10. doi: [10.1177/2047487312465525](https://doi.org/10.1177/2047487312465525) PMID: [23125402](https://pubmed.ncbi.nlm.nih.gov/23125402/)
 22. Singh GM, Danaei G, Pelizzari PM, Lin JK, Cowan MJ, Stevens GA, et al. (2012) The age associations of blood pressure, cholesterol, and glucose: analysis of health examination surveys from international populations. *Circulation* 125 (18): 2204–2211. doi: [10.1161/CIRCULATIONAHA.111.058834](https://doi.org/10.1161/CIRCULATIONAHA.111.058834) PMID: [22492580](https://pubmed.ncbi.nlm.nih.gov/22492580/)
 23. Kelishadi R, Hovsepian S, Qorbani M, Jamshidi F, Fallah Z, Djalalinia Sh, et al. (2014) National and Sub-National Prevalence, Trend, and Burden of Cardiometabolic Risk Factors in Iranian Children and Adolescents, 1990–2013 *Arch Iran Med*17(1): 71–80.
 24. Kelishadi R, Gheiratmand R, Ardalan G, Adeli K, Mehdi Gouya M, Mohammad Razaghi E, et al. (2007) Association of anthropometric indices with cardiovascular disease risk factors among children and adolescents: CASPIAN Study. *Int J Cardiol* 117: 340–348. PMID: [16860411](https://pubmed.ncbi.nlm.nih.gov/16860411/)
 25. Vereecken CA, Todd J, Roberts C, Mulvihill C, Maes L (2006). Television viewing behaviour and associations with food habits in different countries. *Public Health Nutr* 9: 244–50. PMID: [16571179](https://pubmed.ncbi.nlm.nih.gov/16571179/)
 26. Bryant MJ, Lucove JC, Evenson KR, Marshall S (2007). Measurement of television viewing in children and adolescents: a systematic review. *Obes Rev* 8:197–209. PMID: [17444962](https://pubmed.ncbi.nlm.nih.gov/17444962/)
 27. Hinkley T, Crawford D, Salmon J, Okely AD, Hesketh K (2008). Preschool children and physical activity: a review of correlates. *Am J Prev Med* 34:435–441. doi: [10.1016/j.amepre.2008.02.001](https://doi.org/10.1016/j.amepre.2008.02.001) PMID: [18407012](https://pubmed.ncbi.nlm.nih.gov/18407012/)
 28. Jari M, Qorbani M, Motlagh ME, Heshmat R, Ardalan G, Kelishadi R. (2014). A nationwide survey on the daily screen time of Iranian children and adolescents: the CASPIAN-IV study. *International journal of preventive medicine* 5: 224. PMID: [24627751](https://pubmed.ncbi.nlm.nih.gov/24627751/)
 29. Baygi F, Heshmat R, Kelishadi R, Mohammadi F, Motlagh M E, Ardalan G, et al. (2015). Regional Disparities in Sedentary Behaviors and Meal Frequency in Iranian Adolescents: The CASPIAN-III Study. *Iran J Pediatr*, 25: 182.
 30. Ekelund U, Luan J, Sherar LB, Esliger DW, Griew P, Cooper A, et al. (2012) Moderate to vigorous physical activity and sedentary time and cardiometabolic risk factors in children and adolescents. *JAMA* 307:704–12. doi: [10.1001/jama.2012.156](https://doi.org/10.1001/jama.2012.156) PMID: [22337681](https://pubmed.ncbi.nlm.nih.gov/22337681/)
 31. Martinez-Gomez D, Gomez-Martinez S, Ruiz JR, Ortega FB, Marcos A, Veiga OL. (2012) Video game playing time and cardiometabolic risk in adolescents: the AFINOS study. *Med Clin (Barc)* 139:290–2.
 32. Martinez-Gomez D, Rey-Lopez JP, Chillon P, Gomez-Martinez S, Vicente-Rodriguez G, et al. (2010) Excessive TV viewing and cardiovascular disease risk factors in adolescents. The AVENA cross-sectional study. *BMC Public Health* 10:274. doi: [10.1186/1471-2458-10-274](https://doi.org/10.1186/1471-2458-10-274) PMID: [20500845](https://pubmed.ncbi.nlm.nih.gov/20500845/)
 33. Nang EE, Salim A, Wu Y, Tai ES, Lee J, Van Dam RM. (2013) Television screen time, but not computer use and reading time, is associated with cardio-metabolic biomarkers in a multiethnic Asian population: a cross-sectional study. *Int J Behav Nutr Phys Act* 10:70. doi: [10.1186/1479-5868-10-70](https://doi.org/10.1186/1479-5868-10-70) PMID: [23718927](https://pubmed.ncbi.nlm.nih.gov/23718927/)
 34. Byun W, Dowda M, Pate RR (2012) Associations between screen-based sedentary behavior and cardiovascular disease risk factors in Korean youth. *J Korean Med Sci* 27:388–94. doi: [10.3346/jkms.2012.27.4.388](https://doi.org/10.3346/jkms.2012.27.4.388) PMID: [22468102](https://pubmed.ncbi.nlm.nih.gov/22468102/)
 35. Altenburg TM, Hofsteenge GH, Weijs PJ, Delemarre-van de Waal HA, Chinapaw MJ (2012) Self-reported screen time and cardiometabolic risk in obese Dutch adolescents. *PLoS One* 7:e53333. doi: [10.1371/journal.pone.0053333](https://doi.org/10.1371/journal.pone.0053333) PMID: [23285284](https://pubmed.ncbi.nlm.nih.gov/23285284/)
 36. Drenowatz C, Carlson JJ, Pfeiffer KA, Eisenmann JC (2012) Joint association of physical activity/screen time and diet on CVD risk factors in 10-year-old children. *Front Med* 6:428–35. doi: [10.1007/s11684-012-0232-4](https://doi.org/10.1007/s11684-012-0232-4) PMID: [23224418](https://pubmed.ncbi.nlm.nih.gov/23224418/)
 37. Steele RM, van Sluijs EMF, Cassidy A, Griffin SJ, Ekelund U (2009). Targeting sedentary time or moderate- and vigorous-intensity activity. *Am J Clin Nutr* 90:1185–1192. doi: [10.3945/ajcn.2009.28153](https://doi.org/10.3945/ajcn.2009.28153) PMID: [19776141](https://pubmed.ncbi.nlm.nih.gov/19776141/)
 38. Fisher A, Hill C, Webber L, Purslow L, Wardle J (2011). MVPA is associated with lower weight gain in 8–10 year old children. *PLoS One* 6: e18576. doi: [10.1371/journal.pone.0018576](https://doi.org/10.1371/journal.pone.0018576) PMID: [21552554](https://pubmed.ncbi.nlm.nih.gov/21552554/)

39. Verona J, Gilligan LE, Gimenez C, Verona MF, Lombardo SM, Baenz A, et al. (2013) Physical activity and cardiometabolic risk in male children and adolescents: The Balcarce study. *Life Sci* 93:64–8. doi: [10.1016/j.lfs.2013.05.021](https://doi.org/10.1016/j.lfs.2013.05.021) PMID: [23743170](https://pubmed.ncbi.nlm.nih.gov/23743170/)
40. Mitchell JA, Mattocks C, Ness AR, Leary SD, Pate RR, Dowda M, et al. (2009) Sedentary behavior and obesity in a large cohort of children. *Obesity (Silver Spring, Md)* 17:1596–602.
41. Kelishadi R, Heshmat R, Motlagh ME, Majdzadeh R, Keramatian K, Taslimi M, et al. (2012) Methodology and early findings of the third survey of CASPIAN study: A national school-based surveillance of students' high risk behaviors. *Int J Prev Med* 2012; 3:394. PMID: [22783465](https://pubmed.ncbi.nlm.nih.gov/22783465/)
42. Kelishadi R, Marashinia F, Heshmat R, Motlagh ME, Qorbani M, et al. (2013) First report on body image and weight control in a nationally representative sample of a pediatric population in the Middle East and North Africa: the CASPIAN-III study. *ArchMed Sci* 20;9:210–7.
43. Zimmet P, Alberti KGM, Kaufman F, Tajima N, Silink M, Arslanian S, et al. (2007) The metabolic syndrome in children and adolescents—an IDF consensus report. *Pediatr diabetes* 8:299–306. PMID: [17850473](https://pubmed.ncbi.nlm.nih.gov/17850473/)
44. McNamara JR, Schaefer EJ (1987) Automated enzymatic standardized lipid analyses for plasma and lipoprotein fractions. *Clinica Chimica Acta* 166:1–8.
45. Grundy SM, Bilheimer D, Chait A, Clark LT, Denke M, Havel RJ, et al. (1993) Summary of the second report of the National Cholesterol Education Program (NCEP) Expert Panel on detection, evaluation, and treatment of high blood cholesterol in adults (Adult Treatment Panel II). *JAMA* 269:3015–23. PMID: [8501844](https://pubmed.ncbi.nlm.nih.gov/8501844/)
46. Kelishadi R, Ardalan G, Gheiratmand R, Majdzadeh R, Hosseini M, Gouya MM, et al. (2008) Thinness, overweight and obesity in a national sample of Iranian children and adolescents: CASPIAN Study. *Child Care Health Dev* 34:44–54. doi: [10.1111/j.1365-2214.2007.00744.x](https://doi.org/10.1111/j.1365-2214.2007.00744.x) PMID: [18171443](https://pubmed.ncbi.nlm.nih.gov/18171443/)
47. Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III) final report (2002). *Circulation*. 106:3143–421. PMID: [12485966](https://pubmed.ncbi.nlm.nih.gov/12485966/)
48. Caro DH, Cortés D (2012) Measuring family socioeconomic status: An illustration using data from PIRLS 2006. *IERI Monograph Series. Issues and Methodologies in Large-Scale Assessments* 5, 9–33.
49. Salmon J, Campbell KJ, Crawford DA (2006) Television viewing habits associated with obesity risk factors: a survey of Melbourne schoolchildren. *Med J Aust* 184:64–7. PMID: [16411870](https://pubmed.ncbi.nlm.nih.gov/16411870/)
50. American Academy of Pediatrics (2001) children, adolescents, and television. *Pediatrics* 107(2):423–6. PMID: [11158483](https://pubmed.ncbi.nlm.nih.gov/11158483/)
51. Drenowatz C, Carlson JJ, Pfeiffer KA, Eisenmann JC. (2012) Joint association of physical activity/ screen time and diet on CVD risk factors in 10-year-old children. *Fron Med* 6:428–35.
52. Kelishadi R. Childhood overweight, obesity, and the metabolic syndrome in developing countries. *Epidemiol Rev*. 2007; 29:62–76. PMID: [17478440](https://pubmed.ncbi.nlm.nih.gov/17478440/)
53. Liao Y, Harada K, Shibata A, Ishii K, Oka K, Nakamura Y, et al. (2011) Joint associations of physical activity and screen time with overweight among Japanese adults. *Int J Behav Nutr Phys Act* 8:131. doi: [10.1186/1479-5868-8-131](https://doi.org/10.1186/1479-5868-8-131) PMID: [22128879](https://pubmed.ncbi.nlm.nih.gov/22128879/)
54. Sugiyama T, Healy GN, Dunstan DW, Salmon J, Owen N (2008) Joint associations of multiple leisure-time sedentary behaviours and physical activity with obesity in Australian adults. *Int J BehavNutrPhys Act* 5:35.
55. Dunton GF, Berrigan D, Ballard-Barbash R, Graubard B, Atienza AA (2009) Joint associations of physical activity and sedentary behaviors with body mass index: results from a time use survey of US adults. *Int J Obes (Lond)* 33:1427–1436.
56. Andersen LB, Harro M, Sardinha LB, Froberg K, Ekelund U, Brage S, et al. (2006) Physical activity and clustered cardiovascular risk in children: a cross-sectional study (The European Youth Heart Study). *Lancet* 368:299–304. PMID: [16860699](https://pubmed.ncbi.nlm.nih.gov/16860699/)
57. Ekelund U, Luan J, Sherar LB, Esliger DW, Griew P, Cooper A, et al. (2012) Moderate to vigorous physical activity and sedentary time and cardiometabolic risk factors in children and adolescents. *JAMA* 307:704–712. doi: [10.1001/jama.2012.156](https://doi.org/10.1001/jama.2012.156) PMID: [22337681](https://pubmed.ncbi.nlm.nih.gov/22337681/)
58. Chaput JP, Saunders TJ, Mathieu MÈ, Henderson M, Tremblay MS, O'Loughlin J et al. (2013) Combined associations between moderate to vigorous physical activity and sedentary behaviour with cardiometabolic risk factors in children. *Appl Physiol Nutr Metab* 38:477–83. doi: [10.1139/apnm-2012-0382](https://doi.org/10.1139/apnm-2012-0382) PMID: [23668753](https://pubmed.ncbi.nlm.nih.gov/23668753/)
59. Wilmore JH (2001) Dose-response: variation with age, sex, and health status. *Med Sci Sports Exerc* 33: 622–34.

60. Bertrais S, Beyeme-Ondoua JP, Czernichow S, Galan P, Hercberg S, et al. (2005) Sedentary Behaviors, Physical Activity, and Metabolic Syndrome in Middle-aged French Subjects. *Obesity Research* 13: 936–944. PMID: [15919848](#)
61. Bey L, Hamilton M (2003) Suppression of skeletal muscle lipoprotein lipase activity during physical inactivity: a molecular reason to maintain daily low-intensity activity. *J Physiol* 551: 673–682. PMID: [12815182](#)
62. Hamilton MT, Hamilton DG, Zderic TW (2004) Exercise physiology versus inactivity physiology: an essential concept for understanding lipoprotein lipase regulation. *Exerc Sport Sci Rev* 32:161–166. PMID: [15604935](#)
63. Thomson M, Spence JC, Raine K, Laing L (2008) The association of television viewing with snacking behavior and body weight of young adults. *Am J Health Promot* 22:329–335. doi: [10.4278/ajhp.22.5.329](#) PMID: [18517093](#)