

openheart Associations between physical activity, sedentary behaviour and left ventricular structure and function from the Echocardiographic Study of Latinos (ECHO-SOL)

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ABSTRACT

Objective The cross-sectional association between accelerometer-measured physical activity (PA), sedentary behaviour (SB) and cardiac structure and function is less well described. This study's primary aim was to compare echocardiographic measures of cardiac structure and function with accelerometer measured PA and SB.

Methods Participants included 1206 self-identified Hispanic/Latino men and women, age 45–74 years, from the Echocardiographic Study of Latinos. Standard echocardiographic measures included M-mode, two-dimensional, spectral, tissue Doppler and myocardial strain. Participants wore an Actical accelerometer at the hip for 1 week.

Results The mean±SE age for the cohort was 56±0.4 years, 57% were women. Average moderate to vigorous PA (MVPA) was 21±1.1 min/day, light PA was 217±4.2 min/day and SB was 737±8.1 min/day. Both higher levels of light PA and MVPA (min/day) were associated with lower left ventricular (LV) mass index (LVMI)/end-diastolic volume and a lower E/e' ratio. Higher levels of MVPA (min/day) were associated with better right ventricular systolic function. Higher levels of SB were associated with increased LVMI. In a multivariable linear regression model adjusted for demographics and cardiovascular disease modifiable factors, every 10 additional min/day of light PA was associated with a 0.03 mL/m² increase in left atrial volume index (LAVI) (p<0.01) and a 0.004 cm increase in tricuspid annular plane systolic excursion (p<0.01); every 10 additional min/day of MVPA was associated with a 0.18 mL/m² increase in LAVI (p<0.01) and a 0.24% improvement in global circumferential strain (p<0.01).

Conclusions Our findings highlight the potential positive association between the MVPA and light PA on cardiac structure and function.

INTRODUCTION

Lack of physical activity (PA) is common in the US population, and it is associated with multiple detrimental health outcomes.¹

Key questions

What is already known about this subject?

► Hispanic/Latino are the largest ethnic minority in the USA and represent a high-risk population; higher prevalence of cardiovascular disease factors, relatively understudied population and higher rates of sedentary behaviour (SB) and lower rates of physical activity (PA) than other ethnic groups.

What does this study add?

► To our knowledge, this is the first study to examine the association between moderate to vigorous PA and favourable left ventricle myocardial dynamics, remodelling, improved right ventricle function without any association with left ventricular mass in a large cohort of asymptomatic Hispanics/Latinos.

How might this impact on clinical practice?

► These findings may allow for more detailed, patient-specific, risk-stratification and prognosis in healthy individuals by gaining insight into the interpretation of resting echocardiography. This study provides further evidence that formal recommendations on limiting SB in persons of varying age should be developed to provide public health professionals and clinical practitioners with information to improve their effectiveness in promoting PA and health. Physical inactivity should be considered as an important cardiovascular disease risk factor.

Troubling is the even lower trend of PA in Hispanic/Latino men and women. According to 2008 report of the National Health Interview Survey, Hispanics/Latinos had lower leisure-time PA levels than non-Hispanic whites.² Similarly, 2007–2016 National Health and Nutrition Examination Survey (self-reported data) indicate that Hispanics/Latinos are less likely to meet recommended

PA guidelines compared with non-Hispanic whites (33.4% vs 47.6%).³

PA is associated with cardiovascular (CV) health benefits, including prevention of age-related cardiac remodelling, systolic and diastolic dysfunction, adaptive pro-hypertrophic effect, decreased markers of inflammation, improved metabolic health, decreased risk of heart failure (HF) and improved overall survival.^{4–8} Physical inactivity is associated with obesity, diabetes, hypertension and the metabolic syndrome.^{9–11} All these conditions share several underlying mechanisms that result in increased HF risk; insulin resistance, impaired energy availability, endothelial dysfunction, oxidative stress, which ultimately leads to increased myocardial stiffness, left ventricular (LV) hypertrophy (LVH), diastolic dysfunction and an increased propensity for coronary artery disease.^{12–14}

A majority of previous epidemiological studies have been limited by a lack of objective measures on PA and sedentary behaviour (SB). A prior study that used accelerometry data only included a large community-based sample of young adults to middle-aged men and women.^{6,7} Additionally, the Hispanic/Latino population is understudied, with a limited number of large-scale analyses of this population, particularly those that consider the wide array of ethnic, cultural and geographic diversity.

Echocardiographic Study of Latinos (ECHO-SOL), an ancillary echocardiographic study of the Hispanic Community Health Study/Study of Latinos (HCHS/SOL), is a multicentre epidemiological study designed to include a diverse Hispanic/Latino subgroup representation and provided a unique opportunity to explore the cross-sectional association between accelerometer-measured PA, SB, and cardiac structure and function as measured by transthoracic echocardiography (TTE). This study's primary aim was to compare echocardiographic measures of cardiac structure and function with accelerometer measured PA and SB. We hypothesised that higher PA would be favourably associated, and SB would be negatively associated with LV structure and function.

METHODS

Study setting

HCHS/SOL is a prospective population-based longitudinal cohort study (n=16,415, mean age 43 years) of self-identified Hispanic/Latino individuals, 18–74 years of age enrolled from 2008 to 2011.¹⁵ This study was designed to assess chronic disease in Hispanic/Latino individuals living in four cities in the USA: Bronx, New York; Chicago, Illinois; Miami, Florida; and San Diego, California. Participants were recruited in communities surrounding four field centres in the respective cities. The details of the HCHS/SOL sampling method have been previously described.¹⁵

ECHO-SOL is an ancillary study of HCHS/SOL consisting of 1818 participants recruited through a

stratified sampling process representative of the parent study.¹⁶ ECHO-SOL was designed to characterise cardiac structure and function using TTE in a representative sample of Hispanic/Latino individuals living in the USA. The baseline TTE examination for the ECHO-SOL ancillary study was performed from 2011 to 2014.¹⁶ Inclusion criteria for ECHO-SOL included: age 45 years or older; self-reported Hispanic/Latino of Mexican, Puerto Rican, Cuban, Dominican, Central American or South American background; and enrolment 36 months or fewer from the date of the baseline HCHS/SOL visit. ECHO-SOL enrolled, on average, ~80% of eligible participants at each of the study sites. Inclusion and exclusion criteria for ECHO-SOL was previously described.^{16,17}

The current analysis excluded participants who did not wear the accelerometer or were not adherent to the accelerometer protocol (n=465; at least 3 days with 10 hours of wear time each). We excluded those with poor quality data on at least one echocardiographic variable (n=39) and participants with prior CV disease (CVD) for our analysis (n=108). These exclusions yielded an analytic sample of n=1206 (66% of participants). The baseline characteristics of all 1818 ECHO-SOL participants were previously described.¹⁷

STUDY MEASURES

PA measures

The Actical accelerometer (version B-1 model 198-0200-03; Respironics Co, Bend, Oregon) was used to assess PA and SB. The accelerometer data were collected at the baseline examination visit. During the HCHS/SOL baseline clinic visit, participants were fitted with a belt and left the clinic wearing the accelerometer above the iliac crest. Participants were instructed to perform normal daily activities and wear the accelerometer only during awake hours and remove it for sleep, showering and swimming. The Actical device was programmed to measure accelerations in counts per 1 min epochs based on the convention used in previous studies.¹⁸ The time used for analysis started at 05:00 the morning after the initial clinic visit and ended at midnight on the sixth day of wear. Non-wear was defined as consecutive zero counts for at least 90 min.¹⁸ PA threshold levels were defined as follows: SB <100 counts/min, light 100–1534 counts/min, moderate 1535–3961 counts/min and vigorous \geq 3962 counts/min.^{19,20} For the present analysis, data for moderate to vigorous PA (MVPA) were combined, denoted as MVPA, because very few people had a significant amount of vigorous activity. Adherence and accelerometer performance have been previously described.¹⁸ Adherence was defined as \geq 10 hours/day of wear time for at least 3 days of wear. Time spent in PA by intensity level was calculated by summing the minutes in a day where the counts were within each threshold and then averaging across adherent days.

The HCHS/SOL assessed self-reported PA in a typical week using an interviewer-administered, modified Global

PA Questionnaire that included questions on three activity domains (work, transport and leisure) available at www.int/chp/steps/GPAQ/en. Participants were asked to think about activities that lasted at least 10 min in a typical week. We only included accelerometer-measured PA for our analysis.

Echocardiography measures

The average time between the accelerometer data and echocardiographic measures was 815 ± 196 days. A complete description of the echocardiographic measures of cardiac structure and function was previously reported.^{16 17 21} Briefly, a standardised TTE ultrasound examination, including M-mode, two-dimensional, spectral, colour flow and tissue Doppler, was performed by experienced Registered Diagnostic Cardiac Sonographers at each of the four parent study field sites using Philips IE-33 or Sonos 7500 scanners interfaced with a standard 2.5–3.5 MHz phased-array probe, according to American Society of Echocardiography (ASE) recommendations.²² Echocardiograms were analysed and interpreted centrally at Wake Forest School of Medicine (Winston-Salem, North Carolina, USA). Inter-reader and intra-reader reproducibility was assessed and previously reported.¹⁶ Two-dimensional imaging of the LV was performed to obtain the best possible images of the LV endocardium without foreshortening of the LV cavity or echo drop out. Using the apical 4-chamber and 2-chamber views, LV end-diastolic volumes (EDV) and end-systolic volumes (ESV) were derived using the biplane method of discs, as per the ASE recommended methodology.²² The following echocardiographic measured and derived variables were used: LV mass (LVM) was determined by subtracting the LV endocardial cavity volume from the LV epicardial volume and multiplying the resultant myocardial volume by the myocardial density (1.05 g/mL) then indexing based on the body surface area or height^{2,7}. LV geometry was predicated on the measurement of LVM, LV posterior wall thickness at diastole (LVPWd), interventricular septal thickness at diastole and relative wall thickness (RWT) defined as $2 \times (\text{LVPWd} / \text{LV internal diameter at diastole})$. LV geometry was defined as normal (normal LVM and normal RWT) or abnormal with abnormal LV geometry categorised as follows: concentric remodelling (normal LVM and increased RWT), eccentric hypertrophy (increased LVM and normal RWT) and concentric hypertrophy (increased LVM and increased RWT).²² LV ejection fraction (LVEF) = $(\text{EDV} - \text{ESV}) / \text{EDV}$. LVEF could not be ascertained in 4.9% of the cohort because of image quality. Our assessment of LV diastolic dysfunction (LVDD) has been previously published.²¹ Right ventricular (RV) systolic function and tricuspid valve regurgitation velocity (TRV) were assessed according to ASE guidelines. Speckle-tracking strain analysis was performed offline using the TomTec Cardiac Performance Analysis package on two-dimensional images acquired, and methods were previously described.¹⁶

Primary structural and functional measures included LVM indexed to height^{2,7} (LVMI), RWT, left atrial volume (LAV), LAV index (LAVI) and EF, mid-LV segment global circumferential strain (GCS), global longitudinal strain (GLS) and LV diastolic functional measures. Secondary cardiac measures included LV stroke volume (LVSV), lateral peak systolic annular velocity (lat peak S'), RV fractional area change (RV FAC), TRV, tricuspid annulus systolic velocity (RV S' wave), RV velocity-time integral (RV VTI), RVSV and tricuspid annular plane systolic excursion (TAPSE).

Clinical covariates

All measures were taken from the baseline visit (2008–2011). Trained technicians measured each participant's height and weight twice and then averaged these two measures to calculate body mass index ($\text{BMI} = \text{weight (kg)} / \text{height (m)}^2$). Participants with a BMI of $\geq 30 \text{ kg/m}^2$ were categorised as obese. Three seated blood pressure (BP) measures were taken using an automatic sphygmomanometer averaging the last two measures. Hypertension was defined as a systolic BP of 140 mm Hg or greater, diastolic BP of 90 mm Hg or greater, or receiving antihypertensive medication. Participants were classified as having hypercholesterolaemia if they were currently using the cholesterol-lowering medication, had low-density lipoprotein levels $\geq 160 \text{ mg/dL}$, or total cholesterol $\geq 240 \text{ mg/dL}$. Total and high-density lipoprotein (HDL) cholesterol was measured with a direct magnesium/dextran sulfate method. Plasma glucose was measure fasting and after a 2-hour glucose load. Diabetes mellitus was a fasting plasma glucose of 126 mg/dL or greater, 2-hour-post load glucose of 200 mg/dL or greater, glycosylated haemoglobin 6.5% or greater, or use of antihyperglycaemic medications. Self-reported current smoking history referring to cigarette smoking and current alcohol intake referring to alcohol history was collected via questionnaire.²³

Healthy subgroup ($n=460$) was defined as not having hypertension (systolic BP $\geq 140 \text{ mm Hg}$, diastolic BP $\geq 80 \text{ mm Hg}$ or history of drug-treated hypertension), diabetes (history of a diagnosis of diabetes mellitus, fasting glucose $\geq 126 \text{ mg/dL}$ or glycosylated haemoglobin $\geq 6.5\%$), coronary heart disease (self-reported history of revascularisation, myocardial infarction, angina or ECG defined evidence of myocardial infarction) or obesity ($\text{BMI} \geq 30 \text{ kg/m}^2$).

Statistical methods

The primary aim of this study was to compare echocardiographic measures of cardiac structure and function in participants with higher levels of PA and SB. The dependent variables were echocardiographic measures of cardiac structure and function. SB, light PA and MVPA were the primary independent variables. Cohort characteristics at baseline were described using means and SEs for continuous variables and frequencies (percentages) for categorical variables. SB, light PA and MVPA, and all echocardiographic measures were treated as continuous variables.

Hypertension, diabetes and current cigarette smoking were treated as categorical variables. The means of echocardiographic measures of cardiac structure and function were compared across categories of PA (MVPA, light PA) and SB using multivariable linear regression models. Age, sex, hypertension, antihypertensive medications, diabetes, current cigarette smoking and HDL cholesterol were included as covariates in the analysis. Covariates were chosen based on known associations with ventricular size and heart disease, including demographics and anthropometric variables, and variables reflecting comorbidities. Model 1—demographics, model 2—model 1 plus continuous systolic BP and antihypertension medication use, model 3—model 2 plus other CV disease modifiable factors (diabetes mellitus, current smoking and continuous HDL). We divided the light PA, MVPA and SB into quartiles (Q1–Q4) and compared the echocardiographic variables between the highest quartile (Q4) to other quartiles by univariate analysis.

Estimated beta coefficients and p values based on the F statistic were presented. Statistical significance was designated as a p value ≤ 0.01 with no adjustments for multiple comparisons. SAS SURVEY procedures were used, and all reported values were weighted to adjust for sampling probability and non-response bias to make the inference applicable to the ECHO-SOL target population. All analyses were performed using SAS V.9.4 (SAS Institute).

RESULTS

Baseline characteristics of the study population are provided in [table 1](#). The mean age of included participants was 56 years; 57% were women, 28% had diabetes, 50% had hypertension and 18% reported current cigarette smoking. On average, participants accumulated (mean \pm SD) 21 \pm 1.1 min/day of MVPA; 217 \pm 4.2 min/day of light PA and 737 \pm 8.1 min/day of SB. Men accumulated more MVPA compared with women (27 \pm 1.6 vs 17 \pm 1.3 min/day, $p < 0.001$) ([figure 1](#)). Participants aged > 65 years had less MVPA accumulation than participants aged < 55 years (17 \pm 3.0 vs 24 \pm 1.5, $p = 0.03$). Overall, participants aged > 55 years spent less time in both light PA and MVPA than those less than 55 years of age ([figure 2](#)). Online supplemental table 1 shows the prevalent self-reported PA in ECHO/SOL.

Associations between PA and echocardiographic measures

Light PA

Compared with less light PA time (quartile 1), more light PA (quartile 4) was associated with decreased LVMI/EDV ratio ($p < 0.001$) and a decrease in E/e' ratio ($p < 0.001$) (online supplemental table 2). In multivariable linear regression models between light PA and echocardiographic measures (model 3), every 10 min/day of light PA was associated with a 0.03 mL/m² increase in LAVI ($p < 0.01$) and a 0.004 cm increase in TAPSE ($p < 0.01$). There were no statistically significant associations between light PA and other echocardiographic measures ([table 2](#)).

Table 1 Baseline characteristics

Variables	n=1206
Age, years	56.4 \pm 0.36
Female, %	691(57)
Body mass index (kg/m ²)	30.1 (0.1)
Body surface area (m ²)	1.8 (0.1)
Systolic BP, mm Hg	136 (1.0)
Diastolic BP, mm Hg	78 (0.7)
Diabetes, %	343 (28)
Hypertension, %	603 (50)
Hypercholesterolaemia, %	626 (49)
HDL cholesterol, mg/dL	50.4 (0.7)
Alcohol use, %	903 (75)
Current cigarette smoking, %	212 (18)
Echocardiographic measurements	
Left ventricular mass index (LVMI, g/m ^{2.7})	41.5 (0.35)
Relative wall thickness, ratio	0.40 (0.003)
LVMI ^{2.7} /end-diastolic volume	0.52 (0.006)
Left atrial volume index, mL/m ²	23.02 (0.25)
Left ventricular ejection fraction, %	59.80 (0.20)
E/e', ratio	9.96 (0.12)
Global longitudinal strain, %	-17.56 (0.13)
Global circumferential strain, %	-24.33 (0.29)
LV stroke volume, mL	70.30 (0.61)
Lateral peak S velocity, cm/s	8.30 (0.08)
RV peak S velocity, cm/s	11.88 (0.09)
RV fractional area change, %	0.45 (0.004)
TR max velocity, cm/s	206.35 (1.69)
RV velocity-time integral, cm	17.42 (0.15)
TAPSE, cm	2.09 (0.02)
Physical activity and sedentary behaviour	
MVPA, min/day	20.93 \pm 1.1
Light PA, min/day	216.84 \pm 4.2
Sedentary behaviour, min/day	736.50 \pm 8.1

N (%) using weighted row percentages; N's presented are weighted counts.

Data presented in means \pm SE of mean.

BP, blood pressure; E, peak early diastolic transmitral inflow velocities; e', average of mitral early diastolic septal and lateral annular velocities; HDL, high-density lipoprotein; LV, left ventricular; MVPA, moderate to vigorous-intensity PA; n, number; PA, physical activity; RV, right ventricular; TAPSE, tricuspid annular plane systolic excursion; TR, tricuspid regurgitation.

Moderate to vigorous PA

Compared with less MVPA (quartile 1), greater MVPA (quartile 4) was associated with decreased LVMI/EDV ratio ($p < 0.001$), increased LAVI ($p < 0.001$) and decrease in E/e' ($p < 0.001$) ratio (online supplemental table 3). In a multivariable linear regression model between PA and echocardiographic measures (model 3), every

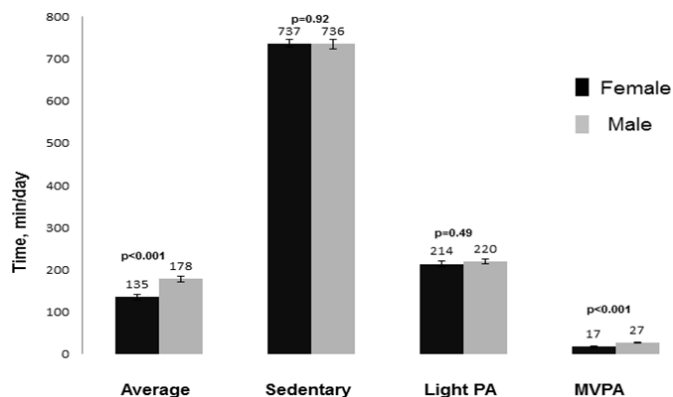


Figure 1 Physical activity and sedentary behaviour by gender. *Data presented in means±SE of mean. The number above the bar shows mean activity in min/day. MVPA, moderate to vigorous-intensity PA; PA, physical activity.

10 min/day of MVPA was associated with a 0.18 mL/m² increase in LAVI (p<0.01) and a 0.24% improvement in GCS (p<0.01). No statistically significant associations were noted between MVPA and other echocardiographic measures (table 3).

SB and echocardiographic measures

Compared with lower SB (quartile 1), higher SB (quartile 4) was associated with increased LVMI (p=0.008) (online supplemental table 4). In multivariable linear regression models between SB and echocardiographic measures (model 3), there were no statistically significant associations between SB and any of the echocardiographic measures (table 4). Online supplemental table 5 shows an unadjusted association between PA and echocardiographic measures.

PA and echocardiographic measures in healthy subgroup

In a multivariable linear regression model between PA and echocardiographic measures (model 3), both MVPA and light PA was associated with increased LAVI, improved GLS, improved E/e ratio, improved RV function (increase in TAPSE); MVPA was associated

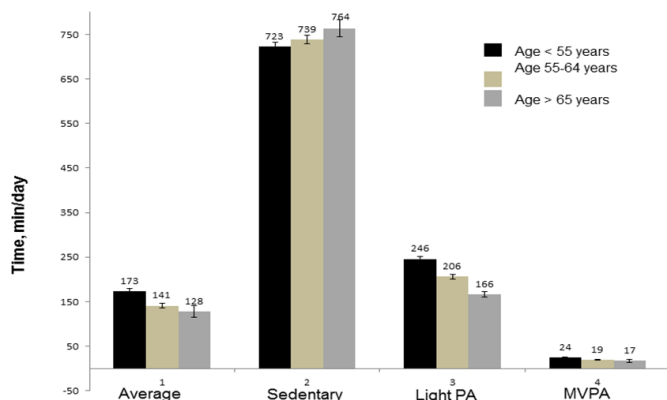


Figure 2 Physical activity and sedentary behaviour by age. Data presented in means±SE of mean. The number above the bar shows mean activity in min/day. MVPA, moderate to vigorous-intensity PA; PA, physical activity.

Table 2 Multivariable linear regression for the association between light PA and echocardiogram parameters

Outcome	Model 1	Model 2	Model 3
LVMI, g/m ^{2.7}	-0.02882	-0.00597	0.01003
LVMI ^{2.7} /EDV	-0.00157	-0.00135	-0.00108
RWT, ratio	0.00014	0.00021	0.00030
LAVI, mL/m ²	0.02431*	0.03206*	0.02840*
EF, %	0.01336	0.01369	0.00728
E/e', ratio	-0.00843	-0.00521	0.00086
GLS, %	-0.02085	-0.01784	-0.01333
GCS, %	-0.02332	-0.02139	-0.01778
SV, mL	-0.02847	-0.03209	-0.03206
RV peak S, cm/s	0.01126	0.01249	0.01287
RV FAC, %	0.00048	0.00051	0.00034
TRV Max, cm/s	0.20262	0.18327	0.13431
TAPSE, cm	0.0037*	0.0040*	0.0037*

M1: demographics (age and sex).

M2: M1+systolic blood pressure, antihypertensive medication use.

M3: M2+other cardiovascular disease modifiable factors (diabetes mellitus, current smoking and continuous high-density lipoprotein). Values are per 10 min/day light PA.

*P value <0.01.

E, peak early diastolic transmitral inflow velocities; EDV, end diastolic volume; e', average of mitral early diastolic septal and lateral annular velocities; EF, ejection fraction; GCS, global circumferential strain; GLS, global longitudinal strain; LAVI, left atrial volume index; LVMI, left ventricular mass index; M, model; PA, physical activity; RV, right ventricular; RV FAC, RV fractional area change; RWT, relative valve thickness; SV, stroke vol; TAPSE, tricuspid annular plane systolic excursion; TRV, tricuspid regurgitation velocity.

with greater SV (online supplemental table 6). SB was associated with worsening E/e, GLS and RV function (decrease in TAPSE). SB was negatively correlated with SV.

We performed additional analysis to find the correlation between MVPA and SB. The correlation between sedentary time and MVPA is very weak with a Pearson correlation coefficient of -0.1 (p=0.26) and a very flat slope on the scatterplot of the two variables.

DISCUSSION

In this large cohort of diverse Hispanic/Latinos, we found that higher light PA was associated with increased LAVI and better RV systolic function after adjustment for potential confounders. Similarly, higher MVPA was associated with greater LV myocardial mechanics as assessed by strain and increased LAVI. Both higher levels of light PA and MVPA (min/day) were associated with decreased LVMI/EDV, indicative of favourable LV remodelling and decrease E/e' ratio, a sign of decrease LA pressure. Higher levels of SB were associated with increased LVM. In the healthy subgroup, both MVPA and light PA were associated with greater LV myocardial mechanics, decrease LA pressure, and better RV function. SB was associated with

Table 3 Multivariable linear regression for the association between MVPA and echocardiogram measures

Outcome	Model 1	Model 2	Model 3
LVMI, g/m ²	0.0411	0.1318	0.1381
LVMI ^{2.7} /EDV	-0.0014	-0.0006	-0.0006
RWT, ratio	-0.0005	-0.0001	0.0001
LAVI, mL/m ²	0.1707*	0.1808*	0.1792*
EF, %	0.0578	0.0448	0.0389
E/e', ratio	0.0037	0.0221	0.0259
GLS, %	-0.069	-0.0521	-0.0539
GCS, %	-0.2477*	-0.2397*	-0.2397*
SV, mL	0.3277	0.3177	0.3174
RV peak S, cm/s	0.0589	0.0590	0.0615
RV FAC, %	0.0021	0.0019	0.0019
TRV Max, cm/s	1.1367	1.1386	1.1597
TAPSE, cm	0.0084	0.0082	0.0078

M1: demographics (age and sex).

M2: M1+systolic blood pressure, antihypertensive med use.

M3: M2+other cardiovascular disease modifiable factors (diabetes mellitus, current smoking and high-density lipoprotein).

Values are per 10 min/day light MVPA.

*P value <0.01.

E, peak early diastolic transmitral inflow velocities; EDV, end diastolic volume; e', average of mitral early diastolic septal and lateral annular velocities; EF, ejection fraction; GCS, global circumferential strain; GLS, global longitudinal strain; LAVI, left atrial volume index; LVMI, left ventricular mass index; M, model; MVPA, moderate to vigorous PA; RV, right ventricular; RV FAC, RV fractional area change; RWT, relative valve thickness; SV, stroke volume; TAPSE, tricuspid annular plane systolic excursion; TRV, tricuspid regurgitation velocity.

worsening LV myocardial mechanics, worsening LA pressure, and RV function.

Although increased PA has been associated with higher LVM, LA size and LV EDV in multiple prior studies,^{5 7 24 25} to our knowledge, this is the first study to examine the association between accelerometry measured MVPA and improved LV myocardial mechanics. While there was no clinically significant association between LVEF and MVPA, LV GCS was significantly more favourable with MVPA. GCS reflects the change in length around the LV perimeter during systole as the myocardium shortens and the LV cavity circumference reduces. During incremental dynamic aerobic exercise, studies reported a progressive increase in GCS due to increased apical circumferential strain (ACS) with trivial effects on basal circumferential strain (BCS).²⁶ In contrast, GCS decreased during the isometric exercise with no effect on ACS and decreased BCS from rest.²⁶ Therefore, exercise modality plays some role in GCS responses, and we did not have a granular breakdown of PA type. Further studies of global and region-specific circumferential strain during incremental exercise are lacking at present, and GCS responses during dynamic and static modalities remain incompletely understood.

Table 4 Multivariable linear regression for the association between sedentary behaviour and echocardiogram measures

Outcome	Model 1	Model 2	Model 3
LVMI, g/m ^{2.7}	0.0626*	0.0485	0.0433
LVMI ^{2.7} /EDV	0.0008	0.0006	0.0005
RWT, ratio	0.0000	-0.0002	0.0000
LAVI, mL/m ²	-0.0066	-0.0066	-0.0035
EF, %	-0.0025	-0.0005	0.0025
E/e', ratio	0.0144	-0.0074	-0.0341
GLS, %	0.0041	0.0035	0.0014
GCS, %	-0.0377	-0.0390	-0.0413
SV, mL	0.0073	-0.0039	0.0001
RV peak S, cm/s	0.0090	0.0075	0.0068
RV FAC	0.0001	0.0001	0.0002
TRV max, cm/s	-0.0457	-0.0459	-0.0330
TAPSE, cm	-0.0010	-0.0011	-0.0010

M1: demographics (age and sex).

M2: M1+systolic blood pressure, antihypertensive med use.

M3: M2+other cardiovascular disease modifiable factors (diabetes mellitus, current smoking and high-density lipoprotein).

Values are per 10 min/day of sedentary behaviour.

*P value <0.01.

E, peak early diastolic transmitral inflow velocities; EDV, end diastolic volume; e', average of mitral early diastolic septal and lateral annular velocities; EF, ejection fraction; GCS, global circumferential strain; GLS, global longitudinal strain; LAVI, left atrial volume index; LVMI, left ventricular mass index; M, model; RV, right ventricular; RV FAC, RV fractional area change; RWT, relative valve thickness; SV, stroke vol; TAPSE, tricuspid annular plane systolic excursion; TRV, tricuspid regurgitation velocity.

Light PA was associated with better RV systolic function. Besides, higher quartiles of MVPA were also associated with better RV systolic function. Although these findings' implications are unclear, even small absolute changes in the very thin-walled RV in disease-free participants may have significant long-term effects. Aaron *et al* found that higher levels of both MVPA and intentional exercise were associated with larger RV stroke volume without a change in RVEF.²⁷ Recent studies of patients with pulmonary arterial hypertension have shown that exercise training is safe and can improve exercise endurance, 6 min walk distance and quality of life.²⁸ Recently, it has been noted that higher cardiorespiratory fitness (CRF) was associated with improved RV systolic function (TAPSE, RV S' wave).²⁹ In addition, less decline in PA over time was associated with better RV systolic function. Structural and functional LV abnormalities may contribute to abnormalities in RV function. LV systolic or diastolic dysfunction leads to elevated LV end-diastolic pressure, which may cause pulmonary venous hypertension and high pulmonary artery systolic pressure, and RV systolic dysfunction. In addition, HF risk factors can adversely affect both RV and LV function such that subclinical abnormalities in RV function may precede and contribute to the

development of HF, especially in HFpEF. Taken together, such exercise-induced adaptive changes in RV may be beneficial with potentially important implications for diseases characterised by pulmonary hypertension and RV dysfunction. These data may be useful in studying the RV-specific benefits of exercise in at-risk populations.

Increases in LA size associated with higher CV fitness and MVPA have been described.⁴⁻⁷ In the general population, however, increased LA size is associated with many pathological conditions, and it is usually the result of two processes, either increased volume or increased cardiac filling pressures. Interestingly, similar to our findings, the Cooper Clinic Longitudinal study observed no apparent association between diastolic dysfunction and LA size among high-fit individuals.⁴ Our findings and those reported by others suggest that LA dilation mechanisms may reflect physiology unrelated to diastolic function (ie, greater cardiac output) with higher PA.⁴⁻⁷

Recently Dallas Heart Study, which included 1368 participants (age, 49 years; 40% men, free of CVD), showed that objective measures of MVPA (assessed by accelerometer) were positively associated with more favourable LV structure and function parameters (assessed cardiac MRI), with better LV contractility, higher SV, lower arterial elastance, and LVH without myocardial injury, independent of traditional CVD risk factors.⁷ Vigorous PA was also positively associated with higher SV and lower effective arterial elastance measures and greater likelihood of both LVH without myocardial injury and malignant LVH. The moderate PA levels and SB were not associated with LV structure and function in the most adjusted analysis. Higher quartiles of MVPA and light PA were associated with favourable LV remodelling and improved diastolic function.

Improved diastolic function, increased LA size, improved LV mechanics (GCS), and RV systolic function further support that physiological adaptation with increased cardiac size characterised by normal cardiac morphology with a normal and enhanced cardiac function. This is similar to the recent Cooper Clinic Longitudinal study, which showed that higher cardiac fitness levels were independently associated with increased LA size, LV internal diameter at diastole, and decreased RWT and decreased E/e' .⁴ Similar to our study, there was no positive association between higher fitness and LVMI. This evidence taken together with our study confirms that staying active is more strongly associated with favourable measures of cardiac structure and function even at the level of light PA. Our improved LV strain, LV remodelling, LA volume, and improved diastolic function align with Dallas heart study results.⁷ These results further support the physiological benefits of long-term aerobic exercise and PA on cardiac structure and function. Importantly prior study also showed that only MVPA but not sedentary time were significantly associated with CRF levels.

While LVM, RV mass, LV volumes tend to decrease with bed rest or deconditioning.³⁰ The chronic consequences of SB (increased obesity or obesity-related

comorbidities, diabetes and hypertension) are generally associated with increased LVM and worse cardiac function.³¹⁻³³ Thus, it is unclear whether and in which direction SB would be related to LV structure and function independently of PA, obesity and obesity-related comorbidities. A recent community-based study that included middle-aged adults free from overt clinical CV disease found that higher sedentary screen time was associated with increased LVMI, mass-to-volume ratio and RWT in Caucasians but not African Americans.³⁴ Similarly, low fitness was independently associated with smaller heart size and concentric remodelling patterns in the Cooper Clinic Longitudinal Study.⁴

Taken together from ours and other studies, it has been noted that PA or higher CRF was associated with greater LV myocardial mechanics (LV strain), decreased LVMI/EDV (indicative of favourable LV remodelling), decrease E/e' ratio (a sign of decrease LA pressure), improved myocardial mechanics, lower arterial stiffness and improved RV function, which further supports the role of long-term aerobic exercise in preventing HF.^{29 35-37} Besides, SB has been associated with higher HF risk and a greater burden of abnormalities in LV structure. Furthermore, low CRF also explained up to 40% of HF's high body mass index associated risk.³⁵ This is very important with HFpEF since there is no proven benefit with pharmacological therapy in HFpEF except exercise training and calorie restriction and obesity is parallel with the HFpEF epidemic.³⁸ While there is a large base of data that supports the CV benefits of PA and the detrimental effect of SB, these effects are broad and pleiotropic. PA is associated with various CV health benefits; decrease in BP, improved metabolic health, improved vascular function, decrease vascular stiffness, improved endothelial function, prevention of age-related cardiac remodelling, systolic and diastolic dysfunction, and adaptive pro-hypertrophic effect.^{4-7 39 40}

It is hypothesised that improved vascular function is due to reduced oxidative stress and subsequent improved endothelial dysfunction from PA. A separate study demonstrated that exercise could return the expected age-associated vascular tone to a more vasodilated state equivalent to that of younger subjects.⁴⁰

Strength and limitations

The sizeable community-based sample, comprehensive assessment of PA (including SB and light PA) and echocardiographic measurements (including diastolic function, strain analysis, RV function) strengthen our investigations. Our study is also unique since this included an understudied high-risk population with a high prevalence of CV disease, risk factors and comorbidities. Most prior studies rely on the self-reported assessment of PA and SB. Using accelerometers to objectively quantify PA and SB reduces measurement error and allows for a more precise categorisation of the degree and amount of PA and SB.

Nonetheless, several limitations merit comment. The cross-sectional nature of this analysis prevents any causal inferences. Although we adjusted all models for multiple

variables, the influence of residual confounding cannot be excluded. The accelerometry data were obtained prior to echocardiography, and that can result in underestimating the strength of the associations between MVPA and cardiac measurements (regression dilution bias). The hip-worn accelerometer is likely to underestimate certain activities such as bicycling and swimming (since they removed it when in the water). Additionally, accelerometry was assessed for only 1 week. While this is standard in the field,⁴¹ and others have shown it may reflect longer-term patterns of PA and SB among women,⁴² a longer measurement period would be an improvement.

Here we chose not to adjust the p value for multiple comparisons because each outcome was selected based on a priori hypothesis and a unique mechanism between exposure and each cardiac parameter. We were interested in controlling the type I error rate per the outcome of interest rather than per group of outcome variables. We only rejected the null hypothesis with a conservative p value of less than 0.01. Nevertheless, the results should be interpreted cautiously and validated in other studies. Several of our findings may be considered hypothesis-generating and warrant replication in subsequent studies. The sample size of ECHO-SOL is modest, limiting type II error. The type of PA (weight-training, cardio or isometric) may influence a different cardiac response (with GCS or other parameters). We did not have a PA-type breakdown for accelerometry data. Besides, accelerometry does not capture resistance training.

CONCLUSION

In a community-based population with multiple risk factors for developing LV dysfunction, both MVPA and light PA were associated with favourable cardiac mechanics after adjusting for major CV risk factors. Future studies of PA and cardiac function with a larger sample and a prospective longitudinal design are needed to assess cardiac function with clinical outcomes better.

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the reliability and freedom from bias of the data presented and their discussed interpretation.

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Supplemental Table 1. Prevalent self-reported physical activity in ECHO SOL

Activity	Prevalence
Vigorous sports /activities	(n=131)
American football	4(2%)
Basketball	22(15%)
Boxing	11(6%)
Lacrosse	1(0.4%)
Soccer	25(17%)
High-intensity aerobics	22(13%)
Hard lap swimming	17(12%)
Running	65(50%)
Weight lifting	38(27%)
Cycling	27 (16%)
Moderate Sports/Activities	(n=356)
Aerobic dance	38 (6%)
Baseball	22(5%)
Brisk walking	259 (75%)
Canoeing/Kayaking	5(2%)
Coaching sports	15(3%)
Light cycling	73(21%)
Weight lifting	53(14%)
Swimming laps	46(12%)
Table tennis	8(2%)
Rock climbing	1(0.2%)
Skating	4(2%)
Hunting/Fishing	20(5%)
Karate/Judo/Other Martial arts	7(1%)
Frisbee play	6(1%)
Stationary exercise	82(19%)

All proportions are weighted to account for the survey design

Supplemental Table 2. Echocardiographic measures with quartiles of light PA

Light PA	Q1: <159 Min/day	Q2: 159-208 Min/day	Q3: 208-273 Min/day	Q4: >273 Min/day	p value Q4 vs. Q1	p value Q4 vs. Q2
LVMI, g/m ^{2.7}	42.5±0.9	41.7±0.80	41.7±0.68	40.7±0.58	0.081	0.300
LVMI ^{2.7} / EDV	0.58±0.02	0.53±0.01	0.51±0.01	0.49±0.01	< 0.001	0.050
RWT, ratio	0.40±0.00	0.40±0.00	0.39±0.00	0.40±0.00	0.456	0.530
LAVI, ml/m ²	22.6±0.21	23.3±0.24	23.9±0.19	23.0±0.15	0.090	0.200
EF, %	60.0±0.34	59.7±0.34	59.9±0.39	59.8 ±0.33	0.673	0.700
E/e', ratio	10.6±0.2	9.6±0.2	9.6±0.2	9.5±0.2	<0.001	0.760
GLS, %	-17.3±0.24	-17.5±0.19	-17.3±0.19	-18.1±0.27	0.012	0.064
GCS, %	-24.0±0.53	-24.4±0.41	-23.7±0.51	-25.0±12.3	0.135	0.280
RV peak S, cm/sec	11.6±0.15	11.8±0.18	12.0±0.16	12.3± 0.19	0.010	0.050
TAPSE, cm	17.2±0.21	17.1±0.27	17.2±0.20	17.7±0.41	0.223	0.262
RV FAC, %	0.44±0.00	0.45±0.00	0.45±0.00	0.45±0.00	0.287	0.870
TRV Max, cm/sec	211.3±3.2	204.0±3.3	203.1±2.6	207.7±3.2	0.427	0.422

*Data presented in means +/- standard errors of mean

PA =physical activity, Q=quartile, LVMI =left ventricular mass index, EDV=end diastolic volume, RWT=relative valve thickness, LAVI=left atrial volume index, EF=ejection fraction, E =peak early diastolic transmitral inflow velocities, e' =average of mitral early diastolic septal and lateral annular velocities, GLS =global longitudinal strain, GCS = global circumferential strain, RV=right ventricular, RVFAC= RV fractional area change, TAPSE = tricuspid annular plane systolic excursion; TRV= tricuspid regurgitation velocity.

p value is calculated based on quartile 4 to other quartiles
p ≤ 0.01 considered significant

Supplemental table 3. Echocardiographic measures with quartiles of MVPA

MVPA	Q1: <5 Min/day	Q2: 5-13 Min/day	Q3: 13-27 Min/day	Q4: >27 Min/day	p value Q4 vs. Q1	p value Q4 vs. Q2
LVMI, g/m ^{2.7}	43.8±0.80	41.7±0.88	41.0±0.68	41.2±0.62	0.127	0.688
LVMI ^{2.7} / EDV	0.58±0.02	0.54±0.01	0.51±0.01	0.48±0.01	< 0.001	0.001
RWT, ratio	0.41±0.00	0.40±0.00	0.40±0.00	0.39±0.00	0.031	0.150
LAVI, ml/m ²	22.7±0.2	22.8±0.2	23±0.2	24.2±0.2	< 0.001	< 0.001
EF, %	60.1±0.36	59.6±0.51	60.3±0.34	59.5±0.40	0.248	0.888
E/e', ratio	10.6±0.2	10.1±0.2	9.4±0.2	9.4±0.2	< 0.001	0.005
GLS, %	-17.6±0.24	-17.3±0.21	-17.8±0.21	-17.6±0.28	0.868	0.300
GCS, %	-23.4±0.6	23.7± 0.4	-24.9± 0.4	25.2 ±0.6	0.010	0.024
RV peak S, cm/sec	11.6±0.18	11.6±0.15	12.1± 0.15	12.2± 0.17	0.010	0.003
TAPSE, cm	17.5±0.24	17.2±0.24	17.1±0.22	17.5±0.42	0.983	0.642
RV FAC, %	0.44±0.01	0.44±0.00	0.45±0.00	0.46±0.00	0.033	0.031
TRV Max, cm/sec	209.1±3.6	208.6±3.4	206.±0 2.6	208.6±3.5	0.920	0.211

*Data presented in means +/- standard errors of mean

MVPA = moderate to vigorous-intensity physical activity; Q=quartile, LVMI =left ventricular mass index, EDV=end diastolic volume, RWT=relative valve thickness, LAVI=left atrial volume index, EF=ejection fraction, E =peak early diastolic transmitral inflow velocities, e' =average of mitral early diastolic septal and lateral annular velocities, GLS =global longitudinal strain, GCS = global circumferential strain, RV=right ventricular, RVFAC= RV fractional area change, TAPSE = tricuspid annular plane systolic excursion; TRV= tricuspid regurgitation velocity.

p value is calculated based on quartile 4 to other quartiles
p ≤ 0.01 considered significant

Supplemental Table 4. Echocardiographic measures with quartiles of sedentary behavior

Sedentary behavior	Q1: < 604 min/day	Q2: 604-748 min/day	Q3: 748-884 min day	Q4: > 884 min/day	p value Q4 vs. Q1	p value Q4 vs. Q2
LVMI, g/m ^{2.7}	41±0.6	40.4±0.6	41.8±0.7	44.0±0.9	0.008	0.002
LVMI ^{2.7} / EDV	0.52±0.01	0.51±0.01	0.53±0.01	0.56±0.01	0.029	0.020
RWT, ratio	0.40±0.00	0.40±0.00	0.40±0.00	0.40±0.00	0.314	0.530
LAVI, ml/m ²	23.5±0.6	23.1±0.19	22.9±0.23	23.2±0.3	0.381	0.851
EF, %	59.8±0.49	59.7±0.36	60.1±0.34	60.0±0.32	0.804	0.552
E/e', ratio	9.6±0.23	10.0±0.18	9.9±0.19	10.0±0.16	0.117	0.837
GLS, %	-17.5±0.25	-17.6±0.26	-17.6±0.21	-17.3±0.18	0.620	0.476
GCS, %	-23.4±0.51	-24.1±0.62	-24.8± 0.40	-25.1±0.51	0.010	0.190
RV peak S, cm/sec	11.5±0.17	11.8±0.19	12.3±0.16	12.0±0.19	0.080	0.364
TAPSE, cm	17.5±0.19	17.6±0.37	17.4±0.23	16.6±0.26	0.022	0.042
RV FAC, %	0.44±0.00	0.45±0.00	0.46±0.00	0.45±0.00	0.357	0.780
TRV Max, cm/sec	205.4±2.4	211.2±3.3	203.5±3.2	205.5±3.6	0.984	0.255

*Data presented in means +/- standard errors of mean

Q=quartile, LVMI =left ventricular mass index, EDV=end diastolic volume, RWT=relative valve thickness, LAVI=left atrial volume index, EF=ejection fraction, E =peak early diastolic transmitral inflow velocities, e' =average of mitral early diastolic septal and lateral annular velocities, GLS =global longitudinal strain, GCS = global circumferential strain, RV=right ventricular, RVFAC= RV fractional area change, TAPSE = tricuspid annular plane systolic excursion; TRV= tricuspid regurgitation velocity.

p value is calculated based on quartile 4 to other quartiles
p ≤ 0.01 considered significant

Supplemental Table 5. Unadjusted Associations Between Physical Activity and Echocardiographic Measures

	Sedentary time/day		MVPA/day		Light PA/day	
	Estimate	p-value	Estimate	p-value	Estimate	p-value
LVMI, g/m ^{2.7}	0.0721	0.003	-0.0002	0.999	-0.0892	0.012
RWT	0.0001	0.367	-0.0013	0.291	-0.0004	0.172
LVMI ^{2.7} /EDV	0.0011	0.004	-0.0007	0.005	-0.0033	<0.001
LAVI	-0.0042	0.481	0.1568	<0.001	0.0081	0.334
EF %	0.0003	0.973	-0.0200	0.810	-0.0057	0.748
E/e'	0.0065	0.268	-0.0696	0.244	-0.0396	<0.001
GLS, %	0.0006	0.345	-0.0279	0.494	-0.0280	0.023
GCS, %	-0.0350	0.011	-0.2244	0.015	-0.0348	0.242
SV, ml	0.0086	0.842	0.6841	0.093	-0.0165	0.812
Lat peak S	-0.0028	0.596	0.0271	0.257	0.0092	0.248
RV peak S	0.0072	0.136	0.0893	0.005	0.0211	0.016
RV FAC	0.0001	0.600	0.0017	0.125	0.0004	0.185
TRV, cm/sec	0.0004	0.959	0.3446	0.507	-0.1817	0.311
RV VTI	-0.0196	0.005	0.0088	0.861	0.0256	0.158
TAPSE, cm	-0.0012	0.137	0.0121	0.026	0.0045	<0.001

***p-value <0.01 considered significant**
Values are per 10 min/day of PA

LVMI =left ventricular mass index, RWT=relative valve thickness, EDV=end diastolic volume, LAVI=left atrial volume index, EF=ejection fraction, E =peak early diastolic transmitral inflow velocities, e' =average of mitral early diastolic septal and lateral annular velocities, LS =global longitudinal strain, GLS =global longitudinal strain, GLS = global circumferential strain, SV =stroke volume, RV=right ventricular, RVFAC= RV fractional area change, TR= tricuspid regurgitation, TAPSE = tricuspid annular plane systolic excursion, MVPA = moderate to vigorous-intensity PA, PA= physical activity

Supplemental Table 6: Multivariable Linear Regression for the Association between MVPA, Light PA, SB and Echocardiographic Measures in Healthy-Subgroup

Outcome	MVPA	Light PA	SB
LVMI, g/m ²	-0.053	0.053	-0.037*
LVMI ^{2.7} / EDV	-0.0015	0.0008	-0.0005
LAVI, ml/m ²	0.0371^	0.0065 ^	-0.0079^
EF %	0.0001	-0.0004	0.0003
E/e' ratio	0.0184^	0.0008*	-0.0022^
GLS, %	-0.0079^	-0.0023^	0.0024^
SV, ml	0.1743^	0.0038	-0.0188^
RV peak S, cm/sec	0.0056^	-0.0026	0.0015
RV FAC,%	0.0000	0.0000	0.0000
TRV Max, cm/sec	0.1234	0.0165	-0.0248*
TAPSE, cm	0.0018^	0.0006^	-0.0006^

Regression model adjusted for demographics (age and sex), systolic blood pressure, antihypertensive med use, other cardiovascular disease modifiable factors (diabetes mellitus, glycosylated hemoglobin, current smoking, and high-density lipoprotein).

*p-value < 0.01

^p value < 0.001

Values are per 10 min/day light MVPA/light PA/SB

MVPA = moderate to vigorous physical activity (PA); SB =sedentary behavior; LVMI =left ventricular mass index, EDV=end diastolic volume, LAVI=left atrial volume index, EF=ejection fraction, E =peak early diastolic transmitral inflow velocities, e' =average of mitral early diastolic septal and lateral annular velocities, GLS =global longitudinal strain, SV =stroke volume, RV=right ventricular, RVFAC= RV fractional area change, TRV= tricuspid regurgitation velocity, TAPSE = tricuspid annular plane systolic excursion