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RESEARCH ARTICLE

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Physical activity guidelines and cardiovascular risk in children: a cross sectional analysis to determine whether 60 minutes is enough

L. M. Füssenich¹, L. M. Boddy², D. J. Green^{2,3}, L. E. F. Graves², L. Foweather², R. M. Dagger⁴, N. McWhannell⁵, J. Henaghan², N. D. Ridgers⁶, G. Stratton⁷ and N. D. Hopkins^{2*}

Abstract

Background: Physical activity reduces cardiovascular mortality and morbidity. The World Health Organisation (WHO) recommends children engage in 60 min daily moderate-to-vigorous physical activity (MVPA). The effect of compliance with this recommendation on childhood cardiovascular risk has not been empirically tested. To evaluate whether achieving recommendations results in reduced composite-cardiovascular risk score (CCVR) in children, and to examine if vigorous PA (VPA) has independent risk-reduction effects.

Methods: PA was measured using accelerometry in 182 children (9–11 years). Subjects were grouped according to achievement of 60 min daily MVPA (active) or not (inactive). CCVR was calculated (sum of z-scores: DXA body fat %, blood pressure, VO_2 peak, flow mediated dilation, left ventricular diastolic function; CVR score $\geq 1SD$ indicated 'higher risk'). The cohort was further split into quintiles for VPA and odds ratios (OR) calculated for each quintile.

Results: Active children (92 (53 boys)) undertook more MVPA (38 ± 11 min, $P < 0.001$), had greater VO_2 peak (4.5 ± 0.8 ml/kg/min $P < 0.001$), and lower fat % (3.9 ± 1.1 %, $P < 0.001$) than inactive. No difference were observed between active and inactive for CCVR or OR ($P > 0.05$). CCVR in the lowest VPA quintile was significantly greater than the highest quintile (3.9 ± 0.6 , $P < 0.05$), and the OR was 4.7 times higher.

Conclusion: Achievement of current guidelines has positive effects on body composition and cardiorespiratory fitness, but not CCVR. Vigorous physical activity appears to have beneficial effects on CVD risk, independent of moderate PA, implying a more prescriptive approach may be needed for future VPA guidelines.

Keywords: Physical activity guidelines, Moderate/vigorous physical activity, Clustered cardiovascular risk

Background

Physical activity (PA) predicts cardiovascular disease (CVD) morbidity and mortality [1] and prevents and/or reduces traditional and emerging cardiovascular (CV) risk factors in healthy and high risk children [2, 3]. The American College of Sports Medicine originally published PA recommendations for young people in 1988, albeit based on their adult recommendations [4]. These guidelines and the supporting evidence base have been

re-evaluated numerous times in an attempt to account for advances in PA and CVD risk factor measurement techniques [5, 6]. Most recently, The World Health Organization (WHO) updated their paediatric PA recommendations [7] to reflect findings from a review by Janssen and LeBlanc [2] and the European Youth Heart Study [8], which suggested that previous guidelines underestimated the activity necessary to reduce CVD risk in young people. The WHO guidelines now suggest children aged 5–17 years accumulate 60 min of moderate-to-vigorous physical activity (MVPA) daily, in addition to everyday physical activities, and that vigorous

* Correspondence: N.d.hopkins@ljamu.ac.uk

²Research Institute for Sport and Exercise Science, Liverpool John Moore's University, Tom Reilly Building, Byrom Street, Liverpool L3 2ET, UK
Full list of author information is available at the end of the article

intensity physical activity (VPA) should be incorporated at least three times per week.

Whilst the updated guidelines address some of the limitations of previous versions [3] and advocate more PA than previously, numerous limitations remain. No study has empirically tested guidelines by comparing CVD risk in children who do achieve them to those who do not, whilst both Andersen et al. and Strong et al. [3, 9] provide comprehensive reviews on the dose response relationship, and make important recommendations for childhood PA, they do not test the validity of current recommendations using empirical data. Secondly, current guidelines are based on self-report PA, which has numerous limitations, and do not include novel CVD risk markers as outcome measures. Andersen et al. [3] addressed these limitations via inclusion of objective PA data, and novel inflammatory markers. However, endothelial and diastolic dysfunction are yet to be included in such analyses despite the crucial role each plays in the development of CVDs [10], their strong prognostic capacity in predicting CV events [11, 12], and clear associations with PA [13–15]. Finally, evidence suggests VPA may have potent effects on CVD risk [16], yet, WHO recommendations on VPA specifically, remain vague. With these limitations in mind, we sought to evaluate whether adherence to current WHO recommendations equates to a reduction in CVD risk in children. Secondly, we aimed to examine if, and in what quantity, VPA provides additional CVD risk benefits beyond moderate PA (MPA).

Methods

Subjects

Data were generated by the REACH Year 6 and the Active City of Liverpool, Active Schools and SportsLinX (A-CLASS) studies (Liverpool, UK). Fourteen schools from areas of social deprivation (IMD >40) participated across the studies; all 9–11 year old children were invited to participate. Following parental consent and medical screening, 233 children (10.8 ± 0.6 years; 100 boys) were studied. All were healthy and not suffering from cardiovascular/metabolic conditions. Informed written parent/guardian consent and child assent were obtained. Ethical approval was obtained from Liverpool John Moores University Ethics Committee. All data collection methods were standardised between the studies unless otherwise stated.

Experimental design

Initially, children visited the laboratory to complete measurements, including anthropometric tests and dual-energy X-ray absorptiometry (DXA), VO_2 peak testing, vascular endothelial function and echocardiography. Assessments were made in a quiet temperature-controlled room at the same time of day, following a morning fast

and avoidance of strenuous PA for 24 h. Subsequently, PA was assessed via accelerometry over 7 consecutive days.

Measurements and post-test analysis

Anthropometry and body composition

Body mass (kg), stature and sitting height (cm) were measured using standard methods. Somatic maturity was estimated by calculating time to peak height velocity (TPHV) using gender specific regression equations [17, 18].

A DXA scan (QDR discovery A, Hologic, MA) was completed according to standardized manufacturer procedures. Participants were scanned in the supine position while wearing t-shirt and shorts.

Vascular function

Following 20 min supine rest, brachial artery diameter, blood flow and shear rate were assessed via high-resolution ultrasonography (Acuson, Aspen, Penn and Terason, T3000, Aloka, UK) prior to, and following, 5 min forearm ischaemia. Methods were identical to those previously described [14] in accordance with best practice guidelines [19].

Left ventricular diastolic function

Following 10 min of quiet rest in the left lateral decubitus position. Left ventricular diastolic function was assessed via echocardiography (Mylab30CV system, ESAOTE, Italy). All system settings including gain, filter, PRF, sector size and depth were adjusted to optimise image quality. Mitral inflow was assessed from the apical four-chamber via a 2 mm sample volume at the tips of the mitral leaflets, parallel with flow, peak early (E) and late/atrial (A) velocities were obtained and E/A ratio reported.

Cardiorespiratory fitness

During each protocol VO_2 and VCO_2 were measured breath-breath via an online gas analysis system (Jaeger Oxycon Pro, Viasys Health Care, Warwick, UK). Heart rate (HR) was monitored continuously (Polar, Kempele, Finland).

A-CLASS study method Peak oxygen uptake (VO_2 peak) was determined during a discontinuous treadmill exercise test which involved walking and running until volitional exhaustion. The test consisted of 3 min stages, followed by a 30-s rest interval. Peak VO_2 was accepted as the highest 15-s averaged oxygen uptake achieved during the test with a respiratory exchange ratio ≥ 1.05 and/or HR ≥ 195 beats.min⁻¹.

REACH study method To account for differences in biological age and limb length, treadmill speeds were individually calibrated by anchoring speeds to set Froude (Fr) numbers as described previously [20]. A continuous

protocol was used until volitional exhaustion occurred. Peak VO_2 was defined as the highest 15-s average oxygen uptake achieved with a respiratory exchange ratio >1.05 and/or HR >199 beats/min⁻¹.

Physical activity

Physical activity was objectively measured for 7 consecutive days using a hip mounted uni-axial accelerometer (GT1M model, ActiGraph, FL, USA) set to 5 s epochs. Children wore the accelerometer during all waking hours, except during water-based activities. Consecutive zero counts >20 min were removed from analysis as non-wear. Minimum wear time for inclusion in data analysis was 9 h/day for any 3 days of the week [21]. Accelerometer data reduction was performed using ActiLife v 6.1.4 (ActiGraph, LLC, 2010–2012). The Evenson cut-points [22] were used to define PA and sedentary intensity thresholds [23]. Total time spent in each PA/SB threshold for each valid day was divided by the total number of valid days. Subjects were then split into groups; those who achieved a daily average of 60 min MVPA (active) and those who did not (inactive [24]).

Statistical analysis

All statistical analysis were performed using SPSS (18.0, Chicago, Illinois) software. Statistical significance was set at $P < 0.05$. Variables were tested for normality when grouped by sex and PA level using the Kolmogorov-Smirnov test. DBP and E/A ratio data were normalised by log transformation and FMD using square root transformation. All analyses were performed at the cohort level initially; additional analyses were then performed by gender.

Gender specific standardized z-scores for percentage body fat, VO_2 peak, FMD, systolic blood pressure and E/A ratio were calculated and inverted where necessary, z-scores were summed to create a composite CVD risk score (CCVR). Pearson's correlation analysis was used to assess relationships between z scores (Table 1). Children with CCVR ≥ 1 SD were defined as 'higher risk' [8]. Differences in individual CVD risk factors and CCVR between the active and inactive group, boys and girls were assessed

using an independent *t*-test, or analysis of covariance (ANCOVA), with TPHV as a covariate (continuous variables), or a chi-square test (discrete variables).

The cohort was split into quintiles according to VPA. Differences in individual CVD risk factors and CCVR across quintiles were assessed using ANCOVA with MPA, sedentary behaviour and TPHV as covariates, or using a chi-square test (discrete variables). Logistic regression was then used to obtain odds ratios for each group (Boys/Girls, Active/Inactive and all VPA quintiles).

Results

Subject characteristics

Of the 233 children initially recruited, 182 children met criteria for accelerometer wear time coupled with adequate vascular measurements. Those who did not meet criteria were removed from analyses, there were no differences between included and excluded children across measured variables. Baseline characteristics are presented in Table 2. Boys had significantly higher VO_2 peak and DBP, whilst girls were significantly closer to PHV and had higher percentage body fat. Additionally, boys engaged in significantly more MVPA and VPA than girls (Table 2).

Active vs. inactive analysis

Inactive children had significantly higher percentage body fat and lower VO_2 peak compared to active children. No other significant differences were found between the two groups (Table 3). The odds for being 'higher risk' were 1.9 (95 % CI: 0.8–4.3) times higher in the inactive group than the active group ($p = 0.126$).

Gender analysis

Boys were more active than girls ($p < 0.001$, Table 3) and active boys engaged in more MVPA than active girls (11 min/day, $p = 0.004$). Active boys had significantly lower percentage body fat and CCVR, and higher VO_2 peak than inactive boys (Table 3). Thirty seven percent of inactive boys were classed as 'at risk' compared to 10 % of the active boys ($p = 0.028$), the odds of being at risk were 5.1 times higher in the inactive boys than the active boys (95 % CI: 1.4 – 19.1, $p = 0.015$). Active

Table 1 Correlations between z scores used to calculate CCVD risk score

Z score	VO_2 max (ml/kg/min)		Fat % (DXA)		SBP (mmHg)		FMD %		E/A ratio	
	<i>r</i>	<i>p</i>	<i>r</i>	<i>p</i>	<i>r</i>	<i>p</i>	<i>r</i>	<i>p</i>	<i>r</i>	<i>p</i>
VO_2 max (ml/kg/min)	1		-0.691	0.000	-0.163	0.036	0.025	0.743	0.173	0.027
Fat % (DXA)	-0.691	0.000	1		0.239	0.001	-0.056	0.448	-0.230	0.002
SBP (mmHg)	-0.163	0.036	0.239	0.001	1		-0.112	0.134	-0.068	0.370
FMD %	0.025	0.743	-0.056	0.448	-0.112	0.134	1		0.072	0.340
E/A ratio	0.173	0.027	-0.230	0.002	-0.068	0.370	0.072	0.340	1	

Fat % percentage of body fat, DXA dual-energy X-ray absorptiometry, SBP systolic blood pressure, VO_2 max peak oxygen uptake, FMD flow mediated dilation, E/A ratio ratio between passive and active filling of the left ventricle (cm/s)

Table 2 Descriptive statistics

	Group	Girls (105)	Boys (77)
Age (years)	10.8 (0.6)	10.7 (0.6)	10.8 (0.6)
Height (cm)	145.0 (8.0)	145.4 (7.8)	144.6 (8.3)
Body mass (kg)	41.7 (10.6)	42.3 (10.2)	40.9 (11.0)
Maturity offset (TPHV)	-2.51 (1.23)	-2.30 (1.44)	-2.79 (.79)
BMI (kg/m ²)	19.6 (3.5)	19.8 (3.3)	19.3 (3.7)
Fat % (DXA)	27.7 (6.7)	29.4 (5.7)	25.2 (7.2) ^a
SBP (mmHg)	106 (11)	105 (11)	108 (12)
DBP (mmHg)	63 (6)	62 (5)	64 (6) ^a
VO ₂ max (ml/kg/min)	46.0 (6.9)	43.5 (6.1)	49.6 (6.4) ^a
FMD %	8.9 (4.1)	8.9 (4.1)	8.9 (4.3)
E/A ratio	2.1 (0.5)	2.1 (0.4)	2.1 (0.5)
CCVD risk	-0.04 (2.94)	0.12 (2.98)	-0.27 (2.89)
MVPA (min)	64 (25)	56 (20)	75 (27) ^a
VPA (min)	25 (13)	22 (12)	30 (13) ^a
Sedentary (min)	605 (148)	608 (80)	600 (209)
MPA/VPA ratio	1.7 (0.6)	1.8 (0.7)	1.7 (0.6)

TPHV time to peak height velocity, BMI body mass index, Fat % percentage of body fat, DXA dual-energy X-ray absorptiometry, SBP systolic blood pressure, DBP diastolic blood pressure, VO₂max peak oxygen uptake, FMD flow mediated dilation, E/A ratio ratio between passive and active filling of the left ventricle (cm/s), CCVD risk composite cardio vascular disease risk score as sum of z-scores, MVPA moderate to vigorous physical activity, VPA vigorous physical activity. All data expressed as mean (SD). ^aSignificant difference between boys and girls <0.05

girls engaged in significantly less sedentary behaviour compared to inactive girls ($p < 0.001$), no further differences were found.

Vigorous physical activity

Comparisons were made between the highest VPA quintile (Q5) and all others (Table 4). CCVR was significantly elevated in Q1. Q1, Q2 and Q4 had significantly higher percentage body fat. VO₂peak was significantly lower in Q1 and Q2. The OR was significantly higher in Q1 than Q5 (OR 4.7, $p < 0.05$, Table 4; Fig. 1). When quintiles were examined by gender, no significant differences were found between quintiles (difference between Q1 and Q5: 6.1 (95 % CI: 0.6 – 59.5, $p < 0.05$) and 7.4 (95 % CI: 0.7 – 80.0, $p < 0.05$), in girls and boys respectively).

Discussion

To our knowledge, this is the first study which used a composite risk score which included novel pre-clinical markers of CVD risk to empirically evaluate the effects of adherence to the current WHO PA guidelines for children, and provide novel information regarding recommendations for VPA. Our findings from the whole cohort indicate no difference in CCVR between active and inactive asymptomatic 9–11 year old children, implying current guidelines may underestimate the PA necessary to reduce CCVR. Although achieving WHO PA recommendations did have beneficial effects on VO₂peak and adiposity. Furthermore, VPA appeared to provide health benefits in addition to those of MPA,

Table 3 Cardiovascular risk factors in the active and inactive

	Group			Girls			Boys		
	Inactive (n = 90)	Active (n = 92)	p	Inactive (n = 66)	Active (n = 39)	p	Inactive (n = 24)	Active (n = 53)	p
Fat %	29.6 (5.9)	25.7(7.0)	>0.001	29.2 (5.9)	29.8 (5.4)	0.90	30.7 (5.8)	22.7 (6.5)	>0.001
SBP	106 (12)	106 (12)	0.89	106 (10)	104 (11)	0.58	108 (11)	108 (12)	0.97
Baseline artery diameter (mm)	3.1 (0.3)	3.1 (0.5)	0.73	3.0 (0.3)	3.0 (0.4)	0.67	3.2 (0.4)	3.1 (0.5)	0.90
FMD%	9.1 (4.1)	8.7 (4.2)	0.45	9.1 (4.1)	8.6 (3.9)	0.65	9.2 (4.0)	8.8 (4.4)	0.44
VO ₂ max	43.70 (6.1)	48.3 (6.9)	>0.001	43.1 (6.3)	44.2 (5.7)	0.16	45.56 (5.2)	51.2 (6.2)	0.01
E/A ratio	2.2 (0.5)	2.1 (0.4)	0.78	2.2 (0.5)	2.2 (0.4)	0.36	2.1 (0.6)	2.2 (0.4)	0.64
CCVD risk	0.36 (2.95)	-0.43 (2.90)	0.40	0.09 (2.93)	0.19 (3.12)	0.87	1.22 (2.92)	-0.85 (2.70)	0.01
At risk (%)	23	13	0.35	18	18	0.99	37	10	0.03
OR CCVD risk	1.9	1.0	0.13	1.0	1.0	0.99	5.1	1.0	0.02
MVPA (min)	45 (10)	83 (21)	>0.001	45 (10)	77 (17)	>0.001	46 (9)	88 (22)	>0.001
VPA (min)	16 (6)	33 (14)	>0.001	16 (6)	30 (15)	>0.001	17 (5)	36 (12)	>0.001
MPA/VPA ratio	1.9 (0.7)	1.6 (0.6)	>0.001	1.9 (0.7)	1.7 (0.7)	0.18	1.9 (0.6)	1.6 (0.5)	>0.01
Sedentary (min)	625 (80)	585 (192)	0.03	629 (84)	573 (59)	>0.001	613(68)	594 (248)	0.53

Inactive = those who did not achieve the recommended 60 min MVPA per day. Active = those who achieved the recommended 60 min MVPA per day
Fat % percentage of body fat, SBP systolic blood pressure, BD baseline diameter of brachial artery in mm, FMD flow mediated dilation, VO₂max peak oxygen uptake, E/A ratio ratio between passive and active filling of the left ventricle (cm/s), CCVD risk clustered cardio vascular disease risk score as sum of z-scores, At risk percentage of children with more than 1SD in the CCVD risk score, OR CCVD risk odds ratio for being at risk compared to the active group, MVPA moderate to vigorous physical activity in minutes/day, VPA vigorous physical activity in minutes/day, sedentary sedentary behaviour in hours/day. All data expressed as mean (SD)

Table 4 Cardiovascular risk factors in quintiles of vigorous physical activity

Whole cohort	Number	VPA	MPA	Sedentary	MPA/VPA ratio	CCVD risk	At risk (%)	OR	Fat %	FMD%	SBP	VO2max	E/A ratio
1	36	11 (2) ^a	27 (10) ^a	632 (91)	2.5 (0.8) ^a	1.31 (3.26) ^a	34.5 % ^a	4.7 ^a	31.9 (1.2) ^a	8.4 (3.8)	106 (12)	42.3 (6.7) ^a	2.0 (0.3)
2	37	17 (2) ^a	34 (8) ^a	613 (64)	2.0 (0.5) ^a	0.23 (2.93)	20.7 %	2.1	29.3 (1.0) ^a	9.5 (4.9)	104 (12)	43.6 (5.3) ^a	2.2 (0.4)
3	36	22 (2) ^a	36 (7) ^a	603 (69)	1.6 (0.3)	-0.40 (2.81)	17.2 %	1.6	27.1 (1.0)	8.3 (3.7)	105 (9)	46.7 (6.7)	2.3 (0.6)
4	37	29 (2) ^a	43 (11) ^a	587 (56)	1.5 (0.4)	-0.06 (2.58)	17.2 %	1.7	27.5 (1.0) ^a	9.1 (4.2)	108 (12)	46.9 (6.9)	2.0 (0.3)
5	36	45 (14)	56 (13)	587 (303)	1.3 (0.3)	-1.22 (2.71)	10.3 %	1	22.4 (1.3)	9.3 (4.0)	106 (11)	51.0 (5.8)	2.1 (0.5)
Boys													
1	15	13 (2) ^a	30 (11) ^a	613 (72)	2.3 (0.8) ^a	1.11 (3.38)	36.4 %	7.4	30.0 (7.0)	8.8 (4.5)	105 (13)	47.5 (6.1)	1.9 (0.3)
2	16	20 (2) ^a	39 (6) ^a	590 (73)	1.9 (0.3) ^a	0.25 (2.63)	18.8 %	3.0	27.4 (6.5)	9.0 (4.0)	106 (11)	47.1 (7.2)	2.3 (0.6)
3	15	28 (3) ^a	42 (11) ^a	586 (22)	1.5 (0.3)	-0.20 (2.68)	15.4 %	2.4	24.7 (7.2)	7.3 (3.7)	111 (13)	50.8 (5.7)	2.1 (0.4)
4	16	36 (2) ^a	51 (9) ^a	675 (42)	1.4 (0.2)	-0.72 (3.04)	14.3 %	2.2	23.7 (6.9)	10.0 (5.3)	108 (10)	49.7 (5.5)	2.2 (0.5)
5	15	50 (9)	63 (14)	530 (48)	1.3 (0.2)	-1.56 (2.56)	7.1 %	1	20.2 (5.4)	9.6 (3.7)	110 (11)	53.0 (6.1)	2.2 (0.4)
Girls													
1	21	10 (2) ^a	25 (8) ^a	652 (99) ^a	2.5 (0.8) ^a	0.51 (3.16)	27.8 %	6.2	29.6 (6.8)	8.9 (4.0)	107 (11)	42.3 (5.9)	2.1 (0.3)
2	21	15 (1) ^a	33 (9) ^a	625 (75)	2.2 (0.6) ^a	0.98 (3.73)	30.0 %	6.9	32.2 (6.3) ^a	8.3 (3.9)	103 (14)	40.7 (5.9)	2.2 (0.5)
3	21	20 (1) ^a	31 (6) ^a	601 (57)	1.6 (0.3)	-0.69 (2.25)	5.6 %	0.9	28.9 (5.3)	10.1 (4.6)	104 (8)	44.0 (5.4)	2.3 (0.5)
4	21	26 (2) ^a	39 (8)	605 (69)	1.5 (0.3)	0.43 (2.71)	20.0 %	4.0	29.3 (4.6)	7.7 (3.8)	106 (9)	45.6 (5.5)	2.0 (0.5)
5	21	37 (18)	45 (13)	559 (69)	1.3 (0.7)	-0.79 (2.62)	5.9 %	1	27.2 (4.8)	9.6 (3.9)	104 (11)	45.3 (6.9)	2.1 (0.3)

VPA vigorous physical activity (min/day), MPA moderate physical activity (min/day), CCVD risk clustered cardiovascular disease risk score as sum of z-scores, At risk % of children with more than 1SD in the CCVD risk score, OR CCVD risk odds ratio for being at risk compared to the active group, Fat % percentage of body fat, FMD% flow mediated dilation, SBP systolic blood pressure, VO2max peak oxygen uptake, E/A ratio ratio between passive and active filling of the left ventricle (cm/s). All data expressed as mean (SD). CCRD risk score consists of Fat%, FMD%, SBP, VO2Max, E/A ratio. ^aSignificant difference compared to most active quintile <0.05

suggesting it necessary to incorporate a more specific recommendation regarding the VPA.

Active boys have 1/5 of the risk compared to inactive boys (Table 3), whereas no differences were found between active and inactive girls. We observed a significant

difference in MVPA between active boys and active girls, (11 ± 5 min, *p* = 0.004; Table 3) and active girls had a similar risk factor profile to inactive girls and inactive boys. This suggests that gender differences in risk factors may not be resultant of gender *per se*, but are possibly

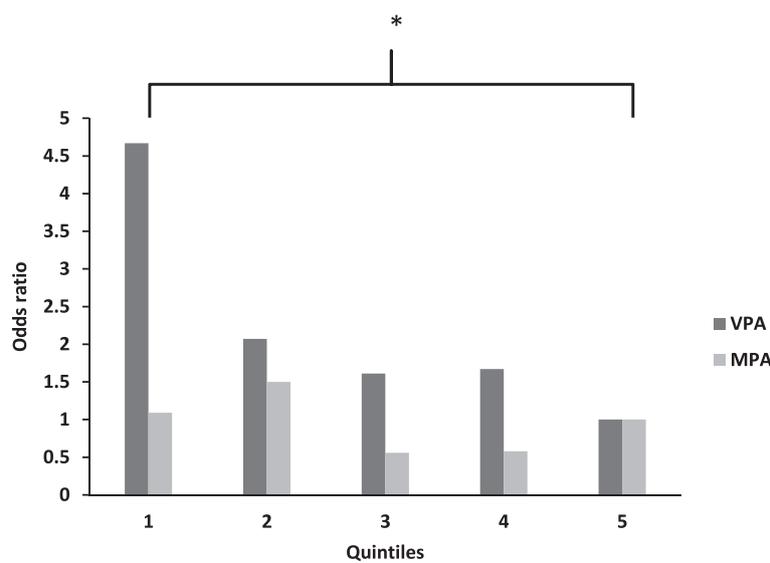


Fig. 1 Odds ratios for being at risk by quintiles of physical activity. *Significant difference between VPA quintile 1 and 5, *P* < 0.05. No significant differences across MPA quintiles. VPA = vigorous physical activity, MPA = moderate physical activity

determined by differences in PA levels [2, 3, 25, 26]. Although active girls engaged in daily MVPA in excess 60 min, this appears insufficient to confer the CVD risk benefits afforded by the active boys. Our data suggest 77 min MVPA was insufficient for CCVR risk reduction, whilst 88 min MVPA resulted in a reduction, supporting previous conclusions from Andersen et al. [8] that PA guidelines should be higher than the current 60 min MVPA/day. Furthermore, our data raise the question of whether children's PA guidelines should be gender specific; little is known about gender specific dose responses to PA, further investigation therefore appears to be warranted.

In agreement with adult literature, which demonstrate that VPA is a more meaningful predictor of cardiometabolic risk than MPA [27], we provide evidence that VPA may afford additional CVD risk reduction in children. CCVR ORs were ~5 times higher in Q1 compared to Q5 ($p < 0.05$), whilst no differences were observed across MPA quintiles (Fig. 1). The largest reduction in risk was evident between Q1 and Q2 (VPA 11 ± 2 vs 17 ± 2 min, respectively, $p > 0.05$; Table 4) and although this risk reduction is non-significant in this small study, we suggest that a reduction in odds ratio of ~50 % is a clinically meaningful finding. On this basis, 17 min of VPA per day (Q2), equating to around 2 h VPA/week is needed to reduce CVD risk in pre-pubertal children.

In contrast to percent body fat and VO_2 peak, there was no significant difference between active and inactive groups or across VPA quintiles for E/A ratio or FMD. Whilst the inclusion of FMD and E/A ratio in analysis provides novel information, these surrogate markers do not appear to enhance the predictive power of the CCVR model in this cohort. One possible interpretation of this finding is that diastolic and endothelial function are not modulated by PA, however a wealth of previous findings contradict this hypothesis [13, 14, 28]. A more plausible hypothesis relates to the dose of activity the inactive children are exposed to; 45 min MVPA/day, including 17 min of VPA, whilst they fall short of WHO guidelines for MVPA, this level of VPA may be adequate to prevent deterioration of endothelial and diastolic function. These data lend support to our recommendations above that 17 min VPA/day is cardioprotective, and previous findings from our group which imply that VPA is more important for endothelial function than other PA intensities [14]. Further research is needed to confirm or refute the role of VPA in the modulation of these variables.

This study advances knowledge of the complex relationship between PA and CVD risk in children, as it investigates, for the first time, the utility of adding novel CVD surrogates, to a composite score of 'pre-clinical' markers to estimate CVD risk. As CVD risk factors tend to cluster in sedentary and obese individuals, stronger associations

between CVD risk and PA may be observed when a composite CVD risk score is generated. Additionally, as we measured PA levels objectively we are confident of greater measurement precision than the self-report PA data from which the current WHO guidelines are derived, nonetheless given the sporadic nature of children's PA patterns, it is plausible that using an epoch length of 5 s may result in an underestimation of VPA. Furthermore, a lack of parity in PA measurement techniques and accelerometer cut points used between this and other studies makes comparison and interpretation difficult. Various confounders including diet, smoking status and socioeconomic status were not accounted for in our analyses as we lack the data to do so. Finally, although our findings lend support to the previous recommendations that 60 min daily MVPA may not be enough for children of this age [8], given the relatively small sample size and limited number of children that achieved guidelines, findings should be interpreted with caution.

Conclusions

In our study of asymptomatic 9–11 year old children, there were no differences between CCVR of children who undertook 60 min MVPA per day in accordance with WHO recommendations, and those who did not. This implies that current recommendations may be an underestimation of the PA necessary to reduce clustered CVD risk. A gender difference between the CVD risk in active and inactive children, raises the possibility that gender specific guidelines may be needed, although much work is needed to determine if these differences are a result of gender specific responses to PA or sex differences in PA level. Finally, VPA appears to provide CCVR benefits beyond those afforded by MPA, with data suggesting that 17 min VPA/day may provide clinically meaningful CVD risk reductions. Taken together these findings suggest that in order to reduce CVD risk, the current guidelines should be updated to increase the amount of MVPA recommended, and to prescribe a daily amount of VPA.

Abbreviations

PA: physical activity; VPA: vigorous physical activity; MPA: moderate physical activity; MVPA: moderate to vigorous physical activity; CVD: cardiovascular disease; WHO: World Health Organisation; CCVR: clustered cardiovascular risk; OR: odds ratio; DXA: dual-energy X-ray absorptiometry; FMD: flow mediated dilation; TPHV: time to peak height velocity.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

LF – Data analysis and interpretation, significant writer of article, review of final draft. LMB- conception and design, revising manuscript, review of final draft. DJG - conception and design, revising manuscript, review of final draft. LEFG - Data collection and analysis, revising manuscript, review of final draft. LF - Data collection and analysis, revising manuscript, review of final draft. RG - Data collection and analysis, revising manuscript, review of final draft.

NM - Data collection and analysis, revising manuscript, review of final draft.
 JH - Data collection and analysis, revising manuscript, review of final draft.
 NDR - Data collection and analysis, revising manuscript, review of final draft.
 GS - conception and design, revising manuscript, review of final draft.
 NDH - conception and design, significant writer, review of final draft. All authors read and approved the final manuscript.

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Author details

¹Department of Physiology, Radboud University Nijmegen Medical Centre, Nijmegen, Netherlands. ²Research Institute for Sport and Exercise Science, Liverpool John Moore's University, Tom Reilly Building, Byrom Street, Liverpool L3 2ET, UK. ³School of Sports Science, Exercise and Health, The University of Western Australia, Crawley, Perth, WA 6009, Australia. ⁴School of Health Sciences, Liverpool Hope University, Liverpool L16 9JD, UK. ⁵Department of Sport and Exercise Science, University of Chester, Chester CH1 4BJ, UK. ⁶Centre for Physical Activity and Nutrition (C-PAN) Research, Deakin University, 221 Burwood Highway, Burwood, VIC 3175, Australia. ⁷Applied Sports Technology Exercise and Medicine Research Centre (A-STEM), Swansea University, Swansea SA2 8PP, UK.

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