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Associations of physical activity, sedentary time, and diet quality with biomarkers of inflammation in children

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ABSTRACT

We investigated the associations of physical activity (PA), sedentary time (ST), and diet quality with biomarkers of inflammation in 390 children (192 girls, 198 boys) aged 6–8 years. PA energy expenditure (PAEE), light PA, moderate PA (MPA), vigorous PA (VPA), moderate-to-vigorous PA (MVPA), and ST were assessed by combined movement and heart rate sensor. Finnish Children Healthy Eating Index was calculated using data from 4 d food records. Body fat percentage (BF%) was measured by dual-energy X-ray absorptiometry. High-sensitivity C-reactive protein (Hs-CRP), leptin, interleukin-6 (IL-6), adiponectin, tumour necrosis factor- α , and glycoprotein acetyls were measured from fasting blood samples. PAEE, MPA, VPA, and MVPA were inversely associated with Hs-CRP ($\beta=-191$ to -139 , 95% CI= -0.294 to -0.024), leptin ($\beta=-0.409$ to -0.301 , 95% CI= -0.499 to -0.107), IL-6 ($\beta=-0.136$ to -0.104 , 95% CI= -0.240 to -0.001) and PAEE, MPA, and MVPA were inversely associated with glycoprotein acetyls ($\beta=-0.117$ to -0.103 , 95% CI= -0.213 to -0.001). ST was directly associated with Hs-CRP ($\beta=0.170$, 95% CI= $0.070-0.269$), leptin ($\beta=0.355$, 95% CI= $0.265-0.445$), and IL-6 ($\beta=0.105$, 95% CI= $0.005-0.205$). VPA was inversely associated with Hs-CRP, leptin, and IL-6 in children with higher BF% ($\beta=-0.344$ to -0.181 , 95% CI= -0.477 to -0.033) but not among children with lower BF% ($\beta=-0.007-0.033$, 95% CI= $-0.183-0.184$). In conclusion, PA was inversely and ST directly associated with circulating levels of biomarkers of inflammation among children. Furthermore, we observed that PA was inversely associated with these biomarkers for inflammation in children with a higher BF%.

Abbreviations: BF%: Body fat percentage; BMI: Body mass index; BMI-SDS: Body mass index standard deviation score; FCHEI: Finnish Children Healthy Eating Index; Hs-CRP: High-sensitivity C-reactive protein; IL-6: Interleukin-6; MET: Metabolic Equivalent of Task; PA: Physical activity; PANIC: Physical Activity and Nutrition in Children Study; ST: Sedentary time; TNF- α : Tumour necrosis factor α .

Highlights

- Systemic inflammation, as indicated by increased circulating concentrations of biomarkers for inflammation, may be important in causal pathways leading to insulin resistance, sub-clinical atherosclerosis, and eventually clinical manifestations of cardiovascular diseases.
- Higher levels of physical activity and lower levels of sedentary time were associated with more favourable inflammatory profile.
- Body fat percentage modified these associations and especially vigorous intensity physical activity was inversely associated with biomarkers of inflammation on children with higher body fat percentage but not in children with lower body fat percentage.

KEYWORDS

Inflammation; biomarkers; youth; exercise; nutrition; obesity

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Introduction

The prevalence of paediatric obesity is increasing in most parts of the world, predisposing children to increased risk for type 2 diabetes and cardiovascular diseases already in childhood (Juonala et al., 2011). Furthermore, few children meet the recommended levels of physical activity (PA) (Cooper et al., 2015; Guinhouya, Samouda, & de Beaufort, 2013) and adhere to the dietary recommendations (Banfield et al., 2016; Eloranta et al., 2011). Physical inactivity (Wahid et al., 2016) and poor diet quality (Rodríguez-Monforte, Flores-Mateo, & Sánchez, 2015) have been found to increase the risk of type 2 diabetes and cardiovascular diseases in adults. Importantly, pathological process leading to these cardiometabolic diseases begin already in childhood (Steinberger et al., 2016). Traditional cardiometabolic risk factors, such as overweight, insulin resistance, and dyslipidaemia, have been associated with increased risk of type 2 diabetes and sub-clinical atherosclerosis and cardiovascular diseases in adulthood (Baigent et al., 2020; Magnussen et al., 2010). However, recent evidence suggests that systemic inflammation, as indicated by increased circulating concentrations of biomarkers for inflammation, may be important in causal pathways leading to insulin resistance, sub-clinical atherosclerosis, and eventually clinical manifestations of cardiovascular diseases (Gleeson et al., 2011).

Obese children have been found to have higher circulating levels of biomarkers of low-grade inflammation than their normal-weight peers (Lund et al., 2020). Furthermore, higher levels of PA and better diet quality, as indicated by a higher consumption of fruits, vegetables, berries, vegetable oils, and fish and low consumption of foods containing high amount of sugar, have been associated with lower levels of traditional cardiometabolic risk factors already in children (Eloranta et al., 2016; Poitras et al., 2016). Even if some of the associations of PA and diet quality with biomarkers for inflammation may be modified by fat mass, it has been suggested that increased PA and improved diet quality are independently associated with decreased systemic inflammation. There are few studies on the associations of PA and diet quality with biomarkers for inflammation in general populations of children and the observed associations have been weak (Poitras et al., 2016). Furthermore, screen time, commonly defined as time spent in TV viewing, and sedentary time (ST) assessed by accelerometers, has been directly associated with circulating levels of high-sensitivity C-reactive protein (hs-CRP) (Carson & Janssen, 2011; Gabel et al., 2016) but not with interleukin 6 (IL-6), adiponectin, or tumour necrosis factor- α (TNF- α) in children and adolescents (Gabel et al., 2016). The results of intervention studies in obese children suggest that exercise training

has a potential to reduce circulating levels of hs-CRP (Han et al., 2019), leptin, and IL-6 (Sirico et al., 2018) and increase circulating levels of adiponectin (García-Hermoso et al., 2017; Sirico et al., 2018). Evidence on the effects of exercise training on individual biomarkers of inflammation is still mixed and most of the studies addressing this issue have been performed among overweight or obese children (García-Hermoso et al., 2017; Han et al., 2019; Sirico et al., 2018). These contradictory observations may be due to differences in exercise intensity or in the reduction of fat mass in response to exercise training (García-Hermoso et al., 2017).

Healthy dietary patterns have been associated with decreased circulating levels of hs-CRP in adults (Norde, Collese, Giovannucci, & Rogero, 2021), but evidence on the role of diet quality indices in systemic low-grade inflammation in children is limited and mixed. Liese et al. found no associations of diet quality indices with biomarkers for inflammation, such as hs-CRP and IL-6, in youth with type 1 diabetes mellitus (Liese et al., 2018). Nevertheless, Saneei et al. observed that a dietary intervention reduced hs-CRP, but had no effect on IL-6, adiponectin, or TNF- α in adolescents with metabolic syndrome (Saneei, Hashemipour, Kelishadi, & Esmailzadeh, 2014). Moreover, some studies have reported a weak if any effect of a combined exercise and diet intervention on hs-CRP and TNF- α (Blüher et al., 2014). Nevertheless, two-week intensive combined diet and exercise intervention decreased circulating levels of leptin, IL-6, and TNF- α and to increase adiponectin in normal weight and obese children and youth aged 8–17 years independent of changes in waist circumference, body weight, or body mass index (BMI) (Roberts, Izadpanah, Angadi, & Barnard, 2013). Interestingly, dietary intervention or a combined dietary and exercise intervention increased serum adiponectin but a moderate intensity exercise intervention alone decreased it over 12 weeks among obese children (Shalitin et al., 2009). In addition, a combined exercise, diet, and behavioural intervention has been found to increase adiponectin and decrease hs-CRP but not IL-6 and leptin in obese children (Nemet, Oren, Pantanowitz, & Eliakim, 2013). There is also some evidence that diet quality modifies the association between screen time and biomarkers for inflammation in adolescents (Arouca et al., 2019). However, little is known about the possible modifying effects of diet quality or body fat percentage (BF%) on the associations of PA with biomarkers for inflammation in general populations of children. Furthermore, there are no previous studies on the associations of PA and diet quality with glycoprotein acetyls in children. Increased level of glycoprotein acetyls, assessed by nuclear magnetic resonance (NMR) spectroscopy, is a composite biomarker

of systemic inflammation that have been independently associated with increased risk of type 2 diabetes, cardiovascular diseases, cardiovascular mortality (Connelly et al., 2017; Kettunen et al., 2018).

We investigated the associations of ST, screen time, PA at different intensities, and diet quality with biomarkers for inflammation, including hs-CRP, leptin, IL-6, adiponectin, TNF- α , and glycoprotein acetyls, in a population sample of children. We also investigated the modifying effect of BF% on these associations. Finally, we studied the modifying effect of diet quality on the associations of ST, screen time, and PA at different intensities with these biomarkers of inflammation.

Methods

Study design and study participants

The present data are from the Physical Activity and Nutrition in Children (PANIC) Study, which is an 8-year physical activity and dietary intervention study and a long-term follow-up study in a population sample of children from the city of Kuopio, Finland (Eloranta et al., 2011). The Research Ethics Committee of the Hospital District of Northern Savo approved the study protocol in 2006 (Statement 69/2006). The parents or caregivers of the children gave their written informed consent, and the children provided their assent to participation. The PANIC study has been carried out in accordance with the principles of the Declaration of Helsinki as revised in 2008.

Altogether 736 children 6–8 years of age from primary schools of Kuopio were invited to participate in the baseline examination in 2007–2009. A total of 512 children, who represented 70% of those invited, participated in the baseline examinations. Six children were excluded from the study at baseline because of physical disabilities that could hamper participation in the intervention or no time or motivation to attend the study. The participants did not differ in sex distribution, age, or BMI standard deviation score (BMI-SDS) from all children who started the first grade in 2007–2009 based on data from the standard school health examinations performed for all Finnish children before the first grade (data not shown). Complete data on variables used in the analyses on the associations of ST, media time, PA, and diet quality with inflammatory biomarkers were available for 390 children (192 girls, 198 boys).

Assessment of body size, body composition, and pubertal status

Body weight was measured twice with the children having fasted for 12 h, emptied the bladder, and standing

in light underwear using a weight scale integrated into a calibrated InBody® 720 bioelectrical impedance device (Biospace, Seoul, South Korea) to an accuracy of 0.1 kg. The mean of these two values was used in the analyses. Stature was measured three times with the children standing in the Frankfurt plane without shoes using a wall-mounted stadiometer to an accuracy of 0.1 cm. The mean of the nearest two values was used in the analyses. BMI was calculated by dividing weight (kg) by stature (m) squared. BMI-SDS was calculated based on Finnish reference data (Saari et al., 2011). The prevalence of overweight and obesity was defined using the cut-off values provided by Cole et al. (2000). Body fat percentage (BF%) was measured by the Lunar® dual-energy X-ray absorptiometry device (GE Medical Systems, Madison, WI, USA) using a standardised protocol.

A research physician assessed pubertal status using a 5-stage scale described by Marshall and Tanner (1969, 1970). Boys were defined as having entered clinical puberty if their testicular volume assessed by an orchidometer was ≥ 4 mL (stage ≥ 2). Girls were defined having entered clinical puberty if their breast development had started (stage ≥ 2).

Assessment of biomarkers for inflammation

Venous blood samples were taken the children having fasted for 12 h. Blood was immediately centrifuged and stored at a temperature of -75°C until biochemical analyses. Plasma hs-CRP was measured using an enhanced immunoturbidimetric assay with the CRP (Latex) High Sensitive Assay reagent (Roche Diagnostics GmbH, Mannheim, Germany) and the limit of quantitation of 0.3 mg/l. Plasma leptin concentration was measured by a competitive radioimmunoassay (Multi-gamma 1261-001, PerkinElmer Wallac Oy, Turku, Finland). Commercially available ELISA kits were employed for the measurement of plasma IL-6 and TNF- α concentrations (Sanquin Reagents, Amsterdam, The Netherlands). Serum high-molecular-weight adiponectin concentration was analysed using an ELISA kit after a specific proteolytic digestion of other multimeric adiponectin forms (Millipore, Billerica, MA, USA). The Nightingale high-throughput NMR metabolomics platform was used to quantify plasma glycoprotein acetyls (Soininen et al., 2015).

Assessment of sedentary time and physical activity

ST and PA were assessed using a combined heart rate and movement sensor (Actiheart®, CamNtech Ltd., Papworth, UK) for a minimum of four consecutive days without

interruption, including two weekdays and two weekend days, analysed in 60 s epochs (Brage et al., 2005; Collings et al., 2017). The combined heart rate and movement sensor was attached to the child's chest with two standard electrocardiogram electrodes (Bio Protech Inc, Wonju, South Korea). The children were asked to wear the monitor continuously, including sleep and water-based activities, and not to change their usual behaviour during the monitoring period. Data on heart rate were cleaned and individually calibrated with parameters from the maximal exercise test and combined with movement sensor data to derive PA energy expenditure (PAEE). Instantaneous physical activity energy expenditure (PAEE), i.e. PA intensity, was estimated using branched equation modelling as explained, in detail earlier (Brage et al., 2007) and summarised as daily PA volume ($\text{kJ}\cdot\text{day}^{-1}\cdot\text{kg}^{-1}$) and time spent at certain levels of standard metabolic equivalents of task (METs) in minutes per day, weighting all hours of the day equally to reduce diurnal bias caused by imbalances in wear-time. Initially, the summarised data included 25 narrowly defined intensity categories. For the present analyses, we re-categorised these intensity categories into a broader format of sedentary time (≤ 1.5 METs), light PA (1.5–4 METs), moderate PA (>4–7 METs), vigorous PA (>7 METs) and MVPA (>4 METs), which have been commonly applied in investigations of PA among children and youth. In order to estimate the time spent sedentary whilst awake, we subtracted average daily sleep duration from total ST. We only included children who had sufficient valid data, i.e. a recording period of at least 48 h of wear data with the additional requirement that enough data were included from all four quadrants of a 24 h day to avoid bias from over-representation of specific parts of days (Brage et al., 2013). This resulted in at least 12 h of wear data from morning (3 am – 9 am), noon (9 am – 3 pm), afternoon / evening (3 pm – 9 pm), and night (9 pm – 3 am). Screen time was assessed by the PANIC Physical Activity Questionnaire filled out at home by the parents or caregivers of the children. Screen time included watching TV and videos, using a computer and playing video games, and using a mobile phone and playing mobile games. Screen time was calculated by summing up times spent in each type of screen time and was expressed in hours per week weighted by the numbers of weekdays and weekend days.

Assessment of diet quality

Food consumption and nutrient intake were assessed by food records administered by the parents on four pre-defined consecutive days, including two weekdays and two weekend days (99.5% of participants) or three

week-days and one weekend day (0.5% of participants), as described previously (Eloranta et al., 2011). The food records were analysed using the Micro Nutrica dietary analysis software, Version 2.5 (The Social Insurance Institution of Finland). We used the Finnish Children Healthy Eating Index (FCHEI) as a measure of overall diet quality (Kyttälä et al., 2014). FCHEI summarises the consumption of vegetables, fruit, and berries; vegetable oils and vegetable oil-based margarine; foods containing high amounts of sugar; fish; and low-fat (<1%) milk based on deciles of these dietary variables in the study population. A higher scores indicate a better diet quality.

Other assessments

Parental education was used as a measure of socioeconomic status. The parents were asked to report in a questionnaire their completed or ongoing educational degrees (vocational school or less, polytechnic or university). The degree of the more educated parent was used in the analyses.

Statistical methods

Statistical analyses were performed by the SPSS statistical software, version 25.0 (IBM corp. Armonk, NY, USA). The characteristics of children between boys and girls were compared using the Student's t-test for normally distributed continuous variables, the Mann-Whitney's U-test for continuous variables with skewed distributions, or the χ^2 -test for categorical variables. The associations of ST, screen time, PA, and diet quality with biomarkers for inflammation were investigated using linear regression analyses adjusted for age and sex. The data were further adjusted for BF% and parental education. To study the modifying effect of BF% and FCHEI, we dichotomised BF% using sex-specific medians and FCHEI using a sample-specific median. The modifying effect of BF% on the associations of ST, media time, PA, and diet quality with biomarkers for inflammation and modifying effect of FCHEI on the associations of ST, media time, and PA with biomarkers for inflammation were investigated using general linear models adjusted for age and sex.

Results

Characteristics of children

Girls were shorter and lighter and had higher BF% than boys (Table 1). Girls also accumulated less screen time, MPA, and VPA, and had lower PAEE than boys. Moreover, girls had higher leptin and lower IL-6 and TNF- α than boys.

Table 1. Characteristics of children.

| | All | Girls | Boys | P |
|--|--------------------------|-------------|-------------|--------|
| Age (years) | 7.6 (0.4) | 7.6 (0.4) | 7.7 (0.4) | 0.171 |
| Stature (cm) | 128.8 (5.5) | 127.6 (5.5) | 129.9 (5.3) | <0.001 |
| Weight (kg) | 25.9 (5.5) ¹ | 25.4 (5.8) | 26.8 (5.5) | 0.011 |
| Body mass index standard deviation score | -0.21 (1.1) | -0.22 (1.1) | -0.19 (1.1) | 0.802 |
| Body fat percentage (%) | 18.9 (10.2) ¹ | 20.3 (9.3) | 15.9 (9.8) | <0.001 |
| Prevalence of overweight and obesity | 8.2 / 3.6 | 9.4 / 3.6 | 7.1 / 3.5 | 0.705 |
| Proportion of children entered to clinical puberty (%) | 2.1 | 2.6 | 1.5 | 0.448 |
| Sedentary time (min/d) | 205 (171) ¹ | 211 (182) | 203 (156) | 0.117 |
| Media time (min/d) | 94.3 (62.1) ¹ | 86.8 (55.7) | 98.6 (60.0) | <0.001 |
| Light physical activity (min/d) | 510 (106) | 518 (110) | 503 (102) | 0.170 |
| Moderate physical activity (min/d) | 81.4 (57.9) ¹ | 73.0 (56.9) | 86.7 (66.8) | <0.001 |
| Vigorous physical activity (min/d) | 17.0 (27.5) ¹ | 12.0 (18.2) | 23.0 (38.5) | <0.001 |
| Moderate-to-vigorous physical activity (min/d) | 102 (83.4) ¹ | 88.6 (73.4) | 128 (92.7) | <0.001 |
| Physical Activity Energy Expenditure (kJ/kg/d) | 97.5 (31.8) | 89.3 (28.3) | 105 (33.1) | <0.001 |
| Finnish Children Healthy Eating Index | 23.1 (7.1) | 23.5 (6.5) | 22.7 (7.7) | 0.294 |
| hs-CRP (mg/L) | 0.3 (0.3) ¹ | 0.29 (0.33) | 0.29 (0.22) | 0.052 |
| Leptin (ng/mL) | 3.7 (3.1) ¹ | 4.3 (3.8) | 3.1 (2.3) | <0.001 |
| Interleukin (IL)-6 (pg/mL) | 0.9 (0.9) ¹ | 0.8 (0.7) | 1.1 (0.9) | 0.005 |
| High-molecular-weight adiponectin (µg/mL) | 8.1 (5.5) ¹ | 7.8 (5.5) | 8.4 (5.5) | 0.363 |
| Tumour necrosis factor (TNF)-α (pg/mL) | 15.8 (31.8) ¹ | 13.4 (27.5) | 17.8 (36.3) | 0.037 |
| Glycoprotein acetyls (mmol/L) | 0.76 (0.11) | 0.76 (0.11) | 0.76 (0.11) | 0.759 |

Data are from the Student *t*-test or Mann-Whitney U test for continuous variables and from the Chi-square test for categorical variables and are displayed as means (SD), medians (IQR), or percentages (%).

Associations of sedentary time, physical activity, and diet quality with biomarkers for inflammation

ST was directly and MPA, VPA, MVPA, and PAEE inversely associated with hs-CRP and IL-6 after adjustment for age and sex (Table 2). ST was directly and LPA, MPA, VPA, MVPA, and PAEE inversely associated with leptin after adjustment for age and sex. MPA, MVPA, and PAEE were inversely associated with glycoprotein acetyls after adjustment for age and sex. However, none of these associations were statistically significant after further adjustment for BF% ($p=0.084-0.895$). FCHEI was not associated with biomarkers for inflammation after

adjustment for age and sex. BF% was directly associated with hs-CRP, leptin, IL-6, and glycoprotein acetyls. Further adjustment for parental education had no effect on these associations.

Modifying effect of body fat percentage on the associations of sedentary time, physical activity, and diet quality with biomarkers for inflammation

ST time directly associated with hs-CRP in children with higher BF% ($\beta=0.160$, 95% CI=0.018-0.301, $p=0.027$) but not among those with lower BF%

Table 2. Associations of sedentary time, physical activity, and diet quality with biomarkers for inflammation in children.

| | Hs-CRP | Leptin | Interleukin-6 | Adiponectin | Tumour necrosis factor-α | Glycoprotein acetyls |
|------------|-------------------------------------|-------------------------------------|-----------------------------------|-----------------------|--------------------------|-----------------------------------|
| Media time | 0.062 (-0.040-0.164) | -0.012 (-0.109-0.086) | 0.014 (-0.088-0.115) | -0.046 (-0.148-0.056) | 0.033 (-0.069-0.134) | -0.057 (-0.159-0.045) |
| ST | 0.170 (0.070-0.269)** | 0.355 (0.265-0.445)*** | 0.105 (0.005-0.205)* | -0.030 (-0.131-0.070) | -0.033 (-0.133-0.068) | 0.100 (0.000-0.201) |
| LPA | -0.070 (-0.171-0.030) | -0.201 (-0.295 to -0.107)*** | -0.036 (-0.136-0.064) | 0.040 (-0.060-0.140) | 0.061 (-0.039-0.161) | -0.041 (-0.141-0.059) |
| MPA | -0.168 (-0.269 to -0.067)** | -0.301 (-0.395 to -0.208)*** | -0.114 (-0.216 to -0.013)* | 0.004 (-0.099-0.106) | -0.030 (-0.132-0.072) | -0.103 (-0.205 to -0.001)* |
| VPA | -0.139 (-0.233 to -0.024)* | -0.318 (-0.414 to -0.222)*** | -0.107 (-0.211 to -0.002)* | -0.053 (-0.158-0.052) | -0.061 (-0.166-0.045) | -0.080 (-0.186-0.132) |
| MVPA | -0.191 (-0.294 to -0.088)*** | -0.372 (-0.465 to -0.279)*** | -0.136 (-0.240 to -0.033)* | -0.015 (-0.120-0.090) | -0.047 (-0.152-0.057) | -0.117 (-0.221 to -0.013)* |
| PAEE | -0.184 (-0.286 to -0.083)*** | -0.409 (-0.499 to -0.318)*** | -0.104 (-0.206 to -0.001)* | -0.022 (-0.128-0.081) | -0.017 (-0.120-0.086) | -0.110 (-0.213 to -0.007)* |
| FCHEI | 0.043 (-0.057-0.144) | 0.000 (-0.096-0.097) | 0.032 (-0.068-0.133) | -0.010 (-0.110-0.091) | 0.015 (-0.085-0.116) | 0.045 (-0.055-0.146) |
| BF% | 0.382 (0.285-0.480)*** | 0.851 (0.797-0.904)*** | 0.131 (0.027-0.235)* | -0.043 (-0.148-0.062) | -0.013 (-0.118-0.091) | 0.302 (0.201-0.403)*** |

Notes: Data are standardised regression coefficient and their 95% confidence intervals from multivariate linear regression analyses adjusted for age and sex. ST, sedentary time; LPA, light physical activity; MPA, moderate intensity physical activity; VPA, vigorous intensity physical activity; PAEE, physical activity energy expenditure; MVPA, moderate-to-vigorous physical activity; FCHEI, Finnish Children Healthy Eating Index; BF%, body fat percentage.

* $p<0.05$; ** $p<0.01$; *** $p<0.001$.

($\beta=0.071$, 95% CI= -0.072 – 0.214 , $p=0.329$, $p=0.055$ for interaction).

VPA was inversely associated with hs-CRP ($p=0.003$ for interaction), leptin ($p=0.034$ for interaction), IL-6 ($p=0.036$ for interaction), and glycoprotein acetyls ($p=0.027$ for interaction) in children with higher BF% but not in children with lower BF% (Table 3). MVPA was inversely associated with hs-CRP in children with higher BF% but not among those with lower BF% ($p=0.019$ for interaction). PAEE was inversely associated with CRP in children with higher BF% but not among those with lower BF% ($p=0.019$ for interaction) (Table 3). PAEE was also inversely associated with adiponectin in children with lower BF% ($\beta=-0.160$, 95% CI= -0.308 to -0.012 , $p=0.034$) but not among those with higher BF% ($\beta=0.088$, 95% CI= -0.057 – 0.233 , $p=0.231$, $p=0.016$ for interaction).

Modifying effect of diet quality on the associations of sedentary time, physical activity, and diet quality with inflammatory biomarkers

MPA was inversely associated with hs-CRP in children with a lower FCHEI ($\beta=-0.254$, 95% CI= -0.397 to -0.112 , $p=0.001$) but not among those with a higher FCHEI ($\beta=-0.084$, 95% CI= -0.229 – 0.061 , $p=0.253$, $p=0.036$ for interaction) after adjustment for age and sex. MVPA was inversely associated with hs-CRP in children with a lower FCHEI ($\beta=-0.283$, 95% CI= -0.427 to -0.139 , $p<0.001$) but not among those with a higher FCHEI ($\beta=-0.092$, 95% CI= -0.240 – 0.056 , $p=0.221$, $p=0.034$ for interaction). PAEE was inversely associated with hs-CRP in children with a lower FCHEI ($\beta=-0.256$, 95% CI= -0.400 to -0.111 , $p=0.001$) but not among those with a higher FCHEI ($\beta=-0.107$, 95% CI= -0.253 – 0.038 , $p=0.147$, $p=0.053$ for interaction). However, the associations of MPA ($\beta=-0.121$, 95% CI= -0.259 – 0.018 , $p=0.088$), MVPA ($\beta=-0.117$, 95% CI= -0.264 – 0.029 ,

$p=0.117$), and PAEE ($\beta=-0.057$, 95% CI= -0.208 – 0.094 , $p=0.456$) with hs-CRP in children with lower FCHEI attenuated markedly and were no longer statistically significant after further adjustment for BF%

Discussion

We found that BF% was strongly and directly associated with circulating levels of biomarkers for inflammation and it also largely explained the associations of ST, MPA, VPA, MVPA, and PAEE with these biomarkers for inflammation in children 6–8 years of age. We also observed that higher levels of ST and lower levels of PA were associated with higher circulating levels of biomarkers for inflammation in children with a higher BF% but not among those with a lower BF%. Moreover, we found that PA was inversely associated with circulating levels of biomarkers for inflammation in children with poorer diet quality but not among those with better diet quality. These inverse associations were mainly observed between PA at least moderate intensity and hs-CRP, leptin, IL-6, and glycoprotein acetyls.

In contrast to the results of some previous studies in children (Poitras et al., 2016), we found inverse associations of PA with hs-CRP, leptin, IL-6 and direct associations of ST with hs-CRP, leptin, and IL-6 in children. Individual biomarkers of inflammation often exhibit a high intra-individual variability (Connelly et al., 2017). Therefore, our findings on the inverse associations of MPA, MVPA, and PAEE with glycoprotein acetyls derived from NMR metabolomics, a composite indicator of systemic inflammation providing novel and more stable (Connelly et al., 2017) evidence on the inverse association between PA and systemic inflammation. However, all of these associations in our study were largely explained by BF%. Our results are supported by evidence suggesting that reductions in body fat mass mediates the positive effects of exercise training on

Table 3. Associations of physical activity with biomarkers for inflammation in children with higher or lower body fat percentage.

| | Hs-CRP | | Leptin | | Interleukin-6 | | Glycoprotein acetyls | |
|------|------------------------------|--|------------------------------|---|-----------------------------|---------------------------------------|-----------------------------|------------------------------|
| | Body fat percentage | | Body fat percentage | | Body fat percentage | | Body fat percentage | |
| | <median | \geq median | <median | \geq median | <median | \geq median | <median | \geq median |
| VPA | 0.031 (-0.122– 0.184) | -0.186 (-0.332 to -0.039)* | -0.007 (-0.146– 0.131) | -0.344 (-0.477 to -0.212)*** | 0.033 (-0.183– 0.117) | -0.181 (-0.328 to -0.033)* | 0.060 (-0.092– 0.213) | -0.143 (-0.291– 0.005) |
| MVPA | -0.070 (-0.221– 0.080) | -0.215 (-0.360 to -0.069)** | | | | | | |
| PAEE | -0.058 (-0.209– 0.092) | -0.193 (-0.336 to -0.050)** | | | | | | |

Note: Data are standardised regression coefficient and their 95% confidence intervals from multivariate linear regression analyses adjusted for age and sex. VPA, vigorous physical activity; MVPA, moderate-to-vigorous physical activity; PAEE, physical activity energy expenditure.

* $p<0.05$; ** $p<0.01$; *** $p<0.001$.

biomarkers for inflammation (García-Hermoso et al., 2017). Nevertheless, MVPA was inversely associated with hs-CRP, IL-6, and adiponectin and ST was directly associated with adiponectin independent of adiposity in Danish children (Nielsen et al., 2016). Furthermore, in contrast to the results of some previous studies (Dong et al., 2017), we found no association between screen time and biomarkers for inflammation. However, we observed that ST was directly associated with biomarkers for inflammation although these associations were mostly explained by BF%. The inconsistent findings may be explained by differences in maturation status and PA levels between study populations and methods used to assess screen time and ST. Furthermore, it is possible that increased ST, not screen time per se, is associated with increased systemic inflammation through increased adiposity. Therefore, these results together suggest that increased levels of PA have potential to decrease systemic inflammation, but it is possible that the associations are mediated by decreased adiposity.

We found that PA at least at moderate intensity was associated with biomarkers for inflammation only in children with a higher BF%. These results agree with our previous observations suggesting that higher levels of MVPA and lower levels of ST were associated with lower insulin resistance only in children with higher levels of adiposity (Haapala et al., 2020) suggesting that increasing PA could decrease systemic inflammation and insulin resistance particularly among overweight and obese children. In contrast to the results of some intervention studies in youth showing that exercise training increased adiponectin in obese children (García-Hermoso et al., 2017), we observed an inverse association between PAEE and adiponectin in children with lower BF% and no statistically significant association in children with higher BF%. Furthermore, Nielszen et al. (2016) suggested that the negative association between MVPA and adiponectin could be due to lower insulin resistance in children with higher levels of MVPA as they also reported an inverse association between insulin resistance and adiponectin (Nielsen et al., 2016). Similarly, Su et al. (2011) reported that individuals with lower levels of insulin and body fat mass also had lower adiponectin than those with higher levels of insulin and fat mass. We have previously reported that children with higher levels of MVPA or PAEE and lower levels of adiposity have lower insulin resistance than other children (Haapala et al., 2020). It is, therefore, possible that the inverse association between PAEE and adiponectin in the present study and between MVPA and adiponectin in the study by Nielszen et al. is due to adaptive mechanisms driven

by insulin resistance (Nielsen et al., 2016). We have observed that PAEE, from all measure of PA in our study, has the strongest association with insulin resistance (Haapala et al., 2020; Väistö et al., 2019) and BF% (Väistö et al., 2019), which may explain why PAEE also had the strongest association with adiponectin in the present study.

We found that BF% was the strongest determinant of biomarkers for inflammation and even if the mechanisms of low-grade inflammation are not fully understood, it has been shown that excess adiposity leads to secretion of several cytokines and adipokines that activate the immune system. Gleeson et al. suggested that both the reduction in visceral fat mass and the anti-inflammatory environment induced by each exercise session might elicit long-term anti-inflammatory effects (Gleeson et al., 2011).

In line with the results of previous observational studies in children, we did not find statistically significant associations between diet quality and biomarkers for inflammation (Liese et al., 2018). One reason for not observing such associations is that children in our population sample were relatively healthy and most of them were normal weight and physically active. However, we found that higher levels of PA were associated with lower levels of systemic low-grade inflammation in children with poorer diet quality. These results suggest that poor diet quality together with low PA levels may increase systemic low-grade inflammation in children, although diet quality did not have independent associations with biomarkers for inflammation. Nevertheless, the associations of PA at different intensity levels with hs-CRP in children with poorer diet quality were partly explained by BF% suggesting that adiposity may also mediate the combined association of diet quality and PA with biomarkers for inflammation.

The strengths of the present study include a relatively large and representative sample of children with valid and reproducible assessment of free-living PA and ST by individually calibrated combined movement and heart rate sensing, diet quality by 4-day dietary records, body composition using whole-body dual-energy X-ray absorptiometry, and several biomarkers for inflammation in a population sample of children. However, we used 60 s epochs in the assessment of ST and PA because of limited memory capacity of the monitor. This may have influenced especially the amount of VPA in the current study (Gao et al., 2018) and weaken the associations of VPA with biomarkers for inflammation. Our study was cross sectional which limits our ability to make causal inferences. Furthermore, our study included mainly Caucasian children and therefore our results cannot be directly generalised to the

samples of different ethnical backgrounds. Systemic low-grade inflammation and its relationship to the pathogenesis of cardiometabolic diseases is multifaceted, and no single optimal biomarker for it exists. Although we studied the associations ST, PA, and diet quality with several possible biomarkers for inflammation, there are many other pro- or anti-inflammatory factors that could have been measured. Finally, the applicability and clinical relevance of these biomarkers for inflammation in identifying children at increased risk of cardiometabolic diseases remains unclear.

In conclusion, we found that higher levels of MPA, VPA, MVPA, and PAEE and lower levels of ST were associated with lower circulating levels of biomarkers of inflammation among children, but these associations were largely explained by BF%. Furthermore, we observed that PA was inversely associated with these biomarkers for inflammation in children with a higher BF%. In addition, our study provides some evidence that higher levels of PA could improve circulating biomarkers of inflammation in children with poorer diet quality. More research on the long-term effects of PA and diet as well as their synergistic effects on biomarkers for inflammation is warranted.

Disclosure statement

No potential conflict of interest was reported by the author(s).

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Availability of data and material

The data that support the findings of this study are available from the corresponding author, [EAH], upon reasonable request.

Author contributions

EAH analysed the data. EAH and JKI drafted the manuscripts. JV, AV, TS, A-ME, UE, US, SB, MA, and TAL collected and processed the data for analyses. MHL, AV contributed to planning the manuscript and interpreting the results and reviewed the manuscript. All authors provided significant intellectual contribution to the manuscript.

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