**The contribution of physical fitness to individual and ethnic differences in risk markers for type 2 diabetes in children: the Child Heart and health Study in England (CHASE)**

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**Running header**

Physical fitness and diabetes risk in children

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**Conflict of interest statement**

We declare that we have no conflicts of interest.

**Author contributions**

Professor Whincup, Professor Owen, Professor Cook and Dr Rudnicka conceptualized and designed the study, and coordinated and supervised data collection. All authors (Dr Nightingale, Dr Rudnicka, Ms Kerry-Barnard, Dr Donin, Dr Brage, Ms Westgate, Professor Ekelund, Professor Cook, Professor Owen and Professor Whincup) interpreted data. Dr Nightingale carried out the initial analyses and drafted the initial manuscript. All authors critically reviewed and revised the manuscript and approved the final manuscript as submitted. All authors agree to be accountable for all aspects of the work.

***Abstract***

**Background:** The relationships between physical fitness and risk markers for type 2 diabetes (T2D) in children and the contribution to ethnic differences in these risk markers have been little studied. We examined associations between physical fitness and early risk markers for T2D and cardiovascular disease in 9-10 year-old UK children.

**Methods**: Cross-sectional study of 1445 9-10 year-old UK children of South Asian, black African-Caribbean and white European origin. A fasting blood sample was used for measurement of insulin, glucose (from which HOMA-insulin resistance (IR) was derived), HbA1c, urate, C-reactive protein (CRP) and lipids. Measurements of blood pressure (BP) and fat mass index (FMI) were made; physical activity was measured by accelerometry. Estimated VO2 max was derived from a submaximal fitness step test. Associations were estimated using multilevel linear regression.

**Results:** Higher VO2 max was associated with lower FMI, insulin, HOMA-IR, HbA1c, glucose, urate, CRP, triglycerides, LDL-cholesterol, BP and higher HDL-cholesterol. Associations were reduced by adjustment for FMI, but those for insulin, HOMA-IR, glucose, urate, CRP, triglycerides and BP remained statistically significant. Higher levels of insulin and HOMA-IR in South Asian children were partially explained by lower levels of VO2 max compared to white Europeans, accounting for 11% of the difference.

**Conclusions:** Physical fitness is associatedwith risk markers for T2D and CVD in children, which persist after adjustment for adiposity. Higher levels of IR in South Asians are partially explained by lower physical fitness levels compared to white Europeans. Improving physical fitness may provide scope for reducing risks of T2D.

***Introduction***

Higher levels of physical fitness in adults are associated with a lower risk of developing type 2 diabetes (T2D) (1;2) and lower rates of cardiovascular disease (CVD) mortality and all-cause mortality (3;4). South Asian adults are at higher risk of developing T2D, stroke and coronary heart disease compared to white Europeans in the UK and other Western countries (5;6). Evidence suggests that ethnic differences in risk markers for T2D (particularly insulin resistance) are apparent in pre-pubertal children with higher levels T2D risk markers in South Asian children compared to white Europeans (7). Ethnic differences in physical fitness are also apparent in childhood; we have shown in a recent report lower levels of physical fitness in South Asians compared to white Europeans (8). These differences in physical fitness could potentially help to explain ethnic differences in risk markers for T2D. We therefore examined associations between physical fitness and risk markers for T2D and CVD in a study of British schoolchildren and examined the contribution of differences in physical fitness to ethnic differences in risk markers for T2D between South Asians and white Europeans. It has been suggested that physical fitness and physical activity could have separate and independent effects on metabolic risk in childhood (9). Given the complex relationship between physical activity and fitness, we also examine the independent associations between physical activity, physical fitness and risk for T2D and CVD.

***Methods***

The Child Heart and Health Study in England (CHASE) was a cross-sectional study of the health of 9-10 year old children of white European, South Asian and black African-Caribbean origin carried out between 2004 and 2007 in London, Birmingham and Leicester. Ethical approval was obtained from the relevant multicentre research ethics committee and informed, written consent was obtained from the parents of all participating children. Full details have been published elsewhere (7;10;11). In brief, the study was based on a random sample of 200 state primary schools, half were drawn from a sampling frame of schools with a high prevalence of UK South Asian children and half were drawn from a sampling frame with a high prevalence of UK black African-Caribbean children. This report is based on the final phase of the study carried out between January 2006 and February 2007 in which children from 81 schools had assessments of physical fitness and physical activity.

A single survey team consisting of three trained Research Nurses and two Research Assistants carried out all assessments in schools. Physical measurements included height, weight, and arm-to-leg bioelectrical impedance, using the Bodystat 1500 monitor (Bodystat Ltd, Isle of Man, UK). Fat mass was derived from bioelectrical impedance using gender- and ethnic-specific equations derived for children of this age group (12). Fat mass index [fat mass (kg)/height(m)5] (FMI) was derived to be uncorrelated with height (r=-0.04) which has been shown previously to provide a more valid measure of body fatness in this multi-ethnic study population (13). Systolic and diastolic BP was measured twice in the right arm using an Omron HEM-907 (Omron Electronics, Milton Keynes, UK). Mean systolic and diastolic BP were adjusted for cuff size using a previously validated method (14).

Participants provided a blood sample after an overnight fast; children who reported having eaten breakfast were excluded from the analysis. Breakfast was provided after the sample was collected. Serum was separated and frozen on dry ice immediately after collection for measurement of insulin. Urate was measured in serum using an enzymatic assay (15). Samples were shipped to a central laboratory within 48 hours of collection. Details of insulin, glucose, glycated haemoglobin (HbA1c), C-reactive protein and blood lipids assays have been previously reported (7;10). Homeostasis model assessment (HOMA) equations (16) were used to derive insulin resistance from fasting insulin and glucose measurements.

An 8-min submaximal step test was performed by participating childrenas previously described (17;18). In brief, participants followed an audible prompt which instructed them to increase their step frequency progressively from 15 to 32.5 body lifts/min on a 150mm high step whilst wearing a combined heart rate (HR) and movement sensor (Actiheart, CamNtech, Papworth, UK). The test was terminated if the participant was unable to sustain the prescribed step frequency. At the end of the test, two minutes of seated recovery was recorded. ECG and acceleration waveforms were recorded at 128 and 32 Hz sampling respectively. VO2max was estimated using a similar method to that used in the Health Survey for England 2008 (19). In brief, predicted workload was regressed against instantaneous HR (expressed above resting HR) and 1 minute recovery HR was extracted using quadratic regression against recovery time. This was combined with resting HR and test duration to calculate the submaximal relationship between workload and HR, which was extrapolated to predicted maximal HR to predict maximal work capacity. This result was converted to VO2 max by adding an estimate of resting metabolic rate and then dividing by the energetic value of oxygen (19).

Objective measures of physical activity were made using an accelerometer recorded at 5 second epochs (GT1M; ActiGraph LLC, Pensacola, FL, USA), worn on the participant’s left hip during waking hours for seven days; the device was removed for water-based activities. Non-wear time was defined as a period of at least 20 consecutive minutes of zero counts, and was excluded from the analyses. Physical activity was summarised as total daily counts. Participants with at least one day of valid data (defined as at least 600 minutes of registered time) were included in the analysis.

Ethnic origin was defined using information from a parental questionnaire on the ethnicity of both parents where available (63%), or using the parentally defined ethnic origin of the child (36%). In a small number of children (1%) where this information was not available, ethnic origin was defined using information on parental and grand-parental place of birth provided by the child, cross-checked with observer assessment of ethnic origin. Children were defined as white European, South Asian, black African-Caribbean and ‘other’ ethnicity.

**Statistical methods**

Statistical analyses were carried out using Stata/SE software (Stata/SE 13 for Windows; StataCorp LP, College Station, TX). All risk markers were inspected for normality; all variables except systolic and diastolic BP followed a log-normal distribution and were therefore log transformed in analyses. A regression calibration method to allow for measurement error in physical activity counts (20) was used to allow for within-child variation by day of the week and variation in the number of days of recording (between 1 and 7), which provides an unbiased average for each child. The majority of participants (83%) wore the ActiGraph for at least 4 days. We examined the shape of associations between VO2 max and risk markers by deciles of VO2 max and did not find any evidence of departures from linearity. Hence, associations between estimated VO2 max and risk markers were assessed using multilevel linear models adjusted for sex, age (in quartiles), ethnic group, month of measurement and height (fitted as fixed effects) and school fitted as a random effect to take into account clustering of children within schools. The effect of additional adjustment for FMI was also examined. We also examined whether associations between VO2 max and risk markers were consistent in boys and girls and ethnic groups. To elucidate whether ethnic differences in estimated risk markers were explained by differences in physical fitness, we examined the effect of adjustment for VO2 max on ethnic differences in risk markers. In separate models we investigated whether associations between physical fitness and risk markers were consistent across tertiles of physical activity.

***Results***

The participation rate for children taking part in this phase of the study was 63%; of those who took part 1979 (89%) provided a fasting blood sample and had complete data. Estimated VO2 max values were available for a total of 1445 participants (50% female with a mean age of 10.0 years [95% reference range 9.2, 10.7 years]) who had provided a fasting blood sample and had measurements of FMI. Of these children, 1083 provided a valid estimate of physical activity. The reasons for missing physical activity data were refusal to wear the device, failure to return the device or providing less than one day with 600 minutes of wear time of the device. Adjusted mean levels of VO2 max and risk markers for T2D and CVD are shown in Table 1. Boys had higher estimated levels of VO2 max than girls; South Asians had lower levels and black African-Caribbeans had higher levels of estimated VO2 max compared to white Europeans.

Associations between physical fitness (estimated VO2 max) and risk markers for T2D and CVD are shown in Table 2. Estimated VO2 max was inversely associated with FMI, fasting insulin, HOMA-IR, HbA1c, fasting glucose, urate, CRP, triglyceride, LDL-cholesterol and systolic and diastolic BP, and positively associated with HDL-cholesterol. Associations between estimated VO2 max and insulin, HOMA-IR, glucose, urate, CRP and triglyceride were attenuated by adjustment for FMI by between 32% and 63%. Further adjustment for fat free mass index did not further attenuate these associations (data available from authors). Associations with HbA1c, HDL- and LDL-cholesterol were completely explained by adjustment for FMI and those with systolic and diastolic blood pressure were attenuated by 31% and 16% respectively. After adjustment for FMI, a one IQR increase in estimated VO2 max was associated with lower fasting insulin, HOMA-IR, fasting glucose, urate, CRP and triglyceride of 7.7%, 7.7%, 0.7%, 2.2%, 20.8% and 5.3% respectively. Similarly, a one IQR increase in estimated VO2 max was associated with decreases in systolic and diastolic BP of 1.7 and 2.7 mmHg respectively. These associations were broadly similar in boys and girls (supplementary Table 1) and in South Asians, black African-Caribbeans, white Europeans and ‘other’ ethnic groups (supplementary Table 2) with the exception of LDL-cholesterol which was inversely associated with VO2 max in all ethnic groups except black African-Caribbeans.

In a subset of 1083 children with objective measures of both physical fitness and physical activity, associations between risk markers for T2D/CVD and physical activity and VO2 max are shown in supplementary Table 3. Associations were stronger between risk markersfor T2D and estimated VO2 max compared to those for overall physical activity. Following adjustment for FMI however, associations between fasting insulin, HOMA-IR and physical activity were stronger than those for estimated VO2 max. To explore whether the associations between physical fitness and risk markers were modified by physical activity level, associations between VO2 max and risk markers were examined by tertiles of physical activity in supplementary Table 4. The inverse associations between estimated VO2 max and fasting insulin, HOMA-IR and triglyceride were strongest among children in the lowest tertile of physical activity and weakest among children in the highest tertile; a graphical representation for fasting insulin is shown in supplementary Figure 1.

Ethnic differences in risk markers for T2D between South Asians and white Europeans and the effect of adjustment for physical fitness are shown in Table 3. South Asians had higher levels of fasting insulin, HOMA-IR, HbA1c, fasting glucose and triglyceride and lower levels of HDL-cholesterol. Adjustment for VO2 max reduced these ethnic differences in insulin, HOMA-IR, glucose and triglyceride by approximately 11%; differences in HbA1c and HDL-cholesterol were reduced by 4% and 13% respectively. The borderline ethnic difference in CRP was reduced by 50% by adjustment for estimated VO2 max. In the subset of 1083 participants with measurements of physical activity, Supplementary Table 5 shows whether adjustment for physical activity counts in addition to VO2 max would further explain the ethnic differences in risk markers for T2D. For fasting insulin, HOMA-IR, triglycerides and HDL-cholesterol adjustment for physical activity as well as VO2 max, further reduced the difference between South Asians and white Europeans by 5-7%.

***Discussion***

***Main findings***

This study provides strong evidence that low levels of physical fitness are associated with risk markers for T2D and CVD in this multi-ethnic population of school children. These associations were attenuated but remained statistically significant after adjustment for measures of body fat, suggesting that associations were at least partly reduced by the effect of adjustment for adiposity, though additional adjustment for fat free mass had no material effect. Moreover, ethnic differences in risk markers for T2D, particularly increased levels of insulin resistance in South Asians compared to white European children, were partially explained by physical fitness (approximately 11%). Stratified analysis suggested that the association between physical fitness and risk markers differed by level of physical activity, with stronger associations being observed at lower levels of physical activity.

***Relation to previous studies***

The finding in the present study that objective measures of physical fitness are inversely associated with risk markers for T2D and CVD in childhood, is consistent with a small number of studies, which have used similar objective methodologies (21). In particular, findings from the EYHS showed that maximal ergometer assessments of cardiorespiratory fitness were inversely related to metabolic risk markers at both 9 and 15 years of age (21;22). These associations have since been demonstrated in prospective studies, providing further evidence of a causal relationship (23). We have previously reported that both physical activity and adiposity are associated with risk markers for T2D and CVD in this study population (20;24), and that these associations are partially, but not wholly, mutually independent.

The present study confirms the association between physical fitness and cardiometabolic risk markers, but is novel in suggesting that the association observed might be altered at different levels of physical activity, with stronger associations between physical fitness and metabolic risk at lower levels of physical activity. This is consistent with a report based on 589 Danish children in the European Youth Heart Study (EYHS) but this was not confirmed in the full EYHS study population, which suggested little or no modifying influence of physical activity (21). Intuitively it would seem plausible that physical fitness- metabolic risk associations might be modified by levels of physical activity, given the established link between activity and fitness, where previous physical activity interventions have been shown to improve physical fitness (25-27). However, this apparent interaction could also reflect differences in the precision of measurement of physical activity and physical fitness, and the possibility of non-linear associations between them. A sensitivity analysis examining these associations in children with at least 6 days of physical activity data (54% of the sample) showed that stronger associations between risk markers and physical fitness persisted at lower levels of physical activity (data available from authors).

This study suggests that adiposity is an important potential mediator of the associations between physical fitness and risk markers for T2D and CVD. Adjustment for objective markers of body fatness (FMI), reduces fitness-metabolic associations by between one-third to two-thirds, and weakens associations with some cardiovascular risk markers even further. Previous studies have been inconsistent about the confounding influence of adiposity (21;28-30), and whether adiposity confounds, mediates or modifies associations between fitness and metabolic risk markers in early life. Randomised controlled trials are needed to further elaborate on the causal nature of these associations. Teasing out the potential interrelationship between these factors may have ramifications for future interventions, which may need to target improvements in both levels of physical activity and fitness, as well as adiposity, in order to improve metabolic and cardiovascular health from an early age.

Previous studies examining the consistency of early physical fitness and T2D/CVD risk marker associations have suggested that these are similar in boys and girls, and show little regional variations amongst populations of European ancestry (31). The present study confirms that associations were similar by sex, and extends knowledge by showing consistent associations in children of more diverse ethnic origin, including those of white European, South Asian, and black African-Caribbean origin. As far as we are aware, the current study is also novel in showing that early ethnic differences in risk markers for T2D, where those of South Asian origin have a worse metabolic profile (32) can be at least partly “explained” by levels of physical fitness. Risk markers for T2D in black African-Caribbean children showed smaller differences from white Europeans, and physical fitness levels were higher compared to white Europeans suggesting this would not help to explain any of the difference.

***Strengths and limitations***

There are a number of strengths and weaknesses worthy of consideration. A strength of the study is the objective measure of physical fitness using a submaximal fitness test to estimate VO2 max (8), which has previously been used in a nationally representative large-scale study including this age group (19). The use of a submaximal fitness test means that VO2 max must be extrapolated, however, this test was chosen for its suitability in large scale studies (19) and to encourage participation amongst a target population with low levels of physical activity (11). This was accompanied by objective assessment of physical activity, using waist worn accelerometry over a 7-day period, which provides a reliable assessment of habitual levels of physical activity (20). While both these measures provide accurate assessment, their comparative accuracy should be considered. The former provides a single measure of a longer-term attribute (indicative of fitness levels over a month or longer), whereas accelerometry provides a measure of a short-term behaviour, which may not fully capture habitual levels of physical activity behaviour. Hence, while the fitness-metabolic or cardiovascular risk marker associations might be considered robust, the predictive value of physical activity may have been underestimated. Specifically fitness is partly a function of longer term sustained physical activity, so it may be representing effects of past activity and/or more intense activity.

Another key strength is the measurement of important early risk markers for T2D (including HOMA insulin resistance, HbA1c, triglyceride and urate) and CVD (including systolic and diastolic blood pressure and LDL cholesterol). In addition, adiposity was measured using bioelectrical impedance with ethnic- and gender-specific equations for the prediction of fat mass in this age group; this method has been shown to provide valid assessments of adiposity in a multi-ethnic population (12). Whilst the response rate for the present study was modest, the sample contained balanced representation of South Asians, black African-Caribbeans and white Europeans and of boys and girls. Furthermore, risk markers were similar in those who completed the fitness test and those that did not (data not presented). Ethnic differences in metabolic risk markers in the CHASE study have previously been reported in the whole study population (32); patterns of ethnic differences were broadly similar in this subset of data.

***Implications***

The results presented in the present study show that ethnic differences in physical fitness account for up to 11% of the difference in insulin resistance and glycaemia markers between South Asians (who had higher levels of these markers) and white Europeans. This has potential implications for the early prevention of T2D in South Asians. Clinical trials are needed to determine whether efforts to improve physical fitness (potentially through increases in vigorous intensity physical activity) could help to improve metabolic health in children, perhaps especially among South Asian children/adolescents in whom risks of insulin resistance and T2D are particularly high (5;6).

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Table 1: Estimated VO2 max and risk markers for type 2 diabetes and cardiovascular disease by sex and by ethnic group

|  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | Geometric mean/mean\* (95% CI) | | | | | | | | | | | |  |
| Boys (n=723) | | Girls (n=722) | | White European (n=389) | | South Asian (n=373) | | Black African-Caribbean (n=346) | | Other (n=337) | | p(ethnic group)‡ |
| Fat mass index (kg/m5) | 2.0 | (1.9, 2.0) | 2.2 | (2.1, 2.3) | 2.1 | (2.0, 2.2) | 2.1 | (2.1, 2.2) | 1.9 | (1.8, 2.0) | 2.2 | (2.1, 2.3) | <0.0001 |
| Estimated VO2 max (ml O2/min/kg)\* | 40.8 | (40.4, 41.2) | 37.9 | (37.5, 38.3) | 39.3 | (38.9, 39.8) | 38.5 | (37.9, 39.0) | 40.3 | (39.8, 40.8) | 39.3 | (38.8, 39.8) | <0.0001 |
| Physical activity counts (x1000)\*† | 437 | (428, 445) | 364 | (356, 373) | 400 | (390, 410) | 378 | (367, 390) | 422 | (411, 432) | 398 | (387, 408) | <0.0001 |
| Insulin (mU/l) | 6.53 | (6.16, 6.92) | 8.05 | (7.59, 8.53) | 6.26 | (5.84, 6.72) | 8.68 | (8.03, 9.39) | 6.64 | (6.16, 7.16) | 7.69 | (7.14, 8.28) | <0.0001 |
| HOMA-IR | 0.82 | (0.78, 0.87) | 1.00 | (0.94, 1.06) | 0.78 | (0.73, 0.83) | 1.08 | (1.01, 1.17) | 0.83 | (0.77, 0.90) | 0.96 | (0.89, 1.03) | <0.0001 |
| HbA1c (%) | 5.29 | (5.26, 5.31) | 5.29 | (5.26, 5.31) | 5.23 | (5.20, 5.26) | 5.33 | (5.29, 5.37) | 5.30 | (5.27, 5.34) | 5.29 | (5.25, 5.32) | <0.001 |
| HbA1c (mmol/mol) | 34.0 | (33.8, 34.3) | 34.1 | (33.8, 34.4) | 33.5 | (33.1, 33.8) | 34.5 | (34.1, 34.9) | 34.2 | (33.8, 34.6) | 34.1 | (33.7, 34.5) | <0.001 |
| Glucose (mmol/l) | 4.51 | (4.49, 4.54) | 4.43 | (4.40, 4.45) | 4.45 | (4.41, 4.48) | 4.51 | (4.47, 4.55) | 4.43 | (4.39, 4.46) | 4.49 | (4.46, 4.53) | 0.002 |
| Urate (mmol/l) | 0.22 | (0.21, 0.22) | 0.23 | (0.22, 0.23) | 0.22 | (0.22, 0.23) | 0.23 | (0.22, 0.23) | 0.21 | (0.20, 0.21) | 0.23 | (0.22, 0.23) | <0.0001 |
| CRP (mg/l) | 0.45 | (0.41, 0.49) | 0.60 | (0.54, 0.65) | 0.48 | (0.43, 0.55) | 0.58 | (0.51, 0.67) | 0.46 | (0.40, 0.53) | 0.55 | (0.48, 0.63) | 0.06 |
| Triglyceride (mmol/l) | 0.80 | (0.78, 0.83) | 0.89 | (0.86, 0.92) | 0.83 | (0.80, 0.86) | 0.96 | (0.92, 1.00) | 0.73 | (0.70, 0.76) | 0.87 | (0.84, 0.91) | <0.0001 |
| HDL cholesterol (mmol/l) | 1.53 | (1.51, 1.56) | 1.46 | (1.44, 1.48) | 1.51 | (1.48, 1.54) | 1.44 | (1.41, 1.48) | 1.55 | (1.51, 1.59) | 1.48 | (1.45, 1.52) | <0.001 |
| LDL cholesterol (mmol/l) | 2.55 | (2.50, 2.60) | 2.52 | (2.48, 2.57) | 2.52 | (2.45, 2.58) | 2.56 | (2.49, 2.63) | 2.50 | (2.44, 2.58) | 2.56 | (2.49, 2.63) | 0.56 |
| Systolic BP (mmHg)\* | 105.2 | (104.3, 106.2) | 103.6 | (102.7, 104.6) | 105.1 | (103.9, 106.3) | 104.6 | (103.3, 105.9) | 102.5 | (101.2, 103.8) | 105.5 | (104.2, 106.7) | <0.001 |
| Diastolic BP (mmHg)\* | 62.9 | (62.1, 63.7) | 63.3 | (62.5, 64.1) | 62.6 | (61.6, 63.6) | 63.8 | (62.6, 64.9) | 62.7 | (61.6, 63.7) | 63.4 | (62.4, 64.5) | 0.32 |

\* Means are shown for untransformed variables

† N = 1083 for physical activity counts

‡ p-value for heterogeneity in association with ethnic group

Geometric means are shown for all other variables which are log transformed

Means presented by sex are adjusted for age (in quartiles), ethnic group, month of measurement height, and school (random effect)

Means presented by ethnic group are adjusted for sex, age (in quartiles), month of measurement height, and school (random effect)

Abbreviations: BP, blood pressure; Chol, cholesterol; Est, estimated; HOMA-IR, HOMA insulin resistance; PA, physical activity

Table 2: Associations between estimated VO2 max and risk markers for type 2 diabetes and cardiovascular disease with additional adjustment for fat mass index

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| Risk markers (N = 1445) | Adjustments | % difference/difference\* (95% CI) for a one IQR increase in estimated VO2 max | | p-value |
| Fat mass index (kg/m5) | Standard | -16.28 | (-18.44, -14.06) | <0.0001 |
| Insulin (mU/l) | Standard | -18.87 | (-22.27, -15.33) | <0.0001 |
| Standard + FMI | -7.69 | (-11.37, -3.85) | <0.001 |
| HOMA Insulin resistance | Standard | -18.80 | (-22.22, -15.23) | <0.0001 |
| Standard + FMI | -7.68 | (-11.40, -3.80) | <0.001 |
| HbA1c (%) | Standard | -0.56 | (-1.02, -0.10) | 0.02 |
| Standard + FMI | -0.34 | (-0.83, 0.15) | 0.17 |
| Glucose (mmol/l) | Standard | -0.99 | (-1.53, -0.45) | <0.001 |
| Standard + FMI | -0.67 | (-1.24, -0.09) | 0.02 |
| Urate (mmol/l) | Standard | -5.85 | (-7.56, -4.10) | <0.0001 |
| Standard + FMI | -2.19 | (-3.99, -0.35) | 0.02 |
| C-reactive protein (mg/l) | Standard | -40.85 | (-46.11, -35.06) | <0.0001 |
| Standard + FMI | -20.84 | (-27.52, -13.54) | <0.0001 |
| Triglyceride (mmol/l) | Standard | -9.79 | (-12.27, -7.24) | <0.0001 |
| Standard + FMI | -5.26 | (-7.93, -2.51) | <0.001 |
| HDL cholesterol (mmol/l) | Standard | 3.78 | (2.18, 5.41) | <0.0001 |
| Standard + FMI | 0.78 | (-0.81, 2.39) | 0.34 |
| LDL cholesterol (mmol/l) | Standard | -3.47 | (-5.34, -1.57) | <0.001 |
| Standard + FMI | -1.80 | (-3.80, 0.24) | 0.08 |
| Systolic blood pressure (mmHg)\* | Standard | -2.46 | (-3.22, -1.70) | <0.0001 |
| Standard + FMI | -1.69 | (-2.48, -0.89) | <0.0001 |
| Diastolic blood pressure (mmHg)\* | Standard | -3.17 | (-3.86, -2.49) | <0.0001 |
| Standard + FMI | -2.66 | (-3.38, -1.94) | <0.0001 |

\* Absolute differences are shown for untransformed variables

Percentage differences are shown for all other variables which are log transformed

Standard adjustment is for sex, age (in quartiles), ethnic group, month of measurement, height, and school (random effect)

Table 3: Ethnic differences in risk markers for type 2 diabetes and cardiovascular disease: effect of adjustment for estimated VO2 max

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| Risk markers (N = 1445) | Adjustments | % difference / difference\* (95% CI), p-value, % reduction in ethnic difference following adjustment for estimated VO2 max | | | |
| South Asian - white European | | | |
| Insulin (mU/l) | Standard | 38.63 | (26.78, 51.59) | <0.0001 |  |
| Standard + VO2 max | 34.52 | (23.32, 46.73) | <0.0001 | 10.6 |
| HOMA Insulin resistance | Standard | 39.34 | (27.41, 52.39) | <0.0001 |  |
| Standard + VO2 max | 35.19 | (23.90, 47.50) | <0.0001 | 10.5 |
| HbA1c (%) | Standard | 1.93 | (1.00, 2.86) | <0.0001 |  |
| Standard + VO2 max | 1.85 | (0.92, 2.78) | <0.0001 | 4.1 |
| Glucose (mmol/l) | Standard | 1.41 | (0.31, 2.51) | 0.01 |  |
| Standard + VO2 max | 1.25 | (0.16, 2.36) | 0.02 | 11.3 |
| Urate (mmol/l) | Standard | 1.76 | (-1.91, 5.57) | 0.35 |  |
| Standard + VO2 max | 0.86 | (-2.72, 4.58) | 0.64 | 51.1 |
| C-reactive protein (mg/l) | Standard | 20.80 | (0.21, 45.63) | 0.05 |  |
| Standard + VO2 max | 10.37 | (-8.09, 32.54) | 0.29 | 50.1 |
| Triglyceride (mmol/l) | Standard | 15.63 | (9.27, 22.36) | <0.0001 |  |
| Standard + VO2 max | 13.92 | (7.74, 20.44) | <0.0001 | 10.9 |
| HDL cholesterol (mmol/l) | Standard | -4.27 | (-7.20, -1.24) | 0.01 |  |
| Standard + VO2 max | -3.72 | (-6.66, -0.68) | 0.02 | 12.9 |
| LDL cholesterol (mmol/l) | Standard | 1.87 | (-1.92, 5.80) | 0.34 |  |
| Standard + VO2 max | 1.31 | (-2.45, 5.21) | 0.50 | 29.9 |
| Systolic blood pressure (mmHg)\* | Standard | -0.47 | (-2.02, 1.09) | 0.56 |  |
| Standard + VO2 max | -0.83 | (-2.37, 0.71) | 0.29 | -76.6 |
| Diastolic blood pressure (mmHg)\* | Standard | 1.19 | (-0.21, 2.59) | 0.10 |  |
| Standard + VO2 max | 0.76 | (-0.61, 2.13) | 0.28 | 36.1 |

\* Absolute differences are shown for untransformed variables

Percentage differences are shown for all other variables which are log transformed

Standard adjustment is for sex, age (in quartiles), ethnic group, month of measurement, height, and school (random effect)