Research Article

Reliable Prediction of Insulin Resistance by a School-Based Fitness Test in Middle-School Children

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Objectives. (1) Determine the predictive value of a school-based test of cardiovascular fitness (CVF) for insulin resistance (IR); (2) compare a “school-based” prediction of IR to a “laboratory-based” prediction, using various measures of fitness and body composition.

Methods. Middle school children (n = 82) performed the Progressive Aerobic Cardiovascular Endurance Run (PACER), a school-based CVF test, and underwent evaluation of maximal oxygen consumption treadmill testing (VO2 max), body composition (percent body fat and BMI z score), and IR (derived homeostasis model assessment index [HOMAIR]). Results. PACER showed a strong correlation with VO2 max/kg (rs = 0.83, P < .001) and with HOMAIR (rs = −0.60, P < .001). Multivariate regression analysis revealed that a school-based model (using PACER and BMI z score) predicted IR similar to a laboratory-based model (using VO2 max/kg of lean body mass and percent body fat).

Conclusions. The PACER is a valid school-based test of CVF, is predictive of IR, and has a similar relationship to IR when compared to complex laboratory-based testing. Simple school-based measures of childhood fitness (PACER) and fatness (BMI z score) could be used to identify childhood risk for IR and evaluate interventions.

1. Introduction

Insulin Resistance (IR) is an independent predictor for the development of hypertension, coronary heart disease, stroke, cancer, and type 2 diabetes, and greater insulin sensitivity is a protective factor against these clinical conditions [1]. IR and type 2 diabetes are increasing in childhood and adolescence [2].

In addition to obesity and genetic predisposition, poor cardiovascular fitness (CVF) is an important independent risk factor for IR [3]. Poor CVF increases a person’s risk for cardiovascular disease, hypertension, and type 2 diabetes [4–6], and improved CVF attenuates the morbidities associated with obesity. As a result, active yet overweight individuals can have lower risk for type 2 diabetes and cardiovascular disease than sedentary normal-weight individuals [7, 8]. Thus, an important public health goal is the improvement of CVF in the population in general, not just obese persons.

Unfortunately, the current gold standard for assessing CVF, maximal oxygen consumption treadmill or bike testing (VO2 max), requires specialized equipment, time, and expert supervision, and is therefore not practical for evaluation of large groups of children in school or community settings. To expand the ability to identify children with IR, a feasible school-based test of childhood CVF shown to be predictive of IR is needed.

FITNESSGRAM, developed in 1982 by The Cooper Institute, is a validated battery of tests that can be used to assess childhood physical fitness in schools. The Progressive Aerobic Cardiovascular Endurance Run (PACER) is a component of the FITNESSGRAM and consists of a multistage progressive 20-meter shuttle test (20MST). The PACER is a valid school-based test of CVF in pediatric populations [9]. Although laboratory-based fitness assessment (VO2 max) has been shown to be predictive of IR, it is not known if a school-based fitness assessment such as the PACER is sufficiently accurate to be predictive of IR.

The objectives of this study are (1) determine the predictive value of the PACER, alone and controlled for body composition, for IR; (2) compare “school-based” and

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“laboratory-based” assessments of childhood fitness and body composition in terms of predicting IR.

2. Materials and Methods

Children (n = 82) from two local middle schools participated in this study. Over a 12-month period (fall of 2006 to fall of 2007), each participant underwent testing at the University of Wisconsin Exercise Science Laboratory after an overnight fast. The procedures were approved by the Human Subjects Committee, and informed written consent was obtained before initiating the testing protocol. Testing included a physical examination, blood work, body composition assessment, and CVF assessment. Within 14 days of this testing, the participants performed the PACER at their schools. Subjects with overt type 2 diabetes were excluded.

Percent body fat (%BF) and lean body mass (LBM) were measured by dual-energy X-ray absorptiometry (DXA). Whole body scans were performed with the Norland XR-36 whole body bone densitometer (Norland Corporation, Ft. Atkinson, WI) following standard calibration, and tissue masses were analyzed with software version 3.7.4/2.1.0. Height was measured on a wall-mounted stadiometer to the nearest 0.5 cm. Weight was measured on a calibrated beam balance platform scale to the nearest 0.1 kg. “Obese” was defined as a BMI ≥ 95th percentile for age and gender.

The gold standard CVF assessment was maximal oxygen consumption (VO2 max), determined by open-circuit spirometry using a progressive treadmill walking protocol to volitional fatigue with a Medical Graphics CPX-D treadmill (St. Paul, MN). The speed of the treadmill was set initially per the subject’s comfort, starting at 0% grade and increasing 2% every minute. Requirements to strictly define whether subjects reached their maximal oxygen consumption by this protocol included at least 2 of the following three criteria: (1) maximal heart rate >200 beats per minute; (2) respiratory exchange ratio (VCO2/VO2) > 1; and (3) a plateau in oxygen consumption. All of the children included in the data analysis met at least 2 of the 3 criteria. VO2 max can be expressed as VO2 max per kg of total body weight (VO2 max/kg) or as VO2 max per kg of LBM (VO2 max/kgLBM). Since the best measurement of fitness is still debated, we analyze our data using both measures of CVF.

The measure of IR for this study was HOMAIR. HOMAIR is a validated equation that adjusts the fasting insulin and glucose to provide a reliable assessment of insulin resistance, particularly in obese children and adolescents [10, 11]. In children, the HOMA (r = 0.91) correlates closely with euglycemic clamp studies [11]. HOMAIR was calculated with the formula: fasting insulin (μIU/mL) × fasting glucose (mmol/L)/22.5. Fasting insulin concentration was determined with the chemiluminescent assay (ARUP Laboratories, Salt Lake City, UT), and glucose concentration was determined by an enzymatic method (Beckman Diagnostics, Fullerton, CA). According to ARUP Laboratories, at a mean insulin level of 14.8 μIU/mL, the standard deviation of the insulin assay is 1 and the coefficient of variation is 6.7%.

The PACER was conducted by school physical education teachers trained in administering the tests according to the FITNESSGRAM protocol [9]. The PACER is a multistage progressive 20-meter shuttle run. Subjects run back and forth along a 20-meter course, and each minute the pace required to run the 20 meters increases. The pace is determined by beeps from a pre-recorded tape or CD. The initial running speed is 8.5 km/hour, and the speed increases by 0.5 km/h every minute. The test is finished when the subject fails to complete the 20-meter run in the allotted time twice [9]. Because the number of laps per minute increases as the running speed increases, the PACER results are “scored” as number of laps successfully completed. The PACER results were also analyzed using running speed at the last completed one-minute stage and the estimated VO2 max/kg using the Leger et al. equation, which is the regression formula utilized by the FITNESSGRAM software [12]. The Leger equation is estimated VO2 max/kg = 31.025 + 3.238 (running speed in km/hr) – 3.248 (age in years) + 0.1536 (running speed × age) [12]. In the regression analysis, the PACER is expressed as number of laps completed.

Body composition, fitness, and insulin resistance measurements were summarized by standard descriptive statistics in terms of mean ± SD. The comparisons of all continuous measurements between subgroups (males versus females) were performed using a two-sample t-test or nonparametric Wilcoxon rank sum test, when data were not normally distributed. The univariate associations between fitness and body composition variables and HOMAIR were examined with the nonparametric Spearman rank correlation analysis. Multivariate linear regression analysis was performed for both a school-based model (with PACER as a measure of fitness and BMI z score as a measure of body composition) and a laboratory-based model (with VO2 max/kgLBM as a measure of fitness and %BF by DXA as a measure of body composition) to assess the independent contributions of fitness and body composition on insulin resistance. Both models were controlled for age and gender. The HOMAIR was log-transformed in the regression analyses to meet the assumption of normality. Variance inflation factors were calculated for each multivariate regression model in order to assess for collinearity. Standard model diagnostic tools (residual plots, etc.) were used to examine model assumptions. The predictive power of the regression models was assessed by performing cross-validation analysis. Cross-validation studies were conducted by randomly selecting sub-datasets which were used to fit the model and then validated on the remaining data. This was repeated 1000 times, and the root mean square error (RMSE) across all 1000 replications of the predicted versus observed values was calculated. A smaller RMSE indicates better prediction. All statistical analyses were performed with SAS software (version 9.1; SAS Institute, Cary, NC). All P values were 2-sided, and P values <.05 were considered statistically significant.
Table 1: Study subject characteristics (n = 82).

|                          | Total  | Males  | Females | P-value† |
|--------------------------|--------|--------|---------|----------|
| **Descriptive characteristics** |        |        |         |          |
| Number                   | 82     | 34     | 48      |          |
| Age (years)              | 13.0 ± 0.7 | 13.1 ± 0.7 | 12.9 ± 0.7 | .5028    |
| Height (cm)              | 162.8 ± 7.7 | 162.2 ± 8.2 | 163.3 ± 7.3 | .2958    |
| Weight (kg)              | 65.0 ± 19.7 | 67.5 ± 19.9 | 63.3 ± 19.5 | .3094    |
| Obese (BMI ≥ 95th percentile) | 26 (32%) | 17 (50%) | 9 (19%) | .0027*    |
| **Body composition assessments** |        |        |         |          |
| BMI                      | 24.4 ± 6.5 | 25.5 ± 6.6 | 23.5 ± 6.3 | .2193    |
| BMI z score              | 1.0 ± 1.1 | 1.2 ± 1.2 | 0.8 ± 1.1 | .1163    |
| Percent body fat         | 28.8 ± 8.6 | 28.3 ± 10.9 | 29.0 ± 6.7 | .7775    |
| Lean body mass (kg)      | 37.41 ± 8.86 | 39.47 ± 9.12 | 36.00 ± 8.48 | .0778    |
| **CVF assessments**      |        |        |         |          |
| VO2 max (mL/kg/min)      | 40.4 ± 9.4 | 43.4 ± 11.3 | 38.3 ± 7.1 | .0239*    |
| VO2 max/kgLBM (mL/kg/min)| 67.8 ± 8.2 | 71.0 ± 7.1 | 65.5 ± 8.2 | .0033*    |
| PACER (laps)             | 25.2 ± 15.1 | 27.7 ± 17.7 | 23.5 ± 12.8 | .2446    |
| PACER speed (km/h)       | 9.4 ± 0.8 | 9.5 ± 0.9 | 9.3 ± 0.7 | .2505    |
| Estimated VO2 max (mL/kg/min) [12] | 37.9 ± 4.5 | 38.4 ± 4.9 | 37.5 ± 4.2 | .3740    |
| **IR assessments**       |        |        |         |          |
| Fasting insulin (μIU/mL) | 15.0 ± 8.0 | 15.2 ± 9.5 | 14.9 ± 6.8 | .8419    |
| Fasting glucose (mg/dL)  | 92.8 ± 6.8 | 95.3 ± 6.5 | 91.2 ± 6.5 | .0071*    |
| HOMAIR                   | 3.5 ± 1.9 | 3.7 ± 2.3 | 3.4 ± 1.6 | .2766    |

†Comparison between males and females.
*P < .05.

3. Results

3.1. Subject Characteristics. Subject characteristics are presented in Table 1 as mean ± SD or number (percentage), and the P value reflects the comparison between males and females. Seventeen of the 34 boys (50%) and 9 of the 48 girls (19%) had a BMI greater than the 95th percentile (P = .003). Regardless of how CVF is expressed (relative to total body weight or lean body mass), boys had a higher CVF than girls. Girls had a higher fasting glucose level than boys (95.3 versus 93.8, P = .007), but no significant difference was seen between boys and girls in the fasting insulin or HOMAIR.

3.2. PACER Validation. The PACER correlated significantly and closely with VO2 max/kg (r = 0.83, 95% CI 0.74 to 0.93, P < .001; Figure 1). The PACER also correlated significantly with VO2 max/kgLBM (r = 0.55, 95% CI 0.37 to 0.68, P < .001). Since previous validation studies have compared the PACER to VO2 max, and not to VO2 max/kgLBM, we used VO2 max/kg for the additional correlations between the PACER and CVF. In the 26 obese children with a BMI greater than the 95th percentile, the PACER was highly correlated with VO2 max/kg (r = 0.74, 95% CI 0.49 to 0.87, P < .001).

The PACER test result has been expressed in multiple ways in the literature. VO2 max/kg correlated strongly with
Table 2: Univariate analysis for HOMAIR as a dependent variable (*n* = 82).  

| Independent variable                  | β (se)     | P value* | r_s† | RMSE** |
|---------------------------------------|------------|----------|------|--------|
| PACER (laps)                          | −0.07 (0.01) | <.0001*  | −0.60 | 1.68   |
| PACER-estimated VO₂ max               | −0.21 (0.04) | <.0001*  | −0.51 | 1.72   |
| VO₂ max                               | −0.13 (0.02) | <.0001*  | −0.70 | 1.56   |
| VO₂ max/kgLBM                         | −0.10 (0.02) | .0001*   | −0.47 | 1.79   |
| BMI                                   | 0.19 (0.02)  | <.0001*  | 0.65  | 1.54   |
| BMI z score                           | 1.03 (0.25)  | <.0001*  | 0.66  | 1.59   |
| Percent body fat                      | 0.14 (0.02)  | <.0001*  | 0.71  | 1.47   |

*P*-value for testing $H_0: \beta = 0$ versus $H_A: \beta \neq 0$.
†Spearman’s rank correlation coefficient.
**Cross-validation Root Mean Square Error (RMSE) on original scale for $m = 1000$ re-samples with sample size $n_1 = 50$ (model fitting) and $n_2 = 32$ (model validation).
*P < .05.

both the running speed (km/hr) at the last completed one-minute PACER stage ($r_s = 0.83$, 95% CI 0.74–0.88, $P < .001$) and the estimated VO₂ max/kg ($r_s = 0.83$, 95% CI 0.75–0.89, $P < .001$). Therefore, regardless of how the PACER is expressed (laps, running speed, or estimated VO₂ max), there was a strong correlation between the PACER and laboratory-based CVF assessment.

3.3. Univariate Analysis. Table 2 shows the results of the univariate analysis between HOMAIR and all the variables of interest. The PACER was significantly negatively correlated with HOMAIR ($r_s = −0.60$, 95% CI −0.75 to −0.44, $P < .001$; Figure 2). Laboratory-based fitness assessments and the body composition assessments were also significantly and strongly correlated with HOMAIR. An analysis using fasting insulin as the dependent variable revealed virtually identical results (data not shown), and therefore we focused additional analyses on HOMAIR only.

3.4. Multivariate Analysis. We used multivariate regression analysis to assess various measurements of fitness and body composition with HOMAIR as the dependent variable (Table 3). Both regression models were adjusted for sex and age. Model 1 utilized laboratory-based measurements of fitness (VO₂ max/kgLBM by maximal treadmill testing) and body composition (%BF by DXA) on insulin resistance (HOMAIR). Since more recent studies investigating the relationship between CVF and IR used VO₂ max/kgLBM, we use VO₂ max/kgLBM as the measure of fitness in the multivariate regression analysis. When we used VO₂ max/kg as the measure of fitness, there was no difference in the results (data not shown).

The laboratory-based method (Model 1) accounted for 57% of the variance in HOMAIR. Both VO₂ max/kgLBM ($P < .001$) and %BF ($P < .001$) were significant independent predictors of insulin resistance by HOMAIR. Model 2 utilized school-based measurements of fitness (PACER) and body composition (BMI z score) on the HOMAIR. This school-based method accounted for 45% of the variance in HOMAIR. The PACER and BMI z score were significant independent predictors of HOMAIR, with slope parameters of −0.012 ($P = .005$) and 0.201 ($P < .001$), respectively. In both multivariate regression models, the variance inflation factors were less than 2.5, which indicate that collinearity did not confound the results.

The ability of the models to predict IR can be compared with the RMSE (a smaller RMSE indicates greater predictive value). The laboratory-based assessment of fitness and body composition (Model 1) had a slightly lower RMSE of 1.42 compared to an RMSE of 1.49 for the school-based assessment (Model 2). Both models had greater predictive value than using BMI z score alone (RMSE of 1.59). Percent body fat, as determined by a DXA scan, had an RMSE of 1.47.

![Figure 2: Relationship between PACER (number of laps) and HOMAIR in a sample of middle school children (*n* = 82).](image-url)
Table 3: Regression coefficients ($\beta$), $P$ value, and coefficient of determination ($R^2$) for the association of log (HOMA\textsubscript{IR}) with CVF and body composition in 2 multiple regression models, after adjusting for sex and age.

| Variables          | $\beta$ (SE) | $P$ value* | $R^2$ | RMSE** |
|--------------------|--------------|------------|-------|--------|
| Model 1            |              |            |       |        |
| VO\textsubscript{2} max/kgLBM | $-0.027$ (0.006) | <.001* | 0.57  | 1.42   |
| %BF                | 0.035 (0.005) | <.001*     |       |        |
| Model 2            |              |            |       |        |
| PACER (laps)       | $-0.012$ (0.004) | .0050* |       | 0.45   |
| BMI z score        | 0.201 (0.057) | .0007*     |       | 1.49   |

* $P$-value for testing $H_0: \beta = 0$ versus $H_A: \beta \neq 0$.
** Cross-validation Root Mean Square Error (RMSE) on original scale for $m = 1000$ re-samples with sample size $n_1 = 50$ (model fitting) and $n_2 = 32$ (model validation).

* $P < .05$.

4. Discussion

Given that CVF is an important predictor of cardiovascular and metabolic health problems independent of weight, a method to accurately and feasibly assess childhood fitness levels in addition to BMI in larger school-based or population-based studies is needed. While a school- or field-based test by definition is easier to use in community settings, the value of school-based assessments of fitness is greatly enhanced if they have been shown to have predictive value for other health outcomes, such as IR. This study shows that the PACER is a valid school-based test of CVF and is predictive of IR, independent of body composition. Moreover, the combination of school-based measurements of both childhood fitness (PACER) and fatness (BMI z score) predicts IR to a degree similar to laboratory-based measurements of fitness (VO\textsubscript{2} max/kgLBM by maximal treadmill testing) and fatness (%BF by DXA). Thus, the PACER can be used to translate findings of laboratory-based studies to large population-based settings to identify children with low fitness levels and/or increased risk for IR. To our knowledge, this study is the first to identify a correlation between a school-based fitness test in children and measurements of IR.

This study showed that CVF, measured either by a laboratory-based or a school-based assessment, is an important predictor of insulin resistance, independent of fatness. Some previous studies have shown that laboratory-based assessment of fitness in children has a highly significant correlation with IR after controlling for body composition [3, 13–17], while others have not [18–20]. A previous study by our group found that VO\textsubscript{2} max/kgLBM (by maximal treadmill testing) was significantly correlated with HOMA\textsubscript{IR} ($r = -0.42$) [3]. In the present study, the laboratory-based assessment of fitness was a significant independent predictor of IR. The new contribution of this study is the correlation of school-based fitness tests in children with measures of IR. The PACER correlated very closely and significantly with HOMA\textsubscript{IR} ($r_s = -0.60$), and after controlling for body composition, the PACER was still independently predictive of IR (Model 2, Table 3).

We compared a laboratory-based method to a school-based method of predicting insulin resistance using multivariate regression analysis. Model 1 utilized laboratory-based assessments of fitness and body composition while Model 2 included school-based assessments of fitness and body composition. The laboratory-based model had an RMSE of 1.42 compared to 1.49 in the school-based model (a smaller RMSE indicates better prediction). It can be reasonably argued that this small difference in predictive value is counterbalanced by the ability to administer the school-based tests to a much larger population of children.

Another valid question is whether assessment of fitness adds substantial value to simply measuring and tracking obesity. The RMSE for BMI z score is 1.59 (Table 2), compared to 1.49 for Model 2. Additionally, a “weight-only” model would not capture the many benefits of fitness testing and fitness promotion that extend beyond the prediction of IR [4, 21, 22].

It is interesting to note that the scatterplot comparing PACER results to HOMA\textsubscript{IR} (Figure 2) shows high variability in IR at low fitness levels, and progressively less variability as the fitness levels improve. This could indicate that the increased IR variability at low fitness levels reflects individual genetic predispositions for obesity-induced IR, and that higher fitness levels can improve insulin sensitivity in all persons. Our group has previously shown that increasing fitness levels in middle school children can lead to improvements in fasting insulin levels [23], and the effect of improved fitness on IR and the risk for type 2 diabetes has been a consistent finding in adults [4, 8]. In this cross-sectional study, the effect of improved fitness levels on IR cannot be determined.

This study has limitations. The HOMA\textsubscript{IR} is a useful approximation of insulin resistance measured by euglycemic clamp studies, but does have a degree of error compared with that gold standard. Correlation analysis can overestimate the relationship between variables and therefore has intrinsic limitations. Additionally, puberty can increase insulin resistance and may be a confounding variable in the relationship between fitness testing and IR. Finally, our multivariate models did not include family history of type 2 diabetes, which would likely improve the prediction of IR by accounting for a child’s genetic predisposition for IR. Future studies hope to incorporate family history in order to improve the predictive value of school-based assessments for IR.

The relative contributions of fitness and fatness to the development of IR continue to be debated, and this study was not designed to address that issue. Regardless of their relative importance, it is clear that improvements in both...
fitness and fatness will yield important health consequences for increasingly obese and sedentary children [22]. Thus, it is essential to develop interventions to reduce insulin resistance that are applicable to the “real world” of children. These interventions will need a field-based metric that can be easily and broadly implemented to translate exercise laboratory study findings to the public health setting.

5. Conclusions

With rates of both obesity and poor fitness increasing in children, development of IR will put increasing numbers of children at risk of developing early metabolic and cardiovascular disease. Precise measurement of IR involves blood studies of insulin levels required to maintain euglycemia after glucose administration (euglycemic clamp study), an impractical procedure in school or community settings. Laboratory-based prediction of IR, through measurement of CVF (by maximal treadmill testing) and body composition (by DXA), is also resource intensive and impractical in nonlaboratory settings. This study shows that (1) the PACER is a valid school-based test of CVF and is predictive of IR; (2) simple measures of both childhood fitness (PACER) and body composition (BMI z score) together accurately predict IR to a greater degree than BMI alone, approximating the degree of prediction from logistically complex and expensive laboratory tests of fitness and body composition. Thus, the PACER offers a valid school-based test of CVF that is predictive of IR and allows findings from exercise laboratory studies to be translated to a school or community setting.

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