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1 **Title:** Novel measures of inflammation and insulin resistance are related to obesity and
2 fitness in a diverse sample of 11-14 year-olds: The HEALTHY Study

3

4 **Running title:** Fitness, BMI, GlycA and LP-IR in US youth

5

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18

19 **Conflict of interest:** JDO is an employee of LabCorp, a commercial supplier of NMR-based
20 diagnostic testing. JBB has been a consultant to LipoScience and Quest Diagnostics under a
21 service agreement with his employer. This provides no direct financial benefit to him.

22

23 **Word count = 3624 words**

24 **ABSTRACT**

25 **Background:** GlycA is a novel serum marker of systemic inflammation. There is no
26 information on GlycA in pediatric populations, how it differs by gender or its association
27 with body mass index (BMI) or fitness. LP-IR is a serum measure of insulin resistance which
28 is related to changes in BMI group in adolescents, but its relationship with fitness is
29 unknown. The current study examined the independent associations between fitness and BMI
30 with GlycA and LP-IR among US adolescents.

31 **Methods:** Participants were 1664 US adolescents from the HEALTHY study with complete
32 6th and 8th grade BMI, fitness and blood data. GlycA and LP-IR were measured by NMR
33 spectroscopy. Three BMI groups and three fitness groups were created. Linear mixed models
34 examined associations between GlycA, LP-IR, fitness and BMI.

35 **Results:** LP-IR decreased between 6th and 8th grade. GlycA increased among girls but
36 decreased among boys. At 8th grade, median GlycA values were 27 (7.6%) $\mu\text{mol/L}$ higher
37 (381 versus 354) for girls than boys. Median GlycA 6th grade values were 9% higher in obese
38 girls than healthy weight girls. Overall there was strong evidence ($p < 0.001$) that GlycA was
39 higher in higher BMI groups. Fitness was negatively associated with GlycA ($r = -0.37$ and -
40 0.35) and LP-IR ($r = -0.34$ and -0.18) at the 6th and 8th grade assessments. As BMI category
41 increased and fitness category decreased, GlycA and LP-IR levels increased. Lowest GlycA
42 was found in the low BMI / high fitness group.

43 **Conclusions:** GlycA was associated with BMI and fitness among in US adolescents. These
44 findings suggest that there are independent effects for BMI and fitness group with both
45 GlycA and LP-IR. Future studies should validate the role of GlycA and LP-IR to evaluate the
46 effects of interventions to modify obesity and fitness in order to improve systemic
47 inflammation and insulin resistance.

48

49 **INTRODUCTION**

50 Obesity and low levels of cardio-respiratory fitness (fitness) are associated with the
51 development of cardiovascular disease (CVD) and type 2 diabetes mellitus (T2DM)¹⁻³. The
52 physiological mechanisms that contribute to these well established associations are
53 dyslipidemia, insulin resistance and inflammation all of which are associated with obesity
54 and low fitness⁴⁻⁷. A number of studies have demonstrated the adverse effects of increased
55 body fat and low fitness in childhood on future health, with childhood obesity being strongly
56 associated with its persistence into adulthood and fitness tracking through childhood into the
57 adulthood⁸⁻¹⁰. A number of adult studies have suggested that there are independent
58 associations between fitness and body mass index in relation to the risk of cardiovascular
59 disease, type 2 diabetes and all-cause mortality^{1,2}. C-reactive protein (CRP), a marker of
60 systemic inflammation and predictor of cardiovascular risk, has been shown to be closely
61 associated with obesity among children and adults¹¹. Thus, there is a need to examine how
62 various relevant metabolic markers, such as dyslipidemia, insulin resistance and
63 inflammation are associated with body mass and fitness among children. A key issue is
64 whether there are associations between markers of dyslipidemia, insulin resistance and
65 inflammation with body mass and fitness in pediatric populations. If associations exists these
66 markers could be considered potential targets for the reduction of future cardiovascular
67 disease risk.

68

69 GlycA, a novel composite measure of systemic inflammation, and the lipoprotein insulin
70 resistance index (LP-IR) are promising new clinical biomarkers measured by nuclear
71 magnetic resonance (NMR) spectroscopy^{12,13}. Both are obtained efficiently and
72 inexpensively from the same *NMR LipoProfile* test spectra acquired on automated clinical
73 NMR analyzers to quantify lipoprotein particles for use in CVD risk management¹⁴. Several

74 recent reports have appeared relating these new markers to CVD and T2DM risk in adults ¹⁵,
75 ¹⁶, but no comparable data are available in children and adolescents.

76

77 Clinical interest in GlycA stems partly from its composite nature, reflecting the integrated
78 concentrations and glycosylation states of several of the most abundant acute-phase proteins
79 in serum, and its much lower intra-individual biological variability compared to CRP and
80 other markers of inflammation ¹². As a result, GlycA may provide a more stable measure of
81 low-grade systemic inflammation that responds more consistently to diverse inflammatory
82 stimuli than individual acute-phase reactants such as CRP. In several adult studies, GlycA
83 and CRP were found to have independent associations with incident CVD of comparable
84 strength, with some evidence of complementarity suggesting a possible adjunctive clinical
85 use ¹⁷⁻²⁰. Similar observations were made relating GlycA to prediction of future T2DM in the
86 Women's Health Study and Dutch PREVEND study ^{15, 16}.

87

88 The insulin resistance biomarker, LP-IR, reflects the lipoprotein derangements of insulin
89 resistance and is derived by combining 6 NMR measures of very-low-density lipoprotein
90 (VLDL), low-density lipoprotein (LDL) and high-density lipoprotein (HDL) particle size and
91 subclass concentration ¹³. Each of these subclass and size parameters has been shown
92 individually to be associated with incident T2DM in the Women's Health Study ²¹, and the
93 composite LP-IR score exhibited robust diabetes prediction in a large multi-ethnic cohort of
94 men and women ²².

95

96 We previously obtained *NMR LipoProfile* spectra from a substantial subset of 6th and 8th
97 grade participants in the ethnically diverse HEALTHY trial, initially to characterize the
98 differences between NMR-measured lipoprotein particle numbers and traditional lipid

99 measures in a large pediatric population ²³. In a subsequent report, changes in relative weight
100 group from 6th to 8th grade were related to lipoprotein particle changes associated with risk of
101 CVD and T2DM, as well as alterations in insulin resistance assessed by LP-IR and insulin
102 and glucose measurements ²⁴. In this paper, we take advantage of the ability, with newly
103 available software, to extract GlycA values from the same *NMR LipoProfile* dataset to
104 address the absence of information in youth regarding relations of body weight and fitness
105 with GlycA and LP-IR. The aims of this study in an ethnically diverse sample of children
106 were to: a) report on levels of GlycA and the change in GlycA as children move from 6th to
107 8th grade; b) examine whether BMI group is associated with GlycA in these children; c)
108 determine if fitness was associated with GlycA, LP-IR and traditional lipid panel variables;
109 and d) examine whether fitness and BMI are independently related to GlycA and/or LP-IR.

110

111 **METHODS**

112 The analyses reported in this paper used information from stored blood from the HEALTHY
113 Study, a National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK) cluster
114 randomized controlled trial that aimed to reduce the prevalence of risk factors for type 2
115 diabetes mellitus among middle school children (6th - 8th grade) ^{25, 26}. Details of the study
116 design and results have been reported in a number of publications ^{25, 26}. Briefly, the study was
117 conducted in 42 middle schools across the US. In order to participate in the study, schools
118 had to have at least 50% of students eligible for free or reduced-price lunch or belonging to
119 an ethnic minority group at increased risk of type 2 diabetes. The intervention had several
120 components including changes to the physical education and cafeteria programs as well as
121 health education and a school wide social marketing campaign ^{25, 27-32}. The study was
122 approved by the Institutional Review Boards at each field center and written parental consent
123 and childhood assent was obtained from all participants ³³. The sample for this study is

124 limited to the 1664 participants who provided parental consent/child assent for ancillary
125 analyses of stored blood, and for whom complete data were available.

126

127 *Procedures*

128 All measures were assessed at baseline (beginning of 6th grade) and follow-up (end of 8th
129 grade). Pubertal status was self-reported using the Pubertal Development Scale³⁴ and
130 converted to pubertal stage groups consistent with the five pubertal stages outlined by Tanner
131³⁵. Household education were determined from parental report and gender and race/ethnicity
132 was self-reported. Height and body mass were measured without shoes using the Prospective
133 Enterprises PE-AIM-101 stadiometer and the SECA Corporation Alpha 882 electronic scale.
134 Body Mass Index (kg/m^2) was calculated and converted to an age and gender specific BMI
135 percentile using CDC 2000 criteria³⁶. For descriptive purposes participants with a BMI <85th
136 percentile were classified as healthy weight; while BMI $\geq 85^{\text{th}}$ percentile but <95th percentile
137 were classified as overweight and those with BMI $\geq 95^{\text{th}}$ percentile were classified as obese.

138

139 Cardiorespiratory fitness was assessed using the 20-meter shuttle test (20-MST)^{37, 38} during a
140 Physical Education class. The test required students to run back and forth between two lines
141 set 20 meters apart. The running pace was determined by audio signals emitted from a pre-
142 recorded CD. The test started at 8.5 km/hr and increased by 0.5 km/hr with each subsequent
143 level. The test was completed when the participant was not able to complete the distance at
144 the stipulated pace on two laps.

145

146 Fasting blood samples were collected from all participants. Standard lipid profiles including
147 HDL-C were measured by CDC-standardized direct assay at the University of Washington³⁹.
148 LDL-C was calculated using the Friedewald equation⁴⁰. Insulin was measured by a two-site

149 immuno-enzymometric assay⁴¹. Fasting insulin (performed using a Tosoh 1800 auto-
150 analyzer) and glucose (performed on a Roche P module auto-analyzer by the hexokinase
151 method) were used to calculate the homeostatic model assessment of insulin resistance
152 (HOMA-IR) according to the formula: $\text{Glucose} \times \text{Insulin} / [\mu\text{U/L}] \times 22.5$ ⁴². Lipoprotein particle
153 profiles were measured by NMR spectroscopy with the LipoProfile-3 algorithm at
154 LipoScience, Inc (Raleigh, NC) on frozen EDTA plasma specimens and LP-IR and GlycA
155 were derived using previously published procedures^{12, 13}.

156

157 *Statistical analysis*

158 Overall, 2367 participants in HEALTHY had samples analyzed by LipoScience. Of these,
159 703 were excluded from the present analysis, most due to imprecise classification of their
160 race or the lack of either a 6th or 8th grade fitness test. Descriptive statistics, including means,
161 standard deviations, and percentages were calculated for those included and excluded from
162 the analysis. Differences between those included and excluded were tested using a
163 generalized linear mixed model which took into account the sources of variability within and
164 between schools. A similar approach was undertaken for baseline characteristics of the
165 analysis sample to examine gender differences.

166

167 Measures in 6th grade, 8th grade and change between 6th and 8th grade are summarized using
168 mean and 95% confidence interval for normally distributed measures (number of laps) and
169 medians and interquartile range (IQR) for parameters not fitting a normal distribution (lipid
170 measures, insulin resistance measures and GlycA) by gender. Median and IQR are also
171 presented for GlycA based on BMI category for normal weight (BMI < 85th percentile),
172 overweight (BMI 85th to 94th percentile) and obese (BMI \geq 95th percentile) and percentiles
173 were calculated by gender and grade for GlycA.

174

175 Generalized linear mixed models were constructed to examine the association between
176 GlycA and BMI category for 6th grade and 8th grade (adjusting for GlycA in 6th grade),
177 separately for girls and boys taking into account the sources of variability within and between
178 schools. These models were adjusted for race/ethnicity, highest household education,
179 intervention group with the 6th grade models also adjusted for 6th grade Tanner stage and the
180 8th grade models adjusted for 8th grade Tanner stage and 6th grade GlycA value. Spearman
181 correlations between GlycA and lipid measures, insulin resistance, BMI percentile, and
182 fitness (number of laps) were calculated for 6th grade and 8th grade with both genders
183 combined since there were no appreciable differences between the genders. The analysis of
184 the 8th grade data, which is adjusted for 6th grade values, allows us to account for the
185 longitudinal nature of the data.

186

187 Level of fitness was classified into quartiles by grade and gender and then further classified
188 as low, medium or high fitness levels. Low fitness level was defined to be those grouped into
189 the first quartile: 6th grade girls 0-11 laps, 6th grade boys 0-12 laps, 8th grade girls 1-12 laps,
190 8th grade boys 1-17 laps. Medium fitness level was defined to be those grouped into the
191 second or third quartile: 6th grade girls 12-23 laps, 6th grade boys 13-30 laps, 8th grade girls
192 13-26 laps, 8th grade boys 18-44 laps. Finally, high fitness was defined as those grouped into
193 the fourth quartile: 6th grade girls 24-57 laps, 6th grade boys 31-75 laps, 8th grade girls 27-79
194 laps, 8th grade boys 45-103 laps. Adjusted means and standard errors were then computed for
195 GlycA, LP-IR, LDL-P and non-HDL-C within BMI classification (normal, overweight,
196 obese) and fitness level. Levels of GlycA, LP-IR, LDL-P and non-HDL-C were then
197 categorized within the three BMI and three fitness groups to create nine subgroups and these

198 groups were tabulated by gender at both 6th and 8th grade. To further facilitate understanding,
199 the GlycA and LP-IR values in these subgroups were then presented graphically.

200

201 All p-values reported within this paper represent findings associated with secondary
202 outcomes from a large cluster randomized controlled trial and these hypotheses were not pre-
203 specified in the trial design. As such, p-values are provided to help facilitate the interpretation
204 of the data only. SAS 9.3 statistical software (SAS Institute, Cary, NC) was used for
205 analyses.

206

207 **RESULTS**

208 Descriptive statistics for 6th grade participants are presented overall and by gender in Table 1.
209 The data in Table 1 present strong evidence that boys recorded a higher number of shuttle run
210 laps than the girls (23.1 versus 18.5, $p < 0.001$) and some evidence that boys had a higher BMI
211 percentile (75.4 versus 72.4, $p = 0.0264$) than the girls.

212

213 Supplemental Table A provides descriptive information on the participants included and
214 excluded from the analyses. These data provide some evidence ($p = 0.0043$) that there was a
215 difference in the ethnicity of the included versus excluded participants with higher
216 proportions of Hispanic (62.7% vs. 40.4%) and White participants (20.7% vs. 12.2%) in the
217 included sample.

218

219 Table 2 provides descriptive information (means and 95% CI or median and inter-quartile
220 range) for fitness, lipids, insulin resistance and GlycA variables at 6th and 8th grade along
221 with the temporal changes (8th - 6th grade value) stratified by gender. The table shows that
222 among both girls and boys non-HDL-C, LDL-C and LP-IR decreased between 6th and 8th

223 grade while HOMA-IR increased. GlycA increased by 16 $\mu\text{mol/L}$ (3.8% based on medians)
224 among girls but decreased by 6 $\mu\text{mol/L}$ (3.1% based on medians) among the boys. At 8th
225 grade the median GlycA values were 27 (7.6%) $\mu\text{mol/L}$ higher (381 versus 354) for girls than
226 boys. Supplementary Table B provides percentiles of GlycA by grade and gender.

227

228 Supplementary Table C provides the medians and inter-quartile ranges for GlycA by BMI
229 group stratified by gender and grade level. The median GlycA value for 6th grade obese girls
230 was 417 $\mu\text{mol/L}$ compared to 351 for healthy weight girls, and as such, GlycA levels are 19%
231 higher in obese girls than healthy weight girls. The data in the table provide strong evidence
232 ($p < 0.001$) that in all sub-groups GlycA is higher in higher BMI groups. Supplementary Table
233 D provides Spearman correlations between all of the variables. The number of shuttle run
234 laps is negatively associated with GlycA ($r = -0.37$ and $r = -0.35$) and LP-IR ($r = -0.34$ and r
235 $= -0.18$) for the 6th and 8th grade associations respectively.

236

237 Figure 1 provides a graphical presentation of levels of adjusted GlycA by BMI and fitness
238 categories stratified by gender and 6th or 8th grade. The figure demonstrates that across all
239 four sub-groups there is evidence that as BMI category increases and fitness category
240 decreases levels of GlycA increase. Furthermore, in all four sub-groups the lowest levels of
241 GlycA are in the low BMI / high fitness group with the highest levels in the high BMI / low
242 fitness group. Comparable patterns are also evident for LP-IR levels, which are graphically
243 presented in Figure 2. The data used to create Figures 1 and 2 are available in Supplementary
244 Table E.

245

246

247

248 **DISCUSSION**

249 This study is the first to provide descriptive information on levels of GlycA, a new NMR-
250 derived marker of systemic inflammation, in adolescents. The data presented in this study
251 have shown, in an ethnically diverse sample of adolescents, that GlycA is associated with
252 BMI group with levels of GlycA higher across BMI groups among both boys and girls in the
253 6th and 8th grades. These data also show that GlycA is inversely correlated with shuttle run
254 laps, a surrogate measure of cardiorespiratory fitness at both 6th and 8th grade. Furthermore,
255 when levels of GlycA were analyzed by the three BMI and three fitness groups there is
256 evidence of a relationship between fitness, BMI group and GlycA with the highest levels of
257 GlycA among obese children in the lowest fitness category. The relationship between fitness,
258 BMI and GlycA was similar for both boys and girls at 6th and 8th grade. A key finding of this
259 study is therefore that GlycA, a new measure of systemic inflammation, is associated with
260 both body mass and fitness in a pediatric population. As such, further examination of GlycA
261 in pediatric populations is warranted to identify associations with future disease risk and the
262 response to changes in fitness and body mass among children.

263

264 There was evidence that LP-IR was negatively correlated with the number of shuttle laps run
265 at both 6th and 8th grade. We have previously reported that LP-IR is associated with BMI
266 group and change in BMI group among boys and girls in the same sample²⁴. Thus, in this
267 paper we have extended these findings by showing that there are independent effects of
268 fitness and BMI group on LP-IR with the lowest levels of LP-IR among the low BMI / high
269 fitness group and the highest levels among the high BMI / low fitness group. These patterns
270 were comparable among boys and girls at 6th and 8th grade. These finding therefore suggest
271 that facilitating increased fitness and lower BMI is likely to be important for achieving lower
272 levels of LP-IR among adolescents, which may reduce overall CVD risk.

273

274 The association reported in this paper between fitness, BMI and GlycA is broadly consistent
275 with previous research, which has shown that fitness is associated with CRP, another measure
276 of systemic inflammation, after accounting for body mass. For example, in a cross-sectional
277 analysis, the fitness levels of young adults were inversely associated with CRP and this
278 association was maintained even after adjustment for body mass index⁴³. Similarly, among
279 adults with type 2 diabetes a change in fitness over 12-months was associated with change in
280 CRP and this association was independent of BMI change⁴⁴. In adolescents, fitness and
281 fatness (assessed via skinfolds) have been independently associated with systemic
282 inflammation, as measured by CRP⁴⁵. There is some evidence that GlycA may be a better
283 marker of shorter-term CVD risk (events occurring within ~6 years) than longer-term risk
284 which may mean that GlycA can serve as an early marker of adult disease risk but it is not
285 currently clear if this is the case for adolescents¹⁷. However, GlycA has a lower level of
286 intra-individual variability than CRP which might make it attractive as a CVD risk marker
287 across time. As CRP was not available in the HEALTHY study, it is not possible to directly
288 compare the associations between CRP and GlycA, fitness and BMI group in this dataset. As
289 such, we are unable to state that one biomarker may be preferable to another, but such an
290 assessment in a future study is warranted. Moreover, it would be useful to assess whether
291 intervention studies that target increased fitness and reduced body mass might yield
292 improvements in GlycA.

293

294 Higher levels of cardio-respiratory fitness have been associated with a reduced risk of
295 developing cardiovascular disease and type 2 diabetes among adults^{1,2}. For example, recent
296 analyses of the CARDIA study have shown that fitness in young adulthood was associated
297 with all-cause mortality approximately 27 years later with each additional minute of exercise

298 test duration associated with a 15% lower hazard of death³. Interestingly the study showed
299 that CRP was much higher in the low fitness group, thereby highlighting the important link
300 between fitness and inflammation. The current study extends this work to show how fitness
301 and obesity are independently associated with GlycA, a novel measure of systemic
302 inflammation in a pediatric population.

303

304 GlycA levels were 7% higher for girls than boys at 8th grade but were comparable between
305 the genders in 6th grade. This finding is broadly consistent with the adult literature in which
306 CRP levels are higher among women than men^{46, 47} despite the lack of a difference in
307 absolute or relative risk of cardiovascular events when compared to men⁴⁸. We presume that
308 this gender difference may represent a hormonal influence upon GlycA, which is independent
309 of its role as a CVD risk marker. The emergence of higher levels of GlycA in 8th grade girls,
310 therefore, might be expected with the progression of puberty, generally occurs between 6th
311 and 8th grade in girls. For example, in this study the proportion of girls classified as Tanner
312 Stage 4 or 5 at 6th grade was 23.5% but by 8th grade this had increased to 94.2%. As previous
313 research has shown that the advancement of puberty is associated with changes in insulin
314 sensitivity⁴⁹ it may also be the case that pubertal hormones exert more direct effects on
315 systemic inflammation, and this influence may persist beyond puberty. Further research into
316 hormonal influences on these markers of inflammation is therefore warranted.

317

318 ***Strengths / limitations***

319 The major strength of this study is the provision of a detailed analysis of GlycA and LP-IR in
320 a large, ethnically diverse sample of young people progressing from 6th to 8th grade. No
321 comparable data exist and as such these data make a unique contribution to the field.

322 However, the study has several limitations to be considered. Firstly, a field based measure of

323 fitness (shuttle run laps) was used in the analyses. Although this measure has been shown to
324 be closely correlated with directly measured oxygen uptake it is less precise than laboratory
325 based methods⁵⁰⁻⁵². Secondly, we used a self-report measure for assignment of pubertal
326 status. Although this measure has been validated, it is not generally considered to be as
327 reliable as clinician assessment of Tanner staging³⁴. Thirdly, the analyses in this paper have
328 focused on GlycA, a new measure of systemic inflammation, but as CRP is not available in
329 this dataset it is not possible to assess how associations may compare to the more widely used
330 marker of systematic inflammation. Fourth, it is important to recognize that the analyses
331 reported in this paper were conducted on a sub-set of participants who provided consent for
332 ancillary analyses and complete data for all variables were available. In particular, the
333 observed difference in the ethnicity of the included versus the excluded participants is a
334 potential weakness. Fifth, it is important to note that consistent with other studies^{1,2} that
335 have examined the association between fitness, body mass and health outcomes we have only
336 assessed and analyzed cardio-vascular fitness and have no data to comment on broader
337 aspects of fitness such as strength. Finally, this is a post hoc analysis of secondary endpoints
338 and our analyses have not been corrected for the multiple comparisons being made, and
339 should therefore be regarded as exploratory.

340

341 **CONCLUSIONS**

342 GlycA, a new measure of systemic inflammation, was associated with BMI and fitness
343 among an ethnically diverse sample of adolescents in the US. Analyses also provided
344 evidence of independent effects for BMI and fitness groups when related to both GlycA and
345 LP-IR, a multivariate insulin resistance score. These findings suggest that reducing body
346 mass and increasing fitness may reduce both systemic inflammation (GlycA) and insulin
347 resistance (LP-IR). Further examination of how body mass, fitness and changes in both of

348 these health indicators is associated with GlycA in pediatric populations is therefore
349 warranted.

350

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356 at the middle schools and school districts that participated in the HEALTHY study.

357 Please see Appendix 1 for a full list of study group members and affiliations. HEALTHY
358 intervention materials are available for download at <http://www.healthystudy.org/>.

359

360 **Conflict of interest:** JDO is an employee of LabCorp, a commercial supplier of NMR-based
361 diagnostic testing. JBB has been a consultant to LipoScience and Quest Diagnostics under a
362 service agreement with his employer. This provides no direct financial benefit to him.

363

364 **REFERENCES**

- 365 1. Lee CD, Blair SN, Jackson AS. Cardiorespiratory fitness, body composition, and all-
366 cause and cardiovascular disease mortality in men. *Am. J. Clin. Nutr* 1999; **69**: 373-
367 380.
- 368 2. Sui X, Hooker SP, Lee IM, Church TS, Colabianchi N, Lee CD *et al.* A Prospective
369 Study of Cardiorespiratory Fitness and Risk of Type 2 Diabetes in Women. *Diabetes*
370 *Care* 2007: 550-5.
371
- 372 3. Shah RV, Murthy VL, Colangelo LA, Reis J, Venkatesh BA, Sharma R *et al.*
373 Association of Fitness in Young Adulthood With Survival and Cardiovascular Risk:
374 The Coronary Artery Risk Development in Young Adults (CARDIA) Study. *JAMA*
375 *internal medicine* 2016; **176**(1): 87-95.
376
- 377 4. Klop B, Elte JW, Cabezas MC. Dyslipidemia in obesity: mechanisms and potential
378 targets. *Nutrients* 2013; **5**(4): 1218-40.
379

- 380
381 5. Lavie CJ, Arena R, Swift DL, Johannsen NM, Sui X, Lee DC *et al.* Exercise and the
382 cardiovascular system: clinical science and cardiovascular outcomes. *Circ Res* 2015;
383 **117**(2): 207-19.
- 384
385 6. Vasconcellos F, Seabra A, Katzmarzyk PT, Kraemer-Aguiar LG, Bouskela E,
386 Farinatti P. Physical activity in overweight and obese adolescents: systematic review
387 of the effects on physical fitness components and cardiovascular risk factors. *Sports*
388 *Med* 2014; **44**(8): 1139-52.
- 389
390 7. Tam CS, Clement K, Baur LA, Tordjman J. Obesity and low-grade inflammation: a
391 paediatric perspective. *Obes Rev* 2010; **11**(2): 118-26.
- 392
393 8. Whitaker RC, Wright JA, Pepe MS, Seidel KD. Predicting obesity in young adulthood
394 from childhood and parental obesity. *N Engl J Med* 1997; **337**: 869-873.
- 395
396 9. Twisk JWR, Kemper HCG, Mechelen WV. Tracking of activity and fitness and the
397 relationship with cardiovascular disease risk factors. *Medicine & Science in Sports &*
398 *Exercise* 2000; **32**(8): 1455-1461.
- 399
400 10. McMurray RG, Harrell JS, Bangdiwala SI, Hu J. Tracking of physical activity and
401 aerobic power from childhood through adolescence. *Med Sci Sports Exerc* 2003;
402 **35**(11): 1914-22.
- 403
404 11. Choi J, Joseph L, Pilote L. Obesity and C-reactive protein in various populations: a
405 systematic review and meta-analysis. *Obes Rev* 2013; **14**(3): 232-44.
- 406
407 12. Otvos JD, Shalaurova I, Wolak-Dinsmore J, Connelly MA, Mackey RH, Stein JH *et*
408 *al.* GlycA: A novel nuclear magnetic resonance biomarker of systemic inflammation.
409 *Clinical Chem* 2015; **61**(5): 714-723.
- 410
411 13. Shalaurova I, Connelly MA, Garvey WT, Otvos JD. Lipoprotein insulin resistance
412 index: a lipoprotein particle-derived measure of insulin resistance. *Metabolic*
413 *Syndrome and Related Disorders* 2014; **2014**(12): 422-429.
- 414
415 14. Jeyarajah EJ, Cromwell WC, Otvos JD. Lipoprotein particle analysis by nuclear
416 magnetic resonance spectroscopy. *Clin Lab Med* 2006; **26**(4): 847-70.
- 417
418 15. Akinkuolie AO, Pradhan AD, Ridker PM, Mora S. Novel protein glycan derived
419 biomarker is associated with incident diabetes. *Circulation* 2013; **2013**(A18807).
- 420

- 421 16. Connelly MA, Gruppen EG, Wolak-Dinsmore J, Matyus SP, Riphagen IJ, Shalaurova
 422 I *et al.* GlycA, a marker of acute phase glycoproteins, and the risk of incident type 2
 423 diabetes mellitus: PREVEND study. *Clin Chim Acta* 2016; **452**: 10-7.
- 424
 425 17. Akinkuolie AO, Buring JE, Ridker PM, Mora S. A novel protein glycan biomarker
 426 and future cardiovascular disease events. *J Amer Heart Assoc* 2014; **3**(5): e001221.
- 427
 428 18. Akinkuolie AO, Glynn RJ, Ridker PM, Mora S. Protein glycan side-chains,
 429 rosuvastatin therapy, and incident vascular events: an analysis from the JUPITER
 430 trial. *Circulation* 2014; **130**: A2509.
- 431
 432 19. Duprez D, Neuhaus J, Otvos J, Neaton JD, Lundgren JD. GlycA, a novel marker of
 433 inflammation, predicts cardiovascular events in HIC-positive patients: results of
 434 SMART study. *Circulation* 2014; **130**(A2509).
- 435
 436 20. Gruppen EG, Riphagen IJ, Connelly MA, Otvos JD, Bakker SJ, Dullaart RP. GlycA, a
 437 Pro-Inflammatory Glycoprotein Biomarker, and Incident Cardiovascular Disease:
 438 Relationship with C-Reactive Protein and Renal Function. *PLoS One* 2015; **10**(9):
 439 e0139057.
- 440
 441 21. Mora S, Otvos JD, Rosenson RS, Pradhan A, Buring JE, Ridker PM. Lipoprotein
 442 particle size and concentration by nuclear magnetic resonance and incident type 2
 443 diabetes in women. *Diabetes* 2010; **59**(5): 1153-60.
- 444
 445 22. Mackey RH, Mora S, Bertoni AG, Wassel CL, Carnethon MR, Sibley CT *et al.*
 446 Lipoprotein particles and incident type 2 diabetes in the multi-ethnic study of
 447 atherosclerosis. *Diabetes Care* 2015; **38**(4): 628-36.
- 448
 449 23. Mietus-Snyder M, Drews KL, Otvos JD, Willi SM, Foster GD, Jago R *et al.* Low-
 450 Density Lipoprotein Cholesterol versus Particle Number in Middle School Children. *J*
 451 *Pediatr* 2013.
- 452
 453 24. Jago R, Drews KL, Otvos JD, Foster GD, Marcus MD, Buse JB *et al.* Effect of
 454 relative weight group change on nuclear magnetic resonance spectroscopy derived
 455 lipoprotein particle size and concentrations among adolescents. *J Pediatr* 2014;
 456 **164**(5): 1091-1098 e3.
- 457
 458 25. Buse J, Hirst K. The HEALTHY study: introduction. *Int J Obes (Lond)* 2009; **33**
 459 **Suppl 4**: S1-2.
- 460
 461 26. The Healthy Study Group. A School-Based Intervention for Diabetes Risk Reduction.
 462 *N Engl J Med* 2010; **363**(5): 445-53.

- 463
464 27. DeBar LL, Schneider M, Ford EG, Hernandez AE, Showell B, Drews KL *et al.* Social
465 marketing-based communications to integrate and support the HEALTHY study
466 intervention. *Int J Obes (Lond)* 2009; **33 Suppl 4**: S52-9.
- 467
468 28. Gillis B, Mobley C, Stadler DD, Hartstein J, Virus A, Volpe SL *et al.* Rationale,
469 design and methods of the HEALTHY study nutrition intervention component. *Int J*
470 *Obes (Lond)* 2009; **33 Suppl 4**: S29-36.
- 471
472 29. Jago R, McMurray RG, Drews KL, Moe EL, Murray T, Pham TH *et al.* HEALTHY
473 Intervention: Fitness, Physical Activity, and Metabolic Syndrome Results. *Med Sci*
474 *Sports Exerc* 2011; **43(8)**: 1513-22.
- 475
476 30. Venditti EM, Elliot DL, Faith MS, Firrell LS, Giles CM, Goldberg L *et al.* Rationale,
477 design and methods of the HEALTHY study behavior intervention component. *Int J*
478 *Obes (Lond)* 2009; **33 Suppl 4**: S44-51.
- 479
480 31. Foster GD, Linder B, Baranowski T, Cooper DM, Goldberg L, Harrell JS *et al.* A
481 school-based intervention for diabetes risk reduction. *N Engl J Med* 2010; **363(5)**:
482 443-53.
- 483
484 32. McMurray RG, Bassin S, Jago R, Bruecker S, Moe EL, Murray T *et al.* Rationale,
485 design and methods of the HEALTHY study physical education intervention
486 component. *Int J Obes (Lond)* 2009; **33 Suppl 4**: S37-43.
- 487
488 33. Jago R, Bailey R. Ethics and paediatric exercise science: Issues and making a
489 submission to a local ethics and research committee. *Journal of Sport Sciences* 2001;
490 **19(7)**: 527-535.
- 491
492 34. Petersen AC, Crockett L, Richards M, Boxer A. A self-report measure of pubertal
493 status: reliability, validity, and initial norms. *Youth Adol* 1988; **17**: 117-133.
- 494
495 35. Tanner JM. *Growth at adolescence*, Blackwell: Oxford, 1962.
- 496
497 36. Centers for Disease Control National Center for Health Statistics. 2000 CDC growth
498 charts for the United States. In. Atlanta Centers for Disease Control, 2009.
- 499
500 37. Leger LA, Lambert J. A maximal multistage 20-m shuttle run test to predict VO₂
501 max. *Eur J Appl Physiol Occup Physiol* 1982; **49(1)**: 1-12.
- 502
503 38. Leger LA, Mercier D, Gadoury C, Lambert J. The multistage 20 metre shuttle run test
504 for aerobic fitness. *Journal of sports sciences* 1988; **6(2)**: 93-101.

- 505
506 39. Willi SM, Hirst K, Jago R, Buse J, Kaufman F, El Ghormli L *et al.* Cardiovascular
507 risk factors in multi-ethnic middle school students: the HEALTHY primary
508 prevention trial. *Pediatric obesity* 2012; **7**(3): 230-9.
- 509
510 40. Fridewald WT, Levy RI, Fredrickson DS. Estimation of the concentration of low-
511 density lipoprotein cholesterol in plasma, without use of the preparative
512 ultracentrifuge. *Clinical Chem* 1972; **18**: 499-502.
- 513
514 41. Marcovina S, Bowsher RR, Miller WG, Staten M, Myers G, Caudill SP *et al.*
515 Standardization of insulin immunoassays: report of the American Diabetes
516 Association Workgroup. *Clinical Chem* 2007; **53**(4): 711-6.
- 517
518 42. Matthews DR, Hosker JP, Rudenski AS, Naylor BA, Treacher DF, Turner RC.
519 Homeostasis model assessment: insulin resistance and beta-cell function from fasting
520 plasma glucose and insulin concentrations in man. *Diabetologia* 1985; **28**(7): 412-9.
- 521
522 43. Williams MJ, Milne BJ, Hancox RJ, Poulton R. C-reactive protein and
523 cardiorespiratory fitness in young adults. *Eur J Cardiovasc Prev Rehabil* 2005; **12**(3):
524 216-20.
- 525
526 44. Balducci S, Zanuso S, Cardelli P, Salvi L, Mazzitelli G, Bazuro A *et al.* Changes in
527 physical fitness predict improvements in modifiable cardiovascular risk factors
528 independently of body weight loss in subjects with type 2 diabetes participating in the
529 Italian Diabetes and Exercise Study (IDES). *Diabetes Care* 2012; **35**(6): 1347-54.
- 530
531 45. Martinez-Gomez D, Eisenmann JC, Warnberg J, Gomez-Martinez S, Veses A, Veiga
532 OL *et al.* Associations of physical activity, cardiorespiratory fitness and fatness with
533 low-grade inflammation in adolescents: the AFINOS Study. *Int J Obes (Lond)* 2010;
534 **34**(10): 1501-7.
- 535
536 46. Lakoski SG, Cushman M, Criqui M, Rundek T, Blumenthal RS, D'Agostino RB, Jr. *et*
537 *al.* Gender and C-reactive protein: data from the Multiethnic Study of Atherosclerosis
538 (MESA) cohort. *Amer Heart J* 2006; **152**(3): 593-8.
- 539
540 47. Ford ES, Giles WH, Mokdad AH, Myers GL. Distribution and correlates of C-
541 reactive protein concentrations among adult US women. *Clinical Chem* 2004; **50**(3):
542 574-81.
- 543
544 48. Cushman M, Arnold AM, Psaty BM, Manolio TA, Kuller LH, Burke GL *et al.* C-
545 reactive protein and the 10-year incidence of coronary heart disease in older men and
546 women: the cardiovascular health study. *Circulation* 2005; **112**(1): 25-31.
- 547

- 548 49. Moran A, Jacobs DR, Jr., Steinberger J, Hong CP, Prineas R, Luepker R *et al.* Insulin
549 resistance during puberty: results from clamp studies in 357 children. *Diabetes* 1999;
550 **48(10)**: 2039-44.
- 551
552 50. Liu NY, Plowman SA, Looney MA. The reliability and validity of the 20-meter
553 shuttle test in American students 12 to 15 years old. *Res Q Exerc Sport* 1992; **63(4)**:
554 360-5.
- 555
556 51. van Mechelen W, Hlobil H, Kemper HCG. Validation of two running tests as
557 estimates of maximal aerobic power in children. *Eur J Appl Physiol Occup Physiol*
558 1986; **55(5)**: 503–506.
- 559
560 52. Boreham CA, Paliczka VJ, Nichols AK. A comparison of the PWC170 and 20-MST
561 tests of aerobic fitness in adolescent schoolchildren. *J Sports Med Phys Fitness* 1990;
562 **30(1)**: 19-23.

563

564 **FIGURE LEGENDS**

565 **Figure 1:** Inflammation (GlycA) in Obesity and Fitness Subgroups

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567 **Figure 2:** Insulin Resistance (LP-IR) in Obesity and Fitness Subgroups

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Table 1: Baseline Characteristics

	Gender			<i>p</i> -value**
	OVERALL	Female (N=897)	Male (N=767)	
	Mean (SD) or N and %	Mean (SD) or N and %	Mean (SD) or N and %	
Age (years)	11.28 (0.55)	11.22 (0.50)	11.35 (0.60)	<.0001
Number of Laps	20.63 (11.60)	18.53 (9.33)	23.09 (13.38)	<.0001
BMI Percentile	73.75 (27.48)	72.37 (27.35)	75.35 (27.57)	0.0264
BMI Category				0.0019
< 85 th Percentile	818 49.2%	471 52.5%	347 45.2%	
85 th – 94 th Percentile	331 19.9%	184 20.5%	147 19.2%	
≥ 95 th Percentile	515 30.9%	242 27.0%	273 35.6%	
Race/Ethnicity				0.3107
Hispanic	1044 62.7%	573 63.9%	471 61.4%	
Black	276 16.6%	151 16.8%	125 16.3%	
White	344 20.7%	173 19.3%	171 22.3%	
Positive Reported 1 st Degree Family History of Diabetes	215 12.9%	115 12.8%	100 13.0%	0.9214
Highest Household Education				0.8817
≤ HS Graduate	837 50.3%	473 52.7%	364 47.5%	
≥ Some college	827 49.7%	424 47.3%	403 52.5%	
6 th Grade Pubertal Status				*
Tanner Stage 1	168 10.1%	49 5.5%	119 15.5%	
Tanner Stage 2	436 26.2%	120 13.4%	316 41.2%	
Tanner Stage 3	669 40.2%	379 42.3%	290 37.8%	
Tanner Stage 4	357 21.5%	315 35.1%	42 5.5%	
Tanner Stage 5	34 2.0%	34 3.8%	0 0	

*Test does not converge due to zero cells.

***p*-values obtained from generalized linear mixed models taking account of sources of variability within and between schools.

Table 2: 6th Grade, 8th Grade and Change (8th-6th) in Fitness, Lipids, Insulin Resistance and GlycA by Gender

	<i>Female</i>						<i>Male</i>					
	<i>6th Grade</i>		<i>8th Grade</i>		<i>Difference (8th – 6th)</i>		<i>6th Grade</i>		<i>8th Grade</i>		<i>Difference (8th – 6th)</i>	
<i>Fitness (# of laps)</i>	18.5	(17.9, 19.2)	21.2	(20.4, 21.9)	2.7	(2.0, 3.3)	23.1	(22.1, 24.0)	33.9	(32.6, 35.3)	10.8	(9.7, 12.0)
<i>Non-HDL-C (mg/dL)</i>	102	(86, 120)	95	(81, 110)	-7	(-18, 4)	105	(88, 124)	93	(79, 111)	-11	(-23, 1)
<i>LDL-C (mg/dL)</i>	83	(71, 99)	79	(67, 93)	-4	(-15, 5)	88	(73, 104)	77	(64, 93)	-10	(-21, -0)
<i>HDL-C (mg/dL)</i>	51	(44, 59)	53	(45, 62)	2	(-3, 7)	51	(44, 60)	48	(41, 56)	-3	(-8, 2)
<i>HOMA-IR</i>	2.62	(1.76, 4.19)	3.35	(2.34, 4.77)	0.61	(-0.42, 1.74)	2.11	(1.36, 3.36)	3.03	(1.94, 4.62)	0.84	(-0.12, 2.12)
<i>LP-IR</i>	32	(20, 49)	28	(16, 42)	-5	(-15, 5)	38	(22, 58)	36	(22, 54)	-1	(-13, 9)
<i>GlycA (μmol/L)</i>	374	(334, 421)	397	(361, 442)	24	(-16, 58)	379	(333, 425)	365	(322, 418)	-10	(-49, 32)

Non-HDL-C, LDL-C, HDL-C, HOMA-IR, LP-IR and GlycA are presented as medians and (25th percentile, 75th percentile) while fitness is presented as mean and 95% confidence interval.

Figure 1: Inflammation (GlycA) in Obesity and Fitness Subgroups

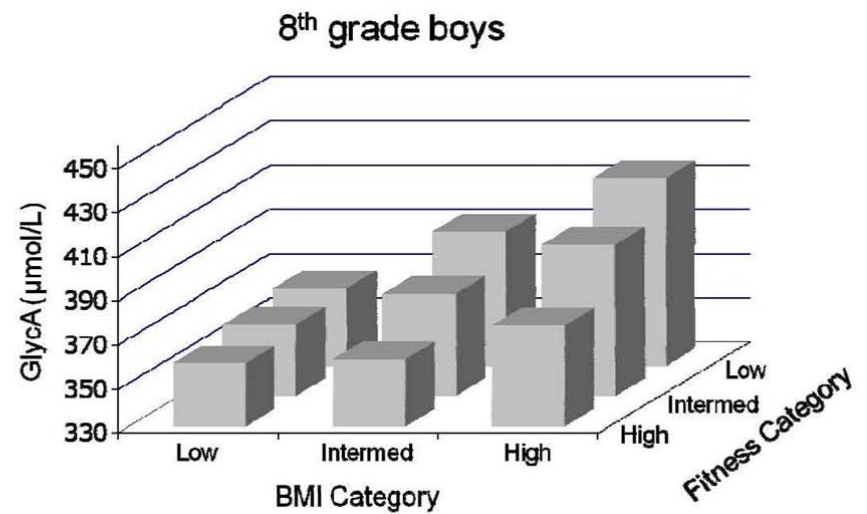
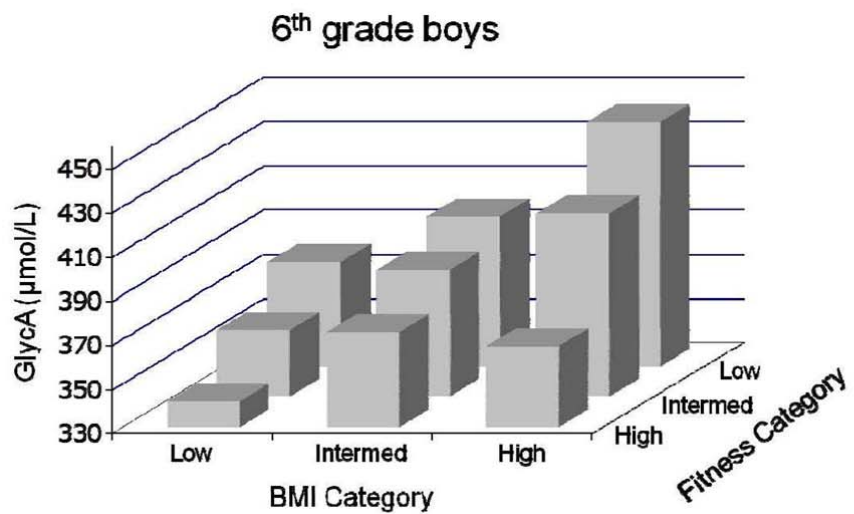
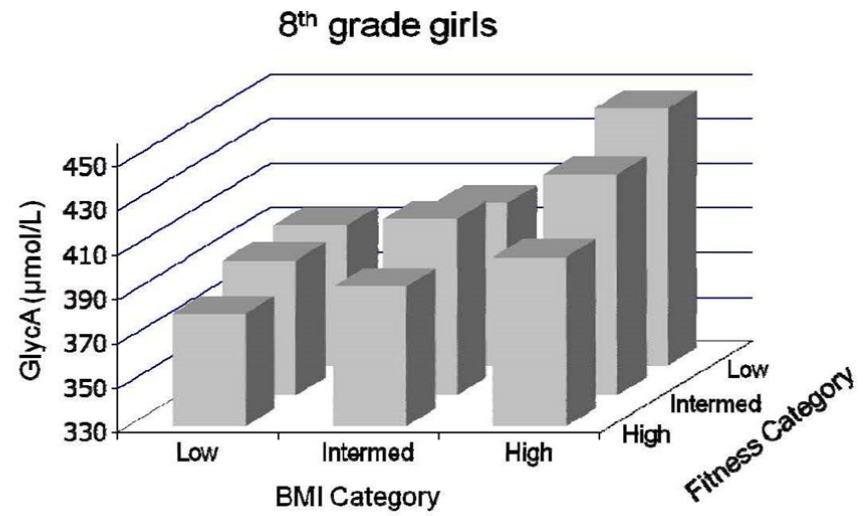
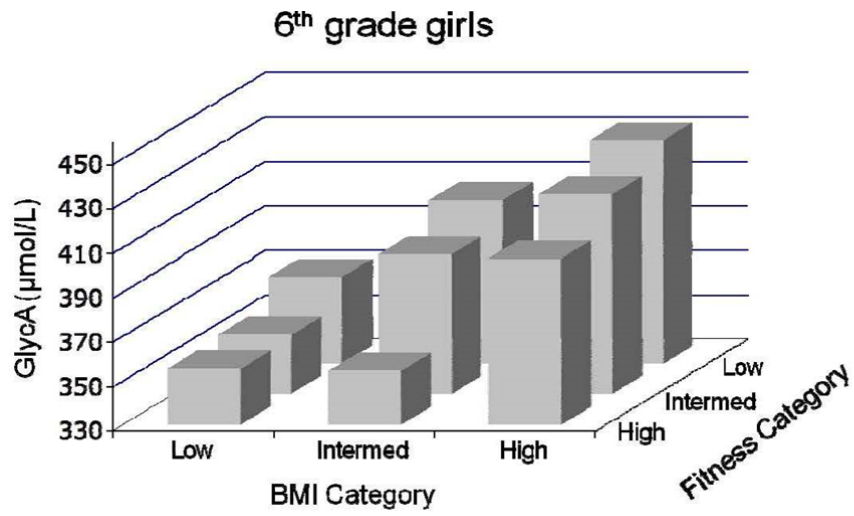


Figure 2: Insulin Resistance (LP-IR) in Obesity and Fitness Subgroups

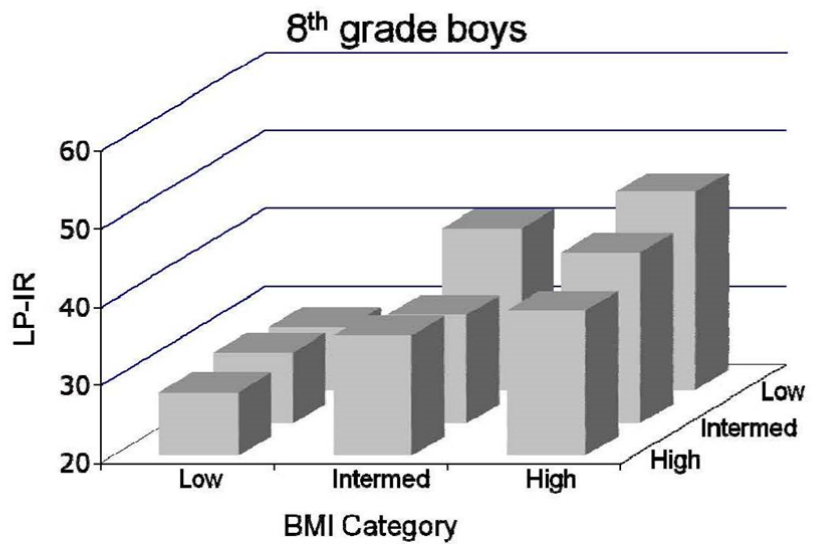
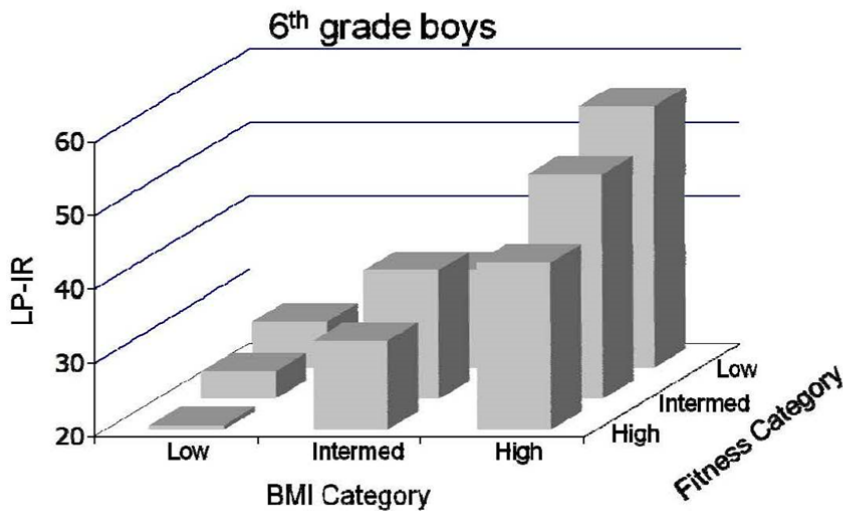
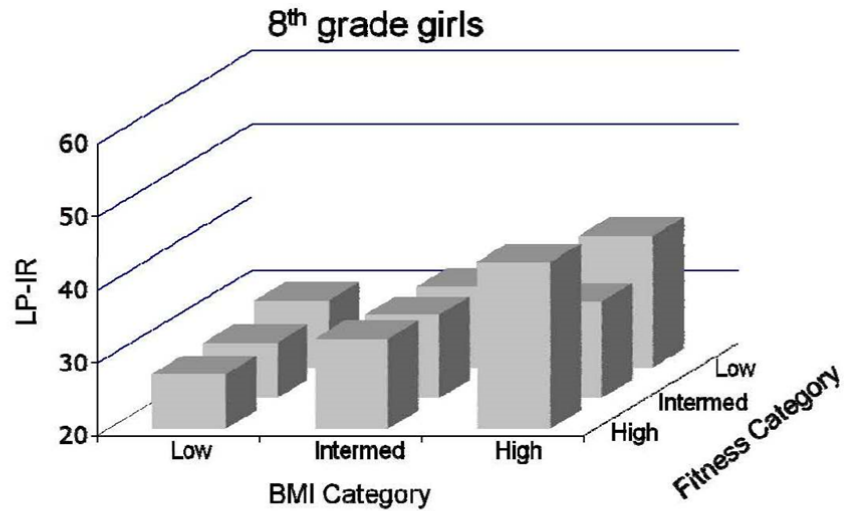
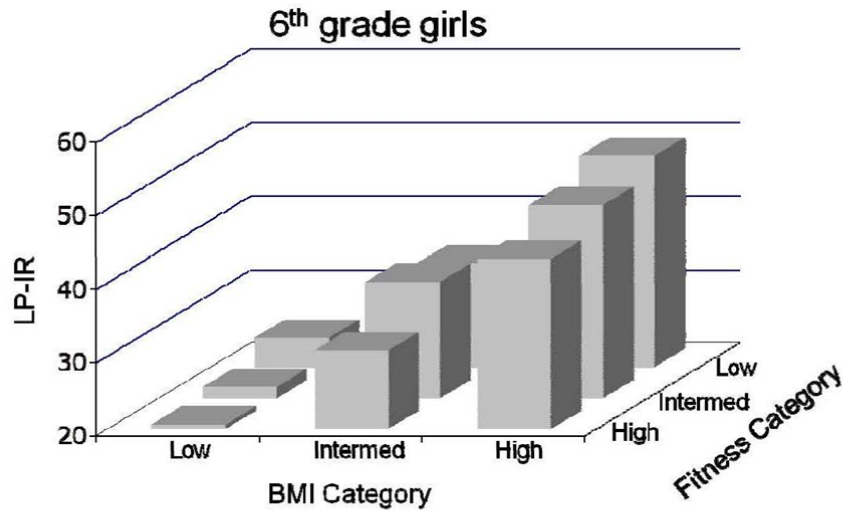


Table A: Comparison of Subjects Included and Excluded from Analysis

	<i>Analysis Status</i>			<i>p-value**</i>
	<i>OVERALL</i>	<i>Included</i>	<i>Excluded</i>	
	<i>Mean (SD) or</i>	<i>Mean (SD) or</i>	<i>Mean (SD) or</i>	
	<i>N and %</i>	<i>N and %</i>	<i>N and %</i>	
<i>Age (years)</i>	11.27 (0.57)	11.28 (0.55)	11.26 (0.60)	0.6780
<i>Gender</i>				0.3834
<i>Female</i>	1288 54.4%	897 53.9%	391 55.6%	
<i>Male</i>	1079 45.6%	767 46.1%	312 44.4%	
<i>Number of Laps</i>	20.48 (11.63)	20.63 (11.60)	20.05 (11.72)	0.0617
<i>Randomization Status</i>				0.3756
<i>Control</i>	1229 51.9%	835 50.2%	394 56.0%	
<i>Intervention</i>	1138 48.1%	829 49.8%	309 44.0%	
<i>Baseline BMI Percentile</i>	72.89 (27.98)	73.75 (27.48)	70.86 (29.05)	0.0792
<i>Baseline BMI Category</i>				0.6775
< 85 th Percentile	1183 50.0%	818 49.2%	365 51.9%	
85 th – 94 th Percentile	462 19.5%	331 19.9%	131 18.6%	
≥ 95 th Percentile	722 30.5%	515 30.9%	207 29.4%	
<i>Race/Ethnicity</i>				0.0043*
<i>Hispanic</i>	1328 56.1%	1044 62.7%	284 40.4%	
<i>Black</i>	402 17.0%	276 16.6%	126 17.9%	
<i>White</i>	430 18.2%	344 20.7%	86 12.2%	
<i>Other</i>	207 8.7%	0 0%	207 29.4%	
<i>Highest Household Education</i>				0.4280
≤ HS Graduate	1136 49.1%	837 50.3%	299 46.1%	
≥ Some college	1177 50.9%	827 49.7%	350 53.9%	
<i>Baseline LDL-C (mg/dL)</i>	87.26 (22.68)	87.50 (22.54)	86.70 (23.02)	0.1813
<i>Baseline HDL-C (mg/dL)</i>	52.81 (12.33)	53.52 (12.96)	52.51 (12.05)	0.4280

*Other Race not used for test

**p-values obtained from generalized linear mixed models taking account of sources of variability within and between schools.

Table B: Percentiles of GlycA ($\mu\text{mol/L}$) by Gender and Grade

	<i>Girls</i>		<i>Boys</i>	
	<i>6th Grade</i>	<i>8th Grade</i>	<i>6th Grade</i>	<i>8th Grade</i>
<i>Minimum</i>	223	251	243	235
<i>1st Percentile</i>	272	283	264	246
<i>5th Percentile</i>	293	309	284	279
<i>10th Percentile</i>	308	330	302	296
<i>25th Percentile</i>	334	361	333	322
<i>50th Percentile</i>	374	397	379	365
<i>75th Percentile</i>	421	442	425	418
<i>90th Percentile</i>	470	487	477	470
<i>95th Percentile</i>	501	519	505	502
<i>99th Percentile</i>	562	599	549	579
<i>Maximum</i>	807	755	591	771

Table C: Medians for GlycA by Weight Category and Gender and Tests for Associations between Category and GlycA

	Girls						
	<i>BMI < 85th Percentile</i>		<i>BMI 85th – 94th Percentile</i>		<i>BMI ≥ 95th Percentile</i>		<i>p-value</i>
	<i>Median</i>	<i>(Q1, Q3)</i>	<i>Median</i>	<i>(Q1, Q3)</i>	<i>Median</i>	<i>(Q1, Q3)</i>	
<i>6th Grade GlycA* (μmol/L)</i>	351	(317, 392)	386	(347, 424)	417	(386, 464)	<.0001
<i>8th Grade GlycA** (μmol/L)</i>	379	(344, 417)	407	(367, 444)	443	(403, 488)	<.0001

	Boys						
	<i>BMI < 85th Percentile</i>		<i>BMI 85th – 94th Percentile</i>		<i>BMI ≥ 95th Percentile</i>		<i>p-value</i>
	<i>Median</i>	<i>(Q1, Q3)</i>	<i>Median</i>	<i>(Q1, Q3)</i>	<i>Median</i>	<i>(Q1, Q3)</i>	
<i>6th Grade GlycA* (μmol/L)</i>	340	(311, 386)	378	(344, 417)	419	(386, 459)	<.0001
<i>8th Grade GlycA** (μmol/L)</i>	341	(306, 377)	369	(332, 417)	415	(379, 463)	<.0001

*Models adjusted for 6th grade Tanner stage, race/ethnicity, highest household education, and intervention group

**Models adjusted for 8th grade Tanner stage, 6th grade value of the GlycA, race/ethnicity, highest household education, and intervention group

Table D: Spearman Correlations for Between GlycA, Lipids, Lipoproteins, Insulin Resistance, Fitness and BMI (not adjusting for school cluster) by Grade

<i>6th Grade</i>								
	<i>LP-IR</i>	<i>HOMA-IR</i>	<i>BMI %ile</i>	<i>HDL-C</i>	<i>LDL-C</i>	<i>LDL-P</i>	<i>Non-HDL-C</i>	<i>Number of Laps</i>
GlycA (μmol/L)	0.41	0.41	0.50	-0.33	0.18	0.39	0.29	-0.37
LP-IR	--	0.49	0.59	-0.69	0.13	0.54	0.37	-0.34
HOMA-IR		--	0.65	-0.41	0.09	0.31	0.24	-0.41
BMI Percentile			--	-0.46	0.18	0.41	0.31	-0.50
HDL-C (mg/dL)				--	-0.03	-0.39	-0.20	0.29
LDL-C (mg/dL)					--	0.68	0.92	-0.10
LDL-P (nmol/L)						--	0.76	-0.28
Non-HDL-C (mg/dL)							--	-0.18
<i>8th Grade</i>								
	<i>LP-IR</i>	<i>HOMA-IR</i>	<i>BMI %ile</i>	<i>HDL-C</i>	<i>LDL-C</i>	<i>LDL-P</i>	<i>Non-HDL-C</i>	<i>Number of Laps</i>
GlycA (μmol/L)	0.31	0.42	0.43	-0.24	0.20	0.35	0.28	-0.35
LP-IR	--	0.43	0.47	-0.72	0.09	0.48	0.28	-0.18
HOMA-IR		--	0.55	-0.30	0.12	0.28	0.24	-0.33
BMI Percentile			--	-0.41	0.19	0.38	0.28	-0.37
HDL-C (mg/dL)				--	-0.04	-0.42	-0.16	0.10
LDL-C (mg/dL)					--	0.70	0.94	-0.15
LDL-P (nmol/L)						--	0.76	-0.23
Non-HDL-C (mg/dL)							--	-0.19

Table E: Adjusted mean (SE) level of measures of inflammation, insulin resistance, and atherogenic lipoproteins by obesity and fitness subgroup

Girls									
<i>6th Grade</i>									
	<i>BMI <85th %ile</i>			<i>BMI 85th – 94th %ile</i>			<i>BMI ≥95th %ile</i>		
	<i>Low Fitness</i>	<i>Med Fitness</i>	<i>High Fitness</i>	<i>Low Fitness</i>	<i>Med Fitness</i>	<i>High Fitness</i>	<i>Low Fitness</i>	<i>Med Fitness</i>	<i>High Fitness</i>
<i>GlycA (μmol/L)</i>	369.13 (8.59)	356.81 (5.34)	355.31 (5.49)	403.43 (9.09)	392.67 (6.88)	354.42 (10.55)	430.16 (6.40)	419.56 (6.54)	404.77 (16.55)
<i>LP-IR</i>	23.96 (0.05)	21.50 (0.02)	20.59 (0.02)	34.14 (0.06)	35.94 (0.03)	30.83 (0.08)	48.78 (0.03)	46.27 (0.03)	43.18 (0.20)
<i>LDL-P (nmol/L)</i>	611.98 (33.79)	646.68 (20.76)	602.41 (21.38)	712.57 (35.84)	755.87 (26.95)	664.56 (41.65)	876.62 (25.04)	866.93 (25.59)	776.56 (65.60)
<i>Non-HDL-C (mg/dL)</i>	99.04 (3.40)	101.17 (2.02)	100.05 (2.10)	108.98 (3.63)	111.12 (2.69)	100.59 (4.24)	115.26 (2.49)	117.46 (2.54)	110.24 (6.74)
<i>8th Grade</i>									
<i>GlycA (μmol/L)</i>	393.05 (11.92)	390.87 (10.67)	381.04 (11.06)	402.69 (13.34)	409.21 (11.35)	392.93 (13.45)	445.79 (11.22)	429.86 (11.21)	406.27 (17.46)
<i>LP-IR</i>	29.23 (0.09)	27.58 (0.07)	27.52 (0.08)	31.12 (0.11)	31.49 (0.08)	32.36 (0.12)	37.97 (0.08)	33.31 (0.08)	42.90 (0.20)
<i>LDL-P (nmol/L)</i>	709.12 (39.57)	718.39 (35.43)	716.93 (36.52)	737.51 (44.60)	759.70 (38.00)	696.38 (44.31)	832.58 (37.67)	731.10 (37.71)	774.29 (58.31)
<i>Non-HDL-C (mg/dL)</i>	100.66 (3.56)	99.76 (3.18)	99.33 (3.29)	99.57 (4.00)	102.59 (3.40)	95.60 (4.01)	104.69 (3.34)	100.15 (3.35)	98.10 (5.21)
Boys									
<i>6th Grade</i>									
	<i>BMI <85th %ile</i>			<i>BMI 85th – 94th %ile</i>			<i>BMI ≥95th %ile</i>		
	<i>Low Fitness</i>	<i>Med Fitness</i>	<i>High Fitness</i>	<i>Low Fitness</i>	<i>Med Fitness</i>	<i>High Fitness</i>	<i>Low Fitness</i>	<i>Med Fitness</i>	<i>High Fitness</i>
<i>GlycA (μmol/L)</i>	377.24 (11.10)	360.40 (5.39)	341.77 (5.39)	397.87 (11.13)	387.21 (6.89)	372.97 (9.71)	440.58 (5.85)	412.96 (5.79)	366.88 (22.78)
<i>LP-IR</i>	26.26 (0.08)	23.71 (0.02)	20.61 (0.02)	33.16 (0.08)	37.44 (0.03)	32.10 (0.06)	55.51 (0.02)	50.49 (0.02)	42.77 (0.35)
<i>LDL-P (nmol/L)</i>	642.09 (54.83)	619.67 (25.71)	610.24 (25.69)	808.29 (54.97)	770.36 (33.56)	744.21 (47.83)	1024.60 (28.10)	857.27 (27.87)	735.31 (113.67)
<i>Non-HDL-C (mg/dL)</i>	96.00 (4.94)	99.80 (2.25)	96.58 (2.25)	111.92 (4.95)	114.36 (3.00)	108.11 (4.30)	127.99 (2.48)	114.85 (2.46)	105.85 (10.33)
<i>8th Grade</i>									
<i>GlycA (μmol/L)</i>	364.98 (11.10)	362.42 (7.80)	358.62 (8.21)	390.55 (14.69)	376.18 (8.86)	360.56 (13.41)	415.15 (8.41)	398.60 (9.05)	376.11 (19.52)

<i>LP-IR</i>	28.13 (0.06)	29.14 (0.03)	28.16 (0.04)	40.69 (0.11)	33.91 (0.04)	35.47 (0.09)	45.39 (0.04)	41.78 (0.04)	38.07 (0.20)
<i>LDL-P (nmol/L)</i>	686.11 (38.31)	685.60 (26.89)	689.00 (28.37)	808.40 (51.11)	708.59 (30.87)	689.42 (46.57)	869.99 (29.33)	834.98 (31.60)	672.09 (67.91)
<i>Non-HDL-C (mg/dL)</i>	89.23 (3.31)	93.30 (2.34)	93.31 (2.47)	100.36 (4.37)	93.29 (2.68)	89.11 (3.98)	103.03 (2.53)	100.43 (2.72)	91.74 (5.76)

Low fitness=Q1 (6th Grade – Girls: 0-11 laps, Boys: 0-12 laps; 8th Grade – Girls: 1-12 laps, Boys: 1-17 laps),

Medium Fitness=Q2 & Q3 (6th Grade – Girls: 12-23 laps, Boys: 13-30 laps; 8th Grade – Girls: 13-26 laps, Boys: 18-44 laps)

High Fitness=Q4 (6th Grade – Girls: 24-57 laps, Boys: 31-75 laps; 8th Grade – Girls: 27-79 laps, Boys: 45-103 laps)

6th grade adjusted for 6th grade Tanner stage, race/ethnicity, SES category and intervention status

8th grade adjusted for 8th grade Tanner stage, race/ethnicity, SES category, intervention status and 6th grade value of the dependent variable (GlycA, LP-IR, LDL-P or Non-HDL-C)