

1 THE ROLE OF INTER-SET REST PERIODS IN 2 HIGH-INTENSITY INTERVAL TRAINING ON 3 ACUTE GLUCOSE AND INSULIN SENSITIVITY IN 4 TYPE I DIABETIC CHILDREN: A RANDOMIZED 5 CROSSOVER STUDY

6 Abstract

7 **Aims:** To investigate the impact of different inter-set rest periods on blood glucose levels
8 and insulin sensitivity (IS) in children with type 1 diabetes.

9 **Study Design:** An experimental randomized crossover trial.

10 **Place and Duration:** Recruitment took place in the public health network of Rio Branco,
11 Acre, Brazil.

12 **Methodology:** Twenty subjects participated in three different HIIT protocols with varying
13 rest intervals. The first protocol consisted of 20 seconds of exercise followed by 10 seconds
14 of rest (HIIT20-10), the second with 20 seconds of rest (HIIT20-20), and the third with 60
15 seconds of rest (HIIT20-60). The exercise used in all protocols was the burpee, performed
16 at maximum effort for a total of 12 minutes, comprising 24 sets. Blood glucose levels were
17 measured through finger-prick tests before exercise, immediately after, and at 15, 30, 60,
18 and 120 minutes post-exercise. For insulin sensitivity, 4 ml of venous blood was drawn
19 before, immediately after, and at 30, 60, and 120 minutes post-exercise, and the TyG Index
20 was used to calculate IS.

21 **Results:** No significant differences were observed between the HIIT20-20 and HIIT20-60
22 groups ($p>0.05$) regarding glucose levels. However, the HIIT20-10 protocol showed
23 significantly lower glucose levels from 15 minutes to 120 minutes post-exercise ($p<0.05$)
24 and improved IS at 60 minutes ($p<0.05$) and 120 minutes ($p<0.01$) after the exercise
25 session.

26 **Conclusion:** The inter-set rest period influences glucose metabolism during HIIT in children
27 with type 1 diabetes.

28
29
30
31
32
33
34
35
36
37
38
Key Words: Metabolic Health, Glucose Metabolism, Type I Diabetes, High-Intensity Interval
Training, Insulin Sensitivity

39

40 1. Introduction

41 Type 1 diabetes (T1D) is a chronic autoimmune disease that destroys the insulin-producing β -
42 cells of the pancreas, leading to insulin deficiency. This condition is a significant public health
43 concern worldwide due to its silent progression and associated complications[1]. T1D is often
44 accompanied by metabolic disorders, including insulin resistance and central fat accumulation,
45 which are predictive factors for cardiovascular disease[2].

46 Consistent evidence shows that a sedentary lifestyle is linked to metabolic impairments, while
47 physical exercise can prevent or reverse these effects[3]. Exercise has been shown to improve
48 the body's systemic responses to various nutrients, including glucose[4]. Specifically, physical
49 activity is recognized for its positive effects on metabolic diseases such as diabetes[5–7]. In
50 contrast, a sedentary lifestyle is associated with the development and worsening of conditions
51 like diabetes[1].

52 Our research group has demonstrated that high-intensity interval training (HIIT) is an effective
53 tool for combating metabolic diseases in both young and older adults with diabetes or metabolic
54 syndrome[8]. HIIT-based programs offer non-pharmacological benefits in treating insulin
55 resistance (IR) and diabetes[9], particularly through the use of HIIT[10].

56 In 2017, the American Diabetes Association recommended 30-60 minutes of moderate-intensity
57 exercise for children and adolescents with T1D[11]. However, the optimal exercise guidelines
58 for improving both acute and long-term glycemic control in youth with T1D are still unclear.
59 Previous studies have shown that fasted HIIT is safe[12] and can consistently increase blood
60 glucose levels during exercise. Intense interval exercise is often linked to higher blood glucose
61 profiles and a lower risk of hypoglycemia during exercise compared to continuous exercise[13].
62 These findings are largely due to increased counter-regulatory hormone secretion, which
63 enhances hepatic glucose production and reduces peripheral glucose disposal[14,15].

64 In healthy adolescents, a single bout of time-efficient HIIT improves glucose tolerance and
65 insulin sensitivity, suggesting HIIT as a viable strategy for managing glycemic control in youth
66 with T1D[12,14]. However, the relationship between exercise stimulus, rest periods, total
67 exercise volume, and intensity-key variables in exercise prescription has not been thoroughly
68 investigated[8]. This results in inconsistent evidence supporting the optimal HIIT dosage for
69 improving glycemic control in children with T1D.

70 This study's findings are particularly significant for the scientific community as they provide
71 insights into how different HIIT protocols may manage glucose levels in children with type 1
72 diabetes. Given the increasing prevalence of diabetes in children, understanding the effects of
73 various exercise regimens on metabolic health is crucial. We hypothesize that exercise induces
74 positive acute adaptations in glucose metabolism and insulin resistance. Thus, we aimed to
75 investigate the impact of different interest rest on blood glucose levels, and insulin sensitivity
76 (IS).

77 2. Methods

78 2.1 Study type, participants and ethic

79 This study employed a randomized crossover design with blinding during the selection process to
80 minimize bias. Children with type 1 diabetes were selected based on specific inclusion criteria, such
81 as a medical history free of cardiovascular complications and no limitations on physical exercise.

82 Participants: During the initial week of the experiment, 20 children (age: 12.8 ± 2.1 years, weight:
83 60.4 ± 16.8 kg, height: 145 ± 22.35 cm, BMI: 28.7 ± 14.8 kg/m²) were randomly assigned to different
84 intervention groups using a 1:1:1:1 ratio. Over the course of the study, each group was reassigned to
85 different interventions weekly, following a randomized sequence. A 6-day washout period between
86 interventions was implemented to mitigate the influence of prior exercise sessions. Figure 1
87 illustrates the crossover method utilized in this study.

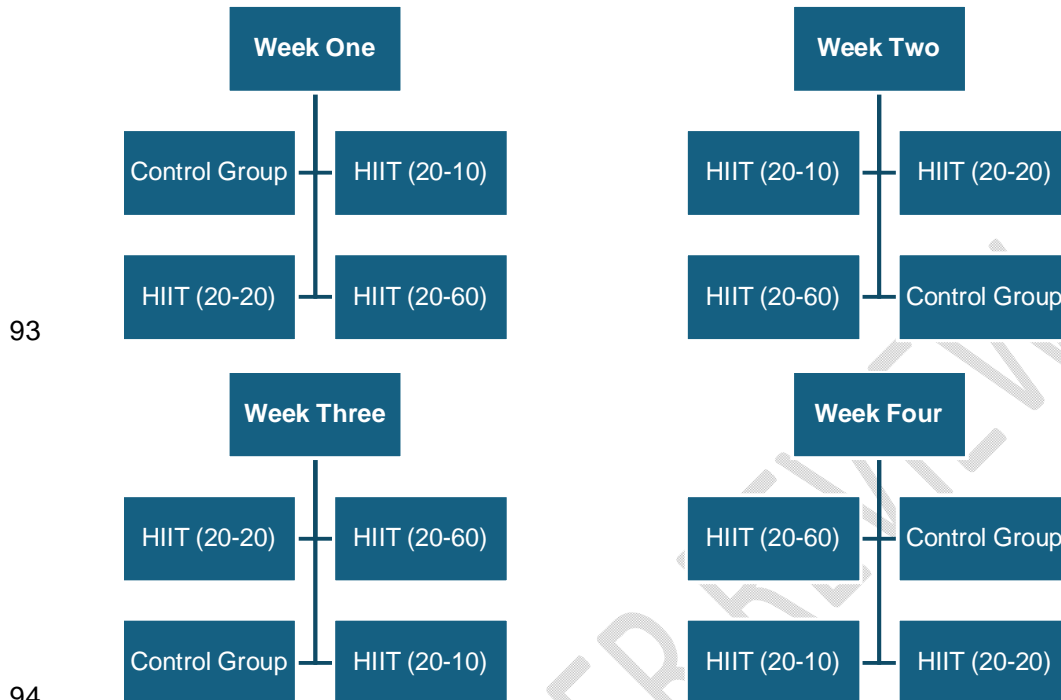
88 Interventions: Three HIIT protocols with varying rest intervals were tested:

89 **HIIT20-10:** 20 seconds of exercise followed by 10 seconds of rest.

90 **HIIT20-20:** 20 seconds of exercise followed by 20 seconds of rest.

91 **HIIT20-60:** 20 seconds of exercise followed by 60 seconds of rest.

92



95 **Figure 1:** The crossover method used to insert the groups into different intervention each week.

96 All protocols involved performing burpees at 90% or higher of maximum heart rate, monitored
97 using a G-Tech (Model ONE) finger oximeter. Each session lasted 12 minutes, consisting of 24
98 sets of exercises. Participants followed a standardized warm-up routine, including stretching and
99 light gymnastics for 5 minutes, followed by the main workout, and ended with a 5-minute cool-
100 down.

101 2.2 Basic Proceedings

102 Blood samples were collected at eight time points: baseline (before the intervention),
103 immediately after the session, and at 5, 15, 30, 60, 90, and 120 minutes post-exercise. Each
104 blood sample was processed twice independently, yielding four measurements per participant.
105 Discrepant results were excluded, and the average of consistent measurements was calculated.

106 2.3 Exercise protocols

107 Three models of high-intensity interval training (HIIT) programs were evaluated to address our
108 problem. All protocols maintained an intensity level at 90% of maximum heart rate, monitored
109 using a G-Tech (Model ONE) finger oximeter. The first protocol, HIIT20-10, consisted of 20
110 seconds of exercise followed by 10 seconds of rest. The second protocol, HIIT20-20, involved
111 20 seconds of exercise followed by 20 seconds of rest. The third protocol, HIIT20-60, comprised
112 20 seconds of exercise followed by 60 seconds of rest. The exercise performed was the burpee,
113 completed over a total duration of 12 minutes, resulting in 24 sets of stimuli. All exercises were
114 performed without equipment, relying solely on body weight.

115 To perform a burpee, start by standing with your feet shoulder-width apart and your arms at your
116 sides. Lower your body into a squat position, placing your hands on the floor in front of you, just
117 inside your feet. With your weight on your hands, kick your feet back to land in a plank position,
118 keeping your body in a straight line from head to heels. You can add an optional push-up at this

119 stage by lowering your chest to the ground and then pushing back up to the plank position.
120 Jump your feet forward to return to the squat position, with your hands still on the ground.
121 Finally, explode up from the squat position into a jump, reaching your arms overhead. Land
122 softly and immediately lower back into the squat position to begin the next repetition.

123 Do a standardized 5-minute gentle warm-up including stretching and gymnastics for 5 minutes
124 each day. This is followed by the main workout, and finally, a 5-minute cool down was
125 performed to promote cool down.

126 **2.4 Blood Sampling Acquisition and Glucose Measurement**

127 A finger blood samples were collected at eight time points: baseline (before the intervention),
128 immediately after the session, and at 5, 15, 30-, 60-, 90-, and 120-minutes post-exercise. Each
129 blood sample was processed twice independently, yielding four measurements per participant.
130 Discrepant results were excluded, and the average of consistent measurements was calculated.

131 For glucose curve assessment, blood glucose levels were measured using a glucometer
132 (ROCHE, Accutrend Plus) at each time point. Additionally, 4 ml of venous blood was drawn at
133 baseline, 60 minutes, and 120 minutes after the intervention to determine serum triglyceride
134 concentrations.

135 **2.5 Insulin Resistance Assessment**

136 The TyG index, calculated as the product of serum triglyceride concentration and fasting blood
137 glucose, was used to assess insulin resistance. The TyG index is a cost-effective and clinically
138 applicable method, with a cut-off point of 4.5, as suggested by Guerrero-Romero et al.[16].

$$3 - \text{TyG} = \frac{\{\text{Try} \times \text{Glu}\}}{2}$$

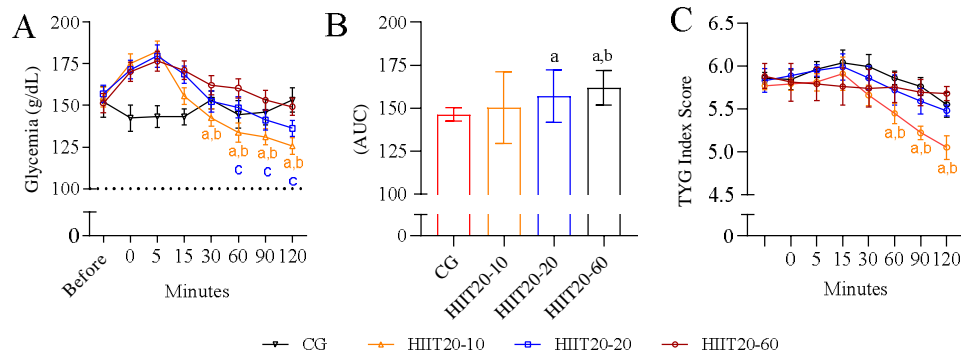
139 **Equation 1:**TyG Index. 3= Tyg Index, Try= Tryglicerides, Glu= Glucose.

140 **2.6 Statistical Procedures**

141 Descriptive data were expressed as mean, percentage, and standard error. The Shapiro-Wilk
142 test was used to assess data normality. A two-way ANOVA with Sidak's post hoc test was
143 employed to determine differences between groups, using the "GraphPad Prism 9.5.1"
144 software. A significance level of $p < 0.05$ was set for all analyses.

145 **3. Results**

146 The anthropometric characteristics of the participants, including BMI and body fat percentage,
147 were comparable across all groups ($p > 0.05$ for all). Analyzing the intragroup data, no
148 significant differences were observed in glucose control and insulin sensitivity (IS) between the
149 HIIT20-60 and the other protocols ($p > 0.05$ for both). However, the HIIT20-20 protocol showed
150 significant differences at 30, 60, 90, and 120 minutes compared to 15 minutes post-exercise (p
151 < 0.05 , 0.05 , 0.01 , and 0.01 , respectively). Notably, the HIIT20-10 protocol resulted in lower
152 glucose levels at 30, 60, 90, and 120 minutes and improved IS when comparing baseline, time
153 0, 5, and 15 minutes with 120 minutes post-exercise ($p < 0.05$, 0.05 , 0.01 , and 0.0001 ,
154 respectively), as well as 15 with 60 minutes post-exercise.



155

156 **Figure 2: Blood Glucose levels and Insulin Sensitivity.** The Figure A represent the blood
 157 glucose concentration along the time of investigation; Figure B the Area Under the Curve of the
 158 Glucose, and C is the Insulin Sensitivity. The ANOVA TWO WAY with SidákPos-Hoc Test set up
 159 at 5% was used to determine the differences between groups. **Legend:** (a,b= difference to
 160 baseline, 5 minutes, and 15 minutes, and c= difference to HIIT20-60 and control).

161

162

163 Discussion

164 This study aimed to investigate the effects of three different rest intervals in high-intensity
 165 interval training (HIIT) on acute blood glucose levels and insulin sensitivity. We found that
 166 shorter rest intervals significantly impacted the body's acute adaptations to exercise, leading to
 167 greater metabolic perturbations and a more pronounced glucose response. Specifically, shorter
 168 rest periods led to a peak in glucose levels at 15 minutes, often coinciding with a high
 169 concentration of insulin in response to the glucose surge. Subsequently, blood glucose levels
 170 decreased at 30, 60, 90, and 120 minutes, demonstrating the expected metabolic curve.
 171 Additionally, the reduction in glucose concentration during HIIT contributed to improved insulin
 172 sensitivity.

173 Although the area under the glucose curve did not differ significantly between the HIIT20-10 and
 174 control groups, the HIIT20-10 protocol exhibited a more substantial decrease in glucose levels,
 175 which likely accounts for the observed improvement in insulin sensitivity. These findings align
 176 with previous studies showing that HIIT can significantly impact blood glucose levels and insulin
 177 resistance. For example, De Teles et al.[17], investigated two HIIT protocols and reported
 178 reductions in blood glucose levels, which are consistent with our results[18].

179 In terms of comparison De teles[19] investigated two HIIT protocols: a long HIIT (2 minutes at
 180 100% VO₂peak followed by 2 minutes of passive rest) and a short HIIT (30 seconds at 100%
 181 VO₂peak followed by 30 seconds of passive rest). They reported reductions in blood glucose
 182 levels of 32.14 mg/dL and 31.40 mg/dL for the long and short HIIT protocols, respectively. These
 183 findings are consistent with our results, which showed reductions of 12.35, 21.35, 23.79, and
 184 29.4 mg/dL at 30, 60, 90, and 120 minutes after the exercise session, respectively.

185 Despite the clinical relevance of these findings, there is no consensus on the optimal
 186 combination of parameters for designing exercise programs for individuals with metabolic
 187 disorders. However, our study highlighted the importance of carefully planning rest intervals
 188 between each stimulus [20,21]. Our study highlights the importance of carefully planning rest
 189 intervals within HIIT protocols, as they significantly influence the metabolic and physiological
 190 outcomes. Longer rest periods may allow for higher exercise overload, while shorter rest
 191 periods may induce greater metabolic stress, impacting glucose metabolism and insulin
 192 sensitivity. This is important because when performing exercises with increased overload, the
 193 rest period needs to be adjusted to maintain the exercise, focusing the stimulus more on the

194 muscular system. Conversely, shorter rest periods suggest less overload but impact the
195 metabolism, as many myokines are overexpressed in this exercise approach [22,23].

196 Interleukin-6 (IL-6), a cytokine often elevated during high-intensity exercise, may play a role in
197 the metabolic adaptations observed in this study. IL-6 is known to regulate the expression of
198 glucagon-like peptide 1 (GLP-1), which has been shown to improve glucose control in diabetic
199 models [24]. Ellingsgaard et al., (2011), demonstrated that high-intensity exercise induces the
200 overexpression of IL-6, which upregulates GLP-1 in L cells and alpha cells, leading to improved
201 glucose control in diabetic mice. Therefore, we suggest that the overexpression of IL-6 during
202 HIIT may contribute to the improved glucose and insulin sensitivity observed in our participants.

203 **What does this article add and practical implications?**

204 This study highlights the potential of high-intensity interval training (HIIT) to acutely improve
205 glycemic control, making it a valuable exercise option for individuals looking to manage their
206 blood glucose levels more effectively. The findings suggest that shorter rest intervals in HIIT
207 protocols may enhance insulin sensitivity and reduce blood glucose concentrations more
208 significantly. These insights can assist fitness professionals and healthcare providers in
209 designing more effective exercise programs for individuals with or at risk of metabolic diseases
210 such as diabetes. Incorporating HIIT into regular exercise routines could provide a time-efficient
211 and effective strategy for improving overall metabolic health. Future research should explore the
212 role of IL-6 and other biomarkers in optimizing HIIT protocols further.

213

214

215 **Conclusion**

216 The manipulation of the inter-set rest period is crucial in promoting different metabolic
217 responses. Shorter recovery durations appear to affect metabolism more profoundly than longer
218 recovery periods, leading to a more significant acute impact on blood glucose levels and insulin
219 sensitivity in children with T1D. However, these findings should be interpreted with caution. The
220 combination of HIIT protocols, medication, and insulin therapy requires close monitoring due to
221 the potential risk of inducing a hypoglycemic state, which could be dangerous.

222

223 **Disclaimer of artificial intelligence**

224

225 The authors hereby declare that generative AI technology, CHAT GPT 4.0, was used for editing
226 the English language of the manuscript. The request was solely to correct spelling, punctuation,
227 and grammar, without altering, adding, omitting information or data, or changing the meaning of
228 the text.

229

230 **Acknowledgements**

231

232 The authors would like to thank to all participants, to SESI, and to Convivence Older Center in
233 Porto Velho, Rondônia, Brazil.

234

235 **Competing interests**

236

237 Authors have declared that no competing interests exist.

238

239 **Authors' contributions**

240

241 The authors JRV-S performed all steps of this manuscript. The all authors performed the
242 administrative tasks as supervision, control visualization and participated in the first wrote
243 version of this manuscript. JRV-S performed all stages, data curation and treatment, written all
244 versions and did all corrections, additionally, was the laboratorial supervisor. Both did the final
245 version and JRV-S did the submission.

246 **Ethical approval and Consent**

247

248 The study adhered to all mandatory ethical requirements as per the Brazilian National Health Council
249 Ethics Law 466/2012 and received approval from the Ethics Committee of the Federal University of
250 Rondônia (approval report number 2066823). All participants and their parents signed informed
251 consent forms after being informed of the study's risks, benefits, and procedures. Participation was
252 voluntary, and subjects could withdraw at any time without any penalties.

253

254

References

- 255 1. García-Hermoso, A.; Ezzatvar, Y.; Huerta-Urbe, N.; Alonso-Martínez,
256 A.M.; Chueca-Guindulain, M.J.; Berrade-Zubiri, S.; Izquierdo, M.; Ramírez-
257 Vélez, R. Effects of Exercise Training on Glycaemic Control in Youths with Type
258 1 Diabetes: A Systematic Review and Meta-Analysis of Randomised Controlled
259 Trials. *Eur J Sport Sci* 2023, 23, 1056–1067.
- 260 2. Gurka, M.J.; Guo, Y.; Filipp, S.L.; Deboer, M.D. Metabolic Syndrome
261 Severity Is Significantly Associated with Future Coronary Heart Disease in Type
262 2 Diabetes. *CardiovascDiabetol* 2018, 1–9, doi:10.1186/s12933-017-0647-y.
- 263 3. Stanford, K.I.; Goodyear, L.J. Exercise and Type 2 Diabetes: Molecular
264 Mechanisms Regulating Glucose Uptake in Skeletal Muscle.
265 *AdvPhysiolEduc* 2014, doi:10.1152/advan.00080.2014.
- 266 4. Ahmad, A.M.; Mahmoud, A.M.; Serry, Z.H.; Mohamed, M.M.;
267 AbdElghaffar, H.A. Effects of Low-versus High-Volume High-Intensity Interval
268 Training on Glycemic Control and Quality of Life in Obese Women with Type 2
269 Diabetes. A Randomized Controlled Trial. *J ExercSci Fit* 2023, 21, 395,
270 doi:10.1016/J.JESF.2023.08.003.
- 271 5. Suzuki, K. Chronic Inflammation as an Immunological Abnormality and
272 Effectiveness of Exercise. *Biomolecules* 2019, doi:10.3390/biom9060223.
- 273 6. Ramos, J.S.; Dalleck, L.C.; Keith, C.E.; Fennell, M.; Lee, Z.; Drummond,
274 C.; Keating, S.E.; Fassett, R.G.; Coombes, J.S. Optimizing the Interaction of
275 Exercise Volume and Metformin to Induce a Clinically Significant Reduction in
276 Metabolic Syndrome Severity: A Randomised Trial. *Int J Environ Res Public*
277 *Health* 2020, doi:10.3390/ijerph17103695.
- 278 7. Da Silva, M.A.R.; Baptista, L.C.; Neves, R.S.; De França, E.; Loureiro,
279 H.; Lira, F.S.; Caperuto, E.C.; Veríssimo, M.T.; Martins, R.A. The Effects of
280 Concurrent Training Combining Both Resistance Exercise and High-Intensity
281 Interval Training or Moderate-Intensity Continuous Training on Metabolic
282 Syndrome. *Front Physiol* 2020, 11, 1–10, doi:10.3389/fphys.2020.00572.
- 283 8. Portela, P.F. de M.; Neto, V.G.C.; Monteiro, E.R.; Santos da Silva, R.; da
284 Silva, V.F.; Nogueira, C.J.; Schutz, S.; Scudese, E.; Salvino, A.K.S.; Valentim-
285 Silva, J.R. HIIT Is Most Effective than MICT on Glycemic Control of Older People
286 with Glucose Metabolism Impairments: A Systematic Review and Metanalysis.
287 *Prim Care Diabetes* 2023, 17.
- 288 9. Sun, S.; Zhang, H.; Kong, Z.; Shi, Q.; Tong, T.K.; Nie, J. Twelve Weeks
289 of Low Volume Sprint Interval Training Improves Cardio-Metabolic Health
290 Outcomes in Overweight Females. *J Sports Sci* 2019,
291 doi:10.1080/02640414.2018.1554615.
- 292 10. Júnior, A.L.A.L.; da Silva, J.M.J.M.; da Silva, V.F.V.F.; Castro,
293 A.C.M.A.C.M.; de Freitas, R.E.R.E.; Cavalcante, J.B.J.B.; Dos Santos,
294 K.M.K.M.; Albuquerque, A.P.A.A.P.A.; Brandão, P.P.P.P.; Bello, M.N.D.M. de

- 295 N.D.; et al. Multimodal HIIT Is More Efficient than Moderate Continuous Training
296 for Management of Body Composition, Lipid Profile and Glucose Metabolism in
297 the Diabetic Elderly. *Int. J. Morpho***2020**, *38*, 392–399, doi:10.4067/S0717-
298 95022020000200392.
- 299 11. Cockcroft, E.J.; Moudiotis, C.; Kitchen, J.; Bond, B.; Williams, C.A.;
300 Barker, A.R. High-Intensity Interval Exercise and Glycemic Control in
301 Adolescents with Type One Diabetes Mellitus: A Case Study. *Physiol Rep***2017**,
302 *5*, doi:10.14814/phy2.13339.
- 303 12. Yardley, J.E. Fasting May Alter Blood Glucose Responses to High-
304 Intensity Interval Exercise in Adults With Type 1 Diabetes: A Randomized, Acute
305 Crossover Study. *Can J Diabetes***2020**, *44*, 727–733,
306 doi:10.1016/j.cjcd.2020.09.007.
- 307 13. Dubl, M.C.; Lavoie, C.; John Weisnagel, S. Glucose or Intermittent High-
308 Intensity Exercise in Glargine/Glulisine Users with T1DM. *Med Sci Sports*
309 *Exerc***2013**, *45*, 3–7, doi:10.1249/MSS.0b013e31826c6ad3.
- 310 14. Riddell, M.C.; Pooni, R.; Yavelberg, L.; Li, Z.; Kollman, C.; Brown, R.E.;
311 Li, A.; Aronson, R. Reproducibility in the Cardiometabolic Responses to High-
312 Intensity Interval Exercise in Adults with Type 1 Diabetes. *Diabetes Res*
313 *Clin Pract***2019**, *148*, 137–143, doi:10.1016/j.diabres.2019.01.003.
- 314 15. Bally, L.; Zueger, T.; Buehler, T.; Dokumaci, A.S.; Speck, C.; Pasi, N.;
315 Ciller, C.; Paganini, D.; Feller, K.; Loher, H.; et al. Metabolic and Hormonal
316 Response to Intermittent High-Intensity and Continuous Moderate Intensity
317 Exercise in Individuals with Type 1 Diabetes: A Randomised Crossover Study.
318 *Diabetologia***2016**, *59*, 776–784, doi:10.1007/s00125-015-3854-7.
- 319 16. Guerrero-Romero, F.; Simental-Mendía, L.E.; González-Ortiz, M.;
320 Martínez-Abundis, E.; Ramos-Zavala, M.G.; Hernández-González, S.O.;
321 Jacques-Camarena, O.; Rodríguez-Morán, M. The Product of Triglycerides and
322 Glucose, a Simple Measure of Insulin Sensitivity. Comparison with the
323 Euglycemic-Hyperinsulinemic Clamp. *Journal of Clinical Endocrinology and*
324 *Metabolism***2010**, doi:10.1210/jc.2010-0288.
- 325 17. Gillen, J.B.; Martin, B.J.; MacInnis, M.J.; Skelly, L.E.; Tarnopolsky, M.A.;
326 Gibala, M.J. Twelve Weeks of Sprint Interval Training Improves Indices of
327 Cardiometabolic Health Similar to Traditional Endurance Training despite a Five-
328 Fold Lower Exercise Volume and Time Commitment. *PLoS One***2016**, *11*,
329 doi:10.1371/journal.pone.0154075.
- 330 18. Holmstrup, M.; Fairchild, T.; Keslacy, S.; Weinstock, R.; Kanaley, J.
331 Multiple Short Bouts of Exercise over 12-h Period Reduce Glucose Excursions
332 More than an Energy-Matched Single Bout of Exercise. *Metabolism***2014**, *63*,
333 510–519, doi:10.1016/j.metabol.2013.12.006.
- 334 19. De Teles, G.O.; Gentil, P.; e Silva, L.R.B.; De Sousa, W.M.; Seguro,
335 C.S.; Rebelo, A.C.S. HIIE Protocols Promote Better Acute Effects on Blood
336 Glucose and Pressure Control in People with Type 2 Diabetes than Continuous
337 Exercise. *Int J Environ Res Public Health***2022**, *19*, doi:10.3390/ijerph19052601.
- 338 20. Braga, J.C.; de Freitas, R.E.; dos Santos, K.M.; Pontes da Silva, R.;
339 Mota da Silva, J.; Junior, A.L.; de Oliveira, G.L.; Oliveir, T.A.P.; Pernambuco,
340 C.S.; Furtado da Silva, V.; et al. Twelve Weeks of High-Intense Interval Training

- 341 Enhance the Neuromuscular and Cardiorespiratory Performance of Elderly.
342 *Open Sports Sci J***2020**, *13*, 42–48, doi:10.2174/1875399x02013010042.
- 343 21. Louzada Júnior, A.; Mota da Silva, J.; Furtado da Silva, V.; Clodoaldo
344 Melo Castro, A.; Eufrásio de Freitas, R.; Braga Cavalcante, J.; Maia dos Santos,
345 K.; Paula Azevedo Albuquerque, A.; Paraguassú Brandão, P.; de Nazaré Dias
346 Bello, M.; et al. Multimodal HIIT Is More Efficient Than Moderate Continuous
347 Training for Management of Body Composition, Lipid Profile and Glucose
348 Metabolism in the Diabetic Elderly. *Int. J. Morphol***2020**, *38*, 392–399.
- 349 22. Wiecek, M.; Szymura, J.; Maciejczyk, M.; Kantorowicz, M.; Szygula, Z.
350 Acute Anaerobic Exercise Affects the Secretion of Asprosin, Irisin, and Other
351 Cytokines - A Comparison between Sexes. *Front Physiol***2018**, *9*, 1782,
352 doi:10.3389/fphys.2018.01782.
- 353 23. Moldoveanu, A.I.; Shephard, R.J.; Shek, P.N. Exercise Elevates Plasma
354 Levels but Not Gene Expression of IL-1beta, IL-6, and TNF-Alpha in Blood
355 Mononuclear Cells. *J Appl Physiol (1985)***2000**, *89*, 1499–1504, doi:8750-
356 7587/00.
- 357 24. Ellingsgaard, H.; Seelig, E.; Timper, K.; Coslovsky, M.; Soederlund, L.;
358 Lyngbaek, M.P.; WewerAlbrechtsen, N.J.; Schmidt-Trucksäss, A.; Hanssen, H.;
359 Frey, W.O.; et al. GLP-1 Secretion Is Regulated by IL-6 Signalling: A
360 Randomised, Placebo-Controlled Study. *Diabetologia***2020**, *63*, 362–373,
361 doi:10.1007/s00125-019-05045-y.
- 362 25. Ellingsgaard, H.; Hauselmann, I.; Schuler, B.; Habib, A.M.; Baggio, L.L.;
363 Meier, D.T.; Eppler, E.; Bouzakri, K.; Wueest, S.; Muller, Y.D.; et al. Interleukin-6
364 Enhances Insulin Secretion by Increasing Glucagon-like Peptide-1 Secretion
365 from L Cells and Alpha Cells. *Nat Med***2011**, *17*, 1481–1489,
366 doi:10.1038/nm.2513.
- 367