

University of Dundee

Exercise management in type 1 diabetes

Riddell, Michael C.; Gallen, Ian W.; Smart, Carmel E.; Taplin, Craig E.; Adolfsson, Peter; Lumb, Alistair N.

Published in:
The Lancet: Diabetes and Endocrinology

DOI:
[10.1016/S2213-8587\(17\)30014-1](https://doi.org/10.1016/S2213-8587(17)30014-1)

Publication date:
2017

Document Version
Peer reviewed version

[Link to publication in Discovery Research Portal](#)

Citation for published version (APA):

Riddell, M. C., Gallen, I. W., Smart, C. E., Taplin, C. E., Adolfsson, P., Lumb, A. N., Kowalski, A., Rabasa-Lhoret, R., McCrimmon, R. J., Hume, C., Annan, F., Fournier, P. A., Graham, C., Bode, B., Galassetti, P., Jones, T. W., Millán, I. S., Heise, T., Peters, A. L., ... Laffel, L. M. (2017). Exercise management in type 1 diabetes: a consensus statement. *The Lancet: Diabetes and Endocrinology*, 5(5), 377-390. [https://doi.org/10.1016/S2213-8587\(17\)30014-1](https://doi.org/10.1016/S2213-8587(17)30014-1)

General rights

Copyright and moral rights for the publications made accessible in Discovery Research Portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from Discovery Research Portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain.
- You may freely distribute the URL identifying the publication in the public portal.

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

1 **Exercise management in type 1 diabetes: a consensus statement**

2 Michael C. Riddell¹, Ian W. Gallen², Carmel E. Smart³, Craig E. Taplin⁴, Peter Adolfsson⁵, Alistair N. Lumb⁶, Aaron
3 Kowalski⁷, Remi Rabasa-Lhoret⁸, Rory McCrimmon⁹, Carin Hume¹⁰, Francesca Annan¹¹, Paul A. Fournier¹², Claudia
4 Graham¹³, Bruce Bode¹⁴, Pietro Galassetti¹⁵, Timothy W. Jones¹⁶, Inigo San Millan¹⁷, Tim Heise¹⁸, Anne Peters¹⁹,
5 Andreas Petz²⁰, and Lori M. Laffel²¹.

6 **Running title: Exercise management in T1D**

7 ¹Muscle Health Research Centre, York University, Toronto, Ontario, Canada
8 ²Royal Berkshire NHS Foundation Trust Centre for Diabetes and Endocrinology, Royal Berkshire Hospital, Reading, UK
9 ³Hunter Medical Research Institute, School of Medicine and Public Health, University of Newcastle, Rankin Park, NSW, Australia;
10 Department of Paediatric Diabetes and Endocrinology, John Hunter Children's Hospital, Newcastle, NSW, Australia
11 ⁴Division of Endocrinology and Diabetes, Department of Pediatrics, University of Washington, Seattle Children's Hospital,
12 Seattle, Washington, USA
13 ⁵Department of Pediatrics, The Hospital of Halland, Kungsbacka, Institute of Clinical Sciences, The Sahlgrenska Academy at
14 University of Gothenburg, Gothenburg, Sweden
15 ⁶Oxford Centre for Diabetes, Endocrinology and Metabolism (OCDEM), Churchill Hospital, Oxford, UK
16 ⁷JDRF, New York, NY, USA
17 ⁸Department of Nutrition & Institut de Recherches Cliniques de Montréal, Faculty of Medicine, Université de Montréal,
18 Montreal, Quebec, Canada
19 ⁹Department of Cardiovascular and Diabetes Medicine, University of Dundee, Dundee, UK.
20 ¹⁰London Medical, London, UK
21 ¹¹Children and Young People's Diabetes Service, University College London Hospitals NHS Foundation Trust, London, UK
22 ¹²School of Sport Science, Exercise, and Health. The University of Western Australia, Perth, Western Australia 6008, Australia
23 ¹³Dexcom, Inc, San Diego, CA, USA
24 ¹⁴Atlanta Diabetes Associates, Atlanta, GA, USA
25 ¹⁵Department of Pediatrics, University of California Irvine, Irvine, California, USA
26 ¹⁶Department of Endocrinology and Diabetes, Perth Children's Hospital and Telethon Kids Institute, The University of Western
27 Australia, Perth, Australia
28 ¹⁷Department of Physical Medicine & Rehabilitation, University of Colorado, School of Medicine, Aurora, CO, USA
29 ¹⁸Profil, Neuss, Germany
30 ¹⁹Keck School of Medicine, University of Southern California, Los Angeles, CA, USA
31 ²⁰StubbaekSkole, Aabenraa, Denmark
32 ²¹Division of Endocrinology, Boston Children's Hospital, Boston, MA Pediatric, Adolescent and Young Adult Section, Joslin
33 Diabetes Center, Boston, MA.
34

35 **Corresponding author:**

36 Michael C. Riddell, PhD
37 347 Bethune College, 4700 Keele Street
38 Toronto, Ontario, M3J 1P3, Canada
39 Telephone: (416) 736-2100 ext. 40493
40 Email: mriddell@yorku.ca
41

42 Word count abstract: 380
43 Word count main text: 5069
44 References: 119
45
46

47 **Abstract**

48 Type 1 diabetes (T1D) is challenging condition to manage for a variety of physiological and behavioural
49 reasons. Regular exercise is important, however management of the different forms of activity is a
50 particular struggle for both the individual with T1D and the health care provider. People with T1D tend
51 to be at least as inactive as the general population, with a large percentage of individuals not
52 maintaining a healthy body mass nor achieving the minimum number of minutes per week of moderate-
53 to-vigorous aerobic activity. Regular exercise can improve health and well-being and can help individuals
54 to achieve their lipid, body composition, fitness and glycaemic goals. However, several additional
55 barriers to exercise may exist for the person with diabetes including fear of hypoglycaemia, loss of
56 glycaemic control, and inadequate knowledge around exercise management. This review provides an up
57 to date consensus on exercise management for individuals with T1D who exercise regularly, including
58 glucose targets for safe and effective exercise, and nutritional and insulin dose adjustments to protect
59 against exercise-related glucose excursions.

60 Introduction

61 Despite tremendous advances since the discovery of insulin almost 100 years ago, type 1
62 diabetes (T1D) remains a challenging disease to manage (1,2). A majority of patients living with T1D are
63 not at a healthy body weight (~60% are overweight or obese), suffer from hypertension (~40%) and/or
64 dyslipidaemia (~60%) (3) and are not engaging enough regular physical activity (4). Regular exercise
65 helps patient achieve a number of goals. It improves the cardiovascular disease risk profile in paediatric
66 patients (5) and reduces HbA1c (-0.3%) in that particular segment of the patient population (6). Body
67 composition, cardiorespiratory fitness, endothelial function and blood lipid profile (i.e. triglycerides,
68 total cholesterol) all improve with regular physical activity in children and young people with T1D (7).
69 These cardiometabolic improvements are all important, given that cardiovascular disease is the leading
70 cause of morbidity and mortality in persons with T1D (8). In adults, both retinopathy and
71 microalbuminuria are less common in those who are more physically active (9). Active adults with T1D
72 tend to have better chance of achieving target HbA1c levels, blood pressure levels and a healthier body
73 mass index when compared to inactive patients (3). Regular exercise also lowers total daily insulin needs
74 (10). Having a high exercise capacity in adulthood with T1D is associated with less risk for coronary
75 artery disease, myocardial ischaemia and stroke if you have diabetes or not (11). In a large cross
76 sectional study of 18,028 adults with T1D, patients who fall in the most active category of physical
77 activity levels (exercising two or more times per week) had better HbA1c levels, a more favourable body
78 mass index, less dyslipidaemia, hypertension and fewer diabetes-related complications (retinopathy,
79 microalbuminuria), compared to those who were less habitually active (3). In general, patients with T1D
80 who are more active tend to have less diabetic ketoacidosis and less risk for developing severe
81 hypoglycaemia with coma (3), except for with older women where this latter relationship is reversed;
82 those most active have higher rates of severe hypoglycaemia (with coma) when compared with those
83 who are inactive (3). However, several barriers may exist for exercise including: a fear of hypoglycaemia;
84 a loss of glycaemic control, lack of time; access to facilities; lack of motivation; issues around body image
85 and a general lack of knowledge around exercise management (12–14).

86 The physical activity prescription for all adults living with diabetes, including those living with
87 T1D, is 150 minutes of accumulated physical activity each week, with no more than 2 days in a row with
88 no activity. Resistance exercise is also recommended two to three times a week. Getting this much
89 exercise is difficult for a large majority of patients; with less than 20% of patients performing aerobic
90 exercise more than two times per week and 60% of the patient population performing no structured
91 exercise at all (3). For children and adolescent, at least 60 minutes of physical activity should be
92 performed per day (15). Physical inactivity and prolonged sitting times increase gradually with age and
93 are linked to high HbA1c levels in youth with T1D (16) and inactivity appears to be more common in
94 females than in males (3).

95 Regular exercise should be encouraged and supported by health care professionals (HCPs), for a
96 number of reasons, but primarily because the overall cardiometabolic benefits outweigh the immediate
97 risks if certain precautions are made. In this review, the basic categories of exercise are described from a
98 physiological perspective as are the starting points for nutritional and insulin dose adjustments to keep
99 patients in a targeted glycaemic range. This review summarizes our consensus on the available

100 strategies that help incorporate exercise safely into the daily T1D management plan for those adults
101 who are regularly engaging in exercise, sport and/or competitive events. It is hoped that these new
102 guidelines for exercise management will improve patient control and engage more individuals with T1D
103 to be more physically active.

104

105 **Search strategy and selection criteria**

106 We searched PubMed.gov and other relevant biomedical databases for articles pertaining to
107 'type 1 diabetes' OR 'insulin-dependent diabetes AND 'exercise' OR 'physical activity'; published
108 between 01/1990 to 07/2016 and filtered for human and restricted to English publications. Additional
109 searches using the search terms 'nutrition' OR 'dietary carbohydrate' OR 'dietary protein' OR 'glycemic
110 index' OR 'hypoglycaemia' OR 'energy expenditure OR 'glycemic control' OR 'management' OR
111 'hypoglycemia' OR 'hyperglycemia' OR 'prevention & control' were conducted for various subtopics
112 within this review.

113

114 **Physiology of Physical Activity and Exercise**

115 Modalities of exercise

116 Understanding the metabolic and neuroendocrine responses to the various types of exercise undertaken
117 by people with T1D is critical for determining appropriate nutritional and insulin management
118 strategies. Exercise is generally classified as *aerobic* or *anaerobic*, depending on the predominant energy
119 systems used to support the activity, although a majority of exercise activities include a mix of energy
120 systems. Aerobic exercise involves repeated and continuous movement of large muscle groups (e.g.
121 walking, cycling, jogging, and swimming) that rely primarily on aerobic energy-producing systems.
122 Resistance (strength) training is a type of exercise using free weights, weight machines, body weight, or
123 elastic resistance bands that rely primarily on anaerobic energy-producing systems. High intensity
124 interval training (HIIT), involves alternating between brief periods of vigorous exercise and recovery
125 periods at low-to-moderate intensity (e.g., from 20 seconds to 4 minute intervals of exercise and rest,
126 for up to ~10 cycles) (17). Both aerobic and resistance type activities are recommended for a majority of
127 people living with diabetes (15,18) and recent guidelines also now incorporate HIIT as a training
128 modality with established benefits for individuals with prediabetes or type 2 diabetes (18). In some
129 studies, HIIT has been shown to be more effective than continuous aerobic training in improving
130 cardiovascular fitness and various parameters related to glucose metabolism including insulin sensitivity
131 and glycaemic control in type 2 diabetes (19). At present, it is unclear what form(s) of exercise are best
132 for improving cardiometabolic control in type 1 diabetes (20).

133 Neuroendocrine and metabolic responses to exercise

134 *Individuals without diabetes*

135 The metabolic responses to different forms of exercise are distinct. However, in almost all forms of
136 exercise, no matter the intensity or duration, blood glucose concentrations are normally held within a
137 tight range (4-6 mmol/l). During aerobic exercise, insulin secretion drops and glucagon secretion rises in
138 the portal vein to facilitate glucose release from the liver to match the rate of glucose uptake into the
139 working muscles (21). Exercise can increase glucose uptake into muscle by up to 50-fold; a phenomenon
140 independent of insulin signalling (22), so the drop in insulin in the circulation does not limit glucose
141 provision to the working body. Although the main determinant of glucose production for aerobic
142 exercise is a rise in glucagon levels, there is also neural control of glucose release and other
143 counterregulatory hormones play a supportive role (23). With increased exercise duration, there is
144 reduced reliance on muscle glycogen as fuel and a greater reliance on lipid oxidation and plasma-derived
145 glucose (24). If insulin levels do not fall during aerobic exercise, the rise in counterregulatory hormones
146 is less effective in promoting hepatic glucose production (21).

147 As the intensity of exercise increases above ~50-60% of maximal oxygen consumption (VO_2max),
148 fat oxidation decreases, particularly in those who are untrained, and CHO are the preferred fuel (25).
149 Prolonged high-intensity exercise is supported by both muscle glycogen and blood glucose utilization
150 with minimal contributions from lipid and protein (26). During predominantly anaerobic activities (27)
151 and during a HIIT session (28), circulating insulin concentrations do not drop as markedly as compared to
152 purely aerobic activities, in part because the duration of activity is typically shorter. High rates of
153 external power output during HIIT increase reliance on muscle phosphagens and glycogen, with lactate
154 levels rising markedly in the circulation (28). Insulin levels increase above baseline levels in early
155 recovery from a HIIT session to offset the rise in glucose caused by the elevations in counterregulatory
156 hormones and other metabolites (27).

157

158 *Dysglycaemia during exercise in individuals with T1D*

159 In T1D, the glycaemic responses to exercise are influenced by the location of insulin delivery, the
160 amount of insulin in the circulation, the pre exercise blood glucose concentration, the composition of
161 the last meal or snack, as well as the intensity and duration of the activity (29) (Figure 1).

162 During aerobic exercise, most individuals with T1D have a drop in glycaemia, unless
163 carbohydrates (CHO) are ingested, because insulin levels cannot be lowered rapidly enough at the start
164 of the activity, and levels may rise in the systemic circulation (30), perhaps because of increased
165 subcutaneous adipose tissue blood flow during exercise (31). Even if basal insulin infusion rates are
166 halved 60-min before the start of exercise in patients on continuous subcutaneous insulin infusion (CSII),
167 circulating free insulin levels do not drop at exercise commencement and levels tend to rise transiently
168 during the activity (32). Higher insulin levels in circulation during exercise promotes increased glucose
169 disposal relative to hepatic glucose production, and may delay lipolysis, another feature that increases
170 the muscles reliance on glucose as a fuel. Hypoglycaemia develops in a majority of patients within ~45
171 minutes of activity (33,34). Trained individuals with T1D have greater reductions in blood glucose
172 concentrations during aerobic exercise when compared to less fit patients (35), possibly because the

173 overall work rate is higher in those more aerobically conditioned. As such, both trained and untrained
174 individuals with T1D typically require increased CHO intake, and/or insulin dose reduction, for prolonged
175 aerobic exercise (see below). High intensity interval sprint training promotes increased oxidative
176 capacity of skeletal muscle in T1D and attenuates the rates of glycogen breakdown (36), which may
177 protect against post-exercise hypoglycaemia, at least in theory. Perhaps in line with this, individuals who
178 are aerobically conditioned have reduced glucose variability compared to those unconditioned (37). Low
179 insulin levels due to aggressive reductions in administration or a skipped insulin dose can cause
180 hyperglycaemia prior to and during aerobic exercise (38) and ketosis may develop, even with mild
181 activity (39).

182 Compared to continuous moderate-intensity aerobic exercise, resistance exercise is associated
183 with better glucose stability (40), although it may cause a modest rise in some individuals (41).
184 Compared to aerobic exercise, performing a HIIT session attenuates the drop in glycaemia (42), as does
185 performing resistance exercise before aerobic exercise (43), possibly because of increases in
186 counterregulatory hormones and various metabolites that limit glucose disposal (44). In situations of
187 brief and intense anaerobic exercise (e.g. sprinting, weight lifting, some competitive sports) (41,45), or
188 during HIIT (28), glucose levels typically rise.

189

190 *Dysglycaemia post-exercise in individuals with T1D*

191 Immediately after aerobic exercise, glucose uptake into muscle drops but overall glucose disposal
192 is still elevated for hours in recovery to help replenish glycogen stores (46). Hypoglycaemia risk is
193 elevated for at least 24 hours in recovery from exercise with the greatest risk for nocturnal
194 hypoglycaemia occurring after afternoon activity (47). As mentioned above, weight lifting, sprinting and
195 intense aerobic exercise can promote elevations in glycaemia that may last for hours in recovery.
196 Although a conservative insulin “correction” post-exercise may be prudent in some situations (48), over-
197 correction with insulin can promote severe nocturnal hypoglycaemia and death (49). HIIT appears to
198 increase risk for nocturnal hypoglycaemia compared to continuous aerobic exercise in some (50), but
199 not all (51,52) studies.

200

201 **Exercise Goals and Glycaemic Targets**

202 Individuals with T1D should perform exercise for a variety of health reasons. The evidence that
203 regular exercise training improves metabolic control in adults with T1D is somewhat limited (20,53),
204 although in youth it appears to be helpful (7). Exercise readiness questionnaires for adults with diabetes
205 who may be at increased risk for adverse events can be found at eparmedx.com. Patient goals for
206 exercise should be considered before making management decisions (e.g. metabolic control and
207 prevention of complications, fitness, weight loss, competition/performance). This is a critical element of
208 the diabetes management plan. For example, exercise for weight loss requires strategies that focus on
209 reducing insulin levels during and after exercise, as opposed to consuming additional CHOs. By contrast,

210 if maximising sports and exercise performance is the primary goal, then sport-specific nutritional
211 guidance is needed and a modified insulin plan to match increased nutritional requirements should be
212 considered (54). For all patients, blood glucose monitoring before, during and after exercise is critical for
213 informing strategies and maintaining stable and safe glycaemia.

214 The appropriate blood glucose concentration for the start of exercise should be individually
215 tailored. Based on consensus, a reasonable starting range for most patients doing aerobic exercise
216 lasting up to an hour is between 7-10 mmol/l. This range balances performance considerations against
217 hypoglycaemia risk. Higher levels may be acceptable in some situations where added protection against
218 hypoglycaemia is needed. Achieving and maintaining circulating glucose in this range is challenging. The
219 glycaemic response to exercise is variable and based on several factors including the duration and
220 intensity of exercise (44,55), the starting level of glycaemia (34), the individual's aerobic fitness (35) and
221 the amount of insulin in circulation (56,57) (Figure 1). Anaerobic and a HIIT session can be initiated with
222 a lower starting glucose level (5-7 mmol/l) since glucose concentrations tend to remain relatively stable,
223 fall to a lesser extent compared to continuous aerobic exercise, or rise slightly (Figure 1). Strategies to
224 cope with a range of glucose concentrations near the start of exercise are provided in Table 1. If glucose
225 level is too high because of insulin omission, risk of ketosis and further hyperglycaemia can occur (39)
226 and work effort probably rises. Although it is unclear if there is an optimal glycaemic range for exercise
227 performance, clinical experience and limited field study investigation (58) suggest that maintaining a
228 concentration between ~6.0-8.0 mmol/l may be ideal.

229

230 **Contraindications and cautions for exercise**

231 While few exercise limitations should be placed on patients, some considerations are important:

232 A. Ketones

233 Elevated blood ketones (≥ 1.5 mmol/l) before a bout of exercise should be addressed prior to
234 the start of the session via insulin administration and/or CHO feeding (Table 1). The cause of
235 elevated ketone levels should be identified (illness, diet manipulation, a recent bout of
236 prolonged exercise, insulin omission, etc.). Both prolonged endurance type activities
237 (marathons, trekking, etc.) and very low CHO diets can elevate blood ketone levels in
238 patients and the HCP should define appropriate levels and provide tailored guidance for
239 each individual. Blood ketone levels of ≥ 3.0 mmol/l should be managed immediately by a
240 qualified HCP (e.g. emergency department, physician, etc.).

241 B. Recent hypoglycaemia

242 Severe hypoglycaemia (defined here as a blood glucose ≤ 2.8 mmol/l or a hypoglycaemic
243 event requiring assistance from another individual) within the previous 24 hours is a
244 contraindication to exercise, due to the significant increased risk of a more serious episode
245 during the exercise (59). Where minor hypoglycaemia (blood glucose 2.9-3.9 mmol/l, with

246 the ability to self-treat) has occurred, the increased risk of a recurrence must be taken into
247 account (60). Vigilance around monitoring should be stressed and exercise should be
248 avoided if the setting is deemed particularly unsafe (e.g. swimming/trekking alone).

249 C. Diabetes-related complications

250 Overall the health benefits of being physically active outweigh the risks of being sedentary
251 for people with diabetes. Those with complications can derive numerous health benefits
252 from lower intensity physical activities, with little risk for any adverse events (61). In those
253 with long-standing disease, or with HbA1c levels well above target, vigorous exercise, heavy
254 weight-bearing activities and competitive endurance events are contraindicated, particularly
255 if the patient has unstable proliferative retinopathy, severe autonomic dysfunction or renal
256 failure (61).

257 D. Failure to be prepared for exercise-associated hypoglycaemia

258 In preparation for exercise, individuals with T1D should be aware of their starting glucose
259 concentrations, have blood glucose monitoring equipment and snacks to treat
260 hypoglycaemia. They should also be advised to wear/carry diabetes identification.

261

262 **Nutritional Management**

263 Goals for nutritional management

264 Nutritional management for people with T1D should incorporate strategies that optimise
265 glycaemic control, while promoting long-term health (62). The main strategies around nutrition for
266 exercise and sport discussed in this section are primarily to maximise athletic performance and are
267 based largely on studies conducted in highly trained healthy individuals without diabetes (63), with
268 limited studies in people with T1D. The application of these strategies must consider the individual's
269 insulin management plan and specific advice targeting nutrition for both athletic performance and
270 glycaemic management (see Glycaemic Management section below). A registered dietitian with
271 specialist diabetes and sports knowledge is the most qualified to help active people with T1D.

272 An individualised meal planning approach is central to improving performance and glycaemic
273 outcomes. Daily CHO intake should relate to the fuel cost of training in the athletic subpopulation and
274 hypoglycaemia prevention for all active people. Balancing insulin dose to CHO intake during exercise is
275 essential. A variety of CHO and insulin adjustment strategies can be used, such as reducing the pre-
276 exercise bolus insulin dose by 30-50% up to 90 minutes before aerobic exercise (64), consuming high
277 glycaemic index (GI) CHO (30-60g /hour) during sport or replacing CHO post-exercise for anaerobic
278 exercise. Personal tolerance of ingested CHO particularly during exercise is a key factor in individualising
279 recommendations. The distribution of macronutrient intake over the day should take into account the
280 timing of exercise so that liver and muscle glycogen stores are maximised before the activity and

281 replenished in early recovery (63). This strategy should include CHO feeding well before exercise (~4
282 hours) and early in recovery (63,65).

283

284 Daily energy and macronutrient balance

285 Athletes with T1D need sufficient energy to meet the demands of their daily activity. These will
286 vary with age, sex, body composition and activity type (66). Total energy requirements differ with
287 individual aims. Predictive equations can be used to estimate resting energy expenditure (67); however
288 they should serve only as a guide as they may over or underestimate actual requirements. An
289 appropriate macronutrient balance and micronutrient intake (63), coupled with a glycaemic control
290 strategy, is required to maximise performance. The optimal macronutrient distribution will vary
291 depending on an individualised assessment and exercise goals. A guide to the distribution of the total
292 daily energy intake is 45-65% CHO, 20-35% fat and 10-35% protein, with higher protein intakes indicated
293 for individuals wanting to lose weight (68).

294 The major nutrients required to fuel performance are CHOs and lipids, while the addition of
295 protein is needed to help foster recovery and maintain nitrogen balance (63,69). Protein requirements
296 range from 1.2- 1.6 g/kg body weight (BW) /day and will vary with training type and intensity and CHO
297 availability (63,70). Higher intakes may be needed for recovery from injury or for individuals on energy
298 restricted diets (71) to maintain lean body mass.

299

300 CHO needs before, during and after exercise

301 Distinction should be made between CHO needs for performance and CHO required for
302 hypoglycaemia prevention (Table 2). CHO requirements will alter insulin management strategies and
303 vice versa. The majority of studies in T1D investigate the amount and distribution of CHO to prevent
304 hypoglycaemia rather than to optimise performance, although the two may be at least partially related
305 (34,64,72,73). As an example, although only 15-20 grams/hr of CHO may be required to prevent
306 hypoglycaemia in people who reduce their insulin levels in anticipation of exercise; this amount of CHO
307 may be insufficient for performance. It has been shown that it is possible to implement larger CHO
308 supplementation (up to 75g/hr) for prolonged competition greater than 2.5 hours (marathons and other
309 endurance type races) without adversely impacting glycaemia as long as insulin dose is titrated
310 appropriately (54). In general, CHO requirements during shorter, intermittent high intensity and
311 anaerobic activities can be much less (Table 2).

312

313 Nutritional needs for recovery

314 Post-exercise nutrition requirements to maximise muscle recovery and muscle protein synthesis
315 have been well studied in the athletic population without diabetes (74). For replenishment of glycogen

316 content after exercise, CHO intake is essential (63). For athletes with T1D, it is important to ensure rapid
317 and adequate replenishment of muscle and liver glycogen stores to help prevent late-onset
318 hypoglycaemia. Glycogen replacement strategies may also be important to help prevent euglycaemic
319 ketosis in exercise recovery (75). Ingesting protein (~20-30 grams) in addition to CHO in the post-
320 exercise period is beneficial for muscle protein synthesis, but it does not appear to facilitate glycogen
321 replenishment, at least in non-diabetic athletes (63).

322

323 Role of high and low GI foods for maintenance of euglycaemia

324 The GI of a CHO-rich food can be used to assist with the selection of CHO type for exercise; with
325 high GI sports drinks and gels providing rapidly released CHO to increase blood glucose levels during
326 endurance events and for the treatment of hypoglycaemia. Low GI foods have been suggested pre-
327 exercise to sustain CHO availability and maintain euglycaemia, while higher GI meals/snacks consumed
328 post-exercise may enhance recovery. Low and moderate GI snacks may also be preferred for long
329 distance activities (like trekking and long distance cycling) at low to moderate workloads. Low GI CHO
330 (isomaltose) consumed 2 hours before a high intensity run showed improved blood glucose responses
331 during exercise compared to a high GI CHO (dextrose) (76). A low GI meal and bedtime snack consumed
332 after evening exercise prevented postprandial hyperglycaemia compared to a high GI meal and snack,
333 with both meal types protective against hypoglycaemia for ~8 hours (77). Protection beyond 8 hours
334 with a snack is lost and hypoglycaemia risk remains significant (77).

335

336 Fluid Replacement

337 Adequate fluid intake before, during and after exercise is necessary to avoid dehydration and for
338 optimal performance (65). Water is the most effective drink for low intensity and short duration sports
339 (i.e. \leq ~45 min), as long as glucose levels are at or above target (\geq 7 mmol/l). Sports beverages containing
340 CHO (6-8%) and electrolytes are useful for athletes with T1D in longer duration, higher intensity exercise
341 as a hydration and fuel source and to prevent hypoglycaemia (34,78). However, it is important to ensure
342 these are not over consumed as this can result in hyperglycaemia. Milk-based drinks containing CHO and
343 protein can assist recovery and prevent delayed hypoglycaemia (73).

344

345 Low-CHO high-fat diets and exercise

346 People with T1D may choose a low-CHO high-fat (LCHF) diet for a variety of reasons. A recent
347 review on LCHF diets and sports performance in subjects without T1D concluded that despite increasing
348 the muscles' ability to utilise fat over time, there was no evidence of performance benefits (79). Long-
349 term studies have yet to be conducted on the health, glycaemia, or performance effects of LCHF diets in
350 T1D. A concern with these diets is that they may impair the capacity for high intensity exercise (80).

351 Variation in CHO intake (i.e. periodisation throughout the training cycle according to fuel needs
352 and performance) has been suggested by some researchers as a way to help promote skeletal muscle
353 adaptation to training (81). Additionally, various exercise-nutrient protocols (i.e. training in a fasted
354 state or withholding CHO intake at meal before or after exercise) are used to manipulate CHO
355 availability. These approaches have not been studied in individuals with T1D where manipulation of
356 dietary CHO as part of training presents unique challenges for insulin therapy and requires careful
357 glucose monitoring.

358

359 Sports nutritional aids and T1D

360 The use of ergogenic aids is a widespread performance enhancement strategy used by athletes.
361 The evidence for ergogenic aids on performance is limited in athletes with T1D.

362 Caffeine intake in athletes without diabetes has shown improvements in endurance capacity
363 and power output (82). Caffeine intake (5-6 mg/kg body mass) before exercise attenuates the drop in
364 glycaemia during exercise in individuals with T1D but may increase late-onset hypoglycaemia risk (83).

365

366 **Glycaemic Management Recommendations**

367 There is high between- and within-patient variability in glucose responses to the various forms
368 and intensities of exercise (Figure 1); therefore glycaemic management is based on frequent glucose
369 monitoring, adjustments to both basal and/or bolus insulin dosing and the consumption of CHOs during
370 and after exercise. These recommendations are intended to serve as a starting point for insulin
371 adjustments and CHO intake that can then be individualised (Figure 2).

372 Clinical management strategies should be built around exercise types and individual aims and
373 implemented, taking into account the factors summarised in Table 3. Generally, sustained aerobic
374 exercise requires more substantial reductions in insulin dose and/or higher CHOs than a shorter-term
375 HIIT session. In stark contrast, brief anaerobic exercise (sprinting, weight lifting) may require increased
376 insulin delivery, which is typically given in early recovery rather than before exercise for obvious safety
377 reasons (48). Strategies for insulin dose adjustments and/or CHO intake during and after planned
378 exercise are presented in Table 4.

379

380 Insulin adjustment for prolonged activities: bolus insulin approaches

381 Pre-exercise meal insulin bolus dose reductions and/or additional CHO consumed during
382 exercise are typically needed to avoid hypoglycaemia during prolonged exercise (>30 minutes)
383 (34,55,64,84–86). Bolus dose reductions require pre-planning and are probably only appropriate for
384 exercise with a predictable intensity performed within 2-3 hours after a meal. As shown in Table 5, the

385 extent of mealtime dose reduction is proportional to both the intensity and duration of the activity. This
386 approach is safe and effective; even reducing the bolus insulin dose by as much as 75% does not appear
387 to increase ketone production during exercise (86).

388 Another strategy is to combine the reduction of the pre-exercise insulin bolus dose (by 75%)
389 with the ingestion of a low GI snack/meal (87). Importantly, this method also reduces the risk of pre-
390 exercise hyperglycaemia. However, protection against hypoglycaemia with this approach is lost if the
391 exercise is performed an hour or more after the snack (87). As such, this combined approach may be
392 preferable only for early postprandial exercise.

393

394 Basal insulin approaches

395 Late postprandial hypoglycaemia (4+ hours after a meal) following aerobic exercise is driven
396 partly by circulating basal insulin concentrations. Elevated insulin sensitivity post-exercise, and perhaps
397 a blunting of glucose counterregulation appear to place individuals at risk for at least 12 hours. Reducing
398 circulating basal insulin levels can ameliorate this risk. For patients on multiple daily insulin injections
399 (MDI), clinical observations and limited experimental data (88) demonstrate that reducing long acting
400 basal (as well as prandial) insulin before exercise reduces hypoglycaemia risk during and after the
401 activity, but may promote hyperglycaemia at other points during the day. Therefore reduction in basal
402 insulin dose for MDI patients should not be routinely recommended but may be a therapeutic option for
403 those having unusual days with considerably more planned activity (e.g. camps, tournaments). In
404 general, basal insulins with a relatively short half-life such as NPH-insulin or insulin detemir seem to lead
405 to less hypoglycaemia in conjunction with exercise when compared to longer basal insulins such as
406 glargine (89), although the mechanism for this is unclear. While ultra-long acting insulins (e.g. insulin
407 degludec with a 25hr half-life) pose similar risks for hypoglycaemia with endurance exercise to that of
408 insulin glargine (90), dose reductions for exercise would have to be implemented at least 48 hours
409 before planned exercise. This is not recommended, as it would compromise overall control.

410 CSII offers flexibility to modify basal infusion delivery and to obtain a relatively quick effect
411 (within ~1-2hrs) (91). Suspension of basal insulin infusion at the onset of 60-min exercise reduces
412 hypoglycaemia risk during the activity, but it may increase post-exercise hyperglycaemia risk (92).
413 Moreover, glucose levels may still drop 2-3 mmol/l over 30-60 minutes even when basal insulin is
414 dramatically reduced (or completely suspended) (64,92,93), due to the lag time in the change in
415 circulating insulin levels. Where practical, a basal rate reduction, rather than suspension, should be
416 attempted well before the start of exercise (60-90 minutes). An 80% basal reduction at the onset of
417 exercise helps mitigate post-exercise hyperglycaemia, compared to basal suspension, and appears to be
418 associated with reduced hypoglycaemic risk both during and after the activity (64). However, the
419 optimal timing of basal rate insulin reductions for aerobic and HIE activities and the maximal safe
420 duration for insulin pump suspension is unclear and remains open to debate. To limit the risk of
421 compromised glycaemic control and ketosis a time limit of <2hours is proposed based on rapid acting
422 insulin pharmacokinetics (91).

423 Post-exercise hyperglycaemia is a common complaint for patients doing intense exercise,
424 particularly if insulin levels are reduced. CSII seems to offer advantages over MDI in managing early
425 post-exercise hyperglycaemia (94) and late-onset post-exercise hypoglycaemia (95), due to the
426 increased flexibility around basal insulin adjustments. Overcorrection of post-exercise hyperglycaemia
427 via repeated insulin dose administration results in increased risk for severe late-onset hypoglycaemia,
428 which may even be fatal (49).

429

430 Strategies to reduce the risk of post-exercise late-onset hypoglycaemia

431 Increased insulin sensitivity lasts up to 24-48 hours following exercise (46). Very few studies
432 have tested various nutrient or insulin dose adjustments to prevent hypoglycaemia after exercise.
433 Nocturnal hypoglycaemia after exercise is a major occurrence for individuals with T1D (96), with
434 increased risk for afternoon exercise (47,97). Immediate increases in post-exercise insulin sensitivity can
435 be accommodated for by reductions in the bolus insulin at the meal after exercise by ~50%, along with a
436 low GI snack at bedtime (77). In one study of 16 youth, a ~20% temporary pump basal rate reduction
437 from bedtime for 6 hours reduced nocturnal hypoglycaemia risk (95). Similarly, in another study of ten
438 males on MDI, a 20% basal rate reduction on the exercise day along with a “free” CHO snack at bedtime
439 (0.4 g CHO/kg body mass) reduced hypoglycaemia risk overnight (88). Individuals at high risk of severe
440 nocturnal hypoglycaemia (e.g., recurrent hypoglycaemia, and those sleeping alone), should take
441 additional preventive measures including blood glucose checks at 2-3AM and/or use a real time CGM
442 system with alarms and automatic pump suspension (98). A snack alone, without changes to basal
443 insulin therapy, does not appear to entirely eliminate nocturnal hypoglycaemia risk (77) and alcohol
444 intake may increase risk (99).

445

446 **Glucose monitoring, CGM and other emerging tools for exercise management**

447 A range of treatment regimens exists for people with T1D. CSII offers better flexibility in basal
448 insulin adjustments and the management of exercise-associated hyperglycaemia (100). CSII is associated
449 with reduced post-exercise hyperglycaemia compared to MDI (94), but can create frustrating challenges
450 for sports requiring pump disconnection (101). CSII can also contribute to a greater sense of being
451 “diseased” for some individuals and may promote stigma (101). Prolonged pump disconnect (> 60
452 minutes) should be managed with reconnecting, testing and re-infusion if necessary, or a change to
453 basal insulin provision by needle. CGM provides comprehensive information on blood glucose levels,
454 real-time trends in glucose levels and rates of glucose change in glucose, which can be used to prevent
455 lows during exercise (102), even in unique settings when self-monitoring of blood glucose (SMBG) is
456 difficult to perform (103). Current sensors are reasonably accurate for exercise (104,105); however, the
457 lag time in glucose equilibrium with the interstitial space and the rapid turnover in glucose during
458 exercise may impact accuracy (i.e. overestimate glucose value when levels are dropping and
459 underestimate it when levels are rising) (106,107).

460 Structured educational sessions can be implemented using downloads of SMBG, CGM and CSII
461 (108). CGM now offers the option to add “followers” who can view glucose levels in real time and
462 potentially alert the patient while he/she is playing sports. Threshold suspension of insulin delivery in
463 CSII may offer additional protection against exercise-associated hypoglycaemia according to some
464 limited data (109). The development of a fully artificial pancreas for exercise remains an elusive goal
465 (110).

466 **Summary**

467 Regular physical activity should be a routine objective for patients with type 1 diabetes for a
468 variety of health and fitness reasons. Considerable challenges remain for the person with T1D, and their
469 HCP team, in exercise/sports management. A number of small observational studies and a limited
470 number of clinical trials have been published to date that help to inform the consensus
471 recommendations here. More studies are needed to determine how to best prevent exercise-associated
472 hypoglycaemia with basal rate insulin dose adjustments and how to manage in the post-exercise
473 recovery period. In general, aerobic exercise is associated with reductions in glycaemia while anaerobic
474 exercise may be associated with a transient rise in glucose levels. Both forms of exercise can cause
475 delayed-onset hypoglycaemia in recovery. A sound understanding of the physiology of different forms of
476 exercise and the variables that can influence glycaemia during exercise and sport should underpin the
477 implementation of safe and effective glycaemic management strategies. For aerobic exercise, reductions
478 in insulin administration before the activity (basal and/or bolus) can help ameliorate hypoglycaemia risk,
479 as can increasing CHO intake to 60 grams per hour or more. For anaerobic exercise, conservative insulin
480 dose corrections may be required, although this too may increase the risk for nocturnal hypoglycaemia,
481 particularly if the exercise is performed late in the day. In all instances, more vigilance around glucose
482 monitoring is needed before, during and after the activity.

483

484 **Contributors**

485 The literature search was conducted by MCR, IWG and CES. All authors (MCR, IWG, CES, CET, PA,
486 ANL, AK, RR-L, RM, CH, FA, PF, CG, BB, PG, TWJ, ISM, TH, AP, AP, and LML) contributed to the original
487 draft of the manuscript. MCR, FA and CES edited the revised manuscript. All authors approved the final
488 submission.

489 **Declaration of interest**

490 The authors declare no relevant conflicts of interests that influence the content of this consensus
491 review.

492

493 Table 1. Pre-exercise blood glucose concentrations and initial glucose management strategies.

Starting blood glucose concentrations	General Recommendations*
Below Target (<5 mmol/l)	<ul style="list-style-type: none"> ▪ Ingest 10–20 g of glucose before starting exercise ▪ Delay exercise until blood glucose > 5mmol/l (90 mg/dL) and monitor closely for hypoglycaemia
Near target (5-6.9mmol/l)	<ul style="list-style-type: none"> ▪ Ingest 10 g of glucose before starting aerobic exercise ▪ Anaerobic exercise and HIIT sessions can be started
Target (7-10mmol/l)	<ul style="list-style-type: none"> ▪ Aerobic exercise can be started ▪ Anaerobic exercise and HIIT sessions can be started but glucose levels may rise
Slightly above target 10.1-15.0 mmol/l)	<ul style="list-style-type: none"> ▪ Aerobic exercise can be started ▪ Anaerobic exercise can be started but glucose levels may rise
Above target (>15 mmol/l)	<ul style="list-style-type: none"> ▪ If the hyperglycaemia is unexplained (not associated with a recent meal), check blood ketones. If ketones are modestly elevated (up to 1.4 mmol/l), exercise should be limited to a light intensity for only a brief duration (<30 minutes) and a small corrective insulin dose may be needed before the exercise begins. If blood ketones are elevated (≥ 1.5mmol/l), exercise is contraindicated and management should be initiated rapidly as per the advice of the HCP/team. ▪ Mild to moderate aerobic exercise may be started if blood ketones are low (<0.6 mmol/l) or if urine ketones are less than 2+. Blood glucose levels should be monitored during exercise to help notify if glucose is rising further. ▪ Intense exercise should be initiated only with caution as it may promote a further rise in glycaemia.

494 *Note: The CHO intake amounts shown here are to help with glucose stability at the start of exercise. For aerobic
 495 activities lasting greater than 30 minutes, additional CHOs will likely be needed (see Table 2). Blood glucose levels
 496 at the start of exercise must also be viewed within a wider context. Factors to consider include directional trends in
 497 glucose concentrations, insulin levels, patient safety and individual patient preferences based on experience. CHO
 498 intake will need to be higher if circulating insulin levels are high at the onset of exercise. See Nutritional
 499 Management section. HIIT= high intensity interval training.

500

501 Table 2 CHO requirements for endurance (aerobic) exercise performance and hypoglycaemia prevention

Situation	Endurance exercise performance (Athletes with and without diabetes)	Hypoglycaemia prevention under low insulin conditions	Hypoglycaemia prevention under high insulin conditions
Pre-exercise meal (low fat, low GI)	A minimum of 1g CHO/kg BW according to exercise intensity and type	A minimum of 1g CHO/kg BW according to exercise intensity and type	A minimum of 1g CHO/kg BW according to exercise intensity and type
Immediately pre-exercise (high GI)	No CHO required for performance	If BG < 5mmol/l ingest 10-20g CHO	If BG < 5mmol/l ingest 20-30g CHO
Up to 30 min duration	No CHO required for performance	If BG < 5mmol/l ingest 10-20g CHO	May require 15-30g CHO to prevent or treat hypoglycaemia
30- 60 min duration	Small amounts of CHO (10-15 g/hr) may enhance performance	<u>Low- moderate intensity (aerobic):</u> Small amounts of CHO (10-15 g/hr) depending on the exercise intensity and BG	May require up to 15-30g CHO/30 min to prevent hypoglycaemia
		<u>High intensity (anaerobic):</u> No CHO required during exercise unless BG is < 5 mmol/l then ingest 10-20g CHO. Replace CHO needs post-exercise.	
60- 150 min duration	30-60g CHO/hr	30-60g CHO/hr to prevent hypoglycaemia and enhance performance	Up to 75 g CHO/hr to prevent hypoglycaemia and enhance performance*
> 150 min duration (Mixture of CHO sources)	60-90g CHO/hr spread across the activity (e.g. 20-30g CHO/20 min) Use CHO sources that utilize different gut transporters (e.g. glucose and fructose)	Follow sports nutrition guidelines (60-90g/hr) with appropriate insulin adjustment for glycaemic management	
Post- exercise meal	1-1.2g CHO/kg body BW	Follow sports nutrition guidelines to maximise recovery with appropriate insulin adjustment for glycaemic management	

502 Note: These guidelines are based on the following references (63,111,112) and on the expert opinion of the
 503 authors. BW= body weight, BG= blood glucose concentration. * Note: CHO consumption at a high rate may cause
 504 gastric upset in some individuals and may contribute to hyperglycaemia during and after the activity. To increase
 505 CHO absorption rate during exercise, and maintain hydration status, sport beverages containing glucose and
 506 fructose may be preferable.

507 Table 3 Examples of factors that need to be considered before making adjustments for exercise.

Factor	Effect
Subcutaneous insulin injection and its adjustments	<ul style="list-style-type: none"> • Difference in injection site and depth affect insulin absorption characteristics (113,114). • Lipodystrophy. • Misunderstanding of insulin pharmacokinetic often leads to inappropriate insulin adjustments, including excessive insulin corrections (stacking), which may be particularly dangerous after exercise. • Rapid acting (30), regular and intermediate acting (115,116), but likely not long acting (117) insulin absorption rates are increased with exercise.
CHO intake	<ul style="list-style-type: none"> • Variation in CHO quantity (including inaccuracy to evaluate intake) and type will impact glycaemic excursions (118).
Self-monitored capillary glucose measurements and CGM	<ul style="list-style-type: none"> • Errors in SMBG sampling or measurement errors (SMBG, CGM) may result in inappropriate insulin dose estimations (119,120). • CGM accuracy, while improving, can be compromised by poor SMBG accuracy and calibrations methods (121). • Lag time in CGM may impact accuracy during exercise (104,106).
Medications/ alcohol	<ul style="list-style-type: none"> • Insulin sensitivity may be impacted (99) as might glucose monitoring tools (120).
Physiological cycles	<ul style="list-style-type: none"> • Diurnal endocrine variation, menstrual cycle and pregnancy impact insulin sensitivity and impact glycaemic patterns (122).
Changes in work and sleep patterns	<ul style="list-style-type: none"> • Require changes in timing of insulin basal dose administration. • Timing of exercise should be considered relative to insulin sensitivity and nocturnal hypoglycaemia risk (47).
Intercurrence illness and stress	<ul style="list-style-type: none"> • May require changes in both basal and bolus insulin dose (123). • Vigorous exercise contraindicated.

508
509

510 Table 4. Therapeutic adjustment options (insulin and/or food intake) to minimize glycaemic excursions for
 511 prolonged aerobic and brief high intensity aerobic/anaerobic exercise.

Adjustment	Prolonged endurance exercise (predominantly aerobic)	Brief intense exercise (aerobic and anaerobic)
Pre-exercise meal bolus dose insulin reduction	Advised when exercise occurs within ~120min of bolus dose The magnitude of reduction vary according to timing, type, duration and intensity of exercise (see Table 5)	Bolus reduction not advised May require additional conservative bolus dose correction if hyperglycaemia develops
Pre-exercise basal insulin dose reduction in (MDI patients) by ~20%	Useful especially if exercise occurs less than every 3 days or if exercise levels are elevated throughout the day May also be useful if on twice daily intermediate insulin	Basal reduction not advised
Basal nocturnal insulin dose reduction (MDI & CSII) following exercise by ~20% to reduce nocturnal hypoglycaemia	Particularly important if the exercise occurred in the afternoon or early evening	Useful for helping limiting post-exercise hypoglycaemia after a HIIT session
Temporary basal rate change (CSII)	Reduce basal rate to as low as total suspension of normal basal during exercise To take into account rapid acting insulin pharmacokinetics, this basal rate reduction should ideally occur well before exercise start (up to 90 minutes before) Normal basal rates can be resumed either at the end of exercise, or later in recovery depending on glucose trends	Increased basal rate may be needed to help prevent/treat hyperglycaemia either during or immediately after exercise
Pre-exercise CHO intake	See Table 2	Not usually needed
Intra-exercise CHO intake	Typically up to 60g/h if no insulin dose adjustments have been made See Table 2 for additional information	Not usually needed
Pre-exercise or post-exercise sprint	May help reduce hypoglycaemia risk	May increase hyperglycaemia risk Consider a prolonged aerobic cool down
Post-exercise CHO intake	Useful to reduce risk of hypoglycaemia and enhance recovery (see Nutritional Management section) May need a specified insulin bolus depending on length and intensity of exercise (may need a reduced insulin to CHO ratio)	Useful to reduce risk of hypoglycaemia and enhance recovery but should be delayed if hyperglycaemia is initially observed (see Nutritional management section) May need a specified insulin bolus strategy (e.g. may need a reduced insulin to CHO ratio)

512 Table 5: Suggested pre-exercise meal bolus percent reduction for exercise started within 90min of a meal.

Exercise intensity	Exercise duration	
	30 minutes	60 minutes
Mild aerobic (~25%VO ₂ max)	- 25%*	- 50%
Moderate aerobic (~50% VO ₂ max)	- 50%	- 75%
Heavy aerobic (70-75% VO ₂ max)	- 75%	N-A
Intense aerobic/anaerobic (>80% VO ₂ max)	No reduction recommended	N-A

513 Notes: Recommendations based on the following references (51,55,72,124); N-A: Not assessed, since the exercise
 514 intensity is typically too high to sustain for 60min for most individuals; * Estimated from the 60min study.

515

516 **References**

- 517 1. Mayor S. Insulin pumps improve glucose control in children with type 1 diabetes, study finds.
518 BMJ. 2015;351:h5998.
- 519 2. McKnight JA, Wild SH, Lamb MJE, Cooper MN, Jones TW, Davis EA, et al. Glycaemic control of
520 Type 1 diabetes in clinical practice early in the 21st century: an international comparison. *Diabet*
521 *Med J Br Diabet Assoc.* 2015 Aug;32(8):1036–50.
- 522 3. Bohn B, Herbst A, Pfeifer M, Krakow D, Zimny S, Kopp F, et al. Impact of Physical Activity on
523 Glycemic Control and Prevalence of Cardiovascular Risk Factors in Adults With Type 1 Diabetes: A
524 Cross-sectional Multicenter Study of 18,028 Patients. *Diabetes Care.* 2015 Aug;38(8):1536–43.
- 525 4. McCarthy MM, Funk M, Grey M. Cardiovascular health in adults with type 1 diabetes. *Prev Med.*
526 2016 Aug 12;91:138–43.
- 527 5. Herbst A, Kordonouri O, Schwab KO, Schmidt F, Holl RW, Germany DI of the GWG for PD. Impact
528 of physical activity on cardiovascular risk factors in children with type 1 diabetes: a multicenter
529 study of 23,251 patients. *Diabetes Care.* 2007 Aug;30(8):2098–100.
- 530 6. Quirk H, Blake H, Tennyson R, Randell TL, Glazebrook C. Physical activity interventions in children
531 and young people with Type 1 diabetes mellitus: a systematic review with meta-analysis. *Diabet*
532 *Med J Br Diabet Assoc.* 2014 Oct;31(10):1163–73.
- 533 7. Quirk H, Blake H, Tennyson R, Randell TL, Glazebrook C. Physical activity interventions in children
534 and young people with Type 1 diabetes mellitus: a systematic review with meta-analysis. *Diabet*
535 *Med J Br Diabet Assoc.* 2014 Oct;31(10):1163–73.
- 536 8. Katz M, Giani E, Laffel L. Challenges and Opportunities in the Management of Cardiovascular Risk
537 Factors in Youth With Type 1 Diabetes: Lifestyle and Beyond. *Curr Diab Rep.* 2015
538 Dec;15(12):119.
- 539 9. Kriska AM, LaPorte RE, Patrick SL, Kuller LH, Orchard TJ. The association of physical activity and
540 diabetic complications in individuals with insulin-dependent diabetes mellitus: the Epidemiology
541 of Diabetes Complications Study–VII. *J Clin Epidemiol.* 1991;44(11):1207–14.
- 542 10. Chimen M, Kennedy A, Nirantharakumar K, Pang TT, Andrews R, Narendran P. What are the
543 health benefits of physical activity in type 1 diabetes mellitus? A literature review. *Diabetologia.*
544 2012 Mar;55(3):542–51.
- 545 11. Pierre-Louis B, Guddati AK, Khyzar Hayat Syed M, Gorospe VE, Manguerra M, Bagchi C, et al.
546 Exercise capacity as an independent risk factor for adverse cardiovascular outcomes among
547 nondiabetic and diabetic patients. *Arch Med Sci AMS.* 2014 Feb 24;10(1):25–32.
- 548 12. Brazeau AS, Rabasa-Lhoret R, Strychar I, Mircescu H. Barriers to physical activity among patients
549 with type 1 diabetes. *Diabetes Care.* 2008 Nov;31(11):2108–9.
- 550 13. Jabbour G, Henderson M, Mathieu M-E. Barriers to Active Lifestyles in Children with Type 1
551 Diabetes. *Can J Diabetes.* 2016 Apr;40(2):170–2.

- 552 14. Lascar N, Kennedy A, Hancock B, Jenkins D, Andrews RC, Greenfield S, et al. Attitudes and Barriers
553 to Exercise in Adults with Type 1 Diabetes (T1DM) and How Best to Address Them: A Qualitative
554 Study. *PLoS One*. 2014 Sep 19;9(9):e108019.
- 555 15. Robertson K, Riddell MC, Guinhouya BC, Adolfsson P, Hanas R. Exercise in children and
556 adolescents with diabetes. *Pediatr Diabetes*. 2014 Sep;15 Suppl 20:203–23.
- 557 16. Galler A, Lindau M, Ernert A, Thalemann R, Raile K. Associations between media consumption
558 habits, physical activity, socioeconomic status, and glycemic control in children, adolescents, and
559 young adults with type 1 diabetes. *Diabetes Care*. 2011 Nov;34(11):2356–9.
- 560 17. Batacan RB, Duncan MJ, Dalbo VJ, Tucker PS, Fenning AS. Effects of high-intensity interval training
561 on cardiometabolic health: a systematic review and meta-analysis of intervention studies. *Br J
562 Sports Med*. 2016 Oct 20;
- 563 18. Colberg SR, Sigal RJ, Yardley JE, Riddell MC, Dunstan DW, Dempsey PC, et al. Physical
564 Activity/Exercise and Diabetes: A Position Statement of the American Diabetes Association.
565 *Diabetes Care*. 2016 Nov 1;39(11):2065–79.
- 566 19. Mitranun W, Deerochanawong C, Tanaka H, Suksom D. Continuous vs interval training on
567 glycemic control and macro- and microvascular reactivity in type 2 diabetic patients. *Scand J Med
568 Sci Sports*. 2014 Apr;24(2):e69-76.
- 569 20. Yardley JE, Hay J, Abou-Setta AM, Marks SD, McGavock J. A systematic review and meta-analysis
570 of exercise interventions in adults with type 1 diabetes. *Diabetes Res Clin Pract*. 2014
571 Dec;106(3):393–400.
- 572 21. Camacho RC, Galassetti P, Davis SN, Wasserman DH. Glucoregulation during and after exercise in
573 health and insulin-dependent diabetes. *Exerc Sport Sci Rev*. 2005 Jan;33(1):17–23.
- 574 22. Ploug T, Galbo H, Richter EA. Increased muscle glucose uptake during contractions: no need for
575 insulin. *Am J Physiol*. 1984 Dec;247(6 Pt 1):E726-731.
- 576 23. Coker RH, Kjaer M. Glucoregulation during exercise : the role of the neuroendocrine system.
577 *Sports Med Auckl NZ*. 2005;35(7):575–83.
- 578 24. Coyle EF. Substrate utilization during exercise in active people. *Am J Clin Nutr*. 1995 Apr;61(4
579 Suppl):968S–979S.
- 580 25. Venables MC, Achten J, Jeukendrup AE. Determinants of fat oxidation during exercise in healthy
581 men and women: a cross-sectional study. *J Appl Physiol Bethesda Md 1985*. 2005 Jan;98(1):160–
582 7.
- 583 26. van Loon LJ, Greenhaff PL, Constantin-Teodosiu D, Saris WH, Wagenmakers AJ. The effects of
584 increasing exercise intensity on muscle fuel utilisation in humans. *J Physiol*. 2001 Oct 1;536(Pt
585 1):295–304.

- 586 27. Fahey AJ, Paramalingam N, Davey RJ, Davis EA, Jones TW, Fournier PA. The effect of a short sprint
587 on postexercise whole-body glucose production and utilization rates in individuals with type 1
588 diabetes mellitus. *J Clin Endocrinol Metab.* 2012 Nov;97(11):4193–200.
- 589 28. Harmer AR, Chisholm DJ, McKenna MJ, Morris NR, Thom JM, Bennett G, et al. High-intensity
590 training improves plasma glucose and acid-base regulation during intermittent maximal exercise
591 in type 1 diabetes. *Diabetes Care.* 2007 May;30(5):1269–71.
- 592 29. Bally L, Laimer M, Stettler C. Exercise-associated glucose metabolism in individuals with type 1
593 diabetes mellitus. *Curr Opin Clin Nutr Metab Care.* 2015 Jul;18(4):428–33.
- 594 30. Mallad A, Hinshaw L, Schiavon M, Dalla Man C, Dadlani V, Basu R, et al. Exercise Effects on
595 Postprandial Glucose Metabolism in Type 1 Diabetes: A Triple Tracer Approach. *Am J Physiol*
596 *Endocrinol Metab.* 2015 Apr 21;ajpendo.00014.2015.
- 597 31. Frayn KN, Karpe F. Regulation of human subcutaneous adipose tissue blood flow. *Int J Obes* 2005.
598 2014 Aug;38(8):1019–26.
- 599 32. McAuley SA, Horsburgh JC, Ward GM, La Gerche A, Gooley JL, Jenkins AJ, et al. Insulin pump basal
600 adjustment for exercise in type 1 diabetes: a randomised crossover study. *Diabetologia.* 2016
601 Aug;59(8):1636–44.
- 602 33. Tansey MJ, Tsalikian E, Beck RW, Mauras N, Buckingham BA, Weinzimer SA, et al. The effects of
603 aerobic exercise on glucose and counterregulatory hormone concentrations in children with type
604 1 diabetes. *Diabetes Care.* 2006 Jan;29(1):20–5.
- 605 34. Riddell MC, Bar-Or O, Ayub BV, Calvert RE, Heigenhauser GJ. Glucose ingestion matched with
606 total carbohydrate utilization attenuates hypoglycemia during exercise in adolescents with IDDM.
607 *Int J Sport Nutr.* 1999 Mar;9(1):24–34.
- 608 35. Al Khalifah RA, Suppère C, Haidar A, Rabasa-Lhoret R, Ladouceur M, Legault L. Association of
609 aerobic fitness level with exercise-induced hypoglycaemia in Type 1 diabetes. *Diabet Med J Br*
610 *Diabet Assoc.* 2016 Jan 16;
- 611 36. Harmer AR, Chisholm DJ, McKenna MJ, Hunter SK, Ruell PA, Naylor JM, et al. Sprint training
612 increases muscle oxidative metabolism during high-intensity exercise in patients with type 1
613 diabetes. *Diabetes Care.* 2008 Nov;31(11):2097–102.
- 614 37. Singhvi A, Tansey MJ, Janz K, Zimmerman MB, Tsalikian E. Aerobic fitness and glycemic variability
615 in adolescents with type 1 diabetes. *Endocr Pract Off J Am Coll Endocrinol Am Assoc Clin*
616 *Endocrinol.* 2014 Jun;20(6):566–70.
- 617 38. Zander E, Bruns W, Wulfert P, Besch W, Lubs D, Chlup R, et al. Muscular exercise in type I-
618 diabetics. I. Different metabolic reactions during heavy muscular work in dependence on actual
619 insulin availability. *Exp Clin Endocrinol.* 1983 Jul;82(1):78–90.
- 620 39. Berger M, Berchtold P, Cüppers HJ, Drost H, Kley HK, Müller WA, et al. Metabolic and hormonal
621 effects of muscular exercise in juvenile type diabetics. *Diabetologia.* 1977 Aug;13(4):355–65.

- 622 40. Yardley JE, Kenny GP, Perkins BA, Riddell MC, Balaa N, Malcolm J, et al. Resistance versus aerobic
623 exercise: acute effects on glycemia in type 1 diabetes. *Diabetes Care*. 2013 Mar;36(3):537–42.
- 624 41. Turner D, Luzio S, Gray BJ, Dunseath G, Rees ED, Kilduff LP, et al. Impact of single and multiple
625 sets of resistance exercise in type 1 diabetes. *Scand J Med Sci Sports*. 2015 Feb;25(1):e99-109.
- 626 42. Guelfi KJ, Ratnam N, Smythe GA, Jones TW, Fournier PA. Effect of intermittent high-intensity
627 compared with continuous moderate exercise on glucose production and utilization in individuals
628 with type 1 diabetes. *Am J Physiol Endocrinol Metab*. 2007 Mar;292(3):E865-870.
- 629 43. Yardley JE, Kenny GP, Perkins BA, Riddell MC, Malcolm J, Boulay P, et al. Effects of performing
630 resistance exercise before versus after aerobic exercise on glycemia in type 1 diabetes. *Diabetes*
631 *Care*. 2012 Apr;35(4):669–75.
- 632 44. Bally L, Zueger T, Buehler T, Dokumaci AS, Speck C, Pasi N, et al. Metabolic and hormonal
633 response to intermittent high-intensity and continuous moderate intensity exercise in individuals
634 with type 1 diabetes: a randomised crossover study. *Diabetologia*. 2016 Apr;59(4):776–84.
- 635 45. Davey RJ, Paramalingam N, Retterath AJ, Lim EM, Davis EA, Jones TW, et al. Antecedent
636 hypoglycaemia does not diminish the glycaemia-increasing effect and glucoregulatory responses
637 of a 10 s sprint in people with type 1 diabetes. *Diabetologia*. 2014 Jun;57(6):1111–8.
- 638 46. Teich T, Riddell MC. The Enhancement of Muscle Insulin Sensitivity After Exercise: A Rac1-
639 Independent Handoff to Some Other Player? *Endocrinology*. 2016 Aug;157(8):2999–3001.
- 640 47. Gomez AM, Gomez C, Aschner P, Veloza A, Muñoz O, Rubio C, et al. Effects of performing
641 morning versus afternoon exercise on glycemic control and hypoglycemia frequency in type 1
642 diabetes patients on sensor-augmented insulin pump therapy. *J Diabetes Sci Technol*. 2015
643 May;9(3):619–24.
- 644 48. Turner D, Luzio S, Gray BJ, Bain SC, Hanley S, Richards A, et al. Algorithm that delivers an
645 individualized rapid-acting insulin dose after morning resistance exercise counters post-exercise
646 hyperglycaemia in people with Type 1 diabetes. *Diabet Med J Br Diabet Assoc*. 2015 Jul 29;
- 647 49. Tanenberg RJ, Newton CA, Drake AJ. Confirmation of hypoglycemia in the “dead-in-bed”
648 syndrome, as captured by a retrospective continuous glucose monitoring system. *Endocr Pract*
649 *Off J Am Coll Endocrinol Am Assoc Clin Endocrinol*. 2010 Apr;16(2):244–8.
- 650 50. Maran A, Pavan P, Bonsembiante B, Brugin E, Ermolao A, Avogaro A, et al. Continuous glucose
651 monitoring reveals delayed nocturnal hypoglycemia after intermittent high-intensity exercise in
652 nontrained patients with type 1 diabetes. *Diabetes Technol Ther*. 2010 Oct;12(10):763–8.
- 653 51. Moser O, Tschakert G, Mueller A, Groeschl W, Pieber TR, Obermayer-Pietsch B, et al. Effects of
654 High-Intensity Interval Exercise versus Moderate Continuous Exercise on Glucose Homeostasis
655 and Hormone Response in Patients with Type 1 Diabetes Mellitus Using Novel Ultra-Long-Acting
656 Insulin. *PLoS One*. 2015;10(8):e0136489.

- 657 52. Iscoe KE, Riddell MC. Continuous moderate-intensity exercise with or without intermittent high-
658 intensity work: effects on acute and late glycaemia in athletes with Type 1 diabetes mellitus.
659 *Diabet Med J Br Diabet Assoc.* 2011 Jul;28(7):824–32.
- 660 53. Kennedy A, Nirantharakumar K, Chimen M, Pang TT, Hemming K, Andrews RC, et al. Does
661 exercise improve glycaemic control in type 1 diabetes? A systematic review and meta-analysis.
662 *PLoS One.* 2013;8(3):e58861.
- 663 54. Adolfsson P, Mattsson S, Jendle J. Evaluation of glucose control when a new strategy of increased
664 carbohydrate supply is implemented during prolonged physical exercise in type 1 diabetes. *Eur J*
665 *Appl Physiol.* 2015 Dec;115(12):2599–607.
- 666 55. Rabasa-Lhoret R, Bourque J, Ducros F, Chiasson JL. Guidelines for premeal insulin dose reduction
667 for postprandial exercise of different intensities and durations in type 1 diabetic subjects treated
668 intensively with a basal-bolus insulin regimen (ultralente-lispro). *Diabetes Care.* 2001
669 Apr;24(4):625–30.
- 670 56. Campbell MD, Walker M, Trenell MI, Luzio S, Dunseath G, Tuner D, et al. Metabolic implications
671 when employing heavy pre- and post-exercise rapid-acting insulin reductions to prevent
672 hypoglycaemia in type 1 diabetes patients: a randomised clinical trial. *PLoS One.*
673 2014;9(5):e97143.
- 674 57. Dubé M-C, Weisnagel SJ, Prud'homme D, Lavoie C. Is early and late post-meal exercise so
675 different in type 1 diabetic lispro users? *Diabetes Res Clin Pract.* 2006 May;72(2):128–34.
- 676 58. Kelly D, Hamilton JK, Riddell MC. Blood glucose levels and performance in a sports camp for
677 adolescents with type 1 diabetes mellitus: a field study. *Int J Pediatr.* 2010;2010.
- 678 59. Graveling AJ, Frier BM. Risks of marathon running and hypoglycaemia in Type 1 diabetes. *Diabet*
679 *Med J Br Diabet Assoc.* 2010 May;27(5):585–8.
- 680 60. Galassetti P, Tate D, Neill RA, Richardson A, Leu S-Y, Davis SN. Effect of differing antecedent
681 hypoglycemia on counterregulatory responses to exercise in type 1 diabetes. *Am J Physiol*
682 *Endocrinol Metab.* 2006 Jun;290(6):E1109-1117.
- 683 61. Riddell MC, Burr J. Evidence-based risk assessment and recommendations for physical activity
684 clearance: diabetes mellitus and related comorbidities. *Appl Physiol Nutr Metab Physiol*
685 *Appliquée Nutr Métabolisme.* 2011 Jul;36 Suppl 1:S154-189.
- 686 62. American Diabetes Association. (4) Foundations of care: education, nutrition, physical activity,
687 smoking cessation, psychosocial care, and immunization. *Diabetes Care.* 2015 Jan;38 Suppl:S20-
688 30.
- 689 63. Thomas DT, Erdman KA, Burke LM. Position of the Academy of Nutrition and Dietetics, Dietitians
690 of Canada, and the American College of Sports Medicine: Nutrition and Athletic Performance. *J*
691 *Acad Nutr Diet.* 2016 Mar;116(3):501–28.

- 692 64. Franc S, Daoudi A, Pochat A, Petit M-H, Randazzo C, Petit C, et al. Insulin-based strategies to
693 prevent hypoglycaemia during and after exercise in adult patients with type 1 diabetes on pump
694 therapy: the DIABRASPORT randomized study. *Diabetes Obes Metab.* 2015 Dec;17(12):1150–7.
- 695 65. American College of Sports Medicine, Sawka MN, Burke LM, Eichner ER, Maughan RJ, Montain SJ,
696 et al. American College of Sports Medicine position stand. Exercise and fluid replacement. *Med
697 Sci Sports Exerc.* 2007 Feb;39(2):377–90.
- 698 66. Loucks AB, Kiens B, Wright HH. Energy availability in athletes. *J Sports Sci.* 2011;29 Suppl 1:S7-15.
- 699 67. Frankenfield D, Roth-Yousey L, Compher C. Comparison of predictive equations for resting
700 metabolic rate in healthy nonobese and obese adults: a systematic review. *J Am Diet Assoc.* 2005
701 May;105(5):775–89.
- 702 68. Murphy CH, Hector AJ, Phillips SM. Considerations for protein intake in managing weight loss in
703 athletes. *Eur J Sport Sci.* 2015;15(1):21–8.
- 704 69. Phillips SM. Dietary protein requirements and adaptive advantages in athletes. *Br J Nutr.* 2012
705 Aug;108 Suppl 2:S158-167.
- 706 70. Phillips SM, Chevalier S, Leidy HJ. Protein “requirements” beyond the RDA: implications for
707 optimizing health. *Appl Physiol Nutr Metab Physiol Appl Nutr Metab.* 2016 May;41(5):565–72.
- 708 71. Mettler S, Mitchell N, Tipton KD. Increased protein intake reduces lean body mass loss during
709 weight loss in athletes. *Med Sci Sports Exerc.* 2010 Feb;42(2):326–37.
- 710 72. Shetty VB, Fournier PA, Davey RJ, Retterath AJ, Paramalingam N, Roby HC, et al. Effect of Exercise
711 Intensity on Glucose Requirements to Maintain Euglycemia During Exercise in Type 1 Diabetes. *J
712 Clin Endocrinol Metab.* 2016 Mar;101(3):972–80.
- 713 73. Hernandez JM, Moccia T, Fluckey JD, Ulbrecht JS, Farrell PA. Fluid snacks to help persons with
714 type 1 diabetes avoid late onset postexercise hypoglycemia. *Med Sci Sports Exerc.* 2000
715 May;32(5):904–10.
- 716 74. Beelen M, Burke LM, Gibala MJ, van Loon LJC. Nutritional strategies to promote postexercise
717 recovery. *Int J Sport Nutr Exerc Metab.* 2010 Dec;20(6):515–32.
- 718 75. van Albada ME, Bakker-van Waarde WM. Recurrent nightly ketosis after prolonged exercise in
719 type 1 diabetes - the need for glycogen replacement strategies. Case report and review of
720 literature. *Pediatr Diabetes.* 2015 Nov 4;
- 721 76. Bracken RM, Page R, Gray B, Kilduff LP, West DJ, Stephens JW, et al. Isomaltulose improves
722 glycemia and maintains run performance in type 1 diabetes. *Med Sci Sports Exerc.* 2012
723 May;44(5):800–8.
- 724 77. Campbell MD, Walker M, Trenell MI, Stevenson EJ, Turner D, Bracken RM, et al. A low-glycemic
725 index meal and bedtime snack prevents postprandial hyperglycemia and associated rises in
726 inflammatory markers, providing protection from early but not late nocturnal hypoglycemia
727 following evening exercise in type 1 diabetes. *Diabetes Care.* 2014 Jul;37(7):1845–53.

- 728 78. Perrone C, Laitano O, Meyer F. Effect of carbohydrate ingestion on the glycemic response of type
729 1 diabetic adolescents during exercise. *Diabetes Care*. 2005 Oct;28(10):2537–8.
- 730 79. Burke LM. Re-Examining High-Fat Diets for Sports Performance: Did We Call the “Nail in the
731 Coffin” Too Soon? *Sports Med Auckl NZ*. 2015 Nov;45 Suppl 1:33–49.
- 732 80. Burke LM, Hawley JA, Angus DJ, Cox GR, Clark SA, Cummings NK, et al. Adaptations to short-term
733 high-fat diet persist during exercise despite high carbohydrate availability. *Med Sci Sports Exerc*.
734 2002 Jan;34(1):83–91.
- 735 81. Bartlett JD, Hawley JA, Morton JP. Carbohydrate availability and exercise training adaptation: too
736 much of a good thing? *Eur J Sport Sci*. 2015;15(1):3–12.
- 737 82. Paluska SA. Caffeine and exercise. *Curr Sports Med Rep*. 2003 Aug;2(4):213–9.
- 738 83. Zaharieva DP, Miadovnik LA, Rowan CP, Gumieniak RJ, Jamnik VK, Riddell MC. Effects of acute
739 caffeine supplementation on reducing exercise-associated hypoglycaemia in individuals with Type
740 1 diabetes mellitus. *Diabet Med J Br Diabet Assoc*. 2015 Jul 14;
- 741 84. Mauvais-Jarvis F, Sobngwi E, Porcher R, Garnier JP, Vexiau P, Duvallet A, et al. Glucose response
742 to intense aerobic exercise in type 1 diabetes: maintenance of near euglycemia despite a drastic
743 decrease in insulin dose. *Diabetes Care*. 2003 Apr;26(4):1316–7.
- 744 85. West DJ, Morton RD, Bain SC, Stephens JW, Bracken RM. Blood glucose responses to reductions
745 in pre-exercise rapid-acting insulin for 24 h after running in individuals with type 1 diabetes. *J
746 Sports Sci*. 2010 May;28(7):781–8.
- 747 86. Bracken RM, West DJ, Stephens JW, Kilduff LP, Luzio S, Bain SC. Impact of pre-exercise rapid-
748 acting insulin reductions on ketogenesis following running in Type 1 diabetes. *Diabet Med J Br
749 Diabet Assoc*. 2011 Feb;28(2):218–22.
- 750 87. West DJ, Stephens JW, Bain SC, Kilduff LP, Luzio S, Still R, et al. A combined insulin reduction and
751 carbohydrate feeding strategy 30 min before running best preserves blood glucose concentration
752 after exercise through improved fuel oxidation in type 1 diabetes mellitus. *J Sports Sci*. 2011
753 Feb;29(3):279–89.
- 754 88. Campbell MD, Walker M, Bracken RM, Turner D, Stevenson EJ, Gonzalez JT, et al. Insulin therapy
755 and dietary adjustments to normalize glycemia and prevent nocturnal hypoglycemia after
756 evening exercise in type 1 diabetes: a randomized controlled trial. *BMJ Open Diabetes Res Care*.
757 2015;3(1):e000085.
- 758 89. Arutchelvam V, Heise T, Dellweg S, Elbroend B, Minns I, Home PD. Plasma glucose and
759 hypoglycaemia following exercise in people with Type 1 diabetes: a comparison of three basal
760 insulins. *Diabet Med J Br Diabet Assoc*. 2009 Oct;26(10):1027–32.
- 761 90. Heise T, Bain SC, Bracken RM, Zijlstra E, Nosek L, Stender-Petersen K, et al. Similar risk of
762 exercise-related hypoglycaemia for insulin degludec to that for insulin glargine in patients with
763 type 1 diabetes: a randomized cross-over trial. *Diabetes Obes Metab*. 2016 Feb;18(2):196–9.

- 764 91. Heinemann L, Nosek L, Kapitza C, Schweitzer M-A, Krinelke L. Changes in basal insulin infusion
765 rates with subcutaneous insulin infusion: time until a change in metabolic effect is induced in
766 patients with type 1 diabetes. *Diabetes Care*. 2009 Aug;32(8):1437–9.
- 767 92. Diabetes Research in Children Network (DirecNet) Study Group, Tsalikian E, Kollman C,
768 Tamborlane WB, Beck RW, Fiallo-Scharer R, et al. Prevention of hypoglycemia during exercise in
769 children with type 1 diabetes by suspending basal insulin. *Diabetes Care*. 2006 Oct;29(10):2200–
770 4.
- 771 93. Admon G, Weinstein Y, Falk B, Weintrob N, Benzaquen H, Ofan R, et al. Exercise with and without
772 an insulin pump among children and adolescents with type 1 diabetes mellitus. *Pediatrics*. 2005
773 Sep;116(3):e348-355.
- 774 94. Yardley JE, Iscoe KE, Sigal RJ, Kenny GP, Perkins BA, Riddell MC. Insulin pump therapy is
775 associated with less post-exercise hyperglycemia than multiple daily injections: an observational
776 study of physically active type 1 diabetes patients. *Diabetes Technol Ther*. 2013 Jan;15(1):84–8.
- 777 95. Taplin CE, Cobry E, Messer L, McFann K, Chase HP, Fiallo-Scharer R. Preventing post-exercise
778 nocturnal hypoglycemia in children with type 1 diabetes. *J Pediatr*. 2010 Nov;157(5):784–788.e1.
- 779 96. Frier BM. Hypoglycaemia in diabetes mellitus: epidemiology and clinical implications. *Nat Rev*
780 *Endocrinol*. 2014 Dec;10(12):711–22.
- 781 97. Davey RJ, Howe W, Paramalingam N, Ferreira LD, Davis EA, Fournier PA, et al. The effect of
782 midday moderate-intensity exercise on postexercise hypoglycemia risk in individuals with type 1
783 diabetes. *J Clin Endocrinol Metab*. 2013 Jul;98(7):2908–14.
- 784 98. Garg SK, Brazg RL, Bailey TS, Buckingham BA, Slover RH, Klonoff DC, et al. Hypoglycemia begets
785 hypoglycemia: the order effect in the ASPIRE in-clinic study. *Diabetes Technol Ther*. 2014
786 Mar;16(3):125–30.
- 787 99. Richardson T, Weiss M, Thomas P, Kerr D. Day after the night before: influence of evening alcohol
788 on risk of hypoglycemia in patients with type 1 diabetes. *Diabetes Care*. 2005 Jul;28(7):1801–2.
- 789 100. Chu L, Hamilton J, Riddell MC. Clinical management of the physically active patient with type 1
790 diabetes. *Phys Sportsmed*. 2011 May;39(2):64–77.
- 791 101. Binek A, Rembierz-Knoll A, Polańska J, Jarosz-Chobot P. Reasons for the discontinuation of
792 therapy of personal insulin pump in children with type 1 diabetes. *Pediatr Endocrinol Diabetes*
793 *Metab*. 2016 Feb 18;21(2):65–9.
- 794 102. Riddell MC, Milliken J. Preventing exercise-induced hypoglycemia in type 1 diabetes using real-
795 time continuous glucose monitoring and a new carbohydrate intake algorithm: an observational
796 field study. *Diabetes Technol Ther*. 2011 Aug;13(8):819–25.
- 797 103. Adolfsson P, Ornhagen H, Jendle J. The benefits of continuous glucose monitoring and a glucose
798 monitoring schedule in individuals with type 1 diabetes during recreational diving. *J Diabetes Sci*
799 *Technol*. 2008 Sep;2(5):778–84.

- 800 104. Bally L, Zueger T, Pasi N, Carlos C, Paganini D, Stettler C. Accuracy of continuous glucose
801 monitoring during differing exercise conditions. *Diabetes Res Clin Pract.* 2016 Feb;112:1–5.
- 802 105. Yardley JE, Sigal RJ, Kenny GP, Riddell MC, Lovblom LE, Perkins BA. Point accuracy of interstitial
803 continuous glucose monitoring during exercise in type 1 diabetes. *Diabetes Technol Ther.* 2013
804 Jan;15(1):46–9.
- 805 106. Davey RJ, Low C, Jones TW, Fournier PA. Contribution of an intrinsic lag of continuous glucose
806 monitoring systems to differences in measured and actual glucose concentrations changing at
807 variable rates in vitro. *J Diabetes Sci Technol.* 2010 Nov;4(6):1393–9.
- 808 107. Taleb N, Emami A, Suppere C, Messier V, Legault L, Chiasson J-L, et al. Comparison of Two
809 Continuous Glucose Monitoring Systems, Dexcom G4 Platinum and Medtronic Paradigm Veo
810 Enlite System, at Rest and During Exercise. *Diabetes Technol Ther.* 2016 Sep;18(9):561–7.
- 811 108. Adolfsson P, Strömngren A, Mattsson S, Chaplin J, Jendle J. Education and individualized support
812 regarding exercise and diabetes improves glucose control and level of physical activity in type 1
813 diabetes individuals. *J Endocrinol Diabetes Obes.* 2015;3(2):1–6.
- 814 109. Danne T, Tsioli C, Kordonouri O, Blaesig S, Remus K, Roy A, et al. The PILGRIM study: in silico
815 modeling of a predictive low glucose management system and feasibility in youth with type 1
816 diabetes during exercise. *Diabetes Technol Ther.* 2014 Jun;16(6):338–47.
- 817 110. Riddell MC, Zaharieva DP, Yavelberg L, Cinar A, Jamnik VK. Exercise and the Development of the
818 Artificial Pancreas: One of the More Difficult Series of Hurdles. *J Diabetes Sci Technol.* 2015
819 Nov;9(6):1217–26.
- 820 111. Burke LM, Hawley JA, Wong SHS, Jeukendrup AE. Carbohydrates for training and competition. *J*
821 *Sports Sci.* 2011;29 Suppl 1:S17-27.
- 822 112. Jeukendrup A. A step towards personalized sports nutrition: carbohydrate intake during exercise.
823 *Sports Med Auckl NZ.* 2014 May;44 Suppl 1:S25-33.
- 824 113. Frid A, Ostman J, Linde B. Hypoglycemia risk during exercise after intramuscular injection of
825 insulin in thigh in IDDM. *Diabetes Care.* 1990 May;13(5):473–7.
- 826 114. Hirsch L, Byron K, Gibney M. Intramuscular risk at insulin injection sites--measurement of the
827 distance from skin to muscle and rationale for shorter-length needles for subcutaneous insulin
828 therapy. *Diabetes Technol Ther.* 2014 Dec;16(12):867–73.
- 829 115. Hildebrandt P. Subcutaneous absorption of insulin in insulin-dependent diabetic patients.
830 Influence of species, physico-chemical properties of insulin and physiological factors. *Dan Med*
831 *Bull.* 1991 Aug;38(4):337–46.
- 832 116. Rönnemaa T, Koivisto VA. Combined effect of exercise and ambient temperature on insulin
833 absorption and postprandial glycemia in type I patients. *Diabetes Care.* 1988 Dec;11(10):769–73.

- 834 117. Peter R, Luzio SD, Dunseath G, Miles A, Hare B, Backx K, et al. Effects of exercise on the
835 absorption of insulin glargine in patients with type 1 diabetes. *Diabetes Care*. 2005
836 Mar;28(3):560–5.
- 837 118. Bell KJ, Smart CE, Steil GM, Brand-Miller JC, King B, Wolpert HA. Impact of fat, protein, and
838 glycemic index on postprandial glucose control in type 1 diabetes: implications for intensive
839 diabetes management in the continuous glucose monitoring era. *Diabetes Care*. 2015
840 Jun;38(6):1008–15.
- 841 119. Breton MD, Kovatchev BP. Impact of blood glucose self-monitoring errors on glucose variability,
842 risk for hypoglycemia, and average glucose control in type 1 diabetes: an in silico study. *J*
843 *Diabetes Sci Technol*. 2010 May;4(3):562–70.
- 844 120. Tonyushkina K, Nichols JH. Glucose meters: a review of technical challenges to obtaining accurate
845 results. *J Diabetes Sci Technol*. 2009 Jul;3(4):971–80.
- 846 121. Castle JR, Jacobs PG. Nonadjunctive Use of Continuous Glucose Monitoring for Diabetes
847 Treatment Decisions. *J Diabetes Sci Technol*. 2016 Feb 15;
- 848 122. Blaak E. Sex differences in the control of glucose homeostasis. *Curr Opin Clin Nutr Metab Care*.
849 2008 Jul;11(4):500–4.
- 850 123. Deane AM, Horowitz M. Dysglycaemia in the critically ill - significance and management. *Diabetes*
851 *Obes Metab*. 2013 Sep;15(9):792–801.
- 852 124. Campbell MD, Walker M, Trenell MI, Jakovljevic DG, Stevenson EJ, Bracken RM, et al. Large pre-
853 and postexercise rapid-acting insulin reductions preserve glycemia and prevent early- but not
854 late-onset hypoglycemia in patients with type 1 diabetes. *Diabetes Care*. 2013 Aug;36(8):2217–
855 24.
- 856
- 857

858 **Figure 1: Blood glucose trends and different forms of exercise.** High patient variability exists in the
859 blood glucose responses to different form of exercise, as denoted by the arrows and grey shading. In
860 general, aerobic exercise lowers glycaemia, anaerobic exercise raises glycaemia and mixed activities is
861 associated with relative glucose stability. The individual responses depend on a number of additional
862 factors including the duration/intensity of the activity; initial blood glucose level; individual fitness;
863 levels of insulin, glucagon, other counterregulatory hormones in circulation; and the nutritional status of
864 the individual.

865

866 **Figure 2: Decision tree for aerobic exercise and mixed aerobic and anaerobic activities lasting 30 min
867 or longer.** This decision tree can serve as a starting point for decision-making for aerobic exercise.

868 Notes: ¹ Mixed activities that include anaerobic bursts of exercise may require less carbohydrate intake
869 and/or less insulin dose reductions compared continuous moderate aerobic activities. If both resistance
870 and aerobic exercise are to be performed, suggest performing resistance first to help attenuate the drop
871 in glycaemia. ² In some situations, increased carbohydrate feeding rather than insulin dose reduction
872 may help improve endurance performance in prolonged activities. ³ In other situations, both bolus and
873 basal insulin dose reductions may be preferred to help limit CHO needs. Consider CGM where patient or
874 parent preference dictates, or with history of nocturnal or severe hypoglycaemia.