

Research Space

Book chapter

Laboratory and field-based data collection (Quantitative)

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1 **Laboratory and Field-Based Data Collection (Quantitative)**
2

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4

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36

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77

78

79 **1. Introduction**

80 Rigorous assessment of sport and exercise measures is a requirement for any scientist
81 aiming to answer a research question. Sport and exercise scientists may strive to answer
82 questions such as, "Does caffeine improve an athlete's performance?", "What are the
83 physiological determinants of endurance running?" and "When can an athlete return to training
84 after injury?". Researchers aim to answer these questions through data collection in
85 experimental studies that are designed to test a hypothesis and provide robust evidence on a
86 topic. This is pertinent as the replicability of findings in sport and exercise research has been
87 questioned (Mesquida et al., 2022). By prioritizing methodological quality in research,
88 researchers can enhance the credibility and trustworthiness of their findings and, in turn,
89 promote the replicability of research findings in the field of sport and exercise science. To help
90 researchers design their studies, there are several guidelines that offer recommendations on
91 appropriate reporting (Consolidated Standards Of Reporting Trials, CONSORT) with some
92 more specific to exercise nutrition (Proper Reporting of Evidence in Sport and Exercise
93 Nutrition Trials, PRESENT) (Betts et al., 2020). While these provide excellent considerations
94 to ensure reporting of the scientific method is complete, they can also be used as guidelines
95 implemented prior to data collection to ensure that the study results are robust.

96

97 Quantitative data collection in sport and exercise research can include different methods
98 including surveys and questionnaires, biomechanical and physiological measures and exercise
99 capacity and performance measures. These data can be obtained in controlled laboratory
100 environments or in an applied setting (*e.g.*, during a race) depending upon the specific research
101 question. Here we aim to focus primarily on practical data collection, such as obtaining
102 measures of physiological responses and exercise performance. Furthermore, fundamental to
103 this is the use of randomised controlled trials, which are often regarded as the cornerstone of
104 any data collection researchers conduct across the field of science. Below, we provide an
105 overview of the essential components that researchers should consider both in the laboratory
106 and field, with emphasis given to collecting data during randomised controlled trials.

107

108 **2. Ethical considerations**

109 **1. Institutional review or ethics committee**

110 Prior to initiating data collection, researchers are required to submit their project to their
111 Institutional Review Committee or Institutional Ethics Committee which is formally

112 designated to review and approve research involving human participants according to ethical
113 principles such as the Declaration of Helsinki developed by the World Medical Association
114 ([https://www.wma.net/policies-post/wma-declaration-of-helsinki-ethical-principles-for-](https://www.wma.net/policies-post/wma-declaration-of-helsinki-ethical-principles-for-medical-research-involving-human-subjects/)
115 [medical-research-involving-human-subjects/](https://www.wma.net/policies-post/wma-declaration-of-helsinki-ethical-principles-for-medical-research-involving-human-subjects/)). Specifically, the primary role of the ethics
116 committee is to safeguard the rights, welfare, and privacy of those participating in research
117 studies. Researchers are required to submit detailed proposals outlining their study objectives,
118 methodology, participant recruitment procedures, and outcome measures to ensure
119 confidentiality and informed consent. The committee then evaluates all aspects of these study
120 proposals to ensure that potential risks to participants are minimized and that the anticipated
121 benefits of the research justify any potential harm. The committee may request some changes
122 to the proposal if they believe that the risk of certain procedures is too high or outweighs the
123 societal benefits. Only once a study has been approved by the ethics committee can a study
124 initiate participant recruitment, following obtention of informed consent (see Section 2.2
125 Informed consent). This ensures the ethical and responsible conduct of research while
126 protecting the rights and well-being of the research participants.

127

128 **2. *Informed consent***

129 Before the collection of any data, it is prerequisite to gain informed consent, ideally in
130 writing, from participants in the study. Non-written consent (i.e., verbal) should be supported
131 by witness statements or audio or video recordings to ensure all parties are covered and avoid
132 disputes as to whether consent was given. All informed consent forms should be stored safely
133 and confidentially (see Section 3.4.2. Data management). To ensure participant well-being, and
134 protect them from harm, informed consent ensures that participants are aware of the aims, the
135 method, and potential outcomes and risks associated with the study. To achieve this,
136 researchers need to provide unbiased, up-to-date, relevant information of their decision to
137 participate in the study and importantly, that participation is completely voluntary, for which
138 they can choose to withdraw at any time without reason and consequence. To help participants
139 decide whether to participate in the study, and understand potential consequences, they should
140 be provided with an information sheet that contains brief and clear information on the essential
141 aspects of the study. The Standards for Ethics in Sport and Exercise Research (Harriss et al.,
142 2022) lists what need to be included in the information sheet (see Table 1).

143

144 It is important to note that any information should be written clearly and be easy to read for
145 a layman. The use of technical and jargon should be avoided, but if required, should be first

146 explained in a plain, accessible language. Researchers may not consider that a lot of the
147 language used in their day-to-day work is in fact technical. Words used throughout this chapter,
148 for example - randomisation, sample size, blinding and validity – may be complex for a
149 participant, and as such, should be avoided to ensure they are fully aware of what is required
150 of them. Given this, researchers should aim to provide information about the study in both
151 written and spoken form. The former can be emailed or sent to participants prior to visiting the
152 data collection site (*e.g.*, the laboratory), so that they have ample opportunity to read all
153 information and be cognisant of what to expect in the study. The latter offers the opportunity
154 to expand on technical areas and provides participants the opening to question and alleviate
155 any concerns.

156

157 While most data collection will sample the general population, researchers may also be
158 interested in sampling other populations that are more vulnerable, including children, the
159 elderly, and those with intellectual impairments. Researchers will therefore need to consider
160 additional ethical concerns and be aware that it may not be possible to gain consent or that they
161 need more time. Passive assent, which can involve a parent or guardian, should be avoided
162 where possible, and every effort should be made to involve the participant in the informed
163 consent process. Explaining the details for informed consent for vulnerable groups are outside
164 the scope of this chapter, but readers are directed towards the UK Research and Innovation
165 guidance (UKRI, 2023).

166

Table. 1 Brief outline of information required for a participant information sheet

#	Information given to participant	Elaboration
1	Researcher details	Names and institutional affiliations
2	The aims of the research	Why is the work being undertaken?
3	Methods of the study	What will participants be asked to do?
4	Sources of funding	Has an organisation funded the study?
5	Conflicts of interest	Would financial or personal consideration compromise the research?
6	Anticipated benefits	What benefits can participants receive?
7	Potential risks	What harms or consequences come from participation?
8	Right to decline	Participants do not have to take part and can do so without consequence

9	Right to withdraw	Participants can withdraw consent without consequence
10	Handling of data	Where will data be stored, shared and accessed
11	Retention of data	How long will data be stored?
12	Contact details	Who can participant's contact if they have questions or complaints?

Note: Content is adapted from Harriss et al. (2022)

167

168 **3. Experimental design**

169 The extent to which the observed results of an experimental study represent the true effect
170 of the intervention depends on the rigour of the methodology. Internal validity is the term used
171 to describe whether the methodology was conducted adequately to answer the research
172 question without substantial bias (Andrade, 2018; Halperin et al., 2015). There is an extensive
173 list of confounding factors which could potentially influence exercise performance (e.g., diet,
174 sleep, fatigue) and, thus, should be considered and/or controlled to various extents depending
175 upon the research question being asked and how they might impact upon the data. External
176 validity relates to how generalisable the current data are to other contexts (Andrade, 2018). For
177 example, a study looking at the effects of a training intervention in elderly individuals with
178 type II diabetes will likely not be entirely generalisable to a young and athletic population.
179 Ecological validity is a sub-section of external validity applied to the real-world, specifically,
180 whether the study can be generalisable to everyday life. For example, a study showing the side-
181 effects of caffeine (e.g., anxiety) on participants in a resting and relaxed state in a seamlessly
182 controlled laboratory may have high internal validity but is in stark contrast to the high-pressure
183 environment of competitive sport, and results may therefore not be directly applicable.
184 Understanding of internal and external validity is vital to design and conduct studies and to
185 understand the limitations of that research. The following sections aims to critically discuss
186 their importance in relation to data collection.

187

188 ***1. Laboratory and field-based research***

189 Most research questions are focused on determining the mechanistic characteristics (e.g.,
190 physiological, psychological, biomechanical, sociological) of sport or the effectiveness of sport
191 and exercise science interventions, both in the field (applied) and laboratory. The advancement
192 and development of cutting edge and portable technologies means that researchers have a
193 plethora of methods through which to answer their research questions in both the laboratory

194 and field. While the laboratory is often the preferred choice, given its high reliability,
195 sensitivity, and ability in which to control several variables, such as temperature and humidity,
196 researchers can conduct research within the field, which offers more ecological validity that
197 can help translate findings into real world scenarios. Nevertheless, both have their own
198 limitations.

199

200 1. *Laboratory-based research*

201 The primary benefit of laboratory research is that more extraneous factors can be controlled
202 compared to field or remote data collection, including the ability to control factors such as the
203 environment (humidity, temperature) and using ‘gold standard’ laboratory equipment to
204 enhance internal validity (in most cases). These added layers of control allow the researcher to
205 be confident that performance measures are not a result of extraneous factor(s). In most
206 laboratories, temperature is controllable through air conditioning systems, and in most cases,
207 humidity will also be constant. It is advised to keep this consistent during data collection both
208 between and within participant procedures, with records being kept for each experimental
209 session. The main drawback of laboratory research is that the environment is largely artificial,
210 especially in sport where athletes often compete in an environment that is constantly changing
211 (e.g., weather, temperature, typography, anxiety from high-pressured environments). As a
212 result, the findings in rigorously controlled laboratories lose generalisability to sport
213 practitioners (*i.e.*, ecological validity). Moreover, demand characteristics could impact the
214 findings whereby participants may behave differently when being observed (Nichols & Maner,
215 2008).

216

217 2. *Field-based research*

218 Field-based research has become a more common approach within sport and exercise
219 sciences due to the ability to increase the ecological validity of the findings. An area of concern
220 with field-based research is selecting an exercise protocol or using equipment that are valid
221 against laboratory or ‘gold standard’ measurements (Halperin et al., 2018). Exercise protocols
222 such as the multistage 20-m shuttle run test (more commonly known as the “bleep test”) have
223 been shown to correlate to a good level with maximal rate of oxygen output ($\dot{V}O_{2max}$) (Léger
224 & Lambert, 1982; Paliczka et al., 1987; Ramsbottom et al., 1988) making it an appropriate
225 surrogate in the field. Considering physiological measures and blood lactate as an example,
226 analysers were traditionally a large benchtop equipment that was not readily portable. The

227 development of a portable handheld device such as the Lactate Pro 2 (Arkray, Japan) has
228 overcome such issues, and research has shown it to be useable in the field, and importantly,
229 valid (Bonaventura et al., 2015) and reliable (Tanner et al., 2010) against ‘gold standard’
230 laboratory analysers. As a result, the findings in studies using field-based measurements and
231 techniques can then make valid inferences to guide practitioners. Despite some successes, in
232 sport sciences, this is perhaps not implemented as often as it should. One example includes
233 electromyography (EMG), which is commonly used to infer muscle hypertrophy with higher
234 versus lower amplitudes, however, it is unknown if this is a causal relationship (Halperin et al.,
235 2018). The use of valid techniques is an area that sport science could improve to help
236 practitioners make informed decisions with participants from a sports performance, but also a
237 health perspective (Abt et al., 2022).

238

239 Another factor to consider in the field is the lack of control versus laboratory settings, such
240 as weather, temperature, and aerodynamics. This is particularly common if data collection is
241 ongoing during a competition. While these extraneous factors could influence the results and
242 compromise internal validity, particularly if the study design is a crossover design and
243 researchers are attempting to determine changes from multiple different treatments (*e.g.*, a
244 supplement study to assess the impact on exercise performance), it can be minimised by
245 conducting the test at the same time of day, season (*i.e.*, summer vs. winter) and in similar air
246 density (*e.g.*, indoor track cycling). The best approach for this type of research is to measure
247 and describe as much as possible so that the reader can interpret the extraneous factors that
248 might have influenced results. Furthermore, the authors themselves may use the measured
249 variables to apply a correction factor to standardise conditions for test performed on different
250 days. For example, one study investigating caffeine on 100-m running performance measured
251 temperature, humidity, atmospheric pressure and wind speed to standardise measurements
252 (Matsumura et al., 2022). While it may reduce the generalisability of the research, if all can be
253 accounted for, the benefit of field-based studies is the increased level of ecological validity,
254 which in turn, usually means greater impact within the given sport of focus.

255

256 3. *Remote data collection*

257 Since the COVID-19 pandemic in 2020, a contemporary approach has been to collect data
258 remotely due to the obvious constraints on face-to-face contact (Souza et al., 2022). This is
259 unique compared to field-based testing as it requires no observer (*i.e.*, researcher) of the data

260 collection process. In the context of sport and exercise research, this could increase the
261 inclusivity and reduce the carbon footprint of research, as well as opening opportunities for
262 multicentre experiments. For example, one study collected 165 data sessions on a cycle
263 ergometer remotely over a 2-month period using the commercially available software
264 TrainingPeaks™ (Bennett et al., 2021). Given that small sample sizes are common in sport
265 science and can cause issues with power (Abt et al., 2020) and difficulties in translating to real-
266 world settings, remote data collection provides an opportunity to sport and exercise researchers
267 in recruiting larger homogeneous and heterogenous samples. For this to grow and become the
268 norm within the discipline, however, attempts to maintain the reliability and validity must be
269 factored into the study design. Like field-based studies, this includes using consistent
270 methodologies and equipment across participants (*e.g.*, software and equipment), and visual
271 inspections of data collection where possible (*i.e.*, raw data checks, virtual observation of
272 experimental trials taking place). An example of this approach was shown by Matta and
273 colleagues (2022) whereby the reproducibility of a 20-min cycling time-trial was assessed
274 using a home-based protocol. Participants completed two exercise trials using their own home
275 setup on a commercially available software platform (Zwift™) and cycle ergometer (and power
276 meter). This type of approach could be adopted for similar studies, except it would be
277 encouraged that the researcher could watch experimental trials being performed virtually using
278 software (*e.g.*, Microsoft Teams™, Zoom™), which the researchers opted against in their study
279 design (Matta et al., 2022). With this addition, there would be little difference between this
280 approach and both laboratory and field study designs providing no complex data collection is
281 required (*e.g.*, blood sampling, physiological measures).

282

283 **2. Randomisation**

284 Randomisation is considered a critical component of an experimental study that ensures
285 each participant has an equal chance of being assigned to a specific treatment group (in a
286 parallel group study; *e.g.*, 4-week of either beta-alanine or placebo supplementation) or
287 intervention order (in a crossover study; *e.g.*, receiving caffeine first then placebo, or placebo
288 first then caffeine). In performing this allocation entirely randomly, we avoid distorting results
289 due to non-random allocation, which could lead to group differences due to baseline
290 characteristics or identical treatment orders for all participants that, in turn, could bias
291 outcomes.

292

293 Where possible, simple randomisation methods should be preferred. Randomisation is
294 as simple as allocating participants to a treatment group (*e.g.*, beta-alanine or placebo) or order
295 (*e.g.*, “caffeine – placebo” or “placebo – caffeine”) using a coin flip or throwing a dice (Schulz
296 & Grimes, 2002). There is a limitation with simple randomisation in that small sample sizes
297 (<200; (Schulz & Grimes, 2002)), which are common in sport and exercise research, may lead
298 to an uneven number of participants allocated to a particular order, or uneven group sizes.
299 Nonetheless, with increasing sample sizes, this chance is diminished. Block randomisation is
300 also often employed, whereby participants are allocated in an equal ratio (*e.g.*, 1:1 or 2:2) to a
301 treatment group or order. Additionally, studies in sport and exercise science often wish to avoid
302 baseline differences in fitness or performance of participants between groups, and can use
303 stratified randomization to do so (Kang et al., 2008). For example, in a study examining the
304 effects of different training protocols (*e.g.*, high-intensity intermittent exercise vs. continuous
305 exercise) on changes in $\dot{V}O_{2max}$, it would be undesirable for the two training groups to differ
306 significantly in their baseline $\dot{V}O_{2max}$ since those with lower baseline values are likely more
307 susceptible to greater training responses (Støren et al., 2017), regardless of the specific training
308 protocol. Thus, participants could be stratified according to groups based upon their baseline
309 $\dot{V}O_{2max}$. One way this could be achieved is allocating participants to chosen groups of baseline
310 $\dot{V}O_{2max}$ (*e.g.*, 45-50; >50-55; >55-60; >60-65 mL·kg⁻¹·min⁻¹) and within each group, an equal
311 number of participants are randomly allocated to each training condition. This helps ensure that
312 baseline $\dot{V}O_{2max}$ does not differ between groups. An obvious limitation is that the researchers
313 are reliant on equal numbers of participants in each sub-group, and that drop-outs may occur
314 more so in one group than another, which is something that cannot be predicted, and may lead
315 to significant baseline differences. If this occurs, researchers should report the differences in
316 baseline and/or number of dropouts for each condition, and exercise caution in their
317 conclusions.

318

319 The randomisation procedure should be performed by somebody not involved in data
320 collection so that there is no knowledge from the participant or researcher about the
321 intervention being administered, a concept termed allocation concealment. This maintains the
322 blinding of the study should it be necessary (see Section 4. Blinding) and minimises the chance
323 of selection bias, an error that occurs if proper randomisation is not performed resulting in
324 skewed or unrepresentative samples. The person undertaking the randomisation may wish to
325 use free online tools such as Randomization Plans: Never the same thing twice!

326 (jerrydallal.com) or [Research Randomizer](https://www.researchrandomizer.com). As much information as possible as to how the
327 randomisation was performed should be included in any subsequent publication to allow
328 readers to evaluate whether proper randomisation was implemented, of whether possible bias
329 has occurred due to improper randomisation which may occur unwittingly (Schulz & Grimes,
330 2002). We direct the reader towards further reading to gain a more in-depth overview of the
331 methods and techniques for randomisation (Kang et al., 2008; Suresh, 2011).

332

333 *Personal view: In our study on caffeine supplementation and exercise performance (Saunders,*
334 *de Oliveira, et al., 2017), block randomisation was performed by someone not involved in data*
335 *collection so that all possible orders in which participants could receive the supplements (6*
336 *different orders to receive three treatments; caffeine, placebo and control) were balanced*
337 *across 42 participants.*

338

339 **3. Blinding**

340 Participant and researcher expectations about the intervention can significantly affect
341 outcomes during data collection. As a result, within randomised controlled trials, a fundamental
342 decision is to consider whether participants, and those conducting data collection, are blinded
343 to the intervention (*i.e.*, they do not know what interventions are being provided). For example,
344 imagine a research study examining whether caffeine improves 5000-m running time compared
345 to placebo. If a participant is aware they received caffeine, and expect it improves performance,
346 they may change how they perform the trial than when they receive placebo (see for example
347 (Hurst, Schiphof-Godart, et al., 2020)). Researchers would therefore be unable to determine if
348 it was caffeine that improved performance or the change in behaviour. Similarly, if a researcher
349 is aware they are administering caffeine to participants, they may change their behaviour during
350 the trial, such as their body language, words used during administering the caffeine, and type
351 of encouragement given during the trial. As a result, even if the participant is unaware they
352 received caffeine, they may perform the trial differently based on the behaviour of the
353 researcher.

354

355 Blinding in research studies generally takes three forms. First, researchers can use a single-
356 blind design, which involves ensuring only participants do not know which intervention has
357 been administered. This will most likely occur when resources are limited and the person
358 conducting the data collection also needs to administer the intervention. Second, a double-blind

359 intervention can be conducted, in which both the participant and researcher administering the
360 intervention are unaware of what has been administered. In this design, a third-party not
361 involved in data collection disguises both the intervention and placebo so that they are identical
362 in appearance. The researcher would then administer the intervention or placebo to the
363 participant, and both would be unaware what had been administered. Finally, in a triple-blind
364 study, to remove any biases relating to how the data is analysed, the person analysing the data
365 following the completion of data collection can also be unaware of which data is related to the
366 intervention or placebo.

367

368 Blinding is more than just keeping the name of the intervention hidden. Blinding relates
369 to the entirety of the study. This includes, but is not limited to, researchers developing the
370 blinding, witnessing other participants receiving the intervention, perceptual cues of the
371 interventions (*e.g.*, taste, colour, smell) and even physiological responses. The latter can be
372 inherently difficult to blind, especially for some interventions that have noticeable
373 physiological responses, such as sodium bicarbonate that can cause gastrointestinal symptoms
374 (McNaughton, 1992; Saunders et al., 2014). If a participant experiences such effects, then the
375 blinding has failed, and any further data collected is likely biased. It is generally considered
376 that successful blinding ensures the results of the study are not subject to bias. Nonetheless, it
377 is possible that participants experience side-effects related to the active ingredient despite
378 having received a placebo, which may be intrinsically linked to expectation and the information
379 provided regarding the intervention. Blinding success can be assessed by directly asking
380 participants which intervention they think was administered and this data can then be analysed
381 using a number of different tools, such as the Bang's Blinding Index (Bang et al., 2004), which
382 can be used to evaluate the blinding of each intervention (*e.g.*, in a caffeine vs. placebo study,
383 you can determine whether blinding was successful both within the caffeine visit and the
384 placebo visit). Bang's Blinding Index provides a value between -1 and 1, with successful
385 blinding considered between -0.30 to 0.30 (Bang et al., 2010). If blinding was unsuccessful,
386 then blinding may have been compromised and influenced the result of the study, something
387 which researchers may wish to consider upon interpretation of the data.

388

389 It can sometimes be impractical or unfeasible to blind participants to an intervention. It
390 would be impossible to blind a participant to, for example, Normatec compression therapy,
391 physiotherapy, or high-intensity interval training (HIIT) since participants know when they are

392 receiving these interventions. As a result, in such studies, it would be necessary for the
393 researcher to understand participant expectations of the intervention and whether they believed
394 it influenced outcomes. This can be achieved via a questionnaire prior to or post study (*e.g.*,
395 asking participants on a Likert-type scale from 1-5 how much they expect it to affect outcomes),
396 or through post-study interviews, and assessing how much they expected the intervention to
397 influence outcomes (Gurton et al., 2022). The results of this data should be considered during
398 the main analyses and can help determine if they influenced outcomes of the intervention.

399

400 While blinding is often regarded as the gold standard during experimental data
401 collection, sometimes researchers may be interested in understanding the effects of an
402 intervention that has already been shown to be beneficial in blinded studies. This design is
403 called open-label, and is arguably best conducted within the field, where outcomes are of
404 interest under real-world conditions. Given that caffeine has shown to be efficacious during
405 double-blind randomised controlled trials (Grgic et al., 2020), it would be useful to understand
406 if these effects are translated to the field, when participants are aware they have received
407 caffeine. There would be no need to blind participants to what they received, and researchers
408 can understand if caffeine improves performance when given openly.

409

410 *Personal view: We conducted a double-blind, randomised controlled trial to determine if an*
411 *acute dose of dietary nitrate improved 5-km running performance (Hurst, Saunders, et al.,*
412 *2020). We purchased the placebo from the supplier “Beet-IT”, who developed a placebo*
413 *product identical in taste, smell and appearance (Gilchrist et al., 2014). To ensure we*
414 *administered the correct intervention to participants, we asked another person to label one*
415 *“X” and another “Y”. During data collection, participants received the X or Y intervention,*
416 *and we were unaware of whether it was the dietary nitrate or placebo until after the study had*
417 *been completed.*

418

419 **4. Confounding Variables**

420 **1. Observers and researchers**

421 One factor that could impact research is the number and/or sex of observers present at
422 data collection, which could enhance or hinder the participant’s performance. Winchester et al.
423 (2012) reported that ratings of perceived exertion, a subjective measure of how hard the
424 participant believes the exercise is, was reduced with both female and male observers when

425 men were completing a run at ~60% peak running speed. This seemed to be due to the changes
426 in affect scores as these were significantly higher compared to a control trial. In another study,
427 van der Meij et al. (2008) reported that testosterone increased in men by 8% when a woman
428 was introduced to the experimental trials versus a 0.5% change when this was a man. Similarly,
429 24 young male handball players' performance was improved in the presence of female versus
430 male observers. In contrast, the exercise performance of women when in the presence of
431 observers appears hampered, although in some cases it was unchanged. Based on this evidence,
432 researchers should be aware of these potential issues and ensure their research environment
433 limits these impacts. This can be achieved using private research spaces or the use of screens
434 to block the viewing of external individuals within open laboratory spaces.

435

436 The number of people observing exercise can influence exercise performance, whether
437 indirectly or directly observing in the environment (Halperin et al., 2015). One study showed
438 that an audience of fifteen individuals directly watching participants perform a 1-RM bench
439 press improved performance compared to either a passive audience of co-actors (not directly
440 watching; 12.9% increase) or a competitive scenario (fewer direct observers; 2% increase). A
441 factor that might mitigate or enhance these responses is whether the observer is known to the
442 participant, where it has been shown that if this is the case, performance may not change,
443 whereas if the additional person is unknown, a reduction in performance may be found (Guerin,
444 1986). It is worth noting that this change is more likely to be seen for complex tasks (*e.g.*, team
445 sport actions) than simple ones (*e.g.*, capacity or stamina tests). This impact is related to the
446 work of Guerin (1983) who suggested only if the additional audience are not known to the
447 participant would this cause uncertainty and the performance might reduce. This contrasts an
448 early theory such as the generalised drive hypothesis (Zajonc, 1965), whereby a participant's
449 performance will be improved simply through the presence of others. Whilst such theories have
450 since been criticised and many are not discussed in this chapter (for full review see (Strauss,
451 2002)), it is worthwhile for a researcher to consider this within their laboratory research project
452 to reduce the interference of observers in the results. Our recommendation would be that
453 researchers standardise the number and sex of the researchers who will be present at all main
454 data collection sessions throughout a project.

455

456 2. *Verbal encouragement*

457 Verbal encouragement is often seen as a key factor to help participants produce their
458 best effort. However, approximately one third of participants can experience a neutral or
459 negative response to verbal encouragement (Midgley et al., 2018). Unfortunately, there is little
460 evidence to guide recommendations with limited literature to date (Midgley et al., 2018). Of
461 the available evidence, Andreacci et al. (2002) reported that verbal encouragement every 20-s
462 and 60-s improved running performance, whilst no effect was found with encouragement every
463 180-s. Therefore, for maximal efforts verbal encouragement in a frequency of every 20-60-s
464 could assist participants performance. During resistance training, verbal encouragement can
465 improve performance, as Weakley et al. (2020) reported improvements in weight lifted during
466 barbell back squats within a group of 12 semi-professional rugby players. Binboğa et al. (2013)
467 reported that those with low conscientiousness significantly improved their maximal voluntary
468 contraction of the *triceps surae*, but reported no improvements in those with high
469 conscientiousness (9.7% vs. 2.4%). Reasons for discrepancies between Weakley et al. (2020)
470 and Binboğa et al. (2013) may be the sample size ($n = 12$ vs $n = 83$) and the different exercise
471 tests (barbell back squat vs. maximal voluntary contractions). Nonetheless, this suggests that
472 for resistance type exercise, verbal encouragement may be beneficial to produce a best effort
473 performance, however, this might be dependent on the level of conscientiousness within
474 individuals.

475 Although most research has focused on positive feedback, there is a small body of
476 research examining negative feedback. Instead of stating “great effort”, “excellent values” and
477 “looking strong”, when researchers state “you're not trying”, “low values” or “you can do
478 better”, this may improve performance (Halperin et al., 2020). This was hypothesised to be due
479 to participants experiencing some level of anger and exerting greater effort due to the
480 suggestion that their initial efforts were lacking. However, caution is advised since negative
481 feedback might not elicit positive effects in the long-term due to effects on motivation and self-
482 efficacy. Since positive feedback improved performance over no feedback (Halperin et al.,
483 2020), this type of feedback should be preferred. Based on this evidence, it is reasonable to
484 suggest that if verbal encouragement is to be offered it should, at the very least, be standardised.
485 Preferably, the level and frequency of encouragement would also match the type of exercise to
486 achieve the desired effect. Equally, it may be intuitive to match the encouragement based on
487 the level of conscientiousness of participants where possible.

488

489 *Personal view: In our studies (Gough et al., 2018; Gough, Rimmer, et al., 2019), we have used*
490 *multiple approaches for verbal encouragement; however, all have been standardised either to*
491 *time or distance based (for both time-to-exhaustion and time-trial tests) at approximately 60-s*
492 *intervals. In one study, the encouragement was provided every 500 m across a 4-km time-trial.*
493 *We also attempt to standardise the phrases used throughout (e.g., ‘good work, keep going’) by*
494 *using a phrase bank for encouragement.*

495

496 3. *Familiarisation or habituation to the exercise protocol*

497 A key element of rigorous study control is whether participants within a research study
498 are familiarised to the exercise protocol. Familiarisation sessions are usually included in
499 experimental designs to reduce the effect of learning. This is especially important when
500 utilizing untrained samples and participants not familiar to the exercise protocol. Participants
501 in research are often unfamiliar with the exact demands of the exercise task being undertaken
502 (e.g., many cyclists may not be familiar with a 4-km cycling time-trial). Including a session
503 whereby participants perform the exercise task to become familiar with it, researchers can
504 reduce the coefficient of variation and increase test-retest reliability between exercise sessions
505 (Stevens & Dascombe, 2015) which avoids confounding the effect of the intervention with
506 learning. The importance of this is highlighted by the work of Stein and colleagues, who first
507 published their study showing that caffeine improved performance, but were forced to retract
508 their article after discovering results were due to data tabulation error (discussed below) and
509 that the effect was due to a lack of a familiarisation to the exercise protocol and a learning
510 effect (Stein et al., 2020a; Stein et al., 2020b).

511

512 While many researchers perform a solitary familiarisation session in which participants
513 are made familiar with the exercise task, this should not be confused with habituation of a
514 participant to an exercise task. That is, a familiarisation offers participants to become familiar
515 with the exercise protocol, whereas habituation is determined when performance does not
516 change after subsequent visits to the laboratory and can be determined via statistical assessment
517 (e.g., the difference between consecutive tests is very small). It is unclear how many
518 familiarisation sessions are required to attain habituation to an exercise protocol, and will be
519 protocol and participant specific, but this would substantially increase study costs and the
520 number of laboratory visits required for the participant. Nonetheless, we consider it essential
521 that at least one familiarisation is performed prior to initiating the main interventions. There

522 are exceptions wherein it may be appropriate not to include a specific familiarisation protocol.
523 This would be specific to when then the sample population being studied is already familiar
524 with the exercise being undertaken. For example, it is common for rowers to perform regular
525 2000 m rowing tests on a rowing ergometer. Similarly, professional football players will likely
526 perform several YoYo Intermittent Recovery Tests throughout a season to determine exercise
527 capacity. In these situations, it would be appropriate to forgo a specific familiarisation session
528 and simply report that the athletes are well acquainted with the exercise test undertaken.

529

530 *Personal view: Our (BS, FM) research laboratory generally aims to include two*
531 *familiarisation sessions to any exercise protocol to ensure participants are well familiarised*
532 *to the exercise protocol. This is what was required of participants in our study on caffeine*
533 *supplementation and exercise performance (Saunders, de Oliveira, et al., 2017), in which*
534 *trained cyclists performed two familiarisation trials of a simulated time-trial before the main*
535 *intervention session.*

536

537 4. Time of day

538 Several aspects of exercise performance appear to be influenced by the time of day at
539 which they are measured, including strength (Grgic et al., 2019) and endurance (Küüsmaa et
540 al., 2016) exercise, with afternoon and evening performance generally superior to that in the
541 morning. Since the time of day at which individuals exercise can influence exercise
542 performance, when participants attend the laboratory for data collection, researchers should
543 strive to ensure that tests are performed at the same time of day for each participant. Although
544 it may be desirable for all participants to perform exercise when performance appears to be
545 optimised, it is highly improbable that all studies can perform data collection during this very
546 limited late afternoon/early evening timeframe. As such, while it appears unnecessary to
547 require all participants in a study to perform exercise at the same time of day (unless this is a
548 specific aim of the study), each participant should attend the laboratory for data collection
549 within a study at the same time according to their own schedule. Once a participant has
550 performed their first visit, all subsequent visits should then be performed at the same or a
551 similar time to avoid potential influence of circadian variation on the outcome measures.

552

553 *Personal view: In our laboratory, we aim to have participants attend the lab at the best time*
554 *of day that suits them. This might be early morning for some, or late evening for others. For*

555 *example, in our study on caffeine and exercise performance (Saunders, de Oliveira, et al.,*
556 *2017), most participants favoured a morning (06:00 – 08:00) or evening (18:00 – 20:00) start*
557 *due to their working day; this also coincided with their usual training hours. All visits were*
558 *subsequently performed within a ± 1 h period of the initial visit for each participant, since it*
559 *was impossible to always begin at exactly the same time.*

560

561 5. *Dietary control*

562 A person's diet strongly influences their health (Willett, 1994) and exercise
563 performance (Burke & Hawley, 2018). Therefore, it is crucial to monitor or control dietary
564 intake of participants enrolled to the study. While it is common to criticize the lack of control
565 over participants' diet or the way in which such control was carried out, generic criticisms stem
566 from the false belief that all studies should approach dietary control in the same way. To reflect
567 on this, the researcher should not assume that dietary control must be done, but rather evaluate
568 whether there is a need for it and, if so, how to implement it. To develop a good experimental
569 design, there must be clarity with respect to the main research question, namely what will be
570 evaluated, and what the primary outcome (dependent variable *e.g.*, $\dot{V}O_{2max}$, power output,
571 force) is.

572

573 Once researchers have determined if monitoring or controlling diet in the study is truly
574 necessary, the next step is to determine how to do it. It is crucial that the way dietary data is
575 collected and evaluated is valid and appropriate for the study aims. Many options exist
576 including the duplicate diet approach, food consumption recording, 24-h dietary recall, dietary
577 record, dietary history and food frequency questionnaires. Detailing each of these is beyond
578 the scope of the current chapter but those wishing to obtain more specific information about
579 each of these dietary assessment methods are directed towards further reading (Shim et al.,
580 2014; Thompson & Subar, 2017). Where possible, dietary assessment should be performed by
581 the same experienced nutritionist to minimise errors and variation, although some errors
582 between actual and estimated/calculated dietary intake are always likely (Stables et al., 2021).
583 From this point, the researcher should aim to determine whether diet should be monitored,
584 replicated or intervened.

585

586 a) **Monitoring:** In this situation, the researcher does not control the participant's diet in
587 any way, but simply measures it via one of several methods available to monitor the quality,

588 composition, or a specific bioactive compound. For example, a study that aims to evaluate
589 carbohydrate consumption in the week leading up to a sports competition may ask a volunteer
590 to record their food consumption via daily food diaries. Or a study that aims to determine the
591 dietary habits and nutritional status of a distinct group of athletes (*e.g.*, endurance runners or
592 CrossFit® athletes). A consideration here is the observer effect; participants may actively make
593 different choices throughout the study to appear healthier or to be more knowledgeable about
594 food choices, meaning the data may not be an accurate representation of their true diet.

595

596 b) Replication: Participants should be requested to maintain their normal dietary intake
597 and avoid major changes throughout their participation in a study. In situations where changes
598 in diet may cause unwanted changes in the primary outcome, participants should be requested
599 to replicate their diet for a period of between 24-72 h. For example, during a crossover study
600 aiming to determine whether sodium bicarbonate supplementation is ergogenic during a 100-
601 km time-trial on a cycle ergometer, it is possible that carbohydrate intake (and other nutrients)
602 impacts performance, which is the primary outcome for the study. As a result, dietary
603 replication may be advisable before every visit so that this does not influence performance and
604 differences can be attributed to the intervention and not to differences in diet. Replication could
605 occur via one of two ways. Firstly, participants could record their dietary intake during the
606 prespecified period (*e.g.*, 24-72 h pre-test) prior to the first main test, and then be asked to
607 repeat this as closely as possible prior to each subsequent visit. The second option would be to
608 provide participants with pre-prepared food prior to each main test. The former option may be
609 more favourable for studies that do not have funds for food purchases but is reliant on
610 participants repeating their food choices closely which may not always be done. The second
611 option certainly provides more study control since the participants are instructed to eat the food
612 provided by the researchers.

613

614 c) Intervention: This related to when the diet is the independent variable, meaning it is
615 the intervention itself. For example, a study that aims to investigate whether a ketogenic diet
616 impacts the performance of rowers in a 2000-m rowing test compared to a carbohydrate-rich
617 diet. Ideally, since the diet is the intervention, strict control over the diet is desired and all food
618 is provided to the participants. Unfortunately, we do not live in an ideal world and many studies
619 would not have the resources to provide this, and thus dietary advice would likely be provided
620 to participants while dietary monitoring would occur throughout the study to ensure

621 participants are adhering to their respective diets. The frequency and method (see below) via
622 which this information is obtained will depend upon the researchers. Some studies may be more
623 mechanistic and acute, for example investigating whether carbohydrate ingestion alongside
624 beta-alanine supplementation aids in the entry of beta-alanine into the muscle. In this case,
625 participants can be provided with a standardised carbohydrate-rich meal with and without beta-
626 alanine on separate occasions to determine whether there are differences in muscle levels of
627 beta-alanine. In this context, it is necessary that the provided meal is standardised according to
628 carbohydrate (and other nutrients) content.

629

630 Sport science studies often prohibit certain foods and drinks in the day(s) prior to
631 exercise tests, including alcohol and caffeine, to avoid any influence on exercise performance.
632 Alcohol can negatively impact performance (Shirreffs & Maughan, 2006), and while caffeine
633 can positively influence exercise performance (Guest et al., 2021), the quantities found in
634 coffee can vary up to 100% even when the same quantities and brewing methods are applied
635 (Desbrow et al., 2012; Desbrow et al., 2007; McCusker et al., 2003). Therefore, it makes sense
636 to ensure participants do not ingest these prior to their laboratory visits as they may interfere
637 with the outcomes of the study. Since carbohydrate intake is known to impact endurance
638 performance (Bergström et al., 1967; Jensen et al., 2020), it may be desirable to monitor or
639 control for this in the lead up to an exercise task. since it is known that this can impact upon
640 endurance performance. Similarly, a debated topic is whether research participants should
641 perform exercise in a fasted or fed state. As with most of these factors, the choice should depend
642 upon the primary aims of the study. If the aim of a study is to determine whether nitrate could
643 be a useful pre-exercise supplement to improve 16-km cycling time-trial performance in
644 competition, then it makes sense to have participants consume a pre-exercise diet that the
645 participant would regularly have. However, if the study is mechanistic in nature, such as
646 whether nitrate supplementation increases the rate of oxygen consumption during 16-km time-
647 trial cycling, then researchers may wish to have participants exercise in a fasted state as an easy
648 method of dietary control. Nonetheless, results of such a study may not be entirely applicable
649 to a real-world scenario where athletes are likely to ingest a pre-exercise meal.

650

651 *Personal view: In our laboratory, we perform studies with dietary supplements to determine*
652 *their influence on exercise performance. In these studies, we try to be as applicable to the real-*
653 *world as possible, and generally simply ask volunteers to maintain their normal dietary*

654 *patterns throughout their participation in the study. Since diet can influence exercise*
655 *performance, we request that participants record their dietary intake in the 24-h prior to the*
656 *first intervention session and ask them to replicate this as closely as possible prior to the*
657 *subsequent sessions. The participants are still required to perform 24-h dietary records prior*
658 *to these subsequent sessions so that we can analyse how closely these were followed.*

659

660 **5. Exercise control**

661 **1. Prior to main sessions**

662 A key component to a sport and exercise research study is the control of exercise prior
663 to experimental trials, which is important since this may have negative or positive effects on
664 the outcome of the experimental trial. Specifically, exercise close to an experimental trial may
665 lead to carry-over fatigue, which could impact exercise performance when a best performance
666 is required. This is common when the participants studied, for example, are triathletes, who are
667 reported to train at least once per day (Korkia et al., 1994). The solution would be to allow
668 exercise prior to an experimental trial, however, ensure that this is standardised and recorded.
669 As previously discussed, (see Section 3.1.2. Field-based research), commercial software can
670 assist with checking adherence to this approach (e.g., StravaTM). This would be stronger than
671 attempting to make certain populations refrain from exercise 24-48 hours prior to a trial when
672 this is highly unlikely in practice. Monitoring exercise can also allow the researcher to prescribe
673 the exercise, such as the intensity and volume that would minimise the impact on the
674 experimental trial. For example, if the aim of a study was to investigate the changes in muscle
675 glycogen during a 3-hour simulated time-trial, the researcher could instruct participants to only
676 complete exercise that will not deplete glycogen stores in the 24-48 hours prior to the
677 experimental trial which will help minimise the impact of this on the 3-hour simulated time-
678 trial.

679

680 **2. Throughout short-term studies**

681 The longer the duration of involvement in a study, the longer biological variability
682 might influence outcome measures. Biological variability is defined as “non-intervention
683 related processes that cause true scores to change” (Swinton et al., 2018). Parallel group designs
684 somewhat account for this, whereby a separate group of participants complete the trial under
685 control conditions (i.e., without the intervention). However, crossover designs, such as those
686 using acute supplements such as caffeine, do not. In this instance, it is recommended that

687 participants complete all main trials in as short a time period as is feasible to avoid substantial
688 changes in biological variability. In previous work (Gough, Deb, et al., 2019; Gough et al.,
689 2018), participants completed the study within a three-week window to minimise the impact
690 of training adaptations, which are typically studied (or periodised) over an 8–12-week period
691 (Solli et al., 2019). Using a short time frame of approximately 2-4 weeks should allow for the
692 influence of training adaptations to be minimal. Of course, this approach also needs to be
693 balanced with the time frame between each experimental trial. Generally, a time frame of
694 between 2-3 days between experiments trials has been used (Gough, Deb, et al., 2019; Gough
695 et al., 2018) in dietary supplement studies, and this ensures that sufficient recovery is provided
696 for the physiological systems to reach homeostasis bearing in mind both the influence of the
697 exercise and the supplement (Siegler et al., 2012; Stanley et al., 2013). A caveat to this would
698 be the exercise task employed in the study. If the study involves longer duration exercise, such
699 as running a half marathon or full marathon, then a longer period of recovery may be required.
700 However, for longer duration exercise a parallel-group design is usually preferred when there
701 are either carryover effects or repeated bout effects (Bacchieri & Della Cioppa, 2007).
702 Additionally, it is important to note, albeit anecdotally, that participants consenting to research
703 can often see the research study as a chance to change other elements of their behaviour such
704 as nutrition and training (*i.e.*, to begin a health kick). It could also have the opposite effect,
705 whereby participants feel because they are being healthy in the study they can be unhealthy
706 outside of it (*i.e.*, a licensing effect) (Chiou et al., 2011). This makes it vital at the outset to
707 explain to participants that the intervention is not intended to support this and that other than
708 what the intervention intends to change, all else should remain consistent (other than typical
709 daily variation).

710

711 3. *Throughout longer-term studies*

712 With advances in technology, it is now possible to monitor factors such as physical
713 activity, sleep, and training, whereby the latter can even be controlled (or prescribed) for long-
714 term intervention studies. In the example of a 12-week training study, training monitoring can
715 be completed using applications such as Strava™ and TrainingPeaks™. Due to the autonomous
716 nature of commercial applications, there is no longer a need to rely on written logs that can
717 also increase the level of error compared to commercial applications that track work completed
718 through global positioning satellites (GPS), power meters or heart rate, although these can still
719 have small error themselves (Rampinini et al., 2015). These platforms, however, only cover a

720 few sports such as running and cycling and rely on expensive equipment (*e.g.*, power meter).
721 In other sports, written training diaries might be a more practical method through which to
722 monitor external influences over a long-term study due to the incompatibility of commercial
723 applications (*e.g.*, swimming). The use of written logs may be a benefit to the study to help
724 reduce participant attrition as reflection can lead to better adherence of the experimental
725 procedures (Pirodda et al., 2019), although the opposite might also be expected due to more time
726 being dedicated to the study. In respect of that point, strategies to reduce the amount of
727 participant attrition is vital in research, as the procedures are usually logistically difficult and
728 time consuming. Equally, it can lead to issues of internal and external validity through those
729 dropping out from the research would change the outcome of the study (*i.e.*, negative response),
730 yet would not be included in analysis (dropouts are typically excluded) (Barry, 2005). There is
731 a statistical concept called intention-to-treat analysis that suggests including every participant
732 that was randomised to a treatment group or order in the analysis, regardless of incomplete
733 data, and more reading on this can be done elsewhere (Gupta, 2011). To counter the problem
734 of dropouts, researchers may also wish to consider financial incentives and/or frequent
735 reporting points to complete studies that are long term as this has been shown to increase
736 participant adherence (Pirodda et al., 2019). Researchers should report, as a minimum, how
737 many participants were initially recruited and how many dropped out, and best practice would
738 be to attempt to identify why the participants dropped out. If the dropout was due to the
739 intervention than this should be discussed and interpreted to reduce internal validity issues.

740

741 *Personal view: In a study investigating 4-weeks of beta-alanine supplementation on cycling*
742 *performance in trained cyclists (Perim et al., 2022), we wanted to ensure that potential changes*
743 *in training did not influence our results. To do this, we monitored participant's training*
744 *volumes for 4 weeks prior to supplementation, and during the 4 weeks of supplementation, and*
745 *compared the two to ensure there were no differences. This was done using the participant's*
746 *own GPS of preference, the data from which was uploaded to Strava™ from where we could*
747 *have access to all the information regarding training.*

748 **4. DATA COLLECTION**

749 **1. Equipment**

750 Equipment used for data collection should be calibrated according to standards or
751 manufacturer recommendations prior to every use. It is recommended that researchers
752 understand what “normal” values are expected for whatever measurement they are making so
753 that they can immediately identify whether an equipment reading is off. It is always worth
754 keeping records of calibration values as these can be a good way to check if the equipment is
755 working correctly and provides an audit trail for accreditation purposes (*e.g.*, BASES
756 laboratory accreditation). It is important to note that researchers should aim to use the same
757 exercise equipment, not just the same make or model, during repeat testing as there may be
758 subtle variability in outcomes. From our own experience, we found that two different exercise
759 ergometers of the same make and model reported differences of ~3%, which is large enough to
760 mask any changes after administering an intervention. This applies to field-based research as
761 well as the laboratory. For example, if a running test is performed on a grass surface, ensure
762 all subsequent tests are performed on the same surface so that changes in performance are not
763 influenced by different floor surfaces.

764

765 **2. Exercise protocol**

766 **1. Exercise protocol validity**

767 The type of exercise protocol that is chosen in a research study is important and can
768 depend upon the specific aims of the study. Sometimes the choice is straightforward, for
769 example, if the aim is to determine the efficacy of caffeine on 100-m sprint performance, then
770 the exercise test should be a 100-m sprint (Matsumura et al., 2022). However, this choice is
771 not always as easy, for example, if the aim is to investigate the effect of beta-alanine on football
772 (soccer) performance as performance during such activities are numerous and difficult to
773 measure (*e.g.*, it can be difficult to determine what a performance improvement in soccer is).
774 Often, researchers will develop a test that replicates the demands of the activity, which in the
775 case of football is the YoYo Intermittent Recovery Test (Krustrup et al., 2003), a running test
776 consisting of 2 x 20 m runs which 10 s active recovery until exhaustion. Such a protocol should
777 resemble performance during the activity that it is attempting to simulate as closely as possible,
778 an aspect called validity, though there are many types of validity with further reading suggested
779 (Currell & Jeukendrup, 2008). YoYo Intermittent Recovery Test performance is strongly
780 correlated to running trends during match play (Krustrup et al., 2003; Krustrup et al., 2006)

781 making it a good surrogate for match performance. Since Saunders et al. (2012) showed a
782 positive effect of beta-alanine supplementation on YoYo Intermittent Recovery test
783 performance, this can then be extrapolated to suggest that beta-alanine may be effective for in-
784 match football performance.

785

786 Sometimes the choice of an exercise test is to determine the underpinning mechanisms
787 of an intervention. For example, Hill et al. (2007) developed a high intensity cycling capacity
788 test that is performed until exhaustion and limited by muscle acidosis. This makes it an
789 excellent model to determine whether increased muscle buffering capacity (which delays
790 acidosis), achieved via beta-alanine supplementation, can improve performance during
791 exercise limited by acidosis. They showed not that beta-alanine is effective for a sport-specific
792 exercise, but that it can improve performance during exercise limited by muscle acidosis.

793

794 Researchers using exercise capacity tests, in which participants perform exercise at a
795 fixed intensity until no longer tolerable (also called a time-to-exhaustion protocol), are often
796 criticised for not considering the ecological validity of the test (*i.e.*, they do not necessarily
797 replicate a real-world situation). This is particularly true for supramaximal intensities in which
798 the participant is instructed to exercise at an intensity well above their usual maximum,
799 meaning that they will fatigue rapidly. Nonetheless, in addition to providing potential
800 mechanistic insights, for many athletes trying to maintain race pace with the leader, this is a
801 true reflection in an applied setting. For example, in road cycling, an end sprint on a climb
802 would likely be supramaximal and close to a time-to-exhaustion test since the athlete will aim
803 to exert themselves maximally and aim to be completely depleted by the finish line. Thus,
804 knowing how long they could realistically maintain such a high intensity, and how this might
805 be improved, could provide valuable information.

806

807 Some studies in sport science evaluate measures of performance or fatigue during
808 exercise to determine how this differs between, for example, sex or ability (McKay et al., 2022).
809 This could be achieved by using specific exercise protocols replicating real-world competition
810 such as a 100-m running sprint or a 4-km cycling time-trial. It is natural to question whether
811 laboratory measurement of a particular sporting activity truly represents the physiological
812 demands of competition, but studies do exist showing that they may not be different. One study
813 showed that physiological responses to a 5-km cycling time-trial were not different when

814 measured in the laboratory or during a competition (Foster et al., 1993). Some exercise
815 protocols have been developed to measure a specific component of exercise capacity. For
816 example, the 30-s cycling Wingate test, in which participants cycle maximally (all-out) against
817 a fixed resistance for 30 s, was developed to measure muscular power and anaerobic exercise
818 capacity (Bar-Or, 1987; Bar-Or et al., 1977). This test can then subsequently be used to
819 determine differences in anaerobic capacity between athletic groups (*e.g.*, endurance vs. sprint
820 cyclists) or whether a nutritional intervention can improve anaerobic capacity (*e.g.*, sodium
821 bicarbonate supplementation).

822

823 *Personal view: In a study performed by our laboratory, we supplemented participants with*
824 *beta-alanine for 24-weeks to see how much muscle carnosine could be increased and whether*
825 *improvements in exercise performance followed suit (Saunders, Painelli, et al., 2017). We used*
826 *the high intensity cycling capacity test employed by Hill et al. (2007) because they had*
827 *previously shown it to be limited by muscle acidosis and improved by 4 weeks of beta-alanine*
828 *supplementation making it an appropriate model for our study. The aim was not to determine*
829 *whether beta-alanine improved a specific sport, but how closely performance improvements*
830 *mimicked muscle changes.*

831

832 2. Exercise protocol reliability

833 The reliability of an exercise protocol is an important consideration, particularly when
834 considering that many intervention effects may be small. For example, supplementation effects
835 are generally 1-3% (Carr et al., 2011; Hobson et al., 2012). It is, therefore, key that the day-to-
836 day variability in performance during the exercise test is minimal, as it may render the test
837 unable to detect intervention changes. Test-retest studies typically have participants perform
838 the same exercise test on two separate occasions, usually following at least one familiarisation,
839 and under the same strict controlled conditions. The performance difference between sessions
840 is then calculated using metrics such as the coefficient of variation (CV), Pearson's correlation,
841 intraclass correlation 95% limits of agreement or typical error (for more reading see (Currell
842 & Jeukendrup, 2008; Hopkins, 2000; Swinton et al., 2018)). The CV is considered an
843 appropriate statistic, easy to interpret as it is expressed as a percentage since it uses the standard
844 deviation as a percentage of the mean, and allows easy comparison between different exercise
845 protocols (Currell & Jeukendrup, 2008). The higher the CV value, the greater the variation
846 between one visit and the next, which is undesirable. Though there is no specific cut-off limit,

847 CVs above 10% are often considered too high rendering the test inadequate. Such high CVs
848 are generally seen in time-to-exhaustion exercise capacity tests performed at low intensities
849 (Currell & Jeukendrup, 2008; Jeukendrup et al., 1996), though high-intensity capacity tests
850 often show more suitable CVs below 10% (Higgins et al., 2014; Saunders et al., 2013). Time-
851 trial tests generally show excellent reliability (<5%) (Currell & Jeukendrup, 2008; Jeukendrup
852 et al., 1996) meaning they are often the preferred choice for intervention studies. Researchers
853 should also be aware that training status positively influences test-retest reliability (Benton et
854 al., 2013), meaning that less trained participants may exhibit higher variability than is desired.
855 Clinical populations may also show different consistency in performance dependent upon their
856 disorder and the exercise test being employed. Anyone initiating data collection should be
857 aware of the reliability of the exercise protocol being used and the expected changes with the
858 intervention under investigation so appropriate decision-making can be made. Furthermore,
859 protocols with large variability may explain equivocal results in some intervention studies, for
860 example, in which the variation of the exercise protocol will likely have masked the small effect
861 of a dietary supplement.

862

863 *Personal view: We previously sought to employ a time-to-exhaustion cycling protocol*
864 *performed at 75% of peak power output to determine the effects of caffeine supplementation*
865 *on performance. Pilot testing with a handful of cyclists revealed a day-to-day variation of*
866 *approximately 30%, similar to that shown by Jeukendrup et al. (1996), which led us to choose*
867 *a time-trial protocol with a smaller variation of ~3% (Oliveira et al., 2017).*

868

869 **3. Blood sampling**

870 Blood samples are often taken in sport science studies to determine a plethora of
871 measures depending upon the aims of the study. Some blood analytes can be measured almost
872 immediately using standard laboratory equipment, such as blood lactate concentration or pH.
873 Other compounds, such as markers of muscle damage or stress (*e.g.*, creatine kinase or lactate
874 dehydrogenase) may be more complex and require blood samples to be collected and
875 adequately stored (see Sample management below) for posterior analysis using intricate
876 analytical techniques and equipment. As with most factors, there are a number of
877 considerations to be addressed regarding blood sampling, the most important being which
878 blood parameters are being analysed as this will affect where blood will be sampled from (*e.g.*,
879 the arm, finger, earlobe), the type needed (venous vs. arterial vs. arterialised) and the amount

880 required. Sampling at different sites may lead to different values for certain measures. For
881 example, many studies may choose to measure blood lactate from the ear (for example, during
882 rowing exercise), but researchers should be aware that results are not directly comparable to
883 those obtained from the fingertip (Feliu et al., 1999). Participant posture can also influence the
884 measurement of many clinical blood measurements depending on whether the participant is in
885 a seated vs. standing position (Lima-Oliveira et al., 2017; Lippi et al., 2015). The type of blood
886 collected may also modify the measure in question, for example, venous blood provides lower
887 glucagon-like peptide-1 concentrations than arterialized blood in the postprandial (*i.e.*, fed)
888 state (Chen et al., 2018). Nonetheless, since many sport science studies are unlikely to take
889 place in a hospital, venous or capillary blood samples are usually preferable. This may not
890 always be an issue, as in the example of blood pH and bicarbonate, which shows high levels of
891 agreement whether sampled from venous or arterial blood (Ayaz et al., 2021; Kelly et al.,
892 2004), meaning venous blood is an acceptable substitute for arterial for these measurements.
893 To avoid any unwanted variability in blood sampling, researchers should aim to always take
894 blood samples from the same site (which should be researched and chosen based upon the study
895 aims and accessibility) with the participants in the same position (standing, seated or supine).
896 Anecdotally, researchers may wish to familiarize their participants to blood sampling since a
897 fear of needles may artificially increase blood lactate or glucose levels, though this fear is likely
898 to subside after multiple exposures.

899

900 *Personal view: In our studies, we often take venous blood samples with participants seated on*
901 *a bike. To ensure sampling differences are not encountered due to postural differences, despite*
902 *cannulation occurring in a supine position, we then sample blood with participants in a seated*
903 *position.*

904

905 **4. Sample and data management**

906 Participant information in research should be confidential, to ensure that the identities
907 and their associated information is protected. Researchers must follow ethical guidelines to
908 ensure that the data collected is handled appropriately and with respect for participants' privacy
909 so it cannot be linked to specific participants. One way to maintain confidentiality is by
910 assigning a unique identifier to each participant, as opposed to directly using their name or
911 other identifying information. This unique identifier is then used in all future data, samples or
912 notes relating to that particular participant, while the identifying information linking the

913 participant to the unique identifier should be kept separate and safe. Some studies may require
914 complete anonymity to protect participant privacy, particularly when sensitive topics are being
915 studied. This may involve removing any identifying information from participant records and
916 using coding systems or anonymous questionnaires to collect data.

917

918 *1. Sample storage*

919 Biological samples such as blood, muscle, sweat, saliva, or other such samples are
920 sensitive materials with potential risk of and for contamination, meaning they need to be
921 handled and stored with the utmost care. Many countries may have governing policies on this
922 type of collection with specific regulations that researchers must adhere to. One example is the
923 United Kingdom with the Human Tissue Act (2004) ([https://www.hta.gov.uk/guidance-
924 professionals/hta-legislation/human-tissue-act-2004](https://www.hta.gov.uk/guidance-professionals/hta-legislation/human-tissue-act-2004)). Samples should be put into appropriate
925 containers and properly labelled with information containing the unique identifier of the
926 participant, the specific moment of collection and potentially the study to which they belong
927 (e.g., CAF001BS, V1A; this might refer to a specific caffeine study [CAF], participant 001
928 with initials BS, Visit 1 [V1] and the first timepoint of data collection [A]). Samples can then
929 be organised into larger airtight containers such as freezer boxes or plastic bags which are
930 subsequently stored at the appropriate temperature other specific conditions to prevent
931 degradation or contamination. The ideal storage conditions will vary depending on the type of
932 sample and analysis to be performed but are often stored at -20°C or -80°C for long-term
933 storage. Organisation of samples within a freezer or similar (e.g., liquid nitrogen) should be
934 detailed in an inventory using a computerized tracking system or manual logbook, and access
935 should be restricted to authorized personnel only. Samples should be stored until analysed and
936 then disposed of correctly (i.e., according to university or company guidelines regarding
937 disposal of contaminated samples).

938

939 *2. Data management*

940 Data management should be considered a critical component of research as it ensures
941 that the information collected is accurate, reliable, and easily accessible. Laboratory books are
942 an essential tool for researchers to document their experimental methods, observations, and
943 results. Tabulation of data is an important step following data collection as it allows researchers
944 to organize and analyse their data more effectively. It is recommended that researchers tabulate
945 their data immediately (into Excel, for example) following a collection session to avoid losing

946 data. This can also help the researchers to evaluate whether there is any issue in data collection
947 by visually inspecting whether data appear normal. Data can then be backed up to secure online
948 storage networks or to portable drives to ensure that it is saved to multiple locations in the
949 (hopefully unlikely) event that a laboratory book is lost, file becomes corrupted, or somebody
950 steals your computer (a favourite excuse of a final year undergraduate student to gain more
951 time). Online storage networks, such as OneDrive or DropBox, may be particularly favourable
952 since they allow remote access from any device. Researchers should also take care to store
953 participant data securely, such as in a locked cabinet or password-protected computer file.
954 Storage and maintenance of data for an appropriate period are necessary to ensure that the data
955 can be accessed and reviewed for future research or audits. The length of this retention period
956 varies depending on the type of data, funding requirements, and the research area but is often
957 considered to be 5 years for sport and exercise science.

958

959 *Personal view: Each student in our laboratory has their own laboratory book in which they*
960 *are to write down all their results and are strongly encouraged to extract any data file*
961 *immediately and back it up, tabulate all data as soon as possible and back it up to an online*
962 *server. The laboratory book should also be stored in a secure location.*

963 **References**

- 964 Abt, G., Boreham, C., Davison, G., Jackson, R., Nevill, A., Wallace, E., & Williams, M.
965 (2020). Power, precision, and sample size estimation in sport and exercise science
966 research. *J Sports Sci*, 38(17), 1933-1935.
967 <https://doi.org/10.1080/02640414.2020.1776002>
- 968 Abt, G., Jobson, S., Morin, J. B., Passfield, L., Sampaio, J., Sunderland, C., & Twist, C.
969 (2022). Raising the bar in sports performance research. *J Sports Sci*, 40(2), 125-129.
970 <https://doi.org/10.1080/02640414.2021.2024334>
- 971 Andrade, C. (2018). Internal, External, and Ecological Validity in Research Design, Conduct,
972 and Evaluation. *Indian J Psychol Med*, 40(5), 498-499.
973 https://doi.org/10.4103/ijpsym.Ijpsym_334_18
- 974 Andreacci, J. L., Lemura, L. M., Cohen, S. L., Urbansky, E. A., Chelland, S. A., & Duvillard,
975 S. P. v. (2002). The effects of frequency of encouragement on performance during
976 maximal exercise testing. *Journal of Sports Sciences*, 20(4), 345-352.
977 <https://doi.org/10.1080/026404102753576125>
- 978 Ayaz, F., Furrukh, M., Arif, T., Ur Rahman, F., & Ambreen, S. (2021). Correlation of
979 Arterial and Venous pH and Bicarbonate in Patients With Renal Failure. *Cureus*,
980 13(11), e19519. <https://doi.org/10.7759/cureus.19519>
- 981 Bacchieri, A., & Della Cioppa, G. (2007). Experimental Design: Fallacy of “Before-After”
982 Comparisons in Uncontrolled Studies. In A. Bacchieri & G. Della Cioppa (Eds.),
983 *Fundamentals of Clinical Research: Bridging Medicine, Statistics and Operations*
984 (pp. 183-199). Springer Milan. https://doi.org/10.1007/978-88-470-0492-4_8
- 985 Bang, H., Flaherty, S. P., Kolahi, J., & Park, J. (2010). Blinding assessment in clinical trials:
986 A review of statistical methods and a proposal of blinding assessment protocol.
987 *Clinical Research and Regulatory Affairs*, 27(2), 42-51.
988 <https://doi.org/10.3109/10601331003777444>
- 989 Bang, H., Ni, L., & Davis, C. E. (2004). Assessment of blinding in clinical trials. *Control*
990 *Clin Trials*, 25(2), 143-156. <https://doi.org/10.1016/j.cct.2003.10.016>
- 991 Bar-Or, O. (1987). The Wingate anaerobic test. An update on methodology, reliability and
992 validity. *Sports Medicine*, 4(6), 381-394. [https://doi.org/10.2165/00007256-](https://doi.org/10.2165/00007256-198704060-00001)
993 [198704060-00001](https://doi.org/10.2165/00007256-198704060-00001)

- 994 Bar-Or, O., Dotan, R., & Inbar, O. (1977). A 30-second all-out ergometric test - its reliability
995 and validity for anaerobic capacity. *A 30-second all-out ergometric test - its reliability*
996 *and validity for anaerobic capacity.*, 13, 326.
- 997 Barry, A. E. (2005). How attrition impacts the internal and external validity of longitudinal
998 research. *J Sch Health*, 75(7), 267-270. [https://doi.org/10.1111/j.1746-](https://doi.org/10.1111/j.1746-1561.2005.00035.x)
999 [1561.2005.00035.x](https://doi.org/10.1111/j.1746-1561.2005.00035.x)
- 1000 Bennett, S., Tiollier, E., Brocherie, F., Owens, D. J., Morton, J. P., & Louis, J. (2021). Three
1001 weeks of a home-based “sleep low-train low” intervention improves functional
1002 threshold power in trained cyclists: A feasibility study. *Plos One*, 16(12), e0260959.
1003 <https://doi.org/10.1371/journal.pone.0260959>
- 1004 Benton, M. J., Raab, S., & Waggener, G. T. (2013). Effect of training status on reliability of
1005 one repetition maximum testing in women. *J Strength Cond Res*, 27(7), 1885-1890.
1006 <https://doi.org/10.1519/JSC.0b013e3182752d4a>
- 1007 Bergström, J., Hermansen, L., Hultman, E., & Saltin, B. (1967). Diet, muscle glycogen and
1008 physical performance. *Acta Physiol Scand*, 71(2), 140-150.
1009 <https://doi.org/10.1111/j.1748-1716.1967.tb03720.x>
- 1010 Betts, J. A., Gonzalez, J. T., Burke, L. M., Close, G. L., Garthe, I., James, L. J., Jeukendrup,
1011 A. E., Morton, J. P., Nieman, D. C., Peeling, P., Phillips, S. M., Stellingwerff, T., van
1012 Loon, L. J. C., Williams, C., Woolf, K., Maughan, R., & Atkinson, G. (2020).
1013 PRESENT 2020: Text Expanding on the Checklist for Proper Reporting of Evidence
1014 in Sport and Exercise Nutrition Trials. *International Journal of Sport Nutrition and*
1015 *Exercise Metabolism*, 30(1), 2-13. <https://doi.org/10.1123/ijsnem.2019-0326>
- 1016 Binboğa, E., Tok, S., Catikkas, F., Guven, S., & Dane, S. (2013). The effects of verbal
1017 encouragement and conscientiousness on maximal voluntary contraction of the triceps
1018 surae muscle in elite athletes. *J Sports Sci*, 31(9), 982-988.
1019 <https://doi.org/10.1080/02640414.2012.758869>
- 1020 Bonaventura, J. M., Sharpe, K., Knight, E., Fuller, K. L., Tanner, R. K., & Gore, C. J. (2015).
1021 Reliability and accuracy of six hand-held blood lactate analysers. *J Sports Sci Med*,
1022 14(1), 203-214.
- 1023 Burke, L. M., & Hawley, J. A. (2018). Swifter, higher, stronger: What's on the menu?
1024 *Science*, 362(6416), 781-787. <https://doi.org/10.1126/science.aau2093>

- 1025 Carr, A. J., Hopkins, W. G., & Gore, C. J. (2011). Effects of acute alkalosis and acidosis on
1026 performance: a meta-analysis. *Sports Medicine*, 41(10), 801-814.
1027 <https://doi.org/10.2165/11591440-000000000-00000>
- 1028 Chen, Y.-C., Edinburgh, R. M., Hengist, A., Smith, H. A., Walhin, J.-P., Betts, J. A.,
1029 Thompson, D., & Gonzalez, J. T. (2018). Venous blood provides lower glucagon-like
1030 peptide-1 concentrations than arterialized blood in the postprandial but not the fasted
1031 state: Consequences of sampling methods. *Experimental Physiology*, 103(9), 1200-
1032 1205. <https://doi.org/https://doi.org/10.1113/EP087118>
- 1033 Chiou, W.-B., Yang, C.-C., & Wan, C.-S. (2011). Ironic Effects of Dietary
1034 Supplementation: Illusory Invulnerability Created by Taking Dietary Supplements
1035 Licenses Health-Risk Behaviors. *Psychological Science*, 22(8), 1081-1086.
1036 <https://doi.org/10.1177/0956797611416253>
- 1037 Currell, K., & Jeukendrup, A. E. (2008). Validity, Reliability and Sensitivity of Measures of
1038 Sporting Performance. *Sports Medicine*, 38(4), 297-316.
1039 <https://doi.org/10.2165/00007256-200838040-00003>
- 1040 Desbrow, B., Henry, M., & Scheelings, P. (2012). An examination of consumer exposure to
1041 caffeine from commercial coffee and coffee-flavoured milk. *Journal of Food*
1042 *Composition and Analysis*, 28(2), 114-118.
1043 <https://doi.org/https://doi.org/10.1016/j.jfca.2012.09.001>
- 1044 Desbrow, B., Hughes, R., Leveritt, M., & Scheelings, P. (2007). An examination of consumer
1045 exposure to caffeine from retail coffee outlets. *Food Chem Toxicol*, 45(9), 1588-1592.
1046 <https://doi.org/10.1016/j.fct.2007.02.020>
- 1047 Feliu, J., Ventura, J. L., Segura, R., Rodas, G., Riera, J., Estruch, A., Zamora, A., &
1048 Capdevila, L. (1999). Differences between lactate concentration of samples from ear
1049 lobe and the finger tip. *J Physiol Biochem*, 55(4), 333-339.
- 1050 Foster, C., Green, M. A., Snyder, A. C., & Thompson, N. N. (1993). Physiological responses
1051 during simulated competition. *Med Sci Sports Exerc*, 25(7), 877-882.
1052 <https://doi.org/10.1249/00005768-199307000-00018>
- 1053 Gilchrist, M., Winyard, P. G., Fulford, J., Anning, C., Shore, A. C., & Benjamin, N. (2014).
1054 Dietary nitrate supplementation improves reaction time in type 2 diabetes:
1055 development and application of a novel nitrate-depleted beetroot juice placebo. *Nitric*
1056 *Oxide*, 40, 67-74.

- 1057 Gough, L. A., Deb, S. K., Brown, D., Sparks, S. A., & McNaughton, L. R. (2019). The
1058 effects of sodium bicarbonate ingestion on cycling performance and acid base balance
1059 recovery in acute normobaric hypoxia. *J Sports Sci*, 37(13), 1464-1471.
1060 <https://doi.org/10.1080/02640414.2019.1568173>
- 1061 Gough, L. A., Deb, S. K., Sparks, S. A., & McNaughton, L. R. (2018). Sodium bicarbonate
1062 improves 4 km time trial cycling performance when individualised to time to peak
1063 blood bicarbonate in trained male cyclists. *J Sports Sci*, 36(15), 1705-1712.
1064 <https://doi.org/10.1080/02640414.2017.1410875>
- 1065 Gough, L. A., Rimmer, S., Sparks, S. A., McNaughton, L. R., & Higgins, M. F. (2019). Post-
1066 exercise Supplementation of Sodium Bicarbonate Improves Acid Base Balance
1067 Recovery and Subsequent High-Intensity Boxing Specific Performance. *Frontiers in*
1068 *Nutrition*, 6, 155. <https://doi.org/10.3389/fnut.2019.00155>
- 1069 Grgic, J., Grgic, I., Pickering, C., Schoenfeld, B. J., Bishop, D. J., & Pedisic, Z. (2020). Wake
1070 up and smell the coffee: caffeine supplementation and exercise performance—an
1071 umbrella review of 21 published meta-analyses. *British Journal of Sports Medicine*,
1072 54(11), 681-688.
- 1073 Grgic, J., Lazinica, B., Garofolini, A., Schoenfeld, B. J., Saner, N. J., & Mikulic, P. (2019).
1074 The effects of time of day-specific resistance training on adaptations in skeletal
1075 muscle hypertrophy and muscle strength: A systematic review and meta-analysis.
1076 *Chronobiology International*, 36(4), 449-460.
1077 <https://doi.org/10.1080/07420528.2019.1567524>
- 1078 Guerin, B. (1986). Mere presence effects in humans: A review. *Journal of Experimental*
1079 *Social Psychology*, 22(1), 38-77. [https://doi.org/https://doi.org/10.1016/0022-](https://doi.org/https://doi.org/10.1016/0022-1031(86)90040-5)
1080 [1031\(86\)90040-5](https://doi.org/https://doi.org/10.1016/0022-1031(86)90040-5)
- 1081 Guest, N. S., VanDusseldorp, T. A., Nelson, M. T., Grgic, J., Schoenfeld, B. J., Jenkins, N.
1082 D. M., Arent, S. M., Antonio, J., Stout, J. R., Trexler, E. T., Smith-Ryan, A. E.,
1083 Goldstein, E. R., Kalman, D. S., & Campbell, B. I. (2021). International society of
1084 sports nutrition position stand: caffeine and exercise performance. *J Int Soc Sports*
1085 *Nutr*, 18(1), 1. <https://doi.org/10.1186/s12970-020-00383-4>
- 1086 Gupta, S. K. (2011). Intention-to-treat concept: A review. *Perspect Clin Res*, 2(3), 109-112.
1087 <https://doi.org/10.4103/2229-3485.83221>

- 1088 Gurton, W. H., Matta, G. G., Gough, L. A., & Hurst, P. (2022). Efficacy of sodium
1089 bicarbonate ingestion strategies for protecting blinding. *European Journal of Applied*
1090 *Physiology*, 122(12), 2555-2563.
- 1091 Halperin, I., Pyne, D. B., & Martin, D. T. (2015). Threats to internal validity in exercise
1092 science: a review of overlooked confounding variables. *Int J Sports Physiol Perform*,
1093 10(7), 823-829. <https://doi.org/10.1123/ijssp.2014-0566>
- 1094 Halperin, I., Ramsay, E., Philpott, B., Obolski, U., & Behm, D. G. (2020). The effects of
1095 positive and negative verbal feedback on repeated force production. *Physiol Behav*,
1096 225, 113086. <https://doi.org/10.1016/j.physbeh.2020.113086>
- 1097 Halperin, I., Vigotsky, A. D., Foster, C., & Pyne, D. B. (2018). Strengthening the Practice of
1098 Exercise and Sport-Science Research. *Int J Sports Physiol Perform*, 13(2), 127-134.
1099 <https://doi.org/10.1123/ijssp.2017-0322>
- 1100 Harriss, D., Jones, C., & MacSween, A. (2022). Ethical standards in sport and exercise
1101 science research: 2022 update. *International Journal of Sports Medicine*, 43(13),
1102 1065-1070.
- 1103 Higgins, M. F., James, R. S., & Price, M. J. (2014). Familiarisation to and reproducibility of
1104 cycling at 110% peak power output. *J Sports Med Phys Fitness*, 54(2), 139-146.
- 1105 Hill, C. A., Harris, R. C., Kim, H. J., Harris, B. D., Sale, C., Boobis, L. H., Kim, C. K., &
1106 Wise, J. A. (2007). Influence of beta-alanine supplementation on skeletal muscle
1107 carnosine concentrations and high intensity cycling capacity. *Amino Acids*, 32(2),
1108 225-233. <https://doi.org/10.1007/s00726-006-0364-4>
- 1109 Hobson, R. M., Saunders, B., Ball, G., Harris, R., & Sale, C. (2012). Effects of β -alanine
1110 supplementation on exercise performance: a meta-analysis. *Amino Acids*, 43(1), 25-
1111 37. <https://doi.org/10.1007/s00726-011-1200-z>
- 1112 Hopkins, W. G. (2000). Measures of reliability in sports medicine and science. *Sports*
1113 *Medicine*, 30(1), 1-15. <https://doi.org/10.2165/00007256-200030010-00001>
- 1114 Hurst, P., Saunders, S., & Coleman, D. (2020). No differences between beetroot juice and
1115 placebo on competitive 5-km running performance: A double-blind, placebo-
1116 controlled trial. *International Journal of Sport Nutrition and Exercise Metabolism*,
1117 30(4), 295-300.
- 1118 Hurst, P., Schiphof-Godart, L., Hettinga, F., Roelands, B., & Beedie, C. (2020). Improved
1119 1000-m running performance and pacing strategy with caffeine and placebo: a

- 1120 balanced placebo design study. *International Journal of Sports Physiology and*
1121 *Performance*.
- 1122 Jensen, R., Ørtenblad, N., Stausholm, M. H., Skjaerbaek, M. C., Larsen, D. N., Hansen, M.,
1123 Holmberg, H. C., Plomgaard, P., & Nielsen, J. (2020). Heterogeneity in subcellular
1124 muscle glycogen utilisation during exercise impacts endurance capacity in men. *The*
1125 *Journal of Physiology*, 598(19), 4271-4292. <https://doi.org/10.1113/jp280247>
- 1126 Jeukendrup, A., Saris, W. H., Brouns, F., & Kester, A. D. (1996). A new validated endurance
1127 performance test. *Med Sci Sports Exerc*, 28(2), 266-270.
1128 <https://doi.org/10.1097/00005768-199602000-00017>
- 1129 Kang, M., Ragan, B. G., & Park, J. H. (2008). Issues in outcomes research: an overview of
1130 randomization techniques for clinical trials. *J Athl Train*, 43(2), 215-221.
1131 <https://doi.org/10.4085/1062-6050-43.2.215>
- 1132 Kelly, A. M., McAlpine, R., & Kyle, E. (2004). Agreement between bicarbonate measured on
1133 arterial and venous blood gases. *Emerg Med Australas*, 16(5-6), 407-409.
1134 <https://doi.org/10.1111/j.1742-6723.2004.00642.x>
- 1135 Korkia, P. K., Tunstall-Pedoe, D. S., & Maffulli, N. (1994). An epidemiological investigation
1136 of training and injury patterns in British triathletes. *Br J Sports Med*, 28(3), 191-196.
1137 <https://doi.org/10.1136/bjism.28.3.191>
- 1138 Krstrup, P., Mohr, M., Amstrup, T., Rysgaard, T., Johansen, J., Steensberg, A., Pedersen, P.
1139 K., & Bangsbo, J. (2003). The yo-yo intermittent recovery test: physiological
1140 response, reliability, and validity. *Med Sci Sports Exerc*, 35(4), 697-705.
1141 <https://doi.org/10.1249/01.Mss.0000058441.94520.32>
- 1142 Krstrup, P., Mohr, M., Nybo, L., Jensen, J. M., Nielsen, J. J., & Bangsbo, J. (2006). The Yo-
1143 Yo IR2 test: physiological response, reliability, and application to elite soccer. *Med*
1144 *Sci Sports Exerc*, 38(9), 1666-1673.
1145 <https://doi.org/10.1249/01.mss.0000227538.20799.08>
- 1146 Küüsmaa, M., Schumann, M., Sedliak, M., Kraemer, W. J., Newton, R. U., Malinen, J.-P.,
1147 Nyman, K., Häkkinen, A., & Häkkinen, K. (2016). Effects of morning versus evening
1148 combined strength and endurance training on physical performance, muscle
1149 hypertrophy, and serum hormone concentrations. *Applied Physiology, Nutrition, and*
1150 *Metabolism*, 41(12), 1285-1294. <https://doi.org/10.1139/apnm-2016-0271> %M
1151 27863207

- 1152 Léger, L. A., & Lambert, J. (1982). A maximal multistage 20-m shuttle run test to predict
1153 VO₂ max. *Eur J Appl Physiol Occup Physiol*, 49(1), 1-12.
1154 <https://doi.org/10.1007/bf00428958>
- 1155 Lima-Oliveira, G., Guidi, G. C., Salvagno, G. L., Danese, E., Montagnana, M., & Lippi, G.
1156 (2017). Patient posture for blood collection by venipuncture: recall for standardization
1157 after 28 years. *Rev Bras Hematol Hemoter*, 39(2), 127-132.
1158 <https://doi.org/10.1016/j.bjhh.2017.01.004>
- 1159 Lippi, G., Salvagno, G. L., Lima-Oliveira, G., Brocco, G., Danese, E., & Guidi, G. C. (2015).
1160 Postural change during venous blood collection is a major source of bias in clinical
1161 chemistry testing. *Clinica Chimica Acta*, 440, 164-168.
1162 <https://doi.org/https://doi.org/10.1016/j.cca.2014.11.024>
- 1163 Matsumura, T., Tomoo, K., Sugimoto, T., Tsukamoto, H., Shinohara, Y., Otsuka, M., &
1164 Hashimoto, T. (2022). Acute Effect of Caffeine Supplementation on 100-m Sprint
1165 Running Performance: A Field Test. *Med Sci Sports Exerc*.
1166 <https://doi.org/10.1249/mss.0000000000003057>
- 1167 Matta, G., Edwards, A., Roelands, B., Hettinga, F., & Hurst, P. (2022). Reproducibility of 20-
1168 min Time-trial Performance on a Virtual Cycling Platform. *Int J Sports Med*, 43(14),
1169 1190-1195. <https://doi.org/10.1055/a-1848-8478>
- 1170 McCusker, R. R., Goldberger, B. A., & Cone, E. J. (2003). Caffeine content of specialty
1171 coffees. *J Anal Toxicol*, 27(7), 520-522. <https://doi.org/10.1093/jat/27.7.520>
- 1172 McKay, A. K. A., Stellingwerff, T., Smith, E. S., Martin, D. T., Mujika, I., Goosey-Tolfrey,
1173 V. L., Sheppard, J., & Burke, L. M. (2022). Defining Training and Performance
1174 Caliber: A Participant Classification Framework. *Int J Sports Physiol Perform*, 17(2),
1175 317-331. <https://doi.org/10.1123/ijsp.2021-0451>
- 1176 McNaughton, L. R. (1992). Bicarbonate ingestion: Effects of dosage on 60 s cycle ergometry.
1177 *Journal of Sports Sciences*, 10(5), 415-423.
1178 <https://doi.org/10.1080/02640419208729940>
- 1179 Mesquida, C., Murphy, J., Lakens, D., & Warne, J. (2022). Replication concerns in sports and
1180 exercise science: a narrative review of selected methodological issues in the field.
1181 *Royal Society Open Science*, 9(12), 220946. <https://doi.org/doi:10.1098/rsos.220946>
- 1182 Midgley, A. W., Marchant, D. C., & Levy, A. R. (2018). A call to action towards an
1183 evidence-based approach to using verbal encouragement during maximal exercise

- 1184 testing. *Clinical Physiology and Functional Imaging*, 38(4), 547-553.
1185 <https://doi.org/https://doi.org/10.1111/cpf.12454>
- 1186 Nichols, A. L., & Maner, J. K. (2008). The good-subject effect: investigating participant
1187 demand characteristics. *J Gen Psychol*, 135(2), 151-165.
1188 <https://doi.org/10.3200/genp.135.2.151-166>
- 1189 Oliveira, L. F. d., Yamaguchi, G., Painelli, V. S. d., Silva, R. P. d., Gonçalves, L. S.,
1190 Gualano, B., & Saunders, B. (2017). Comprehensive reliability analysis of a 16 km
1191 simulated cycling time-trial in well-trained individuals. *Journal of Science and*
1192 *Cycling*, 6(1), 11-17.
- 1193 Paliczka, V. J., Nichols, A. K., & Boreham, C. A. (1987). A multi-stage shuttle run as a
1194 predictor of running performance and maximal oxygen uptake in adults. *Br J Sports*
1195 *Med*, 21(4), 163-165. <https://doi.org/10.1136/bjism.21.4.163>
- 1196 Perim, P., Gobbi, N., Duarte, B., Oliveira, L. F. d., Costa, L. A. R., Sale, C., Gualano, B.,
1197 Dolan, E., & Saunders, B. (2022). Beta-alanine did not improve high-intensity
1198 performance throughout simulated road cycling. *European Journal of Sport Science*,
1199 22(8), 1240-1249. <https://doi.org/10.1080/17461391.2021.1940304>
- 1200 Pirotta, S., Joham, A., Hochberg, L., Moran, L., Lim, S., Hindle, A., & Brennan, L. (2019).
1201 Strategies to reduce attrition in weight loss interventions: A systematic review and
1202 meta-analysis. *Obes Rev*, 20(10), 1400-1412. <https://doi.org/10.1111/obr.12914>
- 1203 Rampinini, E., Alberti, G., Fiorenza, M., Riggio, M., Sassi, R., Borges, T. O., & Coutts, A. J.
1204 (2015). Accuracy of GPS devices for measuring high-intensity running in field-based
1205 team sports. *Int J Sports Med*, 36(1), 49-53. <https://doi.org/10.1055/s-0034-1385866>
- 1206 Ramsbottom, R., Brewer, J., & Williams, C. (1988). A progressive shuttle run test to estimate
1207 maximal oxygen uptake. *Br J Sports Med*, 22(4), 141-144.
1208 <https://doi.org/10.1136/bjism.22.4.141>
- 1209 Saunders, B., de Oliveira, L. F., da Silva, R. P., de Salles Painelli, V., Goncalves, L. S.,
1210 Yamaguchi, G., Mutti, T., Maciel, E., Roschel, H., Artioli, G. G., & Gualano, B.
1211 (2017). Placebo in sports nutrition: a proof-of-principle study involving caffeine
1212 supplementation. *Scand J Med Sci Sports*, 27(11), 1240-1247.
1213 <https://doi.org/10.1111/sms.12793>
- 1214 Saunders, B., Painelli, V. S., LF, D. E. O., V, D. A. E. S., RP, D. A. S., Riani, L., Franchi,
1215 M., Goncalves, L. S., Harris, R. C., Roschel, H., Artioli, G. G., Sale, C., & Gualano,
1216 B. (2017). Twenty-four Weeks of beta-Alanine Supplementation on Carnosine

- 1217 Content, Related Genes, and Exercise. *Med Sci Sports Exerc*, 49(5), 896-906.
1218 <https://doi.org/10.1249/MSS.0000000000001173>
- 1219 Saunders, B., Sale, C., Harris, R. C., Morris, J. G., & Sunderland, C. (2013). Reliability of a
1220 high-intensity cycling capacity test. *J Sci Med Sport*, 16(3), 286-289.
1221 <https://doi.org/10.1016/j.jsams.2012.07.004>
- 1222 Saunders, B., Sale, C., Harris, R. C., & Sunderland, C. (2014). Sodium bicarbonate and high-
1223 intensity-cycling capacity: variability in responses. *Int J Sports Physiol Perform*, 9(4),
1224 627-632. <https://doi.org/10.1123/ijsp.2013-0295>
- 1225 Saunders, B., Sunderland, C., Harris, R. C., & Sale, C. (2012). β -alanine supplementation
1226 improves YoYo intermittent recovery test performance. *Journal of the International*
1227 *Society of Sports Nutrition*, 9(1), 39. <https://doi.org/10.1186/1550-2783-9-39>
- 1228 Schulz, K. F., & Grimes, D. A. (2002). Generation of allocation sequences in randomised
1229 trials: chance, not choice. *The Lancet*, 359(9305), 515-519.
1230 [https://doi.org/10.1016/S0140-6736\(02\)07683-3](https://doi.org/10.1016/S0140-6736(02)07683-3)
- 1231 Shim, J. S., Oh, K., & Kim, H. C. (2014). Dietary assessment methods in epidemiologic
1232 studies. *Epidemiol Health*, 36, e2014009. <https://doi.org/10.4178/epih/e2014009>
- 1233 Shirreffs, S. M., & Maughan, R. J. (2006). The effect of alcohol on athletic performance.
1234 *Curr Sports Med Rep*, 5(4), 192-196. <https://doi.org/10.1007/s11932-006-0046-8>
- 1235 Siegler, J. C., Marshall, P. W., Bray, J., & Towlson, C. (2012). Sodium bicarbonate
1236 supplementation and ingestion timing: does it matter? *J Strength Cond Res*, 26(7),
1237 1953-1958. <https://doi.org/10.1519/JSC.0b013e3182392960>
- 1238 Solli, G. S., Tønnessen, E., & Sandbakk, Ø. (2019). Block vs. Traditional Periodization of
1239 HIT: Two Different Paths to Success for the World's Best Cross-Country Skier. *Front*
1240 *Physiol*, 10, 375. <https://doi.org/10.3389/fphys.2019.00375>
- 1241 Souza, H. L. R., Bernardes, B. P., Prazeres, E. O. d., Arriel, R. A., Meireles, A., Camilo, G.
1242 B., Mota, G. R., & Marocolo, M. (2022). Hoping for the best, prepared for the worst:
1243 can we perform remote data collection in sport sciences? *Journal of Applied*
1244 *Physiology*, 133(6), 1430-1432. <https://doi.org/10.1152/jappphysiol.00196.2022>
- 1245 Stables, R. G., Kasper, A. M., Sparks, S. A., Morton, J. P., & Close, G. L. (2021). An
1246 Assessment of the Validity of the Remote Food Photography Method (Termed Snap-
1247 N-Send) in Experienced and Inexperienced Sport Nutritionists. *Int J Sport Nutr Exerc*
1248 *Metab*, 31(2), 125-134. <https://doi.org/10.1123/ijsnem.2020-0216>

- 1249 Stanley, J., Peake, J. M., & Buchheit, M. (2013). Cardiac parasympathetic reactivation
1250 following exercise: implications for training prescription. *Sports Medicine*, 43(12),
1251 1259-1277. <https://doi.org/10.1007/s40279-013-0083-4>
- 1252 Stein, J. A., Ramirez, M., & Heinrich, K. M. (2020a). Acute Caffeine Supplementation Does
1253 Not Improve Performance in Trained CrossFit((R)) Athletes. *Sports (Basel)*, 8(4).
1254 <https://doi.org/10.3390/sports8040054>
- 1255 Stein, J. A., Ramirez, M., & Heinrich, K. M. (2020b). Retraction: Stein, J.A. et al. The
1256 Effects of Acute Caffeine Supplementation on Performance in Trained CrossFit
1257 Athletes. *Sports* 2019, 7, 95. *Sports (Basel)*, 8(2), 24. [https://www.mdpi.com/2075-](https://www.mdpi.com/2075-4663/8/2/24)
1258 [4663/8/2/24](https://www.mdpi.com/2075-4663/8/2/24)
- 1259 Stevens, C. J., & Dascombe, B. J. (2015). The reliability and validity of protocols for the
1260 assessment of endurance sports performance: an updated review. *Measurement in*
1261 *Physical Education and Exercise Science*, 19(4), 177-185.
- 1262 Støren, Ø., Helgerud, J., Sæbø, M., Støa, E. M., Bratland-Sanda, S., Unhjem, R. J., Hoff, J.,
1263 & Wang, E. (2017). The Effect of Age on the V`O₂max Response to High-Intensity
1264 Interval Training. *Med Sci Sports Exerc*, 49(1), 78-85.
1265 <https://doi.org/10.1249/mss.0000000000001070>
- 1266 Strauss, B. (2002). Social facilitation in motor tasks: a review of research and theory.
1267 *Psychology of Sport and Exercise*, 3(3), 237-256.
1268 [https://doi.org/https://doi.org/10.1016/S1469-0292\(01\)00019-X](https://doi.org/https://doi.org/10.1016/S1469-0292(01)00019-X)
- 1269 Suresh, K. (2011). An overview of randomization techniques: An unbiased assessment of
1270 outcome in clinical research. *J Hum Reprod Sci*, 4(1), 8-11.
1271 <https://doi.org/10.4103/0974-1208.82352>
- 1272 Swinton, P. A., Hemingway, B. S., Saunders, B., Gualano, B., & Dolan, E. (2018). A
1273 Statistical Framework to Interpret Individual Response to Intervention: Paving the
1274 Way for Personalized Nutrition and Exercise Prescription. *Frontiers in Nutrition*, 5,
1275 41. <https://doi.org/10.3389/fnut.2018.00041>
- 1276 Tanner, R. K., Fuller, K. L., & Ross, M. L. (2010). Evaluation of three portable blood lactate
1277 analysers: Lactate Pro, Lactate Scout and Lactate Plus. *European journal of applied*
1278 *physiology*, 109(3), 551-559. <https://doi.org/10.1007/s00421-010-1379-9>
- 1279 Thompson, F. E., & Subar, A. F. (2017). Chapter 1 - Dietary Assessment Methodology. In A.
1280 M. Coulston, C. J. Boushey, M. G. Ferruzzi, & L. M. Delahanty (Eds.), *Nutrition in*

- 1281 *the Prevention and Treatment of Disease (Fourth Edition)* (pp. 5-48). Academic
1282 Press. <https://doi.org/https://doi.org/10.1016/B978-0-12-802928-2.00001-1>
1283 UKRI. (2023). *Research with potentially vulnerable people*.
1284 [https://www.ukri.org/councils/esrc/guidance-for-applicants/research-ethics-](https://www.ukri.org/councils/esrc/guidance-for-applicants/research-ethics-guidance/research-with-potentially-vulnerable-people/)
1285 [guidance/research-with-potentially-vulnerable-people/](https://www.ukri.org/councils/esrc/guidance-for-applicants/research-ethics-guidance/research-with-potentially-vulnerable-people/)
1286 van der Meij, L., Buunk, A. P., van de Sande, J. P., & Salvador, A. (2008). The presence of a
1287 woman increases testosterone in aggressive dominant men. *Horm Behav*, *54*(5), 640-
1288 644. <https://doi.org/10.1016/j.yhbeh.2008.07.001>
1289 Weakley, J., Wilson, K., Till, K., Banyard, H., Dyson, J., Phibbs, P., Read, D., & Jones, B.
1290 (2020). Show Me, Tell Me, Encourage Me: The Effect of Different Forms of
1291 Feedback on Resistance Training Performance. *J Strength Cond Res*, *34*(11), 3157-
1292 3163. <https://doi.org/10.1519/jsc.0000000000002887>
1293 Willett, W. C. (1994). Diet and Health: What Should We Eat? *Science*, *264*(5158), 532-537.
1294 <https://doi.org/doi:10.1126/science.8160011>
1295 Winchester, R., Turner, L. A., Thomas, K., Ansley, L., Thompson, K. G., Micklewright, D.,
1296 & St Clair Gibson, A. (2012). Observer effects on the rating of perceived exertion and
1297 affect during exercise in recreationally active males. *Percept Mot Skills*, *115*(1), 213-
1298 227. <https://doi.org/10.2466/25.07.05.Pms.115.4.213-227>
1299