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2 **The benefits of strength training on musculoskeletal system health: Practical**
3 **applications for interdisciplinary care**
4

5 **Short title:**

6 **The benefits of strength training**
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45 **ABSTRACT**

46 Global health organizations have provided recommendations regarding exercise for the
47 general population. Strength training has been included in several position statements due to
48 its multi-systemic benefits. In this narrative review, we examine the available literature, first
49 explaining how specific mechanical loading is converted into positive cellular responses.
50 Secondly, benefits related to specific musculoskeletal tissues are discussed, with practical
51 applications and training programmes clearly outlined for both common musculoskeletal
52 disorders and primary prevention strategies.

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54 **KEY POINTS:**

- 55 • Strength training confers unique benefits to the musculoskeletal system in common
56 disorders and in healthy people.
- 57 • The application of mechanical loading must be specific in order to obtain the desired
58 positive adaptation
- 59 • Healthcare professionals should promote strength training among the general
60 population due to its multi-systemic and specific musculoskeletal benefits

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63 **1.0 Introduction**

64 The importance of strength with regard to athletic performance has been highlighted within
65 recent reviews [1, 2]. The benefits of increasing muscular strength include a positive
66 influence on rate of force development (RFD) and power [1, 3, 4], improved jumping [1],
67 sprinting [5] and change of direction (COD) performance [6], greater magnitudes of
68 potentiation [1], and enhanced running economy [7]. Strong evidence supports the notion that
69 maximal strength serves as one of the key foundations for the expression of high power
70 outputs and that improving and maintaining high levels of strength are of utmost importance
71 to best capitalise on these associations [8-13].

72 What appears to be discussed less so is the impact of strength training on musculoskeletal
73 health. This is surprising given that within previous literature it has been highlighted that
74 strength training can reduce acute sports injuries by one third, and overuse injuries by almost
75 half [14]. Furthermore, strength training programmes appear superior to stretching,
76 proprioception training, and multiple exposure programmes for sports injury risk reduction
77 [14]. Malone et al. [15] found that players with a higher relative lower body strength (3
78 repetition maximum [RM] trap bar deadlift normalised to bodyweight) had a reduced risk of
79 injury compared to weaker players. In addition, stronger athletes had a better tolerance to
80 both higher absolute workloads and spikes in load than weaker athletes. Despite its apparent
81 effectiveness for the reduction of injury risk, there is still far less coverage regarding the
82 positive effect of strength training on injury risk or occurrence within the scientific literature,
83 which may be due to its poor integration within musculoskeletal rehabilitation [16] and
84 primary prevention strategies for sports injuries [17, 18]. This is further limited by a poor
85 understanding and knowledge of physical activity guidelines among healthcare professionals
86 [19-21], which provides challenges for its integration into sports medicine practice. Indeed, it
87 is not uncommon for healthcare professionals to recommend “strengthening programmes”
88 using 10 or more repetitions per set without a clear indication of the intensity adopted [22,
89 23]. Although most of resistance training modes have demonstrated improvements in strength
90 in inactive/untrained individuals during the first weeks [24], it must be pointed out that
91 “strengthening programmes” and “strength training” are not the same; hence, they cannot be
92 used interchangeably.

93 Strength training is not an exclusive cornerstone of sports performance or injuries. The World
94 Health Organization (WHO) has provided global recommendations for the general population

95 relevant to the prevention of non-communicable diseases. They recommended at least 150
96 minutes of moderate-intensity aerobic physical activity (3-5.9 metabolic equivalent tasks,
97 METs)[25], with muscle strengthening activities involving major muscle groups on two or
98 more days a week [230-233]. The biological principles underlying these global
99 recommendations rely on the unique multi-systemic and multi-dimensional benefits of
100 exercise [26] (see Figure 1), its inexpensive adoption, and natural human responsiveness [27].
101 To mention the most salient point, recent evidence showed that vigorous physical activity has
102 potential anti-tumorigenic properties [28]. In fact, it is associated with larger reductions on
103 all-cause mortality [25] and cancer mortality [29, 30]. Specifically, resistance training alone
104 was associated with 21% lower all-cause mortality [31]. Furthermore, patients with breast,
105 colorectal, and prostate cancer involved in superior levels of exercise following cancer
106 diagnosis, were associated with a 28-44% reduced risk of cancer-specific mortality, a 21-35%
107 lower risk of cancer recurrence, and a 25-48% decreased risk of all-cause mortality [32, 33].

108 In this narrative review, we focus on the available literature related to strength training and
109 musculoskeletal health, with the aim of providing practical recommendations in line with best
110 practice for healthcare professionals involved in orthopaedic and sports medicine. Clear
111 prescription details will be outlined in order to foster the best possible biological adaptations
112 and thus, facilitate the use of strength training within all populations. In doing so, we will
113 first outline the key principles underpinning mechano-transduction to illustrate how the body
114 converts mechanical loading into cellular responses, before finally providing evidence-based
115 recommendations for the safe interdisciplinary application of strength training across
116 different populations.

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124 **FIGURE 1** Multi-systemic benefits of strength training.



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126 **2.0 Strength, mechano-transduction, and the neuroendocrine system**

127 Strength training has been shown to demonstrate a superior, dose-dependent and safe risk
128 reduction strategy for acute and overuse sports injuries [34]. Information regarding the
129 underpinning qualities of muscular strength development and the interaction of both cellular
130 and metabolic processes in response to specific mechanical loading will first be discussed.
131 Strength training's wide application to improved musculoskeletal tissues, and its role in the
132 regulation and prevention of systemic disorders will then be examined.

133 **2.1 Underpinning factors**

134 The development of muscular strength can be broadly divided into morphological and neural
135 factors [10]. The maximal force generated by a single muscle fibre is directly proportional to
136 its cross-sectional area (CSA) (number of sarcomeres in parallel) [35, 36], and by the muscle
137 fibres' composition [2, 9, 10, 37], specifically, type II fibres (IIa/IIx) have a greater capacity
138 to generate power per unit of CSA, than the relatively smaller type I fibres. Architectural

139 features such as longer fascicle length and the pennation angle also affect the force
140 generating capacity of the muscle. Longer fascicle length allow more force production
141 through an optimal length-tension relationship [10]. The number of sarcomeres in series
142 influences a muscle's contractility and the rate at which it can shorten. As pennation angle
143 increases, more sarcomeres can be arranged in parallel, thus improving the muscle force
144 generating capacity [10]. Greater pennation angles are more common in hypertrophied than
145 in normal muscles. In regards to neural factors, the size principle dictates that motor unit
146 (MU) recruitment is related to MU type, and that MUs are recruited in a sequenced manner
147 based on their size (smallest to largest) [38]. Thus, the availability of high-threshold MUs is
148 advantageous for higher force production. Furthermore, a higher rate of neural impulses
149 (firing frequency) and the concurrent activation of multiple motor units (motor unit
150 synchronization) enhance the magnitude of force generated during a contraction. These,
151 together with an effective neurological system and inter-muscular coordination (i.e.,
152 appropriate magnitude and timing of activation of agonist, synergist, and antagonist muscles)
153 permit maximal force production [2, 9, 10, 37, 39, 40]. The development of these specific
154 features underpinning improved force capacity, is determined by the mechanical stimuli
155 applied to the musculoskeletal system. Indeed, the musculoskeletal system not only enables
156 locomotion and the transmission of forces for functional movements, but also provides
157 protection to vital organs. Furthermore, the musculoskeletal system stores and secretes key
158 substances (e.g., amino acids, glucose, myokines, ions, etc.) that regulate whole body
159 metabolism [41, 42].

160 Given their mechanical role, musculoskeletal tissues are capable of responding and adapting
161 to mechanical forces via a process called mechano-transduction [43]. The body converts
162 mechanical loading into cellular responses, which in turn, promotes structural changes in
163 tissue mass, structure, and quality [44]. For example, an appropriate increase in mechanical
164 loading of skeletal muscle results in an augmented skeletal muscle mass (i.e., increased
165 CSA). The same rules apply for bone and tendon properties, which are in large part,
166 dependent on skeletal muscle-derived mechanical loading [41]. Both acute and chronic
167 mechanical stressors may temporarily compromise the body's "allostasis". This refers to the
168 process by which the body responds to stressors and maintains homeostasis [45, 46], with the
169 neuroendocrine system responsible for regulating the maintenance of an optimal
170 catabolic/anabolic state. Dysregulation induced by allostatic overload has been associated
171 with the breakdown of musculoskeletal tissues, inflammation [47, 48], and delayed tissue

172 healing [49]. The neuroendocrine system plays an important role not only in acute exercise
173 performance, but also in tissue growth and remodelling. Relevant to mechano-transduction,
174 the endocrine system secretes hormones into the circulatory system that are generally
175 categorised as catabolic, leading to the breakdown of muscle proteins (e.g., cortisol), or
176 anabolic (e.g., testosterone), leading to the synthesis of muscle proteins [50]. Muscle protein
177 synthesis, recovery, and adaptation are the results of the dynamic interaction between these
178 anabolic and catabolic hormones [51]. Although several factors such as exercise selection,
179 intensity and volume, nutritional intake and training experience appear to influence the acute
180 testosterone response [50-52], it has been shown that compound exercises, such as
181 weightlifting exercises, squats, and deadlifts, are capable of producing larger elevations of
182 testosterone than isolation exercises [52-54]. Furthermore, programmes characterized by
183 moderate load, high total volume load and short rest periods (i.e. hypertrophy schemes) may
184 produce substantial elevations in total testosterone; thus, reinforcing the importance of
185 specific exercise prescription in order to reach the targeted physiological adaptation [51, 52].
186 Similarly, increases in acute cortisol levels tend to be influenced by high volume programs,
187 and not by typical strength training protocols [51, 55], thus altering the testosterone/cortisol
188 ratio [56, 57].

189 Understanding the coupling of the mechanical stimuli into molecular responses appears vital
190 for regenerative medicine applied to musculoskeletal disorders and for primary prevention
191 strategies in a wide range of health issues and medical specialties. Mechanical forces may be
192 manipulated in such a way that maximise the positive body responses within a predictable
193 physiological timeframe, and the next section includes relevant information for
194 interdisciplinary care.

195 ***3.0 Multi-systemic benefits***

196 Physical inactivity increases the risk of type 2 diabetes, cardiovascular diseases (CVD), colon
197 cancer, postmenopausal breast cancer, dementia, and depression [58-60]. Furthermore,
198 physical inactivity is associated with abdominal adiposity, which may carry the detrimental
199 effects of visceral fat and persistent systemic low grade inflammation [61, 62]. It is suggested
200 that the skeletal muscles counteract the harmful effects of inactivity via release of specific
201 myokines, such as myostatin, leukemia inhibitory factor (LIF), interleukin (IL)-6, IL-7,
202 brain-derived neurotropic factor (BDNF), insulin-like growth factor 1 (IGF-1), fibroblast
203 growth factor 2 (FGF-2), follistatin-related protein 1 (FSTL-1) and irisin [63]. Therefore,

204 contracting skeletal muscles may be capable of releasing protective factors into the
205 circulatory system during exercise. This may then mediate metabolic and physiological
206 responses in other organs, such as the adipose tissue, liver, the cardiovascular system, and the
207 brain [63]. Increased energy expenditure via resistance training can lead to a decrease in
208 abdominal fat and specifically visceral fat, improving the catabolism and hydrolysis of very
209 low-density lipoprotein-triglycerides [61]. These changes in body composition decrease
210 inflammatory products; thus, reducing the risk of developing multiple associated chronic
211 diseases such as type 2 diabetes and CVD [31]. Furthermore, resistance training improves
212 mitochondrial function in skeletal muscles, oxidative and glycolytic enzyme capacity, and
213 glucose homeostasis; thus, leading to decreased blood glucose [64] and improved type 2
214 diabetes symptoms [31, 61]. Also, resistance training is associated with reduced treatment
215 side effects in cancer patient [33, 65, 66]. The anti-tumorigenic effects of exercise appear to
216 be related to the suppression of cancer cells growth, restriction of inflammatory signalling
217 pathways in myeloid immune cells, and regulation of acute and chronic systemic
218 inflammatory responses [28, 67, 68].

219 Further benefits of resistance training include a reduction in anxiety (overall mean effect $\Delta =$
220 0.31) [69] and depressive symptoms, with a moderate effect size of 0.66 (95% CI = 0.48-
221 0.83) [70, 71]. Mental health benefits may be underpinned by the social interactions typically
222 experienced during exercise and by the positive expectations toward exercise [72]. However,
223 alterations in the hypothalamic pituitary adrenal (HPA) axis and in the neural circuitry
224 involved in affective, behavioural, and cognitive processes have been documented in anxiety
225 and depression-related disorders [73]. Although still speculative, strength training may affect
226 the HPA axis through modulation of cortisol activity [74] and may have antidepressant
227 effects through circulation of neurotrophins such as brain-derived neurotrophic factor
228 (BDNF) [26] and growth factors such as the insulin-like growth factor (IGF-1) [75].
229 Considering that sleep disturbance is one of the cardinal symptoms of depressive illness, it is
230 not surprising that chronic resistance training in isolation also improves subjective sleep
231 quality and day-time function, with moderate-to-large effect sizes [76].

232 Furthermore, there is strong evidence that exercise, including strength training, delivered
233 within a biopsychosocial approach, is effective for musculoskeletal pain [77-79]. From a
234 neurobiological perspective, it can strengthen central pain inhibitory pathways and the
235 immune system response to potentially nociceptive stimuli [80-85].

236 In regard to coronary heart disease, progressive resistance training provides improvement in
237 cardiorespiratory function comparable to aerobic training alone. When combined, they offer
238 more substantiated improvements in both fitness and strength [86]. Resistance and aerobic
239 training seem to increase the number of a specific subset of stem cells, broadly referred as
240 circulating angiogenic cells (CAC). This enhances the vascular endothelium regeneration and
241 angiogenesis; thus, improving myocardial perfusion and lowering the risk of cardiovascular
242 diseases [26, 87]. Also, systolic and diastolic blood pressure may significantly be lowered by
243 dynamic and isometric resistance training [88].

244

245 ***3.1 The effect of strength training on cartilage health***

246 The connective tissue that lines the ends of bones in all diarthrodial joints is called articular
247 cartilage. Its role is to support and distribute forces generated during joint loading [89]. The
248 articular surface is covered with hyaline cartilage, which is avascular, firm, yet pliable. It
249 adapts its structure under forces but may recover its original shape on the removal of such
250 forces. Of note, the ability of cartilage to repair is somewhat limited, which is mainly the
251 result of its avascularity [90]. Differences in cartilage morphology between individuals
252 cannot be readily explained by variability in mechanical loading history. It seems that
253 mechanical stimulation does not play a significant role in cartilage regulation, with evidence
254 to suggest that cartilage thickness is strongly determined by genetics [91]. Although it has
255 been demonstrated that immobilisation reduces cartilage thickness (range 5-7%) [92], the
256 adaptive functional ability of human cartilage in relation to exercise does not seem to be
257 linear [91]. Interestingly, Hudelmaier et al. [93] found that thigh muscle CSA (which is a
258 modifiable factor) is a good and independent predictor of cartilage morphology in both young
259 and elderly adults. Similarly, Ericsson et al. [94] showed that lower thigh muscle strength
260 four years after partial meniscectomy was associated with more severe radiographic
261 osteoarthritis (OA) in the medial tibiofemoral compartment of the operated and the
262 contralateral knee eleven years later, suggesting that muscle strength can help to preserve
263 joint integrity.

264 For years, changes in the articular surface have been erroneously deemed the only cause of
265 symptoms of patients suffering of OA. Compelling evidence shows the coexistence of
266 multiple comorbidities such as obesity, cardiovascular diseases, diabetes, and metabolic
267 syndrome in OA patients [21, 95]. Metabolic disturbances, chronic low-grade inflammation,

268 and vascular endothelial dysfunction appear to be important factors in OA development and
269 progression [21, 96]. Consistent with these findings, a negative correlation between knee
270 cartilage volume and the concentration of circulating inflammatory cytokines, such as IL- 6
271 and TNF, as well as C-reactive protein (CRP) has been demonstrated [95]. Therefore,
272 contemporary evidence frames the definition of OA within a biopsychosocial model, in
273 which multi-dimensional aspects modulate inflammatory processes and tissue sensitivity [97,
274 98]. Among these potential factors, recent reviews stated that knee extensor muscle weakness
275 is a risk factor for knee OA [98, 99]. Segal et al. [100] found that thigh muscle strength did
276 not predict incident radiographic, but did predict incident symptomatic knee OA. In contrast,
277 Thorstensson et al. [101] showed that reduced functional performance in the lower extremity
278 predicted development of radiographic knee OA 5 years later among people aged 35-55 with
279 persistent knee pain and normal radiographs at baseline. Pietrosimone et al. [102] found that
280 higher levels of quadriceps strength correlated with higher physical activity in knee OA
281 patients ($r = 0.44$; $r^2 = 0.18$).

282 Clinical guidelines for knee OA recommend strength training as one of the key elements of
283 OA management [98, 103]. Indeed, the systematic review and meta-analysis conducted by
284 Juhl et al. [104] showed that more pain and disability reduction occurred with quadriceps
285 specific exercise than general lower limb exercise (standardized mean difference [SMD] 0.85
286 versus 0.39, and 0.87 versus 0.36, for pain and disability respectively). Strength training
287 should be an integral component of OA management together with education, weight loss,
288 increase of lean mass, and improvement of aerobic capacity [103]. Beyond the
289 aforementioned benefits on pain and disability levels, Bricca et al. [105] showed that loading
290 the knee joint (via strength training) was safe and provided no detrimental effects for articular
291 cartilage in people at increased risk of, or with knee OA. Although the dosage is still unclear
292 [106], potential beneficial mechanisms may be related to stiffening of the pericellular and
293 inter-territorial matrix in response to dynamic loading [107], increased cartilage volume and
294 glycosaminoglycan [105], and the protective role of muscle strength against cartilage loss
295 [108].

296

297 ***3.2 The effect of strength training on bone health***

298 Bone tissue regulates metabolic demands on the skeleton largely through calciotropic
299 hormones (vitamin D₃, parathyroid hormone, and calcitonin) [109]. Secondly, it maintains

300 the structure needed to withstand daily loading. These structural functions are determined by
301 genetic factors as well as adaptation mechanisms to the loading environment, which are
302 mediated by osteoprogenitor cells, including stromal cells, osteoblasts, and osteocytes [110,
303 111]. Osteocytes are believed to be the critical mechanical sensor cells. Their stimulation
304 cannot be derived directly from matrix deformation, as the required magnitude of strains is so
305 high that it would cause bone fracture [112, 113]. Therefore, it appears that mechanical
306 loading induces the dynamic flow of the pericellular interstitial fluid in the lacunar-
307 canalicular system. This seems to contribute significantly to osteocyte mechanotransduction
308 and bone remodelling process [114].

309 Improved bone tissue mass provides higher structural strength and better protection against
310 fractures [91]. Hence, failure to maintain a positive bone adaptation needed to withstand daily
311 loading might be used to define osteoporosis [110]. Indeed, according with Wolff's Law, a
312 sufficient stimulus needs to be applied to the bone tissue to promote a specific magnitude of
313 positive adaptation [115]. Contrary to societal misconceptions, bone responds positively to
314 mechanical loads that induce high-magnitude strains at high rates or frequencies [116-118].
315 Indeed, despite being common advice from healthcare professionals, data showed that regular
316 walking has no significant effect on preservation of bone mineral density (BMD) at the spine
317 in postmenopausal women [119]. In contrast, Watson et al. [120] demonstrated the superior
318 benefits of high-intensity resistance and impact training (HiRIT) compared to a low-intensity
319 exercise program (10-15 repetitions at < 60% 1RM) in post-menopausal women with
320 osteopenia and osteoporosis. Specifically, after a first month of safe transition and
321 familiarization, a supervised HiRIT program was completed over an 8-month period, twice-
322 weekly, for 30-minutes. Resistance exercises included compound movements such as a
323 deadlift, overhead press, and back squat, performed in 5 sets of 5 repetitions at an intensity of
324 80-85% 1RM. Impact loading was applied via jumping chin-ups with drop landings. HiRIT
325 was significantly ($p \leq 0.001$) superior compared to the control group for lumbar spine BMD
326 ($+2.9\% \pm 3.0\%$ for exercise group versus $-1.2\% \pm 2.3\%$ for control; 95% CI 2.1% to 3.6%
327 versus -1.9% to -0.4%) and femoral neck BMD ($+0.1\% \pm 2.7\%$ versus $-1.8\% \pm 2.6\%$; 95%
328 CI -0.7% to 0.8% versus -2.5 to -1.0%) and physical function (lumbar and back extensor
329 strength, timed up-and-go test, 5 times sit to stand test, functional reach test, and vertical
330 jump). Furthermore, it did not increase the risk of vertebral fracture, and had a clinically
331 relevant improvement in thoracic kyphosis [121]. Similar results have been reported in a
332 meta-analysis including 1769 postmenopausal women [122]. Combined resistance and

333 impact training (i.e. jumping, skipping, hopping) are estimated to promote clinically
334 significant gains (almost 1.8 and 2.4%) in hip and spine BMD in postmenopausal women
335 [122]. Considering that in the first few years after menopause women lose up to 5% of bone
336 mass annually, smaller changes may be considered a valuable result to counteract the decline
337 in bone mass during the aging process [123]. This further highlights the effectiveness of
338 progressive resistance training combined with high-impact or weight-bearing exercises in
339 increasing BMD at the femoral neck and lumbar spine. The cumulative body of evidence
340 shows that the greatest skeletal benefits to the spine and hip are provided by progressive
341 resistance training [124, 125] and can be achieved with high magnitude of loading (around
342 80-85% 1 RM), performed at least twice a week, targeting large muscles crossing the hip and
343 spine through multi-joint movements (e.g. squats and deadlifts) [126, 127]. Such intervention
344 may show positive changes after 4 or 6 months, although greater magnitudes are expected
345 when the intervention is continued for more than 1 year. Progressive resistance training,
346 combined with weight-bearing impact training, can be implemented among different
347 populations, with men and premenopausal women showing consistently positive adaptations
348 [123, 128-130].

349 The transition from childhood to adolescence is critical for bone mineral accrual. During this
350 phase, growth hormone (GH) and insulin-like growth factor-I (IGF-I) are major contributors
351 to bone growth [131]. Participation in sports that emphasize weight-bearing, high-impact and
352 multiplanar-impact (e.g., soccer and racquet games) exercises promote peak bone mass and
353 geometry [132]. Exposure to mechanical loading has substantial benefits not only in youth. It
354 also appears to translate to greater bone strength over a lifetime [133], with consequent
355 reduced risk of fracture, as well as potential delay in osteoporosis development [134].
356 Consistently, research has showed that youth athletes exposed to high or unusual impact
357 weight-bearing sports with rapid rates of loading have superior bone mass at loaded skeletal
358 sites compared to non-athletes or athletes in non-weight-bearing or lower impact sports [127].
359 For example, Courteix et al. [135] found that elite pre-pubertal female gymnasts displayed
360 significantly ($p \leq 0.05$) higher BMD at mid-radius (+15.5%), distal radius (+33%), L2-4
361 vertebrae (+11%), femoral neck (+15%) and Ward's triangle (+15%) than swimmers and
362 active peers. This further reinforces how bone mineral accrual responds positively to physical
363 activity and specific sites of impact loading. Collectively, the available data strongly suggest
364 to include exercise that is weight-bearing and characterised by impact loading in youth to
365 promote and maintain bone health over one's lifetime [131].

366 Stress fractures in the lower limb account for 80%–90% of all stress fractures, representing
367 between 0.7% and 20% of all sports medicine injuries [136]. The proposed mechanism
368 underpinning stress fractures appears to be related to an imbalance between the rate of stress-
369 induced micro-fractures and the rate at which bone repairs [136]. Although it is important to
370 recognise their multifactorial pathophysiology, Schnackenburg et al. [137] showed a
371 correlation of impaired bone quality, particularly in the posterior region of the distal tibia,
372 and decreased muscle strength with lower limb stress fractures in female athletes. Clark et al.
373 [138] revealed that lower grip strength correlated with higher risk of upper limb fractures
374 (odds ratio 2.10, 95% CI 1.23 to 3.31) in active young people aged 12 to 16 years. They also
375 showed that muscle strength was positively associated with BMD, BMC, or bone area. Popp
376 et al. [139] analysed competitive distance runners with and without a history of stress
377 fracture. Lower cortical bone strength, cortical area and smaller muscle CSA were present in
378 runners with a history of stress fracture. Hoffman et al. [140] found that military recruits who
379 were one standard deviation below the population mean in both absolute and relative
380 strength, had a five times greater risk for stress fracture than stronger recruits. This is
381 probably related to increased BMD associated with greater strength levels.

382

383 ***3.3 The effect of strength training on tendon health***

384 The tendon is a connective tissue that transmits the force exerted by the corresponding
385 muscle to the skeleton [141]. Its key role is to store, recoil, and release energy while
386 maintaining optimal efficiency in power production [142]. Hence, tendon stiffness (i.e., the
387 slope of the force-elongation relationship or the resistance to deformation in response to an
388 applied force) plays a critical role in athletic performance, stretch shortening cycle (SSC)
389 activities, and movement economy [141]. Changes in tendon stiffness are a consequence of
390 periods of increased mechanical loading. Alterations of the tendon material (i.e., increase of
391 Young's modulus) and morphological properties (i.e., increase in CSA) are the two
392 underpinning mechanisms [143]. Excessive mechanical loading is commonly considered an
393 important factor in the development of tendinopathy, which is an umbrella term that indicates
394 a nonrupture injury in the tendon or paratendon that is exacerbated by mechanical loading
395 [144]. Clinical features are activity-related pain, focal tendon tenderness, and reduced load
396 capacity and performance [145, 146]. A disconnection between tendon structure and
397 symptoms in tendinopathy exists [147, 148]; thus, confirming multi-factorial aspects

398 contributing to its occurrence and persistence [149]. Nonetheless, loading protocols have
399 been shown to be effective in the management of this condition [150, 151]. Evidence-based
400 recommendations for an effective stimulus for tendon adaptation in healthy adults suggest
401 high intensity loading (85-90% of maximal voluntary isometric contraction [MVIC]) applied
402 in five sets of four repetitions, with a contraction and relaxation duration of 3s each, and an
403 inter-set rest of 2-minutes [141]. This has been shown to increase maximal strength, tendon
404 stiffness, Young's modulus, and tendon CSA [141, 143, 152, 153]. Eccentric actions are the
405 most commonly used loading schemes in the management of tendinopathies, despite their
406 non-superiority to other loading programmes [154-157]. The load employed is usually less
407 than the concentric 1RM, which is in contrast with the documented benefits of supramaximal
408 eccentric training stimuli [158, 159]. Similarly, in absence of clear supporting evidence,
409 isometric exercise has recently become the latest debated trend in tendon rehabilitation in the
410 initial phase [160-162]. Overall, key factors such as time under tension and load/intensity are
411 missing in most tendinopathy studies [150, 154, 163], thus making unclear which physical
412 adaptation is targeted and limiting the synthesis regarding optimal doses into evidence based
413 recommendations [22]. In fact, the magnitude and duration of the force application on the
414 tendon appear more relevant than the type of contraction [141]. This highlights the need of
415 adequately designed studies to improve knowledge within this field [23].

416 Achilles tendinopathy (AT) is one of the most common tendinopathies with an incidence rate
417 of 2.35 per 1,000 within the general adult population and a prevalence of 36% among
418 recreational runners [164]. Reduced plantarflexor strength has been recognized to be a
419 significant risk factor of AT [165, 166]. Cross-sectional studies confirm large deficits in
420 plantarflexor torque between AT symptomatic subjects and healthy controls [167, 168].
421 Although it may appear intuitive that strength training could be adopted as primary
422 prevention strategy for reducing the risk of tendinopathies, current literature to support this
423 notion is lacking. A recent systematic review found limited evidence for the efficacy of
424 preventative interventions for tendinopathies [169]. Among the studies examined, strength
425 training was employed with much lighter loads and subsequently higher repetition ranges
426 [170]; thus, not meeting evidence based recommendations for an effective stimulus for the
427 tendon [141, 143]. Therefore, further prospective studies are needed in this area.

428 Loading programmes have been shown to positively enhance structural adaptations among
429 patients presenting with tendinopathy [150, 164]. However, Heinemeier et al. [171] found
430 that renewal of adult core tendon tissue is extremely limited especially following

431 adolescence. Kubo et al. [172] revealed that length and CSA of the patellar tendon correlated
432 with increases in body size during growth, whereas Young's modulus was lower in the pre-
433 pubertal phase compared to junior high school students and adults. Waugh et al. [173]
434 demonstrated that dimensional and maturational aspects of Achilles tendon stiffness were
435 underpinned not only by age, but also by body mass and peak force production; thus,
436 reinforcing the correlation between tendon stiffness and muscular force capacity in childhood
437 and adolescence. In this regard, it should be noted that safe improvements in muscular
438 strength are possible in youth of all ages and stages of maturation with resistance training
439 [174]. Concomitant with a reduction in the number of sport-related injuries [175], this
440 reinforces the importance of engagement in youth athletic development programmes in the
441 pre-pubertal years with continuation throughout the later stages of maturation and into
442 adulthood [176, 177].

443

444 ***3.4 The effect of strength training on muscle health***

445 Skeletal muscles are characterized by the myofibres and connective tissue. The myofibres are
446 responsible for the contractile function of the muscle, whereas the connective tissue supply
447 the structure that binds the individual muscle cells together during muscle contraction [178].
448 Both mechanical and metabolic stress can trigger muscle adaptation and growth [143]. A
449 protein kinase called the mechanistic/mammalian target of rapamycin (mTOR) appears
450 crucial in the pathway through which mechanical stimuli regulate protein synthesis and
451 muscle mass [41]. Morphological factors such as CSA, muscle fibre composition, pennation
452 angle, and fascicle length, are important in force production. Loss of skeletal muscle mass,
453 reduced motor unit (MU) discharge rate, and impaired function is primarily associated with
454 aging. This is defined as either sarcopenia (age-related loss of skeletal muscle mass and
455 function) or dynapenia (age-associated loss of muscle strength that is not caused by
456 neurologic or muscular diseases) [179, 180]. The reduction of MU discharge rate and type 2
457 muscle fibres lead consequently to reduced RFD, which is associated with impaired
458 functional capacity during daily tasks (e.g. balance recovery during tripping) [3, 181, 182].
459 Pijnappels et al. [183] showed that the identification of individuals most at risk of falling
460 could be predicted by their maximal leg press push-off force level. In older adults, lower
461 muscle strength is also associated with an increased risk of dementia [184], loss of
462 independence, and mortality [185-188]. However, the rate of strength decline is dependent on

463 age and physical activity levels. Indeed, individuals participating in strength training can
464 significantly attenuate the loss of muscle mass and strength, and their undesirable
465 consequences [189]. Strong evidence suggests that an appropriately designed resistance
466 training program for older adults should include an individualized and periodized approach
467 working toward 2-3 sets of 1-2 multi-joint exercises per major muscle group, achieving
468 intensities of 70-85% of 1RM, 2-3 times per week [126]. Strength training is a feasible and
469 effective strategy to counteract muscle weakness [190], physical frailty, age-related
470 intramuscular adipose infiltration, decline in physical function, risk for falls, and reduction in
471 CSA [189, 191]. These benefits are underpinned by the ability of strength training to
472 countermeasure age-related changes in muscle and central nervous system function.
473 Specifically, strength training is highly effective in improving MU discharge rate, reducing
474 loss of type 2 fibres, enhancing RFD and muscle strength, thus explaining the functional
475 benefits in the older population, especially in frail elderly [3, 181].

476 Overall, strength training increases neural drive, intermuscular coordination, myofibrillar
477 CSA of Type I and II fibres, lean muscle mass, and pennation angle [2, 10, 11]. Not
478 surprisingly, primary prevention strategies recommend the employment of strength training
479 for the reduction of acute sports injuries [15, 34]. Among these, muscle injuries are very
480 common in sports [34, 192], constituting 31% of all injuries in elite football [193]. For
481 example, the Nordic hamstring exercise (NHE) (i.e., a form of supramaximal eccentric
482 loading) has been shown to significantly reduce the risk of hamstring injuries [192, 194-196],
483 with long-term benefits associated with increases in fascicle length and improvements in
484 eccentric knee flexor strength [197]. The systematic review and meta-analysis conducted by
485 van Dyk et al. [198] showed that programmes including the NHE reduced hamstring injuries
486 by 51% in athletes across multiple sports. Zouita et al. [199] showed that strength training
487 reduced the risk of injury in elite young soccer players during one season (estimated total
488 injury rate per 1,000 hours of exposure were: 0.70 for the experimental group and 2.32 for the
489 control group). Of note, approximately 50% of the total injuries sustained were classified as
490 “muscle strains”; thus, demonstrating the protective role of strength training on muscle
491 tissues. Although not thoroughly consistent with strength training prescription over the study
492 period, Harøy et al. [200] showed that a single exercise with different levels of targeting the
493 adductors, reduced the prevalence and risk of groin injuries in semi-professional Norwegian
494 football players by 41%. Considering the economic burden of muscle injuries in elite settings
495 (e.g., a single hamstring injury resulting in ~17days lost from training and competition is

496 estimated to cost about €280,000 in elite soccer clubs) [197] and the importance of muscle tissue health for players' availability and
 497 performance, implementation of accurate strength training schedule during the season appears vital. A summary of the benefits for various
 498 musculoskeletal tissues and disorders are depicted in Table 1.

499 **TABLE 1** Summary of benefits for various musculoskeletal tissues and disorders associated with strength/resistance training

Musculoskeletal tissue	Function	Potential beneficial mechanisms	Specific recommendation	Examples of application for common related conditions
Cartilage	Support and distribution of forces generated during joint loading	Stiffening of the pericellular and inter-territorial matrix Increase of cartilage volume and glycosaminoglycan Protection against cartilage loss	Specific exercise for targeted area appears relevant Inclusion in multidimensional care management Potential benefits	Knee Osteoarthritis Joint loading exercises Optimal programme characteristics not identified yet

			associated to increased CSA	Recommended frequency being 3 times weekly with a duration of at least 12 supervised sessions
Bone	Regulation of metabolic demands Structural maintenance to withstand loading	Increase of bone mineral density, bone mineral content, and bone area	To target large muscles Safe transition towards high loads ($\geq 80\%$ 1RM) Familiarisation with movement patterns Combination with impact loading exercises	Osteopenia and osteoporosis 5 sets of 5 repetitions, maintaining an intensity of 80-85% 1 RM performed at least twice per week

Tendon	<p>Force transmission</p> <p>Storage, recoil and release of energy</p>	Increase of tendon stiffness, Young's modulus and tendon cross sectional area	<p>To adopt muscle contraction intensities higher than 70% of MVC or RM</p> <p>Type of contraction (isometric, concentric, eccentric) not relevant</p> <p>Longer durations (≥ 12 weeks) more effective</p>	<p>Reduction of tendon stiffness and Young's modulus</p> <p>5 sets of 4 repetitions with high intensity loading (85–90% MVIC) with a contraction and relaxation duration of 3 s each, and an inter-set rest of 2 min. To be performed 3 times per week</p>
Muscle	Contraction to produce force and	Increase of myofibrillar cross sectional area (CSA) of Type I/II fibers, lean muscle mass,	Individualised and periodised approach	Sarcopenia

	motion	fascicle length and pennation angle	Multi-joint exercise per major muscle group in elderly Type of contraction relevant for muscle fibers architectural adaptations	2–3 sets of 1–2 multi-joint exercises per major muscle group, with intensities of 70–85% of 1RM, 2–3 times per week
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500 CSA (cross sectional area), RM (repetition maximum), MVC (maximal voluntary contraction), MVIC (maximal voluntary isometric contraction)

501

502

503 **4.0 Strength training: Practical applications**

504 Researchers have challenged the existence of “non-responders” to exercise. Positive
505 adaptations are influenced by multidimensional aspects such as genetic factors, fitness level,
506 training history, nutritional intake, psychological and social states, sleep and recovery, age,
507 weight, and prescribed training workload [27] and therefore the magnitude of adaptations
508 between individuals may differ. Thus, strength training prescription should begin with an
509 accurate subjective and objective examination. This investigates training and injury history,
510 general health status, comorbidities co-existence, single-joint and multi-joint strength
511 evaluation and movement pattern analysis relevant to the potential proposed exercise
512 programme. Clinical tools such as questionnaires and outcome measures may be
513 implemented in the subjective examination to more accurately detect and discuss the
514 significant aspects that may negatively counteract the expected positive adaptations and can
515 be administered at specific timeframes at the judicious discretion of healthcare professionals.
516 For example, specific questionnaires and outcomes measures can be adopted to monitor sleep
517 [201] and stress levels [202, 203] over the course of an intervention. This transdiagnostic
518 approach attempts to understand commonalities and shared mechanisms among different
519 multidimensional aspects and to identify any adverse responses to the planned intervention
520 that may be driven by such factors [204]. This enables a stratified model of care (i.e.
521 personalised medicine) to maximise treatment-related benefits, reduce risk of adverse events
522 and increase healthcare efficiency [205] (see examples in figure 2,3,4).

523

524 **** Insert Figure 2,3,4 about here ****

525

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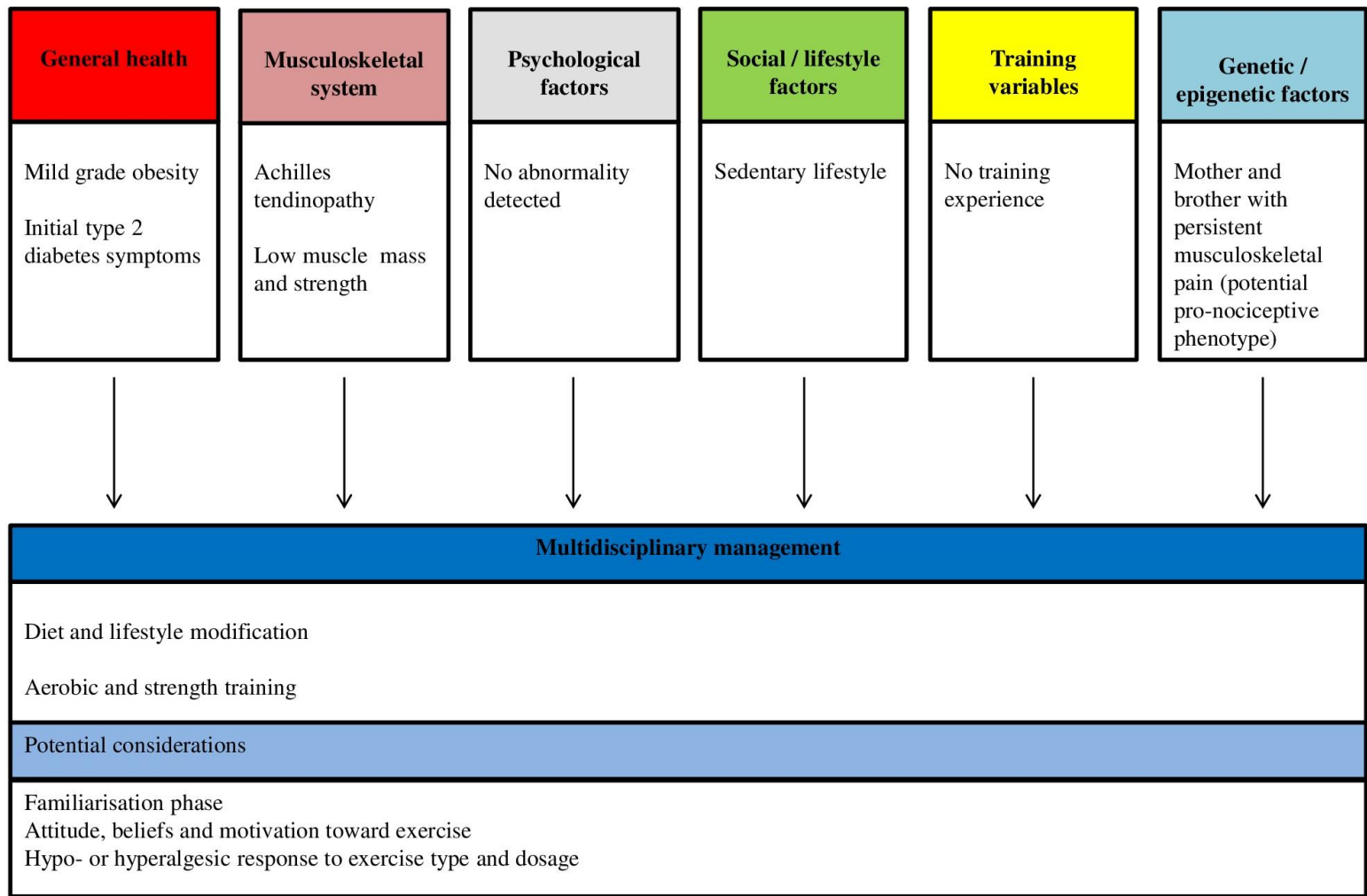
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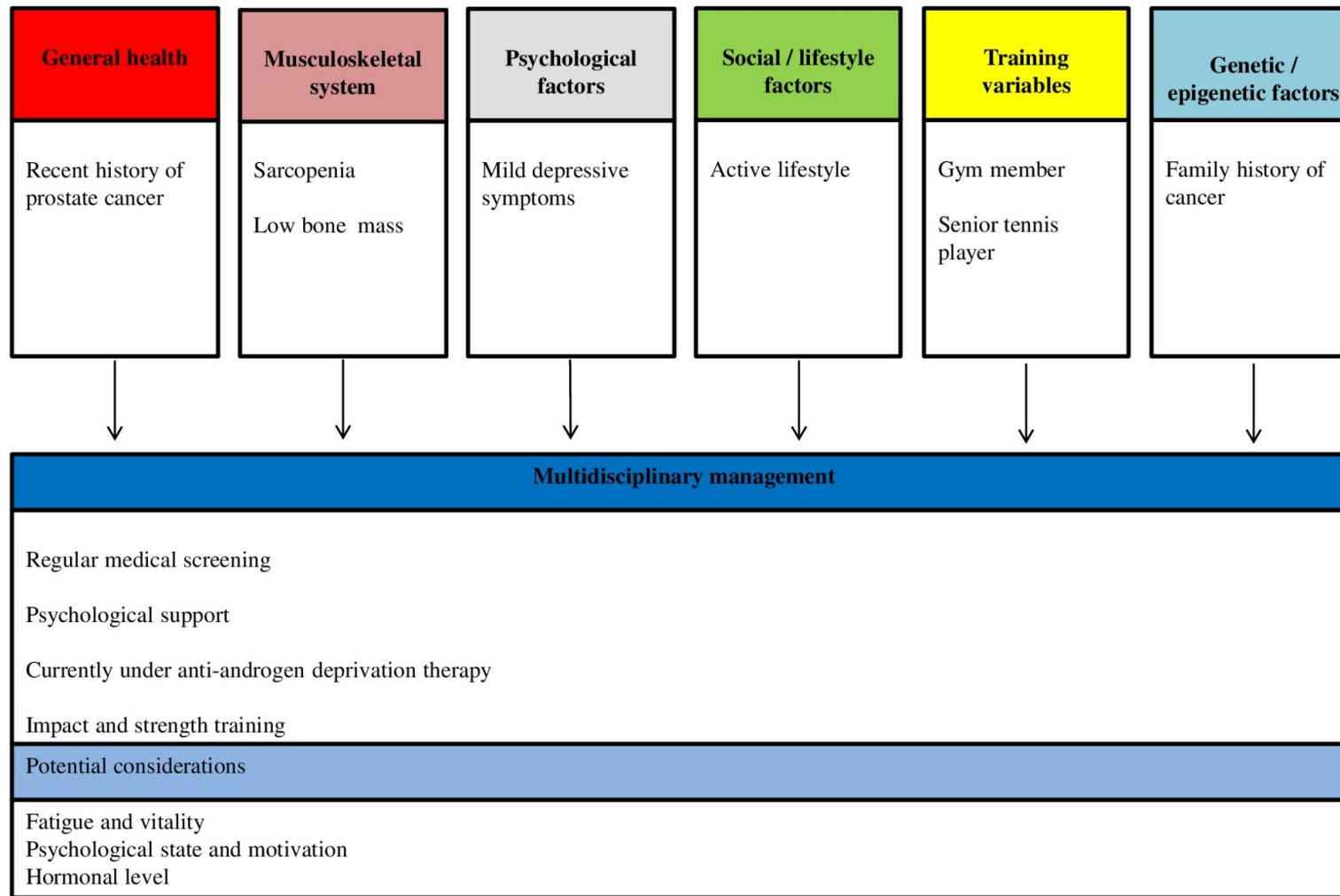
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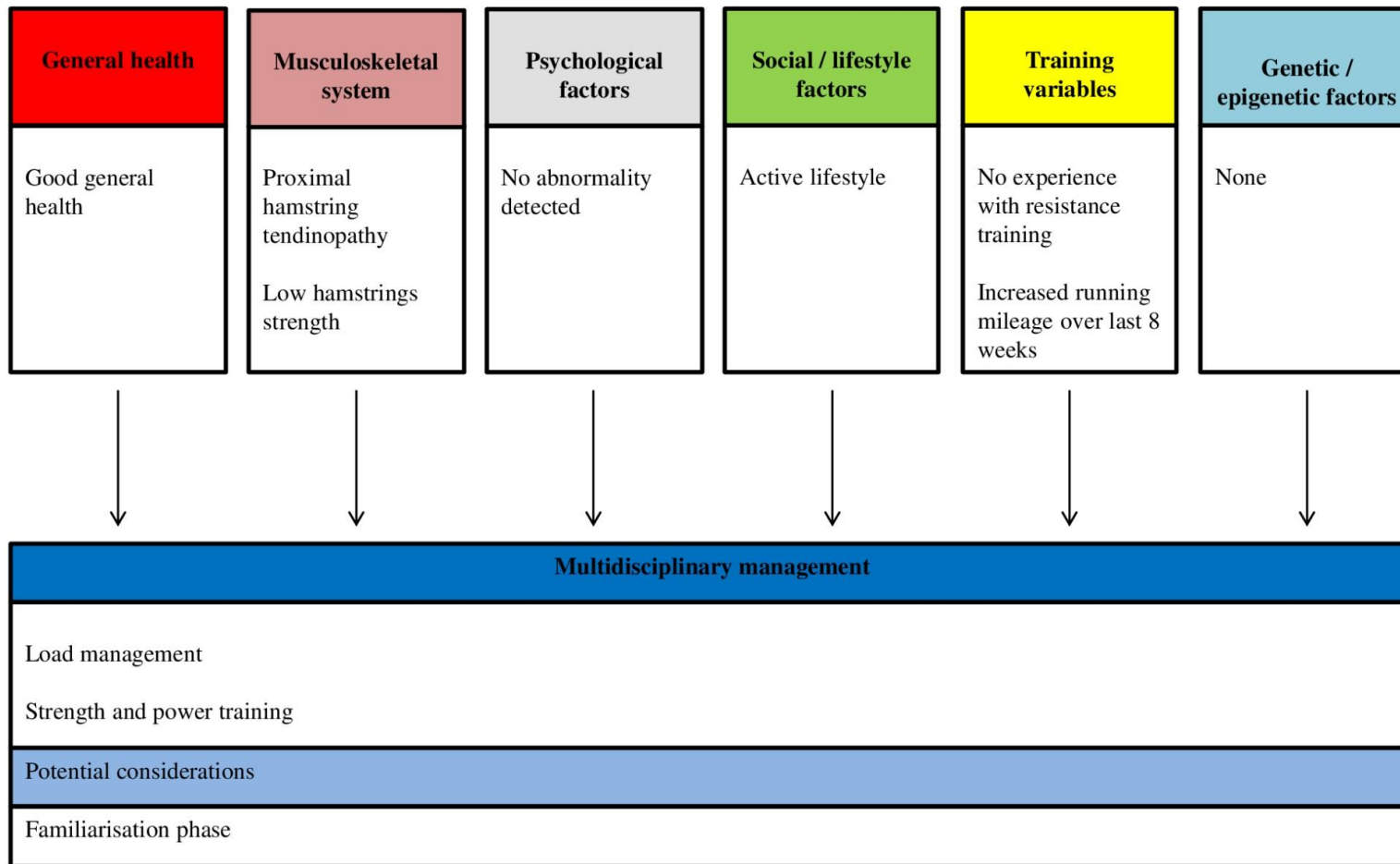
533 **Figure 2** Profile of a middle-aged man with mid-portion Achilles tendinopathy

534



535

536 **Figure 3** Profile of an older man (73 years old) presenting with sarcopenia and a recent history of prostate cancer



537

538 **Figure 4** Profile of a young runner (19 years old) with proximal hamstring tendinopathy preparing for the Marathon

539

540 This process allows a more complete understanding of the person, his/her past and current
541 exposure to loading activities, quality of life, beliefs and attitude towards exercise, relevant
542 impairment in mobility, potential site of loading, adequate skeletal muscle trophism and/or
543 isolated strength deficits that may impair rapid exposure to high-load exercises; thus,
544 requiring a period of familiarization and anatomical adaptation via adoption of different
545 loading schemes. For example, in untrained individuals sensitive to spinal axial loading, who
546 cannot tolerate large external loads, bilateral exercises, such as the back squat can be
547 confidently substituted with unilateral exercises due to similar effectiveness in lower-body
548 strength development, despite relative lower external loading [206]. When the goal is to elicit
549 alterations in skeletal muscle hypertrophy in untrained individuals, current literature [24, 207-
550 209] suggests to train with a high level of effort, irrespective of load. Whereas momentary
551 failure is important during low load training to capitalise on muscular adaptations, this does
552 not provide any additional benefits when training at high resistance training loads. Hence,
553 lighter loads can be initially lifted until failure to maximise MU recruitment, increase muscle
554 size and strength (to a certain extent). With gradual training exposure and increasing
555 resistance training experience, these can be progressed to higher load-lower repetition
556 schemes without momentary failure, thus providing heightened neural impulses to maximise
557 strength gains [208, 210-212].

558 Global recommendations suggest strength training should be performed two or more days per
559 week [230-233]. Maximal strength can be defined as the upper limit of the neuromuscular
560 system to produce force. Force production against an external resistance is an essential
561 trainable ability [213]. It must be noted that in untrained individuals almost any resistance
562 training exercise programme, load and method may increase strength, which is more likely
563 attributable to neural adaptations in response to the new training stimulus [2, 24, 212, 214,
564 215]. However, progressive overload stimuli appear essential to promote further strength
565 adaptations in more experienced individuals [24, 214][234]. For these current evidence
566 indicates that prescription of maximal strength training should involve a load (or intensity) of
567 80-100% of the participant's one RM, utilizing approximately 1-6 repetitions, across 3-5 sets,
568 with rest periods of 3-5 minutes, and a frequency of 2-3 times per week [234]. This implies
569 that loads are determined by percentages of 1RM, with testing potentially challenging when
570 working with load compromised patients and/or pain interference. Therefore, the adoption of
571 an auto-regulated approach (AR), which is based on RM training zones, rate of perceived
572 exertion (RPE) and repetitions in reserve (RIR) [216, 217], may appear more feasible and

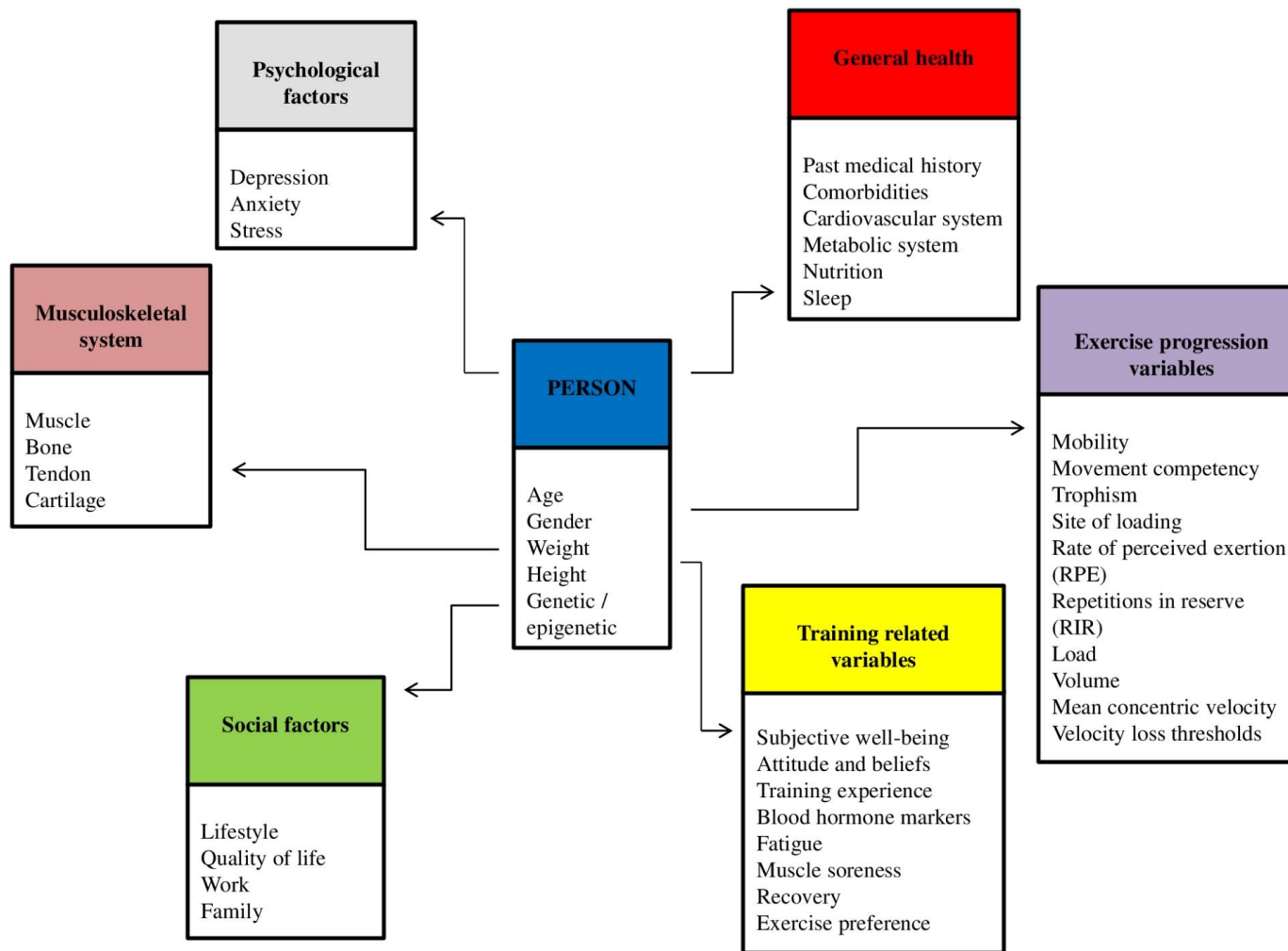
573 clinically advantageous throughout the training cycle. This also accounts for fluctuations in
574 strength capabilities across a training mesocycle [216, 218], which can be influenced by the
575 aforementioned multidimensional aspects. In experienced individuals RPE/RIR scale can be
576 used as a method to assign daily training load, aid in session to session load progression, and
577 monitor individual rates of adaptation [216, 219]. Assessment of movement velocity may
578 also be another valid alternative used to estimate the percentage of loading [220, 221]. This
579 exploits the inverse linear relationship between load and mean concentric velocity (MCV).
580 Indeed, providing that maximal concentric effort is applied during movement, MCV will
581 decrease as magnitude of load increases, thus allowing estimation of relative training loads
582 (%1RM) monitoring movement velocity [222]. In addition, different velocity loss (VL)
583 thresholds across repetitions performed within a set may be also adopted to dictate
584 mechanical and metabolic stress, hormonal responses and neuromuscular fatigue, thus
585 inducing different adaptations. Small to moderate VL threshold (i.e. <20%) are recommended
586 to maximise strength gains in resistance-trained individuals [223, 224]. For clarity of
587 information, example of loading schemes for strength training are depicted in Tables 2.
588 Common subjective and objective variables that contribute to programming and progression
589 decision making are illustrated in Figure 5.

590

591 **Table 2** Suggested strength training variables when employing the traditional percentage
592 fixed loading program (TL) or the auto regulated training (AR).

PROGRAM	REPETITIONS	SETS	LOAD		REST	FREQUENCY
TL	1-6	3-5	@80-100% 1RM		3-5 minutes	2-3 / week
PROGRAM	RM ZONE	SETS	RPE 0-10	RIR	REST	FREQUENCY
AR	1-6	3-5	8-10	0-2	3-5 minutes	2-3 / week

593 TL (traditional loading), AR (auto regulated training), RM (repetition maximum), RPE (rate
594 of perceived exertion), RIR (repetitions in reserve)



595

596 **Figure 5** Graphical representation of common subjective and objective variables that contribute to programming and progression decision
 597 making in strength training

598 **Table 3** Example of a potential strength training session for postmenopausal women with low bone mass (performed at least twice per week for
 599 an ideal duration of at least one year). The length of each phase, exercise selection and the progressions are chosen in accordance to the
 600 participant's weekly evaluation.

Phase 1 - Familiarisation	Exercise	Fixed loading prescription	Auto-regulated training prescription	Impact loading
Training aim				
To ensure safe transition to high-intensity load	Goblet Squat >> Split Squat Romanian Deadlift Box Squat Overhead press >> Press-up	1 sets of 12 repetitions of ~50-60% 1RM 2 min inter set rest	1 sets of 12 RM with RPE 4-6 and RIR 4-5 1 min inter set rest	3 repetitions x 4 sets Snap-downs >> jump to box >> standing broad jump >> depth land >> drop jump

To familiarise with exercises and movement patterns	Bench Press Seated Row >> Bent Over Rows			
Phase 2 – Strength endurance emphasis	Exercise	Fixed Loading prescription	Auto-regulated training prescription	Impact loading
Training aim				
To increase muscle mass, strength and musculotendinous stiffness To facilitate safe	Split Squat >> RFESS >> Box Squat >> Trap-bar Deadlift Romanian Deadlift	3 sets of 8-12 repetitions of ~60-75% 1RM	3 sets of 10RM with RPE 6-7 and RIR 2-3	3 x 20 cm depth land during the first 6 inter-set rest periods

transition to strength training emphasis	Overhead Press>> Press-up or Bench Press Seated Row >> Bent Over Rows	1-2 min inter set rest	2 min inter set rest	2 broad jump during the last 6 inter-set rest periods
Phase 3 – Strength emphasis	Exercise	Fixed Loading prescription	Auto-regulated training prescription	Impact loading
Training aim				
To increase muscle mass, strength, rate of force development and musculotendinous stiffness	Trap-bar Deadlift			4 countermovement jumps during the first 4 inter-set rest periods
To improve motor unit discharge rate	Romanian Deadlift	4 sets of 5 repetitions of > 85%	4 sets of 5RM with RPE 8-9 and	

<p>To reduce loss of type II fibres</p> <p>To increase bone mass, bone mineral content and bone mineral density</p>	<p>Overhead Press or Bench Press</p> <p>Bent Over Rows</p>	<p>1RM</p> <p>3-5 min inter set rest</p>	<p>RIR 1-2</p> <p>3-5 min inter set rest</p>	<p>3 x 3 hurdles jump during the last 4 inter-set rest periods</p>
<p>RM (repetition maximum), RPE (rate of perceived exertion), RIR (repetitions in reserve), RFESS (rear foot elevated split squat); >> = progress to these exercises during next cycle or perform these instead/if preferred and patient/client is competent</p>				

602 Frequency and duration of a strength training program might be variable, although position
603 statements and clinical guidelines for specific disorders and targeted populations are clearly
604 outlined in the available literature [77, 122, 126, 127, 176, 189, 217, 225, 226]. However,
605 significant changes in musculoskeletal tissues are generally evident after eight to twelve
606 weeks, although some studies observed increases in muscle mass after only 2 to 4 weeks [37].
607 This early increase in strength is likely caused by neuromuscular and connective tissue
608 adaptations [227], whereas the early increases in muscle CSA may be the result of oedema
609 [228]. For tendon adaptations, longer durations (≥ 12 weeks) appears to be more effective
610 [141]. Example of a potential strength training session is outlined in Table 3 and further
611 examples can be found in our recent published work [229].

612

613 **5.0 Conclusion**

614 This article has briefly examined the mechanisms underpinning positive adaptations to
615 strength training as well as potential benefits for the musculoskeletal system. An overview of
616 training strategies to target these adaptations have also been discussed in both common
617 musculoskeletal disorders and primary prevention strategies. The concepts expressed in this
618 review may help healthcare professionals in understanding and promoting clear and
619 evidence-based recommendations for strength training in musculoskeletal practice, sports
620 medicine and a wide array of medical specialties. Therefore, shared interdisciplinary
621 recommendations appear vital.

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625 **Conflict of interest**

626 Luca Maestroni, Paul Read, Anthony Turner, Konstantinos Papadopoulos declare that they
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628 **Authorship Contributions**

629 LM and AT: concept, layout, and writing the first version of the manuscript. KP, CB and PR:
630 writing, editing manuscript and tables. PC and TS: controlling and editing manuscript, figures
631 and tables.

632

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