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Beyond FITT: How Density Can Improve the Understanding of the Dose Response Relationship Between Physical Activity and Brain Health

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Please cite as: Herold, F., Zou, L., Theobald, P., Manser, P., Falck, R. S., Yu, Q., LiuAmbrose, T., Hillman, C. H., Kramer, A. F., Erickson, K. I., Cheval, B., Chen, Y., Heath, M.,
Zhang, Z., Ishihara, T., Kamijo, K., Ando, S., Gao, Y., Costello, J. T., Hou, M., Hallgren, M.,
Chen, Z., Moreau, D., Farrahi, V., Raichlen, D. A., Stamatakis, E., Wheeler, M. J., Owen, N.,
Ludyga, S., Budde, H., Gronwald, T. (2024). *Beyond FITT: How Density Can Improve the Understanding of the Dose-Response Relationship Between Physical Activity and Brain Health*. SportRxiv.

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All authors have read and approved this version of the manuscript. This article was last modified on 21. May 2024.

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94 Abstract

Research on physical activity and health, including planned and structured forms such as acute 95 and chronic physical exercise, has focused on understanding potential dose-response 96 relationships. Traditionally, the variables of (i) Frequency, (ii) Intensity, (iii) Time, (iv) and Type 97 98 (known as the FITT principle) have been used to operationalize the dose of physical activity. 99 In this article, we describe the limitations of FITT and propose that it should be complemented 100 by the underappreciated variable density, which defines the temporal distribution of physical 101 activity stimuli within a single bout of physical activity or between successive bouts of physical activity relative to time spent resting (e.g., in napping/sleeping or sedentary behaviors). Using 102 the field of physical activity and brain health as an example, we discuss challenges and 103 104 opportunities for further research to use density to improve our understanding of doseresponse relationships between physical activity and health-related outcomes. 105 106

107 Keywords: physical exercise, sedentary behavior, brain, cognition, personalized interventions
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109 **1. Introduction**

110 Physical activity (PA), which includes planned and structured forms such as acute and chronic physical exercise (see Table 1 for definition), is associated with improved brain health across 111 various age groups, and with different health status [1-4]. Regular engagement in PA is 112 beneficial for brain health at multiple levels [5-8], namely (i) the molecular and cellular level 113 (e.g., expression of brain-derived neurotrophic factor [9–15]), (ii) the functional and structural 114 brain level (e.g., brain activity patterns [16-18] or hippocampal volume [19-21]), (iii) the 115 behavioral level (e.g., better cognitive performance [1, 2, 22-30]), and (iv) the risk of adverse 116 117 health-related events (i.e., lower dementia risk [31–34]). However, the optimal dose of PA, including but not limited to the time point at which PA should be applied or repeated to trigger 118 119 changes in specific health-related outcomes (i.e., brain health), is not fully understood [6, 8, 120 22, 26, 27, 35, 36].

There is currently a need for greater clarity in the definition of the dose of PA (including physical exercise) [37–42]. This extends to the call for a more complete reporting of dose in intervention studies using PA [41, 43–45]. From a practical perspective, elucidating the complex doseresponse relationship of PA and health-related outcomes, comprising the interindividual response variability, is an important prerequisite when aiming to maximize the benefits of PA interventions (e.g., on brain health) by individualizing the PA prescription [37, 38, 40, 45–55].

Traditionally, the dose of PA has been characterized and prescribed using the FITT principle, 127 an acronym representing: (i) Frequency, (ii) Intensity, (iii) Time (also referred to as duration), 128 and (iv) Type of PA [51, 56–68]. The FITT principle can also be used to retrospectively analyze 129 how the dose of free-living PA (e.g., unplanned and unstructured forms of PA) is associated 130 131 with health-related outcomes, which can inform recommendations for a specific amount of PA 132 to maintain or improve health. The FITT principle is also commonly used in systematic reviews and meta-analyses when analyzing the dose-response relationship between PA and measures 133 134 of brain health [26–28, 60]. Some researchers have suggested extending the four elements of 135 the FITT principle by the factors of: (v) Volume (V), which is defined as the total amount of PA spent in a given intensity zone that is typically operationalized as a product of the duration of 136 the acute PA bouts spent in a particular zone of intensity x frequency [57]; and, (vi) Progression 137 (P), which characterizes the gradual and systematic increase of the PA stimulus to maintain 138 overload and, thus, provoke further adaptation(s) [69], into FITT-VP [58, 70]. However, 139 adhering to the FITT-VP principle to prescribe and analyze PA has several disadvantages. 140

First, the FITT-VP principle does not take into account all acute and chronic variables (e.g., movement frequency) that determine the dose of PA (especially of planned and structured forms such as acute and chronic physical exercise) [37, 38, 40, 71]. Second, the FITT-VP principle does not consider the temporal distribution of PA stimuli within a single bout of PA or between successive bouts of PA relative to the time spent resting, which is conceptualized as density (see definition below) [37, 38, 40]. Third, each component of the FITT-VP principle is treated somewhat independently when in reality variables characterizing PA can be interrelated [37, 71] (e.g., intensity is significantly influenced by other variables such as acute duration [72, 73] and movement frequency [e.g., cadence operationalized as revolutions per minute when using a cycle ergometer] [74, 75]).

151 For example, one study provided evidence that exercise intensity influences the duration individuals can spend in a specific exercise intensity zone [72]. In particular, in healthy younger 152 153 adults (i) the maximal duration (i.e., defined in minutes) that the participants were able to spend 154 in a given exercise intensity zone during a constant-load exercise test, and (ii) the physiological 155 responses characterizing distinct duration phases during this performance test show a high interindividual variability, while the relative duration (e.g., operationalized as % of maximal 156 duration) was comparable among participants [72]. These findings suggest that a personalized 157 exercise prescription should consider the individualization of the duration spent in specific 158 exercise intensity zones [72, 73]. 159

Regarding movement frequency, a study in trained cyclists showed that, at the same exercise 160 intensity, cycling at a higher movement frequency (i.e., 120 revolutions per minute on a cycle 161 ergometer) led to higher physical demands (i.e., operationalized by ratings of perceived 162 exertion, peripheral blood lactate concentration, heart rate, indices of heart rate variability [74], 163 or spectral parameters of the electroencephalography [76]) than cycling at a lower movement 164 frequency (i.e., 60 revolutions per minute) [74, 76]. In addition to the acute differences in 165 physiological markers, there is evidence that in trained cyclists endurance training at different 166 movement frequencies (i.e., high vs. low cadence training for four weeks) may differently 167 168 influence specific brain measures [77, 78]. In particular, in trained cyclists endurance training 169 at either high or low cadence produces similar improvements in markers of endurance 170 performance (i.e., maximal oxygen uptake and power at the individual anaerobic threshold) [77, 78]. However, training at high cadence led to more pronounced changes in several brain 171 172 parameters (e.g., reduction in alpha-, beta- and overall-power spectral density [77] or increase in frontal alpha/beta ratio [78] assessed during an incremental exercise test). 173

174 The above-presented examples highlight the complexity of determining or providing a specific 175 dose of PA and suggest that an oversimplification of dose may hinder accurate prediction and 176 optimization of PA interventions on health [37, 38, 40]. This is also supported by the fact that 177 different PA variables converge in the PA-induced stimulus (i.e., external load) that feeds into the response matrix, where it interacts with non-modifiable factors such as age, sex, or genetic 178 predisposition, and (potentially) modifiable non-PA-related factors such as sleep, nutrition, 179 general stress, and environmental factors, and then triggers specific biological processes that 180 181 determine the dose (i.e., defined as (a) specific marker(s) of internal load that are involved in biological processes driving the desired changes in outcomes of interest – see Table 1) [37,
38, 40, 71, 79]. Thus accounting for such interrelations of PA variables must not only be
considered when tailoring, programming, or progressing PA interventions [37, 38, 40, 71, 80]
but also as part of the assessment and analytic approaches used.

Consequently, to advance the understanding of the dose-response relationship of PA with specific domains of health (i.e., brain health [40]), it is necessary to consider additional variables, such as density, which we will show can allow for a more precise determination of

- the dose of PA and provide a more nuanced approach beyond the FITT-VP principle.
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- Table 1. Definition of key terms. PA: physical activity; MET: metabolic equivalent of the task;
 SB: sedentary behavior

Key terms	
Brain Health	can be defined as the optimal development and maintenance of brain integrity which encompasses: (i) structural (e.g., hippocampal volume) and functional (e.g., changes in brain activity) brain parameters; (ii) functions that depend on the integrity of the brain, including but not limited to mental health, cognition, and movement; and (iii) the absence of neurological disorders (e.g., dementia). [81, 82]
Dose	is characterized by three key components: (1) external load (i.e., defined as the work performed by the individual independent of internal characteristics), (2) influencing factors (i.e., all factors [e.g., including environmental factors] that can strengthen or weaken the stimuli of a single bout of PA), and (3) internal load (i.e., defined as the individual and acute physiological, psychological, motor, and biomechanical responses to the external load and the influencing factors during and/or after the cessation of a single bout of PA). Thus, the dose can be operationalized and monitored by using specific indicators of internal load involved in the biological processes that drive the desired changes in outcomes of interest. [37, 40, 79]
Physical Activity (PA)	can be defined as any muscle-induced bodily movement (e.g., in occupational or leisure time) that results in an increase in the energy expenditure above ~1.5 metabolic equivalents of the task (MET; 1 MET = 1 kcal (4.184 kJ) \cdot kg ⁻¹ \cdot h ⁻¹). This includes planned and structured forms such as acute and chronic physical exercise (see the following definition). PA can be divided into acute (single bout/session of) and chronic (multiple bout/session) PA based on temporal characteristics." [81, 83–90] Furthermore, PA can be differentiated based on the domains in which it occurs, including recreation/leisure time (such as household), transportation, education, or occupation [87, 88, 91–95].

Physical Exercise	can be defined as a specific form of PA that is planned, structured, repetitive, and designed to improve or at least maintain the performance in one or more fitness dimensions. Physical exercise can be divided into acute (single bout/session) and chronic (multiple bouts/sessions) based on temporal characteristics, also referred to as physical training [83–86, 88, 89, 91]. In addition, physical exercise is typically performed in recreational/leisure time when it is not part of healthcare service (e.g., rehabilitation) or occupation (e.g., elite athlete). To delimit physical exercise from PA: Physical exercise is always PA, PA is not necessarily physical exercise [96].
Sedentary Behavior (SB)	can be defined as any waking behavior characterized by a low energy expenditure (\leq 1.5 MET) while sitting or lying down [87–89, 92, 97, 98]. SB is ubiquitous, due to rapid changes in human environmental, economic, social, and technological contexts. Scientifically, SB has been identified as a newer component of the activity spectrum, which can adversely impact health [99–102]. SB can be categorized as cognitively active (e.g., reading) and cognitively passive (e.g., watching television) [81, 103]. For many adolescents and adults, daily time spent sedentary is \geq 5 hours per day [104–106].

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194 **2. Method**

Given that the German exercise and training variable "Belastungsdichte" [107] (hereafter 195 referred to as "density"), which has its roots in the field of exercise science, is not well-196 197 recognized internationally, we aimed to improve its accessibility by introducing this variable to the broader scientific community. In this context, we extend the description and application of 198 199 "density" to the field of free-living PA, where it has not previously been applied. As "density" is 200 underappreciated in the scientific community, we opted to perform a narrative review, since 201 there is not a large and specific enough literature base to conduct a systematic review (e.g., 202 on the role of density of PA on brain health).

The author group comprises junior, mid-career, and senior researchers from different disciplines, and cultural and ethnic backgrounds.

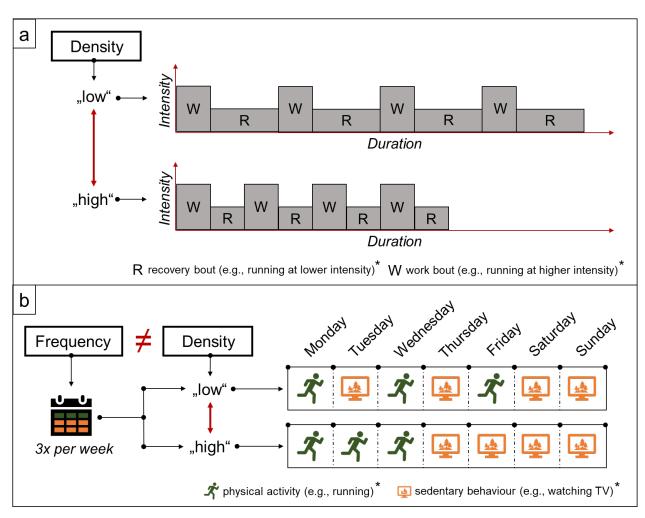
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206 3. Definition of density

Density can be defined as the distribution of PA bout(s) (also referred to as "work bout[s]") or portions thereof over a specific time interval (e.g., within a single bout, day, week, month, or year) in comparison to the time spent resting (also referred to as "rest, recovery or relief bouts") [8, 40, 80, 108]. Assuming the characteristics of work bouts remain similar (i.e., are identical in terms of acute and chronic variables that characterize PA), density is determined by the duration of rest bouts. In other words, density can be modified by changing the duration ofsuch bouts to adjust the work-rest ratio.

In this context, we would like to highlight three important points. First, density is related to the 214 construct of the work-rest ratio, but differs conceptually in that density is associated with 215 216 changing the time spent at rest (i.e., duration of the rest bout[s]), whereas the work-rest ratio can also be adjusted by increasing the duration of the work bout(s). Second, the variables that 217 218 characterize the work bout(s) and the rest bout(s), namely the type of activity, the intensity, and the duration, need to be considered to gain a more nuanced understanding of the influence 219 220 of density and, in turn, the dose-response relationship of PA with measures of brain health. 221 Third, density needs to be further differentiated based on the temporal context, namely (i) in 222 acute density (i.e., in the context of acute PA; see Figure 1 a) and (ii) in chronic density (i.e., in the context of chronic PA; see Figure 1 b) [37]. 223

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Figure 1: (a) Schematic illustration of different acute densities using an acute bout of physical exercise in interval mode as an example. In our example, the number of the work bouts (4x) and rest bouts (4x) is equal whereas the duration of the rest bout in the upper example (i.e., low acute density; the work-rest ratio of 1:2) is twice as long as in the lower one (high acute density; the work-rest ratio of 1:1) resulting in a different acute density and, in turn, dose. In this example, an active rest bout, which is conducted at half of the intensity as the work bout, is selected.

232 The example also illustrates the fact that specific acute variables are interrelated (e.g., acute density, acute duration, and intensity of work and rest intervals). (b) Schematic illustration of 233 the difference between frequency and chronic density in the context of chronic physical 234 activity. The visualization shows that the same frequency (3x physical activity bouts per week) 235 236 can be distributed differently over a week resulting in a different chronic density and, in turn, 237 dose. The asterisk (*) indicates that other acute (i.e., type of physical activity, intensity, and acute duration) and chronic variables (i.e., chronic duration) that characterize the bout(s) of 238 physical activity are assumed to be constant. Please note that we used sedentary behavior 239 240 as an example for the rest bout(s). With regard to acute and chronic physical activity, physical 241 activity at a lower intensity than that of the work bout(s), standing, and sleep can be also encompassed by the rest bout(s), depending on the context. Furthermore, the 242 operationalization of chronic density depends on the period of interest (e.g., day, week, 243 244 month, year).

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3. Operationalization of acute and chronic density

In the following sections, we propose different approaches to operationalize and analyze density considering the temporal context of PA, the availability and accessibility of populationbased datasets, and recent advances in technology to assess PA (i.e., miniaturized wearables to track activities within the 24-hour activity cycle).

251 3.1 Acute density

As illustrated in Figure 1a, acute density can be operationalized by the duration of the rest bout(s) between the successive work bouts (i.e., in seconds or minutes or relative to the duration of the work bout) within a single session of PA. Thus, a modification of acute density can be achieved by decreasing or increasing the duration of the rest bout(s), resulting in a higher acute work-rest ratio (i.e., higher density) or a lower acute work-rest ratio (i.e., lower density), respectively.

258 3.2 Chronic density – Simple analysis approaches

259 The operationalization of chronic density depends on the period of interest (e.g., day, week, 260 month, year). Although chronic density can be operationalized in minutes or hours when several isolated work bouts are performed throughout the day, the operationalization of chronic 261 density is more challenging when longer periods are considered (e.g., week, month, year), 262 especially for unplanned and unstructured forms of PA. To illustrate chronic density in terms 263 of a micro-cycle of one week, consider the following example: if a person is physically active 264 on Monday, Wednesday, and Friday or Monday, Tuesday, and Wednesday, this will result in 265 the same frequency but not the same chronic density within a micro-cycle of one week (see 266 also Figure 1b). More specifically, in the first example shown in Figure 1b, the person is 267 268 physically active on non-consecutive days (i.e., work bouts spread over a week), whereas in the second example, the person is physically active on consecutive days (i.e., work bouts 269 270 performed on three consecutive days).

271 Accordingly, a simple approach to studying the influence of different chronic density patterns on brain health is to characterize different groups of individuals based on their chronic density 272 273 patterns (e.g., a low chronic density group in which individuals performed PA on non-274 consecutive days versus a high chronic density group in which individuals performed PA on 275 consecutive days – see also Figure 1). For chronic physical exercise, the influence of chronic 276 density on specific measures of brain health can be studied by comparing intervention groups 277 that were instructed to perform physical exercise sessions with different chronic densities (e.g., a low chronic density group performing physical exercise sessions on non-consecutive days 278 279 versus a high chronic density group performing physical exercise sessions on consecutive 280 days).

281 3.3 Chronic density – Sophisticated analysis approaches

282 Comparable to other studies analyzing the influence of PA patterns (e.g., intensity, and 283 duration of the acute PA bouts) on health-related outcomes (e.g., cognitive performance or 284 cardiometabolic health), the application of more sophisticated approaches using distributional 285 data analysis [109] or machine learning (e.g., via K-means clustering) [110–113] holds some promise for identifying groups of individuals with distinct chronic density patterns. Despite 286 287 some limitations and challenges (e.g. the need for large sample sizes and, high-dimensional data, the time-consuming nature of training algorithms, and the lack of benchmark data), 288 289 machine learning-based approaches provide several advantages for the purpose of profiling 290 PA patterns (e.g., more accurate classification and prediction, the possibility of a hypothesis-291 free/generating approach) [114–118]. Another advantage of machine learning-based approaches is their capacity to handle large, complex, and high-dimensional datasets [114]. 292 293 The ability and flexibility to handle such datasets make machine learning-based approaches 294 well-suited for analyzing the influence of density on specific markers of brain health because 295 density is a more complex variable than other PA variables (e.g. duration). This assumption is 296 supported by the fact that these approaches have already been successfully applied to 297 elucidate the influence of "micropatterns" of PA including intensity and duration (also referred 298 to as bout length) on health-related outcomes such as mortality [119, 120] and cancer incidence [121]. Thus, extending machine learning-based approaches to density is a promising 299 area for future research to elucidate the influence of different chronic density patterns on 300 301 measures of health in general and brain health in particular.

In the context of brain health, the application of such sophisticated classification and analysis techniques may enable the investigation of specific research questions (e.g., *is a low density* of moderate-intensity PA in older adults more, less, or equally beneficial for brain health than having a high density of moderate-intensity PA?) or to study the association of specific densityrelated PA patterns, such as the stability of density, with measures of brain health. In this context, we propose that the stability of density is characterized by the periodicity and the fluctuations (variability) that are reflected by the degree of randomness of the duration of the rest bouts between successive work bouts within a given time interval (e.g., day, week, month, year). We suggest that, among other approaches [122], the stability of density can be operationalized by measures used to assess fractal dynamics.

312 Fractal dynamics are characterized by the self-affinity (also referred to as self-similarity or scale 313 invariance) of a given signal (e.g., derived from accelerometers) across time scales [123–127]. 314 There is a strong case to be made that fractal dynamics can help to better understand the periodization of chronic physical exercise [128], and several studies have used this approach 315 to analyze physiological data (e.g., frequently applied to heart rate variability data [129–146]) 316 317 or PA patterns [147–150]. In the context of PA, a popular method for assessing fractal 318 dynamics (e.g. of PA [147-150]) is detrended fluctuation analysis (DFA), which is a nonstationary time-series analysis of specific signals (e.g., accelerometer data) that reflects 319 the correlative structure and fractal dimension of signal fluctuations across a range of time 320 321 scales based on a modified root-mean-square analysis [126, 127, 151–153]. For instance, a study using data from 5097 middle-aged adults showed that greater fractal stability of daily PA 322 (i.e., assessed via a thigh-mounted accelerometer over seven days and reflected in a higher 323 DFA scaling exponent) was associated with better verbal fluency performance in males but not 324 in females [150]. Such sex-specific differences are consistent with the evidence suggesting 325 that sex is an important moderate in the relationship between PA and brain health [47, 48, 326 154–160]. However, whether such findings extend to the chronic density of PA remains a 327 328 promising area for further investigations.

329 3.4 Recommendations regarding the assessment of chronic density

To quantify the chronic density of PA, we recommend the application of device-based 330 331 assessments to complement subjective assessments (i.e., questionnaires) for the following 332 reasons. First, popular questionnaires to assess chronic PA such as the International Physical 333 Activity Questionnaire (IPAQ) only quantify the frequency but not the chronic PA density (i.e., 334 neither the long form [161] nor the short form [162] of the IPAQ), although some recently 335 developed questionnaires do collect such information (e.g., Daily Activity Behaviours Questionnaire [163-166]). Second, although subjective assessment tools (e.g., 336 questionnaires) have several advantages (e.g., low burden for participants, cost-effective and 337 convenient administration), they are prone to several sources of bias (e.g., recall bias or social 338 desirability bias) that can confound the estimation of chronic PA patterns [95, 167-169]. 339 Device-based assessment tools can circumvent the above-described limitations of subjective 340 341 assessment tools, but it should be considered that (i) the applied device-based measurement tool needs to be valid and reliable [170-172], and (ii) there is not yet a fully established 342 343 consensus on the application of device-based measurement tools (e.g., placement and sampling frequency of the device) or on the data processing procedures to obtain specific 344

indices of PA (e.g., minimal length of the epochs, filter, cut-off points, non-wear-time definition)
although some recommendations exist [173–175].

347 Furthermore, we recommend combining popular device-based tools such as accelerometers 348 with other sensors (e.g., for environmental light, barometer/altimeter, or geolocation) and 349 digital tools (e.g., smartphones) to allow for the recording of contextual information (e.g., weather via geolocation at specific time point [176] or type of activity conducted during rest 350 351 bout(s) via an accelerometer-triggered e-diary [176–182]). The latter approach is also referred to as ambulatory assessment [81, 177, 183, 184]. In addition, regarding the analysis of chronic 352 353 density in the context of chronic PA, future studies should consider SB and sleep to provide a 354 more holistic understanding of the 24-hour activity cycle on health in general [185–189] and 355 brain health in particular [81, 92, 190–193].

3.5 The potential of density to complement existing analysis approaches of the 24-hour activity cycle

Since density specifies the temporal distance between stimuli within or between successive bouts of PA, it can complement other approaches used to analyze the influence of PA patterns within the 24-hour activity cycle on health-related outcomes, namely (i) timing of PA (e.g., time of day on which the PA has been conducted such as in the morning, afternoon or evening [194–196]) and (ii) compositional data analysis (e.g., using the relative time spent in a specific activity [e.g., PA] in relation to the time spent in other activities [e.g., SB or sleep] instead of absolute times spent in a specific activity for analysis [197–204]).

In terms of the diurnal impact of PA, PA is an important "Zeitgeber" (time cue) for the human 365 366 circadian system [205] and thus a critical factor in sleep health, a mediator of the effects of PA 367 on brain health [5, 206]. In this regard, the findings of a recent systematic review suggest that there is currently no consistent evidence in adults as to whether PA conducted at one time of 368 day (e.g., morning) is associated with more pronounced health benefits than PA performed at 369 a different time of day (e.g., afternoon or evening) [194]. In general, PA is associated with 370 371 better sleep health [207–212], but there is no compelling evidence that PA performed at any particular time of day is superior for promoting sleep health [209, 213, 214] because even 372 373 acute PA conducted in the evening is not typically detrimental for sleep [215-217] if it is not 374 performed too close before bedtime (≤ 1 hour) [215]. To the best of our knowledge, the timing 375 of PA and its direct relationship with measures of brain health so far has received relatively 376 little attention in empirical studies. The findings from one study suggest that, in adolescents, 377 an acute bout of physical exercise in the morning is more effective in improving behavioral 378 measures of brain health (e.g., global reaction time), compared with the afternoon [218]. However, currently (i) there is a lack of studies on the influence of the timing of PA on brain 379 380 health, and (ii) the evidence on the timing of PA on sleep health, an important mediator of the

effects of PA on measures of brain health [5, 206], is less clear. Thus, future research is needed to draw firm conclusions on whether the timing of PA can influence specific measures of brain health differentially [219]. Such future research on the timing of PA is likely to benefit from considering density, which specifies the temporal distance between stimuli within or between successive bouts of PA (e.g., the time between morning and/or evening bouts of PA).

Compositional data analysis has been used to investigate the relationship between PA and 386 387 behavioral measures of brain health in preschoolers [220–222], middle-aged [223], and older adults [224] and has provided valuable insights into the complex relationship between PA and 388 389 brain health. For example, compared to other activities of the 24-hour activity cycle (e.g., SB 390 and sleep), a loss of time spent in moderate-to-vigorous PA appears to be relatively detrimental 391 to cognitive performance (i.e., cognition composite score) in middle-aged adults, given its smaller relative amount in the 24-hour cycle [223]. Notably, in older adults, longer time spent 392 in light-intensity PA was associated with better inhibitory control (i.e., operationalized by Stroop 393 task performance), especially when accumulated in bouts longer than 10 minutes [224]. 394

395 Comparable to compositional data analysis approaches, a promising area for further 396 investigations is to operationalize density as the relative time spent in work bout(s) (e.g., PA in 397 a specific intensity zone) in relation to the time spent in rest bout(s) (e.g., SB or sleep) to further 398 our understanding of the temporal dynamics of PA and their influence on brain health. Such a 399 better understanding of the temporal dynamics of PA is needed to better inform the 400 individualization of PA interventions [225].

401 *3.6 Interim summary*

402 Taken together, chronic density captures information beyond that provided by frequency, because frequency only specifies the number of PA bouts in a given time interval (e.g., day, 403 week, month, year) but not their distribution within that time interval. Given that the dose of PA, 404 which is influenced by the external load and confounding factors in terms of the acute 405 406 psychophysiological responses elicited [37, 40], is an important factor in inducing changes in 407 measures of brain health, including cognition [22, 27], it seems reasonable to assume that acute and chronic PA performed at different densities might differentially influence measures 408 of brain health. This latter assumption is also supported by the fact that density is also related 409 410 to exercise intensity [80, 226, 227] and both acute and chronic density are variables that are important in inducing a specific level of overload and achieving progression [70], both of which 411 412 are well-known and important factors and principles influencing the dose of PA and therefore 413 the desired outcomes [40, 69]. In the next section, we will discuss the role of density in modifying the dose of PA in more detail. 414

416 **4. Density and the dose of physical activity**

Currently, neither the precise dose [6, 8, 22, 26, 27, 35, 36] nor the neurobiological 417 mechanisms that drive the positive effects of acute and chronic PA on brain health are fully 418 understood [5, 6, 23, 40, 228–230]. This knowledge gap extends to the empirical evidence on 419 420 how density may influence the dose and neurobiological mechanisms that drive brain health. 421 However, our assumption that accounting for density is crucial when aiming to elucidate the dose-response relationship between PA and brain health is supported by evidence from (i) 422 acute PA studies on the temporal dynamics of specific markers of brain health and (ii) studies 423 424 on glycemic control and brain health in adults with type 2 diabetes, although the latter cannot be readily generalized to healthy adults. 425

426 *4.1 Temporal dynamics of acute physical activity for brain health*

427 There is some evidence from a meta-analysis that the after-effects of acute physical exercise on cognitive performance are transient, depending on the characteristics of the physical 428 exercises, such as type of physical exercise, intensity, and duration [25]. More specifically, 429 430 according to this meta-analysis, the greatest effects of acute physical exercises on cognitive performance can be expected 11-20 minutes after the cessation of the acute bout of physical 431 432 exercises and diminish with longer delays [25]. However, some studies provide evidence that 433 the after-effects of acute physical exercises on specific behavioral measures of brain health (e.g., executive functions) can even persist for up to 30 minutes in healthy younger adults 434 [231–234], 60 minutes in children [235] and younger adults [236], and 90 minutes in healthy 435 younger adults, [237] or even that in healthy younger adults performing acute physical exercise 436 four hours after learning is more beneficial for improving memory performance and 437 hippocampal pattern similarity (i.e., assessed 48 hours later) as compared to performing acute 438 physical exercise immediately after learning the task [238]. 439

440 Based on the paucity of research in this area, the exact time course and moderators (e.g., 441 acute PA-related factors such as type, intensity, duration, and non-PA-related factors such as 442 age, sex, health status, and fitness level) of the after-effects of acute PA on specific measures of brain health remain somewhat elusive, at least in part due to methodological challenges 443 (e.g., a limited number of follow-up assessments, confounding influence of activities performed 444 445 between cessation of acute PA and cognitive test administration) [23]. However, based on the above-presented evidence, it is reasonable to assume that considering temporal dynamics of 446 PA - conceptualized as density - has a great potential to add to our understanding of the dose-447 448 response relationship of acute PA on specific measures of brain health. More importantly, considering density in future research may help to elucidate the precise time point(s) at which 449 450 the acute PA stimulus needs to be applied or repeated to prolong the acute PA-related benefits 451 on specific measures of brain health. Such information on the appropriate timing to set a PA

stimulus is thus crucial to inform an experimental design and to maximize the effectiveness of
PA interventions (e.g., "just-in-time adaptive PA interventions" [239–241]).

454 Several studies support the notion that the density of the PA can be important in optimizing the 455 effectiveness of PA interventions. For example, two studies in healthy younger adults 456 investigated the effects of two repeated acute bouts of high-intensity interval exercise (HIIE, 457 4x 4-minute work bouts at 90% of $VO_{2 peak}$ interspersed with 3-minute rest bouts at 60% VO_{2} peak) on inhibitory control (i.e., assessed by the Stroop task every 10 minutes after the cessation 458 of each bout of physical exercise for 5x times) [242, 243]. In both studies, a recovery interval 459 of 60 minutes separated the first bout of acute HIIE from the second bout of HIIE, in which the 460 461 Stroop task performance was repeatedly assessed [242, 243]. These studies showed that inhibitory control (i.e., reverse Stroop interference score) improved immediately [242, 243] and 462 10 minutes [243] after exercise cessation after the first and second bout acute bouts of HIIE 463 compared to the pretest. However, only after the first acute bout of HIIE the after-effect did 464 persist up to 40 minutes after exercise cessation [242, 243]. In contrast, the executive 465 performance assessed 10 minutes [242] or 20 minutes [243] after the second bout of HIIE was 466 467 not significantly different from the pretest and was lower than that of the first bout of HIIE when 468 assessments at 20 minutes [242], 30 minutes [242], and 40 minutes [242, 243] (but not 50 minutes [242, 243]) after exercise cessation were considered. Collectively, these observations 469 470 suggest that the acute PA-related effects on inhibitory control were less pronounced in the 471 second bout of HIIE compared to the first bout of HIIE. Hypothetically, such a diminished effect 472 after the second bout of HIIE could be, among other factors, related to the relatively close 473 temporal proximity between the two single bouts of HIIE (i.e., 60 minutes).

474 Based on the observation that the acute PA-induced performance improvements in inhibitory control correlated with changes in blood lactate concentration in both studies [242, 243] and 475 that changes in peripheral blood lactate concentration were significantly lower during and after 476 477 the second bout of HIIE [243], it seems reasonable to speculate that there is a neurobehavioral relationship between both measures [8, 40, 108, 244, 245]. This assumption is supported by 478 the fact that peripheral blood lactate can cross the blood-brain barrier via monocarboxylate 479 480 transporters and be utilized as "fuel" for cognitive processes [246-254], which may further 481 explain the positive associations between acute PA-induced blood lactate increases and cognitive enhancement. Indeed, recent studies have reported that changes in peripheral blood 482 lactate concentration are correlated with acute PA-related improvements in cognitive 483 performance [255–257] although it remains somewhat unclear whether blood lactate changes 484 are a mediator of acute PA-induced benefits on cognitive performance because only one study 485 found evidence in favor of this idea [258] while another did not [259]. 486

In addition, there is evidence that a change in peripheral blood lactate concentration (e.g.,
induced by acute physical exercise [260] or infusion at rest [261]) is associated with a change

in the concentration of serum levels of the brain-derived neurotrophic factor (BDNF), an 489 important neurotrophin involved in processes of PA-related neuroplasticity and brain health [7, 490 12, 15, 262–267]. Notably, in younger healthy adults BDNF changes in response to acute PA 491 492 are correlated with cognitive improvements [268], lending credence to the hypothesis that 493 BDNF is involved in acute PA-induced improvements in behavioral measures of brain health [269]. Such acute PA-triggered effects of BDNF on cognitive performance are likely to be 494 495 transient, as several studies on the kinetics of BDNF have consistently shown that elevated BDNF levels return to baseline 15-60 minutes after exercise cessation (for review, see [9]), 496 497 supporting the notion that temporal dynamics (e.g., density) should be considered when 498 examining the effects of acute PA on brain health.

Regarding the functional brain level, alterations in cerebral blood flow (CBF) are hypothesized 499 500 to mediate the acute effects of PA on behavioral measures of brain health [23]. Indeed, some 501 studies provide evidence that acute PA-induced changes in cerebral blood velocity (CBV), a 502 surrogate for CBF that can be operationalized by monitoring middle cerebral artery velocity via 503 transcranial Doppler ultrasound [270–273], correlate with acute PA-induced improvements in 504 behavioral measures of brain health (i.e., executive functioning operationalized by the 505 antisaccade task) [274, 275]. The acute PA-induced increase in CBV can persist for up to 2 506 hours after exercise cessation depending on several factors (e.g., characteristics of the person and the acute bout of PA, methodological factors - for review see [270]) but typically returns to 507 508 baseline levels relatively shortly after exercise cessation [270, 271] (e.g., 30 minutes - for review see [270]). Comparable to the transient effects of acute PA at the cellular and molecular 509 level (e.g., BDNF), the transient nature of acute PA-related changes at the functional brain 510 511 level (e.g., CBF) urges future research to consider density as a variable to facilitate our 512 understanding of the neurobiological mechanisms mediating the effects of acute PA on brain health, which is currently relatively scant [5, 23, 229]. Such a better understanding of the 513 temporal dynamics at different levels of analysis [5, 23, 40] (e.g., molecular and cellular levels, 514 such as changes in the noradrenergic and dopaminergic systems [230] or functional levels, 515 such as brain activity or connectivity changes [17, 18]) may yield a more robust understanding 516 517 of the potential dose-response relationship, which in turn can help to inform future practical 518 applications better.

A recent study provided direct evidence that acute density can influence the acute PA-related effects on specific behavioral measures of brain health. In particular, this study used a withinsubject crossover design with a pretest-posttest comparison to investigate in healthy younger adults whether the use of different inter-set rest intervals (i.e., 1 minute versus 3 minutes, representing higher and lower acute densities) during an acute bout of low-load resistance exercise (i.e., 40% of a one-repetition maximum, 6x sets of 10x repetitions) can influence acute exercise-induced changes in inhibitory control (i.e., operationalized with the Stroop test) [276].

In this study, it was observed that shorter inter-set rest intervals (i.e., 1 minute - high density) 526 improved inhibitory control (i.e., operationalized by a reverse Stroop interference score) 527 immediately, 10 minutes, 20 minutes, and 30 minutes after exercise cessation, whereas such 528 effects were absent for longer inter-set rest intervals (i.e., 3 minutes - lower acute density). 529 530 Moreover, the improvement in executive functions was greater at 20 and 30 minutes after exercise cessation in the shorter inter-set rest interval condition (i.e., higher acute density) 531 532 compared with the longer inter-set rest interval condition (i.e., lower acute density) [276]. Thus, the findings of the above-presented study provide strong support for the importance of 533 534 considering acute density when investigating the dose-response relationship of acute PA with 535 specific measures of brain health.

536 4.2 Glycemic control and brain health

537 There is growing evidence that type 2 diabetes, which is characterized by impaired glucose 538 control [277] and poses a public health burden due to its high and still growing worldwide prevalence and related health complications [277–280], is associated with significantly poorer 539 540 brain health [281–284]. For instance, there is accumulating evidence that type 2 diabetes is associated with reduced structural and functional brain integrity [285-288], lower cognitive 541 542 performance [285-293], and an increased risk of dementia [294-297]. Given that impaired homeostasis of glucose control is the key feature of type 2 diabetes [277], maintaining "normal" 543 glucose control across the lifespan (e.g., by reducing sedentary behavior and engaging in PA) 544 545 seems to be an important factor in maintaining brain health, especially in later life stages [298]. 546 Indeed, some systematic reviews provide evidence that PA in adults with type 2 diabetes is associated with a positive but weak influence on specific measures of brain health such as 547 cognitive performance, [299–302] although such evidence is not universal, probably due to the 548 heterogeneity of intervention studies in terms of the exercise and training variables 549 550 characterizing the physical exercise interventions [303].

Notably, two small-scaled studies (n = 12 in both studies) in adults with type 2 diabetes showed 551 552 that interrupting 7 hours of sitting with 3 minutes of light-intensity walking every 15 minutes (i.e., high acute density) was more beneficial for specific measures of glucose control (e.g., 553 fasting glucose and duration of the dawn phenomenon [304] or post-breakfast and 21-hour 554 glucose control [305]) than interrupting sitting every 30 or 60 minutes (i.e., low acute density) 555 556 [304, 305]. During the rest periods, the participants had access to a personal computer, 557 internet, and books [304, 305]. Thus, these two small studies in adults with type 2 diabetes 558 provide preliminary evidence that density can influence neurobiological processes (i.e., 559 glucose control) relevant to brain health [298] which, in turn, supports our idea that considering density is crucial for a more nuanced understanding of the dose-response relationship between 560 PA and measures of brain health. However, the higher density in the above-described studies 561

562 [304, 305] is also related to a higher frequency of physical exercise bouts, and thus future high-563 quality studies are needed to (i) disentangle the unique influence of frequency and density on 564 (brain) health-related measures, and (ii) investigate whether different acute and chronic 565 densities of PA might differentially influence specific levels of brain health (e.g., at the 566 molecular and cellular levels such as the release of brain-derived neurotrophic factor).

567 *4.3 Interim summary*

Taken together, the evidence on temporal dynamics of specific markers of brain health in 568 response to acute PA and the glucose control - brain health association corroborates our 569 570 assumption that density is important for advancing our understanding of the dose-response relationship between PA and measures of brain health because it provides crucial information 571 on temporal distribution of PA. More specifically, studying density plays an important role in 572 573 understanding the minimal and optimal dose by providing information on the minimal and 574 optimal time interval (i.e., rest bout) between PA stimuli within a single bout of PA or successive 575 bouts of PA (i.e., work bouts) being required to maintain or improve specific measures of brain health. Such information on the minimal and optimal time intervals for the delivery of a PA 576 577 stimulus holds great potential to inform and optimize intervention approaches aimed at 578 promoting PA, such as "just-in-time adaptive PA interventions" [239-241] (e.g., in the context 579 of breaking up prolonged sitting with acute breaks of PA including physical exercise [306-580 309]).

581

582 **5. Density in relation to other activities of the 24-hour cycle**

583 There is an increasing interest in the scientific community to develop a more holistic 584 understanding of the influence of the 24-hour activity cycle including PA, standing, sedentary 585 behavior (SB), and sleep on health status [185–189] and brain health [81, 92, 190–193].

586 Regarding density, rest bouts are a key construct and may be considered synonymous with, 587 or primary to, time spent in SB when considering waking hours. Epidemiological and 588 experimental evidence shows that sedentary time may influence the relationship between 589 participation in PA and its well-established cardiometabolic health benefits (i.e. highly sedentary individuals may need to do more than the recommended levels of PA to offset the 590 detrimental effects of sedentary behavior) [99-101, 310]. Experimental evidence provides 591 592 compelling insights into the potential for "exercise resistance" [100]. Coyle and colleagues showed that when acute physical exercise was preceded by a prolonged period of SB, 593 postprandial metabolic responses and metabolic benefits were significantly attenuated [311-594 595 313]. More specifically for brain health, the effect of physical exercise on cognitive function is 596 altered by subsequent exposure to prolonged sitting versus breaks in sitting [306], and

emerging evidence shows that different types of SB, namely passive and mentally active SB, 597 could be differentially associated with brain health [81, 103, 314]. For instance, previous 598 599 studies have indicated that mentally active SB (e.g., reading or using a computer) can benefit measures of brain health (for review see [81, 103, 314]). A growing body of evidence suggests 600 601 that the consequences of too much time spent in SB are distinct from those of too little PA with respect to cardiometabolic health [100] and brain health [81, 101, 105]. This reinforces the 602 603 utility of considering SB as a mechanism for the importance of density as a key new element 604 to complement the FITT-VP principle.

605 Given that the duration and the characteristics of the rest bout(s) are the key elements in 606 defining density, considering sleep is important in understanding how the temporal distance 607 between successive bouts of PA can influence measures of brain health, especially when tracking and analyzing free-living PA over longer periods (e.g., a week, month, or year). There 608 609 is growing evidence that sleep (i.e., often operationalized as time in bed) can mediate and/ or moderate the effect of PA on brain health ^[193, 219, 315, 316]. For example, several cross-sectional 610 studies provide evidence that (i) older adults with poor sleep efficiency (i.e., percent of the time 611 612 in bed spent asleep) benefit most from PA in terms of global cognition [317], (ii) sleep efficiency 613 mediates the relationship between PA and working memory, task switching, verbal ability and 614 fluency, and memory recall in a mixed sample of younger and older adults [21], (iii) better 615 subjective sleep quality mediates the relationship between PA and verbal fluency, immediate recall, and delayed recall [318] or working memory [319] in middle-aged and older adults, and 616 (iv) subjective sleep quality and sleep efficiency mediate the relationship between PA level and 617 618 inhibitory control in younger adults [320]. A 6-month intervention study, in which cognitively 619 healthy older adults performed moderate- or high-intensity interval exercise twice a week, 620 reported that participants in the moderate-intensity group, who had poorer sleep efficiency at 621 baseline, showed greater exercise-induced improvements in episodic memory and global cognition [321]. 622

623 Collectively, the above-presented evidence supports the idea that consideration of all activities 624 in the 24-hour activity cycle [81, 92, 190–193, 316] is necessary to improve our understanding of the influence of specific lifestyle-related factors on brain health. This assumption is 625 reinforced by emerging evidence suggesting that (i) other activities of the 24-hour cycle that 626 can contribute to or constitute the rest bout(s), such as free-living standing activity [322] and 627 light-intensity PA [29], are positively associated with behavioral measures of brain health, and 628 (ii) activities such as SB and sleep, which are typical activities of a rest bout(s), interact with 629 630 each other with respect to brain health, as an observational study showed that sleep problems 631 mediated the detrimental associations of passive SB with depression [323]. To this end, complementing the 24-hour activity cycle approach with density may enable even more 632

nuanced insights into its health effects by improving the characterization and thus ourunderstanding of the dose of PA.

635

636 6. The current state of evidence and future directions

637 The role of density as an important variable can be considered helpful when investigating doseresponse relationships of PA with key health-related outcomes (e.g., brain health). For brain 638 health, the current evidence indicates that (i) acute density is typically not considered when 639 analyzing the influence of acute bouts of PA on cognitive performance (e.g., as a moderator 640 variable) [23, 25, 324–326], (ii) chronic density is often not reported in studies investigating the 641 642 influence of chronic PA on brain health [8, 327], (iii) chronic density is absent in moderator analyses in recent systematic reviews and meta-analyses investigating the influence of chronic 643 PA on cognitive performance [22, 26], and (iv) chronic density is typically not mentioned in 644 645 recommendations (e.g. from the World Health Organization) and policies aimed at reducing the risk of cognitive decline and dementia by lifestyle changes (e.g., via PA) [328]. Such an 646 absence of density in the literature, analyses of the dose-response-relationships, and 647 recommendations of official bodies could lead to the assumption that (i) acute and chronic 648 649 density are unimportant variables or (ii) that researchers studying the effects of PA on 650 measures of brain health are unaware of the importance of density.

651 Given that other fields of research have begun to recognize the influence of the distribution of 652 PA across a week (e.g., the "weekend warrior" pattern characterized by ≤2x bouts [329–336] or 1x bout [337] of PA per week) and the interrelated impacts of PA, sleep, and SB [81, 92, 653 100, 101, 186–189, 191–193, 316], density is an excellent candidate determinant of brain 654 655 health effects that should not be overlooked when analyzing the dose-response relationship 656 within the context of PA-related benefits on measures of brain health. To simulate future research, we highlight in the following two sections further directions for observational and 657 intervention studies on the influence of PA density on measures of brain health. 658

659 6.1 Observational studies

Other research fields have started to analyze observational and population-based data in 660 661 adults regarding the influence of achieving the amount of PA recommended by the World 662 Health Organization (i.e., ≥150 minutes of moderate- or ≥75 minutes of vigorous-intensity PA per week [88, 89]) in ≤2x bouts per week (i.e., denoted as "weekend warrior") or ≥3x bouts per 663 week on health-related outcomes such as the risk of mortality [329–331], risk of cardiovascular 664 665 events [336], prevalence and health aspects associated with the metabolic syndrome (e.g., adiposity, hypertension) [332, 334], or risk of mental disorders [333]. Although none of the 666 above-mentioned studies considered chronic density, because they did not account for the 667

temporal distance between the successive bouts of PA into account, all provided evidence that achieving the recommended amount of PA in $\leq 2x$ bouts per week has a comparable influence on health-related outcomes as achieving this amount in $\geq 3x$ bouts per week [329–334, 336].

671 Whether such observation extends to measures of brain health, given the moderating role of 672 the acute and chronic density of PA is considered, is a promising area for further investigations. 673 In this regard, we would like to acknowledge that all activities of the 24-hour activity cycle (i.e., 674 PA, sedentary behavior, and sleep) should be considered for a more nuanced understanding of the dose-response relationship between PA and health in general [185, 186] and brain 675 health in particular [81, 92, 191, 192]. In the context of acute and chronic density, we reiterate 676 677 that the characteristics that define the work bout(s) and rest bout(s) must be considered when analyzing density (i.e., type of activity, intensity, and duration). This assumption is supported 678 by emerging evidence showing that the characteristics of activities that are primarily involved 679 in the rest bout(s) can influence brain health differentially. More specifically, there is evidence 680 that the type of SB can moderate the effects of SB on brain health because cognitively active 681 682 SB (e.g., reading) is positively associated with brain health, whereas cognitively passive SB (e.g., watching TV) did not confer such benefits [81, 103, 314, 338]. 683

In addition, from a public health perspective, a key distinction is made between active and 684 passive (sedentary) occupations [339]. In this context, analyzing the influence of acute and 685 chronic density on measures of brain health might be especially relevant for health-related 686 research in individuals with professions that require performing substantial occupational PA at 687 higher intensities in relatively short time intervals (e.g., construction workers, or farmers) 688 689 versus desk-based workers. Considering density in addition to traditional exercise variables (e.g., FITT-VP principle) may enhance our understanding of the "physical activity paradox" 690 (i.e., occupational PA has less clear or no health benefits compared to leisure-time PA) [340-691 344] and the identification of "sweet spots" (e.g., individualizing leisure time PA 692 recommendations by considering occupational PA levels) [187] which in turn can help to better 693 694 inform future public health interventions. The latter assumption is reinforced by the fact that 695 individuals with a lower socioeconomic position (i.e., lower educational qualifications, 696 occupational class, income, or living in a deprived area), as compared to those with a higher 697 socioeconomic position, showed different characteristics concerning their 24-hour activity cycle since they spent more time standing, moving, and walking but less time sitting during 698 699 weekdays while on weekends these patterns were reversed [345]. Notably, those with higher 700 socioeconomic positions engaged in higher levels of physical exercise-like activities (i.e., 701 running, cycling, and inclined walking) and less time lying regardless of the day of the week. 702 These findings suggest that socioeconomic disadvantages are mirrored in 24-hour activity 703 cycle patterns [345]. Such an observation is of particular relevance for future studies on PA 704 and brain health given that in adults a lower socioeconomic position is negatively associated

with different markers of brain health (e.g., lower cognitive function and higher cognitive decline
[346–356], higher dementia risk [355, 357–360], less favorable brain structure outcomes [353,
354, 360, 361]). Future well-designed research is needed for more robust conclusions in this
direction [362, 363] and may benefit from considering the 24-hour activity cycle [191] including
the density of PA.

710 6.2 Intervention studies

In addition to the examination of density in observational studies, we also recommend that 711 acute and chronic density should be considered in the prescription of PA intervention studies 712 to improve the standardization of reporting, the determination of the dose, and the 713 714 comparability across studies. Although there is evidence that a higher frequency (i.e., 5-7 PA 715 sessions per week), which is probably also mirrored in a higher chronic density, is more 716 beneficial for improving cognitive performance in adults older than 50 years (i.e., double the 717 effect size; 0.69 vs 0.32) than a lower frequency (i.e., 1-2 PA sessions per week) [27], providing 718 information on acute and chronic density can be especially relevant for interventions with lower 719 levels of direct supervision (e.g., home- and technology-based interventions using 720 exergames). For example, in home-based studies using exergames and providing only general 721 supervision, partial direct supervision, or even no supervision (for more information on supervision please see [364, 365]), older adults are typically instructed to achieve a certain 722 723 duration of physical exercise over a week but are often allowed to self-select the frequency of 724 the acute PA bouts [366–373]. Such studies have documented that older participants who are 725 highly motivated can exceed the recommended training frequency and/or perform multiple acute PA bouts throughout the day [368, 373–375]. This may result in insufficient rest time, 726 727 which is perhaps less than optimal for the materialization of adaptation processes (i.e., 728 consolidation). The above theoretical assumption is supported by (i) an experimental study 729 showing that in younger adults too much consecutive computer-based training can be 730 detrimental to learning performance (i.e., accuracy of motion discrimination) [376] and (ii) a 731 systematic review observing that cognitive performance declines when endurance athletes are 732 overreached or overtrained [377]. These latter findings support the assumption that acute and chronic density should be considered when prescribing and monitoring physical exercise 733 734 interventions aimed at promoting brain health.

In particular, acute and chronic density are important variables in the organization of physical exercise, namely the periodization and programming of physical exercise sessions, because they characterize the dose by defining the duration of rest bout(s) within a single bout of physical exercise or between successive bouts of physical exercise (i.e., work bouts). Whereas periodization is the temporal organization (i.e., macro-management) of the characteristics of physical exercise sessions (e.g., purposeful adjustment of variables such as exercise intensity and volume for progression) and application of training principles [37, 40, 128, 378–380],

programming is defined as the micro-management of physical exercise that includes, but is 742 not limited to, the organization of exercise and training variables (e.g., type of physical 743 744 exercise, exercise intensity, exercise duration, and acute and chronic density) [40, 378, 380]. 745 Thus, acute density is especially relevant for programming acute physical exercise sessions 746 in which the physical exercises are performed in interval mode or a set structure because acute density defines the rest duration between the work bouts (e.g., also referred to as intervals or 747 748 repetitions), between interval series or sets, or between different physical exercises [80, 227, 379]. As shown in Figure 1, acute density can be manipulated to alter the acute PA stimulus 749 750 by decreasing or increasing the duration of rest between successive work bouts.

From the perspective of PA promotion, density can also complement newer approaches to 751 foster PA, such as "vigorous intermittent lifestyle physical activity" (VILPA) [381, 382] and 752 "exercise snacks" [381–384]. While VILPA has been empirically defined as vigorous bouts of 753 754 incidental PA lasting up to 1 or 2 minutes [119, 121], the term "exercise snacks" has been more 755 loosely defined as single planned bouts of physical exercise that typically (i) lasts \leq 1 minute, 756 (ii) occur multiple times throughout the day, and (iii) are performed at a vigorous intensity [382-757 384]. Regarding the VILPA and "exercise snacks" concepts, the variable density as a 758 characteristic defining the dose can help to more precisely elucidate the influence of different 759 rest durations between the short work bouts (e.g., performed at the vigorous intensity and conceptualized in the VILPA and "exercise snacks" approach or at other intensities in the 760 761 context of free-living PA such as light- or moderate-intensity PA) on health-related parameters (e.g., brain health). However, it is worth noting that for a purposeful modification of density, the 762 763 interrelation with other exercise variables needs to be considered (e.g., implementation of passive or active rest periods, exercise intensity, and duration of work and rest bouts) [37, 38, 764 765 40, 71, 80].

766

767 **7. Limitations**

In this article, we advocate the extension of the FITT-VP principle from a physiological 768 769 perspective by proposing density as an additional variable that allows for a more fine-grained 770 characterization of the dose of PA. However, the following limitations need to be acknowledged. First, it should be noted that others have already advocated for complementing 771 772 FITT from a psychological perspective by integrating an additional "F" representing "fun" as an umbrella term for psychological factors such as affective valence and enjoyment of PA [385] 773 to reflect that these factors are important determinants of engagement and adherence to PA 774 [386–391]. Second, although we provide in this article a strong theoretical rationale that 775 complementing FITT-VP by the variable density will improve our understanding of the dose-776 777 response relationship between PA and health-related outcomes, we wish to emphasize that 778 the precise characterization or prescription of a specific PA dose will remain a considerable challenge because of the myriad of (i) non-modifiable factors (e.g., age, sex, genetics), (ii) 779 780 potentially modifiable non-PA-related factors (e.g., diet, sleep, stress, environmental conditions), and (iii) modifiable PA-related factors (e.g., type of PA, intensity, duration, 781 movement frequency), which include but are not limited to setting (e.g., home-based or center-782 based, and indoor or outdoor), method of delivery (e.g., in-person or online), level of 783 784 supervision (e.g., no supervision, general supervision, direct supervision) and social interaction individual or group-based), that can influence the dose and individual 785 (e.g., 786 psychophysiological response(s) to PA [37, 38, 40, 45, 54, 71, 364]. In other words, adding 787 density to FITT-VP is another piece of the puzzle to better characterize the dose of PA and, in 788 turn, disentangle its influence on specific health-related outcomes.

789

790 8. Conclusions

791 In summary, we have provided an overview of the implications and the potential of addressing the density of PA as a variable that has been under-recognized when studying the relationship 792 793 between PA and health-related outcomes, using the field of brain health as an example. In 794 view of an increasing interest in understanding the dose of PA including but not limited to 795 "micropatterns" assessed using high-resolution wearable data [119, 120, 392], density is a variable that can complement the traditional concept (i.e., the FITT-VP principle) by 796 considering an additional element - the temporal distribution of PA stimuli within a single bout 797 of PA or between successive bouts of PA relative to the time spent resting. We propose a 798 799 definition for density and approaches for operationalizing it which, in turn, may allow for a more precise determination of the dose of PA for improved health effects and the prevention and 800 801 treatment of chronic disease. Considering that an explicit focus on the density variable has 802 been largely absent from research to date, investing greater effort in understanding it will add 803 fruitful nuance to identifying the dose-response relationship between PA and health-related 804 outcomes (e.g., brain health), and thus has the potential to provide important information on 805 the optimal and minimal beneficial doses of PA.

806	Declarations
807	
808	Authors' Contributions
809	F.H.: conceptualization, writing – original draft, visualization; L.Z., P.T., P.M., R.F., Q.Y., T.L
810	A., C.H., A.F.K., K.E., B.C., Y.C., M.H., Z.Z., T.I., K.K., S.A., Y.G., J.C., M.H., M.H., Z.C., D.M.,
811	V.F., D.R., E.S., M.W. N.O., S.L., H.B.: writing - review & editing. T.G.: writing - review &
812	editing, supervision. All authors read and approved the final version of the manuscript.
813	
814	Acknowledgments
815	The authors have nothing to acknowledge.
816	
817	Ethics approval and consent to participate
818	Not applicable.
819	
820	Consent for publication
821	Not applicable.
822	
823	Availability of data and materials
824	Not applicable.
825	
826	Conflict of interests
827	The authors declare no conflict of interest or competing interests.
828	
829	Funding
830	Not applicable.
831	

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