



Using Free-Living Heart Rate Data as an Objective Method to Assess Physical Activity: A Scoping Review and Recommendations by the INTERLIVE-Network Targeting Consumer Wearables

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Abstract

Wearable technologies open up new avenues for the assessment of individual physical activity behaviour. Particularly, free-living heart rate (HR) data assessed by optical sensors are becoming widely available. However, while an abundance of scientific information and guidance exists for the processing of raw acceleration data, no universal recommendations for the utilization of continuous HR recordings during free-living conditions are available. Towards Intelligent Health and Well-Being: Network of Physical Activity Assessment (INTERLIVE[®]) is a joint European initiative of six universities and one industrial partner. The consortium was founded in 2019 and strives towards developing best-practice recommendations in the context of consumer wearables and smartphones. The aim of this scoping review (following PRISMA-ScR procedures) and recommendations was to provide best-practice protocols for deriving individual physical activity profiles from continuous HR recordings by wearables. The recommendations were developed through an initial scoping review, grey literature searches of promotional material and user manuals of leading wearable manufacturers as well as evidence-informed discussions among the members of the INTERLIVE[®]-network. The scoping review was performed on the generic domains required for physical activity assessment, namely: (1) ‘assessment of maximal heart rate’, (2) ‘determination of basal and/or resting heart rate’ and (3) ‘heart rate-derived intensity zones’, for which we finally included a total of 72, 2 and 11 eligible papers, respectively. Gathering recent knowledge, we provide a decision tree and detailed recommendations for the analysis of free-living HR data to derive individual physical activity profiles. Moreover, we also provide examples of HR-metric calculations that help to illustrate data processing and reporting.

1 Introduction

The development of wearable technology is evolving rapidly [1, 2], providing an easy access to extensive data related to physical activity, fitness, sports performance and health. Wearable devices often comprise of inertial sensors [i.e. inertial measurement units (IMU)] for the estimation of steps by means of acceleration and/or optical sensors [i.e. photoplethysmography (PPG)] that allow for estimations of heart rate (HR) through readings of the pulse wave [3]. In contrast to a 12-lead electrocardiogram (ECG) that directly traces ventricular depolarisation, PPG is based on the absorption and reflection of emitted light by the blood, where the

transmitted or reflected light is modulated by the systolic variations in blood volume that, in turn, is closely linked to HR [3]. Owing to improved battery life and gradual reductions in size and weight [2], wearables are especially advantageous for continuous recordings of bodily functions during free-living conditions over multiple days or even weeks.

Most commonly, collected PPG data are processed through algorithms that are rarely publicly disclosed. Moreover, data quality and validity may differ considerably between devices and are commonly unknown to the user [4]. As such, feedback provided for end-users is typically on the basis of summary data, such as average HR, but it remains unknown how many data points are underlying these data (e.g. non-wear time or periods with low data quality/missing data) [4–7]. While it has been shown that summary data may be sufficient to improve adherence to

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Key Points

Free-living heart rate (HR) data assessed by optical sensors are becoming widely available but guidance on the use to assess free-living physical activity (PA) on the basis of continuous HR is lacking.

Utilizing free-living HR to assess individual PA patterns requires also standardized procedures for the measurement/estimation of maximal HR, basal/nocturnal HR and HR-based intensity zones.

Combining the knowledge retrieved from systematic literature searches and discussions within the INTERLIVE network, this paper provides a decision tree and detailed recommendations for the analysis of free-living HR data to derive individual PA profiles.

regular physical activity and may even lead to improved health-related outcomes, such as body composition [8], these data may not be sufficient for clinical purposes or research settings. This is especially the case when individual activity patterns have to be assessed, which is why for research purposes, mostly devices that provide raw acceleration data have been used [9–11].

While an abundance of scientific information and guidance is available for the processing of raw acceleration data [9–12], it has to be acknowledged that devices providing high-quality data are expensive and may, thus, not be preferred for health promotion purposes. Additionally, other weaknesses of acceleration data regarding the wearing position [9, 11, 13], as well as the detection of low-acceleration type of activities (e.g., cycling, resistance training or uphill versus downhill running) have previously been identified [9]. These constraints somewhat limit the generic use of accelerometers and inertial measurement units, supporting the use of other data assessable by wearable devices. In this context, the use of continuous HR to quantify physical activity patterns during free-living (i.e. time spent at different intensities per day) may be promising. However, to the best of our knowledge, currently no guidance for the standardized intensity classification of free-living HR data exists.

Previous research discussing exercise intensity thresholds has mainly focused on the intensity distribution of individual exercise sessions in an athletic context [14–16]. As a result, several methods were proposed, that are based on important metrics, such as maximal HR (HR_{max}), maximal

oxygen consumption (VO_{2max}) or the HR and VO_2 reserve (i.e. $HR_{max} - \text{resting HR}$ and $VO_{2max} - \text{resting } VO_2$, respectively). For example, the American College of Sports Medicine (ACSM) suggested the classification of very light, light, moderate, vigorous and near to maximal/maximal exercise intensities (Table 1) [17]. However, these models are commonly based on assumed disturbances of the physiological homeostasis and their validity is still a matter of debate [15]. Such an approach would at a minimum require cardiopulmonary exercise testing including breathing gas analysis or the assessment of blood lactate concentrations, both of which are labour intensive. Moreover, free-living physical activity is often characterized by an abundance of light activity which may result in an overrepresentation of the light intensity zone [18], and thus not necessarily providing a sufficient resolution to differentiate between individual physical activity profiles.

Considering the lack of guidance on the use of free-living HR metrics, this paper aims to provide evidence-informed recommendations for the profiling of free-living physical activity patterns on the basis of HR. Further elaborating on the above, this also includes the development of standardized approaches for the assessment or estimation of maximal as well as basal and/or resting HR. The provided guidelines are targeted at researchers and manufacturers, as well as sport and clinical practitioners and aim to facilitate a standardized and harmonized approach for obtaining HR-derived activity profiles assessed by wearables.

2 Background: The Interlive® Network

INTERLIVE® is a joint initiative of the University of Lisbon (Portugal), the German Sport University (Germany), University of Southern Denmark (Denmark), Norwegian School of Sport Sciences (Norway), University College Dublin (Ireland), University of Granada (Spain) and Huawei Technologies Finland. The consortium was founded in 2019 and combines expertise in sports and exercise medicine, health epidemiology, health technology and biostatistics. The main aim of the consortium is to develop best-practice protocols for evaluating the validity of consumer-grade wearables as well as to provide guidance on the utilization of wearable-derived data to foster a widespread use of physical activity indicators. To date, INTERLIVE® has published recommendations for determining the validity of consumer-grade wearable devices for HR [4] and step counts [5] as well as more indirectly derived metrics, such as energy expenditure [7] and maximal oxygen consumption (VO_{2max}) [6].

Table 1 Examples of exercise heart rate (HR)-zones suggested by different authors. NB: these zones are provided for illustration purposes only and are not necessarily endorsed by the INTERLIVE[®]-network

ACSM [9]	ACSM [9]	Roete et al. [41]	Sylta et al. [40]	Jamnick et al. [7]	Rønnestad et al. [39]
%HRR	%HR _{max}	%HR _{max}	%HR _{max}	%HR _{max}	%HR _{max}
<30 (very light)	<57 (very light)	50–59	55–72	65–75 (recovery)	60–82
30–39 (light)	57–63 (light)	60–69	72–82	75–80 (extensive)	
40–59 (moderate)	64–76 (moderate)	70–79	82–87	80–85 (intensive)	83–87
60–89 (vigorous)	77–95 (vigorous)	80–89	87–92	85–92 (training)	88–100
≥90 (near max.)	≥96 (near max.)	90–100	92–97	>92 (interval training)	

3 Methodological Approach

In an initial network-meeting held on 24 October 2022, the process for the development of recommendations for the objective profiling of physical activity on the basis of free-living HR data was discussed. In this meeting, an iterative three-step process for the development of recommendations was agreed upon, consisting of a (1) a scoping review with systematic literature search, (2) a grey literature search of user manuals and other promotional materials of leading wearable manufacturers and (3) evidence-based discussions among the INTERLIVE[®]-network. On the basis of the a priori knowledge of each consortium member, the consortium also agreed on three domains to be targeted during the scoping review that were deemed relevant for assessing free-living physical activity on the basis of HR: (1) methods to assess or predict HR_{max}, (2) methods to assess or predict resting and/or basal HR and (3) methods to determine HR-zones.

The scoping review with systematic literature search was conducted by a sub-group of the INTERLIVE[®] network (M.S., J.F.F. and L.H.). Only papers proposing or validating methods to determine, predict or estimate HR_{max} (i.e. domain 1), basal or resting HR (i.e. domain 2) or HR intensity zones (i.e. domain 3) were included. The systematic literature search was conducted on 6 December 2022 and updated on 26 September 2024, using the checklist for Preferred Reporting Items of Systematic Reviews and Meta-Analysis Protocols extension for Scoping Reviews (PRISMA-ScR). The PubMed/MEDLINE, ISI Web of Science, and SPORTDiscus databases were searched for the previously identified three domains. The search strings were specifically adapted to the search requirements of each database (Online supplementary data, Table S1). Additionally, reference lists of included studies were screened for potentially missing papers. A flowchart of the search process and study selection for the three domains is shown in online supplementary data (Figures S1a to S1c). Papers were eligible when their full text was available, papers were listed in one

of the searched databases and were written in English language. No limit in terms of the publication date was in place.

All results from the online search were saved, imported and further analysed using the Rayyan tool for systematic reviews. The literature search process was performed independently by two authors and included removing duplicates and screening titles, abstracts and full texts. Potential conflicts were resolved by consulting with a third author. Article characteristics such as authors, title, type of paper (e.g. original study, systematic review, narrative review) and main results (e.g. formula for predicting maximal heart rate, proposed heart rate-based intensity zones) were extracted separately for each domain.

In parallel with the scoping review, grey literature searches were also performed by another sub-group of the INTERLIVE[®]-network (M.O.R., A.C. and F.B.O.). This search specifically targeted user manuals, technical documentation and other promotional material of established manufacturers. In this process, we summarized grey literature information for an entry-level as well as medium and high-grade model of selected manufacturers that held large market shares from 2020 to 2022 [19] and have comprehensive manuals and technical documentation of their products publicly available. Namely these manufacturers were Amazfit, Apple, Fitbit, Garmin, Huawei, Polar, Samsung, Suunto and Xiaomi. Devices from the recent product range of each manufacturer were classified into ‘entry-level’ as well as ‘medium and high-grade’ models on the basis of their pricing. The cheapest device in the current product range was categorized as ‘entry-level’, the most expensive as ‘high-grade’, and the device with a price closest to the midpoint between these two as the ‘medium-grade’ model. The extracted data from grey literature included information on the options to extract free-living continuous data (i.e. irrespective of HR that is measured during specific exercise sessions) as well as possible predictions of maximal and/or resting/basal HR. Since continuous HR data are typically displayed in mobile applications, we expanded this grey literature search to the following prominent fitness apps: Zepp (Amazfit and Xiaomi), watchOS 9 (Apple), Fitbit

App, Garmin Connect, Huawei Health, Polar Flow, Samsung Health and Suunto App.

The results of the scoping review and grey literature search were then discussed with the entire consortium in another online meeting held on 21 April 2023. In this meeting, a first draft of recommendations for the objective profiling of physical activity by HR was established, which was further refined by selected members of the network (M.S., J.F.F., L.H. and F.B.O.) and subsequently shared for revisions with the entire consortium.

4 Current State of Knowledge

4.1 Results of the Scoping Review with Systematic Literature Search

We identified a total of 72, 2 and 11 eligible papers for the domains ‘HR_{max}’, ‘basal/resting HR’ and ‘HR-zones’, respectively.

Identified Papers for the HR_{max} Domain

Of the 72 reviewed papers for the HR_{max} domain, 47 attempted to derive or propose unique HR_{max} prediction equations or models (online supplementary data, Tables S2a to S2c), while 25 papers solely aimed at evaluating the validity of already existing equations in different populations (online supplementary data, Table S3). Out of the 47 papers, a total of 106 unique HR_{max} prediction equations were extracted. Of these, 63 equations target healthy non-athletic populations (extracted from 31 papers), 28 equations target athletic populations (extracted from 9 papers), and 15 target diseased populations (extracted from 11 papers). Note that five papers provided equations for multiple populations [20–23].

Identified Papers for the Basal and Resting HR Domain

In contrast to HR_{max}, only limited direct evidence exists on the methods to assess basal and resting HR in children [24] as well as young men [16, 24]. Logan et al. [24], derived resting HR by measuring HR in the morning within 30 min of awakening and compared this to variations in the lowest HR assessed through continuous recordings throughout an entire school day [24]. Depending on the method used, a variance between the morning and day measures of up to 35% was observed. In a similar manner, Davis and Convertino [16] compared nocturnal HR with HR determined in one of the following four conditions: (1) directly after awakening using palpation, (2) after 15 min of rest in a supine position, (3) in a seated position and (4) after 10 min of standing in a quiet room. The lowest HR was observed during the night

but did not statistically differ from the resting HR assessed by palpation immediately after awakening. Both the night and morning condition differed substantially from all other conditions.

Identified Papers for the HR-Zones Domain

Additionally, only few papers were identified that directly addressed the determination of HR-zones to cluster exercise or physical activity on the basis of their intensity. The majority of included papers focused on HR-zones that were assessed in accordance with physiological variables, such as %VO_{2max} [16], measures of the ventilatory/lactate thresholds [25–29], or the point of metabolic acidosis [29, 30]. In addition, studies have compared a percentage range above resting HR compared with a percentage of HR_{max} in cardiac patients [31] or used the HR at the critical power to demarcate heavy from severe exercise intensities in young women [32]. Furthermore, another study compared HR-based indices to global positioning system (GPS)-derived training load in professional soccer players [33]. Finally, we identified two review papers that focused on the applicability of different exercise prescription methods based on HR [15, 34]. The ACSM provides recommendations for the classification of exercise intensity on the basis of HR_{max}, VO_{2max} and heart rate reserve (HRR) [34]. Jamnick et al. concluded that estimating training intensity zones based on maximal anchors, such as a percentage of HR_{max} or VO_{2max}, is inaccurate because this method is not consistent with physiological parameters that delineate these intensity zones [15]. However, this is typically discussed in well-trained athletes and whether this holds true for untrained or sedentary populations remains unknown.

Taken together, the findings of our scoping review indicate an abundance of scientific data on HR_{max} assessment, which may be used to directly derive recommendations. Conversely, the scientific evidence on the recommended methods of assessing resting and/or basal HR appears to be insufficient to draw conclusions. Similarly, only few papers have directly addressed the determination of HR-zones. The majority of these papers clearly outline the physiological challenges associated with the definition of intensity thresholds. Moreover, in all included papers, HR-zones were primarily developed to quantify exercise intensity rather than free-living physical activity.

4.2 Results of the Grey Literature Search Focusing on Information Provided by Manufacturers

A summary of the data retrieved from user manuals of established manufacturers is provided in online supplementary data Table S4a. While 10 models did not specify which equation is used to predict the HR_{max}, the remaining 17

models appear to use the Fox et al. [35] prediction equation ‘220 – age’. This is of particular concern as the lack of scientific basis for this equation was previously shown [36] and numerous population-specific and validated alternatives are available (online supplementary data Tables S2a to S2c). Additionally, some manufacturers, such as Polar, allow users to input the individually assessed HR_{max} manually.

Of the 27 reviewed wearables, only 4 models specified their methodology to derive resting HR. All three included models by Polar require the user to lie supine and breathe calmly for 3–5 min. Additionally, the high grade Amazfit model uses the nocturnal HR measured over at least 5 h to estimate the resting HR. All other models simply instruct the user to wear the device continuously throughout the day without further specification.

Concerning the analysis of continuous HR data, Amazfit, Apple, Fitbit, Samsung and Xiaomi allow for a user-friendly download of continuous data through the corresponding app (online supplementary data Table S4b). Huawei and Garmin allow for an export of these data through an additional developer tool, while Polar and Suunto currently only provide an option to download HR data from individual training sessions. Interestingly, out of the nine manufacturers searched, only five (Amazfit, Fitbit, Garmin, Polar and Xiaomi) [37–39] currently display free-living activity zones on the basis of continuous HR, with considerable inconsistencies among the classifications used (Table 2).

4.3 Special Considerations for Physical Activity Profiling

On the basis of the scoping review with systematic literature search, grey literature search and a priori knowledge of the INTERLIVE®-network, the following considerations concerning the three domains (i.e. ‘ HR_{max} ’, ‘basal/resting HR’ and ‘HR-zones’) provided the foundations for the developed recommendations of profiling free-living physical activity on the basis of continuous HR measures. An additional

overview on variables that require attention is provided in Table 3.

HR_{max} Assessment

The majority of studies reporting HR_{max} rely on HR values obtained from a common incremental VO_{2max} test, such as the Bruce protocol [17]. However, it remains inconclusive whether HR_{max} is affected by the protocol characteristics. In two studies, no differences were observed between the HR_{max} obtained with 1 and 3-min increments [40, 41]. However, Machado et al. [42] reported an optimum increment duration of 2 min. Furthermore, it was shown that HR obtained in traditional incremental tests may be lower (5.76 ± 2.81 bpm) than that obtained from specifically designed 3–4 min all-out performance [43].

Since the optimal protocol for assessing HR_{max} remains unknown, it is reasonable to suggest that HR_{max} should be assessed through a graded maximal exercise test that is in line with the population specific standard exercise testing procedures recommended by the ACSM [17]. It should also be considered that longer increments may lead to premature fatigue and prevent the attainment of HR_{max} [15, 40, 42]. Therefore, shorter stage durations (≤ 2 min) or ramp protocols (with 30–60 s per increment) are preferred. Similar to VO_{2max} testing, a total duration of 8–12 min seems optimal to assure maximal cardiovascular exertion without premature fatigue [15]. Irrespective of the protocol used, secondary criteria to determine maximal voluntary exhaustion may be applied. These commonly include respiratory exchange ratio (RER), subjective ratings of perceived exertion or blood lactate concentrations [15, 17].

In addition to the protocol characteristics, the selection of the exercise mode seems crucial. As HR is dependent on muscle mass involvement [44, 45], treadmill tests are considered a gold-standard. VO_{2max} values achieved using treadmill protocols tend to be up to 20% higher compared with cycling protocols [45]. Importantly, a HR_{max} derived from a treadmill protocol likely best reflects the HR during

Table 2 Examples of free-living HR-zones currently used by leading manufacturers. NB: these zones are provided for illustration purposes only and are not necessarily endorsed by the INTERLIVE®-network

Fitbit %HRR	Zepp (Xiaomi and Amazfit) % HR_{max}	Garmin HR and ACC based*	Polar HR and ACC based*
40–59 (fat burn)	> 50 (relaxed)	Below training	Resting
	50–60 (light)	Warm up	Sitting
60–84 (cardio)	60–70 (intensive)	Easy	Low
	80–90 (aerobic)	Aerobic	Medium
> 85 (peak)	90–99 (anaerobic)	Threshold	High
	100 (maximum)	Maximum	

*Manufacturers use a combination of HR and accelerometer (ACC) data to determine daily physical activity zones; exact HR zones are not specified

Table 3 Factors affecting maximal and basal/resting heart rate

Maximal heart rate		Basal/resting heart rate	
Stable	Transient	Stable	Transient
Age ~ [52, 54, 112, 119–123]	Increment duration ~ [15, 40–43]	Age [24, 84, 86, 136]	Acute infections and disease ↓ [142–144]
Sex ~ [23, 47, 50–52, 54, 55, 58, 66, 79, 112, 124–126]	Stage duration ~ [15, 40–43]	Sex [87, 137]	Psychological stress ↑ [145]
Day-to-Day variability ↓ ↑ [48]	Overall test duration ~ [15, 40, 42]	Day-to-day variability [138]	Environmental conditions [146]
Individual preconditions [17]	Exercise mode:	Time of day [16]	Ambient light [147]
Body mass [55]	Treadmill ↑ [44, 45, 52]	Cardiovascular fitness [139]	Sleep quality [92]
Body fat [22, 55]	Cycle ergometer ↓ [17, 44–46, 52]	Body composition [140]	Sleep phase [92]
Ethnicity ~ [50, 51, 55, 67, 127–130]	Sport specific ↑ [47]	Environmental conditions [95]	Deviations from regular bedtime [148]
Training status ~ [23, 47]	Insufficient recovery & fatigue ↑ [17, 43]	Training status [104, 141]	Body position: [85, 149–151]
Disease status:	Nutritional status [17]	Chronic medication ↓ ↑ [17]	Supine ↓
Cardiovascular ↓ [35, 54, 59, 60, 65, 68–70, 73]	Environmental conditions [17]		Sitting ↑
Chronic fatigue syndrome ↓ [131, 132]	Activated muscle mass [47, 52, 60, 66, 134, 135]		Physical activity: [95, 96, 152]
Obesity ↓ [22]	Activity type ↓ ↑ [47, 52, 60, 66]		Time since
Cerebral palsy ↓ [72]	Transient medication ↓ ↑ ^b [17]		Rest period
Mental retardation ↓ [20]			Intensity
Smoking status ~ [58, 133]			Duration
Chronic medication ↓ ↑ ^a [17]			Food intake ↑ [153–155]
			Nicotine ↑ [156, 157]
			Caffeine ~ [158–161]
			Transient medication ↓ ↑ [17]

↑ Increase, ↓ decrease, ↓ ↑ increase or decrease depending on specific conditions, ~ inconclusive evidence

^aMedication that is taken continuously over a longer period of time to treat chronic disease

^bMedication that is taken only once or over a short period of time to treat acute disease

NB: Appendix A (Table A.1) of the ACSM's 'Guidelines for Exercise Testing and Prescription' [17] provides an overview of common medications and their expected (chronic) influence on the HR kinetics

free-living conditions for most populations and testing e.g. on a cycle ergometer may cause early local muscle fatigue, preventing cardiorespiratory exertion [17]. Cycle ergometers may, however, be considered an alternative for clinical populations that are unable to perform treadmill protocols [17, 46]. Furthermore, for specifically trained athletes a sport specific ergometer may be used to attain the actual HR_{max} [47].

Irrespective of the testing mode, standardisation of the testing conditions is required. For example, fatigue and insufficient recovery from previous exercise may acutely affect the HR response. In this context, Ingjer [43] found that after one or two days of high intensity training only few individuals were able to reach their previously tested HR_{max} . Therefore, no strenuous exercise should be performed a minimum of 24 h before the testing procedure [17]. Detailed recommendations for other factors, such as nutritional status, environmental conditions and individual preconditions, are found elsewhere [17]. Importantly, once the test is performed in standardised conditions, the mean day-to-day-variability of HR_{max} in healthy but untrained populations appears to be as low as 1% [48].

Maximal tests are often labour-intensive and may require extensive equipment. Moreover, without medical clearance these are typically restricted to young and healthy

populations [17]. Thus, prediction models were developed to estimate HR_{max} on the basis of individual characteristics, such as age and sex. Our scoping review revealed that nearly all ($n = 44$) papers that derived prediction equations included age among other variables, while 42 papers used age alone (online supplementary data Table S2a–c). However, even though univariate age-based equations are most frequently used because they are simply applicable and relatively easy to understand, the majority of these equations seem to be prone to large prediction errors [36]. Thus, several studies aimed to develop multivariate ($n = 13$) equations or population-specific univariate equations to increase the predictive capacity by including additional factors, comprising of sex [23, 47, 49–54], resting HR [55–60], HR during submaximal tests [57, 61–65] and type of activity [47, 52, 60, 66]. Additionally, individual studies also considered HR variability [62], body mass and body fat [55], ethnicity/nationality [50, 67], specific diseases [20, 59, 65, 68–72] and medication [73]. As it has been shown that the most common prediction equations are valid for adults but not for children and adolescents [72, 74–80], separate equations for children and adolescents have been proposed, including variables like resting HR, maturity offset, body mass and body fat as additional variables [55, 56, 78].

Basal/Resting HR Assessment

Our scoping review revealed a dramatic lack of scientific guidance on how to assess resting and/or basal HR. Generally, the lowest levels of HR refer to a condition where the metabolic requirements are minimal [81]. Thus, it appears reasonable to take advantage of methods that are used to assess basal and/or resting metabolic rate. Basal metabolic rate was initially defined as the minimal rate of energy expenditure compatible with life [82]. This is typically assessed as the heat production (or VO_2 as the surrogate of energy expenditure) at rest and in a supine position in strictly controlled laboratory environments, including a fasted state with controlled environmental conditions [83]. Basal metabolic rate, in turn, needs to be distinguished from nocturnal metabolism, which typically appears to be lower [83], also indicating that the lowest values of HR are expected to be observed during sleep. In fact, it was shown that the HR assessed by palpation directly after awakening was statistically lower than HR measured during a later time of day [16], indicating that lowest values of HR were obtained nocturnally.

Importantly, basal and nocturnal metabolic rate differ from resting metabolic rate, which is typically assessed in less strict laboratory conditions, i.e. at any given state of rest throughout the day. As such, resting HR appears to be higher than nocturnal or basal HR [84]. Nonetheless, resting HR is considered a key vital sign and is a well-established predictor of all-cause and cardiovascular mortality [85] and commonly included in various studies [85]. It appears that resting HR assessments generally involve a resting period between 5 and 10 min [86–88], while the participants are in a supine position [89–91] and the HR is obtained from the last minute of the measurement.

When assessing nocturnal HR, the effects of sleep quality need to be considered. It has been shown that deep and quiet sleep [i.e. non-rapid eye movement (NREM)] is associated with a lower HR compared with restless and superficial sleep [i.e. rapid eye movement (REM)] [92]. In healthy individuals, good sleep quality includes about four to six cycles per night, with each cycle lasting an average of 90 min [93]. However, it remains unknown how many sleep cycles are required for overnight HR assessments. There are a number of studies that incorporated measures of nocturnal HR variability, with a typical duration of 4 h commencing 30 min after reported bedtime [94–99]. This seems reasonable considering the length of individual sleep phases (i.e. 90 min) and is also in line with data showing that the early phases of sleep appear to be a quiet sleep period [100].

HR-Zone Determination

With the rapid increase of commercially available wearables that allow for continuous free-living HR assessment, individual physical activity profiles may be determined. In athletic populations, HR-zones are often aligned with changes in the metabolism, represented by ventilatory and/or lactate thresholds. Indeed, the majority of studies identified by our scoping review used indices of the metabolism as a determinant of HR-zones [15, 16, 25–28, 30, 33]. Especially in endurance athletes, typically a three-zone model is used to cluster the exercise intensity into light (below the first ventilatory/lactate threshold), moderate (between the first and second ventilatory/lactate thresholds) and severe/vigorous (above the second ventilatory/lactate threshold) [14]. However, while these submaximal anchors were used to describe exercise intensity by numerous studies, the actual validity is still debated [15]. The criticism brought up refers to whether these zones make a demarcation that reflects actual homeostatic disturbances [15]. In fact, numerous methods for the determination of lactate thresholds exist [15] and, thus, errors in the definition of the lactate thresholds can have a detrimental effect on the classification of intensity distributions. Moreover, graded exercise testing to voluntary exhaustion is needed to accurately determine these thresholds [101, 102], thus limiting its utilization to specific populations.

For example, our grey literature search of leading manufacturers revealed various fixed HR zones that are on the basis of percentages of the HR_{max} or HRR and are used for scientific purposes (Table 1) or provided to end-users (Tables 2, 4). However, the underlying assumptions for deriving these zones remain unknown, leading to inconsistencies among different methods. This also further underlines that the suggested zones do not reflect the underlying metabolic fluctuations. That is, large ranges of fixed percentages of HR_{max} were previously associated with both the ventilatory threshold (60–90% of HR_{max}) and the maximal lactate steady state (75–97% HR_{max}) [103]. Thus, it appears that fixed percentages of HR_{max} to prescribe exercise intensity do not demarcate distinct physiological characteristics [30].

In addition to the methodological difficulties outlined above, exercise intensity is also affected by resting HR which, in turn, is dependent on individual factors such as the training status [104]. Consequently, for the same HR_{max} , individuals with a lower resting HR (i.e. trained individuals) will be able to produce more work compared to individuals with a higher resting HR (i.e. untrained individuals). Thus, simply quantifying exercise intensity by percentage of HR_{max} will easily lead to an over- or underestimation of the actual activity performed. To overcome these issues, often the HRR has been used to quantify physical activity in various populations [28, 34]. In fact, guidelines for the quantification of physical activity on the basis of HRR

are also provided by the ACSM, suggesting five zones as follows: (1) near maximal to maximal ($\geq 90\%$ HRR), (2) vigorous (60–89% HRR), (3) moderate (40–59% HRR), (4) light (30–39% HRR) and (5) very light ($< 30\%$ HRR) [17]. However, in an earlier position stand even the ACSM clearly outlines the limitations that are associated with the classification of these zones, also when based on HRR [34]. This is because large inter- and intraindividual differences exist in the relationship between HR_{\max} and $VO_{2\max}$ as well as HRR and $VO_{2\text{reserve}}$ (i.e. $VO_{2\max} - VO_{2\text{rest}}$) [41, 105, 106].

Nevertheless, it appears that the use of HRR seems most appropriate for the purpose of assessing exercise or activity intensity. However, the intensity-zones used so far mainly serve to distinguish high-intensity activities/exercise from moderate- and low-intensity activities/exercise that are expected to lead to an improvement in cardiorespiratory fitness [34]. However, free-living physical activity is characterised by an abundance of low-intensity activity below the first threshold [18]. It is, therefore, questionable whether there is sufficient resolution of these activities on the basis of established intensity-zones. In fact, it is important to bear in mind that even lower intensities in the range of 30–45% $VO_{2\text{reserve}}$ can lead to an increase in cardiorespiratory fitness [107] that may not be adequately represented by commonly suggested HRR zones, especially in untrained or diseased populations. Accordingly, a higher resolution of the low-intensity activities is required to accurately describe free-living physical activity. Moreover, particular settings that outline individual changes in daily activity patterns, rather than describing the intensity of a single exercise session, may benefit from a more detailed depiction of these activities.

Considering the demands on methods to quantify free-living physical activity by continuous HR on the one hand, while bearing in mind the physiological variability on the other hand, it appears that the selection of the preferred method highly depends on its purpose. For example, applying the ACSM guidelines also allows to assess adherence to common physical activity guidelines, such as e.g. outlined by the World Health Organization (WHO) [108]. For healthy adults, although caution should be exercised as these

recommendations are based on self-reported data, 150 min of weekly moderate activity are suggested, where moderate is defined as an energy expenditure of 3–6 metabolic equivalents (METs). According to the ACSM intensity thresholds, this would be reflective of 40–59% HRR [17]. However, if the aim is to compare physical activity patterns inter- and/or intra-individually, dividing HRR into clusters of 10% (i.e. 10, 20, 30% etc.) might be an appropriate approach to display individual activity profiles. This approach would allow for a higher resolution of individual free-living activity patterns by using smaller increments that also allow the capture of small changes in activity intensities. More importantly, this approach does not rely on universal group-based thresholds. In fact, utilizing relatively fine increments for the quantification of continuous HR would also be in line with the subjective rates of perceived exertion, as assessed by the modified rating of perceived exertion (RPE) scale (Borg CR10) [18, 109, 110]. The universal use of the RPE scale, however, is debated, as large inter-individual variability exists, especially among individuals with different fitness levels [111]. Nevertheless, aligning the 10% clusters derived from the HRR with the ten increments of the RPE scale may aid in interpreting the individual activity profiles.

5 Evidence-Informed Recommendations

In Fig. 1, we present a decision tree that illustrates the necessary steps to derive physical activity profiles from free-living HR in prospective data collections or existing data sets. Detailed recommendations on each of these levels are provided in Tables 5, 6 and 7. Importantly, our recommendations may only be applied if the pre-processed continuous HR data (as opposed to daily summary data) can be accessed. Thus, caution is warranted when selecting appropriate devices for the free-living HR assessment. This also includes gathering possible information on the accuracy of the selected device [4–7]. It should be also noted that the handling of continuous HR data requires special attention. However, providing standards for the mathematical and

Table 4 Examples of exercise HR-zones currently used by leading manufacturers. NB: these zones are provided for illustration purposes only and are not necessarily endorsed by the INTERLIVE[®] network

Fitbit %HRR	Zepp (Xiaomi and Amazfit) %HR _{max}	Garmin %HR _{max}	Polar %HR _{max}
40–59 (fat burn)	> 50 (relaxed) 50–60 (light)	50–60	50–60 (very light)
60–84 (cardio)	60–70 (intensive) 80–90 (aerobic)	60–70 70–80	60–70 (light) 70–80 (moderate)
> 85 (peak)	90–99 (anaerobic) 100 (maximum)	80–90 90–100	80–90 (hard) 90–100 (maximum)

statistical processing, such as data extraction, smoothing and outlier detection was beyond the scope of the present paper.

Irrespective of whether a prospective data collection is planned, or existing data are to be analysed retrospectively, we recommend that HR_{max} is assessed through a standardised protocol (Table 5). However, in case the direct assessment of HR_{max} is not possible, e.g. owing to medical reasons, HR_{max} may be predicted on the basis of existing models. Since studies that evaluated the validity of HR_{max} equations revealed that commonly used universal age-based prediction equations, such as Fox [35] or Tanaka et al. [112], tend to over- or underestimate the HR_{max} when applied to

specific populations, it is recommended to choose an equation or model that best represents the target population. An overview on available formulas is presented in online supplementary data Tables S2a–c.

As outlined above, using fixed anchors such as the % of HR_{max} may lead to inconsistencies in deriving distinct HR zones. Therefore, we suggest utilizing the HRR for quantification of individual physical activity. Consequently, it is necessary to assess nocturnal or resting HR. The aim is to define the lowest physiological HR for a given individual. Thus, we recommend assessing nocturnal HR through continuous HR recordings. Only in cases where overnight data are not

Fig. 1 Decision tree for the profiling of free-living physical activity by continuous heart rate (HR) measures on the basis of the calculation of heart rate reserve (HRR). Branch A illustrates the necessary steps that should be planned in prospective data collections. Branch B provides guidance for the analysis of already existing data sets (i.e. retrospective analysis). To select the appropriate equation to estimate HR_{max} , a three-step process is suggested: (1) population (i.e. ‘healthy’, ‘athletes’ or ‘diseased’), (2) age, (3) other specifications, such as further characteristics of the athletic background (i.e. type of sport) or disease (i.e. type of disease)

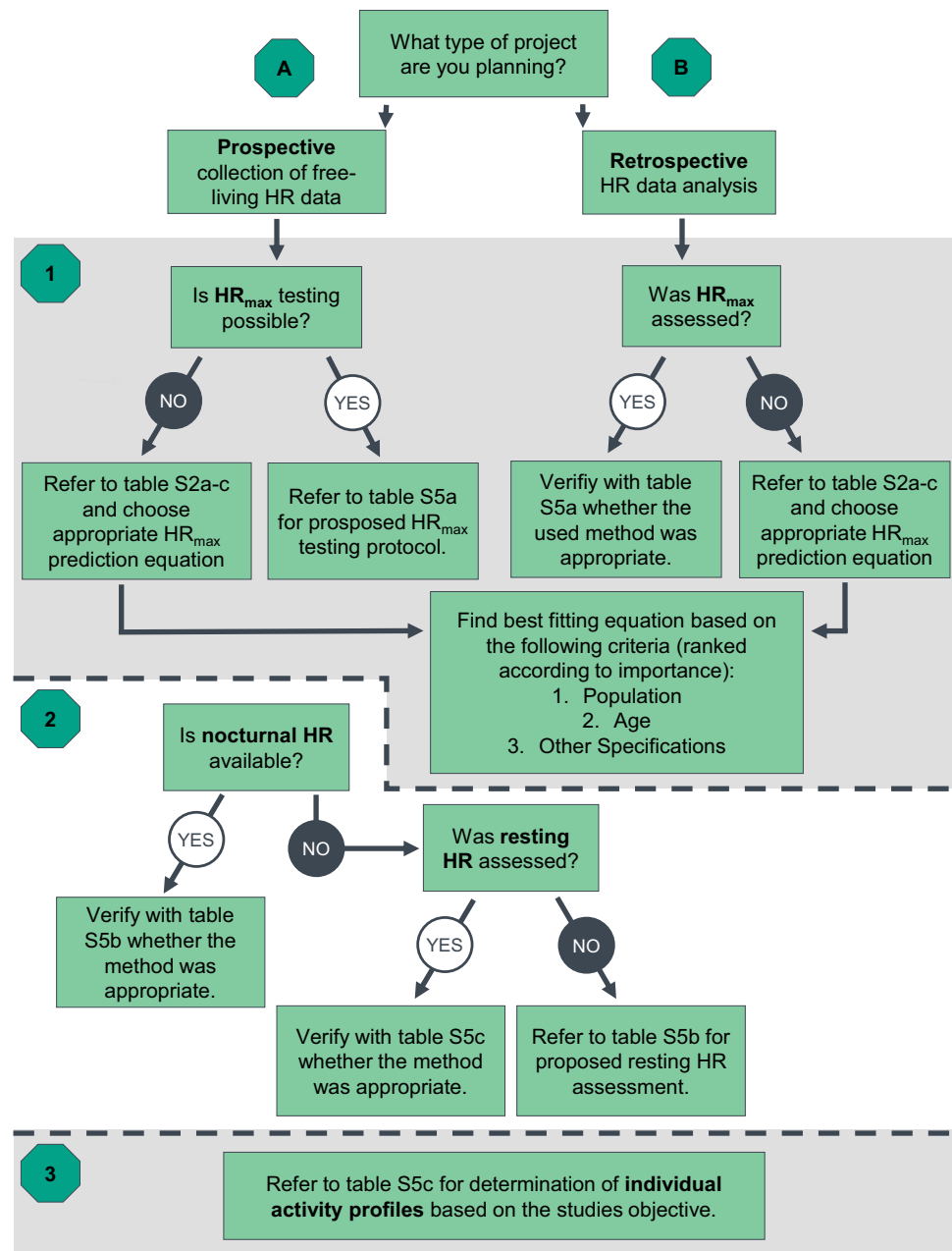


Table 5 Proposed protocol for the assessment of maximal heart rate (HR_{max})

Methodological domains	Methodological variables	Protocol considerations	Reporting considerations
Experimental HR_{max} assessment	Pre-test preparation	<p>Depending on the population of interest, a medical screening is recommended. The Guidelines for Exercise Testing and Prescription of the American College of Sports Medicine provide an overview [17]</p> <p>Participants using regular medication that affects cardiovascular function (e.g. beta blockers) should be asked to continue intake as usual</p> <p>If medically possible, avoid transient medications (e.g. NSAIDs, stimulants, antihistamines, antibiotics, cold medication) for at least 24 h before testing or longer if necessary owing to the half-life of the drug</p> <p>Participants should refrain from intense physical activity 48 h prior to the maximal test</p> <p>Participants should be informed about the testing procedures and preparations. A sufficient amount of sleep (e.g. 7–9 h for adults [162]) prior to the scheduled testing</p> <p>Restrict nutritional intake 3 h prior to the test to avoid gastric complications. Habitual caffeine intake is of no concern</p>	Pre-test standardisation should be reported
	Test standardisation	<p>Testing should be performed in standardised ambient conditions (temperature 20–22 °C, humidity < 60%)</p> <p>We recommend performing the test at a time the individual is habituated with and well-nourished</p> <p>Use a 12-lead ECG or chest strap that has been shown to have an excellent agreement with a gold standard to obtain HR. An overview on suitable devices is presented elsewhere [4]</p> <p>As long as a validated ECG or chest strap is used for the assessment of HR_{max}, sampling rate is of no concern</p>	Report ambient conditions, time of day and the device used for HR recording (including the sampling rate and version of the firmware)
	Maximal test design	<p>An 8–12-min graded exercise test to exhaustion should be performed (duration of each increment ≤ 2 min)</p> <p>Generally, a treadmill test should be preferred. For clinical populations unable to walk or run, a cycle-ergometer test is acceptable. For specifically trained athletic populations, a sport specific ergometer and/or protocol should be preferred</p> <p>Allow for a population-specific warm-up (2–5 min) at self-selected low pace</p> <p>Verify voluntary exhaustion and other maximal criteria according to recommendations provided by the American College of Sports Medicine [17]</p>	<p>Report specifics of the testing protocol (i.e. step length, overall test duration and velocity or power per increment)</p> <p>Report voluntary exhaustion criteria</p>
	Data analysis	<p>Select the highest HR value provided</p> <p>NB: in cases where data are provided beat-by-beat, conversion to HR data for every second is required. Data should be smoothed by 15–30 s rolling averages. The highest 15–30-s rolling average should be considered as HR_{max} [6]</p> <p>If data are obtained by a PPG-based wearable device, it is advised to choose a device that uses a constant high sampling rate. However, in any case, information on the sampling rate has to be obtained prior to selecting a device. Data points that are within the reported sampling rate range should be normalized to the highest sampling rate on the basis of the previous recorded HR value. Missing data (i.e. defined as deviations from a given sampling) should be excluded from the analysis</p>	<p>Report software/application version for data processing (including download and analysis)</p> <p>Report HR_{max} and how it was calculated (e.g. highest 30 s rolling average)</p>

Table 5 (continued)

Methodological domains	Methodological variables	Protocol considerations	Reporting considerations
Prediction of HR_{max}	Selection of prediction model	<p>The selection should be based on the target population. Please refer to supplementary online data, Tables S2a–c for an overview of existing models. For cross-validation of existing models, check supplementary online data, Table S3. Since it is impossible to account for all confounders within the prediction models, we recommend basing the selection mainly on population (healthy, diseases, athletes), age and sex. Some of the available prediction models are activity-specific. However, continuous HR measures during free-living conditions typically include a variety of activities. We recommend models that are specific to walking and/or running since this has been shown to yield the highest values of HR_{max} for the general population.</p>	Report prediction equation and justification

NB: in longitudinal study designs lasting several years, HR_{max} should be updated regularly at pre-determined time points
 HR , heart rate; HR_{max} , maximal heart rate

available, e.g. owing to discomfort that is often experienced by individuals wearing wearable technology while sleeping, should resting HR be assessed. For recommendations on how to assess nocturnal/resting HR, please refer to online supplementary data, Table 7.

The selection of the method applied to quantify physical activity by HRR should be aligned with the individual research question. Adhering to classical definitions as e.g. provided by the ACSM may allow assessment of adherence to common physical activity guidelines and may also facilitate cautious comparisons to studies that have used accelerometry. However, the physiological inconsistencies underlying these thresholds need to be acknowledged. Considering the limitations that has been brought forward concerning intensity zones based on fixed absolute or relative anchors (see special considerations for determining HR-zones in the section “[HR-Zone Determination](#)”), we do feel it is not always desirable to actually report time in specific zones that are linked to metabolic factors but rather provide an individual profile/distribution for each participant. We recommend this to be obtained from HRR, and it can be reported as time spent in arbitrary 10% clusters. Detailed recommendations for the calculation of activity profiles are presented in Table 6. Alternatively, the continuous HR data can be clustered according to the standard intensity zones, for example, as defined by the ACSM: very light, light, moderate, vigorous and near to maximal/maximal exercise intensities (Table 1).

6 HR-Metric Calculation and Reporting

On the basis of the considerations discussed above, this section provides a simplified best-practice example of HR-metric calculation for free-living physical activity profiles. The data represent 24-h of an active and inactive day of the same individual (Fig. 2). Data were collected using a Garmin Vivoactive® 4 smartwatch. The sampling interval was 1 min throughout each 24-h data collection. For nocturnal HR assessment, bedtime and time of awakening were reported by the participant in a sleep diary. HR_{max} was assessed during a graded exercise test to exhaustion on a treadmill using the Firstbeat Bodyguard® 2 2-lead ECG. HR_{max} , nocturnal HR and activity profiles were calculated using MATLAB (R2023a, Mathworks, Inc., USA). Importantly, with this example we are not aiming to provide guidance on data handling (i.e. statistical processing, including handling of missing data) but rather illustrate in a simplified manner the necessary steps described in this paper (i.e. HR_{max} assessment, nocturnal HR assessment and HR-Zones determination) to profile physical activity by free-living HR.

Table 6 Proposed protocol for the assessment of nocturnal and/or resting HR

Methodological domains	Methodological variables	Protocol considerations	Reporting considerations
Assessment of nocturnal HR	Preparation	<p>Participants should be informed about the measurement procedures</p> <p>Avoid assessment of nocturnal HR during periods of psychological and/or physical overload (i.e. acute intense exercise) and acute sickness (e.g. flu)</p> <p>Avoid meals and other substances (e.g. caffeine) for 3 h prior to reported bedtime</p> <p>If medically possible, avoid transient medications (e.g. NSAID's, stimulants, antihistamines, antibiotics, cold medication) for at least 24 h before testing or longer if necessary owing to the half-life of the drug. Medication that is prescribed chronically should be continued as usual</p> <p>During the period of nocturnal HR recording, diaries on physical activity (including exercise) should be kept</p>	Pre-test standardisation should be reported Report physical activity throughout the days of nocturnal HR assessment
	Data collection	<p>Assure that the wearable has been worn during bedtime. Use questionnaires and sleep diaries to control for bedtime</p> <p>If bedtime was not reported, visual inspection of the data is required to verify the device was indeed worn during bedtime (see Data analysis below for details)</p> <p>Wearables do provide information on sleep onset and quality. However, data quality and validity are often not scientifically tested. We suggest not to use those wearable-derived data but rather refer to the two approaches above</p> <p>Select the highest possible sampling rate</p> <p>Since accelerometer-derived data showed that 4 days are sufficient to estimate the physical activity levels of an entire week [11], data should be collected for a minimum of 4 and optimally 7 days</p>	<p>Report the used device and sampling rate (including the firmware)</p> <p>Report whether bedtime was documented and/or controlled</p> <p>Report possible concerns related to sleep quality and well-being</p>
	Data analysis	<p>Use 2-h rolling averages for the duration of reported bedtime. Select the lowest value as nocturnal HR</p> <p>In case bedtime was not reported but the device was worn, 2-h rolling averages should be calculated on the basis of the visually confirmed bedtime</p> <p>NB: some devices may adjust the sampling rate based on the activity performed (e.g. providing data every 5 s up to several minutes). It is advised to choose a device that uses a constant high sampling rate. However, in any case, information on the sampling rate has to be obtained prior to selecting a device. Data points that are within the reported sampling rate range (according to the manufacturer guidelines) should be imputed by using a last value carried forward method on the basis of the highest sampling rate. Missing data (i.e. defined as deviations from a reported sampling rate range) should be excluded from the analysis</p>	<p>Report software/application version for data processing (including download and analysis)</p> <p>Report mean nocturnal HR</p>

Table 6 (continued)

Methodological domains	Methodological variables	Protocol considerations	Reporting considerations
Assessment of resting HR	Test preparation	<p>Assessment of resting HR is only recommended in cases where nocturnal HR data are not available (i.e. the device was not worn during bedtime)</p> <p>Participants should be informed about the testing procedures and preparations. Sufficient sleep (i.e. 7–9 h) should be assured prior to the scheduled testing</p> <p>Habitual food and caffeine intake is of no concern</p> <p>Participants using regular medication that affects cardiovascular function (e.g. beta blockers) should be asked to continue intake as usual</p> <p>Avoid transient medications (e.g. NSAID's, stimulants, antihistamines, antibiotics, cold medication) for at least 24 h before testing or longer if necessary owing to the half-life of the drug</p> <p>Participants should refrain from intense physical activity 48 h prior to the maximal test</p> <p>Participants should avoid any type of activity immediately prior to the testing, including commuting to the lab</p>	Report pre-test standardisation
	Test standardisation	<p>Testing should be performed in standardised ambient conditions (temperature 20–22 °C, humidity <60%)</p> <p>Ambient light conditions are acceptable; however, we recommend avoiding any sources of unnatural light</p> <p>We recommend the test to be performed in close proximity to awakening. As such, the preferred time of the test should be the early morning</p> <p>Use a 12-lead ECG or chest strap that has been shown to have an excellent agreement with gold standard data to obtain HR. An overview on suitable devices is presented elsewhere [4]</p>	Report ambient conditions, time of day and the device used for HR recording (including the sampling rate and version of the firmware)
	Testing procedure	<p>We recommend resting HR to be assessed in supine position for a duration of 10 min</p> <p>As long as a validated ECG or chest strap is used for the assessment of resting HR, sampling rate is not of concern</p>	Report any deviations from the recommended protocol
	Data analysis	<p>We recommend data be analysed by rolling averages over 15–30 s. The lowest rolling average should be considered as resting HR</p> <p>NB: in cases where data are provided beat-by-beat, conversion to HR data for every second is required</p>	Report software/application version for data processing (including download and analysis) Report resting HR

Table 6 (continued)

Methodological domains	Methodological variables	Protocol considerations	Reporting considerations
Alternative resting HR estimation	Data collection	<p>Assure that the wearable has been worn during day time (i.e. from 07:00 am to 10:00 pm)</p> <p>Visual inspection of the data is required to verify the device was indeed worn during daytime (see Data analysis below for details)</p> <p>Select the highest possible sampling rate</p> <p>Since accelerometer-derived data showed that 4 days are sufficient to estimate the physical activity levels of an entire week [11], data should be collected for a minimum of 4 and optimally 7 days</p>	<p>Report the used device and sampling rate (including the firmware)</p> <p>Report factors that may have interfered with an accurate resting HR estimation</p>
	Data analysis	<p>Use a fixed rolling average to find the window for the lowest HR throughout the day. Choose the length of that window on the basis of sampling rate. Since sampling rates tend to vary between few seconds and up to 10 min, the recommended length should be between 5 and 15 min</p> <p>NB: some devices may adjust the sampling rate based on the activity performed (e.g. providing data every 5 s up to several minutes). It is advised to choose a device that uses a constant high sampling rate. However, in any case, information on the sampling rate has to be obtained prior to selecting a device. Data points that are within the reported sampling rate range (according to the manufacturer guidelines) should be imputed by using a last value carried forward method on the basis of the highest sampling rate. Missing data (i.e. defined as deviations from a reported sampling rate range) should be excluded from the analysis</p>	<p>Report software/application version for data processing (including download and analysis)</p> <p>Report mean resting HR</p>

NB: nocturnal HR estimation is recommended as long as HR was recorded throughout the reported bedtime. If that is not the case, resting HR should be assessed. If both nocturnal and resting HR are not available, we propose an alternative procedure as outlined in Table 7. For intra- and inter-individual comparisons, the same approach needs to be applied (i.e. nocturnal or resting HR should be used). In longitudinal study designs, nocturnal and resting HR should be updated regularly at pre-determined time points

HR, heart rate

Table 7 Proposed protocol for determination of heart rate-based individual activity profiles. The selection of an appropriate method should be based on the specific purpose

Methodological domains	Protocol considerations	Reporting considerations
Determining individual activity profiles	<p>Since studies on accelerometer-derived data showed that 4 days are sufficient to estimate the physical activity levels of an entire week [11], we recommend to include the mean of at least 4 full days in order to determine activity profiles for a given week (i.e. when comparing activity levels in longitudinal study designs). A valid day should include a minimum of 10 h [11] of valid data during waking hours or 16 h [9] over the entire 24-h-cycle</p> <p>We recommend to determine individual activity profiles, using HRR. HRR should be calculated as follows: $HRR = HR_{max} - \text{nocturnal HR}$</p> <p>In case nocturnal HR is not available, the assessed resting HR should be used for calculation of HRR. However, for intra- and inter-individual comparisons, the same procedure is required</p> <p>If the purpose is to assess adherence to common physical activity guidelines or an intensity classification based on traditional approaches is desired, HR zones as provided by the American College of Sports Medicine [17] may be determined. For details see Table 1 and Sect. 6</p> <p>If the purpose is to assess inter- or intraindividual changes in activity profiles, we recommend displaying individual activity profiles by the time spent in 10% clusters of the HRR (see Sect. 6), expressed as a percentage (i.e. from 100%). Calculate the activity score based on the mean of HRR counts per 10% cluster (see Sect. 6 for details)</p> <p>NB: Some devices may adjust the sampling rate on the basis of the activity performed (e.g. providing data every 5 s up to several minutes). It is advised to choose a device that uses a constant high sampling rate. However, in any case, information on the sampling rate has to be obtained prior to selecting a device. Data points that are within the reported sampling rate range (according to the manufacturer guidelines) should be imputed by using a last value carried forward method on the basis of the highest sampling rate. Missing data (i.e. defined as deviations from a reported sampling rate range) should be excluded from the analysis</p> <p>Furthermore, often night-time is used to charge the batteries of wearables but this period may include critical data (e.g. for the assessment of nocturnal HR). Depending on the aim of the study, participants should be instructed on preferred time slots for battery charging to not miss important data</p>	<p>Report software/application version for data processing (including download and analysis)</p> <p>Report actual time of data analysed (mean number of valid days)</p> <p>If HR-zones are calculated, report time spent in each HR-zone (and indicate the intensity cut-offs used) or in each 10% cluster</p> <p>If desired, additionally report the activity score as outlined in the main manuscript (see Sect. 6)</p>

HR, heart rate

HR_{max} assessment:

1. HR_{max} was calculated by converting the beat-by-beat HR data provided by the Firstbeat Bodyguard® 2 to beats per minute via the following formula:

$$\text{HR [bpm]} = 60 \div \text{RR Interval [sec]}.$$

2. The highest 30-s rolling average was considered HR_{max}. HR_{max} was defined as 183 bpm (Fig. 2).

Nocturnal HR assessment:

1. Nocturnal HR was assessed by using 2-h rolling averages to detect the lowest mean HR during the reported sleeping window (Fig. 2). Nocturnal HR was thus determined as 55 bpm.

Activity profiling:

1. Missing data were determined by calculating the difference between the expected number of data points for a complete 24-h measurement on the basis of the sampling interval of 1 min compared with the actual number of data points available.
2. The measurement was considered a valid day because more than 16 h of data were available (Fig. 2).
3. HRR was defined as 128 bpm. The physical activity distribution was displayed on the basis of the recommendations provided by ACSM (Fig. 3) as well as in 10% clusters (Fig. 4). The time spent in each zone or cluster (expressed in minutes or % of the total time) should be reported.
4. If desired for easier comparison or quantification of left-right shifts in the activity profile (based on the research question), an activity score may be calculated as follows:

$$\text{Activity Score} = \frac{\sum_{i=1}^n (\text{time in cluster}_i \times i)}{\text{total time}}$$

n = total number of clusters; i = number of the specific cluster (e.g. 1–10)

5. For a user-friendly illustration, the activity profile may be smoothed and presented as the physical activity profile (line), and different days could be presented with different colours for instance (Fig. 5).

7 Discussion and Future Perspectives

Traditionally, free-living physical activity is assessed by accelerometer data, for which an abundance of methodological papers is available [9–12, 113, 114]. However,

while accelerometer data reflect an external load, HR is considered an internal (often referred to as a relative) measure of intensity. Therefore, HR data have typically been used to evaluate the intensity distribution of single exercise sessions in an attempt to optimize exercise prescription in recreational and elite sport settings. However, since the purpose for tracking individual training sessions differs from that of assessing free-living physical activity, existing theories and recommendations on the use of HR for exercise prescription may also not be appropriate for the quantification of free-living activity. Therefore, we feel the guidelines provided within this paper are timely and primarily aimed at facilitating the comparison of free-living activity data derived from continuous HR measures.

Nevertheless, these guidelines may also be utilized by manufacturers. In fact, our grey-literature search on user manuals and promotional materials of leading companies clearly revealed that the utilization of free-living data for the profiling of physical activity is not yet common and uniform (Table 3). While it is likely that such data may be incorporated into algorithms that provide other measures, such as energy expenditure, only few manufacturers actually visualize activity patterns on the basis of HR. Therefore, we also aim to encourage the use of continuous HR measures as an alternative to sole accelerometer-based data.

There are a few limitations that need to be addressed. First, any of the suggested approaches require access to continuous HR data (as opposed to summary data), which is currently granted only by a minority of manufacturers. Furthermore, the sampling rate may vary considerably between manufacturers and wearable models. In an attempt to save battery life, typically the sampling rate is reduced or automatically adjusted on the basis of the type of activity, affecting data quality. In this context, it is important to bear in mind that HR kinetics typically react slower to activity changes [115, 116] compared with the immediate response of acceleration measures and, therefore, naturally lower sampling rates may be sufficient for HR data. However, an appropriate resolution is needed to minimize the shift to lower HR. Furthermore, for a statistically sound analysis an individual estimation of the number of bins would need to be carried out. However, this would again lead to heterogeneous reporting and, therefore, hinder the comparisons between HR data derived from wearables with different sampling rates and would not help to overcome differences in sampling frequency. In line with this, data processing (including the treatment of missing data) requires special attention but was beyond the scope of this paper.

Collectively, manufacturers are encouraged to further improve free-living HR data quality. This also includes transparent reporting on actual sampling rates.

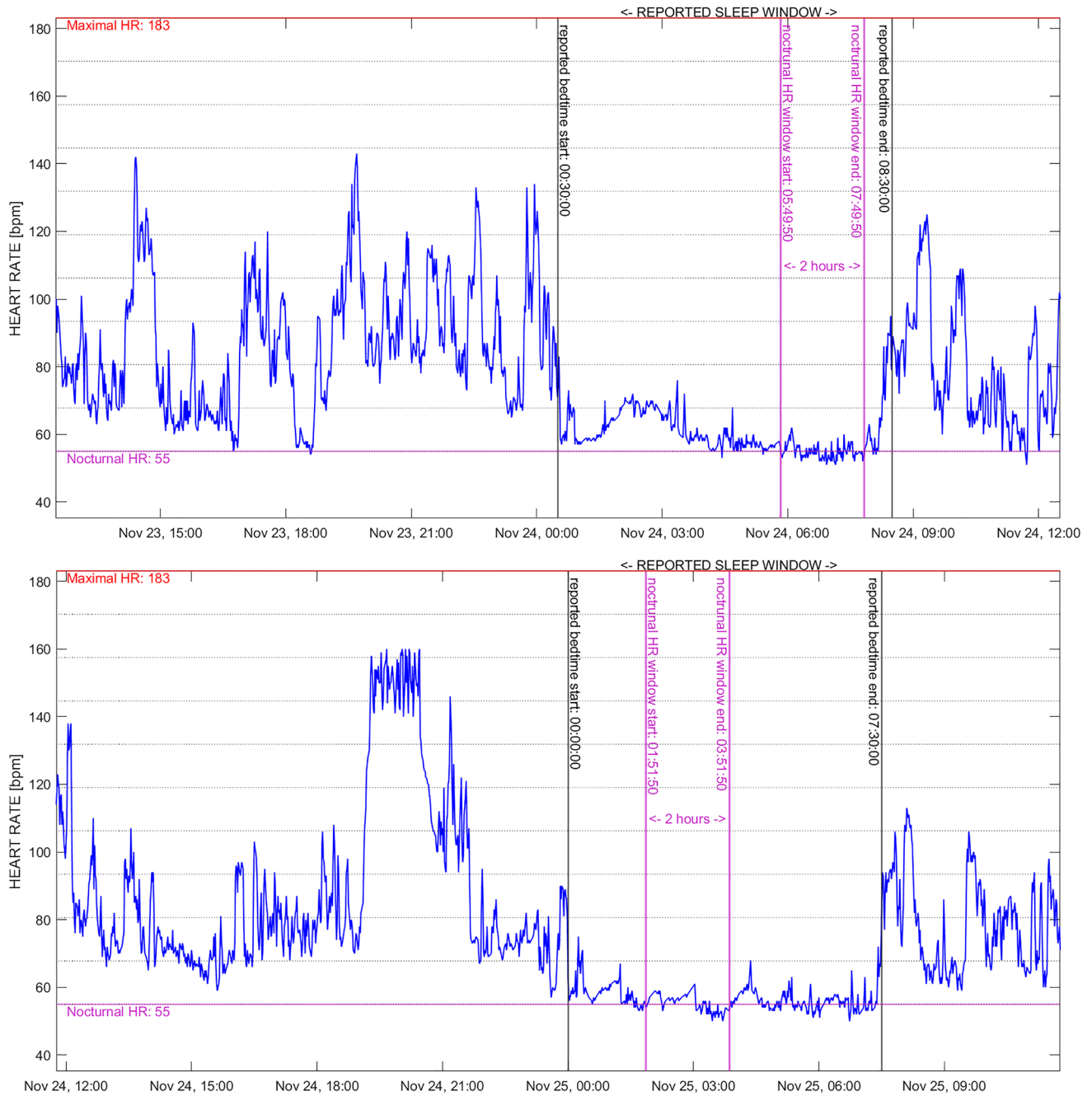


Fig. 2 Continuous free-living heart rate (HR) over 24 h illustrating a less (upper plot) and a more (lower plot) active day

Interestingly, attempts are in place with third-party solutions that allow the set up of customized sampling rates (e.g. the fitrockr software that collaborates with Garmin [117]), allowing access to the continuous HR data as opposed to summary data typically displayed on the user surface of the devices. However, while it is expected that this will provide new horizons for the consistent use of

free-living HR data to quantify individual activity patterns, this will also require further advances in analytical approaches. Examples for assessing the accurate distribution of data collected with very high sampling rates are provided by multivariate/functional data analysis for accelerometer data [12] but the application to HR data requires further studies.

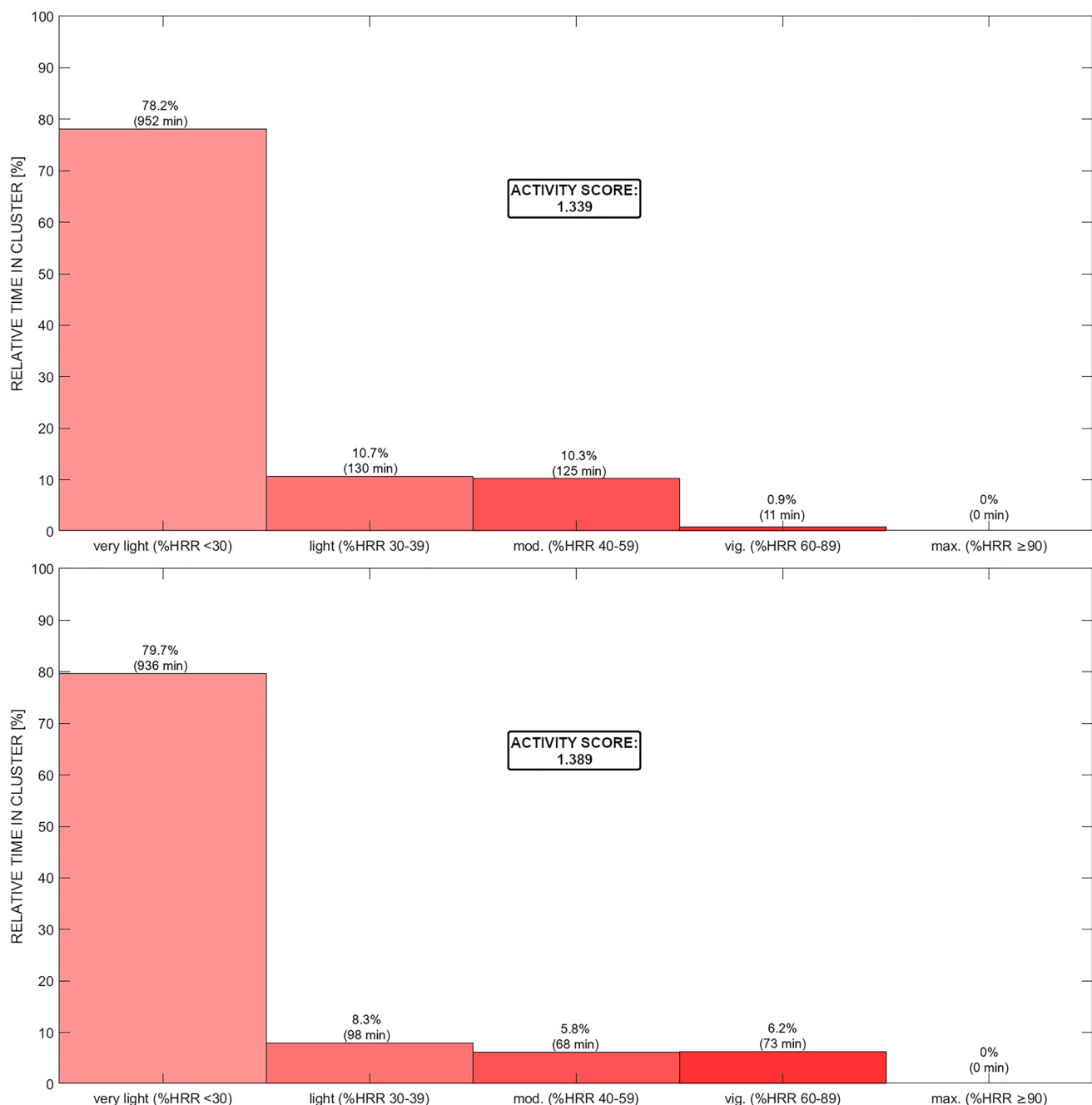


Fig. 3 Free-living heart rate (HR) over 24 h illustrating a less (upper plot) and a more (lower plot) active day on the basis of HR reserve (HRR) and clustered according to the common guidelines provided

by the American College of Sports Medicine. NB: numbers above the bins indicate the relative and absolute time spent in each zone

8 Conclusion and Practical Applications

Combining the information retrieved through a scoping review, grey literature search and the a priori knowledge of the INTERLIVE®-network, we provided detailed recommendations for the analysis of free-living HR data to derive individual physical activity profiles obtained by wearables. In addition, this article provides recommendations on how

to best measure or predict maximal HR and basal/nocturnal/resting HR.

Since it is well-known that higher levels of physical activity at any intensity and reduced sedentary time are associated with substantial reductions in health risks and premature mortality [108, 118], it is of utmost importance to advance on feasible methods to quantify physical activity and reinforce adherence to physical activity and clinical

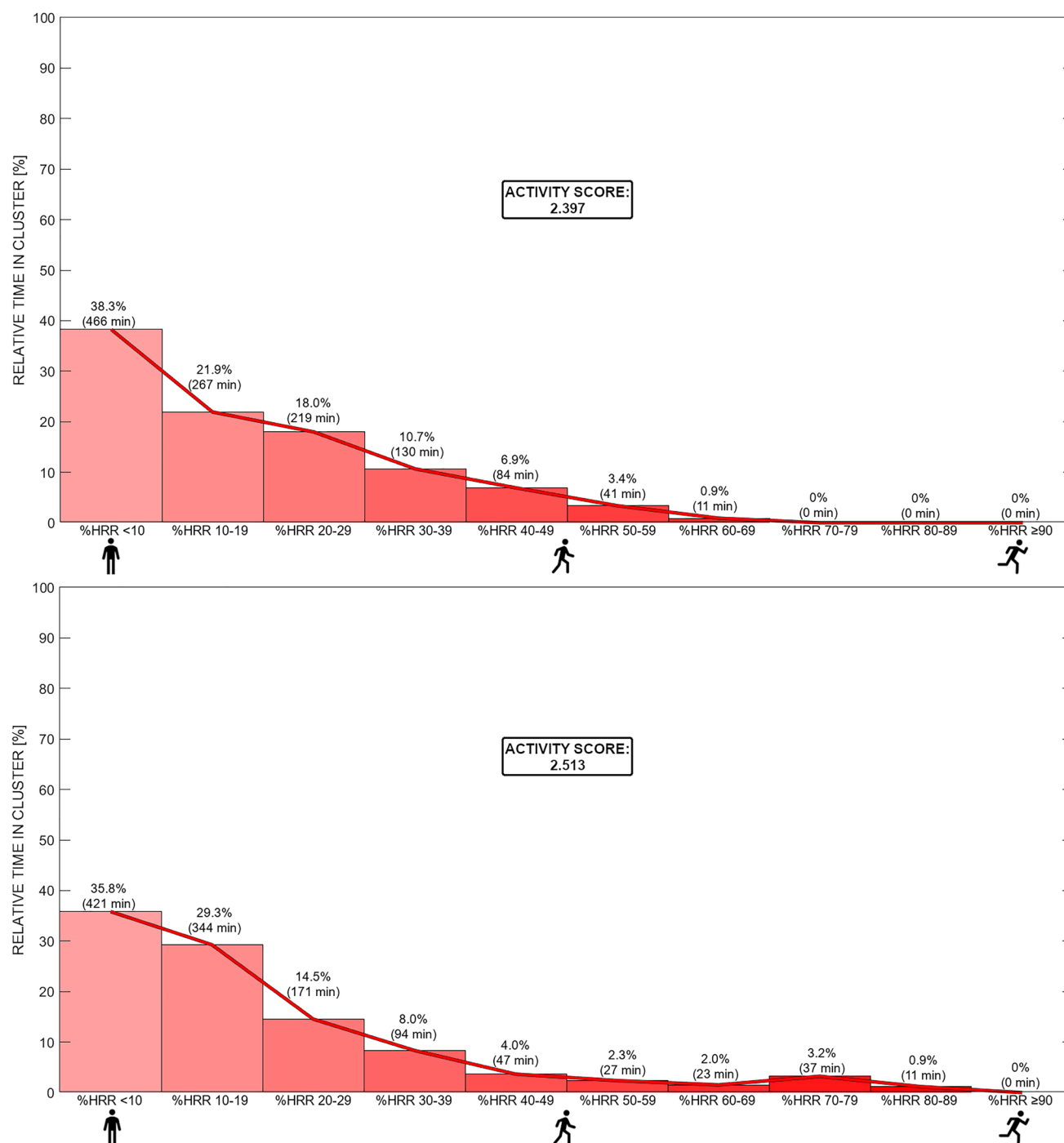


Fig. 4 Free-living heart rate (HR) over 24 h illustrating a less (upper plot) and a more (lower plot) active day, displayed in 10% clusters based on HR reserve (HR) (Cluster 1: 0–10% HRR, [...], Cluster 10:

91–100% HRR). The activity score was calculated to quantify overall physical activity. NB: numbers above the bins indicate the relative and absolute time spent in each cluster

guidelines. In this context, objective data provided by wearable technology open up new opportunities. Especially HR as an internal measure of activity intensity may overcome some of the limitations related to acceleration-derived physical activity, and may therefore be used as an alternative or complementary method. In this article, we provide

a harmonized analytical approach for evaluating physical activity patterns in clinical practice as well as fitness and health settings by free-living HR recordings. Specifically, these recommendations may be applied to various research questions, including randomized controlled trials assessing changes in physical activity patterns, cross-sectional analysis

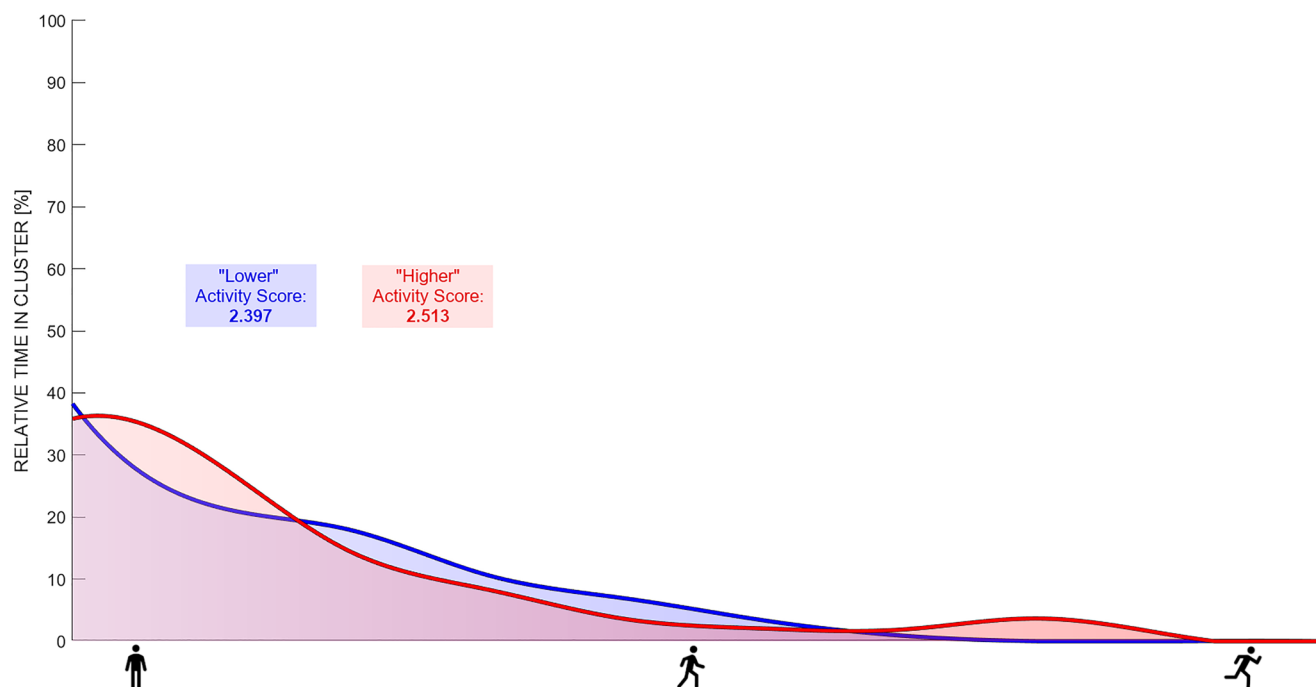


Fig. 5 Free-living heart rate (HR) over 24 h illustrating a less (blue) and more (red) active day, displayed as a smoothed curve for a better comparability with the plots shown in Fig. 4. The activity score was calculated to quantify overall physical activity

comparing physical activity of distinct populations as well as longitudinal observational studies. While these guidelines are directly useful for researchers and manufacturers, end-users may also benefit, being better informed and empowered to understand and use the HR-information that can be derived from their wearables.

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Declarations

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Data availability All underlying data can be found in the online supplementary data.

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References

1. De la Fuente Robles YM, Ricoy Cano AJ, Albín Rodríguez AP, López Ruiz JL, Espinilla EM. Past, present and future of research on wearable technologies for healthcare: a bibliometric analysis using scopus. *Sensors*. 2022;22:8599. <https://doi.org/10.3390/s22228599>.

2. Kuratomi D, Shin C, Duffy VG. Systematic literature review on the advances of wearable technologies. 2023;78–95. https://doi.org/10.1007/978-3-031-48047-8_5.
3. Fortino G, Gravina R, Galzarano S. Wearable computing: from modeling to implementation of wearable systems based on body sensor networks. Wiley-IEEE Press; 2018.
4. Mühlen JM, Stang J, Lykke Skovgaard E, Judice PB, Molina-Garcia P, Johnston W, et al. Recommendations for determining the validity of consumer wearable heart rate devices: expert statement and checklist of the INTERLIVE Network. *Br J Sports Med*. 2021. <https://doi.org/10.1136/bjsports-2020-103148>.
5. Johnston W, Judice PB, Molina García P, Mühlen JM, Lykke Skovgaard E, Stang J, et al. Recommendations for determining the validity of consumer wearable and smartphone step count: expert statement and checklist of the INTERLIVE network. *Br J Sports Med*. 2021;55:780–93. <https://doi.org/10.1136/BJSPOR-RTS-2020-103147>.
6. Molina-Garcia P, Notbohm HL, Schumann M, Argent R, Hetherington-Rauth M, Stang J, et al. Validity of estimating the maximal oxygen consumption by consumer wearables: a systematic review with meta-analysis and expert statement of the INTERLIVE Network. *Sports Med*. 2022;52:1577–97. <https://doi.org/10.1007/S40279-021-01639-Y>.
7. Argent R, Hetherington-Rauth M, Stang J, Tarp J, Ortega FB, Molina-Garcia P, et al. Recommendations for determining the validity of consumer wearables and smartphones for the estimation of energy expenditure: expert statement and checklist of the INTERLIVE network. *Sports Med*. 2022;52:1817–32. <https://doi.org/10.1007/S40279-022-01665-4>.
8. McDonough DJ, Su X, Gao Z. Health wearable devices for weight and BMI reduction in individuals with overweight/obesity and chronic comorbidities: systematic review and network meta-analysis. *Br J Sports Med*. 2021;55:917–25. <https://doi.org/10.1136/BJSPORTS-2020-103594>.
9. Gao Z, Liu W, McDonough DJ, Zeng N, Lee JE. The dilemma of analyzing physical activity and sedentary behavior with wrist accelerometer data: challenges and opportunities. *J Clin Med*. 2021;10:5951. <https://doi.org/10.3390/JCM10245951>.
10. Clevenger KA, Montoye AHK, Van Camp CA, Strath SJ, Pfeiffer KA. Methods for estimating physical activity and energy expenditure using raw accelerometry data or novel analytical approaches: a repository, framework, and reporting guidelines. *Physiol Meas*. 2022. <https://doi.org/10.1088/1361-6579/ac89c9>.
11. Migueles JH, Cadenas-Sanchez C, Ekelund U, Delisle Nyström C, Mora-Gonzalez J, Löf M, et al. Accelerometer data collection and processing criteria to assess physical activity and other outcomes: a systematic review and practical considerations. *Sports Med*. 2017. <https://doi.org/10.1007/s40279-017-0716-0>.
12. Migueles JH, Aadland E, Andersen LB, Brønd JC, Chastin SF, Hansen BH, et al. GRANADA consensus on analytical approaches to assess associations with accelerometer-determined physical behaviours (physical activity, sedentary behaviour and sleep) in epidemiological studies. *Br J Sports Med*. 2022;56:376–84. <https://doi.org/10.1136/BJSPORTS-2020-103604>.
13. Pulsford RM, Brocklebank L, Fenton SAM, Bakker E, Mielke GI, Tsai L-T, et al. The impact of selected methodological factors on data collection outcomes in observational studies of device-measured physical behaviour in adults: a systematic review. *Int J Behav Nutr Phys Act*. 2023;20:26. <https://doi.org/10.1186/s12966-022-01388-9>.
14. Seiler KS, Kjerland GØ. Quantifying training intensity distribution in elite endurance athletes: is there evidence for an “optimal” distribution? *Scand J Med Sci Sports*. 2006;16:49–56. <https://doi.org/10.1111/J.1600-0838.2004.00418.X>.
15. Jamnick NA, Pettitt RW, Granata C, Pyne DB, Bishop DJ. An examination and critique of current methods to determine exercise intensity. *Sports Med*. 2020;50:1729–56. <https://doi.org/10.1007/S40279-020-01322-8>.
16. Davis JA, Convertino VA. A comparison of heart rate methods for predicting endurance training intensity. *Med Sci Sports*. 1975;7:295–8. <https://doi.org/10.1249/00005768-19750740-00010>.
17. American College of Sports Medicine. ACSM’s guidelines for exercise testing and prescription. Philadelphia: Lippincott Williams & Wilkins; 2021. p. 11.
18. Norton K, Norton L, Sadgrove D. Position statement on physical activity and exercise intensity terminology. *J Sci Med Sport*. 2010;13:496–502. <https://doi.org/10.1016/j.jsams.2009.09.008>.
19. Laricchia F. Global smartwatch market share 2020–2022. Counterpoint Technology Market Research. 2023.
20. Fernhall B, Mccubbin JA, Pitetti KH, Rintala P, Rimmer JH, Lynn Millar A, et al. Prediction of maximal heart rate in individuals with mental retardation. *Med Sci Sports Exerc*. 2001;33:1655–60. <https://doi.org/10.1097/00005768-200110000-00007>.
21. Lester M, Sheffield LT, Trammell P, Reeves TJ. The effect of age and athletic training on the maximal heart rate during muscular exercise. *Am Heart J*. 1968;76:370–6. [https://doi.org/10.1016/0002-8703\(68\)90233-0](https://doi.org/10.1016/0002-8703(68)90233-0).
22. Miller WC, Wallace JP, Eggert KE. Predicting max HR and the HR-VO₂ relationship for exercise prescription in obesity. *Med Sci Sports Exerc*. 1993;25:1077–81. <https://doi.org/10.1249/00005768-199309000-00017>.
23. Whyte GP, George K, Shave R, Middleton N, Nevill AM. Training induced changes in maximum heart rate. *Int J Sports Med*. 2008;29:129–33. <https://doi.org/10.1055/S-2007-965783/ID/18>.
24. Logan N, Reilly JJ, Grant S, Paton JY. Resting heart rate definition and its effect on apparent levels of physical activity in young children. *Med Sci Sports Exerc*. 2000;32:162–6. <https://doi.org/10.1097/00005768-200001000-00024>.
25. Strzelczyk TA, Quigg RJ, Pfeifer PB, Parker MA, Greenland P. Accuracy of estimating exercise prescription intensity in patients with left ventricular systolic dysfunction. *J Cardiopulm Rehabil*. 2001;21:158–63. <https://doi.org/10.1097/00008483-200105000-00007>.
26. Goldberg L, Elliot DL, Kuehl KS. Assessment of exercise intensity formulas by use of ventilatory threshold. *Chest*. 1988;94:95–8. <https://doi.org/10.1378/CHEST.94.1.95>.
27. Meyer T, Gabriel HHW, Kindermann W. Is determination of exercise intensities as percentages of VO₂max or HRmax adequate? *Med Sci Sports Exerc*. 1999;31:1342–5. <https://doi.org/10.1097/00005768-199909000-00017>.
28. Weltman A, Snead D, Seip R, Schurrer R, Weltman J, Rutt R, et al. Percentages of maximal heart rate, heart rate reserve and VO₂max for determining endurance training intensity in male runners. *Int J Sports Med*. 1990;11:218–22. <https://doi.org/10.1055/S-2007-1024795>.
29. Hofmann P, Von Duvillard SP, Seibert FJ, Pokan R, Wonisch M, Lemura LM, et al. %HRmax target heart rate is dependent on heart rate performance curve deflection. *Med Sci Sports Exerc*. 2001;33:1726–31. <https://doi.org/10.1097/00005768-200110000-00017>.
30. Katch V, Weltman A, Sady S, Freedson P. Validity of the relative percent concept for equating training intensity. *Eur J Appl Physiol Occup Physiol*. 1978;39:219–27. <https://doi.org/10.1007/BF00421445>.
31. Sebastian LA, Reeder S, Williams M. Determining target heart rate for exercising in a cardiac rehabilitation program: a retrospective study. *J Cardiovasc Nurs*. 2015;30:164–71. <https://doi.org/10.1097/JCN.0000000000000154>.
32. Mielke M, Housh TJ, Hendrix RC, Zuniga J, Camic CL, Schmidt RJ, et al. A test for determining critical heart rate using the

- critical power model. *J Strength Cond Res.* 2011;25:504–10. <https://doi.org/10.1519/JSC.0B013E3181B62C43>.
33. Silva P, Dos Santos E, Grishin M, Rocha JM. Validity of heart rate-based indices to measure training load and intensity in elite football players. *J Strength Cond Res.* 2018;32:2340–7. <https://doi.org/10.1519/JSC.0000000000002057>.
 34. Garber CE, Blissmer B, Deschenes MR, Franklin BA, Lamonte MJ, Lee IM, et al. Quantity and quality of exercise for developing and maintaining cardiorespiratory, musculoskeletal, and neuromotor fitness in apparently healthy adults: guidance for prescribing exercise. *Med Sci Sports Exerc.* 2011;43:1334–59. <https://doi.org/10.1249/MSS.0B013E318213FEFB>.
 35. Fox SM, Naughton JP. Physical activity and the prevention of coronary heart disease. *Prev Med.* 1972;1:92–120. [https://doi.org/10.1016/0091-7435\(72\)90079-5](https://doi.org/10.1016/0091-7435(72)90079-5).
 36. Robergs RA, Landwehr R. The surprising history of the ‘HRmax=220-age’ equation. *J Exerc Physiol.* 2002;5.
 37. Rønnestad BR, Hansen J, Ellefsen S. Block periodization of high-intensity aerobic intervals provides superior training effects in trained cyclists. *Scand J Med Sci Sports.* 2014;24:34–42. <https://doi.org/10.1111/j.1600-0838.2012.01485.X>.
 38. Sylta Ø, Tønnessen E, Seiler S. From heart-rate data to training quantification: a comparison of 3 methods of training-intensity analysis. *Int J Sports Physiol Perform.* 2014;9:100–7. <https://doi.org/10.1123/IJSP.2013-0298>.
 39. Roete AJ, Stoter IK, Lamberts RP, Elferink-Gemser MT, Otter RTA. Introducing a method to quantify the specificity of training for races in speed skating. *J Strength Cond Res.* 2022;36:1998–2004. <https://doi.org/10.1519/JSC.0000000000004271>.
 40. Kuipers H, Rietjens G, Verstappen F, Schoenmakers H, Hofman G. Effects of stage duration in incremental running tests on physiological variables. *Int J Sports Med.* 2003;24:486–91. <https://doi.org/10.1055/S-2003-42020>.
 41. Cunha FA, Midgley AW, Monteiro WD, Farinatti PTV. Influence of cardiopulmonary exercise testing protocol and resting VO₂ assessment on %HRmax, %HRR, %VO₂max and %VO₂R relationships. *Int J Sports Med.* 2010;31:319–26. <https://doi.org/10.1055/S-0030-1248283/ID/27/BIB>.
 42. Machado FA, Kravchychyn ACP, Peserico CS, da Silva DF, Mez-zaroba PV. Effect of stage duration on maximal heart rate and post-exercise blood lactate concentration during incremental treadmill tests. *J Sci Med Sport.* 2013;16:276–80. <https://doi.org/10.1016/j.jsams.2012.08.003>.
 43. Ingjer F. Factors influencing assessment of maximal heart rate. *Scand J Med Sci Sports.* 1991;1:134–40. <https://doi.org/10.1111/j.1600-0838.1991.tb00285.x>.
 44. Muscat KM, Kotrach HG, Wilkinson-Maitland CA, Schaeffer MR, Mendonca CT, Jensen D. Physiological and perceptual responses to incremental exercise testing in healthy men: effect of exercise test modality. *Appl Physiol Nutr Metab.* 2015;40:1199–209. <https://doi.org/10.1139/APNM-2015-0179>.
 45. Beltz NM, Gibson AL, Janot JM, Kravitz L, Mermier CM, Dalleck LC. Graded exercise testing protocols for the determination of VO₂max: historical perspectives, progress, and future considerations. *J Sports Med.* 2016;2016:1–12. <https://doi.org/10.1155/2016/3968393>.
 46. Fletcher GF, Ades PA, Kligfield P, Arena R, Balady GJ, Bittner VA, et al. Exercise standards for testing and training: a scientific statement from the American Heart Association. *Circulation.* 2013;128:873–934. <https://doi.org/10.1161/CIR.0B013E31829B5B44>.
 47. Faff J, Sitkowski D, Ladyga M, Klusiewicz A, Sitkowski D, Ladyga M, et al. Maximal heart rate in athletes. *Biol Sport.* 2007;24:129–42.
 48. Zinner C, Gerspitzer A, Düking P, Boone J, Schiffer T, Holmberg H, et al. The magnitude and time-course of physiological responses to 9-weeks of incremental ramp testing. *Scand J Med Sci Sports.* 2023. <https://doi.org/10.1111/SMS.14347>.
 49. Fairbairn MS, Blackie SP, McElvaney NG, Wiggs BR, Pare PD, Pardy RL. Prediction of heart rate and oxygen uptake during incremental and maximal exercise in healthy adults. *Chest.* 1994;105:1365–9. <https://doi.org/10.1378/chest.105.5.1365>.
 50. Park JH, Jung HC, Jung YS, Song JK, Lee JM. Re-visiting maximal heart rate prediction using cross-validation in population aged 7–55 years. *Int J Environ Res Public Health.* 2022. <https://doi.org/10.3390/ijerph19148509>.
 51. Shargal E, Kislev-Cohen R, Zigel L, Epstein S, Pilz-Burstein R, Tenenbaum G. Age-related maximal heart rate: examination and refinement of prediction equations. *J Sports Med Phys Fitness.* 2015;55:1207–18.
 52. Londeree BR, Moeschberger ML. Effect of age and other factors on maximal heart rate. *Res Q Exerc Sport.* 1982;53:297–304. <https://doi.org/10.1080/02701367.1982.10605252>.
 53. Nikolaidis PT, Rosemann T, Knechtle B. Age-predicted maximal heart rate in recreational marathon runners: a cross-sectional study on Fox’s and Tanaka’s equations. *Front Physiol.* 2018. <https://doi.org/10.3389/fphys.2018.00226>.
 54. Sydó N, Abdelmoneim SS, Mulvagh SL, Merkely B, Gulati M, Allison TG. Relationship between exercise heart rate and age in men vs women. *Mayo Clin Proc.* 2014;89:1664–72. <https://doi.org/10.1016/j.mayocp.2014.08.018>.
 55. Gelbart M, Ziv-Baran T, Williams CA, Yarom Y, Dubnov-Raz G. Prediction of maximal heart rate in children and adolescents. *Clin J Sport Med.* 2017;27:139–44. <https://doi.org/10.1097/JSM.0000000000000315>.
 56. Mahon AD, Marjerrison AD, Woodruff ME, Hanna LE. Evaluating the prediction of maximal heart rate in children and adolescents. *Res Q Exerc Sport.* 2010;81:466–71. <https://doi.org/10.1080/02701367.2010.10599707>.
 57. Tao K, Li J, Li J, Shan W, Yan H, Lu Y. Estimation of heart rate using regression models and artificial neural network in middle-aged adults. *Front Physiol.* 2021. <https://doi.org/10.3389/fphys.2021.742754>.
 58. Whaley MH, Kaminsky LA, Dwyer GB, Getchell LH, Norton JA. Predictors of over- and underachievement of age-predicted maximal heart rate. *Med Sci Sports Exerc.* 1992;24:1173–9.
 59. Magri D, Piepoli M, Gallo G, Corrà U, Metra M, Paolillo S, et al. Old and new equations for maximal heart rate prediction in patients with heart failure and reduced ejection fraction on beta-blockers treatment: results from the MECKI score data set. *Eur J Prev Cardiol.* 2022;29:1680–8. <https://doi.org/10.1093/EURJPC/ZWAC099>.
 60. Keteyian SJ, Kitzman D, Zannad F, Landzberg J, Arnold JM, Brubaker P, et al. Predicting maximal HR in heart failure patients on β -blockade therapy. *Med Sci Sports Exerc.* 2012;44:371–6. <https://doi.org/10.1249/MSS.0b013e318234316f>.
 61. Johnson JH, Prins A. Prediction of maximal heart rate during a submaximal work test. *J Sports Med Phys Fitness.* 1991;31:44–7.
 62. Karavirta L, Tulppo MP, Nyman K, Laaksonen DE, Pullinen T, Laukkanen RT, et al. Estimation of maximal heart rate using the relationship between heart rate variability and exercise intensity in 40–67 years old men. *Eur J Appl Physiol.* 2008;103:25–32. <https://doi.org/10.1007/s00421-007-0667-5>.
 63. Matabuena M, Vidal JC, Hayes PR, Saavedra-Garcia M, Trillo FH. Application of functional data analysis for the prediction of maximum heart rate. *IEEE Access.* 2019;7:121841–52. <https://doi.org/10.1109/ACCESS.2019.2938466>.
 64. Mazzoleni MJ, Battaglini CL, Martin KJ, Coffman EM, Ekaidat JA, Wood WA, et al. A dynamical systems approach for the submaximal prediction of maximum heart rate and maximal oxygen uptake. *Sports Eng.* 2018;21:31–41. <https://doi.org/10.1007/s12283-017-0242-1>.



65. Casillas JM, Joussain C, Gremeaux V, Hannequin A, Rapin A, Laurent Y, et al. A study of the 200-metre fast walk test as a possible new assessment tool to predict maximal heart rate and define target heart rate for exercise training of coronary heart disease patients. *Clin Rehabil*. 2015;29:175–83. <https://doi.org/10.1177/0269215514540922>.
66. Ricard RM, Leger L, Massicotte D. Validity of the “220-Age Formula” to predict maximal heart rate. *Med Sci Sports Exerc*. 1990;22:S96. <https://doi.org/10.1249/00005768-199004000-00574>.
67. Schiller BC, Casas YG, Desouza CA, Seals DR. Maximal aerobic capacity across age in healthy Hispanic and Caucasian women. *J Appl Physiol*. 2001;91:1048–54. <https://doi.org/10.1152/jappl.2001.91.3.1048>.
68. Graettinger WF, Smith DHG, Neutel JM, Myers J, Froelicher VF, Weber MA. Relationship of left ventricular structure to maximal heart rate during exercise. *Chest*. 1995;107:341–5. <https://doi.org/10.1378/chest.107.2.341>.
69. Kirk Hammond H, Lisa Kelly T, Froelicher V. Radionuclide imaging correlates of heart rate impairment during maximal exercise testing. *J Am Coll Cardiol*. 1983;2:826–33. [https://doi.org/10.1016/S0735-1097\(83\)80228-9](https://doi.org/10.1016/S0735-1097(83)80228-9).
70. Fernandes Silva MM, Bacal F, Roque JM, Teixeira-Neto IS, Carvas Junior N, Bocchi EA, et al. Age-related maximum heart rate among ischemic and nonischemic heart failure patients receiving β -blockade therapy. *J Card Fail*. 2012;18:831–6. <https://doi.org/10.1016/j.cardfail.2012.10.007>.
71. Brawner CA, Ehrman JK, Schairer JR, Cao JJ, Keteyian SJ. Predicting maximum heart rate among patients with coronary heart disease receiving beta-adrenergic blockade therapy. *Am Heart J*. 2004;148:910–4. <https://doi.org/10.1016/j.ahj.2004.04.035>.
72. Verschuren O, Maltais DB, Takken T. The 220-age equation does not predict maximum heart rate in children and adolescents. *Dev Med Child Neurol*. 2011;53:861–4. <https://doi.org/10.1111/j.1469-8749.2011.03989.x>.
73. Godlasky E, Hoffman T, Weber-Peters S, Bradford R, Miller N, Kunselman AR, et al. Effects of β -blockers on maximal heart rate prediction equations in a cardiac population. *J Cardiopulm Rehabil Prev*. 2018;38:111–7. <https://doi.org/10.1097/HCR.0000000000000328>.
74. Cicone ZS, Sinelnikov OA, Esco MR. Age-predicted maximal heart rate equations are inaccurate for use in youth male soccer players. *Pediatr Exerc Sci*. 2018;30:495–9. <https://doi.org/10.1123/pes.2017-0281>.
75. Cicone ZS, Holmes CJ, Fedewa MV, MacDonald HV, Esco MR. Age-based prediction of maximal heart rate in children and adolescents: a systematic review and meta-analysis. *Res Q Exerc Sport*. 2019;90:417–28. <https://doi.org/10.1080/02701367.2019.1615605>.
76. Hill M, Talbot C, Price M. Predicted maximal heart rate for upper body exercise testing. *Clin Physiol Funct Imaging*. 2016;36:155–8. <https://doi.org/10.1111/cpf.12201>.
77. Machado FA, Denadai BS. Validade das equações preditivas da frequência cardíaca máxima para crianças e adolescentes. *Arq Bras Cardiol*. 2011;97:136–40. <https://doi.org/10.1590/S0066-782X2011005000078>.
78. Nikolaidis PT. Maximal heart rate in soccer players: measured versus age-predicted. *Biomed J*. 2015;38:84–9. <https://doi.org/10.4103/2319-4170.131397>.
79. Nikolaidis PT, Padulo J, Chtourou H, Torres-Luque G, Afonso J, Heller J. Estimating maximal heart rate with the ‘220-age’ formula in adolescent female volleyball players: a preliminary study. *Hum Mov*. 2018;15:166–70. <https://doi.org/10.1515/HUMO-2015-0007>.
80. Nikolaidis PT. Age-predicted vs. measured maximal heart rate in young team sport athletes. *Niger Med J*. 2014;55:314. <https://doi.org/10.4103/0300-1652.137192>.
81. Johansen CD, Olsen RH, Pedersen LR, Kumarathurai P, Mouridsen MR, Binici Z, et al. Resting, night-time, and 24 h heart rate as markers of cardiovascular risk in middle-aged and elderly men and women with no apparent heart disease. *Eur Heart J*. 2013;34:1732–9. <https://doi.org/10.1093/EURHEARTJ/EHS449>.
82. Mitchell HH. Comparative nutrition of man and domestic animals. Academic Press; 1962.
83. Henry C. Basal metabolic rate studies in humans: measurement and development of new equations. *Public Health Nutr*. 2005;8:1133–52. <https://doi.org/10.1079/PHN2005801>.
84. Speed C, Arneil T, Harle R, Wilson A, Karthikesalingam A, McConnell M, et al. Measure by measure: resting heart rate across the 24-hour cycle. *PLOS Dig Health*. 2023. <https://doi.org/10.1371/journal.pdig.0000236>.
85. Fox K, Borer JS, Camm AJ, Danchin N, Ferrari R, Lopez Sendon JL, et al. Resting heart rate in cardiovascular disease. *J Am Coll Cardiol*. 2007;50:823–30. <https://doi.org/10.1016/J.JACC.2007.04.079>.
86. Greenland P, Daviglus ML, Dyer AR, Liu K, Huang CF, Goldberger JJ, et al. Resting heart rate is a risk factor for cardiovascular and noncardiovascular mortality: the Chicago Heart Association Detection Project In Industry. *Am J Epidemiol*. 1999;149:853–62. <https://doi.org/10.1093/OXFORDJOURNALS.AJE.A009901>.
87. Hsia J, Larson JC, Ockene JK, Sarto GE, Allison MA, Hendrix SL, et al. Resting heart rate as a low tech predictor of coronary events in women: prospective cohort study. *BMJ*. 2009;338:577–9. <https://doi.org/10.1136/BMJ.B219>.
88. Melanson EL, Freedson PS. The effect of endurance training on resting heart rate variability in sedentary adult males. *Eur J Appl Physiol*. 2001;85:442–9. <https://doi.org/10.1007/S004210100479/METRICS>.
89. Jensen MT, Marott JL, Allin KH, Nordestgaard BG, Jensen GB. Resting heart rate is associated with cardiovascular and all-cause mortality after adjusting for inflammatory markers: the Copenhagen City Heart Study. *Eur J Prev Cardiol*. 2012;19:102–8. <https://doi.org/10.1177/1741826710394274>.
90. Esco MR, Olson MS, Williford HN, Blessing DL, Shannon D, Grandjean P. The relationship between resting heart rate variability and heart rate recovery. *Clin Auton Res*. 2010;20:33–8. <https://doi.org/10.1007/S10286-009-0033-2/FIGURES/3>.
91. Carnethon MR, Yan L, Greenland P, Garside DB, Dyer AR, Metzger B, et al. Resting heart rate in middle age and diabetes development in older age. *Diabetes Care*. 2008;31:335–9. <https://doi.org/10.2337/DC07-0874>.
92. Cajochen C, Pischke J, Aeschbach D, Borbély AA. Heart rate dynamics during human sleep. *Physiol Behav*. 1994;55:769–74. [https://doi.org/10.1016/0031-9384\(94\)90058-2](https://doi.org/10.1016/0031-9384(94)90058-2).
93. Patel AK, Reddy V, Shumway KR, Araujo JF. Physiology, sleep stages. StatPearls: StatPearls Publishing; 2023.
94. Nummela A, Hynynen E, Kaikkonen P, Rusko H. High-intensity endurance training increases nocturnal heart rate variability in sedentary participants. *Biol Sport*. 2016;33:7–13. <https://doi.org/10.5604/20831862.1180171>.
95. Myllymäki T, Rusko H, Syväoja H, Juuti T, Kinnunen ML, Kyröläinen H, et al. Effects of exercise intensity and duration on nocturnal heart rate variability and sleep quality. *Eur J Appl Physiol*. 2012;112:801–9. <https://doi.org/10.1007/S00421-011-2034-9>.
96. Myllymäki T, Kyröläinen H, Savolainen K, Hokka L, Jakonen R, Juuti T, et al. Effects of vigorous late-night exercise on sleep quality and cardiac autonomic activity. *J Sleep Res*.

- 2011;20:146–53. <https://doi.org/10.1111/J.1365-2869.2010.00874.X>.
97. Hautala A, Tulppo MP, Mäkilä TH, Laukkanen R, Nissilä S, Huikuri HV. Changes in cardiac autonomic regulation after prolonged maximal exercise. *Clin Physiol*. 2001;21:238–45. <https://doi.org/10.1046/J.1365-2281.2001.00309.X>.
98. Pichot V, Roche F, Gaspoz JM, Enjolras F, Antoniadis A, Minini P, et al. Relation between heart rate variability and training load in middle-distance runners. *Med Sci Sports Exerc*. 2000;32:1729–36. <https://doi.org/10.1097/00005768-200010000-00011>.
99. Hynynen E, Vesterinen V, Rusko H, Nummela A. Effects of moderate and heavy endurance exercise on nocturnal HRV. *Int J Sports Med*. 2010;31:428–32. <https://doi.org/10.1055/S-0030-1249625/ID/31>.
100. Brandenberger G, Buchheit M, Ehrhart J, Simon C, Piquard F. Is slow wave sleep an appropriate recording condition for heart rate variability analysis? *Auton Neurosci*. 2005;121:81–6. <https://doi.org/10.1016/J.AUTNEU.2005.06.002>.
101. Gaskill SE, Ruby BC, Walker AJ, Sanchez OA, Serfass RC, Leon AS. Validity and reliability of combining three methods to determine ventilatory threshold. *Med Sci Sports Exerc*. 2001;33:1841–8. <https://doi.org/10.1097/00005768-200110000-00007>.
102. Binder RK, Wonisch M, Corra U, Cohen-Solal A, Vanhees L, Saner H, et al. Methodological approach to the first and second lactate threshold in incremental cardiopulmonary exercise testing. *Eur J Cardiovasc Prev Rehabil*. 2008;15:726–34. <https://doi.org/10.1097/HJR.0b013e328304fed4>.
103. Iannetta D, Inglis EC, Mattu AT, Fontana FY, Pogliaghi S, Keir DA, et al. A critical evaluation of current methods for exercise prescription in women and men. *Med Sci Sports Exerc*. 2020;52:466–73. <https://doi.org/10.1249/MSS.00000000000002147>.
104. Reimers AK, Knapp G, Reimers CD. Effects of exercise on the resting heart rate: a systematic review and meta-analysis of interventional studies. *J Clin Med*. 2018. <https://doi.org/10.3390/JCM7120503>.
105. Lounana J, Campion F, Noakes TD, Medelli J. Relationship between %HRmax, %HR reserve, %VO2max, and %VO2 reserve in elite cyclists. *Med Sci Sports Exerc*. 2007;39:350–7. <https://doi.org/10.1249/01.MSS.0000246996.63976.5F>.
106. Swain DP, Leutholtz BC. Heart rate reserve is equivalent to %VO2 reserve, not to %VO2max. *Med Sci Sports Exerc*. 1997;29:410–4. <https://doi.org/10.1097/00005768-199703000-00018>.
107. Swain DP, Franklin BA. VO(2) reserve and the minimal intensity for improving cardiorespiratory fitness. *Med Sci Sports Exerc*. 2002;34:152–7. <https://doi.org/10.1097/00005768-200201000-00023>.
108. Bull FC, Al-Ansari SS, Biddle S, Borodulin K, Buman MP, Cardon G, et al. World Health Organization 2020 guidelines on physical activity and sedentary behaviour. *Br J Sports Med*. 2020;54:1451–62. <https://doi.org/10.1136/BJSPOR-2020-102955>.
109. Foster C, Florhaug JA, Franklin J, Gottschall L, Hrovatin LA, Parker S, et al. A new approach to monitoring exercise training. *J Strength Cond Res*. 2001;15:109–15.
110. Williams N. The Borg Rating of Perceived Exertion (RPE) scale. *Occup Med*. 2017;67:404–5. <https://doi.org/10.1093/OCCMED/KQX063>.
111. Chen MJ, Fan X, Moe ST. Criterion-related validity of the Borg ratings of perceived exertion scale in healthy individuals: a meta-analysis. *J Sports Sci*. 2002;20:873–99. <https://doi.org/10.1080/026404102320761787>.
112. Tanaka H, Monahan KD, Seals DR. Age-predicted maximal heart rate revisited. *J Am Coll Cardiol*. 2001;37:153–6. [https://doi.org/10.1016/S0735-1097\(00\)01054-8](https://doi.org/10.1016/S0735-1097(00)01054-8).
113. Pedišić Ž, Bauman A. Accelerometer-based measures in physical activity surveillance: current practices and issues. *Br J Sports Med*. 2015;49:219–23. <https://doi.org/10.1136/BJSPOR-2013-093407>.
114. Ojiambo R, Cuthill R, Budd H, Konstabel K, Casajús JA, González-Agüero A, et al. Impact of methodological decisions on accelerometer outcome variables in young children. *Int J Obes*. 2011;35:S98–103. <https://doi.org/10.1038/ijo.2011.40>.
115. Foster C, Anholm JD, Hellman CK, Carpenter J, Pollock ML, Schmidt DH. Left ventricular function during sudden strenuous exercise. *Circulation*. 1981;63:592–6. <https://doi.org/10.1161/01.CIR.63.3.592>.
116. Foster C, Meyer K, Georgakopoulos N, Ellestad AJ, Fitzgerald DJ, Tilman K, et al. Left ventricular function during interval and steady state exercise. *Med Sci Sports Exerc*. 1999;31:1157–62. <https://doi.org/10.1097/00005768-199908000-00012>.
117. Fitrockr. Health data research & analytics platform for garmin wearables [Internet]. <https://www.fitrockr.com/research/>. Cited 16 June 2023.
118. Ekelund U, Tarp J, Steene-Johannessen J, Hansen BH, Jefferis B, Fagerland MW, et al. Dose-response associations between accelerometry measured physical activity and sedentary time and all cause mortality: systematic review and harmonised meta-analysis. *BMJ*. 2019. <https://doi.org/10.1136/bmj.14570>.
119. Peters CH, Sharpe EJ, Proenza C. Cardiac pacemaker activity and aging. *Annu Rev Physiol*. 2020;82:21–43. <https://doi.org/10.1146/ANNUREV-PHYSIOL-021119-034453>.
120. Hammond HK, Froelicher VF. Normal and abnormal heart rate responses to exercise. *Prog Cardiovasc Dis*. 1985;27:271–96. [https://doi.org/10.1016/0033-0620\(85\)90010-6](https://doi.org/10.1016/0033-0620(85)90010-6).
121. Christou DD, Seals DR. Decreased maximal heart rate with aging is related to reduced {beta}-adrenergic responsiveness but is largely explained by a reduction in intrinsic heart rate. *J Appl Physiol*. 2008;105:24–9. <https://doi.org/10.1152/JAPPLPHYSIOL.90401.2008>.
122. Quan HL, Blizzard CL, Sharman JE, Magnussen CG, Dwyer T, Raitakari O, et al. Resting heart rate and the association of physical fitness with carotid artery stiffness. *Am J Hypertens*. 2014;27:65–71. <https://doi.org/10.1093/AJH/HPT161>.
123. Ghoulil H, Farhani Z, Amara S, Hattabi S, Dridi A, Guelmami N, et al. Normative data in resting and maximum heart rates and a prediction equation for young Tunisian soccer players: a cross-sectional study. *EXCLI J*. 2023;22:670–80. <https://doi.org/10.17179/excli2023-6215>.
124. Engels HJ, Zhu W, Moffatt RJ. An empirical evaluation of the prediction of maximal heart rate. *Res Q Exerc Sport*. 1998;69:94–8. <https://doi.org/10.1080/02701367.1998.10607673>.
125. Gellish RL, Goslin BR, Olson RE, McDonald A, Russi GD, Moudgil VK. Longitudinal modeling of the relationship between age and maximal heart rate. *Med Sci Sports Exerc*. 2007;39:822–9. <https://doi.org/10.1097/MSS.0B013E31803349C6>.
126. Hossack KF, Bruce RA. Maximal cardiac function in sedentary normal men and women: comparison of age-related changes. *J Appl Physiol*. 1982;53:799–804. <https://doi.org/10.1152/jappl.1982.53.4.799>.
127. Egan KJ, Knutson KL, Pereira AC, von Schantz M. The role of race and ethnicity in sleep, circadian rhythms and cardiovascular health. *Sleep Med Rev*. 2017;33:70–8. <https://doi.org/10.1016/J.SMRV.2016.05.004>.

128. Valentini M, Parati G. Variables influencing heart rate. *Prog Cardiovasc Dis*. 2009;52:11–9. <https://doi.org/10.1016/J.PCAD.2009.05.004>.
129. Chandra A, Skali H, Claggett B, Solomon SD, Rossi JS, Russell SD, et al. Race- and gender-based differences in cardiac structure and function and risk of heart failure. *J Am Coll Cardiol*. 2022;79:355–68. <https://doi.org/10.1016/J.JACC.2021.11.024>.
130. Kishi S, Reis JP, Venkatesh BA, Gidding SS, Armstrong AC, Jacobs DR, et al. Race-ethnic and sex differences in left ventricular structure and function: the Coronary Artery Risk Development in Young Adults (CARDIA) Study. *J Am Heart Assoc*. 2015;4: e001264. <https://doi.org/10.1161/JAHA.114.001264>.
131. De Becker P, Roeykens J, Reynders M, McGregor N, De Meirleir K. Exercise capacity in chronic fatigue syndrome. *Arch Intern Med*. 2000;160:3270–7. <https://doi.org/10.1001/ARCHINTE.160.21.3270>.
132. Boneva RS, Decker MJ, Maloney EM, Lin JM, Jones JF, Helgason HG, et al. Higher heart rate and reduced heart rate variability persist during sleep in chronic fatigue syndrome: a population-based study. *Auton Neurosci*. 2007;137:94–101. <https://doi.org/10.1016/J.AUTNEU.2007.08.002>.
133. Bornaards CM, Twisk JWR, Van Mechelen W, Snel J, Kemper HCG. A longitudinal study on smoking in relationship to fitness and heart rate response. *Med Sci Sports Exerc*. 2003;35:793–800. <https://doi.org/10.1249/01.MSS.0000064955.31005.E0>.
134. Fick A. Über die Messung des Blutquantums in den Herzventrikeln. *Phys Med Gesell zu Würzburg*; 1870.
135. Stenberg J, Astrand PO, Ekblom B, Royce J, Saltin B. Hemodynamic response to work with different muscle groups, sitting and supine. *J Appl Physiol*. 1967;22:61–70. <https://doi.org/10.1152/JAPPL.1967.22.1.61>.
136. McMurray RG, Soares J, Caspersen CJ, McCurdy T. Examining variations of resting metabolic rate of adults: a public health perspective. *Med Sci Sports Exerc*. 2014;46:1352–8. <https://doi.org/10.1249/MSS.0000000000000232>.
137. Prabhavathi K, Tamarai Selvi K, Poornima KN, Sarvanan A. Role of biological sex in normal cardiac function and in its disease outcome—a review. *J Clin Diagn Res*. 2014. <https://doi.org/10.7860/JCDR/2014/9635.4771>.
138. Waldeck MR, Lambert MI. Heart rate during sleep: implications for monitoring training status. *J Sports Sci Med*. 2003;2:133.
139. Wilmore JH, Stanforth PR, Gagnon J, Rice T, Mandel S, Leon AS, et al. Heart rate and blood pressure changes with endurance training: the HERITAGE Family Study. *Med Sci Sports Exerc*. 2001;33:107–16. <https://doi.org/10.1097/00005768-200101000-00017>.
140. Kang SJ, Ha GC, Ko KJ. Association between resting heart rate, metabolic syndrome and cardiorespiratory fitness in Korean male adults. *J Exerc Sci Fit*. 2017;15:27–31. <https://doi.org/10.1016/J.JESF.2017.06.001>.
141. Chapman JH. Profound sinus bradycardia in the athletic heart syndrome. *J Sports Med Phys Fitness*. 1982;22:45–8.
142. Williams DWP, Koenig J, Carnevali L, Sgoifo A, Jarczok MN, Sternberg EM, et al. Heart rate variability and inflammation: a meta-analysis of human studies. *Brain Behav Immun*. 2019;80:219–26. <https://doi.org/10.1016/J.BBI.2019.03.009>.
143. Shaffer F, Ginsberg JP. An overview of heart rate variability metrics and norms. *Front Public Health*. 2017. <https://doi.org/10.3389/FPUH.2017.00258/PDF>.
144. Marsland AL, Gianaros PJ, Prather AA, Jennings JR, Neumann SA, Manuck SB. Stimulated production of proinflammatory cytokines covaries inversely with heart rate variability. *Psychosom Med*. 2007;69:709–16. <https://doi.org/10.1097/PSY.0B013E3181576118>.
145. Carter JR, Ray CA. Sympathetic neural responses to mental stress: responders, nonresponders and sex differences. *Am J Physiol Heart Circ Physiol*. 2009. <https://doi.org/10.1152/AJPHEART.01234.2008/ASSET/IMAGES/LARGE/ZH40040987440004.JPEG>.
146. Madaniyazi L, Zhou Y, Li S, Williams G, Jaakkola JJK, Liang X, et al. Outdoor temperature, heart rate and blood pressure in Chinese adults: effect modification by individual characteristics. *Sci Rep*. 2016. <https://doi.org/10.1038/SREP21003>.
147. Gubin DG, Weinert D, Rybina SV, Danilova LA, Solovieva SV, Durov AM, et al. Activity, sleep and ambient light have a different impact on circadian blood pressure, heart rate and body temperature rhythms. *Chronobiol Int*. 2017;34:632–49. <https://doi.org/10.1080/07420528.2017.1288632>.
148. Faust L, Feldman K, Mattingly SM, Hachen DV, Chawla N. Deviations from normal bedtimes are associated with short-term increases in resting heart rate. *NPJ Digit Med*. 2020. <https://doi.org/10.1038/S41746-020-0250-6>.
149. Kristal-Boneh E, Harari G, Weinstein Y, Green MS. Factors affecting differences in supine, sitting, and standing heart rate: the Israeli CORDIS Study. *Aviat Space Environ Med*. 1995;66:775–9.
150. Jones AYM, Dean E. Body position change and its effect on hemodynamic and metabolic status. *Heart Lung J Acute Crit Care*. 2004;33:281–90. <https://doi.org/10.1016/j.hrtlng.2004.04.004>.
151. Hinghofer-Szalkay H, Greenleaf JE. Continuous monitoring of blood volume changes in humans. *J Appl Physiol*. 1987;63:1003–7. <https://doi.org/10.1152/JAPPL.1987.63.3.1003>.
152. Børsheim E, Bahr R. Effect of exercise intensity, duration and mode on post-exercise oxygen consumption. *Sports Med*. 2003;33:1037–60.
153. Lu CL, Zou X, Orr WC, Chen JDZ. Postprandial changes of sympathovagal balance measured by heart rate variability. *Dig Dis Sci*. 1999;44:857–61. <https://doi.org/10.1023/A:1026698800742>.
154. Reed GW, Hill JO. Measuring the thermic effect of food. *Am J Clin Nutr*. 1996;63:164–9. <https://doi.org/10.1093/AJCN/63.2.164>.
155. Compher C, Frankenfield D, Keim N, Roth-Yousey L. Best practice methods to apply to measurement of resting metabolic rate in adults: a systematic review. *J Am Diet Assoc*. 2006;106:881–903. <https://doi.org/10.1016/J.JADA.2006.02.009>.
156. Haass M, Kübler W. Nicotine and sympathetic neurotransmission. *Cardiovasc Drugs Ther*. 1997;10:657–65. <https://doi.org/10.1007/BF00053022>.
157. Robertson D, Tseng CJ, Appalsamy M. Smoking and mechanisms of cardiovascular control. *Am Heart J*. 1988;115:258–63. [https://doi.org/10.1016/0002-8703\(88\)90646-1](https://doi.org/10.1016/0002-8703(88)90646-1).
158. Vlachopoulos C, Hirata K, O'Rourke MF. Pressure-altering agents affect central aortic pressures more than is apparent from upper limb measurements in hypertensive patients: the role of arterial wave reflections. *Hypertension*. 2001;38:1456–60. <https://doi.org/10.1161/HY1201.098767>.

159. Ammar R, Song JC, Kluger J, White CM. Evaluation of electrocardiographic and hemodynamic effects of caffeine with acute dosing in healthy volunteers. *Pharmacotherapy*. 2001;21:437–42. <https://doi.org/10.1592/PHCO.21.5.437.34502>.
160. Mahmud A, Feely J. Acute effect of caffeine on arterial stiffness and aortic pressure waveform. *Hypertension*. 2001;38:227–31. <https://doi.org/10.1161/01.HYP.38.2.227>.
161. Sondermeijer HP, Van Marle AGJ, Kamen P, Krum H. Acute effects of caffeine on heart rate variability. *Am J Cardiol*. 2002;90:906–7. [https://doi.org/10.1016/S0002-9149\(02\)02725-X](https://doi.org/10.1016/S0002-9149(02)02725-X).
162. Hirshkowitz M, Whiton K, Albert SM, Alessi C, Bruni O, DonCarlos L, et al. National Sleep Foundation's updated sleep duration recommendations: final report. *Sleep Health*. 2015;1:233–43. <https://doi.org/10.1016/J.SLEH.2015.10.004>.

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