

Short title: HRV and Mortality

Heart rate variability in the prediction of mortality: A systematic review and meta-analysis of healthy and patient populations

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1. Introduction

Heart rate and its variability plays a role in disease diagnosis and prognosis since the ancient medicus Galen (Billman, 2011). The development of the electrocardiogram (ECG) measurement in 1895 represents a milestone in medical history allowing an accurate measurement of heart rate. Only this allowed the precise calculation of the variability between heartbeats, i.e., heart rate variability (HRV). Early experiments showed that this variability originates from the autonomous nervous system (Akselrod et al., 1981), primarily from rapid vagal (dis)inhibition (Warner and Cox, 1962). At the same time, reduced HRV has been linked to worse prognosis in disciplines as diverse as cardiology (Malik and Camm, 1993) and gynaecology (Freemann R, Garite T, 1981). In the following decades, parameters of HRV were negatively associated to a multitude of somatic diseases and disorders including increased risk for cardiovascular disease (Jarczok et al., 2012; Kristal-Boneh et al., 1995; Mercedes R Carnethon et al., 2002; Schuster et al., 2016) and myocardial infarction (MI) (Thayer and Lane, 2007) as well as to a wide range of psychopathologies including depressive, bipolar and posttraumatic stress disorder (PTSD) (Faurholt-Jepsen et al., 2017; Heiss et al., 2021; Holzman and Bridgett, 2017; Jandackova et al., 2016; Schneider and Schwerdtfeger, 2020; Sigrist et al., 2021). Consequently, an increased risk of mortality was repeatedly reported and summarized for populations with cardiovascular diseases (Fang et al., 2020) and cancer (Xin Zhou), but interestingly there are also reports for general population samples (e.g., Dekker et al., 1997; Huikuri et al., 1998; Mäkikallio et al., 2001; Wulsin et al., 2015).

HRV as a term meanwhile is a theoretical background, represented by many different parameters. There are a lot of different approaches to calculate variability: time-domain bound like the root mean square of successive differences (RMSSD) and the standard deviation of normal-to-normal intervals (SDNN), frequency-domain bound like high frequency power (HF), low frequency power (LF), simple ones like peak to valley and more complex ones like nonlinear dynamics with the theoretical

background of chaos theories (Lombardi, 2000; Shaffer and Ginsberg, 2017). To date, it has not been demonstrated that specific parameters are superior to others across settings. This is understandable on the common background that – all parameters are derived from the RR-intervals and therefore are highly correlating. Nevertheless, it also has been shown that some parameters represent different aspects better than others, like RMSSD and HF reflect vagal compounds (Shaffer and Ginsberg, 2017; Wulsin et al., 2018). It is unclear whether specific parameters such as the SDNN or RMSSD and/or settings (e.g., long-term measurements) have advantages over others and can provide better risk prediction.

Next to these issues, there are many various possible measurement settings (posture such as supine, seated, and period such as during work, leisure time, sport or unstructured) and lengths of measurements (10 sec up to 48 hours), as well as genetic aspects and the influence of existing diseases.

Therefore, the purpose of this meta-analysis is to investigate the predictive power of HRV measures for mortality, i.e., investigating to what extend an observed HRV measure at a baseline can predict the time to event (death) from either all-cause or cardiac mortality. We aim to compare different measures of HRV to investigate if mortality prediction by HRV parameters differs dependent on type of HRV parameter and settings of HRV measurement and investigated population. We chose all-cause mortality and cardiac mortality to compare if HRV is more powerful in prediction of the unspecific all-cause mortality than in specific mortalities, of which cardiac mortality has been researched most often.

To generate valid results, we focused on HRV parameters that have been reported more often, as less common parameters with small number of participants will not add qualitative data to the meta-analysis. A pre-search showed that the most commonly parameters were those which are also recommended by the Task Force of the European Society of Cardiology and the North American Society of Pacing and Electrophysiology (Camm et al., 1996): SDNN, SDANN, TP, VLF, LF and HF, while

other parameters such as SD1, SD2, DFalpha were found only in single studies, thus not fulfilling the criteria for meta-regression.

We hypothesize to observe systematic shorter time-to-event episodes in individuals with lower measures of HRV at study inclusion. We further hypothesize that this principle is independent of various study settings including population type or observation time.

2. Methods

The study was conducted according to the preferred reporting items for systematic reviews and meta-analyses (PRISMA) statement (Moher et al., 2009).

2.1. Search strategy

Figure 1 summarizes the results of the systematic literature search. The Medline database was screened for studies reporting time to event (i.e., mortality) information as outcome and frequency or time domain parameters of HRV as predictor. The terms mortality and cardiac electrophysiology were entered as Medical Subject Heading (MeSH). The term heart rate variability or its variants were screened in title and abstract. Language restriction was set to German or English.

Search terms for MEDLINE (via PubMed)

("mortality"[mesh]) AND

("HRV"[tiab] OR "Heart rate variability"[tiab] OR "Heart rate variabilities"[tiab] OR "Heart period variability"[tiab] OR "Cardiac autonomic activity"[tiab] OR "Cardiac autonomic control"[tiab] OR "Cardiac reactivity"[tiab] OR "Cardiac vagal control"[tiab] OR "Autonomic nervous system activity"[tiab] OR "Parasympathetic nervous system activity"[tiab] OR "Vagal tone"[tiab] OR "Vagus nerve"[Mesh] OR "Vagal activity"[tiab] OR "Vagal reactivity"[tiab] OR "Vagus nerve"[tiab] OR "Parasympathetic nervous system"[tiab] OR "Cardiac electrophysiology"[Mesh]) AND

"Humans"[Mesh] AND

("1997/01/01"[PDAT] : "3000/12/31"[PDAT]) AND

(English[lang] OR German[lang])

The initial search was conducted in June 2019 by the authors CB, EB, MNJ and KW and was updated in March 2021 by MNJ and EB. All empirical studies published between 01.01.1997 and 30.06.2019 were considered. The search was updated in March 2021. Initial database search revealed 231 initial records with two more records identified by snowballing and 21 more captured by updating.

2.2. Study selection

Studies were included into the systematic review and meta-analysis if the following a priori criteria were met:

- (1) study represents original research
- (2) published in English or German
- (3) as an original article or brief report in a journal (i.e., no comments, dissertations, conference abstracts, reviews)
- (4) reports adjusted or unadjusted results from Cox proportional hazard models
- (5) for either all-cause or cardiac mortality
- (6) reports a measure of HRV (see Section 2.4.1)
- (7) in human adults (age ≥ 18 years)

Studies were included if at least one distinguishable subsample was reported that met the inclusion criteria. By applying the inclusion criteria to the information contained in the title and abstract, the pool of records was reduced to 115. After reviewing the full text, a total of 35 samples from 32 publications remained. In addition, two open access samples (MIDUS 2 & MIDUS Refresher,

WHITEHALL II) were analyzed and added to this meta-analysis (see details below). Thus, K=34 studies with S=37 samples containing data from N=38,008 individuals were available.

Abstract screening and selection were carried out independently by the authors MNJ, CB and KW in July 2019. Disagreements were discussed by this triad and resolved on a consensual basis. An overall interrater agreement of 89.2% was assessed for the title/abstract screening in July 2019.

Full texts were screened by MNJ, CB, KW and EB in September 2019 in dyads. As before, disagreements were solved by consensus. Interrater agreement was very high with 91.8% for the full text screening.

At the update in March 2021, disagreements were discussed by MNJ and EB and solved by consensus. The agreement rate for the update was 90.5%.

In case of any missing or inconclusive information in the full text, the corresponding author of the publication was searched for electronically and then repeatedly contacted by MNJ, KW or EMB via email with a friendly request to provide the missing or additional information. If no current address of the corresponding author could be found, the co-authors were approached. Altogether, 13 authors from 13 full-text publications were contacted, of which only two authors responded concerning two of the publications. At this point, we would like to thank all responding authors for their cooperation.

Two publications were excluded to avoid inclusion of duplicate data (i.e. publications reporting data from the same dataset). First, Huikuri et al. (1998) report results from the same dataset as Mäkikallio et al. (2001) but the latter reported more relevant parameters to this systematic review such as unadjusted and adjusted associations. Therefore, we included results reported by Mäkikallio et al. (2001). Second, Cygankiewicz et al. (2015) report results from a subgroup analysis of the population published in Cygankiewicz (2009). Therefore, we included results reported in 2009.

2.3. Individual participant data (IPD)

Two large population studies (MIDUS 2 and MIDUS Refresher: N=1891 & Whitehall II wave 5: N=7870) were available for analysis. The individual studies received local ethical approval at their respective institutions.

MIDUS 2: Open access data from the biomarker project of the second wave (2004-2009) of the *Midlife Development in the U.S.* study (MIDUS 2, P4; N=1255 ICPSR 29282 v9 from Mar 27, 2019 <https://doi.org/10.3886/ICPSR29282.v9>) and the MIDUS Refresher (MIDUS R, P4; N=738; ICPSR 36901 v6 from Nov 18, 2019 <https://doi.org/10.3886/ICPSR36901.v6>) were matched with survival information from the MIDUS core mortality dataset (*MIDUS_Core_MortalityCauseData_N2124_20220310.sav*) until December 2020 (median follow-up = 13.6 years) and the MIDUS refresher mortality dataset (*MIDUS_Refresher_MortalityCauseData_N167_20210305.sav*) until March 2021 (Median follow-up = 6.3 years). The combined median follow-up is 11.8 years with 234 deaths from all-cause mortality recorded (11.1%). According to the study description, institutional review board approval was obtained prior to study start and informed consent was obtained from each participant prior to enrolment in the MIDUS study. Further study details and data are publicly available on the official website after free registration (<https://midus.colectica.org>). Two supine 5-minute ECGs were obtained during rest and HRV measures were derived per 5-minute interval and averaged for this analysis. In cases where only one 5-minute interval was valid or available, the single 5-minute interval was used. The present study included participants with data on all-cause mortality, HRV (RMSSD, SDNN, HF, LF), age (years), sex (male vs. female), and ethnicity (white vs. non-white) at baseline, as well as survival information.

Whitehall II: Data from the fifth (1997–1999) phase of the UK Whitehall II longitudinal population-based cohort study was analyzed with a median follow-up of 17.2 years (N=7870 participants in wave 5) and end of follow-up in June 2015. This ongoing cohort study of subjects initially targeted London-based British civil service office staff, aged 35-55 years (Marmot and Brunner, 2005). The ECG

recordings were collected at the fifth (1997–1999) wave. A 5-minute supine resting 12-lead ECG (Kardiosis™ ECG acquisition module, Tapa, Inc., Turkey and Getemed ECG recorder, Getemed Teltow, Germany) was obtained after five minutes of rest and HRV measures were calculated. The University College London ethics committee approved the study and participants gave informed consent. Whitehall II data, protocols and other meta-data are available to bona fide researchers for research purposes (data sharing policy is available at <https://www.ucl.ac.uk/epidemiology-health-care/data-sharing-faq>). The present study included participants with data on all-cause and cardiac mortality, HRV (RMSSD, SDNN, HF, LF, Total Power [TP]), age (years), sex (male vs. female), and ethnicity (white vs. non-white) at phase 5 (1997–1999), as well as survival information from phases 11 (2012–2013, last updated in June 2015).

For both studies, analyses according to the Cox proportional hazard regression model were performed and HR were extracted with their respective 95% confidence intervals (CI).

2.4. Data extraction and summary statistics

Data on study characteristics, methodological quality and outcomes were extracted independently by two author dyads (MNJ & KW and CB & EB) in accordance with the data extraction sheet. After an initial trial phase including three full texts per author, the items of the extraction sheet were discussed and carefully reviewed for inconclusive items and adjusted if necessary. During the process of study extraction, the sheet was continually adapted. If data were not extractable (i.e., only provided in graphs), authors were contacted and asked for additional information.

The following data were extracted from all included studies:

- (1) names of all authors,
- (2) year of publication,
- (3) year and place of data collection,
- (4) sample size,

- (5) population characteristics such as type of sample (diagnostics), mean age, proportion of females, mean length of follow-up,
- (6) artifact handling,
- (7) excluded subjects,
- (8) total length of heart rate assessment,
- (9) position and paced breathing,
- (10) recorder type and software,
- (11) type of mortality,
- (12) covariates adjustment for in the statistical analysis,
- (13) the HRV parameter type and metric for predicting end points,
- (14) HRs with 95% CI.

Populations were coded into 'clinical' (populations with existing disease like coronary artery disease, cirrhosis, dialysis) and 'non-clinical' (populations characterized as healthy or as general population).

Setting was categorized in 'ambulatory' (mostly 24h measurement), 'supine' (short time measurement), 'sitting' (short time measurement, if explicitly mentioned) and 'resting' (short time measurement without information about posture).

Mortality was categorized as 'all-cause' if it was reported as all-cause and as 'cardiac' if it was reported as cardiac death, arrhythmic death, cardiovascular death, and fatal coronary heart disease.

If HRs were reported from several Cox regression models, priority was given to HRs from covariate unadjusted Cox regression as well as to HR from fully adjusted Cox regression models as reported by the authors of the respective study (see section 2.4.2 Hazard ratios).

If standard errors (SE) were reported instead of standard deviations (SD), the SD was estimated in accordance to an earlier meta-analysis, by using the following formula: $SD = SE \times \sqrt{n}$ (Higgins et al., 2019).

2.4.1. Measures of HRV

All measures of HRV are rooted in the variability observed from intervals between adjacent normal R waves (NN intervals). A detailed description about HRV parameters is summarized by Ginsberg and Shaffer (Shaffer and Ginsberg, 2017). In brief, three broad categories are distinguished: time domain measures enumerating the amount of variability in measurements such as the square root of the mean of the sum of the squares of differences between adjacent R-R intervals (RMSSD), percentage of successive RR intervals that differ by more than 50 ms (pNN50) or standard deviation of all normal-to-normal intervals (SDNN), and the SD of the average normal-to-normal intervals for each 5 min segment (SDANN) in 24h recordings. Frequency-domain measures estimate the distribution of absolute or relative power commonly within four frequency bands: high frequency (HF), low frequency (LF), very low frequency (VLF), and ultra-low frequency (ULF) as well as total power (TP) to capture the power across all bands. Finally, there are nonlinear methods such as sample entropy (SampEn), which is a measure of time series regularity and complexity. The following most frequently reported HRV parameters, which are also recommended by the Task Force of the European Society of Cardiology and the North American Society of Pacing and Electrophysiology (Camm et al., 1996) are selected for the present meta-analysis: SDNN, SDANN, TP, VLF and LF, all measures capturing variability from mixed sources such as sympathetic, parasympathetic, thermoregulation and functional capacity, as well as RMSSD, HF or pNN50, for example, all measures capturing primarily vagal influence. Importantly, the calculation of VLF is not recommended from short segments (Nunan et al., 2010). Thus, studies reporting VLF values were only included from studies that recorded for a sufficient amount of time (50 minutes minimum i.e., 10 cycles of lower frequency band border 0.0033Hz) to resolve this frequency. The following studies report VLF calculated from 5- or 10-minute intervals (Bhogal et al., 2019; Hotta et al., 2005; Lanza et al., 2006; May and Arildsen, 2012; Quintana et al., 1997) or no information of VLF calculation (Badarau et al., 2015) and VLF parameters from these studies were therefore not included.

In addition, SDNN is dependent on the recording length such that comparisons between 5-minute recordings vs. 24h recordings are problematic. Therefore, studies with SDNN recorded for 24h and analyzed as a single segment with 24h were summarized separately (Ablonskytė-Dūdonienė et al., 2012; Lanza et al., 2006; Mäkikallio et al., 2001; Stein et al., 2008).

Other parameters such as SD1, SD2, DFalpha were not considered as they were only found in single studies, thus not fulfilling the criteria for meta-regression.

Both absolute and logarithmically transformed values of HRV have been included in the present meta-analysis. Normalized values have not been included to this meta-analysis due to the fact that these measures are calculated on the basis of parameters such as LF and TP (Burr, 2007). The aim of the present meta-analysis was to associate single parameters of HRV with mortality.

2.4.2. Hazard ratios

Nearly all full texts reported HRs with respective 95% CI. CIs of 99% were transformed to 95%. Similarly, the HR and its variance were log transformed as recommended previously (Parmar et al., 1998). This measure allows for censoring and accommodates variable length of follow-up for each of the included trials. The respective SE was calculated as $\sqrt{(CI_{upper}-CI_{lower}/3.92)^2}$ for 95% CI and $\sqrt{(CI_{upper}-CI_{lower}/5.15)^2}$ for 99% CI in case no SE was reported, but CI were available (Higgins et al., 2019).

The direction of effects was carefully reviewed by MNJ and EBM and conclusively interpreted as a decrease in the HRV parameter per unit of interest. For example, some study authors report cut off values such as the SDNN <20ms vs. ≥20ms and compared groups while others report an increase per millisecond (ms) in SDNN. The latter was inverted using the formula $1/HR$ to correspond to a decrease per ms. As a matter of fact, per unit change is different across studies (cut-offs vs. increment decreases) and is therefore a potential source of heterogeneity. Also, the default inverse variance weighting is replaced with weights per study by their respective study size N (see below).

Several studies presented HR from covariate unadjusted and covariate adjusted Cox regression models. If multiple covariates adjusted Cox regression models were reported, priority was given to the model including age and sex and a minimum of other covariates to enhance between study comparability. Age and sex are known as important determinants of HRV. It would have been best if all models would have been available containing the covariate age and sex only. However, as selection of adjusted covariates widely differed between studies, we also extracted the HR from unadjusted Cox-regression. Meta-Analytic regression tried to estimate the between study effect of different compositions regarding age, sex, population type, etc. (see section 3.2.2. potential effect modifiers).

2.4.3. Risk of bias assessment

We assessed the risk of bias via the following criteria: We added a time window to HRV sampling, with 'one' referring to a well-defined time window (time and situation) and 'zero' referring to a too variable or unclear time window (e.g., between 18 and 24 hours). As a number of studies report only significant and no secondary HRV parameters, HRV parameter selection represented the next criterion. Studies reporting all time-domain and frequency-domain parameters valid for the time window of measurement were assigned one point and those reporting only arbitrarily selected parameters were marked by 0. Quality of artifact control was rated with 'one' if artifact handling was reported and adequate. If too many artifacts (more than 10%) were included or if artifact handling was not reported at all, '0' was assigned. If screening procedures were poorly described (e.g., no flow chart of included and excluded subjects) population bias was assumed and selection bias could not be assessed in the following.

For all criterion points were summed up. The results were rated on a zero (low quality, bias possible) to four (high quality, low probability of bias) range.

2.5. Methodological quality assessment

Leave-one-study-out sensitivity analysis was performed by sequential exclusion of each trial and comparing results graphically, for subgroups (type of mortality all-cause or cardiac, unadjusted- vs. covariate adjusted, HRV parameter) with at least three or more studies. A study was considered to be influential if the point estimates of the leave-one-out-pooled outcomes were not within the 95% CI of the original pooled outcomes.

2.6. Statistical analysis

STATA version 15.1 (Stata Corporation, College Station, TX, USA) with the package *metan* (Version 20210224) was used to perform the meta-analysis. Weighted generalized least squares meta regressions were calculated using SPSS Version 25 (SPSS© Statistics 25, IBM, USA) with the macro *metareg*. HRs and SEs were accordingly extracted from full texts or IPD-datasets and log-transformed. Study weights were set to actual study size, i.e., the reported N of the respective Cox regression model if reported. This method is replacing standard inverse probability weighting, because of the different metrics of reported parameters such as ms, log(ms), or quartiles. These different metrics can be a large source of varying SEs between studies and can bias inverse probability weighting. However, this led to a larger number of studies reporting smaller increments (i.e., per millisecond changes) while studies reporting larger increments (i.e., per quartile change, per SD change) were underrepresented.

Statistical heterogeneity was assessed by using the I^2 statistic. This method quantifies the amount of variation between studies that can be attributed to true variation in effect sizes rather than sampling error. Importantly, I^2 does not depend on the number of studies included in the meta-analysis or the metric of the effect size (Higgins and Thompson, 2002). An I^2 of 25%, 50% and 75% is considered as low, moderate and high heterogeneity, respectively. Heterogeneity was considered to be present if I^2 exceeded 25% (Higgins and Thompson, 2002). If heterogeneity was present, random-effect models were interpreted, if no heterogeneity was present, fixed-effect models were interpreted. A p-value <0.05 was considered to be statistically significant.

2.7. IPD subgroup analysis heart rate correction

To eliminate the potential effect of heart period on HRV, a sub-analysis was performed for the parameters SDNN and RMSSD calculating the coefficient of variation (cvRMSSD and cvSDNN) as suggested by de Geus et al, (2019). For these Cox regression analyses, fixed effect inverse variance weighting was used, calculating HR for the lowest quartile vs. the rest of the respective HRV parameter.

3. Results

3.1. Search results and study selection

A total of 32 studies and two datasets (K=34) summarizing data from 38,008 individuals were analyzed (see Figure 1). From some studies, more than one population sample (S) were extracted because authors separated results by subgroups (Dekker et al., 1997; La Rovere et al., 2003; Liao et al., 2002).

Four major groups were classified: Studies reporting results on all-cause mortality and cardiac mortality each either derived from covariate unadjusted or covariate adjusted Cox proportional hazard models. If authors from included studies reported both, unadjusted and adjusted HRs, both were extracted. We present studies reporting results on A) all-cause mortality from covariate unadjusted Cox proportional hazard models (K=18; S=18; N=17,990) (Ablonskytė-Dūdonienė et al., 2012; Bhogal et al., 2019; Gilliam et al., 2007; Hotta et al., 2005; Kiviniemi et al., 2007; Lanza et al., 2006; Macfarlane et al., 2007; Mäkitallio et al., 2001; Medenwald et al., 2017; “MIDUS - Midlife in the United States, A National Longitudinal Study of Health and Well-being,” n.d., “Whitehall II | Institute of Epidemiology & Health Care - UCL – University College London,” n.d.; Oikawa et al., 2009; Quintana et al., 1997; Shibasaki et al., 2014; Singh et al., 2009; Steeds et al., 2004; van Bommel et al., 2006) and on B) all-cause mortality from covariate adjusted Cox proportional hazard regression models (K=17; S=19; N=31,595) (Bhogal et al., 2019; Carney et al., 2005; Dekker et al., 1997; Gilliam

et al., 2007; Kiviniemi et al., 2007; Kotecha et al., 2019; Lanza et al., 2006; Liao et al., 2002; Macfarlane et al., 2007; Mäkikallio et al., 2001; May and Arildsen, 2012; Medenwald et al., 2017; “MIDUS - Midlife in the United States, A National Longitudinal Study of Health and Well-being,” n.d., “Whitehall II | Institute of Epidemiology & Health Care - UCL – University College London,” n.d.; Oikawa et al., 2009; Shibasaki et al., 2014) as well as studies reporting results on C) cardiac mortality from unadjusted Cox regressions (K=16; S=17; N=16,724) (Ablonskytė-Dūdionienė et al., 2012; Cygankiewicz et al., 2009; Hayano et al., 2001; Kida et al., 2017; Kiviniemi et al., 2007; Kop et al., 2010; La Rovere et al., 2003; Lanza et al., 2006; Macfarlane et al., 2007; Mäkikallio et al., 2001; Medenwald et al., 2017; Nishimura et al., 2010; Oikawa et al., 2009; Stein et al., 2008; van Bommel et al., 2006; “Whitehall II | Institute of Epidemiology & Health Care - UCL – University College London,” n.d.) and on D) cardiac mortality from covariate adjusted Cox regressions (K=15; S=18; N=21,506) (Cygankiewicz et al., 2009; Dekker et al., 1997; Hayano et al., 2001; Kamphuis et al., 2007; Kida et al., 2017; Kiviniemi et al., 2007; La Rovere et al., 2003; Lakusic et al., 2013; Lanza et al., 2006; Liao et al., 2002; Mäkikallio et al., 2001; Medenwald et al., 2017; Nishimura et al., 2010; Oikawa et al., 2009; “Whitehall II | Institute of Epidemiology & Health Care - UCL – University College London,” n.d.).

The sample and study characteristics of the included studies are summarized in Table 1. Most of the included studies (K=13) recruited participants from general population samples (N= 29,716; 78% of total population), followed by populations suffering from cardiac diseases (K=14; N=5935; 16% of total population) such as myocardial infarction and heart failure.

ECG measurement varied by technique and duration as well as artifact correction and calculation of HRV parameters. The setting of ECG measurement was predominantly ambulatory (K=20; 59% of included studies), followed by supine (K=7; 21% of included studies), or at rest (K=6; 18% of included studies). The studies were conducted in Europe (K=19; 56%), North America (K=9; 26%) or Asia (K=6; 18%). The average female proportion was 39% across all studies (range 0% to 75%).

The number of reported HRV parameters of interest varied between one and seven. Twenty-eight out of 37 samples reported SDNN, the most commonly reported parameter, but see discussion section for issues with this parameter.

The results of the methodological quality assessment are listed in Table 2. The time window for HRV analysis was mostly well-defined. Artifact control was poorly described in about two thirds of the studies, while the remaining one third described their procedures in depth. About two thirds of the authors reported HRV parameters only in a selective way. Over two thirds of the studies showed no evidence for population bias. On a scale from 0 (low quality) to 4 (high quality), the average score was 2.2. Only four included studies met all four quality criteria (N=4261; 11%).

3.2. Data analysis

3.2.1. Meta-analysis

An overview of the summary results per HRV parameter of the meta-analysis is presented in Figure 2. All details of the results per included study of the meta-analysis are shown in Table 3. Analysis of the pooled data revealed significantly higher all-cause mortality per unit decrease in SDNN and LF, regardless of the model (unadjusted or covariate adjusted) and for both all-cause and cardiac mortality. The effect size for RMSSD (HR 1.25 95%CI 1.04 – 1.51) and HF (1.56 95%CI 1.37 – 1.77) was highest in unadjusted models for all-cause mortality.

Detailed forest plots are shown in Figure 3-6 for all-cause and cardiac mortality presenting both, covariate unadjusted and covariate adjusted cox proportional hazard models as extracted from the included studies. Due to the high heterogeneity between the studies for most HRV measures, meta-analysis results are reported from random-effect models if $k \geq 2$. Forest plots displaying results from fixed-effects models are shown in Supplemental Figures S1-S4.

3.2.1.1. Correcting for heart rate

To explore the effect of correction for heart rate (i.e., coefficient of variation), individual participant dataset analysis was conducted (i.e. Whitehall and MIDUS). The overall effect did not change

relevantly for models using cvSDNN ($k=2$; $N=5251$ from HR_{SDNN} 1.23 [95%CI: 1.03 – 1.46] to HR_{cvSDNN} 1.21 [95%CI: 1.01 – 1.44]) or cvRMSSD ($k=2$; $N=5251$ from HR_{RMSSD} 1.17 [95%CI: 0.98 – 1.39] to $HR_{cvRMSSD}$ 1.13 [95%CI: 0.95 – 1.34]).

3.2.2. Potential Effect modifiers

Meta regressions on random effects were performed for HRV parameters if $k \geq 9$. Here, only statistically significant effects are reported. Full results are presented in Table 4.

Meta regression of HRs from studies reporting covariate unadjusted models for all-cause mortality revealed the following: Recording length in minutes or as binary indicator (short term vs. rest) as well as mean follow-up time in month modified statistically significant the HRs of all-cause mortality for the HRV parameter LF. Here, studies with shorter recording length (binary; b -0.606, 95%CI -1.134 – -0.078) and shorter mean-follow-up time (b -0.003, 95%CI -0.006 – -0.000) had a lower effect size in studies reporting HRs from covariate unadjusted models.

SDNN effects on cardiac mortality were significantly modified by the type of population and mean follow up time. Here, studies reporting HRs from covariate unadjusted models from non-clinical populations reported smaller effect sizes in SDNN (b -0.814, 95%CI -1.375 – -0.253). Also, studies with shorter recording length (binary; b -0.610, 95%CI -1.186 – -0.033) had a lower effect size in studies reporting HRs from covariate unadjusted models.

Similar, the effect of LF on cardiac mortality were significantly modified by the type of population and mean follow up time. Here, studies reporting HRs from covariate unadjusted models from non-clinical populations reported smaller effect sizes in LF (b -0.837, 95%CI -1.278 – -0.396). Also, studies with shorter mean-follow-up time (b -0.005, 95%CI -0.008 – -0.003) had a lower effect size in studies reporting HRs from covariate unadjusted models.

Meta regression of HRs from studies reporting covariate adjusted models did not reveal any significant effect modifier regarding age (years), female proportion (%), recording length (minutes), continent (Asia vs. rest), decade of study conduction (post 2000 vs. rest), quality of artifact control

(good vs. rest), population (clinical/non-clinical), or length of follow-up (month) for all-cause nor for cardiac mortality (see Table 4).

3.2.3. Sensitivity analysis

The majority of pooled HR values were consistent with the full random effect meta-analysis after sequential exclusion of each study (leave one out) – stratified by covariate unadjusted vs. covariate adjusted studies, type of mortality and HRV-parameter. Two studies were considered to be influential. Studies reporting HRs from covariate unadjusted models, the Whitehall study significantly influenced HR to be lower for TP (K=4; HR 1.61 [1.14 – 2.27] vs. K=3; HR=2.30 [1.75 – 3.02]) and LF (K=11; HR 1.71 [1.51 – 1.93] vs. K=10; HR=1.97 [1.68 – 2.31]) predicting all-cause mortality. Leaving out the results by Gilliam et al. 2007 changed the effect of SDANN predicting all-cause mortality to a significantly smaller effect in covariate adjusted models (K=4; HR 4.83 [1.46 – 16.04] vs. K=3; HR=1.12 [0.94 – 1.33]).

4. Discussion

With the present comprehensive meta-analysis of 32 studies and two IPD-datasets, we aimed to quantify the association between parameters of HRV and all-cause and cardiac mortality measures. The first hypothesis, to observe systematic shorter time to event episodes in individuals with lower measures of HRV at study inclusion, is supported. The second hypothesis, that systematic differences in cardiac and all-cause mortality exist independently of study features such as age, sex, recording length, continent, decade of study conduction, quality of artifact control, population (clinical/non-clinical), or length of follow-up, could also be confirmed. In other words, despite the existing heterogeneity across included populations, study features, and HRV parameters, there appears to be a substantial negative association between measures of HRV and mortality.

4.1. Clinical application

For clinical application, we present a composite HR of 1.56 (95% CI: 1.32–1.85) for a 5-min-RMSSD value within the lowest quartile. The magnitude of this HR is higher than that for shortened sleep

defined as less than 7 h on average per night (relative risk [RR]=1.10, 95%CI 1.06–1.15) (Gallicchio and Kalesan, 2009) and exceeds the 34% higher mortality risk for adults sitting 10 h/day (Chau et al., 2013) as well as the risk of harmful drinkers (RR=1.37, 95%CI 1.35–1.49) (Holman et al., 1996) and the mortality risk of white coat hypertension (HR=1.33, 95%CI 1.07–1.67) (Cohen et al., 2019). This risk is slightly lower compared to the all-cause mortality risk in masked uncontrolled hypertension compared to controlled hypertension (HR=1.80, 95%CI 1.57–2.06) (Pierdomenico et al., 2018). However, fitness is an even stronger predictor of mortality with a two to three times increase of mortality risk (Barry et al., 2018).

The pooled HR for 5-min-RMSSD from unadjusted models from studies that reported HRs comparing the lowest quartile with the other quartiles is shown in Figure 7. The average follow-up time of these studies was 14.5 years. According to this, an individual with a 5-min-RMSSD of the lowest quartile of its reference population would have a 56% higher mortality risk over the next 15 years [95%CI: 32% – 85%]. In these studies, the weighted mean value for the lowest quartile was 12 ms (MIDUS: 9.25 ms; Steeds: 8.3 ms; Whitehall II: 13.3 ms).

From **a clinical perspective**, a five minutes ECG measurement with calculation of SDNN or RMSSD and subsequent quartile classification would be sufficient to identify individuals at risk, without the necessity to assess additional parameters such as a full laboratory profile. For example, we showed in a recent study in 19 different occupational settings with more than 9500 participating employees that RMSSD values below 25 ± 4 ms were associated with an elevated risk across a range of established cardiovascular risk factors (Jarczok et al., 2019b). Because parameters of HRV are a nonspecific correlate of disease risk, it makes it particularly suitable for a prevention setting. Despite the great scientific progress in interventional medicine over the past decades, chronic diseases such as MI and stroke continue to be common diseases causing great suffering and high costs. In 2019, the first nine leading causes for disability-adjusted life years were attributed to non-communicable

diseases (Abbafati et al., 2020). This high number could be reduced through prevention. Although it is widely agreed that prevention is necessary, it is often not transferred into practice (Fineberg, 2013). One problem is communicating potential risk to apparently healthy individuals. This problem is addressed by risk assessments, but commonly they are developed to be disease specific e.g. for coronary heart disease like the Framingham risk score (Wilson et al., 1998) or for diabetes like the Findrisk score (Lindström and Tuomilehto, 2003). Despite the disease specific nature of these scores, they not only show high intercorrelations among each other, but also a medium to high adjusted rank correlation with vagally-mediated HRV (Schuster et al., 2016). The preventive advice from these risk scores overlap to a large extent, as behavioral advice commonly include what the authors would call the hattrick of lifestyle interventions: engage in physical and social activities, eat a healthy diet, and take care of relaxation such as sleep hygiene but also detachment from work (Avery et al., 2012; Dickinson et al., 2006; Galani and Schneider, 2007; O'Connor et al., 2021; Thayer and Lane, 2007). Therefore, HRV is an ideal risk parameter that can be used to measure individual risk for disease in general and, as an outlook, also to measure the effectiveness of lifestyle changes. Interestingly, several of these lifestyle interventions have been reported to improve measures of HRV such as physical activity (Tornberg et al., 2019), relaxation techniques (Lin et al., 2014), healthy diet (Young and Benton, 2018) or slow-paced breathing (Laborde et al., 2022; Sevoz-Couche and Laborde, 2022).

4.2. Effect modifiers

Comparing the HRs of studies conducted with **general populations to ill patients**, the relation between HRV parameter and mortality remains comparable. There is no evidence for any restriction of its use to a special population, as it appears to be valid in health and disease. Furthermore, no differences in effect sizes were found across participants' age. Thus, our results support its possible use as an age-independent risk marker for primary prevention.

The **length of the measurement** is another parameter that can influence the quality of results. Meta-regression did not find any systematic differences in HRs resulting from measurements over 10

seconds compared to 24 hours. From this perspective, a short-term segment seems to be sufficient to predict mortality. Furthermore, artifact handling is easier in shorter measurements, so results are more uniform and are less affected by ‘noise’ caused by artifacts. Shorter measurements were taken almost always at rest within a controlled setting, whereas 24-hour measurements obviously include daily activities which vary highly inter-individually. In conclusion, in the present context of mortality prediction, the length of the measurement can be selected dependent on other local and study requirements. Next to pure risk stratification, a 24h-measurement of HRV can serve as a risk communication tool, including identification of individual resources and strains and improving the knowledge about interactions between body, mind and environment (Jarczok et al., 2021). Based on the data presented in this manuscript, the advantages of a 24h-measurement lie in additional information, not in better risk stratification. This additional information includes insights into circadian rhythms (Jarczok et al., 2019a) that can be disturbed in shift work, but also in ongoing psychosocial stress as well as in diseases like depression (Jarczok et al., 2018). Further, the 24h measurements allow to answer specific work-related questions, e.g., if breaks are taken early enough to prevent exhaustion or if breaks are recreational. This information can’t be obtained from a five minutes measurement.

Whether HRV should be measured during ‘standard’ rest or during **paced breathing**, to standardize the influence of the breathing rate on HRV parameters, is also an important question in this context. In the studies we found, only Medenwald used paced breathing (Medenwald et al., 2017). Interestingly, the HRs reported were not significant. If HRV is a mortality predictor, then the value measured should be quite stable, like a trait. Considering this, the parameter of interest would represent a ‘typical’ HRV of the investigated person. In slow paced breathing, HRV is maximized with immediate beneficial effects (Lehrer and Gevirtz, 2014). Though the maximum potential of an individuals’ HRV is of interest as it tests the limits of his ANS, an HRV measured during their natural breathing pace probably represents the ANS activity that is prevalent most of the time, therefore being more useful in mortality prediction. In addition, some HRV indices (e.g., RMSSD) and some

recording procedures (i.e., detrending and normal distribution, see Lewis et al 2012) are less susceptible to respiratory influences and thus appropriate HRV estimation can enhance the utility for prediction and prevention.

In assessing the significance of HRV in predicting mortality, it has been stated that it would be necessary to correct for **heart rate** or for lifestyle variables like physical activity, especially when this activity occurred during the measurement in 24h-measurements (de Geus et al., 2019). Of course, heart rate and HRV are correlated (Sacha, 2014), as well as physical activity and HRV (Blom et al., 2009; Camillo et al., 2011; May et al., 2017). On the brain's level, the ANS modulates both, heart rate and HRV, in rest and during physical activities (Thayer et al., 2012). On the other hand, physical activity changes settings of brain circuits, not only those of the ANS, but also emotion regulation e.g. in depression (Martinsen, 2009). Coming back to mortality, both heart rate (Lau et al., 2021; Zhang et al., 2016) and physical activity are strongly related to mortality (Leroux et al., 2021). If HRV is also a predictor of mortality, there has to be a correlation between heart rate, HRV and physical activity. Correcting HRV for heart rate and physical activity might remove a part of its predictive power. Yet, sub-analysis of IPD revealed only minor differences in effect size (HR) when comparing unadjusted Cox fixed effect regression models comparing lowest quartile vs. the other quartiles of the HRV parameters RMSSD or SDNN with models using heart rate corrected HRV parameters cvRMSSD or cvSDNN. Thus, correction for heart rate seems not to be mandatory for the prediction of mortality.

The last issue to discuss is the handling of the quality of measurement and **artifacts**. As discussed above, this tends to be a minor problem in short term measurements where artifacts are easily detected and an artifact-free window for HRV calculation can often be detected. In 24-hour measurements, artifacts due to movement as well as extra beats cannot be avoided and therefore have to be addressed. In the studies compared, artifact handling processes were described very differently. Some described it in detail, others did not publish anything about it.

The **mean follow-up time** showed only a small effect on LF in unadjusted analysis models of cardiac mortality. Thus, HRV is a valid predictor of mortality over medium to long time periods. Note that

short periods, order of magnitude of days, were not covered by the included studies as the minimum follow-up period was three months. Although effect sizes were not significantly different between short and long follow-up periods, it is evident from the graphs that studies with long follow up periods around 10 years had smaller CIs than studies covering only months.

Regarding the **different HRV parameters**, our results show similar HR magnitude across most extracted HRV parameters. This is not surprising, since they originate from the same source file of R-R intervals (Wittling and Wittling, 2012). However, the extent to which single parameters represent the ANS activity or vagal activity differs. For example, SDNN captures a wide range of physiological signals from different systems. Especially if HRV measures such as SDNN, but also SDANN, TP or VLF are drawn from 24h periods, they reflect measures of functional capacity rather than autonomic activity (Roach et al., 2004, 1998; Soares-Miranda et al., 2014). The latter is more accurately reflected in vagally-mediated parameters such as RMSSD, HF or LF (Camm et al., 1996; Shaffer and Ginsberg, 2017).

In addition, SDNN results from short term measures can vary largely due to nonstationarity, i.e. depending on the applied detrending method, while results of RMSSD appeared more stable (Tarvainen et al., 2002). This may result in biased estimates, but detrending methods are seldomly reported in the published manuscripts. In sum, for risk prediction, a 5-minute RMSSD measure represents best ANS function and is most robust to be calculated.

4.3. Potential neurobiological underpinnings

HRV represents the peripheral output of the central autonomic network (CAN) (Thayer et al., 2012). The CAN modulates on a moment-to-moment basis not only somatic responses to adapt to internal or external challenges, ultimately to maintain homeostasis (Benarroch, 2014; Bernard, 1867; Thayer and Lane, 2000; Wulsin et al., 2018) but also shapes emotional appraisal (i.e., emotional regulation) and behavioral adaption (Brosschot et al., 2017; Thayer et al., 2021; Thayer and Lane, 2000). Thus,

the function of the CAN is complex and defining an index can be described as measuring the capacity of the body to adapt to environmental challenges (Thayer et al., 2011) and its improper functioning has been suggested to accelerate aging processes and thus increase morbidity and mortality (Thayer et al., 2021). Here, measures of HRV may represent an index of vertical integration and adaption processes that shape brainstem activity and autonomic responses in the body. Therefore it is hardly surprising that HRV is considered to be an indicator for the state of the body and mind like in the case of a higher inflammatory state (Aeschbacher et al., 2017; Jarczok et al., 2014) or work stress (Jarczok et al., 2020, 2013), but also for the risk of disease and accelerated aging (Thayer et al., 2021). The idea that HRV expresses capacity to adapt is also supported by the findings that higher HRV predicts better symptom improvement after psychotherapy for common mental disorders (Balint et al., 2022b, 2022a).

The present results of the meta-analysis can be interpreted based on the concept of the Neurovisceral Integration Model, particularly on the central-autonomic moment-to-moment adaption of somatic responses and emotional appraisal to ultimately balance maintenance of homeostasis and immediate adaption to environmental stimuli (Benarroch, 2014; Thayer et al., 2021; Thayer and Lane, 2000; Wulsin et al., 2018). It has been suggested that healthy aging is associated with significant organ changes of the brain and the heart. Particularly, the regular aging associated functional change on the brains level (i.e. shift in the relative balance between prefrontal cortical thickness and amygdala volume) might be accelerated by continued exposure to stress (Thayer et al., 2021). In brief, ventromedial prefrontal cortex areas (vmPFC) tonically inhibit sub-cortical threat circuits. This path can reduce stress responses and fear behavior in a manner that depends on integrating the external context (e.g. environmental threat) with the internal one (e.g. perceptions of control over the threat) on a moment-to-moment basis. Under conditions of uncertainty or threat, these critical areas of the vmPFC rapidly become hypoactive and the so-called fight-or-flight response can unfold (Thayer et al., 2012). It has been suggested that this fight-or-flight response is actively inhibited by default due to perception of safety (Brosschot et al., 2017; O'Connor et al., 2021), unless

it becomes disinhibited due to the perception of danger (Thayer et al., 2021). Measures of HRV are suggested to index the extent of this vmPFC inhibitory function. With the present meta-analysis, we demonstrate lower values of HRV at baseline to be robustly associated to higher risk of all-cause mortality thus providing indirect support for this neurobiological foundation. These neurobiological and neurophysiological underpinnings of the core association between HRV as a measure of adaptive function and mortality is summarized in the neurovisceral integration model (Thayer et al., 2021; Thayer and Lane, 2009, 2000). Its neurological components have been summarized in a meta-analysis by Thayer et al. (2012).

4.4. Limitations

For clinical use, there is a need for a few and practical numbers like a cut-off for risk/no risk of RMSSD or an HR that applies for a 35-year-old man with a 5-minutes RMSSD of 25ms. Our aim was to calculate this from all studies gathered. Due to the enormous heterogeneity especially regarding the cut offs, this was not possible. We chose to calculate a composite HR at least for the studies reporting HRs for the lowest quartile. It is a limitation that this omits the information included in the other studies. A further limitation is that we selected the unadjusted models for this purpose. Since age and sex are already accounted for in this approach as described above in the discussion section, this seemed reasonable. Of course, part of the information included in HR relies on other variables besides age, and sex, such as preexistent disease, blood pressure and blood parameters like lipids. In a first attempt to guide clinicians, we want to offer here a method that does not need so much additional information. The next step would be to build a complex model like a risk calculator that incorporates more variables. Then, the HR from the covariate adjusted models should be used.

A limitation for the present interpretation of the results is the fact that most studies only report selective HRV parameters. In some cases, nonsignificant parameters were at least described to be not significantly enough to be associated with mortality. This limits the possibility to compare the different parameters across different studies, as many results are obviously not published. To

support further reviews and meta-analyses, for the future, we strongly encourage authors to report all calculated HRV parameters in general, even those without significant effects. This can be done easily in supplementary tables. We strongly recommend reporting a minimum set of HRV parameters such as SDNN, RMSSD, HF, LF, TP for 5-minute recordings and additionally VLF and SDANN for 24h recordings. For example, whereas almost N=29,000 individuals were included from k=14 studies in covariate adjusted SDNN, only 28% of these studies (k=4; N=4343) reported TP, thus representing a massive underreporting of some HRV parameters.

5. Conclusion

HRV represents a nonspecific predictor of mortality with a lower value corresponding to a higher risk of mortality. This association appeared independently of cardiac or all-cause mortality, clinical or non-clinical population, statistical adjustment for covariates, and HRV parameters used continuously or with cut-points. The effect size is comparable to that of masked uncontrolled hypertension. A possible clinical setting could be primary prevention. For future studies, we strongly recommend reporting a minimum set of HRV parameters such as SDNN, RMSSD, HF, LF, TP for 5-minute recordings and additionally VLF for 24h recordings.

6. Supplementary Material

Supplement S1-S4: Detailed forest plots with fixed effect models

S1: Forest plot for all-cause mortality from covariate unadjusted analyses (fixed effect models)

S2: Forest plot for all-cause mortality from covariate adjusted analyses (fixed effect models)

S3: Forest plot for cardiac mortality from covariate unadjusted analyses (fixed effect models)

S4: Forest plot for cardiac mortality from covariate adjusted analyses (fixed effect models)

Legend S1-S4:

Hazard Ratio >1 indicates higher mortality risk with lower HRV (per increment decrease/lowest quartile vs. other quartiles as indicated)

Weights and between-subgroup heterogeneity tests are from fixed effect models and with user-defined weight.

HRV parameters reported by less than two studies are not displayed.

Abbreviations: HRV: heart rate variability; N: number of participants; CI: confidence interval; I^2 :

heterogeneity; $p(I^2)$: P-value from heterogeneity test; SD: standard deviation; GenPop: General population; SDNN: Standard deviation of normal to normal intervals; RMSSD: Square root of the mean of the sum of the squares of differences between adjacent normal to normal intervals; HF-power: High frequency power; LF-power: Low frequency power; VLF-power: Very low frequency power; SDNN24: SDNN calculated from 24 hour measurements; SDANN: Standard deviation of the averages of normal to normal intervals; pNN50: Percentage of successive normal to normal intervals that differ by more than 50 ms; IBI: Inter-beat interval.

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7. Figure Captions

Figure 1: PRISMA flow chart of study selection

Legend:

AC mortality: all-cause mortality, CA mortality: cardiac mortality, IPD: individual participant data,
HRV: heart rate variability.

Figure 2:

Forest Plot for all-cause and cardiac mortality from unadjusted and covariate adjusted random effect
Cox-regression models.

Legend: figure 2

Hazard Ratio >1 indicates higher mortality risk with lower HRV (per increment decrease/lowest
quartile vs. other quartiles, for details of cut-offs see figure 3-6)

Dashes represent 1.000 participants.

HRV parameters reported by less than two studies are not displayed.

Abbreviations: HRV: heart rate variability, k: number of studies; n: number of participants; I^2 :

heterogeneity; $p(I^2)$: P-value from heterogeneity test, CI: confidence interval, all-cause: all-cause
mortality, cardiac: cardiac mortality, cov.-unadj: covariate unadjusted model, cov.-adj.:

covariate adjusted model, SDNN: Standard deviation of normal to normal intervals, RMSSD:

Square root of the mean of the sum of the squares of differences between adjacent normal to

normal intervals, HF-power: High frequency power, LF-power: Low frequency power, VLF-

power: Very low frequency power, SDNN24: SDNN calculated from 24 hour measurements,

SDANN: Standard deviation of the averages of normal to normal intervals, pNN50: Percentage

of successive normal to normal intervals that differ by more than 50 ms, IBI: Inter-beat interval

Figure 3: Forest plot for all-cause mortality from covariate unadjusted analyses

Legend figure 3

Hazard Ratio >1 indicates higher mortality risk with lower HRV (per increment decrease/lowest quartile vs. other quartiles, as indicated). Weights and between-subgroup heterogeneity tests are from random effect models and with user-defined weight.

Dashes represent 1.000 participants.

HRV parameters reported by less than two studies are not displayed.

Abbreviations: HRV: heart rate variability, N: number of participants, CI: confidence interval, I^2 :

heterogeneity; $p(I^2)$: P-value from heterogeneity test, all-cause: all-cause mortality, cardiac:

cardiac population, CAD: coronary artery disease, GenPop: General population, SD: standard

deviation, ms: millisecond, y: years, h: hours, min: minutes, SDNN: Standard deviation of normal

to normal intervals, RMSSD: Square root of the mean of the sum of the squares of differences

between adjacent normal to normal intervals, HF-power: High frequency power, LF-power: Low

frequency power, VLF-power: Very low frequency power, SDNN24: SDNN calculated from 24

hour measurements, SDANN: Standard deviation of the averages of normal to normal intervals,

pNN50: Percentage of successive normal to normal intervals that differ by more than 50 ms, IBI:

Inter-beat interval.

Figure 4: Forest plot for all-cause mortality from covariate adjusted analyses

Legend figure 4

Hazard Ratio >1 indicates higher mortality risk with lower HRV (per increment decrease/lowest quartile vs. other quartiles, as indicated). Weights and between-subgroup heterogeneity tests are from random effect models and with user-defined weight.

Dashes represent 1.000 participants.

HRV parameters reported by less than two studies are not displayed.

Abbreviations: HRV: heart rate variability, N: number of participants, CI: confidence interval, I^2 :

heterogeneity; $p(I^2)$: P-value from heterogeneity test, all-cause: all-cause mortality, cardiac: cardiac population, CAD: coronary artery disease, GenPop: General population, SD: standard deviation, ms: millisecond, y: years, h: hours, min: minutes, SDNN: Standard deviation of normal to normal intervals, RMSSD: Square root of the mean of the sum of the squares of differences between adjacent normal to normal intervals, HF-power: High frequency power, LF-power: Low frequency power, VLF-power: Very low frequency power, SDNN24: SDNN calculated from 24 hour measurements, SDANN: Standard deviation of the averages of normal to normal intervals, pNN50: Percentage of successive normal to normal intervals that differ by more than 50 ms, IBI: Inter-beat interval.

Figure 5: Forest plot for cardiac mortality from covariate unadjusted analyses

Legend figure 5:

Hazard Ratio >1 indicates higher mortality risk with lower HRV (per increment decrease/lowest quartile vs. other quartiles, as indicated). Weights and between-subgroup heterogeneity tests are from random effect models and with user-defined weight.

Dashes represent 1.000 participants.

HRV parameters reported by less than two studies are not displayed.

Abbreviations: HRV: heart rate variability, N: number of participants, CI: confidence interval, I^2 :

heterogeneity; $p(I^2)$: P-value from heterogeneity test, all-cause: all-cause mortality, cardiac: cardiac population, CAD: coronary artery disease, GenPop: General population, SD: standard deviation, ms: millisecond, y: years, h: hours, min: minutes, SDNN: Standard deviation of normal to normal intervals, RMSSD: Square root of the mean of the sum of the squares of differences between adjacent normal to normal intervals, HF-power: High frequency power, LF-power: Low frequency power, VLF-power: Very low frequency power, SDNN24: SDNN calculated from 24 hour measurements, SDANN: Standard deviation of the averages of normal to normal intervals, pNN50: Percentage of successive normal to normal intervals that differ by more than 50 ms, IBI: Inter-beat interval.

Figure 6: Forest plot for cardiac mortality from covariate adjusted analyses

Legend figure 6:

Hazard Ratio >1 indicates higher mortality risk with lower HRV (per increment decrease/lowest quartile vs. other quartiles, as indicated). Weights and between-subgroup heterogeneity tests are from random effect models and with user-defined weight.

Dashes represent 1.000 participants.

HRV parameters reported by less than two studies are not displayed.

Abbreviations: HRV: heart rate variability, N: number of participants, CI: confidence interval, I^2 :

heterogeneity; $p(I^2)$: P-value from heterogeneity test, all-cause: all-cause mortality, cardiac: cardiac population, CAD: coronary artery disease, GenPop: General population, SD: standard deviation, ms: millisecond, y: years, h: hours, min: minutes, SDNN: Standard deviation of normal to normal intervals, RMSSD: Square root of the mean of the sum of the squares of differences between adjacent normal to normal intervals, HF-power: High frequency power, LF-power: Low frequency power, VLF-power: Very low frequency power, SDNN24: SDNN calculated from 24 hour measurements, SDANN: Standard deviation of the averages of normal to normal intervals, pNN50: Percentage of successive normal to normal intervals that differ by more than 50 ms, IBI: Inter-beat interval.

Figure 7: Forest plot of hazard ratios for all-cause mortality from unadjusted analyses (fixed effect models): only studies using lowest quartile of RMSSD.

Legend Figure 7:

Hazard ratio >1 indicates higher mortality risk for lowest RMSSD quartile.

Weights and between-subgroup heterogeneity tests are from fixed effect models and with user-defined weight.

Abbreviations: RMSSD: squares of differences between adjacent R-R intervals, N: number of participants, CI: confidence interval, GenPop: General population, cardiac: cardiac population, y: years, I^2 : heterogeneity; p: P-value from heterogeneity test.

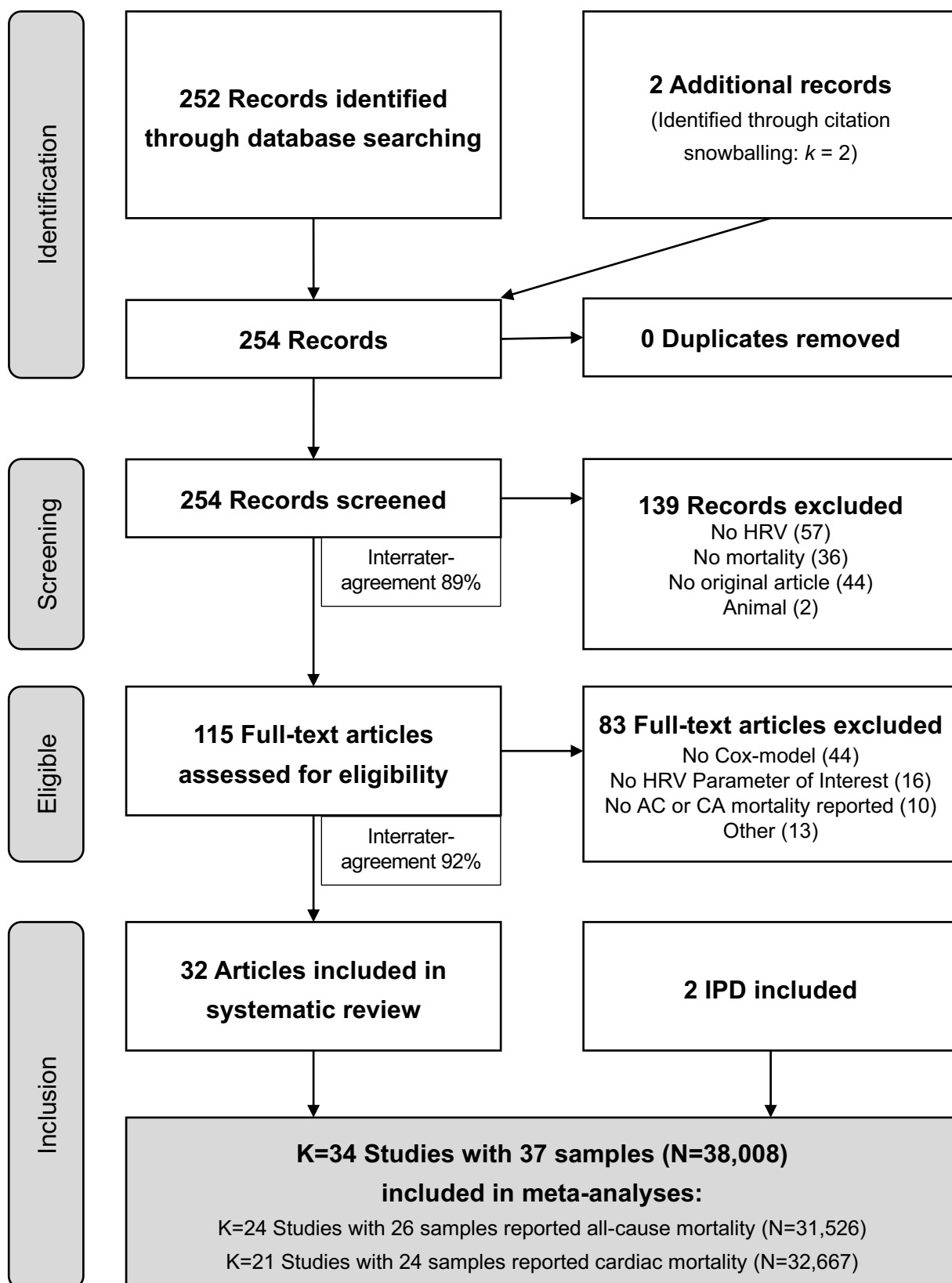
8. Tables

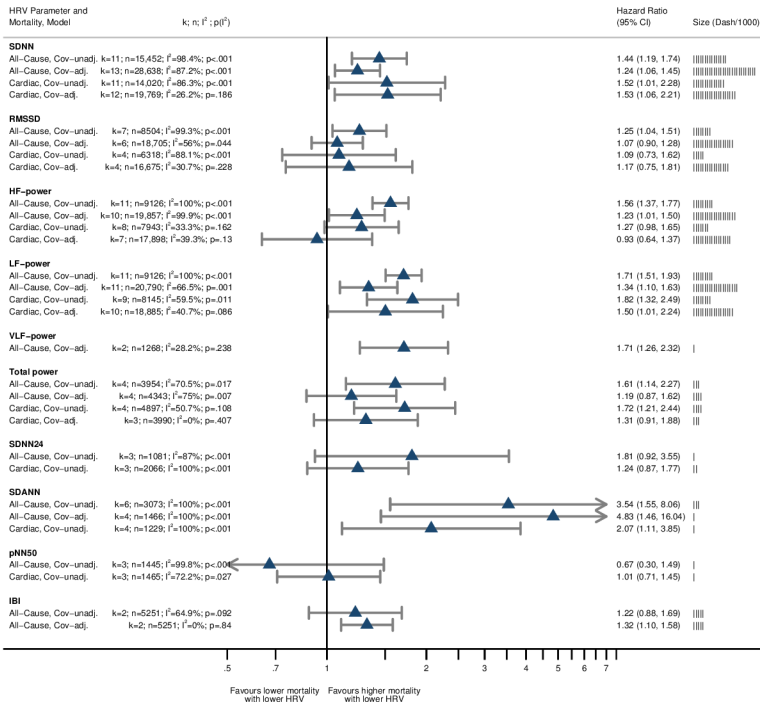
Table 1: Characteristics of included studies

Table 2: Quality and risk of bias rating of included studies

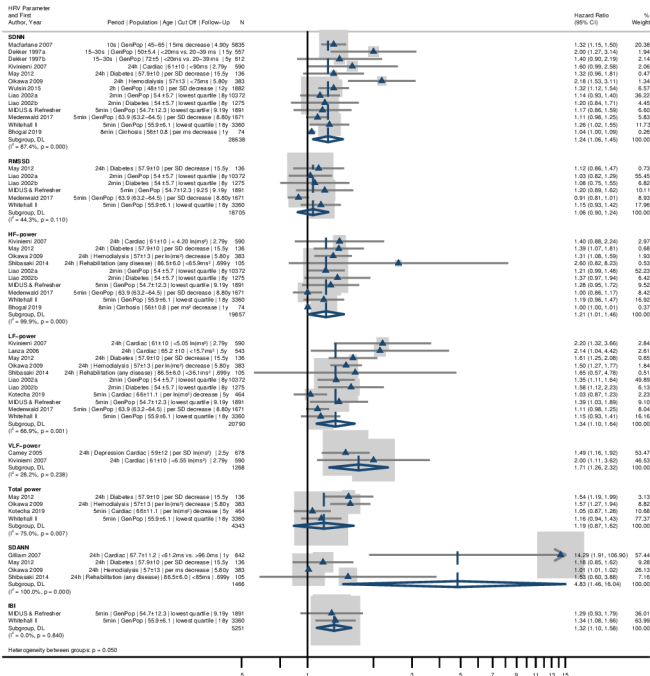
Table 3: Results of Meta-Analysis

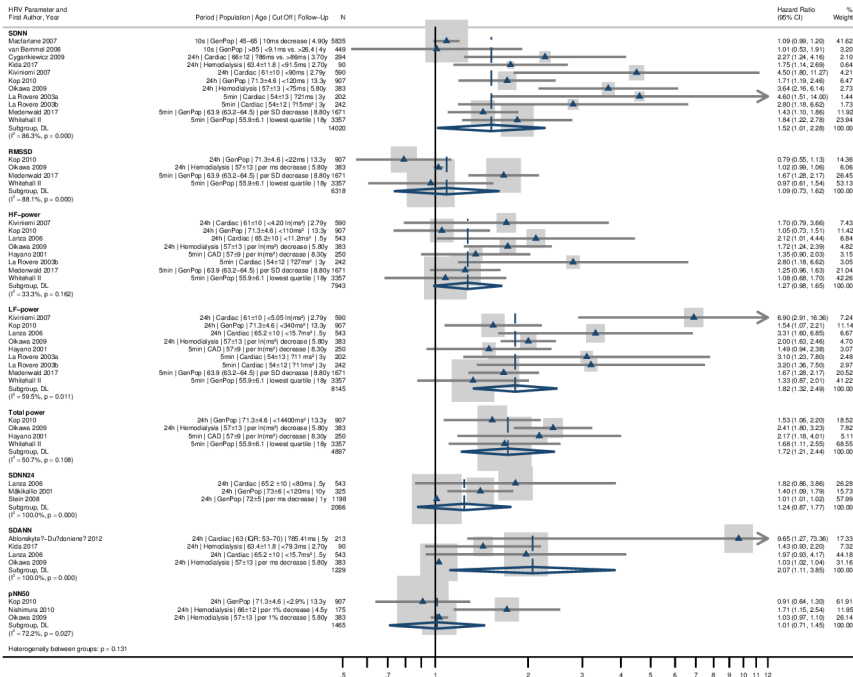
Table 4: Results of Meta-Regressions

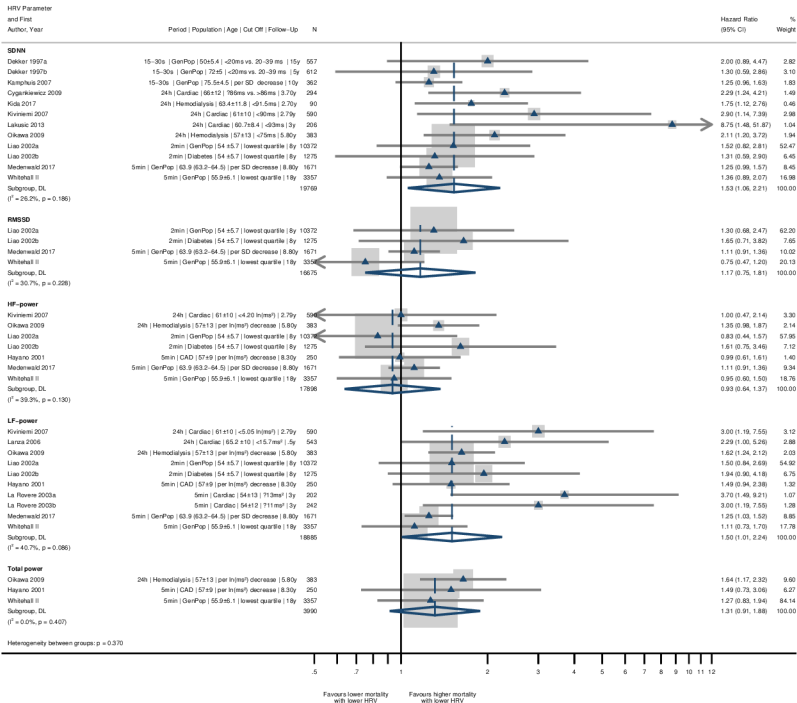




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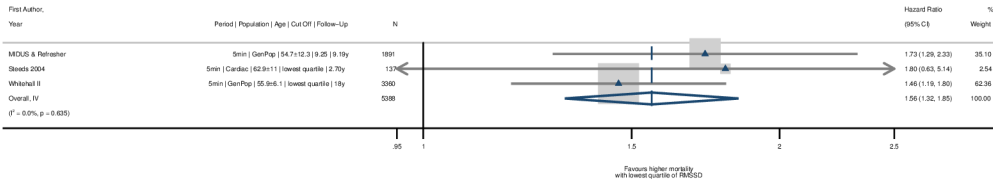


Table 1: Characteristics of included studies

		all-cause mortality; univariate analysis	all-cause mortality; multivariate analysis	cardiac mortality; univariate analysis	cardiac mortality; multivariate analysis			Period / Start of data collection	Proporti on Female (%)								
First author, year of publication	N					Study name	Country			Age (Mean±SD)	Type of Population	Follow up time	controlled variables for multivariate analysis	setting	respiration control	Total duration/analyzed segments (if differing from total duration)	algorithm for frequency domain
Ablonskytė-Dūdionienė et al., 2012	213	SDNN24		SDANN			Lithuania	2003-2009	70	63±13	STEMI Patients	5 years	none	ambulatory	none	24h/24h	n/a
Badarau et al., 2015	116		VLF				Romania	2011-2014	47	62±15	hemodialysis	17.5 months	model with pre-dialysis QRS interval	ambulatory	none	24h/n/a	n/a
Bhagal et al., 2019	74	SDNN, HF, LF, VLF	SDNN, HF, VLF				Italy	2009-2011	66	56 ± 11	Liver cirrhosis	12.3 months	Pugh score	resting	none	10min/8min	n/a
													depression, risk score (age, diabetes mellitus, left ventricular ejection fraction, creatinine level, prior MI, his- tory of pulmonary disease, prior transient ischemic attack or stroke, history of congestive heart failure, Killip class at time of index MI, and treatment with vasodilator)	ambulatory	none	24h/5min; 24h (VLF)	FFT
Carney et al., 2005	678		VLF			ENRICHED Study	USA	1997-2000	40	59±12	Depressed, post-MI	30 months		ambulatory	none	24h/5min; 24h (VLF)	FFT
													age, previous MI, HR>80/min, ventricular premature beats > 10/hour, nonsustained VT, QRS>120 ms, Turbulence onset >=0%, Turbulence slope <=2,5ms/RR, Daytime QT end/RR>0,21	ambulatory	none	24h	n/a
Cygankiewicz et al., 2009	294			SDNN	SDNN	Muerte Subita en Insuficiencia Cardiaca (Sudden Death in Heart Failure [MUSIC]) Study	Spain	2003-2004	32	66±12	heart failure with LVEF > 35% and NYHA II-III	44 months	age	ambulatory	none	15-30 s	n/a
Dekker et al., 1997 A	557		SDNN		SDNN	Zutphen Study (Seven countries study)	Netherlands	1960; 1965; 1970; 1985	0	50±5	General Population	15 years	age	resting	none	15-30 s	n/a
Dekker et al., 1997 B	612		SDNN		SDNN	Zutphen Study (Seven countries study)	Netherlands	1960; 1965; 1970; 1985	0	72±5	General Population	5 years	age	resting	none	15-30 s	n/a
											heart failure NYHA III-IV with cardiac resynchronization therapy					constant recording / 24h	n/a
Gilliam et al., 2007	842	SDANN	SDANN			Heart Failure Heart Rate Variability Registry HF- HRV	USA	2003-2005	24	68±11		11.6 months	age, sex, BMI, DBP	ambulatory	none	24h	n/a
Hayano et al., 2001	250			HF, LF, TP	HF, LF, TP		Japan	1987-1991	28	57±9	CAD	99±23 months	none	supine	15BPM	5min	n/a
Hotta 2005	298	pNNS0, SDNN, RMSSD, LF, HF				Longitudinal Investigation for the Longevity and Aging in Hokkaido County (LILAC) study	Japan	2000-2004	28	80	General Population	4 years	none	supine	none	60min/5min	n/a
Kamphuis et al., 2007	362				SDNN	The Finland, Italy and The Netherlands Elderly (FINE) Study	Netherlands	1989-1991; 2000	0	76±5	healthy (no CVD, no diabetes)	10 years	age	resting	none	15-30s	n/a
Kida et al., 2017	90	SDANN, SDNN	SDNN	SDNN, HF, LF, VLF	SDNN, HF, LF, VLF	multiple risk factor analysis trial (MRFAT)	Japan	2009-2011	29	63±12	hemodialysis	32±31 months	none	ambulatory	none	24h/5min	n/a
Kiviniemi et al., 2007	590	SDNN, HF, LF, VLF	SDNN, HF, LF, VLF	SDNN, HF, LF, VLF	SDNN, HF, LF, VLF		Finland	1996	24	61±10	Post MI	34±14 months	none	ambulatory	none	24h/5min; VLF: 24h	n/a
																24h/5min (HF, LF); 24 h (TP); n/a (SDNN, RMSSD, pNNS0)	n/a
Kop et al., 2010	907			RMSSD, SDNN, pNNS0, HF, LF, VLF, TP		Cardiovascular Health Study (CHS)	USA	1989-1990	59	71±5	Medicare-eligible individuals	13.3 years	none	ambulatory	none		n/a
Kotecha et al., 2019	464		SDNN, LF, TP			The Alternative Risk Markers in Coronary Artery Disease (ARM-CAD) study	Australia	2006-2008	33	66 ± 11	CAD	5 years	age, sex, prior MI, diabetes, smoking, presence of impaired LVF, occlusive CAD, revascularisation, total cholesterol, SBP, BNP	supine	none	5min	n/a
La Rovere et al., 2003 Derivation	202			SDNN, LF	LF		Italy	1991-1995	13	54±13	CHF		none	supine	15BPM	8min/5min	n/a
La Rovere et al., 2003 Validation	242			SDNN, LF, HF	LF		Italy	1996-2001	17	54±12	CHF		none	supine	15BPM	8min/5min	n/a
													age, gender, BMI, number of bypass grafts, type of operation (off/on pump), hypertension, dyslipidemia, diabetes mellitus, previous myocardial infarction, EF before and after operation, functional capacity after operation and time from operation to measuring HRV				
Lakusic et al., 2013	206				SDNN		Croatia	n/a	31	61±8	after CABG	3.0±1.8 years	age, diabetes, worsening UA, ST depression, troponin >0,4ng/ml, ≥10 extrasystoles/h	ambulatory	none	23-2h (range 21-24 h)	n/a
Lanza et al., 2006	543	SDNN24, SDANN, HF, LF, VLF	LF	SDNN24, SDANN, HF, LF, VLF	LF	SPAI (Stratificazione Prognostica dell'Angina Instabile) study	Italy	1997-2001	34	65 ±10	Instable Angina	6 months	age, sex, ethnicity- center, cigaret, triglyceride, BMI, fasting glucose, HDL, hypertension, smoking, and mean heart rate.	ambulatory	none	24h/5min (time domain); 10 min (frequency domain)	FFT
Liao et al., 2002 A	10372		RMSSD, SDNN, HF, LF		RMSSD, SDNN, HF, LF	The Atherosclerosis Risk in Communities (ARIC) Study	USA	1987-1989	58	54 ± 5.7	General Population	8 years	age, sex, ethnicity- center, smoking, and mean heart rate.	supine	none	2min	n/a
Liao et al., 2002 B	1275		RMSSD, SDNN, HF, LF		RMSSD, SDNN, HF, LF	The Atherosclerosis Risk in Communities (ARIC) Study	USA	1987-1989	58	54 ± 5.7	General Population	8 years	age, sex, ethnicity- center, smoking, and mean heart rate.	supine	none	2min	n/a
Macfarlane et al., 2007	5835	SDNN	SDNN	SDNN		West of Scotland Coronary Prevention Study (WOSCOPS)	Scotland	1989 -1995	0	45-65	moderate hypercholesterinemia (LDL)	4.9 years	age, smoking, medical history, lipids, other biochemical measures and ECG variates	supine	none	10s	n/a
Mäkilä et al., 2001	325	SDNN24	SDNN24	SDNN24	SDNN24	elderly in the city of Turku, Finland	Finland	1982-1992	47	73±6	General Population	10 years	age, sex, heart failure, angina pectoris, functional class, previous MI, cardiac medication, ventricular premature beats.	ambulatory	none	24h	n/a
May & Arildsen, 2012	136		RMSSD, SDANN, SDNN, LF, HF, VLF, TP			Horsens, Denmark, 1993-1994	Denmark	1993-1994	40	58±10	Diabetic	15.5 years	age, sex	ambulatory	none	24h/5min	FFT
													age, sex, MetS, history of MI, sport index, plasma level of NT-pro brain natriuretic peptide, HbA1c, LDL, TSH, systolic and diastolic blood pressure, the estimated glomerular filtration rate, regular intake of anti-arrhythmic medication, beta-blockers, digitalis glycosides, glucocorticoids, antiphlogistic medication, thyroid medication, and self-reported malignant neoplasms, thyroid disease, or rheumatic diseases				
Medenwald et al., 2017	1671	RMSSD, SDNN, HF, LF	RMSSD, SDNN, HF, LF	RMSSD, SDNN, HF, LF	RMSSD, SDNN, HF, LF	CARDiovascular Living and Ageing in Halle study (CARLA study)	Germany	2002-2006	47	64±10	General Population	8.8 years	age, LF/HF ratio, diabetes, DBP, Triglycerides	sitting	15BPM	20min/5min	FFT
Nishimura et al., 2010	175			pNNS0	pNNS0		Japan	2000-2002	40	66±12	Hemodialysis patients with left ventricular hypertrophy	4.5±1.9 years		ambulatory	none	24h	n/a
Oikawa et al., 2009	383	RMSSD, pNNS0, SDANN, SDNN, HF, LF, TP	SDANN, SDNN, HF, LF, TP	RMSSD, pNNS0, SDANN, SDNN, HF, LF, TP	SDANN, SDNN, HF, LF, TP		Japan	1997-1999	43	57±13	Chronic hemodialysis patients	5.78±2.47 years	age, LVEF, female gender, diabetes	ambulatory	none	24h	n/a
Quintana et al., 1997	74	HF, LF, VLF, TP					Sweden	n/a	22	61±9	acute MI hospitalisation	36±15 months	none	ambulatory	none	23±2h / per 500RR intervall segments	FFT
Shibasaki et al., 2014	105	HF, LF, SDANN	SDANN, LF, HF				Japan	2007-2011	75	87±6	older adults > 75 years in long term care rehabilitation patients who were implanted with a CONTAK RENEWAL biventricular defibrillator or biventricular pacemaker device	8.9 months	age, sex and cardiovascular risk factors	ambulatory	none	24h/5min	FFT
Singh et al., 2009	436	SDANN				Cardiac Resynchronization Therapy Registry Evaluating Patient Response with RENEWAL Family Devices (CRT RENEWAL)	USA	2004-2007	31	67±12		3 months	none	ambulatory	none	24h/5min	n/a
Steeds et al., 2004	137	RMSSD, SDNN, HF, LF, TP					GB	1998-2000	70	63±11	acute MI	32 months	none	supine	none	256 beats	n/a
Stein et al., 2008	1198			SDNN24		CAST cardiac arrhythmia suppression trial	USA	1989- 1990 + 1992-1993	55	72±5	Cardiovascular Health Study (CHS) cohort, ≥65 years	median of more than	none	ambulatory	none	24h	n/a
Syed et al., 2009	764			pNNS0, RMSSD, SDANN, SDNN		DISPERSE2 (TIMI 33) trial	("14 countries"*)	(10/2004 - 04/2005*)	37	62.3 (50-78)	Patients within 48h of initial event (non-ST-elevation acute coronary syndromes)	3 months	age, diabetes, History of CVD, Morphologic variability	ambulatory	none	24h/5min	"according to Task Force"
van Bemmel et al., 2006	449	SDNN		SDNN	SDNN		Netherlands	1997-1999	66	85	General Population	4 years	none	resting	none	10s	n/a
Wulsin et al., 2015	1882				SDNN	FHS Offspring Cohort	USA	inclusion: 1983-1987	52	48±10	General Population	12 years	sex, age, smoking, HRV x sex	ambulatory	none	2h	n/a
MIDUS II & Refresher	1891	RMSSD, SDNN, HF, LF, IBI	RMSSD, SDNN, HF, LF, IBI			Midlife in the US (MIDUS 2)	USA	2004-2014	56	55±12	General Population	9.5 years	age, sex	resting	none	10min/5min	FFT
Whitehall II	3360	RMSSD, SDNN, HF, LF, TP, IBI	RMSSD, SDNN, HF, LF, TP, IBI	RMSSD, SDNN, HF, LF, TP	RMSSD, SDNN, HF, LF, TP	Stress and Health Study	GB	1997-1999	30	56±6	General Population	18 years	age, sex	resting	none	5min	AR

abbreviations:

CABG coronary artery bypass graft
CAD coronary artery disease
CHF chronic heart failure
CVD cardiovascular disease
HF-power High frequency power
IBI Inter-beat interval
LDL low-density lipoprotein
LF-power low frequency power
LVEF left ventricular ejection fraction
MI myocardial infarction
n/a Not Available
NYHA class New York Heart Association class of heart failure
pNNS0 Percentage of successive normal to normal intervals that differ by more than 50 ms
RMSSD Square root of the mean of the sum of the squares of differences between adjacent normal to normal intervals
SDANN Standard deviation of the averages of normal to normal interval
SDNN Standard deviation of normal to normal intervals
SDNN24 SDNN calculated from 24 hour measurements
STEMI ST-elevation myocardial infarction
VLF-power Very low frequency power

Table 2: Quality and Risk of Bias Rating of Included Studies

First Author, Year	Time Window defined	Quality of Artifact Control	HRV Parameter Selection	Population Bias	Total
Ablonskytė-Dūdonienė 2012	1	1	0	1	3
Badarau 2015	1	0	0	0	1
Bhogal 2019	1	1	0	1	3
Carney 2005	0	0	0	1	1
Cygankiewicz 2009	1	0	1	1	3
Dekker 1997	0	1	1	1	3
Gilliam 2007	1	0	0	0	1
Hayano 2001	1	1	1	1	4
Hotta 2005	0	0	0	1	1
Kamphuis 2007	0	1	1	1	3
Kida 2017	1	0	0	0	1
Kiviniemi 2007	0	0	1	0	1
Kop 2010	0	0	1	1	2
Kotecha 2019	1	0	0	1	2
La Rovere 2003	1	1	0	1	3
Lakusic 2013	1	1	0	0	2
Lanza 2006	1	0	0	1	2
Liao 2002	1	1	0	1	3
Macfarlane 2007	1	0	1	0	2
Mäkikallio 2001	1	0	0	1	2
May 2012	1	0	0	1	2
Medenwald 2017	1	1	1	1	4
MIDUS & Refresher	1	1	1	1	4
Nishimura 2010	1	0	0	1	2
Oikawa 2009	1	1	0	1	3
Quintana 1997	1	0	0	1	2
Shibasaki 2014	1	0	0	0	1
Singh 2009	1	0	0	1	2
Steeds 2004	0	0	0	0	0
Stein 2008	1	0	0	0	1
Syed 2009	1	0	0	1	2
van Bommel 2006	1	1	1	1	4
Whitehall II	1	0	1	1	3
Wulsin 2015	1	0	0	0	1

0: high risk of bias present

1: no or low risk of bias present

Table 3: Results of Meta-Analysis

				Fixed effect						Random effect					Heterogeneit			
		HRV parameter	k	n	Effect size	lower 95%CI	upper 95%CI	z	p#	Effect size	lower 95%CI	upper 95%CI	z	p#	y statistic	I ² **	p\$	Tau ²
All-cause mortality	covariate unadjusted	SDNN	11	15452	1.441	1.294	1.605	6.632	0.000	1.441	1.191	1.744	3.755	<0.001	629.71	98.4%	<0.001	0.029
		SDNN24	3	1081	1.808	1.192	2.743	2.786	0.005	1.808	0.921	3.552	1.72	0.085	15.34	87.0%	<0.001	0.192
		SDANN	6	3073	3.538	1.896	6.604	3.97	0.000	3.538	1.554	8.058	3.01	0.003	190000	100%	<0.001	0.366
		RMSSD	7	8504	1.255	1.047	1.504	2.46	0.014	1.255	1.040	1.514	2.372	0.018	915.62	99.3%	<0.001	0.002
		pNN50	3	1445	0.671	0.303	1.487	-0.984	0.325	0.671	0.303	1.487	-0.984	0.325	916.66	99.8%	<0.001	0.000
		HF	11	9126	1.558	1.375	1.764	6.97	0.000	1.558	1.374	1.765	6.939	<0.001	20000000	100%	<0.001	0.000
		LF	11	9126	1.706	1.506	1.932	8.41	0.000	1.706	1.505	1.934	8.351	<0.001	29000000	100%	<0.001	0.000
		TP	4	3954	1.611	1.344	1.931	5.151	0.000	1.611	1.142	2.272	2.717	0.007	10.17	70.5%	0.017	0.030
	covariate adjusted	SDNN	13	28963	1.236	1.130	1.353	4.603	0.000	1.236	1.065	1.435	2.792	0.005	94.57	86.3%	<0.001	0.018
		SDANN	4	1466	4.834	1.518	15.395	2.666	0.008	4.834	1.457	16.042	2.575	0.010	260000	100%	<0.001	0.061
		RMSSD	6	18705	1.075	0.935	1.235	1.016	0.310	1.075	0.899	1.285	0.793	0.428	11.38	56.0%	0.044	0.009
		HF	10	19857	1.232	1.092	1.390	3.390	0.001	1.232	1.015	1.496	2.107	0.035	13173.15	99.9%	<0.001	0.019
		LF	11	20790	1.338	1.194	1.499	5.000	0.000	1.338	1.095	1.634	2.851	0.004	29.82	66.5%	0.001	0.024
		VLF	2	1268	1.710	1.258	2.325	3.428	0.001	1.710	1.258	2.325	3.428	0.001	1.39	28.2%	0.238	0.000
		TP	4	4343	1.186	1.005	1.401	2.018	0.044	1.186	0.868	1.622	1.071	0.284	12.02	75.0%	0.007	0.030
Cardiac mortality	covariate unadjusted	SDNN	11	14020	1.519	1.340	1.721	6.545	0.000	1.519	1.013	2.278	2.022	0.043	72.73	86.3%	0.043	0.1525
		SDNN24	3	2066	1.241	1.015	1.518	2.107	0.035	1.241	0.873	1.766	1.202	0.229	6463.78	100%	0.229	0.0507
		SDANN	4	1229	2.068	1.275	3.356	2.944	0.003	2.068	1.111	3.850	2.293	0.022	19836.30	100%	0.022	0.1207
		RMSSD	4	6318	1.088	0.836	1.416	0.629	0.529	1.088	0.733	1.616	0.419	0.675	25.29	88.1%	0.675	0.0601
		pNN50	3	1465	1.013	0.809	1.269	0.114	0.909	1.013	0.708	1.451	0.072	0.943	7.19	72.2%	0.943	0.0436
		HF	8	7943	1.274	1.021	1.588	2.149	0.032	1.274	0.984	1.649	1.836	0.066	10.50	33.3%	0.066	0.0187
		LF	9	8145	1.816	1.480	2.228	5.715	0.000	1.816	1.322	2.492	3.688	<0.001	19.73	59.5%	<0.001	0.0639
		TP	4	4897	1.719	1.278	2.312	3.584	0.000	1.719	1.209	2.444	3.019	0.003	6.08	50.7%	0.003	0.0182
	covariate adjusted	SDNN	11	20094	1.515	1.086	2.113	2.445	0.014	1.515	1.018	2.253	2.049	0.040	23.38	48.7%	0.025	0.040
		RMSSD	4	16675	1.167	0.770	1.770	0.727	0.467	1.167	0.751	1.814	0.687	0.492	4.33	30.7%	0.228	0.012
		HF	7	17898	0.934	0.636	1.371	-0.348	0.728	0.934	0.636	1.371	-0.348	0.728	9.88	39.3%	0.130	0.000
		LF	10	18885	1.503	1.074	2.104	2.375	0.018	1.503	1.008	2.243	1.997	0.046	15.17	40.7%	0.086	0.035
		TP	3	3990	1.312	0.914	1.883	1.474	0.140	1.312	0.914	1.883	1.474	0.140	1.80	0.0%	0.407	0.000

Legend:

abbreviations:

- Tau² between-study variance
- CI confidence interval
- HF High frequency power
- HRV heart rate variability
- I²** variation in effect size attributable to heterogeneity
- k number of studies
- LF Low frequency power
- p# p-value of significance test of effect size=1
- p\$ p-value of I² statistic
- pNN50 Percentage of successive normal to normal intervals that differ by more than 50 ms
- RMSSD Square root of the mean of the sum of the squares of differences between adjacent normal to normal intervals
- SDANN Standard deviation of the averages of normal to normal intervals
- SDNN Standard deviation of normal to normal intervals
- SDNN24 SDNN calculated from 24 hour measurements
- TP Total power
- VLF Very low frequency power
- z z statistics

Table 4: Results of Meta-Regressions

		HRV parameter	Effect modifier	k	b	SE	95%CI lower bound	95%CI upper bound	z	p	B	R²
All-cause mortality	Covariate adjusted studies	HF	Mean age (Years)	10	-0.001	0.027	-0.054	0.052	-0.023	0.981	-0.022	0.001
			Proportion female (%)	10	0.002	0.009	-0.015	0.019	0.209	0.834	0.192	0.037
			Recording length (minutes)	10	0.000	0.000	-0.001	0.001	0.402	0.688	0.370	0.137
			Continent (Asia vs. rest)	2/8	0.214	0.646	-1.052	1.479	0.331	0.741	0.304	0.093
			Startdecade (pre- vs. post 2000)	6/4	0.001	0.013	-0.024	0.026	0.074	0.941	0.068	0.005
			Quality of artifact-control (no / yes)	4/6	-0.013	0.245	-0.493	0.468	-0.052	0.958	-0.048	0.002
			Population (clinical vs. non-clinical)	6/4	-0.141	0.298	-0.725	0.444	-0.471	0.638	-0.434	0.188
			Shortterm-recording (no/yes)	4/6	-0.170	0.417	-0.988	0.648	-0.407	0.684	-0.374	0.140
			mean follow-up time (months)	10	-0.001	0.002	-0.005	0.004	-0.243	0.808	-0.223	0.050
		LF	Mean age (Years)	11	0.001	0.024	-0.046	0.048	0.028	0.978	0.018	0.000
			Proportion female (%)	11	0.001	0.008	-0.015	0.018	0.169	0.866	0.110	0.012
			Recording length (minutes)	11	0.000	0.000	0.000	0.001	1.099	0.272	0.721	0.519
			Continent (Asia vs. rest)	2/9	0.138	0.661	-1.157	1.432	0.209	0.835	0.137	0.019
			Startdecade (pre- vs. post 2000)	7/4	-0.006	0.013	-0.030	0.019	-0.435	0.663	-0.285	0.081
			Quality of artifact-control (no / yes)	6/5	0.010	0.231	-0.443	0.463	0.043	0.966	0.028	0.001
			Population (clinical vs. non-clinical)	7/4	-0.251	0.267	-0.775	0.273	-0.940	0.347	-0.616	0.380
			Shortterm-recording (no/yes)	5/6	-0.398	0.360	-1.102	0.307	-1.106	0.269	-0.725	0.526
			mean follow-up time (months)	11	-0.002	0.002	-0.006	0.002	-0.975	0.330	-0.639	0.408
		SDNN	Mean age (Years)	13	0.000	0.025	-0.050	0.049	-0.016	0.987	-0.013	0.000
			Proportion female (%)	13	-0.003	0.004	-0.011	0.005	-0.721	0.471	-0.585	0.342
			Recording length (minutes)	13	0.000	0.000	-0.001	0.001	0.692	0.489	0.561	0.315
			Continent (Asia vs. rest)	1/12	0.574	0.871	-1.132	2.280	0.659	0.510	0.535	0.286
			Startdecade (pre- vs. post 2000)	10/3	-0.005	0.013	-0.030	0.021	-0.361	0.718	-0.292	0.086
			Quality of artifact-control (no / yes)	5/8	-0.099	0.203	-0.497	0.299	-0.487	0.627	-0.395	0.156
			Population (clinical vs. non-clinical)	5/8	-0.144	0.357	-0.844	0.556	-0.404	0.686	-0.328	0.107
			Shortterm-recording (no/yes)	3/10	-0.353	0.518	-1.369	0.663	-0.682	0.496	-0.553	0.305
			mean follow-up time (months)	13	0.000	0.002	-0.004	0.004	0.010	0.992	0.008	0.000
	Covariate unadjusted studies	HF	Mean age (Years)	11	-0.016	0.018	-0.051	0.019	-0.912	0.362	-0.352	0.124
			Proportion female (%)	11	0.009	0.008	-0.007	0.025	1.110	0.267	0.392	0.153
			Recording length (minutes)	11	0.000	0.000	0.000	0.001	0.722	0.470	0.290	0.084
			Continent (Asia vs. rest)	3/8	-0.158	0.380	-0.903	0.586	-0.416	0.677	-0.170	0.029
			Startdecade (pre- vs. post 2000)	6/5	0.002	0.025	-0.046	0.050	0.077	0.938	0.031	0.001
			Quality of artifact-control (no / yes)	7/4	0.082	0.241	-0.390	0.554	0.340	0.734	0.135	0.018
			Population (clinical vs. non-clinical)	7/4	-0.181	0.272	-0.713	0.352	-0.665	0.506	-0.267	0.072
			Shortterm-recording (no/yes)	5/6	-0.210	0.283	-0.764	0.345	-0.740	0.459	-0.297	0.088
			mean follow-up time (months)	11	-0.001	0.002	-0.004	0.002	-0.718	0.473	-0.278	0.077
		LF	Mean age (Years)	11	-0.011	0.020	-0.049	0.027	-0.564	0.573	-0.183	0.033
			Proportion female (%)	11	-0.007	0.011	-0.028	0.015	-0.614	0.539	-0.207	0.043
			Recording length (minutes)	11	0.000	0.000	0.000	0.001	2.242	0.025	0.626	0.392
			Continent (Asia vs. rest)	3/8	-0.284	0.399	-1.066	0.499	-0.711	0.477	-0.230	0.053
			Startdecade (pre- vs. post 2000)	6/5	-0.012	0.029	-0.069	0.045	-0.418	0.676	-0.138	0.019
			Quality of artifact-control (no / yes)	7/4	0.025	0.277	-0.518	0.568	0.089	0.929	0.029	0.001
			Population (clinical vs. non-clinical)	7/4	-0.418	0.264	-0.935	0.099	-1.587	0.113	-0.455	0.207
			Shortterm-recording (no/yes)	5/6	-0.606	0.269	-1.134	-0.078	-2.251	0.024	-0.629	0.395
			mean follow-up time (months)	11	-0.003	0.001	-0.006	0.000	-2.024	0.043	-0.532	0.283
		SDNN	Mean age (Years)	11	-0.004	0.016	-0.036	0.028	-0.250	0.802	-0.099	0.010
			Proportion female (%)	11	0.008	0.005	-0.001	0.017	1.761	0.078	0.643	0.414
			Recording length (minutes)	11	0.000	0.000	0.000	0.001	1.223	0.221	0.451	0.204
			Continent (Asia vs. rest)	2/9	-0.204	0.092	-0.383	-0.024	-2.223	0.026	-0.812	0.659
			Startdecade (pre- vs. post 2000)	6/5	0.274	0.492	-0.690	1.238	0.557	0.578	0.213	0.046
			Quality of artifact-control (no / yes)	6/5	0.017	0.012	-0.007	0.040	1.362	0.173	0.497	0.247
			Population (clinical vs. non-clinical)	5/6	0.200	0.221	-0.233	0.632	0.905	0.365	0.331	0.109
			Shortterm-recording (no/yes)	3/8	-0.388	0.318	-1.011	0.236	-1.220	0.223	-0.451	0.203
			mean follow-up time (months)	11	0.001	0.002	-0.002	0.004	0.926	0.354	0.338	0.114
Cardiac mortality	Covariate adjusted studies	LF	Mean age (Years)	10	0.002	0.030	-0.057	0.060	0.056	0.956	0.023	0.001
			Proportion female (%)	10	-0.001	0.008	-0.016	0.014	-0.120	0.905	-0.050	0.003
			Recording length (minutes)	10	0.000	0.000	0.000	0.001	1.291	0.197	0.534	0.286
			Continent (Asia vs. rest)	2/8	0.045	0.556	-1.044	1.134	0.081	0.935	0.034	0.001
			Startdecade (pre- vs. post 2000)	9/1	-0.008	0.015	-0.038	0.023	-0.494	0.621	-0.205	0.042
			Quality of artifact-control (no / yes)	3/7	0.108	0.235	-0.352	0.569	0.461	0.645	0.191	0.036
			Population (clinical vs. non-clinical)	7/3	-0.471	0.258	-0.976	0.034	-1.827	0.068	-0.756	0.572
			Shortterm-recording (no/yes)	3/7	-0.477	0.368	-1.199	0.244	-1.297	0.195	-0.537	0.289
			mean follow-up time (months)	10	-0.004	0.002	-0.007	0.000	-1.924	0.054	-0.797	0.635
		SDNN	Mean age (Years)	12	-0.002	0.020	-0.040	0.036	-0.101	0.920	-0.042	0.002
			Proportion female (%)	12	-0.002	0.006	-0.013	0.009	-0.342	0.732	-0.140	0.020
			Recording length (minutes)	12	0.001	0.000	0.000	0.001	1.856	0.063	0.762	0.580
			Continent (Asia vs. rest)	2/10	0.297	0.654	-0.986	1.579	0.453	0.650	0.186	0.035
			Startdecade (pre- vs. post 2000)	8/4	0.001	0.013	-0.024	0.026	0.079	0.937	0.032	0.001
			Quality of artifact-control (no / yes)	4/8	-0.035	0.242	-0.509	0.439	-0.146	0.884	-0.060	0.004
			Population (clinical vs. non-clinical)	6/6	-0.328	0.285	-0.887	0.231	-1.151	0.250	-0.472	0.223
			Shortterm-recording (no/yes)	5/7	-0.690	0.371	-1.417	0.036	-1.863	0.063	-0.764	0.584
			mean follow-up time (months)	12	-0.002	0.002	-0.006	0.002	-0.907	0.364	-0.372	0.139
	Covariate unadjusted studies	LF	Mean age (Years)	9	-0.005	0.034	-0.071	0.061	-0.155	0.877	-0.055	0.003
			Proportion female (%)	9	-0.019	0.013	-0.044	0.006	-1.502	0.133	-0.489	0.239
			Recording length (minutes)	9	0.000	0.000	0.000	0.001	1.727	0.084	0.497	0.247
			Continent (Asia vs. rest)	2/7	-0.264	0.504	-1.252	0.724	-0.524	0.600	-0.186	0.035
			Startdecade (pre- vs. post 2000)	8/1	0.008	0.031	-0.052	0.068	0.273	0.785	0.097	0.009
			Quality of artifact-control (no / yes)	4/5	-0.137	0.371	-0.866	0.591	-0.370	0.711	-0.131	0.017
			Population (clinical vs. non-clinical)	6/3	-0.837	0.225	-1.278	-0.396	-3.722	0.000	-0.800	0.639
			Shortterm-recording (no/yes)	4/5	-0.519	0.302	-1.110	0.072	-1.722	0.085	-0.496	0.246
			mean follow-up time (months)	9	-0.005	0.001	-0.008	-0.003	-3.816	0.000	-0.820	0.672
	SDNN	LF	Mean age (Years)	11	-0.012	0.019	-0.050	0.026	-0.613	0.540	-0.208	0.043
			Proportion female (%)	11	0.005	0.006	-0.006	0.016	0.856	0.392	0.246	0.060
			Recording length (minutes)	11	0.000	0.000	0.000	0.001	2.088	0.037	0.587	0.344
			Continent (Asia vs. rest)	2/9	0.649	0.593	-0.512	1.811	1.096	0.273	0.341	0.116
			Startdecade (pre- vs. post 2000)	8/3	0.023	0.016	-0.008	0.054	1.450	0.147	0.397	0.157
			Quality of artifact-control (no / yes)	6/5	0.105	0.318	-0.518	0.727	0.330	0.742	0.107	0.011
			Population (clinical vs. non-clinical)	6/5	-0.923	0.309	-1.528	-0.319	-2.992	0.003	-0.804	0.646
			Shortterm-recording (no/yes)	5/6	-0.610	0.294	-1.186	-0.033	-2.074	0.038	-0.584	0.341
			mean follow-up time (months)	11	-0.001	0.002	-0.005	0.004	-0.221	0.825	-0.074	0.005

Legend: k: number of studies; b: beta coefficient; SE: standard error; CI: confidence interval; z: z statistics; p: p-value of significance; B: standardized beta coefficient; HRV: heart rate variability; HF: high frequency power; LF: Significant p-values (p<.05) are marked in bold.

low frequency power: SDNN: Standard deviation of normal to normal intervals