

Original Paper

# Sociodemographic, Health and Lifestyle, Sampling, and Mental Health Determinants of 24-Hour Motor Activity Patterns: Observational Study

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## Abstract

**Background:** Analyzing actigraphy data using standard circadian parametric models and aggregated nonparametric indices may obscure temporal information that may be a hallmark of the circadian impairment in psychiatric disorders. Functional data analysis (FDA) may overcome such limitations by fully exploiting the richness of actigraphy data and revealing important relationships with mental health outcomes. To our knowledge, no studies have extensively used FDA to study the relationship between sociodemographic, health and lifestyle, sampling, and psychiatric clinical characteristics and daily motor activity patterns assessed with actigraphy in a sample of individuals with and without depression/anxiety.

**Objective:** We aimed to study the association between daily motor activity patterns assessed via actigraphy and (1) sociodemographic, health and lifestyle, and sampling factors, and (2) psychiatric clinical characteristics (ie, presence and severity of depression/anxiety disorders).

**Methods:** We obtained 14-day continuous actigraphy data from 359 participants from the Netherlands Study of Depression and Anxiety with current (n=93), remitted (n=176), or no (n=90) depression/anxiety diagnosis, based on the criteria of the Diagnostic and Statistical Manual of Mental Disorders, fourth edition. Associations between patterns of daily motor activity, quantified via functional principal component analysis (fPCA), and sociodemographic, health and lifestyle, sampling, and psychiatric clinical characteristics were assessed using generalized estimating equation regressions. For exploratory purposes, function-on-scalar regression (FoSR) was applied to quantify the time-varying association of sociodemographic, health and lifestyle, sampling, and psychiatric clinical characteristics on daily motor activity.

**Results:** Four components of daily activity patterns captured 77.4% of the variability in the data: overall daily activity level (fPCA1, 34.3% variability), early versus late morning activity (fPCA2, 16.5% variability), biphasic versus monophasic activity (fPCA3, 14.8% variability), and early versus late biphasic activity (fPCA4, 11.8% variability). A low overall daily activity level was associated with a number of sociodemographic, health and lifestyle, and psychopathology variables: older age ( $P<.001$ ),

higher education level ( $P=.005$ ), higher BMI ( $P=.009$ ), greater number of chronic diseases ( $P=.02$ ), greater number of cigarettes smoked per day ( $P=.02$ ), current depressive and/or anxiety disorders ( $P=.05$ ), and greater severity of depressive symptoms ( $P<.001$ ). A high overall daily activity level was associated with work/school days ( $P=.02$ ) and summer (reference: winter;  $P=.03$ ). Earlier morning activity was associated with older age ( $P=.02$ ), having a partner ( $P=.009$ ), work/school days ( $P<.001$ ), and autumn and spring (reference: winter;  $P=.02$  and  $P<.001$ , respectively). Monophasic activity was associated with older age ( $P=.005$ ). Biphasic activity was associated with work/school days ( $P<.001$ ) and summer (reference: winter;  $P<.001$ ). Earlier biphasic activity was associated with older age ( $P=.005$ ), work/school days ( $P<.001$ ), and spring and summer (reference: winter;  $P<.001$  and  $P=.005$ , respectively). In FoSR analyses, age, work/school days, and season were the main determinants having a time-varying association with daily motor activity (all  $P<.05$ ).

**Conclusions:** Features of daily motor activity extracted with fPCA reflect commonly studied factors such as the intensity of daily activity and preference for morningness/eveningness. The presence and severity of depression/anxiety disorders were found to be associated mainly with a lower overall activity pattern but not with the time of the activity. Age, work/school days, and season were the variables most strongly associated with patterns and time of activity, and thus future epidemiological studies on motor activity in depression/anxiety should take these variables into account.

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## KEYWORDS

actigraphy; functional data analysis; mental health; well-being; activity

## Introduction

The near ubiquitous use of accelerometers in electronic devices ranging from smartphones to wrist-worn wearables provides the biomedical community with a potential richness of data that is useful for the study of health outcomes. Wrist-worn accelerometers have been used for more than 20 years by sleep researchers to estimate sleep and circadian activity rhythms [1,2], as well as by those studying patterns of physical activity [3]. Research indicates that disruptions in circadian activity rhythms, especially daily motor activity patterns, correlate with poor mental [4] and physical [5] health. Burton et al [6] and our recent results showed that a low level of daily motor activity is associated with depressive [7,8] and anxiety [7] disorders. In addition, sociodemographic and lifestyle factors, especially age and BMI, have been linked to disruptions in daily motor activity patterns. Compared with younger persons, older persons have been found to have lower motor activity patterns [9] and earlier bed and rise times, also known as early chronotype [10]. A higher BMI has been associated with a lower daytime activity level and a higher night-time activity level [11,12]. Circadian activity rhythms are also controlled externally by environmental and social cues. For instance, light is an important synchronizer for circadian activity rhythms [13] and has been shown to be effective in the treatment of sleep disorders [14] and affective disorders [15].

Despite the great interest in daily motor activity patterns and their association with health outcomes and other health factors, commonly used methodologies to analyze actigraphy data are limited in the description of circadian rhythms. Often used validated methods aggregate data in daily indices [16], which loses important information that may be a hallmark of circadian impairment. The traditional approach to actigraphy data analysis employs cosinor (ie, based on the mathematical formula of a cosine wave) or modified cosinor analyses that yield information concerning the amplitude of activity, the timing of “peak” activity, and the goodness-of-fit (ie, how close the pattern is to a cosine wave) [16]. Although they are often quite useful in

people with robust activity patterns, these analyses assume the presence of a particular shape of activity (ie, a predictable pattern, such as a cosine waveform) that may be different in individuals with physical or psychological impairments. Functional data analysis (FDA) can be used to model the complete time series of actigraphy data with less restrictive assumptions [17]. Recent studies using functional principal component analysis (fPCA), an FDA technique, have shown that patterns of daily motor activity with specific shapes are associated with psychiatric clinical characteristics (ie, apathy [18], depressive and anxiety symptoms [19], and objectively assessed sleep [19]) in individuals with Alzheimer disease. Another FDA technique that is increasingly being used is function-on-scalar regression (FoSR), which analyzes the relative time-varying associations between each variable of interest and the activity patterns. In addition, this method yields valuable information about the time intervals in which the variables of interest have the greatest influence on activity patterns [20,21]. Banihashemi et al [9] have suggested that older age and higher BMI are linked to lower daytime activity levels and higher BMI and greater symptom severity are associated with nocturnal activity patterns suggestive of sleep disturbances in a population with affective disorders. Those findings are based on first attempts using FDA, and such approaches have not yet been explored in a population with depressive and anxiety disorders. In addition, no studies have extensively assessed the association between actigraphy functional curves and sociodemographic, lifestyle, and sampling factors (eg, season and work/school days). FDA may better capture the complexity and dynamics of daily motor activity to reveal important behavioral biomarkers. This may help us to understand whether intervening on circadian rhythms or sleep (eg, by light therapy or sleep intervention) could be a useful regimen to reduce depressive and anxiety disorders.

In this study, we examined the association between daily activity patterns, assessed using actigraphy and FDA, and (1) sociodemographic (ie, age, sex, partner status, and education level), health and lifestyle (ie, drinking, smoking, chronic

diseases, and BMI), and sampling (ie, season, and work/school day versus nonwork/nonschool day) factors, and (2) psychiatric clinical characteristics (ie, current/remitted depressive and anxiety disorders, severity of depressive, and medication use).

## Methods

### Sample

Participants from the Netherlands Study of Depression and Anxiety (NESDA) were selected to enroll in the Ecological Momentary Assessment (EMA) and Actigraphy substudy (NESDA-EMAA). NESDA is one of the cores sites of the Motor Activity Research Consortium for Health (mMARCH) [22,23], a collaborative network for the application of objective assessment of motor activity, sleep, and mood in population and clinical samples. Details about NESDA have previously been discussed extensively [24]. NESDA was designed to investigate the course of depressive and anxiety disorders over a period of several years and the factors that influence the development and prognosis of such disorders. NESDA participants were initially recruited for baseline assessment between 2004 and 2007 (n=2981) and seen for the fifth time at the 9-year follow-up assessment (2014-2017; n=2069) for a regular follow-up interview, including a psychiatric diagnostic interview. In total, 1776 individuals participated in face-to-face interviews. A total of 367 siblings of NESDA participants who met diagnostic criteria for a depressive or anxiety disorder and had the same biological parents as their sibling(s) were included as participants to NESDA's 9-year follow-up assessment. At the 9-year follow-up, we conducted the EMAA substudy among 384 participants. The NESDA study, including the EMAA component, was approved by the VUmc ethical committee (reference number 2003/183), and all participants gave informed consent for both the regular interview and the EMAA component. A flowchart of the NESDA-EMAA study was previously provided in Difrancesco et al [7]. Eligibility criteria included the following: (1) had a smartphone or were willing to use a smartphone provided by the study, (2) were willing to wear a wrist-worn actigraphy device, and (3) could be enrolled within 1 month of the NESDA interview. Siblings were invited if they did not have a current or past diagnosis of a depressive and/or anxiety disorder or another severe psychiatric disorder (such as psychotic or severe addiction disorder). Participants of the EMAA substudy were provided with a wrist-worn GENEActiv device (Activinsights Ltd) and wore it for 2 weeks on their nondominant wrist. The devices were initialized and set to collect raw activity measures at a frequency of 30 Hz. They also completed questions on their current mood states using EMA [25]. In this paper, we only report on the actigraphy component of the study. Of the 384 participants included in the NESDA-EMAA study, 14 had no available actigraphy data for several reasons, such as technical failure (see [7] for more details), resulting in 370 (96.4%) participants with available data. According to previously published criteria [26], participants' actigraphy data were included in analyses if at least 1 weekday and 1 weekend day of usable data were available, with at least 16 hours recorded per day and per night. The final sample consisted of 359 (359/384, 93.5%) participants

with 13.68 (SD 1.26) valid days; 90.0% (323/359) of participants completed the protocol for 14 days.

### Assessment of Sociodemographic, Health and Lifestyle, and Sampling Factors

Sociodemographic and health and lifestyle factors were assessed at the 9-year follow-up. Sociodemographic factors included age, gender, education level (expressed in years), and partner status. Health and lifestyle factors included BMI (kg/m<sup>2</sup>), number of self-reported chronic diseases under treatment (eg, heart disease, diabetes, stroke, lung disease, osteoarthritis, cancer, ulcer, intestinal problems, liver disease, epilepsy, and thyroid gland disease), number of cigarettes smoked per day, and number of alcoholic drinks consumed per day. Sampling factors were assessed at the 9-year follow-up based on EMA and actigraphy assessment. Sampling factors included whether the actigraphy assessment was performed on a work/school day and the season in which the assessment was performed. Work/school day was identified with information from the EMA assessment. Season was determined based on the date of the actigraphy assessment (eg, 25/09/yyyy=autumn), and winter was used as reference.

### Assessment of Depressive and/or Anxiety Disorders and Clinical Characteristics

As in the previous assessment periods, specially trained clinical research staff conducted the diagnostic interviews at the 9-year follow-up. The Composite International Diagnostic Interview (CIDI; version 2.1) [27] was used to establish diagnoses of depressive disorders (dysthymia and major depressive disorder) and anxiety (social anxiety disorder, panic disorder with and without agoraphobia, agoraphobia, and generalized anxiety disorder) based on the Diagnostic and Statistical Manual of Mental Disorders, fourth edition. For this study, we divided participants into three groups: (1) no lifetime depressive and/or anxiety disorders, (2) remitted depressive and/or anxiety disorders, defined as having a lifetime but not current (6-month) diagnosis, and (3) current depressive or anxiety disorder diagnosed in the past 6 months.

Clinical characteristics that were examined were severity of depressive symptoms and medication use (ie, antidepressant and benzodiazepine use). Severity of depressive symptoms was assessed with the 30-item Inventory of Depressive Symptomatology (IDS) [28]. Antidepressant and benzodiazepine use was based on drug container inspection, and medications were coded according to the Anatomical Therapeutic Chemical (ATC) classification of the World Health Organization. Antidepressant and/or benzodiazepine use was considered present if participants reported using them more than 50% of the time. Antidepressants included selective serotonin reuptake inhibitors (ATC code N06AB), tricyclic antidepressants (ATC code N06AA), and other antidepressants (ATC codes N06AF, N06AG, and N06AX); benzodiazepines included ATC codes N03AE, N05BA, N05CD, and N05CF.

## Statistical Analyses

### *Descriptive Analyses*

Distributions of all variables were checked on normality with Q-Q plots. For descriptive statistics, participants' sociodemographic, health and lifestyle, and sampling factors and clinical characteristics were compared between the three groups (ie, no, remitted, and current depressive and/or anxiety disorders). For normally distributed continuous data, analysis of variance tests were used, and for data with nonnormal distributions, Kruskal-Wallis tests were used. Chi-square tests were used to test differences in frequencies across the three groups. All analyses were performed with the statistical software R (version 1.0.143), and a  $P$  value of  $<.05$  was considered statistically significant.

### *Assessment of Circadian Rhythm Patterns With fPCA*

Raw actigraphy data were processed with open source R package GGIR (version 1.5-18) [29] according to published methods [30,31]. Processing of data included autocalibration, nonwear detection, identification of potentially corrupted data, collapsing of raw data to epoch level, and computation of missing data. Collapsing of raw data to epoch level was done by averaging 5-second data. Minute-to-minute daily actigraphy data were derived per participant by summing these 5-second data; day was defined as the 24-hour clock time interval. As days with at least 16 valid hours were included in the analyses, missing data points were replaced with the participant's data from the same time of day, averaged across the other valid days, to provide a person-specific informed approach.

Daily motor activity patterns were derived from minute-level actigraphy data with the R package *fda* for FDA (version 2.4.8) [32]. First, participants' minute-to-minute actigraphy data was pre-smoothed as linear combinations of a set of nine Fourier basis functions to capture the major trends in daily motor activity (ie, the same procedure as was applied by Zeitzer et al [19] and Gershon et al [33]). Second, fPCA was used to capture the principal directions of daily variation and dimension reduction. fPCA summarized the daily-specific features as the coordinates (called principal component scores) of participants' curves in the basis spanned by the principal components. The first four daily-specific features, referred to as functional principal components in this paper, were considered because together they explained at least 75% of data variability (a similar cutoff was previously used [33]).

The association between the extracted daily functional principal components for each participant and the participant's

sociodemographic, health and lifestyle, and sampling factors and clinical characteristics were tested by using multiple generalized estimating equation (GEE) regressions, with each functional principal component as the outcome. GEE regression was used to account for correlations between repeated days per person. Separate models were run for each clinical characteristic (ie, current/remitted depressive and/or anxiety disorders, IDS score, and antidepressant use) and each model was adjusted for sociodemographic, health and lifestyle, and sampling factors; this was done to avoid collinearity induced by the high correlation between psychiatric clinical characteristics. Multiple testing correction was applied by controlling the false discovery rate [34].

### *Assessment of Time-Varying Associations of the Activity With Sociodemographic, Health and Lifestyle, and Mental Health Determinants With FoSR*

Minute-to-minute daily actigraphy data were aggregated over 10 minutes and averaged over the assessment period for each participant (ie, similar procedure as was used by Goldsmith et al [21]). FoSR was used to study the time-varying associations between participants' sociodemographic, health and lifestyle, and sampling factors and clinical characteristics, with actigraphy data as the outcome (ie, to study the time-varying associations between several factors and activity patterns). For exploratory purposes, FoSR analysis was repeated for each clinical characteristic to account for the high correlation between psychiatric clinical variables. Each model was adjusted for sociodemographic, health and lifestyle, and sampling factors. Data were analyzed with the R script developed by Goldsmith [35].

## Results

The sociodemographic, health and lifestyle, and sampling factors and clinical characteristics of the study participants are shown in Table 1. Of the total sample ( $N=359$ ), 93 and 176 participants, respectively, had current and remitted depressive and/or anxiety disorders, while 90 participants had no current depressive and/or anxiety disorders. The current depressive/anxiety disorder group was heterogeneous in that 38.7% (36/93) had anxiety disorders only, 33.3% (31/93) had depressive and anxiety disorders, and 28.0% (26/93) had depressive disorders only. As expected, individuals with current depressive and/or anxiety disorders scored significantly higher on depressive symptoms ( $P<.001$ ) and used antidepressants more frequently than individuals in the other groups, although no significant differences were found for benzodiazepine use.

**Table 1.** Participants' sociodemographics, health and lifestyle factors, actigraphy assessment and psychiatric characteristics, and medication use (N=359).

Variables	Current depressive and/or anxiety disorders (n=93)	Remitted depressive and/or anxiety disorders (n=176)	No depressive and/or anxiety disorders (n=90)	P value
<b>Sociodemographics</b>				
Age, mean (SD) <sup>a</sup>	50.1 (11.1)	48.2 (13.4)	51.3 (12.5)	.13
Female, n (%) <sup>b</sup>	58 (62.4)	120 (68.2)	50 (55.6)	.12
Education level (years), mean (SD) <sup>a</sup>	12.5 (3.4)	12.7 (2.8)	13.9 (2.9)	<.001
Has a partner, n (%) <sup>b</sup>	45 (48.4)	90 (51.1)	51 (56.7)	.52
<b>Lifestyle factors<sup>a</sup></b>				
BMI, mean (SD)	27.1 (5.1)	26.6 (5.2)	26 (5.4)	.37
Number of chronic diseases, mean (SD)	1.1 (1.2)	1.0 (1.1)	0.6 (0.8)	.008
Number of cigarettes per day, mean (SD)	3.2 (6.5)	2.9 (6.1)	0.8 (2.7)	.004
Number of drinks per day, mean (SD)	0.5 (0.8)	0.7 (1.2)	0.8 (0.8)	.16
<b>Actigraphy assessment characteristics</b>				
Number of actigraphy days, mean (SD) <sup>a</sup>	13.7 (1.0)	13.6 (1.5)	13.8 (0.7)	.48
Measures on work/school days, n (%) <sup>b</sup>	398 (34.1)	769 (37.2)	467 (42.8)	<.001
<b>Season of actigraphy measurement<sup>b</sup></b>				
Winter, n (%)	277 (21.8)	701 (29.3)	369 (29.7)	<.001
Autumn, n (%)	376 (29.5)	584 (24.4)	362 (29.1)	
Spring, n (%)	432 (33.9)	619 (25.8)	260 (20.9)	
Summer, n (%)	188 (14.8)	492 (20.5)	252 (20.3)	
<b>Psychopathology</b>				
Only depressive disorders, n (%)	26 (28.0)	46 (26.1)	N/A <sup>c</sup>	
Only anxiety disorders, n (%)	36 (38.7)	24 (13.6)	N/A	
Depressive and anxiety disorders, n (%)	31 (33.3)	106 (60.2)	N/A	
Inventory of Depressive Symptomatology, mean score (SD) <sup>a</sup>	28.6 (11.4)	20.9 (12.5)	6.0 (4.9)	<.001
<b>Medication use</b>				
Antidepressant users, n (%) <sup>b</sup>	35 (37.2)	34 (19.3)	2 (2.2)	<.001
Benzodiazepine users, n (%) <sup>b</sup>	5 (5.3)	8 (4.5)	0 (0.0)	.10

<sup>a</sup>Analysis of variance tests were performed.

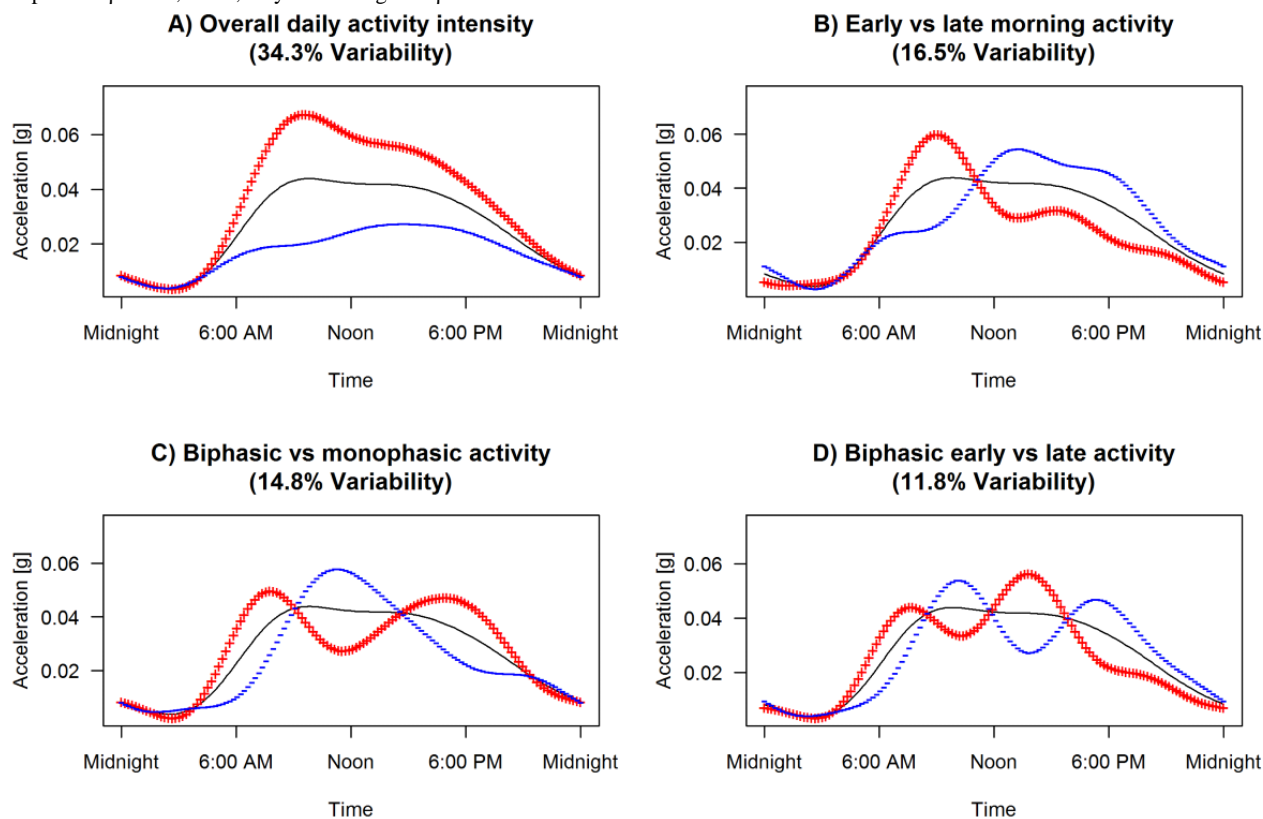
<sup>b</sup>Chi-square tests were performed.

<sup>c</sup>N/A: not applicable.

Four components of daily activity patterns that explained 77.4% of the variability in the data were extracted using fPCA (Figure 1) and interpreted as follows. The first component was the overall daily activity level. The second component described early versus late morning activity and could be indicative of

chronotype. The third component showed a biphasic versus monophasic activity pattern, while the fourth component represented an early versus late biphasic activity pattern. The biphasic pattern showed two cycles of increased activity with subsequent decreased activity.

**Figure 1.** Patterns of daily activity explaining 77.4% of variability in the data (N=359). The black line represents the average daily activity, the red line represents the average daily activity plus 1 SD of the functional principal component analysis (fPCA) score, and the blue line represents the average daily activity minus 1 SD of the fPCA score. (A) High (+) versus low (–) overall daily activity intensity. (B) Early (+) versus late (–) morning activity. (C) Biphasic (+) versus monophasic (–) activity. (D) Early (+) versus late (–) biphasic activity. Determinants associated with a pattern marked with (+) have a positive  $\beta$  value; if not, they have a negative  $\beta$  value.



A low overall daily activity level was associated with a number of sociodemographic, health and lifestyle, and psychopathology variables (Table 2): older age ( $P<.001$ ), higher education level ( $P=.005$ ), higher BMI ( $P=.009$ ), greater number of chronic diseases ( $P=.02$ ), greater number of cigarettes smoked per day ( $P=.02$ ), current depressive and/or anxiety disorders ( $P=.05$ ), and greater severity of depressive symptoms ( $P<.001$ ). A high overall daily activity level was associated with work/school days ( $P=.02$ ) and summer (reference: winter;  $P=.03$ ) (Table 2). Earlier morning activity was associated with older age ( $P=.02$ ), having a partner ( $P=.009$ ), work/school days ( $P<.001$ ), and autumn and spring (reference: winter;  $P=.02$  and  $P<.001$ , respectively) (Table 2). Monophasic activity was associated with older age ( $P=.005$ ; Table 2). Biphasic activity was associated with work/school days ( $P<.001$ ) and summer (reference: winter;  $P<.001$ ) (Table 2). Earlier biphasic activity

was associated with older age ( $P=.005$ ), work/school days ( $P<.001$ ), and spring and summer (reference: winter;  $P<.001$  and  $P=.005$ , respectively) (Table 2).

Age, work/school status, and season were significantly associated with motor activity in the FoSR analyses (Figure 2;  $P<.05$ ). Older age was especially related to lower activity in the late afternoon (around 6 PM). Working or going to school was associated with a higher activity level, with a dip in activity level around noon, compared with not working/going to school (Figure 2). Compared with individuals assessed in winter, those assessed in summer, spring, and autumn had higher activity levels in the morning and those assessed in the summer had higher activity levels late in the afternoon (Figure 2). Average daily motor activity curves by age, work/school status, and season are shown in Figure 3.

**Table 2.** Multivariable associations between daily motor activity patterns and sociodemographics, health and lifestyle factors, actigraphy assessment characteristics, and psychopathology (N=359).<sup>a</sup>

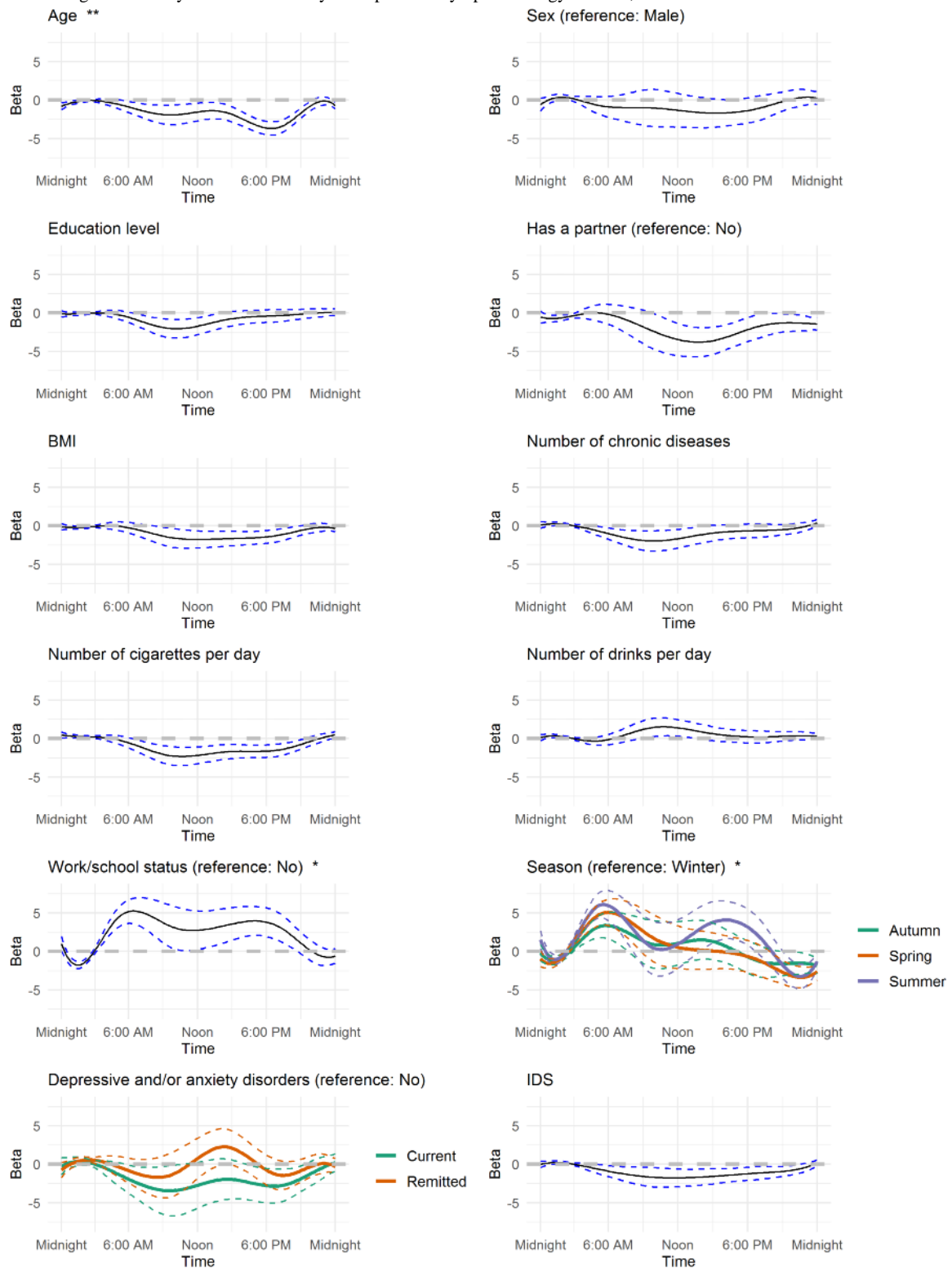
Variables	Overall daily activity intensity			Early vs late morning activity			Biphasic vs monophasic activity			Early vs late biphasic activity		
	$\beta$	SE	<i>P</i>	$\beta$	SE	<i>P</i>	$\beta$	SE	<i>P</i>	B	SE	<i>P</i>
<b>Sociodemographics</b>												
Age	-.15	0.04	<.001	.08	0.03	.02	-.10	0.03	.005	.10	0.03	.005
<b>Sex</b>												
Female	Ref <sup>b</sup>			Ref			Ref			Ref		
Male	-.05	0.08	.65	-.02	0.06	.83	-.03	0.06	.79	-.07	0.06	.34
Education level	-.13	0.04	.005	-.01	0.03	.80	.04	0.03	.30	-.03	0.03	.54
<b>Has a partner</b>												
No	Ref			Ref			Ref			Ref		
Yes	-.17	-0.08	.07	.18	-0.06	.009	-.02	-0.05	.83	.05	-0.05	.496
<b>Lifestyle factors</b>												
BMI	-.12	0.04	.009	.05	0.03	.22	-.01	0.02	.83	.01	0.03	.80
Number of chronic diseases	-.11	0.04	.02	-.01	0.03	.82	-.03	0.03	.47	-.05	0.03	.15
Number of cigarettes per day	-.14	0.05	.02	.03	0.03	.55	-.05	0.03	.19	.02	0.03	.69
Number of drinks per day	.08	0.05	.19	-.05	0.04	.40	-.04	0.03	.34	-.01	0.03	.83
<b>Actigraphy assessment characteristics</b>												
<b>Work/school day</b>												
No	Ref			Ref			Ref			Ref		
Yes	.17	0.06	.02	.28	0.05	<.001	.73	0.05	<.001	.19	0.05	<.001
<b>Season</b>												
Winter	Ref			Ref			Ref			Ref		
Autumn	.10	0.09	.39	.19	0.07	.02	.02	0.07	.83	.14	0.06	.07
Spring	.15	0.09	.21	.33	0.07	<.001	.14	0.07	.08	.26	0.07	<.001
Summer	.29	0.11	.03	.18	0.10	.19	.33	0.07	<.001	.24	0.08	.005
<b>Psychopathology</b>												
<b>Depressive and/or anxiety disorders</b>												
No	Ref			Ref			Ref			Ref		
Remitted	-.06	0.08	.60	-.04	0.08	.79	-.05	0.07	.64	.08	0.07	.39
Current	-.24	0.10	.05	.02	0.08	.83	-.001	0.08	.99	-.01	0.07	.95
IDS <sup>c</sup>	-.14	0.04	<.001	-.05	0.04	.34	-.02	0.03	.70	-.04	0.03	.30
<b>Antidepressant use</b>												
No	Ref			Ref			Ref			Ref		
Yes	-.17	0.11	.22	-.12	0.06	.14	-.08	0.07	.38	-.11	0.08	.33

<sup>a</sup>Multivariable generalized estimating equation models were run and each model was adjusted for sociodemographic, health and lifestyle, and sampling factors. Multiple testing correction was applied using false discovery rate estimation.

<sup>b</sup>Ref: reference.

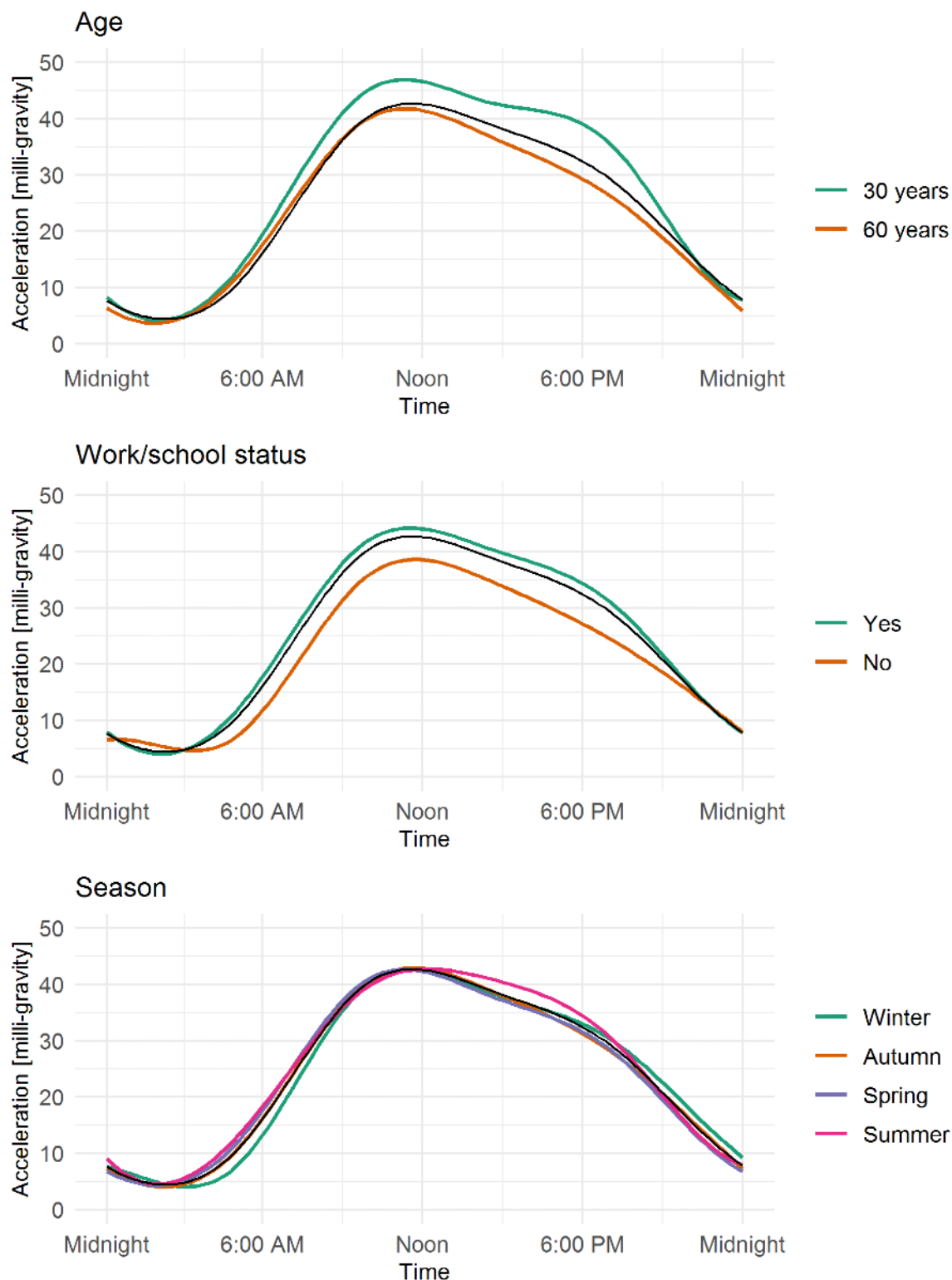
<sup>c</sup>IDS: Inventory of Depressive Symptomatology.

**Figure 2.** Effect of sociodemographics, health and lifestyle factors, and actigraphy and clinical characteristics on time of activity from multivariable function-on-scalar regression analyses. IDS: Inventory of Depressive Symptomatology. \* $P < .05$ ; \*\* $P < .01$ .





**Figure 3.** Average daily motor activity curves by age, work/school status, and season.



## Discussion

### Principal Findings

This is the first study to examine the associations between patterns of daily motor activity and sociodemographic, health and lifestyle, and sampling factors and psychiatric clinical characteristics in individuals with and without depressive and anxiety disorders using FDA. Patterns of daily motor activity

extracted with fPCA seem to reflect commonly studied circadian activity rhythm features such as daily activity level and time-of-day preference for morningness or eveningness [36]. The presence and severity of depressive and anxiety disorders were associated with a low overall activity pattern but had no impact on time of activity. Sociodemographic, health and lifestyle, and sampling factors were independently associated with daily motor activity patterns. FoSR analyses indicated that

age, work/school status, and season of assessment were associated with time of activity.

### Comparison With Previous Literature

The majority of the variability in the data (77.4%) was explained by four principal components that reflect the complexity of activity patterns. In line with previous studies employing fPCA, the first two components were commonly studied actigraphy features related to circadian rhythms: overall level of physical activity [18,19,33] and time-of-day preference for morningness or eveningness [33]. Interpretation of the third and fourth components was less clear. We found a monophasic versus biphasic activity pattern, as previously reported in other studies [19,33], where it was suggested that a biphasic rhythm may reflect napping behavior. This monophasic pattern could also be related to the temporary drop in alertness and performance that often occurs during the early afternoon, referred to as the postlunch dip, which reflects the 12-hour harmonic of the circadian clock [37]. Therefore, the level of daily activity and chronotype seem to be consistent components across studies applying fPCA, demonstrating the generalizability of the extracted components and confirming that these are important features of daily activity patterns.

Sociodemographic factors, especially age, were associated with daily motor activity patterns with a low activity level. This is in line with the results of the study by Takagi et al [38]. This may be due to the age-related decline in physical activity [39]. Aging causes changes in the organism that lead to a gradual loss of function, frailty, disease, and disability [40], and therefore result in decreased physical activity and physical functioning. The sleep-wake cycle also appears to change in the aging process. Our findings showing the association between earlier rhythms and increasing age are consistent with previous research that shows that aging is associated with advanced sleep timing (as reviewed by Duffy et al [41]) and a preference for morningness (as reviewed by Hood et al [42]). The circadian phase of melatonin has also been reported to become earlier with age, as has the timing of the cortisol rhythm [41]. The suprachiasmatic nucleus (SCN), which represents the biological clock of the brain, shows functional changes with age [43] that may be related to disturbances in circadian rhythms.

Work/school status and season, not surprisingly, were also very important to daily activity patterns and circadian rhythms. Circadian rhythms are controlled centrally by the SCN and influenced externally by behavioral/social cues and by light exposure, as reviewed by Duffy and Czeisler [44]. Our finding that assessments performed on autumn, spring, and summer days showed significantly higher overall levels of activity and higher levels of early morning activity than assessments performed on winter days seems to be consistent with the systematic review of Tucker and Gilliland [45], which showed that level of activity varies with seasonality and is the lowest during winter. Also, other factors related to the season can have an impact on daily activity such as weather conditions, which can also explain different circadian patterns across seasons. Also, on work/school days, there appeared to be higher and more sustained levels of daily activity and earlier morning activity compared with on nonwork/nonschool days. Indeed, it

is well known in the literature that modern life habits including night work, shift work [46], jet lag [47], and social jet lag [48] are associated with circadian rhythm disruptions.

We also found that health and lifestyle factors are linked to daily activity patterns. Our results are in line with previous research reporting an association between lower activity levels and higher BMIs in the NIMH Family study [12] and greater numbers of chronic diseases [49]. These results may be suggestive of sedentary behavior, a factor known to relate to weight gain and disabilities. Early morning activity was associated with higher BMIs. This might be because of respiration-related diseases such as apnea, which is known to be more prevalent in persons with a high BMI and can severely disturb sleep [50].

An important aspect of this research was to study the association between timing of daily motor activity and psychopathology. However, we found no association with the timing of the activity. Instead, our functional data-driven models showed similar associations as were found in our previous analyses on NESDA data [7,8], indicating that current depressive and/or anxiety disorders and more severe symptoms were associated with lower physical activity levels but not with a preference for eveningness. These results suggest that the use of daily indices of motor activity may be sufficient when studying the association with psychopathology. On the other hand, we have only evaluated group-level differences; studying differences at the individual level may be important to explore. For instance, by using an idiographic approach (ie, study associations that differ between time points or between individuals), it may be possible to study the dynamics between daily motor activity and depression and/or anxiety [51] and help identify patients in whom activity is strongly predictive of mood. Collecting more empirical data in clinical practice is necessary to establish whether this is a promising approach.

### Strengths of the Study

An important strength of this paper is the use of FDA, which is a useful statistical tool for data exploration and visualization. By providing a graphical representation of motor activity and circadian rhythms, FDA can help to identify specific patterns. This could help to generate new hypotheses that could, in turn, contribute to an improvement of the treatment of circadian disturbances. FDA could also be used to predict future outcomes of treatment. For instance, the study of Zeitzer et al [19] showed that low daytime activity and a late afternoon peak extracted with fPCA are predictive of higher mortality rates in community-dwelling older men, although it remains to be investigated whether fPCA components add to the prediction over traditional actigraphy measures. Daily curves of motor activity could potentially also be used in predictive models to pick up early signs of recovery or nonresponse and, if predictive, could inform clinicians in monitoring treatment response or treatment planning.

### Limitations

This study was limited by several factors. The data were observational and cross-sectional, and thus the associations cannot be inferred to be causal. As the subjects in this study

were individuals participating in the fifth wave of a prospective cohort, there may be a selection bias toward highly motivated individuals. Future studies may investigate whether our results may be replicated in a wider population with and without depression and anxiety. Although actigraphy provides only an indirect assessment of circadian rhythm, it has the advantage of continuously monitoring over a relatively long period of time. Not all functional components were easily interpretable. While the first two components are (possibly) indicative of the overall level of activity and the time-of-day preference for morningness or eveningness, the third and fourth components need to be replicated in order to provide a better interpretation and greater validity.

## Conclusions

Our study showed that features of daily motor activity extracted with fPCA reflect commonly studied factors such as the daily activity level and the time-of-day preference for morningness or eveningness. Age, work/school status, and season were the variables most strongly associated with patterns of daily activity and had time-varying associations with daily motor activity. The presence and severity of depressive and anxiety disorders were associated with a lower overall activity level but not with the timing of activity. Other than psychopathology, sociodemographic, health and lifestyle, and sampling factors were independently associated with a low overall activity pattern.

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According to European law (GDPR), data containing potentially identifying or sensitive patient information are restricted; our data involving clinical participants are not freely available in a public repository. However, data are available upon request via the NESDA Data Access Committee ([nesda@ggzingeest.nl](mailto:nesda@ggzingeest.nl)).

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## Authors' Contributions

SD, FL, and BWJHP formulated the research questions. SD performed the data cleaning and the statistical analyses, interpreted the results, wrote the manuscript, and incorporated feedback from all coauthors. FL contributed to the data cleaning, reviewed and provided feedback on all drafts of the manuscript, and critically interpreted the results. BWJHP reviewed and provided feedback on all drafts of the manuscript and critically interpreted the results. HS provided feedback on the statistical analyses, contributed to the interpretation of results, and revised the manuscript. HR, KRM, VZ, NA, AMvH, and RAS contributed to the interpretation of results and revised the manuscript critically for important intellectual content. All authors approved of the final version of the paper.

## Conflicts of Interest

None declared.

## References

1. Martin JL, Hakim AD. Wrist actigraphy. *Chest* 2011 Jun;139(6):1514-1527 [FREE Full text] [doi: [10.1378/chest.10-1872](https://doi.org/10.1378/chest.10-1872)] [Medline: [21652563](https://pubmed.ncbi.nlm.nih.gov/21652563/)]
2. Ancoli-Israel S, Martin JL, Blackwell T, Buenaer L, Liu L, Meltzer LJ, et al. The SBSM Guide to Actigraphy Monitoring: Clinical and Research Applications. *Behav Sleep Med* 2015;13 Suppl 1:S4-S38. [doi: [10.1080/15402002.2015.1046356](https://doi.org/10.1080/15402002.2015.1046356)] [Medline: [26273913](https://pubmed.ncbi.nlm.nih.gov/26273913/)]
3. Matthews CE, Hagströmer M, Pober DM, Bowles HR. Best practices for using physical activity monitors in population-based research. *Med Sci Sports Exerc* 2012 Jan;44(1 Suppl 1):S68-S76 [FREE Full text] [doi: [10.1249/MSS.0b013e3182399e5b](https://doi.org/10.1249/MSS.0b013e3182399e5b)] [Medline: [22157777](https://pubmed.ncbi.nlm.nih.gov/22157777/)]
4. Lyall LM, Wyse CA, Graham N, Ferguson A, Lyall DM, Cullen B, et al. Association of disrupted circadian rhythmicity with mood disorders, subjective wellbeing, and cognitive function: a cross-sectional study of 91 105 participants from the UK Biobank. *Lancet Psychiatry* 2018 Jun;5(6):507-514. [doi: [10.1016/S2215-0366\(18\)30139-1](https://doi.org/10.1016/S2215-0366(18)30139-1)] [Medline: [29776774](https://pubmed.ncbi.nlm.nih.gov/29776774/)]
5. Abbott SM, Zee PC. Circadian Rhythms: Implications for Health and Disease. *Neurol Clin* 2019 Aug;37(3):601-613. [doi: [10.1016/j.ncl.2019.04.004](https://doi.org/10.1016/j.ncl.2019.04.004)] [Medline: [31256792](https://pubmed.ncbi.nlm.nih.gov/31256792/)]

6. Burton C, McKinstry B, Szentagotai Tătar A, Serrano-Blanco A, Pagliari C, Wolters M. Activity monitoring in patients with depression: a systematic review. *J Affect Disord* 2013 Feb 15;145(1):21-28. [doi: [10.1016/j.jad.2012.07.001](https://doi.org/10.1016/j.jad.2012.07.001)] [Medline: [22868056](https://pubmed.ncbi.nlm.nih.gov/22868056/)]
7. Difrancesco S, Lamers F, Riese H, Merikangas KR, Beekman ATF, van Hemert AM, et al. Sleep, circadian rhythm, and physical activity patterns in depressive and anxiety disorders: A 2-week ambulatory assessment study. *Depress Anxiety* 2019 Oct;36(10):975-986 [FREE Full text] [doi: [10.1002/da.22949](https://doi.org/10.1002/da.22949)] [Medline: [31348850](https://pubmed.ncbi.nlm.nih.gov/31348850/)]
8. Minaeva O, Booij SH, Lamers F, Antypa N, Schoevers RA, Wichers M, et al. Level and timing of physical activity during normal daily life in depressed and non-depressed individuals. *Transl Psychiatry* 2020 Jul 30;10(1):259 [FREE Full text] [doi: [10.1038/s41398-020-00952-w](https://doi.org/10.1038/s41398-020-00952-w)] [Medline: [32732880](https://pubmed.ncbi.nlm.nih.gov/32732880/)]
9. Banihashemi N, Robillard R, Yang J, Carpenter JS, Hermens DF, Naismith SL, et al. Quantifying the effect of body mass index, age, and depression severity on 24-h activity patterns in persons with a lifetime history of affective disorders. *BMC Psychiatry* 2016 Sep 09;16(1):317 [FREE Full text] [doi: [10.1186/s12888-016-1023-2](https://doi.org/10.1186/s12888-016-1023-2)] [Medline: [27612556](https://pubmed.ncbi.nlm.nih.gov/27612556/)]
10. Mitchell JA, Quante M, Godbole S, James P, Hipp JA, Marinac CR, et al. Variation in actigraphy-estimated rest-activity patterns by demographic factors. *Chronobiol Int* 2017;34(8):1042-1056 [FREE Full text] [doi: [10.1080/07420528.2017.1337032](https://doi.org/10.1080/07420528.2017.1337032)] [Medline: [28650674](https://pubmed.ncbi.nlm.nih.gov/28650674/)]
11. Cespedes Feliciano EM, Quante M, Weng J, Mitchell JA, James P, Marinac CR, et al. Actigraphy-Derived Daily Rest-Activity Patterns and Body Mass Index in Community-Dwelling Adults. *Sleep* 2017 Dec 01;40(12):zsx168 [FREE Full text] [doi: [10.1093/sleep/zsx168](https://doi.org/10.1093/sleep/zsx168)] [Medline: [29029250](https://pubmed.ncbi.nlm.nih.gov/29029250/)]
12. Shou H, Cui L, Hickie I, Lameira D, Lamers F, Zhang J, et al. Dysregulation of objectively assessed 24-hour motor activity patterns as a potential marker for bipolar I disorder: results of a community-based family study. *Transl Psychiatry* 2017 Aug 22;7(8):e1211 [FREE Full text] [doi: [10.1038/tp.2017.136](https://doi.org/10.1038/tp.2017.136)] [Medline: [28892068](https://pubmed.ncbi.nlm.nih.gov/28892068/)]
13. Stothard ER, McHill AW, Depner CM, Birks BR, Moehlman TM, Ritchie HK, et al. Circadian Entrainment to the Natural Light-Dark Cycle across Seasons and the Weekend. *Curr Biol* 2017 Feb 20;27(4):508-513 [FREE Full text] [doi: [10.1016/j.cub.2016.12.041](https://doi.org/10.1016/j.cub.2016.12.041)] [Medline: [28162893](https://pubmed.ncbi.nlm.nih.gov/28162893/)]
14. van Maanen A, Meijer AM, van der Heijden KB, Oort FJ. The effects of light therapy on sleep problems: A systematic review and meta-analysis. *Sleep Med Rev* 2016 Oct;29:52-62. [doi: [10.1016/j.smrv.2015.08.009](https://doi.org/10.1016/j.smrv.2015.08.009)] [Medline: [26606319](https://pubmed.ncbi.nlm.nih.gov/26606319/)]
15. Penders TM, Stanciu CN, Schoemann AM, Ninan PT, Bloch R, Saeed SA. Bright Light Therapy as Augmentation of Pharmacotherapy for Treatment of Depression. *Prim. Care Companion CNS Disord* 2016 Oct 20;18(5). [doi: [10.4088/pcc.15r01906](https://doi.org/10.4088/pcc.15r01906)] [Medline: [27835725](https://pubmed.ncbi.nlm.nih.gov/27835725/)]
16. Klerman EB, Wang W, Phillips AJK, Bianchi MT. Statistics for Sleep and Biological Rhythms Research. *J Biol Rhythms* 2017 Feb;32(1):18-25 [FREE Full text] [doi: [10.1177/0748730416670051](https://doi.org/10.1177/0748730416670051)] [Medline: [27836939](https://pubmed.ncbi.nlm.nih.gov/27836939/)]
17. Harezlak J, Ruppert D, Wand M. *Semiparametric Regression with R*. Switzerland: Springer Nature; 2018.
18. Zeitzer JM, David R, Friedman L, Mulin E, Garcia R, Wang J, et al. Phenotyping apathy in individuals with Alzheimer disease using functional principal component analysis. *Am J Geriatr Psychiatry* 2013 Apr;21(4):391-397 [FREE Full text] [doi: [10.1016/j.jagp.2012.12.012](https://doi.org/10.1016/j.jagp.2012.12.012)] [Medline: [23498386](https://pubmed.ncbi.nlm.nih.gov/23498386/)]
19. Zeitzer JM, Blackwell T, Hoffman AR, Cummings S, Ancoli-Israel S, Stone K, Osteoporotic Fractures in Men (MrOS) Study Research Group. Daily Patterns of Accelerometer Activity Predict Changes in Sleep, Cognition, and Mortality in Older Men. *J Gerontol A Biol Sci Med Sci* 2018 Apr 17;73(5):682-687 [FREE Full text] [doi: [10.1093/gerona/glw250](https://doi.org/10.1093/gerona/glw250)] [Medline: [28158467](https://pubmed.ncbi.nlm.nih.gov/28158467/)]
20. Goldsmith J, Zipunnikov V, Schrack J. Generalized multilevel function-on-scalar regression and principal component analysis. *Biometrics* 2015 Jun 25;71(2):344-353 [FREE Full text] [doi: [10.1111/biom.12278](https://doi.org/10.1111/biom.12278)] [Medline: [25620473](https://pubmed.ncbi.nlm.nih.gov/25620473/)]
21. Goldsmith J, Liu X, Jacobson JS, Rundle A. New Insights into Activity Patterns in Children, Found Using Functional Data Analyses. *Med Sci Sports Exerc* 2016 Sep;48(9):1723-1729 [FREE Full text] [doi: [10.1249/MSS.0000000000000968](https://doi.org/10.1249/MSS.0000000000000968)] [Medline: [27183122](https://pubmed.ncbi.nlm.nih.gov/27183122/)]
22. Scott J, Murray G, Henry C, Morken G, Scott E, Angst J, et al. Activation in Bipolar Disorders: A Systematic Review. *JAMA Psychiatry* 2017 Feb 01;74(2):189-196. [doi: [10.1001/jamapsychiatry.2016.3459](https://doi.org/10.1001/jamapsychiatry.2016.3459)] [Medline: [28002572](https://pubmed.ncbi.nlm.nih.gov/28002572/)]
23. 2018. The Motor Activity Research Consortium for Health (mMARCH). URL: <http://mmarch.org/> [accessed 2019-10-07]
24. Penninx BWJH, Beekman ATF, Smit JH, Zitman FG, Nolen WA, Spinhoven P, NESDA Research Consortium. The Netherlands Study of Depression and Anxiety (NESDA): rationale, objectives and methods. *Int J Methods Psychiatr Res* 2008;17(3):121-140 [FREE Full text] [doi: [10.1002/mpr.256](https://doi.org/10.1002/mpr.256)] [Medline: [18763692](https://pubmed.ncbi.nlm.nih.gov/18763692/)]
25. Schoevers RA, van Borkulo CD, Lamers F, Servaas MN, Bastiaansen JA, Beekman ATF, et al. Affect fluctuations examined with ecological momentary assessment in patients with current or remitted depression and anxiety disorders. *Psychol Med* 2020 Apr 01:1-10. [doi: [10.1017/S0033291720000689](https://doi.org/10.1017/S0033291720000689)] [Medline: [32234092](https://pubmed.ncbi.nlm.nih.gov/32234092/)]
26. da Silva IC, van Hees VT, Ramires VV, Knuth AG, Bielemann RM, Ekelund U, et al. Physical activity levels in three Brazilian birth cohorts as assessed with raw triaxial wrist accelerometry. *Int J Epidemiol* 2014 Dec;43(6):1959-1968 [FREE Full text] [doi: [10.1093/ije/dyu203](https://doi.org/10.1093/ije/dyu203)] [Medline: [25361583](https://pubmed.ncbi.nlm.nih.gov/25361583/)]
27. Wittchen HU. Reliability and validity studies of the WHO--Composite International Diagnostic Interview (CIDI): a critical review. *J Psychiatr Res* 1994;28(1):57-84. [doi: [10.1016/0022-3956\(94\)90036-1](https://doi.org/10.1016/0022-3956(94)90036-1)] [Medline: [8064641](https://pubmed.ncbi.nlm.nih.gov/8064641/)]

28. Rush AJ, Gullion CM, Basco MR, Jarrett RB, Trivedi MH. The Inventory of Depressive Symptomatology (IDS): psychometric properties. *Psychol Med* 1996 May;26(3):477-486. [Medline: [8733206](#)]
29. van Hees V. Accelerometer data processing with GGIR. URL: <https://cran.r-project.org/web/packages/GGIR/vignettes/GGIR.html> [accessed 2017-10-01]
30. van Hees VT, Fang Z, Langford J, Assah F, Mohammad A, da Silva ICM, et al. Autocalibration of accelerometer data for free-living physical activity assessment using local gravity and temperature: an evaluation on four continents. *J Appl Physiol* (1985) 2014 Oct 01;117(7):738-744 [FREE Full text] [doi: [10.1152/jappphysiol.00421.2014](#)] [Medline: [25103964](#)]
31. van Hees VT, Gorzelniak L, Dean León EC, Eder M, Pias M, Taherian S, et al. Separating movement and gravity components in an acceleration signal and implications for the assessment of human daily physical activity. *PLoS One* 2013;8(4):e61691 [FREE Full text] [doi: [10.1371/journal.pone.0061691](#)] [Medline: [23626718](#)]
32. Ramsay J, Graves S, Hooker G. fda: Functional Data Analysis. URL: <https://cran.r-project.org/web/packages/fda/index.html> [accessed 2018-04-27]
33. Gershon A, Ram N, Johnson SL, Harvey AG, Zeitzer JM. Daily Actigraphy Profiles Distinguish Depressive and Interepisode States in Bipolar Disorder. *Clin Psychol Sci* 2016 Jul;4(4):641-650 [FREE Full text] [doi: [10.1177/2167702615604613](#)] [Medline: [27642544](#)]
34. Noble WS. How does multiple testing correction work? *Nat Biotechnol* 2009 Dec;27(12):1135-1137 [FREE Full text] [doi: [10.1038/nbt1209-1135](#)] [Medline: [20010596](#)]
35. Goldsmith J. Function-on-Scalar Regression. URL: [http://jeffgoldsmith.com/IWAFDA/shortcourse\\_fosr.html](http://jeffgoldsmith.com/IWAFDA/shortcourse_fosr.html) [accessed 2019-05-24]
36. Kivelä L, Papadopoulos MR, Antypa N. Chronotype and Psychiatric Disorders. *Curr Sleep Med Rep* 2018;4(2):94-103 [FREE Full text] [doi: [10.1007/s40675-018-0113-8](#)] [Medline: [29888167](#)]
37. Monk TH. The post-lunch dip in performance. *Clin Sports Med* 2005 Apr;24(2):e15-23, xi. [doi: [10.1016/j.csm.2004.12.002](#)] [Medline: [15892914](#)]
38. Takagi D, Nishida Y, Fujita D. Age-associated changes in the level of physical activity in elderly adults. *J Phys Ther Sci* 2015 Dec;27(12):3685-3687 [FREE Full text] [doi: [10.1589/jpts.27.3685](#)] [Medline: [26834332](#)]
39. Sallis JF. Age-related decline in physical activity: a synthesis of human and animal studies. *Med Sci Sports Exerc* 2000 Sep;32(9):1598-1600. [doi: [10.1097/00005768-200009000-00012](#)] [Medline: [10994911](#)]
40. Paterson DH, Warburton DE. Physical activity and functional limitations in older adults: a systematic review related to Canada's Physical Activity Guidelines. *Int J Behav Nutr Phys Act* 2010 May 11;7:38 [FREE Full text] [doi: [10.1186/1479-5868-7-38](#)] [Medline: [20459782](#)]
41. Duffy JF, Zitting K, Chinoy ED. Aging and Circadian Rhythms. *Sleep Med Clin* 2015 Dec;10(4):423-434 [FREE Full text] [doi: [10.1016/j.jsmc.2015.08.002](#)] [Medline: [26568120](#)]
42. Hood S, Amir S. The aging clock: circadian rhythms and later life. *J Clin Invest* 2017 Feb 01;127(2):437-446 [FREE Full text] [doi: [10.1172/JCI90328](#)] [Medline: [28145903](#)]
43. Peters R. Ageing and the brain. *Postgrad Med J* 2006 Feb;82(964):84-88 [FREE Full text] [doi: [10.1136/pgmj.2005.036665](#)] [Medline: [16461469](#)]
44. Duffy JF, Czeisler CA. Effect of Light on Human Circadian Physiology. *Sleep Med Clin* 2009 Jun;4(2):165-177 [FREE Full text] [doi: [10.1016/j.jsmc.2009.01.004](#)] [Medline: [20161220](#)]
45. Tucker P, Gilliland J. The effect of season and weather on physical activity: a systematic review. *Public Health* 2007 Dec;121(12):909-922. [doi: [10.1016/j.puhe.2007.04.009](#)] [Medline: [17920646](#)]
46. James SM, Honn KA, Gaddameedhi S, Van Dongen HPA. Shift Work: Disrupted Circadian Rhythms and Sleep-Implications for Health and Well-Being. *Curr Sleep Med Rep* 2017 Jun;3(2):104-112 [FREE Full text] [doi: [10.1007/s40675-017-0071-6](#)] [Medline: [29057204](#)]
47. Vosko AM, Colwell CS, Avidan AY. Jet lag syndrome: circadian organization, pathophysiology, and management strategies. *Nat Sci Sleep* 2010;2:187-198 [FREE Full text] [doi: [10.2147/NSS.S6683](#)] [Medline: [23616709](#)]
48. Wittmann M, Dinich J, Merrow M, Roenneberg T. Social jetlag: misalignment of biological and social time. *Chronobiol Int* 2006;23(1-2):497-509. [doi: [10.1080/07420520500545979](#)] [Medline: [16687322](#)]
49. Durstine JL, Gordon B, Wang Z, Luo X. Chronic disease and the link to physical activity. *Journal of Sport and Health Science* 2013 Mar;2(1):3-11. [doi: [10.1016/j.jshs.2012.07.009](#)]
50. Romero-Corral A, Caples S, Lopez-Jimenez F, Somers V. Interactions Between Obesity and Obstructive Sleep Apnea. *Chest* 2010 Mar;137(3):711-719. [doi: [10.1378/chest.09-0360](#)] [Medline: [20202954](#)]
51. Rosmalen JGM, Wenting AMG, Roest AM, de Jonge P, Bos EH. Revealing causal heterogeneity using time series analysis of ambulatory assessments: application to the association between depression and physical activity after myocardial infarction. *Psychosom Med* 2012 May;74(4):377-386. [doi: [10.1097/PSY.0b013e3182545d47](#)] [Medline: [22582335](#)]

## Abbreviations

**ATC:** Anatomical Therapeutic Chemical

**CIDI:** Composite International Diagnostic Interview

**EMA:** ecological momentary assessment

**FDA:** functional data analysis

**FoSR:** function-on-scalar regression

**fPCA:** functional principal component analysis

**GEE:** generalized estimating equation

**IDS:** Inventory of Depressive Symptomatology

**mMARCH:** Motor Activity Research Consortium for Health

**NESDA:** Netherlands Study of Depression and Anxiety

**NESDA-EMAA:** Ecological Momentary Assessment and Actigraphy substudy of the Netherlands Study of Depression and Anxiety

**SCN:** suprachiasmatic nucleus

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