

Five-year Evolution Patterns of Physical Activity and Sedentary Behavior in Patients with Lower-limb Osteoarthritis and Their Sociodemographic and Clinical Correlates

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ABSTRACT. *Objective.* The present study aimed to identify trajectories of physical activity (PA) components (frequency, duration, intensity, and type) and screen-based sedentary behavior (SB) as well as baseline predictors of each trajectory in patients with hip and/or knee osteoarthritis (OA).

Methods. We included 878 patients with a 5-year follow-up from the KHOALA cohort. PA and SB were measured by the Modifiable Activity Questionnaire. We used group-based trajectory analysis to identify the trajectories of PA components and screen-based SB, and multivariable logistic regression to determine predictors of the identified trajectories.

Results. Two groups of trajectories were identified for each PA component and 3 for SB. High and decreasing PA duration was associated with female sex (OR 0.3, 95% CI 0.1–0.5) as was low and stable, more so than high and decreasing prevalence of weight-bearing activities (OR 0.6, 95% CI 0.4–0.9). Patients with impaired patient-reported outcome measures and obese patients often featured low versus high and decreasing prevalence of weight-bearing activities. Predictors of moderate and high versus low and slightly increasing screen-based SB trajectories were male sex, age < 60 years, single status (OR 1.5, 95% CI 1.1–2.1), obesity (OR 2.1, 95% CI 1.4–3.1), smoking (OR 2.0, 95% CI 1.1–3.7), and less physical jobs. Predictors of moderate and high versus low screen-based SB trajectories were all sociodemographic: male sex, age < 60 years, single status, obesity, smoking, and less physical jobs.

Conclusion. Sociodemographic and clinical predictors of trajectories vary between PA components; they are associated mainly with PA frequency and type. No clinical characteristics were associated with screen-based SB.

Key Indexing Terms: group-based trajectory analysis, osteoarthritis, physical activity, sedentary behavior

Osteoarthritis (OA) is the most common form of arthritis, and knee and/or hip OA has been ranked as the 13th highest contributor to global disability in the Global Burden of Disease study^{1,2}. Being physically active and less sedentary plays a major role in reducing OA symptoms, physical function impairment, and pain, and can improve quality of life (QOL)³. Further, regular physical activity (PA) is associated with decreased risk of cardiovascular mortality in this population⁴. Despite these

potential health benefits, several studies have shown that the recommended levels of PA are less likely to be met by adults with OA than those without OA^{5,6,7}, and the former spend even more time in sedentary behavior (SB) than the general population⁸.

All the above-mentioned studies considered the average level of PA of the sample over a predefined time and so did not consider several distinct patterns that may be present within that sample or whether these behaviors change over time. However, several studies have reported the presence of various subgroups of PA trajectories in the general population⁹, specifically populations such as women¹⁰ and in certain diseases such as rheumatoid arthritis¹¹ or heart disease¹². Indeed, a recent systematic review including 27 longitudinal studies reported the presence of various PA subgroups in the general population, varying most commonly between 3 or 4 trajectory groups⁹. Considering that the course of OA is heterogeneous, with some patients having stable or even improved disease over many years and others showing increasing pain, disability, or structural damage¹³, it is not surprising that PA practice and SB can vary across patients with OA and over time.

However, to date, no studies have examined the presence of common trajectories of PA or SB across representative cohorts of people with OA. In this context, the aims of the present study were to identify and describe trajectories of PA components (i.e.,

Funding for the KHOALA cohort study was obtained from public sources (INSERM, CHU de Nancy, Conseil Régional de Lorraine, Société Française de Rhumatologie) and unrestricted grants from pharmaceutical companies (Expanscience, Genevrier, Grünenthal, Merck & Co Inc., Pfizer, Pierre Fabre Médicaments, Sanofi-Aventis France). Opinions expressed in the present article are those of the authors and do not necessarily reflect those of the sponsors. The study sponsors did not take part in the study design, collection, analysis, and interpretation of data, writing of the report, or the decision to submit the article for publication.

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Accepted for publication February 28, 2020.

frequency, duration, intensity, and type) and screen-based SB over a 5-year follow-up period and to identify baseline predictors for each trajectory in a representative cohort of patients with symptomatic hip and/or knee OA.

MATERIALS AND METHODS

The KHOALA cohort. The Knee and Hip Osteoarthritis Long-term Assessment (KHOALA)¹⁴ cohort is an ongoing, multiregional, population-based cohort. It included 878 patients aged 40–75 years with uni- or bilateral symptomatic hip and/or knee OA [American College of Rheumatology (ACR) criteria] and a Kellgren-Lawrence grade 2 or greater. Patients were recruited from a national prevalence survey conducted in France from April 2007 to March 2009¹⁵ and were followed up every year by the use of a self-reporting questionnaire and a clinical examination at baseline and Years 3 and 5. For the purpose of this study, we used data collected at inclusion and each year of follow-up.

All patients gave their written informed consent to be in the KHOALA study. The ethics committee CPP Est III approved the cohort study (no. 07.01.01) registered at ClinicalTrials.gov (no. NCT00481338).

Outcome measures: Physical activity and sedentary behavior. Patients self-reported PA and SB at inclusion and 1, 2, 3, 4, and 5 years' follow-up by using the Modifiable Activity Questionnaire¹⁶. The questionnaire assesses leisure time PA during the previous 12 months. Patients were asked to indicate the types of PA they had performed, number of months, average number of times per month, and number of minutes spent in an activity. The questionnaire also uses time spent daily watching television and using a computer as an indicator of screen-based SB.

For each participant and each year of follow-up, the frequency of PA was defined as the number of times the patient participated in the PA per week, and the duration as the number of hours per week of PA declared. The intensity was expressed by a continuous variable representing the average metabolic equivalent of task (MET) of PA declared based on the 2011 Compendium of Physical Activities. The type of PA was defined in 2 categories: patients practicing at least 1 weight-bearing (WB) activity and patients practicing only reduced WB (RWB) activities. WB activities such as running or playing tennis are defined as force-generating exercises placing higher mechanical stress on the human skeleton than daily living. RWB activities such as swimming or horseback riding generate load below that associated with activities of daily living. These are activities during which individuals do not support their own weight¹⁷.

Potential predictors. Two different types of predictors were examined at baseline: sociodemographic factors, and clinical and patient-reported outcome measure (PROM) factors.

The sociodemographic factors included sex (female/male), age (40–49, 50–59, ≥ 60 yrs), education level (primary, secondary, or university), marital status (in a couple or single), socioprofessional category (farmer, artisan, trader; executive, intellectual profession; intermediate occupation; employee; without professional activity), smoking status (yes/no), alcohol consumption (yes/no), monthly income (low, intermediate, high), geographical zone in France (north, west, south), and size of the residential area (< 2000, 2000–49,999, or ≥ 50,000 inhabitants). All the above-mentioned variables were treated as categorical variables.

Clinical and PROM factors consisted of affected joint, BMI (normal < 25 kg/m², overweight 25–29.9 kg/m², or obese ≥ 30 kg/m²) based on measured height and weight, Kellgren-Lawrence grade (grade 2, 3, or 4), and comorbidities (Functional Comorbidity Index; FCI)¹⁸. Patients reported the date of first symptoms (month and year) and the duration between symptom onset, and inclusion was calculated accordingly. The overall level of hip or knee pain was measured on a visual analog scale (VAS) with scores ranging from 0 (no pain) to 10 (unbearable pain) by answering the following question: "What is the overall level of your hip/knee pain under any

circumstances in the past 48 hours?" The Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) was used to measure physical function, pain, and stiffness, with scores ranging from 0 (best state) to 100 (worst state)¹⁹. QOL was measured by the vitality dimension of the Medical Outcomes Study 36-item Short Form survey (SF-36) and by the 5 dimensions of the OA Knee and Hip QOL (OAKHQOL) questionnaire: PA, mental health, pain, social functioning, and social support^{20,21}. Scores for both instruments range from 0 (worst state) to 100 (best state). The General Health Questionnaire 28 [scores from 0 (best state) to 84 (worst state)] was used to assess somatic symptoms, anxiety and insomnia, social dysfunction, and severe depression²². The environmental health domain of the World Health Organization QOL (WHOQOL-BREF) scale was used to assess environmental factors²³.

Statistical methods. Statistical analyses involved different steps. First, patient characteristics were described with mean (SD) for quantitative variables and number (percentages) for categorical variables. Second, a semiparametric, group-based trajectory model was used to evaluate subgroups that followed similar trajectories in each PA component and screen-based SB, based on identifying heterogeneous longitudinal polynomial trajectories using the TRAJ procedure of SAS v9.4 (SAS Institute, Inc.)²⁴. The optimal number of groups and degree of polynomial function in each trajectory group were determined by using the Bayesian information criterion (BIC): a lower BIC value indicates a better fitting model. Other criteria such as the proportion of patients in each trajectory group (> 5%) and the clinical interpretability of the identified trajectories were also considered. We fitted 5 models of trajectories: PA frequency, intensity, time, type, and screen-based SB. To check the robustness of the final optimized models, we used the average of the posterior probabilities of group membership for individuals assessing the fit of the models²⁵. Description of missing data for PA components and SB are presented in Supplementary Tables 1 and 2 (available from the authors on request). Missing data were characterized by using the SAS macro %missingPattern²⁶, and their mechanism was assessed by searching for evidence of monotonicity and unit nonresponse. A sensitivity analysis using patients without missing data was also performed. Third, multivariable logistic regression analysis was used to identify baseline predictors of trajectory membership for each model, estimating OR and 95% CI. Factors including sex, age, education level, marital status, socioprofessional category, smoking status, alcohol consumption, monthly income, geographical zone in France, size of the residential area, affected joint, duration between onset of symptoms and inclusion, BMI, FCI, WOMAC, SF-36, VAS, and OAKHQOL questionnaires were tested in bivariate analysis. Because the linearity assumption for logistic regression was violated, data for WOMAC, SF-36, VAS, and OAKHQOL questionnaires were categorized into tertiles. Only factors with a significant association at $P = 0.2$ in bivariate models were entered into multivariable models. We used stepwise variable selection with significance level $P = 0.1$ for entry into the model and $P = 0.05$ for staying in the model. $P < 0.05$ was considered statistically significant. Analyses involved the use of SAS v9.4 (SAS Institute Inc.).

RESULTS

Descriptive analysis. Sociodemographic and clinical data for the 878 patients included are in Table 1. At baseline, the mean (SD) duration of moderate intensity PA was 3.9 (4.5) h/week, frequency was 4.3 (3.6) times/week, and intensity was 4.2 (0.8) MET (Table 1). The most frequent PA was walking for pleasure (29.7%), gardening (22.8%), bicycling for pleasure (8.2%), and swimming (7.9%).

Identified group-based trajectories. We observed no structured missing data pattern, such as monotonicity or unit nonresponse, for outcomes criteria. Accordingly, data were considered

Table 1. Baseline patient characteristics (N = 878).

	N	% or Mean (\pm SD)
Joint		
Hip	222	25.3
Knee	607	69.1
Hip and knee	49	5.6
Sex		
Male	269	30.6
Female	609	69.4
Age at inclusion, yrs	878	62.0 (\pm 8.5)
Education level		
Primary	200	22.9
Secondary	457	52.3
University	216	24.8
Kellgren-Lawrence grade		
Grade 2	443	50.5
Grade 3	259	29.5
Grade 4	176	20.0
Marital status		
In a couple	601	68.7
Single	273	31.3
Geographical zone		
North France	560	67.9
West France	111	13.4
South France	153	18.7
Size of the residential area, no. inhabitants		
< 2000	305	34.7
2000–49,999	377	42.9
\geq 50,000	196	22.3
Socioprofessional category		
Farmer, artisan, trader	119	13.6
Executive, intellectual professional	127	14.5
Intermediate occupation	185	21.1
Employee	378	43.1
Without professional activity	69	7.9
Retired (yes)	546	62.4
Smoking status (yes)	135	15.5
Alcohol consumption (yes)	523	61.7
BMI, kg/m ²		
Normal (< 25)	212	24.1
Overweight (25–29)	329	37.5
Obese (\geq 30)	337	38.4
PA frequency, times/week	745	4.3 (\pm 3.6)
PA intensity (MET)	770	4.2 (\pm 0.8)
PA duration, h/week	625	3.9 (\pm 4.5)
PA type		
Nonweight-bearing activities	459	59.6
Weight-bearing activities	311	40.4
Sedentary behavior, h/day	644	4.1 (\pm 2.3)

MET: metabolic equivalent task; PA: physical activity.

missing at random and were included in the analyses as allowed when using group-based trajectory modeling without requiring imputation, because this model handles missing data by using maximum likelihood estimation²⁵. Restricting the analysis to patients with complete data did not alter the number or size of the observed trajectory pattern groups (Supplementary Figures 1 and 2, available from the authors on request).

Two trajectories were identified for each PA component and 3 for screen-based SB (Figure 1 and Figure 2). With these, the average posterior probability of group membership for individuals assigned to each trajectory was > 0.8 (95% CI 0.82–0.98; Supplementary Table 1, available from the authors on request).

- Trajectories of frequency: (1) moderate and stable frequency ($n = 703$; 80.1%) represented a mean PA frequency of 3 times/week; and (2) high and slightly decreasing frequency ($n = 175$; 19.9%), a frequency of 8 times/week at inclusion, which decreased to 7 times/week at 5-year follow-up.
- Trajectories of intensity: (1) low and quasi-stable level of moderate intensity ($n = 773$; 88%) represented a mean of 4 MET during follow-up; and (2) moderate and slightly decreasing level of moderate intensity ($n = 105$; 12%) represented intensity slightly decreased from 5.5 MET at inclusion to 5 MET at 5-year follow-up.
- Trajectories of duration: (1) moderate and stable duration ($n = 800$; 91.1%) represented a mean duration of 3 h/week; and (2) high and decreasing duration ($n = 78$; 8.9%) represented a duration decreased from 12.9 h/week at inclusion to 3 h/week at 5-year follow-up.
- Trajectories of type: (1) low and stable frequency of WB activities ($n = 549$; 62.5%) represented stable prevalence of WB activities at 9% during follow-up; and (2) high and decreasing frequency of WB activities ($n = 329$; 37.5%) represented decreased prevalence from 89.7% at inclusion to 77.2% at 5-year follow up.
- Trajectories of screen-based SB: (1) low and slightly increasing SB ($n = 500$; 57.1%) represented screen-based SB slightly increased from 2.8 at inclusion to 3.2 h/day at 5-year follow-up; (2) moderate and slightly increasing SB ($n = 290$; 33%) represented screen-based SB slightly increased from 4.9 at inclusion to 5.3 h/day at 5-year follow-up; and (3) high and stable SB ($n = 88$; 10%) represented screen-based SB stable at 8.4 h/day during follow-up.

Predictors of physical activity and screen-based sedentary behavior trajectories. Results of bivariate associations are in Supplementary Tables 3 and 4 (available from the authors on request). Predictors of PA components and SB from multivariable logistic regression are in Table 2 and Table 3, respectively.

For trajectories of intensity, no sociodemographic or clinical factors were statistically significant in the multivariable logistic regression model.

For trajectories of duration, only sex was a predictor for duration trajectories. Females belonged mostly to the moderate and stable trajectories more so than to the high and decreasing group (OR 0.3, 95% CI 0.1–0.5, $P < 0.0001$).

For trajectories of frequency, the probability of belonging to the high and slightly decreasing rather than the moderate and stable trajectory was associated with age (≥ 60 yrs), low monthly income level (OR 0.5, 95% CI 0.3–0.9, $P = 0.02$), and improved WOMAC physical function (OR 0.4, 95% CI 0.3–0.7, $P = 0.0007$; Table 2).

For trajectories of type of PA, the probability of belonging to

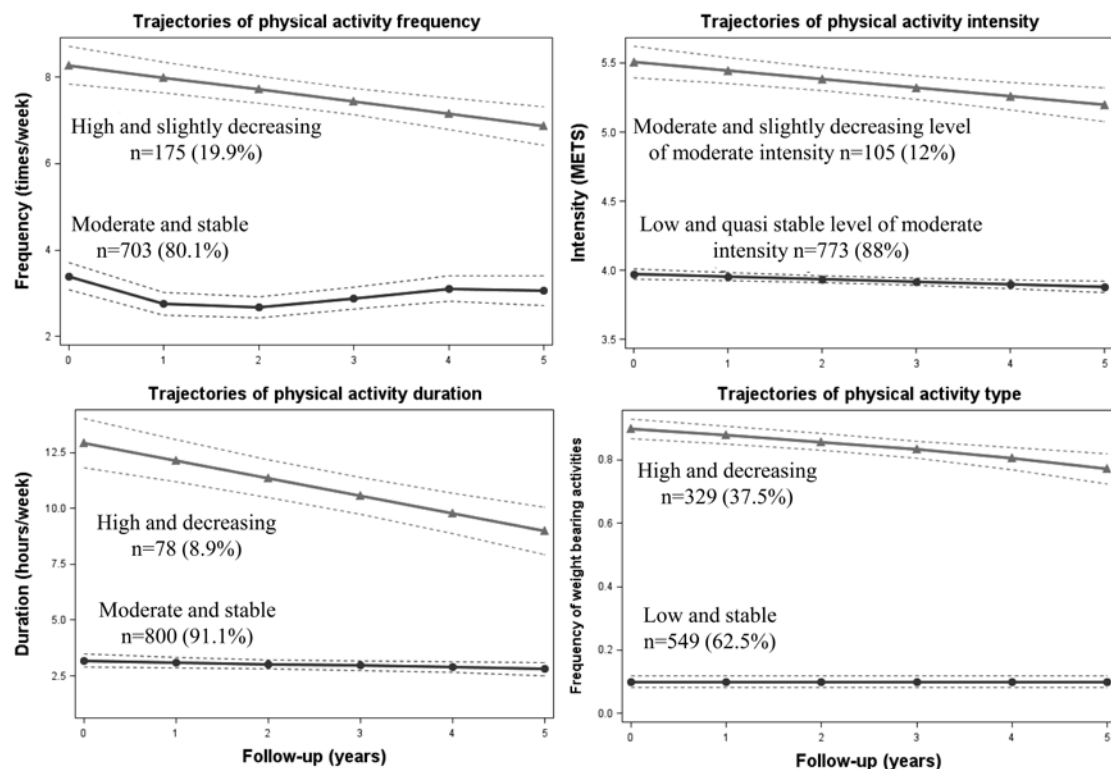


Figure 1. Identified trajectories of physical activity components. MET: metabolic equivalent of task.

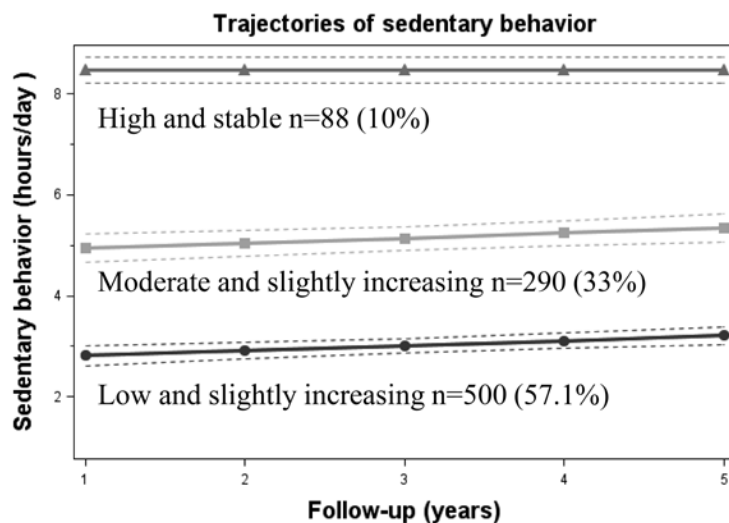


Figure 2. Identified trajectories of sedentary behavior.

the high and decreasing versus stable frequency of WB activities was reduced with female sex (OR 0.6, 95% CI 0.4–0.9, $P = 0.007$) and obesity (OR 0.5, 95% CI 0.3–0.9, $P = 0.007$), and with better SF-36 vitality level (OR 2.2, 95% CI 1.3–3.8, $P = 0.004$), OAKHQOL physical function score (OR 2.1, 95% CI 1.1–3.9, $P = 0.03$), and WOMAC

physical function score (OR 0.5, 95% CI 0.3–0.8, $P = 0.02$], and improved pain VAS level (OR 0.4, 95% CI 0.3–0.7, $P = 0.002$; Table 2).

For trajectories of screen-based SB, the probability of belonging to the high and stable versus low and slightly increasing trajectory was associated with being single (OR 1.8,

Table 2. Factors associated with physical activity frequency and type trajectories derived from the multivariable logistic regression.

	Frequency				Type			
	MSFreq, n = 703 (80.1%) %	HDFreq, n = 175 (19.9%) %	HDFreq (vs MSFreq) OR (95% CI) P*		LSWB, n = 549 (62.5%) %	HDWB, n = 329 (37.5%) %	HDWB (vs LSBW) OR (95% CI) P*	
Sex								0.007
Male	77.3	22.7			52.4	47.6	1	
Female	84.1	15.9			69.5	30.5	0.6 (0.4–0.9)	
Age at inclusion, yrs				0.01				
40–49	91.7	8.3	1		52.8	47.2		
50–59	85.2	14.8	2.2 (0.8–5.9)		65.8	34.2		
≥ 60	79.2	20.8	3.3 (1.3–8.5)		64.9	35.1		
BMI, kg/m ²								0.007
Normal (< 25)	82.5	17.5			55.2	44.8	1	
Overweight (25 to 29)	79.6	20.4			55.6	44.4	1.0 (0.7–1.6)	
Obese (> 30)	84.0	16.0			78.3	21.7	0.5 (0.3–0.9)	
Monthly income				0.02				
Low (< €1220)	75.7	24.3	1		75.7	24.3		
Intermediate (€1220–2440)	82.8	17.2	0.6 (0.4–0.9)		67.2	32.8		
High (≥ €2440)	84.2	15.8	0.5 (0.3–0.9)		56.6	43.4		
WOMAC function				0.0007				0.02
1st tertile (0–20)	80.2	19.8	1		50.5	49.5	1	
2nd tertile (21–44)	77.7	22.3	1.0 (0.7–1.6)		61.9	38.1	0.8 (0.6–1.3)	
3rd tertile (45–100)	88.0	12.0	0.4 (0.3–0.7)		80.6	19.4	0.5 (0.3–0.8)	
WOMAC pain								0.003
1st tertile (0–20)	79.6	20.4			56.0	44.0	1	
2nd tertile (21–40)	79.5	20.5			57.0	43.0	1.7 (1.1–2.7)	
3rd tertile (41–100)	86.9	13.1			81.6	18.4	1.0 (0.5–1.9)	
SF-36 vitality								0.004
1st tertile (0–40)	87.2	12.8			80.7	19.3	1	
2nd tertile (40–60)	80.4	19.6			60.6	39.4	2.2 (1.3–3.6)	
3rd tertile (60–100)	78.1	21.9			52.7	47.3	2.2 (1.3–3.8)	
Pain VAS								0.002
1st tertile (0–21)	81.1	18.9			51.5	48.5	1	
2nd tertile (21–48)	81.7	18.3			64.9	35.1	0.6 (0.4–0.9)	
3rd tertile (48–100)	82.0	18.0			77.9	22.1	0.4 (0.3–0.7)	
OAKHQOL physical activity								0.03
1st tertile (0.625–58)	84.7	15.3			81.1	18.9	1	
2nd tertile (58.1–78.75)	80.8	19.2			64.1	35.9	1.2 (0.7–2.1)	
3rd tertile (79–100)	79.8	20.2			47.9	52.1	2.1 (1.1–3.9)	

*Only factors with a significant association at $P = 0.2$ in the bivariate analysis were entered into the multivariable model. Stepwise variable selection with significance level for entry into the model at $P = 0.1$ and with significance level for staying in the model at $P = 0.05$ was used. HDFreq: high and slightly decreasing frequency; HDWB: high and decreasing prevalence of weight-bearing activities; LSBW: low and stable prevalence of weight-bearing activities; MSFreq: moderate and stable frequency; OAKHQOL: Osteoarthritis Knee and Hip Quality Of Life questionnaire [0 (worst state) to 100 (best state)]; SF-36: Medical Outcomes Study 36-item Short Form survey [0 (worst state) to 100 (best state)]; VAS: visual analog scale; WOMAC: Western Ontario and McMaster Universities Osteoarthritis Index [0 (best state) to 100 (worst state)].

95% CI 1.1–2.9, $P = 0.02$), a smoker (OR 2.0, 95% CI 1.1–3.7, $P = 0.02$), and age 60 years or greater (OR 0.4, 95% CI 0.2–0.9, $P = 0.01$). As well, the probability of belonging to the moderate and slightly increasing versus low and slightly increasing trajectory was associated with being single (OR 1.5, 95% CI 1.1–2.1, $P = 0.01$), and was reduced with female sex (OR 0.7, 95% CI 0.5–1.0, $P = 0.02$) and obesity (OR 2.1, 95% CI 1.4–3.1, $P = 0.002$; Table 3).

DISCUSSION

In this current investigation of the trajectories of PA and SB over time in patients with OA, we report 2 major findings. First, in general, patients with OA were distributed between 2 distinct trajectories for each PA component (a low to moderate stable trajectory including most patients, and a high decreasing trajectory with a few patients), and 3 distinct trajectories of SB (low, moderate, and high level of screen-based SB). Second,

Table 3. Factors associated with sedentary behavior trajectories derived from the multivariable logistic regression.

	LISB, n = 500 (57.1%) %	MISB, n = 290 (33%) %	HSSB, n = 88 (10%) %	MISB (vs LISB)		HSSB (vs LISB)	
				OR (95% CI)	P*	OR (95% CI)	P*
Sex					0.02		
Male	58.0	33.1	8.9	1			
Female	64.2	26.6	9.2	0.7 (0.5–1.0)			
Age at inclusion, yrs							0.01
40–49	55.6	27.8	16.7			1	
50–59	57.4	29.7	12.9			0.8 (0.4–1.9)	
≥ 60	65.5	28.2	6.3			0.4 (0.2–0.9)	
Marital status					0.01		0.02
In a couple	65.9	26.3	7.7	1		1	
Single	55.9	32.5	11.6	1.5 (1.1–2.1)		1.8 (1.1–2.9)	
Socioprofessional category							0.01
Farmer, artisan, trader	52.8	31.5	15.7			1	
Executive, intellectual professional	62.2	27.6	10.3			11.2 (2.5–50.5)	
Intermediate occupation	62.4	29.1	8.5			5.6 (1.2–25.3)	
Employee	69.7	28.6	1.7			5.3 (1.2–22.7)	
Without professional activity	66.7	23.2	10.1			6.3 (1.2–32.5)	
Smoking status (yes)	51.9	32.6	15.6			2.0 (1.1–3.7)	0.02
Alcohol consumption (yes)	61.6	28.7	9.8				
BMI, kg/m ²					0.002		
Normal (< 25)	72.2	20.3	8.2	1			
Overweight (25–29)	61.4	28.9	9.3	1.6 (1.0–2.4)			
Obese (> 30)	57.0	33.5	10.2	2.1 (1.4–3.1)			

*Only factors with a significant association at $P = 0.2$ in the bivariate model were entered into the multivariable model. Stepwise variable selection with significance level for entry into the model at $P = 0.1$ and with significance level for staying in the model at $P = 0.05$ was used. HSSB: high and stable sedentary behavior; LISB: low and slightly increasing sedentary behavior; MISB: moderate and slightly increasing sedentary behavior.

determinants of trajectories differed according to the PA components, confirming that PA involves complex and multidimensional behavior⁴. These determinants were mostly predictors of the frequency and type of PA trajectories. The lack of significance for the association with PA duration despite a significant association in the bivariate analysis may be due to the small sample size of the high and decreasing duration trajectory versus the moderate and stable trajectory (9% vs 91%), and future studies investigating this association are needed to confirm our results. Patients with impaired PROM and obese patients more often featured a lower prevalence of WB activities. Women more often showed a moderate and stable PA duration and lower and stable prevalence of WB activities. Predictors of moderate and higher screen-based SB trajectories were all sociodemographic, including male sex, age < 60 years, single status, obesity, smoking, and less physical jobs.

To our knowledge, this is the first study using longitudinal data to describe the trajectories of PA components and SB over a long time in a well-defined sample of people with hip and/or knee OA. This innovative method, group-based trajectory modeling, has been used in more recent studies investigating PA levels among the general adult population and in specific populations with heart disease and breast cancer, but has not been used in OA^{10,12,27}.

Our results agree with previous findings for older adults

showing the presence of stable and decreasing trajectories of PA, with stable trajectories more prevalent than change trajectories⁹. Our study adds to the existing knowledge by considering the different components of PA in patients with OA.

PA is associated with reduced pain sensitivity among healthy adults²⁸. Despite the potential long-term benefits of PA for patients with hip and/or knee OA, it is often challenging for them to regularly practice because their symptoms can worsen^{29,30}. Our results show that individuals with more impairment, including self-reported pain, functional abilities, and vitality, performed less WB PA. For most, this type of PA is unlikely to be recommended by a health professional because most of the KHOALA patients are followed up by primary care physicians who rarely give advice on types of PA. The perceived functional ability was also a predictor of the frequency of PA, with individuals with higher impairment practicing less frequently. In a systematic review, limited evidence was available to support the association between reduced functional impairment and lower level of PA in hip and knee OA. However, in this review, the PA level definition gave major weight to intensity; as well, the authors did not study PA components and the studies were cross-sectional³¹. However, results of studies describing PA trajectories in the general population have shown that patients with physical difficulties, disabilities, or poor self-rated health were less likely to follow a persistently active trajectory and more likely to follow a

low active or inactive trajectory⁹. Our results complement these studies by showing that inactivity or low activity in the context of OA results in a lower frequency of PA and less WB PA.

Previous studies describing PA trajectories in the general population have shown active trajectories are more prevalent among males than females³². Our results present new elements in patients with OA, showing that women more often practiced moderate and stable durations of PA than other frequencies of PA and RWB PA. Thus, proposing moderate duration and RWB PA could be appropriate for women and may be more successful. Obese patients more often showed a lower than higher prevalence of WB PA. In fact, RWB activities are selected by obese patients for being more tolerable than WB activities, which involve more loading on the joints³³.

Previous work has shown greater pain during PA among OA patients with higher levels of catastrophizing^{34,35}. Without measuring catastrophizing scores but with measuring related psychological measures (anxiety and depression), no relationship with PA components was found. Fatigue was the strongest predictor of reduced subsequent activity in patients with OA in 1 study³⁶. However, vitality, a close concept, did not differ between frequency, duration, or intensity of PA trajectories but was associated with increased frequency of WB PA.

None of the factors predicted the intensity trajectories. Intensity expressed by the metabolic equivalent MET may explain this result. Indeed, MET are defined by the Compendium of Physical Activities³⁷; the compendium does not consider the individual energy cost of PA, which differs especially according to sex and BMI, but instead standardizes the intensities of each PA. However, PA intensity is often of major importance in the outcome criteria chosen to study factors associated with PA level.

SB is detrimental to health, even when recommended levels of PA are reached¹⁵. Predictors of moderate and higher screen-based SB trajectories were all sociodemographic (male sex, age < 60 years, single status, obesity, smoking, and less physical jobs). Some of these characteristics were similar to determinants of a sedentary lifestyle in older people without OA³⁸ and to those of inactive trajectories in the general population³⁹. Our PROM were not associated with screen-based SB trajectories. For older women, mobility impairment, depression, and lack of energy were associated with SB¹⁰. In a previous OA cross-sectional study, SB was related to worse physical function, but no longitudinal studies have explored the effect of PROM impairment on SB⁴⁰. Thus, symptoms and clinical severity do not seem to prevent individuals from being less sedentary.

Our study has limitations. PA was assessed with a self-reporting questionnaire, which may have introduced some measurement error. Although in general, our results show compliance with recommendations for PA components, results from other studies using accelerometers showed reduced levels of PA⁴¹. However, accelerometers are not appropriate for long-term measurements, and they are not suitable to measure some PA such as water-based activities⁴². As well, SB measured only the

time spent in front of the TV and computer and may therefore underestimate the real sedentary time of patients. However, SB is not a single construct; it involves different types of behavior, and screen time is the most-used type of SB studied.

Identifying diverse trajectory groups using group-based trajectory modeling adds to previous knowledge by providing evidence of the heterogeneity of PA and SB. Thus, this study allowed for understanding the variation that occurs in the frequency, duration, intensity, and types of PA and SB during a 5-year follow-up, as well as the factors relating to belonging to specific trajectory classes. Our results allowed for identifying sociodemographic and clinical predictors, mainly of PA frequency and type of PA trajectories, as well as sociodemographic predictors of SB trajectories. This information can help in planning tailored and well-targeted PA promotion strategies and interventions for the OA population, especially those who are sedentary.

REFERENCES

1. Felson DT, Lawrence RC, Dieppe PA, Hirsch R, Helmick CG, Jordan JM, et al. Osteoarthritis: new insights. Part 1: the disease and its risk factors. *Ann Intern Med* 2000;133:635-46.
2. Vos T, Allen C, Arora M, Barber RM, Bhutta ZA, Brown A, et al. Global, regional, and national incidence, prevalence, and years lived with disability for 310 diseases and injuries, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015. *Lancet* 2016;388:1545-602.
3. American Geriatrics Society Panel on Exercise and Osteoarthritis. Exercise prescription for older adults with osteoarthritis pain: consensus practice recommendations. A supplement to the AGS Clinical Practice Guidelines on the management of chronic pain in older adults. *J Am Geriatr Soc* 2001;49:808-23.
4. Barbour KE, Lui LY, Nevitt MC, Murphy LB, Helmick CG, Theis KA, et al. Hip osteoarthritis and the risk of all-cause and disease-specific mortality in older women: a population-based cohort study. *Arthritis Rheumatol* 2015;67:1798-805.
5. de Groot IB, Bussmann JB, Stam HJ, Verhaar JA. Actual everyday physical activity in patients with end-stage hip or knee osteoarthritis compared with healthy controls. *Osteoarthritis Cartilage* 2008;16:436-42.
6. Farr JN, Going SB, Lohman TG, Rankin L, Kastle S, Cornett M, et al. Physical activity levels in patients with early knee osteoarthritis measured by accelerometry. *Arthritis Rheum* 2008;59:1229-36.
7. Rosemann T, Kuehlein T, Laux G, Szecsenyi J. Factors associated with physical activity of patients with osteoarthritis of the lower limb. *J Eval Clin Pract* 2008;14:288-93.
8. Verlaan L, Bolink SN, Van Laarhoven SN, Lipperts M, Heyligers IC, Grimm B, et al. Accelerometer-based physical activity monitoring in patients with knee osteoarthritis: objective and ambulatory assessment of actual physical activity during daily life circumstances. *Open Biomed Eng J* 2015;9:157-63.
9. Lounassalo I, Salin K, Kankaanpää A, Hirvensalo M, Palomäki S, Tolvanen A, et al. Distinct trajectories of physical activity and related factors during the life course in the general population: a systematic review. *BMC Public Health* 2019;19:271.
10. Xue QL, Bandeen-Roche K, Mielenz TJ, Seplaki CL, Szanton SL, Thorpe RJ, et al. Patterns of 12-year change in physical activity levels in community-dwelling older women: can modest levels of physical activity help older women live longer? *Am J Epidemiol* 2012;176:534-43.

11. Demmelmaier I, Dufour AB, Nordgren B, Opava CH. Trajectories of physical activity over two years in persons with rheumatoid arthritis. *Arthritis Care Res* 2016;68:1069-77.
12. Blanchard CM, McSweeney J, Giacomantonio N, Reid RD, Rhodes RE, Spence JC, et al. Distinct trajectories of light and moderate to vigorous physical activity in heart disease patients: results from the Activity Correlates of Ter cardiac hospitalization (ACTION) trial. *J Sci Med Sport* 2014;17:72-7.
13. de Rooij M, van der Leeden M, Heymans MW, Holla JFM, Häkkinen A, Lems WF, et al. Prognosis of pain and physical functioning in patients with knee osteoarthritis: a systematic review and meta-analysis. *Arthritis Care Res* 2016;68:481-92.
14. Guillemin F, Rat AC, Roux CH, Fautrel B, Mazieres B, Chevalier X, et al. The KHOALA cohort of knee and hip osteoarthritis in France. *Joint Bone Spine* 2012;79:597-603.
15. Guillemin F, Rat AC, Mazieres B, Pouchot J, Fautrel B, Euler-Ziegler L, et al. Prevalence of symptomatic hip and knee osteoarthritis: a two-phase population-based survey. *Osteoarthritis Cartilage* 2011;19:1314-22.
16. Vuillemin A, Oppert JM, Guillemin F, Essermeant L, Fontvieille AM, Galan P, et al. Self-administered questionnaire compared with interview to assess past-year physical activity. *Med Sci Sports Exerc* 2000;32:1119-24.
17. Bruyere O. Both weight-bearing and non-weight-bearing exercise improved function in patients with knee osteoarthritis. *Evid Based Med* 2009;14:178.
18. Groll DL, To T, Bombardier C, Wright JG. The development of a comorbidity index with physical function as the outcome. *J Clin Epidemiol* 2005;58:595-602.
19. Bellamy N, Buchanan WW, Goldsmith CH, Campbell J, Stitt LW. Validation study of WOMAC: a health status instrument for measuring clinically important patient relevant outcomes to antirheumatic drug therapy in patients with osteoarthritis of the hip or knee. *J Rheumatol* 1988;15:1833-40.
20. Rat AC, Coste J, Pouchot J, Baumann M, Spitz E, Retel-Rude N, et al. OAKHQOL: a new instrument to measure quality of life in knee and hip osteoarthritis. *J Clin Epidemiol* 2005;58:47-55.
21. Bunevicius A. Reliability and validity of the SF-36 Health Survey Questionnaire in patients with brain tumors: a cross-sectional study. *Health Qual Life Outcomes* 2017;15:92.
22. Goldberg DP, Hillier VF. A scaled version of the General Health Questionnaire. *Psychol Med* 1979;9:139-45.
23. The World Health Organization Quality of Life assessment (WHOQOL): position paper from the World Health Organization. *Soc Sci Med* 1995;41:1403-9.
24. Jones BL, Nagin DS, Roeder K. A SAS Procedure Based on Mixture Models for Estimating Developmental Trajectories. *Sociol Methods Res* 2001;29:374-93.
25. Nagin DS, Odgers CL. Group-based trajectory modeling in clinical research. *Annu Rev Clin Psychol* 2010;6:109-38.
26. Schwartz T, Chen Q, Duan N. 339-2011: Studying missing data patterns using a SAS macro. [Internet. Accessed August 18, 2020.] Available from: support.sas.com/resources/papers/proceedings11/339-2011.pdf
27. Brunet J, Amireault S, Chaiton M, Sabiston CM. Identification and prediction of physical activity trajectories in women treated for breast cancer. *Ann Epidemiol* 2014;24:837-42.
28. Vaegter HB, Handberg G, Graven-Nielsen T. Similarities between exercise-induced hypoalgesia and conditioned pain modulation in humans. *Pain* 2014;155:158-67.
29. Wideman TH, Finan PH, Edwards RR, Quartana PJ, Buenaver LF, Haythornthwaite JA, et al. Increased sensitivity to physical activity among individuals with knee osteoarthritis: relation to pain outcomes, psychological factors, and responses to quantitative sensory testing. *Pain* 2014;155:703-11.
30. Baxter SV, Hale LA, Stebbings S, Gray AR, Smith CM, Tretharve GJ. Walking is a feasible physical activity for people with rheumatoid arthritis: a feasibility randomized controlled trial. *Musculoskeletal Care* 2016;14:47-56.
31. Veenhof C, Huisman PA, Barten JA, Takken T, Pisters MF. Factors associated with physical activity in patients with osteoarthritis of the hip or knee: a systematic review. *Osteoarthritis Cartilage* 2012;20:6-12.
32. Barnett TA, Gauvin L, Craig CL, Katzmarzyk PT. Distinct trajectories of leisure time physical activity and predictors of trajectory class membership: a 22 year cohort study. *Int J Behav Nutr Phys Act* 2008;5:57.
33. Jan MH, Lin CH, Lin YF, Lin JJ, Lin DH. Effects of weight-bearing versus nonweight-bearing exercise on function, walking speed, and position sense in participants with knee osteoarthritis: a randomized controlled trial. *Arch Phys Med Rehabil* 2009;90:897-904.
34. Estévez-López F, Álvarez-Gallardo IC, Segura-Jiménez V, Soriano-Maldonado A, Borges-Cosic M, Pulido-Martos M, et al. The discordance between subjectively and objectively measured physical function in women with fibromyalgia: association with catastrophizing and self-efficacy cognitions. The al-Ándalus project. *Disabil Rehabil* 2018;40:329-337.
35. Vincent HK, George SZ, Seay AN, Vincent KR, Hurley RW. Resistance exercise, disability, and pain catastrophizing in obese adults with back pain. *Med Sci Sports Exerc* 2014;46:1693-701.
36. Murphy SL, Alexander NB, Levoska M, Smith DM. Relationship between fatigue and subsequent physical activity among older adults with symptomatic osteoarthritis. *Arthritis Care Res* 2013;65:1617-24.
37. Ainsworth BE, Haskell WL, Herrmann SD, Meckes N, Bassett DR, Tudor-Locke C, et al. 2011 Compendium of Physical Activities: a second update of codes and MET values. *Med Sci Sports Exerc* 2011;43:1575-81.
38. Heseltine R, Skelton DA, Kendrick D, Morris RW, Griffin M, Haworth D, et al. "Keeping Moving": factors associated with sedentary behaviour among older people recruited to an exercise promotion trial in general practice. *BMC Fam Pract* 2015;16: 67.
39. Aggio D, Papachristou E, Papacosta O, Lennon LT, Ash S, Whincup PH, et al. Trajectories of self-reported physical activity and predictors during the transition to old age: a 20-year cohort study of British men. *Int J Behav Nutr Phys Act* 2018;15:14.
40. Lee J, Chang RW, Ehrlich-Jones L, Kwok CK, Nevitt M, Semanik PA, et al. Sedentary behavior and physical function: objective evidence from the Osteoarthritis Initiative. *Arthritis Care Res* 2015;67:366-73.
41. Dunlop DD, Song J, Semanik PA, Chang RW, Sharma L, Bathon JM, et al. Objective physical activity measurement in the osteoarthritis initiative: are guidelines being met? *Arthritis Rheum* 2011;63:3372-82.
42. Skender S, Ose J, Chang-Claude J, Paskow M, Brühmann B, Siegel EM, et al. Accelerometry and physical activity questionnaires - a systematic review. *BMC Public Health* 2016;16:515.