

EXTENDED REPORT

Rhythmic variations in pain, stiffness, and manual dexterity in hand osteoarthritis

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Objective: To explore circadian variation in pain, stiffness, and manual dexterity in patients with hand osteoarthritis (OA).

Methods: Twenty one patients with hand OA, as defined by ACR criteria (17 women, four men, mean age 62.2 years, range 52–74 years) self rated pain and stiffness on separate 10 cm horizontal visual analogue scales and performed bead intubation coordinometry (BIC) six times each day (on waking up, at bedtime, and every four hours in between) for 10 consecutive days. Each series (using data with the trend removed if there was a significant trend) was analysed for circadian rhythmicity by a cosine vector technique (single cosinor). With individual data expressed as the percentage of the mean, group rhythm characteristics at period 24 hours were summarised for each variable by population mean cosinor analysis.

Results: Individual analyses identified significant circadian rhythms at $p \leq 0.05$ for pain ($n=15/21$), stiffness ($n=16/20$), and dexterity ($n=18/21$), and a significant circadian rhythm on a group basis was identified for pain ($p=0.013$), stiffness ($p<0.001$), and dexterity ($p<0.001$). Pain was least at 1610 and stiffness at 1618. Peak dexterity occurred in mid-afternoon at 1548 and occurred within the 95% confidence interval of least pain (1312–1800) and stiffness (1520–1732).

Conclusions: Dexterity was influenced by the patient's level of pain or stiffness, which changed systematically throughout the day. Similar results have been previously reported in 14 patients with rheumatoid arthritis where peak dexterity occurred at 1544 and at 1528 in 14 age and sex matched healthy controls. The predictability of rhythmic variation in pain, stiffness, and dexterity has implications for scheduling activities of daily living and for timing antirheumatic drug treatment.

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Circadian variation is common in biological systems. Plants, animals, and humans display rhythmic variation at various levels, from the cell to the entire organism. Rhythms that cycle about once a day are called circadian rhythms,¹ those of higher or lower frequency being respectively termed ultradian and infradian rhythms. Several investigators have reported circadian variation in clinical and laboratory variables in patients with either inflammatory or degenerative disorders of the musculoskeletal system.^{2–30} In general, these investigators have used traditional statistical methods to compare within-day variations.³¹ For some years, however, statistical methods based on least squares and cosine vector techniques, which permit the use of real time data, have been used successfully by chronobiologists in mapping the absolute and relative timing of different biological rhythms.^{32–34} We have used such techniques in successfully identifying circadian rhythms in pain, stiffness, and manual dexterity in rheumatoid arthritis (RA),² and knee osteoarthritis (OA) pain.³ Circadian characteristics for these studies and more than 100 variables found in the literature that are pertinent to immune function and disease have been recently summarised in tables.³⁵ The identification of such rhythms has important implications for patients, both in planning their daily activities, and in developing individual therapeutic programmes in relation to circadian variability, which may be more effective (so-called “chronotherapy” programmes).

MATERIALS AND METHODS

A quasiexperimental one group repeated measures design was used. Patients were followed up for 10 consecutive days and made self measurements at home at prespecified time points. The protocol was approved by the University of Western Ontario review board for research involving human subjects.

Patients were recruited by telephone. Consenting outpatients with symptomatic hand OA in the dominant hand were enrolled in the study. To be eligible all patients had to be symptomatic—that is, to have had pain for at least three months in the dominant hand, fulfil American College of Rheumatology (ACR) classification criteria for hand OA,³⁶ be aged between 40 and 75 years, be capable of providing informed consent, be fluent in the English language, and, if taking NSAIDs, have reached a steady state (at least five days of treatment and would continue on the same agent for the duration of the study). Exclusion criteria were amputation of one or more digits, paralysis of either hand, comorbid condition causing pain, stiffness, or functional disability in the hands, pregnant or lactating women, prior orthopaedic surgery on either hand, concomitant musculoskeletal condition affecting the hands, patients requiring the use of narcotic class analgesics, unable to comprehend the visual analogue measurement scales (VAS) or perform bead intubation coordinometry (BIC) after adequate explanation, unavailable to perform self measurement for the next 10 days, requiring intra-articular steroid injections in the next 10 days, and starting physiotherapy in the next 10 days.

At baseline, data collected were age, sex, disease duration in the hands, hand dominance, ACR classification criteria, AUSCAN osteoarthritis hand index score,³⁷ grip strength,³⁸ pinch grip, distribution of hand involvement (for example,

Abbreviations: ACR, American College of Rheumatology; ANOVA, analysis of variance; BIC, bead intubation coordinometry; 95% CI, 95% confidence interval; OA, osteoarthritis; NSAIDs, non-steroidal anti-inflammatory drugs; RA, rheumatoid arthritis; ROC, range of change; VAS, visual analogue scale

homunculus), Doyle index,³⁹ information on current drugs (for example, dose and schedule). Patients self measured pain, stiffness, and manual dexterity each day on waking up and bedtime and about every four hours in between (target times 1000, 1400, 1800, 2200). Patients were instructed specifically to look at a watch or clock and record the actual time (real time) at which they performed self measurement, as it was unlikely over the 10 day period that the measurements could be performed exactly at the time specified. The collection of real time data was critical to the success of the analysis because of the potentially dynamic fluctuation which could occur in the three variables. In reality, some self measurements were performed at nearly every hour of the day and night.

Pain was self rated on a 10 cm horizontal VAS with end markers outside of which were placed terminal descriptors (left=no pain, right=most severe pain I have ever had).^{2,3} Patients were instructed to rate the severity of the pain in their hand at the exact moment of measurement in response to the question, "How much pain are you currently experiencing due to osteoarthritis in your dominant hand?"

Stiffness was self rated on a 10 cm horizontal VAS with end markers, outside of which were placed terminal descriptors (left=no stiffness, right=most severe stiffness I have ever had).^{2,3} Patients were instructed to rate the severity of the stiffness in their hand at the exact moment of measurement in response to the question, "How much stiffness are you currently experiencing due to osteoarthritis in your dominant hand?"

Manual dexterity was measured using a device termed a bead intubation coordinometer (BIC, fig 1).³² This consists of a shallow circular aluminium receptacle with a flat base (11.5 cm) and a vertical side (4.0 cm). The top is closed by a Perspex lid, carrying at its centre, a hollow vertical (8 cm) stainless steel tube. The calibre of the tube is such that it will only accept a series of sand blasted stainless steel precision beads (0.65×0.48 cm) when they are presented with their long axis exactly aligned with the axis of the tube. Twenty five such beads are placed in a shallow plastic bowl to one side (determined by hand dominance) of the coordinometer, and a precision stopwatch is placed on the other side at a distance comfortable to the patient. The test is started with the patient's dominant hand holding the first bead, while the non-dominant hand is on the stopwatch. Simultaneously, the watch is started and bead intubation started. Thereafter, the beads are inserted as fast as possible, one by one, into the tube. Immediately after the insertion of the last bead, the non-dominant hand stops the watch. Patients were instructed that the beads were to be picked up, one at a time, and if a bead was dropped on the table or the floor, the test had to be restarted. The BIC instrument used in this study is a modification of one previously used by one of us (NB) in a field trial in northwest Greenland.⁴⁰ The modifications made were as follows: (a) the base of the coordinometer was enlarged to allow the osteoarthritic hand a better grip, and (b) the beads were sand blasted to allow a better grip when lifting them out of the bowl. Pilot studies with this instrument have suggested that a learning effect occurs in the first 48–72 hours, and that 25 beads produce satisfactory data without tiring the patient.

These exact instruments were used in our previous work on circadian rhythmicity in manual dexterity in patients with RA and proved to be entirely satisfactory.² Subjects were instructed to complete the questionnaire and BIC at or near the time points indicated, and, in any event, record accurately the actual clock time when the observation was made. Subjects waking up after 1000 or going to bed before 2200 would sometimes have fewer than the six designated observation points each day.

Patients were requested to note at the end of the day, the time, dose, and brand of any analgesics taken for any purpose during that day.

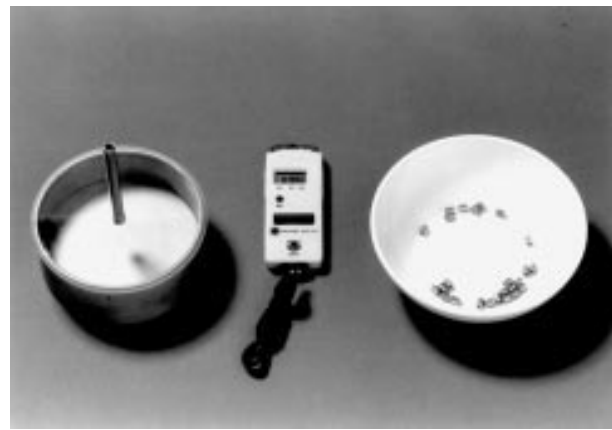


Figure 1 The bead intubation coordinometer.

Time series plots were constructed for each raw data series to view the data for patterns, trends, and outliers. No data were excluded from analysis. If any series showed a significant trend, the trend was statistically removed before analysis for time effects. Individual data series were converted to a percentage of the mean value before constructing overall waveforms for circadian frequencies for the three variables of interest and performing grouped time series analyses.^{33,34} A one way analysis of variance (ANOVA) was used to test grouped data for the effect of time. Each series was analysed for circadian rhythmicity by the least squares method of fitting of cosines (single cosinor method^{33,34}) with periods in the expanded circadian range 14–34 hours,⁴¹ and with 0.1 hours between trial periods. Group rhythm characteristics at precisely 2400 were summarised by population mean cosinor analysis.^{33,34} Rhythm detection was considered statistically significant for $p \leq 0.05$ from a zero amplitude test and borderline significant for $0.05 < p \leq 0.10$. Rhythm characteristics estimated from the single cosinor procedure included the mesor (middle of the fitted cosine, representing a rhythm adjusted mean when data are collected at uneven intervals), the amplitude (the distance from the mesor to the peak or trough (bathypase) of the fitted cosine), and the acrophase (peak of the fitted cosine with reference to local midnight). Because our interest was in peak performance, results were expressed using the bathypase rather than the acrophase, which is the clock time at which pain and stiffness are least and dexterity best. Correlations were also made between 24 hour averages (mesors) of each variable.

RESULTS

Twenty one patients (17 women and four men) with OA of the hand, as defined by the ACR criteria, provided data on self rated pain and stiffness and performed BIC over 10 days (table 1). Mean age was 62.2 years; range 52–74 years. All patients were right handed. Mean grip strength for the dominant hand, over three attempts, was 156 mm Hg. The mean pinch grip for the dominant hand, over three attempts, was 4.7 kg (table 1).

The study was completed over a three month span, patients starting their data collection on different days of the week. All patients were in a steady state: three regularly receiving NSAIDs only, five receiving NSAIDs on a regular schedule and supplementary analgesia with acetaminophen as required, seven taking only analgesics on an "as required" basis, and six having neither NSAIDs nor analgesic treatment (table 1). None of the subjects were taking exogenous steroids for their arthritis or other comorbidities. All patients slept at night, with the average bedtime at 2330 (range 2100–0355) and waking up at 0730 (range 0415–1000). None of the

Table 1 Demographic and disease characteristics

PIP	DIP	CMC	Unilateral/ Bilateral	Patient	Sex	Age	Dominant hand		NSAID medication	Analgesic medication
							Mean grip strength (mm Hg)	Mean pinch strength (kg)		
+	+	-	B	1R	F	52	165	5.5	Nil	Nil
+	-	+	B	3R	F	53	133	2.5	Ibuprofen	Nil
+	+	+	B	4R	F	74	200	4.5	Nyprosyn	Tylenol 2
+	+	+	B	5R	F	62	206	4	Ketoprofen	Nil
+	-	-	B	6R	F	54	123	1.8	Nil	Tylenol 2
+	+	+	B	7R	F	64	174	5.3	Nil	Tylenol
+	+	-	B	8R	F	53	157	5.8	Nil	Nil
+	+	+	B	9R	F	58	89	4.3	Surgam	Tylenol
+	+	+	B	10R	F	69	130	3.7	Nil	Tylenol
+	-	+	B	11R	F	62	94	3.7	Nil	Tylenol
+	+	+	B	15R	F	69	103	4.2	Nil	Tylenol
+	+	-	B	16R	F	69	160	4	Nil	Tylenol
+	-	+	B	17R	F	66	187	4	Nil	Nil
+	+	+	B	18R	F	58	87	4.8	Arthrotec	Tylenol
-	-	+	B	19R	F	57	187	3	Nil	Nil
+	+	+	B	20R	F	59	43	1.7	ASA	Tylenol 3
+	+	+	B	21R	F	71	155	4.7	Ibuprofen	None
+	+	+	B	2R	M	70	189	6.2	Nil	Nil
+	+	+	B	12R	M	62	123	8.2	Nil	Tylenol 3
+	+	+	B	13R	M	71	300+	9	Flurbiprofen	Tylenol 3
+	+	-	B	22R	M	53	273	8.8	Nil	Nil

participants were involved in night work or shift work in the period of the study and none had indulged in transmeridional travel during the course of the study.

Figure 2 shows the overall circadian patterns for pain, stiffness, and dexterity along with the best fitting 24 hour cosine. From data normalised to percentage of the mean, a significant time effect was found by ANOVA for each variable at $p < 0.001$ (table 2).

Pain scores varied from 0 to 93 and 24 hour mean pain values from 1.0 to 70.2 among 21 subjects. All subjects showed within 24 hour variability with a mean range of change (ROC) from lowest to highest values of 42.4 (individual ranges were from 4 to 82). On an individual basis, single cosinor analysis found a significant circadian rhythm at $p \leq 0.05$ for 15 subjects and at $0.05 < p \leq 0.10$ for 17 of the 21 subjects (table 3). Summary by population mean cosinor described a significant circadian rhythm for the group at $p = 0.013$, with an amplitude of 13% and the bathyphase at 1600 (table 2).

One subject failed to complete the stiffness scales. In the remaining 20 subjects, stiffness scores varied from 0 to 96 and

24 hour mean stiffness values from 4.8 to 78.9. All 20 subjects showed within 24 hour variability with a mean ROC of 47.4 (individual ranges 9 to 85). On an individual basis, single cosinor analysis showed a significant circadian rhythm at $p \leq 0.05$ for 16 out of 20 subjects (table 3). Summary by population mean cosinor showed a significant circadian rhythm for the group at $p < 0.001$, with an amplitude of 22% and the bathyphase at 1618 (table 2).

Bead intubation times varied from 40 to 190 seconds and 24 hour mean BIC values varied from 47.1 to 109.0 seconds among 21 subjects. All subjects showed within 24 hour variability with a mean ROC of 46.8 (individual ranges were from 10 to 118 seconds). On an individual basis, single cosinor analysis found a significant circadian rhythm at $p \leq 0.05$ for 18 subjects and at $0.05 < p \leq 0.10$ for 20 of the 21 subjects (table 3). Summary by population mean cosinor described a significant circadian rhythm for the group at $p < 0.001$, with an amplitude of 8% and the bathyphase at 1548 (table 2).

The 24 hour means (mesors) for pain and stiffness showed a positive correlation between each other ($r = 0.76, p < 0.001$),

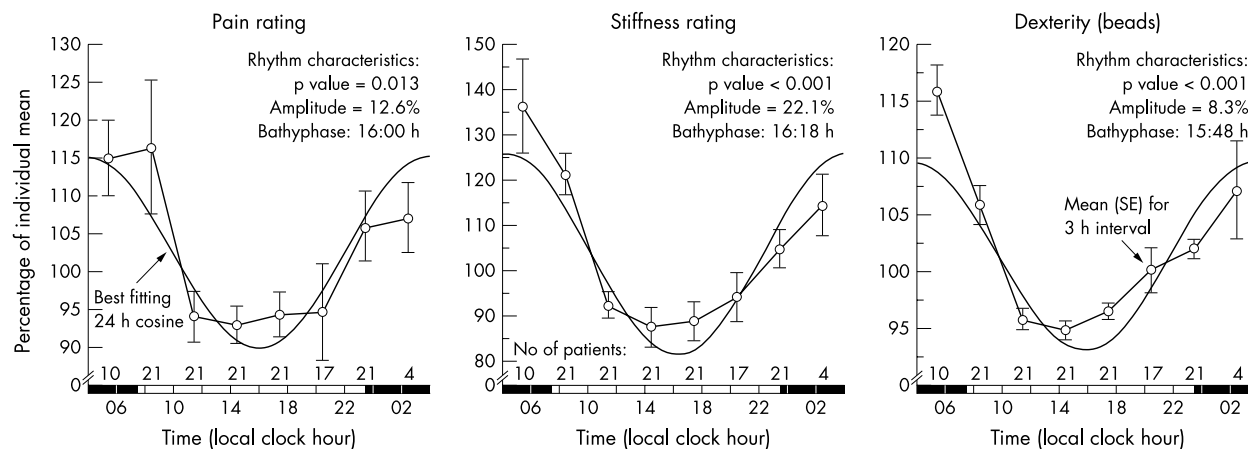


Figure 2 Circadian rhythm for pain, stiffness, and dexterity in patients with osteoarthritis of the hand. Self measurements/ratings were made by 20 or 21 patients every 24 hours during waking for 10 days. Individual values had trends removed and were converted to a percentage of the mean before combining for group analysis by population mean cosinor. For rhythm characteristics p value is from the zero amplitude test; amplitude=half peak trough difference of cosine; bathyphase=lowest point of cosine (referenced from 0000). p Values < 0.001 for each variable from ANOVA for time effect.

Table 2 Group circadian rhythm characteristics for pain rating, stiffness rating, and dexterity

Time effect:	Units analysed:																
	Original									% of mean							
	ANOVA				Cosine period=24.0 h					ANOVA				Cosine period=24.0 h			
Variable	n	F	p Value	p Value	M (SE)	A (SE)	(%A)	bØ (h)	95% CI (h)	F	p Value	p Value	M (SE)	A (SE)	(%A)	bØ (h)	95% CI (h)
Pain	21	1.4	0.217	0.021	31.2 (5.3)	2.9 (1.1)	(9)	16:50	(12:28 to 18:56)	4.6	<0.001	0.013	102 (1)	13 (4)	(12)	16:00	(13:12 to 18:00)
Stiffness	20	5.4	<0.001	<0.001	37.6 (5.3)	6.7 (1.3)	(18)	16:13	(15:08 to 17:16)	22.2	<0.001	<0.001	104 (1)	22 (4)	(21)	16:18	(15:20 to 17:32)
BIC	21	9.4	<0.001	<0.001	75.9 (3.6)	6.4 (0.9)	(8)	15:41	(14:56 to 16:24)	44.2	<0.001	<0.001	101 (0)	8 (1)	(8)	15:48	(15:00 to 16:28)

Data obtained by self measurement every 2–4 h during waking only for 10 days. Any series with a significant trend had the trend removed before analysis for circadian rhythm by the least squares fit of a 24.0 h cosine. Circadian characteristics (M, A, bØ) summarised by population mean cosinor analysis. For rhythm characteristics: p value from zero amplitude test; M = mesor (middle value of fitted cosine); A = amplitude (distance from M to peak or trough of cosine); %A = % amplitude (A as % of M); bØ = bathyphase (lowest point of fitted cosine, referenced from 00:00 h); ANOVA = analysis of variance across six 4 hourly intervals.

Table 3 Individual circadian rhythm characteristics for pain rating, stiffness rating, and dexterity

Patient/ sex	Variable:																					
	Pain rating					Best fitting				Stiffness rating				Best fitting				Dexterity			Best fitting	
	N	Fitted period = 24.0 h				Circadian period				Fitted period=24.0 h				Circadian period				Fitted period=24.0 h			Circadian period	
	p Value	M	A	(%A)	bØ (h)	Hours	p Value	p Value	M	A	(%A)	bØ (h)	Hours	p Value	p Value	M	A	(%A)	bØ (h)	Hours	p Value	
1F	60	0.150	14.2	5.1	(36)	15:42	23.9	0.147	<0.001	18.9	14.0	(74)	15:44	24.1	<0.001	<0.001	70.2	6.5	(9)	15:36	23.7	<0.001
3F	49	<0.001	67.7	7.8	(12)	07:08	23.8	<0.001	<0.001	48.0	18.2	(38)	16:14	24.2	<0.001	<0.001	55.4	6.4	(12)	16:37	23.8	<0.001
4F	60	0.020	2.2	1.3	(60)	17:18	24.3	0.012	0.015	12.6	4.2	(34)	17:59	24.6	0.001	<0.001	67.3	8.2	(12)	15:55	24.0	<0.001
5F	52	0.786	2.3	0.1	(5)	05:12	22.3	0.049	0.041	5.0	1.1	(21)	15:50	24.5	0.012	0.022	64.3	4.2	(7)	15:02	24.7	<0.001
6F	44	0.035	21.2	7.7	(36)	13:10	24.1	0.035	0.005	38.7	17.6	(45)	15:42	23.4	<0.001	0.259	113.8	13.0	(11)	14:49	23.7	0.224
7F	60	0.214	3.5	0.9	(27)	12:44	23.2	0.057	0.327	71.3	1.0	(1)	17:52	20.9	0.112	0.323	70.8	2.7	(4)	17:03	21.6	0.045
8F	59	0.191	6.0	0.4	(7)	12:02	23.0	0.124	0.721	8.0	0.2	(3)	10:38	26.8	0.330	<0.001	47.1	1.7	(4)	09:30	23.5	<0.001
9F	46	0.007	61.7	6.2	(10)	17:02	23.8	0.005	0.126	79.6	4.4	(6)	14:46	24.7	0.017	<0.001	64.3	7.7	(12)	16:05	24.0	<0.001
10F	60	0.138	9.1	2.7	(30)	16:51	23.6	0.100	0.042	17.9	5.2	(29)	14:48	22.8	0.003	<0.001	95.2	10.4	(11)	15:04	23.8	<0.001
11F	60	<0.001	48.8	15.8	(32)	20:32	24.2	<0.001	<0.001	40.0	10.7	(27)	18:48	24.2	<0.001	<0.001	75.3	10.2	(14)	17:15	24.3	<0.001
15F	48	<0.001	40.6	13.6	(34)	18:34	23.9	<0.001	0.011	50.0	13.2	(26)	18:26	25.2	<0.001	<0.001	57.9	7.3	(13)	17:22	24.1	<0.001
16F	58	0.367	36.1	1.9	(5)	01:53	21.8	0.013	–	–	–	–	–	–	–	0.102	77.7	1.8	(2)	14:02	22.4	0.011
17F	56	0.303	13.7	2.2	(16)	22:44	24.6	0.252	0.219	12.3	2.4	(19)	23:33	24.8	0.110	0.088	71.9	2.3	(3)	09:38	23.6	0.069
18F	43	0.050	52.0	8.1	(15)	13:30	23.6	0.023	<0.001	43.0	13.5	(31)	13:33	24.1	<0.001	0.002	102.0	17.8	(17)	14:54	24.1	0.002
19F	50	0.200	11.0	2.7	(24)	10:15	23.2	0.128	0.006	17.5	5.7	(32)	14:19	23.7	0.002	0.095	81.6	7.0	(9)	15:02	23.2	0.005
20F	52	0.002	71.1	7.8	(11)	10:14	24.5	<0.001	0.005	72.7	6.3	(9)	11:56	24.6	<0.001	0.042	103.6	5.9	(6)	11:22	23.3	0.030
21F	60	<0.001	66.2	16.6	(25)	18:01	23.8	<0.001	<0.001	60.3	14.1	(23)	17:39	23.7	<0.001	<0.001	76.4	11.6	(15)	17:46	24.0	<0.001
2M	60	0.022	45.8	4.0	(9)	20:42	23.4	0.014	0.215	52.1	4.4	(8)	16:24	23.8	0.200	0.159	82.7	4.7	(6)	15:16	23.1	0.050
12M	60	0.020	49.2	3.1	(6)	18:10	23.7	0.017	<0.001	59.8	10.0	(17)	16:27	23.9	<0.001	<0.001	73.4	8.1	(11)	16:18	23.8	<0.001
13M	60	0.088	27.2	4.4	(16)	14:46	24.8	0.002	0.034	30.0	5.2	(17)	14:43	24.8	<0.001	0.162	67.7	2.6	(4)	14:39	26.9	0.059
22M	55	0.790	5.6	0.7	(12)	07:32	32.3	0.023	0.049	13.8	4.3	(31)	19:27	23.5	0.022	<0.001	76.4	7.1	(9)	17:28	23.8	<0.001

Data obtained by self measurement every 2–4 h during waking only for 10 days. Any series with a significant trend had the trend removed before analysis for circadian rhythm by the least squares fit of a 24.0 h cosine. Each series was also analysed for best fitting cosine in the circadian domain between 14 and 34 h. For rhythm characteristics: p value from zero amplitude test; M = mesor (middle value of fitted cosine); A = amplitude (distance from M to peak or trough of cosine); %A = % amplitude (A as % of M); bØ = bathyphase (lowest point of fitted cosine, referenced from 00:00h).

indicating that these two measures were proportional within each patient (for example, as one rating increased, so did the other). There was also an inverse correlation between rhythm detection p values and the mesor for pain ($r=-0.58$, $p=0.006$) and stiffness ($r=-0.56$, $p=0.010$), indicating the likelihood of an increased chance of rhythm detection with increasing levels of either of these two ratings (for example, a lower p value was obtained when pain or stiffness were more severe).

DISCUSSION

Pain, stiffness, and physical function are important measures of hand OA. In this study we used conventional 10 cm visual analogue scales to measure pain and stiffness, and a novel instrument (BIC), not previously used in the assessment of patients with hand OA. Performance using the BIC may be altered by the presence of musculoskeletal or neurological disease. However, the patients in this study had no clinical evidence of neurological disease, and thus performance decrements were attributable to the presence of articular disease. As such, BIC became a measure of dominant hand dexterity.

In this study we mapped daily changes in pain, stiffness, and BIC in a group of patients with OA who are typical of those recruited for NSAID studies. It is of note that variability in most, but not all, patients with OA was not random but instead showed a significant adherence to a quasisinusoidal rhythm having a period of about 24 hours. The data suggest that the level of pain, stiffness, and dexterity throughout the day are in part predictable. Furthermore, we noted that dexterity is best about the time that pain and stiffness are least—that is, in the late afternoon.

Analgesic usage data were not interpretable and Environment Canada statistics on temperature and humidity were not obtained. We recognise that the modulating effects of these variables on circadian variation cannot be fully dissected without such information. However, in a previous study of circadian and circaseptan rhythms in pain perception in knee OA we were unable to show any significant relation between fluctuations in pain scores and either analgesic consumption or the aforementioned biometeorological factors.³

We have two prior experiences of using BIC measurement. The first was in a study of 12 medical students in northwest Greenland.⁴⁰ In that study no significant rhythm was detected in BIC, but the field conditions were severe with low ambient temperature, dry skin, and the polished surface of the beads contributing to the difficulty of performing the task. We think that sand blasting the beads was a major contribution to our success in detecting rhythmicity in this study. Our second experience was in hand RA where we noted that manual dexterity in healthy controls was characterised by a rhythm of lower level and smaller amplitude than in patients with RA.² The lower level was accounted for by the faster speed at which healthy subjects completed the task. The smaller amplitude was probably due to the absence of the modulating influences of pain, stiffness, and other aspects of the inflammatory process, which may have made it more difficult for patients with RA to perform BIC at some times of day than others. Because peak dexterity in patients with RA and controls occurred at almost the same time, we postulated that the dexterity profile of patients with RA merely represented a vertical distortion of the normal dexterity rhythm found in healthy controls, and that the distortion might have been mediated by the modulating influences of joint pain and stiffness. In the OA study reported here, we have also found synchronicity between pain, stiffness, and dexterity rhythms. While recognising the limitations of interstudy comparisons, it is of note that peak dexterity in hand OA (1548), RA (1544), and healthy controls (1528) in the two studies occurred within 20 minutes of one another. This reinforces the contention that arthritis symptoms of pain and stiffness vertically distort the level of

performance, but do not horizontally distort the normal phase of the dexterity rhythm.

There are three ways in which the results of this study may find practical application. Firstly, as most subjects show circadian rhythmicity in manual dexterity, and this is consistent from day to day, patients with hand OA may find advantage in planning those activities requiring manual dexterity during the period of the day when manual dexterity is least affected by their arthritis. Secondly, in clinical trials, knowledge of circadian rhythmicity in pain, stiffness, and dexterity can be used to inform protocol development, and both the scheduling and structuring of assessments for these three variables. Finally, the existence of circadian rhythmicity may provide an opportunity for an approach to treatment with analgesic and anti-inflammatory class compounds, using the principles of chronotherapy, where the pharmacokinetic and pharmacodynamic properties of compounds are viewed in the context of circadian rhythmicity, to optimally time drug treatment at an individual patient level. This approach merits formal evaluation using appropriately selected pharmaceutical drugs, and patients showing circadian rhythmicity in pain.

We conclude that pain, stiffness, and dexterity show circadian rhythmicity in hand OA. Assuming that these findings are generalisable to the OA hand population as a whole, there are important implications for scheduling activities of daily living, for measurement in clinical trials, and possibly for the time at which antirheumatic drugs are given.

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