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Nocturnal excretion of a urinary melatonin metabolite among electric utility workers

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Objectives The effects of 60-Hz magnetic field and ambient light exposures on the pineal hormone melatonin were studied among electric utility workers.

Methods Personal exposure was measured at 15-second intervals over 3 consecutive 24-hour periods. Exposure metrics based on magnetic field intensity, intermittence, or temporal stability were calculated for periods of work, home, and sleep. A rate-of-change metric (RCM) was used to estimate intermittence, and the standardized RCM (RCMS = RCM/standard deviation) was used to evaluate temporal stability. The effects of magnetic field exposure on total overnight 6-hydroxymelatonin sulfate (6-OHMS) excretion and creatinine-adjusted nocturnal 6-OHMS (6-OHMS/cr) concentration were analyzed with adjustment for age, month, and light exposure.

Results Magnetic field intensity, intermittence, or cumulative exposure had little influence on nocturnal 6-OHMS excretion. Residential RCMS magnetic field exposures were associated with lower nocturnal 6-OHMS/cr concentrations. In multivariate statistical analyses, the interaction term for geometric mean and RCMS magnetic field exposures at home was associated with lower nocturnal 6-OHMS/cr and overnight 6-OHMS levels. Modest reductions in the mean 6-OHMS levels occurred after RCMS exposures during work. The greatest reductions occurred when RCMS exposures both at work and at home were combined; therefore the effects of temporally stable magnetic fields may be integrated over a large portion of the day.

Conclusions Results from this study provide evidence that temporally stable magnetic field exposures are associated with reduced nocturnal 6-OHMS excretion in humans.

Key terms electromagnetic fields, human, 6-hydroxymelatonin sulfate, 60 Hz, magnetic fields, pineal.
Subjects and methods

The study population comprised 142 male electric power utility workers aged 20 to 60 years. Generation workers [N=29] (utility electricians and operators), distribution workers [N=56] (linemen and substation operators), and a comparison group of utility maintenance and administrative staff [N=57] were studied concurrently over a 1-year period. The mean age was 41 (SD 0.6) years; approximately 90% were Caucasian and non-Hispanic. All the subjects worked a normal daytime shift during their participation in the study. A questionnaire was administered to collect information concerning personal (age, race, body mass index), occupational (job title, employment duration, use of cell phones and other equipment, physical activity, work with chemicals), life-style (tobacco and alcohol use, sleep habits, electrical appliance use, exercise), and medical factors (medication, disease history) that might influence magnetic field exposure or melatonin production. None of the subjects were taking exogenous melatonin.

Exposure assessment

Personal exposure to magnetic fields and ambient light was measured over a period of 3 consecutive workdays, and during the night preceding the first day of work. Twenty-four hour magnetic field and light exposures were recorded at 15-second intervals with EMDEX C meters (19). Light exposure was measured by a Grasby Optronics photometric sensor adapted to the external jack of the EMDEX. The meter was worn in a belt pack with the subject at work and off duty; it was placed beside the bed adjacent to the waist during the worker’s sleep. Calibration logs and recordings of magnetic fields, light, and motion were inspected, and data were excluded if the meter was out of calibration, malfunctioning, or not worn. The participants logged their times at work, at home, and in bed, and exposures were partitioned accordingly. Home exposures were comprised mainly of time spent at the residence in the evening with a small component due to time at home prior to work.

Exposure metrics

Magnetic field and light exposure metrics were calculated for each exposure period and day of study. The arithmetic time-weighted average (TWA) was used to summarize personal light exposure. Magnetic field exposure metrics were selected a priori and included the arithmetic TWA, the geometric TWA, cumulative exposure, and cumulative exposure above 0.2 μT (24, 25). Two other metrics were calculated according to proposed mechanisms of magnetic field action. Exposure to fields with many switching events may have important biological implications (26—28). Therefore, a “rate of change metric” (RCM) based on the root-mean-square variation in successive magnetic field measurements was used to measure the intermittence of exposure (29):

$$\text{RCM} (\mu T/15 s) = \sqrt{\frac{1}{n-1} \sum (M_F^2 - \langle M_F \rangle^2)}$$

where \(M_F^1\) and \(M_F^2\) are successive 15-second magnetic field measurements and \(n\) is the number of measurements within a given exposure period. The RCM provides an estimate of both the variability and the first-lag autocorrelation in a series of measurements. Higher RCM values might indicate greater variability or less autocorrelation between successive readings or both. Others have suggested that temporally stable magnetic fields induce biological effects (30—32). The standardized RCM (RCMS) was therefore derived as follows:

$$\text{RCMS} (\text{per 15 s}) = \text{RCM}/\text{SD}$$

where SD is the standard deviation of the magnetic field measurements in a given period. The RCMS estimates the first-lag autocorrelation. Low RCMS values correspond to relatively small differences between successive measurements and represent magnetic field exposures that are stable over time. Thus low RCMS values should be directly associated with low 6-OHMS levels.

The geometric mean and RCMS magnetic field exposures are summarized in table 1 by worker group and exposure period. In general, the measures of magnetic field intensity correlated well. The geometric mean magnetic field exposures at work were higher for the generation workers than for the comparison workers (P<0.01). For comparison with other studies, the arithmetic means for the workplace exposures were 0.23, 0.32, and 0.15 μT for the distribution, generation, and comparison workers, respectively.

Determination of 6-hydroxymelatonin sulfate

Morning urine samples were collected daily for 4 days to determine the 6-OHMS levels. The base-line sample was obtained prior to the beginning of the workweek. The participants then submitted a morning sample on each of 3 consecutive workdays. Night-time and first morning voids were pooled to provide a total overnight sample. Melatonin production was assessed by a radioimmunoassay of urinary 6-OHMS concentrations (33, 34) (CI-Dtech, Mississauga, Ontario, Canada), which follow a diurnal pattern that is highly correlated with circulating melatonin (35). Total overnight 6-OHMS excretion and the nocturnal 6-OHMS concentration adjusted for creatinine (6-OHMS/cre) were calculated for each day.
Data analyses

Statistical analyses were performed using log-transformed data (log of the reciprocal for RCMS). Magnetic field exposures were compared among the distribution, generation, and comparison groups with a repeated-measures analysis of variance. Analyses for magnetic field effects were adjusted for age, month of participation, and TWA light exposure for the same period using Proc Mixed for repeated measures (SAS Institute Inc, Cary, NC, USA). Additional analyses were performed to evaluate potential confounding by the questionnaire variables; the results were essentially unchanged from those presented in this text. The potential effects of magnetic fields on the 6-OHMS excretion were modeled in 2 ways. First, 6-OHMS excretion was analyzed using each magnetic field metric as a continuous variable with age, month, and light exposure included as covariates in the Proc Mixed analysis, along with “day” and “magnetic fields by day”. Second, magnetic field exposures were divided into quartiles, and the least-squares means (adjusted for age, month, and light exposure) were estimated for the 6-OHMS for each quartile of exposure. The means in the lowest and highest exposure quartiles were compared by Fisher’s least significant difference method.

Results

The overall mean of the overnight 6-OHMS excretion was 22.7 (SD 1.3) µg, a value consistent with previously published data (35, 36). There was a statistically significant association between month of participation and both measures of 6-OHMS excretion (P<0.01); mean levels were higher in winter and lower in summer. Light exposures (TWA or cumulative lux) at home and during commutes from work to home were associated with lower 6-OHMS levels.

Table 1. Summary statistics for the magnetic field exposures of the male electric utility workers by exposure period. (RCMS = standardized rate of change metric).

| Worker group       | Geometric mean (µT) |                      |                  |                      |                      |
|--------------------|---------------------|----------------------|------------------|----------------------|----------------------|
|                    | Work*               | Home*                | Sleep*           | Work*                | Home*                | Sleep*               |
|                    | Mean    | SE     | Mean   | SE     | Mean   | SE     | Mean  | SE     | Mean  | SE     | Mean  | SE     |
| Distribution (N=56) | 0.10    | 0.03   | 0.11   | 0.04   | 0.08   | 0.05   | 0.54  | 0.04   | 0.55*  | 0.03   | 0.50  | 0.04   |
| Generation (N=29)  | 0.22**  | 0.07   | 0.14   | 0.05   | 0.11*  | 0.07   | 0.89  | 0.05   | 0.59   | 0.05   | 0.45*  | 0.04   |
| Comparison (N=57)  | 0.10    | 0.03   | 0.09   | 0.04   | 0.08   | 0.06   | 0.73  | 0.03   | 0.68   | 0.04   | 0.58  | 0.04   |

* Mean and standard error of each exposure metric for days 1, 2, and 3 combined.
* P < 0.05 versus comparison group, ** P < 0.01 versus comparison group.
Table 2. Nocturnal 6-hydroxymelatonin sulfate (6-OHMS) excretion by the quartile of the geometric mean magnetic field exposure. (cr = creatinine)

| Quartile of magnetic field exposure | Nocturnal 6-OHMS / cr concentration (ng/ml) | Total overnight 6-OHMS excretion (µg) |
|-------------------------------------|-------------------------------------------|---------------------------------------|
|                                     | 1  | 2  | 3  | 4  | 1  | 2  | 3  | 4  |
| Work                               | 29.6 | 25.2 | 30.0 | 28.5 | 17.7 | 15.0 | 17.8 | 14.8 |
| Home                               | 27.4 | 27.3 | 31.3 | 25.9 | 16.0 | 16.9 | 16.6 | 15.6 |
| Sleep                              | 25.3 | 31.9 | 30.6 | 26.8 | 15.1 | 17.7 | 17.7 | 16.3 |

* Least-square means based on adjustment for age, month of participation, and mean light exposure during the same period.

Table 3. Nocturnal 6-hydroxymelatonin sulfate (6-OHMS) excretion by the quartile of temporally stable magnetic field exposure. (cr = creatinine)

| Quartile of magnetic field exposure | Nocturnal 6-OHMS / cr concentration (ng/ml) | Total overnight 6-OHMS excretion (µg) |
|-------------------------------------|-------------------------------------------|---------------------------------------|
|                                     | 1  | 2  | 3  | 4  | 1  | 2  | 3  | 4  |
| Work                               | 29.2 | 27.5 | 30.9 | 27.2 | 17.6 | 17.3 | 16.6 | 15.1 |
| Home                               | 32.6 | 28.4 | 30.1 | 24.5** | 18.4 | 15.6 | 18.6 | 14.9 |
| Sleep                              | 29.1 | 23.3 | 29.2 | 28.3 | 17.3 | 16.8 | 16.2 | 15.7 |

* Least-square means based on adjustment for age, month of participation, and mean light exposure during the same period.

Additional analyses were performed to evaluate whether temporally stable magnetic field exposures over a larger portion of the day influenced 6-OHMS excretion. The subjects who were in the lowest quartile of RCMS exposure both at home and at work had mean nocturnal 6-OHMS/cr concentrations that were 39% lower than those in the highest quartile at home and at work (23.3 ng/ml versus 38.2 ng/ml, P=0.02) (figure 1). Similar results were obtained when these analyses were performed using the mean overnight 6-OHMS excretion as the dependent variable (12.9 µg versus 20.5 µg, P=0.03). A similar trend was noted for the subjects with temporally stable magnetic field exposure both at home and during sleep (results not shown).

**P<0.01 for 1st versus 4th quartile.

Discussion

Our findings indicate that the temporal stability of magnetic fields may be important for eliciting biological effects in humans. This hypothesis was based on the findings of Litovitz et al, who measured ornithine decarboxylase (ODC) activity in vitro after exposure to magnetic fields in which the frequency was shifted at various time intervals (30). The ODC activity doubled when the frequency of a 10-µT magnetic field remained stable for intervals of at least 10 seconds (30); this finding suggests that 60-Hz magnetic fields must remain stable over time in order to elicit effects (31, 32).

Based in part on these findings, the RCMS was developed as an estimate of the temporal stability of exposure. Consistent with this hypothesis, low RCMS values were associated with reduced 6-OHMS excretion. Time constants calculated from the lower quartile of the RCMS at work or at home indicated that exposures remaining highly correlated for intervals of at least 3 to 5 minutes on the average (assuming a first-order autoregressive model) were associated with reduced 6-OHMS levels. When analyzed separately, magnetic field intensity, intermittence, or cumulative exposure had little or no influence on...
6-OHMS excretion although the intensities were relatively low. However, the interaction between residential magnetic field intensity and temporal stability was associated with a reduction in both 6-OHMS variables and therefore suggests that the effects of temporally stable magnetic fields are enhanced at higher field strengths.

Our results indicate that the timing of exposure to temporally stable fields may be important for suppressing 6-OHMS excretion. Light exposures that occur at times when people are expected to be at home (ie, near dawn and dusk) influence nocturnal melatonin production (37-39), and the magnetic field suppression of melatonin may be mediated by retinal photoreceptors (40-42). If so, magnetic field exposures may need to coincide with specific periods of photosensitivity for melatonin suppression to occur. In controlled human experiments, magnetic field exposures that occurred prior to the onset of nocturnal melatonin production resulted in a delay in onset and a suppression in peak nocturnal plasma melatonin concentrations (43). Nocturnal melatonin onset usually occurs between 1600 and 2000 hours (2), which corresponds to the time of day when most of the subjects were at home. Other investigators have failed to elicit a reproducible suppression of nocturnal melatonin production in humans using only overnight magnetic field exposures that started at 2300 hours, after the nocturnal melatonin onset (44-46). Similarly, we found no statistically significant reductions in 6-OHMS in association with exposures that occurred only during sleep.

Reductions in mean nocturnal 6-OHMS levels were modest after RCMS magnetic field exposures at work. The greatest reductions in the mean 6-OHMS levels were observed when RCMS exposures at work and at home were combined (figure 1). This finding suggests that the effects of temporally stable magnetic fields are integrated over a longer time span than the approximate 8-hour periods that were used in this study and that exposures occurring during the day influence melatonin production at night. Animal experiments indicate that several weeks of exposure to 50- to 60-Hz electric or magnetic fields over a large portion of the day appear to be the most effective means of suppressing melatonin (47-52) although there are some inconsistencies (53-56). Short-term exposures have been ineffective (57, 58) unless repeated daily for several weeks (59).

Melatonin suppression has been reported for experimental animals after exposure to rapidly switched magnetic fields (27, 28). Our results do not support a role for intermittent exposures in suppressing 6-OHMS excretion. However, intermittent changes in magnetic fields at intervals of less than 15 seconds could not be evaluated. The rate-of-change metric was designed to capture switching events but, like the EMDEX meter, it does not specifically quantify transient exposures. Thus the negative findings for RCM in this study and the positive findings of others (27, 28) suggest that future studies should more carefully characterize exposure to high-frequency transients.

One strength of our study was the ability to measure light exposure and adjust for its effects on melatonin production. However, the light sensor response was matched to that of the human eye and was not maximal at all wavelengths that produce the greatest melatonin inhibition (60). Thus the effects of the measured light on 6-OHMS excretion may have been somewhat attenuated due to the misclassification of exposures.

The reductions in mean nocturnal 6-OHMS excretion associated with RCMS magnetic field exposures in this study (approximately 20-40%) were consistent with those reported elsewhere (15-17), and the results were in general agreement whether nocturnal 6-OHMS concentration or overnight 6-OHMS excretion was used as the outcome variable. Residential, rather than occupational, magnetic field exposures were most strongly associated with a reduction in nocturnal 6-OHMS excretion, which does not support the hypothesis that workplace exposures reduce 6-OHMS levels. However, the mean workplace exposures were lower than those reported by others (18-21), and they were only marginally higher than the mean residential exposures. Thus it was not possible to determine the effects of higher workplace magnetic field exposures on 6-OHMS excretion in our population. The finding that temporally stable magnetic field exposures, as measured by RCMS, are associated with reduced 6-OHMS excretion is unique and requires confirmation. Further work is also needed to determine whether 6-OHMS excretion is chronically suppressed in electric utility workers and to determine whether the effects are due to a reduction in the biosynthesis of melatonin, a phase shift in nocturnal melatonin production, or an increase in melatonin metabolism. Melatonin suppression may serve as a valuable tool for understanding human biological responses to magnetic fields.

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