The impact of shift work induced chronic circadian disruption on IL-6 and TNF-α immune responses

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Abstract

AIM: Sleep disturbances induce proinflammatory immune responses, which might increase cardiovascular disease risk. So far the effects of acute sleep deprivation and chronic sleep illnesses on the immune system have been investigated. The particular impact of shift work induced chronic circadian disruption on specific immune responses has not been addressed so far.

Methods: Pittsburgh-Sleep-Quality-Index (PSQI) questionnaire and blood sampling was performed by 225 shift workers and 137 daytime workers. As possible markers the proinflammatory cytokines IL-6 and TNF-α and lymphocyte cell count were investigated. A medical examination was performed and biometrical data including age, gender, height, weight, waist and hip circumference and smoking habits were collected by a structured interview.

Results: Shift workers had a significantly higher mean PSQI score than day workers (6.73 vs. 4.66; p < 0.001). Day workers and shift workers had similar serum levels of IL-6 (2.30 vs. 2.67 resp.; p = 0.276), TNF-α (5.58 vs. 5.68, resp.; p = 0.841) or lymphocytes count (33.68 vs. 32.99, resp.; p = 0.404). Furthermore there were no differences in cytokine levels (IL-6 p = 0.761; TNF-α p = 0.759) or lymphocyte count (p = 0.593) comparing the sleep quality within the cohorts. When this calculation of sleep quality was stratified by shift and day workers irrespective of their sleep quality day workers and shift workers had similar serum levels of IL-6, TNF-α or lymphocytes count. Multiple linear regression analysis showed a significant correlation of lymphocytes count and smoking habits.

Conclusion: Shift work induces chronic sleep debt. Our data reveals that chronic sleep debt might not always lead to an activation of the immune system, as we did not observe differences in lymphocyte count or level of IL-6 or TNF-α serum concentration between shift workers and day workers. Therefore chronic sleep restriction might be eased by a long-term compensating immune regulation which (in healthy) protects against an overstimulation of proinflammatory immune mechanisms and moderates metabolic changes, as they are known from short-term sleep deprivation or sleep related breathing disorders.

Introduction

Shift-workers are forced to work and sleep against normal chronobiological rhythms: they sleep at times their organism is set to activity and they work when physical and psychical effectiveness is generally low. These contradictory demands induce various indispositions; most frequently sleep disturbances which can cause severe sleep debt [1-6]. Recent research focuses on investigations how these sleep disturbances influence the immune system and if such an activation of the immune system might be linked with cardiovascular disease risks. Activation of inflammatory immune responses is marked by an increase of proinflammatory cytokines, e.g. Interleukin-1beta (IL-1β), IL-6 or Tumor-Necrosis-Factor alpha (TNF-α). Several authors report an augmentation of these cytokines after acute sleep
deprivation and thereby an induction of proinflammatory immune responses [7-9]. Cytokine levels correlate with fatigue and daytime somnolence [10-12]. Therefore Vgontzas et al. call IL-6 and TNF-α fatigue inducing cytokines [12]. Similar to acute sleep disorders people with a chronic sleep debt due to their obstructive sleep apnoe syndrome show elevated cytokine activity [13-16]. Cytokines regulate cell proliferation and differentiation. Thereby proinflammatory cytokine activation induces the recruitment of lymphocytes and neutrophils. Liu et al. demonstrated an increase of lymphocyte and neutrophil cells after acute sleep deprivation [17].

As inflammatory processes are recognized to play a major role in the pathogenesis of atherosclerosis, serum biomarkers are investigated to estimate emerging cardiovascular risk. There is increasing evidence that proinflammatory conditions and sleep disturbances elevate the risk of cerebrovascular and cardiovascular diseases [13,18-25]. This model may explain the higher cardiovascular diseases risk in shift workers [2,26-28].

So far this knowledge of associations between sleep debt and immune responses has not been related to the socioeconomically important factor shift work. We hypothesize that shift work induced chronic circadian disruption does affect proinflammatory immune responses. As possible markers of these changes in the immune system we investigated the proinflammatory cytokines IL-6 and TNF-α and measured the lymphocyte cell count.

Materials and methods

The study was approved by the Research Ethics Committee of the University Schleswig-Holstein Campus Lübeck, Germany (Ref. Az 05-028). Following informed consent, peripheral venous blood was drawn and the Pittsburgh-Sleep-Quality-Index (PSQI) questionnaire was completed by 225 shift workers and 137 daytime workers. A medical examination was performed and biometrical data including age, gender, height, weight, waist and hip circumference and smoking habits were collected by a structured interview. Both cohorts had a similar social background and worked in the industrial sector. Most shift workers worked in a three shift system including night shifts (91%), 9% worked in other schedules like permanent night shift, 24-hours or similar irregular shift schedules.

The mean age of shift workers was 36.4 yrs (SD 9.34) and the mean age of daytime workers 40.1 yrs (SD 7.77; p < 0.001). 86.9% of the shift workers and 75.7% of the day workers were male (p = 0.010). Shift workers and day workers did not differ in their Body Mass Index (BMI; kg/m²), (mean 26.62 versus 26.27 respectively, p = 0.455). Shift workers were significantly more smokers than daytime workers (41.8% vs. 27.0%; p = 0.005).

The PSQI is a validated instrument composed of 19 self-rated questions for the measurement of sleep quality during the previous month. Higher PSQI-Sum-Scores indicate inferior sleep quality. The questionnaire divides into “good sleeper” (PSQI ≤ 6), “poor sleeper” (PSQI 6 - 10) and “people with chronic sleep disorders” (PSQI ≥ 11), but the test does not enable to distinguish between different causes of sleep disorders. For this study a total value of 6 was defined the limit for the diagnosis “disturbed sleep”, meaning values ≥ 6 are a proved sign for relevant sleep disturbances, values < 6 signify none or slight, clinically not relevant sleep disturbances.

Fasting blood samples were withdrawn between 6 to 8 am, promptly centrifuged and aliquots stored at -40°C until analysis. Concentration of IL-6 and TNF-α were measured by ELISA-techniques as recommended by the manufacturer (Immulite, Los Angeles, Germany). Blood cell count was detected immediately after sampling with an automated hematology analyzer.

Statistical test were performed with descriptive methods, Levene's test for equality of variances, and t-tests for mean comparisons if applicable. Independent dichotomous variables were analyzed by Chi-Square-test, metric data by t-test, correlation coefficients were calculated on a ranked basis by spearman-rho procedure. Significance level was defined as p < 0.05. Data were normally distributed. Missing data rate was below 6%.

Results

Shift workers had a significantly higher mean PSQI score than day workers (6.73 vs. 4.66; p < 0.001) and significantly more shift workers had pathologically elevated scores (61.2% vs. 26.7%; p < 0.001).

Day workers and shift workers had similar serum levels of IL-6 (p = 0.276), TNF-α (p = 0.841) or lymphocytes count (p = 0.404) (table 1). Furthermore there were no differences in cytokine levels (IL-6 p = 0.761; TNF-α p = 0.474; lymphocytes p = 0.252).

| Table 1 Influence of shift work and sleep quality by PSQI on IL-6, TNF-α and lymphocytes |
|---------------------------------|-----------------|-----------|---|
| Component | Collective | Mean | SD | p |
|-----------|------------|------|---|---|
| IL-6 | Daytime workers | 2.30 | 1.54 | 0.276 |
| | Shift workers | 2.67 | 3.79 | |
| TNF-α | Daytime workers | 5.58 | 2.91 | 0.841 |
| | Shift workers | 5.68 | 5.19 | |
| Lymphocytes | Daytime workers | 33.68 | 7.64 | 0.404 |
| | Shift workers | 32.99 | 7.49 | |

| Component | PSQI | Mean | SD | p |
|-----------|------|------|---|---|
| IL-6 | PSQI ≤ 6 | 2.47 | 2.50 | 0.761 |
| | PSQI ≥ 6 | 2.58 | 3.77 | |
| TNF-α | PSQI ≤ 6 | 5.53 | 3.49 | 0.759 |
| | PSQI ≥ 6 | 5.68 | 5.41 | |
| Lymphocytes | PSQI ≤ 6 | 33.59 | 7.52 | 0.593 |
| | PSQI ≥ 6 | 33.14 | 7.78 | |
TNF-α \( p = 0.759 \) or lymphocyte count \( p = 0.593 \) within the cohorts comparing their sleep quality (table 1). Neither when this calculations of sleep quality was stratified by shift and day workers, i.e. example, IL-6 concentration in day workers with good sleep quality was 2.31 \( \mu g/ml \) in comparison to 2.67 \( \mu g/ml \) IL-6 in shift workers with good sleep quality (table 2). Comparing IL-6 concentration in respect to sleep quality shift workers with good sleep quality had a similar IL-6 levels (2.67 \( \mu g/ml \)) as their shift working colleges with relevant sleep disturbances (2.72 \( \mu g/ml \)) (table 2).

Multiple linear regression analysis showed a significant correlation of lymphocytes count and smoking habits. In contrast lymphocyte count and BMI, age, sleep quality or shift working status did not correlate. And there was no correlation of IL-6 or TNF-α with any of the above mentioned parameters (table 3).

In a last step cytokine levels were correlated with blood cell counts. Linear regression analysis revealed no significant correlation of the cytokines and monocytes count (IL-6: \( \beta = 0.008; T = 0.153; p = 0.879 \) or TNF-α: \( \beta = -0.025; T = -0.475; p = 0.635 \)) and neutrophils count (IL-6: \( \beta = 0.034; T = -0.542; p = 0.588 \); TNF-α: \( \beta = -0.021; T = -0.343; p = 0.732 \) as well as lymphocytes count (IL-6: \( \beta = 0.002; T = 0.036; p = 0.971 \); TNF-α: \( \beta = 0.014; T = 0.256; p = 0.798 \)).

## Discussion

The PSQI was used to evaluate sleep quality of the participants. Shift workers reported significantly worsened sleep quality compared to day workers. This is in line with many other authors who have demonstrated that shift work induces sleep disturbances and chronic sleep debt [1-6].

To investigate the long-term influence of chronic sleep debt and circadian disruption on the immune system we compared cytokine production and lymphocytes count in a morning blood sample of shift workers with daytime workers. Our results did not indicate an influence of shift work on cytokine levels of IL-6 or TNF-α and lymphocyte cell count. Furthermore there was no association between sleep quality and these indicators of inflammation in this study.

Referring to cell counts Liu et al. reported a short-term effect of a one night sleep deprivation on white blood cells and neutrophils, which we cannot find in our study setting looking for long-term effects [17].

Similarly, literature results on cytokine production are contradictory. E.g. Patel et al. described a direct association between the length of sleep and the increase of IL-6 serum levels or the decline of TNF-α [7]. In contrast Prather et al. showed that self-reported higher sleep debt scores predict elevated cytokine-levels of IL-1β and IL-6 [8]. As well Okun et al. found an association between bad sleep quality of pregnant women and elevation of IL-6 levels but no changes in TNF-α [29]. In their subsequent study in young healthy women IL-6 and TNF-α were not related with PSQI score and sleep duration, consistent with our data [30].

In another approach Vgontzas et al. reported short-term effects of a one night-sleep deprivation, which changed the circadian pattern of IL-6 secretion [11]. And they observed similar effects in diseased people with chronic insomnia [12]. These results have to evoke the considerations that we might have missed possible differences as we have only investigated a single time point and no daytime secretion.

Our results support an association of IL-6 levels and smoking. Literature reveals smoking to be an important confounding factor for proinflammatory reaction and cardiovascular diseases [28,31]. Shift workers smoke more frequently independently from demographic

### Table 2 Influence of sleep quality by PSQI on IL-6, TNF-α and lymphocytes stratified by shift and daytime worker

| Sleep quality | Component | Collective | Mean  | SD    | p      |
|---------------|-----------|------------|-------|-------|--------|
| PSQI < 6      | IL-6      | Daytime    | 2.31  | 1.59  | 0.357  |
|               |           | Shift      | 2.67  | 3.31  |        |
|               | TNF-α     | Daytime    | 5.51  | 3.01  | 0.945  |
|               |           | Shift      | 5.55  | 4.02  |        |
|               | Lymphocytes | Daytime | 33.98 | 7.60  | 0.453  |
|               |           | Shift      | 33.12 | 7.44  |        |
| PSQI ≥ 6      | IL-6      | Daytime    | 2.06  | 0.31  | 0.362  |
|               |           | Shift      | 2.72  | 4.24  |        |
|               | TNF-α     | Daytime    | 5.54  | 2.06  | 0.853  |
|               |           | Shift      | 5.73  | 6.07  |        |
|               | Lymphocytes | Daytime | 32.85 | 8.26  | 0.834  |
|               |           | Shift      | 33.17 | 7.73  |        |

### Table 3 Standardized correlation coefficient (β) with IL-6, TNF-α and lymphocytes in multiple regression analysis

| Component | Factor   | β      | t      | p      |
|-----------|----------|--------|--------|--------|
| IL-6      | BMI      | 0.103  | 1.842  | 0.066  |
|           | Shift work | 0.049  | 0.820  | 0.413  |
|           | Age      | 0.006  | 0.103  | 0.918  |
|           | Smoking  | 0.100  | 1.794  | 0.074  |
|           | Sleep quality | 0.006 | 0.099  | 0.921  |
| TNF-α     | BMI      | 0.025  | 0.447  | 0.656  |
|           | Shift work | 0.010  | 0.168  | 0.866  |
|           | Age      | -0.016 | -0.275 | 0.783  |
|           | Smoking  | -0.022 | -0.400 | 0.690  |
|           | Sleep quality | 0.014 | 0.239  | 0.811  |
| Lymphocytes | BMI    | -0.043 | -0.771 | 0.441  |
|            | Shift work | -0.005 | -0.088 | 0.930  |
|            | Age      | -0.068 | -1.208 | 0.228  |
|            | Smoking  | -0.170 | -3.092 | 0.002  |
|            | Sleep quality | -0.035 | -0.606 | 0.545  |
factors as age, gender or education [26,32-34]. Even after adjusting shift workers and day workers in regards to the variables of age, socioeconomic status, physical activity, smoking, and occupational load the shift workers keep their elevated risk for cardiovascular diseases [35].

An association between shift work, BMI and cardiovascular risks can be suspected [26,28,32]. Therefore one would assume that higher BMI scores predict higher proinflammatory cytokines. Our data do not show any association between sleep disorders and BMI.

Conclusion
In summary, shift workers experience significantly more sleep disorders than day workers. Shift work induces chronic sleep debt. Our data reveals that chronic sleep debt might not always lead to an activation of the immune system, as we did not observe differences in lymphocyte count or level of IL-6 or TNF-α serum concentration between shift workers and day workers. Therefore chronic sleep restriction might be eased by a long-term compensating immune regulation which (in healthy) protects against an overstimulation of proinflammatory immune mechanisms and moderates metabolic changes, as they are known from short-term sleep deprivation or sleep related breathing disorders. We can assume that long-term sleep debt in healthy may not lead to persistent proinflammatory changes in the immune system and to a consecutively higher risk for cardiovascular diseases. Further investigations are required to clarify the role of sleep in the immune system. Thus, most likely the increased cardiovascular disease risk of shift workers is caused from various parameters [28].

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Competing interests
The authors declare that they have no competing interests.

Received: 20 April 2010 Accepted: 5 July 2010 Published: 5 July 2010

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Authors’ contributions
AVM conceived of the study and its design, led the coordination, data collection, data analysis and interpretation of results and drafted the manuscript. SWW participated in the design of the study, data collection, data analysis and interpretation and in performing the statistical analysis. MS Schroeder participated in the design of the study, study coordination, data collection, data analysis and interpretation and in performing the statistical analysis and took part in preparation of the manuscript. ACO participated in the data collection, study coordination, data analysis and interpretation and in performing the statistical analysis and took part in preparation of the manuscript. KJC participated in the design of the study, data collection, data analysis and interpretation and took part in preparation of the manuscript. MS Paalke participated in the design of the study, study
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doi:10.1186/1745-6673-5-18

Cite this article as: van Mark et al.: The impact of shift work induced chronic circadian disruption on IL-6 and TNF-α immune responses. Journal of Occupational Medicine and Toxicology 2010 5:18.

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