

ORIGINAL ARTICLE

Can sleep quality and wellbeing be improved by changing the indoor lighting in the homes of healthy, elderly citizens?

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The study investigated the effect of bright blue-enriched versus blue-suppressed indoor light on sleep and wellbeing of healthy participants over 65 years. Twenty-nine participants in 20 private houses in a uniform settlement in Copenhagen were exposed to two light epochs of 3 weeks with blue-enriched (280 lux) and 3 weeks blue-suppressed (240 lux) indoor light or vice versa from 8 to 13 pm in a randomized cross-over design. The first light epoch was in October, the second in November and the two light epochs were separated by one week. Participants were examined at baseline and at the end of each light epoch. The experimental indoor light was well tolerated by the majority of the participants. Sleep duration was 7.44 (95% CI 7.14–7.74) hours during blue-enriched conditions and 7.31 (95% CI 7.01–7.62) hours during blue-suppressed conditions ($p=0.289$). Neither rest hours, chromatic pupillometry, nor saliva melatonin profile showed significant changes between blue-enriched and blue-suppressed epochs. Baseline Pittsburgh Sleep Quality Index (PSQI) was significantly worse in females; 7.62 (95% CI 5.13–10.0) versus 4.06 (95% CI 2.64–5.49) in males, $p=0.009$. For females, PSQI improved significantly during blue-enriched light exposure ($p=0.007$); no significant changes were found for males. The subjective grading of indoor light quality doubled from participants habitual indoor light to the bright experimental light, while it was stable between light epochs, although there were clear differences between blue-enriched and blue-suppressed electrical light conditions imposed. Even though the study was carried out in the late autumn at northern latitude, the only significant difference in Actiwatch-measured total blue light exposure was from 8 to 9 am, because contributions from blue-enriched, bright indoor light were superseded by contributions from daylight.

Keywords: Actiwatch, circadian rhythm, light for elderly, light intensity, melatonin, pupillometry

INTRODUCTION

Disturbances in circadian rhythms and disturbed sleep are important problems in modern society and the prevalence is increasing (Pallesen, 2014). Both hypersomnia (Jennum, 2014) and insomnia (Bin, 2012) are associated with increased socioeconomic costs related to reduced productivity, in- or outpatient treatment, medications, etc. Poor sleep quality is associated with increased risk of mortality from all causes (Ensrud, 2012), from cardiovascular events (Li, 2014), from stroke (Pan, 2014) and from suicide (Bernert, 2014). Daytime sleepiness is an independent risk factor for stroke and other vascular events (Boden-Albala, 2012), and insomnia is a risk factor for cardiovascular events (Canivet, 2014) and heart failure (Laugsand, 2014). Insomnia is more prevalent in older subjects (Gindin, 2014) and women (Pallesen, 2014; Peretti-Watel, 2009).

Most biological functions have some diurnal variation, e.g. cortisol levels, blood pressure and triglycerides (Skene, 2006). Variability in night-time blood pressure and lipid profiles may account for some of the association between circadian rhythm disturbances and increased risks of cardiovascular events and mortality (Palatini, 2014).

Circadian rhythms are maintained by external cues. The most important zeitgeber is ambient light conditions. In modern life many hours are spent indoors with a markedly lower light intensity than outdoors. During the evening, television and computer are frequently used and during night many bedrooms are polluted by light coming from outdoor sources. These conditions differ from the evolutionary adaptation of man where life was spent outdoors with very low light intensity at night. The lack of a timely sequence of high intensity

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light during the day and too much light at night may affect circadian rhythms and hence influence human mood and health, increasing the risk of hypertension and diabetes (Gordijn, 2012; Lee, 1997; Sharkey, 2013).

The photometer of the human body is located in a small number of cells in the retina (Berson, 2002). These cells, also called intrinsic photosensitive retinal ganglion cells (ipRGC), express a light sensitive photopigment: melanopsin. When illuminated by intense blue light, the ipRGC cell transmit impulses to the brain, in particular the suprachiasmatic nucleus (SCN), which in turn acts as the body's master clock, synchronising the timing of our physiology to the surroundings (Beaulé, 2003; Cajochen, 2005).

The classical photoreceptors, i.e. rods and cones, also contribute to the ipRGC's and studies have shown, that even low levels of light might stimulate the system due to the input from photoreceptors at light levels that are too low to activate melanopsin (Dacey, 2005; Glickman 2002).

Through the connection to the SCN, stimulation of ipRGC's suppresses the release of the hormone melatonin. The ipRGC's also projects to the brain centre for pupillary responses in the pretectal nuclei (Allen, 2011). In mammals, blue light has a significant influence on melatonin release, behaviour and pupil responses (Gamlin, 2007; Gooley 2012; Güler, 2008; Hattar, 2002; Kardon, 2009). In humans, different light regimes have been found to have beneficial effects on sleep parameters in many studies, ranging from younger people during controlled conditions in laboratory settings to elderly in nursing homes (Lockley, 2003; Lockley, 2006; Mishima, 1994; Schochati, 2000; van Someren, 2000; Vitiello, 2004). High intensity white light is used in the treatment of seasonal affective disorder (Gordijn, 2012; Martiny, 2004; Thompson, 1990).

The relation between light, melatonin and sleep has also been investigated for evening light. Evening exposure to blue-enriched light leads to phase-delays with suppression of the evening rise of melatonin and subjective lack of sleepiness (Chang, 2015). The effect on mood may not always be correlated to melatonin as daylight seems to influence alertness though melatonin is nearly absent (Sahin, 2014). Also, a recent study suggests that the melatonin concentration may adapt to long-term changes in light levels (Giménez, 2014).

Whereas the impact of light on the circadian rhythm in human is unquestionable, the optimal light conditions are difficult to define. Both intensity of light, proportions of blue light and timing to the solar cycle has to be considered and in addition, the internal rhythm of the individual subject. In particular, the focus has been on the proportion of blue light due to the major importance of the blue part of the spectrum of visible light for entrainment of the circadian rhythm (Brainard, 2001; Thapan, 2001). It seems that blue light is especially important during the morning and early hours of the day and detrimental late in the evening and

during the night (Chang, 2015; Nuutinen, 2014). With increasing age, blue light is transmitted to a lower degree through the lens of the eye (Brøndsted, 2013; Kessel, 2010; Turner, 2008) which may help explain the increase of sleeping disorders in the elderly population (Kessel, 2011; Kim, 2012; Stone, 2014).

The aim of the present study was to investigate the effect of indoor lighting on sleep and circadian photo-entrainment. In contrast to most studies, where the participants have been elderly in nursing homes and hospitals, the present study investigated the lighting conditions in the private homes of healthy elderly citizens. Participants were randomized in a cross-over design to blue-enriched indoor light versus blue-suppressed light in the morning hours. The primary outcome was sleep duration as reported by the participants, based on the hypothesis that improvement in circadian rhythms leads to a longer sleep.

METHODS

The study was a randomized cross-over design including healthy subjects above 65 years. Randomization to blue-enriched or blue-suppressed light first was performed with a computerized random number procedure, taking into account gender and whether participants shared home or not. The experimental indoor light was applied for 3 weeks in October 2013 followed by 1 week pause, and then the participants were subjected to the other light condition for 3 weeks in November 2013. In Denmark, daylight saving time ends on the last Sunday in October, and this was at the beginning of the one week pause. Due to this shift, sunrise was fairly equal for the two light epochs. The day length was less than 10.15 hours during the last of the 3 weeks in October and less than 8.25 hours during the last week in November.

Recruitment

The study was announced by direct mail to all residents of the chosen community and was followed by information meetings. Those who expressed an interest in participation were invited for initial evaluation and further information at the Department of Ophthalmology at Glostrup Hospital. A total of 38 subjects expressed interest, 2 subjects were too young and further 6 subjects did not continue to informed consent. One subject withdrew from the study before light installation and was substituted by another, leaving 29 participants in 20 households for the study. Subjects were excluded in case of depressions and/or medication considered to influence the circadian rhythm.

The study was approved by a local ethical review board and written informed consent was obtained from all participants. Also, the study was conducted in accordance with the policies of international chronobiological standards (Portaluppi et al., 2010).

Participants were encouraged to spend as much time at home until midday as possible, but strict restrictions on daily life were not imposed and for two male participants the daily routine included continuing working out of house, although at a reduced amount for the study period. All participants lived in uniform houses in a community in the town of Albertslund, in the greater Copenhagen area, Denmark. The sitting-room and dining-room faced South-West and in the opposite direction all walls were windowless. There were some differences in size of the individual houses and some houses were mirrored, i.e. the bedroom could be left or right of the dining-room.

Light conditions

All homes were visited prior to the study epochs to identify potential problems. Despite the uniform architecture of the homes, the visits revealed rather large differences in the indoor daylight levels due to furnishing, wall colors, and floor coverings as well as plants inside and outside the home. The indoor daylight level may have influenced the individual participant's perception and acceptance of the experimental light. However, we assessed the indoor daylight levels to be of little importance for the experiment. A lack of correlation between indoor daylight levels and measured light exposures on Actiwatchs supports this statement (data not shown).

Existing luminaires which were to be replaced by experimental light were hanging pendants over the dining tables and generally these lamps distributed only limited light at the position of the cornea (approximately below 100 lux). The experimental light was Philips Hue light bulbs © (<http://www2.meethue.com/en-us/this-is-hue/>), with 3 bulbs placed in globe-shaped frosted glass pendants in the central area of the house, i.e. where the dining table and/or living room was located and in close connection to the kitchen. This light position was chosen to maximise the hours spent by the participants near the light in each house during meals and other activities, especially in the morning hours.

To increase the light exposure and to avoid contrast problems, an LED uplight was mounted (Philips, SkyRibbon IntelliHUE Wall Washing Powercore © Eindhoven, the Netherlands) to illuminate the ceiling and add some indirect light to the exposure zone. Light from uplights and the HUE light bulbs was set to follow the same light sequence. From 8 am to 1 pm, the intensity of the light system was during the blue-enriched and blue-suppressed conditions controlled to reach approx. 280 and 240 lux, respectively, at the position of the cornea, measured half a meter from the frosted glass pendants. After 1 pm, the light intensity was decreased to approx. 140 lux blue-suppressed in both light epochs. The equivalent total irradiance was 920 mW/m² for blue-enriched, 710 mW/m² for blue-suppressed, and 392 mW/m² for the afternoon set-up (Table 1). From 6 pm until bedtime the setting was approx. 100 lux blue-suppressed.

TABLE 1. Total irradiance from on location measurements of the experimental light for the scenario applied from 8 am to 6 pm among four different houses.

Light condition	Light Intensity mW/m ²
Morning: Blue-enriched	
Mean (SD)	920 (170)
Total range	351
Morning: Blue-suppressed	
Mean (SD)	710 (110)
Total range	225
Afternoon:	
Mean (SD)	392 (48)
Total range	105

The morning scenario were applied from 8 am to 1 pm and were either blue-enriched or blue-suppressed. The afternoon scenario was applied from 1 pm to 6 pm and was blue-suppressed.

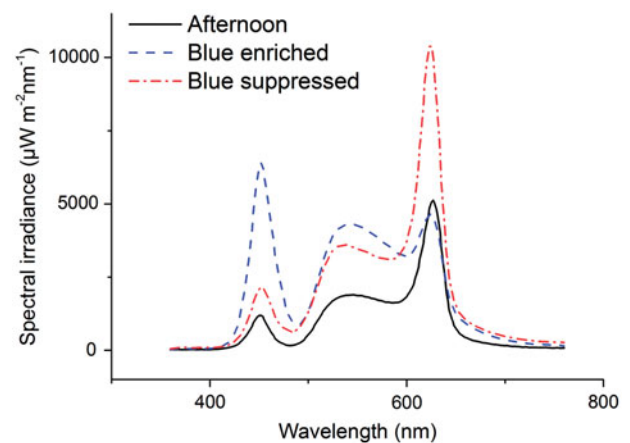


FIGURE 1. Spectral irradiance for blue-enriched, blue-suppressed and afternoon light. The blue-enriched and blue-suppressed light conditions were applied with total irradiance approximately 700–900 mW/m² during morning until 1 pm in the participant's living room. Based on a cross-over design, half of the participants were exposed to blue-enriched light and the other half to blue-deprived light during the first 3-weeks experimental period, and switched to the opposite in the second 3-weeks experimental period. From 1pm, the afternoon light conditions (total irradiance approximately 400 mW/m²) were applied in all cases.

During the night the participants could switch the light off or on when they wanted. If they wanted light turned on during night time, the light level was controlled at 100 lux blue-suppressed. In the bedroom of the homes, blackout curtains were installed. The light settings were measured on location (after sunset) with a handheld spectrometer from UPPrtek (MK350, Zhunan, Taiwan).

The blue-enriched light from the system had a correlated color temperature (CCT) at 5100 K with a wavelength peak (λ_p) at 450 nm (Figure 1). The blue-suppressed light CCT was measured to 2800 K and with a λ_p of 625 nm (Figure 1). The light spectra were chosen to achieve a color rendering index (CRI) above 80.

Ophthalmological examinations

Ophthalmological examinations were performed by the same experienced ophthalmologist in all participants.

The examination included measurement of intraocular pressure, biomicroscopic evaluation of both the anterior and posterior segment of the eye including an evaluation of the status of the lens and any presence of cataract, glaucoma or retinal changes. In addition to the clinical evaluation of the lens, the light transmittance of the lens was measured by an ocular fluorometer (Ocumetrics, Mountain View, CA). The lens was excited by blue light (488 nm) and the emitted auto-fluorescence was detected in the axial direction from the anterior to the posterior surface of the lens. The transmittance of the lens was calculated as the ratio between the signal intensity for the anterior lens divided by the signal from the posterior lens (Brøndsted, 2011). The visual acuity was measured with the ETDRS chart.

Questionnaires

A diary was filled in during the two light epochs. Participants reported the time when they went to bed, when they fell asleep and what time they woke up. The questionnaire also included the number and duration of naps during the day and the hours of television and/or computer usage after 6 pm. Self-reported diary is considered to be a valid method for assessment of sleeping pattern and time (Levenson, 2013).

The Pittsburg sleep index (PSQI) and the Morningness–Eveningness questionnaire were registered at baseline and after both light epochs (Buysse, 1988). Participants were instructed to refer only to the 3 foregoing weeks of experimental light when filling out the questionnaires. Poor sleepers were defined by a PSQI ≥ 5 . The Morningness–Eveningness (MEscore) scheme questionnaire was used to grade participants as morning types, score >59 , or evening types, score <41 , while neutral corresponds to a score in the range 42–58 (Horne, 1976).

At the end of each light period the participants were interviewed to grade their impression of the indoor light during the last 3 weeks. The interview was based on a visual analogue scale (VAS, scale from 1 to 10) and carried out by one of the authors. The aim of the interview was to obtain a subjective evaluation of the change in light and therefore, the participants at the first interview rated the experimental light (either blue-enriched or blue-suppressed as per randomisation) to the light in their home before the study. At the second interview, participants rated blue-enriched versus blue-suppressed light. The participants were encouraged to comment on the light.

Actiwatch

Activity and light exposure was measured with Actiwatch (Actiwatch Spectrum, Respironics, Philips Healthcare, Eindhoven, the Netherlands) worn on the wrist of the participants. Measurements were logged every 30 seconds around the clock in the last 7 days of each light epoch. With Actiwatch, the white light

response is derived from a sensor measuring light in three wavelength bands (red, green and blue). The white light response does not follow the spectral photopic luminous efficiency function, $V(\lambda)$ as described earlier (Figuerio et al., 2013; Price, 2012). Therefore, we corrected the “white light” output of all the Actiwatches against a calibrated illuminance photometer (Hagner® EC1-X, Instr. Nr. 54211, Solna, Sweden) using a side-by-side calibration method (Markvardt, 2015: <http://dx.doi.org/10.1080/15502724.2015.1020948>). Daylight at noon-time during overcast sky condition was used as the reference light-source. The red and blue Actiwatch light output was calibrated towards the average output of all the Actiwatches during the side-by-side calibration test. The Actiwatch calibrated white light output will in the following be termed photopic illuminance.

Chromatic pupillometry

Pupillary responses were measured with a chromatic pupillometer. In short, the pupillary contraction during and following light stimulation was video-filmed by an infrared camera and the pupil diameter expressed relative to the initial, dark adapted, undilated pupil (Herbst, 2011, 2012). Light exposure was 20 seconds of red light (660 nm) at 300 cd/m², followed by a 5 minutes pause, and a new exposure to blue light (470 nm) at 300 cd/m². During exposure to red light, the pupil redilates to some degree, and approximately 5 seconds after the light is turned off the baseline diameter is reached. At the light intensities used, the response to red light reflects cone activation, i.e. the photoreceptors which are also a part of our visual forming system. The pupil response to blue light is maintained both during light on and also after light off. Based on the properties of melanopsin activation the post-exposure response to blue light was chosen as the primary outcome for activation of the circadian system (Kankipati, 2010; Kardon, 2009). The outcome was expressed as the mean pupil contraction in % relative to the dark-adapted pupil-size (100%) and was measured from 0 seconds to 10 seconds and from 10 to 30 seconds after light off.

Melatonin

Saliva samples were collected at the end of each lighting epoch, from approximately 2 hours before estimated dim light onset, with 7 samples every 45 minutes (Aoki, 1998, Sharkey, 2013, Wahnschaffe, 2013). The samples for each participant were classified into 4 groups based on the maximal concentration at any time point: no sample above 2 pg/ml, no sample larger than 4 pg/ml, no sample larger than 8 pg/ml and cases with at least one sample larger than 8 pg/ml. The lower limit of detection from the laboratory was 0.5 pg/ml (Analyses by Stichting Ziekenhuis Gelderse Vallei, Ede, The Netherlands).

Statistics

The data are reported as the mean, SD (standard deviation) or standard errors (SE) and 95% confidence limits (95% CI). Assuming a normal distribution, paired *t*-tests were used to compare blue-enriched and blue-suppressed outcomes. A nonparametric signed rank test confirmed the parametric analysis. Chi-square was used for qualitative data and proportions. The major outcome of the study was the self-reported sleep duration, i.e. the number of hours from falling asleep to awakening in the test periods. Sleep duration, rest-time and other data from the diary and Actiwatch were determined from the mean of the last week of each light epoch. For further analysis, a general linear model was applied to evaluate if the outcomes were significantly affected by explanatory variables (gender, age and transmission of the lens). Backward elimination of non-significant covariates was used for the analysis. Data from married couples were treated statistically as independent observations. Visual acuity was measured on the ETDRS chart, and for comparison, also given as equivalent Snellen decimal.

Actiwatch data from 7 consecutive 24-hour days at the end of each light epoch was used for light-exposure analyses. The statistical analyses were performed with SAS (version 9.4, SAS Institute, Cary, NC) and a *p* value below 5% was considered significant. Separating epoch 1 versus 2 and blue-enriched versus blue-suppressed light conditions, light exposure averages per minute during the 24-hour day were calculated for each experimental day and afterwards this data-set was analysed using the mixed model procedure. The model included epoch and type of light condition as explanatory variables and random effects of experimental day. The same analyses were also performed only using data from the time between 8 am and 1 pm and between 8 am and 9 am.

RESULTS

A total of 29 participants in 20 homes were included in the study. The mean age was 69.7 years (95% CI 68.2 to 71.1) and 16 (55%) were male. The mean age was equal for the two genders, 69.5 years (95% CI 67.5–71.4) for males and 69.9 years (95% CI 67.5–72.4) for females. Nine married couples were included. In accordance with the randomization procedure, there was the same distribution of males and females receiving blue-enriched light in the first or second period; 8 males and 6 females received blue-enriched light in the first period and 8 male and 7 females received blue-enriched light in the second period ($p > 0.8$).

Sleep and rest

The sleep duration during blue enriched light was 7.44 hours (95% CI 7.14–7.74) for blue suppressed light the duration was 7.31 hours (95% CI 7.01–7.62), the difference was not significant ($p = 0.29$, Table 2). No advance

in the phase of sleep timing was found, as the difference between the two light conditions on the diary reported sleep time was less than 5 minutes ($p = 0.85$). The rest duration, i.e. the time from going to bed to awakening in the morning including reading, television etc. in bed, was 8.00 hours (95% CI 7.59–8.41) for blue enriched light and 7.88 hours (95% CI 7.47–8.29) during blue-suppressed light, the difference was not significant ($p = 0.257$, Table 2).

The rest duration measured by Actiwatch was 7.91 hours (95% CI 7.58–8.24) during blue-enriched conditions and 7.75 hours during blue-suppressed (95% CI 7.40–8.11, $p = 0.184$, Table 2). The rest duration measured by Actiwatch was shorter (non-significant) than the diary reported rest (Table 2).

The number of nightly awakenings ranged from 0 to 4 throughout the study period with a mean number of 1.00 during blue-enriched and 1.22 during blue-suppressed conditions ($p = 0.137$). The number of naps during day ranged from 0 to 1, with a mean of 0.33 during blue-enriched and 0.40 during blue-suppressed conditions ($p = 0.241$). Evening use of computer and television was also recorded in the diary. The daily time spent using computers and/or television during the study was 3.51 hours (range 1.56 – 6.63) for the blue-enriched and 3.54 hours (range 2.10 – 6.94) for the blue-suppressed conditions ($p = 0.843$).

Sleep quality

The Pittsburgh Sleep Quality Index (PSQI) at baseline with the participants' habitual lighting conditions was 5.66 (95% CI 4.22–7.09, Table 2). The mean score was significantly different for males and females: 7.62 (95% CI 5.13–10.0) for females versus 4.06 (95% CI 2.64–5.49) for males ($p = 0.0091$, gender-specific data presented in Table 3). Correspondingly, the number of poor sleepers at baseline ($PSQI \geq 5$) was 3 out of 16 men (19%) and 8 out of 13 females (=62%). The PSQI for the blue-enriched and blue-suppressed conditions were not significantly different, neither for all participants ($p > 0.9$) nor when analysed by gender ($p > 0.4$).

The PSQI improved significantly in females after installation of the brighter experimental light. On average, the PSQI score decreased by 2.0 (range 0 to –8, $p = 0.007$) from baseline to blue-enriched light and decreased by 1.76 (range 5 to –7, $p = 0.079$, Table 3) for blue-suppressed light. For males, no significant changes were found from baseline to the experimental light conditions ($p > 0.5$). The PSQI values were also analysed for the comparison of baseline visit to the first light epoch (October 2013), irrespective of light condition, and from baseline to the second light epoch (November 2013). The pattern was as that above, i.e. for females a decrease was found from baseline ($p = 0.018$ to the first period and $p = 0.053$ to the second light period), while no differences were found for males ($p > 0.5$).

TABLE 2. Results for 29 subjects at baseline and after each type of experimental light exposure.

	Baseline	Blue-enriched	Blue-suppressed	<i>t</i> -Value	DF	<i>p</i> value
Sleep and rest						
Diary, sleep hours, mean (SE)		7.44 (0.14)	7.31 (0.15)	1.08		
95% clm		7.14–7.74	7.01–7.62	27†		0.289
Diary, rest hours, mean (SE)		8.00 (0.20)	7.88 (0.20)	1.16		
95% clm		7.59–8.41	7.47–8.29	27†		0.257
Actiwatch, rest hours, mean (SE)		7.91 (0.16)	7.75 (0.18)	1.36		
95% clm		7.58–8.24	7.40–8.11	28		0.184
Sleep Quality						
PSQI, total score						
mean (SE)	5.66 (0.07)*	4.90 (0.70)	4.93 (0.66)	0.13		
95% clm	4.22–7.09	3.47–6.32	3.58–6.28	27		0.901
Poor sleepers						
PSQI ≥ 5, no.	17/29	13/29	11/28 ‡			
%	(59%)	(45%)	(39%)			0.672
Circadian rhythm						
Morningness-Eveningness						
mean (SE)	61.14 (1.36)	59.62 (1.52)	60.29 (1.38)	1.00		
95% clm	58.35–63.92	56.51–62.73	57.47–63.12	28		0.324
Melatonin						
max concentration (pg/ml)		8/7/7/6	8/6/9/5			0.937
≤2/>2≤4/>4≤8/>8						
Pupillometry						
% contraction, 0–10 seconds, mean (SE)	29.92 (1.23)	27.95 (0.95)	29.54 (1.46)	1.26		
95% clm	27.39–32.44	26.00–29.90	26.55–32.53	28		0.217
% contraction, 10–30 seconds, mean (SE)	13.23 (1.61)	11.39 (1.33)	13.54 (1.79)	1.43		
95% clm	9.94–16.52	8.67–14.2	9.87–17.21	28		0.165

Data given as mean, standard error (SE) and 95% confidence limits (95% clm). For paired *t*-tests, the *t*-value, degrees of freedom (DF) and the *p* value are included. *p* values are for the comparison of blue-enriched and blue-deprived light, where all were found non-significant. At baseline, the PSQI index was significantly increased in female gender (*: *p* = 0.009). Data separated by gender are shown in Table 3. Data were missing from one participant for the diary and one participant from the PSQI as indicated by † and ‡.

TABLE 3. PSQI for the subjects at baseline during blue-enriched and blue-suppressed light.

	Baseline	Blue-enriched	Blue-suppressed
Sleep quality by gender			
Male: PSQI, total score			
Mean (SE)	4.06 (0.67)	4.31 (1.04)	4.13 (0.97)
95% clm	2.64–5.49	2.09–6.54	2.06–6.20
Female: PSQI, total score			
Mean (SE)	7.62 (1.14)	5.62 (0.87)	5.85 (0.88)
95% clm	5.13–10.00	3.71–7.52	4.02–7.67
<i>p</i> value for comparison of genders	0.009	0.361	0.198
Male: comparison to baseline			
<i>p</i> value		0.721	0.784
<i>t</i> -Value		0.36	0.28
DF		15	14‡
Female: comparison to baseline			
<i>p</i> value		0.007	0.079
<i>t</i> -Value		3.22	1.92
DF		12	12

Data are presented as the mean, standard error (SE) and 95% confidence limits (95% clm). At baseline the PSQI was significantly higher for women than in men, thereafter the PSQI was similar for both genders. For females, a significant decrease was found from baseline to blue-enriched conditions, for the change to blue-deprived conditions we also found a decrease, but this did not reach statistical significance.

Circadian rhythm

At baseline, the mean MEscore was 61.14 (95% CI 58.35–63.92, Table 2), i.e. moderate morning. As for the PSQI, a significant difference between genders was found at baseline with a mean score of 58.00 (95% CI 52.82–63.18) in women and 63.69 in men (95% CI 60.97–66.41,

p = 0.035). The MEscore was neither significantly different for the comparison of blue-enriched to blue-suppressed conditions for all participants (*p* > 0.3) nor when analysed by gender (*p* > 0.2).

When baseline was compared to blue-enriched or blue-suppressed light, the change was non-significant

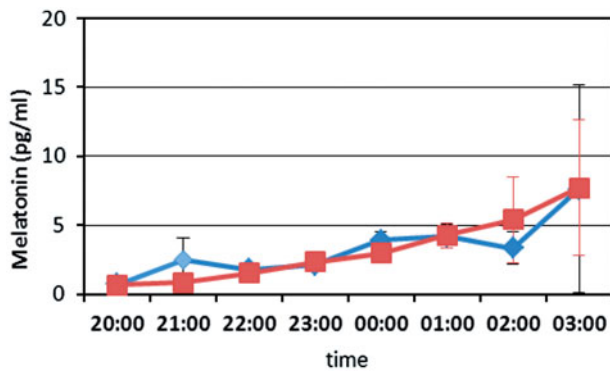


FIGURE 2. Melatonin concentration (mean and error bars) for all subjects during either blue-enriched light (blue) light or blue-suppressed light (red). Data are plotted in hourly intervals and the connecting lines show linear interpolation between data points. No significant differences of melatonin concentrations were found between the two experimental light conditions.

both for males and females ($p > 0.1$). For both genders, the MEscore decreased by 1.5 for the change to blue-enriched indoor light which was just on the border of statistical significance ($p = 0.0517$). For the change to blue-suppressed indoor light there was no significant change in the MEscore ($p = 0.2$). As for the baseline, the MEscore for the last light epoch was significantly lower for women compared to men (56.81 vs 62.41, $p = 0.040$).

Saliva melatonin concentration was measured at the end of each light epoch. No significant difference was found between blue-enriched and blue-suppressed light (Table 2, Figure 2). The variation between subjects was large, ranging from a concentration below 2 pg/ml during all hours with no detectable nightly rise, to a distinct increase during late evening and early night to a level approaching 50 pg/ml. No significant changes were found for the type of light or for the comparison of light epochs ($p > 0.5$).

Chromatic pupillometry did not show any differences in the pupil responses driven by photoreceptors (red light stimulation) or melanopsin containing ganglion cells (blue light stimulation) neither when comparing genders, spectrum of light (blue-enriched or blue-suppressed, $p > 0.3$) nor light epoch ($p > 0.6$).

Light conditions

Several participants reported that they found the experimental light to be brighter or much brighter during the morning hours. Subjective assessments of the pre-experimental light installations support this, revealing little light distributed on the cornea compared to after installation (data not shown). During the last week of each light epoch, the light exposure of the participants was monitored with Actiwatches.

Using data for full 24-hour days, light exposure analyses showed significant difference between the two epoch periods (October and November), with more light during the first epoch ($p < 0.001$). This relates to the analyses of the overall mean differences for red

light, blue light and photopic illuminance. Comparing blue-suppressed to blue-enriched conditions, a significant difference was found for the red light with more red light under the blue-suppressed conditions compared to the blue-enriched ($p = 0.024$). The estimated exposures of blue light ($p = 0.26$) and photopic illuminance ($p = 0.56$) were comparable for the two light conditions (Figure 3). Light exposure data from the Actiwatches, mounted on the wrist of participants living in their own homes, included the experimental indoor lighting as well as light from outside through windows and light received during time spent outside their homes. The daylight intensity was high compared to the imposed electrical light and may account for the lack of difference found for blue light and photopic illuminance exposure between blue-enriched and blue-suppressed light condition (Figure 3).

Limiting the data to the time-of-day from 8 am to 1 pm, the amount of red light was significantly increased during blue-suppressed exposure ($p = 0.0004$), whereas no significant difference was found for blue light ($p = 0.56$) or photopic illuminance ($p = 0.25$), neither when analysing blue-suppressed versus blue-enriched per se nor when the sequence of light-epochs was considered. Analysing only data from 8 am to 9 am the amount of red light was significantly increased during blue-suppressed exposure ($p < 0.0001$) and for this restricted time period, both the blue light and photopic illuminance was significantly decreased ($p < 0.0001$) during blue-suppressed exposure compared to the blue-enriched treatment. The results of the analysis, including estimates of the light exposure per light condition, can be found in Table 4.

Subjective evaluation

Participants were asked to rate the quality of the light on a visual analogue scale ranging from 0 to 10 with 5 being neutral. The participants preferred the experimental light (first epoch) compared to their habitual light, with a mean score of 7.84 (range 2.5–10). There was no preference of blue-enriched or blue-suppressed light, mean score 4.95 (range 1–9). As mentioned, several participants reported that they found the experimental light to be brighter or much brighter during the morning hours. Some participants expressed that the blue-enriched light was acceptable but too bluish and less cozy than their usual indoor light. One participant, in particular, disliked the blue light, while a few other participants favoured the blue-enriched light, feeling much more energetic, whereas the blue suppressed light made them feel less energetic.

Ocular status

The mean visual acuity (logMAR) was 0.03 (95% CI 0.09 to -0.02), corresponding to 1.0 Snellen (total range 0.25 to 1.6). The mean lens transmission was 0.474 (95% CI 0.44–0.53), range 0.21–0.65), i.e. less than half of the blue light was transmitted through the lens to the retina. Out

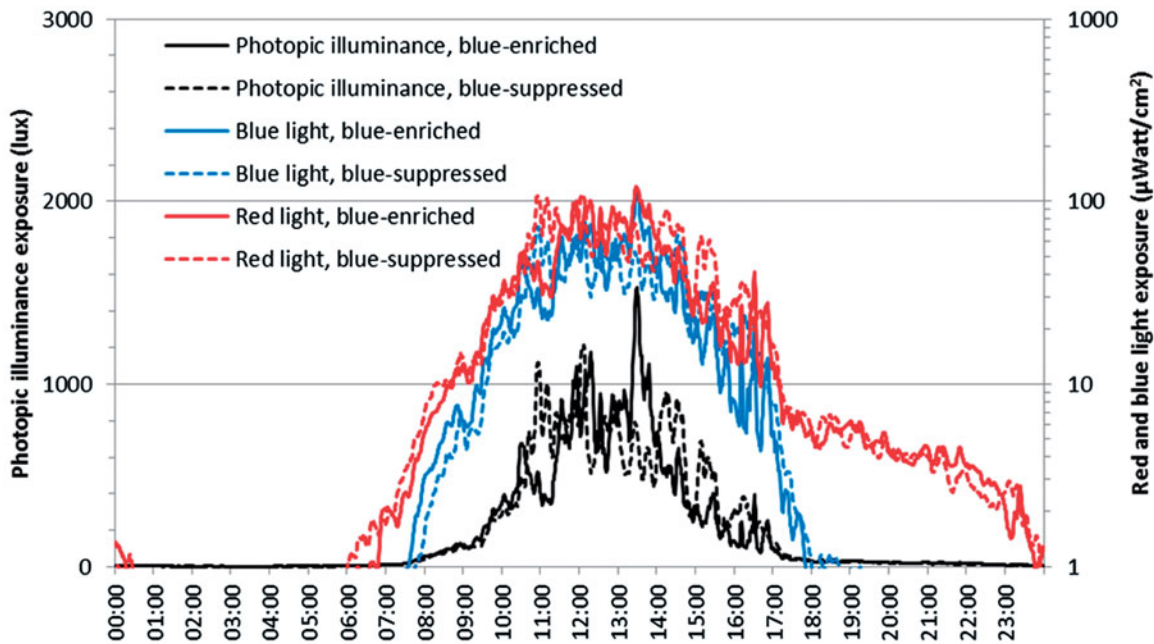


FIGURE 3. Average diurnal light exposure measured with Actiwatch. In the evening, the amount of blue light exposure decreased to almost zero unlike red light exposure (right axis). Significantly more red light exposure was found in the blue-suppressed treatment compared to the blue-enriched treatment ($p=0.024$), while no significant difference was found for the measured blue light ($p=0.26$, right axis) nor photopic illuminance ($p=0.56$, left axis) for the two experimental light conditions.

TABLE 4. Results of the statistical analysis for differences between blue-enriched and blue-suppressed light condition concerning Actiwatch red light, blue light and photopic illuminance for all participants.

Actiwatch measurement analysis	Period 8 am to 1 pm						
	Blue-enriched Estimate	Blue-suppressed Estimate	SE	F-value	DF	<i>p</i> Value	
Actiwatch red light	40.396	45.038	(10.686)	12.48	8385	0.0004	
Actiwatch blue light	28.621	28.050	(7.225)	0.33	8385	0.564	
Photopic illuminance*	444.28	461.25	(115.03)	1.34	8385	0.247	
Actiwatch measurement analysis	Period 8 am to 9 am						
	Blue-enriched Estimate	Blue-suppressed Estimate	SE	F-value	DF	<i>p</i> Value	
	Actiwatch red light	8.892	10.303	(1.139)	32.84	1665	<0.0001
	Actiwatch blue light	5.376	3.592	(0.782)	118.61	1665	<0.0001
Photopic illuminance*	89.005	77.039	(12.111)	24.73	1665	<0.0001	

Red and blue light levels in $\mu\text{Watt}/\text{cm}^2$ (nominal value) and photopic illuminance is the daylight calibrated Actiwatch “white light” output for approximate lux (*). Actiwatch measurements from 8 am to 1 pm time-of-day period and 8 am to 9 am time-of-day period were analysed. 24-hour time-of-day period gave comparable non-significant differences to the 8 am to 1 pm analysis. The Actiwatch recordings for red light showed a significantly lower intensity during blue-enriched light conditions, both during the day from 8 am to 1 pm and in the early morning (from 8 am to 9 am). Only in the early morning (from 8 am to 9 am), the recordings for blue light and photopic illumination showed a significant difference between the light conditions with higher intensity found for the blue-enriched light condition.

of the 58 eyes in 29 participants, 3 eyes had previously been operated for cataract and had an implant intraocular lens. During the study, 2 participants were diagnosed with cataract on both eyes and surgery was recommended, but performed after the study had finished. Glaucoma was diagnosed before the study in 2 eyes of 2 participants, for one of which the diagnosis and anti-glaucomatous treatment was changed as a consequence of the eye examinations. In addition, 1 eye was diagnosed with glaucoma. Retinal changes and a case of corneal scarring were found in additional 4 eyes in 4 participants, one of which was treated for

peripapillary choroidal neovascularization after the study since observation during the study period showed progression of subretinal fluid towards the fovea.

The data were also analysed for additional effects of age, gender, lens transmission. As stated above, gender was significant for PSQI at the baseline. The time of sleep and rest increased significantly with age, both measured with diary and Actiwatch (all p values <0.01), with increases of 0.14 hour per year for rest time and 0.10 hours for sleep time as reported by the diary and 0.11 hours per year as evaluated from the Actiwatch rest

times. The lens transmission was not significant for the outcomes.

DISCUSSION

In the present study, we investigated the effect of blue-enriched versus blue-suppressed indoor lighting in private homes of senior citizens in a randomized cross-over design. We did not find any significant differences between the two indoor lighting conditions in sleep or rest duration measured by diary and Actiwatch, or sleep quality measured with the Pittsburg Sleep Quality Index. Neither did we find significant effects of light conditions on the circadian rhythm parameters, i.e. morning-evening type, melatonin concentration or chromatic pupillometry.

We did, however, find that females had significant worse baseline PSQI score than males, and this improved significantly when the habitual indoor lighting was changed to the brighter blue-enriched experimental light. An improvement was also found for the change to blue-suppressed light, but the difference failed to reach statistical significance. We did not find any changes in PSQI scores for males. This probably reflects a ceiling-effect, as males had normal PSQI scores at baseline and hence no major improvements could be anticipated. The gender differences in sleep quality have been reported for a similar age group (Vitiello, 2004) and for patients with heart disease (Assari, 2013).

Blue-enriched light is considered the optimum for stimulation of the circadian rhythm and thus regulation of sleep (Brainard, 2001). Most studies of lighting conditions in the elderly has been conducted in nursing homes and hospitals, where the participants are subjected to daily routines set by the institution and many participants have severe health problems. Other studies have been conducted in the workplace showing significant effects on concentration and alertness (Mills, 2007; Viola, 2008). In these studies, a high color temperature of 17 000 K has been used and this may be more effective in stimulation the circadian system but also leads to low color discrimination and a bluish appearance of the surroundings which are not acceptable for elderly in their homes. Head to head studies of blue-enriched to blue-suppressed light at approximately the same intensity outside laboratory conditions are not frequent and a direct comparison of our study to previous studies is difficult. The intensity of the experimental light might have been too low to find a difference. When analysing Actiwatch data, we found a significantly increased amount of blue in the morning from 8 to 9, however the amount of blue light was not significantly higher later during the day. This may be due to the active lifestyle of the participants, as a social life with daily activities and also outdoor time are important time keepers. Also the amount of sunlight during the study is important, and unfortunately the weather in Copenhagen was dominated by unusual amount of

clear sky, in particular during the first light epoch. In order to ensure that the participants were exposed to the experimental light for as long as possible each day, our study was placed in the autumn when most people normally spent the majority of the day indoors but we did, however, not impose any restrictions on the daily activities of the participants.

The concentration of melatonin was highly variable, ranging from below 2 pg/ml for all samples in one-fourth of the participants to peak concentrations up to 50 pg/ml with a marked increase during early night. The differences in melatonin concentration between participants could not be explained by the changes in indoor light conditions or any of the other parameters in the study. It is noteworthy that a quarter of the subjects were below 2 pg/ml. Low melatonin concentrations have previously been described in older age groups and a lack of nocturnal rise has been reported both with age and in patients with dementia, a low concentration seems to be associated with a high level during daytime (Aoki, 1998; Cajochen, 2005; Karasek, 2004). Thus, the low melatonin concentration in a large group of our participants is probably related to age.

We performed chromatic pupillometry in all participants at baseline and after each experimental light epoch. Chromatic pupillometry was used as a measurement of activation of intrinsically photosensitive retinal ganglion cells (ipRGCs) by blue light. Stimulation of ipRGCs by blue light is the initial step in photoentrainment of circadian rhythms. We did not find any significant changes during the study suggesting that ipRGCs ability to be activated by blue light was not affected by changes in the indoor electrical lighting conditions.

Overall, the different parameters showed comparable results, both for the sleep duration, sleep quality and circadian rhythm parameters and did not support the concept of beneficial effects of blue light at least not in our healthy, active, elderly participants. A large difference in blue-enriched and blue-suppressed light was achieved for the isolated experimental indoor electrical lighting, with similar light intensities and acceptable color quality (Figure 1). However, the light measurements by the Actiwatch showed a high contribution from outdoor light hiding the contribution of blue light in the blue-enriched condition when analysing all 24-hour data and data between 8 am and 1 pm. This finding was not expected. We had specifically aimed for an autumn test period where the amount of sunlight normally is low in Denmark (latitude 55.7° N). However in 2013 the weather was unusually sunny (153.9 hours of sunshine in October and November 2013 versus an expected average of 145, source: the Danish Meteorological Institute: www.dmi.dk). These meteorological conditions are likely to have influenced our findings. If performed in December–January, the outdoor condition would have been darker, however the Danish tradition for Christmas celebration favours light

candles and reddish illumination and the application of the blue-enriched light would be hard to tolerate. In February, vacation abroad is very popular and due to these reasons, we chose late autumn. However, in the morning between 8 am and 9 am the contribution of daylight was limited and a significant difference between the blue-enriched and blue-suppressed light was found.

We included a detailed ophthalmic evaluation as part of our study design since ocular diseases such as glaucoma, age-related macular degeneration and cataract increases markedly with age and all may influence the visual and non-visual perception of light (Asplund, 2004, Ayaki, 2013; Feigl, 2011; Kankipati, 2011; Kessel, 2010, 2011; Nissen, 2014; Tanaka, 2010). As a consequence of the study, a number of participants were diagnosed with eye disease and were offered treatment for those diseases. This was felt as a positive side-effect of the study by the participants. No participants experienced any adverse events during the study.

The major part of the participants experienced a large subjective improvement in indoor light quality during the study. The amount of blue light was close to zero at evening, regardless of the daytime conditions. The increased light intensity during the mornings and/or low evening light might explain the change in sleep quality for women after initiation of the experimental light.

In conclusion, with a simplified scheme of blue-enriched or blue-suppressed lighting conditions from morning to midday, no difference was found for the sleep duration or other parameters of sleep quality. Daylight intensity was high during the test period and may have blunted responses to the experimental light and the comparison of blue-enriched to blue-suppressed light. The well-known poor sleep quality in females was found to improve by the bright experimental indoor morning light, suggesting that healthy elderly females with decreased sleep quality may benefit from brighter indoor light to improve sleep quality.

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