# Time to wake up: reactive countermeasures to sleep inertia

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Received November 21, 2015 and accepted May 11, 2016 Published online in J-STAGE May 18, 2016

Abstract: Sleep inertia is the period of impaired performance and grogginess experienced after waking. This period of impairment is of concern to workers who are on-call, or nap during work hours, and need to perform safety-critical tasks soon after waking. While several studies have investigated the best sleep timing and length to minimise sleep inertia effects, few have focused on countermeasures -especially those that can be implemented after waking (i.e. reactive countermeasures). This structured review summarises current literature on reactive countermeasures to sleep inertia such as caffeine, light, and temperature and discusses evidence for the effectiveness and operational viability of each approach. Current literature does not provide a convincing evidence-base for a reactive countermeasure. Caffeine is perhaps the best option, although it is most effective when administered prior to sleep and is therefore not strictly reactive. Investigations into light and temperature have found promising results for improving subjective alertness; further research is needed to determine whether these countermeasures can also attenuate performance impairment. Future research in this area would benefit from study design features highlighted in this review. In the meantime, it is recommended that proactive sleep inertia countermeasures are used, and that safety-critical tasks are avoided immediately after waking.

Key words: Body temperature, Caffeine, Countermeasures, Light, Napping, Self-awakening, Shift work, Sleep inertia

Impaired performance and alertness upon waking is known as "sleep inertia"<sup>1, 2)</sup>. Impairment is most severe immediately upon waking and then dissipates, generally returning to baseline levels within  $15-60 \text{ min}^{3-6)}$ . Sleep inertia is a concern for industries in which workers perform safety-critical tasks soon after waking. Motorists are also at risk when driving too soon after waking, for example when following government recommendations to nap if tired during long drives<sup>7, 8)</sup>. Sleep inertia has been a contributing factor in several major accidents and incidents<sup>9-12)</sup>. For example, an air crash involving 158 fatalities resulted from poor decisions made by the Captain who

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had just woken from an in-flight nap<sup>10</sup>. In another example, heavy contact between supply vessels occurred after the Chief Officer over-slept and consequently arrived on the bridge within minutes of waking<sup>11</sup>.

### **Factors that Influence Sleep Inertia**

Sleep inertia is typically measured in laboratory settings using a combination of sleepiness scales and cognitive tests. However, the sensitivity of these cognitive tasks to sleep inertia varies<sup>13, 14</sup>). Consequently, the severity of sleep inertia observed can depend on the task used to measure it. Test batteries are often administered immediately after waking and repeated intermittently for up to one hour. Sleep inertia has been measured under a variety of conditions ranging from a full night's sleep, to short naps at

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different times of day. Much of this research has focussed on establishing the best length and timing of sleep to minimise sleep inertia. Studies suggest that avoiding sleep periods ending during the circadian low<sup>5, 15)</sup>, and keeping naps to less than 30 min<sup>4, 16-18)</sup> to avoid waking from deep sleep<sup>15)</sup> can minimise sleep inertia magnitude. Prior sleep loss can also exacerbate sleep inertia<sup>15, 19</sup>, which is particularly important in shiftwork where workers often experience extended wakefulness and/or sleep loss<sup>20, 21</sup>). Indeed, naps have been suggested as a potentially effective countermeasure to sleepiness following sleep loss and during the circadian  $low^{22-26)}$ . Generally the long-term benefits provided by the nap outweigh the short-term detriments associated with sleep inertia<sup>4, 16, 25)</sup>. The challenge is to maximise the benefits of a nap while minimising and/ or managing sleep inertia. In addition, there may also be individual factors that contribute to overall sleep inertia severity. For example, some sleep medications<sup>27)</sup> or sleep disorders may exacerbate sleep inertia symptoms, although there is currently very little research in this area.

#### **Taking a Reactive Approach**

Research investigating proactive strategies for optimal sleep length and timing to minimise sleep inertia and maximise alertness is important for informing industry guidelines on rest breaks and shift scheduling. In operational environments, however, it is not always feasible to plan the length and timing of a sleep period. For example, when workers are on-call (such as in the emergency services or the military), or for workers such as healthcare professionals who take naps during extended-hours or night shifts. For these workers, the need to process crucial and complex information or to engage in safety critical activities almost immediately after waking could occur at any time. Given this, it is surprising that few studies have directly sought to reduce the effects of sleep inertia through reactive countermeasures. That is, strategies implemented upon wake-up, as opposed to proactive strategies such as planning sleep timing and duration. This review examines the literature on potential reactive countermeasures to sleep inertia including caffeine, light, and temperature, and discusses possible avenues for future research.

# Search Methods

Three electronic databases (PubMed, Science Direct, and Scopus) were searched on July 8, 2015 for the following search terms paired with "sleep inertia": adrena-

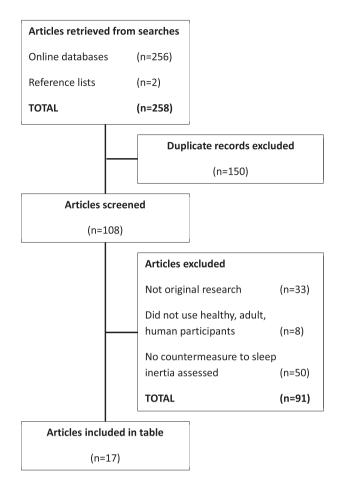


Fig. 1. Flowchart illustrating the structured narrative review selection process for articles populating Table 1.

line, caffeine, countermeasures, epinephrine, light, nap, noise, on-call, self-awakening, shift work, sound, and temperature. This resulted in 256 articles. Two articles were manually retrieved from reference lists. Seventeen articles remained after filtering for duplicates (n=150) and studies not meeting the inclusion criteria outlined below (n=91). Figure 1 illustrates the search and selection process. The large number of duplicate articles was a result of using multiple search terms and multiple databases which each returned similar results. The selected articles are summarised in Table 1. Note that one study is included three times as it investigated three countermeasures<sup>28</sup>.

Given the diversity of methods and the exploratory nature of the many approaches to managing sleep inertia, a traditional systematic review style was not implemented. Rather than limiting the scope of the studies included in the review, a structured narrative approach was chosen to allow collation across a broad range of research in this area<sup>29)</sup>. Many of the studies included in this review have methodological limitations that may have rendered them

Counter- measure	Authors	Prior sleep/ wake protocol	Sleep length	Wake-up timing	Inertia testing points post-sleep	Subjective alertness improved?^	Objective perfor- mance improved?^	Reac- tive?
Caffeine	Reyner & Horne, 1997 <sup>30)</sup>	Sleep restric- tion (5 h TIB)	15 min	14:40	Drive: 0–180 min; 30-min bins KSS: every 200 s	No obvious signs of inertia, but no nap-only comparison group.	No obvious signs of inertia, but no nap- only comparison group.	No
	Van Dongen <i>et al.</i> , 2001 <sup>31)</sup>	88 h extended wakefulness with 2-h naps every 12 h	2 h	04:45 & 16:45	5, 75 min	Not reported	Yes (5 min)	No
	Hayashi et al., 2003 <sup>28)</sup>	Habitual sleep	20 min	13:00	0, 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 55 min	Yes (15 min)	Yes (15 min)	No
	Newman et al., 2013 <sup>32)</sup>	Habitual sleep	1 h & ~6 h	01:00 & 06:00	0, 6, 12, 18 min	Not reported	Yes (18 min)	Yes
Light (post-wake)	Hayashi <i>et al.</i> , 2003 <sup>28)</sup>	Habitual sleep	20 min	13:00	0, 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 55 min	Yes (15 min)	No improvement	Yes
	Santhi <i>et al.</i> , 2013 <sup>13)</sup>	Sleep restric- tion (6.5 h TIB)	6.5 h	06:42–07:27, depending on habitual wake-up	3, 33, 63, 93 min	No improvement	No improvement	Yes
Light (pre-wake)	Van de Werken <i>et al.</i> , 2010 <sup>35)</sup>	Habitual sleep	~8 h	~07:00	1, 15, 30, 45, 60, 90 min	Yes (15 min)	No improvement	No
	Giménez <i>et al.</i> , 2010 <sup>33)</sup>	Habitual sleep	~8 h	~07:00	N/A	Yes (25 min reduction time needed to feel fully awake)	Not reported	No
	Harrison et al., 2011 <sup>34)</sup>	Habitual sleep	90 min	15:30	3 min	No	No	No
	Thompson <i>et al.</i> , $2014^{36}$	Habitual workday sleep	8 h	Habitual workday tim- ing (mean not reported)	5, 35, 75 min	Yes (better on average over whole testing period)	Yes (better on average over whole testing period)	No
Sound	Tassi et al., 1992 <sup>37)</sup>	Habitual sleep	1 h	01:00 & 04:00	0, 40 min. 30-min test divided into 3 min bins.	Not reported	Yes (3 min)	Yes
	Hayashi <i>et al.</i> , 2004 <sup>38)</sup>	Habitual sleep	20 min	14:20	1, 6, 11, 16 min	Yes (1 min)	No control group. High preference music better than low preference (1 min)	Yes
Tempera- ture	Krauchi <i>et al.</i> , 2004 <sup>39)</sup>	Unknown, assume habitual	8 h (noctur- nal sleep) & 2 h (after- noon nap)	07:00 & 18:00	KSS at 0, 30, 60, 90, 120 min	Correlation with distal-proximal skin temperature gradient	Not reported	Yes
	Krauchi et al., 2006 <sup>40)</sup>	Habitual sleep	75 min	Multiple wake- ups across the circadian cycle	KSS at 0, 30, 60, 90, 120 min	Correlation with distal skin temperature	Not reported	Yes
Self- awakening	Kaida <i>et al.</i> , 2003 <sup>41)</sup>	Habitual sleep	15-20 min	14:20	5, 10, 15, 20, 25, 30 min	Yes (5 min)	No Greater P300 ampli- tude (15 min)	No
	Kaida <i>et al.</i> , 2003 <sup>42)</sup>	Habitual sleep	15-20 min	14:20	5, 10, 15, 20, 25, 30 min	Yes (10 min)	No Heart rate increased (3 min before waking)	No
	Ikeda & Hayashi 2010 <sup>6)</sup>	Habitual sleep	Habitual (mean 7.3 h)	Habitual (mean 08:06)	1, 16, 31, 46 min	Yes (fatigue - better on average over whole testing period) No (sleepiness not improved)	Yes (in first 15 min test session)	No
	Ikeda <i>et al.</i> , 2014 <sup>43)</sup>	Habitual sleep	Partial restriction (5 h)	Habitual (range: 05:30-08:30)	Immediately after waking (home study)	No	Yes (only one test point)	No
Face- washing	Hayashi <i>et al.</i> , 2003 <sup>28)</sup>	Habitual sleep	20 min	13:00	0, 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 55 min	Yes (1 min)	No improvement	Yes

Table 1. A summary of papers investigating countermeasures to sleep inertia

Table notes: ^first recorded improvement; KSS: Karolinska sleepiness scale; N/A: not applicable; TIB: time in bed.

ineligible for traditional systematic review. Nevertheless, these studies highlight important research ideas, and point the way for future studies. In this way, this inclusive review is intended as a "call to arms" for future research initiatives.

Studies in Table 1 met the following inclusion criteria: original article; published in peer-reviewed journal; written in English; used healthy, human, adult participants; and directly trialled a countermeasure to sleep inertia or provided evidence for a potential countermeasure. The majority of screened articles were excluded due to not investigating a countermeasure to sleep inertia or not reporting original research (e.g. review papers) (Fig. 1). All studies in the table were laboratory-based; 15 were experimental and two were observational. Sample size ranged from n=8 to n=23 in 14 within-participant studies, and n=16to n=44 in three between-participants studies. Both male and female participants were included in all but three studies, which only included male participants. Outcome measures, testing points, timing of awakening, and prior sleep/wake history, and length of sleep periods varied, and as such, direct comparisons between studies were not possible. Six countermeasure categories were identified: caffeine<sup>28, 30-32</sup>; light<sup>13, 28, 33-36</sup>; sound<sup>37, 38</sup>; temperature<sup>39, 40</sup>; self-awakening<sup>6, 41-43</sup>; and face-washing<sup>28</sup>).

### Caffeine

Caffeine is a readily available and cost effective stimulant used strategically, socially and often habitually<sup>44, 45)</sup>. Caffeine promotes alertness by blocking adenosine receptors. Adenosine, a by-product of cellular energy expenditure, is a neurotransmitter inhibitor which increases with time awake and causes drowsiness<sup>46)</sup>. Caffeine has been shown to improve performance and alertness under conditions of high sleep pressure such as sleep deprivation and during the night<sup>47, 48)</sup>. The search yielded four studies that examined caffeine as a countermeasure to sleep inertia (Table 1). Van Dongen et al.<sup>31)</sup> found that sustained, low dose caffeine counteracted sleep inertia immediately following 2-h naps across 88 h of sleep deprivation. Notably, caffeine was administered hourly for a plasma concentration approximately 3.7 mg/L, equivalent to one-quarter cup of coffee every hour for 66 h. That is, caffeine was not administered directly after the nap, rather, at regular intervals before and after each nap. In this study, performance on a 10-min psychomotor vigilance task (PVT; a measure of behavioural alertness and vigilance, in which the participant is required to respond to stimuli presented

at random intervals as quickly as possible by pressing a button) under placebo conditions deteriorated immediately post-nap, but there was no change in performance pre- to post-nap in the caffeine condition. Testing points over an hour after the nap showed no differences between the two groups. From these results the authors concluded that caffeine had eliminated the effect of sleep inertia on this task. There was evidence to suggest that caffeine affected the sleep architecture of the naps in this study. For example, it took longer for participants to fall asleep, and the amount of time spent in deep sleep was significantly reduced in some of the nap opportunities. However, this effect was not consistent across all naps, yet the reduction of sleep inertia was consistent following all naps. Therefore, the authors concluded that the action of caffeine on sleep inertia was unlikely to be due to the alteration of prior sleep bout.

More recently, Newman et al.<sup>32)</sup> trialled a caffeinated chewing gum (100 mg) administered immediately upon waking and observed improved performance on a 5-min PVT relative to placebo at 12-18 min post-nap. Caffeine administered via gum reaches peak plasma levels more quickly than via a pill<sup>49)</sup>. However, there were no differences observed between conditions until at least 12-18 min post-nap, suggesting that the caffeine gum, as in the pill form<sup>46)</sup>, has a delayed effect, and is therefore unable to attenuate sleep inertia during the initial, most critical period of wakefulness. The caffeine gum may, however, limit the duration of sleep inertia and therefore provide faster recovery post-nap. It is worth noting that data from the placebo condition did not consistently show clear sleep inertia effects (i.e. performance was not always worse immediately after waking). Therefore, it is difficult to determine whether caffeine had an effect on sleep inertia per se, or improved general performance under conditions of mild sleep restriction and high circadian pressure (awoken at 01:00 and 06:00 after a 1-h and 6-h sleep opportunity, respectively). As an alternative mechanism, studies have shown that chewing non-caffeinated gum can improve alertness and cognitive performance<sup>50, 51)</sup>. Therefore, chewing may have acted as a countermeasure in Newman and colleagues' study<sup>32)</sup>, thus reducing sleep inertia effects in both the placebo and caffeine groups. If so, the caffeine effect would be acting on top of the chewing effect, leading to delayed improvements on background performance. Overall, the use of caffeinated gum offers promising results for truncating sleep inertia duration. This study demonstrates the importance of using a placebo gum, and ideally another placebo or control condition, so that it is possible to isolate the effects of chewing and caffeine on sleep inertia. Further study is warranted to determine whether different caffeine dosages may be more effective, or different performance tasks may be more sensitive to detect changes in sleep inertia following reactive caffeine gum administration.

Other studies have administered caffeine by adding it to decaffeinated coffee which was consumed prior to a short daytime nap<sup>28, 30)</sup>. Hayashi and colleagues<sup>28)</sup> administered 200 mg of caffeine (equivalent to 2-3 cups of instant coffee<sup>52, 53)</sup>) prior to a 20-min nap. Despite pre-nap administration, significant improvements in alertness and measured performance in the caffeine condition were not observed until 15 min after the nap. However, during this 15 min period, there were no differences in alertness or performance between the nap-only and no-nap group. This indicated a lack of observable sleep inertia, which may have been due to the experimental design. Mean total sleep time during the 20-min nap opportunity ending at 13:00 was 14.8 min, with no slow wave sleep (SWS, or "deep sleep"). These conditions are not particularly conducive to sleep inertia<sup>4, 16</sup>). This study demonstrates that, when investigating sleep inertia countermeasures, it is important to use experimental circumstances that are most likely to produce sleep inertia. For example, following: scheduled awakenings at night<sup>5, 15</sup>; extended wakefulness or prior sleep  $loss^{15, 19}$ ; or longer naps<sup>4, 16, 18</sup>).

Reyner and Horne<sup>30</sup>, using 150 mg of caffeine in decaffeinated coffee (equivalent to approximately two cups of coffee<sup>52, 53</sup>), found that the combination of caffeine and a 15-min nap was associated with better performance across a 2-h drive than following either countermeasure in isolation. The driving task commenced 5 min after the nap, within the sleep inertia period. However, as investigation of sleep inertia was not a research aim of the study, the data were reported in 30-min bins which do not allow investigation of performance changes across this peak sleep inertia period. The first 30-min bin, however, did not show any obvious signs of sleep inertia (i.e. worse performance) relative to subsequent bins. Interpretation of these results with regards to sleep inertia is further limited as performance in the caffeine-plus-nap group was not directly compared to a nap-only group, only to results from a group in a previous study<sup>54)</sup>. In the previous study<sup>54)</sup>, participants drove for 2 h before receiving caffeine and/or nap, whereas in the follow-up study<sup>30)</sup> they drove for 1 h. Therefore the post-treatment results of these conditions should be compared with this in mind.

There is a government-led road safety campaign, based on the above study, to promote the use of a caffeine-nap combination as a drowsy driving countermeasure<sup>55)</sup>. Further, despite a lack of studies investigating sleep inertia following short naps at night, other government websites promote the use of short naps ( $\leq$ 30 min) with no mention of potential sleep inertia<sup>7, 8)</sup>. Current policies and recommendations to counteract sleepiness while driving require a stronger evidence base with regards to sleep inertia. The studies reviewed highlight the need for experimental and observational testing which allows for the direct assessment of the impact of caffeine in combination with a nap on driving performance immediately after waking at different times of day.

There are several considerations when applying these results in the workplace. In most studies<sup>28, 30, 31)</sup>, caffeine was administered before the nap. Each of these studies vielded improvements post-nap in caffeine compared to placebo groups. In unpredictable scenarios, however, such as working on-call, there may not be sufficient warning or opportunity to use caffeine proactively. Proactive use may only be effective prior to short naps. There is the potential for sleep to be disrupted if caffeine is taken beforehand, resulting in a trade-off between reduced sleep inertia and reduced total sleep or sleep quality<sup>28, 31)</sup>. However, when caffeine was administered post-nap, even in a rapid acting chewing gum format, caffeine effects were not seen until 12-18 min post-nap, which, while useful in some scenarios, may not be fast enough in time-critical situations<sup>32</sup>). Therefore, when used proactively, caffeine has the potential to eliminate sleep inertia effects almost immediately after short naps. Further research into the use of caffeinated gum is needed to explore the potential benefits of this reactive countermeasure beyond the demonstrated reduction in sleep inertia duration.

Another consideration in the application of these results to operational settings is habitual use and individual caffeine sensitivity. Van Dongen and colleagues<sup>31)</sup> employed a two-week caffeine-free wash-out before the study to minimise the effects of individual differences in caffeine tolerance. This study still observed large between-subjects variability in plasma caffeine concentrations. The other studies in Table 1 did not report habitual caffeine use<sup>30</sup>, or allowed participants to drink up to four cups of coffee per day until the experimental period<sup>28, 32)</sup>. This may have influenced the effects of the caffeine, with low-caffeine users more sensitive to the effects of the experimental dosage compared to high-caffeine users<sup>46</sup>). Alternatively, highcaffeine users may have experienced withdrawals in the placebo condition<sup>56</sup>). Not controlling for habitual caffeine use, however, is perhaps more ecologically valid, as participants had a range of habitual caffeine intakes, just as a work force would. Studies of caffeine use in shift workers report high levels of caffeine intake especially for: older workers, those on night and morning shifts, and following reduced sleep<sup>45, 57)</sup>. Although it is important to understand caffeine effects under controlled conditions, it is also important to understand individual differences in caffeine response under natural conditions. Translating results from controlled studies into real-world scenarios is a critical step before recommending caffeine as a countermeasure to sleep inertia. Further, the long-lasting effects of caffeine may disturb subsequent sleep opportunities<sup>58)</sup>, so it may not be suitable for use on some occasions, such as towards the end of a night shift.

# Light

Melatonin is a hormone which helps to regulate sleepwake cycles and is suppressed under conditions of bright light exposure<sup>59</sup>. Light also serves as a *zeitgeber* (timegiver) to entrain our internal body (circadian) rhythms to a 24-h cycle<sup>60, 61)</sup>. Light signals are sent from the eye to a cluster of cells in the brain responsible for our circadian rhythms, the suprachiasmatic nucleus. In this way, light can be used at key times to manipulate circadian timing, leading to changes in sleep patterns<sup>62, 63)</sup>. In addition to its circadian entrainment effects, bright light exposure has also been shown to directly improve alertness and cognitive performance during the day, night, and following sleep restriction<sup>64–68)</sup>. Investigation into optimising the intensity, wavelength and duration of light exposure to manipulate both sleep and performance are ongoing. There is potential for bright light exposure to directly improve alertness and performance during the sleep inertia period.

Few studies have investigated the effect of light immediately following waking on performance and alertness during this period. The first study of this type exposed participants to bright light (2,000 lux) for 1 min after waking from a 20-min nap<sup>28)</sup>. This brief exposure to bright light did not reduce sleep inertia within the first 15 min after waking. It did, however, modestly improve subjective ratings for the following 45 min. More recently, Santhi and co-authors<sup>13)</sup> investigated the effect of four light conditions (dim, blueintermediate, blue-enhanced, and bright blue-enhanced) on a range of cognitive tasks during the sleep inertia period. Participants were woken from a 6.5-h nocturnal sleep and were exposed to light for 4 h. There were no differences between the four conditions in subjective alertness or performance measured every half-hour for the first two hours and then every hour up to 4 hours after waking. The relatively infrequent testing points during the initial sleep inertia period may have limited observations in this protocol. Participants in this study were selected based on a self-reported need for at least 60 min to feel fully alert after waking. Indeed, subjective ratings of sleepiness showed a clear sleep inertia pattern, with highest sleepiness ratings immediately after waking which dissipated across subsequent testing points. However, the performance tasks varied in their sensitivity to sleep inertia. This demonstrates the important methodological consideration of choosing an appropriate task to measure sleep inertia. Regardless of the outcome measured, however, there was no significant effect of light on performance or alertness within 4 h of waking. The improvement in working memory observed 4 h after waking was likely due to light affecting general performance rather than sleep inertia per se. These studies demonstrated that both brief and sustained light exposure after waking did not improve performance during the sleep inertia period. These results suggest that light exposure may be of limited effectiveness during the sleep inertia period. However, it may be worth investigating different light intensities and qualities.

As well as investigating light exposure after waking, some studies have looked at light exposure before waking. This approach assumes a different action pathway of light. Instead of direct alerting properties of bright light exposure during wake, exposure to light before waking applies the theory that light will act to lighten sleep before waking, therefore reducing sleep inertia<sup>35)</sup>. However, Harrison and colleagues<sup>34)</sup> found no difference in measured performance or alertness following a 90-min afternoon nap during which participants were exposed to varying light intensities (0, 1, 80, 6,400 lux). Similarly, Van de Werken et al.<sup>35</sup>, despite observing improvements in subjective alertness, found no differences in post-sleep performance on a simple reaction time and addition task between a dawn simulation condition and control. In the dawn simulation, light was increased to 300 lux in the 30 min prior to wake-up; there was no light before wake-up in the control condition. There were no sleep stage differences between conditions in either the 90-min nap<sup>34)</sup> or during the 30 min of artificial dawn<sup>35)</sup>.

In contrast, modest performance benefits were observed both on cognitive and physical (cycling) tasks following dawn simulation, with no differences in sleep quality as measured by actigraphy<sup>36</sup>). Giménez *et al.*<sup>33</sup> found that participants reported needing less time to feel fully alert after stronger artificial dawn light levels (250 lux vs 0 lux and 50 lux). Together these studies suggest that dawn light may improve subjective alertness. Further studies of the effect of dawn light on objective performance during the immediate wake-up period are necessary to determine the overall efficacy of dawn lighting to reduce subjective and objective sleep inertia.

Light may have an additional action pathway during the sleep inertia period via the cortisol awakening response (CAR). Cortisol is a hormone typically associated with stress response and follows a diurnal pattern with higher levels during the day and lower levels at night<sup>69)</sup>. The CAR refers to the sharp increase in cortisol upon waking in the morning<sup>70)</sup>. This response is greater in the presence of light presented immediately after waking (800 lux for 1 h)<sup>71)</sup>. Less intense light presented before waking (dawn simulation: light gradually increased to 250–300 lux over 30 min before waking) has had mixed effect<sup>35, 72)</sup>. Furthermore, a greater CAR has been associated with increased levels of self-reported arousal<sup>72, 73)</sup> and reduced sleepiness<sup>74</sup>).

The role of cortisol and the CAR in sleep inertia is unknown. A clue to its role comes from the discovery that SWS is associated with lower cortisol<sup>75)</sup>. Deep sleep has also been associated with greater sleep inertia<sup>15, 76)</sup>, although this relationship has not been consistently demonstrated<sup>18, 77)</sup>. The relationship between deep sleep and sleep inertia may be mediated by cortisol, although this has not been directly investigated. If the relationship between cortisol and sleep inertia can be better established, manipulating cortisol through light interventions or exogenous administration may offer an alternative sleep inertia countermeasure. The inclusion of measures of cortisol, alertness, and performance across the sleep inertia period would be beneficial to systematically track inter-relationships between these factors during this time.

Using light after waking would be relatively straightforward to implement in the workplace. For example, strategic exposure to light boxes at work has successfully helped shift workers adjust their circadian rhythm to better match their shifts<sup>78, 79)</sup>. However, a trial of bright light in a chemical plant uncovered several limitations including poor compliance with the interventions and variations in light exposure amongst workers due to different operational tasks<sup>80)</sup>. The effectiveness of such a countermeasure may vary by workplace. However, a light box, or light-emitting glasses<sup>81)</sup> used immediately after waking, could feasibly be implemented as a reactive countermeasure. While using light *before* waking would be unsuitable for implementation in unpredictable sleep/wake environments, it may have an application for scheduled rest periods. For example, a dawn simulation light box could be used in a workplace napping room. An additional caution to be added to the use of bright light as a countermeasure to sleep inertia is the potentially unwanted effect of entrainment. For example, bright light exposure, particularly near the circadian nadir, could delay or advance circadian phase<sup>62, 63)</sup>. This could lead to changes in sleep timing that are counterproductive to the work schedule and ultimately lead to sleep loss, sleepiness and poor performance.

## Sound

Sound such as pink noise (random noise with more low frequency components than white noise) can be used as a sleeping aid by providing a constant, ambient auditory background thereby minimising the impact of random noises on sleep initiation and maintenance<sup>82, 83)</sup>. Conversely, noise has been used to promote alertness under sleep deprivation conditions<sup>84)</sup> or as a positive stressor to improve performance<sup>85)</sup>. This inconsistency in the literature may be due to variation in noise type (e.g. continuous versus intermittent). The effect of noise may also be task dependent, with significant effects more likely observed on simple compared to complex tasks<sup>86–88)</sup>. The search identified two studies which used different sounds in an attempt to reduce sleep inertia.

Tassi and colleagues<sup>37)</sup> exposed participants to continuous noise during a spatial memory test performed after waking. In this study<sup>37)</sup>, spatial memory after waking from a 1-h nap at 01:00 during the no noise condition was worse for up to 15 min compared to the total sleep deprivation condition (no nap, no noise). Performance in the group exposed to 75 dB of pink noise (random noise with more low frequency components than white noise) after waking, did not differ from total sleep deprivation. The authors suggest that noise had an arousing effect which reduced sleep inertia. Noise delivered after a nap ending at 04:00, however, was ineffective and may have exacerbated the effects of partial sleep deprivation under these conditions. Given the conflicting results from this study and the literature generally, more research is required to establish the effect of pink noise exposure under different sleep conditions, and on different tasks. The background noise level of the work place may also impact on the effectiveness of this countermeasure. For example, high levels of ambient noise are typically associated with elevated stress and fatigue in noisy work environments such as factories<sup>89, 90)</sup>. The results of Tassi and colleagues' study<sup>37)</sup> suggest, however, that under certain conditions, exposure to pink noise after waking could minimise sleep inertia.

Hayashi and colleagues<sup>38)</sup> took a different approach, playing excitatory music at 60 dB during a 20-min testing period after waking. Music, especially music that the participant preferred, reduced subjective sleepiness post-nap compared to a no-music control. Similarly, performance on a visual oddball task was improved with high preference music compared to low preference music. Future research in this area would benefit from the inclusion of a no nap control group to confirm that the effects observed target sleep inertia relative to general performance. Reyner and Horne<sup>91)</sup>, under conditions of sleep restriction, found that playing the radio only had a brief, positive effect on alertness and simulated driving performance, and was therefore not suitable to long-distance driving. Playing music may, however, be effective in short-term sleep inertia scenarios.

From the limited literature it appears that sound (noise or music) has the potential to improve performance, at least briefly, under certain conditions. Further research in this area would help to determine the most effective sound to target sleep inertia symptoms. From an operational perspective, implementation in a work setting should be relatively easy. Indeed, background music has been used successfully in both industrial and office settings to improve vigilance and performance at work<sup>92)</sup>. However, rather than background music, sound delivery would likely need to be via headphones, especially in shared sleeping environments. For example, a personal music player with a collection of songs pre-selected by shift workers (high preference music<sup>38)</sup>) could be kept in a napping room for use immediately after waking.

# Temperature

Many studies have investigated the role of thermoregulation in moderating sleep onset and maintenance (for review, see Lack *et al.*<sup>93</sup>). At sleep onset there is a drop in core body temperature (CBT) and an associated increase vasodilation and blood flow to the extremities in order to lose heat. This change in body temperature is thought to promote sleep onset and maintain sleep throughout the nocturnal sleep period<sup>93</sup>). Furthermore, the association between the rate of distal skin temperature increase (measured at the extremities) and sleep onset can be observed at different phases of circadian rhythm in CBT<sup>94</sup>). Manipulation of skin temperature and CBT have been found to improve sleep quality<sup>95</sup>, performance, and maintenance of wakefulness<sup>96</sup>). For example, strategies to warm the extremities (e.g. a hot water bottle at the feet) can help to promote sleep onset<sup>97)</sup>. The role of thermoregulation at sleep offset, however, has received less attention<sup>98)</sup>.

A tantalising preliminary study in this area has demonstrated a relationship between subjective measures of sleep inertia and changes in the distal-proximal skin temperature gradient (DPG)<sup>39)</sup>. In this study, the reduction in subjective sleepiness correlated with a decrease in the DPG. Furthermore, the relationship between distal skin temperature and subjective sleepiness remained in a multiple nap protocol in which sleep and wake opportunities were rapidly alternated (75 min sleep / 150 min wake)<sup>40)</sup>. The authors suggest that actively manipulating the temperature of extremities through simple cooling strategies that evoke distal vasoconstriction and reduce heat loss from the periphery may result in faster sleep inertia dissipation. Van De Werken<sup>35)</sup> also found that dawn light accelerated the DPG change after waking and that this was associated with a decrease in subjective sleepiness, but not objective performance.

From an operational perspective, if changing body temperature only changed subjective feelings, it may lead to a false sense of improved objective alertness and performance after waking<sup>99–104)</sup>. Studies further investigating the relationship between DPG and objective measures are well-warranted, however, as an intervention would be relatively easy to implement in most work places. For example, workers could place their hands and feet in a receptacle filled with cold or iced water to rapidly lose heat from their extremities immediately after waking. Such a countermeasure would also have the advantage of not negatively influencing future sleep opportunities (c.f. caffeine, light).

# Self-awakening

Some studies have observed that waking from deeper stages of sleep (i.e. SWS) is associated with greater sleep inertia<sup>15, 76</sup>). This has led to studies of self-awakening to minimise the chances of waking from deep sleep. Self-awakening refers to spontaneously waking after a set period of time, without the use of an external stimulus such as an alarm. Kaida and colleagues<sup>41, 42</sup>) compared self-awakening to being woken by an experimenter after a 15-min afternoon nap. They reported that self-awakening led to lighter sleep, as scored using hypnagogic scoring<sup>105</sup>), a method which looks at the first two lighter stages of non-rapid eye movement sleep (Stage 1 and Stage 2)<sup>106</sup>) in greater resolution<sup>41</sup>). There were no differences observed across stages according to Rechtschaffen and Kales<sup>106</sup>) criteria<sup>41, 42</sup>). This finding suggests that traditional sleep scor-

ing methods may not be sensitive enough to detect electroencephalographic changes induced by self-awakening techniques. Spectral analysis of these sleep episodes may be another useful method for understanding the mechanisms underlying self-awakening.

Kaida and colleagues<sup>41)</sup> measured physiological arousal levels by evaluating the amplitude of a specific brain response to an audible tone presented immediately after waking and repeated every 5 min for 30 min. There was a greater response after self- compared to forced-awakening at 15 min post-wake, but not at earlier or later testing points. The authors argue that this greater response following self-awakening is indicative of greater physiological arousal<sup>41)</sup>. In a second study<sup>42)</sup>, subjective sleepiness was improved at 10 min post-wake, but improvements in performance on an auditory oddball task were not observed until later. These results should be interpreted with caution as the experimental design (i.e. a short afternoon nap) may not have generated sleep inertia. Indeed, there was no difference in performance between the nap and nonap group, and no immediate subjective ratings of sleep inertia. This study is another example highlighting the importance, when investigating sleep inertia countermeasures, of setting experimental conditions that are likely to produce sleep inertia. For example, Ikeda and colleagues investigated self-awakening following partial sleep restriction<sup>43, 107)</sup>. In contrast to Kaida and colleagues' studies<sup>41, 42)</sup>, improvement was observed on cognitive tasks following self-awakening, but there were no significant differences in subjective sleepiness ratings.

While there is some evidence to support the use of selfawakening to reduce sleep inertia, a potential limitation in applying this technique is the inconsistent success rate of self-awakening. In Kaida and colleagues' studies<sup>41, 42)</sup>, there was a 71-82% success rate for self-awakening within  $\pm 5$ min of the 15-min target, much higher than previous studies cited by the authors which range from  $18-42\%^{41}$ . The study, however, selected participants who were well-rested and claimed they were able to self-awaken from nocturnal sleep. Therefore this approach may be most beneficial for those who find it easier to self-awaken. It would be interesting to further this research under conditions better representing shift work scenarios, e.g. following chronic sleep restriction or extended periods of wakefulness<sup>21)</sup>. In Ikeda and Hayashi's study<sup>6</sup>, for example, participants habitually woke from an alarm, and subsequently only nine of the 30 attempted self-awakenings in the study (30%) were within 30 min of target wake time. As workers are generally required to return to shift at a specific time, it is

unlikely that self-awakening would be a reliable strategy. Furthermore, this method is not a viable countermeasure for unscheduled wake-ups. For naps taken at home without time pressures, however, the self-awakening technique may offer relief from subjective feelings of sleep inertia.

# Face Washing and Common Countermeasures

Anecdotally, people have wake-up routines which reduce sleep inertia. However, while commonly used countermeasures to driver fatigue such as cold air or playing the radio have been investigated<sup>91)</sup>, few studies have examined their use for sleep inertia. Hayashi and colleagues'28) trialled face-washing as a practical, and commonly used, sleep inertia countermeasure following a 20-min afternoon nap. The authors describe an immediate, but short-lived, reduction in subjective sleepiness relative to nap-only, with no differences in performance on a memory search task. These results suggest that face-washing may improve subjective alertness after waking. Further research using different performance measures would be useful to establish whether the null findings for performance were specific to the memory search task used. It is possible that other tasks may be more sensitive to inertia.

In a study examining the time course of sleep inertia, Jewett et al.<sup>3)</sup> found no differences in performance on an addition task between a constant routine condition (lying in semi-recumbent position) and an ambulatory condition in which participants were able to get up, eat breakfast, and shower. Both groups were kept under the same semi-recumbent, fasting conditions until at least 35 min after waking, therefore it is not clear whether these activities would have made a difference if undertaken sooner after waking. Another study reported a positive effect of breakfast on morning mood and memory, but did not report when participants woke, so it is unclear whether it was strictly a study of sleep inertia<sup>108)</sup>. More studies on common countermeasures such as showers, meals, or physical exercise are worthy of investigation. For example, a shower may act to change overall body temperature, and therefore might be more effective than face washing. Empirical research is needed to confirm effective common practices, and to de-bunk ineffective strategies.

# Adrenaline and Operational Scenarios

Finally, when considering sleep inertia in real-world scenarios, adrenaline is often proposed as a natural coun-

termeasure. Adrenaline is a stress hormone released from the adrenal glands<sup>109)</sup>. Adrenaline may play a role in mitigating impairment during the sleep inertia period<sup>110)</sup>. Laboratory studies of sleep inertia are typically conducted in low-stress environments. It is unknown whether a change in stress levels may affect sleep inertia, or the effectiveness of countermeasures. Further, there are no studies of adrenaline and its effect on performance and alertness during the sleep inertia period. This is likely due to constraints associated with collecting blood samples during the sleep inertia period and the short half-life of adrenaline<sup>111)</sup>. Anecdotal and experimental evidence suggest that adrenaline may not be a reliable countermeasure to sleep inertia. Emergency service workers often report habituating to their high pressure roles, no longer feeling the "rush of adrenaline". Reports are supported by a study that described changes to hormone levels following stressful field exercises and military training<sup>112)</sup>. Special Forces soldiers had a different response than non-Special Forces soldiers, suggesting a degree of adaptation to stress. However, this adaptation may not be common for shift workers in less stressful occupations. Whether the hormonal response immediately upon waking in high stress scenarios changes with experience, and/or varies between individuals, is unknown. Nor do we know whether increases in hormones such as adrenaline can restore cognitive function during the sleep inertia period, or whether they may in fact further compromise clear thinking and decision-making<sup>12</sup>). Until there are studies directly investigating the role of adrenaline on cognitive performance during sleep inertia, reliance on this endogenous countermeasure is not recommended.

This area would also benefit from study in the field, which to date, are few<sup>110</sup>. A survey of emergency medical service pilots revealed that sleep inertia had occasionally compromised flight safety<sup>113</sup> and some critical care nurses report avoiding napping on night shift due to concerns about performance impairment upon waking<sup>114</sup>). Taken together, these studies suggest that there is potential for sleep inertia-related safety risks, or at least, that the perception of sleep inertia as a safety issue exists in the field. Several incident investigation reports have cited sleep inertia as a contributing factor<sup>9–12</sup>). Systematic studies of the effect of sleep inertia in real-world situations are essential for progressing our understanding of sleep inertia, and the development of effective countermeasures for practical implementation.

### Conclusions

Examination of the current literature reveals a gap in the evidence-base for the implementation of a reactive countermeasure to sleep inertia which is effective within 15 min of waking. Caffeine is perhaps the closest option, although to target immediate performance and alertness, it needs to be administered prior to a short sleep bout, and in this way, is not a reactive strategy. This countermeasure may be useful, therefore, in situations where napping can be planned. Given that current investigations into light and temperature are in their infancy, further study is warranted to understand the physiological mechanisms underlying these interventions and whether performance can be improved after waking, in line with some encouraging results from subjective measures.

A challenge in identifying an effective sleep inertia countermeasure is that effectiveness may depend on sleep inertia severity. The majority of the studies reviewed assessed sleep inertia following afternoon naps, or upon waking from habitual night time sleep. There were no clear differences in the efficacy of countermeasures between these two scenarios. However, strategies that work under these conditions may not be sufficient in other scenarios. In shift work conditions, for example, the magnitude of sleep inertia may be exacerbated due to factors such as prior sleep loss, time of day, and sleep length. The impact of these factors on the effectiveness of countermeasures has yet to be investigated. Studies investigating sleep inertia countermeasures should consider directly comparing the effect of these factors, or testing countermeasures in scenarios more common to shift work (e.g. a night shift nap).

Identifying a viable countermeasure also involves assessing operational viability. Many of the countermeasures discussed could be implemented in a workplace. Developing a countermeasure that can attenuate the magnitude and/or duration of sleep inertia in both scheduled and unpredictable shift work environments warrants further investigation. In the meantime, it is recommended that proactive sleep inertia countermeasures are used, and that safety-critical tasks are avoided immediately after waking.

## Acknowledgements

The authors have no conflicts of interest to declare.

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