Predicting lapses of attention with sleep-like slow waves.

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Data and Code Availability:
   Raw data: https://osf.io/ey3ca/?view_only=680c39e7065649c3b783a4efec0a1a94
   Code used for analyses: https://github.com/andrillon/wanderIM

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Abstract

Attentional lapses are ubiquitous and can negatively impact performance. They correlate with mind wandering, or thoughts that are unrelated to ongoing tasks and environmental demands. In other cases, the stream of consciousness itself comes to a halt and the mind goes blank. What is happening in the brain that leads to these mental states? To understand the neural mechanisms underlying attentional lapses, we cross-analysed the behaviour, subjective experience and neural activity of healthy participants performing a task. Random interruptions prompted participants to indicate their mental states as task-focused, mind-wandering or mind-blanking. High-density electroencephalography revealed the occurrence of spatially and temporally localized slow waves, a pattern of neural activity characteristic of the transition toward sleep. These slow waves accompanied behavioural markers of lapses and preceded reports of mind wandering and mind blanking. Furthermore, the location of slow waves distinguished sluggish versus impulsive behaviours, mind wandering versus mind blanking. Our results suggest attentional lapses share a common physiological origin: the emergence of local sleep-like activity within the awake brain.

Introduction

The human brain sustains the stream of our conscious experiences. Attention can direct cognitive resources toward the external world and enable the selection and amplification of information relevant to an individual's current behavioural goals[1]. But attention can also turn inward, as is the case when we focus on internally generated task-unrelated thoughts, a phenomenon usually referred to as mind wandering[2]. Recent investigations have also shown that the stream of thoughts can come to a pause, as when individuals who are awake are left with the feeling of an empty mind (mind blanking)[3].

Shifts of attention towards the internal world, invoking mind blanking and mind wandering, can occur spontaneously without our knowledge or will[4]. In fact, a characteristic feature of attention is its fleeting nature and the difficulty to maintain it on a task for long periods of time[1,5]. In this paper, we define lapses of attention as the shift of the focus of thoughts away from the task at hand or environmental demands. The consequences of attentional lapses are very diverse. At the behavioural level, they can result in a lack of responsiveness or sluggish reactions, but they can also result in impulsive responses[6]. Curiously, these behavioural failures can be accompanied by a lack of conscious awareness and the absence of mental activity (mind blanking[3]), or rich, spontaneous mental activity (mind wandering[2]).

It is yet unclear whether these different types of attentional lapses (sluggish vs. impulsive behaviours; mind-blanking vs. mind-wandering) belong to a disparate family of behavioural and phenomenological events[7,8], each of them associated with different physiological causes,[9,10] or whether they can be traced back to common underlying

physiological causes[11]. Previous models of mind wandering have proposed that mind

wandering and mind blanking might arise in distinct neurophysiological states[3,9,10].

However, the fact that both sluggish and impulsive responses increase following sleep

deprivation[12,13] and in individuals with attentional deficits[6,14] implies a common

mechanism. Likewise, sleepiness has been associated with both mind wandering and mind

blanking[15,16] despite these two mental states being phenomenologically distinct[3].

Furthermore, investigations of the sleep onset period (hypnagogia) indicate that subjective

experiences resembling mind wandering (focus on internally generated contents) and mind

blanking (loss of awareness) can both occur at the border between wakefulness and

sleep[17,18]. Interestingly, these studies seem to associate lapses with pressure for sleep,

suggesting an involvement of fatigue.

Indeed, each hour spent awake comes at the cost of mounting sleep pressure. Past research

suggests that the need for sleep might only be dissipated by sleep itself,[19] as sleep plays a

vital role in neural homeostasis[20]. When individuals are prevented from sleeping for

extended periods of time (as in sleep deprivation studies), a subset of brain regions can

start displaying electroencephalographic (EEG) signatures of non-rapid eye-movement

(NREM) sleep in the form of sleep-like slow waves (within the delta (1-4 Hz) or theta (4-

7 Hz) range), despite individuals being behaviourally and physiologically awake[21,22]. These

sleep-like slow waves within wakefulness are referred to as "local sleep" in contrast with

the global whole-brain transition commonly observed at sleep onset[22–25]. Importantly, both

local and global transitions toward sleep are characterised by the same neural signature:

the occurrence of high-amplitude slow waves.

It has been proposed that the multiplication of sleep-like slow waves could perturb brain

functions and cause behavioural lapses[22]. In fact, during sleep, slow waves are associated

with episodes of widespread neural silencing[26], which have been connected to sensory

disconnection, behavioural unresponsiveness and the loss of consciousness[27,28].

Intracranial studies in humans and rodents showed that slow waves during waking are

associated with reduced neuronal firing, resulting in behavioural errors[21,22]. Wake slow

waves can also be detected in human non-invasive recordings[29–31] and here again the

amount of slow waves recorded in a given brain region correlates with the number of errors

performed in a task recruiting this specific brain region[29,30]. These results strongly suggest

that slow waves could explain the behavioural component of attentional lapses[22]. However,

the impact of slow waves on conscious experience, including mind wandering and

blanking, is still unclear.

In a recent review, we proposed that local sleep, manifested as the occurrence of slow

waves in waking, could not only explain the behavioural consequences of attentional

lapses, both regarding sluggish and impulsive responses, but also their phenomenology,

like mind wandering or blanking[11]. We also argued that local sleep is not an extreme

phenomenon, occurring only when individuals are pushed to their limit, but could occur in

well-rested individuals[31] and explain the occurrence of lapses in our everyday lives. To test

this framework, we rely on the detection of slow waves and formulate three different

hypotheses as follows: (i) Can slow waves predict, at the single trial level, both sluggish

and impulsive behaviours in well-rested individuals? (ii) Are slow waves associated with

both mind wandering and blanking? (iii) Does the location of slow waves (i.e. which

electrodes show slow waves) differentiate between sluggishness and impulsivity, mind-

blanking and mind-wandering? Through these hypotheses we will test the idea that local

slow waves could act as a functional switch, transiently perturbing the functioning of a

given cortical network. Accordingly, a common physiological event (local slow waves)

could lead to drastically different outcomes depending on its location within the brain.

To test these hypotheses, we cross-examined the behavioural performance, subjective

reports and physiological data from healthy individuals (N=26) performing an

undemanding Go/NoGo tasks[32] (Figure 1a). We sampled participants' subjective

experience by interrupting them during the task and asking them a series of questions about

their mental states prior to the interruption, including whether they were focusing on the

task, mind-wandering or mind-blanking (Figure 1b). Finally, we recorded their brain

activity using high-density scalp EEG and pupil size as objective proxies for participants'

level of vigilance[33]. Importantly, here participants were neither sleep deprived nor placed

in conditions favouring dozing off (i.e. not in bed or reclined, task requiring constant

attention and responses).

Figure 1. Experimental design and hypotheses.

a: Participants performed both a SART on faces stimuli (NoGo trials: smiling faces, stimuli

shown because of BioRxiv's guidelines) and a SART on digits (NoGo trials: "3").
*Face/Digit presentation was continuous (new face/digit every 0.75-1.25s). **b:** Every 30 to*
70s, participants were interrupted and instructed to report their mental state (see Online
and Supplementary Methods). Most importantly, they were asked to indicate whether they
were focusing on the task (task-focused: ON), thinking about nothing (mind-blanking: MB)
or thinking about something other than the task (mind-wandering: MW). High-density EEG
*and pupil size were continuously recorded throughout the task. **c:** Proportion of mental*
states reported during probes categorized as task-focused (ON, green), mind-wandering
(MW, orange) and mind-blanking (MB, blue) during the tasks with Digits (circles for each
individual participant; filled surfaces for smoothed density plot) and Faces (diamonds and
surfaces with horizontal stripes). Grey diamonds and circles show the average across
participants.

Results

Task performance and subjective experience. The Go/NoGo tests (see Online Methods)
require participants' sustained attention, but our participants declared focusing on the task
only in ~48% of the probes (Face Task: 49.4 ± 4.9%; Digit Task: 47.2 ± 5.1%; mean ±
Standard Error of the Mean (SEM) across N=26 participants; Figure 1c). The rest of the
time, they declared thinking about something else (Mind Wandering (MW); Face: 38.0 ±
4.3%; Digit: 40.9 ± 4.8%; Figure 1c) or thinking about nothing (Mind Blanking (MB);
Face: 12.7 ± 3.0%; Digit: 11.9 ± 2.9%; Figure 1c). These results are well in line with
previous findings[34,35] and highlight the prevalence of attentional lapses. Attentional lapses
were also reflected in participants' accuracy on the Go trials (Face: 3.1 ± 0.4% of miss, i.e.,
errors on Go trials; Digit: 1.7 ± 0.2%; Figure 2a), but more notably on the NoGo trials
(Face: 35.0 ± 2.5% of false alarms (FA), i.e., errors on NoGo trials; Digit: 32.5 ± 2.7%;
Figure 2b).

Figure 2. Low arousal is associated with attentional lapses characterized by different behavioural outcomes

*Proportion of misses (**a**) and false alarms (**b**) in the 20s preceding task-focused (ON, green), mind-wandering (MW, orange) and mind-blanking (MB, blue) during the tasks with*

Digits (circles for each individual participant; filled surfaces for smoothed density plot)

and Faces (diamonds and surfaces with horizontal stripes). The markers' size is

*proportional to the number of reports for each participant (same for **c-e**). Grey diamonds*

and circles show the average across participants, weighted by the number of reports (same

*for **c-e**). **c:** Distribution of reaction times (RT) for Go Trials (left: Face; right: Digit) in the*

20s preceding ON, MW and MB reports. Box plots show the mean (central bar), the lowest

and highest individual data points (end of the whiskers) and the lower and higher quartiles

(edges of the box). On top of each plot are shown the smoothed density plot for the different

*mental states. **d:** Vigilance scores (subjective ratings provided during probes) associated*

*with ON, MW and MB reports. **e:** Discretized pupil size (see Online Methods) in the 20s*

*preceding ON, MW and MB reports. In **a-e**, stars show the level of significance of the effect*

*of mental states (Likelihood Ratio Test, see Online Methods; ***: p<0.005).*

Next, we focused on the behavioural patterns preceding subjective reports of attentional

lapses. Specifically, we examined participants' behaviour in the 20s before the onset of the

probes that led to MW and MB reports (see Online Methods). To quantify the impact of

mental states (i.e., ON, MW or MB) on behaviour, we compared statistical models that

either did or did not include mental states as a predictor of behaviour (see Online Methods

and Supplementary Table 1). A significantly better fit by the model incorporating mental

states, assessed through a Likelihood Ratio Test, was interpreted as evidence for the

influence of mental states. To describe the size and direction of the statistical effects, we

report the estimates (β) of the contrasts of interest (MW vs ON, MB vs ON and MB vs

MW) and their 95% confidence-interval (CI). Accordingly, we found a significant effect

of mental states on misses (model comparison: $\chi^2(2)=36.0$, $p=1.5\times10^{-8}$; Figure 2a).

Specifically, MW and MB were associated with an increase in misses compared to ON

reports (MW vs ON: $\beta=0.011$, CI: [0.005, 0.016]; MB vs ON: $\beta=0.023$, CI: [0.015, 0.032])

and misses were more frequent for MB compared to MW reports (MB vs MW: $\beta=0.013$,

CI: [0.005, 0.021]). False alarms were also modulated across mental states (model

comparison: χ^2(2)=115.9, p<10^{-16}; see Figure 2b), with an increase for both MW and MB

compared to ON (MW vs ON: β=0.21, CI: [0.17, 0.24]; MB vs ON: β=0.17, CI: [0.12,

0.23]), but similar levels of false alarms for MB and MW (MB vs MW: β=-0.028, CI: [-

0.084, 0.028]). Finally, mental states were associated with different patterns of reaction

times (RT; model comparison: χ^2(2)=16.3, p=2.9x10^{-4}; Figure 2c) with slower reaction

times for MB compared to both ON and MW reports (MB vs ON: β=0.019, CI: [0.009,

0.030]; MB vs MW: β=0.022, CI: [0.011, 0.032]; MW vs ON: β=-0.0025, CI: [-0.0096,

0.0045]).

Taken together, these results suggest that MW and MB decrease performance in different

ways. MB is characterized with more misses and slower RT than MW and ON, which is

consistent with the idea that MB induces sluggish mental states. As to MW, the analysis of

misses may ostensibly compatible with an idea of MW as a mild form of MB, however,

the fact that RT in MW is faster than in MB is not easy to reconcile with such an idea.

Instead, this overall pattern is consistent with an idea that MW induces more impulsive

mental states.

The same pattern of results was obtained when normalising misses, false alarms and RT in

the MW and MB conditions by the average obtained in the ON condition for each subject

(see Supplementary Table 2). We also obtained qualitatively the same results when

focusing on a shorter time-window (10s before probe onsets instead of 20s; see

Supplementary Table 3).

Vigilance. Although MW and MB differ according to their phenomenological definition

and associated behaviours, both states seem to occur in a similar context of low vigilance

(as reported by participants themselves during probes). To address this, we quantified the

degrees of correlation between participants' vigilance ratings and each mental state

(comparison between models including or not the information about mental state:

$\chi^2(2)$=144.8, p<10^{-16}; Figure 2d and Supplementary Table 1). Participants reported lower

vigilance ratings for both MW and MB compared to ON (MW vs ON: β=-0.39, CI: [-0.40,

-0.37] and MB vs ON: β=-0.53, CI: [-0.55, -0.50]). Vigilance ratings were even lower for

MB compared to MW (MB vs MW: β=-0.13, CI: [-0.24, -0.02]). We then examined a

classical objective proxy for vigilance: pupil size[33,36]. Pupil size prior to probes (Figure 2e,

N=25 participants here, see Online Methods) was significantly modulated by mental states

(model comparison: $\chi^2(2)$=18.0, p=1.210^{-4}) with MW and MB associated with smaller

pupils than ON probes (MW vs ON: β=-0.29, CI: [-0.43, -0.15]; MB vs ON: β=-0.22, CI:

[-0.45, -0.003]). Pupil size did not differ between MW and MB (MB vs MW: β=0.065, CI:

[-0.16, 0.29]), despite the significant correlation between vigilance ratings and pupil size

(model comparison: $\chi^2(2)$=134.5, p<10^{-16}; see Table S1 for details on the models). This

implies that these two measures of vigilance can be differentially sensitive to mental states.

Slow Waves. To further examine the potential mechanistic link between sleep pressure and

attentional lapses, we set out to detect a marker of sleep pressure in the EEG signal, in the

form of local, sleep-like slow waves. We recently reviewed the rationale behind this

approach[11]. In particular, relying on a local marker of sleep allowed us to test whether

distinct families of attentional lapses can be coherently explained by the occurrence and

spatio-temporal characteristics of slow waves.

Following evidence from [21,22,29,30], we operationally defined slow waves as the occurrence

of large-amplitude waves within the delta ([1-4] Hz) range. We first detected the

occurrence of slow waves in each EEG electrode using an established approach developed

in wakefulness and sleep (see [30,31,37] and Online Methods). Both the temporal profile and

topographical distribution of slow waves detected during the tasks (Figure 3a-b) resemble

the slow waves observed in NREM sleep[37,38]. This is not trivial as our detection algorithm

did not select this specific temporal profile or topographical distributions.

Figure 3. Properties of Slow Waves

a: Average waveform of the slow waves detected over electrode Cz during the behavioural tasks (red, left; N=26 participants). The average waveform of slow waves detected during sleep (blue, right), extracted from another dataset (see Supplementary Methods), is shown for comparison. Slow waves were aligned by their start, defined as the first zero-crossing before the negative peak (see Online Methods). b: Scalp topographies of the density of slow waves (arbitrary units) detected in wakefulness (top) and sleep (bottom). c: Scalp topographies of wake slow-waves properties (1st column: temporal Density; 2nd: peak-to-peak Amplitude; 3rd: Downward Slope (D-Slope); 4th: Upward Slope (U-Slope); see Online Methods) averaged across participants (N=26). d: Scalp topographies for slow-waves Density (1st column), Amplitude (2nd), Downward Slope (D-Slope, 3rd) and Upward Slope (U-Slope, 4th) for the different mental state (ON, MW and MB).

Relationship between global properties of slow waves and vigilance. Next, we checked whether global properties of the slow waves index participants' level of vigilance. We correlated first participants' subjective vigilance ratings with the amount and properties of slow waves. The amount and properties of slow waves were extracted prior to probe onset ([-20, 0]s) and averaged across the entire scalp (all 63 electrodes, see Online Methods). Model comparisons between models with or without slow-wave density, amplitude, downward slope or upward slope indicate a negative correlation between vigilance ratings and slow-waves density ($\chi^2(1)=13.1$, p=3.9×10^{-4}, β=-0.074, CI: [-0.114, -0.034]), amplitude ($\chi^2(1)=33.1$, p=8.5×10^{-9}, β=-0.023, CI: [-0.031, -0.015]), downward slope ($\chi^2(1)=82.1$, p<10^{-16}, β=-2.5×10^{-3}, CI: [-3.1×10^{-3}, -2.0×10^{-3}]), and upward slope ($\chi^2(1)=47.3$, p<10^{-11}, β=-1.7×10^{-3}, CI: [-2.2×10^{-3}, -1.2×10^{-3}]) (see also Figure S1). We then verified that slow-wave properties were negatively correlated with pupil size recorded before each probe (model comparison between models with or without slow-wave density, amplitude, downward slope or upward slope: density: $\chi^2(1)=12.3$, p=4.6×10^{-4}, β=-0.13, CI:

[-0.21, -0.059]; amplitude: $\chi^2(1)=7.4$, $p=6.4\times10^{-3}$, $\beta=-0.0088$, CI: [-0.015, -0.0025];

downward slope: $\chi^2(1)=15.1$, $p=1.0\times10^{-4}$, $\beta=-6.9\times10^{-4}$, CI: [-1.0×10^{-3}, -3.4×10^{-4}]; upward

slope: $\chi^2(1)=17.6$, $p=2.7\times10^{-5}$, $\beta=-7.6\times10^{-4}$, CI: [-1.1×10^{-3}, -4.0×10^{-4}]) (see also Figure S2).

Finally, we confirmed that the amount of slow waves detected increased with time spent

on task (Figure S3), as can be expected when considering the homeostatic regulation of

sleep and local sleep[22].

Relationship between local properties of slow waves and mental states at the probe level.

Local properties of the slow waves (Figure 3c-d) predicted reports of mental states.

Specifically, we examined, for each scalp electrode, whether slow-wave properties

(density, amplitude, downward and upward slopes) prior to probes were predictive of the

mental state reported by participants. To do so we focused on pairwise contrasts (MW vs

ON, MB vs ON and MB vs MW) and performed this analysis for each electrode

independently. A cluster permutation approach ($p_{cluster}<0.05$, Bonferroni corrected cluster

threshold; see Online Methods) revealed that MW (compared to ON) was predicted by an

increase in the number of slow waves and slow wave amplitude over frontal electrodes,

and by an increase in slow waves' downward and upward slope over centro-frontal

electrodes (Figure 4a). MB (compared to ON; Figure 4b) was also predicted by an increase

in slow wave density over frontal areas as well as by the slow-wave slopes (downward and

upward) over centro-parietal electrodes (no significant cluster for slow wave amplitude).

Finally, a direct contrast between MB and MW (Figure 4c) indicated that a reduction of

slow wave amplitude over frontal electrodes but an increase in their upward slope over

parietal electrodes were predictive of MB. No significant clusters were obtained for slow

wave density and downward slope.

To provide more details on the temporal relationship between slow waves and mental

states, we replicated this analysis by splitting the 20s window before probes into four

windows of 5s (Figure S4-6). The contrasts between MW and ON (Figure S4) as well as

MB and ON (Figure S5) show that the properties of slow waves best predict mental states

within the 5s before a probe. In terms of topography, when compared to ON state, MW is

best predicted by the slow wave properties in frontal electrodes (Figure S4) while MB is

best predicted by those in the centro-parietal electrodes (Figure S5).

Figure 4. Local properties of slow waves are predictive of mental states

Locally, based on each individual electrode, we performed mixed-effect analyses,

following with permutation analysis to quantify the impact of slow wave properties on

mental states. 1st column: Density; 2nd: Amplitude; 3rd: Downward Slope (D-slope); 4th:

Upward Slope (U-slope). These slow-wave properties were extracted for each electrode

and used to compute the t-values (shown in each topographical plot) from t-tests on the

following comparisons: (a) MW > ON (b) MB > ON and (c) MB > MW. Black dots denote

significant clusters of electrodes ($p_{cluster}$<0.05 corrected for 12 comparisons using a
Bonferroni approach, see Online Methods).

Relationship between local properties of slow waves and behaviour at the trial level. To

further understand the association between slow waves and attentional lapses, we examined

the influence of slow waves on participants' behaviour at the single-trial level (for all trials

within 20s of a probe onset, independently of mental states). To do so, for each trial and

electrode, we marked the presence or absence of slow waves between stimulus onset and

offset (see Online Methods) and used this as a (binary) predictor of RT, misses and false

alarms (Figure 5). This analysis revealed spatially-specific effects of slow waves on distinct

behavioural outcomes. Namely, slow waves in frontal electrodes co-occurred with faster

reaction times while slow waves in posterior electrodes co-occurred with slower reaction

times (Figure 5a, $p_{cluster}$<0.05, Bonferroni corrected cluster threshold). Likewise, frontal

slow waves were associated with more false alarms (a marker of impulsivity, Figure 5b)

and posterior slow waves with more misses (a marker of sluggishness, Figure 5c).

Figure 5. Local occurrences of slow waves are associated with modulations of
performance

Mixed-Effects Models were used to quantify the correlation between slow-wave occurrence

*and reaction times (**a**), false alarms (**b**) and misses (**c**) at the single-trial level.*
Topographies show the scalp distribution of the associated t-values. Black dots denote
significant clusters of electrodes ($p_{cluster}$<0.05 corrected for 3 comparisons using a
Bonferroni approach, see Online Methods).

Decision modelling. Finally, we implemented an influential model of two-alternative

forced-choice (2AFC) decision making: the diffusion decision model (DDM)[39]. The DDM

decomposes full reaction-time distributions and choice proportions into latent cognitive

processes that are thought to underlie participants' decisions in 2AFC tasks (see Online

and Supplementary Methods and Figure S7). These include the time it takes for participants

to start computing their response (Non-Decision Time or NDT [t]), the speed at which

participants accumulate evidence for the two responses (drift rates for Go [v_{Go}] and NoGo

[v_{NoGo}] responses), the amount of evidence needed to reach a decision (decision threshold

[a]), the participants' initial bias for one of the two responses (decision bias [z]) and the

bias for the accumulation of evidence for one of the two responses (drift bias, [v_{Bias}]). A

hierarchical Bayesian approach was used to fit the DDM to the reaction times obtained in

the Go/NoGo tests[40] so that each parameter (v_{Go}, v_{NoGo}, a, z, t and v_{Bias}) was free to vary by

participant, task and mental states (ON, MW and MB) or slow-wave occurrence (present

vs. absent). Simulations confirmed this hierarchical DDM can successfully predict the

observed data (Figure S8).

Considering the trials that were within 20 seconds from the onset of the probes, we first

estimated these different parameters of the DDM for each mental state (Figure S9). This

analysis shows differences between MW and MB reports in terms of decision bias and

threshold (Figure S9). The lower threshold and higher decision bias observed for MW

(compared to MB) are concordant with the idea that MW is associated with behavioural

impulsivity.

We then used this modelling approach to examine how the occurrences of slow waves

impact the different cognitive processes leading to participants' responses, with a particular

focus being the test of our core hypothesis: the presence of slow waves in frontal areas

leads to impulsivity by disrupting the cognitive mechanisms underlying executive control,

while the presence of slow waves in posterior areas leads to sluggishness by slowing down

the integration of sensory inputs.

We report here the differences in the parameters' estimates in the presence or absence of

slow waves detected for each electrode (Figure 6). The associated scalp topographies

indicate both global and local (electrode-specific) effects of slow waves on DDM

parameters. As global effects (Figure 6d-f), we found first that the presence of slow waves

was associated with a reduction in decision threshold (a; Figure 6d), consistent with the

idea that slow waves facilitate impulsive responses. Second, slow waves were also

associated with longer non-decision times (t; Figure 6e), suggesting that slow waves can

slow down neural processes underlying stimulus encoding and/or motor preparation.

Finally, the presence of slow waves was correlated with an increase in the starting point of

the decision process (prior bias z; Figure 6f), implying shifts in the decision process

towards Go responses.

As local electrode-specific effects, we observed again contrasting results between posterior

and frontal slow waves (Figure 6a-c). Slow waves within posterior electrodes were

associated with a reduction of v_{Go} and v_{Bias}, meaning that evidence accumulation was

slower for Go decisions and the drift bias of the decision process for Go responses was

reduced (Figure 6a,c). Conversely, slow waves within frontal electrodes correlated with a

reduction of v_{NoGo}, indicating that evidence accumulation was slower for NoGo Decisions

(Figure 6b). These modelling results provide a potential explanation on how local slow

wave properties relate to mental states (Figure 4) and single-trial task performance (Figure

5). In other words, posterior slow waves were associated with sluggish responses and

increased misses while frontal slow waves were associated with faster, impulsive responses

and more false alarms. Indeed, slower evidence accumulation in favour of Go responses

would lead to slower reaction times or even misses, whereas slower evidence accumulation

for NoGo responses would lead to faster responses and possibly false alarms.

Figure 6. Global and local effects of the occurrence of slow waves on sub-

components of decision-making

Reaction Times in the Go/NoGo tasks were modelled according to a Hierarchical Drift

Diffusion Model (see Online Methods). a-f: Topographical maps of the effect of slow

waves (i.e. whether or not a slow wave was detected for each trial and for a specific

electrode) on the parameters of decision-making: drift Go [v_{Go}] (a), drift NoGo [v_{NoGo}]

(b), drift bias [v_{Bias}] (c), threshold [a] (d), Non-decision time or NDT [t] (e), decision

bias [z] (f). The effect of slow wave occurrence was estimated with LMEs (see Online

 Methods) and topographies show the scalp distribution of the associated t-values. Black

dots denote significant clusters of electrodes (pcluster<0.05, Bonferroni-corrected, see

Online Methods).

These results suggest that slow waves could represent both a general index of fatigue as

well as a mechanism underlying specific behavioural and subjective consequences of

sleepiness. In other words, regardless of where it happens, slow waves can reflect the global

"sleepiness" of an individual. Meanwhile, at a finer grain, depending on where they occur,

slow waves can disrupt specific cognitive processes carried out by the affected brain

regions. Taken together, we interpret this as strong evidence in support of the idea that slow

waves are a compelling physiological phenomenon, which precedes and co-occurs with

subjective and behavioural aspects of attentional lapses. In addition, our results indicate

that spatio-temporal properties of slow waves can specifically explain distinctive

components of behavioural and mental consequences of attentional lapses in a unified and

quantitative manner.

Discussion

According to both in-lab and real-life studies, humans spend up to half of their waking

lives not paying attention to their environment or any task at hand[34,35]. However, the

ubiquitousness of these attentional lapses remains unexplained. Part of the difficulty in

identifying their neural correlates could be due to their intrinsic diversity[7,8]. We embraced

this challenge to clarify the neural mechanisms underlying attentional lapses by linking

three different levels of explanation: behaviour, phenomenology and physiology. To grasp

the phenomenological diversity of attentional lapses, we distinguish between MW and MB,

where MW is operationally defined as a shift in attention away from environmental

demands and toward spontaneous task-unrelated thoughts, and MB as the absence of

thoughts and an empty mind. We propose that these two types of attentional lapses can be

explained by the occurrence of sleep-like low-frequency high-amplitude waves, where

regional differences in the occurrence of slow waves predicts the type of attentional lapse

as well as its behavioural profile. These slow waves have been previously linked to the

transition toward sleep when they are visible across the scalp[38] or to the concept of local

sleep when they occur in a subset of electrodes[11,22,25,29,30] and in the context of an awake

individual.

The concept of local sleep builds upon a recent questioning of the classical view of sleep

as an all-or-nothing phenomenon[22–24,41,42]. Although sleep is orchestrated at the scale of the

entire brain, some of the key neural mechanisms underlying sleep are implemented and

regulated at the level of local cortical circuits[24,25]. Consequently, when the pressure for

sleep increases, an awake brain can start displaying local sleep-like patterns of activity such

as delta or theta waves[21,22,29,43]. These bouts of local sleep-like activity, in the form of large-

amplitude slow waves, are both time-dependent (i.e. increase with time spent awake) and

use-dependent (i.e. depend on the level of activation of a given brain region)[29,30]. The

occurrence of slow waves has been linked to perturbations of information processing and

task-related errors or attentional lapses in animal and human intracranial data[21,22]. A similar

relationship between local sleep and behavioural errors has been evidenced when detecting

slow waves in non-invasive scalp recordings of sleep-deprived[29,30] or well-rested[31]

individuals.

Importantly, our study proposes a novel link between sleep-like slow waves in normal

wakefulness and both behaviour and subjective experience. Specifically, we showed that,

within the same individual, slow waves temporally precede different types of attentional

lapses. At the behavioural level, we observed that sluggish responses (slow responses and

misses) were associated with an increase in slow waves over posterior electrodes (Figure

5a,c). Conversely, impulsive responses (hasty responses and false alarms) were associated

with an increase in slow waves over frontal electrodes (Figure 5a-b). At the

phenomenological level, compared to task-focused (ON) reports, mind wandering (MW)

was preceded by an increase in slow-wave density, amplitude, upward and downward

slopes over frontal electrodes (Figure 4a). Compared to ON reports, mind blanking (MB)

was also preceded by an increase in slow-wave density over frontal electrodes, but we

additionally observed an increase in slow-wave upward and downward slopes over

posterior electrodes (Figure 4b). A direct comparison between MW and MB reports shows

that MB reports were associated with slow waves with steeper upward slopes over posterior

electrodes and smaller amplitude over frontal electrodes (Figure 4c). Finally, focusing on

shorter time windows (Figure S4-6), we showed that slow waves best predict mental states

(MW or MB vs ON) in the last few seconds before a probe, in line with previous

findings[44,45].

These results imply a spatial and temporal relationship between markers of local sleep and

behavioural errors: local sleep events, as measured through the presence of slow waves,

occurring at the right time (i.e. during stimulus presentation) and in the right place (i.e. in

the brain regions involved in the task) are predictive of distinct behavioural and

phenomenological aspects of attentional lapses[11].

Importantly, our results are largely consistent with previous findings on the neural

correlates of attentional lapses. Most of these studies focused on mind wandering, although

often defined as any mental state that is not on task (i.e. attentional lapses, which include

MW and MB as defined here). Early fMRI studies showed that mind wandering in this

sense was associated with the activation of the Default Mode Network (DMN)[46,47].

Relevant to our local sleep interpretation, the DMN is also suggested to be involved in

dream generation during sleep, consistent with the idea that the DMN supports a broad

array of experiences that are decoupled from the environment[48]. Furthermore, lesions

within the DMN are associated with a decrease in both mind wandering[49] and dreaming[50].

However, recent findings suggest a complex relationship between the DMN and

spontaneous experiences. Unlike what was initially hypothesized, the DMN is now

considered to be involved in both task-unrelated and task-related processes[44,51], depending

for example on environmental demands or the vividness of individuals' experiences[52,53].

Based on our results, we speculate that local sleep plays a key role in the complex

relationship between the DMN and spontaneous experiences. Previous studies have indeed

shown that a state of low alertness could induce the phasic activation of the DMN[54]. We

speculate that slow waves occurring within the DMN could lead to episodes of mind

wandering by disconnecting individuals from their environment[55]. Indeed, slow waves

during sleep have been proposed to be responsible for disconnecting sleepers from their

environment[27,56], as they are accompanied by a phenomenon of neuronal silencing that can

disrupt the processing of external inputs. If slow waves trigger specific occurrences of mind

wandering, other mechanisms could be responsible for the stability of these episodes[57].

Further investigations, including source localization or simultaneous recording of EEG and

fMRI[58,59], promise a deeper understanding of the mechanisms underlying these attentional

lapses.

Our work might also help reconcile previous findings on the EEG correlates of mind

wandering. A seminal study reported a reduction of alpha and/or beta oscillations during

mind wandering,[60] whereas others have reported an increase[61–63] . However, the relation

between alpha and sleep is complex: alpha power is low both when participants are fully

alert and, on the contrary, when approaching sleep onset[64,65]. Perhaps, the divergent results

obtained in previous studies regarding mind wandering and alpha oscillations could be

explained by different baseline levels of alpha oscillations in these studies. Likewise, past

pupilometry results on attentional lapses have been similarly inconclusive. While most

studies found a dampening of stimulus-locked increases in pupil size during mind

wandering (e.g. [47,66,67]), results diverge for baseline pupil size, which has been reported as

either increasing (e.g. [68,69]) or decreasing (e.g. [47,70,71]). When distinguished from mind

wandering, mind blanking has also been associated with reduced pupil size compared to

task-focused states[10,66]. Our results largely align with the latter results, with both mind

wandering and mind blanking being characterized by a decrease in pre-probe pupil size

(Figure 2e), which goes together with low vigilance ratings (Figure 2d) and an increase in

slow wave occurrence (Figure 3-4). In addition, the complex relations between pupil size

and mind wandering may be partly explained by the fact that pupil size does not index only

arousal but also correlates with motivation[72] and cognitive load[73], which also correlate with

mind wandering[2]. In contrast, the sleep-like nature of wake slow waves would make them

an unambiguous marker of sleepiness. Furthermore, slow waves are detected at the

electrode level and can therefore indicate how brain regions respond differently to sleep

pressure.

Other than mind wandering and blanking, we foresee that the spatio-temporal properties of

slow waves might predict other types of spontaneous experiences. For example, previous

research focusing on the hypnagogic period at sleep onset has shown that slow-wave like

activities are predictive of the occurrence of spontaneous imagery[74], the intensity of

thoughts[58] and can discriminate between different contents of spontaneous experiences[75].

Likewise, during sleep, it has been reported that local modulation of slow wave power is

predictive of the occurrence and contents of dreams[76]. Local sleep, therefore, might be

related to the type and content of spontaneous experiences not only during wakeful states,

as we showed here, but also during sleep-wake transitions and sleep.

Our results also speak to the broader question of how different brain regions participate in

shaping the stream of consciousness. Slow waves, considered as a perturbation of the

normal wake activity of local cortical networks, could reflect the functions of these

networks under normal situations. For example, we observed that slow waves in frontal

regions were associated with false alarms (Figure 5b), which aligns well with the role of

frontal cortices in executive functions and response inhibition[77]. Conversely, slow waves

in the back of the brain were associated with misses (Fig 5c), which is consistent with the

involvement of parietal cortices in sensorimotor integration[78]. At the phenomenological

level, frontal slow waves were associated with mind wandering, whereas posterior slow

waves were associated with mind blanking. Thus our results could speak to the debate on

the respective involvement of frontal and posterior cortices in supporting different

conscious states[79,80]. Our results suggest that a perturbation of frontal cortices leads to

unconstrained thoughts (mind wandering) rather than the loss of awareness, but that

awareness decreases when posterior regions momentarily go offline, a pattern similar to

the neural correlates of dreaming during sleep or spontaneous thoughts during wakeful

rest[58,76]. However, frontal and posterior slow waves do not only differ by their location but

also in terms of spatial expanse: slow waves in frontal electrodes appears more focal,

whereas slow waves in posterior electrodes is more widespread (Figure S10). Thus, the

loss of awareness during mind blanking could reflect a tendency for posterior slow waves

to involve a broader fronto-parietal network. This is in line with theories attributing an

essential role of fronto-parietal connections in the emergence of consciousness[81–83].

We speculate that the slow waves we report here are generated by similar neural

mechanisms as slow waves in sleep. Our speculation rests on the profile of slow waves

characterized in (i) time (Figure 3a) and (ii) space (Figure 3b,c), and the relationships

between slow wave properties and (iii) time spent on task (Figure S3), (iv) subjective

vigilance and (v) pupil size. While these five lines of evidence are correlational, we note

that together, they imply a high degree of similarity between slow waves in waking and

sleep. This, in turn, allows us to interpret wake slow waves as local sleep. Importantly, as

our study builds on convergent lines of indirect evidence to argue that the slow waves we

measured in waking participants are sleep-related, this interpretation is tentative and we do

not wish to suggest that the presence of slow-frequency oscillations in wake EEG is by

itself sufficient to establish a relation to sleep. Here, for example, we paired these

observations with objective (pupil size) and subjective (vigilance ratings) markers of

fatigue and checked that slow waves increased with the time spent on task. Future studies

could use direct evidence from intracranial recordings or sleep deprivation to more solidly

establish this interpretation. Intracranial recordings in particular can confirm whether wake

slow waves are accompanied by episodes of neuronal silencing, as for sleep slow waves,

and would help the identification of the exact neural mechanisms that generate the slow

waves we report here.

As our task involved the continuous presentation of visual stimuli, we could not fully

disentangle the occurrence of slow waves and task-related events. Yet, we showed that

slow waves differ from typical responses evoked by stimuli or participants' responses

(Figure S11). Future studies could include task-free resting state sessions to compare the

occurrence and properties of slow waves occurring during a task and without a task.

Relatedly, further investigation could help determine to which extent our findings

generalize to everyday situations (e.g., driving, attending a lecture, reading, etc).

It is worth noting that the tasks used here (SARTs) are rather undemanding and could

favour sleepiness and local sleep compared to more difficult and engaging experimental

paradigms. Generalising our findings to different experimental contexts or more

naturalistic settings would need further experimental validation[11]. From our findings, we

predict that slow waves would predict the occurrence of attentional lapses and mind

wandering in situations where participants feel rather sleepy and to a lesser extent or not at

all in situations where participants are well awake, motivated and highly engaged. We also

speculate that slow waves are not the only mechanism that underlies sensory decoupling in

wakefulness and sleep[56]. Other factors, such as the neuromodulation of brain activity, may

be critical in clarifying the total set of mechanisms underlying mind wandering[9,84] and

blanking, and more broadly attentional lapses.

In conclusion, we show here that attentional lapses occurring in the context of an

undemanding task are accompanied by sleep-like slow waves, even when participants are

well-rested. Furthermore, the location of slow waves is predictive of certain behavioural

and phenomenological properties of these lapses. Thus, we propose that these slow waves

reflect local intrusions of sleep within waking and constitute a mechanistic and proximate

cause to explain attentional lapses. Identifying a proximate mechanism of attentional lapses

could inspire novel applications leveraging brain-machine interfaces in educational or

professional settings.

581 **Online Methods**

Participants. Thirty-two (N=32) healthy adults were recruited and participated in this

study. Six individuals were not included in our analyses because of technical issues during

recordings or an abnormal quality of physiological recordings assessed through a post-hoc

visual inspection of the data. The remaining 26 participants (age: 29.8 ± 4.1 years, mean $\pm$

standard-deviation; 10 females) were included in all analyses except for one individual for

whom we do not have eye-tracking data.

Experimental Design and Stimuli. Participants were seated in a dimly lit room with their

chin resting on a support at approximately 60cm from a computer screen. All task

instructions and stimuli were displayed and button responses were collected via the

Psychtoolbox extension[85] for Matlab (Mathworks, Natick, MA, USA).

The experimental design consisted of two modified Sustained Attention to Response Tasks

(SARTs)[32] in which participants were instructed to pay attention to a series of pictures of

human faces in the Face SART blocks or digits in the Digit SART blocks. The order of

Face or Digit blocks was pseudo-randomized for each participant. Each block lasted

approximately 12 to 15 minutes. Participants were allowed to rest between blocks.

Participants performed 3 Face SART blocks and 3 Digit SART blocks for a total duration

of 103 ± 19.7 minutes (mean $\pm$ standard-deviation) from beginning to end. Each type of

the Face and Digit SART was preceded by a brief training session (27 trials) on each SART.

During this SART training session, feedback on the proportion of correct trials and average

reaction times was provided to participants. Participants were encouraged to prioritize

accuracy over speed.

Face and digit stimuli were presented continuously, each stimulus appearing for a duration

of 750 to 1250ms (random uniform jitter). Face stimuli were extracted from the Radboud

Face Database[86] and consisted of 8 faces (4 females) with a neutral facial expression and

one smiling female face. Digits from 1 to 9 were displayed with a fixed font size. For the

Face SART, participants were instructed to press a button for all neutral faces (Go trials)

but to avoid pressing the response button for the smiling face (NoGo trials). The order of

faces was pseudo-randomized throughout the entire task (i.e. we permuted the presentation

order every 9 stimuli and we did not present twice the same stimuli in a row). For the Digit

SART, participants were instructed to press a button for all digits except the digit "3"

(NoGo trials), with the order of the digits pseudo-randomized as well.

During the SART, we stopped the presentation of stimuli at random times (every 30 to 70s,

random uniform jitter) with a sound and the word "STOP" displayed on the screen. These

interruptions allowed us to probe the mental state of the participants with a series of 8

questions (including 1 conditional question; see Supplementary Methods). In particular,

we instructed participants to report their attentional focus "just before the interruption".

Participants had to select one of the four following options: (1) "task-focused" (i.e.

focusing on the task, ON), (2) "off-task" (i.e. focusing on something other than the task,

which we define here as mind wandering MW), (3) "mind blanking" (i.e. focusing on

nothing), (4) "don't remember". As the 4th option accounted for only 1.1% of all probes

(i.e. less than 1 probe per participant on average) and since previous studies do not always

distinguish between these options (e.g. [87]), we collapsed the 3rd and 4th options as mind-

blanking (MB) in all analyses. We also instructed participants to rate their level of

vigilance, reflecting "over the past few trials", with a 4-point scale (Figure 2d; from

1="Extremely Sleepy" to 4="Extremely Alert"). Each of the 12-15 min SART blocks

included 10 interruptions (in total, 30 interruptions for each SART task and 60 interruptions

per participant). Participants were informed of the presence of interruptions and the nature

of each question before starting the experiment. The mental state categories (ON, MW and

MB) were also explained to participants orally and in writing.

Physiological Recordings and Preprocessing. High-density scalp electroencephalography

(EEG) was recorded using an EasyCap (63 active electrodes) connected to a BrainAmp

system (Brain Products GmbH). A ground electrode was placed frontally (Fpz in the 10-

20 system). Electrodes were referenced online to a frontal electrode (AFz). Additional

electrodes were placed above and below the left and right canthi respectively to record

ocular movements (electrooculogram, EOG). Two electrodes were placed over the deltoid

muscles to record electrocardiographic (ECG) activity. EEG, EOG and ECG were sampled

at 500Hz. Eye-movements and pupil size on one eye were recorded with an EyeLink 1000

system (SR Research) with a sampling frequency of 1000Hz. The eye-tracker was

calibrated at the start of each recording using the EyeLink acquisition software.

The EEG signal was analysed in Matlab with a combination of the SPM12, EEGlab[88] and

Fieldtrip[89] toolboxes. The raw EEG signal was first high-pass filtered above 0.1 Hz using

a two-pass 5th-order Butterworth filter. A notch filter was then applied (stop-band: [45, 55]

Hz, 4th-order Butterworth filter) to remove line noise. Electrodes that were visually

identified as noisy throughout the recording were removed and interpolated using

neighbouring electrodes. Finally, the continuous EEG data was segmented according to

probe onsets on a 64s window ([-32, 32] s relative to the probe onset); the average voltage

over the entire window (64s) was then removed for each electrode and probe.

Pupil size was analysed with custom functions in Matlab and corrected for the occurrence

of blinks (see [33] and Supplementary Methods). Pupil size was averaged over the stimulus

presentation window for each trial (window length: 0.75 to 1.25s). Pupil size values in

Figure 2e were computed by averaging the pupil size in all trials within 20s preceding the

probe onset and then by discretizing them into 5 bins across all probes for each participant

and task, to normalize pupil size across participants[33].

Behavioural Analyses. Go trials were considered incorrect (Miss) if no response was

recorded between stimulus onset and the next stimulus onset. Conversely, NoGo trials were

considered incorrect (false alarm) if a response was recorded between stimulus onset and

the next stimulus onset. Reaction times were computed from the onset of the stimulus

presentation. Trials with reaction times shorter than 300ms were excluded from all analyses

(so not considered correct or incorrect). In all subsequent analyses, we focused only on

trials within 20s from probe onset. The choice of a 20s-window allowed the inclusion of 2

NoGo trials for each probe, while focusing on trials that are relatively close to the probe

onset and subsequent subjective reports. Since previous studies sometimes examined

shorter pre-probe time windows (e.g. [44]), we also analysed the data with shorter time

windows (Table S3 and Figure S4-6).

Detection of Slow Waves. The detection of sleep-like slow waves in waking was based on

previous algorithms devised to automatically detect slow waves during NREM sleep[30,37].

First, the preprocessed EEG signal was re-referenced to the average of the left and right

mastoid electrodes (TP9 and TP10) to match the established guidelines for sleep

recordings[90]. Then, the signal was down-sampled to 128Hz and band-pass filtered within

the delta band. A type-2 Chebyshev filter was used to reach an attenuation of at least 25

dB in the stop-band ([0.1, 15] Hz) but less than 3 dB in the pass-band ([1, 10] Hz). All

waves were detected by locating the negative peaks in the filtered signal. For each wave,

the following parameters were extracted: start and end point (defined as zero-crossing

respectively prior the negative peak of the wave and following its positive peak), negative

peak amplitude and position in time, positive peak amplitude and position in time, peak-

to-peak amplitude, downward (from start to negative peak) and upward (from negative to

positive peak) slopes.

Slow waves in sleep typically have a larger negative peak compared to their positive peak

(Figure 3a) and are predominantly observed over fronto-central channels[37,38] (Figure 3b).

This contrasts with artefacts in the EEG signal caused by blinks, which typically have a

large positive component and are more frontally distributed. To reduce the false detection

of these artefacts as candidate slow waves, we excluded waves with a positive peak above

75µV. We also excluded waves within 1s of large-amplitude events (>150µV of absolute

amplitude). Finally, we discarded all waves that were shorter than 143ms in duration

(corresponding to a frequency > 7Hz). We then selected the waves with the highest

absolute peak-to-peak amplitude (top 10% computed for each electrode independently) as

local sleep slow waves. This 10% threshold was selected based on the visual examination

of the distribution of the amplitudes at the subject level (see Figure S12 for the

corresponding topography). The mean ± SEM of the threshold voltage across subjects was

30.7 ± 1.5 µV. Finally, we compared slow waves with neural activity related to task events

(stimulus onsets and motor responses). To do so, we computed the event related potentials

(ERPs) by averaging, across participants, the EEG signal band passed around 0.1 and 30Hz

and referenced to the mastoids. These ERPs were locked either on slow-waves' start,

stimulus onset (for face and digit stimuli separately) or motor responses. The corresponding

temporal profiles are shown in Figure S11a-c for electrode Cz.

Hierarchical Drift Diffusion Modelling. Hierarchical Bayesian Drift Diffusion Modelling

(HDDM) was used to extend our analysis beyond simple behavioural metrics and examine

the impact of mental states (Figure S9) and slow waves (Figure 6) on the sub-processes of

decision making. The DDM is a sequential-sampling model of 2AFC decision making that

can be considered an extension of signal detection theory into the time-domain, accounting

for full reaction time distributions as well as choice behaviour[39]. The HDDM package[40] in

Python 2.8 was used to fit the drift-diffusion model to the SART data. DDM parameters

were estimated using a hierarchical Bayesian method that uses Markov-chain Monte Carlo

(MCMC) sampling to generate full posterior distributions of model parameters. The

following DDM parameters were estimated: the drift rate for Go trials (v_{Go}), the drift rate

for NoGo trials (v_{NoGo}), the decision threshold (a), the decision bias (z) and the non-decision

time (t). Drift bias (v_{Bias}) was computed by taking the difference between the absolute

values of v_{Go} (positive) and v_{NoGo} (negative), where greater values indicate stronger v_{Go}

drift bias (Figure S7). To examine whether the model could reproduce key patterns in the

SART data, posterior predictive checks were undertaken by simulating 100 datasets from

the joint posteriors of model parameters and comparing these to the observed data[91] (Figure

S8).

To estimate HDDM parameters 8,000 samples from the posterior were generated with

MCMC methods and the initial 2,000 were discarded as burn-in to reduce autocorrelation.

HDDM models were first fit on the trials preceding subjective reports to examine the

influence of mental states on decision parameters (Figure S9). HDDM parameters were

free to vary by state and task. From the estimated models, we extracted the subject-level

point-estimates of parameters as the mean of each individual's posterior distribution for a

given task (Face and Digit) and mental state (ON, MW or MB). We then fitted HDDM

models so as to examine the influence of slow waves (event present vs. absent; Figure 6).

To do so we considered each EEG electrode separately. For a given electrode, a trial was

flagged as being associated with slow waves if the onset of a slow wave was detected for

this electrode during stimulus presentation (i.e. between stimulus onset and offset).

Parameters were also free to vary by task (Digit vs. Face). When examining the impact of

mental states or slow waves on HDDM parameters, we included trials within 20s of a probe

onset (allowing to include 2 NoGo trials for each probe). From the estimated models, we

extracted the subject-level point-estimates of parameters as the mean of each individual's

posterior distribution for a given task (Face and Digit) and slow waves (present or absent).

Statistical comparisons were performed on the subject-level point-estimates.

Statistical Analyses. Statistics were performed using Linear Mixed-Effects modelling

(LMEs). In all models, subject identity was coded as a categorical random effect. The task

type (Digit or Face SART) was used as a categorical fixed effect in all analyses. Several

fixed effects were independently tested in our different analyses: mental state (categorical

variable: ON, MW and MB; Figure 2-4) or slow waves (binary variable: present/absent,

Figure 5). LMEs were run to predict different variables of interest: behavioural variables

(misses, false alarms, reaction times) or physiological variables (pupil size, presence or

properties of slow waves). We also used LMEs to estimate the effect of mental states or

slow waves on the point-estimates derived from the HDDM models (Figure S9 and Figure

6). Model comparisons were performed using a Likelihood Ratio Test to estimate the

influence of multi-level categorical variables such as mental states. In practice, we

compared a model including mental state (along possibly other random and fixed effects)

with a model not including mental state as a predictor. All models and model comparisons

are described in Supplementary Table 1. In the Results section, we report the Likelihood

Ratio Test as χ^2(df), where χ^2 is the Likelihood Ratio Test statistic and df the degrees of

freedom[92,93]. When several model comparisons were performed for the same analysis using

the Likelihood Ratio Test, a Bonferroni correction was applied to the statistical threshold.

To indicate the magnitude and direction of the effects, we report the estimates (β) and

confidence interval (CI) for the contrasts of interest (MW vs. ON, MB vs ON, MB vs MW).

All models performed are described in Supplementary Table 1. For topographical maps,

clusters were identified using a cluster-permutation approach. In practice, for each

electrode, we extracted the t-values and p-values for the effects of interest. Clusters of

neighbouring electrodes were defined as electrodes with p-values below 0.025 (cluster

alpha). Once the clusters were defined, a comparison was performed with permuted

datasets and we used a Monte-Carlo p-value threshold of 0.05 to identify the significant

clusters ($p_{cluster}$, see Supplementary Methods for details). This threshold was corrected for

multiple comparisons when several non-independent cluster permutations were performed

(e.g. Figure 4-6, S2-4) in order to keep the type-1 error rate constant across analyses.

Acknowledgments

TA and NT were supported by Australian Research Council Discovery Projects (DP180104128 and DP180100396) and National Health Medical Research Council Ideas Projects (APP1183280). TA was supported by a Long-Term Fellowship from the Human Frontier Science Program (LT000362/2018-L). JW was supported by Australian Research Council Discovery Early Career Researcher Awards (DE170101254). We thank Devangna Tangri for her help in data collection and Giulio Bernardi for sharing his slow-wave detection algorithm. We also thank our reviewers, Dr Tristan Bekeinschtein, Dr Jonathan Smallwood and a third anonymous reviewer for their constructive comments and criticisms.

Author Contributions

Design: TA, TM, JW & NT. Data collection: TA & TM. Analyses: TA & AB.

Manuscript: TA, AB, JW & NT.

Data Availability

All raw data and code generated for this study are publicly available:

Raw data: https://osf.io/ey3ca/?view_only=680c39e7065649c3b783a4efec0a1a94
Code used for analyses: https://github.com/andrillon/wanderIM

References

1. Chun, M. M., Golomb, J. D. & Turk-Browne, N. B. A Taxonomy of External and Internal Attention. *Annual Review of Psychology* **62**, 73–101 (2011).

2. Smallwood, J. & Schooler, J. W. The Science of Mind Wandering: Empirically Navigating the Stream of Consciousness. *Annual Review of Psychology* **66**, 487–518 (2015).

3. Ward, A. F. & Wegner, D. M. Mind-blanking: when the mind goes away. *Frontiers in Psychology* **4**, (2013).

4. Seli, P., Risko, E. F., Smilek, D. & Schacter, D. L. Mind-Wandering With and Without Intention. *Trends in Cognitive Sciences* **20**, 605–617 (2016).

5. VanRullen, R. Perceptual Cycles. *Trends in Cognitive Sciences* **20**, 723–735 (2016).

6. Johnson, K. A. *et al.* Response variability in Attention Deficit Hyperactivity Disorder: Evidence for neuropsychological heterogeneity. *Neuropsychologia* **45**, 630–638 (2007).

7. Seli, P. *et al.* Mind-Wandering as a Natural Kind: A Family-Resemblances View. *Trends in Cognitive Sciences* **22**, 479–490 (2018).

8. Christoff, K. *et al.* Mind-Wandering as a Scientific Concept: Cutting through the Definitional Haze. *Trends Cogn. Sci. (Regul. Ed.)* **22**, 957–959 (2018).

9. Mittner, M., Hawkins, G. E., Boekel, W. & Forstmann, B. U. A Neural Model of Mind Wandering. *Trends in Cognitive Sciences* **20**, 570–578 (2016).

10. Unsworth, N. & Robison, M. K. Tracking arousal state and mind wandering with pupillometry. *Cognitive, Affective, & Behavioral Neuroscience* **18**, 638–664 (2018).

11. Andrillon, T. *et al.* Does the Mind Wander When the Brain Takes a Break? Local

Sleep in Wakefulness, Attentional Lapses and Mind-Wandering. *Frontiers in*

Neuroscience **13**, (2019).

12. Drummond, S. P. A., Paulus, M. P. & Tapert, S. F. Effects of two nights sleep

deprivation and two nights recovery sleep on response inhibition. *Journal of Sleep*

Research **15**, 261–265 (2006).

13. Doran, S. M., Van Dongen, H. P. & Dinges, D. F. Sustained attention performance

during sleep deprivation: evidence of state instability. *Arch Ital Biol* **139**, 253–267

(2001).

14. Metin, B. *et al.* ADHD performance reflects inefficient but not impulsive information

processing: A diffusion model analysis. *Neuropsychology* **27**, 193–200 (2013).

15. Poh, J.-H., Chong, P. L. H. & Chee, M. W. L. Sleepless night, restless mind: Effects

of sleep deprivation on mind wandering. *Journal of Experimental Psychology:*

General **145**, 1312–1318 (2016).

16. Madiouni, C., Lopez, R., Gély-Nargeot, M.-C., Lebrun, C. & Bayard, S. Mind-

wandering and sleepiness in adults with attention-deficit/hyperactivity disorder.

Psychiatry Research **287**, 112901 (2020).

17. Goupil, L. & Bekinschtein, T. A. Cognitive processing during the transition to sleep.

Arch Ital Biol **150**, 140–154 (2012).

18. Nielsen, T. Microdream neurophenomenology. *Neuroscience of Consciousness* **2017**,

(2017).

19. Cirelli, C. & Tononi, G. Is sleep essential? *PLoS Biol.* **6**, e216 (2008).

20. Tononi, G. & Cirelli, C. Sleep and the Price of Plasticity: From Synaptic and Cellular

Homeostasis to Memory Consolidation and Integration. *Neuron* **81**, 12–34 (2014).

21. Nir, Y. *et al.* Selective neuronal lapses precede human cognitive lapses following

sleep deprivation. *Nature Medicine* **23**, 1474–1480 (2017).

22. Vyazovskiy, V. V. *et al.* Local sleep in awake rats. *Nature* **472**, 443–7 (2011).

23. Krueger, J. M., Nguyen, J. T., Dykstra-Aiello, C. J. & Taishi, P. Local sleep. *Sleep*

Medicine Reviews **43**, 14–21 (2019).

24. Rector, D. M., Schei, J. L., Van Dongen, H. P. A., Belenky, G. & Krueger, J. M.

Physiological markers of local sleep. *European Journal of Neuroscience* **29**, 1771–

1778 (2009).

25. D'Ambrosio, S. *et al.* Sleepiness as a Local Phenomenon. *Frontiers in Neuroscience*

13, (2019).

26. Steriade, M. *Neuronal Substrates of Sleep and Epilepsy*. vol. 1 (Cambridge

University Press, 2003).

27. Tononi, G. & Massimini, M. Why does consciousness fade in early sleep? *Ann. N. Y.*

Acad. Sci. **1129**, 330–334 (2008).

28. Andrillon, T., Poulsen, A. T., Hansen, L. K., Leger, D. & Kouider, S. Neural Markers

of Responsiveness to the Environment in Human Sleep. *Journal of Neuroscience* **36**,

6583–6596 (2016).

29. Hung, C.-S. *et al.* Local experience-dependent changes in the wake EEG after

prolonged wakefulness. *Sleep* **36**, 59–72 (2013).

30. Bernardi, G. *et al.* Neural and Behavioral Correlates of Extended Training during

Sleep Deprivation in Humans: Evidence for Local, Task-Specific Effects. *Journal of*

Neuroscience **35**, 4487–4500 (2015).

31. Quercia, A., Zappasodi, F., Committeri, G. & Ferrara, M. Local Use-Dependent

Sleep in Wakefulness Links Performance Errors to Learning. *Frontiers in Human*

Neuroscience **12**, (2018).

32. Robertson, I. H., Manly, T., Andrade, J., Baddeley, B. T. & Yiend, J. 'Oops!':

performance correlates of everyday attentional failures in traumatic brain injured and

normal subjects. *Neuropsychologia* **35**, 747–758 (1997).

33. van Kempen, J. *et al.* Behavioural and neural signatures of perceptual decision-

making are modulated by pupil-linked arousal. *eLife* **8**, (2019).

34. Killingsworth, M. A. & Gilbert, D. T. A Wandering Mind Is an Unhappy Mind.

Science **330**, 932–932 (2010).

35. Seli, P., Carriere, J. S. A., Levene, M. & Smilek, D. How few and far between?

Examining the effects of probe rate on self-reported mind wandering. *Frontiers in*

Psychology **4**, (2013).

36. Joshi, S., Li, Y., Kalwani, R. M. & Gold, J. I. Relationships between Pupil Diameter

and Neuronal Activity in the Locus Coeruleus, Colliculi, and Cingulate Cortex.

Neuron **89**, 221–234 (2016).

37. Riedner, B. A. *et al.* Sleep homeostasis and cortical synchronization: III. A high-

density EEG study of sleep slow waves in humans. *Sleep* **30**, 1643–57 (2007).

38. Siclari, F. *et al.* Two Distinct Synchronization Processes in the Transition to Sleep: A

High-Density Electroencephalographic Study. *SLEEP* (2014)

doi:10.5665/sleep.4070.

39. Ratcliff, R., Smith, P. L., Brown, S. D. & McKoon, G. Diffusion Decision Model:

Current Issues and History. *Trends in Cognitive Sciences* **20**, 260–281 (2016).

40. Wiecki, T. V., Sofer, I. & Frank, M. J. HDDM: Hierarchical Bayesian estimation of

the Drift-Diffusion Model in Python. *Frontiers in Neuroinformatics* **7**, (2013).

41. Nir, Y. *et al.* Regional slow waves and spindles in human sleep. *Neuron* **70**, 153–69

(2011).

42. Siclari, F. & Tononi, G. Local aspects of sleep and wakefulness. *Current Opinion in

Neurobiology* **44**, 222–227 (2017).

43. Rector, D. M., Topchiy, I. A., Carter, K. M. & Rojas, M. J. Local functional state

differences between rat cortical columns. *Brain Research* **1047**, 45–55 (2005).

44. Sormaz, M. *et al.* Default mode network can support the level of detail in experience

during active task states. *Proceedings of the National Academy of Sciences* **115**,

9318–9323 (2018).

45. Ellamil, M. *et al.* Dynamics of neural recruitment surrounding the spontaneous

arising of thoughts in experienced mindfulness practitioners. *NeuroImage* **136**, 186–

196 (2016).

46. Mason, M. F. *et al.* Wandering minds: the default network and stimulus-independent

thought. *Science* **315**, 393–5 (2007).

47. Mittner, M. *et al.* When the Brain Takes a Break: A Model-Based Analysis of Mind

Wandering. *The Journal of Neuroscience* **34**, 16286–16295 (2014).

48. Fox, K. C. R., Nijeboer, S., Solomonova, E., Domhoff, G. W. & Christoff, K.

Dreaming as mind wandering: evidence from functional neuroimaging and first-

person content reports. *Frontiers in Human Neuroscience* **7**, (2013).

49. Philippi, C. L. *et al.* Lesion network mapping demonstrates that mind-wandering is

associated with the default mode network. *J Neurosci Res* **99**, 361–373 (2021).

50. Zellner, M. R. Dreaming and the Default Mode Network: Some Psychoanalytic

Notes. *Contemporary Psychoanalysis* **49**, 226–232 (2013).

51. Konu, D. *et al.* A role for the ventromedial prefrontal cortex in self-generated

episodic social cognition. *NeuroImage* **218**, 116977 (2020).

52. Turnbull, A. *et al.* Left dorsolateral prefrontal cortex supports context-dependent

prioritisation of off-task thought. *Nature Communications* **10**, (2019).

53. Turnbull, A. *et al.* The ebb and flow of attention: Between-subject variation in

intrinsic connectivity and cognition associated with the dynamics of ongoing

experience. *NeuroImage* **185**, 286–299 (2019).

54. Tagliazucchi, E. & Laufs, H. Decoding Wakefulness Levels from Typical fMRI

Resting-State Data Reveals Reliable Drifts between Wakefulness and Sleep. *Neuron*

82, 695–708 (2014).

55. Kam, J. W. Y. *et al.* Slow Fluctuations in Attentional Control of Sensory Cortex.

Journal of Cognitive Neuroscience **23**, 460–470 (2011).

56. Andrillon, T. & Kouider, S. The vigilant sleeper: neural mechanisms of sensory

(de)coupling during sleep. *Current Opinion in Physiology* (2019)

doi:10.1016/j.cophys.2019.12.002.

57. Smallwood, J. Distinguishing how from why the mind wanders: A process–

occurrence framework for self-generated mental activity. *Psychological Bulletin* **139**,

519–535 (2013).

58. Perogamvros, L. *et al.* The Phenomenal Contents and Neural Correlates of

Spontaneous Thoughts across Wakefulness, NREM Sleep, and REM Sleep. *Journal*

of Cognitive Neuroscience 1–12 (2017) doi:10.1162/jocn_a_01155.

59. Hawes, S. *et al. Sleeping While Awake: The Intrusion of Neural Activity Associated*

with Sleep Onset in the Awake Human Brain.

http://biorxiv.org/lookup/doi/10.1101/2020.06.04.133603 (2020)

doi:10.1101/2020.06.04.133603.

60. Braboszcz, C. & Delorme, A. Lost in thoughts: Neural markers of low alertness

during mind wandering. *NeuroImage* **54**, 3040–3047 (2011).

61. Compton, R. J., Gearinger, D. & Wild, H. The wandering mind oscillates: EEG alpha

power is enhanced during moments of mind-wandering. *Cognitive, Affective, &*

Behavioral Neuroscience **19**, 1184–1191 (2019).

62. Macdonald, J. S. P., Mathan, S. & Yeung, N. Trial-by-Trial Variations in Subjective

Attentional State are Reflected in Ongoing Prestimulus EEG Alpha Oscillations.

Frontiers in Psychology **2**, (2011).

63. Arnau, S. *et al.* Inter-trial alpha power indicates mind wandering. *Psychophysiology*

57, (2020).

64. Hori, T., Hayashi, M. & Morikawa, T. Topographical EEG changes and the

hypnagogic experience. in *Sleep onset: Normal and abnormal processes.* (eds.

Ogilvie, R. D. & Harsh, J. R.) 237–253 (American Psychological Association, 1994).

65. Jagannathan, S. R. *et al.* Tracking wakefulness as it fades: Micro-measures of

alertness. *NeuroImage* **176**, 138–151 (2018).

66. Unsworth, N. & Robison, M. K. Pupillary correlates of lapses of sustained attention.

Cognitive, Affective, & Behavioral Neuroscience **16**, 601–615 (2016).

67. Smallwood, J. *et al.* Pupillometric Evidence for the Decoupling of Attention from

Perceptual Input during Offline Thought. *PLoS ONE* **6**, e18298 (2011).

68. Smallwood, J. *et al.* Insulation for Daydreams: A Role for Tonic Norepinephrine in

the Facilitation of Internally Guided Thought. *PLoS ONE* **7**, e33706 (2012).

69. Franklin, M. S., Broadway, J. M., Mrazek, M. D., Smallwood, J. & Schooler, J. W.

Window to the Wandering Mind: Pupillometry of Spontaneous Thought While

Reading. *Quarterly Journal of Experimental Psychology* **66**, 2289–2294 (2013).

70. Grandchamp, R., Braboszcz, C. & Delorme, A. Oculometric variations during mind

wandering. *Frontiers in Psychology* **5**, (2014).

71. Konishi, M., Brown, K., Battaglini, L. & Smallwood, J. When attention wanders:

Pupillometric signatures of fluctuations in external attention. *Cognition* **168**, 16–26

(2017).

72. Varazzani, C., San-Galli, A., Gilardeau, S. & Bouret, S. Noradrenaline and Dopamine

Neurons in the Reward/Effort Trade-Off: A Direct Electrophysiological Comparison

in Behaving Monkeys. *Journal of Neuroscience* **35**, 7866–7877 (2015).

73. Kahneman, D. & Beatty, J. Pupil Diameter and Load on Memory. *Science* **154**, 1583–

1585 (1966).

74. Germain, A. & Nielsen, T. A. EEG power associated with early sleep onset images

differing in sensory content. *Sleep Research Online* **4**, 83–90 (2001).

75. Noreika, V. *et al.* Intrusions of a drowsy mind: neural markers of phenomenological

unpredictability. *Frontiers in Psychology* **6**, (2015).

76. Siclari, F. *et al.* The neural correlates of dreaming. *Nature Neuroscience* **20**, 872–878

(2017).

77. Simmonds, D. J., Pekar, J. J. & Mostofsky, S. H. Meta-analysis of Go/No-go tasks

demonstrating that fMRI activation associated with response inhibition is task-

dependent. *Neuropsychologia* **46**, 224–232 (2008).

78. Lynch, J. C. The functional organization of posterior parietal association cortex.

Behavioral and Brain Sciences **3**, 485–499 (1980).

79. Boly, M. *et al.* Are the Neural Correlates of Consciousness in the Front or in the Back

of the Cerebral Cortex? Clinical and Neuroimaging Evidence. *J. Neurosci.* **37**, 9603–

9613 (2017).

80. Odegaard, B., Knight, R. T. & Lau, H. Should a Few Null Findings Falsify Prefrontal

Theories of Conscious Perception? *The Journal of Neuroscience* **37**, 9593–9602

(2017).

81. Dehaene, S., Changeux, J.-P., Naccache, L., Sackur, J. & Sergent, C. Conscious,

preconscious, and subliminal processing: a testable taxonomy. *Trends in Cognitive*

Sciences **10**, 204–211 (2006).

82. Mashour, G. A., Roelfsema, P., Changeux, J.-P. & Dehaene, S. Conscious Processing

and the Global Neuronal Workspace Hypothesis. *Neuron* **105**, 776–798 (2020).

83. Dehaene, S., Kerszberg, M. & Changeux, J. P. A neuronal model of a global

workspace in effortful cognitive tasks. *Proc Natl Acad Sci U S A* **95**, 14529–34

(1998).

84. O'Callaghan, C., Walpola, I. C. & Shine, J. M. Neuromodulation of the mind-

wandering brain state: the interaction between neuromodulatory tone, sharp wave-

ripples and spontaneous thought. *Phil. Trans. R. Soc. B* **376**, 20190699 (2021).

85. Brainard, D. H. The Psychophysics Toolbox. *Spat Vis* **10**, 433–436 (1997).

86. Langner, O. *et al.* Presentation and validation of the Radboud Faces Database.

Cognition & Emotion **24**, 1377–1388 (2010).

87. Van den Driessche, C. *et al.* Attentional Lapses in Attention-Deficit/Hyperactivity

Disorder: Blank Rather Than Wandering Thoughts. *Psychological Science* **28**, 1375–

1386 (2017).

88. Delorme, A. & Makeig, S. EEGLAB: an open source toolbox for analysis of single-

trial EEG dynamics including independent component analysis. *J. Neurosci. Methods*

134, 9–21 (2004).

89. Oostenveld, R., Fries, P., Maris, E. & Schoffelen, J.-M. FieldTrip: Open source

software for advanced analysis of MEG, EEG, and invasive electrophysiological

data. *Comput Intell Neurosci* **2011**, 156869 (2011).

90. Iber, C., Ancoli-Israel, S., Chesson, A. & Quan, S. The AASM Manual for the

Scoring of Sleep and Associated Events: Rules, Terminology and Technical

Specifications. (2007).

91. Gelman, A., Meng, X.-L. & Stern, H. Posterior predictive assessment of model

fitness via realized discrepancies. *Statistica sinica* 733–760 (1996).

92. Bates, D., Mächler, M., Bolker, B. & Walker, S. Fitting Linear Mixed-Effects Models

Using **lme4**. *Journal of Statistical Software* **67**, (2015).

93. Harrison, X. A. *et al.* A brief introduction to mixed effects modelling and multi-

model inference in ecology. *PeerJ* **6**, e4794 (2018).

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2   **Supplementary Information:**
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4   **Predicting lapses of attention with sleep-like**
5   **slow waves**.
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23   **Supplementary Methods**

24   **Supplementary Table 1 to 3**

25   **Supplementary Figures 1 to 12**

Supplementary Methods

Participants. Prior to their participation in the protocol, all 26 participants but one filled in online surveys on Qualtrics (N=25). They reported normal levels of sleepiness (Epworth Sleepiness Scale: 14.6 ± 4.7; mean ± standard-deviation) and mind wandering (Mind Wandering Questionnaire[1]: 3.6 ± 0.91) in their everyday lives. Participants were not instructed to follow a particular schedule before the experiment (in particular, our protocol did not involve a sleep restriction procedure). The timing of the experiment was determined by participants' availability and included both morning and afternoon sessions.

Experimental Design and Stimuli. Face stimuli were divided in two parts vertically (half-left and half-right faces) which were flickered on the screen at different frequencies (12 and 15 Hz, counterbalanced across participants). Similarly, the digits were inserted in a Kanizsa illusory square (Fig. 1a) whose right and left parts also flickered at different frequencies (12 and 15 Hz). This flickering was introduced to elicit Steady State Visual Evoked Potentials (SSVEPs) in the EEG signal. The detailed analysis of this aspect of our dataset will be reported elsewhere. As the flickering occurred at a high rate, participants did not report a negative effect on their ability to perform the SART.

Experience Sampling. Following task interruptions (probes), participants were asked to answer a series of 8 questions in the following fixed order: (1) 'Were you looking at the screen?' (response: yes / no); (2) 'Where was your attention focus?' (response: on-task / off-task / blank / don't remember); (3) 'What distracted your attention from the task?' (response: Something in the room / personal / about the task); (4) 'How aware were you of your focus?' (response: from 1, I was fully aware, to 4, I was not aware at all); (5) 'Was your state of mind intentional?' (response: from 1, entirely intentional, to 4, entirely unintentional); (6) 'How engaging were your thoughts?' (response: from 1, not engaging, to 4, very engaging); (7) 'How well do you

think you have been performing?' (response: from 1, not good, to 4, very good); (8) 'How alert

have you been?' (response: extremely alert / alert / sleepy / extremely sleepy). Question 3 was

displayed only if participants answered off-task in question 1. In this report, we focus only on

questions (2) and (8).

Physiological Recordings and Preprocessing. The raw pupil size was corrected for the

occurrence of blinks as in[2]. The timings of blinks were obtained through the EyeLink

acquisition software. For each of these blinks, the pupil size was corrected by linearly

interpolating the average signal preceding the blink onset ([-0.2, -0.1]s) and following the blink

offset ([0.1, 0.2]s). The corrected signal was then low-pass filtered below 6Hz (two-pass

Butterworth filter at the 4th order). Finally, for blinks longer than 2s, data points between -0.1s

prior to blink onset and 0.1s after blink onset were considered missing.

Detection of Slow Waves. In sleep, according to established guidelines[3], only waves with peak-

to-peak amplitude exceeding 75µV are defined as slow waves. In wakefulness, similar slow

waves can be observed but with smaller amplitudes. Accordingly, previous studies relied on a

relative rather than absolute threshold[4–6]. Here, we defined slow waves as the waves with

absolute peak-to-peak amplitude within the top 10% of all the waves detected for a given EEG

electrode and a given participant. On average, the detection threshold was 30 µV (average

across N=26 participants and across all electrodes).

Figure 3a shows the average waveform of the slow waves detected on electrode Cz as well as

the average waveform of waves detected during sleep recording in another published dataset

(N=15 participants)[7]. To compute the average waveform of sleep slow waves, we applied the

same algorithm to epochs of 20s scored as NREM2 and NREM3. Only slow waves with peak-

to-peak amplitude over 75µV were considered.

We also compared the average waveform of slow waves with task-related Event Related

Potentials (ERPs). To do so, we averaged the EEG signal time-locked to stimuli or response

onsets at the electrode level. Figure S11a-c shows the average ERP across participants for

electrode Cz as well as the scalp topographies of the voltage observed across the scalp at the

time of the peak for electrode Cz. Finally, we also compared single-trial voltage values between

the average waveforms for slow waves and stimulus-locked or response-locked ERPs (Figure

S11d-f, data shown for one participant). Overall, these analyses showed a clear difference

between the amplitude, waveform and topographies of slow waves compared to task-related

ERPs.

Drift Diffusion Modelling. The Drift Diffusion Model (DDM) assumes that a decision

variable noisily accumulates evidence from a starting point (z) with drift rate (v) towards one

of two boundaries that represent choice alternatives (i.e. 'Go' or 'No-go'; see Figure S7). The

decision threshold (a) is the distance between the two boundaries and represents the amount of

evidence that must be accumulated before a decision is made. Once the decision variable

crosses a decision boundary, a response is made. Five parameters were fitted using a DDM

approach: the participants' initial bias for one of the two responses (decision bias, z), drift rates

for Go and NoGo responses (v_{Go} and v_{NoGo}), the decision threshold (a), the non-decision time

parameter (t). The last parameter, t, captures extra-decisional components, including stimulus

encoding, response preparation and execution. We also extracted the drift rate bias (v_{Bias}) as

the difference between the absolute values of v_{Go} (positive) and v_{NoGo} (negative). We performed

a model selection based on the Deviance Information Criteria (DIC), which assess goodness of

fit while accounting for model complexity in hierarchical models[8]. With posterior predictive

checks, we confirmed that our DDM was able to generate simulated behavioural data that are

similar to the recorded data (Figure S8; based on the Fz model). For this check, we simulated

100 datasets based on the posterior distributions of HDDM parameters.

Statistics. A cluster-permutation approach (derived from [9]) was applied to identify significant

clusters in topographical maps. Candidate clusters were defined as neighbouring electrodes

with a p-value below a threshold (called "cluster alpha") of 0.025. For each candidate cluster,

we computed the sum of the t-values for all the electrodes belonging to the cluster (which we

will refer to as the "cluster statistics"). We then created permuted datasets by permuting the

labels of the predictor within each subject, each task and each electrode (N=1,000

permutations). For each of these permuted datasets, we also identified the candidate cluster

with maximal absolute cluster statistics. The cluster statistics from permutations formed a null

distribution, against which we compared the cluster statistics from the real dataset. Clusters

(real and permuted) with positive and negative cluster statistics were compared separately. A

Monte-Carlo p-value was derived from this comparison ($p_{cluster}<0.05$ means that a negative

cluster has a cluster statistics below the 5th percentile of the permuted distribution and that a

positive cluster has a cluster statistics above the 95th percentile of the permuted distribution).

In cases where several cluster-permutations were performed in the same analysis (Fig. 5 and

6), we corrected the Monte-Carlo p-values of the real clusters with the Bonferroni method.

References:

1. Mrazek, M. D., Phillips, D. T., Franklin, M. S., Broadway, J. M. & Schooler, J. W. Young and restless: validation of the Mind-Wandering Questionnaire (MWQ) reveals disruptive impact of mind-wandering for youth. *Frontiers in Psychology* **4**, (2013).

2. van Kempen, J. *et al.* Behavioural and neural signatures of perceptual decision-making are modulated by pupil-linked arousal. *eLife* **8**, (2019).

3. Iber, C., Ancoli-Israel, S., Chesson, A. & Quan, S. The AASM Manual for the Scoring of Sleep and Associated Events: Rules, Terminology and Technical Specifications. (2007).

4. Hung, C.-S. *et al.* Local experience-dependent changes in the wake EEG after prolonged wakefulness. *Sleep* **36**, 59–72 (2013).

5. Bernardi, G. *et al.* Neural and Behavioral Correlates of Extended Training during Sleep Deprivation in Humans: Evidence for Local, Task-Specific Effects. *Journal of Neuroscience* **35**, 4487–4500 (2015).

6. Quercia, A., Zappasodi, F., Committeri, G. & Ferrara, M. Local Use-Dependent Sleep in Wakefulness Links Performance Errors to Learning. *Frontiers in Human Neuroscience* **12**, (2018).

7. Legendre, G., Andrillon, T., Koroma, M. & Kouider, S. Sleepers track informative speech in a multitalker environment. *Nature Human Behaviour* **3**, 274–283 (2019).

8. Spiegelhalter, D. J., Best, N. G., Carlin, B. P. & van der Linde, A. Bayesian measures of model complexity and fit. *Journal of the Royal Statistical Society: Series B (Statistical Methodology)* **64**, 583–639 (2002).

9. Maris, E. & Oostenveld, R. Nonparametric statistical testing of EEG- and MEG-data. *J. Neurosci. Methods* **164**, 177–190 (2007).

Supplementary Table 1. Summary of Linear Mixed-Effects Models

Figure	Predicted Variable X	Level	Predictor Of Interest	Model 0	Model 1
2a-c	False Alarms, Misses, Reaction Times	Probe	Mental State (MS)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + MS + (1 | Subject)
2d-e	Vigilance Scores, Pupil Size	Probe	Mental State (MS)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + MS + (1 | Subject)
N/A	Pupil Size	Probe	Vigilance Ratings (Vig)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + Vig + (1 | Subject)
N/A	Vigilance Scores	Probe	Slow Wave (SW) Properties (density, amplitude, upward slope and downward slope; average across electrodes)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + SW + (1 | Subject)
N/A	Pupil Size	Probe	SW Properties (density, amplitude, upward slope and downward slope; average across electrodes)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + SW Properties + (1 | Subject)
4*	Mental State (binary contrast: MW vs ON; MB vs ON and MB vs MW).	Probe	SW Properties (density, amplitude, downward slope and upward slope per electrode)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + SW Properties + (1 | Subject)
5*	False Alarms, Misses, Reaction Times	Trial	SW (presence or absence)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + SW + (1 | Subject)
6*	a, t, z, v_{Go}, v_{NoGo}, vBias	Subject	SW (presence or absence)	X ~ 1 + Task + (1 | Subject)	X ~ 1 + Task + SW + (1 | Subject)

142   *: *Analyses corrected for multiple comparison (see Online and Supplementary Methods).*

144 **Supplementary Table 2. Effect of normalisation on behavioural analyses**

		Normalised	Before Normalisation
Misses (%) in Go trials	State	$\chi^2(2)$=33.4 (***)	$\chi^2(2)$=36.0 (***)
	MW vs ON	**-0.0096 [-0.015, -0.0041]**	**-0.011 [-0.016, -0.0052]**
	MB vs ON	**-0.023 [-0.032, -0.015]**	**-0.023 [-0.032, -0.015]**
	MB vs MW	**-0.014 [-0.022, -0.0055]**	**-0.013 [-0.021, -0.0047]**
False alarms (%) in NoGo trials	State	$\chi^2(2)$=88.5 (***)	$\chi^2(2)$=115.9 (***)
	MW vs ON	**-0.24 [-0.29, -0.19]**	**-0.20 [-0.24, -0.17]**
	MB vs ON	**-0.20 [-0.26, -0.19]**	**-0.17 [-0.23, -0.12]**
	MB vs MW	0.064 [-0.014, 0.14]	0.028 [-0.028, 0.084]
RT (s) in Go trials	State	$\chi^2(2)$=33.4 (***)	$\chi^2(2)$=16.9 (**)
	MW vs ON	-0.0005 [-0.013, 0.011]	-0.0025 [-0.0096, 0.0045]
	MB vs ON	**0.039 [0.022, 0.057]**	**0.019 [0.0088, 0.030]**
	MB vs MW	**0.040 [0.022, 0.058]**	**0.022 [0.011, 0.032]**

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The analyses of the effect of mental states (model comparisons and post-hoc contrasts) were consistent
even after normalizing Misses, False alarms and Reactions Times (RT) for each participant by the
average obtained on ON probes ("Normalised" column). The "State" row shows the result of the
likelihood ratio test (chi-squared value). Stars denote the corrected significance level (Bonferroni
*correction for 6 likelihood ratio tests: ***: p<0.001, **: p<0.01, *: p<0.05). The estimates of the post-*
hoc contrasts (MW vs ON, MB vs ON, MB vs MW) obtained from the mixed effect models are also
reported, with the corresponding 95% confidence intervals. Bold fonts signal the significant results
(p<0.05 for model comparisons, or confidence intervals excluding 0 for the post-hoc contrasts).

Supplementary Table 3. Effect of different pre-probe window sizes on behavioural analyses

		20s before Probes	10s before Probes
Misses (%) in Go trials	State	$\chi^2(2)=36.0$ (***)	$\chi^2(2)=31.3$ (***)
	MW vs ON	**-0.011 [-0.016, -0.0052]**	**-0.012 [-0.019, -0.0037]**
	MB vs ON	**-0.023 [-0.032, -0.015]**	**-0.033 [-0.045, -0.021]**
	MB vs MW	**-0.013 [-0.021, -0.0047]**	**-0.021 [-0.033, -0.0096]**
False alarms (%) in NoGo trials	State	$\chi^2(2)=115.9$ (***)	$\chi^2(2)=55.97$ (***)
	MW vs ON	**-0.20 [-0.24, -0.17]**	**-0.20 [-0.25, -0.15]**
	MB vs ON	**-0.17 [-0.23, -0.12]**	**-0.14 [-0.22, -0.062]**
	MB vs MW	0.028 [-0.028, 0.084]	0.061 [-0.018, 0.14]
RT (s) in Go trials	State	$\chi^2(2)=16.9$ (**)	$\chi^2(2)=8.42$ (ns)
	MW vs ON	-0.0025 [-0.0096, 0.0045]	0.0026 [-0.0066, 0.012]
	MB vs ON	**0.019 [0.0088, 0.030]**	**0.020 [0.0064, 0.034]**
	MB vs MW	**0.022 [0.011, 0.032]**	**0.018 [0.0038, 0.031]**

The analyses of the effect of mental states (model comparisons and post-hoc contrasts) were replicated using a shorter window of 10s (right column) instead of 20s before probe onsets (left column). Likelihood Ratio Tests are reported for the effect of State on the correctness for Go and NoGo trials as well as the Reaction Times (RT) for Go Trials. Stars denote the corrected significance level (Bonferroni correction for 6 comparisons: ***: p<0.001, **: p<0.01, *: p<0.05). The estimates for the post-hoc contrasts and the corresponding 95% confidence intervals are also shown. Bold fonts signal the significant results (p<0.05 for model comparisons, or confidence intervals excluding 0 for the post-hoc contrasts).

Supplementary Figure 1. Sleepiness is associated with more, larger and steeper slow waves

Slow waves detected within 20s of probes' onset were separated according to participants' vigilance ratings (from 1 (extremely sleepy) to 4 (extremely alert)) provided after each probe. The density (a), amplitude (c), downward slope (D-Slope, e) and upward slope (U-Slope, f), averaged across participants and electrodes, are shown with error-bars showing the standard-error-of-the-mean across participants. Topographies of the average density (b), amplitude (d), downward slope (f) and upward slope (h) are also shown for the different vigilance levels.

Supplementary Figure 2. A reduction in pupil size is associated with more, larger and steeper slow waves

Slow waves detected within 20s of probes' onset were separated according to participants' pupil size (binned, see Methods) computed on the same window. The density (a), amplitude (c), downward slope (D-Slope, e) and upward slope (U-Slope, f), averaged across participants and electrodes, are shown with error-bars showing the standard-error-of-the-mean across participants. Topographies of the average density (b), amplitude (d), downward slope (f) and upward slope (h) are also shown for the different pupil size bins.

Supplementary Figure 3. Slow waves increase with time spent on task

(a) Topographies of the temporal density of slow waves (slow waves per minute, averaged across all 26 subjects). An increase in the number of slow waves, maximal over central electrodes, can be seen from the first to the last experimental block. (b-d) Slow wave density for electrode Cz (b), vigilance ratings (c) and pupil size (d) averaged within each block. Connected larger dots show the average across participants (error-bars: SEM). Individual semi-transparent circles show individual subjects.

MW > ON

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Supplementary Figure 4. Spatiotemporal dynamics of the effect of slow-waves properties on mental states (MW vs ON).

*Mixed-Effects Models were used to quantify the impact of slow-wave properties (**a**: Density; **b**: Amplitude; **c**: Downward Slope (D-Slope); **d**: Upward Slope (U-Slope)) on mental states as in Figure 4. Slow waves were detected on 4 different windows: [-20, -15]s, [-15, -10]s, [-10, -5]s and [-5, 0]s before probe onsets. Slow-wave properties were extracted for each electrode and used to predict the MW vs ON contrast. Topographies show the scalp distribution of the t-values associated with each slow-wave parameter and electrode. Black dots denote significant clusters of electrodes ($p_{cluster}<0.05$ corrected for 48 comparisons (Figure S4-6) using a Bonferroni approach).*

Supplementary Figure 5. Spatiotemporal dynamics of the effect of slow-waves properties on mental states (MB vs ON).

Same format as in Supplementary Figure 4.

Supplementary Figure 6. Spatiotemporal dynamics of the effect of slow-waves properties on mental states (MB vs MW).

Same format as in Supplementary Figure 4.

Supplementary Figure 7. Drift Diffusion Modelling

a: The Go/NoGo tasks were modelled according to the Drift Diffusion Model (DDM, see Online and Supplementary Methods). The following parameters were estimated: threshold (a), non-decision time or NDT (t), bias (z), drift rate for Go trials (v_{Go}), drift rate for NoGo trials (v_{NoGo}). We then also computed the drift bias: $abs(v_{Go})-abs(v_{NoGo})$. The figure shows a graphical representation of these parameters. Note that here, drift rates for NoGo trials are negative. b-c: Graphical representation of decision processes using the parameters obtained by the DDM for trials with (SW+) or without (SW-) slow waves. The presence or absence of slow waves was determined for each electrode (FCz (b; frontal) and Oz (c; posterior)). Note the reduction in decision threshold, drift rates and bias associated with slow waves but the increase in NDT.

Supplementary Figure 8. HDDM Fit to Behavioural Data

Posterior predictive checks of Go/No-Go DDM fit to the behavioural data. Observed data for all participants (black bars) are plotted underneath model-predicted RT distributions and No-Go choice proportions (red lines). Positive distribution represents the normalised frequency of reaction times (RT) from Go responses. Negative bin at RT=-1 represents the proportion of No-Go responses.

Supplementary Figure 9. Impact of mental states on HDDM parameters.

Hierarchical Drift Diffusion Modeling (HDDM, see Methods) was applied to the Reaction
Times obtained in the Face (top) and Digit (bottom) SART. The parameters were fitted for each
task and mental state. Each panel shows the distribution of the estimated variable for
individual participants (drift for NoGo and Go trials, drift bias, threshold, non-decision time
(NDT) and decision bias; see Methods). For each plot, coloured areas show the smoothed
distribution of individual data points (see Methods). Diamonds and circles show individual
estimates for the Face and Digit SART respectively. Box plots show the 1st and 3rd quartiles
(edges) as well as the median (middle bar). Blue boxes around individual plots and stars next
to the titles indicate variables with significant state-effects (model comparison, see Methods;
*: p<0.05, Bonferroni correction for 12 comparisons). Significant differences between MW
and MB (for threshold and decision bias) are highlighted with red brackets.

Supplementary Figure 10. Spatial expanse of slow waves

Four seeds electrodes were selected along the scalp midline from the front (Fz) to back (Oz).
For each slow wave detected in these seed electrodes, we computed the probability that slow
waves were also observed in the other electrodes. The average co-occurrence probability
averaged across participants (N=26) is shown for each seed electrode. Note that slow waves
detected over Fz tend to co-occur with other slow waves only in a limited number of
neighboring frontal electrodes whereas occipital slow waves (Oz) tend to co-occur with other
slow waves in both frontal and posterior electrodes (more widespread).

Supplementary Figure 11. Slow waves compared to stimulus and response-locked activity

Event-related potentials averaged over electrode Cz and across participants for slow waves (a), stimulus-locked activity (b) and response-locked activity (c). Slow waves' EEG time courses are aligned with the first negative crossing before the negative peak. Stimulus-locked responses are aligned on the onset of the digit (full line) or face (dotted line) stimuli. Response-locked responses are aligned to the onset of participants' responses. Shaded areas show the SEM across participants (N=26). Insets show the topography of the average voltage at the times shown by the vertical dotted lines (a: 0.05s; b: 0.2s; c: -0.05s), which approximates the absolute maximum of each ERP. (d-f) Voltage of the event-related potentials per trial for one participant. Each row corresponds to (d) an individual slow wave, (e) a stimulus presentation (face or digit), (f) a motor response. We limited the number of rows in (e) and (f) to match the number of detected slow waves (N=120) in (d).

Supplementary Figure 12. Scalp topography of the average and standard-deviation of the slow-wave detection threshold

Only slow waves whose amplitude is within the top-10% of the distribution for each electrode and participant were analysed. The left topography shows the value, in voltage, of this threshold for each electrode, averaged across participants. The right topography shows the standard deviation (SD) of the threshold values across participants.