

SUBMITTED VERSION

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How much is left in your "sleep tank"? Proof of concept for a simple model for sleep history feedback

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Abstract: Technology-supported methods for sleep recording are becoming increasingly affordable. Sleep history feedback may help with fatigue-related decision making - Should I drive? Am I fit for work? This study examines a "sleep tank" model (SleepTank™), which is analogous to the fuel tank in a car, refilled by sleep, and depleted during wake. Required inputs are sleep period time and sleep efficiency (provided by many consumer-grade actigraphs). Outputs include suggested hours remaining to "get sleep" and percentage remaining in tank (Tank%). Initial validation was conducted using data from a laboratory-based simulated nightshift study. Ten, healthy males (18-35y) undertook an 8h baseline sleep opportunity and daytime performance testing (BL), followed by four simulated nightshifts (2000h-0600h), with daytime sleep opportunities (1000h-1600h), then an 8h night-time sleep opportunity to return to daytime schedule (RTDS), followed by daytime performance testing. Psychomotor Vigilance Task (PVT) and Karolinska Sleepiness Scale were performed at 1200h on BL and RTDS, and at 1830h, 2130h 0000h and 0400h each nightshift. A 40-minute York Driving Simulation was performed at 1730h, 2030h and 0300h on each nightshift. Model outputs were calculated using sleep period timing and sleep efficiency (from polysomnography) for each participant. Tank% was a significant predictor of PVT lapses ($p < 0.001$), and KSS ($p < 0.001$), such that every 5% reduction resulted in an increase of one lapse, or one point on the KSS. Tank% was also a significant predictor of %time in the Safe Zone from the driving simulator ($p = 0.001$), such that every 1% increase in the tank resulted in a 0.75% increase in time spent in the Safe Zone. Initial examination of the correspondence between model predictions and performance and sleepiness measures indicated relatively good predictive value. Results provide tentative evidence that this "sleep tank" model may be an informative tool to aid in individual decision-making based on sleep history.

37 **Abstract**

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39 history feedback may help with fatigue-related decision making – Should I drive? Am I fit for work?
40 This study examines a “sleep tank” model (SleepTank™), which is analogous to the fuel tank in a car,
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61

62 **1. Introduction**

63

64 Sleep loss results in increased likelihood of error and accident, in the workplace and on the roads
65 (Rajaratnam and Arendt 2001, Rogers, Hwang et al. 2004, Dorrian, Tolley et al. 2008). In Australia, in
66 any workplace covered by the Work Health and Safety Act (SafeWorkAustralia 2011), fatigue
67 management is a reciprocal responsibility for employers and employees, whereby the employers
68 have a duty of care to provide a safe workplace, and the employees have a responsibility to be fit for
69 work. Decisions about whether an employee is safe to start or to continue work are frequently self-
70 reported, and in some circumstances (e.g. truck cab, driver-only train cab) solo operators may only
71 have their own insight to rely on (Dorrian, Lamond et al. 2003). This leads us to consider the
72 information that people use to make decisions about whether they are sufficiently fit to start, or to
73 continue work.

74

75 Sleepiness is often used as an indicator of fatigue-related impairment, and in many circumstances, it
76 tracks performance measures (Kaida, Takahashi et al. 2006). However, this is not always the case
77 (Van Dongen, Maislin et al. 2003). Further, measuring sleepiness is not necessarily measuring
78 awareness of risk. Performance ratings often track sleepiness more closely than they track
79 performance (Dorrian, Lamond et al. 2000, Dorrian, Lamond et al. 2003, Dorrian, Roach et al. 2007;
80 Paech, Banks et al. 2016). Given these inconsistencies, an alternative to solely relying on self-
81 assessment could involve integration of technology-supported methods for sleep recording. Indeed,
82 consumer-grade sleep technologies (CST) are becoming increasingly affordable (Ko, Kientz et al.
83 2015) and may provide data that can scaffold personal monitoring systems to inform decisions
84 relating to fitness for work. While the proliferation of such devices supports their potential, current
85 barriers to effective use of CST include a lack of validation against research-grade actigraphy and
86 polysomnography, which is the gold-standard for sleep measurement.

87

88 Current international best practice in workplace fatigue management involves a Fatigue Risk
89 Management System (FRMS) (Dawson and McCulloch 2005, Gander, Hartley et al. 2011, Cabon,
90 Deharvenge et al. 2012, Gander, Mangie et al. 2014), which includes specific policy, education and
91 awareness training, fatigue monitoring systems with feedback, procedures for reporting,
92 investigating and recording fatigue-related incidents and accidents, and evaluation processes and
93 mechanisms for testing the impact of any fatigue-interventions (Gander, Hartley et al. 2011). In this
94 context, validated CSTs may assist with fatigue monitoring, however, it is critical to consider the
95 form of the feedback. At present, CSTs provide information such as prior sleep duration, timing, and
96 sometimes an indicator of sleep quality or efficiency. In operational settings, a worker needs to be
97 able to use sleep history information to assess what this may mean for their fatigue level when
98 starting work. Further, it is often necessary to project fatigue assessments into the future – towards
99 the end of my shift, will I still be fit for work, or for my commute?

100

101 Techniques to transform sleep history to indicate fatigue likelihood are already in use as part of
102 FRMS in a number of industries including, for example, rail (TransportCanada 2011) and healthcare
103 (SA_Ambulance 2008). These incorporate a simple calculation based on the allocation of “fatigue
104 likelihood points” according to the amount of sleep in the prior 24 and 48h, and the number of
105 hours that the employee has been awake (Dawson and McCulloch 2005). The calculation may be
106 performed by the worker (TransportCanada 2011), or may be automated based on simple inputs
107 (SA_Ambulance 2008), and is followed by a series of actions to take should the employee reach
108 critical thresholds indicating that fatigue, at levels of operational concern, is likely.

109

110 Another approach that has received much attention in the literature for transforming sleep history
111 into estimation of fatigue likelihood (or performance impairment, alertness, or effectiveness)
112 involves biomathematical modelling (Dinges 2004, Hursh, Redmond et al. 2004, Mallis, Mejdal et al.
113 2004, Van Dongen, Mott et al. 2007, Dawson, Noy et al. 2011). Frequently, the input is simply work

114 hours, from which sleep (and then the outcomes) are estimated (Kandelaars, Fletcher et al. 2006).
115 Sleep estimation is primarily based on the two-process model (Borbély 1982), which includes a
116 circadian oscillator, representing the sinusoidal 24h rhythm, and a homeostatic function, which
117 increases with wake, and decreases during sleep. These models are used as part of FRMS in industry
118 at an aggregate level to predict fatigue hotspots across rosters, target roster areas for
119 countermeasure application, and to compare potential rosters (Mallis, Banks et al. 2017).

120

121 A current area of development for biomathematical models is their application in the context of
122 predicting individual fatigue likelihood (Van Dongen, Mott et al. 2007, Dorrian, Darwent et al. 2012).
123 One challenge for individual modelling is the estimation of the circadian component, especially in
124 the context of irregular work hours and/or time zone changes, as are frequent in 24h industries
125 including aviation. The circadian system adapts slowly to sleep in new time zones, and is influenced
126 strongly by differences in light exposure (Mallis, Banks et al. 2017). In contrast, the homeostatic
127 function is relatively simple to model if sleep history is known. Further, since there is a circadian
128 rhythm to sleep length and quality (Van Dongen and Dinges 2005), measures of sleep timing, length,
129 and efficiency (as can be estimated from CSTs), these measures, to some extent, have an implicit
130 circadian signal. This study examines a model that mathematically transforms sleep history (timing,
131 length, efficiency), using our understanding of homeostatic component only of the two-process
132 model, that could be built-in to CSTs to provide useful feedback to aid in fatigue-related decision-
133 making and forward planning. This “sleep tank” model (SleepTank™¹) was invented by the second
134 author and enhances the homeostatic component of the two-process model with factors to account
135 for the effects of prolonged sleep restriction (Belenky, Wesensten et al. 2003, Van Dongen, Maislin
136 et al. 2003, Hursh, Redmond et al. 2004) and individual differences in sleep requirement. For this
137 study, the sleep tank was calibrated to an average sleep requirement of 8 hrs of sleep per day.

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¹ SleepTank™ is a trademark of the Institutes for Behavior Resources, Inc.

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2.0 Method

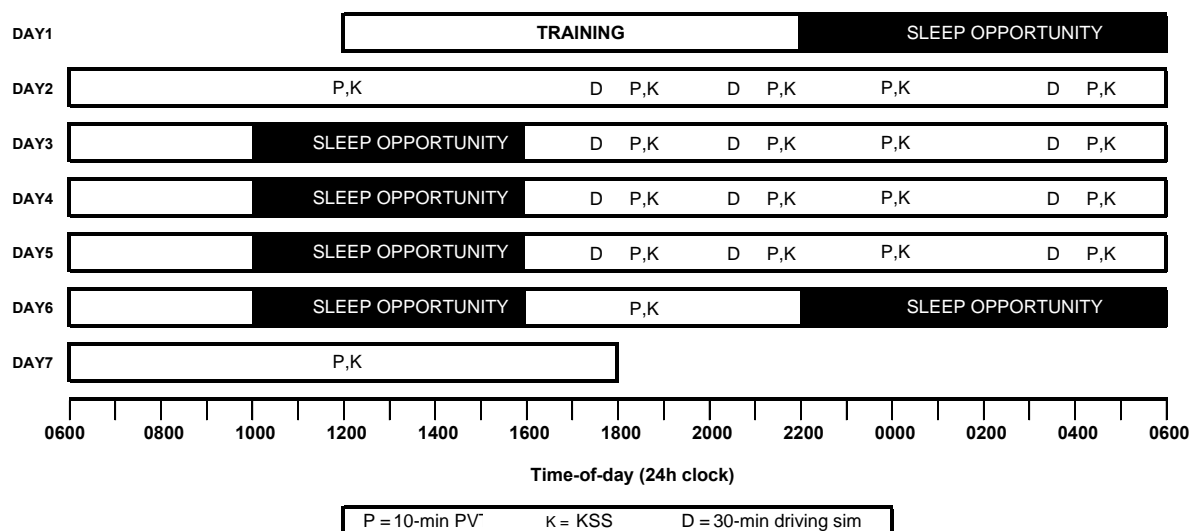
Initial validation of the “sleep tank” model (SleepTank™) was conducted using data from a simulated nightshift study. This study was approved by the University of South Australia Human Research Ethics Committee (0000033621) and was conducted in accordance with the Declaration of Helsinki.

2.1 Participants

Ten, healthy males (18-35y) were recruited using flyers and social media. Interested participants underwent screening. Inclusion criteria included good physical (confirmed by a general health questionnaire and fasting blood screen) and mental health (assessed by clinical history and Beck Depression Inventory) (Beck, Steer et al. 1996), a score between 22-43 on the Composite Morningness Questionnaire (Horne and Ostberg 1976), and habitual nightly sleep time between 7 and 9 h (confirmed by sleep diaries and wrist actigraphs, Actiwatch 2, Philips Respironics, Bend, OR). Exclusion criteria included reported: (a) sleep disorder (general health questionnaire) or sleep disturbance (>6 on the Pittsburgh Sleep Quality Index) (Buysse, Barzansky et al. 2003); (b) food allergy/ intolerance; (c) restrained eaters; (d) BMI >30 kg/m²; (e) use of prescription or over-the-counter medications known to affect glucose metabolism (Grant, Coates et al. 2017) or sleep; or (f) engagement in night shift work, trans-meridian travel, smoking, illicit drug use, excessive alcohol consumption (>2 standard drinks/day) or caffeine consumption (>2 cups/day) in the three months prior to the study. Participants were instructed to abstain from alcohol and caffeine in the week prior to the study (with compliance checked using a 3-day food diary completed on non-consecutive days).

164 **2.2 Procedure**

165 Participants (in groups of three or four) stayed in the Centre for Sleep Research laboratory at the
 166 University of South Australia for 7 days. This period included an 8 h baseline sleep opportunity
 167 (2200h-0600h) and daytime performance testing (BL), followed by four simulated nightshifts (2000h-
 168 0600h), with daytime sleep opportunities (1000h-1600h), then an 8h night-time sleep opportunity
 169 (2200h-0600h) to return to daytime schedule (RTDS), before a final period of daytime performance
 170 testing. Psychomotor Vigilance Task (PVT) and Karolinska Sleepiness Scale (KSS) were performed at
 171 1200h on BL and RTDS, and at 1830h, 2130h 0000h and 0400h each simulated nightshift. A 40-
 172 minute York Driving Simulation was performed at 1730h, 2030h and 0300h on each simulated
 173 nightshift (Figure 1). On BL, Day4 and RTDS, sleep was monitored using polysomnography.



174 **Figure 1. Protocol Diagram** – Time-of-day (24h clock) is on the x-axis, with day of study on the y-axis. Black
 175 bars indicate sleep opportunities and grey bars indicate simulated night shifts. BL=Baseline, Shift1-4=simulated
 176 night shifts, RTDS=return to daytime schedule. PVT=Psychomotor Vigilance Task, KSS=Karolinska Sleepiness
 177 Scale.
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180 The laboratory was sound-attenuated. Light levels were maintained at <50 lux during wake periods
 181 and <0.03 lux during sleep opportunities. Temperature was kept constant at 23 ± 1°C. No access to
 182 time cues was permitted throughout the protocol (i.e. clocks, mobile phones, television, internet).
 183 When not completing study tasks, participants were allowed to watch DVDs, play board games, or
 184 listen to music. Meals were controlled according to estimated energy requirement, calculated for

185 each participant given the sedentary laboratory protocol (Harris and Benedict 1918). Participants
186 were on one of two eating schedules (Grant, Coates et al. 2017). In one, participants had a meal
187 during their simulated night shifts, and in the other, energy was redistributed to times outside of
188 1900 to 0700h. Importantly, for all participants, 24h energy consumption was kept constant. Since
189 this was a proof-of-concept analysis to investigate whether the model would track average
190 performance and sleepiness across this schedule, eating groups were not looked at separately.
191 Moreover, differences in eating patterns are consistent with the workplace, where employees
192 differentially distribute their food consumption around the clock (Banks, Dorrian et al. 2014).

193

194 **2.3 Psychomotor Vigilance Task (PVT)**

195 The PVT was delivered via a hand-held response box. Participants were required to respond as
196 quickly as possible to a stimulus in the form of a red millisecond counter by pressing a response
197 button with the thumb of their dominant hand. When the button is pressed the millisecond counter
198 stops, the number representing the response time in milliseconds. The stimulus was displayed at
199 random intervals between 2 and 10 milliseconds, across the 10-minute task duration. This task is
200 sensitive to the effects of sleep loss and has a 1-3 trial learning curve (Dorrian, Rogers et al. 2005).
201 The variable for analysis in this manuscript was the average number of PVT lapses, defined as
202 response times greater than 500 msec.

203

204 **2.4 Karolinska Sleepiness Scale (KSS)**

205 Sleepiness was measured using the KSS, which is a 9-point scale ranging from 1=extremely alert to
206 9=extremely sleepy-fighting sleep. This scale has been used extensively and has been validated
207 against polysomnography and performance measures (Kaida, Takahashi et al. 2006).

208

209 **2.5 Driving Simulator**

210 Driving performance was measured using the York highway driving simulator (York Computer
211 Technologies, Kingston, ON). The simulator was run on a computer with a steering wheel and
212 accelerator and brake pedals. The simulated drive was a two-lane country road with road markings,
213 signs, and oncoming cars. Speed was displayed at the bottom of the screen and participants were
214 instructed to follow the road rules including a maximum speed limit of 100 km/h. This simulator has
215 demonstrated sensitivity in studies of sleep restriction (Arnedt, Wilde et al. 2000), with minimal
216 practice effects (De Valck, De Groot et al. 2003). The variable for analysis was the percentage of
217 time spent in safe zone, which was defined as within 10km/h of the speed limit and within 0.8m of
218 the centre of the lane.

219

220 **2.6 Polysomnography (PSG)**

221 Sleep period time and efficiency were measured using PSG with standard electrode placements
222 (C3/A2, C4/A1) during the baseline sleep, one of the 6 h daytime sleep opportunities, and during the
223 RTDS sleep.

224

225 **2.7 The “Sleep Tank”**

226 The “sleep tank” (SleepTank™), analogous to the fuel tank in a car, is refilled by sleep, and depletes
227 during wake. Required inputs are sleep period time and efficiency. Maximum tank size represents
228 the sleep-fuel required to remain awake for four days. The model focuses on the homeostatic
229 process of the two-process model. It does not include a circadian factor (i.e. it will have a known
230 residual error due to this rhythmic component). This “simplification” is deliberate to enable
231 immediate and continuous feedback from basic sleep inputs.

232

233 **2.8 Statistical Analyses**

234 In order to investigate changes in performance and sleepiness metrics across the simulated
235 nightshift protocol. Mixed effects ANOVA specified dependent variables of PVT Lapses, KSS, and

236 driving simulator %time in safe zone with fixed effects of shift (1-4), trial (PVT, KSS = 1830h/ 2130h
237 /0000h /0400h; %safe zone = 1730h/ 2030h/ 0300h) and shift* trial with a random effect of
238 subjectID on the intercept. In order to investigate the relationship between model predictions and
239 performance and sleepiness metrics, mixed effects regression specified dependent variables of PVT
240 Lapses, KSS, and driving simulator %time in safe zone with a fixed effect of the percentage left in the
241 “sleep tank” (Tank%) with a random effect of subjectID on the intercept. In order to estimate an
242 effect size for correlations between Tank% and performance and sleepiness metrics, time series
243 correlations were conducted for each individual. Since distributions of *r*-values are skewed, they
244 were transformed using Fisher’s *r*-*z* transformation, the average and standard error (sterr) across
245 participants was calculated, and then values were converted back to *r*-values using the inverse
246 Fisher function.
247

248 **3.0 Results**

249

250 Performance and sleepiness were significantly worse during the last test session of the shift
 251 compared to earlier trials, and sleepiness was significantly worse during the first shift compared to
 252 shifts 2-4 ($p < 0.01$, Table 1).

253

254 **Table 1. Performance Changes During Simulated Night Work** – Mixed effects ANOVA for PVT Lapses, KSS and
 255 driving simulator %time in safe zone with fixed effects of shift (1-4), trial (PVT, KSS = 1830h/ 2130h /0000h
 256 /0400h; %safe zone = 1730h/ 2030h/ 0300h) and shift* trial with a random effect of subjectID.

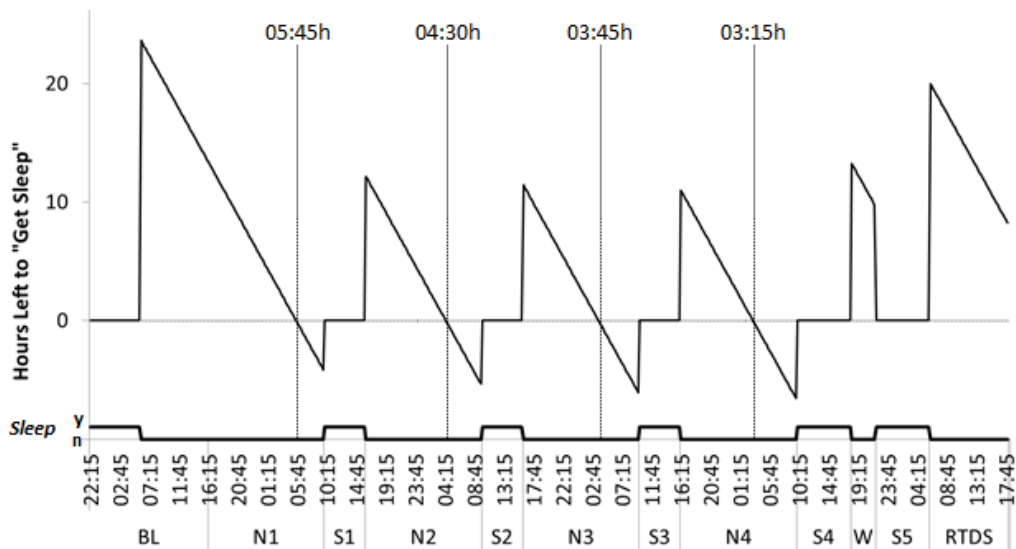
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	Shift			Trial			Shift*Trial			Post-hoc
	F	df	p	F	df	p	F	df	p	p<0.01
PVT	0.09	3,134.0	0.966	23.96	3,134.0	<0.001	0.25	9,134.0	0.985	1830h, 2130h, 0000h < 0300h
KSS	6.63	3,134.0	<0.001	62.02	3,134.0	<0.001	1.42	9,134.0	0.183	1830h < 0000h, 0300h Shift1 > Shifts2-4
%Safe Zone	2.04	3,99.0	0.113	10.24	2,99	<0.001	0.87	6,99.0	0.518	1730h, 2030h > 0300h

258

259

260 Average total sleep time during the baseline night was 7.11 h (SEM=0.05 h), during daytime sleep
 261 opportunities was 5.51 h (SEM=0.06), and during the return to daytime schedule night was 6.79 h
 262 (SEM=0.05 h). Average sleep efficiency was 89.5 % (SEM=0.6), 92.4% (SEM=1.1), and 84.9%
 263 (SEM=0.7) for each of these sleep opportunities respectively. Figure 2 illustrates the suggested
 264 hours left to “get sleep” for the laboratory study protocol. Model outputs were calculated using
 265 sleep period timing and sleep efficiency (for BL, day sleeps and RTDS using polysomnographic
 266 recordings) for each participant. These were then compared to study metrics of performance and
 267 sleepiness. On waking at BL, there is >20h “in the tank,” with a latest advisable bedtime of 5:45am.
 268 After waking from subsequent daytime sleep periods, the starting value “in the tank” is lower, with
 269 latest advisable bedtimes moving earlier in the shift across multiple nights.

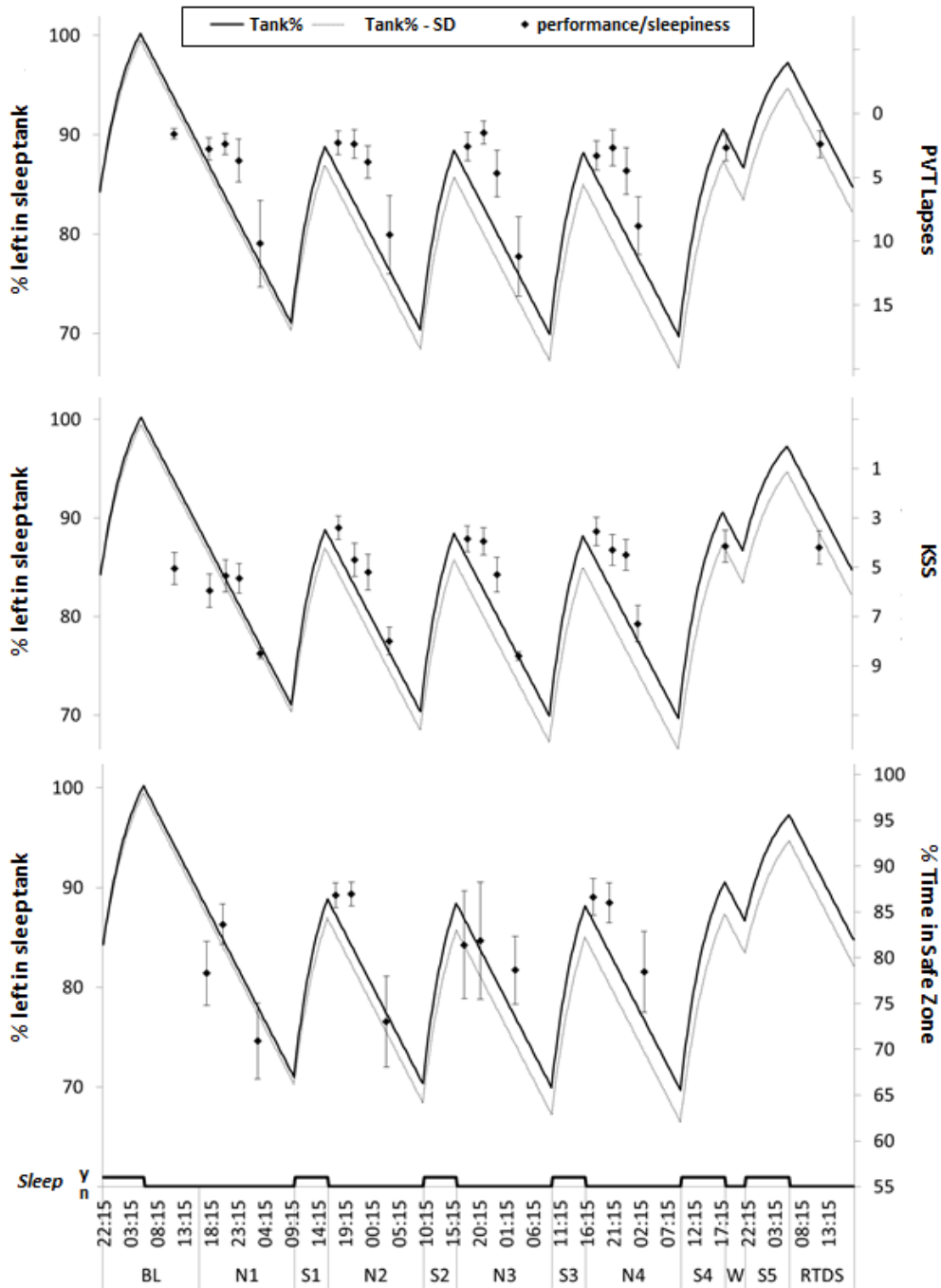


270 **Figure 2. Model Hours Left to “Get Sleep” Metric** – Time-of-day (24h clock) is on the x-axis, hours left to “get
 271 sleep” (an estimate of the latest advisable bedtime given what remains in the “Sleep Tank”) on the y-axis.
 272 Sleep opportunities are shown along the x-axis. BL=Baseline, N1-4=nightshifts, S1-4=Sleep opportunities,
 273 RTDS=return to daytime schedule. Times for $y=0$ (latest advisable bedtimes) are indicated.
 274

275 Figure 3 displays the percentage remaining in the tank (Tank%), which is highest on waking, with
 276 longer, more efficient sleep periods filling the tank to a greater level. Tank% was a significant
 277 predictor of PVT lapses ($\beta=-0.44$, $\text{sterr}=0.06$, $t=-6.87$, $p<0.001$), and KSS ($\beta=-0.20$, $\text{sterr}=0.03$, $t=-7.68$,
 278 $p<0.001$), such that every 5% reduction resulted in an increase of one lapse, or one point on the KSS.
 279 Tank% was also a significant predictor of %time in the Safe Zone ($\beta=0.75$, $\text{sterr}=0.22$, $t=3.40$,
 280 $p=0.001$), such that every 1% increase in the tank resulted in a 0.75% increase in time spent in the
 281 Safe Zone.

282
 283 On average, time series correlations between Tank% and performance and sleepiness were
 284 moderate (PVT lapses $r_{\text{Lag}0}=-0.50$, $\text{sterr}=0.08$, $R^2=0.25$; KSS $r_{\text{Lag}0}=-0.54$, $\text{sterr}=0.08$, $R^2=0.30$; %Time in
 285 Safe Zone $r_{\text{Lag}0}=0.45$, $\text{sterr}=0.17$, $R^2=0.20$).

286
 287



288

289

290 **Figure 3. "Sleep Tank", Performance and Sleepiness** – Time-of-day (24h clock) is on the x-axis, with % left in
 291 the sleep tank on the y-axis. Sleep opportunities are shown along the x-axis. BL=Baseline, N1-4=nightshifts, S1-
 292 S4=Sleep opportunities, RTDS=return to daytime schedule. PVT Lapses (upper), KSS (middle) and %Time in the
 293 Safe Zone from the driving simulator (lower) are superimposed over model output.

294

295

296

297 **4.0 Discussion**

298

299 Initial examination of the correspondence between model predictions and performance and
300 sleepiness measures from a four-night simulated nightshift protocol indicated relatively good
301 predictive value, with percentage left in the “Sleep Tank” significantly predicting performance
302 lapses, subjective sleepiness and safe driving during a 40-minute driving simulation. The model
303 explained an average of 20-30% of the variance across participants. Performance and sleepiness
304 were worst at the trials closest to model-indicated latest advisable bedtimes. Not only did this
305 simple model map onto the performance and sleepiness low points during the night shifts, but also
306 onto the recovery points after the final daytime sleep and return to daytime schedule.

307

308 As expected, during the study, performance and sleepiness were significantly impaired towards the
309 circadian low on each night shift (Dorrian, Lamond et al. 2000, Lamond, Dorrian et al. 2001).
310 Interestingly, there was no significant cumulative impairment across nights, which may be expected
311 across multiple nightshifts (Folkard and Tucker 2003). Sleepiness was worse on the first night in the
312 series, which likely reflects the extended wakefulness that often accompanies transition to night
313 shift (Tepas, Walsh et al. 1981). The relative stability in performance, and improvement in
314 sleepiness across consecutive nights is consistent with previous laboratory studies (Dorrian, Lamond
315 et al. 2003) and may reflect the ideal sleeping conditions in the laboratory, since sleep length and
316 quality are likely to be lower in sleeping environments where light, temperature, and noise
317 frequently cause sleep disturbance (Åkerstedt and Gillberg 1981b, Åkerstedt 1991).

318

319 Indeed, this study was a first-step proof-of-concept in a controlled laboratory environment, with
320 young, healthy males on basic performance tests. This clearly limits generalisability to the
321 workplace where employees include females, as well as people who are older, people who are
322 experiencing health complaints, such as gastrointestinal and cardiovascular disease, which are

323 higher amongst shiftworkers (Knutsson 1989, Costa 1997, Lowden, Moreno et al. 2010), and where
324 countermeasure use (e.g. caffeine) is common. The controlled light exposure during the study is
325 also different to the variable light exposure often experienced by nightshift workers, which often
326 includes a bright pulse of morning light during the commute home (Eastman, Stewart et al. 1994).
327 This is a particularly important reason for trialling this work in the field, since the “sleep tank” model
328 only picks up circadian changes implicitly via changes in sleep, and it is necessary to investigate how
329 model predictions map onto performance in the more chaotic workplace environment.

330

331 Therefore, critical next steps include examining the model with different shift schedules in the
332 laboratory and the field, and using polysomnographic sleep recording to estimate sleep period time
333 and sleep efficiency (as in the current study) as well as actigraphic estimates of these measures.
334 Results provide tentative evidence that this “sleep tank” model may be an informative tool to aid in
335 individual decision-making based on sleep history. Field validation will allow us to be ready to pair
336 the model with CSTs as they are validated against research-grade sleep measurement tools.

337

338 **4.1 Summary**

339 Increasingly, people are gaining access to information about their sleep. Using this information to
340 make evidence-based decisions relating to fatigue safety is not always straightforward. Arithmetic
341 transformation of sleep duration and quality into an intuitive “sleep tank” (SleepTankTM), which
342 includes suggestions such as the number of hours until more sleep is critical, may assist individual
343 deliberation about fitness for work at that moment, and across a coming shift. Further validation is
344 necessary, however initial findings are promising. Following validation of the model (and the
345 devices) “sleep tank” calculations could be added to consumer-grade actigraphs and/or sleep
346 monitoring apps to help people to map the performance and safety implications of their recent
347 sleep history. Such a device could display time left in the tank (hours), the latest advisable sleep time
348 and a sleep tank gauge, along with advice on how naps might refill the “tank”.

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