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## **Different underlying mechanisms for high and low arousal in probabilistic learning in humans**

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## 1 **ABSTRACT**

2 Humans are uniquely capable of adapting to highly changing environments by updating  
3 relevant information and adjusting ongoing behaviour accordingly. Here we show how this  
4 ability —termed cognitive flexibility— is differentially modulated by high and low arousal  
5 fluctuations. We implemented a probabilistic reversal learning paradigm in healthy participants  
6 as they transitioned towards sleep or physical extenuation. The results revealed, in line with  
7 our pre-registered hypotheses, that low arousal leads to diminished behavioural performance  
8 through increased decision volatility, while performance decline under high arousal was  
9 attributed to increased perseverative behaviour. These findings provide evidence for distinct  
10 patterns of maladaptive decision-making on each side of the arousal inverted u-shaped curve,  
11 differentially affecting participants' ability to generate stable evidence-based strategies, and  
12 introduces wake-sleep and physical exercise transitions as complementary experimental  
13 models for investigating neural and cognitive dynamics.

14  
15

## 16 **INTRODUCTION**

17 Making mistakes is inherent to learning and the accomplishment of any task. We make  
18 mistakes every day, even when faced with the same task repeatedly. Our ability to learn from  
19 these errors and flexibly adapt ongoing behaviour according to changes in the environment is  
20 critical for our survival. This ability —termed cognitive flexibility— depends on our innate  
21 capacity to establish associations between stimuli (S), responses (R), and outcomes (O), as well  
22 as to integrate previously acquired knowledge and skills into effective strategies for coping  
23 with similar future demands.<sup>1</sup> Here, we implement a Probabilistic Reversal Learning (PRL)  
24 task to study the modulatory effect of low and high arousal on cognitive flexibility —  
25 participants continue to perform as they fall asleep or with increasing physical exercise— to  
26 map either side of the Yerkes-Dodson Curve (1908).<sup>2</sup>

27 Cognitive flexibility is often studied using PRL tasks, typically assigning probabilistic  
28 reinforcement contingencies to abstract S-R associations, that are later abruptly reversed,  
29 requiring participants to learn new S-R reinforcement contingencies by trial and error to  
30 overcome prepotent ones<sup>3</sup>. Efficient performance relies on learning from the reinforcement  
31 received<sup>4</sup>, the estimation of the likelihood that a reversal may occur,<sup>5,6</sup> and the continuous  
32 integration of a history of choices and reinforcements.<sup>7</sup> Indeed, evidence from both human and  
33 animal studies suggests that different high- and low-order strategies or series of rules are  
34 adopted during reversal learning, leading to maladaptive response patterns when the external  
35 pressures change or when the internal milieu varies.<sup>7,8</sup> Parsing the microstructure of learning  
36 derived from trial-by-trial responses enables the dissociation of the cognitive processes and  
37 behavioural strategies that drive subjects' choices during reversal learning. Here we propose  
38 that arousal fluctuations may differentially modulate cognitive flexibility leading to distinct  
39 maladaptive behavioural patterns of performance.<sup>9</sup>

40 Fluctuations in arousal and alertness (hereafter described jointly as “arousal”) occur  
41 constantly across the day but are exacerbated during transitions toward strained states such as  
42 sleep<sup>10</sup> or physical extenuation,<sup>11</sup> where arousal levels change drastically in a progressive and  
43 nonlinear manner.<sup>12,13</sup> These arousal fluctuations play a crucial role in modulating cognition,

44 facilitating or hindering certain cognitive processes and performance to internal and external  
45 stimuli.<sup>14,15,16,17,18</sup>

46 The interaction between arousal and cognition has been traditionally approached from  
47 the perspective proposed by Yerkes and Dodson in 1908.<sup>2</sup> According to their famous inverted  
48 U-shaped law, the optimal level of cognitive performance in complex tasks is reached at  
49 moderate levels of arousal, whereas deviations from this optimal arousal point, below or  
50 beyond, result in cognitive performance impairments. Though reductionist, the inverted U-  
51 shaped law represents a useful minimal framework to characterize the neural and cognitive  
52 dynamics of many physiological states across the arousal spectrum. Among these physiological  
53 states, researchers have paid special attention to reduced arousal states, including sleep stages,<sup>19</sup>  
54 sedation,<sup>20</sup> sleep deprivation,<sup>21</sup> motivation<sup>22</sup> and fatigue.<sup>23</sup>

55 Sleep can be used as the gold standard model of transition toward low arousal.<sup>10</sup> This  
56 area looking at the interaction between homeostasis and cognitive function is understudied due  
57 to the complexity of capturing dynamically metastable states like mild sedation<sup>24,25</sup> and  
58 drowsiness.<sup>17</sup> When falling asleep, individuals manifest a wide range of changes, from  
59 physiological to phenomenological, that are categorized into several well-described sleep  
60 stages.<sup>26</sup> One of these stages is drowsiness, a transitional stage of consciousness between  
61 attentive wakefulness and light sleep, characterized by a progressive and nonlinear loss of  
62 responsiveness to external stimuli which does not immediately imply unconsciousness.<sup>27,28,29</sup>  
63 Drowsiness, as well as similar reduced arousal states, has been repeatedly associated with an  
64 impairment of cognitive processing, and particularly the capacity to deal with conflicting  
65 information,<sup>18</sup> attentional performance,<sup>30</sup> and perceptual decision-making.<sup>31</sup> However, in  
66 drowsiness, and even during highly reduced arousal states, pre-attentive and early bottom-up  
67 attentive processing can still be accomplished with and without conscious awareness.<sup>17,32,33</sup>

68 The transition towards the other side of the arousal spectrum (i.e., heightened arousal  
69 states) has received even less attention.<sup>34</sup> The absence of a theoretical model for progressive  
70 physiological transitions towards high arousal states, has also contributed to a lack of advance  
71 in the field. Here, we consider endurance physical exercise as a useful experimental model of  
72 arousal transition upwards, with many commonalities with sleep transition. A single bout of  
73 endurance physical exercise (e.g., running or cycling) up to physical extenuation involves a  
74 complex transition encompassing a wide range of changes (e.g., neural, motor, endocrinal,  
75 phenomenological, etc.), that are also categorized into several well-described stages, from  
76 resting, through the aerobic and the anaerobic thresholds, up to the limit where the individual  
77 has to stop.<sup>35</sup> This highly fluctuating transition has been also associated with changes in  
78 cognitive processing to internal and external stimuli.<sup>36,37,38</sup> In particular, high-order top-down  
79 processes that govern goal-directed behaviour in changing environments (i.e., cognitive  
80 control) appear to benefit from increases in the level of arousal<sup>39</sup> up to a certain exercise  
81 intensity. Further intensity increments approaching and exceeding the anaerobic threshold  
82 seem to hinder cognitive performance,<sup>36,37,38,40</sup> in line with the Yerkes-Dodson law prediction.

83 Sleep and physical exercise provide complementary perspectives on the cognitive  
84 dynamics, and experimental models, when the arousal level is altered. However, and despite  
85 the fact that both sides of the arousal spectrum exhibit similar cognitive performance  
86 impairments, they cannot be treated as mirroring states in terms of cognitive performance  
87 without a fine-grained differentiation of the behavioural dynamics that lead to these global

88 impairments. Furthermore, the theoretical differences in the transitions towards sleep or  
89 complete (physical) exhaustion have to be considered in the assumptions and interpretations of  
90 this and future studies. Thus, it is crucial to ask when arousal is altered (increased or decreased),  
91 which specific processes of cognitive flexibility and information processing are affected, and  
92 whether low and high arousal states are characterized by different strategic behaviours  
93 underlying decision-making. It should be understood that the physiological processes  
94 underlying the change in performance seen in different Dodson-Yerkes experiments since 1908  
95 are different at each side of the curve, and it should be expected that these changes in arousal  
96 modulate differently the cognitive abilities. Here, we use a PRL task to disentangle the  
97 behavioural dynamics of cognitive flexibility as they get modulated by ongoing fluctuations in  
98 arousal levels and to further delineate the microstructure of learning derived from trial-by-trial  
99 responses to conflicting evidence. In particular, we manipulated arousal level to facilitate  
100 natural transitions to low alertness, from awake to asleep; or to elicit high arousal, instructing  
101 participants to exercise during 60 minutes at the highest intensity and effort possible without  
102 reaching premature exhaustion. During both arousal modulations, participants performed a  
103 PRL task, requiring the adaptation of behaviour following changes in reinforcement and  
104 punishment, as well as the maintenance of strategic response patterns in the face of misleading  
105 (probabilistic) feedback.

106 Based on the premises that (1) drowsiness hinders the extraction of task-relevant  
107 information from external stimuli and its integration, fragmenting specific aspects of cognition  
108 while preserving crucial executive control processes;<sup>18,31,33,41</sup> (2) drowsiness has been  
109 associated with more liberal decision-making;<sup>17,30,31</sup> (3) moderate-to-high intensity endurance  
110 exercise leads to a selective enhancement of executive control processes while lower and higher  
111 intensities result in an impairment or minimal effect;<sup>40,42,43</sup> and (4) high arousal promotes  
112 habitual responding and reduced engagement of complex cognitive strategies;<sup>44,45,46</sup> predicted  
113 that behavioural performance would be enhanced in moderate-intensity physical exercise,  
114 while drowsiness and high-intensity exercise would lead to diminished performance in light of  
115 the inverted U-shaped Yerkes-Dodson Law. Specifically, we hypothesized that reduced arousal  
116 states would be associated with an impairment of performance (compared to baseline), which  
117 would be attributed to a tendency to apply a simple strategy (win-stay/lose-shift) instead of  
118 using an integrated history of choices and outcomes to drive performance (probabilistic  
119 switching behaviour). In contrast, while we also expected an impairment of performance during  
120 heightened arousal states, we hypothesized it would be attributed to a failure to disengage from  
121 ongoing behaviour (perseveration). In addition, we hypothesized that altered arousal states  
122 might reduce the ability of participants to apply a proper higher order strategy, resulting in wide  
123 periods of time-on-task in which participants would perform the task simply responding to the  
124 tones (i.e., automatic rule) but without applying any strategy (i.e., higher order rule). All these  
125 hypotheses, together with the analysis plan, were pre-registered after data collection.<sup>9</sup>

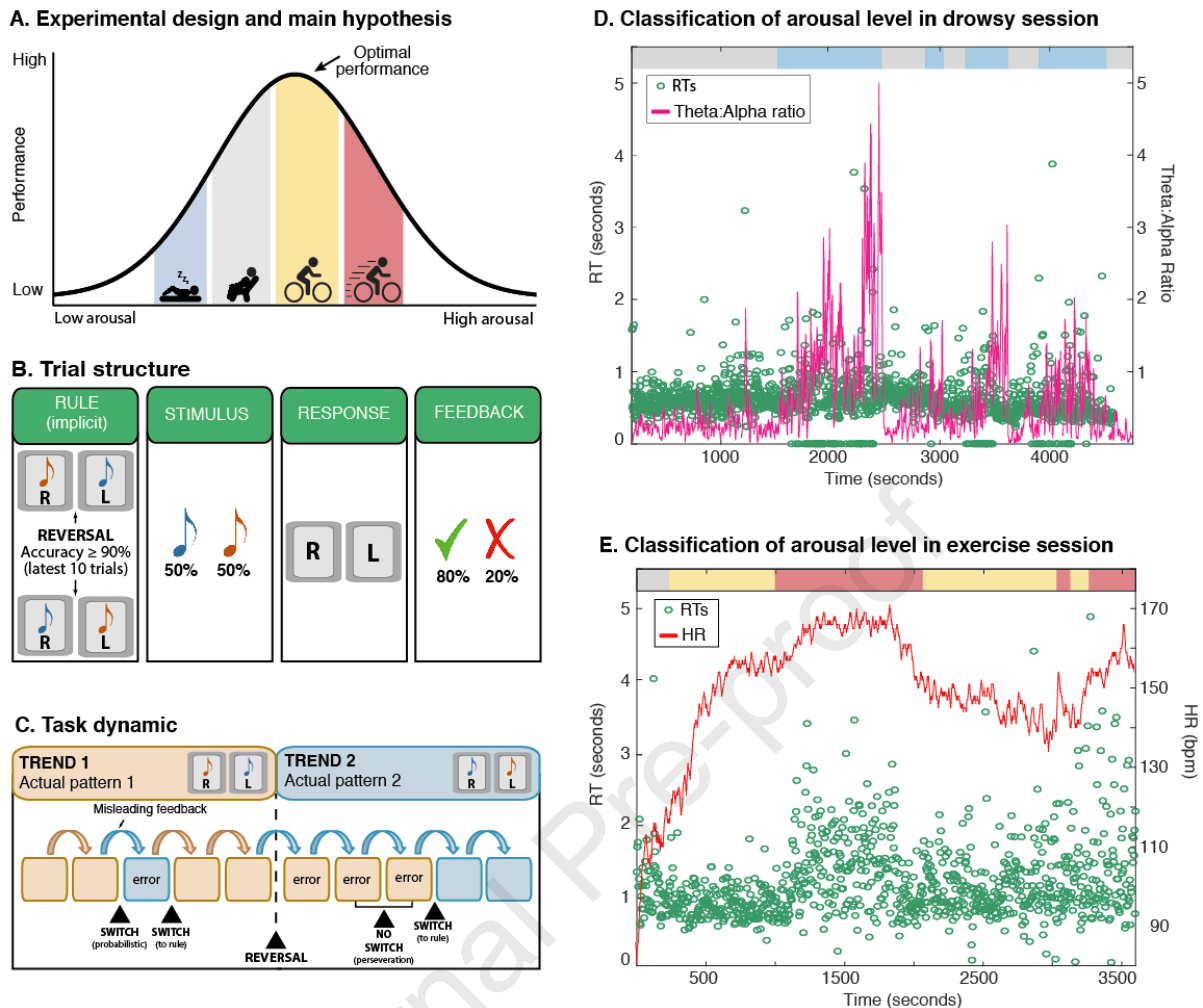
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## 127 **RESULTS**

128 To investigate the modulatory effect of arousal fluctuations on cognitive flexibility, a PRL task  
129 was carried out with human participants (n=100) while they were transitioning towards  
130 drowsiness or physical extenuation. Participants were instructed to associate an auditory  
131 stimulus (S) —high pitch sound or low pitch sound— with a response (R) button —left or right.

132 In this auditory version of the PRL task, each S-R association leads to an auditory outcome (O)  
133 —correct (ding sound) or incorrect (white noise)— which participants use to assess their  
134 choice, and apply this knowledge to guide the next choices. Indeed, participants were explicitly  
135 told that there was a rule connecting each of the auditory sounds to a corresponding button  
136 (e.g., the low pitch sound could correspond to the left button, and the high pitch sound to the  
137 right button or vice-versa), which they had to figure out based upon instructive feedback they  
138 would receive after each R. Additionally, they were instructed on two key issues: 1) the S-R  
139 rule might switch after a certain amount of time —becoming the opposite of what it was  
140 previously— and that no specific indication whether such a switch had occurred would be  
141 provided; 2) although the majority of the time the feedback would be truthful, sometimes it  
142 could be false and in essence mislead to them. Therefore, the task entails the use of, at least,  
143 two rules to success, as participants have to press a button after each auditory stimulus (i.e.,  
144 automatic rule) and to use an integrated history of S-R-O associations to determine the correct  
145 S-R association (i.e., high order rule). Once participants reach 90% accuracy or greater on the  
146 latest 10 trials, the implicit abstract S-R association is reversed, and participants have to infer  
147 the new association from the feedback received. The number of responses needed to attain a  
148 reversal (RAR) of the abstract association is used as the main index of performance. We  
149 hypothesized<sup>9</sup> that reduced arousal states would lead to reductions in behavioural performance  
150 compared to baseline arousal state; while heightened arousal states would lead to improved  
151 performance relative to baseline, but only to an optimal point (i.e., moderate arousal) after  
152 which the performance will be deteriorated with further increases in arousal level (see figure  
153 1A). These hypotheses were formulated in line with the famous psychology inverted u-shaped  
154 law originally attributed to Yerkes and Dodson (1908)<sup>2</sup> relating arousal modulation  
155 performance in complex tasks, but later more formally defined by Broadhurst (1958)<sup>47</sup> and  
156 Brown (1961).<sup>48</sup>

157 Note that, as a probabilistic task, the feedback provided is not always truthful nor  
158 reliable and misleads the participant 20% of the time (see figure 1B). Thus, the participant  
159 could correctly apply the S-R association and press the correct button in response to the  
160 auditory stimulus, and still receive negative feedback, thus indicating an incorrect choice. This  
161 scenario of conflicting evidence can lead participants to two different maladaptive response  
162 patterns (see figure 1C) while performing the task: 1) switching the pattern choice across trials  
163 with little (i.e., one negative feedback against the choice) or no evidence (i.e., no feedback  
164 against the choice) of an actual rule change (probabilistic switching); or 2) sticking with the  
165 previous choice despite having strong evidence (i.e., two or more negative feedbacks against  
166 the choice) of an actual rule change (perseveration). Relying on these response patterns lead to  
167 poor performance,<sup>7</sup> as the optimal strategy in this task is to stick with the previous choice with  
168 zero or one negative feedback against the choice, and to switch the pattern choice if two or  
169 more consecutive negative feedbacks against the choice happen.



170  
 171 **Figure 1. Experimental design and arousal level classification:** A) Schematic representation of the  
 172 experimental design and main hypotheses. Arousal level was endogenously manipulated by facilitating the natural  
 173 transition of participants from awake to sleep, or instructing them to exercise during 60' at the highest intensity  
 174 and effort they could maintain without reaching premature exhaustion. Notice that half of the participant  
 175 transitioned towards drowsiness, while the other half transitioned towards physical exertion. A probabilistic  
 176 reversal learning task was assessed continuously during the arousal modulation. Optimal performance of the task  
 177 was expected at moderate arousal state (exercising at moderate intensity), while lower (drowsiness) and higher  
 178 (exercising at high-intensity) arousal state were expected to result in task performance deterioration. B) In this  
 179 auditory version of the probabilistic reversal learning paradigm, an auditory stimulus was presented on each trial,  
 180 and participants had to associate the sound with a response button, left or right. After that, auditory feedback was  
 181 provided according to the ongoing implicit rule. Notice that the feedback provided was not always truthful nor  
 182 reliable, and attempted to mislead the participant 20% of the time. C) Task trials were grouped into sequences of  
 183 trials following a particular rule (trend) where a particular sound was implicitly associated with a response button  
 184 (e.g., high pitch sound with the left button, and low pitch sound with the right button). Participants were instructed  
 185 to infer the rule from the provided feedback to assess their previous choice and apply the knowledge of their  
 186 accuracy to guide the next choices, knowing that the rule might change after a certain time. Based on the feedback  
 187 received, participants could make probabilistic or perseverative errors in the following trials. D) Automatic  
 188 classification of arousal during a drowsy session (representative participant). The pink line depicts changes in the  
 189 theta:alpha ratio (occipital electrodes cluster) during the pre-trial period (2 seconds before the auditory stimulus  
 190 onset). The horizontal bars on top represent trials classified as baseline (grey) or low arousal (blue). The variability  
 191 in the reaction times (green circles) closely follows the changes in theta:alpha ratio. Notice that circles on the  
 192 horizontal axis (reaction time equal to zero) were non-responsive trials, usually during low arousal (drowsy)  
 193 periods but also observed during exercise periods. E) Automatic classification of arousal during a physical  
 194 exercise session (representative participant). The red line depicts changes in the heart rate during the pre-trial

195 period (2 seconds before sound onset), and the horizontal bars on top represent trials classified as baseline (grey),  
196 moderate (yellow) or high arousal (red). Similar to the low arousal session, the reaction times (green circles)  
197 fluctuates with the changes in heart rate.

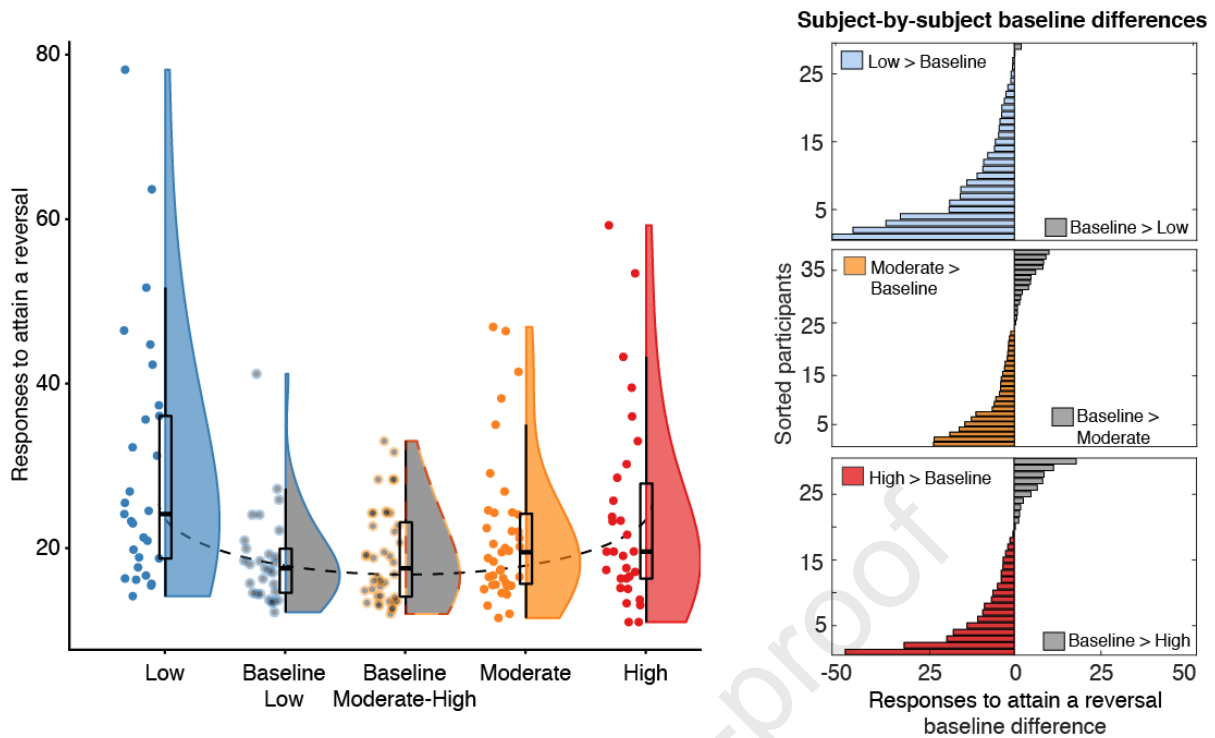
198

### 199 **Arousal modulates probabilistic information during a stream of conflicting evidence.**

200 First, we calculate the average RAR per participant in each arousal state (low, baseline sitting,  
201 baseline cycling, moderate, high). To account for the dependencies potentially generated by  
202 any procedural differences between Experiments, we fitted RAR using hierarchical linear  
203 mixed-effects modelling, with arousal as fixed effect, and participant nested into Experiment  
204 as random effects. The model showed a strong effect of arousal on RAR,  $F(3, 113.02) = 11.59$ ,  
205  $p < 0.001$ ,  $\beta = 0.61$  (details on testing model assumptions can be found in the supplementary  
206 material), indicating that the processing of probabilistic information that allows the detection  
207 of changing patterns in a stream of conflicting evidence was modulated by the arousal level.  
208 Next, we checked for non-linearity in the relationship between arousal and RAR, to test the  
209 famous u-shaped curve. As expected, we found that the quadratic ( $AIC = 1243.6$ ;  $BIC = 1262.3$ ;  
210  $R^2 = 0.40$ ) outperformed linear fitting ( $AIC = 1264.8$ ;  $BIC = 1280.4$ ;  $R^2 = 0.23$ ), confirming a  
211 possible curvilinear pattern (U shaped) of the effect of arousal on RAR (see figure 2), with a  
212 reliable increase in the number of responses required by the participants to complete a trend  
213 reversal (i.e., decrease of performance) as the level of arousal progress towards the extremes  
214 of the defined arousal range, confirming, for reversal learning, convergence with the Yerkes-  
215 Dodson law, later reformulated by Broadhurst in 1958.<sup>47</sup>

216 Splitting the comparisons to its specific baselines per arousal condition (i.e., sitting  
217 baseline compared to low arousal in the drowsiness condition; cycling baseline compared to  
218 moderate and high arousal in the exercise condition) yielded a reliable increase of RAR in low  
219 arousal,  $t(124.62) = 5.67$ ,  $p < 0.001$ ,  $\beta = 1.02$ , and high arousal state,  $t(117.93) = 2.57$ ,  $p =$   
220  $0.011$ ,  $\beta = 0.45$ , compared with their corresponding baselines. Notably, baseline performance  
221 did not differ across arousal conditions (see supplementary figure 1). Contrary to what we  
222 expected, moderate arousal state was not associated with a decrease of RAR (the expected peak  
223 in performance), relative to baseline ( $t(114.85) = 1.61$ ,  $p = 0.11$ ,  $\beta = 0.25$ ). Moreover, we did  
224 not find evidence for a potential dual-task confounding effect in the heightened arousal  
225 conditions (see supplementary material). In sum, these findings provide evidence for an  
226 impairment in the processing of probabilistic information when the arousal level is altered,  
227 regardless of the side of the arousal spectrum.





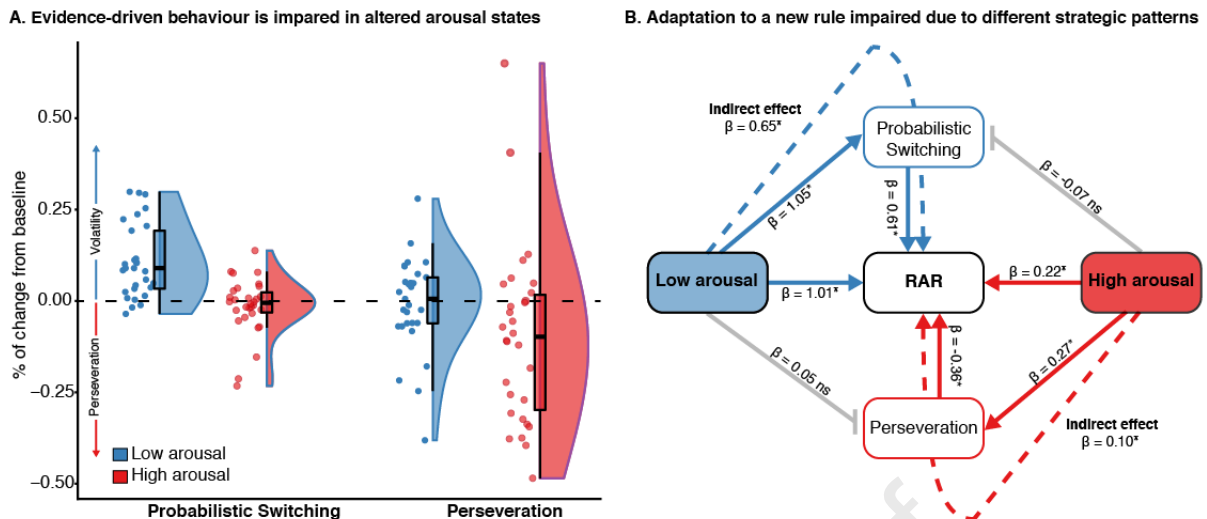
228  
 229 **Figure 2. Number of responses needed to attain a trend reversal as a function of the arousal state.** A) Violins  
 230 and overlaid box plots of mean responses to reverse across arousal states. In box plots, middle black mark indicates  
 231 the median, and bottom and top edges indicate 25th and 75th percentiles, respectively. The upper and lower  
 232 whiskers indicate the maximum value of the variable located within a distance of 1.5 times the interquartile range  
 233 above the 75th percentile and below the corresponding distance to the 25th percentile value. Surrounding the  
 234 boxes (shaded area) is a rotated kernel density plot, which is comparable to a histogram with infinitely small bin  
 235 sizes. Jittered dots represent the averaged response to reverse score for each participant in each arousal state.  
 236 Linear mixed-effects model analysis revealed a reliable quadratic fitting between arousal and task performance,  
 237 outlined by the dashed line. Low and high arousal states were associated with a worse task performance relative  
 238 to their own baseline arousal states. Moderate arousal state was not associated with the expected optimal  
 239 performance as no differences were found with the baseline arousal state. B) Baseline differences of each  
 240 participant across altered arousal states are represented by the bars (grey bars indicate that these participants  
 241 needed more trials to attain a trend reversal in the baseline compared with the altered arousal states; blue, yellow  
 242 and red bars depict that these participants needed more trials to attain a trend reversal when arousal level was  
 243 altered -increased or decreased- compared with baseline arousal state). Participants are sorted by performance  
 244 difference between baseline and the arousal state. Upper and bottom panels show a consistent impairment of task  
 245 performance across participants in low and high arousal states. Non-reliable differences were found between  
 246 moderate and baseline arousal.

247  
 248 **Different underlying mechanisms explain decreased performance in low and high arousal**  
 249 **states**

250 In the analysis above, performance under high and low arousal states was compared  
 251 irrespective of the strategy participants may have used to solve the task. To test for the  
 252 hypotheses of the differential mechanism driving changes in performance for each arousal side  
 253 of the u-shaped curve, we calculated: a) probabilistic switching, as the proportion of trials when  
 254 the participants change the pattern choice with little or no evidence (i.e., zero or one negative  
 255 feedback against the choice); and b) perseveration, the likelihood of sticking with the previous  
 256 choice despite strong evidence (i.e., receiving two or more negative feedbacks in a row) that  
 257 the pattern has changed. Probabilistic switching and perseveration are proportion indices of

258 strategic behaviour based on the probability of switching when negative feedback is provided.  
259 Thus, they range between 0 and 1, allowing comparison across arousal states while accounting  
260 for potential experimental differences (e.g., number of trials). We hypothesized that the  
261 impairment of performance in low arousal would be primarily attributed to an increase in  
262 probabilistic switching, relative to the baseline arousal state; and in contrast, the observed  
263 impairment of performance in high arousal state will be primarily due to an increase in  
264 perseverative behaviour. To test these hypotheses, we fitted probabilistic switching and  
265 perseveration (separately for low and high arousal states) using the hierarchical linear mixed-  
266 effects model structure defined previously. The analyses revealed that, while the probabilistic  
267 switching increased consistently across subjects during low arousal state compared with  
268 baseline arousal,  $F(1,56) = 14.78$ ,  $p < 0.001$ ,  $\beta = 1.01$ ,  $R^2 = 0.21$ , no reliable differences were  
269 observed in perseveration between these arousal states ( $F < 1$ ). On the other hand, high arousal  
270 states led to a reliable increase in perseverative behaviour compared to the baseline state,  $F$   
271  $(1,67) = 9.12$ ,  $p = 0.035$ ,  $\beta = 0.34$ ,  $R^2 = 0.12$ , with no reliable differences observed in  
272 probabilistic switching ( $F < 1$ ). These results suggest that altered arousal states lead to distinct  
273 maladaptive decision-making patterns that affect participants' ability to generate stable  
274 evidence-based strategies, although evidence-driven responses were present (see figure 3A).

275 To further prove that the impairment in performance in low and high arousal states  
276 could be attributed to the different maladaptive behavioural patterns, we carried on a mediation  
277 analysis separately for each arousal state (low, high). We first confirmed that probabilistic  
278 switching and perseveration have an effect on the RAR, while controlling for the arousal state  
279 (see figure 3B). These results, together with the previous analyses where we found an effect of  
280 arousal state on probabilistic switching and perseveration, revealed a full mediation between  
281 these variables. As figure 3B illustrates, the regression coefficient between arousal and RAR,  
282 and the regression coefficient between probabilistic switching and RAR were statistically  
283 reliable, showing a full mediation of probabilistic switching on the effect of low arousal on  
284 RAR. The bootstrapped standardized indirect effect of low arousal on RAR, mediated by  
285 probabilistic switching, was 0.65 ( $p < 0.001$ ), and the 95% confidence interval ranged from  
286 0.29 to 1.07. A similar fully mediation effect was observed in high arousal state, showing that  
287 the effect of high arousal on behavioural performance was fully mediated via the perseverative  
288 behaviour. The bootstrapped standardized indirect effect was 0.10 ( $p = 0.014$ ), and the 95%  
289 confidence interval ranged from 0.14 to 0.24. As predicted, participants showed an impairment  
290 of performance during low arousal state, relative to baseline arousal, which was primarily  
291 attributed to an increase of probabilistic switching (i.e., changing pattern choice with little or  
292 no evidence of an actual rule change). In contrast, while participants also showed an  
293 impairment of performance during high arousal state, relative to the baseline arousal, it was not  
294 attributed to an increase in probabilistic switching, but to an increase in perseverative behaviour  
295 (i.e., sticking with the previous choice despite consecutive negative feedbacks).  
296



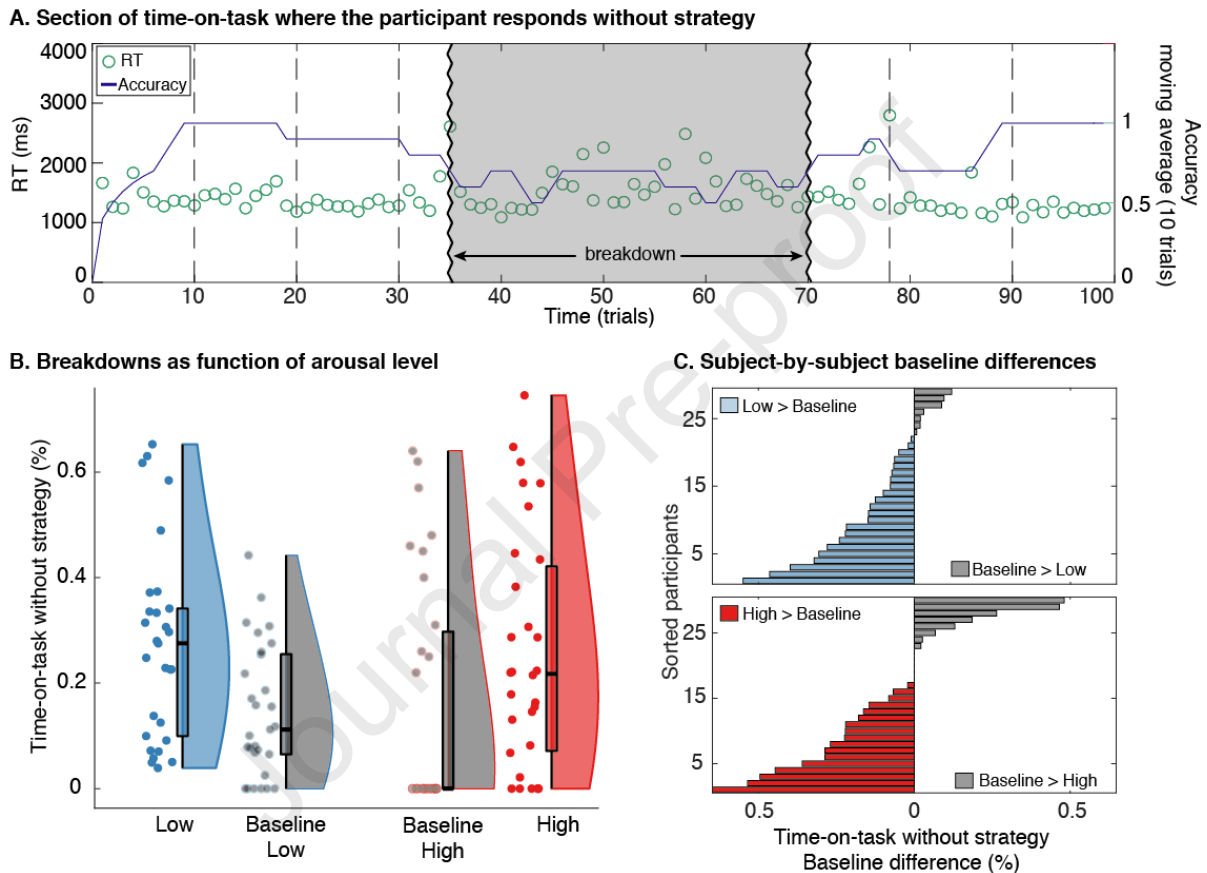
297  
 298 **Figure 3. Maladaptive behavioural patterns across participants in low and high arousal states.** A) Violins  
 299 and overlaid box plots of the percentage of change from baseline to low (blue) and high (red) arousal states in  
 300 probabilistic switching and perseveration. In box plots, middle black mark indicates the median, and bottom and  
 301 top edges indicate 25th and 75th percentiles, respectively. The upper and lower whiskers indicate the maximum  
 302 value of the variable located within a distance of 1.5 times the interquartile range above the 75th percentile and  
 303 below the corresponding distance to the 25th percentile value. Surrounding the boxes (shaded area) is a rotated  
 304 kernel density plot, which is comparable to a histogram with infinitely small bin sizes. Jittered dots represent the  
 305 averaged response to reverse score for each participant in each arousal state. B) Mediation model diagram to  
 306 illustrate that the general impairment in task performance found in low and high arousal states was mediated by  
 307 different maladaptive behavioural patterns. Dashed lines (indirect effects) represent the effect of low (blue) and  
 308 high (red) arousal on task performance (indexed by the averaged responses to attain a trend reversal) through  
 309 probabilistic switching and perseveration, respectively. Solid lines depict direct effects between variables. Grey  
 310 lines represent the absence of a direct effect of low arousal on perseveration and high arousal on probabilistic  
 311 switching. Notice that a direct effect of an independent variable (arousal) onto the mediator (probabilistic  
 312 switching, perseveration) is a prerequisite for mediation being possible. Standardized  $\beta$  regression coefficients  
 313 are indicated in each effect (\* depicts  $p < 0.05$ ). Accordingly, the values of all effects are expressed as the number of  
 314 standard deviations from the mean. For example, the direct effect of high arousal on RAR ( $\beta = 0.22$ ) implies that  
 315 a standard deviation change of 1 in the arousal variable would result in a standard deviation increase of 0.22 in  
 316 RAR.

317

### 318 **Arousal disrupts the reversal strategy**

319 To maximise performance in the task, a good strategy is to not fall for the false feedback and  
 320 stand your ground until the next feedback, as well as switch to the second consecutive feedback.  
 321 The fact that participants sometimes needed an unreasonable high number of responses to attain  
 322 a reversal in low and high arousal states suggests the existence of sections of time on task in  
 323 which they responded to the tones but could not apply the strategy rules (see fig 4A). These  
 324 sections without clear strategic behaviour, that we call breakdowns, have been often neglected  
 325 in previous studies using PRL tasks as failures of compliances or “bad participant”. The  
 326 transient on/off nature of these breakdowns may provide valuable insight into the behavioural  
 327 dynamics of participants in different states of arousal. We hypothesized that breakdowns  
 328 sections would increase in low and high arousal states, relative to a baseline arousal state. First,  
 329 we traced the sections of the task (more than 20 trials) in which participants did not attain a  
 330 reversal. Second, we calculated the proportion of time these sections represented to the total  
 331 time-on-task, and finally, we implemented a hierarchical linear mixed-effects model with the

332 structure defined in previous analyses, separately for each arousal state (low, high), with the  
 333 number of breakdowns as the index of performance. As hypothesized, low and high arousal  
 334 states lead to longer breakdown sections compared with baseline arousal state ( $t(127.99) =$   
 335  $3.40, p < 0.001, \beta = 0.13$ ;  $t(121.69) = -2.97, p = 0.003, \beta = 0.11$ ). Subject-by-subject results  
 336 (fig 4C) show a consistent increase of breakdowns across participants in low arousal state.  
 337 Although high arousal states also showed a reliable increase of breakdowns as a group, this  
 338 effect was less systemic, with half of the participants showing the opposite effect, no difference  
 339 or no breakdowns.  
 340



341  
 342 **Figure 4. Behavioural strategy breaks as arousal changes.** A) Automatic classification of a section of time  
 343 where a representative participant responded without a clear behavioural strategy. The green circles show RTs  
 344 and the blue line shows the ongoing accuracy of the task (10-points moving average). The grey shaded area flanked  
 345 by the zigzagging vertical lines depicts the section of time classified as a breakdown. B) Violins and overlaid box  
 346 plots of the averaged percentage of time-on-task without strategy across participants in low and high arousal states,  
 347 compared with their respective baseline states. In box plots, the middle black mark indicates the median, and  
 348 bottom and top edges indicate 25th and 75th percentiles, respectively. The upper and lower whiskers indicate  
 349 the maximum value of the variable located within a distance of 1.5 times the interquartile range above the  
 350 75th percentile and below the corresponding distance to the 25th percentile. Surrounding the boxes (shaded area)  
 351 is a rotated kernel density plot, which is comparable to a histogram with infinitely small bin sizes. Jittered dots  
 352 represent the averaged percentage of time-on-task without a strategy of each participant in each arousal state.  
 353 Linear mixed-effects model analyses revealed that low and high arousal states lead to longer periods of breakdown  
 354 relative to the baseline arousal state. Interestingly, violin plots show a considerable number of participants who  
 355 had no breakdowns at baseline arousal states, something that completely disappears in low arousal state (all  
 356 participants had breakdowns), and that is reduced in high arousal state. C) Baseline differences of each participant  
 357 in low and high arousal states represented by horizontal bars (grey bars indicate that these participants spent more  
 358 time performing the task without a particular strategy in the baseline arousal state compared with the altered

359 arousal states; blue and red bars depict that these participants were applying behavioural strategies less time when  
360 arousal level was altered (increased or decreased) than in baseline arousal state. Participants are sorted by  
361 performance difference between baseline and the arousal state. Both panels show a consistent impairment of task  
362 performance across participants in low and high arousal states.

363

## 364 **DISCUSSION**

365 In the present study, we facilitated natural transition of healthy participants towards the borders  
366 of non-pharmacological arousal states (drowsiness, physical exertion) to investigate the  
367 behavioural dynamics of cognitive flexibility. In line with our pre-registered hypotheses,<sup>9</sup> the  
368 findings revealed a quadratic-like pattern (inverted U-shape) of the effect of arousal  
369 fluctuations on cognitive performance. As the level of arousal progressed towards the extremes  
370 of the defined arousal range reversal learning performance decreased, in agreement with the  
371 predictions of the Yerkes-Dodson law (1908).<sup>2</sup> Although cognitive flexibility diminished in  
372 both under high and low arousal states, different maladaptive behavioural patterns drove this  
373 performance impairment. As predicted, the performance decline exhibited by our participants  
374 under drowsy states was primarily attributed to a more decision volatility (i.e., shifting pattern  
375 choice with little or no evidence of reinforcement contingencies change). In contrast,  
376 participants also showed a decline in performance during high arousal state but attributed to  
377 increased perseverative behaviour (i.e., sticking with a particular pattern choice despite having  
378 strong evidence that the contingencies have changed). Our findings also revealed that most  
379 participants undergo prolonged periods of time-on-task in which they seem unable to apply any  
380 specific higher order strategy. These breakdown periods, which can last for several minutes,  
381 are more frequent and sustained during high or low arousal. In short, our results provide solid  
382 evidence for distinct maladaptive decision-making patterns under altered arousal states,  
383 differentially affecting the participants' ability to generate stable evidence-based strategies.

384 Arousal fluctuations thus seem to elicit a distinctive behavioural distortion of cognitive  
385 flexibility as further indicated by the microstructure of learning derived from trial-by-trial  
386 responses to negative feedback. Healthy participants under high arousal exhibited normal  
387 acquisition of S-R reinforcement contingencies but perseverative response patterns when  
388 contingencies were reversed. This failure to disengage from ongoing behaviour is a  
389 translational phenomenon strongly linked to impulsivity and compulsivity,<sup>49</sup> and prevalent in  
390 numerous neuropsychiatric and medical conditions.<sup>7,50,51</sup> For instance, patients with lesions that  
391 include ventral prefrontal cortex and orbitofrontal cortex,<sup>52</sup> as well as chronic cocaine users<sup>53</sup>  
392 and patients with schizophrenia,<sup>54</sup> show normal acquisition of S-R contingencies but are  
393 severely impaired when those S-R reinforcement contingencies are abruptly reversed,  
394 exhibiting perseverative responding to the previously reinforced S-R contingency. Altogether,  
395 these findings suggest that high arousal undermines healthy individuals' capacity to engage in  
396 complex cognitive strategies driving them to rely on habitual response patterns, which,  
397 paradoxically, might also enhance behavioural control in terms of response inhibition.<sup>46</sup> Our  
398 findings not only further the understanding of the processes underlying automatized behaviour  
399 and habitual response tendencies, but high arousal may be used as a model to inform both  
400 impulsive and compulsive aspects of psychopathology.

401 In contrast, healthy participants under low arousal seemed unable to maintain the  
402 learned S-R reinforcement contingency and started to deviate from the evidence, revealing a

403 volatile pattern of behaviour. Since a crucial aspect of the PRL experimental design was the  
404 existence of a 20% of misleading feedback, to maximise performance, individuals should not  
405 fall for the false feedback and —ideally— stand their ground until the next feedback. Further  
406 and as part of a successful strategy, they should switch if two or more consecutive feedbacks  
407 are given against the previously reinforced choice pattern. Consequently, adaptive behaviour  
408 during the task requires a balance between both types of behaviour (stability and flexibility).  
409 Those participants under low arousal fell repeatedly for the misleading feedback, switching  
410 prematurely after negative feedback. Furthermore, they showed increased decision volatility  
411 by spontaneously switching even without any negative feedback. This volatile pattern of  
412 cognitive flexibility has been linked to serotonin<sup>55</sup> and dopamine systems,<sup>56</sup> and is observed in  
413 patients with major depression,<sup>57,58,59</sup> often linked to either an oversensitivity to punishment or  
414 an impaired control over negative feedback.<sup>60,61</sup> It is reasonable to speculate that low arousal  
415 levels render individuals more sensitive in updating S-R reinforcement contingencies, rather  
416 than increase sensitivity to punishment as in major depression. Moreover, low arousal may  
417 increase volatility by decreasing attentional resources, leading to spontaneous explorations,  
418 higher RT variability and periodic omissions (see supplementary figure 2).

419 The fragmentation of cognitive control due to changes in arousal has been primarily  
420 shown in sleep deprivation<sup>62,63,64,65,66</sup> and not in spontaneous fluctuations of alertness as we  
421 show in this study. The increased volatility in the PRL with low arousal suggests a decrease in  
422 cognitive control that is different from an increase in perseverative behaviour seen in high  
423 arousal. Indeed, we have previously shown that decreased levels of arousal can fragment or  
424 reconfigure specific aspects of cognition while preserving crucial executive control processes  
425 such as the capacity to detect and react to incongruity,<sup>18</sup> the efficiency in perceptual decision  
426 making,<sup>31</sup> and the precision of conscious access.<sup>17</sup> Here, we add further evidence showing that  
427 individuals under reduced arousal state, although struggling to maintain stable evidence-based  
428 decision-making patterns, are able to learn new S-R reinforcement contingencies,  
429 demonstrating flexibility of the human brain to adapt to increasing levels of endogenous  
430 (arousal) noise. The evidence of cognitive and —indirectly— neural reconfiguration of  
431 cognitive control networks suggests compensatory mechanisms elicited by the change in  
432 arousal.

433 Upon further examining the microstructure of learning derived from trial-by-trial  
434 performance of the PRL task, we uncovered the existence of prolonged periods of time-on-task  
435 in which participants did not seem to apply any particular high-order behavioural strategy.  
436 Although these breakdown periods emerged regardless of the arousal level, they were prevalent  
437 under low and high arousal states, lasting from few to several minutes. Remarkably, the  
438 transient on/off nature of these breakdowns suggests that extreme arousal levels alternate  
439 between different metastable cognitive states. The first state can be defined by a relatively  
440 successful application of the reinforcement information where participants can navigate the  
441 uncertainty of the PRL, while in the other metastable state they seem to only apply the simple  
442 auditory-motor S-R rule to respond to the auditory tones but are unable to use choice history  
443 to develop a successful strategy.

444 In the context of this study, arousal as a biological construct defined by the homeostatic  
445 regulatory capacity of the system and its responsiveness,<sup>67</sup> helps to link drowsiness and  
446 increased alertness during physical exercise in a common framework where the predictions of

447 the Yerkes-Dodson inverted U-shaped law can be experimentally tested. Despite the obvious  
448 difference at the biological, neural and psychological level between both sides of the curve, the  
449 common decrease in performance highlights the commonalities between the extremes in human  
450 performance, adding to the fact that both —sleep and physical exertion— emerge as natural  
451 transitions from a similar state (resting) traversing different stages, and exhibit nonlinear  
452 dynamics and hysteresis processes in their transitions.<sup>12</sup> Thus, drowsiness and physical exertion  
453 provide complementary perspectives on cognitive dynamics when the arousal level is altered.  
454 The present findings point out their differences in the cognitive fragmentation leading to a  
455 general decline in task performance.

456 Transitions towards drowsiness or physical exertion entail changes in levels of arousal,  
457 which are in turn associated with a wide range of alterations (e.g., neural, motor, endocrinal,  
458 phenomenological, etc.) that might cause the cognitive fragmentation described in the present  
459 study. For instance, during a single bout of aerobic exercise, as intensity increases from low to  
460 high, there is a release of epinephrine and, to lesser extent norepinephrine, into the blood from  
461 the adrenal medulla.<sup>15</sup> This exercise-induced increase in brain concentrations of  
462 catecholamines has been proposed as a physiological mechanism underlying cognitive  
463 performance during and after physical exercise.<sup>15</sup> Similarly, when falling asleep, we experience  
464 a cascade of changes in almost every system of the organism, including the somatic and  
465 autonomic nervous systems,<sup>12</sup> which might be playing a crucial role in cognitive processing.  
466 The extent to which each of the changes that occur during these transitions (drowsiness and  
467 physical exertion) are responsible for the cognitive adaptations we report here is something  
468 that future studies might reveal, for example, combining measurements of the autonomic  
469 nervous system and brain functioning, which would make it possible to gain more insight into  
470 the underlying physiological mechanisms involved in arousal-related changes in cognition.  
471 These inferences of this study are hence mediated by physiological processes that might  
472 partially explain the cognitive modulations in an independent manner if dissociated from  
473 arousal changes.

474 Though the Yerkes-Dodson law was not initially formulated to be a general rule to  
475 apply to all psychology subfields (learning, motivation, emotion, etc.), through the years, and  
476 with the pressure to find common mechanisms in psychology, the findings initially defined for  
477 learning were further extended and reinterpreted as a law about the relationship between  
478 arousal and other physiological constructs to perceptual and cognitive performance.<sup>68</sup> Despite  
479 this overgeneralization from its genuine formulation and its reductionist nature, our findings  
480 rely on such inverted U-shaped law as a basic useful theoretical framework, providing an  
481 attractive theoretical model to characterize the neural, cognitive and behavioural dynamics  
482 involved in the impact of arousal fluctuations in a wide range of physiological states and  
483 neuropsychiatric conditions.

484 Our findings bring some generalizations about the need to extend the traditional  
485 framework of understanding the interplay between cognitive dynamics and arousal through the  
486 prism of the homeostatic steady-state dynamics using pharmacological interventions<sup>34</sup> or  
487 transient alterations of emotional state.<sup>69</sup> In addition to this classical approach, we believe that  
488 drowsiness and physical exertion provide fruitful —naturally occurring— alterations of the  
489 arousal level with a preserved capacity to behaviourally respond, which can be utilized to study  
490 the modulation of neural function and cognitive processing. In the traditional steady-state

491 approach, such natural fluctuations of the arousal level may be undetected,<sup>70</sup> hindering or  
492 distorting cognitive and neural markers of crucial aspects of information processing.<sup>17</sup>  
493 Pharmacological and lesion perturbations of the brain are regarded as causal in cognitive  
494 neuroscience and regarded as stronger in their explanatory power than conditions relying on  
495 stimuli or psychological modulations. Arousal is an internally modulated change that can be  
496 used to study cognition and may be regarded in the strong causality range due to its partial  
497 independence from psychological processes.<sup>18</sup> The cases of drowsiness and physical exertion  
498 as causal models to study the neural mechanism of cognitive flexibility may prove to be very  
499 useful in the exploration of how cognition is fragmented or remain resilient under (reversible)  
500 perturbations of arousal<sup>17,33,71</sup> Our findings highlight that further research should focus on the  
501 rapidly changing dynamics of brain function and cognitive processing that appear to capture  
502 key dynamics relevant to our behavioural and perhaps even phenomenological experience, as  
503 we drift into strained physiological states.

504

## 505 **MATERIALS AND METHODS**

### 506 **Participants**

507 A total sample of 100 participants of an age range between 18 and 40 years old was included  
508 in the present study. All participants reported normal binaural hearing, no visual impairment  
509 and no history of cardiovascular, neurological or psychiatric disease. They were asked to get a  
510 normal night rest on the day previous to testing, and not to consume stimulants like coffee or  
511 tea on the day of the experiment. We report how we determined our sample size, all data  
512 exclusions, all inclusion/exclusion criteria, whether inclusion/exclusion criteria were  
513 established prior to data analysis, all manipulations, and all measures in the study.

514 The first experiment (herein Experiment#1) consisted of 35 participants (15 female; age  
515 range 18-40). In addition to the general aforementioned inclusion criteria, only easy sleepers,  
516 as assessed by the Epworth Sleepiness Scale (ESS),<sup>72</sup> were selected to increase the probability  
517 that participants fell asleep. Recruited participants were considered healthy with relatively high  
518 ESS scores but not corresponding to a condition of pathological sleep such as hypersomnia  
519 (i.e., scores 7–14). They were recruited via the Cambridge Psychology SONA system. Note  
520 that the target sample size was 50 participants transitioning towards drowsiness. However, after  
521 collecting the first 35 we decided to make slight modifications to the experimental protocol by  
522 increasing the time of the drowsy blocks to obtain a higher proportion of trials in low arousal.  
523 For this reason, we decided to collect a second sample (Experiment#2) which consisted of 15  
524 participants (11 female; age range 18-40), where we included these key modifications to the  
525 experimental protocol (see Procedure section for more details). Inclusion criteria and  
526 recruitment processes were similar to Experiment#1.

527 The third experiment (herein Experiment#3) consisted of 50 participants (6 female; age  
528 range 19-39). Additionally to the common inclusion criteria, only individuals who reported at  
529 least 8 hours of cycling or triathlon per week were selected. Well-trained cyclists were selected  
530 because they are used to maintaining the pedalling cadence at high intensity during long periods  
531 of time. Furthermore, they are able to keep a fixed posture over time, which notably reduces  
532 movement artefacts. They were recruited from the University of Granada (Spain) through  
533 announcements on billboards and previous databases.



534 All participants from the three experiments gave written informed consent to participate  
535 in the study and received a remuneration of 10€ per hour (i.e., approximately 30€ per  
536 participant). The Cambridge Psychology Ethics Committee and the University of Granada  
537 Ethics Committee approved the study (CPREC 2014.25; 287/CEIH/2017).

538

### 539 **Experimental task**

540 A modified version of the probabilistic reversal learning paradigm was used in all three  
541 experiments, which was characterized by employing auditory stimuli and an abstract rule (see  
542 figure 1B-C). In this task, participants learnt to choose one of two randomly presented tones  
543 by receiving instructive auditory feedback tones after each response, indicating either a correct  
544 or incorrect choice. When participants reached a 90% accuracy in the last 10 trials,  
545 reinforcement/punishment contingencies were reversed so that the previously reinforced tone  
546 was punished and vice versa. Within each reversal trend, a 20% probabilistic error trial was  
547 included in which “wrong” feedback was given for correct choices, even though the  
548 reinforcement contingencies had not changed. Participants were instructed to infer the rule  
549 from the feedback received, knowing that sometimes it might be misleading and that the rule  
550 might change after a certain time (see supplementary material for more details on the task  
551 instructions). The stimuli were binaurally presented at a random time interval (between 1000  
552 and 1500 ms) during 500 ms. They had to respond to both targets by pressing a button with  
553 their right or left hand.

554

### 555 **Procedure**

556 In Experiment#1, participants were fitted with an EGI electrolyte 129-channel cap (Electrical  
557 Geodesics, Inc. systems) after receiving the task instructions and subsequently signing the  
558 informed consent. The whole session was completed in a comfortable adjustable chair with  
559 closed eyes. Task instructions were to respond as fast and accurately as possible, reducing body  
560 movements as possible and keeping the eyes closed. In the beginning, the back of the chair was  
561 set up straight and the lights in the room were on. Participants were asked to remain awake  
562 with their eyes closed whilst performing the first block (awake block) of the task which  
563 consisted of 480 trials, lasting 30 min approximately. Then, the chair was reclined to a  
564 comfortable position, the lights were turned off and participants were offered a pillow and a  
565 blanket. They were explicitly told that they were allowed to fall asleep during this part of the  
566 task and that the experimenter would wake them up by making a sound (i.e. knocking on the  
567 wall) if they missed 5 consecutive trials. This block (drowsy block) also consisted of 480 trials.  
568 Then, the sequence of two blocks (awake-drowsy) was repeated. In total, participants  
569 completed 1920 trials divided into 4 blocks of 480 trials each one. The whole session lasted for  
570 3 hours approximately.

571 In Experiment#2, the procedure was similar to the Experiment#1 except for the time to  
572 fall asleep that was increased to get a higher amount of low-arousal (i.e., drowsy) trials.  
573 Participants completed a total of 2120 trials, divided into 4 blocks. The order of the blocks was  
574 the same for all participants and followed the same sequence as in Experiment#1: awake-  
575 drowsy-awake-drowsy. Awake blocks had 100 trials each one, while drowsy blocks consisted  
576 of 960 trials each one. The session lasted for 3 hours approximately.

577 In Experiment#3, upon arrival to the laboratory, participants were seated in front of a  
578 computer in a dimly illuminated, sound-attenuated room with a Faraday cage. They received  
579 verbal and written instruction about the experiment and were prepared for electrophysiological  
580 measurement. They were fitted with a 64-channel high-density actiCHamp EEG system (Brain  
581 Products GmbH, Munich, Germany) and a Polar RS800CX heart rate (HR) monitor (Polar  
582 Electro Oy, Kempele, Finland). Notice that EEG data was acquired but was not used to test the  
583 hypotheses of this study, and will be reported elsewhere. The whole session consisted of 4  
584 different blocks. The first one was an adaptation (non-exercise) block in which participants  
585 performed 100 trials while resting in a comfortable chair. Then, they got on a cycle-ergometer  
586 and completed 100 trials while warming-up at light intensity. Subsequently, they completed a  
587 self-paced 60' time-trial (i.e., high-intensity exercise) while performing the task, resulting in  
588 850 trials approximately (the number of trials slightly varied as a function of the reaction time  
589 of participants). In line with previous experiments from our laboratory,<sup>73,74,75</sup> in the self-paced  
590 time-trial participants were instructed to achieve the highest average power (watts) during the  
591 60' time-trial exercise, and were allowed to modify the power load during the exercise. They  
592 were encouraged to self-regulate effort in order to optimize physical performance without  
593 reaching premature exhaustion. That self-regulation yielded fluctuations of effort during the  
594 60' exercise period, which allowed us to study the effect of arousal on the management of  
595 probabilistic information. Once the 60' time-trial block was finished, participants completed  
596 the last block while cooling down at light intensity, which was also composed of 100 trials. All  
597 participants completed the blocks in the same order, lasting around 3 hours.

598

### 599 **Arousal classification**

600 To capture the arousal fluctuations during the transitions towards drowsiness or physical  
601 exertion at the single-trial level, we implemented two different analytical approaches which  
602 were pre-registered after data collection.<sup>9</sup>

603 In Experiment#1 and Experiment#2, the arousal level was endogenously manipulated  
604 by facilitating the natural transition from awake to sleep. This transition reduces arousal and  
605 yields a considerable proportion of drowsy yet responsive trials as seen in previous experiments  
606 from our laboratory.<sup>17,30,71</sup> This way, we were able to study the effect of arousal (i.e. baseline  
607 arousal [awake] trials vs. low-arousal [drowsy] trials) on the management of probabilistic  
608 information. Given that awake-sleep transition is characterized by a decreasing alpha range  
609 activity, together with an increasing theta range activity (Hori et al., 1994), progression of  
610 drowsiness was quantified by the spectral power of respective EEG frequency bands<sup>i</sup>. We  
611 computed the spectral power of EEG frequency oscillations for each trial from -2000 ms to 0  
612 ms in respect to the onset of a target tone using continuous wavelet transform, set from 3 cycles  
613 at 3 Hz to 8 cycles at 40 Hz. Theta (4-6 Hz) and alpha (10-12 Hz) power were then averaged

---

<sup>i</sup> Deviation from pre-registration. Originally, we aimed to use the automated offline method developed by Jagannathan and collaborators based on frequency and sleep grapho elements to detect EEG micro variations in alertness and characterize awake and drowsy trials.<sup>76</sup> However, our PRL task design, especially the pretrial duration, which was limited to 2 seconds, did not fit the task features recommended by Jagannathan and collaborators (e.g., 4 seconds pretrial duration) for a reliable characterization of awake and drowsy trials. So we decided to classify awake/drowsy trials based on theta:alpha ratio, as seen in previous experiments from our laboratory.<sup>17,30,71</sup>

614 individually for each trial across central (E36, E104) and occipital (E75, E70, E83) electrodes  
615 for theta and alpha rhythms respectively. Finally, theta/alpha ratio was computed and smoothed  
616 with a 4-point moving average resulting in a single “sleepiness” value per trial. Visual  
617 inspection of theta/alpha ratio and RT dynamics of each participant confirmed the presence of  
618 clear sleepiness-related fluctuations during the experimental session, especially during drowsy  
619 blocks. Those participants who did not show clear fluctuations of the theta:alpha ratio were  
620 removed from final analyses (5 subjects). Then, each trial for each participant was initially  
621 categorized as drowsy (top 33% of lower theta-upper alpha ratio scores) or alert (lowest 33%).  
622 Further, following the sleep hysteresis physiology criteria<sup>77</sup> isolated awake trials within  
623 prolonged periods of drowsy ( $\geq 10$  trials) were considered as drowsy to account for the gradual  
624 homeostatic change during the sleep transition. In addition, the first 100 trials of each block  
625 (awake and drowsy) were considered as awake trials.

626 In Experiment#3, the arousal level was endogenously manipulated by facilitating the  
627 natural transition from a resting state to high-intensity physical exercise. This transition  
628 increases the arousal level progressively, with continuous fluctuations that affect cognitive  
629 performance as seen in previous studies from our laboratory.<sup>40,75,78,79</sup> We captured these arousal  
630 fluctuations at a single trial level (moderate arousal trials, high arousal trials) by using the HR  
631 response. To address the intersubject variability, HR data were transformed into differential  
632 scores relative to the HRmax estimated using the equation of Tanaka et al., (2001)<sup>80</sup>, a reliable  
633 and well-established method to calculate HRmax in healthy individuals. Then, moderate and  
634 high arousal trials were characterized based on percentage relative to HRmax. HR between  
635 60% and 80% of HRmax were considered as moderate arousal, while HR higher than 80%  
636 HRmax were considered as high arousal. Due to technical issues with HR monitoring, 4  
637 subjects were removed for further analyses.

638

### 639 **Behavioural data analysis**

640 In probabilistic reversal learning paradigms, participants are instructed to infer an abstract rule  
641 from the feedback they receive, knowing that sometimes it might be misleading and that the  
642 rule might change. Since a reversal is triggered when a high-level accuracy is reached, the  
643 number of responses needed to attain a reversal is considered one of the main indices of  
644 performance. To delineate the microstructure of learning derived from trial-by-trial responses  
645 we considered the likelihood of switching the pattern choice across trials as a function of the  
646 amount of consecutive negative feedback received. The likelihood of switching was considered  
647 the main index of strategic behaviour, and was divided into 2 different strategies: *i)*  
648 Probabilistic switching: the proportion of trials when the participants change the pattern choice  
649 with little (one negative feedback against the choice) or no evidence (no feedback against the  
650 choice) of an actual rule change; *ii)* Perseveration: likelihood that participants stay with the  
651 seemingly incorrect choice even after receiving two or more negative feedbacks in a row).

652 The number of breakdown sections was also used as an index of performance. We  
653 defined a breakdown as a section of time in which participants ‘lose’ the task, and do not follow  
654 any strategy, being unable to reach a change of trend during more than 20 consecutive trials.  
655 RT, accuracy, and omissions were also checked as secondary indices of behavioural  
656 performance.

657 Participants with overall accuracy under 60% or less than 3 reversals attained during  
 658 the baseline period were excluded (i.e., 4 subjects from Experiment 1; 2 subjects from  
 659 Experiment 2; 6 subjects from Experiment 3).

660

## 661 **Statistics**

### 662 *Single-subject analysis*

663 In order to test the hypotheses, we took a set of strategies. We first captured the direction of  
 664 effects for each of the key performance variables (i.e., RAR, RT, accuracy, omissions, and  
 665 switching likelihood), and contrasted them for each participant, obtaining an indication of the  
 666 direction and strength of the effects per participant. Descriptive and distribution measures, as  
 667 well as single-subject statistics, were used as guidance of the variability of effect size in single  
 668 variables, and for guiding the previously defined exploratory hypotheses. Per participant, effect  
 669 sizes were calculated and depicted for each of the key performance variables to check the effect  
 670 size of individual differences across arousal states.<sup>ii</sup>

671

### 672 *Group analysis*

673 To investigate the management of probabilistic information as a function of arousal, we  
 674 conducted mixed-effects analyses including data from the three experiments collapsed into a  
 675 single dataset with RAR as the main index of performance. In face of the diversity of samples'  
 676 characteristics and experiment features, we fit RAR using hierarchical linear mixed-effects  
 677 modelling, as implemented in the lme4 R package.<sup>81</sup> We treated RAR as obeying to a  
 678 hierarchical data structure with arousal as fixed effect, and participant (level 2) nested into  
 679 experiment (level 1) as random effects. This random part was common to all models. We tested  
 680 the specific hypothesis by using the same approach based on multilevel linear mixed-effects  
 681 modelling. Different variables (i.e., probabilistic switching, perseveration, breakdowns, RT  
 682 variability and omissions) were analysed in a multilevel data structure, with the fixed (arousal)  
 683 and random effects (experiment/participant) adjusted to the specific hypothesis tested.

684 Models were compared using the Akaike Information Criterion (AIC), and a likelihood  
 685 ratio test. Notice that AIC does not assume that the true model is among the set of candidates  
 686 (and is just intended to select the one that is closest to the true one). In our case, fitting decisions  
 687 were not about the truthiness of models, but to include or not a given factor. For model  
 688 comparisons performed to identify the best-fitting model, a relatively lenient 0.010 p-value  
 689 criterion was adopted.

690 Causal mediation analyses were conducted to estimate the proportional direct and  
 691 indirect effects of arousal on task performance through probabilistic switching and  
 692 perseveration strategies (mediators) using the “mediation” package in R<sup>iii</sup>.<sup>82</sup> This method  
 693 allowed us to assess a confidence interval of the mediation effect itself using rigorous sampling  
 694 techniques with fewer assumptions of the data. The average causal mediation effect was

---

<sup>ii</sup> Deviation from pre-registration. Spearman rank-order correlation tests and Bayes factors were finally not performed to estimate the degree of association between switch likelihood as a function of consecutive negative feedbacks and arousal states. We will check the slope and effect.

<sup>iii</sup> Deviation from pre-registration. The mediation analysis was not initially included in the pre-registration, however, we decided to run it in order to test whether the impairment in performance in low and high arousal states could be attributed to the different maladaptive behavioural patterns.

695 determined using a nonparametric bootstrapping method (bias-corrected and accelerated; 1000  
696 iterations) and reported as standardized  $\beta$  regression coefficients for direct comparison with  
697 each other. Confidence intervals were obtained using a quasi-Bayesian approximation.

698

### 699 **Pre-registration**

700 The hypotheses and analyses plan were pre-registered in the OSF repository after data  
701 collection but prior to data observation and analysis (<https://osf.io/tzw6d>). All deviations from  
702 the pre-registered procedures and analysis plans are transparently identified in the manuscript.

703

### 704 **Data and code**

705 Data and codes used for the analyses presented here are available at the OSF repository  
706 (<https://osf.io/xk379/>). Raw data may be found at <http://doi.org/10.5281/zenodo.4559813>.

707

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## CRedit author statement

**Luis F. Ciria:** Conceptualization, Methodology, Software, Data curation, Writing, Reviewing and Editing. **Marta Suárez-Pinilla:** Data curation, Software and Reviewing. **Alex Williams:** Data curation, Software and Reviewing. **Sridhar R. Jagannathan:** Software and Reviewing. **Daniel Sanabria:** Conceptualization, Methodology, Writing, Reviewing, Editing and Supervision. **Tristán Bekinschtein:** Conceptualization, Methodology, Writing, Reviewing, Editing and Supervision.

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