1	Reactivating Positive Personality Traits During Sleep
2	Promotes Positive Self-Referential Processing
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37 Abstract

- 38 Positive self-view is evident by a bias in favor of positive self-referential processing, as
- 39 individuals tend to endorse positive characteristics over negative ones when making self-
- 40 judgments. While research suggests that a positivity bias can contribute to psychological well-
- 41 being, it remains unclear how to enhance positive self-referential processing. Here, we reported
- 42 an integrated training procedure that aimed at enhancing individuals' positive self-referential
- 43 processing. Specifically, participants engaged in a cue-approach training task (CAT) during
- 44 wakefulness where they gave speeded motor responses to positive personality traits. In a
- 45 subsequent nap, we unobtrusively re-played half of the trained positive traits during participants'
- 46 slow-wave sleep to reactivate memories associated with these positive traits (targeted memory
- 47 reactivation, TMR). Upon awakening, we found that CAT+TMR enhanced participants' positive
- 48 self-referential processing, as evidenced by faster endorsement of positive traits. Further analysis
- 49 revealed that this enhancement was associated with specific brainwave patterns during sleep:
- 50 delta (1–4 Hz) traveling waves moving from posterior to anterior brain regions. These findings
- 51 demonstrate the potential benefits of integrated wakeful cue-approach training and sleep-based
- 52 memory reactivation in strengthening positive self-referential processing.
- 53 Keywords: self-positivity, cue-approach training, targeted memory reactivation, traveling wave,
- 54 self-referential processing, slow-wave sleep

55

56 Introduction

57 People often perceive themselves through rose-tinted lenses, exhibiting a positivity bias (Taylor & Brown, 1988; Zell et al., 2020). This positivity bias is evident in self-referential judgments, as 58 59 people preferentially choose positive personality traits to describe themselves and have better 60 memories for positive traits compared to negative ones (Taylor & Brown, 1988; Watson et al., 61 2007; Guenther & Alicke, 2010; Romero et al., 2016; Dainer-Best et al., 2017; Collins & Winer, 62 2023). This positive self-referential bias is commonly associated with lower levels of depressive 63 symptoms (e.g., self-doubt, worthlessness), and is crucial for mental well-being, especially when 64 facing self-threatening information (Sowislo et al., 2013). While the psychological benefits of positive self-referential processing is well-established (Taylor & Brown, 1988; Colombo et al., 65 66 2020; Orth et al., 2022; Weisenburger et al., 2023), a significant gap exists in understanding how 67 to effectively enhance this process (Orth et al., 2022; Hoffmann et al., 2023). To address this gap, 68 we integrated two procedures that may enhance positive self-referential processing: (1) wakeful 69 cued-approach training (CAT, Schonberg et al., 2014), and (2) a sleep-based targeted memory

reactivation procedure (TMR, Oudiette and Paller 2013).

71 The CAT task prompts participants to give speeded motor responses to cued stimuli, 72 ultimately increasing positive evaluations or preference toward these trained stimuli (Schonberg 73 et al., 2014; Salomon et al., 2018; Schonberg & Katz, 2020; Itzkovitch et al., 2022). While CAT 74 has been used to alter preferences for various stimuli, such as food, abstract patterns, images (for 75 a review, see Salomon et al. 2018), its impact on higher-order social-cognitive processes such as 76 self-referential processing remains unexplored. Complementing the wakeful CAT, the TMR aims 77 to promote memory consolidation during post-training sleep, a phase vital for stabilizing newly 78 acquired memories. During sleep, covert, repeated memory reactivation contributes to memory 79 consolidation, notably during non-rapid eye movement (NREM) sleep characterized by the <480 Hz slow-wave activity (Diekelmann & Born, 2010; Rasch & Born, 2013; Klinzing et al., 2019; 81 MacDonald & Cote, 2021; Brodt et al., 2023). TMR entails replaying memory-related sensory 82 cues to sleeping participants, further strengthening episodic memories or even changing 83 subjective preferences during NREM sleep (Creery et al., 2015; Hu et al., 2015; Schreiner & 84 Rasch, 2015; Cairney et al., 2016; Ai et al., 2018; Lewis & Bendor, 2019; Abdellahi et al., 2023; 85 for a meta-analysis of TMR, see Hu et al., 2020). Here, in the context of self-referential 86 processing, we hypothesize that the integration of wakeful CAT and sleep-based TMR could 87 change how individuals perceive and endorse positive personality traits as self-descriptive. 88 Specifically, while CAT primes the brain to be more receptive to specific positive traits, TMR 89 works to consolidate these traits during sleep, potentially leading to more robust and enduring 90 positive self-referential judgments. Therefore, we tested the joint impact of CAT and TMR on 91 the positive self-referential processing, particularly focusing on how they influence the 92 immediate and long-term endorsements and retention of positive personality traits.

During NREM sleep, cardinal neural oscillations such as slow oscillations (<1 Hz), delta waves (1–4 Hz) and the 12-16 Hz spindles are instrumental in mediating memory reactivation and consolidation (Born & Wilhelm, 2012; Rasch & Born, 2013; Antony et al., 2017; Klinzing et al., 2019; Schreiner et al., 2021; Petzka et al., 2022; Brodt et al., 2023). Specifically, in TMR, researchers repeatedly found that the cue-elicited delta and sigma EEG power changes predicted TMR benefits (Lehmann et al., 2016; Blume et al., 2017; Göldi et al., 2019; Denis & Payne, 2023; Liu et al., 2023; Schechtman et al., 2023; Xia et al., 2023). While these findings have

100 significantly advanced our understanding, they remained silent on how propagation of sleep EEG 101 oscillations may contribute to memory consolidation. The propagation of EEG oscillations across 102 different brain regions, known as traveling waves, has been increasingly recognized for their 103 significance in linking brain function to behavior (Muller et al., 2018). Importantly, slow 104 oscillations and spindles have been observed to manifest as robust traveling waves propagating 105 across the cortices (Muller et al., 2018). Such traveling waves may coordinate cross-region 106 information flow and neural communications during sleep, which can be crucial for reactivating 107 and consolidating memory traces (Massimini et al., 2004; Murphy et al., 2009; Hangya et al., 108 2011; Kurth et al., 2017). However, to date, how delta slow-wave activity as traveling waves 109 may modulate memory is not yet understood. Given that the TMR employs external sensory cues 110 to trigger internal memory reactivation, it is plausible that upon processing auditory cues such as 111 the spoken positive traits, the delta slow waves would exhibit a forward trajectory from the 112 posterior to the anterior frontal brain regions, contributing to effective processing of positive 113 traits and thus influencing positive self-referential processing. Indeed, recent research suggests 114 that a forward patterns of traveling wave could aid in the bottom-up processing of stimuli in a 115 wakeful state (Alamia et al., 2023). We thus hypothesize that during TMR, the forward slow 116 traveling wave activity could align sensory processing with semantic processing of positive traits

117 during sleep, thereby enhancing positive self-referential processing.

118 Here, we employed an adapted version of the well-established self-referential encoding 119 task (SRET) to quantify participants' self-referential processing (Derry & Kuiper, 1981; Dainer-120 Best et al., 2017, 2018; Collins & Winer, 2023, for procedure and tasks, see Figure 1). In 121 addition to this SRET, we assessed participants' recall of self-referential traits from the SRET in 122 a free recall task, and self-referential preferences in a probe task. To examine the immediate and 123 possible long-term effects of TMR, we measured participants' self-referential processing twice: 124 immediately after the TMR and one-week later. Our findings revealed that the integration of 125 CAT and TMR facilitated the endorsement speed of positive personality traits immediately after 126 sleep. Moreover, analysis of cue-elicited EEG showed that the strength of 1–4 Hz forward 127 traveling waves predicted the endorsement speed of positive traits during immediate test and the 128 endorsement of positive traits one-week later.





130 Figure 1 An overview of experimental design and main tasks

131 (A) The task flow illustrates the baseline tests (phase 1), CAT and post-CAT tests (phase 2),

132 sleep-based TMR (phase 3), and post-TMR tests (phase 4), followed by a delayed tests phase

133 after one week (n = 35). (B) Modified SRET, in which participants made speeded binary

endorsement task to determine whether a personality trait was descriptive of oneself, followed by

rating the accuracy of specific traits in describing themselves within the same trial (i.e.,

endorsement level). After completing the baseline SRET, participants performed a self-

referential free recall task. In both the post-TMR and the one-week delay phases, participants
completed the free recall task and the SRET with binary endorsements while omitting ratings. (C)

An exemplar trial of CAT, in which participants either passively viewed positive traits presented

140 visually and aurally (i.e., NoGo trials) or pressed a button when they saw a white circle appear

141 immediately after the positive trait onset (i.e., Go trials). The average GSD (go-signal-delay) was

142 900 ms. (D) Probe test, participants were presented with pairs of positive Go and NoGo traits and

- 143 were asked to select which trait was more self-descriptive. Note that Go and NoGo traits in each
- 144 pair were matched on baseline self-descriptive ratings (see Methods for a full description of the
- 145 procedure and experimental tasks).

146 **Results**

147 Awake CAT promoted self-referential preferences

148 First, to examine whether CAT promoted the preferences of positive Go traits, we analyzed the

149 proportion of trials in which participants preferred Go traits over NoGo traits as better self-

- 150 descriptive in the probe task (Figure 1D), using a generalized linear mixed model with
- 151 participant factor as a random effect (GLMM, see Method for specific model). In each Go/NoGo
- 152 pair, both traits had comparable initial endorsement level based on the baseline SRET rating
- 153 phase. Consistent with previous CAT research (Salomon et al., 2018), we found that participants
- 154 were more likely to choose Go over NoGo traits despite their comparable baseline endorsement
- 155 level: mean proportion $\Box = \Box 53.3\%$ (vs. chance level 50%), odds ratio (OR) $\Box = \Box 1.24$, 95% CI
- 156 [1.09, 1.41], p = 0.001. This result suggested that the CAT specifically increased participants'
- 157 self-referential choice of the Go traits in the probe task.

158 Awake CAT + sleep TMR enhanced positive self-referential endorsement and speed

159 Having established the effectiveness of CAT in enhancing the preferences of positive Go traits,

- 160 we next examined how TMR may further influence positive self-referential processing.
- 161 Specifically, we analyzed two outcome variables from the SRET task, including binary
- 162 endorsement choice, and reaction times (RTs) when endorsing positive traits. Note that we
- 163 applied false discovery rate (FDR) corrections for all multiple comparisons.

164 To examine positive self-referential endorsements change, we ran a GLMM using 165 baseline endorsement rating value as a covariate, TMR condition (Go-cued, Go-uncued, and 166 NoGo-uncued) and time (baseline, post-TMR, and delay) as fixed effects, and participant factor 167 as a random effect to predict endorsement choices (yes, no) of positive traits. Our results revealed a significant TMR effect, $\Box^2(2) = 8.02$, p = 0.018. Particularly, participants endorsed 168 more Go-cued traits than NoGo traits (p = 0.026). No differences were found between other 169 170 conditions (all ps > 0.12). However, neither time nor the TMR by time interaction were significant, $\Box^2(2) = 2.52$, p = 0.284, $\Box^2(4) = 2.80$, p = 0.592, respectively. 171

172 Given that choice speed could indicate preferences (Konovalov & Krajbich, 2019), we 173 analyzed item-level RTs when participants endorsed positive traits via an LMM including TMR 174 conditions and time as fixed effects. The results showed a significant main effect of time, F(2,175 (35) = 8.37, p = 0.001, indicating that RTs became faster from baseline to post-TMR and delay (all ps < 0.001), while no difference was found between post-TMR and delay (p = 0.126). No 176 177 significant main effects were observed for the TMR condition, F(2, 4921) = 0.73, p = 0.480. 178 Notably, we found a significant TMR by time interaction, F(4, 4922) = 3.19, p = 0.013 (Figure 179 2A). Post-hoc comparisons revealed that in the post-TMR, participants were significantly faster 180 in endorsing Go-cued traits than NoGo-uncued traits (p = 0.022). In contrast, this pattern was not 181 observed in the delay testing (p = 0.414). Other comparisons across different time points did not 182 yield any significant differences in RTs (Go-uncued versus NoGo-uncued, Go-cued versus Go-183 uncued traits, all ps > 0.1). Taken together, these results suggest that the CAT+TMR jointly

184 facilitated endorsement speed for Go-cued positive traits compared to NoGo-uncued traits.

To address the concern that the observed RT differences might be solely due to the influence of CAT, we conducted an additional LMM in two additional independent behavioral samples to investigate if RT differences in endorsing positive traits were present between Go and NoGo trait words. These samples consisted of one group undergoing active CAT with 'Go' training trials and another group exposed to passive CAT without such active components (See SOM for behavioral sample details). In this analysis, we included CAT conditions (Go vs. NoGo), time (baseline, post-CAT, and delay), and group (active vs. passive) as fixed effects,

- 192 with participants as random effects, focusing on positive endorsement RTs. The results only
- showed a significant main effect of time, F(2, 75) = 12.62, p < 0.001, with faster endorsements
- after CAT and during the delay tests compared to the baseline (all ps < 0.001). However, we
- 195 observed no significant main effects or interactions specifically attributable to CAT (all
- 196 ps > 0.35). These findings, therefore, suggest that it is the combination of CAT and TMR, rather
- 197 than CAT alone or solely repetition of positive trait words, that promotes the endorsement speed
- 198 of positive traits.





200 Figure 2 Behavioral results across time in the SRET tasks

201 (A). Fitted values for the interaction effect of TMR conditions and time on predicting log-

transformed RTs for endorsing positive traits during the SRET. Error bars indicate 95%

203 confidence intervals (CIs). (B). Fitted values for the interaction effect of TMR conditions and

time on predicting recall percentage for positive traits during the SRET. * = p < 0.05.

205 Overall enhancement of positive self-referential memory recall following CAT and TMR

206 Recognizing the established efficacy of TMR in enhancing memory performance (Hu et al.,

207 2020), our study specifically investigated whether integrating CAT with TMR would improve

- 208 the recall of positive traits in self-referential memory tests. We used a GLMM including TMR
- 209 (Go-cued, Go-uncued, NoGo-uncued) and time (baseline, post-CAT, post-TMR, delay) as fixed
- 210 effects, and participant factor as a random effect to predict percentage of positive traits
- 211 participants recalled as self-referential. Results revealed a significant time effect, $\Box^2(3) = 72.70$,
- 212 p < 0.001, with post-hoc comparisons indicating that compared to baseline, there were
- significantly higher recalls of positive traits at post-CAT, post-TMR and delay tests (ps <0.001,
- Figure 2B). Moreover, recall declined from the post-CAT to the delay (p = 0.011) and from the
- 215 post-TMR to the delay (p < 0.001), while there was no difference between post-CAT and post-
- TMR recalls (p = 0.063). Furthermore, we found no significant TMR effect, $\chi^2(2) = 1.04$, p = 0.063
- 217 0.595, nor any significant TMR by time interaction, χ^2 (6) = 2.29, *p* = 0.891, indicating that CAT 218 or TMR did not selectively change recall performance. Lastly, when examining the recall of
- negative traits, the time effect was not significant, $\chi^2(3) = 3.35$, p = 0.341, suggesting that the
- 1219 negative traits, the time effect was not significant, $\chi^{2}(5) = 5.55$, p = 0.541, suggesting th
- 220 increased recall was specific to positive traits.

221 To ascertain whether CAT+TMR jointly contributed to the overall enhancement of 222 positive self-referential memory, we also contrasted the current CAT+TMR group with the 223 abovementioned two additional independent behavioral samples (active CAT and passive CAT 224 groups, see SOM for behavioral sample details). A subsequent GLMM, accounting for earlier 225 recall as a covariate, showed a significant group effect, $\gamma^2(2) = 9.29$, p = 0.010. Participants who 226 received both TMR and CAT exhibited superior recall of positive traits compared to the passive 227 CAT group (p = 0.008), but not higher than the active CAT group (p = 0.085), while no 228 significant difference was found between the active and passive CAT groups (p = 0.229, all FDR 229 corrected). Together, these result highlights that combining CAT and TMR had long-lasting 230 impact in facilitating self-referential recall of positive traits.

Together, the findings demonstrated that CAT shifted preferences towards positive traits,
 while combining CAT and subsequent TMR effectively enhanced positive self-referential
 processing by accelerating RTs when for endorsing positive traits.

234 Auditory processing of positive traits during sleep TMR

235 To first validate the processing of spoken positive traits during sleep, we quantified cue-elicited 236 event-related potentials (ERPs) and time-frequency resolved EEG power changes during the 237 TMR. Consistent with prior TMR research (Schreiner et al., 2015; Lehmann et al., 2016; Antony 238 et al., 2018; Schechtman et al., 2021; Abdellahi et al., 2023; Guttesen et al., 2023; Liu et al., 239 2023; Schechtman et al., 2023; Xia et al., 2023), cue-elicited ERPs showed two positive clusters 240 over frontal-central electrodes (F1/2, FC1/2, C1/C2, Fz, Cz) from 0.29 to 0.52 seconds and from 241 1.04 to 1.40 seconds (two-tailed *t*-test, cluster-based permutation-corrected p < 0.005). In 242 addition, the time-frequency analysis also identified two significant positive clusters over frontal-243 central electrodes; the delta-theta-alpha band (1 to 12 Hz, 0 to 1.66 seconds), and the sigma-244 beta band (10 to 30 Hz, 0.3 to 1.42 seconds, two-tailed t-test, cluster-based permutation-245 corrected p < 0.001, Figure 3A, B). We next examined whether EEG power changes within these 246 clusters may be associated with changes in RTs and choices during the positive self-referential 247 processing. However, we did not find significant associations (all $p_{\text{corrected}} > 0.562$ for delta-theta-

248 alpha; and > 0.364 for sigma-beta.



- 250 Figure 3 Cue-elicited power changes did not predict post-TMR endorsements or positive
- endorsement speed
- 252 (A) Grand averaged ERPs across frontal-central electrodes (F1/2, FC1/2, C1/2, Fz, Cz). Shaded
- area indicates significant time point when comparing ERPs against zero. Top right panel presents
- 254 group average scalp topography of ERPs in response to TMR cues; with black circles
- 255 highlighting the electrodes used in the ERP analysis. (B) Contour plot depicting the temporal and
- 256 spectral characteristics of the significant clusters. Cluster a represents the low-frequency delta-
- theta-alpha band (1–12 Hz), and cluster b represents the sigma-beta band (10–30 Hz), with both
- clusters showing significant changes across the TMR time course (cluster-based permutation-
- 259 corrected p < 0.001).

260 Cue-elicited slow travelling waves predicted post-TMR positive traits endorsement speed

261 Next, we investigated how traveling waves during TMR might influence post-TMR positive self-

- 262 referential processing. Given that TMR benefits would begin with effective sensory processing
- 263 (posterior cortex), followed by high-level memory processing (see Liu et al. 2023), we
- 264 hypothesized that the forward traveling waves propagating from posterior to anterior cortex
- would be critical here. Building upon methodologies used in previous research (Alamia et al.,
- 266 2023), we analyzed the strength of directionality in both forward (from posterior to anterior brain
- regions) and backward (from anterior to posterior brain regions) traveling waves within the first
- 268 2s following the TMR cue onset, using EEG signals from midline electrodes (POz, Pz, CPz, Cz,
- 269 Fz, FPz) during SWS in the 1–4 Hz frequency band (Figure 4A-C, see Method). The strength of
- 270 forward and backward traveling waves was then utilized to predict item-level binary
- endorsement choices and RTs for endorsing positive traits, across immediate and a one-week
- delay test time, with False Discovery Rate (FDR) corrections for multiple comparisons.



274 Figure 4 Slow travelling waves after TMR cue onset

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275 (A). EEG signals from midline electrodes (0–2000 ms post-TMR cue) underwent two-

276 dimensional Fourier transform (2D-FFT) decomposition, yielding power spectra. Baseline-

277 corrected spectra using shuffled data delineate backward (anterior to posterior) and forward

278 (posterior to anterior) traveling waves (detailed methodology in Alamia et al (2023). (B). Scalp

diagram illustrating slow wave directionality. Forward waves are indicated by arrows pointing to

frontal electrodes, backward waves to posterior electrodes. Asterisks mark analyzed electrode

281 locations. (C) Demonstrations of forward (left panel) and backward (right panel) traveling waves.

We used (G)LMM with intensities of traveling waves as a fixed effect, alongside the number of

trait repetitions during TMR and baseline endorsement ratings as covariates, and participant as a

random effect, to predict post-TMR endorsements and RTs when endorsing positive traits.

- 285 Results revealed that during the post-TMR test, neither forward nor backward traveling waves
- significantly predicted endorsement probabilities ($p_{corrected} > 0.272$, Figure 5A&B). Notably, forward traveling waves negatively predicted RTs when endorsing positive trats in the immediate
- forward traveling waves negatively predicted RTs when endorsing positive trats in the immediate test ($p_{corrected} = 0.002$, Figure 5C), but not in the one-week delay ($p_{corrected} = 0.766$, Figure 5G).
- Backward traveling waves showed no significant associations ($p_{corrected} = 0.686$, Figure 5D&H).
- 290 Interestingly, forward traveling waves also significantly predicted the endorsement of positive
- traits after one week ($p_{corrected} = 0.026$, Figure 5E), suggesting a delayed effect on positive self-
- 292 referential processing. Backward waves, however, did not demonstrate a significant prediction

 $(p_{\text{corrected}} = 0.060, \text{Figure 5F})$. Together, our results showed that the forward traveling waves elicited by positive traits during sleep can predict post-sleep positive self-referential processing.



Figure 5 Predictive impact of delta traveling waves on endorsement probabilities and RTs in

297 post-TMR and delay phases of the SRET task.

298 Predictions from forward and backward traveling waves on (A, B) endorsement probabilities in

- 299 post-TMR phase. (C, D) RTs in post-TMR phase. (E, F) endorsement probabilities in delay
- 300 phase. (G, H) RTs in delay phase. In panel (C, D, G, H), each data point corresponds to the fitted
- 301 value from a single trial within the LMM. Where data points overlap, they present a darker shade.
- 302 In panel (A, B, E, F), we excluded raw data points due to their binary (zero or one) nature.
- 303 Shaded area indicates 95% confidence intervals (CIs). TW: traveling wave. SRET: self-

304 referential encoding task. ** = p < 0.01. * = p < 0.05.

305 Discussion

306 By combining wakeful cue-approach training (CAT) and sleep-based targeted memory

- 307 reactivation (TMR), we found that this integrated procedure effectively enhanced participants'
- 308 positive self-referential processing. We first used CAT to heighten participants' preferences for
- 309 specific "Go" positive traits, extending the existing CAT research. Following this, TMR was
- 310 employed to re-play a subset of these Go traits during participants' SWS sleep, further enhancing
- 311 their accessibility and consequently promoting positive self-referential processing. The CAT and
- 312 TMR expedited endorsement of these Go-cued positive traits, although it did not selectively alter
- 313 self-referential memory immediately after sleep TMR. After one week delay, we observed a
- 314 general increase in positive memory recall, rather than TMR-specific changes. Additionally, the
- 315 presence of 1–4 Hz forward slow travelling waves during TMR was associated with enhanced
- 316 positive self-referential processing, indicating an important role of cross-regional forward neural

317 communications in driving behavioral benefits. These new findings contributed to our318 understanding of how to modulate and enhance positive self-referential processing.

319 We first found that the CAT successfully increased participants' likelihood to choose 320 Go over NoGo traits as self-descriptive in the probe task, demonstrating CAT's efficacy in 321 influencing self-referential choices. This finding extends the known effects of CAT on 322 consumables such as snacks (Salomon et al., 2018; Schonberg & Katz, 2020), revealing its 323 capability to shape high-level self-referential processing. Following the CAT phase, we replayed 324 a subset of the trained positive traits during sleep to determine the cumulative impact of CAT 325 and TMR on self-referential processing. Behaviorally, we found that participants exhibited faster 326 endorsement of positive traits when these traits were trained during the wakeful CAT and 327 subsequently reactivated during the SWS (Go-cued), compared to untrained traits (NoGo-328 uncued). This contrast highlights the joint benefits of CAT and TMR in facilitating the speed of positive self-referential processing. Apart from positive endorsement speed, we also observed 329 330 that participants endorsed more Go-cued positive traits than NoGo-uncued traits as self-331 descriptive independent of testing time. Previous CAT research has indicated that the CAT can 332 improve stimulus salience, effectively making these traits more prominent during the waking 333 state (Schonberg et al., 2014; Schonberg & Katz, 2020). TMR during post-training sleep, on the 334 other hand, further promoted memory reactivation and consolidation via cueing, improving the 335 accessibility and retrieval efficiency of the cued stimuli (Walker & Stickgold, 2006; Diekelmann 336 & Born, 2010; Klinzing et al., 2019; Lewis & Bendor, 2019; Brodt et al., 2023). In the context of 337 our study, the TMR following the CAT likely further augments the accessibility of the trained 338 traits, thereby speeding up their endorsements.

339 The accelerated endorsement of positive traits and the overall higher endorsement 340 probabilities for CAT+TMR traits can be partially explained by neural oscillations during sleep 341 and TMR. In contrast to power spectral analysis that often concentrates on regional oscillations, 342 the concept of traveling waves encompasses a wider array of neural characteristics. These 343 include both spatial propagation and frequency property, offering a more comprehensive view of 344 the spatial-temporal dynamics of brain activity during sleep (Massimini et al., 2004; Muller et al., 345 2018; Zhang et al., 2018; Halgren et al., 2019). Our study observed that the 1-4 Hz delta forward 346 traveling waves, moving from posterior to anterior brain regions within 2 seconds after cue onset, 347 were significantly associated with post-TMR endorsement RT and the endorsement of positive 348 traits following a one-week delay. This finding, for the first time, suggests that delta forward 349 traveling waves are pivotal in memory consolidation during sleep, broadening the existing 350 knowledge of the neural mechanisms supporting sleep-mediated memory consolidation 351 (Massimini et al., 2004). Previous studies have indicated the thalamic origin of delta waves 352 during sleep (Adamantidis et al., 2019) and its integral role in memory consolidation (Schreiner 353 et al., 2022). Our results contribute to this body of knowledge by demonstrating that the delta 354 waves, as traveling waves, may also contribute to memory consolidation during exogenous 355 memory reactivation. Furthermore, the forward direction of delta traveling waves implies the 356 bottom-up processing of external cues in TMR sleep. Specifically, re-playing spoken positive 357 traits during sleep, may initiate basic auditory processing at the posterior brain regions that 358 advanced to high-level semantic and self-referential processing at the frontal regions. This 359 forward propagation of delta slow waves may effectively integrate the positive traits into one's 360 self-scheme, contributing to positive self-referential processing. Although the exact 361 neurocomputing processing remains elusive, parallels can be drawn from awake-state studies. A

noteworthy study highlighted the bottom-up processing of visual stimulus is facilitated by alphaband forward traveling waves (Alamia et al., 2023). These waves spatiotemporally organize
distributed brain areas, enabling efficient processing of external stimuli. Consequently, we
proposed that the sleeping brain processes spoken positive traits by coordinating different brain
regions through traveling waves, particularly the forward delta traveling waves that propagated
from posterior to anterior brain regions.

368 When evaluating participants' self-referential memories using a free recall task, we did 369 not find significant main effects of CAT or TMR. This result may stem from the experimental 370 design, where participants engaged in the free recall task twice prior to sleep. Repeated recall 371 may induce fast memory consolidation that makes the self-referential memories less susceptible 372 to TMR (Antony et al., 2017; Liu et al., 2023). Notably, a week later, participants showed an 373 overall enhanced memories for positive traits compared to the baseline, regardless of CAT or 374 TMR manipulations. Supporting this, broader sleep and TMR studies indicate memory 375 enhancement can often be observed over extended periods (Rakowska et al., 2021; Barner et al., 376 2023). Even more intriguingly, we found that this non-selective, general memory enhancement 377 was only observed in the CAT+TMR group, but not in the other two groups (CAT only, or 378 passive CAT). These findings suggest that the post-CAT sleep and TMR may further enhance 379 the overall positive self-referential memories, regardless of cueing. Indeed, previous TMR 380 research suggested that memory reactivation during sleep may have generalized benefits: in 381 addition to enhancing cue-specific memories, TMR also strengthened uncued memories that 382 shared the same context as the cued memories, leading to overall benefits of both cued and 383 uncued memories (Schechtman et al. 2023; see also Oudiette et al., 2013 for TMR generalization 384 effects).

385 Future directions and limitations shall be discussed. First, our study follows most prior 386 research in administering the TMR during the NREM sleep, given the established link between 387 NREM sleep and TMR benefits (see Lewis and Bendor 2019; Hu et al. 2020). However, research 388 also pinpoints the role of REM sleep in modulating emotional memory and vocabulary learning 389 (Batterink et al., 2017; Hutchison et al., 2021). Future research could investigate how TMR 390 during REM sleep, and how the REM-related neural activity may impact the consolidation of 391 self-referential memories. Second, while positive self-referential processing is linked with mental 392 wellness (Wisco, 2009; Lou et al., 2019; Collins & Winer, 2023), our study did not examine how 393 our procedure may impact outcomes that bear direct clinical relevance, such as depression-394 related symptoms. Future research is warranted to investigate whether enhancing positive self-395 referential processing may directly alleviate depressive symptoms (Orth et al., 2022; Hobbs et al., 396 2023). Third, while our research question concerns self-referential memory, we did not include 397 non-personal traits as a control condition. Future studies could consider including a control task, 398 specifically designed to disentangle non-self-referential memory from self-referential memory in 399 understanding the CAT and TMR effects.

In conclusion, our study presents a novel approach in enhancing positive self-referential
 processing by combining wakeful motor training and sleep-based memory reactivation. In
 addition to behavioral benefits, our findings underscore the importance of cue-related forward
 delta traveling waves in predicting the speed of endorsing positive traits, establishing a direct
 connection between traveling waves induced by TMR and positive self-referential processing.
 By reinforcing positive self-referential processing through CAT+TMR, it may be possible to

406 alter maladaptive cognitive biases or restore self-esteem, contributing to improved mental health407 outcomes.

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422 **Conflict of interest statement**

- 423 None
- 424

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686 Data and code availability

The data and analytical code supporting the study's findings are available at the Open Science
 Framework repository: https://shorturl.at/bEG23.

689 Experimental model and study participant details

690 Participants

- 691 Our final sample included 35 participants with valid behavioral and EEG data (8 males, Mage ±
- $SD = 20.83 \pm 2.20$ years), which is comparable to recent TMR studies (e.g., Schechtman et al.
- 693 2023). Nine additional participants had inadequate number of cues (< =3 rounds) due to
- relatively short slow-wave sleep (SWS). To ensure signal-to-noise ratio in the EEG analyses,
- 695 experiments for these participants were terminated following the TMR, and data from these nine
- 696 participants were not included in subsequent analyses. An additional participant was excluded
- because he or she reported hearing the cues during sleep. To facilitate sleep in the lab, we asked
- 698 participants to wake up one hour earlier than their usual waking time and to avoid consuming
- 699 caffeinated drinks on the day prior to and of the experiment. Participants were pre-screened
- regarding any current or history of sleep, psychiatric, or neurological disorders and had normal
- or corrected-to-normal vision. Participants received monetary compensation for their time (250
- RMB, ~36 USD), and gave written consent prior to the experiment. The study was approved by
- the Human Research Ethics Committee of the University of Hong Kong.

704 Materials

- All experimental procedures were implemented in E-Prime® 3.0 (Psychology Software Tools,
- Inc., Sharpsburg, Pennsylvania, USA). A pilot group of 20 participants rated personality traits
- 707 (two characters trait words) on a scale from 1 (extremely negative) to 9 (extremely positive). We
- selected 60 positive personality trait adjectives (e.g., 'clever', $M \pm SD = 6.92 \pm 0.44$) and 60
- negative personality trait adjectives (e.g., 'lazy', $M \pm SD = 3.00 \pm 0.44$; see SOM for the
- 710 complete list of personality traits). Each spoken trait lasted around 1 second (range: 0.72-1.08s,
- 711 $M \pm SD = 0.91 \pm 0.08s$). During the TMR phase, we used a neutral trait (valence rating: 4.9) as a
- control word.

713 Method details

714 Task overview

715 Participants attended two lab sessions, scheduled approximately one week apart. In the first

session, participants arrived to the lab at approximately 12:00 pm (exact arrival times ranged

between 11:30 am to12:30 pm), where they read and signed consent forms and were set up with

EEGs. Subsequently, a series of four task phases began in which participants completed a

- number of tests, beginning with baseline tests in the first phase, followed by CAT and post-CAT
- tests in the second phase, sleep-based TMR in the third phase, and post-TMR tests in the fourth
- 721 phase. In the preliminary baseline phase, participants completed computer-based personality
- questionnaires, serving as a cover story for the personality trait words (hereafter, traits) presented
- to them in the following SRET. During the SRET, participants rated the extent to which specific

traits described themselves. Participants then completed a self-referential free recall test. In the

- second phase, participants manually responded to positive traits (i.e., Go traits), prompted by
- visual and aural cues presented on screen and from a nearby loudspeaker (CAT). Participants
- then completed a free recall test and a probe test, in which they were presented with Go and
- NoGo trait word pairs and asked to select the trait word that was more self-descriptive. In the third phase, half of the positive traits were aurally re-played to sleeping participants during slow
- third phase, half of the positive traits were aurally re-played to sleeping participants during slowwave sleep (SWS). Then, in the fourth phase, participants completed the same free recall test,
- 730 wave sleep (SwS). Then, in the fourth phase, participants completed the same free fecan test, 731 probe test, and SRET. In the second lab visit (~ 7 days later), participants completed the same
- free recall test, probe test and SRET as previously completed in the final phase of the first visit to
- received in the possible long-term TMR effects. Thus, they completed four self-referential free
- recall tests (baseline, post-CAT, post-TMR, delay), three SRETs (baseline, post-TMR, delay),
- and three probe tasks (post-CAT, post- TMR, delay).

736 Baseline tasks

737 Participants completed preliminary computer-based personality questionnaires, including the

- 738 Rosenberg Self-Esteem Scale (RSES) (Rosenberg, 1965), Narcissistic Personality Inventory
- (NPI) (Raskin & Hall, 1981), Big Five Inventory (BFI) (John et al., 1991), Beck Depression
- 740 Inventory-II (BDI-II) (Beck et al., 1996), State-Trait Anxiety Inventory (STAI state and STAI
- trait) (Spielberger, 1983), and Barratt Impulsiveness Scale (BIS-11) (Patton et al., 1995).
- 742 Completing these questionnaires served as a cover story for the subsequent self-referential
- recoding task (SRET): participants were told that the personality traits that would be presented
- in the SRET were from their questionnaire data (for descriptives, see Table S1).

745 In the SRET (see Figure 1B), a cross symbol was presented on a computer screen at the 746 beginning of each trial for 0.5 seconds, followed by the presentation of the sentence 'I think this 747 word is applicable to me' in the center of the screen for another 0.5 seconds. After 1.2 to 1.4 748 seconds, participants were presented with a random word, given visually in written form and 749 aurally from a speaker, from a selection of 120 adjectives for 0.8 seconds. After, participants 750 were shown a blank screen for another 0.8 seconds and then were prompted to select if a trait 751 word applied to them within 2.5 seconds by moving the mouse cursor continuously. The spatial 752 location of 'Yes' and 'No' responses were counterbalanced (upper left/upper right or upper 753 right/upper left). Following a 'Yes' response, participants were asked to rate the extent to which 754 a trait word applied to them on a scale ranging from "slightly accurate" to "extremely accurate". 755 covertly equating to values from 1 to 50; following a 'No' response, participants were asked to 756 rate the extent to which a trait word did not apply to them on a scale ranging from "slightly 757 inaccurate" to "extremely inaccurate", covertly equating to values from -50 to -1.

Following the SRET, participants completed a self-referential free recall task. Unlike previous free recall tasks wherein participants wrote down as many traits as possible, here, participants were asked to recall only the traits they had been presented with and they endorsed (i.e., "yes" response) during the previous SRET. Participants typed each recalled trait on a computer one at a time. Therefore, performance during this version of the recall task reflected self-referential memories.

764 Traits selection in the probe task

For each participant, we ranked all 60 positive traits in ascending order based on their baseline

rating and least self-descriptive, to 60, being the

highest rating and most self-descriptive). We next equally divided these 60 traits into 'Go' and

⁷⁶⁸ 'NoGo' trials, forming 30 Go-NoGo pairs for each participant. We chose traits for each pair

based on each trait rating's rank orders (i.e., from 1 to 60), to ensure that the Go and NoGo traits

had comparable baseline ratings (p = 0.64, for details, see SOM and Figure S1A). For the post-

- TMR probe task, these Go/NoGo pairs were further categorized into cued (Go-cued) and uncued
 (Go-uncued) conditions, with each condition having 15 trait pairs. Full details for the trait
- allocations in the CAT and the probe task are provided in Supplementary Figure 1.
- anocations in the CAT and the probe task are provided in Supplementary Figure 1.

774 CAT and post-CAT tests

775 Following baseline assessments, participants completed a cue-approach training (CAT) task (see 776 Figure 1C). For each CAT trial, a positive trait was presented visually and aurally for 1.2 777 seconds. For Go trials, each trait was paired with a delayed Go cue that required participants to 778 press a button as quickly as possible before the trait's offset. To maintain participants' attention, 779 we used an adaptive response window. Specifically, the go-signal-delay (GSD, the delay 780 between trait onset and Go-cue onset) was approximately 0.9 second. If the participants gave a 781 timely response (i.e., button press before the offset of the trait), the GSD was increased by 17 ms 782 to increase task difficulty. If participants failed to make a button press before the offset of the 783 trial, the GSD was reduced by 50 ms to reduce task difficulty (Salomon et al., 2018; Schonberg 784 et al., 2014). Conversely, for NoGo trials, participants merely viewed and listened to the traits 785 without any behavioral responses. All 60 positive traits were presented randomly in each of the 786 five blocks during the CAT, resulting in a total of 300 trials. Participants could take a 0.5-1-787 minute break between blocks. While previous CAT research adopts over 10 blocks of training 788 (Salomon et al., 2018), we chose to only include 5 blocks so as to avoid ceiling effect in 789 subsequent memory recall. This CAT task was followed by a 5-minute working memory task, 790 serving as distractions.

Following the working memory task, participants proceeded to a 3-minute post-CAT selfreferential free recall task, which was identical to the baseline task. Subsequently, a post-CAT probe task was administered to evaluate the impact of CAT.

794 In the probe task (see Figure 1D), participants were presented with Go and NoGo traits in 795 pairs and were asked to choose which trait would be more self-descriptive. Within each trial, the 796 Go and NoGo traits were matched on baseline endorsement ratings, so that preferential choices 797 of Go traits would indicate the CAT training effects. The positions of the Go/NoGo traits per pair 798 were randomly assigned to the upper-left/right or upper-right/left sides of the monitor in the first 799 block, and were swapped in the second block. Each trial started with a fixation cross (1 second), 800 followed by the side-by-side presentation of two traits. Participants selected the trait that would 801 best describe them by clicking a push button below the trait within 2.5 seconds. The chosen trait 802 was then highlighted by a button-press shaped image for 0.5 seconds. If participants exceeded 803 the 4-second response time, a prompt would appear during the confirmation phase, urging them 804 to respond quickly. We excluded trials with response times exceeding 3 seconds, accounting for 805 potential mouse delays.

806 Nap targeted memory reactivation (TMR)

807 Participants took a 90-minute nap in a quiet, darkened sleep chamber. Background white noise

- 808 (at ~38 dB) was played to participants throughout the duration of the nap via a loudspeaker
- 809 placed near the bed. Participants' brain and physiological activities were continuously monitored
- 810 during the map. Upon participants entered SWS for at least 2 minutes, we presented spoken
- 811 positive traits (the same spoken traits presented during the SRET and CAT tasks) at
- 812 approximately 40 dB. Note that the spoken traits (~40 dB) were played against the background 812 white poise ($_{28}$ dB) yet remained subtle to avoid arousel and waking participants up
- 813 white noise (~38 dB), yet remained subtle to avoid arousal and waking participants up.

814 The TMR began with playing a neutral trait (~0.6 s) for three times, ensuring that the 815 auditory stimulation would not wake participants up. We started playing the spoken traits if 816 participants did not show signs of arousal or changes in NREM sleep stage. During each round 817 of the TMR, half of the positive Go traits (i.e., 15 traits) were played together with the neutral 818 trait as a control word. Each trait last for about 1 second, with a randomized interstimulus 819 interval of 5–6 seconds. TMR continued as participants remained in the SWS, with a minimal 820 repetition of three rounds of stimulation, resulting in at least $3 \times 16 = 48$ trials for TMR-related

EEG analyses.

Specifically, participants were exposed to spoken traits once they entered a sustained SWS period lasting at least 2 minutes. The TMR procedure was discontinued after 30 minutes, or earlier if EEG recordings indicated micro-arousal or full awakening. If no SWS was detected within the first 40 minutes, the presentation of spoken traits commenced during the N2 sleep stage. After a total sleep session of 90 minutes, participants were awakened if they were in the N1 or N2 sleep stages, or we waited until they transitioned to these stages before awakening them. A brief 5-minute break was provided upon awakening to mitigate the effects of sleep

829 inertia.

830 Post-TMR tests

831 Participants completed the self-referential free call task, probe task, and SRET task. Here, the

832 probe task instructions were identical to the post-CAT probe task, but with randomized 'Go' and

833 'NoGo' trait positions. The SRET was similar to the baseline SRET except that participants only

834 made a Yes/No binary response to each trait, omitting the rating part.

835 One-week delayed tests

836 Participants returned to the lab about one week later to complete the delayed tests in the

following order: (1) a 3-minute self-referential free call task; (2) a probe task; (3) a SRET task.

838 The tasks were identical to the tasks in the post-TMR. Participants were not informed of the

839 delayed tasks ahead of the time. Upon completing all tasks, participants were debriefed and paid.

840 **Quantification and statistical analysis**

841 Behavioral data analysis

- 842 Statistical analyses were carried out using R (Version 4.2.1., R Core Team (2020). We
- 843 performed (G)LMMs fitted via 'glmer' and 'lmer' functions of the 'lme4' R package (Bates et al.
- 844 2014 June 23) to analyze the CAT- and TMR-induced behavioral changes. For statistical
- significance testing, we used Type III Analysis of Variance with the Satterthwaite approximation

- 846 method for the LMM and Type II Wald Chi-Square tests for the GLMM. We followed up
- significant effects with post-hoc comparisons in *emmeans* (Lenth et al., 2022) to derive the
- 848 estimated marginal means from each model. Model predictions were visualized with the
- 849 'plot_model' function from the sjPlot package (Lüdecke, 2023). Unless otherwise stated, we
- 850 used the False Discovery Rate (FDR) method to adjust for multiple comparisons to control for
- false-positive results. The significance threshold (alpha level) was set at 0.05.

852 Self-referential preference choices in the probe task

Following previous CAT research (Botvinik-Nezer et al., 2020; Salomon et al., 2018; Schonberg
et al., 2014), we ran generalized linear mixed models (GLMMs) to compare the odds of choosing
Go traits against the chance level (50%, log odds = 0; odds ratio = 1) during post-CAT phase.
Given the alternation of Go/NoGo positions (left and right) in two blocks, we included Go

- 857 position as a covariate in our model. The GLMM was defined as:
- 858 Preference Choice (Go/NoGo) ~ 1 + Position +(1|Subject ID)

859 Self-referential endorsement in the SRET

860 We employed a generalized linear mixed model (GLMM) to examine how TMR conditions (Go-861 cued, Go-uncued, and NoGo-uncued) influenced participants' endorsement for positive traits 862 across time (baseline, post-TMR and delay). We used baseline endorsement rating as a covariate 863 and participant as random effect. The model was defined as:

864 Endorsement choice (Yes/No) ~ 1 + Baseline endorsement rating + Time × TMR
 865 Condition + (1+ TMR Condition + Time |Subject ID)

Subsequently, we employed a linear mixed model (LMM), incorporating the same factors
as used in the preceding GLMM for binary choice outcomes to analyze RTs when endorsing
positive traits except that we removed TMR as random slope due to singular fitting:

RTs (via log-transformed) ~ 1 + Baseline endorsement rating + Time × TMR Condition +
 (1 + Time |Subject ID)

In addition, we also ran a GLMM to examine the endorsement changes for negative traits,
 using time as a fixed effect:

873 Endorsement choice (Yes/No) ~ 1 + Baseline endorsement rating + Time + (1 + Time |
874 Subject ID)

Lastly, in order to assess whether CAT alone influenced the response speed during the endorsement of positive traits, we employed another LMM in two additional behavioral samples. These samples comprised one group that underwent only CAT training (referred to as the 'active' group) and another group that received no CAT training (referred to as the 'passive' group). Detailed information on these two behavioral samples can be found in the Supplementary Online Material (SOM). The LMM incorporated several fixed effects: group (active vs. passive), time (baseline, post-CAT, delay), and CAT (Go vs. NoGo). Additionally, the baseline endorsement

rating was included as a covariate. The model also accounted for random effects at theparticipant level:

RTs (via log-transformed) ~ 1 + Baseline endorsement rating + Group × Time × CAT
 Condition + (1 + Time |Subject ID)

886 Self-referential memories in the free recall task

To understand how TMR affect self-referential memories across time, we ran a GLMM using
TMR (Go-cued, Go-uncued, and NoGo-uncued), and time (baseline, post-CAT, post-TMR, and
delay) as fixed effects, baseline endorsement rating as covariate, participant as random effect.
Time was removed from random slope given singular fitting issue. The model was defined as
follows:

- Recall outcome (Yes/No) ~ 1 + Baseline endorsement rating + Time × TMR Condition +
 (1 + TMR Condition | Subject ID)
- Additionally, a GLMM was applied to analyze negative traits, using time as a fixed effect:
- 895 Recall outcome (Yes/No) ~ 1 + Baseline endorsement rating + Time + (1 | Subject ID)

Finally, incorporating data from two additional samples — one with only CAT training and another with no training (see SOM for details regarding behavioral samples)— we expanded our analysis to encompass three distinct groups. To assess delayed recall across these groups, we employed a GLMM on delayed recall performance with baseline and post-CAT recall as covariate, training groups (i.e., both TMR and CAT trained, only CAT trained, no CAT trained) as fixed effect:

- Recall outcome (Yes/No) ~ 1 + Baseline recall + Post-CAT recall + Baseline
 endorsement rating + Group + (1 + Group | Subject ID)
- 904 **EEG data analysis**
- 905 EEG data pre-processing

- 906 Continuous EEGs were recorded using a 63-channel customized cap with passive Ag/AgCl
- 907 electrodes via a BrainAmp amplifier with a 1000 Hz sampling rate (Brain Products, Gilching,
- 908 Germany). Electrodes were positioned according to the International 10–10 system. The ground
- 909 electrode was located at AFz, with FCz as the on-line reference electrode. Impedances were kept
- 910 below 20 k Ω . We placed one electro-oculography (EOG) electrode under participants' left eyes
- and bipolar electromyography (EMG) electrodes on their chins to monitor eye movements and
- 912 muscle activity during sleep.
- 913 EEG data were pre-processed using custom-written scripts and the MATLAB Toolbox
- EEGLAB (Delorme & Makeig, 2004). First, nap EEG data were down-sampled to 250 Hz,
- 915 notch-filtered at 50 Hz, and then re-referenced to the averaged mastoids. Second, EEG data were
- band-pass filtered at 0.5 to 40 Hz. While EOG and EMG data were used for sleep staging, these
- 917 data were not used in the time-frequency analysis.

918 Offline Sleep Stage Scoring

919 Sleep stages, including N1, N2, Slow-Wave Sleep (SWS), and Rapid Eye Movement (REM),

920 were scored using EEG (Channel C4), EOG, and EMG patterns. This process employed

algorithms from the YASA open-source Python Toolbox (Vallat & Walker, 2021). Consistent

922 with YASA guidelines, the EEG data were initially re-referenced to FPz before conducting the

923 staging analysis. Table 1 presents the sleep staging results for 34 participants (One participants

924 only reserved 28 min EEG data including TMR stage).

925 **Table 1**

Total time	Wake	N1	N2	N3	REM
90.10	7.76	7.18	36.00	27.00	12.36
±	<u>+</u>	<u>+</u>	±	±	±
1.27	1.19	0.63	1.56	1.83	1.27

926 Sleep stages parameters (mean $\Box \pm \Box$ SEM, in minutes, N = 34).

927

928 EEG time-frequency analysis

Before analyzing cue-elicited time-frequency EEG power changes, the cue-elicited EEG data

930 were epoched into -1.5 to 5.5 second segments, relative to the onset of each cued trait word. This

931 long epoch ensured that we had enough edge artifact-free segments for each clean epoch to

assess TMR benefits (-1 to 3 seconds). Epochs with artefacts were visually inspected and

933 removed. Time-frequency decomposition was performed in the Fieldtrip open-

934 source MATLAB toolbox (Oostenveld et al., 2011). We used 3 to 10 cycles in a step of 0.5 Hz

Morlet wavelet and baseline corrected using z-transformation of all trials from -1 to -0.1 seconds

relative to the cue onset. Following previous sleep and TMR studies (Mölle et al., 2011; Wilhelm

et al., 2020; Xia et al., 2023; Züst et al., 2019), we calculated the mean EEG power over frontalcentral channel (F1/2, FC1/2, C1/2, Fz, Cz) to ensure the robustness of results. The calculated time-frequency decompositions were then down-sampled to 50 Hz. To investigate cue-elicited

940 EEG activity, we employed the rigorous cluster-based one-sample permutation t-test (cluster-

941 thresholding p at 0.001) to identify the significant cluster against zero across all participants in

942 the time-frequency domain (Maris & Oostenveld, 2007).

943 Traveling wave analysis

We employed a traveling wave analysis approach similar to that used by Alamia et al. 2023. This
involved calculating values for spontaneous slow backward (from the anterior to posterior
regions) and forward (from posterior to anterior regions) traveling waves using 2D Fast Fourier
Transform (FFT) on the time-electrode EEG signals. The power measured in the upper right and
lower right quadrants corresponded to the amount of backward and forward propagating waves,

respectively, as shown in Figure 4A. Specifically, we used EEG data from the interval after cue onset [0, 2000] ms during NREM stage 3 (based on YAS staging result) across midline positions

951 (POz, Pz, CPz, Cz, Fz, FPz) to create time-electrode EEG representations. To establish a baseline,

we shuffled the electrodes and repeated this process. For the slow wave range (1–4 Hz), we

953 identified the maximum values in the 2D-FFT spectra in both the actual data (*BW* and *FW*) and

954 the shuffled data (*BWss* and *FWss*). The magnitude of the backward and forward traveling waves,

955 expressed in decibels [dB], was calculated as:

956 Backward =
$$10 \times \log_{10} BW/BW_{ss}$$
; Forward = $10 \times \log_{10} FW/FW_{ss}$ (1)

957 Brain-behavior association analysis

958 To establish a direct link between TMR-induced behavioral changes and TMR-elicited EEG 959 activity, we extracted the averaged power within the identified significant positive clusters, and 960 also calculated mean traveling waves for each participant at each item level. Then we performed 961 a series of LMMs using EEG power and traveling waves to predict post-TMR SRET 962 performance metrics, including endorsement choices and positive endorsement RTs. All EEG 963 metrics were centered before being included as fixed effects. We used GLMM to predict 964 endorsement choice (Yes/No) and LMMs to predict RTs for endorsing positive traits. The models were defined as: 965

966 (1) Post-TMR endorsement choice (Yes/No) ~ 1 + Positive delta-theta-alpha cluster/
967 Positive sigma-beta cluster/ Forward traveling wave / Backward traveling wave + Baseline
968 Choice + Repetition + (1|Subject ID).

969 (2) Delay endorsement choice (Yes/No) ~ 1 + Positive delta-theta-alpha cluster/ Positive
 970 sigma-beta cluster/ Forward traveling wave / Backward traveling wave + Baseline Choice +
 971 Repetition + (1|Subject ID).

972 (3) Post-TMR RTs for endorsing positive traits ~ 1 Positive delta-theta-alpha cluster/
973 Positive sigma-beta cluster/ Forward traveling wave / Backward traveling wave + Baseline RTs
974 + Repetition + (1|Subject ID).

- 975 (4) Delay RTs for endorsing positive traits ~ 1 + Positive delta-theta-alpha cluster/
- 976 Positive sigma-beta cluster/ Forward traveling wave / Backward traveling wave + Baseline RTs
- 977 + Repetition + (1|Subject ID).