

This is a self-archived version of an original article. This version may differ from the original in pagination and typographic details.

Author(s): Waselius, Tomi; Xu, Weiyong; Sparre, Julia Isabella; Penttonen, Markku; Nokia, Miriam S.

Title: -Cardiac cycle and respiration phase affect responses to the conditioned stimulus in young adults trained in trace eyeblink conditioning

Year: 2022

Version: Accepted version (Final draft)

Copyright: © 2022, Journal of Neurophysiology

Rights: In Copyright

Rights url: http://rightsstatements.org/page/InC/1.0/?language=en

Please cite the original version:

Waselius, T., Xu, W., Sparre, J. I., Penttonen, M., & Nokia, M. S. (2022). -Cardiac cycle and respiration phase affect responses to the conditioned stimulus in young adults trained in trace eyeblink conditioning. Journal of Neurophysiology, 127(3), 767-775. https://doi.org/10.1152/jn.00298.2021

- 1 Cardiac cycle and respiration phase affect responses to the conditioned
- 2 stimulus in young adults trained in trace eyeblink conditioning
- 3 Authors: Tomi Waselius[#], Weiyong Xu, Julia Sparre, Markku Penttonen & Miriam S. Nokia[#]
- 4 Affiliations for all authors: Department of Psychology, University of Jyvaskyla, Finland and
- 5 Centre for Interdisciplinary Brain Research, University of Jyvaskyla, Finland
- 6 Running head: Bodily rhythms affect eyeblink classical conditioning
- 7 Address for corresponding author: Tomi Waselius, tomi.waselius@jyu.fi, Department of
- 8 Psychology, P.O. Box 35, 40014 University of Jyvaskyla, Finland
- 10 [#] T. W. and M. S. N. contributed equally to this work.

11

Abstract

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

30

31

32

33

Rhythms of breathing and heartbeat are linked to each other as well as to rhythms of the brain. Our recent studies suggest that presenting conditioned stimulus during expiration or during the diastolic phase of the cardiac cycle facilitates neural processing of that stimulus and improves learning in a conditioning task. To date, it has not been examined whether utilizing information from both respiration and cardiac cycle phases simultaneously allows even more efficient modulation of learning. Here we studied whether the timing of the conditioned stimulus to different cardiorespiratory rhythm phase combinations affects learning in a conditioning task in healthy young adults. The results were consistent with previous reports: Timing the conditioned stimulus to diastole during expiration was more beneficial for learning than timing it to systole during inspiration. Cardiac cycle phase seemed to explain most of this variation in learning at the behavioral level. Brain evoked potentials (N1) elicited by the conditioned stimulus and recorded using electroencephalogram were larger when the conditioned stimulus was presented to diastole during expiration than when it was presented to systole during inspiration. Breathing phase explained the variation in the N1 amplitude. To conclude, our findings suggest that non-invasive monitoring of bodily rhythms combined with closed-loop control of stimulation can be used to promote learning in humans. The next step will be to test if performance can also be improved in humans with compromised cognitive ability, such as in older people with memory impairments.

Keywords

34 breathing, heartbeat, event-related potential, learning

New & Noteworthy

- We report for the first time that the rhythms of breathing and the beating of the heart have a
- 38 phase combination that is indicative of a neural state beneficial for cognition. This suggests
- 39 that bodily rhythms not only modulate cognition but that this phenomenon can be non-
- 40 invasively harnessed to improve learning in humans.

41

Introduction

42

43 Bodily rhythms like cardiac cycle and respiration usually vary at their own pace without 44 much conscious thought put into them. When we are in a relaxed psychophysiological state, 45 breathing and heartbeats synchronize [1-3]. The beat-to-beat intervals of the heart are longest 46 at the end of expiration [4] and shorter during inspiration [5]. This is called respiratory sinus 47 arrhythmia (RSA). In addition, the last heartbeat of each expiration delays the onset of the 48 following inspiration, the "working phase" of breathing, so that the cardiac "working phase", 49 systole, precedes inspiration onset by 150–500 ms [6,7]. This phenomenon is termed 50 cardioventilatory coupling. Especially RSA is thought to increase efficiency and stability in 51 pulmonary gas exchange [2] and to decrease energy use caused by heartbeats [8]. 52 Nevertheless, the root cause and outcome of cardiorespiratory synchrony is somewhat unclear 53 [6]. 54 Interestingly, cardiac cycle and respiration also synchronize with electrophysiological 55 rhythms of the brain [9], which in turn affect neural processing of external information. 56 Temporal correlations exist, for example, between cardiac cycle and the hippocampal theta 57 oscillation (3–12 Hz) in rodents [10,11]. Theta oscillation is crucially involved in memory 58 formation during spatial [12] and non-spatial tasks [13,14] as it, for example, regulates the 59 firing of hippocampal principal cells. In addition to cardiac cycle, also the respiration rhythm 60 couples with brain oscillations [15]. For example, electrophysiological oscillations in the 61 limbic system are entrained by nasal breathing in humans at the delta (0.5–4 Hz), theta (4–8 62 Hz) and beta (13–30 Hz) frequency bands [16]. In addition, hippocampal sharp-wave ripples 63 (SWRs, 100–200 Hz), crucial for memory consolidation [17], are entrained by respiration in 64 mice [18].

Further and most importantly, brain responses to external stimuli and consequent behavior
such as startle eyeblinks and premotor reaction times [19,20] and even associative learning
[21] are modulated by the cardiac cycle phase. Regarding respiration, associative learning is
enhanced when the significant stimuli are presented during expiration [22]. However, to our
knowledge, there are no studies considering the combined effect of cardiac cycle and
respiration phases on brain responses nor behavior. It is possible that utilizing combined
information from these bodily rhythms might allow even more efficient modulation of
behavior to the desired direction. Hence, we investigated the combined effect of cardiac cycle
and respiration phase on learning in an associative task called trace eyeblink conditioning
(TEBC). Participants were trained while watching a documentary film, using a tone as a
conditioned stimulus (CS) and an air puff towards the corner of the right eye as an
unconditioned stimulus. The presentation of the CS was fixed to a certain phase of the cardiac
cycle (systole, SYS or diastole, DIA) and respiration (expiration, EXP or inspiration, INS) for
each participant. In addition to conditioned responses also electroencephalogram (EEG) was
recorded. Our previous data [21] showed mixed effects of cardiac cycle phase on CS-evoked
brain responses in humans and in rabbits while we have not examined the effect of respiration
phase. Based on our previous behavioral results [21, 22], we hypothesized that timing the CS
to systole during inspiration would be less than optimal for learning TEBC whereas
presentation of the CS to the diastole during expiration would be most favorable for learning
(see Figure 1B).

Materials and Methods

Participants

Participants were recruited via student email lists. All participants gave informed written consent to this study and were free to discontinue participation in the experiment at any point. All participants received a reward (a movie ticket or a gift card) even if they discontinued the experiment at some point (however, no one did). The study was approved by the University of Jyväskylä Ethical Committee. A total of 59 young adults (12 males; aged 20–30 years: mean 23.4 years, standard error of mean 0.4 years) took part in the study. All participants were healthy with no history of psychiatric or neurological illnesses. They were not taking medication affecting the central nervous system, and they had no disabilities in hearing or vision.

97 Physiological recordings

Recording electrodes were attached after participants had signed the written consent.

Respiration was recorded and monitored during the experiment with a reusable fabric belt (RESPA00000, Spes Medica, Italy), which was fastened on top of the clothes on the lower chest area. Heart rate was recorded using three electrocardiogram (ECG) electrodes (Kendall, H92SG); one electrode was placed on top of the right clavicle, one on the left lower ribs, and the grounding electrode on the back of the neck. Electromyography (EMG) to determine eyeblinks was recorded using two electrodes (70010-K/12, Ambu, Ballerup, Denmark) that were attached on top of the participant's right eye muscles (orbicularis oculi). EEG was recorded using a 128-channel EGI Sensor Net (Electrical Geodesics Inc., Hydrogel GSN 128, 1.0). All signals were high-pass filtered (0.16 Hz) and low-pass filtered (250 Hz) online and

108 recorded with NeurOne Tesla (with Analog Out Option, Bittium Biosignals Ltd., Finland) at 109 a 1-kHz sampling rate. 110 Experimental procedure 111 We chose to use a between-subjects design with four groups (INS-SYS, INS-DIA, EXP-SYS, 112 and EXP-DIA) to keep the paradigm simple and use just one CS and one US. We do 113 acknowledge that a within-subjects design would be more powerful as there is quite a lot of 114 inter-individual variability in TEBC, and this is something that should be addressed in 115 follow-up studies. 116 The outline of the experimental procedure is presented in Figure 1A. The participants sat in a 117 chair in front of a TV screen (Asus VG236 series H, 23"; distance: approximately 100 cm). 118 They were informed that the aim of the study was to record physiological responses to 119 different types of stimuli while their attention was to be directed at a silent film depicting 120 landscapes and animals. The participants were instructed to pay attention to the film and told 121 that there would be questions considering the content of the footage after the recording 122 session. They were also instructed to sit comfortably in the chair and not pay attention to the 123 disturbing stimuli. In other words, the participants were led to believe that the idea was to 124 study the disturbance caused by beeping sounds and air puffs on their attention towards the 125 film. 126 Trace eyeblink conditioning 127 The conditioned stimulus (CS) was a 200-ms, 440-Hz, 66-dB tone delivered via a 128 loudspeaker situated in the lower right-hand corner of the room. The unconditioned stimulus 129 (US) was an air puff (0.2 bar source pressure, 100 ms) targeted at the right eye and it was

delivered via a plastic tube attached to modified safety goggles. Note that the air pressure was

131 low and none of the participants reported that the air puff hitting the eye was unbearable. 132 During conditioning trials, a 600-ms trace interval separated the tone-CS offset and the 133 airpuff-US onset. The presentation of the stimuli used for conditioning was controlled by 134 custom software running on an Arduino-based device (ABD). 135 First, five US-alone trials with an inter-trial interval (ITI) of 5 s were presented to make sure 136 that the participants felt comfortable enough to proceed with the experiment. After this, 5 137 minutes of resting data were recorded, followed by five CS-alone trials to determine baseline 138 eyeblink rate. Then, 50 CS+US classical conditioning trials were presented either at 139 inspiration-systole, inspiration-diastole, expiration-systole or at expiration-diastole. Last, five 140 CS-alone trials were presented as an extinction training block. A random ITI of 20-40 s was 141 applied throughout the experiment. 142 To time the classical conditioning trials, the respiration, cardiac cycle, and EMG signals were 143 conveyed to a custom script running in LabVIEW (National Instruments). Signals were 144 sampled at 1 kHz. At each time point, the last second of respiration, ECG and EMG signals 145 were analyzed. EMG was evaluated for spontaneous eyeblinks, that is, the signal had to stay 146 below a set amplitude threshold to proceed with presenting the conditioning trial. The 147 respiration signal was analyzed in two consecutive 500-ms windows. To trigger a trial, the 148 signal amplitude during the latter 500-ms time window had to cross a set absolute threshold 149 value (peak for inspiration, trough for expiration) and the signal had to either rise 150 (inspiration) or fall (expiration) at a certain rate between the two consecutive time windows. 151 In addition, R-peaks were detected from the ECG and used for timing the trial either at 152 systole (immediately) or diastole (delayed from R-peak). Note that the threshold values for 153 the EMG and for the respiration peak and rise (inspiration) and for the trough and fall 154 (expiration) were set individually for each participant during the 5-minute baseline recording

prior to conditioning. As a result, when the participant was not spontaneously blinking, and respiration and cardiac cycle were at desired phases, LabVIEW sent a TTL pulse to the ABD, which then presented the actual conditioning stimuli. In addition, whenever voluntary movement of the participant was visible either in the monitoring video or in the breathing signal, the trial presentation was manually halted. Also, any changes in the overall breathing baseline were taken into account and ABD controlling LabView parameters were adjusted accordingly.

Two minutes of spontaneous breathing and ECG without any external stimuli were recorded after the conditioning session to visually confirm online that the experimental manipulation had not changed the respiration pattern overall and that the respiration belt signal quality had remained similar to that recorded before experimental manipulations. The whole procedure lasted about 40 minutes depending on the random ITI.

Questionnaire

After the experiment, participants answered background questions about age, sex, and handedness and five questions concerning the silent film (e.g., "What equipment did the man in the film use for travelling in the snow?") and an open question about the disruptive stimuli. Questions about the film were asked to find out if participants had been concentrating on the film because attention has a serious impact on learning in classical conditioning [23]. Participants also answered seven true/false questions about the occurrence of the disruptive stimuli (e.g., air puffs occurred immediately after beeping sounds). These questions were asked to find out how conscious the participants became of the CS-US association. For the complete questionnaire (translated into English for reporting purposes), please see Appendix 1.

178 Data analysis

Conditioned responses

The conditioned responses (CR) performed by each participant were analyzed offline using MATLAB (The MathWorks Inc.). First, the EMG signal was low pass filtered (40 Hz) and the absolute value of the signal was derived. Then the mean amplitude of the rectified EMG signal during a 500-ms pre-US period (MEANpre) was calculated. In addition, the mean of the standard deviation of the signal amplitude during the 500-ms pre-CS period (SDpre) was determined. Learned responses were detected from a 200-ms time window immediately preceding the US. To qualify as a learned response, the rectified EMG signal amplitude had to exceed the following threshold: MEANpre + 2 * SDpre. For statistical analysis, trials were grouped into blocks of five trials and the proportion (%) of conditioned responses per block was calculated. These measures were used as dependent variables when analyzing learning. Further, to create a simple measure of the outcome of TEBC, we determined the highest proportion (%) of conditioned responses during any given 10-trial block (50 trials, 5 blocks). This measure is referred to as the best performance in the TEBC task.

Event-related potentials (ERPs)

EEG data were analyzed using the MNE python [24]. First, EEG channels were visually inspected and bad channels were interpolated using spherical spline interpolation method [25]. Then fast independent component analysis (ICA) was applied to remove any eyeblink and cardiac artifacts related components [26]. Our previous study has shown that after applying ICA to remove cardiac related components, sometimes also referred to as heart-evoked potentials, the cardiac-related signal in EEG is virtually flat (see Figure 4 in Waselius et al. 2018). Then a band-pass filter of 0.1–30 Hz (zero phase finite impulse response filter with a Hamming-window) was applied to the continuous EEG recordings. After filtering, the

EEG signal was re-referenced to the common average. Then the EEG data were segmented into epochs spanning from −100 to 500 ms relative to the onset of the CS. The EEG epochs were manually checked to exclude any trials that were contaminated by movement-related artifacts or other high-amplitude noise. EEG epochs exceeding 100 μV peak-to-peak amplitudes were excluded from further analysis. Finally, the event-related potentials (ERP) were obtained by averaging EEG epochs around the CS over all paired conditioning trials.

Next, the ERP data were grand averaged across all participants. Two major ERP components were evident: An auditory N1, which peaked around 112 ms, and auditory P2, which peaked around 189 ms after the onset of the CS. In addition, the center of activities for both N1 and P2 peaks were around the channels number 6, 7 and 106 (128-channel EGI Sensor Net), which are located around the center of the head (see Figure 3A). This pattern (vertex negative-positive potentials) is consistent with our previous study [21] and other studies [27, 28]. Based on this, auditory N1 and P2 mean amplitudes were extracted from each participant for further statistical analysis from a 30-ms time window around the grand average N1 (112 ms) and P2 (189 ms) peaks from channels number 6, 7 and 106.

Statistics

One way analysis of variance (ANOVA) and independent samples t-test were used to examine differences between groups in single variables. Repeated-measures (rm) ANOVA was used to analyze changes across training and differences between the groups in conditioned responding: Five-trial averages (blocks, 10) were used as a within-subjects factor and respiration phase (2) and cardiac cycle phase (2) as between-subjects factors. Univariate ANOVA was used to examine the effects of respiration phase (2) and cardiac cycle phase (2) on single variables: best performance, N1 and P2 amplitude. Cohen's d or partial eta squared (η 2) are reported for statistically significant differences.

226 Results

227

241

242

243

244

228 Of the 59 participants, 56 answered correctly to all the questions about the film content 229 (questions 1–5 of Appendix 1) and the rest of them had only one missing answer. 230 Respectively, only 27 of 59 participants answered correctly to the questions about how the 231 disruptive stimuli were presented (questions 7–13 of Appendix 1). This indicates that the 232 participants were generally well concentrated on watching the film and not on the 233 conditioning stimuli. 234 Participants trained during expiration-diastole made more conditioned responses than those 235 trained during inspiration-systole 236 Fifty-one out of 52 participants made conditioned responses at some point during the TEBC 237 and were included in the analyses (EXP-DIA: n = 13, EXP-SYS: n = 13, INS-DIA: n = 12, 238 INS-SYS: n = 13) (see Figure 2). Participants in all groups responded (i.e., blinked their eye) 239 at an equal rate (mean \pm standard error of mean: 10 % \pm 2 percentage units) to the CS during 240 the CS-alone trials (one way ANOVA: F [3, 47] = 1.146, p = 0.340). To test the effects of

Participants concentrated on watching the documentary film

245 423] = 8.051, p < 0.001, η_p^2 = 0.146) a statistically significant main effect of cardiac cycle

phase (2) and cardiac cycle phase (2) as between-subjects factors and block (10) as the

within-subjects factor. In addition to the statistically significant main effect of block (F [9,

breathing (EXP: n = 26 vs. INS: n = 25) and cardiac cycle phase (DIA: n = 25 vs. SYS: n = 25)

26) on TEBC, we analyzed the conditioned response data with rm ANOVA using respiration

- 246 phase (F [1, 47] = 6.109, p = 0.017, $\eta_p^2 = 0.115$) was detected. Interactions were not
- 247 statistically significant (within-subjects: F [9, 423] = 0.434–0.963, p = 0.461–0.889; between
- subjects: F[1, 47] = 0.471, p = 0.496) nor was the main effect of breathing phase (F[1, 47] =

- 0.852, p = 0.361). Next, to test our hypothesis directly, an independent samples t-test was
- used to analyze the difference in conditioned responses to CS during all blocks (10, average)
- between EXP-DIA and INS-SYS. Conditioned responding was higher in the EXP-DIA (63 %
- \pm 18 percentage units) than in the INS-SYS group (44 % \pm 19 percentage units), t (24) =
- 253 2.588, p = 0.016, Cohen's d 1.015.
- To further analyze the outcome of TEBC we determined a measure of best performance for
- each participant as the highest proportion (%) of conditioned responses during any given 10-
- trial block. Univariate ANOVA indicated a significant difference in best performance
- between participants trained at systole vs. diastole (cardiac cycle phase: F [1, 47] = 7.667, p =
- 258 0.008, $\eta_p^2 = 0.140$; respiration phase, F [1, 47] = 0.357, p = 0.553; interaction, F [1, 47] =
- 0.008, p = 0.930). To directly test our hypothesis, we performed a comparison between just
- 260 the EXP-DIA (82 % \pm 17 percentage units) and the INS-SYS (62 % \pm 22 percentage units)
- 261 groups using independent samples t-test which indicated a significant difference: t (24) =
- 262 2.734, p = 0.012, Cohen's d = 1.073.
- 263 To summarize, participants in all groups readily acquired the conditioned response and those
- trained during diastole made more conditioned responses than those trained at systole.
- Specifically, and in accordance with our hypothesis, participants trained during expiration-
- diastole made more conditioned responses than those trained during inspiration-systole.
- 267 The conditioned stimulus evoked a larger N1 response in participants trained during
- 268 expiration than in those trained during inspiration
- 269 High-quality EEG data were recorded from 40 participants with valid behavioral data (10 in
- each group, see Figure 3). In analyzing the EEG data, we followed the same logic as for the
- 271 conditioned responses: Univariate ANOVA revealed a significant effect of respiration phase

on the N1 amplitude (F [3, 36] = 12.219, p = 0.001, η_p^2 = 0.253; cardiac cycle phase: F [3, 272 273 36] = 0.632, p = 0.432; interaction: F [3, 36] = 0.737, p = 0.396) but not on the P2 amplitude 274 (F [3, 36] = 0.567, p = 0.456; cardiac cycle phase: F [3, 36] = 3.144, p = 0.085; interaction: F[3, 36] = 0.698, p = 0.409). To follow up on our direct hypothesis of better learning in the 275 276 expiration-diastole group compared to inspiration-diastole group, we performed independent 277 samples t-test on the N1 and P2 amplitudes. The N1 amplitude was larger in the EXP-DIA 278 compared to the INS-SYS group (t [18] = 2.766, p = 0.013, Cohen's d 0.135) but there was 279 no difference in P2 amplitude (t [18] = 1.626, p = 0.121). To conclude, N1 responses were 280 largest in the EXP-DIA group and overall larger N1 responses were evoked when the CS was 281 presented during expiration rather than during inspiration.

282

Discussion

283

284

285

286

287

288

289

290

291

292

293

294

295

296

297

298

299

300

301

302

303

304

305

306

Respiration rhythm and cardiac cycle are known to synchronize to each other [6], to modulate brain activity [11, 15] and to affect, for example, perception and learning [29, 30]. However, it is unknown whether combinations of respiration and cardiac cycle phases modulate learning. Here, healthy young adults were trained in trace eyeblink classical conditioning, timing the conditioned stimulus based on four combinations of respiration and cardiac cycle phases (inspiration-systole, inspiration-diastole, expiration-systole, expiration-diastole; see Figure 1). Based on results of our previous studies [21, 22], we assumed that the diastolic phase during expiration would be a beneficial phase for stimulus presentation when learning, whereas systolic phase during inspiration would be less beneficial for learning. As expected, timing the CS onset to diastole during expiration resulted in more frequent conditioned responding compared to timing the CS onset to systole during inspiration. Further, conditioned responding was overall more frequent if the CS was timed to diastole than to systole. Parallel differences were also observed in electrophysiological brain responses evoked by the CS: The N1 response was larger in amplitude when the CS occurred during expiration, and especially when it occurred during expiration and diastole. Together these results support our main assumption and our previous findings [21, 22] that bodily rhythms can be used to facilitate learning in humans. Most importantly, our current study indicates that learned behavior can be modulated by the combinatory phases of breathing and the cardiac cycle. Overall, participants in our study acquired the conditioned eyeblink very fast, within the first few training blocks. As hypothesized, in our participants trained exclusively during the "resting states" of the heart and respiratory muscles (expiration-diastole), performance of a learned motor response was more likely compared to that in participants trained in the "working phase" of these organs

(inspiration-systole). Further, the phase of the cardiac cycle was the main factor explaining
this difference. This result is in contrast with our earlier finding indicating no effect of
cardiac cycle phase on learning in humans [21]. However, this could be explained by the
further development of the conditioning paradigm in terms of triggering the trials to systole
or diastole, which was more accurate in the current experiment. Namely, the delay from the
R-peak was individually adjusted to suit each participant's heart rate instead of using a set
delay for all participants. We also did not detect a main effect of breathing phase on
conditioned responding, again in contrast with our earlier finding [22]. However, it could be
that as in the current experiment the timing of the CS hinged on the R-peak, the phases of the
respiration (EXP vs. INS) are not directly comparable to those in our earlier study. Namely,
in our current study, the onset of the CS was delayed until the next heartbeat within the
expiration or the inspiration phase of breathing while in the Waselius et al. 2019 study a CS
was triggered immediately as the desired breathing phase was detected. In any case, putting
all evidence together, it seems that the neural state during diastole and expiration might be
most favorable for acquiring an auditory CS-somatosensory US association and then
performing a learned motor response. This conclusion is in line with all our findings, current
and previous [21, 22]. It is also in line with a report of faster reactions to and higher saliency
evaluations of auditory startle stimuli when presented during expiration rather than during
inspiration [31]. Further support comes from studies reporting greater startle eyeblink
responses to auditory stimuli presented at diastole than systole [19, 20].
Our current results suggest that respiration and cardiac cycle phases affect learning itself and
not just the performance of the conditioned response, as 1) there is no difference between
groups during the CS-alone treatment or the very first conditioning trials and 2) there is a
clear distinction in the probability of a conditioned eyeblink once it reaches a plateau (see
· · · · · · · · · · · · · · · · · · ·

Figure 2). According to Prokasy's theory (1984), during eyeblink conditioning the
participants first learn an association between the conditioned stimulus and the unconditioned
stimulus. Then they learn to shut their eye before the irritating air puff, that is, they learn to
perform the motor conditioned response. Over time, with extended training, the conditioned
eyeblink is adjusted temporally so that it optimally protects the eye from the flow of air [32].
Considering this, it seems that in our current experiment the effects of the neural state
indicated by the phases of the bodily rhythms center on the acquisition of the CS-US
contingency and the motor conditioned response taking place early in training and not so
much on the later phases of the process when the CR is further adjusted. Learning the CS-US
association during trace eyeblink conditioning is considered to be hippocampus-dependent
[33–35] because of the gap between the CS-offset and the US onset while the simpler version
of the task where the two stimuli partially overlap relies solely on the cerebellum responsible
for motor learning [36, 37]. Thus, it is possible that the neural state indicated by diastole
during expiration is related to a more efficient acquisition of the CS-US association, perhaps
involving the hippocampus, and to a more reliable execution of the conditioned motor
response governed by the cerebellum.
As anticipated based on the behavioral results, electrophysiological brain responses evoked
by the conditioned stimulus also differed between the experimental groups in our study.
Specifically, the N1 component of the ERP responses was largest in the participants trained
in diastole during expiration and an overall larger N1 was evoked when the CS was presented
during expiration rather than during inspiration (Figure 3). This suggests that the CS evoked
more synchronous neural activity and was possibly perceived as more surprising or salient
[38] during expiration (and diastole). It should be noted that earlier studies have not
addressed how respiration might modulate auditory ERPs, but it has been suggested that

auditory startle stimuli are subjectively rated more intense if presented to mid-expiration [31]. Regarding the effects of cardiac cycle phase, in our earlier study [21] the N1 of the ERP was in fact higher in amplitude during systole than diastole. However, Schulz and colleagues (2020) found the N1 to be higher in amplitude during diastole than systole when studying the effects of the cardiac cycle phase on responses to auditory startle stimuli [20]. Clearly, the modulation of neural responses by bodily rhythms should be explored in more detail and considered in data analysis. One might claim, for example, that a fixed inter-stimulus interval could result in the stimulus being presented repeatedly in the same phase of respiration and/or cardiac cycle, depending on the rate. This might then affect the amplitude of the different ERP components. When it comes to the mechanism behind the link between bodily rhythm phases, brain function and behavior, not a lot is known. It does seem clear that the spontaneous rhythms of the brain relate to the rhythms of breathing and heartbeat [9], but a deeper mechanistic explanation of the anatomical and functional connections is missing. One of the most studied phenomena is the link between respiration, olfaction, and related brain activity. Several studies report that respiration rhythm and phase are connected to brain oscillatory activity [15,16,18,39–41]. In most studies, the connection seems to be limited to nasal respiration [16] and to crucially depend on the function of the olfactory bulb neurons [18]. Some of the respiration-driven brain rhythms might even be separate from the traditional brain rhythms, such as theta paced by subcortical structures [14]. Specifically, sniffing rodents are reported to display hippocampal oscillations that occur at the theta-frequency and couple with respiration but are not theta [39]. Since oscillations are the foundation of information transfer in the brain [42–44], it is obvious that rodents largely dependent on their olfactory senses could benefit from this coupling of respiration and brain oscillations [45]. As an

355

356

357

358

359

360

361

362

363

364

365

366

367

368

369

370

371

372

373

374

375

376

377

evolutionarily close relative to rodents, humans might also possess this characteristic.

Interestingly, olfaction and spatial memory seem to be congruent in humans: A good sense of smell is linked to good navigating ability [46]. Curiously, here and in our previous studies the "sniffing phase", meaning inspiration, has emerged as the *less* optimal phase for associative learning of a causal connection between an auditory cue and an aversive somatosensory stimulation [22]. This might be related to the observations that hippocampal sharp-wave ripples (SPW-Rs) known to be crucial for memory consolidation [47, 48] and reflect a state in which rabbits acquire eyeblink conditioning better [47, 48] are more likely to occur during expiration than inspiration in mice [18]. Whether hippocampal SPW-Rs are more likely to occur during expiration in the diastolic phase of the cardiac cycle in humans is not known and should be studied in the future. Further, it should be tested whether limiting breathing to the oral route would possibly abolish the link between cardioventilatory rhythms and conditioned responding [16]. In sum, we suggest that the findings of our current experiment could be in part explained by the link between bodily rhythms and brain oscillatory activity, especially by that between expiration and SPW-Rs.

Limitations

There are some limitations in this study that we want to point out. Most obvious is the somewhat small sample size and small effect sizes. Nevertheless, the effect sizes in the ERP and behavioral results were large in our previous studies with similar group sizes and experimental set-up. Thus, we were confident that a similar group size should be large enough to detect possible effects in our current study as well. In addition, one must remember that associative learning in TEBC is by no means representative of all learning. In fact, in our current follow-up studies we have employed a different type of associative memory task, more closely resembling real-life situations. Moreover, here bodily rhythms modulated

auditory stimulus processing in a way that might or [22] might not be similar to how these rhythms modulate processing of visual or somatosensory stimuli in different settings. Last, we have previously reported [22] that participants have high RSA during this kind of an experimental setup but we did not take into account if the cardioventilatory coupling was strong or not and whether it showed fluctuations during the experiment [4]. In future studies, it might be worth investigating if the level of cardioventilatory coupling has an additional effect on learning.

Conclusion

To summarize, this study is the first to demonstrate that both breathing, and heartbeat rhythms influence the brain processing of external stimuli and learning about those stimuli. Learned responding is more likely when an auditory conditioned stimulus is presented during the resting phase of the heartbeat when breathing out instead of during the working state of the heartbeat when breathing in. In addition, the N1 component of the auditory-evoked potential is larger when the stimulus is presented during expiration (and diastole) than when it occurs during inspiration (and systole). These findings suggest that non-invasive measurement of cardiorespiratory rhythms combined with closed-loop control of stimulation can be utilized to promote learning in humans. The next step will be to test if performance can also be improved in humans with compromised cognitive ability [49, 50]. Further, it will be interesting to test the effects of bodily rhythm phases on learning tasks more closely resembling real-life situations.

423 References

- 424 1. Brecher GA, Hubay CA. Pulmonary blood flow and venous return during spontaneous
- 425 respiration. Circ Res 3: 210–214, 1955.
- 426 2. Yasuma F, Hayano J-I. Respiratory sinus arrhythmia: why does the heartbeat synchronize with
- 427 respiratory rhythm? *Chest* 125: 683–690, 2004.
- 428 3. Ludwig C. Beitrage zur Kenntniss des Einflusses der Respirationsbewegungen auf den Blutlauf im
- 429 Aortensysteme. Arch Anat Physiol Leipzig 13: 242–302, 1847.
- 430 4. Galletly DC, Larsen PD. Relationship between cardioventilatory coupling and respiratory sinus
- 431 arrhythmia. *Br J Anaesth* 80: 164–168, 1998.
- 432 5. Hirsch JA, Bishop B. Respiratory sinus arrhythmia in humans: how breathing pattern modulates
- 433 heart rate. *Am J Physiol* 241: H620–9, 1981.
- 434 6. Elstad M, O'Callaghan EL, Smith AJ, Ben-Tal A, Ramchandra R. Cardiorespiratory
- 435 interactions in humans and animals: rhythms for life. Am J Physiol Heart Circ Physiol 315: H6–
- 436 H17, 2018.
- 437 7. Coleman WM. The psychological significance of bodily rhythms. *Journal of Comparative*
- 438 *Psychology* 1: 213–220, 1921. https://doi.org/10.1037/h0067228.
- 439 8. Grossman P, Taylor EW. Toward understanding respiratory sinus arrhythmia: relations to cardiac
- vagal tone, evolution and biobehavioral functions. *Biol Psychol* 74: 263–285, 2007.
- 441 9. Klimesch W. The frequency architecture of brain and brain body oscillations: an analysis. Eur J
- 442 Neurosci 48: 2431–2453, 2018.
- 443 10. Komisaruk BR. Synchrony between limbic system theta activity and rhythmical behavior in rats. J
- 444 *Comp Physiol Psychol* 70: 482–492, 1970.
- 445 11. Pedemonte M, Goldstein-Daruech N, Velluti RA. Temporal correlations between heart rate,
- medullary units and hippocampal theta rhythm in anesthetized, sleeping and awake guinea pigs.
- 447 Autonomic Neuroscience 107: 99–104, 2003. https://doi.org/10.1016/s1566-0702(03)00132-2.
- 448 12. O'Keefe J, Recce ML. Phase relationship between hippocampal place units and the EEG theta
- 449 rhythm. *Hippocampus* 3: 317–330, 1993.
- 450 13. Nokia MS, Waselius T, Mikkonen JE, Wikgren J, Penttonen M. Phase matters: responding to
- and learning about peripheral stimuli depends on hippocampal θ phase at stimulus onset. Learning
- 452 & Memory 22: 307–317, 2015. https://doi.org/10.1101/lm.038166.115.
- 453 14. **Buzsáki G**. Theta oscillations in the hippocampus. *Neuron* 33: 325–340, 2002.
- 454 15. Tort ABL, Brankačk J, Draguhn A. Respiration-Entrained Brain Rhythms Are Global but Often
- Overlooked. Trends in Neurosciences 41: 186–197, 2018.
- 456 https://doi.org/10.1016/j.tins.2018.01.007.
- 457 16. Zelano C, Jiang H, Zhou G, Arora N, Schuele S, Rosenow J, Gottfried JA. Nasal Respiration
- Entrains Human Limbic Oscillations and Modulates Cognitive Function. *J Neurosci* 36: 12448–
- 459 12467, 2016.

- 460 17. **Buzsáki G**. Hippocampal sharp wave-ripple: A cognitive biomarker for episodic memory and planning. *Hippocampus* 25: 1073–1188, 2015. https://doi.org/10.1002/hipo.22488.
- 462 18. **Liu Y**, **McAfee SS**, **Heck DH**. Hippocampal sharp-wave ripples in awake mice are entrained by respiration. *Sci Rep* 7: 8950, 2017.
- 464 19. Schulz A, Reichert CF, Richter S, Lass-Hennemann J, Blumenthal TD, Schächinger H.
- Cardiac modulation of startle: effects on eye blink and higher cognitive processing. *Brain Cogn* 71:
- 466 265–271, 2009.
- 467 20. Schulz A, Vögele C, Bertsch K, Bernard S, Münch EE, Hansen G, Naumann E, Schächinger
- 468 H. Cardiac cycle phases affect auditory-evoked potentials, startle eye blink and pre-motor reaction
- times in response to acoustic startle stimuli. *Int J Psychophysiol* 157: 70–81, 2020.
- 470 21. **Waselius T, Wikgren J, Halkola H, Penttonen M, Nokia MS**. Learning by heart: cardiac cycle reveals an effective time window for learning. *J Neurophysiol* 120: 830–838, 2018.
- 472 22. Waselius T, Wikgren J, Penttonen M, Nokia MS. Breathe out and learn: Expiration-contingent
- stimulus presentation facilitates associative learning in trace eyeblink conditioning.
- 474 *Psychophysiology*: e13387, 2019. https://doi.org/10.1111/psyp.13387.
- 475 23. Clark RE, Manns JR, Squire LR. Classical conditioning, awareness, and brain systems. *Trends Cogn Sci* 6: 524–531, 2002.
- 477 24. Gramfort A, Luessi M, Larson E, Engemann DA, Strohmeier D, Brodbeck C, Goj R, Jas M,
- 478 Brooks T, Parkkonen L, Hämäläinen M. MEG and EEG data analysis with MNE-Python. Front
- 479 Neurosci 7: 267, 2013.
- 480 25. **Perrin F, Pernier J, Bertrand O, Echallier JF**. Spherical splines for scalp potential and current density mapping. *Electroencephalogr Clin Neurophysiol* 72: 184–187, 1989.
- 482 26. **Hyvärinen A**. Fast and robust fixed-point algorithms for independent component analysis. *IEEE Trans Neural Netw* 10: 626–634, 1999.
- 484 27. **Carpenter AL**, **Shahin AJ**. Development of the N1–P2 auditory evoked response to amplitude rise time and rate of formant transition of speech sounds. *Neurosci Lett* 544: 56–61, 2013.
- 486 28. Näätänen R, Sams M, Alho K, Paavilainen P, Reinikainen K, Sokolov EN. Frequency and
- location specificify of the human vertex N1 wave. Electroencephalogr Clin Neurophysiol 69: 523–
- 488 531, 1988.
- 489 29. Park H-D, Correia S, Ducorps A, Tallon-Baudry C. Spontaneous fluctuations in neural
- responses to heartbeats predict visual detection. *Nat Neurosci* 17: 612–618, 2014.
- 491 30. Perl O, Ravia A, Rubinson M, Eisen A, Soroka T, Mor N, Secundo L, Sobel N. Human non-
- olfactory cognition phase-locked with inhalation. *Nat Hum Behav* 3: 501–512, 2019.
- 493 31. Münch EE, Vögele C, Van Diest I, Schulz A. Respiratory modulation of intensity ratings and
- psychomotor response times to acoustic startle stimuli. *Neurosci Lett* 711: 134388, 2019.
- 495 32. Prokasy WF. Acquisition of Skeletal Conditioned Responses in Pavlovian Conditioning.
- 496 *Psychophysiology* 21: 1–13, 1984.
- 497 33. Kim JJ, Clark RE, Thompson RF. Hippocampectomy impairs the memory of recently, but not
- remotely, acquired trace eyeblink conditioned responses. *Behavioral Neuroscience* 109: 195–203,
- 499 1995.

- 500 34. **Moyer JR Jr**, **Deyo RA**, **Disterhoft JF**. Hippocampectomy disrupts trace eye-blink conditioning in rabbits. *Behav Neurosci* 104: 243–252, 1990.
- 502 35. Solomon PR, Vander Schaaf ER, Thompson RF, Weisz DJ. Hippocampus and trace
- conditioning of the rabbit's classically conditioned nictitating membrane response. Behav Neurosci
- 504 100: 729–744, 1986.
- 505 36. **Medina JF**, **Nores WL**, **Ohyama T**, **Mauk MD**. Mechanisms of cerebellar learning suggested by eyelid conditioning. *Curr Opin Neurobiol* 10: 717–724, 2000.
- 507 37. **Thompson RF**, **Steinmetz JE**. The role of the cerebellum in classical conditioning of discrete behavioral responses. *Neuroscience* 162: 732–755, 2009.
- 509 38. **Friston K**. The free-energy principle: a rough guide to the brain? *Trends Cogn Sci* 13: 293–301, 2009.
- 511 39. Nguyen Chi V, Müller C, Wolfenstetter T, Yanovsky Y, Draguhn A, Tort ABL, Brankačk J.
- Hippocampal Respiration-Driven Rhythm Distinct from Theta Oscillations in Awake Mice. J
- 513 *Neurosci* 36: 162–177, 2016.
- 514 40. Ito J, Roy S, Liu Y, Cao Y, Fletcher M, Lu L, Boughter JD, Grün S, Heck DH. Whisker barrel
- 515 cortex delta oscillations and gamma power in the awake mouse are linked to respiration. *Nat*
- 516 *Commun* 5: 3572, 2014.
- 517 41. Karalis N, Sirota A. Breathing coordinates limbic network dynamics underlying memory
- 518 consolidation. 2018. doi: http://dx.doi.org/10.1101/392530, 2018.
- 519 42. Buzsáki G. Neural syntax: cell assemblies, synapsembles, and readers. Neuron 68: 362–385, 2010.
- 520 43. **Fell J, Axmacher N**. The role of phase synchronization in memory processes. *Nat Rev Neurosci* 12: 105–118, 2011.
- 522 44. Fries P. A mechanism for cognitive dynamics: neuronal communication through neuronal
- 523 coherence. *Trends Cogn Sci* 9: 474–480, 2005.
- 524 45. Tsanov M, Chah E, Reilly R, O'Mara SM. Respiratory cycle entrainment of septal neurons
- mediates the fast coupling of sniffing rate and hippocampal theta rhythm. Eur J Neurosci 39: 957–
- 526 974, 2014.
- 527 46. Dahmani L, Patel RM, Yang Y, Chakravarty MM, Fellows LK, Bohbot VD. An intrinsic
- association between olfactory identification and spatial memory in humans. *Nat Commun* 9: 4162,
- 529 2018.
- 530 47. Nokia MS, Mikkonen JE, Penttonen M, Wikgren J. Disrupting neural activity related to awake-
- state sharp wave-ripple complexes prevents hippocampal learning. Frontiers in Behavioral
- 532 *Neuroscience* 6: 2012. https://doi.org/10.3389/fnbeh.2012.00084.
- 533 48. Nokia MS, Penttonen M, Wikgren J. Hippocampal ripple-contingent training accelerates trace
- eyeblink conditioning and retards extinction in rabbits. *J Neurosci* 30: 11486–11492, 2010.
- 535 49. Jacobson SW, Stanton ME, Dodge NC, Pienaar M, Fuller DS, Molteno CD, Meintjes EM,
- Hoyme HE, Robinson LK, Khaole N, Jacobson JL. Impaired delay and trace eyeblink
- 537 conditioning in school-age children with fetal alcohol syndrome. Alcohol Clin Exp Res 35: 250–
- 538 264, 2011.

539 50. **Woodruff-Pak DS**. Eyeblink classical conditioning differentiates normal aging from Alzheimer's disease. *Integr Physiol Behav Sci* 36: 87–108, 2001.

Figure captions

541

542

543

544

545

546

547

548

549

550

551

552

553

554

555

556

557

558

559

560

561

562

563

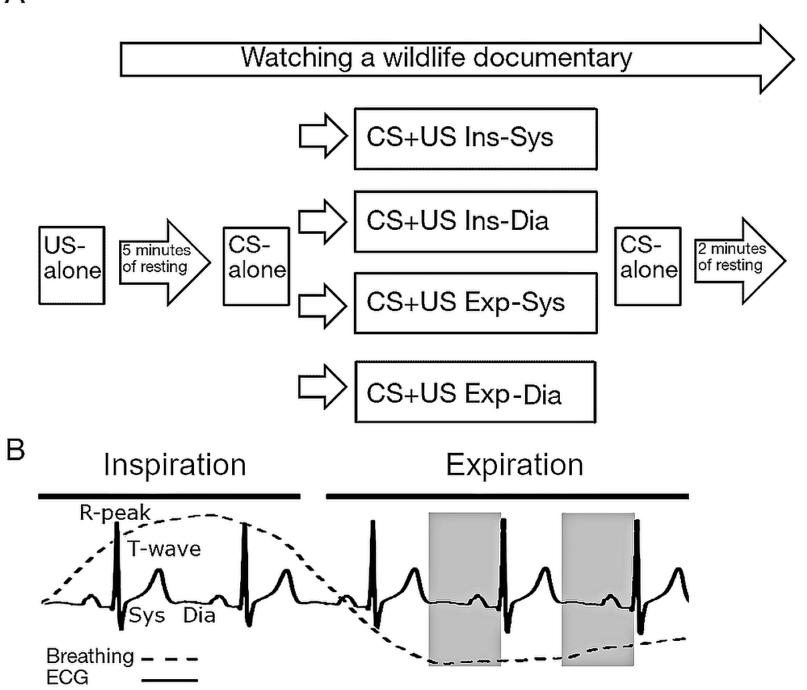
564

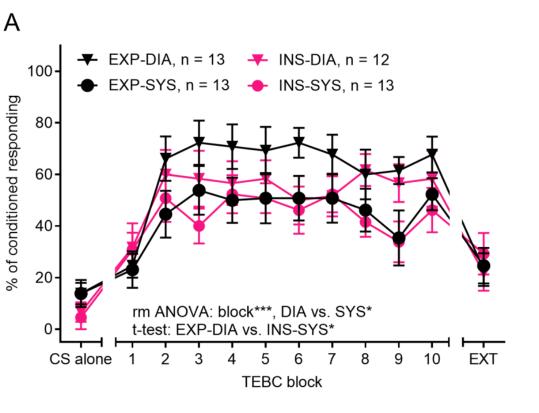
Figure 1. Experimental design and examples of breathing and ECG signals used for timing the conditioning trials. (A) After presenting five US-alone trials the participants started to watch a documentary film. Then there was a five-minute resting period followed by five CS-alone trials. Next, 50 pairs of CS+US trials were timed to a certain cardiorespiratory phase (Ins-Sys, Ins-Dia, Exp-Sys or Exp-Dia). At last, five CS-alone trials were presented as an extinction training (EXT) followed by a two-minute rest period before the experiment ended. (B) Breathing signal and ECG were recorded and followed online to time conditioning trials to a certain cardiorespiratory phase. During the systolic phase (Sys) of the cardiac cycle, the ECG shows the QRS complex, reflecting ventricular depolarization, and the T wave, reflecting ventricular repolarization. Between the end of the T-wave and next R-peak, is the diastolic phase (Dia). The diastolic phase during expiration is marked with grey bars for demonstrating the hypothesized optimal phases for stimulus presentation. Figure 2. Participants trained during expiration-diastole (EXP-DIA) made more conditioned responses than those trained during inspiration-systole (INS-SYS). (A) The percentage of conditioned responses per 5-trial block was used as a measure of learned behavior. There was no difference in responding to the tone-CS prior to conditioning (CS alone, one way ANOVA). Participants in all groups learned the trace eyeblink conditioning task (TEBC, main effect of block) and participants trained in diastole made more conditioned responses than those trained in systole (rm ANOVA, main effect of cardiac cycle). Further, in accordance with our hypothesis, participants trained at EXP-DIA made more conditioned responses during TEBC than those trained at INS-SYS (independent samples t-test). (B) Conditioned responding at best was higher in participants trained at diastole than at systole

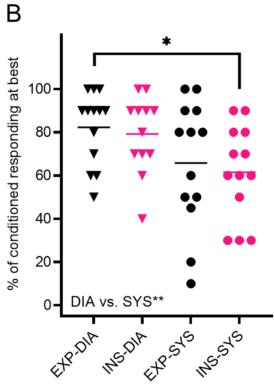
(univariate ANOVA) and higher in participants trained at EXP-DIA than those trained at

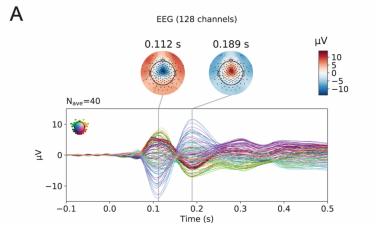
566 *	
900 *	* p < 0.010, *** p \leq 0.001. Vertical lines in panel A indicate standard error of mean.
567 H	Iorizontal lines in panel B refer to the mean.
568 F	Figure 3. The conditioned stimulus evoked a larger N1 response in participants trained
569 d	during expiration than in those trained during inspiration. (A) Joint plot of the grand-
570 a	verage ERP waveform and topographic maps (depicted at the N1 and P2 peak) of all
571 p	articipants with valid data ($n = 40$). The butterfly plot of the ERP waveform is spatially
572 c	olored by the channel locations. (B) Left: ERP waveform at region of interest (channels 6, 7
573 a:	nd 106) for the four groups separately (n = 10 in each group). Right: Topographic maps of
574 tv	wo major components for the four groups separately: the auditory N1 peaks at around 112
575 m	ns and auditory P2 component peaks around 189 ms. N1 amplitude was larger in participants
576 tr	rained at EXP than at INS (univariate ANOVA) and larger in participants trained at EXP-
577 D	DIA than at INS-SYS (independent samples t-test). Asterisks refer to statistically significant
578 d	ifferences between groups in N1 amplitude: * p < 0.050, *** p \leq 0.001.
579	

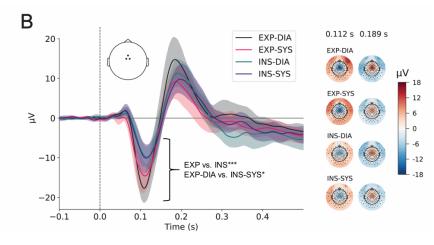
581	Acknowledgements
582	We thank Jan Wikgren, Lauri Viljanto, Viki-Veikko Elomaa and Petri Kinnunen for technical
583	help in building the recording systems. Special thanks to Jan Wikgren for helping with the
584	manuscript.
585	Grants
586	The work was supported by the Academy of Finland grant number 321522 to MSN.
505	
587	Disclosures
588	The authors declare no conflict of interest.
589	
590	











APPENDIX 1: A FULL QUESTIONNAIRE FOR PARTICIPANTS

The researcher fills in: ID	Date	Time	_
Please fill in the missing informa	tion and answei	the following questi	ons.
Age:			
Sex:			
Handedness:			
1) What equipment did the man in	the film use for t	ravelling in the snow?	
2) Were there any light phenomena	a in the film?		
3) Name two to three animals you	saw in the film.		
4) Did the fishermen catch any fish	1?		
5) What large bird appeared at the	beginning of the	second film?	
6) When did the air puff occur?			
			PLEASE TURN!

Proposition	True	False
1. The air puff occurred right <i>before</i> the beep.		
2. The air puff occurred right <i>after</i> the beep.		
3. The beep occurred right <i>before</i> the air puff.		
4. The beep occurred right <i>after</i> the air puff.		
5. The beep and the air puff always occurred very close to each other.		
6. The beep and the air puff occurred close to each other only occasionally.		
7. The beep predicted the air puff.		

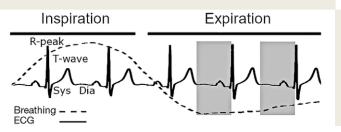


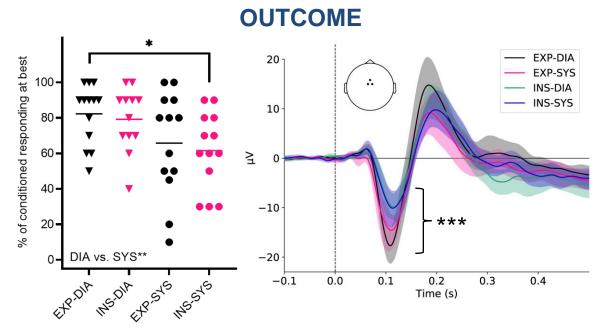
Bodily rhythms affect conditioning



METHODS

Breathing (EXPiration vs. INSpiration) and cardiac cycle phase (DIAstole vs. SYStole) were monitored to trigger eyeblink conditioning trials in four different bodily states while participants were watching a wildlife documentary. Conditioned eyeblinks and electroencephalogram were recorded.





CONCLUSION

Breathing and heartbeat have a phase combination (diastole during expiration) that is indicative of a neural state beneficial for conditioning.