

Title: Cross-species real time detection of trends in pupil size fluctuation

Authors: Sharif I. Kronemer¹, Victoria E. Gobo¹, Joshua B. Teves¹, Diana C. Burk², Somayeh Shahsavarani^{1,3}, Catherine R. Walsh¹, Javier Gonzalez-Castillo¹, Peter A. Bandettini^{1,4}

Affiliations: [1] Section on Functional Imaging Methods, Laboratory of Brain and Cognition, National Institute of Mental Health (NIMH), National Institutes of Health (NIH), Bethesda, Maryland (MD), United States of America (USA), [2] Laboratory of Neuropsychology, NIMH, NIH, Bethesda, MD, USA, [3] Mortimer B. Zuckerman Mind Brain Behavior Institute and Department of Biomedical Engineering, Columbia, University, New York (NY), NY, USA, [4] Functional Magnetic Resonance Imaging Core Facility, NIMH, NIH, Bethesda, MD, USA

Abstract:

Pupillometry is a popular method because pupil size is an easily measured and sensitive marker of neural activity and associated with behavior, cognition, emotion, and perception. Currently, there is no method for monitoring the phases of pupillary fluctuation in real time. We introduce *rtPupilPhase* – a software that automatically detects trends in pupil size in real time, enabling novel implementations of real time pupillometry towards achieving numerous research and translational goals.

Main Text:

Pupillometry – the measure of pupil size – has grown in popularity with evidence that pupil size is an easily measured marker of neurophysiological state. Neuromodulatory networks, including the locus coeruleus-noradrenergic and basal forebrain-cholinergic systems, are linked with changes in pupil size (Figure 1A)^{1,2}. These networks are also involved in widespread regulation of cortical and subcortical regions^{3,4}. Therefore, pupillary dynamics may reflect both large-scale brain activity and behavioral, cognitive, emotive, and perceptual states that emerge from central neural mechanisms. For example, pupil size has been shown to reveal states of locomotion, conscious awareness, and arousal⁵⁻⁷.

The neural underpinnings of pupil size explain *pupillary unrest*: fluctuations in pupil size under constant environmental luminance (Figure 1B)^{8,9}. The light-independent trend of pupillary unrest includes four main phases: (1) *dilation*, (2) *peak* (i.e., local maximum), (3) *constriction*, and (4) *trough* (i.e., local minimum; Figure 1B). Pupil phase is sensitive to neural activity, thus a real time method for detecting trends in pupil size is valuable in various experimental and translational contexts (Figure 1B)^{1,4,6,10,11}. However, pupillometry is commonly recorded passively and analyzed post hoc, which restricts experimental design and the interpretation of results. Real time pupillometry would enable new task types (e.g., closed-loop paradigms) and novel applications (e.g., self-regulation of arousal state via pupil size feedback¹²). Despite the potential advantages, there is currently no published procedure for the real time detection of pupil phase.

To address this deficit, we introduce *rtPupilPhase* – a software that automatically detects dilation, peak, constriction, and trough pupil phase events in real time. *rtPupilPhase* also identifies pupil phase-independent or *random* events, which we used to evaluate the performance of *rtPupilPhase*. Monitoring pupil phase in real time is challenging, in part, because pupil size change is sluggish, undergoes baseline shifts, and the regular occurrence of behaviors that occlude the pupil (e.g. blinking and whisking in rodents). *rtPupilPhase* mounts these challenges and achieves the real time detection of pupil phase events in four main stages (Figure 1C; see *Real Time Detection of Pupil Phase Events* Methods section). In summary, *rtPupilPhase* aggregates intervals of live-streamed pupil size data, models these data intervals, and records detected pupil phase events determined by recurrently updated, individual-specific pupil size and pupil size derivative (i.e., pupil size samples x_1 - x_2 , x_2 - x_3 , etc.) thresholds.

We evaluated *rtPupilPhase* using data from healthy, adult human participants (N = 8) who completed a passive fixation task integrated with *rtPupilPhase* under constant luminance and simultaneous pupillometry recording (see *Fixation Task* and *Human Pupillometry Acquisition* Methods

sections). rtPupilPhase detected hundreds of pupil phase events in real time per participant (Supplementary Figure 1). We extracted epochs of pupil size data centered on these pupil phase events (see *Human Pupil Size, Blink, and Saccade Fraction Epoch Extraction Methods* section). The resulting pupil size epoch timecourses demonstrated that rtPupilPhase reliably located pupil phases with high-temporal precision at the event (Supplementary Figure 2), participant (Supplementary Figure 3A), and group levels (Figure 1A; Supplementary Figure 4A). In addition, the pupil size timecourses significantly differed ($p < 0.05$; cluster-based permutation testing; see *Cluster-Based Permutation Testing Statistical Analysis* section) between pupil phase and random events, particularly in the interval immediately preceding the pupil phase event (Supplementary Figure 4A).

To assess the accuracy of rtPupilPhase, we compared the real time detected pupil phase events against the *true* pupil phase events, as determined by post hoc evaluation of the pupillometry data (see *Pupil Phase Detection Accuracy Statistical Analysis* section). rtPupilPhase achieved an average pupil phase event-level accuracy of 88.16 (dilation), 79.26 (peak), 86.90 (constriction), and 73.37 percent (trough), all statistically greater than the random event accuracy of approximately 50 percent ($p < 0.0005$; Supplementary Figure 5A, B). These results affirm that rtPupilPhase correctly identifies pupil phases near the moment of their true occurrence in real time.

We also explored the rate of blinks and saccades preceding the real time detected pupil phase events, as eye movements are known to affect pupil size¹³. Blink and saccade occurrence epochs centered on the pupil phase events detected with rtPupilPhase were extracted. The resulting epoch timecourses averaged within event type and across participants revealed increases in blink and saccade fraction preceding the event time, each event type with different periods of peak change (Supplementary Figure 6C, E). To determine if the performance of rtPupilPhase was explained by these eye movements, we removed all pupil size epochs with blinks *or* saccades in the 0.5 seconds preceding the real time detected pupil phase events. Nonetheless, the pupil size timecourse trends among the remaining event epochs were unchanged across pupil phases (Supplementary Figure 6A). This result supports that rtPupilPhase detects pupil size trends independent of eye movements.

Next, we aimed to evaluate if rtPupilPhase could be used on pupillometry from non-human animals. As a proof of concept, we developed *sim-rtPupilPhase* – a software that *simulates* rtPupilPhase offline using previously collected pupil data (see *Human Simulated Real Time Detection of Pupil Phase Events Methods* section). We verified that *sim-rtPupilPhase* replicates the performance of rtPupilPhase using the previously described human pupil data. Although, achieving an exact duplication of the rtPupilPhase result with *sim-rtPupilPhase* was not possible due to the variable online sampling rate of the human pupillometry recordings (see *Human Pupillometry Acquisition Methods* section). Still, a similar number of pupil phase events were detected with *sim-rtPupilPhase* as with rtPupilPhase (Supplementary Figure 1B). Crucially, the extracted epochs centered on the pupil phase events detected with *sim-rtPupilPhase* reproduced the trends and statistics across pupil size, blink, and saccade fraction timecourses from events detected with rtPupilPhase (Figure 2A versus B; Supplementary Figure 3A versus B; Supplementary 4A versus B; Supplementary 6A versus B, C versus D, and E versus F).

After validating on humans, we implemented *sim-rtPupilPhase* to estimate the performance of rtPupilPhase on previously acquired pupil data from mice ($N = 5$; ¹⁴) and rhesus macaques ($N = 2$; data newly published here). As with the human data set, *sim-rtPupilPhase* detected hundreds of pupil phase events for each non-human animal (Supplementary Figure 1C, D). Likewise, the trend of the pupil size epoch timecourses centered on these events was consistent with the human event pupil size timecourses detected with rtPupilPhase on the individual animal (Supplementary Figure 3C, D) and group levels (Figure 2C, D; Supplementary Figure 4C, D). These non-human animal results were achieved with minimal adjustment to *sim-rtPupilPhase* and rtPupilPhase parameters despite major differences among the data sets, including pupil physiology, experimental setup, and recording system (Table 1; see *Mouse and Monkey Simulated Real Time Detection of Pupil Phase Events Methods* sections). In the future, *sim-rtPupilPhase* can be used as a tool for optimizing the rtPupilPhase parameters for testing alternative data sets and achieving unique research goals.

In conclusion, real time pupillometry is a nascent approach with exciting applications. In a major advance, rtPupilPhase is a novel method for monitoring light-independent trends in pupil size at rest and linked with behavioral, cognitive, emotive, and perceptual states across species. An immediate use of rtPupilPhase includes developing closed-loop paradigms where the real time detection of pupil phase events triggers experimental stimuli. These types of paradigms are powerful to efficiently target pupil phases of interest and deduce casual relationships. Thus, rtPupilPhase introduces novel opportunities to study and implement the pupil-neurophysiology link towards achieving broad experimental and translational goals.

Methods

Participants. Healthy, adult humans ($N = 8$; mean age = 25.1 years; age standard deviation [SD] = 4.01 years; males = 1; mean education = 17.0 years; education SD = 1.9 years) were recruited from the local Bethesda, Maryland, USA community. The Institutional Review Board of the National Institute of Mental Health approved the recruitment and consent protocols. Inclusion criteria included: (1) being between 18 and 65 years old at the time of experimentation, (2) a healthy physical examine within a year of the study session, and (3) able to give informed consent. Exclusion criteria included: (1) no previous nor current histories of neurologic or psychiatric disorders, (2) poor vision that cannot be corrected, and (3) head injuries (e.g., loss of consciousness for >30 minutes and three or more concussive injuries). Each participant underwent a health exam completed by a nurse practitioner prior to their study session.

Fixation Task. The fixation task was coded in Python and run with PsychoPy (v2022.2.4; Open Science Tools Ltd.). The task consisted of a single display: a central fixation point (a plus sign; 0.51×0.51 degrees) on a solid gray screen. Participants were instructed to maintain central fixation throughout the task. Participants completed five 10-minute fixation task blocks with a ~2-minute break between each block. Integration of rtPupilPhase with the fixation task enabled the covert detection of pupil phase and random events (see *Real Time Detection of Pupil Phase Events* section) and accommodates future development of alternative experimental designs involving rtPupilPhase.

Human Pupillometry Acquisition. Human, head-fixed pupillometry and eye tracking was acquired with a desktop mounted EyeLink 1000 Plus (recorded eye: right eye; offline sampling rate = 1000 Hz; Version 5.50; SR Research, Inc.). The online sampling rate of the pupil size data was variable (mode sampling rate ≈ 60 Hz or 17 milliseconds between samples). During pupillometry acquisition, participants were instructed to place their head in a chin and forehead rest system (SR Research Head Support; SR Research, Inc.) to maintain a fixed head position and a constant distance between the participant and pupillometer.

Human Testing Equipment and Facility. Human testing was conducted in a single 1.5-hour study session in a windowless, temperature-controlled behavioral testing room. The lighting in the testing room was set to a consistent level for all participants and maintained throughout the entire study session. The fixation task was administered on a behavioral laptop (MacBook Pro; 13-inch; 2560×1600 pixels, 2019; Mac OS Catalina v10.15.7, Apple, Inc.)¹⁵. The EyeLink 1000 Plus software ran on a Dell desktop computer (Dell OptiPlex XE2, Dell, Inc.). The behavioral laptop and the fixation task were programmed to communicate with the EyeLink desktop via an Ethernet connection. Participants viewed the fixation task on a VIEWPixx display monitor (1920×1200 pixels; VPixx Technologies, Inc.). The behavioral laptop monitor was mirrored to the display monitor via a DVI cable. Participants were positioned approximately 56 centimeters from the center of the display monitor, measured from the nasion.

Mouse Pupillometry Data Set. The mouse pupillometry data were acquired from a previously published experiment (<https://zenodo.org/records/7968402>)¹⁴. In summary, *in vivo* wide-field optical mapping, pupil size, locomotion, and whisking data were recorded from five adult male Thy1-jRGECO1a transgenic mice over multiple recording sessions. Throughout these sessions, the mice were head-fixed and free to exhibit spontaneous behaviors such as walking and running on a wheel without any experimental stimuli. The mouse pupil size was recorded with a BFS-U3-16S2M-CS USB 3.1 Blackfly S Monochrome Camera and sampled at ~ 20 Hz. Using the tracking software DeepLabCut¹⁶, pupil size was monitored by identifying eight circumferential points, then fitting a circle to these points to estimate pupil diameter. For the current experiment, eight 10-minute recording sessions were selected for each mouse. Any session in which significant portions of the pupil data were missing was not selected for inclusion in the mouse data set.

Monkey Pupillometry Data Set. The monkey pupillometry data consisted of two adult (~7 years old), male rhesus macaques who completed a variant of the visual reinforcement learning task previously published in ¹⁷ with simultaneous pupillometry recording. The pupillometry data were acquired in non-contiguous trials lasting several seconds (~2-10 seconds), with each data segment corresponding to a single task trial. One study session was selected per monkey, each including more than 1500 trials of the learning task. Only trials with pupillometry recordings lasting longer than 5 seconds were selected for each monkey. This criterion resulted in the inclusion of 184 and 202 trials for each monkey, respectively, or approximately 25 minutes of pupillometry data per monkey.

Head-fixed pupillometry was acquired using the MATLAB-based (MathWorks, Inc.) Monkeylogic toolbox (Version 2.2; <https://monkeylogic.nimh.nih.gov>; ¹⁸) and Arrington Viewpoint eye-tracking system (recorded eye: left eye; sampling rate: 1000 Hz; Arrington Research, Inc.) run on a desktop computer (Dell OptiPlex 9020; Dell, Inc.). An infrared camera was positioned ~40 centimeters from the monkey. Pupil size was acquired in arbitrary units, linked to the voltage output of the eye tracking system. The pupillometry data reported in the current study were not previously published.

Target Pupil Phase Events. Pupil size fluctuations consist of four main phases: (1) dilation, (2) peak, (3) constriction, and (4) trough (Figure 1B). The *dilation* phase is when pupil size increases. The *peak* phase is when the pupil has reached a local maximum (i.e., the moment of a phase switch from a period of dilation to constriction). The *constriction* phase is when pupil size decreases. The *trough* phase is when the pupil has reached a local minimum (i.e., the moment of a phase switch from a period of constriction to dilation). During pupillary unrest – the spontaneous fluctuation of pupil size under constant environmental luminance – pupil size cycles among these four pupil phases, in part, according to neurophysiological state and other factors, including eye movements (e.g., blinks)¹³.

Real Time Detection of Pupil Phase Events. Real time monitoring methods are available for various physiologic signals, for example, neural and cardiac potentials^{19,20}. Meanwhile, there are limited real time monitoring methods for the pupil, in part, because pupil physiology presents unique challenges for developing real time methods. First, pupil size undergoes large baseline shifts over seconds, for example, linked to states of arousal and locomotion. These low-frequency drifts limit the efficacy of applying constant pupil size thresholds for detecting pupil phases. Second, pupil size trends do not conform to a consistent profile or shape (e.g., the stereotyped membrane potential change of a neuronal action potential). Third, spontaneous and evoked pupil size changes are slow – unfolding over 100s to 1000s of milliseconds. Therefore, at the sampling rates commonly used in pupillometry data acquisition (>20 Hz), there are small differences in pupil size between adjacent data samples. As a result, a derivative approach (i.e., pupil size sample $x_1 - x_2$, $x_2 - x_3$, etc.) to determine pupil phase is limited. Finally, the pupillometry data stream is subject to repeated artifacts and interruption due to behaviors that occlude the pupil, including blinks, drooping eyelid (e.g., due to drowsiness), and whisking in rodents. *rtPupilPhase* – our method for detecting the phases of pupillary fluctuation in real time – overcomes these unique challenges of pupil physiology for developing real time pupillometry methods.

rtPupilPhase is a Python-based software that we integrated with a fixation task (see *Fixation Task* section). The main goal of *rtPupilPhase* is to analyze live streamed pupil size data and detect the real time occurrence of target pupil phase events: dilation, peak, constriction, and trough (Figure 1B; see *Target Pupil Phase Events* section). As a control condition, *rtPupilPhase* also selects pupil phase-independent or *random* events. Here, we use the random events to measure the performance of *rtPupilPhase*. Future applications of *rtPupilPhase* can suppress the selection of random events or implement these events as a control or baseline condition against the pupil phase events.

The *rtPupilPhase* method is summarized in four main stages (Figure 1C; see Table 1 for a list of the method parameters):

Stage 1: Monocular, pupil size data is live streamed (see *Human Pupillometry Acquisition* section) and stored in brief, continuous intervals of pupil size data called a *pupil sample*. The pupil sample is filled with sequential pupil size data points until it comprises a total of 0.1 seconds of pupil

size data (i.e., ~6 pupil size data points at ~60 Hz sampling rate). For the human pupillometry acquisition system, blinks, and other interruptions in the pupil data stream (e.g., lost eye tracking due to head movement or drooping eyelid) results in a pupil size value of 0. Any pupil size value in the pupil sample equal to 0 is replaced with the designation ‘not-a-number’ or NaN.

Stage 2: The pupil sample is simultaneously added to a *search window* and *baseline window*. Subsequently, the pupil sample is cleared, and a new pupil sample is built (see Stage 1). Thus, overtime, the search and baseline windows grow, comprising continuous pupil size data that are sequentially added in increments of pupil sample (i.e., 0.1 seconds). The search window is analyzed for pupil phase events (see Stage 3). The baseline window is used to recurrently update the pupil size and pupil size derivative thresholds that determine the occurrence of pupil phase events within the search window (see Stage 4). If one or more samples in the current search window includes a NaN, the search window is cleared, and a new search window built. The current search window is also reset if the *previous* search window contained at least one NaN sample. With this approach, rtPupilPhase limits the influence of blinks and other common human pupillometry recording artifacts in the real time detection of pupil phase events by only considering pupil data intervals absent these artifactual occurrences.

Stage 3: When the search window comprises a minimum of two pupil samples (i.e., at least two cycles of Stages 1 and 2 have iterated without resetting the search window), the search window data is demeaned (i.e., the mean of all the pupil size data in the search window is subtracted from all search window samples). Next, the demeaned search window pupil size data is fit with a quadratic model (x-coordinates = 0 to the number of samples in the search window minus one; NumPy function *polyfit*; <https://numpy.org>). The resulting fitted trace is a model of the pupil size data in the search window. The final pupil size value from the fitted model timecourse, corresponding with the final sample in the search window, is recorded for later use to guide determining pupil phase (see Stage 4). Stages 1 through 3 iterate until a target pupil phase event is detected (see Stage 4) or if the search window exceeds 5 seconds of cumulative pupil size data. In either case, the search window is emptied and built anew from subsequent pupil samples (see Stages 1 and 2).

Stage 4: When there is at least one previous fitted model (i.e., Stage 3 has iterated two or more times), the previous and current final pupil size values from the fitted model timecourses are compared to determine the occurrence of a target pupil phase event. A peak event is detected when the final pupil size value of the current search window is *less than* the final pupil size value from the previous search window (final value of search window_i < final value of search window_{i-1}) *and* the final pupil size value from the current search window is *greater than* the pupil size 75-percentile for all peak events detected in the previous baseline window (see details below). A trough event is found when the final pupil size value of the current search window is *greater than* the final pupil size value from the previous search window (final value of search window_i > final value of search window_{i-1}) *and* the final pupil size value from the current search window is *less than* the pupil size 25-percentile for all trough events found in the previous baseline window (see details below). The dilation and constriction events are detected when the difference between the final pupil size values from the fitted model timecourses of the current and previous search windows (final value of search window_i – final value of search window_{i-1}) is *greater than* the pupil size derivative 99-percentile or is *less than* the pupil size derivative 1-percentile, respectively, the derivative percentile thresholds evaluated across all valid pupil size samples in the previous baseline window (see details below).

In the initial iterations of Stage 4, the pupil size and pupil size derivative thresholds (Table 1) are set to default constants (peak pupil size threshold = 0; trough pupil size threshold = 0; dilation pupil size derivative threshold = 50; constriction pupil size derivative threshold = -50). Subsequently, these thresholds are updated when the baseline window is filled – approximately every 5 seconds. At least half of the baseline window samples must be valid pupil size values (i.e., non-NaN) to accept the current baseline window for analysis to update the pupil size and pupil size derivative thresholds. If the baseline window is rejected, the threshold values remain the same and the baseline window is cleared and filled again (see Stage 2). However, when a valid baseline window is confirmed, the baseline window pupil size data is demeaned, and all NaN pupil size samples are removed from the baseline window. The pupil

size threshold is determined by finding all peak and trough events (SciPy function *find_peaks*; <https://scipy.org>) in the demeaned baseline window and then calculating the pupil size 75-percentile among peaks and pupil size 25-percentile of among troughs. Prior to finding troughs in the baseline window, the pupil size data are inverted (i.e., multiplied by negative 1), so that finding peaks in the inverted data identifies troughs in the original, uninverted data set. The pupil size at each trough event is calculated with the original data. The pupil derivative thresholds are calculated by finding the derivative across all valid pupil size samples in the baseline window (NumPy function *diff*; <https://numpy.org>) and then calculating the 99 and 1-percentile of all derivative values.

When a pupil phase event is detected, the search window is cleared, and the procedure begins anew from Stage 1. An inter-event interval (IEI) of 3 seconds is set so that rtPupilPhase will not log a so-called *accepted* pupil phase event unless the IEI has been exceeded. Events detected within the IEI are logged but designated as *not* accepted. When the IEI is exceeded for any detected pupil phase event, the search window is *not* cleared, instead additional pupil samples are added to the search window until a pupil event is detected that exceeds the IEI or the maximum length of the search window (5 seconds) is exceeded, in either case, the search window is reset. We implemented the IEI in the current demonstration of rtPupilPhase to limit the number of events analyzed to manage the computational load and so that the data samples contained within the pupil size, blink, and saccade fraction epochs are non-overlapping with any other event epoch. Future application of rtPupilPhase can modify the IEI duration or eliminate it altogether (i.e., set the IEI to 0 seconds).

Human Simulated Real Time Detection of Pupil Phase Events. Simulated rtPupilPhase or *sim-rtPupilPhase* is a MATLAB-based (MathWorks, Inc.) software designed to simulate the rtPupilPhase method in a live recording using previously acquired pupil data. *sim-rtPupilPhase* was developed as a proof of concept to assess the performance of rtPupilPhase on pupil data from non-human animals. In future applications, *sim-rtPupilPhase* can serve as a tool for optimizing the rtPupilPhase parameters to achieve specific experimental goals. In summary, *sim-rtPupilPhase* loads previously collected pupillometry data and streams the data to rtPupilPhase method (see *Real Time Detection of Pupil Phase Events* section; Figure 1C). The pupil data was modeled (see Stage 3 of the rtPupilPhase method) with the MATLAB function *fit* (x-coordinates = 1 to the number of samples in the search window; 0 to the number of samples in the search window minus one was also tested and achieved similar results [data not shown]; MathWorks, Inc.). To directly compare the performance between rtPupilPhase and *sim-rtPupilPhase* on the human pupillometry data set, the pupil size data were down sampled from its offline sampling rate of 1000 to 60 Hz (i.e., the approximate online sampling rate of the human pupillometry recordings; see *Human Pupillometry Acquisition* section). Otherwise, the *sim-rtPupilPhase* parameters (e.g., the pupil size and pupil size derivative thresholds) were set to exactly as those used in the rtPupilPhase live study (Table 1).

Mouse Simulated Real Time Detection of Pupil Phase Events. All pupil size samples concurrent with a locomotion event or with a value less than 0 (e.g., indicative of blinking or whisking) were replaced in the pupil data with NaN (see *Mouse Pupillometry Data Set* section). Otherwise, all *sim-rtPupilPhase* parameters and procedures were exactly as those used in the human *sim-rtPupilPhase* testing (Table 1; see *Human Simulated Real Time Detection of Pupil Phase Events* section).

Monkey Simulated Real Time Detection of Pupil Phase Events. To accommodate the brief pupillometry data segments (<10 seconds) acquired in monkeys (see *Monkey Pupillometry Data Set* section), several adjustments were made to the *sim-rtPupilPhase* parameters (Table 1). First, the baseline window duration was adjusted from 5 seconds, as used in humans and mice, to 0.5 seconds. In addition, where in humans and mice, all event types were considered together to determine if the IEI threshold was breached, in the monkey data set, each pupil phase event was considered relative to its own event type to determine if the IEI was exceeded (i.e., events of different type could fall within the IEI). This adjustment was made to help increase the number of accepted pupil phase events detected in the monkey

pupillometry data. Another adaptation made in the monkey sim-rtPupilPhase procedure was the initial removal of artifactual periods within each trial using the human pupil size preprocessing procedure (see *Monkey Pupillometry Data Set* and *Human Pupil Size, Blink, and Saccade Fraction Epoch Extraction* sections). Preprocessing was necessary because the monkey pupil data contained numerous artifacts, including those linked to eye movements and task stimuli. Otherwise, all sim-rtPupilPhase parameters and procedures were exactly as those used in human sim-rtPupilPhase (Table 1; see *Human Simulated Real Time Detection of Pupil Phase Events* section).

Statistical Analysis

Human Pupil Size, Blink, and Saccade Fraction Epoch Extraction. Epoch extraction involved selecting segments of pupil size, blink, and saccade occurrence data centered on the pupil phase and random event times that were detected by rtPupilPhase and sim-rtPupilPhase. Prior to extracting the pupil size epochs, the pupillometry data were preprocessed for each participant, including the removal of blink and other artifactual events (e.g., drooping eyelid) that result in blank periods in the pupil size data stream (*stublinks.m* available at <http://www.pitt.edu/~gsiegle>)²¹. The preprocessing procedure also outputs a binary blink vector of the same length as the inputted pupil size data – blink data samples indicated with 1 and non-blink samples with 0. Similarly, a saccade binary vector was made using an established saccade detection method^{22,23} – saccade data samples indicated with 1 and non-saccade samples with 0. Note that the saccade binary vector was inclusive of microsaccades. Therefore, each participant contributed a preprocessed pupil size, blink binary, and saccade binary vector that extended the entire pupillometry recording session.

The preprocessed pupil size, blink binary, and saccade binary data were cut into 5001-millisecond epochs centered on each pupil phase and random event (i.e., 2.5 seconds before and after the event time) detected with either rtPupilPhase or sim-rtPupilPhase. The epoch was not extracted if 2.5 seconds before or after the event time exceeded the length of the pupillometry recording. The extracted pupil size epochs were demeaned (i.e., the mean pupil size across the entire epoch was subtracted from all epoch samples). After extracting all epochs, the mean across epochs was calculated within participant and events detected with rtPupilPhase and sim-rtPupilPhase, so that each participant contributed a rtPupilPhase and sim-rtPupilPhase mean pupil size, blink, and saccade fraction timecourse per pupil phase and random event. To interrogate the sensitivity of rtPupilPhase to eye movements (e.g., blinks and saccades), a subset of pupil size epochs with neither blinks nor saccades in the 0.5 seconds prior to the event time were selected and averaged separately within participant and rtPupilPhase and sim-rtPupilPhase methods (Supplementary Figure 6A, B). The participant-level saccade timecourses were smoothed using the MATLAB *smooth* function (window size = 100; MathWorks, Inc).

Subsequent group-level analyses involved finding the mean across participant pupil size, blink, and saccade fraction timecourses within event type and rtPupilPhase and sim-rtPupilPhase detection methods (Supplementary Figure 4A, B; Supplementary Figure 6A, B). An alternative group-level analysis first calculated the z-score across participant timecourses and then calculated the mean of the z-scored timecourses across participants within event and method type (Figure 2A, B; Supplementary Figure 3A, B). The standard error of the mean was calculated across participants. For visualization of the pupil size, blink, and saccade fraction timecourses, only the 1.5 seconds before and after the event were displayed (Figure 2A, B; Supplementary Figures 2B; 3A, B; 4A, B; 6).

Mouse Pupil Size Epoch Extraction. The mouse pupillometry data were cut into 5001-millisecond epochs (101 samples at ~20 Hz sampling rate; see *Mouse Pupillometry Data Set* section) centered on each pupil phase and random event detected with sim-rtPupilPhase (i.e., 2.5 seconds before and after the event time). The epoch was not extracted if 2.5 seconds before or after the event time exceeded the length of the pupillometry recording. Next, the pupil size epochs were demeaned (i.e., the mean pupil size across the entire epoch was subtracted from all epoch samples). After extracting all epochs, the mean across epochs was calculated within mouse and event type, so that each mouse contributed a

single mean pupil size timecourse for each event. Group-level epoch analyses involved finding the mean across mouse pupil size timecourses within event type (Supplementary Figure 4C). An alternative group-level analysis first calculated the z-score across mouse pupil size timecourses and then calculated the mean of the z-scored timecourses across mice (Figure 2C; Supplementary Figure 3C). The standard error of the mean was calculated across mice. For visualization of the pupil size timecourses, only the 1.5 seconds before and after the event were displayed (Figure 2C; Supplementary Figure 3C; Supplementary Figure 4C).

Monkey Pupil Size Epoch Extraction. Prior to extracting the monkey pupil size epoch timecourses, the pupil data were preprocessed using the same method applied on the human pupillometry data (see *Human Pupil Size, Blink, and Saccade Fraction Epoch Extraction* section). The monkey pupillometry data were cut into 1001-millisecond epochs centered on the pupil phase and random events (i.e., 0.5 seconds before and after the event time) that were detected by *sim-rtPupilPhase* (see *Monkey Simulated Real Time Detection of Pupil Phase Events* section). The epoch duration was reduced for the monkey pupillometry data relative to the 5001-millisecond epochs extracted from the human and mouse data sets because the monkey data were acquired in short segments (see *Monkey Pupillometry Data Set* section). Therefore, cutting longer epochs exceeded the bounds of the pupillometry data acquired in each data segment. Accordingly, epochs were not extracted if 0.5 seconds before or after the event time exceeded the length of the pupillometry recording. Next, the pupil size epochs were demeaned (i.e., the mean pupil size across the entire epoch was subtracted from all epoch samples). Unique to the monkey data set, the demeaned epochs were also detrended using the MATLAB *detrend* function (MathWorks, Inc.) to correct for a low-frequency drift in pupil size related to the task the monkeys were engaged. The mean across all detrended epochs within monkey and pupil phase and random events was calculated, so that each monkey contributed a single mean pupil size timecourse for each event. Group-level epoch analyses involved finding the mean across all monkey-mean pupil size epoch timecourses within event type (Supplementary Figure 4D). An alternative group-level analysis first calculated the z-score across monkey epoch timecourses and then calculated the mean of the z-scored epochs across monkeys for each pupil phase and random event (Figure 2D; Supplementary Figure 3D). The standard error of the mean was calculated across monkeys.

Cluster-Based Permutation Testing. Human pupil size timecourses for the pupil phase events were statistically tested against the random event pupil size timecourse with cluster-based permutation testing (number of permutations = 250; $p < 0.05$). The permutation analysis method employed in this study is a modified version of the method shared in the Mass Univariate ERP Toolbox (https://openwetware.org/wiki/Mass_Univariate_ERP_Toolbox)²⁴. This approach was also used in a previous experiment to statistically analyze pupil size timecourses⁵. In summary, the cluster-based permutation testing method involves creating positive and negative null statistical distributions by participant-level permutation between the compared event types (e.g., peak versus random event). Subsequently, a one-sample *t*-test was performed on the non-permuted data, identifying statistically significant positive and negative clusters based on the null distributions. Clusters were defined by temporal adjacency (i.e., statistically significant data samples contiguous in time). Four cluster-based permutation analyses were evaluated, each for the *rtPupilPhase* and *sim-rtPupilPhase* human pupil size timecourses: (1) dilation versus random, (2) peak versus random, (3) constriction versus random, and (4) trough versus random events (Supplementary Figure 4A, B). The mouse and monkey data were not subjected to statistical testing due to low individual animal sample sizes.

Pupil Phase Event Detection Accuracy. The accuracy of *rtPupilPhase* in detecting pupil phase events was calculated based on accepted events (i.e., those that exceeded the IEI; see *Real Time Detection of Pupil Phase Events* section). Event accuracy was evaluated by comparing the real time detected pupil phase event versus the *true* pupil phase. The true pupil phase was determined by post hoc analysis of the pupillometry data. In summary, first, the human pupillometry data were preprocessed, including

removing blinks and artifactual samples (see *Human Pupil Size, Blink, and Saccade Fraction Epoch Extraction* section). Next, the preprocessed pupil data were smoothed using the MATLAB *smoothdata* function (Savitzky-Golay filter; window size = 100; MathWorks, Inc.) to eliminate high-frequency signals. The true peak and trough pupil phase events were identified using the MATLAB *findpeaks* function (MathWorks, Inc.). For trough events, prior to implementing *findpeaks*, the pupil size data were inverted (i.e., multiplied by negative 1). Finally, the top 75 percent of peaks and troughs were selected based on their prominence. For two participants, the prominence percent threshold was adjusted to 90 percent because visual inspection determined that lower thresholds resulted in the selection of peak and trough events with low prominence. Finally, peak and trough event binary indices were made by setting all samples within 0.25 seconds before and after each peak and trough event to an index value of 1, while all other samples outside this range were set to 0.

The true dilation and constriction pupil phase events were found by calculating the slope at every sample of the pupil size data using the MATLAB *gradient* function (MathWorks, Inc.). A dilation event binary index was created by setting all samples from the gradient output with a positive value to 1, indicating a positive slope or dilation, and a negative value to 0, indicating a negative slope or constriction. Conversely, the constriction binary index was created by setting all the samples from the gradient output with a negative value to 1 and a positive value to 0.

Finally, the accuracy of the real time detected pupil phase events was assessed by comparing each event sample to the corresponding true pupil phase event index. An index value of 1 at the queried event sample indicated agreement between the real time and true pupil phase events. The real time detected event accuracy percentage was calculated within participant and pupil phase event type detected with *rtPupilPhase* by determining the total number of pupil phase events that were accurately detected (i.e., in agreement with the true pupil phase event indices) divided by the total number of detected pupil phase events. The result was multiplied by 100 to convert from rate to percentage units. The baseline or chance accuracy rate for each pupil phase event was determined by comparing the random event samples to each of the true pupil phase event indices (e.g., random event versus true dilation event index, random event versus true peak event index, etc.). Finally, to assess if the real time pupil phase event accuracy was greater than the random accuracy percentage, group-level paired *t*-tests across participants were conducted to statistically compare pupil phase and random event accuracies (Holm-Bonferroni corrected; $p < 0.05$). A total of four *t*-tests were completed: (1) dilation versus random, (2) peak versus random, (3) constriction versus random, and (4) trough versus random events (Supplementary Figure 5).

Data and Code Availability

All code and data are available at <https://github.com/nimh-sfim/rtPupilPhase>.

Author Contributions

S.I.K. contributed to conceptualization, methodology, software, formal analysis, investigation, data curation, visualization, supervision, project administration, and writing (original draft); V.E.G. contributed to methodology, software, formal analysis, investigation, and writing (review and editing); J.B.T. contributed to methodology, software, and writing (review and editing); D.C.B. contributed to formal analysis, resources, and writing (review and editing); S.S. contributed to formal analysis, resources, and writing (review and editing); C.R.W. contributed to software, data curation, and writing (review and editing); J.G-C. contributed to conceptualization, methodology, supervision, and writing (review and editing); P.A.B. contributed to conceptualization, methodology, supervision, funding acquisition, and writing (review and editing).

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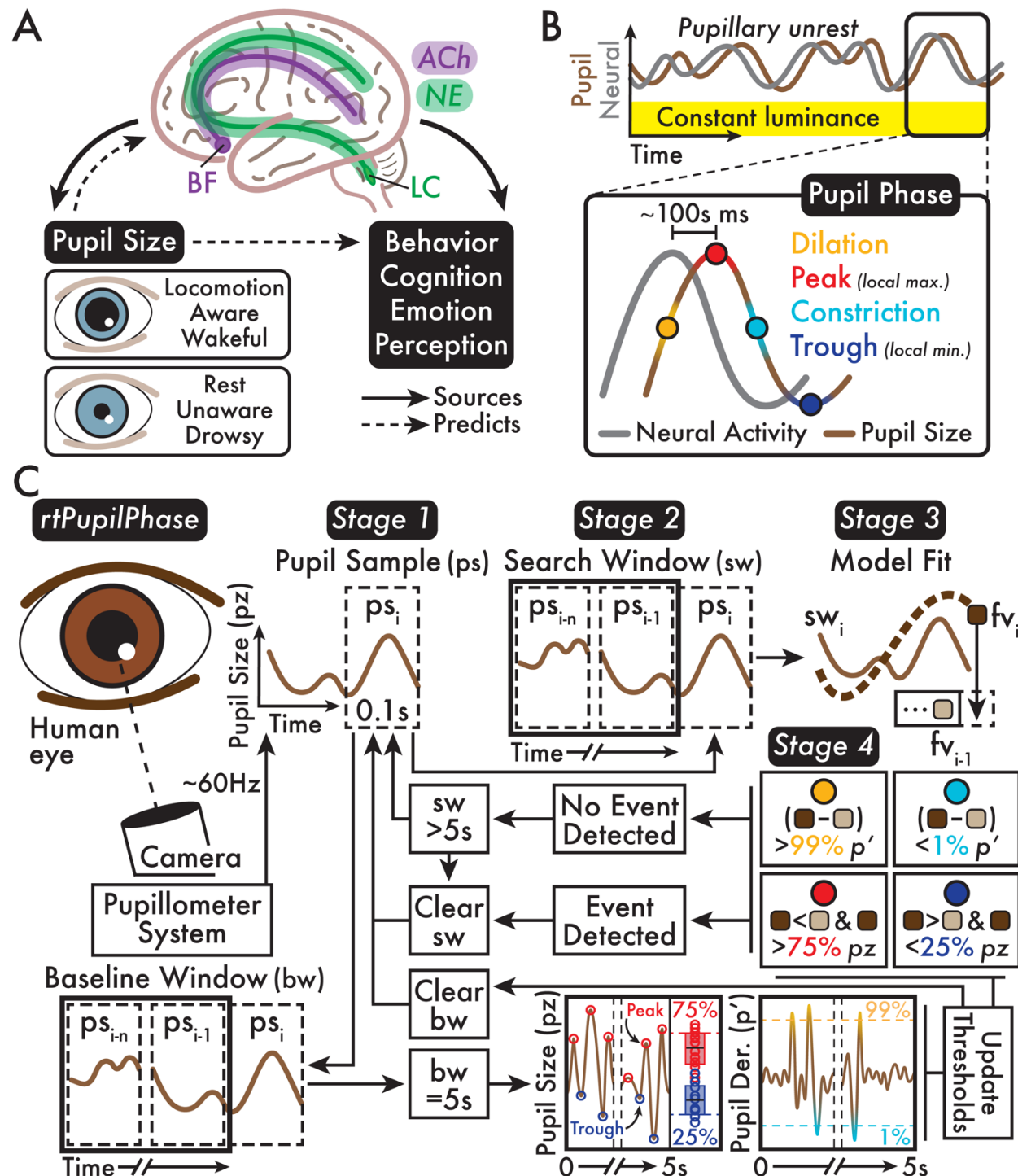


Figure 1. Neural sources of pupil size fluctuation, the phases of pupillary unrest, and the *rtPupilPhase* method schematic summary. **(A)** Major neuromodulatory networks, including the basal forebrain (BF) cholinergic system (ACh) and the locus coeruleus (LC) noradrenergic system (NE), broadly modulates subcortical and cortical activity. In addition, spontaneous and evoked ACh and NE neural activity are linked with pupil size. Therefore, pupil size can predict both large-scale brain activity and behavioral, cognitive, emotive, and perceptual states. For example, in humans, dilated pupils indicate locomotion, awareness, and wakefulness, while constricted pupils indicate rest, unawareness, and drowsiness. **(B)** *Pupillary unrest* is the spontaneous fluctuation of pupil size independent of environmental luminance. These trends in pupil size or *pupil phase* are a lagging (up to 100s of milliseconds [ms]) indicator of neural activity. The light-independent trend of pupillary unrest consists of four main phases: (1) *dilation*,

(2) *peak* or local maximum (max), (3) *constriction*, and (4) *trough* or local minimum (min). (C) The *rtPupilPhase* method is summarized in four stages. *Stage 1*: Live streamed pupillometry data (~60 Hz sampling rate; see *Human Pupillometry Acquisition Methods* section) is gathered into *pupil samples* (ps) of 0.1-second intervals. *Stage 2*: The ps is added to a *search window* (sw) and *baseline window* (bw). *Stage 3*: The sw data is modeled with a quadratic fit. The final pupil size value (fv) from the fitted sw model is recorded. *Stage 4*: A pupil phase event is determined by comparing the fv from the current and previous sw and pupil size and pupil size derivative thresholds that are updated approximately every 5 seconds (s) according to analysis of the bw. Dilation: fv_i minus fv_{i-1} is *greater than* the dilation pupil derivative threshold (p'); peak: fv_i is *less than* fv_{i-1} and fv_i is *greater than* the peak pupil size (p_z) threshold; constriction: fv_i minus fv_{i-1} is *less than* the constriction pupil derivative threshold (p'); trough: fv_i is *greater than* fv_{i-1} and fv_i is *less than* the trough pupil size threshold. The sw is reset whenever an event was detected or the sw exceeds 5 s. See the *Real Time Detection of Pupil Phase Events Methods* section for full details.

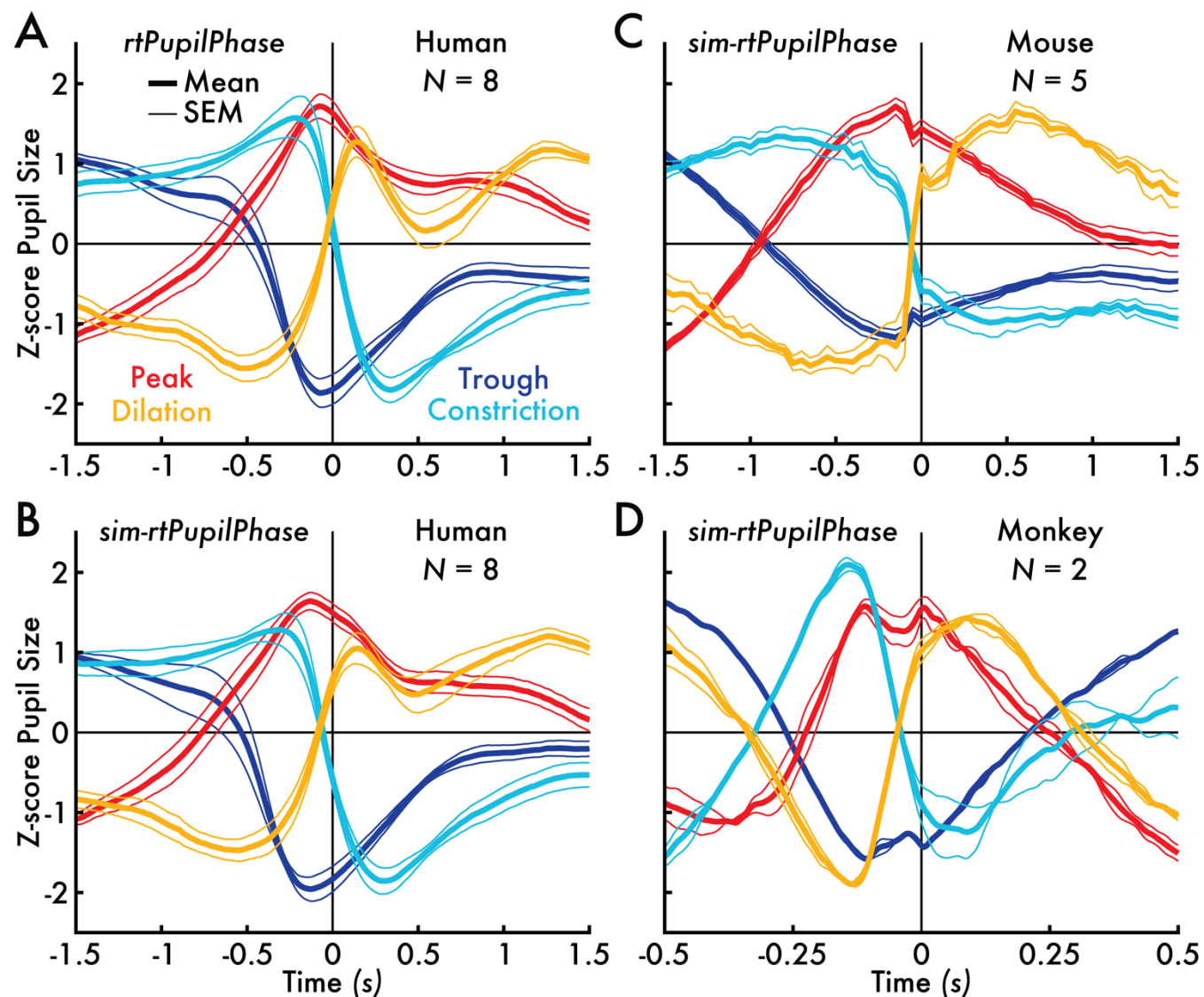
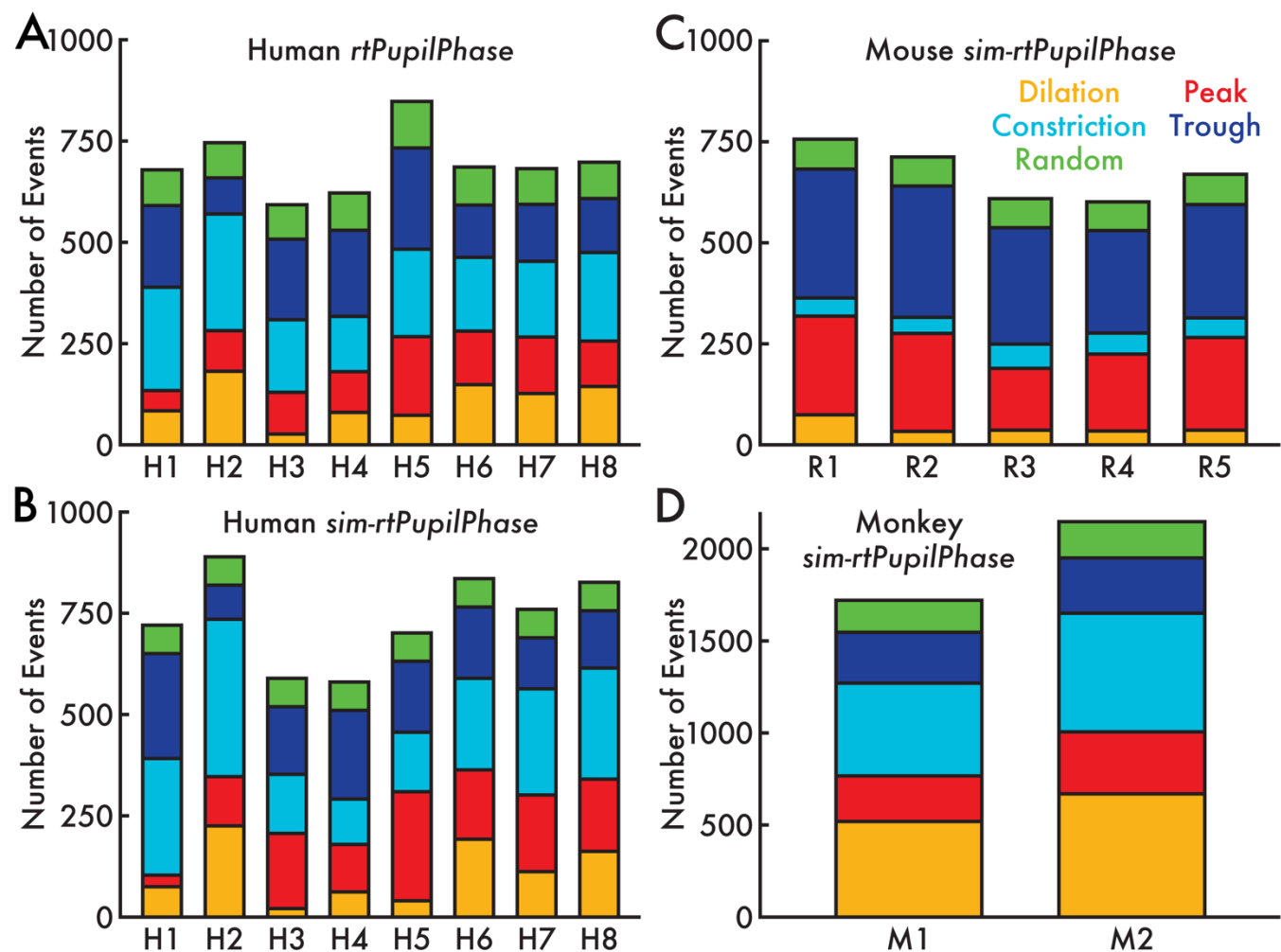


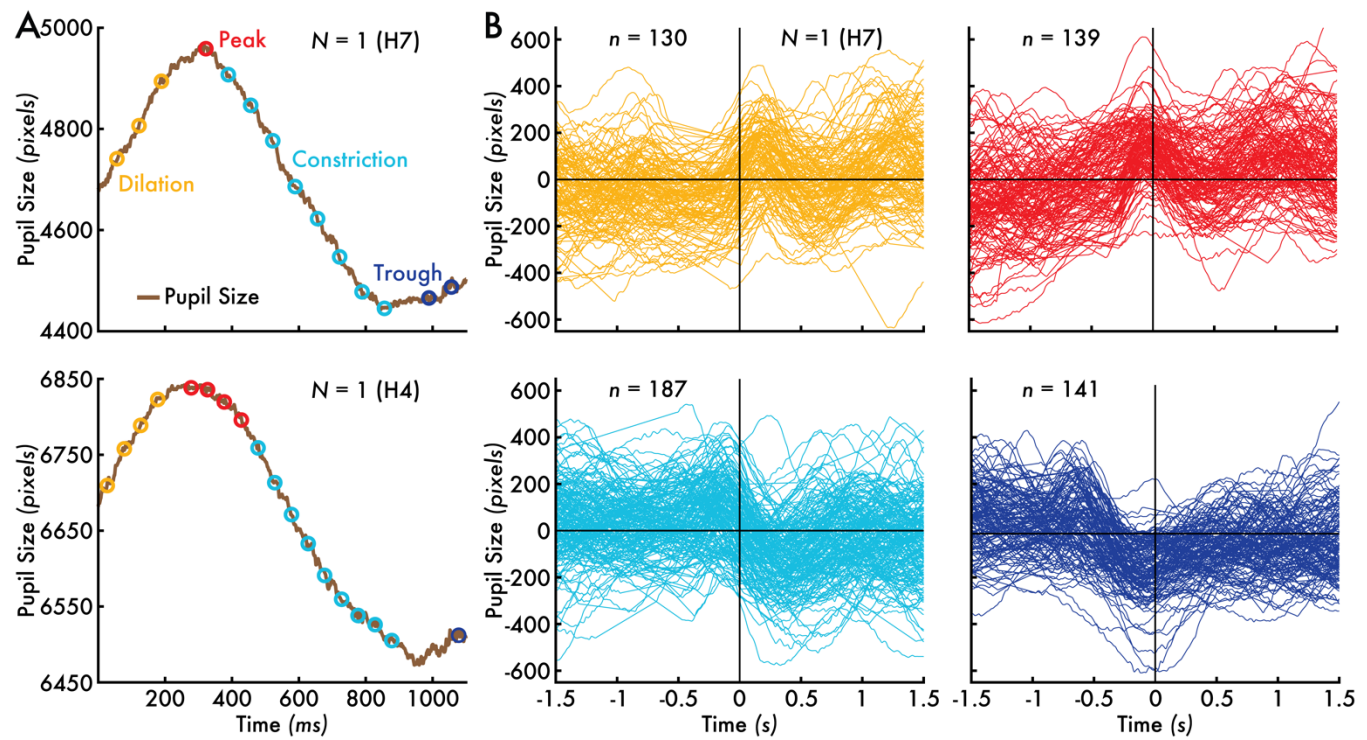
Figure 2. Human, mouse, and monkey mean z-score pupil size timecourses centered on pupil phase events detected with *rtPupilPhase* and *sim-rtPupilPhase*. **(A)** The *rtPupilPhase* human ($N = 8$) mean z-score pupil size timecourses. **(B)** The *sim-rtPupilPhase* human ($N = 8$) mean z-score pupil size timecourses. **(C)** The *sim-rtPupilPhase* mouse ($N = 5$) mean z-score pupil size timecourses. **(D)** The *sim-rtPupilPhase* monkey ($N = 2$) mean z-score pupil size timecourses. The pupil phase event time is 0 seconds (s). The group mean pupil phase event pupil size timecourse is shown in the thicker trace, bounded by thinner traces that depicts the standard error of the mean (SEM). The trace color corresponds with the pupil phase event type.

| | <i>Sampling Rate</i> | <i>Pupil Sample Duration</i> | <i>Max Search Window Duration</i> | <i>Baseline Window Duration</i> | <i>Inter-Event Interval</i> | <i>Pupil Phase Percentile Thresholds</i> |
|---|----------------------|------------------------------|-----------------------------------|---------------------------------|-----------------------------|---|
| Human (N = 8) <i>rtPupilPhase</i> | ~60 Hz | 0.1 seconds | 5 seconds | 5 seconds | 3 seconds | <i>Peak: 75%</i> <i>Trough: 25%</i> <i>Dilation: 99%</i> <i>Constriction: 1%</i> |
| Human (N = 8) <i>sim-rtPupilPhase</i> | 1000 Hz* | 0.1 seconds | 5 seconds | 5 seconds | 3 seconds | <i>Peak: 75%</i> <i>Trough: 25%</i> <i>Dilation: 99%</i> <i>Constriction: 1%</i> |
| Mouse (N = 5) <i>sim-rtPupilPhase</i> | ~20 Hz | 0.1 seconds | 5 seconds | 5 seconds | 1 seconds | <i>Peak: 75%</i> <i>Trough: 25%</i> <i>Dilation: 99%</i> <i>Constriction: 1%</i> |
| Monkey (N = 2) <i>sim-rtPupilPhase</i> | 1000 Hz | 0.1 seconds | 5 seconds | 0.5 seconds | 1.5 seconds | <i>Peak: 75%</i> <i>Trough: 25%</i> <i>Dilation: 99%</i> <i>Constriction: 1%</i> |

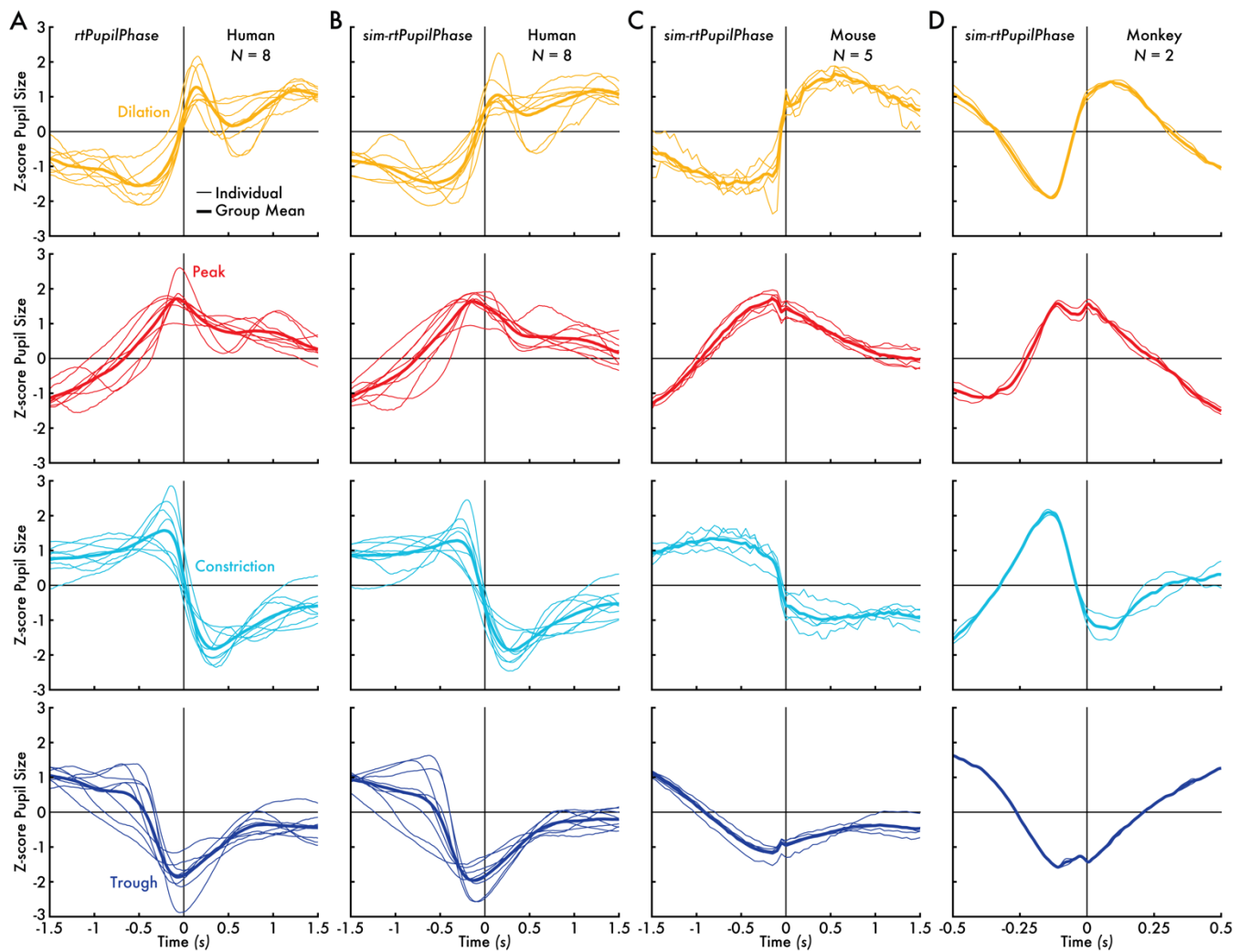
Table 1. *rtPupilPhase* and *sim-rtPupilPhase* parameters. For the monkey data set, the *sim-rtPupilPhase* baseline window duration and inter-event interval were adjusted to 0.5 and 1.5 seconds, respectively. The percentile (%) thresholds are applied on pupil size and pupil derivative data from the baseline window and used to detect pupil phase events. *Human *sim-rtPupilPhase* data were down sampled from 1000 to 60 Hz to match the approximate online sampling rate while testing *rtPupilPhase* on humans. The variable online sampling rate of the human pupillometry recordings (see *Human Pupillometry Acquisition* section) meant there was variability in the duration of the pupil sample, max search window, and baseline window. See the *Human*, *Mouse*, and *Monkey Simulated Real Time Detection of Pupil Phase* and *Real Time Detection of Pupil Phase* sections for details.



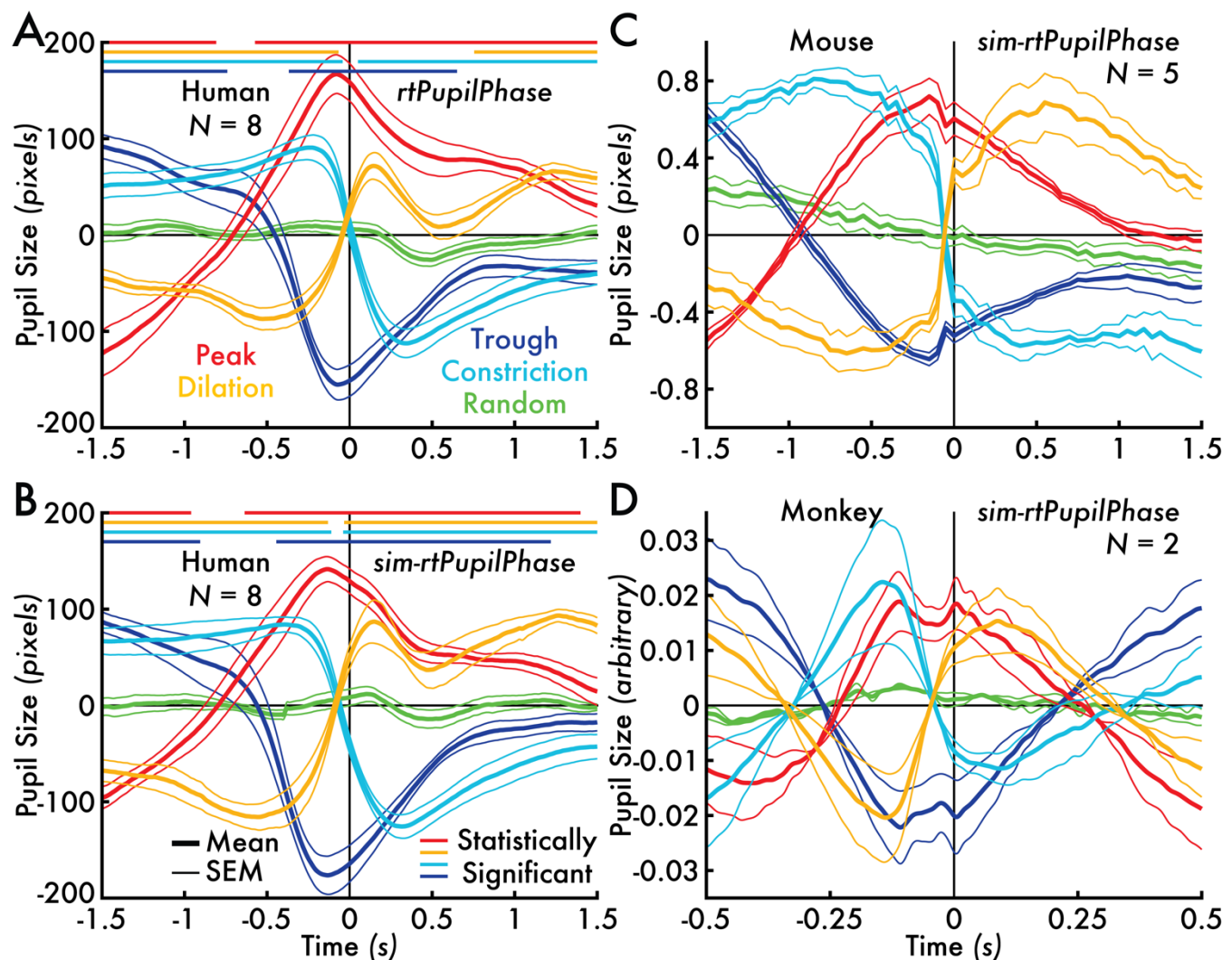
Supplementary Figure 1. Number of pupil phase and random events detected across individual humans, mice, and monkeys. **(A)** Human (H; N = 8) rtPupilPhase detected pupil phase and random events. **(B)** Human (N = 8) sim-rtPupilPhase detected pupil phase and random events. **(C)** Mouse (R; N = 5) sim-rtPupilPhase detected pupil phase and random events. **(D)** Monkey (M; N = 2) sim-rtPupilPhase detected pupil phase and random events. The depicted events are the same as those reported in Figure 2 and Supplementary Figures 2-4, 6. Modulating the parameters of rtPupilPhase (Table 1) can adjust the number of detected events. The bar color specifies the event type.



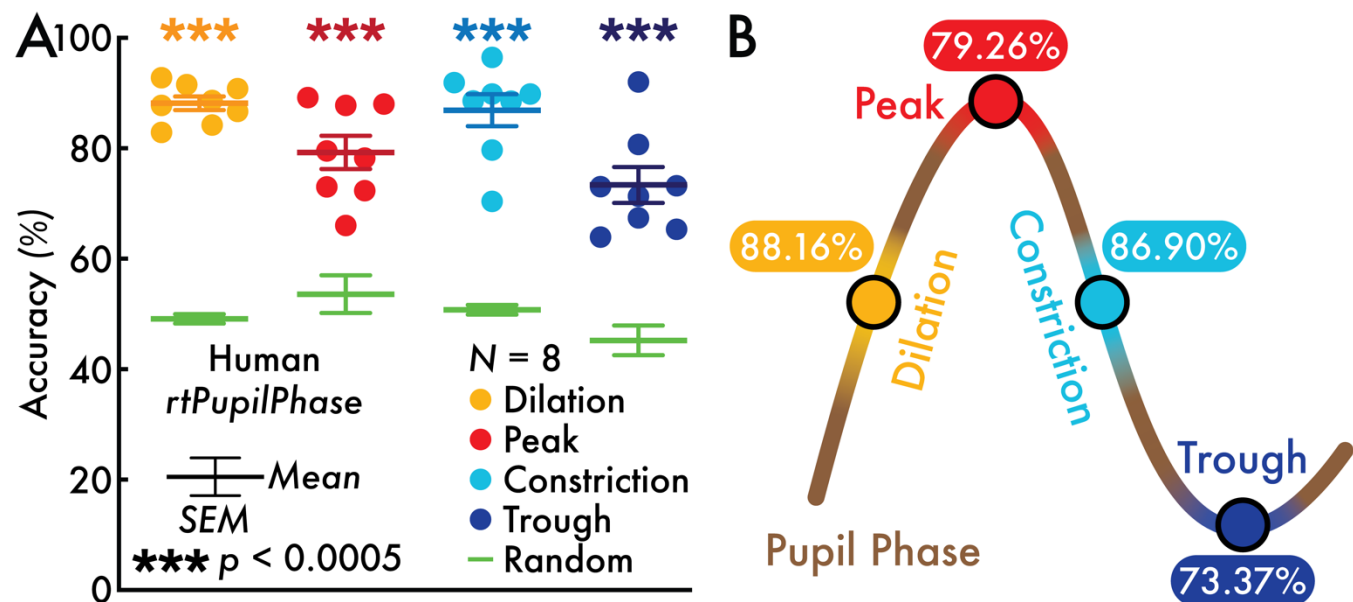
Supplementary Figure 2. Event-level *rtPupilPhase* performance. (A) Pupil size trace interval (1100 milliseconds [ms]) from two human participants (H4, H7; see Supplementary Figure 1). Open circles indicate the time of real time pupil phase events detected with *rtPupilPhase*, ignoring the inter-event interval that determines accepted events (see *Real Time Detection of Pupil Phase Events* Methods section). (B) Individual participant (H7) pupil size timecourses for each extracted epoch centered on the pupil phase event detection time (0 seconds [s]). The open circle and trace colors specify the pupil phase event type. N = number of participants; n = number of pupil phase event epochs.



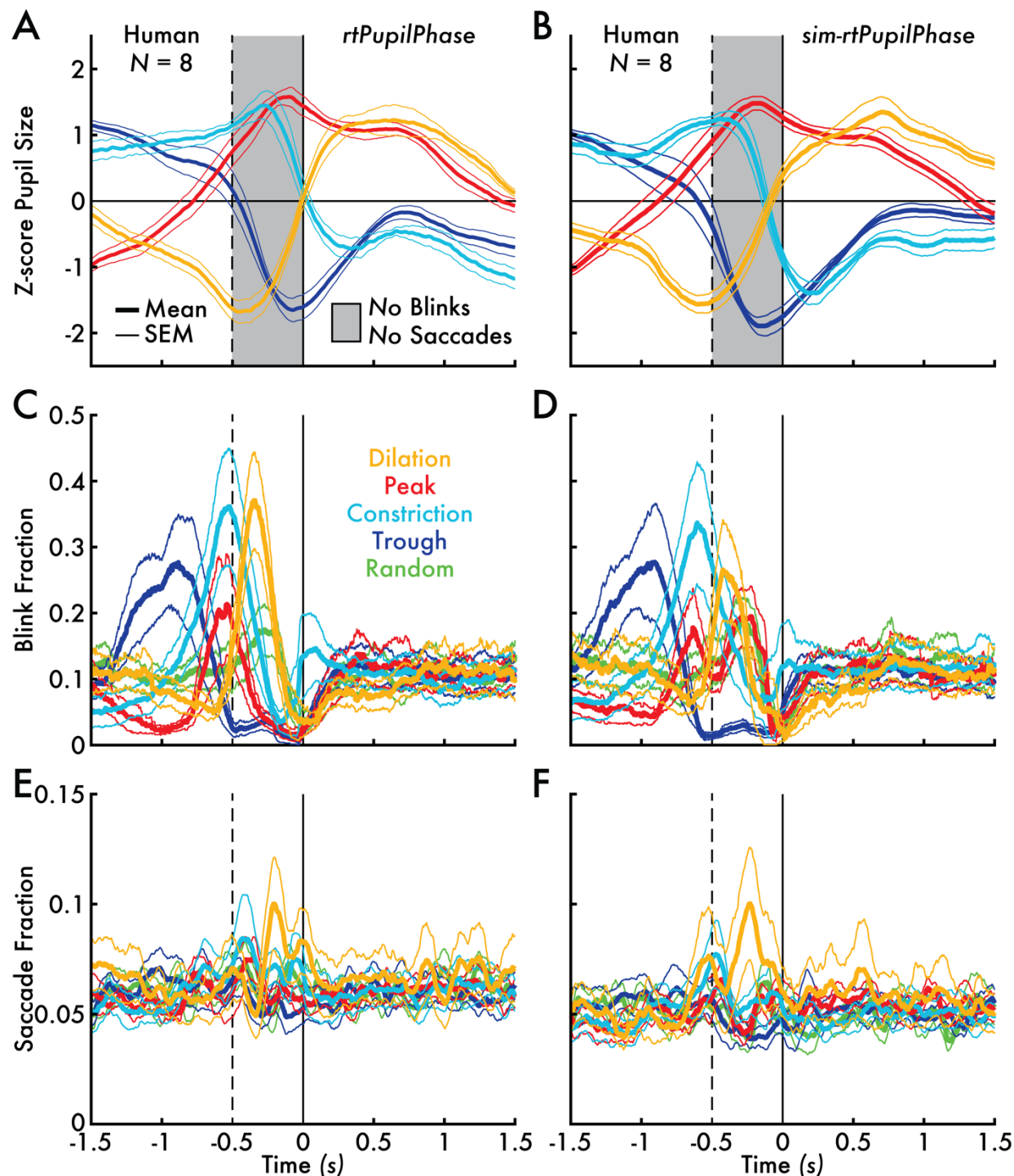
Supplementary Figure 3. Individual human, mouse, and monkey pupil size epoch timecourses for pupil phase events detected with *rtPupilPhase* and *sim-rtPupilPhase*. (A) The *rtPupilPhase* human (N = 8) individual and group mean z-score pupil size timecourses. (B) The *sim-rtPupilPhase* human (N = 8) individual and group mean z-score pupil size timecourses. (C) The *sim-rtPupilPhase* mouse (N = 5) individual and group mean z-score pupil size timecourses. (D) The *sim-rtPupilPhase* monkey (N = 2) individual and group mean z-score pupil size timecourses. The thinner lines show the individual animal (i.e., human, mouse, and monkey) pupil size timecourses. The thicker lines show the mean pupil size timecourse across all individuals within the human, mouse, and monkey data sets (i.e., the same z-score pupil size timecourses shown in Figure 2). The pupil phase event time is 0 seconds (s). The trace color specifies the pupil phase event type.



Supplementary Figure 4. Human, mouse, and monkey pupil size timecourses centered on pupil phase events detected with *rtPupilPhase* and *sim-rtPupilPhase* in native recording units. **(A)** The *rtPupilPhase* human ($N = 8$) mean pupil size timecourses. **(B)** The *sim-rtPupilPhase* human ($N = 8$) mean pupil size timecourses. **(C)** The *sim-rtPupilPhase* mouse ($N = 5$) mean pupil size timecourses. **(D)** The *sim-rtPupilPhase* monkey ($N = 2$) mean pupil size timecourses. The group mean pupil phase event pupil size timecourse is shown with the thicker line, bounded by thinner traces that depicts the standard error of the mean (SEM). The pupil phase event time is 0 seconds (s). The trace color corresponds with the pupil phase event. The horizontal-colored lines near the top of subplots (A) and (B) depict the intervals where a statistically significant difference (cluster-based permutation testing, $p < 0.05$) was found between the pupil phase and random event (see *Cluster-Based Permutation Testing Methods* section). The statistical line color indicates the tested pair (orange = dilation versus random event; red = peak versus random event; cyan = constriction versus random event; blue = trough versus random event). Statistics were not completed on the mouse and monkey data sets due to low individual animal sample size.



Supplementary Figure 5. *rtPupilPhase* detected pupil phase event performance accuracy. **(A)** The pupil phase accuracy percentage for each human participant (each circle corresponds with a participant; N = 8) is shown for the *rtPupilPhase* real time detected pupil phase events versus the true pupil phase event (see *Real Time Pupil Phase Event Detection Accuracy Methods* section). The random event accuracy relative to the true pupil phase event is shown in green (individual participant accuracy is not shown). The random accuracy percentage served as the baseline to statistically test against the pupil phase event accuracy (participant-level paired *t*-test; see *Real Time Pupil Phase Event Detection Accuracy Methods* section). All pupil phase event accuracies were statistically greater than the random event accuracy (Holm-Bonferroni corrected; $p < 0.05$; *** $p < 0.0005$). The mean and standard error of the mean (SEM) are depicted with error bars. **(B)** Summary of the *rtPupilPhase* real time detected pupil phase event mean accuracy for each pupil phase event averaged across participants – the same mean values shown in subplot (A).



Supplementary Figure 6. Human pupil size timecourses without blinks and saccades and blink and saccade fraction timecourses centered on pupil phase and random events detected with *rtPupilPhase* and *sim-rtPupilPhase*. (A) The *rtPupilPhase* human (N = 8) mean z-score pupil size timecourses excluding epochs with blinks or saccades between -0.5 and 0 seconds (s; exclusion interval is shown with a gray bar). (B) The *sim-rtPupilPhase* human (N = 8) mean z-score pupil phase event pupil size timecourses excluding epochs with blinks or saccades between -0.5 and 0 s (exclusion interval is shown with a gray bar). (C) The *rtPupilPhase* human mean blink fraction timecourses. (D) The *sim-rtPupilPhase* human mean blink fraction timecourses. (E) The *rtPupilPhase* mean human saccade fraction timecourses. (F) The *sim-rtPupilPhase* mean human saccade fraction timecourses. The pupil phase event time is 0 s. The vertical dashed line shown in subplots (C), (D), (E), and (F) depict the 0.5 s

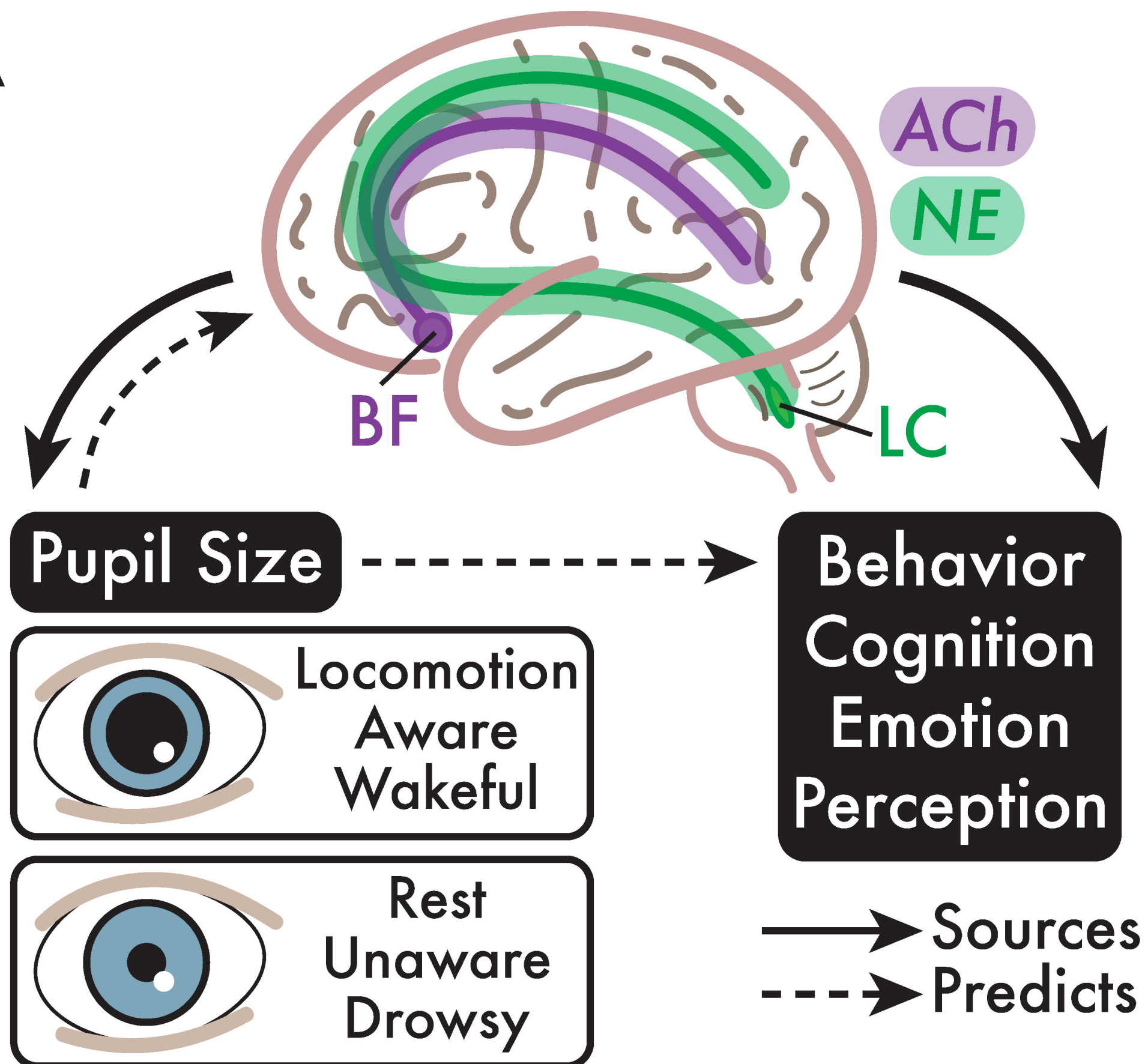
interval used to exclude pupil size epochs if any blink or saccade event were present (see subplots A and B). The group-level mean pupil size, blink, and saccade fraction timecourses for each pupil phase event is shown with the thicker line, bounded by thinner traces that depicts the standard error of the mean (SEM). The trace color corresponds with the pupil phase and random event type.

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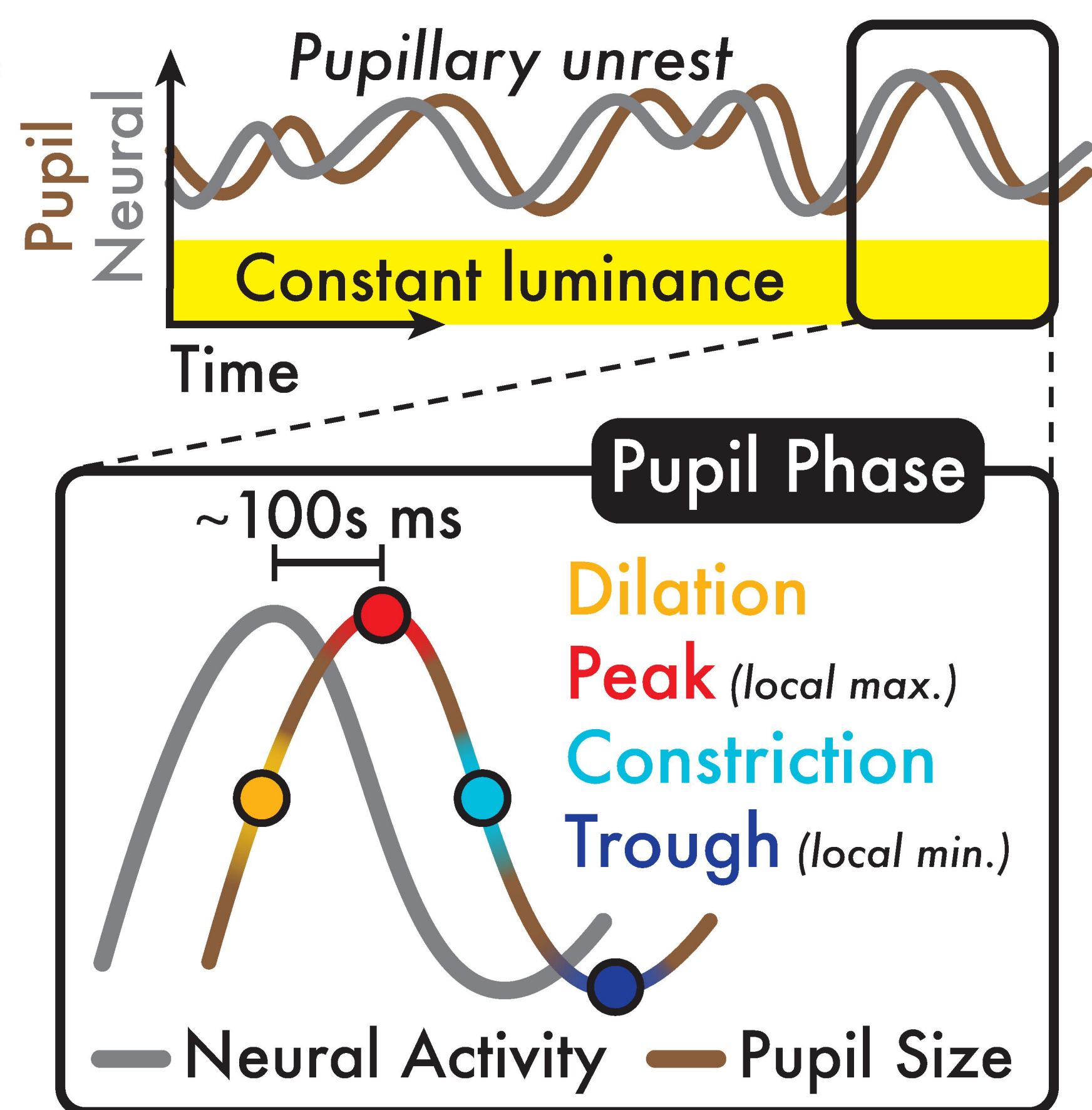
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A



B



C

