Indexing brain state-dependent pupil dynamics with simultaneous fMRI and optical fiber calcium recording

Patricia Pais-Roldán, Kengo Takahashi, Filip Sobczak, Yi Chen, Xiaoning Zhao, Hang Zeng, Yuanyuan Jiang, and Xin Yu

*High-Field Magnetic Resonance Department, Max Planck Institute for Biological Cybernetics, 72076 Tuebingen, Germany; aGraduate Training Centre of Neuroscience, International Max Planck Research School, University of Tuebingen, 72074 Tuebingen, Germany; bMedical Imaging Physics, Institute of Neuroscience and Medicine, Forschungszentrum Juelich, 52425 Juelich, Germany; and cAthinoula A. Martinos Center for Biomedical Imaging, Massachusetts General Hospital, Harvard Medical School, Charlestown, MA 02129

Edited by Marcus E. Raichle, Washington University in St. Louis, St. Louis, MO, and approved February 7, 2020 (received for review June 10, 2019)

Pupillometry, a noninvasive measure of arousal, complements human functional MRI (fMRI) to detect periods of variable cognitive processing and identify networks that relate to particular attentional states. Even under anesthesia, pupil dynamics correlate with brain-state fluctuations, and extended dilations mark the transition to more arousable states. However, cross-scale neuronal activation patterns are seldom linked to brain state-dependent pupil dynamics. Here, we complemented resting-state fMRI in rats with cortical calcium recording (GCaMP-mediated) and pupillometry to tackle the linkage between brain-state changes and neural dynamics across different scales. This multimodal platform allowed us to identify a global brain network that covaried with pupil size, which served to generate an index indicative of the brain-state fluctuation during anesthesia. Besides, a specific correlation pattern was detected in the brainstem, at a location consistent with noradrenergic cell group 5 (A5), which appeared to be dependent on the coupling between different frequencies of cortical activity, possibly further indicating particular brain-state dynamics. The multimodal fMRI combining concurrent calcium recordings and pupillometry enables tracking brain state-dependent pupil dynamics and identifying unique cross-scale neuronal dynamic patterns under anesthesia.

Neural activity fluctuates at low frequencies (<0.1 Hz) to mediate arousal or attentional states during rest (1, 2). Studying such resting-state (rs) fluctuations at different scales makes it possible to identify the neuromodulatory schemes underlying the spontaneous transitions between different brain states (3–7). Functional MRI performed during rest (rs-fMRI) can be used to extract the temporal dynamics from different neuronal populations across the brain (8–10). The temporal correlation between remote areas can be used to map baseline functional connectivity patterns that portray so-called resting-state networks, which have been associated with particular neuropsychological states and have a potential prognostic value in clinical disorders (11–13). Alternatively, transitions between distinct brain states (e.g., sleep or attention) can be tracked as direct changes in the global fMRI signal, which has also been well-reported in animal brains under general anesthesia with rs-fMRI (4, 14–22). Several human and nonhuman primate studies have demonstrated that the global fMRI signal accurately reflects the momentary level of attention or arousal (1, 23, 24), suggesting a critical role of the ubiquitous spontaneous oscillations in orchestrating the transition between brain states. Additionally, human experiments where pupillometry was added to the fMRI measurement have identified some potential neural correlates of arousal (25–27), yet the mechanisms underlying spontaneous transitions toward varying brain states remain only partially understood.

Recently, experiments performed in rats have revealed key global signatures that precede brain-state changes during anesthesia by measuring the fMRI signal in parallel to the calcium-dependent neuronal and astrocytic activity (4), which indicates the potential of multimodal fMRI platforms to be applied in behaving rodents to track different brain states. However, despite ongoing efforts to set up an awake rodent multimodal fMRI platform (28–30), the potential stress and motion-related artifacts (31, 32), summed to the lack of a robust behavioral arousal measurement inside the MRI scanner, have prevented multimodal fMRI investigations of the brain state in awake rodents. Eye blinks, unconscious eye movements, and pupil-size variations can report the level of arousal (1) and condition of task performance (7, 33, 34) and reflect the level of cognitive processing (35), attention (36), fear (37), or reward anticipation (38). Animal experiments performed at the bench (e.g., electrophysiological studies) have allowed accurately testing the relationship between pupil dilations and neural activity specific to particular brain areas. Pupil fluctuations have been repeatedly linked to the activity of neurons in the noradrenergic locus coeruleus (LC) (39, 40) and mark cortical state changes (33, 34, 41–44), which places the pupil as an affordable arousal indicator to merge with other brain imaging schemes. These studies have linked pupil dilations to desynchronized cortical activity and depolarized states during walking-related arousal (33, 34, 41).

Significance

Resting-state fMRI (rs-fMRI) performed in animals allows investigating the spatial correlation patterns of brain function, which presents diverse oscillatory features in varied brain states, even under anesthesia. However, the link between global fMRI signal oscillations and fluctuations in the level of arousal, which can be indexed by spontaneous pupil dilations, is seldom investigated in rodent brains. Here, we established pupillometry with rs-fMRI and concurrent cortical calcium recordings in anesthetized rats, enabling brain-state fluctuation analysis across multiple scales. A unique positive correlation with pupil dilations was detected near noradrenergic cell group 5 (A5) in the ventral brainstem, which was associated with a specific coupling between spectrally different calcium waves, highlighting unique brain-state dynamics.


The authors declare no competing interest.

This article is a PNAS Direct Submission.

Published under the PNAS license.

Data deposition: All fMRI datasets, as well as the synchronized calcium recordings and pupil-size vectors, reported in this paper have been deposited in Zenodo at https://doi.org/10.5281/zenodo.3677925.

1To whom correspondence may be addressed. Email: xy9@mhg.harvard.edu.

This article contains supporting information online at https://www.pnas.orglookup/suppl/doi:10.1073/pnas.1909937117/-/DCSupplemental.

www.pnas.org/cgi/doi/10.1073/pnas.1909937117

PNAS Latest Articles | 1 of 8
Pupillometry can be especially informative during anesthesia, where motor-driven behavioral measurements are not accessible to the experimenter. Kum et al. (43) showed that anesthetic washout (i.e., emergence from anesthesia) is accompanied by burst activity and mydriasis in rats, suggesting that lighter unconscious states during anesthesia could be tracked by measuring the pupil size.

Despite existing rodent studies combining electrophysiological measurements with pupillometry, to date there are no studies merging pupil dynamics with the activity of neuronal populations and concurrent whole-brain fMRI in rats. The goal of this work was to merge the bench measures with fMRI to create a platform that allows investigating the link between spatiotemporal changes in brain activity and transitory brain states that can be tracked with pupillometry. Here, GCaMP-mediated calcium signal changes were recorded from cortical neurons in parallel to whole-brain fMRI in anesthetized rats (4, 21) subjected to concurrent pupillometry. Here, GCaMP-mediated calcium signal changes were recorded from cortical neurons in parallel to whole-brain fMRI in anesthetized rats (4, 21) subjected to concurrent pupillometry. Based on the rs-fMRI signal and its correlation with pupil dilations, we identified the presence of oscillatory brain states under anesthesia, which were specifically represented by neuronal calcium dynamics. Besides tackling the technical challenge of performing pupillometry in small-animal MR scanners, we demonstrate the potential of this platform to investigate the mechanisms underlying brain-state transitions across multiple scales.

Results

Concurrent fMRI and Neuronal Calcium and Pupil Dynamics Track Brain-State Changes in the Anesthetized Rat. We used a small MRI-compatible camera to track pupil dynamics from the rat eye in parallel to fMRI and obtain a measure of the brain-state fluctuation during anesthesia (SI Appendix, Fig. S1). The measure of pupil-size changes was introduced in a multiscale analysis workflow as illustrated in Fig. 1. Three main recordings were obtained in parallel: 1) video recording of the pupil, 2) optical fiber calcium recording (in the cingulate cortex, as well as in the barrel cortex in some rats), and 3) whole-brain fMRI imaging. First, the vector representing the pupil-size fluctuation was generated from the acquired video (45) (Fig. 2, Top trace, SI Appendix, Fig. S2, and Movie S1) (see SI Appendix, Methods for details). This vector of pupil dynamics was correlated with each fMRI voxel time course to obtain a pupil–fMRI correlation map (Fig. 3A). In addition, another correlation map was created by comparing the fMRI time courses with the first derivative of the pupil size (1st der.; i.e., pupil dilations) (Fig. 3B). Second, the pupil dilation–fMRI correlation map, considered a pupil-relevant spatial template, was spatially correlated with the original 4-dimensional fMRI (45) (i.e., a time-varying volume) to create a specific spatiotemporal correlation time course that provided an fMRI index of the brain state (Fig. 4A and Movie S2), similar to what has been shown in previous rs-fMRI arousal studies on nonhuman primates (1). Third, the pupil dynamics were studied in relation to the calcium signal fluctuation from the cortex (45), in particular the 2- to 3-Hz activity band, which showed the strongest power throughout the calcium spectrum (Figs. 2, Bottom, and 5C and SI Appendix, Fig. S3, Lower Right) (the spectral analysis of the calcium signals is explained in SI Appendix, Methods). Finally, the time course of the fMRI-inferred brain-state index was compared with the calcium dynamics from the cingulate cortex (Fig. 4D and E). SI Appendix, Fig. S3 shows a summary pipeline with the main steps followed in this work. The relationship between the fMRI signals and the fMRI-inferred brain-state time course, as well as the pupil and calcium dynamics, is exemplified in Movie S2. Fig. 2 shows the brain-state fluctuation detected with the multimodal platform in a representative animal under anesthesia (another example is shown in SI Appendix, Fig. S4). Alternation between brain states was evidenced by the transition from constricted to dilated pupil or vice versa, which showed a strong correlation with the whole-brain fMRI signal fluctuation and with the neuronal calcium signal oscillation. It is noteworthy that the brain-state dynamics varied across different trials. This dynamicity can be presented by the index of pupillary unrest (PUI), which measures the variability in pupil size per run (SI Appendix, Methods) (46). The PUI was positively correlated with the SD of the pupil size, showing 43% of the total variance of the pupil size (n = 71 of 10 rats) with a PUI above 1,000, denoting salient low-frequency oscillatory patterns (SI Appendix, Fig. S5). Together, the acquired measures indicate that the multimodal platform merging fMRI, pupillometry, and calcium imaging enables a cross-scale investigation of the varying brain states in anesthetized animals.

Correlation Analysis of Whole-Brain fMRI with Pupil Dynamics in Anesthetized Rats. Simultaneous fMRI and pupillometry allowed us to compute 2 pupil-correlated fMRI maps in the anesthetized...
state. The map in Fig. 3A shows the voxel-wise correlation between the pupil-size time course and the fMRI signals, indicating a negative correlation in most of the cerebral cortex, thalamus, septal nuclei, and superior colliculus, some of which have been previously identified as neural correlates of pupil dilations during wakefulness [i.e., cingulate cortex or superior colliculus (47, 48)]. While the pupil size correlates with the arousal level during wakefulness (26, 40), specific brain regions relevant to the switch between arousal states can be potentially assessed by correlating the fMRI signal with the first-order derivative of the pupil-size dynamic time course (26, 38, 49); therefore, we also performed correlation analysis between blood oxygen level-dependent (BOLD) fMRI and 1d-pupil (e.g., the purple trace in Fig. 2). The resulting 1d-pupil-fMRI map showed that the areas exhibiting negative correlation with the pupil size were also negatively correlated with 1d-pupil and, in addition to it, an area in the brainstem was found positively correlated with the 1d-pupil time course, which was not detected in the pupil-size correlation map (Fig. 3). The resulting 1d-pupil-fMRI map showed that the areas exhibiting negative correlation with the pupil size were also negatively correlated with 1d-pupil and, in addition to it, an area in the brainstem was found positively correlated with the 1d-pupil time course, which was not detected in the pupil-size correlation map (Fig. 3). The positively correlated brainstem overlapped with the rostral A5 area (Fig. 3 C and D), which contains primarily noradrenergic cells and projects to the major subcortical arousal nuclei through the reticular formation pathway (50). It is worth noting that no significant positive correlation was observed in other subcortical nuclei involved in arousal regulation, for example the LC (39) or the central and mediodorsal thalamic nuclei (4, 51).

To study the temporal relationship between the fMRI and the pupil dynamics, the whole-brain fMRI time courses were also correlated with a lagged version of the pupil size or 1d-pupil. This analysis resulted in 2 sets of maps (SI Appendix, Fig. S6A), which indicated that the areas correlating negatively with the pupil size did so at short lags (i.e., faster hemodynamic response functions [HRFs]), with longer lags reducing their correlation value, while the same areas were well-correlated with 1d-pupil at all lags. Importantly, the positive correlation in the brainstem, only present in the 1d-pupil correlation map, was mainly observed at short lags (using fast HRFs). In order to investigate the temporal dynamics of the positively correlated area in the brainstem with respect to the negative global signal in the context of brain-state regulation, we averaged the fMRI time courses from both functional clusters and performed cross-correlation analysis with respect to the pupil size or to 1d-pupil. The positive cluster (brainstem) was most correlated with 1d-pupil at a 0-s lag (pink wave in SI Appendix, Fig. S6B; first derivative). In contrast, the negative cluster exhibited its maximum correlation 5 s later, in agreement with the voxel-wise correlation results. The results of this analysis indicated that the brainstem activates during pupil

![Fig. 2. Tracking brain-state changes with pupillometry, fMRI, and calcium imaging. The traces, map, and spectrum show an example of the measures obtained concurrently during a 15-min scan in an anesthetized rat. Note the negative correlation between 2-3-Ca and the tv-fMRI-bs, and between 2-3-Ca and the pupil diameter. The inset over the power spectrum decomposition of the calcium signal (Bottom) shows an averaged cross-correlation between pupil-size changes and 2-3-Ca ±SD across trials. a.u., arbitrary units.](image-url)
dilation but not necessarily after the pupil has increased its size, which coincides with the globally reduced BOLD fMRI signal that fluctuates with pupil-size changes. To represent a brain state more timely correlated with pupil dynamics, we are using the 1d-pupil-fMRI correlation map as a template in the successive steps of our analysis.

**Pupil-fMRI-Derived Index to Measure the Brain State.** As shown in Fig. 4 and Movie S2, the momentary pupil-linked brain state can be inferred from each acquired rs-fMRI volume by calculating its spatial correlation with the 1d-pupil-fMRI correlation map. Concatenation of the spatial correlation values produces a vector of the time-varying fMRI-inferred brain state (tv-fMRI-bs) (1). To specify the neuronal correlates of the inferred brain-state fluctuation, the power profile of the 2- to 3-Hz calcium fluctuation (2-3-Ca) was cross-correlated with the tv-fMRI-bs and with the varying pupil size, showing strong correlation features with both signals (Fig. 4 E and G). The tv-fMRI-bs was positively correlated with the pupil dynamics (Fig. 4 B and C) but showed a negative correlation with the 2-3-Ca oscillation (Fig. 4 D and E); cross-validation is shown in SI Appendix, Fig. S7). In addition, the neuronal calcium recordings from cingulate and barrel cortex in our anesthetized rats covaried (negatively) with pupil-size changes, showing a decrease of the 2-3-Ca upon dilations (Fig. 4 G and H and SI Appendix, Fig. S8), consistent with the negative correlation observed from the same cortical region in the pupil-fMRI correlation map. The salient correlation features observed across multiple trials between the tv-fMRI-bs, the calcium oscillation from the cortex, and the pupil dynamics present a generalized 3-way interaction scheme to elucidate brain-state changes in anesthetized rats (Fig. 4 F).

**Identification of Brain State-Dependent Pupil Dynamics under 2 Distinct Cortical States under Anesthesia.** The calcium signal recorded from the cortex exhibited 2 main components: a slowly fluctuating baseline (<0.5 Hz) and oscillations within the frequency range typical of the electroencephalogram (EEG) (strongest at 2 to 3 Hz, as shown in Figs. 2 and 5C and SI Appendix, Fig. S3). In particular, the 2- to 3-Hz calcium transients have been verified with simultaneous local field potential and multunit activity recordings showing a strong correlation (Fig. 5 B, E, and G). Analysis of the neuronal activity at both frequency ranges provided an additional dynamic feature to depict the neurophysiological state of animals. A subset of trials showed strong cross-frequency coupling between the baseline and 2-3-Ca (Fig. 5H, red dots) (21); however, ~40% of the trials demonstrated poor correlation or uncoupled cross-frequency calcium dynamics (Fig. 5H, black dots), similar to the dissociation of phase and amplitude coupling observed in previous EEG studies (52, 53). The 1d-pupil-fMRI correlation maps acquired from both groups independently showed a negative correlation through most of the brain (Fig. 5I), similar to the correlation map acquired from all trials together (Fig. 3B). However, the positive correlation in the brainstem was significantly higher in the calcium-uncoupled group (Fig. 5F). Interestingly, the SD of the pupil was also higher in calcium-uncoupled trials (SI Appendix, Fig. S9C, dots outside the red box), indicating a brain state-dependent pupil dynamic behavior. In all, the assessment of cross-frequency coupling from the calcium data suggested the existence of distinct modes of brain-state dynamics under anesthesia, also supported by the opposite correlation between the calcium baseline and the tv-fMRI-bs in the 2 identified groups (SI Appendix, Fig. S9D), which is consistent with the EEG observed during brain-state switches under anesthesia (54, 55). Altogether, our data indicate that the cross-frequency coupling of the calcium signal constitutes a potential marker to further differentiate between periods of stronger or weaker pupil control during the dynamic brain state.

**Discussion**

In this work, simultaneous whole-brain fMRI and optical fiber-mediated neuronal calcium recordings were acquired to specify brain state-dependent pupil dynamics in anesthetized rats. Cross-correlation...
analysis of the whole-brain fMRI with pupil dilation, in particular the first derivative of pupil size, demonstrated global negative correlation throughout the brain but specific positive correlation with an area overlapping with noradrenergic cell group A5 located in the brainstem. This ventral pontine activation during pupil dilation was more salient in specific brain states showing uncoupled 2- to 3-Hz and baseline calcium dynamics than in brain states with cross-frequency coupling. In summary, we identified the time-varying fMRI-inferred brain states with concurrent neuronal calcium dynamics, providing distinct correlation patterns to the pupil dynamics in the anesthetized rat brain. Our results suggest that the pupil dynamics observed during anesthesia in rats are directly linked to the global fMRI signal fluctuation (21, 23, 24, 56), as well as to the cortical population activity observed as neuronal calcium transient oscillations, in particular, the 2-3-Ca (Fig. 2, Bottom graph, Fig. 4 G and H, and SI Appendix, Fig. S8) (34). A relationship between pupil size and cortical activity, presumably linked to arousal, has been reliably observed in mice during wakefulness and during non-REM (rapid eye movement) sleep (34, 41, 44). These animals also transitioned between periods of high-amplitude and low-frequency neuronal firing coupled to constricted pupils and periods of low-population activity (i.e., more desynchronized states) linked to pupil dilations, in agreement with our observations. Also, the switch from desynchronized (low-amplitude fast activity) to synchronized (high-amplitude slow waves) EEG in rats anesthetized with isoflurane (57) or urethane (58) has been previously associated with the transition from pupil dilation to constriction, and vice versa (43).

We report 2 findings based on the pupil dynamics–fMRI correlation analysis. One is the global negative fMRI signal related to the pupil dilations (Fig. 3). The observation of cortical regions anticorrelated with the pupil dynamics is in agreement with the arousal-based rs-fMRI signal correlation patterns obtained from unanesthetized nonhuman primates with eye-opening/closing tracking (1). The reduced global BOLD signal observed during pupil dilations in our anesthetized animals coincided with suppressed calcium transient activity from the cortex (2- to 3-Hz calcium fluctuations; Fig. 4D), which is in agreement with previous reports of lightly anesthetized rats (21). However, in contrast to the previously reported anticorrelation between cortical and subcortical functional dynamics during vigilant brain states (1, 59–62), the pupil-based fMRI correlation maps in our study showed a rather global negative correlation feature spread over both cortical and subcortical regions (Fig. 3), including the thalamus, which has been reported to be anticorrelated with the cortex in rs-fMRI studies at varied arousal states (63, 64). Wang et al. (4) observed a positive BOLD response in the midline thalamic nuclei coupled to specific astrocytic calcium transients preceding a massive global decrease of the BOLD signal in anesthetized rats. The lack of detection of this fast-positive thalamic BOLD signal from the pupil-based fMRI correlation map may be caused by the largely varied indices of pupil unrest across the different trials (SI Appendix, Fig. S5). In addition, the massive global negative BOLD signal may overshadow the phasic positive contribution from the thalamus. A future study integrating concurrent astrocytic calcium signal recording into the multimodal fMRI platform will attempt to further investigate the anticorrelation features of the cortex and thalamus relevant to the brain state.

The second finding is the robust positive correlation observed in a location compatible with noradrenergic cell group A5 from the 1d-pupil–fMRI correlation map (Fig. 3D). The neurons of this ventral pontine nucleus send projections to several midbrain areas involved in arousal and vital functions, including the periaqueductal gray and the parabrachial nucleus (50, 65), and are suppressed during REM-like states induced by the acetylcholine agonist...
Calcium coupling-dependent cortical states and associated pupil-fMRI patterns. (A) The calcium signal can be decomposed into slow (baseline) or relatively fast fluctuations (e.g., 2 to 3 Hz). (B) Magnified in a red rectangle are 25 s extracted from a 30-min recording of a GCaMP-based calcium fluorescent signal (in blue) and local field potentials (LFP, in black) recorded simultaneously in a representative animal in the absence of stimulation. (C) Power spectrum of the calcium signals of animals undergoing fMRI (average of 64 trials ±SD). (D) Average calcium peak. (E) Average multiunit activity (MUA) during calcium spikes (the gray curve represents the average calcium spike, for comparison). (F) Average LFP peak. (G) Correlation between LFP and calcium spikes. (H) Cross-correlation between the 2- to 3-Hz band and the baseline (red/black dots: coupled/uncoupled trials). (I) Left) Maps showing the voxel-wise correlation between LFP and calcium spikes. (J) Right) Two coronal slices covering the brainstem and a sagittal slice are magnified from both maps to show the differences in A5 correlation values between both groups. The third vertical panel shows the results of a 2-group t-test run on the 1d-pupil-fMRI correlation map, for trials with uncoupled (left map) or coupled (right map) calcium signals. (K, Right) Two group t-test of the 1d-pupil-fMRI correlation map, for trials with uncoupled (left map) or coupled (right map) calcium signals. (L, Right) Two group t-test of the 1d-pupil-fMRI correlation map, for trials with uncoupled (left map) or coupled (right map) calcium signals.

Carbachol and the alpha-2 receptor clonidine (66), which supports its potential role related to arousal-state fluctuation and pupil-size regulation. Similar to the LC, the A5 noradrenergic area may play an alternative regulatory role in arousal, as suggested from the multiple studies associating noradrenaline with brain-state modulation (67–69). Although brain-state fluctuation and pupil dynamics have been reported to be regulated through noradrenergic projections from the LC (39, 40), we observed no inhibition (the fMRI signal of the A5 area increases in parallel with pupil dilations; SI Appendix, Fig. S6B), and there was a higher correlation of the calcium baseline with dissociated. Interestingly, in those uncoupled trials, the pupil-correlation patterns in the LC or its projection areas. This fact may indicate that the temporal features of pupil dynamics did not serve as a strong linear regressor to fit the neuronal activity-coupled fMRI signal particular to the LC in the anesthetized brain (70, 71). Importantly, to date, the existence of a causal link between the LC and pupil modulation remains to be clarified (49). Further strategies (e.g., noradrenergic-specific cell targeting) will be needed to confirm and characterize the unexplored relationship between the A5 group and pupil dynamics. Additionally, similar to the specific astrocytic-dependent fMRI activation pattern deciphered in anesthetized rats (4), knowledge about the specific dynamic signaling underlying pupil dilations may be needed to identify functional nuclei that may modulate pupil dynamics with particular firing and neurovascular coupling patterns. Also noteworthy is that the A5 area is mostly positively correlated with the first derivative of the pupil rather than the direct pupil size (Fig. 3C), which indicates that brainstem activation is more relevant to the change than to the maintenance of the brain state. Cross-correlation of the fMRI signals with 1d-pupil indicated that brainstem activation occurs earlier than global inhibition (the fMRI signal of the A5 area increases in parallel with pupil dilations; SI Appendix, Fig. S6B), and suggests that the latter might be triggered by the first one; however, a causal link between A5 activation and global cortical/subcortical suppression should be investigated (e.g., with optogenetic experiments).

The concurrent calcium signal acquired with fMRI represents a key advantage to measuring cross-modal brain dynamics. The GCaMP6f-mediated calcium transients, detected with optical fibers, constitute a reliable indicator of neuronal activity (Fig. 5 B, D, E, and G), and their spectrogram presents power fluctuation estimates similar to EEG (4, 21). In the present study, 60% of the trials showed a tight coupling between baseline and 2-3-Ca fluctuations, while in 40% of them the calcium dynamics were dissociated. Interestingly, in those uncoupled trials, the pupil-based fMRI correlations were stronger throughout the brain and there was a higher correlation of the calcium baseline with the tv-fMRI-bs (SI Appendix, Fig. S9D). We observed different coupling from different trials of the same animal and through several animals, which rules out the influence of the optical fiber tip location on the calcium dynamics (i.e., excludes the possibility
that the calcium baseline is determined by the proximity of the implanted optical fiber to a vessel). Also, the brain state-dependent cross-frequency coupling features cannot be simply caused by the blood effect (i.e., the BOLD signal) on the optical fiber-mediated calcium recording. In particular, the largely varied global fMRI signal fluctuation did not match the specific distribution of the calcium cross-frequency coupling, in contrast to the pupil dynamic changes (SI Appendix, Fig. S9C). This suggested that calcium (un)coupling is associated with brain-state dynamics and does not merely emerge as a result of physiological noise. Specifically, the increased correlation between baseline calcium fluctuations and the fMRI-inferred brain-state vector may be related to spontaneous astrocytic activity, which is known to mark the transition between brain states (4, 55). A previous study also reported an increase of ultrasonic electrophysiological activity in conjunction with suppression of higher-frequency bands during arousal switches in mice (55). Our work, involving trials with diverse brain-state dynamics (SI Appendix, Fig. S5), has identified different calcium-related conditions that are in agreement with the findings encountered separately in previous reports (21, 55, 56). It is likely that changes in the calcium coupling indicate a shift in the mode of cortical processing (72), however, the meaning and underlining cause of the coupling-dependent states need to be further investigated. The desynchronization between both calcium bands could be associated with arousal pathways emerging at the ventral pons (likely noradrenergic group A5) that may trigger pupil dilations while enhancing global reduction of population activity (2-3-Ca), mostly in cortical neurons. This global regulatory scheme may be necessary to prevent cortical activation from external sources during internal arousal. The dissociation between 2- to 3-Hz and baseline oscillations may reflect a distinct recruitment of both generators (the ultraslow and the 2-3-Ca oscillator) during the arousal fluctuation; however, the source of those cortical dynamics is not sufficiently understood and, hence, any hypothesis remains speculative. Given that all subjects of our study were anesthetized, the degree to which these brain-state changes can be investigated from the data obtained here is limited. Future studies conducted in awake animals should confirm whether these observations can be extrapolated to more aroused brain states that occur during conscious awareness.

Inferring brain-state changes (e.g., fluctuations in the level of arousal) during anesthesia might seem counterintuitive. Arousal has been defined, electrophysiologically, as a state of depolarized neuronal membranes and desynchronized firing (73, 74), also characterized by wide pupils (33, 75), and it is usually assessed during quiet wakefulness or in the context of behavioral performance. Extremely low and extremely high arousal levels yield poorer task performance compared with middle-arousal states, a phenomenon described as an inverted-U relationship (33, 76, 77). Under anesthesia, the brain state is restricted by a maximum arousal level (arousal) during anesthesia might seem counterintuitive. Arousal pathways emerging at the ventral tegmental area, a procedure which has been shown to lead to emergence from general anesthesia (81). Although the meaning of brain-state oscillations remains debatable, together our results suggest that anesthesia is not an isolated unconscious condition but a dynamic oscillatory brain state that can be traced at different scales.

Conclusion

Here, we identified a qualitatively negative fMRI spatial pattern that was correlated with pupil dilation and served as a template to produce a brain-state fMRI index during anesthesia. The neurobiological relevance of the linkage between the time-varying fMRI-inferred brain state and the pupil dynamics was verified with concurrent recording of neuronal calcium oscillations in the cingulate cortex. In addition, cross-correlation analysis of the calcium baseline and the 2- to 3-Hz power fluctuation revealed 2 distinct neurophysiological states based on the cross-frequency (un)coupling conditions of the calcium signals. While the global negative pupil-fMRI correlation was present under both states, a positive correlation of the ventral pons was highly linked with uncoupled neuronal calcium dynamics. Altogether, we demonstrate that the pupil size can be tracked in animals undergoing fMRI, allowing to correlate whole-brain activation patterns with particular brain state-dependent pupil dynamics.

Materials and Methods

Ten adult rats were used to acquire parallel pupillometry, cortical calcium recordings, and whole-brain fMRI. The study was performed in accordance with the German Animal Welfare Act (TierSchG) and Animal Welfare Laboratory Animal Ordinance (Versuchstierhaltungsdirektive der Verwaltungspräsidenten, Berlin). This is in full compliance with the guidelines of the European Union (EU) Directive on the protection of animals used for scientific purposes (2010/63/EU). The study was reviewed by the ethics commission (S15 TierSchG) and approved by the state authority (Regierungspräsidium, Tübingen, Baden-Württemberg, Germany). Analysis of the multimodal data included temporal correlation between the pupil dynamics and the fMRI signals, spatial correlation between each fMRI volume and the pupil–fMRI correlation map, and cross-correlation measures between the whole-brain fMRI, the calcium signal at 2 to 3 Hz or baseline, and the pupil size/pupil dilations. A scheme summarizing the main analysis pipeline can be found in SI Appendix, Fig. S3. For details concerning surgical procedures, calcium signal acquisition, fMRI sequence, and pupil measures, see SI Appendix.

ACKNOWLEDGMENTS. This work emerged as part of a PhD thesis supported by the Max Planck Society and the Graduate Training Centre of Neuroscience in Tuebingen (P. Pais-Roldán. “Multimodal assessment of coma and delirium: fMRI-based methods to study consciousness in the rat,” 2019, http://dx.doi.org/10.15496/publikation-34456). The research benefited from funding from the NIH Brain Initiative grants (RFA/NS11327B-01 and R01 MH111438-01), an S10 instrument grant (S10 RR020099-01) to the Massachusetts General Hospital/Harvard-MIT Program in Health Sciences and Technology Martinos Center, Deutsche Forschungsgemeinschaft (DFG, Germany Research Foundation) grant (YU 2153/1-3), and Bundesministe-rium fuer bildung und forschung (BMBF, Federal Ministry of Education and Research) grant (01GQ1702). We thank Dr. N. Avdievitch and Ms. H. Schulz for technical support, Mr. H. Huang for initial guidance on computer-based pupil-detection algorithms, Dr. E. Weiler, Ms. M. Pitscheider, and Ms. S. Fischer for animal protocol and maintenance support, and the teams of Mr. J. Boldt and Mr. O. Holder for mechanical and electrical support.


Pais-Roldán et al.


36. C. E. Byrum et al, A5 cells are silenced when REM sleep-like signs are elicited by pontine stimulation. J. Neurosci. 34, 7608-7618 (2014).


