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Astrocytes: integrators of arousal state and sensory context

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The integration of external information with the internal state of the body is central to the survival of virtually every multicellular organism. However, a complete picture of the mechanisms that govern this process is lacking. In this opinion article, we synthesize evidence demonstrating that astrocytes sense the momentary arousal state – through neuromodulator release – as well as the sensory inputs – through local synaptic activity – and respond to them with changes in calcium (Ca2+) signaling. We hypothesize that astrocytes integrate sensory signals with the internal state and that this process is necessary to secure optimal behavior. Finally, we argue that dysfunctional astrocytic Ca2+ signaling could be an underlying factor in disorders characterized by disrupted sensory processing.

How are external signals and internal state information merged in the brain?
The brain is constantly integrating signals conveying information about the inner states of the body with external sensory inputs from the outside world to produce appropriate, context-dependent behaviors [1–4]. This process ensures both efficient behavioral shifts when conditions change and the selection of relevant information for memory consolidation to guide future behavior.

The current model for this integration process suggests that sensory information is encoded by fast and spatially constricted neuronal synaptic transmission, while internal state information is relayed brain-wide by slowly acting neuromodulators. In this opinion article, we argue that astrocytes, an often-overlooked type of brain cell in systems neuroscience, also contribute to the integration of these two information sources, adding a new dimension to our understanding of how external and internal information is merged in the brain.

We first provide a short overview of the nature and characteristics of astrocytic Ca2+ signals. Second, we present evidence, mainly from rodent models, demonstrating that astrocytes respond to both arousal and sensory inputs with changes in their Ca2+ signaling. Next, we formulate a general hypothesis about the integration of arousal and sensory information in astrocytes. Finally, we discuss this hypothesis in light of recent data and outline outstanding questions and future experiments that can extend our knowledge about the role of astrocytes in cognitive functions.

Astrocytic Ca2+ signals
Astrocytic Ca2+ signaling has been characterized in a wide range of species and preparations, including human tissue [5], ferrets [6], rodents [7–15], zebrafish [16], and flies [17,18]. Surges in intracellular Ca2+ concentration are predominantly caused by Ca2+ release from internal stores, triggered by G protein-coupled receptor-mediated activation of the inositol trisphosphate (IP3) pathway. However, other IP3-independent mechanisms, such as Ca2+-permeable ion channels, exchangers, and transporters, are likely to be also involved (reviewed in [19,20]). The spatiotemporal

Highlights
A growing body of literature suggests that astrocytic calcium (Ca2+) signaling regulates various aspects of neuronal activity and behavior.

We propose that astrocytes are in a unique position to integrate internal and external signals via astrocytic Ca2+ signaling.

Astrocytes respond to internal arousal signals by detecting neuromodulator release, such as noradrenaline and acetylcholine, and elevating their intracellular Ca2+ concentration.

Astrocytes also respond to external sensory inputs by sensing local synaptic activity that leads to Ca2+ transients.

Manipulation of astrocytic Ca2+ signaling results in altered acquisition of sensory information.

We hypothesize that arousal can amplify weak sensory input at the level of astrocytes.

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The organization of 
Ca\(^{2+}\) signals in astrocytes is diverse (reviewed in [20–22]). Spatially, events range from local responses in fine astrocytic processes to responses extending throughout the cell, with the latter often occurring in many astrocytes in parallel [8–10,23] or spreading from one astrocyte to another [12,24]. Similarly, the time course of responses varies notably, from hundreds of milliseconds to minutes. Unfortunately, understanding of the underlying mechanisms and functional role of this diversity remains limited. A growing body of literature suggests that astrocytic 
Ca\(^{2+}\) signaling regulates neuronal activity and behavior, and the phylogenetic conservation of 
Ca\(^{2+}\) signaling in astrocytes suggests that this signaling has important functions. However, the question of when and why astrocytic 
Ca\(^{2+}\) signals are evoked remains to be answered. Interestingly, astrocytes express receptors for both neuromodulators and neurotransmitters [25], indicating that they are able to sense changes in arousal state as well as sensory input.

**Astrocytes signal arousal state**

The ability of animals to interact with external events highly depends on their body’s internal state, including their momentary arousal level [1–4] (Box 1).

In mammals, periods of heightened arousal correlate with increased noradrenaline (NA) and acetylcholine (ACh) signaling [26–28]. Recent in vivo experiments in awake mice have reignited research into NA-mediated astrocytic 
Ca\(^{2+}\) signaling [8,9] and there is strong evidence now supporting the notion that astrocytes reflect arousal changes by NA-driven global 
Ca\(^{2+}\) signaling [19–22,29], confirming early ex vivo data [30–32] and further evidenced by the cellular expression pattern of noradrenergic receptors [33,34]. Electrical or optogenetic stimulation of NA-producing locus coeruleus neurons leads to a sharp increase in astrocytic 
Ca\(^{2+}\) in the somatosensory cortex of awake and anesthetized mice [23,35].

Astrocytic 
Ca\(^{2+}\) signaling has been investigated in both externally triggered (e.g., air-puff whisker stimulation, tail stimulation, forced locomotion) and internally generated (e.g., spontaneous locomotion bouts) arousal-increasing contexts. Concretely, salient sensory stimulation [7–9,36–38] and self-initiated locomotion bouts [10,11,39,40] elicit NA-mediated 
Ca\(^{2+}\) signaling in large cortical astrocytic networks through activation of the alpha\(_{1}\) noradrenergic receptor. Although most studies on NA and astrocytic signaling have been done in rodents, there is evidence from zebrafish and *Drosophila* to support that this type of signaling is evolutionarily conserved [16,18].

The close association between arousal and astrocytic signaling is corroborated by experiments showing that astrocytic 
Ca\(^{2+}\) signaling is notably reduced during natural sleep and anesthesia compared with wakefulness [41–45]. Interestingly, large synchronized astrocytic 
Ca\(^{2+}\) events extending across the soma and processes are consistently associated with awakening from sleep [41–43], and these 
Ca\(^{2+}\) events are most likely to be induced by NA release [46]. While most

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**Box 1. The internal state of arousal**

In psychology and related fields, arousal is defined as a state of physiological activation associated with sensory stimulation or a state of excitement linked to an emotion. While arousal is sometimes conceptualized as a distinct state, in reality arousal comprises a spectrum ranging from states of lower to higher arousal. Arousal level is indexed by biological measures such as pupil diameter, heart and breathing rate, and shifts in brain activity patterns. Essential regulators of arousal in the nervous system are neuromodulators such as NA and ACh, which are increased in the aroused brain [1–4]. An inherent challenge when evaluating sensory input-related astrocytic 
Ca\(^{2+}\) responses in awake behaving animals is, therefore, the potential influence of concomitant arousal. In this opinion article, as a simplification, we differentiate between non-arousing sensory stimulation, such as the display of visual gratings and subtle whisker stimulation, and arousal-inducing salient sensory stimulation, such as startle-related air-puff stimulation and forced locomotion. In future experiments, monitoring arousal fluctuations during sensory experiences is essential and will allow researchers to disentangle the contributions of arousal and sensation and provide important new insight into the interaction between the two.
research on astrocytes and arousal has focused on NA, another important regulator of arousal, ACh, can also elicit astrocytic Ca\(^{2+}\) signaling [47,48]. Collectively, the literature strongly supports the idea that astrocytes respond to state-related neuromodulators and that astrocytic Ca\(^{2+}\) signaling is a general correlate of arousal.

**Astrocytes respond to sensory input**

Mounting evidence indicates that astrocytes can respond to sensory inputs *in vivo* [6,9,12–15,24,39,48–51] and local synaptic transmission *ex vivo* [13–15,20,50,52–55]. Curiously, astrocytic responses to sensory stimulation *in vivo* appear to depend on the arousal state of animals. For instance, astrocytes in the visual cortex of mice exhibit robust visual evoked Ca\(^{2+}\) signals during locomotion (i.e., increased arousal) [9,37,39]. Interestingly, these Ca\(^{2+}\) signals are larger than signals triggered by arousal alone, suggesting an interaction between sensory inputs and arousal in astrocytes [9,34]. By contrast, during periods of quiescence, visual evoked Ca\(^{2+}\) is weak or nonexistent [9,37,39]. These findings suggest that high arousal is necessary for sensory activated astrocytes, but sensory inputs still trigger astrocytic Ca\(^{2+}\) signals during anesthesia, a state characterized by low arousal [6,12–14,24,45–47]. Furthermore, after pharmacological blocking of arousal-related astrocytic Ca\(^{2+}\) signals in awake mice, visual input-driven Ca\(^{2+}\) transients in the soma [51] and whisker stimulation-induced Ca\(^{2+}\) events in the processes [15] are still observed. This aligns with the idea that local sensory evoked synaptic activity is adequate to drive astrocytic Ca\(^{2+}\) signaling independent of the arousal state. Together, these results suggest that astrocytic signaling reflects both sensory signals and arousal levels.

**Astrocytes may integrate arousal state and sensory information**

We hypothesize that astrocytes integrate external sensory signals (carried as synaptic activity in neuronal circuits) and internal arousal state information (relayed by neuromodulators such as NA and ACh) through differential Ca\(^{2+}\) signaling (Figure 1). We postulate that arousal-related signaling shifts astrocytes into a distinct activity mode, which increases the probability of them exhibiting Ca\(^{2+}\) signals in response to sensory inputs. Hence, if the sensory input-driven synaptic activity is insufficient to elicit measurable Ca\(^{2+}\) signaling alone, neuromodulators such as NA and ACh can push astrocytes above a theorized activation threshold. Conversely, if sensory input is strong enough, it can trigger astrocytic Ca\(^{2+}\) signaling without notable arousal. Intriguingly, our proposal is in congruence with recent preliminary work showing that the probability that astrocytic Ca\(^{2+}\) signals originating in the processes will propagate to the soma increases during states of higher arousal [56], suggesting that arousal primes astrocytes to be more responsive to sensory input.

While the precise mechanism of the proposed integration is currently unresolved, it is tempting to speculate that global arousal signaling is primarily mediated by Ca\(^{2+}\) release from internal stores through the IP\(_3\) pathway and local sensory signaling is initially mediated by IP\(_3\)-independent Ca\(^{2+}\) transients, such as via the activation of Ca\(^{2+}\)-permeable ion channels. It is plausible that these two different signaling pathways converge; arousal-related intracellular Ca\(^{2+}\) released from internal stores can increase the conductance of Ca\(^{2+}\)-permeable ion channels and thereby amplify sensory related signaling, while local Ca\(^{2+}\) transients mediated by ion channels can stimulate Ca\(^{2+}\) release from internal stores through Ca\(^{2+}\)-induced Ca\(^{2+}\) release [57]. Although this mechanistic explanation is likely to be overly simplified, it captures the core elements of our proposed hypothesis and thus seems a reasonable starting point for the guidance of future experiments aiming to decipher the subcellular mechanisms of the integrated astrocytic Ca\(^{2+}\) signal.

Finally, we propose that, downstream of the integrated Ca\(^{2+}\) signal, astrocytes can feed back to neural circuits to modulate sensory processing and, ultimately, behavior. Astrocyte-to-neuron signaling remains largely enigmatic, but we propose that it can occur as a result of direct
gliotransmitter release (e.g., ATP [53]) and/or through indirect regulation of glutamate and GABA uptake [58,59] or extracellular potassium ion levels [60]. Through these mechanisms, astrocytes can both enhance and dampen synaptic activity, and future experiments are needed to determine whether and when this occurs (see Outstanding questions).

Predictions and evidence for the behavioral role of astrocytic Ca\(^{2+}\) signaling

The most direct support for our hypothesis that astrocyte Ca\(^{2+}\) serves as a molecular substrate for integration of arousal and sensory information comes from studies manipulating astrocytic Ca\(^{2+}\) signaling in behaving animals. Here, we interpret the results of such studies through the lens of our hypothesis by discussing how they align with specific predictions.

Prediction 1

Attenuation of astrocytic Ca\(^{2+}\) signaling will result in impaired integration of information conveyed by sensory evoked neuronal and arousal states (i.e., necessity).

A novel tool that allows astrocytic Ca\(^{2+}\) signaling manipulation is iβARK, an inhibitory peptide that attenuates G\(_{q}\) G protein-coupled receptor-dependent Ca\(^{2+}\) elevations [36]. Interestingly, mice with brain-wide expression of iβARK in astrocytes exhibit deficient sensory adaptation to repeated obnoxious stimuli [36]. In this experimental context, the sensory input remains unchanged while the arousal level of the animals is likely to diminish as they habituate to repeated stimulation [61,62]. Thus, the deficient behavioral adaptation of iβARK-expressing mice suggests that astrocytic Ca\(^{2+}\) signaling secures a behavioral response that matches the subjects’ arousal state with incoming sensory information. Furthermore, attenuation of Ca\(^{2+}\) signaling led to impaired memory performance in tasks that relied on consolidating sensory information [36]. Specifically, this deficit was obvious in a subtle object exploration task (i.e., object location recognition), while performance in a task with salient novelty (i.e., object recognition) remained unaffected. Similarly, although using different tests, memory processes are affected in another mouse model with reduced Ca\(^{2+}\) signals in astrocytes: namely, mice lacking the IP\(_{3}\) type 2 receptor (IP\(_{3}\)R\(_{2}\) KO)
These mice exhibit subtle memory impairments for newly acquired sensory information and deficient long-term plasticity in the hippocampus and somatosensory cortex [64–69].

Notably, the findings from βARK and IP3R2 KO manipulations are in congruence with results from mice expressing an artificial Ca2+ pump that dampens Ca2+ signaling in both the soma and processes of astrocytes [59,70,71]. Such manipulation in the striatum leads to reduced neuronal excitability and excessive self-grooming [59]. Interestingly, NA-deficient mice also exhibit increased self-grooming [72]. A parsimonious interpretation of these results is that astrocytic Ca2+ signaling in response to arousal, signaled by NA release, and the accumulation of sensory information from the periphery (e.g., the orofacial area) is important for the termination of self-grooming [73]. Similar evidence comes from zebrafish, where astrocytic Ca2+ signaling increases after repeated bouts of futile swimming to control when an animal ultimately gives up [16]. These studies suggest that normal astrocytic Ca2+ signaling is necessary to ensure context-dependent sensory guided behaviors.

It is important to note that none of the currently available astrocytic Ca2+ loss-of-function tools permits specific attenuation of either arousal-related or synaptic activity-induced astrocytic Ca2+ transients. Therefore, there are currently no straightforward means to dissociate the two sources of Ca2+ signaling in astrocytes and establish a causal description of their roles.

Prediction 2
Artificial induction of astrocytic Ca2+ signaling that resembles naturally occurring dynamics will facilitate sensory information accumulation, leading to improved behavioral performance (i.e., sufficiency).

Studies have shown that artificial enhancement of astrocytic Ca2+ signaling during sensory guided learning promotes memory formation and retention [74–76]. These findings support the prediction that elevation of Ca2+ signaling will prime astrocytes to sense local synaptic activity and enhance sensory information accumulation, causing improved behavioral performance. However, an important limitation of these studies is the inherently artificial nature of the evoked Ca2+ signal. The highly complex spatiotemporal dynamics of astrocytic Ca2+ signals cannot currently be replicated with optogenetic or chemogenetic manipulations [77]. Finally, while currently available data appear to confirm specific predictions derived from our hypothesis, the multifaceted role of astrocytic Ca2+ signaling is undoubtedly more complex than outlined here, and many open questions remain to be addressed (see Outstanding questions).

Astrocytic Ca2+ signaling and pathophysiological conditions
Abnormal astrocytic Ca2+ signaling appears to be a pathophysiological component of various conditions, including disorders associated with disrupted sensory processing and arousal [78,79]. Sensory hypersensitivity and hyperarousal are hallmarks of autism spectrum disorders and fragile X and Rett syndromes [80–82]. Conversely, sensory hyposensitivity and hypovigilance are associated with major depression and dementia [83–86]. Interestingly, astrocytes derived from individuals with autism spectrum disorder or Rett or fragile X syndrome exhibit exacerbated Ca2+ signaling ex vivo [87–89], and mice deficient in astrocytic Ca2+ signaling exhibit long-range functional connectivity changes in vivo consistent with those seen in patients with major depressive disorder [90]. Moreover, recent reports suggest that astrocytic Ca2+ signaling is less responsive to NA and sensory inputs in two different models of Alzheimer’s disease [91,92]. Interestingly, some antidepressant medications, such as ketamine, could act through astrocytes [93]. It is therefore tempting to speculate that distorted astrocytic Ca2+ signaling is a key component of the cellular pathophysiology underlying sensory and arousal-related deficits in various brain disorders, and astrocytes could be a valid treatment target.
Concluding remarks and future perspectives
Considering astrocytes as integral players in cognitive function adds a new layer to the mechanistic understanding of information processing in the brain. While neurons can deliver fast and precise information, and neuromodulators provide long-lasting and long-ranging signals to tune brain activity, astrocytic Ca^{2+} signaling is somewhere in the middle: slow yet sufficiently fast to regulate behavior; diffused yet defined; and gradual yet punctual. It is these properties that enable astrocytes to participate in the integration of external sensory information carried as synaptic activity in sensory neuronal circuits, with momentary internal arousal state relayed by neuromodulators. According to our hypothesis, astrocytic Ca^{2+} signaling is central in fine-tuning the dynamic range of sensory processing, permitting the integration of subtle sensory inputs during aroused states while also responding to strong sensory inputs in the absence of notable arousal. This optimum sensory dynamic range facilitates appropriate behavioral responses and drives memory formation.

Astrocytes are increasingly appreciated as a vastly heterogeneous cell-type population between and within brain areas [94–97]. Astrocytic Ca^{2+} signaling is likely to depend on the specific astrocyte subtype and is adapted to respond to local neuronal activity patterns and neuromodulation. While most studies in this context have focused on astrocytic Ca^{2+} signaling in the neocortex, technological progress now allows the investigation of astrocytic activity in subcortical areas such as the hippocampus [98,99] and the striatum [69]. Furthermore, microglia, the resident immune cells of the CNS, interact with astrocytes and neurons [100] and can regulate neuronal activity in an arousal-dependent manner [101,102]. Future work will undoubtedly widen current knowledge of the diversity and function of state-dependent glial cell signaling.

There is growing recognition that tackling some of the fundamental questions that neuroscience is grappling with requires large datasets collected using advanced tools in a streamlined manner. In view of this perspective, there has been a move towards centralized brain observatories and large collaborations (e.g., Allen Brain Observatory, International Brain Laboratory, BRAIN Initiative) [103]. While glial cells are yet to be part of the focus of such large-scale initiatives, placing astrocytes and other glial cells in their remit would allow these cells’ roles in brain function and behavior to be uncovered. Crucially, well-defined hypotheses – for instance, the proposed roles of astrocytic Ca^{2+} dynamics in controlling behavior, as presented here – are essential to make the most out of the new age of neuroscience research.

Much remains to be learned about how astrocytic Ca^{2+} signaling is regulated by, and in turn influences, neuronal activity. However, two things are certain: (i) communication between neurons and astrocytes is vastly more complicated than previously thought; and (ii) continuing technological progress is allowing detailed dissection of this communication. It is our hope that future work, guided by specific hypotheses, will significantly expand current understanding of how astrocytes regulate internal state-dependent sensory processing.

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Declaration of interests
The authors declare no interests.
References

1. Raveli, S.W. et al. (2022) The emergence and influence of internal states. Neuron 110, 2545–2570
36. Nagy, J. et al. (2021) Specific and behaviorally consequential astrocyte G\alpha i GPCR signaling attenuation in vivo with 1APR. Cell 109, 2256–2274
42. Bojanowski, L. et al. (2020) Astrocytic Ca^2+ signaling is reduced during sleep and is involved in the regulation of slow wave sleep. Nat. Commun. 11, 3240
43. Tsunematsu, T. et al. (2021) Region-specific and state-dependent astrocyte Ca^2+ dynamics during the sleep-wake cycle in mice. J. Neurosci. 41, 5440–5452
44. Vadiyanathan, T. et al. (2021) Cortical astrocytes independently regulate sleep depth and duration via separate GPCR pathways. Elife 10, e63229
Trends in Neurosciences, June 2023, Vol. 46, No. 6


55. Panahian, A. et al. (2011) Astrocytes are endogenous regulators of basal transmission at central synapses. Cell 146, 785-798


63. Liu, J. et al. (2022) Distinct roles of astroglia and neurons in synaptic plasticity and memory. Mol. Psychiatry 27, 873-885


68. Lu, X. et al. (2022) OXO-astress induces extrasynaptic calcium signaling attenuation in vivo with CaEx Fluorescent dyes. J. Neurosci. 41, 4556-4574


70. Lustberg, D.J. et al. (2012) Norepinephrine and dopamine contribute to distinct repetitive behaviors induced by novel odorant stress in male and female mice. Horm. Behav. 64, 1050-1056


74. Iwai, Y. et al. (2021) Transient astrocytic GABA signaling underlies remote memory enhancement. Front. Neural Circuits 15, 658343


81. Schiller, C.E. et al. (2013) Remitted major depression is characterized by reduced prefrontal cortex reactivity to reward loss. J. Affect. Disord. 151, 756-762


85. Allen, M. et al. (2022) Astrocytes derived from ASD individuals alter behavior and destabilize neuronal activity through aberrant Ca2+ signaling. Mol. Psychiatry 27, 2470-2484


89. Lines, J. et al. (2022) Astrocyte-neuronal network interplay is disrupted in Alzheimer’s disease mice. Glia 70, 368-378

90. Åbjørsbråten, K.S. et al. (2022) Impaired astrocytic Ca2+ signaling in awake-behaving Alzheimer’s disease transgenic mice. Elife 11, e50555


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