Microglial TNF α controls GABA_AR plasticity, slow waves and memory consolidation during sleep

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ABSTRACT: Microglia sense the changes in their environment. How microglia actively translate these changes into suitable cues to adapt brain physiology is unknown. We reveal an activity-dependent regulation of cortical inhibitory synapses plasticity by microglia, driven by purinergic signaling acting on P2RX7 and mediated by microglia-derived TNF α . We demonstrate that sleep induces this microglia-dependent inhibitory plasticity by promoting synaptic enrichment of GABA_ARs. We further show that in turn, microglia-specific depletion of TNF α alters slow waves during NREM sleep and blunts sleep-dependent memory consolidation. Together, our results reveal that microglia orchestrate sleep-intrinsic plasticity of inhibitory synapses, ultimately sculpting sleep slow waves and memory.

Microglia, the immune cells of the brain, tune neuronal networks in the healthy brain by finely modulating synapses (Kettenmann et al., 2013). They can sculpt developing circuits and remodel neuronal connectivity in adulthood by adjusting synapse density, function and plasticity (Favuzzi et al., 2021; Nguyen et al., 2020; Parkhurst et al., 2013; Schafer et al., 2012; Wang et al., 2020). Microglia accomplish these functions either by direct interaction with synaptic elements (Miyamoto et al., 2016; Schafer et al., 2012) or through the release of factors (Lewitus et al., 2016; Parkhurst et al., 2013). Many of the latter, such as prostaglandin, BDNF, IL-1 β or TNF α , control synaptic plasticity and were independently shown to regulate sleep (Porkka-Heiskanen, 2013). Recent work highlights the ability of microglia to control sleep duration (Corsi et al., 2021; Liu et al., 2021); however, whether and how microglia-released factors shape sleep via modulation of synaptic plasticity remains unknown.

Synapse plasticity in the sleeping brain likely supports crucial functions of sleep. In the cortex, spine turnover during sleep is associated to learning and memory (Li et al., 2017; Yang et al., 2014) and, in parallel, wake-associated strengthening of excitatory synapses is downscaled during sleep (Vyazovskiy et al., 2008a; Diering et al., 2017). Synaptic inhibition is critically involved in sleep generation and sleep oscillations (Dixon et al., 2015; Funk et al., 2017; Niethard et al., 2018) but whether and how inhibitory synapses are dynamically modulated during sleep remains poorly understood. Here, we uncover the molecular pathway underlying sleep-intrinsic microglia-dependent modulation of synaptic GABA_AR as well as its impact on sleep slow waves and sleep-dependent memory consolidation.

Daily modulation of GABA_ARs in a sleep- and microglia-dependent manner

We first analyzed the modulation of synapses in the frontal cortex across the 24h light/dark cycle by measuring the synaptic content of neurotransmitter receptors (fig. 1 and fig. S1 and S2). Mice sleep more during the light phase and spend most of the dark phase awake. Therefore, we compared brains of mice at Zeitgeber time 18 (ZT18; dark/middle of wake phase) and at ZT6 (light/middle of sleep phase; fig. 1a). We focused on cortical layer 1 (L1) which is a key node for wide-scale cortical computation (Schuman et al., 2021). Consistent with the well-established downscaling of excitatory synapses during sleep (Diering et al., 2017), synaptic accumulation of the AMPA receptor subunit GluA2 was decreased at ZT6 as compared to ZT18 (fig. S1). Yet, analysis of L1 inhibitory synapses revealed an increased enrichment of synaptic, but not extra-synaptic, GABA_AR- γ 2 and - α 1 subunits at ZT6 (fig. 1a-c and fig. S2b-d), as well as an enhanced proportion of inhibitory synapses containing a GABA_AR cluster (fig. S2e). In contrast, in layer 5 (L5) the synaptic content of GABA_AR γ 2 did not significantly differ between ZT6 and ZT18 (fig. 1e and fig. S2f). These results reveal a novel form of regulation of GABAergic synapses across the light/dark cycle with synaptic enrichment of GABA_ARs in L1 during the light phase (ZT6), which likely contributes to upregulation of inhibitory transmission in the upper cortex during this phase (Bridi et al., 2020).

To discriminate between a sleep-intrinsic or time of day-dependent regulation of synaptic GABA_AR, mice were forced to stay awake during their normal sleep period (sleep deprivation from ZT0 to ZT6, SD6; fig. 1a). The synaptic content of GABA_AR was not different between mice at ZT18 and mice sleep-deprived in the light

phase (SD6; fig. 1d), showing that modulation of synaptic GABA_AR in L1 is driven by sleep-dependent mechanisms.

We have previously shown that microglia control the accumulation of receptors at inhibitory synapses in the spinal cord (Cantaut-Belarif et al., 2017). This prompted us to investigate the involvement of microglia in the modulation of synaptic GABA_AR over the sleep/wake cycle. Strikingly, microglia depletion by feeding mice with the CSF1R antagonist PLX3397 (PLX) (Elmore et al., 2014) (fig. S2a, b) completely prevented the changes in synaptic GABA_AR content between ZT6 and ZT18 (fig. 1c and fig. S2b,e). This shows that the sleep-dependent modulation of synaptic GABA_AR in L1 requires microglia.

Modulation of synaptic GABA_AR by microglial P2RX7 -TNF α signaling via CaMKII

We next identified the molecular actors of this novel microglia-dependent synaptic regulation (fig. 2), first in an *ex-vivo* system before validating them *in vivo*. During sleep, excitatory synaptic plasticity is triggered by NMDAR-dependent dendritic calcium spikes in L1 (Li et al., 2017). We thus selected an NMDA-induced GABAergic plasticity protocol, known as inhibitory long-term potentiation (iLTP). This form of plasticity, known to drive potentiation of excitatory synapses (Lee et al., 1998), also leads to an upregulation of synaptic GABA_ARs (Petrini et al., 2014) in pyramidal neurons specifically at somatostatin interneurons inputs (SOM-IN) (Chiu et al., 2018), which are mainly located in L1 (Tremblay et al., 2016) (fig. S3). In agreement with the SOM-IN topographical organization, NMDA-induced iLTP in brain organotypic slices led to specific synaptic enrichment of GABA_ARs in L1 but not in L5 somatic synapses (fig. 2a, b and fig. S4b-d). This protocol thus mimics *ex-vivo* the synaptic regulations occurring during the sleep/wake cycle so we used it to identify the actors of the L1-restricted sleep-related synaptic enrichment of GABA_ARs (fig. 1). We further demonstrated that this form of GABA_AR plasticity was completely abolished when microglia were depleted by PLX (fig. 2a, b and fig. S4a, c). We ruled out possible secondary effects of PLX by showing the same effect upon microglia depletion using Mac1-Saporin (SAP) (fig. S4a) or inactivation using minocycline (fig. 2b).

Microglia produce a broad repertoire of signaling molecules that regulate synaptic function (Kettenmann et al., 2013). TNF α , which is mostly if not exclusively produced by microglia in the brain (Zeisel et al., 2018), controls basal synaptic strength (Santello et al., 2011), neurotransmitter receptors' dynamics and homeostatic synaptic plasticity (Stellwagen and Malenka, 2006). We thus hypothesized that microglial TNF α controls the activity-dependent regulation of synaptic GABA_AR. Indeed, neutralization of TNF α by specific antibodies (fig. 2c) as well as conditional microglia-specific TNF α depletion (fig. 2d and fig. S5a) prevented the enrichment of synaptic GABA_AR upon iLTP. Soluble TNF α derives from the cleavage of a membrane form of TNF α by the TNF α -converting enzyme (TACE) (Black et al., 1997). Both forms can signal through TNFR1 whereas TNFR2 is only activated by membrane TNF α (Holbrook et al., 2019). The increase of synaptic GABA_ARs on iLTP-treated slices was prevented by TAPI-1, a TACE inhibitor, and by a TNFR1- but not by a TNFR2-neutralizing antibody (fig. S5b). Finally, we showed that recombinant TNF α and activation of TNFR1 were sufficient to increase synaptic accumulation of GABA_AR α 1 (fig. S5c-f). These experiments show that microglial soluble TNF α acting through TNFR1 mediates the activity-dependent regulation of GABA_AR in L1.

We next identified the signaling pathway between neuron and microglia leading to TNF α release upstream of GABA_ARs regulation. We first demonstrated that the well-established neuronal CX₃CL1-microglial CX₃CR1 signaling, with known roles in synaptic plasticity (Rogers et al., 2011), is not involved (fig. S4e). Microglia behavior is finely tuned by ATP via an array of purinergic receptors (Madry and Attwell, 2015). Microglial cells rapidly react to ATP following the activation of neurons by glutamate and NMDA (Badimon et al., 2020; Dissing-Olesen et al., 2014; Li et al., 2012), and the stimulation of microglial P2RX7 promotes the release of TNF α (Suzuki et al., 2004). We thus investigated whether NMDA-induced release of ATP causes microglia-mediated modulation of synaptic GABA_AR. Indeed, apyrase, a promoter of ATP hydrolysis, PPADS, a broad P2X antagonist, or A740003, a specific P2RX7 antagonist, prevented the increase of synaptic GABA_AR upon iLTP (fig. 2e). Moreover, BzATP, a P2RX7 agonist (Surprenant et al., 1996) was sufficient to increase postsynaptic GABA_AR α 1 in L1 (fig. 2f and fig. S6a,b) but not in L5 (fig. S6c). Of note, BzATP had no effect when microglia were depleted or when TNF α was specifically inactivated in microglia (fig. 2f). Thus, ATP/P2RX7 signaling acts upstream of microglial TNF α release to modulate synaptic accumulation of GABA_AR.

Finally, we explored the intracellular signaling downstream of microglial TNF α . CaMKII α is a central neuronal mediator of postsynaptic plasticity whose activity is triggered by Ca²⁺/calmodulin and can be prolonged in a Ca²⁺-independent manner by its autophosphorylation at Thr286 (Bayer and Schulman, 2019). Upon iLTP, Thr286-autophosphorylated CaMKII α leads to insertion of GABA_ARs at synapses (Marsden et al., 2010). We reasoned that TNF α may modulate CaMKII α Thr286-phosphorylation, as observed in non-neuronal cells (Defer et al., 2007). Indeed, Thr286-phosphorylation of CaMKII was increased in L1 upon iLTP (fig. 2g) and this increase was abolished by microglial depletion and by blocking P2RX7 or TNF α signaling (fig. 2h). In agreement, BzATP, which upregulates the synaptic content of GABA_ARs via a microglial relay, was sufficient to enhance CaMKII Thr286-phosphorylation (fig. 2g, h). The finding that CaMKII Thr286 phosphorylation, and therefore its activity, is gated by microglial TNF α downstream neuronal activity emphasizes a cardinal position of microglia in fine-tuning synapses.

Collectively, our results support a bidirectional neuron-microglia crosstalk underlying activity-driven GABAergic potentiation in L1 (fig. 2i): ATP released downstream neuronal activity activates microglial P2RX7 with concomitant release of TNF α which modulates CaMKII autophosphorylation and thereby enrichment of synaptic GABA_ARs.

Sleep-dependent modulation of synaptic GABA_AR driven by P2RX7 and microglial TNF α

Having shown that microglial P2RX7/TNF α pathway controls L1 activity-dependent GABA_AR plasticity via CaMKII autophosphorylation *ex-vivo*, we examined the involvement of this pathway in sleep-dependent modulation of synaptic GABA_AR across the light/dark cycle (fig. 1). We first showed that CaMKII Thr286-phosphorylation, but not total CaMKII levels, was increased in L1 at ZT6 (fig. 3a, b and fig. S7a). Notably, such regulation was not found in L5 (fig. S7b). Next, we showed that Thr286-phosphorylation of CaMKII, as well as the synaptic content of GABA_ARs in L1, were not enhanced at ZT6 in mice with microglia-

specific TNF α depletion (micTNF α -KO, fig. S5a) and in P2rx7-KO mice (fig. 3a-c and fig. S7c, d). This shows that P2RX7 and microglial TNF α drive daily fluctuations in CaMKII Thr286-phosphorylation and are required for sleep-dependent GABA_AR synaptic upregulation in L1 during the light phase.

NREM slow waves shaped by microglial TNF α

The results above show that microglial TNF α is required for a novel sleep-dependent regulation of inhibitory synapses in cortical L1. Sleep is an alternation of REM and NREM sleep periods which are hallmarked by major EEG oscillatory events (Adamantidis et al., 2019). During NREM sleep, slow waves in the delta frequency band (0.1-4Hz), quantified as slow wave activity (SWA), result from the synchronous alternation of active (up) and silent (down) states of cortical neurons. Cortical GABAergic inhibition is a major actor of NREM sleep slow waves (Hay et al., 2021; Lemieux et al., 2015). In particular, the activation of somatostatin positive interneurons, that mainly target L1, triggers down-states (Funk et al., 2017; Niethard et al., 2018). Because microglial TNF α modulates synaptic GABA_AR at L1 during sleep, we anticipated that it could also shape SWA. Sleep was analyzed using EMG and epidural EEG recordings, and we concentrated our analysis in the frontal cortex where slow waves are predominant (Vyazovskiy et al., 2006). We first showed that microglial TNF α has limited effects on sleep-wake patterns as shown by the lack of major alterations in the amounts of wake, NREM and REM sleep between micTNF α -KO and tCTL mice along a light/dark cycle (fig. 4a). Of note, the daily amount of REM sleep and number of REM bouts was increased on micTNF α -KO (table S1). This phenotype agrees with a suppressive effect of TNF α on REM sleep when injected into the basal forebrain (Terao et al., 1998). We thus anticipate that microglial TNF α may control REM by acting at the basal forebrain, which is out of the scope of this study and therefore not further explored.

We then analysed the spectral density of the EEG during sleep (figure 4b). In agreement with a role of microglial TNF α in the regulation of SWA, we found a shift in the EEG spectral density towards lower frequency activity during NREM sleep with a slower peak frequency in the delta range in micTNF α -KO as compared to tCTL ($p=0.02539$). There were no detectable differences in the other frequency ranges nor in REM sleep (fig. 4b). Finally, we explored how microglial TNF α shapes the properties of individual slow waves during NREM sleep. Slow waves were identified in the epidural EEG as alternation of large transient negative and positive deflections in the 0.1-4 Hz filtered EEG (fig. 4c). We verified that these deflections correspond to up and down states respectively (fig. S8; Nir et al. 2011). Remarkably, the maximum ascending slope of the slow waves, which coincides with the onset of the cortical positive deflection (downstate; fig. S8c), was decreased and the duration of the SW down state (fig. S8a, b) was increased in mice depleted of microglial TNF α as compared to controls (fig. 4d). These results indicate that microglial TNF α shapes slow waves during NREM sleep by favouring transition into the down states.

Microglial TNF α tunes sleep-dependent memory consolidation.

Slow waves are known to play a causal role for the consolidation of memory during sleep (Fattinger et al., 2017; Marshall et al., 2006). Our previous results show that microglial TNF α exquisitely shapes slow waves during NREM sleep, leading us to anticipate a crucial role of microglial TNF α in the memory consolidation

processes during sleep. To test this hypothesis, we compared the consolidation of a complex motor learning task known to be sleep-dependent (Nagai et al., 2017) in micTNF α -KO and tCTL mice. In this learning paradigm, mice learnt to run on top of a complex wheel attached to an accelerating rotarod in a first 20-trial session (S1). After one day of ad libitum sleep, performance on the complex wheel was assessed on a second 20-trial session by measuring the latency to fall (S2; fig. 5a, b).

We first confirmed that the learning, evaluated as the improvement of performance within each session, was not different between tCTL and micTNF α -KO (fig. 5c). This shows that lack of microglial TNF α did not alter complex motor learning, which is known to be sleep-independent (Nagai et al., 2017). We further showed that it does not impair locomotor activity and does not induce anxiety-like behavior as assessed in an open-field task (fig. S9). We next measured the improvement of performance in the complex motor learning task between S1 and S2 and found that it was higher in tCTL as compared with micTNF α -KO (fig. 5d). Finally, we tested the memory consolidation by comparing performance at the beginning of the second session (First S2) with either the last or the mean performance of the first session (Last S1 or Mean S1). In agreement with our initial hypothesis, memory consolidation was impaired in mice lacking microglial TNF α as compared to controls (fig. 5e). Remarkably, in the complex motor learning task, the improvement of performance between sessions and the memory consolidation are both known to be sleep-dependent processes (Nagai et al., 2017).

Discussion

Microglia, the principal immune cells of the brain, are now acknowledged as instrumental for the perception of the external environment (Thion et al., 2018). How microglia actively translate their external sensing into suitable cues to adapt circuitries in the healthy brain is yet unknown. Our results favour a model in which microglia sense neuronal activity through an ATP/P2RX7 signalling pathway and respond to it by releasing TNF α . Microglial TNF α then gates the phosphorylation of neuronal CaMKII that modulates GABA_AR content at cortical synapses (fig. 2). During sleep, this pathway is likely recruited by calcium spikes on apical dendrites of pyramidal neurons (Li et al., 2017), leading to strengthening of inhibitory synapses by upregulation of synaptic GABA_AR content (figs. 1 and 3). In line with a prominent role of inhibition in the generation of slow waves (Funk et al., 2017; Hay et al., 2021; Lemieux et al., 2015; Niethard et al., 2018; Zucca et al., 2017) microglial TNF α shapes slow waves by controlling transition into down states (fig. 4) and is thereby involved in sleep-dependent memory consolidation (fig. 5).

TNF α has long been known as a sleep factor. Administration of exogenous TNF α promotes SWA and NREM sleep (Fang et al., 1997; Yoshida et al., 2004) whereas inhibition of endogenous TNF α reduces NREM sleep (refs in Rockstrom et al., 2018). This is in apparent contradiction with our results showing no difference in NREM sleep amount in mice with microglia-specific TNF α depletion. Yet, to increase NREM sleep, TNF α was injected either directly in the brain (Yoshida et al., 2004) or intraperitoneally (Fang et al., 1997) at concentrations probably higher than the physiological concentration (Garré et al., 2017), with putative widespread diffusion and off-target and indirect effects. Finally, the studies using TNF α knock-out have not used cell specific and inducible inactivation and it is thus difficult to discriminate between microglial specific effect to indirect effects due non-microglial TNF α and/or secondary developmental alterations. Finally, TNF α is also a major mediator of inflammation, which induces sleep dysfunction by yet elusive mechanisms (Irwin, 2019). Our work now provides possible molecular and cellular links between sleep and inflammation.

Our results were obtained from experiments performed both *in vivo* and in organotypic slices. The latter system closely mimics the *in vivo* brain by preserving tissue architecture and cellular composition. In these slices microglia retain their 3D ramified morphology and their functional properties (De Simoni et al., 2008; Delbridge et al., 2020; Weinhard et al., 2018). They further conserve their ability to regulate synapses (Cantaut-Belarif et al., 2017; Pascual et al., 2012). This indicates that functional interactions between neurons and microglia are conserved in organotypic slices. Indeed, in this work, we have identified microglial TNF α , ATP/P2RX7 and CaMKII as molecular actors of synaptic GABA_ARs regulation in organotypic slices and we have further confirmed their role *in vivo* in the regulation of GABA_ARs plasticity during the sleep/wake cycle.

Excitatory synapses are scaled down during sleep, through removal of AMPA receptors, to compensate for potentiation due to ongoing learning during wake (Vyazovskiy et al., 2008a). Microglia contribute to downscaling during sleep by eliminating excitatory synapses (Choudhury et al., 2019), a behaviour presumably tuned down during wake by noradrenaline (Liu et al., 2019; Stowell et al., 2019). More recently, microglial CX3CR1 signalling was shown to differentially regulate excitatory synaptic transmission along the light/dark cycle (Corsi et al., 2021). We now show that microglia tune a sleep/wake regulation of inhibitory synapses

restricted to cortical layer 1, via mobilization of the P2RX7/TNF α signalling and regulation of synaptic GABA_AR content. Altogether, microglia are arising as active players in excitatory and inhibitory synapse remodelling along the sleep/wake cycle, likely to underlie daily oscillations in synaptic strength (Bridi et al., 2020; Tononi and Cirelli, 2019; Vyazovskiy et al., 2008b).

Cortical inhibition is involved in the control of slow wave activity during NREM sleep (Funk et al., 2017; Zucca et al., 2017; Zielinski et al., 2019), with a prominent role in the onset of down states (Chen et al., 2012; Lemieux et al., 2015). Such control is likely achieved through a combination of different inhibitory networks. Thalamic drive onto L1 inhibitory neurogliaform cells induces transition into down states (Hay et al., 2021). In addition, SOM-IN firing precedes entry into down states (Niethard et al., 2018) and accordingly their stimulation increases the slope of slow waves and triggers down-states (Funk et al., 2017). We now show that in agreement with their ability to modulate inhibitory synapses in the upper layers of the cortex, microglia are genuine regulators of slow waves during NREM sleep. Together with evidence of astrocytic involvement in slow wave activity (Szabó et al., 2017; Vaidyanathan et al., 2021), our work offers insight into how local glial modulation of neuronal networks tunes brain oscillations during sleep. One of the major physiological role of sleep is to allow memory consolidation (Klinzing et al., 2019), which critically depends on slow waves (Fattinger et al., 2017; Marshall et al., 2006). We now demonstrated that mice lacking TNF α display weaker consolidation than control mice, a process that putatively relies on the molecular microglia-neuron signaling herein described and downstream impact on neuronal dynamics during NREM sleep.

Finally, this work adds to the yet limited knowledge of microglia functions in the healthy adult brain (Parkhurst et al., 2013; Wang et al., 2020) and establishes microglia as genuine regulators of inhibitory synapses plasticity, brain oscillations and memory in the healthy brain. Noteworthy, microglial regulation likely occurs through the control of CaMKII α , which is a cardinal regulator of synaptic plasticity (Bayer and Schulman, 2019). It occurs on SOM interneurons inputs, which are involved in the control of complex behaviours such as sleep, but also decision making and learning (Adler et al., 2019). Finally, this work demonstrates that microglia tune slow waves and support memory consolidation during sleep (Klinzing et al., 2019). We anticipate a far wider involvement of microglia in other forms of plasticity and higher brain functions.

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Fig. 1 –Plasticity of GABA_AR in the light/dark cycle is sleep- and microglia-dependent. **a**, Experimental groups: mice at dark phase (ZT18), mice at light phase (ZT6) and mice submitted to sleep deprivation (dashed) in the light phase (SD6). Vertical bars: time of perfusion. **b**, Representative images showing enrichment of GABA_ARγ2 (cyan) at cortical L1 inhibitory synapses in the light phase (ZT6). Arrowheads: GABA_ARγ2 clusters at gephyrin⁺VGAT⁺ synapses in control (CTL) or PLX3397-treated mice (PLX). Scale bars, 5 and 1 μm. Dashed box corresponds to CTL at ZT18. **c**, **d**, Mean intensity of GABA_ARγ2 clusters at gephyrin⁺VGAT⁺ synapses (synaptic) and at extrasynaptic sites normalized to ZT18. n = 48 to 65 fields of view (FOVs) from 4–5 mice per group. *p < 0.05, nested t-test; **p < 0.01 compared with ZT18 and ##p < 0.01 compared with SD6, nested one-way ANOVA followed by Sidak's multiple comparison test. **e**, Left: Representative confocal images of VGAT and GABA_ARγ2 in cortical L5. Yellow line delineates soma identified by NeuN staining. Scale bar, 5 μm. Right: Mean intensity of GABA_ARγ2 at somatic VGAT⁺ clusters in L5 normalized to ZT18. n = 60 FOVs from 5 mice per group.

Fig. 2 – Microglial P2X7R-TNFα signaling drives GABA_ARs synaptic enrichment through CaMKIIα phosphorylation.

a, Representative confocal images showing increase of GABA_ARα1 (cyan) at gephyrin⁺ clusters (arrowheads) upon NMDA-induced inhibitory long-term potentiation (iLTP: 2 min 20 μM NMDA/10 μM CNQX plus 20 min recovery) in organotypic slices cortical L1 (CTL: control; bsl: baseline). **b-e**, Mean intensity of GABA_ARα1 clusters at gephyrin⁺ cluster normalized to CTL at bsl. n = FOVs/ independent experiments: (b) n = 44 to 69/ 5 to 6; (c) n = 47 to 53/ 5; (d) n = 66 to 102/ 5 to 7; (e) n = 49 to 68/ 7 to 9. *p<0.05, **p<0.01 and ***p<0.001, nested one-way ANOVA followed by Sidak's multiple comparison test. iLTP-induced synaptic GABA_AR enrichment is abolished by: **b**, microglia depletion/inactivation (PLX: PLX3397; SAP: Mac1-saporin; minoc: minocycline); **c**, neutralization of TNFα (nTNFα); **d**, microglia-specific TNFα deletion through 4-hydroxy-tamoxifen (4-OHT)-induced recombination on CX3CR1^{CreERT2/+}:TNF^{fl/fl} but not on a TNF^{fl/fl} background; **e**, ATP hydrolysis (apy), P2XR antagonist (PPADS) and P2X7R antagonist (A74). **f**, Left: Confocal images of GABA_ARα1 (cyan) at gephyrin⁺ clusters (arrowheads) in bsl and upon BzATP treatment. Right: Mean intensity of GABA_ARα1 clusters at gephyrin⁺ clusters normalized to CTL at bsl. n = 49 to 63/ 5 to 6. *p<0.05, **p<0.01, nested one-way ANOVA followed by Sidak's multiple comparison test. **g**, Left: Thr286-phosphorylated CaMKII is enhanced in L1 at the induction phase of plasticity (iLTP0' or BzATP0'). Scale bar, 5 μm. Right: Mean intensity of Thr286-phosphorylated CaMKII puncta normalized to bsl. n = 25 to 33 FOVs from 3 independent experiments. *p<0.05, nested t-test. **h**, Left: Western blot analysis showing iLTP0'-induced CaMKII Thr286-phosphorylation. Right: Ratio between Thr286-phosphorylated CaMKII and total CaMKII normalized to the respective iLTP-free control. n = 5 to 9 independent experiments. *p<0.05 compared with respective control, Kruskal-Wallis test followed by Dunn's multiple comparisons test. **i**, Model. ATP released downstream NMDA-induced neuronal activity activates microglial P2RX7, which triggers the release of microglial TNFα. TNFα signaling gates CaMKIIα autophosphorylation which controls the enrichment of synaptic GABA_ARs in pyramidal neurons.

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Fig. 4 - Microglial TNF α modulates slow waves during NREMS. **a**, Amounts of vigilance states over 24h reported by 2h segments. **b**, Average spectral density (top) of tCTL and micTNF α -KO (lines: means. Shaded area: SEM) and distribution of peak frequency of delta oscillations. The peak delta frequency is significantly lower in micTNF α -KO mice (Mann-Whitney $W = 157$, $p = 0.025$). **c**, Left: Examples of EEG and 0.1-4Hz filtered EEG during NREMS from a transgenic control tCTL; the positive and negative peaks of the delta-filtered signal are indicated by orange and blue points, respectively. Ticks on grey lines indicate large positive deflections corresponding to Slow Waves (SW). Right: average SW for tCTL showing duration (d) and maximum slope (s). **d**, The micTNF α -KO mice exhibit significantly shorter peak onset slope and longer SW duration than tCTL mice. Mann-Whitney $W=42$ $p=0.005$ and $W=153$, $p=0.37$ respectively

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