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Brief Communications

GABAergic Projections from the Medial Septum Selectively Inhibit Interneurons in the Medial Entorhinal Cortex

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The medial septum (MS) is required for theta rhythmic oscillations and grid cell firing in the medial entorhinal cortex (MEC). While GABAergic, glutamatergic, and cholinergic neurons project from the MS to the MEC, their synaptic targets are unknown. To investigate whether MS neurons innervate specific layers and cell types in the MEC, we expressed channelrhodopsin-2 in mouse MS neurons and used patch-clamp recording in brain slices to determine the response to light activation of identified cells in the MEC. Following activation of MS axons, we observed fast monosynaptic GABAergic IPSPs in the majority (>60%) of fast-spiking (FS) and low-threshold-spiking (LTS) interneurons in all layers of the MEC, but in only 1.5% of nonstellate principal cells (NSPCs) and in no stellate cells. We also observed fast glutamatergic responses to MS activation in a minority (<5%) of NSPCs, FS, and LTS interneurons. During stimulation of MS inputs at theta frequency (10 Hz), the amplitude of GABAergic IPSPs was maintained, and spike output from LTS and FS interneurons was entrained at low (25–60 Hz) and high (60–180 Hz) gamma frequencies, respectively. By demonstrating cell type-specific targeting of the GABAergic projection from the MS to the MEC, our results support the idea that the MS controls theta frequency activity in the MEC through coordination of inhibitory circuits.

Key words: gamma; interneuron; lamina organization; medial entorhinal cortex; medial septum; theta

Introduction

Spatial firing by neurons in the medial entorhinal cortex (MEC) is associated with theta frequency (4–11 Hz) oscillations in network activity (Mitchell and Ranck, 1980; Chrobak and Buzsáki, 1998; Colgin et al., 2009; Newman et al., 2013). Theta modulation of the MEC has lamina- and cell type-specific organization. For example, grid cells in layer II fire action potentials that prescess relative to the theta rhythm, whereas grid cells in layer III fire action potentials at a fixed phase relative to the theta rhythm (Hafting et al., 2008; Mizuseki et al., 2009). The phase and power of theta and associated nested gamma oscillations also vary between the lamina of the MEC (Chrobak and Buzsáki, 1998; Mizuseki et al., 2009; Quilichini et al., 2010). The medial septum (MS), which contains interacting populations of glutamatergic, GABAergic, and cholinergic projection neurons, is critical to theta activity and grid firing. Inactivation of the MS reversibly abolishes theta frequency oscillations and grid firing patterns (Mitchell et al., 1982; Brandon et al., 2011; Koenig et al., 2011). However, while anatomical studies establish direct projections from the MS to the MEC (Meibach and Siegel, 1977; Mitchell et al., 1982), we know very little about the synaptic targets of these projections or their cellular mechanisms of action.

To address this, we investigate synaptic responses of MEC neurons in brain slices following optical activation of axons from MS neurons expressing channelrhodopsin 2 (ChR2). We find that a high proportion of GABAergic interneurons across all layers of the MEC receive monosynaptic GABAergic inputs from the MS. Activation of GABAergic input from the MS at theta frequencies entrains low-frequency (25–60 Hz) and high-frequency (60–180 Hz) action potential firing by low-threshold-spiking (LTS) and fast-spiking (FS) interneurons respectively.

Materials and Methods

Surgical procedure. Experiments used C57BL/6J Ola Hsd (RRID:IMSR_JAX:000664) 8- to 10-week-old male mice, were approved by the University of Edinburgh animal welfare committee, and were performed under a UK Home Office project license. Animals were anesthetized with isoflurane and mounted in a stereotaxic frame (David Kopf Instruments). Adeno-associated virus (AAV) expressing either ChR2 conjugated with Venus fluorescent protein (pACAGW-ChR2-Venus-AAV serotype 2/1; generated by Vector Biolabs, from Plasmid 20071, AddGene) or ChR2 conjugated with mCherry [AAV-hSyn-CHR2(H134R)-mCherry serotype 2, University of North Carolina Vector Core, Chapel Hill, NC] was injected through a craniotomy 0.3 mm lateral to the midline and 0.6 mm caudal to bregma. Two injections of 400 nl were made at

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The Journal of Neuroscience, December 10, 2014 • 34(50):16739 – 16743 • 16739
Electrophysiological recordings. Three to 5 weeks after virus injection, horizontal brain slices were prepared, and whole-cell patch-clamp recordings were made from neurons in layers II–VI of the MEC, as described previously (Nolan et al., 2007). Slices were cut at ~4°C in solution with the following composition (in mM): NaCl 86, NaH2PO4 25, glucose 20, CaCl2 2, and MgCl2 1. For maintenance and recording the extracellular solution had the following composition (in mM): NaCl 124, NaH2PO4 1.2, KCl 2.5, NaHCO3 25, Glucose 20, CaCl2 0.5, and MgCl2 1. For maintenance and recording the extracellular solution had the following composition (in mM): K gluconate 130; KCl 10, HEPES 10, MgCl2 2, EGTA 0.1, Na2ATP 2, Na2GTP 0.3 Na phosphocreatine 10, biocytin 5.4. All recordings took place at 36 ± 2°C.

ChR2 was activated by 470 nm light, with irradiance of ~145 mW/mm², from an LED (Thorlabs) attached to the epifluorescence port of the microscope (BX-51; Olympus). Light stimuli had a duration of 10 ms and were repeated every 20 s for a minimum of five sweeps. Recordings used Multiclamp or AxoClamp amplifiers (Molecular Devices), and data were sampled at 20 kHz and digitized using a Digidata 1320A (Molecular Devices). Series resistance was ≤30 MΩ. Bridge balance and pipette capacitance neutralization were applied in current clamp, and series resistance compensation in voltage-clamp. An experimentally determined liquid junction potential of 12.7 mV was not corrected for. The reversal potential of IPSPs was estimated by fitting a polynomial function to plots of the mean IPSP amplitude as a function of baseline membrane potential (between ~80 and ~50 mV). Pharmacological agents were obtained from Abcam and were added to the standard extracellular solution at the following concentrations (in μM): NBQX 5 and D-APV 50 to block ionotropic glutamate receptors (iGluRs); picrotoxin 50 to block GABAA receptors (GABAARs); and atropine 1, mecamylamine 60, and methylacetylcholine 0.05 to block acetylcholine receptors (AChRs).

Cell identification. We used a hierarchical scheme to distinguish stellate cells (SCs), nonstellate principal cells (NSPCs), FS interneurons, and LTS interneurons. SCs were first identified by location in layer II, a sag response of <0.7 (SCs, 0.58 ± 0.01; NSPCs, 0.86 ± 0.01; LTS interneurons, 0.87 ± 0.02; FS interneurons, 0.87 ± 0.16), and input resistance of <90 MΩ (SCs, 38.4 ± 3.1 MΩ; NSPCs, 222.9 ± 10.6 MΩ; LTS interneurons, 276.0 ± 25.2 MΩ; FS interneurons, 94.9 ± 19.7 MΩ; cf. Garden et al., 2008; Pastoll et al., 2013). Putative interneurons were next distinguished from NSPCs by an afterhyperpolarization (AHP) of ≥10 mV (SCs, 5.8 ± 0.5 mV; NSPCs, 0.6 ± 0.4 mV; LTS interneurons, 13.3 ± 0.6 mV; FS interneurons, 21.6 ± 4.0 mV; Jones and Bühl, 1993; Sills et al., 2012; Couey et al., 2013). LTS interneurons were then identified by a rheobase of ≤120 pA (SCs 315.1 ± 32.1 pA; NSPCs, 107.1 ± 51.1 pA; LTS interneurons, 72 ± 6.5 pA; FS interneurons, 327.4 ± 60.6 pA).
remaining neurons were classified as FS interneurons. With these criteria neuronal firing properties during injection of current steps sufficient to trigger spiking for a duration of 3 s were similar to previous descriptions (e.g., median interspike interval: SCs, 108.0 ± 6.5 ms; NSPCs, 168.6 ± 8.89 ms; LTS interneurons, 61.7 ± 6.1 ms; FS interneurons, 19.7 ± 4.2 ms; p < 1 × 10⁻¹⁷, ANOVA; p < 0.005 for FS vs SC or NSPC, and LTS vs NSPC, Tukey’s test; mean action potential half-width (in ms): SCs, 0.60 ± 0.02; NSPCs, 0.78 ± 0.02; LTS interneurons, 0.52 ± 0.03; FS interneurons, 0.29 ± 0.06 ms; p < 1 × 10⁻²⁴, ANOVA; p < 0.005 for FS vs SC, NSPC, or LTS, and LTS vs NSPC, Tukey’s test; cf. Jones and Bühl, 1993; Pastoll et al., 2013).

To further evaluate our electrophysiological classification of interneurons, we examined filled cells for the presence of dendritic spines, which are typically found in principal cells, but not interneurons (Canto et al., 2008). After recording, a subset of slices was placed in 4% paraformaldehyde (PFA) overnight, washed three times with 0.01 M PBS and PBT 0.5%, washed in PBS, and mounted with Mowiol solution. Dendritic spines were present in electrophysiologically identified SCs (n = 2 of 2) and NSPCs (n = 4 of 4), and were absent in FS interneurons (n = 3 of 3) and LTS interneurons (n = 3 of 3).

Immunohistochemistry. Animals were transcardially perfused with 4% PFA in PBS. Sections were prepared and processed for immunohistochemistry as described previously (Nolan et al., 2007) and with the following primary antibodies: goat anti-chAT (1:200; RRID:AB_2079751, Millipore) for labeling cholinergic neurons; mouse anti-parvalbumin (PV; 1:2000; RRID:AB_10000343, Swant) and anti-GAD 67 (1:200; RRID: AB_2278725, Millipore) for labeling GABAergic neurons; and NeuroTrace 640/660 fluorescent Nissl stain (1:2000; RRID:nlx_152414, Life Technologies). The following secondary antibodies were used: donkey-key goat far red 647 (1:500; RRID:AB_141844, Invitrogen); goat anti-mouse Alexa Fluor-488 (1:800; RRID:AB_141167, Invitrogen). Imaging of labeled sections used a Nikon A1R confocal microscope with a 26.8 µm pinhole diameter and a 4.95 µm optical slice. MS mCherry or Venus-positive cells coexpressing ChAT, GAD-67, or PV were identified on a single focal plane and visually inspected for double labeling.

Data analysis and statistics. Electrophysiological data were analyzed using IGORpro (WaveMetrics), Clampfit (Molecular Devices), and Spike2 (CED). Rise and decay time constants were calculated by fitting a double exponential function to synaptic response waveforms. Normality of groups was assessed with the Shapiro–Wilk test. Comparisons between groups with Gaussian distributed data used t test or ANOVA followed by Tukey’s post hoc comparison. The Wilcoxon signed rank test, Kruskal–Wallis ANOVA, and Kolmogorov–Smirnov (KS) test were used for comparing nonparametric data. For quantification of fluorescence, the mean signal density in each layer was normalized to the peak signal density across all layers.

Results
To assess the identity of MS neurons projecting to the MEC, we injected the retrograde tracer CTB into the MEC (n = 4 mice). Within the MS, 41.7 ± 0.7% of cells labeled with CTB (72 of 168 cells) were positive for GAD67, the most common GAD isoform in the MS (Castañeda et al., 2005), and 6.6 ± 1.9% (13 of 204 cells) were positive for PV (data not shown), while 18.1 ± 3.7% of GAD67 (72 of 423 cells) and 6.9 ± 2.8% of PV cells (13 of 233 cells) were CTB positive. Thus, a large fraction of the projection from the MS to the MEC originates from GABAergic neurons, with a subset positive for PV. A further 13.2 ± 2.9% of CTB cells (26 of 204) were colabeled with ChAT, while 14.1% of ChAT cells (26 of 207 cells) were CTB positive, indicating that cholinergic neurons also project from the MS to the MEC.

To investigate the functional organization of this projection, we injected AAV expressing ChR2 conjugated with either mCherry or Venus into the MS (Fig. 1A). With our injection protocol, we found labeled cell bodies in 67.0 ± 5.2% of the area of the MS (n = 4 mice; Fig. 1B). Of neurons labeled with viral reporter, 55.0 ± 0.04% of cells (223 of 402 cells) were labeled with antibodies against GAD67, 9.5 ± 3.4% were positive for PV (24 of 284 cells), and a further 4.8 ± 1.4% were positive for ChAT (13 of 284 cells), while 61.0 ± 7.2% (223 of 357 cells), 33.1 ± 7.8% (24 of 71 cells), and 16.3 ± 6.0% (13 of 116 cells), respectively, of GAD67, PV, and ChAT cells were colabeled with viral reporter. Within the MEC, we observed a consistent laminar organization of labeled axons (Fig. 1C). We quantified this for horizontal MEC slices 2.6 mm from the dorsal surface of the brain. Here, fluorescence was significantly higher in layers II and V compared with upper layers I and VI (p = 0.0002, ANOVA; p < 0.05 for significant differences between layers, Tukey’s test; Fig. 1C). A similar organization was seen in more dorsal and ventral slices (data not
rons in layer II, received GABAergic inputs (least half of FS interneurons in layers III–VI, and all FS interneurons, LTS interneurons, SCs, or NSPCs (similar, whereas the response latency (1.7\(\tau\)) was abolished by picrotoxin (14.3 \(\pm\) 7.2 pA; \(p < 0.001\), Kruskal-Wallis ANOVA; Fig. 4A). Furthermore, the latency of GABAergic IPSPs was both short (3.3 \(\tau\)) and invariant (SD, 0.33 \(\pm\) 0.12 ms; range, 0.13–0.94 ms; Fig. 4C). Together, these data indicate that the MS sends monosynaptic inhibitory projections primarily to interneurons in the MEC. Glutamatergic responses are detected less frequently and do not appear to distinguish between NSPCs and interneurons, while SCs do not appear to respond to glutamatergic or GABAergic input from the MS.

We next sought to establish whether theta frequency activation of GABAergic input from the MS can affect spike output from neurons in the MEC. We first asked whether synaptic responses accurately follow stimulation at 10 Hz. We found that IPSPs on FS and LTS neurons were maintained during 2 s of stimulation (Fig. 4B–D). To determine whether IPSPs could entrain firing by interneurons, we examined responses during injection of constant current with amplitude just sufficient to sustain action potential firing (Fig. 4E–G). We observed that light stimulation induced pauses in spike firing followed by periods of rhythmic spiking, with little effect on the overall spike rate (control, 43.2 \(\pm\) 14.5 Hz; light stimulation, 47.7 \(\pm\) 15.9 Hz; \(p = 0.42\), paired \(t\) test). The duration of the pause in spiking was not significantly different for FS and LTS interneurons (14.4 \(\pm\) 2.5 and 21.7 \(\pm\) 10.0 ms; \(p = 0.17\), unpaired \(t\) test) and was relatively invariant (SD: FS interneurons, 4.3 \(\pm\) 1.3 ms; LTS interneurons, 2.6 \(\pm\) 1.1 ms; \(p = 0.11\), unpaired \(t\) test), while the interval between the first and second spikes was typically in the low gamma band for LTS (25–43 Hz) and in the high gamma band (50–163 Hz) for FS interneurons (34.8 \(\pm\) 5.3 and 118.2 \(\pm\) 26.5 ms respectively; \(p = 0.046\), unpaired \(t\) test), reflecting the intrinsic spike frequency of each cell type. These data demonstrate that MS GABAergic inputs to the MEC can follow 10 Hz stimulation and, in doing so, can entrain gamma frequency spike trains in MEC interneurons.

**Discussion**

We demonstrate that GABAergic projections from the MS synapse onto both FS and LTS interneurons in all layers of the MEC,
but rarely synapse onto excitatory cells. Projections from the MS generate inhibitory GABA<sub>a</sub> receptor-mediated postsynaptic responses that are maintained during stimulation at theta frequency and can entrain activity of responding cells. We observed glutamatergic responses to MS activation much less frequently and in a similar proportion of responding interneurons and NSPCs. We did not observe glutamatergic or GABAergic responses in layer II SCs. The specificity of connectivity between the MS and MEC imposes constraints on mechanisms for generation of theta oscillations and spatial firing by neurons in the MEC.

Our finding of preferential targeting of inhibitory inputs to interneurons in the MEC is consistent with anatomical findings in the hippocampus (Freund and Antal, 1988). Thus, while the hippocampus and MEC differ in their laminar organization, principles for inhibitory control of each structure by the MS may be similar. It is important to consider the degree to which our analysis might underestimate connectivity through incomplete infection of all projection neurons in the MS. This is unlikely to explain the absence of inhibitory responses of NSPCs and SCs, as NSPCs and SCs are intermingled with interneurons, and inhibitory responses were observed in the majority of interneurons in all layers. Thus, our approach clearly activates inhibitory axons from the MS. Our finding that glutamatergic projections from MS to MEC are relatively rare is consistent with the lower percentage of glutamatergic projections from MS to the hippocampus (Colom et al., 2005; Henderson et al., 2010). However, while our injections likely infect glutamatergic neurons in the MS, as they are found adjacent to GABAergic neurons (Colom et al., 2005), our data nevertheless do not rule out the possibility that glutamatergic projections are more difficult to activate. Our finding that glutamatergic and GABAergic synaptic responses are absent in SCs is consistent with low levels of connectivity found with rabies virus tracing of inputs to SCs (Rowland et al., 2013).

What implications does the specificity of targeting of MS GABAergic projections to the MEC have for generation of theta rhythms in entorhinal–hippocampal circuits? A majority of interneurons across all layers of the MEC and subregions of the hippocampus have similar preferred firing phases with respect to the theta rhythm (Mizuseki et al., 2009). Because GABAergic fibers appear to specifically target interneurons in the hippocampus (Freund and Antal, 1988) as well as the MEC, GABAergic projections from the MS may coordinate theta firing by interneurons throughout the entorhinal–hippocampal circuit. Consistent with this idea, firing of GABAergic neurons in the MS is phase locked to hippocampal theta and on average precedes firing of hippocampal interneurons (Hangya et al., 2009). A possibility suggested by our results is that this GABAergic input from the MS entrains rebound firing of entorhinal interneurons rather than simply switching off their activity. Given recent evidence that SCs communicate exclusively via inhibitory interneurons (Couey et al., 2013; Pastoll et al., 2015), control of inhibitory networks by GABAergic projections from the MS may be well suited to contribute to roles of septal GABAergic neurons in spatial behaviors (Dwyer et al., 2007; Pang et al., 2011).

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