Modulation of olfactory bulb network activity by serotonin: synchronous inhibition of mitral cells mediated by spatially localized GABAergic microcircuits

Loren J. Schmidt and Ben W. Strowbridge

Department of Neurosciences, Case Western Reserve University School of Medicine, Cleveland, Ohio 44106, USA

Although inhibition has often been proposed as a central mechanism for coordinating activity in the olfactory system, relatively little is known about how activation of different inhibitory local circuit pathways can generate coincident inhibition of principal cells. We used serotonin (5-HT) as a pharmacological tool to induce spiking in ensembles of mitral cells (MCs), a primary output neuron in the olfactory bulb, and recorded intracellularly from pairs of MCs to directly assay coincident inhibitory input. We find that 5-HT dysynaptically depolarized granule cells (GCs) only slightly but robustly increased the frequency of inhibitory postsynaptic inhibitory currents in MCs. Serotonin also triggered more coincident IPSCs in pairs of nearby MCs than expected by chance, including in MCs with truncated apical dendrites that lack glomerular synapses. That serotonin-triggered coincident inhibition in the absence of elevated GC somatic firing rates suggested that synchronized MC inhibition arose from glutamate receptor-mediated depolarization of GC dendrites or other (non-GC) interneurons outside the glomerular layer. Tetanic stimulation of GCL afferents to GCs triggered robust GC spiking, coincident inhibition in pairs of MCs, and recruited large-amplitude IPSCs in MCs. Enhancing neurotransmission through NMDARs by lowering the external Mg^{2+} concentration also increased inhibitory tone onto MCs but failed to promote synchronized inhibition. These results demonstrate that coincident MC inhibition can occur through multiple circuit pathways and suggests that the functional coordination between different GABAergic synapses in individual GCs can be dynamically regulated.

Results

Direct and indirect effects of serotonin in olfactory bulb neurons

Although coherent inhibitory inputs have been proposed to underlie spike synchrony in principal cells in the OB during sensory responses (Freeman 1972; Kashiyadani et al. 1999; Galán et al. 2006; Kay and Stopfer 2006), relatively few studies have
directly assayed the inhibitory local circuits responsible for synchronous inhibition. Typically, electrical stimulation methods are employed to either activate glomerular inputs (Aungst et al. 2003; Schoppa 2006a,b) or focally activate circuits within the plexiform layers (Nakashima et al. 1978; Isaacson and Strowbridge 1998; Friedman and Strowbridge 2003). Utilizing in vitro OB slices, we used a pharmacological approach to activate large populations of MCs to probe circuits responsible for inhibitory synchronization.

In rat horizontal OB brain slices, bath application of 200 μM 5-HT significantly increased the frequency of spontaneous MC spiking assayed in cell-attached recordings (from 3.0 ± 1 to 20.3 ± 4.7 Hz, P < 0.001, paired t-test, n = 8) (Fig. 1A–C). This depolarizing effect appeared to be mediated by postsynaptic 5-HT receptors on MCs as we found a similar increase in spontaneous spiking when 5-HT was tested after ionotropic glutamate and GABAA receptors were blocked using an antagonist cocktail containing APV, NBQX, and gabazine (from 4.3 ± 2.3 to 20.6 ± 5.9 Hz, P < 0.01, n = 11) (Fig. 1C). Although previous studies have identified effects of 5-HT on neurons within the glomerular layer (Hardy et al. 2005; Petzold et al. 2009), the depolarization of MCs we observed by 5-HT persisted in dissected OB slices that lack a glomerular layer (Fig. 1D; as in Friedman and Strowbridge 2003), suggesting a direct effect on post-synaptic receptors located on the MC somata or dendritic processes contained in the external plexiform layer. These results are consistent with Hardy et al. (2005) who showed intracellular depolarization following activation of 5-HT2Rs on MCs, though the location of the postsynaptic 5-HT receptors was not determined in that study. In similar experiments recording from tufted cells (TCs, fast synaptic transmission blocked), we found heterogeneous results with three of four cell-attached recordings from TCs remaining silent before and after application of 200 μM 5-HT. The one spontaneously active TC increased its firing rate in 5-HT from 4.1 to 13.1 Hz. One previous study (Liu et al. 2012) found direct depolarization of external TCs following activation of 5-HT2ARs. Given the possible heterogeneous effects of 5-HT on different subclasses of TCs, we focused the present study on 5-HT effects on MCs.

In cell-attached MC recordings, bath application of 200 μM 5-HT elicited a rapid increase in spontaneous firing that subsided with a time constant of 44 ± 9 sec (Fig. 1E). We also found rapid onset kinetics in intracellular recordings under whole-cell current-clamp conditions (from 69.2 ± 0.9 to –58.7 ± 2.2 mV, P < 0.001, n = 9, 200 μM 5-HT tested in APV, NBQX, and gabazine) (Fig. 1F). We took advantage of the rapid activation and desensitization kinetics following 5-HT applications in this study to determine the network consequences of brief (2–3 min) periods of induced spiking in populations of MCs.

We first asked if 5-HT also activated GABAergic GCs that form reciprocal dendrodendritic synapses with mitral and tufted cells (Fig. 2A,B; Rall et al. 1966; Isaacson and Strowbridge 1998). In 12 out of 14 GCs tested, 200 μM 5-HT depolarized the membrane potential. The mean membrane potential change following 5-HT application in GCs was 3.4 ± 0.7 mV, n = 14. Serotonin also increased the apparent frequency of spontaneous EPSPs (Fig. 2C) though many of these synaptic events were of small amplitude and therefore difficult to analyze. Blocking ionotropic glutamate and GABA_A receptors with APV, NBQX, and gabazine abolished most spontaneous synaptic activity in GCs and occluded the depolarizing effect of 5-HT (significantly different than the response to 5-HT without synaptic blockers, P < 0.002, unpaired t-test).

Figure 1. Serotonin directly depolarizes olfactory bulb mitral cells. (A) Cartoon of intracellular mitral cell (MC) recording configuration with synaptically connected granule cells (GCs). (B) Example cell-attached MC recording with fast synaptic transmission blocked (5 μM NBQX, 25 μM D-APV, 10 μM gabazine, GBZ). Bath application of serotonin (5-HT, 200 μM) increased the frequency of spontaneous MC action currents from 4.3 ± 2.3 to 20.6 ± 5.9 Hz. (C) Plot of increase in MC firing frequency following 5-HT application in control conditions (left, n = 8 experiments, [***] P < 0.001, paired t-test) and when fast synaptic transmission was blocked with NBQX + APV + GBZ (Blockers) prior to 5-HT application (right, n = 11 cell-attached recording experiments, [**] P < 0.001, paired t-test). (D) Plot of increase in MC firing following 5-HT application in dissected OB slices without glomerular layers. Same conditions as indicated in C. (†) P < 0.02, (‡) P < 0.05, paired t-test. (E) Plot of mean ± SEM of MC firing rate in 14 cell-attached recordings in NBQX + APV + GBZ. (F) Time course of MC depolarization following bath application of 5-HT (n = 9 MCs recorded in NBQX + APV + GBZ).
Indirect excitation of granule cells by serotonin. (A) Diagram of recording configuration. (B) Example granule cell (GC) response to a 50-pA depolarizing current step. (C) Example intracellular GC recording before and during 5-HT application. Horizontal line at −74 mV in both traces. (D) Blockade of fast synaptic responses with NBQX + APV + GBZ occludes depolarizing effect of 5-HT on GC. Horizontal line at −68 mV in all three traces. Mean membrane potential change in synaptic blocker recordings within 5 min of 5-HT application (0 out of 14 GCs tested). We confirmed the low incidence of 5-HT-induced spiking in GCs by performing 22 cell-attached recordings under the same conditions where 5-HT reliably induced MC spiking (as in Fig. 1B,C). The same small fraction of GCs (3/22) was spontaneously spiking before and after application of 200 μM 5-HT (Fig. 2E). Of the three out of 22 GCs that were spontaneously active before 5-HT, two increased their firing rates and one decreased firing following 5-HT treatment. We confirmed that we could detect spiking in cell-attached GC recordings by electrically stimulating in the granule cell layer following 5-HT application (Fig. 2E). Results from the GC cell-attached recording are summarized in Figure 2H and suggest that 5-HT-driven depolarization of OB principal cells synaptically excites GCs, resulting in a relatively small amplitude sustained somatic depolarization that remained subthreshold in nearly all GCs tested.

Inhibitory feedback to mitral cells evoked by serotonin

Despite only rarely triggering somatic spiking activity in GCs, 5-HT effectively increased the frequency of spontaneous inhibitory postsynaptic responses recorded in MCs (Fig. 3A1). The average frequency of spontaneous IPSCs recorded in MCs increased from 19.8 ± 1.9 to 47.0 ± 3.6 Hz following 3-min treatment with 200 μM 5-HT (137.4% increase, P < 0.0001, n = 61; IPSCs recorded as inward currents as a consequence of reversed Cl− gradient). The kinetics of 5-HT-evoked increase in inhibitory tone paralleled closely the time course of elevated MC spiking recorded in cell-attached recordings, including the rapid desensitization phase after ~3-min exposure to 200 μM 5-HT (Fig. 3A2). Blockage of GABA_A receptors with gabazine (GBZ, 10 μM) occluded the effect of 5-HT on spontaneous synaptic currents in MCs (n = 3 MCs) (Fig. 3A2).

Serotonin appeared to increase GABAergic inhibition of MCs via AMPA receptor-mediated depolarization of GCs. Blockade of NMDA receptors with APV (25 μM) lowered the basal IPSC frequency recorded in MCs (P < 0.05, n = 7 cells but failed occlude the transient increase in IPSC frequency following 220 ms occlude the depolarizing effect of 5-HT on GC. Horizontal line at −68 mV in all three traces. Mean membrane potential change in synaptic blocker recordings within 5 min of 5-HT application (0 out of 14 GCs tested). We confirmed the low incidence of 5-HT-induced spiking in GCs by performing 22 cell-attached recordings under the same conditions where 5-HT reliably induced MC spiking (as in Fig. 1B,C). The same small fraction of GCs (3/22) was spontaneously spiking before and after application of 200 μM 5-HT (Fig. 2E). Of the three out of 22 GCs that were spontaneously active before 5-HT, two increased their firing rates and one decreased firing following 5-HT treatment. We confirmed that we could detect spiking in cell-attached GC recordings by electrically stimulating in the granule cell layer following 5-HT application (Fig. 2E). Results from the GC cell-attached recording are summarized in Figure 2H and suggest that 5-HT-driven depolarization of OB principal cells synaptically excites GCs, resulting in a relatively small amplitude sustained somatic depolarization that remained subthreshold in nearly all GCs tested.
Serotonin and coincident inhibition

In addition to increasing the overall frequency of spontaneous IPSCs, 5-HT triggered synchronous inhibition in pairs of closely spaced MCs but only infrequently in more distant pairs of MCs (Fig. 4A). Synchronous IPSCs following 5-HT treatment did not result from activity in inhibitory local circuits in the glomerular layer because we observed similar synchronous inhibitory responses in a near pair of MCs in which both cells had apical dendrites truncated before the glomerular layer (Fig. 4B). We used the clipped cross-intensity function (CIF) (Hahnloser 2007) to quantify the frequency of synchronous IPSCs within a given onset latency window (2-msec duration) and the frequency of coincident events expected by chance (“Shuf”). In the paired recording between visualized MCs without apical dendritic tufts, 5-HT triggered a statistically significant increase in CIF over the CIF independent of previously described effects of 5-HT on periglomerular circuits (Petzold et al. 2009). Bath applications of 5-HT had no effect on the rate of miniature IPSCs recorded in 1 μM TTX, suggesting that the primary action of this transmitter was not related to mechanisms regulating GABA release from interneurons (Fig. 3E). The results summarized in Figure 3E–G are consistent with the hypothesis that the primary effect of 5-HT in these experiments is on principal neurons and that the increase in inhibitory tone following bolus applications of 5-HT arises from AMPAR-mediated excitation of GABAergic interneurons outside the glomerular layer.

Serotonin appeared to depolarize MCs by binding to postsynaptic 5-HT2 receptors. The 5-HT2R antagonist cinanserin (20 μM) (Peroutka and Snyder 1981) completely occluded the effect of bath application of 200 μM 5-HT (IPSC frequency 14.2 ± 4.5 Hz before 5-HT vs. 10.2 ± 3.1 Hz in 5-HT, P < 0.05, paired t-test, n = 5). By contrast, the 5-HT1R antagonist pindolol (25 μM) (Tricklebank et al. 1984) failed to block 5-HT-mediated increase in IPSC frequency (12.1 ± 3.1 Hz before 5-HT vs. 67.5 ± 14 Hz in 5-HT, P < 0.001, n = 5). Similarly, the 5-HT3/4R antagonist SDZ 205–557 (10 μM) (Buchheit et al. 1991; Eglen et al. 1993) did not prevent the 5-HT-mediated increase in MC IPSC frequency (9.4 ± 2.9 Hz before 5-HT vs. 31.7 ± 8.2 Hz in 5-HT, P < 0.02, n = 6). Also consistent with an action on 5-HT2Rs, we found the 5-HT2AR-selective agonist TCB2 (10 μM) (Fox et al. 2010) triggered a statistically significant increase in IPSC frequency (16.5 ± 2.8 Hz in control vs. 21.0 ± 3.0 Hz in TCB2, P < 0.02, n = 5).

**Synchronous inhibition onto subpopulations of mitral cells**

5-HT (Fig. 3C). Blockade of AMPARs with NBQX (5 μM) lowered the spontaneous IPSC frequency in MCs (P < 0.005, n = 7 cells) and also prevented 5-HT from increasing inhibitory tone (Fig. 3D). Serotonin triggered a similar increase in spontaneous inhibitory tone (160%) in dissected OB slices lacking glomerular layers (P < 0.0001) (Fig. 3E,F), suggesting that this response was similar to that seen in the presence of glutamate receptor antagonists. (***) P < 0.01, paired t-test, n.s., P > 0.05.

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**Figure 3.** Serotonin triggers disynaptic inhibition onto mitral cells. (A1) Example MC voltage–clamp recordings illustrating the increase in spontaneous IPSC frequency during 5-HT application (Cl– reversal potential reversed because of CsCl internal solution). (A2) Gabazine (GBZ, 10 μM) blocked spontaneous and 5-HT triggered inward currents. (B) Plot of the time course of the spontaneous IPSC frequency during 5-HT application. Each symbol represents mean ± SEM from 61 MCs. (C) Plot of the increase in MC IPSC frequency with 5-HT after NMDARs were blocked with 25 μM APV. Example traces in APV and APV + 5-HT shown in inset. (D) Plot of the effect of 5-HT on MC IPSC frequency after AMPARs were blocked with 5 μM NBQX. Example traces shown in inset. (E) Summary of the increase in MC IPSC frequency with bath application of 5-HT in 61 MCs recorded in OB brain slices in control conditions (left bars) and in 1 μM TTX (right bars). (***) P < 0.001, paired t-test, n.s., P > 0.05. (F) Plot of MC IPSC frequency increase with 5-HT in five experiments with dissected OB brain slices lacking glomerular layers. (***) P < 0.001, paired t-test. (G) Summary of the increase in MC IPSC frequency with 5-HT in the presence of glutamate receptor antagonists. (***) P < 0.01, paired t-test, n.s., P > 0.05.
expected by chance during 5-HT treatment ($P < 0.0002$) (Fig. 4C). The synchronous MC inhibition assayed by CIF only persisted for $\sim 1\mathrm{~min}$, paralleling the rapid desensitization of the 5-HT-evoked increase in IPSC frequency (see Fig. 3B). We also found a statistically significant increase in MC IPSC CIF in a different visualized pair of MCs in which each neuron contacted a different glomerulus (70 $\mu\text{m}$ between cell bodies, 0.22 IPSC CIF, $P < 0.001$, not shown). Over a set of 13 paired recordings between nearby MCs (cell bodies separated by $<200\ \mu\text{m}$, mean separation $= 83.2 \pm 13\ \mu\text{m}$), 5-HT reliably triggered more synchronous IPSCs than expected by chance (actual CIF was significantly larger than CIF computed following shuffling inter-IPSC intervals, $P < 0.005$) (Fig. 4D). We also observed a smaller, but still statistically significant, increase in IPSC synchrony in eight pairs of MCs separated by between 200 and 1000 $\mu\text{m}$ and no increase in IPSC synchrony above chance in five MC pairs separated by $>1000\ \mu\text{m}$ (Fig. 4D).

The lack of IPSC synchrony in distant pairs of MCs was not due to variation in effectiveness of 5-HT in driving inhibitory local circuits, since 5-HT elevated mean IPSC frequencies similarly in three paired recording configurations (Kruskal–Wallis ANOVA, $P > 0.05$) (Fig. 4E). When plotted directly against MC separation (Fig. 4F), 5-HT-driven “excess” IPSC synchrony (actual CIF—expected CIF estimated by shuffling inter-event intervals) decreased sharply with distance; most paired recordings separated by $<150\ \mu\text{m}$ had excess IPSC synchrony. By contrast, fewer paired recordings separated by $>150\ \mu\text{m}$ had more IPSC synchrony than expected by chance (near 0 excess CIF). These results suggest that despite the long length of MC dendritic processes (100s of microns) (Shepherd et al. 2004), coordinated inhibition only reliably occurred in pairs of closely spaced MCs under our experimental conditions. We also observed a large variation in degree of inhibitory coupling even between pairs of nearby MCs, as illustrated by the range of excess CIF estimates shown in Figure 4F. Serotonin triggered a statistically significant, but transient, increase in excess IPSC CIF that was extinguished rapidly ($\sim 1\mathrm{~min}$ after the IPSC frequency was maximal) (Fig. 4G). Mitral cell excess IPSC CIF was not correlated with the mean IPSC frequency in that episode ($R^2 = 0.20, n = 26$ episodes from 13 MC paired recordings with $<200\ \mu\text{m}$ separation). Together, these results suggest that the spatial extent of inhibitory local circuits revealed by 5-HT is largely restricted to pairs of MCs separated by $<150\ \mu\text{m}$ and is most prominent in pairs separated by $<80\ \mu\text{m}$.

Next, we tested whether other experimental treatments that increased the frequency of spontaneous IPSCs also promoted synchronous inhibition in nearby pairs of MCs. Reducing extracellular Mg$^{2+}$ concentration eliminated the tonic blockade of NMDARs (Ascher and Nowak 1987) and enhanced GC-mediated reciprocal dendrodendritic inhibition (Isaacson and Strowbridge 1998; Schoppa et al. 1998; Chen et al. 2000). Treatment with Mg$^{2+}$-free ACSF increased the frequency of spontaneous IPSCs in MCs ($P < 0.005, n = 8$ experiments) (Fig. 5A–D). Despite the robust increase in inhibitory tone evoked by low Mg$^{2+}$ ACSF, we observed no increase in frequency of synchronous IPSCs assayed through cross-correlation analysis in two example experiments (Fig. 5B) and when IPSC CIF was assayed in four experiments ($P > 0.05$) (Fig. 5D). Even when the CIF analysis was restricted to large-amplitude IPSCs ($>200\ \text{pA}$), that might reflect GABA release events triggered by dendritic spikes in GCs, we did not observe increased IPSC synchronization in low Mg$^{2+}$ ACSF ($P > 0.05$ actual CIF vs. shuffled interval controls, paired $t$-test). These results demonstrate that the synchronous inhibition of MCs triggered by 5-HT was not simply a result of enhanced inhibitory tone. Enhancing excitatory synaptic drive onto interneurons using low Mg$^{2+}$ ACSF elevated the frequency of spontaneous IPSCs, and also triggered many more large-amplitude IPSCs than 5-HT, without promoting synchronous IPSCs in nearby pairs of MCs. Figure 5E summarizes the effects of two different experimental manipulations that increased IPSC frequency (5-HT and low Mg$^{2+}$ ACSF) on synchronous MC inhibition. Only 5-HT increased inhibitory tone and also promoted synchronous inhibition, an effect that was robust over a wide range of CIF analysis window durations from 0.5 to 10 msec. The excess CIF evoked by 5-HT increased as the analysis window was lengthened from 0.5 to 4 msec and then was asymptotic, suggesting that the jitter associated with mechanisms underlying coincident inhibition of MCs was $<4\mathrm{~ms}$. This estimate includes the jitter associated with transmitter release at both inhibitory synapses and the temporal variance associated the processes that link the two release sites.

Finally, we asked if tetanic stimulation in the GCL (Fig. 6A, B) evoked synchronous inhibition in MCs. GCL stimulation often triggered spiking in GCs (e.g., spiking triggered in 12 of 13 cell-attached GC recordings) (Fig. 2E); the mean firing rate was 7.8 Hz $\pm$ 2.2 Hz in five GCs that responded to GCL stimulation throughout the analysis window employed below (100–1100 msec following}
serotonin and coincident inhibition

Discussion

We make three principal conclusions in this study. First, activation of large ensembles of MCs with 5-HT recruits inhibitory local circuits through a 5-HT2R- and AMPAR-dependent mechanism that forms divergent synapses on nearby but not distant pairs of principal neurons. Serotonin-driven inhibitory circuit activation functioned without frequent somatic spiking in GCs. Second, enhancement of dendrodendritic inhibition by removing extracellular Mg2+ ions increases inhibitory tone on MCs but fails to recruit coincident IPSCs on nearby pairs of MCs beyond the frequency expected by chance. Finally, tetanic stimulation of afferents in the GCL triggers robust synchronous inhibition of MCs, even in MC pairs where 5-HT triggered only modest coincident inhibition. These results suggest that different local circuits may regulate inhibition onto nearby MCs than clusters of principal cells that span multiple glomerular modules.

Although a variety of stimulation methods have been used to increase inhibitory tone onto MCs (Rall and Shepherd 1968; Nakashima et al. 1978; Jahr and Nicoll 1982), our results demonstrate that the current is strong evidence that through very different local circuit pathways. Enhancing currents through NMDARs by applying Mg2+-free ACSF increased the frequency of spontaneous IPSCs recorded in MCs but failed to recruit coincident inhibition even in closely spaced pairs of principal cells. This treatment likely facilitated inhibition by directly activating reciprocal dendrodendritic

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**Figure 4.** Serotonin promotes synchronous inhibitory synaptic responses in pairs of nearby mitral cells. (A) Example intracellular responses in near pairs of MCs shown in left panel that include an IPSC recorded in both cells (arrows). IPSC onset latency differed by 0.8 msec in the two MCs, 60-μm spacing between mitral cell bodies). Example recordings in right panel from a distant MC pair with no synchronous IPSCs (740-μm spacing between MCs). (B) Reconstructions of dual MC recordings following intracellular labeling with 150 μM Alexa594. Picture represents collage from multiple maximal Z-stack projections acquired using live two-photon imaging. Neither MC had an intact apical dendrite; arrows indicate primary dendritic processes directed toward the glomerular layer that were truncated at the surface of the brain slice. Calibration bar, 50 μm. (EPL) external plexiform layer, (MCL) mitral cell layer, (GCL) granule cell layer. (C) Inset, example dual voltage-clamp recordings from two MCs that received synchronous IPSCs during 5-HT treatment. Arrow indicates example synchronous IPSC recorded in both MCs (IPSC onset latency difference = 0.4 msec in example shown). Plot of increase in clipped cross-intensity function (CIF) (2-msec analysis window) following 5-HT application. Gray bars represent results from bootstrap shuffles. 5-HT bar represents 140 sec following start of 5-HT application. Post bar represents 200 sec following 5-HT application. (***) *P* < 0.01, paired Student’s *t*-test, *n* = 4 episodes in each condition. Mean ± S.E.M. (D) Plot of MC IPSC synchrony estimated by clipped cross-intensity function (CIF) in near (~200-μm separation between cell bodies), mid (200 to 1000-μm separation), and distant (>1000-μm separation) pairs of MCs. Gray columns represent average CIF computed from IPSC interval shuffles in each paired recording experiment. (****) *P* < 0.005, paired *t*-test. Number of paired recording experiments analyzed indicated at base of each bar. (E) Plot of mean IPSC frequency in control (Cift) and 5-HT in paired MC recordings from the same three groups of experiments presented in D. (****) *P* < 0.005, paired *t*-test. Mean IPSC frequencies in 5-HT did not differ significantly between the three groups (*P* = 0.05, Kruskal–Wallis ANOVA, same MC paired recordings analyzed in D). (F) Plot of excess IPSC CIF (Actual—mean from interval shuffles) vs. mitral cell body separation for 25 paired recording experiments. Dashed line represents no increase in frequency of coincident IPSCs over the frequency expected by chance (excess IPSC CIF = 0). (G) Summary of the increase in excess IPSC CIF following 5-HT in 13 MC pairs separated by ~200 μm; post bar indicates excess CIF at the same time window indicated in C. (*) *P* < 0.05, paired *t*-test.

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The last stimulus). We assayed coincident inhibition in pairs of MCs separated by ~100 μm (range 60–250 μm), a separation distance in which 5-HT only infrequently triggered synchronous inhibition (example traces presented in Fig. 6C). IPSC synchrony was significantly elevated in six of eight paired recordings following a tetanic stimulation (100–1100 msec following the last GCL stimulation, *P* < 0.05); the group mean CIF also was significantly greater following GCL stimulation than CIF computed from shuffled IPSC times (160% increase over bootstrap, *P* < 0.0005) (Fig. 6D). In the same set of eight paired recording experiments, 5-HT triggered only modest IPSC synchrony and a 15% increase in IPSC CIF (*P* < 0.05). When analyzed individually, only two of eight paired recordings had a significant elevation in IPSC CIF beyond the level expected by chance in 5-HT. Four of the six paired recordings that showed no increase in IPSC synchrony in 5-HT had statistically significant increases in IPSC CIF following tetanic stimulation. Excess IPSC CIF also was significantly increased over control conditions following GCL stimulation (*P* < 0.0002) but not following 5-HT (*P* > 0.05; *n* = 8 MC pairs) (Fig. 6E). There was no statistically significant difference in mean IPSC frequency triggered by GCL stimulation and bath application of 5-HT (*P* > 0.05) (Fig. 6F).

One advantage of assaying inhibitory synchrony using the CIF method is the ability to quantitatively determine the minimum test window duration required to capture coincident IPSCs. Simply expanding the duration of the sliding analysis window will always yield more detected coincident events. However, subtracting the bootstrap estimate of the frequency expected by chance for that window generated a metric (excess IPSC CIF) that can be compared across different conditions. As shown in Figure 6G, this metric increased in parallel as the window duration was increased in both 5-HT and GCL stimulation conditions (open and filled symbols, respectively). Both curves were well fit by a model with each pair of near-coincident IPSCs originating from the same interneuron that had a synaptic release jitter of 2 ± 0.5 msec (mean ± SD) (solid line in Fig. 6G; Kay and Wong 1987; Sabatini and Regehr 1999). More complex models of coincident inhibition arising from near-synchronous excitation of multiple interneurons included the jitter associated with IPSC responses on granule cells (1.1–2.3 msec) (Balu et al. 2007) and the jitter associated with EPSP-spike coupling (1.4–4.7 msec) (Schoppa 2006b; Rodríguez-Molina et al. 2007). Even using the EPSP and spike coupling mechanisms described in prior studies with relatively low jitter required significantly longer sliding analysis windows to capture coincident events than we found in our measurements (time constant significantly greater than either 5-HT or GCL stimulation conditions, *P* < 10−4) (dashed line in Fig. 6G). Using the estimate of the jitter associated with EPSP-spike coupling in OB GCs (4.7 msec) (Schoppa 2006b) required even longer duration analysis windows (dotted line in Fig. 6G). These results suggest that following 5-HT and GCL stimulation, the coincident inhibition we observe in MCs likely arises from divergent output from individual interneurons.

Finally, we asked if we could differentiate the spontaneous IPSCs triggered by 5-HT, 0 Mg2+ ACSF, and GCL stimulation. As shown in the plots in Figure 6H, 5-HT triggered IPSCs with a very similar distribution of amplitudes as control conditions (*P* > 0.05, K-S test). Reducing extracellular Mg2+ ions results in a uniform shift toward larger amplitude IPSCs (different from control, *P* < 10−8, K-S test) while GCL stimulation skewed the amplitude distribution toward large-amplitude IPSCs (different from control, *P* < 10−8, different from 0 Mg2+, *P* < 10−8, K-S test). While the uniform enhancement of IPSC amplitude in 0 Mg2+ ACSF could arise from multiple mechanisms, the selective increase in spontaneous large-amplitude IPSCs following GCL stimulation is consistent with the preferential recruitment of a population of IPSCs that was not strongly active under control conditions.
synapses, an inhibitory pathway effectively driven by currents through NMDARs (Isaacson et al. 1998; Isaacson 2001). The absence of coincident inhibition through NMDARs (Isaacson and Strowbridge 1998; Schoppa et al. 1998) and at two time points following application of 0 Mg2+ ACSF (gray curve, separate experiment). (C) Plot of spontaneous IPSC frequency in the four experimental conditions shown in A for eight MCs. (** P < 0.005. (D) Plot of IPSC CIF at two times following application of 0 Mg2+ ACSF in four MC paired recordings (<200-μm cell body separation). Gray bars represent average CIF computed from 10,000 IPSC interval shuffles (Shuf), n.s., P > 0.05. (E) Plot of the relationship between duration of synchronous IPSC detection window and excess CIF (Actual—expected from IPSC interval shuffles) in near MC paired recordings (<200-μm cell body separation). Serotonin (5-HT, black symbols) increased IPSC CIF significantly above expected values for all coincidence detection windows while 0 Mg2+ (gray symbols) did not increase CIF beyond expected values at any coincidence window duration tested. Comparisons at all window durations tested were statistically significant in 5-HT experiments (P < 0.01, paired t-test).

**Figure 5.** Low Mg2+ external solution increase spontaneous MC inhibition without promoting synchronous IPSCs in paired MC recordings. (A) Example traces in one MC under control conditions and at two time points following application of 0 Mg2+ ACSF. (B) Example cross-correlograms computed from spontaneous IPSC onset times illustrating frequent coincident inhibitory inputs promoted by 5-HT (black curve, peak at lag = +0.2 msec) with no correlated inhibitory input triggered by 0 Mg2+ ACSF (gray curve, separate experiment). (C) Plot of spontaneous IPSC frequency in the four experimental conditions shown in A for eight MCs. (** P < 0.005. (D) Plot of IPSC CIF at two times following application of 0 Mg2+ ACSF in four MC paired recordings (<200-μm cell body separation). Gray bars represent average CIF computed from 10,000 IPSC interval shuffles (Shuf), n.s., P > 0.05. (E) Plot of the relationship between duration of synchronous IPSC detection window and excess CIF (Actual—expected from IPSC interval shuffles) in near MC paired recordings (<200-μm cell body separation). Serotonin (5-HT, black symbols) increased IPSC CIF significantly above expected values for all coincidence detection windows while 0 Mg2+ (gray symbols) did not increase CIF beyond expected values at any coincidence window duration tested. Comparisons at all window durations tested were statistically significant in 5-HT experiments (P < 0.01, paired t-test).

electrical shock, we sought to minimize IPSC synchronization mediated by synaptic inputs directly activated by the tetanic stimulus itself.

Activation of MCs with bath 5-HT triggered widespread spiking in MCs and also recruited coincident inhibition in groups of nearby principal cells, primarily in MC pairs separated by <80 μm. The disynaptic depolarization we report in GCs following 5-HT is consistent with an increase in inhibitory tone in MCs mediated by GABA release at dendrodendritic synapses formed between granule and MC dendrites (Isaacson and Strowbridge 1998; Schoppa et al. 1998). However, coincident IPSCs could potentially arise from inhibitory cell types in the glomerular layer. Petzold et al. (2009) described an action of 5-HT on glomerular-layer inhibitory circuits mediated by 5-HT2C receptors on periglomerular interneurons that could contribute to inhibitory tone in MCs despite the electrotonically distant location of these synaptic inputs. Presumably divergent connections formed by GABAAergic periglomerular cells underlie coincident IPSCs recorded in the subclass of TCs located near the glomerular layer (external tufted cells) (Hayar et al. 2005). Several lines of evidence suggest that the modulation of inhibitory tone we observe in MCs is distinct from this effect in the glomerular layer. First, we observe similar increases in MC IPSC frequency in slices in which the glomerular layer was surgically removed. Second, we find a similar, statistically significant increase in IPSC synchronization in MCs with apical dendrites that do not arborize in the same glomerulus. Finally, modulation of periglomerular inhibitory circuits appears to rely primarily on 5-HT2C receptors while modulation of inhibitory tone can be evoked by the selective 5-HT2AR agonist TCB2. Petzold et al. (2009) found no effect on glomerular-layer circuit function when they tested the same 5-HT2AR agonist, a result that is consistent with the high density of 5-HT2ARs found primarily in the MC layer using immunohistochemistry (Hamada et al. 1998) and in situ hybridization (Pompeiano et al. 1994). While there is a consensus in the literature regarding 5-HT2CR expression in the glomerular layer. Petzold et al. 1994; Abrahams et al. 1995; Clemett et al. 2000; Petzold et al. 2009), several of these studies also found 5-HT2CR expression in deeper OB layers.

Unlike inhibitory responses triggered by tetanic GCL stimulation and low Mg2+ ACSF, the IPSC amplitude distribution was not significantly affected by 5-HT. This result suggests that 5-HT affected primarily inhibitory local circuits that were already active spontaneously (or triggered IPSCs that had similar amplitudes to spontaneous IPSCs). Surprisingly, the widespread excitatory synaptic drive elicited by 5-HT did not lead to frequent somatic spikes in GCs. Instead, intracellular recordings revealed these interneurons were typically only modestly depolarized by glutamatergic EPSPs during 5-HT and did not spike spontaneously. However, the low jitter we find associated with coincident MC inhibition driven by both 5-HT and GCL stimulation likely results
Serotonin and coincident inhibition

There are at least two possible mechanisms that could explain 5-HT-stimulated synchronous inhibition in the absence of frequent GC somatic spiking. Serotonin-triggered coincident inhibition could arise from dendritic spikes in GCs that failed to trigger somatic action potentials or from spiking in an as-yet-unidentified inhibitory local interneuron that targets MC cell bodies and/or proximal dendrites. Several studies have reported dendritically localized spiking in GCs, though often involving primarily low-threshold Ca\(^{2+}\) spikes (Egger et al. 2003, 2005). Spontaneous fast Na\(^+\)-based dendritic spikes have been reported in frog GCs under basal (unstimulated) conditions (Zelles et al. 2006). If also present in mammals, spontaneous “D-spikes” could function to synchronize multiple dendrodendritic release sites in one GC, enabling coincident inhibition of nearby pairs of principal cells. The sensitivity of coincident inhibition in S-HT to AMPAR antagonists suggests that these responses were likely recruited selectively by excitatory inputs with fast kinetics. Although we rarely observed spikelets in somatic intracellular recordings from GCs under basal or S-HT-stimulated conditions that might correspond to dendritic D-spikes (Zelles et al. 2006), these events might not be easily detectable given the frequent synaptic inputs mammalian GCs normally receive in S-HT.

Coincident inhibition of MCs also could arise from other (non-GC) interneurons that target MC somata or proximal dendrites, explaining the bias toward synchronous inhibition in near but not far MC pairs. At present, there is little evidence for axo-dendritic or axosomatic inhibition of MCs apart from periglomerular circuits. A wide variety of morphologically defined interneuron subtypes has been identified in the EPL and GCL that could potentially form axonal connections with MCs (Schneider and Macrides 1978; Pressler and Strowbridge 2006; Kosaka and Kosaka 2011; Pressler et al. 2013). Recent studies (Kato et al. 2013; Miyamichi et al. 2013) described a new class of parvalbumin (PV)-immunoreactive interneuron located in the external plexiform layer that could potentially mediate the synchronous IPSCs we observe in MCs in S-HT. Alternatively, one or more subtypes of large “short-axon” interneurons in GCL, such as Blanes cells (Pressler and Strowbridge 2006) and Golgi cells (Pressler et al. 2013), may contact MCs in addition to GCs.

Although our experimental results cannot definitively determine whether coincident inhibition in S-HT arises from dendritic

from spiking in common presynaptic interneurons. Although the blockade of 5-HT-stimulated inhibition on MCs by AMPAR antagonists argues that the primary site of 5-HT action was not on interneurons, our results cannot exclude additional modulatory actions of metabotropic receptors that affect interneuron function.
D-spikes that couple multiple dendrodendritic GABA release sites within individual GCs or from divergent axonal input from another interneuron class, several lines of evidence are suggestive of the latter hypothesis. First, GCs are known to synapse along the length of MC secondary dendrites and, therefore, could potentially provide synchronous inhibition to widely separated pairs of MCs—which we did not observe. In principal, cable attenuation could explain the bias we found toward synchronous inhibition only in closely spaced MC pairs. However, using focal uncaging of GABA, Lowe (2002) recorded IPSCs originating from dendrodendritic synapses on lateral dendrites $\approx 200 \mu m$ from the cell body that were often larger than spontaneously occurring IPSPs. Based on this prior uncaging study, we expected that divergent GC inputs should be detectable in our experimental conditions in MC pairs separated by $300–400 \mu m$. The ability of GCL stimulation to recruit coincident inhibition in a set of paired MC recordings that showed little synchronous inhibition in 5-HT (Fig. 6D,E) also is suggestive of a role for two distinct inhibitory interneuron subtypes outside the glomerular layer. Inhibitory circuits recruited during 5-HT treatment typically linked clusters of MCs separated by $< \approx 100 \mu m$ (likely only one or a few glomerular modules) while local circuits activated by GCL stimulation extended over larger clusters of MCs and included more glomerular modules. The significantly different IPSC amplitude distributions following 5-HT treatment and GCL stimulation also are suggestive of a role for non-GC interneurons.

Multiple investigators have hypothesized that the inhibitory local circuits formed between MCs and GCL interneurons operate on different spatial scales relative to the size of glomerular columns (Egger and Urban 2006, Balu et al. 2007). The smallest spatial scale is likely to be the reciprocal dendrodendritic synapse formed between MC and GC dendrites (Isaacson and Strowbridge 1998; Halabisky et al. 2000) which may function to gate AP propagation along the MC secondary dendrite (Chen et al. 2000). Unexpectedly, we found that widespread activation of these synapses following low Mg$^{2+}$ ACSF triggered robust MC inhibition but little inhibitory synchrony, suggesting that dendrodendritic synapses may function to coordinate MCs on smaller spatial scales than assayed in this study. The other unexpected finding from this study was that activation of large ensembles of MCs did not generate inhibitory synchrony on the same spatial scale as MC dendritic processes, many of which span millimeters (Shepherd et al. 2004). Instead, we find coincident inhibition was restricted to a spatial scale similar to the size of a glomerular column. Given previous work demonstrating the ability of correlated inhibitory synaptic input to drive synchronous firing in MCs (Galán et al. 2006; Schoppa 2006a), a primary prediction from our study is that activation of inhibitory local circuits driven disynaptically by 5-HT should promote to spike synchrony in MCs with same or neighboring glomerular modules. Unfortunately, it is not possible to directly test this prediction using the experimental paradigm used in this study because of the direct depolarizing effects of 5-HT on MCs. In principle, optogenetic methods that selectively depolarize the same interneuron population that was disynaptically excited by 5-HT could test this hypothesis though additional studies will be required to identify the appropriate interneuron subclass.

Materials and Methods

Slice preparation

Horizontal OB slices (300 $\mu m$ thick) were prepared from P14–P19 Sprague-Dawley rats, either sex, as previously described (Halabisky et al. 2000; Friedman and Strowbridge 2003; Balu et al. 2007). Slices were incubated in ACSF at 35°C for 30 min and then maintained at room temperature. All experiments were carried out in accordance with the guidelines approved by the Case Western Reserve University Animal Care and Use Committee.

Electrophysiology

All recordings were performed in a submerged recording chamber. Most recordings were performed at 34°C–35°C; experiments visualizing GC dendritic arbors using two-photon imaging were performed at 30°C. Slices were continuously perfused with an extracellular solution containing (in mM): 124 NaCl, 5 KCl, 1.23 Na$_2$HPO$_4$, 1.2 MgSO$_4$, 26 NaHCO$_3$, 10 dextrose, 2.5 CaCl$_2$, equilibrated with 95% O$_2$/5% CO$_2$ (pH 7.3) at a flow rate of 1.5–2.0 mL/ min. Whole-cell patch-clamp electrophysiological recordings were made using AxoPatch 1B or 1D amplifiers (Axon instruments) and borosilicate glass pipettes ($2–8 \Omega$). All data were digitized at 5 or 10 kHz by an ITC-18 computer interface (Instrutech) and a Windows 7 personal computer. Slices were imaged using IR/DIC optics on an upright fixed stage Zeiss Axioskop 1FS microscope and cells were identified based upon morphology, soma location within a slice, and intrinsic membrane properties. For current-clamp whole-cell and cell-attached recordings, the internal solution contained (in mM): 140 K methylsulfate, 4 NaCl, 10 HEPES, 0.2 EGTA, 4 MgATP, 0.3 Na$_2$GTP, 10 phosphate-Tris. A CsCl-based internal solution was used for all MC voltage-clamp recordings that reversed the Cl$^-$ gradient; it contained (in mM): 115 CsCl, 4 NaCl, 10 HEPES, 1 EGTA, 25 TEA-Oh, 0.5 QX-314, 4 MgATP, 0.3 Na$_2$GTP, 10 phosphate-Tris. Tris. All internal solutions were equilibrated to a pH of 7.3 and an osmolarity of $\approx 290$ mmol/kg. Serotonin phosphocreatine was prepared for use daily from powder (Sigma Aldrich). All other drugs were purchased from Tocris and were prepared in aliquots and frozen at $\approx 20^\circ$C until use. Drugs were added to the bath by changing the source of external solution. Evoked responses were obtained using a constant current stimulus isolation unit (A-360, World Precision Instruments) and a monopolar tungsten electrode (FHC) placed in the GC layer. Neurons filled with Alexa594 through the patch pipette were imaged using a custom two-photon laser scanning system (Balu et al. 2007).

OB slice microdissection

Horizontal OB slices lacking a glomerular layer were prepared by first making two cuts perpendicular to the mitral cell layer (MCL) that extended through the entire slice, leaving a dissected slice $\sim 2$ mm wide. The glomerular layer was then separated from the remaining tissue by making a third cut, parallel to the MCL, through the EPL (Friedman and Strowbridge 2003). All microdissection cuts were performed in a submerged recording chamber while visualizing the slice through a 5× objective. Dissected slices were allowed to recover for at least 20 min before attempting recordings. Dissected slices were stained with methylene blue dissolved in ACSF at the conclusion of each experiment to visualize the OB laminar structure. Only recordings from slices in which the glomerular layer was completely removed were analyzed.

Data analysis

Data analysis was performed with custom Matlab 2012b (Mathworks) scripts. Inhibitory and excitatory postsynaptic currents (IPSCs, EPSCs), and action potentials were automatically detected as previously described (Larimer and Strowbridge 2008). Briefly, traces were smoothed via Savitsky–Golay filter and the instantaneous derivative was calculated. Events were defined by a cumulative dV/dt or dI/dt that exceeded a set threshold, and event times were defined by the first deviation toward threshold. Average frequencies of spikes or synaptic events were determined by taking the reciprocal of the average inter-event interval. All data are presented as mean $\pm$ S.E.M. unless otherwise noted. The paired Student’s t-test was used to determine statistical significance, unless otherwise specified. Autocorrelation and cross-correlation were
performed by detecting events and convolving a 2-msec Gaussian kernel with the binary event times (Galan et al. 2006).

The clipped cross-intensity function (CIF) (Hahnloser 2007) was computed based on detected IPSC event times and used to determine whether the incidence of synchronous events was greater than expected by chance. To calculate the CIF for two response channels, we computed the total number of coincident events (event onset latencies that differed by less than the synchronous window cutoff of 2-msec events, except for the analyses in Figures 5 and 6 that used varying duration windows) and divided by the total number of events on the channel with the fewest events. For example, one paired MC recording episode that contained 200 detected IPSCs in MC1 and 250 IPSCs in MC2, with 50 IPSCs that each occurred within 2 msec on both channels, would have an IPC CIF of 0.15 (30/200). Two advantages of the CIF method, compared with correlation-based methods for detecting synchrony, are the ability to quantitatively define event synchrony over variable coincidence stringencies and the ability to directly compute estimates of synchrony expected by chance. Unless noted, all analyses were based on the average CIF computed over three control (pre 5-HT) and two consecutive 30-sec episodes acquired between 140 and 210 sec after switchover to ACSF containing 5-HT. We computed CIF expected by chance for each episode by permuting the inter-event intervals contained in each channel; bootstrap CIF estimates were computed over 10^6 to 10^7 iterations. Bar plots labeled shuffled CIF reflect the mean of the bootstrap runs computed from each episode. “Excess” CIF was computed by subtracting the mean bootstrap CIF from the CIF computed from the actual event timing. Probability estimates obtained from individual episodes were computed from cumulative distribution functions based on the bootstrap estimates.

Estimates of excess IPC CIF from interneurons that form divergent synaptic connections with MCs and clusters of interneurons synchronized by common excitatory input (Fig. 6G) were generated by a common template of 100 exactly synchronous events across two MCs and distributed with Poisson inter-event intervals. For the divergent interneuron model (solid line in Fig. 6G), a separate IPSC releases jitter values drawn from a Gaussian distribution with a mean of 2.0 msec and a SD of 0.5 msec (Kay and Wong 1987; Sabatini and Regehr 1999) was added to each simulated MC channel. We constructed two models of nondivergent interneurons coupled by common excitatory inputs. The nondivergent model with lowest jitter (dashed line in Fig. 6G) was made by including glutamate release jitters of 1.1 ± 0.2 msec (Balu et al. 2007) on each interneuron channel as well as the jitter associated with EPSP-spike coupling in neocortical neurons (1.4 ± 0.9 msec) (Rodriguez-Molina et al. 2007). Using the less reliable EPSP-spike coupling reported in OB GCs (4.7 ± 1.5 msec) (Schoppe 2006b) and the jitter associated with dendrodendritic EPSs onto GCs (2.3 ± 3.2 msec) (Balu et al. 2007) required much longer sliding windows to capture synchronous IPSCs (dotted line in Fig. 6G).

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