Calcium Signaling in Mitral Cell Dendrites of Olfactory Bulbs of Neonatal Rats and Mice During Olfactory Nerve Stimulation and β-Adrenoceptor Activation

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Synapses formed by the olfactory nerve (ON) provide the source of excitatory synaptic input onto mitral cells (MC) in the olfactory bulb. These synapses, which relay odor-specific inputs, are confined to the distally tufted single primary dendrites of MCs, the first stage of central olfactory processing. β-adrenergic modulation of electrical and chemical signaling at these synapses may be involved in early odor preference learning. To investigate this possibility, we combined electrophysiological recordings with calcium imaging in olfactory bulb slices prepared from neonatal rats and mice. Activation of ON–MC synapses induced postsynaptic potentials, which were associated with large postsynaptic calcium transients. Neither electrical nor calcium responses were affected by β-adrenergic agonists or antagonists. Immunochemical analysis of MCs and their tufted dendrites revealed clear immunoreactivity with antibodies against α1A (Cav2.1, P/Q-type) and α1B (Cav2.2, N-type), but not against α1C (Cav2.2, L-type) or α1D (Cav3.1, L-type) calcium channel subunits. Moreover, nimodipine, a blocker of L-type calcium channels, had no effect on either electrical or calcium signaling at ON–MC synapses. In contrast to previous evidence, we concluded that in neonatal rats and mice (P5–P8), mitral cells do not express significant amounts of L-type calcium channels, the calcium channel type that is often targeted by β-adrenergic modulation. The absence of β-adrenergic modulation on either electrical or calcium signaling at ON–MC synapses of neonatal rats and mice excludes the involvement of this mechanism in early odor preference learning.

Early odor preference learning is observed in neonatal rats when they form a preference to an odor that is paired with a reinforcing stimulus (Sullivan et al. 1988; Sullivan and Wilson 1994; Wilson and Sullivan 1994). This form of learning depends on noradrenergic neurotransmission in the main olfactory bulb (MOB) that receives noradrenergic input exclusively from the locus coeruleus (Shipley et al. 1985; McLean et al. 1989; McLean and Shipley 1991). Both α and β-adrenoceptors are localized in the MOB (Wanaka et al. 1989; Nicholas et al. 1993a,b; Woo and Leon 1995; Day et al. 1997; Winzer-Serhan et al. 1997; Yuan et al. 2003) and might therefore contribute differentially to early odor preference learning. Although work in vitro (Trombley 1992, 1994; Trombley and Shepherd 1992) and in vivo (Jiang et al. 1996) has indicated norepinephrine (NE)-mediated disinhibition via α-adrenoceptors, a role of α-adrenoceptors in early odor preference learning has only been recently proposed (McLean et al. 2003). On the other hand, activation of β-adrenoceptors has been shown to promote learning-dependent behavioral and neural changes (Wilson and Sullivan 1994; Coopersmith and Leon 1995). Early odor preference learning in the MOB is blocked by β-adrenoceptor antagonists; and the effect of reinforcing stimulus required for early odor preference learning can be provided by application of β-adrenoceptor agonists. On the basis of these findings, it has been proposed that β-adrenoceptor activation is both necessary and sufficient for early odor preference learning (Sullivan et al. 2000; but see McLean et al. 2003).

In a recent cellular model for early odor preference learning, Yuan et al. (2003) proposed that β-adrenoceptor activation triggers an intracellular cAMP cascade, whereas ON input triggers an influx of Ca2+ in MC dendrites. The coincident activation of the cAMP cascade and increased Ca2+ levels facilitates Ca2+/calmodulin phosphorylation of cAMP response element (CRE)-binding protein (CREB) and CRE-regulated gene transcription. In this model, the cAMP-UCS (unconditioned stimulus) and Ca2+-CS (conditioned stimulus) pathways are hypothesized to converge at the level of CREB phosphorylation/dephosphorylation. There are several possible scenarios, however, for an interaction of the two pathways at an earlier stage. First, the Ca2+ pathway may enhance the cAMP cascade. This possibility has been excluded because odor input itself did not change the cAMP level. Furthermore, odor plus stroking (tactile stimulation) did not provide an additional increase to the cAMP level compared with stroking alone (Yuan et al. 2003). The second possibility for an interaction of the cAMP and the Ca2+ pathways at an early stage is the direct enhancement of the odor-induced Ca2+ signaling via the β1-adrenoceptor.

In both the hippocampus and the amygdala, the modulation of Ca2+ signaling by β-adrenoceptors has been proposed to play a critical role during long-term memory formation (Gray and Johnston 1987; Huang et al. 1993, 1996). The pairing of CS and UCS inputs during fear conditioning leads to Ca2+ entry through both NMDA receptors and L-type Ca2+ channels in lateral amygdala principal neurons (Bauer et al. 2002). Activation of β-adrenoceptors has been reported to facilitate glutamate release in the amygdala by increasing presynaptic Ca2+ influx (Huang et al. 1993, 1996). In hippocampal pyramidal cells, NE and β-adrenoceptor agonists increase the activity of postsynaptic L-type calcium channels (Gray and Johnston 1987); the activation of L-
type \( \text{Ca}^{2+} \) channels, but not NMDA receptors, leads to sustained nuclear CREB phosphorylation and CREB-driven gene expression (Hardingham et al. 2001).

In the current study, we combined intracellular recordings and \( \text{Ca}^{2+} \) imaging with immunocytochemical analysis to test whether \( \beta \)-adrenergic modulation on both electrical synaptic transmission between the ON and MCs and synthetically induced \( \text{Ca}^{2+} \) signals in MC dendritic tufts of neonatal rats and mice. We therefore exclude the possibility of an early interaction model for early odor preference learning.

**RESULTS**

We used olfactory slices prepared from both mouse and rat pups. Using both of these species, we took into account the fact that most behavioral work is done in rats, whereas the number of studies using genetic approaches in mice is increasing. Although data from different species were not pooled, no obvious differences were found in data between rats and mice. The morphology of postnatal development of rat MCs has been previously described (Malun and Brunjes 1996; Treloar et al. 1999). In agreement with these studies, dye-filled mouse MCs between postnatal day 5 (P5) and P8 exhibit a distal dendritic tuft less developed than that in adults, and occasion ally also exhibit a secondary apical dendrite. Figure 1 shows a dye-filled P6 mouse MC with a single apical dendrite along with a schematic drawing indicating the patch-clamp electrode and placement of the stimulation electrode on the ON. A single electrical stimulus to the ON induced an excitatory postsynaptic potential (EPSP) recorded at the soma of MCs. The EPSP gradually increased in amplitude with increasing stimulus intensity and eventually reached threshold for triggering action potentials (Fig. 2A).

Calcium imaging revealed that both sub- and supra-threshold EPSPs were associated with a transient rise in \( [\text{Ca}^{2+}] \), in the tufted dendritic terminal. Maps of these \( \text{Ca}^{2+} \) signals indicated that the amplitude of \( \text{Ca}^{2+} \) transients was relatively homogenous over the whole tuft, but slightly smaller in the adjacent apical dendrite segment (Fig. 2C).

The \( \text{Ca}^{2+} \) transients associated with synaptic transmission between ON and MCs may be caused by \( \text{Ca}^{2+} \) influx through NMDA receptors or by activation of voltage-gated calcium channels (VGCCs). To investigate these possibilities, we applied the NMDA receptor antagonist D-APV, which reduced, but did not abolish, the EPSPs and associated calcium transients (Fig. 3). Both signals were abolished with the additional application of an AMPA receptor blocker, NBQX (data not shown), indicating that, in the absence of NMDA receptor activation, activation of AMPA receptors suffices to produce a \( \text{Ca}^{2+} \) elevation via activation of VGCCs. The effect of D-APV demonstrates a significant contribution of NMDA receptors to the compound EPSP. The reduction of the \( \text{Ca}^{2+} \) signal by D-APV could be accounted for by the reduced postsynaptic depolarization (see Fig. 1). The close correlation between the amplitudes of postsynaptic membrane depolarization and \( \text{Ca}^{2+} \) signals (Fig. 1) suggests...
that the bulk of Ca\(^{2+}\) influx associated with the EPSP is mediated by VGCCs. However, we cannot exclude the possibility that a component of the Ca\(^{2+}\) transients was caused by flux through NMDA receptors or calcium released from intracellular sources.

We then studied the effects of \(\beta\)-adrenoceptor agonists and antagonist on ON stimulation-induced glomerular responses in neonatal rats (Fig. 4). Three drugs were used as follows: isoproterenol (1–10 µM, a standard selective \(\beta\)-adrenoceptor agonist, \(n = 4\)), xamoterol (1 µM, a partial \(\beta_1\)-adrenoceptor agonist, \(n = 2\)) and betaxolol (1 µM, a selective \(\beta_1\)-adrenoceptor antagonist, \(n = 4\)). None of the above drugs exerted significant effects on the ON–MC EPSPs or on the associated Ca\(^{2+}\) signals. We confirmed these results in olfactory slices from neonatal mice by testing the effect of isoproterenol on ON stimulation-induced glomerular responses either sub- or supra-threshold for action potential induction. Isoproterenol did not affect the electrical or the Ca\(^{2+}\) signaling in MCs in either condition.

Our results were surprising, as previous work conducted in other laboratories suggested that L-type calcium channels are present in MCs (Trombley 1992; Tanaka et al. 1995; Wang et al. 1996; Davila et al. 2003), and it is generally assumed that these channels are modulated by \(\beta\)-adrenoceptor activation in other cortical structures (Gray and Johnston 1987; Huang et al. 1993). Because at least \(\beta_1\)-adrenoceptors are expressed in MCs (Yuan et al. 2003), we reinvestigated the expression of Ca\(^{2+}\) channel subtypes in MCs using immunocytochemical and pharmacological measures. Immunocytochemistry revealed clear immunoreactivity with antibodies against \(\alpha_1\)C (Cav2.1, P/Q-type) and \(\alpha_1\)B (Cav2.2, N-type), but not against L-type \(\alpha_1\)C (Cav1.2)/\(\alpha_1\)D (Cav1.3) Ca\(^{2+}\) channel subunits in MCs and their tufted dendrites (Fig. 6). Whereas \(\alpha_1\)A and \(\alpha_1\)B antibodies clearly labeled MC soma and glomeruli (also granule cell soma for \(\alpha_1\)B; Fig. 6A,B), no obvious staining was observed in mitral cell soma and glomeruli by L-type VGCC subunit \(\alpha_1\)C antibody (Fig. 6C). \(\alpha_1\)D immunocytochemistry revealed very weak immunoreactivity throughout the MOB (Fig. 6D). The external and internal plexiform layers, however, were heavily stained by the \(\alpha_1\)C antibody, suggesting that granule cell dendrites (or MC lateral dendrites, see Discussion) express L-type \(\alpha_1\)C subunits of VGCCs. The lack of immunoreactivity for \(\alpha_1\)C (Cav1.2) and \(\alpha_1\)D (Cav1.3) subunits suggest that L-type Ca\(^{2+}\) channels are absent or poorly expressed in the cell body and apical dendrites of MCs. To investigate this issue further, we tested the effect of nimodipine (an L-type Ca\(^{2+}\) channel blocker) on electrical and Ca\(^{2+}\) signaling mediated by ON–MC synapses. Nimodipine (10 µM) had no significant effect on either the ON–MC EPSPs or the associated Ca\(^{2+}\) signals (Fig. 7A,B). Nimodipine also did not affect field EPSPs induced by ON stimulation recorded in the glomerular layer (Fig. 7C,D).

**DISCUSSION**

Given the critical role of \(\beta\)-adrenoceptor activation in early odor preference learning, it is surprising that the physiological mechanisms associated with \(\beta\)-adrenoceptor activation in MCs of the MOB have not been elucidated in detail. The present study revealed that \(\beta\)-adrenoceptor activation interferes...
neither with synaptic transmission between ON and MCs nor with Ca²⁺ signaling in the distally tufted apical dendrites of MCs.

Involvement of Adrenoceptors in Early Odor Preference Learning
In an early hypothesis accounting for the role of noradrenergic mechanisms in early odor preference learning. Wilson and Sullivan (1994) proposed that NE may act on granule cells to disinhibit MCs and modify the synaptic efficacy between the MCs and granule cells. Other studies suggested that the disinhibitory effect of NE on MCs is mediated through α-adrenoceptors (Trombley 1992, 1994; Trombley and Shepherd 1992; Jiang et al. 1996). More recently, Yuan et al. (2003) proposed that NE via β₁-adrenoceptors mediates the production of Ca²⁺ signaling induced in lateral dendrites via L-type calcium channels. Preliminary experiments (n = 3 cells; data not shown) revealed, however, that nimodipine also had no effect on Ca²⁺ transients induced in lateral dendrites by back-propagating action potentials that were induced through depolarization of the cell body. These results are consistent with two previous studies showing little effect of an L-type Ca²⁺ channel blocker, nifedipine, on either mitral cell self-excitation (Friedman and Strowbridge 2000) or lateral dendroexcitatory inhibition (Isaacson and Strowbridge 1998).

In summary, our results exclude the role of β-adrenoceptor activation in directly regulating Ca²⁺ signals in MCs induced by the ON stimulation. Thus, our results suggest that the CAMP-UCS and Ca²⁺-CS pathways may converge solely at the stage of CREB phosphorylation in early odor preference learning.

β-Adrenoceptor Activity, Electrical and Calcium Signaling at ON–MC Synapses
Our data demonstrated that β-adrenoceptor activation did not affect EPSPs recorded from MCs by ON stimulation, which is consistent with a previous study (Hayar et al. 2001). Furthermore, β-adrenoceptor activation did not affect Ca²⁺ signals induced by ON stimulation and recorded in MC apical dendrites. There are no previous studies that have tested the effect of β-adrenoceptor activation on Ca²⁺ signaling in MCs in slices. Previous work by Trombley (1992, 1994) on cultured olfactory neurons suggested that NE inhibits synaptic transmission from MCs to granule cells by reducing Ca²⁺ currents via α-adrenoceptors. Our working hypothesis—the β-adrenoceptor activation may modulate Ca²⁺ signaling in MC dendrites—was motivated by findings reported in other systems, in which the up-regulation of Ca²⁺ signals through L-type VGCCs by β-adrenoceptor activation is involved in learning and memory (Huang et al. 1993, 1996; Bauer et al. 2002). The lack of β-adrenoergic modulation of Ca²⁺ signals in the mitral cell apical dendrite may be explained by three possibilities as follows: (1) β-adrenoceptors do not colocalize with L-type VGCCs, (2) β-adrenoceptors colocalize with L-type VGCCs, but do not interact with them, or (3), L-type calcium channels are absent in the apical dendritic tufts of MCs. To test the above possibilities, we performed immunocytochemical and pharmacological experiments to locate VGCCs, especially L-type VGCCs, in the olfactory bulb, and to test their role in the ON stimulation-induced Ca²⁺ changes.

In contrast to P/Q (α1A) and N (α1B) subunits, which were densely stained in MC soma and glomeruli, the L-type (α1C and α1D) antibody did not stain either of these structures. The lack of immunoreactivity for L-type calcium subunits in MC soma and glomeruli, in which the MC sends its apical dendrite, may explain the lack of effect of β-adrenoceptor activation on Ca²⁺ signaling in MC tufted dendrites in glomeruli. To support the immunocytochemical evidence, we tested the effect of nimodipine (a L-type VGCC blocker) on electrical signaling induced by ON-MC synapses. Nimodipine had no significant effect on either the EPSPs or the associated calcium signals. Whereas these electrophysiological data do not exclude the possibility of an involvement of L-type calcium channel in activities of MCs that were not tested in the present study, they confirm the lack of involvement of L-type VGCCs in the Ca²⁺ signaling in MCs during ON stimulation. The immunoreactivity for α1C subunits of VGCCs in the external plexiform layer may indicate the presence of L-type VGCCs in MC lateral dendrites. Preliminary experiments (n = 3 cells; data not shown) revealed, however, that nimodipine also had no effect on Ca²⁺ transients induced in lateral dendrites by back-propagating action potentials that were induced through depolarization of the cell body. These results are consistent with two previous studies showing little effect of an L-type Ca²⁺ channel blocker, nifedipine, on either mitral cell self-excitation (Friedman and Strowbridge 2000) or lateral dendroexcitatory inhibition (Isaacson and Strowbridge 1998).

In summary, our results exclude the role of β-adrenoceptor activation in directly regulating Ca²⁺ signals in MCs induced by the ON stimulation. Thus, our results suggest that the CAMP-UCS and Ca²⁺-CS pathways may converge solely at the stage of CREB phosphorylation in early odor preference learning.
MATERIALS AND METHODS

Slice Preparation

Experiments were performed on slices obtained from Sprague-Dawley rats and ICR mouse pups of both sexes aged between P5 and P8. Briefly, animals were anesthetized in ice and decapitated. The brain with the two olfactory bulbs was removed and glued to the stage of a Vibroslicer (VT 1000S, Leica). The brain was positioned so that the bulbs were in approximately the same horizontal plane with the most basal part of the brain. Slices (250–300-μm thick) were cut and allowed to recover at 32°C for 1 h, and then at room temperature for up to 8 h. After at least 1 h of incubation, slices were transferred into a recording chamber and were perfused with artificial cerebrospinal fluid (ACSF; composition in mM: NaCl 118, NaHCO₃ 25, NaH₂PO₄ 1, KCl 10, MgCl₂ 1, CaCl₂ 2, and glucose 10). For electrophysiological and fluorescence recordings, slices were placed in an immersion-type perfusion chamber mounted on the stage of an upright microscope (Axioskope 2, Carl Zeiss) and visualized using a 40× water-immersion lens.

Electrophysiology

Patch-clamp experiments were performed in whole-cell configuration. Patch electrodes (3.5–4.5 MΩ resistance) were pulled from borosilicate glass using a two-stage vertical puller (PP-810, Narishige) and contained (in millimolars) KCl 9, KOH 10, K-gluconate 120, MgCl₂ 3.48, NaCl 4, HEPES 10, sucrose 17.5, Na₂ATP 4, Na₃GTP 0.4, and 0.1 Oregon Green 488 BAPTA-1 (pH 7.25) osmolarity 300 mOsm. Extracellular field recordings were recorded with a glass pipette (filled with ACSF, 1.0–2.0 MΩ) pulled from borosilicate glass and placed in the center of a glow-mercury. Glass pipettes (0.8–1.5 MΩ) pulled from borosilicate glass and filled with ACSF were used for electrical stimulation (negative current pulse, 5–100 μA, 300 μs).

Imaging

Fluorescence of Oregon Green was excited by epi-illumination (488 nm) with light provided by a monochromator, and detected by a cooled 12-bit charge coupled device (CCD) under the control of Till Vision software (Till Photonics). A filter set consisting of a dichroic beam splitter (DCLP 505 LP) and an emission filter (535 ± 25 nm) was used. Fluorescence from intracellularly loaded dye equilibrated throughout the cell within 30–40 min of commencing whole-cell configuration. Changes of [Ca²⁺], were expressed as relative fluorescence changes (ΔF/ΔF, where F is the fluorescence before stimulus and ΔF is the evoked change in fluorescence). For traces showing the time course of ΔF/ΔF, fluorescence was measured as the integral over the entire tufted dendrite. Color-coded maps of ΔF/ΔF were obtained using custom-made macros under Image-Pro Plus (Media Cybernetics). For traces showing the time course of ΔF/ΔF, fluorescence was expressed as the integral over the entire tufted dendrite. Color-coded maps of ΔF/ΔF were obtained using custom-made macros under Image-Pro Plus (Media Cybernetics).

Statistics

Summary data are expressed as mean ± SEM; and n represents the number of experiments performed on different MCs. Significance was determined using Student’s unpaired t test, unless stated otherwise.

Immunocytochemistry

P7 rats and mice were anesthetized with an overdose of pentobarbital sodium and perfused transcardially with ice-cold saline, followed by 4% paraformaldehyde in 0.1 M PB (pH 7.4) for 30 min. Brains were removed and post-fixed in the same fixative for 1 h, then transferred to 30% sucrose solution overnight. The olfactory bulbs were sectioned with a cryostat at 30 μm. Olfactory bulb sections were either collected free-floating, or mounted onto subbed slide glass. After incubation with 10% normal goat serum for 30 min at room temperature, sections were incubated overnight in one of the following antibodies: a1A (P/Q-type), a1B (N-type), a1C, and a1D (L-type) (diluted 1:200, Chemicon) in PBS containing 0.3% Triton X-100 and 2% normal goat serum. After washing with PBS, sections were incubated with a secondary antibody (Alexa488, diluted 1:1000, Molecular Probes). Immunoreactions were observed under a confocal microscope (Olympus Fluoview). To establish the specificity of the immunostaining, a negative control test (incubation without primary antibody) or a preabsorption test was performed. All labeling interpreted as specific immunoreactivity was absent in these controls.

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