Development of Spontaneous Miniature EPSCs in Mouse AVCN Neurons During a Critical Period of Afferent-Dependent Neuron Survival

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INTRODUCTION

Synaptic activity has two basic forms: evoked and spontaneous. For the auditory system, evoked synaptic activity refers to transmission that occurs in response to acoustic stimuli from the environment. Spontaneous activity, defined here as synaptically driven signal, can be further divided into two categories: action-potential-dependent and -independent. Action potential-dependent spontaneous activity, elicited by the invasion of action potentials into the presynaptic terminal, can be as large in size as evoked synaptic responses due to the release of many transmitter vesicles. In contrast, action potential-independent synaptic transmission gives rise to small postsynaptic responses (miniature events). These responses are mostly due to the release of a single transmitter vesicle that may reside in a different pool than the ones responsible for evoked release (Frerking et al. 1997; Sara et al. 2005).

Spontaneous activity in developing neurons is important in maintaining cell survival in the visual system (Catsicas et al. 1992; Chytrova and Johnson 2004; Galli-Resta et al. 1993), and some studies suggest that this may be due to neurotrophic factors co-released during neurotransmission (reviewed in Altar and DiStefano 1998; Davies 1994). Similarly, in the auditory system, conductive hearing loss does not result in cell death in the cochlear nucleus of chickens (Tucci et al. 1987) or monkeys (Doyle and Webster 1991), indicating that spontaneous activity originating from the hair cells or ganglion neurons may be sufficient to maintain central auditory neurons. Interestingly, conductive hearing loss leads to an increase in excitatory responses of ventral cochlear nucleus to sound stimuli given to the contralateral ear, and the spontaneous firing rate of those neurons increases (Sumner et al. 2005). Roles of spontaneous synaptic activity in the auditory system also have been implicated in studies using deaf animals. Single-unit recordings from the auditory nerve in deaf cats showed that spontaneous spikes exist, although hair cells are lacking and most ganglion cells degenerate (Ryugo et al. 1998). These studies did not distinguish between action-potential-dependent and -independent spontaneous synaptic activity and may include both.

Since miniature synaptic events were first observed at neuromuscular junctions (Fatt and Katz 1950, 1952), they have been considered as nonfunctional neuronal activity due to their randomness of occurrence and small amplitude. This view has been challenged recently (reviewed in Otsu and Murphy 2003). Sharma and Vijayaraghavan (2003) provided evidence showing that miniature activity can be regulated by nicotinic receptors on the presynaptic terminals to such a level that elicits action potentials in the postsynaptic neuron, influencing signaling across synapses. Miniature synaptic activity is able to maintain dendritic spines of hippocampal neurons via activation of AMPA receptors (McKinney et al. 1999) and regulate dendritic protein synthesis in cultured hippocampal neurons (Sutton et al. 2004). These studies suggest that miniature synaptic activity may play important roles that have been poorly understood. Whether miniature synaptic activity plays any roles in central auditory neurons is unknown.

In both birds and mammals, deprivation of auditory inputs in young animals results in many changes in the structure and function of cochlear nucleus neurons, followed by neuronal death in the cochlear nucleus of chickens (Tucci et al. 1987) or monkeys (Ryugo et al. 1998). These studies did not distinguish between action-potential-dependent and -independent spontaneous synaptic activity and may include both.

Methods

Slice preparation and electrophysiological recording

Brain slice experiments (200–250 μm thickness) were performed from C57BL/6 mice at ages P7, P14, and P21, using methods modified from Tzounopoulos et al. (2004) and Brew and Forsythe (1995). Briefly, after decapitation, a brain block containing the brain stem and the cerebellum was dissected out in ice-cold bicarbonate-buffered solution containing (in mM) 230 sucrose, 3 KCl, 1.2 KH₂PO₄, 20 NaHCO₃, 1.2 CaCl₂, 5 MgSO₄, 3 HEPES, and 10 dextrose with pH of 7.4 and osmolarity around 290 mOsm/l. The brain block was then glued to a vibratome platform (Technical Products, Saint Louis, MO), which was filled with the same solution, and kept at 4°C until use. The brain block was then stored in a cooled room (Technical Products, Saint Louis, MO) until use. The brain block was then cut using a vibratome (Technical Products, Saint Louis, MO) into 200–250 μm thick slices, which were then placed in a 0.5-ml chamber mounted on a Zeiss Axioskop FS (Zeiss, Germany) with a ×40 water-immersion objective and infrared, differential interference contrast optics and continuously superfused with artificial cerebrospinal fluid (ACSF) at a rate of 2–3 ml/min. The ACSF contained (in mM) 130 NaCl, 3 KCl, 1.2 KH₂PO₄, 20 NaHCO₃, 2.4 CaCl₂, 1.3 MgSO₄, 3 HEPES, and 10 dextrose and was constantly gassed with 95% O₂-5% CO₂, with pH of 7.4 and osmolarity ~290 mOsm/l. All recordings were performed at a temperature of 34–36°C, controlled by a feedback temperature controller (Warner Instrument, Hamden, CT). Patch pipettes were pulled to 1–2 μm tip diameter using calibrated (75 μm) glass (VWR, West Chester, PA) and a vertical pipette puller (David Kopf Instruments, Tujunga, CA) and coated with silicone elastomer (Sylgard; Dow Corning, Midland, MI). Electrodes had resistances between 3 and 7 MΩ when filled with an internal solution containing (in mM) 140 K-gluconate, 4.5 MgCl₂, 4.4 tris-phosphocholine, 9 HEPES, 5 EGTA, 4 Na-ATP, and 0.3 tris-GTP, pH 7.3 adjusted by KOH (1 M), and osmolarity of ~290 mOsm/l. A liquid junction potential of 10 mV was corrected.

Voltage-clamp experiments were performed with an Axopatch 200B and current-clamp experiments with an AxoClamp 2B amplifier (Molecular Devices, Union City, CA). Development of firing properties of AVCN neurons was examined using data obtained with the AxoClamp 2B amplifier. The fast current-clamp configuration of the AxoPatch 200B was also used to examine firing properties to determine cell types before acquiring voltage-clamp data, but current-clamp data obtained using the AxoPatch 200B were not included in analysis of firing properties because of known distortions of action potentials by the patch-clamp amplifier (Magistretti et al. 1996). Series resistance was compensated by 80–90% in voltage-clamp recordings. Data were low-pass filtered at 5–10 kHz and digitized with an ITC-16 (Instrutech, Port Washington, NY) at 20–50 kHz for both on and off-line analysis. Data acquisition was done using Axograph software, version 4.5 (Molecular Devices).

All chemicals and drugs were obtained from Sigma (St Louis, MO) except SR 95531 hydrobromide (gabazine) and sucrose, which were obtained from Tocris (Ballwin, MO) and B. J. Baker (Phillipsburg, NJ), respectively. Drugs were bath-applied.

Recording and analysis of mEPSCs

Immediately after obtaining whole cell configuration, firing patterns in response to current injection were examined under the fast current-clamp configuration to determine cell types; bushy cells fire one to three brief spikes in response to prolonged depolarizing current injections, whereas stellate cells fire spikes throughout the duration. Recording mode was then switched to voltage clamp, and spontaneous activity was chart-recorded at a holding potential of ~60 mV. Besides glutamatergic excitatory inputs from the auditory nerve, bushy cells also receive inhibitory inputs from a variety of sources including vertical neurons in the dorsal cochlear nucleus, the trapezoid nucleus, and the superior parvoalveolar nucleus. Stellate cells also receive inhibitory inputs from vertical neurons as well as local interneurons. Glycine is the major inhibitory transmitter at these synapses (reviewed in Ferragamo and Oertel 2001). Spontaneous excitatory postsynaptic currents (sEPSCs) were isolated in the presence of strychnine (1 μM) and gabazine (10 μM), which block glycinegic and GABAergic responses, respectively. Adding tetrodotoxin, a voltage-gated sodium channel blocker, does not change sEPSCs at mouse AVCN synapses, indicating that sEPSCs represent miniature EPSCs (mEPSCs) (Oleskevich and Wahlmsley 2002).

To detect mEPSCs, we slid a template through the chart recordings. The parameters of the template, including amplitude, 10–90% rise time, and decay time constant (tau), were defined based on an average of real events as well as on previous reports for the same type of neurons (Gardner et al. 1999). The detection threshold is 2.5 to 4 times the noise SD. Statistics were performed using Excel (Microsoft, Redmond, WA) and Statview (Abacus Concepts, Berkeley, CA), and graphs were made in Igor (Wavemetrics, Lake Oswego, OR). Means ± SD are reported.

Surgery: cochlear ablation

The procedures were the same as reported previously (Mostafapour et al. 2000, 2002). Mice were anesthetized with inhaled isoflurane until areflexic. This level of anesthesia was maintained throughout the surgical procedure. An incision was made inferior to the pinna, and
the tympanic membrane was identified. Middle ear mesenchyme was aspirated, ossicles were removed, and the basal turn of the cochlea was visualized. Using a 30-gauge needle, the bony wall between the oval window and the round window was penetrated, and the contents were aspirated using a fine glass pipette. The modiolus was also destroyed. The skin incision was closed using cyanoacrylate glue. Animals were returned to their mothers and the rest of the litter within 2 h. Temporal bones were examined by light microscopy after obtaining brain slices for electrophysiology experiments or after perfusion of animals in immunohistochemistry experiments to confirm the ablation. The procedures were approved by the University of Washington Animal Use Committee.

**Immunohistochemistry**

Mice that received unilateral cochlear ablations at P7 (n = 2) and P14 (n = 2) were overdosed with sodium pentobarbital and perfused transcardially with 4% paraformaldehyde at 1 wk after surgery. After perfusion, the brains were removed and postfixed for an additional 2 h at room temperature (RT), then serially dehydrated in ethanol, cleared in methyl salicylate, and embedded in paraaffin. A one-in-five series of 10-μm coronal sections was mounted on slides. Sections were deparaffinized and rinsed in 0.01% Triton-X in PBS (PBST) for 15 min. Antigen retrieval was performed in 10 mM citric acid (pH 6.0) in a steamer for 25 min and then cooled for 10 min on ice. Slides were rinsed again in PBST for 15 min and then blocked for 1.5 h at RT in 10% normal serum, 0.1% Triton-X, and 0.5% bovine serum albumin in PBS. Sections were next incubated overnight at 4°C with the primary antibodies: mouse anti-synaptic vesicle 2 (SV2, 1:100) and rabbit anti-microtubule associated protein 2 (MAP2, 1:500, Chemicon International, Temecula, CA). The SV2 antibody was generously provided by Dr. Sandra Baijalieh and has been characterized previously (Buckley and Kelly 1985). The next day, tissue was rinsed for 30 min in PBST and incubated in Alexa Fluor 488 conjugated goat anti-mouse and Alexa Flour 594 conjugated goat anti-rabbit (1:400, Molecular Probes) for 4 h at RT. Slides were then rinsed in PBS for 2 h and coverslipped with Glycergel mounting medium (Dako, Carpinteria, CA). Controls omitting the primary antibodies were included with every trial. Photomicrographs were taken using a CoolSnap HQ Digital Camera (Photometrics, Tucson, AZ) under a ×63 objective on a Zeiss Axioplan 2 microscope. Images were captured using Slidebook 4.0 software (Intelligent Imaging Innovations, Santa Monica, CA). All photomicrograph figures were prepared using Photoshop 7.0 (Adobe Systems Incorporated, San Jose, CA).

**RESULTS**

**Electrophysiological identity of AVCN neurons**

There are two main neuronal cell types in the AVCN, bushy cells and stellate cells. They can be distinguished from each other by morphology or electrophysiology. bushy cells have short bushy-like dendrites with round cell bodies innervated by large endbulb of Held auditory terminals. Stellate cells have long branched dendrites and receive bouton-like glutamatergic synapses from auditory terminals (reviewed in Young and Oertel 2004). In electrophysiological recordings, bushy cells fire only one or a few action potentials in response to prolonged depolarizing current injections, a hallmark of auditory neurons encoding timing information (reviewed in Oertel 1999; Trussell 1999). In contrast, stellate cells fire multiple action potentials in response to prolonged depolarizing current injections. Intracellular sharp electrode recordings have shown that the electrophysiological identity of AVCN neurons is established as early as age P7 (Wu and Oertel 1987). We confirmed this observation using whole cell recordings under current-clamp configuration and further characterized the firing properties of AVCN neurons so that determination of cell type can be made in the following experiments.

Data were collected from a total of 108 AVCN neurons (38 cells were recorded by using AxoClamp 2B and 70 cells by AxoPatch 200B). Bushy cells and stellate cells were distinguished from each other by their firing patterns in response to prolonged current injection (Fig. 1). Both cell types possess a voltage sag in response to hyperpolarizing current injections, likely reflective of the presence of nonselective cation channels (I_Na). Bushy cells fired single action potentials in response to threshold current (defined here as the minimum current needed to elicit action potentials; Fig. 1A). At more depolarizing current steps, a few action potentials (1–3) were elicited and followed by a steady state potential plateau (data not shown). Stellate cells fired multiple large action potentials distributed across the duration of the current injection (Fig. 1B). To investigate the profile of firing patterns, we plotted the number of action potentials against the amplitude of the injected currents relative to the neuron’s threshold current for all individual neurons. Bushy cells showed an almost flat function (Fig. 1C, - - - -); note that several overlapped - - - are present in each graph), whereas stellate cells showed an increase in number of action potentials with increasing current amplitude, and at more depolarizing current steps, some stellate cells showed fast adaptation, especially at P7 (Fig. 1C, n = 4, 8, and 6 bushy cells for P7, P14, and P21, respectively; n = 8, 8, and 4 stellate cells for P7, P14, and P21, respectively). Another apparent difference between the two cell types is neuronal excitability. Threshold currents for bushy cells were 0.55 ± 0.17, 0.48 ± 0.16, and 0.29 ± 0.09 nA for P7, P14, and P21, respectively. Threshold currents for stellate cells were 0.11 ± 0.06, 0.10 ± 0.08, and 0.11 ± 0.10 nA for P7, P14, and P21, respectively. At the same age, stellate cells were significantly more excitable than bushy cells (Fig. 1D). Furthermore, stellate cells possessed a stable excitability over the ages tested, whereas bushy cells appeared to become more excitable with age; threshold current at P21 is significantly smaller than at P7/P14, and there is no significant difference between threshold currents between P7 and P14 (Fig. 1D).

**Development of mEPSCs in AVCN bushy cells**

Figure 2 shows development of mEPSCs of AVCN bushy cells in mice at ages P7, P14, and P21. mEPSCs were blocked by CNQX (20 μM), a specific AMPA receptor (AMPA) antagonist, indicating that mEPSCs are mediated by AMPARs (data not shown). The frequency of mEPSCs increased dramatically in bushy cells from P7 to P14/P21 (Fig. 2A). The examples were chosen to match the average frequency of mEPSCs in pooled data from each age group. Shown on the right (Fig. 2A) are three averaged traces from events of single cells normalized to their peaks. The examples were chosen to closely match the averaged 10–90% rise time and decay tau of pooled data. Figure 2, B—D, shows the histogram distributions of mEPSC amplitude (bin width of 5 pA), 10–90% rise time (bin width of 20 μs), and decay tau (bin width of 100 μs) of all mEPSCs from all cells (n = 6, 6, 9 cells for P7, P14, and P21, respectively). These histograms show the ranges and distributions of mEPSC parameters across the ages tested. mEPSCs at
P7 appeared to have broader distributions in 10–90% rise time and decay tau than those at P14/P21. Due to the large variation in mEPSC frequency in different cells, the distributions are possibly biased toward the cells with the largest number of events. Ideally, equal numbers of events from each cell should be pooled to make the histograms. However, in some cells, only a few events can be recorded, even in a prolonged period of recording, so we decided to plot all events in these histograms and then present another group of data to show the means and SDs of each parameter (Fig. 2, E–H).

We observed significant changes in the frequency and kinetics but not the amplitude of mEPSCs across the ages tested (Fig. 2, E–H). Frequency of mEPSCs increased significantly from P7 to P14 or P21 (Fig. 2E, ANOVA post hoc Fisher's test \( P < 0.05 \) and 0.01, respectively). No significant difference was detected in mEPSC frequency between P14 and P21. mEPSCs of bushy cells recorded from P21 mice show a significantly shorter 10–90% rise time than those from P7 animals. The difference in 10–90% rise time between P7 and P14 is not significant, although the trend was suggestive (\( P = 0.086 \)). Decay tau of mEPSCs gets smaller over age. A significant difference was detected by ANOVA (\( P < 0.05 \)), and post hoc Fisher's tests show that decay tau at P14 or P21 is significantly smaller than at P7, and no significant difference was detected between P14 and P21. Numeric data are shown in Table 1.

To determine whether dendritic filtering exists in bushy cells at ages tested, we plotted mEPSC 10–90% rise time against decay tau in bushy cells for the three age groups (Fig. 2, I–K). Due to the cable properties of dendrites, synaptic inputs recorded at the soma are expected to be attenuated in amplitude (Rall 1967; Rall et al. 1967). The time course of synaptic events also changes as a function of input location; the further from the soma a synaptic event occurs, the slower its time course is (Rall et al. 1967; Rinzel and Rall 1974). Therefore if dendritic filtering exists, one would expect that the synaptic events (in our case mEPSCs) with long rise time would also have long decay time. Thus a positive correlation is indicative of dendritic filtering (Gardner et al. 1999). No correlations between mEPSC 10–90% rise time and decay tau were found; linear regression analysis gave rise to correlation coefficients close to zero (\( R = -0.02, 0.09, \) and 0.05 for P7, P14, and P21, respectively). This indicates a lack of dendritic filtering in AVCN bushy cells at the ages tested, consistent with the fact that excitatory inputs from the auditory nerve impinge onto the somata and not the dendrites of bushy cells (Cant and Morest 1979; Lenn and Reese 1966).

Development of mEPSCs in AVCN stellate cells

Figure 3 shows the development of mEPSCs in AVCN stellate cells (\( n = 10, 7, 5 \) cells for P7, P14, and P21, respectively). Recordings of mEPSCs of AVCN stellate cells from mice age P7, P14, and P21 show mEPSCs of similar number and amplitude (Fig. 3A, left). Similar mEPSC kinetics were also seen across ages, as shown by the three averaged traces from events of single cells, normalized to their peaks (Fig. 3A, right). The 10–90% rise time and decay tau at P7 appeared to have a broader range than at P14 or P21. However, no significant differences were detected in averaged mEPSC frequency, amplitude, 10–90% rise time, or decay tau among different age groups (Fig. 3, E–H; Table 1; ANOVA \( P > 0.05 \)). As stated previously, all mEPSC events from all cells in...
each age group were plotted in the distribution histograms. Because the number of events in individual cells varies widely, the distributions are biased toward the cells with the largest number of events. In contrast, to generate the pooled data, we obtained a trace by averaging all events in each cell. We then measured the parameters (amplitude, rise time, and decay tau) of the averaged trace, and considered each measurement as one data point. We presented the means ± SD of the values obtained from the averaged traces (equal to the number of cells) in each age group. Therefore one should not estimate the means and SDs based on the distribution histograms. Relatively strong positive correlations between 10–90% rise time and decay tau are seen in stellate cells at all ages tested (Fig. 3, I–K; $R = 0.32, 0.24$, and $0.28$ for P7, P14, and P21, respectively), suggesting of dendritic filtering. These data are consistent with the fact that excitatory inputs from the auditory nerve impinge onto the soma, not the dendrites, of bushy cells.

FIG. 2. Development of spontaneous miniature excitatory postsynaptic currents (mEPSCs) in AVCN bushy cells. A, left: 3 original chart recordings of mEPSCs of AVCN bushy cells from mice ages P7, P14, and P21. Cells were voltage clamped at –60 mV. Strychnine (1 µM) and gabazine (10 µM) were present in the perfusion solution to block glycine and GABAA receptors. Right: averaged traces from mEPSCs of 3 individual cells, normalized to their peaks. B–D: histogram distributions of mEPSC amplitude (bin width of 5 pA), 10–90% rise time (bin width of 20 µs), and decay time constant (decay tau, bin width of 100 µs) of all events of all cells ($n = 6, 6, 9$ cells for P7, P14, and P21, respectively) at the 3 different ages. Labels indicating age at the far left in A apply to histograms in B–D. E: frequency of mEPSCs increases significantly from P7 to P14 or P21. No significant difference was detected in mEPSC frequency between P14 and P21. F: amplitude of mEPSCs shows no difference across ages tested. G: mEPSCs of bushy cells recorded from P21 mice rise significantly faster than those of P7 mice as evidenced by briefer 10–90% rise time of mEPSCs. The difference between P7 and P14 is not significant ($P = 0.086$). H: decay tau of mEPSCs gets progressively smaller from P7 to P21. Significant differences were detected by ANOVA ($P < 0.05$), and post hoc Fisher’s tests show that decay tau at P14 is significantly smaller than that at P7, and decay tau at P21 is also significantly smaller than at that P7, but no significant difference was detected between P14 and P21. Note that to generate graphs F–H, an averaged trace was obtained for each cell by averaging all detected events in that single cell. Amplitude, 10–90% rise time, and decay tau of the averaged mEPSCs were measured, then data were pooled in each age group. Means ± SD are shown in E–H. *$P < 0.05$, **$P < 0.01$ (ANOVA post hoc Fisher’s test). I–K: plots of 10–90% rise time against decay tau of mEPSCs in AVCN bushy cells at ages P7, P14, and P21. No correlations between mEPSC 10–90% rise time and decay tau were found in AVCN bushy cells at the ages tested, indicating a lack of dendritic filtering, consistent with the fact that excitatory inputs from the auditory nerve impinge onto the soma, not the dendrites, of bushy cells.
TABLE 1. Comparisons of mEPSC properties in bushy and stellate cells at different ages

<table>
<thead>
<tr>
<th></th>
<th>P7</th>
<th>P14</th>
<th>P21</th>
<th>ANOVA P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bushy cell, (N)</td>
<td>6</td>
<td>6</td>
<td>9</td>
<td>0.0190*</td>
</tr>
<tr>
<td>Frequency (Hz)</td>
<td>0.2 ± 0.3</td>
<td>7.3 ± 5.5**</td>
<td>5.7 ± 4.4*</td>
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<tr>
<td>Amplitude (pA)</td>
<td>−61 ± 28</td>
<td>−80 ± 30</td>
<td>−77 ± 25</td>
<td>0.4293</td>
</tr>
<tr>
<td>10–90% rise time (ms)</td>
<td>0.18 ± 0.08</td>
<td>0.13 ± 0.01</td>
<td>0.12 ± 0.01*</td>
<td>0.0684</td>
</tr>
<tr>
<td>Decay tau (ms)</td>
<td>0.47 ± 0.12</td>
<td>0.26 ± 0.13**</td>
<td>0.19 ± 0.05**</td>
<td>0.0002**</td>
</tr>
<tr>
<td>Stellate cell (N)</td>
<td>10</td>
<td>7</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Frequency (Hz)</td>
<td>0.8 ± 1.3</td>
<td>1.7 ± 2.6</td>
<td>1.7 ± 2.0</td>
<td>0.5864</td>
</tr>
<tr>
<td>Amplitude (pA)</td>
<td>−58 ± 16 (n = 9)</td>
<td>−52 ± 18</td>
<td>−53 ± 17</td>
<td>0.7868</td>
</tr>
<tr>
<td>10–90% rise time (ms)</td>
<td>0.18 ± 0.07 (n = 9)</td>
<td>0.18 ± 0.05</td>
<td>0.20 ± 0.10</td>
<td>0.8473</td>
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<tr>
<td>Decay tau (ms)</td>
<td>0.48 ± 0.34 (n = 9)</td>
<td>0.46 ± 0.22</td>
<td>0.53 ± 0.37</td>
<td>0.9127</td>
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</tbody>
</table>

Means ± SD are shown. n: number of cells; *P < 0.05, **P < 0.01, ANOVA, and post hoc Fisher’s tests (between P14 and P7 or P21 and P7); no significant difference was detected in any of the parameters between P14 and P21 in both cell types at any age.

FIG. 3. Development of mEPSCs in AVCN stellate cells (n = 10, 7, 5 cells for P7, P14, and P21, respectively). A–D: for figure legends, see Fig. 2. E–H: no significant differences are detected in mEPSC frequency, amplitude, 10–90% rise time, or decay tau among different age groups (ANOVA P > 0.05). I–K: plots of 10–90% rise time against decay tau of mEPSCs in AVCN stellate cells at ages P7, P14, and P21. Positive correlations between mEPSC 10–90% rise time and decay tau are seen in stellate cells at the ages tested. This is suggestive of dendritic filtering in these neurons, consistent with the fact that excitatory inputs from the auditory nerve impinge onto their dendrites.

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Effects of cochlea ablation performed at P7 on development of mEPSCs (recorded at P14)

We performed cochlea ablations at P7 and examined mEPSCs at P14, when AVCN bushy cell death had reached its maximum (Mostafapour et al. 2000). Physiological identity of the two main neuron types is preserved after cochlea ablation (data not shown), consistent with previous reports on rats (Francis and Manis 2000). Figure 4 shows mEPSC recordings obtained from surviving bushy cells (n = 7 cells). Figure 4A shows examples of an original chart recording of mEPSCs and an averaged trace (normalized to the peak of the trace for P14 bushy cells in Fig. 2). CNQX (20 μM) completely blocked mEPSCs in surviving bushy cells, indicative of unaltered pharmacology of mEPSCs (data not shown). Histogram distributions of mEPSC amplitude (bin width of 5 pA), 10–90% rise time (bin width of 20 μs), and decay tau (bin width of 100 μs) show similar patterns to the bushy cells of the same age (P14) in Fig. 2. A slightly broader range of mEPSC amplitude is seen for activity-deprived bushy cells (Fig. 4B). Unpaired t-test detected no difference in mEPSC frequency, amplitude, 10–90% rise time, and decay tau between control cells at P14 (same data as in Fig. 2, E–H) and activity-deprived cells at the same age (Fig. 4, E–H). mEPSCs of the experimental group appeared to have larger variability (SD bars) in frequency, amplitude, 10–90% rise time, and decay tau.

A major difference between mEPSCs recorded from activity-deprived bushy cells and normal cells is the relationship between rising time and decay tau. A fairly strong positive correlation (Fig. 4I, R = 0.31) between mEPSC 10–90% rise time and decay tau is seen in activity-deprived bushy cells as compared with normal P14 bushy cells (Fig. 2I, R = 0.09; z test, P < 0.001). This observation suggests dendritic filtering in surviving bushy cells, probably resulting from added synaptic inputs on their dendrites. To further examine effects of cochlea ablation on the expression patterns of synaptic contacts, we examined the distribution pattern of synaptic vesicle protein 2 (SV2), a transmembrane protein that is commonly present at synaptic vesicles (Buckley and Kelly 1985). The tissue was double labeled with microtubule-associated protein 2 (MAP2), one of the most abundant proteins in neurons that is highly expressed in dendrites. Unilateral cochlea ablation was performed at P7 and animals were allowed to survive for 7 days. The contralateral side of each slice was used as a within-slice control. We observed a ring of SV2 labeling around the somata in the contralateral bushy cells (Fig. 5A) that was mostly absent on the ipsilateral side (Fig. 5B). However, the overall density of SV2 labeling in the neuropil appeared qualitatively similar between sides.

In contrast to bushy cells, cochlea ablation resulted in no significant changes in mEPSCs of stellate cells (Fig. 6, n = 9 cells). Figure 6A shows examples of an original chart recording of mEPSCs and an averaged trace (normalized to the peak of the trace for P14 stellate cells in Fig. 3). Histogram distributions of mEPSC amplitude, 10–90% rise time, and decay tau show similar patterns to the control cells of the same age (P14) in Fig. 3 (Fig. 6, B–D). Unpaired t-test detected no difference in mEPSC frequency, amplitude, 10–90% rise time, and decay tau between control cells at P14 (same data as in Fig. 3, E–H) and activity-deprived cells at the same age (Fig. 6, E–H, P > 0.05). mEPSCs of the experimental group appeared to have larger variability (SD bars) in frequency, amplitude, 10–90% rise time, and decay tau. Furthermore, the correlation coefficient (R = 0.30) between mEPSC 10–90% rise time and decay tau is comparable to that of stellate cells at P14 in normal animals (Fig. 6D; z test, P > 0.05).

Effects of cochlea ablation performed at P14 on development of mEPSCs (recorded at P21)

Synaptic inputs may be less susceptible to activity modulation after the critical period. To test this hypothesis, we examined the development of mEPSCs if cochlea ablation is performed after the critical period, we performed cochlea...
ablation at P14, and recorded mEPSCs in AVCN neurons at P21. We observed similar patterns in histogram distribution of mEPSC amplitude, 10–90% rise time, and decay tau in activity-deprived bushy cells to those of control cells at the same age (P21; Fig. 7). Unpaired t-test detected no difference in mEPSC frequency, amplitude, 10–90% rise time, and decay tau between control cells at P21 (same data as in Fig. 2, E–H) and activity-deprived cells at the same age (Fig. 7, E–H). A stronger positive correlation (R = 0.14) between mEPSC 10–90% rise time and decay tau is seen in activity-deprived bushy cells.
bushy cells than control ($R = 0.05$; Fig. 7I; $z$ test $P < 0.01$). A ring of SV2 labeling around the somata of bushy cells was seen in both the contralateral and the ipsilateral AVCN, but the SV2 labeling on the ipsilateral somas appeared less continuous than on the contralateral side (Fig. 8). Similarly, cochlea ablation resulted in no changes in frequency, amplitude, and time courses of mEPSCs of stellate cells (Fig. 9, $n = 6$ cells). The histograms (Fig. 9, B–D) show similar patterns to those of the stellate cells of the same age (P21) in normal animals (Fig. 3). Unpaired $t$-test detected no difference in mEPSC frequency, amplitude, 10–90% rise time, and decay tau between control cells at P21 (same data as in Fig. 3, E–H) and activity-deprived cells at the same age (Fig. 9, E–H). A significant increase in correlation coefficient ($R = 0.41$, compared with $R = 0.28$ for P21 stellate cells under control conditions) between mEPSC 10–90% rise time and decay tau was seen (Fig. 9I; $z$ test, $P < 0.01$). MAP2 and SV2 staining in stellate cells is not shown due to difficulties in identifying stellate cells in the stained slices.

**DISCUSSION**

We investigated the development of spontaneous miniature excitatory synaptic currents (mEPSCs) in AVCN neurons in mice at ages P7, P14, and P21. We showed that the frequency of mEPSCs in bushy cells increases significantly with little changes in amplitude and that mEPSCs become progressively briefer. In contrast, stellate cells lack significant changes in mEPSCs during these ages. We also examined the effects of unilateral cochlea ablation on development of mEPSCs. Cochlea ablation performed at P7 and P14 did not alter mEPSC frequency, amplitude or kinetics but did lead to the emergence of dendritic filtering in bushy cells. Stellate cells also show no changes in mEPSCs compared with age-matched controls except for an increase in dendritic filtering when the surgery was performed at P14.

**Development of mEPSCs in AVCN neurons and correlation to their morphology**

Synapses at AVCN neurons are present at birth. During the first weeks in postnatal development, auditory nerve endings impinging onto bushy cells in mouse AVCN show substantial changes in morphology (Limb and Ryugo 2000). At P7, auditory nerve inputs to AVCN bushy cells are bouton like and lack any branches at the terminals. At P14, the nerve endings increase substantially in size (10- to 15-fold larger than P7).

Most nerve endings are cup-shaped at this age, but some of them have branches at the terminals and can be identified as nascent endbulbs. Assuming that an increase in size of nerve endings is associated with an increase in number of synapses and thus more release sites, the dramatic change in nerve
endings size could well account for the sharp increase identified here in mEPSC frequency in bushy cells from P7 to P14 (35-fold) because the frequency of spontaneous EPSCs seems to be correlated with the number of auditory inputs (Gardner et al. 1999). Four weeks after birth (P28), the auditory nerve endings on AVCN bushy cells form several branches that are similar in caliber to their parent branch and can be clearly identified as nascent endbulbs. However, the overall coverage of the cell bodies by the nerve terminals appears similar to that of P14 mice (Limb and Ryugo 2000). This may explain why we observed that the frequency of mEPSCs in bushy cells does not further increase from P14 to P21. The increased mEPSC frequency, especially from P7 to P14, indicates the maturation of some presynaptic elements such as number of synapses, activity level of individual release sites or a combination of both. Future anatomical measures are needed to further determine which is the case.

Along with morphological maturation of presynaptic terminals and postsynaptic neurons (Limb and Ryugo 2000), AMPAR-mediated mEPSCs at the endbulb of Held synapses in mouse AVCN bushy cells become significantly faster in their time courses across the critical period (Fig. 2). This is consistent with previous reports on development of endbulb of Held synaptic physiology in the cochlear nucleus in rats (Bellingham et al. 1998) or chickens (Brenowitz and Trussell 2001). The faster time course of mEPSCs parallels, and may account in part for, the development of faster evoked EPSCs (eEPSCs) in rat AVCN bushy cells (Bellingham et al. 1998) or chicken nucleus magnocellularis neurons (Brenowitz and Trussell 2001). The decay time constants in P21 bushy cells are half as large those reported in the Gardener et al. study (1999). This discrepancy may be due to the following reasons. The mouse strain is different; we used C57BL/6, whereas Gardener et al. (1999) used CBA and ICR. The recording temperature we used (34–36°C) is slightly higher than the temperature they used (34°C). Synaptic events are expected to have faster time course at a higher temperature within the physiological range. The age of animals may be a factor, too. We used P21 mice, and Gardener et al. (1999) used mice age between P18 and 25. The age difference may also account for the observation that the SD of the decay tau we obtained (0.05 ms) is much smaller than that Gardener et al. (1999) obtained (0.15 ms). Interestingly, in aging DBA mice, mIPSCs in AVCN bushy cells showed reduced frequency and a slower time course compared with young animals (Wang and Manis 2005), precisely the opposite to what was observed in developing animals.

Research on the morphological development of auditory nerve endings impinging on AVCN stellate cells at the ages we studied is lacking. Based on our electrophysiological observations, auditory nerve endings onto stellate cells are not expected to show substantial morphological changes from the first to the third week after birth. This may indicate that maturation of the excitatory inputs to stellate cells occurs either prior to P7 or after the third postnatal week. Morphological studies are needed to test these alternative hypotheses.

mEPSCs in surviving bushy cells after cochlea ablation

Cochlea ablation results in degeneration of the auditory nerve terminals, but the overall strength of synaptic inputs onto surviving cochlear nucleus neurons seems to be preserved, likely due to reinnervation of cochlear nucleus neurons by fibers from sources other than the ipsilateral auditory nerve. In rats, when the auditory nerve is transected, auditory nerve terminals start degenerating within 24 h after the surgery, and by 5–9 days, most degenerating terminals disappear. What is interesting is that electron microscopy confirms reinnervation of AVCN neurons during this time probably due to reoccupation of nearby intact terminals containing transmitter vesicles (Gentschev and Sotelo 1973) or an increase in axonal sprouting from medial olivocochlear neurons (Illing et al. 2005). In cats, cochlea ablation results in degeneration of endbulbs of Held, the auditory nerve endings on AVCN bushy cells, but leaves other types of synapses intact (Cant and Morest 1979). In addition, hearing deprivation in mice leads to a significant increase in nonauditory nerve endings impinging on AVCN neurons and a decrease in auditory nerve endings, but the overall cell membrane covered by the auditory plus the nonauditory nerve endings is unchanged (Trune and Morgan 1988). Although these structural changes have been known for some time, physiological studies are lacking to determine if there are functional changes in synaptic transmission after manipulations of the auditory afferent inputs.

Assuming a similar time course of degeneration of the auditory nerve terminals after cochlea destruction among the rat, cat, and the mouse, presumably most or all auditory terminals at the synapses in the AVCN would disappear in ~1 week. This would lead one to predict much weaker spontaneous synaptic activity in activity-deprived neurons compared with neurons in normal hearing animals of the same age. However, one may not observe a significant difference in spontaneous synaptic activity in activity-deprived neurons if there is a compensatory reaction via reinnervation or regeneration of synapses from other sources including the AVCN neuron itself (auto-synapses). If reinnervation and/or regeneration of inputs occur, one prediction is that mEPSCs of similar properties to normal neurons should be recorded in the surviving neurons 1 or 2 wk after cochlea ablation at which time maximal cell death (30–70% of the total) in AVCN has occurred (Mostafapour et al. 2000).

Our results support the occurrence of reinnervation and/or regeneration. Seven days after cochlea ablation performed either at P7 or P14, mEPSCs in AVCN bushy cells are maintained at a similar level in terms of frequency, amplitude, and time course to that of AVCN neurons of normal animals of the same age. Seven days after cochlea ablation performed at P7, a positive correlation between mEPSC 10–90% rise time and decay tau was detected, indicative of synaptic inputs onto dendrites and emergence of dendritic filtering in surviving bushy neurons. A positive correlation was also seen when the surgery was performed at P14, suggesting that at least some events occurred at locations other than the soma. Assuming that the frequency of mEPSCs at single synapses is not changed by cochlea ablation, we speculate the new synapses formed and impinged on dendrites of surviving bushy cells. These physiological observations were supported by our observations of the expression patterns of SV2. Cochlea ablation led to a reduction of SV2 expression around the somas, especially when the surgery was performed at P7 (Fig. 5), whereas the overall density of SV2 expression in the neuropil appeared to be similar to the control. Future anatomical measures using electronic microscopy are needed to determine the
exact location of synaptic vesicles. In addition, the origin of the newly added synaptic inputs that impinge onto bushy cells as a complimentary reaction to cochlea ablation remains largely unknown. Some previous studies suggest that the contralateral bushy cells are one of the sources (Jackson and Parks 1988; Parks et al. 1990; Sumner et al. 2005). It would be also interesting to examine the time course for the effects of cochlea ablation on mEPSCs in AVCN bushy cells. Furthermore, because bushy cells also receive nonglutamate receptor-mediated synaptic inputs, it is unclear whether changes, if any, in nonglutamate synaptic vesicles after cochlea ablation contribute to this observation. Such changes have been seen in the motor system, for example, in developing mammalian motor neurons, inhibitory synaptic inputs mediated by GABA and glycine receptors play important roles in neuronal survival (Banks et al. 2005) and in axotomized motor neurons, GABAergic inputs change their sign from inhibitory to excitatory (Nabekura et al. 2002).

mEPSCs and the critical period

Various approaches have been taken to study the mechanisms underlying the critical period for mouse AVCN neurons. For example, bcl-2, a gene that functions as an anti-apoptotic factor and promotes cell survival, has been shown to modulate cell death in the mouse cochlear nucleus. Deletion of the gene renders mature animals (≥P27) susceptible to afferent deprivation, whereas overexpression of the gene protects AVCN neurons in young (P5/P8) animals (Mostafapour et al. 2000, 2002). Using gene microarray analysis, Harris et al. (2005) found that during and after the critical period, AVCN cells in mice differentially express genes shown to be involved in neuronal survival (Banks et al. 2005) and in axotomized motor neurons, GABAergic inputs change their sign from inhibitory to excitatory (Nabekura et al. 2002).

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