A Developmental Shift from GABAergic to Glycinergic Transmission in the Central Auditory System

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GABAergic and glycinergic circuits are found throughout the auditory brainstem, and it is generally assumed that transmitter phenotype is established early in development. The present study documents a profound transition from GABAergic to glycinergic transmission in the gerbil lateral superior olive (LSO) during the first 2 postnatal weeks. Whole-cell voltage-clamp recordings were obtained from LSO neurons in a brain slice preparation, and IPSCs were evoked by electrical stimulation of the medial nucleus of the trapezoid body (MNTB), a known glycinergic projection in adult animals. GABAergic and glycinergic components were identified by blocking transmission with bicuculline and strychnine (SN), respectively. In the medial limb of LSO, there was a dramatic change in the GABAergic IPSC component, decreasing from 78% at postnatal day 3 (P3)–P5 to 12% at P12–P16. There was an equal and opposite increase in the glycinergic component during this same period. Direct application of GABA also elicited significantly larger amplitude and longer duration responses in P3–P5 neurons compared with glycine-evoked responses. In contrast, MNTB-evoked IPSCs in lateral limb neurons were more sensitive to SN throughout development. Consistent with the electrophysiological observations, there was a reduction in staining for the β2,3-GABAA receptor subunit from P4 to P14, whereas staining for the glycine receptor-associated protein gephyrin increased. Brief exposure to baclofen depressed transmission at excitatory and inhibitory synapses for ~15 min, suggesting a GABAA-mediated metabotropic signal. Collectively, these data demonstrate a striking switch from GABAergic to glycinergic transmission during postnatal development. Although GABA and glycine elicit similar postsynaptic ionotropic responses, our results raise the possibility that GABAergic transmission in neonates may play a developmental role distinct from that of glycine.

Key words: GABAA; glycine; inhibition; GABAB; development; gerbil; lateral superior olive

In contrast to the literature on the developmental plasticity of excitatory synapses, little is known about activity-dependent mechanisms at inhibitory terminals. In the adult nervous system, GABA and glycine are each known to hyperpolarize the postsynaptic membrane by gating chloride channels (Bormann et al., 1987). However, during early development, GABA or glycine often produces a membrane depolarization that is accompanied by an influx of calcium (Connor et al., 1987; Ben-Ari et al., 1989; Ito and Cherubini, 1991; Obrist and Van den Pol, 1995; Lo et al., 1998). In the developing lateral superior olive (LSO), a binaural nucleus in the ventral auditory brain stem, inhibitory synapses can depolarize cells directly (Kandler and Friauf, 1995) or elicit a hyperpolarization followed by a rebound depolarization (Sanes, 1993). Thus, the transient depolarizing influence of inhibitory terminals could allow them to use calcium-dependent mechanisms to play a role in the stabilization or elimination of excitatory terminals (Connold et al., 1986; Cash et al., 1996).

Inhibitory afferents to the LSO project from the medial nucleus of the trapezoid body (MNTB), a glycinergic nucleus that is activated by sound to the contralateral ear (Guinan et al., 1972a,b; Moore and Caspary, 1983; Spangler et al., 1985; Wenthold et al., 1987, 1990; Zook and DiCaprio, 1988; Sanes and Siverls, 1991; Schwartz, 1992). Anatomically, LSO is innervated by terminals that contain flattened or pleomorphic vesicles that stain positively with both GABA and glycine antibodies (Helfert et al., 1989, 1992). In adult animals, LSO neurons encode interaural level differences by integrating excitatory potentials driven by the ipsilateral ear and inhibitory potentials driven by the contralateral ear (Boudreau and Tsuchitani, 1970; Caird and Klinke, 1983; Harnischfeger et al., 1985; Sanes and Rubel, 1988).

The inhibitory terminals from MNTB appear to be dynamic during early development, displaying a physical reduction in their arbor size and a reduction in the number of functional inhibitory afferents per postsynaptic neuron (Sanes and Siverls, 1991; Sanes, 1993). Furthermore, when inhibitory transmission is disrupted during development, both the morphology and physiology of LSO neurons are affected (Sanes et al., 1992; Sanes and Takács, 1993; Aponte et al., 1996; Kotak and Sanes, 1996; Sanes and Hafidi, 1996). For example, denervation of the glycinergic afferents to LSO or strychnine rearing result in the upregulation of functional NMDA receptors, thus enhancing excitatory transmission (Kotak and Sanes, 1996). Therefore, we are interested in inhibitory synaptic mechanisms that might contribute to presynaptic and postsynaptic maturation. In the present report, we present physiological and anatomical evidence for a major transition from GABAergic to glycinergic inhibitory transmission during early development and provide evidence that early GABAergic transmission may activate metabotropic receptors.
MATERIALS AND METHODS

Brain slice physiology. Postnatal gerbils (Meriones unguiculatus) at postnatal day 3 (P3)–P16 were used to generate brain slices through the LSO, MNTB, and ipsilateral cochlear nucleus afferents (Sanes, 1993). Transverse vibratome sections of 300 μm were cut in cold (−8°C) oxygenated artificial CSF (ACSF), preincubated at room temperature for 2 hr in a holding chamber, and transferred to the recording chamber where ACSF was superfused at 7 ml/min at room temperature. The ACSF contained (in mM): 130 potassium gluconate, 0.6 EGTA, 10 HEPES, 2 MgCl₂, 5 KCl, 2 ATP, and 0.3 GTP, pH 7.2. Whole-cell voltage-clamp recordings were obtained with pipettes containing (in mM): 127.5 cesium borosilicate glass microcapillaries, and they had a resistance of 4–6 MΩ. Recordings were fabricated from 1.5-mm-inner diameter boroscopic glass microcapillaries, and they had a resistance of 4–6 MΩ.

For current-clamp recordings, the composition of the internal pipette solution was (in mM): 130 potassium glutamate, 0.6 EGTA, 10 HEPES, 2 MgCl₂, 5 KCl, 2 ATP, and 0.3 GTP, pH 7.2. Whole-cell voltage-clamp recordings were obtained with pipettes containing (in mM): 127.5 cesium glutamate, 0.6 EGTA, 10 HEPES, 2 MgCl₂, 5 KCl, 2 ATP, 0.3 GTP, and 5 QX-314, pH 7.2. The resting potential was checked immediately after breaking the cell membrane, and neurons with a resting potential of −40 mV or better were used in this analysis. Custom-designed software running on a 486 personal computer platform was used for programmed stimulus delivery, data acquisition, and analysis (Sanes, 1993). Bipolar stimulating electrodes were placed on the MNTB and the afferent pathway from the ipsilateral cochlear nucleus at the lateral edge of LSO. Incremental voltage pulses (200 μsec) were delivered to each set of afferents at 0.5 Hz, and the maximum amplitude IPSC was determined. The data were divided into three age ranges: P3–P5, P8–P11, and P12–P16.

The IPSCs were recorded at holding potentials of −20–0 mV in the presence of 6-cyano-7-nitroquinazoline-2,3-dione (CNQX; 20 μM) and d(-)2-amino-5-phosphonopanoic acid (AP-5; 50 μM; h = 35) (Research Biochemicals, Natick, MA), or kynurenic acid (KYN; 5 mM; h = 36). The glycineergic IPSC component was blocked with strychnine (SN; 2 μM; Sigma, St. Louis, MO), and the GABAergic IPSC component was blocked with bicuculline (BIC; 10 μM; Sigma). For GABA (5 mM; 15 sec) or glycine (5 mM; 15 sec) exposure, slices were first incubated in KYN. To avoid long-lasting alterations in intracellular environment, only a single agonist dose was tested per slice. Voltage-clamp (n = 4) and current-clamp (n = 4) recordings were also performed to examine the effects of a GABA_A antagonist, baclofen (50 μM), on synaptic function and membrane properties. The data include whole-cell recordings from 101 LSO neurons from 87 brain slices.

Immunocytochemistry. Gerbils at P4 (n = 3) and P14 (n = 3) were anesthetized with sodium pentobarbital and perfused transcardially with cold saline nitrite solution (0.9% sodium chloride and 0.1% sodium nitrite) and 4% paraformaldehyde and 0.1% glutaraldehyde in 0.12M PBS, pH 7.4. After 30–60 min on ice, brains were removed and immersed overnight in cold fixative. Brains were then washed in PBS and transferred to 30% sucrose in buffer overnight. The brains were then placed in OCT mounting compound (Miles, Elkhart, IN) and frozen in ethanol over dry ice. Serial cryostat sections (30 μm) of the brainstem cut in the coronal plane were used for immunohistochemical staining with monoclonal antibodies directed against the glycinergic IPSC component was blocked with strychnine (SN; 2 μM; Sigma, St. Louis, MO), and the GABAergic IPSC component was blocked with bicuculline (BIC; 10 μM; Sigma). For GABA (5 mM; 15 sec) or glycine (5 mM; 15 sec) exposure, slices were first incubated in KYN. To avoid long-lasting alterations in intracellular environment, only a single agonist dose was tested per slice. Voltage-clamp (n = 4) and current-clamp (n = 4) recordings were also performed to examine the effects of a GABA_A antagonist, baclofen (50 μM), on synaptic function and membrane properties. The data include whole-cell recordings from 101 LSO neurons from 87 brain slices.

RESULTS

Several anatomical findings indicate that the medial (high-frequency) and lateral (low-frequency) limbs of the gerbil LSO differ from one another, including the density of glycine receptors and MNTB afferents (Sanes et al., 1987; Sanes and Siverls, 1991). Therefore, we have analyzed the electrophysiological data obtained from medial and lateral limb neurons separately.

Inhibitory currents in the medial limb

Ipsilaterally and MNTB-evoked IPSCs were recorded in 44 of 45 medial limb neurons. Figure 1A shows a P4 neuron in which 10 μM BIC reduced the major IPSC component, whereas the remainder was almost eliminated by 2 μM SN. An identical result was obtained when the sequence of antagonist exposure was reversed in a second neuron (Fig. 1B). In contrast, BIC marginally decreased IPSC amplitude in a P14 neuron, whereas SN...
eliminated the remaining component (Fig. 1C). Once again, a similar trend was observed when SN was used first, followed by BIC in a separate neuron (Fig. 1D). In P4 (n = 2) and P14 (n = 2) neurons, reversal of the BIC effect was followed by SN application in the same neuron, producing complementary reduction of the IPSCs, which is consistent with the recordings shown in Figure 1. A summary of MNTB-evoked IPSC amplitudes is shown in Figure 2. The decreasing contribution of GABAergic transmission with age is evident when the data are plotted as the amplitude of pharmacologically isolated IPSCs (Fig. 2A) or as a percent of the total IPSC amplitude (Fig. 2B). The total calculated conductance for MNTB-evoked IPSCs did not vary significantly with age (5 ± 1.1 nS at P3–P5, 7.2 ± 1.1 nS at P8–P11, and 4.9 ± 2.1 nS at P12–P16; mean ± SEM; ANOVA, p > 0.3, df = 33). In some neurons, a small amplitude current persisted after exposure to both antagonists (Fig. 1A). This small remaining synaptic current (not blocked by BIC and SN) generally reversed at a holding potential more negative than −30 mV (n = 3). The percentage reductions in ipsilaterally evoked IPSCs by BIC or SN were similar to those observed for MNTB stimulation at all ages (data not shown).

An important question that arises is why we failed to observe significant GABAergic inhibition in previous current-clamp studies (Sanes and Hafidi, 1996; Kotak and Sanes, 1996). To assess possible “masking” of GABAergic synaptic potentials, we performed additional current-clamp recordings while sequentially applying pharmacological agents (n = 8, 4 each at P4 and P10). As shown for two different neurons in Figure 3, SN application eliminated the IPSPs but also unmasked contralateral EPSPs (Fig. 3A) or mixed responses (Fig. 3B). Addition of KYN (5 mM) abolished any excitatory potentials and revealed SN-insensitive IPSPs. These IPSPs were then blocked by BIC.
To assess the relative postsynaptic sensitivity to GABA or glycine, P3–P5 medial limb neurons were exposed to either of the two transmitters (5 mM; 15 sec), and the holding current was monitored at a holding potential of −90 mV. For this experiment, we restricted analyses to the early ages when GABAergic transmission predominated (P3–P5). Figure 4A shows representative currents obtained from each agonist and illustrates the greater sensitivity to GABA than glycine in P3–P5 medial limb neurons. Although BIC eliminated IPSCs elicited by either agonist, GABA application produced a significantly larger-amplitude and longer-lasting current (−3 min). Recordings were made in the presence of KYN (5 mM) at a holding potential of −90 mV.

A summary of the data (Fig. 4B) indicates that GABA induced larger-amplitude and longer-duration currents than those elicited by glycine. In two P4 lateral limb neurons that did not display IPSCs, GABA application also produced sizable responses (321 and 340 pA, 110 and 140 sec).

In voltage-clamp recordings, the reversal potential of MNTB-evoked IPSCs were relatively depolarized in P3–P5 neurons and gradually shifted close to the calculated EC_{50} with increasing postsynaptic age (Fig. 5). An identical observation was made for ipsilaterally evoked IPSCs (data not shown). When E_{IPSC} was compared in neurons from P7 animals before and after BIC application, it remained unchanged (control E_{IPSC} = −41 ± 1.7 mV; E_{IPSC} after BIC treatment = −40 ± 1.9 mV, mean ± SEM; n = 3). However, it should be noted that hyperpolarizing IPSPs were observed in current-clamp recordings, even at P3–P5.

**Inhibitory currents in the lateral limb**

MNTB or ipsilateral stimulation elicited IPSCs in 22 of 26 LSO lateral limb neurons, although all neurons displayed bilateral excitatory synaptic currents. Although BIC decreased IPSC amplitude in P3–P5 and P12–P16 neurons (ipsilateral, 20 of 26; MNTB, 22 of 26 neurons), the addition of SN nearly eliminated the major IPSC component at both ages (Fig. 6A). In three P4 animals, recordings were obtained from medial and lateral limb neurons in the same slice. Although BIC eliminated most of the IPSC in the medial limb neuron, it only blocked a minor fraction of the IPSC recorded in the lateral limb. Figure 6B presents a summary of the GABAergic and glycineergic synaptic components recorded in lateral limb neurons and illustrates that sensitivity to BIC and SN did not change with age. BIC reduced the MNTB-evoked IPSCs by 24 ± 6% at P3–P5 and 15 ± 5% at P12–P16 (t = −0.9; df = 21; p > 0.3). A similar trend was observed for ipsilaterally evoked IPSCs (data not shown). Lateral limb neurons also differed from medial limb neurons in that they did not exhibit depolarized IPSC reversal potentials in young animals. At P3–P5, MNTB-evoked E_{IPSC} was −48 ± 3.1 mV for lateral limb neurons compared with −31 ± 2.2 mV for medial limb neurons (t = 4.5; df = 31; p < 0.0001).
Effect of baclofen
Because GABAergic transmission appeared to be prominent in neonatal animals, we examined whether a GABA B mechanism was present in P5–P6 medial limb neurons. Baclofen elicited a reversible (10–15 min) decrease in ipsilaterally evoked EPSP amplitude (~80%) and in MNTB-evoked IPSP amplitude (~50%). In two neurons there was a 20% decrease in input resistance, although no change was noted in membrane potential, input resistance, or spike threshold in the other two neurons (Fig. 7). In two voltage-clamp recordings, a similar baclofen-evoked reduction in EPSCs and IPSCs was observed.

Immunocytochemical localization of GABA A and glycine receptors
At P4, the LSO was stained relatively intensely with GABA A receptor antibody and appeared darker compared with adjacent areas (Fig. 8A). There was no apparent difference in the staining pattern between the medial and lateral limbs. At higher magnification, the neurons showed punctate staining on the somatic periphery and darkly stained neuropil, although a few cells had patchy staining throughout the soma (Fig. 9A). At P14, there was a considerable reduction in the intensity of staining within the LSO (Fig. 8C) compared with the staining at P4. At higher magnification, there was a clear reduction in the neuropil staining, although a few cells with intense staining were observed occasionally (Fig. 9C).

In contrast, gephyrin staining at P4 was much less intense in the medial limb of LSO (Figs. 8B, 9B) compared with GABA A receptor staining. At P14, there was a dramatic increase in the intensity of staining in the medial limb (Figs. 8D, 9D) compared with P4. Although the intensity of somatic staining remained at the same level in both medial and lateral limbs, neuropil of the medial limb was stained darker compared with that of the lateral limb. This feature made the medial limb appear more darkly stained than the lateral limb at P14 (Fig. 8D). A large number of positively stained cells were seen in the lateral limb of LSO at P4, and a few faintly stained cells could be identified in the medial limb. From the morphology they did not appear to be the principal cells (Fig. 9B). Much of this somatic gephyrin staining may be located intracellularly. A striking feature was the gephyrin-positive axons running within the nucleus (Fig. 9B) that were not apparent in P14 animals.

DISCUSSION
The major finding in this study is a switch from GABAergic to glycineergic transmission in the medial limb of the gerbil LSO during the first 2 postnatal weeks. This change could arise from an alteration of the MNTB neurotransmitter and/or the LSO receptor. Alternatively, transient GABAergic projections to LSO may be activated by stimuli delivered to the MNTB region (e.g., axons of passage) and the ipsilateral pathway. Below, we argue in favor of a transition of existing projections.

Development of glycineergic transmission in the medial limb
There is a striking developmental transformation from GABAergic to glycineergic transmission in the medial limb of the LSO (Figs. 1, 2). This conclusion is based on the fact that ~80% of ipsilaterally and MNTB-evoked IPSCs were blocked by BIC in
subunit was intense throughout the LSO at P4 but declined by P14. A similar finding was reported previously in the rat (Fritschy et al., 1994). In contrast, gephyrin staining was low at P4, particularly in the LSO medial limb, but increased by P14. These results are consistent with a recent report that the α1 glycine receptor subunit appears gradually in the rat LSO during development (Friauf et al., 1997).

Although BIC and SN are well established blockers of GABA_{A} and glycine receptors, respectively, a certain degree of cross-reactivity is possible. Trombley and Shepherd (1994) have shown that 30 μM SN can antagonize GABA-evoked currents in the olfactory bulb, although 3–10 μM BIC had no effect on glycine-mediated currents. It is unlikely that substantial cross-reactivity contributed to the present results because (1) there is a complementary decrease of IPSC amplitude when BIC or SN is applied separately to the same neuron; that is, the percent IPSC amplitude remaining after BIC exposure (glycineric component) is equal to the percent IPSC amplitude antagonized by SN exposure; (2) sensitivity of the two LSO limbs to BIC or SN differs in the same slice; and (3) the concentrations of SN (2 μM) and BIC (10 μM) were within the limits that exhibit selective antagonism (Dichter, 1980; Trombley and Shepherd, 1994).

Early development of GABA-containing neurons has been found throughout the nervous system (Lauder et al., 1986), and transient expression of GABA in the spinal cord has been particularly well studied (Obata et al., 1978; Reitzel et al., 1979; Maderdrut et al., 1986; Ma et al., 1992; Mitchell and Redburn, 1996). For example, in the chick spinal cord, there is a decrease in GABA-positive neurons and a complementary increase in glycine-positive neurons during embryonic development (Berki et al., 1995), and it was suggested that neurons may change transmitter phenotype. Recent immunohistochemical staining for GABA in the MNTB and LSO of neonatal ferrets (Henkel and Brunso-Bechtold, 1998) and gerbils (S. Korada and I. R. Schwartz, unpublished observations) suggests that it is expressed in neonates.

If MNTB afferents change transmitter, then this may reflect a process of differentiation such as has been described in the sympathetic nervous system. The adrenergic or cholinergic identity of sympathetic terminals is regulated by the local environment, electrical activity, and the postsynaptic target (Walicke et al., 1977; Habecker and Landis, 1994; Guidry and Landis, 1995; Reissman et al., 1996). Furthermore, the co-release of neurotransmitters may be more common than suspected although difficult to verify when many afferent pathways are present (Johnson, 1994).

There are several reasons why we failed to detect IPSPs in current clamp after SN exposure in previous studies. At times, a very small postsynaptic potential did remain (Sanes and Hafidi, 1996), but we did not consider this to be a significant synaptic component. It is possible that a poor space clamp obscured these small remaining PSpS, perhaps because GABAergic IPSPs originate at distal dendrites. In other cases (Fig. 3, left), SN application revealed an underlying excitatory current that obscured the GABAergic IPSC. In the present study, the pipette solution used for voltage-clamp studies (e.g., Cs and QX-314) and the presence of ionotropic glutamate receptor antagonists, permitted full analysis of inhibitory synaptic events. In this regard it is interesting that the staining pattern for GABA_{A} receptors indicates that they are primarily localized in the neuropile at P4 (Fig. 9A) and do not surround the LSO somata as is found for gephyrin at P14 (Fig. 9D). Gephyrin appears to induce clustering of GABA_{A} receptor

![Figure 7](image)

**Figure 7.** Baclofen causes a long-lasting depression of synaptic transmission in the LSO. **A.** In current-clamp recording from a P6 medial limb neuron, subthreshold ipsilateral stimulation (IPSI) and MNTB stimulation elicited sizable EPSPs and IPSPs, respectively (top left). Responses of this neuron to a depolarizing and hyperpolarizing current injection are also shown (right). Five minutes after a 45 sec exposure to baclofen (Bac; 100 μM), both the EPSP and IPSP remained depressed (left middle), whereas the current-evoked responses were unchanged (right middle). Full recovery was seen ~45 min after the initial exposure period. **B.** Plot showing replication of this experiment but with only 10 sec baclofen exposure periods (Bac; 100 μM) in three P6 medial limb LSO neurons. Control EPSPs (filled circles) and IPSPs (open circles) were depressed by ~50% or more 1 min after the brief baclofen exposure (Bac). The evoked synaptic potential amplitudes reached nearly complete recovery by ~15 min.

P3–P5 animals, whereas the IPSCs became primarily SN-sensitive after P11. Furthermore, GABA-evoked currents were larger and of longer duration in P4 neurons compared with glycine-evoked currents (Fig. 4). Finally, we observed a comparable change in staining with antibodies to GABA_{A} receptor and gephyrin (Figs. 8, 9). Staining for the β_{2,3} GABA_{A} receptor...
subunits in the retina (Sassoe-Pogentto and Wässle, 1997). Thus, its absence from the medial limb at P4 suggests that the transiently expressed GABA<sub>A</sub> receptors may be uniformly distributed on the postsynaptic membrane.

**Implications for LSO function**

It is not clear how much GABAergic transmission remains in the adult gerbil LSO or what its role may be. GABA-evoked inhibition is present in ~70% of adult chinchilla LSO neurons, although BIC only blocked contralateral auditory-evoked inhibition in 1 of 15 cells (Moore and Caspary, 1983). Ipsilateral inhibition has been described in vivo and in vitro (Brownell et al., 1979; Wu and Kelly, 1995), and contralateral excitation has been described for gerbil lateral limb neurons (Kil et al., 1995). However, the contralateral cochlear nucleus afferents did not apparently arborize within the medial limb from P3 onward (Kil et al., 1995). In P21–P45 mouse LSO neurons in vitro, BIC- or picrotoxin-sensitive inhibitory transmission is observed, and contralaterally evoked excitatory responses are observed in the presence of strychnine (Wu and Kelly, 1995).

The medial and lateral limbs of the gerbil LSO exhibit several distinct anatomical properties. In the medial limb, neurons have more narrow dendritic arbors (Sanes et al., 1990), there is a greater density of [3H]SN binding (Sanes et al., 1987), and there is a distinct complement of glial markers (Hafidi et al., 1994, 1996). The present data demonstrate two major differences in the development of inhibition in the lateral limb. First, there is no transition from GABAergic to glycinergic inhibition (Fig. 6), and the possibility that such a switch could have occurred in the lateral

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**Figure 8.** A, GABA<sub>A</sub> receptor staining (β<sub>2,3</sub>) at P4 shows similar intensity in both medial (m) and lateral (l) limbs of LSO. B, GlyR (gephyrin) staining in the LSO at P4 showing a stained cell (thin arrow) and axons (thick arrow) running across the nucleus. C, GABA<sub>A</sub> receptor staining at P14 showing the decreased immunoreactivity in LSO compared with adjacent periolivary areas. D, GlyR immunoreactivity at P14 shows similar levels of intensity in the neurons of medial limb (thick arrow) and lateral limb (thin arrow). Note the intensely labeled neuropil in the medial limb. Scale bar, 100 μm.
log₁₀ [Clout]/[Clin] beled neuropil (arrowheads). Therefore, inhibitory projections from the MNTB to the lateral limb must develop in a different manner from the medial limb. It does suggest that lateral limb neurons may display a distinct pattern of differentiation, because their functional properties differ from those of medial limb neurons.

Possible significance of GABAergic transmission

A number of studies indicate that GABAergic transmission is an important signal during development. Results from in vitro experiments suggest that GABAergic signaling can modulate process outgrowth (Michler-Stuke and Wolff, 1987; Spoerri, 1988; Mattis and Kater, 1989; Behar et al., 1996), synaptogenesis (Corner and Ramakers, 1992; Redburn, 1992), and GABA<sub>Ac</sub> receptor expression (Frieder and Grimm, 1985; Hablitz et al., 1989; Montpied et al., 1991; Kim et al., 1993; Liu et al., 1997; Poulter et al., 1997). Thus, neonatal GABAergic transmission in the LSO may influence the transition to glycinergic transmission.

Our previous studies have shown that manipulations designed to decrease inhibitory transmission in the LSO have a major impact on the development of structure and function (Sanes and Chokshi, 1992; Sanes et al., 1992; Aponte et al., 1996; Sanes and Hafidi, 1996). Contralateral cochlear ablation and SN rearing cause a reduction in MNTB-evoked inhibition and an unexpected enhancement of ipsilaterally evoked excitation (Kotak and Sanes, 1996). Because one manipulation uses SN, we concluded that glycinergic inhibition plays an important role in neuronal maturation. Contralateral cochlear ablation, which functionally denervates the MNTB, should have affected GABAergic and glycinergic transmission, whereas SN rearing should attenuate glycinergic transmission only. The relative efficacy of each treatment is not known, and it is possible that contralateral ablation is less effective at attenuating glycinergic transmission than SN treatment.

One possibility is that glycinergic and GABAergic transmission exert a similar influence on the maturation of postsynaptic neurons. For example, GABA or glycine can depolarize neurons during early development and cause influx of calcium (Conner et al., 1987; Ben-Ari et al., 1989; Ito and Cherubini, 1991; Obrietan and Van den Pol, 1995; Boehm et al., 1997; Lo et al., 1998). In the rat, both inhibitory potentials and glycine are almost exclusively known for glycinergic systems. Activation of presynaptic GABA<sub>B</sub> receptors inhibit transmission in neonatal rat hippocampus (Gaiarsa et al., 1995). The electrical properties of postsynaptic LSO neurons remained primarily unchanged after baclofen exposure, suggesting that ionotropic GABA<sub>B</sub> receptors are not significantly involved (Dutar and Nicoll, 1988). Rather, the prolonged time course of synaptic depression (10–15 min) indicates that metabotropic GABA<sub>B</sub> receptors may be located on afferent terminals (Bowery, 1989). Baclofen-sensitive GABA<sub>B</sub> mechanisms have been shown to modulate second messenger pathways (Tremblay et al., 1995; Barthel et al., 1995; Zhang et al., 1997). The GABAergic system appears to be one of several metabotropic pathways in the developing LSO, and we have previously described long-lasting effects of glutamate and serotonin (Kotak and Sanes, 1995; Fitzgerald and Sanes, 1997).

In the open field, low-frequency LSO neurons encounter minor level differences but may exhibit a sensitive response to time differences (Joris and Yin, 1990; Finlayson and Caspary, 1991). Therefore, inhibitory projections from the MNTB to the lateral limb may participate in temporal processing, as suggested by intracellular recordings from a brain slice preparation (Sanes, 1990). Although this discussion does not explain why inhibition in the lateral limb must develop in a different manner from the medial, it does suggest that lateral limb neurons may display a distinct pattern of differentiation, because their functional properties differ from those of medial limb neurons.
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