Synaptic Plasticity in Morphologically Identified CA1 Stratum Radiatum Interneurons and Giant Projection Cells

Brian R. Christie, 1* Kevin M. Franks, 2
Jeremy K. Seamans, 2 Karin Saga, 2
and Terrence J. Sejnowski 2, 3

1 University of South Dakota School of Medicine, Vermillion, South Dakota
2 Computational Neurobiology Laboratory, Howard Hughes Medical Institute, Salk Institute for Biological Studies, La Jolla, California
3 Department of Biology, University of California at San Diego, La Jolla, California

ABSTRACT: Long-term potentiation (LTP) of synaptic efficacy was examined in interneurons and giant cells in the stratum radiatum region of the hippocampal CA1 subfield. Cells were visually selected using differential interference contrast (DIC) optics and filled with biocytin while being recorded using whole-cell patch-clamp techniques. Electrophysiological criteria, including spike height, width, and degree of spike adaptation shown to sustained depolarization, proved inadequate for differentiating interneurons from giant cells. We found that cells in the stratum radiatum, however, could be reliably differentiated using DIC optics or following intracellular staining. The response of the two cell types to tetanic stimulation further dissociated them. Long-term potentiation, dependent on the activation of NMDAR (N-methyl-D-aspartate receptors), could reliably be induced in interneurons with stimuli administered at 200 Hz, but not 100 Hz. Giant cells, in contrast, exhibited NMDA receptor-dependent LTP in response to 100-Hz stimuli, but not the 200-Hz stimuli. LTP induction in interneurons also appeared temperature-dependent, being much more robust at 34°C than at room temperature. The LTP in both cell types required postsynaptic calcium influx, and was not due to the passive propagation of LTP induction in neighboring pyramidal cells. These results suggest that different cell types within the hippocampal formation may preferentially alter synaptic connectivity in a frequency-specific manner. Hippocampus 2000;10:673–683. © 2000 Wiley-Liss, Inc.

KEY WORDS: LTP; LTD; calcium; NMDA; hippocampus

INTRODUCTION

Long-term potentiation of synaptic efficacy is a leading candidate mechanism for learning and memory processes. In numerous studies, LTP has been reliably induced at the asymmetric spines formed onto hippocampal principal cells, i.e., the excitatory projection neurons of the hippocampal formation. The induction of LTP at synapses onto the inhibitory neurons of the hippocampal formation is more controversial (reviewed in Freund and Buzsaki, 1996; McBain et al., 1999). LTP-inducing conditioning stimulation (100 Hz, 1 s) has been reported to produce LTP (Buzsaki and Eidelberg, 1981; Taube and Schwartzkroin, 1987; Stelzer et al., 1994; Cowan et al., 1998), long-term depression (McMahon and Kauer, 1997), or no change (Maccaferri and McBain, 1996) in stratum radiatum interneurons. These same stimuli have been shown to produce LTP in stratum oriens interneurons, but not in interneurons located near the border of the stratum radiatum and stratum molecular (Ouardouz and Lacaille, 1995).

A critical concern for elucidating LTP in interneurons is their morphological identification. It was recently shown that a population of projection cells, known as giant cells, exists in the stratum radiatum region of the hippocampal formation (Gulyas et al., 1998). Giant cells exhibited an NMDAr-dependent form of LTP when depolarized to −20 mV and synaptically stimulated at 1 Hz (Maccaferri and McBain, 1996; Gulyas et al., 1998), whereas identified interneurons did not (Maccaferri and McBain, 1996). In many studies it is unclear whether interneurons or projection cells were being studied, and histological identification may be critical for differentiating these cells to determine their capacity to exhibit LTP.

There are also a number of molecular and structural differences between interneurons and principal cells.
that suggest that the rules for the induction of synaptic plasticity may differ for the two cell types. Both principal cells and interneurons have NMDA receptors and voltage-gated Ca\(^{2+}\) channels; however, interneurons appear to lack both CaMKII\(\alpha\) and calcineurin (Sik et al., 1998). Once thought integral for LTP expression, it is now clear that activation of CaMKII\(\alpha\) and calcineurin by calcium entry through the NMDA receptor is not essential for NMDA-dependent LTP (Wang and Kelly, 1996; Hinds et al., 1998). A second difference is that, compared to principal cells, interneurons are relatively devoid of spines (Amaral and Witter, 1989; Van Groen and Wyss, 1990; Sik et al., 1998; Gulyas et al., 1998), and interneuron synaptic plasticity does not appear to be synapse-specific (McMahon and Kauer, 1997; Cowan et al., 1998). The possibility that interneurons may express a form of LTP that is not dependent on NMDA activation has not been explored, and the only stimulus paradigms that have been used are ones that reliably induce NMDA-dependent LTP and LTD in principal cells. The present experiments compare the effects of different forms of tetanic stimulation in morphologically identified interneurons and giant cells located in the stratum radiatum layer of the hippocampal CA1 subfield to address some of these issues.

### MATERIALS AND METHODS

#### Slice Preparation

Hippocampal slices (350–400 \(\mu m\)) were prepared on a Vibratome (Oxford series 1000) from Sprague-Dawley rats (postnatal days 14–35). Slices were dissected in a chilled medium (pH 7.4) containing (in mM) 110.0 choline chloride, 2.5 KCl, 1.25 NaH\(_2\)PO\(_4\), 25.0 NaHCO\(_3\), 0.5 CaCl\(_2\), 7.0 MgCl\(_2\), 25.0 dextrose, 1.3 ascorbate, and 2.4 pyruvate, continuously bubbled with 95% \(O_2\)/5% \(CO_2\). To maximize the number of CA1 neurons that extend their dendritic trees in the same plane as the slice, the brains were placed on their temporal poles, and oriented at an angle of 25–30° for sectioning. Cut slices were then incubated at 32°C in artificial cerebrospinal fluid (ACSF; pH 7.4), continuously bubbled with 95% \(O_2\)/5% \(CO_2\), and composed of (in mM): 125.0 NaCl, 2.5 KCl, 1.25 NaH\(_2\)PO\(_4\), 25.0 NaHCO\(_3\), 2.0 CaCl\(_2\), 1.3 MgCl\(_2\), and 10.0 dextrose, for at least 20 minutes.

#### Electrophysiological Methods

Slices were stored in a circulating perfusion chamber and maintained at room temperature for a maximum of 6 h. Individual slices were transferred to the recording chamber as required, and experiments were performed in ACSF (see above) maintained at either 23°C or 34°C. Individual neurons were examined using a 60× Olympus water immersion objective and DIC optics (Christie et al., 1996). Whole-cell recording pipettes (5–7 M\(\Omega\)) were pulled from borosilicate glass (VWR Scientific) and filled with (in mM): 140.0 KmeSO\(_4\), 10.0 HEPES, 4.0 NaCl, 0.1 EGTA, 4.0 Tris-ATP, 0.3 Tris-GTP, and 14.0 phosphocreatine (pH 7.4, KOH). Biocytin (0.2–0.4%) was routinely added to the internal solution to allow for subsequent histological analysis. Input resistance was 200–500 M\(\Omega\) in interneurons and 100–300 M\(\Omega\) in giant cells. Cells were periodically monitored to ensure that series resistance did not change over the course of an experiment. Series resistance was always less than 40 M\(\Omega\) in both cell types. Cells located in the upper half of the stratum radiatum region of the CA1 hippocampal subfield were specifically targeted for this study. Once a cell was selected, a small (~5-\(\mu m\) tip) bipolar stimulating electrode was placed in the distal third of the stratum radiatum, approximately 150–500 \(\mu m\) lateral to the soma. A tight seal (>1 G\(\Omega\)) was initially formed, and then whole-cell recordings were made by briefly applying negative pressure to rupture the cell membrane in the electrode tip. The amplifier (Axoclamp-2B) was then immediately switched from voltage-clamp to current-clamp mode, and the resting membrane potential of the neuron was recorded. Data were filtered at 2 kHz and digitized at 5–10 kHz, using an A/D interface (National Instruments) and analyzed using custom written software (Lee Campbell). Stimulus strengths were adjusted to 30% of that required to illicit a maximal EPSP.

#### Stimulation Protocols

Stimulus isolation units (Getting Instruments) provided constant current stimulation (110 \(\mu s\) duration, 15–200 \(\mu A\) intensity) via the stimulating electrode (A-M Systems) located in the stratum radiatum, as described above. Stimuli were applied every 15 s (0.25 Hz) to assess basal synaptic transmission. For LTP induction, a total of 200 pulses was administered at either 100 Hz or 200 Hz. Stimuli were presented as 4 trains of 50 pulses, 100-Hz and 200-Hz stimuli as 2 trains of 100 pulses. In all experiments, excitatory postsynaptic potential (EPSP) data are expressed as a percentage relative to the initial 10-min baseline period. All statistical data were computed as an average of 20 responses per slice over a 5-min period 25–30 min post-conditioning, except where otherwise noted. Data are expressed at mean \(\pm\) SEM.

#### Histology

Slices were fixed immediately after electrophysiological recording in 4% paraformaldehyde and refrigerated for at least 24 h. Following several washes in phosphate-buffered saline (PBS), the sections were then washed in PBS containing 10% methanol and 1% hydrogen peroxide. Biocytin-injected cells were then visualized, using an avidin-biotinylated horseradish peroxidase (HRP) complex reaction (Vectastain ABC kit, Vector Labs) and precipitated with hydrogen peroxide in diamino benzidine. The reaction was intensified with a solution containing nickel ammonium sulfate (1%) and cobalt chloride (1%). The slices were then mounted on gelatin-coated slides, dehydrated, and coverslipped for post hoc microscopic analysis.
RESULTS

Identification of Stratum Radiatum Cell Types: Morphology

The identification of the different cell types was a major concern for this series of experiments. Figure 1 shows examples of both a bipolar interneuron and a giant cell using DIC optics. Note how prominent the thick bifurcating apical dendrites of the giant cell are, while little or no dendritic process is evident around the smooth spherical surface of the interneuron soma. During the course of experimentation, we found that the shape of the soma, as well as an absence of discernible dendritic processes (especially the thick bifurcating dendrite that typifies giant cells), were the best indicators for classifying cells as interneurons using DIC optics. In most cases, interneuron somata appeared as round or ovoid spheres. Conversely, the larger pyramidal-shaped soma and thick bifurcating dendrite have become our main criteria for identification of giant cells in the upper stratum radiatum region with DIC optics.

Cells were filled with biocytin (0.2–0.4%) so that they could be morphologically identified using light microscopy. Figure 1C,D shows the distinct morphologies of the somata and main dendrites arising from interneurons and giant cells in the stratum radiatum. Note how two main dendrites extend in the apical direction away from the soma of the giant cell in the right, while in the interneuron, numerous smaller dendritic processes extend laterally in both directions. As depicted in Figure 1E,F, interneurons and giant cells could be further differentiated by the absence or presence of spiny processes on the dendrites (Gulyas et al., 1998; Parra et al., 1998). In the present experiments, all cells classified as interneurons were relatively devoid of dendritic spines. These results are in agreement with works that morphologically described stratum radiatum neurons (Lacaille and Schwartzkroin, 1988; McBain and Dingledine, 1993; Valtschanoff et al., 1993; Williams et al., 1994; Maccabery and McBain, 1996; Gulyas et al., 1998). Our results indicate that post hoc histological analysis is an extremely useful, but not essential, means to establish neuronal identity. The DIC optics used in the present study provides an equally reliable means to differentiate giant cells from interneurons in living tissue. Figure 2 presents some examples of the cell types examined in the present work. A key feature for this study is that all of the cell bodies and the majority of dendrites of each cell were confined mainly to the stratum radiatum of the hippocampal CA1 subfield.

Identification of Stratum Radiatum Cell Types: Electrophysiology

A commonly used method for distinguishing interneurons from principal cells electrophysiologically is to examine neuronal responses to depolarizing current injections. Interneurons are identified in this manner as those cells that 1) exhibit trains of nonaccommodating action potentials in response to depolarizing current injection, and/or 2) have short-duration action potentials which are followed by a deep fast afterhyperpolarizing potential (fAHP) (Schwartzkroin and Mathers, 1978; Lacaille et al., 1989; Buhl et al., 1995; Cowan et al., 1998).

In the present study, the only prominent feature of the electrophysiological profile of interneurons that was not commonly observed in giant cells was the presence of a pronounced fAHP (Fig. 3). Attempts to dissociate the two cell types on the basis of spike width and amplitude were inconclusive. While giant cells typically had broader and larger amplitude action potentials than interneurons, it was often the case that both cell types had action potentials of equivalent duration and amplitude (Fig. 3A,B). Similarly, both identified giant cells, and interneurons exhibited a wide range of firing patterns in response to depolarizing current injections. Giant cells commonly exhibited some degree of spike frequency accommodation (Fig. 3C,D). However, adaptation was not observed in all cells (Fig. 3E). The interneurons recorded in this study displayed a regular firing pattern for the most part (Fig. 3F,J), although some cells exhibited burst firing (Fig. 3H) or were strongly accommodating (Fig. 3G) to the same stimuli. These findings are similar to those for anatomically identified basket and bistratified cells in the hippocampus (Buhl et al., 1996). These results indicate that it is difficult to dissociate stratum radiatum giant cells from interneurons on the basis of their firing characteristics alone.

Stimulation (200 Hz) Induces LTP in Interneurons But Not Giant Cells

LTP induction was attempted at both 23°C (room temperature) and 34°C. As shown in Figure 4, the stimuli administered to interneurons at 200 Hz were ineffective at inducing LTP at room temperature (30 ± 13%; t(9) = -2.19, P = 0.051), while LTP was reliably induced by these same stimuli at 34°C (91 ± 29%; t(9) = 3.1, P < 0.05). At 23°C only 6/10 cells exhibited LTP greater than 20%, while 3 cells exhibited <20% change, and one cell exhibited >20% LTD. With the perfusion maintained at 34°C, LTP (>20%) was observed in 8/10 neurons. One neuron in this group exhibited >20% LTD. A direct comparison of the two groups revealed that cells tested at 34°C showed a significantly greater long-lasting LTP than that exhibited by those tested at 23°C (t(9) = 2.6, P < 0.05).

Stratum radiatum giant cells tested at 23°C, like interneurons, failed to exhibit a significant degree of long-lasting LTP (5 ± 8%, n = 6; P > 0.05; Fig. 4). Only one cell showed >20% LTP, one showed smaller LTD, and the remaining four did not exhibit any degree of plasticity. LTP was not larger at 34°C in giant cells (19 ± 13%, n = 12; P > 0.05). Six cells did not exhibit any change in synaptic efficacy, while 5/12 exhibited LTP (>20%) and one exhibited LTD (>20%). There was no significant difference in degree of plasticity observed between the two cell types at 23°C. However, interneurons showed significantly greater LTP than giant cells when tested at 34°C (t(20) = 2.359, P < 0.05).

Conditioning Stimulation (100 Hz) Induces LTP in Giant Cells But Not Interneurons

Given the previous results, all subsequent experiments were performed at 34°C. As shown in Figure 6, synaptic efficacy was increased markedly in giant cells (51 ± 18%, t(10) = 2.8, P < 0.05). Individually, 6/11 cells exhibited LTP >20%, while alterations in
synaptic efficacy in the remaining cells were <20%. No cells exhibited LTD. In contrast to the giant cells, radiatum interneurons showed smaller increases in synaptic efficacy on average (26 ± 14%, n = 12; P > 0.05). Five cells exhibited >20% in synaptic efficacy, while six showed >20% change, and the remaining cell exhibited LTD >20%.

FIGURE 1. Anatomical characterization of interneurons and giant cells in the hippocampal CA1 subfield stratum radiatum. A,B: Differential interference contrast (DIC) photomicrographs of whole-cell pipettes attached to cell bodies in the CA1 stratum radiatum, taken with a 60× water immersion lens. The large bifurcating apical dendrites of giant cells were easily viewed under these conditions (B); the dendrites of interneurons were much more difficult to distinguish (A). C,D: Photomicrographs of the cells depicted in A and B, following the expression of biocytin. Dendrites are now discernible, extending laterally in both directions in the interneuron (C), while a number of secondary dendrites are now also visible coming off the two main apical dendrites and the basal portion of the soma of the giant cell (D). Dotted areas indicate the regions enlarged and shown in E and F. E,F: Like pyramidal cells, the second-order dendrites of giant cells are covered in spines, while both the main and second-order dendrites of the interneurons examined in the present study appear aspiny. The CA1 pyramidal cell layer is located at the top of all photomicrographs, out of the field of view. Scale bars = 25 μm.

LTP Induction in Hippocampal Giant Cells and Interneurons Is Calcium-Dependent and Not Passively Propagated

Research into the mechanisms responsible for the induction of LTP suggests that an increase in postsynaptic Ca²⁺ plays an important role
FIGURE 2. Camera lucida tracings of representative giant cells and nonpyramidal cell types recorded from in the present study. A: All giant cells recorded in the present study exhibited a stereotypic somatic and dendritic morphology. The large soma was always located in the upper half of the stratum radiatum (S.R.), and dendritic processes always descended into the stratum lacunosum-moleculare (S.L.-M.). Typically the dendritic processes bifurcated at the level of the soma (1, 3, 4); however, in a few cells (2), the bifurcation occurred 50–100 μm from the soma. These cells more closely resembled pyramidal cells; however, they invariably gave rise to axonal process that bifurcated at the level of the stratum oriens (S.O.), extending away towards both the fimbria-fornix and the subiculum. In many cases, numerous axon collaterals could be seen localized to the stratum pyramidale (S.P.) and/or S.O. layers (1, 2, 4). B: Interneurons, on the whole, tended to be a more morphologically heterogeneous population of cells, with smaller somata than those of giant cells. The main common feature of these cells was that the majority of their dendritic processes were confined to the S.R. region. Many cells exhibited a bipolar appearance, with numerous small dendrites extending away both apically and basally (1). Dense axonal processes (not shown) extended into the S.P. and S.O. from many of these cells. Cells with similar somatic morphology to 1 could also extend numerous dendritic processes laterally across the S.R. (2, 3). These cells tended to localize their axonal processes around the S.P. region. A few recordings were also obtained from cells located closer to the S.L.-M. (4). These cells had numerous small dendritic processes extending laterally across the S.R. Axonal processes in these cells were mainly localized to the S.R. and S.P. layers.
To determine whether the LTP induced in these cells was an NMDAr-mediated or VGCC-mediated calcium entry into the postsynaptic cell, we included calcium chelator EGTA (10 mM) in the whole-cell pipette to severely attenuate postsynaptic calcium levels (Lynch et al., 1983; Malenka et al., 1988; Yeckel et al., 1999). Neither morphologically identified interneurons nor giant cells exhibited LTP under these conditions (Fig. 6). Giant cells exhibited a modest depression (−12 ± 18, n = 5, P > 0.05), with only 1/5 cells showing a change in synaptic efficacy greater than 20%. No interneuron tested exhibited a significant change in the slope of EPSPs recorded following the application of the tetanus under these conditions (7 ± 3%, n = 5, P > 0.05). These experiments provide evidence against the possibility that the LTP observed in giant cells and interneurons was being passively propagated from pyramidal cells (Maccarreri and McBain, 1995). Since LTP was blocked under these conditions, it is unlikely that the previous results reflect pyramidal cell LTP being expressed disynaptically.

**Role of NMDAr in Stratum Radiatum Neuronal Synaptic Plasticity**

We next attempted to ascertain the mechanisms involved in LTP genesis in interneurons and giant cells. Although the LTP induced in pyramidal cells with 200-Hz stimulation consists of both an NMDAr-mediated component and a VGCC-mediated component (Grover and Teyler, 1990a; Grover, 1998), the contribution of the NMDAr appears greater in young animals like those used in this study (Shankar et al., 1998). We found that both hippocampal stratum radiatum interneurons and giant cells exhibited NMDAr-mediated responses when DNQX (20 μM) and Bicuculline (2 μM) were added to the ACSF (data not shown).

To examine the role of NMDA-receptor activation in stratum radiatum neuronal LTP, whole-cell recordings were made in the presence of the NMDA-receptor antagonist DL-APV (1-amino-5-phosphonovaleric acid). As shown in Figure 7, the 200-Hz conditioning stimulation did not induce LTP in interneurons when APV was present (0.1 ± 6%; n = 6; t_{(5)} = 0.7, P > 0.05). Individually, LTP (>20%) was observed in only one cell; LTD (>20%) was apparent in another; and the remaining four cells did not exhibit any significant changes.

In giant cells, LTP was preferentially induced by the 100-Hz stimuli in the previous experiments. However, the addition of APV prevented the induction of LTP in giant cells (13 ± 16%, n = 6). In some cells, we attempted to induce LTP following APV exposure. However, even following a 30-min washout in normal ACSF, we were unable to induce LTP in either cell type. Given the long time course of these experiments, this may reflect a washout of some critical intracellular component that is essential for LTP induction in these cells.

We also examined the effects of NMDA-receptor blockade in giant cells to see if we could elicit an NMDAr-independent form of LTP with the 200-Hz conditioning stimuli, similar to that observed in pyramidal cells (Grover and Teyler, 1990a, 1995; Grover, 1998). Surprisingly, a long-lasting form of LTD developed in these cells rather than the expected LTP (−34 ± 9%, n = 7; t_{(6)} = 3.7, P < 0.05). This finding, along with the failure of the 200-Hz stimuli to generate LTP in control conditions, clearly differentiates giant cells from pyramidal neurons (Fig. 8).
FIGURE 4. Temperature- and frequency-specific synaptic plasticity in stratum radiatum interneurons and giant cells. A,B: Averaged evoked responses recorded from morphologically identified interneurons (A) and giant cells (B) prior to (gray) and following (black) the application of 200-Hz tetanic stimulation at either 23°C (A1,B1) or 34°C (A2,B2). C: Individual responses recorded from interneurons administering the 200-Hz conditioning stimuli at either 23°C (open circles) or 34°C (solid circles). D: Individual responses recorded from giant cells administered the 200-Hz conditioning stimuli at either 23°C (open triangles) or 34°C (solid triangles). E,F: Averaged group responses for all interneurons (E) and giant cells (F) receiving the 200-Hz conditioning stimulation at the two temperatures. A significant degree of LTP was only produced in interneurons in ACSF maintained at 34°C (91 ± 29%, n = 10; t_{0.05} = 3.1, P < 0.05) and not in giant cells maintained at the same temperature (19 ± 13%, n = 12; P > 0.05).
DISCUSSION

Differentiating Stratum Radiatum Giant Cells From Interneurons

The present experiments confirm previous reports that differentiate hippocampal stratum radiatum giant cells and interneurons on the basis of several morphological differences (Lacaille et al., 1989; van Groen and Wyss, 1990; Maccaferri and McBain, 1996; Gulyas et al., 1998). The large soma and bifurcating apical dendrites of giant cells were easily visible using DIC optics, and could often be followed for several hundred microns as they extended towards the stratum moleculare region. The dendrites of interneurons, in contrast, could usually not be followed for more than 50 mm. Second, the somata of giant cells were typically triangular in appearance and approximately 20 μm in size. The somata of the interneurons used in this study were either round or ovoid in appearance, and appeared much smaller (10–15 μm) under DIC optics. Finally, the biocytin-stained interneurons used in this study possessed dendritic processes whose primary and secondary branches were almost utterly devoid of visible spines. Giant cells, in contrast, possessed dendritic processes that were heavily covered in spiny processes. Gulyas et al. (1998) suggested that these morphological features indicate that giant cells are likely excitatory cells and not inhibitory. It needs to be established whether these cells are GAD-positive to determine whether these cells are excitatory or inhibitory.

Electrophysiologically, the large fAHP differentiated interneurons from giant cells, which lack a large fAHP. There were no differences in action potential height and width that could be used to reliably differentiate the cell types. Both interneurons and giant cells could respond with adapting or nonadapting trains of action potentials, though a small population of interneurons also responded with bursts of action potentials (see also Lacaille et al., 1988; Williams et al., 1994). This variability in the responsiveness of these cells makes it difficult to differentiate between them on the basis of electrophysiological criteria alone, and we were only fully confident of a cell’s classification when confirmed visually.

FIGURE 5. Giant cells, but not interneurons, exhibited LTP following 100-Hz conditioning stimulation. A,B: Ensemble averages of evoked responses obtained prior to (i) and following (ii) the application of the 100-Hz stimuli in a morphologically identified interneuron (A) and giant cell (B). C,D: Percentage change in recorded synaptic EPSPs from an interneuron (C) and a giant cell (D) prior to and following the application of the 100-Hz stimuli. E,F: Averaged responses of all interneurons (E) administered the 100-Hz stimuli do not exhibit a significant degree of LTP (26 ± 14%, n = 12), while those obtained from giant cells (F) are significantly potentiated (40 ± 16%, t(9) = 2.5, P < 0.05).
Frequency-Specific LTP and LTD in Stratum Radiatum Giant Cells and Interneurons?

The application of high-frequency conditioning stimuli served to further differentiate the two cell types. A significant degree of LTP was only produced in interneurons following the application of a 200-Hz stimulus, and not following 100-Hz stimuli. A novel aspect of the LTP observed with the 200-Hz stimulation was that it was induced at physiological temperatures, but not at room temperature. In contrast, LTP was not induced in giant cells with the 200-Hz stimuli, but was observed following the application of tetanic stimulation at 100 Hz.

Another difference in the two cell types was apparent following the administration of the 200-Hz conditioning stimuli, while NMDAr were blocked with APV. Under these conditions, LTP was prevented in stratum radiatum interneurons, while a significant degree of LTD was induced in giant cells under NMDAr blockade. This difference suggests that 200-Hz stimulation may simultaneously induce an NMDAr-dependent LTP and NMDAr-independent LTD in giant cells. In contrast to interneurons, but like CA1 pyramidal cells (Reymann et al., 1989; Grover and Teyler, 1990b), neither LTP nor LTD were induced when giant cells were administered the 100-Hz stimuli in the presence of APV.

The LTP observed in giant cells and interneurons did not appear to be passively propagated from nearby pyramidal cells. The term “passive propagation” refers to LTP observed in the secondary components of EPSPs evoked in stratum oriens-alveus inhibitory neurons but mediated by pyramidal neurons (Macciaferri and McBain, 1995). The inclusion of the calcium chelator EGTA in the intracellular medium prevented the induction of monosynaptic LTP in both cell types. This manipulation suggests that the observed LTP is also calcium-dependent.
Implications of Stratum Radiatum Neuronal LTP for Hippocampal Functioning

Taken together, these findings provide further evidence that cells in the stratum radiatum region are capable of exhibiting synaptic plasticity. The ability of interneurons to exhibit long-term alterations in synaptic efficacy has a number of implications for the modulation of hippocampal circuits. Tsodyks et al. (1997) showed that increasing the strength of the inhibitory drive onto interneurons can increase network stability, and paradoxically increase the firing rate of interneurons. Since LTP of EPSPs and IPSPs may occur concurrently in a population of cells receiving a given input, Perez et al. (1999) suggested that LTP of interneurons could enhance the detection of populations of neurons that are being excited from those that are not. Interneuron LTP would result in more inhibition, to downstream neurons, reducing their output. Giant cells may be strategically placed in the stratum radiatum to signal output regions that hippocampal pyramidal cell afferents are currently being inhibited. Future experiments will attempt to determine if these giant cells are the recipients of local circuit inhibitory connections as well.

Acknowledgments

The authors thank G. Buszaki, J. Kauer, J.-C. Lacaille, J. Magee, C. McBain, and C. Stevens for helpful comments during the course of these experiments. We also thank Lee Campbell for providing the data acquisition and analysis software and Bryan Nielsen for technical support.
REFERENCES


