Progressively increasing (augmenting) responses are elicited in thalamocortical systems by repetitive stimuli at ~10 Hz. Repeated pulse trains at this frequency lead to a form of short-term plasticity consisting of a persistent increase in depolarizing synaptic responses as well as a prolonged decrease in inhibitory responses. In this study, we have investigated the role of thalamocortical (TC) and neocortical neurons in the initiation of thalamically and cortically evoked augmenting responses. Dual intracellular recordings in anesthetized cats show that thalamically evoked augmenting responses of neocortical neurons stem from a secondary depolarization (mean onset latency of 11 msec) that develops in association with a diminution of the early EPSP. Two nonexclusive mechanisms may underlie the increased secondary depolarization during augmentation: the rebound spike bursts initiated in simultaneously recorded TC cells, which precede by ~3 msec the onset of augmenting responses in cortical neurons; and low-threshold responses, uncovered by hyperpolarization in cortical neurons, which may follow EPSPs triggered by TC volleys. Thalamic stimulation proved to be more efficient than cortical stimulation at producing augmenting responses. Stronger augmenting responses in neocortical neurons were found in deeply located (<0.8 mm, layers V–VI) regular-spiking and fast rhythmic-bursting neurons than in superficial neurons. Although cortical augmenting responses are preceded by rebound spike bursts in TC cells, the duration of the self-sustained postaugmenting oscillatory activity in cortical neurons exceeds that observed in TC neurons. These results emphasize the role of interconnected TC and cortical neurons in the production of augmenting responses leading to short-term plasticity processes.

Key words: augmenting responses; thalamus; neocortex; plasticity; dual intracellular recordings; EPSP
neurons largely depend on the postinhibitory rebound spike bursts characterizing the LT-type responses in TC cells. However, the duration of the self-sustained postaugmenting oscillatory activity in cortical neurons exceeds that observed in TC neurons. These results emphasize the role of interconnected TC and cortical neurons in the production of augmenting responses, which lead to short-term plasticity processes. In a companion paper (Bazhenov et al., 1998b) computer simulations of thalamocortical augmenting responses are used to explore the underlying mechanisms.

MATERIALS AND METHODS
Experiments were conducted on adult cats of either sex (n = 39), some anesthetized with pentobarbital (35 mg/kg, i.p.), others maintained under ketamine–xylazine anesthesia (10–15 mg/kg and 2–3 mg/kg, i.m.). Similar results on augmenting responses were obtained under both types of anesthesia (each figure legend mentions the experimental condition). In addition, tissues to be excised and pressure points were infiltrated with a local anesthetic (lidocaine, 2%). The depth of general anesthesia was continuously monitored by recording sleep-like EEG patterns (spindle and slow oscillations). Additional doses of anesthetics were administered at the slightest tendency toward lower-amplitude, faster-frequency EEG waves. The heart rate was monitored by means of electrocardiogram and kept constant (90–110 beats/min). Body temperature was maintained at 37–39°C. Once the EEG indicated that anesthesia induced sleep-like patterns, the animals were paralyzed with gallamine triethiodide and artificially ventilated by maintaining the end-tidal CO2 concentration at 3.5–3.8%. The stability of intracellular recordings was ensured by cisternal drainage, hip suspension, bilateral pneumothorax, and covering the whole hemisphere with a warm solution of agar (4% in 1% saline).

Stimulation and recordings. Repetitive stimuli (trains of five pulses at 10 Hz), with variable intensities (0.02–0.3 mA) and durations (0.05–0.2 msec), were delivered through stereotaxically inserted coaxial electrodes into relay [ventralateral (VL)], association [lateroposterior (LP)], and rostral intralaminar [centrolateral (CL)] thalamic nuclei, as well as in prefrontal (areas 4 and 6) and suprasylvian (areas 5 and 7) cortices, depending on the sites of recordings. Intracellular recordings were performed in conjunction with recording of field potentials from neocortical motor and association areas 4, 6, 5, 7, and 21. In the thalamus, we recorded intracellularly from VL and RE nuclei. For intracellular recordings and staining, we used glass micropipettes filled with a solution of 2.5–3 M potassium acetate and 2% neurobiotin (DC resistance, 30–50 MΩ). A high-impedance amplifier with active bridge circuitry was used to record the membrane potential (Vm) and inject current into the cells. Intracellular activities were recorded, together with field potentials, on an eight-channel tape with a bandpass of 0–9 kHz, digitized at 20 kHz for off-line computer analysis.

Histology. At the end of experiments, the animals were given an intravenous lethal dose of pentobarbital and perfused intracardially with physiological saline, followed by 4% paraformaldehyde and 1% glutaraldehyde. The brain was removed, stored in formalin with 30% sucrose, and finally sectioned at 80 μm, processed with the avidin–biotin standard kit, mounted on gel-dipped slides, and coverslimped. Reconstructions of different types of cortical neurons (see Fig. 12) were performed from series of adjacent sections. The difference between the depth of cortical cells estimated by intracellular staining and by micromanipulator readings was <15% (Steriade et al., 1993a; Contreras and Steriade, 1995).

RESULTS
We present the results in the following order. First, intracellular recordings from single cortical neurons show the selectivity of augmenting responses in different cortical areas, evoked by stimulation of the appropriate dorsal thalamic nucleus, the secondary depolarizing component that selectively develops during augmentation, and the dependency of augmenting response amplitude on the immediate history of the neuronal network. Next, results from dual intracellular recordings from thalamus and cortex demonstrate that the secondary depolarizing component, which characterizes augmenting responses in neocortical neurons, is preceded by postinhibitory rebound bursts in the simultaneously impaled TC cell. Finally, dual intracellular recordings and staining of cortical neurons reveal the shorter latencies and higher propensity to thalamically evoked augmenting in deeply lying, compared with more superficial, regular-spiking and fast rhythmic-bursting cells.

Database and neuronal identification
The following results are based on intracellular recordings of 320 cortical neurons from motor pericruciate areas 4 and 6 and association suprasylvian areas 5, 7, and 21, from 189 thalamic neurons from VL nucleus, and from 9 RE cells. Neurons retained for analysis were recorded for at least 20 min (but up to 2 hr). The resting Vm was more negative than −60 mV in cortical neurons (mean, −66 ± 2.4 mV) and more negative than −55 mV in TC cells (mean, −60 ± 1.8 mV). Action potentials were overshooting. Of those 518 cortical and thalamic neurons, we performed 59 simultaneous recordings from cortical area 4 and thalamic VL or RE neurons and 27 simultaneous recordings from two cortical neurons located in areas 4 and 5 or 7.

(1) Cortical neurons belonged to different electrophysiologically defined classes: regular-spiking (68%), intrinsically bursting (12%), and fast rhythmic-bursting (20%). The former two classes have been described in previous in vitro (Connors et al., 1982; McCormick et al., 1985) and in vivo (Núñez et al., 1993) studies. The fast rhythmic-bursting cells have been described in superficial layers of visual cortex (Gray and McCormick, 1996) as well as in superficial and deep layers of motor and association areas (Steriade, 1997; Steriade et al., 1998), and they produce depolarization-dependent high-frequency (400–600 Hz) spike bursts recurring rhythmically at 30–40 Hz. Cortical neurons received short-latency excitation from appropriate thalamic nuclei, and some of them were formally identified as corticothalamic by antidromic invasion (see Figs. 3, 4). (2) All neurons recorded from dorsal thalamic nuclei belonged to the TC class, as demonstrated by powerful LT rebound spike bursts deactivation by membrane hyperpolarization (Deschénes et al., 1984; Jahnsen and Linás, 1984) and backfiring from the projection cortical area. (3) RE neurons, recorded from the rostral lateral sector of the RE nucleus, were identified by their depolarizing envelope during spindles and the accelerating-decelerating pattern of their spike bursts (Domingh et al., 1986; Steriade et al., 1986).

Buildup of thalamically elicited augmenting responses in cortical neurons
We first investigated the topographical relation between the stimulated site in the dorsal thalamus and the type of evoked responses in different neocortical areas. Although augmenting responses are localized in cortical territories that are topographically related to the thalamic stimulus, and the field potentials are depth-negative (Morison and Dempsey, 1942; Spencer and Brookhart, 1961), cortical recruiting responses elicited by the thalamic ventromedial (VM) nucleus, which projects widely over the cerebral cortex, are negative at the surface, because the VM nucleus exerts depolarizing actions onto layer I (Glenn et al., 1982). The present study mainly dealt with depth-negative field potentials associated with depolarizing responses in cortical neurons, characteristic of augmenting responses. Occasionally, however, prevalent depth-positive augmentation was observed (see VL-evoked field responses in Fig. 1 and CL-evoked recruiting in area 5, simultaneously with CL-evoked augmenting in area 21; see Fig. 4). In fact, augmenting responses should not be considered as sharply distinct from recruiting ones: both types
depend on the prevalent (although not exclusive) laminar projections of different thalamic nuclei. The VM and some intralaminar nuclei preferentially project to layer I but also have deeper projections, and specific relay nuclei have, in addition to their major projections to middle layers, superficial projections arising from small-sized neurons (Jones, 1985; Steriade et al., 1997).

During spontaneously occurring spindles, intracellularly recorded cortical activities and field potentials from related thalamic nuclei were time-locked (see Fig. 8). Stimulation of various thalamic nuclei, while recording intracellularly from the same cortical neuron, showed that clear-cut augmenting responses were selectively induced by stimulating the thalamic nucleus projecting to the recorded cortical area. Figure 1 shows that LP stimulation elicited intracellular and field potential augmenting responses in suprasylvian area 7 but not in area 4, whereas VL stimulation elicited recruiting responses with highest amplitudes in area 4. Cortical stimulation, even when applied close (∼2 mm) to the intracellularly recorded area 7 neuron, did not elicit augmentation but, rather, a prolonged hyperpolarization.

As shown in Figures 1 and 2, cortical augmenting responses developed from a secondary depolarizing component that appeared from the second thalamic stimulus and continued until the last (fifth) stimulus in the 10 Hz train. This feature was observed in all analyzed cortical neurons (n = 120). In a majority of them (n = 92), the increased amplitude of the secondary depolarization (initiated at 7–16 msec; mean, 11 ± 0.8 msec) was associated with a diminished amplitude of the early depolarization. In Figure 2, the LP-evoked early depolarization in area 7 neuron diminished in amplitude by 65% at the second stimulus, and the VL-evoked early depolarization in area 6 neuron diminished by 37%. In both neurons, incremental responses grew from the secondary depolarization, from the second (Fig. 2A) or third response (Fig. 2B), continuing up to the end of the pulse train.

The study of thalamically evoked augmenting responses in cortical neurons at different membrane potentials (n = 27) provided further evidence for the selective development of augmenting responses from a secondary depolarization. The layer V corticothalamic cell in Figure 3 (identified by backfiring from CL nucleus) displayed a monosynaptic EPSP at 2.5 msec, evoked by the first CL stimulus in the pulse train at 10 Hz. The response latency did not change from rest (∼64 mV) to a hyperpolarized level (∼72 mV). At a depolarized level (∼55 mV), the cell also

Figure 1. Specificity of augmenting responses in different thalamocortical systems. Barbiturate anesthesia. Three simultaneously recorded sites (each with three superimposed traces) are (from top to bottom) field potentials from depth of area 4 and depth of area 7 and intracellular activity of area 7 neuron. The LP-evoked (10 Hz) augmenting responses in area 7 neuron are expanded below (arrow). Below, averages (n = 3) of responses to different stimuli in the pulse train at 10 Hz (numbers of responses to LP stimuli correspond to those in the above panel) showing that augmenting already occurred with the second stimulus and selectively developed from a late depolarization whose onset latency was at ∼13 msec. Repetitive LP stimulation induced augmenting responses in both neuron and field potentials recorded from area 7 (and virtually no response in area 4); VL stimulation induced recruiting responses with highest amplitudes in area 4. Stimulation of cortical area 7 evoked negligible augmentation in area 7 cell and field potentials and produced steady hyperpolarization (∼6 mV) in area 7 cell. Time and voltage calibration in the bottom right panel is valid for all other panels (with the exception of expanded averages).
Figure 2. Cortical augmenting responses to thalamic repetitive stimuli (10 Hz) develop from a late depolarization. Barbiturate anesthesia. A, LP-evoked augmenting responses in field potentials and intracellularly recorded neuron from suprasylvian area 7. Three superimposed traces. Below, Average (n = 3) of cellular responses to the first and second stimuli in the 10 Hz train (left) and to all five stimuli in the pulse train (right). Note that the amplitude of the early EPSP (latency, ~5 msec) diminished at the second stimulus and that a secondary depolarization (onset latency, ~12 msec), leading to action potential, appeared from the second stimulus. B, Precruciate area 6 cell, displaying augmentation with 10 Hz VL stimuli. Average (n = 3) of responses shows that, compared with the response evoked by the first stimulus in the train (thick trace), the amplitude of the early EPSP (latency, 1.7 msec) evoked by the second stimulus diminished (as in A), and late augmented depolarization (latency, ~7–8 msec) occurred starting with the third response (asterisk at right). Note multiple EPSPs building up the secondary depolarization.
Figure 3. Thalamocortical augmenting responses in area 7 corticothalamic cell at different $V_m$ levels. Barbiturate anesthesia. Stimulation (10 Hz) of rostral intralaminar CL nucleus. Inset, Neuronal identification by CL stimulus: at $-55$ mV (top traces), antidromic spikes (latency, 0.45 msec) followed by orthodromic responses (latency, 2.5 msec); at $-64$ mV (bottom traces), antidromic spikes failed and synaptic responses survived. Three traces show (left from top to bottom) augmenting responses at a depolarized, resting and hyperpolarized level. The first and fifth responses in the train are expanded below in the same order (from left to right: depolarization, rest, hyperpolarization). Under steady depolarization (+0.5 nA), the augmented response to the fifth stimulus displayed a delayed appearance of the first action potential but a greater number of spikes in the secondary augmented response (compared with that of the 1st response). Also note self-sustained (1 sec) oscillatory response after stimulation, visible at the resting and hyperpolarized levels.
fired an antidromic spike (latency, 0.45 msec) in advance of the synthetically driven spike train. By contrast, the augmenting response started at 8.5 msec (simultaneously with a decrease in the early EPSP), at rest as well as at a hyperpolarized level, and consisted of a burst with a frequency of 300–400 Hz.

Augmenting responses were enhanced when preceded by spontaneously occurring spindle oscillations \((n = 42)\). In the example depicted in Figure 4, the area 7 corticothalamic neuron was recorded simultaneously with field potentials from the more anterior area 5 and more posterior area 21. Interestingly, the initially depth-negative field augmenting responses in area 21 contrasted with depth-positive (surface-negative) augmenting responses in area 5. This emphasizes that the same thalamic site (in this case the rostral intralaminar CL nucleus) may have different laminar projections to different cortical areas. In the intracellularly recorded area 7 neuron, the CL-evoked depolarizing augmenting responses had a pattern similar to that of spontaneous spindles. When occurring after a spontaneous spindle wave, the augmented responses displayed the same features (including the diminished amplitude of the primary EPSP) but the amplitude of the secondary depolarization increased and led to more action potentials than when the testing CL pulse train was delivered during interspindle lulls (Fig. 4; see Discussion).

Augmenting responses were followed by oscillatory activities within the frequency range of the testing pulse train \((n = 215)\). The frequency of self-sustained activity ranged from 8 to 11 Hz \((mean 10 \pm 1.3 Hz)\), and its duration was \(-1 sec\) (see Figs. 3, 4, 7, 8).

**Development of cortical augmenting responses after the low-threshold rebound bursts in TC neurons**

Dual intracellular recordings were made from cortical area 4 and related VL neurons \((n = 32)\). Without exception, the initiation of augmenting responses in area 4 neurons lagged the action potentials of spike bursts crowning the LT-augmented responses in TC cells from the VL nucleus. This relation resulted from analyses using spike-triggered averages, the reference being the first action potential in the augmented thalamic response.

Thalamic stimulation within the VL nucleus induced monosynaptic EPSPs, occasionally preceded by antidromic activation, in TC cells. The excitation was followed by a biphasic IPSP, described in vitro (Hirsch and Burnod, 1987; Crunelli et al., 1988) and in vivo (Paré et al., 1991). During the first, C1 \(-\)dependent IPSP, the fast \((-100 Hz)\) subthreshold oscillations in VL cells, originating in deep cerebellar nuclei (Timofeev and Steriade, 1997), were obliterated (Fig. 5). The second, 0.1 sec-delayed stimulus in the 10 Hz pulse train reached the cell during the late part of the IPSP and triggered an LT-type augmented response. The temporal relations between TC and cortical neurons during augmenting responses are illustrated in Figure 5. Similar results were found with both ketamine–xylazine (Fig. 5) and barbiturate anesthesia (data not shown). The responses of thalamic and cortical neurons shared similarities but also exhibited some differences: (1) in Figure 5, spike-triggered averages indicate that, from the second to the fifth augmented responses, the first action potentials in the postinhibitory rebound spike bursts of TC cells (marked by asterisk) preceded by \(-3 m sec\) the onset of augmented secondary depolarizations in cortical neurons; and (2) however, although the number of spikes in the rebound bursts of TC cells progressively increased from the second to the fifth response, the spike trains in the simultaneously recorded cortical neurons did not increase in parallel to TC responses. Thus, in some cases, the maximum number was reached in the third cortical response. The augmented depolarization of the cortical neuron in Figure 5, occasionally superimposed by single spikes, progressively diminished from the onset to the end of incremental responses. In a sample of 27 cortical and thalamic neurons, we determined the evolution of number of spikes, from the first to the fifth response evoked by thalamic repetitive stimuli. The probability of discharge to the first stimulus in the pulse train at 10 Hz was \(p = 0.52\) for TC cells and \(p = 0.31\) for cortical cells. For the second and third responses, the average number of spikes increased by 275 and 438\% in TC cells, and by 238 and 429\% in cortical cells. For the fourth and fifth responses, however, TC cells continued to increase their discharges (546 and 584\% compared with the first stimulus), but cortical neurons reached a saturation level and increased comparatively less or not at all the number of action potentials (477 and 435\%).

We measured the area \((millivolts \times milliseconds)\) of secondary depolarization in cortical neurons as a function of the number of action potentials in the rebound spike bursts of TC cells during different successive stimuli in pulse trains eliciting augmenting responses \((n = 7)\). In Figure 6, the area of secondary depolarization in area 4 neuron (Fig. 6b) was stippled to distinguish it from the early (Fig. 6a) excitatory component elicited by VL stimulation. The averaged augmenting responses to the second and third thalamic stimuli show that the onset of the postinhibitory rebound spike burst in the VL neuron preceded by \(-3 m sec\) the onset of the secondary depolarization in the cortical neuron. And the plots indicate a progressive increase in the area of secondary depolarization, parallel to the increased number of action potentials of the VL cell in response to successive stimuli in the pulse train \((r = 0.8; p < 0.001)\).

**Persistent oscillations in cortical neurons after thalamically evoked augmenting responses**

All 320 recorded neocortical neurons displayed activities that outlasted the last stimulus in the 10 Hz pulse train. The self-sustained rhythmic oscillations, at \(-10–12 Hz\), were analyzed from both cortical field potentials and intracellularly recorded activities of TC and cortical neurons. (1) Of those 320 neurons, 58 (18\%) displayed only one postaugmenting rebound (Figs. 5, 6; also see Fig. 9, *Thalamic stimulation*). However, even when there was a single rebound intracellularly, multiple oscillatory cycles occurred in cortical field potentials (Fig. 5), suggesting that other cortical neurons were repetitively depolarized. (2) Of 59 simultaneous recordings from cortical and thalamic VL or RE neurons showing postaugmenting oscillatory activities, the rhythmic self-sustained activity in 16 neurons (27\%) had a similar duration in the thalamic and coupled cortical cells (see Fig. 8, *Evoked*). (3) In the majority of cortical neurons, rhythmic waves lasted 0.2–0.6 sec longer than those recorded simultaneously in thalamic cells \((n = 32; 54\%)\). The duration of self-sustained cortical activity was longer than thalamic activity (Fig. 7). Even when the self-sustained oscillation was equally long in TC and cortical neurons after thalamically evoked augmenting responses, the oscillations outlasting spindles lasted longer in the cortical cell than in the TC cell (Fig. 8).

**Thalamic stimulation is more efficient than cortical stimulation in eliciting augmentation**

The demonstration that the secondary depolarization in cortical neurons follows the rebound spike bursts of TC neurons during augmenting (Fig. 6) suggests that thalamic events are mainly
Figure 4. Increased thalamocortical augmenting responses when preceded by spontaneous spindle waves. Barbiturate anesthesia. The three traces depict (from top to bottom) field potentials from the depth of suprasylvian areas 5 and 21 and intracellularly recorded corticothalamic neuron from area 7 (antidromic spikes evoked by threshold CL stimulation are depicted below; latency, 0.4 msec; note IS-SD break). Responses to two CL pulse trains at 10 Hz are illustrated in the top panel: the left one was preceded by the first wave in a spontaneously (spont.) occurring spindle sequence. Augmenting responses to the two pulse trains applied to CL nucleus are expanded below.
responsible for the development of incremental cortical responses. We then supposed that thalamic stimulation will produce more powerful augmenting responses than cortical stimulation, because, in the former case, GABAergic RE neurons are set into action more synchronously. This would induce stronger hyperpolarizations and rebound spike bursts, which are necessary for the LT-type augmenting in TC cells, with direct consequences for cortical augmentation. The comparison between thalamic and 

Figure 5. LT augmenting responses in thalamocortical cell precede cortical responses to 10 Hz thalamic stimuli. Ketamine–xylazine anesthesia. Two superimposed traces in the top panel depict simultaneous recordings of field potentials from area 4 and dual intracellular recordings from area 4 and VL thalamus. Stimuli applied to the VL nucleus. Below, Expanded responses to the first and fifth stimuli in the train. Inset, Average triggered by the first VL action potential (asterisk) in the second to fifth augmented responses. Small deflections in intracellularly recorded cortical neuron, coincident with action potentials in VL cell (visible in bottom panels), are attributable to capacitive coupling.
Figure 6. During thalamic-evoked augmenting responses, the depolarization area in cortical neuron increases as a function of number of action potentials in the rebound spike bursts of thalamocortical cell. Ketamine–xylazine anesthesia. Top two traces, Dual intracellular recordings from VL and area 4 neurons. Below, Average of second and third responses in thalamic and cortical cells. The area of secondary depolarization (b) in the response of cortical neuron is marked by dots. Left plot, Area of secondary depolarization of cortical cell as a linear function of number (Figure legend continues)
Cortical rhythmic stimulation was investigated in five cell pairs. In each case, augmentation was much stronger when evoked by thalamic stimuli. Figure 9 shows that, in the same cell pair, thalamic stimulation, close to the VL-recorded neuron (1–2 mm apart), produced LT-type augmenting in the TC cell, with progressively increasing number of action potentials in the rebound bursts, preceding augmentation in the area 4 neuron. In contrast, cortical stimulation close to the recorded area 4 neuron (1–2 mm apart) produced smaller-amplitude hyperpolarizations in the TC cell, leading to only one rebound spike burst at the end of the pulse train. Cortical stimuli triggered shorter-latency responses in the cortical neuron; however, the difference between the response to the first stimulus and the responses to subsequent stimuli in the pulse train was less dramatic than in the case of thalamic stimulation.

**Differences between various types of cortical neurons during thalamically evoked augmenting responses**

Dual simultaneous recordings from cortical neurons \((n = 27)\) were performed to assess the specificity of augmentation in different areas as a function of the stimulated thalamic nucleus and to detect possible differences between superficially (above 0.8 mm) and deeply (below 0.8 mm) located pyramidal cells (see Fig. 12). Most dual cortical recordings were performed in precruciate area 4 and suprasylvian areas 5–7, where the boundary between the lower part of layer III (area 4) or layer IV (areas 5–7) and the
Figure 8. Spontaneously occurring spindle oscillation and thalamically evoked augmenting responses in dual intracellular recordings from VL thalamus and area 4. Barbiturate anesthesia. Parts marked by horizontal lines are expanded at right (arrows). Note LT rebound spikes in TC cell preceding the spindle-related depolarization in cortical cell, during both spontaneous and postaugmenting spindles.
Figure 9. Differences between thalamically and cortically evoked augmenting responses in dual simultaneous recordings of thalamic (VL) and cortical (area 4) neurons. Ketamine–xylazine anesthesia. A, VL-evoked augmenting responses. At right, expanded responses to first four stimuli. Inset, Averaged responses (n = 5). B, Same neurons, but stimuli applied to area 4.
upper part of layer V is at ~0.8 mm below the surface of the cortex [Hassler and Muhs-Clement (1964), their Figs. 13, 19, 30, 31].

Simultaneous intracellular recordings from cortical areas 5 and 4 (Fig. 10) revealed that augmenting responses were largely restricted to area 5 when stimulating the thalamic LP nucleus (which projects heavily to that suprasylvian area), whereas the area 4 neuron showed negligible, if any, augmentation. The area 5 neuron displayed a depolarizing response of increased amplitude during the hyperpolarization elicited by thalamic LP stimulation. This aspect is not usually observed in other cortical neurons, which generally showed augmenting responses over a depolarizing envelope (Figs. 3–8; also see the sustained hyperpolarization in Fig. 5, similar to that in Fig. 10). In the case of LP-evoked responses in area 5 neuron, the LT response deinnactivated by membrane hyperpolarization was preceded by a small depolarizing response, probably an EPSP of thalamic origin (Fig. 10, bottom panel with averaged responses). The VL stimulation elicited augmentation in the intrinsically bursting cell recorded from area 4, in contrast with no sign of augmentation observed in area 5 (see averaged responses).

Figure 10. Specificity of augmenting responses in different thalamocortical systems: LP-evoked augmenting in area 5 and VL-evoked augmenting in area 4. Dual intracellular recordings from neurons in areas 5 (regular-spiking) and 4 (intrinsically bursting). Ketamine–xylazine anesthesia. Note that LP-evoked augmenting in area 5 developed as LT-type responses during hyperpolarization, similarly to LT augmentation in TC cells (see Discussion).
Figure 11. Comparison between LP-evoked responses in two, simultaneously recorded, neurons from area 7: Intra 1 at a depth of 0.65 mm and Intra 2 at 0.9 mm (see morphological features of these intracellularly stained pyramidal neurons in Fig. 12). The average of 10 responses is shown below at two time scales to compare the latencies of excitatory responses. Note shorter latency of the secondary depolarization, characterizing augmenting responses, in the deeply lying pyramidal cell.
We analyzed the degree of augmentation in cortical neurons that were simultaneously recorded from superficial (above 0.8 mm) and deep (below 0.8 mm) layers. The typical example shown in Figure 11 illustrates two adjacent pyramidal neurons from area 7 that were stained (Fig. 12) and found to be located at depths of 0.65 mm (Fig. 11, *Intra 1*) and 0.9 mm (Fig. 11, *Intra 2*), respectively. Although both neurons displayed augmenting responses to stimulation of the thalamic LP nucleus, the secondary depolariz-
ing responses had a shorter latency in the deeply lying cell (∼10 msec), compared with the latency of the same component (16–18 msec) in the more superficial neuron. Precursor activity in the deeply lying cell was also observed in the self-sustained, postaugmenting rhythmic waves.

Finally, we compared the augmenting responses in a sample of regular-spiking cells (n = 5) and fast rhythmic-bursting cells (n = 5) from superficial layers II–IV and deep layers V–VI. Fast rhythmic-bursting cells were identified by responses to depolarizing current pulses (for details, see Steriade et al., 1998). Figure 13 illustrates superimpositions of thalamically evoked augmenting responses in two neurons of each type, recorded more superficially than 0.8 mm and deeper than 0.8 mm. Deeply lying regularly-spiking as well as fast rhythmic-bursting neurons displayed stronger augmenting responses, with shorter latencies, than more superficially located neurons.

DISCUSSION

We report four major findings: (1) thalamically evoked augmenting responses of cortical neurons mainly result from the selective enhancement of a secondary depolarization, whereas the early excitation is simultaneously reduced in a majority (>75%) of tested neurons; (2) deeply lying (below 0.8 mm) regular-spiking and fast rhythmic-bursting neurons display a greater propensity and shorter latency for augmenting responses than neurons of both categories located more superficially; (3) the augmented secondary depolarization of cortical neurons is preceded by LT-type rebound spike bursts in simultaneously recorded TC cells; and (4) postaugmenting, self-sustained oscillations, within the same frequency range as responses to pulse trains (10 Hz) applied to thalamic nuclei, persist for longer periods in cortical than in TC cells.

Multiple mechanisms underlying cortical augmenting responses

In the majority of cortical neurons, thalamically elicited augmentation resulted from a selective increase of the secondary depolarization associated with a reduction in the early EPSP (Figs. 2, 3). This aspect is the intracellular counterpart of earlier local field potential data and extracellular unit recordings from somatosensory [Steriade and Morin, 1981], their Fig. 6] and suprasylvian association [Steriade, 1991], his Fig. 4] cortices. In those studies, the probability of single action potentials and, relatedly, the amplitude of the early depth negativity of field potentials evoked by appropriate thalamic stimuli were reduced or abolished at the second stimulus at 10 Hz, whereas the secondary depth negativity was simultaneously enhanced and accompanied by spike trains at high frequencies. The diminution in amplitude and associated discharges related to the early cortical EPSP during thalamically evoked augmenting responses may be ascribed to two, nonexclusive mechanisms: (1) a decrease in input resistance of cortical neurons as a result of the action of local GABAergic neurons (I. Timofeev and M. Steriade, unpublished data; also see below); and (2) the IPSPs in TC cells may prevent the triggering of monosynaptically elicited action potentials in many cortically projecting cells. The increase in the cortical secondary depolarization could contribute, in addition to the intrathalamic mechanisms [Steriade and Timofeev, 1997; Timofeev and Steriade, 1998], to the appearance of a secondary depolarization in related TC neurons [Steriade and Deschênes, 1984], their Fig. 13].

It has been suggested (Purpura et al., 1964; Creutzfeldt et al., 1966) that the increased secondary depolarization in cortical neurons results from the attenuation of hyperpolarizing potentials during repetitive stimulation. Typical augmenting responses occur, however, in cortical neurons even when hyperpolarization cannot be detected in response to the first and subsequent stimuli as well as when hyperpolarizing potentials are not diminished or may even increase (Figs. 5, 10, 11). Thus, factors other than the reduction in hyperpolarization may account for the increased secondary depolarization of augmented responses evoked by thalamic stimulation. There are at least three such factors: augmenting-related postinhibitory spike bursts in TC cells, intrinsic membrane properties of cortical neurons that are uncovered by hyperpolarization, and particular types of cortical neurons that exhibit a high propensity for rhythmic, high-frequency spike bursts. These factors are discussed below.

(1) The present data demonstrate that in all simultaneously recorded TC and cortical neurons (n = 32) the first action potential in the LT rebound bursts of TC cells precedes by short latencies (∼3 msec) the augmented depolarization in cortical neurons. This result points to thalamic incremental responses as a major source for cortical augmentation. We considered here only the LT-type augmenting responses that are deinactivated by membrane hyperpolarization. The temporal relation between the other (HT) type of augmenting potentials in TC cells (Steriade and Timofeev, 1997) and target cortical cells remains to be investigated. The proportion of TC neurons generating HT-type augmenting responses in animals with intact thalamocortical loops is much smaller (∼10%) than in decorticated animals [Steriade and Timofeev, 1997]. This is probably attributable to the fact that the corticothalamic feedback drives GABAergic RE neurons, with obvious inhibitory influences on TC cells and prevalent appearance of LT-type augmenting responses. The role of TC-cell LT rebound bursts in priming augmenting cortical responses is also suggested by the facilitation of cortical augmentation when preceded by spontaneous spindles (Fig. 4). Compared with control epochs, the spindles are associated with powerful IPSPs in TC cells, followed by rebound spike bursts that are transferred to cortex, where they produce the enhancement of the secondary depolarization, typical for augmentation.

(2) In view of incremental responses arising from a hyperpolarized membrane in cortical cells (Figs. 5, 10, 11), we propose that another factor promoting augmentation is the activation of inhibitory cortical neurons driven by thalamic repetitive stimuli. Abundant evidence shows that TC axons contact the different varieties of local circuit inhibitory interneurons (Jones, 1975, 1981). During sleep spindles, the naturally occurring phenomenon mimicked by augmenting responses, the rhythmic volleys from TC cells produce powerful inhibitory effects on cortical pyramidal cells, as revealed by the transformation of reversed IPSPs, with Cl−-filled pipettes, into robust bursts resembling paroxysmal depolarizing shifts during seizures (Contreras et al., 1997a). Thus, the strength of thalamic inputs during spindles, leading to bisynaptic inhibition of pyramidal cells, is more effective than expected solely based on reduced discharge frequencies of pyramidal tract neurons during resting sleep (Evarts, 1964; Steriade et al., 1974). Thalamically induced hyperpolarization of cortical neurons is not exclusively attributable to GABAergic inhibition. Indeed, under in vivo conditions, with significant spontaneous neuronal activity, the GABA_A-mediated monophasic IPSP in neocortical neurons (Pollen and Lux, 1966; Contreras et al., 1997b) shuts off discharges in a large proportion of cells, thus contributing to a disfacilitation phenomenon during which K⁺ currents dominate the membrane behavior (Contreras et al.,
Figure 13. Superimposed responses of superficial (above 0.8 mm) and deep (below 0.8 mm) regular-spiking and fast rhythmic-bursting cells during augmenting responses elicited by thalamic stimulation. In each case, responses from two different cells of each type are superimposed. The high-frequency burst of a fast rhythmic-bursting cell at a depth of 1.2 mm is expanded at right (arrow).
Roles played by the thalamus and neocortex in augmenting responses

Thalamic repetitive stimulation produces stronger augmenting responses in cortical neurons than does cortical stimulation (Fig. 9) because of intrathalamic incremental responses (Steriade and Timofeev, 1997) consisting of repetitive spike bursts that are transferred from TC to target cortical neurons. The inclusion of the thalamus in this circuit explains the differences between cortical augmenting responses in the presence of the thalamus and those recorded in somatosensory cortex after destruction of the corresponding thalamic nuclei (Morin and Steriade, 1981). A decrease in the inhibitory phase after the first thalamic stimulus in a pulse train at 10 Hz can be produced by brainstem reticular activation. Under these experimental conditions, the postinhibitory rebound is produced before the second stimulus is delivered, and the augmenting responses to the second as well as following stimuli are decreased in amplitude or abolished [Steriade and Morin (1981), their Fig. 5]. This result emphasizes the role of postinhibitory rebound bursts in TC cells in the production of cortical augmentation. However, the fact that augmentation also occurs in thalamically lesioned animals and that the shortened latency of rebound bursts similarly leads to diminished augmenting responses together indicate that inhibitory processes in the cerebral cortex also contribute to augmentation (Steriade and Morin, 1981). This conclusion is congruent with recent intracellular data (Castro-Alamancos and Connors, 1996b) and modeling studies (Bazhenov et al., 1998b).

The possible contribution of intracortical circuits to the production of augmenting responses is further strengthened by the presence of self-sustained, postaugmenting oscillatory responses that are longer-lasting in cortex than in related thalamic foci (Fig. 7). This was also apparent in dual intracellular recordings, in which the duration of spontaneously occurring spindle sequences in TC cells was shorter than in cortical cells (Fig. 8). In agreement with field potential studies in rat neocortex (Kandel and Buzsáki, 1997), these observations suggest that complex intracortical circuits have a major influence on the incoming thalamocortical inputs and can amplify oscillatory activity arising in the thalamus. Moreover, cortical responses could lead to the spread of oscillatory activity to remote cortical foci by corticothalamocortical loops (Bazhenov et al., 1998b) that return to areas outside those where the original pathways originate (Kato, 1990). The self-sustained oscillatory activity that follows augmenting responses may lead to paroxysmal activity that is largely generated intracortically. This is a form of cortical short-term plasticity, because the repetition of cortical pulse trains at 10 Hz led in deeply lying intrinsically bursting cells, recorded from the homotopic cortical area in the contralateral hemisphere of athalamic animals, to (1) a progressive depolarization of cortical neurons, (2) an increase in the depolarization area of synaptically evoked responses, (3) an increased number of action potentials to testing stimuli, and (4) self-sustained seizures [Steriade et al. (1993b), their Fig. 14]. The role of intrinsically bursting layer V cells in augmenting responses was also emphasized by Castro-Alamancos and Connors (1996b). The fast rhythmic-bursting cortical neurons recorded from layers V–VI, some of them with identified thalamic projections (Steriade et al., 1998), represent another strong candidate for such processes (Fig. 13). The role of corticothalamic neurons in rhythmic oscillations was also emphasized by Kao and Coulter (1997).

Thalamocortical augmenting responses are modulated by behavioral states of vigilance in naturally sleeping and aroused cats (Steriade et al., 1969) and rats (Castro-Alamancos and Connors, 1996c); they are maximal during epochs in which the animals are still and when they display spontaneous spindle oscillations within the frequency range of evoked responses, whereas they are suppressed during strong arousal. These data, together with the previously mentioned results on the development from augmenting responses to seizures, are consistent with the preferential occurrence of some types of paroxysmal activities during drowsiness and light sleep (Steriade, 1974).

Concluding remarks

Augmenting responses, whose study is relevant for short-term plasticity processes in both normal and pathological states, can be generated in either the thalamus in the absence of the cortex or the cerebral cortex of athalamic animals. However, the preservation of reciprocal corticothalamic circuits in intact-brain preparations ensures, first, the production of postinhibitory rebound spike bursts in TC cells that are transferred to cortex and are followed by secondary depolarizations, which are typical for augmenting responses in cortical cells, and, second, the amplification of thalamofugal volleys by intracortical excitatory and disinhibitory circuits.

REFERENCES


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