

DYNAMIC PROPERTIES OF CORTICOTHALAMIC EXCITATORY POSTSYNAPTIC POTENTIALS AND THALAMIC RETICULAR INHIBITORY POSTSYNAPTIC POTENTIALS IN THALAMOCORTICAL NEURONS OF THE GUINEA-PIG DORSAL LATERAL GENICULATE NUCLEUS

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Abstract—The properties of postsynaptic potentials evoked by stimulation of cortical, retinal and GABAergic thalamic afferents were examined *in vitro* in thalamocortical neurons of the guinea-pig dorsal lateral geniculate nucleus. Brief trains of stimulation (2–10 stimuli) delivered to corticothalamic fibers led to a frequency-dependent increase in excitatory postsynaptic potential amplitude associated with an increase in activation of both *N*-methyl-D-aspartate and non-*N*-methyl-D-aspartate glutamate receptors. In addition, repetitive stimulation of corticothalamic fibers also gave rise to a slow excitatory postsynaptic potential that was blocked by local application of the glutamate metabotropic receptor antagonist α -methyl-4-carboxyphenylglycine. In contrast, repetitive stimulation of optic tract fibers resulted in monosynaptic excitatory postsynaptic potentials that did not potentiate and were not followed by the generation of a slow excitatory postsynaptic potential.

Repetitive activation of the optic radiation also evoked both GABA_A and GABA_B receptor-mediated inhibitory postsynaptic potentials. These inhibitory postsynaptic potentials exhibited frequency-dependent depression during repetitive activation. The presence of frequency-dependent facilitation of corticothalamic excitatory postsynaptic potentials and frequency-dependent decrement of inhibitory postsynaptic potentials, as well as the ability of corticothalamic fibers to activate glutamate metabotropic receptors, suggests that sustained activation of corticothalamic afferents *in vivo* may result in postsynaptic responses in thalamocortical cells that are initially dominated by GABAergic inhibitory postsynaptic potentials followed by prominent monosynaptic excitatory postsynaptic potentials as well as a slow depolarization of the membrane potential. Therefore, the corticothalamic system may inhibit or enhance the excitability and responsiveness of thalamocortical neurons, based both on the spatial and temporal features of thalamocortical interactions. © 1999 IBRO. Published by Elsevier Science Ltd.

Key words: facilitation, depression, metabotropic glutamate receptor.

The processing and transmission of visual information through the dorsal lateral geniculate nucleus (LGNd) is under the influence of many different aspects of the neuronal circuit in which these cells are embedded, including the electrophysiological properties of each cell type, the properties of synapses and synaptic networks, as well as the status of various modulatory neurotransmitters (reviewed

in Refs 31, 49 and 52). One prominent feature of the dorsal lateral geniculate nucleus, and indeed the thalamus in general, is the presence of a massive axonal innervation arising from layer VI pyramidal cells in the cerebral cortex (reviewed in Refs 22 and 49). These axons densely innervate thalamocortical neurons in all layers of the LGNd as well as GABAergic neurons, both local and in the thalamic reticular nucleus.^{35,45,46}

Previous electrophysiological investigations of the corticothalamic pathway have demonstrated that these axons exhibit a strong monosynaptic excitatory influence on thalamocortical, thalamic reticular, and local GABAergic interneurons.^{1,2,10,24,26} The ability of corticothalamic inputs to strongly activate GABAergic neurons within the thalamus also results in the generation of robust di-synaptic inhibitory postsynaptic potentials (IPSPs) in thalamocortical

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Abbreviations: 1S,3R-ACPD, 1S,3R-1-amino-1,3-cyclopentane-dicarboxylic acid; ACSF, artificial cerebrospinal fluid; AMPA, α -amino-3-hydroxy-5-methylisoxazole-4-propionate; APV, 2-amino-5-phosphonovaleric acid; CNQX, 6-cyano-7-nitroquinoxaline-2,3-dione; EPSP, excitatory postsynaptic potential; IPSP, inhibitory postsynaptic potential; LGNd, dorsal lateral geniculate; α -MCPG, (\pm) α -methyl-4-carboxyphenylglycine; NRT, thalamic reticular nucleus; OT, optic tract; PGN, perigeniculate nucleus.

cells, giving the corticothalamic system the ability to either directly excite, or indirectly inhibit, the output of the thalamus.

Previously it has been suggested that the spatial organization of excitatory and inhibitory connections may be important in determining whether or not the cortex exerts an overall excitatory or inhibitory influence on thalamocortical activity.^{37,56} For example, the experiments of Tsumoto *et al.*⁵⁶ demonstrated with cross-correlation techniques that activity in cortical neurons may be followed by either excitatory or inhibitory influences in cells of the LGNd. Another important variable in determining the influence of the descending corticothalamic influence is the temporal pattern of activation of these fibers and the effects of this on excitatory and inhibitory neurotransmission. Repetitive activation of the corticothalamic pathway results in a marked frequency-dependent facilitation in the excitatory influence,^{10,24,26} as well as the generation of a slow excitatory postsynaptic potential (EPSP) that follows the monosynaptic fast EPSPs and that may be several seconds in duration.³⁴ In other neuronal systems, frequency-dependent facilitation has been demonstrated to result from both increases in transmitter release by the excitatory terminals (reviewed in Refs 15, 28 and 62) as well as decreases in the release of GABA from di-synaptically activated inhibitory neurons.^{8,9,40} Indeed, dual intracellular recordings from pairs of GABAergic perigeniculate and thalamocortical neurons have revealed that repetitive activation of perigeniculate nucleus (PGN) to thalamocortical GABAergic synapses results in frequency-dependent depression.^{25a}

In hippocampal pyramidal cells, frequency-dependent depression of GABAergic transmission has been demonstrated to result from the activation of presynaptic GABA_B receptors through the release of GABA itself (reviewed in Ref. 54). Present evidence suggests that this presynaptic action of GABA may arise through inhibition of Ca²⁺ currents in the presynaptic membrane.^{27,61} Similarly, in the mammalian thalamus the activation of GABA_B receptors also potently inhibits the function of GABAergic terminals, apparently through a presynaptic mechanism.^{47,57}

The generation of a prolonged slow EPSP following the repetitive activation of corticothalamic fibers has important functional consequences for the influence of this pathway on the generation of thalamocortical activity. Thalamocortical neurons depolarize by 10–20 mV in the transition from slow wave sleep to waking or rapid eye movement sleep.²⁰ Previous investigations have suggested that the activation of brainstem and hypothalamic systems may underlie this depolarization of thalamocortical neurons (reviewed in Refs 31, 51 and 52). However, the ability of the activation of corticothalamic fibers to generate a slow EPSP that is identical to that associated with activation of these

brainstem inputs suggests that the activation of the corticothalamic pathway may also underlie the depolarization of thalamocortical cells with arousal.³⁴ Previously, we have suggested that the corticothalamic slow EPSP may be mediated through the activation of glutamate metabotropic receptors, since the activation of these receptors with a selective agonist generated a similar response³⁴ and the corticothalamic pathway is believed to be at least in part glutamatergic.³⁴ However, pharmacological examination of this possibility with antagonists to glutamate metabotropic receptors remained to be performed. Portions of this work have appeared in abstract form.⁵⁹

EXPERIMENTAL PROCEDURES

Male and female guinea-pigs (Hartley; $n=56$) were deeply anaesthetized with sodium pentobarbital (35 mg/kg i.p.) and killed by decapitation, as previously described.^{32,33} The thalamus was rapidly dissected and was then sectioned parasagittally at 400 μm in the plane of either the corticothalamic or the optic tract fibers using a Vibratome (Pelco). Slices were maintained in an interface chamber (Fine Science Tools) at $35.5 \pm 1^\circ\text{C}$ and bathed with an artificial cerebrospinal fluid solution (ACSF) containing 126 mM NaCl, 2.5 mM KCl, 1.2 or 2 mM MgSO₄, 26 mM NaHCO₃, 1.25 mM NaH₂PO₄, 2 mM CaCl₂ and 10 mM glucose, saturated with 95% O₂/5% CO₂ to a final pH of 7.4. Pharmacological antagonists were applied either in the bathing medium or locally with the "pico-drop" method in volumes of 5–20 μl . The glutamate receptor antagonists D- or DL-2-amino-5-phosphonovaleric acid (APV, 25–50 μM ; Sigma) and 6-cyano-7-nitroquinoxilane-2,3-dione (CNQX, 10–25 μM ; Research Biochemicals) were used to block, respectively, N-methyl-D-aspartate (NMDA) and non-NMDA receptors. GABA receptor antagonists bicuculline methiodide (30–500 μM , Research Biochemicals), picrotoxin (100 μM) and 2-Hydroxysaclofen (0.5–2 mM, Research Biochemicals) were used to block GABA_A or GABA_B receptors, respectively.

Electrical stimulation (10–400 μA , 50–100 μs duration) was generated using a constant current generator (World Precision Instruments) and passed through a concentric bipolar stimulating electrode placed in either the corticothalamic or optic tracts. The stimulus trains were generated using a multichannel programmable pulse generator (Master-8, A.M.P.I.). No effort was made to remove the thalamic reticular nucleus (NRT) from the slice preparation.

Intracellular recording microelectrodes were made from thin-walled glass capillary tubes (1 mm o.d., 0.67 mm o.d., World Precision Instruments) and pulled on a Brown-Flaming micropipette puller (Sutter Instruments). The micropipettes were filled with either 4 M potassium acetate or 4 M potassium acetate and 50 mM lidocaine N-ethyl bromide quaternary salt (QX-314, Research Biochemicals Inc.), and had final tip resistances of 40–65 M Ω . Intracellular recordings were performed using an Axoclamp-2A amplifier (Axon Instruments).

The use of QX-314

QX-314 is a quaternary derivative of lidocaine bromide salt which blocks voltage-dependent sodium channels when included in the intracellular recording electrode solution.³⁹ In experiments without QX-314, EPSPs activated action potentials at depolarized potentials and low-threshold Ca²⁺ spikes and bursts of action potentials at more hyperpolarized membrane potentials. Therefore to examine facilitation of EPSPs without the confounding influence of the

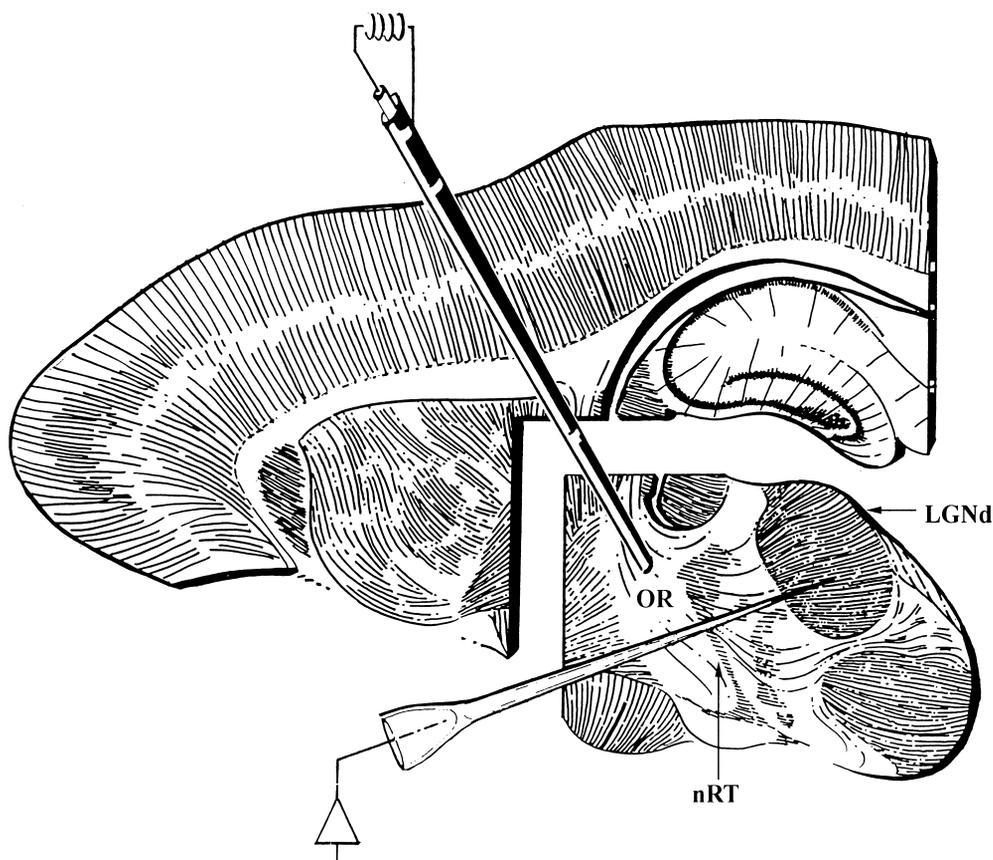


Fig. 1. A schematic summary diagram illustrating the parasagittal orientation of the thalamus and dorsal lateral geniculate (LGNd) used in this study. The stimulating electrode is located in the cortical white matter and was located at least 1–2 mm outside of the LGNd. Figure drawn by Thierry Bal.

activation of Na^+ and low-threshold Ca^{2+} channels, we routinely included QX-314 in the recording micropipette and depolarized the thalamocortical cells with the intracellular injection of current to membrane potentials where the low threshold Ca^{2+} current is inactivated (i.e. positive to -65 mV). QX-314 was allowed to diffuse into the cell with either no holding current or with a slight depolarizing current (0.1–0.5 nA). The effective block by QX-314 took place over a period of 10–20 min and was judged to have occurred when an action potential could not be elicited by depolarizing current injections, in response to EPSPs evoked by corticothalamic stimulation, or upon activation of a rebound low threshold Ca^{2+} spike through the injection of a hyperpolarizing current pulse. In addition to blocking the voltage-dependent sodium channels, QX-314 also prevents GABA_B receptor activation of potassium currents in the recorded cell.³⁹

Isolation and stimulation of fiber tracts

The path taken by corticothalamic fibers between the visual cortex and the LGNd has been described in detail in both rat⁶⁰ and cat.^{45,46} The course of the guinea-pig corticothalamic projection from the primary visual cortex to the LGNd was determined here using biocytin as an anterograde tract tracer.²¹ From these tracing experiments it was determined that the corticothalamic fibers coursed anteriorly and then in a posterior and lateral direction from the visual cortex to the LGNd. Owing to a dominant anterior to posterior direction at the level of the thalamus, parasagittal sections were determined to be the optimal

plane for the formation of geniculate slices for stimulation of corticothalamic fibers.

Sections in which the optic tract (OT) was stimulated were cut parallel to the plane of the OT as it travels along the ventrolateral and lateral margin of the diencephalon. Stimulation of the OT was made at least 1 mm outside the lateral geniculate whereas stimulation of the corticothalamic fibers was made 1–2 mm anterior to the border of the LGNd (see Fig. 1). Stimulus intensity was typically set at one and a half to twice the threshold required to resolve an EPSP in the second or third stimulus in a train of five stimuli. The stimulus intensity ranged from 60–400 μA for the activation of corticothalamic afferents within the optic radiation and 10–100 μA for the activation of OT axons. Increasing stimulus intensity above threshold lead to a progressive increase in amplitude of the evoked corticothalamic EPSP, whereas increasing stimulus intensity in the OT led to clear “jumps” in EPSP amplitude, presumably corresponding to the activation of individual fibers within the optic tract.⁴²

RESULTS

Optic radiation stimulation evokes both excitatory and inhibitory postsynaptic potentials

Intracellular recordings were obtained from 60 thalamocortical neurons in the LGNd. A representative sample of 15 of these cells exhibited an average input resistance of $69 \pm 12 \text{ M}\Omega$, resting membrane potential of -65 mV ($\pm 3.3 \text{ mV}$) and were capable

Cortical-tract Stimulation

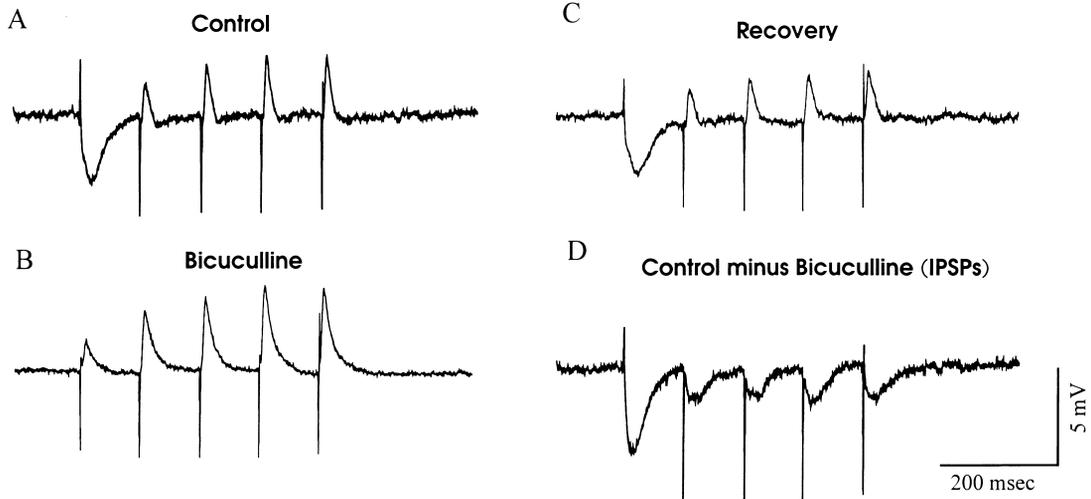


Fig. 2. Repetitive stimulation of optic radiation evokes both EPSPs and IPSPs (A; Control). Application of the GABA_A receptor antagonist bicuculline methiodide (B; 50–100 μ M; Bicuculline) reversibly (C; Recovery) blocks IPSP activity and reveals that corticothalamic EPSPs are facilitated by repetitive stimulation. The difference between the control and the bicuculline conditions reveals that evoked IPSPs undergo depression during repeated activation (D; Control minus bicuculline). In these experiments action potentials and postsynaptic GABA_B IPSPs were prevented by including 50 mM QX-314 in the recording electrode. Membrane potentials were -53 ± 2 mV.

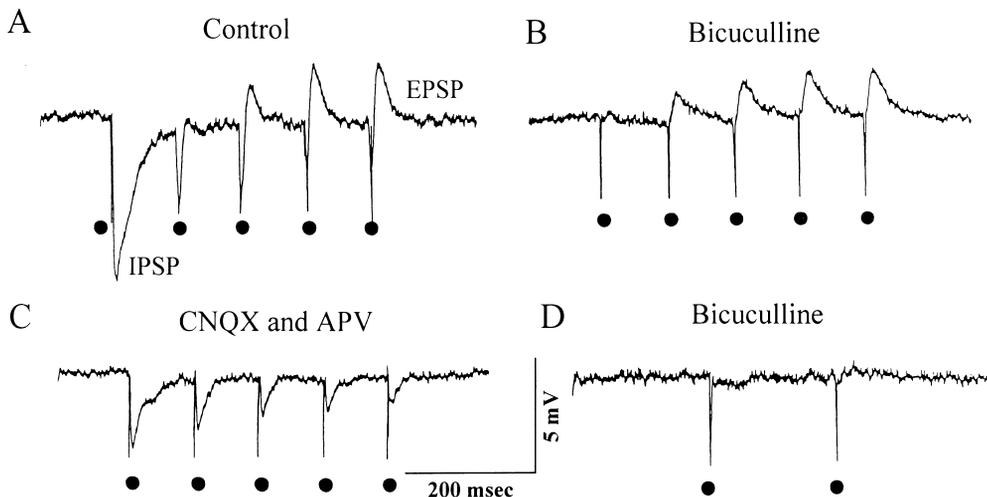


Fig. 3. Repetitive activation modulates the amplitude of both EPSPs and IPSPs. As in the previous figure repeated activation of the optic radiation or the corticothalamic white matter evoked both EPSPs and IPSPs (A, Control). Application of bicuculline methiodide (50–100 μ M) blocked the IPSPs. The isolated corticothalamic EPSPs were facilitated during repeated activation. Thus, corticothalamic EPSP facilitation does not require the modulation of IPSPs (B, Bicuculline). After washout of bicuculline methiodide, application of glutamate antagonists CNQX (20 μ M) and APV (20 μ M) blocked the evoked corticothalamic EPSPs. Isolated GABA_A IPSPs were depressed by repeated activation (C, CNQX and APV). These IPSPs were abolished by the local application of the GABA_A antagonist bicuculline (500 μ M in the micropipette; D). Membrane potential was -54 ± 1 mV and the recording electrode contained 50 mM QX-314.

of generating repetitive trains of action potentials upon depolarization.

Repetitive stimulation at 10 Hz of fibers lying in the optic radiation approximately 0.5 to 2 mm anterior to the LGNd (see Fig. 1) resulted in both IPSPs and EPSPs in thalamocortical neurons in normal bathing solution and recorded with microelectrodes

containing either 4 M potassium acetate or 50 mM QX-314 dissolved in 4 M potassium acetate (Figs 2A and 3A). Initially, the evoked postsynaptic responses were dominated by the presence of a large IPSP, although this quickly gave way to prominent EPSPs during the stimulus train (Figs 2A and Fig. 3A). The EPSPs presumably arise from the

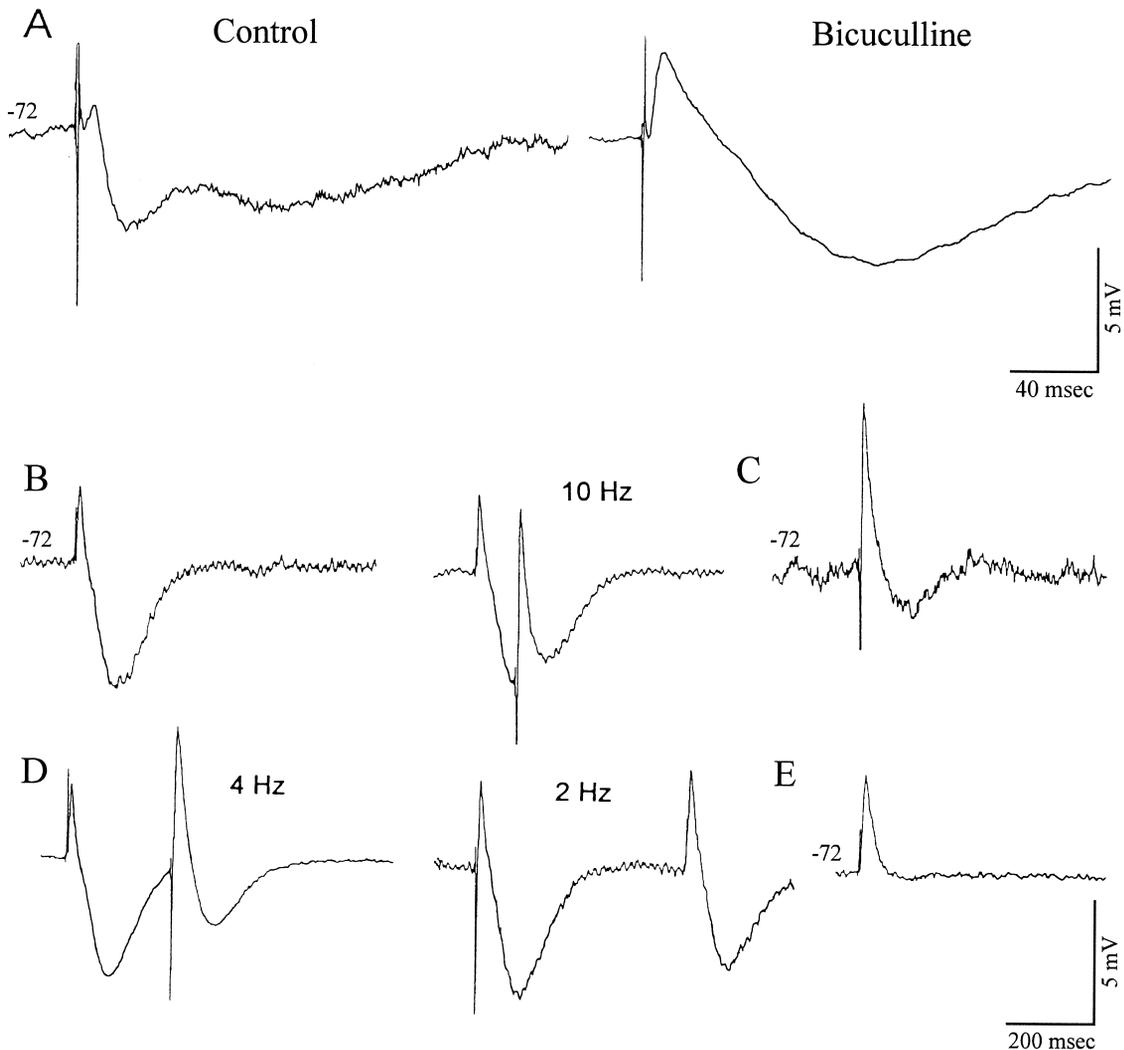


Fig. 4. GABA_B receptor-mediated IPSPs exhibit depression. Stimulation of the optic radiation evokes both EPSPs and IPSPs (Control A,B). At higher temporal resolution the IPSPs appear biphasic (Control, A). Application of the GABA_A antagonist bicuculline methiodide abolished the early component of the IPSP (A, Bicuculline) but did not diminish the slower second component of the IPSP. (B) The slow, bicuculline resistant, IPSPs showed frequency-dependent depression. Comparison of the second of the two evoked IPSPs with either the first IPSP or a single IPSP (first trace in B) demonstrates that the slow IPSP is depressed by repeated activation. (C) The subtraction of a 10 Hz paired IPSP from a single IPSP (first trace in B) also demonstrates prominent depression. (D) The depression at 10 Hz was greater than that observed at the lower stimulation rates of 2 and 4 Hz. (E) With bicuculline present, the local application of the GABA_B antagonist 2-hydroxysaclofen (1 mM) abolished the slow IPSP. Thus the slower second component of the IPSP is mediated by GABA_B receptor activation. Recordings were made with 4 M potassium acetate in the recording electrodes. The resting membrane potentials were -72 ± 1 mV (A–E). Bicuculline methiodide was present for all except A, control.

activation of corticothalamic fibers that form monosynaptic connections with the intracellularly recorded thalamocortical cell, while the IPSPs may arise from either the activation of neurons or axons of the thalamic reticular nucleus that innervate the recorded neuron or through the excitation of local GABAergic LGNd interneurons owing to the activation of the corticothalamic input.

The changes in amplitude and time-course of the evoked IPSPs and EPSPs occurring during the response to the stimulus train suggested that there was both a frequency-dependent decrease in IPSP

amplitude and a frequency-dependent increase in corticothalamic EPSP amplitude (Fig. 2, control). The following experiments were performed to examine this hypothesis.

Optic radiation-evoked excitatory postsynaptic potentials strongly facilitate in the absence of inhibitory postsynaptic potentials

The block of GABA_A and GABA_B receptors revealed isolated EPSPs in response to stimulation of the optic radiation (Fig. 2B, bicuculline). GABA_A

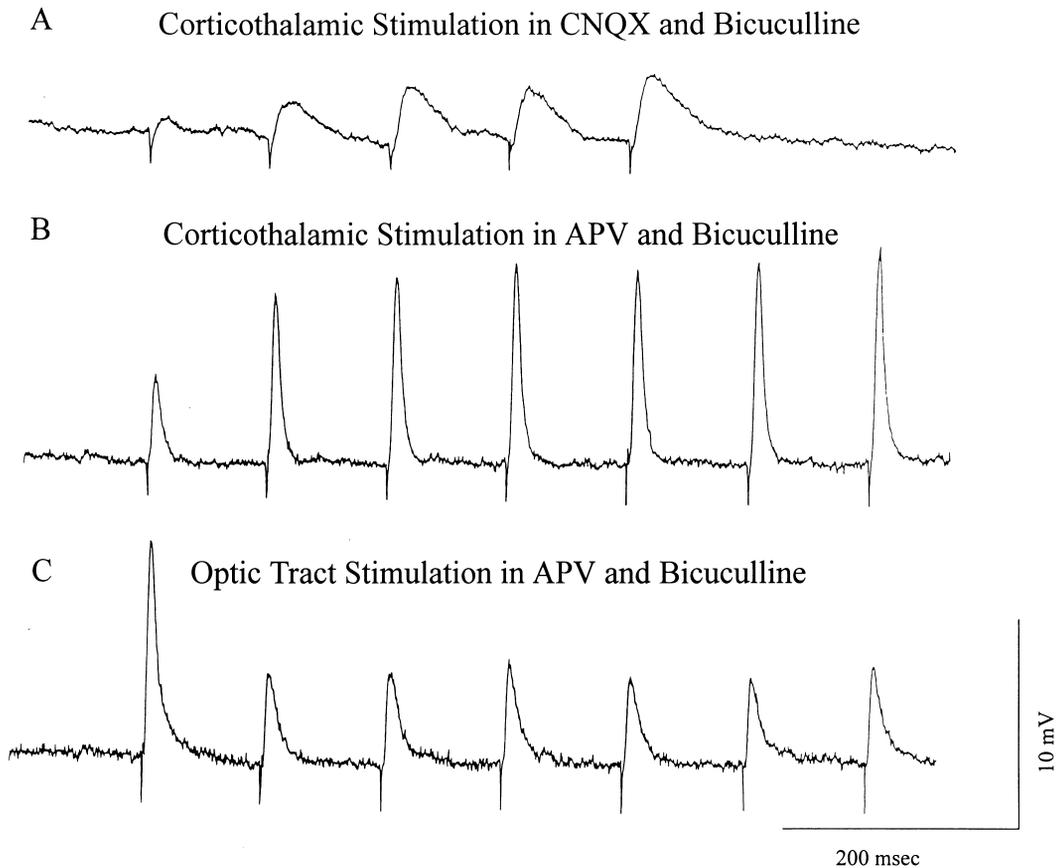


Fig. 5. Repetitive activation of the corticothalamic fibers revealed that both NMDA (A) and non-NMDA (B) receptors mediate frequency-dependent facilitation. In contrast, optic tract-evoked EPSPs undergo depression in response to repetitive activation (C). EPSPs were isolated from IPSPs by including 50 mM QX-314 in the recording electrode and bicuculline methiodide (50–100 μ M) in the ACSF. (A) NMDA EPSPs were isolated by including CNQX (12–25 μ M) in the bathing medium and excluding Mg^{2+} . These NMDA receptor-mediated EPSPs exhibited marked frequency-dependent facilitation. (B) Isolation of AMPA/kainate receptor-mediated EPSPs by including APV (100–500 μ M) and bicuculline (50–100 μ M) in the bathing medium. As with the NMDA receptor-mediated EPSPs, excitatory PSPs mediated by AMPA/kainate receptors also exhibit facilitation. (C) Repetitive stimulation of optic tract axons results in a decrease in EPSP amplitude between the first and second evoked responses, followed by a relatively maintained amplitude at this frequency. The membrane potentials were -61 ± 1 mV.

receptor-mediated IPSPs were blocked by either local (500 μ M in micropipette) or bath (30–100 μ M) application of the GABA_A receptor antagonist bicuculline methiodide ($n = 46$). GABA_B receptor-mediated IPSPs were blocked either through the inclusion of 50 mM QX-314 in the recording electrode (Fig. 2; $n = 35$) or through the bath application of 2-hydroxysaclofen (1–2 mM; $n = 10$). The EPSPs evoked by repetitive stimulation of the optic radiation (following the block of GABA_A and GABA_B receptor-mediated IPSPs) exhibited strong facilitation (80–500%; $n = 15$; Fig. 2B, bicuculline). Examining the difference between the postsynaptic responses before and after the local application of bicuculline revealed that the evoked IPSPs show marked depression during repetitive stimulation, particularly between the first and second IPSPs (Fig. 2D, Control minus bicuculline). In order to directly test this hypothesis,

EPSPs were blocked with the application of CNQX (250 μ M in micropipette; 12–25 μ M in bath) and APV (100–500 μ M in micropipette; 20–50 μ M in bath; Fig. 3C). IPSPs, evoked in this circumstance (with direct electrical stimulation placed in the NRT anterior to the LGNd) exhibited marked depression (30–70%) during repetitive stimulation (Fig. 3C; $n = 21$). These IPSPs were mediated largely by GABA_A receptors, since they were blocked by the local application of bicuculline (Fig. 3D).

The activation of corticothalamic fibers in normal solution could also result in the activation of fast EPSPs followed by both fast GABA_A receptor and slow GABA_B receptor-mediated IPSPs in neurons recorded with 3 M KAC-containing microelectrodes (Fig. 4A, control). Local application of bicuculline resulted in block of the fast component, resulting in an enhancement of both the fast EPSP and the slow IPSP (Fig. 4A, bicuculline).⁷ Paired stimulation of

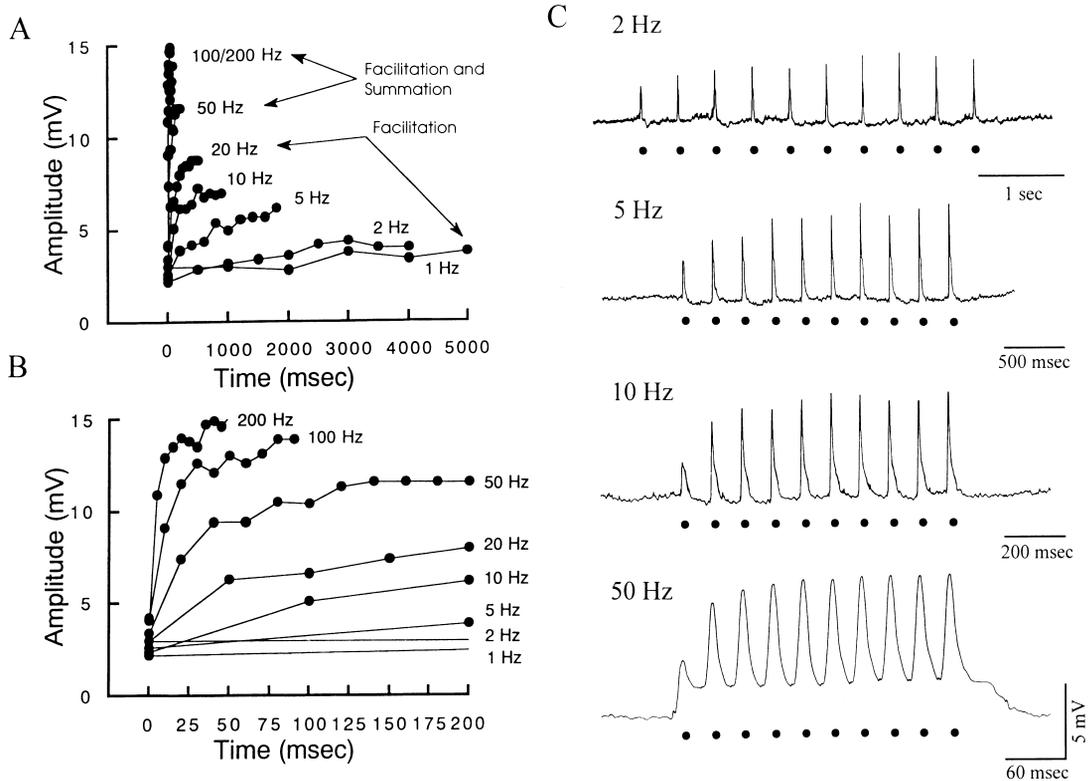


Fig. 6. Frequency dependence of facilitation and summation. Corticothalamic fibers were activated at different rates of stimulation (1–200 Hz). A, B and C show these different frequencies expanded at different time scales. Graphs A and B show that even at frequencies as low as 1 Hz corticothalamic EPSPs facilitate and this facilitation increases markedly with increases in the frequency of stimulation. At stimulation rates above 20 Hz both facilitation and temporal summation occur (C, 50 Hz). These EPSPs were isolated from IPSPs through the inclusion of QX-314 (50 mM) in the intracellular micropipette and the local application of bicuculline (0.5–1 mM in micropipette). Membrane potential for this cell was -56 ± 2 mV.

the corticothalamic inputs revealed a marked potentiation of the EPSP evoked by the second stimulus, and a decrement in the amplitude of the second slow IPSP, even if there was little or no overlap in the evoked PSP sequences (e.g., Fig. 4D, 2 and 4 Hz). Local application of the GABA_B antagonist 2-hydroxysaclofen (1 mM in bath) blocked the bicuculline-insensitive IPSP (Fig. 4E), demonstrating that this event was a GABA_B receptor-mediated IPSP.

Excitatory postsynaptic potentials mediated by either N-methyl-D-aspartate or α -amino-3-hydroxy-5-methylisoxazole-4-propionate/kainate glutamate receptors exhibit facilitation

Repetitive stimulation of optic radiation fibers in the presence of the α -amino-3-hydroxy-5-methylisoxazole-4-propionate (AMPA)/kainate receptor antagonist CNQX (25 μ M) and following the removal of Mg²⁺ from the bathing medium (Fig. 5A; $n = 12$) revealed EPSPs, presumably mediated by NMDA receptors, that exhibited strong facilitation (66–290%). Similarly, repetitive activation of EPSPs by stimulation of the optic radiation following the bath application of the NMDA receptor

antagonist D-APV/DL-APV (25–50 μ M, $n = 18$; normal Mg²⁺) also revealed EPSPs that exhibit marked facilitation (78–505%; Fig. 5B). Application of both CNQX and APV abolished these fast EPSPs (not shown; see Fig. 3C).

Repetitive stimulation of the OT in the presence of bicuculline resulted in EPSPs that did not exhibit facilitation, and in fact, typically exhibited a reduction in amplitude from the first to second EPSP (57–88%; Fig. 5C).⁴¹ Similar results were obtained even when NMDA receptors were not blocked ($n = 4$). In normal bathing medium, stimulation of OT fibers resulted in typical EPSP–IPSP sequences (see Ref. 7). Repetitive stimulation of these fibers resulted in depression of both the EPSP and IPSP amplitude (not shown).

Frequency dependence and time-course of facilitation

The facilitation of isolated monosynaptic corticothalamic EPSPs is frequency dependent. The delivery of repetitive trains of 10 stimuli to the optic radiation at various frequencies revealed facilitation even at frequencies as low as 1 Hz ($n = 46$; Fig. 6). This facilitation increased markedly with

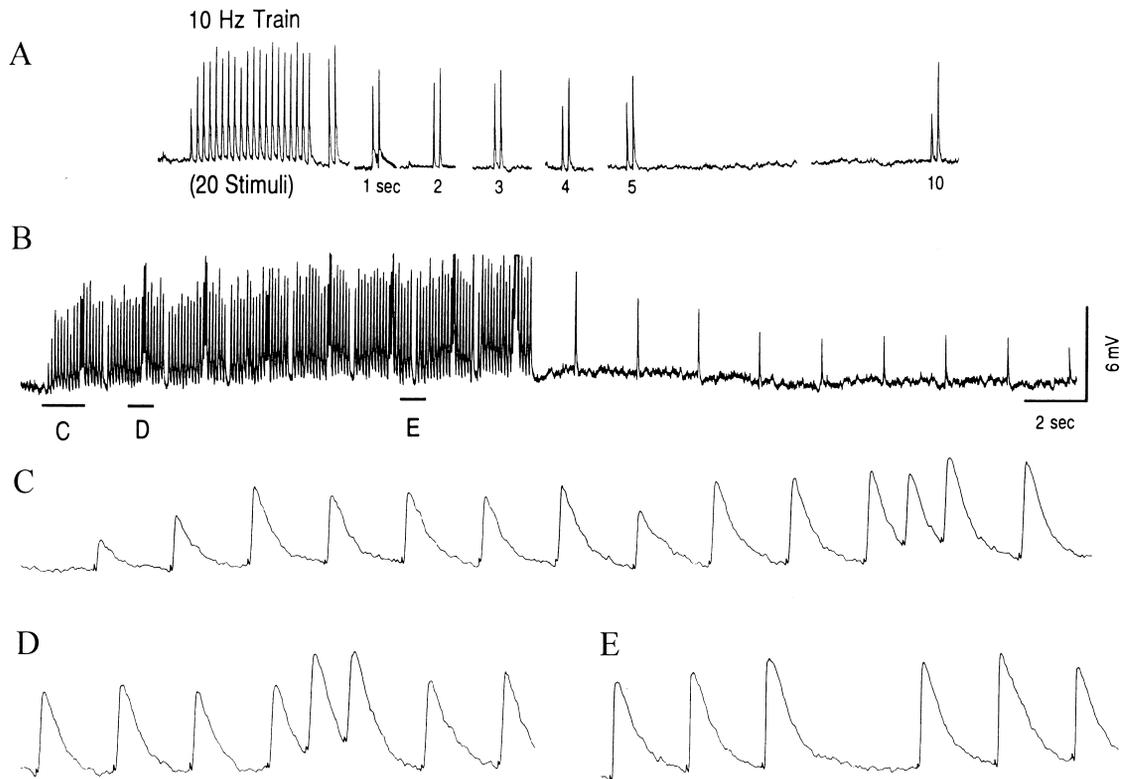


Fig. 7. Facilitation decays with a prolonged time-constant. (A) Following the induction of facilitation with the delivery of 20 stimuli at 10 Hz, a pair of identical stimuli were applied at varying intervals. Facilitation persists for approximately 10 s and therefore decays slowly. (B) Owing to the slow build-up and decay of facilitation, the addition or deletion of single stimuli do not have marked effects on this property (B–E). Corticothalamic EPSPs were isolated by including 50 mM QX-314 in the micropipette and with the local application of bicuculline methiodide (0.5–1 mM in micropipette).

increases in frequencies from 1–20 Hz (Fig. 6A). Increasing the frequency of stimulation to 50 Hz or higher gave rise to temporal summation of EPSPs as well as facilitation of EPSP amplitude (Fig. 6B, C).

The time-course of the persistence of facilitation was examined by applying a train of 20–200 stimuli followed at various intervals by a single test stimulus or a pair of stimuli ($n=9$; Fig. 7). These experiments revealed that facilitation of monosynaptic corticothalamic inputs persists for approximately 10 s (Fig. 7A, B). The slow time-course of the build-up and decrement of facilitation is illustrated by the relative lack of effect of the insertion or deletion of single stimuli in a prolonged train (Fig. 7B–E).

Frequency dependence of GABAergic inhibitory postsynaptic potentials

Previously we have proposed that the strong activation of GABA_B receptors in thalamocortical cells may require the prolonged and high-frequency discharge of GABAergic neurons in the thalamus.^{3,25} Here we examined this hypothesis further through the repetitive stimulation in the region of the thalamic reticular nucleus following the block of

EPSPs with bath application of CNQX (25 μ M) and APV (50 μ M). Under these circumstances, repetitive stimulation at 500 Hz resulted in a fast, presumed GABA_A receptor mediated IPSP followed by a slow, presumed GABA_B receptor-mediated IPSP (Fig. 8A). Applying either one, two, five, 10, 20, or 30 stimuli in the repetitive train resulted in varying degrees of activation of both fast and slow IPSPs in thalamocortical neurons (Fig. 8A, B). With only one electrical stimulus, a prominent fast IPSP is evoked and this is followed by a small slow IPSP (Fig. 8A, B). Increasing the number of stimuli to five results in a substantial enlargement in both the fast and slow IPSPs (Fig. 8A, B). However, increasing the number of stimuli further to 10, 20, or 30 results in a larger enhancement of the slow IPSP over that of the fast IPSP ($n=4$, Fig. 8A, B).

In contrast to the potent effects of 500 Hz stimulation, repetitive stimulation at 50 Hz failed to yield strong activation of slow, presumed GABA_B receptor-mediated IPSPs, even following the delivery of 30 stimuli (Fig. 8C, D). Similarly, the delivery of stimuli at 50 Hz also reduced the amplitude of the evoked fast IPSP in comparison with that obtained in the same neurons following stimulation at 500 Hz (Fig. 8).

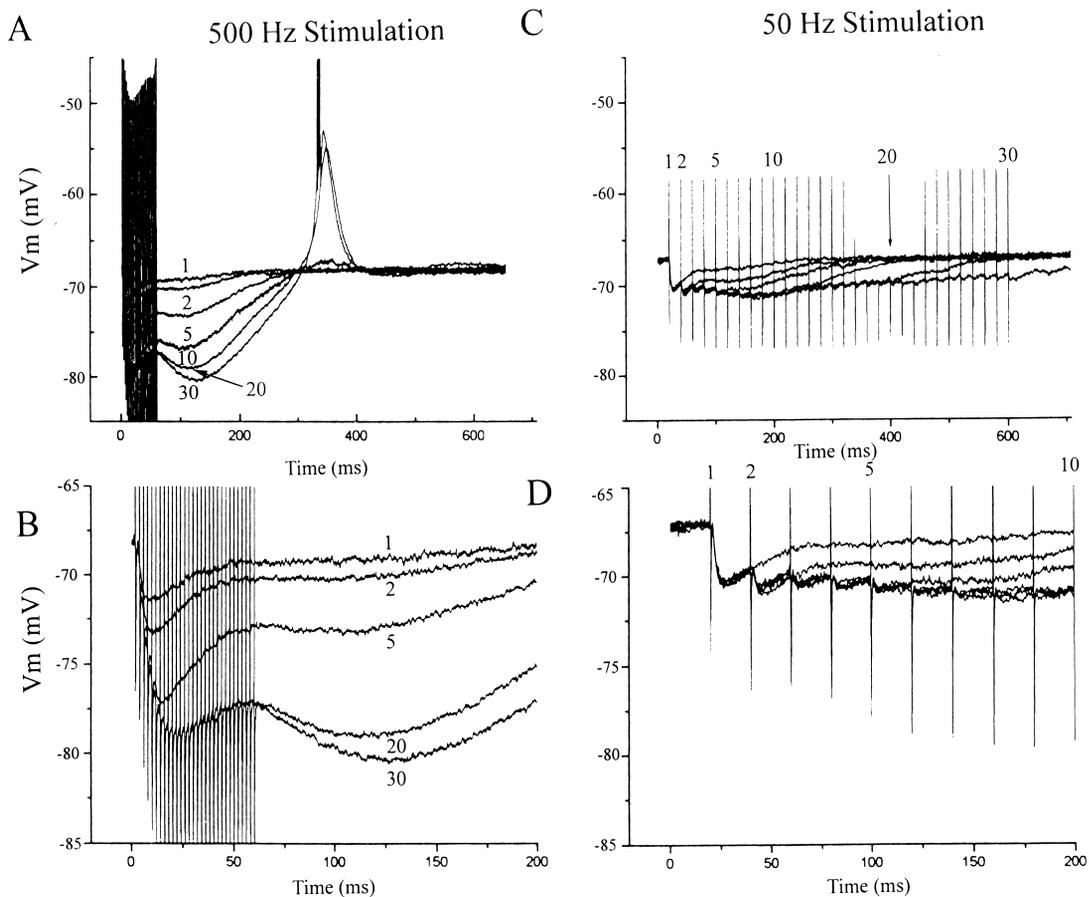


Fig. 8. Frequency of stimulation of NRT inputs to thalamocortical cells determines the degree to which GABA_B receptors are activated. (A) Activation of fibers in the optic radiation at a frequency of 500 Hz results in robust fast and slow IPSPs in thalamocortical neurons (expanded in B for detail). Increasing the number of shocks delivered from one to two, five, 10, 20 and 30 results in an increase in the amplitude of both the early and late IPSPs. (C) In contrast, with stimulation at 50 Hz, increasing the number of stimuli does not markedly increase the peak amplitude of the early or late IPSP (expanded in D for detail).

Optic radiation stimulation evokes a slow excitatory postsynaptic potential through glutamate metabotropic receptors

Repetitive stimulation of the optic radiation could evoke a slow EPSP following the train of fast EPSPs and IPSPs ($n = 40$; Fig. 9A). This slow EPSP was elicited following the bath application of the muscarinic antagonist scopolamine (1 μ M), the α_1 -adrenoceptor antagonist prazosin (1 μ M), the H₁ antagonist diphenhydramine (10 μ M) and the serotonergic antagonist methylsergide (1 μ M). We have previously suggested that this slow EPSP may be mediated by the activation of glutamate metabotropic receptors, since it is occluded by maximal application of the exogenous agonist (1S,3R)-1-amino-1,3-cyclopentanedicarboxylic acid (1S,3R-ACPD).³⁴ Here we tested this hypothesis further through the local application of the glutamate metabotropic receptor antagonist α -methyl-4-carboxyphenylglycine (α -MCPG).¹² Local application of α -MCPG (5–10 mM in micropipette)

resulted in a substantial reduction in the response of thalamocortical neurons to 1S,3R-ACPD ($55.2 \pm 14.8\%$ S.D.; $n = 8$; 5–10 μ M in micropipette) as well as the amplitude of the slow corticothalamic EPSP (Fig. 9A, B; $57.1 \pm 14.0\%$ S.D.; $n = 33$). This effect of α -MCPG was reversible (Fig. 9). Close examination of the fast EPSPs and IPSPs evoked by corticothalamic stimulation reveals little change following the application of α -MCPG (see Fig. 9C).

DISCUSSION

The thalamus and cerebral cortex reciprocally innervate one another in a point-to-point manner that preserves, in the case of the lateral geniculate nucleus, retinotopic order.^{34,56} In the cat, single fibers from the visual cortex densely innervate the A and C layers of the LGNd³⁷ and form synaptic terminals on both thalamocortical and local GABAergic interneurons.^{23,36,45} These fibers also

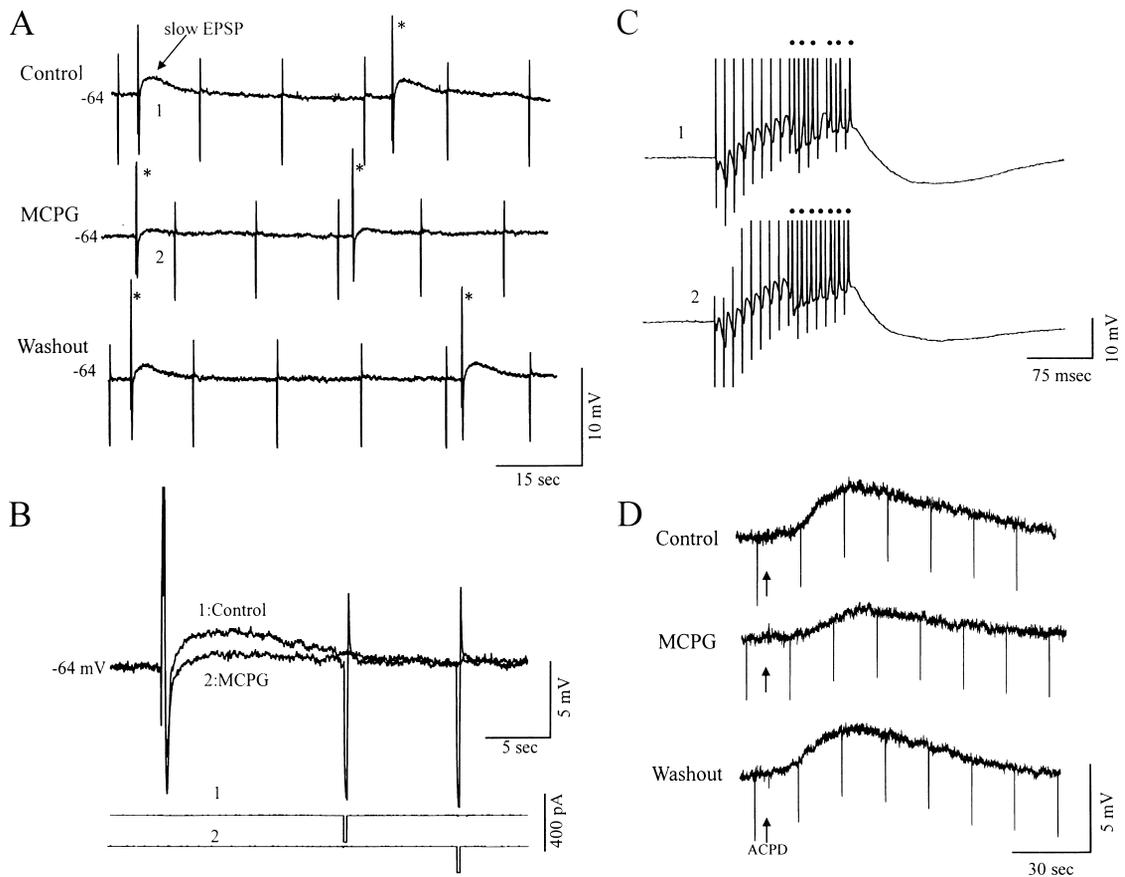


Fig. 9. Repetitive stimulation of the corticothalamic fibers results in a slow EPSP that is reduced by MCPG. (A) Repetitive electrical stimulation (15 stimuli at 100 Hz) results in a slow EPSP that lasts approximately 10 s. Local application of the glutamate metabotropic receptor antagonist MCPG results in a substantial reduction in the amplitude of the slow EPSP. This effect is reversible (washout). (B) Overlay of the slow EPSP before and after application of MCPG. (C) Expansion of the fast EPSPs and action potentials generated in response to the train of electrical stimuli before and after application of MCPG. (D) Local application of MCPG (10 mM in micropipette) results in a substantial reduction in the response of a thalamocortical cell to the local application of ACPD.

give rise to axon collaterals within the PGN and innervate these GABAergic neurons.^{35,37}

Corticothalamic neurons exert a monosynaptic excitatory influence on thalamocortical and perigeniculate/thalamic reticular neurons,^{1,6,10,24,26} perhaps through the release of glutamate or aspartate (reviewed in Ref. 31). Repetitive activation of corticothalamic inputs to thalamocortical cells, as well as to layer IV neurons in the primary visual cortex, results in frequency-dependent facilitation of the postsynaptic excitatory response^{5,10,14,24,26,42,43,59} as well as prolonged (seconds) increases in excitability of thalamocortical cells.^{10,34}

In the present study, we demonstrate that the repetitive stimulation of axons in the optic radiation results in EPSPs and IPSPs in thalamocortical neurons. The EPSPs presumably arise from activation of corticothalamic axons, since axon collaterals between thalamocortical neurons are extremely rare.^{16,37,47} The IPSPs examined in the present study may arise from either the activation of thalamic reticular neurons or their axons (either

directly or indirectly) or through the activation of local GABAergic interneurons within the LGNd.^{7,25} Pharmacologically, the corticothalamic EPSPs are able to activate both non-NMDA and NMDA receptors,^{11,24,48} while the evoked IPSPs are mediated through both GABA_A and GABA_B receptors. Repetitive stimulation of pharmacologically-isolated EPSPs resulted in marked facilitation at stimulation frequencies of 1 Hz or greater and this facilitation decayed with a time-constant of several seconds, a time-course that is most similar to synaptic augmentation.¹⁵ Previously, it has been suggested that facilitation of the corticothalamic pathway may result from the relief of voltage-dependent block of NMDA receptor activation.¹⁰ However, our results indicate that this is not the mechanism of frequency-dependent facilitation of corticothalamic synapses. This facilitation was present even following the pharmacological block of NMDA receptors or the removal of Mg²⁺ from the bathing medium (see Fig. 5). Instead, our results are most consistent with an increase in release of

neurotransmitter from corticogeniculate terminals during facilitation, as has been demonstrated in other neuronal systems (reviewed in Refs 15 and 28). Detailed investigations of the cellular mechanisms of synaptic facilitation have repeatedly demonstrated a crucial role of the dynamics of Ca^{2+} mediated processes in presynaptic terminals (reviewed in Refs 10 and 62).

In addition to facilitation of corticogeniculate synaptic transmission, repetitive stimulation of isolated IPSPs, presumably representing the activation of thalamic reticular axons, resulted in synaptic depression. Similarly, dual intracellular recordings from monosynaptically-coupled pairs of PGN and LGNd neurons have demonstrated that repetitive discharge of PGN neurons results in depression of the postsynaptic IPSP.^{25a} In the hippocampus, depression of GABAergic IPSPs following repetitive stimulation results largely through the presynaptic inhibition of GABA release owing to the activation of GABA_B receptor (reviewed in Ref. 54), which appears to operate mainly through a decrease in Ca^{2+} conductances.^{27,61} Similarly in thalamocortical neurons, activation of presynaptic GABA_B receptors results in a substantial decrease in the release of GABA from NRT terminals^{47,57} and block of GABA_B receptors results in a reduction in paired pulse depression of GABA release.⁵⁷ Another possible mechanism for the depression of IPSPs observed here is the accumulation of Cl^- in the postsynaptic cell,⁵⁵ although the contribution of this mechanism to depression of IPSPs in thalamocortical neurons remains to be examined.

Together, the process of synaptic facilitation of corticogeniculate synapses and depression of GABAergic pathways imparts on the corticogeniculate pathway the ability to exhibit predominantly inhibitory or excitatory effects, depending upon the frequency of stimulation. At low frequencies (e.g., < 1 Hz) activation of the corticogeniculate pathway may result in prominent IPSPs in thalamocortical neurons owing to excitation of thalamic reticular and intra-nuclear GABAergic neurons. However, at higher frequencies of activation, this inhibitory response gives way to pronounced and direct EPSPs. An additional factor to consider is the ongoing activity of thalamic reticular and intrathalamic interneurons, even in the absence of corticogeniculate activation. The spontaneous or evoked activity of these cell types will likely result in a baseline level of presynaptic inhibition of GABA release. Activation of corticothalamic axons in this case may, therefore, result in prominent excitatory effects from the very beginning of repetitive stimulation.

Corticogeniculate fibers activate a slow excitatory postsynaptic potential

We have previously demonstrated that the repetitive

stimulation of corticogeniculate fibers in the guinea-pig LGNd results not only in the generation of fast, monosynaptic EPSPs, but also in the generation of a slow EPSP that has a latency to peak of several seconds and that may persist for up to 1 min.³⁷ This slow EPSP persists following the block of muscarinic, α and β adrenergic, serotonergic, and H_1 and H_2 histaminergic receptors and is occluded by the maximal stimulation of glutamate metabotropic receptors. Here we demonstrate that the antagonism of glutamate metabotropic receptors results in a substantial reduction of the corticogeniculate slow EPSP, confirming that this event is mediated through the activation of these receptors.

Eight different subtypes of metabotropic glutamate receptor have been cloned and have been segregated into three different groups based upon the postsynaptic responses they induce, their sequence similarities, and pharmacology (for review see Refs 38 and 44). In the thalamus, metabotropic glutamate receptor 1 has been demonstrated by both *in situ* hybridization^{30,50} as well as immunocytochemistry.^{4,19,29,58} Examination at the electron microscopic level has revealed that metabotropic receptor 1a, a splice variant of metabotropic glutamate receptor 1, is prominent in the distal, cortico-recipient, zones of thalamocortical neurons⁵⁸ and pharmacological analysis of corticothalamic slow EPSPs suggest that they are mediated by group I metabotropic receptors.¹¹ These results suggest that the corticothalamic synapses operate through the activation of both ionotropic AMPA/kainate and NMDA receptors and glutamate metabotropic receptors. Electron microscopic results have suggested that glutamate metabotropic receptors may be localized adjacent to synaptic terminals, and are therefore activated by "spill over" of glutamate,^{4,19,58} thereby presumably requiring a more intense release of glutamate for their activation. This possibility remains to be carefully examined in the corticogeniculate pathway.

Recent results suggest that there may exist an important glutamatergic input to thalamocortical neurons from the brainstem.¹³ Whether or not these glutamatergic synapses may also activate metabotropic receptors, and therefore contribute to the ascending control of thalamocortical activity through modulation of the membrane potential of these cells, is unknown. Since these brainstem inputs also synapse onto the dendrites of thalamocortical cell and appear morphologically similar to corticogeniculate synapses it is likely that metabotropic glutamate 1a receptors are also adjacent to these brainstem inputs.

Functional properties of corticogeniculate transmission

Despite numerous *in vivo* investigations, the

functional role of the massive corticothalamic projection is still not well understood. In general, inactivation of the cerebral cortex results in a decrease in excitability of thalamocortical neurons, a result which is consistent with the direct excitatory effects of this pathway^{17,18} (reviewed in Ref. 53). However, other studies have also found increases in responsiveness of some thalamic neurons following inactivation of the cerebral cortex, suggesting that the functional role of this pathway may be complex. One important feature in determining the excitatory or inhibitory influence of the corticogeniculate path is the spatial distribution of afferent fibers between these two structures.⁵⁶ Here we also demonstrate that the temporal structure of activation of corticothalamic pathway is important in determining the functional influence of layer VI on thalamocortical responsiveness. At low frequencies of activation, the inhibitory influence of activation of thalamic reticular or local interneurons are likely to dominate. However, upon higher frequency

stimulation, or repetitive stimulation, these inhibitory influences should undergo frequency-dependent depression. At the same time, the monosynaptic corticothalamic EPSPs would undergo synaptic facilitation, resulting in a switch from predominately inhibitory to excitatory interactions between the cortex and thalamus (see Figs 2 and 3). Repeated activation of corticothalamic inputs may also result in a slow depolarization of thalamocortical cells through the activation of glutamate metabotropic receptors. This slow depolarization will result in an enhancement of the tonic, single spike mode of action potential generation in thalamocortical cells and a suppression of intrinsic and network-based low-frequency oscillations.

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