Involvement of Serotonin in the Excitation of Phrenic Motoneurons Evoked by Stimulation of the Raphe Obscurus

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Short-latency averaged responses in the C5 phrenic nerves to electrical stimulation (2.5-80 μA; 5-80 Hz; 150 μsec pulse duration) of raphe pallidus (RP) and raphe obscurus (RO) were investigated in anesthetized, paralyzed, and artificially ventilated cats. The responses to stimulation of RO were excitatory, whereas a mixture of inhibitory and excitatory responses of lesser magnitude were observed after stimulating in RP. The maximal response was obtained from the ventral part of RO and consisted of early and delayed excitatory responses that were of equal magnitude in both left and right C5 phrenic nerve roots. The mean latency for the early response was 2.5 ± 0.1 msec and for the delayed response was 7.0 ± 0.2 msec. Both responses were elicited during inspiratory phase stimulation, but only the delayed response was present during expiratory phase stimulation. The stimulus threshold of the early response was 5 μA; the delayed response was elicited at currents as small as 2.5 μA. Early and delayed responses were affected in different ways by increasing stimulus current and by increasing stimulus frequency. Intravenous administration of serotonin receptor antagonists methysergide (0.1-0.7 mg/kg), metergoline (33-244 μg/kg), and cinanserin (1.5-9.0 mg/kg) caused significant dose-related reductions in the magnitude of the delayed response, but did not significantly affect the early response. These data suggest that the early and delayed excitatory responses are mediated by different neuronal pathways. The early response does not involve serotonin release, while the later response is mediated at least in part by activation of a serotonergic pathway. The early excitatory response may be mediated by descending bulbospinal inspiratory axons that decussate in the midline before proceeding to the spinal cord to activate phrenic motoneurons. However, the delayed excitatory response probably occurs as a result of activation of serotonergic cell bodies in RO. These cell bodies likely project to medullary respiratory nuclei, which then project to the spinal cord to evoke phrenic motoneurons. These findings are important in that they identify a specific respiratory motor response mediated by a specific neurotransmitter and a brain stem nucleus.

Evidence from recent anatomical and physiological studies indicates that the caudal raphe nuclei, consisting of the raphe obscurus (RO), raphe pallidus (RP), and raphe magnus (RM), may be involved in neural control of respiration. Retrograde and anterograde tracing techniques have revealed that the caudal raphe nuclei project to nuclei known to be involved in respiratory control. These nuclei include the phrenic motor nucleus (PMN) in the spinal cord and the nuclei of the tractus solitarius, which includes the dorsal respiratory group in the medulla. Holtman et al. (1984a) showed that RO, RP, and RM project into the area of the PMN in the cervical spinal cord. In addition, the RM was also shown to project to the nuclei of the tractus solitarius (Basbaum et al., 1978).

Using neurochemical and immunohistochemical techniques, the neurotransmitter serotonin, which is localized primarily in cell bodies of the raphe nuclei (Dahlström and Fuxe, 1964; Poitras and Parent, 1978; Wiklund et al., 1981), has been shown to be present in several nuclei involved with respiratory control. Oliveras et al. (1977) found a high serotonin content in micropunches from the ventral horn of the cervical spinal cord at C5 and C6, the location of the PMN column. Furthermore, serotonin-containing varicosities, which are thought to represent nerve terminals, have been found surrounding phrenic motoneurons (Holtman et al., 1984b). Serotonin-containing varicosities have also been identified in the area of the nuclei of tractus solitarius (Maley and Elde, 1982), as well as in nucleus ambiguus (Steinbusch, 1981), the latter constituting a portion of the medullary ventral respiratory group. Significant amounts of serotonin have been found in micropunches from the area of the nuclei of tractus solitarius and also in nucleus ambiguus (Paliovits et al., 1974).

Stimulation of the caudal raphe nuclei produces changes in respiratory activity. As early as 1939, Pitts and colleagues reported changes in respiratory activity in the spontaneously breathing cat induced by electrical stimulation of RO, RP, and RM. Stimulation within RO and RM resulted in mixed effects, observed as either a tonic inspiration or tonic expiration, whereas stimulation within the RP resulted in tonic inspiration. Sessle and colleagues (1981) reported that stimulation of RM inhibits inspiration as brief stimuli resulted in periods of apnea. Changes in overall phrenic nerve activity have also been observed during electrical stimulation of the caudal raphe nuclei. Phrenic nerve activity was increased by stimulation of RP and decreased by stimulation of RM (Drechsel et al., 1983). Holtman and coworkers (1986) showed that stimulation of RO causes an increase in phrenic nerve activity. Changes in phrenic nerve activity were also seen after microinjection of L-glutamate into the RO. This finding indicates that activation of RO cell bodies and not axons of passage (Goodchild et al., 1982) is responsible...
for the increase in the phrenic nerve activity. Lalley (1985) recorded intracellularly from phrenic motoneurons and found that stimulation of RO causes inhibition, but that stimulation of the RP causes excitation of phrenic motoneuron spontaneous activity.

In summary, data from prior studies indicate that serotonergic neurons of caudal raphe nuclei may be involved in respiratory control. However, as only single sites of stimulation within each raphe nucleus typically were studied and the role of serotonin in mediating changes in respiratory activity was not addressed, we felt that a more complete evaluation needed to be done. In the present study, we have systematically examined the effects of stimulation of the caudal raphe nuclei on short-latency phrenic nerve responses and determined whether the neurotransmitter serotonin mediates the observed responses. We have focused on the RO and RP and characterized the short-latency phrenic nerve responses elicited by electrical stimulation throughout the entire rostrocaudal and dorsoventral aspects of these nuclei. Furthermore, the effects of several serotonin antagonists (methysergide, metergoline, and cinanserin) have been evaluated for their abilities to block the evoked phrenic nerve responses to stimulation of RO.

Materials and Methods

General surgical procedures

Twenty-four cats of either sex, ranging in weight from 1.8 to 3.5 kg, were anesthetized initially with Althesin (20 mg/kg, i.m., Glaxo Vet, England), followed by chloralose-urethane (30 and 150 mg/kg, i.v.). Supplemental doses of chloralose-urethane were administered to maintain anesthesia. A femoral artery and vein were cannulated for measurement of blood pressure and administration of drugs, respectively. Blood pressure was continuously recorded on a Gould/Brush polygraph. The trachea was also cannulated and end-tidal CO₂ measured (Beckman LB-2), filtered (1 Hz-10 kHz), and displayed on an oscilloscope; the osclilloscope signal was sent to a signal-averaging computer (Princeton Applied Research) for analysis. The neural signals were also full-wave-rectified, integrated using a "leaky integrator" circuit, and the integrated output continuously recorded on the polygraph to monitor inspiratory output.

Electrical stimulation

The overlying muscles of the head and neck were dissected, and a limited occipital craniotomy was performed to expose the caudal cerebellum and medulla. The dura was cut and retracted, and the caudal half of the cerebellum was removed by aspiration, thereby exposing the dorsal surface of the medulla. A stimulating electrode (stainless steel, monoconductor, 200-500 kΩ) was positioned perpendicular to the medullary surface and then visually aligned with the midline using the obex as a reference point. A stimulator (Grass S88) connected to a constant-current isolation unit (Grass P51) delivered square-wave pulses at points throughout the RO and RP. Parameters for electrical stimulation were in the range of 2.5-80 μA, 5-80 Hz, with stimulus pulses of 150 μsec duration. Generally, stimuli were applied throughout the inspiratory phase as well as the expiratory phase of the respiratory cycle. At the end of each experiment, stimulation sites were marked by passing a current of 100 μA for 5 sec. The brain stem was then removed and placed in 10% formalin in 0.9% NaCl. Fifty-micron sections were cut either in a sagittal or a transverse plane. Sections were stained with cresyl violet and Luxol fast blue. Lesion sites were identified using a microprojector (Bausch & Lomb).

Data analysis

Short-latency responses in the phrenic nerves to electrical stimulation within RO and RP were analyzed using signal-averaging techniques. The trigger to the averager was generated by the stimulator. The parameters for averaging were 200 sweeps, 1024 bins with 40 μsec/bin. We simultaneously averaged both right and left phrenic nerve responses on line. After acquisition of each pair of averages, the averaging computer memory was sent to an LM2 laboratory computer for storage on floppy disk for subsequent plotting and analysis. The evoked responses were quantified by determining the areas under each response (peak) using a digitizing tablet (Numonics Co.). In order to compare data between cats, the areas were converted to the percentage of maximal response for the current and frequency response curves, and to the percentage of control (vehicle only) for dose–response curves in the pharmacological experiments.

Pharmacological studies

Three serotonin antagonists (Fuller, 1980) were evaluated for their effects on the evoked phrenic nerve averaged responses to stimulation within RO: methysergide (Sandox, East Hanover, NJ), metergoline (Farmitalia, Milan, Italy), and cinanserin (E. J. Squibb, Princeton, NJ). These compounds have been found to be selective serotonin antagonists (McCall, 1984; Proudfoot and Anderson, 1974). Methysergide and cinanserin were dissolved in 0.9% NaCl; metergoline was dissolved in 95% ethanol.

Cumulative dose–response curves were generated by intravenous administration of each drug. Initially, the effect of vehicle control on evoked responses in phrenic nerve was evaluated. Neither 0.9% NaCl nor 95% ethanol caused significant changes in evoked responses. After administration of the respective vehicle control, successive doses of each drug were administered at 5-15 min intervals. The effects of the drug on evoked responses in phrenic nerve were determined just prior to administration of the next dose of drug. The level of phrenic nerve activity was observed to decrease in some cats after the higher doses of serotonin antagonists. When this decrease occurred, inspired CO₂ was increased to maintain phrenic nerve activity at pre-drug injection control level. All drug responses were converted to percentage of control, the latter being the evoked response obtained after administration of the respective vehicle in which each drug was dissolved.

Statistical analysis

Current and frequency response curves, as well as the dose–response curves, were evaluated by regression analysis (Snedecor and Cochran, 1967). The Student's t test for paired data was used for all other comparisons. The level of statistical significance for each test was p < 0.05. In all cases, values are reported as means ± SEM.

Results

Anatomical mapping of the averaged responses elicited in the phrenic nerve by electrical stimulation of raphe obscurus and raphe pallidus

We began our study by determining what types of averaged phrenic nerve responses could be evoked by electrical stimulation during inspiration within the RO and RP. Responses to stimulation (50 μA, 20 Hz, 150 μsec pulse duration) were mapped along the entire rostrocaudal and dorsoventral extents of the RO and RP as defined by Taber and colleagues (1960). Stimulations were made at 5, 4, 3, 2, and 1 mm rostral to the obex, at the obex, and at 1 mm caudal to the obex. In each track, the stimulations were performed in 500 μm increments from the dorsal surface of the medulla. This procedure was performed in six cats, and a representative map is presented on a mid sagittal section of the medulla in Figure 1. The extents of RO and RP are represented in this figure by dashed lines; RO can be seen to begin just caudal to the obex and to extend to about 5 mm rostral to the obex. The RP begins near the obex and extends rostrally for 5 to 6 mm. Rostrally, the RO and RP can be seen...
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The averaged responses elicited in the phrenic nerves were mixed. Electrical stimulation produced excitation, inhibition, or no response. An example of an excitatory response is seen at +4 mm and 4500 μm. The inhibitory responses were more numerous and can be seen at +2 mm and 4000 μm, +1 mm and 3500 μm, or obex and 2500 μm. In some instances, the electrical stimulations produced essentially no response, as can be seen at +5 mm and 5000 μm, +3 mm and 4500 μm, or +1 mm and 3000 μm.

The type of response elicited in the phrenic nerves by electrical stimulation within RO was clearer. The predominant responses were excitatory. Examples can be observed in all of the tracks at various depths along the entire rostrocaudal extent of the RO. Although excitatory responses were elicited in the phrenic nerves by stimulating at several sites within RO, the maximal excitatory response in the six cats tested consistently occurred at +2 or +3 mm rostral to the obex and at 2500 or 3000 μm deep from the dorsal surface of the medulla. Figure 1 shows the excitation, consisting of an early and a delayed excitatory response at +3 mm and 3000 μm.

We focused our attention on this site within RO to characterize more completely the nature of the early and delayed excitatory response. To determine the mediolateral extent of the excitatory response, electrical stimulation (50 μA, 20 Hz, 150 μsec pulse duration) was performed within the midline and at 1 and 2 mm on either side of the midline. Stimulations were again performed at 500 μm intervals from the surface of the medulla. These mediolateral stimulation experiments were done in four cats, and a representative map of the results from one animal is presented on a cross section of the medulla in Figure 2. In this plane, the maximal excitatory response occurred in the midline within RO at 2500 μm (Fig. 2). In the four cats tested, the depth of the maximal delayed excitatory response occurred at 2500 or 3000 μm. The site of the maximal excitatory response in each animal was always dorsal to the inferior olive. The distance rostral from the obex, where the maximal excitatory response occurred, ranged from 2.0-2.2 mm. Evidence of excitation was also present at 1 and 2 mm lateral to the midline. At 1 mm lateral, the response was present at 3500 μm. Although present in these lateral tracks, the excitatory response was centered in the midline and was much diminished at 2 mm lateral to the midline.

Electrophysiological characterizations of the averaged excitatory response elicited in the phrenic nerve by electrical stimulation of the raphe obscurus

Following anatomical characterization of the averaged excitatory response elicited by electrical stimulation of the RO, the averaged excitatory response was characterized electrophysiologically. First, we investigated whether early and delayed excitatory responses occurred in both right and left phrenic nerves, and whether the responses could be elicited when stimulation was applied during expiration. This was done in four cats and representative results from one animal are presented in Figure 3. This figure shows that early and delayed excitatory responses are clearly present in a single-sweep record as well as in the averaged record (cf. A and B, Fig. 3). The latency, form, and magnitude of both early and delayed responses were similar in the left and right C5 phrenic nerves. Both excitatory responses were present when electrical stimulation was applied during inspiration; however, only the delayed response could be elicited in each of the phrenic nerves when stimulation was applied during expiration (Fig. 3).

Clear separation between the early and delayed excitatory

Figure 1. Representative map in the midsagittal plane of the averaged C5 phrenic nerve responses elicited by electrical stimulation of the raphe obscurus and pallidus. Stimulations were made during inspiration at -1 to +5 mm from obex and at 500 μm increments from the dorsal surface of the medulla. Stimulus parameters: 50 μA, 20 Hz, and 150 μsec. PT, Pyramidal tract; RM, raphe magnus; RO, raphe obscurus; RP, raphe pallidus.
Figure 2. Representative map in the transverse plane of the averaged C5 phrenic nerve responses elicited by electrical stimulation, both within and lateral to the raphe obscurus and pallidus. Stimulations were made during inspiration in the midline and at 1 and 2 mm on each side of the midline at 500 μm increments from the dorsal surface of the medulla. Stimulus parameters: 30 μA, 20 Hz, and 130 μsec. DMV, Dorsal motor nucleus of the vagus; PT, pyramidal tract; RO, raphe obscurus; RP, raphe pallidus; TS, tractus solitarius; XII, hypoglossal nucleus.

responses in the phrenic nerves allowed for calculations of onset and peak latency times. Results of these calculations in 19 cats are presented in Table 1. The average latency-to-onset was 2.5 ± 0.1 msec for the early response and 7.0 ± 0.2 msec for the delayed response. The peak of the early response occurred about 1 msec (latency-to-peak of 3.6 ± 0.1 msec) after its onset; this brief time is indicative of the sharpness of this peak. The peak of the delayed response occurred about 3 msec after its onset (a latency-to-peak of 10.2 ± 0.2 msec); this longer time is indicative of the greater width of this peak compared with the earlier one.

The effect of altering stimulus intensity (2.5–80 μA) and stimulus frequency (5–80 Hz) on the early and delayed excitatory responses was tested. These tests were performed on 10 and 9 cats, respectively, and the results are presented in Figures 4 and 5. Magnitudes of both the early and delayed responses increased with current intensity (Fig. 4). The threshold for the early response was approximately 5 μA, while the delayed response could be elicited at currents as small as 2.5 μA. The maximum for the early response occurred at 40–80 μA; the delayed response continued to increase for currents as high as 80 μA. Magnitudes of the early and delayed responses were also dependent on stimulus frequency, but neither response was linearly related to stimulus frequency. The early response was maximal at 5–10 Hz, reached a minimum at 40 Hz, and then began to increase at 80 Hz. The magnitude of the delayed response was linearly related to frequency up to 20 Hz, but decreased at 40 and 80 Hz.

Pharmacological characterization of averaged responses elicited in phrenic nerve by electrical stimulation of raphe obscurus

In order to determine if either or both components of the excitatory response elicited in the phrenic nerve by electrical stimulation of RO was mediated by serotonin, we first tried blocking the response with methysergide. Methysergide was administered intravenously in successive doses at 5 min intervals to five cats; representative results from one cat are presented in Figure 6. Successive doses of methysergide caused a dose-related reduction in the magnitude of the delayed response, which was essentially abolished after a dose of 0.7 mg/kg. However, following this, a small-amplitude, delayed response could be evoked when stimulus intensity was increased to 100 μA. Methysergide had no significant effect on the magnitude of the early response. For all cats tested with methysergide, the cumulative dose–response curve is shown in Figure 7A. One can see that only the delayed excitatory response is significantly affected by the drug. The slope of the early excitatory response curve was not significantly
different from zero. Two other serotonin receptor antagonists, metergoline and cinanserin, were also tested for their abilities to block the delayed excitatory response. Successive doses of metergoline were administered intravenously at 10 min intervals to five cats, and the results are shown in Figure 7B. Metergoline also caused a significant dose-related reduction in the magnitude of the delayed excitatory response. The slope of the early excitatory response did not differ significantly from zero, indicating that metergoline also had no dose-related effect on the early response. Successive doses of cinanserin were administered at 15 min intervals to four cats, and the results are shown in Figure 7C. Cinanserin also caused a significant reduction (although much smaller) in the magnitude of the delayed response, but had no significant effect on the early response. Whereas methysergide and metergoline at high enough doses could abolish the delayed excitatory response evoked at 50 μA, cinanserin only reduced the response to 85% of control. The maximal reduction occurred at the 6.0 mg/kg dose; larger doses of cinanserin did not produce any greater reductions in the magnitude of the delayed response. The reduction at 9.0 mg/kg was not significantly different from that at 6.0 mg/kg. Doses larger than 9.0 mg/kg were not possible, as they produced cardiotoxicity characterized by severe hypotension and arrhythmias, often resulting in death of the animal.

| Latency-to-onset (msec) | Latency-to-peak (msec) |
|------------------------|------------------------|
| Early response         | 2.5 ± 0.1 (1.9–2.8)    |
|                        | 3.6 ± 0.1 (2.8–4.6)    |
| Delayed response       | 10.0 ± 0.2 (5.6–8.3)   |
|                        | 10.2 ± 0.2 (9.0–11.6)  |

Values are means ± SEM; n = 19. Values in parentheses are ranges. Stimulus parameters for evoking responses were 50 μA, 20 Hz, and 150 μsec.

Figure 3. Representative responses in simultaneously recorded right and left C5 phrenic nerves elicited by electrical stimulation of the raphe obscurs during inspiration and expiration. A, Single sweep responses; B, averaged responses. Control records are without stimulation. Stimulus parameters (A, B): 50 μA, 20 Hz, and 150 μsec.

Figure 4. Averaged C5 phrenic nerve responses elicited by electrical stimulation of the raphe obscurs during inspiration as a function of stimulus intensity. Values are means ± SEM; n = 10 cats at each point. Stimulus parameters: 20 Hz, 150 μsec. Early response, open circles; delayed response, filled circles.
Figure 5. Averaged C5 phrenic nerve responses elicited by electrical stimulation of the raphe obscurus during inspiration as a function of stimulus frequency. Values are means ± SEM; n = nine cats at each point. Stimulus parameters: 50 μA, 150 μsec. Early response, open circles; delayed response, filled circles.

Discussion

The results of the present study indicate that inspiratory phase stimulation of RO and RP produces short-latency responses in the phrenic nerves. The responses in the RO were excitatory, while a mixture of inhibitory and excitatory responses of smaller magnitude were observed in the RP. The maximal response obtained was from the ventral part of RO; it consisted of early and delayed excitatory responses that were of equal magnitude in both the left and right C5 phrenic nerve roots. This response appears to be similar to the one reported by Pitts (1943) with electrical stimulation in the midline.

Several findings indicate that the early and delayed responses are mediated, at least in part, by different neuronal pathways. Evidence for this includes (1) the different onset latencies of the responses; (2) different current and frequency response relationships; (3) presence of the delayed response, but not the early response, during expiratory-phase stimulation; and (4) reduction in magnitude of the delayed but not the early response by serotonin receptor antagonists.

The early response probably results from stimulation of bulbospinal inspiratory axons that cross the midline in this part of the RO before proceeding to the spinal cord, where they monosynaptically excite phrenic motoneurons (Lipski et al., 1983). This conclusion is supported by both neurophysiological and neuroanatomical evidence. First, the mean latency-to-onset of 2.5 ± 0.1 msec and mean latency to peak of 3.6 ± 0.1 msec for this response are consistent with direct stimulation of inspiratory bulbospinal axons. Mean axonal conduction velocity of inspiratory bulbospinal axons is approximately 55 m/sec (Dick and Berger, 1985). Mean axonal conduction velocity of phrenic motoneuronal axons is also approximately 55 m/sec (Berger, 1979). These values lead to a calculated mean axonal conduction time of about 0.7 msec for the 95 mm distance between the point of stimulation in the RO and the phrenic nerve recording site. If a utilization time of 0.25 msec (see Dick and Berger, 1985) for generation of the descending volley, a chemical synaptic delay of 0.5 msec (Eccles, 1968) and a delay of 0.3 msec to initiate the phrenic motoneuron axonal spike from the start of the EPSP are included (see Berry and Pentreath, 1976), then the observed latency-to-onset and latency-to-peak of the early phrenic response are consistent with direct activation of inspiratory bulbospinal axons. Second, decussating axons from bulbospinal inspiratory neurons have been identified in the midline, where they monosynaptically excite phrenic motoneurons (Lipski et al., 1983). Evidence for this includes (1) the different onset latencies of the responses; (2) different current and frequency response relationships; (3) presence of the delayed response, but not the early response, during expiratory-phase stimulation; and (4) reduction in magnitude of the delayed but not the early response by serotonin receptor antagonists.

The delayed response in the phrenic nerves elicited by stimulation of RO is of greater interest. Whereas the early response is most likely caused by stimulation of axons of passage, the delayed response probably results from activation of cell bodies of RO. This conclusion is supported by previous findings of Holtman et al. (1986). These investigators used L-glutamate to activate cell bodies from a part of the RO similar to that examined in the present study. Microinjection of L-glutamate re-
serotonin via the ventral and ventrolateral funiculi to the ventral motoneurons in the spinal cord (Dahlström and Fuxe, 1965; addition, serotonergic nerve terminals are found surrounding and Fuxe, 1965; Martin et al., 1978). These projections ter-
minate in the area of the PMN (Holtman et al., 1984a). In
consistent with the delayed excitatory response observed in the present study. Our pharmacological data also suggest that activation of cell bodies in RO is responsible for the delayed response in the phrenic nerve. Serotonin is found primarily in cell bodies of raphe nuclei, and is thought to be a neurotransmitter used by these neurons (Dahlström and Fuxe, 1964; Po-
terus and Pareit, 1978, Wiklund et al., 1981). Therefore, activation of cell bodies of RO would be expected to release serotonin and activate serotonin receptors. All three serotonin receptor antagonists used in this study caused inhibition of the delayed response, strongly suggesting that activation of serotonergic cell bodies of RO is responsible for at least part of the delayed response. The presence of the delayed response to stimulation lateral to midline may be due to activation of neural processes arising from RO cell bodies that enter the reticular formation (Fox et al., 1976).

Our pharmacological data indicate that a serotonergic synapse is involved in mediating the delayed response. Methysergide, metergoline, and cinanserin are selective and competitive sero-
tonin receptor antagonists (Leysen et al., 1982; Peroutka et al., 1981) that have been shown to block the facilitatory actions of serotonin (McCall, 1984; McCall and Aghajanian, 1980). How-
ever, some caution must be used in drawing conclusions about actions of drugs in the CNS following their systemic adminis-
tration. Drugs may have actions at several sites, which will all contribute to a net final effect. Although a serotonergic pathway is involved in the delayed response, it is difficult to determine conclusively from our data whether the delayed response is mediated by a monosynaptic projection directly onto phrenic motoneurons or by projections to other respiratory nuclei that then project to phrenic motoneurons.

Evidence that the delayed response is mediated to some extent by a direct monosynaptic serotoninergic projection onto phrenic motoneurons is found in both neuroanatomical and physiolog-
ical studies. The RO has been shown to send serotoninergic projec-
tions via the ventral and ventrolateral funiculi to the ventral horn of the spinal cord (Basbaum and Fields, 1979; Dahlström and Fuxe, 1965; Martin et al., 1978). These projections ter-
minate in the area of the PMN (Holman et al., 1984a). In
addition, serotoninergic nerve terminals are found surrounding motoneurons in the spinal cord (Dahlström and Fuxe, 1965; Steinbusch et al., 1978), including phrenic motoneurons (Hol-
man et al., 1984b). Furthermore, serotonin has been shown to facilitate motoneuron excitability in the spinal cord (Myslinski and Anderson, 1978; White and Neuman, 1980).

The delayed excitatory response may also be mediated by activation of medullary respiratory nuclei, which then project to and excite phrenic motoneurons. Again, there is both neu-
ronanatomical and neurophysiological evidence supporting this possibility. Serotonin nerve terminals have been found in the nuclei of the tractus solitarius (Maley and Elde, 1982), which contains the dorsal respiratory group neurons, and in the area of the nucleus ambiguous (Steinbusch, 1981), which constitutes part of the ventral respiratory group. Neurons from both of these areas project to and excite phrenic motoneurons (Lipski et al., 1983; Merrill, 1974). Furthermore, serotonin has been shown to cause an increase in the firing rate of inspiratory neurons in the dorsal respiratory group (Sessle and Henry, 1985). Additional evidence indicating that the delayed response is mediated through medullary respiratory nuclei comes from findings in the present study. The onset latency of 7.0 msec for the delayed response is not consistent with what would be expected of a monosynaptic serotoninergic spinal cord projection. A conduction velocity of about 10 m/sec is too fast for these small diameter axons of the raphe–spinal monosynaptic pathway is predicted by this onset latency. Serotonergic axons in the spinal cord are both unmyelinated and finely myelinated, and have been shown to have diameters averaging about 1 μm (Dahlström and Fuxe, 1965; Ruda and Gobel, 1980). This predicted conduction velocity of 10 m/sec is too fast for these small diameter axons of the raphe–spinal serotonergic pathway (Wes-
sendorf et al., 1981). Although the onset latency is too short to explain a serotoninergic raphe–spinal pathway, it could explain a slowly conducting serotonergic pathway via the dorsal and/or ventral respiratory group neurons, which then project by rapidly conducting medullary inspiratory axons to the PMN. The distance from RO to the dorsal or ventral respiratory groups is approximately 3.5 mm (Berman, 1968). The latency between these nuclei would be 3.5 msec, assuming a pathway with con-
duction velocity of 1 m/sec. Using this latency, we predict that the overall onset latency would be approximately 7.0 msec for the disynaptic pathway between RO and the PMN. This is con-
sistent with the experimentally determined onset latency of the delayed response observed in the present study.

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**Figure 7.** Cumulative dose-response curves of the effects of serotonin antagonists on averaged C5 phrenic nerve responses elicited by electrical stimulation of the raphe obscurus during inspiration. A, Methysergide; B, metergoline; C, cinanserin. Stimulus parameters: 50 μA, 20 Hz, and 150 μsec. Early response, open circles; delayed response, filled circles. Values are means ± SEM; n = three to five cats at each point.
The delayed response may not be totally mediated by release of serotonin. This conclusion is based on two findings. First, the blockade of the delayed response by serotonin antagonists could be partially reversed by increasing current intensity. Second, cinanserin, which has been shown in other studies to be effective in antagonizing the facilitatory effects of serotonin (McCall, 1984; McCall and Aghajanian, 1980), was able to block only a small portion of the delayed excitatory response. These data suggest that other neurotransmitters may mediate a portion of the delayed response. Other putative neurotransmitters, such as substance P, thyrotropin-releasing hormone, and met-enkephalin, have been identified in cell bodies of RO (Hökfelt et al., 1978; Hunt and Lovick, 1982; Johansson et al., 1981; Lovick and Hunt, 1983).

In conclusion, we have found in this study the presence of a serotonergic pathway, probably arising from cell bodies within RO and mediating an excitation of phrenic motoneurons. To the best of our knowledge, this constitutes the first demonstration of a specific respiratory motor response associated with a specific neurotransmitter and a brain stem nucleus.

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