To the Editor:—We appreciate the three articles in the April 2004 issue of ANESTHESIOLOGY regarding GW280430A.1-3 Although remarkable advances in developing intermediate and short-acting muscle relaxants were realized, anesthetists have not yet been provided with a substance comparable to succinylcholine in terms of its rapid onset and ultrashort-acting pharmacodynamic profile.

However, this aim should not be lost. A survey in Germany4 revealed that, despite its undesirable side effects, succinylcholine is still the most used drug for both rapid sequence inductions5 and for elective case induction.6 An overwhelming majority (76.6%) of respondents answered that they would appreciate a nondepolarizing substitute for succinylcholine if a similar pharmacodynamic profile was preserved. Assuming that this is not an isolated German viewpoint, a substance replacing succinylcholine would be highly desirable.

The developers and the researchers have a great responsibility when introducing a new drug into clinical practice, particularly in neuromuscular blocking drugs. Dr. Caldwell addresses this issue in his editorial when he compares the side effects of rapacuronium and GW280430A.8 Because we were involved in the clinical evaluation of rapacuronium,7-11 we would like to comment on some relevant aspects of the side effect profiles of both drugs and on the drug approval processes. First, in clinically relevant concentrations, rapacuronium potentiates bronchoconstriction most probably by destabilization of the balance between M2 and M3 muscarinic receptors.12 In contrast, GW280430A seems to release histamine5 and therefore may possibly induce bronchoconstriction. Second, although many antihistaminic drugs and prophylactic strategies are available, an effective treatment to rebalance the muscarinic effects of rapacuronium was and is still missing. Third, because rapacuronium did not release histamine,13 because different M2 versus M3 muscarinic effects of muscle relaxants were unknown at that time, and because clinical symptoms of the pulmonary side effects differed from those seen during typical bronchoconstriction,12 the clearly described dose-dependent pulmonary side effects (from 10.7% with 1.5 mg/kg rapacuronium to 18.5% with 2.5 mg/kg rapacuronium)10 may have been questioned—unfortunately until patients were badly harmed. Therefore, we agree with Dr. Caldwell that the recent experience with rapacuronium must be considered during the trials with GW280430A, e.g., by in addition investigating its effects on M2 and M3 receptors. The fiasco with rapacuronium, however, must not induce pessimism if new drugs and especially GW280430A may have the potency to improve anesthesia practice.

GW280430A was, of course, not compared with rapacuronium, but it was also not compared to succinylcholine.1-3 Regardless, the hope that GW280430A will be a substitute for succinylcholine has been advanced6 with this first presentation. Expectations that this new drug will approximate the rapid onset of succinylcholine may in high doses, high injection speeds, and, therefore, the risk for high incidences of side effects. The presentations7-5 primarily suggest that GW280430A may be an ultrashort-acting rather than a rapid-onset muscle relaxant.

Unfortunately, preclinical and clinical trials to approve new drugs are expensive, and, in this context, the substance to be replaced is already very cheap. Nevertheless, we (and many other anesthetists14-16) would like to encourage the recent attempts to develop better muscle relaxants (or reversal drugs, e.g., Org 2596917) to improve safety and efficiency of neuromuscular treatment during anesthesia.

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Anesthesiology 2005; 102:862–3

In Reply—I am grateful to Drs. Geldner and Blobner and Dr. Lien et al. for their responses to my editorial.1 I agree with Drs. Geldner and Blobner that succinylcholine still has a significant clinical role, and that anesthesia providers would likely see it replaced by a nondepolarizing drug with a similar time course of action. I agree also that we should not be pessimistic about prospects for new drugs, but history and pulmonary disease.Bronchospasm or any other difficulty with ventilation was not encountered over the course of the volunteer trial. This problem has been encountered rarely in the thousands of patients who have received mivacurium, which has a greater propensity to release histamine than GW280430A.

The bronchospasm noted after administration of rapacuronium is likely not due to histamine release. Bronchospasm has occurred in patients receiving rapacuronium with no evidence of histamine release.6 The bronchospasm after administration of rapacuronium is likely caused by its antagonism of the muscarinic M2 receptor.7 Non-depolarizing neuromuscular blocking agents can interact with the M3 receptors that exist in the airways (M1, M2, and M3). Their antagonism of the M3 receptor causes bronchodilation by inhibiting vagally induced bronchoconstriction. Antagonism of the M2 receptors, which are located presynaptically at postganglionic parasympathetic nerve endings, results in an increased release of acetylcholine that subsequently binds to M3 receptors, causing bronchoconstriction. The affinity of rapacuronium for the M2 receptor is 15 times its affinity for the M3 receptor.7

As shown in experiments in cats,6 GW280430A is a very weak inhibitor of muscarinic receptors in general, with nearly the same safety ratio for this side effect as mivacurium. In cats, the muscarinic blocking dose (ED90) of GW280430A is more than 25 times its ED95 for neuromuscular block.8 A closer look at the data in the study of Heerdt et al.3 in dogs shows a complete lack of effect of GW280430A on airway pressures in the dog, even at doses of 50 times the ED95. Nevertheless, GW280430A will have to be further tested for its relative affinity for the muscarinic receptors of the airways, as will all other nondepolarizing relaxants that may be introduced into clinical practice. Based on the data published to date,2,9 there is no reason to anticipate that GW280430A may even rarely cause life-threatening bronchospasm.

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In Reply.—The three reports by our group in the April (2004) issue of ANESTHESIOLOGY1–3 on the new ultrashort-acting nondepolarizing relaxant GW280430A (AV430A) have evoked invited commentary by Caldwell2 and observations by Geldner and Blochner. Both anticipation and caution were expressed by these commentators regarding the possible future clinical performance of AV430A.

To add to the discussion, I would offer a rather optimistic viewpoint based on additional data in preparation for publication. I am certainly biased because of my closeness to the development of AV430A. Nevertheless, I do believe that future results will strengthen the candidacy of AV430A as a replacement for succinylcholine or as an excellent alternative.

AV430A is a representative of the new class of nondepolarizing relaxants, which we have called asymmetric mixed-ion channel blockers.4 These compounds are inactivated by two entirely chemical (nonenzymatic) mechanisms both in vivo and in vitro: cysteine adduction and pH-sensitive hydrolysis.5,6 The cysteine adduction reaction is notably manipulatable.5,6 In AV430A, both reactions are fast (combined T 1/2 of approximately 1–2 min) such that the duration of block is ultrashort, similar in duration to or even shorter than succinylcholine in cats, dogs, and three different primate species.1,3

Dr. Savarese has been a consultant to Glaxo Wellcome. Research Triangle Park, North Carolina, and is one of the inventors and patent holders of GW280430A and related compounds on behalf of Cornell University, New York, New York.

Notable characteristics of this series include high potency, easy antagonism by anticholinesterases, and rapid antagonism by cysteine (see last paragraph). AV430A is particularly nonaggressive,1 an important contrast vis-à-vis rapacuronium, and does not block sympathetic ganglia.3 AV430A is approximately four times weaker than mivacurium with respect to release of histamine in the rhesus monkey.1 These autonomic data suggest that the only possible side effect of AV430A that might appear in the clinical dose range is a rather mild tendency to cause minor symptoms and signs suggestive of slight degrees of histamine release, exemplified by facial erythema, brief decrease in blood pressure, and increase in heart rate.

AV430A and its related series were first studied in 1997. Despite encouraging results in human volunteers,2 the development process has been delayed. First, a corporate decision by Glaxo Wellcome (Research Triangle Park, NC) to discontinue the development of products for the field of anesthesia, and then the merger of Glaxo Wellcome and Smith Kline, halted the project for 4 yr until the AV430A series was acquired by Avera Pharmaceuticals (San Diego, CA) in 2002. In 2003, a second study of a new formulation of AV430A in volunteers was completed and reported in 2004. The kinetic and safety data are encouraging.7,8 The failure of rapacuronium and its removal from the market, as a result of the side effect of bronchospasm, will place greater scrutiny on the possible airway effects of all new relaxants. An ultrashort-acting agent such as AV430A, which will no doubt be given for tracheal intubation, will bear the greatest scrutiny. Bronchospasm after rapacuronium, although noted in clinical trials, was far more serious in
clinical practice with rapacuronium than foreseen before its release. This is the crux of the problem: Minimal effects of AV430A on the airway must be shown to provide convincing data regarding the safety of AV430A. Caldwell and Geldner and Blobner both allude to this.

Following is a discussion of some of the data regarding AV430A that are already published, which suggest that AV430A should be safe as far as airway effects are concerned. Minimal airway effects have already been found in dogs in doses up to 50 × ED₉₀. The nonnagalytic properties of AV430A found in cats mean that it is unlikely to have any blocking effects on M₂ or M₃ receptors in the human airway. This indicates minimal possibility of bronchospasm on this basis, in contrast with the severe responses noted after rapacuronium. Jooste et al. have shown that mivacurium has minimal interactions with M₂ and M₃ receptors where the actions of rapacuronium are clearly demonstrable. Because mivacurium, which is a predecessor of AV430A, has approximately the same large safety ratio (> 25) as does AV430A regarding lack of interaction with muscarinic receptors in cats, the imbalanced blocking property on muscarinic receptors in the human airway, which is the likely mechanism of rapacuronium-induced bronchospasm, is a most unlikely clinical scenario in the case of AV430A.

What about histamine release? Both Caldwell and Geldner and Blobner express concern about this, but in the rhesus monkey, the safety ratio of AV430A for this side effect is approximately four times greater than that of mivacurium. This results in the following comparison in humans: mivacurium, at 2.5–5 × ED₉₀ (0.20–0.25 mg/kg), causes greater symptomatology of histamine release when injected over 15 s than does AV430A when injected over only 5 s at 3–4 × ED₉₀ (0.5 mg/kg). Since the original study in human volunteers by Belmont et al. in 1998, AV430A has been reformulated. This reformulation has further improved the safety ratio for histamine release to approximately 4 × ED₉₀ in humans. As a reminder, this “safety ratio” is defined as the dose required to cause an average decrease in blood pressure of 10%, divided by the ED₉₀ for neuromuscular blockade: ED Hist/ED₉₀.

Both Caldwell and Geldner and Blobner caution that anesthesiologists might increase dosage to cause faster onset and thereby increase the possibility of histamine release by AV430A. We have already compared the onset and duration of AV430A in the human larynx and the thumb via via a via succinylcholine, rapacuronium, rocuronium, mivacurium, and cisatracurium. The data are not yet published but suggest that the onset of AV430A is as fast as that of succinylcholine and faster than those of the others. So why administer AV430A at a dosage higher than 0.4–0.5 mg/kg? Onset of block after AV430A does not get any faster at dosages higher than this and is probably circulation limited, as pointed out by Caldwell. The onset is fast in all species—dogs, cats, monkeys, and humans. Consequently, Geldner and Blobner, who believe that “GW280430A may be an ultrashort-acting rather than a rapid-onset muscle relaxant” are most likely incorrect. AV430A is clearly ultrashort and very rapid in onset in all studies. figure 1 shows a mechanomyograph recording of the response to AV430A (0.4 mg/kg) in the adductor pollicis in a healthy human volunteer during nitrous oxide–oxygen–fentanyl–propofol anesthesia. The pattern of block seen in figure 1 was noted in every volunteer subject of the more than 100 humans treated so far. The dose of 0.4 mg/kg is approximately 3 × ED₉₀. Doses up to 5 × ED₉₀ caused minimal side effects. A 5-s bolus dose of AV430A (0.4 mg/kg) was given at the arrow. Two control train-of-four (TOF) responses are followed by the elicited twitch in the thumb at 0.15 Hz. Twitch is abolished within approximately 80 s. Recovery begins at approximately +6 min (time scale at top). The time scale changes at and after this point. At +8 min, TOF shows appearance of T₂ and T₄ is at approximately 25% of control. At +10 min, TOF is 45% and T₄ is 75% of control. At +12 min, TOF is 85% and T₄ is 95% of control. At +13.5 min, TOF is 95% of control (control TOF is at the far left). The time from 5% T₂ to a TOF of 90% is 6 min. Heart rate (upper record) and blood pressure (lower record) show no change. Time and experience will tell, as clinical studies in patients are undertaken, whether early data from volunteers accurately predict the performance of AV430A in practice. The dosage recommended to achieve certain clinical endpoints, e.g., intubation of the trachea within 60 s, among others, must be defined by these future studies.

In volunteer studies thus far, at dosages as high as 0.8–0.9 mg/kg (approximately 6 × ED₉₀ or 2 × intubating dosage) where AV430A is given as a rapid (5 s) bolus, the manifestations of histamine release after AV430A are rather mild (facial flushing and brief decrease in blood pressure, not requiring treatment). There has been no bronchospasm. This suggests that clinicians could give these very large dosages safely, particularly by injecting them a little more slowly (such as over 15 s). The side effect of histamine release as caused by AV430A, because it is four times weaker than it is in mivacurium, may constitute a minor concern in future clinical practice. Time again will tell. Only after a couple of years of experience in thousands of administrations will the pattern be fully defined.

Nevertheless, we can be reassured by this data. If, as Kopman et al. have suggested, a dose of 2.0 or 2.5 × ED₉₀ of AV430A is enough for good to excellent intubating conditions within 60 s (this dose would be in the range 0.5 ± 0.05 mg/kg), there may very well be minimal side effects.

The chemical pathways of inactivation of AV430A are, in my opinion, its most promising feature. The chemical breakdown will ensure no prolonged neuromuscular blockade. Problems with atypical pseudocholinesterase will not be at issue. Cysteine, given intravenously as a “reversal drug” or “rescue agent,” will rapidly inactivate AV430A. Complete recovery from 100% twitch inhibition can be induced in monkeys with exogenous cysteine within 1–2 min. In the future, anesthesiologists may have the choice of spontaneous recovery (12–14 min) from AV430A via endogenous cysteine or induced recovery by giving additional cysteine, e.g., in case of an airway emergency.
Cysteine, given within 2–3 min after injection of AV430A, should abolish complete paralysis within 1–2 min. The latter treatment with cysteine may shorten the total duration of action in humans to an estimated 5 min to return of full neuromuscular function such as cough, normal vital capacity, and head lift.

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In the current cases, the local anesthetic solution must have reached the cranial parts of the lumbar plexus because sufficient anesthesia and analgesia of the inguinal region was observed in all five patients. Recent investigations of ultrasound-guided posterior lumbar plexus block in pediatric patients revealed a greater extent of anesthesia and analgesia compared with adults.

It was not the aim of our study to investigate posterior lumbar plexus block for inguinal hernia repair in children, and we agree with Dr. Jöhr that it should not be the first choice for this surgical procedure. However, in our opinion, it might be a useful alternative and still represents a peripheral nerve block.

Dr. Jöhr also had concerns about the strategy of applying small volumes in the current setting. Nevertheless, one of the proven benefits of ultrasound-guided techniques is the decreased need of local anesthetics compared with traditional approaches.

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(Accepted for publication November 30, 2004.)

Of Mice and Men: Should We Extrapolate Rodent Experimental Data to the Care of Human Neonates?

To the Editor—Over the years, Dr. Olney et al.1 have enlightened the scientific community with their research on the N-methyl-D-aspartate receptor and its role in human disease. Recently, Dr. Olney’s group has examined the effects of commonly used anesthetics and anticonvulsants on the developing brain. In several high-profile scientific journals, they reported that the administration of these drugs, including ketamine, ethanol, phencyclidine, nitrous oxide, isoflurane, propofol, barbiturates, diazepam, and other anticonvulsants all increase apoptotic neurodegeneration in developing rat brain.2–5 We and others have replicated some of the studies with ketamine, and there is no doubt regarding the scientific validity of their findings.6–7

The direct applicability of these experimental findings to the clinical practice of pediatric anesthesia and critical care, however, should be questioned. Likewise, the implication that all types of anesthetic and sedative agents may have similar potentially deleterious effects on neuronal development in neonates is premature and inappropriate. Multiple lines of evidence cast doubt on the clinical relevance of these experimental paradigms, as recently reviewed in our Special Article published in the August 2004 issue of ANESTHESIOLOGY.8 Dr. Olney et al.9 eloquently provided a counterpoint editorial in the same issue of ANESTHESIOLOGY, which mainly questioned our assertion that repeated large doses of ketamine were responsible for mediating the neurodegenerative changes noted in neonatal rat brains. Similar doses and durations of administration, particularly in the absence of surgical stimulation, are never used in pediatric anesthesia.

Recent data from the Neurotoxicology division at the National Center for Toxicological Research (Jefferson, Arkansas) further confirm our findings that the single- or smaller-dose regimens of ketamine do not increase neuronal apoptosis in the brains of 7-day-old (P7) rat pups.7 Scallet et al.9 examined the effect of various dosing regimens on the different assays for neuroapoptosis in P7 rat pup brains. They reported no differences in the appearance of apoptotic neurons in rat pups receiving saline, seven doses of ketamine (10 mg/kg) over 9 h, and a single dose of ketamine (20 mg/kg), whereas rat pups receiving higher doses, seven doses of ketamine (20 mg/kg) over 9 h, had significant increases in neuronal apoptosis. Scallet et al. found that the magnitude and pattern of neuronal changes were similar to those reported earlier.2,6 Unlike Olney et al., these investigators also measured blood concentrations of ketamine to show that the repeated low-dose (10 mg/kg) and single-dose regimens resulted in the blood concentrations achieved in clinical practice, whereas the repeated high-dose regimen (20 mg/kg) resulted in sevenfold higher ketamine concentrations. Furthermore, Scallet et al. verified the presence of neurodegeneration with several histochemical methods, including cu-
Fig. 1. Wechsler Preschool and Primary Scale of Intelligence score measures at 4 and 8 yr after surgical repair of complex congenital heart defects. Population norms are 100 ± 10. There were no significant differences between the study groups and the population norms. Data compiled from Bellinger et al.21,22 and de Ferranti et al.23 WISC-III = Wechsler Intelligence Scale for Children–III.

We agree with Drs. Olney and Todd that human clinical studies are needed to truly examine the potential neurotoxic effect of anesthetic drugs in the developing human brain.9,19,20 Given the ethical and societal constraints, randomized controlled trials mirroring the experimental designs used in rodent models are not possible. We examined children at 4 and 8 yr after the surgical repair of congenital heart defects as neonates using a standardized anesthetic regimen including high-dose barbiturates and opiates.21–23 Despite the known neurologic sequelae of cardiopulmonary bypass and deep hypothermic circulatory arrest,24 their performance IQ scores were not significantly different between the study groups and population norms. Data compiled from Bellinger et al.21,22

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In Reply.—In their letter to the editor, Soriano et al. state that in our recent counterpoint editorial,1 we “mainly questioned [their] assertion that repeated large doses of ketamine were responsible for mediating the neurodegenerative changes noted in neonatal rat brains.” This is not what we questioned, nor do we agree that repeated large doses of ketamine can cause extensive neuroapoptosis. What we mainly questioned was whether repeated large doses are necessary, or whether neuroapoptosis can be triggered by a single low dose of ketamine. We then presented evidence that a single subcutaneous dose of ketamine (20, 30, or 40 mg/kg)—one that does not fully immobilize, anesthetize, or abolish pain responses in infant mice—does trigger a significant increase in the rate of neuroapoptosis. The scientifically appropriate response would be for Soriano et al. to administer these single subanesthetic doses of ketamine to infant mice and, using the same methods we used, to see whether they could reproduce our findings. Instead, Soriano et al. imply that it requires “repeated large doses” to trigger neuroapoptosis and argue that ketamine is safe for pediatric anesthesia because such large “doses and durations . . . are never used in pediatric anesthesia.”

To bolster their claim that a single dose is ineffective, they cite a recent report of Scallet et al.2 in which a single dose of ketamine at 20 mg/kg did not trigger apoptosis in infant rats, although repeated 20-mg/kg doses did. Because, in our single-dose experiments, 20 mg/kg was the threshold dose for triggering apoptosis, it is not surprising or very meaningful that one laboratory would report a barely significant result and another would report a barely insignificant effect at this dose. What is surprising is that instead of directly acknowledging and discussing the implications of our finding that a single dose of ketamine at 20, 30, or 40 mg/kg does trigger apoptosis in a dose-dependent manner, Soriano et al. continue to promote their original position that ketamine is safe because, in their hands, a single dose as high as 75 mg/kg does not trigger neuroapoptosis. It is difficult to reconcile this position with their introductory statement that “there is no doubt regarding the scientific validity” of our findings.

Soriano et al. suggest that we should have measured blood ketamine concentrations in our mouse experiments. However, Soriano et al. did not measure ketamine blood concentrations in their rodent experiments, and anesthesiologists do not routinely measure, much less rely on, ketamine blood concentrations to determine depth of anesthesia. We reported that a single dose of ketamine, in the range of 20–40 mg/kg, that does not fully immobilize, anesthetize, or render an infant mouse insentient to pain does trigger neuroapoptosis in the infant mouse brain. This is a message that is not difficult to understand. Presumably, we can all agree that regardless of ketamine blood concentrations, it would be unacceptable to perform surgery on an infant mouse, or infant human, whose depth of anesthesia is such that the infant is squirming around, failing the extremities, and responding to skin pinch by vigorous antalgic movements.

Soriano et al. continue to argue that the neuroapoptosis response to anesthetic drugs is due to hypoxia/ischemia. How is this possible in light of our demonstration3 that arterial oxygen saturation remains in the 97–99% range over a 4-h period after a dose of ketamine that triggers neuroapoptosis within this same time interval? Soriano et al. postulate that the oxygen saturation fleetingy decreased to brain-damaging levels during intervals between our sampling time points but abruptly resumed normal levels at each time point (15, 30, 60, 120, 180, 240 min) just before we drew blood. We doubt that the readership of Anesthesiology will be persuaded by this argument, especially because we are talking about a subanesthetic dose of ketamine, a drug that reputedly, even at anesthetic doses, does not compromise cardiorespiratory function.

Even if severe hypoxia/ischemia did occur, it could not account for the neuroapoptosis response to ketamine because 4–6 h after ketamine administration, an increase in apoptotic profiles is evident both as a caspase-3 activation response and as ultrastructurally confirmed apoptotic morphology. However, when one intentionally induces hypoxia/ischemia and examines the developing brain 4–6 h later, there is no increase in apoptotic profiles, either by caspase-3 activation or ultrastructural criteria. It is illogical to argue that anesthesia-induced apoptosis is caused by hypoxia/ischemia if one cannot demonstrate that intentionally induced hypoxia/ischemia reproduces the anesthesia-induced apoptosis phenomenon. What one does find in the brain 4–6 h after hypoxia/ischemia, as we have demonstrated previously,4 and also very recently,5 is excitotoxic neurodegeneration. (See Young et al.6 for a detailed presentation of evidence directly addressing and clarifying this issue.) Soriano et al. challenge our position by citing works from other laboratories that they believe contradict our observations. We have examined the cited works, some of which are in vitro studies, and find that these works either support our position or are irrelevant to the issue in contention. We stand by our own observations, which are based on a three-decade-long direct investigation of the specific issue in contention: in vitro excitotoxic versus apoptotic neurodegeneration in the developing brain.7–10

Regarding the nutritional deprivation issue, we stated1 that in a “typical” experiment, we expose infant rodents to a single dose of saline or an apoptogenic anesthetic drug and, without returning the pups to the maternal nest, kill them 4–8 h later. Because both the control and experimental pups are exposed to the same degree of maternal/nutritional deprivation during this 4- to 8-h period, nutritional deprivation cannot explain the higher rate of neuroapoptosis in the experimental pups. In an apparent effort to refute this interpretation, Soriano et al. note that in one study we killed animals not only at 4 and 8 h but also at 12, 16, 24, and 48 h, and determined, using a staining procedure that detects cumulative neuronal degeneration, that apoptosis became increasingly more prominent at 12, 16, and 24 h. We do not understand how this reference to our earlier comprehensive evaluation of the apoptotic response to large doses of MK80111 refutes our current interpretations pertaining to “typical” experiments focusing on the very early response to low subanesthetic doses of ketamine.

Soriano et al. point out that our most recent findings12 pertaining to threshold conditions for inducing developmental neuroapoptosis were conducted in mice and suggest that species differences between rats and mice and between rodents and humans may be of paramount importance. We have tested rats and mice and find no appreciable differences between these species, but we agree that differences between rodents and primates may be very important. Of course, species differences can go in either direction—humans may be less vulnerable or they may be more vulnerable than rodents.

Soriano et al. conclude that only human studies can provide the final answer. We do not contest the importance of human experiments, but such experiments will require many years to complete and, because of design limitations, may provide equivocal results that defy interpretation. Therefore, we recommend that the issue be addressed in nonhuman primate studies designed to test the sensitivity of the primate brain to anesthesia-induced developmental neuroapoptosis. If the pri-
mate brain proves sensitive. Additional studies designed to evaluate the neurobehavioral consequences of graded degrees and controlled patterns of apoptotic neurodegeneration in the developing nonhuman primate brain would be informative. Such studies would provide the anesthesiology community with reliable information and guidance in the conduct of much-needed human studies.

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Richard B. Weiskopf, M.D., served as Handling Editor for this letter, the following letter by Willis and Martin, and the reply by Rathmell et al.

To the Editor:—I read with interest the article by Rathmell et al. regarding cervical transforaminal injection of steroids. An important safety issue not addressed was the need to minimize complications by avoiding excessive sedation. Hodges et al. reported two cases of nerve injury after cervical epidural steroid injections, both performed in heavily sedated patients using fluoroscopy. Excessive sedation may result in the inability of the patient to experience and report pain and paresthesias at the time of spinal cord or nerve root contact. In addition, some recommend that cervical injections should only be performed by experienced and well-trained practitioners. 3

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(Accepted for publication December 20, 2004.)

A Safer and More Effective Intervention for Radiculopathic Pain

To the Editor:—We thank Dr. Rathmell et al. for emphasizing the potential hazards of transforaminal injections. 1 It is clear that these injections should be performed by individuals who are fully trained in advanced imaging and interventional techniques. Moreover, the practitioner must be capable of managing any adverse sequelae.

Although the treatment of radiculopathic pain with the local injection of corticosteroid is appreciated, we believe that the best treatment of radiculopathic pain is by the application of pulsed radiofrequency current to the involved dorsal root ganglion. In our extensive experience with pulsed radiofrequency, we have found the results of treatment to be superior to those of conventional corticosteroid injections in both effectiveness and duration. 2,3

Furthermore, pulsed radiofrequency application exposes patients to less risk for the following reasons: (1) Cannula placement can be performed based on advanced imaging and interventional techniques. Moreover, the practice of pulsed radiofrequency can be repeated as indicated, without fear of accumulating medicinal toxicity.

The only potential disadvantage to the use of pulsed radiofrequency versus injection therapy is the requirement of a larger cannula (20–22 vs. 26 gauge) that could cause more tissue trauma. Regardless, the take-home message is well elucidated by the authors. Spinal interventional techniques should only be performed by practitioners who have demonstrated expertise in neuraxial imaging for interventional treatment modalities.

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In Reply:—Critical to the safety of cervical transforaminal injection of steroids is an understanding of the anatomy of the cervical intervertebral foramina and their contents, coupled with disciplined and accurate imaging. We thank both Dr. Gajraj and Drs. Willis and Martin for emphasizing that cervical transforaminal injection should be performed by experienced and well-trained practitioners. Indeed, the radiographic anatomy of the cervical spine is difficult to master. We have all watched talented physicians-in-training get confused by small changes in alignment on the x-ray image caused by positioning or rotation of the neck and stray dangerously off-course during needle placement. Image-guided injection in the cervical spine requires advanced and extended training under the guidance of an experienced practitioner; as we have emphasized before, this is not something that can be mastered through a weekend cadaver course. 1

Even with the best technique in skilled hands, minimal changes in needle direction and depth can lead the tip into contact with neural structures. After the needle is in proper position, the volume of the injectate itself can cause painful neural compression. We have emphasized the need to maintain an awake and responsive patient when performing cervical epidural steroid injection via a translaminar route as the only safe means to avoid injury, 2 and we thank Dr. Gajraj for raising this point because it is equally relevant to any type of neural blockade.

As for Drs. Willis and Martin’s advocating pulsed radiofrequency treatment of the dorsal root ganglion as a superior technique for treating cervical radicular pain, we point out that there is little evidence to support their assertion. Small, randomized trials of conventional radiofrequency treatment do not result from actual tissue destruction caused by voltage fluctuations in the area of treatment that induce long-term changes in the dorsal horn of the spinal cord. 3 The appeal of pulsed radiofrequency treatment is immediately clear: a simple treatment that imparts long-term pain relief without tissue destruction. However, we do not have even a single randomized trial that compares the efficacy of pulsed radiofrequency to any type of control treatment or to conventional radiofrequency treatment. We hope that the evidence will soon appear to support the unbridled zeal of practitioners for this new treatment. We urge those like Drs. Willis and Martin who have extensive experience with these techniques to conduct the randomized trials we need to demonstrate the effectiveness (or lack thereof) of pulsed radiofrequency treatment.

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Endotracheal Tube Damage during Head and Neck Surgeries as a Result of Harmonic Scalpel® Use

To the Editor.—The Harmonic Scalpel® (Ethicon, Somerville, NJ) and laser are precise cutting and coagulating surgical devices. These devices are widely used worldwide for endoscopic and open surgical procedures. We describe our experience with the Harmonic Scalpel® during the anesthetic management of head and neck oncurgery.

A 58-yr-old male with postradiation recurrence of tongue and soft palate carcinoma was scheduled for wide excision and functional neck dissection with the Harmonic Scalpel® under general anesthesia. After induction of general anesthesia, direct laryngoscopy was performed and an 8 mm cuffed endotracheal tube (ETT) was introduced nasotracheally under vision. Maintenance of anesthesia was achieved with opioids, muscle relaxants, and positive pressure ventilation with 66% nitrous oxide, 34% oxygen, and 0.8% halothane delivered by anesthesia ventilator. The patient remained stable and was closely monitored while the surgeons proceeded with dissection using the Harmonic Scalpel®. While the soft palate lesion was being dissected, the ventilator suddenly emitted the low airway pressure alarm and stopped functioning. The surgical team also noticed bubbling of blood inside the oral cavity. Damage to the ETT was suspected and the head end of the operating table was immediately lowered to prevent aspiration of blood that had pooled in the oropharynx. The fraction of inspired oxygen was increased to 60% and manual ventilation was attempted after thorough suctioning of the oral cavity. However, this was unsuccessful due to the leak (fig. 1), and a fresh ETT of the same size was introduced with the aid of a tube exchanger. The patient was reversed at the end of surgery and the ETT was retained overnight. The postoperative period was uneventful.

The blade of the Harmonic Scalpel® vibrates at 55,000 Hz. It cuts and coagulates tissue at temperatures much lower than either lasers or traditional electrocautery. As a result, the risk of airway fires should be reduced. However, as shown here, it is clear that the Harmonic Scalpel®—like a laser—can cause accidental damage to the ETT when used in the pharynx.

The existing literature abounds with techniques to protect the airway.
To the Editor—I wish to report a little-known and potentially dangerous mechanism of interference between Extraneal® (Baxter Healthcare Corporation, Deerfield, IL) peritoneal dialysis solution and the Accu-Chek® (Roche Diagnostics, Basel, Switzerland) blood glucose test strip monitor. Extraneal® is being integrated into continuous ambulatory peritoneal dialysis regimens because of its increased ultrafiltration and its extralong dwell (time a solution resides in the abdomen). Unfortunately, this new solution interferes with most modern capillary glucose strip–based measuring devices, including the Accu-Chek® brand. The test strip devices can dangerously overestimate the true blood glucose, potentially leading to erroneous treatment and hypoglycemia. A full year before the Food and Drug Administration approved Extraneal® in the United States in December 2002, a case series in a British diabetes journal highlighted this concern. Several patients in that report experienced symptomatic hypoglycemia although their glucose strip machine reported a normal or even increased blood sugar. This year, a case report published in Diabetes Care, the journal of the American Diabetes Association, described a patient on Extraneal® who fell into a hypoglycemic coma secondary to this interference in capillary blood glucose measurement. As anesthesiologists, we must be aware of this potentially lethal monitoring malfunction, especially because during anesthesia, there may be no other clinical warnings of hypoglycemia.

The mechanism behind this interference is quite interesting. Icodextrin, a starch-derived, water-soluble glucose polymer, is the osmotically active colloid in the Extraneal® formulation. Up to 40% of indwelling icodextrin is systemically absorbed and then metabolized by α-amylase into several oligosaccharides, including maltose, maltotriose, and maltotetrose. Although the serum metabolite concentration peaks at the end of the long dwell (approximately 12 h after infusion into the peritoneum), metabolites remain in the circulation for a full 7 days after the last dwell. Many handheld blood glucose monitors use a glucose dehydrogenase–based method to determine glucose concentration, and both maltose and maltotriose interfere with this test, leading to a falsely increased reported value. In contrast, laboratory-based blood glucose–quantifying machines typically use the glucose oxidase-based method, which does not interfere with icodextrin metabolites, thus providing an accurate measurement. In conclusion, before relying on handheld glucose monitors, we must be sure that the specific monitor is compatible with icodextrin-based peritoneal dialysis.

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To the Editor.—Placement of a labor epidural or combined spinal-epidural in advanced labor is technically challenging. Regular painful contractions often make it difficult for the parturient to remain still during epidural placement, and this may increase the chance of an accidental dural tap or nerve injury. Decreasing the intensity and frequency of uterine contractions during neuraxial placement in this setting may be advantageous. Previous reports show that nitroglycerin produces rapid effective uterine relaxation.1–5 Nitroglycerin to facilitate the placement of a labor epidural has not previously been reported. This case describes the use of nitroglycerin in the setting of advanced labor to facilitate the placement of a labor epidural.

A 35-yr-old, healthy, 90-kg, gravida 3, para 2 parturient admitted to labor and delivery in advanced labor requested an epidural for pain relief. She had two previous uncomplicated normal vaginal deliveries without the use of a labor epidural. A recent cervical examination showed her cervix to be dilated 8 cm, with the fetal head at +1 station. She was moving and uncooperative during contractions, which occurred every 45–60 s. She had not responded to 100 μg intravenous fentanyl given 5 min previously. With difficulty, we managed to position her in a sitting position to administer the epidural after a sterile preparation of her back with a 10% povidone-iodine solution and 1% lidocaine skin infiltration, we attempted to insert the epidural catheter using a 17-gauge Tuohy needle. However, she kept moving and was uncooperative during and between uterine contractions. After informing the obstetrician and the patient that we were going to administer medication to help ease the painful contractions, we administered three sprays (400 μg per spray dose) of sublingual nitroglycerin (Nitrolingual™; Pumpspray; First Horizon Pharmaceutical Corporation, Alpharetta, GA). This produced a temporary decrease in her uterine contractions (reduction in peak uterine pressures and an increased between-contraction interval as measured by external tocodynamometer) and resulted in some transient pain relief. It was then possible to perform the combined spinal–epidural during the interval between contractions. The patient experienced no hypotension or cardiovascular disturbances after administration of the nitroglycerin and resumed her normal uterine contraction pattern within a few minutes. The patient was delivered of a healthy baby vaginally 2 h later, with 1- and 5-min Apgar scores of 8 and 9, respectively.

Reducing contraction pain during placement of a labor epidural is potentially beneficial. However, the risks of uterine tocolysis must be balanced with the potential benefit of safer epidural placement and labor analgesia. Although there have been no studies demonstrating increased dural puncture or neural damage after epidural placement in an uncooperative and moving parturient, most clinical anesthesiologists believe that a relation must exist. Decreasing the intensity and frequency of uterine contractions during neuraxial placement in this setting should be potentially advantageous. Remifentanil has been described in this setting to improve analgesia and facilitate the insertion of a labor epidural.6 However, potent narcotics have potential adverse effects, in particular maternal apnea, dysphoria, and emesis. Nitroglycerin is a safe, effective uterine tocolytic commonly used in labor, with a rapid onset and brief half-life.2,7 Nitroglycerin has minimal, short-lived cardiovascular effects compared with β-adrenergic tocolytics. Although the safety of nitroglycerin during obstetric emergencies seems high, with no adverse maternal or neonatal outcomes,2 maternal hypotension and hemodynamics changes are possible, especially if high doses are given.8 No more than three metered sprays are recommended within a 15-min period.9 A number of studies and case reports describe the use of nitroglycerin in achieving rapid uterine relaxation.2–5,10 Nitroglycerin has been used as a tocolytic to reduce uterine hyperactivity,10 assist reduction of an inverted uterus,11 facilitate intrapartum external cephalic version,12 and manage preterm labor contractions.9 Nitroglycerin can be administered via a number of routes (intravenous, sublingual, or ointment); however, bioavailability is highly variable between subjects because of a pronounced first-pass metabolism. After sublingual administration, bioavailability is approximately 38%.14 Nitroglycerin may be useful in a setting where advanced labor and parturient movement during uterine contractions makes the placement of an epidural difficult and potentially dangerous. It exposes the mother and fetus to minimal risk and, in selected patients, may offer potential benefits justifying its use in this setting. However, physicians should remember that this is ‘off-label’ use of nitroglycerin.9 Both the risks and the benefits must be considered, and the patient and her obstetrician must be consulted before uterine tocolytics are administered in this setting.

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To the Editor—Complications due to internal jugular vein (IJV) cannulation are infrequent and rarely life threatening. However, inadvertent carotid artery puncture can lead to serious problems in patients who have atheromas or bleeding disorders or who are undergoing full anticoagulation therapy, as for cardiopulmonary bypass. An external vascular ultrasound technique, using a vascular probe positioned on the neck, has previously been described as an aid for IJV cannulation. We have found useful in our practice an alternative method, using a transesophageal probe.

During the past year, we selected 50 cardiac patients for whom jugular vein cannulation could present a risk. Patients who required monitoring with a transesophageal probe and who had carotid artery disease, previous carotid artery surgery, and difficult anatomy, such as unclear landmarks or no palpable venous pulse, were selected. The mean age of the patients was 72.3 ± 8.8 yr (median, 74 yr).

Transesophageal echocardiographic monitoring is performed with use of a multiplane transducer and a Sonos 5500 imaging system (Hewlett Packard, Andover, MA). Induction of anesthesia and tracheal intubation are performed before insertion of the echo probe. A small towel is placed under the patient’s shoulders. The head is then extended and turned slightly to the side opposite the cannulation, and the patient’s arms are placed by his or her sides. The patient is positioned in a 25° head-down position. The ultrasound monitor is placed in front of the operator.

The transesophageal echo probe is inserted 12–20 cm from the teeth. The tip is directed along the pharyngeal lateral wall, which is why we call this method transpharyngeal. The probe is then rotated laterally 15–20° until the cervical vascular bundle is seen. The view is a mirror image of that obtained from conventional vascular ultrasonography (fig. 1).

A needle covered with a plastic hood is used to search the skin surface of the neck for the best site of cutaneous insertion to find the IJV, which is not pulsating and does not compress. The ultrasound probe is kept stable on a trolley. The operating field, the operator, and the devices are then prepared as usual under sterile conditions.

The skin puncture site is that nearest the IJV, and the puncturing needle is directed away from the carotid artery. The needle is inserted at a 60° angle to the neck axis and directed to the middle of the vein while it is observed on the monitor. Insertion of the needle through the vein may be seen directly, or it may be seen indirectly as movement of the vein wall (fig. 1). Aspiration of blood in the syringe confirms the proper needle position. As the probe is left in place, no adjustment is needed to maintain the correct view.

The IJV puncture was successful in 100% of the patients studied. No carotid punctures or other immediate complications occurred. Conventional ultrasound-guided cannulation of the IJV significantly improves the success rate, decreases the access time, and reduces the complication rate of cannulation. Meta-analyses and a systematic review of control data from literature were performed and suggested an advantage of ultrasonography in complicated cases and when access problems were anticipated. At our institution, the widespread use of transesophageal echocardiography, with its ability to provide a view of the vascular bundle of the neck, offers the anesthesiologist a simple way to aid in central venous cannulation, without any additive cost.

Potential advantages of transpharyngeal ultrasonography in comparison with conventional ultrasonography in intubated patients undergoing transesophageal echocardiographic monitoring are as follows: Direct compression on the IJV by the external probe is not needed; other professionals do not need to be involved in the procedure; the operator’s hands are free; and the ultrasound probe, still working in the patient, can be used for other purposes.

Further studies are needed to assess the reliability of this procedure for IJV cannulation, to determine its proper indications, and to compare this technique with other methods.

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Fig. 1. Transpharyngeal, short-axis view of the neck vessels. The direction of the needle is away from the carotid artery (CA), and the tip of the needle is inside the internal jugular vein (IJV).
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