GW280430A

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.7 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 histamine9 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 anesthetica 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 advanced14 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 presentations1–3 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 anesthetists15–16) would like to encourage the recent attempts to develop better muscle relaxants (or reversal drugs, e.g., Org 25969) to improve safety and efficiency of neuromuscular treatment during anesthesia.

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GW280430A: Pharmacodynamics and Potential Adverse Effects

To the Editor.—We read with interest Dr. Caldwell’s recent editorial1 accompanying the articles published on GW280430A.2–4 Although we agree with his assessment of its potential advantages, we have several concerns with respect to his comments regarding its adverse effects.

GW280430A is a potent, nondepolarizing neuromuscular blocking agent with a fast onset of effect and a duration of action similar to that of succinylcholine.5–4 When administered as a rapid intravenous bolus to volunteers in doses greater than or equal to 2.8 times the ED<sub>95</sub>, it...
may cause histamine release and subsequent hemodynamic changes, a decrease in blood pressure, and an increase in heart rate. As reported by Belmont et al., such changes were observed in 25% of volunteers receiving a dose of 2.8 times the ED₉₅ and 75% of volunteers receiving administration of 2.8 times the ED₉₅ of mivacurium. An average increase in HR of 7% occurs with doses lower than those recommended for mivacurium.5 Doses of 2.5 times the ED₉₅ of GW280430A do not cause histamine release even when administered over 5 s. Furthermore, because 90% suppression of T₁ occurs within 1.3 min after administration of 0.36 mg/kg GW280430A, tracheal intubation may likely be accomplished in 60 s when administered as a bolus (5 s) into a rapidly flowing intravenous line. In the study of mivacurium,6 the neuromuscular blocking agent was injected three times more slowly (15 s).

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 like to 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 Nondepolarizing neuromuscular blocking agents can interact with two of the three muscarinic 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,4 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.4 A closer look at the data in the study of Heerdt et al.5 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 all other nondepolarizing relaxants that may be introduced into clinical practice. Based on the data published to date,4,5 there is no reason to anticipate that GW280430A may even rarely cause life-threatening bronchospasm.

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Fig. 1. The mean maximal heart rate (HR) and blood pressure changes after bolus administration of GW280430A2 and mivacurium. Decreases in HR occur at doses of 2.5 times the ED₉₅ of mivacurium. An average increase in HR of 7% occurs with administration of 2.8 times the ED₉₅ of GW280430A, and an average increase of 14% occurs after administration of 3.8 times the ED₉₅ of GW280430A. Decreases in MAP occurred at a dose of 3.8 times the ED₉₅. The average maximal decrease at this dose of GW280430A was 18%. At all doses, GW280430A was administered as a bolus (5 s) into a rapidly flowing intravenous line. To put this further into perspective, as demonstrated in figure 1, the average maximal heart rate and blood pressure changes after administration of GW280430A are less than those observed after administration of mivacurium, when mivacurium is injected slowly (over 15 s). Doses of 2.5 times the ED₉₅ of GW280430A do not cause histamine release even when administered over 5 s. Furthermore, because 90% suppression of T₁ occurs within 1.3 min after administration of 0.36 mg/kg GW280430A, tracheal intubation may likely be accomplished in 60 s with doses lower than those recommended for mivacurium.

We did not enroll volunteers with a history of cigarette smoking or recent experience (the withdrawal of rapacuronium) amply demonstrate that it is prudent to exercise caution when evaluating the clinical potential of experimental drugs.

Before proceeding, I must make an apology for and clarify confusing language in my editorial. The text as written, ‘Is there evidence of such a potential problem with GW280430A?,’ is clearly open to misinter-
pretation. I did not intend to suggest that GW280403A produced bronchospasm; there is no evidence for that. Rather, my intent was to emphasize that any adverse effect observed during clinical trials may assume far greater significance when the drug goes into general use. The experience with rapacuronium was a dramatic illustration of this point. In the case of GW280403A, the adverse effects raising concern are its propensity to release histamine and to produce hypotension.2

I differ with Dr. Lien et al. in the interpretation of the significance of the adverse hemodynamic effects of GW280403A. For example, take their statement “As is typical of cardiovascular effects due to histamine release, these hemodynamic changes were all self-limited and required no pharmacologic treatment.” I do not believe that the word typical can be used to characterize histamine-related effects. These effects can vary from trivial, localized, cutaneous flushing to life-threatening cardiovascular collapse. The fact that in a small sample of healthy young volunteers, who can increase heat rate to compensate for histamine-mediated vasodilation, a blood pressure decrease of nearly 40% was observed is a matter for concern.2 The obvious question is how severely might this degree of histamine release manifest in patients with much less cardiovascular reserve.

In addition, the fact that the blood pressure changes required no pharmacologic treatment in this limited subject population cannot be used to predict safety in the general population of patients. The subjects studied were all healthy young volunteers in whom a period of hypotension might be tolerable. Allowing the persistence of hypotension without treatment would not be an option for many patients undergoing anesthesia.

Will the doses of GW280403A causing histamine release and hypotension overlap with the doses that might be used in clinical practice? Dr. Lien et al. suggest not by speculating that GW280403A in dose of 0.36 mg/kg (2 × ED95) will provide good intubating conditions in 60 s. Because the onset time of 0.36 mg/kg GW280403A ranged up to 2 min in this small, healthy, and homogeneous group of subjects, it is difficult to believe that intubation within 60 s could be reliably achieved in the general patient population with this dose. It is likely that larger doses will be required to facilitate rapid tracheal intubation. As I described in my editorial, there is ample evidence that clinicians tend to increase doses of neuromuscular blocking drugs to achieve better and faster intubating conditions.3–5 Therefore, it is feasible that clinicians might use GW280403A in the dose range 3–4 × ED95, and the work by Dr. Lien et al. shows that at these doses, histamine release and hypotension can occur.2

Finally, it is not justified to infer the cardiovascular safety of GW280403A by using 15-yr-old historic comparisons with mivacurium.6 There is no way to ensure the comparability of the study procedures or subjects. The weakness of this comparison is further compounded by the use of “average maximal changes” as the variables for comparison. This averaging masks larger changes in individual subjects, and it is these that have the most clinical significance.

In summary, I do not believe that bronchospasm is a significant clinical issue with GW280403A, but histamine release and hypotension most assuredly are.

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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. Caldwell7 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 × ED95.3 The nonvagolytic properties of AV430A found in cats1 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.10 have shown that mivacurium has minimal interactions with M2 and M3 receptors9 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,7 the imbalanced blocking property on muscarinic receptors in the human airway, which is the likely mechanism of rapacuronium-induced bronchospasm,10 is a most unlikely clinical scenario in the case of AV430A.

What about histamine release? Both Caldwell4 and Geldner and Blobner express concern about this, but in the rehus monkey, the safety ratio of AV430A for this side effect is approximately four times greater than that of mivacurium.1 This results in the following comparison in humans: mivacurium, at 2.5–4 × ED95 (0.20–0.25 mg/kg), causes greater symptomatology of histamine release when injected over 15 s11 than does AV430A when injected over only 5 s at 3–4 × ED95 (0.5 mg/kg).7 Since the original study in human volunteers by Belmont et al.7 in 1998, AV430A has been reformulated. This reformulation has further improved the safety ratio for histamine release to approximately 4 × ED50 in humans.7 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 ED50 for neuromuscular blockade: ED Hist/ED50.1

Both Caldwell4 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 evoked twitch response, 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 × ED95 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).7 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,7 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.12 have suggested, a dose of 2.0 or 2.5 × ED95 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.13 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.

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Fig. 1. A recording of evoked twitch (at 0.15 Hz) and train-of-four (2 Hz for 2 s every 10 s) of the adductor pollicis in an anesthetized human volunteer. AV430A (0.4 mg/kg) was given intravenously at the arrow. Time scale (minutes) at top. Note change of time scale and change of evoked response from single twitch to train-of-four at minute 7. Onset of block is 80 s. Recovery to train-of-four of 95% is 13.5 min.
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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**The Right Thing in the Right Place: Lumbar Plexus Block in Children**

To the Editor—I read with interest the article “Lumbar Plexus in Children” by Lukas Kirchmair et al. published in the August 2004 issue of *Anesthesiology*. The authors have nicely shown that sonography of the lumbar plexus in children is feasible, they have clarified the anatomical understanding, and they have given additional information on the depth of the lumbar plexus in pediatric patients.

In addition, they applied their technique to five children scheduled to undergo inguinal hernia repair and concluded that all lumbar plexus blocks provided effective anesthesia and analgesia of the inguinal region during surgery and for postoperative pain relief. Although I am unable to check the clinical effects in these five individual patients, I have considerable doubt that lumbar plexus block is a suitable anesthetic technique for inguinal hernia repair and would not recommend it for the following reasons:

First, the iliopsoas and iliopectineal nerves arise from the roots T12 and L1, and the ilioinguinal nerve arises from L1. These two nerves are usually not densely blocked by injections made at an L4–L5 level. Even the genitofemoral nerve, arising from L1 and L2, is often missed.

Second, a successful lumbar plexus block is followed by a profound motor blockade of all knee extensor and most hip adductor muscles. This is clearly an undesirable side effect after inguinal hernia repair. Avoiding this by using a small volume and a low concentration of the local anesthetic does not seem to be a valuable strategy, especially when nerves far away from the injection site must be blocked.

Third, clinically, in awake adolescents, using the posterior lumbar plexus block with an injection at the L4–L5 level in combination with a sciatic nerve block, incisions coming high up, close to the inguinal ligament, are often troublesome. The ilioinguinal and genitofemoral nerves are often not sufficiently blocked.

In summary, lumbar plexus block can be used in children. However, its use should be restricted to indications where this technique is clearly effective, e.g., for lower limb procedures involving parts of the hip or extensive knee surgery, and where no other less invasive alternative exists.

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**Reference**


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In Reply—we thank Dr. Jörh for his comments on our article published in the August 2004 issue of *Anesthesiology*. He pointed out important anatomical features regarding lumbar plexus anatomy that might raise concerns about the indications of ultrasound-guided posterior lumbar plexus blocks in our series of pediatric patients.

The lumbar plexus is derived from the ventral rami of the roots L1–L5. The latter and its branches are arranged in a potential space within the substance of the psoas major muscle, the so-called psoas compartment. The iliopsoas, ilioinguinal, and genitofemoral nerves are the most cranial branches and provide innervation of the inguinal region. They arise from the ventral rami of the roots T12–L1, L1, and L1–L2, respectively. Therefore, a posterior lumbar plexus block by means of an approach at the level L4–L5 would probably result in an incomplete or missing block of these nerves, as mentioned by Dr. Jörh.

In adults, the iliopsoas and ilioinguinal nerves approximately exit the psoas major muscle at L1–L2, and the genitofemoral nerve usually exits at L4. Nevertheless, the detailed topographic anatomy and arrangement of these nerves within the psoas major muscle in children is unknown.

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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.

To the Editor—Over the years, Dr. Olney et al. 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.6–9 We and others have replicated some of the studies with ketamine, and there is no doubt regarding the scientific validity of their findings.6–9

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 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. 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 rats 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 from the population norms (fig. 1). Certainly their neurologic outcome was no worse than that of peer groups with similar congenital heart defects repaired later in life and exposed to anesthetic agents after the “vulnerable” neonatal and infant periods of brain development at issue in the study of Newburger et al.25 The longer operative repair (and anesthetic exposure) was postponed after infancy in this study population, the greater the cognitive impairment was. These findings suggest that even under extreme conditions encountered in the conduct of pediatric anesthetics, neonatal exposure to anesthetic agents does not necessarily confer substantial neurologic deficits and is, at worst, only one factor among many potentially influencing brain development.

Regardless of this interpretation of such data, the controversy initiated by the studies of Olney and others should provoke clinical investigators to examine the long-term neurocognitive effects of prolonged exposure to anesthetic drugs in pediatric patients, to help sort out the differences between mice and men. Further knowledge in these areas is required before changes in clinical practices in neonatal and pediatric anesthesia can be recommended; such changes may not be necessary or may even be deleterious to long-term neurologic outcome. We are all indebted to Olney et al. for their contribution in raising these issues.

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In Reply.—In their letter to the editor, Soriano et al. state that in our recent counterpoint editorial, 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. We never suggested 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 subcatastic 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. [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 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, flailing 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 demonstration 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 fleeting 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 cardio-respiratory 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, and also very recently, is excitotoxic neurodegeneration. (See Young et al. 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.

Regarding the nutritional deprivation issue, we stated that a ‘typical’ experiment, we expose infant rodents to a single dose of saline or an apoptogenetic 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 MK801 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 findings 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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To the Editor.—I read with interest the article by Rathmell et al.1 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.2 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

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.

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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.5 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.2,5 Furthermore, pulsed radiofrequency application exposes patients to less risk for the following reasons: (1) Cannula placement can be performed based solely on osseous anatomy and electrophysiologic stimulation results. Therefore, no injection of any material is requisite. (2) The patient is exposed to no systemic sequelae of medication or radiocontrast administration. (3) Unlike injection therapies, 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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(Accepted for publication December 20, 2004.)
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 Drs. 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.

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 of the dorsal root ganglion as a superior technique for blockade.

Critical to the safety of cervical transforaminal injection

References


(Accepted for publication December 20, 2004.)

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 oncoscupry.

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 anesthetic, direct laryngoscopy was performed and an 8 mm cuffed endotracheal tube (ETT) was introduced nasopharyngeally under vision. Maintenance of anesthetic 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 oropharynx.

The existing literature abounds with techniques to protect the airway from damage to the endotracheal tube. However, it is always a good practice to maintain an awake and responsive patient when performing transforaminal injection of steroids. 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.

Critical to the safety of cervical transforaminal injection...
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.1 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 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.2 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.3 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, maltohexaose, 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 maltohexaose 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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(Accepted for publication November 11, 2004.)
Nitroglycerin to Facilitate Insertion of a Labor Epidural

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 facil-
tate the placement of a labor epidural has not previously been re-
ported. 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 oc-
curred every 45–60 s. She had not responded to 100 µg intravenous
fentanyl given 5 min previously. With difficulty, we managed to pos-
tion 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 inform-
ing 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 (Ni-
trolingual® Pumpspray; First Horizon Pharmaceutical Corporation, Al-
pharetta, 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 tocomynamom-
eter) 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 cardiovascu-
lar 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 anesthesiolo-
gists 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
recommended within a 15-min period.9 However, potent narcotics have 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 adminis-
tered in this setting.

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(Accepted for publication November 24, 2004.)
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 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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