

Chronic Local Therapy for Brainstem Tumors

To the Editor:

Opinions have been reported about the value and safety of chronic infusions for brainstem tumor therapy. These opinions may influence planning for clinical trials for a disease that relentlessly claims the lives of approximately 200 children yearly in the United States alone.

Median survival for children with diffuse pontine gliomas is approximately 10 months (1). Trials with small numbers of patients from single institutions have generated the consensus that neither chemotherapy nor radiation prolongs survival and that new approaches are needed (8, 9).

Local therapy seems to be a logical response (7). Two strategies are available. Both have demonstrated efficacy in animal studies. Both have provided baseline safety data via human trials in supratentorial tumors (5). The methods differ in their rate of intratumoral delivery. Fast infusions pump at rates of 1 $\mu\text{l}/\text{min}$ or more. Convective gradients distribute molecules homogeneously several centimeters from the point of delivery. Slow infusions release drugs from biodegradable implants or from pumps at rates of 1 $\mu\text{l}/\text{h}$ or less. Drugs diffuse several millimeters from the point of delivery in normal tissues (6, 11, 12, 16).

Conjectures about the clinical irrelevance of slow infusions derive from the opinion that distributive capacities are too small to invade the tumor volumes generally encountered in clinical practice. Some have reported that the modest "statistical" gains seen in clinical trials with slow infusions further attest to the irrelevance of this strategy (7, 23).

Laboratory models have demonstrated the safety of fast brainstem infusions in normal tissues (15, 19, 21). Questions remain about clinical safety in abnormal tissue. Elegant and technically challenging studies in a rat supratentorial glioma model have demonstrated that intracranial pressure increases with increasing tumor volumes as well as the volume and flow rate of a fast infusion (4). Data in *Table C1* show the lethal effects of increasing infusion rates in the brainstems of normal mice and rats. The data on rat infusions have been published (6). We think that it is important to add the unpublished mouse data to further illustrate the stark relationship between morbidity and increasing flow rates. Brainstem infusions of saline in 180- to 220-g Fischer rats at 10 $\mu\text{l}/\text{h}$ were lethal in four of five rats in 7 days. Although a child's brainstem is significantly larger, fast chronic infusion in a brainstem potentially challenged by increased pressures from a glioma could lead to unintended events.

Fast infusions with targeted toxins have shown promise in Phase I trials with supratentorial tumors (13, 17, 20). Increased intracranial pressure, whether related to drug toxicity or tumor necrosis, seemed to be manageable. However, there is a lack of information about the management of increased pressure in the brainstem (2). Children are a vulnerable population (14). Trials will require an independent committee to monitor

TABLE C1. Survival and infusion flow rates in rat and mice brainstems

Species and pump (μl)	Flow rate ($\mu\text{l}/\text{h}$)	No. of animals surviving			
		Day 2	Day 7	Day 14	Day 30
Mouse					
200	8	0/3			
200	1	1/3	0/3		
200	0.5	2/3	1/3	1/3	1/3
200	0.25	2/3	2/3	2/3	2/3
Rat					
2000	10	3/5	1/5	0/5	
2000	5 ^a	9/10	5/10	3/10	3/10
2000	2.5 ^a	10/10	9/10	9/10	9/10
200	1	5/5	5/5	5/5	5/5

^a Treated with 1 mg dexamethasone per day intraperitoneally for 3 days postsurgery.

treatment effects (10, 18). With the information at hand, the committee could not distinguish potentially toxic effects of increased pressure from the progression of the disease.

Clinical trials can begin with slow infusions. Theoretical equipoise, the null hypothesis, states that there will be no difference between the infusion of saline, drug escalation, and palliative treatment for a child with a pontine glioma. Infusions at 0.42 $\mu\text{l}/\text{h}$ of 1 ml of saline (90 d) have been performed with no apparent toxicity (23). Regardless of the initial skepticism about volumes of distribution from slow infusions, the multiple reports of efficacy shown in human trials demonstrate the clinical relevance of this approach in neurosurgical oncology (3, 22, 24–26). Short-term methodological safety has been documented. Side effects, which frequently blunt systemic drug trials, have been greatly reduced. Clinical equipoise considers both the risks and benefits of a new therapy. The safety and efficacy of slow infusions have been achieved with first-generation chemotherapeutics such as carmustine and cisplatin. Randomized controlled trials can continue the search for progress using faster infusion rates and newer generations of chemotherapeutics, including targeted toxins.

Michael Guarnieri
Benjamin S. Carson, Sr.
Baltimore, Maryland

- Abbott R, Goh KYC: Brainstem gliomas, in Albright L, Pollack I, Adelson D (eds): *Principles and Practices of Pediatric Neurosurgery*. New York, Thieme Medical Publishers, 1999, pp 629–640.
- Avellino AA, Carson BS Sr: Increased intracranial pressure, in Maria BL (ed): *Current Management in Child Neurology*. New York, B.C. Decker, 2002, pp 481–486.

3. Brem H, Piantadosi S, Purger PC, Walker M, Selker R, Vick NA, Black K, Sisti M, Brem S, Mohr G, Muller P, Morawetz R, Schold SC: Placebo-controlled trial of safety and efficacy of intraoperative controlled delivery by biodegradable polymers of chemotherapy for recurrent gliomas. *Lancet* 345:1008–1012, 1995.
4. Bruce JN, Falavigna A, Johnson JP, Hall JS, Birch BD, Yoon JT, Wu EX, Fine RL, Parsa AT: Intracerebral clysis in a rat glioma model. *Neurosurgery* 46:683–691, 2000.
5. Carson BS Sr, Guarnieri M: Local therapy for brain tumors. *Adv Clin Neurosci* 12:89–99, 2002.
6. Carson BS, Wu QZ, Tyler B, Sukay L, Raychaudhuri R, DiMeco F, Clatterbuck R, Olivi A, Guarnieri M: New approach to tumor therapy for inoperable areas of the brain: Chronic intraparenchymal drug delivery. *J Neurooncol* 60:151–158, 2002.
7. Dunn IF, Black PMcL: The neurosurgeon as a local oncologist: Cellular and molecular neurosurgery in malignant glioma therapy. *Neurosurgery* 52: 1411–1424, 2003.
8. Freeman CR: Hyperfractionated radiotherapy for diffuse intrinsic brain stem tumors in children. *Pediatr Neurosurg* 24:103–110, 1996.
9. Freeman CR, Perilongo G: Chemotherapy for brain stem gliomas. *Childs Nerv Syst* 15:545–553, 1999.
10. Freidman LM, Furberg CD, DeMets DL: *Fundamentals of Clinical Trials*. New York, Springer-Verlag, 1998.
11. Fung LK, Ewend MG, Aills A, Sipos EP, Thompson R, Watts M, Colvin OM, Brem H, Saltzman WM: Pharmacokinetics of interstitial delivery of carmustine, 4-hydroperoxycyclophosphamide, and paclitaxel from a biodegradable polymer implant in the monkey brain. *Cancer Res* 58:672–684, 1998.
12. Kroin JS, Penn RD: Intracerebral chemotherapy: Chronic microinfusion of cisplatin. *Neurosurgery* 10:349–354, 1982.
13. Laske DW, Youle RJ, Oldfield EH: Tumor regression with regional distribution of the targeted toxin TF-CRM107 in patients with malignant brain tumors. *Nat Med* 3:1362–1368, 1997.
14. Levine RJ: *Ethics and Regulations of Clinical Research*. New Haven, Yale University Press, 1988.
15. Lonsner RR, Walbridge S, Garmestani K, Butman JA, Walters HA, Vortmeyer AO, Morrison PF, Brechbiel MW, Oldfield EH: Successful and safe perfusion of the primate brainstem: In vivo magnetic resonance imaging of macromolecular distribution during infusion. *J Neurosurg* 97:905–913, 2002.
16. Lum JT, Nguyen T, Felpel LP: Drug distribution in solid tissue of the brain following chronic local perfusion utilizing implanted osmotic minipumps. *J Pharmacol Methods* 12:141–147, 1984.
17. Mardor Y, Roth Y, Lidar Z, Jonas T, Preffer R, Maier SE, Faibel M, Na D, Hadani M, Orenstein A, Cohen JS, Ram Z: Monitoring response to convection-enhanced taxol delivery in brain tumor patients using diffusion-weighted magnetic resonance imaging. *Cancer Res* 61:4971–4973, 2001.
18. Meinert CL: *Clinical Trials: Design, Conduct, and Analysis*. New York, Oxford University Press, 1986.
19. Occhiogrosso G, Edgar MA, Sandberg DI, Souweidane MM: Prolonged convection-enhanced delivery into the rat brainstem. *Neurosurgery* 52:388–394, 2003.
20. Rand RW, Kreitman RJ, Patronas N, Varricchio F, Pastan I, Puri RJ: Intratumoral administration of recombinant circularly permuted interleukin-4-*Pseudomonas* exotoxin in patients with high-grade gliomas. *Clin Cancer Res* 6:2157–2165, 2000.
21. Sandberg DI, Edgar MA, Souweidane MM: Convection-enhanced delivery into the rat brainstem. *J Neurosurg* 96:885–891, 2002.
22. Sheleg SV, Korotkevich EA, Zhavrid EA, Muravskaya GV, Smeyanovich AF, Shanko YG, Yurkshtovich TL, Bychkovsky PB, Belyaev SA: Local chemotherapy with cisplatin-depot for glioblastoma multiforme. *J Neurooncol* 60:53–59, 2002.
23. Storm PB, Clatterbuck RE, Liu YJ, Johnson RM, Gillis EM, Guarnieri M, Carson BS Sr: A surgical technique for safely placing a drug delivery catheter into the pons of primates: Preliminary results of carboplatin infusion. *Neurosurgery* 52:1169–1177, 2003.
24. Valtonen S, Timonen U, Toivanen P, Kalimo H, Kivipelto L, Heiskanen O, Unsgaard G, Kuurme T: Interstitial chemotherapy with carmustine-loaded polymers for high-grade gliomas: A randomized double-blind study. *Neurosurgery* 41:44–49, 1997.
25. Wenig BL, Werner JA, Castro DJ, Sridhar KS, Garewal HS, Kehrl W, Pluazanska A, Arndt O, Costantino PD, Mills GM, Dunphy FR, Orenberg EK, Leavitt RD: The role of intratumoral therapy with cisplatin/epinephrine injectable gel in the management of advanced squamous cell carcinoma of the head and neck. *Arch Otolaryngol Head Neck Surg* 128:880–885, 2002.
26. Westphal M, Hilt DC, Bortey E, Delavault P, Olivares R, Warnke PC, White IR, Jääskeläinen J, Ram Z: A phase 3 trial of local chemotherapy with biodegradable carmustine (BCNU) wafers (Gliadel wafers) in patients with primary malignant glioma. *Neuro-oncology* 5:79–88, 2003.

In Reply:

Drs. Guarnieri and Carson raise important issues regarding local therapy for brainstem tumors (1). Convection-enhanced drug delivery is rapidly being validated as a safe and useful strategy for delivery of antitumor compounds to brain tumors. This has led to examination of its potential application for tumors in more anatomically hazardous areas, including the brainstem.

Morbidity from local intratumoral infusion can come from either direct drug toxicity, which is influenced by local drug concentration and duration of infusion, or mass effect, which is influenced by volume delivered and preexisting mass effect from the tumor itself. With brainstem lesions, Drs. Guarnieri and Carson are particularly concerned with infusion rates and volumes associated with convection-enhanced delivery and present evidence that increased flow rates can lead to significant morbidity in rodent models. Their work has led them to advocate for more chronic infusions at considerably lower flow rates than those that produce convective forces. Their letter highlights the difficulties in interpreting experimental studies in animals, in which brain volumes are a fraction of human brain volumes.

Given the limitations of animal models, I agree with the need to examine local delivery parameters more fully in clinical trials incorporating careful empirical protocol designs. Intratumoral infusion studies will need to focus on three variables: 1) drug concentration, 2) infusion rate, and 3) duration of infusion (which along with infusion rate determines volume of distribution). It would be prudent to begin these studies slowly, using safe parameters that could be increased gradually. These types of study designs are similar to what has traditionally been used when testing new systemic chemotherapies.

Given the limited treatment options available for patients with malignant brainstem tumors, it is reasonable to propose that these trials begin now. Experienced teams of investigative neurosurgeons, neuro-oncologists, and radiologists will be needed to devise reasonable treatment protocols with careful dose and infusion escalation parameters that will ultimately determine whether convection-enhanced or chronic slow infusion can be efficacious for brainstem pathology. These studies would be greatly assisted by noninvasive radiological imaging to monitor treatment end points for safety and response. Monitoring may be the greatest challenge, because early clin-

ical trials with convection-enhanced delivery have been hindered by the difficulties in interpreting radiographic imaging studies of these heterogeneous tumors, which have often undergone previous surgical, radiotherapeutic, and chemotherapeutic intervention.

Jeffrey N. Bruce
New York, New York

1. Bruce JN, Falavigna A, Johnson JP, Hall JS, Birch BD, Yoon JT, Wu EX, Fine RL, Parsa AT: Intracerebral clysis in a rat glioma model. *Neurosurgery* 46:683–691, 2000.

DOI: 10.1227/01.NEU.0000117119.32806.AF

The Impact of Provider Volume on Mortality after Intracranial Tumor Resection and Outcome and Cost of Craniotomy Performed to Treat Tumors in Regional Academic Referral Centers

To the Editor:

There is growing evidence that patients with intracranial neoplasms should be treated in highly specialized neurosurgical centers. This statement was clearly confirmed by the authors of both referenced articles (1, 2), who found direct correlation between the volume of the neurosurgical activity for brain tumors and early outcome, namely, mortality rate and length of in-hospital stay. Several years ago, we reached a similar conclusion in our study of the same problem. However, despite publication in a MEDLINE-cited journal (3), our results seem to be obscured from the neurosurgical community, and therefore, it may be of interest to highlight our study briefly.

We performed a population-based study of the outcome after surgical removal of primary intracranial tumors in adult citizens of St. Petersburg, Russia. Patients with metastatic and recurrent neoplasms had been excluded from the analysis. The early outcome in all patients (n = 307) who were treated during 1993 was identified retrospectively, with an emphasis on postoperative morbidity and mortality. There were 148 men and 159 women (mean age, 48 yr). At the time of hospital admission, the condition of 54.7% of the patients was considered to be compensated, 31.9% subcompensated, and 13.4% decompensated. Gliomas were encountered most frequently (n = 134), followed by meningiomas (n = 87), pituitary adenomas (n = 39), and acoustic schwannomas (n = 15). Surgical treatment was performed in 13 different neurosurgical departments. The mean volume of the surgical activity, defined as number of brain tumor removals per year, was 23.6 (range, 3–104 operations). The mean morbidity rate constituted 19.5% (range, 6.7–66.7%). The mean mortality rate was 9.4% (range, 0–33.3%). A strong dependence of the morbidity and mortality rates on the surgical volume was found (Fig. C1). The mathematically defined breakpoint of both curves corresponded to the surgical volume of 41 cases/year. Compar-

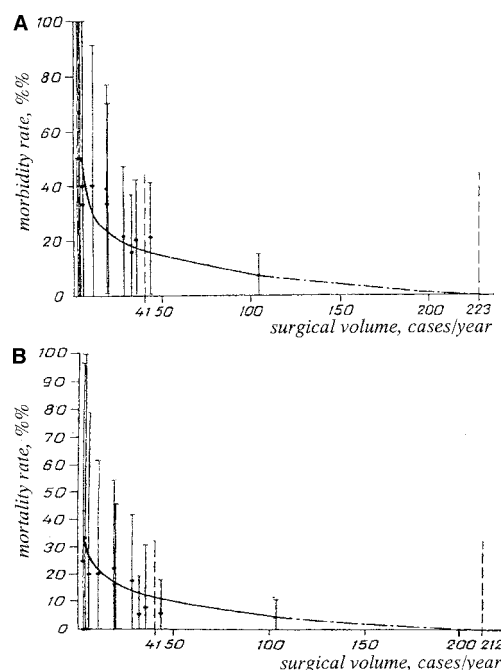


FIGURE C1. Graphs depicting the morbidity rate (A) and mortality rate (B) after surgical removal of intracranial tumors versus the volume of surgical activity of the neurosurgical department (adapted from, Tigliev GS, Ulitin AY, Chernov MF: The dependence of the results of the surgical treatment of patients with primary intracranial tumors on the volume of surgical activity of the neurosurgical department [exemplified by Saint Petersburg, Russia] [in Russian]. *Zh Vopr Neurokhir Im N N Burdenko* 2:44–46, 1999 [3]).

ison of the low-volume hospitals (surgical volume <41 cases/yr) and high-volume hospitals (surgical volume >41 cases/yr) revealed relatively worse clinical condition of the patients and higher frequency of critical tumor location in the latter group, whereas the mean age did not differ significantly. The morbidity curve reached its minimal value at the surgical volume of 223 cases/year, whereas the mortality curve reached its minimal value at 212 cases/year.

It seems that our results, which are in complete concordance with the conclusions of the referenced authors, may carry some additional information to the interested reader. First, not only mortality rate but also morbidity rate after removal of brain tumors depends on the volume of surgical activity, which probably reflects the lower average length of in-hospital stay at the high-volume centers found by Long et al. (2). Second, in our study, better results in high-volume neurosurgical departments were obtained despite the relatively worse clinical condition of patients and more frequent critical tumor location. Therefore, better outcome in hospitals with higher surgical volume seems not to correspond to more preferable patient or tumor characteristics. Third, we found that not only the results of surgery for brain tumors but also the quality of the adjunctive neuro-oncological treatment (chemotherapy) is lower in low-volume hospitals (data not shown here).

Another point that seems to be extremely interesting is that nearly the same surgical volume corresponded to mathematically defined breakpoints of both morbidity and mortality curves in our study (41 cases/yr) and defined for categorization of low- and high-volume hospitals by Long et al. (2) (50 cases/yr). Moreover, the curves we constructed reached their minimal values at a surgical volume of more than 200 cases/year, which corresponds to a “high volume of surgical activity” (as defined by authors of both referenced articles) associated with the better outcome. If such similar data were obtained in such different sociological and medical communities, can we consider this similarity as an objective tendency that may be used in the future for better organization of the neurosurgical and neuro-oncological services?

In conclusion, it seems that treatment of patients with intracranial neoplasms in the neurosurgical centers with a high volume of surgical activity can provide lower morbidity and mortality rates compared with low-volume hospitals. However, the dependence of other important outcomes of brain tumor management (for example, radicality of tumor removal, long-term patient survival, and quality of life) on the surgical volume is still unknown. Therefore, I am looking forward to further contributions on this topic.

Mikhail F. Chernov
Tokyo, Japan

-
1. Cowan JA Jr, Dimick JB, Leveque J-C, Thompson BG, Upchurch GR Jr, Hoff JT: The impact of provider volume on mortality after intracranial tumor resection. *Neurosurgery* 52:48–54, 2003.
 2. Long DM, Gordon T, Bowman H, Etzel A, Burleyson G, Betchen S, Garonzik IM, Brem H: Outcome and cost of craniotomy performed to treat tumors in regional academic referral centers. *Neurosurgery* 52:1056–1065, 2003.
 3. Tigliev GS, Ulitin AY, Chernov MF: The dependence of the results of the surgical treatment of patients with primary intracranial tumors on the volume of surgical activity of the neurosurgical department (exemplified by Saint Petersburg, Russia) [in Russian]. *Zh Vopr Neirokhir Im N N Burdenko* 2:44–46, 1999.

In Reply:

We appreciate the comments of Dr. Chernov and his previous contributions, which described a volume-outcome effect after surgical removal of primary intracranial tumors in St. Petersburg, Russia. His work reminds us that variation in outcome after the delivery of complex neurosurgical care is not unique to healthcare systems in the United States.

As mentioned in our article (1), “high” caseload volume has been shown to be a consistent marker for superior outcomes after complex surgery. Increasing hospital volume alone will not guarantee better outcomes (even though, on average, these high-caseload centers have better results). We therefore would refrain from applying specific caseload thresholds but rather would focus on the organizational structures and processes of care that improve outcome, including intensive care unit physician staffing, low nurse-to-patient ratios, access to neuroimaging, optimization of

comorbid conditions, and a multidisciplinary approach to care. These organizational components may currently be found more often at high-volume centers. Whether they are unique to high-volume centers is unknown.

John A. Cowan, Jr.
B. Gregory Thompson
Julian T. Hoff
Ann Arbor, Michigan

-
1. Cowan JA Jr, Dimick JB, Leveque J-C, Thompson BG, Upchurch GR Jr, Hoff JT: The impact of provider volume on mortality after intracranial tumor resection. *Neurosurgery* 52:48–54, 2003.

DOI: 10.1227/01.NEU.0000117120.32806.22

Far Posterior Subtemporal Approach to the Dorsolateral Brainstem and Tentorial Ring: Technique and Clinical Experience

To the Editor:

It was with great interest that I read the description of the far posterior subtemporal approach (4). It closely resembles another modification of the subtemporal approach, namely, the transpetrous supratentorial-anterior approach, which has been used in our clinic since 1992 in more than 70 cases of various tumors of the tentorial notch area (1, 5).

The positioning of the patient, direction of the surgical access, and main steps of the approach we used are the same as described by the authors. We strongly agree with the importance of the head tilt below the horizontal position (which we prefer to do after opening of the dura), mobilization of the vein of Labbé, use of the lumbar or ventricular drain (to our mind, the latter is preferable in cases with associated hydrocephalus), mediotemporal retraction for the visualization of the tentorial notch, and laterally directed tentoriotomy, starting from the tentorial edge and proceeding parallel to the superior petrosal sinus. In addition, it should be emphasized that the increase of the retraction force on the basitemporal lobe must be as gradual as possible, with steady evacuation of the cerebrospinal fluid (CSF) after early opening of the basal subarachnoid cisterns. I need not mention the advantages of adequate neuroanesthesia, which provides optimal brain compliance. All of these measures are of paramount importance for the prevention of complications caused by temporal lobe retraction.

Although the risk of injury to the vein of Labbé should not be underestimated, it should also not be overemphasized. We did not observe this complication in any case in our series, because it seems that careful mobilization of the vessel from the subarachnoid space and dural duplication provides sufficient mobility for safe retraction of the temporal lobe and further surgical manipulations.

However, it should be noted that the amount of retraction necessary during the subtemporal approach to the region of the tentorial notch depends on the individual peculiarities

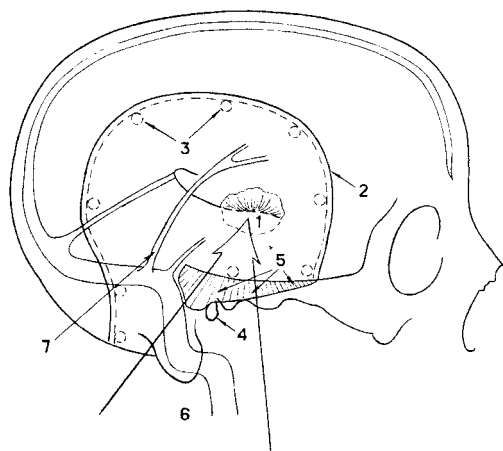


FIGURE C2. Scheme of the transpetrous supratranstentorial-anterior approach: tumor in the region of the tentorial notch (1), skin incision (2), burr holes (3), external acoustic meatus (4), resected posterolateral portion of the petrous bone (5), direction of the surgical access (6), and vein of Labbé (7) (adapted from, Chernov MF: Basal meningiomas with both supra and infratentorial extension: Clinical picture, diagnosis, management [in Russian]. St. Petersburg, Russian Polenov Neurosurgical Institute, 1995 [dissertation] [1]).

of the neuroanatomy, namely, the angle between the tentorium and the upper plane of the petrous bone. The smaller the angle, the larger the retraction force needed. In this way, the avoidance of complications can be provided by resection of the posterolateral part of the petrous bone up to the level of the external acoustic meatus (Fig. C2). Such limited resection is not associated with the risk of damage of the intrapetrous structures but increases the angle of the surgical access 60% on average (3) and therefore should be kept in mind if the retraction force on the temporal lobe during the subtemporal approach seems to be too high. Although further extension of the resection on the glenoid fossa has been proposed (2), it seems to be unnecessary in the vast majority of cases.

Finally, the authors have provided a good description of the modified subtemporal approach, which represents one of the main routes to the region of the tentorial notch. The only concern I have is about the proposed name of the approach. Because the lateral border between the temporal and occipital lobes is assumed to be the preoccipital notch, which lies a bit posterior to the termination of the vein of Labbé, it seems that “midsubtemporal” rather than “far posterior subtemporal” is a more appropriate definition of the access.

Mikhail F. Chernov
Tokyo, Japan

1. Chernov MF: Basal meningiomas with both supra- and infratentorial extension: Clinical picture, diagnosis, management [in Russian]. St. Petersburg, Russian Polenov Neurosurgical Institute, 1995 (dissertation).

2. Goel A: Extended lateral subtemporal approach for petroclival meningiomas: Report of experience with 24 cases. *Br J Neurosurg* 13:270–275, 1999.
3. Sindou MP, Fobe JL: Removal of the roof of the external auditory meatus in approaching the tentorial notch through a low temporal craniotomy: Technical note. *J Neurosurg* 74:520–522, 1991.
4. Smith ER, Chapman PH, Ogilvy CS: Far posterior subtemporal approach to the dorsolateral brainstem and tentorial ring: Technique and clinical experience. *Neurosurgery* 52:364–369, 2003.
5. Tigliev GS, Chernov MF: The characteristics of the surgical treatment of basal meningiomas with supra- and subtentorial spread [in Russian]. *Zh Vopr Neirokhir Im N N Burdenko* 1:3–6, 1998.

In Reply:

We thank Dr. Chernov for his thoughtful comments regarding the surgical approach to the posterior lateral brainstem region (1). He is indeed correct that it closely resembles the transpetrous supratranstentorial-anterior approach. As pointed out by Dr. Chernov, as familiarity with this approach is gained, one can mobilize the vein of Labbé. This is one of our most feared potential complications; however, if handled with care, the vein can remain intact and complications associated with venous manipulation can be avoided.

Christopher S. Ogilvy
Paul H. Chapman
Boston, Massachusetts

1. Smith ER, Chapman PH, Ogilvy CS: Far posterior subtemporal approach to the dorsolateral brainstem and tentorial ring: Technique and clinical experience. *Neurosurgery* 52:364–369, 2003.

DOI: 10.1227/01.NEU.0000117121.32806.6B

Seizures after Aneurysmal Subarachnoid Hemorrhage Treated with Coil Embolization

To the Editor:

I read with interest the article by Byrne et al. (1) on seizures after coil embolization of ruptured aneurysms and its relevance to current driving regulations in the United Kingdom. The authors successfully demonstrated that the actual rate of epilepsy after subarachnoid hemorrhage in the selected group is lower than previously thought. The interventional technique of treating aneurysms undoubtedly has the advantage of eliminating the inherent risk of epilepsy caused by craniotomy, which is added to that resulting from the subarachnoid hemorrhage itself. However, the authors' cohort over 10 years may perhaps have a selection bias whereby the patients selected also may be at low risk for epilepsy. Considering an increasing trend in the United Kingdom to treat ruptured aneurysms primarily by interventional means, more patients who are at high risk for epilepsy who previously would have been considered unsuitable are now likely to be considered for coiling. Therefore, the current and future patients treated interventionally may not necessarily follow the same pattern as demonstrated by the authors. It would be worthwhile to analyze whether the same low rate of epilepsy

continues over the next 1 or 2 decades before any definitive conclusions can be drawn.

In our center, we do not use prophylactic anticonvulsant therapy in any patients with subarachnoid hemorrhage, and we agree entirely with the authors' conclusion of not administering prophylactic anticonvulsant medications. We also share an experience of a low incidence of epilepsy similar to that noticed by the authors but including the aneurysms treated by surgery. My own personal experience of the last 100 cases of surgically treated acutely ruptured aneurysms shows a 3% incidence of postoperative epilepsy (unpublished observations). Up to half of these patients are those with intracerebral clots, middle cerebral territory aneurysms, multiple aneurysms, and giant or difficult aneurysms considered technically unsuitable for coiling. All of these factors, I believe, are epileptogenic themselves. Unfortunately, the results achieved in any single-institute or single-operator study may not reflect those of the entire population so as to warrant any changes in driving guidelines on a national level.

Recanalization and rebleeding from coiled aneurysms is another important factor that needs to be considered before a person with an apparently protected aneurysm is allowed to drive. It was recently shown that aneurysm recanalization is the major limitation of current Guglielmi detachable coil treatment (2). Another recently published study has also clearly established that initially incompletely occluded aneurysms, aneurysms treated in the acute phase after rupture, and the length of follow-up are independent significant predictors of major aneurysm recurrences that may possibly lead to a rebleed (3). It is reasonable to believe that a rebleed during driving may carry significant morbidity and mortality, making it equally or perhaps more hazardous than a stand-alone seizure. Because most workers would prefer to treat recanalized or recurred aneurysms before they rebleed, the precise rebleeding rate after acutely coiled aneurysms may never be known. Therefore, unless multicenter studies unequivocally confirm combined rates of long-term angiographic recanalization or recurrence rate as well as epilepsy after coiling lower than those currently considered acceptable for epilepsy alone, it would be hard to justify modifying the current driving legislation in our country for coiled cases only, as suggested by the authors.

Kishor A. Choudhari
Belfast, United Kingdom

In Reply:

Dr. Choudhari makes several interesting observations about current practice and advice to patients about driving motor vehicles after aneurysmal subarachnoid hemorrhage. His comments are well received and welcome. We entirely accept his points concerning selection bias and the limitations of single-center studies such as ours (and his) (1). But the implications of the possibility of rebleeding of treated aneurysms require analysis. The experience of the past decade has shown that the vast majority of patients are protected against rebleeding by coil embolization, despite an appreciable angiographic failure rate. The question is: does the latter warrant imposing driving restrictions more onerous than those for patients managed by surgical clipping?

Dr. Choudhari argues that current evidence is insufficient (in terms of epilepsy and rebleeding risks) to relax the driving restrictions "for coiled cases alone." But the United Kingdom Driver and Vehicle Licensing Authority restrictions imposed, at the time our article was published, were more restrictive for aneurysmal subarachnoid hemorrhage patients managed by coil embolization than for those treated by craniotomy and clipping (for aneurysms other than at the middle cerebral artery). Guidance updated in January 2000 states that a Group 1 driver (i.e., nonprofessional) treated by embolization should "cease driving until full clinical recovery and undergo annual medical review for 4 years," whereas the guidance for a patient treated by craniotomy with no deficit states "driving permitted when clinically recovered from craniotomy" (i.e., without subsequent reviews). We are pleased to report that, after our publication, the advice for patients treated by coil embolization has been amended so that patients are now required only to "cease driving until clinically recovered" (2).

The lack of clear thinking on this issue is exemplified by the advice to patients who sustain aneurysmal subarachnoid hemorrhage and whose aneurysm is not treated by either clipping or coiling. Such patients are required to stop driving for 6 months only. We assume that the authority has taken the view that rebleeding after 6 months is infrequent and does not pose a risk to the patient or other road users. We would argue that to impose more onerous driving restrictions on embolized patients requires evidence to show that this form of management makes patients more liable to rebleeding or seizures than those treated by aneurysm clipping or managed conservatively.

James V. Byrne
Jane Adcock
Oxford, England

1. Byrne JV, Boardman P, Ioannidis I, Adcock J, Traill Z: Seizures after aneurysmal subarachnoid hemorrhage treated with coil embolization. *Neurosurgery* 52:545-552, 2003.
2. Murayama Y, Nien YL, Duckwiler G, Gobin YP, Jahan R, Frazier J, Martin N, Viñuela F: Guglielmi detachable coil embolization of cerebral aneurysms: 11 years' experience. *J Neurosurg* 98:959-966, 2003.
3. Raymond J, Guilbert F, Weill A, Georganos SA, Juravsky L, Lambert A, Lamoureux J, Chagnon M, Roy D: Long-term angiographic recurrences after selective endovascular treatment of aneurysms with detachable coils. *Stroke* 34:1398-1403, 2003.

1. Byrne JV, Boardman P, Ioannidis I, Adcock J, Traill Z: Seizures after aneurysmal subarachnoid hemorrhage treated with coil embolization. *Neurosurgery* 52:545-552, 2003.
2. Drivers Medical Group, Driver and Vehicle Licensing Agency: *At a Glance Guide to the Current Medical Standards of Fitness to Drive*. Swansea, Driver and Vehicle Licensing Agency, 2003.

DOI: 10.1227/01.NEU.0000117122.32806.B0

Factors Related to Hydrocephalus after Aneurysmal Subarachnoid Hemorrhage

To the Editor:

We read with interest the recent article by Dorai et al. (1). We were somewhat surprised by their finding that there was a highly significant ($P < 0.001$) increase in shunt-dependent hydrocephalus for patients experiencing aneurysmal subarachnoid hemorrhage who underwent endovascular obliteration of their aneurysms as opposed to those patients undergoing microsurgical repair.

It seems that there was no statistical discounting of the fact that the endovascular and microsurgical groups were drastically different in nature, making their comparison by use of a χ^2 test invalid. The differences in these two groups in terms of presenting grade, Fisher score, and aneurysm location, for example, are not detailed other than to say that "Of patients treated solely with endovascular methods, 38% demonstrated admission Hunt and Hess grades of IV or V, compared with only 12% of patients who underwent surgical treatment."

It has been shown in multiple reports (including this one) that, overwhelmingly, the major predictors of shunt-dependent hydrocephalus after aneurysmal subarachnoid hemorrhage are Hunt and Hess grade, Fisher grade, and presence of intraventricular blood (2, 4). If the endovascular group had many more patients with higher grades on presentation, this must be accounted for in the analysis. This can be readily performed with a multivariate logistic regression; a simple χ^2 test is not appropriate for this comparison.

Presenting the data in this way is particularly confusing given the fact that the results run counter to those of the few other reports on this subject. In two other prospective randomized studies, patients undergoing microsurgery were found to have a higher incidence of shunt dependence after aneurysmal subarachnoid hemorrhage compared with the endovascularly treated group, and in one other prospective study, no difference between the groups was found (2, 3, 5).

Although the authors identify the multifactorial nature of shunt dependence in the Discussion and acknowledge that their statistical result with regard to endovascular as opposed to microsurgical treatment is probably attributable to their not discounting for these other predictive factors, leaving the statistical result (endovascular versus surgery, $P < 0.001$) in the abstract without clarification is misleading. We, too, would welcome a prospective trial to study this issue.

Jonathan L. Brisman
Alejandro Berenstein
New York, New York

1. Dorai Z, Hyman LS, Kopitnik TA, Samson D: Factors related to hydrocephalus after aneurysmal subarachnoid hemorrhage. *Neurosurgery* 52:763-771, 2003.
2. Gruber A, Reinprecht A, Bavinszki G, Czech T, Richling B: Chronic shunt-dependent hydrocephalus after early surgical and early endovascular treatment of ruptured intracranial aneurysms. *Neurosurgery* 44:503-512, 1999.

3. Koivisto T, Vanninen R, Hurskainen H, Saari T, Hernesniemi JA, Vapalahti MP: Outcomes of early endovascular versus surgical treatment of ruptured cerebral aneurysms: A prospective randomized study. *Stroke* 31:2369-2377, 2000.
4. Vale FL, Bradley EL, Fisher WS III: The relationship of subarachnoid hemorrhage and the need for postoperative shunting. *J Neurosurg* 86:462-466, 1997.
5. Vanninen R, Koivisto T, Saari T, Hernesniemi JA, Vapalahti MP: Ruptured intracranial aneurysms: Acute endovascular treatment with electrolytically detachable coils—A prospective randomized study. *Radiology* 211:325-336, 1999.

DOI: 10.1227/01.NEU.0000117123.32806.F9

Anatomic and Clinical Study of the Orbitopterional Approach to Anterior Communicating Artery Aneurysms

To the Editor:

We read with great interest the article by Andaluz et al. (1) showing that modifications to the pterional approach, consisting of removing the lateral wall and roof of the orbit, widen the exposure and narrow the distance between the surgeon and the operative target. Andaluz et al. nicely present the clinical application of this technical nuance for aneurysms of the anterior communicating artery. We agree that a wider exposure brings the anatomy closer to the surgeon's hands, reducing the length of the instruments, making it easier to manipulate the instruments, and, most importantly, reducing the amount of retraction on the brain.

We previously reported similar results using a different technique (2). Using a robotic microscope and a computerized system to identify cartesian coordinates, we measured the angle of attack (i.e., the corridor between the dura and brain) as a traditional pterional approach was enlarged into an orbitozygomatic approach with an additional maxillary osteotomy and extension. Four different targets were identified during the different approaches (anterior clinoid process, maximal exposure of A2, maximal exposure of M2, and carotid bifurcation). Using these targets as vertices, we defined three triangles that allowed us to calculate the area under the microscope (i.e., working area). Using titanium miniplates to fixate bony osteotomies allowed us to make progressive exposures while maintaining the same amount of brain retraction. Angles were measured for the same increment (10 degrees) as in the angle between the brain and dura (i.e., the projection plane) when the pterional approach was extended into the orbit.

We believe that the concept of a spatial cone developed by Sindou et al. (4) and used by Schwartz et al. (3) and Andaluz et al. (1) is not as critical as the working area. Intuitively, the base of the cone will be wider as bony resection increases. The working area, however, corresponds to the area under the microscope. The larger the working area, the easier it becomes to visualize more distant targets. Using different triangular areas (lateral, superior, and medial), we simulated different tilting movements of the microscope as practiced during surgery.

In conclusion, using a different methodology, Andaluz et al. confirmed that the orbitopterional approach significantly wid-

ens the surgical corridor. They concluded that widening is the key to decreasing complications during surgery for aneurysms of the anterior communicating artery. Their findings and ours are consistent with the primary principles underlying cranial base surgery: wide exposure, proximity to anatomy, and minimal retraction. The authors have shown the clinical impact and relevance of modifying the pterional approach for the treatment of aneurysms of the anterior communicating artery.

L. Fernando Gonzalez
Joseph M. Zabramski
Phoenix, Arizona

1. Andaluz N, van Loveren HR, Keller JT, Zuccarello M: Anatomic and clinical study of the orbitopterional approach to anterior communicating artery aneurysms. *Neurosurgery* 52:1140–1149, 2003.
2. Gonzalez LF, Crawford NR, Horgan MA, Deshmukh P, Zabramski JM, Spetzler RF: Working area and angle of attack in three cranial base approaches: Pterional, orbitozygomatic, and maxillary extension of the orbitozygomatic approach. *Neurosurgery* 50:550–557, 2003.
3. Schwartz MS, Anderson GJ, Horgan MA, Kellogg JX, McMenomey SO, Delashaw JB: Quantification of increased exposure resulting from orbital rim and orbitozygomatic osteotomy via the frontotemporal transsylvian approach. *J Neurosurg* 91:1020–1026, 1999.
4. Sindou M, Emery E, Acevedo G, Ben-David U: Respective indications for orbital rim, zygomatic arch and orbito-zygomatic osteotomies in the surgical approach to central skull base lesions: Critical, retrospective review of 146 patients. *Acta Neurochir (Wien)* 143:967–975, 2001.

In Reply:

It was with great pleasure that we read the letter from Drs. Gonzalez and Zabramski regarding our article (2). In their letter, they reference their cadaveric study (3) that compared the pterional approach with two consecutive maneuvers (i.e., orbitozygomatic and maxillary osteotomies) to expand its exposure. In a very compelling and elegant way, they demonstrated, through data collected with the assistance of a robotic microscope and a computerized stereotactic system, a statistically significant increment in the working area afforded by the addition of an orbitozygomatic osteotomy to the standard pterional approach. However, the addition of a maxillary osteotomy did not significantly increase exposure. This work clearly highlights the value of cranial base approaches, particularly for the treatment of aneurysms of the posterior circulation. In relation to our work, a 10-degree increment in the “angle of attack” was found with the addition of an orbital osteotomy.

In our study, using a different methodology but a similar concept, we focused on the anterior communicating artery region. We achieved an 11-degree increment in the projection plane (equal to the angle of attack by Gonzalez et al.) by taking the midpoint of the anterior communicating artery as a reference. This figure was found not only by Gonzalez et al. (3) but also by Alaywan and Sindou (1), using different landmarks. Unfortunately, we erroneously deleted this datum from the work of Gonzalez et al. during one of our revisions.

In addition to the foregoing, our work on the orbitopterional approach also focused on the extent of exposure afforded in the axial or “circumferential” plane, in which this approach

also demonstrated a statistically significantly improved exposure, and the depth of the surgical field, which decreased but was not statistically significant. What is important is that we could find a clinical correlate of our laboratory findings in a series of 40 patients.

We agree with Gonzalez and Zabramski that what matters is the “working area,” which we translate as “useful surgical area exposed” or “effective surgical exposure.” However, its quantification is not an easy task (1–4). The availability of computerized frameless stereotaxy yields a more refined estimate of what is gained in exposure by cranial base approaches. However, other conditions inherent to cadaveric work, such as brain rigidity and degree of brain atrophy, still compromise data acquisition. For these reasons, we decided to rely primarily on fixed, unmodifiable, and reproducible points (i.e., bony landmarks). Under those premises, the “spatial cone” concept (5) was the most accurate to complement our data.

We envision the ideal comparison between approaches as a volumetric analysis of truncated pyramids with polyhedral bases, with a special interest in the top of the pyramid. The incremental presence of computer-assisted devices in cranial base laboratories may some day bring this near-science-fiction concept into real numbers.

Working area, angle of attack, projection angle, field of view angle, cone of approach, and surgical vector are concepts that convene under the same philosophy of cranial base surgery, that is, minimal brain retraction, better exposure, illumination, and instrument maneuverability. We again thank Drs. Gonzalez and Zabramski for their warm remarks, and we apologize for not including their work in our article.

Norberto Andaluz
Harry R. van Loveren
Jeffrey T. Keller
Mario Zuccarello
Cincinnati, Ohio

1. Alaywan M, Sindou M: Fronto-temporal approach with orbito-zygomatic osteotomy: Surgical anatomy. *Acta Neurochir (Wien)* 104:79–83, 1990.
2. Andaluz N, van Loveren HR, Keller JT, Zuccarello M: Anatomic and clinical study of the orbitopterional approach to anterior communicating artery aneurysms. *Neurosurgery* 52:1140–1149, 2003.
3. Gonzalez LF, Crawford NR, Horgan MA, Deshmukh P, Zabramski JM, Spetzler RF: Working area and angle of attack in three cranial base approaches: Pterional, orbitozygomatic, and maxillary extension of the orbitozygomatic approach. *Neurosurgery* 50:550–557, 2002.
4. Schwartz MS, Anderson GJ, Horgan MA, Kellogg JX, McMenomey SO, Delashaw JB: Quantification of increased exposure resulting from orbital rim and orbitozygomatic osteotomy via the frontotemporal transsylvian approach. *J Neurosurg* 91:1020–1026, 1999.
5. Sindou M, Emery E, Acevedo G, Ben-David U: Respective indications for orbital rim, zygomatic arch and orbito-zygomatic osteotomies in the surgical approach to central skull base lesions: Critical, retrospective review of 146 patients. *Acta Neurochir (Wien)* 143:967–975, 2001.

DOI: 10.1227/01.NEU.0000117124.32806.37

Trigeminal Neuralgia Associated with a Primitive Trigeminal Artery Variant: Case Report

To the Editor:

Tamura et al. (5) present a case of trigeminal neuralgia associated with a primitive (persistent) trigeminal artery (PTA) variant. They conclude that although the PTA variant is frequently associated with intracranial aneurysms, it is extremely rare for the variant to lead to trigeminal neuralgia. They also stress that during microvascular compression surgery, surgeons should be careful to prevent injury to the perforating arteries arising from the PTA variant.

It was Quain (1) who illustrated, as an autopsy study, the first case involving a PTA, and later Sutton (4) did the same as an angiographic study. In 1959, Saltzman (3) proposed an angiographic classification for the PTA. He described Type I as a variation in which the vertebrobasilar system distal to the anastomosis is supplied by the PTA with an incomplete filling of the posterior communicating artery. Type II has bilateral filling of the superior cerebellar arteries by the PTA, whereas the posterior cerebral arteries receive their blood from the posterior communicating arteries. There is a combination of the two types (intermediate type) in which the posterior cerebral artery receives its blood through the posterior communicating artery and the PTA supplies the superior cerebellar arteries on both sides and the posterior cerebral artery on the opposite side. The case presented by Tamura et al. is presumed to be that of a Saltzman Type II.

Which type of PTA may cause trigeminal neuralgia? In one of our recent studies, we found it useful to classify the PTA by its relationship to the abducens nerve, distinguishing the lateral (petrosal) and medial (sphenoidal) variations (2). When the trigeminal artery courses laterally to the abducens nerve, the artery arises from the posterolateral aspect of the C4 segment of the cavernous carotid and crosses underneath the nerve. The abducens nerve may be displaced superiorly by the PTA. This petrosal variation of the PTA pierces the dura just medial to the sensory root of the trigeminal nerve. This type of PTA may compress the trigeminal nerve. When the PTA courses medial to the abducens nerve, the artery arises from the posteromedial aspect of the C4 segment of the cavernous carotid artery and pierces the dura of the dorsum sellae (sphenoid variation). In cases of PTA with trigeminal neuralgia, the artery should be the lateral (petrosal) type, which will affect the sensory root of the trigeminal nerve. In conclusion, according to Saltzman's classification, which is based on the blood supply of the PTA (3), and our classification, which is based on the relationship of the PTA with the cranial nerves (2), the case reported by Tamura et al. may be categorized as Saltzman Type II/lateral (petrosal) type PTA.

Ibrahim M. Ziyal
Osman E. Özcan
Ankara, Turkey

1. Quain R: *The Anatomy of the Arteries of the Human Body and Its Applications to Pathology and Operative Surgery, with a Series of Lithographic Drawings*. London, Taylor & Walton, 1844.

2. Salas E, Ziyal IM, Sekhar LN, Wright D: Persistent trigeminal artery: An anatomic study. *Neurosurgery* 43:557-562, 1998.
3. Saltzman GF: Patent primitive trigeminal artery studied by cerebral angiography. *Acta Radiol* 51:329-336, 1959.
4. Sutton D: Anomalous carotid basilar anastomosis. *Br J Radiol* 23:617-619, 1950.
5. Tamura Y, Shimano H, Kuroiwa T, Miki Y: Trigeminal neuralgia associated with a primitive trigeminal artery variant: Case report. *Neurosurgery* 52:1217-1220, 2003.

In Reply:

We thank Drs. Ziyal and Özcan for their comments on our article (1). The PTA is a most common anomalous vessel with carotid-basilar anastomoses. Conversely, the PTA variant reported by Teal et al. (2) directly supplied the territory of the distal anteroinferior cerebellar artery and/or superior cerebellar artery without a basilar artery anastomosis. In our patient, indeed, the posterior cerebral arteries were filled by the posterior communicating arteries, similar to Saltzman Type II. As Ziyal and Özcan point out, it is thought that the lateral (petrosal) type of PTA running near the trigeminal nerve may cause trigeminal neuralgia. We consider, however, that the PTA variant will be different embryologically from the PTA connected with the basilar artery.

Yoji Tamura
Toshihiko Kuroiwa
Osaka, Japan

1. Tamura Y, Shimano H, Kuroiwa T, Miki Y: Trigeminal neuralgia associated with primitive trigeminal artery variant: Case report. *Neurosurgery* 52:1217-1220, 2003.
2. Teal JS, Rumbaugh CL, Bergeron RT, Scanlan RT, Segall HD: Persistent carotid-superior cerebellar artery anastomosis: A variant of persistent trigeminal artery. *Radiology* 103:335-341, 1972.

DOI: 10.1227/01.NEU.0000117125.32806.7E

Brain Metastases Treated with Radiosurgery Alone: An Alternative to Whole Brain Radiotherapy?

To the Editor:

We read with great interest the recent article by Hasegawa et al. (3) and the comments that followed regarding the treatment of brain metastases with stereotactic radiosurgery (SRS) alone. The article is an excellent addition to the growing body of evidence regarding the use of SRS without whole-brain radiation therapy (WBRT). The management of brain metastases has been evolving with the broader acceptance and application of SRS. WBRT, which has long been the standard palliative treatment for patients with brain metastases, seems to have been superseded by the results of Radiation Therapy Oncology Group 95-08, in which an advantage in overall survival was noted in a significant percentage of patients with one to three brain metastases who were treated with both SRS and WBRT (7). With the establishment of SRS + WBRT as the putative "standard" therapy, the next question that arises is whether WBRT is necessary as an adjunct to SRS.

The present study reaches many of the same conclusions that have been published, including those reached in a retrospective, multi-institutional study in which 268 patients were treated with SRS alone and 301 received SRS + WBRT (6). In that series, after adjustment for known prognostic factors, there was not a significant difference in overall survival. Hasegawa et al. noted that patients treated with SRS alone relapsed elsewhere in the brain at a gross rate of 38%. In another retrospective series, patients receiving SRS alone for cerebral metastases who survived for 1 year were free from relapse in the brain only 28% of the time (5). Despite the high rate of new lesions developing in patients treated with SRS alone, however, overall survival seems to be equivalent to SRS + WBRT, because salvage therapies are fairly effective and patients' extracranial disease is frequently the cause of death (5, 6).

The primary argument for use of SRS alone in the treatment of cerebral metastases is to limit the neurocognitive side effects of radiation therapy. Although there is evidence that WBRT can produce negative neurocognitive sequelae, the available data are not compelling. The two most frequently quoted articles on the topic evaluated a total of 18 affected patients and were published in the late 1980s (1, 2). The patients in the larger series were treated in the late 1970s to the mid-1980s with radiation regimens that were generally not representative of present standards. In fact, 75% of the patients received daily fractions of 5 Gy or greater for some or all of their therapy (2). Despite the widespread perception that WBRT inevitably results in worsened neurocognitive function, the contrary argument can be made that the exclusion of WBRT in patients with brain metastases can lead to inferior cognitive outcomes. Recently published data provide evidence that when patients treated with SRS alone relapse, they are frequently symptomatic (71%), and the majority experience a neurological deficit (59%) (4).

Unfortunately, any retrospective analysis of such a complex issue is intriguing at best but cannot be definitive as a result of inherent selection bias. The authors correctly conclude that a randomized trial including a prospective quality of life and neurocognitive evaluation is the best way to answer the questions raised.

We are pleased to have the opportunity to inform your readership about an open Phase III randomized trial addressing the exact issues previously raised. The American College of Surgeons Oncology Group (ACOSOG) is a relatively recently formed cooperative trial group. ACOSOG is funded by the National Cancer Institute to conduct prospective, randomized clinical trials evaluating surgical therapies in the management of patients with malignant tumors. ACOSOG has activated study Z0300, "a phase III randomized trial of the role of WBRT in addition to radiosurgery in the management of patients with one to three cerebral metastases." Z0300 is the first study opened by the group's Central Nervous System Organ Site Committee. Z0300 is designed to evaluate the role of WBRT in patients with one to three brain metastases treated with SRS. The primary end point of the study is overall survival, because the previous retrospective analyses referenced all showed unavoidable selection

bias. Perhaps equally important, however, are secondary end points that include prospective evaluation of quality of life and neurocognitive function. ACOSOG Z0300 is the first multi-institutional trial to prospectively evaluate the neurocognitive effects of WBRT in which one arm does not receive WBRT. This feature should allow accurate conclusions to be drawn regarding the role of WBRT in the optimal management of patients with one to three brain metastases for not only survival but also quality of life and neurocognitive effects.

The first patient in the trial was enrolled in December 2002, and 34 patients have been enrolled at the time of this writing. At present, more than 20 sites around the country are open to accrue patients. Other institutions are in the process of opening the trial, but additional sites that are interested in helping to answer this important scientific question are encouraged to participate. Further information regarding the ACOSOG Central Nervous System Organ Site Committee or this trial can be obtained by contacting ACOSOG (www.acosog.org) directly or Dr. Anthony Asher, Chair, Central Nervous System Organ Site Committee, ACOSOG (asher@cnsa.com).

Stuart H. Burri
Anthony Asher
Charlotte, North Carolina
Mark Shaffrey
Charlottesville, Virginia

1. Asai A, Matsutani M, Kohno T, Nakamura O, Tanaka H, Fujimaki T, Funada N, Matsuda T, Nagata K, Takakura K: Subacute brain atrophy after radiation therapy for malignant brain tumor. *Cancer* 63:1962-1974, 1989.
2. DeAngelis LM, Delattre JY, Posner JB: Radiation-induced dementia in patients cured of brain metastases. *Neurology* 39:789-796, 1989.
3. Hasegawa T, Kondziolka D, Flickinger JC, Germanwala A, Lunsford LD: Brain metastases treated with radiosurgery alone: An alternative to whole brain radiotherapy? *Neurosurgery* 52:1318-1326, 2003.
4. Regine WF, Huhn JL, Patchell RA, St. Clair WH, Strottmann J, Meigooni A, Sanders M, Young B: Risk of symptomatic brain tumor recurrence and neurologic deficit after radiosurgery alone in patients with newly diagnosed brain metastases: Results and implications. *Int J Radiat Oncol Biol Phys* 52:333-338, 2002.
5. Sneed PK, Lamborn KR, Forstner JM, McDermott MW, Chang S, Park E, Gutin PH, Phillips TL, Wara WM, Larson DA: Radiosurgery for brain metastases: Is whole brain radiotherapy necessary? *Int J Radiat Oncol Biol Phys* 43:549-558, 1999.
6. Sneed PK, Suh JH, Goetsch SJ, Sanghavi SN, Chappel R, Buatti JM, Regine WF, Weltman E, King VJ, Breneman JC, Sperduto PW, Mehta MP: A multi-institutional review of radiosurgery alone vs. radiosurgery with whole brain radiotherapy as the initial management of brain metastases. *Int J Radiat Oncol Biol Phys* 53:519-526, 2002.
7. Sperduto PW, Scott C, Andrews D, Schell MC, Flanders A, Werner-Wasik M, Demas W, Ryu JK, Gaspar LE, Bahary J, Souhami L, Rotman M, Curran WJ: Stereotactic radiosurgery with whole brain radiation therapy improves survival in patients with brain metastases: Report of Radiation Therapy Oncology Group Phase III Study 95-08. *Int J Radiat Oncol Biol Phys* 54s:3, 2002.

In Reply:

We strongly support the proposed randomized trial organized by the American College of Surgeons. We hope to participate in this study with the goal of answering questions regarding the optimal use of radiation techniques for patients

with brain metastasis, particularly as they apply to quality of life (1).

Douglas Kondziolka
Pittsburgh, Pennsylvania

-
1. Hasegawa T, Kondziolka D, Flickinger JC, Germanwala A, Lunsford LD; Brain metastases treated with radiosurgery alone: An alternative to whole brain radiotherapy? *Neurosurgery* 52:1318–1326, 2003.

DOI: 10.1227/01.NEU.0000117126.32806.A5

Intrasphenoidal Encephalocele Associated with Cerebrospinal Fluid Fistula and Subdural Hematomas: Technical Case Report

To the Editor:

We read the article of Fraioli et al. (1) with interest. The authors obtained excellent results treating an intrasphenoidal encephalocele associated with a CSF fistula using a transsphenoidal procedure. There is one technical question that we would like to ask Dr. Fraioli. There is general agreement that dural repair is performed with greater success with a viable graft. If the graft material is fat, the sphenoid mucosa should be removed to allow the revascularization of the fat by feeders coming from the bone. Putting a synthetic dura mater between the bone and the fat compromises the vitality of the graft. Therefore, to support the viable graft, might it not be better to put the synthetic dura mater behind the fat and, in the same session, inflate the pouch?

Diego Mazzatenta
Ernesto Pasquini
Giorgio Frank
Bologna, Italy

-
1. Fraioli B, Conti C, Lunardi P, Liccardo G, Fraioli MF, Pastore FS: Intrasphenoidal encephalocele associated with cerebrospinal fluid fistula and subdural hematomas: Technical case report. *Neurosurgery* 52:1487–1490, 2003.

In Reply:

Our primary purpose in the presented technique of plasty (1) was not to obtain a viable graft. We considered that 1) the CSF leakage was so remarkable that a bilateral subdural hematoma was induced. This required a sealing method able to promptly establish a satisfactory mechanical obstruction, like the inflated pouch described in our article. The amount of rhinoliquorrhea also advised us against using a previously published technique (2) for transmucosal closure of sphenoidal CSF fistulae: we considered this method especially suitable for leaks after transsphenoidal surgery for pituitary adenomas despite an accurate intraoperative plasty; 2) in our opinion, once a prolonged arrest of CSF leakage is obtained, the mechanisms of spontaneous fibrosis should contribute to consolidate the sealant apparatus. This probably happened in our

patient, because the favorable result is still lasting (after 3 yr of follow-up).

Bernardo Fraioli
Francesco Saverio Pastore
Mario Francesco Fraioli
F. Contratti
Rome, Italy

-
1. Fraioli B, Conti C, Lunardi P, Liccardo G, Fraioli MF, Pastore FS: Intrasphenoidal encephalocele associated with cerebrospinal fluid fistula and subdural hematomas: Technical case report. *Neurosurgery* 52:1487–1490, 2003.
2. Fraioli B, Pastore FS, Floris R, Vagnozzi R, Simonetti G, Liccardo G, Giuffrè R: Computed tomography-guided transsphenoidal closure of postsurgical CSF fistula: A transmucosal needle technique. *Surg Neurol* 48:409–412, 1997.

DOI: 10.1227/01.NEU.0000117127.32806.EC

Autotransplantation of Human Carotid Body Cell Aggregates for Treatment of Parkinson's Disease

To the Editor:

I offer the following comments regarding the interesting article by Arjona et al. (1) concerning the bilateral autotransplantation of carotid body (CB) cell aggregates into the striatum in six patients with advanced Parkinson's disease (PD). A moderate neurological improvement was observed in five patients and no change in one. Clinical improvement was better during the first months after the surgery than in the following months. These results confirm the efficacy of autotransplants of CB cell aggregates suggested previously by other authors (2, 4).

In my opinion, this modest improvement was the result of a reduced number of translated CB cells and survival of the graft. First, PD has the highest frequency in older patients, and the regional cerebral blood flow is reduced in the neostriatum (3) and mesencephalic structures (6) of these patients. Second, in situ, all donor tissues of catecholamines (adrenal medulla, fetal mesencephalic tegmentum, cervical sympathetic ganglion, and CB) are normally very vascularized (5), and once they are implanted into the striatum, they release solely noradrenaline and dopamine (Fig. C3). Likewise, it seems that the main product of CB cell aggregates is dopamine (2, 4, 5). Third, in donor tissues, the biosynthesis of catecholamines is related directly to its angioarchitecture and intravascular concentration of the L-tyrosine and molecular oxygen (5, 8). Dopaminergic and/or noradrenergic cells are situated in direct contact with the basement of the fenestrated capillaries, and through these contacts, the catecholaminergic cells receive L-tyrosine, oxygen, and other essential nutrients. Fourth, the cell population in the CB decreases with the age of the patients, especially after age 50 years, because of atrophy or sclerosis (5). Therefore, CB is not a donor tissue acceptable to treat PD, especially for patients with advanced PD.

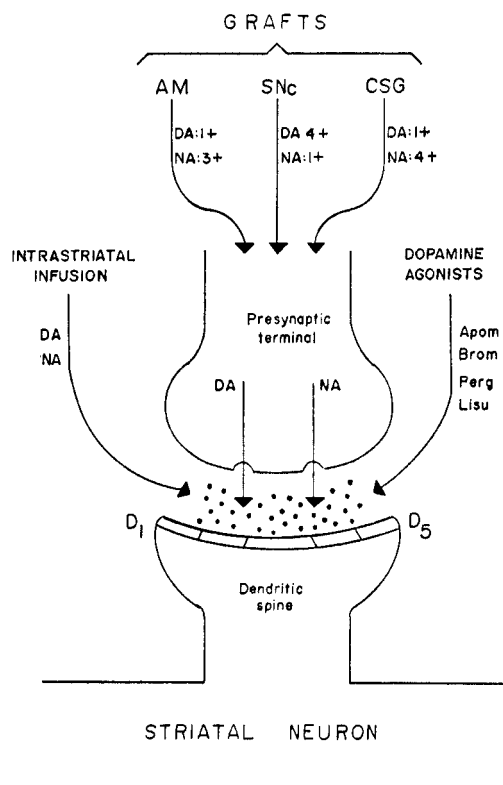


FIGURE C3. Medical (dopamine agonists) and neurosurgical (grafts and intrastriatal infusion) treatments for PD in at least five (D1 to D5) dopamine receptors. DA, dopamine; NA, noradrenaline; Apom, apomorphine; Brom, bromocriptine; Perg, pergolide; Lisu, lisuride; AM, adrenal medulla; SNc, embryonic or fetal substantia nigra; CSG, cervical sympathetic ganglion.

For these reasons, unlike the conclusions of Arjona et al., I have doubts about the security and viability of this therapeutic approach for the treatment of PD, because it is possible that in some PD patients, we may find during the surgery a CB with scarce glomus or epithelioid cells and, by contrast, abundant conjunctive tissue. Moreover, I believe that all catecholamine-producing grafts implanted into the striatum must be revascularized, because a rapid and efficient revascularization of these grafts (Fig. C3) is an essential prerequisite to improve function and prolong survival of the grafts.

Thereby, on the basis of clinical data suggesting that PD is initiated in the intraparenchymal territory of the posterior perforating arteries caused by atherosclerotic plaques located at the mouths of these collateral branches, we proposed two surgical procedures to treat PD (6, 7): 1) omental transplantation on the interpeduncular fossa to revascularize the dopaminergic nuclei and surrounding structures in the early stages of PD, and 2) dual catecholamine-producing tissue and omental transplantation in moderate or advanced stages of PD (i.e., implantation of donor tissues into the neostriatum by a

transinsular pathway and omental transplantation on the anterior perforated space and insular cortex).

Hernando Rafael
Mexico City, Mexico

1. Arjona V, Mínguez-Castellanos A, Montoro RJ, Ortega A, Escamilla F, Toledo-Aral JJ, Pardal R, Méndez-Ferrer S, Martín JM, Pérez M, Katati MJ, Valencia E, García T, López-Barneo J: Autotransplantation of human carotid body cell aggregates for treatment of Parkinson's disease. *Neurosurgery* 53:321–330, 2003.
2. Barinaga M: Unusual cells may help treat Parkinson's disease. *Science* 279:1301, 1998.
3. Leenders KL, Salmon EP, Tyrrell P, Perani D, Brooks DJ, Sager H, Jones T, Marsden D, Frackowiak RS: The nigrostriatal dopaminergic system assessed in vivo by position emission tomography in healthy volunteers subjects and patients with Parkinson's disease. *Arch Neurol* 47:1290–1298, 1990.
4. Luquin MR: Implants of carotid body cells as a treatment alternative for Parkinson's disease [in Spanish]. *Neurología* 14:373–376, 1999.
5. Rafael H: Tejidos donadores de Catecolaminas: Una revisión. *Diagnóstico (Perú)* 34:42–49, 1995.
6. Rafael H: Mesencephalic ischemia and Parkinson's disease. *J Neurol Neurosurg Psychiatry* (in press).
7. Rafael H, Mego R: Ablative surgery and deep brain stimulation for Parkinson's disease. *Neurosurgery* 45:199–200, 1999.
8. Rafael H, Moromizato P, Ayulo V: The dopamine transporter gene and PD in a Chinese population. *Neurology* 52:429–430, 1999.

DOI: 10.1227/01.NEU.0000117128.32806.08

Arctic Explorers

To the Editor:

I very much enjoyed reading the September issue of the Journal, and especially all the rare pictures of Arctic exploration. However, I think it is a misconception that Richard E. Byrd was the first to fly over the North Pole in 1926. For years, he and Floyd Bennett were credited as being the first, just 2 days before Roald Amundsen and Umberto Nobile passed over the Pole in a dirigible. Seventy years later, it was reported that Byrd never reached the Pole, and the credit now belongs to Amundsen. This is documented in the permanent exhibit in the American Museum of Natural History in New York, where pictures of Byrd were replaced by pictures of Amundsen in 1996. Oddly enough, Amundsen always tried to diminish the substantial contributions of his archenemy Nobile, only to lose his life while flying to rescue him from a crash near Spitsbergen in 1928. A few years previously, Amundsen had had a tumor resected from his thigh in San Francisco by Dr. Söyland (Norwegian), who also implanted radioactive material in the wound. Amundsen was only 56 years old when he died, but he looked 20 years older. I wonder whether he was suicidal. He was a former medical student and must have sensed what was coming. In 1998, Helge Ingstad, who discovered the Viking settlements in L'Anse aux Meadows in Newfoundland, told me that years before, Bennett had actually admitted that he and Byrd never passed over the North Pole. Apparently, the powerful Byrd family effectively prevented such rumors from spreading by threatening to sue any com-

pany who dared publish the truth. This includes the memoirs of the Norwegian-American aviator and Arctic explorer Bernt Balchen, who made the first flight over the South Pole with Byrd in 1929. It is a sad fact that whereas Peary, Cook, Byrd, Amundsen, Nansen, and other leaders of polar ventures became immortal, their invaluable companions remained in the shadows and were often forgotten.

Finally, I want to tell you a personal story about Amundsen. When I visited Baffin Island in 1998, I took a trip to Gjoa Haven on King William Land, where Amundsen and his crew spent almost 2 years from 1903 to 1905 trying to navigate the Northwest Passage. Amundsen had introduced strict rules of conduct: his men were not to be engaged in intimacies with the ladies of the Arctic. He detested the local customs of wife swapping and Inuit men offering their wives and daughters "for the price of a rusty nail." He also warned them that many of the local women might have contracted syphilis from vis-

iting whalers. Amundsen would frequently disappear for several days to make weather observations and look for the magnetic North Pole.

In Gjoa Haven, I stayed in the small Hotel Amundsen, where I met an elderly Inuit who looked a little different, with a big nose and fair skin. When he realized that I was a Norwegian, he said:

"My grandfather was Norwegian too."

"What was his name?" I said.

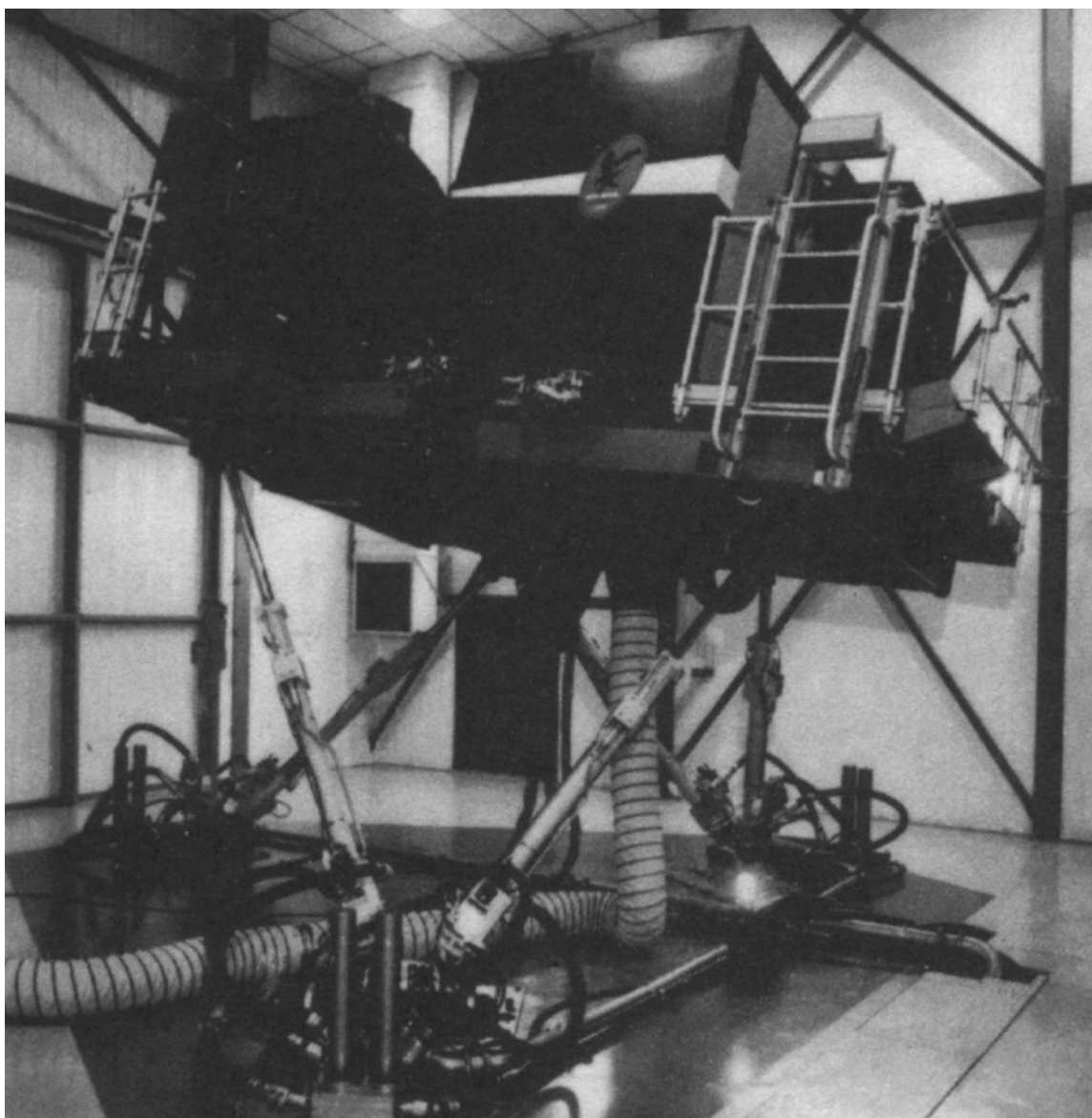
"Same as mine; Amundsen."

Roald Amundsen, I believe, found the magnetic North Pole.

Harald Fodstad

New York, New York

DOI: 10.1227/01.NEU.0000117129.32806.41



The Canadian CAE-Link Corporation developed the motion platform pictured, which simulates flying in a Black Hawk helicopter. The platform, from the early 1990s, rested on pneumatic legs and was run by an array of computers that could process 670,000 instructions per second. Also see pages 883 and 1041. (Courtesy, CAE-Link Corp.)