Endoscopic Versus Traditional Craniofacial Resection for Patients with Sinonasal Tumors Involving the Anterior Skull Base

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Objectives. With the advent of microdebriders and image guidance systems, endoscope-assisted surgery is now more widely used for the treatment of tumors involving the base of the skull. The aim of this study was to analyze the clinical features of tumors involving the anterior skull base and to evaluate the treatment outcomes according to the surgical approach, which included the traditional craniofacial resection (TCFR) and the endoscopic craniofacial resection with craniotomy (ECFR).

Methods. Forty-six patients who underwent craniofacial resection from 1989 through 2006 at Seoul National University Hospital and Seoul National University Bundang Hospital were included in this study. Demographics, histology, surgical management, surgical outcomes, complications, and morbidity were analyzed.

Results. The number of malignant and benign lesions was 40 and 6 cases respectively. The most common diagnosis was olfactory neuroblastoma occurring in 41% of the cases followed by squamous cell carcinoma and malignant melanoma. Thirty-six patients underwent TCFR, while ECFR was performed with or without adjunctive chemotherapy or radiotherapy in 10 patients. The overall five-year survival rate for patients with malignant tumors of the anterior skull base was 47.4%. Out of 19 patients with olfactory neuroblastomas, 10 patients had TCFR and six among them died of their disease. Nine patients underwent ECFR, and none of them died of their disease. The ECFR group had lower morbidity and cosmetic deformity than did the TCFR group.

Conclusion. The ECFR may be considered as an alternative option for the treatment of selected tumors with anterior skull base invasion. This approach offers the advantages of avoiding facial incisions with comparable treatment results.

Key Words. Nose neoplasms, Skull base, Endoscopic craniofacial resection

INTRODUCTION

Sinoasal malignancies involving the anterior skull base (ASB) are challenging due to the complex anatomy and important structures that may be involved (1). Many surgical techniques for approaching this area have been developed for complete resection of these tumors. Since Ketcham (2) first described the combined transcra-
nasal tumors. The clinical features were analyzed and the surgical complications and morbidity were compared according to the surgical technique used, which included the traditional craniofacial resection (TCFR) or the endoscopic craniofacial resection with craniotomy (ECFR) (11).

MATERIALS AND METHODS

A retrospective chart review was performed on 46 patients who were diagnosed with sinonasal tumors with involvement of the ASB that underwent craniofacial resection between January 1989 through December 2006 at the Seoul National University Hospital and Bundang Hospital. The patient records were reviewed for demographic data, clinical presentation, operative procedures, postoperative course, histopathological findings, immediate/ delayed complications, morbidity and mortality. Twenty-four patients were male and 22 patients were female. The average age of the patients was 43 yr, with a range of 5 to 70 yr. The mean follow-up period was 31 months, with a range of 2 to 202 months. The preoperative evaluation consisted of sinonasal endoscopy, CT/MR scans and screening for distant metastasis: bone/liver scans or positron emission tomography (PET). Patients who had resectable lesions with no systemic metastasis underwent surgery. Up until 1998, 18 cases of TCFR were performed; since 1999, in cases without involvement of the orbit, skin or facial bones, ECFR (10 cases) was performed; in patients with involvement, TCFR (18 cases) was performed. In cases with malignant tumors, ECFR was performed in 9 patients with ONB; for benign tumors, ECFR was performed in 1 case with a meningioma. Our Institutional Review Board approved the protocol for this retrospective analysis.

Endoscopic craniofacial resection with craniotomy

In all patients, CSF drainage was performed to minimize injury to the frontal lobes during the retraction. For large tumors where the origin could not be defined, the procedure began with margin clarification after tumor volume reduction with the use of a microdebrider. The incision had a minimum margin of 1 cm from the tumor attaching site. In cases where en bloc resection was feasible, intranasal structures were included in the extent of the resection after an accurate delineation of the tumor margin with the use of an endoscope. A safe margin around the tumor was ensured with a 1 cm incision. In the areas more apart from the tumor margin a frozen section biopsy was done around the region of the incision site. Then, subperiosteal dissection was performed to identify the nasal septum, skull base and orbit. In areas such as ethmoid sinus where the subperiosteal dissection was difficult, the microdebrider was first used to reduce the mass volume and then the mucosal dissection followed. For visualization, the septum was removed and the frontal or sphenoid sinus was opened. An osteotomy was created at the lesion and the tumor was delivered through the craniotomy site. A pericranial flap was used for reconstruction. No facial incision was made with this surgical technique in contrast to the TCFR.

Adjunctive treatment for ONB

Ten patients underwent TCFR, all were classified as Kadish type C. Among them, eight patients had postoperative radiotherapy, and chemotherapy was provided to one patient. In the nine patients that had ECFR, six (Kadish type A: 1, B: 1, C: 4 cases) underwent neoadjuvant chemotherapy and postoperative radiotherapy. Two patients (Kadish type C) had postoperative radiotherapy only. One patient with Kadish type C who showed partial remission after neoadjuvant chemotherapy, was reluctant to receive surgery, and had radiotherapy instead. However, there was no improvement after the radiotherapy, and finally surgery with chemotherapy was performed. The chemotherapy regimen consisted of etoposide (75 mg/m²/day, days 1-5), ifosfamide (1,000 mg/m²/day, days 1-5) and cisplatin (15 mg/m²/day, days 1-5). At first, two cycles of chemotherapy were administered and additional cycles were planned according to the response to the first two chemotherapy cycles.

The initial symptoms, tumor involvement and the histopathological findings were reviewed and the morbidity and complications were assessed according to the surgical technique. The Mann-Whitney U test and the Kaplan Meier method were used to identify the surgical outcomes and morbidity of the patients receiving the TCFR compared to those who had an ECFR. The data were analyzed using SPSS 12.0 for Windows (SPSS Inc., Chicago, IL, USA). P-values ≤ 0.05 were considered statistically significant.

RESULTS

Clinical features of the tumors

The patient presentation most commonly included a history of nasal obstruction and unilateral epistaxis. The average duration from symptom development to diagnosis was 4.3 months. The sites commonly involved included the nasal cavity, ethmoid sinus and cribriform plate. In addition, the sphenoid sinus, frontal lobe

### Table 1. Involved sites at diagnosis

| Involved sites                        | No. * |
|--------------------------------------|-------|
| Nasal cavity                         | 27    |
| Ethmoid sinus                        | 23    |
| Cribriform plate & frontal floor      | 23    |
| Orbit                                | 14    |
| Maxillary sinus                      | 11    |
| Sphenoid sinus                       | 11    |
| Frontal sinus                        | 5     |
| Frontal lobe                         | 4     |
| Others (Masticator space, Cavernous sinus) | 2     |

*The numbers are not mutually exclusive.
and cavernous sinus were also involved (Table 1).

A list of the histopathological diagnoses is given in Table 2 and 3. The patients with benign tumors were followed with no evidence of disease except one patient with a solitary fibrous tumor that underwent malignant transformation (Table 3). For the 40 patients with a malignant tumor, 18 were followed with no evidence of disease, and the survival outcomes depended on the histopathological type of tumor (Table 4).

Table 2. Histopathological types of malignant tumors

| Pathology               | TCFR | ECFR | Total |
|-------------------------|------|------|-------|
| Olfactory neuroblastoma | 10   | 9    | 19    |
| Squamous cell carcinoma | 7    | 7    | 14    |
| Malignant melanoma      | 3    | 3    | 6     |
| Rhabdomyosarcoma        | 2    | 2    | 4     |
| Chondrosarcoma          | 2    | 2    | 4     |
| Basal cell carcinoma    | 2    | 2    | 4     |
| Undifferentiated carcioma | 1  | 1    | 2     |
| Adenoid cystic carcioma | 1    | 1    | 2     |
| Malignant meningioma    | 1    | 1    | 2     |
| Undifferentiated sarcoma | 1  | 1    | 2     |
| Non-keratinizing carcioma | 1  | 1    | 2     |
| Total                   | 31   | 9    | 40    |

TCFR: traditional craniofacial resection; ECFR: endoscopic craniofacial resection with craniotomy.

Table 3. Summary of benign tumors

| Pathology                     | Sex/age | Involved sites | Treatment | F/U (mo) | State |
|-------------------------------|---------|----------------|-----------|----------|-------|
| Angiofibroma                  | M/16    | N, middle      | TCFR      | 5        | NED   |
|                               |         | cranial fossa  |           |          |       |
| Angiofibroma                  | M/16    | S              | TCFR      | 72       | NED   |
| Angiofibroma                  | M/13    | N,S            | TCFR      | 95       | NED   |
| Ossifying fibroma             | M/16    | E, S, cavernous| TCFR      | 202      | NED   |
|                               |         | sinus          |           |          |       |
| Solitary fibrous tumor        | M/38    | N, M, O, E, C | TCFR      | 29       | F/L   |
| Meningioma                    | F/62    | E, C           | ECFR      | 16       | NED   |

N: nasal cavity; S: sphenoid sinus; M: maxillary sinus; O: orbit; E: ethmoid sinus; C: cribiform plate & frontal floor; TCFR: traditional craniofacial resection; ECFR: endoscopic craniofacial resection with craniotomy; NED: no evidence of disease; F/L: follow-up loss.

Table 4. Follow-up of patients treated with craniofacial resection of malignant tumors

| Pathology                      | No. | NED | AWD | DOD | DOC | F/L |
|--------------------------------|-----|-----|-----|-----|-----|-----|
| Olfactory neuroblastoma        | 19  | 11  | 3   | 2   | 2   | 1   |
| Squamous cell carcinoma        | 7   | 3   | 4   |     |     |     |
| Malignant melanoma             | 3   | 3   |     | 3   |     |     |
| Rhabdomyosarcoma               | 2   | 1   | 1   |     |     |     |
| Chondrosarcoma                 | 2   | 1   |     |     |     |     |
| Basal cell carcinoma           | 2   | 1   | 1   |     |     |     |
| Undifferentiated carcinoma     | 1   |     |     |     |     | 1   |
| Adenoid cystic carcinoma       | 1   | 1   |     |     |     |     |
| Malignant meningioma           | 1   |     |     |     |     |     |
| Undifferentiated sarcoma       | 1   |     |     |     |     | 1   |
| Non-keratinizing carcinoma     | 1   |     |     |     |     |     |
| Total                          | 40  | 18  | 6   | 10  | 2   | 4   |

NED: no evidence of disease; AWD: alive with disease; DOD: died of disease; DOC: died from other causes; F/L: follow-up loss.

Treatment outcomes

Thirty-one patients out of 40 patients who were diagnosed with malignant tumors, underwent a TCFR, of which 17 patients (55%) developed disease recurrence; 11 had a local recurrence (35%), 4 had brain metastases (13%) and 2 had spinal metastases (6%). One patient (11%) out of 9 who underwent an ECFR had a local recurrence. For malignant tumors, the follow-up period ranged from 2 to 116 months, with a mean of 25.4 months. The overall 5-yr survival rate for the sinonasal tumors involving the anterior skull base was 47.4%.

Complications and morbidity

There were nine complications related to the surgery. Eight occurred after the TCFR, and 1 case of an epidural hematoma occurred after an ECFR. Complications related to the TCFR included CSF leakage, meningitis, hematoma, and flap necrosis at the facial incision site. Seven cases with complications were managed by surgery such as a craniotomy and wound revision; one patient expired due to complications and one patient had persistent facial paralysis. Another two patients with complications were treated with conservative antibiotic management.

A subset of the patients who underwent an ECFR was selected and compared to a group that underwent a TCFR. The groups
were compared with respect to operation time, intraoperative blood loss, length of hospital stay and length of ICU stay. The ECFR group had a shortened length of stay and operation time; these differences were statistically significant (P-value: 0.016 and 0.002 respectively). With respect to the length of the ICU stay and the operation time, the differences did not reach statistical significance (P-value: 0.679 and 0.059 respectively) (Fig. 1).

Postoperative course
In 19 patients with ONB, 10 patients underwent a TCFR, while nine patients underwent an ECFR. For the patients with a TCFR, the mean follow-up period was 28.8 ± 34.7 months and six of the 10 patients died of their disease (Table 5). For the patients who had an ECFR, the mean follow-up period was 20.7 ± 13.4 months and all nine patients are still alive (Table 6).

**DISCUSSION**

The incidence of sinonasal cancer has been reported to be one in 100,000 per year; it accounts for approximately 3% of all the upper aerodigestive tract malignancies and less than 1% of all cancers (12). In particular, the incidence of tumors involving the anterior skull base is too low to analyze the demographic data and survival outcome according to the histopathological type. Squamous cell carcinomas and olfactory neuroblastomas are known to be the two most common malignancies followed by primary salivary gland tumors and sarcomas; our findings confirm this (13). ONB is known to have a good prognosis. Although several adjunctive treatments have been developed, surgery is considered to be the standard treatment and postoperative radiotherapy is recommended to improve the local control rate. The standard treatment may vary according to the histopathological type of tumor. However in many cases, there is no standard treatment established to date.

Previously, craniofacial resections were associated with a high rate morbidity and mortality related to the surgical technique (2, 14, 15). However, with progress in surgical methods and reconstruction approaches such as the pericranial flap, the complication rate has decreased dramatically (16-18). Recently, noninvasive surgical techniques have created a great deal of interest. The endoscope assisted CFR has been developed and can be used...
clinically. Endoscopic surgery has been used in rhinosinusitis surgery since the 1980s. As the knowledge and experience with endoscopic surgery has improved, and the image guidance systems and surgical instruments for endoscopic surgery have evolved, endoscopic surgery has been increasingly used for the treatment of benign tumors (19). As experience with benign tumor surgery accumulated, endoscopic surgery was adopted for the treatment of malignant tumors (5-9, 20). Buchmann et al. (1) reported that endoscopic instruments and techniques not only allowed excellent visualization of the tumor, but also greatly aided the accurate microscopic resection, resulting in good patient outcomes and reduced surgery related morbidity. In 2006, Paolo et al. (8) reported that the major exclusion criteria for an ECFR were as follows: 1) tumors involving the lacrimal tract, 2) tumor infiltration of the hard palate, 3) tumors that involve the posterior wall of the sphenoid sinus, and 4) tumor invasion of all but the medial wall of the maxillary sinus. However, the indications for an ECFR have not yet been established. The development of imaging diagnostic tools as well as adjuvant treatments has expanded the role of the ECFR. The development of imaging diagnostic tools has enabled early detection and accurate preoperative evaluation. In addition, neoadjuvant/adjuvant chemotherapy and radiotherapy has aided in improving treatment results (1, 21).

In our study, for the malignancies, ONB was the only histopathological type where the ECFR was used. This is due to the limitations of the endoscopic approach and the advent of adjuvant therapy for ONB. The endoscopic approach is limited in lesions involving the orbit, skin and facial bones. Consequently, an ONB, which is commonly located at the interface between the superior nasal cavity and the anterior cranial fossa, is a better indication for an ECFR than squamous cell carcinoma, which commonly occurs in the maxillary sinus and is invasive to adjacent structures. The ONB originates from neuroendocrine cells similar to small cell lung cancer, and small cell lung cancer is known to respond to chemotherapy agents such as cisplatin. There is one report that confirms that ONB is responsive to cisplatin (22). Therefore, at our institution, a combined modality treatment has been adopted, in which initial cisplatin based neoadjuvant chemotherapy (VIP) is provided to reduce the tumor volume and surgery is planned to minimize the functional and cosmetic deformities. In addition, in patients with Kadish type C disease, postoperative radiotherapy is recommended (21, 23, 24). The development of adjuvant therapy has supplemented the limitations of endoscopic resection and allows for a more expanded use of endoscopy with other histopathological tumor types.

The results of this study showed that the patients who had an ECFR had a tendency to have reduced morbidity and a good survival outcome. However, in most ECFR cases, neoadjuvant chemotherapy and postoperative radiotherapy were performed, and the follow-up period was relatively short, in contrast to the TCFR cases, where neoadjuvant chemotherapy was not usually performed. Therefore, further long term follow-up studies are needed to evaluate the outcomes of the ECFR compared to the TCFR with respect to oncological safety and survival.

In conclusion, the ECFR may have the advantages of reducing the surgery related morbidity and mortality compared to the TCFR. Therefore, it should be considered as an alternative treatment option for selected sinonasal tumors involving anterior skull base.

REFERENCES

1. Buchmann L, Larsen C, Pollack A, Tawfik O, Sykes K, Hoover LA. Endoscopic techniques in resection of anterior skull base/paranasal sinus malignancies. Laryngoscope. 2006 Oct;116(10):1749-54.
2. Ketcham AS, Wilkins RH, Van Buren JM, Smith RR. A combined intracranial facial approach to the paranasal sinuses. Am J Surg. 1963 Nov;106:698-703.
3. Kraus DH, Shah JP, Arbit E, Galicich JH, Strong EW. Complications of craniofacial resection for tumors involving the anterior skull base. Head Neck. 1994 Jul-Aug;16(4):307-12.
4. Richtsmeier WJ, Briggs RJ, Koch WM, Eisele DW, Loury MC, Price JC, et al. Complications and early outcome of anterior craniofacial resection. Arch Otolaryngol Head Neck Surg. 1992 Sep;118(9):913-7.
5. Yuen AP, Fung CF, Hung KN. Endoscopic cranial resection of anterior skull base tumor. Am J Otolaryngol. 1997 Nov-Dec;18(6):431-3.
6. Thaler ER, Kotapka M, Lanza DC, Kennedy DW. Endoscopically assisted anterior cranial skull base resection of sinonasal tumors. Am J Rhinol. 1999 Jul-Aug;13(4):303-10.
7. Devaiah AK, Larsen C, Tawfik O, O’Boynick P, Hoover LA. Esthesioneuroblastoma: endoscopic nasal and anterior craniofacial resection. Laryngoscope. 2003 Dec;113(12):2086-90.
8. Castelnuovo PG, Beli E, Bignami M, Battaglia P, Sberze F, Torrei G. Endoscopic nasal and anterior craniofacial resection for malignant nasoethmoid tumors involving the anterior skull base. Skull Base. 2006 Feb;16(1):15-8.
9. Lund V, Howard DJ, Wei WI. Endoscopic resection of malignant tumors of the nose and sinuses. Am J Rhinol. 2007 Jan-Feb;21(1):89-94.
10. Lee CH, Jung HW, Rhee CS, Park HJ, Hah JH, Min YG, et al. Anterior and middle skull base surgery: SNUH Experience. Korean J Otolaryngol-Head Neck Surg. 1998 Feb;41(2):218-225.
11. Har-El G, Casiano RR. Endoscopic management of anterior skull base tumors. Otolaryngol Clin North Am. 2005 Feb;38(1):133-44.
12. Rhee CS, Won TB, Lee CH, Min YG, Sung MW, Kim KH, et al. Adenoid cystic carcinoma of the sinonasal tract: treatment results. Laryngoscope. 2006 Jun;116(6):982-6.
13. Ogusthorpe JD, Patel S. Craniofacial approaches to tumors of the anterior skull base. Otolaryngol Clin North Am. 2001 Dec;34(6):1123-42.
14. Cheesman AD, Lund VJ, Howard DJ. Craniofacial resection for tumors of the nasal cavity and paranasal sinuses. Head Neck Surg. 1986 Jul-Aug;8(6):429-35.
15. Shah JP, Sundaresan N, Galicich J, Strong EW. Craniofacial resections for tumors involving the base of the skull. Am J Surg. 1987 Oct;154(4):352-8.
16. Levine PA, Scher RL, Jane JA, Persing JA, Newman SA, Miller J, et al. The craniofacial resection-eleven-year experience at the University of Virginia: problems and solutions. Otolaryngol Head Neck Surg. 1989 Dec;101(6):665-9.
17. Lund VJ, Harrison DF. Craniofacial resection for tumors of the nasal cavity and paranasal sinuses. Am J Surg. 1988 Sep;156(3 Pt 1):187-90.
18. McCutcheon IE, Blacklock JB, Weber RS, DeMonte F, Moser RP, Byers M, et al. Anterior transcranial (craniofacial) resection of tumors of the paranasal sinuses: surgical technique and results. Neurosurgery. 1996 Mar;38(3):471-9.
19. Bockmühl U, Minovi A, Kratzsch B, Hendus J, Draf W. Endonasal micro-endoscopic tumor surgery: state of the art. Laryngorhinootologie. 2005 Dec;84(12):884-91.

20. Roh HJ, Batra PS, Citardi MJ, Lee J, Bolger WE, Lanza DC. Endoscopic resection of sinonasal malignancies: a preliminary report. Am J Rhinol. 2004 Jul-Aug;18(4):239-46.

21. Oskouian RJ Jr, Jane JA Sr, Dumont AS, Sheehan JM, Laurent JJ, Levine PA. Esthesioneuroblastoma: clinical presentation, radiological, and pathological features, treatment, review of the literature, and the University of Virginia experience. Neurosurg Focus. 2002 May 15;12(5):e4.

22. Kim DW, Jo YH, Kim JH, Wu HG, Rhee CS, Lee CH, et al. Neoadjuvant etoposide, ifosfamide, and cisplatin for the treatment of olfactory neuroblastoma. Cancer. 2004 Nov 15;101(10):2257-60.

23. Simon JH, Zhen W, McCulloch TM, Hoffman HT, Paulino AC, Mayr NA, et al. Esthesioneuroblastoma: the University of Iowa experience 1978-1998. Laryngoscope. 2001 Mar;111(3):488-93.

24. Polin RS, Sheehan JP, Chenelle AG, Munoz E, Larner J, Phillips CD, et al. The role of preoperative adjuvant treatment in the management of esthesioneuroblastoma: the University of Virginia experience. Neurosurgery. 1998 May;42(5):1029-37.