Proton Therapy for Skull Base Chordomas: An Outcome Study from the University of Florida Proton Therapy Institute

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Abstract

Objectives Skull base chordoma is a rare, locally aggressive tumor located adjacent to critical structures. Gross total resection is difficult to achieve, and proton therapy has the conformal advantage of delivering a high postoperative dose to the tumor bed. We present our experience using proton therapy to treat 33 patients with skull base chordomas.

Design Retrospective outcomes study.

Setting University of Florida Proton Therapy Institute; 2007 to 2011.

Participants A total of 33 patients with skull base chordomas received postoperative three-dimensional conformal proton therapy. The patients were 79% male and 6% diabetic; 27% had received a gross total resection.

Main Outcome Measures The gross tumor/tumor bed received a dose between 77.4 CGE and 79.4 CGE. Local control and overall survival were tracked, and radiation toxicity was assessed using a modified Radiation Therapy Oncology Group/European Organization for Research and Treatment of Cancer Late Radiation Morbidity Scoring Scheme.

Results Median follow-up for all patients was 21 months. Local control and overall survival rates at 2 years were 86% and 92%, respectively. Grade 2 toxicity was observed in 18% of our cohort in the form of unilateral hearing loss partially corrected with a hearing aid. No grade 2 or higher optic or brainstem toxicities were observed.

Conclusions Proton therapy is an effective treatment modality for skull base chordomas.

Keywords

- head and neck tumors
- chordoma
- outcomes
- proton therapy
- particle therapy

Introduction

Chordoma is a rare aggressive tumor that arises from the remnants of the notochord, which is the precursor to the nucleus pulposus or intervertebral disc.¹ According to Surveillance Epidemiology and End Results data, the annual incidence is ~0.08 for 100,000 people.² Chordomas, although rarely metastatic, typically occur in a location adjacent to critical structures. The anatomical distribution of chordomas has been found to be 32.8% spinal, 32% cranial, and 29.2%

sacral.^{3,4} Local progression of a base of skull chordoma can lead to severe neurologic deficits or death. The average time to death in an untreated skull base chordoma is between 6 and 24 months.⁵ Symptoms can be insidious, leading to a delay in diagnosis; Volpe et al showed that the median time from identification of symptoms to diagnosis was 14.6 months.⁶

The mainstay of treatment for a skull base chordoma is maximal tumor debulking. A gross total resection, however, is difficult to achieve, often leaving residual tumor. Local

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recurrence with surgery alone has been found to be at least 58% and is associated with a significant postsurgical toxicity and mortality.⁷ Adjuvant radiotherapy (RT) is critical to achieve tumor control.

Adjuvant photon RT has been used extensively in the past. Historically, conventional opposed photon fields treating to a median dose of 60 Gy yielded local control rates from 23 to 39% and survival rates from 35 to 51% at 5 years.^{8,9} Debus et al used more conformal photon techniques and achieved a higher median dose to 66.6 Gy, resulting in a 5-year local control rate of 50%.¹⁰ While suggesting a dose–response relationship, it appears that photon RT is limited by its conformality to achieve the necessary dose. Stereotactic radiosurgery (SRS) for skull base chordomas has shown improved or comparable local control for small tumors or recurrent disease. A study from Japan reported on 30 patients with skull base chordomas with a mean volume of 19.7 cm³ treated to a mean margin dose of 14 Gy. The 5-year overall local control rate of 76% correlated only with a tumor volume of greater or less than 20 cm³ on univariate and multivariate analyses.¹¹ Martin et al treated 18 chordoma patients with SRS to a mean marginal dose of 16.5 Gy and achieved a 5-year local control rate of 53%.¹²

Proton RT has an advantage over photon-based RT by using the Bragg peak to deliver a high conformal tumor dose to a target volume with very steep dose falloff, thus minimizing irradiation to adjacent critical structures. Physicians at the Massachusetts General Hospital (Boston, MA) have the most extensive experience using proton therapy for skull base chordomas.¹³ Their 5- and 10-year local control rates were 73% and 54%, respectively, using a tumor dose ranging from 66 to 83 CGE. We report our experience at the University of Florida Proton Therapy Institute (UFPTI) treating base of skull chordomas with protons.

Methods and Materials

A total of 33 patients with histologically confirmed skull base chordoma were enrolled from 2007 through 2011 in a prospective study approved by the institutional review board. Inclusion criteria were a primary site arising from the base of skull (sphenoid, clivus, petrous, or basiocciput), age 18 years or older at the time of consent, Karnofsky performance status ≥ 50, neurologic function allowing the patient to cooperate with treatment, and surgery ranging from biopsy to gross total resection before RT. ►Table 1 lists the patient and tumor characteristics for our study population.

Exclusion criteria included the following: prior RT to the head and neck, evidence of metastatic disease, serious medical or psychiatric illness that would preclude treatment compliance or informed consent, and pregnant or breastfeeding women. Only 27% of our patients had a gross total resection of the tumor.

For the simulation and treatment setup, each patient's head was supported on a carbon fiber Base of Skull (BoS) frame (Qfix, Avondale, PA) and immobilized with a customized thermal mask. The BoS frame is fabricated to ensure uniform attenuation of the proton beam over its entire

Table 1 Tumor and treatment characteristics

Characteristic	Number (%) of patients
Presenting cranial nerve deficit	
VI	11 (33)
XII	4 (12)
Brainstem involvement	
Yes	11 (33)
No	22 (67)
Optic pathway involvement	
Yes	3 (9)
No	30 (91)
Resection	
GTR	9 (27)
STR	22 (67)
Biopsy Only	2 (6)

Abbreviations: GTR, gross total resection; STR, subtotal resection.

surface area. The design also allows maximum flexibility in field arrangements in conjunction with a robotic couch and a 360-degree proton gantry. Patient setup was conducted with care to ensure shoulder clearance and minimize the air gaps (and the lateral penumbra) of the treatment fields. Computed tomography (CT) images with and without intravenous contrast were acquired at 1-mm spacing from the vertex to the base of skull. The CT image sets were registered with magnetic resonance imaging (MRI) data sets to guide the delineation of the target volumes and critical structures.

►Fig. 1 shows a patient with a clivus chordoma treated with proton therapy. The prescriptions were 50.4 CGE to the initial planning target volume (PTV 50.4) and 73.8 CGE to the boost volume (PTV 73.8). Because of the tendency of the target volume to wrap around the brainstem, and the target's close proximity to the visual apparatus, a patch-field technique unique to proton treatment is often used as part of the field arrangement. In the example shown, the initial target volume, PTV 50.4, was treated with five fields including a pair of patch fields. In the first pair, the left anterior oblique through field (LAO-T) was intentionally blocked to avoid the brainstem, missing the posterior aspect of the target as a result. The residual target was treated, or “patched,” with a left posterior oblique patch field (LPO-P), which was feasible because of the finite range characteristics of the proton beam. The 50% dose level of the distal falloff of the LPO-P was adjusted to match the 50% dose corresponding to the field edge of the LAO-T beam. Ideally, the dose gradient of the distal falloff should closely match that of the lateral penumbra to achieve a homogeneous dose at the junction and within the target volume. In reality, there could be hot and cold spots near the junction. The second pair, composed of the right anterior oblique (RAO-T) through beam and the right posterior oblique patch (RPO-P) beam, somewhat mirrored the first so as to minimize the uncertainties and the impact of the

Fig. 1 A patient with a clivus chordoma. Five treatment fields were used for the planning target volume (PTV) 50.4 CGE (delivered dose: 54 CGE). (A) Axial view showing the two pairs of patch fields: First pair is left anterior oblique-through (LAO-T) and left posterior oblique-patch (LPO-P); second pair is right anterior oblique-through (RAO-T) and right posterior oblique-patch (RPO-P). The through beams (LAO-T and RAO-T) are laterally blocked to avoid the brainstem. The residual target volumes are covered by the patch fields (LPO-P and RPO-P) coming from the posterior. (B) Sagittal view showing the 5th superior anterior oblique (SAO) beam. This beam angle allows the brainstem to be blocked while essentially covering the whole target. Also shown is the boost target volume, PTV 73.8 CGE.

hot and cold spots that could arise at the junctions. Note that both the RAO and LAO fields were selected so that the chiasm and optic nerves could be blocked. To further improve the target coverage, a superior anterior oblique (SAO) field was added with an angle that allowed the brainstem to be shielded (►Fig. 1B). The boost volume was treated with a similar field arrangement.

►Fig. 2 shows the isodose in color wash and the dose-volume histogram of the composite plan. The dose plan was designed with high priority to limit the dose to the critical structures. The 0.1 cm³ doses to the brainstem, spinal cord, chiasm, and optic nerves as well as the expanded volume

(5 mm for spinal cord and 3 mm for brainstem, chiasm, and optic nerves) are tabulated in ►Table 2. The dose levels achieved were well under the established dose tolerances for these structures. Despite our conservative approach, adequate doses for the target volumes were also achieved. Nearly 100% of the initial target volume received the treatment dose of 50.4 CGE; 90% of the boost volume received the treatment dose of 73.8 CGE. To minimize the risk of necrosis, the hot spot within the target (2% volume) was limited to 114% of the prescription dose. Daily treatments were delivered with a digital online image guidance system, and a setup accuracy >2 mm was achieved.

Fig. 2 (A–C). Isodose shown in color wash for the composite plan. The prescription dose to the planning target volume (PTV) was 50.4 CGE for the initial target volume and 73.8 CGE for the boost volume. (D) Dose-volume histogram distribution for the composite plan. Note: BS + 3 = brainstem plus 3-mm expansion.

Table 2 Dose constraint results

Constraint	Number (%) of patients who met the goal
PTV D95 = 100%	29 (88)
Brainstem (center) 0.1 cm ³ <55 CGE	29 (88)
Cord 0.1 cm ³ <50 CGE	33 (100)
Chiasm 0.1 cm ³ <55 CGE	31 (94)
Right optic nerve 0.1 cm ³ <55 CGE	32 (97)
Left optic nerve 0.1 cm ³ <55 CGE	33 (100)

Abbreviation: PTV, planning target volume.

Patients received a median PTV dose of 74 CGE (range: 70–79 CGE). Patients were followed for a median of 21 months (range: 3–58 months). Skull base MRs were obtained every 6 months.

Toxicities were scored using a modified Radiation Therapy Oncology Group/European Organization for the Research and Treatment of Cancer Late Radiation Morbidity Scoring Scheme.

All statistical computations were accomplished with SAS and JMP software (SAS Institute, Cary, NC). The Kaplan-Meier product limit provided estimates of local control and overall survival.¹⁴

Results

With regard to surgical toxicities, 10 patients experienced a cerebrospinal fluid leak, 4 patients had postoperative meningitis, and 4 patients experienced a new cranial nerve deficit after resection.

The local control and overall survival rates at 2 years were 86% and 92%, respectively (►Fig. 3). One patient progressed through treatment and was put on salvage immunomodulator therapy. Three patients (9%) had an in-field recurrence after treatment. No patient experienced a marginal recur-

Fig. 3 Kaplan-Meier curves for overall survival and freedom from recurrence rates at 2 years.

rence or distant metastases. One patient died from disease progression.

The only radiation-related grade 2 or higher toxicity was unilateral hearing loss partially corrected by a hearing aid, which was observed in 18% of our patients. Endocrine toxicity data were largely unavailable to us because many of our patients were located in different states or internationally. There were no grade 2 or higher brainstem or visual toxicities.

Discussion

Skull base chordomas are locally aggressive and infiltrative tumors surrounded by critical structures and often cause debilitation or death.¹⁵ Surgery is the primary treatment, but the high likelihood of residual tumor requires optimal adjuvant RT. Our outcomes when treating this disease with adjuvant proton therapy are similar to those reported by investigations at other proton therapy institutions.

Maximal debulking is critical to control of a skull base chordoma, but, as seen in our experience, gross total resection is difficult to achieve.¹⁶ At the University of Arkansas, gross total resection was achieved in 10 of 25 patients (40%).¹⁷ Gay et al reported a 47% rate of near-total resection among 60 patients with skull base chordomas and chondrosarcomas.¹⁸ Although most of our patients underwent multiple operations, our gross total resection rate was 27%. Thus adjuvant proton therapy was predominantly delivered in the setting of gross residual disease. Furthermore, 30% of our patients experienced a cerebrospinal fluid leak, and 12% had postoperative meningitis following surgery. These surgical toxicities are similar to other reported series.^{17,18}

As with other studies,^{13,19,20} our volume definition was based on preoperative and postoperative MRI and CT images. Our normal-tissue constraints were also similar, as shown in ►Table 2. We saw only in-field recurrences and achieved a local control rate of 86% at 2 years, which makes our findings comparable with the published data. Hug et al reported a series of 33 patients with chordoma treated at Loma Linda University to a median dose of 70.7 CGE with a 5-year local control of 59%.¹⁹ A series from Massachusetts General Hospital in which 621 patients were treated with a mixture of photons and protons to between 66 and 83 CGE showed 5- and 10-year local control rates of 73% and 54%, respectively.¹³ Ares et al reported on their experience using the spot-scanning technique of proton delivery to treat 42 patients with skull base chordomas and observed a 5-year local control rate of 81% with this highly conformal technique.²⁰

Because the main site of recurrence is local and the salvage of local recurrences is unlikely, particularly in previously resected patients, survival is directly correlated with local control. Other institutions have reported 5-year survival rates of 59%, 73%, and 81%.^{13,19,20}

At 2 years, our patients have not experienced any brainstem or optic toxicities, and other series have shown low toxicity rates as well. Debus et al reported a 5.5% rate of brainstem symptoms in 19 of 348 evaluable patients. The median time to symptoms was 17 months. On multivariate analysis, there was a statistically significant increase in the

rate of brainstem symptoms when the volume of brainstem receiving 60 CGE was $>0.9 \text{ cm}^3$.^{21,22} Of our patients, 88% of patients received $<55 \text{ CGE}$ to 0.1 cm^3 of the brainstem. Pai et al reported on endocrine toxicity for 107 patients with skull base chordomas and chondrosarcomas. The 5- and 10-year rates for hyperprolactinemia were 72% and 84% and for hypothyroidism were 30% and 63%.³ These endocrinopathies were observed when the minimum dose to the pituitary exceeded 50 CGE $D_{\min} >50$. At present, we do not have the endocrine laboratory results for our patients.

Conclusions

Our experience treating patients with skull base chordomas with proton therapy is promising in terms of control of this locally aggressive tumor. In view of the high dose delivered and the proximity of adjacent critical structures, the toxicity profile is acceptable. With longer follow-up, we expect our 5- and 10-year local control and survival rates to approach those reported by other investigators.

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