Subacute Facial Nerve Paralysis Following Cyberknife Treatment for Vestibular Schwannoma

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Abstract
Facial nerve paralysis may occur along the natural progression of an untreated vestibular schwannoma, as a surgical complication following vestibular schwannoma resection, or as a late sequela of stereotactic radiation. However, facial paralysis occurring as a subacute complication of CyberKnife (Accuray, Inc., Sunnyvale, CA, USA) stereotactic radiation has not previously been reported. A 47-year-old male presented with a right vestibular schwannoma measuring 3.1 cm in greatest dimension with House-Brackmann grade I/VI facial function and American Academy of Otolaryngology – Head and Neck Surgery class a hearing. After deferring surgery, he elected to undergo CyberKnife stereotactic radiation to a total dose of 18 Gy in three stages. Within three months, the patient noticed gradual ipsilateral facial paralysis that progressed to House-Brackmann grade VI/VI over the course of four weeks. The patient's facial function was assessed continuously for one year with no volitional function or resting tone. He thus underwent a hypoglossal-facial transfer, restoring facial function to House-Brackmann IV/VI, with durable results at a follow-up length of five years. To our knowledge, this is the first reported case of subacute facial paralysis following CyberKnife stereotactic radiation for vestibular schwannoma.

Keywords: Facial Nerve Paralysis; Cyberknife Treatment; Vestibular Schwannoma

List of abbreviations: MRI: Magnetic Resonance Imaging; IAC: Internal Auditory Canal; WRS: Word Recognition Score

Introduction
Facial nerve paralysis may occur in the setting of vestibular schwannoma as a sequela of the natural history of an untreated tumor, as a complication of surgery, or as a late complication of stereotactic radiation. Estimates of facial nerve preservation following surgical resection, corresponding to grades I or II on the House-Brackmann scale [1], have been estimated at roughly 60% immediately postoperatively to 80-90% at long-term follow-up [2,3]. On the other hand, stereotactic radiation has been associated with an overall facial nerve preservation rate of 96.2% at long-term follow-up [4]. In particular, CyberKnife (Accuray, Inc., Sunnyvale, CA, USA) has been associated with excellent facial nerve preservation [5], with one series noting 100% preservation of facial function in 60 patients [6].

While facial nerve neuropathy in the setting of long-term outcomes following CyberKnife radiosurgery has been documented [7], facial nerve paralysis has not previously been reported as a subacute complication of stereotactic radiation for vestibular schwannoma. Here, we report a case of complete facial paralysis occurring within 3 months following CyberKnife stereotactic radiation for vestibular schwannoma.

Case Presentation
Informed verbal consent was obtained by the patient reported in this study.

A 47-year-old male presented to the neurology clinic with a chief complaint of episodic vertigo. The patient had no hearing complaints, and facial nerve function was noted to be I/VI. Magnetic resonance imaging (MRI) of the internal auditory canals (IAC) was performed, revealing a right-sided vestibular schwannoma measuring 3.1 by 2.7 by 2.4 cm, with extension into the IAC (Figure 1A). Subsequently, the patient was referred to the neurotologic clinic, where an audiogram revealed a right-sided mild high-
frequency hearing loss with a word recognition score (WRS) of 100%, corresponding to American Academy of Otolaryngology – Head and Neck Surgery hearing class A [8].

Following a discussion of management options, including surgery, stereotactic radiation, and observation, the patient deferred surgery and elected to proceed with stereotactic radiation. He was referred to the neurosurgical clinic for CyberKnife evaluation, and was deemed an acceptable candidate.

Over the course of three days, the patient underwent CyberKnife stereotactic radiation to his right vestibular schwannoma with a dose of 18 Gy delivered to the 78% isodose line using a 15mm collimator. The maximal dose within the center of the treatment volume was 23.1 Gy, and the volume treated was 10.4 cm³ (Figure 2). Dexamethasone 8 mg was administered orally after each of the three treatment sessions.

Serial imaging performed thereafter revealed a peak tumor size of 3.8 cm in greatest dimension. The patient developed transient papilledema and radiological evidence of hydrocephalus, which were managed medically. Eight months following stereotactic radiation, papilledema and hydrocephalus had both resolved, and oral dexamethasone was discontinued after a taper. However, facial function remained at House-Brackmann grade VI.
One year following the onset of facial paralysis, the patient presented again to the neurotologic clinic. He endorsed no serviceable hearing on the affected side, and audiological testing confirmed a WRS of 0% accompanying a mild sloping to profound sensorineural loss. Examination confirmed persistent House-Brackmann grade VI paralysis, and the patient elected to undergo hypoglossal-facial anastomosis for dynamic reanimation of his paralyzed face. By 18 months following surgery, the patient had restoration of House-Brackmann grade IV facial function, which remained stable at the last follow-up of 5 years postoperatively. Repeat MRI performed approximately two and a half years following stereotactic radiation revealed no progression in tumor size (Figure 1D).

Discussion

Facial nerve paralysis is recognized as a potential complication of surgical management or a sequela of stereotactic radiation that carries significant implications for quality of life following treatment [9]. Rates of anatomic preservation of the facial nerve following microsurgery range from 93-97% in large series [2,10]. Normal or near-normal facial function following surgery, defined as grade I or II on the House-Brackmann scale, is achieved in close to 90% of patients [2].

Delayed facial palsy, defined as deterioration of facial nerve function more than three days following treatment, has been reported as an uncommon sequela of microsurgery for vestibular schwannoma. Rates of delayed facial palsy range from 5 to 26%, but the prognosis is excellent, with a vast majority of patients noting full recovery to immediate postoperative function within one year [11,12]. Potential etiologies for delayed facial palsy include reactivation of herpes simplex virus and varicella zoster virus [13], and because of this, routine antiviral therapy following vestibular schwannoma resection has been advocated [14].

Facial nerve dysfunction following stereotactic radiation is relatively uncommon, with reported rates ranging from 0% to 7% [6,15,16]. Facial nerve dysfunction in these cases is hypothesized to result from compromised microvasculature in addition to progressive inflammation and fibrosis, a theory corroborated by intraoperative findings reported at the time of surgical salvage [17]. Overall, however, facial nerve paralysis following stereotactic radiation, including the CyberKnife system, is uncommon if facial nerve function is intact prior to treatment [4,7,18-20].

In the present case, a patient with a 3.1 cm vestibular schwannoma presenting with House-Brackmann grade I facial function experienced progressive, ultimately complete facial paralysis beginning fewer than three months following stereotactic radiation with CyberKnife. To our knowledge, this is the first such report in the literature. After observation for one year, the patient was noted to have no facial function, remaining at House-Brackmann grade VI. Function was restored to House-Brackmann grade IV following a hypoglossal-facial anastomosis.

Various authors have suggested that cranial nerve dysfunction, including facial paralysis, following stereotactic radiosurgery correlates strongly with minimum radiation dose as well as total radiation dose [21]. Hypofractionation has additionally been associated with a lower incidence of cranial nerve dysfunction following stereotactic radiation [22]. Although hypofractionation was indeed employed in the present case, a significant factor predisposing to a poorer outcome following stereotactic radiation was the size of the patient’s tumor (3.1 cm).

Conclusion

To our knowledge, this is the first report of subacute-onset facial nerve paralysis following CyberKnife stereotactic radiation. Facial paralysis, whether subacute or delayed, remains a known complication of stereotactic radiation for vestibular schwannoma. The possibility of such—as embodied by the present case—carries implications for patient counseling when determining the optimal management of a vestibular schwannoma.

References

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