Recurrent and residual juvenile angiofibromas

I Tyagi, R Syal, A Goyal

Abstract
Introduction: In the surgical management of juvenile nasopharyngeal angiofibromas the possibility of recurrences and residual tumours is always there. This study was undertaken to predict the prognostic factors determining recurrence of juvenile nasopharyngeal angiofibroma and to find out the usual sites of these tumours.

Material and methods: The medical records of 95 patients with histologically proven juvenile nasopharyngeal angiofibroma were reviewed retrospectively. The commonest surgical approach used was a combined transpalatal and transmaxillary approach with a lazy S incision. A conservative lateral infratemporal approach was used in three cases.

Results: Complete removal of the juvenile nasopharyngeal angiofibroma was achieved in 78 (82 per cent) of the cases in a single operation. A residual tumour was found in 17 (18 per cent) cases and recurrences occurred in 13 (13.7 per cent) cases.

Conclusions: Extensions into the pterygoid fossa and basisphenoid, erosion of the clivus, intracranial extensions medial to the cavernous sinus, invasion of the sphenoid diploe through a widened pterygoid canal, feeders from the internal carotid artery, a young age and a residual tumour were risk factors found associated with recurrence of juvenile nasopharyngeal angiofibroma.

Key words: Angiofibroma, Child; Nasopharynx, Recurrence; Otorhinolaryngological Surgical Procedures

Introduction
Juvenile nasopharyngeal angiofibroma is a histologically benign yet locally aggressive vascular tumour. It is considered to be the most common benign neoplasm of the nasopharynx accounting for 0.05 per cent of head and neck tumours. The highest incidence is found in India and Egypt. It commonly affects young pubescent males. Juvenile nasopharyngeal angiofibroma usually originates from the region of the sphenopalatine foramen just posterior to the middle turbinate. Although benign, the tumour causes pressure erosion of the bones and exhibits submucosal spread. Surgery has been the overwhelming choice for primary treatment of juvenile nasopharyngeal angiofibroma and the selection of proper approaches depends primarily upon the extensions of the tumour. The surgical management of juvenile nasopharyngeal angiofibroma is complicated by tumour vascularity, pattern of vascular supply, young age of patients, difficult surgical access, and complex loco regional anatomy. As a result the possibility of incomplete removal of the angiofibroma is always there. Recurrence is the other conspicuous feature of the natural history of juvenile nasopharyngeal angiofibroma. The situation was summed up accurately by Lafargue that some angiofibromas do not recur while others operated on by the same method recur with astonishing rapidity. This study was undertaken to help predict the prognostic factors determining recurrence of juvenile nasopharyngeal angiofibroma and to find out the usual sites of these tumours.

Material and methods
The medical records of patients with histologically proven large juvenile angiofibroma who received treatment in our institution between 1992 and 2002 were reviewed retrospectively. Fisch’s staging of juvenile nasopharyngeal angiofibroma, revised by Andrews et al. in 1989, with a few modifications (large extradural extensions of juvenile nasopharyngeal angiofibroma were included in stage IVa) was used for classification in this study (Table I). As our institute is a referral centre the majority of cases operated on were in stage III and IV as shown in Table II. Stage I juvenile nasopharyngeal angiofibroma were not included in this study. A battery of investigations like contrast enhanced computed tomography (CT) scan, magnetic resonance imaging (MRI) (in cases with intracranial extensions) and angiography (selected cases only) were reviewed retrospectively in patients in...
the study group. Pre-operative embolisation of feeding vessels and the tumour bed by polyvinyl alcohol particles was performed in 25 cases.

A transpalatal surgical approach was used in 15 cases of stage II juvenile nasopharyngeal angiofibroma. Stage IIIa juvenile nasopharyngeal angiofibroma having infratemporal, pterygopalatine and orbital extensions in 62 cases were removed in this case series by combining a transpalatal and transmaxillary approach with a lazy S incision, only in three cases of stage IIIa where the angiofibroma was extending up to the temporal fossa, was a conservative lateral infratemporal approach (type D1 approach) used. Small intracranial extensions of juvenile nasopharyngeal angiofibroma, that is stage IIIb tumours, were removed along with extra cranial part in 10 cases. Defects in the bone through which the intracranial extensions had occurred were widened. Just by performing a blunt dissection and applying gentle traction, intracranial extensions were separated from the dura. Intracranial pressure was kept low by giving mannitol during this procedure. Even stage IVa juvenile nasopharyngeal angiofibromas were removed by this approach in two cases. Staged frontotemporal craniotomy was performed to remove residual tumours in five cases with stage IVa tumours. A second operation was performed to remove residual tumour in 14 (82 per cent) cases in spite of complete removal. In four cases, three to four operations were performed to remove recurrences. Incidence of recurrent and residual tumours by stage is shown in Table III.

Discussion

In this case series the incidence of residual angiofibromas was 18 per cent while recurrences occurred in 13.7 per cent of cases. These results are good when compared to other studies as shown in Table IV. Risk factors found associated with recurrence of juvenile nasopharyngeal angiofibromas were extensions into the pterygoid fossa and basisphenoid, invasion of the sphenoid diploic through a widened pterygoid canal (Figure 1a), intracranial extensions medial to the cavernous sinus (Figure 1b), erosion of the clivus (Figure 1c), feeders from the internal carotid artery (Figure 1d), a young age and an evident residual tumour. Pre-operative embolisation was not found to be associated with an increased incidence of residual and recurrent tumours. The majority of recurrences occurred within six months of the first operation.

Residual tumours

In this case series, in 17 cases there was evident residual tumour on the first post-operative CT scan done after six weeks of surgery. Residual tumours were found in the region of the temporal and infratemporal fossa in four cases (23.5 per cent), in the

<table>
<thead>
<tr>
<th>Stage</th>
<th>Characteristics</th>
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<tbody>
<tr>
<td>I</td>
<td>Tumour limited to the nasal cavity and nasopharynx</td>
</tr>
<tr>
<td>II</td>
<td>Extensions into pterygopalatine fossa, maxillary, sphenoid, ethmoid sinuses</td>
</tr>
<tr>
<td>IIIa</td>
<td>Extensions into orbit or infratemporal fossa without intracranial extensions</td>
</tr>
<tr>
<td>IIIb</td>
<td>Stage IIIa with small extradural extensions of intracranial (parasellar) involvement</td>
</tr>
<tr>
<td>IVa</td>
<td>Large extradural intracranial or intradural extensions</td>
</tr>
<tr>
<td>IVb</td>
<td>Involvement of the cavernous sinus, pituitary, or optic chiasm</td>
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</table>

<table>
<thead>
<tr>
<th>Stage</th>
<th>Number of cases</th>
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<tbody>
<tr>
<td>I</td>
<td>Not included</td>
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<tr>
<td>II</td>
<td>10</td>
</tr>
<tr>
<td>IIIa</td>
<td>65</td>
</tr>
<tr>
<td>IIIb</td>
<td>10</td>
</tr>
<tr>
<td>IVa</td>
<td>7</td>
</tr>
<tr>
<td>IVb</td>
<td>3</td>
</tr>
<tr>
<td>Total</td>
<td>95</td>
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<table>
<thead>
<tr>
<th>Stage of JNA</th>
<th>No. of cases operated</th>
<th>Residual tumour n (%)</th>
<th>Recurrence n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>II</td>
<td>10</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>IIIa</td>
<td>65</td>
<td>9 (13.8)</td>
<td>8 (12)</td>
</tr>
<tr>
<td>IIIb</td>
<td>10</td>
<td>0</td>
<td>2 (20)</td>
</tr>
<tr>
<td>IVa</td>
<td>7</td>
<td>5 (71.4)</td>
<td>3 (43)</td>
</tr>
<tr>
<td>IVb</td>
<td>3</td>
<td>3 (100)</td>
<td>0</td>
</tr>
</tbody>
</table>

JNA = juvenile nasopharyngeal angiofibroma
region of the sphenoid sinus, the base of the pterygoids, the clivus and the vidian canal in five cases (29.4 per cent), the medial aspect of the cavernous sinus in three cases (17.6 per cent) and large residual tumours were present in the remaining five cases (29.4 per cent). Since the majority of cases in this case series were operated on by a combined transpalatal and transmaxillary approach, the infratemporal fossa and temporal regions were associated with an increased incidence of residual tumours. Jones et al.\textsuperscript{10} (1986) also observed that anterior approaches were associated with an increased incidence of residual tumours in the temporal and infratemporal region but tumour regrowth was not observed from these regions. Out of 13 cases in which recurrences occurred, in seven cases (54 per cent) there was evidence of residual tumours. In cases in which recurrences occurred, residual tumours were found in the region of the pterygoid fossa, sphenoid sinus, clivus, vidian canal, medial aspect of the cavernous sinus and the foramen ovale. Small evident residual tumours in these areas had regrown substantially by the time a second operation was performed in these seven cases. A post-operative CT scan should be performed after six weeks as an early scan may give a false impression of a residual tumour due to the pooling of secretions.

**Extensions into the pterygoid fossa**

Radkowski et al.\textsuperscript{12} observed that extensions of juvenile nasopharyngeal angiofibroma posterior to the pterygoid plates into the pterygoid muscles, a muscular bed similar to the infratemporal fossa, makes complete surgical excision difficult and increased risk of skull base erosion makes them prone to recurrences. Although the majority of angiofibroma cases usually extend laterally through the widened sphenopalatine foramen, sometimes angiofibroma may extend posterior to the pterygoid plates along the base of the skull involving the pterygoid fossa and may then extend laterally into the infratemporal fossa (retromaxillary spread) (Figure 2). This occurred in two cases in this present case series and both presented with residual and recurrent tumours (Figure 3). Juvenile nasopharyngeal angiofibroma extensions in the pterygoid fossa, a potential site of recurrent and residual tumours,\textsuperscript{12} were taken care of in this study by extending the limit of the transpalatal incision on the involved side into the retromaxillary region and if required, the medial pterygoid plate was removed to expose the extensions of the angiofibroma in the pterygoid fossa (Figures 4 and 5). The angiofibroma was then separated from surrounding soft tissues under direct vision. Since no major muscle is attached to the medial pterygoid plate so removal of the medial pterygoid plate produces less morbidity as compared to the lateral pterygoid plate, which is usually removed in lateral infratemporal approaches. Liu et al.\textsuperscript{15} also found a transpalatal approach combined with a labiogingival groove incision most appropriate to remove angiofibromas from the pterygoid fossa. Recurrences in their series were 16.7 per cent. Duvall et al.\textsuperscript{13} also recommended a combination of transantral, sublabial and transpalatal approaches, using a continuous palatobuccal (lazy S) incision around the maxillary tuberosity into the gingivobuccal sulcus to give a wide exposure to the pterygoid plates and the pterygomaxillary fossa.

**Intracranial extensions medial to the cavernous sinus**

Stage III and IV juvenile nasopharyngeal angiofibroma cases with involvement of the sphenoid sinus and extensions through the lateral wall of the sphenoid which were medial to the cavernous sinus were also prone to recurrent and residual tumours (23 per cent of cases). McGahan et al.\textsuperscript{16} advocated radiotherapy for extensions of juvenile nasopharyngeal angiofibroma which spread medial to the cavernous sinus and labelled them unresectable. Paris et al.\textsuperscript{17} (2001) used a transantral and lateral rhinotomy approach to remove such extensions and found it difficult to access the sphenoid sinus. Zhang et al.\textsuperscript{18} used lateral infratemporal approaches in stage III and IV juvenile nasopharyngeal angiofibroma and found that extensions of angiofibroma medial to the cavernous sinus were inaccessible by lateral approaches. Danesi et al.\textsuperscript{6} (2000) observed that tumours extending intracranially through an orbital fissure were lateral to the cavernous sinus and were accessible through either a lateral skull base or anterior approach like a transantral incision whereas tumours extending through the lateral wall of the sphenoid which were medial to the cavernous sinus were more readily removed through anterior approaches like a transpalatal approach with minimum morbidity. The sphenoid sinus and intracranial extensions of angiofibroma medial to the cavernous sinus were more readily removed through anterior approaches like a transpalatal one with minimum morbidity.
Lateral spread of angiofibroma into the infratemporal fossa (a) through the widened sphenopalatine fossa (b) retromaxillary spread.

Fig. 1
(a) CT scan showing dilatation of the vidian canal, (b) a residual tumour in the region of the sphenoid sinus and medial to the cavernous sinus; the patient had a recurrence, (c) MRI showing erosion of the clivus by angiofibroma, marked with arrow, (d) feeders from the internal carotid artery as is clear from the tumour blush when contrast was injected through the internal carotid artery.

Fig. 2
Lateral spread of angiofibroma into the infratemporal fossa (a) through the widened sphenopalatine fossa (b) retromaxillary spread.
sinus were easily accessible with a combined transpalatal and transmaxillary approach used in this case series. In spite of better exposure, in 23 per cent of cases of recurrence there was evident residual tumour in the sphenoid sinus eroding the lateral wall of the sphenoid and extending medial to the cavernous sinus. To remove the intracranial extensions of an angiofibroma through the lateral wall of the sphenoid that was medial to the cavernous sinus, tumours were displaced inferiorly from the roof of the sphenoid sinus. The plane between the tumour and the dura or thin bony remnant were identified. The bony defect in the lateral wall of the sphenoid was widened. The tumour was then gently separated from the dura. Usually there was no infiltration of neural or vascular components. Venous bleeding usually encountered from the cavernous sinus dura was controlled with surgicel kept in place for a few minutes by cottonoid. We are now drilling the base of the sphenoid sinus to expose the base of the greater wing of the sphenoid followed by drilling in the area of the vidian canal which we feel can significantly decrease the incidence of recurrent and residual tumours.

Age and feeders from the internal carotid artery
A young age (95 per cent of recurrences occurred in patients below 18 years of age) and feeders from the internal carotid artery were other factors found to be associated with the aggressive behaviour of an angiofibroma and a higher incidence of recurrences. Definite feeders from the internal carotid artery or tumour blush after injecting contrast in the internal carotid artery were found in 10 cases. Five such cases had a recurrence of the angiofibroma (50 per cent). No correlation between the incidence of recurrence of an angiofibroma and pre-operative embolisation was found.

Invasion of sphenoid diploe
Lloyd et al. found two types of involvement of the sphenoid: simple pressure erosion (40 per cent) and deep invasion and expansion of the sphenoid (60 per cent). Ninety-three per cent of the recurrences occurred in the latter group. In this present case series, on retrospective analysis imaging the evidence of invasion of the sphenoid diploe through a widened pterygoid canal (Figure 6 a and b) was found in...
84.6 per cent of cases (11 out of 13) of recurrences. Howard et al.\textsuperscript{20} also found that 93 per cent of recurrences occurred in patients with imaging evidence of invasion of the sphenoid diploe through a widened pterygoid canal. They also found that the mean number of recurrences was proportional to the degree of invasion of the greater wing of the sphenoid. Similarly, in the present case series the incidence of recurrence in stage IVa tumours was 43 per cent compared to 20 per cent in stage IIIb (Table III).

**Surgical approach**

In the present case series no correlation between surgical approaches and recurrences of angiofibroma was found. The majority of cases (96.9 per cent) in the present series were operated on by a combined transpalatal and transmaxillary approach by lazy S incision. In only 3.1 per cent (three cases) of cases in the present case series was a conservative lateral infratemporal approach (type D1 approach) used. Out of these three cases, in one case there was a large residual intracranial extension which was removed by staged frontotemporal craniotomy and in one case in spite of complete removal there was a recurrence. Retrospectively, when the MRI scan of this patient was reviewed there was imaging
evidence of invasion of the sphenoid diploe (Figure 6b) which we feel should be drilled to avoid recurrences. A combined transpalatal and transmaxillary approach avoids sequelae like neuralgia, hearing loss, trismus and cosmetic defects that are associated with lateral approaches.6 Lateral approaches should be used only if either lateral extensions of the angiofibroma are very large, extending up to the temporal region, or if there are posterior extensions through the foramen ovale (Figure 6c). These extensions are not easily accessible by a combined transpalatal and transmaxillary approach and usually residual tumour remains if this approach is used. No cosmetic deformity, facial scar or facial asymmetry was produced with a combined transpalatal and transmaxillary approach as ethmoidectomy, nasal septal trauma, facial osteotomies and metal plate fixation, which are used in other anterior approaches and are usually considered to produce such defects,21 were avoided in this approach. This is very significant considering that the majority of juvenile nasopharyngeal angiofibromas are young and growth of the maxillary growth continues after the age of 20 years and approximately 40 per cent of maxillary growth occurs after the age of 12 years.22 Fagan et al.21 used facial translocation procedures along with infratemporal fossa approaches to remove angiofibromas and found a 37.5 per cent recurrence rate and patients usually suffered from cosmetic defects along with retarded facial growth. They observed that recurrences occurred in spite of adequate visualisation of the tumour and suggested lack of encapsulation and submucosal spread as risk factors of incomplete resection. This further justifies relatively conservative surgical approaches used by us as meticulous removal not wide exposure is required for not only complete removal of the angiofibroma but also to prevent recurrences.

Conclusions

Extensions into the pterygoid fossa, clivus and basisphenoid, intracranial extensions medial to the cavernous sinus, evident residual tumours, invasion of the sphenoid diploe through a widened pterygoid canal, feeders from the internal carotid artery and young age were risk factors found associated with recurrence of juvenile nasopharyngeal angiofibroma. A combined transpalatal and transmaxillary approach with lazy S incision followed by nibbling of the medial pterygoid plate can expose the above mentioned areas with minimum morbidity. The majority of recurrences occurred in cases where there was imaging evidence of invasion and expansion of the sphenoid diploe through the pterygoid canal. Small lobules of angiofibromas remain in the cancellous bone of the sphenoid and pterygoid canal which should be regularly drilled to decrease the incidence of recurrences.

References

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Address for correspondence:
Dr Rajan Syal,
Neuro-otology Unit,
Department of Neuro-surgery,
Sanjay Gandhi Post Graduate Institute of Medical Sciences,
Raibarely Road,
Lucknow (UP) – 226 014 India.

E-mail: rajans@sgpgi.ac.in

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