Solitary fibrous tumours of the lacrimal sac: A case study and comparison

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Abstract
Solitary fibrous tumours are rare benign spindle cell neoplasms that arise uncommonly in the lacrimal sac but can occur in numerous other locations. Due to their rarity, heterogeneity of clinical presentations and histological appearances, they may easily be missed or misdiagnosed. In this paper we report a case of this rare tumour and compare our findings with the other 4 available cases in the English literature.

Case report
A 62 year old Chinese male presented with a 3-month history of right epiphora and blurring of vision. On examination, right eye hyperglobus (Figure 1) was noted and pupillary examination showed a relative afferent pupillary defect. On palpation there was firmness over the right lower eyelid. Flexible nasendoscopy was unremarkable and no mass was noted in the inferior meatus. MRI and CT orbits revealed a 3.6cm mass centred in the right lacrimal sac with extension into the proximal right nasolacrimal duct. Mass effect and indentation of medial aspect of right globe was noted, displacing it superolaterally, and there was no signal abnormality or enhancement in adjacent orbital fat (Figures 3-5). He underwent an endoscopic biopsy and histology showed a spindle cell tumour (Figure 2) with immunohistochemical staining positive for CD34. Decision was made for surgical resection of the tumour. Patient underwent a right lateral rhinotomy and excision of right lacrimal sac tumour. Intraoperatively, a 3cm right lacrimal sac tumour was noted with a clear plane between tumour and right periorbita laterally. The tumour did not breach the skull base (Figure 6a and 6b).

Histopathology revealed a cellular spindle cell lesion featuring fascicles and storiform architecture with occasional foci of staghorn vessels, perivascular hyalinization and keloidal collagen bundles. There was mild to moderate atypia, focal necrosis and a mitoses rate up to 6-7 per 10 HPF.

Postoperatively the patient’s hyperglobus improved (Figure 7), but he developed a right ectropion which was surgically corrected at a later date. The patient was last reviewed 16 months post-operatively, and flexible nasendoscopy showed no recurrence (Figure 8).

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Received: May 05, 2018; Accepted: May 21, 2018; Published: May 24, 2018
Caroline LCY (2018) Solitary fibrous tumours of the lacrimal sac: A case study and comparison

Figure 3. MRI T2 weight image of right lacrimal sac tumour, coronal cut

Figure 4. MRI T1 weight image of right lacrimal sac tumour, coronal cut

Figure 5: MRI T1 weight image of right lacrimal sac tumour, axial cut

Figures 6a & 6b: Intraoperative photo of right lacrimal sac tumour
Discussion

Solitary fibrous tumours (SFT) are a rare benign spindle cell neoplasm of mesenchymal origin. It was initially described in the pleura but now is recognized anywhere in the body including soft tissue and viscera of the abdomen, pelvis and trunk. It can also arise in the head and neck region, which represents 6% of all SFTs [1]. The sinonasal tract and orbit were the most common extracranial head and neck sites involved, accounting for 30% and 25% of head and neck SFTs [1].

To the best of our knowledge, we have found 4 case reports of solitary fibrous tumour in the lacrimal sac between 1999 and 2017 [2-4]. Table 1 lists the clinical features and Table 2 lists the histological findings by case.

SFTs may present across varied age groups – in these five patients (including our case) the median age at diagnosis was 44 years old with a range of 23 to 67 years. Out of the 5 patients, 2 were male and 3 were female. Due to the rarity of these tumours it is difficult to conclude if there is a gender predilection. However, based on literature of other SFTs, no gender predominance was noted [5].

Three cases presented with epiphora, four cases presented with a medial canthal mass, while in our case both were present. In the cases presenting with a mass, they had a prolonged history ranging from 3 to 12 months in duration, suggesting that this tumour is slow growing. Recurrent dacryocystitis was not a consistent feature across these cases.

The diagnosis of SFTs is challenging as it does not have distinct histopathological features [2]. Microscopically, SFTs typically consist of ovoid to spindle cells that are haphazardly arranged with varying cellular densities in admixture with stromal collagen bundles [1]. However due to histological diversity, SFTs can mimic other tumours such as mesotheliomas and sarcomas [6,7]. Hence immunohistochemical staining has proven useful in establishing the diagnosis. Comparing the 4 cases of lacrimal sac SFTs, immunohistochemistry was positive for CD-34 and vimetin in all cases and negative for S-100 protein and desmin. Mitoses do not commonly exceed 3/10 HPF [8].

A multi-institutional retrospective case series of 88 cases of head and neck SFTs, analysed that size and mitoses rate of >4/10 HPF were associated with risk of recurrence [1]. In cases of lacrimal sac SFTs reported so far, mitoses were not commonly found, however, in our case mitoses rate of 6-7/10 HPF was noted. Based on the few reported cases alone, it is difficult to predict the prognosis of lacrimal sac SFTs, however in all cases reported thus far, the patient has remained recurrence free up to 1 year after complete resection [2].

Tumour recurrence is also associated with incomplete excision as reported in the first 2 cases of lacrimal sac SFTs, in which their patients had undergone excision 4 months to 7 years prior to re-presenting. In our case, histology showed that the tumour extended to the superior margin of the specimen, which corresponds to the skull base. However intra-operatively, the tumour was noted to be abutting but not breaching the skull base. In addition, further drilling of the skull base was done. Our patient has been recurrence free up to 16 months post-op.
Table 1. Clinical features of lacrimal sac solitary fibrous tumours

<table>
<thead>
<tr>
<th>Author</th>
<th>Rumelt 2003</th>
<th>Kurdi 2014</th>
<th>Woo (case 1)</th>
<th>Woo (case 2)</th>
<th>Lee 2018</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>67</td>
<td>44</td>
<td>23</td>
<td>34</td>
<td>62</td>
</tr>
<tr>
<td>Gender</td>
<td>Male</td>
<td>Female</td>
<td>Male</td>
<td>Female</td>
<td>Female</td>
</tr>
<tr>
<td>Presenting Symptoms</td>
<td>Epiphora</td>
<td>Epiphora/ Medial canthal mass</td>
<td>Medial canthal mass</td>
<td>Medial canthal mass</td>
<td>Epiphora/ Blurring of vision/ Medial canthal mass</td>
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</tbody>
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Table 2. Histological and immunohistochemical features of lacrimal sac solitary fibrous tumours

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<tbody>
<tr>
<td>Mitoses</td>
<td>No mitotic activity</td>
<td>No mitotic activity</td>
<td>Low mitotic activity</td>
<td>Low mitotic activity</td>
<td>Mitotic rate 6-7/10 HPF</td>
</tr>
<tr>
<td>Necrosis</td>
<td>Absent</td>
<td>Absent</td>
<td>Not reported</td>
<td>Not reported</td>
<td>Present (&lt;5%)</td>
</tr>
<tr>
<td>Margins</td>
<td>Clear (tumour within the sac submucosa)</td>
<td>Not reported</td>
<td>Clear</td>
<td>Clear</td>
<td>Tumour extended to superior margin (skull base)</td>
</tr>
<tr>
<td>CD-34</td>
<td>Positive</td>
<td>Positive</td>
<td>Positive</td>
<td>Positive</td>
<td>Not available</td>
</tr>
<tr>
<td>Vimentin</td>
<td>Positive</td>
<td>Positive</td>
<td>Positive</td>
<td>Positive</td>
<td>Not available</td>
</tr>
<tr>
<td>S-100 protein</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td>Not available</td>
</tr>
<tr>
<td>Desmin</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td>Not available</td>
</tr>
</tbody>
</table>

References

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