Orbital solitary fibrous tumor: a clinicopathologic study from a Chinese tertiary hospital with a literature review

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Purpose: To report the clinical features, imaging manifestations, histopathology, and immunohistochemical features of several cases of orbital solitary fibrous tumors (SFTs) in a Chinese tertiary hospital, and to undertake a literature review of this rare disease.

Methods: A non-comparative retrospective review of clinical presentations, imaging manifestations, histopathology, and immunohistochemical features as well as the management and disease outcomes of patients with orbital SFT was conducted along with a review of orbital SFT cases in the literature.

Results: This study includes two male and two female patients, with an average age of 53 years. Common presentations among these patients included a palpable subcutaneous mass, swelling of the eyelid, proptosis, diplopia, and vision disturbance. Three patients (cases 2–4) underwent imaging scans. All patients had complete surgical excisions and the tissue was subjected to pathological analysis. One patient (Case 4) experienced a recurrence with malignant transformation and received a re-excision surgery and postoperative radiotherapy. All patients remain alive and well after a minimum follow-up of 12 months (range 12–34 months).

Conclusion: Despite its rare occurrence, we suggest that the possibility of orbital SFTs needs to be considered when a painless, slowly growing orbital mass is identified. Typical characteristic magnetic resonance imaging features of orbital SFTs are iso- or hypointense signals on T1 and T2-weighted images, with marked enhancement. A positive cluster of differentiation 34 (CD34) staining is an important diagnostic clue favoring SFT. Some orbital SFTs are infiltrating, aggressive, or recur with malignant transformation. Therefore, regular long-term follow-up after complete excision is mandatory.

Keywords: orbit, CD34, malignant transformation, MR

Introduction
The solitary fibrous tumor (SFT) is a rare spindle cell tumor originating from the mesenchyme and was first described in pleural tissue in 1931.1 A recent study has reported that SFT is found in extrapleural locations such as lung, liver, pericardium, peritoneum, mediastinum, nasal cavities, thyroid, parotid gland, and the orbit.2 SFT in the orbit is extremely rare. Since the first report of an orbital SFT in 1994,3,4 only 90 cases have been described in the English literature to date.5–15 Most orbital SFTs are described as isolated case reports. Herein, we present four orbital SFT cases and conduct a review of the literature discussing the presentation, imaging, histopathology, management, and outcomes of this rare disease.
Methods
Clinical data
A non-comparative retrospective review was conducted for patients presenting to our hospital with a pathological diagnosis of orbital SFT between 2014 and 2017. The study was approved by the Human Subjects Ethics Subcommittee of the First Affiliated Hospital of Zhejiang University, and it adhered to the tenets of the Declaration of Helsinki. Written informed consent for the publication of case details was obtained from all patients in our study. Simultaneously, a review of the literature was done using PubMed to search combinations of the terms “orbital solitary fibrous tumor.”

Histopathology and immunohistochemistry
Surgical samples were collected and embedded in paraffin for histological and immunohistochemical analyses. Immunohistochemical studies for cluster of differentiation 34 (CD34), cluster of differentiation 99 (CD99), B-cell lymphoma 2 (Bcl-2), smooth muscle actin (SMA), S-100 protein, Ki-67, cytokeratin (CK), Desmin, and cluster of differentiation 117 (CD117) were undertaken on paraffin sections in all samples. All antibodies were obtained from OriGene (Rockville, MD, USA).

Results
Case 1
A 37-year-old woman was referred to our department with a complaint of swelling in the right lower eyelid for 4 years. A focal, firm, nontender swelling with well-defined margins was palpated in the inferomedial side of her right orbit. The mass was completely excised via a subciliary approach. Gross examination of the surgical specimen disclosed a well-encapsulated 25×20 mm oval mass. On histopathological study, the tumor was composed of haphazardly arranged spindle cells with bland nuclei and inconspicuous nucleoli (Figure 1). Immunohistochemical analysis showed that the tumor cells exhibited diffuse immunoreactivity for CD34 and CD99 and was approximately 5% positive for Ki-67 (Table 1). These observations were consistent with the diagnosis of SFT.

This patient is still alive and well, without local relapse after 27 months.

Case 2
A 51-year-old man came to our department with a palpable subcutaneous mass in the right lacrimal sac area which was first noticed 6 months earlier. Findings on ophthalmic examination were normal, except for a movable mass without tenderness.

Figure 1  Histopathological and immunohistochemical staining of tumors in our cases.
Notes: (A) The tumor is composed of haphazardly arranged spindle cells with bland nuclei and inconspicuous nucleoli. (Case 1; HE staining; original magnification: ×200). (B) Variously collagenous and numerous endothelium-lined vascular channels are noted. (Case 2; arrow shows an occasional vascular channel with a staghorn appearance; HE staining; original magnification: ×200). (C) Tumor cells with oval nuclei, conspicuous nucleoli, and mitotic figure are noted. (recurrent sample of Case 4; HE staining; original magnification: ×200). (D) Tumor cells are positive for CD34. (recurrent sample of Case 4; original magnification: ×200). (E) The tumor cells are positive for Bcl-2 (Case 2; original magnification: ×200). (F) The tumor cells are positive for Ki-67 (>10%), which indicates malignant transformation (recurrent sample of Case 4; original magnification: ×200).
Abbreviations: HE, hematoxylin and eosin; CD34, cluster of differentiation 34; Bcl-2, B-cell lymphoma 2.
Orbital magnetic resonance (MR) imaging showed an approximately 20×15 mm, well-circumscribed circular tumor in the subcutaneous region of the right lacrimal sac area, presenting with slight low signal intensity on T1 and T2 images, heterogeneous intermediate signal intensity on fat-suppression images, and diffusely heterogeneous enhancement after intravenous gadolinium administration (Figure 2).

The mass was completely removed via a medial canthus split approach. Histopathologically, the tumor was composed of spindle cells and the stroma was variably collagenous. The tumor showed numerous endothelium-lined vascular channels, some of which showed an occasional staghorn appearance (Figure 1). Immunohistological study showed positivity for CD34 and Bcl-2, and weak positivity for Ki-67 (<5%; Table 1 and Figure 1). These findings support the diagnosis of SFT.

This patient is alive and has kept well for 34 months.

### Case 3

A 65-year-old woman was referred to our department, and presented with a 1-year history of a painless, palpable nasal mass in the left orbit. A 20×15 mm, well-circumscribed mass

<table>
<thead>
<tr>
<th>Case No.</th>
<th>CD34</th>
<th>CD99</th>
<th>Bcl-2</th>
<th>SMA</th>
<th>S-100</th>
<th>CK</th>
<th>Desmin</th>
<th>CD117</th>
<th>Ki-67</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case 1</td>
<td>+</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>+ (5%)</td>
</tr>
<tr>
<td>Case 2</td>
<td>+</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>+ (&lt;5%)</td>
</tr>
<tr>
<td>Case 3</td>
<td>+</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>+ (&lt;5%)</td>
</tr>
<tr>
<td>Case 4</td>
<td>+</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>+ (&lt;5%)</td>
</tr>
<tr>
<td>2nd operation</td>
<td>+</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>−</td>
<td>+ (&gt;10%)</td>
</tr>
</tbody>
</table>

**Abbreviations:** CD, cluster of differentiation; Bcl-2, B-cell lymphoma 2; SMA, smooth muscle actin; CK, cytokeratin; +, positive; −, negative.

Figure 2 MRI scan of Case 2 shows a well-circumscribed circular mass in the subcutaneous region of the right lacrimal sac area.

**Notes:** (A) T1 image, (B) T2 image, and (C) post-contrast T1 image.

**Abbreviation:** MR, magnetic resonance.
was palpated in the nasal region of the left upper lid, and the rest of the ophthalmic examination was unremarkable.

Orbital MR imaging revealed a well-circumscribed, 20×15 mm subcutaneous mass in the superomedial side of her left orbit, with homogeneous isointense signals on T1 images and hypointense signals on T2 images. The lesion showed strong gadolinium enhancement (Figure 3).

Complete removal of the lesion was undertaken via an upper eyelid medial split approach. The mass was approximately 22×17×15 mm in size. Histopathological examination revealed the tumor was composed of uniform spindle cells containing fusiform nuclei, indistinct nucleoli, and a pale eosinophilic cytoplasm in a collagenous matrix. Small- to medium-sized endothelium-lined vascular channels were diffusely scattered. Moreover, immunohistochemistry showed strong positivity for CD34 staining and weak positivity for Ki-67 staining (<5%), consistent with a diagnosis of SFT (Table 1).

The patient is alive and healthy, without local relapse for at least 26 months.

Case 4

A 60-year-old man was referred to our department with complaints of proptosis and diplopia in his left eye for 2 months. He had a history of hepatocellular carcinoma and had undergone a transcatheter arterial chemoembolization operation 2 years before this presentation.

Ophthalmic examination showed an 8-mm proptosis, diffuse conjunctival congestion, and chemosis of his left eye as well as limitation of extraocular movement in all gazes. The visual acuity of the left eye had decreased to 1/20, and could not be corrected. A fundus examination showed papilloedema and venous congestion.

The orbital MR showed a giant, well-circumscribed mass in the muscle cone of the left orbit that could not be separated from the optic nerve and pushed the optic nerve to the nasal side. The lesion had two types of imaging features on MR. One presented with low signal intensity on T1 and T2 images, and the other presented with isointense signal intensity on T1 and T2 images. The latter presented a diffusely
heterogeneous enhancement after intravenous administration of gadolinium (Figure 4).

The tumor was removed by lateral orbitotomy. A part of the mass was grayish white with a size of 18×15×15 mm, whereas the rest of the mass was wine-red with a size of 3.0×2.5×2.5 mm. Histopathological study revealed haphazardly arranged spindle cells with dense collagenous fibrosis. No mitoses or areas of necrosis were identified. The tumor contained numerous blood vessels, ranging from capillary to medium-sized vessels. Immunohistochemistry showed strong positivity for CD34 staining and weak positivity for Ki-67 staining (<5%), which confirmed our diagnosis of SFT (Table 1).

The mass recurred 14 months after the operation. Orbital MR imaging revealed a well-circumscribed 37×34×23 mm mass in the muscle cone of the left orbit with isointense signal intensity on T1 images, and isointense to slight hyperintense signal intensity on T2 images. A post-contrast T1-weighted image revealed a heterogeneous enhancement of the lesion (Figure 5).

An extensive resection was done. Preoperatively, the recurrent mass was identified in the muscle cone and was found to be adherent to the optic nerve. Gross examination revealed the tumor was grayish white with a size of 35×30×25 mm. Histopathologically, the tumor was composed of spindle cells, a section of which had oval nuclei and conspicuous nucleoli. Mitotic figures were noted, with a transition into a high-grade component (Figure 1). Moreover, these tumor cells showed strong and diffuse positivity for CD34 staining and >10% cells were positive on Ki-67 staining, suggesting malignant degeneration of the tumor (Table 1 and Figure 1).

The patient was referred to the department of radiation oncology for radiotherapy. There was no evidence of recurrence during a 12-month follow-up period.

Discussion

The SFT, first identified by Klemperer and Rabin in 1931, is a rare tumor which is described as a mesenchymal neoplasm originating in the pleural structures.1 Subsequently, it has also been reported in extrapleural locations, such as the upper airway tract, nasal and paranasal sinuses, parotid and salivary glands, thyroid, lung, mediastinum, pericardium, peritoneum, liver, spine, soft tissues, and orbit.16–21 An orbital SFT was first described by Westra et al and Dorfman et al in 19943,4 and, since then, approximately 90 cases have been reported in ophthalmologic and pathologic journals.5–15

Clinical features

Orbital SFT has been documented across a wide age range from 9 to 76 years, predominating in the fifth decade, with no sex predilection according to previous reports.22–25 In this study, the four cases described include two men and two women, with a mean age of 53 (range 37–65 years), which was consistent with previous reports (Table 2).

In a study reported by Romer et al, 25 cases were identified in the left orbit and 14 in the right orbit.26 In our study,
two cases were found in the right orbit and two in the left (Table 2).

According to the reviews from Le et al, orbital SFTs may occur in a variety of areas in the orbital space: 20% in the superomedial orbital quadrant, 15% in the medial orbit, 13% in the superotemporal area, 9.5% in the lacrimal area, and 7% in the inferotemporal area. In our study, the tumor in Case 1 was in the lower eyelid, and in the lacrimal sac area in Case 2. Tumors in cases 3 and 4 were located in the superomedial orbital quadrant and muscle cone, respectively.

Orbital SFT commonly presents as a slowly progressive, unilateral, painless proptosis and may be associated with eyelid swelling, vision disturbances, a palpable mass, tearing, and ptosis. In the present study, a palpable subcutaneous mass was the most frequent initial feature. Proptosis and vision disturbances were observed in Case 4 (Table 2). Orbital SFTs were less commonly associated with optic nerve dysfunction and extraocular muscle impairment depending on their size and location.

**MR imaging presentation**

On imaging, an orbital SFT presents as well-defined soft tissue masses with heterogeneously or homogeneously strong enhancement on computed tomography (CT) and MR images. Furthermore, internal calcifications and necrosis have been reported occasionally. Compared with the cerebral cortex or extraocular muscle, the signal intensity of SFTs seen on MR images, in general, were isointense on T1-weighted images.

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Age (yrs)</th>
<th>Gender</th>
<th>Affected side</th>
<th>Location</th>
<th>Main clinical presentation</th>
<th>Clinical history</th>
<th>Treatment</th>
<th>Time of recurrence (mo)</th>
<th>Follow-up (mo)</th>
<th>Status</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>37</td>
<td>F</td>
<td>R</td>
<td>Lower eyelid</td>
<td>A palpable subcutaneous mass</td>
<td>4 yrs</td>
<td>Total excision</td>
<td>–</td>
<td>27</td>
<td>Alive</td>
</tr>
<tr>
<td>2</td>
<td>51</td>
<td>M</td>
<td>R</td>
<td>Lacrimal area</td>
<td>A palpable subcutaneous mass</td>
<td>6 mos</td>
<td>Total excision</td>
<td>–</td>
<td>34</td>
<td>Alive</td>
</tr>
<tr>
<td>3</td>
<td>65</td>
<td>F</td>
<td>L</td>
<td>Superomedial orbital quadrant</td>
<td>A palpable subcutaneous mass</td>
<td>1 yr</td>
<td>Total excision</td>
<td>–</td>
<td>26</td>
<td>Alive</td>
</tr>
<tr>
<td>4</td>
<td>60</td>
<td>M</td>
<td>L</td>
<td>In the muscle cone</td>
<td>Proptosis, vision disturbance</td>
<td>2 mos</td>
<td>Total excision</td>
<td>14</td>
<td>14</td>
<td>Alive with relapse</td>
</tr>
<tr>
<td>2nd operation</td>
<td>60</td>
<td>M</td>
<td>L</td>
<td>Local recurrence</td>
<td>Proptosis, vision disturbance</td>
<td>–</td>
<td>Total excision</td>
<td>–</td>
<td>12</td>
<td>Alive</td>
</tr>
</tbody>
</table>

**Notes:** (A, D) T1 image, (B, E) T2 image, and (C, F) post-contrast T1 image.

**Table 2: Summary of clinical features of orbital SFT in the present series**

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Age (yrs)</th>
<th>Gender</th>
<th>Affected side</th>
<th>Location</th>
<th>Main clinical presentation</th>
<th>Clinical history</th>
<th>Treatment</th>
<th>Time of recurrence (mo)</th>
<th>Follow-up (mo)</th>
<th>Status</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>37</td>
<td>F</td>
<td>R</td>
<td>Lower eyelid</td>
<td>A palpable subcutaneous mass</td>
<td>4 yrs</td>
<td>Total excision</td>
<td>–</td>
<td>27</td>
<td>Alive</td>
</tr>
<tr>
<td>2</td>
<td>51</td>
<td>M</td>
<td>R</td>
<td>Lacrimal area</td>
<td>A palpable subcutaneous mass</td>
<td>6 mos</td>
<td>Total excision</td>
<td>–</td>
<td>34</td>
<td>Alive</td>
</tr>
<tr>
<td>3</td>
<td>65</td>
<td>F</td>
<td>L</td>
<td>Superomedial orbital quadrant</td>
<td>A palpable subcutaneous mass</td>
<td>1 yr</td>
<td>Total excision</td>
<td>–</td>
<td>26</td>
<td>Alive</td>
</tr>
<tr>
<td>4</td>
<td>60</td>
<td>M</td>
<td>L</td>
<td>In the muscle cone</td>
<td>Proptosis, vision disturbance</td>
<td>2 mos</td>
<td>Total excision</td>
<td>14</td>
<td>14</td>
<td>Alive with relapse</td>
</tr>
<tr>
<td>2nd operation</td>
<td>60</td>
<td>M</td>
<td>L</td>
<td>Local recurrence</td>
<td>Proptosis, vision disturbance</td>
<td>–</td>
<td>Total excision</td>
<td>–</td>
<td>12</td>
<td>Alive</td>
</tr>
</tbody>
</table>

**Abbreviations:** F, female; M, male; R, right; L, left; yr, years; mos, months; –, not applicable.
images and isointense to hypointense on T2-weighted images. Sometimes, a hypointense signal intensity can be seen within the lesion on both T1- and T2-weighted images, reflecting fibrous tissue with high collagen content.\textsuperscript{19,23,26,28–32} Moreover, a hyperintense signal can be observed within the lesion on T2-weighted images in some cases, causing signal intensity heterogeneity on T2-weighted images. These T2 hyperintense signals are related to internal hemorrhage, cystic degeneration, or relatively fresh fibrosis.\textsuperscript{19,32,33} Variations of the signal intensity on MR images reflected the differences of cellular components, collagen, fibroblasts, and degenerative change contained within individual tumors.

Homogeneous or heterogeneously attenuated enhancement is reported to be the most prominent feature of SFT that is revealed with CT and MR imaging, which is attributed to the prominent vascular channels within the tumor.\textsuperscript{19,23,31,34–37} Of the three lesions examined by MR in this study, one showed homogeneously marked enhancement (Case 3), and the rest showed heterogeneously marked enhancement (Cases 2 and 4). On the MR scan of Case 4, signal-intensity tubular structures were noted on T1, T2, and post-contrast T1-weighted MR images, which might represent fast-flow vessels within the tumor (Figure 5).

**Histopathologic and immunohistochemical features**

As clinical signs and radiological features are not distinctive, histological examination plays an important role in the diagnosis of orbital SFT. Tumors appeared as regular round masses, of hard consistency, and were homogeneously grayish white upon sectioning. All tumors in the present study were firm; neither cystic spaces nor necrosis were found. Tumors in cases 1, 2, and 3 were grayish white in color, whereas the mass in Case 4 was wine-red. Histological study revealed alternating cellular and hypocellular areas with spindle-shaped cells presenting an ovoid and irregular nucleus, inconspicuous nucleoli, and a scant, undefined eosinophilic cytoplasm.\textsuperscript{38} Spindle-shaped cells grow in a haphazard manner in a variable cellular stroma, which is described as a “patternless pattern” or keloid-like hyalinization. The tumor showed thick bands of collagen interspersed between the tumor cells, branching thin-walled vessels of varying caliber, and a “staghorn” vascular pattern, similar to a hemangiopericytoma (HPC). These features are consistent with those observed in our cases.

Microscopic features alone are insufficient to validate SFT. The key to identify SFT is to conduct immunohistochemical analysis.\textsuperscript{25,30} Strong and consistent positivity of CD34 is an important diagnostic marker of SFT, which demonstrates strong CD34 reactivity in 90–100% of cases,\textsuperscript{2,3} and the expression of CD34 may be lost in tumors that undergo malignant transformation.\textsuperscript{40} Meanwhile, SFT has been documented to exhibit strong and diffuse positivity with vimentin, CD99, and Bcl-2.\textsuperscript{38} Epithelial membrane antigen (EMA) and SMA might occasionally be detected, whereas SFT is usually negative for S-100 and CK.\textsuperscript{41}

Due to its histological similarities to other spindle-shaped cell tumors, SFT has been confused histologically with other lesions, such as HPC, fibrous histiocytoma, schwannoma, and menigioma. HPC displays a mixture of spindle-shaped tumor cells with oval nuclei and a small cytoplasm mixed with a network of thin-walled blood vessels or sinusoid-like spaces. HPC cells show high reactivity to the CD34 antibody and vimentin, but lack immunoreactivity to EMA.\textsuperscript{42} Fibrous histiocytoma usually consists of fibroblasts and prominent round histiocytic cells in a characteristic storiform pattern. Immunohistochemical staining consistently reveals that histiocytes are CD68 positive but are usually CD34 negative. Schwannomas display a characteristic morphology of alternating Antoni A and Antoni B areas and Verocay bodies. Immunohistochemical staining shows strong reactivity to neural markers such as S-100 protein, which is negative in a SFT.\textsuperscript{25} Fibrous menigioma often shows characteristic psamomma bodies, which are absent in SFTs. Besides, compared to SFT, a fibrous menigioma is usually better stained by an EMA antibody, but only mildly or focally positive on CD34 and Bcl-2 staining.\textsuperscript{41}

**Malignant transformation and aggressive behavior**

Most orbital SFTs are described in isolated case reports; thus their clinical behavior is currently unknown. It is generally believed that SFTs behave in a benign fashion and follow a non-aggressive course.

A few of the orbital SFT cases reported have displayed malignant histological features, defined by hypercellularity, pleomorphism, increased tumor size (≥5 cm), mitotic counts >4 per 10 high-power fields, and necrosis.\textsuperscript{29,38,44–47} Immunohistochemically, the expression of CD34 may be lost in tumors that undergo malignant transformation in cases of orbital SFTs.\textsuperscript{40,45} However, these pathologic factors do not always correlate with the clinical behavior of the tumor. Our literature search reviewed that only two of the five documented malignant orbital SFTs cases exhibited aggressive behavior, one of which had a recurrence whereas the other one had a local invasion.\textsuperscript{7,25,44,48,49} Some tumors with malignant
histologic findings remained indolent, whereas other tumors with completely benign histologic findings behaved aggressively. The aggressive behavior of orbital SFTs is defined by a local invasion, recurrence, and distant metastases.7,29,44,46,50–54 Nineteen of the 21 recurrent orbital SFTs were histologically benign. Young and Hardy reported a histologically benign orbital SFT case that invaded intracranial tissues.54 Malignant transformation of a recurrent SFT has been described in lung, pleura, and thigh SFT cases.55–57 A malignant variant of a recurrent SFT has been reported only in four cases, to our knowledge (Table 3). The intervals of malignant evolution varied from 6 months to 9 years after initial tumor excision.44,58–60 Here, we report the fifth case (Case 4) of a recurrent orbital SFT with malignant transformation. We report the fifth case (Case 4) of a recurrent orbital SFT with malignant transformation. We report the fifth case (Case 4) of a recurrent orbital SFT with malignant transformation.

**Table 3** Summary of reported recurrent orbital SFT with malignant transformation

<table>
<thead>
<tr>
<th>Source, year</th>
<th>Age</th>
<th>Gender</th>
<th>Affected side</th>
<th>Initial diagnosis</th>
<th>Initial treatment</th>
<th>Times of recurrence</th>
<th>Final diagnosis</th>
<th>Final treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carrera et al61</td>
<td>64</td>
<td>M</td>
<td>L</td>
<td>SFT</td>
<td>Total excision</td>
<td>4</td>
<td>Malignant SFT</td>
<td>Re-excision</td>
</tr>
<tr>
<td>Manousaridis et al60</td>
<td>53</td>
<td>F</td>
<td>R</td>
<td>SFT with focal malignant transformation</td>
<td>Total excision</td>
<td>1</td>
<td>Malignant SFT</td>
<td>Re-excision</td>
</tr>
<tr>
<td>Wang et al61</td>
<td>38</td>
<td>M</td>
<td>R</td>
<td>SFT</td>
<td>Total excision</td>
<td>2</td>
<td>Malignant SFT</td>
<td>Re-excision</td>
</tr>
<tr>
<td>Blandamura et al62</td>
<td>75</td>
<td>NM</td>
<td>R</td>
<td>SFT</td>
<td>Excision + radiotherapy</td>
<td>1</td>
<td>SFT with focal malignant transformation</td>
<td>Malignant SFT</td>
</tr>
<tr>
<td>Case 4 in the present study</td>
<td>60</td>
<td>M</td>
<td>L</td>
<td>SFT</td>
<td>Total excision</td>
<td>1</td>
<td>Malignant SFT</td>
<td>Re-excision + radiotherapy</td>
</tr>
</tbody>
</table>

Abbreviations: F, female; M, male; R, right; L, left; NM, not mentioned; SFT, solitary fibrous tumor.

Malignant transformation of a recurrent SFT has been reported only in four cases, to our knowledge (Table 3). The intervals of malignant evolution varied from 6 months to 9 years after initial tumor excision.44,58–60 Here, we report the fifth case (Case 4) of a recurrent orbital SFT with malignant transformation, which showed progressive histological dedifferentiation. Two types of malignant transformation have been reported previously: one is malignant or high-grade transformation from benign, low-grade, or intermediate-grade SFT, and the other is de novo occurrence of a malignant SFT.61 Malignant transformation of an SFT may have a worse consequence and need close follow-up.

**Treatment and prognosis**

Orbital SFTs are so rare that most published cases were individual case reports without documented extended follow-up; therefore, it is difficult to predict the true clinical course of these tumors.

Radical surgical resection seemed to be the most important prognostic factor. The extent of resection might be the most predictive factor for recurrence. In Chen et al's research, all four patients who underwent gross total removal did not have recurrence, whereas five of the six patients who had subtotal removal had a recurrence with a median recurrence time of 48 months.62 However, there are many factors that could prevent total resection, such as location, size, blood loss, and adhesion. Therefore, resecting as much tumor content as possible should be the goal of the treatment. In Case 4, the patient had a recurrence and malignant transformation. Therefore, a complete re-excision of the tumor and postoperative radiation therapy were carried out. Postoperative radiotherapy can be used as an adjuvant therapy when the lesion is not completely removed or in recurrent and malignant cases. Although there has been no sign of recurrence during a 12-month follow-up in this case, the prognosis should be guarded, and long-term follow-up is warranted.

**Conclusion**

Orbital SFTs should be suspected when confronting painless, slow-growing orbital masses although it is a rare entity. High CD34 expression is an important diagnostic marker favoring SFT. Some orbital SFTs are infiltrating, aggressive, or recurrent with malignant transformation. We report the fifth case (Case 4) of a recurrent orbital SFT with malignant transformation. Further research with a larger patient pool and extended follow-up is necessary to verify the clinical outcome of these tumors.

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**Disclosure**

The authors report no conflicts of interest in this work.

**References**

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