Orbital granulocytic sarcoma

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Abstract

Aim—Orbital granulocytic sarcoma is a localised tumour composed of cells of myeloid origin. Histological diagnosis can be difficult in patients with poorly differentiated orbital tumours and no evidence of systemic leukaemia. The naphthol AS-D chloracetate esterase (Leder stain) and immunohistochemical stains for lysozyme and MAC387 were used to determine the staining characteristics of these tumours. A case series of seven patients with orbital granulocytic sarcoma is presented.

Methods—Seven patients with orbital granulocytic sarcoma were studied. Haematoxylin and eosin, Leder, and lysozyme stained sections were available in seven cases. Unstained formalin fixed paraaffin embedded sections of seven cases were available for immunohistochemical evaluation using the avidin-biotin-complex technique for MAC387.

Results—The mean age of presentation of the orbital tumour was 8.8 years. Four patients presented with an orbital tumour before any systemic manifestations of leukaemia. In two cases the diagnosis of the orbital tumour and systemic leukaemia was made simultaneously. There was one case of established systemic myeloid leukaemia in remission with the subsequent development of orbital granulocytic sarcoma. Six of seven cases (86%) were positive for the Leder stain. Five of seven cases (71%) showed positive immunoreactivity with lysozyme. The immunohistochemical stain for MAC387 was positive in all seven cases (100%) including one case that was negative for both lysozyme and Leder stains.

Conclusions—Orbital granulocytic sarcoma is a tumour that affects children and young adults and must be differentiated from other orbital tumours that typically affect this age group. When a patient has a history of leukaemia and then develops orbital granulocytic sarcoma the diagnosis is usually not difficult. However, when the orbital tumour precedes the development of systemic leukaemia the diagnosis can be difficult for both clinician and pathologist. In addition, these tumours are often poorly differentiated making diagnosis more difficult. The differential diagnosis includes malignant lymphoma, rhabdomyosarcoma, and neuroblastoma. Histological features of myeloid differentiation such as eosinophilic granules within the cytoplasm may be helpful in establishing a diagnosis. However, these findings are often absent and not specific to granulocytic sarcoma. The naphthol-AS-D chloracetate esterase stain (Leder stain) greatly improved the ability of the pathologist to identify cells that contain myeloperoxidase. Similarly, immunohistochemical staining using anti-lysozyme antibodies for detection of lysozyme has proved useful in the identification of myeloid cells. Each of these stains in combination or alone is helpful when positive; however,
We present seven well documented cases of orbital granulocytic sarcoma. We studied the clinical features and compared the use of the Leder stain and two immunohistochemical markers for myeloid cells—lysozyme and MAC387.

Materials and methods
Seven cases of well documented orbital granulocytic sarcoma were studied at the McGill Ophthalmic Pathology Laboratory. Three of the cases were kindly provided by Dr R Font from the Baylor College of Medicine. The other four cases were obtained from the McGill Ophthalmic Pathology Registry. Cases were included in the study only if the myeloid origin of the tumour could be established by either positive Leder or lysozyme staining or the documentation of acute myeloid leukaemia in the bone marrow or peripheral blood. Clinical information was assessed for all seven cases. Histological features were evaluated using haematoxylin and eosin stained sections for degree of differentiation, mitotic figures, malignant features, and the presence of cytoplasmic eosinophilic granules. Paraffin blocks were obtained in all seven cases. The Leder stain was available for all seven cases. The Leder stain was considered positive if more than 5% of the cells stained positive. Immunohistochemical stains for lysozyme, MAC387, LCA, CD20 (L26), CD 43 (MT-1), CD 45 (LCA), desmin, and cytokeratin (CAM 5.2) were performed using the standard avidin-biotin-complex technique previously described. Appropriate positive and negative controls were used for all immunohistochemical stains. Stains were considered positive if staining could be unequivocally demonstrated in tumour cells. The anti-lysozyme, MAC387, CD 20, CD 45, and desmin antibodies were obtained from Dako Corporation, Carpinteria, CA, USA. Cytokeratin antibodies were obtained from Becton Dickinson Immunocytochemistry Division, San Jose, CA, USA. CD 43 antibodies were obtained from Biotest Diagnostics Denville, NJ, USA. Five cases of orbital lymphoma were obtained and stained with lysozyme and MAC387 as negative controls.

Results
Clinical data are summarised in Table 1. The age of presentation ranged from 5 months to 14 years with a mean age of 8.8 years. The orbital tumour preceded the development of systemic leukaemia in four cases. The mean time to development of systemic leukaemia was 5 months in these patients. In two cases the diagnosis of the orbital tumour and systemic leukaemia was made simultaneously. There was one case of established leukaemia in remission with development of orbital granulocytic sarcoma 8 months later.

Selected case reports
ORBITAL TUMOUR PRECEDES THE DEVELOPMENT OF LEUKAEMIA

Patient 1
A nine year old Latin-American boy was admitted to the hospital for rapidly progressive proptosis of the right globe. Two weeks before admission his right eye became painful. The patient had no previous illnesses.

Ophthalmic examination revealed right upper and lower lid oedema with proptosis and chemosis of the right eye (Fig 1A).

Computed tomography (CT) revealed a large irregular mass in the right superior orbit approximately 2 cm in diameter. Systemic examination, including a complete blood count and chest x ray, was negative.

An excisional biopsy revealed a greenish tumour, 20 mm in diameter. Histopathological examination revealed an infiltrating neoplasm composed of poorly differentiated cells with round or reniform nuclei, prominent nucleoli, and many mitotic figures. The scant to moderate cytoplasm contained eosinophilic granules. The Leder, lysozyme, and MAC387 stains were positive (Fig 1B, C, D). A diagnosis of orbital granulocytic sarcoma was made.

Eleven months after the onset of the orbital tumour, the patient developed acute myelogenous leukaemia with blast cells in the peripheral blood. The patient underwent polychemotherapy.

ORBITAL TUMOUR AND LEUKAEMIA DISCOVERED SIMULTANEOUSLY

Patient 2
A 5 month old Spanish boy was admitted to the paediatric unit for evaluation of rapidly progressive proptosis of the left globe. Five days before admission his left eye became proptotic with left upper lid fullness and epiphora.

The initial ophthalmic examination revealed unilateral left globe proptosis with downward and outward displacement of the eye without any signs of inflammation. A firm mass measuring approximately 5 x 5 mm could be palpated in the superior nasal quadrant of the left orbit.

A complete physical examination was unremarkable. CT revealed a homogeneous mass with uniform contrast enhancement located in the left superior orbit, involving the lacrimal gland.

The Leder stain was positive. Immunohistochemistry was negative for actin-desmin and L26 (CD 20), and positive for LCA (CD 45), lysozyme, and MAC387.

A bone marrow biopsy demonstrated a monomorphic hypercellular proliferation of
A 14 year old boy with acute myeloid leukaemia was treated with polychemotherapy. He was in remission for 8 months when he developed rapidly progressive proptosis of the right globe. Ophthalmic examination revealed left upper and lower lid erythema and oedema. The left eye was proptotic with conjunctival chemosis. Palpation revealed a firm mass in the superotemporal left orbit.

CT demonstrated a large homogeneous mass in the left orbit. A biopsy was performed. Histological examination revealed a monotonous proliferation of poorly differentiated cells. The nuclei were round to oval with nuclei indented. There were prominent nucleoli and moderate mitotic activity. There is scant cytoplasm with no eosinophilic granules. The Leder stain was positive. A diagnosis of orbital granulocytic sarcoma was made. The patient was again treated with chemotherapy and has since been in remission.

**Discussion**

Granulocytic sarcoma of the orbit can present in infancy to 61 years of age. Despite this broad age range of presentation, these tumours...
Granulocytic sarcoma can occur in a variety of locations throughout the body including the orbit. The most common sites of involvement are bone, soft tissues, lymph nodes, skin, and kidney. Bone involvement is usually subperiosteal and most frequently located in the vertebrae, sternum, orbit, and cranium. In Liu et al’s study of 23 necropsy cases of granulocytic sarcoma, exophthalmos was present in seven cases and was the most common observable sign of tumour nodule.

Certain paediatric populations are known to have a higher incidence of orbital granulocytic sarcoma. Reports in the literature indicate that African, Middle Eastern, East Asian, and Latin American children have a disproportionately higher incidence. This series also supports geographic variation of this condition as all of the cases were of Spanish descent. This study also revealed a slight male predominance which is in accordance with previously published literature.

The clinical features of orbital granulocytic sarcoma can vary considerably. Proptosis is the most common presenting clinical sign. In fact, Zimmerman and Font found 29 of 33 patients with orbital granulocytic sarcoma presented with signs of proptosis. Grossniklaus and Wojno described a case presenting as periorbital cellulitis which represented myeloblastic invasion of eyelid dermis. Various other presentations have been described including lacrimal gland involvement, eyelid tumour, post-traumatic, iris tumour, uveitis, conjunctival mass, and scleral mass.

The most puzzling feature of orbital granulocytic sarcoma is its variable temporal onset relative to the development of systemic leukaemia. The occurrence of the orbital granulocytic sarcoma before the development of systemic leukaemia is not uncommon and was demonstrated in 29 of 33 cases (88%) in the largest study of orbital granulocytic sarcoma by Zimmerman and Font. Numerous other instances of this phenomenon can be found in the literature. The subsequent development of leukaemia usually occurs within 1 year. The AFIP series by Zimmerman and Font showed that the vast majority of cases had haematological evidence within 2 months. However, there are reports of prolonged intervals between the development of granulocytic sarcoma and systemic leukaemia varying from 2 to 16 years. In our series, the development of orbital granulocytic sarcoma occurred before any evidence of systemic leukaemia in four of seven cases (57%) with a mean time to development of leukaemia in either the peripheral blood or bone marrow of 5 months.

The differential diagnosis of orbital granulocytic sarcoma can be challenging, particularly when there is no evidence of systemic leukaemia. It is most frequently confused with malignant lymphoma, but can also be confused with rhadomyosarcoma, neuroblastoma, and other poorly differentiated tumours of childhood. Therefore, it cannot be overemphasised that any tumour of the head and neck region, including the orbit, can potentially represent infiltration by malignant myeloid cells.

In patients with known systemic leukaemia CT may help to distinguish between granulocytic sarcomas from haematomas and abscesses, which are possible complications of leukaemia. However, in patients who develop granulocytic sarcoma before leukaemia, CT or magnetic resonance imaging features are not specific enough to distinguish granulocytic neoplasms from other tumours.

To establish the diagnosis of orbital granulocytic sarcoma a biopsy is often required. There is a correlation between the degree of differentiation of these tumours and the accuracy of the initial diagnosis. Blastic tumours are correctly diagnosed 38% of the time, poorly differentiated tumours 47% of the time, and well differentiated tumours 54% of the time. Typically, these tumours are composed of a monotonous invasive proliferation of malignant cells. The nuclei can vary from round in undifferentiated cells, to indented in more differentiated cells. Nucleoli are frequently prominent and the number of mitotic figures is variable. Histological differentiation between granulocytic sarcoma and lymphoma may be facilitated if the tumour reveals cells with large nuclei with blastic chromatin and prominent nucleoli as this finding is rare in lymphomas. The amount of cytoplasm and the presence of eosinophilic granules varies depending on the degree of differentiation of the cells. The cytoplasm is scant with no granules in undifferentiated cells whereas in more differentiated cells the cytoplasm may be more abundant and exhibit eosinophilic granules. Granulocytic differentiation when present is clearly helpful in establishing the diagnosis. However, Niemen et al found in their series that only 20% of these tumours have a significant degree of differentiation.

Electron microscopy has been shown to be useful in establishing the diagnosis of granulocytic sarcoma. However, it is rarely used and since the development of the Leder stain and immunohistochemistry, electron microscopy is not required to establish the diagnosis.

The development of the Leder and immunohistochemical stains have enhanced the ability of the pathologist to establish the diagnosis of granulocytic sarcoma. The Leder stain identifies cells with esterase activity. This stain can be used on formalin fixed paraffin embedded sections. The Leder stain has been shown to
be positive in approximately 75% of cases in two retrospective series on granulocytic sarcoma. The results using the Leder stain in our study are similar to those previously published. A positive Leder stain is supportive of the diagnosis of granulocytic sarcoma but a negative stain does not rule out the diagnosis.

The immunohistochemical stain using anti-lysozyme antibody identifies cells that contain the enzyme lysozyme. The use of this marker in identifying tumours of myeloid origin has been established. The lysozyme stain has been shown to be positive in 60% to 89% of cases in previous studies of granulocytic sarcoma. More importantly, Neimen et al have shown that 98% of cases were identified positive for either Leder or lysozyme stains. Because one of the cases in our series gave negative results with both the Leder and lysozyme stains, we examined the use of monoclonal antibody MAC387. All seven cases stained with MAC387 demonstrated immunoreactivity. For the vast majority of cases, the combination of Leder and lysozyme stains will establish the diagnosis of granulocytic sarcoma. However, in cases of extremely poorly differentiated tumour in which the Leder and lysozyme stains give negative results, but clinical suspicion is high, MAC387 may prove useful in establishing the diagnosis.

Conclusions
Orbital granulocytic sarcoma may develop before, during, or after the occurrence of leukaemia in the peripheral blood and bone marrow. Histological diagnosis of granulocytic sarcoma can be difficult in children with no evidence of leukaemia, so a high index of suspicion by both the clinician and the pathologist is important in order to make the diagnosis. Special stains and immunohistochemistry play an important role in the diagnosis. The use of lysozyme and Leder stains in combination is useful in establishing the diagnosis of granulocytic sarcoma. MAC387 stain may provide additional information in cases in which the Leder and lysozyme stains give negative results and clinical suspicion is high.

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