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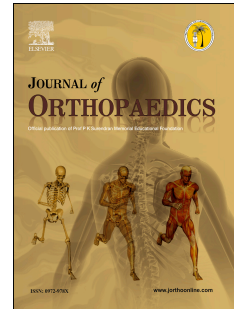
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CLEAR CELL SARCOMA- a review

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CLEAR CELL SARCOMA- a review

Abstract:

Clear cell sarcoma (CCS) previously known as malignant melanoma (MM) of the soft tissue, although, similar in morphology to MM, contemporary histopathologic and cytogenetic techniques have made this diagnosis obsolete, as it is now possible to distinguish between CCS and MM. CCS is often diagnosed in young adults with median age of 25 years. Overall mortality is generally poor, and the 5-year survival is between 40 - 60 %. Hence, early diagnosis and radical surgery are key in the treatment of this extremely rare malignancy of the soft tissue comprising only about 1 % of all sarcomas. This article present an overview of this rare malignancy.

Keywords: sarcoma; clear cell sarcoma; malignant melanoma of the soft tissue; treatment of clear cell sarcoma

Introduction

Clear cell sarcoma (CCS) is an extremely rare and aggressive subtype of sarcoma with melanocytic differentiation both immunohistochemically, ultrastructural and genomic^{1,2}. It comprises approximately 1 % of all diagnosed sarcomas³. Furthermore, the prognosis is poor due to predilection for metastasis at an early stage to the lymphatic system and the lungs⁴. CCS is thought to derive from the neural crest cells⁵. As a whole, sarcomas arise from the mesenchymal tissues, such as bone, muscle, fat, connective tissue, blood vessels and nerves⁵. Hence, corresponding tissue origin of clear cell sarcoma, which was first described by Enzinger et al in 1965⁶. The name CCS originates from the initial histopathological findings of clear cells, as the cytoplasm of the tumor cells may appear clear⁶. The term malignant melanoma of the soft tissue was previously used, however, contemporary histopathologic and cytogenetic techniques have made this diagnosis obsolete, as pathologists are able to distinguish between CCS and malignant melanoma today⁷. In 95 % of the cases, the CCS arises in the lower extremities, particularly the foot and ankle. It presents as an indolent, growing and painless mass situated in the deep soft tissue or beneath the fascia^{4,8-10}. CCS occurs at all ages, but is most often observed in the third decade¹¹⁻¹⁴. The mean age at the time of diagnosis is approximately 25 years and CCS is more common in Caucasians than in African Americans or Asians, without any gender predilection¹¹⁻¹⁴. The aetiology of CCS is virtually unknown, and the exact cause and the mechanisms of formation remain to be elucidated. Genetic defects occurring on account of certain chromosomal exchanges are thought to play a key role in the formation. Hence, CCS is repeatedly seen in relation to specific genetic transformations. Only a few known risk factors are described *e.g.* exposure to chemicals (vinyl chloride, arsenic), chronic tissue irritation (lymphedema, foreign body implants) and radiation⁵.

Diagnosis

Compared to the aforementioned, identifying and diagnosing CCS is challenging. Quick and accurate diagnosis is dependent on the level of experience of the doctor at the initial examination and fast referral to specialized units with relevant experience, such as tertiary sarcoma centres are imperative. The mean time from onset of symptoms to confirmation of the diagnosis is often several months, which could be the result of an initial suspicion of a benign tumor^{13,14}. This prolonged time to relevant examinations and to diagnosis may lead to a more advanced disease at the time of diagnosis and therefore a more sinister prognosis^{5,14,15}.

Radiological examination

Hitherto, the most relevant initial investigation in the diagnosis of CCS is magnetic resonance imaging (MRI). Currently, no exact radiological criteria to differentiate between malignant and benign tumors in the soft tissue exist^{7,16-18}. However, today a combination of different imaging tools is used to provide a tentative diagnosis, while the final diagnosis of CCS is made, based on the resulting or concomitant pathological examination.

MRI is the leading and most established modality to identify the primary tumor and its relations to adjacent structures¹⁷. In addition, conventional ultrasound and computed tomography (CT) may be utilized as supplements and primary investigation in the few cases of contraindications to the MRI or because these modalities are more widely available.

When diagnosing CCS, it is important to consider obtaining the best possible preoperative imaging simultaneously, as surgery is the primary and possible curative treatment and further delays in terms of planning radical surgery, may render the prognosis even worse. As such the MRI produces a clear image of the tumor, and its relations to local structures *e.g.* nerves and major blood vessels, which may result in reduced per operative morbidity due to better planning (*Figure 1*).

***** *Figure 1* near here *****

As CCS typically disseminate by haematogenous spread to the lungs or via the lymphatic system, Positron emission tomography–computed tomography (PET-CT) scan may be used to evaluate patients' status in terms of radical surgery.

Pathological examination

The final diagnosis of CCS is pathological. Macroscopically the CCS appears to be well-defined solid tumors, with mat grey color that frequently infiltrate tendons and aponeuroses. Microscopically findings are small compact nests with uniform neoplastic cells divided into variable sized clusters by fibrous septa along the tendons and aponeuroses^{2,6}. The tumor cells typically appear with a clear eosinophilic cytoplasm and a basophilic nucleolus (*Figure 2*). The eosinophilic cytoplasm is due to the accumulation of glycogen, which can be visualized using periodic (PAS) stain method. The tumor cells show none or a minimal rate of mitosis in concordance with the slow growth rate of CCS. Scattered giant cells with 10-16 nucleoli and scattered necrosis may also be identified in CCS².

***** *Figure 2* near here *****

Immunohistochemical studies of CCS show great similarity to the characteristics of malignant melanoma, including a strong expression of S-100, HMB-45, Melan-A and microphthalmia-associated transcription factor (MITF) in 81-97 % of the cases^{2,19}. Clear cell sarcoma is associated the chromosomal fusion of the Ewing's Sarcoma oncogene (EWS) and the cellular transcription factor (ATF1) into EWS/ATF1 oncogene²⁰. This chromosomal fusion due to (t[12; 22] [q13; q12]) can be detected with cytogenetic investigations, *e.g.* polymerase chain reaction (PCR) and

fluorescence in situ hybridization (FISH) in 90% of the cases²⁰. In addition, to the before mentioned translocation, a fusion between Ewing Sarcoma Breakpoint Region 1 (EWSR1) and Camp Responsive Element Binding Protein 1 (CREB1) is also known to give rise to CCS²¹. Studies have shown that EWSR1/ATFI fusion protein is able to bind and activate melanocyte specific MITF via SRY-related HMG-box 10 (SOX10) transcription factor, which results in the expression of the melanocytic phenotype, thus the malignant melanoma is a differential diagnosis^{22,23}.

Differential diagnosis

The diagnosis of CCS may be difficult and differential diagnosis are plenty and include melanoma, epithelioid malignant peripheral nerve sheath tumor, cellular blue naevus, melanotic schwannoma, paraganglioma-like dermal melanocytic tumor, synovial sarcoma (monophasic type), paraganglioma, epithelioid sarcoma and carcinomas²⁴. In comparison to melanoma although clinically distinctive, the initial pathological examination may be ambiguous, hence bear in mind that CCS is more deeply located, in close relation to tendons or aponeuroses and with no epidermal involvement. In contrast, a primary cutaneous nodular melanoma is usually situated in the dermis and tends to be more pleomorphic and enlarging more rapidly than CCS. Remember that melanoma metastasis can be found in deeper tissue (*Figure 1*). Melanotic schwannoma are rare and most often occurs in patients with Carney syndrome, together with spotty skin and myxomas. These schwannomas are situated in paraspinal or nerve plexus-related locations and frequently contains calcifications and very abundant melanin pigment²⁴.

Malignant peripheral nerve sheath tumor is, as indicated in the name, associated with a large peripheral nerve or neurofibromatosis and is histologically characterized by significant myxoid stroma (*i.e.* pale to light basophilic stroma) in haematoxylin and eosin stains, hyperchromatic

nuclei, brisk mitotic activity, negativity for HMB-45, melanocytic markers and cytogenetic abnormalities³. In contrast to CCS, paraganglioma-like dermal melanocytic tumor is a tumor of the extremities of only females, which presents as a dermal nodule with distinct histological appearance, large epithelioid cells with distinct large nuclei, prominent nucleoli and absence of genomic EWS rearrangement.

Treatment

Margin-free surgery is the primary treatment of CCS^{8,9,18,25}. Recommendations are surgical excision with a sufficient amount of surrounding healthy tissue. Most authors recommend a margin of one centimetre or more, although this is a matter of ongoing debate^{8,9,15}. Adjuvant treatment as discussed below is recommended in cases, where it is not possible to meet the recommendations of margins²⁶. Amputation following CCS is rarely used in contemporary medicine¹⁵. One third of patients if undergoing sentinel node procedure or subsequently lymph node dissection present with lymphatic metastasis and these procedures are discussed in regards of locoregional disease control⁸. The procedures seem to predict recurrent disease as well as survival but are not yet widely implemented.

Radiation therapy

Adjuvant radiation therapy can be used as a postoperative adjuvant treatment of CCS²⁷. It is commonly used in delicate situations, where the recommended margin of one centimetre of healthy tissue cannot be achieved and additional surgery is not feasible or refused by the patient^{18,28,29}.

It is unclear whether adjuvant radiotherapy has a beneficial effect on survival²⁷. It is recommended in cases of high grade (grade 2–3) and large (>5 centimetres) soft tissue sarcomas (level IIB)³⁰.

Grading are enumerated in *Figure 3*.

***** *Figure 3* near here *****

Clear cell sarcoma, however, typically has low mitotic activity and necrosis is only seen occasionally in these tumors, classifying most CCSs as grade 1 tumors²⁹. In cases with negative surgical margins, a radiation dose of 50 Gy is recommended in fractions of 1.8 to 2.0 Gy. If positive margins are present a postoperative radiation regime of 60-66 Gy is recommended²⁹.

Chemotherapy

If complete excision with sufficient margin is achieved, adjuvant treatments are not necessary²⁶. Chemotherapy is indicated in patients with metastatic disease but the efficiency still remains unclear, although it is being used as palliative treatment^{25,26,31}. Hence, in a retrospective study of 24 patients, investigating the response rate of metastatic CCS disease and the treatment with chemotherapy the authors found that treatment with anthracycline based chemotherapy was the only therapy with partial effect on metastatic CCS, as the response rate was 4 %¹³. Both cisplatin and ifosamide based protocols only managed stable disease in two patients with a progression free period of 11 weeks¹³. Similar findings, in another study in which advanced CCS disease was studied in 11 patients with a combination of doxorubicin, dacarbazine and ifosfamide, as only two patients achieved partial response or stable disease and it lasted less than six months³².

Postoperative control

An extensive follow-up program is advised and instituted in most countries, due to the high risk of metastases and relapse. The purpose is to exclude recurrence and metastases, in addition to address occupational health issues, practical needs, and to support self-care and negate long-term sequelae of the treatment.

The following is recommended. Histopathological low grade tumors (grade 1) are to be followed once every three months the first year, once every four months for two-three years and hereafter twice a year until the fifth postoperative year³³. Clinical monitoring is especially important when it comes to local recurrence and clinical examination should be supplemented with MRI on any suspicion³⁴. Patients with high grade tumors (grades 2-3) are recommended to follow a rigid scheme of controls every three months for the first three years and then twice a year through the fifth year³³. Even with a negative clinical examination, it is still recommended to have regular MRI of previous tumor site twice a year. The exact role of PET-CT and chest x-ray is not fully established but often used on suspicion in order to exclude metastases³⁵. Individualized control programs can occur in up to ten years postoperatively. Patients that have received chemotherapy should also be examined biochemically with liver, kidney, and heart function parameters because of the risk of side-effects.

Prognosis

The prognosis of CCS is poor because of the tendency to metastasize at an early disease-state and due to the rate of recurrence^{8,15,18}. Local recurrence rate may be as high as 84 % and 30 % of patients have metastases at the time of diagnosis^{7,15,36}. Negatively associated prognostic factors are necrosis, the number of mitosis, resection margin, anatomic location and tumor size^{11,15,26}. Tumor size, at the time of surgery, seems to be the key factor and is most strongly correlated to survival. Tumors larger than five centimetres are proven to recur more often because of micro metastasis^{11,19,25}.

Conclusion

CCS is a rare malignant disease with very low incidence of the soft tissue with numerous differential diagnoses. The incidence is very low and the prognosis is poor. Primary treatment is

radical surgical extirpation and an extensive postoperative follow-up program. The role of adjuvant therapy *i.e.* radiation and chemotherapy remains to be elucidated and larger prospective studies are required to establish the optimal treatment.

ACCEPTED MANUSCRIPT

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Figure 1:

MRI scan: An 80-year-old woman with subcutaneous soft tissue tumor in the lower extremities with relation to both tendons, skin and vital blood vessels. Primary differential diagnosis was sarcoma. A biopsy was performed. Histopathological examinations raised doubt regarding the diagnosis; was it malignant melanoma or clear cell sarcoma. Subsequently verified as a metastasis of malignant melanoma.

Figure 2

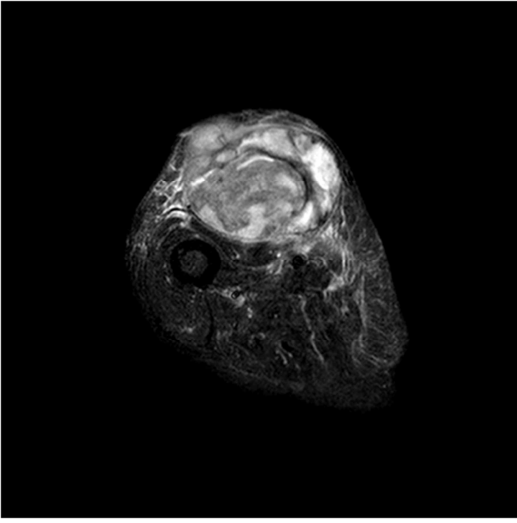
CCS, hematoxylin-eosin stain: There are irregular nests of cells separated by fibrous septa. Cells are fusiform and not pleomorphic. They have regular vesicular nuclei and prominent nucleoli with moderate to abundant eosinophilic or clear cytoplasm.

Figure 3:

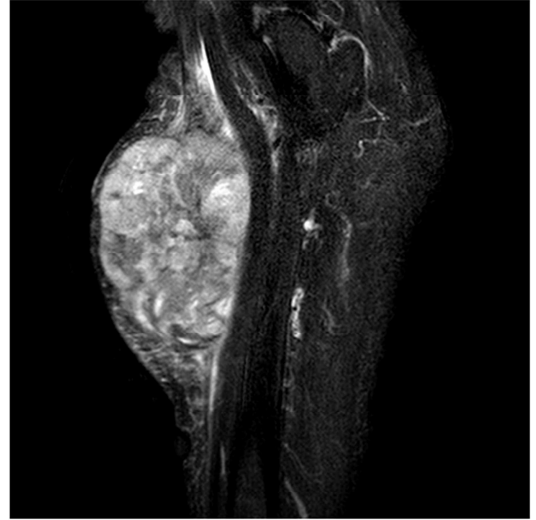
FNCLCC grading

Figure 1

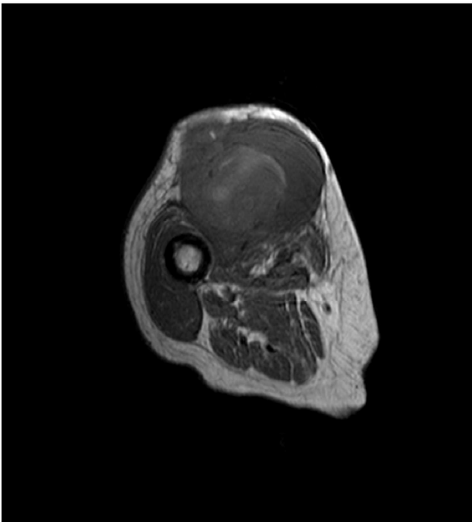
Horizontal section with contrast



Longitudinal section with contrast



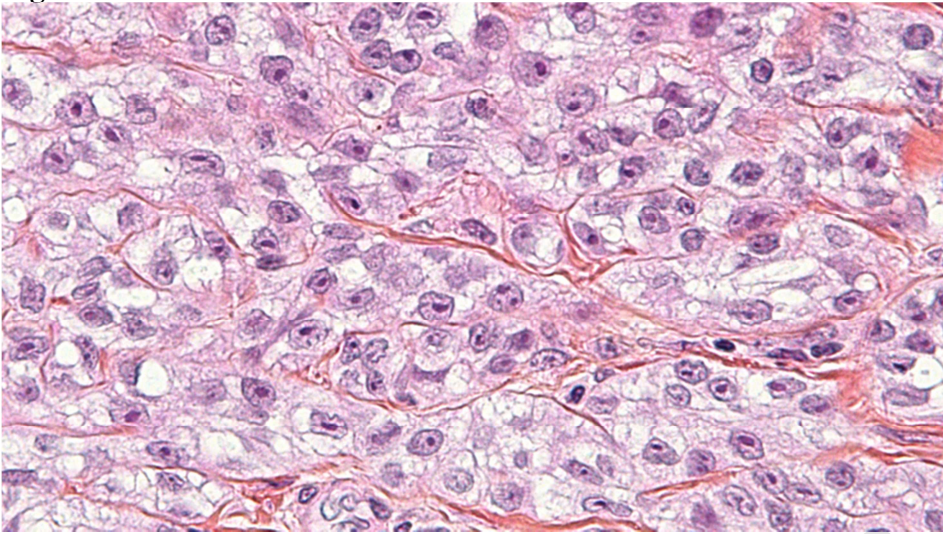
Horizontal section without contrast



Longitudinal section without contrast



ACCE

Figure 2:**Figure 3:**

Fédération Nationale des Centres de Lutte Contre le cancer; High Power Field

Tumor differentiation	Definition
Score 1	Sarcomas that closely resemble normal adult mesenchymal tissues
Score 2	Sarcomas for which histologic typing is certain
Score 3	Embryonal and undifferentiated sarcomas, synovial sarcoma and sarcomas of uncertain differentiation
Mitotic count	
Score 1	0-9 mitosis/ 10 hpf
Score 2	10-19 mitosis/ 10 hpf
Score 3	20≥ mitosis/ 10 hpf
Tumor necrosis	
Score 0	No necrosis
Score 1	<50% tumor necrosis
Score 2	≥50% tumor necrosis
Histologic grade	Tumor differentiation + mitotic count + tumor necrosis
Grade 1 (Low grade)	Total score: 2 or 3
Grade 2 (Intermediate grade)	Total score: 4 or 5
Grade 3 (High grade)	<50% tumor necrosis