



Imaging and Pathological Features of Alveolar Soft Part Sarcoma: Analysis of 16 Patients

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Abstract

Context Alveolar soft part sarcoma (ASPS) is a rare soft tissue tumor most commonly occurring in deep intramuscular plane of lower extremities of adolescents and young adults. It is a highly vascular, slow growing tumor with malignant potential having lung as the most common site of metastases at the time of presentation.

Aims The aim is to review the imaging findings of ASPS and determine characteristic imaging features of this rare tumor.

Materials and Methods Sixteen patients having histopathological diagnosis and preoperative imaging of ASPS attending the dedicated sarcoma clinic at our institute were included in the study. The demographic, clinical, and imaging data were retrieved from the case records and then evaluated for characteristic imaging features which may raise suspicion of ASPS.

Results The patients ranged from 3 to 72 years of age and with a slight male preponderance. Of the eight CECTs evaluated, 62.5% tumors showed well-defined lobulated margins, 87.5% cases showed intense enhancement with presence of feeder vessels. On CEMRI of 10 patients, 70% had well circumscribed lobulated margins with intense enhancement and tortuous flow voids in most of them. All cases showed T2 hyperintense signal. Fourteen of 16 (87.5%) patients had metastatic disease with lung as the most common site (92.8%).

Conclusion ASPS is a rare soft tissue sarcoma seen in children and young adults. Imaging may mimic a vascular malformation due to the presence of tortuous feeders. Misdiagnosis at an early stage may lead to later metastatic presentation of the disease, thus emphasizing the need to suspect it on imaging.

Keywords

- ▶ alveolar soft part sarcoma
- ▶ soft tissue sarcoma
- ▶ imaging
- ▶ feeding vessels
- ▶ metastatic disease

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Introduction

Soft tissue sarcomas (STS) are a rare malignancy with undifferentiated pleomorphic sarcoma (previously called malignant fibrous histiocytoma), liposarcoma, and leiomyosarcoma being the most frequent histologies in adults.¹ Alveolar soft part sarcoma (ASPS) comprises <1% of all STS with literature limited to case reports and series.² As per the 5th WHO classification of soft tissue tumors updated in 2020, ASPS falls under the malignant category of “tumors of uncertain differentiation”³ with extremities being the most common site of occurrence. It mostly occurs in children and young adults in the age group of 15 to 35 years.⁴ Nearly two-thirds of adults and one-third of the children have metastases to lungs, lymph nodes, and brain at the time of diagnosis. Metastases are known to occur after long tumor free intervals, approximately 38% of metastases occurring after 10 years of diagnosis with or without local site recurrence.^{4,5} Patients of ASPS usually have a poor prognosis. The mainstay of treatment remains total surgical excision and metastatectomy as it is unresponsive to conventional chemotherapy. Various targeted immunotherapies have been tried; however, no standard medical treatment protocol exists.⁶

Its typical histopathological appearance is arrangement of large epithelioid tumor cells in a pseudo“alveolar” pattern surrounded by sinusoidal vascular channels thus giving its name. ASPS is characterized by the unbalanced translocation t(X; 17)(p11; q25), which creates the ASPSCR1–TFE3 fusion protein, having angiogenic properties.^{7,8}

The imaging features often mimic vascular tumors and malformations due to its highly vascular nature and feeders, which may delay appropriate work-up.^{4,6} Further, it requires review by pathologists having expertise in sarcomas and application of special immunohistochemical or genetic markers for exact characterization. Hence, the initial suspicion must be raised on imaging. Contrast-enhanced MRI serves as a major tool in the work-up of soft tissue tumors to suggest the diagnosis and assess resectability. ASPS are usually located in the deep intramuscular compartment of an extremity, demonstrating lobulated well-defined margins showing hyperintense signal on T2WI and intense enhancement. Feeder vessels within and in the periphery of the tumor are typical.^{4,9–12} Metastases most commonly to the lung are present at the time of presentation.^{4,9,10} This study was conducted to review and determine the imaging features of ASPS patients visiting dedicated sarcoma clinic of our tertiary care center.

Materials and Methods

This retrospective study was conducted on patients with histopathological diagnosis of ASPS presenting to dedicated sarcoma clinic of our institute during the last 3 years (2016 to 2019), after obtaining approval from institute’s ethics committee, with waiver for informed consent. The demographic, clinical, imaging, and pathological details were obtained from the patients’ institutional case records.

Inclusion criteria: Histopathological diagnosis of ASPS in patients presenting to sarcoma clinic, confirmed by sarcoma pathologist of our institute. Of these patients, the ones with availability of at least one baseline cross-sectional imaging (computed tomography [CT] or magnetic resonance imaging [MRI] or positron emission tomography computed tomography [PET-CT]) of the primary tumor and adequate clinical details such as age and gender of the patient were included.

Exclusion criteria: Non availability of baseline CT/MRI/PET-CT.

Histopathology: The pathological diagnosis of ASPS was established by pathologist (A.B.) with an experience in sarcomas. In the patients, who underwent biopsy or resection outside, the slides were reviewed by the same sarcoma pathologist of our institute, before deciding on treatment plan.

Imaging analysis: Imaging details in the form of ultrasonography (USG), CT, MRI, or PET-CT were obtained from the patients’ files. For the patients’ whose imaging was available and was done outside, the films were reviewed by the radiologist (E.D.). The imaging features which were recorded, included: size, site, margins, attenuation/intensity/echogenicity/enhancement pattern/avidity based on the imaging modality, surrounding and internal vascularity, feeders in vicinity, intratumoral fat/necrosis/calcification, surrounding structural infiltration, distant metastases, and treatment response.

Definitions

Margins: The mass was categorized to have lobulated or circumscribed margins when the entire mass could be delineated with no infiltration or irregularity along the outline. However, few lobulations could be seen. Infiltrative margins were noted when there was evident involvement of surrounding muscles, bones or vessels at the site of irregular margins.

Enhancement characteristics were classified as intense, moderate, no or minimally enhancing as compared with surrounding muscles and vessels.

Internal imaging parameters were assessed on CT as iso/hypo/hyperdense and signal characteristics on T1WI and T2WI as equal or iso/hypo/hyperintense as compared with surrounding muscles—that is, same/low/higher than muscles, respectively.

Note was made of presence or absence of intratumoral and peripheral feeder vessels and distant metastases.

PET-CT was evaluated for similar findings as in CT. In addition, the SUV value and presence of any distant metastases >1 cm in size were also recorded.

Follow up: All the cases were followed up till February 2020 in the outpatient visits by interviews with the patients themselves or the attendants.

Statistical analysis: The collected data was transformed into variables, coded and entered in Microsoft Excel. Data were analyzed and statistically evaluated using SPSS-PC-17 version.

Quantitative data was expressed in ranges and mean while qualitative data were expressed in percentage.

Results

A total of 17 patients of ASPS reported to dedicated sarcoma clinic of our tertiary care center, one was excluded due to nonavailability of imaging. The age ranged from 3 to 72 years (mean 29.2 ± 15.2 years) and slight male preponderance (male: female 5:3).

Of these 16 patients, six patients had only MRI, four had only CT, one had MRI and PET-CT, one had MRI and CT, two had only PET-CT, one had USG, PET-CT as well as MRI and one had USG and MRI.

Size and site: The primary tumor size ranges from 3.6 to 17 cm (mean 8.29 cm).

Most common location was extremities (10/16 cases, 62.5%) with 80% (8/10) occurring in lower extremities. Other rare sites included were head and neck (2/16, 12.5%) and chest wall, intrathoracic or intra-abdominal. One patient presented with metastases from ASPS but without a known primary.

Histopathology: All the cases showed typical histomorphology of alveolar STS in the form of epithelioid appearing cells arranged in distinct nesting or organoid pattern with central dyscohesive cells giving appearance of pseudoalveolar pattern (►Fig. 1a). In one of the cases, the tumor was seen predominantly in nesting and solid pattern. Cells were generally monomorphic, polygonal shaped with distinct cytoplasmic boundaries (►Fig. 1b). The nucleus was round with finely dispersed chromatin with prominent nucleoli in most of them. Occasional mitotic figures were noted. Diastase treated periodic acid Schiff (PAS) stain highlights rhomboid-shaped or rod-shaped intracytoplasmic PAS positive inclusions (►Fig. 1c). All the cases demonstrated nuclear immunoreactivity on immunohistochemistry (IHC) for TFE-3.

Imaging: Two patients had undergone baseline ultrasound which revealed a circumscribed lobulated homogeneously hypoechoic mass showing multiple enlarged tortuous vessels within. Four patients had CECT of the local part and four patients had PET CT, hence a total of eight CTs were evaluated for primary tumor.

On CECT, the primary masses had circumscribed lobulated margins (five of eight patients, 62.5%), showed intense

enhancement on post contrast scans (seven of eight patients, 87.5%) with multiple feeder vessels within and in the periphery of the tumor (seven of eight patients, 87.5%). None of the tumor showed necrosis and only one case had the presence of calcification (►Table 1).

PET CT was available for four patients and the primary sites and metastatic disease was avid with an SUV of >2.5 .

MRI of 10 patients was evaluated. Most of the patients showed circumscribed lobulated margins (seven-tenths, 70%), iso to intermediate signal on T1WI (six-tenths, 60%) and hyperintense signal on T2WI (ten-tenths, 100%). Intense or moderate enhancement was seen on gadolinium-enhanced MRI with presence of multiple flow voids in 9 of 10 patients (90%) (►Table 2). The main role of MRI was to assess the resectability of tumor and 7 of 10 cases were found resectable on MRI. Out of these seven, only four patients could undergo surgery as rest had metastatic disease otherwise.

Metastatic disease: Fourteen of 16 (87.5%) patients had presence of metastases at the time of presentation in our clinic. Most common metastatic site was lung in 13 of 14 patients (92.8%). Two patients with lung metastases underwent metastatectomy for lung nodules. Next most common site of metastasis was bones (4/14, 28.6%). Five patients had metastases to more than one site which included liver (three), adrenal (one), and brain (two), and lymph nodes (one).

Follow up: Most of the cases were operated outside hospitals. Recurrence post-surgery occurred in six of 16 (37.5%) patients; revision surgery was done in one and surgical excision with positive margins was obtained in another patient. Medical treatment was given as first line treatment in seven of 16 patients for locally unresectable disease, extensive metastases, or unknown primary tumor.

Till last follow-up all 16 patients were alive and had stable disease.

Discussion

STS are a rare malignancy accounting for 1% of all cancers. Under the WHO 5th edition of classification of soft tissue tumors, ASPS is a rare tumor belonging to the malignant

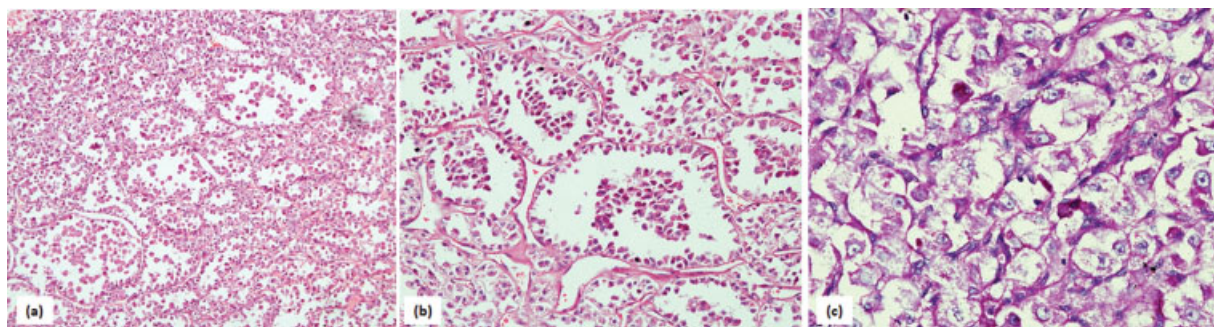


Fig. 1 Histopathology of alveolar soft part sarcoma: (a) Low power picture showing polygonal appearing cells arranged in distinct nesting or organoid pattern with central dyscohesive cells giving appearance of pseudoalveolar arrangement. (b) Cells are generally monomorphic, polygonal shaped with distinct cytoplasmic boundaries. The nucleus is round with finely dispersed chromatin with prominent nucleoli in most of them. Occasional mitotic figures are noted. (c) Diastase treated periodic acid Schiff (PAS) stain highlight rod-shaped intracytoplasmic PAS positive inclusions.

Table 1 Imaging characteristics on CT

Case number	Site	Margin	Size (cm)	Enhancement	Feeders	Necrosis	Ca ²⁺	Local infiltration	Nodes	Mets	Treatment received till last follow-up
1	Leg	Circumscribed	4.2	No	-	-	-	-	-	Lung	Resection with metastatic resection
2	RP	Circumscribed	9.1	Intense	+	-	+	-	-	-	Operated and then LFU
5	Chest wall	Infiltrative	4.7	Intense	+	-	-	Muscle, pleura, ribs	-	Lung	LFU
6	Leg	Circumscribed	9	Intense	+	-	-	-	-	Lung	Stable, On follow-up, no treatment received
7	Chest wall	Infiltrative	11	Intense	+	-	-	Bone	-	Liver	On pazopanib
8	Gluteal	Circumscribed	9	Intense	+	-	-	-	-	Lung	On sunitinib, crizotinib
12	Scalp	Infiltrative	8	Intense	+	-	-	Bone	-	Brain and lung	LFU
15	Leg	Circumscribed	7.7	Intense	+	-	-	-	-	Lungs	VATS with metastatectomy

Abbreviations: CT, computed tomography; Ca²⁺, calcification; LFU, lost to follow-up; mets, metastasis; RP, retroperitoneum; VATS, video-assisted thoracoscopic surgery.

Table 2 Imaging characteristics on MRI

Case number	Site	Margin	Size (cm)	Enhancement	T1 signal	T2 signal	Feeder	Local infiltration	Node	NV bundle involve	Mets	Treatment provided
1	Leg	Circumscribed	4.2	No	Iso	Hyper	-	-	-	-	Lung	WLE + RT metastatectomy
3	Orbit	Circumscribed	3.6	Intense	Hyper	Hyper	+	-	-	-	-	Excision f/b recurrence
4	Leg	Infiltrative	15	Intense	Hypo	Hyper	+	Bone	-	-	Lung, bone	Non resectable; pall CT
6	Leg	Circumscribed	9	Intense	Hetero	Hyper	+	-	-	-	Lung	Stable mets
9	Shoulder	Circumscribed	5	Intense	Hyper	Hyper	+	-	-	-	Lung	WLE Stable mets
10	Leg	Circumscribed	4.6	Intense	Hyper	Hyper	+	-	-	-	Lung	WLE; Stable mets
13	Leg	Circumscribed	7.7	Intense	Iso to intermediate	Hyper	+	-	-	-	Lung, adrenal, bone	Chemo
14	scapula	Infiltrative	5.4	moderate	Iso	Hyper	+	Muscles	-	-	Lung	Partial scapulectomy; stable lung mets
15	Leg	Circumscribed	14.4	Intense	Iso to intermediate	Hyper	+	-	-	-	Lung	Chemo; VATS +metastatectomy
16	Leg	Infiltrative	17	Intense	Iso	Hyper	+	Bone, muscle	+	-	Lung, liver, bone	CT + RT

Abbreviations: f/b, followed by; hetero, heterogeneous echotexture; hyper, hyperintense; hypo, hypointense; Iso, isointense; mets, metastases; NV, neurovascular; pall CT, palliative chemotherapy; RT, radiotherapy; VATS, video-assisted thoracoscopic surgery; WLE, wide local excision.

group of tumors of uncertain differentiation.³ The common occurrence in the skeletal muscle raises speculations of myogenic origin; however, inability to consistently demonstrate IHC markers for the same raises uncertainty of cell of origin. This may be attributed to its origin from myxoid precursors without differentiation.¹³ Histologically, tumor is known to have a homogenous appearance across the mass and between different patients. It has a characteristic appearance of cells arranged in pseudoalveolar or organoid pattern between fibrovascular septae and delicate vessels. In addition, there are intracytoplasmic rod-shaped inclusions which are better highlighted by PAS stain and are diastase resistant. Ultrastructurally, it shows membrane bound rhomboid shaped crystals that are composed of periodic lattice-work of rigid fibrils with dimension of 5 to 7 nm and periodicity of 10 nm. On IHC, the cells are usually negative for epithelial markers like cytokeratins and EMA, neural markers like glial fibrillary acidic protein, and neuroendocrine markers such as synaptophysin. Nuclear reactivity for MyoD1 is variable.¹⁴ In addition, most of the cases showed nuclear immunoreactivity for TFE-3 immunostain. An unbalanced translocation t(X; 17)(p11; q25), which creates the ASPSCR1-TFE3 fusion protein is classically seen in these cases.^{7,8} Rarely this translocation may also be found in some pediatric renal cell carcinomas and periendothelial epithelioid cell tumors.¹⁰ There are reports where histologically renal cell carcinomas mimicked appearance of ASPS due to fascicular arrangement of cells in some areas; however, absence of any renal primary ruled out the diagnosis.¹⁰ Endothelial and periepithelial tumors can be distinguished as these push intact and uninvolved endothelium.⁷

The common presentation of these tumors is as a slowly growing painless mass. Due to their highly vascular nature and feeder vessels, they are often misdiagnosed as arteriovenous malformations (AVMs). Later, these tumors present with distant metastasis due to the diagnostic delay as histopathological examination or surgery is almost never done in the latter. Pennacchioli et al showed that around 9% (3/33) of their patients were initially thought to have vascu-

lar malformations on MRI, hence delaying the diagnosis.⁶ AVMs usually lack any soft tissue component in contrast to sarcomas; they are hypointense on T1 and T2 and show early arterial feeders and early venous washout.^{4,6} Though conventional angiography was not done in any patient in our study, findings such as arterial feeders with prolonged staining of the tumor and late washout indicate soft tissue vascular tumor rather than an AVM.¹⁴

Another commonly confused entity with ASPS can be hemangiomas which are benign vascular tumors. In a review by Viry et al of six children with ASPS, one case of orbital ASPS was initially suspected to be a hemangioma and embolized. However, hemangiomas are known to occur in a much younger age group than ASPS and are usually present since birth. These show characteristic involution with age and can often contain a fatty component on imaging. Also, they enhance in the early arterial phases compared with soft tissue tumors which enhance later in the late arterial or early venous phases.⁴

Small case series or reviews with maximum of 25 cases of ASPS are present in literature due to the rarity of the disease.⁹ Most of the cases of this rare tumor occur in children and young adults. The age group of the cases in our study ranged from 3 years to 72 years with the mean age of 29.25 years of age. The mean age reported in other studies ranged from 18.5 to 36 years.^{9,11} These are seen to occur in a much younger age group than other more common sarcomas like liposarcoma, GIST, and leiomyosarcoma which peak in the fifties.

In contrast to available literature, we had more males with ASPS (six females and 10 males); however, due to small sample size, it would not be correct to say that ASPS has specific gender predilection. It is postulated that due to the characteristic translocation involving X chromosome, ASPS is twice more probable to occur in females.^{4,9,10}

The most common site of occurrence in our study was found to be extremities (10 of 16 cases) which is in concordance with other authors^{4,9-11} (→ **Figs. 2** and **3**). Other tumor locations found in our cohort were retroperitoneum, orbit, chest wall,

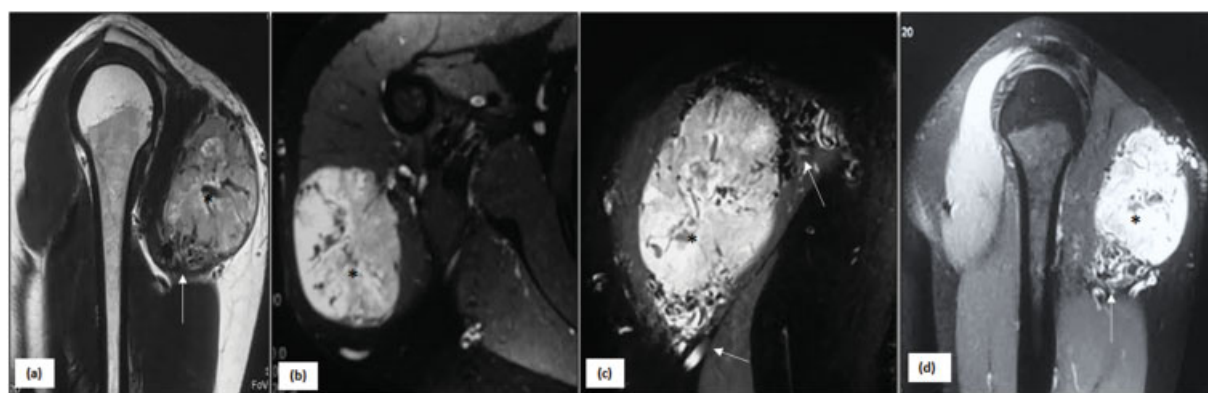


Fig. 2 ASPS of shoulder in a 17-year-old male: Sagittal T1 (a), axial (b), coronal (c), and sagittal (d) T1 PC FS MR images of right shoulder shows a well-defined mass along posterior aspect of shoulder (*asterisk*) infiltrating the deltoid muscle with no infiltration of neurovascular bundle or surrounding bones. The mass is hyperintense on T1 (a) as well as T2-weighted sequences and shows significant homogeneous enhancement (b-d). Multiple flow voids are seen within the mass along with tortuous feeding vessels in vicinity (*arrows*). ASPS, alveolar soft part sarcoma.

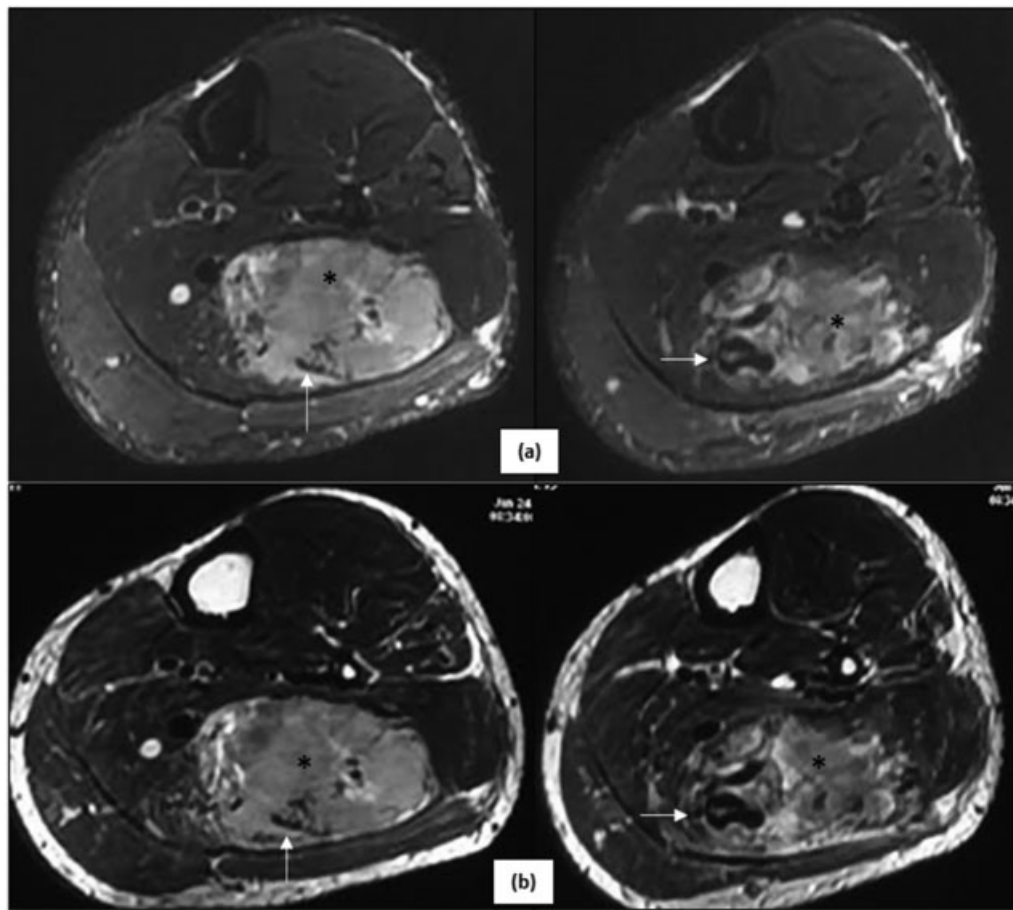


Fig. 3 ASPS in an 18-year-old male patient: Axial T2 FS (a) and axial T2 (b) MR images reveals presence of a well-defined mass (asterisk) in the posterolateral aspect of the left calf appearing hyperintense on T2WI. Multiple tortuous vessels are seen in its vicinity (arrows) and within the mass.

and scalp. Uncommon sites reported by other authors ranged from paraspinal location, tongue, larynx, and buccal space.^{4,13} Head and neck are uncommon sites, however, particularly frequently encountered in children (► Fig. 4).

Ultrasound is often done as a first-time imaging modality to evaluate when a patient presents with a soft tissue swelling which has a wide differential diagnosis ranging from developmental, inflammatory, traumatic, or neoplastic causes. Based on the clinical history and imaging

appearance it is important to exclude an abscess or hematoma. Two of our patients had ultrasound available which showed a well marginated hypoechoic solid mass with multiple large vessels. Daly et al evaluated six patients of ASPS and ultrasound findings revealed a hypoechoic or mixed echotexture mass with well-defined margins and hypervascular nature with large vessels. On spectral analysis, high velocity systolic and diastolic flow was detected.¹⁵ Ultrasound findings are not diagnostic,

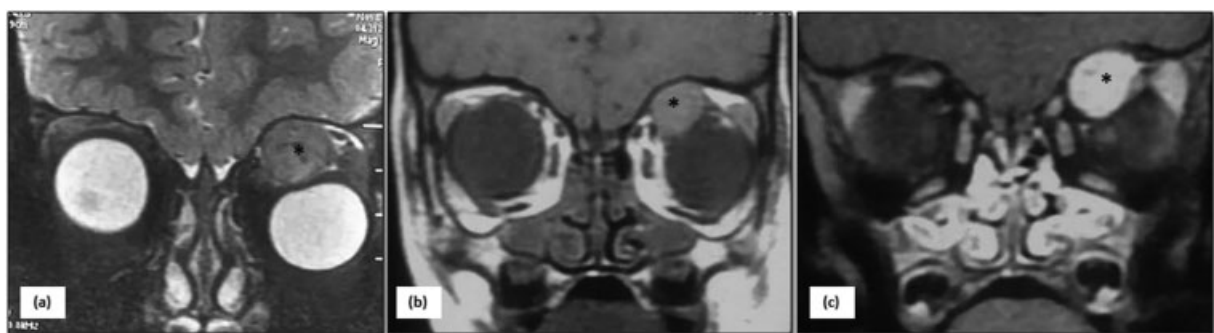


Fig. 4 ASPS of orbit in a 3-year-old boy: coronal T2 (a), T1 (b), and T1 PC (c) MRI of orbits show presence of a well-defined mass in the extraconal compartment of left orbit along the superomedial aspect (asterisk). The mass is hyperintense on T2 (a) as well as T1- (b) weighted sequence and shows homogeneous intense enhancement (c). No intracranial extension was present. ASPS, alveolar soft part sarcoma.



Fig. 5 ASPS of calf with lung metastases in a 29-year-old male patient: plain radiographs AP and lateral views (a) show soft tissue swelling in the region of calf along posterolateral aspect (*asterisk* in a) with permeative lytic changes and cortical irregularity in fibula (*arrow* in a). Subsequently performed MRI selected images-sagittal T2 FS (b), axial T2 (c) showed mass in the posterior aspect of calf (*asterisk* in b, c) with tortuous feeding vessels in the vicinity (*thin arrow* in b). This mass is infiltrating and eroding fibula (c) and is associated with noncontiguous metastatic lesions in tibia (*thick arrow* in b). Axial CT thorax (d) revealed presence of multiple metastatic lesions of variable sizes, diffusely scattered in both lung fields. ASPS, alveolar soft part sarcoma.

however, may raise the suspicion of a soft tissue tumor and prompt further imaging.

On contrast-enhanced CT and MR of the primary mass, most of the cases in our study showed well-defined lobulated margins. Well circumscribed nature of the tumor could be attributed to its indolent slow growing nature. Intense enhancement and feeder vessels were also a commonly seen feature. Intense enhancement, presence of feeders, and multiple flow voids were the most consistent findings across many studies^{4,9-12,16} (→ **Figs. 2, 3, 4, and 5**). Necrosis was not seen, and calcification was only seen in one case. Presence of necrosis is related to tumor size and thus variable results were found in different case series.²

Metastases is common at the time of presentation with lung being the most frequently reported site. This is probably due to the high vascularity and early microvascular dissemination of the tumor.¹⁴ Lung was the most common site with 13 of 14 patients harboring metastatic nodules (→ **Figs. 5 and 6**). This was in keeping with findings reported by other authors.⁴ Other sites of metastatic disease were brain, liver, adrenal, and lymph nodes in our study. Lymph nodes is an unusual site of spread of sarcoma and angiosarcoma is the most common type to harbor nodal metastases (13.5–45%). The same study showed the incidence of nodal spread in ASPS to be 0 to 12.5%.¹⁷ Lymph nodal metastasis was seen in only one of 16 cases in the present study. Overall metastatic disease reported in other studies

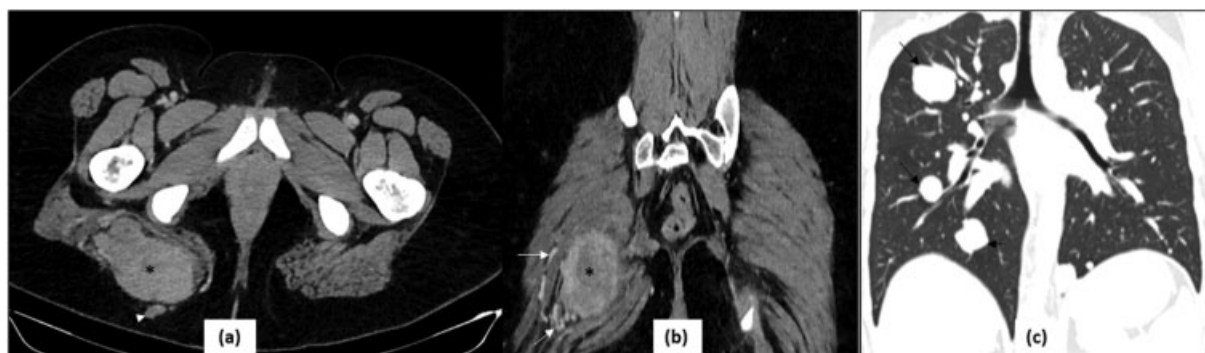


Fig. 6 ASPS in right gluteal region in a 20-year-old girl: Axial (a) and coronal (b) reformatted CT images demonstrate a circumscribed intensely enhancing homogeneous mass (*asterisk*) in the right gluteal region with peripheral feeding vessels (*arrows* in b), infiltrating the gluteal maximus muscle. A smaller satellite lesion is also seen in its vicinity in the subcutaneous plane (*arrowhead* in a). Though the mass appears resectable, the patient could not undergo surgery as she had lung metastases at the time of presentation, as seen on coronal (c) reformatted lung window of chest (*arrows* in c). ASPS, alveolar soft part sarcoma.

Table 3 Summary of imaging characteristics of ASPS seen on different studies

Author	n	Modalities	Primary tumor										Metastasis	
			Age (mean) M:F	Size (mean, in cm)	m/c site	Well circumscribed margin	Flow void/ feeder	Intense Contrast enhancement	Necrosis	Iso-high SI on T1	High SI on T2	Presence	m/c site	
Kim et al, 2005 ¹³ R	5 (only head and neck cases)	CT, MRI, DSA	13.6 2:3	7.2	N.A	5/5	5/5	5/5	—	5/5	5/5	5/5	1/5	N.A
Viry et al, 2013 ⁴ R	6 (children and adolescent 7–17 y)	CT, MRI, USG	Range 7–17 2:4	6	Extremity (4/6)	4/6	6/6	6/6	2/6	4/6	6/6	6/6	5/6	Lungs (5/5)
Sood et al, 2014 ² R	25 5 primary 25 follow-up	CT, MRI, PETCT	Mean-25 14:11	10.2	Extremity (17/25)	3/5	5/5	5/5	3/5	3/3	3/3	23/25	23/25	Lungs (100%)
McCarville et al, 2014 ¹⁰ R	22	CT, MRI	Mean 15 y 8 mo 10:12	5.9	Extremity (16/22)	19/22	11/16	6/16	6/16	14/18	15/17	11/22	11/22	Lungs (11/11)
Li et al, 2014 ¹¹ R	14	MRI	36 9:5	9.8	Extremity (10/14)	0/14	10/14	14/14	—	14/14	14/14	8/14	8/14	—
Tian et al, 2016 ¹⁶ R	14	CT, MRI	29 9:5	9.1	Extremity (6/14)	3/14	12/14	12/14	—	6/9	9/9	13/14	13/14	Lung
Cui et al, 2017 ¹² R	12	MRI	25 5:7	6.8	Extremity (9/12)	—	1/6	12/12	—	—	—	6/12	6/12	Lungs (6/6)
Crombé et al, 2019 ⁹ R	25	MRI	Mean 18.5 9:16	6.6	Extremity (20/25)	0/25	24/25	24/25	10/24	20/25	23/23	14/25	14/25	Lungs (13/14)
Present study, 2020 R	16	USG, CT, MRI, PET-CT	29.2 5:3	8.3	Extremity (10/16)	9/16	15/16	15/16	—	9/10	10/10	14/16	14/16	Lungs (13/14)

Abbreviations: ASPS, alveolar soft part sarcoma; m/c, most common; M:F, male: female; N, subjects included; PET, positron emission tomography; R, retrospective; USG, ultrasonography.

ranged from 50 to 65%^{6,18} as compared with 88% in our study. This can be attributed to the fact that ours is a referral center and thus receives patients in advanced stages mostly. ASPS are indolent slow growing malignant tumors. In spite of this fact, they are usually metastatic at presentation.¹⁹ Metastatic lesions may remain stable for years on continued chemotherapy.²⁰ Pulmonary metastatectomy is the preferred treatment in the absence of metastases at other sites.²¹ ► **Table 3** summarizes the imaging features of ASPS seen in different studies.

It is important here to discuss the various differential diagnosis of an avidly enhancing mass with feeder vessels. The possibility of a hypervascular metastasis from a renal or thyroid primary should always be kept in mind. Also, angiomatous tumors like angiosarcoma, hemangiopericytoma could also have a similar imaging appearance.¹⁴ However, these angiomatous tumors often show blood fluid levels on MRI which are not described in ASPS.²²

Retrospective study design with small sample size of 16 patients due to rare nature of this sarcoma is one of the limitations of our study. Also, the imaging protocols were not standardized due to most imaging being done from outside hospitals and referred to our tertiary care center for review and further management. The imaging and histopathology were reviewed by single radiologist and pathologist from multidisciplinary sarcoma team; thus, interobserver variability was not evaluated in this study. Also, as the cohort of cases belonged to a tertiary care center, more patients with metastatic disease were seen.

In conclusion, ASPS is a rare STS seen in children and young adults having an indolent course. It should be suspected in cases showing an intensely enhancing intramuscular mass with feeder vessels. Imaging may mimic a vascular malformation or tumor due to large number of feeders. Misdiagnosis at an early stage may lead to later metastatic presentation of the disease.

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