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·病例报道·

Contrast-enhanced ultrasonic manifestations of anterior mediastinal extraskelatal osteosarcoma: a case report 前纵隔骨外骨肉瘤超声造影表现 1 例

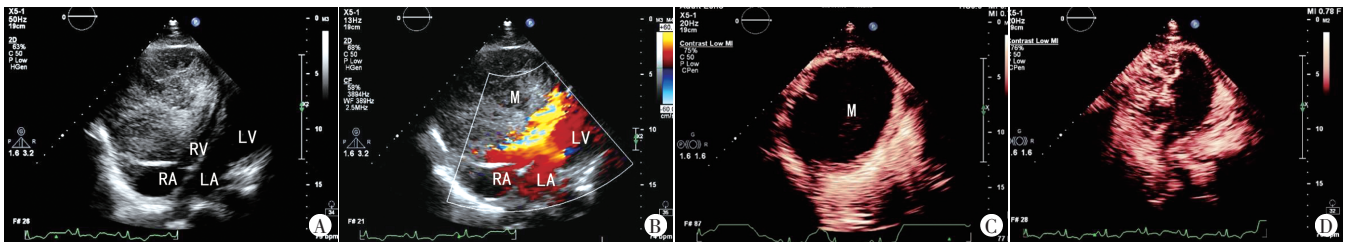
严定芳 谢超 贺赞 张文君

[中图法分类号]R445.1

[文献标识码]B

患者男,71岁,纵隔肉瘤术后11个月余,因“呼吸困难”就诊。胸部平扫+增强CT检查:前下纵隔肉瘤术区见一巨大肿块影,边界不清,较大层面大小约11.0 cm×10.0 cm×9.4 cm,内密度不均,明显不均匀强化,边缘强化为主;心包少量积液,双侧胸腔少量积液。CT提示:前下纵隔肉瘤术后复发。经胸超声心动图检查:前纵隔见一大约11.7 cm×11.0 cm×9.8 cm不均质肿块(图1A),边界不清,形态规则,内部回声不均匀,大部分呈低弱回声区,右心明显受压,右心腔狭小;CDFI于肿块内未探及明显血流信号(图1B)。结合病史超声提示:前纵隔恶性肿瘤术后复发可能。为进一步明确肿块性质及其与心脏关系,

行左心声学造影检查:纵隔见一巨大肿块,边界不清,形态规则,内呈不均匀性增强,以边缘增强为主,中心见大片状无增强区(图1C、D)。左心声学造影提示:前下纵隔肉瘤术后复发,肿瘤坏死区较大。后于全麻下行纵隔肿瘤切除术,术中见:肿块巨大,范围由右室流出道水平至膈肌水平,双侧胸膜至膈神经水平均累及,与心脏界限不清(图2A),无法判定心肌是否受侵,切开右室面上方肿块,其内见大片出血坏死(图2B)。术后组织病理学检查:富于巨细胞的肿瘤组织伴大片坏死,可见脉管内瘤栓(图2C)。病理诊断:纵隔高级别肉瘤伴大片坏死,考虑骨肉瘤,可见脉管内瘤栓。免疫组化检查:CD34(-),Ki-67(70%+),



A:二维超声示前纵隔巨大不均质肿块,边界不清,形态规则,内大部分呈低弱回声,右心腔明显受压;B:CDFI示前纵隔肿块内未探及明显血流信号;C、D:左心声学造影示前纵隔肿块边缘成分不均匀性增强,中心大片无增强

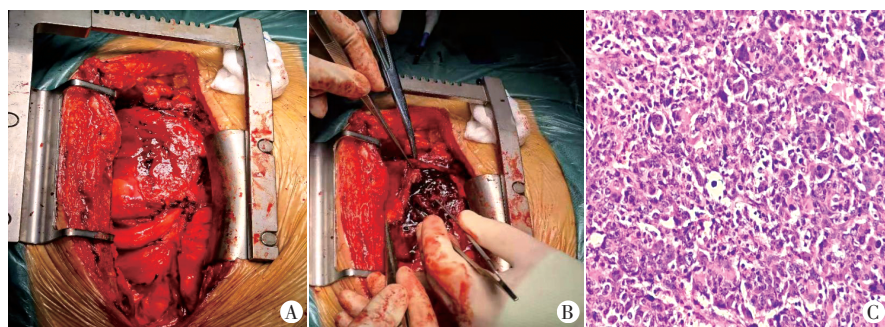
图1 前纵隔骨外骨肉瘤常规超声及超声造影图(RV:右室;LV:左室;RA:右房;LA:左房;M:肿块)

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A: 术中见肿块巨大, 双侧胸膜至膈神经水平均累及, 与心脏界限不清; B: 切开右室面上方肿块, 其内见大片出血坏死; C: 病理图显示富于巨细胞的肿瘤组织伴大片坏死, 可见脉管内瘤栓 (HE 染色, $\times 40$)

图2 前纵膈骨外骨肉瘤术中大体观及病理图

STAT6(-), SATB2(+), CD68(多核巨细胞+), CD163(部分+), SMA(部分+), CK(P,-)。患者术后常规行抗感染治疗, 未行放、化疗, 随访半年未见肿瘤复发和远处转移。

讨论: 原发性纵膈骨外骨肉瘤 (extramedullary osteosarcoma, ESOS) 是一种罕见的起源于间充质细胞并向骨、类骨及软骨分化的软组织恶性肿瘤, 约占所有软组织肉瘤的 1%~2%, 占所有骨肉瘤的 2%~4%^[1-2]。纵膈 ESOS 平均发病年龄约 50 岁^[3], 发病机制目前尚未明确, 可能与肺部并发非结核分枝杆菌感染有关^[4]。前纵膈 ESOS 患者早期常无阳性临床体征, 多因进行性增大的软组织肿块伴胸闷、咳嗽、呼吸困难而就诊。本例纵膈骨肉瘤术后 11 个月因呼吸困难发现复发。本文首次报道前纵膈 ESOS 的超声及超声造影表现, 二维超声示肿块形态规则, 边界不清, 回声不均匀; 超声造影表现为肿块内部呈明显不均匀增强, 周边组织实质性成分明显增强, 增强程度不低于邻近心肌且与邻近心肌分界不清, 中心见大片状无增强区, 提示肿瘤为多血管肿瘤并中心大片状液化坏死可能。本例手术病理诊断为 ESOS, 为高度恶性肿瘤, 与超声造影表现相符。

纵膈 ESOS 容易局部复发及发生远处转移, 最常见的转移

部位为肺及邻近软组织, 患者 5 年生存率 24%~46%, 合并转移的纵膈 ESOS 患者平均生存期仅为 8 个月^[5]。本例前纵膈 ESOS 在第一次手术 11 个月后再次复发, 且术中探查发现双侧胸膜至膈神经水平均受侵犯, 具有高度恶性的生物学行为。

总之, 虽然纵膈 ESOS 超声表现尚无特征性, 但常规超声检查方便、快捷, 且超声造影能实时反映肿块组织内血供分布情况, 仍可作为评估其血供及远期疗效随访的重要影像学手段; 当纵膈内出现非转移性的软组织肿块, 图像上可见点状或片状钙化斑, 且肿块未与周围骨骼相连时需警惕纵膈 ESOS 的可能。

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