Case Report

Dedifferentiated Liposarcoma of the Retroperitoneum with Extensive Leiomyosarcomatous Differentiation and β-Human Chorionic Gonadotropin Production

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Dedifferentiated liposarcomas may display a variety of “heterologous” lines of differentiation, including osseous, vascular, skeletal, and/or smooth muscular. There have been six previously reported examples of leiomyosarcomas associated with high levels of serum human chorionic gonadotropin (hCG) production, comprised of cases originating from the retroperitoneum, spermatic cord, small intestine, and uterus. This report describes the first example of a dedifferentiated liposarcoma that combined both of the aforementioned features: extensive heterologous (leiomyosarcomatous) differentiation and β-hCG production (maximum serum levels 1046 mIU/ml, reference < 5 mIU/ml). The tumor, which originated in the retroperitoneum in the region of the right kidney, was rapidly progressive and ultimately fatal within three months of its diagnosis. In addition to characteristic morphologic features, lipogenic and smooth muscle differentiation were confirmed with immunohistochemical stains for MDM2 and smooth muscle actin, respectively. The tumor also displayed diffuse immunoreactivity for β-hCG in both primary and metastatic sites. This case further expands the clinicopathologic spectrum of lipogenic tumors.

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1. INTRODUCTION

The term “tumor dedifferentiation” and its underlying concepts, as used by contemporary surgical pathologists, are based on the paradigm originally outlined by Dahlin and Beabout in low-grade chondrosarcomas [1]. Tumor dedifferentiation is characterized by the “emergence” of an undifferentiated component (phenotypically unrecognized as to histogenesis) from an otherwise low-grade/borderline neoplasm, with well-defined morphologic demarcations between the aforementioned areas and a relatively worsened clinical course. The phenomenon of tumor dedifferentiation has subsequently been described in a wide variety of bone, soft tissue, and epithelial and neural neoplasms [2].

Tumor dedifferentiation is probably most well characterized in lipogenic tumors. Although it had been previously noted that well-differentiated and pleomorphic sarcomas may be admixed [3, 4], dedifferentiated liposarcoma (DDLs) was formally described by Evans in 1979 [5]. In a series of 55 liposarcomas, 8 cases (7 retroperitoneal, 1 spermatic cord) were noted to display “distinct areas of well-differentiated liposarcoma and of cellular spindle cell or pleomorphic sarcoma without recognizable lipogenesis” [5]. None of the latter cases displayed a mitotic count of >5/10 high power fields, and the cellular areas were often sharply demarcated from the well-differentiated areas [5]. Subsequent reports indicated that (a) dedifferentiation in well-DDLS is probably not anatomic site specific but time dependent [6], (b) most retroperitoneal tumors previously classified as malignant fibrous histiocytomas are actually DDLS [7], (c) most examples of DDLS present as de novo lesions rather than as a complication of a preexisting well-differentiated liposarcoma [8], (d) DDLS may be classified into high and low grade depending on mitotic activity and cellularity, although
the prognostic significance of this gradation is unclear [8–11].

As with their clinical spectrum, the pathologic spectrum of DDLs has been similarly expanded. Newer morphologic observations have included a potentially poorly defined interface between the DDLs and its well-differentiated component [8], whorled, meningothelial-like structures [8, 12–14], a prominent inflammatory component [9], a micronodular growth pattern [9], a hemangiopericytomatic vascular pattern [8, 9], and heterologous differentiation [8, 9, 14–20].

Chondroid metaplasia, rhabdomyosarcomatous, and smooth muscular differentiation have been described in well-differentiated and myxoid liposarcomas [2, 21–24]; the term lipoleiomyosarcoma is commonly used for morphologic ad-mixtures of well-differentiated liposarcoma and leiomyosarcoma. [23]. However, the probability for histologic misclassi-fication is highest when high-grade DDLs displays heterolo-gous differentiation [4, 5, 8, 9, 11–16]. Lines of heterolo-gous differentiation that have been described in DDLs in-clude leiomyosarcomatous, osteochondrosarcomatous, angiosarcomatous, and rhabdomyosarcomatous and do not ap-pear to have independent prognostic significance [4, 5, 8, 9, 11–16, 25].

This report describes an example of a retroperitoneal DDLs with extensive leiomyosarcomatous differentiation that was accompanied by high levels of serum beta-human chorionic gonadotropin (β-hCG). This composite of find-ings has, to our knowledge, not been previously reported.

2. CLINICAL HISTORY

In 2006, a 67-year-old Caucasian female presented with episodic right upper quadrant abdominal pain and associ-ated nausea. Her past medical history was significant for an infiltrating ductal breast carcinoma (treated with rad-i-cal mastectomy followed by 4 cycles of adriamycin in 1999), a stable plasma cell dyscrasia (multiple myeloma diagnosed by a bone marrow biopsy and serum protein electropho-sis in 2003, treated with high dose chemotherapy, and which was associated with a mild normocytic anemia in the ensu-ing period), and a hysterecomy/salpingo-oophoretomy (for benign indications). A computed tomographic (CT) scan of the abdomen showed a heterogeneously enhancing mass extending from the inferior right renal pole exophyti-cally, with significant associated inflammatory changes and some peripheral coarse calcifications. Portocaval and mesen-teric adenopathy were also present. Serum levels of alpha-fetoprotein, carcinoembryonic antigen, CA15-3, and CA27-29 were within normal limits. Notably, a serum β-hCG, requested inadvertently, measured 210.3 mIU/ml (reference value for nongestational adults <5 mIU/L).

Given the high suspicion for a renal cell carcinoma, the right kidney was resected as well as all grossly identifiable perinephric tumor. Intraoperatively, the mass was noted to be prominently adhered to the surrounding retroperitoneal tissue. A follow-up serum β-hCG, which was measured 26 days after the first measurement and 25 days after her sur-gical procedure, measured 1046 mIU/ml. An abdominal CT scan performed shortly thereafter revealed multiple hypo-dense liver masses measuring up to 5 cm, with an increase in the aforementioned portocaval adenopathy. Biopsy of one of the hepatic masses confirmed the presence of metastatic dis-ease. The patient declined further therapeutic measures and expired 2 months later. An autopsy was not performed.

3. MATERIALS AND METHODS

Twenty sections of the retroperitoneal mass were routinely processed for microscopic evaluation. The liver biopsy was similarly processed. For immunohistochemistry, 4–5 μ-thick sections were cut and mounted on a glass slide, depara-finized and rehydrated. Appropriate negative and positive controls were included. All assays were carried out in an Axiom 36 autostainer (Lab Vision Corporation, Fremont, Calif, USA). The following markers were evaluated in the liver biopsy and a representative section of the retroperitoneal mass, in parallel: β-hCG (clone CG04+CG05, predi-luted, Lab Vision), keratin cocktail comprised of clones AE1/AE3 (Signet Corporation, Dedham, Mass, USA) and LP34 (DakoCytomation, Carpinteria, Calif, USA), dilution 1 : 200, Smooth muscle actin: SMA (clone1A4, prediluted, Lab Vision), Estrogen receptor-alpha: ER-alpha (clone ID5, dilution 1 : 50, DakoCytomation), Progesterone receptor: PR (clone PgR 636, prediluted, DakoCytomation), Desmin (clone D33, prediluted, Lab Vision), Epithelial membrane antigen (EMA) (clone E29, prediluted, ThermoFisher Sci-entific, Fremont, Calif, USA), MDM2 (clone IF2, dilution 1 : 100, Lab Vision), and placental-like alkaline phosphatase: PLAP (Clone SP15, prediluted, Lab Vision).

4. PATHOLOGIC FEATURES

4.1. Macroscopic

The kidney and associated perinephric tissue measured in ag-gregate 21 × 10 × 7 cm. Sectioning revealed an inferior pole mass measuring 9.5 cm in maximum dimension, which com-pressed and distorted the renal pelvis and the tissue in this re-gion. The cut surface of the mass was largely firm and solid, tan-white in color, with hemorrhagic areas but no grossly identifiable necrosis.

4.2. Morphologic features

The retroperitoneal tumor displayed the characteristic mor-phologic features of DDLs. In the well-differentiated com-ponent, large coarse fibrotic bands were observed irregularly, coursing through lobules of well-differentiated adipocytes. Within the fibrous bands, the perivascular regions and within the adipocytic lobules were scattered cells with large irregu-lar and hyperchromatic nuclei (Figure 1). The mitotic index in these regions was low. The dedifferentiated component comprised approximately 80% of the tumor mass. The inter-face between the well-differentiated and dedifferentiated components was variable: the interface was well demarcated in some regions (Figure 2), and highly irregular/infiltrative in others. The central portion of the mass was necrotic, while the surrounding areas, comprising approximately 10% of the mass were hypocellular and hyalinized. However, atypical
Michael J. Russell et al.

4.3. Immunophenotypic features

The retroperitoneal tumor and liver mass displayed an identical immunoprofile, consistent with the latter being a metastatic deposit from the former. Both displayed diffuse immunoreactivity for desmin, SMA, β-hCG (Figure 5), and MDM2 (Figure 6). The tumors were negative for ER, PR, EMA, PLAP, and keratins.

5. DISCUSSION

hCG, a glycoprotein with a molecular weight of 38,000, is comprised of two subunits—α and β—that are noncovalently bound to form a heterodimer [26, 27]. The α subunit of hCG is identical to the α subunits of luteining hormone, follicle stimulating hormone, and thyrotropin; the β subunit therefore provides hormonal specificity [26, 27]. The measurement of serum β-hCG has its greatest clinical utility in the diagnosis of gestation, congenital disorders, and germ cell neoplasia. However, it has long been recognized that a wide variety of other nongerm cell neoplasms, predominantly carcinomas, may be associated with elevated serum β-hCG, in which it appears to function as an autocrine growth factor by inhibiting apoptosis [28, 29]. The expression of hCG has also been found to be an adverse prognostic factor in some specific histotypes such as small cell lung carcinoma,
transitional cell carcinoma of the urinary bladder, and prostatic and colorectal adenocarcinoma [30–33]. Rare examples of osteosarcoma [28, 34–36], chondrosarcoma [37], and leiomyosarcoma [38–43] have been associated with elevated serum β-hCG. Amongst the six previously reported examples of β-hCG-producing leiomyosarcomas, two originated in the retroperitoneum [42, 43], two in the spermatic cord [39, 41], one in the uterus [40], and one in the small intestine [38]. Notably, five of the six patients succumbed to their disease within one year of their diagnoses. However, the sarcomas were high grade; thus the independent prognostic significance of β-hCG remains unclear.

In the present case, a DDLS with extensive leiomyosarcomatous differentiation was associated with β-hCG production, as evidenced by demonstration of β-hCG expression in the tumor and its metastatic deposit by immunohistochemistry, and an elevation of serum β-hCG that was temporally related to tumor progression. The diagnostic utility of MDM2, a recently described marker reportedly useful for distinguishing DDLS from their histologic mimics [44], was confirmed, as this marker helped to establish, in addition to morphologic features, underlying lipogenic differentiation.

β-hCG production by DDLS has not been previously reported to our knowledge and is presumed to represent “aberrant” differentiation in a high-grade tumor. Nonetheless, this finding expands the clinicopathologic spectrum of lipogenic tumors in general. Furthermore, it suggests that liposarcoma should be a differential consideration when elevated serum β-hCG is associated with a radiographic mass lesion, especially when the mass is retroperitoneal.

REFERENCES


