Carcinosarcoma of Gallbladder with Metastases in Liver and a Lymph Node

Authors
AK Kapoor, Akanksha Singh, Pratibha Singh
RML Mehrotra Pathology Pvt. Ltd., Nirala Nagar, Lucknow, India
Corresponding Author
Ashok Kumar Kapoor
RML Mehrotra Pathology Pvt Limited, B 171, Nirala Nagar, Lucknow, India
Email: drashokkapoor2016@gmail.com

Abstract
Carcinosarcomas are highly malignant neoplasms where pleomorphic epithelial cells comingle with areas showing mesenchymal differentiation. Herewith, we describe the clinicopathological features of a case of carcinosarcoma gallbladder (CSGB). The patient was a female, aged 60 years. Immunohistochemical (IHC) examination revealed strong positivity of adenocarcinomatous tissue for cytokeratin while sarcomatous tissue showed vimentin antigen positivity. Findings of current case suggested differentiation of pluripotent tumor cells both as epithelial and mesenchymal components. She was diagnosed as a case of CSGB metastasizing in liver and a regional lymph node.

Keywords: Epithelial mesenchymal differentiation in a malignant neoplasm.

Introduction
Clinically, the patients with this tumor may present as pain in right hypochondrium. Rarely, pain may be severe and may be associated with vomiting. The patient may present as a case of acute pancreatitis. Few patients may also develop obstructive jaundice due to a stone in common bile duct. Tumor may locally invade surrounding organs, e.g. liver, stomach and transverse colon. Tumor may arise from a single pluripotent stem cell which later differentiates into epithelial and mesenchymal cells. Herewith, we report a case of carcinosarcoma gallbladder (CSGB) with metastases both in liver and a lymph node.

Case Report
A female, aged 60 years complained of recurrent episodes of pain in right hypochondrium and vomiting for the last 1 year. She also had tenderness in right side of upper abdomen. Liver function tests revealed serum total bilirubin 0.8 mg/dl. SGPT was 60 U/L and S. alkaline phosphatase was 120 U/L. S. albumin was 3.6 gm/dl and S. globulin was 2.4 gm/dl. SGOT was 53 U/L. Later, cholecystectomy was done. Gallbladder measured 10×5×4 cms. It contained multiple sludge-balls. Outer surface was smooth. Cut surface of gallbladder showed an irregular tumor reaching the neck of gallbladder. Tumor measured 7×3×1 cms. Sections were taken both from the neck of the gallbladder and tumor tissue.
Another piece consisted of excised wedge-shaped liver tissue, measuring 3×2.5×2 cms. Sections were taken. Microscopic examination showed tumor tissue. Tumor consisted of proliferated cuboidal or columnar epithelial cells forming irregular papillary structures or acini (figure 1a). Tumor cells had ovoid hyperchromatic nuclei and light-stained cytoplasm. Tumor cells showed pleomorphism. Tumor necrosis was seen. In addition, tumor showed areas of chondrosarcomatous differentiation (figure 1b). Lymphocytic infiltration and fibrosis were seen. Section from the neck of gallbladder showed structure of a lymph node showing metastatic tumor with capsular invasion (figure 1c). Section from liver also showed tumor tissue (figure 1d). At places, liver parenchyma showed pools of mucin containing tumor cells. Metastatic tumors showed both adenocarcinomatous and sarcomatous tissue. Immunohistochemical examination of tumor tissue was done using monoclonal anti-cytokeratin and anti-vimentin antibodies. Adenocarcinomatous tissue showed strong positivity for cytokeratin (figure 1e) and sarcomatous tissue showed vimentin antigen positivity in 60% of tumor cells (figure 1f). Tumor was diagnosed as carcinosarcoma gallbladder (CSGB) with metastases both in liver and a lymph node.

**Figure-1**

![Figure 1](image_url)

- a) Shows irregular acini and papillary structures (HE × 100).
- b) Shows chondrosarcomatous component of tumor (HE × 100).
- c) Metastatic tumor lymph node (HE × 100).
- d) Metastatic tumor liver (HE × 100).
- e) Epithelial component of tumor shows strong positivity for cytokeratin antigen (× 100).
- f) Mesenchymal component shows strong positivity for vimentin (× 100).
Discussion
Most important feature of the current case was the detection of metastatic tumors both in liver and a regional lymph node. Invasion of a venule might have resulted in metastatic tumor in liver through portal vein. Spread of tumor to a regional lymph node might have occurred through invasion of an afferent lymphatic. Another interesting feature was the morphology of tumor in metastases; tumor showed carcinosarcomatous tissue both in liver and in the regional lymph node. In another study, a sarcomatous thrombus alone was detected in portal vein1. Lymphatic spread of CSGB is rare but occasionally it may occur2. Current case of CSGB was a female. Most of the cases of this tumor are females. In addition, ovariectomized estrogen-free LXRβ⁻/⁻ mice failed to develop preneoplastic lesions3. These observations suggest role of estrogen in gallbladder carcinogenesis. Another interesting feature was the detection of multiple sludge-balls in our case. Sludge-balls are known to be the precursors of gall stones; these structures develop due to bile stasis. Role of bile stasis has also been suggested in an earlier study4. Pancreatico-biliary reflux due to maljunction may also be associated with biliary carcinogenesis. Several risk factors may be involved in gallbladder carcinogenesis, e.g. chronic inflammation by Helicobacter species or Salmonella typhi5,6. In addition, p53 gene mutations may trigger gallbladder carcinogenesis7. Role of survivin, an apoptosis-inhibiting protein has been suspected in gallbladder carcinogenesis. Furthermore, increased survivin expression was observed in poorly differentiated malignant tumor when compared with well differentiated gallbladder carcinoma8. Most of the patients with CSGB die within 6 months after surgery. The patients with tumors <5 cm had longer survival as compared to those with bigger tumors9. Targeted therapies using anti-angiogenic and anti HER 2/neu agents may be beneficial to CSGB patients7.

Conclusion
Carcinosarcoma is a rare malignancy of gallbladder. Exact etiopathogenesis of this tumor is not known. A 60 years old female had a gallbladder tumor. Tumor showed the features of both adenocarcinoma and chondrosarcoma. Moreover, carcinosarcomatous tissue metastasized to liver as well as to a regional lymph node. In spite of surgery, prognosis of carcinosarcoma is poor. However, longer survival may occur in a few patients.

Financial support and sponsorship: Nil
Conflicts of interest: There are no conflicts of interest.

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