Case Report

Neuroendocrine Small-Cell Carcinoma of the Gallbladder: Case Report

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ARTICLE INFO

Article history:
Received: 19 December, 2019
Accepted: 1 January, 2020
Published: 11 January, 2020

Keywords:
Gallbladder tumors
neuroendocrine tumors
small-Cell neuroendocrine carcinoma

ABSTRACT

Gallbladder neuroendocrine carcinoma is a very rare entity, accounting for <0.2% of all neuroendocrine tumours [1]. Of the cases described in the literature, the initial presentation is usually locally advanced or even with disseminated disease [2]. In this paper we report the case of a 43-year-old female patient with an incidental diagnosis of gallbladder neuroendocrine carcinoma. Although clinical presentation is the most common, it is a very rare condition that poses a therapeutic and prognostic challenge.

Introduction

Primary Neuroendocrine Tumors (NET) of the gallbladder are rare, and while well-differentiated NETs have an indolent course and overall good prognosis, poorly differentiated neuroendocrine carcinomas (NEC) have a poor outcome [3]. Histological types of gallbladder NET can be divided in well-differentiated carcinoid tumors, small-cell carcinomas (grade 1) and large-cell neuroendocrine carcinomas (grade 2) and mixed adeno-neuroendocrine carcinomas (MANEC) [3, 4]. Of these types, carcinoid tumors have a more favorable course where small-cell neuroendocrine and undifferentiated carcinomas have a poor prognosis [5-7]. In this case report we pretend to illustrate the presentation, diagnosis, evolution and outcome of a small-cell NEC of the gallbladder and the struggle to optimize the treatment to this aggressive form of NET.

Case Presentation

A 43-year-old female presented to the emergency department with abdominal pain, localized in the right upper quadrant for the last 24 hours.

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T3N1Mx. The patient underwent a thoraco-abdomino-pelvic CT scan, which revealed a hepatic lesion with 8cm-diameter in segment 4 with arterial enhancement, consistent with a secondary location. No other suspicious lesions where identified (Figure 1).

**Figure 1**: Initial assessment after the histopathology diagnosis, showcasing liver metastasis.

An FDG-PET was also performed and revealed a hepatic secondary location in segment 4, corresponding to the same site on the CT-scan, and metastatic lymphatic invasion involving the hepatic hilum. The patient was proposed for systemic chemotherapy. Besides QT, disease progression occurred and reevaluation with FDG-PET after 6 weeks of QT showed a progressing liver metastasis and the appearance of multiple bone metastasis, with the patient dying at 9 months after the initial diagnosis. (Figure 2).

**Figure 2**: Post QT assessment that reveals the progressing of liver infiltration.

**Discussion**

This case illustrates the rapid evolution of the NEC of the gallbladder, with a poor prognosis, short-term survival and little to no response to chemotherapy. Gallbladder carcinomas prognosis is dependent not only on local staging but also of the histological type – so adenocarcinomas and other well-differentiated tumors have a better outcome than small-cell neuroendocrine and other poorly differentiated tumors. The literature compares most NET of the gallbladder behavior to gallbladder adenocarcinoma, especially the poorly differentiated ones, poor prognosis, usually with late-stage presentation. In the specific case of small-cell neuroendocrine carcinoma, the few cases described in the literature – the behavior of these tumors seems to be even more aggressive than the gallbladder adenocarcinoma with shorter survival after diagnosis and even worst response to chemotherapy. So, although surgical radical resection is the paramount treatment it is rarely doable or of any clinical benefit at all.

**Conclusion**

Gallbladder neuroendocrine carcinomas are initially at an advanced stage and do not respond to chemotherapy regimens. Rarely with the possibility of radical surgical therapy and poor prognosis. The median survival is 4-6 months despite the therapeutic measures instituted [8]. Nevertheless, a recognition of a neuroendocrine type carcinoma is important to evaluate prognosis and also to prevent and treat other paraneoplastic syndromes as well as a possible different chemotherapy scheme.

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