

Aggressive clinical course of large cell neuroendocrine carcinoma of ampulla of vater

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ABSTRACT

A 78-year-old male patient with history of right hemicolectomy due to adenocarcinoma was admitted by the complaint of epigastric discomfort. Laboratory data showed increase in liver biochemistries (aspartate aminotransferase (AST): 159 IU/L, alanine aminotransferase (ALT): 235 IU/L, alkaline phosphatase (ALP): 350 IU/L, gamma glutamyl transferase (GGT): 911 IU/L, total bilirubin: 1.55 mg/dl and direct bilirubin: 0.82 mg/dl). Endoscopic retrograde cholangiopancreatography (ERCP) done after the gastrointestinal (GI) upper endoscopy was compatible with tumoral lesion, and biopsy confirmed 'neuroendocrine carcinoma'. Pylorus-preserving pancreaticoduodenectomy (PPPD) had been performed with R0 resection. Pathologic evaluation revealed 1.5 cm tumor of large cell neuroendocrine carcinoma (LCNEC). Five months later, biopsy of suspicious lesions in liver had been documented as 'high grade neuroendocrine carcinoma metastasis'. He was referred to oncology for chemotherapy but unfortunately he had expired three months later. Large cell neuroendocrine carcinoma (LCNECs) of ampulla of Vater may have aggressive clinical course despite radical resections involving lymph node dissections. Small tumor size and lymph node negativity are not reliable factors for this tumor type.

Keywords: Ampulla of vater; large cell type; neuroendocrine tumor.

Cite this article as: Sonmez RE, Buyuker F, Girgin RB, Eren TT, Bas G, Alimoglu O. Aggressive clinical course of large cell neuroendocrine carcinoma of ampulla of vater. *North Clin Istanbul*

Neuroendocrine tumors (NETs) of ampulla of Vater are very rare neoplasms of digestive tract with a prevalence less than 5% of all duodenal malignancies, and less than 2% of gastrointestinal neuroendocrine tumors [1]. In 2010, (WHO) World Health Organization diagnostic criteria classified NETs as; NET G1 (Low grade), G2 (Intermediate grade), and G3 (High grade or neuroendocrine carcinoma (NEC)), and mixed adenoneuroendocrine carcinoma [MANEC] [2].

Neuroendocrine tumors are yellowish-grayish in color with a nodular or polypoid appearance. G1 and G2 NETs are mostly located in the mucosa or submucosa, whereas G3 NETs (NECs) appear as ulcerated masses

invading deeper layers of GI system. NECs are histologically classified as LCNEC (Large cell neuroendocrine carcinoma) and SCNEC (Small cell neuroendocrine carcinoma). Large-cell neuroendocrine carcinomas are composed of medium-large cell sized tumor cells, having nuclear to cytoplasmic ratio lower than that of small-cell neuroendocrine carcinomas [3].

As a general review; NECs having tumor size greater than 2 cm, muscularis propria invasion with/or involvement of lymph nodes have a tendency to make distant metastasis. Surgery holds the mainstay in treatment for these tumors. Also, some authors mention the reliability of more conservative treatment (endoscopic interven-



Received: January 23, 2019 Accepted: May 14, 2019 Online: July 11, 2019

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tions) for the management of early stage NECs of ampulla of Vater.

There are only few studies discussing the clinical course of LCNECs in literature [4]. This case study is about a 78-year-old male patient diagnosed with LCNEC of ampulla of Vater.

CASE REPORT

A 78-year-old man had been presented with jaundice and discomfort on the epigastric region of abdomen. He had a history of right-hemicolectomy three years ago due to colonic adenocarcinoma, and had completed his adjuvant chemotherapy with six cures of capecitabine plus oxaliplatin. On his admission, laboratory data showed an increase in his liver biochemistries (aspartate aminotransferase (AST): 159 IU/L (normal values <35), alanine aminotransferase (ALT): 235 IU/L (normal values <41), alkaline phosphatase (ALP): 350 IU/L (normal values <129), gamma glutamyl transferase (GGT): 911 IU/L (normal values <55), total bilirubin: 1.55 mg/dl (normal values <1.0) and direct bilirubin: 0.82 mg/dl (normal values <0.2). Intra-extra hepatic biliary ducts

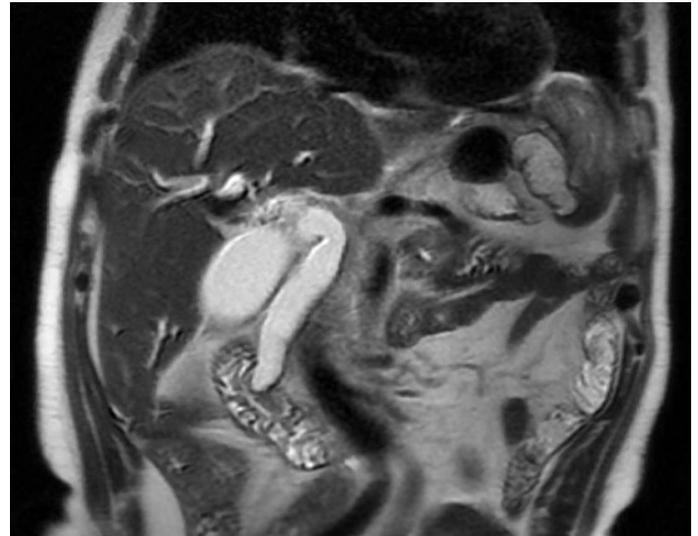


FIGURE 1. Blunt ending of common bile duct with 1 cm nodular lesion on periampullar region.

were dilated, and distal common bile duct had a blunt ending appearance on hepatobiliary ultrasonography (US), and 1 cm nodular lesion was observed on periampullary region by magnetic resonance cholangiopancreatography (MRCP) evaluation (Fig. 1).

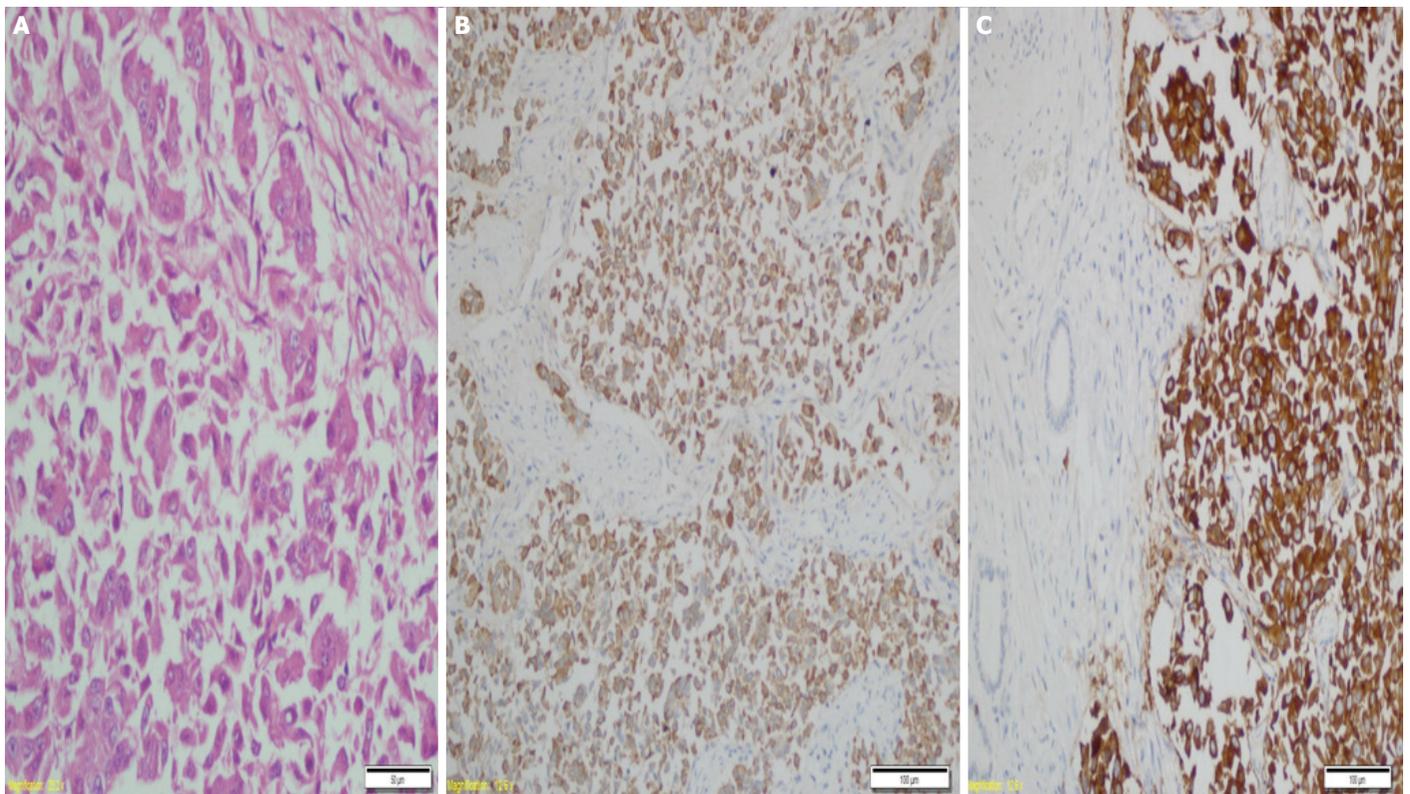


FIGURE 2. (A) Tumor cells with extensive eosinophilic cytoplasm, some with prominent nucleoli (H&E, x400). (B, C) Immunohistochemical chromogranin A and synaptophysin expression in tumor cells (chromogranin A, x200; synaptophysin, x200).

He was hospitalized to our General Surgery department for further assessment. On his endoscopic retrograde cholangio pancreatography (ERCP), there was 'bulging' tumor appearance invading orifice of papilla. Preoperative biopsy of the suspected lesion was 'neuroendocrine tumor' (NET).

Pylorus-preserving pancreaticoduodenectomy (PPPD) with regional lymph node dissection was performed. Pathologic diagnosis of the specimen was LCNEC (G3) (Fig. 2). Ki-67 labeling index was 80%. Surgical margins were clear of tumor cells. Size of the tumor was 1.5 cm, and a total of 17 lymph nodes were dissected. And none of the lymph nodes were found positive.

The patient didn't receive adjuvant therapy. Five months after the operation, he was screened by IV contrast-enhanced computerized tomography (CT) due to increase in enzymes showing liver biochemistry. Multiple metastatic lesions located on both lobes of the liver, with the biggest one approaching nearly 4 cm in segment 8 was found on CT scan. Biopsy of one of the lesions was reported as 'neurocarcinoma metastasis'. He was referred to oncology department to receive chemotherapy. But the disease had an aggressive clinical course despite of oncotherapy, and the patient had expired three months later.

DISCUSSION

Neuroendocrine tumors are uncommon neoplasms of digestive tract. Even, ampulla of Vater NECs comprise less frequent percent in this group. Although advancement in technology eased the way for management by means of improvement in radiology (Multislice CT, magnetic resonance imaging

(MRI), endoscopy and endoscopic US (EUS)), preoperative diagnosis of NECs is often challenging [5].

LCNECs are rare and very aggressive subtype of NETs. Since they are not common, clinical features and treatment strategies are not established yet. According to some studies, tumor recurrence and poor survival rates are correlated with liver metastasis [6].

Previous studies claim that 50% of NEC of ampulla of Vater boulder lymph node metastasis. Their recommendation is pancreaticoduodenectomy (PD) together with lymph node dissection for better overall survival [7]. On the other hand; recent literature emphasize more on the histopathologic features of the tumor (differentiation grade, small or large cell type) rather than 'TNM' or

'ENETS' (European Neuroendocrine Tumor Society) staging systems in predicting overall course and treatment strategy [8].

We performed PD with lymph node dissection for our case. According to WHO classification system, it was a G3 (Grade 3) tumor having 80% Ki-67 labeling index. Oncology preferred close follow-up with conservative approach instead of giving oncotherapy. In a study presented by Naoya Imamura et al., adjuvant chemotherapy after radical resection performed for LCNECs had been found with better overall survival without any relapses on the long term. In 'The NANETS Consensus Guidelines for the Diagnosis and Management of Poorly Differentiated (High-Grade) Extrapulmonary Neuroendocrine Carcinomas' which was conducted by Strosberg JR et al, adjuvant chemotherapy (4-6 cycles of cisplatin or carboplatin and etoposide) is recommended regarding post-operative treatment of poorly differentiated NECs [9]. Considering the tumor characteristics of our case, maybe it could be a better choice to give adjuvant therapy to prevent distant organ metastasis.

Conclusion

We presented our clinical experience of LCNEC of ampulla of Vater. These uncommon tumors may have a very aggressive clinical behaviour despite of small tumor size and/or lymph node negativity. Especially for tumors equal or larger than 1 cm and having high percent Ki-67 labeling index, radical surgery with adjuvant therapy may increase overall survival.

Informed Consent: Written informed consent was obtained from the patient for the publication of the case report and the accompanying images.

Conflict of Interest: No conflict of interest was declared by the authors.

Financial Disclosure: The authors declared that this study has received no financial support.

Authorship Contributions: Concept – RES, FB; Design – RES, RBG; Supervision – GB, OA; Funding – RBG; Materials – RBG, TTE; Data collection and/or processing – FB, TTE; Analysis and/or interpretation – RES, FB; Literature review – GB, OA; Writing – RES, TTE; Critical review – GB, OA.

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