Gastroblastoma, a biphasic neoplasm of stomach: A case report

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ABSTRACT

INTRODUCTION: Gastroblastoma is a rare gastric biphasic tumor with both epithelial and mesenchymal components. To the best of our knowledge only eight cases have been reported in the English literature till date.

PRESENTATION OF CASE: We report a case of a 29-year-old female, hospitalized for epigastric pain with poor general condition. An upper gastrointestinal endoscopy showed a polyoid mass in the stomach near the gastric cardia suspicious of gastrointestinal stromal tumor. The patient underwent atypical proximal gastrectomy with splenectomy. Detailed histopathological examination of the resected specimen revealed the diagnosis of gastroblastoma. After six months, the patient developed loco-regional recurrence for which surgical debulking was performed.

DISCUSSION: Gastroblastoma is predominantly seen in young adults with non-specific complaints. They appear as submucosal lesion in the stomach mimicking gastrointestinal stromal tumor. Preoperative diagnosis is often difficult. Surgical resection remains the mainstay of treatment. On histology, they consist of mesenchymal component which stain positively for vimentin and CD10 and epithelial component which is positive for cytokeratin on immunohistochemistry.

CONCLUSION: Gastroblastoma is a malignant tumor with risk of local recurrence after curative resection.

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1. Introduction

Gastroblastoma is a recently described gastric tumor characterized by presence of both epithelial and mesenchymal components. This biphasic tumor was first reported to be a distinct entity by Miettinen et al. in 2009 [1]. Till date, only eight cases have been reported in English literature. Due to the rarity of this disease, the etiopathogenesis, malignant potential and appropriate treatment for this disease remains unknown.

We report a case of gastroblastoma with aggressive behavior in a young lady who developed local recurrence within six months of curative treatment. Additionally, we performed a literature review so as to provide better understanding of the clinico-pathological characteristics, therapeutic approaches and prognosis of this rare type of gastric tumor. This case has been reported in line with the SCARE criteria [2].

2. Case description

A 29-year-old lady presented with complaints of epigastric pain for eight months. The pain was dull aching in nature, paroxysmal, unrelated to food and without radiation. Patient also had a single episode of hematemesis two days prior to the admission.

Upper gastrointestinal endoscopy revealed the presence of a 7-cm polyoid submucosal lesion near the gastric cardia on the posterior wall of the stomach towards the greater curvature. The lesion was located about 30 cm from the incisors. Endoscopic biopsy was inconclusive. Laboratory tests showed hypochromic microcytic anemia (Hb – 6 g/dl, MCV – 66.7 fl, MCHC – 27 g/dl). Liver and renal function tests were normal. Serum CEA level was normal and serum CA 19-9 level was 12 U/ml.

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Abdominal CT scan showed the presence of a solid-cystic tumor of 7 cm arising from the posterior wall of the stomach close to the gastric cardia (Fig. 1).

Intra operatively, the tumor was located on the posterior gastric wall occupying the proximal stomach with encroachment of the splenic hilum. Atypical partial gastrectomy with splenectomy was performed due to the extension to the splenic hilum. Postoperative recovery was uneventful.

On macroscopic examination, the tumor was 7 × 4 × 4 cm in size. On cut section, there were focal areas of necrosis and hemorrhage. The tumor showed infiltration of all the layers of the gastric wall with invasion of the splenic hilum. Six enlarged lymph nodes were found at the splenic hilum and two enlarged lymph nodes were located at the gastric fundus.

On microscopic examination, the tumor was predominantly located in the submucosa with focal invasion of the mucosa, muscular layer, serosa and splenic hilar fat at places. The tumor consisted of two components – mesenchymal and epithelial (Fig. 2). The predominant component was mesenchymal, consisting of ovoid cells arranged in layers, with regular nuclei and scanty cytoplasm (Fig. 3). There were 21 mitoses per 10 high power fields. The epithelial component was focally distributed, consisting of lightly-reduced glands, lined by cuboidal to columnar cells with atypical hyperchromatic nuclei and infrequent mitosis (Fig. 4). Both components were of blastematic immature type. Peritumoral vascular emboli were present. Two of the six nodes of the splenic hilum were metastatic while both of the lymph nodes at the fundus were free of tumor.

On immunohistochemistry, the tumor was uniformly positive for vimentin, CD 99 (Mic 2) and focally positive for CD 10 (Fig. 5). Staining with anti-cytokeratin antibodies, anti-chromogranin, anti-synaptophysin and anti-C-kit was negative. Based on the above
histopathological findings, final diagnosis of gastroblastoma was made.

At six months of follow up, patient developed loco-regional recurrence in the retro-gastric area for which surgical debulking was performed. However, the patient died one month after debulking due to massive pulmonary embolism.

3. Discussion

Gastroblastoma is a very rare epithelio-mesenchymal biphasic gastric tumor. The etiopathogenesis of this tumor is not very clear, however, it is believed to develop from a totipotent stem cell [3]. It predominantly occurs in young adults with a mean age of 22 years [1]. Among the eight previously reported cases, there were six males and two females. Similar to these cases, our patient was a young girl in her third decade of life. Clinically, the symptoms are nonspecific and dominated by atypical epigastric pain, impaired general condition, and abdominal mass if the tumor is of large size [4]. In two previously reported cases, gastroblastoma manifested as gastrointestinal bleed similar to that reported in the present case (Table 1).

On upper gastrointestinal endoscopy, it appears as a bulging mass lifting the gastric mucosa [5]. The overlying gastric mucosa may be normal or ulcerated. If the tumor is developing exoluminally, the tumor may not be visible on endoscopy and even deep-seated biopsies do not allow the diagnosis [5]. Endoscopic ultrasound is recommended in such cases to visualize the tumor and perform deep biopsies safely [5]. This aspect was noted in our observation. Out of the eight reported cases, gastroblastoma was located in the antrum in five patients, along the greater curvature in two cases and in one patient the location was not specified. In our case the tumor was present along the greater curvature in the proximal stomach.

Surgical resection with clear margins has been the preferred treatment of choice. Like gastrointestinal stromal tumors, laparoscopic approach may be useful for small tumors less than 5 cm located in the anterior surface of the stomach away from the gastroesophageal junction. Gastroblastomas are lymphophic tumors indicating the importance of a lymph node dissection [1]. In our case, this dissection was not performed because of the preoperative diagnosis of a stromal tumor.

Diagnostic confirmation is possible only by the histopathological examination. Macroscopically, they can have varied size ranging from 3.8 to 15 cm. The tumor consistency is variable. The cut surface often shows areas of hemorrhage or necrosis as seen in the present case [1]. It can appear as ulcerated, polypoidal, intramural or exophyctic lesion [1].

In our case, the tumor was predominantly located in the submucosa with focal areas of infiltration in to the gastric mucosa leading to ulcerations. The tumor also seeped through the muscular and serosal layer to involve the splenic hilum.

Histologically, gastroblastoma shows biphasic cell proliferation involving mesenchymal and epithelial cells. The epithelial component is organized mainly in nests, cords and tubules composed of glands with reduced light, carpeted in places by the cubo-cylindrical cells, the hyper-chromatic nuclei and the slightly eosinophilic cytoplasm, and discrete nuclei. In some cells, the nucleus and cytoplasm were condensed and clearly bordered by separate cells. The nuclei of glandular structures were dark and elongated [1]. The mesenchymal cells were arranged in short fascicles or layers formed by oval cells with scant cytoplasm with inconspicuous nuclei and regular nuclei. No signs of structural differentiation such as fences or nuclear vacuolation were noted. Both components showed blastemal immature appearance.
Gastroblastoma variably combines both cell components, in fact, the mesenchymal component is often predominantly observed in 4 cases in the literature [4].

The mitotic index varied from 0 to 30.

Immunohistochemically, as per the original description by Miettinen M et al. [1], mesenchymal cells express vimentin and CD10 while epithelial cells express cytokeratins. However, subsequent case studies on gastroblastoma have reported positivity for additional markers as summarized in Table 2. In the present case, the mesenchymal cells showed positivity for vimentin but negative reaction for CD 10, and epithelial cells express some CD vimentin with CD10 positive response of mesenchymal and epithelial cells for CD 99. The tumor showed negative reaction for markers of gastrointestinal stromal tumors (CD117), neuroendocrine tumors (synaptophysin, chromogranin), synovial sarcoma, and leiomyosarcoma.

The differential diagnosis includes inflammatory myofibroblastic tumors, teratoma, gastrointestinal stromal tumor (GIST), synovial sarcoma, and carcinosarcoma. The literature review shows that lymph node metastases have been reported in one case, as in our patient who developed local and distant lymph node metastases [4]. Furthermore, Wey et al. [6] have reported a case of gastroblastome with lymph node metastasis and at distance. No standard therapy has been established for this tumor. The intra-peritoneal chemotherapy could reduce the loco-regional recurrence and peritoneal dissemination [7]. One reported case had received neo-adjuvant chemotherapy but without response [6]. Another one had received post-operative radiotherapy [11].

Gastroblastoma appears to have low malignant potential as recurrence after curative resection has not been reported in the literature. Our case is the first report of local recurrence after curative resection. The follow-up periods were from 3 months to 14 years. No cases of recurrence were reported during the reported monitoring period. The prognosis depends on several parameters including the size of the tumor, the degree of parietal invasion, mitotic index, and lymph node invasion [7].

Gastroblastoma seems to have a low malignant potential. The average survival was 48.3 months [4], with a range from 3 months to 14 years. In our case, the survival was 7 months.

4. Conclusion

Gastroblastomas is a distinct clinico-pathological entity due to its clinical, radiological, histopathological and immunohistochemical profile. Our case is the first reported case in Tunisia and the ninth observation described in the literature. More data are needed to understand the biological and clinical behavior of the tumor. Despite the development of the diagnostic, morphological, immune-histochemical and anatomo-pathological techniques, the diagnosis is often difficult.

Conflict of Interest

The authors have no conflict of interest to declare.

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Ethical approval

This case study was approved by the Ethics Committee of Hopital Fattouma Bourguiba.
Consent

Written informed consent was obtained from the patient

Authors’ contributions

Study concept or design – OT, HA,
Data collection – HA, TK, OT
Data interpretation – MA, HA, RG
Literature review – HA, RS, AZ,
Drafting of the paper – HA, RG, TK
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