Endoscopic resection of a periamillary gangliocytic paranglioma of the duodenum: A case report

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\textbf{ABSTRACT}

\textbf{INTRODUCTION:} The gangliocytic paranglioma (GP) is an extremely rare neuroendocrine tumour originating from the second part of the duodenum. Generally GP shows benign clinical behaviour. The GP is typically characterized by consisting of three tumour components: the epithelioid, the spindle-shaped and the ganglion-like cells.

\textbf{PRESENTATION OF CASE:} We present a female patient at the age of 65, who underwent a routine gastroscopy due to known gastro-oesophageal reflux. Accidentally a 2 × 1.5 cm sized mass of unknown entity was revealed in the duodenum. The magnet resonance tomography neither detected distant metastasis nor any local lymphadenopathy. After endosonographically guided punctures of the submucosal mass, a malignant tumour could not be diagnosed thus the decision to perform an endoscopical resection was made and successfully conducted. Immunohistochemical examination revealed a total resected GP.

\textbf{DISCUSSION:} In literature malignant transformation with distant metastasis and local recurrences has been described. Furthermore the clinical manifestation and location varies. The GP has often been misdiagnosed as a neuroendocrine tumour (NET) G1.

\textbf{CONCLUSION:} Due to published cases of metastasising GPs, we recommend a long term follow-up. In a non-metastatic stage the endoscopic resection should be the therapy of choice in order to prevent unnecessary major surgical interventions. In this case report we will discuss the clinical appearance, behaviour and differential diagnosis of GP.

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1. Introduction

The GP is a rare submucosal gastrointestinal tumour with mostly benign clinical behaviour. The tumour was first described by Dahl et al. in 1957 [1]. Generally the GP is characterized by consisting of three tumour components: spindle-shaped cells with Schwannian differentiation, ganglion-like cells and neuroendocrine cells [2,3]. These cells can be immunoreactive to pancreatic, synaptophysin, chromogranin and are able to express S-100, NSE, pancreatic polypeptide, somatostatin and vasoactive intestinal peptide [2]. The spindle cells are characterized by forming fascicles or envelope nerve cells and axons expressing the S100 protein. The ganglion cells are scattered as single elements or appear aggregated in clusters [4].

The GP typical occurs in the 5th decade of life [5]. Men are more often affected than women (1:8:1). A literature review by Okubo et al. revealed that approximately 90% of the published cases of GPs originate from the duodenum [2]. These tumours are classified as neoplasia of the digestive tract by the World Health Organisation [4]. In literature the GP has been described as appearing in the esophagus, jejunum, pylorus and even in the retroperitoneal space [6]. The main symptoms are mostly abdominal pain, discomfort, anaemia and upper gastrointestinal bleeding. To diagnose a GP can be extremely challenging. Due to the submucosal location of GP, an endoscopic biopsy often times presents negative results [4]. Moreover a GP can be confirmed by computer tomography, ultrasound imaging, magnet resonance tomography and detection of neuronal and neuroendocrine markers such as synaptophysin, chromogranin A, CD56, S-100, NSE and somatostatin in the blood [7]. Additionally the diagnosis can be supported by immunohistochemical findings such as oestrogen, progesterone receptors, pancreatic polypeptide, somatostatin and vasoactive intestinal polypeptide (VIP) [7]. Therapy of choice is the local endoscopic excision. In case of lymph node or distant metastasis, a pancreaticoduodenectomy is recommended [8].

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2. Presentation of case

In November 2015 a 65-year-old woman was treated in our hospital due to a submucosal tumour of uncertain origin. Her medical history included a diverticulosis and rheumatoid arthritis. The patient suffered from a hiatus hernia and gastro-oesophageal reflux. Therefore she routinely underwent a gastroscopy at her local hospital, where a 2 × 1.5 cm sized mass was found in the second part of the duodenum. After performing magnetic resonance tomography with cholangiopancreatography distant metastasis and lymphadenopathy were not detected (Fig. 1).

Due to the submucosal location several attempts to obtain a histopathological diagnose of the tumour were unsuccessful. Under suspicion of a malignant tumour such as carcinoma of the duodenal papilla the patient was advised to undergo an open surgery exploration. She rejected the proposal, yet she came to our hospital for a second opinion and further diagnostic.

The physical examination did not reveal any pathological findings. Furthermore the laboratory test demonstrated normal levels of C-reactive protein, leucocytes, CA72-4 and CA19-9. The tumour marker CEA was slightly increased (4.1 μg/l). We performed a gastroduodenoscopy with ultrasound guided puncture and endosonographic imaging of the submucosal tumour. The mass was anechoic and seemed to be located in all wall layers. Swollen lymph nodes were not detected. The histopathological examination revealed a tumour arising from the submucosa, covered by mucous membrane and surrounded by smooth muscle. Due to the lack of distant metastasis and lymphadenopathy, the decision for an endoscopic removal of the tumour was made. Under analgesic sedation we performed a gastroduodenoscopy. The mass of unclear entity was located approximately 2 cm above the Papilla vateri. After submucosal injection with adrenalin and mucosal dissection the tumour was successfully removed via cautery snare. The mass seemed to originate from the muscularis propria. The defect of the mucosa was covered with two clips (Fig. 3). The post interventional course was without complications and the patient left the hospital within the next 2 days (Fig. 2).

2.1. Pathological findings

The histopathological examination reveals a GP. Fig. 4 shows a 19 × 15 × 10 mm large tumour with yellow-grey cut surface as a submucosal mass with trilinear differentiation.

2.2. Immunohistochemical findings

Immunohistochemical findings are imaged in Fig. 5. Neuronal, gangliocytic and nest-like epithelioid structures are imaged in Picture 1. The S100 expression of spindle cell with Schwann-like differentiation is imaged in Picture 2. The neuroendocrine cell and ganglion cells show synaptophysin expression is demonstrated in Picture 3. The VIP expression is shown in Picture 4.

As follow-up diagnostic we performed a gastroduodenoscopy 3 month after the endoscopic removal of the GP. No recurrence was
Fig. 3. Images of the gastroduodenoscopy. The tumour was located approximately 2 cm above the Papilla vateri (A). After submucosal injection with adrenalin (B) and mucosal dissection (C) the tumour was successfully removed via cautery snare. The defect of the mucosa was covered with two clips (D).

Fig. 4. The GP is shown with yellow-grey cut surface as a submucosal mass with trilinear differentiation. (HE staining, Magnification 4×).

revealed. Furthermore we recommended performing a gastroduodenoscopy as follow-up diagnostic annually.

3. Discussion

The periampullary GP is an extremely rare tumour entity. Its exact origin still remains unclear. The GP is typically characterized by consisting of three tumour components: the epithelioid, the spindle-shaped and the ganglion-like cells [7,3]. The GP is described as being a tumour with benign clinical behaviour. In a literature review Okubo et al. retrieved 192 patients with GP, 90% of them being located in the duodenum [2]. In a follow up period from 12 to 96 months no death from GP was observed. Furthermore neither Rafiullah nor Yang observed any relapses of duodenal GPs in their case reports during a long term follow up of 3 years [9,10]. Moreover Okubo et al. performed a multi-institutional retrospective study in japan [7]. No local recurrence or distant metastasis were detected in 12 cases of mainly endoscopically removed duodenal GP. In contrast Li et al. published a case report of a 47-year-old male patient suffering from a GP, which was removed via pancreaticoduodenectomy [3]. Four months after the removal computer tomography imaging revealed distant metastasis in the liver and pelvic cavity. The tumour did not show remarkable sensitivity to chemo- or radiotherapy. The patient died after tumour progress with fever and ascites. Also Armin et al. published a case of a GP that developed liver metastasis spreading to the retropancreatic space [6].

The main symptoms of a GP are gastrointestinal bleeding and abdominal pain. This tumour entity typically arises from the second part of the duodenum [7]. In literature the description of its clinical manifestation and location varies. Eeden et al. published a
case of 47 year old male patient revealing a GP in the appendix. This tumour was mimicking an acute appendicitis. Therefore he underwent an appendectomy [11]. Furthermore Lin et al. reported a case of an asymptomatic \(4 \times 3.8 \times 2.5 \text{ cm} \) sized GP located in the right lower lobe of the lung [12]. And Yang et al. diagnosed as well as removed a 1.6 cm sized tumour in the anterior mediastinum. Later immunohistochemical examination revealed a GP surrounded by normal thymic tissue [13]. The GP was even detected in the superior mediastinum and the esophagus [14]. Vural et al. treated a 17-year-old boy due to low back pain and bilateral sciatica [15]. An intradural-extraduillary mass at the level of the L4, exiting through intervertebral foramen into the right psoas muscle, was diagnosed. The tumour was removed and a GP was found. Moreover Mahdavi et al. published a case report of a 55-year-old Caucasian woman suffering from a cystic teratoma with nodules consisting of GP [16]. Furthermore Sinkre et al. treated a 44-year-old woman with a GP in the nasopharynx. The tumour caused headaches and symptoms of abdominal fullness [17].

The GP is a rare neuroendocrine tumour, which has often been misdiagnosed as a neuroendocrine tumour (NET) G1. Both tumour entities are, according to the WHO classification, “neuroendocrine tumours of the ampullary region” [4]. But a literature review by Okubo et al. showed that mitosis and prominent Ki-67 immunoreactivity, important markers for diagnosing of NET G1, were not detected in any published cases of GP [7]. Also an immunohistochemical examination comparing the GP and the NET G1 in the beforehand mentioned review indicated a significant higher positive reactivity for progesterone, estrogen receptors and pancreatic polypeptide (PP) in the GP. Accordingly Armin et al. assumed that PP, which was detected in the serum of a patient suffering of metastasised GP, is a possible suitable marker for the differentiation of GP from NET [6]. In literature NET G1 in comparison to GP is described as a being a tumour entity with a higher risk of metastasising, primarily to lymph nodes and the liver [4]. It is therefore mandatory to clearly differentiate the GP from NET G1 due to different therapeutic approaches.

4. Conclusion

The GP is a rare neuroendocrine tumour entity typical arising from the second part of the duodenum showing mostly benign clinic-
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