Endoscopically well-differentiated neuroendocrine tumors, also called carcinoid tumors, in the duodenum are rare. The therapeutic approach is highly dependent on both tumor size and depth of invasion; for tumors smaller than 1.0 cm and without penetration of the muscularis propria, endoscopic resection is considered as the method of choice [1].

A 65-year-old woman with a histologically proven neuroendocrine tumor in the duodenal bulb was referred for further evaluation. Upper gastrointestinal endoscopy (Fig. 1) revealed a single, slightly elevated, round lesion that was covered by normal mucosa with a central depression, located in the posterior duodenal bulb. Endoscopic ultrasonography (Fig. 2) revealed a 10-mm lesion without penetration into the muscularis propria. There were no signs of regional lymph node metastasis.

Endoscopic resection with the cap technique (Fig. 3) was carried out. After the resection, an arterial bleeding was noted, which was successfully controlled with a hypertonic saline and epinephrine injection and placement of four metal clips (Fig. 4). Macroscopically, the tumor was completely removed (Fig. 5), and this was confirmed histologically (Fig. 6). Immunohistochemical staining was strongly positive for synaptophysin and chromogranin. Recovery was uneventful and the patient was discharged the following day after a second-look endoscopy.

The present case illustrates that endoscopic en-bloc resection with the cap technique is an effective method for the curative treatment of carcinoid tumors in the narrow area of the duodenal bulb. Laparoscopic techniques may be considered as an alternative only in cases where endoscopy is deemed unsuitable [2].

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