



## IMAGE OF THE MONTH

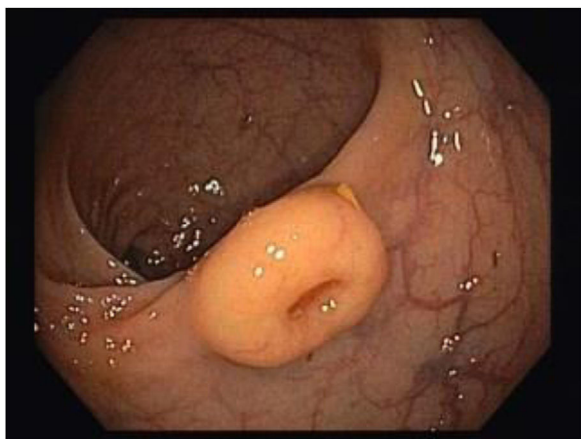
### Rectal neuroendocrine tumor – Case report

#### Tumor neuroendocrino rectal: un caso clínico

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A 60-year-old woman performed a screening colonoscopy, which showed, in the midrectum, a 12 mm, sessile, subepithelial lesion covered with yellow discolored mucosa, with a central depression (Fig. 1). Conventional biopsies showed non-specific inflammatory infiltrate. Rectum endoscopic



**Figure 1** Colonoscopy: In the midrectum, a 12 mm, sessile, subepithelial lesion covered with yellow discolored mucosa, with a central depression.

ultrasound showed an hypoechoic lesion with 12 mm, homogeneous, well delimited, originating in the muscularis of the mucosa with no lymph nodes or muscularis propria involvement (Fig. 2). Thoraco-abdomino-pelvic computed tomography showed no locoregional/distant metastasis. An endoscopic mucosal resection of the lesion in a single fragment was performed. Histopathology showed a 12 × 8 × 9 mm, well-differentiated neuroendocrine tumor (NET) G1 (WHO classification, <2 mitoses/10 HPF, Ki-67<2%), with no linfovascular invasion and with free margins. One-year later, the patient repeated colonoscopy with biopsies which showed no evidence of recurrence.

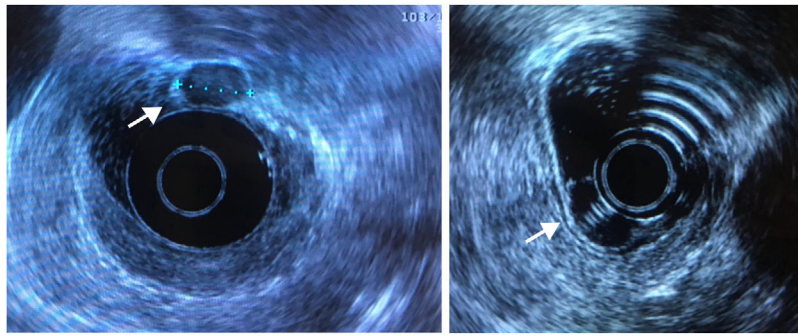
Rectal NETs (R-NET) represent approximately one-third of all gastroenteropancreatic NETs.<sup>1–3</sup> Due to the widespread use of screening colonoscopy, the incidence of R-NET has been increasing. The majority are small, well-differentiated, limited to the submucosal layer, with a good prognosis. Endoscopic ultrasound is essential to assess tumor size, depth of invasion and presence of lymph node invasion, in order to determine the appropriate treatment strategy.<sup>1,2,4</sup> This case underscores the importance of early identification and optimal management of these tumors.

#### Author contributions

Carolina Simões and Sofia Carvalhana were responsible for elaborating the manuscript. Leonor Xavier de Brito, Luís

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**Figure 2** Rectum endoscopic ultrasound: hypoechoic lesion with 12 mm (arrow), homogeneous, well delimited, originating in the muscularis of the mucosa with no lymph nodes or muscularis propria involvement.

Carrilho Ribeiro and Rui Tato Marinho were responsible for the critical review of the manuscript.

The manuscript, including related data, figures and tables have not been previously published and are not under consideration for publication elsewhere.

### Informed patient consent

Obtained.

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The authors have received no funding.

### Conflicts of interest

The authors have no conflicts of interest to report.

### References

1. Kunz PL, Reidy-Lagunes D, Anthony LB, Bertino EM, Brendtro K, Chan JA, et al. Consensus guidelines for the management, treatment of neuroendocrine tumors. *Pancreas*. 2013;42:557, <http://dx.doi.org/10.1097/MPA.0b013e334a41828>.
2. Caplin M, Sundin A, Nillson O, Baum RP, Klose KJ, Kelestimur F, et al. ENETS Consensus Guidelines for the management of patients with digestive neuroendocrine neoplasms: colorectal neuroendocrine neoplasms. *Neuroendocrinology*. 2012;95:88–97, <http://dx.doi.org/10.1159/000335594>.
3. Raposo André T, Brito M, Geraldes Freire J, Moreira A. Rectal and anal canal neuroendocrine tumours. *J Gastrointest Oncol*. 2018;9:354–7, <http://dx.doi.org/10.21037/jgo.2017.10.01>.
4. Rodrigues A, Castro-Poças F, Pedroto I. Neuroendocrine rectal tumors: main features and management. *GE Port J Gastroenterol*. 2015;22:213–20, <http://dx.doi.org/10.1016/j.jpgge.04.008>.