An 88-year-old man with a past medical history of atrial flutter and congestive heart failure presented with rectal bleeding. Owing to his reluctance to undergo colonoscopy, he instead underwent computed tomography colonography, which showed one lesion in the splenic flexure and one lesion in the distal sigmoid colon, suggestive of colonic neoplasms (Fig. 1). The patient later did undergo colonoscopy, which showed multiple yellowish, polypoid lesions dispersed throughout the colon (Fig. 2a–c), some with erosions (Fig. 2d). Several biopsies were taken from different segments of the colon.

The biopsies all showed an extensive diffuse infiltration of the lamina propria by neoplastic lymphoid cells of intermediate/large size, with irregular nuclei, fine chromatin, and scant cytoplasm (Fig. 3a). The mitotic index was moderate. Immunohistochemistry showed positivity for CD45, CD20, CD5, and cyclin D1, consistent with mantle cell lymphoma, blastoid variant (Fig. 3b). A computed tomography scan showed multiple enlarged mesenteric lymph nodes and splenomegaly. As the patient did not have any obstructive symptoms, he was referred for treatment with chemotherapy.

Multiple lymphomatous polyposis (MLP) is an extremely rare form of primary gastrointestinal lymphoma that is characterized by multiple polyps formed by neoplastic lymphoid cells [1]. Most cases of MLP are found in mantle cell lymphoma. The blastoid variant of mantle cell lymphoma is so-called because of its blastoid cytology with large round-to-oval nuclei and prominent nucleoli, in contrast to the much more common classic variant of mantle cell lymphoma, which has nuclei of small-to-medium size with irregular contours [2, 3]. Most of the cases of MLP previously reported in the literature have been in classic mantle cell lymphoma with typical cytology and low proliferation index.

Competing interests: None

References
Fig. 3  Histology of the colonic biopsies showing: a extensive diffuse infiltration of the lamina propria by neoplastic lymphoid cells of intermediate/large size, with irregular nuclei, fine chromatin and scant cytoplasm (hematoxylin and eosin [H&E] stain, original magnification × 100); b positive immunostaining of lymphoid cells with cyclin D1, consistent with mantle cell lymphoma (original magnification × 200).

Corresponding author
Eduardo Rodrigues-Pinto, MD
Gastroenterology Department,
Centro Hospitalar São João
Al. Prof. Hernâni Monteiro
4200-319 Porto
Portugal
Fax: +351-22-5513601
edu.gil.pinto@gmail.com