Secondary rectal linitis plastica as first manifestation of urinary bladder carcinoma

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Abstract

Secondary rectal linitis plastica is a very rare malignancy with poor prognosis. Diagnosis is difficult because of nonspecific clinical and endoscopic findings and negative biopsies in most cases owing to the fact that the mucosa is frequently unaffected. We herein describe a 68-year-old man who presented with a six-month history of tenesmus and constipation. Endoscopy revealed a narrow distal rectum with an indurated, cobblestone appearance of mucosa. Multiple biopsies and fine-needle aspiration were negative for malignancy. He underwent rectal extirpation with total cystectomy and lymph nodes dissection. Histology demonstrated secondary rectal linitis plastica due to a poorly differentiated urinary bladder carcinoma. We emphasize the endoscopic and endosonographic features and the difficulty to establish a preoperative diagnosis of secondary rectal linitis plastica.

Keywords

Secondary rectal linitis plastica, urinary bladder carcinoma

Introduction

Linitis plastica is characterized by the diffuse infiltration of the submucosal and muscularis propria layers of a hollow organ with cancer cells, resulting in wall thickening that makes the involved organ constricted, inelastic and rigid. Although it is most commonly seen in the stomach, other organs, including small intestine, colon and rectum, are occasionally involved [1]. Rectal linitis plastica (RLP) is a rare tumor with an incidence of 1 in 1,000 colorectal carcinomas. Secondary RLP is more common than a primary one, having as a predominant cause for its development, previous gastric linitis plastica.

Case report

A 68-year-old man presented to the Department of Endoscopy with a six-month history of anal outlet obstruction, constipation and tenesmus. The patient had no relevant past medical history. Digital examination revealed a circumferential rectal narrowing with a firm, mass-like area, palpable all around the rectal wall and extending to a few centimeters of the anal margin. Laboratory tests, including urinalysis, were unremarkable. Subsequent endoscopy demonstrated a narrow distal rectum, with the overlying mucosa being indurated, non-ulcerated, with a cobblestone appearance (Fig. 1). Several biopsies and fine-needle aspiration were taken, but none was conclusive for diagnosis. T2-weighted MRI showed a double-layered thickening of the rectal wall with an inner iso-intense circumferential thickening of the submucosa and outer hypo-intense circumferential thickening of the muscular rectal wall, as well as a thickened bladder wall. Transrectal ultrasonography showed circumferential infiltration of the...
submucosa, whereas the other layers were normal (Fig. 2); an image compatible with “linitis plastica”. The patient underwent rectal extirpation and total cystectomy with lymph node dissection. Macroscopic examination of a sagittal section of resected rectum demonstrated an expanded submucosal layer due to infiltration with cancer cells, with intact mucosa and muscular layer (Fig. 3). Histological examination of the resected bladder showed a grade III papillary transitional cell carcinoma. In the rectum clusters of tumor cells of a poorly differentiated adenocarcinoma were observed. Immunostaining was positive for both CK7 and CK20; PSA was negative. Combining the cytokeratine profile histology and transrectal ultrasonography findings, the diagnosis of a primary urinary bladder tumor with hematogenous spread to rectal submucosal layer was established.

The patient received postoperative radiation followed by gemcitabine (100 g/m² on days 1, 8, 15) and cisplatin (70 mg/m² on day 2) for a 28-day cycle for six months, but he died eight months later due to disseminated peritoneal carcinomatosis.

Discussion

RLP secondary to urinary bladder carcinoma is very rare [5,6]. Dressen et al described two patients who presented with changed bowel habits [5]. All diagnostic tests were inconclusive. In both patients pelvic MRI images revealed double-layered thickening of the rectal wall with an inner iso-intense circumferential thickening of the submucosal and hypo-intense circumferential thickening of the muscular rectal wall and a thickened bladder wall. Surgery disclosed a grade III papillary transitional cell carcinoma with infiltration of rectal wall and prostate gland in the first patient. In the second patient the RLP was due to recurrent transitional cell carcinoma of the urinary bladder. Gleeson et al described the EUS features and tissue diagnosis of secondary RLP in three patients with recurrent urinary bladder cancer [6].

Our patient was asymptomatic from the urinary tract and secondary RLP was caused by hematogenous spread of a small urinary bladder carcinoma. Our inability to establish a preoperative diagnosis, despite the fact that endoscopic and endorectal pictures were compatible with RLP, is quite common. There is usually a long delay between the onset of
symptoms and final diagnosis [5-7], caused by the fact that RLP can mimic various diseases such as inflammatory bowel disease, stenosis related to diverticulosis, ischemia, radiation-induced strictures, solitary rectal ulcer, endometriosis and malignant lymphoma [1,8-11]. Moreover, mucosal biopsies are rarely positive, as in our case, because the disease is predominantly located in the submucosal and muscularis propria. With the advent of interventional endosonography, EUS-guided fine needle aspiration is performed more and more often, but both sensitivity and accuracy of the method are low [12,13]. Surgical deep biopsies may be needed if previous biopsies are negative.

The endoscopic picture, associated with the endorectal ultrasonography findings, was compatible with RLP in our patient. Endoscopic features of RLP include luminal stenosis and indurated folds with an infiltrated or cobblestone appearance. Large cerebroid folds and ulcerations are present only in advanced cases [1,3,6]. Dumontier et al reported the main EUS finding of secondary RLP, which consists of a circumferential wall thickening, predominantly affecting the submucosal and muscularis propria layers [7].

In general, prognosis is poor because of the late recognition of the disease and the absence of any effective treatment. Even in patients with early diagnosis and limited RLP who were treated surgically, long-term survival was extremely low [14] Lymphatic invasion with lymph nodes metastases and peritoneal carcinomatosis are usually present in 50-80% of cases at the time of diagnosis and account for the poor prognosis [14].

References