Rectal Carcinoma with Heterotopic Bone: Report of a Case

Yuichi Nagao a Shohei Shimajiri b Takefumi Katsuki a Yoshifumi Nakayama a Koji Yamaguchi a
Departments of aSurgery 1 and bPathology and Cell Biology, School of Medicine, University of Occupational and Environmental Health, Kitakyushu, Japan

Key Words
Rectum · Adenocarcinoma · Osseous metaplasia

Abstract
Heterotopic bone is rarely present in malignant tumors of the gastrointestinal tract. We herein report a case of rectal adenocarcinoma with heterotopic bone. A 46-year-old Japanese male presented to our hospital with abdominal distension and constipation. Colonoscopic examination showed an ulcerated polypoid tumor of the rectum which nearly obstructed the rectal lumen. Abdominal computed tomography showed a tumor of the rectum with calcified deposits. Low anterior resection with lateral lymph node dissection was performed under the tentative diagnosis of rectal cancer. Histological examination of the resected specimen showed mucinous carcinoma of the rectum with heterotopic bone. One of the metastatic lymph nodes dissected also showed heterotopic bone. In the present report, we describe this rare tumor and briefly review the pertinent literature regarding rectal cancer with heterotopic bone.

Introduction
Heterotopic bone is very rarely seen in malignant conditions of the gastrointestinal tract. Dukes [1] first reported that gastrointestinal tumors with heterotopic ossification were low-grade malignancies with no tendency to spread by the venous or lymphatic systems. We herein present a case of heterotopic bone present in rectal cancer and in one of the metastatic lymph nodes.

Case Report
A 46-year-old male went to a neighboring hospital with a 1-month history of abdominal distension and constipation. Digital examination of the rectum revealed a palpable tumor. Colonoscopic
examination showed an ulcerated polypoid tumor almost obstructing the rectal lumen. Endoscopic biopsy of the rectal tumor led to classification of the tumor as adenocarcinoma. The patient was referred to us for further examination and treatment. Laboratory data, including tumor markers (carcinoembryonic antigen and carbohydrate antigen 19-9) were within normal limits. Abdominal computed tomography showed a rectal tumor with calcified deposits, without any sign of metastatic lymph nodes (fig. 1). Therefore, a low anterior resection with lateral lymph node dissection was performed under the tentative diagnosis of rectal cancer with calcification.

The resected specimen showed a large polypoid tumor (9 × 7 cm) with superficial ulceration situated 3 cm from the distal resection margin and almost encircling the rectum (fig. 2). Histologically, the tumor was a mucinous carcinoma extending through the muscularis propria and invading the adjacent perirectal adipose tissue with lymphatic permeation. Abundant fibrovascular stroma was present, and well-formed bony trabeculae were noted mainly in the stroma (fig. 3a). These consisted of irregular islands of mineralized osteoid bone rimmed by a layer of scattered osteoblasts. No necrosis was seen within the tumor. The surgical margins were free of tumor tissue. Eleven of the 20 lymph nodes dissected had metastatic carcinoma, and one of them showed heterotopic bone (fig. 3b). The patient’s postoperative course was uneventful, and he was doing well with no signs of recurrence 3 months after the operation.

Discussion

Rectal cancer is a common disease, but coincidental heterotopic bone is rarely seen. Heterotopic bone has been reported in primary and metastatic gastrointestinal tumors including carcinoma of the stomach, appendix and colorectum. It has also been observed in adenoma of the rectum and carcinoid tumors of the stomach [1–4]. In the landmark paper, Dukes reported that the incidence of ossification of rectal cancer was less than 0.4% [1]. To the best of our knowledge, there have been only ten cases of heterotopic bone in rectal adenocarcinoma reported in the English language literature [1, 2, 4–11]. Unlike the present case, these cases showed no heterotopic bone in the metastatic lymph nodes.

Some clinicopathologic features of rectal adenocarcinoma with heterotopic bone have been reported. Dukes reported the following features concerning rectal carcinoma with heterotopic bone: (1) long duration of symptoms indicating slow growth of the tumor; (2) histologically low-grade malignancy with no tendency to spread through the veins or lymphatics; (3) presence of a necrotic area within the tumor [1]. Although none have been previously reported for rectal cancer, several cases of heterotopic ossification in local recurrences or in metastatic sites have been reported in addition to heterotopic ossification in the primary sites [1, 4, 5]. In our case, eleven of the 20 lymph nodes dissected showed metastatic deposits, and one of them had heterotopic bone.

The mechanism of heterotopic bone formation within gastrointestinal adenocarcinoma is not completely understood. The most plausible explanation for heterotopic ossification in tumors of epithelial origin is a metaplastic process of the stromal mesenchymal cells, wherein they transform into osteoblasts. Supporting this theory, heterotopic bone is more likely to be found in necrotic tumors, the mucin pool, and the tumor stroma [12]. In addition, a number of other theories exist. For example, Randall et al. [3] suggested that metastatic colonic carcinoma can promote heterotopic ossification, and that alkaline phosphatase is intimately associated with bone formation under these pathological conditions, because immunostaining for alkaline phosphatase is seen not only in osteoblast-like cells, but also in the atypical membrane of the cancer cells next to areas of the bone. Rosenbaum et al. [13] suggested that large quantities of gamma-carboxyglutamic acid may be associated with the calcification of bone matrix in a
Heterotopic ossification is metaplasia of fibroblasts induced by diffusible factors released from rapidly dividing epithelial cells of the tumor or by direct contact with epithelial cells of the tumor. Imai et al. [15] reported that bone morphogenetic protein produced by gastrointestinal tract glandular tumor cells might play an important role in heterotopic ossification. In the present case, no necrosis was evident, but the histological type was mucinous adecarcinoma.

Heterotopic bone is rarely present in the gastrointestinal tract, including benign and malignant epithelial tumors. Further studies are therefore necessary to determine the factors that play a role in the development of heterotopic bone formation in gastrointestinal tumors, and to confirm the impact of this bone formation on patient outcome.

**Fig. 1.** Abdominal computed tomography showed a tumor in the rectum with calcified deposits (arrow), without any sign of metastatic lymph nodes.
Fig. 2. The resected specimen contained a large polypoid tumor (9 × 7 cm) with superficial ulceration.

Fig. 3. a Histologically, the tumor was a mucinous adenocarcinoma with well-formed bony trabeculae in the stroma (arrows). Abundant fibrovascular stroma was present. b Eleven of the 20 lymph nodes dissected had metastatic carcinoma, and one of them showed heterotopic bone (arrows).

References


