

Disseminated Carcinomatosis of Bone Marrow Due to Sigmoid Colon Cancer

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Received February 08, 2015; Revised February 24, 2015; Accepted March 03, 2015

Abstract Introduction: Disseminated carcinomatosis of the bone marrow (DCBM) caused by colorectal cancer is rarely seen. DCBM is often accompanied with diffuse intravascular coagulation (DIC). **Presentation Of Case:** A 72-year-old male patient with a history of right nephrectomy three decades ago for referred benign disease, underwent sigmoid colon resection for cancer causing bowel obstruction. The preoperative evaluation revealed elevated values of CA 19-9 and PSA, with no metastatic lesions in abdominal CT. On eighth postoperative day, the patient developed fever with anemia and thrombocytopenia. Bone marrow examination revealed disseminated carcinomatosis caused by colon cancer. The patient deceased on the twenty seventh postoperative day due to the rapid progression of the accompanied disseminated intravascular coagulation. **Discussion:** Prognosis for patients with cancer who develop disseminated carcinomatosis of bone marrow (DCBM) is poor. **Conclusion:** In our case, the rapid deterioration of the patient gave us no option for adjuvant chemotherapy. The outcome was fatal. A 72-year-old male patient with a history of right nephrectomy for benign disease underwent sigmoid colon resection for bowel obstruction due to cancer. Tumor markers CA 19-9 and PSA were elevated, with no metastatic lesions in abdominal CT of admission. On eighth postoperative day, the patient developed severe anemia and thrombocytopenia. Bone marrow examination revealed disseminated carcinomatosis caused by colon cancer. The patient deceased on the twenty seventh postoperative day due to the rapid progression of the accompanied disseminated intravascular coagulation.

Keywords: sigmoid colon, cancer, obstruction, disseminated carcinomatosis, disseminated intravascular coagulation

Cite This Article: Christos Konstantinidis, Panagiotis Varsos, and Sotirios Kypouris, "Disseminated Carcinomatosis of Bone Marrow Due to Sigmoid Colon Cancer." *American Journal of Medical Case Reports*, vol. 3, no. 4 (2015): 102-104. doi: 10.12691/ajmcr-3-4-4.

1. Introduction

Disseminated carcinomatosis of the bone marrow (DCBM) caused by colorectal cancer is rarely seen. In most cases, DCBM is derived from gastric cancer. The prognosis of the disease is poor, especially when it is complicated with disseminated intravascular coagulation (DIC). We report a fatal case of early and rapid manifestation of DCBM accompanied by DIC, which was developed after surgically treatment of sigmoid colon cancer causing bowel obstruction.

2. Case Report

A 72-year-old male patient was admitted to our hospital complaining for abdominal discomfort and constipation. He had a medical history of right nephrectomy for cystic lesions. Physical examination showed moderate abdominal distension. Hematological and blood biochemical examinations were normal, while tumor markers CEA, CA19-9 and PSA had elevated values

(13.04 ng/ml, 623.37 U/ml and 28.71 ng/ml respectively) (Table 1). Abdominal CT revealed a mass from the wall of the sigmoid colon which was causing obstruction of the lumen (Figure 1, Figure 2). The patient underwent nine days after his admission a scheduled resection of the sigmoid colon, after the placement of a stent at the left ureter, with primary end-to-end anastomosis. The histopathological examination of the specimen revealed middle grade adenocarcinoma (Figure 3), without penetration of the wall of the colon but with extensive lymphatic invasions (positive 10 of 10 lymph nodes and other more lymphatic blocks) (Figure 4). The postoperative period was uneventful until the fifth postoperative day, when the complete blood cell account revealed mild thrombocytopenia (~100,000/uL), which worsened rapidly the following days. On eighth postoperative day, the patient developed fever with anemia and severe thrombocytopenia. The total amount of platelets dropped to levels less than 20,000/uL. Despite blood infusions the amount of platelets remained at extremely low levels. Additional abdominal CT revealed lesions at the body of lumbar vertebra while chest CT demonstrated diffuse peripheral pulmonary emboli (Figure 5), laboratory test for HITTS were negative for

pathological findings. A subsequent bone marrow examination revealed diffuse neoplastic infiltration caused by moderate differentiated adenocarcinoma (CK20+, CK7-, CD10-), indicative of disseminated carcinomatosis

of the bone marrow from colon cancer (Figure 6, Figure 7). Due to patient's rapid deterioration, treatment with chemotherapy wasn't an option. The patient deceased twenty seven days after initial surgical operation.

Table 1. Laboratory Data on Admission

Complete blood cell count Chemistry Coagulation				
WBCs:	14,570/Ul	Glucose:	193 mg/dL	PT (sec): 13 aPTT (sec): 25 INR: 1.2 Immunology AFP: 1.64 ng/ml CEA: 13.04 ng/ml Ca19-9: 623.37 U/ml PSA: 28.71 ng/ml Free PSA: 1.89 ng/ml PSA ratio: 0.07
RBCs:	5,170 x 10 ³ /uL	BUN:	42 mg/dL	
Hemoglobin:	14.6 g/dL	Creatinine:	1.0 mg/dL	
Hematocrit:	44.7 %	K:	4.3 meq/L	
MCV:	86.6 fL	Na:	139 meq/L	
MCH:	28.2 pg	SGOT:	24 IU/L	
MCHC:	32.6 g/dL	SGPT:	11 IU/L	
Platelets:	189 x 10 ³ /uL	ALP:	305 IU/L	
Neutrophils	88.2 %	γ-GTP:	17 IU/L	
Lymphocytes	7.9 %	Amylase:	23 IU/L	
Monocytes	2.6 %	T-BIL:	1.01 mg/dL	
Eosinophils	0.2 %	D-BIL:	0.45 mg/dL	
Basophils	0.4 %			

3. Discussion

Patients with disseminated carcinomatosis of the bone marrow complain for back pain and bleeding tendencies, as was with our patient, presented as purpura. The peripheral blood examination reveals severe anemia with thrombocytopenia, elevated alkaline phosphatase and lactase dehydrogenase. Bone marrow examination demonstrates a diffusely infiltration rather than a nodular pattern.

A review of the literature shows that most studies related to disseminated carcinomatosis of the bone marrow (DCBM) were conducted in Japan. In these studies, the origin of DCBM is advanced gastric cancer, in up to 90% of cases. In the contrary, colorectal cancer remains a rare cause of DCBM, with rectal cancer slightly more frequent. In one study, only 27 cases of colorectal cancer are reported, as an origin of DCBM, between 1984 and 2013 in Japan [1].

Prognosis of disseminated carcinomatosis of bone marrow (DCBM) accompanied with diffuse intravascular coagulation (DIC) in patients with cancer is poor. Standard anticoagulant therapy is considered to be ineffective for DIC caused by malignancies [2]. Other treatment options are administration of fresh frozen plasma and platelet replacement, with or without anticoagulation therapy, which also has little impact in patients survival, who usually die within 1–3 weeks [3].

Targeting towards the underlying cancer would offer better survival rates. Various chemotherapy regimens are used for this purpose with different outcomes [4-12]. Although survival is improved in most cases, patients with colorectal cancer and DCBM do not survive more than 100 days [1]. Yoshida et al [13] report the early start of chemotherapy after surgery in patients with colorectal cancer and synchronous metastases, aiming at preventing tumor growth. The effect and safety of such an approach needs further confirmation with prospective studies.

Unfortunately in our case, the rapid deterioration of the patient gave us no option for adjuvant chemotherapy with a better outcome. His general condition was severe on the fourteenth postoperative day and thereafter, making any administration of chemotherapy prohibitive. The general condition of the patient didn't improve and he, finally, deceased on the twenty seventh postoperative day.



Figure 1. Abdominal CT scans in series indicating the sigmoid colon cancer, obstructing the lumen of the bowel (red arrows).



Figure 2. Abdominal CT scans in series indicating the sigmoid colon cancer, obstructing the lumen of the bowel (red arrows).

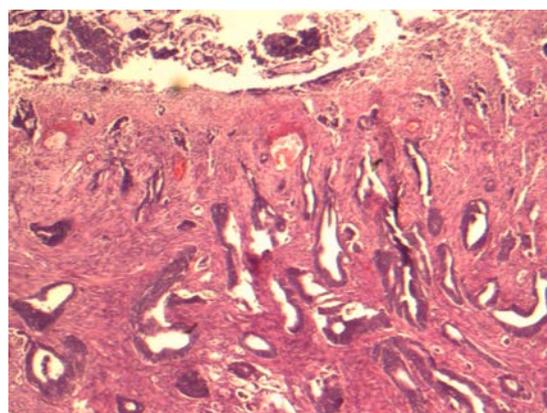


Figure 3. Histopathological specimen examination of the sigmoid colon cancer (H&E X 25)

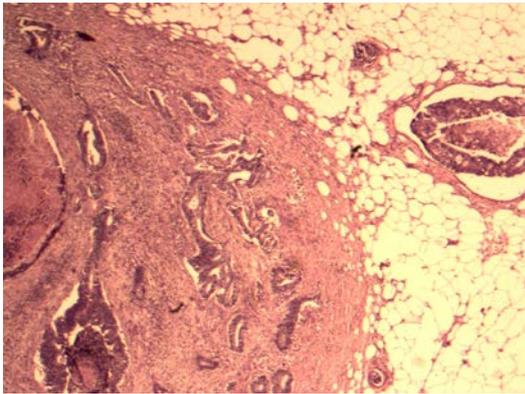


Figure 4. Histopathological examination of infiltrated lymph node from the sigmoid colon cancer (H&E X 25)



Figure 5. Chest CT scan indicating diffuse peripheral pulmonary emboli.

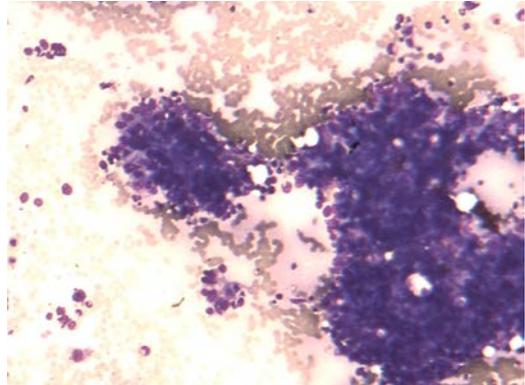


Figure 6. Bone marrow examination of the ilium that reveals a cluster of abnormal cells (stain X 100)

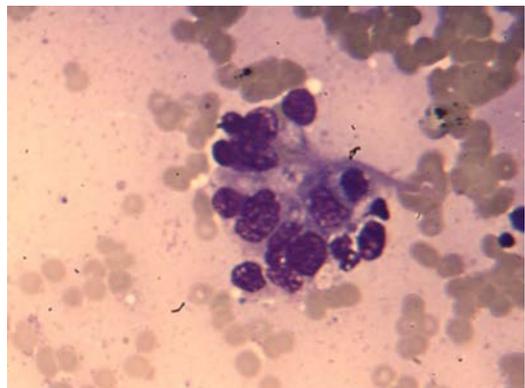


Figure 7. A different imagination of the same bone marrow examination of the ilium that reveals abnormal cells accompanied by mucous (stain X 400)

Conflict of Interest

The authors declare that there is no conflict of interest.

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