Anorectal and gastric peripheral T-cell lymphoma, unspecified in a non-AIDS patient

Eun-A Eum, M.D., Hawk Kim, M.D., Young-Min Kim, M.D.,
Soon Joo Woo, M.D., Joon Hyun Cho, M.D.,
Young Joo Min, M.D. and Jae-Hoo Park, M.D.

Departments of Internal Medicine and Pathology,
Ulsan University Hospital, University of Ulsan College of Medicine, Ulsan, Korea

Anorectum is a rare location for malignant lymphoma. Involvement of is rare even for the lymphoma associated with acquired immune deficiency syndrome (AIDS), and AIDS has a relatively increased frequency of anorectal lymphoma. Most lymphomas in AIDS patients are of a B-cell origin, and T-cell lymphoma of the gastrointestinal tract is extremely rare. We report here on a case of anorectal and gastric peripheral T-cell lymphoma, unspecified (PTCLu) in a non-AIDS patient. A previously healthy 29-year-old man presented with hematochezia and tenesmus that he had suffered with for the previous 2 months. Sigmoidoscopy showed anal and rectal submucosal tumor. Multiple round-shaped, flat and elevated lesions were noted on the gastric antrum and body as well. He underwent excisional biopsy for the anal mass and the diagnosis was PTCLu. Biopsies of the gastric lesions gave the same diagnosis. There was no lymphoma involved in the bone marrow. At admission, no antibodies against human immunodeficiency virus were detected. He underwent systemic chemotherapy and upfront autologous stem cell transplantation.

Key Words : Rectum, Peripheral T-cell lymphoma unspecified, Non-AIDS

INTRODUCTION

The gastrointestinal tract is the most common site of extranodal lymphomas, but the anorectum is not a common location for gastrointestinal lymphoma. Colorectal lymphoma comprised between 10% to 20% of the primary gut lymphomas in a large, published series. Anorectal lymphoma is so rare that its frequency among the cases of gastrointestinal lymphoma has been reported as only 4~6% in North America and 2% in Japan. For patients suffering with acquired immune deficiency syndrome (AIDS), there is a marked increase in gastrointestinal non-Hodgkin’s lymphomas (NHL); these are of a higher histologic grade and they originate from B-cells, they are more aggressive and are primary of an extra nodal origin.

The relative frequency of peripheral T-cell lymphoma (PTCL) of NHL in the gastrointestinal tract is about 2% in Korea. Most lymphomas with a primary or major location in the anorectum have been reported in AIDS patients; however, peripheral T-cell lymphoma, unspecified (PTCLu) of the rectum is quite rare. Therefore, rectal PTCLu in non-AIDS patient seems to be an extremely uncommon finding. Moreover, concurrent extranodal involvement of both the stomach and anorectum is also very rare. In some limited studies, PTCLu in the colon has manifested as either a diffuse or focal segmental lesion with extensive mucosal ulceration. We report here on an unusual case of concurrent rectal and gastric PTCLu without extensive mucosal ulceration in a non-AIDS patient.

CASE REPORT

A 29-year-old man was referred to our hospital with
complaints of hematochezia and tenesmus for 2 months. He underwent colonofiberscopic examination without biopsy by a primary clinician. The colonofiberscopic examination showed a huge multiple fungating submucosal mass at the anorectal region (Figure 1A). On admission, the patient complained hematochezia, tenesmus and abdominal discomfort, but there was no specific findings noted on the abdominal physical examination. The vital signs revealed a blood pressure of 100/70 mmHg, a pulse rate of 68/min, a respiratory rate of 22/min and his temperature was 36.5°C. The laboratory test showed a white blood cell count of 9,110/μL (84.1% polymorphonuclear cells and 9.2% lymphocytes), a hemoglobin level of 12.3 g/dL, a hematocrit of 36.8%, a platelet count of 412,000/μL, an erythrocyte sedimentation rate of 10 mm/hour and a C-reactive protein level of 0.03 mg/dL (normal range: 0–0.5). The partial thromboplastin time and prothrombin time were within normal ranges. The results of liver function tests were as follows protein 8.7 g/dL, albumin 4.5 g/dL, bilirubin 1.6 mg/dL, aspartate transaminase 26 IU/L, alanine transaminase 53 IU/L, alkaline phosphatase 51 IU/L, cholesterol 138 mg/dL and lactic dehydrogenase (LDH) 514 IU/L. The renal function tests were normal. The serum electrolytes showed sodium 138 mEq/L, potassium 4.9 mEq/L, chloride 100 mEq/L, calcium 9.2 mg/dL and phosphate 4.8 mg/dL. Urinalysis was negative for occult blood and there was a trace of protein. No antibodies against syphilis, hepatitis B, hepatitis C or human immuno-deficiency virus were detected. Under the clinical impression of anal and rectal submucosal tumor, the patient underwent excisional biopsy. The histologic findings of the specimen showed dense infiltration of angulated lymphoid cells into the lamina propria together with perivascular infiltrations (Figure 2A). On immunohistochemistry, these lymphoid cells were positive for CD3 (Figure 2B), lymphocyte common antigen, Ki-67 and CD45RO, but they were negative for CD20 and CD56. These morphologic features and the immunophenotype profiles were compatible with PTCLU. For further evaluation, he underwent...
Figure 3. Pelvic CT scan on admission (A) and after 3 cycles of chemotherapy (B). Wall thickening from the lower rectum to the rectosigmoid junction, and multiple lymphadenopathies in perirectal area, sigmoid mesocolon and paraaortic area, and enlarged nodes along hepatoduodenal ligament are all noted (A). After 3 cycles of chemotherapy, the rectal wall thickening together with the perirectal lymphadenopathies completely resolved (B).

gastrofiberscopy and bone marrow biopsy. Gastrofiberscopic examination showed mottled or patched hyperemic mucosa and round-shaped, flat elevated lesions on the gastric antrum (Figure 1B). Biopsy of the gastric lesion gave the same diagnosis, but there was no lymphoma that involved the bone marrow. The chest CT scan was normal: the abdominal–pelvic CT scan showed wall thickening from the lower rectum to the rectosigmoid junction, and there were multiple lymphadenopathies in the perirectal area, sigmoid mesocolon and paraaortic area, and enlarged nodes were noted along the hepatoduodenal ligament.

Finally, the patient was diagnosed as suffering with rectal and gastric PTCLu. The Ann Arbor stage was IV, and the age–adjusted international prognostic index was 3. He was in the high risk group. He was treated with six cycles of combination chemotherapy including cyclophosphamide, adria- mycin, vincristine, etoposide and prednisone (CHOEP). After two cycles of chemotherapy, the initially observed rectal wall thickening disappeared on the follow-up abdominal–pelvic CT scan; we observed markedly decreased size of the lymph nodes of the paraaortic and portocaval space. On follow-up gastrofiberscopic biopsy, a few lymphoid cells had infiltrated to the lamina propria, and these were probably reactive lymphocytes. He underwent autologous stem cell transplantation after six cycles of systemic CHOEP chemotherapy. Regular follow-up was done with CT, gastroscop ic biopsy and colonofiberscopic biopsy every 3 months. He has been in complete remission for over 300 days after autologous stem cell transplantation.

**DISCUSSION**

Lymphomas arising primarily in the rectum are rare occurrences that represented only 0.1~0.3% of all malignant rectal lesions according to the study data from a large population. This is probably because usually there is no lymphoid tissue in the anorectum. Anorectal and perianal lymphoma occurs in an even smaller subset of patients. There are now increasing numbers of reported cases of anorectal lymphomas that were caused by acquired immunodeficiency syndrome. Most cases have been homosexual men, suggesting anorectal trauma as a port of entry for a possible infectious, oncogenic virus. Primary anorectal lymphoma in nonimmunocompromised and heterosexual patients is a rare event with only a few isolated cases being reported. Most primary gastrointestinal lymphomas have been of a B-cell origin. Only a few cases are reported to be positive for T-cell markers. The most common site of involvement for gastrointestinal T-cell lymphoma is the small bowel, and other sites are exceptionally rare.

PTCLu is rather uncommon: it comprises approximately 9.4% of all NHLs, but it is more common in East Asia. PTCL has been subclassified in various ways and usually to include anaplastic large T-cell lymphoma, angioimmunoblastic T-cell lymphoma (AILD), PTCLu and intestinal T-cell lymphoma (ITCL). The PTCLu group showed the second highest response rate, but also the highest relapse rate, and this suggests that while responses are good, they are somewhat brief in duration. Lopez Guillermo et al. concluded that the outcome for PTCLu patients was basically unfavourable with less than half reaching complete remission and most patients eventually relapsed. It is generally assumed that PTCLu probably constitutes a heterogeneous group encompassing several different subtypes that are currently not recognized, and this may partly account for the variation in the findings. According to the recent literature, PTCL involving the colon corresponds to a diffuse ulcerative form of colorectal lymphoma, although this is very rare. The radiologic differentiation of PTCL involving the colon from inflammatory bowel disease is very difficult.
Moreover, pathologic differentiation of PTCL from inflammatory bowel disease is also difficult. In this case, colonofiberscopic examination showed a huge multiple fungating submucosal mass at the anorectal region, and the gastrofiberscopic examination showed mottled or patched hyperemic mucosa and round-shaped, flat elevated lesions on the gastric antrum. These are unusual findings for PTCL involving the colon.

A case of intestinal T-cell lymphoma involving the rectum was previously reported in Korea. However, although the T-cell markers were expressed, the exact subtype was not defined and whether the patient was non-AIDS or not was not clearly mentioned in that case. Another difference of that previous case was the involvement of the entire gastrointestinal tract from the esophagus to the rectum rather than skipping anorectal and gastric involvement, which was true in seen in our case. Our case had some characteristic findings. The patient showed concurrent anorectal and gastric involvement, he was not an AIDS patient, and there were no extensive ulceration in spite of PTCLu.

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