MULTIFOCAL SYNCHRONOUS SIGNET RING CELL CARCINOMA IN ULCERATIVE COLITIS- AN UNCOMMON PRESENTATION

Manjot Kaur¹, Kanwaljit Kaur²

¹Professor, Department of Surgery, Adesh Medical College.
²Assistant Professor, Department of Pathology, Fortis Hospital.

ABSTRACT

BACKGROUND

Ulcerative colitis is associated with an increased risk for developing colorectal cancer and precancerous dysplastic epithelial changes. We present a case report of patient with ulcerative colitis of 22 years duration who presented with signet ring cell carcinoma lesion at splenic flexure and adenocarcinoma at anorectal junction and Linitis plastica of descending colon and paracolic abscesses. Almost all the complications in one patient makes worthy of presentation.

KEYWORDS

IBD (Inflammatory Bowel Disease), PSC (Primary Sclerosing Cholangitis), Signet Ring Cell Carcinoma, Dysplasia, Adenocarcinoma, Linitis Plastica.


BACKGROUND

Colon is the state of third highest incidence of new malignancies and second only to lung cancer in mortality. The ulcerative colitis colorectal carcinoma link is well established. The factors that govern the risk of this most serious complication are duration of the disease and extent of involvement of the colon.

Incidence of colon cancers in patients with extensive ulcerative colitis is 5% - 10% with disease duration of 20 years; 15% at 25 years and 30% at 35 years. Peak age of manifestation of ulcerative colitis is 20 - 40 years with no gender preponderance.[1] Chronic bowel infections, allergy to certain food articles, autoimmune factors, gender preponderance and emotional disturbances[2] are enlisted among various etiological factors; 10% - 15% of ulcerative colitis have a first degree relative with IBD. The extent of involvement of colon has a relation with occurrence of carcinoma colon.

Signet ring cell carcinoma has worse prognosis than mucinous adenocarcinoma.[3,4] Mucinous and signet ring cell carcinoma are of much higher incidence in patients with extensive ulcerative colitis than in general population.

Case Report

A 42-year-old male (Mohan Singh) with long history (22 years) of ulcerative colitis (diagnosed on biopsy) of ulcerative colitis on irregular treatment presented with septic shock. On examination he was severely anaemic, dehydrated and febrile. Abdomen was distended with generalised tenderness. On rectal examination, hard circumferential growth at anorectal junction was felt which was friable and bled on touch. Plain x-ray abdomen showed markedly distended caecum, ascending colon and transverse colon with delayed movement of contrast across splenic flexure, loss of haustrations in descending colon with strictures and acute narrowing in lower rectum. Biopsy of anorectal growth was taken, which was suggestive of adenocarcinoma.

Exploratory laparotomy revealed grossly dilated caecum, ascending colon and hepatic flexure with dense adhesions and pus pockets around splenic flexure. Descending colon was thick and tabular, sigmoid colon and rectum were collapsed. Liver was normal. Multiple significantly enlarged lymph nodes were present along the mesentery. Proctocolectomy with end ileostomy was done. Gross examination of specimen showed ulcerofungating growth (8.5 cm to 6.5 cm) at splenic flexure obstructing the lumen and another fungating growth at lower end rectum extending up to anorectal junction with multiple lymph nodes, which were reactive (Image 1).

Microscopic examination revealed that signet ring cell carcinoma at splenic fixtures with linitis plastica of descending colon, while growth at anorectal region was adenocarcinoma (Image 2 and 3).

Post-operative recovery was uneventful. Health of the patient improved remarkably, 2 years followup uneventful.

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Corresponding Author:
Manjot Kaur,
#145, D-Block,
Ranjit Avenue,
Amritsar-143001, Punjab.
E-mail: manjot888@gmail.com
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Image 1
Colorectal carcinoma represents an important and dreaded complication of long-standing IBD. The overall incidence of neoplasia in patients with pancolitis is 1% per year after 10 years so that cumulative risk of cancer is 10% by 20 years duration of disease.[5] Various risk factors of colorectal cancer in ulcerative colitis are enlisted as anatomic extent of the disease, duration of disease, age at onset of disease, primary sclerosing cholangitis. Disease activity, chromosomal instability, microsatellite instability are well documented pathways of ulcerative colitis related colorectal cancer.[6-7]

The cancer tends to be more advanced, because they are difficult to be detected at an early stage. Approximately, 35% are Duke's C and D category lesions. Certain distinguishing features of colon cancer in ulcerative colitis is that it occurs at a younger age, is more anaplastic, arises from flat mucosa, infiltrates broadly and is more often synchronous. Because of the potential impact on life expectancy and quality of life, optimising strategies are being suggested to reduce the risk of developing colorectal cancer in patients with IBD. These include colonoscopic surveillance, chemoprevention and surgical prophylaxis.

Colonoscopic Surveillance

The goal of this program is to identify cancer on their precursors (dysplasia) at an early stage, in which they are curable. Surveillance colonoscopy has been recommended by the American Society of Gastrointestinal Endoscopy after seven years of disease in patients with pancolitis and after ten years of disease in left-sided colitis. Colonoscopy and biopsy for dysplasia should be performed every year.

Chemoprophylaxis

Aspirin and non-steroidal anti-inflammatory drugs (5-aminosalicylates) may offer a chemoprotective effect. Several studies have reported a colorectal risk reduction in patients with IBD taking folate supplements.[8,9] Interestingly, the use of ursodiol has shown 74% reduction in risk of colorectal carcinoma in patients with PSC-ulcerative colitis.[10]

Surgical Prophylaxis

The most effective way of preventing colorectal cancer in ulcerative colitis patients is to carry out proctocolectomy once dysplasia has been identified, because the risk of cancer in presence of dysplasia is 30%. Nowadays, ideal technique is sphincter conserving approach. But in our patient as there was growth at anorectal junction, so proctocolectomy with end ileostomy had to be carried out.

REFERENCES