Adenocarcinoma of the Colon in a 10-year-old child

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Summary
Colonic adenocarcinoma is rare in children, more so in the first decade of life. Owing to the non-specificity and vagueness of symptoms, most patients in this age group present with advanced malignancy. The authors report a case of adenocarcinoma of the colon in a 10-year-old female child presenting with a history of abdominal pain, distention and constipation. Barium enema highlighted a narrowing of the distal ascending colon with laparotomy revealing a growth involving the caecum and ascending colon. Histology confirmed it as a poorly differentiated mucin secreting adenocarcinoma. A right hemicolectomy with end to end ileo-transverse anastomosis was performed. Four months later, she presented with symptoms of intestinal obstruction. Intraoperatively, widespread metastatic nodules with recurrent tumour were noted. It is important for clinicians to be aware that colorectal carcinoma does occur in children, and age alone should not be a basis for its exclusion. This case is presented to draw attention to a disease in which the prognosis, by wider recognition of its occurrence in childhood, might be improved.

Introduction
Colorectal carcinoma is a common visceral malignancy in the adult age group. Few cases have been reported in children and adolescents. Clinical features in childhood are non-specific and therefore tend to be ignored or misinterpreted (1). Only about 400 cases involving subjects less than 20 years were reported in the literature by 1988, with less than 12 patients below the age of 10 years and the youngest nine months old at the time of diagnosis (2-11). Genetic instability without hereditary nonpolyposis colorectal cancer (HNPCC) is a major mechanism of causation (12,13). In contrast to adult patients, childhood adenocarcinoma is associated with a poorer prognosis and a high index of suspicion is required for early diagnosis.

Case Report
A ten year old girl presented to our facility with a four months history of abdominal pain associated with distension, constipation, anorexia, weight loss and post-prandial emesis. She had received various treatments with only partial relief, for the symptoms and results of stool investigations repeatedly showing pus cells. Her general condition was good. She was not pale and had no jaundice, lymphadenopathy or dehydration. Her vital signs were as follows: pulse 78/minute, respiratory rate 18/minute, blood pressure 110/80 mmHg and temperature 36.5°C.

Examination of the abdomen revealed a poorly circumscribed and immobile mass in the right iliac fossa. The bowel sounds were present and normal. The liver, spleen and rectum were normal as was the rest of the systemic examination.

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CASE REPORT

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Figure 1: Free flow of contrast up to the ascending colon

Figure 2: "Apple core" lesion of colon carcinoma with narrowing of distal ascending colon

Figure 3: Right iliac fossa mass on breaching the abdomen
on auscultation. Appropriate antibiotic treatment was initiated and the rest of her post operative recovery was uneventful. At pathology, the total length of resected gut was 35 cm with the tumour measuring 9 x 6 cm. The tumour involved the entire thickness of the wall circumferentially constricting the caecum and ascending colon. Enlarged mesenteric nodes were evident. No evidence of polyps was noted. The growth was a poorly differentiated mucin secreting adenocarcinoma at histology. It involved the whole colonic wall. The proximal and distal margins of resection were free of tumour. There was also involvement of mesentry, and blood vessels. Five lymph nodes were identified, with level II lymph node metastasis. The pathological stage was denoted T3, N2, Mx (Dukes stage C). The patient was then referred to an oncologist and was due to start on a course of chemotherapy: Folinic acid (leucovorin), 5-Fluorouracil, Oxaliplatin (FOLFOX regimen). However, she was readmitted four months later with features of intestinal obstruction. She had been unwell for a duration of two weeks prior to this admission. She was sick looking and the abdomen distended with an ill-defined right iliac fossa mass. Plain radiographs showed multiple lengthy air–fluid levels within the small intestines. Because of persistent obstructive symptoms, the abdomen was explored. The intraoperative findings this time included widespread metastatic nodules on the greater omentum, small intestines, the entire peritoneal lining and the liver. A 15cm diameter recurrent mass was found at the site of the previous anastomosis with dilated small bowel loops (fig. 5). Excision of the tumour was performed with primary ileo-transverse anastomosis and omentectomy. The patient showed poor response post operatively and died 1 week later.

**Discussion**

Adenocarcinoma of the colon and rectum is the most common cancer of the gastrointestinal tract. Generally it is a disease of older age groups beginning after the age of 40 to 45 years and peaking between 75 and 80 years (14,15). Colorectal cancer is extremely rare in childhood and adolescence, the incidence being 1.3 – 2 cases per million population (8,14,16).

Contrary to adult patients in whom a predisposing history of familial polyposis syndromes, hereditary nonpolyposis syndromes, ulcerative colitis, or colorectal cancer in relatives can be sought, most paediatric patients develop de novo carcinoma in a previously normal colon (6,8,17-19). In a category of patients with colorectal carcinoma at a very young age but who do not have a family history of HNPCC, results suggest that they harbour tumours with a defect characteristic of mismatch repair.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Adult</th>
<th>Children</th>
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<tbody>
<tr>
<td>Incidence</td>
<td>Intestinal obstruction in 16-35%</td>
<td>Intestinal obstruction in 70%</td>
</tr>
<tr>
<td>M:F ratio</td>
<td>1:1</td>
<td>2:1</td>
</tr>
<tr>
<td>Stage at presentation</td>
<td>50-60% in Dukes stage C &amp; D</td>
<td>60-80% in Dukes stage C &amp; D</td>
</tr>
<tr>
<td>Primary site</td>
<td>70-75% in rectosigmoid region</td>
<td>30-35% in rectosigmoid region</td>
</tr>
<tr>
<td>Histopathology</td>
<td>5% mucinous</td>
<td>50% mucinous</td>
</tr>
<tr>
<td>Surgical resectability</td>
<td>90-95%</td>
<td>40%</td>
</tr>
<tr>
<td>Ovarian metastasis</td>
<td>4%</td>
<td>22%</td>
</tr>
<tr>
<td>CEA level</td>
<td>Important tumour marker</td>
<td>Not reliable</td>
</tr>
<tr>
<td>Delay in diagnosis</td>
<td>15%</td>
<td>60%</td>
</tr>
<tr>
<td>Prognosis</td>
<td>Better</td>
<td>Poor</td>
</tr>
</tbody>
</table>

Table I-Colorectal adenocarcinoma: Differences between Adults and Children (1,5,16,20,22,23)
(MMR) deficiency and that many of these have germline mutations of an MMR gene (12,13).

In children and adolescents, it is associated with a significantly worse survival rate likely because of delay in the diagnosis, advanced stage of the disease at presentation, with approximately 60-80% of patients at Dukes stage C or D and poor histological differentiation of the malignancy (1,5-8,20).

Symptoms are related to the primary site of the tumour within the large bowel. In children and adolescents, as with our patient, these may be vague abdominal discomfort, changes in bowel habit, anaemia and gastrointestinal bleeding. These vague symptoms and decreased awareness among clinicians contribute to a delay in diagnosis (1,21).

The differences in duration of symptoms, primary site, pathological findings, stage and prognosis between adults and children is striking as seen in Table I. Histologically, mucin-producing or signet ring adenocarcinoma is the predominant cell type occurring in 50% of paediatric cases as compared to 5% in adults (5,6,22). The mucin absorbs water, swells, and invades tissues, thus promoting tumour growth and spread of malignant cells. It also interferes with the immune recognition of carcinoma cells by the mucopolysaccharide coating (11,16,20,24).

Patients may test positive for occult blood in stool. However, this has not proven to be of significant value in the treatment of paediatric patients (4). Levels of carcinoembryonic antigen (CEA) should be determined despite the fact that fewer than 75% of colon carcinomas in children produce it. It is a useful tool in identifying recurrent disease after resection (23,25).

Surgery is the mainstay of treatment. Radical resection of the involved colon and its mesentery, usually in the form of a hemicolectomy, is the treatment of choice. The role of adjuvant chemotherapy in children and adolescents is controversial. Adjuvant chemotherapy has not been reported to result in cure in metastatic disease. Similarly, radiation has little to offer except when the tumour involves the rectosigmoid and anal area (1).

**Conclusion**

Based on age alone, cancer of the colon should not be excluded as a clinical diagnosis. We emphasize that the possibility of a malignant colorectal tumour should be considered for any child with signs and symptoms of intestinal obstruction, intractable abdominal pain, alteration in bowel habits and gastrointestinal bleeding. This case is presented to draw attention to a disease in which the prognosis might be improved, by the wider recognition of its occurrence in childhood.

**Acknowledgements**

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**References**

genetic polymorphisms. Mutation Research 2007;635:118-45