Base of tongue varices associated with portal hypertension

P Jassar, M Jaramillo, D A Nunez

Abstract
A symptomatic case of tongue base varices in a patient with portal hypertension secondary to liver cirrhosis is presented. There are no previously documented cases in the world literature. Oesophageal varices may not be the only source of expectorated blood in a patient with portal hypertension. (Postgrad Med J 2000;76:576–577)

Keywords: portal hypertension; lingual; tongue; varicose vein

Case report
An 82 year old woman with known portal hypertension secondary to cirrhosis of the liver was referred to the otolaryngology outpatient department with a two month history of daily haemoptysis and bloodstained pharyngeal secretions; this occurred mostly on early morning coughing. There was no history of weight loss, dysphagia, dysphonia, or throat pain. She had already been investigated for a pulmonary cause of the haemoptysis and none was found. In keeping with her history, liver function tests and the prothrombin time were abnormal. She also had a history of well controlled essential hypertension. Chest radiography showed cardiomegaly, but the jugular venous pressure was not raised and there was no other clinical manifestation of heart failure.

Indirect laryngoscopy revealed varicose vessels in the tongue base, mainly on the left side (see fig 1). These appeared friable and one area revealed a propensity to bleed on examination. The rest of the ear, nose, and throat examination was normal.

After perioperative cover with fresh frozen plasma, vitamin K, and tranexamic acid she underwent ablation of the varicosities using a 15 watt continuous carbon dioxide laser under general anaesthesia (see fig 2). She had an uneventful recovery and has remained symptom free.

Anatomy
The dorsal lingual veins drain the tongue base through two or more tributaries. These course inferiorly, join together and form the lingual vein, which accompanies the lingual artery. These vessels pass between genioglossus and hyoglossus and the vein empties into the internal jugular vein just above the level of the greater cornu of the hyoid bone.1

PORTOSYSTEMIC ANASTOMOSIS
Because the portal system has no valves, portal hypertension results in shunting of blood through anastomotic communications with the systemic venous system. Recognised connections are2:

(1) Lower oesophageal veins—the branches of the left gastric vein anastomose profusely with the azygos and hemiazygos veins. Increased shunting results in oesophageal varices within the mucosa of the lower third of the oesophagus.

(2) Periumbilical veins—veins contained in the falciform ligament anastomose with superior and inferior epigastric veins of the anterior abdominal wall. Excessive dilatation of these veins are evident as caput medusa.

(3) Haemorrhoids—the superior rectal vein anastomoses with the middle rectal vein which drains into the internal iliac vein. In addition the middle rectal veins anastomose with the inferior rectal veins which drain into the internal pudendal vein. Increased shunting results in internal and external haemorrhoids.
Cefuroxime induced lymphomatoid hypersensitivity reaction

S A M Saeed, M Bazza, M Zaman, K S Ryatt

Abstract
An 84 year old women developed erythematous blotchy erythema and purpuric rashes over the lower limbs three days after being started on intravenous cefuroxime for acute diverticulitis. A skin biopsy specimen showed a mixed infiltrate of lymphoid cells and eosinophils; many of the lymphocytes were large, pleomorphic, and showed a raised mitotic rate. Immunohistochemistry showed the infiltrate to be T cell rich, with all the large cells being CD30 positive. Typical mycosis fungoides cells, marked epidermotropism, and Pautrier's abscesses were not seen. The rash disappeared 10 days after cessation of cefuroxime and the patient remained asymptomatic 15 months later. This apparent cutaneous T cell lymphoma-like reaction is best described as lymphomatoid vascular reaction. The drug induced immune response with an atypical cutaneous lymphoid infiltrate mimics a cutaneous pseudolymphoma.

Case report
An 84 year old women, with established diverticula disease, presented as an emergency with a short history of fever, acute abdominal pain, and diarrhoea. On examination she was found to be toxic, febrile, dehydrated, with a tachycardia of 110/min and blood pressure 80/40 mm Hg. The left iliac fossa was tender. Haemoglobin, full biochemistry profile, and serum amylase, chest radiography, plain abdominal radiography, and an abdominal ultrasound scan were normal. Her white cell count was raised at 20.4 × 10^9/l. The patient was started on intravenous (IV) fluids, IV metronidazole 500 mg eight hourly, and IV cefuroxime 750 mg three times a day in a sequential manner. Three days later and after five doses of IV cefuroxime, purpuric and erythematous macular rashes developed over the lower limbs. Cefuroxime was considered to be responsible and was discontinued; metronidazole was continued for another week. The rashes were treated with flucinolone acetonide 0.00625% cream. Further investigations at this stage showed normal immunoglobulins, antinuclear factor, antineutrophilic cytoplasmic antibodies, and autoantibody profile. A haematoxylin and eosin stained section of representative skin biopsy showed a mixed dermal infiltrate of lymphoid cells and eosinophils, marked red blood cell extravasation indicative of ongoing small vessel damage, minimal focal vacuolar interface

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change and some basement membrane hyalini-
sation, both within the papillary dermis of the
inflamed skin and in the immediate vicinity. The
lymphoid cells were predominantly lym-
phocytic, with many of these being large and
polymorphic and showing hyperchromatic and
irregular nuclei and mitoses (fig 1). Epidermo-
tropism was only mild and focal and there were
no Pautrier’s abscesses. Immunohistochemistry
showed the infiltrate to be T cell rich, with
virtually all of the large cells being CD30
positive (fig 2). Direct immunofluorescence
showed fine linear deposition of IgG along the
dermoepidermal junction.

Pyrexia, diarrhoea, tachycardia, and hypo-
tension improved rapidly with supportive
measures and the rashes began to fade quickly
and cleared within seven days. She was
completely asymptomatic and clear of all
rashes when reviewed more than 15 months
after the initial episode.

Discussion

Cephalosporins are widely used for the treat-
ment of septicaemia, pneumonia, meningitis,
biliary and urinary tract infection, and peri-
titis. Both the old and new third generations
of cephalosporins are generally well tolerated at
standard recommended dosage. The sequen-
tial cefuroxime regimen was chosen as it has
been shown not only to be an effective

Table 1 Comparison between microscopical features of Mycosis fungoides and
lymphomatoid vascular reaction

<table>
<thead>
<tr>
<th>Microscopical characteristic</th>
<th>Mycosis fungoides</th>
<th>Lymphomatoid vascular reaction</th>
</tr>
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<tbody>
<tr>
<td>Epidermotropism</td>
<td>Marked and diffuse</td>
<td>Minimal and focal</td>
</tr>
<tr>
<td>Pautrier’s microabscesses</td>
<td>Usually present</td>
<td>Usually absent</td>
</tr>
<tr>
<td>Eosinophils</td>
<td>Minimal or absent</td>
<td>Moderate</td>
</tr>
<tr>
<td>Basilar vaculopathy</td>
<td>Usually absent</td>
<td>Usually present</td>
</tr>
<tr>
<td>Marked spongiosis, dermal oedema, and keratinocyte necrosis</td>
<td>Usually absent papillary dermal fibrosis rather than oedema is the rule</td>
<td>Usually present</td>
</tr>
<tr>
<td>Intraepidermal population of cells</td>
<td>Markedly atypical cytomorphology</td>
<td>Mildly atypical cytomorphology</td>
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</tbody>
</table>

Figure 1 High power view, haematoxylin and eosin stain, demonstrating large, pleomorphic lymphoid cells with mitotic activity and lymphocytes in the background (original magnification × 40).

Figure 2 CD30 immunohistochemical stain demonstrating CD30 positive cells in the infiltrate (original magnification × 20).

Table 1 Comparison between microscopical features of Mycosis fungoides and lymphomatoid vascular reaction

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Colonic carcinoma after ureterosigmoidostomy

A Huang, G A D McPherson

Abstract
Urinary carcinogens promote late malignant transformation of the colon after a ureterosigmoidostomy. An unusual case is presented where, despite the early removal of the latter and hence cessation of urine flow, a colonic carcinoma developed at the site of previous anastomosis. The importance of surveillance of all patients who have undergone this procedure to avoid an iatrogenic cancer is emphasised. (Postgrad Med J 2000;76:579–581)

Keywords: rhabdomyosarcoma; ureterosigmoidostomy; colonic carcinoma

Colonic carcinoma arising from the site of a functioning ureterosigmoidostomy anastomosis is a recognised late complication. If the anastomosis is subsequently taken down with no further urine flow into the bowel, the risk of neoplasia is reduced. We describe a rare case where a colonic carcinoma developed coincidentally at the site of a previous ureterosigmoidostomy after a long latent period.

Case history
A 41 year old man presented with rectal bleeding without bowel habit changes. At the age of 3 he underwent cystectomy with ureterosigmoidostomy formation for a rhabdomyosarcoma. Due to recurrent pyelonephritis the ureters were divided near the sigmoid colon and implanted in an ileal conduit seven years later. Two short segments of ureters were left attached to the colon.

Physical examination was normal and a barium enema revealed a lesion in the sigmoid colon. At laparotomy a circumferential cancer was found at the exact site of previous ureteric implantation (fig 1). Histological examination revealed a moderately differentiated adenocarcinoma with two short segments of residual

and focal exocytosis of cells in the epidermis overlying the inflamed dermal papillae observed in this case are well recognised features of delayed type hypersensitivity reaction. Such a delayed type reaction with a cutaneous T cell lymphoma-like morphology has been described as lymphomatoid vascular reaction.16 Most of the large lymphocytic cells in the infiltrate stained CD30 positive. In the context of neoplasia, CD30 is positive in Reed-Sternberg cells of Hodgkin's disease, the cells of lymphomatoid papulosis and in anaplastic large cell lymphoma (Ki1 lymphoma). This has been shown to carry an excellent prognosis.10 11

The close temporal association of starting cefuroxime and the appearance of the rash, the cutaneous T cell lymphoma-like histology with more features of a lymphomatoid vascular reaction than mycosis fungoides and rapid resolution of rash on cessation of the suspected drug, all favour lymphomatoid hypersensitivity reaction to cefuroxime as the most likely diagnosis. This reaction pattern, not previously described with cephalosporins, appears to be a benign reaction, with there being no relapses for more than a year after cessation of the original offending drug. Our patient remains well and clear of her rash as well as showing no sign of developing any cutaneous or systemic lymphomatous pathology 15 months after her discharge.

We are grateful to Dr D Slater, Consultant Histopathologist, Walsall Manor Hospital for their advice on skin histology.

ureter (fig 2). Four of the 14 lymph nodes recovered contained metastatic deposits (Dukes C) and he received adjuvant chemotherapy with 5-fluorouracil and folinic acid.

Discussion
Colonic carcinoma developing at the site of ureteric implant was first described by Hammer in 1929. The incidence is 100 to 550 times that of the general population with an overall lifetime risk of 5%; if the diversion is performed before the age of 25 years, the risk increases to 7000-fold. The latency is between six to 50 years after the procedure with the mean time at 21 years; the median age at diagnosis is 33 years.

Urine in direct contact with colonic epithelium plays a pivotal part in the initiation of carcinogenesis at the suture line. Stewart proposed that dietary nitrates excreted in urine come into the presence of high concentrations of secondary amines when diverted into the colon, with resultant bacterial activation of carcinogenic N-nitroso compounds. Constant faecal stream does not appear to be a prerequisite as carcinomas have been described arising from isolated colonic loops used as a neobladder. Other theories of carcinogenesis including surgical and mechanical trauma, excess concentrations of electrolytes, and chronic irritation resulting in malignant transformation have not been proved.

In the present case the ureterosigmoidostomy had been defunctioned many years previously and the subsequent development of a colonic carcinoma was likely to be a clinical coincidence. It could be argued that there was a causal link between the ureteric implantation and the bowel carcinoma, especially as the latter developed at such young age. However as there was no urine flow during this period this is unlikely.

Lifelong surveillance is recommended for all patients who undergo ureterosigmoidostomy. Starling et al suggested that annual colonoscopy with faecal occult blood test should be started soon after ureterosigmoidostomy, with subsequent alternative urinary diversion if recurrent polyps, cancer, or dysplasia were found. A more complex regimen of a faecal occult blood test every three months after two years, an excretory urogram yearly after five years, and sigmoidoscopy or colonoscopy every five years has also been proposed.

Registries of patients are often undertaken in large centres but the long latency to cancer development with subsequent patient movement makes tracking of these patients difficult. Patients and their physicians should therefore be fully informed of the risks associated with this procedure so that appropriate surveillance could be arranged. Hospital specialists should have a high index of suspicion when a patient presents with a history of urinary diversion and hence the possibility of colonic malignancy. Failure to do so would prevent the early detection of an iatrogenic bowel cancer.

WEB SITE REVIEW

International Herpes Management Forum (www.IHMF.org)

This large site has the most comprehensive collection imaginable of monographs, original papers and pictures on herpesvirus infections. Not just herpes simplex virus and varicella but cytomegalovirus, Epstein-Barr virus, human herpes virus 6, 7, and 8 are all exhaustively covered.

Complete monographs can be downloaded from the Library section or alternatively there are shorter management guidelines on such topics as herpesvirus infections in pregnancy and clinical implications of latency. There is also a whole section entitled Molecular Biology—all you ever wanted to know but were afraid to ask!

The site’s welcome page implies that it will be useful to visitors ranging from specialists to patients but the format is heavily weighted in favour of health care professionals. Most patients would need a good medical dictionary at hand and the format is bland and rather uninviting for the casual visitor.

For specialists there is a journal club and there are regularly updated details of international virology meetings on another of the main sections. Also featured is a world map with links to herpesvirus organisations across the globe.

An Acrobat reader and PowerPoint are required to utilise large parts of the site but anyone with these facilities can download entire lecture presentations of high quality.

The site is sponsored by a pharmaceutical company but is not heavily promotional. Highly recommended.

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Colonic carcinoma after ureterosigmoidostomy

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