Familial adenomatous polyposis (FAP) is one of two well described forms of hereditary colorectal cancer. The primary cause of death from this syndrome is colorectal cancer which inevitably develops usually by the fifth decade of life. Screening by genetic testing and endoscopy in concert with prophylactic surgery has significantly improved the overall survival of FAP patients. However, less well appreciated by medical providers is the second leading cause of death in FAP, duodenal adenocarcinoma. This review will discuss the clinicopathological features, management, and prevention of duodenal neoplasia in patients with familial adenomatous polyposis.

FAMILY ADENOMATOUS POLYPOSPHYS

FAP is an autosomal dominant disorder caused by a germline mutation in the adenomatous polyposis coli (APC) gene. FAP is characterised by the development of multiple (>100) adenomas in the colorectum. Colorectal polyposis develops by age 15 years in 50% and age 35 years in 95% of patients. The lifetime risk of colorectal carcinoma is virtually 100% if patients are not treated by colectomy.1

Patients with FAP can also develop a wide variety of extraintestinal findings. These include cutaneous lesions (lipomas, fibromas, and sebaceous and epidermoid cysts), desmoid tumours, osteomas, occult radio-opaque jaw lesions, dental abnormalities, congenital hypertrophy of the retinal pigment epithelium, and nasopharyngeal angiofibroma. In addition, FAP patients are at increased risk for several malignancies, such as hepatoblastoma, pancreatic, thyroid, biliary-tree, and brain tumours.1

Other gastrointestinal manifestations commonly found in FAP patients are duodenal adenomas, and gastric fundic gland and adenomatous polyps. Of concern, duodenal cancer is the second leading cause of death after colorectal cancer in these individuals.

EPIDEMIOLOGY OF DUODENAL POLYPS AND CANCER

After the colorectum, the duodenum is the second most commonly affected site of polyph development in FAP (fig 1).2 3 Duodenal adenomas can be found in 30–70% of FAP patients2–4 and the lifetime risk of these lesions approaches 100%.4 5

Duodenal/periampullary adenocarcinoma is the leading cause of death in FAP after colorectal cancer.4 These patients have a 100–330-fold higher risk of duodenal cancer compared with the general population.6 7 Of note, duodenal cancer is rare in the population, with an incidence of 0.01–0.04%.9 Estimates of the cumulative risk of developing duodenal cancer in FAP range from 4% at age 70 years to 10% at age 60 years.10 11 Recently, a large prospective five nation study set the cumulative incidence rate of duodenal cancer at 4.5% by age 57 years. The median age of duodenal cancer development was 52 years (range 26–58).4

UPPER GASTROINTESTINAL POLYP DISTRIBUTION AND TYPE

Polyps can be found throughout the duodenum, but the second and third portion and the periampullary region are the most commonly affected sites. This pattern probably reflects exposure of duodenal mucosa to bile acids,12 suggesting a role for these compounds in duodenal carcinogenesis.13 Most polyps in the duodenum are adenomas whereas polyps in the stomach are usually benign non-adenomatous fundic gland lesions. However, approximately 10% of gastric polyps are adenomas.1 12 Interestingly, Japanese and Korean FAP patients have a 3–4 times higher risk of gastric cancer compared with the general population.14 15 Whereas no increased risk has been found in Western countries.8 Besides polypoid neoplasia, flat adenomas can be found in the duodenum of approximately 30% of FAP patients and careful follow up of these lesions is recommended.16

GENOTYPE-PHENOTYPE CORRELATION IN DUODENAL POLYPOSPHYS

The cause of FAP is germline mutation of the APC gene. The APC gene is a tumour suppressor gene with 15 exons that encodes a 2843 amino acid protein with a molecular weight of 309 kDa. One third of all germline mutations occur in codons 1061 and 1309 (fig 2).1
Subsequent mutations in tumour suppressor genes (for example, p53 and SMAD4) and oncogenes (for example, K-Ras) lead to neoplastic progression of the adenoma-carcinoma sequence. Also, expression of important cell regulatory proteins is changed. One of these is cyclooxygenase 2 (COX-2), which is increasingly expressed in consecutive stages of the adenoma-carcinoma sequence.

The adenoma-carcinoma sequence, first identified for colorectal tumorigenesis, has been observed in the setting of duodenal carcinogenesis in patients with both FAP and sporadic disease. Spigelman and colleagues found a strong association between duodenal adenomas and duodenal cancer, showing that villous histology, moderate or severe dysplasia, and the presence of stage IV duodenal polyps were associated with malignant change. Also, case reports of duodenal carcinoma development in or near adenomas have been described. Moreover, Kashiwagi et al noted p53 overexpression in 25% of tubular, 72% of tubulovillus/villous adenomas, and 100% of duodenal carcinomas, and K-Ras codon 12 mutations have been detected in duodenal adenomas and carcinomas. In addition, SMAD4 mutations play a role in polypl development in the upper intestine in mice. Lastly, Resnick and colleagues demonstrated that transforming growth factor α (TGF-α) expression was greater in duodenal carcinomas than in adenomas, and that epidermal growth factor receptor (EGF-R) expression correlated with the degree of dysplasia in duodenal adenomas. These studies reveal that additional molecular alterations drive the transition of adenoma into carcinoma.

COX-2 is known to be an important mediator of colorectal neoplasia progression but expression of COX-2 has not been extensively studied in duodenal or upper gastrointestinal adenomas. Shirvani and colleagues found constitutive COX-2 expression in normal duodenum and oesophagus and significantly higher levels in oesophageal dysplastic tissues. Furthermore, these investigators showed that COX-2 expression in Barrett’s oesophagus increased in response to pulses of acid or bile salts. COX-2 expression is also elevated in gastric cancers.

CLASSIFICATION OF DUODENAL POLYPOSIS

The most useful system for rating the severity of duodenal polyposis was developed by Spigelman and colleagues. This classification describes five (0–IV) stages. Points are accumulated for number, size, histology, and severity of dysplasia of polyps (table 2). Stage I indicates mild disease whereas stages III–IV imply severe duodenal polyposis (fig 4). Approximately 70–80% of FAP patients have stage II or stage III duodenal disease, and 20–30% have stage I or stage IV disease. The estimated cumulative incidence of stage IV duodenal disease however is 50% at age 70 years.
Several investigators have shown that duodenal polyposis slowly progresses. One study followed 114 FAP patients for 51 months and found progression of polyps in size (26%), number (32%), and histology (11%).44 When individuals are followed for longer, duodenal polyps advance in Spigelman stage. Heiskanen and colleagues5 reported worsening polyposis over time followed for longer, duodenal polyps advance in Spigelman stages. One study followed 114 FAP patients for 51 months, and found that the median interval for progression by one stage was 4-11 years. Another group reported a stage change in 42% of patients with an average time of evolution by one stage of 3.9 years. The risk of developing stage III or IV disease exponentially increases after age 40 years.45

The Spigelman classification also correlates with risk of duodenal malignancy. Stages II, III, and IV disease are associated with a 2.3%, 2.4%, and 36% risk of duodenal cancer, respectively.46

**MANAGEMENT**

**Surveillance**

As noted, duodenal polyposis is invaluative over time. Consequently, surveillance of the upper gastrointestinal tract for the development of neoplasia by end and side viewing scopes is recommended by most authorities. One long term upper tract surveillance study of 114 FAP patients failed to prevent the development of duodenal adenocarcinoma in six patients.45 These findings emphasise the need to adjust the frequency of surveillance and to entertain surgical treatment with increasing severity of disease. Recommendations concerning the age of initiation of upper tract surveillance are not uniform. Some propose that screening for upper gastrointestinal disease should start at the time of FAP diagnosis.46 The NCCN (National Comprehensive Cancer Network), after review of all case reports of duodenal cancer in FAP patients, recommended a baseline upper gastrointestinal endoscopic examination at 25–30 years of age.47 Guidelines for continued endoscopic surveillance after baseline examination have been developed according to Spigelman stage by several authorities.46

In general, recommendations include stage 0 every 4 years; stage I every 2–3 years; stage II every 2–3 years; stage III every 6–12 months with consideration for surgery; and stage IV strongly consider surgery (table 3).

**Endoscopic treatment**

Endoscopic treatment options for duodenal lesions include snare excision, thermal ablation, argon plasma coagulation,
and photodynamic therapy (PDT). Most reports of endoscopic therapy use snare excision. However, duodenal adenomas are often flat non-polypoid structures and, therefore, difficult to remove using conventional snare excision. For these cases, prior submucosal saline/adrenaline infusion may facilitate removal and reduce the risk of haemorrhage and perforation. In addition, thermal ablation, argon plasma coagulation, or PDT may be suitable.

PDT is a non-thermal technique relying on the combined effect of a low power activating light and a photosensitising drug that is selectively retained within neoplastic tissue with minimal retention in surrounding normal tissue. Few reports of PDT for adenomas in the gastrointestinal tract exist. Loh and colleagues successfully applied PDT for resection of colorectal adenomas: 7/9 treated adenomas were eradicated. Others have used PDT for resection of neoplastic lesions in the upper gastrointestinal tract but results are disappointing (table 4).

Endoscopic treatment of duodenal neoplasia for Spigelman stage II and III polyposis has been pursued by some investigators. However, the benefit of this approach in eradicating duodenal polyposis is difficult to justify but may be useful in individual cases. Literature reports of endoscopic treatment for FAP patients with duodenal/ampullary polyps are summarised in table 4. These publications reveal that endoscopic treatment is usually insufficient to guarantee a polyp-free duodenum and fraught with complications. Recurrence rates of adenomatous tissue in duodenum of FAP patients treated endoscopically range from 50% to 100%. Lower recurrence was reported by Norton and colleagues but their study population also included patients with sporadic duodenal lesions. In

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**Table 2** Spigelman classification for duodenal polyposis in familial adenomatous polyposis

<table>
<thead>
<tr>
<th>Criterion</th>
<th>Points</th>
</tr>
</thead>
<tbody>
<tr>
<td>Polyp number</td>
<td>1–4</td>
</tr>
<tr>
<td>Polyp size (mm)</td>
<td>1–4</td>
</tr>
<tr>
<td>Histology</td>
<td>Tubular</td>
</tr>
<tr>
<td>Dysplasia</td>
<td>Mild</td>
</tr>
</tbody>
</table>

Stage 0, 0 points; stage I, 1–4 points; stage II, 5–6 points; stage III, 7–8 points; stage IV, 9–12 points.

**Table 3** Recommendations for management of duodenal polyposis in familial adenomatous polyposis, adjusted to the Spigelman stage of duodenal polyposis

<table>
<thead>
<tr>
<th>Spigelman stage</th>
<th>Endoscopic frequency</th>
<th>Chemoprevention</th>
<th>Surgery</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stage 0</td>
<td>4 years</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Stage I</td>
<td>2–3 years</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Stage II</td>
<td>2–3 years</td>
<td>+/-</td>
<td>No</td>
</tr>
<tr>
<td>Stage III</td>
<td>6–12 months</td>
<td>+/-</td>
<td>+/-</td>
</tr>
</tbody>
</table>

Figure 4  Spigelman stages of duodenal polyposis. (A) Stage I. (B) Stage II. (C) Stage III. (D) Stage IV.
Summary, endoscopic treatment appears useful in individual cases but follow up remains necessary and surgical intervention is often indicated in patients with more severe polyposis.

**Surgery**

Surgical options utilised to treat duodenal polyposis include local surgical treatment (duodenotomy with polypectomy and/or ampullectomy), pancreas and pylorus sparing duodenectomy, and pancreaticoduodenectomy. There are no randomised studies published to help guide surgical selection.

Publications of local surgical treatment with duodenotomy for duodenal polyposis in FAP patients are summarised in table 5. This surgery has proven insufficient to guarantee a cure. More radical surgery, in the form of classical pancreaticoduodenectomy, has been indicated for patients with severe polyposis (stage IV), failed endoscopic or local surgical treatment, and carcinoma development. Others recommend consideration of surgery in patients with stage III polyposis.

**Pharmacological treatment**

Non-steroidal anti-inflammatory drugs (NSAIDs) regress colorectal adenomas in FAP patients. The value of these agents for duodenal polyposis regression is unclear. Studies of duodenal adenoma regression have primarily utilised sulindac (NSAID) and selective COX-2 inhibitors (table 7). Low recurrence rates of polyposis have been reported with these procedures (table 6). The specific choice of procedure appears related to local expertise and the site of polyp involvement.

Use of endoscopic retrograde cholangiopancreatography to evaluate biliary duct involvement in patients with ampullary lesions or those with laboratory test perturbations has been suggested to direct appropriate surgery. In the final analysis, the morbidity and mortality of these surgeries must be weighed against the risk of developing duodenal adenocarcinoma.

### Table 4 Endoscopic treatment for duodenal neoplastic lesions

<table>
<thead>
<tr>
<th>Author</th>
<th>Treatment</th>
<th>Follow up</th>
<th>Patients</th>
<th>Outcome</th>
<th>Postoperative</th>
</tr>
</thead>
<tbody>
<tr>
<td>Soravia</td>
<td>Endoscopic resection</td>
<td>4–34 months (mean 18)</td>
<td>6 FAP</td>
<td>Recurrence of duodenal adenomas in all 5 surviving patients</td>
<td>1 patient died of acute pancreatitis after endoscopic ampullectomy</td>
</tr>
<tr>
<td>Bertoni</td>
<td>Snare polypectomy</td>
<td>18 months</td>
<td>2 FAP</td>
<td>Recurrence in 1 patient, successfully retreated</td>
<td>1 oozing-type haemorrhage and 1 mild pancreatitis, controlled by conservative measures</td>
</tr>
<tr>
<td>Morpurgo</td>
<td>Snare polypectomy (3 cases)</td>
<td>6–24 months (mean 19)</td>
<td>5 FAP</td>
<td>Recurrence in 3 patients</td>
<td>No postoperative complications</td>
</tr>
<tr>
<td>Alarcon</td>
<td>Snare polypectomy and thermal contact ablation</td>
<td>14–83 months (mean 43.5)</td>
<td>3 FAP</td>
<td>Recurrence in 3 patients</td>
<td>NS</td>
</tr>
<tr>
<td>Heiskanen</td>
<td>Snare excision (5), YAG laser coagulation</td>
<td>0.4–15.1 years (median 6.8)</td>
<td>6 FAP</td>
<td>No significant difference in survival rate compared to polypectomy</td>
<td>Patient treated with YAG laser developed mild pancreatitis</td>
</tr>
<tr>
<td>Norton</td>
<td>Ampullary ablative therapy</td>
<td>1–134 months (median 24)</td>
<td>59 FAP, 32 sporadic</td>
<td>Return to normal histology in 66.7% of sporadic and 34% of FAP lesions</td>
<td>12 patients had mild complications, 3 severe complications: duodenal stenosis, 1 postoperative pseudocyst, 1 necrotising pancreatitis</td>
</tr>
<tr>
<td>Norton</td>
<td>Snare excision of papilla</td>
<td>2–32 months (median 9)</td>
<td>15 FAP, 11 sporadic</td>
<td>Recurrence rate of duodenal adenomas is 10%</td>
<td>2 minor bleedings, 4 mild pancreatitis, 1 duodenal perforation</td>
</tr>
<tr>
<td>Milroy</td>
<td>PDT with ALA or Photofrin</td>
<td>1–134 months (median 24)</td>
<td>4 FAP patients with duodenal polyps</td>
<td>Superficial necrosis and no polyp reduction after PDT with ALA</td>
<td>Mild skin photosensitivity using Photofrin</td>
</tr>
<tr>
<td>Regula</td>
<td>PDT with ALA</td>
<td>2 duodenal adenomas, 3 ampullary carcinomas</td>
<td>6 FAP</td>
<td>No significant difference in survival rate compared to polypectomy</td>
<td>Side effects included mild skin photosensitivity, nausea/vomiting, and transient increases in AST</td>
</tr>
<tr>
<td>Loh</td>
<td>PDT with HpD or Photofrin</td>
<td>3–50 months (median 5.5)</td>
<td>8 patients with 9 colorectal adenomas</td>
<td>7 adenomas successfully eradicated</td>
<td>No local complications</td>
</tr>
<tr>
<td>Abulafi</td>
<td>PDT with HpD</td>
<td>10 patients with ampullary carcinoma unsuitable for surgery</td>
<td>5 FAP</td>
<td>Remission for 8–12 months in 3 patients with small tumours. In 4 patients with small tumours bulk was reduced. No improvement in patients with extensive disease</td>
<td></td>
</tr>
</tbody>
</table>

NS, not stated; PDT, photodynamic therapy; ALA, 5-aminolaevulinic acid; HpD, haematoporphyrin derivate or Photofrin; FAP, familial adenomatous polyposis.
Table 5 Local surgical treatment (duodenotomy with polypectomy and/or ampullectomy) for duodenal neoplastic lesions

<table>
<thead>
<tr>
<th>Author</th>
<th>Treatment</th>
<th>Follow up</th>
<th>Patients</th>
<th>Outcome</th>
<th>Postoperative</th>
</tr>
</thead>
<tbody>
<tr>
<td>Soravia†</td>
<td>Duodenotomy with polypectomy (1) or ampullectomy (4)</td>
<td>4–34 months (mean 18)</td>
<td>5 FAP</td>
<td>Recurrence in 4 patients. 1 patient died of cancer</td>
<td>1 transient duodenal fistula</td>
</tr>
<tr>
<td>Morpurgo†</td>
<td>Transduodenal ampullectomy (1) or polyp excision (1)</td>
<td>6–24 months (mean 19)</td>
<td>2 FAP</td>
<td>Recurrence in 1 patient</td>
<td>1 severe pancreatitis</td>
</tr>
<tr>
<td>Alarcon†</td>
<td>Local resection</td>
<td>8–33 months (mean 20.2)</td>
<td>5 FAP</td>
<td>Recurrence in 4 patients. 1 had progressive metastatic adenocarcinoma</td>
<td>NS</td>
</tr>
<tr>
<td>Heiskanen†</td>
<td>Duodenotomy</td>
<td>0.4–15.1 years (median 6.8)</td>
<td>15 FAP</td>
<td>No significant difference in Spigelman stage preoperative and at latest endoscopy</td>
<td>No postoperative complications</td>
</tr>
<tr>
<td>Penna†</td>
<td>Duodenotomy with polypectomy</td>
<td>5–36 months (mean 13.3)</td>
<td>12 FAP</td>
<td>Recurrence in 12 patients</td>
<td>NS</td>
</tr>
<tr>
<td>Penna†</td>
<td>Duodenotomy with polypectomy</td>
<td>36–72 months (mean 53)</td>
<td>6 FAP</td>
<td>Recurrence in 6 patients</td>
<td>1 cholecystectomy for cholecystitis, 2 duodenal fistulas</td>
</tr>
<tr>
<td>de Vos tot</td>
<td>Duodenotomy with polypectomy</td>
<td>4–13 months (mean 11)</td>
<td>8 FAP</td>
<td>Recurrence in 6 patients</td>
<td>1 minor morbidity†</td>
</tr>
<tr>
<td>de Vos tot</td>
<td>Duodenotomy with polypectomy</td>
<td>5–103 months (mean 29)</td>
<td>22 FAP</td>
<td>Recurrence in 17 patients. 1 death from metastatic disease</td>
<td>1 minor morbidity†</td>
</tr>
<tr>
<td>Ruo†</td>
<td>Duodenotomy with polypectomy</td>
<td>35 months</td>
<td>1 FAP</td>
<td>Gastric cancer arising from a polyp at 35 months</td>
<td>No postoperative complications</td>
</tr>
<tr>
<td>Farnell§</td>
<td>Transduodenal local excision</td>
<td>10 years</td>
<td>53 sporadic and FAP patients</td>
<td>Recurrence rate of 32% at 5 years and 43% at 10 years of follow up</td>
<td>3 pancreatitis, 3 leaks, 2 delayed gastric emptying, 2 ileus, 1 fluid overload</td>
</tr>
</tbody>
</table>

*That is, wound infection, atelectasis, or urinary tract infection.
NS, not stated; FAP, familial adenomatous polyposis.

Table 6 Pancreaticoduodenectomy and pylorus or pancreas preserving duodenectomy for duodenal neoplastic lesions

<table>
<thead>
<tr>
<th>Author</th>
<th>Treatment</th>
<th>Follow up</th>
<th>Patients</th>
<th>Outcome</th>
<th>Postoperative</th>
</tr>
</thead>
<tbody>
<tr>
<td>Soravia†</td>
<td>Pancreaticoduodenectomy</td>
<td>NS</td>
<td>1 FAP</td>
<td>Unknown</td>
<td>NS</td>
</tr>
<tr>
<td>Morpurgo†</td>
<td>Pancreaticoduodenectomy</td>
<td>NS</td>
<td>4 FAP</td>
<td>No recurrence reported</td>
<td>Increased number of bowel movements. One patient required pouch excision and end ileostomy to control diarrhoea. 3 patients experienced weight loss, 1 patients had episodes of pancreatitis</td>
</tr>
<tr>
<td>Alarcon§</td>
<td>Pancreas sparing duodenectomy</td>
<td>40–50 months (mean 45.7)</td>
<td>3 FAP</td>
<td>No recurrence. Two of three patients had a small tubular adenoma in the duodenal bulb.</td>
<td>1 pancreatic fistula, 1 upper GI haemorrhage</td>
</tr>
<tr>
<td>Penna†</td>
<td>Pancreaticoduodenectomy</td>
<td>9–108 months (mean 42)</td>
<td>7 FAP</td>
<td>Severe duodenal polyposis. Only 1 patients with duodenal cancer survived &gt;4 years</td>
<td>1 pancreatic fistula, 1 upper GI haemorrhage</td>
</tr>
<tr>
<td>de Vos tot</td>
<td>Pancreaticoduodenectomy</td>
<td>7–96 months (mean 47)</td>
<td>23 FAP</td>
<td>Recurrence in 3 of 9, 3 died of metastatic disease</td>
<td>5 minor morbidity†, 12 major morbidity†, 1 patient died of postoperative complications</td>
</tr>
<tr>
<td>de Vos tot</td>
<td>Pancreaticoduodenectomy</td>
<td>2–15 months (mean 11)</td>
<td>6 FAP</td>
<td>Recurrence in 17 patients. 1 death from metastatic disease</td>
<td>1 minor morbidity†, 3 major morbidity†</td>
</tr>
<tr>
<td>de Vos tot</td>
<td>Pancreaticoduodenectomy</td>
<td>7–93 months (mean 45)</td>
<td>12 FAP</td>
<td>Recurrence in 3 of 9, 3 died of metastatic disease</td>
<td>1 minor morbidity†, 4 major morbidity†</td>
</tr>
<tr>
<td>Knowles‡</td>
<td>Pancreaticoduodenectomy</td>
<td>7–18 months (mean 70.5)</td>
<td>7 FAP</td>
<td>1 patient developed jejunal adenomas 12 years after operation</td>
<td>1 patient developed pancreatic ascites</td>
</tr>
<tr>
<td>Chung†</td>
<td>Duodenal adenectomy</td>
<td>0.5–3 years (mean 2.1)</td>
<td>4 FAP</td>
<td>No recurrence</td>
<td>1 gastric retention, 1 pancreatic fistula</td>
</tr>
<tr>
<td>Kalady§</td>
<td>Duodenectomy</td>
<td>10 years</td>
<td>3 FAP</td>
<td>1 had polyp recurrence in jejunum at 5 years of follow up</td>
<td>1 postoperative wound infection, 1 biliary leak</td>
</tr>
<tr>
<td>Ballaud†</td>
<td>Pancreaticoduodenectomy</td>
<td>24 and 28 months 2 FAP</td>
<td>No recurrence</td>
<td>NS</td>
<td>10 leaks, 4 delayed gastric emptying, 1 delirium tremens, 3 abscesses. 1 patient died from bleeding and sepsis related to hepaticojunostomy leak. Morbidity was higher after pancreas sparing duodenotomy</td>
</tr>
<tr>
<td>Farnell§</td>
<td>Pancreaticoduodenectomy</td>
<td>0.3–16 years (mean 5.6)</td>
<td>25 FAP</td>
<td>No recurrences</td>
<td></td>
</tr>
</tbody>
</table>

*That is, wound infection, atelectasis, or urinary tract infection.
†That is, anastomotic leakage, fistula formation, wound abscess, sepsis, or pancreatitis.
NS, not stated; FAP, familial adenomatous polyposis.
300 mg/day with calcium and calciferol revealed no effect on duodenal polyps in 15 patients who completed six months of treatment with sulindac.66

Richard and colleagues67 treated eight FAP patients with residual small periampullary polyps with sulindac 300 mg/day for at least 10 months. Sulindac was discontinued in three patients due to side effects. Follow up endoscopy was performed every six months or at discontinuation of treatment. None of the patients showed regression of polyps; three patients developed large polyps and one an infiltrating carcinoma while on this drug.

A large randomised trial by Phillips and colleagues,68 with statistical power to detect small differences, investigated the effect of the specific COX-2 inhibitor celecoxib on duodenal polyp number and total polyp area. A 14% decrease in polyp number was found after six months of celecoxib 800 mg/day (n = 32) compared with placebo (n = 17) which was not statistically significant. Paired assessment of endoscopic videotapes, however, revealed a significant difference (p = 0.033), although no effect on polyp area was noted.

Winde and colleagues69 preformed a prospective, controlled, non-randomised phase II dose finding study for sulindac. These investigators compared effects of sulindac suppositories (n = 28) with placebo (n = 10) on rectal and upper gastrointestinal adenomas in patients that underwent colectomy. They found complete or partial reversal of rectal polyps but no effects on duodenal and papillary adenomas.

Table 7 Familial adenomatous polyposis patients treated with sulindac, celecoxib, or refecoxib for duodenal adenomas

<table>
<thead>
<tr>
<th>Author</th>
<th>Treatment (dose/day)</th>
<th>Type of study</th>
<th>Duration</th>
<th>Patients</th>
<th>Outcome</th>
<th>Side effects</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nugent</td>
<td>Sulindac 400 mg</td>
<td>Randomised controlled clinical trial</td>
<td>6 months</td>
<td>11</td>
<td>Number of polyps ↓ in 5 patients (p = 0.12 vs placebo). Second evaluation: effect on small polyps (≤2 mm) (p = 0.02)</td>
<td>1 patient with indigestion</td>
</tr>
<tr>
<td>Debinski</td>
<td>Sulindac 300 mg</td>
<td>Randomised controlled clinical trial</td>
<td>6 months</td>
<td>15</td>
<td>No effect</td>
<td>No adverse events reported</td>
</tr>
<tr>
<td>Seow-Choen</td>
<td>Sulindac 300 mg</td>
<td>Clinical trial</td>
<td>10–24 months</td>
<td>5</td>
<td>No regression of small residual polyps; 3 patients developed large polyps; 1 breakthrough carcinoma</td>
<td>2 patients with abdominal cramp. 1 patient with upper GI bleeding 1 patient with allergic reaction. 1 patient with symptoms of dyspepsia 2 patients with mild gastritis due to NSAID</td>
</tr>
<tr>
<td>Richard</td>
<td>Celecoxib 800 mg</td>
<td>Randomised controlled clinical trial</td>
<td>6 months</td>
<td>30</td>
<td>Number of polyps ↓ compared to placebo (p = 0.03)</td>
<td>No effect on small polyps</td>
</tr>
<tr>
<td>Phillips</td>
<td>Celecoxib 25 mg</td>
<td>Randomised controlled clinical trial</td>
<td>6 months</td>
<td>6</td>
<td>Improvement in 2 patients with stage III polyposis; no effect in 4 patients; no effect in ursodeoxycholic acid group</td>
<td>No recurrence of duodenal polyps</td>
</tr>
<tr>
<td>Maclean</td>
<td>Refecoxib 50–300 mg</td>
<td>Prospective, controlled, non-randomised phase II dose finding study</td>
<td>Up to 4 years</td>
<td>xx</td>
<td>No effect on upper GI polyps</td>
<td></td>
</tr>
<tr>
<td>Park Day</td>
<td>Refecoxib 25 mg</td>
<td>Randomised controlled clinical trial</td>
<td>6 months</td>
<td>6</td>
<td>Improvement in 2 patients with stage III polyposis; no effect in 4 patients; no effect in ursodeoxycholic acid group</td>
<td>No recurrence of duodenal polyps</td>
</tr>
<tr>
<td>Theodore</td>
<td>Sulindac 300–400 mg</td>
<td>Case report</td>
<td>5 and 14 years</td>
<td>2</td>
<td>Sulindac normalised adenomatous ampulla and induced elimination of moderate dysplasia</td>
<td>No effect on gastric and small intestinal polyps</td>
</tr>
<tr>
<td>Windle</td>
<td>Sulindac 300–400 mg</td>
<td>Case report</td>
<td>4.5–5 years</td>
<td>2</td>
<td>No effect on gastric and small intestinal polyps</td>
<td></td>
</tr>
</tbody>
</table>

*The control group was treated with calcium and calciferol. †The control group was treated with ursodeoxycholic acid. NSAID, non-steroidal anti-inflammatory drug.

In conclusion, the results of NSAID and other compounds on regression or prevention of duodenal adenomas in FAP appear disappointing, although regression of small adenomas may occur.70

MOLECULAR MECHANISMS OF CHEMOPREVENTION WITH NSAIDS

Studies of chemoprevention/regression of duodenal polyps in FAP have primarily utilised NSAIDs. The action of these agents has been divided into COX dependent, mediated through inhibition of the COX enzymes, and COX independent, caused by direct actions of NSAIDs on different molecular mechanisms.

COX dependent mechanisms

NSAIDs are best known for inhibitory effects on COX-1 and COX-2, key enzymes in the conversion of arachidonic acid to prostaglandins (PGs) (fig 5). COX-1 expression occurs in...
most tissues whereas COX-2 is expressed in response to growth factors, lipopolysaccharide, cytokines, mitogens, and tumour promoters. PGs are involved in cellular functions such as angiogenesis and cell proliferation. Therefore, inhibition of PG synthesis could explain part of the antineoplastic effects of NSAIDs. Also, COX-2 inhibition has antiangiogenic effects, as confirmed by several different studies. COX-2 inhibition may also induce apoptosis, mainly via inhibition of PGE₂, and inhibit invasive properties of cancer cells. COX-2 was induced by coculture and promoted invasion in vitro that was inhibited by NSAIDs or RNAi against COX-2.

**CONCLUSIONS AND FUTURE DIRECTIONS**

With improvement in the management of colorectal disease and increased life expectancy, duodenal polyposis and malignancy have emerged as major health problems in patients with FAP. Although most patients eventually develop duodenal polyps, these lesions occur at later age and have lower potential for malignant change compared with colonic polyps. Moreover, duodenal adenomas seem less responsive to chemoprevention with NSAIDs than colonic counterparts.

Currently, the main treatment options for duodenal polyposis are frequent surveillance and targeted endoscopic treatment, adjusted by severity of duodenal lesions. However, these modalities alone cannot guarantee a polyp-free duodenum. In patients with severe disease, duodenotomy or duodenectomy may be necessary. Drug therapy of duodenal adenomas would be appropriate treatment but...
most published reports find no significant effect of NSAIDs or COX-2 inhibitors on duodenal adenoma regression.

Increasing insights into the molecular changes during the adenoma-carcinoma sequence in the duodenum may point to future treatment strategies. Duodenal mucosa is exposed to different environmental factors than that in the colon. Low pH and bile acids may affect growth of control and malignant potential of duodenal tumours.12 13 34 Little is known about the role of potential molecular targets for chemoprevention, including COX-2, PPARβ, TGF-β receptor type II, EGF-R, and inducible nitric oxide synthase. More powerful chemopreventive/regressive regimens could result from combinations of NSAIDs or COX-2 inhibitors with other drugs, such as selective inhibitors of receptor tyrosine kinases or EGF-R. Further study is needed to understand the molecular changes in duodenal adenoma development and identify molecular targets for chemoprevention and regression of duodenal polyposis.

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